For JEE & Other Engineering Entrance Examinations

CHEMISTRY MCQ. AULTIPLE-CHOICE-OUESTION BANK

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Preface

This book is meant for the IIT-JEE and other examinations in which multiple-choice questions are asked. The book is divided into five parts. The first three parts cover physical, inorganic and organic chemistry, topicwise. In each of these parts a chapter is devoted to assertion–reason questions. In the fourth part miscellaneous questions have been given. In the fifth part a number of test papers have been given. The last two parts form a very important part of a student's preparation.

In a book of this nature, a balance has to be achieved between the number of questions and the quality of the questions, especially because it is relatively easy to frame a very large number of multiple-choice questions. The questions in this book have been selected keeping three things in mind. First, the questions are such that they really test the understanding of the subject. Second, among themselves, the questions cover all the concepts. Third, the number of questions has been kept large enough to offer meaningful practice to the students. Wherever required, hints have been given. In various competitive entrance examinations including the IIT-JEE, questions with multiple correct options are also asked sometimes. We have included questions of this type also. It should be remembered that in such questions marks are awarded only if all the correct options are chosen and no incorrect option is chosen.

We would like to thank the editors and production staff at Bharati Bhawan for their efforts in bringing out this revised edition of the book. Suggestions from readers for the improvement of the book are always welcome.

Authors

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Part 1

Physical Chemistry

1

Atomic Structure

• Type 1 •

Choose the correct option. Only one option is correct.

1.	A Cr atom in its ground state has a 3d 5 4s 1 configuration and a Cu atom
	a 3d 10 4s 1 configuration. This is because a shell which is half-filled or
	completely filled is particularly

- (a) strongly exchange-destabilized
- (b) weakly exchange-stabilized
- (c) strongly exchange-stabilized
- (d) weakly exchange-destabilized

2.	A	d	-sh	ıell	conta	ining	four	unpa	ired	elec	ctro	ns	can	excl	nange
												_		_	

(a) four electrons

- (b) three electrons
- (c) sixteen electrons
- (d) six electrons

3. Based on quantum mechanical theory, the electronic configuration of Pd(Z=46) is

(a) $4d^9 5s^1$

(b) 4d ¹⁰

(c) $4d^8 5s^1$

(d) $4d^{10} 5s^2$

4. When a gold sheet is bombarded by a beam of α -particles, only a few of them get deflected whereas most go straight, undeflected. This is because

- (a) the force of attraction exerted on the α -particles by the oppositely charged electrons is not sufficient
- (b) a nucleus has a much smaller volume than that of an atom
- (c) the force of repulsion acting on the fast-moving α -particles is very small
- (d) the neutrons in the nucleus do not have any effect on the α -particles

- 5. Which of the following statements is incorrect for anode rays?
 - (a) They are deflected by electric and magnetic fields.
 - (b) Their *e*/*m* ratio depends on the gas in the discharge tube used to produce the anode rays.
 - (c) The e/m ratio of anode rays is constant.
 - (d) They are produced by the ionization of the gas in the discharge tube.
- 6. Rutherford's α-particle scattering experiment led to the conclusion that
 - (a) mass and energy are related
 - (b) the mass and the positive charge of an atom are concentrated in the nucleus
 - (c) neutrons are present in the nucleus
 - (d) atoms are electrically neutral
- 7. The radius of $_{13}^{27}$ Al will be
 - (a) 1.2×10^{-15} m

(b) 27×10^{-15} m

(c) 10.8×10^{-15} m

- (d) 3.6×10^{-15} m
- 8. The density of the nucleus of an atom is
 - (a) $2.4 \times 10^{15} \text{ kg m}^{-3}$
- (b) $2.4 \times 10^{19} \text{ kg m}^{-3}$
- (c) $2.4 \times 10^{17} \text{ kg m}^{-3}$
- (d) $2.4 \times 10^{14} \text{ kg m}^{-3}$
- 9. In an X-ray experiment, different metals are used as the target. In each case, the frequency v of the radiation produced is measured. If Z = atomic number, which of the following plots will be a straight line?
 - (a) v against Z

(b) $\frac{1}{y}$ against Z

(c) \sqrt{v} against Z

- (d) v against \sqrt{Z}
- **10.** In Moseley's equation $[\sqrt{v} = a(Z b)]$, which was derived from the observations made during the bombardment of metal targets with X-rays,
 - (a) *a* is independent of but *b* depends on the metal
 - (b) both a and b depend on the metal
 - (c) both a and b are independent of the metal
 - (d) b is independent of but a depends on the metal
- 11. Which of the following reactions led to the discovery of the neutron?
 - (a) ${}_{6}^{14}C + {}_{1}^{1}p \rightarrow {}_{7}^{14}N + {}_{0}^{1}n$
- (b) ${}_{5}^{11}B + {}_{1}^{2}D \rightarrow {}_{6}^{12}C + {}_{0}^{1}n$
- (c) ${}_{4}^{9}\text{Be} + {}_{2}^{4}\text{He} \rightarrow {}_{6}^{12}\text{C} + {}_{0}^{1}\text{n}$
- (d) ${}_{4}^{8}\text{Be} + {}_{2}^{4}\text{He} \rightarrow {}_{6}^{11}\text{C} + {}_{0}^{1}\text{n}$

12.	Let m	p be the mas	s of a p	roton, m _n	that	of a neutro	M_1	that of a $^{20}_{10}$ Ne	
	nucleus and M_2 that of a $^{40}_{20}$ Ca nucleus. Then								
	(a)	$M_2 = 2M_1$			(b)	$M_1 < 10(m_{\rm p}$	+ m _n)	
	(c)	$M_2 > 2M_1$			(d)	$M_1 = M_2$			
13.	the sa comm this sy	me mass as a	an elect	ron, but o Calculate t	ppos he va	site charge) o	orbiti ydbe	ticle which has R_{∞} ng round their R_{∞}	
	` /		()	•	` '	5.0	,		
14.	electro		nucleus	in the 1s				distance of an atom (a_0 = the	
		$1.5a_0$ and a_0		,	(b)	a_0 and $5a_0$			
	(c)	$1.5a_0$ and $0.5a_0$	a_0		(d)	a_0 and $0.5a_0$			
15.		late the mass of hydrogen				that the firs	t line	e in the Lyman	
	(a)	1.66×10^{-27}	kg		(b)	8.3×10^{-28} J	kg		
	(c)	6.68×10^{-27}	kg		(d)	3.34×10^{-27}	kg		
16.	The en	nergy of a 700)-nm ph	oton is					
	` '	1.77 eV			` '	2.47 eV			
	(c)	700 eV			(d)	3.57 eV			
17.		W radio trans ns per second			ıt a f	requency of	880 I	Hz. How many	
	(a)	1.71×10^{21}			(b)	1.71×10^{30}			
	(c)	6.02×10^{23}			(d)	2.85×10^{26}			
18.	The ra	atio of the e/m	values	of a proto	n an	d an α-parti	cle is		
	(a)	2:1	(b) 1:	1	(c)	1:2	(d)	1:4	
19.	(a) (b)	n of the follov A proton an A proton an	d a neu d deute	tron rium	lentio	cal values of	e/m?		
		Deuterium a		-					
20		An electron		-				1 (. /2	
20.		n of the follow	_	ın arrange			-	iue of e/m?	
	(a)	$n < \alpha < p < \epsilon$	=		(D)	e	11		

(d) $p < n < \alpha < e$

(c) n

(a) A positron(c) An α-particle

(a) $\frac{c^2}{v}$

(c) $\frac{mc^2}{h}$

21. Which particle among the following will have the smallest de Broglie

(b) A photon

(d) A neutron

(b) $\frac{hv}{mc}$

(d) νλ

wavelength, assuming that they have the same velocity?

22. The velocity of the de Broglie wave is given by

23. de Broglie wavelength is related to applied voltage as

	(a) $\lambda = \frac{12.3}{\sqrt{h}} \text{ Å}$	(b) $\lambda = \frac{12.3}{\sqrt{V}} \text{ Å}$	
	(c) $\lambda = \frac{12.3}{\sqrt{E}} \text{ Å}$	(d) $\lambda = \frac{12.3}{\sqrt{m}} \text{ Å}$	
24.	Find the de Broglie wavelengt 20 -m s ^{-1} wind.	n of a 1-mg grain o	of sand blown by a
	(a) 3.3×10^{-29} m	(b) 3.3×10^{-21}	m
	(c) 3.3×10^{-49} m	(d) 3.3×10^{-42}	m
25.	The momentum of a hydrogen a	tom is given by	
	(a) $p = \frac{hv}{c}$	(b) $p = \frac{hv}{mc^2}$	
	(c) $p = \frac{mv}{h}$	(d) $p = mvr$	
26.	For an electron, the product <i>vr</i> will be independent of the	(velocity × principa	l quantum number)
	(a) principal quantum numb	er	
	(b) velocity of the electron		
	(c) energy of the electron		
	(d) frequency of its revolution	n	
27.	Assume that the potential energy zero. Then its energy in the first e		in its ground state is
	(a) 13.6 eV	(b) 27.2 eV	
	(c) 23.8 eV	(d) 10.2 eV	
	The ratio of $E_2 - E_1$ to $E_4 - E_3$ fequal to	or the hydrogen ato	om is approximately
	(a) 10 (b) 15	(c) 17	(d) 12

29. The ionization energy of a hydrogen atom is 13.6 eV. The energy of the third-lowest electronic level in doubly ionized lithium (Z = 3) is

30. What is the most probable distance from the nucleus at which a 2p

electron will be found in the hydrogen atom?

(b) -54.4 eV

(d) -13.6 eV

(a) −28.7 eV

(c) -122.4 eV

		53 pm 212 pm			106 pm 26.5 pm		
21		•	l to oveito e l		•	a amound atata ta i	10
31.		excited state i		nydrogen	atom mom n	s ground state to i	ıs
	(a)	12.1 eV		(b)	10.2 eV		
	(c)	0.85 eV		(d)	12.75 eV		
32.	atom	is 2.34×10^{1} tion is	one of the l ⁴ Hz. The (b) 4	ines in the quantum (c)	number n_2	eries of a hydroge which causes th (d) 5	en is
33		ne spectra of	two elemen	ts are not i	dentical hec		
00.		the elements					
	(b)	they have di	ifferent mas	s numbers			
	(c)	their outerm	nost electron	s are at di	fferent energ	y levels	
	(d)	they have di	ifferent vale	ncies			
34.		avelength of	the third lin	e of the Ba	almer series f	or a hydrogen ato	m
	is	21			100		
	(a)	$\frac{21}{100R_{\infty}}$		(b)	$\frac{100}{21R_{\infty}}$		
	(-)	$\frac{21R_{\infty}}{100}$		(4)	$\frac{100R_{\infty}}{21}$		
	(c)	100		(a)	21		
35.	In wh	ich of the foll	owing trans	itions will	the waveler	gth be minimum?	•
	(a)	n = 6 to n = 4	1	(b)	n = 4 to $n =$	2	
	(c)	n = 3 to n = 1	1	(d)	n = 2 to $n =$	1	
36.	In wh	ich of the foll	owing is the	radius of	the first orbi	it minimum?	
		A hydrogen			A tritium at		
	(c)	Triply ionize	ed berylliun	n (d)	Doubly ion	ized helium	
37.		the electron the number o				he $n = 4$ to the $n =$:1
	(a)			(b)			
	(c)	3		(d)	4		

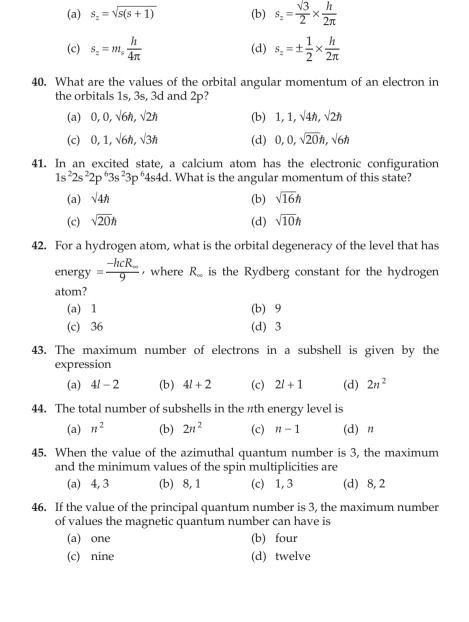
(a) n

(c) m₁

38. The quantum number not obtained from the Schrödinger wave equation

39. If a magnetic field is applied to the electron of a hydrogen atom in the *z*-direction, the *z*-component of the spin angular momentum is given by

(b) l(d) m_e



47. The correct set of quantum numbers for the unpaired electron of a

49. Four sets of values of quantum numbers (n, l, m and s) are given below. Which of these does not provide a permissible solution of the wave

(b) 2, 1, -1, $+\frac{1}{2}$

(d) $3, 0, 0, \pm \frac{1}{2}$

(c) photons

chlorine atom is (a) $2, 0, 0, +\frac{1}{2}$

(c) $3, 1, -1, \pm \frac{1}{2}$

(a) electrons

equation?

48. The Pauli exclusion principle is not applicable to

(b) positrons

(d) protons

	(a) 3	$5, 2, -2, \frac{1}{2}$			(b)	$3, 3, 1, -\frac{1}{2}$		
	(c) 3	$\frac{1}{2}$, 2, 1, $\frac{1}{2}$			(d)	3, 1, 1, $-\frac{1}{2}$		
50.		of the follow of an atom?		sets of qua	intum	numbers rep	presents the	highest
	(a) n	l=4, l=0, m	n=0,	$s = +\frac{1}{2}$	(b)	n = 3, l = 0, n	$m = 0, s = +\frac{1}{2}$	
	(c) n	l = 3, l = 1, n	n = 1,	$s = +\frac{1}{2}$	(d)	n = 3, l = 2, 1	$m = 1, s = +\frac{1}{2}$	
51.	The fou	r quantum :	numl	pers of the	valenc	e electron of	potassium a	ire
	(a) 4	$\frac{1}{2}$	(b)	$4, 0, 0, \frac{1}{2}$	(c)	$4, 1, 0, \frac{1}{2}$	(d) 4, 4, 0,	$\frac{1}{2}$
52.	The follatom.	owing sets	of q	uantum nı	umbers	s represent f	our electron	s in an
	(i) $n = 4$	4, <i>l</i> = 1 (ii)	n = 4,	l = 0 (iii)	n = 3,	l=2 (iv) $n=1$	= 3, l = 1	
		ontext, whi				presents the		reasing
	(a) (iv) < (ii) < (i	iii) < ((i)	(b)	(ii) < (iv) < ((i) < (iii)	
	(c) (i) < (iii) < (ii	i) < (i	v)	(d)	(iii) < (i) < (i)	(v) < (ii)	
53.	The tota	l number of	orbit	als in a she	ll with	principal qua	antum numb	er n is
	(a) 2	n	(b) 2	2n ²	(c)	n^2	(d) $n + 1$	
54.	If $m = n$ then	nagnetic qu	iantui	m number	and l	= azimuthal	quantum n	umber,
	(a) n	n = l + 2			(b)	$m = 2l^2 + 1$		
	(c) l	$=\frac{m-1}{2}$			(d)	l = 2m + 1		

55. Which of the following sets of quantum numbers represents the 19th

56. Which option gives the values of the quantum numbers for the 21st

(b) $4, 1, -1, +\frac{1}{2}$

(d) 3, 2, -2, $+\frac{1}{2}$

electron of chromium ($\check{Z} = 24$)?

electron of scandium (Z = 21)?

(a) $4, 0, 0, +\frac{1}{2}$

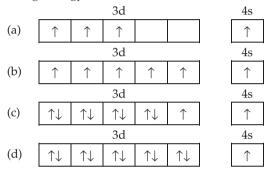
(c) $3, 2, 2, +\frac{1}{2}$

	(a)	$3, 1, 1, +\frac{1}{2}$	(b)	$3, 2, 2, +\frac{1}{2}$
	(c)	$3, 2, -2, -\frac{1}{2}$	(d)	$3, 2, 2, -\frac{1}{2}$
57.	The e	lectronic configuration of Fe 3+	is	
	(a)	$[Ar]3d^54s^1$	(b)	$[Ar]3d^64s^0$
	(c)	$[Ar]3d^{5}4s^{0}$	(d)	$[Ar]3d^64s^2$
58.	The n	umber of unpaired electrons ir	n Mn	$^{4+}(Z=25)$ is
	(a)	four	(b)	two
	(c)	five	(d)	three
59.	The c	onfiguration [Ar]3d 10 4s 2 4p 1 is	simi	ilar to that of
		carbon		oxygen
	(c)	nitrogen	(d)	aluminium
60.		ng the following, the configura	tion	$1s^2 2s^2 2p^5 3s^1$ is valid for the
		ground state of fluorine		excited state of fluorine
	(c)	excited state of neon	(d)	excited state of the O_2^- ion
61.	After	<i>n</i> p orbitals are filled, the next of	orbita	al filled will be
	(a)	(n+1)s	(b)	(n+2)p
	(c)	(n+1)d	(d)	(n+2)s
62.		value of the magnetic mome eton. The ion is	nt o	f a particular ion is 2.83 Bohr
	(a)	Fe ²⁺	(b)	Ni ²⁺
	(c)	Mn ²⁺	(d)	Co ³⁺
63.	Which	h of the following violates the I	Pauli	exclusion principle?
	(a)	$\uparrow\downarrow$ $\uparrow\downarrow$	(l	$\uparrow \qquad \uparrow \qquad \uparrow$
	(c)	$\uparrow\uparrow$ \uparrow \uparrow	(0	d)

64. Which of the following violates the Aufbau principle?

	2s	2p		2s		2p	
(a)	$\uparrow\downarrow$	$\uparrow\downarrow$ \uparrow	(b)	$\uparrow\downarrow$	$\uparrow\downarrow$	$\uparrow\downarrow$	\uparrow
	2s	2p		2s		2p	
(c)	$\uparrow \uparrow$	$\uparrow\downarrow$ $\uparrow\downarrow$ \uparrow	(d)	↑	$\uparrow\downarrow$	↑	\uparrow

65. Which of the following electronic configurations have the highest exchange energy?



- 66. Which of the following sets of orbitals is arranged in the correct order of increasing energy?
 - (a) 3d < 4s < 4p < 6s < 4d
- (b) 2s < 3d < 4p < 4f < 1s
- (c) 4s < 3d < 4p < 5s < 4d
- (d) 1s < 2s < 2p < 4d < 3f

67. The number of spherical nodes in 3p orbitals is

(a) one

(b) three

(c) two

(d) zero

68. In which of the following orbitals is there zero probability of finding the electron in the xy plane?

- (a) p_x
- (b) d_{1/2}
- (c) $d_{x^2-y^2}$ (d) p_z

69. Which of the following electronic configurations have zero spin multiplicity?



(b)

- (c)
- (d)

70. The radial distribution functions for all orbitals is given by

(a) n+l

(b) n-l-1

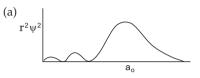
(c) n-l-2

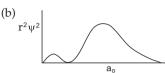
(d) n-l+1

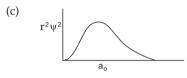
- 71. The angular distribution functions of all orbitals have
 - (a) *l* nodal surfaces

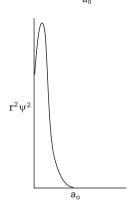
- (b) l-1 nodal surfaces
- (c) n + l nodal surfaces
- (d) n-l-1 nodal surfaces
- **72.** Which of the following radial distribution graphs correspond to l = 2 for the H atom?

(d)

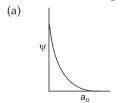


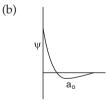


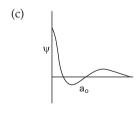


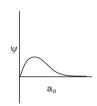


73. Which of the following graphs correspond to one node?









- **74.** Which of the following statements is incorrect with reference to the Zeeman effect?
 - (a) In a magnetic field, the energy of a particular atomic state depends on the values of m_l and n.

(d)

(c) The Zeeman effect involves the splitting of a spectral line of frequency v₀ into three components whose frequencies are

$$v_1 = v_0 - \frac{e}{4\pi m} B$$
, $v_2 = v_0$ and $v_3 = v_0 + \frac{e}{4\pi m} B$,

where *B* is the magnetic field applied on the spectral line.

- (d) From the Zeeman effect, one can calculate the e/m ratio for an electron.
- **75.** The wave function for a hydrogen atom with its electron in the 2p state varies with direction as well as distance from the nucleus. What is the probability of a 2p electron, for which $m_l = 0$, existing on the xy plane?
 - (a) 0
- (b) 1
- (c) $2a_0$
- (d) a_0

1 - 13

• *Type 2* •

Choose the correct options. More than one option is correct.

- **76.** Which of the following is true for Thomson's model of the atom?
 - (a) The radius of an electron can be calculated using this model.
 - (b) In an undisturbed atom, the electrons will be at their equilibrium positions, where the attraction between the cloud of positive charge and the electrons balances their mutual repulsion.
 - (c) When the electrons are disturbed by collision, they will vibrate around their equilibrium positions and emit electromagnetic radiation. The frequency of this radiation is of the order of magnitude of the frequency of electromagnetic radiation, typical of these electrons.
 - (d) It can explain the existence of protons.
- 77. From the α -particle scattering experiment, Rutherford concluded that
 - (a) α -particles can come within a distance of the order of $10^{-14}\,\mathrm{m}$ of the nucleus
 - (b) the radius of the nucleus is less than 10^{-14} m
 - (c) scattering follows Coulomb's law
 - (d) the positively charged parts of the atom move with extremely high velocities
- 78. Rutherford's scattering formula fails for very small scattering angles because
 - (a) the full nuclear charge of the target atom is partially screened by its electron

(b) Frequency of revolution $\propto \frac{1}{n^3}$

transition inside a hydrogen atom can be

(a) $\frac{h}{4\pi}$ (b) $\frac{h}{\pi}$ (c) $\frac{h}{2\pi}$

(c) Radius of orbit $\propto n^2 Z$

(d) Force on electron $\propto \frac{1}{n^4}$

	(b)	the impact parameter between the $\alpha\text{-particle}$ source and the nucleus of the target is very large compared to the size of the nucleus								
	(c)	the kinetic energy of the α -particles is large								
	(d)	the gold foil is very thin								
79.		n of the following transitions are allowed in the normal electronic ion spectrum of an atom?								
	(a)	$2s \rightarrow 1s$ (b) $2p \rightarrow 1s$ (c) $3d \rightarrow 2p$ (d) $5p \rightarrow 3s$								
80.	Choos	se the correct statements.								
	(a)	Every object emits radiation whose predominant frequency depends on its temperature.								
	(b)	The quantum energy of a wave is proportional to its frequency.								
	(c)	Photons are quanta of light.								
	(d)	The value of the Planck constant depends on energy.								
81.	The m	nathematical expression for the uncertainty principle is								
	(a)	$\Delta x \Delta p \ge \frac{h}{4\pi}$ (b) $\Delta E \Delta t \ge \frac{h}{4\pi}$								
	(c)	$\Delta x \Delta p \ge \frac{h}{p}$ (d) $\Delta E \Delta t \ge \frac{h}{p}$								
82.	To wh	nich of the following is Bohr's theory applicable?								
	(a)	He^+ (b) Li^{2+}								
	(c)	Tritium (d) Be ³⁺								
83.	Bohr's	s theory is not applicable to								
	(a)	He (b) Li ²⁺								
	(c)	He^{2+} (d) the H atom								
84.	Choos	se the correct relations on the basis of Bohr's theory.								
	(a)	Velocity of electron $\propto \frac{1}{n}$								

85. The change in orbital angular momentum corresponding to an electron

(d) $\frac{h}{8\pi}$

- 86. The magnitude of the spin angular momentum of an electron is given by
 - (a) $S = \sqrt{s(s+1)} \frac{h}{2\pi}$
- (b) $S = s \frac{h}{2\pi}$

(c) $S = \frac{\sqrt{3}}{2} \times \frac{h}{2\pi}$

- (d) $S = \pm \frac{1}{2} \times \frac{h}{2\pi}$
- 87. Choose the correct configurations from among the following.
 - (a) $Cr(Z = 24) : [Ar]3d^5 4s^1$
- (b) $Cu(Z = 29) : [Ar]3d^{10} 4s^{1}$
 - (c) Pd(Z = 46): [Kr]4d 10 4s 0
- (d) Pt(Z = 78): [Xe]4d ¹⁰ 4s ²
- 88. The configuration [Ar]3d ¹⁰4s ²4p ² is similar to that of
 - (a) boron

(b) oxygen

(c) sulphur

- (d) aluminium
- 89. In which of these options do both constituents of the pair have the same magnetic moment?
 - (a) Zn^{2+} and Cu^{+}

- (b) Co^{2+} and Ni^{2+}
- (c) Mn^{4+} and Co^{2+}
- (d) Mg^{2+} and Sc^{+}
- 90. Which of the following ions are diamagnetic?
 - (a) He_2^+
- (b) Sc³⁺
- (c) Mg^{2+} (d) O_2^{2-}
- 91. The ground-state electronic configuration of the nitrogen atom can be represented as
 - (a)



- (c)

- (d)
- 92. Choose the correct statements from among the following.
 - (a) A node is a point in space where the wave function (Ψ) has zero amplitude.

(b)

- (b) The number of peaks in radial distribution is n l.
- (c) Radial probability density $\rho_{n,l}(r) = 4\pi r^2 R_{n,l}^2(r)$.
- (d) Ψ^2 represents the atomic orbital.

Answers

- 1. c
- **2.** d
- 3. b
- 4. b
- **5.** c

- 6. b
- 7. d

- **11.** c
- **12.** b
- 8. c
- 9. c
- **10.** d

- 17. b
- **13.** a
- **14.** a
- **15.** d

- 16. a
- 18. a
- **19.** c
- **20.** a

21. c	22. b	23. b	24. a	25. a
26. a	27. c	28. b	29. c	30. b
31. d	32. d	33. c	34. b	35. c
36. c	37. b	38. d	39. d	40. a
41. c	42. b	43. b	44. d	45. d
46. c	47. c	48. c	49. b	50. d
51. b	52. a	53. c	54. c	55. a
56. b	57. c	58. d	59. d	60. c
61. a	62. b	63. c	64. d	65. d
66. c	67. a	68. d	69. c	70. b
71. a	72. c	73. b	74. b	75. a
76. b, c	77. a, b, c	78. a, b	79. b, c, d	80. a, b, c
81. a, b	82. a, b, c, d	83. a, c	84. a, b, d	85. b, c
86. a, c	87. a, b, c	88. b, c	89. a, c	90. b, c, d
91. a, d	92. a, b, c, d			

Hints to More Difficult Problems

2.
$$n_{C_2} = 4_{C_2} = \frac{4!}{2!2!} = 6.$$

7.
$$r = 1.2 \times 10^{-15} A^{\frac{1}{3}} \text{m}.$$

For Al, $A = 270.$
 $\therefore r = 1.2 \times 10^{-15} (27)^{\frac{1}{3}} \text{m} = 3.6 \times 10^{-15} \text{ m}.$

8.
$$\rho = \frac{Au}{\frac{4}{3}\pi \left[1.2 \times 10^{-15} A^{1/3}\right]^3} = \frac{u}{(1.33\pi (1.2 \times 10^{-15})^3)}$$
$$= \frac{1.66 \times 10^{-27} \text{ kg}}{1.33\pi (1.2 \times 10^{-15})^3 \text{m}^3} = 2.4 \times 10^{17} \text{ kg m}^{-3}.$$

- **9.** Using Moseley's equation $v = a(Z b)^2$, $\sqrt{v} = \sqrt{a(Z b)} = \sqrt{a} \ Z \sqrt{ab}$ which is in the form of y = mx + c. Therefore, a plot of \sqrt{v} against Z is a straight line.
- 13. The reduced mass is given by

$$\mu = \frac{m_e m_e^+}{m_e + m_e^+} = \frac{m_e^2}{2m_e} \text{ (mass of electron = mass of positron)}$$

$$\mu = \frac{1}{2} m_e$$

 \therefore Rydberg constant = $R_{\infty}/2$

16.
$$E = \frac{hc}{\lambda} = \frac{1242 \text{ eV.nm}}{\lambda} = \frac{1242 \text{ eV.nm}}{700 \text{ nm}} = 1.77 \text{ eV.}$$

17.
$$E = hv = (6.626 \times 10^{-34} \text{ Js}) (880 \text{ s}^{-1})$$

= 5.831 × 10⁻³¹ J.

No. of photons emitted = $\frac{1}{5.831} \times 10^{+31} = 1.71 \times 10^{30}$ per second.

18. For a proton
$$\binom{1}{1}H$$
, $\frac{e}{m} = \frac{1}{1}$.

For an
$$\alpha$$
-particle (${}_{2}^{4}$ He), $\frac{e}{m} = \frac{2}{4} = \frac{1}{2}$.

$$\frac{\binom{e}{m}_p}{\binom{e}{m}_q} = \frac{1/1}{1/2} = 2:1.$$

21. The de Broglie equation is
$$\lambda = \frac{h}{p} = \frac{h}{mv}$$
.

Here *h* and *v* are constant. So $\lambda \propto \frac{1}{m}$.

Since the α -particle has the highest mass among the given entities, it has the smallest de Broglie wavelengths.

22. Momentum =
$$p = \frac{hv}{c}$$
.

$$\therefore mv = \frac{hv}{c} \Longrightarrow v = \frac{hv}{mc}.$$

23.
$$E = eV = \frac{1}{2} mv^2 = \frac{p^2}{2m}$$

where V = applied voltage.

We know that $\lambda = \frac{h}{p}$.

$$2meV = p^2 \Rightarrow p = \sqrt{2meV}$$
.

$$\therefore \quad \lambda = \frac{h}{\sqrt{2meV}} = \frac{12.3}{\sqrt{V}} \text{ Å.}$$

26. We know that $v \propto \frac{1}{n}$ and n = principal quantum number. Therefore, vn will be independent of the principal quantum number.

27.
$$-PE = 2KE = 2 \times (+13.6) = 27.2 \text{ eV}.$$

$$\therefore$$
 PE = -27.2 eV

PE with respect to ground state = 27.2 eV which we assume to be zero.

$$KE = -\frac{13.6}{n^2} \text{ eV}.$$

For the first excited state, n = 2.

$$\therefore KE = -\frac{13.6}{2^2} \text{ eV} = -3.4 \text{ eV}$$

Total energy = (27.2 - 3.4) eV = 23.8 eV.

- **31.** $E_n = -\frac{13.6}{n^2} Z^2$, where n = 1 for ground state, 2 for first excited state, and so on. Z = 1 for hydrogen.
- **34.** $\frac{1}{\lambda} = R_{\infty} \left(\frac{1}{n_1^2} \frac{1}{n_2^2} \right)$ for the H atom.

For the Balmer series.

 $n_1 = 2$ and $n_2 = 5$ (for the third line).

$$\frac{1}{\lambda} = R_{\infty} \left(\frac{1}{2^2} - \frac{1}{5^2} \right) = \frac{21}{100} R_{\infty}.$$

$$\therefore \quad \lambda = \frac{100}{21} R_{\infty}.$$

- 37. $N = \text{number of lines emitted} = \frac{1}{2} n(n-1)$ = $\frac{1}{2} \times 4(4-1) = 6$.
- **42.** Here n = 3, which corresponds to 3s, 3p, 3d orbitals which have a degeneracy = 1 + 3 + 5 = 9.
- **45.** l = 3 corresponds to s, p, d and f orbitals. Maximum multiplicity for the f orbitals (7 electrons)

$$=2s+1=2\times\frac{7}{2}+1=8.$$

Minimum multiplicity for the f orbital (1 electron)

$$=2s+1=2\times\frac{1}{2}+1=2.$$

- **49.** n = l is not permissible.
- **50.** The state corresponds to the 3d atomic orbitals.
- 55. The 19th electron corresponds to the 4s orbital. For this orbital,

$$n = 4, l = 0, m = 0, s = \frac{1}{2}$$
.

56. The 21st electron corresponds to 3d ¹. For this orbital,

$$n = 3, l = 2, m = 2, s = +\frac{1}{2}$$
.

- 57. $Fe(Z = 26) = [Ar] 3d^{6}4s^{2}$ $Fe^{3+}(23 \text{ electrons}) = [Ar] 3d^{5}$
- 58. Mn^{4+} (21 electrons) = [Ar] $3d^3$ contains three unpaired electrons.
- 61. Follow the Aufbau principle.

62. The spin-only formula is given by

$$\mu_{\text{spin only}} = \sqrt{s(s+1)} \text{ BM}$$

The value of μ = 2.83 BM corresponds to the presence of two unpaired electrons. So the ion is Ni ²⁺(3d ⁸).

- 66. Use the energy-level diagram.
- 67. For the 3p orbital, n = 3, l = 1. The number of spherical nodes = n - l - 1 = 3 - 1 - 1 = 1.
- **69.** Spin multiplicity = $(2\Sigma s + 1)$.
- 82. All the species have one electron each. So Bohr's theory is applicable.
- **83.** Because He and He²⁺ have two electrons and no electron respectively, Bohr's theory is not applicable to them.
- **88.** Boron $(1s^22s^22p^1)$ as well as aluminium $(1s^22s^22p^63s^23p^1)$ have an s^2p^1 configurations, and so does [Ar] $(3d^{10} 4s^2 4p^1)$.
- **89.** Zn^{2+} and Cu^{+} have a 3d $\operatorname{^{10}}$ configuration, n=0. Mn $\operatorname{^{4+}}$ and Co^{+} have a 3d $\operatorname{^{3}}$ configuration, n=3.
- **90.** The number of unpaired electrons is zero. So $\mu = 0$ BM.

Ц

Nuclear Chemistry

• Type 1 •

Choose the correct option. Only one option is correct.

1.	In radioa	ctive decay	the emitted	d electrons	come from	m the

- (a) innermost shell of the atom
- (b) K shell of an atom
- (c) outermost shell of the atom
- (d) decay of neutrons in the nucleus

2. In the radioactive decay $^{232}_{92}X \longrightarrow ^{220}_{89}Y$, how many α- and β-particles are ejected from X and Y?

(a)
$$5 \alpha$$
 and 5β

(b)
$$3 \alpha$$
 and 3β

(c)
$$3 \alpha$$
 and 5β

(d)
$$5 \alpha$$
 and 6β

3. In the radioactive decay
$${}^{A}_{Z}X \longrightarrow {}^{A}_{Z+1}Y \longrightarrow {}^{A-4}_{Z-1}Z \longrightarrow {}^{A-4}_{Z-1}Z^*$$

the sequence of the radiation emitted is

(a)
$$\alpha$$
, β , γ (b) β , α , γ

(b)
$$\beta$$
, α , γ

(d)
$$\beta$$
, γ , α

4. In nature, which of the following nuclear reactions lead to the formation of tritium?

(a)
$${}_{3}^{6}\text{Li} + {}_{0}^{1}\text{n} \rightarrow {}_{2}^{4}\text{He} + {}_{1}^{3}\text{H}$$
 (b) ${}_{5}^{10}\text{B} + {}_{1}^{1}\text{p} \rightarrow {}_{5}^{8}\text{B} + {}_{1}^{3}\text{H}$

(b)
$${}_{5}^{10}B + {}_{1}^{1}p \rightarrow {}_{5}^{8}B + {}_{1}^{3}H$$

(c)
$${}_{5}^{11}B + {}_{1}^{2}D \rightarrow {}_{5}^{10}B + {}_{1}^{3}H$$

(c)
$${}^{11}_{5}B + {}^{2}_{1}D \rightarrow {}^{10}_{5}B + {}^{3}_{1}H$$
 (d) ${}^{9}_{4}Be + \gamma \rightarrow {}^{6}_{2}He + {}^{3}_{1}H$

5. Which of the following nuclear reactions in nature lead to the formation of radiocarbon?

(a)
$${}^{16}_{8}\text{O} + {}^{14}_{7}\text{N} \to {}^{14}_{6}\text{C} + {}^{16}_{9}\text{F}$$
 (b) ${}^{14}_{7}\text{N} + {}^{1}_{0}\text{n} \to {}^{14}_{6}\text{C} + {}^{1}_{1}\text{H}$ (c) ${}^{14}_{7}\text{N} + {}^{1}_{0}\text{n} \to {}^{14}_{6}\text{C} + {}^{2}_{1}\text{H}$ (d) ${}^{14}_{7}\text{N} + {}^{3}_{1}\text{H} \to {}^{14}_{6}\text{C} + {}^{2}_{2}\text{He}$

(b)
$${}^{14}_{7}N + {}^{1}_{0}n \rightarrow {}^{14}_{6}C + {}^{1}_{1}H$$

(c)
$${}^{14}_{7}N + {}^{1}_{0}n \rightarrow {}^{14}_{6}C + {}^{2}_{1}H$$

(d)
$${}_{7}^{14}N + {}_{1}^{3}H \rightarrow {}_{6}^{14}C + {}_{2}^{3}H\epsilon$$

6. Which of the following nuclei are β +-emitters? (a) Antinutrino (b) Potassium-40 (c) Radon-222 (d) Chlorine-34 7. Which of the following radioactive elements are soluble in water? (a) Radium (b) Radon (c) Technetium (d) Tritium 8. During the transformation of ${}_{c}^{a}$ X to ${}_{d}^{b}$ Y, the number of β-particles emitted (a) $\frac{a-b}{4}$ (b) $d + \frac{a-b}{2} + c$ (c) $d + \left[\frac{a-b}{2}\right] - c$ (d) 2c - d + a - b9. The nucleus of an electrically neutral atom undergoes radioactive decay. It will remain neutral after the decay if the process is a (a) β^- -decay (b) β⁺-decay (d) neutron decay (c) γ-decay 10. Consider the following decay. $_{7}^{A}X \rightarrow _{7+1}^{A}Y + _{1}^{0}e + \overline{\nu}$ X is unstable because (a) of its large nuclide (b) its nucleus has excess energy (c) the nuclide has more neutrons than protons (d) the nuclide has more protons than neutrons 11. A radioactive nuclide emits γ -rays due to the (a) emission of an electron from its orbital (b) nuclear transition from a higher state to a lower state (c) presence of more neutrons than protons (d) presence of less neutrons than protons 12. Which of the following combinations will give the most stable nuclei? (a) Odd Z and odd N(b) Even Z and even N(c) Odd Z and even N (d) Even Z and odd N13. $_{13}^{27}$ Al is a stable isotope. It is expected to disintegrate by (a) α-emission (b) β -emission (c) β⁺-emission (d) proton emission

(a) α -particles

(a) N > 126, Z > 83, A < 208

(c) Z > 82, N < 126, A > 108

(c) γ-rays

15. Which of the following combinations would make the nucleus unstable?

(b) β -particles

(b) N > 126, Z > 83, A > 209

(d) Z = 83, N = 126, A > 209

(d) Positrons

14. In radioactive decay, which of the following move the fastest?

	. ,		. ,	
16.	For st	able nuclei, the average bindin	g en	ergy per nucleon lies between
	(a)	7 and 9 MeV	(b)	10 and 12 MeV
	(c)	2 and 4 MeV	(d)	5 and 7 MeV
17.	The m	nost stable nuclei found in the l		0 0.
	` '	Cu	(b)	Fe
	(c)	С	(d)	Pb
18.	Which	n of the following radioactive s	eries	is artificial?
	(a)	Thorium series (4n)	(b)	Neptunium series $(4n + 1)$
	(c)	Uranium series $(4n + 2)$	(d)	Actinium series $(4n + 3)$
19.	²¹⁹ ₈₄ Rn series	is a member of the actinium s	serie	s. Another member of the same
	(a)	²³⁵ ₉₁ Pa	(b)	²³² ₉₀ Th
	(c)	$^{235}_{92}$ U	(d)	$\frac{227}{88}$ Ra
20.	Amor	ng the following, which has the	long	gest half-life?
	(a)	²³² ₉₀ Th	(b)	²³⁷ ₉₃ Np
	(c)	$^{238}_{92}$ U	(d)	$_{92}^{235}$ U
21.		nalysis of a rock shows that the is $Pb/U = 0.25$. If $t_{1/2}$ for the ise of the rock (in years) is	ne re react	lative number of ^{206}Pb and ^{238}U ion $^{238}\text{U} \rightarrow ^{206}\text{U}$ is 4×10^9 years,
	(a)	$\frac{2.303}{0.693} (4 \times 10^{9}) \log \frac{5}{4}$	(b)	$\frac{2.303}{0.693} (4 \times 10^{9}) \log \frac{1}{4}$
	(c)	$\frac{2.303}{0.693} (4 \times 10^{9}) \log 4$	(d)	$\frac{2.303}{0.693} (4 \times 10^9) \log \frac{4}{5}$
22.	A cert ${}_{Z}^{A}X$ is l	tain radioactive isotope ${}_Z^AX(t_{\frac{1}{2}} = x_{\frac{1}{2}})$ kept in a sealed vessel, how mu	= 10 o ach i	days) decays to $_{Z-2}^{A-4}$ Y. If 1 mol of on will accumulate in 20 days?
	(a)	22.4 L		11.2 L
	(c)	16.8 L	(d)	33.6 L
23.		ioelement decays by two paral are λ_1 and λ_2 . The effective de		eactions, the decay constants for constant (λ) of the nuclide is

24. The time of decay for a nuclear reaction is given by $t = 4t_{1/2}$. The relation between the mean life (*T*) and time of decay (*t*) is given by

(a) $\lambda = \lambda_1 / \lambda_2$

(c) $\lambda = \lambda_1 + \lambda_2$

(a) $2T \ln 2$

(c) $2T^4 \ln 2$

25.

26.

27.

28.

29.

30.

(b) $\lambda = \lambda_1 - \lambda_2$

(d) $\lambda = \frac{1}{\lambda_1} + \frac{1}{\lambda_2}$

(b) 4T ln 2 (d) $\frac{1}{T^2} \ln 2$

A freshly prepared radioelement has a half-life of 2 hours. It emits radiation whose intensity is 64 times the permissible safe level. The minimum time after which it would be possible to work with this sample is								
(a)	3 hours	(b)	9 hours					
(c)	24 hours	(d)	12 hours					
The activity of a sample of a radioactive nuclide (100 X) is 6.02 curies. Its disintegration constant is 3.7×10^4 s $^{-1}$. The initial mass of the sample is								
(a)	$1 \times 10^{-14} \text{ g}$	(b)	$1 \times 10^{-6} \text{ g}$					
(c)	$1 \times 10^{-15} \text{ g}$	(d)	$1 \times 10^{-3} \mathrm{g}$					
	alf-life of a radioactive sample will remain undecayed after		2 <i>n</i> years. What fraction of this ars?					
(a)	$\frac{1}{2}$	(b)	$\frac{1}{\sqrt{2}}$					
(c)	$\frac{1}{\sqrt{3}}$	(d)	2					
One g	ram of ²²⁶ Ra has an activity of	nearl	ly 1 Ci. The t_{15} of 226 Ra is					
_	1620 years		12.5 years					
(c)	140 days	(d)	4.5×10^9 years					
	tple of a radioisotope (t_{1_2} = 3 dayse was left. What was the initial r		as taken. After 12 days, 3 g of the of the sample?					
(a)	112 g (b) 136 g	(c)	12 g (d) 48 g					
	tte of decay of a radioactive satter time t_2 . The mean life of thi		is given by R_1 at time t_1 , and R_2 ioactive sample is					
(a)	$T = \frac{R_1}{R_2} \times \frac{t_2}{t_1}$	(b)	$T = (t_1 - t_2) / \ln (R_2 / R_1)$					
(c)	$T = (t_2 - t_1) / \ln (R_2 / R_1)$	(d)	$T = \frac{\ln{(R_2/R_1)}}{t_1 - t_2}$					

liquid is
(a) 160

31. The count rate of 200 mL of a radioactive liquid sample is x. Some of the liquid is now discarded. The count rate of the remaining liquid is found to be x/20 after four half-lives. The volume in millilitres of the remaining

(c) 40

(d) 10

(b) 80

32. Which of the following is a β^+ emitter?

	(a)	⁴⁹ ₂₀ Ca	(b)	8 5 B	(c)	²⁰⁸ ₈₂ Pb	(d)	$^{94}_{36}{ m Kr}$	
33.	energ	alf-life of a y of the β-pa mple emit er	rticle	emitt	ample of ²¹⁰ ted is 0.34 N	RaE is 5 IeV. At	days an what rate	nd the a e in wat	verage ts does
	(a)	2.0			(b)	0.1			
	(c)	1.5			(d)	1.0			
34.	undec	of the radioa cayed after o ays will be							
	(a)	64			(b)	20			
	(c)	46			(d)	80			
35.	35. Assuming that 226 Ra ($t_{1/2} = 1.6 \times 10^3$ yr) is in secular equilibrium with 238 U ($t_{1/2} = 4.5 \times 10^9$ yr) in a certain mineral, how many grams of Ra will be present for every gram of 238 U in this mineral?							n with Ra will	
	(a)	3.7×10^{-7}			(b)	3.4×10	7		
	(c)	3.4×10^{-7}			(d)	3.7×10	7		
36.		n of the follo		g proc	cesses resul	ts in an	increase	in the	atomic
	(a)	Alpha emis	sion		(b)	Electron	n capture		
	(c)	Beta emissio	on		(d)	Positro	n emissio	n	
37.	Which	n of the follow	wing	proces	sses causes	the emis	sion of ar	n X-ray?	
	(a)	Alpha emis	sion		(b)	Gamma	a emissio	n	
	(c)	Positron em	issio	n	(d)	Electron	n capture		
38.	With	increasing nu	ıclea	r size,					
			e for	ce bet	ween the pr	otons in	icreases a	nd so d	oes the
	(b)	the repulsiv				otons de	ecreases a	and the	energy
	(c)	the attractive energy of the			ween the pi	otons in	icreases a	nd so d	oes the
	(d)	the attractive energy of the			ween the pr	otons de	ecreases a	nd so d	oes the

- **39.** Temperature has no effect on rate of decay because
 - (a) nuclear energies are very high
 - (b) nuclear energies are low
 - (c) radioactive decay is a statistical process
 - (d) radioactive decay follows exponential law
- 40. Nuclear isomers are nuclei with the same
 - (a) number of electrons, protons and electrons, and have the same half-life
 - (b) number of protons and neutrons but their energy levels are different
 - (c) number of protons and neutrons, and the same energy levels
 - (d) value of half-life and average life
- 41. Sometimes the ejection of an α -particle does not completely stabilize the nucleus. In such a case more α -particles may be emitted. α -decay
 - (a) raises the N/P ratio and is often followed by β -emission
 - (b) lowers the N/P ratio and is often followed by positron emission
 - (c) raises the N/P ratio and is often followed by neutron emission
 - (d) lowers the N/P ratio and is often followed by γ -ray emission
- **42.** The age of a specimen, t, is related to the daughter/parent ratio D/P by the equation

(a)
$$t = \frac{1}{\lambda} \ln \frac{D}{P}$$

(b)
$$t = \frac{1}{\lambda} \ln \left(1 + \frac{P}{D} \right)$$

(c)
$$t = \frac{1}{\lambda} \ln \left(1 + \frac{D}{P} \right)$$

(d)
$$t = \frac{1}{\lambda} \ln \left(2 + \frac{P}{D} \right)$$

- **43.** A $^{235}_{92}$ U nucleus absorbs a neutron and forms $^{236}_{92}$ U. This new nucleus is
 - (a) very stable and does not undergo any further nuclear reaction
 - (b) unstable and almost at once explodes into two fragments
 - (c) unstable and forms ²³⁹₉₄Pu
 - (d) unstable and the fragment product undergoes nuclear fusion
- 44. The mechanism of nuclear fission is explained on the basis of
 - (a) magic number of the nucleus (b) binding energy
 - (c) liquid-drop model
- (d) viscosity of the nucleus
- **45.** As the mass number A increases, the binding energy per nucleon of the nucleus
 - (a) decreases
 - (b) varies in a way that depends upon the actual value of A

- (c) remains unchanged
- (d) increases
- 46. During a nuclear fission reaction,
 - (a) a heavy nucleus bombarded by thermal neutrons, splits
 - (b) two light nuclei combine to give a heavier nucleus
 - (c) a heavy nucleus splits into two fragments by itself
 - (d) a light nucleus bombarded by thermal neutrons, splits
- **47.** Two deuterium nuclei in a deuterium vapour state at room temperature do not combine to form a helium nucleus because
 - (a) He ²⁺ nucleus is unstable
 - (b) coulombic repulsion does not allow the nuclei to come close together
 - (c) a deuterium nucleus is more tightly bound than a helium nucleus
 - (d) it is not energetically favourable
- **48.** A heavy nucleus has larger N/Z ratio because
 - (a) a neutron has large mass compared to a proton and an electron
 - (b) mean life of neutron is greater than that of a proton
 - (c) a neutron exerts electric repulsion
 - (d) the nucleus has larger number of neutrons
- **49.** During nuclear fission, the final state has lesser energy than the reactant. The nuclear reaction takes place even though intermediate state has greater energy than the initial one and no energy is supplied externally. This process is called
 - (a) barrier penetration
 - (b) tunnelling
 - (c) formation of compound nucleus
 - (d) binding energy
- **50.** The fission fragments reduce their N/Z ratio further via
 - (a) positive beta decay
- (b) negative beta decay

(c) alpha decay

- (d) gamma decay
- 51. As fission fragments decay
 - (a) it gives rise to stable heavy nucleus
 - (b) it gives light nuclei with an emission of energy
 - (c) an additional energy in the form of kinetic energy of β-particles, antinutrino and photons are released
 - (d) an additional energy is absorbed and then it decays to give β and $\gamma\text{-rays}$

52.		n of the following nuclei will 1–100 eV?	abso	orb fast neutron having energy				
	Given, fast neutron (energy ≈ 2 MeV) and slow neutron (energy ≈ 0.04 eV).							
	(a)	^{235}U	(b)	^{238}U				
	(c)	^{233}U	(d)	$_{1}^{2}H$				
53.				s of neutron is increased fast, the using a metal moderator which				
	(a)	Be	(b)	Zn				
	(c)	Cd	(d)	carbon rod				
54.		n and other stars, where temp dominantly by	erat	ure is about 10 ⁷ K, fusion takes				
	(a)	proton-nitrogen cycle	(b)	proton-proton cycle				
	(c)	proton-deuterium cycle	(d)	proton-lithium cycle				
55.	When	the temperature inside the st ss can continue to produce stab	ar in ole	creases, it produces $^{12}C + \gamma$. The				
	(a)	Fe(A = 56)	(b)	N(A = 14)				
	(c)	Zn(A=65)	(d)	U(A = 238)				
56.	(over			ther may come close enough temperature of about 10 ⁸ K) to				
	(a)	thermonuclear fission	(b)	nuclear fusion				
	(c)	breeder reaction	(d)	positron emission				
57.		tter stars, where the temperature cycle is known as	ıre is	about 10 ⁸ K, fusion takes place				
	(a)	proton-carbon cycle	(b)	proton-neutron cycle				
	(c)	carbon-deuterium cycle	(d)	nitrogen-oxygen cycle				
58.	On w	_	ne pi	rinciple of radiocarbon dating is				

- (a) The radioactive ^{14}C content of a sample of dead animal or plant tissue decreases steadily, while its ^{12}C content remains unchanged. Hence the ratio of ^{14}C to ^{12}C of the sample indicates that elapsed since the death of the organism.
- (b) The rate of disintegration of ¹⁴C is faster than ¹²C.
- (c) The ratio of ¹⁴C to ¹²C is not a fixed quantity.
- (d) ¹⁴C is radioactive and ¹²C is non-radioactive.

• *Type* 2 •

Choose the correct options. More than one option is correct.

- **59.** Which of the following is used as a moderator in a nuclear reactor?
 - (a) Heavy water

(b) Graphite

(c) Beryllium

- (d) Sodium
- **60.** Which of the following statements is correct?
 - (a) ²³⁸U has only a small cross section for the capture of slow neutron
 - (b) ²³⁵U has a very large cross section for the capture of slow neutron
 - (c) Neutrons are accelerated by the moderators
 - (c) All of these
- **61.** When fission occurs, several neutrons are released and the fission fragments are beta radioactive because
 - (a) the neutron/proton ratio required for stability, decreases with increasing A
 - (b) some of the excess neutrons are released directly
 - (c) some of the neutrons change to proton by beta decay in the fission fragments
 - (d) all of these
- **62.** Which of the following statments is correct?
 - (a) Radiocarbon is produced by the cosmic rays following the nuclear reaction $^{14}N(n, p)$
 - (b) All living things do not contain radiocarbon
 - (c) The proportion of radiocarbon in living matter is constant but decreases after death, which permits the remains to be dated
 - (d) All of these
- 63. In the decay process

$$A \xrightarrow{-\alpha} B \xrightarrow{-\beta} C \xrightarrow{-\beta} D$$

- (a) A and B are isobars
- (b) A and D are isotopes
- (c) B, C and D are isobars
- (d) A and C are isotones
- **64.** Which of the following are β -emitters?
 - (a) Carbon-14

(b) Cobalt-60

(c) Tritium-3

- (d) Free neutron
- **65.** Which of the following are α -emitters?
 - (a) Polonium-212

(b) Radium-226

(c) Helium-5

(d) Tritium

- 66. Which of the following do not occur?
 - (a) ${}^{40}_{20}$ Ca + ${}^{1}_{0}$ N $\rightarrow {}^{40}_{10}$ K + ${}^{1}_{1}$ H
- (b) ${}_{12}^{24}\text{Mg} + {}_{2}^{4}\text{He} \rightarrow {}_{14}^{27}\text{Si} + {}_{0}^{1}\text{n}$
- (c) ${}^{113}_{49}\text{Cd} + {}^{1}_{0}\text{N} \rightarrow {}^{112}_{49}\text{Cd} + {}^{0}_{1}\text{e}$ (d) ${}^{43}_{20}\text{Ca} + {}^{4}_{2}\text{He} \rightarrow {}^{46}_{21}\text{Sc} + {}^{1}_{1}\text{H}$
- 67. Which of the following make up an isotonic triad?
 - (a) ${}^{14}_{6}C$, ${}^{16}_{8}O$, ${}^{15}_{7}N$

- (b) ${}^{76}_{32}$ Ge, ${}^{77}_{33}$ As, ${}^{75}_{31}$ Ga
- (c) ⁴⁰₁₀Ar, ⁴⁰₁₀K, ⁴⁰₂₀Ca
- (d) $^{233}_{92}$ U, $^{232}_{90}$ Th, $^{239}_{94}$ Pu
- 68. Stable nuclides cannot be obtained for
 - (a) Z = 43, N = 35

(b) Z = 61, N = 89

(c) A = 8 or 5

- (d) A > 209
- **69.** Which of the following processes are feasible?
 - (a) ${}_{0}^{1}n \rightarrow {}_{1}^{1}p + {}_{0}^{0}e + \overline{\nu}$
- (b) ${}_{1}^{1}p \rightarrow {}_{0}^{1}n + {}_{1}^{0}e + v$

(c) ${}_{1}^{1}H + {}_{1}^{0}e \rightarrow {}_{0}^{1}n$

- (d) ${}_{1}^{1}p + {}_{0}^{1}n \rightarrow {}_{1}^{2}D + v$
- 70. Which of the following nuclei are stable?
 - (a) ${}^{60}_{28}$ Ni

(b) 11₆C

(c) ⁷₂Li

- (d) ⁸₄Be
- 71. A nuclide has mass number A and atomic number Z. During a radioactive process, if
 - (a) both A and Z decrease, the process is called α -decay
 - (b) A remains unchanged and Z decreases by one, the process is called β⁺-decay or K-electron capture
 - (c) both A and Z remain unchanged, the process is called γ -decay
 - (d) both *A* and *Z* increase, the process is called nuclear isomerism
- 72. Which of the following nuclei are doubly magic?
 - (a) ${}_{2}^{4}$ He
- (b) ${}^{16}_{8}$ O
- (c) ${}^{208}_{82}$ Pb (d) ${}^{238}_{92}$ U
- 73. A nuclide X undergoes α -decay and another nuclide Y, β ⁻ decay. Which of the following statements are correct?
 - (a) The β -particles emitted by Y may have widely different speeds.
 - (b) The α -particles emitted by X may have widely different speeds.
 - (c) The α -particles emitted by X will have almost the same speed.
 - (d) The β -particles emitted by Y will have the same speed.
- 74. Which of the following statements is correct?
 - (a) When an electron is emitted by an atom and its nucleus gets de-excited as a result, the process is called internal conversion.

- (b) Pair production is a process which involves the creation of a positron-electron pair by a photon of energy 1.02 MeV.
- (c) Neutrons are emitted in the electron-capture process.
- (d) Electron capture and β^+ -emission are identical processes.

75. Which of the following statements are correct?

- (a) In many α and β -decay processes, γ -radiation is emitted.
- (b) The nuclear isomers produced by γ-ray bombardment have the same atomic and mass number but differ in their life-times (except that the ground states may be stable).
- (c) The wavelength and absolute magnitude of the difference between the spins of the initial and final states are the same for γand X-rays.
- (d) A nucleus in an excited state may give up its excitation energy and return to the ground state by the emission of electromagnetic γ-radiation.

Answers

1. d	2. b	3. b	4. a	5. b
6. b	7. b	8. c	9. c	10. c
11. b	12. b	13. b	14. c	15. b
16. a	17. b	18. b	19. c	20. a
21. a	22. c	23. c	24. b	25. d
26. c	27. b	28. a	29. d	30. b
31. a	32. b	33. d	34. a	35. c
36. c	37. d	38. a	39. a	40. b
41. a	42. c	43. b	44. c	45. b
46. a	47. b	48. d	49. a	50. b
51. c	52. b	53. c	54. b	55. a
56. b	57. a	58. a	59. a, b, c	60. a, b
61. b, c	62. a, c	63. b, c	64. a, b, c, d	65. a, b
66. c, d	67. a, b	68. a, b, c, d	69. a, b	70. a, c
71. a, b, c	72. a, b, c	73. a, c	74. a, b, d	75. a, b, c, d

Hints to More Difficult Problems

2.
$$^{232}_{92}X \rightarrow ^{220}_{89}Y + x^{4}_{2}He^{2+} + y - ^{0}_{-1}e$$
.

Equating the mass numbers on both sides of the equation, we get $232 = 220 + 4x + y \Rightarrow x = 3$.

Equating the atomic numbers on both sides of the equation, we have $92 = 89 + 2x - y = 89 + 2 \times 3 - y = 89 + 6 - y \Rightarrow y = 3$.

Thus, three α and three β -particles are emitted.

- 3. ${}_{Z}^{A}X \rightarrow {}_{Z+1}^{A}Y + {}_{-1}^{0}e(\beta^{-}), \ {}_{Z+1}^{A}Y \rightarrow {}_{Z-1}^{A-4}Z + {}_{2}^{4}He^{2+}(\alpha) \text{ and } {}_{Z-1}^{A-4}Z \rightarrow {}_{Z-1}^{A-4}Z + \gamma$ Therefore, the sequence of radiation is β , α , γ .
- 8. It may be stated that

$$_{c}^{a}X \rightarrow _{d}^{b}Y + x_{2}^{4}He + y_{-1}^{0}e$$

where $x = \text{no. of } \alpha\text{-particles emitted}$

and $y = \text{no. of } \beta\text{-particles emitted.}$

$$a = b + 4x \quad \text{or} \quad x = \frac{a - b}{4} \tag{1}$$

and
$$c = d + 2x - y$$
. (2)

Substituting the value of x from Equation (1) in Equation (2), we get

$$c = d + \left(\frac{a - b}{4}\right)2 - y$$

- $y = \text{no. of } \beta\text{-particles emitted} = d + \left[\frac{a-b}{2}\right] c$
- 9. During γ -radiation, the mass number and atomic number remain unchanged. So the nuclide remains neutral.

13.
$$^{29}_{13}\text{Al}$$
 \longrightarrow $^{29}_{14}\text{Si}$ + $^{0}_{-1}\text{e}$ $N/Z = \frac{16}{13} = 1.23$ $N/Z = \frac{15}{14} = 1.07$ (required for stability)

- **15.** The N/Z ratio in option (b) is not conductive to stability.
- **19.** $^{219}_{84}$ Rn and $^{235}_{92}$ U belong to the 4n + 3 series.

21. Given
$$\frac{Pb}{U} = 0.25 \Rightarrow 1 + \frac{Pb}{U} = 1 + 0.25 \Rightarrow \frac{U + Pb}{U} = 1.25$$
 or $\frac{N_0}{N} = 1.25 = \frac{5}{4}$.
 $t = \frac{2.303}{\lambda} \log \frac{N_0}{N} y$ and $\lambda = \frac{0.693}{t_{1/2}} = \frac{0.693}{4 \times 10^9} y^{-1}$.
 $\therefore t = \frac{2.303}{0.693} (4 \times 10^9) \left(\log \frac{5}{4}\right) y$.

24.
$$t_{1/2} = \frac{\ln 2}{\lambda} = T \ln 2$$
 $\left[T = \frac{1}{\lambda} \right].$ (1)

Given,
$$t = 4t_{1/2} \Rightarrow t_{1/2} = \frac{t}{4}$$
 (2)

From Equations (1) and (2),

$$\frac{t}{4} = T \ln 2 \Rightarrow t = 4T \ln 2.$$

25. We know that
$$t = t_{1/2} \frac{\ln (N_0 / N)}{\ln 2}$$
.
 $t = 2 \frac{\ln 64}{\ln 2} = 2 \frac{\ln 2^6}{\ln 2} = 12 \text{ hrs.}$

26. Activity =
$$\lambda N$$
.

$$6.02 \times 3.7 \times 10^{10} = 3.7 \times 10^{4} \frac{m}{100} \times 6.02 \times 10^{23} \Rightarrow m = 1 \times 10^{15} \text{ g}.$$

27. We know that
$$t = t_{\frac{1}{2}} \frac{\ln (N_0/N)}{\ln 2}$$
.

$$n = 2n \frac{\ln (N_0/N)}{\ln 2} \Rightarrow \ln 2 = \ln \left(\frac{N_0}{N}\right)^2 \Rightarrow \frac{N_0}{N} = \sqrt{2} \Rightarrow \frac{N}{N_0} = \frac{1}{\sqrt{2}}$$

29. Using
$$t = t_{1/2} \frac{\ln (N_0/N)}{\ln 2}$$
, we get

$$12 = 3 \frac{\ln (N_0/N)}{\ln 2} \Rightarrow 4 \ln 2 = \ln (N_0/N)$$

$$\Rightarrow \frac{N_0}{N} = 2^4 \Rightarrow N_0 = 2^4 N = 2^4 \times 3 = 48 \text{ g.}$$

30. Let
$$R_0$$
 = initial activity.

Then $R_1 = R_0 e^{-\lambda t_1}$ and $R_2 = R_0 e^{-\lambda t_2}$ for state 2.

$$\frac{R_2}{R_1} = \frac{e^{-\lambda t_2}}{e^{-\lambda t_1}} = e^{\lambda(t_1 - t_2)} = e^{(t_1 - t_2)/T} \qquad \left[\lambda = \frac{1}{T}\right]$$

or
$$R_2 = R_1 e^{(t_1 - t_2)/T}$$
.

$$T = (t_1 - t_2)/\ln (R_2/R_1).$$

31. Initial count rate for 1 mL of liquid = $\frac{x}{200}$.

After 4 half-lives,

count rate for 1 mL of liquid =
$$\frac{1}{2^4} \times \frac{x}{200}$$
.

Let the volume of the remaining liquid = V mL.

$$\therefore$$
 count rate of this liquid $V \times \frac{x}{2^4 \times 200} = \frac{x}{20}$

or
$$V = 160 \text{ mL}$$

33. Using the equation

$$P = E\left(\frac{dN}{dt}\right),\,$$

where P = power

and E = energy

$$\left|\frac{dN}{dt}\right| = \lambda N,$$

we get

$$P = 0.34 \times 10^{6} \times 1.6 \times 10^{-19} \times \frac{0.693}{5 \times 86400} \times \frac{4 \times 10^{-3}}{210} \times 6.02 \times 10^{23}$$

$$= 1 \text{ W}. \qquad (1 \text{ day} = 86.400 \text{ s})$$

- **34.** Equal fractions decay in equal periods of time. Therefore, if a fraction 0.8 remains undecayed after 1 day, a fraction $(0.8)^2 = 0.64$ will remain undecayed after 2 days.
- **35.** Using the equation for secular equilibrium, $\lambda_1 N_1 = \lambda_2 N_2$, where the subscript 1 corresponds to ²³⁸U and 2 to ²²⁶Ra, we get

$$N_2 = \frac{\lambda_1}{\lambda_2} N_1 \Rightarrow \frac{m_2}{226} \times 6.02 \times 10^{23} = \frac{\frac{0.693}{4.5 \times 10^9}}{\frac{0.693}{1.6 \times 10^3}} \times \frac{m_1}{238} \times 6.02 \times 10^{23}.$$

$$\frac{m_2}{m_1} = \frac{1.6 \times 10^3}{4.5 \times 10^9} \times \frac{226}{238} = 3.4 \times 10^{-7}.$$

- **37.** During electron capture, the proton in the nucleus captures an electron from the K shell, creating a vacancy. This vacancy is filled by the electron in the L shell, with the emission of X-rays.
- **42.** $N = N_0 e^{-\lambda t}$

where N = parent remaining = P

and N_0 = parent + daughter = P + D.

$$\begin{split} P &= (P+D)e^{-\lambda t} \\ \frac{P}{P+D} &= e^{-\lambda t} \Longrightarrow \ln \frac{P+D}{P} = \lambda t \\ \Longrightarrow t &= \frac{1}{\lambda} \ln \left(1 + \frac{D}{P} \right). \end{split}$$

- 43. Forms energetic compound nucleus
- 44. Greater cross-section area
- 49. Based on "Mass defects" and "Binding energy"
- 54. Cadmium is a good neutron absorber

63. Example

$$\overset{232}{_{90}}Th \overset{-\alpha\left(\overset{4}{_{2}}He^{2+}\right)}{\overset{228}{_{88}}Ra} \overset{228}{\overset{-\beta\left(\overset{0}{_{-1}}e\right)}{\overset{228}{_{89}}}Ac} \overset{228}{\overset{-\beta\left(\overset{0}{_{-1}}e\right)}{\overset{228}{_{90}}}Th}$$

The emission of one α - and two β -particles produces isotopes (A and D). The emission of n β -particles produces isobars (B, C and D).

- **65.** Usually when A > 209 and Z > 83, the nucleus is an α -emitter.
- 67. All the nuclei in (a) have 8 neutrons and in (b) 44 neutrons.
- **72.** The magic numbers of nuclides are 2, 8, 20, 28, 50, 82 and 126 (for either *N* or *Z*, or both *N* and *Z*).

 ${}_{2}^{4}$ He(N = 2, Z = 2), ${}_{8}^{16}$ O(N = 8, Z = 8) and ${}_{82}^{208}$ Pb(N = 126, Z = 82) are doubly magic.

Chemical Bonding

• Type 1 •

Choose the correct option. Only one option is correct.

- 1. An ionic bond can be formed between two atoms when
 - (a) one of them has a low ionization energy and the other a high electron affinity
 - (b) both the atoms have low values of ionization energy
 - (c) both the atoms have high values of ionization energy
 - (d) both the atoms have low values of electron affinity
- 2. Which of the following is required for the formation of an ionic bond?
 - (a) An electron from the more electronegative element should be transferred to the less electronegative element.
 - (b) The total energy of the resulting molecule should be less than the total energy of the reactants.
 - (c) The lattice energy of the resultant molecule should be as low as possible.
 - (d) The ionic potentials of the reactants should be identical.
- 3. The cohesive energy of an ionic crystal is the energy
 - (a) liberated during its formation from individual neutral atoms
 - (b) absorbed during its formation from individual neutral atoms
 - (c) liberated during the formation of positive ions
 - (d) absorbed during the formation of negative ions
- **4.** Among the following compounds, which has the maximum number of sp-hybridized C atoms?
 - (a) $(CN)_2$
 - (b) CH=C=CH—CN

- (c) $HC \equiv C CH_2CH_2 = C = C = CH_2$
- (d) HC≡C—CN
- 5. Which of the following statements is incorrect?
 - (a) NH₃ is more basic than PH₃.
 - (b) NH₃ has a higher boiling point than that of HF.
 - (c) N_2 is more inert than P_4 .
 - (d) The dipole moment of NH₃ is less than that of SO₂.
- 6. An ionic solid is a poor conductor of electricity because
 - (a) ions do not conduct electricity
 - (b) the charge on the ions is uniformly distributed
 - (c) ions occupy fixed positions in solids
 - (d) ions have uniform fields of influence
- 7. According to Fajans' rules, ionic bonds are formed when cations have
 - (a) low positive charge and large size
 - (b) low positive charge and small size
 - (c) high positive charge and large size, and anions have a small size
 - (d) a low positive charge and large size, and anions have a small size
- 8. A covalent bond is formed between two atoms when
 - (a) one or more pairs of electrons are shared by the two atoms
 - (b) the two atoms are of the same size
 - (c) the electronegativities of the two atoms differ considerably
 - (d) the electron affinities of the two atoms are the same
- 9. What conclusions can be drawn from the following reactions?

$$H_2 + 4.5 \text{ eV} \longrightarrow H + H$$

 $H + 13.6 \text{ eV} \longrightarrow p^+ + e^-$

- (a) It is more difficult to break up an H₂ molecule than it is to break up a hydrogen atom.
- (b) It is easier to break up an H₂ molecule than it is to break up a hydrogen atom.
- (c) The average energy of formation of H and p + are the same.
- (d) Electron and proton attraction in an H₂ molecule as well as an H atom are the same.
- **10.** In an H_2^+ ion
 - (a) one electron is bound to two protons
 - (b) two electrons are bound to two protons
 - (c) three electrons are bound to two protons
 - (d) none of these happens

11. AlCl₃ is covalent while AlF₃ is ionic. This can be justified on the basis of

(a) the valence-bond theory

	(b)	Fajans' rules	3						
	(c)	the molecula	ar-or	bital theory					
	(d)	(d) hydration energy							
12.		ng of the follo CaCl ₂	-	g molecules, v SnCl ₂		h is the most NaCl		e? CuCl	
13.	Amor	ng the followi	ng, t	which has ma	xim	um hardness	?		
		NaF	_	Al_2O_3				${\rm TiO_2}$	
14.	Which	n of the follow	ving	has the highe	est b	ond energy?			
	(a)	F_2	(b)	Cl_2	(c)	Br_2	(d)	I_2	
15.	Which energ		owir	ng pairs hav	e ne	early identic	al va	alues of bond	
	(a)	O_2 and H_2	(b)	N_2 and CO	(c)	F_2 and I_2	(d)	O ₂ and Cl ₂	
16.	 6. The types of bonds present in CuSO₄·5H₂O are (a) electrovalent and covalent (b) electrovalent and coordinate covalent (c) electrovalent, covalent, coordinate covalent and hydrogen bonds (d) covalent and coordinate covalent 								
17.	Which	n of the follow	ving	is the most ic	onic?				
	(a)	P_4O_{10}			(b)	MnO			
	(c)	CrO_3			(d)	Mn_2O_7			
18.	(a) (b) (c)	ng LiCl, BeCl ₂ LiCl < BeCl ₂ LiCl > BeCl ₂ LiCl < BeCl ₂ LiCl > BeCl ₂	2 > B0 2 < B0 2 < B0	$Cl_3 > CCl_4$ $Cl_3 < CCl_4$ $Cl_3 < CCl_4$	he co	ovalent bond	char	acter varies as	
19.		ng LiCl, BeCl ₂ onic characte			comp	oounds with	the g	reatest and the	
		LiCl and Cs		pectively are	(b)	NaCl and Li	iCl		
	. ,	CsCl and Na				CsCl and Be			
20.	The o	verlapping pols	owei order	rs (overlap in	tegra	als) of 2s, 2p,	2sp ²	2 , 2sp^3 and 2sp	
	(a)	$2s > 2p > 2s_1$	$>^3 > 1$	$2sp^2 > 2sp$	(b)	$2p > 2s > 2s_1$	$2^3 > 1$	$2sp^2 > 2sp$	
	(c)	$2sp^3 > 2sp^2$	> 2sı	0 > 2s > 2p	(d)	$2sp^3 > 2sp^2$	> 2s ₁	p > 2p > 2s	

21. Among the following species, which has the maximum number of

22. According to Fajans' rules, covalent bond formation is favoured when

(b) SO₄²⁻

(d) MnO_4^-

resonating structures?

(a) large cation and a small anion(b) large cation and a large anion

(a) PO_4^{3-}

(c) AsO_3^{3-}

there is a

		small cation and a small anion small cation and a large anion							
23.	(a)	ohorus shows five six	s a maximum ((b)	ey of seven three				
24.	(a)	ng the follow: BF ₃ SF ₆	ing, which hav	(b)	ating structu PCl ₅ IF ₇	ires?			
25.		n of the follow Al ₂ Cl ₆	wing molecule (b) Hg ₂ Cl ₂			centred (d)			
26.	eleme comp	ent X has 6 ound formed	nt A has 3 electrons in it by A and X is (b) A_2X_3	ts outer	most shell.	The fo	rmula of the		
27.	In a m (a) (b) (c)	netallic crysta valence ele own kernels valence elec valence elec	nl the ctrons remain	within te a sea (lized bet	the fields of mobile elewern the two	of influ	ence of their		
28.	Polari adjace staten (a) (b) (c)	zation involently placed nents is corre Maximum charge. Minimum p A large ca polarization	ves the distocation. In ext? polarization is polarization is ation is likel	rtion of this cor is broug brought y to b	the shape atext, which the about by a cring about	n of the sy a cation of a high	tion of high of low radius.		

29.	The b	onds present in N ₂ O ₅ are							
	(a)	only ionic	(b)	covalent and coordinate					
	(c)	only covalent	(d)	covalent and ionic					
30.		ement X forms compounds not form XCl ₅ . X is	s of forr	mula XCl_3 , X_2O_5 and Mg_3X_2 but					
	(a)	aluminium	(b)	phosphorus					
	(c)	nitrogen	(d)	boron					
31.	-	drous AlCl ₃ is covalent but AlCl ₃ has a plane triangul							
	(b)	the ionization energy of A	l is very	low					
	(c)	the hydration energy of A	lCl ₃ is ve	ery high					
	(d)	the hydration energy of A energy of aluminium	lCl ₃ com	npensates for the high ionization					
32.		he molecular sizes of ICl and Br_2 are nearly the same, but the boiling oint of ICl is about 39°C higher than that of Br_2 . This is because							
	(a)	the bond energy of I—Cl i	s greate	r than that of Br—Br					
	(b)	the ionization energy of io	dine is l	ess than that of bromine					
	(c)	ICl is polar while Br ₂ is no	npolar						
	(d)	the size of iodine is greate	r than th	nat of bromine					
33.		h of the following oxyacids) bonds?	of sulp	hur contain no sulphur-sulphur					
	(a)	$H_2S_2O_4$	(b)	$H_2S_2O_5$					
	(c)	$H_2S_2O_7$	(d)	$H_2S_2O_3$					
34.		n of the following oxya	icids of	phosphorus are monoprotic					
	(a)	H_3PO_4	(b)	H_3PO_3					
	(c)	H_3PO_2	(d)	$H_4P_2O_7$					
35.	Which	h of the following has great	er bond	length?					
	(a)	P—O (b) S—O	(c)	Cl—O (d) O=O					
36.	Amor	ng following alcohols, which	h has the	e lowest boiling point?					
	(a)	n-Butyl alcohol	(b)	Isobutyl alcohol					
	(c)	sec. Butyl alcohol	(d)	tert. Butyl alcohol					
37.	Which charac		arrange	d in order of increasing covalent					
	(a)	KCl < CaCl ₂ < AlCl ₃ < SnC	Cl ₄ (b)	$SnCl_4 < AlCl_3 < CaCl_2 < KCl$					

 $\text{(c)} \quad AlCl_3 < CaCl_2 < KCl < SnCl_4 \quad \text{(d)} \quad CaCl_2 < SnCl_4 < KCl < AlCl_3 \\$

38. Which of the following factors is the most responsible for increase in

39. In which of the following structures is the energy requirement maximum

(b) F—H----O

(d) O-H----N

boiling point as we move from He to Xe?(a) Decrease in ionization energy(b) Increase in electronegativity(c) Decrease in polarizability(d) Increase in polarizability

for the formation of a hydrogen bond?

(a) F—H----F (c) O—H----O

40.	Hydrogen bonding is exhibited by									
	(a)	all substance	es containing H	atoms						
	(b)	molecules in	n which hydroge	en is bo	onded to F,	O or N				
	(c)	molecules in bonded to C		drogen	is bonded	to F and the other is				
	(d)	all substance	es containing H	and O	atoms					
41.			es are pressed ving forces help			n to form one cube. ner?				
	(a)	Hydrogen bond formation								
	(b)	v) Van der Waals forces								
	(c)	Covalent att	raction							
	(d)	Dipole inter	action							
42.		naximum pos rule can partic		of hyd	rogen bond	ls in which a water				
	(a)	four	(b) three	(c)	two	(d) one				
43.	The d	ensity of wate	er is greater thai	n that o	of ice becaus	se of				
	(a)	dipole-dipol	le interaction							
	(b)	hydrogen bo	onding							
	(c)	dipole-indu	ced dipole inter							
	(d)	c) dipole–induced dipole interaction d) covalent bond formation								
	(- /	covalent bor	-	action						
44.	The f	ollowing mo	nd formation	ie sam		r weight. Which of				
44.	The f	ollowing mo	nd formation lecules have thest boiling point?	ie sam		· ·				
44.	The f them (a)	ollowing mo has the highe CH ₃ CH ₂ CO	nd formation lecules have thest boiling point?	ne sam ? (b)	e molecula	H ₂ CH ₂ OH				
	The f them (a) (c)	ollowing mo has the highe CH ₃ CH ₂ CO	nd formation lecules have the st boiling point? 2H 2-CH2-CH3	ne sam ? (b) (d)	e molecula CH ₃ CH ₂ CI CH ₃ CH ₂ CO	H ₂ CH ₂ OH OCH ₃				
	The f them (a) (c)	ollowing mo has the highe CH ₃ CH ₂ CO CH ₃ CH ₂ —C nitrophenol i orthonitropl	nd formation lecules have the st boiling point? 2H 0—CH2CH3 s steam volatile	te sam? (b) (d) but pa	e molecula CH ₃ CH ₂ CI CH ₃ CH ₂ CO aranitropher ular hydro	H ₂ CH ₂ OH OCH ₃ nol is not because gen bonding while				
	The f them (a) (c) Ortho	ollowing mo has the highe CH ₃ CH ₂ CO CH ₃ CH ₂ —C nitrophenol i orthonitropl	nd formation lecules have the st boiling point? 2H 0—CH2CH3 s steam volatile the nol has intra	te sam? (b) (d) but pa	e molecula CH ₃ CH ₂ CI CH ₃ CH ₂ CO aranitropher ular hydro	H ₂ CH ₂ OH OCH ₃ nol is not because gen bonding while				

- (b) both ortho- and paranitrophenol have intramolecular hydrogen bonding
- (c) orthonitrophenol has intermolecular hydrogen bonding and paranitrophenol has intramolecular hydrogen bonding
- (d) Van der Waals forces are dominant in orthonitrophenol
- **46.** The H bond in solid HF can be best represented as
 - (a) H-F--H-F--H-F
 - (b) H _ _ _ H _ _ H
 - (c) H F....H F....H
 - F H. F
- 47. For which of the following crystalline substances does the solubility in water increase upto 32°C and then decrease rapidly?
 - (a) CaCl₂·2H₂O

(b) Na₂SO₄·10H₂O

(c) FeSO₄·7H₂O

- (d) Alums
- **48.** Which of the following is true?
 - (a) Bond order $\propto \frac{1}{\text{bond length}} \propto \text{bond energy}$
 - (b) Bond order \sim bond length $\sim \frac{1}{\text{bond energy}}$
 - (c) Bond order $\propto \frac{1}{\text{bond length}} \propto \frac{1}{\text{bond energy}}$
 - (d) Bond order

 bond length

 bond energy
- **49.** Which of the following has been arranged in order of decreasing bond length?

 - (a) P O > Cl O > S O (b) P O > S O > Cl O

 - (c) S O > Cl O > P O (d) Cl O > S O > P O
- 50. During the formation of a molecular orbital from atomic orbitals, the electron density is
 - (a) minimum in the nodal plane
 - (b) maximum in the nodal plane
 - (c) zero in the nodal plane
 - (d) zero on the surface of the lobe

- 51. Which of the following represents correctly the order of filling of the molecular orbitals of B_2 , C_2 and N_2 ?
 - (a) $\pi_{2p_{\nu}} = \pi_{2p_{\nu}} < \sigma_{2p_{\nu}} < \pi_{2p_{\nu}}^* = \pi_{2p_{\nu}}^* < \sigma_{2p}^*$
 - (b) $\sigma_{2p_{a}} < \pi_{2p_{a}} = \pi_{2p_{a}} < \pi_{2p_{a}}^{*} = \pi_{2p_{a}}^{*} < \sigma_{2p_{a}}^{*}$
 - (c) $\sigma_{2p_{2}} = \pi_{2p_{2}} = \pi_{2p_{2}} < \pi_{2p_{2}}^{*} = \pi_{2p_{2}}^{*} < \sigma_{2p_{2}}^{*}$
 - (d) $\pi_{2p_{1}} < \pi_{2p_{2}} < \sigma_{2p_{2}} > \pi_{2p_{2}} < \pi_{2p_{2}}^{*} = \sigma_{2p_{2}}^{*}$
- **52.** The oxygen molecule is paramagnetic because
 - (a) the bonding electrons outnumber the antibonding electrons in the molecular orbital
 - (b) it contains unpaired electrons in the antibonding molecular orbitals
 - (c) it contains unpaired electrons in the bonding molecular orbitals
 - (d) the number of bonding electrons equals that of the antibonding electrons in the molecular orbitals
- 53. Which of the following have been arranged in increasing order of bond order as well as bond dissociation energy?
 - (a) $O_2^{-2} < O_2^{-} < O_2^{+} < O_2$
- (b) $O_2^{-2} < O_2^- < O_2 < O_2^+$
 - (c) $O_2 < O_2^+ < O_2^{2-} < O_2^-$
- (d) $O_2^+ < O_2^{2-} < O_2^- < O_2$
- **54.** Which of the following statements are not correct?
 - (a) Hybridization is the mixing of atomic orbitals.
 - (b) sp²-hybrid orbitals are formed from two p-atomic orbitals and one s-atomic orbital.
 - (c) dsp²-hybrid orbitals are all at 90° to one another.
 - (d) d²sp³-hybrid orbitals are directed towards the corners of a regular octahedron.
- 55. Which of the following has a pyramidal shape?
 - (a) PCl₃
- (b) SO₃
- (c) CO_3^{-2} (d) NO_3^{-1}
- 56. The compound in which carbon uses its sp³-hybrid orbitals for bond formation is
 - (a) HCO₂H

(b) (H₂N)₂CO

(c) HCHO

- (d) CH₃CHO
- 57. The hybridization of carbon involved in the C—C single bond in the molecule CH≡C—CH=CH₂ is
 - (a) sp^3-sp^2

(b) $sp^{3}-sp^{3}$

(c) $sp-sp^2$

(d) sp^2-sp^2

58. Which of the following has been arranged in increasing order of size of

(b) $sp^3 < sp^2 < sp$

(d) $sp^2 < sp < sp^3$

(b) sp²-hybridized

the hybrid orbitals? (a) $sp < sp^2 < sp^3$

(c) $sp^2 < sp^3 < sp$

59. The silver atom in $AgCl_2^-$ is

(a) sp³-hybridized

	(c)	sp-hybridize	ed		(d)	unhybridize	ed	
60.	(a) (b) (c)	napes of PCl ₄ ⁺ square plana tetrahedral, tetrahedral, trigonal bipy	ar, te see-s squa	trahedral ar saw and trig are planar ar	nd see onal b nd per	-saw pipyramidal ntagonal bipy	ram/	
61.	Amor	ng the followi	ng, t	he molecule	that i	s linear is		
	(a)	CO_2	(b)	NO_2	(c)	SO ₂	(d)	ClO ₂
62.		s not isostruct HgCl ₂			(c)	C_2H_2	(d)	$ZnCl_2$
63.		nolecule MX ₃ by M (atomic				ent, the sign	na bo	onding orbitals
		pure p				sp hybrid		
	(c)	sp ² hybrid			(d)	sp ³ hybrid		
64.		the context of carbon, which of the ler of electronegativity.				owing is arra	ange	d in the correct
	(a)	$sp > sp^2 > sp$	o ³		(b)	$sp^3 > sp^2 > s$	sp	
	(c)	$sp^2 > sp > sp$	o ³		(d)	$sp^3 > sp > sp$	p ²	
65.	PCl ₅ t	ındergoes						
	(a)	$sp^3 d_{z^2}$ -hybr	idiza	ation	(b)	$sp^{3} d_{x^{2}-y^{2}}-h$	ybri	dization
	(c)	$sp^3 d_{xy}$ -hybr	idiz	ation	(d)	sp 3 d _{yz} -hybi	ridiza	ation
66.		2s-2s, 2p-2 ases in the ord		nd 2p-2s o	rbital	s overlap, t	he l	oond strength
	(a)	p-p > s-s > 1	p–s		(b)	p-p > p-s >	s–s	
	(c)	s-s > p-p > 1	p–s		(d)	s-s > p-s > 1	р–р	
67.	The h	ybridization o	of ca	rbon in grap	hite i	S		
	(a)	$sp^2 - sp^2$			(b)	$sp^2 - sp^2 + 1$	2π	
	(c)	$sp^{2} - sp^{2} + \frac{3}{3}$	$\frac{\pi}{3}$		(d)	$sp^2 - sp^3 +$	$\frac{\pi}{2}$	

68.		n of the following pairs of eleations respectively?	emei	nts form oxide polyanions and
	(a)	Si and Al	(b)	Cu and Si
	(c)	Al and B	(d)	Ti and As
69.	Which	n of the following kinds of hybr	ridiz	ation are possible in $> C = C = ?$
	(a)	sp^2-sp^2	(b)	sp^2-sp
	(c)	sp-sp ³	(d)	sp-sp
70.	The h in the		liam	ond, graphite and acetylene are
	(a)	sp^3 , sp , sp^2	(b)	sp^3 , sp^2 , sp
	(c)	sp, sp^2, sp^3	(d)	sp^2, sp^3, sp
71.	Which	n of the following pairs are isos	truc	tural?
	(a)	CH ₃ ⁻ and CH ₃ ⁺	(b)	NH ₄ ⁺ and NH ₃
	(c)	SO ₄ ²⁻ and BF ₄ ⁻	(d)	NH ₂ and BeF ₂
72.	How	many sigma and pi bonds are pr	esen	t in tetracyanoethylene?
	(a)	Nine σ and nine π	(b)	Five π and nine σ
	(c)	Nine σ and seven π	(d)	Eight σ and eight π
73.	The sl	hape of XeF ₄ is		
	(a)	tetrahedral	(b)	square planar
	(c)	pyramidal	(d)	nearly linear
74.	The sl	hape of XeOF ₄ is		
	(a)	square pyramidal	(b)	square antiprismatic
	(c)	distorted octahedral	(d)	pentagonal bipyramidal
75.	The st	tructure of XeO ₂ F ₂ is		
		plane triangular		trigonal bipyramidal
	(c)	square planar	(d)	tetrahedral
76.		e atom in XeOF ₂ involves the h	-	
		sp ³		sp ³ d
	(c)	sp^3d^2	(d)	sp^3d^3
77.		olatility of HF is low because o	f	
		its low polarizability		
		the weak dispersion interaction	n be	etween the molecules
	` '	its small molecular mass		
	(a)	its strong nyarogen honding		

78.	. The shapes of ${\rm IF}_5$ and ${\rm IF}_7$ are respectively									
	(a) (b)	tetragonal py octahedral ar			nal bipyram	idal				
	(c)	trigonal bipy			ntiprismatic					
	` '	distorted squ		•	-					
79.	The N	—O—N bond	d angle is ma	ximum i	n					
	(a)	NO_2^+		(b)	NO ₂					
	(c)	NO_2^-		(d)	N_2O_3					
80.	_	n of the follow of bond angle		ements c	correctly rep	resen	t a decreasi	ing		
	(a)	$NH_3 > PH_3 >$	· AsH ₃	(b)	$NH_3 > H_2O$	$> F_2$)			
	(c)	$NO_2^+ > NO_2^-$	> NO ₂	(d)	$CH_4 > NH_3$	$> H_2$	0			
81.	(a) (b) (c)	n of the follow 1,4-dichlorob cis-1,2-dichlorost trans-1,2-dich trans-1,2-dich	penzene proethene hloroethene		ent?					
82.		n of the follow CCl ₄	ving has max (b) CH ₃ Cl	-	pole momen CH ₂ Cl ₂		CHCl ₃			
83.	(a) (b) (c) (d)	napes of XeF ₆ , octahedral, to square pyran square plana see-saw, T-sh ctively.	rigonal bipyr midal, pentaş ar, planar per	ramidal a gonal bip ntagonal	yramidal and and square a	d oct	ahedral			
84.		n of the follow HF	ving is the lea (b) HBr	nst polar? (c)		(d)	HCl			
85.	Which mome (a) (b) (c)	n of the follow	ving has been 3F > CH ₃ Br > Cl > CH ₃ Br > 3Br > CH ₃ I >	CH ₃ I CH ₃ I CH ₃ F	ed in order c			ole		
86.		n of the follow	_	_		(4)	NH.			

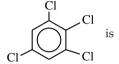
- **87.** The experimental value of the dipole moment of HCl is 1.03 D. The length of the H—Cl bond is 1.275 Å. The percentage of ionic character in HCl is
 - (a) 43

(b) 21

(c) 17

- (d) 7
- **88.** The dipole moment of is 1.5 D.

The dipole moment of



- (a) 0 D
- (b) 1.5 D
- (c) 2.86 D
- (d) 2.25 D
- 89. Which of the following pairs of structures are resonance forms?
 - (a) $: N N \equiv N$ and : N = N = N
 - (b) $: N N \equiv N$: and : N N = N:
 - (c) $N N \equiv N$ and N N N
 - (d) All of these
- **90.** Among the following structures, which is not a permissible resonance form?
 - (a) + N O: CH₃ N O: CH₃

(b) $CH_2 = N = Q$: CH_3

(c) $CH_2 = N - O^{\frac{1}{2}}$ CH_3

- (d) $\bar{C}H_3 N = O$ CH_3
- **91.** Which of the following molecules is the most polar?
 - (a) CH₃NH₂

(b) (CH₃)₃CCl

(c) CH₃NO₂

(d) (CH₃)₃CH

- **92.** The I_3^- ion has
 - (a) five equatorial lone pairs on the central I atom and two axial bonding pairs in a trigonal bipyramidal arrangement

- (b) five equatorial lone pairs on the central I atom and two axial bonding pairs in a pentagonal bipyramidal arrangement
- (c) three equatorial lone pairs on the central I atom and two axial bonding pairs in a trigonal bipyramidal arrangement
- (d) two equatorial lone pairs on the central I atom and three axial bonding pairs in a trigonal bipyramidal arrangement
- **93.** Which of the following sets of characteristics leads to the increase in solubility of ionic substances?
 - (a) High dipole moment, strong attraction by an ion and large solvation energy
 - (b) Low dipole moment, weak attraction by an ion and high solvation energy
 - (c) High dipole moment, strong attraction by an ion and low solvation energy
 - (d) High dipole moment, weak attraction by an ion and large solvation energy
- 94. Among the following species, which has the minimum bond length?(a) B₂(b) C₂
 - (c) F_2 (d) O_2^-
- 95. SnCl₄ is a covalent liquid because
 - (a) electron clouds of the Cl⁻ ions are weakly polarized to envelop the cation
 - (b) electron clouds of the Cl ¯ ions are strongly polarized to envelop the cation
 - (c) its molecules are attracted to one another by strong van der Waals forces
 - (d) Sn shows inert-pair effect
- **96.** The melting point of AlF_3 is $1040^{\circ}C$ and that of SiF_4 is $-77^{\circ}C$ (it sublimes) because
 - (a) there is a very large difference in the ionic character of the Al—F and Si—F bonds
 - (b) in AlF₃, Al $^{3+}$ interacts very strongly with the neighbouring F $^-$ ions to give a three-dimensional structure but in SiF₄ no interaction is possible
 - (c) the silicon ion in the tetrahedral SiF₄ molecule is not shielded effectively from the fluoride ions whereas in AIF₃, the Al³⁺ ion is shielded on all sides
 - (d) the attractive forces between the ${\rm SiF_4}$ molecules are strong whereas those between the ${\rm AlF_3}$ molecules are weak

• Type 2 •

(c) high solubilities in polar solvents and low solubilities in nonpolar

Choose the correct options. More than one option is correct.

(a) high melting points and low boiling points (b) high melting points and nondirectional bonds

97. Most ionic compounds have

		solvents							
	(d)	three-dimen electricity in				ires, and are	good	l conductor	s of
98.	Which	n of the follow	ving	substances	are ex	pected to be	cova	lent?	
	(a)	$BeCl_2$			(b)	$SnCl_4$			
	(c)	ZnS			(d)	$ZnCl_2$			
99.	Which	n of the follow	ving	have a three	e-dime	ensional netw	vork	structure?	
	(a)	SiO ₂			(b)	$(BN)_x$			
	(c)	P ₄ (white)			(d)	CCl_4			
100.	To wh	ich of the fol	lowi	ng species i	s the o	ctet rule not	appl	icable?	
	(a)	BrF_5			(b)	SF ₆			
	(c)	IF ₇			(d)	CO			
101.	Which	n of the follow	ving	do not exist	t?				
	(a)	SH_6			(b)	HFO_4			
	(c)	FeI_3			(d)	HClO ₃			
102.	-	pecies which nagnetic are	conf	tain an odd	numb	per of valence	e ele	ctrons and	are
	(a)	NO			(b)	NO ₂			
	(c)	ClO ₂			(d)	N_2O_4			
103.	Amon	g the followi	ng, t	he elements	whicl	h show inert-	pair	effect are	
	(a)	Bi			(b)	Sn	_		
	(c)	Pb			(d)	C			
104.	Which	n of the follow	ving	have an (18	s + 2)-e	lectron confi	gura	tion?	
	(a)	Pb^{2+}	(b)	Cd^{2+}	(c)	Bi ³⁺	(d)	SO_4^{2-}	
105.	Which	n of the follow	ving	species con	tain co	valent coord	inate	e bonds?	
	(a)	AlCl ₃	(b)	CO	(c)	$[Fe(CN)_6]^{4-}$	(d)	N_3^-	
106.	Which	n of the follow	ving	oxyacids of	sulph	ur contain S-	–S b	onds?	
	(a)	$H_2S_2O_8$	(b)	$H_2S_2O_6$	(c)	$H_2S_2O_4$	(d)	H_2SO_5	

- **107.** Which of the following factors are responsible for van der Waals forces?
 - (a) Instantaneous dipole-induced dipole interaction
 - (b) Dipole–induced dipole interaction and ion–induced dipole interaction
 - (c) Dipole-dipole interaction and ion-induced dipole interaction
 - (d) Small size of molecule
- **108.** Which of the following are true?
 - (a) Van der Waals forces are responsible for the formation of molecular crystals.
 - (b) Branching lowers the boiling points of isomeric organic compounds due to van der Waals forces of attraction.
 - (c) In graphite, van der Waals forces act between the carbon layers.
 - (d) In diamond, van der Waals forces act between the carbon layers.
- **109.** Which of the following statements are correct?
 - (a) The crystal lattice of ice is mostly formed by covalent as well as hydrogen bonds.
 - (b) The density of water increases when heated from 0°C to 4°C due to the change in the structure of the cluster of water molecules.
 - (c) Above 4°C the thermal agitation of water molecules increases. Therefore, intermolecular distance increases and water starts expanding.
 - (d) The density of water increases from 0°C to a maximum at 4°C because the entropy of the system increases.
- **110.** Intermolecular hydrogen bonding increases the enthalpy of vaporization of a liquid due to the
 - (a) decrease in the attraction between molecules
 - (b) increase in the attraction between molecules
 - (c) decrease in the molar mass of unassociated liquid molecules
 - (d) increase in the effective molar mass of hydrogen-bonded molecules
- 111. Which of the following molecules have intermolecular hydrogen bonds?
 - (a) KH_2PO_4

(b) H_3BO_3

(c) $C_6H_5CO_2H$

(d) CH₃OH

- 112. Intramolecular hydrogen bonds occur in
 - (a) 2-chlorophenol
 - (b) salicylic acid
 - (c) the enol form of acetylacetone
 - (d) paranitrophenol

113.	Which	n of the follow	ving	are diama	gnetic?				
	(a)	C_2			(b)	O_2^{2-}			
	(c)	Li ₂			(d)	N_2^+			
114.	Which	n of the follow	ving	are param	agnetic	?			
	(a)	B_2			(b)	O_2			
	(c)	N_2			(d)	He_2			
115.	Which	n of the follow	ving	species ha	ve a bo	nd order of 3	3?		
	(a)	CO			(b)	CN ⁻			
	(c)	NO ⁺			(d)	O_2^{+}			
116.	Amor	ng the followi	ng, t	he species	with or	ne unpaired	electr	on are	
	(a)	O_2^+	(b)	NO	(c)	O_2^-	(d)	B_2	
117.	Which	n of the follow	ving	pairs have	identic	cal values of	bond	order?	
	(a)	N_2^+ and O_2^+			(b)	F ₂ and Ne ₂			
	(c)	O_2 and B_2			(d)	C_2 and N_2			
118.	Which	n of the follow	ving	is correct?					
	(a)	During N ₂ ⁺ bonding mo				on each is	remo	oved from	the
	(b)	During O ₂ ⁺ antibonding				on each is	remo	oved from	the
	(c)	During O ₂ molecular o			electro	n each is ad	ded t	to the bond	ding
	(d)	During CN molecular o			e electr	on each is ac	lded	to the bond	ding
119.	Which	n of the follow	ving	species are	e linear	?			
	(a)	ICl ²⁻			(b)	I^{3-}			
	(c)	N_3^-			(d)	ClO ₂			
120.	The st	ructure of Xe	eF ₆ is						
	(a)	pentagonal	bipy	ramidal	(b)	distorted od	ctahe	dral	
	(c)	capped octa	hedı	al	(d)	square pyra	amida	al	
121.		n of the follow	_	have dipo	le mom	ent?			
		nitrobenzen				<i>p</i> -chloroniti			
	(C)	<i>m</i> -dichlorob	enze	ene	(d)	o-dichlorob	enzer	ne –	

1. a	2. b	3. a	4. d	5. b
6. c	7. d	8. a	9. b	10. a
11. b	12. c	13. b	14. b	15. c
16. c	17. b	18. c	19. d	20. d
21. b	22. d	23. c	24. a	25. c
26. b	27. b	28. a	29. b	30. c
31. d	32. c	33. c	34. c	35. a
36. d	37. a	38. d	39. b	40. b
41. a	42. a	43. b	44. a	45. a
46. c	47. b	48. a	49. b	50. c
51. a	52. b	53. b	54. c	55. a
56. d	57. c	58. a	59. c	60. b
61. a	62. b	63. c	64. a	65. a
66. b	67. c	68. a	69. b	70. b
71. c	72. a	73. b	74. a	75. b
76. b	77. c	78. a	79. a	80. a
81. b	82. b	83. c	84. c	85. a
86. b	87. c	88. b	89. a	90. b
91. c	92. c	93. a	94. b	95. b
96. b	97. a, b, c, d	98. a, b	99. a, b	100. a, b, c
101. a, b, c	102. a, b, c	103. a, b, c	104. a, c	105. b, c
106. b, c	107. a, b, c	108. a, b, c	109. a, b, c, d	110. b, d
111. a, b, c, d	112. a, b, c	113. a, b, c	114. a, b	115. a, b, c
116. a, b, c	117. a, c	118. a, b, d	119. a, b, c	120. b, c
121. a, b, c, d				

Hints to More Difficult Problems

- 1. A low ionization energy helps the formation of cations and a high electron affinity that of anions.
- The energy required to form the hydrogen atom is greater than that required to form the H₂ molecule, as H₂ molecules are held by strong forces of attraction.
- **12.** Among these pairs, the difference in electronegativity between Na and Cl atoms is the greatest.
- **17.** The least charge on Mn favours ionic character.

- **18.** Apply Fajans' rules (a large positive charge, and a small cation favour covalency).
- 23. Consider Sidgwick's covalency rule.

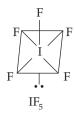
34.
$$O = P \stackrel{\longleftarrow}{\leftarrow} H$$
 (monoprotic acid)

- 37. Apply Fajans' rules.
- **40.** Hydrogen bonding is facilitated by the high electronegativity and small size of the atoms concerned.
- **44.** The tendency to form hydrogen bonds is greater in the —COOH group than in ethers, alcohols and ketones.
- **50.** The amplitude of the wave function is zero, and hence the probability of finding an electron in the region concerned is almost zero.
- 51. From considerations of energy
- 55. Ag has no unshared electrons.
- **63.** The molecule MX_3 has the general formula AX_3 . The number of unshared electrons on M is zero and it is, therefore, sp²-hybridized.

69.
$$sp^2 sp > C = C =$$







$$F$$
 F
 F
 F
 F
 F

(tetragonal pyramidal)

(pentagonal bipyramidal)

79.
$$O = N = C$$

(lp-lp > lp-bp > bp-bp) where lp stands for lone pair and bp for bond pair.

- **80.** The increasing size and decreasing electronegativity of the central atom permit the bonding electrons to be drawn out further, thus decreasing repulsion between bonding pairs. This can also be seen by the energy difference i of the s and p orbitals of the central atoms.
- 84. The smallest difference in electronegativity is between H and I. One may also apply the H-annay and Smith equation percentage of ionic character = $16 \mid (X_A X_B) \mid +3.5, \mid (X_A X_B) \mid$, where $X_A X_B$ is the electronegativity difference between the atoms.

87.
$$\mu_{cal} = e \times l = (4.802 \times 10^{-10} \text{ esu})(1.275 \times 10^{-8} \text{ cm})$$

= 6.12 D (1 Debye = 10^{-18} esu cm).

Percentage of ionic character = $\frac{\mu_{obs}}{\mu_{cal}} \times 100 = \frac{1.03}{6.12} \times 100 \approx 17$.

88.



The μ 's of 2Cl and 5Cl are vectorically cancelled.

$$\mu^{2} = \mu_{1}^{2} + \mu_{2}^{2} + 2\mu_{1}\mu_{2}\cos\theta$$

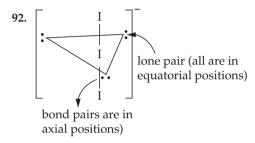
= $(1.5)^{2} + (1.4)^{2} + 2 \times 1.5 \times 1.5\cos 120^{\circ}$.

 $\therefore \quad \mu = 1.5 \text{ D}.$

89. The structures shown in option (a) are resonance forms, since they have the same atomic positions and the same number of electrons.

16 valence eletrons 16 valence eletrons (net charge = -1) (net charge = -1)

- **90.** The structure shown in (b) has 10 electrons surrounding nitrogen, but the octet rule limits nitrogen to 8 electrons.
- 91. The Lewis structure of CH_3NO_2 has a formal charge of +1 on nitrogen, making it more electron-attracting than the other structures.



- **94.** Among the species, C₂ has the maximum bond order (=2) and the maximum bond energy.
- 96. Structural factors
- 98. Apply Fajans' rules and also consider the effects of polarization.
- **100.** BrF_5 (14 electrons), SF_6 (12 electrons), IF_7 (14 electrons)
- 102. NO (5+6=11 valence electrons) $NO_2 (5+12=17 \text{ valence electrons})$ $ClO_2 (7+12=19 \text{ valence electrons})$ All molecules have odd-electrons and are paramagnetic.



2-Chlorophenol

Salicylic acid

enol form of acetylacetone

(All molecules have intramolecular hydrogen bonds.)

- 113. From the molecular orbital theory, we know that these species have a bond order equal to zero. The number of unpaired electrons they possess is also zero. Hence they are diamagnetic.
- **114.** From the molecular orbital theory, we know that B_2 and O_2 have two unpaired electrons each. Hence they are paramagnetic.
- 119. The species in options (a), (b) and (c) are sp-hybridized and are, therefore, linear.

Gaseous State

• Type 1 •

Choose the correct option. Only one option is correct.

(a) high pressure and low temperature(b) low pressure and high temperature(c) high pressure and high temperature(d) low pressure and low temperature2. The energy of an ideal gas depends only on its

1. A gas behaves like an ideal gas at

(a) pressure

(c) number of moles

3.	The quantity $pV/(k_BT)$ represents the							
	(a)	number of molecules in the gas						
	(b)	mass of the gas						
	(c)	number of moles of the gas						
	(d)	translation energy of the gas						
4.		ressure p of a gas is plotted against its absolute temperature T for ifferent constant volumes, V_1 and V_2 . When $V_1 > V_2$, the						
	(a)	curves have the same slope and do not intersect						
	(b)	curves must intersect at some point other than $T = 0$						
	(c)	curve for V_2 has a lower slope than that for V_1						

(b) volume

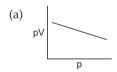
(d) temperature

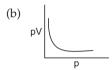
5. Which of the following graphs is not a straight line for an ideal gas? (a) $n \to T$ (b) $T \to p$ (c) $n \to \frac{1}{T}$ (d) $n \to \frac{1}{p}$

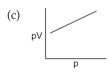
(d) curve for V_1 has a lower slope than that for V_2

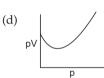
- 6. The density of nitrogen is maximum at
 - (a) stp

- (b) 273 K and 2 atm
- (c) 546 K and 1 atm
- (d) 546 K and 4 atm
- 7. A closed vessel is maintained at a constant temperature. It is first evacuated, and then an ideal gas is introduced into it continuously. The pressure of the gas inside the vessel will
 - (a) remain constant
 - (b) first increase and then decrease
 - (c) first increase and then remain constant
 - (d) make it burst after some time
- 8. Which of the following is a Boyle plot at very low pressure?









- 9. 1.0 litre of N_2 and $\frac{7}{8}$ litre of O_2 at the same temperature and pressure were mixed together. What is the relation between the masses of the two gases in the mixture?
 - (a) $M_{N_0} = 3M_{O_0}$

(b) $M_{N_2} = 8M_{O_2}$

(c) $M_{N_a} = M_{O_a}$

- (d) $M_{N_2} = 16M_{O_2}$
- **10.** The density of a gas A is twice that of a gas B at the same temperature. The molecular weight of gas B is thrice that of A. The ratio of the pressures acting on A and B will be
 - (a) 1:6
- (b) 7:8
- (c) 2:5
- (d) 1:4
- 11. A volume V of a gas at a temperature T_1 and a pressure p is enclosed in a sphere. It is connected to another sphere of volume V/2 by a tube and stopcock. The second sphere is initially evacuated and the stopcock is closed. If the stopcock is opened the temperature of the gas in the second sphere becomes T_2 . The first sphere is maintained at a temperature T_1 . What is the final pressure p_1 within the apparatus?

 - (a) $\frac{2pT_2}{2T_2 + T_1}$ (b) $\frac{2pT_2}{T_2 + 2T_1}$ (c) $\frac{pT_2}{2T_2 + T_1}$ (d) $\frac{2pT_2}{T_1 + T_2}$

12.	Two vessels A and B have volumes V and $4V$ respectively. Both vessels contain some water. The pressure in the space above water is p_1 for vessel A and p_2 for vessel B. What will be the relation between p_1 and p_2 ?							
	(a)	$p_1 = 4p_2$			(b)	$4p_1 = p_2$		
	(c)	$p_1 = p_2$			(d)	$p_1 = 16p_2$		
13.	Air contains 23% oxygen and 77% nitrogen by weight. The percentage of O_2 by volume is							tage of
	(a)	28.1			(b)	20.7		
	(c)	21.8			(d)	23.0		
14.	To wapplic		foll	owing g	aseous	mixtures is	Dalton's la	w not
	(a)	Ne + He + S	O_2		(b)	$NH_3 + HCl$	+ HBr	
	(c)	$O_2 + N_2 + C$	O_2		(d)	$N_2 + H_2 + C$)2	
15.		xture contain or density of			NO ₂ in	the ratio 2	:1 by volum	e. The
	` '	45.4				49.8		
	(c)	32.6			(d)	38.3		
16.	An open vessel at 27°C is heated until ³ %th of the air in it has been expelled. Assuming that the volume remains constant, calculate the temperature at which the vessel was heated.							
	(a)	307°C			(b)	107°C		
	(c)	480°C			(d)	207°C		
17.	0.0006	density of li o cm ⁻³ at 10 cules in 1 litre	0°C a	nd 1 atn	n, then	the volume	of water vap	oour is water
	(a)	$6 \mathrm{cm}^3$			(b)	60 cm^3		
	(c)	0.6 cm^3			(d)	0.06 cm^3		
18.	A gas	has a density	y of 2	.68 g L ⁻¹	at stp. Id	lentify it.		
	(a)	NO_2	(b)	Kr	(c)	COS	(d) SO_2	
19.		the reaction C(s	s) + H	₂ O(l)	→ CO(g)	$+ H_2(g)$		
			ne of	the gases			m 48.0 g of ca	rbon.
		179.2 L				89.6 L		
	. ,	44.8 L			` '	22.4 L		
20.	which	n has a vapou	ır der	sity of 36		ight in N ₂ O) ₄ (a dimer o	f NO ₂)
	(a)	27.7	(b)	67.7	(c)	37.7	(d) 25.7	

- Gaseous State 1 - 5921. A preweighed vessel was filled with CO₂ at stp and weighed. It was then evacuated, filled with SO₂ at the same temperature and pressure and again weighed. The weight of the CO₂ will be (a) the same as that of the SO₂ (b) twice that of the SO₂ (c) half that of the SO₂ (d) one-fourth that of the SO₂ 22. The vapour pressure of water at 80°C is 355 mm of Hg. A 100-mL vessel contains water saturated with O₂ at 80°C, the total pressure being 760 mm of Hg. The contents of the vessel were pumped into a 50-mL vessel at the same temperature. What is the partial pressure of O_2 ? (a) 1115 mm (b) 810 mm (c) 405 mm (d) 355 mm 23. A bottle of dry ammonia and one of dry hydrogen chloride are connected through a long tube. The stoppers at both ends of the tube are opened simultaneously. The white ammonium chloride ring first formed will be (a) at the centre of the tube (b) near the hydrogen chloride bottle (c) near the ammonia bottle (d) throughout the length of the tube **24.** At stp, 0.48 g of O_2 diffused through a porous partition in 1200 seconds. What volume of CO₂ will diffuse in the same time and under the same conditions? (a) 286.5 mL (b) 346.7 mL (c) 112.2 mL (d) 224.8 mL 25. At what temperature will the molar KE of 0.3 mol of He be the same as that of 0.4 mol of argon at 400 K?
 - - (a) 700 K
- (b) 500 K
- (c) 800 K
- (d) 400 K
- **26.** Indicate which of the following statements are correct.
 - (a) At constant temperature, the KE of all gas molecules is the same.
 - (b) At constant temperature, the KE of different molecules is different.
 - (c) At constant temperature, the KE is greater for heavier gas
 - (d) At constant temperature, the KE is less for heavier gas molecules.
- 27. The molecular velocities of two gases at the same temperature are u_1 and u_2 and their masses are m_1 and m_2 respectively. Which of the following expressions are correct?

(a)
$$\frac{m_1}{u_1^2} = \frac{m_2}{u_2^2}$$

(b)
$$m_1u_1 = m_2u_2$$

(c)
$$\frac{m_1}{u_1} = \frac{m_2}{u_2}$$

(d)
$$m_1 u_1^2 = m_2 u_2^2$$

becomes (a) u/2

(c) 4u

(a) smaller average speed

(c) greater kinetic energy

28. The rms speed of N₂ molecules in a gas is *u*. If the temperature is doubled and the nitrogen molecules dissociate into nitrogen atoms, the rms speed

29. Consider a mixture of SO_2 and O_2 kept at room temperature. Compared to the oxygen molecule, the SO_2 molecule will hit the wall with

(b) 2*u*

(d) 14u

(b) greater average speed

(d) greater mass

30.	The translational kinetic energy of an ideal gas depends only on its							
	(a)	pressure	(b)	force				
	(c)	temperature	(d)	molar mass				
31.	A bubble of gas released at the bottom of a lake increases to eight times its original volume when it reaches the surface. Assuming tha atmospheric pressure is equivalent to the pressure exerted by a column of water 10 m high, what is the depth of the lake? (a) 90 m (b) 10 m							
	(c)	70 m	(d)	80 m				
32.	. Choose the correct arrangement. The symbols have their usual meaning							
	(a)	$\overline{u} > u_p > u_{ m rms}$	(b)	$u_{\rm rms} > \overline{u} > u_p$				
	(c)	$u_p > \overline{u} > u_{\rm rms}$	(d)	$u_p > u_{\rm rms} > \overline{u}$				
33.	33. At what temperature is the rms speed of hydrogen molecules the sa that of oxygen molecules at 1327°C?							
		173 K	` '	100 K				
	(c)	400 K	(d)	523 K				
34.	The compressibility of a gas is less than unity at stp. Therefore,							
	(a)	$V_{\rm m} > 22.4 \text{ L}$	(b)	$V_{\rm m}$ < 22.4 L				
	(c)	$V_{\rm m}$ = 22.4 L	(d)	$V_{\rm m}$ = 44.8 L				
35.	. At low pressure, the van der Waals equation is reduced to							
	(a)	$Z = \frac{pV_{\rm m}}{RT} = 1 - \frac{ap}{RT}$	(b)	$Z = \frac{pV_{\rm m}}{RT} = 1 + \frac{b}{RT}p$				
	(c)	$pV_{\rm m} = RT$	(d)	$Z = \frac{pV_{\rm m}}{RT} = 1 - \frac{a}{RT}$				
36.	The temperature at which real gases obey the ideal gas laws over a wice range of pressures is called							
	(a)	critical temperature	(b)	inversion temperature				
	(c)	Boyle temperature	(d)	reduced temperature				

1-61

- 37. However great the pressure, a gas cannot be liquefied above its (a) Boyle temperature (b) inversion temperature (c) critical temperature (d) room temperature 38. At high temperature and low pressure, the van der Waals equation is (a) $\left(p + \frac{a}{V_{\rm m}^2}\right)(V_{\rm m}) = RT$ (b) $pV_{\rm m} = RT$ (d) $\left(p + \frac{a}{V_{\rm m}^2}\right) (V_{\rm m} - b) = RT$ (c) $p(V_{\rm m} - b) = RT$ 39. Real gases approach ideal gas behaviour at (a) low temperature and low pressure (b) low temperature and high pressure (c) high temperature and high pressure (d) high temperature and low pressure 40. In the corrections made to the ideal gas equation for red gases, the reduction in pressure due to attractive forces is directly proportional to (a) n/V(b) *nb* (d) n^2/V^2 (c) n^2/V^2b 41. Consider the following gases and their corresponding values of a given in brackets. $CH_4(2.25)$, $N_2(1.35)$, $O_2(1.36)$ and CO(1.46)Which of them will liquefy with great difficulty? (a) CH₄ (b) N_2 (c) O₂ (d) CO
 - **42.** Generally, the greater the value of the van der Waals constant *b* of a gas, the greater is the size of its molecule. Which of the following is an exception to this rule?
 - (a) $H_2(b = 0.0267)$

(b)
$$He(b = 0.0241)$$

(c) $CO_2(b = 0.0427)$

(d)
$$Ne(b = 0.0169)$$

- **43.** Consider the equation $Z = \frac{pV}{RT}$. Which of the following statements is correct?
 - (a) When Z > 1, real gases are easier to compress than the ideal gas.
 - (b) When Z = 1, real gases get compressed easily.
 - (c) When Z > 1, real gases are difficult to compress.
 - (d) When Z = 1, real gases are difficult to compress.

44. The van der Waals equation of state is

$$\left(p + \frac{a}{V^2}\right)(V - nb) = nRT.$$

The pressure exerted by individual gas molecules on the walls of the container depend on the

- (a) frequency of the collisions of the molecules with the walls as well as the momentum imparted by the molecules to the walls.
- (b) frequency of molecular collision
- (c) mean free path of the molecules
- (d) momentum and critical pressure of the gas molecules
- **45.** Which of the following statements is incorrect?
 - (a) It is not possible to compress a gas at a temperature below T_c .
 - (b) At a temperature below T_c , the molecules are close enough for the attractive forces to act, and condensation occurs.
 - (c) No condensation takes place above T_c .
 - (d) The kinetic energy of the gas molecules is higher above T_c, and the attraction between them decreases.
- **46.** The van der Waals equation for n = 1 mol may be expressed as

$$V_{\rm m}^3 - \left(b + \frac{RT}{p}\right)V_{\rm n}^2 + \frac{aV_{\rm n}}{p} - \frac{ab}{p} = 0,$$

where $V_{\rm m}$ is the molar volume of the gas. Which of the following is correct?

- (a) For a temperature less than $T_{c'}$ V has three real roots.
- (b) For a temperature less than $T_{\rm c}$, V has one real and two imaginary roots.
- (c) For a temperature equal to $T_{\rm c}$ all three roots of V are real and identical.
- (d) All of these
- 47. At the critical point,

(a)
$$\left(\frac{\partial p}{\partial V_{\rm m}}\right)_T = 1$$
 and $\left(\frac{\partial^2 p}{\partial V_{\rm m}^2}\right)_T = 0$

(b)
$$\left(\frac{\partial p}{\partial V_{\rm m}}\right)_T = 0 \text{ and } \left(\frac{\partial^2 p}{\partial V_{\rm m}^2}\right)_T = 0$$

(c)
$$\left(\frac{\partial T}{\partial V_{\rm m}}\right)_p = 0$$
 and $\left(\frac{\partial^2 T}{\partial V_{\rm m}^2}\right)_p = 0$

(d)
$$\left(\frac{\partial p}{\partial V_m}\right)_T = 0$$
 and $\left(\frac{\partial p}{\partial V_m}\right)_T = 0$

where $V_{\rm m}$ is the molar volume of the gas.

- **48.** The density of sea-water is 1.03 g cm^{-3} and that of freshwater 1.00 g cm^{-3} . The height of a seawater column and the pressure exerted by it can be
 - (a) 0.10 m and 0.1 atmosphere
- (b) 1.0 m and 1 atmosphere
- respectively.
- (c) 10.0 m and 1.0 atmosphere (d) 12.0 m and 2 atmospheres
- 49. Let p and p_s be the partial pressure and saturated partial pressure of water respectively. Then the relative humidity is given by
 - (a) $\frac{p_{\rm s} + p}{n} \times 100$

(b) $\frac{p}{p_0} \times 100$

(c) $\frac{p_s}{p} \times 100$

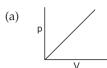
- (d) $(p + p_s) \times 100$
- **50.** Which of the following statements is correct?
 - (a) In effusion, molecular flux is directly proportional to molecular speed and inversely proportional to the square root of the molecular mass.
 - (b) In diffusion as well as effusion, ΔS (change in entropy) tends to become minimum.
 - (c) In effusion, molecular flux as well as molecular speed are inversely proportional to the square root of molecular mass.
 - (d) All of these
- **51.** At Boyle temperature,
 - (a) the effects of the repulsive and attractive intermolecular forces just offset each other
 - (b) the repulsive intermolecular forces are greater than the attractive intermolecular forces
 - (c) the repulsive intermolecular forces are less than the attractive intermolecular forces
 - (d) $b \frac{a}{RT} > 0$

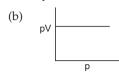
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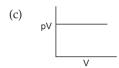
Choose the correct option. More than one option is correct.

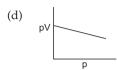
- 52. Which of the following quantities is the same for all ideal gases at the same temperature?
 - (a) The kinetic energy of 1 mol
 - (b) The kinetic energy of 1 g
 - (c) The number of molecules in 1 mol
 - (d) The number of molecules in 1 g

53. Which of the following graphs represent Boyle's law?

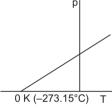


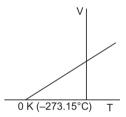






- 54. The value of the molar gas constant is
 - (a) $8.3145 \times 10^3 \text{ J (kg mol)}^{-1} \text{ K}^{-1}$
 - (b) 1.987 cal mol K⁻¹
 - (c) 0.083145×10^3 dm³ bar mol⁻¹ K⁻¹
 - (d) $0.083145 \text{ dm}^3 \text{ bar mol}^{-1} \text{ K}^{-1}$
- 55. What conclusion would you draw from the following graphs?





- (a) As the temperature is reduced, the volume as well as the pressure increase.
- (b) As the temperature is reduced, the volume becomes zero and the pressure reaches infinity.
- (c) As the temperature is reduced, the volume as well as the pressure decrease.
- (d) A point is reached where, theoretically, the volume as well as the pressure become zero.
- **56.** Which of the following statements are correct on the basis of Charles's law?
 - (a) The volume of an ideal gas can never be zero.
 - (b) The pressure of an ideal gas can be zero.
 - (c) At zero pressure, all molecular motion ceases in a gas, and it does not exert any pressure on the walls of the container.
 - (d) It is not possible to attain absolute zero.

57. Which of the following expressions is correct on the basis of the ideal gas equation?

(a)
$$pV = \frac{N}{N_A}RT$$
 (b) $pV = Nk_BT$ (c) $pV = \frac{\rho}{m}RT$ (d) $pV = \frac{Mk_BT}{m}$

- **58.** Which of the following statements are correct?
 - (a) Helium diffuses at a rate 8.65 times as much as CO does.
 - (b) Helium escapes at a rate 2.65 times as fast as CO does.
 - (c) Helium escapes at a rate 4 times as fast as CO₂ does.
 - (d) Helium escapes at a rate 4 times as fast as SO₂ does.
- 59. According to the kinetic theory of gases,
 - (a) the pressure exerted by a gas is proportional to the mean square speed of the molecules
 - (b) the pressure exerted by a gas is proportional to the root mean square speed of the molecules
 - (c) the root mean square speed is inversely proportional to the temperature
 - (d) the mean translational kinetic energy of the molecule is directly proportional to the absolute temperature
- **60.** Indicate the correct statement for equal volumes of $N_2(g)$ and $CO_2(g)$ at 298 K and 1 atm.
 - (a) The average translational KE per molecule is the same for N_2 and CO_2 .
 - (b) The rms speed remains constant for both N_2 and CO_2 .
 - (c) The density of N_2 is less than that of CO_2 .
 - (d) The total translational KE of both N_2 and CO_2 is the same.
- **61.** A gas can be easily liquefied
 - (a) when its inversion temperature equals the Boyle temperature
 - (b) under reversible adiabatic expansion
 - (c) under pressure when it is cooled to below the critical temperature
 - (d) at low pressure and above the critical temperature
- **62.** Which of the following is correct for critical temperature?
 - (a) It is the highest temperature at which liquid and vapour can coexist.
 - (b) Beyond this temperature, there is no distinction between the two phases and a gas cannot be liquefied by compression.
 - (c) At this temperature, the surface tension of the system is zero.
 - (d) At this temperature, the gas and the liquid phases have different critical densities.

Answers

1. b	2. d	3. a	4. d	5. d
6. b	7. c	8. d	9. c	10. a
11. a	12. a	13. b	14. b	15. d
16. d	17. c	18. c	19. a	20. a
21. c	22. b	23. b	24. a	25. d
26. a	27. d	28. b	29. d	30. c
31. c	32. b	33. b	34. b	35. a
36. c	37. c	38. b	39. d	40. d
41. b	42. d	43. c	44. a	45. a
46. d	47. b	48. c	49. b	50. a
51. a	52. a, c	53. b, c	54. a, d	55. c, d
56. a, c, d	57. a, b	58. b, d	59. b, c	60. a, c, d
61. b, c	62. a, b, c			

Hints to More Difficult Problems

3.
$$pV = nRT$$
. (i) $N = nN_A$. (ii) In Equations (i) and (ii),
$$pV = \frac{N}{N_A}RT = N\frac{R}{N_A}T$$

$$pV = Nk_BT \Rightarrow N = \frac{pV}{k_BT}.$$

where N = number of molecules.

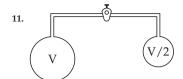
4.
$$pV = nRT$$
 or $p = \left(\frac{nR}{V}\right)T$.
For a $p-T$ graph, the slope $= \frac{nR}{V}$. (The equation is in the form $y = mx$.)
Thus, the slope $\propto \frac{1}{V}$.

6. Density =
$$\frac{pM}{RT}$$
.

The density of a given mass of gas $\propto p \propto \frac{1}{T}$.

9. 22.4 L of
$$N_2$$
 at stp = 28 g of N_2
1 L of N_2 at stp = 1.25 g of N_2 and 22.4 L of O_2 at stp = 32 g of O_2 .
7/8 litre of O_2 at stp = 1.25 g of O_2 .
 $\therefore M_{N_2} = M_{O_2}$.

Gaseous State 1-67



Let the total number of molecules of the gas be n, of which n_1 are in the larger sphere and n_2 in the smaller sphere after the stopcock is opened

$$n = n_1 + n_2 \text{ and } pV = nRT$$

$$\frac{pV}{RT_1} = \frac{p'V}{RT_1} + \frac{p'V}{2T_2R}$$

$$p' = \frac{2pT_2}{2T_2 + T_1}.$$

13. Percentage of O₂ by volume =
$$\frac{V_{\text{O}_2}}{V_{\text{O}_2} + V_{\text{N}_2}} = \frac{22.4 \left(\frac{23}{32}\right)}{22.4 \left(\frac{23}{32}\right) + 22.4 \left(\frac{27}{28}\right)} = 20.7.$$

(The molecular weights of O₂ and N₂ are 32 and 28 respectively.)

15.
$$V_{\text{N}_2\text{O}_4} = 100 \times \frac{2}{3} \text{ and } V_{\text{NO}_2} = 100 \times \frac{1}{3}$$

By the law of mixtures,

mass of N_2O_4 + mass of NO_2 = mass of mixture.

The vapour density of NO_2 is 23 and that of N_2O_4 is 4.

$$\frac{200}{3} \times 46 + \frac{100}{3} \times 23 = 100 \times d_{\text{mix}}$$

where d_{mix} is the vapour density of the mixture.

 \therefore vapour density d = 38.3.

16. Using
$$n_1 T_1 = n_2 T_2$$
, $T_2 = \frac{n_1}{n_2} T_1 = \frac{n}{(1 - 3/8)n} \times 300 = \frac{n}{(5/8)n} \times 300 = 480 \text{ K.}$
 $\therefore T_2 = 207^{\circ}\text{C.}$

17. Volume occupied =
$$\frac{0.0006 \text{ cm}^{-3} \times 10^{3} \text{cm}^{3}}{1.0 \text{ g cm}^{-3}} \times 1$$

= 0.6 cm³.

18.
$$1 L = 2.68 g$$

 $22.4 L = 2.68 \times 22.4 = 60.$
The molecular weight corresponds to that of COS.

22. Volume of $O_2 = 100$ mL and $p_{O_2} = 760 - 355 = 405$ mm. Since the temperature is constant, using Boyle's law we get

 $100 \text{ mL} \times 405 \text{ mm} \equiv 50 \text{ mL} \times p'_{O_2}$

:.
$$p'_{O_2} = 810 \text{ mm}$$

24. Volume of
$$O_2$$
 diffused = $\frac{22400 \times 0.48}{32}$ = 336 mL.

Let the volume of CO_2 diffused be x mL.

Rate of diffusion of $O_2 = \frac{336}{1200}$ mL s⁻¹.

Rate of diffusion of $CO_2 = \frac{x}{1200}$ mL s⁻¹.

$$\frac{r_{\text{O}_2}}{r_{\text{CO}_2}} = \frac{V_{\text{O}_2}/t}{V_{\text{CO}_2}/t} = \sqrt{\frac{M_{\text{CO}_2}}{M_{\text{O}_2}}}$$
or
$$\frac{\frac{336}{1200}}{\frac{x}{1200}} = \sqrt{\frac{44}{32}}.$$

$$\therefore x = 286.5 \text{ mL}$$

28.
$$(U_{\text{rms}})_1 = \sqrt{\frac{3RT_1}{M_1}}$$
 for N_2 molecule, mol. wt. $M_1 = 28$.

$$(U_{\text{rms}})_2 = \sqrt{\frac{3RT_2}{M_2}}$$
 for N atom, $M_2 = 14$.

$$\frac{(U_{\rm rms})_1}{(U_{\rm rms})_2} = \frac{\sqrt{\frac{3RT_1}{M_1}}}{\sqrt{\frac{3RT_2}{M_2}}} = \sqrt{\frac{3RT_1}{M_1} \times \frac{M_2}{3RT_2}} = \sqrt{\frac{T_2 \times 14}{28 \times 2T_2}} = \sqrt{\frac{1}{4}} = \frac{1}{2}.$$

$$(U_{\rm rms})_2 = 2(U_{\rm rms}).$$

31. Let the original volume of the bubble = V and the final volume = 8 V.

Let p be the atmospheric pressure and p_1 the pressure at the bottom of the lake.

Using Boyle's law, $p \times 8V = p_1 \times V \Rightarrow 8p = p_1$.

 p_1 = atmospheric pressure + pressure due to water in lake.

 \therefore pressure of water in lake = 7p.

Since p = pressure exerted by 10 m of water, the depth of the lake = 70 m.

Gaseous State 1-69

33.
$$U_{\text{rms}} = \sqrt{\frac{3RT}{M}}$$
.

According to the question, $\left(\sqrt{\frac{3RT_1}{M_1}}\right) = \left(\sqrt{\frac{3RT_2}{M_2}}\right)_{O_2}$.

 $\sqrt{\frac{T_1}{2}} = \sqrt{\frac{1600}{32}}$ or $\frac{T_1}{2} = \frac{1600}{32}$.

 $\therefore T_1 = 100 \text{ K}$.

35. The van der Waals equation for n moles of a real gas is given by

$$\left(p + \frac{n^2 a}{V^2}\right)(V - nb) = nRT \text{ or } \left(p + \frac{a}{V_{\rm m}^2}\right)(V_{\rm m} - b) = RT,$$

where $V_{\rm m}$ = molar volume = V/n.

At low pressure, $V_{\rm m}$ is high and so b can be neglected.

$$\begin{split} &\operatorname{The}\left(p+\frac{a}{V_{\mathrm{n}}^{2}}\right)V_{\mathrm{m}}=RT & \operatorname{or} & pV_{\mathrm{m}}=\frac{a}{V_{\mathrm{m}}}=RT \\ &\Rightarrow pV_{\mathrm{m}}=RT-\frac{a}{V_{\mathrm{m}}} & \Rightarrow \frac{pV_{\mathrm{m}}}{RT}=Z=1-\frac{a}{RTV_{\mathrm{m}}} \cdot \\ &Z=1-\frac{ab}{RT} \left(\because V \propto \frac{1}{p}\right). \end{split}$$

38. For *n* moles of a real gas, the van der Waals equation becomes

$$\left(p + \frac{a}{V_{\rm m}^2}\right)(V_{\rm m} - b) = RT$$

At high temperature and low pressure, $V_{\rm m}$ is large in comparison to b and $\frac{a}{V_{\rm m}^2}$ is negligibly small in comparison to p. Hence the above equation

is reduced to $pV_m = RT$.

- **41.** The less the value of a_i , the weaker is the intermolecular attraction.
- 43. Under this condition the intermolecular forces of attraction become weak and the molecules do not come close to each other. Hence real gases are difficult to compress.
- 61. (b) The expression for reversible adiabatic expansion is

$$\left(\frac{T_2}{T_1}\right)^{\frac{3}{2}} = \frac{V_1}{V_2} \Longrightarrow T^{\frac{3}{2}} \quad \propto \frac{1}{V} \quad \text{or} \quad V \propto \frac{1}{T^{\frac{3}{2}}}$$

So, on expansion, the gas cools.

(c) Greater intermolecular force of attraction

The Mole Concept

• *Type 1* •

1. Common salt obtained from sea-water contains 96% NaCl by mass. The approximate number of molecules present in 10.0 g of the salt is

2. When burnt in air, a 12.0-g mixture of carbon and sulphur yields a mixture of CO₂ and SO₂, in which the number of moles of SO₂ is half that

of CO₂. The mass of the carbon the mixture contains is

(b) 10²²

(d) 10²⁴

(b) 5.14 g

(d) 1.54 g

(b) $6.0 \times 10^{-22} \text{ mL}$

(d) $9.0 \times 10^{-23} \,\mathrm{mL}$

Choose the correct option. Only one option is correct.

(a) 10^{21}

(c) 10^{23}

(a) 4.08 g (c) 8.74 g

(a) $3.0 \times 10^{-23} \text{ mL}$

(c) $3.0 \times 10^{-21} \text{ mL}$

	(0)	0.7.18	(01)	110 1 8		
3.		experiment, it is found that 2.0 X_2O_5 . The number of moles of X_2O_5		g of pure X produces 3.6769 g of		
	(a)	0.04	(b)	0.06		
	(c)	0.40	(d)	0.02		
4.	. How many moles of $MgIn_2S_4$ can be made from 1.00 g of magnesium (of atomic mass = 24.0), 1.00 g of indium (of atomic mass = 114.8) and 1.00 g of sulphur (of atomic mass = 32.0)?					
	(a)	6.74×10^{-4}	(b)	3.1×10^{-2}		
	(c)	4.17×10^{-2}	(d)	8.7×10^{-3}		
5.		ensity of water at 4° C is $1.0 \times 10^{\circ}$		$^{-3}$. The volume occupied by		

6. When 0.5 mol of BaCl₂ is added to 0.2 mol of Na₃PO₄, the number of

7. A gaseous mixture contains $CO_2(g)$ and $N_2O(g)$ in a 2:5 ratio by mass. The ratio of the number of molecules of $CO_2(g)$ and $N_2O(g)$ is

(b) 0.20

(d) 0.15

(b) 2:5

moles of Ba₃(PO₄)₂ formed is

(a) 0.10

(c) 0.40

(a) 5:2

	(c)	1:2	(d)	5:4
8.	weigh			$_2$ Y ₃ and X ₃ Y ₄ . If 0.20 mol of X ₂ Y ₃ s 92.8 g, the atomic weights of X
	(a)	16.0 and 56.0	(b)	8.0 and 28.0
	(c)	56.0 and 16.0	(d)	28.0 and 8.0
9.				phite, the volume of the gases of moles of CO produced at stp.
	(a)	$\frac{1}{11.2}$		$\frac{28}{22.4}$
	(c)	$\frac{1}{22.4}$	(d)	$\frac{14}{22.4}$
10.	Sulph	ur trioxide is prepared by the f	ollo	wing two reactions.
		$S_8(s) + 8O_2(g) \longrightarrow 8$	SSO ₂	(g)
		$2SO_2(g) + O_2(g)$	> 2S0	$O_3(g)$
	How	many grams of SO ₃ are produc	ed fr	rom 1 mol of S ₈ ?
	(a)	1280.0	(b)	640.0
	(c)	960.0	(d)	320.0
11.		nntity of aluminium has a mas number of magnesium atoms?		54.0 g. What is the mass of the
	(a)	12.1 g	(b)	24.3 g
	(c)	48.6 g	(d)	97.2 g
12.		atomic weight of carbon is tadro constant will be	aken	to be 6 amu, the value of the
	(a)	$12.04 \times 10^{23} \text{ mol}^{-1}$	(b)	$3.01 \times 10^{23} \text{ mol}^{-1}$
	(c)	$1.5 \times 10^{23} \text{mol}^{-1}$	(d)	$6.02 \times 10^{23} \text{ mol}^{-1}$
13.	The cl	narge on 1 gram ion of Al ³⁺ is		
	(a)	$\frac{1}{27}N_{A}e$ coulomb	(b)	$\frac{1}{3} \times N_A e$ coulomb
	(c)	$\frac{1}{9}N_{\rm A}e$ coulomb	(d)	$3 \times N_{\rm A} e$ coulomb

14. How many moles of HCl will be present in 100 mL of a solution of

specific gravity 1.08, containing 20% HCl by mass?

	` ′	0.30			()	0.00		
	(c)			11: 6	` '	0.12		
15.		ensity in grar of methane a				e containing	an equa	l number of
		1.03				1.10		
	(c)	0.94			(d)	1.20		
16.		weights of e						
	(a)	$\frac{1}{2}$			(b)	$\frac{1}{1}$		
	(c)	1 16			(d)	$\frac{15}{16}$		
17.	n mol 1 atm	of N ₂ and 0.0 and 27°C. Fin	05 m nd <i>n</i> .	ol of Ar an $(R = 0.082)$	e enclos L atm	sed in a vess mol K ⁻¹ .)	sel of cap	pacity 2 L at
	(a)	0.30	(b)	0.10	(c)	0.03	(d) 0.0	06
18.	112.0 1.15 g liquid	mL of NO_2 mL ⁻¹ . Calcu NO_2 .	at st _] late t	p was liq he volum	uefied, e of and	the density the number	of the li	iquid being cules in the
	(a)	0.10 mL and	1 3.01	$\times 10^{22}$	(b)	0.20 mL and	d 3.01 × 1	10^{21}
	(c)	0.20 mL and	l 6.02	$\times 10^{23}$	(d)	0.40 mL and	d 6.02×	10 ²¹
19.	The m	nass of 1×10	²² mc	olecules of	CuSO ₄ .	5H ₂ O is		
	(a)	4.144 g			(b)	8.288 g		
	(c)	2.648 g			(d)	5.295 g		
20.		miconductor BaO ₂ and Cu						involving
	(a)	1:2:4				1:2:3		
	(c)	3:2:1			(d)	1:1.5:2.5		
21.	a mix	of iodine and ture of ICl an	d ICl	3. How ma	any mo			
	(a)				U			
		1.0 mol of IC			-			
	` '	0.5 mol of IO 0.5 mol of IO			0			
						20 111102		
22.	The n	umber of mo	Iecul.	es in 100 r	n L. ot ().(リス-N H ₂ S()。	1S	

(b) 6.02×10^{18}

(d) 6.02×10^{22}

(a) 6.02×10^{20}

(c) 6.02×10^{21}

25. Under the same conditions, two gases have the same number of

(b) 5 mol of R

(d) 13 mol of R

(b) $1.0 \text{ g of nitrogen } (N_2)$

(d) $1.0 \text{ g of water (H}_2\text{O})$

23. For the reaction $2P + Q \rightarrow R$, 8 mol of P and 5 mol of Q will produce

24. Which of the following contains the greatest number of atoms?

(a) 8 mol of R

(c) 4 mol of R

(a) 1.0 g of butane (C_4H_{10})

(c) 1.0 g of silver (Ag)

molecules. They must

	(a)	be noble gases		
	(b)	have equal volumes		
	(c)	have a volume of 22.4 dm ³ ea	ich	
	(d)	have an equal number of ator	ns	
26.	N_2O	nolar mass of N_2O as well as C contains n molecules of gas. T under the same conditions is	O ₂ is	s 44 g mol ⁻¹ . At 25°C and 1 atm, number of molecules in 2.0 L of
	(a)	$\frac{n}{4}$	(b)	$\frac{n}{8}$
	(c)	4 <i>n</i>	(d)	n
27.	(a) (b) (c)	flasks P and Q of the same cap ctively at 27°C and 1 atmosphe the same number of atoms as half the number of atoms as Q twice the number of atoms as gas of the same weight as Q d	ric p Q do Q do Q do	pes es
28.	acid a		roxid s	tely with an excess of sulphuric le. The ratio of the numbers of $3:2$
		2:1		9:4
29.	The n	umber of molecules of water ir	ı 333	g of Al ₂ (SO ₄) ₃ .18H ₂ O is
		$18.0 \times 6.02 \times 10^{23}$		$9.0 \times 6.02 \times 10^{23}$
		18.0	(d)	36.0
30.	300 k appro	C. The number of moles of eximately	NO ₂	ining NO_2 and N_2O_4 is 38.3 at in 100 g of the mixture is
	` '	0.44 33.4	` '	4.4 3.34
	(c)	33.4	(u)	J.J 4

31. The atomic weights of two elements A and B are 40 and 80 respectively. If x g of A contains y atoms, how many atoms are present in 2x g of B?

(c) *y*

(d) 2y

(b) $\frac{y}{4}$

(a) molecules present in 1 mL of a gas at stp
(b) molecules in 1 gram mole of a gas at stp
(c) atoms present in 1 mL of a gas at stp
(d) atoms present in 1 gram mole of a gas at stp

32. The Loschmidt number is the number of

	` '	1	0		0 1		
33.		of NaCl is of this solution	dissolved in 1 L on is	of pu	re water. The	e number of ion	s in
	(a)	6.02×10^{19}		(b)	1.2×10^{22}		
	(c)	1.2×10^{20}		(d)	6.02×10^{20}		
34.	than	the combine	ss of a $_{1}^{2}$ H atom is ed mass of a $_{1}^{1}$ l he missing mass	H ato			
	(a)	1.2 MeV		(b)	2.2 MeV		
	(c)	0.22 MeV		(d)	2.8 MeV		
35.	The re	est mass of 1	mol of neutrons	$(m_n =$	1.675×10^{-27}	kg) is	
	(a)	1.800×10^{-3}	kg	(b)	1.008×10^{-4}	kg	
	(c)	1.080×10^{-3}	kg	(d)	1.008×10^{-3}	kg	
36.		392 mg of H s of H_2SO_4 are	I_2SO_4 , 1.204 × 10 ³ e left?	²¹ mol	ecules are re	moved. How m	any
	(a)	2.0×10^{-3}	(b) 1.2×10^{-3}	(c)	4.0×10^{-3}	(d) 1.5×10^{-3}	
37.	Which	h of the follow	wing is arranged	in ord	der of increas	sing weight?	
	(a)		Fe < 0.0105 equ n < 0.006 g atom		nt of H ₂ C ₂ O ₂	$_{4}.2H_{2}O < 6.02 \times 1$	l0 ²¹
	(b)		Fe < 6.02 × 10 equivalent of H			< 0.006 g atom	of
	(c)	0.0105 e Ag < $6.02 \times$	quivalent of 10^{21} atom of Zn	H ₂ 0< 0.62	C ₂ O ₄ .2H ₂ O < 5 g of Fe	0.006 g atom	of
	(d)	0.0105 equi of Ag < 6.02	valent of $H_2C_2C_2$ $\times 10^{21}$ atoms	0 ₄ .2H ₂ 0	O < 0.625 g c	of Fe < 0.006 g a	tom
38.	and 3	300 K. The	$_4$ (g) and C_3H_8 (g) weight of the g of the numbers	as mi	xture in the	vessel is 0.613	
	(a)	1.54	(b) 1.68	(c)	1.44	(d) 1.60	

• Type 2 •

Choose the correct options. More than one option is correct.

39.	Which of the following expressions is correct ($n = no.$ of moles of the gas,
	N_A = Avogadro constant, m = mass of 1 molecule of the gas, N = no. of
	molecules of the gas)?

(a) $n = mN_A$

(b) $m = nN_A$

(c) $N = nN_A$

- (d) $m = mn/N_A$
- **40.** In which of the following pairs do 1 g of each have an equal number of molecules?
 - (a) N₂O and CO

(b) N_2 and C_3O_2

(c) N₂ and CO

- (d) N₂O and CO₂
- **41.** Among the following, which solutions contain equal numbers of millimoles?
 - (a) 100 mL of 0.05 M H₂SO₄
- (b) 200 mL of 0.05 M NaOH
- (c) $100 \text{ mL of } 0.010 \text{ M Na}_2\text{C}_2\text{O}_4$
 - (d) 200 mL of 0.025 M KOH
- **42.** 1 mol of ${}^{14}_{7}$ N ${}^{-3}$ ions contains
 - (a) $7N_A$ electrons

(b) $7N_A$ protons

(c) $7N_A$ neutrons

(d) $14N_A$ protons

39. b, c

40. c, d

- 43. 11.2 L of a gas at stp weighs 14.0 g. The gas could be
 - (a) N_2O

(b) NO₂

(c) N₂

(d) CO

Answers

- **1.** c **2.** b **4.** d **3.** a **5.** a 7. b **10**. b **6.** a 8. c **9.** c **11.** c **12.** b **13.** d **14.** b **15.** a **17.** c **16.** d **18.** b **19.** a **20.** b **21.** b **22.** a **23.** c **24.** a **25.** b **26.** d 27. b **28.** a **29**. b **30.** a **31.** c **32.** a **33.** c **34.** b 35. d
- **36.** a **41.** a, d
- **37.** b **42.** b, c
- **38.** a **43.** c, d

Hints to More Difficult Problems

- 4. The number of moles of Mg, In, and S are respectively $\frac{1}{24}$, $\frac{1}{114.8}$ and $\frac{1}{32}$ = 0.0417, 0.0087 and 0.031. The number of moles of the limiting reagent is $0.0087 = 8.7 \times 10^{-3}$.
- 5. $N = nN_A$. $1 = n \times 6.02 \times 10^{23}$ or $n = 1.66 \times 10^{-24}$ g or $wt = n \cdot (Mol. wt.) = 1.66 \times 10^{-24} \text{ mol} \times 18.0 \text{ g mol}^{-1}$ $= 3.0 \times 10^{-23} \text{ g}.$
- 8. Molecular weight of $X_2Y_3 = \frac{32.0 \text{ g}}{0.2 \text{ mol}} = 160 \text{ g mol}^{-1}$.

Molecular weight of $X_3Y_4 = \frac{92.8 \text{ g}}{0.4 \text{ mol}} = 232 \text{ g mol}^{-1}$.

Let atomic weight of X be a and that of Y be b. Then 2a + 3b = 160 and 3a + 4b = 232.

Solving, we get a = 56 and b = 16.

9.
$$CO_2(g) + C(s) \longrightarrow 2CO(g)$$

According to the question,

$$1 - x + 2x = 1.5 \implies x = 0.5 \text{ L} \implies 2x = 1.0 \text{ L} = 1/22.4 \text{ mol}.$$

- 13. 1 gram ion means 1 mole.
 - $\therefore \text{ charge} = 3 \times N_A \times e \text{ coulomb}.$
- **14.** Wt. of HCl solution = $100 \times 1.08 = 108.0$ g

Wt. of HCl =
$$\frac{20}{100} \times 108.0 = 21.6 \text{ g}$$

- \therefore no. of moles of HCl = $\frac{21.6}{36.5} \approx 0.6$.
- **16.** Let the total weight be x.

The fraction of the total pressure $=\frac{\frac{1}{2x}}{\frac{1}{2}x + \frac{1}{30}x} = \frac{15}{16}$.

18.
$$W_{\text{NO}_2} = \frac{112.0 \text{ mL} \times 46.0 \text{ g mol}^{-1}}{22400 \text{ mL mol}^{-1}} = 0.23 \text{ g}.$$

$$V_{\text{NO}_2, l} = \frac{0.23 \text{ g}}{1.15 \text{ g mL}^{-1}} = 0.20 \text{ mL}.$$

$$N = mN_A = \frac{0.23}{46} \times 6.02 \times 10^{23} = 3.01 \times 10^{21}.$$

- **22.** $100.0 \text{ mL of } 0.02 \text{ H}_2\text{SO}_4 = 100 \times 0.02 \text{ m eq.} = 2 \text{ m eq.} = 1 \text{ m mol}$ = $1 \times 10^{-3} \times 6.02 \times 10^{23} = 6.02 \times 10^{20} \text{ molecules.}$
- 31. No. of moles of A = $\frac{x}{40}$.

Number of atoms of A =
$$\frac{x}{40} \times N_A = y$$
 (say) or $x = \frac{40y}{N_A}$.

No. of moles of B = $\frac{2x}{80}$.

No. of atoms of B =
$$\frac{2x}{80}N_A = \frac{2}{80} \times \frac{40y}{N_A}N_A = y$$

33. Molarity of NaCl = $\frac{5.85}{58.5}$ = 0.1 M.

 $1 \text{ mL of NaCl} = 10^{-4} \text{ mol.}$

1 mole of NaCl = 6.02×10^{23} molecules of NaCl.

But NaCl molecules are dissociated into two ions (Na + and Cl -).

1 mol of NaCl = $6.02 \times 10^{23} \times 2$ ions.

$$1 \times 10^{-4}$$
 mole of NaCl = $1 \times 10^{-4} \times 6.02 \times 10^{23} \times 2$ ions = 1.2×10^{20} ions.

- **34.** Energy = 0.0024×931.5 MeV = 2.2 MeV.
- 35. Mass of 1 mole = $1.675 \times 10^{-27} \times 6.02 \times 10^{23}$ kg = 1.008×10^{-3} kg
- **36.** No. of moles of $H_2SO_4 = \frac{0.392}{98} = 0.004$.

$$N = nN_{\rm A} \implies n = \frac{N}{N_{\rm A}} = \frac{1.204 \times 10^{21} \text{ molecules}}{6.02 \times 10^{23} \text{ molecules}} = 0.002.$$

No. of moles of H_2SO_4 left = $0.004 - 0.002 = 0.002 = 2.0 \times 10^{-3}$.

38. Let the number of moles of C_2H_4 and C_3H_8 be x and y respectively. Using the formula

$$pV = nRT$$
,

1 atm \times 0.820 L = (x + y) mol \times 0.082 L-atm K⁻¹ mol⁻¹ \times 300 K

or
$$x + y = \frac{1}{30}$$
. (i)

Again,
$$28x + 44y = 0.613$$
. (ii) Solving, we get $\frac{y}{x} = 1.54$.

43. Calculate mol. wt. = $\frac{W}{V} \times 22.4 = \frac{14.0}{11.2} \times 22.4 = 28 \text{ g mol}^{-1}$.

This molecular weight corresponds to N₂(g) and CO(g).

Calculations Based on Chemical Equations and Eudiometry

• Type 1 •

1. 1.00×10^{-3} mol of Ag⁺ and 1.00×10^{-3} mol of CrO₄²⁻ react together to form solid Ag₂CrO₄. Calculate the amount of Ag₂CrO₄ formed

2. 1.00 g of $Cr_2O_7^{2-}$ is oxidized in an acidic solution by an excess of SO_2 to form HSO_4^- and Cr^{3+} . What is the minimum number of moles of H^+ that

must be produced for this reaction to occur ($Cr_2O^{2-} = 216$)?

(b) 0.166 g

(d) 1.66 g

(b) 0.0282

(d) 0.0268

Choose the correct option. Only one option is correct.

 $(Ag_2CrO_4 = 331.73 \text{ g mol}^{-1}).$

(a) 0.268 g

(c) 0.212 g

(a) 0.0231(c) 0.0322

(a) 0.236

3.	In an acidic solution, I change produced if, in the same proceed reduce H ₃ AsO ₄ to H ₃ AsO ₃ (I = 12)	ess, 1.5×10^{22} electrons	ny grams of I_2 are ons are used up to
	(a) 1.6 g	(b) 6.4 g	
	(c) 4.8 g	(d) 3.2 g	
4.	What would be the weight of t 8.0 g of ammonium chloride?	the slaked lime requ	uired to decompose
	(a) 5.53 g (b) 2.12 g	(c) 15.52 g	(d) 7.62 g
5.	The number of moles of $Cr_2O_7^{2-1}$ $N_2H_5^+$ through the reaction $N_2H_5^+ + Cr_2O_7^{2-1} \longrightarrow$		0.136 equivalent of

(c) 0.136

(d) 0.488

(b) 0.087

6. On being strongly heated, 2.76 g of Ag₂CO₃ yields a residue weighing

7. An impure sample of silver weighing 2.50 g is dissolved in HNO₃ and the silver is precipitated to yield 2.50 g of AgCl. What is the percentage by

weight of silver in the original sample (Ag = 108, Cl = 35.5)?

(b) 1.44 g (d) 4.16 g

(b) 100.00

(a) 3.48 g

(c) 2.16 g

(a) 75.26

	(c)	50.26			(d)	88.45		
8.		uric acid is prater in the pr				nur dioxide re	acts	s with oxygen
		$2SO_{2}(g) +$	O ₂ (g	$(3) + 2H_2O(1) -$	\longrightarrow	$2H_2SO_4(aq)$		
						O_2 and a large of H_2SO_4 that can		ccess of water, be obtained?
	(a)	5.6			(b)	11.2		
	(c)	2.4			(d)	1.4		
9.	passe		ion (of KI. Calcula				gas evolved is odine liberated
	(a)	7.7 g			(b)	15.4 g		
	(c)	12.7 g			(d)	25.4 g		
10.	to for (Fe =)	m FeS. Wha 55.85 g mol ⁻¹	at fr	action of the	ori ')?	ginal weight		ether and react left unreacted
	(a)	0.225			. ,	0.425		
	(c)	0.875			(d)	0.575		
11.		FeS_2 is burnt ne after the re			ir. C	alculate the pe	erce	ntage of N ₂ by
	(a)	81.94			(b)	9.89		
	(c)	8.17			(d)	89.26		
12.	liquor contai evapo (a)	containing	4% lids	(by weight).) ca Cal- apor (b)	ustic soda to culate the we	р	rentrates weak roduce a lye t of the water
13.	with o	calcium hydro containing 20	oxid	e. How many	/ litro amm	es of a solution nonia can be pi	n (s	g mol ⁻¹ reacts pecific gravity ared using this
	(a)	12.0 L	(b)	9.0 L	(c)	18.0 L	(d)	4.5 L

14. Calculate the number of millilitres (at stp) of hydrogen sulphide needed to precipitate cupric sulphide completely from 100 mL of a solution

15. Aluminium metal is prepared by the electrolysis of a solution of Al₂O₃ in molten cryolite (Na₃AlF₆). Assuming that all of the aluminium comes from Al₂O₃, how much of the latter would be needed for each tonne of

(b) 14.2

(d) 12.4

(b) 2.88 tonne

(d) 1.44 tonne

containing 0.75 g of CuCl₂ in a 1-L solution.

aluminium produced (Al = 27, O = 16)?

(a) 21.4

(c) 41.2

(a) 1.88 tonne

(c) 1.68 tonne

16.	2PbS	$+3O_2 \longrightarrow 2PbO + 2SO_2$		
	3SO ₂ -	$+ 2HNO_3 + 2H_2O \longrightarrow 3H_2SO$	4 + 2	NO
		eding to the above sequence of PbS produce?	of re	actions, how much H ₂ SO ₄ will
	(a)	245.2 g	(b)	490.4 g
	(c)	484.6 g	(d)	409.5 g
17.		inium dissolves according to th	e rea	
		$Al + 3H^+ \longrightarrow Al$	3++	$\frac{3}{2}$ H ₂
		1 L of H_{2} at stp. The percentage		0.50-g sample of the alloy gives luminium by weight in the alloy
	(a)	90	(b)	85
	(c)	78	(d)	96
18.	much			gentite (Ag ₂ S) by weight. How ed to obtain 1.00 g of pure solid
	(a)	45.7 g	(b)	67.4 g
	(c)	57.4 g	(d)	87.6 g
19.	perce the or		ng 7 tely	ms of carbon per molecule, the 70. The gram molecular mass of 365.0
	` '	415.0	. ,	667.0
20.	What		trog	en dioxide that can be produced
	(a)	4.60 g	(b)	2.30 g
	(c)	3.22 g	(d)	6.44 g

21. Calculate the volume required of a 20.0% HCl solution of density $1.20~g~mL^{-1}$ to prepare 363.0 g of AsCl $_3$ according to the equations (As = 75, Cl = 35.5)

$$2KMnO_4 + 16HCl \rightarrow 2KCl + 2MnCl_2 + 5Cl_2 + 8H_2O$$
 and
$$2As + 3Cl_2 \rightarrow 2AsCl_3$$

is

(a) 2.56 L

(b) 0.73 L

(c) 1.46 L

(d) 2.92 L

22. 1 L of an acidified solution containing 31.6 g of KMnO₄ is decolourized by passing SO_2 through it. How much iron pyrites (FeS₂) has to be roasted to produce the necessary amount of SO_2 (K = 39, Mn = 55, S = 32, Fe = 56)?

(a) 30.0 g

(b) 7.5 g

(c) 15.0 g

(d) 45.0 g

23. 2.0 g of dolomite was heated to a constant weight of 1.0 g. Calculate the total volume of the CO_2 produced at stp (Ca = 40, Mg = 24, C = 12, O = 16) by this reaction.

(a) 482.4 mL

(b) 502.6 mL

(c) 492.8 mL

(d) 428.6 mL

24. 1 g of a mixture of NaHCO₃ and Na₂CO₃ is heated to 150°C. The volume of the CO₂ produced at stp is 112.0 mL. Calculate the percentage of Na₂CO₃ in the mixture (Na = 23, C = 12, O = 16).

(a) 20

(b) 46

(c) 84

(d) 16

25. 2 g of impure CaCO₃ reacts with HCl to produce 410 mL of CO₂ at 1 atmospheric pressure and 27°C. Calculate the percentage purity of the CaCO₃ used.

(a) 83.5

(b) 97.5

(c) 87.5

(d) 73.5

26. 60 g of NaOH is converted into NaCl and NaClO₃ by the action of Cl₂. The Cl₂ is produced by the reaction between MnO_2 and concentrated HCl. The amount of MnO_2 required for the process (Mn = 55, Na = 23) is

(a) 70.95 g

(b) 25.65 g

(c) 65.25 g

(d) 75.45 g

27. The atomic weight of Cu is 63.546. There are only two naturally occurring isotopes of copper, ⁶³Cu and ⁶⁵Cu. The natural abundance of the ⁶³Cu isotope is approximately

(a) 20%

(b) 70%

(c) 30%

(d) 80%

28.	obtaine	- 1	ea c	opper. The volume of the $N_2(g)$
	(a)	15.0 mL	(b)	7.5 mL
	(c)	30.0 mL	(d)	45.0 mL
29.	yieldir the cor		stea ire.	ely burnt in an excess of oxygen, im. Calculate the percentages of $30\% \text{ CH}_4$ and $70\% \text{ C}_2\text{H}_4$
		75% CH_4 and 25% C_2H_4		50% CH ₄ and 50% C ₂ H ₄
30.	1 mol o	of a gaseous aliphatic compo ess of oxygen. The contraction	und	$C_nH_{3n}O_m$ is completely burnt in
	(a)	$\left(1 + \frac{1}{2}n - \frac{3}{4}m\right)$	(b)	$\left(1 + \frac{3}{4}n - \frac{1}{4}m\right)$
		$\left(1 - \frac{1}{2}n - \frac{3}{4}m\right)$		$\left(1 + \frac{3}{4}n - \frac{1}{2}m\right)$
31.	mixed the ga solutio	with 100.0 mL of oxygen gas, s after the explosion was 90	caus).0 n	nd containing C, H and O was sing an explosion. The volume of nL. On treatment with a KOH ne was observed. Calculate the roduced.
	(a)	1.0	(b)	3.0
	(c)	2.0	(d)	4.0
32.	O ₂ and Calcula		as a ginal	C ₂ H ₂ is mixed with 100.0 mL of fter the combustion is 10.5 mL. mixture.
	(b)	15 mL of CO and 25 mL of C_2	H_2	

33. What is the volume of air required for the complete combustion of 20.0 L

34. 200.0 mL of oxygen is added to 100.0 mL of a mixture containing CS_2 vapour and CO, and the total mixture is burnt. After combustion, the volume of the entire mixture is 245.0 mL. Calculate the volume of the

(b) 240 L

(d) 192 L

(b) 125.0 mL

(d) 100.0 mL

(c) 10 mL of CO and 30 mL of C_2H_2 (d) 20 mL of CO and 20 mL of C_2H_2

of methane?

(a) 840 L(c) 240 L

oxygen that remains. (a) 67.5 mL

(c) 200.0 mL

35. 80 mL of oxygen is added to 50 mL of a mixture of H₂, C₂H₂ and CO, after which the total mixture is burnt. The volume of the cooled mixture after combustion measures 65 mL. This is reduced to 15 mL by treatment with a KOH solution. Calculate the volume of each gas in the original mixture.

36. 30 mL of a gaseous hydrocarbon requires 90 mL of O₂ for complete combustion, 60 mL of CO₂ being formed in the process. The molecular

(b) C₂H₀

(a) 20 mL of H₂, 20 mL of C₂H₂, 10 mL of CO
(b) 10 mL of H₂, 20 mL of C₂H₂, 20 mL of CO
(c) 15 mL of H₂, 15 mL of C₂H₂, 20 mL of CO
(d) 20 mL of H₂, 25 mL of C₂H₂, 5 mL of CO

formula of the hydrocarbon is

(a) C₂H₂

	(c)	C_4H_{10}			(d)	C_2H_4		
37.	burnt 112.5 (a)		of th	ne cooled mi	xture	of gases aft original mix 60	th 150 mL of O ₂ arer the combustion ture is	
38.	carbo origin aqueo of the (a)	n, hydrogen al room temp	and perat	loxygen is ure, its volu	heat me b g 88 r (b)	ed. After becomes 100	and containing or being cooled to t mL. Treatment wi ne empirical formu	he ith
39.	oxyge potas formi	en. The obse h, there was ala of the hyd	rved a fu roca	contraction orther contra orthon?	was	$12^{1}/2V$, and of $2V$. Wh	ed with an excess on treatment winat is the molecule (d) C_2H_4	ith
40.	contra took hydro (a)	action of 60 n	nĽ w	as observed	l, and added (b)	l a further c	120 mL of oxygen. contraction of 60 n the formula of t	nL
41.	of me		Cal				ely reduced to 1.7 the original samp	
	(a)	0.85 g	(b)	0.55 g	(c)	0.75 g	(d) 0.95 g	

42.	A gaseous mixture of ethene and ethyne measuring 50 mL is mixed with 150 mL of O_2 and burnt. After combustion and cooling, the volume of the mixture of gases is 112.5 mL . Calculate the percentage by volume of ethene in the mixture.						
	(a) 75	(b) 60					
	(c) 50	(d) 80					
43.	50.0 mL of a gaseou	s mixture of H ₂ and HCl is exposed to a sodium					

- amalgam. The volume decreases to 42.5 mL. If 100.0 mL of the same mixture is added to 50.0 mL of gaseous ammonia and then exposed to water, what will be the volume of the final mixture?
 - (a) 35.0 mL

(b) 50.0 mL

(c) 15.0 mL

- (d) 70.0 mL
- 44. A gaseous organic compound containing C, H and N, which is completely burnt in an excess of oxyen, produces
 - (a) $x \text{ vol. of } CO_2(s) + \frac{y}{2} \text{ vol. of } H_2O(g) = \frac{z}{2} \text{ vol. of } NO_2(g)$
 - (b) $x \text{ vol. of } CO_2(g) + \frac{y}{2} \text{ vol. of } H_2O(l) \frac{z}{2} \text{ vol. of } N_2(g)$
 - (c) $x \text{ vol. of } CO_2 + \frac{y}{2} \text{ vol. of } H_2O(1)$
 - (d) $x \text{ vol. of } CO_2(s) + \frac{y}{2} \text{ vol. of } H_2O(s) + z \text{ vol. of } N_2(g)$
- 45. Three volumes of a gaseous hydrocarbon containing carbon, hydrogen and sulphur is burnt in an excess of oxygen to yield three volumes of CO₂, three volumes of SO₂ and six volumes of water vapour. The formula of the compound is
 - (a) C_6H_6S
- (b) C_4H_4S (c) CH_4S
- (d) C₂H₄S

• *Type 2* •

Choose the correct options. More than one option is correct.

- **46.** Calculate the amount of lime (CaO) produced by heating 100 g of 90% pure limestone.
 - (a) 50.4 g

(b) 0.98 mol

(c) 0.90 mol

- (d) 56.0 g
- **47.** 2 mol of CO₂ is required to prepare
 - (a) 336 g of NaHCO₃
- (b) 168 g of NaHCO₃
- (c) 462 g of Ca(HCO₃)₂
- (d) 162 g of Ca(HCO₃)₂

- **48.** 1.5 g of oxygen is produced by heating KClO₃. How much KCl is produced in the reaction?
 - (a) $4.15 \times 10^{-2} \text{ mol}$
- (b) 4.33 g
- (c) $1.78 \times 10^{-2} \text{ mol}$
- (d) 1.33 g
- **49.** Which of the following gases are absorbed by an ammoniacal cuprous chloride solution?
 - (a) NO

(b) CO

(c) O_3

- (d) C_2H_2
- **50.** 50 millilitres of CO is mixed with 20 mL of oxygen and sparked. After the reaction, the mixture is treated with an aqueous KOH solution. Choose the correct option.
 - (a) The volume of the CO that reacts = 40 mL.
 - (b) The volume of the CO_2 formed = 40 mL.
 - (c) The volume of the CO that remains after treatment with KOH = 10 mL.
 - (d) The volume of the CO that remains after treatment with KOH = 20 mL.

Answers

1. b	2. a	3. d	4. a	5. c
6. c	7. a	8. a	9. d	10. b
11. a	12. c	13. b	14. d	15. a
16. b	17. a	18. c	19. b	20. d
21. c	22. a	23. c	24. d	25. a
26. c	27. b	28. a	29. d	30. d
31. c	32. a	33. d	34. b	35. a
36. d	37. c	38. b	39. a	40. b
41. a	42. c	43. d	44. b	45. c
46. a, c	47. a, d	48. c, d	49. b, d	50. a, b, c

Hints to More Difficult Problems

- 1. The reaction is $2Ag^+ + CrO_4^{2-} \longrightarrow Ag_2CrO_4$. Using the limiting-reagent concept, no. of moles of $Ag_2CrO_4 = 0.5 \times 10^{-3} \times 331.73 = 0.166$ g.
- 2. The reaction is

$$Cr_2O_7^{2-} + 3SO_2 + 5H^+ \longrightarrow 3HSO_4^- + 2Cr^{3+} + H_2O$$

No. of moles of $H^+ = \frac{1.0}{216} \times 5 = 0.0231$ $[Cr_2O_7^{2-} = 2 \times 52 + 7 \times 16 = 216]$

4.
$$2NH_4Cl + Ca(OH)_2 \longrightarrow CaCl_2 + 2NH_3 + 2H_2O$$

 $2 \times 53.5 \text{ g}$ $(40 + 34) \text{ g}$
Wt. of $Ca(OH)_2$ required = $\frac{74}{107} \times 8 = 5.53 \text{ g}$.

7. Since the weights of the sample and the precipitated AgCl are equal, the required amount is given by the

weight percentate of silver in AgCl =
$$\frac{108}{108 + 35.5} \times 100 = 75.25$$

9.
$$MnO_2 + 4HCl \longrightarrow MnCl_2 + 2H_2O + Cl_2$$

$$Cl_2 + 2KI \longrightarrow 2KCl + I_2$$

The number of moles of MnO₂ equals that of I₂.

Thus, 87 g liberates 254 g of I₂.

Therefore 8.7 g of MnO₂ corresponds to 25.4 g of I₂.

10. Use the limiting-reagent concept.

12. 100 kg of weak liquor (feed) contains 4 kg of caustic soda. Let the quantity of the lye be x kg. Then the amount of caustic soda in the lye = 0.25 x. However caustic soda does not take part in the evaporation

$$\therefore$$
 0.25 $x = 4 \Rightarrow x = 16 \text{ kg}$

The weight of the water that evaporates = 100.0 - 16.0 = 84.0 kg

17. $1.5 \times 22400 \text{ mL of H}_2 \text{ at stp} = 27 \text{ g of Al}$ 560 mL of H₂ at stp = 0.45 g of Al

:. Percentage of Al =
$$\frac{0.45}{0.50} \times 100 = 90$$

18. Ag₂S has two atoms of Ag.

Therefore, 216 g of Ag corresponds to 248 g of Ag₂S.

1 g of Ag is contained in =
$$\frac{248}{216}$$
 g of Ag₂S

wt. of the ore =
$$\frac{248}{216} \times \frac{100}{2} = 57.4 \text{ g}$$

19. Gram molecular mass of carbon

$$= \frac{\text{no. of atoms} \times \text{At. mass of C} \times 1 \text{ amu} \times N_{\text{A}}}{\text{percentage of carbon}}$$

$$= \frac{20 \times 12 \times 1.66 \times 10^{-24} \text{ g} \times 6.02 \times 10^{23} \text{ mol}^{-1}}{70/100}$$

$$= 365.0 \text{ g mol}^{-1}$$

20. No. of moles of NO =
$$\frac{4.2 \text{ g}}{30}$$
 = 0.14

No. of moles of
$$O_2 = \frac{3.2 \text{ g}}{32} = 0.10$$

$$2NO(g) + O_2(g) \longrightarrow 2NO_2(g)$$

Using the limiting reagent concept,

$$W_{NO_2} = 0.14 \text{ mol} \times 46 \text{ g mol}^{-1}$$

= 6.44 g

22.
$$4\text{FeS}_2 + 11\text{O}_2 \longrightarrow 2\text{Fe}_2\text{O}_3 + 8\text{SO}_2$$

 $2\text{KMnO}_4 + 5\text{SO}_2 + 2\text{H}_2\text{O} \longrightarrow \text{H}_2\text{SO}_4 + 2\text{MnSO}_4 + 2\text{H}_2\text{SO}_4$
 $\frac{8}{5}(2\text{KMnO}_4) \equiv 8\text{SO}_2 \equiv 4\text{FeS}_2$

$$\frac{16}{5}$$
 mol of KMnO₄ \equiv 4 mol of FeS₂

No. of moles of KMnO₄ =
$$\frac{31.6}{158}$$
 = 0.2

0.2 mol of KMnO₄
$$\equiv \frac{4}{16/5} \times 0.2$$
 mol of FeS₂

$$= \frac{1}{4} \text{ mol of FeS}_2 = \frac{1}{4} \times 120 \text{ g}$$

$$= 30.0 \text{ g}$$

23.
$$CaCO_3 \longrightarrow CaO + CO_2$$
 $MgCO_3 \longrightarrow MgO + CO_2$
 100.0 g 56.0 g 84.0 g 40.0 g
 $x \text{ g}$ $\frac{56}{100} x \text{ g}$ $(2.0 - x) \text{ g}$ $\frac{40.0}{84.0} (2.0 - x) \text{ g}$

According to the question,

$$\frac{56.0}{100.0}x + \frac{40.0}{84.0}(2.0 - x) = 1$$
 $\therefore x = 1.0 \text{ g}$ $2.0 - x = 1.0 \text{ g}$

Total no. of moles of
$$CO_2 = \frac{1.0}{100.0} + \frac{1.0}{84.0} = 0.022$$

= $0.022 \times 22.4 \text{ L} = 492.8 \text{ mL}$

24.
$$2\text{NaHCO}_3 \xrightarrow{150^{\circ}\text{C}} \text{Na}_2\text{CO}_3 + \text{CO}_2 + \text{H}_2\text{O}$$

$$\frac{n_{\text{NaHCO}_3}}{n_{\text{CO}_2}} = \frac{2}{1}$$

$$n_{\text{NaHCO}_3} = 2n_{\text{CO}_2} = 2 \times \frac{112}{22400} = 0.01 \text{ mole}$$

$$W_{\text{NaHCO}_3} = 0.01 \times 84 = 0.84 \text{ g}$$

$$W_{\text{Na}_2\text{CO}_3} = 1.00 - 0.84 = 0.16 \text{ g}$$
% $\text{Na}_2\text{CO}_3 = 16$

25.
$$n_{\text{CO}_2} = \frac{\text{pV}}{\text{RT}} = \frac{(1 \text{ atm})(0.410 \text{ L})}{(0.082 \text{ L atm mol}^{-1} \text{ K}^{-1})(300 \text{ K})} = 0.0167 \text{ mol}$$

$$1 \text{ mol of CO}_2 \text{ accounts for 1 mol of CaCO}_3 = 100 \text{ g of CaCO}_3$$

$$0.0167 \text{ mol of CO}_2 = 1.67 \text{ g of CaCO}_3$$
Percentage purity of CaCO₃ = $\frac{1.67}{2.00} \times 100 = 83.5$

26.
$$[MnO_2 + 4HCl \longrightarrow MnCl_2 + 2H_2O + Cl_2] \times 3$$

 $3Cl_2 + 6NaOH \longrightarrow 5NaCl + NaClO_3 + 3H_2O$
 $6NaOH \Longrightarrow 3MnO_2$
 $6 \times 40 \text{ g} = 3 \times 87 \text{ g}$
 $3 \text{ mol of } MnO_2 \text{ accounts for } 12 \text{ mol of } HCl \text{ and } 6 \text{ mol of } NaOH.$
 $\frac{60}{40} \text{ mol of } NaOH \text{ accounts for } 0.75 \text{ mol of } MnO_2$
 $= 0.75 \text{ mol} \times 87 \text{ g mol}^{-1}$
 $= 65.25 \text{ g}$

30.
$$C_n H_{3n} O_m(g) + \left(n + \frac{3}{4}n - \frac{m}{2}\right) O_2(g) \longrightarrow nCO_2(g) + \frac{3}{2}n H_2O(1)$$

Contraction of volume = $1 + n + \frac{3}{4}n - \frac{m}{2} - n = \left(1 + \frac{3}{4}n - \frac{m}{2}\right)$

31.
$$C_x H_y O_z(g) + \left(x + \frac{y}{4} - \frac{z}{2}\right) O_2(g) \longrightarrow x C O_2(g) + \frac{y}{2} H_2 O(l)$$

10.0 mL $10\left(x + \frac{y}{4} - \frac{z}{2}\right) mL$ 10 x mL zero

Volume of CO_2 produced = $10x = 20 \Rightarrow x = 2$

33.
$$CH_4(g) + 2O_2(g) \longrightarrow CO_2(g) + 2H_2O(1)$$

 $20.0 \text{ L} \quad 40.0 \text{ L}$
 $20.8 \text{ L} \text{ of } O_2 = 100.0 \text{ L of air}$
 $40.0 \text{ L of } O_2 = 192.0 \text{ L of air}$

36.
$$C_xH_y(g) + \left(x + \frac{y}{4}\right)O_2(g) \longrightarrow xCO_2(g) + \frac{y}{2}H_2O(1)$$

 $mL \longrightarrow 30.0 \quad 30.0\left(x + \frac{y}{4}\right) \quad 30.0x$

According to the question $30.0 x = 60.0 \Rightarrow x = 2$

and
$$30.0\left(x + \frac{y}{4}\right)O_2 \text{ mL} = 90.0 \text{ mL } O_2$$

or
$$y = 4$$

Therefore, the molecular formula of hydrocarbon = C_2H_4 .

43.
$$Na/Hg + 2HCl(g) \longrightarrow 2NaCl + H_2 + 2Hg$$

2 mol 1 mol

Contraction in volume = 50.0 - 42.5 = 7.5 mL of H₂(g)

1 mol of $H_2(g) \equiv 2 \text{ mol of } HCl(g)$

7.5 mL of $H_2(g) \equiv 15$ mL of HCl(g)

Volume of $H_2 = 50.0 - 15.0 = 35.0 \text{ mL}$ Volume of H_2 in 100.0 mixture $= \frac{100 \times 35}{50} = 70.0 \text{ mL}$

44. The reaction is

$$C_xH_yN_z(g) + \left(x + \frac{y}{4}\right)O_2(g) \longrightarrow xCO_2(g) + \frac{y}{2}H_2O(l) + \frac{1}{2}zN_2(g)$$

45. The reaction is

$$C_xH_yS_z(g) + \left(x + \frac{y}{4} + z\right)O_2(g) \longrightarrow xCO_2(g) + \frac{y}{2}H_2O(g) + zSO_2(g)$$

50. The reaction is

$$\begin{array}{cccc} 2CO(g) \ + \ O_2(g) & \longrightarrow & 2CO_2(g) \\ 2 \ Vol & 1 \ Vol & 2 \ Vol \\ 40.0 \ mL & 20.0 \ mL & 40.0 \ mL \end{array}$$

Volume of CO(g) that reacts = 40.0 mL

Volume of $CO_2(g)$ formed = 40.0 mL

Volume of CO(g) removed after treatment with KOH

= 50 mL - 40 mL = 10.0 mL

The Liquid State

• Type 1 •

Choose the correct option. Only one option is correct.

- 1. Calculate the radius of a capillary tube if water rises to a height of 12.5 cm within it, assuming the angle of contact between water and glass to be 0° ($g = 10 \text{ m s}^{-2}$, $\rho = 1 \times 10^{3} \text{ kg m}^{-3}$).
 - (a) 0.22 mm
- (b) 0.18 mm
- (c) 0.11 mm
- (d) 0.44 mm
- 2. Which of the following statements is correct?
 - (a) Surface tension is the energy per unit area of a free liquid surface.
 - (b) The unit of surface tension is N m⁻² and its dimensions are $ML^{\circ}T^{-1}$.
 - (c) If a soap bubble is filled with more air, the pressure inside it increases.
 - (d) The angle of contact between a solid and a liquid is a property of the shape of the solid.
- 3. When a glass plate is immersed in mercury, the surface near the plate
 - (a) becomes concave
 - (b) is depressed
 - (c) is flat
 - (d) is first concave and then depressed
- **4.** The dimensions $ML^{\circ}T^{-2}$ correspond to
 - (a) coefficient of viscosity
- (b) moment of inertia
- (c) surface tension
- (d) surface area
- 5. There is a depression in the surface of the liquid in a capillary when
 - (a) the cohesive force is smaller than the adhesive force
 - (b) the cohesive force is greater than the adhesive force

- (c) there is an equilibrium between the cohesive and the adhesive force
- (d) none of these is true
- **6.** A soap bubble of radius *r* is filled with air until the radius is doubled. If the surface tension of the soap solution is *S*, the work done in the process is
 - (a) $16\pi r^2 S$
- (b) $4\pi r^2 S$
- (c) $24\pi r^2 S$ (d) $12\pi r^2 S$
- 7. Which of the following correctly represents the relation between capillary rise h and capillary radius r?

(a)





(c)



(d)



- 8. What is the excess pressure inside a spherical soap bubble of radius 5 cm if the surface tension of the soap film is 3.5×10^{-2} N m⁻¹?
 - (a) 2.8 Pa

(b) 1.4 Pa

(c) 5.6 Pa

- (d) 11.2 Pa
- **9.** Which of the following will happen when two soap bubbles of radius r_1 and r_2 ($r_2 > r_1$) are connected by a piece of tubing?
 - (a) Both bubbles will collapse.
 - (b) The bubbles will change in size.
 - (c) The bubbles will not change in size.
 - (d) The angle of contact will decrease.
- **10.** When soap is added to water,
 - (a) the angle of contact will increase
 - (b) the surface tension of water will increase
 - (c) the angle of contact will decrease
 - (d) nothing will happen
- 11. The radius of a soap bubble is 5 cm. The surface tension of the soap film is 3.5×10^{-2} N m⁻¹. What is the work done in blowing the bubble?
 - (a) 4.4×10^{-2} J

(b) $1.1 \times 10^{-2} \text{ J}$

(c) 1.1×10^{-3} J

(d) $2.2 \times 10^{-2} \text{ J}$

12. Consider the relation

$$\ln \frac{p}{p_0} = \frac{2SM}{PRT} \times \frac{1}{r}$$

where p_0 and p are the vapour pressures over the liquid in bulk and over a curved surface respectively, the radius of curvature being r. Choose the correct option from among the following.

- (a) The smaller the droplet of a liquid, the higher is its vapour pressure.
- (b) The bigger the droplet of a liquid, the higher is its vapour pressure.
- (c) The smaller the droplet of a liquid, the lower is its vapour pressure.
- (d) None of these
- When water is in contact with a waxed glass surface, the angle of contact will
 - (a) be zero

- (b) be greater than 90°
- (c) be less than 90°
- (d) depend on the amount of wax
- 14. Surface tension does not vary with
 - (a) temperature

- (b) vapour pressure
- (c) the size of the surface
- (d) concentration
- **15.** The surface tension of CHCl $_3$ at 20°C is 27.4×10^{-3} N m $^{-1}$. The density of H $_2$ O at 20°C is 0.9982 kg m $^{-3}$ and that of CHCl $_3$ at the same temperature is 1.595 kg m $^{-3}$. If the rise of CHCl $_3$ and water in a capillary tube of a certain diameter is 2.33 cm and 9.9 cm respectively, calculate the surface tension of water.
 - (a) $82.76 \times 10^{-3} \text{ N m}^{-1}$
- (b) $72.86 \times 10^{-3} \text{ N m}^{-1}$
- (c) $78.68 \times 10^{-3} \text{ N m}^{-1}$
- (d) $68.72 \times 10^{-3} \text{ N m}^{-1}$
- 16. Among the following molecules, which has the least surface tension?
 - (a) Benzene

(b) Acetic acid

(c) Diethyl ether

- (d) Chlorobenzene
- **17.** The film of a soap bubble has a
 - (a) small thickness and there is air both inside and outside the film
 - (b) small thickness and there is no air inside and outside the film
 - (c) large thickness and there is air only inside the film
 - (d) small thickness and there is air only outside the film
- 18. The pressure inside a soap bubble is
 - (a) greater than that outside by an amount 2S/R (S =surface tension and R =radius of bubble)
 - (b) less than that outside by an amount 4S/R

19. For which of the following pairs is the angle of contact the maximum?

(c) Methylene iodide with glass (d) Mercury with glass **20.** A capillary tube of radius 0.20 mm is dipped vertically in water. Calculate the height of the water column raised in the capillary tube (surface tension of water = 0.075 N m⁻¹, density of water = 1.0×10^3

(b) Water with paraffin

(b) 17.5 cm(d) 5.5 cm

(c) greater than that outside by an amount 4*S/R*(d) less than that outside by an amount 2*S/R*

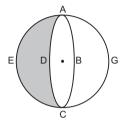
(a) Water with glass

kg m⁻³, g = 10 m s⁻²). (a) 10.0 cm

(c) 7.5 cm

21. A 5 cm-long needle is floating on water. What can be the maximum mass of the needle (surface tension of water = 0.075 N m^{-1} , $g = 10 \text{ m s}^{-2}$)? (b) 1.25 g (a) $0.75 \, g$ (c) 0.50 g(d) 1.00 g 22. When dipped in a liquid vertically, a capillary tube records a rise of 5 cm. When the tube is held in the liquid inclined at an angle of 45° to the vertical, the approximate length of the tube up to which the liquid rises is (a) 9.0 cm (b) 8.0 cm (c) 7.0 cm (d) 10.0 cm 23. The surface tension of a soap solution is 30×10^{-3} N m⁻¹. The work done in stretching a bubble of this solution of surface area $5 \text{ cm} \times 5 \text{ cm}$, to an area of $10 \text{ cm} \times 10 \text{ cm}$, is (b) $6.0 \times 10^{-4} \text{ J}$ (a) $4.5 \times 10^{-4} \text{ J}$ (c) $4.5 \times 10^{-5} \text{ J}$ (d) $7.5 \times 10^{-4} \text{ J}$ 24. Calculate the force exerted by water by across a line of length 10 cm drawn along the surface (surface tension of water = 0.075 N m^{-1}). (a) $7.5 \times 10^{-3} \text{ N}$ (b) $7.5 \times 10^{-4} \text{ N}$ (d) $7.5 \times 10^{-3} \text{ N}$ (c) $7.5 \times 10^{-2} \text{ N}$ **25.** Which of the following factors is responsible for surface tension? (a) Symmetrical force distribution in the liquid. (b) Asymmetrical force distribution in the liquid. (c) The symmetrical force distribution in solid. (d) All of these. 26. A small drop of liquid takes a nearly spherical shape, because (a) for a given volume, a sphere assumes the smallest surface area (b) for a given volume, a sphere assumes the smallest surface tension

- (c) the drop has a symmetrical distribution of force
- (d) the drop has the least adsorption power
- **27.** Imagine a diametric cross-section ABCD of the liquid drop in the figure, which divides it into two hemispheres. The surfaces of the two hemispheres touch each other along the periphery ABCD.



The forces acting on the surface are

- (a) F_1 due to the surface tension of the surface ABCDG in contact
- (b) F_2 due to the air outside the surface ABCDE
- (a) F_3 due to the liquid inside the surface ABCDE
- (d) all of these
- 28. For the figure associated with Q.27

(a)
$$F_1 + F_2 + F_3 = 0$$

(b)
$$F_1 + F_2 = F_3$$

(c)
$$F_1 - F_2 = F_3$$

(d)
$$F_1 = F_2 + F_3$$

- 29. Which of the following statements is correct?
 - (a) The formation of a water droplet is exothermic and the breaking up of one endothermic.
 - (b) The formation of a water droplet is endothermic and the breaking up of one exothermic.
 - (c) The formation as well as the breaking up of a water droplet are exothermic.
 - (d) None of these.
- 30. If two soap bubbles of different radii are connected by a tube,
 - (a) the angle of contact between the soap bubbles and water decreases
 - (b) the surface energy of the system becomes zero
 - (c) air flows from the smaller to the bigger bubble
 - (d) the total pressure of the system remains the same
- 31. Bernoulli's principle is a consequence of the law of conservation of
 - (a) angular momentum
- (b) mass

(c) energy

(d) momentum

- The Liquid State 1-95 32. The viscous force acting between two layers of a liquid may be worked out from the formula $\frac{F}{A} = -\eta \frac{dv}{dx}$, where the symbols have their usual meanings. F/A is called (a) tangential stress (b) tangential volume (c) density stress (d) bulk modulus 33. For laminar flow, all the particles of the liquid move parallel to the tube, and the velocity increases regulary from (a) zero at the wall to maximum at the wall
 - (b) zero at the wall to maximum at the centre
 - (c) $-\infty$ at the wall to ∞ at the centre
 - (d) ∞ at the wall to $-\infty$ at the centre
- 34. The relation betwen turbulent and laminar flow is provided by the Reynolds number, which is equal to
 - (a) $\frac{2Rv\rho}{n}$

(b) $2\eta \frac{Rv}{\rho}$

(c) $\frac{Rv\rho}{n}$

(d) $2\eta \frac{\rho}{r^{2}}$

where R = radius of capillary tube, v = velocity of liquid, $\rho = \text{density}$ of liquid, η = viscosity of liquid.

- 35. When a body falls through a viscous medium, its velocity
 - (a) first increases and then decreases
 - (b) first increases and then becomes constant, and this constant velocity is called critical velocity
 - (c) first increases and then becomes constant, and this velocity is called terminal velocity
 - (d) increases continuously
- **36.** A wooden plate of area 5 m² floating on the surface of a river is made to move horizontally with a speed of 1 m s⁻¹ by applying a tangential force. If the river is 1 m deep and the water in contact with the plate is stationary, calculate the tangential force (viscosity = 10^{-3} poise).
 - (a) $5.0 \times 10^{-2} \text{ N}$

(b) $5.0 \times 10^{-4} \text{ N}$

(c) $5.0 \times 10^{-6} \text{ N}$

- (d) $5.0 \times 10^{-3} \text{ N}$
- 37. The SI unit and dimensions of the coefficient of viscosity are respectively
 - (a) $Ns^{-1}m^{-1}$ and $ML^{-2}T^{-2}$ (b) Nsm^{-2} and $ML^{-1}T^{-1}$
 - (c) $Ns^{-2}m^{-2}$ and $ML^{-1}T^{-1}$ (d) $Ns^{-1}m^{-2}$ and $ML^{-2}T^{-1}$
- 38. Which of the following molecules has the highest viscosity?

(a) Acetone

(b) Benzene

(c) Ethanol

- (d) Carbon tetrachloride
- **39.** Which of the following statements is correct?
 - (a) The viscosities of most liquids decrease with increasing temperature, and those of gases increase with temperature.
 - (b) The viscosities of most liquids increase with temperature, and those of gases decrease with increasing temperature.
 - (c) The viscosities of liquids as well as gases decrease with increase in temperature.
 - (d) The viscosities of liquids as well as gases increase with increase in temperature.
- 40. The reciprocal of viscosity is called
 - (a) resistance

(b) Reynolds number

(c) fluidity

- (d) surface tension
- **41.** A certain volume of heptane (density = 0.7 g cm^{-3}) flows through a viscometer in 60 s at 20°C while the same volume of water requires 100 s at the same temperature. Calculate the absolute viscosity of heptane if that of water is 1.0×10^{-2} poise.
 - (a) 5.0×10^{-3} poise
- (b) 4.2×10^{-3} poise
- (c) 6.5×10^{-3} poise
- (d) 1.2×10^{-3} poise
- 42. The viscosity of a liquid is related to the temperature by the relation
 - (a) $\eta = \frac{a}{T} + b$

- (b) $\eta = a + bT + cT^2$
- (c) $\log \eta = \frac{a}{T} + bT^2$
- (d) $\ln \eta = \frac{a}{RT} + b$
- **43.** At a high temperature, a mesomorphic milky fluid changes sharply into a clear ordinary liquid. Such a substance is known as a
 - (a) paracrystalline substance
- (b) liquid crystal
- (c) transition liquid
- (d) colloid
- **44.** Under the influence of an electric field, liquid crystals exhibit an optical phenomenon called dynamic scattering, in which
 - (a) transparent liquid crystals become opaque
 - (b) transparent liquid crystals become solid crystals
 - (c) liquid crystals become colloids
 - (d) liquid crystals become superconductors

• *Type 2* •

Choose the correct options. More than one option is correct.

- **45.** Which of the following statements is correct?
 - (a) The angle of contact between water and glass is 0° .
 - (b) The angle of contact betwen mercury and glass is 100°.
 - (c) The angle of contact between mercury and glass is 140°.
 - (d) A liquid in equilibrium cannot sustain tangential stress.
- **46.** When a capillary tube is dipped into a liquid, the liquid neither rises nor falls in the capillary. Choose the correct options.
 - (a) The surface tension of the liquid must be zero.
 - (b) The angle of contact must be 90°.
 - (c) The surface tension may be zero.
 - (d) The angle of contact may be 90°.
- 47. Viscosity is a property of
 - (a) liquids

(b) gases

(c) solids

- (d) all of these
- 48. The viscosity of a liquid molecule depends on
 - (a) the volume of the liquid
- (b) the temperature of the liquid
- (c) the surface area of the liquid (d) the structure of the molecule
- 49. Thermotropic liquid crystals may be

47. a, b

(a) sematic

(b) nematic

(c) chlolestic

46. c, d

(d) lyotropic

49. a, b, c

Answers

1. c	2. a	3. b	4. c	5. b
6. c	7. b	8. a	9. b	10. c
11. d	12. a	13. b	14. c	15. b
16. c	17. a	18. c	19. d	20. c
21. a	22. c	23. a	24. d	25. b
26. a	27. d	28. b	29. b	30. c
31. c	32. a	33. b	34. a	35. c
36. d	37. b	38. c	39. a	40. c
41. b	42. d	43. b	44. a	45. a, c, d

48. b, d

Hints to More Difficult Problems

1. $2\pi r \cos \theta S = \pi r^2 \rho g h$

or
$$r = \frac{2S}{h\rho g}\cos\theta = \frac{2 \times 0.075 \text{ N m}^{-1} \times 1}{12.5 \times 10^{-2} \text{ m} \times 1.0 \times 10^{3} \text{ kg m}^{-3} \times 10.0 \text{ m s}^{-2}}$$

 \therefore r = 0.11 mm.

7. From the relation $r = \frac{2T}{h\rho g}\cos\theta$, it is clear that $r \propto \frac{1}{h}$. So the plot is hyperbolic.

8.
$$p = \frac{4S}{r} = \frac{4 \times 3.5 \times 10^{-2} \text{ N m}^{-1}}{5 \times 10^{-2} \text{ m}} = 2.8 \text{ N m}^{-2} = 2.8 \text{ Pa}.$$

- 11. Following Q. 8, $p = 2.8 \text{ N m}^{-2}$. Work done = $p \times \pi r^2 = 2.8 \text{ N m}^{-2} \times \frac{22}{7} (0.05 \text{ m})^2$ = $2.2 \times 10^{-2} \text{ J}$.
- **15.** Using the equation $\frac{S_1}{S_2} = \frac{d_1 l_2}{d_2 l_1}$,

$$S_1 = \frac{(27.4 \times 10^{-3} \text{ N m}^{-1}) (0.9982 \text{ kg m}^{-3}) (9.9 \times 10^{-2} \text{m})}{(1.595 \text{ kg m}^{-3}) (2.33 \times 10^{-2} \text{ m})}$$
$$= 72.86 \times 10^{-3} \text{ N m}^{-1}.$$

- 20. $h = \frac{2S \cos \theta}{r \rho g} = \frac{2 \times 7.5 \times 10^{-2} \text{ N m}^{-1} \times 1}{(0.20 \times 10^{-3} \text{ m})(10^{3} \text{ kg m}^{-3})(10 \text{ m s}^{-2})}$ = 7.5 cm.
- **21.** $S \times 2l = mg$

or
$$m = \frac{2Sl}{g} = \frac{2 \times 7.5 \times 10^{-2} \times 5.0 \times 10^{-2}}{10}$$

= 0.75 g.

- 23. Work done = $2(A_2 A_1) \times S$ = $2(10 \times 10 - 5 \times 5) \times 10^{-4} \text{ m}^2 \times 30 \times 10^{-3} \text{ N m}^{-1}$ = $4.5 \times 10^{-4} \text{ J}$.
- 27. Each hemispherical surface pulls the other due to surface tension.
- **30.** Excess pressure = $\frac{4S}{r}$ or pressure $\propto \frac{1}{r}$. Therefore, air flows from the smaller to the bigger bubble.
- **35.** The viscous force and the buoyant force together balance the weight, and thus the body moves with constant velocity.

- **36.** $F = -\eta A \frac{dv}{dx} = 10^{-3} \text{ N s m}^{-2} \times 5 \text{ m}^2 \times \text{m s}^{-1} \text{ m}^{-1} \text{ (1 poise} = \text{N s m}^{-2}\text{)}$ = 5.0 × 10⁻³ N.
- 39. For gases, viscosity $\eta = \sqrt{\frac{RTM}{\pi}} \frac{1}{N_A \pi \sigma^2} (\sigma = \text{molecular diameter})$. $\therefore \quad \eta \propto \sqrt{T}$.

The viscosity of gases increases with temperature.

Oxidation-Reduction

• *Type 1* •

1. Among the following molecules, in which does bromine show the

 ${\it Choose the correct option. Only one option is correct.}$

	maxin	num oxidatio	n num	ber?				
	(a)	$Hg_2(BrO_3)_2$			(b)	Br-Cl		
	(c)	$KBrO_4$			(d)	Br_2		
2.	A = +2 $compo$ (a)	2, that of B ound is ABC ₂			(b)	-2 . A possil $A_2(BC_3)_2$		on number of ormula of the
	(c)	$A_3(BC_4)_2$			(a)	$A_3(B_4C)_2$		
3.	The oxidation number of nitrogen varies from							
	(a)	-3 to +5			(b)	-1 to +1		
	(c)	-3 to -5			(d)	-5 to $+1$		
4.		n of the folloer in their co			do n	ot exhibit a	posi	itive oxidation
	(a)	I	(b) E	Br	(c)	Cl	(d)	F
5.	Iron s	hows an oxid	lation 1	number of	+1 in			
	(a)	$[Fe(H_2O)_6]^2$	+		(b)	Fe ₄ [Fe(CN) ₆	$]_3$	
	(c)	$[Fe(H_2O)_5N]$	O]SO ₄		(d)	$[\mathrm{FeBr}_4]^-$		
6.	Phosp	horus shows	an ox	idation nui	nber	of +3 in		
	(a)	orthophosp	horic a	cid	(b)	orthophosp	horo	us acid
	(c)	metaphosph	noric a	cid	(d)	pyrophosph	oric	acid
	1-100							

(a) +3

(c) -3

number of nitrogen in compound X?

7. A metal ion M^{3+} loses three electrons to produce another cation. The oxidation number of the metal in the cation is

8. A sample of 2.5 mol of hydrazine (N₂H₄) loses 25 mol of electrons on being converted to a new compound X. Assuming that there is no loss of nitrogen in the formation of the new compound, what is the oxidation

(b) +4

(d) +6

	(a)	-1	(b)	-2
	(c)	+3	(d)	+4
9.		nich of the following has the ged in increasing order?	oxic	dation number of oxygen been
	(a)	$BaO_{2} < KO_{2} < O_{3} < OF_{2}$	(b)	$OF_2 < KO_2 < BaO_2 < O_3$
	(c)	$BaO_2 < O_3 < OF_2 < KO_2$	(d)	$KO_2 < OF_2 < O_3 < BaO_2$
10.	Which	n of the following agents is the	mos	t oxidizing?
	(a)	O_3	(b)	$KMnO_4$
	(c)	H_2O_2	(d)	$K_2Cr_2O_7$
11.	Which	n of the following agents is the	mos	t reducing?
	(a)	Mg	(b)	Na
	(c)	K	(d)	Br_2
12.	Which	n of the following agents is the	mos	t reducing?
	(a)	HNO ₂	(b)	H_2S
	(c)	H ₂ SO ₃	(d)	SnCl ₂
13.	In the	reaction $3Br_2 + 6CO_3^{2-} + 3H_2O$	\rightarrow 51	$Br^- + BrO_3^- + 6HCO_3^-$
	(a)	bromine is oxidized and the c	arbo	nate radical is reduced
	(b)	bromine is reduced and the ca	arboı	nate radical is oxidized
	(c)	bromine is neither reduced no	or ox	idized
	(d)	bromine is both reduced and	oxid:	ized
14.			ns w	vill there be no change in the
		tion number of nitrogen? $HNO_3 + 2H_2SO_4 \rightarrow NO_2^+ + H_3$	O+ +	JUCO-
		$2N_2O_4 + 2KI \rightarrow 2KNO_3 + 2NC$		-
		$2KHN_2 + N_2O \rightarrow KN_3 + KOH$	_	
		$6K_3[Fe(CN)_6] + Cr_2O_3 + 10KO_3$		*
	(a)	$6K_4[Fe(CN)_6] + 2K_2CrO_4 + 5H$		
		16 (/03 / 2 - 4	_	

reducing power?

acts as

(a) Zn < Al < Mg < Na

(a) an oxidizing agent(b) a reducing agent(c) a catalyst

(c) Na < Mg < Al < Zn

15. Which of the following shows agents arranged in order of increasing

16. In the reaction $2Ag + 2H_2SO_4 \rightarrow Ag_2SO_4 + 2H_2O + SO_2$, sulphuric acid

(b) Al < Mg < Zn < Na

(d) Mg < Na < Zn < Al

	(d)	an acid as well as an oxidant		
17.		nich of the following coordin s have an oxidation number of		n compounds do the transition
	(a)	$[Cr(H_2O)_4Cl_2]Cl.2H_2O$	(b)	$[Fe(CO)_5]$
	(c)	$[(H_2O)_5Cr -\!$	(d)	$K_2[Cr(CN)_2O_2(O_2)NH_3]$
18.	oxida (a) (b) (c)	h of the following has been tion number of nitrogen? $NH_3 < N_2O_5 < NO < N_2 \\ NO_2^+ < NO_3^- < NO_2^- < N_3^- \\ NH_4^+ < N_2H_4 < NH_2OH < N_2O_1^+ < NO_2^- < NaN_3^- < NO_2^+ < N_2O_2^+ < NaN_3^+ < Na$		ranged in order of increasing
19.		al and a strong alkaline medi		an acidic, a slightly alkaline or are respectively $(M = molecular)$
	(a)	M/5, M/2, M	(b)	M/5, M/3, M/2
	(c)	M/5, M/3, M	(d)	M/3, M, M/5
20.	The e	quivalent weight of $Cr_2O_7^{2-}$ in a	an ac	ridic medium is
	(a)	M/2	(b)	M/3
	(c)	M/9	(d)	M/6
21.		quivalent weight of Na ₂ S ₂ O ₃ in $2Na_2S_2O_3 + I_2 \rightarrow Na_2S$		
	is	λ.((1.)	M /0
	(a)		` ′	M/8
	(c)	M/0.5	(d)	<i>M</i> /2
22.	In the	conversion $NH_2OH \rightarrow N_2O$, the	ne eg	uivalent weight of NH ₂ OH is
	(a)	M/4	(b)	M/2
	(c)	M/5	(d)	M

23. What will be the value of the equivalent weight of KBrO₃ in the following ionic equation?

$$2BrO_3^- + 12H^+ + 10e \rightarrow Br_2 + 6H_2O$$

(a) M/4

(b) M/6

(c) M/10

(d) M/5

24. In the redox reaction

$$xMnO + yPbO_2 + zHNO_3 \rightarrow HMnO_4 + Pb(NO_3)_2 + H_2O$$

- (a) x = 2, y = 5, z = 10
- (b) x = 2, y = 7, z = 8
- (c) x = 2, y = 5, z = 8
- (d) x = 2, y = 5, z = 5

25. In the redox reaction

$$xKMnO_4 + yNH_3 \rightarrow KNO_3 + MnO_2 + KOH + H_2O$$

(a) x = 4, y = 6

(b) x = 8, y = 3

(c) x = 8, y = 6

(d) x = 3, y = 8

26. In the redox reaction

$$x$$
CrCl₃ + y H₂O₂ + z NaOH \rightarrow Na₂CrO₄ + NaCl + H₂O

- (a) x = 2, y = 4, z = 10
- (b) x = 2, y = 6, z = 5
- (c) x = 2, y = 3, z = 8
- (d) x = 2, y = 3, z = 10

27. In the ionic reaction

$$x Br O_3^- + y Cr^{3+} + z H_2 O \rightarrow Br_2 + H Cr O_4^- + H^+$$

- (a) x = 6, y = 10, z = 20
- (b) x = 6, y = 10, z = 11
- (c) x = 6, y = 10, z = 22
- (d) x = 6, y = 8, z = 22

28. In the ionic equation

$$xCH_3CH_2OH + yI_2 + zOH^- \rightarrow CHI_3 + HCO_2^- + I^- + H_2O$$

- (a) x = 1, y = 4, z = 6
- (b) x = 1, y = 6, z = 4
- (c) x = 1, y = 8, z = 12
- (d) x = 1, y = 8, z = 8
- 29. In the ethylene molecule the two carbon atoms have the oxidation numbers
 - (a) -1, -1

(b) -2, -2

(c) -1, -2

- (d) +2, -2
- **30.** The oxidation numbers of C-1 and C-2 in propene ($CH_3CH=CH_2$) are respectively
 - (a) -1, -2

(b) -1, -1

(c) -2, -1

- (d) +2,-1
- **31.** The oxidation number of carbon in HCOOH is
 - (a) +2
- (b) -2
- (c) +4
- (d) zero

32. The oxidation number of the carbox	ylic carbon atom in	CH ₃ COOH is
---	---------------------	-------------------------

(a) +2

(b) +4

(c) +1

(d) +3

33. When methane is burnt in oxygen to produce CO₂ and H₂O, the oxidation number of carbon changes by

(a) -8

(b) zero

(c) +8

(d) +4

• Type 2 •

Choose the correct options. More than one option is correct.

- **34.** The oxidation number of Cr = +6 in
 - (a) FeCr₂O₄

(b) KCrO₃Cl

(c) CrO₅

- (d) [Cr(OH)₄]⁻
- 35. The oxidation number of carbon is zero in
 - (a) HCHO

(b) CH₂Cl₂

(c) $C_6H_{12}O_6$

- (d) $C_{12}H_{22}O_{11}$
- **36.** Which of the following are not redox reactions?
 - (a) $Mg + N_2 \rightarrow Mg_3N_2$
 - (b) $K_4[Fe(CN)_6] + H_2SO_4 + H_2O \rightarrow K_2SO_4 + CO + FeSO_4 + (NH_4)_2SO_4$
 - (c) $I_2 + 3Cl_2 \rightarrow ICl_3$
 - (d) $CuSO_4 + NH_3 \rightarrow [Cu(NH_3)_4]SO_4$
- 37. Which of the following are redox reactions?
 - (a) $NaIO_3 + NaHSO_3 \rightarrow NaHSO_4 + Na_2SO_4 + I_2 + H_2O$
 - (b) $FeCl_3 + K_4[Fe(CN)_6] \rightarrow KCl + Fe_4[(Fe(CN)_6]_3]$
 - (c) $AgCl + Na_2S_2O_3 \rightarrow Na_3[Ag(S_2O_3)_2] + NaCl$
 - (d) NaBiO₃ + MnSO₄ + HNO₃ \rightarrow $HMnO_4 + Bi(NO_3)_3 + NaNO_3 + Na_2SO_4 + H_2O$
- 38. Which among the following are examples of autoredox reactions?
 - (a) $P_4 + OH^- \rightarrow H_2PO_4^- + PH_3$ (b) $S_2O_3^{2-} \rightarrow SO_4^{2-} + S$
 - (c) $H_2O_2 \rightarrow H_2O + O_2$
- (d) $AgCl + NH_3 \rightarrow [Ag(NH_3)_2]Cl$

- **39.** The oxidation number of S = +6 in
 - (a) peroxomonosulphuric acid (Caro's acid)
 - (b) peroxodisulphuric acid (Marshall's acid)
 - (c) pyrosulphuric acid (oleum)
 - (d) sodium thiosulphate (hypo)
- **40.** Which of the following have been arranged in order of decreasing oxidation number of sulphur?
 - (a) $H_2S_2O_7 > Na_2S_4O_6 > Na_2S_2O_3 > S_8$
 - (b) $SO^{2+} > SO_4^{2-} > SO_3^{2-} > HSO_4^{-}$
 - (c) $H_2SO_5 > H_2SO_3 > SCl_2 > H_2S$
 - (d) $H_2SO_4 > SO_2 > H_2S > H_2S_2O_8$

Answers

1. c	2. c	3. a	4. d	5. c
6. b	7. d	8. c	9. a	10. a
11. c	12. b	13. d	14. a	15. a
16. d	17. d	18. c	19. c	20. d
21. a	22. b	23. d	24. a	25. b
26. d	27. c	28. a	29. b	30. c
31. a	32. d	33. c	34. b, c	35. a, b, c, d
36. b, d	37. a, d	38. a, b, c	39. a, b, c	40. a, c

Hints to More Difficult Problems

- 9. $BaO_2 < KO_2 < O_3 < OF_2$ -1 -0.5 0 +2
- 10. Among these, O_3 has the greatest value of standard redox potential $E^{\circ}(=+2.07 \text{ V})$.
- 13. Br_2 is reduced to Br^- (oxidation number decreases from zero to -1) and Br_2 is oxidized to BrO_3^- (oxidation number increases from zero to +5).
- **16.** H₂SO₄ is a diprotic acid. It is also an oxidant because the oxidation number of S in the given reaction decreases from 6 to 4.
- 18. Oxidation number $\frac{\underline{N}H_4^+ < \underline{N}_2H_4 < \underline{N}H_2OH < \underline{N}_2O}{-1} < \frac{\underline{N}}{+1} = 0$

19.
$$KMnO_4 \xrightarrow{\text{Acidic}} Mn^{2+} E = \frac{M}{7-2} = \frac{M}{5}$$
 $KMnO_4 \xrightarrow{\text{medium}} MnO_4 \xrightarrow{\text{slightly alkaline}} MnO_2 E = \frac{M}{7-4} = \frac{M}{3}$
 $KMnO_4 \xrightarrow{\text{strongly alkaline}} MnO_4^{-2-} E = \frac{M}{7-6} = M$

20.
$$Cr_2O_7^{2-} + 14H^+ + 6e \longrightarrow 2Cr^{+3} + 7H_2O$$

 $E_{K_2Cr_2O_7} = \frac{M}{6}$ (no. of electrons involved = 6)

21.
$$2S_2O_3^{2-} \longrightarrow S_4O_6^{2-} + 2e$$

 $E_{Na_2S_2O_3} = \frac{2M}{2} = M$

22. The oxidation numbers of N in NH_2OH and N_2O are -1 and 1 respectively.

$$2N = -2$$

$$2NH_2OH \longrightarrow N_2O$$

$$E = \frac{Mol. \text{ wt.}}{\Delta 0.N} = \frac{2M}{2 - (-2)} = \frac{2M}{4} = \frac{M}{2}$$

23.
$$2BrO_3^- + 12H^+ + 10e \rightarrow Br_2 + 6H_2O$$

$$E_{KBrO_3} = \frac{\text{mol. wt.}}{\text{no. of electrons gained}} = \frac{2BrO_3^-}{10}$$
$$= \frac{BrO_3^-}{5} = \frac{KBrO_3}{5} = \frac{M}{5}$$

where M stands for mol. wt. of KBrO₃

31. The oxidation no. of HCOOH = 0 or $1 + x - 4 + 1 = 0 \Rightarrow x = +2$, where x is the oxidation number of carbon in HCOOH.

33.
$$CH_4(g) + 2O_2(g) \rightarrow CO_2(g) + 2H_2O(1)$$

The oxidation number of C changes by $4 - (-4) = +8$

34. The total oxidation number of KCrO₃Cl = $0 \Rightarrow 1 + x - 6 - 1 = 0 \Rightarrow x = +6$, where x is the oxidation number of Cr.

The oxidation number of

Oxidation number of Cr = +6.

- **36.** There is no change in the oxidation number of the element concerned during the course of the reaction.
- **38.** In an autoredox reaction, an element or compound is oxidized as well as reduced.

39.
$$H \longrightarrow O \longrightarrow S O_3 H$$
, $S O_3 H \longrightarrow O \longrightarrow S O_3 H$, $x = +6$ $x = +6$

Modern Concepts of Acids and Bases

• Type 1 •

Ch

00	se the o	correct option	n. O1	ıly one	option is	correct.		
1.		ng the follow tor character?		Lewis a	acids, whi	ch has the	maximum	electror
	(a)	PCl ₃			(b)	PCl ₂ R		
	(c)	PF ₃			(d)	PClR ₂		
2.	(a) (b) (c)	n of the follow $C_6H_5CO_2H$: H_3O^+ and $C_6H_3NH_3^+$ and H_3O^- and H_3O^- and H_3O^-	and ()H ⁻ ıd CI	C ₆ H ₅ CC		se conjugate	pair?	
3.	Which	n of the follow	ving	species	s cannot ex	ist in an aqı	ueous solut	ion?
	(a)	NO_3^-		_	(b)	NH ₄ ⁺		
	(c)	NH_2^-			(d)	NH_3		
4.	Amor	ng the followi	ng, t	he stro	ngest conj	ugate base i	S	
	(a)	NO_3^-			(b)	Cl-		
	(c)	SO ₄ ²⁻			(d)	CH ₃ COO	=	
5.	The co	onjugate acid	of th	ne HPC	0_3^{2-} ion is			
	(a)	PO ₃ ³⁻			(b)	$H_2PO_3^-$		
	(c)	$H_2PO_4^-$			(d)	HPO_4^{2-}		
6.	Of the	e given anion	s, the	strong	gest Bronst	ed base is		
		ClO-				ClO ₃	(d) ClO	$_4^-$
	1-100							

7.		CH ₂ COOH may behave			
		only as an acid			
		only as a base			
		as an acid as well as a base neither as an acid nor as a bas	:e		
8.	` /	n of the following species cannot		a Bronsted b	pase?
•		O^{2-}		CH ₃ ⁺	
	` '	CH₄	. ,	PH ₃	
9.		n of the following can act as a B		o de la companya de l	not as a Lewis acid?
		OH-		AlCl ₃	
	` '	FeCl ₃	` /	NH ₃	
10.	. ,	formation of Al ₂ Cl ₆ , each AlC	. ,	9	
		hydrolyses	13 111		
		acts as a Bronsted acid			
	(c)	behaves both as a Bronsted ac	cid a	nd as a Brons	ted base
	(d)	behaves both as a Lewis acid	and	as a Lewis ba	ise
11.	Amor	ng the following, which is the s	tron	gest protonic	acid?
	(a)	$[Fe(H_2O)_6]^{3+}$	(b)	$[Fe(H_2O)_6]^{2}$	+
	(c)	H_2O	(d)	BF_3	
12.	Amor	ng the following, which is the s	tron	gest Lewis ac	id?
	(a)	BF_3	(b)	BCl ₃	
	(c)	BBr_3	(d)	BI_3	
13.	Amor	ng the following, which is the n	nost	basic toward	BMe ₃ ?
	(a)	Me_3N (b) Et_3N	(c)	O_2	(d) SO ₃
14.	Amor	ng the following, which is the s	tron	gest acid?	
	(a)	ClO ₃ (OH)	(b)	ClO ₂ (OH)	
	(c)	SO(OH) ₂	(d)	$SO_2(OH)_2$	
15.	Amor	ng the following, which has the	higl	nest dielectric	constant?
	(a)	H_2O		H_2SO_4	
	(c)	CH ₃ CH ₂ OH	(d)	CH ₃ COCH ₃	
16.		n of the following has been arra strength?	nge	d correctly in	order of increasing
	(a)	$CH_3CO_2H < O_2NCH_2CO_2H <$	NC	$CH_2CO_2H < H$	HOCH ₂ CO ₂ H

 $(b) \quad CH_3CO_2H < HOCH_2CO_2H < NCCH_2CO_2H < O_2NCH_2CO_2H \\$

- (c) HOCH₂CO₂H < NCCH₂CO₂H < CH₃CO₂H < O₂NCH₂CO₂H
- (d) $O_2NCH_2CO_2H < NCCH_2CO_2H < HOCH_2CO_2H < CH_3CO_2H$
- 17. Which of the following has been arranged correctly in order of increasing acidic strength?
 - (a) $CH_3CO_2H < HCO_2H < HO_2CCO_2H < CH_3CH_2CO_2H$
 - (b) CH₃CH₂CO₂H < CH₃CO₂H < HO₂CCO₂H < HCO₂H
 - (c) $HO_2CCO_2H < HCO_2H < CH_3CO_2H < CH_3CH_2CO_2H$
 - (d) $CH_3CH_2CO_2H < CH_3CO_2H < HCO_2H < HO_2CCO_2H$
- 18. Among the following, which is the strongest acid?
 - (a) CH₃CHOH | F

(b) CH₂CH₂OH

(c) CH₂CH₂OH Cl

- (d) CH₃CH₂OH
- **19.** Which of the following has been arranged correctly in order of increasing basic strength?
 - (a) $OH^- < HC \equiv C^- < NH_2^- < CH_3CH_2^-$
 - (b) $HC \equiv C^{-} < CH_{3}CH_{2}^{-} < NH_{2}^{-} < OH^{-}$
 - (c) $OH^- < NH_2^- < HC = C^- < CH_3CH_2^-$
 - (d) $NH_2^- < HC = C^- < OH^- < CH_3CH_2^-$
- **20.** Which of the following has been arranged correctly in order of increasing basic strength?
 - (a) $NH_3 < p-O_2NC_6H_4NH_2 < o-O_2NC_6H_4NH_2 < C_6H_5NH_2$
 - (b) $C_6H_5NH_2 < NH_3 < o-O_2NC_6H_4NH_2 < p-O_2NC_6H_4NH_2$
 - (c) $o-O_2NC_6H_4NH_2 < p-O_2NC_6H_4NH_2 < C_6H_5NH_2 < NH_3$
 - (d) $NH_3 < C_6H_5NH_2 < p-O_2NC_6H_4NH_2 < o-O_2NC_6H_4NH_2$
- 21. In a particular Lewis acid-base reaction, boric acid
 - (a) accepts a pair of electrons from the oxide ion present in a water molecule
 - (b) donates a pair of electrons to the hydrogen ion present in a water molecule
 - accepts a pair of electrons from the hydroxide ion present in a water molecule
 - (d) donates a pair of electrons to the hydroxide ion present in a water molecule

22.		n negative radical of the follow ous solution independently?	ing	compounds does not exist in an
	(a)	NaCl	(b)	NaNH ₂
	(c)	K ₂ SO ₄	(d)	CaCl ₂
23.		n of the following pairs representation of the following pairs representation of the following pairs are seen to be seen as the following pairs are seen as the following pairs represent	ents	the strongest acid and strongest
	(a)	H_3O^+ and OH^-	(b)	NH ₄ and NH ₂
	(c)	ClO_4^- and H_2F^+	(d)	NO_3^- and $HC \equiv C^-$
24.	Which water		n ord	der of increasing acid strength in
	(a)	$[Al(OH)_6]^{3+} < [Fe(H_2O)_6]^{3+} < [Fe(H_$	Fe(F	$(H_2O)_6]^{2+}$
	(b)	$[Fe(H_2O)_6]^{3+} < [Fe(H_2O)_6]^{2+} <$	[Al($H_2O)_6]^{3+}$
	(c)	$[Fe(H_2O)_6]^{2+} < [Fe(H_2O)_6]^{3+} <$	[Al($H_2O)_6]^{3+}$
	(d)	$[Fe(H_2O)_6]^{2+} < [Al(H_2O)_6]^{3+} <$	[Fe($(H_2O)_6]^{3+}$
25.	Which	n of the following is not a Lewi	s aci	d-base reaction?
	(a)	$BrF_3 + F^- \rightarrow [BrF_4]^-$	(b)	$AlCl_3 + BF_3 \rightarrow AlCl_3.BF_3$
	(c)	$I_2 + I^- \longrightarrow I_3^-$	(d)	$KH + H_2O \rightarrow KOH + H_2$
26.	Which	n of the following is arranged in	n ord	der of increasing proton affinity?
	(a)	$HS^- < I^- < NH_2^- < F^-$	(b)	$F^- < I^- < NH_2^- < HS^-$
	(c)	$I^- < F^- < HS^- < NH_2^-$	(d)	$NH_{2}^{-} < HS^{-} < F^{-} < I^{-}$
27.	Which	n of the following is the most st	rong	gly basic in water?
	(a)	CO ₃ ²⁻	(b)	ClO ₄
	(c)	NO_3^-	(d)	O ²⁻
28.	Which	n of the following is the weakes	st aci	id in H ₂ SO ₄ ?
	(a)	ClO_4^- (b) NO_3^-	(c)	HSO_4^- (d) CN^-
29.	Amor	ng the following, which is the s	trong	gest Lewis acid?
	(a)	$B(t-Bu)_3$	(b)	$BeCl_2$
	(c)	$B(n-Bu)_3$	(d)	BF ₃
30.	Which	n of the following is the most b	asic	towards B(CH ₃) ₃
	(a)	4 -CH $_3$ C—H $_4$ N	(b)	$(C_2H_5)_3N$
	(c)	NH ₂	(d)	2-CH ₂ C _E H ₄ N

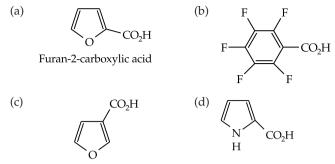
- **31.** The enthalpies of reaction of trimethylboron with NH₃, CH₃NH₂, (CH₃)₂NH and (CH₃)₃N are –58, –74, –81 and –74 kJ mol ⁻¹ respectively. What is the reason for trimethylamine being out of line?
 - (a) Resonance stabilization
 - (b) Steric repulsion
 - (c) Solvation factor
 - (d) Lattice energy considerations
- 32. Which of the following is arranged in order of increasing acidity?
 - (a) $[Mn(H_2O)_6]^{2+} < [Na(H_2O)_6]^{+} < [Sc(H_2O)_6]^{3+} < [Ni(H_2O)_6]^{2+}$
 - (b) $[Ni(H_2O)_6]^{2+} < [Sc(H_2O)_6]^{3+} < [Na(H_2O)_6]^{+} < [Mn(H_2O)_6]^{2+}$
 - (c) $[Na(H_2O)_6]^+ < [Mn(H_2O)_6]^{2+} < [Ni(H_2O)_6]^{2+} < [Sc(H_2O)_6]^{3+}$
 - $(d) \quad \left[Sc(H_2O)_6\right]^{3+} < \left[Ni(H_2O)_6\right]^{2+} < \left[Mn(H_2O)_6\right]^{2+} < \left[Na(H_2O)_6\right]^{4+}$
- 33. Among the following, which is the weakest Lewis base?
 - (a) CH₃

(b) NH_2^-

(c) OH-

- (d) F-
- 34. H_3PO_2 and $H_4P_2O_7$ are respectively
 - (a) tribasic and tetrabasic acids
 - (b) dibasic and tetrabasic acids
 - (c) monobasic and tetrabasic acids
 - (d) tribasic and dibasic acids
- **35.** Which of the following statements is incorrect?
 - (a) HNO₂ is a stronger acid than CH₃CO₂H.
 - (b) NO_2^- is a weaker base than CN^- .
 - (c) NH₂ does not exist in an aqueous solution.
 - (d) NH₂ ion exists in an aqueous solution.
- **36.** Which of the following represents an ordered arrangement of the most acidic through amphoteric to the most basic compounds?
 - (a) $BaO < Al_2O_3 < B_2O_3 < CO_2 < SO_3 < Cl_2O_7$
 - (b) $Cl_2O_7 < SO_3 < CO_2 < B_2O_3 < Al_2O_3 < BaO$
 - (c) $Al_2O_3 < B_2O_3 < CO_2 < Cl_2O_7 < SO_3 < BaO$
 - (d) $SO_3 < Cl_2O_7 < B_2O_3 < CO_2 < BaO < Al_2O_3$
- 37. Which of the following compounds is the most likely to be amphoteric?
 - (a) Li₂O
- (b) PbO₂
- (c) MgO
- (d) P_4O_{10}

38. Which of the following carboxylic acids is the least acidic?



Furan-3-carboxylic acid

- 39. Which of the carboxylic acids is the most acidic?
 - (a) Acetic acid

(b) Formic acid

(c) Benzoic acid

- (d) Cyclopentane carboxylic acid
- **40.** A transition element forms several binary oxides, and the oxide with the metal in the
 - (a) highest oxidation state is the most acidic
 - (b) lowest oxidation state is the most acidic
 - (c) highest oxidation state is amphoteric
 - (d) highest oxidation state is basic

• *Type* 2 •

Choose the correct options. More than one option is correct.

- 41. Which of the following cannot act as a Bronsted base?
 - (a) NH₃

(b) BF₃

(c) H_3O^+

(d) H₂O

- **42.** Which of the following compounds are not bases when dissolved in water?
 - (a) K₂O

(b) Na_2O_2

(c) NO_2

(d) Cl_2O_7

- **43.** Which of the following compounds/ions can act as a Bronsted acid as well as a Bronsted base?
 - (a) HCO₃

(b) K_2CO_3

(c) H_2SO_4

(d) $H_2PO_4^-$

- **44.** Which of the following acids are likely to form zwitterions?
 - (a) $H_2N \cdot CH \cdot CO_2H$ ĊН
 - (b) $HO_2C \cdot CH_2 \cdot CH \cdot CO_2 \cdot H$ NH_2
 - (c) $H_2N \cdot CH_2CH_2CH_2CH_2 \cdot CH \cdot CO_2H$ NH₂

(d) HO—CH
$$_2$$
 · CH · COOH NH $_2$

- 45. Which of the following species can be formed in acid-base reactions as Bronsted bases as well as Bronsted acids?
 - (a) $[Fe(H_2O)_6]^{3+}$ (b) HPO_4^{2-} (c) HSO_4^{-} (d) NH_4^{+}
- **46.** On the basis of Bronsted concepts, which of the following are correct?
 - (a) $CH_3CH_2OH + HCl \rightleftharpoons CH_3CH_2O^+H_2 + Cl^-$
 - (b) $CH_3CH_2OH + NH_2^- \rightleftharpoons CH_3CH_2O^+H_2 + NH^{2-}$
 - (c) $CH_3CO_2H + CH_3O^- \rightleftharpoons CH_3CO_2^- + CH_3OH$
 - (d) $CH_3CO_2H + CH_3NH_2 \rightleftharpoons CH_3COOH_2^+ + CH_3NH_2^-$
- 47. Which of the following statements are correct?
 - (a) HO is a stronger base than NH₂.
 - (b) H₂O is a stronger base than CH₃OH.
 - (c) CH₃COO is a stronger base than HCO₂H.
 - (d) A Lewis acid is a compound that can accept a pair of electrons.
- 48. Which of the following species can act as a Bronsted base as well as a Lewis base?
 - (a) NO_3^-
- (b) CN⁻
- (c) NH₃
- (d) BF₃
- 49. Which of the following are Lewis acids?
 - (a) $(CH_3)_3B$
- (b) : CH₂
- (c) S
- (d) CH₃⁺
- 50. On the basis of the Lewis concept, which of the following are correct?
 - (a) $Ag^+ + 2NH_3 \rightarrow [H_3N Ag NH_3]^+$
 - (b) $Br_2 + FeCl_3 \rightleftharpoons Br \stackrel{+}{Br} FeCl_3$

(c)
$$ROH + ZnCl_2 \rightleftharpoons R \stackrel{+}{\longrightarrow} CH$$

 ZnC

(d)
$$NH_4^+ + OH^- \rightleftharpoons NH_3 + H_2O$$

- **51.** Which of the following have been arranged correctly in order of increasing acidic strength?
 - (a) HF < HCl < HBr < HI
 - (b) $H_3PO_2 < H_3PO_3 < H_3PO_4$
 - (c) H₃PO₄ < H₂SO₄ < HClO₄
 - (d) $NH_3 < PH_3 < AsH_3 < BiH_3$
- **52.** Which of the following have been arranged correctly in order of increasing basic strength?
 - (a) LiOH < NaOH < KOH < RbOH
 - (b) $Al(OH)_3 < Mg(OH)_2 < NaOH$
 - (c) $NH_3 < NH_2OH < N_2H_4$
 - (d) $HF < H_2O < NH_3$
- **53.** Which of the following have been arranged correctly in order of increasing basic strength?
 - (a) $C_6H_5N(CH_3)_2 < C_6H_5NHCH_3 < C_6H_5NH_2$
 - (b) $NH_2^- < OH^- < F^-$
 - (c) $(CH_3)_3N < CH_3NH_2 < (CH_3)_2NH$
 - (d) $NH_2OH < N_2H_4 < NH_3$
- 54. Which of the following salts are acidic in aqueous solutions?
 - (a) $(NH_4)_2SO_4$

(b) NaHCO₃

(c) CuSO₄

- (d) AgNO₃
- 55. Which of the following salts are alkaline in aqueous solutions?
 - (a) NaCN

(b) $K_4[Fe(CN)_6]$

(c) CH₃CO₂Na

(d) FeCl₃

Answers

 1. c
 2. b
 3. d
 4. d
 5. b

 6. a
 7. c
 8. b
 9. d
 10. d

 11. a
 12. c
 13. a
 14. a
 15. b

16. b	17. d	18. a	19. a	20. c
21. c	22. b	23. a	24. c	25. b
26. c	27. d	28. a	29. c	30. a
31. b	32. c	33. d	34. c	35. d
36. b	37. b	38. d	39. b	40. a
41. b, c	42. c, d	43. a, d	44. a, b, c, d	45. b, c
46. a, c	47. b, c, d	48. a, b, c	49. a, b, c, d	50. a, b, c
51. a, c, d	52. a, b, d	53. a, c, d	54. a, c, d	55. a, c

Hints to More Difficult Problems

- Acetic acid is the weakest of these.
- 6. Among the oxyacids of chlorine, HClO is the weakest.
- During the formation of Al₂Cl₆, one AlCl₃ group is an electron-pair donor and the other is an electron-pair acceptor.
- 12. The strengths of boron halides as Lewis acids are in the order $BBr_3 > BCl_3 > BF_3$. This order is the opposite of what is expected on steric grounds as well as from considerations of electronegatively. It can be explained in terms of boron-halogen π -bonding.
- **18.** The compound listed first is the most acidic because the electron-withdrawing fluorine is the closest to the bond that holds the acidic hydrogen.
- **22.** The amide ion (NH₂⁻) is a stronger base than OH⁻ and therefore reacts completely with water as follows:

$$NH_2^-(aq) + H_2O(1) \longrightarrow NH_3(aq) + OH^-(aq)$$

For this reason the amide ion does not exist in aqueous solutions.

- 24. The weakest acid is the Fe $^{2+}$ complex because of its relatively large radius and low charge. The increase of charge to +3 increases the acid strength. The greater acidity of Al $^{3+}$ can be explained by its small radius.
- **25.** Both the reactants are electron-deficient.
- **29.** B(n-Bu) has no sterific effect.
- **30.** It is easier to remove a proton from 4-methylpyridine, and the electron left behind when the proton is removed is delocalized onto the electronegative nitrogen atom. In 4-methylpyridine, the electrons can be delocalized only onto the carbon atom. So nitrogen will be a good electron-pair donor for B(CH₃)₃.
- 32. Sc³⁺ has the smallest radius.

- 36. Cl_2O_7 forms a strong oxoacid and alkaline earth metals form strong bases with water.
- **42.** NO₂ and Cl₂O₇ form acids (HNO₂ and HClO₄) upon reacting with water.
- **45.** HPO_4^{2-} and HSO_4^{-} can donate a proton to form PO_4^{3-} and SO_4^{2-} respectively. They can also accept a proton to form $H_2PO_4^{-4}$ and H_2SO_4 respectively.

10

Volumetric Analysis

• Type 1 •

1. Calculate the equivalent weight of Na₂CO₃ when it is titrated against HCl

2. 5 mL of N/50 Ba(OH)₂ solution was poured into a 2.5-L flask containing some CO₂. The flask was then tightly corked. The CO₂ was completely absorbed by the Ba(OH)₂ solution. The percentage of CO₂ in the flask was

(b) 53

(d) 212

(b) 0.45

(d) 0.065

Choose the correct option. Only one option is correct.

in presence of phenolphthalein.

(a) 106(c) 26.5

(a) 0.045

(c) 4.5

3.	Calculate the equivalent weight of Na ₂ CO ₃ when it is titrated against HCl in the presence of methyl orange.			
	(a)	5.3	(b)	53
	(c)	10.6	(d)	106
4.		mole fraction of a solute in an a solution is	aque	ous solution is 0.50, the molality
	(a)	1.55	(b)	0.5
	(c)	55.55	(d)	5.55
5.	Carbo	onate-free sodium hydroxide is	prep	pared by using
	(a)	NaHCO ₃	(b)	$Ba(OH)_2$
	(c)	Na ₂ CO ₃	(d)	${\rm BaSO_4}$
6.		lost from H ₃ PO ₄) against a sta		sphoric acid (when the first H ⁺ rd NaOH solution, the indicator

	(a)	methyl orange	(b)	phenolphthalei	in
	(c)	1-naphtholphthalein	(d)	thymolphthale	in
7.	H ⁺ io	g the second stage of titration on is lost from $H_2PO_4^-$) again tor used is			
	(a)	methyl red	(b)	methyl orange	
	(c)	phenolphthalein	(d)	bromo-cresol g	reen
8.	NaNC	O_3 is estimated by			
	(a)	reducing NaNO ₃ by zinc in a titrated against a standard aci		kaline medium	to NH ₃ which is
	(b)	reducing NaNO ₃ by zinc in a titrated against a standard aci		idic medium to	NaOH which is
	(c)	reducing NaNO ₃ by zinc in a is titrated against a standard l			o NaNO ₂ which
	(d)	none of these			
9.	A 15-v	volume sample of an H ₂ O ₂ solu	ıtion	is equivalent to	
	(a)	5.30 N	(b)	1.77 N	
	(c)	2.68 N	(d)	7.50 N	
10.	acid i	pyrolusite is made to complet $_{2}$ pyrolusite is made to complet $_{2}$ $_{3}$ $_{4}$ $_{5}$ $_{6}$ $_{7}$ $_{7}$ $_{1}$ $_{2}$ $_{3}$ $_{4}$ $_{5}$ $_{7}$ $_{7}$ $_{1}$ $_{2}$ $_{3}$ $_{4}$ $_{5}$ $_{7}$ $_{7}$ $_{7}$ $_{1}$ $_{2}$ $_{3}$ $_{4}$ $_{7$			
	(a)	43.5 (b) 16	(c)	8 (d	.) 32
11.	100 m	L of N/5 NaOH will neutralize	9		
	(a)	0.06184 g of H ₃ BO ₃	(b)	$0.1855 \text{ g of H}_3\text{B}$	SO_3
	(c)	1.2368 g of H ₃ BO ₃	(d)	0.03092 g of H_3	BO_3
12.	Durin	g the titration of KMnO ₄ , the se	oluti	on is made acidi	ic with
	(a)	dilute H ₂ SO ₄	(b)	dilute HCl	
	(c)	concentrated H ₂ SO ₄	(d)	concentrated H	ICl
13.	$0.5~{\rm g}$ of an iron wire was dissolved in 75 mL of bench ${\rm H_2SO_{4}}$, and the volume was increased to 250 mL with water. 25 mL of this solution reduced $8.5~{\rm mL}$ of of $N/10~{\rm KMnO_4}$ solution. The amount of pure iron present was (atomic weight of Fe = 56)				
		0.500 g		0.252 g	
	(c)	0.238 g	(d)	0.476 g	
14.	Using it with	a standard KMnO $_4$ solution, the	ne fe	rric ion is estima	nted by reducing

(b) an FeSO₄ solution

(d) an HgCl₂ solution

(a) mercury

(c) an SnCl₂ solution

- 15. When we say that an H_2O_2 solution is a "10-volume" solution, we mean that, when fully decomposed by heat, it yields
 - (a) ten times its volume of oxygen measured at 273 K and 1 atmospheric pressure
 - (b) ten times its volume of oxygen measured at 298 K and 1 atmospheric pressure
 - (c) five times its volume of oxygen measured at 273 K and 10 atmospheric pressure
 - (d) its volume of oxygen measured at 273 K and 1 atmospheric pressure
- 16. Hydrazine reacts with KIO₃ in a concentrated HCl medium. This reaction may be represented as

$$N_2H_4 + IO_3^- + 2H^+ + Cl^- \longrightarrow ICl + N_2 + 3H_2.$$

Choose the correct option.

- (a) 1 mL of 0.25-M KIO₃ = 0.0008013 g of $N_2H_4 = 0.003253$ g of $N_2H_4.H_2SO_4$
- (b) 1 mL of 0.25-M $KIO_3 = 0.008013 g$ of $N_2H_4 = 0.03254 g$ of N₂H₄.H₂SO₄
- (c) 1000 mL of 1-M KIO₃ \equiv 0.032 g of N₂H₄ \equiv 0.130 g of N₂H₄.H₂SO₄
- (d) 1 mL of 0.5-M KIO₃ \equiv 0.016 g of N₂H₄ \equiv 0.0065 g of N₂H₄.H₂SO₄
- 17. In the estimation of hydroxylamine based upon the reduction of ferric solutions in the presence of H₂SO₄, by boiling and subsequent titration in the cold condition with a standard KMnO4 solution, which of the following reactions will occur.
 - (a) $NH_2OH + Fe^{3+} \rightarrow N_2H_4 + Fe^{2+} + H^+ + H_2O$
 - (b) $NH_2OH + Fe^{3+} \rightarrow N_2O + Fe^{2+} + H^+ + H_2O$
 - (c) $NH_2OH + Fe^{3+} \rightarrow NO_2 + Fe^{2+} + H^+ + H_2O$
 - (d) $NH_2OH + Fe^{3+} \rightarrow NH_3 + Fe^{2+} + H^+ + H_2O$
- 18. Copper is estimated iodometrically by titrating a solution of copper sulphate against a standard solution of
 - (a) potassium permanganate (b) potassium iodate
 - (c) potassium persulphate
- (d) sodium thiosulphate
- 19. A sample of 1.0 g of solid Fe₂O₃ of 80% purity is dissolved in a moderately concentrated HCl solution which is reduced by zinc dust. The resulting solution required 16.7 mL of a 0.1-M solution of the oxidant. Calculate the number of electrons taken up by the oxidant (the atomic weights of Fe and O are 56 and 16 respectively).
 - (a) 2

(b) 4

(c) 6

(d) 5

20. What volume of approximately 1.00 M HCl should be mixed with how

(b) 667 mL and 333 mL

(d) 250 mL and 750 mL

much 0.25 M HCl in order to prepare 1.00 L of 0.50 M HCl?

21. What volume of 0.25 M Ca(NO₃)₂ will contain 2.0 mol of nitrate ion?

(a) 667 mL and 333 mL(c) 500 mL and 500 mL

	(a)	0.5 L	(b)	1.0 L
	(c)	1.5 L	(d)	2.0 L
22.	the re (a)	volume of 0.1 M H_2SO_4 will be action $5H_2SO_4 + 8NaI \rightarrow 4Na_2S$ 70.0 L 25.0 L	(b)	aired to produce 17.0 g of H_2S by $4I_2 + H_2S + 4H_2O$? 50.0 L 5.0 L
23.	The 100 ($\rho = d$	relation between molarity ($N_1 = 1$) relation between molarity ($N_2 = 1$) relation, $N_3 = 1$	Л) a ılar v	and molality (<i>m</i>) is given by weight of solute)
	(a)	$m = \frac{1000 M}{1000 \rho - M_1}$	(b)	$m = \frac{1000 \ \rho \ M}{1000 \ \rho - MM_1}$
	(c)	$m = \frac{1000 \ MM_1}{1000 \ \rho - MM_1}$	(d)	$m = \frac{1000 M}{1000 \rho - MM_1}$
24.	The n	umber of iodine atoms present	in 5	0 mL of a 0.1 M KI solution is
	(a)	6×10^{23}	(b)	12×10^{23}
	(c)	3×10^{21}	(d)	6×10^{22}
25.	Calcu (densi	late the mass of anhydrous I ity = 1.2 g/mL^{-1}) which has 37	% H	-
	(a)	4.44 g	(b)	4.44 mg
	(c)	$4.44 \times 10^{-3} \text{ g}$	(d)	0.444 μg
26.		volume of $0.4 \text{ M FeCl}_3.6\text{H}_2\text{O}$ at of Fe = 55.85)?	will	contain 600 mg of Fe ³⁺ (atomic
	(a)	49.85 mL	(b)	26.85 mL
	(c)	147.55 mL	(d)	87.65 mL
27.		L of 10% NaOH (w/V) is added ant solution becomes	ed to	125 mL of 10% HCl (w/V). The
	. ,	alkaline		strongly alkaline
	(c)	acidic	(d)	neutral
28.		nple of Na ₂ CO ₃ .H ₂ O weighing ₄ solution. The resulting solution		g is added to 200 mL of a 0.1 N $$
		acidic	(b)	strongly acidic
	(c)	alkaline	(d)	neutral

- **29.** 0.2 mol of HCl and 0.1 mol of barium chloride were dissolved in water to produce a 500 mL solution. The molarity of the Cl⁻ ions is
 - (a) 0.06 M

(b) 0.09 M

(c) 0.12 M

- (d) 0.80 M
- 30. Equal volumes of $0.50\,\mathrm{M}$ HCl, $0.25\,\mathrm{M}$ NaOH and $2.75\,\mathrm{M}$ NaCl are mixed. The molarity of the NaCl solution is
 - (a) 0.75 M

(b) 1.00 M

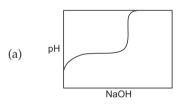
(c) 0.50 M

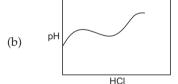
- (d) 2.00 M
- **31.** 25 mL of 2 N HCl, 50 mL of 4 N HNO $_3$ and x mL of 5 M H $_2$ SO $_4$ are mixed together and the total volume is made up to 1 L with water. 50 mL of this acid mixture exactly neutralizes 25 mL of a 1 N Na $_2$ CO $_3$ solution. The value of x is
 - (a) 25 mL

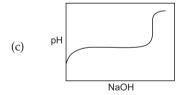
(b) 40 mL

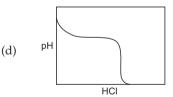
(c) 60 mL

- (d) 50 mL
- **32.** Which of the following indicates the titration of a weak base against a strong acid?



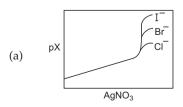


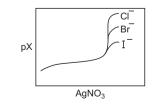


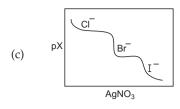


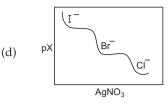
33. Which of the following represents the titration curve for 100 mL of 0.1 M Cl⁻, Br⁻ and I⁻ solutions against 0.1 M AgNO₃? (X = Cl⁻, Br⁻, I⁻)

(b)









- **34.** Which of the following processes are used to remove temporary as well as permanent hardness in water?
 - (a) Boiling the water
 - (b) Making the water react with lime
 - (c) Treating the water with a Ca(OH)₂ solution and then with HCl
 - (d) Boiling the water with a calculated quantity of Na_2CO_3 solution and then with HCl
- 35. 1 L of tap water contains 20 mg of Ca $^{2+}$ and 12 mg of Mg $^{2+}$ ions. What is the volume of a 2 N Na $_2$ CO $_3$ solution required to soften 1000 L of tap water?
 - (a) 100 L

(b) 1 L

(c) 1000 L

- (d) 10 L
- **36.** The most suitable indicator for the titration of a mixture of HCl and H_3PO_4 against NaOH will be
 - (a) methyl red and methyl orange
 - (b) methyl red and bromothymol blue
 - (c) methyl orange and phenolphthalein
 - (d) only methyl red
- 37. How many grams of copper will be replaced in 2 L of a 1.50 M CuSO₄ solution if the latter is made to react with 27.0 g of aluminium? (Cu = 63.5, Al = 27.0)
 - (a) 190.50 g

(b) 95.25 g

(c) 31.75 g

- (d) 10.65 g
- 38. The pH range of an indicator is given by
 - (a) pH = pK + 1

(b) pH = pK - 1

(c) pH = pK

- (d) $pH = pK \pm 1$
- **39.** The strength of a mixture of HCl and $\rm H_2SO_4$ is 0.1 N. On treatment with an excess of an AgNO $_3$ solution, 20 mL of this acid mixture gives 0.1435 g of AgCl. The strength of the $\rm H_2SO_4$ is
 - (a) 24.5 g L^{-1}

(b) 2.45 g L^{-1}

(c) 49 g L^{-1}

(d) 4.9 g L^{-1}

(a) 0.75 g of KCl

with a strong base?

(a) Methyl red

(c) 0.1 mol of KCl

40. To prepare a 0.5 M KCl solution from 100 mL of 0.40 M KCl, we need to

41. Which of the following indicators is used in the titration of a weak acid

(b) 20 mL of water

(d) 0.2 mol of KCl

(b) Cresol red

		Phenolphthalein	. ,	Methyl orange
42.	Which	•	are	used in the titration of KMnO ₄
		Starch		Phenolphthalein
	(c)	$K_2Cr_2O_7$	(d)	No indicator is necessary
43.	titrati	n of the following indicators (ron of $Na_2S_2O_3$ with I_2 ? Starch (external indicator)	atur	e indicated) are used during the
	(b)	Starch (internal indicator)		
	(c)	K ₂ CrO ₄ (external indicator)		
	(d)	Fluorescein (internal indicato	r)	
44.		der a pure H_2SO_4 (98% by y of 1.8. The molarity of the H_2		ght) solution having a specific is
	(a)	36 M	(b)	18 M
	(c)	9 M	(d)	72 M
45.	Why titration		d as	standard solutions in acid-base
	(a)	If we use them, the pH at the	equi	valence point will always be 7.0.
	(b)	They can be used to titrate str	ong	as well as weak bases.
	(c)	Strong acids form more stable	e solı	utions than weak bases do.
	(d)	Salts of strong acids and strong	ng ba	ses do not hydrolyse.
46.		is the volume of a sample of 0. of 0.20 M K_2CO_3 in the follow $K_2CO_3 + 2HNO_3 \longrightarrow 2KNO_3$	ing 1	
	(a)	40 mL		10 mL
		20 mL	. ,	80 mL
47.	The m in C ₆ H		of I ₂	in C_6H_6 is 0.2. The molality of I_2
	(a)	0.32	(b)	3.2
	(c)	0.032	(d)	0.48

48. Calculate the molality of a 1-L solution of 90% $\rm H_2SO_4$ (w/v), given that the density of the solution is 1.80 g mL $^{-1}$.

49. A sample of H₂SO₄ (density 1.8 g mL⁻¹) is 90% by weight. What is the volume of the acid that has to be used to make 1 L of 0.2 M H₂SO₄?

(b) 8.6 (d) 10.8

(b) 10 mL

(d) 18 mL

(a) 10.2

(c) 1.02

(a) 16 mL

(c) 12 mL

50.	Calculate the weight of the $BaCl_2$ needed to prepare 250 mL of a solution having the same concentration of Cl^- ions as that in a solution of KCl of concentration 80 g L $^{-1}$ (Ba = 137.4, Cl = 35.5).							
	(a)	27.92 g			(b)	14.50 g		
	(c)	22.52 g			(d)	11.46 g		
51.	water mL of	: Calculate the this solution	ne vo		HCl	gas at stp tha		aOH in 1 L of l neutralize 50
		224 mL				56 mL		
	(c)	112 mL			(d)	448 mL		
52.	NaOI	H to give a so			conce	entration of 0.0		0 mL of 0.1 M in H_2SO_4 ?
	` '	400 mL			` '	50 mL		
	(c)	200 mL			(d)	100 mL		
53.		naximum vol						. What will be com these two
	(a)	250 mL			(b)	750 mL		
	(c)	500 mL			(d)	1000 mL		
54.	NaOI		ne o	f the base r				on of 0.112 M 25.6 mL. The
			H_2F	$PO_4^- + OH^$	\longrightarrow	$HPO_4^{2-} + H_2C$)	(P = 31)
	The p	ercentage pu	rity (of KH ₂ PO ₄ is				
	(a)	46.32			(b)	79.32		
	(c)	78.42			(d)	88.45		
55.		L of a solution			ıtrali	zes 49.35 mL	of 4.	0 N HCl. The
			CO	$^{2-}_{3}$ + 2H $^{+}$ ——	→ C($O_2 + H_2O$		
		lensity of thi O_3 in it is	s Na	₂ CO ₃ solutio	n is	1.25 g mL $^{-1}$.	The	percentage of
	(a)	47.7	(b)	37.7	(c)	26.7	(d)	16.7

L-126	Chemistry	MCQ
56.	solution till all the NH ₃ (g) evolved	poiled with 100 mL of a 0.2 N NaOH and the remaining solution was diluted as neutralized using 10 mL of a 0.1 N rity of the $(NH_4)_2SO_4$ sample is (c) 47.4 (d) 79.8
57.		vas dissolved in water to make 1 L of required 10 mL of a 0.2 N AgNO ₃ Identify the alkali metal.
	(a) Na	(b) Li
	(c) K	(d) Cs
58.		have concentrations of 5N and 2N ons of A and B required to make 1 L of ther will be
	(a) $\frac{1}{4}$ L of A + $\frac{3}{4}$ L of B	(b) $\frac{2}{3}$ L of A + $\frac{1}{3}$ L of B
	(c) $\frac{1}{3}$ L of A + $\frac{2}{3}$ L of B	(d) $\frac{1}{5}$ L of A + $\frac{1}{2}$ L of B
59.	KMnO ₄ reacts with oxalic acid acco	ording to the equation
	$2MnO_4^- + 5C_2O_4^{2-} + 16H^+$	$\rightarrow 2Mn^{2+} + 10CO_2 + 8H_2O$
	Here 20 mL of 0.1-M KMnO ₄ is equ	
	(a) 120 mL of 0.25 M H ₂ C ₂ O ₄	

- 120 mL of 0.25 M $H_2C_2O_4$ (b) 150 mL of 0.10 M $H_2C_2O_4$
- (c) $50 \text{ mL of } 0.10 \text{ M H}_2\text{C}_2\text{O}_4$
- (d) $50 \text{ mL of } 0.20 \text{ M H}_2\text{C}_2\text{O}_4$
- 60. What would be the normality of a 0.1 M K₂Cr₂O₇ solution used as a precipitating agent for Pb ²⁺?
 - (a) 0.1 N
- (b) 0.6 N
- (c) 0.4 N
- (d) 0.2 N
- 61. A solution of H₂O₂ is titrated against a solution of KMnO₄. The reaction is $2MnO_4^- + 5H_2O_2 + 6H^+ \longrightarrow 2Mn^{2+} + 5O_2 + 8H_2O$

If it requires 46.9 mL of 0.145 M KMnO₄ to oxidize 20.0 g of H₂O₂, the mass percentage of H₂O₂ in this solution is

- (a) 2.9
- (b) 29
- (c) 21
- (d) 4.9

• *Type 2* •

Choose the correct options. More than one option is correct.

- **62.** Which of the following are primary standard substances?
 - (a) Na₂CO₃.10H₂O
- (b) NaOH
- (c) $Na_2B_4O_7.10H_2O$
- (d) KMnO₄

- **63.** Which of the following statements are correct?
 - (a) The point at which an equivalent amount of the titrant is added is called the equivalence point.
 - (b) The point at which the reaction is observed to be complete is called the end point.
 - (c) At the end point of a reaction there is no change in the properties of the solution.
 - (d) At the equivalence point of a reaction the stoichiometric amount of the titrant is not added.
- **64.** 100 mL of a $0.1 \,\mathrm{M}\,\mathrm{SO_4^{2-}}$ solution is
 - (a) 10 millimoles

- (b) 5 millimoles
- (c) 20 milliequivalents
- (d) 40 milliequivalents
- **65.** Which of the following will be present in the solution formed when 50 mL of 0.1 M HCl is mixed with 50 mL of 0.1 M NaOH?
 - (a) 4.5 m mol of H^+
- (b) $0.05 \text{ m} \text{ mol of OH}^-$

(c) 0.05 M NaCl

- (d) $10^{-7} \,\mathrm{M} \,\mathrm{of} \,\mathrm{H}^+\mathrm{ion}$
- **66.** Which of the following statements are correct?
 - (a) During the titration of a strong acid against a strong base, the pH at the equivalence point will be neutral.
 - (b) During the titration of a weak acid against a strong base, the pH at the equivalence point will be alkaline.
 - (c) During the titration of a weak base with a strong acid, the pH at the equivalence point will be acidic.
 - (d) During the titration of a weak acid against a weak base, the pH at the equivalence point will be neutral.
- 67. During the titration of a mixture of Na₂CO₃ and NaHCO₃ against HCl,
 - (a) phenolphthalein is used to detect the first end point
 - (b) phenolphthalein is used to detect the second end point
 - (c) methyl orange is used to detect the second end point
 - (d) methyl red is used to detect the first end point
- 68. 1 mol of H₂SO₄ will exactly neutralize
 - (a) 2 mol of ammonia
- (b) 1 mol of $Ba(OH)_2$
- (c) 0.5 mol of Ba(OH)₂
- (d) 2 mol of KOH
- **69.** At the end point there is a sharp change of colour in the indicator. This happens because the
 - (a) pH at the end point changes sharply
 - (b) structure of the indicator changes
 - (c) colour of indicator is adsorbed by water
 - (d) dissociation constants of acids and bases differ by ten

71. b, d

- 70. '20 volumes' of H_2O_2 is equal to
 - (a) $20\% H_2O_2$ by mass
- (b) $6\% H_2O_2$ by mass

(c) 1.764 N

- (d) 3.528 N
- 71. A solution of $Na_2S_2O_3$ is standardized iodometrically against 0.1262 g of KBrO₃. This process requires 45 mL of the $Na_2S_2O_3$ solution. What is the strength of the $Na_2S_2O_3$?
 - (a) 0.2 M
- (b) 0.1 M
- (c) 0.05 N
- (d) 0.1 N

Answers

1. a	2. a	3. b	4. c	5. b
6. a	7. c	8. a	9. c	10. b
11. c	12. a	13. d	14. c	15. a
16. a	17. b	18. d	19. c	20. a
21. b	22. c	23. d	24. c	25. a
26. b	27. c	28. d	29. d	30. b
31. a	32. d	33. a	34. d	35. b
36. b	37. b	38. d	39. b	40. a
41. c	42. d	43. a	44. b	45. b
46. d	47. b	48. a	49. c	50. a
51. c	52. d	53. b	54. b	55. d
56. a	57. c	58. c	59. c	60. d
61. a	62. a, c	63. a, b	64. a, c	65. c, d
66. a, b, c	67. a, c	68. a, b, d	69. a, b	70. b, d

Hints to More difficult Problems

 With phenolphthalein as indicator the only action is the conversion of the sodium carbonate to the bicarbonate.

$$\begin{array}{c} Na_{2}CO_{3} + HCl \rightarrow NaCl + NaHCO_{3} \\ 106 \text{ g} \end{array}$$

Equivalent weight of $Na_2CO_3 = \frac{106}{1} = 106$

2. Ba(OH)₂ + CO₂ \rightarrow BaCO₃ + H₂O 2000 mL of 1 N Ba(OH)₂ = 22400 mL of CO₂ 5 mL of N/50 Ba(OH)₂ = 1.12 mL of CO₂ Percentage by volume in air = $\frac{1.12}{2500} \times 100 = 0.045$.

3. When methyl orange is used as an indicator, Na₂CO₃ is converted into NaCl.

$$Na_2CO_3 + 2HCl \rightarrow 2NaCl + CO_2 + H_2O$$

 106 g 2×36.5

Equivalent weight of $Na_2CO_3 = \frac{106}{2} = 53$.

4.
$$x_1 = \frac{n_1}{n_1 + n_2}$$
 or $\frac{n_1 + n_2}{n_1} = \frac{1}{x_1}$.
 $1 + \frac{n_2}{n_1} = \frac{1}{x_1} = \frac{1}{0.5} = 2 \Rightarrow \frac{n_1}{n_2} = 1$
or $\frac{w_1 \times m_2 \times 1000}{m_1 \times w_2 \times 1000} = 1 \Rightarrow \left(\frac{w_1 \times 1000}{m_1 \times w_2}\right) \left(\frac{m_2}{1000}\right) = 1$.
Molality $\times \frac{18}{1000} = 1 \Rightarrow \text{molality} = \frac{1000}{18} = 55.55$.

- **10.** 1 milliequivalent of oxalic acid $\equiv 0.008$ g of oxygen. 20 milliequivalents of oxalic acid $\equiv 0.16$ g of oxygen.
 - \therefore percentage of available oxygen by weight = $\frac{0.16}{1.00} \times 100 = 16$.
- 13. 25 mL of solution = 8.5 mL of N/10 KMnO₄ solution. 250 mL of solution \equiv 8.5 milliequivalents of KMnO₄ solution $\equiv 8.5 \times 10^{-3} \times 56$ g of Fe = 0.476 g of Fe.
- **19.** Weight of pure $Fe_2O_3 = 1 \times \frac{80}{100} = 0.8 g$

$$Fe_2O_3 + 6HCl = 2FeCl_3 + 3H_2O$$

$$FeCl_3 + H_2 \xrightarrow{Zn \text{ dust}} FeCl_2 + HCl$$

 $Fe^{2+} + \text{ oxidant} \longrightarrow Fe^{3+} + \text{ reductant}$

$$Fe^{2+}$$
 + oxidant $\longrightarrow Fe^{3+}$ + reductant

Equivalent weight of
$$Fe_2O_3 = \frac{Molecular\ weight}{2} = \frac{160}{2} = 80\ g$$

Milliequivalents of Fe_2O_3 = milliequivalents of oxidant

$$\frac{0.8}{80} \times 10^3 = 16.7 \times 0.1(x) \Rightarrow x = 6$$

where *x* is the number of electrons involved.

22. 1 mol of $H_2S = 5$ mol of H_2SO_4 . $0.5 \text{ mol of } H_2SO_4 \equiv 2.5 \text{ mol of } H_2SO_4.$

Volume of
$$H_2SO_4 = \frac{2.5 \text{ mol}}{0.10 \text{ mol L}^{-1}} = 25.0 = 25.0 \text{ L}.$$

30. Let the volume of each electrolyte = x mL. Hence total volume = 3x mL x mL of 0.25 M NaOH reacts with x mL of 0.50 M HCl to produce x mL of 0.25 M NaCl solution.

Total number of milli moles of NaCl = 0.25x + 2.75x = 3.00x.

(Initially, the concentration of NaCl was 2.75 M)

$$\therefore$$
 Molarity of NaCl = $\frac{m \text{ mol}}{\text{mL}} = \frac{3.00 \text{ x mol}}{3 \text{ x mL}} = 1.00 \text{ M}.$

31. Use the normality equation.

Let N_1 = normality of acid mixture.

Let volume of $H_2SO_4 = x$ mL.

Then
$$(25 \times 2 + 50 \times 4 + x \times 5 \times 2)$$
 mL $(N) = 1000$ mL (N_1)

$$(50 + 200 + 10x) \text{ mL}(N) = 1000 \text{ mL } N_1$$
(1)

Secondly $[50 \text{ mL}(N_1)]_{acid \text{ mixture}} = [25 \text{ mL}(N)]_{alkali}$

$$N_1 = \frac{1}{2} N \tag{2}$$

From Equations (1) and (2),

$$(250 + 10x) \text{ mL}(N) = 1000 \text{ mL} (N/2)$$

$$\Rightarrow$$
 250 + 10 x = 500 mL

$$\Rightarrow x = 25.0 \text{ mL}.$$

35. Milliequivalents of Ca $^{2+}$ + milliequivalents of Mg $^{2+}$ = milliequivalents of N₂CO₃.

$$\frac{20}{20} + \frac{12}{12} = V \times 2.$$

 \therefore V = 1 mL for 1 L of tap water.

For 1000 L of tap water,

$$V = 1000 \text{ mL} = 1 \text{ L}.$$

37. 2 L of 1.5-M $CuSO_4 = 3 \text{ mol of } CuSO_4$

$$\begin{array}{c} 2\text{Al} + 3\text{CuSO}_4 \rightarrow \text{Al}_2(\text{SO}_4)_3 + 3\text{Cu} \\ 2\text{ mol} & 3\text{ mol} \end{array}$$

2 mol of Al \equiv 3 mol of Cu.

$$27 \text{ g} = 1 \text{ mol of Al} = \frac{3}{2} \times 63.5 \text{ g of Cu} = 95.25 \text{ g}.$$

39.
$$N_{\text{HCl}} + N_{\text{H}_2\text{SO}_4} = 0.1.$$
 (1)

1 equivalent of AgCl = 1000 mL (N).

 10^{-3} equivalent of AgCl = 1 mL (N)

$$[0.1435 \text{ g AgCl} = 1 \times 10^{-3} \text{ equivalent of AgCl}]$$

$$1 \text{ mL} (N) = 20 \text{ mL} \times N_{\text{HCI}} \Rightarrow N_{\text{HCI}} = 0.05.$$
 (2)

From Equations (1) and (2),

$$N_{\rm H_2SO_4} = 0.05 \ N.$$

Strength of $H_2SO_4 = 0.05 \times 49 = 2.45 \text{ gL}^{-1}$.

48. The sample of H_2SO_4 is 90% by volume

$$W_{\text{H}_2\text{SO}_4} = 900 \text{ g in } 1000 \text{ mL}$$

Volume of solution = 1000.0 mL

Weight of solution = $1000.0 \times 1.80 = 1800$ g.

Weight of water = 1800 - 900 = 900 g.

Molality =
$$\frac{900 \times 1000}{98 \times 900}$$
 = 10.2 mol kg⁻¹.

51.
$$N_{\text{NaOH}} = \frac{4 \text{ g}}{40 \text{ gram equivalent}} \times \frac{1}{1 \text{ L}} = 0.1 \text{ N}.$$

1 milliequivalent of NaOH \equiv 22.4 mL of HCl(g) at stp.

 50×0.1 milliequivalent of NaOH = $22.4 \times 50 \times 0.1$ mL of HCl (g) at stp = 112 mL of HCl (g) at stp.

52.
$$x \times 0.1 \times 2 - 50 \times 0.1 = (x + 50) \times 0.05 \times 2.$$

 $0.2x - 5 = 0.1x + 5 \Rightarrow 0.1x = 10 \Rightarrow x = 100 \text{ mL}.$

56. Milliequivalents of NaOH taken = 20.

Milliequivalents of H₂SO₄ = milliequivalents of NaOH reacted

$$= \frac{10 \times 0.1}{25} \times 250 = 10.$$

Milliequivalents of NaOH reacted = 20 - 10 = 10

1milliequivalentNaOH = 0.066 g of $(NH_4)_2SO_4$.

10 milliequivalents NaOH = 0.66 g of $(NH_4)_2SO_4$.

Percentage of
$$(NH_4)_2SO_4 = \frac{0.66}{0.70} \times 100 = 94.3$$
.

71. Eq. wt. of KBrO₃⁻ = $\frac{1}{6}$ of its mol. wt. = $\frac{1}{6} \times 167$ g equiv. ⁻¹

$$N_{\text{Na}_2\text{S}_2\text{O}_3} = \frac{0.1262 \text{ g}}{\frac{1}{6} \times 167 \text{ g equiv.}^{-1}} \times \frac{1}{0.045 \text{ L}} = 0.1 \text{ N}$$

For $Na_2S_2O_3$, molarity = normality, and so normality = 0.1 N and molarity = 0.1 M

11

Thermodynamics and Thermochemistry

• Type 1 •

1. Consider an ideal gas that occupies 2.50 dm³ at a pressure of 3.00 bar. If the gas is compressed isothermally at a constant pressure $p_{\rm ext}$, so that the final volume is 0.500 dm³, calculate the smallest possible value of $p_{\rm ext}$ and

2. Calculate the work done when 1 mol of an ideal gas is compressed reversibly from 1.00 bar to 5.00 bar at a constant temperature of 300 K.

(b) 15 bar and 750 J

(d) 10 bar and 375 J

(b) +18.02 kI

Choose the correct option. Only one option is correct.

the work done using p_{ext} . (a) 20 bar and 100 I

(c) 30 bar and 150 J

(a) -14.01 kI

	()		(-)	
	(c)	4.01 kJ	(d) -8.02 kJ	
3.			mol of an ideal gas is expande n ³ at a constant temperature of 300 k	
	(a)	7.78 kJ	(b) −1.73 kJ	
	(c)	11.73 kJ	(d) -4.78 kJ	
4.	work	done by the system, the proces	s decreases by the same amount as thess is (c) adiabatic (d) isolated	ıe
5.	(a) (b) (c)	heat is supplied to an ideal gas as will do positive work gas will do negative work kinetic energy of the gas will gas will not obey the law of control of the supplied in 1-132	conservation of energy	

pressure is
(a) negative

(c) infinity

(a) -1.435 cal g⁻¹

(c) 80.0 cal g^{-1}

6. The molar heat capacity of water in equilibrium with ice at constant

7. Calculate the enthalpy change when 1.0 g of water is frozen at 0°C $(\Delta H_f^{\circ} = 1.435 \, \text{kcal mol}^{-1})^3$?

(b) zero

(d) $40.45 \text{ kJ K}^{-1} \text{ mol}^{-1}$

(b) -80.0 cal g^{-1} (d) -55.6 cal g^{-1}

8.	The hydrogen-bond energy of water at 25°C is about							
	(a)	5 kcal mol	1		(b)	23 kcal m	10^{-1}	
	(c)	0.1 kcal mol	l ⁻¹		(d)	100 kcal ı	mol ⁻¹	
9.	eleme		ned	sets represe at 25°C	ent th and			al states of the ressure, with
		C(diamond						
		C(diamond	_	_	(1)			
		C(graphite)	_					
	(d)	C(graphite)	, Br ₂ (1)), I ₂ (s)				
10.	50 mL of water takes 5 minutes to evaporate from a vessel on a heater connected to an electric source which delivers 400 W. The enthalpy of vaporization of water is							
	(a)	40.3 kJ mol	-1		(b)	43.2 kJ m	ol -1	
	(c)	16.7 kJ mol	-1		(d)	180.4 kJ r	nol ⁻¹	
11.	The mathematical expression for the standard enthalpy of sublimation is given by							
	_	ΔH° (sublim	ation)	$=\Delta H^{\circ}(\text{fusion})$	n) –	2ΔH°(vap	orizatio	n)
	(b) ΔH° (sublimation) = ΔH° (fusion) – ΔH° (vaporization)							
	(c) ΔH° (sublimation) = ΔH° (fusion) + ΔH° (vaporization)							
	(d)	ΔH° (sublim	ation)	$=\Delta H^{\circ}(\text{coml})$	ousti	on) + ΔH°	(dissoci	ation)
12.	basis	of enthalpy o			droc	arbons is	the mos	st stable on the
	` '	1-Butene			` '	trans-2-B		
	(c)	2-Methylpro	opene		(d)	cis-2-Bute	ene	
13.		n of the foli lpy of format		g are exoth	iermi	ic compoi	unds w	rith respect to
	(a)	$C_2H_2(g)$	(b)	HI(g)	(c)	HBr(g)	(d)	$I_2(s)$

- **14.** Which among the following has maximum standard enthalpy of formation?
 - (a) Glucose

(b) Sucrose

(c) Hexane

(d) Ethanol

- **15.** In which of the following reactions is $\Delta H = \Delta U$?
 - (a) $H_2(g) + I_2(g) \rightarrow 2HI(g)$
 - (b) $KI(aq) + I_2(s) \rightarrow KI_3(aq)$
 - (c) $6NaOH(aq) + 3Cl_2(g) \rightarrow 5NaCl(aq) + NaClO_3(aq) + 3H_2O(l)$
 - (d) $N_2O_4(g) \rightarrow 2NO_2(g)$
- **16.** Which of the following salts should cause maximum cooling when 1 mol of it is dissolved in the same volume of water?
 - (a) NaCl; $\Delta H^{\circ} = 5.35 \text{ kJ mol}^{-1}$
 - (b) KNO₃; $\Delta H^{\circ} = 53.5 \text{ kJ mol}^{-1}$
 - (c) KOH; $\Delta H^{\circ} = -56.0 \text{ kJ mol}^{-1}$
 - (d) HBr; $\Delta H^{\circ} = -83.3 \text{ kJ mol}^{-1}$
- 17. The standard enthalpies of formation of H⁺ and OH⁻ ions in water are zero and -229.6 kJ mol⁻¹ respectively. The standard enthalpy of formation of liquid water is -285.6 kJ mol⁻¹. Then the enthalpy of neutralization of HCl(aq) by KOH(aq) is
 - (a) $229.6 \text{ kJ mol}^{-1}$

(b) $-173.4 \text{ kJ mol}^{-1}$

(c) 56.0 kJ mol^{-1}

(d) $-56.0 \text{ kJ mol}^{-1}$

18. Given that $\Delta H_{\rm f}^{\circ}({\rm CO_2}, {\rm g}) = -393.509 \text{ kJ mol}^{-1}$, $\Delta H_{\rm f}^{\circ}({\rm H_2O}, {\rm l}) = -285.830 \text{ kJ mol}^{-1}$, $\Delta H_{\rm f}^{\circ}({\rm C_2H_5OH}, {\rm l}) = -277.800 \text{ kJ mol}^{-1}$.

Calculate the ΔH° (combustion) of liquid ethanol. The following is the required reaction.

$$C_2H_5OH(1) + 3O_2(g) \rightarrow 2CO_2(g) + 3H_2O(1)$$

- (a) -1366.82 kJ mol⁻¹
- (b) $-278.82 \text{ kJ mol}^{-1}$
- (c) $-2264.66 \text{ kJ mol}^{-1}$
- (d) -1155.78 kJ mol⁻¹
- **19.** Given that the standard enthalpies of combination of C(s), $H_2(g)$ and $CH_4(g)$ are -393.5 kJ mol^{-1} , -285.8 kJ mol^{-1} and -890.4 kJ mol^{-1} respectively at 298 K. Calculate the standard enthalpy of formation of methane $[CH_4(g)]$.
 - (a) $-724.42 \text{ kJ mol}^{-1}$
- (b) −74.81 kJ mol⁻¹
- (c) $-114.82 \text{ kJ mol}^{-1}$
- (d) -194.62 kJ mol⁻¹
- **20.** The molar enthalpies of combustion of isobutane and n-butane are $-2870 \text{ kJ mol}^{-1}$ and $-2878 \text{ kJ mol}^{-1}$ respectively at 298 K and 1 atm. Calculate ΔH° for the conversion of 1 mol of n-butane to 1 mol of isobutane.

(a) -8 kJ mol^{-1}

- (b) $+8 \text{ kJ mol}^{-1}$
- (c) $-5748 \text{ kJ mol}^{-1}$

- (d) $+5748 \text{ kJ mol}^{-1}$
- **21.** Equal volumes of 1 M HCl and 1 M H₂SO₄ are neutralized by a 1 M NaOH solution, and x and y kJ/equivalent of heat are liberated respectively. Which of the following relations is correct?
 - (a) x = 2y

(b) x = 3y

(c) x = 4y

- (d) $x = \frac{1}{2}y$
- **22.** The enthalpies of hydrogenation of unsaturated hydrocarbons usually lie in the range
 - (a) 28–32 kcal mol⁻¹
- (b) 128–132 kcal mol⁻¹
- (c) 224–250 kcal mol⁻¹
- (d) 100–130 kcal mol⁻¹
- **23.** In which of the following thermochemical changes is ΔH° always negative?
 - (a) Enthalpy of solution
- (b) Enthalpy of hydrogenation
- (c) Enthalpy of reaction
- (d) Enthalpy of transition
- **24.** For which of the following processes does the enthalpy change represent the enthalpy of formation of NaOH?
 - (a) $Na^+(aq) + OH^-(aq) \longrightarrow NaOH(s)$
 - $\text{(b)} \quad Na(s) + \frac{1}{2}\,O_2(g) + \frac{1}{2}\,H_2(g) \, \longrightarrow \, NaOH(s)$
 - (c) $2Na(s) + 2H_2O(l) \longrightarrow 2NaOH(aq) + H_2(g)$
 - (d) All of these
- **25.** For the reaction $CO(g) + \frac{1}{2}O_2(g) \to CO_2(g)$; $\Delta H^{\circ} = -67650$ cal at 25°C.

Calculate ΔH° at 100°C, given that the required molar heat capacities are as follows.

$$C_p(CO, g) = 6.97 \text{ cal } ^{\circ}\text{C}^{-1}$$

 $C_p(CO_2, g) = 8.97 \text{ cal } ^{\circ}\text{C}^{-1}$
 $C_p(O_2, g) = 7.00 \text{ cal } ^{\circ}\text{C}^{-1}$

(a) -54.6 cal

(b) -67650.4 cal

(c) -67684.4 cal

- (d) -67762.5 cal
- **26.** The heat of atomization of $PH_3(g)$ is 228 kcal mol⁻¹ and that of $P_2H_4(g)$ is 355 kcal mol⁻¹. The energy of the P—P bond is
 - (a) 102 kcal mol⁻¹

(b) 51 kcal mol⁻¹

(c) 26 kcal mol^{-1}

(d) $204 \text{ kcal mol}^{-1}$

- 27. For a reversible process, the total change in entropy of the universe is
 - (a) $\Delta S(\text{system}) + \Delta S(\text{surroundings})$
 - (b) $\Delta S(\text{system}) \Delta S(\text{surroundings})$
 - (c) zero
 - (d) negative
- **28.** The molar enthalpy of fusion of water is 6.01 kJ mol⁻¹. The entropy change of 1 mol of water at its melting point will be
 - (a) 22 kJ mol⁻¹

(b) 109 kJ mol⁻¹

(c) 44 kJ mol⁻¹

- (d) 11 kJ mol⁻¹
- **29.** The enthalpy of fusion per gram and the corresponding molecular weights are given for five substances

	1	2	3	4	5
$\Delta H_{\rm fusion}^{\circ}({\rm cal~g}^{-1})$	100	40	80	40	60
Mol. wt. (g mol ⁻¹)	20	30	40	80	30

Which of the following pairs contains substances with the same molar enthalpy of fusion?

(a) 3 and 4

(b) 2 and 3

(c) 2 and 4

- (d) 3 and 4
- 30. The enthalpy of solution of a substance is given by
 - (a) $\Delta H_{\text{soln.}}^{\circ} = U \text{ (lattice energy)} + \Delta H_{\text{hydration}}^{\circ}$
 - (b) $\Delta H_{\text{soln.}}^{\circ} = U$ (lattice energy) $\Delta H_{\text{hydration}}^{\circ}$
 - (c) $\Delta H_{\text{soln.}}^{\circ} = -U \text{ (lattice energy)} + \Delta H_{\text{hydration}}^{\circ}$
 - (d) $\Delta H_{\text{soln.}}^{\circ} = -U \text{ (lattice energy)} \Delta H_{\text{hydration}}^{\circ}$
- 31. Which of the following statements is incorrect?
 - (a) There is nothing to surround a universe.
 - (b) The Clausius inequality rule states that $\int \delta s/T \le \Delta s$ for any process.
 - (c) In reversible expansion the system expands/contracts against maximum possible pressure.
 - (d) Gibbs free energy is not a path function.
- **32.** Given $C_p/C_v = \gamma$. A gas goes from the initial state (p_1, V_1, T_1) to a final state (p_2, V_2, T_2) through an adiabatic process. The work done by the gas is

(a)
$$\frac{nR(T_1 - T_2)}{\gamma - 1}$$

(b)
$$\frac{p_2V_2 - p_1V_1}{\gamma - 1}$$

(c)
$$\frac{p_1V_1 - p_2V_2}{\gamma + 1}$$
 (d) $\frac{n\gamma R(T_2 - T_1)}{\gamma - 1}$

- **33.** The molar heat capacity (C_m) of an ideal gas
 - (a) cannot be negative
 - (b) must be equal to either C_v or C_p
 - (c) must lie in the range $C_v < C_p$
 - (d) may have any value between $-\infty$ and $+\infty$
- **34.** Which of the following statements is correct?
 - (a) For every process in an isolated system the entropy increases.
 - (b) The sum of two path functions is also a path function.
 - (c) In a relativistic endothermic process, the total mass of the system increases.
 - (d) ΔH is always equal to $C_p \Delta T$.
- **35.** The mathematical formulation of the first law of thermodynamics, in differential form, is
 - (a) $\Sigma dU = 0$ (isolated system)
 - (b) dU = dq dw (closed system)
 - (c) $dU = dq dw + \Sigma \mu_i dn_i$ (open system)
 - (d) all of these
- **36.** Calculate the final temperature of a sample of a monoatomic gas that is expanded reversibly and adiabatically from 500 mL at 300 K to 2.00 L.
 - (a) 119 K
- (b) 150 K
- (c) 300 K
- (d) 20 K

37. For the reaction

$$\begin{split} 2C_6H_5CO_2H(s) + 13O_2(g) & \longrightarrow 12CO_2(g) + 6H_2O(g) \\ \Delta U^\circ &= -772.7 \text{ kJ mol}^{-1} \text{ at 298 K. Calculate } \Delta H^\circ. \end{split}$$

- (a) $+760.3 \text{ kJ mol}^{-1}$
- (b) $-760.3 \text{ kJ mol}^{-1}$
- (c) +670.3 kJ mol⁻¹
- (d) -790.3 kJ mol⁻¹
- **38.** The change in entropy when the pressure of a perfect gas is changed isothermally from p_1 to p_2 is
 - (a) $\Delta S = nR \ln(p_1 + p_2)$
- (b) $\Delta S = nR \ln(p_2/p_1)$
- (c) $\Delta S = nR \ln(p_1/p_2)$
- (d) $\Delta S = nR \ln \left(\frac{p_1 + p_2}{p_2} \right)$
- **39.** The enthalpy of neutralization of the reaction between CH₃COOH(aq) and NaOH(aq) is −13.2 kJ equiv. ⁻¹ and that of the reaction between H₂SO₄(aq) and KOH(aq) is −13.7 kJ equiv ⁻¹. The enthalpy of dissociation of CH₃COOH(aq) is
 - (a) -0.5 kJ equiv. $^{-1}$
- (b) $+0.5 \text{ kJ equiv.}^{-1}$

- (c) $-26.9 \text{ kI equiv.}^{-1}$ (d) +13.45 kJ equiv. -1
- 40. 80 kJ of heat is given to 36 g of water. Then the
 - (a) number of H⁺ and OH⁻ ions are $1.2044 \times 10^{24} \times \frac{80}{20}$ each
 - (b) number of water molecules that remain in solution is

$$\frac{36}{18} \times 6.02 \times 10^{23} \times 80$$

- (c) the ratio of H⁺ to OH⁻ $\left(\frac{H^+}{OH^-}\right)$ ions = $\frac{80 \times 36}{6.02 \times 10^{23}}$
- (d) numbers of H $^+$ and OH $^-$ ions are $1.2044 \times 10^{24} \times 80$ each
- **41.** The enthalpy of formation of $CO_2(g)$ at 298 K is -394 kJ mol⁻¹. Consider the following four reactions. (Given $\Delta H_{\text{comb}}^{\circ}(C_6H_6, l) = -3268 \text{ kJ mol}^{-1}$.)

$$(i) \ \frac{1}{6} \, C_6 H_6(l) + 1.25 \, O_2(g) \, \longrightarrow \, CO_2(g) + \frac{1}{2} \, H_2O(l)$$

(ii)
$$C(s) + \frac{1}{2}O_2(g) \longrightarrow CO(g)$$

(iii)
$$C(s) + 2O(s) \longrightarrow CO_2(g)$$

(iv)
$$C(s) + O_2(g) \longrightarrow CO_2(g)$$

In which of these reactions will 394 kJ of heat be evolved?

(a) (i)

(b) (ii) and (iv)

(c) (iv)

- (d) (ii), (iii) and (iv)
- 42. During an adiabatic process, the pressure of a gas is found to be proportional to the cube of its absolute temperature. The ratio $C_v/C_v = \gamma$ for the gas is
 - (a) $\frac{3}{2}$
- (b) $\frac{7}{2}$ (c) $\frac{5}{3}$
- **43.** C_v is always greater than C_v for a gas. Choose the correct option.
 - (a) When a gas absorbs heat at constant pressure, its volume remains unchanged.
 - (b) No work is done by a gas at constant volume.
 - (c) For the same change in temperature, the internal energy of a gas changes by a smaller amount at constant volume than at constant
 - (d) The mass of a gas at constant pressure is greater than that at constant volume.
- 44. 1 gram equivalent of H₂SO₄ is treated with 112 g of KOH for complete neutralization. Which of the following statements is correct?
 - 13.7 kcal of heat is evolved with the formation of 87 g of K₂SO₄, leaving no KOH.

- (b) 27.4 kcal of heat is evolved with the formation of 87 g of K₂SO₄, leaving 4 gram equivalent of KOH.
- (c) 15.7 kcal of heat is evolved with the formation of 1 gram equivalent of K_2SO_4 , leaving 56 g of KOH.
- (d) 13.7 kCal of heat is evolved with the formation of 87 g of K₂SO₄, leaving 1 gram equivalent of KOH.
- **45.** 20 g of CH₄ is burnt with 150 g of O₂, producing 675 kJ of heat. Assume that the amount of oxygen required is in accordance with the equation

$$CH_4(g) + 2O_2(g) \longrightarrow CO_2(g) + 2H_2O(l)$$

After the reaction, an excess of Cl_2 is added to the reaction mixture and 77 g of CCl_4 is obtained.

- (a) The enthalpy of combustion of methane is -900 kJ mol⁻¹ and the amount of O_2 remaining unused is 16 g.
- (b) The enthalpy of combustion of methane is -900 kJ mol^{-1} and the amount of O_2 remaining unused is 48 g.
- (c) The enthalpy of combustion of methane is -900 kJ mol⁻¹ and the amount of O_2 remaining unused is 54 g.
- (d) The enthalpy of combustion of methane is -920 kJ mol⁻¹ and the amount of O_2 remaining unused is 32 g.
- 46. The first law of thermodynamics introduces the concepts of conservation of
 - (a) heat
 - (b) energy and equivalence of heat and work
 - (c) work
 - (d) energy and equivalence of temperature and work
- **47.** A system is taken from state A to state B along two different paths 1 and 2. The heat absorbed and work done by the system along these paths are Q_1 and Q_2 and W_1 and W_2 respectively. Then

(a)
$$Q_1 = Q_2$$

(b)
$$Q_1 + W_1 = Q_2 + W_2$$

(c)
$$W_1 = W_2$$

(d)
$$Q_1 - W_1 = Q_2 - W_2$$

- **48.** For a gas, $C_p/C_v = \gamma$. The molecular mass of the gas is M. Its specific heat capacity at constant pressure is
 - (a) $\frac{\gamma R}{M(\gamma 1)}$

(b)
$$\frac{\gamma}{RM}$$

(c)
$$\frac{M}{R(\gamma-1)}$$

(d)
$$\frac{\gamma RM}{\gamma + 1}$$

- **49.** Which of the following statements is incorrect when H₂SO₄ gradually goes into solution with different number of moles of water?
 - (a) In the reaction $H_2SO_4(l) + n_1H_2O \rightarrow H_2SO_4(n_1H_2O)$, ΔH is known as the integral enthalpy of solution.

- (b) The difference between the integral enthalpies of solution of two differently specified concentrations of H2SO4 gives the enthalpy of dilution.
- (c) The slope of a plot of the integral enthalpy of solution against the number of moles of H₂SO₄ per mole of H₂O gives the integral enthalpy of solution at infinite dilution.
- (d) The slope of the plot of the integral enthalpy of solution against mol of H₂SO₄/mol of H₂O gives the differential enthalpy of solution.
- Calculate the difference between ΔH and ΔU when 1.0 mol of grey tin (density = 5.75 g cm^{-3}) changes to white tin (density = 7.31 g cm^{-3}) at 10.0bar (at 298 K, $\Delta H = +2.1$ kJ; atomic weight of Sn = 119.0).

 - (a) -8.8 I (b) -4.4 I
- (c) -2.2 J
- (d) +4.4 J
- 51. If water at 0°C, kept in a container with an open top, is placed in a large evacuated chamber,
 - (a) all the water will freeze
 - (b) all the water will vaporize
 - (c) part of the water will vaporize and the rest will freeze
 - (d) ice, water and water vapour will be formed and reach equilibrium at the triple point
- **52.** The standard enthalpy of formation of gaseous H_2O at 298 K is -241.82kJ mol⁻¹. Calculate $\Delta \dot{H}^{\circ}$ at 373 K given the following values of the molar heat capacities at constant pressure.

Molar heat capacity of $H_2(g) = 33.58 \text{ jK}^{-1} \text{ mol}^{-1}$

Molar heat capacity of $H_2(g) = 28.84 \text{ jK}^{-1} \text{ mol}^{-1}$

Molar heat capacity of $O_2(g) = 29.37 \text{ jK}^{-1} \text{ mol}^{-1}$

Assume that the heat capacities are independent of temperature.

- (a) $-242.6 \text{ kI mol}^{-1}$
- (b) $-485.2 \text{ kJ mol}^{-1}$
- (c) $-121.3 \text{ kJ mol}^{-1}$
- (d) $-286.4 \text{ kJ mol}^{-1}$
- 53. The enthalpy of neutralization of oxalic acid $(H_2C_2O_4)$ by a strong base is -25.4 kcal mol⁻¹. The enthalpy of neutralization of the reaction between a strong acid and strong base is -13.7 kcal equiv⁻¹. The enthalpy of dissociation of $H_2C_2O_4 \rightleftharpoons 2H^+ + C_2O_4^{2-}$ is
 - (a) $1.0 \text{ kcal mol}^{-1}$
- (b) $2.0 \text{ kcal mol}^{-1}$
- (c) 18.55 kcal mol⁻¹
- (d) 11.7 kcal mol⁻¹
- 54. The temperature of 25 mL of an HCl solution increases by 10°C when 25 mL of NaOH is added to it. What will be the increase in the temperature of 50 mL of this HCl solution when 50 mL of the same solution of NaOH is added to it?

the thermodynamic processes have proper analogs.

(b) 25°C

(d) 35°C

(a) The carnot cycle is not specifically meant for gases and can always be applied to mechanical or electromagnetic systems so long as

(b) The Carnot cycle can be carried out for nonthermodynamic systems such as those involving paramagnetic substances and

(d) The efficiency of a Carnot cycle is independent of the working substance employed because all reversible engines are necessarily

(a) 50°C

(c) 12.5°C

surface films.

55. Which of the following statements is incorrect?

(c) An ideal Carnot cycle has 100% efficiency.

less efficient than the Carnot heat engine. **56.** Which of the following has a positive enthalpy of formation? (b) COCl₂(l) (a) $NH_3(g)$ (c) $S_2Cl_2(s)$ (d) HN₃ 57. Which of the following has the greatest enthalpy of combustion per gram? (a) $H_2(g)$ (b) CH₂OH(l) (c) CH₃CH₂OH(l) (d) $C_2H_6(g)$ **58.** Which of the following statements is incorrect? (a) The specific heat capacity of a substance is greater in the solid state than in the liquid state. (b) The specific heat capacity of a substance is greater in the liquid state than in the solid state. (c) The latent heat of vaporization of a substance is greater than that of fusion. (d) The internal energy of an ideal gas is a function of its temperature. **59.** If we consider U as a function of S and V, then (a) $dU = \left(\frac{\partial U}{\partial S}\right)_{R} dS + \left(\frac{\partial U}{\partial V}\right)_{R} dV$ (b) $\left(\frac{\partial U}{\partial S}\right)_{R} = T$ (c) $\left(\frac{\partial U}{\partial V}\right)_{a} = -p$ (d) all of these **60.** Which of the following is an inexact differential? (b) $dq \times \frac{1}{T}$ (a) dq (c) C_n (d) H

- **61.** In which of the following processes does the entropy decrease?
 - (a) The dissolving of sodium chloride in water
 - (b) The evaporation of water
 - (c) The conversion of CO₂(g) into dry ice
 - (d) When a dozen marbles are taken out of a small bag and dropped on the ground
- **62.** A gas obeys the equation of state pV = nRT + nbp where b is a constant. If n moles of the gas expand from V_1 to V_2 reversibly at a constant temperature *T*, the work done is given by
 - (a) $-nRT \ln[(V_2 nb)/(V_1 nb)]$
 - (b) $nRT \ln[(V_2 nb)/V_2 nb)]$
 - (c) zero
 - (d) $-nRT \ln[(V_1 nb)/(V_2 nb)]$
- 63. Calculate the enthalpy change when 50 mL of 0.01 M Ca(OH)₂ reacts with 25 mL of 0.01 M HCl. Given that $\Delta H_{\text{neut}}^{\circ}$ of a strong acid and strong base is 140 kcal equivalent.
 - (a) 14.0 cal

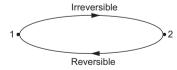
(b) 35 cal

(c) 10.0 cal

- (d) 7.5 cal
- 64. The enthalpy of neutralization of a strong acid by a strong base is -57.32 kJ mol⁻¹. The enthalpy of formation of water is -285.84 kJ mol⁻¹. The enthalpy of formation of the hydroxyl ion is
 - (a) +228.52 kJ mol⁻¹
- (b) $-114.26 \text{ kJ mol}^{-1}$
- (c) $-228.52 \text{ kJ mol}^{-1}$
- (d) +114.26 kJ mol⁻¹
- 65. A current of 10 A from a 12-V battery is passed through a heating element in a calorimeter for 300 s. If the temperature of the calorimeter rises by 5.5 K, the calorimeter constant is

 - (a) 4.5 kJ K^{-1} (b) 45.0 kJ K^{-1} (c) 450 kJ K^{-1} (d) 6.5 kJ K^{-1}

66. In a cyclic process given below



(a)
$$\triangle S = 0 > \int_{1}^{2} \frac{dq_{rev}}{T}$$

(b)
$$\Delta S = 0 > \int_{1}^{2} \frac{dq_{rev}}{T} + \int_{1}^{2} \frac{dq_{irr}}{T}$$

(c)
$$\Delta S = 0 > \int_{1}^{2} \frac{dq_{irr}}{T} + \int_{2}^{1} \frac{dq_{rev}}{T}$$

(c)
$$\triangle S = 0 > \int_{1}^{2} \frac{dq_{irr}}{T} + \int_{2}^{1} \frac{dq_{rev}}{T}$$
 (d) $\triangle S = 0 > \int_{2}^{1} \frac{dq_{irr}}{T} + \int_{2}^{1} \frac{dq_{rev}}{T}$

- 67. The total entropy change for irreversible process is given by
 - (a) $\Delta S_{\text{total}} = \Delta S_{\text{sys}} + \Delta S_{\text{surr}} = 0$
 - (b) $\Delta S_{\text{total}} = \Delta S_{\text{sys}} + \Delta S_{\text{surr}} = nR \ln V_2 / V_1$
 - (c) $\Delta S_{\text{total}} = \Delta S_{\text{sys}} + \Delta S_{\text{surr}} = nR \ln V_2 / V_1 + C_7 dT$
 - (d) $\Delta S_{\text{total}} = \Delta S_{\text{svs}} + \Delta S_{\text{surr}} = nR \ln V_1/V_2$
- **68.** Which of the following conditions for entropy is incorrect?
 - The state of equilibrium corresponds to maximum entropy of the universe
 - The system maintained at constant entropy and constant volume will attain the equilibrium at a state of minimum energy
 - necessary condition $(dS)_{U,V} < 0$, $(dU)_{S,V} > 0$ sufficient condition dS > 0, dU > 0.
 - (d) The necessary condition $(dS)_{U,V} > 0(dU)_{S,V} < 0$ sufficient condition dS < 0, dU < 0.
- **69.** In a reversible adiabatic change ΔS is
 - (a) infinity

(b) zero

(c) equal to $C_n dT$

- (d) equal to $nR\ln V_2/V_1$
- 70. In an adiabatic irreversible expansion, the external pressure is constant and the work of expansion is given by

(a)
$$(-W_{irr}) = P_{ext} \left(\frac{nRT_2}{p_2} - \frac{nRT_1}{p_1} \right)$$
 (b) $(-W_{irr}) = P_{ext} (nRT_2 - nRT_1)$

- (c) $(-W_{irr}) = P_{ext} (nRT_2 + nRT_1)$ (d) none of these
- 71. When ideal gases are mixed together, the molar entropy of mixing is given by
 - (a) $\Delta \overline{S}_{mix} = -R \sum_{i=1}^{N} y_i \ln y_i$
- (b) $\Delta \overline{S}_{mix} = -nR \sum_{i=1}^{N} y_i \ln y_j$
- (c) $\Delta \overline{S}_{mix} = +R \sum_{i=1}^{N} y_i \ln y_i$ (d) $\Delta \overline{S}_{mix} = -nRT \sum_{i=1}^{N} y_i \ln y_i$
- 72. The direction of a spontaneous process for a system at constant pressure and temperature is given by
 - (a) $d(U TS + PV) \ge 0$
- (b) $d(U TS + PV) \le 0$
- (c) $d(U TP + VS) \le 0$
- (d) $d(U TS + Vn) \le 0$

• Type 2 •

Choose the correct options. More than one option is correct.

73. Which of the following expression represents Gibbs-Helmholtz equation?

(a)
$$\Delta G^{\circ} = \Delta H^{\circ} + T \left[\frac{\partial (\Delta G^{\circ})}{\partial T} \right]_{n}$$
 (b) $\left[\frac{\partial (\Delta G^{\circ})}{\partial T} \right]_{n} = -\frac{\Delta H^{\circ}}{T^{2}}$

(b)
$$\left[\frac{\partial(\Delta G^{\circ})}{\partial T}\right]_{v} = -\frac{\Delta H^{\circ}}{T^{2}}$$

(c)
$$\left[\frac{\partial(\triangle G^{\circ}/T)}{\partial(1/T)}\right]_p = \triangle H^{\circ}$$

(d)
$$\left[\frac{\partial(\triangle G^{\circ}/T)}{\partial T}\right]_p = -\triangle H^{\circ}$$

74. In an isothermal irreversible expansion of an ideal gas

(a)
$$\Delta U = 0$$

(b)
$$-wq = nRT[1 - P_2/P_1]$$

(c)
$$\Delta H \neq 0$$

- (d) all of these
- 75. The differential form of thermodynamic energies U, H and G can be represented by

(a)
$$dU = Tds - pdv$$

(b)
$$dH = Tds + Vdp$$

(c)
$$dG = -SdT + Vdp$$

- (d) none of these
- 76. Which of the following mathematical relations are correct for an ideal gas?

(a)
$$\left(\frac{\partial H}{\partial V}\right)_T = 0$$

(b)
$$\left(\frac{\partial H}{\partial p}\right)_T = 0$$

(c)
$$C_p - C_V > R$$

(d)
$$\left(\frac{\partial C_V}{\partial V}\right)_T = 0$$

77. Which of these mathematical expressions is a perfect differential?

(a)
$$\int_{1}^{2} dU = \Delta U$$

(b)
$$\int_{1}^{2} \delta W = \Delta W$$

(c)
$$\int_{1}^{2} \delta q = \Delta q$$

(d)
$$dU = \delta q + \delta W$$

- 78. An adiabatic process is one in which
 - (a) all energy is transferred as heat
 - (b) no energy is transferred as heat
 - (c) the temperature of a gas decreases in a reversible adiabatic expansion
 - (d) $dU \neq dW$
- 79. When the gas is ideal and the process is isothermal
 - (a) $p_1V_1 = p_2V_2$ (b) $U_1 = U_2$ (c) $H_1 \neq H_2$ (d) W = 0

- **80.** Which of the following statements are correct?
 - (a) The work done by the system on the surroundings is negative.
 - (b) The work done on the system by the surroundings is positive.
 - (c) The heat absorbed by the system from the surroundings is positive.
 - (d) The heat absorbed by the surroundings from the system is negative.
- **81.** Which of the following is a path function as well as an extensive property?
 - (a) Temperature

- (b) Internal energy
- (c) Molar heat capacity
- (d) Heat capacity
- 82. For ideal diatomic gases,
 - (a) $C_n = (7/2)R$

(b) $C_v = (3/2)R$

(c) $C_V = (5/2)R$

- (d) $C_V = (3/2)R$
- 83. When a solid melts there will be
 - (a) an increase in enthalpy
 - (b) a decrease in free energy
 - (c) no change in enthalpy
 - (d) a decrease in internal energy
- 84. Which of the following are thermodynamically stable?
 - (a) C(diamond)

(b) C(graphite)

(c) P₄(white)

- (d) $P_4(red)$
- 85. Which of the following are endothermic compounds?
 - (a) $N_2O_4(g)$

(b) $NO_2(g)$

(c) $N_2O_4(l)$

- (d) N₂H₄(l)
- **86.** The standard molar enthalpy of CO₂ is equal to
 - (a) the standard molar enthalpy of combustion of gaseous carbon
 - (b) the standard molar enthalpy of combustion of carbon (graphite)
 - (c) the sum of the standard molar enthalpies of formation of CO and O_2
 - (d) -394 kJ mol^{-1}
- 87. Which of the following statements are correct?
 - (a) The entropy of an isolated system increases in an irreversible process.
 - (b) The entropy of an isolated system remains unchanged in a reversible process.

- (c) Entropy can never decrease.
- (d) ΔS (system) as well as ΔS (surroundings) are negative quantities.
- 88. Which of the following statements are correct?
 - (a) When $\Delta G = 0$, the system is at equilibrium.
 - (b) When $\Delta G < 0$, the process will be spontaneous.
 - (c) When ΔG is negative, the process is said to be exergonic.
 - (d) When ΔG is positive, the process is said to be endergonic.
- **89.** Which of the following relations are correct?

(a)
$$\Delta G = \Delta H - T \Delta S$$

(b)
$$\Delta G = \Delta H + T \left[\frac{\partial (\Delta G)}{\partial T} \right]_p$$

(c)
$$\Delta G = \Delta H + T \Delta S$$

(d)
$$\Delta G = \Delta H + \Delta nRT$$

90. In the following table, which of the options are correct?

	ΔΗ	ΔS	Nature of reaction
(a)	(-)	(+)	Spontaneous at all temperatures
(b)	(+)	(-)	Nonspontaneous regardless of temperature
(c)	(+)	(+)	Spontaneous only at high temperature
(d)	(-)	(-)	Spontaneous only at low temperature

Answers

1. b	2. c	3. b	4. c	5. a	
6. c	7. b	8. a	9. d	10. b	
11. c	12. c	13. c	14. b	15. a	
16. b	17. d	18. a	19. b	20. a	
21. d	22. a	23. b	24. b	25. d	
26. b	27. c	28. a	29. d	30. a	
31. b	32. a	33. d	34. c	35. d	
36. a	37. b	38. c	39. b	40. a	
41. c	42. a	43. b	44. d	45. c	
46. b	47. d	48. a	49. c	50. b	
51. c	52. a	53. b	54. b	55. d	
56. d	57. a	58. b	59. d	60. a	
61. c	62. a	63. b	64. c	65. d	
66. c	67. b	68. c	69. b	70. a	
71. a	72. b	73. a, c	74. a, b	75. a, b, c	

76. a, b, d	77. a, d	78. b, c	79. a, b	80. a, b, c, d
81. b, d	82. a, c	83. a, b	84. b, c	85. a, b, d
86. b, d	87. a, b, c	88. a, b, c, d	89. a, b	90. a, b, c, d

Hints to More Difficult Problems

1. For a compression to occur, $p_{\rm ext}$ must be at least as large as the final pressure of the gas. The final pressure of the gas is

$$P_{\rm f} = \frac{p_{\rm i}V_{\rm i}}{V_{\rm f}} = \frac{(3.0 \text{ bar})(2.50 \text{ dm}^3)}{0.50 \text{ dm}^3} = 15.0 \text{ bar}$$

This is the smallest possible value of $p_{\rm ext}$ that can be applied to compress the gas isothermally from 2.50 dm³ to 0.5 dm³. The work done involving the value of $p_{\rm ext}$ is

$$W = -p_{\text{ext}}\Delta V = -(15 \text{ bar})(-0.5 \text{ dm}^3) = 7.5 \text{ dm}^3 \text{ bar}$$
$$= (7.5 \text{ dm}^3 \text{ bar})(10^{-3} \text{ m}^3 \text{dm}^{-3})(10^5) \text{ Pa bar}^{-1}$$
$$= 750 \text{ Pa} \cdot \text{m}^3 = 750 \text{ J}.$$

6.
$$C_{p,m} \underset{T \to 0}{\underset{Lt}{C}} \left(\frac{\partial H}{\partial T} \right)_p = \infty$$

10. Work done in 1 s = 400 J.

Work done in 300 s = 120,000 J = 120 kJ.

$$\Delta H^{\circ}_{\text{vap}} = \frac{120 \text{ kJ}}{50/18 \text{ mol}} \text{ (assume density of water} = 1.00 \text{ g cm}^{-3}\text{)}$$

= 43.2 kJ mol⁻¹.

- **15.** $\Delta n = 0$, and hence $\Delta H = \Delta U$.
- **20.** $\Delta H_{\text{iso}}^{\circ} = -2878 (-2870) = 8 \text{ kJ mol}^{-1}.$
- **26.** Bond dissociation energy of $PH_3(g) = 228 \text{ kcal mol}^{-1}$.

P—H bond energy =
$$\frac{228}{3}$$
 = 76 kcal mol⁻¹

Bond energy of $4(P-H) + (P-P) = 355 \text{ kcal mol}^{-1}$ or, $4 \times 76 + (P-P) = 355 \text{ kcal mol}^{-1}$ $\therefore P-P \text{ bond energy} = 51 \text{ kcal mol}^{-1}$

33. By definition, $C_m = \frac{1}{n} \frac{\Delta Q}{\Lambda T}$.

Set ΔQ and ΔT to get the result.

36. Use the equation
$$\frac{T_2}{T_1} = \left(\frac{V_1}{V_2}\right)^{\frac{2}{3}}$$
.

39.
$$\Delta H_{\text{neut}}^{\circ} \text{ (obs)} = \Delta H_{\text{neut}}^{\circ} \text{ (actual)} + \Delta H_{\text{diss}}^{\circ}$$

42.
$$T^{\gamma}p^{1-\gamma} = \text{constant}$$
 or $p \propto T^{(\gamma/\gamma-1)}$. Given, $p \propto T^3$.

$$\therefore \quad \frac{\gamma}{\gamma - 1} = 3 \Longrightarrow \gamma = \frac{3}{2} \cdot$$

44. Use the principle of enthalpy of neutralization.

48. Specific heat capacity =
$$\frac{\text{molar heat capacity}}{\text{molar mass}}$$
.

50.
$$V_{\text{grey tin}} = \frac{119.0}{5.75} = 20.696.$$

$$V_{\text{white tin}} = \frac{119.0}{7.31} = 16.279.$$

$$p\Delta V = (10 \times 10^{5} \text{ Pa}) (-20.696 + 16.279) \times 10^{-6} \text{ m}^{3}.$$

= -4.4 kJ

52. The reaction is
$$H_2(g) + \frac{1}{2}O_2(g) \longrightarrow H_2O(g)$$

$$\Delta C_{p}^{\circ} = C_{p,m}^{\circ}(H_{2}O, g) - \left\{C_{p,m}^{\circ}(H_{2}, g) + \frac{1}{2}C_{p,m}^{\circ}(O_{2}, g)\right\}$$
$$= 33.58 - \left\{28.84 + \frac{1}{2}(29.37)\right\}$$

$$= -9.94 \text{ JK}^{-1} \text{ mol}^{-1}$$
.

Using Kirchhoff's equation,

$$\Delta H^{0}(373 \text{ K}) = \Delta H^{\circ}(298 \text{ K}) + (T_{2} - T_{1})\Delta C^{\circ}_{p}$$
$$= -241.82 + (373 - 298) \times (-9.94)$$
$$= -242.6 \text{ kJ mol}^{-1}$$

53. $\Delta H_{\text{neut}}^{\circ}(\text{obs}) = \Delta H_{(\text{neut})}^{\circ} (\text{actual}) + \Delta H_{\text{diss}}^{\circ}$. Since the acid is diprotic,

$$\Delta H_{\text{diss}}^{\circ} = -25.4 - (-13.7 \times 2)$$
$$= 27.4 - 25.4$$
$$= 2.0 \text{ kcal mol}^{-1}.$$

- 58. Lower molar mass.
- **59.** Differentiate U = f(S, V) and compare with the equation dU = TdS pdV.
- **60.** *q* is not a state function.

63. Number of milliequivalent of $Ca(OH)_2 = 50 \times 2 \times 0.01 = 1.00$.

Number of milliequivalents of HCl = $25 \times 0.01 = 0.25$.

0.25 milliequivalents of HCl will be neutralized by 0.25 milliequivalents of $Ca(OH)_2$.

1 equivalent corresponds to 140.0 kcal

Therefore, 0.25×10^{-3} equivalent corresponds to $140.0 \times 0.25 \times 10^{-3}$ kcal = 35 cal.

64. (a)
$$H^{+}(aq) + OH^{-}(aq) \longrightarrow H_{2}O(1)$$
, $\Delta H^{\circ} = -57.32$ kJ mol⁻¹

(b)
$$H_2(g) + \frac{1}{2} O_2(g) \longrightarrow H_2O(l), \Delta H^{\circ} = -285.84 \text{ kJ mol}^{-1}$$

$$(a)-(b)=H^{+}(aq)+OH^{-}(aq) \xrightarrow{} H_{2}(g)+\frac{1}{2} \underset{\text{zero}}{O_{2}(g)} \Delta H^{\circ}=+228.52 \text{ kJ mol}^{-1}$$

$$\Delta H^{\circ} = \Sigma \Delta H_n^{\circ} - \Sigma \Delta H_R^{\circ}$$

$$228.52 = (0+0) - (x+0)$$

$$\therefore x = -228.52 \text{ kJ mol}^{-1}.$$

- **69.** For adiabatic changes q = 0
- 77. Internal enegry (U) is a state function.
- **80.** Use the law of acquisition.

Chemical Equilibrium

• Type 1 •

Choose the correct option. Only one option is correct.

- 1. Which of the following is true at chemical equilibrium?
 - (a) $(\Delta G)_{T, p}$ is minimum and $(\Delta S)_{U, V}$ is also minimum.
 - (b) $(\Delta G)_{T,V}$ is minimum and $(\Delta S)_{U,V}$ is maximum.
 - (c) $(\Delta G)_{T, V}$ is maximum and $(\Delta S)_{U, V}$ is zero.
 - (d) $(\Delta G)_{T, p}$ is zero and $(\Delta S)_{U, V}$ is also zero.
- 2. In the reaction, $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$,

(a)
$$K_p = K_x(RT)^2$$

(b)
$$K_p = \frac{K_N}{(RT)^2}$$

(c)
$$K_c = \frac{K_p}{(RT)^{-2}}$$

(d)
$$K_p = \frac{K_x}{(RT)^2}$$

- 3. The equilibrium of the reaction $N_2(g)+3H_2(g)\rightleftharpoons 2NH_3(g)$ will shift to the product side when
 - (a) $K_p > 1$

(b)
$$Q < K_p$$

(c) $Q = K_p$

(d)
$$Q < 2K_p$$

4. Which of the following expressions is correct?

(a)
$$K_p = K_c \left(\frac{RT}{\Sigma n}\right)^{\Delta n}$$

(b)
$$K_p = K_x \left(\frac{p}{\Sigma n}\right)^{\Delta n}$$

(c)
$$K_p = K_N \left(\frac{p}{\Sigma n}\right)^{\Delta n - 1}$$

(d)
$$K_p = K_c \left(\frac{p}{\Sigma n}\right)^{-\Delta n + 1}$$

5. It is found that the equilibrium constant increases by a factor of four when the temperature is increased from 25°C to 40°C. The value of ΔH° is

6. In which of the following systems will the equilibrium be shifted to the

7. The equilibrium constants for the following reactions at 1400 K are given.

(b) 20.5

(d) 8.4

right if the volume is increased at constant temperature?

 $2H_2O(g) \rightleftharpoons 2H_2(g) + O_2(g); \ K_1 = 2.1 \times 10^{-13}$ $2CO_2(g) \rightleftharpoons 2CO(g) + (g); \ K_2 = 1.4 \times 10^{-12}$ Then the equilibrium constant *K* for the reaction $H_2(g) + CO_2(g) \rightleftharpoons CO(g) + H_2O(g)$

(b) 171.67 kJ mol⁻¹

(d) 71.67 kJ mol⁻¹

(b) $A_2(g) + 3O_2(g) \rightleftharpoons 2AO_3$

(d) $A_2(g) + 4B_2(g) \rightleftharpoons 2AB_4(g)$

(a) $25.46 \text{ kJ mol}^{-1}$

(c) 89.43 kJ mol⁻¹

(c) $A_2(g) \rightleftharpoons 2A(g)$

is

(a) 2.04

(c) 2.6

(a) $A_2(g) + B_2(g) \rightleftharpoons 2AB(g)$

	` '				` '		
8.	is 4.0	at 1000 K. Who K, the percenta 67) + F en e	$H_2O(g) \rightleftharpoons CC$ quimolar mi) ₂ (g) xtur	+H ₂ (g) es of CO and equilibrium i 33	$ m H_2O$ are heated to $ m s$
9.	mol o		ith 3	B mol of B in	a 1-I		In an experiment a quilibrium 3 mol of
	(c)	2.5 mol			(d)	3.5 mol	
10.		ne reaction AF are of p . Then	B(g)	\rightleftharpoons A(g) + B(g	g), A	B is 33% dis	ssociated at a total
	(a)	$p = K_p$			(b)	$p = 4K_p$	
	(c)	$p = 3K_p$			(d)	$p = 8K_p$	
l1.	value	ne equilibrium of K_p is four the les of A formed	mes	the reaction that of the t	AB(total	$g) \rightleftharpoons A(g) + F$ pressure. Ca	B(g), the numerical lculate the number
	(a)	0.1	(b)	0.09	(c)	0.05	(d) 0.9
12.	For th	e reaction A(g	\Rightarrow	B(g) + C(g),			
	(a)	$K_p = \alpha^3 p$			(b)	$K_p = \alpha^2 (K_p +$	p + 1)
	(c)	$K_p = \alpha^2 (K_p + \mu$))		(d)	$K_p = \alpha^2 \left(\frac{K_p}{p} \right)^{\frac{1}{2}}$	$\left(\frac{p}{p}\right)$

- 13. Consider the reactions
 - (i) $CO(g) + H_2O(g) \rightleftharpoons CO_2(g) + H_2(g)$; K_1
 - (ii) $CH_4(g) + H_2O(g) \rightleftharpoons CO(g) + 3H_2(g)$; K_2
 - (iii) $CH_4(g) + 2H_2O(g) \rightleftharpoons CO_2(g) + 4H_2(g)$; K_3

Which of the following is correct?

(a) $K_3 = K_1/K_2$

(b) $K_2 = K_1^2 / K_2^3$

(c) $K_3 = K_1 K_2$

- (d) $K_3 = K_1 \sqrt{K_2}$
- 14. For the equilibrium of the reaction

$$HgO(s) \rightleftharpoons Hg(g) + \frac{1}{2}O_2(g)$$

(a) $K_p = \frac{2}{2^{3/2}} p^{3/2}$

(b) $K_p = \frac{2}{2^{1/2}} p^{1/2}$

(c) $K_p = \frac{1}{2^{2/3}} p^{3/2}$

- (d) $K_p = \frac{1}{2^{2/3}} p$
- 15. The total pressure observed at equilibrium in the dissociation of solid ammonium carbamate at a certain temperature is 2.0 atm. The equilibrium constant K_p is
 - (a) 4.185
- (b) 1.185
- (c) 2.276
- (d) 1.072
- 16. XY_2 dissociates as $XY_2(g) \rightleftharpoons XY(g) + Y(g)$. When the initial pressure of XY_2 is 600 mm of Hg, the total pressure developed is 800 mm of Hg. K_n for the reaction is
 - (a) 200
- (b) 50
- (c) 100
- (d) 150

17. For the reaction

$$H_2(g) + I_2(g) \rightleftharpoons 2HI(g)$$

the equilibrium constant K_v depends on the

(a) total pressure

- (b) catalyst used
- (c) amount of H₂ and I₂
- (d) temperature
- 18. When pressure is applied to the equilibrium system ice(s) \rightleftharpoons H₂O(l), which of the following will happen?
 - (a) More ice will be formed.
 - (b) Water will evaporate.
 - (c) More water will be formed.
 - (d) Equilibrium will not be reached.
- 19. In which of the following reactions is the standard reaction entropy positive and does ΔG° decrease sharply with increasing temperature?

 - $\text{(a)} \quad M(s) + \frac{1}{2} \, O_2(g) \, \to \, MO(s) \qquad \text{(b)} \quad \frac{1}{2} \, C(s) + \frac{1}{2} \, O_2(g) \, \to \, \frac{1}{2} \, CO_2(g)$

 - (c) $C(s) + \frac{1}{2}O_2(g) \rightarrow CO(g)$ (d) $CO(g) + \frac{1}{2}O_2(g) \rightarrow CO_2(g)$

20. The standard free energy of formation of NH₃(g) is -16.48 kJ mol⁻¹. The value of the equilibrium constant (R = 8.314 JK ⁻¹ mol⁻¹) is

21. Which of the following is correct for the equilibrium of the reaction $C(s) + H_2O(g) \rightleftharpoons CO(g) + H_2(g)$

(b) 3×10^5 (d) 9×10^5

(b) $p_{\rm H_2} \propto \sqrt{p_{\rm H_2O}}$

(a) 6×10^{5}

(c) 1×10^5

(a) $p_{\rm H_2} \propto p_{\rm H_2O}$

	(c)	$p_{\rm H_2} \propto p_{\rm H_2O}^2$	(d)	$p_{\rm H_2} \propto \frac{p_{\rm H_2O}}{p_{\rm CO}}$
22.	are si volur some (a) (b) (c)		in a ater	an equilibrium box at constant introduced into the vessel. After
23.	point of (a)		(g) + (b)	on of helium at the equilibrium $Cl_2(g)$, the degree of dissociation PCl_5 will increase Cl_2 will increase
24.	N ₂ (g) (a) (b) (c)	onstant pressure, the presence $+3H_2(g) \rightleftharpoons 2NH_3(g)$ will reduce the formation of NH_3 increase the formation of H_2 increase the formation of both	-3	f argon at the equilibrium of $\label{eq:hard_eq} \text{and } \mathbf{H}_2$
25.	molai (a)	ne equilibrium of the reaction r mass of N_2O_4 is 77.70 g. The p 28.4 22.4	erce (b)	$O_4(g) \rightleftharpoons 2NO_2(g)$, the observed ntage dissociation of N_2O_4 is 46.7
26.	2. Cal (a)		ion (b)	e reaction $N_2O_4(g) \rightleftharpoons 2NO_2(g)$ is of N_2O_4 at a pressure of 0.5 atm. 50

- 27. 1 mol of $N_2O_4(g)$ at 300 K is kept in a closed container under 1 atm. It is heated to 600 K, upon which 20% by mass of $N_2O_4(g)$ decomposes to $NO_2(g)$. The resultant pressure is
 - (a) 1.2 atm

(b) 2.4 atm

(c) 2.0 atm

- (d) 1.0 atm
- **28.** A reaction at equilibrium involving 2 mol each of PCl₅, PCl₃ and Cl₂ is maintained at 250°C and a total pressure of 3 atm. The value of K_p is
 - (a) 2

(b) 3

(c) 4

- (d) 1
- 29. The equilibrium constant for the reaction

$$CO(g) + H_2O(g) \rightleftharpoons CO_2(g) + H_2(g)$$

is 3.0 at 723 K and 1 atm pressure. The initial amount of water gas is 60 g and that of steam 90 g. At equilibrium the number of moles of CO₂ is

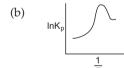
(a) 4.5

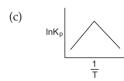
(b) 1.5

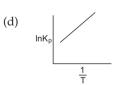
(c) 2.5

- (d) 3.5
- **30.** At the equilibrium of the reaction $2X(g) + Y(g) \rightleftharpoons X_2Y(g)$, the number of moles of X_2Y at equilibrium is affected by the
 - (a) temperature and pressure
 - (b) temperature only
 - (c) pressure only
 - (d) temperature, pressure and catalyst used
- 31. Which of the following graphs represents an exothermic reaction?









32. K_c for the equilibrium of

$$CH_3CO_2H(l) + C_2H_5OH(l) \rightleftharpoons CH_3CO_2C_2H_5(l) + H_2O(l)$$

is 4. How much ester is formed if 2 mol of the acid and 1 mol of the alcohol are taken initially to bring the reaction to equilibrium?

(a) 3.15

(b) 0.85

(c) 1.25

(d) 0.25

33. Consider the dissociation $N_2O_4(g) \rightleftharpoons 2NO_2(g)$. If we start with n_0 moles of $N_2O_4(g)$, then the extent of the reaction is given by

(a)
$$\frac{\zeta_{\text{eq}}}{n_0} = \left(\frac{K_p}{K_p + 4p}\right)$$

(b)
$$\frac{\zeta_{\text{eq}}}{n_0} = \left(\frac{K_p + 4p}{K_p}\right)^{\frac{1}{2}}$$

(c)
$$\frac{\zeta_{\text{eq}}}{n_0} = \left(\frac{4K_p}{p}\right)^{\frac{1}{2}}$$

(d)
$$\frac{\zeta_{eq}}{n_0} = \left(\frac{K_p}{K_p + 4p}\right)^{\frac{1}{2}}$$

34. The equilibrium constant for the decomposition of water

$$[H_2O(g) \longrightarrow H_2(g) + \frac{1}{2}O_2(g)]$$

is given by

(a)
$$K = \frac{\alpha^3 p^{1/2}}{(1+\alpha)(2-\alpha)^{1/2}}$$

(b)
$$K = \frac{\alpha^{3/2} p^{1/2}}{(1-\alpha)(2+\alpha)^{1/2}}$$

(c)
$$K = \frac{\alpha^3 p^{\frac{1}{2}}}{\sqrt{2}}$$

(d)
$$K = \frac{\alpha^3 p^{3/2}}{(1-\alpha)(2+\alpha)^{1/2}}$$

- **35.** In the equilibrium $H_2O(g) \rightleftharpoons H_2(g) + \frac{1}{2}O_2(g)$, the extent of dissociation of water when p = 1 atm and $K = 2.08 \times 10^{-3}$ is approximately
 - (a) 2%
- (b) 0.2%
- (c) 20%
- (d) 1%

36. In the van't Hoff equation

$$\frac{d \ln K}{dT} = \frac{\Delta H^{\circ}}{RT^2}$$

- (a) when $\frac{d \ln K}{dT}$ < 0, the reaction is exothermic
- (b) when $\frac{d\ln K}{dT}$ < 0, the reaction is endothermic
- (c) the slope of the graph is positive throughout
- (d) the slope of the graph increases and then decreases
- 37. When $-\ln K$ is plotted against $\frac{1}{T}$ using the van't Hoff equation, a straight line is expected with a slope equal to
 - (a) $\Delta H^{\circ}/RT$

(b) $-\Delta H^{\circ}/R$

(c) $\Delta H^{\circ}/R$

- (d) $R/\Delta H^{\circ}$
- **38.** In a gas-phase reaction $2A + B \rightleftharpoons 3C + 2D$ it was found that when 1.00 mol of A, 2.00 mol of B, and 1.00 mol of D were mixed and allowed to come to equilibrium at 25°C, the resulting mixture contained 0.90 mol of C at a total pressure of 1.00 bar. The value of K_v for the reaction is
 - (a) 6.86

(b) 4.86

(c) 68.6

(d) 10.86

- **39.** In the expression $-\frac{\Delta G}{T} = -\frac{\Delta H}{T} + \Delta S$
 - (a) when the reaction is exothermic, $-\frac{\Delta H}{T}$ corresponds to a positive change of entropy of the surroundings, and favours the formation of the products
 - (b) when the reaction is endothermic, $-\frac{\Delta H}{T}$ corresponds to a negative change of entropy of the surroundings, and favours the formation of the products
 - (c) when the reaction is exothermic, $-\frac{\Delta H}{T}$ corresponds to a negative change of entropy of the surroundings, and favours the formation of the reactants
 - (d) when the reaction is indothermic, $-\frac{\Delta H}{T}$ corresponds to a positive change of entropy of the surroundings, and favours the formation of the reactants
- **40.** Predict the effect of a tenfold increase in pressure on the equilibrium composition of the reaction $3N_2(g) + H_2(g) \rightleftharpoons 2NH_3(g)$.
 - (a) A thousandfold increase in K_x
 - (b) A hundredfold increase in K_x
 - (c) A tenfold decrease in K_x
 - (d) A hundredfold decrease in K_x
- **41.** On the basis of ΔG° , predict which of the following oxides can be reduced to the corresponding metal easily even in the absence of carbon.
 - (a) Ag₂O

(b) CuO

(c) Al_2O_3

- (d) Fe_2O_3
- 42. At constant pressure, the presence of inert gases
 - (a) reduces the dissociation of PCl₅
 - (b) increases the dissociation of PCl₅
 - (c) does not affect the degree of dissociation of PCl₅
 - (d) steps up the formation of PCl₅
- 43. At constant pressure, the addition of argon
 - (a) reduces the formation of ammonia from nitrogen and hydrogen
 - (b) increases the formation of ammonia from nitrogen and hydrogen
 - (c) does not affect the equilibrium of the reaction in which ammonia is formed from nitrogen and hydrogen
 - (d) reduces the dissociation of ammonia

- 44. Consider the reactions
 - (i) $PCl_5(g) \rightleftharpoons PCl_3(g) + Cl_2(g)$
 - (ii) $N_2O_4(g) \rightleftharpoons 2NO_2(g)$

The addition of an inert gas at constant volume

- (a) will increase the dissociation of PCl_5 as well as N_2O_4
- (b) will reduce the dissociation of PCl_5 as well as N_2O_4
- (c) will increase the dissociation of PCl₅ and step up the formation of NO₂
- (d) will not disturb the equilibrium of the reactions
- **45.** What is the effect of the reduction of the volume of the system for the equilibrium $2C(s) + O_2(g) \rightleftharpoons 2CO(g)$?
 - (a) The equilibrium will be shifted to the left by the increased pressure caused by the reduction in volume.
 - (b) The equilibrium will be shifted to the right by the decreased pressure caused by the reduction in volume.
 - (c) The equilibrium will be shifted to the left by the increased pressure caused by the increase in volume.
 - (d) The equilibrium will be shifted to the right by the increased pressure caused by the reduction in volume.
- **46.** In the following equilibrium

$$\frac{1}{2}$$
 N₂(g) + O₂(g) \rightleftharpoons NO₂(g) + heat

- (a) the equilibrium will shift to the right due to decrease in the concentration of $\ensuremath{N_2}$
- (b) the equilibrium will shift to the right due to an increase in temperature
- (c) the equilibrium will shift to the left due to an increase in volume
- (d) the equilibrium will shift to the left due to an increase in the concentration of ${\rm O}_2$
- **47.** Consider the following reaction.

$$C(S, diamond) \rightleftharpoons C(S, graphite) + heat$$

Choose the correct option.

- (a) An increase in temperature will shift the equilibrium to the right, and so will an increase in pressure.
- (b) An increase in temperature will shift the equilibrium to the left and so will an increase in pressure.
- (c) An increase in temperature will shift the equilibrium to the left and one in pressure to the right.
- (d) Any increase in temperature and pressure will not shift the equilibrium.

48. Consider the reaction

$$2CO(g) + O_2 \rightleftharpoons 2CO_2(g) + heat$$

Which of the following is incorrect?

- (a) The addition of CO and removal of CO₂ at constant volume will shift the equilibrium to the right.
- (b) The addition of O_2 and decrease in volume will shift the equilibrium to the right.
- (c) The addition of CO and increase in temperature at constant volume not determinable.
- (d) The addition of a catalyst and decrease in temperature will shift the equilibrium to the left.
- **49.** At 400 K, the following equilibrium is established.

$$H_2(g) + S(s) \rightleftharpoons H_2S(g); K_{eq.} = 7.0 \times 10^{-2}$$

If 0.1 mol of hydrogen and 1.0 mol of sulphur are heated to 400 K in a 1 L vessel, the partial pressure of H_2S at equilibrium is

- (a) 28 atm
- (b) 0.33 atm
- (c) 2.3 atm
- (d) 0.933 atm
- **50.** Which of the following equations is valid for a reversible process in a state of equilibrium?
 - (a) $\Delta G = -RT \ln K_n$

- (b) $\Delta G = RT \ln K_p$
- (c) $\Delta G^{\circ} = -RT \ln K_p$
- (d) $\Delta G^{\circ} = RT \ln K_p$
- **51.** Consider the following expression

$$\left(\frac{\partial \ln K_x}{\partial p}\right)_T = -\frac{\Delta n}{p/p^{\circ}}$$

and apply it to the formation of NH3 and indicate the correct statement.

- (a) K_x increases with increasing pressure
- (b) K_x increases with decreasing pressure
- (c) K_x decreases with increasing pressure
- (d) none of these

(K_x is the equilibrium constant in term of mole-fraction)

- **52.** Consider the expression $\triangle G = -RT \ln K_p + RT \ln Q_p$ and indicate the correct statement at equilibrium
 - (a) $\Delta G = 0$ $Q_p > K_p$ the equilibrium reaction will shift from left to right
 - (b) $\Delta G = 0$, $Q_p = K_p$ $Q_p > K_p$ the equilibrium reaction will shift from left to right
 - (c) $\Delta G = \infty$, $Q_p < K_p$ the equilibrium reaction will shift from right to left
 - (d) $\Delta G < 0$, $Q_p > K_p$ do -

where Q_p and K_p term refer to reaction quotient and equilibrium constant at constant pressure respectively.

- **53.** During the mixing of two ideal gases the Free Energy of mixing, ΔG_{mixing} is given by
 - (a) $\triangle G_{mixing} = nRT(x_A \ln x_A x_B \ln x_B)$
 - (b) $\triangle G_{mixing} = nRT(x_A \ln x_A + x_B \ln x_B)$
 - (c) $\triangle G_{mixing} = nRT(x_A + x_B) \ln x_A \cdot x_B$
 - (d) $\triangle G_{mixing} = nRTx_A \cdot x_B(\ln x_A + \ln x_B)$

(where x_A , x_B are the mole-fractions of the gases A and B respectively)

- **54.** A plot of the Gibbs energy of a reaction-mixture against the extent of the reaction is
 - (a) minimum at equilibrium
 - (b) zero at equilibrium
 - (c) equal to $\triangle H T \triangle S$ at equilibrium
 - (d) maximum at equilibrium
- **55.** In which of the following equilibrium, the equilibrium constant *K* is not greater than one?
 - (a) $HCl(g) + NH_3(g) \rightleftharpoons NH_4Cl(s)$
 - (b) $Fe(s) + H_2S(g) \rightleftharpoons FeS(s) + H_2(g)$
 - (c) $2H_2O_2(1) + H_2S(g) \rightleftharpoons H_2SO_4(1) + 2H_2(g)$
 - (d) $2Al_2O_3(s) + 3Si(s) \rightleftharpoons 3SiO_2(s) + 4Al(s)$
- **56.** By which of the following reactions, the equilibrium constant is related to temperature?

(a)
$$\ln K_2 - \ln K_1 = \frac{\Delta H^{\circ}}{R} \int_{T_1}^{T_2} d\left(\frac{1}{T}\right)$$

(b)
$$\ln K_2 - \ln K_1 = -\frac{\Delta H^{\circ}}{R} \int_{1/T_1}^{1/T_2} d\left(\frac{1}{T^2}\right)$$

(c)
$$\ln K_2 - \ln K_1 = -\frac{\Delta H^{\circ}}{R} \int_{T_1}^{T_2} d\left(\frac{1}{T}\right)$$

(d)
$$\ln K_2 - \ln K_1 = -\frac{\Delta H^{\circ}}{R} \int_{1/T_2}^{1/T_1} d\left(\frac{1}{T}\right)$$

- 57. Which of the following expressions is incorrect?
 - (a) $\left(\frac{\partial \ln K_p}{\partial p}\right)_T = 0$

(b)
$$\left(\frac{\partial \ln K_c}{\partial p}\right)_T = 0$$

(c)
$$\left(\frac{\partial \ln K_x}{\partial p}\right)_T = -\frac{\Delta n}{(p/p^\circ)}$$

(d) All of these

• *Type 2* •

Choose the correct options. More than one option is correct.

- 58. Which of the following statements are incorrect?
 - (a) For a closed system, *S* is always maximum at equilibrium.
 - (b) The addition of a reactant gas to an ideal-gas reaction mixture shifts the equilibrium in such a way that some of the added gas is used up.
 - (c) In any closed system, *G* is always minimum at equilibrium.
 - (d) In the limit $T \to 0$, ΔG° approaches ΔH° .
- **59.** The expression for the equilibrium constant for an ideal-gas reaction mixture is given by

(a)
$$\frac{d \ln K_c^0}{dT} = \frac{\Delta U^0}{RT^2}$$

(b)
$$\left(\frac{\partial \ln K_x^0}{\partial T}\right)_p = \frac{\Delta S^0}{RT^2}$$

(c)
$$\left(\frac{\partial \ln K_x^0}{\partial p}\right)_T = -\frac{\Delta n/\text{mol}}{p}$$

(d)
$$\frac{d \ln K_p^0}{dT} = \frac{\Delta H^0}{RT^2}$$

60. The condition for spontaneity in a chemical reaction is

(a)
$$(\Delta G)_{T,p} \leq 0$$

(b)
$$(\Delta U)_{S,V} \leq 0$$

(c)
$$(\Delta H)_{S, p} \leq 0$$

(d)
$$(\Delta S)_{U,V} \ge 0$$

61. In which of the following reactions is $K_p < K_c$?

(a)
$$CO(g) + Cl_2(g) \rightleftharpoons COCl_2(g)$$

(b)
$$CO(g) + 3H_2(g) \rightleftharpoons CH_4(g) + H_2O(g)$$

(c)
$$2BrCl(g) \rightleftharpoons Cl_2(g) + Br_2(g)$$

- (d) $I_2(g) \rightleftharpoons 2I(g)$
- **62.** The dissociation of phosgene, which occurs according to the reaction $COCl_2(g) \rightleftharpoons CO(g) + Cl_2(g)$

is an endothermic process. Which of the following will increase the degree of dissociation of COCl₂?

- (a) Adding Cl₂ to the system
- (b) Adding helium to the system
- (c) Decreasing the temperature of the system
- (d) Reducing the total pressure
- **63.** The equilibrium of which of the following reactions will not be disturbed by the addition of an inert gas at constant volume?

(a)
$$H_2(g) + I_2(g) \rightleftharpoons 2HI(g)$$

(b)
$$N_2O_4(g) \rightleftharpoons 2NO_2(g)$$

- (c) $CO(g) + 2H_2(g) \rightleftharpoons CH_3OH(g)$
- (d) $C(s) + H_2O(g) \rightleftharpoons CO(g) + H_2(g)$
- **64.** A box contains CO(g), Cl₂(g) and COCl₂(g) in equilibrium at 1000 K. The removal of CO(g) will
 - (a) decrease the concentration of COCl₂
 - (b) increase the concentration of Cl₂
 - (c) increase the concentration of COCl₂
 - (d) reduce the concentration of CO as well as Cl₂
- **65.** An industrial fuel, 'water gas', which consists of a mixture of H₂ and CO can be made by passing steam over red-hot carbon. The reaction is

$$C(s) + H_2O(g) \rightleftharpoons CO(g) + H_2(g), \Delta H = +131 \text{ kJ}$$

The yield of CO and H_2 at equilibrium would be shifted to the product side by

- (a) raising the relative pressure of the steam
- (b) adding hot carbon
- (c) raising the temperature
- (d) reducing the volume of the system
- **66.** For the equilibrium $2SO_2(g) + O_2(g) \rightleftharpoons 2SO_3(g)$, $\Delta H = -198$ kJ, the equilibrium concentration of SO_3 will be affected by
 - (a) doubling the volume of the reaction vessel
 - (b) increasing the temperature at constant volume
 - (c) adding more oxygen to the reaction vessel
 - (d) adding helium to the reaction vessel at constant volume
- **67.** The following reaction attains equilibrium at high temperature.

$$N_2(g) + 2H_2O(g) + heat \rightleftharpoons 2NO(g) + 2H_2(g)$$

The yield of NO is affected by

- (a) increasing the nitrogen concentration
- (b) decreasing the hydrogen concentration
- (c) compressing the reaction mixture
- (d) none of these

68.
$$N_2(g) + 3H_2(g) \xrightarrow{\text{catalyst}} 2NH_3 + \text{heat}$$

In this reaction, the direction of equilibrium will be shifted to the right by

- (a) increasing the concentration of nitrogen
- (b) compressing the reaction mixture
- (c) removing the catalyst
- (d) decreasing the concentration of ammonia

69. The dissociation of ammonium carbamate may be represented by the equation

$$NH_4CO_2NH_2(s) \rightleftharpoons 2NH_3(g) + CO_2(g)$$

 ΔH° for the forward reaction is negative. The equilibrium will shift from right to left if there is

- (a) a decrease in pressure
- (b) an increase in temperature
- (c) an increase in the concentration of ammonia
- (d) an increase in the concentration of carbon dioxide

		Answers	<u> </u>	
1. d	2. c	3. b	4. b	5. d
6. c	7. d	8. b	9. d	10. d
11. d	12. c	13. c	14. a	15. b
16. c	17. d	18. c	19. c	20. a
21. b	22. c	23. b	24. a	25. d
26. a	27. b	28. d	29. b	30. c
31. d	32. d	33. d	34. b	35. a
36. a	37. c	38. a	39. a	40. b
41. a	42. b	43. a	44. d	45. a
46. c	47. b	48. d	49. c	50. c
51. a	52. b	53. b	54. a	55. d
56. c	57. d	58. a, b, c	59. a, c, d	60. a, b, c, d
61. a, b	62. b, d	63. a, b, c, d	64. a, b	65. a, c
66. a, b, c	67. a, b, c	68. a, b, d	69. b, c, d	

Hints to More Difficult Problems

2. For the reaction $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$, $\Delta n = -2$.

$$K_p = K_c (RT)^{\Delta n} = K_c (RT)^{-2}$$

$$\Rightarrow K_c = \frac{K_p}{(RT)^{-2}}.$$

5. Using the equation

$$\log \frac{(K_p)_{40^{\circ}\text{C}}}{(K_p)_{25^{\circ}\text{C}}} = \frac{\Delta H}{2.303R} \left(\frac{1}{T_1} - \frac{1}{T_2} \right),$$
we get $\log 4 = \frac{\Delta H}{2.303 \times 8.314} \left(\frac{1}{273 + 25} - \frac{1}{273 + 40} \right).$

$$\therefore \Delta H = 71.67 \text{ kJ mol}^{-1}.$$

- 6. Apply the Le Chatelier principle.
- 9. A(g) + B(g) \rightleftharpoons 2C(g) a 3 0 a-x 3-x 2x From the question, $2x = 3 \Rightarrow x = 1.5$

$$3 = \frac{(2x)^2}{(a-x)(3-x)}$$

Substituting x = 1.5, we get a = 3.5.

10. AB(g)
$$\rightleftharpoons$$
 A(g) + B(g)
$$\begin{array}{cccc}
1 & 0 & 0 \\
-\frac{1}{3} & +\frac{1}{3} & \frac{1}{3} \\
\frac{2}{3} & \frac{1}{3} & \frac{1}{3}
\end{array}$$

$$(\Sigma n)_{\text{eqlm}} = \frac{2}{3} + \frac{1}{3} + \frac{1}{3} = \frac{4}{3} \cdot K_p = \frac{p_{\text{A}}p_{\text{B}}}{p_{\text{AB}}} = \frac{\frac{1}{3}}{\frac{2}{3}} p^{\frac{1}{3}} \frac{p^{\frac{1}{3}}}{\frac{4}{3}} p = \frac{1}{8} p.$$

$$\therefore p = 8K_p.$$

15.
$$NH_4COONH_2(s) \Rightarrow 2NH_3(g) + CO_2(g)$$

$$\frac{2}{3}p \qquad \frac{1}{3}p$$

$$K_p = p_{NH_3}^2 p_{CO_2} = \left(\frac{2}{3}p\right)^2 \left(\frac{1}{3}p\right) = \frac{4}{27}p^3 = \frac{4}{27} \times (2)^3 = 1.185.$$

- 17. Use the van't Hoff equation.
- **20.** Use the formula $\Delta G^{\circ} = -2.303 \ RT \log K_p$ and assume that $T = 298.15 \ K$. For questions 22, 23 and 24 use the Le Chatelier principle.

25.
$$\alpha = \frac{M_{\text{Th}} - M_{\text{obs}}}{M_{\text{obs}}(n-1)}$$

Molar mass of $N_2O_4 = 92$ g mol⁻¹.

Here, n = 2.

$$\alpha = \frac{92.00 - 77.70}{77.70(2 - 1)} = 0.184 = 18.4\%.$$

31. Use the van't Hoff equation

$$\frac{d \ln K_p}{dT} = \frac{\Delta H^{\circ}}{RT^2} \quad \text{or } \ln K_p = -\frac{\Delta H}{R} \frac{1}{T} + C.$$

35. For the equilibrium $H_2O(g) \longrightarrow H_2(g) + \frac{1}{2}O_2(g)$

$$K_p = \frac{\alpha^{3/2} p^{1/2}}{\sqrt{2}} p = 1$$
 atm.

$$\alpha = (\sqrt{2} \cdot k_p)^{\frac{2}{3}} = 0.0205 \approx 2\%.$$

40. For the equilibrium $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$,

$$K_p = \frac{K_x}{p^2}, \ p = 10.$$

$$K_x = K_p(10)^2 = 100K_p$$
.

Questions 42, 43 and 44 can be answered using the equation

$$K_p = \frac{n_{\text{C}}^c \cdot n_{\text{D}}^d}{n_{\text{A}}^a \cdot n_{\text{B}}^b} \left(\frac{p}{\Sigma n}\right)^{\Delta n}$$

At constant volume the equilibrium will not be disturbed.

54.
$$\left(\frac{\partial G_x}{\partial G_{Eq}}\right)_{p, T} = \Delta G$$
 (minimum)

- 56. Use Van't Hoff equation
- **57.** K_v , K_c are independent of pressure but K_x is related to pressure
- **61.** For reactions (a) and (b), $\Delta n < 1$. Questions 57 to 62 may be answered by applying the Le Chatelier principle.

Chemical Kinetics

• Type 1 •

Choose the correct option. Only one option is correct.

1. The decomposition of nitrogen pentoxide can be represented as

$$N_2O_5(g) \longrightarrow 2NO_2(g) + \frac{1}{2}O_2(g)$$

The rate of the reaction can be expressed as

(a)
$$-\frac{d[N_2O_5]}{dt} = 2\frac{d[NO_2]}{dt} = \frac{1}{2}\frac{d[O_2]}{dt} = k[N_2O_5]$$

(b)
$$\frac{1}{2} \frac{d[N_2O_5]}{dt} = \frac{1}{2} \frac{d[NO_2]}{dt} = \frac{d[O_2]}{dt} = k[N_2O_5]$$

(c)
$$-\frac{d[N_2O_5]}{dt} = \frac{1}{2}\frac{d[NO_2]}{dt} = 2\frac{d[O_2]}{dt} = k[N_2O_5]$$

(d)
$$-\frac{d[N_2O_5]}{dt} = \frac{1}{2}\frac{d[NO_2]}{dt} = \frac{1}{2}\frac{d[O_2]}{dt} = k[N_2O_5]$$

2. The decomposition of azomethane $(C_2H_6N_2)$ at a certain temperature in the gas phase follows first-order kinetics. The following are some data for the reaction $C_2H_6N_2(g) \rightarrow C_2H_6(g) + N_2(g)$.

Time (min) 0 15 30 48 75 $[C_2H_6N_2]$ (M) 0.36 0.30 0.25 0.19 0.13

The value of the rate constant is

(a) $0.007 \, \text{min}^{-1}$

(b) $0.014 \, \text{min}^{-1}$

(c) $0.042 \, \text{min}^{-1}$

(d) $0.028 \, \text{min}^{-1}$

3. The instantaneous rate of disappearance of the MnO_4^- ion in the following reaction is $4.56\times10^{-3}\,M$ s $^{-1}$.

$$2MnO_4^- + 10I^- + 16H^+ \longrightarrow 2Mn^{2+} + 5I_2 + 8H_2O$$

The rate of appearance of I₂ is

- (a) $1.14 \times 10^{-3} \,\mathrm{M \, s}^{-1}$
- (b) $5.7 \times 10^{-3} \,\mathrm{M s^{-1}}$
- (c) $4.56 \times 10^{-4} \,\mathrm{M \, s^{-1}}$
- (d) $1.14 \times 10^{-2} \,\mathrm{M \, s^{-1}}$
- **4.** For a reaction $I^- + OCl^- \rightarrow IO^- + Cl^-$, in an aqueous medium, the rate of the reaction is given by

$$\frac{d[IO^-]}{dt} = k \frac{[I^-][OC1^-]}{[OH^-]}$$

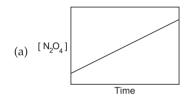
The overall order of the reaction is

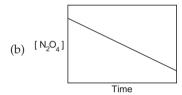
(a) -1

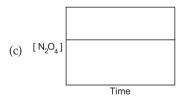
(b) 1

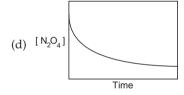
(c) zero

- (d) 2
- 5. Which of the following graphs best describes the rate at which N_2O_4 decomposes to NO_2 if the reaction is first-order in N_2O_4 ?









6. The molecularity of the reaction

$$2H_2(g) + O_2(g) \longrightarrow 2H_2O(l)$$

is

- (a) 2
- (b) 3
- (c) undefined (d) 1
- (c) underined (d) 1
- 7. The unit of the rate of reaction is the same as that of the rate constant for a
 - (a) zero-order reaction
- (b) first-order reaction
- (c) second-order reaction
- (d) half-order reaction
- 8. The rate constant for a zero-order reaction is
 - (a) $k = \frac{c_0}{2t}$

(b) $k = \frac{c_0 - c_t}{t}$

(c) $k = \ln \frac{c_0 - c_t}{2t}$

(d) $k = \frac{c_0}{c_t}$

9. The half-life periods of decomposition of PH3 for different initial

37.5

84

(c) $\frac{1}{2}$

(d) 2

79

84

pressures are given below.

The order of the reaction is

707

84

10. The half-life for a zero-order reaction equals

(b) 0

p(torr.)

 $t_{1/2}(\min.)$

(a) 1

	(a)	$\frac{1}{2} \frac{k}{a^2}$	(b)	$\frac{a^2}{2k}$	(c)	$\frac{2k}{a}$	(d)	$\frac{a}{2k}$
	where	<i>a</i> is the initia	al cor	ncentration.				
11.	doubl		conce	entration of A				of the reaction s. The order of
	(a)	0	(b)	$\frac{1}{2}$	(c)	1	(d)	2
12.		ate constant to on when the i						e initial rate of
	(a)	100 M			(b)	$1 \times 10^{-2} \mathrm{M}$		
	(c)	1.0 M			(d)	0.1 M		
13.		ntration is i						nen the initial order of the
	(a)	1	(b)	4	(c)	0	(d)	2
14.	The sa	aponification	of et	hyl acetate is	a			
	(a)	zero-order			(b)	half-order		
	(c)	second-orde	er		(d)	third-order		
	reaction	on.						
15.		he rate beco						rease fourfold order of the
	(a)	2.0	(b)	3.5	(c)	2.5	(d)	1.5
16.	What	ate expression changes in the ction to incres	ne in	itial concentr	atior	ns of A and B	g) is will	rate = $kC_A^2C_B^{1/2}$. cause the rate
	(a)	$C_{\rm A} \times 2$; $C_{\rm B} \times$	2		(b)	$C_{\rm A} \times 2$; $C_{\rm B} \times$	4	
	(c)	$C_{\rm A} \times 1$; $C_{\rm B} \times$	4		(d)	$C_{\rm A} \times 4$; $C_{\rm B} \times$	1	

17. Calculate the half-life of the first-order reaction

$$C_2H_4O(g) \longrightarrow CH_4(g) + CO(g)$$

if the initial pressure of $C_2H_4O(g)$ is 80 mm and the total pressure at the end of 20 minutes is 120 mm.

(a) 40 min

(b) 120 min

(c) 20 min

- (d) 80 min
- **18.** In a particular reaction the time required to complete half of the reaction was found to increase 16 times when the initial concentration of the reactant was reduced to one-fourth. What is the order of the reaction?
 - (a) 1

(b) 4

(c) 2

- (d) 3
- **19.** For a first-order reaction, the time required for 99.9% of the reaction to take place is nearly
 - (a) 10 times that required for half the reaction
 - (b) 100 times that required for two-thirds of the reaction
 - (c) 10 times that required for one-fourth of the reaction
 - (d) 20 times that required for half of the reaction
- **20.** The hydrolysis of ethyl acetate was carried out separately with 0.075 M HCl and 0.075 M H₂SO₄. Which of the following relations is correct?
 - (a) $k_{\text{H}_2\text{SO}_4} >> k_{\text{HCl}}$

(b) $k_{H,SO_4} = 2k_{HCl}$

(c) $k_{\text{HCl}} > k_{\text{H,SO}_4}$

- (d) $k_{H_2SO_4} = 8k_{HCl}$
- **21.** For a reaction following first-order kinetics, which of the following statements are correct?
 - (a) The time taken for the completion of 50% of the reaction is twice $t_{\frac{1}{2}}$.
 - (b) A plot of the reciprocal of the concentration of the reactants against time gives a straight line.
 - (c) The degree of dissociation is equal to $1 e^{-kt}$.
 - (d) A plot of $[A]_0/[A]$ versus time gives a straight line.
- 22. The reaction of NO with Cl_2 follows the equation

$$2NO(g) + Cl_2(g) \longrightarrow 2NOCl$$

The following data were collected.

Initial concentration of NO $\pmod{L^{-1}}$	Initial concentration of $Cl_2(\text{mol L}^{-1})$	Initial rate of formation of NOCl $(\text{mol L}^{-1} \text{ s}^{-1})$
0.10	0.10	2.53×10^{-6}
0.10	0.20	5.06×10^{-6}
0.20	0.10	10.10×10^{-6}
0.30	0.10	22.80×10^{-6}

The value of the rate constant is

(a) 2.53×10^{-6}

(b) 5.06×10^{-6}

(c) 10.10×10^{-6}

(d) 22.80×10^{-6}

- $L^2 \text{ mol}^{-2} \text{ s}^{-1}$.
- 23. For the reaction

$$CH_3COCH_3 + Br_2 \xrightarrow{H^+} CH_3COCH_2Br + H^+ + Br^-$$

the following data were collected.

[CH ₃ COCH ₃]	$[Br_2]$	$[H^+]$	Rate of disappearance of Br_2 (M s ⁻¹)
0.15	0.025	0.025	6.0×10^{-4}
0.15	0.050	0.025	6.0×10^{-4}
0.15	0.025	0.050	1.2×10^{-3}
0.20	0.025	0.100	3.2×10^{-3}
0.20	0.025	0.025	8.0×10^{-3}

The overall order of the reaction is

(a) 1.5

(b) 4

(c) 5

- (d) 2
- **24.** Consider the reaction X + Y → products. If the initial concentration of X is increased to four times its original value, keeping the concentration of Y constant, the rate of reaction increases fourfold. When the concentrations of both X and Y become four times their original values, the rate of reaction becomes sixteen times its original value. The observed rate law is
 - (a) $k[X]^2[Y]^2$

(b) $k[X]^{1}[Y]^{2}$

(c) $k[X]^{1}[Y]^{1}$

- (d) $k[X]^2[Y]^1$
- **25.** An endothermic reaction has a positive internal energy change ΔU . In such a case, what is the minimum value that the activation energy can have?
 - (a) ΔU

- (b) $\Delta U = \Delta H + \Delta nRT$
- (c) $\Delta U = \Delta H \Delta nRT$
- (d) $\Delta U = E_a + RT$
- 26. The activation energy of a reaction may be lowered by
 - (a) decreasing the temperature
 - (b) increasing the temperature
 - (c) adding a catalyst
 - (d) reducing the potential energy

- **27.** In an exothermic reaction $X \to Y$, the activation energy is 100 kJ mol^{-1} of X. The enthalpy of the reaction is -140 kJ mol^{-1} . The activation energy of the reverse reaction $Y \to X$ is
 - (a) 40 kJ mol⁻¹

(b) 340 kJ mol⁻¹

(c) 240 kJ mol⁻¹

- (d) 100 kJ mol⁻¹
- **28.** The rate constant, the activation energy and the pre-exponential factor of a chemical reaction at 25°C are 8.0×10^{-4} s⁻¹, 112 kJ mol⁻¹ and 4×10^{15} s⁻¹ respectively. The value of the rate constant as $T \rightarrow \infty$ is
 - (a) $8 \times 10^{16} \,\mathrm{s}^{-1}$

(b) $4 \times 10^4 \,\mathrm{s}^{-1}$

(c) $4 \times 10^{15} \,\mathrm{s}^{-1}$

- (d) $112 \times 10^{12} \,\mathrm{s}^{-1}$
- **29.** Which of the following is correct?
 - (a) Total collision rate ∞ mean speed ∞ absolute temperature
 - (b) Total collision rate $\propto \frac{1}{\text{mean speed}} \propto \frac{1}{\text{absolute temperature}}$
 - (c) Total collision rate \propto mean speed \propto (absolute temperature) $^{1/2}$
 - (d) Total collision rate \propto (mean speed) $^2 \propto$ (absolute temperature) 3
- **30.** It is often stated that, near room temperature, a reaction rate doubles if the temperature increases by 10°C. Calculate the activation energy of a reaction that obeys this rule exactly.
 - (a) 12.4 kcal

(b) 24.8 kcal

(c) 6.2 kcal

- (d) 49.6 kcal
- **31.** Which of the following statements is correct for the activation energy of a reaction?
 - (a) It increases with increase in temperature.
 - (b) When the activation energy is zero the rate constant is temperature-dependent.
 - (c) It decreases with decrease in temperature.
 - (d) It is nearly independent of temperature, over a wide range.
- 32. In the reaction

$$S + C \xrightarrow{K_2} X$$

$$X + R \xrightarrow{K'_2} products + C$$

where S = substrates, C = catalyst, X = intermediate complex and R = other substrates.

The rate of formation of products is given by $v = K_2' = [X]$ [R]. The rate of accumulation of X is given by

(a)
$$\frac{d[C]}{dt} = K'[C][S]$$

(b)
$$\frac{d[R]}{dt} = K'[X][R]$$

(c)
$$\frac{d[X]}{dt} = 0$$

(d)
$$\frac{d[X]}{dt} \ge 1$$

- **33.** When the activation energies of the forward and reverse reactions are equal, then
 - (a) $\Delta U = 0$, $\Delta S = 0$

- (b) $\Delta U = \infty$, $\Delta S = 0$
- (c) $\Delta G = 0$, $\Delta U = 0$
- (d) only $\Delta U = 0$
- **34.** A drop of a solution (volume = 0.05 mL) contains 6×10^{-7} mol of H⁺. If the rate of disappearance of H⁺ is 6.0×10^{5} mol L⁻¹ s⁻¹, how long will it take for the H⁺ in the drop to disappear?
 - (a) 8.0×10^{-8} s

(b) 2.0×10^{-8} s

(c) 6.0×10^{-6} s

- (d) 2.0×10^{-2} s
- 35. The reaction $2A \longrightarrow B$ is first-order in A with a rate constant of $2.8 \times 10^{-2} \, s^{-1}$. How long will it take for A to decrease from 0.88 M to 0.14 M?
 - (a) $50 \, s$

(b) 76 s

(c) 66 s

- (d) 44 s
- 36. Inversion of sucrose occurs under
 - (a) general acid-base catalysis
- (b) general acid catalysis
- (c) general base catalysis
- (d) specific acid catalysis
- **37.** In the Arrhenius equation $k = A \exp(-E_A/RT)$, the rate constant
 - (a) decreases with increasing activation energy and increases with temperatue
 - (b) increases with activation energy and temperature
 - (c) decreases with activation energy and temperature
 - (d) increases with activation energy and decreasing temperature
- **38.** Nickel as a catalyst is said to be geometrically effective when ethylene is adsorbed on it because
 - (a) the closest distance between the nickel atoms is greater than the C—C distance in ethylene
 - (b) the closest distance between the nickel atoms is less than the C—C distance in ethylene
 - (c) the closest distance between the nickel atoms is equal to the C—C distance in ethylene
 - (d) the closest distance between the nickel atoms is less than that between the nickel and carbon atoms after adsorption

39. The rate of the reaction $A \longrightarrow B$ is given by

(a)
$$-\frac{1}{V}\frac{dn_{A}}{dt} = \frac{1}{V}\frac{dn_{B}}{dt}$$

(b)
$$\frac{p}{RT} \frac{dn_A}{dt} = -\frac{p}{RT} \frac{dn_B}{dt}$$

(c)
$$-\frac{1}{[A]}\frac{dn_{A}}{dV} = +\frac{1}{[B]}\frac{dn_{B}}{dV}$$

(d)
$$-\frac{dV_{\rm A}}{dt} = +\frac{dV_{\rm B}}{dt}$$

where n_A and n_B are the number of moles of A and B and V_A and V_B are their volumes, respectively.

40. Consider the dehydrogenation of 2-propanol,

$$CH_3CHOHCH_3(l) \longrightarrow CH_3COCH_3(l) + H_2(g)$$

The following data were obtained about this reaction. (Assume the volume of the reaction mixture to be 1 dm³.)

Time (min) 0 5 10 15 [Acetone] mol
$$L^{-1}$$
 0 0.3 0.6 0.9

The initial rate of the reaction was

- (a) $0.03 \text{ mol dm}^{-3} \text{ min}^{-1}$
- (b) $0.06 \text{ mol dm}^{-3} \text{ min}^{-1}$
- (c) $0.09 \text{ mol dm}^{-3} \text{ min}^{-1}$
- (d) $0.12 \text{ mol dm}^{-3} \text{ min}^{-1}$

41. The decomposition of nitrogen pentoxide is a first-order reaction

$$\left(N_2O_2 \longrightarrow 2NO_2 + \frac{1}{2}O_2\right) \cdot$$

The rate of this reaction is given by

$$\frac{-dc_{\text{N}_2\text{O}_5}}{dt} = \frac{1}{2} \frac{dc_{\text{NO}_2}}{dt} = 2 \frac{dc_{\text{O}_2}}{dt} = K_1 c_{\text{N}_2\text{O}_5}$$

Then

$$\begin{split} &-\frac{dc_{\mathrm{N_2O_5}}}{dt} = k_1 c_{\mathrm{N_2O_5}} = k_1 c_{\mathrm{N_2O_5}} \\ &-\frac{dc_{\mathrm{NO_2}}}{dt} = 2k_1 c_{\mathrm{N_2O_5}} = k_2' c_{\mathrm{N_2O_5}} \\ &-\frac{dc_{\mathrm{O_2}}}{dt} = \frac{1}{2} \; k_1 c_{\mathrm{N_2O_5}} = k_1'' c_{\mathrm{N_2O_5}} \end{split}$$

Choose the correct option.

(a)
$$k_1 = 2k_1' = k_1''$$

(b)
$$k_1 = k_1' = k_1''$$

(c)
$$4k_1 = 2k'_1 = k''_1$$

(d)
$$4k_1 = k' = 2k_1''$$

42. The rate expression for the second-order reaction $A + B \longrightarrow products$ is given by

 $K_2 t = \frac{1}{a - b} \ln \frac{b(a - x)}{a(b - x)}$

Bear in mind the following data.

$$t = 0 \qquad a \qquad b$$

$$t = t \qquad a - x \qquad b - x$$

43. A sample of ammonia decomposes on a tungsten wire at 1135 K. At

221

100

constant volume, the following results were obtained.

210

(b) third-order

(d) first-order

232

200

(b) 0.33 torr s^{-1}

255

400

277

600

where a =concentration of A and b = concentration of B.

zero-order (c) half-order

Pressure (torr)

(a) 0.22 torr s^{-1}

Time (s)

When $a \gg b$, this expression will become

The rate constant for the reaction is

	(c)	0.11 torr s^{-1}	(d)	0.44 torr s^{-1}
44.	(a) (b) (c)	nit of the rate constant depend temperature of the reaction activation energy of the reacti molecularity of the reaction order of the reaction		on the
45.	1.0 × 1 pondi	10^{-2} dm ³ mol ⁻¹ s ⁻¹ and $3.0 \times$ ing energies of activation of th 0.0 kJ mol^{-1} respectively, what	10 ⁻² e pai	reactions were found to be dm ³ mol ⁻¹ s ⁻¹ . If the corres- rallel reactions are 60.0 kJ mol ⁻¹ the apparent overall energy of
	(a)	130.0 kJ mol ⁻¹	(b)	67.5 kJ mol ⁻¹
	(c)	$100.0 \text{ kJ mol}^{-1}$	(d)	65.0 kJ mol^{-1}
46.	56.0 k			gaseous compound initially at $8.0~\mathrm{kP_{a'}}$ the half-life is $170~\mathrm{s}$. The
	(a)	0	(b)	2
	(c)	1	(d)	$\frac{1}{2}$
47.	Consi	der the reaction mechanism		
		$A_2 \rightleftharpoons 2A$	fast)	
		$A + B \longrightarrow P$	(slov	w)
	where	e A is the intermediate. The rat		
	(a)	$k_2[A][B]$	(b)	$k_2 k^{\frac{1}{2}} [A_2]^{\frac{1}{2}} [B]$
	(c)	$k_2 k^{\frac{1}{2}} [A][B]$	(d)	$k_2 k^{\frac{1}{2}} [A]^2 [B]$

48. Consider the following reactions at 300 K.

 $A \longrightarrow B$ (uncatalysed reaction)

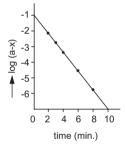
$$A \xrightarrow{\text{catalyst}} B \text{ (catalysed reaction)}$$

The activation energy is lowered by 8.314 kJ mol⁻¹ for the catalysed reaction. The rate of this reaction is

- (d) 28 times (a) 15 times (c) 22 times (b) 38 times that of the uncatalyzed reaction.
- 49. Aqueous NH₄NO₂ decomposes according to the first-order reaction $NH_4NO_2(aq) \longrightarrow N_2(g) + 2H_2O(l)$

After 20 minutes the volume of N₂ collected during such a reaction is 20 mL, and that collected after a very long time is 40 mL. The rate constant for the reaction is

- (a) 1.435×10^{-2} min⁻¹
- (b) $3.466 \times 10^{-2} \text{ min}^{-1}$
- (c) $3.465 \times 10^{-2} \text{ min}^{-1}$
- (d) 6.93 min⁻¹
- 50. The conversion of vinyl allyl ether to pent-4-enol follows first-order kinetics. The following plot is obtained for such a reaction.



The rate constant for the reaction is

(a) $4.6 \times 10^{-2} \text{ s}^{-1}$

(b) $1.2 \times 10^{-2} \text{ s}^{-1}$

(c) $2.3 \times 10^{-2} \text{ s}^{-1}$

- (d) $8.4 \times 10^{-2} \text{ s}^{-1}$
- 51. $N_2O_2(g) \longrightarrow 2NO$ is a first-order reaction in terms of the concentration of $N_2O_2(g)$. Which of the following is valid, $[N_2O_2]$ being constant?

 - (a) $[NO] = [N_2O_2]_0 e^{-kt}$ (b) $[NO] = [N_2O_2]_0 (1 e^{-kt})$ (c) $[NO] = [N_2O_2]_0 (e^{-kt} 1)$ (d) $[NO] = [N_2O_2]_0 (1 e^{-kt})$
- **52.** The reaction $A(g) + 2B(g) \rightarrow C(g) + D(g)$ is an elementary process. In an experiment involving this reaction, the initial partial pressures of A and B are $p_A = 0.60$ atm and $p_B = 0.80$ atm respectively. When $p_C = 0.20$ atm, the rate of the reaction relative to the initial rate is
 - (a) $\frac{1}{6}$
- (b) $\frac{1}{12}$ (c) $\frac{1}{36}$
 - (d) $\frac{1}{18}$

- 53. Which of the following statements is correct?
 - (a) The order of a reaction must be a positive integer.
 - (b) A second-order reaction is also bimolecular.
 - (c) The order of a reaction increases with temperature.
 - (d) All bimolecular reactions are of the second order.
- **54.** For the reaction A + B \longrightarrow C the rate constant for the second-order forward reaction is $k_2 = 10.00$ exp. $\left(-\frac{90500}{RT}\right)$ dm³ mol⁻¹ s⁻¹. The preexponential factor and activation energy are, respectively,
 - (a) 10^{10} dm³ mol⁻¹ s⁻¹ and -90.50 kJ mol⁻¹
 - (b) $\log 10 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1} \text{ and } -45.25 \text{ kJ mol}^{-1}$
 - (c) $10 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1} \text{ and } 90.50 \text{ kJ mol}^{-1}$
 - (d) $10 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1} \text{ and } -90.25 \text{ kJ mol}^{-1}$
- **55.** On thermal decomposition, 2-nitropropane yields propylene, and the rate constant is given by the relationship

$$k = 1.11 \times 10^{11} \text{ exp.} \left(-\frac{164438}{8.314T} \right) \text{s}^{-1}.$$

What is the half-life of this reaction at 300°C?

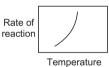
(a) 5964 s

(b) 6964 s

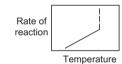
(c) 9654 s

(d) 4964 s

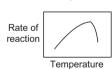
56. (I)



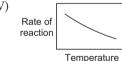
(II)



(III)



(IV)



Which of these graphs represents an enzyme reaction?

- (a) II
- (b) III
- (c) I
- (d) IV
- 57. What happens when the temperature of a solution is increased from 25°C to 65°C?
 - (a) The rate of the reaction remains unchanged and the rate constant *k* decreases.
 - (b) The rate of the reaction increases and the rate constant k decreases.
 - (c) The rate of the reaction decreases and so does the rate constant k.
 - (d) The rate of the reaction increases and so does the rate constant k.

58. The rate law for the reaction

$$2H_2(g) + 2NO(g) \longrightarrow N_2(g) + H_2O(g)$$

is
$$\frac{d[N_2]}{dt} = k[H_2][NO]^2$$
.

Which of the following mechanisms is consistent with the rate law?

(a)
$$H_2(g) + 2NO(g) \xrightarrow{k_1} N_2O(g) + H_2O(g)$$

(b)
$$H_2(g) + N_2O(g) \xrightarrow{k_2} N_2(g) + H_2O(g)$$

(c)
$$H_2O(g) + NO(g) \xrightarrow{k_3} H_2(g) + NO_2(g)$$

$$(d) \ \ H_2(g) + 2NO(g) \xrightarrow{\quad k_4 \quad } \ N_2(g) + H_2O(g) + \frac{1}{2} \, O_2(g)$$

59. In the decomposition of $N_2O_5(g)$, i.e.,

$$2N_2O_5(g) \xrightarrow{k_{obs}} 4NO_2(g) + O_2(g)$$

the observed rate law is given by

$$\frac{d[O_2]}{dt} = k_{\text{obs}}[N_2O_5].$$

Which of the following proposed mechanisms is consistent with the rate law?

(a)
$$N_2O_5(g) \xrightarrow{k_1} N_2O_3(g) + O_2(g)$$

(b)
$$N_2O_5(g) \xrightarrow{k_2} NO_2(g) + NO_3(g)$$

$$(c) \quad NO_2(g) + NO_3(g) \xrightarrow{\quad k_3 \quad} NO(g) + NO_2(g) + O_2(g)$$

(d)
$$NO_3(g) + NO(g) \xrightarrow{k_4} 2NO_2(g)$$

• Type 2 •

Choose the correct options. More than one option is correct.

- **60.** Which of the following statements are correct?
 - (a) The order of a reaction is the sum of the components of all the concentration terms in the rate equation.
 - (b) The order of a reaction with respect to one reactant is the ratio of the change of logarithm of the rate of the reaction to the change in the logarithm of the concentration of the particular reactant, keeping the concentrations of all other reactants constant.
 - (c) Orders of reactions can be whole numbers or fractional numbers.

- (d) The order of a reaction can only be determined from the stoichiometric equation for the reaction.
- **61.** Which of the following statements are correct?
 - (a) The rate of the reaction involving the conversion of ortho-hydrogen to parahydrogen is $-\frac{d[H_2]}{dt} = k[H_2]^{3/2}$.
 - (b) The rate of the reaction involving the thermal decomposition of acetaldehyde is k[CH₃CHO] ^{3/2}.
 - (c) In the formation of phosgene gas from CO and Cl_2 , the rate of the reaction is $k[CO][Cl_2]^{1/2}$.
 - (d) In the decomposition of H_2O_2 , the rate of the reaction is $k[H_2O_2]$.
- **62.** Which of the following isomerization reactions is of the first order?
 - (a) cyclopropane ----- propane
 - (b) cis-but-2-ene $\longrightarrow trans$ -but-2-ene
 - (c) vinyl allyl ether $\longrightarrow pent-4$ -enal
 - (d) $CH_3NC \longrightarrow CH_3CN$
- **63.** Which of the following reactions is of the first order?
 - (a) The decomposition of ammonium nitrate in an aqueous solution
 - (b) The inversion of cane-sugar in the presence of an acid
 - (c) The acidic hydrolysis of ethyl acetate
 - (d) All radioactive decays
- **64.** Which of the following are examples of unimolecular reactions?
 - (a) $O_3 \rightarrow O_2 + O$
 - (b) CH_2 CH_2 CH_3CH CH_2
 - (c) $NO + O_3 \rightarrow NO_2 + O_2$
 - (d) $O + NO + N_2 \rightarrow NO_2 + N_2$
- 65. The calculation of the pre-exponential factor is based on the
 - (a) idea that, for a reaction to take place, the reactant species must come together
 - (b) calculation of the molecularity of the reaction
 - (c) idea that the reactant species must come together, leading to the formation of the transition state which then transforms into the products
 - (d) calculation of the order of the reaction

66. Which of the following are examples of pseudo-unimolecular reactions?

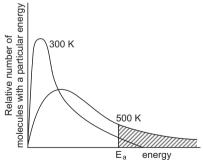
(a)
$$CH_3CO_2C_2H_5 + H_2O \xrightarrow{H^+} CH_3CO_2H + C_2H_5OH$$

(b)
$$C_{12}H_{22}O_{11} + H_2O \xrightarrow{H^+} C_6H_{12}O_6 + C_6H_{12}O_6$$
(glucose) (fructose)

(c)
$$CH_3COCl + H_2O \longrightarrow CH_3CO_2H + HCl$$

(d)
$$CH_3CO_2C_2H_5 + H_2O \xrightarrow{OH^-} CH_3CO_2H + C_2H_5OH$$

- **67.** In which of the following ways does an activated complex differ from an ordinary molecule?
 - (a) It is quite unstable and has no independent existence.
 - (b) ΔH°_{f} is probably positive.
 - (c) The system has a greater vibrational character.
 - (d) The system has no vibrational character.
- 68. The basic theory behind Arrhenius's equation is that
 - (a) the number of effective collisions is proportional to the number of molecules above a certain threshold energy
 - (b) as the temperature increases, so does the number of molecules with energies exceeding the threshold energy
 - (c) the rate constant is a function of temperature
 - (d) the activation energy and pre-exponential factor are always temperature-independent
- **69.** The distribution of molecular kinetic energy at two temperature is as shown in the following graph.



Which of the following conclusions are correct?

- (a) The number of molecules with energy $E_{\rm a}$ or greater is proportional to the shaded area for each temperature.
- (b) The number of molecules with energy $E_{\rm a}$ or less is proportional to the shaded area for each temperature.

- (c) The number of molecules with energy E_a is the mean of all temperatures.
- (d) The graph follows the Maxwell-Boltzmann energy distribution law.
- **70.** In Arrhenius's equation, $k = A \exp\left(-\frac{E_a}{RT}\right)$. A may be termed as the rate constant at

Answers

- (a) very low temperature
- (b) very high temperature
- (c) zero activation energy
- (d) the boiling temperature of the reaction mixture

			_	
1. c	2. b	3. d	4. b	5. d
6. c	7. a	8. b	9. a	10. d
11. b	12. c	13. c	14. c	15. d
16. b	17. c	18. d	19. a	20. c
21. c	22. a	23. d	24. c	25. a
26. c	27. c	28. c	29. c	30. a
31. d	32. c	33. d	34. b	35. c
36. d	37. a	38. a	39. a	40. b
41. b	42. d	43. c	44. d	45. b
46. a	47. b	48. d	49. b	50. c
51. d	52. a	53. d	54. c	55. a
56. b	57. d	58. a	59. b	60. a, b, c
61. a, b, c, d	62. a, b, c, d	63. a, b, c, d	64. a, b	65. a, c

Hints to More Difficult Problems

70. b, c

2. Use the formula $k = \frac{2.303}{t} \log \frac{a}{a - x}$.

66. a, b, c **67.** a, c **68.** a, b, c **69.** a, d

8. For a zero-order reaction, $-\frac{d[A]}{dt} = k$. $-\int_{C_0}^{C_t} d[A] = k \int_{t=0}^{t=0} dt.$ $C_0 - C_t = kt \implies k = \frac{C_0 - C_t}{t}.$

10.
$$k = \frac{C_0 - C}{t}$$
.

At
$$t_{1/2}$$
, $C = \frac{1}{2}C_0$ and $t = t_{1/2}$ or $k = \frac{C_0 - \frac{1}{2}C_0}{t_{1/2}} = \frac{\frac{1}{2}C_0}{t_{1/2}}$.
 $\therefore t_{1/2} = \frac{C_0}{2k} = \frac{a}{2k}$ $[C_0 = a]$.

15. Rate =
$$KC^n$$
 or $r = KC^n$. (1)

$$8r = K(4C)^n. (2)$$

Dividing Equation (2) by Equation (1), we get

$$2^3 = 2^{2n}$$
 or $2n = 3 \implies n = 1.5$.

17.
$$C_2H_4O(g) \longrightarrow CH_4(g) + CO(g)$$
At $t = 0$ p_0 0 0
At $t = t$ $p_0 - p$ p

According to the question,

 $p_0 - p + p = 120 \text{ mm}$ or $p_0 + p = 120 \text{ mm}$ or p = 120 - 80 = 40 mm.

For a first-order reaction,

$$k = \frac{1}{t} \ln \frac{p_0}{p_0 - p} = \frac{1}{20} \ln \frac{80}{80 - 40} = \frac{1}{20} \ln 2.$$
 (1)

But we know that
$$t_{1/2} = \frac{\ln 2}{k}$$
. (2)

From Equations (1) and (2),

$$k = \frac{1}{20} k t_{1/2} \implies t_{1/2} = 20 \text{ min.}$$

19.
$$k = \frac{1}{t} \ln \frac{100}{100 - 99.9} = \frac{1}{t} \ln \frac{100}{0.1}$$

or $\frac{\ln 2}{t_{1/2}} = \frac{1}{t} \ln 10^3$
or $\frac{\log 2}{t_{1/2}} = \frac{1}{t} \times \log 10^3 = \frac{3}{t}$.
 $t_{1/2} = \frac{\log 2}{3} \times t = \frac{0.30103}{3} \times t \approx 0.10t$.

$$\therefore \quad t = 10 \ t_{\frac{1}{2}}.$$

21.
$$k = \frac{1}{t} \ln \frac{a}{a - x} \Rightarrow \ln \frac{a}{a - x} = kt$$
.

$$\Rightarrow \ln \frac{a - x}{a} = -kt \Rightarrow a - x = ae^{-kt}$$
or $x = a(1 - e^{-kt}) \Rightarrow \frac{x}{a} = (1 - e^{-kt})$.

28. When
$$T \rightarrow \infty$$
 then $k = A$.

$$\therefore k = 4 \times 10^{15} \text{ s}^{-1}.$$

33.
$$\overrightarrow{E_a} - \overleftarrow{E_a} = \Delta U$$
. When $\overrightarrow{E_a} = \overleftarrow{E_a}$ then $\Delta U = 0$.

34.
$$[H^+] = \frac{6 \times 10^{-7} \text{ mol}}{0.05 \times 10^{-3} \text{ L}} = 1.2 \times 10^{-2} \text{ M}$$

or
$$r = \frac{\Delta x}{\Delta t}$$
 or $\Delta T = \frac{\Delta x}{r} = \frac{1.2 \times 10^{-2} \text{ M}}{6 \times 10^{5} \text{ M s}^{-1}}$.

$$\therefore t = 2 \times 10^{-8} \text{ s.}$$

40. Rate =
$$\frac{dC}{dt} = \frac{0.6 - 0.3}{10 - 5}$$
 or $\frac{0.9 - 0.6}{15 - 10} = 0.06 \text{ mol dm}^{-3} \text{ s}^{-1}$.

45.
$$E_a = \frac{K_2 E_a + K_2' E_a'}{K_2 + K_2'} = \frac{1.0 \times 10^{-2} \times 60 + 3.0 \times 10^{-2} \times 70}{1.0 \times 10^{-2} + 3.0 \times 10^{-2}}$$

= 67.5 kJ mol⁻¹

48.
$$k_{\text{uncat}} = A[-\exp(E_{\text{uncat}}/RT)]$$

$$k_{\text{catl}} = A[-\exp(E_{\text{uncatl}}/RT)]$$

$$k_{\text{uncatl}} = \exp[-(E_{\text{uncatl}} - E_{\text{catl}}/RT]]$$

$$k_{\text{catl}} = \exp(-8.314 \times 10^3 / 8.314 \times 300) = e^{-3.33}$$

or
$$\frac{k_{\text{catl}}}{k_{\text{uncatl}}} = e^{3.33} \approx 28$$

53. Represent the reaction in terms of an elementary reaction.

58. Experimental finding

14

Ionic Equilibrium

• *Type 1* •

1. If the H⁺ ion concentration of a solution is increased to ten times its initial

Choose the correct option. Only one option is correct.

	value,	its pH will					
	(a)	increase by o	ne		(b)	remain unchange	ed
	(c)	decrease by	one		(d)	increase by ten	
2.	50 mL				M N (b)	n a solution prepa NaOH and 25 mL o 0.05 M 0.10 M	
3.	20.00 i				mL (b)	t results from th of 0.01 M HCl. 10.53 8.35	e addition of
4.	solutio		1.01		he re	NaOH is added ar sulting solution is 11.95 (d)	nd the resulting 12.95
5.	(a)	H of milk lies 6.6 and 6.9 7.0 and 7.5	betv	ween	()	2.60 and 4.40 7.35 and 7.45	
6.	(a) (b) (c)	of the follow $K_{\rm a}$ (weak acid $K_{\rm a}$ (strong ac $K_{\rm a}$ (weak acid $K_{\rm a}$ (weak acid	d) · l id) · d) · l	$K_{\rm b}$ (conjugate $K_{\rm b}$ (conjugat $K_{\rm b}$ (weak base	e we e) = <i>l</i>	$ (x_{w}) = K_{w} $ $ (x_{w}) = K_{w} $	

dissociation for a weak acid?

(a) 1.0 M

7. Which of the following concentrations has the largest degree of

(b) 0.5 M

	(c)	0.10 M	(d)	0.01 M
8.	A give	en weak acid (0.01 M) has pK_a	= 6. 7	Γhe pH of the solution is
	(a)	3	(b)	4
	(c)	5	(d)	6
9.	Two v	veak solutions are isohydric w	hen i	their
	(a)	hydrogen-ion concentrations	are t	he same before mixing
	(b)	hydrogen-ion concentrations	are t	he same before and after mixing
	(c)	degrees of dissociation are the	e san	ne
	(d)	chemical properties are the sa	me	
10.	An acconcer	ridic solution (0.1 M) of a salt ntration of H_2S is, given $K_1K_2 =$	is s 10 -2	aturated with H_2S (0.1 M). The
	(a)	10^{-22}	(b)	10^{-19}
	(c)	10^{-20}	(d)	10^{-18}
11.	Which NaOH		an b	be titrated with HCl as well as
	(a)	Glycine	(b)	Pyruvic acid
	(c)	Triethylamine	(d)	Aniline
12.	The p	H of 10^{-10} M $\mathrm{H_2SO_4}$ will be alm	ost	
	(a)	4	(b)	7
	(c)	6	(d)	0
13.	How hydro	much water should be addec gen-ion concentration equal to	d to	10.0 g of acetic acid to give a \times 10 ⁻³ M (given p K_a = 4.74)?
	(a)	4 L	` ′	6 L
	(c)	5 L	(d)	3 L
14.		H of the new solution 5.0.		ent water is added to it to make hydrogen-ion concentration is
	(a)	tenfold	(b)	sevenfold
	(c)	thousandfold	(d)	hundredfold
15.				y mixing 2.0 mL of a strong acid strong base (NaOH) of pH 10.0.
	(a)	2.5	(b)	3.5
	(c)	4.5	(d)	6.5

16.	16. The dissociation constant of monobasic acids A, B and C are 10^{-4} , 10^{-4} and 10^{-10} respectively. The concentration of each is 0.1 M. Which of t following has been arranged in order of increasing pH?				
	(a) A	A < B < C	(b) $C < A < B$		
	(c) 1	B < C < A	(d) $B < A \approx C$		

- 17. Among the following, which causes the greatest change in pH on addition to 50 mL of a 0.2 M malonic acid solution?
 - (a) 25 mL of 0.02 M malonic acid

(c) B < C < A

- (b) 25 mL of 0.02 M NaOH solution
- (c) 25 mL of 0.02 M HCl solution
- (d) 50 mL of 0.2 M acetic acid
- 18. An acetic acid and sodium acetate buffer has pH = 5.36. The ratio of concentrations [OAc]/ [HOAc] is (pK_a) of acetic acid = 4.76)
 - (b) 4:3 (a) 6:1 (d) 4:1 (c) 1:1
- 19. Calculate the pH at the equivalence point in the titration of 25 mL of 0.10 M formic acid with a 0.1 M NaOH solution (given that pK_a of formic acid = 3.74).
 - (a) 4.74 (b) 8.74 (c) 8.37 (d) 6.06
- 20. A buffer solution is used in the
 - (a) preparation of chrome alum
 - (b) removal of PO₄³⁻ ions in qualitative analysis
 - (c) precipitation of Zn(OH)₂ from ZnSO₄
 - (d) determination of the ionic product of water
- **21.** Which of the following mixture solutions has pH = 1.0?

(a)
$$100 \text{ mL} \frac{N}{10} \text{ HCl} + 100 \text{ mL} \frac{M}{10} \text{ NaOH}$$

(b)
$$55 \text{ mL} \frac{M}{10} \text{ HCl} + 45 \text{ mL} \frac{M}{10} \text{ NaOH}$$

(c)
$$10 \text{ mL} \frac{M}{10} \text{ HCl} + 90 \text{ mL} \frac{M}{10} \text{ NaOH}$$

(d)
$$75 \text{ mL} \frac{M}{5} \text{ HCl} + 25 \text{ mL} \frac{M}{5} \text{ NaOH}$$

- 22. Fear or excitement generally causes one to breathe rapidly and results in the decrease of CO₂ concentration in one's blood. In what way does this change the pH of blood?
 - (a) It increases.

- (b) It decreases.
- There is no change in the pH. (d) The pH level is adjusted at 7.

		Ionic Equilibrium	1-185			
23.	The pK_a of acetyl salicyclic acid (aspirin) is 3.5. The pH of gastric juice in the human stomach is about 2–3 and that in the small intestine about 8.0. Aspirin will be					
	(a)	unionized in the small intestine and in the stomach				
	(b)	completely ionized in the small intestine and in the stomach				
	(c)	ionized in the stomach and almost unionized in the small int	estine			
	(d)	ionized in the small intestine and almost unionized in the sto	omach			
24.		$^{\circ}$ C, the dissociation constants of CH ₃ COOH and NH ₃ in aq ons are almost the same. The pH of a solution of 0.01 M CH ₃				

is 4.0 at 25°C. The pH of a 0.01 M NH₃ solution at the same temperature is

(a) 3.0

(b) 4.0

(c) 10.0

(d) 11.0

- 25. The pH of blood does not appreciably change by a small addition of acid or a base because blood
 - (a) contains serum protein which acts as a buffer
 - (b) contains iron as a part of the molecule
 - (c) can be easily coagulated
 - (d) is a body fluid
- **26.** What will be the K_h value of a weak base (BOH) of which a 0.1 M solution has a pH of 8, when it is half neutralized with 0.1 M HNO₃?

(a) 10^{-8}

(b) 10^{-4}

(c) 10^{-6}

(d) 10^{-7}

- 27. For the precipitation of cations of group IV in qualitative analysis, the medium is made alkaline before passing H₂S gas. The medium is made alkaline because this helps
 - (a) suppress the ionization of H₂S
 - (b) increase the ionization of H₂S
 - (c) increase the ionization of metal salts
 - (d) reduce the ionization of metal salts
- 28. The solubility of AgCl in 0.1 M NaCl will
 - (a) increase
 - (b) decrease
 - (c) remain unchanged
 - (d) AgCl will dissociate completely.
- **29.** The solubility of a salt A_2B_3 is 1.0×10^{-3} M. Its solubility product is

(a) 1.08×10^{-13}

(b) 1.08×10^{-15}

(c) 1.08×10^{-10}

(d) 1.08×10^{-17}

this solid is

(a) $4 \times 10^{-4} \,\mathrm{M}$

(c) $2 \times 10^{-2} \,\mathrm{M}$

(a) $1 \times 10^{-4} \,\mathrm{M}$

(c) $1 \times 10^{-2} \,\mathrm{M}$

30. M_2SO_4 (M^+ is a monovalent metal ion) has a K_{sp} of 3.2×10^{-5} at 25°C. The maximum concentration of SO_4^{2-} ions possible in a saturated solution of

31. The pH of an aqueous solution of Ba(OH)₂ is 10.0. If the $K_{\rm sp}$ of Ba(OH)₂ is 1×10^{-9} then the concentration of Ba²⁺ ions in the solution is

32. The $K_{\rm ep}$ of Mg(OH), is 1×10^{-12} . 0.01 M MgCl₂ will precipitate at the

(b) $5 \times 10^{-6} \text{ M}$ (d) $8 \times 10^{-3} \text{ M}$

(b) $1 \times 10^{-6} \text{ M}$ (d) $1 \times 10^{-1} \text{ M}$

	limiti	ng pH	7.2		0 2	•	1
	(a)	3	(b) 9	(c)	12	(d)	8
33.	M(OF	H_{x} has a K_{sp}	of 4×10^{-9} and its	solu	bility is 10^{-3} l	M. Tł	he value of x is
	(a)		(b) 1	(c)		(d)	
34.	Wher precip	equal volu pitation of Ag	umes of the for $Cl(K_{sp} = 1.8 \times 10^{-5})$	llow -10) w	ing solution vill occur with	s ar	e mixed, the
	(a)	$10^{-4} \mathrm{M} (\mathrm{Ag})$	$^{\scriptscriptstyle +}$) and $10^{ \scriptscriptstyle -4}\mathrm{M}$ (Cl	_)			
	(b)	$10^{-5}\mathrm{M}$ (Ag	$^{+}$) and $10^{-5}\mathrm{M}$ (Cl	_)			
	(c)	$10^{-5} \mathrm{M} (\mathrm{Ag})$	$^{+}$) and $10^{-6}\mathrm{M}$ (Cl	_)			
	(d)	$10^{-10}{ m M}$ (Ag	$ m g^{+}$) and $10^{-10} m M$ (C	Cl ⁻)			
35.	Why H ₂ S is	does only As passed thro	³⁺ get precipitated ugh an acidic solu	d as . ition	As ₂ S ₃ and not containing A	: Zn ² s ³⁺ a	²⁺ as ZnS when and Zn ²⁺ ?
	(a)	The solubili	ty product of As ₂	S_3 is	less than that	of Z	inS.
	(b)	Enough As	3+ ions are present	t in a	n acidic medi	um.	
	(c)	A zinc salt o	does not ionize in	an a	cidic medium	1.	
	(d)	The solubili	ty product chang	es in	the presence	of ar	n acid.
36.	of 10	⁻⁶ M. Which o	ing NH ₄ Cl and NI of the following h ded to an equal v	ydro	oxides may be	pre	cipitated when
	(a)	$Ba(OH)_2(K_s)$	$_{\rm sp} = 1.0 \times 10^{-4}$)	(b)	$Ca(OH)_2(K_s)$	sp = 2	2.2×10^{-5})
	(c)	$Mg(OH)_2$ (K	$X_{\rm sp} = 3.0 \times 10^{-12}$	(d)	$Fe(OH)_2 (K_s)$	$_{p} = 8.$	$.0 \times 10^{-16}$)
37.	conta	is the maximining 0.15 M S^{2-}] _{H₂S} = 4 × 1	num possible con HCl and 0.10 M 0^{-21} .	H ₂ S	? Given that	$K_{\rm sp}(1)$	is in a solution NiS) = 2×10^{-21}
	(a)	0.65 M	(b) 0.45 M	(c)	0.10 M	(d)	0.15 M

38. 50 mL of a solution containing 10^{-3} mol of Ag $^+$ is mixed with 50 mL of a 0.1 M HCl solution. How much Ag $^+$ remains in solution ($K_{\rm sp}$ of AgCl = 1.0×10^{-10})?

39. At a certain temperature, a saturated solution of Zn(OH)2 has a pH of

8.63. The value of $K_{\rm sp}$ of ${\rm Zn}({\rm OH})_2$ at this temperature is

(b) 2.5×10^{-7} (d) 2.5×10^{-10}

(a) 2.5×10^{-9}

(c) 2.5×10^{-8}

	(a)	4.0×10^{-18}	(b)	3.9×10^{-28}
	(c)	3.9×10^{-17}	(d)	2.0×10^{-19}
	(a) (b) (c) (d) Which	A salt of a strong acid and a set A salt of a strong acid and a set A salt of a strong acid and a set A salt of a weak acid and a set A salt of a weak acid and a weak acid an	trong veak rong eak l	g base base base pase
		•		
	(c)	$(NH_4)_2CO_3$	(d)	HCOONH ₄
42.	0.50 L			eutralized with 0.20 M NaOH in lution will be (given that pK_a for
	(a)	12.67	(b)	7.87
	(c)	8.87	(d)	7.00
43.	have	alent amounts of aqueous solu a dissociation constant of 5 dysis of the salt formed by ther	$5.0 \times$	s of a weak acid and a weak base 10^{-7} each. The percentage of
	(a)	40	(b)	
	(c)	50	(d)	25
44.		on obtained by mixing equ		lutes, calculate the pH of the volumes of $N/10$ NaOH and
	(a)	7.6	(b)	12.4
	(c)	1.6	(d)	13.4
45.		, for the fluoride ion at 25°C ofluoric acid in water at this ten		0.83, the ionization constant of ature is
	(a)	1.74×10^{-5}	(b)	3.52×10^{-3}
	(c)	6.76×10^{-4}	(d)	5.38×10^{-2}

 $Fe(OH)_2 = 1.6 \times 10^{-14}$].

(a) 8.9×10^{-10}

(a) 0.06

46. Calculate the molar solubility of $Fe(OH)_2$ at a pH of 8.00 [K_{sp} of

47. Calculate the molar solubility of AgCl in a 1-L solution which contains

(b) 0.016

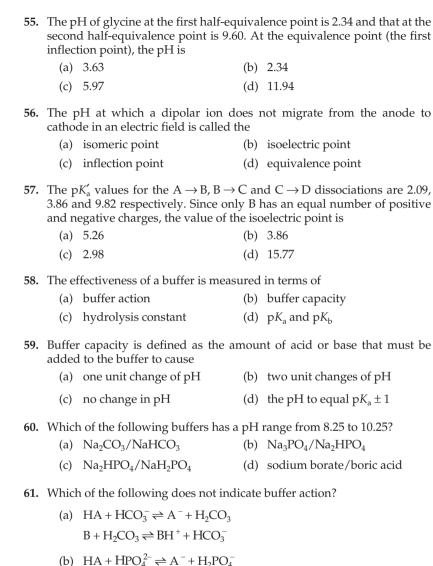
10.0 g of CaCl₂ [$K_{sp}(AgCl) = 1.6 \times 10^{-10}$].

(c) 0.010

(b) 8.9×10^{-11}

(d) 0.16

	(c)	8.9×10^{-9}		(d)	8.9×10^{-12}		
48.			ic acid are 6.5×1 0.01 M solution of			resp	ectively. What
	(a)	9.6×10^{-6}		(b)	1.4×10^{-1}		
	(c)	1.3×10^{-6}		(d)	1.3×10^{-8}		
49.			of a solution con $10^{-7} \times 10$ and K_2	tainin (HCC	(9.1 M) + (1.0 M) = (1.0	O_3^- ar $^{-11}$].	nd 0.2 M CO ₃ ²⁻
	(a)	3.18		(b)	10.62		
	(b)	6.62		(d)	9.31		
50.	Calcu = 1.0 > 1 × 10	< 10 ^{−5} change	at which an	acid the c	indicator w oncentration	of the	K_{acid} (indicator) he indicator is
	(a)	5		(b)	11		
	(c)	3		(d)	8		
51.	At wh	nat pH will a	1×10^{-4} M solut	tion o	f an indicato	r will	$K_{\rm b}$ (indicator)
	$=1\times1$	10^{-11} change	colour?				
	(a)	7.0		(b)	3.0		
	(c)	5.5		(d)	11.0		
52.	indica	itor is red a	icator has a K_a and the basic for the colour of the	m is	blue. Calcul	ate tl	he pH change
	(a)	1.20		(b)	0.80		
	(c)	0.20		(d)	1.40		
53.	Calcu K ₂ (HC	late the pH CO_3^-) = 4.8×1	of a 0.01 M Nal 0^{-11}].	HCO ₃	solution [K	1(H ₂ C	$(CO_3) = 4 \times 10^{-7},$
	(a)	9.38	(b) 6.38	(c)	8.38	(d)	7.38
54.	Which	n of the follow	wing statements i	is corı	rect for glycir	ne?	
	(a)	It behaves a	s a base when tit	rated	with HCl.		
	(b)	It behaves a	s an acid when t	itrated	d with NaOH	I.	
	(c)	It forms the	zwitterion NH ₃ C	CO ₂ CO	OO⁻.		
	(d)	All of these					



 $B + H_2PO_4^- \rightleftharpoons BH^+ + HPO_4^{2-}$

(d) $HA + CH_3COO^- \rightleftharpoons A^- + CH_3COOH$ $B + CH_3COOH \rightleftharpoons BH^+ + CH_3COO^-$

(c) $HA + PO_4^{3-} \rightleftharpoons HPO_4^{2-} + A^{-}$ $B + HPO_4^{2-} \rightleftharpoons PO_4^{3-} + BH$

- 62. Blood plasma is maintained at a pH of 7.4 largely by the
 - (a) HCO₃⁻/H₂CO₃ buffer
 - (b) HPO_4^{2-}/PO_4^{3-} buffer
 - (c) HCO_3^-/H_2CO_3 and $HPO_4^{2-}/H_2PO_4^-$ buffers
 - (d) HPO₄²⁻/H₃PO₄ and haemoglobin buffers
- **63.** The $[HCO_3^-]/[H_2CO_3]$ ratio in the blood (pH = 7.4) is approximately (p K_a = 6.1 for H_2CO_3)
 - (a) 20

(b) 16

(c) 24

- (d) 10
- **64.** In the titration of a weak diprotic acid (H_2A) with a strong base (NaOH), $[H^+]$ is given by
 - (a) $\sqrt{K_{a_1}c_a}$

(b) $K_a \sqrt{c_{a_1}}$

(c) $K_a c_{a_1}$

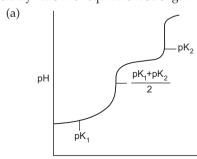
(d) K_{a_1}

(where K_a is the dissociation constant for the first stage and c_a is the concentration of the weak acid).

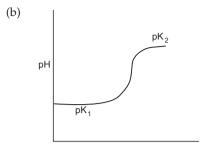
- **65.** During the titration of a weak diprotic acid (H₂A) against a strong base (NaOH), the pH of the solution half-way to the first inflection point and that at the first inflection point are given respectively by
 - (a) pK_1 and $pK_1 + pK_2$
- (b) $\sqrt{K_1 c_a} \text{ and } \frac{pK_1 + pK_2}{2}$
- (c) pK_1 and $\frac{pK_1 + pK_2}{2}$
- (d) $pK_1 + pK_w$ and $\frac{pK_1 + pK_2}{pK_w}$

where K_1 and K_2 are the first and second dissociation constants of H_2A , and c_a is the initial concentration of the weak acid.

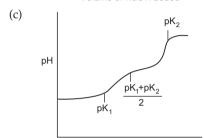
66. Which of the following curves indicates the titration of a weak diprotic acid by NaOH of equivalent strength?



Volume of NaOH added



Volume of NaOH added



 $\begin{array}{c|c} & & & & \\ \text{Volume of NaOH added} \\ \text{(d)} & & & & \\ \hline \text{pH} & & & & \\ \hline \text{pK}_1 + \text{pK}_2 & & \\ \hline \text{pK}_1 & & & \\ \hline \end{array}$

Volume of NaOH added

- 67. Which of the following represents hydrolysis?
 - (a) $HCO_3^- + H_2O \rightleftharpoons CO_3^{2-} + H_3O^+$
 - (b) $HCO_3^- + H_2O \rightleftharpoons H_2CO_3 + OH^-$
 - (c) $H_3BO_3 + H_2O \rightleftharpoons H_2BO_3^- + H_3O^+$
 - (d) $H_2PO_4^- + H_2O \rightleftharpoons HPO_4^{2-} + H_3O^+$
- **68.** The concentration of a sample of H_2CO_3 is 1.0×10^{-3} M. Which of the following is correct?
 - (a) $1.0 \times 10^{-3} \text{ M} = [\text{H}_2\text{CO}_3] + [\text{HCO}_3^-] + 2[\text{CO}_3^{2-}]$
 - (b) $1.0 \times 10^{-3} \text{ M} = [\text{H}_2\text{CO}_3]$
 - (c) $1.0 \times 10^{-3} \text{ M} = [\text{H}_2\text{CO}_3] + [\text{HCO}_3^-] + [\text{CO}_3^{2-}]$
 - (d) $1.0 \times 10^{-3} \text{ M} = [\text{H}_2\text{CO}_3] + [\text{HCO}_3^-]$
- 69. The inflection point of a titration curve is the point at which

73.

- (a) the second derivative is zero (b) pH = 0(c) pH = 7(d) the first derivative is zero • Type 2 • Choose the correct options. More than one option is correct. **70.** Which of the following mixtures constitute a buffer? (a) CH₃COOH + CH₃COONa (b) Na₂CO₃ + NaHCO₃ (c) NaCl + HCl (d) $NH_4Cl + (NH_4)_2SO_4$ 71. Which of the following mixtures constitute a buffer? (a) Na₂CO₃ + HCl (b) NaOH + CH₃COOH (c) $NH_3 + CH_3COONH_4$ (d) NaOH + BaCl₂ 72. In which of the following pairs of solutions is there no effect on the pH upon dilution? (a) $0.1 \text{ M NH}_3 \text{ and } 0.1 \text{ M (NH}_4)_2 \text{SO}_4$ (b) 0.1 M NaH₂PO₄ and 0.1 M Na₂HPO₄ (c) 0.1 M HCl and 0.01 M NaOH (d) 0.1 M KCl and 0.1 M HCl Which of the following will suppress the ionization of phthalic acid in an aqueous solution? (a) KCl (b) H₂SO (c) HNO₃ (d) NaOH 74. Which of the following buffers have a pH greater than 7? (a) $NaHCO_3 + Na_2CO_3$ (b) CH₂COOH + CH₂COONa
 - (c) $Na_3PO_4 + NaH_2PO_4$ (d) $NH_3 + (NH_4)_2SO_4$ 75. Which of the following mixtures can act as a buffer?
 - (a) NaOH + CH₃COONa (1 : 1 molar ratio)
 - (b) CH₃COOH + NaOH (2 : 1 molar ratio)
 - (c) CH₃COOH + NaOH (3 : 1 molar ratio)
 - (d) CH₃COOH + NaOH (1:1 molar ratio)
 - 76. When HCl(g) is passed through a saturated solution of common salt, pure NaCl is precipitated because
 - (a) HCl is highly soluble in water
 - (b) the ionic product [Na $^+$][Cl $^-$] exceeds its solubility product (K_{sp})
 - (c) the K_{sp} of NaCl is lowered by the presence of Cl⁻ions
 - (d) HCl causes precipitation

Answers

1. c	2. b	3. a	4. d	5. a
6. a	7. d	8. b	9. a	10. c
11. a	12. b	13. d	14. c	15. b
16. a	17. b	18. d	19. c	20. b
21. d	22. c	23. d	24. b	25. a
26. c	27. b	28. b	29. a	30. c
31. d	32. b	33. d	34. a	35. a
36. d	37. b	38. c	39. c	40. a
41. b	42. c	43. d	44. b	45. c
46. b	47. a	48. c	49. b	50. a
51. b	52. a	53. c	54. d	55. c
56. b	57. c	58. b	59. a	60. d
61. c	62. c	63. a	64. a	65. c
66. a	67. b	68. c	69. a	70. a, b
71. a, b	72. a, b	73. b, c	74. a, b	75. b, c
76. b, d				

Hints to More Difficult Problems

2.
$$H^+ = \frac{(25 \times 0.3 - 25 \times 0.1) \text{ meq}}{100 \text{ mL}} = 0.05 \text{ M}.$$

6.
$$HA + H_2O \rightleftharpoons H_3O^+ + A^-$$
, $K_a(HA) = \frac{[H_3O^+][A^-]}{[HA]}$. (1)

$$A^- + H_2O \rightleftharpoons OH^- + HA^-, \quad K_b(A^-) = \frac{[OH^-][HA]}{[HA^-]}.$$
 (2)

Multiplying Equation (1) by Equation (2),

$$K_{a}(HA) \times K_{b}(A^{-}) = [H_{3}O^{+}][OH^{-}] = K_{w}.$$

 $\textbf{7.} \ \ The degree of dissociation increases with dilution.$

8.
$$pH = \frac{1}{2}pK_a - \frac{1}{2}\log C = \frac{1}{2} \times 6 - \frac{1}{2}\log(0.01)$$

= 3 + 1 = 4.

- **11.** Glycine (H₂N · CH₂CO₂H) contains an acidic group as well as a basic group.
- **15.** Number of milliequivalents of HCl = 2×10^{-3} . We know that pH + pOH = 14

⇒
$$10 + \text{pOH} = 14$$
.
 $\text{pOH} = 10^{-4} \text{ M}$.
 $3 \text{ mL of } 10^{-4} \text{ M NaOH} = 3 \times 10^{-4}$.

Number of milliequivalents of NaOH = 3×10^{-4} .

The resulting solution is acidic.

$$[H^+] = \frac{(2 \times 10^{-3} - 3 \times 10^{-4} \text{ m. eq.}}{(2+3) \text{ mL}} = 3.4 \times 10^{-4} \text{ M}$$

or
$$pH = -log[H^+] = -log(3.4 \times 10^{-4}) = 3.5.$$

18. Use the Henderson equation

$$pH = pK_a + \log \frac{[OAC^-]}{[HOAC^-]}$$

19. At the equivalence point, 0.10 M of HCOONa (sodium formate) is formed. This problem is solved by the considering the concept of hydrolysis equilibria.

$$pH = \frac{1}{2}pK_w + \frac{1}{2}pK_a + \frac{1}{2}\log C$$
$$= \frac{1}{2} \times 14 + \frac{1}{2} \times 3.74 + \frac{1}{2}\log 0.1 = 8.37.$$

26. $K_{\rm a} \cdot K_{\rm b} = K_{\rm w}$

Under this condition,

$$pH = pK_a = 8 \implies K_a = 10^{-8}$$
 or
$$10^{-8} \cdot K_b = 10^{-14} \implies K_b = 10^{-6}.$$

- **29.** $K_{\rm sp}$ for A_2B_3 equilibria = $108~{\rm S}^5 = 108 \times (1.0 \times 10^{-3})^5 = 1.08 \times 10^{-13}$.
- **34.** The precipitation will occur when $Q > K_{\rm sp}$. $Q = 10^{-4} \times 10^{-4} = 10^{-8} > K_{\rm sp} (1.8 \times 10^{-10})$.

38.
$$Ag^+ + Cl^- \longrightarrow AgCl(s)$$

 $\downarrow \qquad \qquad \downarrow$
 $1 \times 10^{-3} \text{ mol} \qquad 5 \times 10^{-3} \text{ mol}$

$$\{[HCl] = 50 \text{ mL} \times 0.1 \text{ mol L}^{-1}$$

= 5.0 m mol = 5.0 × 10⁻³ mol.

$$[H^+] = [Cl^-] = 5.0 \times 10^{-3} \text{ mol}$$
.

 $[Cl^{-}] = 4 \times 10^{-3}$ mol after the reaction.

$$[Ag^+] = \frac{1.0 \times 10^{-10}}{4.0 \times 10^{-3}} = 2.5 \times 10^{-8}.$$

44.
$$[OH^{-}] = \frac{x \times 0.1 - x \times 0.05}{2x} = 2.5 \times 10^{-2}.$$

 $pOH = 2 - 0.4 = 1.6 \implies pH = 12.4.$

45.
$$pK_b = 10.83$$
, $log K_b = -10.83 = \overline{11}.17$
 $\Rightarrow K_b = 1.479 \times 10^{-11}$.
 $K_a \cdot K_b = K_w$
or $K_a \times 1.479 \times 10^{-11} = 1.0 \times 10^{-14} \Rightarrow K_a = 6.76 \times 10^{-4}$.

48. The hydrolysis of $C_2O_4^{2-}$ is represented by

$$C_2O_4^{2-} + H_2O \Rightarrow HC_2O_4^{-} + OH^{-}$$

 $K_h = \frac{K_w}{K_2} = \frac{1.0 \times 10^{-14}}{6.1 \times 10^{-5}} = \frac{x^2}{0.01}$.

$$\therefore x = 1.3 \times 10^{-6}$$
.

49.
$$HCO_3^- + H_2O \longrightarrow H_3O^+ + CO_3^{2-}$$

$$K_2 = \frac{[H_3O^+][CO_3^{2-}]}{[HCO_3^-]} = 4.8 \times 10^{-11}.$$

$$[H_3O^+] = 4.8 \times 10^{-11} \times \frac{[HCO_3^-]}{[CO_3^{2^-}]} = 4.8 \times 10^{-11} \left(\frac{0.1}{0.2}\right).$$

$$pH = -log[H_3O^+] = -log[4.8 \times 10^{-11} \times 0.5]$$

= 10.62.

50. Use the equation

$$pH = pK_{In} + log \frac{[In^{-}]}{[HIn]} \cdot$$

During change of colour, i.e., at the neutral point,

$$[In^{-}] = [HIn]$$
 or $pH = pK_{In} = -log (1.0 \times 10^{-5}).$

52. Consider an indicator of the type HIn which is ionized as

$$HIn \rightleftharpoons H^+ + In^-$$

According to the question, 20% of the final form of the indicator is blue (acidic) and 80% red (basic).

$$K_{\text{In}} = \frac{[\text{H}^+][\text{In}^-]}{[\text{HIn}]}$$
, where K_{In} is the indicator constant.

$$[H^+] = K_{In} \frac{[In^-]}{[HIn]} = 1 \times 10^{-5} \left(\frac{80}{20}\right) = 4 \times 10^{-5}.$$

$$pH = 4.4.$$

For 80% blue,
$$[H^+] = 1 \times 10^{-5} \left(\frac{20}{80}\right)$$
.
 $pH = 6 - 0.4 = 5.6$.
 $\Delta pH = 5.6 - 4.4 = 1.2$.

57. The pH at which the dipolar ion does not migrate in an electric field is called the isoelectric point, denoted by pI.

The isoelectric point pI =
$$\frac{2.09 + 3.86}{2}$$
 = 2.975 \approx 2.98.

60.
$$pK_a(H_3BO_3) = 9.25$$
. $pH = pK_a \pm 1$, i.e., 8.25 and 10.25.

63.
$$pH = pK_a + log \frac{[HCO_3^-]}{[H_2CO_3]}$$

or $7.4 = 6.1 + log \frac{[HCO_3^-]}{[H_2CO_3]}$
or $\frac{[HCO_3^-]}{[H_3CO_2]} = 20$.

64. For
$$H_2A \rightleftharpoons H^+ + HA^-$$
,

$$K_{a_1} = \frac{[H^+][HA^-]}{[H_2A]} \approx \frac{[H^+]^2}{c_a}$$
.

$$\therefore \quad [\mathbf{H}^+] = \sqrt{K_{\mathbf{a}_1} c_{\mathbf{a}}}.$$

65. At this point, half of the H_2A has been converted to HA^- so that $[HA] \approx [H_2A]$, giving $K_{a_1} = [H^+]$ $pH = pK_1$.

75. Both will give a mixture of CH₃COOH and CH₃COONa which serve as buffers.

15

Electrochemistry

• Type 1 •

1. Which of the following aqueous solutions remains neutral after

2. Which of the following aqueous solutions remains acidic after

(b) AgNO₃(d) NaCl

(b) KMnO₄

Choose the correct option. Only one option is correct.

electrolysis?
(a) CuSO₄

electrolysis?
(a) K₂Cr₂O₇

(c) K₂SO₄

	(c)	CH₃COONa	(d)	CuCl ₂
3.	Which batter	ě.	s is o	correct for a solid-state lithium
	(a)	-	-	lymer material that permits the ions, and the battery is not
	(b)			ymer material that permits the and the battery is rechargable.
	(c)	The voltage of the battery car	ı go	up to 12 V.
	(d)	The anode is made of what is	kno	wn as an insertion compound.
4.	How	many faradays are required to	redu	ice 1 mol of BrO ₃ to Br ⁻ ?
	(a)	3	(b)	5
	(c)	6	(d)	4
5.	How	many moles of Zn ²⁺ carry a ch	arge	of 2×10^{-5} C?
	(a)	1×10^{-4}	(b)	4×10^{-10}
	(c)	4×10^{-5}	(d)	1×10^{-10}
		1-197		

(a) 1.08×10^{5} C

(a) 1.50

(c) 2.50

(c) 1.08×10^4 electrons

7. How many kilojoules of energy is expended during the passage of a

8. A metal wire carries a current of 1 A. How many electrons move past a

(b) 0.11 F

(d) 1.51 V

(b) 0.15

(d) 9.65

6. A steady current of 3.0 A for 1 hour corresponds to a passage of

current of 0.1 A for 100 s under a potential of 150 V?

	point	in the wire in one second?		
	(a)	6.02×10^{23}	(b)	3.12×10^{18}
	(c)	3.02×10^{23}	(d)	6.24×10^{18}
9.	a curr	electrolysis of aqueous NaCl, sent of 1.0 A through the cell to M of NaOH?	for h	ow long would you have to pass vert 1.0 L of a 1 M NaCl solution
	(a)	35.5 hours	(b)	23.8 hours
	(c)	15.6 hours	(d)	26.8 hours
10.	How 1 mL	much charge is required to p s ⁻¹ by the electrolysis of molter	rodu n Na	ace hydrogen gas at the rate of Cl?
	(a)	8.6 C	(b)	18.4 C
	(c)	4.3 C	(d)	1.4 C
11.	of Cl (a) (b) (c)	umber of electrons gained and and 65.4 g of Zn^{2+} respectively 3.01×10^{23} and 3.01×10^{23} 6.02×10^{23} and 3.01×10^{23} 6.02×10^{23} and 6.02×10^{23} and 6.02×10^{23} 3.01×10^{23} and 6.02×10^{23}	l lost y are	during the electrolysis of 35.5 g
12.		mium metal can be plated out according to the following equ $CrO_3(g) + 6H^+ + 6e \rightarrow 0$	atior	m an acidic solution containing a (atomic weight of $Cr = 52$). $+3H_2O$
	How	many grams of chromium will	be p	plated by 22400 C?
	(a)	2.96	(b)	2.16
	(c)	21.6	(d)	23.2
13.	NaCl			400 mL of a 2.0 M solution of I of the solution after the current
	(a)	12.98	(b)	12.13
	(c)	10.48	(d)	9.24

- **14.** After the electrolysis of a sodium chloride solution with inert electrodes for a certain period of time, 600 mL of a 1 N NaOH solution was left. During the same time 31.80 g Cu was deposited in a copper voltameter in series with the electrolytic cell. Calculate the percentage of NaOH obtained (atomic weight of Cu = 63.6).
 - (a) 40

(b) 50

(c) 60

- (d) 25
- 15. During the purification of copper by electrolysis
 - (a) the anodes used are made of copper ore
 - (b) pure copper is deposited on the cathodes, with the evolution of hydrogen at the cathodes
 - (c) the impurities such as Ag, Au, Zn and Fe go into solution
 - (d) the voltage is carefully controlled to prevent the deposition of Zn and Fe at the cathode
- **16.** The density of copper is 8.94 g mL^{-1} . Find the charge needed to plate an area of $10 \times 10 \text{ cm}^2$ to a thickness of 10^{-2} cm using a CuSO₄ solution as electrolyte (atomic weight of Cu = 63.6).
 - (a) $2.7 \times 10^4 \text{ C}$

(b) $8.8 \times 10^4 \,\text{C}$

(c) $18.3 \times 10^4 \text{ C}$

- (d) 1.7×10^4 C
- 17. Calculate the volume of gas liberated at the anode at stp during the electrolysis of a CuSO₄ solution by a current of 1 A passed for 16 minutes 5 seconds.
 - (a) 224 mL

(b) 56 mL

(c) 112 mL

- (d) 448 mL
- **18.** Acidified water was electrolysed using an inert electrode. The volume of gases liberated at stp was 168 mL. The amount of electricity passed through the acidified water was
 - (a) 96,500 C

(b) 9,650 C

(c) 965 C

(d) 168 C

19. For the cell reaction

$$2Hg(1) + 2Ag^{+}(aq) \rightarrow 2Ag(s) + Hg_{2}^{2+}(aq)$$

it is given that

$$E_{Ag^{+}/Ag}^{0} = 0.800 \text{ V}, \text{ [Ag^{+}]} = 10^{-3} \text{ M},$$

 $E_{Hg^{-2+}/Hg}^{0} = 0.785 \text{ V} \text{ and [Hg}_{2}^{2+}] = 10^{-1} \text{ M}.$

- (a) The forward reaction is spontaneous.
- (b) The backward reaction is spontaneous.
- (c) $E_{\text{cell}} = 1.585 \text{ V}$
- (d) $E_{\text{cell}} = 3.170 \text{ V}$

20. Consider the cell reaction

$$Mg(s) + Cu^{2+}(aq) \rightarrow Cu(s) + Mg^{2+}(aq)$$

If the standard reduction potentials of Mg $^{2+}$ /Mg(s) and Cu $^{2+}$ /Cu(s) are -2.37 V and +0.34 V respectively, E° for the cell is

(a) -2.71 V

(b) +2.71 V

(c) -2.03 V

- (d) +2.03 V
- **21.** If all species are in their standard states, which of the following is the strongest oxidizing agent?
 - (a) Br⁻

(b) Zn^{2+}

(c) Fe²⁺

- (d) Co³⁺
- **22.** During the electrolysis of a concentrated brine solution, the amount of chlorine gas produced by the passage of 2F of electricity is
 - (a) 0.25 mol

(b) 0.50 mol

(c) 1.00 mol

- (d) 2.00 mol
- **23.** Fe³⁺(aq) + e \rightarrow Fe²⁺; $E^0 = +0.77 \text{ V}$

Al
$$^{3+}$$
(aq) + 3e \rightarrow Al(s); $E^0 = -1.66 \text{ V}$

$$Br_2(aq) + 2e \rightarrow 2Br^{-}(aq); E^0 = +1.08 \text{ V}$$

Considering the above data, state which of the following represents the correct order of reducing power.

(a) $Br^- < Fe^{2+} < Al$

(b) $Fe^{2+} < Al < Br^{-}$

- (c) $Al < Br^- < Fe^{2+}$
- (d) $Al < Fe^{2+} < Br^{-}$

24. $E_{\text{Na}^+/\text{Na(s)}}^0 = -2.71 \text{ V},$

$$E_{\rm Mg^{2+}/Mg(s)}^{0} = -2.37 \text{ V},$$

$$E_{\text{Fe}^{2+}/\text{Fe(s)}}^0 = -0.44 \,\text{V}$$
 and

$$E_{\rm Cr}^{0_{3+}}/C_{\rm r(s)} = -0.41 \text{ V}.$$

Based on this data, state which of the following is the weakest reducing agent.

(a) Na +

(b) Na

(c) Cr

- (d) Fe^{2+}
- **25.** Given that $E_{Ag^+/Ag}^0 = +0.80 \text{ V}$ and $E_{Zn^{2+}/Zn}^0 = -0.76 \text{ V}$, which of the following is correct?
 - (a) Ag + can be reduced by H₂.
 - (b) Ag can oxidize H₂ into H⁺.
 - (c) Zn^{2+} can be reduced by H_2 .
 - (d) Ag can reduce Zn²⁺.

26. For the cell reaction

$$Cu^{2+}(aq)(c_1) + Zn(s) \rightarrow Zn^{2+}(aq)(c_2) + Cu(s)$$

the change in free energy ΔG at a given temperature is a function of

(a) $\ln c_1$

(b) $\ln (c_2/c_1)$

(c) $\ln (c_1 + c_2)$

- (d) $\ln(c_2)$
- 27. A gas X at 1 atm is bubbled through a solution containing a mixture of 1 M Y⁻ and 1 M Z⁻ at 25°C. If the order of reduction potential is Z > Y > X,
 - (a) Y will oxidize X and not Z
 - (b) Y will oxidize Z and not X
 - (c) Y will oxidize X as well as Z
 - (d) Y will reduce as well as X and Z
- 28. The value of the reaction quotient, Q, for the cell

$$Zn(s) | Zn^{2+}(0.01 \text{ M}) | | Ag^{+}(1.25 \text{ M}) | Ag(s) \text{ is}$$

(a) 156

(b) 125

(c) 1.25×10^{-2}

- (d) 6.4×10^{-3}
- **29.** The logarithm of the equilibrium constant, $\log K_{\rm eq}$, of the net cell reaction of the cell, $X(s) | X^{2+} | Y^+ | Y(s)$ (given $E_{\rm cell}^0 = 1.20 \text{ V}$), is
 - (a) 47.2

(b) 40.5

(c) 21.4

- (d) 12.5
- **30.** The temperature coefficient of a cell whose operation is based on the reaction $Pb(s) + HgCl_2(aq) \rightarrow PbCl_2(aq) + Hg(l)$ is

$$\left(\frac{\partial E}{\partial T}\right)_p = 1.5 \times 10^{-4} \text{ V K}^{-1}.$$

The amount of heat released during the operation is

(a) 8.4 kJ mol^{-1}

(b) 16.4 kJ mol^{-1}

(c) 6.5 kJ mol^{-1}

- (d) 4.5 kJ mol^{-1}
- 31. Calculate the useful work done during the reaction

$$Ag(s) + \frac{1}{2}Cl_2(g) \rightarrow AgCl(s)$$

given that $E_{\text{Cl}_2/\text{Cl}}^0 = +1.36 \text{ V}$ and $E_{\text{AgCl, Ag, Cl}}^0 = +0.220 \text{ V}$ if $p_{\text{Cl}_2} = 1$ and T = 298 K.

(a) 110 kJ mol^{-1}

(b) 220 kJ mol⁻¹

(c) 55 kJ mol^{-1}

(d) 100 kJ mol^{-1}

32. Determine the electrode potential for

Pt,
$$H_2(g)$$
 | $H^+(aq)$
 $p_{H_2} = 1$ atm | $a_{H_1}^+ = 0.1$

(a) 0

(b) 0.0592 V

(c) -0.0592 V

(d) 0.1184 V

33. Calculate the equilibrium constant of the reaction

$$Cd^{2+}(aq) + Zn(s) \rightarrow Zn^{2+}(aq) + Cd(s)$$

if $E_{CA^{2+}/CA}^{0} = -0.403 \text{ V}$ and $E_{Zn^{2+}/Zn}^{0} = -0.763 \text{ V}$.

(a) $K = 1.45 \times 10^{12}$

(b) $K = 4.25 \times 10^{14}$

(c) $K = 1.45 \times 10^{18}$

(d) $K = 14.4 \times 10^{11}$

34. Given that at 25°C,

for
$$Cr^{3+}(aq) + e \rightarrow Cr^{2+}(aq)$$
, $E^0 = -0.424 \text{ V}$
 $Cr^{2+}(aq) + 2e \rightarrow Cr(s)$, $E^0 = -0.900 \text{ V}$
find E^0 at 25°C for $Cr^{3+}(aq) + 3e \rightarrow Cr(s)$.

- (a) -0.74 V
- (b) +0.74 V (c) -1.324 V (d) -0.476 V

35. Calculate the emf of the following cell at 25°C.

 $Ag(s) \mid AgNO_3(0.01 \text{ mol kg}^{-1}) \parallel AgNO_3(0.05 \text{ mol kg}^{-1}) \mid Ag(s)$

(a) -0.414 V

(b) 0.828 V

(c) 0.414 V

(d) 0.0414 V

36. Consider the given data.

Half-cell reaction

Standard reduction potential, E^0 (volts)

- 1. $Cr_2O_7^{2-} + 14H^+ + 6e \rightleftharpoons 2Cr^{3+} + 7H_2O$
- 1.33

2. $\operatorname{Cr}^{3+} + 3e \rightleftharpoons \operatorname{Cr}(s)$

-0.74

3. $Cu^+ + e \rightleftharpoons Cu(s)$

0.52

4. $Cu^{2+} + 2e \rightleftharpoons Cu(s)$

0.34

The numerical value of the standard cell potential for the reaction

$$2Cr^{3+} + 3Cu^{2+}(aq) \rightleftharpoons 2Cr^{3+}(aq) + 3Cu(s)$$

is

- (a) -1.08 V (b) -0.40 V (c) 1.08 V (d) 0.34 V

37. For the electrochemical cell

$$\overset{\ominus}{Ag}\mid AgCl(s)\text{, }KCl(aq)\parallel AgNO_{3}(aq)\mid \overset{\oplus}{Ag}$$

the overall cell reaction is

- (a) $Ag^+ + KCl(aq) \longrightarrow AgCl(s) + K^+$
- (b) $Ag(s) + AgCl(s) \longrightarrow 2Ag(s) + \frac{1}{2}Cl_2(g)$

(c)
$$AgCl(s) \longrightarrow Ag^+ + Cl^-$$

(d)
$$Ag^+ + Cl^- \longrightarrow AgCl(s)$$

- 38. When $\Delta G = -100$ kJ mol⁻¹ and n = 1, the potential of the cell is
 - (a) 2 V
- (b) 1 V
- (c) 10 V
- (d) 5 V

39. Given the half-cell reactions

(i) Fe²⁺(aq) + 2e
$$\longrightarrow$$
 Fe(s); $E^0 = -0.44 \text{ V}$

(ii)
$$2H^{+}(aq) + \frac{1}{2}O_{2}(g) + 2e \longrightarrow H_{2}O(1); \quad E^{0} = +1.23 \text{ V}$$

 E^0 for the reaction

$$Fe(s) + 2H^{+} + \frac{1}{2}O_{2}(g) \longrightarrow Fe^{2+}(aq) + H_{2}O(1)$$

is

(a)
$$+1.67 \text{ V}$$
 (b) -1.67 V (c) -0.77 V (d) $+0.77 \text{ V}$

40. Given the half-cell reactions

$$Cu^{+}(aq) + e \longrightarrow Cu(s)$$
, $E^{0} = +0.52 \text{ V}$
 $Cu^{2+}(aq) + e \longrightarrow Cu^{+}(aq)$, $E^{0} = +0.16 \text{ V}$

the equilibrium constant for the disproportionation reaction

$$2Cu^{+}(aq) \longrightarrow Cu(s) + Cu^{2+}(aq)$$
 at 298 K is

(a) 6×10^4

(b) 6×10^6

(c) 1.2×10^6

- (d) 1.2×10^{-6}
- 41. Which of the following statements are incorrect for an electrode concentration cell?
 - (a) The emf is usually small.
 - (b) The emf of the cell decreases continually during its operation as the concentrations in the two compartments approach each other.
 - (c) When the concentrations of the ions in the two compartments are the same, *E*(emf of cell) becomes zero.
 - (d) When the concentrations of the ions in the two compartments are the same, E^0 of the cell becomes zero.
- **42.** Which of the following statements is incorrect?
 - (a) E⁰ changes sign whenever a half-cell reaction is reversed.
 - (b) Changing the stoichiometric coefficients of a half-cell reaction does not affect the value of E^0 because electrode potential is an intensive property.
 - (c) The half-cell reactions are reversible.
 - (d) Changing the stoichiometric coefficients of a half-cell reaction does affect the value of E^0 because electrode potential is an extensive property.

43. The following concentration cell is a step-up cell.

 $Zn(s) | Zn^{2+}(aq, 0.20 \text{ M}) | KCl(sat'd) | Zn^{2+}(aq, 2.0 \text{ M}) | Zn(s)$

Choose the correct option.

- (a) Zn²⁺ reacts with Cl⁻ to form ZnCl₂.
- (b) Zn(s) reacts with K(s) to form an intermetallic compound.
- (c) Reduction occurs in the more concentrated compartment and oxidation takes place in the more dilute, according to Le Chatelier's principles
- (d) Oxidation takes place in the more concentrated compartment and reduction in the more dilute, following Le Chatelier's principles.
- 44. A quantity of 0.300 g of Cu was deposited from a CuSO₄ solution by passing a current of 3.00 A through it for 304 s. The value of the faraday constant is
 - (a) 9.66×10^4 C

(b) 9.66×10^3 C

(c) 9.66×10^5 C

- (d) 9.66×10^{2} C
- **45.** A solution containing H ⁺ and D ⁺ ions is in equilibrium with a mixture of H₂ and D₂ gases at 25°C. If the partial pressures of both the gases are 1.0 atm, calculate the ratio

$$[D^+]/[H^+]$$
 $(E_{D^+/D}^0 = -0.003 \text{ V}).$

(a) 1.2

(b) 1.1

(c) 0.11

- (d) 1.0
- 46. A galvanic cell is constructed as follows. A half-cell consists of a platinum wire immersed in a solution containing 1.0 M of Sn ⁺ and 1.0 M of Sn⁴⁺, and another half-cell has a thallium rod immersed in a 1.0 M solution of Tl +.

Given
$$\operatorname{Sn}^{4+}(aq) + 2e \longrightarrow \operatorname{Sn}^{2+}(aq); \quad E^0 = +0.13 \text{ V}$$

 $Tl^{4+}(aq) + e \longrightarrow Tl(s); E^{0} = -0.34 \text{ V},$

what is the cell voltage if the Tl + concentration is increased tenfold?

- (a) 0.411 V
- (b) 4.101 V
- (c) 0.492 V
- (d) 0.222 V
- 47. The cell formed by the two-redox couple ox_1/red_1 and ox_2/red_2 may be denoted by

$$red_1, ox_1 \parallel red_2, ox_2.$$

The cell reaction is

- (a) $\operatorname{red}_1 + \operatorname{ox}_1 \longrightarrow \operatorname{red}_2 + \operatorname{ox}_2$ (b) $\operatorname{red}_2 + \operatorname{ox}_2 \longrightarrow \operatorname{red}_1 + \operatorname{ox}_1$
- (c) $\operatorname{red}_1 + \operatorname{ox}_2 \longrightarrow \operatorname{ox}_1 + \operatorname{red}_2$ (d) $\operatorname{red}_1 + \operatorname{red}_2 \longrightarrow \operatorname{ox}_1 + \operatorname{ox}_2$
- 48. Given the cell reactions

$$AgCl(s) + e \longrightarrow Ag(s) + Cl^{-}(aq); \quad E^{0} = +0.22 \text{ V}$$

 $Ag^{+}(aq) + e \longrightarrow Ag(s); \quad E^{0} = +0.80 \text{ V}$

the solubility product of AgCl(s) at 298 K is

(a)
$$1.6 \times 10^{-18}$$

(b) 1.6×10^{-5}

(c)
$$1.6 \times 10^{-12}$$

(d) 1.6×10^{-10}

49. For the half-cell reaction

$$H^+(aq) + e \longrightarrow \frac{1}{2} H_2(g)$$

- (a) $E(H^+/H_2) = -59.2 \text{ mV} \times \text{pH}$
- (b) $E^{0}(H^{+}/H_{2}) = -59.2 \text{ mV} \times \text{pH}$
- (c) $E^{0}(H^{+}/H_{2}) = pH \log[H^{+}]$
- (d) $E(H^+/H_2) = -pH \log[H^+]$
- **50.** In an H_2 - O_2 fuel cell,
 - (a) the cell reaction is $2H_2O(1) \longrightarrow 2H_2(g) + O_2(g)$
 - (b) the cell reaction is $2H_2(g) + O_2(g) \longrightarrow 2H_2O(l)$
 - (c) the cell voltage is 2.0 V
 - (d) chemical energy is stored
- **51.** The unit of ionic mobility is

(a)
$$m^{-2}V^{-1}s^{-1}$$

(b) $m^2V^{-1}s^{-1}$

(c)
$$m^{-2}V s^{-1}$$

(d) $m^2V^{-2}s^{-1}$

- 52. Drift speed of an ion is directly proportional to
 - (a) the strength of the applied field
 - (b) the strength of the applied field and the charge on the ion
 - (c) the strength of the applied field, the viscosity and the charge on the ion
 - (d) the cell constant of the conductivity cell
- 53. Which of the following statements is incorrect?
 - (a) Specific conductance of an electrolytic solution decreases with dilution
 - (b) Conductance of an electrolytic solution increases with dilution
 - (c) Equivalent conductance of an electrolytic solution increases with dilution
 - (d) Conductance of an electrolytic solution decreases with dilution
- 54. Which of the following compounds shows maximum value of equivalent conductance in a fused state?
 - (a) MgCl₂
- (b) BeCl₂
- (c) CaCl₂
- (d) SrCl₂
- 55. Kohlrausch given the following relation for strong electrolyte

$$\Lambda = \Lambda_0 - B\sqrt{C}.$$

Which of the following equality holds?

- (a) $\Lambda = \Lambda_0$ as $c \to 1$
- (b) $\Lambda = \Lambda_0 \text{ as } c \to 0$
- (c) $\Lambda = \Lambda_0 \text{ as } c \to \infty$
- (d) $\Lambda = \Lambda_0 \text{ as } c \rightarrow \sqrt{B}$
- **56.** Which of the following statements is incorrect?
 - (a) There is no corelation between ionic size and conductance.
 - (b) Ions of normal fatty acids and ionic conductance attain a constant value.
 - (c) The ionic conductivity at infinite dilution is the highest for H⁺ ion.
 - (d) The ionic conductivity of OH ion at infinite dilution is greater than that of H + ion.
- **57.** The ionic conductance at an infinite dilution is related to
 - (a) the speed of the ions which move under the influence of an applied potential gradient
 - (b) the conductance ratio and the concentration of the solution
 - (c) equivalent and molar conductance
 - (d) weak van der Waals forces
- 58. Which of the following is arranged in order of increasing ionic conducrtance?
 - (a) $NH_4^+ < Ag^+ < Na^+ < Li^+$ (b) $Na^+ < NH_4^+ < Ag^+ < Li^+$
 - (c) $\text{Li}^+ < \text{Na}^+ < \text{Ag}^+ < \text{NH}_4^+$ (d) $\text{Ag}^+ < \text{Li}^+ < \text{Na}^+ < \text{NH}_4^+$
- 59. Which of the following compounds of 0.1 N concentration has smallest Λ_0 ?
 - (a) NaCl

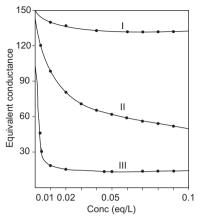
(b) NiSO₄

(c) CaCl₂

- (d) Na₂SO₄
- **60.** If ρ is the resistance in ohm of a centimeter cube, generally called the specific resistance of the substance constituting the conductor, the resistance *r* of the layer containing "a" cubes is given by
 - (a) $\frac{1}{r} = \frac{1}{0} + \frac{1}{0} + \dots$
- (b) $\frac{1}{r} = \frac{1}{\Omega a} + \frac{1}{\Omega a} + \dots$

(c) $r = a/\rho$

- (d) $r = \rho + \rho + ...$
- 61. A graph was plotted between the equivalent conductance of various electrolytes (CH₃COOH, NiSO₄ and HCl) and their concentrations in equivalents per litre.



which of the followings represents the correct setting?

- (a) I (NiSO₄), II (CH₃COOH), III (HCl)
- (b) I (HCl), II (NiSO₄), III (CH₃COOH)
- (c) I (CH₃COOH), II (HCl), III (NiSO₄)
- (d) I (HCl), II (CH₃COOH), III (NiSO₄)
- **62.** Which of the following statements is incorrect?
 - (a) Increase in temperature increases the equivalent conductance.
 - (b) Conductance ratio decreases with increase in temperature.
 - (c) The solvent with low dielectric constant and the equivalent conductance decrease with increase in concentration.
 - (d) The solvent with high dielectric constant and the equivalent conductance decrease with increase in concentration.
- **63.** The expression $\alpha = \Lambda/\Lambda_0$ will not hold good for the electrolyte
 - (a) HOCN

(b) H_2SO_4

(c) CH₃CH₂OH

(d) CH₃CO₂H

- **64.** The mathematical expression for law of independent migration of ions and Ostwald's dilution law are given by
 - (a) $\Lambda = \Lambda_m^0 BC^{1/2}$

(b) $\Lambda_0 = F(U_+ + U_-)$

(c) $\Lambda_m^0 = v_+ \lambda_+ + v_- \lambda_1$

(d) $\frac{\Lambda_0}{\Lambda_m} = \frac{1}{\Lambda_m^0} + \frac{\Lambda_m c}{K_a(\Lambda_m^0)^2}$

- **65.** The conductance of a solution of an electrolyte is equal to that of its specific conductance. The cell constant of the conductivity cell is equal to
 - (a) resistance

(b) faraday

(c) zero

(d) unity

- 66. The equivalent conductance of an acetic acid solution of concentration 0.01 M is 1.65 m S m² mol⁻¹ at 298 K. The dissociation constant of acetic acid at this temperature is (given, $\Lambda_0 = 39.0 \text{ m S m}^2 \text{ mol}^{-1}$)
 - (a) 1.8×10^{-4} (b) 4.0×10^{-10} (c) 1.8×10^{-5} (d) 1.8×10^{-6}

- 67. The ionic mobility of H⁺ ions are much higher than the other ions. These high values are the results of hydrogen bonding. In water, the proton is hydrated and its movement can be represented as

(a)
$$\stackrel{\text{H}}{\longrightarrow} O - H \cdots O - H \cdots H - O \cdots H - O$$

(b)
$$\stackrel{\text{H}}{\longrightarrow} O - H \cdots O - H \cdots H - O \cdots H - O$$

(c)
$$\stackrel{\text{H}}{\longrightarrow} O - \stackrel{\text{H}}{\longrightarrow} O -$$

- **68.** In the equation $\Lambda = \Lambda_m^0 Bc^{1/2}$, the constant *B* depends upon
 - (a) $c^{1/2}$

(b) stoichiometry of the electrolyte

(c) resistance

- (d) specific conductivity
- 69. Which of the following is arranged in increasing order of ionic mobility?
 - (a) $I^- < Br^- < Cl^- < F^-$
- (b) $F^- < Cl^- < Br^- < I^-$
- (c) $F^- < I^- < CI^- < Br^-$ (d) $F^- < CI^- < I^- < Br^-$
- 70. Ionic mobility principle (utilized for purifying and identifying proteins) is based on
 - (a) electroosmosis

(b) electrophoresis

(c) electrolysis

- (d) isoelectronic point
- **71.** If the voltage across the electrode is 1 V, then $c(\lambda_0^+ + \lambda_0^-)$ represents the
 - (a) current

(b) molar conductance

(c) conductance

(d) ionic mobility

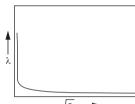
- 72. The specific conductance (κ) is given by
 - (a) $(\lambda_0^+ + \lambda_0^-)$

(b) $Vc(\lambda_0^+ + \lambda_0^-)$

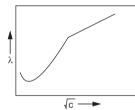
(c) $c(\lambda_0^+ + \lambda_0^-)$

- (d) $F(u_+ + u_-)$
- 73. The specific conductance of a saturated AgCl solution is found to be $1.86\times10^{-6}\,\mathrm{S}\,\mathrm{cm}^{-1}$ and that for water is $6.0\times10^{-8}\,\mathrm{S}\,\mathrm{cm}^{-1}$. The solubility of AgCl is $(\Lambda_0=137.2\,\mathrm{S}\,\mathrm{equiv}^{-1}\,\mathrm{cm}^2)$
 - (a) $1.7 \times 10^{-3} \text{ mol L}^{-1}$
- (b) $1.3 \times 10^{-5} \text{ mol L}^{-1}$
- (c) $1.3 \times 10^{-4} \text{ mol L}^{-1}$
- (d) $1.3 \times 10^{-6} \text{ mol L}^{-1}$
- **74.** Which of the following solutions of KCl will have the highest value of specific conductance?
 - (a) 0.01 M
- (b) 0.1 M
- (c) 1.0 M
- (d) 0.5 M
- **75.** The variation of equivalent conductance of a weak electrolyte with $(concentration)^{1/2}$ is represented as

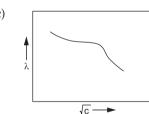
(a)



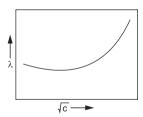
(b)



(c)



(d)



- **76.** The equivalent conductance of strong electrolyte increases with dilution because of
 - (a) intermolecular interaction between the ions
 - (b) increase in number of ions per unit volume
 - (c) increase in degree of dissociation
 - (d) lower enthalpy of hydration
- 77. The ionic mobility of ions at infinite dilution is related to ionic conductance by
 - (a) $\Lambda_0 = F\kappa$

(b) $\Lambda_0 F = U_+ + U_-$

(c) $\Lambda_0 = U_+ + U_-$

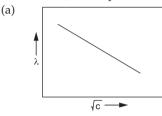
(d) $\Lambda_0 = F(U_+ + U_-)$

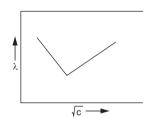
- 78. The increase in equivalent conductance of a weak electrolyte with dilution is due to
 - (a) increase in degree of dissociation and decrease in ionic mobility
 - (b) decrease in degree of dissociation and decrease in ionic mobility
 - (c) increase in degree of dissociatin and increase in ionic mobility
 - (d) decrease in degree of dissociation and increase in ionic mobility

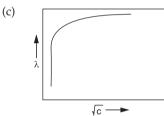
(b)

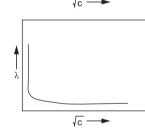
(d)

79. The variation of equivalent conductance of strong electrolyte with $(concentration)^{1/2}$ is represented by









- **80.** The relation among conductance (κ) , specific conductance (G) and cell constant (l/A) is
 - (a) $G = \kappa \frac{l}{A}$ (b) $G = \kappa \frac{A}{l}$ (c) $G\kappa = \frac{l}{A}$ (d) $G = \kappa Al$

- 81. During conductivity measurement (based on Wheatstone bridge principle) using conductivity cell, alternating current (AC) is used because a direct current would lead to
 - (a) association and ionization
 - (b) electrolysis and polarization
 - (c) polymerization and polarization
 - (d) polarization
- 82. During conductivity measurement we use platinized platinum electrodes which
 - (a) catalyze the union of hydrogen and oxygen which tend to be liberated by the successive pulses of the current
 - (b) eliminates the polarization of emf
 - (c) both (a) and (b)
 - (d) none of these

- **83.** The value of molar conductance of HCl is greater than that of NaCl at a given temperature and concentration because
 - (a) ionic mobility of HCl is greater than that of NaCl
 - (b) the dipole moment of NaCl is greater than that of HCl
 - (c) NaCl is more ionic than HCl
 - (d) HCl is Bronsted acid and NaCl is a salt of a strong acid and strong base
- 84. Which of the following statements is correct?
 - (a) The conductivity of a solution decreases with increasing viscosity and ion size.
 - (b) Molar conductivities of the alkali metal ions decrease from Li $^{\scriptscriptstyle +}$ to Cs $^{\scriptscriptstyle +}$
 - (c) Stokes formula is given by $f = 6\pi\eta Ga$
 - (d) According to the Grotthuss mechanism, there is an effective motion of a proton and neutron that involves the rearrangement of bonds in a group of water moelcules.
- 85. Aniline hydrochloride ($C_6H_5NH_3Cl$) is a salt of a weak base and strong acid. The hydrolysis equilibria

$$K_h = \frac{h^2 c}{(1-h)}.$$

On the basis of conductivity, h is given by

(a)
$$h = \frac{\Lambda - \Lambda_1}{\Lambda_2 - \Lambda}$$

(b)
$$h = \frac{\Lambda - \Lambda_1}{\Lambda_2 - \Lambda_1}$$

(c)
$$h = \frac{\Lambda_2 + \Lambda_1}{\Lambda}$$

(d)
$$h = \frac{\Lambda + \Lambda_1}{\Lambda_2 - \Lambda_1}$$

Where Λ , Λ_1 , Λ_2 are equivalent conductance of hydrolysed salt, unhydrolysed salt and HCl respectively, K_h = hydrolysis constant, c = concentration of $C_6H_5NH_3Cl$ and h = degree of hydrolysis.

• *Type 2* •

Choose the correct options. More than one option is correct.

- 86. Which of the following statements is correct for ionic mobility?
 - (a) It depends upon the voltage across the electrodes.
 - (b) Lower the voltage, higher is the velocity.
 - (c) Ionic velocity per unit electric field strength is constant.
 - (d) All of these.

- 87. Which of the following units is correctly matched?
 - (a) SI units of conductivity $(k) \rightarrow$ siemens per metre (S m⁻¹)
 - (b) SI units of molar conductivity \rightarrow siemens squared per mol (S m² mol⁻¹)
 - (c) SI unit of ionic mobility \rightarrow m V⁻¹s⁻¹
 - (d) All of these
- 88. According to Grotthuss model
 - (a) there is no coordinated motion of a proton along a chain of water molecules.
 - (b) there is a rapid hopping between neighbouring sites of water molecules.
 - (c) the system H +(H2O)4 has low activation energy
 - (d) none of these
- 89. Oxygen and hydrogen gas are produced at the anode and cathode during the electrolysis of dilute aqueous solutions of
 - (a) Na_2SO_4

(b) AgNO₃

(c) H_2SO_4

- (d) NaOH
- 90. Which of the following statements are correct?
 - (a) The electrolysis of aqueous NaCl produces hydrogen gas at the cathode and chlorine gas at the anode.
 - (b) The electrolysis of a dilute solution of sodium fluoride produces oxygen gas at the anode and hydrogen gas at the cathode.
 - (c) The electrolysis of concentrated sulphuric acid produces SO₂ gas at the anode and O₂ gas at the cathode.
 - (d) After the electrolysis of aqueous sodium sulphate, the solution becomes acidic.
- **91.** Which of the following cell reactions correctly represent the electrolysis of water?
 - (a) $2H^+ + 2e \rightarrow H_2(g)$
- (b) $2H_2O + 2e \rightarrow H_2(g) + 2(OH^-)$
- (c) $4(OH^{-}) \rightarrow 2H_{2}O + O_{2}(g) + 4e$ (d) $2H_{2}O \rightarrow O_{2}(g) + 4H^{+} + 4e$
- **92.** Which of the following statements are correct?
 - (a) The electrolysis of concentrated H_2SO_4 at 0–5°C using a Pt electrode produces $H_2S_2O_8$.
 - (b) The electrolysis of a brine solution produces NaClO₃ and NaClO.
 - (c) The electrolysis of a CuSO₄ solution using Pt electrodes causes the liberation of O₂ at the anode and the deposition of copper at the cathode.
 - (d) All electrolytic reactions are redox reactions.

- 93. How much charge must be supplied to a cell for the electrolytic production of 245 g NaClO₄ from NaClO₃? Because of a side reaction, the anode efficiency for the desired reaction is 60%.
 - (a) 6.43×10^5 C

(b) 6.67 F

(c) 6.43×10^6 C

- (d) 66.67 F
- 94. In an electrochemical process, a salt bridge is used
 - (a) to maintain electroneutrality in each solution
 - (b) to complete the circuit so that current can flow
 - (c) as an oxidizing agent
 - (d) as a colour indicator
- **95.** Consider the cell

$$Cd(s) \mid Cd^{2+}(1.0 \text{ M}) \parallel Cu^{2+}(1.0 \text{ M}) \mid Cu(s)$$

If we wish to make a cell with a more positive voltage using the same substances, we should

- (a) increase $[Cd^{2+}]$ as well as $[Cu^{2+}]$ to 2.0 M
- (b) reduce only [Cd²⁺] to 0.1 M
- (c) increase only [Cu²⁺] to 2.0 M
- (d) decrease $[Cd^{2+}]$ as well as $[Cu^{2+}]$ to 0.1 M
- **96.** A concentration cell is a galvanic cell in which
 - (a) the electrode material and the solutions in both half-cells are composed of the same substances
 - (b) only the concentrations of the two solutions differ
 - (c) $\Delta E_{coll}^0 = 0$
 - (d) the Nernst equation reduces to $\Delta E_{\text{cell}} = -\left(\frac{0.0592}{n}\right) \log Q$ at 25°C
- 97. Which of the following statements are correct?
 - (a) A reaction is spontaneous from left to right if $K_{eq} > Q$, in which case $\Delta E_{\text{cell}} > 0$.
 - (b) A reaction occurs from right to left if $K_{eq} < Q$, in which case $\Delta E_{\rm cell} < 0$.
 - (c) If the system is at equilibrium, no net reaction occurs.
 - (d) ΔE_{cell} is temperature-independent.
- 98. Which of the following are concentration cells?
 - (a) $Pt, H_2(g) \mid HCl \mid H_2(g), Pt$ (b) $Cd, (Hg) \mid Cd^{2+} \mid (Hg), Cd^{2+} \mid (Hg), Cd^{2+} \mid (Hg) \mid Cd^{2+} \mid (Hg), Cd^{2+}$

(c)
$$Zn(s) \mid Zn^{2+} \mid Cu^{2+} \mid Cu$$

(d)
$$\operatorname{Ag}(s)$$
, $\operatorname{AgCl}(s) \mid \operatorname{HCl} \mid \operatorname{HCl} \mid \operatorname{AgCl}(s)$, $\operatorname{Ag}(s)$

Which of the following expressions represent the emf of the above cell at 25°C?

(a)
$$E = \frac{0.0592}{2} \log \frac{(a_{Pb}^{2+})_2}{(a_{Pb}^{2+})_1}$$

(b)
$$E = \frac{0.0592}{2} \log \frac{(a_{Pb}^{2+})_1}{(a_{Pb}^{2+})_2}$$

(c)
$$E = \frac{0.0592}{2} \log \frac{\left[K_{sp}(PbI_2)\right]^{1/3}}{\left[K_{sp}(PbSO_4)\right]^{1/2}}$$

(d)
$$E = \frac{0.0592}{2} \log \frac{K_{sp}(PbI_2)}{K_{sp}(PbSO_4)}$$

100. Which of the following represent electrodes of the second kind?

- (a) $Ag(s) | AgCl(s), Cl^{-}$
- (b) $Cu(s) | CuSO_4(aq), SO_4^{2-}$
- (c) $Hg(l) \mid Hg_2Cl_2(s), Cl^-$ (d) $Pb(s) \mid PbSO_4(s), SO_4^{2-}$

Answers

1. c	2. d	3. b	4. c	5. d
6. b	7. a	8. d	9. d	10. a
11. a	12. b	13. b	14. c	15. d
16. a	17. b	18. c	19. c	20. b
21. d	22. c	23. a	24. a	25. a
26. b	27. a	28. d	29. b	30. c
31. a	32. c	33. a	34. a	35. a
36. c	37. d	38. b	39. a	40. c
41. c	42. d	43. c	44. a	45. b
46. a	47. c	48. d	49. a	50. b
51. b	52. a	53. d	54. b	55. b
56. d	57. a	58. c	59. b	60. a
61. b	62. d	63. b	64. c	65. d

66. c	67. a	68. b	69. d	70. b
71. a	72. c	73. b	74. c	75. a
76. b	77. d	78. c	79. a	80. b
81. b	82. c	83. a	84. a	85. b
86. a, c	87. a, b	88. a, b, c	89. a, b, c, d	90. a, b
91. b, c	92. a, b, c, d	93. a, b	94. a, b	95. b, c
96. a, b, c, d	97. a, b, c	98. a, b, d	99. a, c	100. a, c, d

Hints to More Difficult Problems

- K₂SO₄ is a salt of a strong acid and strong base. So the electrolysis of K₂SO₄(aq) is the electrolysis of water.
- 4. $BrO_3^- + 6H^+ + 6e \longrightarrow Br^- + 3H_2O$
- 7. Energy = $0.1 \times 150 \times 100 \text{ J} = 1.5 \text{ kJ}$.
- **10.** 1 equivalent = $11200 \text{ mL mol}^{-2} = 1 \text{ F mol}^{-1} = 96500 \text{ C mol}^{-1}$.

As we know Q = nF where n = no. of moles.

$$Q = \frac{1 \text{ mL}}{11200 \text{ mL mol}^{-1}} \times 96500 \text{ C mol}^{-1}$$
$$= 8.6 \text{ C}.$$

13. After electrolysis aqueous NaCl is converted into aqueous NaOH.

The quantity of electricity passed

$$= \frac{0.250 \times 35 \times 60}{96500 \text{ C}} \text{ A s} = 5.44 \times 10^{-3} \text{ F}$$

The number of equivalents of OH ion formed

$$=5.44\times10^{-3}$$
.

:. Molarity of NaOH =
$$\frac{5.44 \times 10^{-3}}{0.400 \text{ L}}$$
 eq.
= 1.36×10^{-2} .

$$pOH = -log(1.36 \times 10^{-2}) = 2.00 - 0.13 = 1.87.$$

$$\therefore$$
 pH = 12.13.

16.
$$W = \frac{A}{nF} It$$
 or $\rho \times V = \frac{A}{nF} Q$.

$$Q = \frac{\rho \times \text{area} \times \text{thickness}}{A} \times nF$$

$$=\frac{8.94\times10\times10\times10^{-2}\times2\times96500}{63.6}=2.7\times10^{-4}\text{ C}.$$

18.
$$2H_2O \longrightarrow 2H_2(g) + O_2(g)$$

$$3x = 168$$
.

$$x = 56 \text{ mL}.$$

$$V_{\rm H_2} = 2x = 112 \text{ mL}.$$

$$V_{O_2} = x = 56 \text{ mL}.$$

11200 mL of H_2 at stp = 1 F.

112 mL of H_2 at stp = 0.01 F.

5600 mL of O_2 at stp = 1 F.

56 mL of O_2 at stp $\equiv 0.01$ F.

Amount of electricity passed = $0.01 \text{ F} = 0.01 \times 96500 = 965 \text{ C}$.

We get identical results whether we consider H₂ or O₂.

20. The cell is represented as

$$Mg(s) | Mg^{2+} || Cu^{2+} | Cu(s)$$

$$E^{0} = E_{Cu^{2+}}^{0} | Cu - E_{Mg^{2+}}^{0} | Mg$$

$$= 0.34 - (-2.37) = 2.71 \text{ V}.$$

28. The cell reaction is

$$Zn(s) \longrightarrow Zn^{2+}(0.01) M$$

 $[Ag^{+}(1.25 M) + e \longrightarrow Ag(s)] \times 2$

$$Zn(s) + 2Ag^{+}(1.25) M \rightarrow Zn^{2+}(0.01 M) + 2Ag(s)$$

$$Q = \frac{[Zn^{2+}]}{[Ag^+]^2} = \frac{0.01}{(1.25)^2} = 6.4 \times 10^{-3}.$$

29.
$$\Delta G^0 = -nE^0 F = -RT \ln K_{eq}$$

 $\Rightarrow \ln K_{eq} = \frac{nE^0 F}{RT} = \frac{nE^0}{RT/F}$.
 $\log K_{eq} = \frac{nE^0}{2.303 \frac{RT}{F}} = \frac{nE^0}{0.0592}$ (at 25°C)

$$=\frac{2\times1.20}{0.0592}=40.5.$$

31. For the cell reaction

$$Ag(s) + \frac{1}{2}Cl_2(g) \longrightarrow AgCl(s)$$

$$E^{0} = -1.14 \text{ V}$$

or
$$E = E^0 - \frac{0.0592}{1} \log [Cl_2]^{\frac{1}{2}}$$
.

In standard conditions, $p_{Cl_2} = 1$.

$$\therefore \log[Cl_2]^{\frac{1}{2}} = 0.$$

Useful work =
$$-W_{\text{max}} = \Delta G = -nEF$$

= $(-1) \times (-1.14) \times 96500 \times 10^{-3} \text{ kJ mol}^{-1}$
= 110 kJ mol^{-1} .

34. Adding the first two reactions, we get the third equation and using the free-energy concept, we have

$$\Delta G_1^0 + \Delta G_2^0 = \Delta G_3^0 \quad (n = \text{no. of electrons involved}).$$

$$-n_1 E_1^0 F - n_2 E_2^0 F = -n_3 E_3^0 F.$$

$$E_3^0 = \frac{n_1 E_1^0 + n_2 E_2^0}{n_3} = \frac{1 \times (-0.424) + 2 \times (-0.900)}{3}$$

$$= -0.741 \text{ V}.$$

35. It is a concentration cell. The emf of a concentration cell is

E = +0.0592 log
$$\frac{C_1}{C_2}$$
 at 25°C and $n = 1$
= +0.0592 log $\frac{0.01}{0.05}$ = -0.414 V.

- **39.** Equation (ii) equation (i) = 1.67 V
- **40.** $E^0 = 0.52 0.16 = 0.36 \text{ V}.$

$$\log K_{\text{eq}} = \frac{nE^0}{0.0592} = \frac{1 \times 0.36}{0.0592} = 6.081.$$

$$K_{eq} = 1.20 \times 10^{6}$$
.

46. The cell is represented as

$$\begin{array}{c} -0.34\,\text{M} & +0.13\,\text{M} \\ Tl(s) \mid Tl^+(1.0\,M) \mid \mid Sn^{\,4+}(1.0\,M), Sn^{\,2+}(1.0\,M) \mid Pt \\ \end{array}$$
 The cell reaction is
$$\begin{array}{c} Tl(s) & \longrightarrow Tl^+ + e \times 2 \\ & Sn^{\,4+} + 2e & \longrightarrow Sn^{\,2+} \\ \end{array}$$
 Overall reaction
$$\begin{array}{c} 2Tl(s) + Sn^{\,4+} & \longrightarrow 2Tl^+ + Sn^{\,2+} \\ \end{array}$$

$$E = (E_{\text{Right}}^{0} - E_{\text{Left}}^{0}) - \frac{0.0592}{2} \log \frac{[\text{TI}^{+}]^{2}[\text{Sn}^{2+}]}{[\text{Sn}^{4+}]}$$

$$= 0.47 \text{ V} - 0.0296 \log (10)^{2} \quad [\text{TI}^{+} \text{ concentration increases tenfold}]$$

$$= 0.47 - 0.0592 = 0.411 \text{ V}.$$

48.
$$\log K_{\rm sp} = \frac{nE^0}{0.0592}$$

= $\frac{1 \times (-0.58)}{0.0592} = -9.797 = \overline{10}.203$ (here $E^0 = -0.58$ V and $n = 1$).

$$K_{\rm sp} = 1.6 \times 10^{-10}$$
.

49.
$$E_{\text{H}^+/\text{H}_2} = E_{\text{H}^+/\text{H}_2}^0 - \frac{0.0592}{1} \log \frac{p_{\text{H}_2}^{\frac{1}{2}}}{[\text{H}^+]}$$

= 0 + 0.0592 log [H^+] = -0.0592 pH
= -59.2 mV × pH.

- 51. Ionic velocity is in ms $^{-1}$ and electric field strength is V m $^{-1}$, so ionic mobility is (m s $^{-1}$)/V m $^{-1}$ or m 2 s $^{-1}$ V $^{-1}$
- 54. BeCl₂ is covalent
- 56. Smallest ion possesses maximum mobility
- 58. Water molecules carries in its hydration sphere based on Stokes formula
- **59.** Bi-bivalent nature
- 61. Uni-univalent strong electrolyte
- 64. H₂SO₄ is a strong electrolyte

67.
$$K = \frac{c\Lambda^2}{\Lambda_0(\Lambda_0 - \Lambda)}$$

 $\Lambda = 1.65 \text{ m S m}^2 \text{mol}^{-1}, \text{ c} = 0.01 \text{ M}$ $\Lambda_0 = 39.05 \text{ m S m}^2 \text{mol}^{-1}$

72. Electrical property of colloids

73.
$$I = \frac{V}{R} = VG = \frac{V \kappa A}{l} = V \kappa = V c (\lambda_0^+ + \lambda_0^-)$$

75.
$$S(\text{Solubility}) = \frac{1000(\kappa_{\text{Agcl}} - \kappa_{\text{H}_2\text{O}})}{\Lambda_0} = \frac{1000(1.86 \times 10^{-6} - 6.0 \times 10^{-8})}{137.2}$$

= 1.3 × 10⁻⁵ mol L⁻¹

79.
$$F(U_+ + U_-)c = G\kappa = c(\lambda_0^+ + \lambda_0^-)$$

 $F(U_+ + U_-) = \lambda_0^+ + \lambda_0^- = \Lambda_0$

84. Large surface area of the finely divided platinum.

93.
$$ClO_4^- + 2H^+ + 2e \longrightarrow ClO_3^- + H_2O$$

Eq. wt. of $NaClO_4 = \frac{23 + 35 + 64}{2} = 61.25$.

No. of equivalents of NaClO₄ = $\frac{245}{61.25}$ = 4 = 4.0 F.

The anode efficiency is 60%.

No. of faradays =
$$\frac{4.0}{60} \times 100 = 6.67$$
 F.

$$6.67 \text{ F} = 6.67 \times 9.65 \times 10^{4} \text{ C} = 6.43 \times 10^{5} \text{ C}.$$

16

Colligative Properties of Solutions

• Type 1 •

1. The relative lowering of the vapour pressure of an aqueous solution containing a nonvolatile solute is 0.0125. The molality of the solution is

(b) 0.50 (d) 0.40

> 0.075 M CuSO₄

 $> 0.05 \text{ M KNO}_3$

 $> 0.05 \text{ M KNO}_3$

 $> 0.04 \text{ M BaCl}_2$

Choose the correct option. Only one option is correct.

(a) 0.70

(c) 0.80

2.	The v	apour pressure of a deliquescer	nt su	ıbstance is
	(a)	equal to the atmospheric pres	sure	
	(b)	equal to that of the water vapor	our i	n air
	(c)	greater than that of the water	vapo	our in air
	(d)	less than that of the water vap	our	in air
3.	Which	n of the following aqueous solu	ition	s has the highest boiling point?
	(a)	0.1 M KNO_3	(b)	$0.1 \text{ M Na}_3 PO_4$
	(c)	0.1 M BaCl ₂	(d)	$0.1 \text{ M K}_2\text{SO}_4$
4.	Which	O .	inge	d in order of decreasing freezing

(a) $0.05 \text{ M KNO}_3 > 0.04 \text{ M CaCl}_2 > 0.140 \text{ M sugar}$

(b) $0.04 \text{ M BaCl}_2 > 0.140 \text{ M sucrose} > 0.075 \text{ M CuSO}_4$

(c) $0.075 \text{ M CuSO}_4 > 0.140 \text{ M sucrose} > 0.04 \text{ M BaCl}_2$

(d) $0.075 \text{ M CuSO}_4 > 0.05 \text{ M NaNO}_3 > 0.140 \text{ M sucrose}$

Chen	mstry MCQ
Among the following, the so pressure is	olution which shows the highest osmotic
(a) 0.05 M NaCl	(b) 0.10 M BaCl ₂
(c) 0.05 M FeCl_3	(d) $0.05 \text{ M Na}_2\text{SO}_4$
$C_{12}H_{22}O_{11}$ will be in the order (a) NaCl > $C_{12}H_{22}O_{11}$ > BaC	2
	Among the following, the so pressure is (a) 0.05 M NaCl (c) 0.05 M FeCl ₃ The osmotic pressures of each

- (c) $NaCl > BaCl_2 > C_{12}H_{22}O_{11}$
- (d) $C_{12}H_{22}O_{11} > NaCl > BaCl_2$
- 7. 10 g of glucose (π_1) , 10 g of urea (π_2) and 10 g of sucrose (π_3) are dissolved in 250 mL of water at 300 K $(\pi = \text{osmotic pressure of solution})$. The relationship between the osmotic pressures of the solutions is
 - (a) $\pi_1 > \pi_2 > \pi_3$

(b) $\pi_3 > \pi_1 > \pi_2$

(c) $\pi_2 > \pi_1 > \pi_3$

- (d) $\pi_2 > \pi_3 > \pi_1$
- 8. When mercuric iodide is added to an aqueous solution of KI, the
 - (a) freezing point is raised
 - (b) freezing point is lowered
 - (c) boiling point does not change
 - (d) freezing point does not change
- **9.** Which of the following pairs of solutions can be expected to be isotonic at the same temperature?
 - (a) 0.1 M urea and 0.1 M NaCl
 - (b) 0.1 M urea and 0.2 M MgCl₂
 - (c) 0.1 M NaCl and 0.1 M Na₂SO₄
 - (d) 0.1 M Ca(NO₃)₂ and 0.1 M Na₂SO₄
- **10.** If a solute undergoes dimerization and trimerization, the minimum values of the van't Hoff factors are
 - (a) 0.50 and 1.50

(b) 1.50 and 1.33

(c) 0.50 and 0.33

- (d) 0.25 and 0.67
- 11. At 323 K, the vapour pressure in millimetres of mercury of a methanol-ethanol solution is represented by the equation $p = 120X_A + 140$, where X_A is the mole fraction of methanol. Then the

value of
$$\lim_{X_A \to 1} \frac{p_A}{X_A}$$
 is

(a) 120 mm

(b) 140 mm

(c) 260 mm

(d) 20 mm

12. The van't Hoff factors i for an electrolyte which undergoes dissociation

13. A compound X undergoes tetramerization in a given organic solvent.

(c) 0.125

(d) 2.0

and association in solvents are respectively
(a) greater than 1 and greater than 1
(b) less than 1 and greater than 1
(c) less than 1 and less than 1
(d) greater than 1 and less than 1

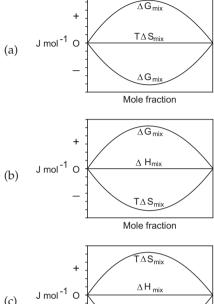
(b) 0.25

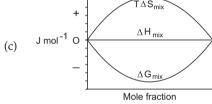
The van't Hoff factor *i* is

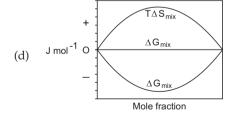
(a) 4.0

14.	4. A solution containing $0.8716 \text{ mol } L^{-1}$ of sucrose at 298 K is iso-osmotic with a solution of sodium chloride containing $0.5 \text{ mol } L^{-1}$ of NaCl. The degree of dissociation of NaCl is			
	(a)	0.743	(b)	0.894
	(c)	0.876	(d)	0.943
15.	Calcu 40.685	late the ebullioscopic constant 5 kJ mol^{-1} .	for w	vater. The heat of vaporization is
	(a)	$0.512~\mathrm{K~kg~mol}^{-1}$	(b)	1.86 K kg mol ⁻¹
	(c)	5.12 K kg mol ⁻¹	(d)	3.56 K kg mol ⁻¹
16.	solution 0.17 H	on containing 100 g of benz	ene	stance if the freezing point of a and 0.2 g of the substance is oscopic constant of benzene is
	(a)	70.46	(b)	85.66
	(c)	60.23	(d)	178.25
17.	An $K_b^{H_2O} =$ point of	equeous solution of NaCl = $0.512 \text{ K kg mol}^{-1}$ and $K_f^{\text{H}_2\text{O}} = 1.00 \text{ of this solution is}$	free 86 K	zes at -0.186°C. Given that kg mol ⁻¹ , the elevation in boiling
	(a)	0.0585 K	(b)	0.0512 K
	(c)	1.864 K	(d)	0.0265 K
18.	is 1.39 substa	$95 \mathrm{K}$ and that in the freezing poince is $1.280 \mathrm{K}$. Explain the diff	int c feren	
	(a)	solution	soiut	ion as well as in the benzene
	(b)	forms complexes in solution		
	(c)	associates in the benzene solu	ution	
	(d)	dissociates in the aqueous solution	solu	ation and not in the benzene

19. Which of the following represents correctly the changes in thermodynamic properties during the formation of 1 mol of an ideal binary solution.







- 20. Which of the following statements is correct for a binary solution?
 - (a) A solution in which heat is evolved exhibits positive deviations from Raoult's law.
 - (b) A solution in which heat is absorbed shows negative deviations from Raoult's law.
 - (c) When one component in solution shows negative deviation from Raoult's law, the other exhibits positive deviation.

- (d) When one component in solution shows positive deviation from Raoult's law, so does the other.
- **21.** The normal boiling point of toluene is 110.7°C, and its boiling point elevation constant is 3.32 K kg mol⁻¹. The enthalpy of vapourization of toluene is nearly
 - (a) 17.0 kJ mol^{-1}

(b) 34.0 kJ mol⁻¹

(c) 51.0 kJ mol^{-1}

- (d) 68.0 kJ mol^{-1}
- 22. Consider the following statements.
 - 1. Isotonic solutions have the same molar concentration at a given temperature.
 - 2. The molal elevation constant K_b is characteristic of a solvent, and is independent of the solute added.
 - 3. The freezing point of a 0.1 M aqueous KCl solution is more than that of a 0.1 M aqueous $AlCl_3$ solution.

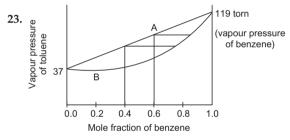
Which of these statements is correct.

(a) 1 and 2

(b) 2 and 3

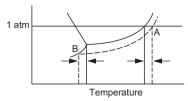
(c) 1 and 3

(d) 1, 2, and 3



Choose the correct option.

- (a) A represents vapour composition and B liquid composition.
- (b) *A* as well as *B* prepresent liquid composition.
- (c) Both *A* and *B* represent vapour composition
- (d) A represents liquid composition and B vapour composition.
- **24.** The phase diagrams for a pure solvent (represented by the solid line) and a corresponding solution (containing a nonvolatile solute and represented by the dashed lines) are shown below.

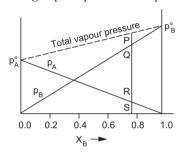


Choose the correct option.

- (a) $A = \Delta T_b m$ and $B = \Delta T_f m$
- (b) $A = \Delta T_f m$ and $B = \Delta T_b m$
- (c) $A = \Delta T_f$ and $B = \Delta T_b$
- (d) $A = \Delta T_b$ and $B = \Delta T_f$

where $T_{\rm f}$, $T_{\rm b}$ and m stand for freezing-point temperatures, boiling-point temperature and molality.

25. Consider the following vapour-pressure composition graph.



SP is equal to

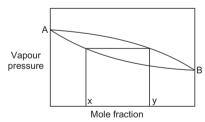
(a)
$$PQ + RS$$

(b)
$$PQ + QR + RS$$

(c)
$$SR + SQ$$

(d)
$$PQ + QR$$

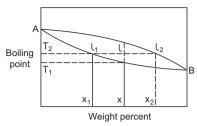
26. Consider the following graph.



Which of the following statements is correct?

- (a) A liquid mixture of composition x is in equilibrium with a vapour mixture of composition y.
- (b) A liquid mixture of composition x is in equilibrium with a vapour mixture of composition x.
- (c) $x = y^2$
- (d) $y = x^2$

27. Consider the following graph pertaining to distillation.



The distillate-to-residue ratio is

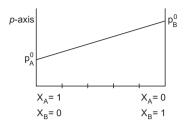
- (a) $\frac{l_1}{l_1 + l_2}$ (b) $\frac{lx_1}{lx_2}$
- (c) $\frac{ll_1}{ll_2}$
- (d) $\frac{ll_2}{ll_1}$

• *Type 2* •

Choose the correct options. More than one option is correct.

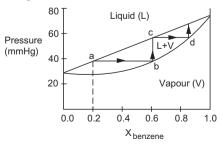
- 28. When a solute is added to a pure solvent, the
 - (a) vapour pressure of the solution becomes lower than that of the pure solvent
 - (b) rate of evaporation of the pure solvent is reduced
 - solute does not affect the rate of condensation
 - (d) rate of evaporation of the solution is equal to the rate of condensation of the solution at a lower vapour pressure than that in the case of the pure solvent
 - 29. According to Raoult's law the relative decrease in the solvent vapour pressure over the solution is equal to
 - (a) the mole fraction of the solvent
 - (b) the mole fraction of the solute
 - (c) the number of moles of the solute
 - (d) i times the mole fraction of the solute which undergoes dissociation or association in the solvent (i = van't Hoff factor)
 - **30.** Which of the following combinations are correct for a binary solution, in which the solute as well as the solvent are liquid?
 - (a) C_6H_6 and $C_6H_5CH_3$; $\Delta H_{soln} > 0$; $\Delta V_{sol} = 0$
 - (b) CH_3COCH_3 and $CHCl_3$; $\Delta H_{soln} < 0$; $\Delta V_{sol} < 0$
 - (c) H_2O and HCl; $\Delta H_{soln} > 0$; $\Delta V_{sol} < 0$
 - (d) H_2O and C_2H_5OH ; $\Delta H_{soln} > 0$; $\Delta V_{sol} > 0$

31. The following is a graph plotted between the vapour pressures of two volatile liquids against their respective mole fractions.

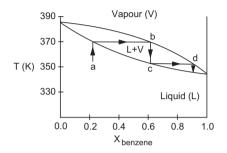


Which of the following combinations are correct?

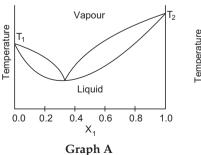
- (a) When $x_A = 1$ and $x_B = 0$, then $p = p_A^0$.
- (b) When $x_B = 1$ and $x_A = 0$, then $p > p_A^0$.
- (c) When $x_A = 1$ and $x_B = 0$, then $p < p_B^0$.
- (d) When $x_B = 1$ and $x_A = 0$, then $p = p_B^0$.
- **32.** Which of the following statements are correct for a binary solution which shows negative deviation from Raoult's law?
 - (a) The negative deviation from linearity diminishes and tends to zero as the concentration of the solution component approaches unity.
 - (b) When solutions form, their volumes are smaller than the sum of the volumes of their components.
 - (c) Heat is released during the formation of the solution.
 - (d) Heat is evolved during the formation of the solution.
- 33. A binary liquid (AB) shows positive deviation from Raoult's law when
 - (a) $p_A > p_A^0 X_A^{liq}$ and $p_A > p_B^0 X_B^{liq}$
 - (b) Intermolecular forces: A–A, B–B > A–B
 - (c) $\Delta V_{\text{mix}} > 0$
 - (d) $\Delta H_{mix} > 0$
- **34.** A graph is plotted between the vapour pressure and mole fraction of a solution containing benzene and toluene. Choose the correct options.

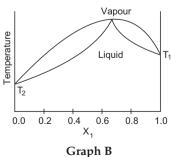


- (a) At the point *a*, the mole fraction of toluene is 0.80.
- (b) $b \rightarrow c$ represents condensation.
- (c) $c \rightarrow d$ represents vapourization.
- (d) $c \rightarrow d$ represents vapourization as well as condensation.
- **35.** A graph is plotted with the temperature of a solution containing benzene and toluene as a function of mole fraction. Choose the correct options.



- (a) $a \rightarrow b$ represents evaporation.
- (b) $b \rightarrow c$ represents condensation.
- (c) $c \rightarrow d$ represents evaporation.
- (d) $c \rightarrow d$ represents condensation.
- **36.** The azeotropic solutions of two miscible liquids
 - (a) can be separated by simple distillation
 - (b) may show positive or negative deviation from Raoult's law
 - (c) are supersaturated solutions
 - (d) behave like a single component and boil at a constant temperature
- **37.** The following two graphs are plotted between the temperature and mole fraction of the components of two different azeotropic mixtures. $(X_1 = \text{mole fraction of solvent.})$



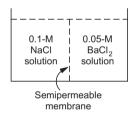


Choose the correct options.

- (a) Graph A represents minimum boiling point and maximum vapour pressure.
- (b) Graph B represents maximum boiling point and minimum vapour pressure.
- (c) Graph A represents maximum boiling point and minimum vapour pressure.
- (d) Graph B represents minimum boiling point and maximum vapour pressure.

38. A substance effloresces

- (a) due to the formation of a crust on its crystal surface
- (b) when its vapour pressure is greater than that of the water vapour in air
- (c) till it melts
- (d) when all of the above happen
- 39. Study the following figure, and choose the correct options.



- (a) There will be no movement of any solution across the membrane.
- (b) BaCl₂ will flow towards the NaCl solution.
- (c) NaCl will flow towards the BaCl₂ solution.
- (d) The osmotic pressure of 0.1 M NaCl is higher than the osmotic pressure of 0.05 M BaCl₂, assuming complete dissociation of the electrolyte.
- **40.** In which of the following pairs of solutions will the values of the van't Hoff factor be the same?
 - (a) 0.05 M K₄[Fe(CN)₆] and 0.10 M FeSO₄
 - (b) 0.10 M K₄[Fe(CN)₆] and 0.05 M FeSO₄(NH₄)₂SO₄.6H₂O
 - (c) 0.20 M NaCl and 0.10 M BaCl₂
 - (d) 0.05 M FeSO₄(NH₄)₂SO₄.6H₂O and 0.02 M KCl.MgCl₂.6H₂O

Answers

1. a	2. d	3. b	4. a	5. b
6. b	7. c	8. a	9. d	10. c
11. c	12. d	13. b	14. a	15. a
16. c	17. b	18. d	19. c	20. d
21. b	22. d	23. d	24. d	25. c
26. a	27. b	28. a, b, c, d	29. b, d	30. b, d
31. a, d	32. a, b, c	33. a, b, c	34. a, b, c	35. a, b, c
36. b, d	37. a, b	38. a, b	39. b, d	40. b, d

Hints to More Difficult Problems

1. As we know,

$$\frac{p^0 - p}{p^0} = X_1 = \text{mole fraction of solute.}$$

The ratio $(p^0 - p)/p^0$ is the relative lowering of vapour pressure, which is equal to 0.0125 here.

$$X_1 = 0.0125.$$

The relation between the mole fraction and molality is

$$\left(\frac{1}{X_1} - 1\right) = \frac{1000}{m \times 18}$$
 (mol. wt. of H₂O = 18)
or $\left(\frac{1}{0.0125} - 1\right) = \frac{1000}{m \times 18} \implies m = 0.70$.

- 3. The total molarity of all the ions is maximum in Na_3PO_4 (0.1 × 3 = 0.3 M). So it has the highest boiling point.
- 6. In an aqueous solution, BaCl₂ produces the maximum number of ions.
- 8. The number of molecules decreases due to the formation of a complex.

$$HgI_2 + 2Kl \longrightarrow K_2[HgI_4]$$

9. Ca(NO₃)₂ and Na₂SO₄ produce the same number of ions with the same molarity.

13.
$$4A \rightleftharpoons (A)_4$$
 $i = \frac{1 - \beta + \beta/4}{1} = 1 - \frac{3}{4}\beta$. $\beta = \text{degree of association} = 1 = (100\%)$. $\therefore i = 1 - \frac{3}{4} = 0.25$.

15.
$$K_b = \frac{RT_0^2 M}{1000\Delta H_{\text{vapour}}} = \frac{8.314 \times (373.15)^2 \times 18}{1000 \times 40.685} \times 10^{-3}$$

= 0.512 K kg mol⁻¹.

17. $\Delta T_b = K_b \times \text{molality}$ and $\Delta T_f = K_f \times \text{molality}$. For the same solution, the molality is the same,

or
$$\frac{\Delta T_b}{\Delta T_f} = \frac{K_b}{K_f}$$

or $\frac{\Delta T_b}{0.186} = \frac{0.512}{1.86} \Rightarrow \Delta T_b = 0.0512 \text{ K}.$

20. Molecule-solvent interaction

21.
$$K_b = \frac{RT_0^2 M_1}{1000\Delta H_{\text{vap}}} \Rightarrow \Delta H_{\text{vap}} = \frac{RT_0^2 M_1}{1000K_b}$$
.
We know that
$$T_0 = 110.7 + 273.15 = 383.15 \text{ K.}$$

$$M_1 = \text{mol mass of toluene}$$

$$= 92 \text{ g mol}^{-1}.$$

$$K_b = 3.32 \text{ K kg mol}^{-1}.$$

$$\therefore \Delta H_{\text{vap}} = \frac{8.314 \times (383.15)^2 \times 92}{1000 \times 3.32 \times 10^3} = 34 \text{ kJ mol}^{-1}.$$

- **30.** Both pairs form non-ideal solutions.
- 33. Intermolecular force of attraction
- **36.** They form a nonideal solution.
- **40.** Both pairs in (b) and (d) produce the same total number of ions after dissociation.

17

Solid-State Chemistry

• Type 1 •

1. Argon crystallizes in a structure in which the atoms are located at the positions (0, 0, 0), $(0, \frac{1}{2}, \frac{1}{2})$, $(\frac{1}{2}, 0, \frac{1}{2})$ and $(\frac{1}{2}, \frac{1}{2}, 0)$. The unit cell is

2. Potassium metal crystallizes in the form of a body-centred cubic structure. The number of nearest-neighbour atoms for each potassium

(b) six

(d) eight

(b) body-centred cubic

(d) hexagonal close packed

Choose the correct option. Only one option is correct.

(a) simple cubic

atom in the solid is

(a) four

(c) twelve

(a) 6

(c) 18

(c) face-centred cubic

3.	the fo		1	a compound that crystallizes in I lattice with a molecule at each
	(a)	one	(b)	two
	(c)	four	(d)	six
4.	NiO a ion is	adopts a	rock-salt structure. The co	ordination number of the Ni 24
	(a)	two	(b)	four
	(c)	twelve	(d)	six
5.	Consi	der the	structure of rock-salt. Ho	w many Na + ions occupy the

1-231

(b) 12

(d) 24

second–nearest neighbour location of an Na + ion?

(a) Cubic

(c) Orthorhombic

6. Which type of crystals contain the maximum number of Bravais lattices?

7. A compound contains two types of atoms—X and Y. It crystallizes in a

(b) Triclinic

(d) Tetragonal

	cubic lattice with atoms X at the corners of the unit cell and atoms Y at the body centres. The simplest possible formula of this compound is				
		X ₈ Y		X ₂ Y	
		XY		XY_8	
8.	Which	n of the following solids is amo	rpho	ous?	
		Fe metal	_	Fused quartz	
	(c)	Wurtzite	(d)	NiAs	
9.		n of the following expressions parameter <i>a</i> ?	is c	correct for a CsCl unit cell with	
	(a)	$r_{\text{Cs}^+} + r_{\text{Cl}^-} = 2a$	(b)	$r_{\rm Cs}^+ + r_{\rm Cl}^- = \frac{a}{\sqrt{2}}$	
	(c)	$r_{\rm Cs^+} + r_{\rm Cl^-} = \frac{\sqrt{3}}{2} a$	(d)	$r_{\text{Cs}^+} + r_{\text{Cl}^-} = \frac{3a}{2}$	
10.	by th	ne the construction of an MX_2 se removal of half the Cs^+ ination around each Cl^- . What An antifluorite (Na_2O) structu	ion:	ture from the bcc CsCl structure s so that there is tetrahedral as structure of MX ₂ ?	
			пе		
		A fluorite (CaF ₂) structure			
		A rutile (TiO ₂) structure	_		
		A pyrolusite (MnO ₂) structure			
11.		nolecules NaCl, MgO and Ni its 4 : 4 coordination because	O sh	now 6:6 coordination, but BeO	
	(a)	BeO is covalent	(b)	BeO is ionic	
	(c)	BeO is amphoteric	(d)	BeO is polymeric	
12.	c = 9.4			al are $a = 5.62$ Å, $b = 7.41$ Å and mutually perpendicular to each	
	(a)	tetragonal	(b)	orthorhombic	
	(c)	monoclinic	(d)	trigonal	
13.	In dia	mond, the coordination number	er of	carbon is	
	(a)	four and its unit cell has eight	carl	oon atoms	
	(b)	four and its unit cell has six ca	arboı	n atoms	
	(c)	six and its unit cell has four ca	arbo	n atoms	
	(d)	four and its unit cell has four	carb	on atoms	

- 14. In NaCl, the centre-to-centre nearest-neighbour distance of ions is
 - (a) $\frac{1}{4}a$

(b) $\frac{\sqrt{3}}{2}a$

(c) $\frac{1}{2} a \sqrt{2}$

(d) $\frac{1}{2}a$

where a is the lattice parameter.

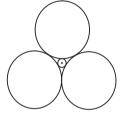
- 15. Calculate the ionic radius of a Cs^+ ion, assuming that the cell-edge length for CsCl is 0.4123 nm and that the ionic radius of a Cl^- ion is 0.181 nm.
 - (a) 0.176 nm

(b) 0.231 nm

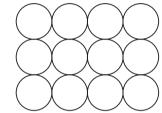
(c) 0.352 nm

- (d) 0.116 nm
- **16.** Which of the following figures represents the cross-section of an octahedral site?

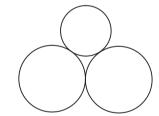




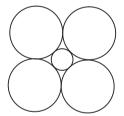
(b)



(c)



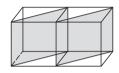
(d)



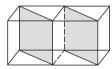
- 17. Zinc sulphide exists in two different forms—zinc blende and wurtzite. Both occur as 4:4 coordination compounds. Choose the correct option from among the following.
 - (a) Zinc blende has a bcc structure and wurtzite an fcc structure.
 - (b) Zinc blende has an fcc structure and wurtzite an hcp structure.
 - (c) Zinc blende as well as wurtzite have a hcp structure.
 - (d) Zinc blende as well as wurtzite have a cpp structure.
- **18.** Consider the radii 0.095 nm (Na $^+$), 0.181 nm (Cl $^-$), 0.074 nm (Zn $^{2+}$), 0.184 nm (S $^{2-}$), 0.068 nm (Ti $^{4+}$), 0.140 nm(O $^{2-}$), 0.169 nm (Cs $^+$). Choose the correct option from among the following. (Use radius ratio rules.)
 - (a) Na⁺ ions are packed in octahedral holes between the planes of close-packed Cl⁻ ions.

- (b) Zn^{2+} ions are packed in tetrahedral holes.
- (c) Cs⁺ ions are packed in a simple cubic array of Cl⁻ ions.
- (d) All of these
- 19. Every atom or ion that forms an fcc unit cell is surrounded by
 - (a) six octahedral holes and eight tetrahedral holes
 - (b) eight octahedral holes and six tetrahedral holes
 - (c) six octahedral holes and six tetrahedral holes
 - (d) eight octahedral holes and four tetrahedral holes
- 20. Which of the following figures represents a 110 plane?

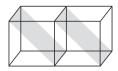
(a)



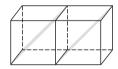
(b)



(c)



(d)



21. Which of the following expressions is correct for an NaCl unit cell with lattice parameter *a*?

(a)
$$r_{\text{Na}^+} + r_{\text{Cl}^-} = \frac{a}{\sqrt{2}}$$

(b)
$$r_{\text{Na}^+} + r_{\text{Cl}^-} = 4a$$

(c)
$$r_{\text{Na}^+} + r_{\text{Cl}^-} = \frac{a}{4}$$

(d)
$$r_{\text{Na}^+} + r_{\text{Cl}^-} = \frac{\sqrt{3}}{4} a$$

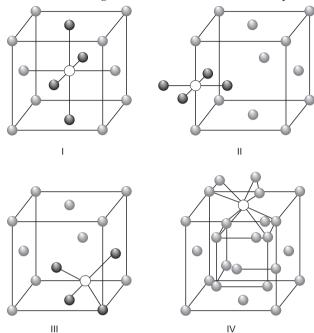
- **22.** Consider the structure of CsCl (8 : 8 coordination). How many Cs⁺ ions occupy the second-nearest neighbour locations of a Cs⁺ ion?
 - (a) 8

(b) 24

(c) 6

- (d) 16
- 23. The addition of a CaCl₂ crystal to a KCl crystal
 - (a) lowers the density of the KCl crystal
 - (b) raises the density of the KCl crystal
 - (c) does not affect the density of the KCl crystal
 - (d) increases the Frenkel defects of the KCl crystal

24. Consider the following fcc unit cells choose the correct option.



- (a) I and II represent tetrahedral holes.
- (b) II, III and IV represent tetrahedral holes.
- (c) I and II represent octahedral holes.
- (d) I, II and IV represent octahedral holes.
- 25. Iron crystallizes in a bcc system with a lattice parameter of 2.861 Å. Calculate the density of iron in the bcc system (atomic weight of Fe = 56, $N_A = 6.02 \times 10^{23} \text{ mol}^{-1}$).
 - (a) 7.92 g mL^{-1}

(b) 8.96 g mL^{-1}

(c) 2.78 g mL^{-1}

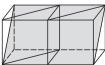
- (d) 6.72 g mL^{-1}
- 26. Bragg reflection can occur only for
 - (a) $\lambda \leq 2d$

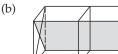
(b) $\lambda \ge 2d$

(c) $\lambda > 2d$

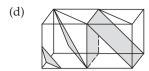
- (d) none of these
- 27. The shaded area in which of the following figures represents a 101 plane?











- **28.** Nickel crystallizes in an fcc unit cell with a cell-edge length of 0.3524 nm. Calculate the radius of the nickel atom.
 - (a) 0.1624 nm
- (b) 0.1246 nm
 - (c) 0.2164 nm
- (d) 0.1426 nm
- 29. In which of the following pairs of structures are tetrahedral as well as octahedral holes found?
 - (a) bcc and fcc

(b) hcp and simple cubic

(c) hcp and ccp

- (d) bcc and hcp
- 30. The effective nuclear charges of Na+ and F- ions are 6.50 and 4.50 respectively. If $r_{Na}^+ + r_F^- = 231$ pm, calculate the radii of Na ⁺ and F ⁻ ions.
 - (a) $r_{\text{Na}^+} = 94.5 \text{ pm}$, $r_{\text{F}^-} = 46.5 \text{ pm}$. (b) $r_{\text{Na}^+} = 94.5 \text{ pm}$, $r_{\text{F}^-} = 136.5 \text{ pm}$.
 - (c) $r_{\text{Na}^+} = 136.5 \text{ pm}, r_{\text{F}^-} = 94.5 \text{ pm}.$ (d) $r_{\text{Na}^+} = 36.0 \text{ pm}, r_{\text{F}^-} = 92.0 \text{ pm}.$
- 31. KCl crystallizes in the same type of lattice as does NaCl. Given that $r_{\text{Na}} / r_{\text{Cl}} = 0.55$ and $r_{\text{Na}} / r_{\text{K}} = 0.74$, calculate the ratio of the side of the unit cell for KCl to that for NaCl.
 - (a) 1.122
- (b) 1.224
- (c) 1.414
- (d) 0.732
- **32.** The relation between the *d*-spacing formula and the Bragg equation for a cubic crystal for first-order reflection is

 - (a) $\sin \theta = \frac{\lambda}{2a} (h^2 + k^2 + l^2)^2$ (b) $\sin \theta = \frac{\lambda}{2a} (h^2 + k^2 + l^2)^{1/2}$
 - (c) $\sin \theta = \frac{2a}{\lambda} (h^2 + k^2 + l^2)^{1/2}$ (d) $\sin \theta = \frac{2a}{\lambda} (h^2 + k^2 + l^2)^{1/2}$
- 33. The orientation of a lattice plane is determined according to its
 - (a) Scattering factor
- (b) Bravais lattice
- (c) nearest neighbour
- (d) Miller indices
- 34. Perovskite is a mineral with the formula CaTiO₃. Which of the positive ions in the crystal is more likely to be packed in the octahedral holes?
 - (a) Ti 4+
- (b) Ti²⁺
- (c) Ca 2+
- (d) O_2^+
- **35.** Which of the following statements is incorrect?
 - (a) A substitutional solid solution is one in which atoms of the solute metal occupy some locations that solvent metal atoms are expected to occupy.
 - (b) An interstitial solid solution is one in which the solute atoms occupy the interstices (holes) between the solvent atoms.

metals and metalloids.

36. The unit cell of $CO_2(s)$ is (a) fcc (b)

37. Because of anisotropy

38.

39.

40.

41.

42.

rod-like pieces

(d) Cu₃Zn is an intermetallic compound.

(b) bcc

(c) An intermetallic compound is a compound formed between

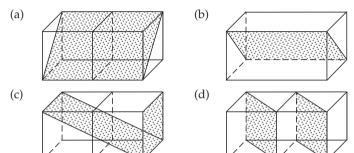
(a) mica cleaves into thin sheets and asbestos cleaves into long,

(c) linear

(d) hcp

(b)	mica cleaves into long, rod-like pieces and asbestos cleaves into thin sheets						
(c)	mica as well as asbestos cleave into thin sheets						
(d)	mica as well as asbestos cleave into long, rod-like pieces						
In Na	Cl the centres of two n	earest like-cl	narged ions a	re at a distan	ce of		
(a)	$\frac{1}{2}a\sqrt{2} \qquad \qquad \text{(b)} \frac{1}{2}a$	(c)	$\frac{\sqrt{3}}{2}a$	(d) $\frac{1}{\sqrt{2}} 2a$			
from 6	each other.						
Which struct	n of the following co ure?	mpounds re	present an i	nverse 2:3	spinel		
(a)	$Fe^{III}(Fe^{II}Fe^{III})O_4$	(b)	PbO ₂				
(c)	Al_2O_3	(d)	Mn_3O_4				
	ompounds have a stru ounds are said to have						
(a)	the smaller cations oc larger anions that of t			uoride ions a	nd the		
(b)	the larger cations occ smaller anions that of			ioride ions a	nd the		
(c)	Each fluoride ion is surrounded by four calcium ions in a tetrahedral arrangement.						
(d)	Fluoride ions occupy all the eight octahedral holes.						
Which	n of the following pairs	have fcc lat	tice structures	s?			
(a)	Rock salt and Wurtzit	te (b)	Rock salt an	d sphalerite			
(c)	CsCl and rutile	(d)	Polonium ar	nd fluorite			
Which of the following statements is incorrect for sphalerite, a form of ZnS?							
(a)	It has the same structure as diamond, except that the alternate atoms are zinc and sulphur.						
(b)	Because the sulphide than four or eight sulphide						

- (c) Because the sulphide ion is larger than the zinc ion, only four rather than six or eight sulphide ions can be packed around a zinc ion.
- (d) ZnS is polar covalent.
- **43.** Which of the following figures represents the $(\overline{1}\ 0\ 1)$ plane?



- 44. Which of the following statements is correct?
 - (a) The crystal structure of rock salt is an fcc array of anions in which the cations occupy all the octahedral holes (or vice versa)
 - (b) The sphalerite crystal structure is an expanded fcc anion lattice with cations occupying one type of tetrahedral hole.
 - (c) In the flourite crystal structure, cations occupy half the cubic holes of a primitive cubic array of anions.
 - (d) All of these.
- **45.** If the three interaxial angles defining the unit cell are all equal in magnitude, the crystal cannot be
 - (a) rhombohedral

(b) cubic

(c) hexagonal

- (d) tetragonal
- 46. Which of the following statements is correct for the NaCl lattice?
 - (a) An Na⁺ ion is placed at a distance $\frac{1}{6}a$ directly above each Cl⁻ ion (where a is the cubic unit-cell edge length).
 - (b) An Na $^+$ ion is placed at a distance $\frac{1}{4}a$ directly above each Cl $^-$ ion.
 - (c) An Na⁺ ion is placed at a distance of $\frac{1}{2\sqrt{2}}a$ directly above each Cl⁻ ion.
 - (d) An Na⁺ ion is placed at a distance $\frac{1}{\sqrt{2}}a$ directly above each Cl⁻ ion.
- **47.** Which of the following statements is correct?
 - (a) A rutile (TiO₂) structure consists of an hcp anion lattice with cations occupying half the octahedral holes.

- (b) The Wurtzite structure is derived from an expanded hcp anion array with cations occupying one type of octahedral holes.
- (c) In the fluorite structure (CaF₂), anions occupy both types of tetrahedral holes in an expanded fcc lattice of cations.
- (d) All the these
- **48.** When the radius ratios lie between 0.732 and 0.414, the arrangement of the crystal is
 - (a) tetrahedral

(b) octahedral

(c) linear

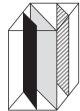
- (d) cubic
- **49.** An fcc cubic cell contains eight X atoms at the corners of the cell and six Y atoms at the faces. What is the empirical formula of the solid?
 - (a) X_3Y_4

(b) X₃Y

(c) XY₃

- (d) X_4Y_3
- 50. The spinel structure (AB_2O_4) consists of an fcc array of O^{2-} ions in which the
 - (a) A cation occupies one-eighth of the tetrahedral holes and the B cations the octahedral holes
 - (b) A cation occupies one-fourth of the tetrahedral holes and the B cations the octahedral holes
 - (c) A cation occupies one-eighth of the octahedral holes and the B cation the tetrahedral holes
 - (d) A cation occupies one-fourth of the octahedral holes and the B cations the tetrahedral holes
- **51.** Which of the following statements is incorrect for the diamond structure?
 - (a) Each atom has 4 nearest neighbours and 12 next-nearest neighbours.
 - (b) It is relatively empty.
 - (c) The maximum proportion of the available volume which may be filled by hard spheres is only 0.34.
 - (d) The maximum proportion of the available volume which may be filled by hard spheres is only 0.46.
- 52. In an NaCl structure,
 - (a) 4 corners are shared
 - (b) 6 corners and 2 edges are shared
 - (c) 3 edges are shared
 - (d) 12 edges are shared

53. If the shaded plane





intercepts the a axis at a/2, the b axis at b/2, and lies parallel to the c axis, then its Miller indices are

(a) $\left(\frac{1}{2} \ \frac{1}{2} \ 0\right)$

(b) $\left(0\,\frac{1}{2}\,\,\frac{1}{2}\right)$

(c) (2 2 0)

- (d) (2 1 0)
- **54.** For a given crystal the lattice parameter *a* is 318 pm. The *d*-spacing for a (III) plane is
 - (a) 225 pm

(b) 184 pm

(c) 318 pm

- (d) 390 pm
- 55. All the positions in an fcc lattice are occupied by A atoms, and the bcc octahedral hole in it by a B atom of appropriate size. The formula of the compound is
 - (a) A_2B
- (b) AB₄
- (c) A₄B
- (d) A_4B_3
- 56. The zinc blende structure results when the Zn atoms occupy one
 - (a) fcc lattice and the S atoms another
 - (b) bcc lattice and the S atoms another
 - (c) bcc lattice and the S atoms an fcc lattice
 - (d) sc lattice and the S atoms another
- **57.** A metal of density 7.5×10^3 kg m⁻³ has an fcc crystal structure with lattice parameter a = 400 pm. Calculate the number of unit cells present in 0.015 kg of the metal.
 - (a) 6.250×10^{22}

(b) 3.125×10^{23}

(c) 3.125×10^{22}

- (d) 1.563×10^{22}
- **58.** The molar volume of KCl and NaCl are 37.46 mL and 27.94 mL respectively. The ratio of the unit cube edges of the two crystals is
 - (a) 1.296

(b) 1.116

(c) 1.341

- (d) 0.950
- **59.** When the radius ratio $r_{\rm M}/r_{\rm X}$ of a crystal lies in the range 0.414–0.732, the coordination number of M and its corresponding crystal is
 - (a) 4, TiO₂
- (b) 6, ZnS
- (c) 8, CsCl
- (d) 6, TiO₂
- **60.** The ratio of the volume of a tetragonal lattice to that of a hexagonal lattice is

	(a)	$\frac{\sqrt{3}}{2}abc$	(b) $\frac{2}{\sqrt{3}}$	(c)	$\frac{2}{\sqrt{3}} \frac{a^2}{b} c$	(d) -	$\frac{\sqrt{3}}{2} \frac{a^2}{bc}$	
61.	If a n becau	netal has a se	bcc crysta	l structure,	the coord	ination 1	number	is 8,
	(a)	each atom layer below		our atoms in ne in its own		above it	, four i	n the
	(b)	each atom layer below		our atoms in e in its own l	,	above it	, four i	n the
	(c)	two atoms layer below		atoms in the			n, four	in the
	(d)	each atom t layer below		ght atoms ir ne in its own		above it	, eight i	in the
62.	volum (a)	c lattice has ne of the lattion 10.8 mL 8.6 mL		ng all the em (b)			te the	molar
63.	(a) (b) (c)	ep structure, first and thi first and fou second and first, third a	rd layers a urth layers fourth lay	are repeated ers are repea	nted			
64.	equal (a)	5.66 <i>a</i> ³ 2.66 <i>a</i> ³		pied per sphe (b)				res of
65.	hexag (a)	n of the follo onal close-pa Each sphere Each sphere	acked laye e is surrou	r? nded by six.		or a two	-dimens	sional

(d) Each void is surrounded by three spheres. 66. Which of the following mechanisms is incorrect for a stoichiometrically pure compound AB?

(c) Each sphere has three voids.

(a) There is a displacement of A atoms to interstitial sites, leaving an equal number of A-type vacancies in the structure.

(b) There is a displacement of B atoms to interstitial sites, accompanied by the formation of an equal number of B-type vacancies.

- (c) An equal number of A- and B-type vacancies are created by the migration of atoms to the crystal surface.
- (d) An equal number of interstitial A and B atoms, and A- and B-type vacancies, are created.
- 67. Suppose there are N atoms in a crystal with N_i interstitial positions in its structure. If there are n Frenkel defects in the crystal then

 - (a) $n = (NN_i) \exp(-E_i/2k_BT)$ (b) $N = (nN_i)^{1/2} \exp(-E_i/2k_BT)$
 - (c) $n = (NN_i)^{1/2} \exp(-E_i/2k_BT)$ (d) $n = (NN_i)^{1/2} \exp(+E_i/2k_BT)$

where E_i is the energy required to remove an atom from a lattice site to an interestitial position.

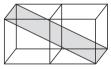
- **68.** Crystals may be coloured by
 - (a) the introduction of chemical impurities
 - (b) X-ray, γ-ray and electron bombardment
 - (c) introducing an excess of the metal
 - (d) all these methods
- **69.** The radius ratio (γ_M/γ_X) for cation-anion and anion-anion contact for a tetrahedral arrangement of anions around a cation is
 - (a) 0.414

(b) 0.225

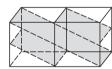
(c) 1.00

- (d) 0.732
- 70. Which of the following shaded areas of the figures represent (002) planes?

(a)



(b)



(c)



(d)



- 71. The numbers of tetrahedral and octahedral holes in a ccp array of 100 atoms are respectively
 - (a) 200 and 100

(b) 100 and 200

(c) 200 and 200

- (d) 100 and 100
- 72. Copper crystallizes in an fcc form and sodium in a bcc form. The coordination numbers of Cu and Na are respectively
 - (a) 12 and 12

(b) 12 and 8

(c) 8 and 12

(d) 6 and 8

- **73.** In AgCl, the silver ion is displaced from its lattice position to an interstitial position. Such a defect is called a
 - (a) Schottky defect

(b) Frenkel defect

(c) Wadsley defect

- (d) colour centre
- 74. A solid solution of CdBr₂ in AgBr contains
 - (a) Schottky defects
 - (b) Frenkel defects
 - (c) Frenkel as well as Schottky defects
 - (d) colour centres
- 75. NaCl shows Schottky defects and AgCl Frenkel defects. Their electrical conductivity is due to the
 - (a) motion of ions and not the motion of electrons
 - (b) motion of electrons and not the motion of ions
 - (c) lower coordination number of NaCl
 - (d) higher coordination number of AgCl
- **76.** If a crystal contains a total of N atoms, and n Schottky defects are produced by removing n cations and r anions from the interior of the crystal, then

(a)
$$n = N \exp(-E_p/2k_BT)$$

(b)
$$N = n \exp(-E_p/2k_BT)$$

(c)
$$n = N \exp(-E_p/k_BT)$$

(d)
$$n = N \exp(E_p/k_BT)$$

when E_p is the energy of the formation of a pair, k_B is the Boltzmann constant and T is the temperature in kelvin.

- 77. Which of the following statements is correct for a close-packed structure?
 - (a) Each octahedral void is surrounded by six spheres and each sphere is surrounded by six octahedral voids.
 - (b) Each octahedral void is surrounded by six spheres and each sphere is surrounded by three octahedral voids.
 - (c) Each octahedral void is surrounded by six spheres and each sphere is surrounded by eight octahedral voids.
 - (d) Each octahedral void is surrounded by eight spheres and each sphere is surrounded by six octahedral voids.
- 78. The coordination number of the fcc structure for metals is 12, since
 - each atom touches four others in the same layer, three in the layer above and three in that below
 - (b) each atom touches four others in the same layer, four in the layer above and four in that below
 - (c) each atom touches six others in the same layer, six in the layer above, and six in that below

80.

81.

82.

83.

84.

85.

86.

above and six in that below

(a) isotropic and supercooled liquids(b) anisotropic and supercooled liquids(c) isoenthalpic and superheated liquids(d) isotropic and superheated solids

79. Amorphous solids may be classified as

(d) each atom touches three others in the same layer, six in the layer

In the face-centred cubic lattice structure of gold, the closest distance between gold atoms (the length of the cubic unit cell being <i>a</i>) is					
(a)	$\frac{1}{4}a\sqrt{2}$		(b)	$\frac{1}{4}(a\sqrt{2})$	
(c)	$a\sqrt{2}$		(d)	$\frac{1}{2}\frac{a}{\sqrt{2}}$	
How :		lral sites per spl	nere a	re there in a	cubic close-packed
(a)	four	(b) two	(c)	one	(d) six
which and a	the gold ator	ms occupy the l	attice	points at the	in a cubic lattice in the corners of a cube face. The formula of
(a)	Au ₃ Cu		(b)	$AuCu_3$	
(c)	Au_4Cu		(d)	$AuCu_2$	
In the	structure of d	liamond, the car	bon a	toms appear	at
(a)	0 0 0 and $\frac{1}{2}$	$\frac{1}{2} \frac{1}{2}$	(b)	$\frac{1}{4}$ $\frac{1}{4}$ $\frac{1}{4}$ and	$\frac{1}{2} \frac{1}{2} \frac{1}{2}$
(c)	0 0 0 and $\frac{1}{4}$	$\frac{1}{4}$ $\frac{1}{4}$	(d)	0 0 0 and	$\frac{3}{4} \frac{3}{4} \frac{3}{4}$
					tetragonal with 16 bes the basis consist
(a)	8	(b) 16	(c)	4	(d) 2
Which		wing statements	s is c	orrect for a	cubic close-packed
(a)	There are thr	ree sites.	(b)	There are to	wo types of sites.
(c)	There are for	ur types of sites.	(d)	There are si	ix types of sites.
					the solid state are rays are scattered by
(a)	electrons		(b)	positrons	
(c)	protons		(d)	neutrons	

- 87. All spinel structures have a
 - (a) ccp array of anions
 - (b) simple cubic structure
 - (c) bcc array of anions
 - (d) hexagonal close-packed array of anions
- **88.** Which of the following statements are correct in the context of point defects in a crystal?
 - (a) AgCl has anion Frenkel defects and CaF₂ has Schottky defects.
 - (b) AgCl has cation Frenkel defects and CaF₂ has anion Frenkel defects.
 - (c) AgCl as well as CaF₂ have anion Frenkel defects.
 - (d) AgCl as well as CaF₂ have Schottky defects.
- **89.** A solid is made of two elements, A and B. The atoms of A are arranged in a ccp structure and those of B occupy all the tetrahedral sites. The formula of the compound is
 - (a) AB_2
- (b) A₂B
- (c) A_2B_3
- (d) A_3B_2
- **90.** Barium titanate has a pyrolusite structure (a cubic lattice), with barium ions occupying the corners of the unit cell, oxide ions the face centres and titanium ions the centres. Assuming that Ti ⁴⁺ ions occupy the holes of the BaO lattice, what type of hole and what fraction of such holes do these ions occupy?
 - (a) 25% of the octahedral holes
- (b) 75% of the tetrahedral holes
- (c) 50% of the tetrahedral holes
- (d) 100% of the octahedral holes
- **91.** Three elements P, Q and R crystallize in a cubic solid lattice. The P atoms occupy the corners, Q atoms the cube centres and R atoms the edges. The formula of the compound is
 - (a) PQR
- (b) PQR₂
- (c) PQR₃
- (d) PQ₃R

• *Type 2* •

Choose the correct options. More than one option is correct.

- **92.** Which type of crystals contain one Bravais lattice?
 - (a) Hexagonal

- (b) Triclinic
- (c) Rhombohedral
- (d) Monoclinic
- 93. Which of the following crystals have 8:8 coordination?
 - (a) NH₄Cl

(b) AlFe

(c) MnO

(d) NH₄Br

94.	Which	n of the following crystals have	e 6 : 6	coordination?
	(a)	NH_4I	(b)	MgO
	(c)	MnO	(d)	ZnS
95.	Which	n of the following crystals have	e 4 : 4	coordination?
	(a)	HgS	(b)	NH ₄ F
	(c)	SiC	(d)	NaCl
96.		luorite structure shows 8:4 c have fluorite-like lattices?	oord	ination. Which of the following
	(a)	CaF ₂	(b)	SrCl ₂
	(c)	BaF ₂	(d)	ThO ₂
97.		rutile) shows 6 : 3 coordination le-like structure?	n. W	hich of the following solids have
	(a)	MnO_2	(b)	ZnS
	(c)	KCl	(d)	SnO ₂
98.	Which	n of the following structures ha	ave la	ayered lattices?
	(a)	Cadmium iodide	(b)	Ice
	(c)	Graphite	(d)	Diamond
99.	Which	n of the following crystals show	w 4 :	2 coordination?
	(a)	CaF ₂	(b)	BeF_2
	(c)	SiO_2	(d)	PbO ₂
100.	The h	cp and ccp structure for a give	n ele	ment would be expected to have
	(a)	the same coordination number	er	
	(b)	the same density		
		the same packing fraction		
	(d)	all the above		
101.	In an	NaCl structure, all the		
	(a)	octahedral sites are occupied		
		tetrahedral sites are unoccup		
		octahedral as well as the tetra		-
	(d)	octahedral as well as the tetra	hedi	ral sites are unoccupied
102.		e e		rrect for the rock-salt structure?
		The catalog dual sites are small		
	(b)	sites are empty.	иртес	l by cations and the tetrahedral
	(c)	The radius ratio is 0.732.		

(d) The radius ratio is 0.999.

- **103.** Which of the following compounds represent a normal 2:3 spinel structure?
 - (a) $Mg^{II}Al_2^{III}O_4$

 $\text{(b)} \quad \text{Co}^{\text{II}}\text{(Co}^{\text{III}}\text{)}_2\text{O}_4$

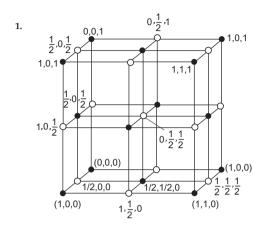
(c) $Zn(TiZn)O_4$

- (d) Ni(CO)₄
- **104.** Which of the following statements are correct?
 - (a) The coordination number of each type of ion in a CsCl crystal is eight.
 - (b) A metal that crystallizes in a bcc structure has a coordination number of twelve.
 - (c) A unit cell of an ionic crystal shares some of its ions with other unit cells.
 - (d) The length of the unit cell in NaCl is 552 pm (given that r_{Na^+} = 95 pm and r_{Cl^-} = 181 pm.

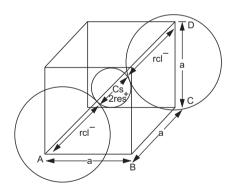
Answers

1. c	2. d	3. b	4. d	5. b
6. c	7. c	8. b	9. c	10. b
11. a	12. b	13. a	14. d	15. a
16. d	17. b	18. d	19. a	20. a
21. a	22. c	23. a	24. c	25. a
26. a	27. a	28. b	29. c	30. b
31. a	32. b	33. d	34. a	35. c
36. a	37. a	38. a	39. a	40. a
41. b	42. b	43. b	44. d	45. c
46. b	47. d	48. b	49. c	50. a
51. d	52. d	53. c	54. b	55. c
56. a	57. c	58. b	59. b	60. b
61. a	62. d	63. b	64. a	65. c
66. d	67. c	68. d	69. b	70. c
71. a	72. b	73. b	74. c	75. a
76. a	77. a	78. b	79. a	80. b
81. c	82. b	83. c	84. a	85. b
86. a	87. a	88. b	89. b	90. a
91. c	92. a, b, c	93. a, b, d	94. a, b	95. a, c
96. a, b	97. a, d	98. a, c	99. b, c	100. a, c
101. a, b	102. a, c	103. a, b	104. a, c	

Hints to More Difficult Problems



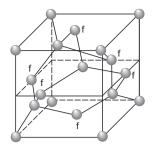
- **4.** The coordination number of Na in NaCl and that of Cl in the same compound is six, which is same as that of the Ni ion in NiO.
- 9. CsCl has a bcc structure.



$$AD^{2} = AC^{2} + CD^{2}$$

$$= (a\sqrt{2})^{2} + a^{2} = 3a^{2} \Rightarrow AD = a\sqrt{3}.$$
Also,
$$AD = r_{Cl}^{-} + 2r_{Cs}^{+} + r_{Cl}^{-} = a\sqrt{3} \Rightarrow r_{Cs} + r_{Cl}^{-} = \frac{\sqrt{3}}{2}a$$

13. The space lattice of diamond is fcc. The primitive basis has two identical atoms (0, 0, 0), $(\frac{1}{4}, \frac{1}{4}, \frac{1}{4})$ associated with each point of the fcc lattice. (see the figure.). Thus the conventional unit cube contains eight atoms. Also diamond is tetrahedral.



15. CsCl has a bcc lattice. So $d_{\text{body}} = a\sqrt{3}$

or
$$d_{\text{body}} = \sqrt{3} \times 0.4123 \text{ nm} = 0.7141 \text{ nm}.$$

The sum of the ionic radii of Cs⁺ and Cl⁻ ions is half this distance, i.e.,

$$r_{\text{Cs}}^+ + r_{\text{Cs}}^- = \frac{d_{\text{body}}}{2} = \frac{0.7141}{2} \text{ nm} = 0.3571 \text{ nm}.$$

- **16.** The cross-section of an octahedral hole is represented by (a). The octahedral holes are surrounded by six atoms arranged at the corners of an octahedron.
- 19. Focus on the atoms in the centre of the top face of the unit cell.

25.
$$\rho = \frac{ZM}{N_{\Delta}a^3}$$
 For bcc, $Z = 2$.

$$\rho_{Fe} = \frac{(2) \times 56.0 \text{ g mol}^{-1}}{(6.02 \times 10^{23} \text{ mol}^{-1})(2.861 \times 10^{-8}) \text{ cm}^3} = 7.92 \text{ g cm}^{-3}.$$

28. For an fcc lattice,

$$d_{\text{face}} = a\sqrt{2}$$
 or
$$4r_{\text{Ni}} = a\sqrt{2}.$$

$$r_{\text{Ni}} = \frac{a\sqrt{2}}{4} = \frac{\sqrt{2}}{4} \times 0.3524$$

= 0.1246 nm.

31.
$$\frac{r_{\text{Na}}^+}{r_{\text{Na}}^-} = 0.55$$
 or $1 + \frac{r_{\text{Na}}^+}{r_{\text{Cl}}^-} = 1.55$ or $\frac{r_{\text{Na}}^+ + r_{\text{Cl}}^-}{r_{\text{Cl}}^-} = 1.55$.

Given that
$$\frac{r_{\text{Na}}^+}{r_{\text{K}}^+} = 0.74$$

or
$$\frac{r_{\text{K}^+}}{r_{\text{Na}^+}} = \frac{r_{\text{K}^+}}{r_{\text{Na}^+}/0.55} = \frac{0.55}{r_{\text{Na}^+}/r_{\text{K}^+}} = \frac{0.55}{0.74} = 0.74$$

or
$$1 + \frac{r_{K}^{+}}{r_{Na}^{+}} = 1 + 0.74.$$

$$\frac{r_{\text{K}^+} + r_{\text{Cl}^-}}{r_{\text{Na}^+} + r_{\text{Cl}^-}} = \frac{1.74}{1.55}$$
or
$$\frac{a_{\text{KCl}}/2}{a_{\text{NaCl}}/2} = \frac{1.74}{1.55} = 1.122$$

32. For a cubic crystal,

$$d_{hKl} = \frac{a}{\sqrt{h^2 + k^2 + l^2}} \tag{1}$$

where a = lattice parameter

hkl = Miller indices.

Use Bragg's equation

$$n\lambda = 2d_{hkl}\sin\theta$$
.

For first-order reflection, n = 1.

$$\therefore \quad \lambda = 2d_{hkl}\sin\theta. \tag{2}$$

From Equations (1) and (2),

$$\sin \theta = \frac{\lambda}{2a} (k^2 + k^2 + l^2)^{\frac{1}{2}}$$

- **36.** Solid CO₂ adopts a rock-salt structure.
- **39.** The inverse spinel is represented by the formula $B(AB)O_4$ containing A^{2+} and B^{3+} ions.
- 45. In a hexagonal lattice,

$$\alpha = \beta = 90^{\circ}; \gamma = 120^{\circ}.$$

- 48. Apply the radius ratio rules.
- 50. Consider the positions where a cation can be surrounded by four atoms.

53. Miller indices
$$hkl = \frac{a}{2}, \frac{b}{2}, \frac{c}{\infty} = (2, 2, 0)$$

54.
$$d_{hkl} = \frac{a}{\sqrt{h^2 + k^2 + l^2}} \implies d_{110} = \frac{a}{\sqrt{1^2 + 1^2 + 1^2}} = \frac{a}{\sqrt{3}}$$

 $d_{11} = \frac{318}{\sqrt{3}} = 184 \text{ pm.}$

57.
$$V_1 = (400 \times 10^{-12} \text{ m})^3 = 64 \times 10^{-30} \text{ m}^3.$$

$$V_2 = \frac{\text{mass}}{\text{density}} = \frac{0.015 \text{ kg}}{7.5 \times 10^{-3} \text{ m}^{-3}}$$

$$= 2 \times 10^{-6} \text{ m}^3,$$

where V_1 is the volume of the unit cell and V_2 that of the metal sample.

$$\therefore$$
 no. of unit cells = $\frac{2 \times 10^{-6} \text{ m}^3}{64 \times 10^{-30} \text{ m}^3} = 3.125 \times 10^{22}$.

58.
$$\frac{d_{\text{KCl}}}{d_{\text{NaCl}}} = \left(\frac{37.46}{27.94}\right)^{\frac{1}{3}} = 1.116.$$

- 59. Apply the radius ratio rules.
- **60.** The volume of a lattice is given by

$$V = abc(1 - \cos^2\alpha - \cos^2\beta - \cos^2\gamma - 2\cos\alpha\cos\beta\cos\gamma)^{\frac{1}{2}}.$$

$$V_{\text{tetraconal}} = a^2c \text{ (because } a = b \neq c, \alpha = \beta = \gamma = 90^\circ).$$

$$V_{\text{hexagonal}} = a^2 c \times \frac{\sqrt{3}}{2}$$
 (because $a = b \neq c$, $\alpha = \beta = 90^\circ$, $\gamma = 120^\circ$).

$$\therefore \quad \frac{V_{\text{tetragonal}}}{V_{\text{hexagonal}}} = \frac{a^2 c \times 2}{a^2 c \sqrt{3}} = \frac{2}{\sqrt{3}}.$$

62. Volume =
$$a^3 = (400 \times 10^{-12} \text{ m})^3 = 64 \times 10^{-24} \text{ cm}^3$$

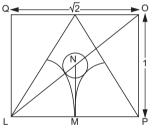
$$V_{\text{total}} = V N_{\text{A}} = 64 \times 10^{-24} \times 6.02 \times 10^{23} \approx 38.4.$$

Molar volume =
$$\frac{1}{4} V_{\text{total}} = \frac{1}{4} \times 38.4 = 9.6 \text{ mL}.$$

- **63.** Draw the structure.
- **69.** The required tetrahedron is shown in the figure. The right triangle LMN is similar to the right triangle LPO.

$$\therefore \quad \frac{LM}{LN} = \frac{LP}{LO} = \sqrt{\frac{2}{\sqrt{3}}} .$$

If *r* is the radius of the sphere representing the void and *R* the radius of the spheres in a closest-packed structure then



$$\frac{LM}{LN} = \frac{R}{R+r} = \sqrt{\frac{2}{3}} .$$

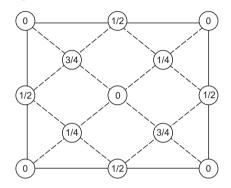
$$r = 0.225 R \Rightarrow \frac{r}{R} = 0.225.$$

- **71.** The number of octahedral voids (holes) in a close-packed structure is equal to the number of spheres (atoms). Similarly, the number of tetrahedral voids (holes) in a close-packed structure is twice the number of spheres (atoms).
- **72.** For an fcc or a ccp structure, the coordination number is 12 and for a bcc structure the coordination number is 8.

81. The number of octahedral voids belonging to one sphere is given by the following ratio.

$$\frac{\text{Number of octahedral voids around the sphere}}{\text{Number of spheres around the void}} = \frac{6}{6} = 1.$$

83. The space lattice of diamond is fcc. The primitive basis has two identical atoms at 0, 0, 0 and $\frac{1}{4}$, $\frac{1}{4}$, $\frac{1}{4}$ associated with each point of the fcc lattice as shown in the figure.



- 89. See the hint to Q. 72.
- **91.** The compound has a perovskite structure.
- **93.** Such coordination is exhibited by compounds which have bcc latice structures.
- 102. Draw the structure and apply radius ratio rules.

$$\Pi$$
 Π

103. Normal spinels are represented by formula AB₂O₄.

18

Surface Chemistry and Colloids

• Type 1 •

(a) Physical adsorption occurs at very low temperature and

(b) The magnitude of chemisorption decreases with rise in temperature and physisorption increases with rise in temperature.

(c) Chemisorption is irreversible and physisorption is reversible.(d) In physisorption, the activation energy of desorption is very low and in chemisorption, the activation energy of desorption is very high.

2. For the adsorption of a gas on a solid, the plot of $\log(x/m)$ versus $\log p$ is

(c) n

(d) $\frac{1}{1}$

Choose the correct option. Only one option is correct.1. Which of the following statements is incorrect?

linear with slope equal to

(a) k

chemisorption occur at all temperatures.

(b) $\log k$

(a) increase with decrease in temperature.(b) increase with increase in temperature.

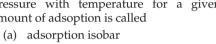
				11	
	(where, <i>p</i> = presadsorbed)	ure of gas, $m = m$	ass of the adsorbe	nt, x = mass of the	gas
3.	Which of the fol of physisorption	0 0	ecules have maxim	um value of entha	ılpy
	(a) C_2H_6	(b) Ne	(c) H_2O	(d) H_2	
4.	Which of the fol	lowing gas is ads	sorbed most by act	ivated charcoal?	
	(a) CO_2	(b) N ₂	(c) CH ₄	(d) Ar	
5.			the evolution of at of substance ads	heat. So according orbed should	g to

1-253

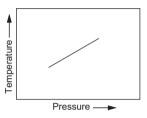
- (c) decrease with decrease in temperature.
- (d) decrease with increase in temperature.
- 6. In physical adsorption, the gas molecules are held by solid surfaces by
 - (a) strong chemical forces
- (b) van der Waals forces

(c) metallic bonds

- (d) gravitational forces
- 7. At low pressure, the fraction of the surface covered follows
 - (a) zero-order reaction
- (b) second-order reaction
- (c) first-order reaction
- (d) fractional order
- 8. The curve showing the variation of pressure with temperature for a given amount of adsoption is called



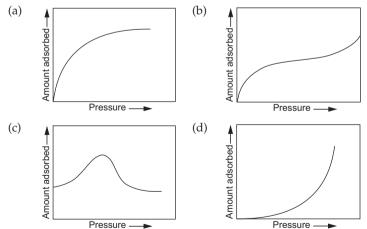
- (b) adsorption isotherm
- (c) adsorption isostere
- (d) adsorption isochore



- 9. At high pressure, the entire surface gets covered by a monomolecular layer of the gas follows
 - (a) three-halved order
- (b) second-order

(c) first-order

- (d) zero-order
- 10. Which of the following curves do not correspond to adsorption isotherms?



- 11. Which of the following statements is correct?
 - (a) Physical adsorption is a multilayer phenomena.
 - (b) Chemical adsorption is the function of adsorbate only.

- (c) The extent of physical adsorption increases with increase in pressure of the adsorbate and ultimately attain a limited value.
- (d) Physical adsorption is a monolayer phenomena.

12. Indicate the correct statement.

- (a) In chemisorption, there is no disruption of bonding in an adsorbed molecule.
- (b) The rate of decomposition of the substance adsorbed on a surface depends on the surface coverage.
- (c) In heterogeneous catalytic reaction no surface reaction occurs.
- (d) Increase in surface area of catalyst reduces the surface phase reactions.
- **13.** Which of the following statements is incorrect?
 - (a) Adsorption always leads to a decrease in enthalpy and entropy of the system.
 - (b) Adsorption arises due to unsaturation of valence forces of atoms or molecules on the surface.
 - (c) Adsorption increases with rise in the temperature.
 - (d) Adsorption decreases the surface energy.
- 14. During adsorption
 - (a) $T \triangle S$ is positive
- (b) $\triangle H T \triangle S$ is negative

(c) $\triangle H$ is positive

- (d) $T \triangle S$ and $\triangle G$ become zero
- 15. Which of the following statements is not correct?
 - (a) Decrease of temperature and increase of pressure, both tend to cause increase in the magnitude of adsorption of a gas on a solid.
 - (b) The easily liqueficable gases adsorb more on solid.
 - (c) Greater the surface area per unit mass of the adsorbent, the greater is its capacity of adsorption.
 - (d) None of these.
- 16. The range of colloidal dispersions lie between
 - (a) less than 1 mµ
- (b) greater than 0.5μ

(c) 0.5 µ to 1 mµ

- (d) none of these
- 17. If the disperse phase and the dispersion medium both are liquid, the colloidal solution is classified as
 - (a) emulsion

(b) foam

(c) gel

- (d) supercooled liquid
- **18.** Separation of colloidal particles from those of molecular dimension by means of electric current is known as
 - (a) Electroosmosis
- (b) Electrophoresis

(c) Electrodialysis

- (d) Electrolysis
- 19. The Brownian movement is due to
 - (a) enthalpy change during the formation of colloids
 - (b) attractive forces between the colloidal particles and the molecules of dispersion medium
 - (c) the impact of moelcules of the dispersion medium on the colloidal particles
 - (d) the movement of positively charged colloidal particle to negatively charged particle
- 20. Gold number is the index for
 - (a) protective power of lyophilic colloid
 - (b) purity of gold
 - (c) metallic gold
 - (d) electroplated gold
- 21. Lyophobic colloids are
 - (a) reversible colloid
- (b) irreversible colloid
- (c) protective colloids
- (d) proteins
- 22. The critical micelle concentration (CMC) is
 - (a) the concentration at which micellization starts
 - (b) the concentration at which the true solution is formed
 - (c) the concentration at which one molar electrolyte is present per 1000 g of the solution
 - (d) the concentration at which $\Delta H = 0$
- **23.** Which type of molecules form micelles?
 - (a) Nonpolar molecules
- (b) Polar molecules
- (c) Surfactant molecules
- (d) Salt of weak acid and weak base
- **24.** Fog is a colloidal solution of
 - (a) liquid particles dispersed in gas
 - (b) gaseous particles dispersed in a liquid
 - (c) solid particles dispersed in a liquid
 - (d) solid particles dispersed in gas
- **25.** A gel is
 - (a) a liquid mass of a lyophilic sol in which all the dispersion medium has penetrated into the sol particles.
 - (b) like an emulsion which is stabilized by adding emulsifying agent
 - (c) a semirigid mass of a lyophobic sol in which all the dispersion medium has penetrated into the sol particles

26.

27.

28.

29.

30.

(d)	a semirigid mass of a lyophilic sol in which all the dispersion medium has penetrated into the sol particles.					
Addit	ion of lyophil	ic soln. to the em	ulsio	n, forms		
(a)	a protective i	film around the d	lispe	rsed phase		
(b)	a protective i	film around the c	lispe	rsion medium	L	
(c)	an aerosol					
(d)	true solution					
Durin	g Micelle forn	nation				
(a)	$\triangle H = +ve$	$\Delta S = +ve$	(b)	$\Delta H = -ve$	$\Delta S = -ve$	
(c)	$\Delta H = -ve$	$\Delta S = +ve$	(d)	$\triangle H = +ve$	$\Delta S = -ve$	
Micell	es form only					
(a)	below the critical micelle concentration (CMC) and below the krafft temperature (KT)					
(b)	above the CN	MC and below th	e KT			
(c)	above the CN	MC and above the	e KT			
(d)	below the CN	MC and above th	e KT			
Micell	es are					
(a)	emulsion cur	n gel	(b)	associated co	lloids	
(c)	adsorbed cat	alysts	(d)	ideal solution	ns	
Which	n of the follow	ing ions have mi	nimı	ım value of flo	occulating power?	
(a)	PO_4^{3-}		(b)	SO_4^{2-}		
(c)	SO ₃ ²⁻		(d)	NO_3^-		

- **31.** The isoelectric-point of a colloidally dispersed material is the pH value at which
 - (a) the dispersed phase migrate in an electric field
 - (b) the dispersed phase does not migrate in an electric field
 - (c) the dispersed phase has pH equal to 7
 - (d) the dispersed phase has pH equal to zero
- **32.** Which of the following statements is incorrect?
 - (a) Emulsions are prepared by shaking two liquid components, say oil and water and adding some emulsifying agent.
 - (b) Water-in-oil emulsions are formed when the emulsifying agent at the interface is chiefly in the water phase.
 - (c) Water-in-oil emulsions are formed when the emulsifying agent at the interface is chiefly in the oil phase.
 - (d) Gems and gels mixed together to give emulsion.

- 33. Blue colour of the sky and red colour of the sunsets are due to
 - (a) scattering of light from the sun
 - (b) scattering of light from particles of dust in the atmosphere
 - (c) refraction of blue light by impurities in sea water
 - (d) scattering of light due to ozone layer.
- **34.** The arsenious soln. is negatively charged. The maximum power of precipitating it, is in
 - (a) Na₂SO₄

(b) Na₃PO₄

(c) AlCl₃

- (d) $Mg(NO_3)_2$
- **35.** Which of the following statements is correct for Tyndall effect?
 - (a) Scattering and polarizing of light by small suspended particles is called Tyndall effect.
 - (b) Tyndall effect of colloidal particles is due to dispersion of light.
 - (c) Tyndall effect is due to refraction of light.
 - (d) Zig-zag motion of suspended particles.
- **36.** The coagulating power of an effective ion carrying the charge opposite to the sol particles is given by
 - (a) Brownian movement
- (b) Gold number

(c) Tyndall effect

- (d) Hardy-Schulz law
- 37. Milk is an emulsion in which
 - (a) a liquid is dispersed in a liquid
 - (b) a solid is dispersed in a liquid
 - (c) a gas is dispersed in a liquid
 - (d) lactose is dispersed in a liquid
- 38. Lyophilic solutions are more stable than lyophobic solution because
 - (a) the colloidal particles have positive charge
 - (b) the colloidal particles have negative charge
 - (c) the colloidal particles are solvated
 - (d) there is strong electrostatic repulsions between the negatively charged colloidal particles
- 39. Hardy-Schulz law states that
 - (a) larger the size of the coagulating ions, greater its coagulating power, having opposite sign of solution
 - (b) solution must have zero gold number
 - (c) disperse phase and dispersion medium must be of the same sign
 - (d) micelles coagulate in presence of surfactants

40. Cationic, anionic and non-ionogenic surfactants are respectively

(a)
$$C_{17}H_{35}CO_2Na$$
, $C_{16}H_{33}$ —Cl and $C_nH_{2n+1}(OCH_2CH_2)_nOH$

(b) $C_nH_{2n}(OCH_2CH_2)_mOH$, $C_{15}H_{31}CO_2Na$ and $C_{18}H_{37}(NH_3)Cl^{-1}$

(c)
$$C_{16}H_{33}$$
— $C1$, $C_{17}H_{35}CO_2Na$ and $C_nH_{2n+1}(OCH_2CH_2)_mOH$

(d)
$$C_{18}H_{37}^{+}NH_3Cl^{-}$$
, $C_{16}H_{33}$ —Cl and $C_nH_{2n+1}OH$

- 41. Peptization involves
 - (a) precipitation of colloidal particles
 - (b) disintegration of colloidal aggregates
 - (c) evaporation of dispersion medium
 - (d) impact of molecules of the dispersion medium on the colloidal particles
- 42. A freshly prepared Fe(OH)₃ precipitate is peptized by adding FeCl₃ solution. The charge on the colloidal particles is due to preferential adsorption of

 - (a) Br^-ion (b) $Fe^{3+}ion$ (c) OH^-ion (d) $Ba^{2+}ion$
- 43. Which of the following is not the property of hydrophilic solutions?
 - (a) High concentration of dispersed phase can be easily obtained.
 - (b) Coagulation is reversible.
 - (c) Viscosity and surface tension are nearly the same as that of water.
 - (d) The charge of the particle depends on the pH of the medium and it may be positive, negative or zero.

• Type 2 •

Choose the correct options. More than one option is correct.

- 44. At CMC, the surfactant molecules undergoes
 - (a) dissociation

- (b) aggregation
- (c) micelle formation
- (d) all of these
- 45. Emulsions are normally prepared by shaking vigorously the two components together with some kinds of emulsifying agent to stabilize the product. The emulsifying agents may be a

(a) soap

- (b) surfactant
- (c) lyophilic solution
- (d) none of these
- **46.** Surfactant molecules or ions can cluster together as *micelles*, which
 - (a) are colloid-sized clusters of molecules
 - (b) due to their hydrophobic tails tend to congregate
 - (c) due to their hydrophilic heads provide protection
 - (d) none of these
- 47. Micelle systems are used in
 - (a) detergents

- (b) magnetic separation process
- (c) petroleum recovery
- (d) all of these
- 48. Which of the following statements are correct?
 - (a) Non-ionic surfactant molecules cluster together in clumps
 - (b) Ionic surfactants tend to disrupt by electrostatic repulsions between head groups
 - (c) Micelles look like flattened-spherical structure at CMC
 - (d) All of these
- **49.** Emulsion can be destroyed by
 - (a) the addition of an emulsifier which tends to form an emulsion of the same type
 - (b) electrophoresis with a high potential
 - (c) freezing
 - (d) all of these
- **50.** Which of the following statements is correct?
 - (a) If the mutual affinity between the dispersed phase and the dispersion medium is small, the system will be lyophobic.
 - (b) If the mutual affinity between the dispersed phase and dispersion medium is great, the system will be lyophilic.
 - (c) In a system, when water is the dispersion medium, the system may be hydrophobic or hydorphilic.
 - (d) None of these
- 51. In which of the following reactions colloids are prepared by the double decomposition method?

(a)
$$2H_3AsO_4 + 3H_2S \rightarrow As_2S_3 + 6H_2O$$

sol

(b)
$$3K_4[Fe(CN)_6] + 4FeCl_3 \rightarrow Fe_4[Fe(CN)_6]_3 + 12 KCl_{sol}$$

(c)
$$Hg(CN)_2 + H_2S \rightarrow HgS + 2HCN$$

(d) All of these

- **52.** Colloidal solution is prepared by
 - (a) reduction method
- (b) dissociation method
- (c) hydrolysis method
- (d) none of these
- 53. Which of the following statements is correct for electrophoresis?
 - (a) Colloids are uncharged particles and do not migrate towards the electrodes when electric field is applied.
 - (b) In electrophoresis, solution migrates either to the anode or to the cathode depending on the positively or negatively charged solution.
 - (c) Electrophoresis is a useful method for finding the charge of a solution.
 - (d) All of these.

Answers	;
---------	---

1. b	2. d	3. c	4. a	5. a
6. b	7. c	8. c	9. d	10. c
11. a	12. b	13. c	14. b	15. d
16. c	17. a	18. b	19. c	20. a
21. b	22. a	23. c	24. a	25. d
26. a	27. a	28. c	29. b	30. a
31. b	32. d	33. b	34. c	35. a
36. d	37. a	38. c	39. a	40. c
41. b	42. b	43. c	44. b, c	45. a, b, c
46. a, b, c	47. a, c	48. a, b, c, d	49. b, c	50. a, b, c
51. a, b, c, d	52. a, b, c	53. b, c		

Hints to More Difficult Problems

- **2.** Apply $x/m = Kp^{1/n}$
- 3. H₂O can be liquefied easily
- 6. Weak intermolecular forces
- 12. Greater area of the surface
- 14. Spontaneous reaction
- 19. Zig-zag motion of suspended particles
- 21. Rest are lyophilic

- It is the minimum concentration at which surfactant molecules undergo aggregation.
- 24. Fog is liquid dispersed in gas, a class of colloidal system
- 26. Surface phase reaction
- 30. Greater magnitude of charge
- **32.** Gems are solid and gels are semi-solid. This mixture will not form emulsion.
- 34. Greater charge on Al
- 38. Form stable colloids due to solvation
- **42.** Solution particle adsorbs common ion present in the medium.
- **43.** Solvent hating character.
- **44.** All are the same processes.

19

Assertion-Reason Questions

These questions consist of an *assertion* in column 1 and a *reason* in column 2. Use the following key to choose the appropriate answer.

- (a) If the *assertion* as well as the *reason* are correct, and the *reason* is the correct explanation of the *assertion*.
- (b) If the *assertion* as well as the *reason* are correct, but the *reason* is not the correct explanation of the *assertion*.
- (c) If the *assertion* is correct but the *reason* is not.
- (d) If the *assertion* is incorrect but the *reason* is correct.

Assertion

Reason

- We cannot measure position and momentum with arbitrary precision.
- Bohr's theory gave reasonably correct spectroscopic results for hydrogenic species even though it used classical mechanics for its derivation.
- **3.** An ideal gas follows a hyperbolic relation between *p* and *V* in isothermal conditions.
- Spectral wave numbers bear a linear relationship with the square of the principal quantum number.
- **5.** All collisions between reactants do not yield the desired product.

There is a deep fundamental flaw in the very measuring process.

The semiclassical approach is valid to some extent for spectroscopic derivations in the context of hydrogenic species.

The percentage change in p added to that in V equals zero (T = constant).

This is a direct consequence of Bohr's energy condition.

An activation complex can form only when the required steric and energy conditions are met.

- **6.** N_2F_2 never obeys the ideal gas law.
- 7. The *T-P* curve in the Joule— Thomson experiment does not represent an isenthalpic process.
- By appropriately choosing a catalyst, we can significantly increase yield.
- Alpha-emission is classically forbidden.
- A photochemically induced reaction does not yield products based on the equilibrium constant.
- The nucleus of gold is stable even though there is a very strong Coulomb repulsion between the protons.
- **12.** For every process in an isolated system, the entropy increases.
- **13.** An ideal Carnot cycle has 100% efficiency.
- **14.** Lead is most effective in shielding radiation.
- 15. Henry's law and Raoult's law are not independent, i.e., one can be derived from the other.
- **16.** A salt bridge is essential for the working of a galvanic cell.
- Electronic penetration is observed in almost all multielectronic species.
- **18.** H⁺ and OH⁻ react very rapidly to yield H₂O.
- K-shell electron capture is detected by analysing the wavelength of the X-ray photon emitted.

It has a very high viscosity.

The enthalpy remains constant during the entire Joule–Thomson expansion.

Catalysts increase the rate constant to a large extent.

It involves quantum tunnelling.

A photochemical energy distribution is not a classical Boltzmann distribution.

The inverse square Coulomb force is exactly balanced by another inverse square force which is very powerful—the nuclear force.

 $\Delta S \ge \int \delta q_{\rm r} / T$

All steps of the Carnot cycle are reversible, and are assumed to be frictionless.

It is very stable, and many radioactive reactions finally yield lead.

The partial pressure is directly proportional to the mole fraction of the concerned species for ideal solutions.

A salt bridge greatly enhances ionic mobility, μ_{ion} .

The radial distribution function shows multiple peaks of varying amplitude.

The equilibrium constant K_{eq} for the reaction is very high (~10¹⁵).

The wavelength of the X-ray photon is characteristic of the daughter element and not the parent element.

- **20.** Tin-plating of iron is not recommended as a guard against corrosion.
- **21.** A fuel cell gives much more energy than a conventional Galvanic cell.
- **22.** Activity is not an experimentally measurable quantity.
- 23. Bond dissociation energy refers to energy changes during the gaseous state.
- 24. In deriving the Nernst equation, we directly equate electric work with change in free energy.
- **25.** The term "molecularity of the overall reaction" is meaningless.
- **26.** The Zeeman effect could not be predicted by the Bohr model.
- **27.** A reversible process is associated with maximum work.
- **28.** The entropy of a system always increases after a spontaneous process.
- The van't Hoff equation in the conventional form cannot be used for macromolecular solutes.
- **30.** The volume of a hexagonal ice lattice is $a^2c \sin 120^\circ$.
- Phenolphthalein is used as an indicator during the titration of oxalic acid against sodium hydroxide.

Once the surface of the tin-plating is scratched, there is rapid oxidation of iron because of the formation of an electrolytic couple.

The exchange current density of a fuel cell is very low.

Activity is just a type of effective mole fraction and $a = p/p^0$.

Bonds cannot break in the liquid and solid phases.

This is just an approximation; we neglect the P-V work of the system.

Molecularity refers to the order of the rate-determining step.

The Zeeman effect is the result of the interaction of the magnetic field *B* and the quantized angular momentum, *L*.

In a reversible expansion the system expands/contracts against maximum possible pressure.

The Clausius inequality rule states that

$$\int_{\text{cycle}} \delta q/T \le \Delta S \text{ for any process.}$$

The van't Hoff equation is the first term of the virial-like equation

$$\Pi = [B] RT \{1 + B[\vec{B}] + ...\}$$
 where $[B] = c/M$ ($c = \text{mass concentration}$, $M = \text{molecular mass}$, $B = \text{virial coefficient}$).

$$a = b \neq c$$
; $\alpha = \beta = 90^{\circ}$, $\gamma = 60^{\circ}$

The pH range of phenolphthalein is from 8 to 9.6.

32. CH₃NO₂ is less polar than CH₃NH₂.

33. The reaction of isopropyl alcohol with HF and that with HCl are endothermic and exothermic respectively.

34. In the Nernst equation, the stoichiometric coefficients of a balanced chemical reaction do not matter.

 The efficiency of a Carnot cycle is independent on the working substance employed.

36. ΔH is not always equal to $C_p \Delta T$.

37. A catalyst can be used almost infinite times for a particular chemical reaction

38. The NaCl molecule has 8 corners and 12 edges shared.

39. The standard electrode potential of alkali metals are uniform and close to 3.0 V.

40. Quasi-crystals form when certain molten alloys cool very slowly.

In CH_3NO_2 , the formal charge on N is ± 1 .

 ΔH° is negative in the former case and positive in the latter.

The change in the form of K is taken care of by the premultiplying factor of n (number of moles).

All irreversible engines are necessarily less efficient than the Carnot engine.

 C_p is a polynomial function of temperature.

A catalyst is not consumed during a chemical reaction.

NaCl has a 6:6 coordination system.

All alkali metals have very low hydration energy.

Quasi-crystals have short-range as well as long-range order in their arrangements.

Answers

1. c	2. c	3. b	4. d	5. a
6. c	7. b	8. d	9. a	10. a
11. c	12. d	13. c	14. b	15. b
16. c	17. a	18. b	19. b	20. a
21. c	22. d	23. a	24. d	25. c
26. b	27. a	28. c	29. a	30. c
31. a	32. d	33. c	34. a	35. b
36. a	37. d	38. d	39. c	40. d

Matching-Type Questions (Chapterwise)

Atomic Structure—I

1.	Energy	of a	massless	particle
1.	LHEIEV	OI a	1110331033	particle

2. Work function

ψψ*

4. Number of lobes in a 3d orbital other than 3d₂²

5. Energy and angular momentum

6. Mutual repulsion of atomic electrons

7. Mass times velocity times radius

8. The frequency distribution of the radiation emitted from a black body

9. Probability of finding an electron in a given region

10. Black-body radiation

(a) $a^2 + b^2$

(b) Four

(c) E = pc

(d) Connected to orbital motion

(e) \overrightarrow{L}

(f) Minimum energy needed to liberate an electron from the surface of a metal

(g) Quanta

 $(h) \int_{x_1}^{x_2} |\psi|^2 dx$

(i) Hund rule

(j) Temperature-dependent

Atomic Structure—II

1. Pauli exclusion principle

2. Nodal plane

3. Number of angular nodes

4. Number of radial nodes

(a) Lower energy state

(b) $\sqrt{l(l+1)} \hbar$

(c) $\frac{\sqrt{3}}{2}i$

(d) Higher energy state

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- 5. The $3d_{z^2}$ orbital looks like a cone having angles
- (e) The plane dividing regions of opposite sign of the wave function
- **6.** Magnitude of the spin angular momentum of an electron
- (f) $R_{\infty}hc$
- 7. Magnitude of the orbital angular momentum of an electron
- (g) 55° and 125°

8. Bound-state energy

- (h) l
- 9. Unbound-state energy
- (i) n l 1

10. Ionization energy

(j) Explained by determinant

Nuclear Chemistry

1. 14C

- (a) Gamma rays
- **2.** Survival probability of a radioactive nucleus
- (b) K-electron capture

3. Breeder reactor

- (c) Neptunium series
- 4. De-excitation of nuclei
- (d) $N_0 N_0 e^{-\lambda t}$
- **5.** Addition of an electron and a positron
- (e) Shortest half-life

6. 241 Pu $\rightarrow ^{209}$ Bi

(f) Proton-proton cycle

7. Neutron decay

(g) Annihilation

8. Nuclear fusion

(h) Production of the same fissile material as it uses

9. Emission of β^+

(i) β^- emitter

10. ²¹²Po

(j) Antineutrino

Chemical Bonding

- Maximum number of carbon atoms arranged linearly in the molecule CH₃C≡CH₂=CH₂
- (a) 12 pentagons and 20 hexagons

- **2.** Number of π bonds in hex-1,3-diene-5-yne
- (b) Graphite

3. A C₆₀ molecule has

- (c) Neopentane
- 4. Species having zero dipole moment
- (d) ClO₂

5. All carbon atoms are sp²-hybridized

- (e) An electron is removed from the $\pi^* MO$
- 6. Intramolecular hydrogen bonding
- (f) IF₆
- 7. Species not obeying the octet rule
- (g) An electron is removed from the $\sigma\,MO$
- 8. Expanded octet structure
- (h) $Ni(CN)_4^{2-}$

9. $O_2 \to O_2^+$

(i) B₂

10. $N_2 \rightarrow N_2^+$

- (j) Four
- **11.** Species having two unpaired electrons
- (k) I_3^-

12. Linear structure

- (l) C_{60}
- (m) 2-Nitrophenol
 - (n) Ethyl acetoacetate (enol form)
 - (o) PCl₅
- (p) 20 pentagons and 18 hexagons

The Gaseous State

- **1.** No gas can liquefy above T_c
- (a) Supercritical fluid

2. Boyle temperature

- (b) $\frac{1}{C_{p, m}} \left(\frac{2a}{RT} b \right)$
- 3. Inversion temperature
- (c) Unrealistic oscillations in the van der Waals isotherms

- 4. Van der Waals loops
- (d) The equal-area rule for the replacement of the van der Waals loops by horizontal straight lines
- 5. Maxwell construction
- (e) The temperature at which the Joule–Thomson coefficient changes its sign
- 6. Graham's law of diffusion
- (f) The temperature at which $\frac{dZ}{dp} = 0$
- 7. Dense fluid phase above $T_{\rm c}$ and $p_{\rm c}$
- (g) Molecules have high thermal energy, and intermolecular forces are not powerful enough to make molecules stick together
- 8. Joule-Thomson coefficient
- (h) $\frac{u_{\text{rms }1}^2}{u_{\text{rms }2}^2} = \frac{M_2}{M_1}$

The Liquid State

(Viscosity and Surface Tension)

- 1. Superfluid helium
- 2. Viscosity
- 3. Newtonian flow
- 4. Coefficient of viscosity
- 5. Energy flux
- 6. Momentum of flux

- (a) $\frac{dT}{dz}$
- (b) $-\eta \frac{dv_x}{dz}$
- (c) $\frac{dv}{dz}$
- (d) Migration of linear momentum down a velocity gradient
- (e) Flow of liquid through a series of layers
- (f) A liquid phase that flows without viscosity

Oxidation-Reduction

- 1. $Cr_2O_7^{2-} \to CrO_5$
- 2. $BrO_3^- + Br^- + H^+ \rightarrow Br_2$
- 3. $I^- + I_2 \rightarrow I_3^-$
- **4.** MnO₄⁻ in acidic medium
- 5. MnO₄ in alkaline medium
- **6.** MnO₄⁻ in neutral or slightly alkaline medium
- 7. $2MnO_4^- + 5C_2O_4^{2-} + 16H^+ \rightarrow 2Mn^{2+} + 10CO_2 + 8H_2O$
- **8.** Oxidation number(s) of Fe in Fe₃O₄
- 9. Oxidation number(s) of Pb in Pb₃O₄
- **10.** $Fe_3O_4 + Al \rightarrow Al_2O_3 + Fe$

- (a) $\frac{M}{5}$
- (b) $\frac{M}{1}$
- (c) $\frac{M}{3}$
- (d) 24 electrons
- (e) $-\frac{1}{3}$
- (f) +2, +4
- (g) Change in oxidation number of the metal is zero
- (h) Change in oxidation number of carbon is one
- (i) Disproportionation reaction
- (i) +2, +3

Modern Concepts of Acids and Bases

- 1. Species having no conjugate base
- (a) $Be(OH)_2$
- 2. Species that can act neither as a conjugate acid nor as a conjugate base
- (b) N₂H₄

- 3. Species that can act both as a conjugate acid and a conjugate
- (c) BeCl₂
- 4. Species that can act both as a Brønsted base and a Lewis base
- (d) Base

5. Monoprotic acid(s)

- (e) Acid
- 6. Compound having amphoteric character
- (f) $Ca(HCO_3)_2$
- 7. Compound that does not act as a Lewis acid
- (g) HCO₃

8. Protophilic solvent

- (h) C_6H_6
- 9. In a nitrating mixture, HNO₃ acts as a(n)
- (i) SO_4^{2-}
- 10. Salts do not exist in the solid state
- (j) : CH_2
- (k) : CN⁻
- (l) H_3BO_3 and H_3PO_2

Volumetric Analysis (Titration)

- 1. Titration of FeSO₄ versus KMnO₄
- 2. Titration of a mixture of Na₂CO₃ and NaHCO₃
- 3. Titration of Na₂C₂O₄ versus NaOH
- 4. Titration of boric acid and NaOH solution
- 5. Titration of CuSO₄ solution and Na₂S₂O₃ solution
- **6.** Titration of Mohr salt versus $K_2Cr_2O_7$
- 7. Titration of NH₂OH versus $KMnO_4$
- 8. Estimation of CuSO₄ by producing CuSCN
- 9. Titration of AgNO₃ versus NaCl
- 10. Estimation of a mixture of Na₂C₂O₄ and NaHC₂O₄ versus a strong base

- (a) Phenolphthalein indicator
- (b) Gravimetrically
- (c) No indicator
- (d) Phenolphthalein and methyl orange indicators
- (e) K₂CrO₄ indicator
- (f) Acid-base and redox titration
- (g) Barium diphenylamine sulphonate indicator
- (h) Methyl red indicator
- (i) Starch solution indicator
- (j) Methyl orange indicator

Thermodynamics and Thermochemistry

- 1. Clausius inequality
- 2. Work done in adiabatic process
- **3.** Work done in isothermal reversible expansion
- **4.** Entropy of all perfectly crystalline substances
- 5. Criteria of spontaneity
- 6. Spontaneous process
- 7. $\Delta_{\text{sub}}H^{\circ}$
- **8.** *dq*_{rev}
- 9. $\frac{dq_{\text{rev}}}{T}$
- **10.** $\frac{\partial U}{\partial V}\Big]_S$

- (a) Perfect differential
- (b) Zero when $T \rightarrow 0$
- (c) Imperfect differential
- (d) –p
- (e) $dS \ge \frac{dq}{T}$
- (f) $\Delta_{\text{fus}}H^{\circ} \Delta_{\text{vap}}H^{\circ}$
- (g) $dS_{U, V} \ge 0$ and $dU_{S, V} \le 0$
- (h) $\frac{p_1V_1 p_2V_2}{\gamma 1}$
- (i) $-nRT \ln \frac{V_2}{V_1}$
- (j) $\Delta S > 0$
- (k) $\Delta S < 1$
- (l) $\Delta_{\text{fus}}H^{\circ} + \Delta_{\text{vap}}H^{\circ}$

Chemical Equilibrium

- 1. Reaction quotient
- **2.** Equilibrium constant (K_{eq})
- 3. $\frac{\partial K}{\partial p}$
- **4.** Extent of dissociation for $A(g) \rightleftharpoons 2B(g)$
- 5. $N_2O_4(g) \rightleftharpoons 2NO_2(g)$
- 6. $CO(g) + 2H_2(g) \rightleftharpoons CH_3OH(g)$

- $(a) \left(\frac{1}{1+4p/K}\right)^{1/2}$
- (b) High temperature and low pressure will favour the formation of the product
- (c) Low temperatue and high pressure will favour the formation of the product
- (d) $\prod^{J} a_{J}^{\nu_{J}}$
- (e) Zero
- (f) $\left(\prod^{J} a_{J}^{\nu_{J}}\right)_{eq}$

Chemical Kinetics

- 1. Acidic hydrolysis of ethyl acetate
- (a) mol $L^{-1} s^{-1}$
- **2.** Alkaline hydrolysis of ethyl acetate
- (b) Studied by polarimeter
- **3.** Unit of *A* (Arrhenius parameter)
- (c) Second-order reaction
- 4. Time required for the concentration of a reactant to fall to 1/e of its initial value
- (d) Saponification
- 5. Reciprocal of rate constant
- (e) Straight line
- 6. Decomposition of CH₃CHO
- (f) First-order kinetics
- 7. $S_2O_8^{2-} + 2I^- \implies 2SO_4^{2-} + I_2$
- (g) Same as that of rate constant
- **8.** Unit of rate constant of zero-order reactions
- (h) Time constant
- **9.** Inversion of cane sugar
- (i) Fractional order
- **10.** Plot of $t_{1/2}$ versus concentration
- (j) Half-life constant
- (k) Second-order kinetics
- (l) Third-order kinetics

Ionic Equilibrium

- 1. Autoprotolysis constant
- (a) Boric acid and borax

2. The pH of 0.1-M HA $(K_a = 10^{-5} \text{ mol L}^{-1})$

- (b) 1.0
- 3. The pH range 6.8–9.2
- (c) Two buffers
- 4. The pH of 0.1-M NH₄Ac is calcualted by the equation
- (d) $pH = pK_a + log \frac{[salt]}{[base]}$
- 5. Sodium tetraborate in water
- (e) 3.0
- 6. The pH of a mixture of 0.1 mol CH_3NH_2 ($K_b = 5 \times 10^{-4}$ mol L^{-1}) and 0.08 mol HCl
- (f) Three buffers
- 7. The pH of a mixture of 75 mL of a 0.2-N HCl solution and 25 mL of a 0.2-N NaOH solution
- (g) Decrease in solubility
- 8. Effectiveness of a buffer
- (h) $a_{H,O^{+}} a_{OH^{-}}$
- 9. Selection of indicators is based on
- (i) Alkaline

- 10. Addition of KBr to AgBr
- (j) 10.1

11. H₃PO₄ can have

- (k) Buffer capacity
- (l) pK_{In}
- 12. The pH value of a 0.1-M NH_4Cl solution is determined by using the equation
- (m) pH = $\frac{1}{2}$ p K_w + $\frac{1}{2}$ p K_a + $\frac{1}{2}$ log c
- (n) $pH = \frac{1}{2} pK_w \frac{1}{2} pK_a \frac{1}{2} pK_b$
- (o) Increase in solubility
- (p) $pH = \frac{1}{2}pK_w \frac{1}{2}pK_b + \frac{1}{2}\log c$

Electrochemistry

1. Electrochemical cell

(a) $\frac{NE^{\circ}F}{2.303RT}$

2. Electrolytic cell

- (b) Action potential
- 3. Electrolyte-concentration cell
- (c) Electrochemical process
- 4. Electrode-concentration cell
- (d) Acidic cell

5. log *K*

- (e) $\frac{dE^{\circ}}{dT}$
- **6.** The p.d. across the membrane of a neurone when a signal is propagating
- (f) Kohlrausch's law

7. Rusting of iron

(g) Law of independent migration of ions

8. Weak electrolytes

(h) Electrodes have different concentrations

9. Strong electrolytes

- (i) $\frac{\Delta S^{\circ}}{ZF}$
- 10. Temperature coefficient of a cell
- (j) Not a rechargeable battery

11. Lead storage battery

(k) Nonspontaneous reaction is driven by an external source of current **12.** $\Lambda_{\rm m} = \Lambda_{\rm m}^{\circ} - Kc^{1/2}$

- (l) Electrolytes have different concentations
- (m) Molar conductance falls sharply to low values as concentration increases
- (n) Molar conductance varies slightly with concentration
- (o) Chemical reactions do electrical work

Colligative Properties of Solutions

- 1. The vapour pressure of $Na_2CO_3\cdot 10H_2O$ is greater than the water vapour pressure of the atmosphere
- (a) Beckmann method
- 2. The vapour pressure of CaCl₂·6H₂O is lesser than the water vapour pressure of the atmosphere
- (b) Complexation
- **3.** Absorption of water vapour from the atmosphere without dissolution
- (c) $p = p_A^* p_B^* / p_A^* + (p_B^* p_A^*) y_A$
- **4.** 10% glucose (w/v) and 10% urea (w/v) solutions are
- (d) Positive $\Delta_{\rm mix}H$, $\Delta_{\rm mix}V$ and positive minimum boiling azeotrope
- **5.** Composition of vapour in a binary solution
- (e) Negative $\Delta_{mix}H$, $\Delta_{mix}V$ and negative maximum boiling azeotrope

6. Azeotropic mixture

- (f) Precipitation
- **7.** The elevation of freezing point takes place during
- (g) Boiling without changing
- 8. During mixing of H₂O and HCl
- (h) Effervescence
- **9.** During mixing of H₂O and CH₃CO₂C₂H₅
- (i) Isotonic solutions
- **10.** Most accurate method for the measurement of molar mass
- (j) $y_A = x_A p_A^* / p_B^* + (p_A^* p_B^*) x_A$
- 11. 34.2% sucrose (w/v) and 6% urea (w/v) solutions are
- (k) $y_A = p_A^* p_B^* / p_A^* + (p_B^* p_A^*) y_A$

- **12.** Total vapour pressure of a mixture of two binary liquids
- (l) Not isotonic solutions
- (m) Landsberger's method
 - (n) Deliquescence
 - (o) Hygroscopic

Solid-State Chemistry

1. Layer structures

- (a) ZnS
- 2. Materials having the conductivity range 10^{-6} – 10^{-4} S m⁻¹
- (b) Colours in alkyl halides

3. Supercooled liquid

- (c) Body-centred-cubic structures
- 4. Materials having conductivity range 10^{-10} – 10^{-20} S m $^{-1}$
- (d) NaCl
- **5.** Substance having Schottky defects
- (e) AgBr
- **6.** Substance having a fluorite structure
- (f) Insulators
- 7. Substance having a rutile structure
- (g) Semiconductors
- **8.** Substance having Frenkel defects
- (h) Coordination numbers 6, 3
- Devices that converts electrical energy to mechanical strains and vice versa
- (i) Graphite and CdI₂
- **10.** Substance having both Schottky and Frenkel defects
- (j) Glass

11. Creation of F-centres

- (k) Coordination numbers 8, 4
- **12.** Even value of h + k + l

(l) Piezoelectric crystals

Surface Chemistry and Colloids—I

1. Physical adsorption

(a) Synthetic detergents and proteins

2. During adsorption

(b) Multilayer phenomenon

3. Isoelectronic points

(c) Synthetic rubber and vulcanization

4. Chemical adsoption

(d) Freundlich adsopriton isotherm

5. Colloidal electrolytes

(e) Langmuir adsorption isotherm

- 6. Macromolecular colloids
- (f) ΔG° , ΔH° , ΔS° are negative
- 7. Plot of $\log \frac{x}{m}$ against pressure under low pressure
- (g) Monolayer phenomenon
- 8. Plot of $\log \frac{x}{m}$ against pressure under any pressure
- (h) No migration of ions
- (i) ΔG° and ΔH° are negative, and ΔS° is positive

Surface Chemistry and Colloids—II

1. Emulsifying agent

(a) Colloid-size clusters of molecules

2. Gel

(b) Surfactant

3. Flocculation

(c) An orderly arrangement of micelles

4. Coagulation

(d) Peptization

5. Schultze-Hardy rule

(e) A semirigid mass of a lyophilic sol having a network structure

6. Micelles

- (f) Reversible aggregation of colloidal particles
- 7. Lyotropic mesomorph
- (g) Hydrophobic colloids flocculate most efficiently by ions of opposite charge and high charge number

 \leftrightarrow d

- **8.** Conversion of solution to sol by adding electrolytes
- (h) Irreversible aggregation of colloidal particles

Answers

Atomic Structure—I

$1. \leftrightarrow c$	$2. \leftrightarrow \mathbf{f}$	$3. \leftrightarrow a$	$4. \leftrightarrow \mathbf{b}$	5.

$$\textbf{6.} \leftrightarrow i \hspace{1cm} \textbf{7.} \leftrightarrow e \hspace{1cm} \textbf{8.} \leftrightarrow j \hspace{1cm} \textbf{9.} \leftrightarrow h \hspace{1cm} \textbf{10.} \leftrightarrow g$$

Atomic Structure—II

1.
$$\leftrightarrow$$
 j 2. \leftrightarrow e 3. \leftrightarrow h 4. \leftrightarrow i 5. \leftrightarrow g 6. \leftrightarrow c 7. \leftrightarrow h 8. \leftrightarrow a 9. \leftrightarrow d 10. \leftrightarrow f

Nuclear Chemistry

- $\textbf{1.} \leftrightarrow i$
- $\mathbf{2.} \leftrightarrow \mathbf{d}$
- $3. \leftrightarrow h$
- $\mathbf{4.} \leftrightarrow \mathbf{a}$
- $5. \leftrightarrow g$

- $6. \leftrightarrow c$
- 7. ↔ j
- $8. \leftrightarrow f$
- $9. \leftrightarrow b$
- **10.** ↔ e

Chemical Bonding

- $1. \leftrightarrow j$
- $2. \leftrightarrow j$
- $3. \leftrightarrow a$
- $4. \leftrightarrow c, h$
- $5. \leftrightarrow b, 1$

- $\textbf{6.} \leftrightarrow m \text{, n}$
- $7. \leftrightarrow d$
- $8. \leftrightarrow f, o$
- $9. \leftrightarrow e$
- 10. \leftrightarrow g

- 11. \leftrightarrow i
- $\textbf{12.} \leftrightarrow k$

The Gaseous State

- $1. \leftrightarrow g$ $6. \leftrightarrow h$
- $\mathbf{2.} \leftrightarrow \mathbf{f}$ $\mathbf{7.} \leftrightarrow \mathbf{a}$
- $3. \leftrightarrow e$ $8. \leftrightarrow b$
- $4. \leftrightarrow c$
- $\mathbf{5.} \leftrightarrow \mathbf{d}$

- The Liquid State
 - $1. \leftrightarrow f$
- $\mathbf{2.} \leftrightarrow \mathbf{d}$
- $3. \leftrightarrow e$
- $4. \leftrightarrow b$
- $5. \leftrightarrow a$

 $\textbf{6.} \leftrightarrow c$

Oxidation-Reduction

- $\textbf{1.} \leftrightarrow g$
- $\textbf{2.} \leftrightarrow i$
- $\textbf{3.} \leftrightarrow e$
- $\textbf{4.} \leftrightarrow a$
- $5. \leftrightarrow b$

- $\textbf{6.} \leftrightarrow c$
- $7. \leftrightarrow h$
- 8. \leftrightarrow j
- $9. \leftrightarrow f$
- $\textbf{10.} \leftrightarrow d$

Modern Concepts of Acids and Bases

- 1. \leftrightarrow j 6. \leftrightarrow a
- $\mathbf{2.} \leftrightarrow \mathbf{i}$ $\mathbf{7.} \leftrightarrow \mathbf{c}$
- $3. \leftrightarrow g$ $8. \leftrightarrow b$
- $\mathbf{4.} \leftrightarrow \mathbf{k}$ $\mathbf{9.} \leftrightarrow \mathbf{e}$
- $5. \leftrightarrow l$ $10. \leftrightarrow g$

- Volumetric Analysis
 - $\textbf{1.} \leftrightarrow c$
- $\mathbf{2.} \leftrightarrow \mathbf{d}$
- $3. \leftrightarrow a$
- $\textbf{4.} \leftrightarrow h$
- $5. \leftrightarrow i$

- $6. \leftrightarrow g$
- $7. \leftrightarrow c$
- $8. \leftrightarrow b$
- $9. \leftrightarrow e$
- $\textbf{10.} \leftrightarrow f$

Thermodynamics and Thermochemistry

- **1.** ↔ e **6.** ↔ j
- 2. \leftrightarrow h 7. \leftrightarrow l
- **3.** ↔ i **8.** ↔ a
- $4. \leftrightarrow b$ $9. \leftrightarrow b$
- $\mathbf{5.} \leftrightarrow \mathbf{g}$ $\mathbf{10.} \leftrightarrow \mathbf{d}$

- Chemical Equilibrium
 - $1. \leftrightarrow d$
- $\textbf{2.} \leftrightarrow f$
- $3. \leftrightarrow e$
- $\mathbf{4.} \leftrightarrow \mathbf{a}$
- $5. \leftrightarrow b$

 $6. \leftrightarrow c$

Chemical Kinetics

- $\textbf{1.} \leftrightarrow f$
- $\mathbf{2.} \leftrightarrow \mathbf{c, d}$
- $3. \leftrightarrow g$
- $4. \leftrightarrow h, j$
- $5. \leftrightarrow h, j$

- $6. \leftrightarrow i$
- $7. \leftrightarrow k$
- $8. \leftrightarrow a$
- $9. \leftrightarrow b$
- 10. \leftrightarrow e

Ionic Equilibrium

- **1.** ↔ h
- $2. \leftrightarrow e$
- $3. \leftrightarrow a$ $8. \leftrightarrow k$
- $4. \leftrightarrow n$
- $5. \leftrightarrow i$

- $6. \leftrightarrow j$ $11. \leftrightarrow f$
- 7. \leftrightarrow b 12. \leftrightarrow p

- $9. \leftrightarrow 1$
- 10. \leftrightarrow g

- Electrochemistry
 - $1. \leftrightarrow o$
- $\textbf{2.} \leftrightarrow k$
- $3. \leftrightarrow l$
- $\textbf{4.} \leftrightarrow h$
- $5. \leftrightarrow a$

- $\textbf{6.} \leftrightarrow b$
- $7. \leftrightarrow c$
- $\textbf{8.} \leftrightarrow m$
- $\textbf{9.} \leftrightarrow n$
- 10. \leftrightarrow e, i

- 11. \leftrightarrow d
- 12. \leftrightarrow g

Colligative Properties of Solutions

- $\textbf{1.} \, {\leftrightarrow} \, h$
- $\mathbf{2.} \leftrightarrow \mathbf{n}$
- $3. \leftrightarrow 0$
- $4. \leftrightarrow l$
- $5. \leftrightarrow j$

- $\mathbf{6.} \leftrightarrow \mathbf{g}$ $\mathbf{11.} \leftrightarrow \mathbf{i}$
- $7. \leftrightarrow b$ $12. \leftrightarrow c$
- $8. \leftrightarrow e$ $9. \leftrightarrow d$
- **10.** \leftrightarrow a

- Solid-State Chemistry
 - $1. \leftrightarrow i$
- $\textbf{2.} \leftrightarrow g$
- $3. \leftrightarrow j$
- $\textbf{4.} \leftrightarrow f$
- $5. \leftrightarrow d$

- $\textbf{6.} \leftrightarrow k$
- 7. ↔ h
- $\textbf{8.} \leftrightarrow a$
- $9. \leftrightarrow 1$
- 10. \leftrightarrow e

- 11. \leftrightarrow b
- **12.** \leftrightarrow c
- Surface Chemistry and Colloids—I
 - $\mathbf{2.} \leftrightarrow \mathbf{f}$
- $3. \leftrightarrow h$
- $4. \leftrightarrow b$
- $5. \leftrightarrow a$

- $\mathbf{1.} \leftrightarrow \mathbf{g}$ $\mathbf{6.} \leftrightarrow \mathbf{c}$
- $7. \leftrightarrow d$
- $8. \leftrightarrow e$

Surface Chemistry and Colloids—II

- $1. \leftrightarrow b$
- **2.** ↔ e
- $3. \leftrightarrow f$
- $4. \leftrightarrow h$
- $\textbf{5.} \leftrightarrow g$

- **6.** ↔ a
- $7. \leftrightarrow c$
- 0. (/ 1
- $8. \leftrightarrow d$

Matrix-Matching-Type Questions (Mixed)

Matrix A

1. Expansion of gas to fill a cylinder

(a) The work done is zero

2. $E_{\text{Fe}^{2+}|\text{Fe}}^{\circ} = -0.44 \text{ V}$ and

(b) A spontaneous process

$$E^{\circ}(O_2 + 4H^+ + 4e = 2H_2O) = +1.23 \text{ V}$$

3. Uniform mixing of Br₂ vapour and N₂ gas

(c) $\Delta S = -R\Sigma(x_i \ln x_i)$

4. $N_2O_4(g) \rightleftharpoons 2NO_2(g)$

(d) From colourless to brown

Matrix B

1. Minimum half-life period

(a) Polonium-212

2. $\log \lambda = A \log E + B$ is the mathematical form of Geiger–Nuttal rule

(b) Neutron decay

3. Beta negative decay

(c) 4n, 4n + 2 and 4n + 3 series

4. Beta positive decay or electron capture

(d) $\Delta m = m_{\text{parent}} - m_{\text{daughter}} > 0$

Matrix C

1. Transition between two atomic energy levels

(a) X-rays

2. Electron emission from a material

(b) Photoelectric effect

3. Moseley's law

- (c) Hydrogen spectrum
- **4.** Change of photon energy into kinetic energy of electrons
- (d) β-decay

Matrix D

1. Steam engine

(a) Energy conversion

2. Galvanic cell

(b) Redox reactions

3. Lead storage cell

(c) A primary cell

4. Leclanché cell

(d) A secondary cell

Matrix E

1. $(CH_3)_3CBr + H_2O \rightarrow$

- (a) A pseudo-unimolecular reaction
- 2. $HCOOH(aq) + Br_2(aq) \rightarrow$
- (b) A first-order reaction
- 3. $H_2(g) + I_2(g) \rightarrow 2HI(g)$
- (c) A fractional-order reaction
- 4. $CH_3CHO(g) \rightarrow CH_4(g) + CO(g)$
- (d) A zero-order reaction

Matrix F

- **1.** Zinc blende has a hexagonal close-packed structure
- (a) The crystal has a 6 : 6 coordination
- **2.** Wurzite has a cubic close-packed structure
- (b) Space occupancy = 74%
- The crystal having an fcc structure and maintaining equal numbers of cation and anion vacancies
- (c) The crystal has alternate tetrahedral void

- 4. Anisotropic properties
- (d) The crystal has the coordination number 4 and possesses Frenkel defects

Matrix G

Lyophilic sol

(a) Isoelectric point

2. Tyndall cone

(b) Self-stabilized sol

- Process of converting a suspension into a colloidal solution by the addition of electrolyte
- (c) Coagulation
- **4.** Precipitation of a colloidal solution by the addition of electrolyte
- (d) Peptization

Matrix H

- 1. Depression of freezing point
- (a) The vapour pressure of a solution is less than that of the pure solvent
- 2. Elevation of boiling point
- (b) Only solvent molecules solidify at the freezing point
- 3. Elevation of freezing point
- (c) Addition of K₄[Fe(CN)₆]
- **4.** The van't Hoff factor may be less than or greater than one
- (d) During complexation

Matrix I

- 1. Negative deviation from Raoult's law
- (a) Benzene-methanol system
- **2.** Positive deviation from Raoult's law
- (b) Acetone-ethanol system
- 3. Minimum boiling azeotrope
- (c) Water-hydrochloric acid system
- 4. Maximum boiling azeotrope
- (d) $\Delta H_{\text{mix}} > 0$ and $\Delta V_{\text{mix}} > 0$

Matrix J

- 1. Change in Gibbs free energy
- (a) An extensive property

2. Enthalpy change

(b) Efficiency of energy conversion

3. Entropy change

(c) $dw_{\text{add. max}}$

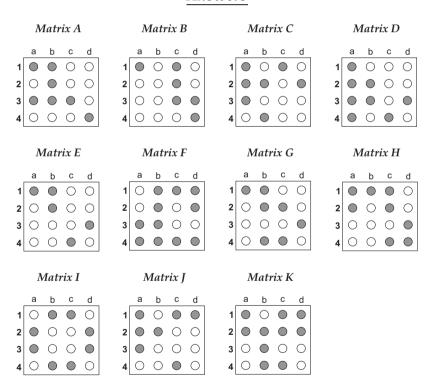
4. dG

(d) $\Delta H - nF \frac{\partial E}{\partial T}\Big]_p$

Matrix K

- **1.** Equivalent conductivity of a weak (a) Kohlrausch's law electrolyte at infinite dilution
- **2.** Molar conductivity at infinite dilution (Λ_m°)
- (b) $v_+ \lambda_+ + v_- \lambda_-$
- 3. Limiting molar conductivity
- (c) $\Lambda_{\rm m} + Kc^{1/2}$
- 4. Parameter which changes with dilution
- (d) Solubility product

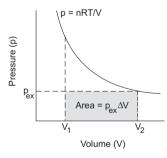
Answers

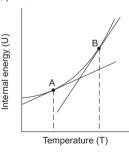


Comprehension-Type Questions

- In thermodynamics, the work done by a gas when it expands against a constant external pressure, $p_{\rm ex}$, is equal to the area $p_{\rm ex}\Delta V$. The work done by a perfect gas when it expands reversibly and isothermally is equal to
 - $-nRT \int_{V_{1}}^{V_{2}} \frac{dV}{V}$. But due to irreversible expansion at the same final pressure, the

work done is equal to $p_{\rm ex}\Delta V$. The work done is greater in reversible process than in irreversible process. The internal energy thus produced in the system is plotted against temperature at constant volume. The slope will be the heat capacity. A similar graph can be plotted between enthalpy (H) and temperature (T). The internal energy during adiabatic process is $C_V\Delta T$. In isothermal process, $p \propto 1/V$, and in adiabatic process, $p \propto 1/V^{\gamma}$.





- 1. The energy transferred as heat to a system at constant volume is equal to
 - (a) ΔU
- (b) IVt
- (c) ΔH
- (d) dq + dw

- 2. From the graph,
 - (a) the heat capacity is lesser at *B* than at *A*
 - (b) $C_p < C_V$

- (c) the heat capacity is greater at *B* than at *A*
- (d) the heat capacity at *B* is equal to that at *A*
- 3. Indicate the correct statement for ideal gases.
 - (a) $C_{v,m} C_{V,m} = R$
 - (b) The slope of the graph of enthalpy versus temperature is steeper than that of the graph of internal energy versus temperature
 - (c) The slope of the graph of internal energy versus temperature is steeper than that of the graph of enthalpy versus temperature
 - (d) $C_{p, m}/C_{V, m} = \gamma = \frac{4}{3}$ for a gas with nonlinear polyatomic molecules
- **4.** Which of the following is the correct expression for adiabatic process?
 - (a) $PV^{\gamma} = \text{constant}$
- (b) $VT^{C_{V,m}/R} = \text{constant}$

(c) $\Delta U = W_{ad}$

- (d) All of these
- **5.** The molar heat capacity of water in equilibrium with ice at constant pressure is
 - (a) zero

- (b) infinity (∞)
- (c) 40.45 kJ K⁻¹ mol⁻¹
- (d) 75.48 J K⁻¹ mol⁻¹
- **6.** In a given process on an ideal gas, dw = 0 and dq < 0. Then, for the gas, the
 - (a) temperature will decrease
- (b) volume will increase
- (c) pressure will remain constant (d) temperatue will increase
- The colligative property of a solution is a property that depends only on the number of solute particles present, not on their identity. An ideal solution is a solution in which all components obey Raoult's law (i.e., $p_A = x_A p_A^{\circ}$) throughout the composition range. The vapour pressure of a binary volatile mixture is $p = p_B^{\circ} + (p_A^{\circ} p_B^{\circ})x_A$. The composition of the vapour is given by $y_A = x_A p_A^{\circ} / \{p_B^{\circ} + (p_A^{\circ} p_B^{\circ})\}x_A$ and $y_B = 1 y_A$. The total vapour pressure of a mixture is $p = p_A^{\circ} p_B^{\circ} / p_A^{\circ} + (p_B^{\circ} p_A^{\circ})y_A$. Azeotrope is a mixture that boils without change in composition. In colligative properties, the elevation of boiling point is given by $\Delta T = K_b m$ and the depression of freezing point by $\Delta T = K_f m$. During dissociation of ionic electrolytes, the van't Hoff factor equals

$$i = 1 + (n-1)\alpha = \frac{M_{\text{th}}}{M_{\text{obs}}}.$$

During association of electolytes,

$$i = 1 - \beta + \frac{\beta}{n} = \frac{M_{\text{th}}}{M_{\text{obs}}}.$$

Here α and β are the degrees of dissociation and association, respectively, of electrolytes.

- 7. The vapour-phase compositions in two binary liquid mixtures follow
 - (a) Boyle's law

(b) Dalton's law

(c) Raoult's law

(d) Henry's law

- 8. The mole fraction of a solute is 0.4. The relative lowering of vapour pressure is (a) 60% (b) 80% (c) 40% (d) 20% The van't Hoff factor is 3 for (a) Na_2SO_4 (b) $Al_2(SO_4)_2$ (c) CaCl₂ with 80% disociation
- (d) K₄[Fe(CN)₆] with 50% dissociation
- 10. The most accurate method for the measurement of molar mass is (a) osmotic pressure (b) ebullioscopy
 - (c) cryoscopy (d) Raoult's law
- 11. The freezing point of a solution containing 0.2 g of acetic acid in 20.0 g benzene is lowered by 0.45°C. The degree of association of acetic acid is (given that K_f for benzene is 5.12 K kg mol⁻¹)
- (a) 0.985 (b) 0.945 (c) 0.845 (d) 0.795 12. In a 0.2-molal aqueous solution of a weak acid HX, the degree of ionization is 0.3. If K_f for water is 1.86 K kg mol⁻¹, the freezing point of
 - the solution is nearly equal to (b) -3.56°C (a) -0.480° C (c) −0.260°C (d) 0.48°C
 - According to the nature of the forces which are holding the gaseous molecular species over the surface of the solid, adsorption can be classified as physisorption and chemisorption. In physisorption, the molecules are held by van der Waals forces. This process has $\Delta H_{\text{ads}}^{\circ} = -20 \text{ kJ mol}^{-1}$; occurs at low temperatures; is reversible, having almost zero activation energy; shows multilayer phenomena; is a function of only the adsorbate; adsorption increases with pressure. On the other hand, in chemisorption, the molecules are held by strong chemical forces. This process has $\Delta H_{\rm ads}^{\circ} = -200 \text{ kJ mol}^{-1}$; occurs at high temperatures, having activation energy; shows unilayer phenomena; is a function of both the adsorbate and adsorbent; adsorption decreases with pressure.

Easily liquefiable gases and consequently larger cohesive force are more easily adsorbed. Adsorption is accompanied by decrease in entropy. The amount of gas adsorbed (x) on the mass of the adsorbent (m) is the function of both pressure and temperature. The variation of amount of gas adsorbed per unit mass with pressure is expressed by empirical equation given by Freundlich as $\frac{x}{m} = kp^{1/n}$ where k, n are constants depending upon the nature of adsorbent. This equation is also applicable in solution. At high pressure of gases, deviation occurs. In order to rectify the deviation, Irving Langmuir modified the above equation as

$$\frac{x}{m} = \theta = \frac{bp}{1 + bp},$$

where $b = \frac{K_a}{K_d}$, $K_a = \text{rate constant for adsorption}$, $K_d = \text{rate constant for adsorption}$

- 13. Physical adsorption is
 - (a) a monolayer phenomenon
- (b) a multilayer phenomenon
- (c) both (a) and (b)
- (d) none of these
- **14.** In the Freundlich adsorption isotherm, when $\frac{1}{n} = 0$, $\frac{x}{m}$ is
 - (a) $\frac{1}{k}$
- (b) zero
- (c) k
- (d) infinity

- 15. In the adsorption isotherm,
 - (a) $\frac{x}{m} = f(p)$

(b) $\frac{x}{m} = f(T)$

(c) $\frac{x}{m}$ = constant

- (d) $\frac{x}{m} = f(V)$
- **16.** Which of the following is true during adsorption?
 - (a) ΔH° is negative.
- (b) ΔG° is negative.
- (c) ΔS° is negative.
- (d) All of these
- 17. At equilibrium in adsorption,

(a)
$$\Delta S^{\circ} = \frac{\Delta H^{\circ}}{T}$$

(b)
$$\Delta G^{\circ} = 0$$

(c)
$$\Delta G^{\circ} < 0$$

(d)
$$\Delta G^{\circ} = \infty$$

- **18.** When k = 1, the surface coverage is
 - (a) 25%
- (b) 75%
- (c) 50%
- (d) 100%
- **19.** The surface coverage, θ , is related to the pressure and the equilibrium constant by the Langmuir isotherm,

(a)
$$\theta = \frac{bp}{1 + bp}$$

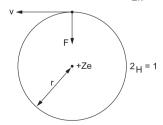
(b)
$$\theta = bp$$

(c)
$$\theta = \frac{1 + bp}{bp}$$

(d)
$$\theta = \frac{b}{v}$$

• The Bohr model is represented by the figure shown, where v = tangential speed. The attractive coulomb force provides the centripetal acceleration v^z/r . The angular momentum, $\overrightarrow{L} = \overrightarrow{r} \times \overrightarrow{p} = mvr$. Thus Bohr's postulate is $mvr = \frac{nh}{2\pi}$.

When the electron and the nucleus are separated by an infinite distance corresponding to $n = \infty$, we have E = 0. We might begin with the electron and electron is separated by an infinite distance and then bring the electron closer to the nucleus until it is in the orbit in a particular state n. Since this state has less energy than E = 0, with which we began. We have 'gained' an amount of energy equal to E_n . Conversely,



if we have an electron in a state n, we can 'take the atom apart' by supplying an energy E_n . This energy is known as the *binding energy* of the state n. If we supply more energy than $E_n = -13.6$ eV, the excess energy will appear as the kinetic energy of free electrons.

The excitation energy of an excited state n is just the energy above the ground state, $E_n - E_1$. Thus for the first excited state, n = 2, has excitation energy = -3.4 - (-13.6) = 10.2 eV. Similarly, the second excited state is 12.1 eV. The photon appears with energy $hv = E_{n_1} - E_{n_2}$ or $hc/\lambda = hc\overline{v} = E_{n_1} - E_{n_2}$. The Balmer formula $\lambda = 364.5$ nm $(n_1^2/n_1^2 - 4)$. There are five spectral series only, Lyman series appears in the absorption spectra, all series are present in the emission spectrum.

20.				for the hydr he first excite				approximately
	(a)	0.27 Å	(b)	1.27 Å	(c)	2.12 Å	(d)	3.12 Å
21.	The w	vavelength of	the	first line of th	е Ва	lmer series is	app	roximately
	(a)	212 nm	(b)	654 nm	(c)	120 nm	(d)	446 nm
22.	If the will b		nth	orbit is direc	tly p	roportional to	o E_n^x	, the value of x
	(a)	+2	(b)	+1	(c)	-2	(d)	-1
23.	The a	ngular mome	entur	n in lowest o	rbit i	s		
	(a)	2	(b)	1	(c)	0	(d)	∞
24.				eing the veloc ergy of the <i>n</i> tl			in th	e nth shell and
	(a)	+2	(b)	-2	(c)	-1	(d)	+1
25.	If $\frac{1}{r^n}$	$\times Z^y$ ($Z = atom$	nic n	umber) the va	alue	of <i>y</i> will be		
	(a)	2	(b)	3	(c)	4	(d)	1
26.	additi weak (b) CF ratio. S conjug (2:1) equati log [s] a little change	on of small an acid $CH_3COH + Na$ $Similarly$, the gate acid (NH molar ratio). The solutions, $pH = p$, $alt]$ (for basic change in pH	noun DOH DOH basic $_4$ Cl) The $_1$ $K_a + 1$ buff I. Rar addit	t of acid or all and a salt (2:1 molar r buffer consist (b) NH ₃ + HC pH of a buffe log [salt] (for er). Addition of acid or a	<pre>cali. 7 of atio) cs of (1 (2 : r sol r ac of a s H = p</pre>	The acidic buffits conjugate (c) CH_3CON (a) week base, 1 molar ratio; ution is given cidic buffer) mall amount of $K_a \pm 1$ or pOF	fer co bas a + H NH ₃) (c) I n by and of aci H = p	in pH by the onsists of (a) the consists of (a) the consists of (a) the consists of (a) the consists of the consist of the co
		$dpH/dn_{acid or}$		control do	(b)	dpH/pK_a or	pK_b	
		$dn_{\text{acid or base}}/$				$dpH/pK_a + p$		

- 27. A buffer solution is one which has
 - (a) reserve acidity
 - (b) reserve alkalinity
 - (c) a pH equal to 7
 - (d) reserve acidity and reserve alkalinity
- **28.** 0.1 mol of CH₃NH₂ ($K_b = 5 \times 10^{-4} \text{ mol L}^{-1}$) is mixed with 0.08 mol of HCl and diluted to 1 L. What will be the H⁺ ion concentration of the resulting solution?
 - (a) $8 \times 10^{-2} \text{ mol L}^{-1}$
- (b) $8 \times 10^{-11} \text{ mol L}^{-1}$
- (c) $1.6 \times 10^{-11} \text{ mol L}^{-1}$
- (d) $8 \times 10^{-5} \text{ mol L}^{-1}$
- 29. The pH of a buffer solution depends upon
 - (a) concentration of salt
 - (b) concentration of either acid or base
 - (c) the ratio of salt and acid or salt and base
 - (d) pK_a or pK_b
- **30.** Which pair will show the common-ion effect?
 - (a) $MgCl_2$ and $Mg(NO_3)_2$
- (b) KCl and HCl
- (c) $C_6H_5NH_2$ and $C_6H_5NH_3Cl$ (d) AuCl₃ and Au(OH)₃
- 31. A buffer solution can be prepared from a mixture of
 - (a) sodium acetate and acetic acid in water
 - (b) sodium acetate and hydrochloric acid in water
 - (c) ammonia and ammonium chloride in water
 - (d) all of these
 - The feasibility of a reaction is determined by ΔG . Consider a gaseous reaction of the type

$$n_1 A + n_2 B \rightleftharpoons n_3 C + n_4 D.$$

 ΔG is related to the reaction quotient as

$$\Delta G = \Delta G^{\circ} + RT \ln Q$$

For the above reaction.

$$\Delta G = \Delta G^{\circ} + RT \ln \frac{C_{\mathrm{C}}^{n_3} \cdot C_{\mathrm{D}}^{n_4}}{C_{\mathrm{A}}^{n_1} \cdot C_{\mathrm{B}}^{n_2}}$$

At equilibrium, $\Delta G = 0$.

$$\therefore \quad \Delta G^{\circ} = -RT \ln K_p \quad \text{or} \quad \Delta G = -RT \ln K_p + RT \ln Q = RT \ln \frac{Q}{K_p}.$$

 ΔG° is related to ΔH° as

$$\Delta G^{\circ} = \Delta H^{\circ} - T\Delta S^{\circ}$$
 (at standard conditions).

The relation between K_v and K_n are

$$K_{p} = \frac{n_{\rm C}^{n_{3}} \cdot n_{\rm D}^{n_{4}}}{n_{\rm A}^{n_{1}} \cdot n_{\rm B}^{n_{2}}} \left(\frac{P}{\Sigma n}\right)^{\Delta n},$$

where Σn = total number of moles after the addition of inert gases. For the reaction

$$n_1A + n_2B \rightleftharpoons n_3C + n_4D$$
,

we have

$$K_p = \frac{p_{\text{C}}^{n_3} \cdot p_{\text{D}}^{n_4}}{p_{\text{A}_1}^{n_4} \cdot p_{\text{B}}^{n_2}} \quad \text{and} \quad K_c = \frac{c_{\text{C}}^{n_3} \cdot c_{\text{D}}^{n_4}}{c_{\text{A}}^{n_4} \cdot c_{\text{B}}^{n_2}} \cdot$$

 K_p and K_c are related by

$$K_p = K_c \left(\frac{c^{\circ} RT}{p^{\circ}} \right)^{\Delta n}.$$

- **32.** The unit of K_c
 - (a) depends on Δn
 - (b) depends upon the speed of reaction
 - (c) depends on the concentration
 - (d) is always mol L⁻¹
- 33. For the feasibility of reaction,
 - (a) $Q = K_n$

(b) $Q < K_n$

(c) $Q > K_v$

- (d) all of these are true
- **34.** Addition of inert gases on equilibrium $PCl_5(g) \rightleftharpoons PCl_3(g) + Cl_2(g)$ at constant volume
 - (a) will shift the equilibrium to the right
 - (b) will shift the equilibrium to the left
 - (c) the equilibrium will remain unchanged
 - (d) will explode the reaction
- **35.** ΔG_f° of $N_2O_4(g)$ and $NO_2(g)$ are respectively -100 kJ mol^{-1} and -50 kJ mol^{-1} . An equilibrium mixture of $N_2O_4(g)$ and $NO_2(g)$ is enclosed in a vessel at a pressure of 20 atm and a temperature of 25°C. In which direction will the reaction $2NO_2(g) \rightleftharpoons N_2O_4(g)$ proceed?
 - (a) Forward direction
 - (b) Reverse direction
 - (c) The reaction will remain in equilibrium.
 - (d) Data are insufficient.
- 36. The spontaneous reaction is not easily possible if
 - (a) both ΔH and ΔS are negative
 - (b) both ΔH and ΔS are positive
 - (c) ΔH is negative and ΔS is positive
 - (d) ΔH is positive and ΔS is negative

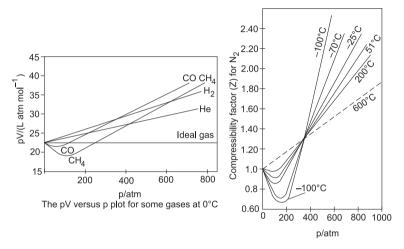
• The rate of reaction is the change of concentration of reactant or product with time. The rate law for the reaction $aA + bB \rightarrow cC + dD$ the rate law is rate = $k[A]^a[B]^b$. The rate of reaction is calculated by knowing k, a and b. The rate laws are determined experimentally. The order of reaction is always defined in terms of reactants.

During the collisions among two A and two B molecules, doubling the number of either type of molecule increases the number of collisions to eight. The species temporarily formed by the reactant molecules as a result of the collision before they form the product is called the *activated complex*. The temperature-dependent rate constant is given by the Arrhenius equation.

In many cases, the sum of a series of simple reactions are called elementary steps or elementary reactions because they represent the progress of the overall reaction at the molecular level. The sequence of elementary steps that leads to product formation is called the *reaction mechanism*. The number of molecules reacting in an elementary step determines the molecularity of a reaction.

- 37. For a zero-order reaction the plot of the rate of reaction versus [A] gives
 - (a) a zigzag line
 - (b) a straight line
 - (c) a straight line with negative slope
 - (d) none of these
- **38.** On which of the following quantities does the rate constant of a reaction depend?
 - (a) Concentrations of reactants(b) Nature of reactions(c) Temperature(d) All the above
- 39. Instantaneous rate of reaction is measured from the
 - (a) graph of time versus concentration
 - (b) molecularity of reaction
 - (c) integration method
 - (d) reaction mechanism of a reaction
- 40. The order of reaction is always defined in terms of
 - (a) product concentration
 - (b) rate constant of a reaction
 - (c) reactant concentration
 - (d) ratio of the product concentration to the reactant concentration
- **41.** During the collision between A and B molecules, doubling the A and B molecules increases the number of collisions to
 - (a) 4 (b) 8 (c) 32 (d) 16
- **42.** In terms of collision theory the rate of reaction is
 - (a) proportional to the number of molecular collision per second
 - (b) frequency of the molecular collisions
 - (c) depend on concentration of reactants
 - (d) all of these

- 43. The number of molecules reacting in an elementary step of a reaction may be
 - (a) 1
- (b) $\frac{1}{2}$ (c) $\frac{3}{2}$
- (d) 5
- The compressibility factor, *Z*, is defined by the relation $Z = \frac{pV}{nRT}$. For ideal gases, Z = 1 at all temperatures and pressures. The deviation from unity is an index of deviation from ideal behaviour.



Z may be calculated from the above equation. Plot of $pV \rightarrow p$ and $Z \rightarrow p$ is shown. Actually p = 0 to p = 100 atm Z changes only from 1.00 to 1.02. Beyond 100 atm Z increases rapidly with pressure and attains value above Z = 1. This temperatue at which a real gas obeys ideal gas law over an appreciable range of pressure is called the Boyle temperature or Boyle point (T_B) . Above T_B the gas shows only positive deviations from ideality, so Z > 1.

After making correction on pressure and volume on ideal gas equation, van der Waals derived the following eqn of state for a real gas for n moles $(p + n^2a/v^2)(v - nb) = nRT$, where a and b are the van der Waals constants. This equation is valid for wider range of pressure.

- **44.** The compressibility factor (*Z*) for a real gas at a very high temperature and a very low pressure is
 - (a) $Z \rightarrow \infty$
- (b) Z > 1
- (c) Z < 1
- (d) Z = 1
- **45.** The compressibility factor (Z) for a real gas at a moderately low pressure is given by
 - (a) $\frac{pV}{nPT}$

(b) $1 + \left(\frac{bp}{nRT}\right)$

(c)	$1 - \left(\frac{bp}{nRT}\right)$			(d)	$1 + \left(\frac{pV}{nRT}\right)$				
The B	oyle tempera	ture	$(T_{\rm B})$ is given	by					
(a)	$\frac{a}{bR}$	(b)	$\frac{a}{R}$	(c)	$\frac{ab}{R}$	(d)	$\frac{bR}{a}$		
	Identify the conditions of pressure and temperature at which a real gas shows maximum deviation from ideality.								
(a)	12 atm, 273 l	K		(b)	7 atm, 273 K				
(c)	12 atm, 373 l	K		(d)	7 atm, 373 K				
	n of the foll tion? (Use the			s the	e least interr	nole	cular force of		
(a)	He	(b)	CO	(c)	CH_4	(d)	CO_2		
	nich of the fo he second fig			tures	s the maximu	ım c	lip N ₂ shows?		
(a)	−25°C	(b)	−70°C	(c)	200°C	(d)	−100°C		
Which value		ing	gases are eas	ily co	ondensed and	l has	the maximum		
(a)	NH_3	(b)	H_2O	(c)	SO ₂	(d)	CO ₂		
(a) NH_3 (b) H_2O (c) SO_2 (d) CO_2 The atmosphere contains both $^{12}CO_2$ and $^{14}CO_2$. These $^{12}CO_2$ and $^{14}CO_2$ are consumed by plants and animals. The concentration of ^{14}C in living organism remains almost constant for longer time. When plants or animal dies, they are no longer in a position to take up CO_2 from the atmosphere. ^{14}C is assimilated by animal or plant during its life time, after that the species begins to decay by emitting β -particles, which have a half-life of 5760 years. The decay rate is 15.3 counts per minute per gram of carbon. Now measuring the decay rate of a sample of dead matter (wood, fossils etc.) at any period									

51. ¹⁴C is produced in the atmosphere by the nuclear reaction of

(a)
$${}^{14}C(n, p)$$

 $t = \frac{2.303}{\lambda} \log \frac{N_0}{N}.$

46.

47.

48.

49.

50.

(b) ${}^{14}N(n, p)$

(c)
$$^{15}N(n, p)$$

(d) ${}^{12}C(n, p)$

52. The radioactivity due to 14 C isotope ($t_{1/2} = 5760$ years) of a sample of wood from an ancient tomb was found to be nearly half that of fresh wood. The tomb is, therefore, about

from the date, it is possible to calculate the life of a living body which had died. The half-life is calculated by using the equation $N = N_0 e^{-\lambda t}$ or

(a) 5760 years old

(b) 2880 years old

(c) 11520 years old

(d) 4320 years old

53.	3. A piece of wood recovered in an excavation has 25.6% as much as 14 C as ordinary wood has when did this piece get burried? ($t_{1/2}$ of 14 C is given 5760 years)								
	-	22649.6 yea	rs		(b)	5662.4 year	'S		
		2831.2 years				11324.8 yea			
E4				arwandad tha				atur in 1060 fa	211
34.		scovery of ca				ei prize ioi c	пеш	stry in 1960 fo	Л
		Willard Lib				Otto Hahn			
	(c)	Hideki Yuk	,			Hans Bethe	9		
55	The a	ge of fossils i	s deta	ermined by					
55.		¹⁴ C		⁴⁰ K	(a)	²³⁸ U	(4)	all of those	
	(a)	C	(D)	K	(c)	U	(a)	all of these	
•	cell from a salt concernation $W_{\rm e,\ max}$ measu $E=E^{\circ}$ concernequiliblike A H ⁺ (aq pH, vi	om the zinc as bridge (an intrated solution in the solution of the solution o	nd en inve on of are to the end of the end o	ter into the corted U-tube; agar-agar + he same. The related to em mf of a cell is e Nernst equal to the calculate the one has to evention). From modynamic	opper) is KC1/ k max f as Δ is det ation compone the param	electrode. To used. Salt be KNO_3/NH_4 ! imum electrode. The electrode is also used to electrode electro	o compridge NO ₃ . 'ical we extend the Notes calculated and a good shift we constitute the Notes calculated and a good shift we constitute the Notes can be shifted as a good shift we constitute the Notes can be shifted as a good shift was constituted as	ctrons leave the plete the circular consists of The mobility of ork is given been to freaction. The reaction depends of the control of the co	of of of at at all
56.	The e	lectrical ener	gy of	a Daniell cel	ll pro	duces due to)		
		movement		ions and ani	ons				
	()	increase in							
	(c)	decrease in							
	(d)	decrease in	ΔG						
57.		implest refer	ence (electrode is t					
	` '	SHE				calomel ele			
	(c)	O ₂ electrode	e		(d)	quinhydro	ne ele	ectrode	
58.	The le	east strongly	redu	cing metal is					
	(a)	copper			(b)	silver			
	(c)	platinum			(d)	gold			
59.	The n	nost strongly	redu	cing metal is	3				
	(a)	sodium	(b)	zinc	(c)	calcium	(d)	iron	

- **60.** The standard potential of a Daniell cell is +1.10 V. The equilibrium constant for cell reaction is
 - (a) 1.5×10^{12}

(b) 1.5×10^{37}

(c) 1.5×10^{22}

- (d) 1.5×10^{30}
- **61.** When a calomel electrode is coupled with hydrogen electrode, the pH of the cell is given by
 - (a) $pH = \frac{E + E_{cal}}{0.0592}$

(b) $pH = \frac{E - E_{cal}}{0.0592}$

(c) $pH = \frac{E_{cal}}{0.0592}$

- (d) $pH = \frac{2E + E_{cal}}{0.0592}$
- **62.** The temperature coefficient is given by
 - (a) $\frac{dE^{\circ}}{dT} = \frac{\Delta S^{\circ}}{n}$

(b) $\frac{dE^{\circ}}{dT} = \frac{\Delta S^{\circ}}{F}$

(c) $\frac{dE^{\circ}}{dT} = \frac{\Delta S^{\circ}}{nF}$

- (d) $\frac{dE^{\circ}}{dT} = \frac{\Delta G^{\circ}}{nF}$
- **63.** The solubility, S, of a sparingly soluble 1 : 1 salt is related to the solubility product, K_s , by
 - (a) $S = K_s^2$

(b) $S^{1/2} = K_s$

(c) $S = K_s^{3/2}$

62. c

61. a

(d) $S = K_s^{1/2}$

Answers

1. b	2. c	3. a, b, d	4. d	5. b	6. a
7. b	8. c	9. d	10. c	11. b	12. a
13. b	14. c	15. a	16. d	17. a	18. c
19. a	20. c	21. b	22. d	23. c	24. b
25. d	26. a	27. d	28. b	29. c, d	30. c
31. d	32. a	33. b	34. c	35. a	36. d
37. b	38. a	39. a	40. c	41. d	42. d
43. a	44. d	45. b	46. a	47. c	48. a
49. d	50. b	51. b	52. a	53. d	54. a
55. a	56. d	57. a	58. d	59. c	60. b

63. d

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Integer-Answer-Type Questions

• Set I •

- 1. Calculate the coordination number of carbon in diamond.
- 2. Calculate the change in the oxidation state of Mn when $KMnO_4$ reacts with concentrated HCl to produce the Mn^{2+} ion.
- 3. Calculate the value of the equilibrium constant, by using the relationship between K and ΔG° . Given $E^0 = 0.028$ V.
- **4.** Compute the time required to pass 36000 C through an electroplating bath using a current of 5 A.
- 5. 20.0 mL of 0.2 M NaOH is added to 50.0 mL of 0.2 M CH₃COOH to give 70.0 mL of the solution. Calculate the additional volume of the 0.2 M NaOH required to make the pH of the solution 4.74 (p K_a for CH₃COOH = 4.74.)
- 6. For a reaction 3A → B + C with a constant rate at every concentration of A, find the order of the reaction with respect to A.
- 7. The difference between wavelengths for the first lines of the Balmer series and the Lyman series is 59.3 nm for a hydrogen-like ion. Calculate the value of Z. (Given $R_H = 109,678 \text{ cm}^{-1}$.)
- 8. 25.4 g of iodine and 14.2 g of chlorine are made to react completely to yield a mixture of ICl and ICl $_3$. Calculate the ratio of the moles of ICl and ICl $_3$ formed.

• Set II •

- 1. The dichromate ion reacts with an NaOH solution to produce the chromate ion. Calculate the change in the oxidation number of chromium during this transformation.
- 2. Indicate the overall order of the following reaction.

$$3I^{-} + S_2O_8^{2-} \rightarrow I_3^{-} + 2SO_4^{2-}$$

- **3.** 10.0 A of current flowed for 1.0 hour through water containing a little sulphuric acid. How many litres of gas were formed at both electrodes at 27°C and 740 mm pressure?
- **4.** Calculate the ratio of the value of any colligative property for a $K_4[Fe(CN)_6]$ solution (assuming complete dissociation) to that of the corresponding property for a sucrose solution.
- **5.** How much calcium (in g) is present in Ca(NO₃)₂ that contains 1.4 g of nitrogen?
- 6. Calculate the resonance energy of NO₂.
 - N O, N = O, N = N and O = O are 222, 607, 946 and 498 kJ mol⁻¹ respectively. Given, ΔH of NO₂ is 134 kJ mol⁻¹.
- 7. 1.5 g of an organic acid was dissolved in water and the volume made up to 300 mL. 10.0 mL of this acid required 12.3 mL of $^{\rm N}\!/_{10}$ NaOH for complete neutralization. If the molecular weight of the acid be 122.0, find the basicity of the acid.
- **8.** Calculate the pH of a 0.1 M CH₃COONH₄ solution. Given, pK_a of CH₃COOH is 4.74 and pK_a of NH₃ is 4.74.

• Set III •

- The chromate ion reacts with an HCl solution to produce the dichromate ion. Calculate the change in the oxidation number of chromium during this transformation.
- 2. How many litres of water should be added to 10.0 g of acetic acid to give a hydrogen ion concentration equal to 1.0×10^{-3} M. ($Ka = 1.8 \times 10^{-5}$.)
- 3. Calculate the order of reaction for the gas-phase reaction $2NO_2 + O_3 \rightarrow N_2O_5 + O_2$, which has the rate constant $K = 2.0 \times 10^4 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1} \text{ at } 300 \text{ K}$.
- 4. The lattice of crystelline CoCl₂ is body-centred tetragonal with 16 formula units per unit cell. How many molecules does the basis consist of?
- An aqueous solution containing an ionic salt with molality equal to 0.1892 freezes at -0.704°C. Calculate the van't Hoff factor of the ionic salt. (K_f for water = 1.86 K/m.)
- **6.** Calculate the value of the reaction quotient Q for the cell $Zn(s) \mid Zn^{2+}(0.01 \text{ M}) \parallel Ag^{+}(0.05 \text{ M}) \mid Ag(s)$.
- 7. 10.0 g of $CaCO_3$ is placed in a 10.0-L vessel at 1100 K. Calculate the approximate amount of $CaCO_3$ unreacted at this temperature, given that K_p for the reaction $CaCO_3(s)$ $CaO(s) + CO_2(g)$ is 0.060 at 1100 K.
- **8.** A gas is kept at 1 atm. It is compressed to $\frac{1}{4}$ th of its initial volume. Calculate the pressure applied .

• Set IV •

- 1. Metallic iron can exist in the β form. (bcc, cell dimension = 2.90 Å) and the gamma (γ) form (fcc, cell dimension = 3.68 Å). The β form can be converted into the γ form by applying high pressures. Calculate the ratio of the density of the β form to that of the γ form.
- **2.** The emf of the cell Ni(s) | Ni $^{2+}$ (0.10 M) || HCl (a = ?) | H₂ (g, 1 atm) | Pt(s) at 25°C is 0.1615 V. Calculate the pH of HCl solution, given $E_{Ni^{2+}/Ni}^0 = -0.25$ V.
- **3.** The rate of decomposition of a gas at a certain temperature is 5.14 and 7.25 in some units for 20% and 5% decomposition respectively. Calculate the order of the reaction.
- **4.** 0.05 M HA is titrated against a 0.05 M NaOH solution. Find the pH at the equivalence point. The dissociation constant for the acid HA is 5.0×10^{-6} .
- **5.** The formula weight of an acid is 123 g mol⁻¹. 100 mL of a solution of this acid containing 39.0 g of the acid per litre was completely neutralized by 95.0 mL of aqueous NaOH containing 40.0 g of NaOH per litre. Calculate the basicity of this acid.
- 7. 10.0 mL of a mixture of nitrogen and oxygen was mixed with 20 mL of hydrogen and the mixture then exploded. The volume after explosion was found to be 21.0 mL. Calculate the volume of nitrogen gas.
- 8. Water rises to a height of 6 cm in a capillary tube of radius r. If the radius is made $\frac{2r}{3}$, calculate the capillary rise in cm.

Answers

Set I

Set II

Set III

1. 8 **2.** 0 **6.** 0 **7.** 3 **3.** 3

4. 2

5. 5

8. 1

1. 0 **2.** 2 **3.** 7

4. 5

5. 2

6. 8

7. 3

8. 7

1. 0

2. 3

3. 2

4. 8

5. 2

6. 4

7. 9

8. 4

Set IV

1. 1 **6.** 4 **2.** 2 **7.** *7* **3.** 2 **8.** 9 **4.** 9

5. 3

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Numerical Problems

• Objective Type •

1. Calculate the concentration of I₃ ions in a standard solution of iodine in 0.5 M KI, making use of the following standard electrode potentials:

2. The boiling point of *n*-hexane at 760 Torr is 68.9°C. The critical

 $I_2 + 2e \rightleftharpoons 2I^-$; $E^\circ = +0.5355 \text{ V}$ and $I_3^- + 2e \rightleftharpoons 3I^-$; $E^\circ = +0.5365 \text{ V}$. The molarity of I^- in the 0.5-M KI solution can be assumed to be

(b) 0.48 mol L^{-1}

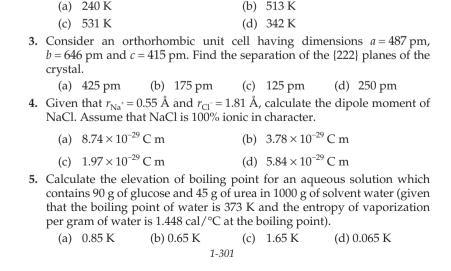
(d) $0.24 \text{ mol } L^{-1}$

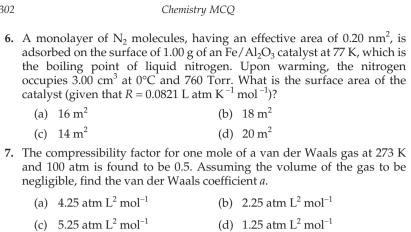
Choose the correct option. Only one option is correct.

 0.5 mol L^{-1} .

(a) 0.12 mol L⁻¹
 (c) 0.88 mol L⁻¹

temperature of *n*-hexane is





- 8. Calculate the Rydberg constant (R_{∞}) if He⁺ ions are known to have a wavelength difference between the first lines of the Balmer and Lyman series equal to $\Delta\lambda = 133.7$ nm.
 - (a) $1.1 \times 10^7 \,\mathrm{m}^{-1}$

(b) $2.2 \times 10^7 \text{ m}^{-1}$

(c) $1.1 \times 10^9 \text{ m}^{-1}$

(d) $1.1 \times 10^5 \,\mathrm{m}^{-1}$

9. A solution contains oxalate and hydrogenoxalate ions but not any other material with either reducing or acidic properties. A sample of the solution decolorizes 48.0 mL of a 0.1-N KMnO₄ solution, and another sample of the same solution of equal volume neutralizes 18.0 mL of a 0.08-N KOH solution. What fraction of the oxalate ions present is protonated to hydrogenoxalate ions?

(a) 0.2

- (b) 0.6
- (c) 0.4

(d) 0.8

- 10. The time required for the 20% completion of a first-order reaction at 27°C is 1.5 times that required for its 30% completion at 37°C. If $A = 3 \times 10^{9} \text{ s}^{-1}$, calculate the time required for the 40% completion at 47°C. Also calculate the activation energy, E_a , for the reaction.
 - (a) 15 s: 88.8 kJ mol⁻¹

(b) 29 s: 76.8 kJ mol⁻¹

(c) 29 s: 90.8 kJ mol⁻¹

(d) 19 s: 67.6 kJ mol⁻¹

11. The emf of the cell

 $Hg(1) | Hg_2SO_4 | H_2SO_4(1 M) | PbSO_4(s) | Pb(s)$

is -0.932 V. The cell reactions are as follows:

$$Hg_2SO_4 + 2e \rightleftharpoons 2Hg(1) + SO_4^{2-}; E^\circ = +0.62 \text{ V}$$

 $PbSO_4 + 2e \rightleftharpoons Pb(s) + SO_4^{2-}; E^\circ = -0.36 \text{ V}$

Given that K_s for Hg₂SO₄ is 1.6×10^{-8} (mol L⁻¹)³, calculate the solubility product of PbSO₄ at 25°C.

(a) $7.0 \times 10^{-12} \, (\text{mol L}^{-1})^2$

(b) $3.0 \times 10^{-11} \text{ (mol L}^{-1})^2$

(c) $6.6 \times 10^{-7} \text{ (mol L}^{-1})^2$

(d) $1.8 \times 10^{-9} \text{ (mol L}^{-1})^2$

12. LiBr, NaBr, KBr and RbBr have the same crystal structure shown by X-ray diffraction. The first three have the fcc structure. The structure of

(b) body-centred cubic(d) both (a) and (c)

RbBr is

(a) face-centred cubic

(c) simple cubic

13.	The conductivity of pure water at 298 K is $5.55 \times 10^{-8} \Omega^{-1} \mathrm{cm}^{-1}$. The molar conductivities of H ⁺ and OH ⁻ ions at infinite dilution are 350 $\Omega^{-1} \mathrm{cm}^2 \mathrm{mol}^{-1}$ and 200 $\Omega^{-1} \mathrm{cm}^2 \mathrm{mol}^{-1}$ respectively. Calculate K_{w} of pure water at 298 K.						
	(a) 1.0	$\times 10^{-12} (mo)$	ol L^{-1}) ²	(b)	1.0×10^{-14} ($\text{mol } L^{-1})^2$	
	(c) 1.0	$\times 10^{-16} (mo$	$(1 L^{-1})^2$	(d)	1.0×10^{-18} ($\text{mol } L^{-1})^2$	
14.	chloride. 'reduced b	What volumy 1 g of 119 S	me of a decin	orma	dichromate	K ₂ Cr ₂ O ₇ to stannie solution would b	ic e
	(a) 525				168 mL		
	(c) 18 i	mL		(d)	336 mL		
15.	5. The heat of neutralization of a weak acid in a normal solution by a strong alkali is 13 385 cal. Assuming that the acid is 14% dissociated in the normal solution, calculate the heat of dissociation of the acid. (a) 236 cal (b) 366 cal (c) 428 cal (d) 192 cal						
16.	mol L ⁻¹ a	nnd 7.32×1 f 20 g aceti	$10^{-5} \text{ mol } L^{-1}$	respe	ctively. Cald	e acid are 1.80×10^{-5} culate the pH of issolved in one litr (d) 4.5	a
17	()	`	,			. ,	4
17.	n_2 are primodel to	ncipal quar be valid. T	ntum numbers The time perio	s of the	ne two states he electron i	$_{1} \rightarrow n_{2}$, where n_{1} and s. Assume the Boh in the initial state in between n_{1} and n_{2}	ır is
	(a) $n_1 =$	$=4n_2$ (k	b) $n_1 = 2n_2$	(c)	$2n_1 = n_2$	(d) $n_1 = n_2$	
18.	an electro	n must be a	ccelerated thr	ough	a potential d		٦,
10	(a) 2 µ		b) 15 μV		10 μV	(d) 5 μV ere are carbon atom	
19.						$\left(\frac{1}{4} \frac{1}{4} \frac{1}{4}\right), \left(\frac{1}{4} \frac{3}{4} \frac{3}{4}\right)$	
						of this crystal if th	
	density of	diamond is	$\mathrm{s}~3.515~\mathrm{g}~\mathrm{cm}^{-3}$?			

(a) 356.6 pm

(c) 556.8 Å

(a) 0.544 V

(c) 0.816 V

20. K_a for acetic acid is 1.8×10^{-5} mol L⁻¹. A decinormal solution of acetic acid is also 0.1-molar with respect to sodium acetate. Calculate the cell potential between a hydrogen electrode in this solution and a decinormal calomel electrode, assuming complete dissociation of the 0.1-M sodium

21. For the reaction $2NO + H_2 \rightarrow 2NOH$, the following reaction mechanisms

acetate solution. E° of the calomel electrode is +0.337 V.

(b) 35.66 cm

(d) 0.568 nm

(b) 0.326 V

(d) 0.618 V

	have been proposed.								
	Mech	anism A.		ν					
	Step I	:	NO	$+ H_2 \stackrel{K}{\rightleftharpoons}$	NOH	[2]			
	Step I	I: NO	OH ₂ -	$+NO \xrightarrow{k} 2$	NOH]			
	Mech	anism B.		K'					
	Step I	:		$2NO \stackrel{K'}{\Longrightarrow}$	N_2C) 2			
	Step I	I:	N ₂ C	$O_2 + H_2 \xrightarrow{k'}$	2NO	Н			
		e basis of thon is proport			isms	we observe	that	the rate	of the
	(a)	$[NO][H_2]$			(b)	$[NO][H_2]^2$			
	(c)	$[NO]^2[H_2]^2$			(d)	$[NO]^2[H_2]$			
22.	If one	mole of a mo	nator	nic gas $\left(\gamma = \frac{5}{2}\right)$	$\frac{5}{3}$ is n	nixed with or	ne mo	le of a di	iatomic
	$gas\Big(\gamma$	$=\frac{7}{5}$, calcula	te th	e value of γ f	or the	e mixture.			
	(a)	2.50			(b)	1.50			
	(c)	1.67			(d)	2.00			
23.	anoth	ctivity of a er 140 days, mple is foun	the a	activity redu					
	(a)	6000 Bq			(b)	9000 Bq			
	(c)	24 000 Bq			(d)	3000 Bq			
24.	energ	nergy of a ph y of the pho length of the	ton l	be E. If λ_1 ar	$nd \lambda_2$	are respecti	vely	the de E	Broglie
	$\frac{\lambda_1}{\lambda_2}$ wil	ll be proporti E°	onal	to					
	(a)	E°	(b)	$E^{1/2}$	(c)	E^{-1}	(d)	E^{-2}	

• Subjective Type •

Solve the following problems.

25. The standard free energy change for the reaction

$$H_2(g, 1 \text{ atm}) + \frac{1}{2} O_2(g, 1 \text{ atm}) \rightarrow H_2O(l)$$

is -240 kJ at 298 K.

(i) Write the galvanic cell which represents the cell reaction

$$H_2(g, 1 \text{ atm}) + \frac{1}{2}O_2(g, 1 \text{ atm}) \rightarrow H_2O(l).$$

(ii) Determine the standard reduction potential for the electrode $OH^-, O_2 \mid Pt$, given that the cell potential for the reaction

$$H_2O(l) \rightarrow H^+ + OH^-$$
 is $E^\circ = -0.6 \text{ V}$.

26. The freezing point of an aqueous solution of $0.1892 \text{ mol kg}^{-1} \text{ KCN}$ is -0.704°C . The freezing point, however, becomes -0.53°C on adding $0.095 \text{ mol Hg}(\text{CN})_2$. Assume that the complex is formed according to the equation

$$Hg(CN)_2 + nCN^- \rightarrow [Hg(CN)_{n+2}]^{n-}$$
.

Find the formula of the complex formed.

27. Auric hydroxide dissolves in hydrochloric acid according to the thermochemical equation

$$Au(OH)_3 + 4HCl \xrightarrow{\Delta H} HAuCl_4 + 3H_2O; \Delta H^\circ = -23 \text{ kcal}$$

and in hydrobromic acid according to the equation

Au(OH)₃ + 4HBr
$$\xrightarrow{\Delta H}$$
 HAuBr₄ + 3H₂O; ΔH° = -36.8 kcal.

On mixing 1 mol HAuBr₄ with 4 mol HCl, there is a heat of absorption of 510 cal. What percentage of bromoauric acid has been transformed into chloroauric acid in the process?

28. The emf of cell

$$Ag(s) | AgNO_3(0.01 \text{ M}) | | AgNO_3(0.001 \text{ M}) | Ag$$

is 0.0579 V at 25°C. The 0.001-M $AgNO_3$ solution is completely dissociated. Calculate the degree of dissociation and the concentration of Ag^+ in the 0.01-M $AgNO_3$ solution.

- **29.** An ideal gas is initially at a temperature T and with a volume V. Its volume is increased by ΔV due to the increase in temperature, ΔT , pressure remaining constant. The quantity δ is defined by $\delta = \Delta V/V\Delta T$. Establish the relation between δ and T.
- **30.** Given that the second dissociation constant for H_2SO_4 is 10^{-2} mol L^{-1} , calculate the pH of a 0.1-M H_2SO_4 solution.

- **31.** Calculate the cell potential of a Daniell cell having 1.0 M Zn²⁺ and originally having 1.0 M Cu²⁺ after sufficient NH₃ has been added to the cathode compartment to make the NH₃ concentration 2.0 mol L⁻¹ (given that $E_{\rm Zn}^{2+}|_{\rm Zn} = -0.76$ V, $E_{\rm Cu}^{2+}|_{\rm Cu} = +0.34$ V, and $K_{\rm forward}$ for Cu(NH₃)₄²⁺ is 1.0×10^{12} mol⁻⁴ L⁴).
- **32.** AB crystallizes in the rock-salt structure, with A: B=1:1. The shortest distance between A and B is $Y^{1/3}$ nm and the formula mass of AB is 6.022 Y u, where Y is any arbitrary constant. Find the density of crystal AB.
- 33. For the galvanic cell

$$\begin{split} &Ag(s) \mid AgCl(s), NaCl(0.2\ M) \mid NaBr(0.02\ M), AgBr(s) \mid Ag, \\ &calculate the emf of the cell at 25°C and assign correct polarity, given that \\ &K_s(AgCl) = 2.8 \times 10^{-10}\ (mol\ L^{-1})^2\ and\ K_s(AgBr) = 3.3 \times 10^{-13}\ (mol\ L^{-1})^2. \end{split}$$

- **34.** Find the quantum number n corresponding to the excited state of an He⁺ ion if on transition to the ground state the ion emits two photons in succession with the wavelengths 108.5 nm and 30.4 nm respectively.
- 35. What amount of heat is to be transferred to oxygen in an isobaric heating process so that the gas may perform 2 J work (given that $\gamma = 1.4$ for O_2)?
- **36.** Calculate the order of the rate-determining step in the pyrolysis of $B_2H_6(g)$ at 100°C from the following data.

Concentration of B_2H_6	Rate of increase of pressure			
(in 10^2 mol L^{-1})	(in 10 ⁴ mol L ⁻¹ h ⁻¹)			
2.153	7.40			
0.433	0.73			

37. In the reaction

$$2N_2O_5 \rightarrow 4NO_2 + O_2$$

the rate can be expressed as

(i)
$$-\frac{d[N_2O_5]}{dt} = k_1[N_2O_5]$$

(ii)
$$\frac{d[NO_2]}{dt} = k_2[N_2O_5]$$

(iii)
$$\frac{d[O_2]}{dt} = k_3[N_2O_5]$$

How are k_1 , k_2 and k_3 related?

- **38.** 5 mol of an ideal gas expands isothermally and reversibly from a pressure of 10 atm to 2 atm at 300 K. What is the largest mass which can be lifted through a height of 1 m in this expansion?
- 39. Hydrogen iodide is very strongly adsorbed on gold but slightly adsorbed on platinum. Assume that the adsorption shows the Langmuir isotherm and predict the order of the HI decomposition reaction on each of the two metal surfaces.

40. KCN, which crystallizes in the rock-salt structure, has a density of $1.52 \,\mathrm{g \ cm^{-3}}$. Calculate d_{100} for the unit cell.

Angriore

	717	130013	
1. d	2. b	3. c	4. b
5. b	6. a	7. d	8. a
9. b	10. d	11. c	12. c
13. b	14. d	15. b	16. b
17. b	18. d	19. a	20. d
21. d	22. b	23. c	24. b
25. (ii) +0.643 V	26. $[Hg(CN)_4]^{2-}$	27. 3.7	28. 96%; $9.6 \times 10^{-3} \text{ mol L}^{-1}$
29. $\delta = \frac{1}{T}$	30. 0.9	31. 0.70 V	32. 5 kg m ⁻³
33. –0.0549 V	34. 5	35. 7 J	36. $\frac{3}{2}$
37. $2k_1 = k_2 = 4k_3$	38. 205 kg	39. 0; 1	40. 6.57 Å

Hints to More Difficult Problems

1.
$$I_2(s) + 2e \Rightarrow 2I^-; E^\circ = +0.5355 \text{ V}$$

$$3I^- \Rightarrow I_3^- + 2e; E^\circ = -0.5365 \text{ V}$$

$$I_2(s) + I^- \Rightarrow I_3^-; E^\circ = -0.0010 \text{ V}$$

As we know, $\Delta G^{\circ} = -nE^{\circ}F = -RT \ln K_{\text{eq}}$.

$$\therefore \log K_{\text{eq}} = \frac{nE^{\circ}}{0.0592} = \frac{2(-0.0010)}{0.0592} = 3.34 \times 10^{-2}.$$

$$K_{eq} = 0.925.$$

$$I_2(s)$$
 + $I^- \implies I_3^ K_{eq} = 0.925 = \frac{x}{0.5 - x}$
- 0.5 - x $\therefore x = 0.24 \text{ mol L}^{-1}$

- 0.5 - x x \therefore $x = 0.24 \text{ mol L}^{-1}$. **2.** We know that $T_b = \frac{2}{3} T_{cr}$ where T_b is the boiling point at 1 atm and T_c the critical temperature.

$$T_c = \frac{3}{2} T_b = \frac{3}{2} (273.15 \text{ K} + 68.9 \text{ K}) \approx 513 \text{ K}.$$

3.
$$\frac{1}{d_{hkl}^2} = \frac{h^2}{a^2} + \frac{k^2}{b^2} + \frac{l^2}{c^2} = \frac{2^2}{(487 \text{ pm})^2} + \frac{2^2}{(646 \text{ pm})^2} + \frac{2^2}{(415 \text{ pm})^2}$$
$$= 2^2 (4.216 \times 10^{-6} + 2.396 \times 10^{-6} + 5.8055 \times 10^{-5}) \text{ pm}^{-2}$$
$$= 64.667 \times 10^{-6} \text{ pm}^{-2}.$$

$$\therefore \quad \frac{1}{d_{hkl}} \approx 8 \times 10^{-3} \text{ pm}^{-1}.$$

Therefore, the separation of the {222} planes is given by

$$d_{hkl} = \frac{1}{8} \times 10^3 \text{ pm} = 125 \text{ pm}.$$

4. The radius of NaCl is given by

$$r_{\text{NaCl}} = r_{\text{Na}^+} + r_{\text{Cl}^-}$$

= $(0.55 + 1.81) \text{ Å} = 2.36 \text{ Å} = 2.36 \times 10^{-10} \text{ m}.$

the dipole moment of NaCl will be

$$\mu = e \times r = (1.602 \times 10^{-19} \text{ C}) \times (2.36 \times 10^{-10} \text{ m})$$

 $\approx 3.78 \times 10^{-29} \text{ C m}.$

5.
$$K_{\rm b} = \frac{RT_{\rm b}^2}{1000L_{\rm vap}} = \frac{RT_{\rm b}}{1000L_{\rm vap}/T_{\rm b}} = \frac{RT_{\rm b}}{s}$$

= $\frac{0.002 \times 373}{1.448} = 0.52 \text{ K kg mol}^{-1}$.

$$\Sigma n = n_{\text{glucose}} + n_{\text{urea}} = \frac{90}{180} + \frac{45}{60} = 1.25$$
; $m = 1.25 \text{ mol kg}^{-1}$.

$$\Delta T_{\rm b} = K_{\rm b} \cdot m = 0.52 \text{ K kg mol}^{-1} \times 1.25 \text{ mol kg}^{-1} = 0.65 \text{ K}.$$

6.
$$n = \frac{pV}{RT} = \frac{1.00 \text{ atm} \times (3 \times 10^{-3}) \text{ L}}{0.0821 \text{ L atm K}^{-1} \text{ mol}^{-1} \times 273 \text{ K}} = 1.3385 \times 10^{-4} \text{ mol}.$$

$$N = nN_A = (1.3385 \times 10^{-4}) \text{ mol} \times (6.02 \times 10^{23}) \text{ mol}^{-1} = 8.06 \times 10^{19}.$$

$$A = 8.06 \times 10^{19} \times 0.2 \times 10^{-18} \text{ m}^2 = 16.12 \text{ m}^2$$

$$A = 8.06 \times 10^{19} \times 0.2 \times 10^{-18} \text{ m}^2 = 16.12 \text{ m}^2.$$
7. $Z = \frac{pV}{nRT} \implies V = Z \cdot \frac{nRT}{p} = 0.111 \text{ 93 L}.$

For negligible volume,
$$\left(p + \frac{a}{(V/n)^2}\right) \cdot \frac{V}{n} = RT$$
.

Upon proper substitution, we have a = 1.253 atm L² mol⁻².

8. The first Lyman line for He⁺

$$\overline{v} = R_{\infty} Z^{2} \left(\frac{1}{m^{2}} - \frac{1}{n^{2}} \right) = R_{\infty} \cdot 4 \cdot \left(\frac{1}{1} - \frac{1}{4} \right) = \frac{3}{R_{\infty}}$$

The first Balmer series for He⁺

$$\bar{v} = R_{\infty} 2^2 \left(\frac{1}{2^2} - \frac{1}{3^2} \right) \text{ and } \lambda_A = \frac{1}{3} \cdot \frac{1}{R_{\infty}}, \lambda_B = \frac{9}{5} \cdot \frac{1}{R_{\infty}}$$

$$\Delta \lambda = \lambda_{\rm B} - \lambda_{\rm A} = \left(\frac{9}{5} - \frac{1}{3}\right) \frac{1}{R_{\infty}}$$

$$\therefore 133.7 \times 10^{-9} \text{ m} = \frac{22}{15} \cdot \frac{1}{R_{\infty}} \implies R_{\infty} = 1.097 \times 10^{7} \text{ m}^{-1}.$$

9. From the KMnO₄ titration, amount of e reacted = $48 \text{ mL} \times 0.1 \text{ mol L}^{-1} = 4.80 \text{ mmol}$.

$$C_2O_4^{2^-} \rightleftharpoons 2CO_2 + 2e$$

Amount of oxalate reacted = $\frac{4.80}{2}$ mmol = 2.40 mmol.
From the KOH titration,
amount H⁺ in sample = 18 mL × 0.08 mol L⁻¹ = 1.44 mmol.

Hence, the fraction of oxalate protonated is $\frac{1.44}{2.40} = 0.60$.

10.
$$K_{300 \text{ K}} = \frac{1}{t_1} \ln \frac{100}{100 - 20}$$
 and $K_{310 \text{ K}} = \frac{1}{t_2} \ln \frac{100}{100 - 30}$
Given $t_1 = 1.5t_2$; $\frac{K_{300 \text{ K}}}{K_{310 \text{ K}}} = 0.417$
 $\ln \frac{K_2}{K_1} = \frac{E_a}{R} \left(\frac{1}{T_1} - \frac{1}{T_2} \right) \Rightarrow \ln 0.417 = -\frac{E_a}{R} \left(\frac{1}{T_1} - \frac{1}{T_2} \right)$
 $E_a = 67.6 \text{ kJ mol}^{-1}$
 $\ln k = \ln A - \frac{E_a}{RT} = \ln (3 \times 10^9) - \frac{67.6 \times 10^3}{8.314 \times 320}$
 $k = 0.027 \text{ s}^{-1}$
 $t = \frac{1}{0.027 \text{ s}^{-1}} \ln \frac{100}{100 - 40} = 19 \text{ s}.$

11. The cell reactions are as follows:

Anode:
$$2Hg(l) + SO_4^{2-} \rightarrow Hg_2SO_4(s); E^{\circ} = -0.62 \text{ V}$$

Cathode: $PbSO_4(s) + 2e \rightarrow Pb(s) + SO_4^{2-}; E^{\circ} = -0.36 \text{ V}$
 $2Hg(l) + PbSO_4(s) \rightarrow Pb(s) + Hg_2SO_4; E^{\circ} = -0.98 \text{ V}$
 $PbSO_4 = Pb^{2+} + SO_4^{2-} \text{ and } Hg_2SO_4 = Hg_2^{2+} + SO_4^{2-}$

Using the Nernst equation,

$$E = (E_{\text{PbSO}_4/\text{Pb}, \text{SO}_4^{2-}}^2 - E_{\text{Hg}_2\text{SO}_4/\text{Hg}, \text{SO}_4^{2-}}^2) - \frac{0.0592}{2} \log \frac{[\text{Hg}_2\text{SO}_4]}{[\text{PbSO}_4]}$$

$$-0.932 = -0.98 - \frac{0.0592}{2} \log \frac{[\text{Hg}^+]^2[\text{SO}_4^{2-}]}{[\text{Pb}^2^+][\text{SO}_4^{2-}]}$$

$$-0.932 = -0.98 - \frac{0.0592}{2} \log \frac{K_s(\text{Hg}_2\text{SO}_4)}{K_s(\text{PbSO}_4)}$$

$$= -0.98 - \frac{0.0592}{2} \log \frac{1.6 \times 10^{-8} \text{ (mol L}^{-1})^3}{K_s(\text{PbSO}_4)}$$

$$K_{\rm s}({\rm PbSO_4}) = 6.6 \times 10^{-7} \; ({\rm mol} \; {\rm L}^{-1})^2.$$

12. In the RbBr molecule, the Rb⁺ and Br⁻ ions have equal (36) number of electrons. Thus these two ions diffract (or 'reflect') X-rays in the similar way. As these ions are identical, an RbBr crystal appears to be simple cubic, whereas the other three crystals have the face-centred cubic structure.

13.
$$\Lambda = \frac{1000 \,\mathrm{K}}{c} = \frac{1000 \,\times 5.55 \,\times 10^{-8} \,\Omega^{-1} \,\mathrm{cm}^{-1}}{55.5 \,\mathrm{mol} \,\mathrm{cm}^{-3}} = 10^{-6} \,\Omega^{-1} \,\mathrm{cm}^2 \,\mathrm{mol}^{-1}.$$

$$\Lambda^{\circ} = \lambda_{\mathrm{H}^{+}}^{\circ} + \lambda_{\mathrm{OH}^{-}}^{\circ} = (350 + 200) \,\Omega^{-1} \,\mathrm{cm}^2 \,\mathrm{mol}^{-1} = 550 \,\Omega^{-1} \,\mathrm{cm}^2 \,\mathrm{mol}^{-1}.$$

$$\alpha = \frac{\Lambda}{\Lambda^{\circ}} = \frac{10^{-6}}{550} = 1.8 \times 10^{-9}.$$

$$H_2 O \rightleftharpoons H^{+} + OH^{-} \qquad K_{\mathrm{w}} = [\mathrm{H}^{+}][\mathrm{OH}]$$

$$K_{\mathrm{w}} = (1.8 \times 10^{-9} \times 55.5)^2 \,(\mathrm{mol} \,\mathrm{L}^{-1})^2 = 1.0 \times 10^{-14} \,(\mathrm{mol} \,\mathrm{L}^{-1})^2.$$

14. 1 equivalent Sn = 1 equivalent $K_2Cr_2O_7$

According to question $\frac{1000}{119/4}$ mL N = $V \times N$

$$\therefore$$
 $V = 336 \text{ mL}.$

15. We know that

$$\Delta H_{\rm diss}^{\rm o}$$
 + $\Delta H_{\rm neut}$ (strong acid and strong base) = $\Delta H_{\rm neut}$ (weak acid and strong base)

or
$$\Delta H_{\text{diss}}^{\circ} - 13700 \text{ cal} = -13385 \text{ cal}.$$

$$\therefore \qquad \Delta H_{\rm diss}^{\circ} = +315 \, {\rm cal.}$$

$$\Delta H_{\text{diss}}^{\circ}$$
 per equivalent = $\frac{315 \text{ cal}}{86} \times 100 = 366 \text{ cal}$

(HA
$$\rightleftharpoons$$
 H⁺ + A⁻ $\alpha = 14\%$ and $c = 1$ N)

16. $CH_3COOH \rightleftharpoons H^+ + CH_3COO^- \quad C_6H_5COOH \rightleftharpoons C_6H_5COO^- + H^+$

$$K = \frac{[H^{+}][CH_{3}COO^{-}]}{[CH_{3}COOH]}$$
 (1)
$$K = \frac{[H^{+}][C_{6}H_{5}COO^{-}]}{[C_{6}H_{5}COOH]}$$
 (2)
$$[CH_{3}COOH] = \frac{20}{60}$$

$$[C_{6}H_{5}COOH] = \frac{10}{122}$$

$$[H^{+}] = [CH_{3}COO^{-}]$$

$$[H^{+}] = [C_{6}H_{5}COO^{-}]$$

From (1),
$$[H^+]^2 = 1.80 \times 10^{-5} \times \frac{20}{60} \text{ (mol L}^{-1})^2$$
. (3)

From (2),
$$[H^+]^2 = 1.80 \times 10^{-6} \times \frac{10}{122} \text{ (mol L}^{-1})^2$$
. (4)

From (3) and (4) we get $[H^+] = 3.46 \times 10^{-3} \text{ mol L}^{-1}$. \therefore pH = 2.46.

17.
$$T = \frac{2\pi r}{v}$$
 or $T \propto \frac{r}{v}$

Now, $r \propto n^2$ and $v \propto \frac{1}{n}$.

$$\therefore T \propto n^3 \implies T_1 = 8T_2 \implies n_1 = 2n_2.$$

18. The de Broglie wavelength is given by

$$\lambda = \frac{h}{(2m_e eV)^{1/2}} \cdot \lambda^2 = \frac{h^2}{2m_e eV} \cdot$$

Then the electron must be accelerated through the potential difference

$$V = \frac{h^2}{2\lambda^2 m_e e} = \frac{(6.626 \times 10^{-34} \text{ J s})^2}{2(550 \text{ nm})^2 (9.109 \times 10^{-31} \text{ kg}) (1.602 \times 10^{-19} \text{ C})}$$

$$= \frac{43.903 876 \times 10^{-68} \text{ J}^2 \text{ s}^2}{2(550 \times 10^{-9} \text{ m})^2 (14.592 618 \times 10^{-50} \text{ kg C})}$$

$$= \frac{43.903 876 \times 10^{-18} (\text{kg m}^2 \text{ s}^{-2})^2 \text{ s}^2}{2(302 500 \times 10^{-18} \text{ m}^2) (14.592 618 \text{ kg C})}$$

$$= 0.000 004 97 \text{ kg m}^2 \text{ s}^{-2} \text{ C}^{-1}$$

$$= 4.97 \times 10^{-6} \text{ J C}^{-1} \approx 5 \times 10^{-6} \text{ V}$$

$$= 5 \text{ µV}.$$

19. According to the question, the number of atoms per unit cell is Z = 8.

Using the formula
$$\rho = \frac{Z \cdot M}{N_{\text{A}} \cdot a^3}$$
, we have
$$3.515 \text{ g cm}^{-3} = \frac{8 \times 12 \text{ g mol}^{-1}}{(6.022 \times 10^{23} \text{ mol}^{-1}) \times a^3}.$$

$$\therefore$$
 $a \approx 356.6 \text{ pm}.$

20. $HAc \rightleftharpoons H^+ + Ac^-$ and $NaAc \rightarrow Na^+ + Ac^-$

$$K_{\rm a} = \frac{[{\rm H}^+][{\rm Ac}^-]}{[{\rm HAc}]} = \frac{[{\rm H}^+] \times 0.1 \text{ mol } {\rm L}^{-1}}{0.1 \text{ mol } {\rm L}^{-1}} = 1.8 \times 10^{-5} \text{ mol } {\rm L}^{-1}.$$

$$\therefore$$
 [H⁺] = 1.8 × 10⁻⁵ mol L⁻¹.

$$E_{\text{cell}} = E_{\text{cal}}^{\circ} - E_{\text{H}_2}^{\circ} = 0.337 \text{ V} - \frac{0.0592}{1} \log(1.8 \times 10^{-5}) \text{ V}$$

= $(0.337 + 0.281) \text{ V} = 0.618 \text{ V}.$

21. For Mechanism A we have

$$v = k[NOH_2][NO] \text{ and } K = \frac{[NOH_2]}{[NO][H_2]}.$$

$$\therefore \qquad [NOH_2] = \frac{v}{k[NO]} = K[NO][H_2].$$

$$\therefore \qquad v = Kk[NO]^2[H_2].$$

$$\therefore \qquad v \propto [NO]^2[H_2]. \qquad (1)$$

For Mechanism B we obtain

$$v' = k'[N_2O_2][H_2]$$
 and $K' = \frac{[N_2O_2]}{[NO]^2}$.

$$: [N_2O_2] = \frac{v'}{k'[H_2]} = K'[NO]^2.$$

$$: v' = K'k'[NO]^2[H_2].$$

$$: v' \propto [NO]^2[H_3]. \tag{2}$$

Now, from (1) and (2) it is clear that the rate of the given reaction is proportional to $[NO]^2[H_2]$ for either of the two mechanisms proposed.

22. Let U_1 and U_2 be the internal energies per mole of the monatomic gas and the diatomic gas respectively.

Thus
$$U_1 = \frac{3}{2}RT$$
; $U_2 = \frac{5}{2}RT$
or $U = \frac{U_1 + U_2}{2} = \frac{\frac{3}{2}RT + \frac{5}{2}RT}{2} = 2RT$ (Total mole = 2)
or $C_v = \frac{\partial U}{\partial T}\Big|_v = 2R$.

We know that $C_p - C_v = R$ or $C_p - 2R = R$ or $C_p = 3R$.

$$\therefore \quad \gamma = \frac{C_p}{C_v} = \frac{3R}{2R} = \frac{3}{2} = 1.5.$$

- **23.** As the activity of the sample reduces from 6000 Bq to 3000 Bq in 140 days, it is clear that the half-life of the sample is $t_{1/2} = 140$ days.
 - \therefore in $2t_{1/2} = 280$ days, the activity becomes one-fourth of the initial activity.
 - :. the initial activity of the radioactive sample equals

$$4 \times 6000 \text{ Bq} = 24\ 000 \text{ Bq}.$$

24. Since the kinetic energy of the proton is E and its de Broglie wavelength is λ_1 , we have

$$\lambda_1 = \frac{h}{(2m_p eV)^{1/2}} = \frac{h}{(2m_p E)^{1/2}}.$$
 (1)

Again, the wavelength of the photon having an energy equal to E is given by

$$\lambda_2 = \frac{hc}{E} \,. \tag{2}$$

Now, from (1) and (2) we find that

$$\frac{\lambda_1}{\lambda_2} = \frac{\frac{h}{(2m_{\rm p}E)^{1/2}}}{\frac{hc}{E}} = \frac{E}{\sqrt{2m_{\rm p} \cdot cE^{1/2}}} = \frac{E^{1/2}}{\sqrt{2m_{\rm p} \cdot c}}.$$

As the mass of a proton (m_p) and the speed of light in vacuum (c) are fundamental constants, we conclude that

$$\frac{\lambda_1}{\lambda_2} \propto E^{1/2}$$
.

25. (i) The galvanic cell is

Pt |
$$H_2(g, 1 \text{ atm}) | NaOH(1 \text{ M}) | O_2(g, 1 \text{ atm}) | Pt$$

LHE (ox):
$$H_2 + 2OH^- \rightarrow 2H_2O + 2e$$

$$\mbox{RHE (red):} \quad \frac{1}{2}\,\mbox{O}_2 + \mbox{H}_2\mbox{O} + 2\mbox{e} \rightarrow \mbox{ 2(OH}^-) \label{eq:condition}$$

Net reaction:
$$H_2(g) + \frac{1}{2}O_2(g) \rightarrow H_2O(l)$$

(ii) Pt
$$\mid$$
 H₂(g, 1 atm) \mid H⁺ \mid OH⁻, O₂(g, 1 atm) \mid Pt

LHE (ox):
$$H_2(g) \rightarrow 2H^+ + 2e$$

$$\label{eq:RHE} \text{RHE (red):} \quad \frac{1}{2}\,O_2 + H_2O + 2e \rightarrow \ 2(OH^-)$$

Net reaction:
$$H_2 + \frac{1}{2}O_2 + H_2O \rightarrow 2H^+ + 2(OH^-)$$

For the reaction $H_2O(1) \rightleftharpoons H^+ + OH^-$,

$$\Delta G^{\circ} = -nFE^{\circ} = -(1) (96\ 500\ \text{C mol}^{-1})(-0.6\ \text{V}) = 57.9\ \text{kJ mol}^{-1}.$$

For the reaction $H_2 + \frac{1}{2}O_2 + 2H_2O \rightarrow H_2O + 2H^+ + 2OH^-$,

$$\Delta G_r^{\circ} = \Delta_{\text{formation}} G_{\text{H}_2\text{O}}^{\circ} + 2\Delta_{\text{ions}} G_{\text{H}_2\text{O}}^{\circ}$$

$$= 57.9 \times 2 - 240 = -124.2 \text{ kJ}$$

$$E_{\text{cell}}^{\circ} = -\frac{\Delta G^{\circ}}{nF} = \frac{124.2 \times 10^{3}}{2 \times 96500} = 0.643 \text{ V}.$$

Thus,
$$E_{\text{cell}}^{\circ} = E_{\text{red}}^{\circ} \mid \text{OH}^{-} \mid \text{O}_{2}$$
, $\text{Pt} - E_{\text{H}^{+} \mid \text{H}_{2}}^{\circ} = E_{\text{cell}}^{\circ} \mid \text{OH}^{-} \mid \text{O}_{2} \mid \text{Pt} = 0.643 \text{ V}$.

Equation:	Hg(CN) ₂	+	nCN ⁻	$\longrightarrow [Hg(CN)_{n+2}]^{n-}$
Initial amounts:	0.095 mol		0.1892 mol	0 mol
Changes to reach equilibrium:	-0.095 mol		-0.095 <i>n</i> mol	+0.095 mol
Amounts at equilibrium:	0 mol	(0.	.1892–0.095 <i>n</i>) n	nol 0.095 mol

26. The equilibrium composition of the given reaction is given below.

Hence, the total molality obtained after the addition of Hg(CN)₂ is

$$m_{\text{total}} = m(\text{K}^+) + m(\text{CN}^-) + m([\text{Hg}(\text{CN})_{n+2}]^{n-})$$

= 0.1892 mol kg⁻¹ + (0.1892 – 0.095n) mol kg⁻¹ + 0.095 mol kg⁻¹
= (0.4734 – 0.095n) mol kg⁻¹.

Since the freezing point of pure water is 0°C and that of the aqueous solution of KCN is -0.704°C, the depression of freezing point of water on adding KCN is $\Delta T_{\rm f.KCN} = 0$ °C - (-0.704°C) = 0.704°C.

Also, the van't Hoff factor for KCN is $i \approx 2$ as the solute dissociates almost completely.

: the freezing-point constant is given by

$$K_{\rm f} = \frac{\Delta T_{\rm f, KCN}}{m({\rm KCN}) \cdot i} = \frac{0.704}{0.1892 \times 2} \,{\rm K \ kg \ mol}^{-1} \approx 1.86 \,{\rm K \ kg \ mol}^{-1}.$$

Now, the depression of freezing point of water on adding $Hg(CN)_2$ is given by

$$\Delta T_{\rm f, Hg(CN)_2} = K_{\rm f} \cdot m_{\rm total}$$

∴
$$0^{\circ}\text{C} - (-0.53^{\circ}\text{C}) = (1.86 \text{ K kg mol}^{-1}) \cdot (0.4734 - 0.095n) \text{ mol kg}^{-1}$$

⇒ $0.53 \text{ K} = (0.880524 - 0.1767n) \text{ K}$

⇒ $0.1767n = 0.880524 - 0.53 = 0.350524$.

∴ $n = \frac{0.350524}{0.1767} = 1.9837... \approx 2$.

Therefore the formula of the complex formed is found to be $[Hg(CN)_4]^{2-}$.

27. The ratio of the quantity of bromoauric acid transformed into chloroauric acid to the total quantity of bromoauric acid employed is equal to the ratio of the observed heat of evolution to the molar heat of the reaction

$$HAuBr_4 + 4HCl \rightarrow HAuCl_4 + 4HBr$$
.

The molar heat of this reaction is obtained as follows by subtracting equation (2) from equation (1), giving equation (3):

$$Au(OH)_3 + 4HCl = HAuCl_4 + 3H_2O; \Delta H^{\circ} = -23.00 \text{ kcal}$$
 (1)

$$Au(OH)_3 + 4HBr = HAuBr_4 + 3H_2O; \Delta H^{\circ} = -36.8 \text{ kcal}$$
 (2)

$$HAuBr_4 + 4HCl = HAuCl_4 + 4HBr; \Delta H^{\circ} = +13.8 \text{ kcal}$$
 (3)

Let x% bromoauric acid be transformed into chloroauric acid.

$$\therefore \quad \frac{x}{100} = \frac{0.51 \text{ kcal}}{1.38 \text{ kcal}} \quad \Rightarrow \quad x \approx 3.7.$$

28. The emf of concentration cell is

$$E = -\frac{0.0592}{n} \log \frac{a_{\rm L}}{a_{\rm R}}$$
 or
$$0.0579 = -\frac{0.0592}{1} \log \frac{0.001\alpha_1}{0.01\alpha_2} \cdot$$
 or
$$0.0579 = -\frac{0.0592}{1} \log \frac{0.001 \times 1}{0.01\alpha_2}$$

$$\alpha_2 = 0.96 = 96\%.$$

$$[Ag^+] = 0.01 \text{ mol } L^{-1} \times \alpha_2 = 9.6 \times 10^{-3} \text{ mol } L^{-1}.$$

29. Use the ideal gas equation, i.e. pV = nRT. (1)

Differentiating both sides w.r.t. T at constant pressure, we get

$$p\frac{dV}{dT} = nR. (2)$$

Dividing (2) by (1),

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$$\frac{1}{V}\frac{dV}{dT} = \frac{1}{T}$$
 or $\delta = \frac{1}{T}$.

30. The dissociation of H_2SO_4 takes place in the following steps.

Step I:
$$H_2SO_4 \xrightarrow{K_1} H^+ + HSO_4^-$$

Step II: $HSO_4^- \xrightarrow{K_2} H^+ + SO_4^{2-}$

Now, consider the equilibrium composition of Step II as shown below.

Equation:	HSO ₄ —	\rightarrow H ⁺	+ SO ₄ ²⁻
Initial amounts:	0.1 mol	0 mol	0 mol
Changes to reach equilibrium	-x mol	+x mol	+x mol
Amounts at equilibrium:	(0.1 - x) mol	x mol	x mol

Hence, we have

$$K_{2} = \frac{[H^{+}][SO_{4}^{2-}]}{[HSO_{4}^{-}]} = \frac{(x \text{ mol } L^{-1})(x \text{ mol } L^{-1})}{(0.1 - x) \text{ mol } L^{-1}}.$$

$$\vdots \qquad 10^{-2} \text{ mol } L^{-1} = \frac{x^{2} (\text{mol } L^{-1})^{2}}{(0.1 - x) \text{ mol } L^{-1}}$$

$$\Rightarrow \qquad x^{2} = 10^{-2}(0.1 - x) = 0.001 - 0.01x$$

$$\Rightarrow \qquad x^{2} + 0.01x - 0.001 = 0.$$

$$\vdots \qquad (x - 0.027)(x + 0.037) \approx 0.$$

Since *x* cannot be negative, $x \approx 0.027$.

$$(H^{+}]_{total} = [H^{+}]_{H_{2}SO_{4}} + [H^{+}]_{HSO_{4}^{-}}$$

$$= (0.1 + x) \text{ mol } L^{-1}$$

$$= (0.1 + 0.027) \text{ mol } L^{-1} = 0.127 \text{ mol } L^{-1}.$$

$$\therefore \qquad pH = -\log 0.127 = -(-0.896 \text{ } 19...) \approx 0.9.$$

31. The cell is $Zn(s) \mid Zn^{2+}(1 \text{ M}) \parallel Cu^{2+}(1 \text{ M}) \mid Cu(s) + NH_3(excess)$

$$Cu^{2+} + 4NH_3 \implies Cu(NH_3)_4^{2+} \qquad K_f = \frac{[Cu(NH_3)_4^{2+}]}{[NH_3]^4[Cu^{2+}]}$$
Initially, 1.0 M excess 0
At equilibrium, $(1-x)$ 2.0 M x 1.0 × 10⁻¹² = $\frac{x}{2^4(1-x)}$

$$\frac{x}{1-x} = 16 \times 10^{-12} \implies \frac{1-x}{x} = \frac{1}{16 \times 10^{-12}} \implies \frac{1}{x} - 1 = \frac{1}{16 \times 10^{-12}}.$$

$$\frac{1}{x} = \frac{1}{16 \times 10^{-12}} \implies x = 16 \times 10^{-12}.$$

$$E_{cell} = (E_R^{\circ} - E_L^{\circ}) - \frac{0.0592}{2} \log \frac{[Zn^{2+}]}{[Cu^{2+}]} V$$

$$= 1.10 \text{ V} - 0.0296 \log \frac{1.0}{16 \times 10^{-12}} V$$

$$= 1.10 \text{ V} - 0.0296(-\log 2^4 + 12) \text{ V}.$$

$$\therefore E_{cell} = 0.70 \text{ V}.$$

32. Since the crystal is fcc, the edge length of the unit cell is $a = 2 \times$ shortest distance between A and $B = 2Y^{1/3}$ nm.

As we know,
$$\rho = \frac{Z \cdot M}{N_{\text{A}} \cdot a^3} = \frac{4 \times 6.022 \text{ y u mol}^{-1}}{(6.022 \times 10^{23} \text{ mol}^{-1})(2 \text{ y}^{1/3} \times 10^{-9} \text{ m})^3}$$
$$= 0.5 \times 10^4 \text{ u m}^{-3} = 0.83 \times 10^{-23} \text{ kg m}^{-3}.$$

33.
$$E_{\text{cell}} = \frac{0.0592}{1} \log \frac{K_{\text{s}}(\text{AgBr/Br}^{-})}{K_{\text{s}}(\text{AgCl/Cl}^{-})} \text{ V}$$
$$= \frac{0.0592}{1} \log \frac{3.3 \times 10^{-13}/0.02}{2.8 \times 10^{-10}/0.2} \text{ V}$$

Since $E_{\rm cell}$ is negative, the reverse reaction is spontaneous.

34.
$$\lambda_1 = 108.5 \text{ nm} = 108.5 \times 10^{-9} \text{ m}$$

 $\lambda_2 = 30.4 \text{ nm} = 30.4 \times 10^{-9} \text{ m}$

Let the excited state be n_2 . The electron falls first from n_2 to n_1 and then from n_1 to the ground state.

$$\frac{1}{\lambda_2} = Z^2 R_{\infty} \left(\frac{1}{1^2} - \frac{1}{n_1^2} \right)$$
or
$$\frac{1}{30.4 \times 10^{-9} \text{ m}} = 2^2 \times (1.097 \times 10^7 \text{ m}^{-1}) \times \left(\frac{1}{1^2} - \frac{1}{n_1^2} \right)$$
or
$$n_1 = 2.$$
Again,
$$\frac{1}{\lambda_1} = Z^2 R_{\infty} \left(\frac{1}{2^2} - \frac{1}{n_2^2} \right)$$
or
$$\frac{1}{108.5 \times 10^{-9} \text{ m}} = 2^2 \times (1.097 \times 10^7 \text{ m}^{-1}) \times \left(\frac{1}{2^2} - \frac{1}{n_2^2} \right)$$
or
$$n_2 = 5.$$

35. We know that $\Delta U = \frac{p\Delta V}{v-1}$.

At constant pressure,
$$dw = p\Delta V$$
. (1)

From (1) and (2),

$$\Delta U = \frac{dw}{\gamma - 1}$$
 or $\Delta U = dq - dw$ or $dq = \Delta U + dw = \frac{dw}{\gamma - 1} + 1$.

$$dq = \frac{\gamma}{\gamma - 1} dw = \frac{1.4}{1.4 - 1} \times 2 J = \frac{2.8}{0.4} J = 7 J.$$

36. Let the order of the rate-determining step in the pyrolysis of $B_2H_6(g)$ be n. Then the rate law for the reaction is

$$v = k[B_2H_6]^n.$$

Hence, from the given data we obtain

$$7.40 \times 10^4 \text{ mol L}^{-1} \text{ h}^{-1} = k(2.153 \times 10^2 \text{ mol L}^{-1})^n$$
 (1)

and

$$0.73 \times 10^4 \text{ mol L}^{-1} \text{ h}^{-1} = k(0.433 \times 10^2 \text{ mol L}^{-1})^n$$
 (2)

Now, dividing (1) by (2),

$$\frac{7.40}{0.73} = \left(\frac{2.153}{0.433}\right)^n$$

$$\therefore \qquad \log \frac{7.40}{0.73} = \log \left(\frac{2.153}{0.433}\right)^n$$

$$\Rightarrow \qquad \log 10.137 = n \log 4.972.$$

$$\therefore \qquad n = \frac{\log 10.137}{\log 4.972} \approx \frac{1.006}{0.696}$$

$$\approx 1.4454... \approx 1.5 = \frac{3}{2}.$$

37. The rate law of the given reaction is

$$\operatorname{rate} = -\frac{1}{2} \frac{d[N_2O_5]}{dt} = \frac{1}{4} \times \frac{d[NO_2]}{dt} = \frac{d[O_2]}{dt} = k[N_2O_5].$$

$$\frac{d[N_2O_5]}{dt} = 2k[N_2O_5] = k_1[N_2O_5] \quad \text{or} \quad 2k = k_1 \quad \text{or} \quad 4k = 2k_1.$$

$$\frac{d[NO_2]}{dt} = 4k[N_2O_5] = k_2[N_2O_5] \quad \text{or} \quad 4k = k_2 \quad \text{or} \quad 4k = k_2.$$

$$\frac{d[O_2]}{dt} = k[N_2O_5] = k_3[N_2O_5] \quad \text{or} \quad k = k_3 \quad \text{or} \quad 4k = 4k_3.$$

The relation is $2k_1 = k_2 = 4k_3$.

38. Work done by the system $(w) = -nRT \ln \frac{p_1}{p_2}$. $w = -2.303 \ nRT \log \frac{p_1}{p_2} = -2.303 \times 5 \times 8.314 \times 300 \log \frac{10}{2}$ $= -2.0075 \times 10^4 \text{ J}.$

Let *m* be the mass which can be lifted through a height of 1 m.

$$w = mgh$$
 or 2.0075 × 10 ⁴ J = m × 9.8 × 1 J
∴ $m = 205$ kg (approx.)

39. Rate =
$$v = K\theta = \frac{kKp}{1 + Kp}$$

- (a) Adsorption on gold is $\theta \approx 1$, and v = K = constant. Hence, it is a zero-order reaction.
- (b) Adsorption on platinum is $\theta \approx K_p$ as $K_p < 1$. So v = kKp, and the reaction is of the first order.
- 40. The formula weight of KCN is equal to

$$M_r = 39.1 + 12 + 14 = 65.1.$$

: the molar mass is given by

$$M = 65.1 \text{ g mol}^{-1}$$
.

- : molar volume, $V_{\rm m} = \frac{M}{\rho} = \frac{65.1 \text{ g mol}^{-1}}{1.52 \text{ g cm}^{-3}} \approx 42.8289 \text{ cm}^3 \text{ mol}^{-1}.$
- : the volume of a formula unit is

$$V = \frac{V_{\rm m}}{N_{\rm A}} = \frac{42.8289~{\rm cm^3~mol^{-1}}}{6.022 \times 10^{23}~{\rm mol^{-1}}}$$

$$\approx 7.112~07 \times 10^{-23}~{\rm cm^3}$$

$$= 71.1207 \times 10^{-24}~{\rm cm^3}.$$

Since the rock-salt structure is a face-centred cubic crystal containing four formula units, the volume of the unit cell equals

$$a^3 = 4V = 4(71.1207 \times 10^{-24} \text{ cm}^3)$$

= 284.4828 × 10⁻²⁴ cm³.

$$a = (284.4828 \times 10^{-24} \text{ cm}^3)^{1/3}$$

$$= (284.4828)^{1/3} \times 10^{-8} \text{ cm}$$

$$= (284.4828)^{1/3} \times 10^{-10} \text{ m}$$

$$= (284.4828)^{1/3} \text{ Å}.$$

$$\therefore \qquad d_{100} = \frac{a}{(1^2 + 0^2 + 0^2)^{1/2}} = (284.4828)^{1/3} \text{ Å}.$$

$$\therefore \qquad \ln d_{100} = \frac{1}{3} (\ln 284.4828) \text{ Å} \approx \frac{5.6507}{3} \text{ Å} \approx 1.883 \text{ Å}.$$

$$\therefore \qquad d_{100} = (\exp 1.883) \text{ Å} \approx 6.57 \text{ Å}.$$

Part 2

Inorganic Chemistry

1

Periodic Table

• Type 1 •

1. The correct order of increasing radius of the elements Si, Al, Na and P is

(b) P, Si, Al, Na

Choose the correct option. Only one option is correct.

(a) Si, Al, P, Na

	(c)	Al, Si, P, Na			(d)	Al, P, Si,	Na		
2.	Which radii?	n of the follo	wing	series of	elemen	ts have n	early th	e same a	atomi
	(a)	F, Cl, Br, I			(b)	Na, K, R	b, Cs		
	(c)	Li, Be, B, C			(d)	Fe, Co, N	Ji, Cu		
3.	In wh	ich of the foll	owir	ng pairs is	the seco	ond atom	larger th	nan the f	irst?
	(a)	Br, Cl			(b)	Na, Mg			
	(c)	Sr, Ca			(d)	N, P			
4.		n of the follow racter?	wing	compour	nds may	one expe	ect to be	strongly	y ionio
	(a)	$BaCl_2$	(b)	CsCl	(c)	$CaCl_2$	(d)	$SrCl_2$	
5.		rder of increa s ² 2s ² 2p ⁶ 3s ²					tronic co	onfigura	tions
				(iv) 1					
	is	1		(/		1			
	(a)	(ii), (iv), (iii)	, (i)		(b)	(i), (ii), (i	ii), (iv)		
	(c)	(i), (iii), (ii),	(iv)		(d)	(iv), (iii),	(ii), (i)		
6.	The h	ighest ionizat	ion e	energy is e	exhibited	d by			
	(a)	halogens			(b)	alkaline	earth me	etals	
	(c)	transition m	etals		(d)	noble ga	ses		
				2	-3				

element with the maximum electron affinity?

other as a liquid under normal conditions?

(a) $1s^2 2s^2 2p^6$

(a) I_2 and F_2

(c) $1s^2 2s^2 2p^6 3s^1$

7. Which among the following electronic configurations represents the

8. In which of the following pairs does one element exist as a solid and the

(b) $1s^2 2s^2 2p^5$

(b) I₂ and Br₂

(d) $1s^2 2s^2 2p^6 3s^2 3p^5$

	(c)	Br ₂ and Hg	(d)	Na and Cs
9.	Amor	ng the following, the element w	hich	has the configuration
	(a)	[Ne] $3s^{1} 3p^{2}$	(b)	[Ne] $3s^2 3p^5$
	(c)	[Ne] $3s^2 3p^4$	(d)	[Ne] $3s^2 3p^6 3d^5 4s^1$
	has th	e highest electron affinity.		
10.		n of the following sets of atom oup 16?	ic n	umbers corresponds to elements
	(a)	8, 16, 32, 54	(b)	16, 34, 54, 86
	(c)	8, 16, 34, 52	(d)	10, 16, 32, 50
11.		ng the following electronic conf ement with the highest ionizati		ations, the one corresponding to nergy is
	(a)	[Ne] $3s^2 3p^1$	(b)	$[Ar] 3d^{10} 4s^2 4p^2$
	(c)	[Ne] $3s^2 3p^2$	(d)	[Ne] $3s^2 3p^3$
12.	Of will	9	ts is	the first ionization energy the
	(a)	Lead (b) Carbon	(c)	Silicon (d) Tin
13.	Amor	ng the following elements, whi		as the highest ionization energy?
		Nitrogen	. ,	Chromium
	(c)	Neon	(d)	Manganese
14.	The the or		CO ₃ ,	SrCO ₃ and MgCO ₃ decreases in
	(a)	$BaCO_3 > SrCO_3 > MgCO_3 > C$	aCO	3
	(b)	$CaCO_3 > SrCO_3 > MgCO_3 > B$	aCO	3
	(c)	$MgCO_3 > CaCO_3 > SrCO_3 > B$	aCO	3
	(d)	$BaCO_3 > SrCO_3 > CaCO_3 > M_3$	gCO	3
15.	Whicl	n of the following has the maxi	mun	n number of unpaired electrons?
	(a)	Ti ³⁺	(b)	V^{3+}
	(c)	Fe ³⁺	(d)	Fe ²⁺

16.		aufbau princolaced in the	ciple had not be	en fo	ollowed, Ca(2	Z=20) would have
	(a)	s-block		(b)	p-block	
	(c)	d-block		(d)	f-block	
17.		ng the equation	· .	the o	one whose Δl	H° value equals the
	(a)	$Ca^+(g) \longrightarrow$	$Ca^{2+}(g) + e$	(b)	$Ca(g) \longrightarrow$	$Ca^+(g) + e$
	(c)	$Ca(s) \longrightarrow$	$Ca^+(g) + e$	(d)	$Ca(g) \longrightarrow$	$Ca^{2+}(g) + 2e$
18.	Which	n of the follow	ving oxides is neu	ıtral?	•	
	(a)	SnO_2		(b)	CO	
	(c)	Al_2O_3		(d)	Na ₂ O	
19.	Which	n of the follow	ving is the least st	able	?	
	(a)	PbI_4	(b) KCl	(c)	GeI_4	(d) SnI_4
20.	Amor has a the	ng the following ground-state	ing, the element electronic config	with gurat	the lowest a ion of $(n-1)$	tomic number that d^6ns^2 is located in
	(a)	fifth period		(b)	sixth period	l
	(c)	fourth perio	d	(d)	third period	l
21.	Which	n of the follow	ving properties d	oes n	ot depend or	n periodicity?
		Atomic weig		` /	Atomic nun	
	` '	Ionization e	07		Electronega	•
22.			ty of the followin			
	(a)	C < N < Si <	P	` /	Si < P < C <	
	(c)	N < C < P < S	Si	(d)	C < Si < N <	:P
23.	Which	n of the follow	ving is arranged i	n ord	der of increas	ing radius?
	(a)	$K^+(aq) < Na$	$^{+}(aq) < Li^{+}(aq)$			
	(b)	$K^+(aq) > Na$	$^{+}(aq) > Zn^{2+}(aq)$			
	(c)	$K^+(aq) < Li^+$	$T(aq) < Na^+(aq)$			
	(d)	$Li^+(aq) < Na$	$a^+(aq) < K^+(aq)$			
24.	The th	nird ionization	n energy is maxir	num	for	
	(a)	nitrogen		(b)	phosphorus	3
	(c)	aluminium		(d)	boron	
25.			_			llowing elements is
	` '	S < O < Se <			O < C < S <	
	(C)	O < S < Se <	C	(a)	C < O < S <	Se

- **26.** Which of the following arrangements shows the correct order of decreasing paramagnetism?
 - (a) N > Al > O > Ca
- (b) N > O > Al > Ca
- (c) O > N > Al > Ca
- (d) O > N > Ca > Al
- **27.** A large difference between the fourth and fifth ionization energies indicates the presence of
 - (a) 5 valence electrons in an atom
 - (b) 6 valence electrons in an atom
 - (c) 4 valence electrons in an atom
 - (d) 8 valence electrons in an atom
- **28.** Which of the following has been arranged in order of increasing covalent character?
 - (a) $BaCl_2 < AlF_3 < BeCl_2 < LiCl$
 - (b) KCl < NaCl < BeCl₂ < PbCl₄
 - (c) $NH_4Cl < BCl_3 < CCl_4 < AlF_3$
 - (d) KCl < CaCl₂ < AlCl₃ < SiCl₄
- 29. Electron affinity is defined as the
 - energy required to remove an electron from an isolated gaseous atom
 - (b) energy released when an electron is added to an isolated atom in the gaseous state
 - (c) energy required to add an electron to an isolated atom in the gaseous state
 - (d) ability of an atom to attract an electron
- 30. Which of the following halides is stable to hydrolysis?
 - (a) SiCl₄
- (b) PF₃
- (c) NCl₃
- (d) NF₃
- **31.** Which of the following is arranged in order of increasing thermal stability?
 - (a) $HgCl_2 < BeCl_2 < CaCl_2 < BaCO_3$
 - (b) $BaCl_2 < CaCl_2 < BeCl_2 < HgCl_2$
 - (c) $CaCl_2 < BeCl_2 < BaCO_3 < HgCl_2$
 - (d) $BeCl_2 < BaCl_2 < HgCl_2 < CaCl_2$
- **32.** The correct order of increasing electron affinity of the following elements is
 - (a) C < Be < N < O < F
- (b) Be < C < O < N < F
- (c) Be < C < N < O < F
- (d) Be < C < N < F < O

33. Which of the following has been arranged in order of the increasing

(b) $Pb^{2+} < Ge^{2+} < Sn^{2+}$

(d) $Cu^{2+} < Au^{2+} < Ag^{2+}$

stability of the +2 oxidation state of the ions?

(a) $Ca^{2+} < Ba^{2+} < Sr^{2+}$

(c) $Ge^{2+} < Sn^{2+} < Pb^{2+}$

34.	(a)	h of the following is arranged i Al < Mg < C (graphite) < B	n ord	der of increasing density?
		B < Al < Mg < C (graphite)		
		C (graphite) $<$ Al $<$ B $<$ Mg		
		Mg < C (graphite) < B < Al		
35.	config	guration of Group 13 elements	?	e correct outer-shell electronic
	(a)	$ns^2 nd^1$	(b)	$ns^2 np^6$
	(c)	$ns^2 np^1$	(d)	$ns^2 np^3$
36.	Amor	ng the following, which has the	e low	est enthalpy of fusion?
	(a)	Fluorine	(b)	Hydrogen
	(c)	Chlorine	(d)	Helium
37.	Which	h of the following is arranged i	n or	der of increasing density?
		Ag < Au < Fe < Cu		Cu < Au < Ag < Fe
	(c)	Fe < Cu < Ag < Au	(d)	Au < Ag < Cu < Fe
38.	What Group	0	of t	he outer shell of the elements of
	(a)	$ns^2 np^4$ (b) $ns^2 np^6$	(c)	$ns^2 np^2$ (d) ns^2
39.	Which	h of the following is arranged i	n or	der of increasing boiling point?
	(a)	$H_2O < CCl_4 < CS_2 < CO_2$	(b)	$CO_2 < CS_2 < CCl_4 < H_2O$
	(c)	$CS_2 < H_2O < CO_2 < CCl_4$	(d)	$CCl_4 < H_2O < CO_2 < CS_2$
40.	Which	h of the following is arranged i	n or	der of increasing density?
	(a)	$Ne < Cl_2 < N_2 < O_2$	(b)	$N_2 < O_2 < Ne < Cl_2$
	(c)	$Cl_2 < Ne < O_2 < N_2$	(d)	$Cl_2 < N_2 < O_2 < Ne$
41.	Which	h of the following is arranged i	n de	creasing order of size?
	(a)	$Mg^{2+} > Al^{3+} > O^{2-}$	(b)	$O^{2-} > Mg^{2+} > Al^{3+}$
	(c)	$Al^{3+} > Mg^{2+} > O^{2-}$	(d)	$Mg^{2+} \approx Al^{3+} > O^{2-}$
42.	Which	h of the following compounds	has a	a positive enthalpy of solution?
	(a)	LiF	(b)	LiCl
	(c)	LiBr	(d)	LiI

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43.		h of the following ation energy?	is arranged in order of increasing second
	(a)	C < N < O < F	(b) $F < C < N < O$
	(c)	C < N < F < O	(d) F < O < N < C
44.	The o	rder of increasing ion	ic radius of the following is
	(a)	$K^+ < Li^+ < Mg^{2+} < A$	1^{3+}
	(b)	$K^+ < Mg^{2+} < Li^+ < A$	1 ³⁺
	(c)	$Li^+ < K^+ < Mg^{2+} < A$	l^{3+}
	(d)	$Al^{3+} < Li^+ < Mg^{2+} < 1$	K ⁺

- 45. Which of the following compounds has a negative enthalpy of solution?
 - (a) KCl

(b) KBr

(c) KF

- (d) KI
- **46.** Which of the following statements is incorrect?
 - (a) The second ionization energy of sulphur is greater than that of chlorine.
 - (b) The third ionization energy of phosphorus is greater than that of aluminium.
 - (c) The first ionization energy of aluminium is approximately the same as that of gallium.
 - (d) The second ionization energy of boron is greater than that of carbon.
- 47. Which of the following electronic configurations corresponds to the element with the highest electron affinity?
 - (a) $[Ar] 3d^{10} 4s^2 4p^3$
- (b) $[Ar] 3d^{10} 4s^2 4p^4$
- (c) $[Ar] 3d^{10} 4s^2 4p^5$
- (d) [Ar] $3d^{10} 4s^2 4p^6$
- 48. The ionization energies of which of the following pairs of elements increase in a regular and integral manner?
 - (a) Nitrogen and phosphorus
- (b) Oxygen and fluorine
- (c) Boron and aluminium
- (d) Sodium and magnesium
- 49. Among the following elements, the second ionization energy is maximum for
 - (a) boron

(b) beryllium

(c) magnesium

- (d) aluminium
- **50.** Arrange N, O and S in order of decreasing electron affinity.
 - (a) S > O > N

(b) O > S > N

(c) N > O > S

(d) S > N > O

51. Arrange the elements with the following electronic configurations in

increasing order of electron affinity.

than that of oxygen.

	(i) 1	$s^2 2s^2 2p^5$	(ii)	$1s^2 2s^2 2p$	2 4
	(iii) 1	$s^2 2s^2 2p^3$	(iv)	1s ² 2s ² 2p	p ⁶ 3s ² 3p ⁴
	(a)	i < ii < iii < iv		(b)) iv < iii < ii < i
	(c)	iii < ii < iv < i		(d)) ii < iii < i < iv
52.	Which soluti		pair	s of compo	ounds has positive enthalpies of
	(a)	NaF and NaCl		(b)) NaBr and NaI
	(c)	NH ₄ Cl and NH ₄ F		(d)) CaF ₂ and CaCl ₂
53.	Amor	ng the following elem	men	ts, which h	has the least electron affinity?
	(a)	Phosphorus		(b)) Sulphur
	(c)	Oxygen		(d)) Nitrogen
54.	The b	pasicity of the hydr	oxid	les of the	following alkali metals is of the
	(a)	Li > Na > Rb > Cs		(b)) $Na > Li > Rb > Cs$
	(c)	Cs > Rb > Na > Li		(d)) $Rb > Cs > Na > Li$
55.		pasic strengths of the strengths of the strengths of the strengths are strengths.	he ł	nydroxides	s of the following alkaline earth
		Be > Mg > Sr > Ba) Mg > Be > Ba > Sr
	(c)	Ba > Sr > Mg > Be		(d)) Sr > Be > Mg > Ba
56.	The so	olubilities of Na ₂ SO	₄ , Be	SO ₄ , MgSC	${\sf O}_{\!\scriptscriptstyle 4}$ and ${\sf BaSO}_{\!\scriptscriptstyle 4}$ follow the order
	(a)	$BeSO_4 > MgSO_4 > 1$	Na ₂ S	$SO_4 > BaSC$	O_4
	(b)	$Na_2SO_4 > BeSO_4 >$	Mgs	$SO_4 > BaSC$	O_4
	(c)	$BeSO_4 > Na_2SO_4 >$	Mgs	$SO_4 > BaSC$	\mathcal{O}_4
	(d)	$MgSO_4 > BeSO_4 > 1$	Na ₂ S	$SO_4 > BaSC$	\mathcal{O}_4
57.	The b	asic character of Mg	gO, S	SrO, K ₂ O at	nd NiO increases in the order
	(a)	$K_2O < SrO < MgO$	< N	iO (b)	$) NiO < MgO < SrO < K_2O$
	(c)	MgO < NiO < SrO	< K	$_{2}O$ (d)) K ₂ O < MgO < NiO < SrO
58.	Which affinit		is	arranged	in order of increasing electron
	(a)	C < O < N < B		(b)) N < O < B < C
	(c)	O < C < B < N		(d)) N < B < C < O
59.		n of the following st The magnitude of			rrect? ctron affinity of sulphur is greater

than that of oxygen.

chlorine

approximately the same.

(b) The magnitude of the second electron affinity of sulphur is less

(c) The first electron affinities of bromine and iodine are

(d) The first electron affinity of fluorine is greater than that of

60. Which of the following pairs of molecules have the identical bond dissociation energies? (b) N₂ and CO (a) F_2 and H_2 (d) HF and O₂ (c) F_2 and I_2 61. Which of the following arrangements is correct in terms of electron affinity? (a) Chlorine > fluorine > iodine > bromine (b) Bromine < iodine < chlorine < fluorine (c) Bromine < iodine < fluorine < chlorine (d) Bromine ≈ iodine > chlorine < fluorine **62.** Among the following oxides, which has the maximum lattice energy? (a) MgO (b) CaO (c) SrO (d) BaO 63. Among the following molecules, which has the maximum lattice energy? (a) KBr (b) NaCl (c) NaF (d) LiF 64. Which of the following is arranged in increasing order of hydration energy? (a) $Cu^{2+} < Fe^{3+} < Fe^{2+} < A1^{3+}$ (b) $Fe^{2+} < Cu^{2+} < Al^{3+} < Fe^{3+}$ (c) $Al^{3+} < Fe^{3+} < Fe^{2+} < Cu^{2+}$ (d) $Fe^{3+} < A1^{3+} < C11^{2+} < Fe^{2+}$ 65. Among the following, which has the maximum hydration energy? (b) NH₄⁺ (c) F (d) H⁺ (a) OH⁻ **66.** Which of the following is arranged in order of increasing melting point? (a) Zn < Cu < Ni < Fe(b) Fe < Ni < Cu < Zn(c) Ni < Fe < Zn < Cu(d) Cu < Zn < Fe < Ni**67.** Which of the following is arranged in order of decreasing melting point? (b) Al > Si > P > S(a) S > P > Si > Al(c) Si > Al > S > P(d) P > S > Al > Si

Periodic Table 2-11

68.	Among the following, which has the (a) Sulphur (b) Phosphorus		-
69.	Which of the following is arranged i (a) Mg > Ca > Hg > Zn (c) Ca > Mg > Zn > Hg	n ord (b)	
70.	Which of the following pairs inclu covalent radii? (a) Hydrogen and helium	des (b)	
71.	(c) Nitrogen and oxygen Among the following metals, the mo (a) osmium (b) iridium	ost de	•
72.	Which of the following sets has elem (a) B, C, N, O (c) Cr, Mn, Fe, Cu	(b)	with identical covalent radii? Al, Si, P, S Li, Be, B, C
73.	Among the following, the third ioniz (a) aluminium (c) magnesium • Type	(b) (d)	boron beryllium
Choo	ose the correct options. More than one		
74.	Paramagnetism is exhibited by elem (a) only an odd number of electr (b) only an even number of electr (c) the d-shell partially filled (d) none of these	ons	the atoms of which have
75.	In halogens, which of the following (a) Bond length (b) Electronegativity (c) The ionization energy of the edition of the ionization power		
76.		iv)] (b)	

77.	Which among the following are isostructural pairs?			
	(a)	NF_3 and $H_3O^{\scriptscriptstyle +}$	(b)	NO ₃ and BF ₃
	(c)	NF ₃ and NO ₃	(b)	NF ₃ and NH ₃
78.	Which	n of the following statements a	re co	rrect?
	(a)	F is the most electronegative element.	e aı	nd Cs the most electropositive
	(b)	The electronegativity of halog	gens	decreases from F to I.
	(c)	The electron affinity of Cl is electronegativities are in the n		her than that of F though their se order.
	(d)	The electron affinity of noble	gase	s is almost zero.
79.	Which	n of the following statements a	re tri	ue?
	(a)	Mendeleev's periodic law is elements.	s bas	sed on the atomic numbers of
	(b)	The table presented by Mend	eleev	did not have the zero group.
	(c)	Each group in Mendeleev's subgroups.	perio	odic table was divided into two
	(d)	Mendeleev's periodic table series.	cons	sists of ten horizontal rows or
80.	Which	n of the following have a diago	nal r	relationship?
	(a)	Li and Mg	(b)	B and Mg
	(c)	Be and Al	(d)	Be and Na
81.	Which	n of the following belong to a t	riad?	
		Osmium	(b)	Platinum
	(c)	Iridium	(d)	Palladium
82.	Which	n of the following are correct?		
	(a)	The configuration of M o($Z =$	42) i	$s [Kr] 4d^5 5s^1$.
	(b)	The configuration of $Pd(Z = 4)$	6) is	$[Kr] 4d^8 5s^2$.
	(c)	The configuration of $Pd(Z = 4)$	l6) is	$[Kr] 4d^{10} 5s^{0}.$
	(d)	The configuration of $Pt(Z = 7)$	8) is	$[Xe] 4d^9 6s^1.$
83.	Which table?		e pre	esent in group 16 of the periodic
	(a)	Sulphur	(b)	Arsenic
	(c)	Tellurium	(d)	Silicon

84. Which of the following species are isoelectronic with Ne?

(a) N³⁻ (c) Al³⁺ (b) Mg²⁺ (d) Ca²⁺

85.		nich of the following arents or ions shown correct		ders of electron affinity of the	ıe
	(a)	$S > O^-$	(b)	$O > S^-$	
	(c)	$O^- > S^-$	(d)	$N^- > P$	
86.		n of the following salts do NaCl		dergo hydrolysis? KCN	
		KClO ₄		Na ₂ CO ₃	
87.	Which (a) (b) (c)	h of the following stateme Liquid oxygen sticks to t Silver perchlorate is solu The atomic size of silver	ents are co the poles of the in wat is less tha	orrect? of magnet. ter.	of
88.	(a)	h of the following halides AlF_3 AgCl	(b)	le in water? AgI AgF	
89.		h of the following pairs ha Pd and Pt Al and Ga	(b)	ximately the same atomic radii Al and Mg Na and Ne	?
90.	(a)	onal relationships are show Be and Al Mg and Al	(b)	Li and Mg B and P	
91.	(a) (b) (c)	onic bonds X ⁺ Y ⁻ are form electron affinity of Y is h ionization energy of X is lattice energy of XY is hi lattice energy of XY is lo	igh low gh	the	
92.	(a) (b) (c)	th of the folloiwng stateme. The melting point of bor. The melting point of bor. BF $_3$ is a weaker electron trans-1,2-Dichloro-2-pen.	on is greaton is less acceptor t	tter than that of carbon. than that of carbon. than BBr_3 .	
93.		h of the following species AgOH		nown? SH ₆	

(d) PI₅

(c) PbI₄

- **94.** Which of the following pairs has elements that do not belong to the same period?
 - (a) Mg and Sb

(b) Ca and Zn

(c) Na and Ca

- (d) Ca and Cl
- 95. Which of the following pairs do not show the inert-pair effect?
 - (a) Cu and Au

(b) Si and Ge

(c) Tl and Pb

(d) Bi and Sn

		<u>Answers</u>		
1. b	2. d	3. d	4. b	5. a
6. d	7. d	8. b	9. b	10. c
11. d	12. d	13. c	14. d	15. c
16. c	17. b	18. b	19. a	20. c
21. a	22. b	23. a	24. a	25. b
26. b	27. c	28. d	29. b	30. d
31. a	32. c	33. c	34. d	35. c
36. d	37. c	38. c	39. b	40. b
41. b	42. a	43. c	44. d	45. c
46. a	47. c	48. a	49. a	50. a
51. c	52. c	53. d	54. c	55. c
56. b	57. b	58. d	59. b	60. c
61. c	62. a	63. d	64. b	65. d
66. a	67. c	68. d	69. c	70. c
71. b	72. c	73. b	74. a, c	75. b, c, d
76. c, d	77. a, b	78. a, b, c, d	79. b, c	80. a, c
81. a, b, c	82. a, c, d	83. a, c	84. a, c	85. a, b
86. a, c	87. a, b, d	88. a, d	89. a, c	90. a, b
91. a, b	92. b, c	93. a, b, c, d	94. a, c, d	95. a, b

Hints to More Difficult Problems

- 1. The effective nuclear charge is the largest for sodium and the least for phosphorus.
- **2.** These are transition metals with the $(n-1)d^{1-10}ns^{1-2}$ configuration, which is associated with transition contraction.
- 4. Apply Fajans' rules.
- 8. Van der Waals forces

Periodic Table 2-15

- 12. Greatest size, and hence lowest value of ionization energy
- 14. Lattice energy considerations
- **19.** PbI₄ disproportionates easily: PbI₄ \rightarrow PbI₂ + I₂.
- 22. The smaller the size, the greater is the electronegativity
- **23.** The reason for this apparent anomaly is that the ions are hydrated in solution. Since Li⁺ is very small, it is heavily hydrated. This makes the radius of the hydrated ion large. In contrast K⁺ (aq.) is the least hydrated because it is the largest.
- 28. Apply Fajans' rules.
- **30.** The electronegative fluorines in NF₃ balance the electron distribution around N to such an extent that the dipole moment is only 0.2 D.
- 31. The size of the cations, and lattice energy considerations
- **33.** Inert-pair effect
- 37. Close-packed structures
- Hydrogen-bonding in water, and molar mass for the rest of the molecules
- 43. The size of the elements and electronic configuration
- 54. Greater size and lower ionization energy
- 56. Lattice-energy considerations
- **60.** The low dissociation energy of F_2 arises from the repulsion between the unpaired electrons on the two atoms which are themselves small
- 61. Electron-electron interaction and effect of p-orbital repulsion
- 63. Lattice energy $U^- \propto Z_+ Z_ \propto \frac{1}{r}$
- 64. The smaller the size, the greater is the lattice energy.
- 68. Al has the most packed structure.
- 69. Hg has the weakest metallic bond, and Ca the strongest.
- 73. Stable electronic configuration
- 77. Both have pyramidal structures
- **80.** On moving across a period, the charge on the ions increases and the size decreases, causing the polarizing power to increase. On moving down a Group, the size increases and the polarizing power decreases. On moving diagonally these two effects partly cancel each other. So there is no marked change in properties.

- **85.** Energy is evolved when 1 electron is added to an O or S atom forming O^- and S^- ions, but a substantial amount of energy is absorbed when 2 electrons are added to these atoms to form O^{2-} and S^{2-} ions.
- 93. Weak bond strength and coordination unsaturation
- **95.** The inert-pair effect is shown by elements of groups 13, 14, 15 and 16, which have elements of electronic configuration ns^2np^1 , ns^2np^2 , ns^2np^3 and ns^2np^4 respectively.

Hydrogen and Oxygen

• Type 1 •

Choose the correct option. Only one option is correct.

- The hydrogen molecule is not very reactive under normal conditions because
 - (a) hydrogen has a large number of isotopes
 - (b) the corresponding ΔS° value is very low
 - (c) the bond dissociation energy of the hydrogen molecule is high
 - (d) the hydrogen molecule has a low activation energy
- 2. The ionization constant of protium in water $(H_2O \rightleftharpoons H^+ + OH^-)$ is 1.0×10^{-14} and that in heavy water $(D_2O \rightleftharpoons D^+ + OD^-)$ is 3.0×10^{-15} . H_2O dissociates about
 - (a) three times as much as D_2O does.
 - (b) thirty times as much as D_2O does.
 - (c) 0.3 times as much as D_2O does.
 - (d) 300 times as much as D₂O does.
- 3. The order of the heats of fusion of T_2 , D_2 and H_2 is
 - (a) $T_2 > D_2 > H_2$

(b) $H_2 > T_2 > D_2$

(c) $D_2 > T_2 > H_2$

- (d) $D_2 = T_2 > H_2$
- 4. The higher density of water than that of ice is due to
 - (a) dipole-dipole interaction
 - (b) dipole-induced dipole interaction
 - (c) hydrogen bonding
 - (d) all of these

- 5. In parahydrogen, the electron spins are in the
 - (a) same direction but the proton spins are in opposite directions
 - (b) opposite directions but the proton spins are in the same direction
 - (c) same direction as the proton spins
 - (d) opposite direction to the proton spins
- Among the following, the latent heats of fusion and vaporization are maximum in
 - (a) H₂
- (b) D₂
- (c) T₂
- (d) H_2^+
- 7. A dilute solution of H_2O_2 can be concentrated by
 - (a) drying it over anhydrous CaCl₂
 - (b) drying it over concentrated H₂SO₄
 - (c) drying it over anhydrous MgSO₄
 - (d) heating it under reduced pressure
- 8. Which of the following mixtures is known as the Fenton reagent?
 - (a) $TiCl_4$ and $Al(C_2H_5)_3$
- (b) FeSO₄ and H₂O₂
- (c) FeCl₃ and H₂O₂
- (d) CH₃COONH₄ and H₂O₂
- 9. Which of the following is the true structure of H_2O_2 ?
 - (a) H O-Ö:

(b) H

- $(c) \overset{H}{\circ} \overset{\bullet}{\circ} \overset{\bullet}{\circ} \\ H$
- (d) H—O=O—H
- 10. Hydrogen gas is generally prepared by the
 - (a) reaction of granulated zinc with dilute H₂SO₄
 - (b) reaction of zinc with concentrated H₂SO₄
 - (c) reaction of pure zinc with dilute H₂SO₄
 - (d) action of steam on red-hot coke
- 11. Superoxides are
 - (a) stronger oxidizing agents than peroxides
 - (b) weaker oxidizing agents than peroxides
 - (c) weaker reducing agents than oxygen
 - (d) stronger reducing agents than H₂O₂
- **12.** Which of the following statements is correct? NaO₂ has a
 - (a) graphite-like structure.
 - (b) rock-salt (NaCl)-like structure.

		pyrite (FeS ₂)-like structure. fluorite (CaF ₂)-like structure.		
12		- and parahydrogen differ in		
15.		atomic number	(b)	mass number
	(c)	nuclear spin	(d)	all of these
14.	In ozo	one, the central oxygen atom us	ses	
	(a)	roughly sp^2 orbitals for σ -bon	ding	5
	(b)	sp orbitals for π -bonding		
	(c)	sp^3 orbitals for σ -bonding		
	(d)	pd_z^2 orbitals for σ - and π -bond	ling	
15.	The st	ructure of ozone involves		
	(a)	delocalized three-centre σ -box	ndin	g
	(b)	delocalized three-centre π -box	ndin	g
	(c)	delocalized three-centre $\sigma\!\!$ - as	well	as π-bonding
	(d)	localized π -bonding		
16.	The p	ale blue colour of ozone gas is	due	to the intense absorption of
		infrared radiation		cosmic radiation
		red light		blue light
17.		e is a powerful oxidizing agent less oxidizing than F ₂		
		less oxidizing than O_2		more oxidizing than F_2 less oxidizing than H_2O_2
10		e oxidizes PbS to	(61)	rece extending than 11202
10.		PbO ₂	(b)	Pb ₃ O ₄
		PbO	. ,	PbSO ₄
19.	Ozone	e reacts with a KOH solution to	pro	duce
		O ₂ and K ₂ O	•	O ₂ and K ₂ O ₂
	(c)	O ₂ only	(d)	KO_3
20.		olume of oxygen obtained by t	the d	decomposition of 4 L of O ₃ at stp
	is (a)	3 L	(b)	9 L
		6 L		2 L
21.	Ozone			
		an allotrope of oxygen	(b)	an isomer of oxygen
	(c)	an isotone of oxygen	(d)	isostructural with H ₂ O ₂

2-20	Chemistry MCQ						
22.	Which of the following compounds produces holes in the ozone layer in the upper atmosphere?						
	(a)	Sulphur dioxide	(b)	Carbon dioxide			
	(c)	Freon	(d)	Carbon monoxide			
23.	3. Which of the following statements is correct?						
	(a) The ionization energy of hydrogen is the same as the fir- ionization energy of helium.						
	(b) The electronegativity of hydrogen is the same as that of carbon.						
	(c) Hydrogen gas is liberated at the anode during the electrolysis of molten lithium hydride.						

- (d) Hydrogen gas is liberated at the cathode during the electrolysis of molten lithium hydride.
- **24.** The ozonolysis of C = C produces

(a)
$$\supset C = O + O = C \subset$$

$$(b)$$
 $C = C = O$

$$C - C = O$$

$$(d) \rightarrow C - OH + CO_2$$

- 25. Ozone can be detected by using
 - (a) Na

(b) Ag

(c) Hg

- (d) Ar
- **26.** A pale blue gas with a fish-like odour restores the colour of a blackened lead painting. The gas is also used as a rocket fuel. It is
 - (a) O_2

(b) NH₃

(c) SO_2

- (d) O₃
- 27. Water undergoes self-ionization to a small extent to give
 - (a) H^+ and OH^-

(b) OH⁺ and H⁻

(c) H_3O^+ and OH^-

- (d) $H_3O_2^+$ and OH^-
- 28. In which of the following compounds does deuteration take place easily on treatment with D_2O ?
 - (a) CH₃CH₂OH

(b) CH₃—CH₃

(c) CH₃CH₂—O—CH₂CH₃

- (d) CH₃CH₂Cl
- 29. Which of the following pairs have an almost identical geometry?
 - (a) H_3O^+ and AlF_3

(b) NH₃ and H₃O⁺

(c) H₂O and ICl₂

(d) H_2O and NO_2^+

- 30. Which of the following compounds has not yet been prepared?
 - (a) D_2S

(b) T₂S

(c) D₂O

- (d) H₂Po
- 31. Which of the following nuclear reactions is used to prepare tritium?

- (a) ${}^{8}_{4}Be + {}^{2}_{4}He \longrightarrow {}^{3}_{1}H + {}^{9}_{5}B$ (b) ${}^{2}_{1}D + {}^{2}_{1}D \longrightarrow {}^{3}_{1}H + {}^{1}_{1}H$ (c) ${}^{3}_{3}Li + {}^{1}_{0}n \longrightarrow {}^{3}_{1}H + {}^{4}_{2}He$ (d) ${}^{10}_{5}B + {}^{1}_{0}n \longrightarrow 2 \, {}^{4}_{2}He + {}^{3}_{1}H$
- 32. Acetone exhibits keto-enol tautomerism:

$$(CH_3)_2C = O \longrightarrow CH_3 - C = CH_2$$

Which of the following products is obtained when acetone is treated with an excess of D₂O for a sufficient time in the presence of a small amount of a dilute NaOH solution?

(a) OD
$$CH_3 - C = CH_2$$

(b) OH
$$CH_2D-C=CH_2$$

(c)
$$O$$
 \parallel CD_3-C-CD_3

(d) OD
$$CH_3 - C = CHD$$

- 33. Which of the following is incorrect?
 - (a) $NH_4^+ + D_2O \rightleftharpoons NH_3D^+ + HDO$
 - (b) $PH_3 + D_2O \rightleftharpoons PD_3 + H_2O$
 - (c) $CH_3NH_2 + D_2O \rightleftharpoons CH_2NHD + HDO$
 - (d) $CH_3OH + D_2O \rightleftharpoons CH_3OD + HDO$
- 34. Most covalent compounds have a very low solubility in water because of the
 - (a) low hydration energy due to the weak interaction between the polar water molecules and the nonpolar covalent molecules, which is insufficient to break the hydrogen bonds between the water molecules
 - (b) low dielectric constant of water
 - (c) high dipole moment of water
 - (d) high density of water
- 35. Zeolite is
 - (a) hydrated sodium aluminium silicate—Na₂Al₂Si₂O₈⋅xH₂O
 - (b) hydrated ferric oxide
 - (c) sodium hexametaphosphate
 - (d) sodium tetraborate

- 36. Hard water is not suitable for washing clothes because
 - (a) it contains Na₂SO₄ and KCl
 - (b) it hydrolyses soap
 - (c) it gives a precipitate with soap
 - (d) water forms micelles with soap
- 37. The degree of hardness of water is usually expressed in terms of
 - (a) parts per million by weight of MgSO₄
 - (b) grams per litre of CaCO₃ and MgCO₃ actually present
 - (c) parts per million by weight of CaCO₃ regardless of whether it is actually present
 - (d) parts per million of CaCO₃ actually present in water
- 38. Hydrogen is obtained as a by-product in the
 - (a) electrolysis of water
 - (b) manufacture of caustic soda
 - (c) Bosch process
 - (d) Lane process
- 39. Calgon, which is used as a water softener, has the formula
 - (a) $Na_4[Na_2(PO_3)_6]$
- (b) $Na_2[Na_4(PO_3)_6]$
- (c) $Na_2[Na_4(PO_4)_5]$
- (d) $Na_2[Na_2(PO_4)_6]$
- 40. Heavy water freezes at
 - (a) 270.7 K

(b) 276.1 K

(c) 269.2 K

- (d) 276.8 K
- 41. Heavy water is usually prepared by the
 - (a) reaction of D₂ with NH₄ OH
 - (b) oxidation of D₂ with ozone
 - (c) prolonged multistage electrolysis of a 0.5 M NaOH solution using nickel electrodes
 - (d) prolonged electrolysis of the solution of a mixture of $\rm Ba(OH)_2$ and $\rm H_2SO_4$

• *Type 2* •

Choose the correct options. More than one option is correct.

- 42. Water may be softened using
 - (a) sodium aluminium silicate
- (b) Graham's salt
- (c) an ion-exchange resin
- (d) trisodium phosphate

(b) using anhydrous CoCl₂ which changes colour from blue to pink

(b) an oxidant of rocket fuel

(d) a bleaching agent

(a) using anhydrous CuSO₄ which changes colour

43. The presence of water can be inferred by

(c) the use of hydrated CuSO₄

45. Which of the following statements are correct?

(d) smell

44. Hydrogen peroxide is used as(a) a reducing agent

(c) an oxidizing agent

(a) Hydrogen, like alkali metals, is electropositive. (b) Like alkali metals, hydrogen shows an oxidation state of +1 in its compounds. (c) Like alkali metals, hydrogen acts as a strong reducing agent. (d) Hydrogen shows an oxidation state of -1 in covalent hydrides. **46.** Which of the following statements are correct? (a) Like halogens, hydrogen combines with nonmetals to form covalent compounds. (b) Hydrogen as well as halogens have a tendency to accept electrons. (c) The oxide of hydrogen is neutral, as is the case with the oxides of halogens, e.g., Cl₂O₇. (d) Hydrogen and halogens form hydride ions with equal ease. 47. In which of the following does hydrogen exist in a negative oxidation state? (a) HCl (b) CaH₂ (d) Li[AlH₄] (c) B_2H_6 **48.** Which of the following statements are correct? (a) Water is amphoteric. (b) Water acts as an oxidizing agent. (c) Water acts as a reducing agent. (d) Sodium hydride is insoluble in water. **49.** The decomposition of H_2O_2 is retarded by (a) acetanilide (b) glycerol (c) sodium bicarbonate (d) oxalic acid **50.** Which of the following statements are correct? (a) Dihydrogen is neutral to litmus. (b) Dihydrogen dissociates into hydrogen atoms only on being heated to above 2000 K.

51.

52.

53.

54.

ions become larger.

56. 10-volume H_2O_2 is equivalent to

(a) 10%

(c) $30 \, \text{g/L}$

(a) Ca^{2+} ions

(c) Mg^{2+} ions

(c) The bond order of superoxides is 1.5.

55. Which of the following statements are correct?(a) H₂O₂ is a pale blue viscous liquid.

(d) H₂O₂ has an 'open-book' structure.

(c)	The reactivity of hydrogen towards halogens decreases in the order $Cl > Br > I > F$.								
(d)	Dihydrogen can reduce CuO and ZnO to their metals.								
Which of the following metals form saline hydrides?									
(a)	•								
(c)	Na (d) Mg								
Which	of the following statements are correct?								
(a)	At 0 K, hydrogen gas contains 100% parahydrogen.								
(b)	At high temperature, hydrogen contains about 75% orthohydrogen.								
(c)	Parahydrogen has lower energy than orthohydrogen.								
(d)	At high temperature, hydrogen contains about 75% parahydrogen.								
Which of the following statements are correct in the context of peroxide ions?									
(a)	They are oxidizing agents.								
(b)	They are salts of a weak diprotic acid (H ₂ O ₂).								
(c)	They react with acids to give H_2O_2 .								
(d)	The bond order of the O_2^{2-} ion is 1.5.								
Which	of the following statements are correct?								
(a)	Peroxide and superoxide ions are larger than oxide ions.								

(b) The stability of peroxides and superoxides increases as the metal

(d) Many peroxides are coloured due to the presence of a superoxide.

(b) $3\% H_2O_2(W/V)$

(d) 1.786 N

(b) H⁺ions

(d) OH-

(b) H₂O₂ can act as an oxidizing as well as a reducing agent.
(c) In H₂O₂ the two hydroxyl groups lie on the same plane.

57. On treatment of hard water with zeolite, sodium ions get exchanged with

- **58.** Which of the following statements are correct?
 - (a) There is a layer of ozone in the upper atmosphere which absorbs harmful UV radiation from the sun, thus protecting life on earth.
 - (b) Freons produce holes in the upper atmosphere.
 - (c) Oxides of nitrogen and halogens can damage the ozone layer.
 - (d) Ozone is prepared by the action of a silent electric discharge upon an oxygen molecule.

Answers							
1. c	2. a	3. a	4. c	5. d			
6. c	7. d	8. b	9. c	10. a			
11. a	12. a	13. c	14. a	15. b			
16. c	17. a	18. d	19. d	20. c			
21. a	22. c	23. c	24. a	25. c			
26. d	27. c	28. a	29. b	30. b			
31. b	32. c	33. b	34. a	35. a			
36. c	37. c	38. b	39. b	40. d			
41. c	42. b, c, d	43. a, b	44. a, b, c, d	45. a, b, c			
46. a, b	47. b, d	48. a, b, c	49. a, b	50. a, b, d			
51. a, c	52. a, b, c	53. a, b, c	54. a, b, c, d	55. a, b, d			
56. b, c, d	57. a, c	58. a, b, c, d					

Hints to More Difficult Problems

- 3. Decreasing atomic mass
- **11.** Higher reduction potential
- **17.** $E_{O_2/O^{2-}}^0 = +2.07 \text{ V}$
- **24.** O_3 attacks the C=C double bond.
- **26.** Exothermicity
- 28. The more polar CH₃CH₂OH causes D₂O to react or exchange.
- 31. Apply the minimum-energy concept.
- **33.** PH_3 does not contain labile hydrogen. The electronegativity values of P and H are nearly the same.
- 38. $4OH^- \longrightarrow 2H_2O + O_2 + 4e$
- 40. From the partial phase diagram of water

- 47. (b) and (d) are metal hydrides.
- 48. $2H_2O + 2e \longrightarrow 2OH^- + H_2$ (H_2O is an oxidizing agent) $2H_2O \longrightarrow 4H^+ + O_2 + 4e$ (H_2O is a reducing agent) $H_2O \rightleftharpoons H^+ + OH^-$ (H_2O is amphoteric) $10^{-7} \text{ mol/L} \quad 10^{-7} \text{ mol/L}$
- 49. Acetanilide and glycerol are negative catalysts.
- 51. Li and Na form ionic hydrides.
- 56. $2H_2O_2 \longrightarrow 2H_2O_+ O_2$ 68 g 22400 mL at stp $10\text{-vol.} \ H_2O_2$ means that 1 vol. of $H_2O_2 \equiv 10 \text{ vol.}$ of O_2 at stp or 1 mL of $H_2O_2 \equiv 10 \text{ mL}$ of O_2 at stp 22400 mL of O_2 at stp = 68 g of H_2O_2 10 mL of O_2 at stp $= 3 \times 10^{-2} \text{ g}$ of H_2O_2 Thus 1 mL of H_2O_2 soln $= 3 \times 10^{-2} \text{ of } H_2O_2$ 100 mL of H_2O_2 soln = 30 g of H_2O_2 = 100 mL of H_2O_2 soln = 30 g of H_2O_2 i.e. 3% W/V

Ц

Alkali Metals and Alkaline Earth Metals

• Type 1 •

1. The magnitude of enthalpy of formation of alkali metal halides decreases

Choose the correct option. Only one option is correct.

(a) iodide > bromide > chloride > fluoride
(b) bromide > iodide > fluoride > chloride
(c) fluoride > chloride > bromide > iodide
(d) fluoride > chloride > iodide > bromide

in the order

2.	Alkali	metals react	netals react with hydrogen forming ionic hydrates. The reactivity						
	of the alkali metals with hydrogen decreases in the order								
	(a)	Li > Na > K	> Rb	> Cs	(b)	Na > K > Rb	> Cs	s > Li	
	(c)	Rb > Cs > Li	> Na	a > K	(d)	Cs > Rb > K	> Na	a > Li	
3.	Which of the following reacts with nitrogen to form a nitride which finally hydrolyses to ammonia?								h
	(a)	K			(b)	Li			
	(c)	Na			(d)	Cs			
3.	Which	ch of the following pairs cannot exist together in solution?							
	(a)	Na ₂ CO ₃ and NaOH			(b)	Na ₂ CO ₃ and NaHCO ₃			
	(c)	NaHCO ₃ and NaOH			(d)	NaHCO ₃ and NaCl			
5.	Among the following, which has the least ionic mobility?								
	(a)	Na +	(b)	K ⁺	(c)	Li +	(d)	Cs +	
6.	The la	ttice energies	of th	ne oxides of N	Иg, (Ca, Sr and Ba	follo	w the order	
	(a)	BaO > SrO >	· CaC) > MgO	(b)	CaO > BaO	> SrC	O > MgO	

(d) MgO > CaO > SrO > BaO

(c) MgO > SrO > CaO > BaO

7.		hen an alkali metal is introduced into a flame, it imparts a distinctive lour to it. The colour arises from						
		electronic transitions in species formed momentarily in the flame						
		the electronic transition 3s ¹ –		-				
		proton-proton transition		- 1				
		d–d transition						
8	Which	n of the following is used in ph	otoe	lectric cells?				
0.	(a)	Na	(b)					
	(c)	Li	` '	Cs				
9.	When	heated to 800°C, NaNO ₃ gives						
٠,		NaNO ₂ + O ₂		$Na_2O + O_2 + N_2$				
	(c)	$Na + N_2 + O_2$	(a)	$NaN_3 + O_2$				
10.				nmonia is blue, and is a strong				
		ing agent, due to the presence sodium atoms	OI					
		sodium hydride						
		sodium amide						
	` '	solvated electrons and solvate	ed m	etal ions				
11		netallic lustre of sodium is expl						
11.	(a)	fcc lattice structure	anic	a by its				
	` '	the effusion of sodium ions						
		the oscillation of its mobile va	lenc	e electrons				
	(d)	the band theory of metals						
12.	Which	n of the following forms the lea	st io	nic chloride?				
	(a)	Ве		Ca				
	(c)	Mg	(d)	Sr				
13.	Lithiu	ım shows a diagonal relationsh	nip w	rith				
		sodium	_	silicon				
	(c)	nitrogen	(d)	magnesium				
14.		m thiosulphate may be prepar n of them is called Spring's rea		y any of the following reactions. ?				
	(a)	$Na_2SO_3 + S \longrightarrow Na_2S_2O_3$						
	(b)	$Na_2S + I_2 + Na_2SO_3 \longrightarrow Na_2$	S_2O_3	+ 2NaI				
	(c)	$2Na_2S + Na_2CO_3 + 4SO_2 \longrightarrow$	3Na	$a_2S_2O_3 + CO_2$				
	(d)	$S + 6NaOH \longrightarrow Na_2S_2O_3 + 2$	2Na ₂	$S + 3H_2O$				

- **15.** Which of the following statements is correct?
 - (a) Sodium reacts with liquid ammonia to give hydrogen.
 - (b) Sodamide is decomposed by water, producing ammonia.
 - (c) Sodamide is decomposed by water, producing nitrogen.
 - (d) Sodamide is used as a reducing as well as a dehydrating agent.
- 16. Lithium differs from alkali metals due to
 - (a) the low hydration energy of Li+
 - (b) its b.c.c. structure
 - (c) its extremely high electropositivity
 - (d) its small atomic and ionic size
- 17. Sodium thiosulphate is used in photography to
 - (a) convert metallic silver to a silver salt
 - (b) reduce silver bromide to a silver salt
 - (c) remove unreduced silver
 - (d) remove undecomposed AgBr as a soluble silver thiosulphate complex
- 18. Sodium carbonate reacts with SO₂ in an aqueous medium to give
 - (a) NaHSO₃
- (b) NaHSO₄
- (c) Na₂SO₃
- (d) Na_2SO_4
- 19. In a nuclear reactor, molten sodium is used to
 - (a) absorb neutrons for controlling the chain reaction
 - (b) absorb the heat generated by a nuclear reaction
 - (c) slow down fast neutrons
 - (d) extract the radioisotopes produced in the reactor
- 20. When hydrated magnesium chloride (MgCl₂·6H₂O) is heated,
 - (a) MgO is formed
 - (b) anhydrous MgCl₂ is formed
 - (c) Mg(OH)HCl is formed
 - (d) Mg(OH)Cl is formed
- 21. Anhydrous magnesium chloride can be prepared by heating MgCl₂.6H₂O
 - (a) with concentrated HCl
 - (b) in an atmosphere of nitrogen
 - (c) in a current of dry HCl gas
 - (d) in an atmosphere of hydrogen
- **22.** In the Solvay process
 - (a) an ammoniacal brine solution is carbonated with CO₂, forming NaHCO₃ which on decomposition at 150°C produces Na₂CO₃

- (b) a sodium amalgam reacts with water to produce NaOH which gives Na₂CO₃ on reacting with CO₂
- (c) a brine solution is made to react with BaCO₃ to produce Na₂CO₃
- (d) brine is carbonated with CO₂ forming NaHCO₃ which on decomposition at 150°C produces Na₂CO₃
- 23. When a saturated solution of magnesium sulphate is treated with NH₄Cl and NH₃, followed by the addition of disodium hydrogen phosphate, a white precipitate of
 - (a) $Mg_2P_2O_7$

(b) $Mg_3(PO_4)_2$

(c) Mg(NH₄)PO₄

(d) Mg(NH₄)HPO₄

is formed.

- 24. The order of the solubility of the sulphates of alkaline earth metals in water is
 - (a) Be > Mg > Ca > Sr > Ba
- (b) Mg > Be >> Ba > Ca > Sr
- (c) Be > Ca > Mg > Ba >> Sr (d) Mg > Ca > Ba >> Be > Sr
- 25. The solubility of the fluorides and hydroxides of alkaline earth metals increases on descending the group because the
 - (a) lattice energy of the compounds increases more rapidly than the hydration energy
 - (b) lattice energy of the compounds decreases more rapidly than the hydration energy
 - (c) size of the metals decreases on descending the group
 - (d) ionization energy of the metals increases on descending the group
- 26. On descending Group 2, the ions of the corresponding metals become larger. Therefore, the
 - (a) lattice energy and the hydration energy of the compounds decrease
 - (b) lattice energy of the compounds decreases and the hydration energy of the compounds encreases
 - (c) lattice energy and the hydration energy of the compounds increase
 - (d) lattice energy of the compounds increases and the hydration energy of the compounds decreases
- **27.** Which of the following is the least thermally stable?
 - (a) MgCO₃

(b) CaCO₃

(c) SrCO₃

- (d) BeCO₃
- 28. Magnesium is obtained by the electrolysis of fused magnesium chloride with sodium chloride. Sodium chloride helps
 - (a) lower the conductivity of MgCl₂

(b) raise the conductivity of MgCl₂ (c) lower the melting point of MgCl₂ (d) change the solid-state structure of MgCl₂ 29. Magnesium is obtained by the electrolysis of fused magnesium chloride using (a) a nickel cathode and a graphite anode (b) a nickel container as cathode and an iron anode (c) an iron container as cathode and a graphite anode (d) a lead cathode and a platinum anode 30. The reducing property of alkali metals follows the order (a) Na < K < Rb < Cs < Li(b) K < Na < Rb < Cs < Li(c) Li < Cs < Rb < K < Na (d) Rb < Cs < K < Na < Li31. An aqueous solution of magnesium sulphate and sodium carbonate gives (a) MgCO₃ (b) MgCO₃.Mg(OH)₂ (d) Mg(HCO₃)₂ (c) $Mg(OH)_2$ 32. An aqueous solution of magnesium sulphate and sodium bicarbonate gives (a) $MgSO_4.Mg(OH)_2$ (b) MgCO₃.Mg(OH)₂ (d) MgCO₃ (c) Mg(OH)₂ 33. Group 2 metals are (a) harder and have higher cohesive energies and melting points than group 1 metals (b) softer and have lower cohesive energies and melting points than group 1 metals (c) softer and have lower cohesive energies and higher melting points than group 1 metals (d) harder and have higher cohesive energies and lower melting points than group 1 metals **34.** The electrolysis of carnallite (KCl⋅MgCl₂⋅6H₂O) yields (a) potassium ions (b) magnesium ions (c) magnesium and chlorine (d) potassium and chlorine 35. Calcium chloride reacts with ethyl alcohol to produce (a) $CaCl_2 \cdot 2C_2H_5OH$ (b) CaCl₂·6C₂H₅OH (c) CaCl₂·4C₂H₅OH (d) CaCl₂·8C₂H₅OH 36. The chlorine available in bleaching powder is estimated by

(b) permanganometric titration

(a) acid-base titration

	(c)	dichrometri	c titration	(d)	iodometric	titrat	ion		
37.	A mixture of MgCl ₂ and calcined magnesia is called								
	(a)	Sorel cemen	nt	(b)	Portland cer	ment			
	(c)	a double sal	lt	(d)	dental ceme	ent			
38.	The cyanamide ion $[N=C=N]^{2-}$ is isoelectronic with								
	(a)	CO ₂ , and is	bent	(b)	CO ₂ , and is	linea	r		
	(c)	N ₂ O, and is	V-shaped	(d)	N_3^- , and is p	yran	nidal		
39.	Calcium cyanamide is produced by heating								
	(a)	CaC ₂ in an e	electric fur	nace in an a	tmosphere of	nitro	ogen at 110	00°C	
	(b)	calcium oxide with nitrogen and carbon dioxide at 1000°C							
	(c)	calcium nitrate with calcium oxide at 1500°C							
	(d)	calcium oxid	de with an	nmonia and	then with car	rbon	at 2000°C		
40.	Calciu	ım is obtaine	d by the						
	(a)	roasting of limestone							
	(b)	reduction of	f calcium c	chloride by c	arbon				
	(c)	electrolysis of an aqueous solution of calcium chloride							
	(d)	electrolysis	of molten	calcium chlo	oride				
41.	Hydr	ides as well a	s halides o	of alkaline ea	arth metals te	nd to	o polymeri	ze	
	(a)	strontium		(b)	calcium				
	(c)	beryllium		(d)	magnesium				
42.	Which	n of the follow	wing is not	t known to e	xist?				
	(a)	BaO_2	(b) BeO	(c)	SrO ₂	(d)	CrO_5		
43. In case of alkaline earth metals, which of the following					ing	increases	with		
		c number?		1 1 .1					
		The solubili	-	-					
		The solubili	-	suipnates					
		Electronega Ionization e	•						
44.	A salt is soluble in water if its (a) hydration energy is more than its lattice energy								
		hydration energy is more than its lattice energy hydration energy is less than its lattice energy							
) hydration energy is equal to its lattice energy							
	(d)								
	(32)	hydration e							
45.	Whicl	n of the follow	wing is not	t a peroxide?	•				
		KO_2	(b) CrO		Na_2O_2	(d)	BaO_2		

- 46. The bicarbonates of group 2 metals are
 - (a) easily oxidized by air
 - (b) stable only in solution
 - (c) unstable in solution
 - (d) stable to heat in the solid state
- 47. Which of the following statements about alkali metals is correct?
 - (a) They can be extracted by the reduction of their oxides.
 - (b) They can be displaced from the aqueous solutions of their salts by other metals.
 - (c) They can be isolated by the electrolysis of the aqueous solutions of their salts.
 - (d) They can be isolated by the electrolysis of their molten salts.
- **48.** All group 2 metals dissolve in liquid ammonia to produce a bright blue colour. The colour is due to
 - (a) a change in the structure of the ammonia
 - (b) d-d transition
 - (c) the spectrum of the solvated electrons
 - (d) an electronic transition from a lower to a higher energy state
- **49.** The hydration energies of group 2 ions are approximately four times as much as those of group 1 ions. This is due to their
 - (a) smaller size and greater nuclear charge
 - (b) greater size and lower nuclear charge
 - (c) smaller size and lower nuclear charge
 - (d) greater size and greater nuclear charge
- **50.** The lattice energies of the carbonates of Mg, Ca, Sr and Ba follow the order
 - (a) $MgCO_3 > CaCO_3 > SrCO_3 > BaCO_3$
 - (b) $CaCO_3 > SrCO_3 > BaCO_3 > MgCO_3$
 - (c) $BaCO_3 > SrCO_3 > CaCO_3 > MgCO_3$
 - (d) $SrCO_3 > MgCO_3 > CaCO_3 > BaCO_3$
- 51. Which of the following is arranged in order of increasing melting point?
 - (a) Ca < Be < Mg < Sr
- (b) Be < Ca < Sr < Mg
- (c) Mg < Sr < Ca < Be
- (d) Sr < Mg < Be < Ca
- **52.** Which of the following statements is incorrect?
 - (a) The ionization energy of Be²⁺ is high and its compounds are covalent.
 - (b) The ionization energy of Be²⁺ is low and its compounds are covalent.

- (c) The compounds formed by Mg, Ca, Sr and Ba are predominantly divalent and ionic.
- (d) The ionization energy of Be ²⁺ is low, and its compounds are ionic.
- 53. For compounds with the same anion, the hydration energies of Na^+, K^+, Rb^+, Cs^+ and Li^+

follow the order

- (a) $Cs^+ > Rb^+ > K^+ > Na^+ > Li^+$ (b) $Li^+ > Na^+ > K^+ > Rb^+ > Cs^+$
- (c) $K^+ > Na^+ > Li^+ > Cs^+ > Rb^+$ (d) $Li^+ > K^+ > Na^+ > Cs^+ > Rb^+$
- **54.** The raw material used in the Solvay process for the manufacture of sodium carbonate comprises
 - (a) sodium chloride and carbon dioxide
 - (b) ammonia and carbon dioxide
 - (c) sodium chloride, limestone and ammonia
 - (d) sodium chloride, limestone and carbon dioxide
- 55. The extents of hydration of the ions of Be, Mg, Ca, Sr and Ba follow the order
 - (a) $Ca^{2+} > Ba^{2+} > Sr^{2+} > Be^{2+} > Mg^{2+}$
 - (b) Be²⁺ > Mg²⁺ > Ca²⁺ > Sr²⁺ > Ba²⁺
 - (c) Ba²⁺ > Sr²⁺ > Ca²⁺ > Mg²⁺ > Be²⁺
 - (d) $Sr^{2+} > Ba^{2+} > Ca^{2+} > Be^{2+} > Mg^{2+}$

• *Type 2* •

Choose the correct options. More than one option is correct.

- 56. Sodium thiosulphate is prepared by
 - (a) boiling an Na₂SO₃ solution with S in an acidic medium
 - (b) boiling an NaOH solution with S
 - (c) neutralizing an H₂SO₄ solution with NaOH
 - (d) boiling Na₂SO₃ with S in an alkaline medium
- 57. The oxide of which of the following metals is amphoteric?
 - (a) Pb

(b) Mg

(c) Ca

- (d) Al
- **58.** Which of the following are good conductors of electricity in the molten state?
 - (a) BeCl₂
- (b) CaCl₂
- (c) SrCl₂
- (d) MgCl₂

59. Chlorides of which of the following metals crystallize from an aqueous

61. In the extraction of which of the following metals is amalgamation used?

(c) K

(b) K

(d) Cs

(b) Au

(d) Fe

(b) Na

60. Which of the following elements exhibit photoelectric effect?

solution as hydrates?

(a) Li

(a) Na

(c) Li

(a) Ag

(c) Cu

(d) Mg

62.		The fluxes that can be used for the removal of basic impurities like CaO and FeO from an ore are			
	(a)	SiO_2	(b)	$Na_2B_4O_7 \cdot 10H_2O$	
	(c)	$MgCO_3$	(d)	CaCO ₃	
63.	Alkal	ine earth metals are			
	(a)	more reactive		less reducing	
	(c)	more oxidizing	(d)	less basic	
	than a	ılkali metals.			
64.	Which	n of the following statements a	re co	rrect?	
	(a)	Alkali metals are better remetals.	duci	ng agents than alkaline earth	
	(b)	SF ₆ is well known but SH ₆ is a	not k	nown to exist.	
		BCl ₃ is a stronger Lewis acid	than	BF_3 .	
	(d)	Boron forms B ³⁺ ions.			
65.	Na ₂ SO	O ₄ is soluble in water while BaS	5O ₄ is	s sparingly soluble because	
	(a)	the lattice energy of $BaSO_4$ is	more	e than its hydration energy	
	(b)	the hydration energy of Na ₂ S	O ₄ is	less than its lattice energy	
	(c)	the hydration energy of Na ₂ S	O ₄ is	more than its lattice energy	
	(d)	d) The hydration energy and the lattice energy have no role to play in the solubility of a substance.			
66.	Which	n of the following statements a	re co	rrect for group 1 metals?	
	(a)	They all have one electron in shell containing eight electron		outer shell preceded by a closed	
	(b)		n wł	s are generally ionic and exist as nich as many ions of opposite ssible.	
	(c)	Their compounds are generathe anions are coloured.	ılly v	water soluble and white, unless	

(d) The reactivity increases from lithium to caesium.

- 67. Which of the following statements is correct for alkali metals?
 - (a) They can form covalent molecules such as Li₂, Na₂ and K₂. About 1% of these are in the vapour state.
 - (b) C_6H_5Li is highly reactive.
 - (c) They form compounds with the liberation of heat.
 - (d) They all adopt ccp structures.

68. Which of the following statements is correct?

- (a) When one electron is removed from a group 2 metal, the ratio of nuclear charge to the number of orbital electrons is increased, so that the remaining electrons are more tightly held.
- (b) The energy needed to remove the second electron from a group 2 metal is nearly double that required for the first electron.
- (c) When ionic compounds are formed by group 2 metals, energy is released.
- (d) When ionic compounds are formed by group 2 metals, energy is absorbed.

69. Which of the following statements is correct?

- (a) The solubility of a group 2 salt depends upon the lattice energy of the solid and the hydration energy of the ions.
- (b) The solubilities of group 2 fluorides and hydroxides have opposite trends.
- (c) The solubilities of most group 2 salts decrease with an increase in the atomic weight of the corresponding metal.
- (d) The solubilities of group 2 fluorides and hydroxides increase with molecular weight.

70. Which of the following statements is correct for group 2 metals?

- (a) On descending the group the metal ions become larger, and so the lattice energy as well as the hydration energy decrease.
- (b) The solubility of their salts increases as the lattice energy decreases, and decreases with hydration energy.
- (c) For a substance to dissolve, the hydration energy must exceed the lattice energy.
- (d) None of these.

71. Which of the following statements is incorrect for alkali metals?

- (a) They do not exist in combination with other elements or radicals.
- (b) They are extremely reactive and electropositive.
- (c) All of them adopt a bcc structure.
- (d) All of them adopt an fcc structure.

- 72. Magnesium may be called a refractory because it
 - (a) has high melting point
 - (b) is a good conductor of heat
 - (c) is chemically inert as well as an electrical insulator
 - (d) forms Grignard's reagent
- 73. Which of the following is correct reaction which is feasible.

(a)
$$Zn + NaOH + NaNO_3 \longrightarrow Na_2ZnO_2 + N_2 + H_2O$$

- (b) $Al + NaOH + H_2O \longrightarrow NaAlO_2 + H_2$
- (c) $Br_2 + Na_2CO_3 \longrightarrow NaBr + NaBrO_3 + CO_2$
- (d) $Sn + NaOH + H_2O \longrightarrow Na_2SnO_3 + H_2$
- 74. Potassium iodide acts as a reducing agent when treated with
 - (a) an acidified K₂Cr₂O₇ solution
 - (b) an acidified KMnO₄ solution
 - (c) a CuSO₄ solution
 - (d) a lead acetate solution
- 75. Which of the following statements is correct for compounds of group 2 metals?

Answers

- (a) The number of molecules of water of crystallization increases with the size of the metal ions.
- (b) The number of molecules of water of crystallization increases as the size of the metal ions decreases.
- (c) The number of molecules of water of crystallization decreases as the size of the metal ions increases.
- (d) None of these

1. c	2. a	3. c	4. d	5. c
6. d	7. a	8. d	9. b	10. d
11. c	12. a	13. d	14. b	15. b
16. d	17. d	18. a	19. b	20. d
21. c	22. c	23. c	24. a	25. b
26. a	27. d	28. c	29. c	30. a
31. b	32. d	33. a	34. c	35. c
36. d	37. a	38. b	39. a	40. d
41. c	42. b	43. a	44. a	45. a
46. b	47. d	48. c	49. a	50. a

51. c	52. d	53. b	54. c	55. b
56. b, d	57. a, d	58. b, c, d	59. a, d	60. a, b, d
61. a, b	62. a, b	63. b, d	64. a, b, c	65. a, c
66. b, c, d	67. a, b, c	68. a, b, c	69. a, b, c	70. a, b, c
71. a, d	72. a, b, c	73. b, c, d	74. a, b, c	75. b, c

Hints to More Difficult Problems

- 1. Fluorides are the most stable and iodides the least.
- **2.** Size factor, i.e. $r_{L_1} + < \ldots, < r_{C_8} +$
- The smallest ion (Li⁺) is the least mobile, because it forms big clusters of ions in water.
- **12.** Beryllium is extremely small in size and, according to Fajans' rules, small highly charged ions tend to form covalent compounds.
- **13.** See the answer to Q. 80 of Chapter 1.
- 23. $MgSO_4 + NH_3 + Na_2HPO_4 \longrightarrow Mg(NH_4)PO_4 \downarrow + Na_2SO_4$ white ppt
- 27. BeCO₃ has the lowest lattice energy.
- This is the increasing order of the standard reduction potential of alkali metals.

34.
$$MgCl_2 \longrightarrow Mg^{2+} + 2Cl^-$$

$$\downarrow \qquad \qquad \downarrow \qquad \qquad \downarrow$$
 $Mg \qquad Cl_2$
(cathode) (anode)

KCl is unaffected by the applied voltage and amperage.

36.
$$Ca(OCl)Cl + H_2O \longrightarrow Ca(OH)_2 + Cl_2$$

 $Cl_2 + 2KI \text{ (excess)} \longrightarrow I_2 + KCl$
 $I_2 + KI \longrightarrow KI_3 \text{ or } K^+ + I_3^-$
 $I_3^- + 2S_2O_3^{2^-} \longrightarrow S_4O_6^{2^-} + 3I^-$
using a starch indicator.

38.
$$O = C = O$$
 (linear)

41. Be
$$X$$
 Be X Be X X X X X X

The coordination number of Be in these chains is not exactly four. The deviations from the ideal tetrahedral angle for a four-coordinate Be atom

depend on the nature of the bridging group, and are related to the presence or absence of lone pairs on the bridging group.

45. KO_2 is a superoxide ($K^+O_2^-$), and the rest are peroxides.

- 47. Because of their degree of reactivity.
- **50.** The smaller the size of the cation, the greater is the lattice energy.
- **51.** The melting point of a metal depends on how closely packed a structure it has.
- **53.** The smaller the size of the cation, the greater is the hydration energy.
- 57. Pb as well as Al are amphoteric because they react with acids as well as alkalis to produce H_2 gas.
- 58. They are ionic salts in the fused state.
- **59.** The smaller the size of the cation, the greater is the hydration power. The corresponding chlorides form hydrated salts.
- 62. They are acidic.
- **63.** Alkaline earth metals have lower standard reduction potentials than alkali metals.
- 67. The ions of alkali metals have small radii.
- 74. KI is a reducing agent,

$$2I^- \rightarrow I_2 + 2e$$

and $Cr_2O_7^{2-}/H^+$, MnO_4^-/H^+ and Cu^{2+} are all oxidizing agents. So, I^- reduces them, and is itself oxidized.

Boron

• *Type 1* •

Choose the correct option. Only one option is correct.

1.	Which Lewis	n of the follow acid?	ving compor	ınds is ar	n important	catalyst as	well as a
	(a)	Al_2S_3		(b)	S_4N_4		
	(c)	N_2H_4		(d)	BF_3		
2.	Amor	phous boron	is prepared l	y heating	g B ₂ O ₃ with		
	(a)	Mn		(b)	Hg		
	(c)	Mg		(d)	SiO_2		
3.	Which	n of the follow	ving ions doe	es not exis	st in an aque	ous solutio	on?
	(a)	Pb ²⁺	(b) Sn ⁴⁺	(c)	B^{3+}	(d) Tl+	
4.		ng the follow pair effect?	ing, which s	hows the	e correct ord	ler of mag	gnitude of
	(a)	B < Al < Ga	< In < Tl	(b)	Ga > In > T	Cl < B > Al	
	(c)	Ga < In < Tl	< B < Al	(d)	Be \approx Al $>$ C	Sa > Tl > Ir	ı
5.	Boron	has an extre	mely high me	elting poi	nt because c	of	
	(a)	the strong va	an der Waals	forces be	etween its at	oms	
	(b)	the strong b	inding forces	in the co	valent polyı	mer	
	(c)	its ionic crys	tal structure				
	(d)	allotropy					
6.	Ortho	boric acid co	ntains				
	(a)	pyramidal B	O_3^{3-} units	(b)	linear BO ₃	units	
	(c)	T-shaped BC	O_3^{3-} units	(d)	triangular l	BO_3^{3-} units	i
		-		2-40	Ü		

Boron 2-41

- 7. A boron carbide rod is used in a nuclear reactor because boron (¹⁰B) has a very
 - (a) low area of cross-section for capturing neutrons
 - (b) high area of cross-section for capturing neutrons
 - (c) low area of cross-section for capturing protons
 - (d) high area of cross-section for capturing neutrinos
- 8. Which of the following statements is incorrect?
 - (a) Boron carbide is used as an abrasive.
 - (b) Boron is used to increase the hardenability of steel.
 - (c) Boron sesquioxide, B_2O_3 , is used in the manufacture of borosilicate glass.
 - (d) Orthoboric acid undergoes intramolecular hydrogen bonding.
- 9. On hydrolysis, diborane produces
 - (a) $H_3BO_2 + H_2O_2$

(b) $H_3BO_3 + H_2$

(c) $B_2O_3 + O_2$

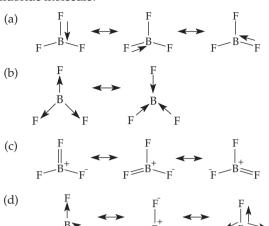
- (d) $H_3BO_3 + H_2O_2$
- 10. Which of the following statements is correct?
 - (a) Borosilicate glass is not heat resistant.
 - (b) Peroxoborate is stable to high heat.
 - (c) Borosilicate glass has a lower coefficient of thermal expansion and is easier to work with than normal soda glass.
 - (d) Soda-free glass fibre is made from boric oxide, sodium silicate and phosphorus.
- 11. Boron nitride is a
 - (a) white solid with a diamond-like structure
 - (b) slippery white solid with a layered structure similar to that of graphite
 - (c) covalent liquid and is structurally similar to carbon monoxide
 - (d) soft low-melting solid with a rock-salt-like structure
- 12. B_2H_6 reacts with $(CH_3)_3N$ to produce
 - (a) $BH_3^+N(CH_3)_3$

(b) $B_2H_6^+N(CH_3)_2CH_3\cdot BH_3$

(c) $(CH_3)_3 NBH_3$

- (d) $BH_3N(CH_3)_2CH_3BH_3$
- 13. The colour of the borax bead is due to the formation of a/an
 - (a) glass-like metal metaborote bead
 - (b) hard boric oxide crystal
 - (c) opaque metal hexaborate bead
 - (d) glass-like metal orthoborate bead

14. Which of the following structures correctly represents the boron trifluoride molecule?



15. By which of the following reactions is borazine prepared?

(a)
$$B_2H_6 + NH_3$$
 (excess) — low temperature

(b)
$$B_2H_6 + NH_3$$
 (excess) high temperature

(c)
$$B_2H_6 + NH_3 \xrightarrow{\text{(ratio 2NH}_3 : 1B_2H_6)}$$
 high temperature

- (d) None of these
- 16. Which of the following pairs contains structurally dissimilar substances?
 - (a) Borazine and benzene
 - (b) Diborane and hydrazine
 - (c) NaCl and NiO
 - (d) Graphite and cadmium iodide
- **17.** Which of the following statements is incorrect in the context of the B—F bond in BF₃?
 - (a) All the three B—F bond lengths are equal and each of them is shorter than the sum of the covalent radii of boron and fluorine.
 - (b) The bond energy of the B—F bond is very high, higher than for any other single bond.
 - (c) The unusual shortness and strength of the B—F bond may be explained by a $p\pi$ - $p\pi$ interaction between boron and fluorine atoms.

Boron 2-43

- (d) The unusual shortness and strength of the bonds may be explained by a $p\pi$ -d π interaction between the atoms of boron and fluorine.
- **18.** Which of the following statements is incorrect in relation to the structure of diborane?
 - (a) The terminal B—H bond distances are the same as the bond distances measured in compounds which are not electrondeficient, and which have no bridge structure.
 - (b) The terminal B—H bond is a 2-centre 3-electron bond.
 - (c) The terminal B—H bond is a 2-centre 2-electron bond.
 - (d) The bridge $B^{H_{B}}$ is a 3-centre 2-electron bond.
- 19. Which of the following statements is incorrect?
 - (a) B(OH)₃ partially reacts with water to form H₃O⁺ and [B(OH)₄]⁻, and behaves like a weak acid.
 - (b) B(OH)₃ behaves like a strong monobasic acid in the presence of sugars, and this acid can be titrated against an NaOH solution using phenolphthalein as an indicator.
 - (c) B(OH)₃ does not donate a proton and hence does not form any salt with NaOH.
 - (d) B(OH)₃ reacts with NaOH, forming Na[B(OH)₄].
- **20.** Which of the following statements is incorrect?
 - (a) B_2H_6 is not an electron-deficient molecule.
 - (b) The dipole moment of BF_3 is zero.
 - (c) The hydroboration of alkenes and subsequent oxidation with H₂O₂ and NaOH leads to the cis hydration of the carbon-carbon double bond with apparent anti-Markovnikov addition to give alcohols.
 - (d) BF₃ and BrF₃ molecules have different shapes.
- **21.** Which of the following structures does not represent the compound given in parentheses?

- 22. Among the boron halides, which is the strongest Lewis acid?
 - (a) BBr_3

(b) BCl₃

(c) BI₃

- (d) BF₃
- 23. Which of the following statements is correct in the context of diborane (B_2H_6) ?
 - (a) There are 12 valence electrons—three from each of the two boron atoms and six from the six hydrogen atoms.
 - (b) Two of the six hydrogen atoms form two bridges between two boron atoms.
 - (c) The two bridging hydrogen atoms are in a plane perpendicular to the rest of the molecule and prevent rotation between the two boron atoms.
 - (d) All of these.

• *Type 2* •

Choose the correct options. More than one option is correct.

- **24.** Which of the following compounds are acidic?
 - (a) $B(OH)_3$

(b) $Al(OH)_3$

(c) BF₃

(d) SiO_2

Boron 2-45

25.	Which	n of the following compounds:	react	with BF ₃ ?
		Ethers		B_2O_3
	(c)	Al_2Cl_6	(d)	NH_3
26.	Which	n of the following species have	tetra	hedral structures?
	(a)	B_2H_6	(b)	$B_3N_3H_6$
	(c)	BH_4^-	(d)	$AlCl_4^-$
27.	Which	n of the following compounds	react	with B ₂ H ₆ ?
	(a)	Cl_2	(b)	CO
	(c)	NH_3	(d)	(CH ₃) ₃ N
28.	Which	n of the following compounds	conta	ain boron?
	(a)	Borax	(b)	Colemanite
	(c)	Cristoballite	(d)	Kernite
29.	Which	n of the following compounds	can b	be made from borax?
	(a)	H_3BO_3	(b)	B_2O_3
	(c)	$LiBH_4$	(d)	BCl ₃
30.	Which	n of the following statements a	re co	rrect?
	(a)	Boric acid is a hydrogen-bone	led r	nolecule.
	(b)	Boric acid combines with Cubead test.	uO t	o give metaborate in the borax
	(c)	Al ₂ O ₃ is more acidic than B ₂ O	₃ .	
	(d)	Al ₂ O ₃ is amphoteric and B ₂ O ₃	is ac	cidic.
		Anstna	rc	

		1111300613	<u>'</u>	
1. d	2. c	3. c	4. a	5. b
6. a	7. b	8. d	9. b	10. c
11. b	12. c	13. a	14. a	15. c
16. b	17. d	18. b	19. c	20. a
21. d	22. a	23. d	24. a, c	25. a, b, c, d
26. c, d	27. a, b, c, d	28. a, b, d	29. a, b, c, d	30. a, b, d

Hints to More Difficult Problems

3. Boron forms covalent compounds.

8.

11.

- **12.** B₂H₆ is electron-deficient and (CH₃)₃N contains a lone pair of electrons. Hence both react together according to the Lewis acid-base concept.
- 22. Certain properties of the BX_3 adducts with donor molecules suggest that the donor-boron bonds may themselves increase in strength in the order $BF_3 < BCl_3 < BBr_3$.
- **25.** BF₃ is a Lewis acid and the other compounds are Lewis bases. Hence it reacts with all of them.
- 27. Same explanation as given for the answer to Q. 25
- **28.** The compounds shown in options (a), (b) and (d) are minerals containing boron; cristobalite has the formula SiO₂.

Carbon

• Type 1 •

Choose the correct option. Only one option is correct.

- 1. In the carbon family the elements other than carbon do not form $p\pi$ - $p\pi$ bonds because the atomic orbitals are too
 - (a) small and diffuse to undergo effective lateral overlap
 - (b) large and diffuse to undergo effective lateral overlap
 - (c) large and far too less diffuse to overlap linearly
 - (d) small to overlap both laterally and linearly
- 2. The stability of the tetrahalides of carbon decreases in the order

 - (a) $CF_4 > CCl_4 > CBr_4 > CI_4$ (b) $CCl_4 > CBr_4 > CF_4 > CI_4$
 - (c) $CI_4 > CCI_4 > CBr_4 > CF_4$ (d) $CBr_4 > CF_4 > CCI_4 > CI_4$
- 3. Graphite has a layered structure. The distance between the layers is
 - (a) 135 pm
- (b) 435 pm
- (c) 225 pm
- (d) 335 pm

- 4. The interlayer distance in graphite is
 - (a) very small, the layers being tightly packed
 - (b) many times larger than the covalent radius of carbon
 - (c) more than twice the covalent radius of carbon
 - (d) the same as the covalent radius of carbon
- 5. Carbon forms a large number of compounds due to its
 - (a) tetravalency
- (b) variable valency
- (c) large chemical affinity
- (d) property of catenation
- **6.** Which of the following statements is incorrect for graphite?
 - (a) Its density is lower than that of diamond.

- (b) It has a layered structure and the bonding between the layers is very weak.
- (c) Its layers are very tightly packed, almost without any space between them.
- (d) It cleaves easily between the layers which accounts for the remarkable softness of the crystals.
- 7. Which of the following statements is correct?
 - (a) Graphite is thermodynamically more stable than diamond.
 - (b) Diamond is thermodynamically more stable than graphite.
 - (c) Graphite has such a high thermodynamical stability that diamond spontaneously changes into graphite in ordinary conditions.
 - (d) Graphite and diamond have equal thermodynamic stability.
- 8. Among the following, the tendency for catenation decreases in the order
 - (a) C > Si > Ge = Sn > Pb
- (b) Si > C > Ge = Sn > Pb
- (c) C > Ge > Sn > Si > Pb
- (d) C > Pb > Sn > Ge > Si
- - (a) low temperature and very high pressure
 - (b) high temperature and low pressure
 - (c) high temperature and very high pressure
 - (d) low temperature and low pressure
- **10.** In the carbon family the melting points of the elements decrease on descending the group because the interatomic bonds become
 - (a) stronger as the size of the atom increases
 - (b) weaker as the size of the atom decreases
 - (c) stronger as the size of the atom decreases
 - (d) weaker as the size of the atom increases
- 11. Which of the following elements forms only three hydrides?
 - (a) C

(b) Si

(c) Ge

(d) Pb

- **12.** Which of the following is the most ionic?
 - (a) CCl₄

(b) PbCl₂

(c) PbCl₄

(d) SiCl₄

- 13. Carbon atoms in diamond are bonded to each other in a
 - (a) linear configuration

(b) planar configuration

(c) octahedral configuration

(d) tetrahedral configuration

Carbon 2-49

	(a)	Aluminium carbide as well as beryllium carbide produce methane gas on treatment with water.			
	(b)	On reacting with water, acetylene while magnesium c		um carbide (CaC_2) produces de (Mg_2C_3) gives propyne.	
	(c)	Calcium carbide has a lattice cell is elongated in one directi		ilar to that of NaCl, but the unit	
	(d)	All of these			
15.	CS ₂ re	eacts with Cl ₂ to produce			
	(a)	CCl ₄ and S ₂ Cl ₂	(b)	CCl ₂ and SCl ₄	
	(c)	CCl ₄ and SCl ₂	(d)	CCl ₄ and SCl ₆	
16.	Which	n of the following metal carbid	es m	ay be called a methanide?	
	(a)	CaC ₂	(b)	Mg_2C_3	
	(c)	Al_4C_3	(d)	BaC_2	
17.	Calciu	ım carbide is manufactured by	hea	ting	
	(a)	calcium with carbon			
	(b)	lime with coke			
	(c)	calcium with carbon monoxic	le		
	(d)	calcium chloride with coke			
18.	In car	bon-60 all carbon atoms are			
	(a)	sp ² -hybridized with a truncat	ted i	cosahedron shape	
	(b)	sp ³ -hybridized with a square	anti	prism shape	
	(c)	sp 2-hybridized with a diamor	nd sł	nape	
	(d)	(d) sp ² -hybridized with a graphite-like shape			
19.	Carbo	on-60 contains			
	(a)	20 pentagons and 12 hexagon	S		

14. Which of the following statements is correct?

(b) 12 pentagons and 20 hexagons(c) 30 pentagons and 30 hexagons(d) 24 pentagons and 36 hexagons

(a) rutile (TiO₂)

(c) silica (SiO₂)

(a) FeCO₃

(c) CaCO₃

metal on strong heating?

20. The crystal structure of calcium carbide resembles that of

21. Which of the following metal carbonates produces the corresponding

(b) fluorite (CaF₂)

(b) Li₂CO₃

(d) Ag_2CO_3

(d) rock salt (NaCl)

- 22. C_{60} can be regarded as a huge ball made up of
 - (a) several conjugated alkene units rather than an aromatic molecule
 - (b) graphite units
 - (c) several aromatic benzene molecules
 - (d) several tetrahedrons
- 23. The correct order of increasing carbon-oxygen bond length among CO, CO_3^{2-} and CO_2 is
 - (a) $CO_3^2 < CO_2 < CO$
- (b) $CO_2 < CO_3^{2-} < CO$
- (c) $CO < CO_3^{2-} < CO_2$
- (d) $CO < CO_2 < CO_3^{2-}$
- 24. CO_3^{2-} and SO_3^{2-} can be distinguished from each other by the production of CO₂ and SO₂ respectively upon using
 - (a) baryta water

- (b) limewater
- (c) acidified dichromate
- (d) sulphamic acid
- 25. On hydrolysis, magnesium carbide produces
 - (a) acetylene
- (b) butvne
- (c) propane (d) propyne
- **26.** Which of the following statements is incorrect?
 - (a) The carbon dioxide molecule behaves as a nonpolar molecule even though two of its resonating structures,

$$\stackrel{-}{O}$$
— $\stackrel{+}{C}$ = $\stackrel{+}{O}$ and $\stackrel{+}{O}$ = $\stackrel{-}{C}$ — $\stackrel{-}{O}$, are dipolar.

- (b) Carbon dioxide is the anhydride of the unstable dibasic acid $O=C(OH)_2$
- (c) The carbon dioxide molecule is linear because the carbon atom utilises its sp orbitals to form σ -bonds.
- (d) The carbon atom is sp ²-hybridized in the CO₂ molecule as well as the molecule of its hydrate H₂CO₃.
- 27. Which of the following statements is correct?
 - (a) An aqueous solution of sodium carbonate is neither acidic nor basic.
 - (b) An aqueous solution of sodium carbonate is alkaline because, being a base, the carbonate ion picks up the hydrogen ion from water releasing a hydroxide ion, making the solution alkaline.

$$CO_3^{2-} + H_2O \rightleftharpoons HCO_3^- + OH^-$$

(c) An aqueous solution of sodium carbonate is acidic because, being very electropositive, the sodium ion picks up the hydroxide ion from water to release a hydrogen ion, making the solution acidic

$$2Na^+ + 2H_2O \longrightarrow 2NaOH + 2H^+$$

(d) An aqueous solution of sodium carbonate is acidic because the carbonate ion reacts with water to form carbonic acid.

$$CO_3^{2-} + 2H_2O \rightleftharpoons H_2CO_3 + 2OH^{-1}$$

Carbon 2-51

28.		n of the follov CaNCN	_	compounds Al ₄ C ₃		ed as an abra CaC ₂		SiC
29		stalline hydra			` ,	-	()	
۷,۰	-	0°C and 1 at				a at		
		100°C and v		_		nraccura		
	(c)	100°C and 1						
	` '	0°C and ver			ssurc			
20						L		
30.		on monoxide : dries up	is po	isonous beca	use 1	t		
		reduces the	orga	nic matter of	tissu	ies		
	(c)						iency	y of oxygen in
	(d)	combines w	ith th	ne O ₂ present	in b	lood to form	CO_2	
31.	Solid	CO ₂ is produ	ced a	s white snov	v by			
	(a)	cooling the	gas b	elow its inve	ersior	n temperature	е.	
		cooling the	_	-		-		
		expanding t	_	_	_			
	(d)			oansion of the n orifice in t			s by	allowing it to
32.	In the	carbon mono	oxide	molecule, ca	arbor	n and oxygen	are	linked by
	(a)	only σ-bond	ls			only π -bond		
	(c)	σ- as well as	s π-bo	onds	(d)	no σ- and π -	-bon	ds
33.	Sodiu	m hydroxide	reac	ts with carbo	n mo	onoxide to pr	oduo	ce
	(a)	CO_2			(b)	(COONa) ₂		
	(c)	HOOC·COO	ONa		(d)	HCOONa		
34.	ordina	ary condition	s?	g solutions	can a	ıbsorb carboı	n mo	onoxide under
		Baryta wate		1				
		A dilute am Limewater	mon	ia solution				
	` '	An ammoni	acal	cuprous chlo	ride	solution		
35.		ructure of Si		1				
00.		that of a lay		lattice	(b)	linear		
		tetrahedral			(d)	square plana	ar	
36.	Which	n of the follow	ving	is an industr	ial fu	iel?		
	(a)	Water gas (CO+	H_2)	(b)	Producer ga	ıs (C	$O + N_2$

- (c) Coal gas $(CO + H_2 + CH_4 + CO_2)$
- (d) All of these
- 37. "Synthesis gas" is a mixture of
 - (a) carbon monoxide and hydrogen
 - (b) carbon monoxide and nitrogen
 - (c) carbon monoxide and steam
 - (d) methane and hydrogen
- 38. Carbon suboxide (C_3O_2) may be obtained by heating
 - (a) malonic acid with P_4O_{10}
- (b) malic acid with P_4O_{10}
- (c) oxalic acid strongly
- (d) maleic acid with P₄O₁₀
- **39.** In graphite, the hybridization state of each carbon atom and the π -bond order of each carbon-carbon bond are, respectively,
 - (a) sp and $\frac{1}{2}$

(b) sp² and $\frac{1}{3}$

(c) sp^3 and 1

- (d) sp² and $\frac{3}{2}$
- 40. The diamond molecule contains
 - (a) sp²-hybridized carbon atoms connected by single bonds
 - (b) sp²-hybridized carbon atoms connected by double bonds
 - (c) sp³-hybridized carbon atoms connected by single bonds
 - (d) sp³- and sp²-hybridized carbon atoms connected by single bonds

• *Type 2* •

Choose the correct options. More than one option is correct.

- 41. Graphite is a
 - (a) bad conductor of heat
 - (b) good conductor of electricity
 - (c) good conductor of heat
 - (d) good insulator
- 42. Which of the following are used as moderators in nuclear reactors?
 - (a) Graphite

(b) Paraffin

(c) Heavy water

- (d) None of these
- 43. When oxalic acid is heated with concentrated H₂SO₄, it produces
 - (a) CO

(b) SO₂ and CO₂

(c) CO and SO₃

(d) CO_2

44.	Metna	ine is obtained	by the hydrolysi	S OI			
	(a)	Be_2C		(b)	B_4C		
	(c)	Li_2C_2		(d)	Al_4C_3		
45.	Which	n of the followin	ng ions are regar	ded	as ionic ca	arbides?	
	(a)	C 4-		(b)	C_2^{2-}		
	(c)	C ₃ ⁴⁻		(d)	C_4^{3-}		
46.	Which	n of the followin	ng halides of car	bon	are solids?	•	
	(a)	CF_4		(b)	CCl_4		
	(c)	CBr_4		(d)	CI_4		
47.	Which	n of the followin	ng oxides of carb	on a	are stable?		
	(a)	CO		(b)	CO_2		
	(c)	C_3O_2		(d)	C_2O_3		
48.	 (a) The name buckminsterfullerene was given to C₆₀. (b) The comon name for C₆₀ is 'bucky ball'. (c) C₆₀ has a geodesic dome structure. (d) Solid C₆₀ has a cubic close-packed structure. 						
49.	Which	n of the followin	ng molecules ha	ve z	ero dipole	moment?	
	(a)	CS_2			CO_2		
	(c)	CCl ₂		(d)	CH_2Cl_2		
50.	Buckr	ninsterfullerene	e is prepared by				
	(a)	the pulsed lase	er vapourization	of §	graphite		
	(b)		arbon by resistiv		_		
	(c)		c discharge bet ium at 100 torr	wee	en carbon	electrodes in a tube	
	(d)	none of these					
			Answer	rs			
1. k)	2. a	3. d		4. c	5. d	
6. 0	2	7. a	8. a		9. c	10. d	
11. c		12. b	13. d		14. d	15. a	
16. c		17. b	18. a		19. b	20. d	
21. (_	22. a	23. d		24. c	25. d	
26. c	ı	27. b	28. d		29. a	30. c	

31. d	32. c	33. d	34. d	35. c
36. d	37. a	38. a	39. b	40. a
41. b, c	42. a, b, c	43. a, d	44. a, d	45. a, b, c
46. c, d	47. a, b, c	48. a, b, c, d	49. a, b	50. a, b, c

Hints to More Difficult Problems

- 7. The free energy of formation of graphite is 2.9 kJ mol⁻¹ lower than that of diamond at 300 K and 1 atm pressure.
- 9. Apply the Le Chatelier principle.
- 12. Apply Fajans' rules.
- **16.** Al_4C_3 reacts with water to produce methane (CH₄).

$$Al_4C_3 + 12H_2O \longrightarrow 4Al(OH)_3 + 3CH_4$$

Therefore, it may be called a methanide.

- **19.** Use Euler's formula for any C_n cluster (n even and greater than 22). Such clusters have at least one closed hollow cage consisting of 12 pentagons and (n 20)/2 hexagons. For C_{60} n = 60. Then (60 20)/2 = 20 (number of hexagons).
- **24.** SO₂ turns acidified K₂Cr₂O₇ solution green whereas CO₂ will have no effect on such a solution.
- 31. Consider the Joule-Thomson effect.

32.
$$: C \underset{\pi}{\overset{\pi}{=}} \sigma O :$$

38.
$$HO_2CCH_2CO_2H \xrightarrow{P_4O_{10}} O=C=C=C=O+2H_2O$$
Malonic acid $150^{\circ}C$ carbon suboxide

- **41.** The continuous π system in each layer of graphite makes it a good conductor of electricity.
- **42.** C_2O_3 does not exist.
- **49.** Both CS₂ and CO₂ have a linear structure.

The vector sum of the individual dipole moments, and hence the dipole moments of the molecules, are zero.

Silicon

• <u>Type 1</u> •

(b) tridynite

(d) carborundum

Choose the correct option. Only one option is correct.

1. Silicon carbide (SiC) is known as

(a) quartz

(c) corundum

2.	Very _l	pure silicon is prepared by		
	(a)	reducing pure silicon tetrachl	oride	e with magnesium
	(b)	decomposing K ₂ [SiF ₆]		
	(c)	heating SiO ₂ with KF		
	(d)	the electrolysis of SiO_2 in SiF_4		
3.	semice (a) (b) (c)	pure silicon is an insulator, lonductor when doped with a group 1 and a group 12 elemegroup 13 and a group 15 elemegroup 12 and a group 16 elemegroup 4 and a group 5 elemer	ent re nent i	respectively respectively
4.	Which	n of the following bonds has th	e hig	ghest bond energy?
		Si—C (b) Si—H	_	•
5.	Glass	is		
	(a)	a superheated liquid	(b)	a supercooled liquid
	(c)	an organosilicon polymer	(d)	a crystalline semisolid
6.	Glass	is soluble in		
	(a)	PbF ₄	(b)	$B(OH)_3$
	(c)	HF	(d)	concentrated HNO ₃
		2-55		

7.	Silicor	n shows a diagonal relation wi	th	
	(a)	magnesium	(b)	phosphorus
	(c)	carbon	(d)	boron
8.	Which	n of the following halides easil	y un	dergoes hydrolysis?
	(a)	AgCl	(b)	NF ₃
	(c)	SiCl ₄	(d)	CF_4
9.	Silicat	e minerals are classified accord	ding	to the manner of linking of
	(a)	SiO_4^{6-} tetrahedral units	(b)	SiO ₄ ⁴⁻ tetrahedral units
	(c)	$(Si_2O_7^{2-})_n$ units	(d)	$(SiO_3)_n^{4n}$ triangular units
10.	Which	n of the following is a purely a	cidic	oxide?
	(a)	SiO_2	(b)	SnO_2
	(c)	PbO	(d)	MnO_2
11.	dimer	nsional-network solid because		cule while SiO ₂ is a three-
	(a)	CO_2 molecules are held by s SiO_2 is an ionic solid	tron	g van der Waals forces whereas
	(b)	CO_2 has a structure like the structure	at o	f CaC_2 , and SiO_2 has a rutile
	(c)	bonds between the carbon an while silicon cannot form suc	d ox h do	due to the formation of double ygen atoms by 2p-2p π -bonding uble bonds with oxygen because ent; thus SiO_2 forms an infinite
	(d)			π -bonding and SiO_2 molecules thus SiO_2 forms an infinite
12.	Which	n of the following molecules is	coor	dinatively saturated?
	(a)	AlCl ₃	(b)	SiCl ₄
	(c)	CCl ₄	(d)	PbI_4
13.	Which	n of the following species is no	t coo	rdinatively saturated?
	(a)	SiF ₆ ²⁻	(b)	CH ₄
	(c)	PCl ₆	(d)	SnF_5^-
14.		a ground-glass stopper gets ning an NaOH solution. The r		k in the neck of a glass bottle n is that
	(a)	there are particles of dirt in be	etwe	en
	(b)	a solid silicate is formed in be	etwe	en by the reaction of the SiO ₂ of

glass with NaOH

Silicon 2-57

(d)	glass contains a boron compo the NaOH solution	ound	which forms a precipitate with
Which	n of the following orthosilicates	s is k	nown as willemite?
(a)	$Zn_3[SiO_4]_2$	(b)	$Mn_2[SiO_4]$
(c)	$W_2[SiO_4]$	(d)	$Zn_2[SiO_4]$
	is easily hydrolysed by wat olysis in water because	er w	rhereas CCl ₄ is stable towards
(a)	CCl ₄ exists as a molecule whe	reas	SiCl ₄ is ionic
(b)	the C—Cl bond is stronger th	an th	ne Si—Cl bond
(c)	silicon has a 3d orbital avait water whereas carbon has no		e for further coordination with bital for bonding
(d)	silicon is more electropositive	thai	n carbon
	on shows strong catenation ation because	whi	le silicon shows little or no
(a)	silicon is a metalloid and carb	on is	s a nonmetal
(b)	silicon forms ionic compour compounds	nds v	whereas carbon forms covalent
(c)	the Si—Si bond is stronger tha	an th	e C—C bond
(d)	the C—C bond is stronger that	n th	e Si—Si bond
Which	n of the following molecules is	not a	a donor?
(a)	$(C_2H_5)_2NH$	(b)	$(C_2H_5)_3N$
(c)	$(SiH_3)_3N$	(d)	C_2H_5OH
	ybridization state of nitrogen and N are	atom	s in the molecules (CH ₃) ₃ N and
(a)	respectively sp ³ and sp ²	(b)	respectively sp ² and sp ³
	both sp ²		both sp ³
	one pair of electrons in the ₃ N and (SiH ₃) ₃ N are accommod		rogen atoms of the molecules I respectively in the
(a)	sp ³ orbital and sp ² orbital	(b)	sp ³ orbital and p orbital
			p orbital in both the cases
The sl	hapes of the molecules of (CH_3) ₃ N a	and (SiH ₃) ₃ N are respectively
	pyramidal	, ,	1 ,
(b)	planar triangular		
(c)	planar triangular and pyrami	dal	
(d)	pyramidal and planar triangu	ılar	

(c) solid Na_2CO_3 is formed in between by reaction of the CO_2 of air

and NaOH

15.

16.

17.

18.

19.

20.

21.

- 22. The oxidation numbers of Si in $Si_3O_9^{6-}$ and $Si_6O_{18}^{12-}$ respectively are
 - (a) -4, -4

(b) +4, +2

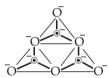
(c) +4.+4

- (c) +4, +6
- 23. Which of the following statements is correct for silicon? It
 - (a) forms molecular halides that are not hydrolysed.
 - (b) forms strong but unconjugated multiple bonds of the $p\pi$ -d π variety, especially with O and N.
 - (c) does not undergo coordination number expansion
 - (d) forms an oxide (SiO₂) that is amphoteric and has a GaAs structure.
- 24. Which of the following represents a pyrosilicate structure?

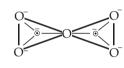
(a)



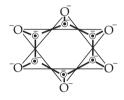
(b)



(c)



(d)



- **25.** Which of the following statements is incorrect?
 - (a) The octahedral SiF_6^{2-} ion is the only halogenocomplex of silicon, and the bonding in it involves sp^3d^2 hybridization.
 - (b) Fluorosilicic acid, H₂SiF₆, known only in solution, is a strong acid.
 - (c) Silicon is more electropositive than carbon.
 - (d) The formula of tetramethyl silane is $(CH_3)_4SiH_4$.
- 26. Silicones are a group of organosilicon polymers containing
 - (a) Si—O—Si linkages
- (b) O-Si-O linkages
- (c) Si—C—Si linkages
- (d) Si—Si—O linkages
- 27. Which of the following statements is incorrect?
 - (a) The hydrolysis of (CH₃)₃SiCl gives a disiloxane.
 - (b) The hydrolysis of (CH₃)₂SiCl gives a chain compound.
 - (c) The hydrolysis of CH₃SiCl₃ gives a cross-linked polymer.
 - (d) The hydrolysis of $(CH_3)_2SiCl_2$ gives $(CH_3)_2SiO_2$.

Silicon 2-59

28.	Which silicat		ıs do	not represent cyclic and chain
	(a)	$Si_2O_7^{2-}$ and $(SiO_3)_n^{2n-}$	(b)	$S_3O_9^{6-}$ and $(Si_4O_{11})_n^{6n-}$
	(c)	$Si_2O_7^{2-}$ and $(Si_2O_5)_n^{2n-}$	(d)	$Si_2O_7^{7-}$ and $(SiO_3)_n^{2n-}$
29.	Which	n of the following statements is	inco	orrect in the context of silicones?
	(a)	They are more stable to heat t	han	other polymers.
	(b)	•	ellen	t, are good electrical insulators,
	(c)	The Si—O bond energy is hig		01 1
	` '	The Si—O bond energy is low		
30.				NaOH in the presence of air to
	produ			
	(a)	$Na_2SiO_3 + H_2$	(b)	$Na_2SiO_3 + Na_2CO_3$
	(c)	$Na_2SiO_2 + H_2$	(d)	$Na_2SiO_4 + O_2$
		• <u>Type</u>	<u>2</u> •	
Choo	se the o	correct options. More than one	opti	ion is correct.
31.	Pure S	SiO ₂ occurs as		
	(a)	quartz	(b)	cristobalite
	(c)	colemanite	(d)	siderite
32.	Which	n of the following have cyclic s	ilicat	te structures?
	(a)	Si ₃ O ₉ ⁶⁻	(b)	SiO ₄ ²⁻
	(c)	$Si_6O_{18}^{12-}$	(d)	$(Si_4O_{11}^{6-})_n$
33.	Which	n of the following elements for	m hy	drides?
	(a)		-	Ge
	(c)	Pb	(d)	
34.	Which	n of the following halides are h	ydro	plysed by water?
	(a)	SiCl ₄	(b)	CCl ₄
	(c)	AlCl ₃	(d)	CF_4
35.	Which	n of the following molecules ha	ive a	V-shaped structure?
		SnCl ₄		SnCl ₂
		COS		PbCl ₂

26. a

31. a, b

27. c

32. a, c

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2. a	3. b	4. c	5. b
7. d	8. c	9. b	10. a
12. c	13. d	14. b	15. d
17. d	18. c	19. a	20. b
22. c	23. b	24. c	25. d
	7. d 12. c 17. d	2. a 3. b 7. d 8. c 12. c 13. d 17. d 18. c	7. d 8. c 9. b 12. c 13. d 14. b 17. d 18. c 19. a

28. b

Hints to More Difficult Problems

33. a, b, c, d

6. Glass contains silica. It dissolves in HF forming $H_2[SiF_6]$ (hydrofluorosilicic acid).

29. d

34. a, c

30. c

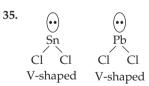
35. b, d

- SiCl₄ is readily hydrolysed. Si can use a d orbital to form a five-coordinate intermediate to produce SiCl₆².
- 12. CCl₄ does not undergo coordination number expansion.
- 18. Trisilylamine or $(SiH_3)_3N$ involves $p\pi$ -d π bonding, because the bond forms between full p orbitals and an empty d orbital. This shortens the N–Si bond lengths. Since nitrogen does not have a lone pair of electrons, the molecule is not a donor.

22. In Si₃O₉⁶⁻,
$$3x - 18 = -6 \Rightarrow 3x = 12 \Rightarrow x = +4$$

In Si₆O₁₈¹²⁻, $6x - 36 = -12 \Rightarrow 6x = 24 \Rightarrow x = +4$

34. Both use a d orbital and undergo coordination-number expansion.



Nitrogen and Phosphorus

• *Type 1* •

Choose the correct option. Only one option is correct.1. The nitrogen molecule is isoelectronic with

		•		
	(a)	CO^- , CN^+ and NO_2^+	(b)	CO, CN ⁻ and NO ⁺
	(c)	CO^+ , N_2O and O_2^{2-}	(d)	O_2^+ , O_2^- and CO^+
2.	N ₂ is p	prepared commercially by		
	(a)	the fractional distillation of lice	quefi	ed air
	(b)	heating ammonium dichroma	ite	
	(c)	heating a mixture of ammonia	um c	hloride and sodium nitrite
	(d)	the serpeck process		
3.	Which	n of the following is an ionic co	mpo	ound?
		NI_3 (b) NF_3	-	NCl ₃ (d) BiF ₃
4.	Active	e nitrogen can be made by passir	ng an	electric spark through N ₂ gas at
	(a)	very low pressure (2 mm of H	Ig)	
	(b)	high pressure		
	(c)	very low temperature		
	(d)	ordinary pressure		
5.	Which	n of the following properties do	oes n	itrogen not exhibit?
	(a)	Low boiling point	(b)	Hydrogen bonding
	(c)	Catenation	(d)	Supporter of life
6.	Nitro	us oxide is prepared by the the	rmal	decomposition of
	(a)	dinitrogen tetroxide	(b)	ammonium nitrate
	(c)	ammonium nitrite	(d)	diazomethane
		2-61		

- 7. Which of the following statements is correct in the context of the N₂O molecule?
 - (a) The N—N bond is longer than the N—O bond.
 - (b) The N—N bond is as long as the N—O bond.
 - (c) The N—N bond has no π character.
 - (d) The N—N bond is shorter than the N—O bond.
- 8. Which of the following pairs consists of species of equal bond order?
 - (a) N₂ and CO

(b) N₂ and CN⁺

(c) CO^- and N_2^+

- (d) N_2 and NO^+
- 9. Which of the following statements is incorrect for nitric oxide?
 - (a) It is an anhydride of nitrous acid.
 - (b) It is a paramagnetic molecule.
 - (c) It does not dimerize.
 - (d) It has a dipole moment of 0.12 D.
- 10. The bond length of N—O is 1.15 Å, which is intermediate between those of a
 - (a) single bond and a double bond between N and O
 - (b) double bond and a triple bond between N and O
 - (c) carbon-carbon single bond and a carbon-carbon double bond
 - (d) carbon-carbon double bond and a carbon-carbon triple bond
- 11. Nitric oxide has 11 outer electrons. Among these, one unpaired electron occupies
 - (a) an antibonding π_{2p} orbital (b) an antibonding π_{2s}^* orbital
 - (c) An antibonding π_{2p}^* orbital (d) a bonding π_{2p} orbital
- 12. The resonance structure of NO is represented by
 - (a) $\vec{N} \vec{O} = \vec{N} = \vec{O}$ (b) $\vec{N} = \vec{O} =$
 - (c) $: \overset{+}{N} = \overset{-}{O}: \longrightarrow : \overset{+}{N} = \overset{-}{O}: \longrightarrow : \overset{+}{N} \overset{+}{N} \overset{-}{N} \overset{+}{N} \overset{-}{N} \overset{+}{N} \overset{-}{N} \overset{+}{N} \overset{-}{N} \overset{+}{N} \overset{+}$

- 13. The NO molecule
 - (a) often acts as a one-electron donor, in contrast to most ligands which donate two electrons
 - (b) often acts as a two-electron donor as is true for most ligands
 - (c) often acts as a three-electron donor, in contrast to most ligands which donate two electrons
 - (d) does not act as a donor

- 14. N_2O_3 is
 - (a) an acidic oxide, and the anhydride of HNO₂
 - (b) an acidic oxide, and the anhydride of H₂N₂O₂
 - (c) a neutral oxide, and the anhydride of HNO₃
 - (d) a basic oxide, and the anhydride of HNO₂
- 15. N_2O_3 can be obtained by cooling (below $-30^{\circ}C$) an equimolecular mixture of
 - (a) N₂O₅ and NO

(b) NO and NO₂

(c) N_2 and O_3

- (d) NO₂ and O₃
- **16.** Dinitrogen tetroxide (N_2O_4) has
 - (a) two unpaired electrons and is paramagnetic
 - (b) two unpaired electrons and is diamagnetic
 - (c) one unpaired electron and is paramagnetic
 - (d) no unpaired electron and is diamagnetic
- 17. Which of the following oxides of nitrogen is a mixed anhydride of two acids?
 - (a) N_2O

(b) NO

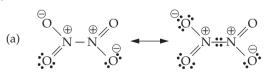
(c) N_2O_4

- (d) NO_3
- 18. Which of the following molecules does not undergo self-ionization?
 - (a) N_2O_4 (l)

(b) $H_2O(l)$

(c) NH₃(l)

- (d) $HNO_3(l)$
- 19. Which of the following statements is correct for the N₂O₄ molecule?
 - (a) The N—N bond is very short and strong.
 - (b) The N—N bond is very short and weak.
 - (c) It is a neutral oxide.
 - (d) It is a paramagnetic molecule.
- **20.** Which of the following structures correctly represents the resonance hybrid of N_2O_4 ?



(b)
$$N-N$$
 N $N-N$ $N-N$ N N N N N

$$(d) \quad \vdots \overset{\circ}{\circ} \oplus \overset{\ominus}{\circ} \overset{\circ}{\circ} \vdots \qquad \vdots \overset{\circ}{\circ} \overset{\ominus}{\circ} \oplus \overset{\ominus}{\circ} \overset{\circ}{\circ} \overset{\circ}{\circ} \vdots \\ \vdots \overset{\circ}{\circ} \overset{\circ}$$

- 21. During the formation of the N₂O₄ dimer from two molecules of NO₂, the odd electrons, one in each of the nitrogen atoms of the NO₂ molecules, get paired to form a
 - (a) weak N—N bond, two N—O bonds become equivalent and the other two N—O bonds become nonequivalent
 - (b) weak N—N bond and all the four N—O bonds become equivalent
 - (c) weak N—N bond and all the four N—O bonds become nonequivalent
 - (d) strong N—N bond and all the four N—O bonds become equivalent
- 22. Dinitrogen pentoxide (N₂O₅), a colourless solid, is prepared by
 - (a) heating NH₄NO₂ with an excess of oxygen
 - (b) dehydrating HNO₃ with CaO
 - (c) dehydrating HNO₃ with P₄O₁₀
 - (d) heating a mixture of HNO₂ and Ca(NO₃)₂
- **23.** Arrange NO₂⁺, NO₂ and NO₂⁻ in order of increasing N—O bond length.
 - (a) $NO_2^+ < NO_2 < NO_2^-$
- (b) $NO_2^- < NO_2^+ < NO_2$
- (c) $NO_2 < NO_2^- < NO_2^+$
- (d) $NO_2^+ = NO_2^- < NO_2$
- **24.** Which among the following is the least basic?
 - (a) NCl₃

(b) NBr₃

(c) NI₃

(d) NF₃

- 25. Which among the following is a deliquescent ionic solid?
 - (a) BrF₅

(b) S_4N_4

(c) N_2O_5

(d) NO

- **26.** Which of the following statements is correct?
 - (a) NO₂ is linear.
 - (b) NO₂⁺ is pyramidal.
 - (c) N_2O_5 is represented as $NO_2^+NO_3^-$.
 - (d) N₂O₅ reacts with concentrated H₂SO₄ to produce NO₂⁺HSO₄⁻.
- 27. Which of the following structures is the correct representation of $N_2O_5\, \rm in$ the gaseous phase?

(a)
$$N - O - N$$
 (b) $N - O - N$ (c) $N - O - N$ (c)

- **28.** An aqueous solution of nitrous acid (HNO₂), free of salts, can be obtained from the reaction
 - (a) $Ba(NO_2)_2 + H_2SO_4 \longrightarrow$
 - (b) $NaNO_2 + H_2SO_4 \xrightarrow{cold}$
 - (c) $NH_4NO_2 + H_2SO_4 \xrightarrow{\Delta}$
 - (d) $KNO_3 + HNO_3 \longrightarrow$
- 29. In which of the following reactions does HNO₂ act as an oxidizing agent?
 - (a) $Cr_2O_7^{2-} + H^+ + NO_2^- \longrightarrow Cr^{3+} + NO_3^- + H_2O$
 - (b) $Cl_2 + HNO_2 \longrightarrow HCl + HNO_3$
 - (c) $SO_2 + HNO_2 \longrightarrow H_2SO_4 + NO_4$
 - (d) $O_3 + H^+ + NO_2^- \longrightarrow NO_3^- + O_2$
- 30. In which of the following reactions does HNO₂ act as a reducing agent?
 - (a) $FeSO_4 + H_2SO_4 + HNO_2 \rightarrow Fe_2(SO_4)_3 + NO + H_2O$
 - (b) $KMnO_4 + H_2SO_4 + HNO_2 \rightarrow K_2SO_4 + MnSO_4 + HNO_3 + H_2O$
 - (c) $KI + H_2SO_4 + HNO_2 \rightarrow K_2SO_4 + NO + I_2 + H_2O$
 - (d) $SnCl_2 + HCl + HNO_2 \rightarrow SnCl_4 + NO + H_2O$
- 31. HNO₂ reacts with secondary amines to produce
 - (a) diazo salts
- (b) nitrile

(c) nitrogen

(d) nitrosoamines

32. Urea reacts with HNO₂ to produce

(a) N_2O_4

	(a)	CO, N ₂ and	H_2O		(b)	CO ₂ and N ₂		
	(c)	CO ₂ , NO an	d NO	O_2	(d)	CO, N ₂ O an	d H ₂	O
33.	In the	e nitrite ion NO ₂ , the bond orders of the two N—O bonds are						
	(a)	2.0 and 1.0			(b)	both 1.5		
	(c)	2.5 and 1.5			(d)	both 2.5		
34.	In din	itrogen pento	oxide	e, the central	NON	I bond angle	is clo	ose to
	(a)	180°			(b)	120°		
	(c)	90°			(d)	134°		
35.	N_2O_5	reacts with ic	odine	to produce				
	(a)	I_2O_6			(b)	IO_3		
	(c)	I_2O_5			(d)	$I_2O_5 + N_2O$		
36.	The fi	rst ionization	enei	rgy of NO is	less t	han that of		
	(a)	N_2			(b)	O_2		
	(c)	Xe			(d)	all of these		
37.	In the	nitrous oxid	e mo	lecule, the N	—N	as well as the	e N—	O bond have
	(a)	some doubl	e bor	nd character				
		some triple						
		a d orbital ii		ed in bondir	ng			
	(d)	none of thes	se					
38.	Which	n of the follow	wing	pairs has a p		-triangle stru		?
	(a)	NO_3^- and N	H_3		(b)	NO ₂ ⁺ and BI	F ₃	
	(c)	NO_3^- and CO_3^-	O_3^{2-}		(d)	NO_3^- and Xe	eO_3	
39.	In wh	_	ollow	ing reactions	wil	l HNO ₃ not a	act a	s an oxidizing
	(a)	$HNO_3 + H_2$	SO ₄ -	\longrightarrow				
	(b)	$HNO_3 + FeS$	5O ₄ +	$H_2SO_4 \longrightarrow$				
	(c)	KI + HNO ₃	\longrightarrow					
	(d)	Au + HNO ₃	. —	>				
40.	Which HNO		ving	metals are pa	assiv	e to treatmen	t witl	h concentrated
		Ni	(b)	Cr	(c)	Co	(d)	All of these
41.	The re	eduction of N	IO ₃ i	n an alkaline	med	lium with zin	nc giv	res

(b) N_2O_5 (c) NH_3

(d) NH₄⁺

42.	vv nici	n of the following factors is res	pons	ible for the passivity of	or a metar:
	(a)	The formation of a self-protect the metal over its surface,	ting	coating of the basic ca	rbonate of
	(b)	The formation of a thin lay surface, preventing the action			etal on its
	(c)	The formation of a thin layer surface, preventing the action			etal on its
	(d)	None of these		-	
43.	The re	eaction of HNO_3 with P_4O_{10} give	es		
	(a)	N_2O_5	(b)	NO ₂	
	(c)	N_2O_4	(d)	N_2O	
44.	The n	itrogen fixation of atmospheric	nitr	ogen utilises the	
	(a)	Fischer–Tropsch process	(b)	Frasch process	
	(c)	Haber process	(d)	Solvay process	
45.	The ca	atalytic oxidation of NH ₃ yields	S		
	(a)	N_2O_5	(b)	NO	
	(c)	NO_2	(d)	N_2	
46.	A goo	od yield is obtained when HNC) ₃ is 1	manufactured by the	
	(a)	Ostwald process		Haber process	
	(c)	Birkeland–Eyde process	(d)	Hall process	
47.	Conce	entrated HNO ₃ acts as an oxidi	zing	agent with	
	(a)	P ₄ (b) NaBr	(c)	Sn (d) all o	of these
48.		eaction of zinc with very dilute			ion of
		N_2O_5		N_2O	
	(c)	NH_4NO_3	(d)	NH_4NO_2	
49.		HNO_3 is a colourless liquid, bun. This is due to the slight deco			ns slightly
		NO and NO ₂	_	NO ₂ and O ₂	
	(c)	NO and O ₂	(d)	N_2O_5 and O_2	
50.	The a	ctive species in the nitration of	an a	romatic organic comp	ound is
	(a)	NO_2	(b)	NO_3^-	
	(c)	NO ₂ ⁺	(d)	NO_2^-	
51.	Gold	and platinum react with aqua	regia	to form soluble comp	lexes. The
	comp	lexes with oxidation number +	3 and	d +4, respectively, are	
		$H_2[AuCl_5]$ and $H_2[PtCl_6]$		H[AuCl ₄] and H ₂ [PtC	0-
	(c)	$H[AuCl_4]$ and $H_4[PtCl_8]$	(d)	H ₃ [AuCl ₆] and H[PtC	Cl ₅]

52.		n of the following nitrates does $Zn(NO_3)_2$		give O ₂ on heating? KNO ₃
		$Pb(NO_3)_2$		NH ₄ NO ₂
	. ,		` ′	
53.		0		tes gives two gaseous products?
		NH ₄ NO ₃	` '	NH ₄ NO ₂
	(c)	$Cu(NO_3)_2$	(a)	NaNO ₃
54.		IO_2^- ion is determined volumet		
	(a)		` '	KClO ₃
	(c)	$Na_2S_2O_3$	(d)	CuSO ₄
55.	Which	n of the following species is pla	nar?	
	(a)	NO_3^-	(b)	N_2O_4
	(c)	N_2O_5	(d)	All of these
56.	The n	itrite ion is angular, and in it		
	(a)	the nitrogen atom has three spelectrons	o ² or	bitals and contains a lone pair of
	(b)	the nitrogen atom has three selectron	p ² o	rbitals, and contain an unpaired
	(c)	the nitrogen atom has three spelectrons	p ork	pitals, and contains a lone pair of
	(d)	the nitrogen atom has three sof electrons	sp ³ c	orbitals, and contains a lone pair
57.	Each o	of the N—O bonds in the NO ₃	ion l	has a bond order of
		0.33		1.33
	(c)	1.50	(d)	1.00
58.	Upon reacting with tin, hot concentrated HNO ₃ produces			
	(a)	$Sn(NO_3)_2$	(b)	H_2SnO_3
	(c)	SnO ₂	(d)	$Sn(NO_3)_4$
59.	Calcium reacts with nitrogen gas to produce a white ionic solid, which on hydrolysis gives the gas (or gases)			
	-	NH ₃		N_2O
	(c)	NO_2	(d)	NH ₃ and NO
60.	manu	n of the following catalyst facture of ammonia by the Hal Finely divided platinum toge	oer p	
	(44)	arriaca pianimini toge		a mener promoter

Finely divided nickel together with a platinum promoter Finely divided iron together with a molybdenum promoter

(d) Finely divided palladium together with a zinc promoter

61.	Which heatin	_	ds d	oes not produce ammonia on
	(a)	NH ₄ Cl	(b)	$(NH_4)_2SO_4$
	(c)	NaNH ₄ HPO ₄ ·4H ₂ O	(d)	NH ₄ NO ₂
62.	For th	e catalytic oxidation of ammo	nia ii	nto nitric oxide by the oxygen of
	(a)			e reaction, and thereafter the ed at the requisite level for the
	(b)	no heat is required to initi exothermic	ate	the reaction as the reaction is
		efficient cooling is required at no heat is required to carry or		0 1
63.	Which	n of the following gases can be	liqu	efied easily?
	(a)	He	` '	NH_3
	(c)	H_2	(d)	H_2O
64.	(a) (b) (c)	ation of sodium in liquid amm solvated electrons and solvate solvated amide ions solvated azide ions solvated sodium atoms	_	and the second s
65.	Amm	onia gas is dried over		
	` '	P_4O_{10}	` '	CaO
	(c)	H_2SO_4	(d)	CaCl ₂
66.	Which	n of the following cations does	not	form a complex with ammonia?
	(a)	Co ³⁺	(b)	Ni ²⁺
	(c)	Pb ²⁺	(d)	Zn^{2+}
67.		n of the following compounds an ammonia solution?	give	es a black precipitate on reacting
	(a)	$ZnSO_4$ (b) Hg_2Cl_2	(c)	FeCl ₃ (d) NiSO ₄
68.	Hydra	azine is manufactured by the		
		Raschig process		Deacon process
		Dow process		Diazo process
69.		onia reacts with NaOCl to pro		
		N ₂ H ₄ , NaCl and N ₂ O N ₂ H ₄ and NH ₂ Cl		NO ₂ , HN ₃ and NH ₂ Cl N ₂ H ₄ and NaCl
	(c)	1 N21 14 alla 1 N1 12C1	(u)	1 V21 14 allu 1 VaC1

- 70. Sodium reacts with liquid ammonia to produce
 - (a) $NaNH_2 + N_2 + H_2$
- (b) $NaNH_2 + H_2$
- (c) $NaNH_2 + N_2H_4$
- (d) $NaNH_2 + N_2$
- 71. By reacting with which of the following reagents does ammonia produce a brown precipitate?
 - (a) Schiff's reagent
- (b) Fehling's solution
- (c) Grignard reagent
- (d) Nessler's reagent
- 72. Sodium azide is prepared by the reaction between
 - (a) N_2H_4 and H_2

- (b) NH₃ and NaOCl
- (c) N₂O and NaNH₂
- (d) NH₂OH and N₂H₄
- 73. Hydroxylamine is prepared by reducing nitrites with
 - (a) $C_6H_5NO_2$
 - (b) SO₂ in the presence of NaHSO₃
 - (c) SO₂ in the presence of Na₂SO₄
 - (d) Na₂S₂O₃
- 74. Among the following, which is the weakest base?
 - (a) NH₂
- (b) N_2H_4
- (c) NH₂OH (d) C₂H₅NH₂
- **75.** The resonance structure of the azide ion is

(a)
$$: \stackrel{\cdot}{N} = \stackrel{\cdot}{N} = \stackrel{\cdot}{N} : \stackrel{\cdot}{N} = \stackrel{\cdot}{N} : \stackrel{\cdot}{N} \stackrel{2-}{\longrightarrow} : \stackrel{\cdot}{N} - \stackrel{2+}{N} = \stackrel{2-}{N} : \stackrel{\cdot}{N} : \stackrel{N} : \stackrel{\cdot}{N} : \stackrel{\cdot}{N} : \stackrel{\cdot}{N} : \stackrel{\cdot}{N} : \stackrel{\cdot}{N} : \stackrel{\cdot}{N} :$$

(b)
$$: N = N = N : \longrightarrow : N = N : N : N : N = N : N : N : N = N$$

(c)
$$: \stackrel{\oplus}{N} - \stackrel{\ominus}{N} = \stackrel{\ominus}{N} : \longrightarrow : \stackrel{\bullet}{N} - \stackrel{\bullet}{N} = N$$
:

- **76.** The azide ion is
 - (a) asymmetrical and bent
- (b) isoelectronic with O₃
- (c) symmetrical and linear
- (d) pyramidal

- 77. The azide ion has
 - (a) 20 outer electrons and is isoelectronic with Br₂O
 - (b) 18 outer electrons and is isoelectronic with NO₂⁻
 - (c) 16 outer electrons and is isoelectronic with CO₂
 - (d) 14 outer electrons and is isoelectronic with H₂O₂

78. On combustion, hydrazine gives

		N_2		N_2O_5				
	(c)	N_2O_4	(d)	NH ₂ OH				
79.	Which (a)	The addition of the extra electhat one electron must occup	of the following statements is correct for HN ₃ (hydrogen azide)? The addition of the extra electron from the hydrogen atom means that one electron must occupy an antibonding molecular orbital, and hence the lengths of the two N—N bonds are different.					
	(b)	It uses eight electrons for sign	na bo	onding.				
	(c)	It uses six electrons for pi bon	ding	5 .				
	(d)	The bond orders of the two is 3.0.	nitro	gen-nitrogen bonds are 2.5 and				
80.	Which	n of the following polymers are	obt	ained by heating E-caprolactum?				
	(a)	Nylon-6	(b)	Nylon-66				
	(c)	Polyvinyl chloride	(d)	Teflon				
81.		n of the following molecules is the tendency to act as a donor?		le, not hydrolysed by water, and				
	(a)	NBr ₃	(b)	NCl ₃				
	(c)	NF ₃	(d)	NBr ₃				
82.		is hydrolysed by water to prod NH_2NH_2 and HCl		NH ₂ Cl and HOCl				
	(c)	NH ₄ ⁺ OH ⁻ and HOCl	(d)	NH ₂ OH and HOCl				
83.	(a) (b) (c)	gen is unable to form pentahali presence of 2s and 2p orbitals absence of 3d orbitals absence of 3p and 3d orbitals absence of 3s, 3p and 3d orbit		because of the				
84.	Which	n of the following molecules is	plan	ar around nitrogen?				
	(a)	$N(SiH_3)_3$ (b) $Si(NH_3)_3$	(c)	$P(NH_3)_3$ (d) $C_6H_5N(CH_3)_2$				
85.	percer (a)	ng the following, which is intage of nitrogen and is easily $CaCN_2$ KNO_3	assii (b)	fertilizer that has the highest milated in soil? (NH ₄) ₂ SO ₄ Urea				
0.5			` ′					
86.	prepa	red by the reaction between		$(H_2PO_4)_2$ and $7CaSO_4$, and is				
	(a)	$Ca_3(PO_4)_2 \cdot CaF_2$ and H_3PO_4	(b)	$Ca_3(PO_4)_2 \cdot CaCl_2$ and H_2SO_4				
	(c)	Ca ₃ (PO ₄) ₂ ·CaF ₂ and HCl	(d)	Ca ₃ (PO ₄) ₂ ·CaCl ₂ and CaSO ₄				

87. Triple superphosphate is represented by the formula $4Ca(H_2PO_4)_2$, and is produced by the reaction between
(a) $3Ca(H_2PO_4)_2$ and $NaHF_2$
(b) $4Ca_3(PO_4)_2$ and $CaCl_2$
(c) $Ca_3(PO_4)_2 \cdot CaF_2$ and $6H_3PO_4$
(d) AlPO ₄ and CaH ₂ PO ₄
88. White phosphorus is more reactive than the nitrogen molecule because the

- (a) electronegativity of phosphorus is low.
- (b) ionization energy of phosphorus is greater than that of N₂
- (c) P—P bond in phosphorus is weaker than the N≡N bond in nitrogen
- (d) P—P—P bond angle is 120° whereas N₂ is linear
- 89. The most reactive and least reactive forms of phosphorus are respectively
 - (a) red and white phosphorus
 - (b) white and black phosphorus
 - (c) scarlet and red phosphorus
 - (d) white and red phosphorus
- 90. In which of the following properties does white phosphorus resemble red phosphorus?
 - (a) Burning in the presence of air
 - (b) Solubility in an organic solvent
 - (c) Fluorescence and phosphorescence
 - (d) Reaction with concentrated NaOH to produce PH₃
- 91. White phosphorus reacts with calcium to form a certain compound which, on hydrolysis, produces
 - (b) P_2H_4 (a) PH₃
- (c) P_4O_6
- (d) P_4O_{10}

- 92. On hydrolysis, PCl₅ would produce
 - (a) HPO₃
- (b) H₃PO₄
- (c) H₃PO₃
- (d) $H_4P_2O_7$

- 93. In the solid state, PCl₅ exists as
 - (a) $[PCl_4]^-$ and $[PCl_6]^+$ ions
- (b) covalent PCl₅ molecules only
- (c) $[PCl_4]^+$ and $[PCl_6]^-$ ions
- (d) $[PCl_3]^{2-}$ and $[PCl_5]^{2+}$ ions
- 94. Which of the following pentahalides is highly reactive and unstable, and is not possible to isolate?
 - (a) PCl₅

(b) SbCl₅

(c) AsCl₅

(d) PBr₅

- 95. PCl₅ reacts with NH₄Cl to form
 - (a) phosphonium chloride and NH₃
 - (b) phosphonitrile chloride polymers (NPCl₂),
 - (c) phosphorus triammine and Cl₂
 - (d) phosphorus pentammine and N₂
- **96.** In P_4O_{10} , the
 - (a) second bond in P=O is formed by $p\pi$ -d π back bonding
 - (b) P=O bond is formed by $p\pi$ - $p\pi$ bonding
 - (c) P=O bond is formed by $d\pi$ - $d\pi$ bonding
 - (d) P=O bond is formed by $d\pi$ - $d\pi$ - 3σ back bonding
- 97. Phosphorus is obtained by the reduction of phosphate rock using
 - (a) coke and silica at high temperature
 - (b) Al at high temperature
 - (c) Fe₂O₃ and coke at high temperature.
 - (d) silica at high temperature
- 98. The basic character of NH₃, PH₃, AsH₃ and SbH₃ decreases in the order
 - (a) $SbH_3 > PH_3 > AsH_3 > NH_3$
 - (b) $NH_3 > PH_3 > AsH_3 > SbH_3$
 - (c) $NH_3 > SbH_3 > PH_3 > AsH_3$
 - (d) $AsH_3 > PH_3 > SbH_3 > NH_3$
- 99. In the reaction

$$P_4 + 3KOH + 3H_2O \longrightarrow 3KH_2PO_2 + PH_3$$

phosphorus is

- (a) only oxidized
- (b) only reduced
- (c) oxidized as well as reduced
- (d) neither oxidized nor reduced
- **100.** The number of short P—O bonds in P_4O_{10} is
 - (a) one

(b) two

(c) three

- (d) four
- **101.** Arrange ZnO, MgO, P₄O₆ and SO₃ in order of increasing acidic strength.

- 102. Which of the following oxoacids of phosphorus are mono-, di- and triprotic acids respectively?
 - (a) HPO_3 , $H_4P_2O_7$, H_3PO_3
- (b) HPO₃, H₆P₄O₁₃, H₃PO₄
- (c) H₃PO₂, H₃PO₃, H₃PO₄
- (d) HPO₃, H₅P₃O₁₁, H₃PO₄

103.		ig the follow: ·H bonds?	ing r	nolecules	s, which	contains the	e max	imum number
	(a)	$H_4P_2O_7$	(b)	H_3PO_2	(c)	H_3PO_3	(d)	H_3PO_4
104.	The n	umber of P—	O—	P bonds i	n cyclic	metaphosph	oric a	cid is
	(a)	zero	(b)	two	(c)	three	(d)	four
105.		n of the follow basic acid as			of phosp	phorus is a r	educi	ng agent and a
	(a)	H_3PO_2	(b)	HPO_3	(c)	H_3PO_3	(d)	$H_4P_2O_5$
106.	Which	n of the follow	ving	is a cyclic	c phosph	ate?		
		$Na_5P_3O_{10}$				$Na_6P_4O_{13}$		
	(c)	$Na_7P_5O_{16}$			(d)	$Na_5P_5O_{15}$		
107.	The pl	hosphate gro	up c	an be esti	mated q	uantitatively	y by p	recipitating
	(a)	$Mg_2(NH_4)_2F$	O_4		(b)	$Mg(NH_4)P$	O_4	
	(c)	$Mg_3(PO_4)_2$			(d)	Mg(NH ₄)H	IPO_4	
108.		n of the follo s a dibasic ac		g oxoacid	ls of pho	sphorus is	a red	ucing agent as
	(a)	H_3PO_3			(b)	HPO_3		
	(c)	H_3PO_2			(d)	H_3PO_4		
109.	Which	of the follow	ving	oxacids o	of phosp	horus contai	ins a I	P—P bond?
		Hypophosp				Tripolypho	•	
	(c)	Pyrophosph	oric	acid	(d)	Hypophos	phori	c acid
110.		reactive, has						he most stable, d conductor of
	(a)	White phosp	phor	us	(b)	Red phosp	horus	
	(c)	Black phosp	horu	ıs	(d)	Scarlet pho	spho	rus
				• <u>Ty</u>	<i>pe</i> 2 •			
Choo	Choose the correct options. More than one option is correct.							

choose the correct options. More than

111. Nitrogen is prepared by heating

- (a) a mixture of CuO and NH_3
- (b) microcosmic salt, $NaNH_4HPO_4\cdot 4H_2O$
- (c) barium azide
- (d) a mixture of NH₄Cl and NaNO₃

112.		h of the following statements are correct for the nitrogen molecule?					
	(a)	The bond order is 2.0.					
	(b)	It is a good ligand.					
	(c)	It is used in the Serpeck process.					
	(d)	It easily reacts with magnesium ev	en at room temperature.				
113.	Nitro	us oxide is prepared by					
	(a)	heating a mixture of NH ₄ Cl and N	aNO ₃				
	(b)	heating a mixture of NH ₄ Cl and N	aNO ₂				
	(c)	the hydrolysis of Mg ₃ N ₂					
	(d)	heating a mixture of nitric oxide as	nd sulphur dioxide				
114.	Which	h of the following statements are co	rrect?				
	(a)	The addition of an electron to NO	forms stable NO				
	(b) The removal of an electron from NO forms stable NO $^{\scriptscriptstyle +}$.						
	(c)	The bond length in NO is greater t	han that in the NO ⁺ ion.				
	(d)	The bond order of NO + is 3.0.					
115.	Which	h of the following statements are co	rrect?				
	(a)	The ionization energy of NO is gre	eater than that of N_2 .				
	(b)	The second ionization energy of nitrogen is less than that of oxygen.					
	(c)	The odd electron in NO is mantibonding π^* molecular orbital t	ore easily removed from the o form the NO^+ ion.				
	(d)	The electronegativity of nitrogen is	s greater than that of oxygen.				
116.	The m	nolecule NO ₂					
	(a)	is diamagnetic					
	(b)	dimerizes to N ₂ O ₄ under suitable of	conditions				
	(c)	is paramagnetic					
	(d)	can be converted into liquid NO_2					
117.	Which	h of the following are not explosive	s?				
	(a)	Calcium cyanamide (b)	Cellulose nitrate				
	(c)	Cellulose (d)	Nitrogen trifluoride				
118.	Amm	nonia acts as a reducing agent when	treated with				

(b) H_2S

(b) NO₂

(d) BrF₃

119. Which of the following is used as a nonaqueous solvent?

(d) NaOCl

(a) CuO (c) HI

(a) NH₃

(c) SO₂

120.		n of the foli			as	ligands in	the preparation o	f
	(a)	NH_3	(b)	PH_3	(c)	$(C_6H_5)_3P$	(d) O ₂	
121.	Hydra	azine is a red	ucin	g agent when	trea	ed with		
	(a)	I_2	(b)	$FeSO_4$	(c)	$CuSO_4$	(d) SnCl ₂	
122.	Which of the following statements are correct for the P ₄ molecule?							
	(a)	The P—P ho	nd a	distances are	eana	1		

- (b) The P—P—P bond angles are 60°.
- (c) It reacts with nitrogen to form a phosphorus-nitrogen polymer.
- (d) The electron affinity of phosphorus is negative.
- **123.** Which of the following statements are correct for the P_4O_6 molecule?
 - (a) The four phosphorus atoms are arranged in a tetrahedral form.
 - (b) The six oxygen atoms are situated along the edges of a tetrahedron.
 - (c) Each oxygen atom is bonded to two adjacent phosphorus atoms.
 - (d) The structure of P₄O₆ is derived from that of PCl₃.
- **124.** Which of the following statements are correct for the P_4O_{10} molecule?
 - (a) The lone pairs on each of the four phosphorus atoms form a coordinate bond with an oxygen atom.
 - (b) Their orbitals exhibit back bonding.
 - (c) It is hydrolysed by water, forming H₄P₂O₇.
 - (d) It has no structural similarity with As₄O₁₀.
- 125. Which of the following oxoacids of phosphorus do not contain a P—P bone?
 - (b) $H_4P_2O_7$ (c) $H_4P_2O_6$ (d) $H_3P_3O_9$ (a) HPO₃

Answers

		11115000		
1. b	2. a	3. d	4. a	5. d
6. c	7. d	8. a	9. a	10. b
11. c	12. b	13. c	14. a	15. b
16. d	17. c	18. d	19. a	20. b
21. b	22. c	23. a	24. d	25. c
26. c	27. b	28. a	29. c	30. b
31. d	32. a	33. b	34. a	35. c
36. d	37. b	38. c	39. a	40. d
41. c	42. b	43. a	44. c	45. b

46. a	47. d	48. c	49. b	50. c
51. b	52. d	53. c	54. a	55. d
56. a	57. c	58. b	59. a	60. c
61. d	62. a	63. d	64. a	65. b
66. c	67. b	68. a	69. d	70. b
71. d	72. c	73. b	74. c	75 . b
76. c	77. c	78. a	79. a	80. a
81. c	82. c	83. b	84. a	85. d
86. b	87. c	88. c	89. d	90. d
91. a	92 . b	93. c	94 . c	95. b
96. a	97. a	98. b	99. c	100. d
101. b	102. c	103. b	104. c	105. a
106. d	107. b	108. a	109. d	110. c
111. a, c	112. b, c	113. a, d	114. b, c, d	115. b, c
116. b, c, d	117. a, c	118. a, d	119. a, b, c, d	120. a, b, c, d
121. a, c	122. a, b	123. a, b, c	124. a, b	125. a, b, d

Hints to More Difficult Problems

- 3. Larger electronegativity difference between Bi and F
- 7. The N—N bond has a double-bond character, which shortens the bond length.
- $8.\ N_2$ as well as CO have a bond order equal to three. See the molecular-orbital energy diagram.
- 11. See the molecular-orbital energy diagram.
- **26.** N_2O_5 is called nitronium nitrate ($NO_2^+NO_3^-$).
- 29. The oxidation number of N is reduced from +3 to +2.
- **30.** The oxidation number of N increases from +3 to +5.
- 31. These are Liebermann nitroso reactions.
- **39.** In option (a) both HNO_3 and H_2SO_4 are oxidizing agents. In option (b), (c) and (d), $FeSO_4$, KI and Au are all reducing agents. So all of them react with HNO_3 .
- **40.** All the metals form a self-protecting oxide layer over themselves.
- 41. $4Zn + NO_3^- + 7(OH^-) \longrightarrow NH_3 + 4ZnO_2^{2-} + 2H_2O$
- **59.** $3Ca + N_2 \longrightarrow Ca_3N_2$ $Ca_3N_2 + 6H_2O \longrightarrow 3Ca(OH)_2 + 2NH_3$

- **63.** The high value of the van der Waals constant a facilitates liquefaction of $H_2O(g)$.
- 65. NH₃ and CaO are both basic. They do not react with each other.

67.
$$Hg_2Cl_2 + 2NH_3 \longrightarrow Hg(NH_2)Cl + Hg + NH_4Cl$$

black

68.
$$2NH_3 + NaOC1 \xrightarrow{\text{Raschig}} N_2H_4 + NaCl + H_2O$$

80.

$$\begin{array}{c}
O \\
-CH_2
\end{array}$$
NH \longrightarrow ····· CO $-\left[NH - (CH_2)_6 - CO\right]_{\overline{n}} NH - CH_2$
- caprolactum

nylon-6

∈- caprolactum

91.
$$P_4 + 6Ca \xrightarrow{heat} 2Ca_3P_2$$

 $Ca_3P_2 + 6H_2O \longrightarrow 3Ca(OH)_2 + 2PH_3$

97.
$$2Ca_3(PO_4)_2 + 6SiO_2 + 10C \longrightarrow P_4 + 6CaSiO_3 + 10CO$$

- **98.** The lower the value of pk_b , the greater will be the basic strength.
- 99. $\overset{0}{P_4} \longrightarrow NaH_2\overset{+1}{PO_2}$ (oxidised) $P_4 \longrightarrow PH_3$ (reduced)
- **101.** Polarization effect

107.
$$MgCl_2 + Na_2HPO_4 + NH_3 \longrightarrow Mg(NH_4)PO_4 \downarrow + 2NaCl$$

112. : N
$$\equiv$$
N : (lone pair of electrons \longrightarrow good ligand)
Al₂O₃ + 3C + N₂ $\xrightarrow{\text{Serpeck process}}$ 2AlN + 3CO \uparrow

116. (b) $2NO_2(g) \rightleftharpoons N_2S_4(g)$ (c) NO_2 contains an unpaired electron (d) $NO_2(g) \longrightarrow NO_2(l)$ using the process of liquefaction (Joule-Thomson effect)

120. All contain a lone pair of electrons

Sulphur

• Type 1 •

1. Which of the following oxoacids of sulphur contains sulphur-sulphur

(b) H₂S₂O₇(d) H₂S₂O₆

(b) S_4N_4

(d) S_3N_8

Choose the correct option. Only one option is correct.

2. The best known sulphur-nitrogen compound is

3. When H₂S gas is passed into aqueous sulphur dioxide,

double bonds? (a) $H_2S_2O_8$

(c) $H_2S_2O_3$

(a) S_2N_4

(c) S_8N_3

	(a)	H ₂ S is converted into a yellow precipitate of sulphur							
	(b)	SO ₂ is converted into a yellow	SO ₂ is converted into a yellow precipitate of sulphur						
	(c)	a clear solution of H ₂ SO ₄ is fo	rme	d					
	(d)	SO_2 as well as H_2S are converted into a yellow precipitate of sulphur							
4.	In thic	osulphuric acid $(H_2S_2O_3)$, the to	WO SI	ulphur atoms have the oxidation					
	(a)	+2 and -2	(b)	+4 and -2					
	(c)	+6 and +2	(d)	+4 and +2					
5.	5. In the cyclo-S ₈ molecule of rhombic sulphur, all the S—S bond length and all the S—S—S bond angles are respectively (give approximativalues)								
	(a)	204 pm and 105°	(b)	102 pm and 120°					
	(c)	204 pm and 180°	(d)	102 pm and 60°					

6.	Which of the following allotropic forms of sulphur is the most stathermodynamically?						ıble		
	(a)	Orthorhomb	oic		(b)	β-monoclin	nic		
	(c)	γ-monoclini	С		(d)	Plastic sulp	hur		
7.	Pure I	H ₂ S is prepar	ed by	y the actior	n of con	centrated H	ICl on		
		FeS ₂				Cu ₂ S			
	(c)	FeS			(d)	Sb_2S_3			
8.	Fe ₂ (SC	$O_4)_3$ can be co	nver	ted into Fe	SO ₄ by	the action o	of		
	(a)	H_2SO_4			(b)	HNO_3			
	(c)	H_2SO_3			(d)	O_3			
9.		dration of cor	ncent	trated H ₂ SO			of P ₄ O	₁₀ produces	3
		SO_2				$H_2S_2O_8$			
	(c)	SO ₂ and SO ₃	3		(d)	SO_3			
10.		rong heating,		SO ₄) ₃ gives					
		SO ₂ and SO ₃				Fe_2O_3 and Se_2O_3			
	()	FeO and SO	0			Fe ₂ O ₃ and S	_		
11.		umber of S—	S bo	nds in the	-	-	ohur t	rioxide is	
	(a) (c)	three one				two zero			
12		that cannot b	0.00	llected ove					
14.	_	N ₂	e co.	nected ove		PH ₃			
	` '	O_2			` '	SO ₂			
13.	Which	n of the follov	vino	acids is no	nt a perc	oxo acid?			
10.			_	$H_2S_2O_8$	•		(d)	H ₂ SO ₅	
14.	Which	n of the follo	wing	oxoacids	of sulp				huı
		bond?							
	(a)	$H_2S_2O_6$	(b)	$H_2S_2O_7$	(c)	$H_2S_2O_8$	(d)	$H_2S_2O_3$	
15.	Which	n of the follow	ving	reactions of	does no	t produce S	O_2 ?		
	(a)	$Ag + H_2SO_4$	(con	ac.) $\xrightarrow{\Delta}$					
	(b)	CaSO ₄ + C -	Δ	\rightarrow					
	(c)	$FeS_2 + O_2$ —	Δ	>					
	(d)	$Zn + H_2SO_4$	(dil`	$\stackrel{\Delta}{\longrightarrow}$					

(d) I_2O_5

(c) KIO₄

17.		eating with sulphur, Na ₂ SO ₃ pr		
		$Na_2S_2O_3$. ,	Na_2S_5
	(c)	$Na_2S_2O_7$	(d)	$Na_2S_4O_6$
18.	Which state?	n of the following molecules of	exist	s as a cyclic trimer in the solid
	(a)	SO_3		O_3
	(c)	SO ₂	(d)	S_4N_4
19.	In wh	ich of the following pairs do be	oth t	he species have a similar shape?
	(a)	SO_4^{2-} and SO_3^{2-}	(b)	SO ₂ and NH ₃
	(c)	SO_4^{2-} and $S_2O_3^{2-}$	(d)	SO_3 and $S_2O_8^{2-}$
20.	Urea	reacts with SO_3 in the presence	of H	I ₂ SO ₄ to produce
	(a)	NH ₂ SO ₃ H (sulphamic acid)	(b)	NH ₂ OH
	(c)	$H_2N \cdot CS \cdot NH_2$ (thiourea)	(d)	$(NH_4)_2SO_4$
21.	The st	ructure of the SO ₃ molecule in	the g	gaseous phase contains
	(a)	only σ -bonds between sulphu	ır an	d oxygen
	(b)	σ -bonds and a (p-p π) bond be	etwe	en sulphur and oxygen
	(c)	σ -bonds and a (p-p π) bond be	etwe	en sulphur and oxygen
	(d)	σ bonds, and a (p-p π) and a oxygen	(p-	$\mathrm{d}\pi$) bond between sulphur and
22.	In pre	paring a standard solution of P FeSO ₄ · (NH ₄) ₂ SO ₄ · 6H		r's salt,
	a few	millilitres of H ₂ SO ₄ is added to		solution. The added H ₂ SO ₄
	(a)	prevents the reduction of the	salt	
	(b)	prevents the oxidation of the	salt	
	(c)	makes the solution homogener formed by hydrolysis, into so		by converting insoluble Fe(OH) ₂ e FeSO ₄
	(d)	neutralizes the ammonia form	ned b	y hydrolysis
23.	K ₄ [Fe	(CN) ₆] reacts with concentrated	d H ₂ S	SO ₄ to produce
	(a)	SO ₂	(b)	CO
	(c)	$(CN)_2$	(d)	$K_3[Fe(CN)_6]$
24.	Conce	entrated H ₂ SO ₄ reacts with PCl		
	(a)	SO_3	(b)	H_3PO_4
	(c)	SO ₂ Cl ₂	(d)	SOCl ₂

16. KIO₃ reacts with SO₂ in an acidic medium to produce

(b) I₂

(a) I_4O_9

- 25. The structure of thioxyl chloride is
 - (a) tetrahedral with one corner of the tetrahedron occupied by a lone pair of electrons, the sulphur atom being at the centre.
 - (b) square planar with the sulphur atom at the centre
 - (c) square pyramidal
 - (d) trigonal bipyramidal
- 26. The structure of the dithionate ion is

(a)
$$\begin{bmatrix} O & O & O \\ O & S & S & O \\ O & O & O \end{bmatrix}^{2^{2}}$$
(b)
$$\begin{bmatrix} O & O & O \\ O & S & S & O \\ O & O & O \end{bmatrix}^{2^{2}}$$
(c)
$$\begin{bmatrix} S \\ | S \\ O & O & O \end{bmatrix}^{2^{2}}$$
(d)
$$\begin{bmatrix} O & S & S & O \\ O & S & S & O \\ O & O & O & O \end{bmatrix}^{2}$$

27. The structure of thiosulphuric acid is

- 28. The hydrolysis of 1 mol of peroxodisulphuric acid produces
 - (a) 2 mol of sulphuric acid
 - (b) 2 mol of peroxomonosulphuric acid
 - (c) 1 mol each of sulphuric acid and peroxomonosulphuric acid
 - (d) 1 mol each of sulphuric acid, peroxomonosulphuric acid and hydogen peroxide
- **29.** The oxidation number of sulphur in the cyclic trimer of sulphur trioxide is
 - (a) +3
- (b) +2
- (c) +4
- (d) +6
- **30.** When SO₂ gas is passed into an acidified K₂Cr₂O₇ solution, the oxidation number of chromium changes from
 - (a) +3 to +6

(b) +6 to +3

(c) +12 to +3

(d) +6 to -3

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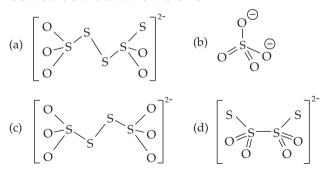
31. In the standardization of Na₂S₂O₃ using K₂Cr₂O₇ by iodometry, the equivalent weight of K2Cr2O7 is

- (a) (molecular weight)/2
- (b) (molecular weight)/6
- (c) molecular weight)/3
- (d) the same as the molecular weight
- **32.** The structure of oleum is

(a)
$$HO = S = OH$$
 (b) $HO = S = OH$ (c) $HO = S = S = OOH$ (d) $HO = S = OOH$ (d) $HO = S = OOH$

- 33. Which of the following reagents is commonly used for the detection of the sulphide ion?
 - (a) 1-Nitroso-2-naphthol
 - (b) Potassium ferricyanide
 - (c) Disodium hydrogen phosphate
 - (d) Sodium nitroprusside
- 34. Which of the following compounds is required in the "brown ring test" for the nitrate ion in an aqueous medium of concentrated H₂SO₄?
 - (a) FeCl₃
- (b) $FeSO_4$ (c) $Fe_2(SO_4)_3$ (d) $Na_2S_2O_3$

35. The structure of the tetrathionate ion is



- **36.** In which of the following oxoacids of sulphur are the two sulphur atoms not in the same oxidation state?
 - (a) $H_2S_2O_8$ (b) $H_2S_2O_3$
- (c) $H_2S_2O_6$ (d) $H_2S_2O_7$

- 37. Which of the following acids is known as dithionic acid?
 - (a) $H_2S_2O_6$

(b) $H_2S_2O_7$

(c) $H_2S_3O_7$

- (d) $H_2S_4O_6$
- **38.** The structure of peroxodisulphuric acid is

(b)
$$HO - S - S - S - S - OH$$

- 39. Which of the following reactions does not occur?
 - (a) $H_2SO_4 + HNO_3 \longrightarrow HSO_4^- + NO_2^+ + H_3O^+$
 - (b) $KIO_3 + SO_2 + H_2O \longrightarrow I_2 + KHSO_4 + H_2SO_4$
 - (c) $NaHSO_3 + Na_2CO_3 \longrightarrow Na_2SO_4 + H_2O + CO_2$
 - (d) $2SO_2 \longrightarrow SO^{2+} + SO_3^{2-}$
- **40.** Which of the following statements is correct?
 - (a) SF₆ does not react with water.
 - (b) OF₆ is d²sp³-hybridized.
 - (c) $S_2O_3^{2-}$ is a linear ion.
 - (d) There is no π -bonding in SO_4^{2-} .
- 41. At high temperature, CaSO₄ reacts with carbon to produce a mixture of
 - (a) CO₂ and SO₂

- (b) CO₂, SO₃ and CaO
- (c) CO₂, SO₂ and CaS
- (d) CaO, SO₂ and CO₂
- 42. Ca(OH)₂ reacts with SO₂ to produce
 - (a) $Ca(HSO_3)_2$

(b) CaSO₃

(c) CaS

(d) CaO

Sulphur 2-85

(b) CS₂

(d) SO₂

44. Elements of Group 16 (except polonium) are called chalcogens because

many metallic ores, mainly as oxides and sulphides

(d) these elements exist in different allotropic forms

(a) these elements, particularly sulphur and oxygen, are present in

(b) a large number of acids contain these elements, particularly

45.	The stability of the hydrides of oxygen, sulphur, selenium and tellurium decreases in the order						
	(a)	$H_2Te > H_2Se > H_2S > H_2O$					
	(b)	$H_2O > H_2S > H_2Se > H_2Te$					
	(c)	$H_2S > H_2O > H_2Se > H_2Te$					
	(d)	$H_2O \approx H_2S < H_2Te < H_2Se$					
46.	Which	n of the following gases dissolv	es ir	n liquid SO ₂ ?			
	(a)	Не	(b)	H_2S			
	(c)	SO ₃	(d)	CO			
47.	On str	rong heating, aqueous sodium	bisu	alphite produces			
	(a)	Na ₂ SO ₄	(b)	$Na_2S_2O_8$			
	(c)	NaHSO ₄	(d)	$Na_2S_2O_5$			
48.	Sodiu	m pyrosulphate, Na ₂ S ₂ O ₇ , can	be n	nade by heating			
	(a)	NaHSO ₄ strongly					
	(b)	NaHSO ₃ strongly					
	(c)	a mixture of Na ₂ S ₂ O ₃ and SO ₂	2				
	(d)	a mixture of Na ₂ SO ₃ and exce	ess of	of sulphur			
49.		O, the bond angle H—O—H iond angles are pretty close to 9		4°28′ but in H ₂ S, H ₂ Se and H ₂ Te This suggests that			
	(a)	oxygen uses sp ² -hybrid orbit orbitals for bonding with the		while S, Se and Te use sp ³ -hybrid lrogen atoms			
	(b)	oxygen uses sp ³ -hybrid orbi atoms while S, Se and Te use	tals almo	to bond with the two hydrogen ost pure p orbitals			
	(c)	2					
	(d)	all the atoms use pure p orbit	als to	to bond with the hydrogen atoms			

43. Which of the following oxides have acidic properties?

sulphur and oxygen

(c) these elements mainly form anions

(a) N_2O

(c) NO

- **50.** H₂S is far more volatile than water because
 - (a) the hydrogen atoms are strongly bonded to sulphur
 - (b) H₂S forms intramolecular hydrogen bonds whereas H₂O forms intermolecular hydrogen bonds
 - the oxygen atom is more electronegative than the sulphur atom, and hence H₂O forms intermolecular hydrogen bonds
 - (d) H₂S forms intermolecular hydrogen bonds whereas H₂O forms intramolecular hydrogen bonds
- 51. H₂S is less stable than water because
 - (a) H₂O molecules form hydrogen bonds but H₂S molecules do not
 - (b) the bonding orbitals of sulphur are larger and more diffuse than those of oxygen, and hence their overlap with the 1s orbitals of the hydrogen atoms is less effective
 - the bonding orbitals of sulphur are smaller and less diffuse than those of oxygen, and hence their overlap with the 1s orbitals of the hydrogen atom is less effective
 - (d) the bonding orbitals of sulphur are smaller and more diffuse than those of oxygen, resulting in a less effective overlap with the 1s orbitals of the hydrogen atoms

• Type 2 •

Choose the correct options. More than one option is correct.

52.	Which	of the	following	are correctly	matched?

- (a) Orpiment \longrightarrow As₂S₂
- (b) Realgar \longrightarrow As₄S₄
- (c) Extraction of sulphur → Frasch process
- (d) Extraction of $SO_2(g) \longrightarrow Raschig process$

53.	SO_2	gas	may	be	prod	luced	ŀ	Эy
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- (a) burning S in air
- (b) roasting ZnS
- (c) heating $Fe_2(SO_4)_3$
- (d) hydrolyzing Na₂SO₄
- 54. Which of the following may act as an oxidizing as well as a reducing agent?
 - (a) H_2S
- (b) H_2SO_4 (c) SO_2
- (d) Na_2SO_3
- 55. Which of the following may form acid salts as well as normal salts?
 - (a) NaHSO₄
- (b) H₂S
- (c) H_2SO_4
- (d) NaHSO₃

- **56.** In which of the following reactions does concentrated H_2SO_4 act as an oxidizing agent?
 - (a) $FeSO_4 + H_2SO_4 \longrightarrow$
- (b) $ZnO + H_2SO_4 \longrightarrow$
- (c) $KBr + H_2SO_4 \longrightarrow$
- (d) NaCl + $H_2SO_4 \longrightarrow$
- 57. Which of the following oxoacids of sulphur contain a peroxo-link?
 - (a) $H_2S_2O_3$

(b) H_2SO_5

(c) $H_2S_2O_6$

- (d) $H_2S_2O_8$
- 58. Which of the following species have (zero) dipole moment?
 - (a) SO_2

(b) CS₂

(c) SO_4^{2-}

- (d) H_2SO_3
- **59.** Which of the following allotropes of sulphur exist as S_8 molecules with a puckered-ring structure assuming a crown conformation?
 - (a) γ-Monoclinic

(b) β-Monoclinic

(c) α-Rhombic

- (d) γ-Rhombic
- **60.** Which of the following statements are correct for the SO_4^{2-} ion?
 - (a) It is tetrahedral.
 - (b) All the S—O bond lengths are equal, and shorter than expected.
 - (c) It contains four σ-bonds between the S and the O atoms, two π-bonds delocalized over the S and the four O atoms, and all the S—O bonds have a bond order of 1.5.
 - (d) It is square planar.

Answers

1. c	2. b	3. d	4. 6	5. a
6. a	7. d	8. c	9. d	10. b
11. d	12. d	13. c	14. a	15. d
16. b	17. a	18. a	19. c	20. a
21. d	22. c	23. b	24. c	25. a
26. b	27. c	28. c	29. d	30. b
31. b	32. b	33. d	34. b	35. c
36. b	37. a	38. d	39. c	40. a
41. c	42. b	43. d	44. a	45. b
46. c	47. d	48. a	49. b	50. c
51. b	52. a, b, c	53. a, b	54. c, d	55. b, c
56. a, c	57. b, d	58. b, c	59. a, b, c	60. a, b, c

Hints to More Difficult Problems

4. $H = O_{+4} / S^2$

The cyclic trimer of SO₃ does not contain an S—S bond.

12. SO₂ reacts with water in the following manner.

$$SO_2 + 2H_2O \longrightarrow SO_4^{2-} + 4H^+ + 2e$$

16.
$$2IO_{3}^{-} + 12H^{+} + 10e \longrightarrow I_{2} + 6H_{2}O$$

 $SO_{2} + 2H_{2}O \longrightarrow SO_{4}^{2-} + 4H^{+} + 2e \times 5$
 $2IO_{3}^{-} + 5SO_{2} + 4H_{2}O \longrightarrow I_{2} + 5SO_{4}^{2-} + 8H^{+}$

- **28.** $H_2S_2O_8 + H_2O \longrightarrow H_2SO_4 + H_2SO_5$
- **29.** The oxidation number of S in SO_3 is +6. This is true for the trimer also.
- **30.** $Cr_2O_7^{2-} + H^+ + 3SO_2 \longrightarrow 2Cr^{3+} + 7H_2O + 3SO_4^{2-}$. The oxidation number of Cr in $Cr_2O_7^{2-}$ is +6 and that in Cr^{3+} is +3.
- 31. $Cr_2O_7^{2-} + 14H^+ + 6e \longrightarrow 2Cr^{3+} + 7H_2O$ Eq wt $(K_2Cr_2O_7) = \frac{\text{mol. wt}}{6}$ (six electrons are gained in the process)
- 33. The sulphide ion, S^{2-} , reacts with sodium nitroprusside to give a dark red complex.

$$S^{2-} + Na_2[Fe(CN)_5NO] \longrightarrow Na_2[Fe(CN)_5NOS] + 2e$$

dark red complex

- 36. See the answer to Q. 4.
- **40.** Because of the high S—F bond strength, coordination saturation is stable. The lack of polarity of the molecule is also a factor.
- **43.** $SO_2 + 2H_2O \longrightarrow H_2SO_4 + 2H^+ + 2e$
- 55. H_2S as well as H_2SO_4 form two series of salts. For example, Na_2S and $NaHSO_4$, Na_2SO_4 .

Sulphur 2-89

Feroxide
$$O_3$$
H O_3 S O_3 H O_3 S O_5 O O_5 H O_3 S O_5 O O_5 H O_3 S O_5 O O_5 O

The Halogens

• Type 1 •

Choose the correct option. Only one option is correct.

1.	Among the following, the properties of which pair of halogens are more
	similar than those of the other pairs?

(a) Fluorine and bromine

(b) Fluorine and chlorine

(c) Chlorine and bromine

(d) Fluorine and astatine

- 2. The colour of halogens progressively deepens from fluorine to iodine because
 - (a) halogens of higher atomic number absorb light of longer wavelength since the difference in energy between the ground state and excited state decreases as the atomic number increases.
 - (b) fluorescence and phosphorescence become more intense as the atomic numbers of halogen increases
 - (c) the standard electrode potential increases from I₂ to F₂.
 - (d) halogens of higher atomic number absorb light of shorter wavelength since the difference in energy between the ground state and excited state increases as the atomic number increases.
- **3.** Which of the following has been arranged in order of increasing bond energy?

(a)
$$Cl_2 < I_2 < Br_2 < F_2$$

(b)
$$F_2 < I_2 < Cl_2 < Br_2$$

(c)
$$F_2 < Br_2 < Cl_2 < I_2$$

(d)
$$I_2 < F_2 < Br_2 < Cl_2$$

4. Which of the following pairs of halogens have approximately identical bond energy?

(a) F_2 and Br_2

(b) F_2 and I_2

(c) F₂ and Cl₂

(d) Cl_2 and I_2

5.	The	reaction

$3ClO^{-}(aq) \longrightarrow ClO_{3}^{-}(aq) + 2Cl^{-}(aq)$	3ClO - (a	$aq) \longrightarrow$	ClO_3^-	(aq) -	+ 2C1 -	(aq)
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is an example of

(a) oxidation

- (b) reduction
- (c) disproportionation
- (d) decomposition
- 6. The F—F bond is weak because
 - (a) the repulsion between the nonbonding pairs of electrons of two fluorine atoms is large
 - (b) the ionization energy of the fluorine atom is very low
 - (c) the length of the F—F bond much larger than the bond lengths in other halogen molecules
 - (d) the F—F bond distance is small and hence the internuclear repulsion between the two F atoms is very low
- 7. Fluorine is prepared by the electrolysis of
 - (a) molten NaF
 - (b) a molten mixture of Ca₃(PO₄)₂ · CaF₂ and cryolite
 - (c) a solution of KHF₂ in HF
 - (d) a solution of KHF₂ in KF
- 8. Fluorine may be prepared by the electrolysis of a molten mixture of KHF₂ and KF. The anode is made of
 - (a) copper

(b) iron

(c) graphite

- (d) ungraphitized carbon
- 9. Fluorine shows only one oxidation state (-1) because it has
 - (a) a high electronegativity
 - (b) no d orbital available for bonding
 - (c) a small covalent radius
 - (d) a low bond energy
- 10. Fluorine reacts with water to produce
 - (a) HF and H_2O_2

(b) HF, O_2 and F_2O_2

(c) F^- , O_2 and H^+

- (d) F_2O and F_2O_2
- 11. Fluorine reacts with H₂S to produce
 - (a) SF_2

(b) SF₄

(c) S and HF

- (d) SF_6
- **12.** Fluorine reacts with dilute NaOH and concentrated NaOH to respectively produce
 - (a) OF_2 and O_3

(b) F_2O_2 and NaF

(c) HF and O_2

(d) OF₂ and NaF

13.	Fluor	ine reacts with aqueous KClO ₃	to p	roduce
	(a)	KClO ₄	(b)	KClO ₂
	(c)	KCl	(d)	KClO
14.	In wh	ich of the following reactions i	s Cl ₂	(gas) produced?
	(a)	$NaCl + K_2Cr_2O_7 + H_2SO_4$ (con	c.) —	\longrightarrow
	(b)	$KCl + Br_2 \longrightarrow$		
	(c)	$Ca(OCl)Cl + H_2O \longrightarrow$		
	(d)	$NaOCl + NH_3 \longrightarrow$		
15.	NaHS	SO_4 reacts with F_2 to produce m	nainl	y
	(a)	$Na_2S_2O_8$	(b)	$Na_2S_2O_5$
	(c)	$Na_2S_2O_7$	(d)	$Na_2S_2O_3$
16.	Cl ₂ re		conc	entrated NaOH to respectively
	(a)	NaClO and NaClO ₂	(b)	NaClO ₃ and NaClO
	(c)	NaClO and NaClO ₃	(d)	NaCl and NaClO ₄
17.	Chlor	ine can be manufactured by nce of a CuCl ₂ catalyst at 450°C	the C. Th	oxidation in air of HCl in the process is known as the
	(a)	Deacon process	(b)	Nelson process
	(c)	Chloride process	(d)	Solvay process
18.		ine acts as an oxidizing agent v		
	` '	O_3	` /	$Fe_2(SO_4)_3$
	(c)	FeSO ₄	(d)	$KMnO_4$
19.		n of the following will act as th		0
		Cr(OH) ₂		ClO ₃ (OH)
		PO(OH) ₃		SO(OH) ₂
20.		tandard reduction potentials of		_
	` ,	$F_2 > Cl_2 > I_2 > Br_2$		$Cl_2 > F_2 > I_2 > Br_2$
	(c)	$I_2 > Br_2 > Cl_2 > F_2$	(d)	$F_2 > Cl_2 > Br_2 > I_2$
21.		n of the following has the high HCl (b) HF		nolar enthalpy of vapourization? HI (d) HBr
22.	The el	lectrolysis of brine produces		
		only Cl ₂	(b)	NaOH and NaClO ₃
	(c)	Cl ₂ and NaOH	(d)	NaCl and NaClO

23. Which of the following reactions does not produce bromine?

	(a)	$NaBr + MnO_2 + H_2SO_4 \longrightarrow$		
	(b)	$NaBrO_3 + NaBr + H_2SO_4$	>	
	(c)	$MgBr_2 + Cl_2 \longrightarrow$		
	(d)	$NaBr + I_2 \longrightarrow$		
24.	Bromi	ne reacts with hot and concent	trate	d Na ₂ CO ₃ to produce
	(a)	$NaBr + NaBrO + CO_2$	(b)	$NaBr + NaBrO_4 + CO_2$
	(c)	$NaBr + NaBrO_3 + CO_2$	(d)	$NaBrO + NaBrO_3 + CO_2$
25.	Which	n of the following reactions doe	es no	et liberate iodine?
	(a)	$KI + MnO_2 + H_2SO_4 \longrightarrow$		
	(b)	$NaIO_3 + NaHSO_3 \longrightarrow$		
	(c)	$CuSO_4 + KI \longrightarrow$		
	(d)	$NaIO_3 + NaHSO_4 + Cl_2$		
26.	a com			neated with concentrated H_2SO_4 , xidation number of $Cr = +6$. The
	(a)	$Cr_2(SO_4)_3$	(b)	CrO ₂ Cl ₂
	(c)	K_2CrO_4	(d)	KCrO ₃ Cl
27.		n of the following halogens is e		
	(a)	-	(b) (d)	Cl ₂
20	` '	Br ₂	` ,	_
28.	accept		inas	is the strongest fluoride-ion
	(a)	SbF_5 (b) AlF_3	(c)	IF ₇ (d) CaF ₂
29.		cid strengths of HF, HCl, HBr a	and I	HI increase in the order
	` '	HCl < HBr < HI < HF	` '	HBr < HCl < HI < HF
20	. ,	HF < HCl < HBr < HI		HF < HBr < HCl < HI
30.		leaching action of chlorine occu moisture		sunlight
	` '	pure oxygen		pure sulphur dioxide
31.	Which	n of the following gases shows	blea	ching action due to reduction?
	(a)	H_2O_2	(b)	H_2S
	(c)	SO ₂	(d)	NH_3

- **32.** The boiling points of HF, HCl, HBr and HI follow the order (a) HF > HCl > HBr > HI (b) HF > HI > HBr > H
 - (c) HI > HBr > HCl > HF
- (b) HF > HI > HBr > HCl(d) HCl > HF > HBr > HI
- 33. In which of the following reactions is bromine liberated?
 - (a) $KBr(aq) + I_2 \xrightarrow{\Delta}$
- (b) $HI(aq) + KBr(aq) \xrightarrow{\Delta}$
- (c) $KBr(aq) + F_2 \xrightarrow{\Delta}$
- (d) $KBr(aq) + SO_2 \xrightarrow{\Delta}$
- **34.** Which of the following hydrogen halides has high dielectric constant and low viscosity?
 - (a) HCl
- (b) HBr
- (c) HI
- (d) HF
- 35. In which of the following reactions is HF liberated?
 - (a) KHF₂ $\xrightarrow{\Delta}$

(b) $F_2 + KCl \xrightarrow{\Delta}$

(c) $PH_4F \xrightarrow{\Delta}$

- (b) $F_2 + NaOH \longrightarrow$
- **36.** Among the following, which acts as the strongest acid in reactions where HNO₃ behaves like a base?
 - (a) HI

(b) H₃PO₃

(c) HF

- (d) HIO₄
- 37. By which of the following reactions is HF prepared industrially?

(a)
$$CaF_2 + H_2SO_4 \xrightarrow{\Delta}$$

(b)
$$Na_3AlF_6 + Al_2O_3 + H_2SO_4 \xrightarrow{\Delta}$$

- (c) $F_2 + HNO_3 \xrightarrow{\Delta}$
- (d) All of these
- **38.** Hydrofluoric acid cannot be stored in glass vessels because it reacts with glass to form
 - (a) Na₂SiO₃ and F₂
- (b) Na₂SiF₆

(c) SiF₂

- (d) $Na_4[SiF_6]$
- 39. Which of the following reactions does not produce HBr?
 - (a) $KBr + H_3PO_3 \xrightarrow{\Delta}$
 - (b) $P(red) + Br_2 + H_2 \xrightarrow{\Delta}$

(c) NaBr +
$$H_2SO_4$$
(conc.) $\xrightarrow{\Delta}$

(d)
$$Br_2 + NaOH \xrightarrow{\Delta}$$

- **40.** The boiling point of HF is unexpectedly higher than those of HCl, HBr and HI because of
 - (a) the greater polarity of the H—F bond
 - (b) the intermolecular hydrogen bond formed between H and F
 - (c) the intramolecular hydrogen bond formed between H and F as a result of the HF molecule becoming highly associated
 - (d) all of these
- 41. The bond dissociation energy of HF is approximately twice that of
 - (a) HCl

(b) HBr

(c) F₂

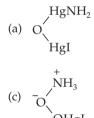
- (d) HI
- **42.** Arrange CCl₄, AlCl₃, PCl₅ and SiCl₄ according to ease of hydrolysis.
 - (a) $CCl_4 < SiCl_4 < PCl_5 < AlCl_3$
 - (b) AlCl₃ < CCl₄ < PCl₅ < SiCl₄
 - (c) CCl₄ < AlCl₃ < PCl₅ < SiCl₄
 - (d) $CCl_4 < AlCl_3 < SiCl_4 < PCl_5$
- **43.** Arrange HF, HCl, HBr and HI in decreasing order of bond dissociation energy.
 - (a) HF > HCl > HBr > HI
- (b) HI > HBr > HCl > HF
- (c) HI > HF > HCl > HBr
- (d) HBr > HCl > HF > HI
- 44. The increasing order of the dipole moments of halogen acids is
 - (a) HI < HBr < HCl < HF
- $(b) \quad HF < HCl < HBr < HI$
- (c) HBr < HI < HCl < HF
- $(d) \quad HCl < HF < HBr < HI$
- **45.** The ionic character of the metal-halogen bond (M—X) in metal halides decreases in the order
 - (a) M Br > M Cl > M F > M I
 - (b) M-I > M-Br > M-Cl > M-F
 - (c) M—Cl > M—Br > M—I > M—F
 - (d) M F > M Cl > M Br > M I
- **46.** Which of the following is arranged in order of increasing ionic character?
 - (a) $PbCl_2 < SnCl_4 < KCl < MgCl_2$ (b) $SnCl_4 < PbCl_2 < KCl < MgCl_2$
 - (c) $SnCl_4 < PbCl_2 < MgCl_2 < KCl$ (d) $PbCl_2 < SnCl_4 < MgCl_2 < KCl$

- 47. Which of the following shows the arrangement of halide ions in increasing order of reducing property?
 - (a) $Cl^- < F^- < I^- < Br^-$
- (b) $F^- < Cl^- < Br^- < I^-$
- (c) $Br^- < I^- < F^- < Cl^-$
- (d) $I^- < Br^- < Cl^- < F^-$
- 48. The correct order of arrangement of the acids ClOH (I), BrOH (II) and IOH (III) in order of decreasing acid strength is
 - (a) I > III > II

(b) II > III > I

(c) I > II > III

- (d) III > II > I
- 49. Nessler's reagent is used to detect NH₃. This reagent is prepared by mixing
 - (a) KI (excess) + HgCl₂ + KOH
 - (b) KCl + HgI₂ + KOH
 - (c) KI (excess) + $HgCl_2 + NH_4^+OH^-$
 - (d) KI (excess) + Hg₂Cl₂ + I₂ solution
- 50. Ammonia reacts with Nessler's reagent to give a brown precipitate known as iodide of Millon's base. This base is believed to have the structure







$$(d) \ \ O \bigg\backslash \\ HgI$$

- **51.** Which of the following statements is correct?
 - (a) The length of the O—O bond in O_2F_2 is greater than that in H_2O_2 .
 - (b) The length of the O—O bond in O_2F_2 is less than that in H_2O_2 .
 - (c) O_2F_2 does not contain the peroxide bond -O-O.
 - (d) The O—O bond distance is the same in H_2O_2 and O_2F_2 .
- 52. Cl₂O (gas) dissolves in a KOH solution to produce
 - (a) Cl_2O_7

(b) KCl

(c) KClO

- (d) KClO₃
- 53. Which of the following molecules has the highest bond angle and is V-shaped?
 - (a) OF_2

(b) Cl₂O

(c) H₂S

(d) Br₂O

- **54.** The structure of OF_2 is
 - (a) tetrahedral with each of two positions occupied by a lone pair of electrons
 - (b) tetrahedral with one position occupied by a lone pair of electrons
 - square planar with two positions each occupied by a lone pair of electrons
 - (d) a pentagonal bipyramid with three positions occupied by a lone pair of electrons each
- 55. ClO₂ reacts with NaOH to give a mixture of
 - (a) NaClO₂ and NaClO₄
- (b) NaClO₂ and NaClO₃
- (c) NaClO and NaClO₂
- (d) NaClO₃ and NaClO₄
- **56.** ClO_2 reacts with O_3 to give
 - (a) Cl_2O_4

(b) Cl₂O

(c) Cl_2O_6

- (d) ClO₄
- 57. Among the following oxoacids of chlorine, acidity increases in the order
 - (a) HClO < HClO₂ < HClO₃ < HClO₄
 - (b) HClO₄ < HClO₃ < HClO₂ < HClO
 - (c) HClO₂ < HClO₄ < HClO₃ < HClO
 - (d) HClO₃ < HClO < HClO₄ < HClO₂
- **58.** Among the following conjugate bases of oxoacids of chlorine, which arrangement shows the correct order of increasing hydration energy and basic character?
 - (a) $ClO^- < ClO_2^- < ClO_3^- < ClO_4^-$
 - (b) $ClO_4^- < ClO_3^- < ClO_2^- < ClO^-$
 - (c) $ClO_3^- < ClO_4^- < ClO_2^- < ClO^-$
 - (d) $ClO_4^- < ClO_3^- < ClO^- < ClO_2^-$
- **59.** The hybridization state of the iodine atom in ICl_2^- is
 - (a) $d^2 sp^3$
- (b) dsp^2
- (c) sp^2
- (d) sp^3d
- **60.** Which of the following statements is correct regarding the ClO_3 and Cl_2O_6 molecules?
 - (a) They are paramagnetic and diamagnetic respectively.
 - (b) They are diamagnetic and paramagnetic respectively.
 - (c) Both are diamagnetic.
 - (d) Both are paramagnetic.
- **61.** The structure of Cl_2O_7 is
 - (a) tetrahedral

- (b) square pyramidal
- (c) pentagonal bipyramidal
- (d) trigonal bipyramidal

62.	. Which of the following is not the salt of periodate?				
	` '	KIO_4		$Na_4I_2O_9$	
	(c)	$Pb_4(IO_5)_2$	(d)	Ag_5IO_6	
63.		ybridization states of the chlor are respectively	ine a	atom in the ions ClO_4^- , ClO_3^- and	
	(a)	sp^3 , dsp^2 and sp^2	(b)	dsp^2 , sp^3 and sp^2	
	(c)	sp ³ in all the species	(d)	sp^3 , sp^2 and sp	
64.	Which	n of the following is not param	agne	etic?	
	(a)	NO ₂	(b)	O_2^-	
	(c)	ClO ₂	(d)	Cl ₂ O ₇	
65.	Which	n of the following interhalogen	s do	es not exist?	
		BrF ₅	(b)		
	(c)	IF ₇	(d)	BrF ₃	
66.	Which	n of the following is not linear?			
	(a)	IBrF ⁻	(b)	ICl_2^-	
	(c)	I_3^+	(d)	I_3^-	
67.	Unde	r anhydrous conditions CCl ₄ re	eacts	with HF to produce	
	(a)	CF ₄	(b)	CHCl ₃	
	(c)	CH_2Cl_2	(d)	CCl ₂ F ₂	
68.	In spi		mol	ecule, ClO ₂ does not dimerize	
	(a)	the odd electron is delocalized	b		
	(b)	the odd electron is localized in	n the	chlorine atom	
		the two Cl—O bonds do not h		-	
	(d)	of $p\pi$ - $p\pi$ bonding in the chlor	ine a	tom	
69.	Which	n of the following statements is	corr	rect?	
		I_2O_4 exists as $IO^+ \cdot IO_3^-$.			
		I_4O_9 exists as $I^{3+} \cdot (IO_3^-)_3$.			
		On decomposition, I ₄ O ₉ gives	I_2O_5	·	
	(d)	All of these			
70.	In the to form			oes a disproportionation reaction	
	(a)	KClO ₄ and KClO ₃	(b)	KCl and O ₂	

(d) KClO₄ and Cl₂

(C) KClO₄ and KCl

71. When an aqueous solution of sodium fluoride is electrolysed the gas

(b) O₂

(d) O_2 and F_2

liberated at the anode is

(a) H₂(c) F₂O

72. The shape of H_5IO_6 is

	(a)	irregular tetrahedral	(b)	square pyramidal
	(c)	pentagonal bipyramidal	(d)	octahedral
73.	The co	ommon form of periodic acid is	HIC	$O_4 \cdot 2H_2O$ or H_5IO_6 . This is called
	(a)	metaperiodic acid	(b)	dimesoperiodic acid
	(c)	mesoperiodic acid	(d)	paraperiodic acid
74.	On str	cong heating, the acid H ₅ IO ₆ gi	ves	
	(a)	I_2O_4	(b)	I_2O_5
	(c)	I_2O_7	(d)	HIO_4
75.	Which	n of the following statements is	corr	rect for the CsBr ₃ molecule?
	(a)	It contains Cs $^{3+}$ and Br $^-$ ions.		
		It is a complex compound.		
		It contains Cs ⁺ and Br ₃ ⁻ ions.		
	(d)	It contains Cs ⁺ , Br ₂ and Br ⁻ ic	ns ir	n its crystal lattice.
76.				halide ions, have two or more
		of which at least one is nitrogo		
		interhalogen compounds	. ,	nitriles
		pseudonitriles		pseudohalides
77.		ybridization states of the centra spectively	al ato	oms of the ions I_3^- , ICl^- and ICl_4^-
	(a)	sp^2 , dsp^2 , pd^3	(b)	sp^3d , sp^3d and sp^3d^2
	(c)	sp^3d , sp^3d , dsp^2	(d)	sp, sp, dsp ²
78.		ybridization state of the centra rule are respectively	al ato	om of BrF ₅ and the shape of the
	(a)	sp ³ d, T-shaped	(b)	sp ³ d ² , octahedral
	(c)	sp ³ d ² , square pyramidal	(d)	sp ³ d ³ , trigonal bipyramidal
79.		ybridization state of the centrule are respectively	al at	om of IF ₇ and the shape of the
	(a)	sp ³ d ³ , pentagonal bipyramid	al	
	(b)	sp ³ d ³ , square antiprismic		
	(c)	sp ³ d ² , octahedral		
		sp ³ d ³ , square based pyramid	al	
	` '	1 1 1 1 1		

80.	0. Which of the following is a pseudohalide ion?							
	(a)	N_3^-	(b)	ONC -				
	(c)	SCN ⁻	(d)	All of these				
81.	Which	n of the following is not a ps	eudoha	lide ion?				
	(a)	SeCN ⁻	(b)	TeCN ⁻				
	(c)	SCN ⁻	(d)	$S_4 N_4^{2-}$				
82.		n of the following compound Cl_2N_3 (b) BrCN		t an "interpseudohalogen"? CICN (d) ICN				
83.	Which	n of the following are not ps	eudoha	logens?				
		Cyanogens	(b)	Thiocyanogens				
	(c)	Azidocyanogens	(d)	Azidocarbon disulphides				
84.	Which	n of the following is the mos	st electro	onegative?				
	(a)	N_3^-	(b)	Br ⁻				
	(c)	Ι-	(d)	NCS-				
85.	Which charac		d in dec	reasing order of pseudohalogen				
	(a)	$(SeCN)_2 > (CN)_2 > (SCN)_2$	(b)	$(SCN)_2 > (CN)_2 > (SeCN)_2$				
	(c)	$(CN)_2 > (SeCN)_2 > (SCN)_2$	(d)	$(SeCN)_2 > (SCN)_2 > (CN)_2$				
		• Typ	1e 2 •					
		<u> </u>	<u>,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,</u>					
Choo	se the	correct options. More than o	one opti	on is correct.				
86.	Which halide	n of the following will display?	ace the l	halogen from the solution of the				
	(a)	Br ₂ added to an NaCl solut	tion					
	(b)	Cl ₂ added to a KBr solution	n					
	(c)	Cl ₂ added to an NaF soluti	on					
	(d)	Br ₂ added to a KI solution						
87.	Chlor	ine is produced by the						
	(a)	electrolysis of an aqueous s						
	(b) action of concentrated HCl on MnO ₂							

(c) action of concentrated $\rm H_2SO_4$ on NaCl in the presence of $\rm MnO_2$

(d) evaporation of sea water

88. Which of the following exist? (b) FCl₃ (a) ICl₃ (c) IF₇ (d) BrF₃ 89. Chlorine behaves as an oxidizing agent upon reaction with (a) $Fe_2(SO_4)_3$ (b) O₃ (d) NaNO₂ (c) $Na_2S_2O_3$ 90. Which of the following compounds react with fluorine? (a) NaCl (b) KF (c) B_2O_3 (d) Al_2O_3 91. Which of the following are used in the preparation of Nessler's reagent? (b) KI (a) Hg₂Cl₂ (c) HgCl₂ (d) NaOH 92. By which of the following structures may Cl₂O₆ be represented? (a) $ClO_2^+ClO_4^-$ (d) $ClO_3 ClO_3^+$ 93. By which of the following reactions is HClO₄ prepared? (a) $HClO_3 \xrightarrow{\Delta}$ (b) KClO₄ + HCl $\xrightarrow{\Delta}$ (c) $NH_4Cl + KClO_3 \xrightarrow{\Delta}$ (d) $POCl_3 + HNO_3 \xrightarrow{\Delta}$ **94.** Which of the following molecules are angular? (a) F_2O_2 (b) Cl₂O (d) ClO₂ (c) KI₃ 95. Which of the following are V-shaped? (a) ClF_2^+ (b) BrF_2^+ (d) Cl₂F⁺ (c) ICl₂⁺ 96. Which of the following halides form infinite chains? (a) BeF₂ (b) BeCl₂

(d) I_2Cl_6

(c) AlCl₃

86. b, d

96. a, b

91. b, c, d

87. a, b, c

97. a, b, c, d

92. a, c

97. Which of the following halides hydrolyses easily? (a) BCl₃ (b) SiCl₄ (c) PCl₃ (d) PCl₅ **98.** Which of the following halides are stable to hydrolysis? (a) NH₄Cl (b) CCl₄ (d) SnCl₄ (c) SF₆ 99. Which of the following are bent? (a) F_3^- (b) Br₂⁺ (c) I_3^- (d) I_3^+ 100. Which of the following molecules undergoes disproportionation under suitable conditions? (c) NaCl (d) NaOCl (a) HClO₂ (b) HClO₄ Answers **1.** c **2.** a **3.** d **4.** b **5.** c **7.** c 8. d 9. b **6.** a **10.** c **11.** d **12.** d **13.** a **15.** a **14.** c **20.** d **16.** c **17.** a **18.** c **19.** b **21.** b **22.** c **23.** d **24.** c **25.** d **26.** b **27.** d 28. a **29.** c **30.** a **31.** c **32.** b **33.** c **34.** d **35.** a **37.** a **38.** b **36.** c **39.** c **40**. b **41.** d **42.** d **44.** a **45.** d **43.** a **47.** b **46.** c **48.** c **49.** a **50.** a **51.** b **52.** c **53.** d **54.** a **55.** b **56.** c **57.** a **58.** b **59.** d **60.** c 65. b **61.** a **62.** c **63.** c **64.** d **66.** c **67.** d **68.** a **69.** d **70.** c **71**. b **72.** d **73.** d 74. b **75.** c 77. b **76.** d **78.** c **79.** a **80.** d **81.** d **82.** a **83.** c **84.** a 85. b

88. a, c, d

93. a, b

98. b, c

89. c, d

94. b, d

99. b, d

90. a, b, c, d

95. a, b, c, d

100. a, d

Hints to More Difficult Problems

- 1. They are of similar size.
- 4. The low dissociation energy of F₂ arises from the repulsion between the unpaired electrons in the two atoms. Because of the large size of I₂, the degree of dissociation is low.
- 5. The oxidation number of Cl in ClO $^-$ is +1, +5 in ClO $^-$ and -1 in Cl $^-$.
- **26.** This is the basis of the chromyl chloride test.
- **29.** Size and bond length increase in this order, and pK_a decreases. Therefore, acid strength increases.
- **32.** The boiling point of HF is highest due to hydrogen bonding, i.e., HF is the associated molecule $(HF)_n$. In the rest of the halides, boiling point increases with molar mass—they do not undergo hydrogen bonding because of their large size and low electronegativity.
- 34. HF is a liquid. See the answer to Q. 32.
- 39. (a) NaBr + $H_2SO_4 \longrightarrow NaHSO_4 + HBr$ (b) $2HBr + H_2SO_4 \longrightarrow 2H_2O + Br_2 + SO_2$

Here HBr is produced in reaction (a), and is a reducing agent. HBr reduces H_2SO_4 to SO_2 and is itself oxidized to Br_2 so instead of getting HBr we get a mixture of Br_2 and SO_2 .

- **42.** PCl₅ readily undergoes coordination number expansion to a maximum covalence of six.
- 45. Apply Fajans' rules.
- **47.** Use the values of the respective standard reduction potentials.
- **49.** Nessler's reagent is a mixture of $K_2[HgI_4]$ and KOH. It is prepared by the following sequence of reactions.

$$2KI + HgCl_2 \longrightarrow 2KCl + HgI_2$$

 $HgI_2 + 2KI \longrightarrow K_2[HgI_4] \xrightarrow{+KOH}$ Nessler's reagent

- **57.** The oxidation number of Cl in $HClO_4$ is +7 (maximum), the ClO_4^- ion is the largest in size and the pK_a value of $HClO_4$ is the least, meaning that it has the greatest K_a (dissociation constant). This accounts for the maximum acid strength of $HClO_4$.
- 58. Use the Brönsted-Lowry concept.
- 70. $4KClO_3 \longrightarrow 3KClO_4 + KCl$

This is a disproportionation reaction.

74.
$$2H_5IO_6 \xrightarrow{100^{\circ}C} \phantom{2HIO_4} \phantom{200^{\circ}C} \phantom{2H_2O} \phantom{2H_2O}\phantom{2H_2O}\phantom{2H_2O}\phantom{2H_2O$$

86. Order of electronegative effect of halogens

89.
$$4Cl_2 + S_2O_3^{2-} + 5H_2O \longrightarrow 8Cl^- + 2SO_4^{2-} + 10H^+$$

 $Cl_2 + NO_2^- + H_2O \longrightarrow 2Cl^- + NO_3^- + 2H^+$

The oxidation number of Cl_2 changes from 0 to -1. Hence, Cl_2 behaves as an oxidizing agent.

100.
$$ClO_2^- \longrightarrow ClO_2 + Cl^-$$
 oxidation number
 $+1$ -1 $+5$ oxidation number
 $+1$ -1 $+5$ oxidation number

10

The Noble Gases

(b) Argon II

(d) Helium I

Choose the correct option. Only one option is correct.

1. Which of the following is a superfluid?

(a) Krypton I

(c) Helium II

2.	Heliu	m II is the most extraordinary liquid with						
	(a)	zero viscosi	zero viscosity and very high heat conductivity					
	(b) zero viscosity and low heat conductivity							
	(c)	(c) very high viscosity and zero heat conductivity						
	(d)	very high v	very high viscosity and very high heat conductivity					
3.								cavities of the sare known as
	(a)	polyhydrate	es		(b)	supercooled	l soli	ds
	(c)	stoichiomet	ric co	mpounds	(d)	clathrates		
4.	XeF ₄ r	eacts violent	ly wi	th water to g	give			
	(a)	$Xe + O_2$			(b)	$XeO_3 + O_2 +$	HF	
	(c)	$Xe + O_2 + H$	$F + \lambda$	GeO_3	(d)	$XeOF_3$		
5.	In clas	thrate atoms	or m	olecules the	bond	formed is		
	(a)	metallic			(b)	covalent		
	(c)	ionic			(d)	They do no	t forr	n bonds.
6.	Solid	XeF ₆ exists as	3					
	(a)	XeF ₄ ⁺ and F ₂	- 2		(b)	XeF ₅ ⁺ and F	_	
	(c)	XeF ₇ and F	+		(d)	Xe ⁴⁺ and F	1–	
7.	The fl	uoride of xer	on v	vith zero dip	ole n	noment is		
		XeF ₆		-			(d)	XeF ₂

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8.		gases are difficult to liquefy b	ecau	use their
		dispersion forces are large		
		dispersion forces are small		
		ionization energies are low		
		affinity energies are high		
9.		lement which has the highest f		•
	(a)	hydrogen	` '	xenon
	(c)	fluorine	(d)	helium
10.	Which	n of the following statements is	s inco	correct for helium?
	(a)	It has a positive Joule–Thoms	on c	coefficient above 40 K.
	(b)	Its spontaneous expansion ca		-
	(c)	It has to be compressed befor	e it c	can liquefy.
	(d)	It has a negative Joule-Thoms	son c	coefficient above 40 K.
11.	Xenor	n trioxide (XeO ₃) has a		
	(a)	T-shaped structure		
	(b)	square-pyramidal structure		
	(c)	trigonal pyramidal structure		
	(d)	irregular tetrahedral structur	e	
12.	Which	n of the following noble gases i	is the	e most polarized?
	(a)	Radon	(b)	Krypton
	(c)	Xenon	(d)	Helium
13.	Which	n of the following noble gases i	is the	e least polarized?
	(a)	Radon	(b)	Krypton
	(c)	Xenon	(d)	Helium
14.	Which		noble	e gases are used to produce laser
		He and Kr	(b)	Ar and Rn
	(c)	Kr and Ar	(d)	He and Ne
15.	XeO ₄	contains		
	(a)		ainii	ng four electron pairs form a
	(b)	three π -bonds, and the rentrigonal bipyramid	naini	ing five electron pairs form a
	(c)	three π -bonds, and the rensquare-planar structure	naini	ing four electron pairs form a
	(d)	four π -bonds, and the rem	ainir	ng three electron pairs form a

pyramid

- **16.** The structure of XeO_3F_2 is
 - (a) square pyramidal
- (b) pentagonal bipyramidal
- (c) trigonal bipyramidal
- (d) octahedral
- **17.** The structure of XeO₄ is
 - (a) pyramidal

(b) square planar

(c) tetrahedral

(d) that of an irregular tetrahedron

- 18. XeO_3F_2 contains
 - (a) three π -bonds, and the remaining five electron pairs form a trigonal bipyramid
 - (b) two π -bonds, and the remaining six electron pairs form an octahedron
 - (c) four π -bonds, and the remaining four electron pairs form a tetrahedron
 - (d) five electron pairs and three lone pairs
- **19.** The structure of XeF_6 is
 - (a) square pyramidal
- (b) that of a distorted octahedron

(c) pyramidal

(d) trigonal bipyramidal

- **20.** XeF_6 has
 - (a) a tetrahedral structure with one lone pair
 - (b) a trigonal bipyramidal structure with two lone pairs
 - (c) a capped octahedral structure with one lone pair
 - (d) a capped octahedral structure with two lone pairs
- **21.** The structure of XeO_2F_2 is
 - (a) square pyramidal
- (b) trigonal bipyramidal

(c) octahedral

(d) tetrahedral

- 22. XeO₃ contains
 - (a) four π -bonds, and the remaining four electron pairs form a tetrahedron with one corner occupied by a lone pair
 - (b) six electron pairs and two lone pairs
 - (c) two π -bonds, and two corners of a tetrahedron occupied by a lone pair
 - (d) three π -bonds and the remaining four electron pairs form a tetrahedron with one corner occupied by a lone pair
- 23. XeO_2F_2 contains
 - (a) four π -bonds and two σ -bonds
 - (b) two π -bonds, and the remaining five electron pairs form a trigonal bipyramid with one equatorial position occupied by a lone pair

- (c) one π -bond, and the remaining six electron pairs form an octahedron with one position occupied by a lone pair
- (d) three π -bonds, and the remaining four electron pairs form a trigonal bipyramid with two equatorial positions occupied by a lone pair

24. XeOF₄ contains

- (a) six electron pairs forming an octahedron with two positions occupied by lone pairs
- (b) two π -bonds and the remaining six electron pairs, forming an octahedron
- (c) three π -bonds, and the remaining four electron pairs forming tetrahedron
- (d) one π -bond, and the remaining six electron pairs forming an octahedron with one position occupied by a lone pair

25. XeF₄ contains

- (a) four electron pairs and four lone pairs
- (b) six electron pairs and two lone pairs
- (c) three electron pairs and three lone pairs
- (d) two electron pairs and six lone pairs
- 26. Helium is suitable for low-temperature gas thermometry because of its
 - (a) high transition temperature
 - (b) real behaviour
 - (c) low boiling point and near-ideal behaviour
 - (d) high boiling point and high polarizability

27. XeO_6^{4-} contains

- (a) eight electron pairs and no lone pairs
- (b) three electron pairs and three lone pairs
- (c) two electron pairs and six lone pairs
- (d) four electron pairs and four lone pairs

28. The structure of XeF_2 is

- (a) planar triangular
- (b) linear

(c) square planar

- (d) pyramidal
- 29. In which of the following pairs of noble gases is there a large difference in van der Waals radii?
 - (a) Kr and Xe

(b) He and Ne

(c) Ne and Ar

(d) Ar and Kr

- **30.** The reaction of Xe with an excess of F₂ at high pressure and 25°C yields
 - (a) XeF₂
- (b) XeF₄
- (c) XeF₆
- (d) XeF_3

(c) $XeOF_4$ (d) $[XeO_6]^{4-}$

(a) Helium-5 and helium-6 are radioactive nuclides with shorthalf-

	(c)	Helium is the most abundant noble gas in the atmosphere.					
	(d)	Helium-4 has path.	a low molecu	ılar v	viscosity and	a lar	ge mean free
33.	XeF ₄ 1	reacts with NO ₂	to produce				
	(a)	$NO_2F + Xe$		(b)	$N_2 + Xe$		
	(c)	$N_2O + XeO_3$		(d)	$NO_2F_2 + XeC$	O_3	
34.	XeF ₄ 1	eacts with BCl ₃	to produce				
	(a)	BF ₃ and XeCl ₄	-	(b)	BF ₃ , Cl ₂ and	Xe	
	(c)	BF ₃ only		(d)	Cl ₂ only		
35.	Whicl	n of the followin	g statements is	s cor	rect?		
	(a)	Noble gases ha	ive very high i	oniza	ation energy.		
	(b)	Noble gases ha	ive positive ele	ectroi	n affinity.		
	(c)	Fluorides and	oxides of xeno	n are	relatively sta	ble.	
	(d)	Noble gases ca below their inv					peratures and
36.	The fi	rst noble-gas co	mpound, prep	ared	by Bartlett ar	d Lo	hman, was
	(a)	$O_2^+[Pt F_6]^-$ (b) XeF ₆	(c)	$Xe^{+}[Pt F_6]^{-}$	(d)	XeF ₂
37.	Which	n of the followin	g statements is	s inco	orrect?		
	(a)	XeF ₂ is a powe	rful reducing a	igent			
	(b)	XeF ₂ is obtained pressure.	ed by the direc	t rea	ction betweer	r F ₂ a	nd Xe at higł
	(c)	XeF ₂ undergoe	s alkaline hyd:	rolys	is to give O_2 a	nd X	e.
	(d)	XeF ₂ contains f	our electron p	airs a	and four lone	pairs.	•
38.	The sl	low hydrolysis o	of XeF ₄ produc	es			
	(a)	XeO ₂ F ₂		(b)	XeO_3		
	(c)	$XeOF_4$		(d)	$[XeO_6]^{4-}$		
39.	Deep-	sea divers breat	he using a mix	cture	of		
	-	O ₂ and H ₂	O		O ₂ and Kr		
		O ₂ and He			O ₂ and Ar		

31. The slow hydrolysis of XeF₆ gives

(a) XeO_2F_2

(b) XeO₃

(b) ${}_{2}^{4}$ He is obtained from the decay of ${}_{1}^{3}$ H.

32. Which of the following statements is correct?

- **40.** Which of the following statements is correct for helium?
 - (a) Among the noble gases, it has the maximum enthalpy of vapourization.
 - (b) Its van der Waals constants (*a* and *b* values) are large due to strong intermolecular attraction.
 - (c) Liquid helium can be produced above critical temperature and below inversion temperature.
 - (d) The inversion temperature (T_i) of helium is negative at 0°C as well as 100°C at 1 atm.

41. XeF₂ contains

- (a) three electron pairs from a trigonal bipyramid, and three lone pairs in the equatorial position
- (b) four electron pairs from a trigonal bipyramid, and three lone pairs in the equatorial position.
- (c) five electron pairs from a trigonal bipyramid, and three lone pairs in the equatorial position
- (d) three electron pairs from a pentagonal bipyramid, and five lone pairs from a trigonal bipyramid

	Answers			
1. c	2. a	3. d	4. c	5. d
6. b	7. c	8. b	9. d	10. a
11. c	12. c	13. d	14. d	15. a
16. c	17. c	18. a	19. b	20. c
21. b	22. d	23. b	24. d	25. b
26. c	27. a	28. b	29. b	30. c
31. b	32. a	33. a	34. b	35. c
36. c	37. a	38. b	39. c	40. d
41. c				

Hints to More Difficult Problems

The molecule is symmetrical and so the vector sum of the individual dipole moments is zero.

9. Small size and filled shell (1s²)

10.
$$\mu_{JT} = -\left(\frac{\partial T}{\partial p}\right)_{H}$$

- **12.** The van der Waals radius is the greatest and the van der Waals interaction between the atoms maximum for Xe.
- **13.** The van der Waals radius is the smallest and the van der Waals interaction between the atoms minimum for He.

11

Aluminium

• Type 1 •

Choose the correct option. Only one option is correct.

- Aluminium and gallium have nearly the same covalent radius because of the
 - (a) greater shielding effect of the s electrons of gallium atoms
 - (b) poor shielding effect of the s electrons of gallium atoms
 - (c) poor shielding effect of the d electrons of gallium atoms
 - (d) greater shielding effect of the d electrons of gallium atoms
- **2.** The electrolytic reduction of alumina to aluminium by the Hall-Heroult process is carrried out in the presence of
 - (a) NaCl
 - (b) fluorite
 - (c) cryolite which induces the mixture to melt at a temperature lower than the melting point of alumina
 - (d) cryolite which raises the melting point of alumina
- 3. Aluminium is obtained by the
 - (a) reduction of alumina with coke
 - (b) electrolysis of alumina dissolved in molten cryolite
 - (c) reduction of alumina with chromium
 - (d) heating cryolite and alumina
- 4. Bauxite is purified by
 - (a) Hall's process

- (b) Baeyer's process
- (c) Serpeck's process
- (d) all of these
- 5. Anhydrous aluminium chloride is prepared by heating aluminium
 - (a) with concentrated HCl
- (b) with dry chlorine gas

	(c)	with a dilute HCl solution	(d)	carbide with dry HCl gas
6.	Alum	inium burns in dinitrogen at hi	igh t	emperature to produce
	(a)	aluminium nitride	(b)	aluminium azide
	(c)	aluminium nitrate	(d)	alumina
7.	Alum	inium dissolves in aqueous soc	lium	hydroxide to produce
	(a)	Na_3AlO_6	(b)	NaAl(OH) ₄
	(c)	$NaAl_2O_4$	(d)	all of these
8.	Thern	nite is a mixture of iron oxide a	nd	
	(a)	zinc powder	(b)	sodium metal
	(c)	coke	(d)	aluminium powder
9.		aluminothermite process, alum		
		a reducing agent		an oxidizing agent
	(c)	a common solder	(d)	a flux
10.		educing powers of Al, Ga In an		
	` '	In > Ga > Al > Tl Tl > In > Ga > Al		Al > Tl < In > Ga
			` ′	Al > Ga > In > Tl
11.		n of the following forms only co		
	(a) (c)	Thallium Boron	. ,	Aluminium Gallium
10	. ,		` '	
14.		inium chloride ionizes in an aq its dissociation constant is low		as solution because
	` '	it forms a dimer in an aqueou		ution
		it is a Lewis acid		
	(d)	its total hydration energy ex	ceec	ls the ionization energy of the
		system		· · · · · · · · · · · · · · · · · · ·
13.	Which	n of the following metals does i	not s	how an inert-pair effect?
	(a)	Indium (b) Aluminium	(c)	Gallium (d) Thallium
14.	Dural	umin is an alloy of Al with		
		Mn and Mg		Mg and Ni
	(c)	Mg, Mn and Cu	(d)	Mg, Cr and Mn
15.		n of the following statements is		
		The hydroxides of boron and		-
	(b)	The hydroxide of boron is amphoteric.	basi	c while that of aluminium is
	(c)	The hydroxide of aluminium	is m	ore acidic than that of boron
	(d)	•		ic while that of aluminium is
	(4)	11.5 11, 010/110C O1 DO1011 15	aciu	uint of arainmualli 15

amphoteric.

- 16. The salts of aluminium and zinc can be distinguished from each other by
 - (a) the flame test
 - (b) the borax bead test
 - (c) the cobalt nitrate test
 - (d) treatment with an excess of an NaOH solution
- 17. Which of the following reactions is not a part of the Goldschmidt aluminothermic process?
 - (a) $Fe_2O_3 + 2Al \longrightarrow Al_2O_3 + 2Fe$
 - (b) $Cr_2O_3 + 2Al \longrightarrow Al_2O_3 + 2Cr$
 - (c) $3Mn_3O_4 + 8Al \longrightarrow 4Al_2O_3 + 9Mn$
 - (d) $3ZnO + 2Al \longrightarrow Al_2O_3 + 3Zn$
- 18. In the Al₂Cl₆ dimer,
 - (a) all the Al—Cl bonds are equivalent
 - (b) three Al—Cl bonds are equivalent and three are not
 - (c) two Al—Cl bonds are equivalent and four are not
 - (d) two Al—Cl bonds are not equivalent and four are equivalent
- **19.** Which of the following gases is evolved when aluminium is boiled in a concentrated alkali solution?
 - (a) Only O₂

(b) A mixture of O₂ and H₂

(c) Only H₂

- (d) All of these.
- 20. Aluminium is thermodynamically stable because
 - (a) a thin layer of aluminium oxide is formed on its surface, protecting it from further attack
 - (b) it has a low hydration power
 - (c) its ionization energy is greater than that of copper
 - (d) it has a low standard reduction potential
- 21. During the extraction of aluminium,
 - (a) a graphite-lined steel tank serves as an anode and a graphite rod as a cathode
 - (b) a graphite-lined steel tank acts as a cathode and a graphite rod as an anode
 - (c) cryolite acts as an anode as well as a cathode
 - (d) a graphite rod acts as an anode and molten aluminium as a cathode
- 22. At low temperatures,
 - (a) AlCl₃ exists as a close-packed lattice of Cl⁻ and Al³⁺, occupying tetrahedral holes

- (b) AlCl₃ exists as a close-packed lattice of Cl⁻ and Al³⁺, occupying trigonal holes
- (c) AlCl₃ exists as a close-packed lattice of Cl⁻ and Al³⁺, occupying octahedral holes
- (d) AlCl₃ reacts with Al to produce AlCl
- 23. Aluminium hydride is best made by the reaction between
 - (a) Li[AlH₄] and AlCl₃
- (b) Li and CaH₂
- (c) LiCl and Li[AlH₄]
- (d) Li[AlH₄] and Cl₂
- **24.** In the structure of α -AlCl₃, each Al participates in
 - (a) four bridges

(b) six bridges

(c) three bridges

- (d) twelve bridges
- **25.** Which of the following statements is incorrect?
 - (a) Cryolite is added to alumina to lower the fusion temperature and make the melt a good conductor.
 - (b) Aluminium forms an $[AlF_6]^{3-}$ ion.
 - (c) Anhydrous aluminium chloride cannot be prepared by heating hydrated aluminium chloride (AlCl₃ · 6H₂O) because of the hydrolysis of AlCl₃.
 - (d) AlCl₃ is a high-melting solid while AlF₃ is a low-melting volatile solid.

• Type 2 •

Choose the correct options. More than one option is correct.

- **26.** Which of the following alloys contain aluminium?
 - (a) Devarda's alloy

(b) Invar

(c) Rolled gold

- (d) Type metal
- 27. Which of the following double salts form acidic solutions when dissolved in water?
 - (a) Fusion mixture

(b) Alum

(c) Mohr's salt

- (d) Carnallite
- 28. Potash alum is used as a
 - (a) disinfectantant
 - (b) water softener
 - (c) mordant in the textile industry
 - (d) fibre in the polymer industry

(a) Fluorspar(c) Mica

(b) Feldspar

(d) Carborundum

29. Which of the following minerals contain aluminium?

30.	Alum	inium is not purified by		
	(a)	Baeyer's process	(b)	Ostwald's process
	(c)	Hoope's process	(d)	Serpeck's process
31.		general formula of alums is ving combinations are alums?	XIY	$\chi^{\text{III}}(SO_4)_3 \cdot 12H_2O$. Which of the
	(a)	K^{I} and Al^{III}	(b)	K^{I} and Cr^{III}
	(c)	$\mathrm{NH_4^+}$ and $\mathrm{Al}^{\mathrm{III}}$	(d)	Na $^{\scriptscriptstyle +}$ and Tl $^{\rm III}$
32.	Which	n of the following statements ar	re co	rrect?
	(a)	All alums crystallize in thisomorphous series of double		octahedral form and produce s.
	(b)	All alums are double salts a crystallization.	and	have large values of water of
	(c)	Aluminium sulphate is useful	as a	mordant in dying and printing.
	(d)	Alums are used in coagulatio tanning of leather.	n, tł	ne purification of water, and the
33.	Alum	inium reacts with hot concentra	ated	NaOH solution to produce
	(a)	$NaAlO_2 + H_2$	(b)	$Na_3AlO_2 + H_2$
	(c)	$Na_3AlO_3 + H_2$	(d)	$Na_2AlO_4 + H_2$
34.	Two	lifferent forms of alumina are k	cnov	vn. These are
	(a)	α -Al ₂ O ₃	(b)	γ -Al ₂ O ₃
	(c)	β -Al ₂ O ₃	(d)	δ -Al ₂ O ₃
35.	Which	n of the following trivalent cation	ons o	can form alums?
	(a)	Mn^{3+}	(b)	Co ³⁺
	(c)	Fe ³⁺	(d)	Cr 3+
36.	Which	n of the following statements ar	re co	rrect?
	(a)	AlCl ₃ reacts with Grignard's r	eage	ent to form trialkyl aluminium.
	(b)	Triethyl aluminium is used to	pre	pare the Ziegler–Natta catalyst.
	(c)			ers and has three-centre bonds a Al and C in the Al—C—Al
	(d)	AlCl is covalent and very stab	le.	

Aluminium 2-117

			_	
1. c	2. c	3. b	4. d	5. b
6. a	7. b	8. d	9. a	10. d
11. c	12. d	13. b	14. c	15. d
16. c	17. d	18. d	19. c	20. a
21. b	22. c	23. a	24. b	25. d
26. a, c	27. b, c	28. a, b	29. b, c	30. a, b, d
31. a, b, c	32. a, b	33. a, c	34. a, b	35. a, b, c, d
36. a, b, c				

Answers

Hints to More Difficult Problems

- 10. Consider the standard electrode potentials.
- 11. Being a nonmetal, boron forms covalent compounds.
- 13. The 3s² electron is not inert in Al, in contrast to the other metals in its series.
- **15.** $B(OH)_3 + 2H_2O \rightleftharpoons H_3O^+ + [B(OH)_4]^-$ Al(OH)₃ does not undergo an acidic reaction in water.
- **16.** Upon conducting the charcoal-block oxidizing-flame test using $Co(NO_3)_2$, we get coloured beads. Al gives blue $Al_2O_3 \cdot CoO$ and Zn gives green $ZnO \cdot CoO$. This is how salts of Al and Zn can be distinguished.

Ч

12

Tin and Lead

• <u>Type 1</u> •

Choose the correct option. Only one option is correct.1. Tin is extracted from its ore, cassiterite, by

	(a)	electrolytic reduction	(b)	carbon-monoxide reduction
	(c)	carbon reduction	(d)	the aluminothermic process
2.	Tin is	not refined by		
	(a)	liquation	(b)	zone refining
	(c)	poling	(d)	any of these
3.		rately concentrated nitric stannic acid, which is represe		reacts with tin to produce
		$H_2Sn_5O_{11}$		$H_2Sn_5O_2$
	(c)	$H_2Sn_4O_{10}$	(d)	$H_2Sn_4O_6$
4.	Tin (I	I) oxide is not prepared by		
	(a)	boiling a stannous chloride	solutio	on with Na ₂ CO ₃
	(b)	heating tin oxalate (SnC ₂ O ₄)	in the	absence of air
	(c)	heating tin hydroxide in air		
	(d)	heating tin (IV) oxide in air		
5.	The g	eometry of SnCl ₂ is		
	(a)	linear	(b)	pyramidal
	(c)	V-shaped	(d)	plane triangular
6.		n of the following crystals is concentrated HCl?	forme	ed when SnCl ₂ · 2H ₂ O is treated
	(a)	$SnCl_4 \cdot 5H_2O$	(b)	$HSnCl_3 \cdot 2H_2O$
	(c)	H ₂ SnCl _c · 8H ₂ O	(d)	H ₂ SnCl ₄ · 4H ₂ O

- 7. Tin (II) sulphide dissolves in yellow ammonium sulphide to give (a) thiostannate $[(NH_4)_2SnS_3)]$ (b) thiostannite $[(NH_4)_2SnS_4)]$ (d) a mixture of SnS₂ and H₂SnS₆ (c) tin (IV) sulphide 8. The shape of SnCl₄ is (a) square planar (b) irregular tetrahedral (c) square pyramidal (d) tetrahedral **9.** Which of the following is correct? (a) $SnS_2 + Na_2S \longrightarrow Na_2SnS_3$ (b) $SnS_2 + (NH_4)_2S \longrightarrow (NH_4)_2SnS_3$ (c) $SnS_2 + NaOH \longrightarrow Na_2SnO_3 + Na_2SnS_3 + H_2O$ (d) $SnS_2 + HNO_3 \longrightarrow Sn(NO_3)_2 + H_2SO_4 + SO_2 + H_2O$ **10.** Which of the following statements is incorrect? (a) An acidified solution of tin (II) chloride becomes cloudy on dilution with water. (b) Hydrogen sulphide reacts with an acidified solution of tin (II) chloride to produce a white precipitate. (c) Tin (IV) sulphide dissolves in yellow ammonium sulphide to produce a clear solution, which on acidification gives a yellow precipitate and hydrogen sulphide. (d) The type of allotropy in which two allotropes are equally stable at the transition temperature is called enantiotropy. 11. Which of the following compounds is a solid? (a) SnI_4 (b) CCl₄ (d) SiBr₄ (d) None of these 12. Among the following, the most stable dihalide is (b) SiX₂ (a) CX₂ (c) GeX₂ (d) SnX_2
 - **13.** Which of the following statements is incorrect?
 - (a) Anhydrous stannous chloride is prepared by heating hydrated stannous chloride crystals in a stream of hydrogen chloride.
 - (b) Due to hydrolysis by water, stannous chloride forms a white precipitate of basic stannous chloride [Sn(OH)Cl \cdot H₂O].
 - (c) The hydrolysed product of stannous chloride does not go into solution in the presence of HCl.
 - (d) The hydrolysed product of stannous chloride goes into the solution in the presence of dilute HCl.
 - 14. Which of the following statements is incorrect for tin (IV) chloride?
 - (a) It is obtained by passing hydrogen chloride over heated tin

(b)	It is a colourless liquid whydrolysis.	nich	fumes in moist air owing to			
(c)	dilute, and in the presence of a little water it is possible to obtain					
(d)	It is ionic, and presumably co	ntair	ns $[Sn(H_2O)_4]^{4+}$ ions.			
Tin pr	oduces a low crackling sound	on b	ending. This is known as tin			
			cry			
(c)	plague	(d)	sound			
		or th	e tin that exhibits enantiotropy			
(a)	Grey tin (α) $\stackrel{500 \text{ K}}{\longleftarrow}$ britt	le tir	$434 \text{ K} \longrightarrow \text{ white tin } (\beta)$			
(b)	Brittle tin $\stackrel{286.5 \text{ K}}{\longleftarrow}$ white t	in (β	$) \stackrel{200 \text{ K}}{\longrightarrow} \text{grey tin } (\alpha)$			
(c)	Grey tin (α) $\stackrel{286.5 \text{ K}}{\longleftarrow}$ whit	e tin	$(\beta) \xleftarrow{434 \text{ K}} \text{ brittle tin}$			
(d)	White tin (β) $\xrightarrow{400 \text{ K}}$ gre	y tin	286.5 K brittle tin			
Which	n of the following halides is the	e leas	it stable?			
	· ·		SnBr ₄ (d) SnCl ₄			
Massi	cot is prepared by heating					
		(b)	cerussite			
(c)	minium	(d)	litharge			
In wh	ich of the following acids is lea	ıd ea	sily soluble?			
(a)	Sulphuric acid	(b)	Nitric acid			
(c)	Hydrochloric acid	(d)	Acetic acid			
Comn	non solder is an alloy of					
		` '	Sn and Pb			
(c)	Sn and Cu	(d)	Sn and Zn			
		<i>a</i> >				
		. ,	Cu, Sn and Pb			
			Fe, Sn and Pb			
			basic lead carbonate			
	(c) (d) Tin pr (a) (c) Which amon (a) (b) (c) (d) Which (a) Massi (a) (c) In wh (a) (c) Comm (a) (c) Type (a) (c) The le (a)	hydrolysis. (c) It is not completely hydroly dilute, and in the presence of solid SnCl ₄ · 5H ₂ O. (d) It is ionic, and presumably co Tin produces a low crackling sound (a) poison (c) plague Which of the following is correct for among its three allotropic forms? (a) Grey tin (α) 500 K britt (b) Brittle tin 286.5 K white the complete white tin (β) 400 K grey tin (α) White tin (β) PbI ₂ Which of the following halides is the (a) PbI ₄ (b) PbI ₂ Massicot is prepared by heating (a) tin in the presence of CO ₂ (c) minium In which of the following acids is lead (a) Sulphuric acid (c) Hydrochloric acid Common solder is an alloy of (a) Sn and Sb (c) Sn and Cu Type metal is an alloy of (a) Zn, Sb and Pb (c) Sb, Sn and Pb	(c) It is not completely hydrolysed dilute, and in the presence of a lit solid SnCl ₄ · 5H ₂ O. (d) It is ionic, and presumably contain Tin produces a low crackling sound on b (a) poison (b) (c) plague (d) Which of the following is correct for the among its three allotropic forms? (a) Grey tin (α) 500 K brittle tin brittle tin complete tin complet			

23.	Which of the following compounds is known as chrome red, in which the oxidation number of $Cr = +6$?				n the				
		PbCrO ₃ · Pb			(b)	PbCrO₄ · Cr	O_3		
		PbCrO₄ · Pb		, =		PbCrO ₄ · Pb			
24.	Which	n of the follow	wing of chi	compounds	is kr .r]3d	nown as chroi	ne y	ellow in w	hich
	(a)	$PbCrO_4$			(b)	PbCr ₂ O ₇			
	(c)	$PbCrO_4 \cdot Cr$	$_{2}O_{3}$		(d)	Pb_2CrO_4			
25.	Lead	sulphate is so	oluble	e in a solutio	n of o	concentrated			
	(a)	sulphuric ac	cid		(b)	hydrochlori	c aci	d	
	(c)	ammonium	acet	ate	(d)	lead nitrate			
26.	In lea	d storage bat	teries	s, we use					
	(a)	Pb as the ca	thod	e and PbO ₂ a	s the	anode			
	(b)	PbO ₂ as the	cath	ode and Pb a	s the	anode			
	(c)	a lead plate	as th	ne cathode an	ıd Sn	O_2 as the ano	de		
	(d)	one lead die	oxide	plate as the	catho	ode and anotl	ner a	s the anod	e
27.		ead accumula on used in th			dary	cell and is re	char	gable. The	acid
	(a)	H ₂ SO ₄ and 0	CH_3C	CO_2NH_4	(b)	H_2SO_4			
	(c)	H ₂ SO ₄ and l	PbSC	O_4	(d)	H ₂ SO ₄ and 0	CH ₃ C	CO_2H	
28.	The le	east melting r	netal	among the f	ollov	ving is			
		tin		carbon		germanium	(d)	lead	
29.	The re	ed and yellov	v for	ms of lead (II	() oxi	de are knowr	ı resj	pectively a	.S
		litharge and				litharge and	-		
	(c)	massicot an	d vei	rmilion	(d)	red lead and	d hea	ıvy spar	
30.	Whicl	n of the follow	ving	compounds	disp	roportionates	easi	ily?	
		$PbCrO_4$		Pb_3O_4		PbCl ₂		PbI ₄	
31.	On de	escending the	groi	up of which l	ead i	is a member,	the		
		_	_	_		of lead decrea			
		-				of lead increa			
		tendency of							
		-		-		increases u	p to	Ge and	then
22	Mhial		vina	ic known ac	mos	nic gold?			
34.		n of the follov As ₂ O ₃	vnig	15 KIOWII dS		PbS			
	(a) (c)					PbI_2			
	(८)	21102			(u)	- 012			

			ar.	2			
	(a)	$PbCl_6^{2-}$ (b) Pb(OH) ₆	(c)	Sn(OH) ₆	(d)	all of these
40.		xpansion of the			-		
		perchlorate, Ph			periodate, F		
		acetate, Pb(CH			acetate, Pb(-	
39.		nly stable lead (O	0.0\
	(d)	lead acetate an	id NaOH soluti	ion			
		lead acetate an	_				
	(b)	lead carbonate	and CO ₂				
	(a)	lead acetate an	nd Na ₂ CO ₃ solu	tion			
38.	White	lead (basic lead	d acetate) is pre	pare	d by the reac	tion b	etween
	(c)	Lead carbonate	e		Diethyl lead		
٠,,		Lead tetraaceta	0 0		Tetraethyl l	ead	
37.	` '	is used as an an	-	` '	9	` "/	4
	-		o) PbI ₂			(d)	PbCl ₂
36.		ite solid insolub vely to the gold					ater responds
	` '	addition of tin					
		removal of imp		b			
	` /	removal of tin			er		
		conversion of l					
35.	"Softe	ening of lead" m	neans the				
		self-reduction	•		self-oxidation		
J4.		is extracted fron lead chamber _l	•	(b)	Pattinson p	rocess	
24			-	(u)	100 - 2100	\mathcal{O}_3	
		$Pb(OH)_2 \cdot 2PbO_3 \cdot PbSO_4$			$PbO \cdot 2PbC$		$H_3COO)_2 \cdot PbS$
33.		n of the followin	~			Dl _e (CT	LCOO) DI-C
	XA71 · 1	(4) (1)	. 1	1	1 12		

• *Type 2* •

Choose the correct options. More than one option is correct.

41. In which of the following reactions can $SnCl_2/Sn^{2+}$ act as a reducing agent?

(a)
$$SnCl_2 + I_2 \longrightarrow SnCl_4 + SnI_4$$

(b) $SnCl_2 + HgCl_2 \longrightarrow SnCl_4 + Hg_2Cl_2$

(a) $MnSO_4 + PbO_2 + HNO_3 \longrightarrow HMnO_4 + PbSO_4 + Pb(NO_3)_2 + H_2O$

(b) $Cr(OH)_3 + KOH + PbO_2 \longrightarrow K_2CrO_4 + K_2PbO_2 + H_2O$

50. Which of the following show the right products?

(c) $Pb_3O_4 + HCl \longrightarrow PbCl_2 + Cl_2 + H_2O$ (d) $Pb + H_2O + O_2 \longrightarrow Pb(OH)_4$

(c) $MnO_4^- + H^+ + Sn^{2+} \longrightarrow Mn^{2+} + Sn^{4+} + H_2O$

Answers

1. c	2. b	3. a	4. d	5. c
6. b	7. a	8. d	9. b	10. b
11. a	12. d	13. c	14. a	15. b
16. c	17. a	18. b	19. d	20. b
21. c	22. a	23. d	24. a	25. c
26. a	27. b	28. a	29. b	30. d
31. a	32. c	33. a	34. c	35. c
36. d	37. b	38. a	39. b	40. d
41. a, b, c, d	42. a, b, d	43. b, c, d	44. a, b	45. a, c, d
46. a, c	47. a, b, d	48. a, b, c	49. a, c, d	50. a, b, c

Hints to More Difficult Problems

- 7. $SnS + (NH_4)_2S_2 \longrightarrow (NH_4)_2SnS_3$
- **12.** Due to the inert-pair effect.
- 17. Pb^{4+} oxidizes I^{-} to I_2 . $PbI_4 \longrightarrow PbI_2 + I_2$
- **25.** $PbSO_4(s) + 2CH_3COONH_4 \longrightarrow Pb(CH_3COO)_2 + (NH_4)_2SO_4$ insoluble soluble
- 28. Tin does not have a packed structure.
- **30.** See the hint to Q. 17.
- 31. Due to the strong inert-pair effect in lead
- **34.** (a) $2PbS + 3O_2 \longrightarrow 2PbO + 2SO_2$
 - (b) $PbS + 2O_2 \longrightarrow PbSO_4$
 - (c) $PbS + 2PbO \longrightarrow 3Pb + SO_2$
 - (d) $PbS + 2PbSO_4 \longrightarrow Pb + 2PbO + 3SO_2$ Reactions (b) and (d) play a minor role.
- 44. The oxidation number of Sn is +2 in SnCl₂ and +4 in SnCl₄.

45. Like SnCl₂, all are V-shaped.

13

Copper, Silver and Gold

• Type 1 •

Choose the correct option. Only one option is correct.

- 1. The second and third ionization energies of copper are much lower than those of alkali metals. This accounts for the
 - (a) existence of colourless diamagnetic ions and complexes in the I oxidation state
 - (b) existence of coloured paramagnetic ions and complexes in the II and III oxidation states
 - (c) linear geometry of CuO
 - (d) formation of various alloys
- 2. Matte contains
 - (a) Cu₂S, FeS and silica
- (b) Cu₂S, FeO and silica
- (c) Cu₂S, CuO and silica
- (d) Cu₂S, Cu₂O and silica
- **3.** By which of the following reactions is blister copper obtained?
 - (a) $Cu_2S + FeO \longrightarrow 2Cu + FeO$
 - (b) $Cu_2S + FeS \longrightarrow 2Cu + FeS_2$
 - (c) $Cu_2S + 2Cu_2O \longrightarrow 6Cu + SO_2$
 - (d) $Cu^{2+} + Fe \longrightarrow Fe^{2+} + Cu$
- **4.** The most stable oxidation states of Group 11 metals in their compounds are
 - (a) Cu(+2), Ag(+1) and Au(+3)
 - (b) Cu(+2), Ag(+1) and Au(+1)
 - (c) Cu(+1), Ag(+2) and Au(+3)
 - (d) Cu(+2), Ag(+2) and Au(+2)

(d) self-reduction of the oxide and sulphide of copper7. In an aqueous solution, Cu(+1) salts are unstable because

(b) the change in free energy of the overall reaction is zero
(c) they disproportionate easily to the Cu and Cu(+2) states
(d) they disproportionate easily to the Cu(+2) and Cu(+3) states

(a) $CuAl_6(PO_4)_4(OH)_8 \cdot 4H_2O$

6. Copper is extracted from its sulphide ore by(a) the carbon reduction process(b) a displacement reaction(c) electrolytic reduction

(a) Cu(+1) has a 3d ¹⁰ configuration

(c) Cu₅FeS₄

5. Which of the following compounds is known as bornite or peacock's ore?

(b) CuFeS₂

(d) YBa₂Cu₃O₇

8.	The c	rystal structure of copper is a				
	(a)	simple cubic lattice	(b)	bcc lattice		
	(c)	ccp lattice	(d)	hcp lattice		
9.	A few	Cu(I) salts are coloured. The c	colou	ar arises from		
	(a)	d-d transition				
	(b)	charge-transfer spectra				
	(c)	the large wavelengths of the	rays	absorbed by the solutions		
	(d)	none of these				
10.		n of the following reaction metrically?	ns	is used to estimate copper		
	(a)	$Cu^{2+} + 4NH_3 \longrightarrow [Cu(NH_3)]$	4]2+			
	(b)	$2Cu^{2+} + 4I^{-} \longrightarrow 2CuI + I_2$				
	(c) $2Cu^{2+} + 4CN^{-} \longrightarrow CuCN + (CN)_2$					
	(d)	$2Cu^{2+} + SO_3^{2-} + 2SCN^- + H_2C$		\rightarrow 2CuSCN + H ₂ SO ₄		
11.	When	a CuSO ₄ solution is treated w	ith I	$K_4[Fe(CN)_6]$, a brown precipitate		
	(a)	$Cu_2[Fe(CN)_6]_2$	(b)	$Cu[Fe(CN)_6]$		
	(c)	$Cu_2[Fe(CN)_6]$	(d)	$Cu_2[Cu(CN)_6]$		
	is obta	ained.				
12.				an excess of KCN, a colourless onfiguration is obtained. The salt		
	(a)	$K_2[Cu_2(CN)_4]$	(b)	$K_2[Cu(CN)_6]$		
	(c)	$K_2[Cu(CN)_4]$	(d)	$K_4[Cu(CN)_{\epsilon}]$		

13. CuSO₄ reacts with an excess of hypo solution to produce a Cu^I complex

	(a)	$Cu_2S_2O_3$			(b)	$Na_3[Cu(S_2C)]$	$(0,0)_{3}$		
	(c)	$Na_4[Cu_6(S_2C)]$	$(3)_{5}$		(d)	$Cu_4[Na_6(S_2)]$	$O_3)_5$		
14.	Durin	g the extracti	on o	f copper	from cha	lcopyrites, i	ron is	removed	l as
	(a)	$Fe_3(PO_4)_2$	(b)	Fe_2O_3	(c)	$Fe_2(SiO_3)_3$	(d)	$FeSiO_3$	
15.	Bliste	r copper is							
	(a)	electrolytica	ılly re	efined co	pper				
	(b)	a mixture of	f imp	ure copp	er and s	ilver			
	(c)	copper cont	ainin	g 2% im	purity				
	(d)	present in the	ne an	ode muc	l in an el	ectrolytic pro	ocess		
16.	A Cus	SO ₄ solution	reacts	s with ar	Na ₂ CO ₃	solution to	produ	ice	
		CuCO ₃			(b)	$CuCO_3 \cdot Cu$	ı(HC	$O_3)_2$	
	(c)	$CuCO_3 \cdot Cu$	(OH))2	(d)	Cu ₂ O			
17.	Which	n of the follow	ving	statemer	nts is cor	rect? Cu (II)	acetat	e is	
	(a)	paramagnet	ic, di	meric ar	nd hydra	ted			
	(b)	paramagnet	ic, m	onomeri	c and hy	drated			
	(c)	diamagnetic	c, din	neric and	l not hyd	rated			
	(d)	paramagnet	ic, te	trameric	and hyd	rated			
18.	Which	n of the follow	ving	is polym	eric in th	e vapour sta	ate?		
	(a)	Cu(CH ₃ CO	$)_2$		(b)	CuF_2			
	(c)	CuCl			(d)	$CuSO_4$			
19.	The re	eactivity of co	ppei	is low b	ecause o	f its			
	(a)	high enthal	by of	sublima	tion and	low ionizati	on en	ergy	
	(b)	high enthal	py of	sublima	tion and	high ionizat	ion e	nergy	
	(c)	low enthalp	y of	sublimat	ion and l	nigh ionizati	on en	ergy	
	(d)	low enthalp	y of	sublimat	ion and l	ow ionizatio	n ene	ergy	
20.	The m	nelting point	of co	pper is h	igher tha	n that of zin	c bec	ause	
	(a)	copper has a	a bcc	structur	e				
	(b)	the atomic v	olun'	ne of cop	per is hig	gher			
	(c)	the d electro	ons o	f copper	are invol	ved in meta	llic bo	onding	
	(d)	the s as we bonding	ell as	d electi	rons of c	copper are i	nvolv	red in me	etallic
21.	Whicl	n of the follow	wing	compou	nds is a s	uperconduc	tor?		
		CaTiO ₃	J	1		$Cu_3(AsO_3)_2$		CH ₃ COC)) ₂
		YBa ₂ Cu ₈ O ₇₋	-x			YBa ₂ Cu ₃ O ₇		-	

- **22.** The ionization energy of copper is higher than that of potassium though both have a 4s configuration because the d electrons in copper
 - (a) form a poor shield, making copper smaller
 - (b) form a poor shield, making copper bigger
 - (c) are strongly shielded, making copper smaller
 - (d) are strongly shielded, making copper bigger
- 23. When a solution of KI is added to a sample of Cu²⁺ ions
 - (a) I^- oxidizes Cu^{2+} to CuI, and is reduced to I_2
 - (b) I^- reduces Cu²⁺ to CuI, and is oxidized to I_2
 - (c) I^- oxidizes Cu^{2+} to Cu, and is reduced to I_2
 - (d) I reduces Cu²⁺ to Cu, and is oxidized to I₂
- 24. The melting points of Cu, Ag and Au follow the order
 - (a) Cu > Ag > Au

(b) Cu > Au > Ag

(c) Au > Ag > Cu

- (d) Ag > Au > Cu
- 25. Which of the following statements is correct?
 - (a) The tetrahedral $[CuCl_4]^{2-}$ ion is green.
 - (b) The tetrahedral $[CuCl_4]^{2-}$ ion is blue, and the square-planar $[CuCl_4]^2$ ion is red.
 - (c) The tetrahedral $[CuCl_4]^{2-}$ ion is orange, and the square-planar $[CuCl_4]^{2-}$ ion is yellow.
 - (d) The tetrahedral as well as square-planar $[{\rm CuCl_4}]^{2-}$ ions are bright yellow.
- 26. When heated to above 800°C, Cu(NO₃)₂ yields
 - (a) $N_2 + CuO$

(b) $Cu_2O + CuO + N_2O$

(c) $Cu + N_2$

- (d) Cu₂O
- **27.** The blue colour of a cupric salt solution in water is due to the formation of a hydrated cupric ion. The hydrated ion is
 - (a) octahedral with two long bonds trans to each other and four short bonds trans to each other
 - (b) tetrahedral with two long bonds trans to each other and two short bonds cis to each other
 - (c) octahedral with two long bonds cis to each other and four short bonds trans to each other
 - (d) octahedral with two short bonds cis to each other and four long bonds trans to each other
- 28. Silver bromide dissolves in a hypo solution to produce
 - (a) $Ag_2S_2O_3$

(b) $[Ag(S_2O_3)_2]^{-}$

(c) $[Ag(S_2O_3)_2]^{3-}$

(d) $[Ag_6(S_2O_3)_5]^{4+}$

29.		is insoluble in water but re on because	adily	soluble in a dilute ammonia				
	(a)	NH ₃ is a better solvent than water						
	(b)	NH ₃ is a stronger base than w	NH ₃ is a stronger base than water					
	(c)	Ag + forms a complex with N	H ₃ to	produce [Ag(NH ₃) ₂] +				
	(d)	the dipole moment of water is	s hig	her than that of NH ₃				
30.	metal	is known as	e inv	volving KCN, air and an active				
		Pattinson's process						
		the amalgamation process						
		the McArthur–Forrest process	S					
	(d)	Parke's process						
31.	Which	n of the following ions disprop	ortio	nates in water?				
	(a)	Au ³⁺	(b)	$[Au(CN)_4]^-$				
	(c)	[AuCl ₄]	(d)	Au ⁺				
32.	Gold	and platinum dissolve in aqua	regia	a to produce respectively				
			_	H[AuCl ₄] and H ₂ [PtCl ₆]				
	(c)	$H_2[AuCl_6]$ and $H[PtCl_4]$	(d)	H ₂ [AuCl ₆] and [PtCl ₆]				
33.	Chlor	oauric acid reacts with sodium	hyd	roxide to produce				
	(a)	$Au(OH)_3$	(b)	$[Au(OH)_4]^{2-}$				
	(c)	Au_2O_3	(d)	K[AuCl ₄]				
34.	The m	netallic radius of gold is almos	t ide	ntical with that of silver because				
	(a)	transition metal contraction						
	(b)	the same crystal structure of s	silvei	and gold				
	(c)	the high electropositive chara	cter	of gold in comparison to silver				
	(d)	the effect of lanthanide contra	action	n in gold				
35.		chloride fuses with sodium ca		-				
	(a)	Ag_2CO_3		Ag_2O				
	(c)	AgOH	(d)	Ag				
36.		is extracted from Ag ₂ S by						
		fusing it with KCl, and electron	olysi	ng the melt				
	(b)	reducing it with zinc						

(c) treating it with sodium cyanide followed by zinc

(d) roasting it and reducing the resultant product by smelting

37.	Silver	is refined by		
		cupellation	(b)	poling
	(c)	the van Arkel method	(d)	liquation
38.	The or	res of Ag and Au are concentra	ted	using their solubility in
	(a)	HCl	(b)	HNO_3
	(c)	H ₂ SO ₄	(d)	KCN
39.	Zinc is	s used to extract silver		
	(a)	by solvent extraction from mo	olten	lead in Parke's process
		by solvent extraction from mo		
		by carbon monoxide reductio		*
	(d)	by solvent extraction from mo	olten	iron in the LD process
40.		electrolytic refining of silver, t		
		Zn, Ag and Au		Zn, Cu, Ag and Au
	()	Au		Cu, Ag and Au
41.		n of the following silver salts is		
		AgClO ₄		Ag_2SO_4
	(c)	AgF	(d)	AgNO ₃
42.	comp			sing an excess of KCN, soluble which have 10 d electrons in Ag
	(a)	K ₂ [Ag(CN) ₄] and K ₃ [Ag(CN) ₄]	
	(b)	K[Ag(CN) ₂] and K[Au(CN) ₂]		
	(c)	K ₂ [Ag(CN) ₃] and K ₂ [Au(CN) ₃	.]	
	(d)	$K_4[Ag_6(CN)_{10}]$ and $K_4[Au_6(CN)_{10}]$	J) ₁₆]	
43.		n of the following reactions wi lting point?	ll oc	cur on heating AgNO ₃ to above
		$2AgNO_3 \longrightarrow 2Ag + 2NO_2 +$		
		$2AgNO_3 \longrightarrow 2Ag + N_2 + 3C$	2	
		$2AgNO_3 \longrightarrow 2AgNO_2 + O_2$		
	(d)	$2AgNO_3 \longrightarrow Ag + NO + O_2$		
44.		AgNO ₃ is strongly heated, the	_	
		N ₂ O and N ₂		N_2O_5 and O_2
	(c)	N_2O_3 and O_2	(d)	NO_2 and O_2
45.	In pho	otography AgBr is mainly used	as	
	` '	a fixer		an emulsion
	(c)	a light-sensitive material	(d)	a developer

- **46.** AgCl dissolved in an excess of solutions of NH₃, KCN and Na₂S₂O₃ produces complex ions. They are, respectively,
 - (a) $[Ag(NH_3)_2]^+$, $[Ag(CN)_2]^-$ and $[Ag(S_2O_3)_2]^{3-}$
 - (b) $[Ag(NH_3)_2]^{2+}$, $[Ag(CN)_2]^{3-}$ and $Ag_4(S_2O_3)_5]^{2-}$
 - (c) $[Ag(NH_3)_2]^{2+}$, $[Ag(CN)_2]^+$ and $[Ag_2(S_2O_3)_2]^{2-}$
 - (d) $[Ag(NH_3)_4]^+$, $[Ag(CN)_4]^{3-}$ and $[Ag_2(S_2O_3)_2]^{2-}$
- 47. Gold (III) chloride exists as a planar Au₂Cl₆ molecule in
 - (a) only the solid state
 - (b) only the vapour state
 - (c) only the liquid state
 - (d) the solid as well the vapour state

• *Type 2* •

Choose the correct options. More than one option is correct.

- **48.** Which of the following are correctly matched?
 - (a) Turquoise \longrightarrow (CuAl₆PO₄)₄(OH)₈ · 4H₂O
 - (b) Peacock ore \longrightarrow Cu₄FeS₂
 - (c) Malachite \longrightarrow CuCO₃ · Cu(OH)₂
 - (d) Chalcopyrites \longrightarrow CuFeS₂
- **49.** Which of the following are correctly matched?
 - (a) Brass \longrightarrow Cu, Sn
 - (b) Nickel silver \longrightarrow Cu, Ni, Zn
 - (c) Phosphor bronze \longrightarrow Cu, Sn, P
 - (d) Fool's gold \longrightarrow CuS₂
- **50.** Which of the following statements are correct in connection with the extraction of silver?
 - (a) Silver is obtained as a by-product in the extraction of copper, lead and zinc.
 - (b) Silver is obtained from the anode slime formed in the electrolytic refining of copper and zinc.
 - (c) Zinc is used to extract silver by solvent extraction from molten lead in Parke's process.
 - (d) None of these

- 51. Which of the following does not disproportionate?
 - (a) Cu +
- (b) Au^{3+}
- (c) Cu²⁺
- (d) Au+
- 52. In which of the following complexes do copper ions show an oxidation state of +1?
 - (a) $[Cu(CN)_4]^{3-}$ (b) $[CuCl_2]^{-}$ (c) $[CuCl_3]^{2-}$ (d) $[CuCl_4]^{3-}$
- 53. Which of the following are correctly matched?
 - (a) Schweitzer's reagent ----- An ammoniacal solution of cupric hydroxide
 - (b) Bordeaux mixture \longrightarrow CuSO₄ and Ca(OH)₂
 - (c) Semiconductor $\longrightarrow YBa_2Cu_3O_7$
 - (d) Horn silver → AgNO₃
- 54. During the extraction of Ag and Au using a KCN solution, cyanide ions react with metal ions as
 - (a) a reducing agent
- (b) a complexing agent
- (c) an oxidizing agent
- (d) a Lewis acid

Answers

1. b	2. a	3. c	4. a	5. c
6. d	7. c	8. c	9. b	10. d
11. c	12. c	13. c	14. d	15. c
16. c	17. a	18. c	19. b	20. c
21. d	22. a	23. b	24. b	25. c
26. d	27. a	28. c	29. c	30. c
31. d	32. b	33. a	34. d	35. d
36. c	37. a	38. d	39. a	40. c
41. b	42. b	43. c	44. d	45. c
46. a	47. d	48. a, c, d	49. b, c	50. a, b, c
51. b, c	52. a, b, c, d	53. a, b	54. a, b	

Hints to More Difficult Questions

- 4. Value of effective nuclear charge.
- 6. $Cu_2S + 2Cu_2O \longrightarrow 6Cu + SO_2$ $2Cu^+ \rightleftharpoons Cu^{2+} + Cu$

$$K_{\rm eq} = \frac{[{\rm Cu}^{2+}]}{[{\rm Cu}^{+}]} = 1.6 \times 10^{6}$$

The high value of K_{eq} indicates that the reaction will proceed from left to right.

$$\begin{aligned} \textbf{13.} \quad & \text{CuSO}_4 + \text{Na}_2 \text{S}_2 \text{O}_3 \longrightarrow & \text{Na}_2 \text{SO}_4 + \text{CuS}_2 \text{O}_3 \\ & & \text{6CuS}_2 \text{O}_3 + \text{Na}_2 \text{S}_2 \text{O}_3 \longrightarrow & \text{Na}_4 [\text{Cu}_6 (\text{S}_2 \text{O}_3)_5] + \text{Na}_2 \text{S}_4 \text{O}_6 \end{aligned}$$

28.
$$AgBr + 2Na_2S_2O_3 \longrightarrow Na_3[Ag(S_2O_3)_2] + NaBr$$

29. Small cations may polarize a complex ion so much that the cation and the anion are pulled apart.

$$AgCl + 2NH_3 \longrightarrow [Ag(NH_3)_2]^+Cl^-$$

31. $3Au^+ \longrightarrow 2Au + Au^{3+} \quad K = 1 \times 10^{10}$

The high value of *K* indicates that disproportionation occurs.

- **52.** (a) In Cu(CN)₄³⁻, $x 4 = -3 \implies x = +1$.
 - (b) In $CuCl_2^-$, $x 2 = -1 \implies x = +1$.
 - (c) In CuCl₃²⁻, $x 3 = -2 \implies x = +1$.
 - (d) In CuCl₄³⁻, $x 4 = -3 \implies x = +1$.

In all these cases, *x* is the oxidation state of Cu.

14

Zinc and Mercury

• Type 1 •

1. The electronic configurations of Zn and Hg are respectively

Choose the correct option. Only one option is correct.

(b) Hg

(a) $3d^{10} 4s^2$ and $4d^{10} 5s^2$ (b) $3d^{10} 4s^2$ and $4f^{14} 4d^{10} 5s^2$ (c) $3d^{10} 4s^2$ and $4f^{14} 5d^{10} 6s^2$ (d) $4f^{14} 5d^{10} 5s^2$ and $4f^{14} 5d^{10} 7s^2$

+1 in its compounds?

(a) Mg

3.	Zinc a	and mercury are respectively the	
	(a)	last member of the 3d and the first member of the 5d transit series	ion
	(b)	first member of the 4d and the last member of the 5d transit series	ion
	(c)	last member of the 3d and the last member of the 5d transit series	ion
	(d)	last member of the 3d and the last member of the 6d transit series	ion
4.	The m	nercury (I) ion is always	
	` '	[Hg—Hg] (b) [Hg—Hg] ²⁺ [Hg—Hg] ⁴⁺ (d) Hg ²⁺	
5	Zinc i	s extracted from 7nS by the	

(a) calcination of ZnS followed by hydrogen reduction at 400°C
 (b) calcination of ZnS followed by carbon dioxide reduction
 2-134

2. Which of the following metals shows a well-established oxidation state of

(c) Zn

(d) Cd

	(c)	roasting of ZnS followed by muffle furnace	alun	ninium reduction at 1200°C in a
	(d)	roasting of ZnS followed by c in a smelter	arbo	n monoxide reduction at 1200°C
6.	(a)	urest zinc is made by Mond's process the van Arkel method		zone refining poling
7.	Which	n of the following is known as	ʻphil	osopher's wool'?
	(a)	$ZnSO_4 \cdot 7H_2O$	(b)	$ZnS + BaSO_4$
	(c)	ZnO	(d)	ZnCO ₃
8.		n of the following structures do Wurtzite		nO have? Rutile
	(c)	Spinel	(d)	Fluorite
9.	(a) (b) (c)	of the following mixtures is k Anhydrous FeSO ₄ and O ₃ Anhydrous MgCl ₂ and LiAlf- Anhydrous ZnCl ₂ and concer Anhydrous Na ₂ SO ₄ and conce	I ₄ ntrate	ed HCl
10.	(a) (b) (c)	ulated zinc is made by displacing Zn from a ZnSO ₄ s zone refining pouring molten zinc into mol pouring molten zinc into wat	ten r	
11.	Lithop	oone is a white pigment, and is	a m	ixture of
	(a)	BaSO ₄ and ZnCO ₃	(b)	BaSO ₄ and ZnS
	(c)	BaS and ZnO	(d)	ZnCl ₂ and ZnS
12.	(a)	SO_4 solution is boiled with an SO_4 Solution SO_4 Solu	(b)	CO_3 solution to produce $ZnCO_3$ $ZnCO_3 \cdot Zn(OH)_2$
13.	(a)	nized iron pipes are made by ' zinc in molten iron iron in molten zinc	(b)	lipping' iron in molten nickel iron in molten magnesium
14.	ZnSO	4 is boiled with Na ₂ CO ₃ to pro-	duce	
		$ZnCO_3 \cdot Zn(OH)_2$	` ′	ZnCO ₃
	(c)	$ZnCO_3 \cdot ZnSO_4$	(d)	$ZnCO_3 \cdot Na_2SO_4$

(b) rock-salt structure

(d) spinel structure

15. ZnO, a white solid, is a covalent molecule, and adopts a

(a) diamond structure

(c) bcc structure

				±	
16.	(a) (b) (c)	n of the following pairs are ison Blue vitriol and white vitriol White vitriol and Mohr's salt Calgon and microcosmic salt White vitriol and epsom salt	morp	phous?	
17.	gravir (a)	ing the following, which is the numeric determination of zinc? $ZnCO_3$ $Zn_2P_2O_7$	(b)	thermally stable, and used in the ${ m Zn(NO_3)_2}$ ${ m ZnNH_4PO_4}$	
18.	(a)	ormula of basic zinc acetate is (CH ₃ COO) ₂ Zn · ZnCO ₃ (CH ₃ COO) ₂ Zn · Zn ₄ O		$(CH_3COO)_2Zn \cdot ZnO$ $(CH_3COO)_2Zn \cdot Zn(OH)_2$	
19.	white (a) (b) (c)				
20.	of Gro	onic radii of Group 12 metals Z oup 2 metals because Zn, Cd an 10 d electrons which shield th 10 d electrons which shield th	nd H ie nu	clear charge poorly	

(c) 10 d electrons which have a large radius ratio(d) 10 d electrons which have a large exchange energy

21. The structure of zinc blend is similar to that of diamond,

(b) CdCl₂

zinc

(a) ZnCl₂

one-fourth by zinc

22. Which of the following is covalent?

(a) with half the positions occupied by sulphur and one-fourth by

(d) with one-fourth of the positions occupied by sulphur and

(c) SnCl₂

(d) HgCl₂

(b) with half the positions occupied by sulphur and half by zinc(c) with one-fourth of the positions occupied by sulphur and half by

24. Which of the following is arranged in order of decreasing thermal

25. Which of the following oxides can exist in yellow as well as red forms?

(b) HgCl₂(d) NH₄Cl

(b) Cd > Hg > Zn

(d) Hg > Cd > Zn

23. Which of the following oxides does not sublime on heating?

(a) HgO

(c) As_2O_3

(a) Zn > Hg > Cd

(c) Zn > Cd > Hg

stability?

	(a)	ZnO	(b)	PbO	(c)	CdO	(d)	HgO
26.		ZnO on head							The colour is
	(a)	The number transition te				S 7	with tem _]	perature	and is zero at
	(b)	The number absolute zer		defects	increases	S 7	with tem _]	perature	and is zero at
	(c)	The number absolute zer		defects	decrease	s v	with tem	perature	and is zero at
	(d)	The number transition te				s v	with tem	perature	and is zero at
27.		n of the follong point?	wing	g are a	rranged	in	the corre	ect orde	r of increasing
	(a)	$ZnCl_2 < ZnI$	3r ₂ <	ZnI ₂ <	ZnF_2				
	(b)	$ZnF_2 < ZnI_2$	< Zr	$Br_2 < 2$	$ZnCl_2$				
	(c)	$ZnF_2 < ZnC$	$l_2 < 2$	ZnBr ₂ <	$<$ ZnI $_2$				
	(d)	$ZnCl_2 < ZnI$	$F_2 < Z_2$	ZnBr ₂ <	$<$ ZnI_2				
28.	The sa	alts of which	of th	e follo	wing met	als	s are exte	nsively l	nydrated?
	(a)	Pb			(b)	Cd		
	(c)	Hg			(d)	Zn		
29.	The ic		of G	roup 1	2 (Zn, Cd	a	nd Hg) h	ave a co	mplete d shell,
	(a)	behave like	supe	rcond	uctors				
	(b)	are very hig	h me	elting s	solids				
	(c)	do not beha	ve lil	ke tran	sition me	ta	ls		
	(d)	behave like	semi	condu	ctors				
30.	Zinc r	eacts with ve	ry d	ilute ni	itric acid 1	Ю	produce		
		$NH_4NO_2 + Z$	_				$ZnO_2^{2-} +$		
	(c)	$N_2O + N_2 +$	Zn ²⁺		(d)	NH ₄ NO	$l_3 + Zn^{2+}$	

31. Mercury vapours are (a) tetratomic

(c) monatomic

32. The metals of Group 12 are softer than other transition metals because

(a) high enthalpy of sublimation, ionization energy and enthalpy of

(a) Group 12 metals have a cage-like structure
(b) Group 12 metals have high ionization energies
(c) s as well as d electrons take part in metallic bonding
(d) d electrons do not take part in metallic bonding

33. Group 12 metals are relatively noble because of their

(b) hexatomic

(d) diatomic

		hydration		
	(b)	low enthalpy of sublimation hydration	, ior	ization energy and enthalpy of
	(c)	high enthalpy of sublimation enthalpy of hydration	on a	nd ionization energy, and low
	(d)	high enthalpy of sublimatic enthalpy of hydration	n, lo	ow ionization energy and high
34.	Which	Č 1	is us	sed as an electrode of the second
	(a)	$Hg_2(NO_3)_2$	(b)	HgCl ₂
	(c)	HgS	(d)	Hg ₂ Cl ₂
35.	Which	n of the following is arranged i	n or	der of increasing melting point?
	(a)	$HgF_2 < HgCl_2 < HgI_2 < HgBr_2$	2	
	(b)	$HgF_2 < HgCl_2 < HgBr_2 < HgI_2$	2	
	(c)	$HgI_2 < HgBr_2 < HgCl_2 < HgF_2$	2	
	(d)	$HgBr_2 < HgI_2 < HgCl_2 < HgF_2$	2	
36.	All m	ercurous compounds are		
	(a)	diamagnetic in the solid state	as w	vell as in solution
	(b)	paramagnetic in the solid stat	te as	well as in solution
	(c)	diamagnetic in the solid state	and	paramagnetic in solution
	(d)	paramagnetic in the solid state	te an	d diamagnetic in solution
37.	Which water		mer	cury is the least hydrolysed by
	(a)	HgF ₂	(b)	HgCl ₂
	(c)	$HgBr_2$	(d)	HgI_2
38.		$ m l_2$ (calomel) and $ m HgCl_2$ (corrost ammonia to produce respecti		sublimate) react separately with

39. Mercury is the only metal which is a liquid at room temperature. This is

van der Waals forces among the mercury molecules

van der Waals forces among the mercury molecules

(c) weak metallic bonding because of its d ¹⁰s ² configuration, and low

(d) strong metallic bonding due to its d ¹⁰s ² configuration and strong

(a) Hg(NH₂)Cl + Hg and Hg(NH₂)Cl(b) Hg(NH₂)Cl and Hg(NH₂)Cl + Hg

(a) high viscosity of mercury(b) large surface tension of mercury

due to the

41.

42.

43.

44.

45.

(c) $Hg(NH_2)Cl + HgCl_2$ and $Hg(NH_2)Cl + Hg_2Cl_2$

(d) $Hg(NH_2)Cl + Hg$ and $Hg(NH_2)Cl + Hg$

40. When an excess of SnCl₂ is added to an HgCl₂ solution,

(b) the composition $Hg_2ONH_2 \cdot HI$

(a)	Hg_2Cl_2	(b)	Hg					
(c)	Sn (d) $[SnCl_6]^{4-}$							
is pro	duced.							
Corro	sive sublimate is made by hea	ting						
(a)	Hg ₂ SO ₄ and NaCl	(b)	Hg ₂ Cl ₂ and I	Hg ₂ O)			
(c)	HgSO ₄ and NaCl	(d)	HgSO ₄ and I	HgS				
Calon	nel is made by treating							
(a)	Hg ₂ (NO ₃) ₂ with HCl							
(b)	Hg(NO ₃) ₂ with NaCl							
(c)	Hg ₂ (NO ₃) ₂ and Hg with aqua	regi	a					
	HgSO ₄ with NaCl							
Nessl	er's reagent is a mixture of							
	HgCl ₂ and KI (excess)							
(b)	HgCl ₂ , KI (excess) and liquor	amn	nonia					
	HgCl ₂ , KI (excess) and KOH							
	K ₂ [HgI ₄] and KI							
	ng the following, Cd (b) Zn	(a)	Na	(d)	Fo			
` '	· /	(C)	INa	(u)	1.6			
does not form amalgams.								
Millon's base has								
(a) a three-dimensional framework of Hg ₂ N with OH ⁻ and water molecules occupying spacious cavities and channels								

- (c) a three-dimensional framework of HgN₂ with OH⁻ and NH₃ molecules occupying spacious cavities and channels
- (d) a perovskite structure
- **46.** SO₂ gas is passed through an aqueous suspension of corrosive sublimate (HgCl₂), in order to produce
 - (a) HgSO₄ and HCl
- (b) Hg₂Cl₂

(c) Hg_2SO_4

- (d) HgSO₄
- 47. HgCl₂ is a solid containing
 - (a) an angular Cl-Hg-Cl molecule
 - (b) a linear Cl—Hg—Cl molecule
 - (c) a T-shaped Cl—Hg—Cl molecule
 - (d) a pyramidal Cl—Hg—Cl molecule
- 48. $ZnSO_4$ reacts with the excess of a KCN solution to produce the complex ion
 - (a) $[Zn(CN)_6]^{4-}$ with an octahedral structure
 - (b) $[Zn(CN)_4]^{2-}$ with a square-planar structure
 - (c) [Zn(CN)₂] with a linear structure
 - (d) $[Zn(CN)_4]^{2-}$ with a tetrahedral structure
- **49.** Which of the following statements is correct?
 - (a) In $[Zn(NCS)_4]^{2+}$, the ligand is bonded through S, but in $[Cd(SCN)_4]^{2+}$ the ligand is bonded through N.
 - (b) In $[Zn(NCS)_4]^{2+}$, the ligand is bonded through N, but in $[Cd(SCN)_4]^{2+}$ the ligand is bonded through S.
 - (c) In $[Zn(NCS)_4]^{2+}$ as well as $[Cd(SCN)_4]^{2+}$, the ligands are bonded through S.
 - (d) In $[Zn(NCS)_4]^{2+}$ as well as $[Cd(SCN)_4]^{2+}$, the ligands are bonded through N.
- **50.** Which of the following has been arranged in order of increasing bond strength?
 - (a) $Zn_2^{2+} < Hg^{2+} < Cd^{2+}$
- (b) $Cd_2^{2+} < Hg_2^{2+} < Zn_2^{2+}$
- (c) $Zn_2^{2+} < Cd_2^{2+} < Hg_2^{2+}$
- (d) $Hg_2^{2+} \ll Cd_2^{2+} \ll Zn_2^{2+}$

• Type 2 •

Choose the correct options. More than one option is correct.

- **51.** Which of the following statements are correct?
 - (a) ZnSO₄ dissolves in liquor ammonia to form a tetrahedral complex.
 - (b) Mercurous and cuprous ions are represented respectively as Hg₂²⁺ and Cu_2^{2+} .
 - (c) ZnS is precipitated from a ZnSO₄ solution in an alkaline medium by H₂S.
 - (d) Granulated zinc easily reacts with dilute sulphuric acid, producing hydrogen gas.
- **52.** HgCl₂ is prepared by the reaction between
 - (a) HgS and aqua regia
- (b) Hg and heated chlorine
- (c) Hg₂Cl₂ and heated mercury (d) Hg₂Cl₂ and dilute HCl
- 53. Which of the following statements are correct?
 - (a) Mercury (II) oxide exists in two different colours due to particle size.
 - (b) Mercury (II) oxide is thermally unstable.
 - (c) Mercurous ion is included in Group I among the analytical groups.
 - (d) The critical angle of mercury is 110°.
- 54. Which of the following are correctly matched?
 - (a) $HgS \longrightarrow Vermilion$
- (b) $Hg_2Cl_2 \longrightarrow Laxative$
- (c) $HgCl_2 \longrightarrow Chlorophyll$ (d) $K_2HgI_4 \longrightarrow Nessler's$ reagent
- **55.** Which of the following statements are correct?
 - (a) The first ionization energy of zinc is greater than that of mercury.
 - (b) The first ionization energy of zinc is less than that of mercury.
 - (c) Zn^{2+} is more reducing than Hg^{2+} .
 - (d) Zn is more reducing than Hg.
- 56. Zinc is the only metal in Group 12 which shows amphoteric properties, and is soluble in alkalis forming
 - (a) $Na_2[Zn(OH)_4]$

- (b) $Na[Zn(OH)_3 \cdot (H_2O)_3]$
- (c) $Na[Zn(OH)_3 \cdot H_2O]$
- (d) $Na_4[Zn(OH)_6]$
- 57. Zinc can form four-coordinated complexes with various ligands, represented by the formulae
 - (a) $[Zn(CN)_4]^{4-}$

(b) $[Zn(CN)_4]^{2-}$

(c) $[Zn(NCS)_4]^{2+}$

(d) $[Zn(NH_3)_4]^{2+}$

- 58. Zinc is used in
 - (a) galvanization

(b) sherardizing

(c) dry batteries

- (d) the preparation of alloys
- **59.** Which of the following statements are correct for ZnO?
 - (a) It is an intrinsic semiconductor.
 - (b) It is an intrinsic superconductor.
 - (c) It shows grain boundary defects.
 - (d) When ZnO is heated with cobalt nitrate on a charcoal block in an oxidizing flame, it forms cobalt zincate, also known as Rinman's green.
- **60.** For the equilibrium

$$Hg_2^{2+}(aq) \rightleftharpoons Hg(l) + Hg^{2+}(aq)$$

$$[H\sigma^{2+}]$$

$$K_{\rm eq} = \frac{[{\rm Hg}^{2+}]}{[{\rm Hg}_2^{2+}]} = 6 \times 10^3 \text{ at } 25^{\circ}{\rm C}$$

choose the correct options.

- (a) The Hg (I) ion disproportionates to the Hg (II) ion and Hg.
- (b) The addition of a Cl⁻ ion shifts the equilibrium to the right.
- (c) The equilibrium is displaced to the right by the removal of the ${\rm Hg}^{2+}$ ion.
- (d) The equilibrium is shifted to the left by the removal of the Hg^{2+} ion.

1. c	2. b	3. c	4. b	5. d
6. b	7. c	8. a	9. c	10. d
11. b	12. b	13. c	14. a	15. a
16. d	17. c	18. c	19. d	20. a
21. b	22. d	23. a	24. c	25. d
26. b	27. a	28. d	29. c	30. d
31. c	32. d	33. c	34. d	35. d
36. a	37. d	38. a	39. c	40. b
41. c	42. a	43. c	44. d	45. a
46. b	47. b	48. d	49. b	50. c
51. a, c, d	52. a, b, c	53. a, b, c	54. a, b	55. b, d
56. a, b, c	57. b, c, d	58. a, b, c, d	59. a, d	60. a, b, c

Hints to More Difficult Problems

- 2. The oxidation state of mercury in Hg_2^{2+} (mercurous ion) is +1.
- 11. It is made by the reaction between BaS and $ZnSO_4$. BaS + $ZnSO_4 \longrightarrow BaSO_4 + ZnS$
- 16. ZnSO₄ · 7H₂O and MgSO₄ · 7H₂O
- 17. $ZnSO_4 + Na_2HPO_4 + NH_3 \longrightarrow Zn(NH_4)PO_4 + Na_2SO_4$ $2Zn(NH_4)PO_4 \stackrel{\Delta}{\longrightarrow} Zn_2P_2O_7(s) + 2NH_3 \uparrow + H_2O \uparrow$
- 22. HgCl₂ is more polarized than the other compounds.
- **24.** The size of the covalent radius increases from Zn to Hg, and the melting point decreases.
- 28. The smaller the ion, the greater is the tendency to form a hydrated salt.
- 34. Hg | Hg₂Cl₂, Cl⁻, the cell reaction is Hg₂Cl₂I(s) + 2e \longrightarrow 2Hg(l) + 2Cl⁻
- 37. HgI_2 is the most covalent, and has the least tendency to hydrolyse.
- 40. $2\text{HgCl}_2 + \text{SnCl}_2 \longrightarrow \text{Hg}_2\text{Cl}_2 + \text{SnCl}_4$ $\text{Hg}_2\text{Cl}_2 + \text{SnCl}_2 \longrightarrow 2\text{Hg} + \text{SnCl}_4$
- 48. $ZnSO_4 + 4KCN \longrightarrow K_2[Zn(CN)_4] + K_2SO_4$ $K_2[Zn(CN)_4] \longrightarrow 2K^+ + Zn(CN)_4^{2^-}$ (tetrahedral)
- 51. (a) $ZnSO_4 + 4NH_3 \longrightarrow [Zn(NH_3)_4]SO_4$
 - (c) $ZnSO_4 + H_2S \xrightarrow{NH_4Cl + NH_3} ZnS \downarrow$
 - (d) $Zn + H_2SO_4 \longrightarrow ZnSO_4 + H_2$
- **60.** (a) $\mathrm{Hg_2^{2+}}$ easily disproportionates to Hg and Hg $^{2+}$ as indicated by the K_{eq} value. For options (b) and (c), consider Le Chatelier's principle.

Iron

• Type 1 •

Choose the correct option. Only one option is correct.

1.	The	chemical	processes	in	the	production	of	iron	from	haematite	ore
	invo	lve									

- (a) reduction
- (b) oxidation
- (c) reduction followed by oxidation
- (d) oxidation followed by reduction

2. In the manufacture of iron, the principal reaction in the zone of heat absorption of the blast furnace is

$$(a) \quad 2C(s) + O_2(g) \longrightarrow \ 2CO_2(g)$$

$$\text{(b)} \quad CO_2(g) + C(s) \longrightarrow 2CO(g)$$

(c)
$$2C(s) + O_2 \longrightarrow 2CO(g)$$

(d)
$$3C(s) + O_2(g) \longrightarrow C_3O_2(g)$$

3. In the extraction of iron, Fe₂O₃ is reduced by

(a) carbon

- (b) carbon dioxide
- (c) carbon monoxide
- (d) calcium carbonate

4. Among the following, the maximum amount of carbon is present in

(a) pig iron

(b) wrought iron

(c) steel

(d) stainless steel

5. When hard steel is heated to bright redness and then allowed to cool slowly, it gets softened. The process is called

(a) annealing

(b) hardening

(c) quenching

(d) tempering

6.		mild steel is heated to a high, it becomes hard and brittle. T		perature and suddenly cooled in rocess is called
	(a)	hardening	(b)	annealing
	(c)	quenching	(d)	tempering
7.		ng the following salts of iron ous solution?	, wh	ich is the most unstable in an
	(a)	$FeCl_3 \cdot 6H_2O$	(b)	$Fe_2(SO_4)_3 \cdot 9H_2O$
	(c)	$K_4[Fe(CN)_6]$	(d)	FeI ₃
8.	An Fe	Cl ₃ solution reacts with sodiur	n hy	droxide to produce
		$Fe_2O_3 \cdot nH_2O$	-	Fe_3O_4
		FeO and FeCl ₃		Fe ₂ O ₃ and FeO
9		manufacture of iron from hae		
9.		slag		flux
	` '	matrix	` '	reducing agent
10	` '	n of the following compounds		
10.		FeBr ₂ (b) FeI ₃		FeCl ₃ (d) Fe ₂ O ₃
11	Which	n of the following statements is	cor	rect?
	(a)	_		e drawn off through the same
	(b)	Molten slag and molten iro openings.	n aı	re drawn off through separate
	(c)	Slag floats on molten iron, the	us pr	otecting iron from reduction.
	(d)	Pig iron is soft and brittle.		
12.	The ir	on group contains		
	(a)	six metals	(b)	three metals
	(c)	nine metals	(d)	eight metals
13.	Iron r		ence	of O ₂ at a high temperature to
	(a)	$Fe_3(PO_4)_2$	(b)	FePO ₄
	(c)	Fe ₃ P	(d)	$Fe_3(PO_3)_2$
14.	Iron is	s not attacked by		
	(a)	concentrated NaOH	(b)	dilute NaOH
	(c)	dilute H ₂ SO ₄	(d)	steam
15.	In Fe ₂	(CO) ₉ ,		
	(a)	two bridging CO groups are j	oine	d with two iron atoms
	(b)	five bridging CO groups are j	oine	d with two iron atoms

		(c) three bridging CO groups are joined with two iron atoms(d) one bridging CO group is joined with two iron atoms				
16.	The st	ructure of Fe(CO) ₅ is				
	(a)	pentagonal bipyramidal	(b)	irregular tetrahedral		
	(c)	irregular octahedral	(d)	trigonal bipyramidal		
17.	Which	n of the following is nonstoichi	ome	tric and metal deficient?		
	(a)	FeO	(b)	Fe_3O_4		
	(c)	Fe_2O_3	(d)	All of these		
18.	Which with I	n of the following has a cubic cl Fe ²⁺ ions occupying all octahed	lose- Iral l	packed arrangement of O ²⁻ ions noles?		
		Fe(OH) ₃		FeO		
	(c)	Fe(OH) ₂	(d)	Fe_2O_3		
19.		H) ₂ dissolves in a concentrated lex with the formula	Na(OH solution giving a blue-green		
	(a)	$Na_4[Fe_2(OH)_6]$	(b)	$Na_2[Fe(OH)_6]$		
	(c)	$Na_4[Fe(OH)_6]$	(d)	$Na_4[Fe(OH)_4]$		
20.	Ferro	cene is a compound of				
	(a)	one Fe $^{2+}$ and two $C_5H_5^-$ ions				
	(b)	two Fe $^{2+}$ and two $C_5H_6^-$ ions				
	(c)	one Fe $^{3+}$ and two $C_6H_5^-$ ions				
	(d)	two Fe $^{2+}$ and two $C_5H_5^-$ ions				
21.	Norm	ally FeCl ₃ · 6H ₂ O consists of				
	(a)	trans- $[Fe(H_2O)_2Cl_2] \cdot Cl \cdot 4H_2Cl_2$)			
	(b)	trans-[Fe(H_2O) ₆ Cl]Cl ₂				
	(c)	$\textit{trans-}[Fe(H_2O)_4Cl_2]Cl \cdot 2H_2O$				
	(d)	$\textit{trans-}[Fe(H_2O)_3Cl_2]Cl \cdot 3H_2O$				
22.	In con	ncentrated HCl, FeCl ₃ · 6H ₂ O fo	rms	an		
		$[FeCl_6^{2-}]$ ion		[FeCl ₄] - ion		
	(c)	[FeCl ₄] + ion	(d)	$[Fe(H_2O)_6]^{3+}$ ion		
23.	The ra	nte of reaction of iron filings wi	th o	xygen may be increased by		
	(a)	increasing the concentration of	of ox	ygen		
	(b)	decreasing the amount of iror	ı			
	(c)	decreasing the pressure of oxy	_			
	(d)	lowering the temperature of i	ron			

Iron 2-147

(d) FeO₄²⁻

(c) FeO

(b) ferrate ion Fe VIO₄²⁻
 (d) ferrate ion Fe IIIO₄⁵⁻

30. If Cl2 is passed into an alkaline solution of hydrated ferric oxide, a

31. The electronic configuration of Fe ³⁺ and Mn ²⁺ are respectively

24.	Which		ısed	to form a neutral ferric chloride
	(a)	Adding an excess of NH ₃ to a	n Fe	Cl ₃ solution
		Adding an excess of NaOH to		-
	(c)	Adding one to two drops of N	NH_3	to an FeCl ₃ solution
	(d)	Adding dilute HCl to an FeC of an NaOH solution	Cl ₃ sc	lution followed by the addition
25.		presence of Fe ³⁺ ions is detect a blood-red colour due to the f		by the addition of SCN^- which ation of
	(a)	[Fe(SCN) ₃ OH] ²⁻ and some [F	e(SC	$N)_3]^{2-}$
	(b)	$[Fe(SCN)_2(H_2O)_3]^{2+}$ and some	e [Fe	(SCN) ₄] -
	(c)	$[Fe(SCN)_3(H_2O)_5]^{2+}$ and some	Fe(S	6CN) ₃
	(d)	[Fe(SCN)(H ₂ O) ₅] ²⁺ and some	Fe(S	CN) ₃ and [Fe(SCN) ₄] ⁻
26.		ddition of F ⁻ to the blood-red s rless due to the formation of	oluti	on of $[Fe(SCN)(H_2O)_5]^{2+}$ turns it
	(a)	$[Fe(H_2O)_5F_3]$	(b)	$[Fe(H_2O)_5F]^{2+}$
	(c)	[FeF ₆] ³⁻	(d)	$[FeF_4]^{-1}$
27.	On fu	sion with Na ₂ CO ₃ , Fe ₂ O ₃ gives		
		Na ₂ FeO ₄		Fe_3O_4
	(c)	NaFeO ₂	(d)	Na_3FeO_3
28.	A mix	xed oxide of iron, Fe ₃ O ₄ , is repr	esen	ted by
	(a)	$Fe_2^{II}Fe^{III}O_4$		
	(b)	Fe ^{II} Fe ₂ ^{III} O ₄ or Fe ^{III} (Fe ^{II} Fe ^{III})C) ₄	

 $\begin{array}{ll} \text{(c)} & Fe_2^{III}Fe^{II}O_4 \\ \text{(d)} & Fe^{III}Fe_2^{II}O_4 \end{array}$

(a) Fe_3O_4

(a) ferric ion Fe³⁺

(c) ferrite ion Fe VIO₄²⁻

29. On ignition to 1400°C, Fe₂O₃ gives black

(b) Fe

red-purple solution is formed containing the

(a) [Ar] 3d ⁶ 4s ² and [Ar] 4d ⁶ 4s ² (b) [Ar] 3d ⁵ 4s ⁰ and [Ar] 3d ⁵ 4s ¹

- (c) $[Ar] 3d^5 4s^0$ and $[Ar] 4d^5 6s^0$ (d) $[Ar] 3d^5 4s^0$ and $[Ar] 3d^5 4s^0$ 32. Which of the following statements is correct for [Fe(H₂O)₅NO]²⁺, which is formed during the brown-ring test for the nitrate radical? (a) The colour of the brown ring is not due to charge transfer spectra. (b) In the complex mentioned, the oxidation number of iron is +2. (c) The magnetic moment of iron in the complex mentioned is 3.9 Bohr magneton. (d) The magnetic moment of iron in the complex mentioned is 2.83 Bohr magneton. 33. Among the following compounds, which is thermally the most stable? (b) Fe_2O_3 (c) Fe_3O_4 (d) FeCO₃ 34. In the ferrocene $[Fe(C_5H_5)_2]$ molecule the bonding is regarded as π -bonding involving the lateral overlap of (a) the d_{yz} and d_{yz} orbitals on Fe with a delocalized aromatic orbital from each cyclopentadienyl ring (b) p_x and p_y orbitals on Fe with a delocalized aromatic orbital from each cyclopentadienyl ring (c) d_{yz} and d_{yz} orbitals on Fe with a localized aromatic orbital from each cyclopentadienyl ring
- **35.** The alloy nichrome contains
 - (a) Ni, Cr, Fe and Mn

cyclopentadienyl ring

- (b) Cr, Ni, Cu and Zn
- (c) Ni, Cr, Fe and Zn
- (d) Ni, Cr, Fe and C
- **36.** The concentration of chromite (FeCrO₄) is carried out by
 - (a) gravity separation
- (b) froth floation
- (c) magnetic separation
- (d) roasting
- 37. The densities of Pb, Fe and Al are in the order
 - (a) Pb > Fe > Al

(b) Fe > Al > Pb

(c) Pb > Al > Fe

- (d) Al > Fe > Pb
- 38. Concentrated nitric acid passivates iron due to the
 - (a) formation of Fe(NO₃)₂, which does not react with iron, on the surface of the iron

(d) d_{z^2} orbitals on Fe with a delocalized aromatic orbital from each

- (b) formation of a protective coating of the oxide of iron which does not react with nitric acid, on the surface of the iron
- (c) formation of a stable complex on the surface of the iron
- (d) decomposition of nitric acid into NO₂ and O₂

(d) $FeSO_4(NH_4)_2SO_4 \cdot 6H_2O$

(b) FeO and SO₃

(d) Fe₂O₃ and SO₂

(b) Fe₃O₄

41.		us and ferric salts can be best the reagent	dist	tinguished	from eacl	n other	by
	(a)	Fe(SCN) ₃	(b)	NH ₄ CNO			
	(c)	$K_4[Fe(CN)_6]$	(d)	dilute HN	O_3		
42.	Acidi	fied KMnO ₄ is decolourized by					
	(a)	the ferric ammonium alum					
	(b)	Mohr's salt					
	(c)	haematite					
	(d)	a neutral ferric chloride soluti	on				
43.	On str SO ₃ ?	rong heating, which of the following	lowi	ng produce	s a mixtuı	re SO ₂ a	nd
	(a)	$Fe_2(SO_4)_3 \cdot 9H_2O$					
	(b)	$(NH_4)_2SO_4 \cdot Fe_2(SO_4)_3 \cdot 24H_2O_4$)				
	(c)	FeSO ₄ · 7H ₂ O					
		FeS ₂					
44.		reacts with K_3 [Fe(CN) ₆] to bull's blue. This precipitate is on			recipitate	known	as
	(a)	K Fe II [Fe II (CN) $_6$]	(b)	K Fe III [Fe I	$^{II}(CN)_6$]		
	(c)	K Fe III [Fe II (CN) ₆]	(d)	K Fe II [Fe II	$[(CN)_6]$		
45.		reacts with $K_4[Fe(CN)_6]$ to ian blue. Prussian blue is	give	a blue pi	recipitate	known	as
	(a)	$Fe_4^{III}[Fe^{III}(CN)_6]$	(b)	Fe ₄ ^{II} [Fe ^{II} (C	$(N)_6$		
	(c)	K Fe III [Fe II (CN) ₆]	(d)	K Fe II[Fe IV	$(CN)_6$		
46.	When	heated with concentrated H ₂ S	O ₄ , I	$K_4[Fe(CN)_6]$	gives		
	(a)	$(CN)_2$ and CO_2	(b)	$(CN)_2$ and	CO		
	(c)	CO	(d)	CO_2			

40. Which of the following is used to estimate iron volumetrically using a

39. On strong heating, Fe₂(SO₄)₃ gives
(a) Fe₂O₃ and SO₂

KMnO₄ or a K₂Cr₂O₇ solution?

(c) Fe_2O_3 and SO_3

(a) FeCl₃ · 6H₂O

(c) $Fe_2(SO_4)_3 \cdot 9H_2O$

- 47. Rusting of iron is
 - (a) an electrochemical process
- (b) an electroanalytical process
- (c) a photochemical process
- (d) all of these
- **48.** Rusting of iron is prevented by
 - (a) galvanizing
 - (b) electroplating the iron with a thin layer of tin
 - (c) converting the outer layer of the iron into iron phosphate
 - (d) all of these
- 49. In which of the following reactions is potassium ferrocyanide produced?
 - (a) The reaction between K₃[Fe(CN)₆] and an excess of FeSO₄
 - (b) The reaction between FeSO₄ and an excess of KCN
 - (c) The reaction between Fe₂(SO₄)₃ and an excess of KCN
 - (d) The reaction between Fe₂O₃ and an excess of KCN
- 50. In an aqueous solution, it is easier to oxidize
 - (a) $[Fe^{II}(CN)_6]^{4-}$ ions than $[Fe^{III}(CN)_6]^{3-}$ ions
 - (b) $[Fe^{II}(CN)_6]^{4-}$ ions than $Fe^{IV}O_4^{4-}$ ions
 - (c) $[Fe^{II}(H_2O)_6]^{2+}$ ions than $[Fe^{III}(H_2O)_6]^{3+}$ ions
 - (d) $[Fe^{II}(CN)_6]^{4-}$ ions than $[Fe^{III}(H_2O)_6]^{3+}$ ions

• Type 2 •

Choose the correct options. More than one option is correct.

- **51.** Which of the following statements are correct?
 - (a) The process of producing a hard coating of iron nitride on the surface of steel is called nitriding.
 - (b) The process of producing a thin coating of hardened steel on the surface of mild steel is called hardening.
 - (c) Quenched steel is produced by heating steel to redness and allowing it to cool slowly.
 - (d) Stainless steel is produced by heating wrought iron in molten chromium.
- **52.** Which of the following statements are correct?
 - (a) Cast iron cannot be permanently magnetized.
 - (b) Steel cannot be permanently magnetized.
 - (c) Steel can be permanently magnetized.
 - (d) Spiegeleisen is an alloy of iron, zinc and antimony.

Iron 2-151

- **53.** By which of the following reactions can Fe ^{III} be detected?
 - (a) $FeCl_3 + 3NH_4SCN \longrightarrow Fe(SCN)_3 + 3NH_4Cl$
 - (b) $\operatorname{FeCl}_3 + \operatorname{K}_3[\operatorname{Fe}(\operatorname{CN})_6] \longrightarrow \operatorname{KFe}[\operatorname{Fe}(\operatorname{CN})_6] + 2\operatorname{KCl} + \frac{1}{2}\operatorname{Cl}_2$
 - (c) $FeCl_3 + K_4[Fe(CN)_6] \longrightarrow KFe[Fe(CN)_6] + 3KCl$
 - (d) $FeCl_3 + 3KSCN \longrightarrow Fe(SCN)_3 + 3KCl$
- 54. In which of the following does iron have an oxidation state of zero?
 - (a) $[Fe(H_2O)_5NO]SO_4$
- (b) $Fe(CO)_5$

(c) Fe(CNS)₃

- (d) $Fe_2(CO)_9$
- **55.** Which of the following reactions are used to prepare $K_4[Fe(CN)_6]$?
 - (a) $FeSO_4 + 6KCN \longrightarrow K_4[Fe(CN)_6] + 2K_2SO_4$
 - (b) $2K_3[Fe(CN)_6] + H_2 + 2KOH \longrightarrow 2K_4[Fe(CN)_6] + 2H_2O$
 - (c) $2K_3[Fe(CN)_6] + K_2SO_4 + 2H_2O \longrightarrow 2K_4[Fe(CN)_6] + H_2SO_4 + H_2O_2$
 - (d) $2K_3[Fe(CN)_6] + 2KOH + H_2O_2 \longrightarrow 2K_4[Fe(CN)_6] + 2H_2O + O_2$
- **56.** Which of the following statements are correct?
 - (a) Potassium hexacyanoferrate (III) is orange while potassium hexacyanoferrate (II) is yellow.
 - (b) Fe (II) and Fe (III) form complexes with the CN⁻ ion but not with NH₃.
 - (c) Ferric salts are more stable than ferrous salts.
 - (d) The ferric ammonium alum is a complex salt.
- 57. Which of the following statements are correct?
 - (a) All Fe ²⁺ salts are pale green.
 - (b) In an aqueous solution, the Fe $^{2+}$ ion can readily be oxidized to the Fe $^{3+}$ ion even by atmospheric O₂.
 - (c) Fe ²⁺ salts decolourize an acidified KMnO₄ solution.
 - (d) Fe $^{2+}$ salts turn acidified $K_2Cr_2O_7$ green.
- 58. Which of the following statements are correct?
 - (a) Ferric salts do not react with NO.
 - (b) The formula of sodium nitroprusside is $Na_2[Fe(CN)_5NO]$.
 - (c) Turnbull's blue is paler than Prussian blue.
 - (d) The structure of the $[Fe(CN)_6]^{4-}$ ion is due to d^2sp^3 hybridization.
- **59.** Which of the following statements are correct?
 - (a) $[Fe(CN)_6]^{4-}$ is diamagnetic but $[Fe(CN)_6]^{3-}$ is paramagnetic.
 - (b) Fe³⁺ ions always form tetrahedral complexes.
 - (c) In a compound with an octahedral structure, the d_{xy} and d_{yz} orbitals of a metal ion should be vacant.
 - (d) $[Fe(H_2O)_6]^{3+}$ is more paramagnetic than $[Fe(CN)_6]^{3-}$.

- **60.** Which of the following statements are correct?
 - (a) The element present in invar are Fe, Co and Ni.
 - (b) $[Fe(CN)_6]^{4-}$ is a low-spin complex while $[Fe(H_2O)_6]^{3+}$ is a high-spin complex.
 - (c) Fe³⁺ usually forms octahedral complexes that have low spin.
 - (d) In the octahedral geometry of a metallic compound, the $d_{x^2-y^2}$ and d_z^2 orbitals of the metal should be vacant.

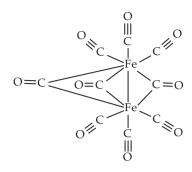
<u>Answers</u>					
1. c	2. b	3. c	4. a	5. a	
6. c	7. d	8. a	9. b	10. c	
11. b	12. c	13. a	14. b	15. c	
16. d	17. d	18. b	19. c	20. a	
21. c	22. b	23. a	24. c	25. d	
26. c	27. c	28. b	29. a	30. b	
31. d	32. c	33. b	34. c	35. d	
36. c	37. a	38. b	39. c	40. d	
41. c	42. b	43. c	44. d	45. c	
46. c	47. a	48. d	49. b	50. a	
51. a, b	52. a, c	53. a, c, d	54. b, d	55. a, d	
56. a, b, c	57. a, b, c, d	58. a, b, c, d	59. a, d	60. a, b, c, d	

Hints to More Difficult Problems

1. In the following formulae, the superscripts denotes oxidation number

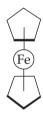
- 2. The reaction is endothermic.
- 3. See the answer to Q. 1.
- 7. Fe³⁺ oxidizes I⁻ to I₂.

15.



17. The fact that they are nonstoichiometric is related to their structures, which are similar. The ccp structures differ only in the arrangement of Fe²⁺ and Fe³⁺ in the holes in the octahedral and tetrahedral crystals.

20.



28. Fe₃O₄ is an inverse (2:3) spinel represented by B(AB)O₄. All spinel structures have a ccp array of anions.

31.
$$Fe(Z = 26) = [Ar]3d^{6}4s^{2}$$

 $Fe^{3+} = [Ar]3d^{5}$
 $Mn(Z = 25) = [Ar]3d^{5}4s^{2}$
 $Mn^{2+} = [Ar]3d^{5}$

37. Density increases with molar mass.

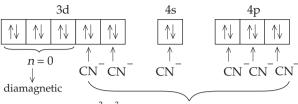
40.
$$MnO_4^- + 8H^+ + 5Fe^{2+} \longrightarrow Mn^{2+} + 5Fe^{3+} + 4H_2O$$

$$Cr_2O_7^{2-} + 14H^+ + 6Fe^{2+} \longrightarrow 2Cr^{3+} + 6Fe^{3+} + 7H_2O$$
The Fe²⁺ ion is produced from FeSO₄ · (NH₄)₂SO₄ · 6H₂O.

43. $FeSO_4 \cdot 7H_2O \xrightarrow{\Delta} FeSO_4 + 7H_2O$ $2FeSO_4 \xrightarrow{\Delta} Fe_2O_3 + SO_2 \uparrow + SO_3 \uparrow$

54. (a) In Fe(CO)₅,
$$x + 5 \times 0 = 0 \Rightarrow x = 0$$
.
(b) In Fe₂(CO)₉, $2x + 9 \times 0 = 0 \Rightarrow x = 0$.
Here x is the oxidation state of iron.

59. (a) In $[Fe(CN)_6]^{4-}$ the electronic structure is as follows.

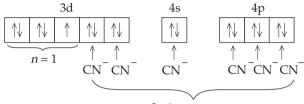


 $\mbox{d}^2 \, \mbox{sp}^3 \, \mbox{hybridization}$ octahedral geometry

$$\mu_{\rm spin} = \sqrt{n(n+2)} \, \, {\rm BM}$$

$$\mu_{spin}=0$$

The electronic structure of $Fe(CN)_6^{3-}$ is as follows.



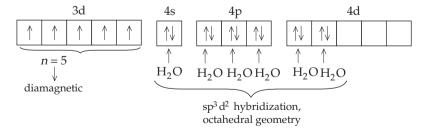
d² sp³ hybridization, octahedral geometry

$$\mu_{spin} = \sqrt{n(n+2)} = \sqrt{3} = 1.73 \text{ BM}$$

Therefore, $[Fe(CN)_6]^{4-}$ is diamagnetic and $[Fe(CN)_6]^{3-}$ paramagnetic.

(d) In $[Fe(CN)_6]^{3-}$, CN^- is a strong ligand. It forces electrons to pair up. But in $[Fe(H_2O)_6]^{3+}$, H_2O is a weak ligand and cannot force electrons to pair up. So n = 5, and hence it is more paramagnetic than $[Fe(CN)_6]^{3-}$.

The electronic structure of $[Fe(H_2O)_6]^{3+}$ is as follows.



Here
$$n = 5$$
.
 $\mu_{\text{spin}} = \sqrt{n(n+2)} = \sqrt{5(5+2)} = 5.92 \text{ BM}$

16

Coordination Chemistry

• Type 1 •

Choose the correct option. Only one option is correct.

- 1. Which of the given statements is not true for the following reaction? $[Cu(H_2O)_4]^{2+} + 4NH_3 \rightleftharpoons [Cu(NH_3)_4]^{2+} + 4H_2O$
 - (a) It is a ligand-substitution reaction.
 - (b) NH_3 is a relatively strong-field ligand while H_2O is a weak-field ligand.
 - (c) During the reaction, there is a change in colour from light blue to dark blue.
 - (d) $\left[Cu(NH_3)_4\right]^{2+}$ has a tetrahedral structure, and is paramagnetic.
- 2. Which of the following is a hexadentate ligand?
 - (a) 2,2-Dipyridyl
 - (b) Ethylenediamine
 - (c) Ethylenediaminetetracetate ion
 - (d) Biphenyl
- 3. Dimethyl glyoxime is an example of a
 - (a) monodentate ligand
- (b) bidentate ligand
- (c) tridentate ligand
- (d) hexadentate ligand

- 4. Ferrocene is
 - (a) bis (cyclopentadienyl) iron (II)
 - (b) dicyclopentadienyl iron (0)
 - (c) dicylopentadienyl iron (III)
 - (d) bis (cyclopentyl) iron (II)
- 5. The IUPAC name of the complex $[Ni(C_4H_7O_2N_2)_2]$ formed from the reaction of Ni $^{2+}$ with dimethyl glyoxime is

	(a)	bis (methyl gloxal) nickel (II)				
	(b) bis (dimethyl oxime) nickelate (IV)					
	(c)	bis (2,3-butanediol dioximato) nic	kel (II)		
	(d)	bis (2,3-butanedione dioximate	to) n	ickel (II)		
6.		n of the following complex ion per (EAN) rule?	ns ob	peys Sidwick's Effective Atomic		
	(a)	[Fe(CN) ₆] ³⁻	(b)	$\left[Cu(CN)_4\right]^{3-}$		
	(c)	$[Cr(NH_3)_6]^{3+}$	(d)	$[Ni(en)_3]^{2+}$		
7.	The ef	ffective atomic number for the	com	plex ion $[Pd(NH_3)_6]^{4+}$ is		
	(a)		(b)			
	(c)	36	(d)	50		
8.	Which	n of the following complex ions	s vio	lates the EAN rule?		
	(a)	$[Fe(CO)_5]$. ,	$[Cr(NH_3)_6]^{3+}$		
	(c)	$[Fe(CN)_6]^{4-}$	(d)	$[Mn(H_2O)_6]^{2+}$		
9.	The ef	ffective atomic number for the	com	plex ion $[Ni(CN)_4]^{2-}$ is		
	(a)	36	(b)	86		
	(c)	34	(d)	18		
10.	The II	JPAC name for the complex co	mpo	ound Li[AlH ₄] is		
	(a)	lithium aluminium hydride				
	(b)	hydrido aluminium lithium (l	III)			
	(c)	lithium tetrahydridoaluminat	e (II	I)		
	(d)	lithium tetrahydridoaluminat	e (I)			
11.	The II	JPAC name for the coordination	on co	ompound [CuCl ₂ (CH ₃ NH ₂) ₂] is		
	(a)	dichlorobis (dimethylamine)	copp	er (II)		
	(b)	dichlorobis (methylamine) co	pper	· (II)		
	(c)	dimethylamine copper (II) ch	lorid	e		
	(d)	bis (dimethylamine) copper (l	II) ch	lloride		
12.	The II	JPAC name for $Ba[BrF_4]_2$ is				
	(a)	barium bromofluorite (III)				
		bis (tetrafluorobromium) bari				
		barium bis (tetrafluorobroma		V)		
	(d)	barium tetrafluorobromate (II	Π)			
13.	The II	JPAC name for the coordination	on co	ompound Na ₃ [Ag(S ₂ O ₃) ₂] is		
	(a)	sodium bis (argentothiosulph	ate)	(I)		

(b) sodium bis (thiosulphato) argentate (I)

		sodium silver hyposulphate (I sodium silver thiosulphate (I))	
14.	The o	oxidation number of Co in [Co(e	n) ₃] ₂	$_2(SO_4)_3$ is
		+2		+4
	(c)	+3	(d)	+6
15.	The II	UPAC name for $K[SbCl_5C_6H_5]$ is	3	
		luteroantimonate (V)		
	(b)	potassium pentachloro (pheny	vl) a	ntimonate (V)
	(c)	potassium phenylchloroantim	ona	te (V)
	(d)	potassium benzalantimony (II	I) ch	nloride
16.	The II	UPAC name for $K_2[Cr(CN)_2O_2(C)]$	$O_2)N$	NH ₃] is
	(a)	potassium amminedicyanodic	xop	eroxochromate (VI)
	(b)	potassium amminedicyanotet	roxc	ochromium (III)
		potassium amminedicyanochi		
	(d)	potassium amminocyanodiper	roxc	ochromate (VI)
17.	The st	tructures of Ni(CO) ₄ and Ni(PP	h ₃) ₂ (Cl ₂ are
	(a)	square planar		
		tetrahedral and square planar	resp	pectively
		tetrahedral		
	(d)	square planar and tetrahedral	resp	pectively
18.	The h Fe(CN	hybridization states of the central N_6^{4-} and $Co(NO_2)_6^{3-}$ are	al at	oms in the complexes $Fe(CN)_6^3$
	(a)	d^2sp^3 , sp^3 and d^4s^2 respective	ely	
	(b)	d^2 sp ³ , sp ³ d and sp ³ d ² respect	tive	ly
	(c)	d^2sp^3 , sp^3d^2 and dsp^2 respect	tive	ly
		all d^2sp^3		•
19.	Which	h of the following complexes is	not	easily oxidized?
	(a)	Ni(CO) ₄	(b)	$[Mn(CN)_6]^{5-}$
	(c)	$Cr(CO)_4$	(d)	Fe(CO) ₅
20.	Amor	ng the following complexes, wh	ich i	is the most stable?
	(a)	[Ni(CN) ₄] ⁴⁻	(b)	$[Pd(CN)_4]^{4-}$
	(c)	[Ni(CN) ₄] ³⁻	(d)	[Fe(CO) ₅]
21.	Which	h of the following is colourless?		

(b) [Ti(NO₃)₄]

(d) [Fe(CN)₆] ⁴⁻

(a) $[Ti(H_2O)_6]^{3+}$

(c) $[Cr(NH_3)_6]^{3+}$

22. The oxidation number and coordination number of Pt in the coordination

(b) 4 and 5

compound Na[Pt(C₂H₄)Cl₃] are respectively

(a) 4 and 4

	()		(-)	
	(c)	2 and 3	(d)	2 and 4
23.	The m	agnetic moment of a complex	ion i	s 1.73 BM. The ion is
	(a)	$[Co(NH_3)_6]^{2+}$	(b)	$[MnF_6]^{3-}$
	(c)	[Fe(CN) ₅ NO ⁺]	(d)	$[Mn(CN)_6]^{4-}$
24.	Which	n of the following is not param	agne	etic?
	(a)	$[Cu(NH_3)_4]SO_4$	(b)	$[Mn(H_2O)_6]^{2+}$
	(c)	$[Ag(NH_3)_2]$ Cl	(d)	[NiCl ₄] ²⁻
25.	The IU	JPAC name for [Co(NH ₃) ₆] [C	r(CN	[) ₆] is
	(a)	hexaammine cobalt (III) hexa	cyan	ochromate (III)
	(b)	hexacyanochromium cobalt h	nexaa	immine (VI)
	(c)	hexaammine cobalt (III) hexa	cyan	ochromium (VI)
	(d)	hexacyanochromium (III) hex	kaam	mine cobalt (III)
26.	The IU	JPAC name for [Be ₄ O(CH ₃ CO	O),[]	is
		basic beryllium acetate (II)	70=	
	(b)	hexa-µ-hexakis (acetato) bery	lliun	n (II)
	(c)	hexa- μ -acetato (O, O')- μ_4 -oxo	-tetra	aberyllium (II)
	(d)	$hexaacetato-\mu-oxo-beryllate$ (II)	
27.	The II	JPAC name for [Co(NCS)(NH	₃) ₅]C	l_2 is
	(a)	pentaammine (thiocyanato-N	I) cob	oalt (III) chloride
	(b)	pentaammine (thiocyanato-S)) cob	alt (III) chloride
	(c)	pentaammine (isothiocyanato	o-N,	S) cobalt (III) chloride
	(d)	pentaammine (mercapto-N) o	cobal	t (III) chloride
28.	The II	JPAC name for K ₂ [OsCl ₅ N] is		
	(a)	potassium pentachloroazidoo	osma	te (VIII)
	(b)	potassium pentachloroazoosi	nate	(VI)
	(c)	potassium pentachloronitrido	osm	ate (VI)

29. Among the following aquated metal ions, which has the highest degree

(b) $[Fe(H_2O)_6]^{2+}$ (d) $[Zn(H_2O)_2]^{2+}$

(d) potassium nitroosmate (III)

of paramagnetism? (a) $[Cr(H_2O)_6]^{3+}$

(c) $[Cu(H_2O)_6]^{2+}$

30.	Which of the following is a low-spin (spin-paired) complex?			
	(a)	$[Co(NH_3)_6]^{2+}$	(b)	$[Fe(C_2O_4)_3]^{3-}$
	(c)	$[Ni(NH_3)_6]^{2+}$	(b)	$[Co(NH_3)_6]^{3+}$
31.	Which	n of the following is a high-spir	n (sp	in-free) complex?
	(a)	$[Co(NH_3)_6]^{3+}$	(b)	$Fe(CN)_6$] ⁴⁻
	(c)	$[CoF_6]^{3-}$	(d)	$[Zn(NH_3)_6]^{2+}$
32.	precip	errous ion in a given sample is pitate on the addition of a potas pitate has the constitutional for	ssiun	cted by the formation of a white n ferrocyanide solution to it. The
	(a)	K_2 Fe II [Fe II (CN) ₆]	(b)	$K_2Fe^{III}[Fe(CN)_6]$
	(c)	K Fe III [Fe II (CN) ₆]	(d)	$KFe^{II}[Fe^{III}(CN)_6]$
33.	The fo	ormula of the complex potassi	um t	richloro (ethylene) platinate (II)
	(a)	$K[PtCl_3(C_2H_4)]$	(b)	$K_2[PtCl_3(C_2H_4)_3]$
	(c)	$K_4[PtCl_3(C_2H_4)]$	(d)	$K_3[Pt_2Cl_3(C_2H_4)_3]$
34.	The fo	ormula of the complex sodium	hyd	ridotrimethoxyborate (III) is
	(a)	$Na_4[BH_2(OCH_3)_3]$	(b)	$Na_2[BH(OCH_3)_3]$
	(c)	$Na[BH_2(OCH_3)_3]$	(d)	$Na[BH(OCH_3)_3]$
35.	The fo	ormula of the complex triammi	netri	i (nitrito-N) cobalt (III) is
	(a)	$[Co(ONO_2)_3(NH_3)_3]$	(b)	[Co(NO2)3(NH3)3]
	(c)	$[Co(ONO)_3(NH_3)_3]$	(d)	$[Co(NO_2)(NH_3)_3]$
36.	(a) (b) (c)	JPAC name for [(CO ₅)Mn—Me bis (pentacarbonylmanganate bis (pentacarbonyl dimangan bis (pentacarbonyl manganes decacarbonyldimanganate (V	e) (V] ese) e)	
37.	(a) (b) (c) (d)	JPAC name for $[Pt(py)_4][PtCl_4$ tetrakis (pyridine platinum (I tetrapyridine tetrachlorodipla tetrachlorotetrapyridine dipla tetrakis (pyridine) platinum (I) tet itinu itinu IV) t	m (IV) m (II) etrachloroplatinum (IV)
38.		UPAC name for [(NH ₃) ₅ Cr–OH		
	(a)	μ-hydroxo-bis (pentaammine	dich	aromium) (5+) ion
	(b)	μ-hydroxo-bis (decaammine d	dichr	romium (5+) ion

- (c) μ-hydroxo-bis (octaammine chromium) (5+) ion
- (d) μ-hydroxo-bis (pentammine chromium) (III)
- 39. The IUPAC name for $[Co(ONO)(NH_3)_5]SO_4$ is
 - (a) pentaammine (nitrito-O) cobalt (III) sulphate
 - (b) pentaammine (nitroso-N) cobalt (III) sulphate
 - (c) pentaammine (nitro-O) cobalt (III) sulphate
 - (d) pentaammine (nitroyl-O) cobalt (III) sulphate
- **40.** The IUPAC name for $[(CO)_3Fe(CO)_3Fe(CO)_3]$ is
 - (a) tri-µ-hexacarbonyl iron (II)
 - (b) tri-μ-carbonyl-bis (tricarbonyl iron)
 - (c) tri-µ-carbonyl-bis (tricarbonyl di-iron)
 - (d) tri-µ-manocarbonyl iron (III)
- **41.** The hybridization states of the central atoms in the complex ions $[FeF_6]^{3-}$, $[Fe(H_2O)_6]^{3+}$ and $[Ni(NH_3)_6]^{2+}$ are
 - (a) $3s4p^34d^2$, $4d3s4d^2$ and $4d^45s^2$ respectively
 - (b) all $3d^24s4p^3$
 - (c) all $4s4p^34d^2$
 - (d) $3s4p^34d^2$, $4d3s4p^3$ and $4p^45d^2$ respectively
- **42.** The formation of the complex ion $[Co(NH_3)_6]^{3+}$ involves the sp 3 d 2 hybridization of Co $^{3+}$. Therefore the complex ion should have
 - (a) an octahedral geometry
 - (b) a tetrahedral geometry
 - (c) a square-planar geometry
 - (d) a square-antiprismic geometry
- 43. [Co(NH₃)₅NO₃]SO₄ and [Co(NH)₅SO₄]NO₃ exhibit
 - (a) coordinate isomerism
- (b) linkage isomerism
- (c) ionization isomerism
- (d) optical isomerism
- **44.** What type of isomerism do the forms $[Co(NH_3)_6][Cr(CN)_6]$ and $[Cr(NH_3)_6][Co(CN)_6]$ exhibit?
 - (a) Coordination position isomerism
 - (b) Coordination isomerism
 - (c) Ligand isomerism
 - (d) Polymerization isomerism
- **45.** How many isomers can $CrCl_3 \cdot 6H_2O$ have?
 - (a) Six

(b) Four

(c) Three

(d) Two

46. Which of the following will have three isomeric forms?

 $[Cr(NO_3)_3(NH_3)_3] \ (I), \ K_3[Co(C_2O_4)_3] \ (II), \ K_3[Co(C_2O_4)_2Cl_2] \ (III) \ and \ [Co(en)_2ClBr] \ (IV) \ where \ en = ethylene \ diamine.$

(a) IV and III

(b) IV and

(c) III and II

- (d) I and II
- **47.** Which among the following pairs of complex compounds is an example of linkage isomerism?
 - (a) $[PtCl_2(NH_3)_4]Br_2$ and $[PtBr_2(NH_3)_4]Cl_2$
 - (b) $[Cu(NH_3)_4][PtCl_4]$ and $[CuCl_4][Pt(NH_3)_4]$

(c)
$$\left[(NH_3)_4 Co \underbrace{NH_2}_{O_2} Co(NH_3)_2 Cl_2 \right]^{2+}$$
 and $\left[(NH_3)_3 ClCo \underbrace{NH_2}_{O_2} CoCl(NH_3)_3 \right]^{2+}$

- (d) $[Co(NO_2)(NH_3)_5]Cl$ and $[Co(ONO)(NH_3)_5]Cl_2$
- **48.** Which of the following statements is correct?
 - (a) The $[Ni(CN)_4]^{2-}$ ion has tetrahedral geometry and is diamagnetic.
 - (b) The $[Ni(CN)_4]^{2-}$ ion has a square-planar geometry and is paramagnetic.
 - (c) The $[Ni(CN)_4]^{2-}$ ion has a square-planar geometry and is diamagnetic.
 - (d) The [Cu(NH₃)₄]²⁺ ion has a tetrahedral geometry and is diamagnetic.
- **49.** Which of the following statements is correct?
 - (a) Most four-coordinated complexes of Ni ²⁺ ions are square planar rather than tetrahedral.
 - (b) The $[Fe(H_2O)_6]^{3+}$ ion is more paramagnetic than the $[Fe(CN)_6]^{3-}$ ion.
 - (c) Square-planar complexes are more stable than octahedral complexes.
 - (d) The $[Fe(CN)_6]^{4-}$ ion is paramagnetic but $[Fe(CN)_6]^{3-}$ ion is diamagnetic.
- **50.** Which of the following statements is true?
 - (a) $Ni(CO)_4$ and $[NiCl_4]^{2-}$ are diamagnetic, and $[Ni(CN)_4]^{2-}$ is paramagnetic.

- (b) $[NiCl_4]^{2-}$ and $[Ni(CN)_4]^{2-}$ are diamagnetic, and $[Ni(CO)_4]$ is paramagnetic. (c) $[Ni(CO)_4]$ and $[Ni(CN)_4]^{2-}$ are diamagnetic, and $[NiCl_4]^{2-}$ is paramagnetic. (d) $[Ni(CO)_4]$ is diamagnetic, and $[NiCl_4]^{2-}$ and $[Ni(CN)_4]^{2-}$ are paramagnetic.
- 51. How many geometrical isomers are possible for the square-planar complex $[Pt(NO_2)(py)(NH_3)(NH_2OH)]NO_2$?
 - (a) Four

(b) Five

(c) Eight

- (d) Three
- 52. Tetrahedral complexes of the types Ma₄ and Ma₃b (where M stands for a metal, and a and b are achiral ligands) do not show optical isomerism because they have
 - (a) a C_n axis of symmetry
 - (b) a plane of symmetry and hence are achiral
 - (c) a centre of symmetry
 - (d) nonsuperimposable mirror images
- 53. Which of the following types of octahedral complexes exhibit geometrical isomerism (M stands for a metal, and a and b are achiral ligands)?
 - (a) $[Ma_6]$

(b) [Ma₅b]

(c) $[M(aa)_3]$

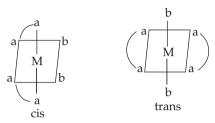
- (d) $[Ma_1b_2]$
- 54. Which of the following types of square-planar complexes can show geometrical isomerism (M stands for a metal, and a and b are achiral ligands)?
 - (a) Ma₄

(b) Ma₃b

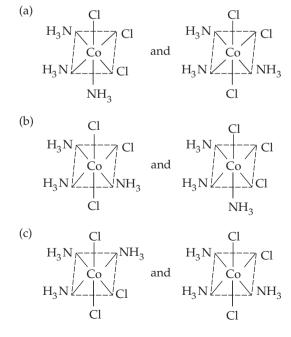
(c) Ma₂b₂

- (d) Mab₃
- 55. The tris (ethylenediamine) cobalt (III) cation, $[Co(en)_3]^{3+}$, can have
 - (a) three stereoisomers, all chiral and optically active
 - (b) two chiral stereoisomers (enantiomers)
 - (c) three stereoisomers, all achiral
 - (d) two stereoisomers, both achiral
- **56.** Which of the following statements is correct?
 - (a) $[Co(NH_3)_6]^{2+}$ is oxidized to diamagnetic $[Co(NH_3)_6]^{3+}$ by the oxygen in air.
 - (b) $[Fe(CN)_6]^{3-}$ is stable but $[FeF_6]^{3-}$ is unstable.
 - (c) $[Co(H_2O)_6]^{2+}$ is easily oxidized to $[Co(H_2O)_6]^{3+}$.
 - (d) None of these

57. Which of the following statements is correct regarding the chirality (optical isomerism) of the cis and trans isomers of the type M(aa)₂b₂ (M stands for a metal, a and b are achiral ligands, and aa is a bidentate ligand)?

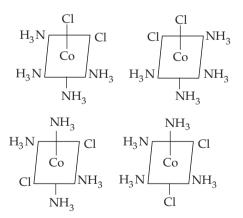


- (a) The trans form is achiral and optically inactive while the cis form is chiral and exists in two enantiomeric forms.
- (b) The cis as well as the trans form are achiral and optically inactive.
- (c) The trans form is chiral and exists in two enantiomeric forms while the cis form is achiral and optically inactive.
- (d) The cis as well as the trans form are chiral, and each of them exists in two enantiomeric forms.
- 58. Which of the following pairs of structures represents facial and meridianal isomers (geometrical isomers) respectively?



(d) Cl
$$NH_3$$
 H_3N Cl Cl Cl NH_3 NH_3 NH_3 NH_3 NH_3 NH_3 NH_3 NH_3

- **59.** Which of the following statements is correct regarding the stereoisomerism of the complexes referred to (M stands for a metal, and a, b, c, and d are achiral ligands)?
 - (a) In the cis and trans isomers of the octahedral complex $M(a_2b_4)$, the two ligands occupy the positions 1, 2 and 1, 6 respectively.
 - (b) The facial and meridianal isomers of the octahedral complex $M(a_3b_3)$ have three identical ligands (a or b) at positions 1,2,3- and 1,2,6-respectively.
 - (c) The octahedral complex M(abcd $\stackrel{\frown}{ee}$) where $\stackrel{\frown}{ee}$ is a bidentate ligand should have twelve stereoisomers.
 - (d) The facial as well as the meridianal isomers of the octahedral complex of the type Ma_3b_3 are optically active.
- **60.** Consider the following arrangements of the octahedral complex ion $[Co(NH_3)_4Cl_2]^+$.



Which of the following statements is incorrect?

- (a) I and II are enantiomers.
- (b) II and III are cis and trans isomers respectively.
- (c) III and IV are trans and cis isomers respectively.
- (d) II and IV are identical structures.

• Type 2 •

Choose the correct options. More than one option is correct.

61.	. Which of the following will produce a white p	precipitate upon reacting
	with AgNO ₃ ?	

(a) $[Co(NH_3)_6]Cl_3$

(b) Co(NH₃)₃Cl₃]

(c) $K_2[Pt(en)_2Cl_2]$

- (d) [Fe(en)₃]Cl₃
- 62. In which of the following does the central atom exhibit an oxidation state of ± 2 ?
 - (a) $K_2[Ni(CN)_4]$

(b) $K_4[Fe(CN)_6]$

(c) $[Fe(C_2O_4)_3]^{3-}$

- (d) $[Cu(NH_3)_4]^{2+}$
- 63. In which of the following are the chemical formula and the name correctly matched?
 - (a) K[Pt(NH₃)Cl₅]—potassium amminepentachloroplatinate (IV)
 - (b) [Ag(CN)₂] —dicyanoargentate (I) ion
 - (c) $K_3[Cr(C_2O_4)_3]$ —potassium trioxalatochromate (III)
 - (d) Na₂[Ni(EDTA)]—Sodium ethylenediaminetetra-acetonickel (I)
- **64.** Which of the following are π -bonded organometallic compounds?
 - (a) Ferrocene

- (b) Diethyl zinc
- (c) Ethylmagnesium iodide
- (d) Dibenzene chromium
- 65. Which of the following exhibit geometrical isomerism (M stands for a metal, and a and b are achiral ligands)?
 - (a) Ma_2b_2 (b) Ma_4b_2
- (c) Ma₅b
- (d) Ma₆

Answers

1. d	2. c	3. b	4. b	5. d
6. b	7. a	8. d	9. c	10. c
11. b	12. d	13. b	14. c	15. b
16. a	17. c	18. d	19. b	20. d
21. b	22. d	23. d	24. c	25. a
26. c	27. a	28. c	29. b	30. d
31. c	32. a	33. a	34. d	35. b
36. c	37. a	38. d	39. a	40. b
41. c	42. a	43. c	44. b	45. c
46. b	47. d	48. c	49. d	50. c
51. d	52. b	53. d	54. c	55. b

56. a	57. a	58. a	59. d	60. a
61. a, d	62. a, b, d	63. a, b, c	64. a, d	65. a, b

• General Hints •

Read the following rules of nomenclature in order to answer questions 4, 5, 10, 11, 12, 13, 15, 16, 25, 27, 28, 33, 34, 35, 36, 37, 38, 39, 40, and 63.

- The symbol of the central atom is placed first in the formula of coordination compounds followed by those of anionic ligands in alphabetical order and then those of neutral ligands, again in alphabetical order.
- 2. Abbreviations of complicated organic formulae may be used in formulae.
- 3. The formula for a complex molecule or ion is enclosed in square brackets.
- **4.** In names, that of the central atom is placed after the names of the ligands.
- The neutral ligand H₂O is called aquo, and the neutral ligand NH₃ ammine.
- **6.** For names, ligands are listed in alphabetical order regardless of the charge on the ligand or the number of ligands.
- 7. The names of anionic ligands end in 'O' (-ido, -ito, and ato commonly). The number of ligands of each kind is indicated by the prefixes di-, tri-, tetra-, penta-, hexa-, and so on, unless the prefix could be misinterpreted as a part of the name of the ligand. In such cases the prefixes bis, tris, tetrakis and so on, are used. For example, diammine for 2NH₃ but bis (methylamine) for 2CH₃NH₂, since dimethylamine is (CH₃)₂NH. The following are the exceptions for the names of anionic ligands (omit -id of -ido).

F—fluoro Cl—chloro Br—bromo I—iodo
$$O^2$$
—oxo OH^- —hydroxo O_2^2 —peroxo S^2 —thio CN^- —cyano.

- **8.** The oxidation number of the central atom is indicated by the corresponding Roman numeral in parenthesis after its name.
- **9.** Formulae and names may be supplemented by the italicized prefixes *cis, trans, fac* (facial), *mer* (meridianal) and so on.
- **10.** Names of metal complex anions end in -ate. Complex cations and neutral molecules, e.g., NO and CO, do not have a distinguishing suffix.
- **11.** A ligand, e.g., SCN⁻, may be attached to the rest of the molecule through different atoms in different cases. These molecules may be distinguished as follows.
 - M-SCN stands for thiocyanato-S and M-NCS stands for thiocyanato-N

12. Bridging groups are indicated by prefixing the Greek letter μ to the names of the groups. Two or more bridging groups of the same kind are indicated by di- μ , and so on. If a bridging group bridges more than two metals, use μ_3 , μ_4 , etc.

• Effective Atomic Number Rule (For questions 6, 7, 8 and 9) •

The Effective Atomic Number (EAN) provides a rough guide for bonding in coordination compounds. Quite a few, but not all, metals achieve the EAN of a noble gas through coordination. The EAN of a metal ion is calculated by adding the electrons of the metal ion to those shared with it through coordination. In [Co(NH₃)₆]³⁺, the cobalt (III) ion has 24 electrons plus 6 electron pairs from NH₃ molecules, making up a total of 36 electrons—the configuration of krypton.

EAN of cobalt ion in $\left[\text{Co(NH}_3)_6\right]^{3+}$ = $(27-3)+6\times2=36$ = atomic number of Kr

• Coordination Compounds •

Coordination compounds are formed when the available empty bonding orbitals of a metal are used for the formalion of coordinate bonds. The coordination number (CN) and geometry are determined in part by size and charge effects, but also to a great extent by the orbitals available for bonding. The common hybridized orbitals encountered in coordination compounds are given in Table 1.

See Tables 1 and 2 for answering Questions 18, 23, 41, 42, 48, 49 and 50.

Coordination no.	Hybridized orbital	Configuration	Examples
2	sp	Linear	$Ag(NH_3)_2^+$
3	sp ²	Trigonal planar	$Ag(PR_3)_3^+$
4	sp ³	Tetrahedral	$Ni(CO)_4$, $Zn(NH_4)_4^{2+}$
4	dsp ²	Planar	$Ni(CN)_4^{2-}$, $Pt(NH_3)_4^{2+}$
5	$d_z^2 sp^3$ or $d^3 sp$	Trigonal bipyramidal	CuCl ₅ ³⁻ , [Ni(PEt ₃) ₂ Br ₃]
6	d^2sp^3	Octahedral	$Co(NH_3)_6^{3+}$, $PtCl_6^{2-}$

Table 1 Hybrid orbitals for metal complex

- **1.** Those complexes in which there are no low-energy d-orbitals available for bond formation, e.g., $\operatorname{Ni}(\operatorname{NH}_3)_6^{2+}$ or $[\operatorname{Fe}(C_2O_4)_3]^{3-}$ (five unpaired electrons), are called "ionic" complexes.
- 2. Whether the electron populations of the d-orbitals are "low-spin" (or spin-paired) or high-spin (or spin-free) determine the magnetic properties of the molecule.
- **3.** Covalent octahedral complexes and tetrahedral complexes are formed when d ²sp ³ and sp ³ orbitals respectively come into play. (See Table 2.)

 Table 2
 Electronic configurations of some metal complexes according to the valence bond (VB) theory

Ion or Complex	Configuration	High- or low-spin
Cr ³⁺	3d 4s 4p ↑ ↑ ↑	High-spin,
$Cr(NH_3)_6^{3+}$	$\uparrow \uparrow \uparrow \uparrow \boxed{\uparrow \downarrow \uparrow \downarrow \uparrow \downarrow \uparrow \downarrow \uparrow \downarrow \uparrow \downarrow}$	paramagnetic
Co ³⁺ , Fe ²⁺	Covalent (VB) $\uparrow \downarrow \uparrow \uparrow \uparrow \uparrow$	
$Co(NH_3)_6^{3+}$, $Fe(CN)_6^{4-}$	$\uparrow\downarrow\uparrow\uparrow\uparrow\uparrow\uparrow\uparrow$ $\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow$	Low-spin, diamagnetic
Co ²⁺	Covalent (VB) $\uparrow \downarrow \uparrow \downarrow \uparrow \uparrow \uparrow \uparrow$	
$Co(NH_3)_6^{2+}$	Ionic (VB) $\uparrow \downarrow \uparrow \downarrow \uparrow \uparrow \uparrow \uparrow$	High-spin,
Zn ²⁺	1) 1) 1) 1) 1	paramagnetic
$ZnCl_4^{2-}$	$\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow$ $Ionic (VB)$	
$Zn(NH_3)_6^{2+}$ Fe ³⁺	$\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow\uparrow\downarrow$	
Fe(C ₂ O ₄) ₃ ³⁻ Ni(NH ₃) ₆ ²⁺	$ \uparrow \downarrow \uparrow \uparrow \uparrow \uparrow \downarrow \uparrow \uparrow$	High-spin, paramagnetic

• Hints to Specific Problems •

- 2. A hexadentate ligand will contain six donor atoms.
- 3. Dimethyl glyoxime contains two donor atoms.
- **6.** In $Cu(CN)_4^{3-}$, Cu has an oxidation number of 1. Z = 29 for Cu. CN^- has one electron pair. Hence $EAN = (29 1) + 4 \times 2 = 36$ (Kr)
- 7. In $[Pd(NH_3)_6]^{4+}$ (Z = 46 for Pd), oxidation number of Pd = +4. EAN = $(46 - 4) + 6 \times 2 = 54$
- 8. In $[Mn(H_2O)_6]^{2+}$ (Z = 25 for Mn), oxidation number of Mn = +2.

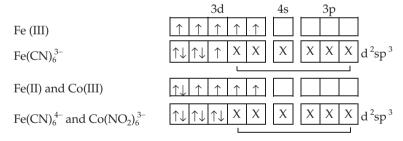
$$\therefore$$
 EAN = $(25 - 2) + 6 \times 2 = 35$

This value does not correspond to the atomic number of any noble gas. So the EAN rule is violated.

18. In Fe(CN) $_6^{3-}$, the oxidation number of Fe = +3 and so it has a 3d 5 configuration.

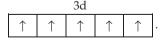
In $Fe(CN)_6^{4-}$, the oxidation number of Fe = +2 and so it has a 3d 6 configuration.

In $Co(No_2)_6^{3-}$ the oxidation number of Co = +3 and so it has a 3d 6 configuration.

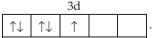


- **19.** The CN ion is a strong ligand.
- 21. There is no charge-transfer transition in Ti 4+.
- 23. In $[Mn(CN)_6]^{4-}$, Mn is in the +2 oxidation state.

The configuration of Mn^{+2} is $3d^{5}$ or



The configuration of $[Mn(CN)_6]^{4-}$ is



n = 1 (no. of unpaired electron)

The unpaired electrons in the d orbitals pair up because CN^- has a strong ligand character.

$$\mu_{\text{spin}} = \sqrt{n(n+2)}$$
 BM = $\sqrt{(1+2)} = \sqrt{3}$ BM = 1.73 BM

- **29.** The configuration of Fe²⁺ is 3d⁶. It contains four unpaired electrons more than any of the other metal ions shown.
- **30.** Co exists as Co^{3+} in $[Co(NH_3)_6]^{3+}$. The configuration of Co^{3+} is $3d^6$

The configuration of $[Co(NH_3)_6]^{3+}$ is

30					
$\uparrow\downarrow$	$\uparrow\downarrow$	$\uparrow\downarrow$			١.

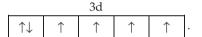
This complex is spin-paired.

The unpaired electrons in the d orbitals pair up because NH_3 has a strong ligand character.

31. Co exists as Co^{3+} in $[CoF_6]^{3-}$. The configuration of Co^{3+} is $3d^6$ or



The configuration of CoF_6^{3-} is



The unpaired electrons in the d orbitals do not tend to pair up because F^- is weak ligand. (Ligands which cause a small degree of crystal-field splitting are said to be weak.)

Hence, this complex is spin-free.

- 44. Exchange of Co and Cr in the coordination sphere
- **47.** Co is linked with —N and with —O respectively in the two compounds shown.

17

Analytical Chemistry

• *Type 1* •

1. Which among the following nitrite salts is sparingly soluble in water?

(b) AgNO₂

Choose the correct option. Only one option is correct.

(a) KNO₂

(c) AgBr

	(c)	$Ba(NO_2)_2$	(d)	NH ₄ NO ₂
2.		eating, a white amorphous incooling, turns white again. T		nic compound becomes yellow alt may be
	(a)	PbCO ₃	(b)	$MgCO_3$
	(c)	ZnCO ₃	(d)	K ₂ CO ₃
3.	Which	n of the following does not sub	lime	on heating?
	(a)	NH ₄ Cl	(b)	As_2O_3
	(c)	HgCl ₂	(d)	Sb_2S_3
4.	Which	n of the following does not resp	ond	to the chromyl chloride test?
	(a)	$HgCl_2$	(b)	KCl
	(c)	CaCl ₂	(d)	NaCl
5.		eing heated with concentrated O_7 produces a gas. The gas is	H ₂ S0	O ₄ , a mixture of solid NaCl and
		SO ₂	(h)	CrO ₂ Cl ₂
	` '	-		
	(C)	HCl	(a)	SO_3
6.	Which	_	aln	nost insoluble in concentrated
	(a)	AgI	(b)	AgF

2-171

(d) AgCl

reagent in the presence of a CCl₄ solvent. The reagent is

8. The hottest part of the flame of a Bunsen burner is the

(a) an AgNO₃ solution

(c) a Ba(OH)₂ solution

(a) blue zone

7. Bromide and iodide ions can be detected in a mixture of NaCl, NaBr and NaI by shaking an aqueous solution of the mixture with a particular

(b) an FeSO₄ solution

(d) chlorine water

(b)	zone of complete combustion				
(c) zone of partial combustion					
(d) All parts of the flame are equally hot.					
drops	of dilute KMnO ₄ solution loses	s col	our. The salt may be a		
	•	` '	carbonate		
(c)	bicarbonate	(d)	nitrate		
		nic :	salt solution produces a white		
(a)	Hg_2^{2+}	(b)	Hg^{2+}		
(c)	Zn^{2+}	(d)	Cd^{2+}		
(a)	Al^{3+}		Cu ²⁺		
(b)	Ba ²⁺	(d)	Zn^{2+}		
		st a	KMnO ₄ solution, the indicator		
(a)	methyl orange	(b)	potassium ferrocyanide		
(c)	KI/starch	(d)	no indicator is used		
		nd te	st, a salt produces a pink bead.		
(a)	an iron salt	(b)	a copper salt		
(c)	a nickel salt	(d)	a manganese salt		
punge mercu	ent gas is produced which turr crous nitrate solution black. Th	is mo	oist red litmus paper blue and a		
(a)	CO ₃ ²⁻	(b)	NH_4^+		
(c)	Sn ⁴⁺	(d)	NO_2^-		
	(c) (d) A salt drops (a) (c) On troprecip (a) (c) In a cligives a (a) (b) In the used i (a) (c) On be The sa (a) (c) When pungemercu contai (a)	(d) All parts of the flame are equal A salt heated with dilute H ₂ SO ₄ ardrops of dilute KMnO ₄ solution loses (a) sulphite (c) bicarbonate On treatment with HCl, an inorgaprecipitate. The salt may contain (a) Hg ₂ ²⁺ (c) Zn ²⁺ In a charcoal cavity test, a colourles gives a green precipitate. The salt contain (a) Al ³⁺ (b) Ba ²⁺ In the titration of oxalic acid again used is (a) methyl orange (c) KI/starch On being subjected to the borax-beat The salt is (a) an iron salt (c) a nickel salt When the solution of an inorganic spungent gas is produced which turn	(c) zone of partial combustion (d) All parts of the flame are equally he as alt heated with dilute H ₂ SO ₄ and so drops of dilute KMnO ₄ solution loses cole (a) sulphite (b) (c) bicarbonate (d) On treatment with HCl, an inorganic precipitate. The salt may contain (a) Hg ₂ ²⁺ (b) (c) Zn ²⁺ (d) In a charcoal cavity test, a colourless salgives a green precipitate. The salt contain (a) Al ³⁺ (b) (b) Ba ²⁺ (d) In the titration of oxalic acid against a used is (a) methyl orange (b) (c) KI/starch (d) On being subjected to the borax-bead terms alt is (a) an iron salt (b) (b) a nickel salt (d) When the solution of an inorganic salt in pungent gas is produced which turns memoreurous nitrate solution black. This excontains (a) CO ₃ ²⁻ (b)		

15. Which of the following reagents may be used to identify ${\rm Sr}^{2+}$ ions in the presence of Ca $^{2+}$ ions in a solution?

(b) (NH₄)₂S₂O₈(d) (NH₄)₂CO₃

(a) NH₄Cl

(c) $(NH_4)_2SO_4$

16.	. Which of the following reagents may be used to identify Ca ²⁺ ions in the presence of Ba ²⁺ ions in a hot dilute acetic acid medium?			
	(a)	$(NH_4)_2CO_3$	(b)	NH ₄ Cl
	(c)	$AgNO_3$	(d)	$(NH_4)_2C_2O_4$
17.		n of the following metal ions ting flame of a Bunsen burner y		its volatile chloride) turns the w-green?
	(a)	Cr ³⁺	(b)	Zn^{2+}
	(c)	Fe ³⁺	(d)	Ba ²⁺
18.		norganic salt turns a borax bea a reducing flame, then the meta		ue in an oxidizing flame as well esent in the salt is
		Co	(b)	
	(c)	Cr	(d)	Ni
19.		ım cannot be separated by pas	sing	
	(a)	Ca^{2+} and Hg_2^{2+}	(b)	Cu ²⁺ and Cd ²⁺
	(c)	Zn^{2+} and Sn^{4+}	(d)	Co ³⁺ and Cu ²⁺
20.		ulphides of which of the follow ammonium sulphide?	ing g	groups of elements are soluble in
		As, Sb and Sn		As, Cd and Sn
	(c)	Cd, Cu and Bi	(d)	Hg, Cu and Cd
21.		and Na ₂ SO ₄ can be distinguish		2
		Gutzeit's test the flame test		Marsh's test
	, ,			the chromyl chloride test
22.	white	ging heated with SiO_2 and confirmed which give a white dened glass rod. The salt may be	epos	ntrated H_2SO_4 , a salt produces it on coming in contact with a
		CaSO ₄		NaCl
	(c)	CaF ₂	(d)	KBr
23.		ganic compound containing su s treatment, S is converted to	ılphı	ur is oxidized by fuming HNO ₃ .
	(a)	SO ₂	(b)	SO ₃
	(c)	H_2S	(d)	H_2SO_4

(c) after testing for Group 2 but before doing so for Group 3

25. The green colour formed by adding KOH to KMnO₄ is due to the

26. Na₂SO₃ and Na₂S can be distinguished from each other by using

(b) Mn²⁺(d) MnO₂

24. In salt analysis, the interference due to PO_4^{3-} is removed

(d) before proceding with the basic radicals

(a) after testing for Group 1(b) after testing for Group 2

formation of
(a) MnO_4^{2-}

(c) Mn_2O_7

(a) concentrated H₂SO₄

	(b)	an acidified KMnO ₄ solution			
	(c)	an acidified K ₂ Cr ₂ O ₇ solution	l		
	(d)	a sodium nitroprusside solut	ion		
27.	passe water	d successively through an ac	cidifi lutio	ICl and the gaseous product is ed $K_2Cr_2O_7$ solution and limenchanges from orange to green, salt mixture contains	
	(a)	CO ₃ ²⁻	(b)	SO ₃ ²⁻	
	(c)	S ²⁻	(d)	CO_3^{2-} as well as SO_3^{2-}	
28.	The a	ddition of solid NH ₄ Cl to an a	queo	us solution of ammonia	
	(a)	reduces the concentration common-ion effect	of th	ne hydroxide ions due to the	
	(b)	increases the concentration common-ion effect	of t	he hydroxide ions due to the	
	(c)	increases the dissociation of Cl—OH	f am	monia due to the formation of	
	(d)	does not change the concentr	atior	of the hydroxide ions	
29.	As_2S_3	dissolves in yellow ammonium	n sul	phide to form	
	(a)	$(NH_4)_3AsS_3$	(b)	$(NH_4)_3AsS_4$	
	(c)	$(NH_4)_2AsS_4$	(d)	$(NH_4)_4AsS_3$	
30.	30. On treatment with KI, mercuric chloride gives a product X which is soluble in an excess of KI to form another product Y. X and Y are, respectively,				
	(a)	HgI ₂ and HgI ₃	(b)	HgI ₂ and K ₂ HgO ₃	
	(c)	HgI ₂ and K ₂ HgI ₄	(d)	Hg ₂ I ₂ and K ₂ HgI ₄	

31. The yellow precipitate formed by passing ammonia gas into Nessler's

32. The purple colour obtained by adding a sodium nitroprusside solution to

a sodium sulphide solution is due to the formation of

(b) NH₂O—Hg—HgI

(b) Na₄[Fe(CN)₅NOS]

(d) NH₂—Hg—I

reagent is due to the formation of

(c) NH₂—Hg—O—Hg—I

(a) HgI_4^{2-}

(a) $Na_2[Fe(CN)_6S]$

	(c)	Na ₂ [Fe(CN)	6NOS]		(d)	KFe[Fe(CN)	₆ NO	S]
33.	3. An excess of an NaOH solution is added gradually to an aqueous solution of ZnSO ₄ . Which of the following happens?							
	(a)	A white prexcess of Na		te is forme	ed w	which does a	not d	lissolve in the
	(b)	A green pre the excess o			l wh	ich dissolves	on t	he addition of
	(c)	A white pronounce NaOH.	ecipitat	te is forme	ed w	hich dissolv	es in	the excess of
	(d)	No observal	ole char	nge takes p	lace.			
34.	ions a		, it is n	ecessary to	boil	the solution	of th	Group 3 metal ne salt mixture o convert
	(a)	Co ²⁺ to Co ³	+		(b)	Fe ²⁺ to Fe ³⁺		
	(c)	Mn ²⁺ to Mn	O_4^-		(d)	Cr ³⁺ to CrO	$_{4}^{2-}$	
35.	Which		lowing	pairs of	com	pounds can	exi	st together in
	(a)	FeSO ₄ and I	b(NO ₃	2)2	(b)	Na ₂ CO ₃ and	l Na(OH
	(c)	NaHCO ₃ an	d NaO	Н	(d)	AgNO ₃ and	NaC	21
36.	colloi		n with	the evolut				duces a yellow The gas turns
	(a)	$NaHCO_3$	(b) N	Ia_2SO_3	(c)	$Na_2S_2O_3$	(d)	$Na_2S_2O_7$
37.								btained which the presence of
	(a)	Fe ²⁺	(b) P	b ²⁺	(c)	Hg^{2+}	(d)	Cd ²⁺
38.	of alu Whic	ıminium, zind	and m wing g	nagnesium	prod	duce residue	s of s	It nitrate), salts specific colour. mposition and

41.

42.

43.

44.

39. In a borax-bead test, a colourless bead becomes coloured on being heated with a colourless metal salt. This happens due to the formation of

40. An insoluble salt (A) is readily soluble in concentrated ammonium

(a) $CoAl_2O_4 \longrightarrow Thenard's blue$

(c) the orthoborate of the metal(d) the hexaborate of the metal

(d) All of these

(b) boric oxide

(b) $CoZnO_2 \longrightarrow Rinmann's green$ (c) $CoMgO_2 \longrightarrow Pale pink$

(a) the borate and metaborate of the metal

	e solution. The resulting soluti v precipitate which is insoluble		n treatment with K ₂ CrO ₄ gives a nineral acid. Identify A		
-	PbSO ₄		Ag_2SO_4		
(c)	$HgSO_4$	(d)	CaSO ₄		
The p	resence of phosphate interferes	wit	h the detection of		
(a)	Pb ²⁺	(b)	Ca ²⁺		
(c)	Sn ²⁺	(d)	Bi ³⁺		
A salt is heated with dilute H_2SO_4 and subsequently treated with a few drops of dilute $K_2Cr_2O_7$, upon which the entire solution turns green. The salt may be a					
(a)	chloride	(b)	nitrate		
(c)	sulphide	(d)	sulphate		
	n of the following statement is anic salts in an oxidizing flame		orrect for the borax-bead test of		
(a)	The formation of a green be indicates the presence of Cu ²	ead v	which turns blue upon cooling he salt.		
(b)	The formation of a blue bead indicates the presence of Co ²⁴		ich retains its colour on cooling he salt.		
(c)	The formation of a dark yellobecomes green upon cooling salt.	ow b indi	pead in the hot condition which cates the presence of Cr ³⁺ in the		
(d)	The formation of a green bear indicates the presence of Mn ²	nd in	hot as well as cold conditions the salt.		
	n of the following pairs of cous solutions are mixed togethe		ounds precipitates when their		
(a)	AlCl ₃ and ZnSO ₄	(b)	FeSO ₄ and BaCl ₂		
(c)	Cu(NO ₃) ₂ and NH ₄ Cl	(d)	NaF and AgNO ₃		

45. Which of the following is insoluble in dilute acids but soluble in alkalis?

46. The constituents of which of the following pairs of ions can be separated

(a) PbS(c) FeS

using a concentrated NaOH solution?

(b) CdS

(d) Sb_2S_3

		Al ³⁺ and Sn ²⁺	. ,	Al ³⁺ and Fe ³⁺
	. ,	Al ³⁺ and Zn ²⁺		Zn ²⁺ and Pb ²⁺
47.	The brown ring test is performed for the detection of the			
	` '	nitrite ion		sulphite ion
	(c)	bromide ion	(d)	nitrate ion
48.	using	n of the following metal ions dimethyl glyoxime in an amm Ni ²⁺	onia	be detected and estimated by cal medium? Co ²⁺
	` '	Cd ²⁺		Mg ²⁺
	. ,			<u> </u>
49.	. Which of the following reagents can one use to distinguish betweer Na ₂ CO ₃ and Na ₂ SO ₃ ?			
		Limewater	(b)	Baryta water
	(c)	Acidified K ₂ Cr ₂ O ₇ solution	(d)	H ₂ SO ₄ solution
50.	50. Cu ²⁺ and Cd ²⁺ are detected in a mixture of their solutions by using			
	(a) KCN and H ₂ S			
	(b)	HCl and H ₂ S		
	(c)	K ₃ [Fe(CN) ₆] and H ₂ S		
	(d)	concentrated NH ₃ and H ₂ S		
51.	When a reagent A reacts with Fe ³⁺ the solution turns red due to the formation of a compound B. The reagent causes no change in colour when it reacts with Fe ²⁺ in the pure state. A and B are respectively			
		K ₄ [Fe(CN) ₆] and Fe ₄ [Fe(CN) ₆ . NH ₄ CNS and [Fe(SCN)] ²⁺	l3	
	` '	/-	. 1	
		K ₃ [Fe(CN) ₆] and K ₂ Fe[Fe(CN]	<i>1</i> 61	
		Na ₂ HPO ₄ and FePO ₄	_	
52.	An aqueous solution of $FeSO_4 \cdot Al_2(SO_4)_3 \cdot 24H_2O$ and chrome alum i heated with an excess of Na_2O_2 and filtered.			
	(a) A green filtrate and a brown residue			
		(b) A yellow filtrate and a green residue		
		(c) A yellow filtrate and a brown residue		
		(d) A colourless filtrate and a green residue		
	are obtained.			

- 53. During the separation of Group IV metal ions as insoluble carbonates by the addition of a saturated solution of $(NH_4)_2CO_3$ to the salt sample in an ammoniacal medium containing an excess of NH₄Cl, magnesium is not precipitated either as Mg(OH)₂ or MgCO₃ because
 - (a) the concentrations of OH⁻ and CO₃²⁻ are so low due to the common-ion effect that the ionic product values of Mg(OH)2 and MgCO₃ cannot exceed their respective solubility product values
 - (b) the solubility products of Mg(OH)₂ and MgCO₃ decrease due to the common-ion effect
 - (c) the solubility products of Mg(OH)₂ and MgCO₃ increase due to the common-ion effect
 - (d) all of these
- 54. Which of the following changes occur when a solution containing Mn²⁺ and Cr³⁺ is heated with an NaOH solution and H₂O₂?
 - (a) Mn(OH)₂ and Cr(OH)₃ precipitates which are formed initially dissolve due to the formation of Na₂MnO₄ and Na₂CrO₄.
 - (b) Soluble yellow Na₂CrO₄ and a brown precipitate of hydrated manganese dioxide are formed.
 - (c) A grey-blue gelatinous precipitate of Cr(OH)3 and a white precipitate of Mn(OH), are formed.
 - (d) Soluble Na₂MnO₄ and a grey-blue precipitate of Cr(OH)₃ are formed.
- 55. For the precipitation of Group IV metal ions as their insoluble carbonates, a saturated (NH₄)₂CO₃ solution is added to the salt sample in an ammoniacal medium, as a group reagent. But Na₂CO₃ or K₂CO₃ should not be used as the group reagent because
 - (a) they will prevent the precipitation of Group IV metal carbonates by forming soluble complex salts
 - (b) this would interfere with the Group V tests the presence or absence of Na + and K +
 - (c) they will prevent the precipitation of Group IV metal ions by forming soluble metal hydroxides
 - (d) they will precipitate out Mg 2+ ions of Group V (if present) along with Group IV metal carbonates
- **56.** When a solution of a salt in dilute HNO₃ is treated with an ammonium molybdate solution, a yellow precipitate is obtained, indicating the presence of

(a) NO_2^-

(c) PO₄³⁻

(b) SO₃²⁻
 (d) AsO₃³⁻

• *Type* 2 •

57. On treatment with dilute H₂SO₄, an inorganic salt produces a gas which gives a milky precipitate when passed through limewater. The salt may

(b) NaNO₃

(d) CH₃CO₂Na

Choose the correct options. More than one option is correct.

be

(a) Na₂CO₃

(c) Na_2SO_3

58.		n of the following radicals : bitated by passing H ₂ S through	0.30 M HCl solution will be	
	(a)	Cu ²⁺		Sb ³⁺
	(c)	Cd ²⁺	(d)	As^{3+}
59.		H_2S is passed through an pitate is obtained. The salt may		moniacal solution of a salt, a
	(a)	nickel salt	(b)	cobalt salt
	(c)	manganese salt	(d)	zinc salt
60.	Which	n of the following cations can b	e de	tected by the flame test?
	(a)	K^+	(b)	Ba ²⁺
	(c)	Sr ²⁺	(d)	Mg^{2+}
61.		ing heated with concentrated I on, a salt solution gives a yello		0_3 and an ammonium molybdate ecipitate. The salt may be
	(a)	Na_2HPO_4	(b)	As_2O_3
	(c)	FeSO ₄	(d)	BaCl ₂
62.		presence of which of the follow H_2SO_4 ?	wing	radicals can be detected using
	(a)	Chloride	(b)	Nitrate
	(c)	Sulphide	(d)	Sulphite
63.		eatment with H_2S gas, the solut precipitate. The salt may be a	ion c	of a salt in dilute HCl produces a
	(a)	copper salt	(b)	lead salt
	(c)	magnesium salt	(d)	nickel salt
64.		ammoniacal solution, a salt prosect through it. The salt may be		es a black precipitate when H ₂ S
	(a)	nickel salt	(b)	mercury salt
	(c)	cobalt salt	(d)	lead salt

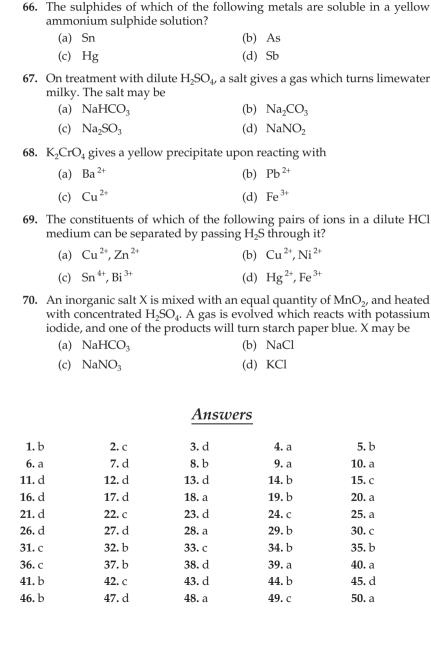
(b) KBrO₃

(d) $Na_2B_4O_7 \cdot 10H_2O$

65. Which of the following can be a primary standard?

(a) $Na_2S_2O_3 \cdot 5H_2O$

(c) $K_2Cr_2O_7$



51. b	52. c	53. a	54. b	55. b
56. c	57. a, c	58. a, b, c, d	59. a, b, c, d	60. a, b, c
61. a, b	62. c, d	63. a, b	64. a, c	65. b, c, d
66. a. b. d	67. a. b. c	68. a. b	69. a. b. d	70. b. d

Hints to More Difficult Problems

- 1. Polarization in the Ag + ion
- 2. Structural defect
- HgCl₂ is a covalent compound. There is little interaction between the Hg and Cl atoms in the HgCl₂ molecule.
- 9. SO₂ reduces the violet KMnO₄ to the almost colourless Mn²⁺.
- **12.** KMnO₄ itself is coloured.
- NaCl reacts positively to the chromyl chloride test whereas Na₂SO₄ does not.
- 26. Na₂S turns a dark purple-red upon reacting with Na₂[Fe(CN)₅NO].
- 40. A green chromic salt is formed.
- **44.** The insoluble BaSO₄ is formed.
- **49.** On treatment with dilute HCl, Na_2CO_3 and Na_2SO_3 produce CO_2 and SO_2 respectively. SO_2 turns an acidified $K_2Cr_2O_7$ solution green whereas CO_2 does not change the colour of the solution.
- 57. CO₂ as well as SO₂ turn limewater milky.
- 58. Cu²⁺ and Cd²⁺ belong to Group IIA. Sb³⁺ and As³⁺ belong to Group IIB.
- 67. CO₂ as well as SO₂ turn limewater milky.

18

Assertion-Reason Questions

The questions consist of an *assertion* in column 1 and a *reason* in column 2. Use the following key to choose the appropriate answer.

- (a) If the *assertion* as well as the *reason* are correct, and the *reason* is the correct explanation of the *assertion*.
- (b) If the *assertion* as well as the *reason* are correct, but the *reason* is not the correct explanation of the *assertion*.
- (c) If the assertion is correct but the reason is not.
- (d) If the reason is correct but the assertion is not.

Assertion Reason

- **1.** FeI₃ cannot exist in an aqueous solution.
- Fe $^{3+}$ oxidizes I $^-$ to I₂ easily.
- **2.** An aqueous solution of LiCl is slightly basic.
- Li⁺ polarizes H₂O molecules slightly.
- **3.** Graphite is chemically more reactive than diamond.
- Diamond is very hard but graphite is soft.
- **4.** AlF₃ is almost insoluble in anhydrous HF but dissolves when NaF is added to it.
- NaF produces free F⁻.
- 5. SF_6 exists but SH_6 does not.
- $d\pi$ -p π bonding cannot take place in SH₆
- **6.** Many peroxides are coloured.
- Diamagnetic superoxides are often found as impurities in these peroxides.
- 7. The stability of peroxides and superoxides increases on passing from Li to Cs
- The electropositive character of the elements in the periodic table increases on moving down a group.

8. The reaction $O + 2e \rightarrow O^{2-}$ is endothermic and yet a large number of compounds containing the oxide ion do exist.

This reaction is exergonic.

9. The compound (CF₃)₃N shows almost no basic behaviour even though (CH₃)₃N does.

There is no hydrogen bonding in $(CF_3)_3N$.

10. The bond in CIF is 5% shorter and that in BrF 7% shorter than the sum of the respective single-bond radii.

The ionic character is greater in BrF.

11. When CO₂ is passed through limewater, the solution turns milky but with an excess of CO₂, the solution becomes clear again.

Excess CO₂ changes the suspension to a colloidal solution

12. When tritium is produced commercially, helium is also produced.

The nuclear reaction to produce tritium is ${}^{6}\text{Li}_{3} + {}^{1}\text{n}_{0} \rightarrow {}^{3}\text{H}_{1} + {}^{4}\text{He}_{2}$

13. NF_3 is not a Lewis base.

The dipole moment of NF_3 is not zero.

14. SO₃ is a much stronger Lewis acid than CO₂.

The conjugate acid of SO_3 is much stronger than that of CO_2 .

15. $S_2O_8^{2-}$ is more oxidizing than SO_4^{2-} .

The oxidation number of sulphur in $S_2O_8^{2-}$ is more than that sulphur in SO_4^{2-} .

16. HF has low volatility.

bonding in HF.

hydrogen

17. BaSO₄ is less soluble than CaSO₄.

BaSO₄ is much heavier than CaSO₄.

18. NH₃ has a higher boiling point than HF.

The N atom is attached to three H atoms, and the molecule exhibits hydrogen bonding.

19. N_2H_4 cannot reduce $S_2O_3^{2-}$.

 $S_2O_3^{2-}$ is converted to $S_4O_6^{2-}$.

There is extensive

20. In liquid NH₃, Na and K are attracted towards the anode.

Na and K are converted to Na and K.

21. Borazole is aromatic in nature.

Nitrogen contributes π -electrons to the system.

22. The coordination number of N is seldom, if ever, 5 whereas that of P, which is in the same group, is 5 in a large number of compounds.

The first ionization energy of nitrogen is more than that of phosphorus.

23. The azide ion is thermodynamically unstable but easily exists under normal conditions.

The azide ion is kinetically inert.

24. Sodium is not produced commercially by the electrolysis of brine.

Sodium reacts violently with water.

25. Fluorine cannot be produced electrolytically.

Water is oxidized at a much higher potential than fluorine, and any fluorine produced will rapidly react with water.

26. NaIO₃ is converted to I₂ by using SO₂ rather than H_2O_2 or Sn²⁺.

SO₂ is very cheap.

27. Many metals produce coloured beads with borax.

Borax forms a glassy structure on heating, which optically influences light in a way that is characteristic of the metal involved.

28. Mercuric salts produce a red precipitate with an iodide ion solution but the precipitate dissolves in an excess of the iodide. In an excess of the iodide, colourless [HgIO₃]²⁻ is formed.

 Orthophosphoric acid is added to the Zimmermann–Reinhardt reagent during the dichrometric titration of ferrous salts. Orthophosphoric acid reduces the potential of the iron couple, thus aiding the oxidation of a ferrous salt.

30. A permanganate is intensely coloured.

The major contribution to the colour is by the d-d transition in manganese.

31. When a ferrous salt is added to a potassium hexacyano-ferrate (II) solution, a white precipitate is formed in the complete absence of air whereas a pale blue precipitate is formed under ordinary atmospheric conditions.

The white precipitate is due to the formation of K_2 Fe[Fe(CN)₆] and the pale blue precipitate due to the formation Fe[Fe(CN)₆].

32. The cyanide radical is a pseudohalogen.

The cyanide radical undergoes reactions similar to those of halogens

33. All solids tend to have defects.

Defects cause nonstoichiometry in solids.

- **34.** The ionization energy of Cu is more than that of K though both have a 4s ¹ configuration.
- **35.** Ketene and diazomethane are structurally similar.
- **36.** The bond energy of NH_3 is less than that of PH_3 .
- Crystals of NaHCO₃ and KHCO₃ show hydrogen bonding of different kinds.
- **38.** Graphite electrodes are not used in the electrolytic synthesis of fluorine.

The 18-electron shell of Cu shields nuclear charge more effectively than the shell of K.

They are isoelectronic.

Increasing size and decreasing electronegativity of the central atom permit the bonding electrons to be drawn out further, lowering the repulsion between the bonding pairs.

In NaHCO₃, the bicarbonate ions are linked in an infinite chain while in KHCO₃, a dimeric anion is formed.

Fluorine forms a highly explosive gunpowder-like mixture with graphite, which can explode even in darkness.

Answers	j
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1. a	2. d	3. b	4. a	5. a
6. c	7. a	8. c	9. b	10. a
11. c	12. a	13. b	14. b	15. c
16. a	17. c	18. d	19. a	20. a
21. b	22. b	23. a	24. b	25. c
26. a	27. c	28. c	29. a	30. c
31. a	32. a	33. b	34. c	35. b
36. a	37. a	38. c		

19

Matching-Type Questions (Chapterwise)

Periodic Table

1.	'Bismuth pentachloride'	does not
	exist due to	

2.
$$[Xe]4f^15d^16s^2$$

3.
$$[Xe]4f^75d^16s^2$$

- 8. Disproportionation
- 9. Diagonal pairs
- **10.** Ground-state electronic configuration is $(n-1)d^6ns^2$

(a) Promethium

(b)
$$PbI_4 \rightarrow PbI_2 + I_2$$

(c)
$$2\text{FeI}_3 \rightarrow 2\text{FeI}_2 + \text{I}_2$$

- (j) Close to the radius of the lutetium trivalent ion
- (k) Chromium
- (l) Manganese
- (m) Zinc
- (n) B and Si

Hydrogen and Oxygen

1. H₂ adsorber

- (a) H_2O and NH_2^-
- 2. Repeated electrolysis of alkaline water
- (b) Absorption of red light

- 3. Oxide that gives $\rm H_2O_2$ on reacting $\;\;$ (c) A moderator as well as a coolant with HCl
- 4. Superoxide ions5. Compound that is deuterated
- (d) sp 2 Hybridization with σ bonding
- **5.** Compound that is deuterated easily
- (e) Tritium

6. Identical geometry

(f) Na₂[Na₄(PO₃)₆]

7. D₂O

(g) Ca(OH)₂

8. O₃ is slightly bluish

- (h) Presence of superoxide ions
- **9.** Water is softened by treating water with
- (i) CH₃CH₂OH
- **10.** The central elements in ozone and SO₂
- (j) Stronger oxidizing agents than peroxide ions

- 11. $\Delta_{\text{fus}}H$ is maximum for
- (k) BaO_2
- 12. Many peroxides are coloured
- (l) D_2O
- (m) Pd
- (n) MnO₂
- (o) PbO₂
- (p) CH₃COCH₃
- (q) NH_3 and H_3O^+
- (r) In the presence of N₂
- (s) Absorption of violet light

Alkali Metals and Alkaline-Earth Metals—I

ConversionName of process1. NaCl \rightarrow NaOH(a) Down process2. NaCl \rightarrow Na $_2$ CO $_3$ (b) Spring reaction3. S \rightarrow Na $_2$ S $_2$ O $_3$ (c) Polymerization4. NaPO $_3$ \rightarrow Graham's salt(d) Solvay process5. NaCl \rightarrow Na(e) Castner–Kellner process

Alkali Metals and Alkaline-Earth Metals—II

1.	An amphoteric hydroxide	(a) Mg(OH)Cl
2.	The most basic hydroxide	(b) Mg and Cl ₂
3.	Setting of cement	(c) Chlorophyll
4.	Electrolysis of carnallite produces	(d) Decrease in the hydration energy of the metal ions down the group
5.	Estimation of Cl ₂ available in bleaching power	(e) Decrease in the lattice energy of the compounds more rapidly than that in the hydration energy
6.	Insoluble salt having an 8:4 coordination	(f) Hydration and gel formation
7.	Biomolecule containing Mg	(g) Ra(OH) ₂
8.	MgCl ₂ ·6H ₂ O on strong heating produces	(h) Iodometry
9.	Decrease in the solubility of carbonates of group 2 elements	(i) Be(OH) ₂

10. Increase in the solubility of fluorides and hydroxides of group 2 elements down the group

down the group

- (j) CaF₂
- (k) ATP
 - (l) Mg(OH)HCl·4H₂O
- (m) Haemoglobin
- (n) Iodimetry
- (o) Mg, K, Cl₂

Boron

- 1. Boron is used in the preparation of (a) Hydrolysis
- 2. Boron with NaOH produces

 (b) The terminal B—H bond is a 2-centre 3-electron bond containing 12 valence electrons
- 3. Orthoboric acid (c) Ammonolysis
- 4. Reaction of borax with water gives (d) Glass
- 5. $B_2H_6 \rightarrow H_3BO_3$ (e) $B(OH)_3$ and $B(OH)_4^-$

- **6.** $B_2H_6 \rightarrow borazine$
- 7. B₂H₆
- **8.** $B(OH)_3$

- (f) Hydrogen-bonded together to form 2D sheets with hexagonal symmetry
- (g) Na₃BO₃ and H₂
- (h) Triangular BO₃³⁻ units
- (i) Polymer
- (j) Na₂B₄O₇ and H₂
- (k) Pyramidal BO₃³⁻ units
- (l) The terminal B—H bond is a 2-centre 3-electron bond containing 16 valence electrons
- (m) Hydrogen-bonded together to form 3D sheets with hexagonal symmetry

Carbon

- 1. $C(graphite) \rightarrow C(diamond)$
- 2. $Be_2C \rightarrow CH_4$
- 3. Magnesium carbide \rightarrow propene
- 4. $K_4[Fe(CN)_6] \rightarrow CO$
- 5. Malonic acid \rightarrow C₃O₂
- 6. $CO \rightarrow HCOONa$
- 7. Heated CaC₂ and N₂ are required in the preparation of
- 8. Diamond-type lattice
- 9. $(SiH_3)_3N$ is planar
- 10. $(CH_3)_3N$ is pyramidal

- (a) Calcium nitride
- (b) Dehydration
- (c) α -Sn
- (d) $p\pi$ – $d\pi$ overlap between N and Si
- (e) No π bonding
- (f) Hydrolysis
- (g) Addition of concentrated H₂SO₄
- (h) High temperature and high pressure
- (i) Polymerization
- (j) Alkali fusion
- (k) Calcium cyanamide
- (l) Germanium
- (m) pπ–dπ overlap between N—N and Si—Si
- (n) Neither σ nor π bonding

Silicon

- **1.** A coordinately saturated compound/ion
- **2.** A coordinately unsaturated compound/ion
- 3. No donor property
- 4. $(SiH_3)_3N$
- 5. (CH₃)₃N
- 6. Silicones
- 7. $(CH_3)_3SiCl \rightarrow$ hexamethyldisiloxane
- 8. SiCl₄ + CH₃MgCl
- 9. $SiF_4 + HF$
- 10. A cyclic silicate

- (a) Si-O-Si linkages
- (b) Ca₃[Si₃O₉]
- (c) Hydrolysis and condensation
- (d) SiF_6^{2-}
- (e) CCl₄
- (f) Both sp³ and sp² orbitals
- (g) (SiH₃)₃N
- (h) Decomposition
- (i) sp² Orbitals only
- (j) Si-Si linkage
- (k) Si—C bond formation
- (l) SnF_5^-
- (m) Be₃Al₂[Si₆O₁₈]
- (n) sp ³ Orbitals only

Nitrogen

- 1. $HNO_3 \rightarrow N_2O_5$
- 2. $N_2O_5 \rightarrow NO + NO_2 + O_2$
- 3. $Zn + OH^- + NO_3^- \rightarrow NH_3$
- **4.** Detection of NO₃⁻ ions in the presence of NO₂⁻ ions
- $5. \quad NH_3 \rightarrow N_2H_4$
- **6.** $CaCN_2 \rightarrow urea$
- 7. $NH_3 \rightarrow N_2$
- 8. $NH_2OH \xrightarrow{H^+} N_2O + NH_4^+$
- 9. $NCl_3 \rightarrow NH_4OH + HOCl$

- (a) Hydrolysis
- (b) Raschig process
- (c) Oxidation by NaOCl
- (d) High pressure
- (e) Dehydration
- (f) Nitration followed by ring test
- (g) Decomposition
- (h) Decomposition by urea
- (i) Reduction

10. $NH_3 + CO_2 \rightarrow urea$

- (j) Decomposition by urea followed by ring test
- (k) Disproportionation
- (l) Addition reaction

Phosphorus

- 1. A reducing agent as well as a diprotic acid
- 2. A reducing agent as well as a monoprotic acid
- 3. A cyclic phosphate
- 4. $Mg(NH_4)PO_4 \rightarrow Mg_2P_2O_7$
- 5. $P_4 \rightarrow H_3PO_4$
- **6.** $Ca_3(PO_4)_2 \cdot CaF_2 + H_2SO_4$
- 7. $Ca_3(PO_4)_2 \cdot CaF_2 + H_3PO_4$
- 8. PCl₅

- (a) Superphosphate
 - (b) Triple superphosphate
 - (c) Ionic in the solid state
- (d) H_3PO_3
- (e) $H_6P_4O_{13}$
- (f) CuSO₄
- (g) H₃PO₂
- (h) Ignition
- (i) AgNO₃
- (j) Covalent in the liquid state

Sulphur

- 1. $Fe_2(SO_4)_3.9H_2O \rightarrow$ $Fe_2O_3 + SO_3 + H_2O$
- **2.** $C_6H_{12}O_6 \to C$
- 3. $SO_3 \rightarrow S_3O_0$
- 4. $H_2S_2O_8 \rightarrow H_2SO_4 + H_2SO_5$
- 5. $FeSO_4 \cdot 7H_2O \rightarrow Fe_2O_3 + SO_2 + SO_3$ (e) Dehydration
- 6. $KMnO_4 + H_2SO_4 + H_2S \rightarrow$ $K_2SO_4 + MnSO_4 + S + H_2O$

- (a) Oxidation of the reducing agent
- (b) Hydrolysis
- (c) Mild heating
- (d) Strong heating
- (f) Cyclization

The Halogens—I

- 1. $ClO^{-}(aq) \rightarrow ClO_{3}^{-}(aq) + Cl^{-}(aq)$
- (a) N_3^-

2. $I_2 + KI$

(b) Used in photography

3. AgBr/hydroquinone

(c) $(CN)_2$

4. A pseudohalide ion

- (d) BrF₃
- **5.** Not an interpseudohalogen compound
- (e) The product has a linear structure

6. A pseudohalogen

- (f) Disproportionation
- (g) ICl_2^-

The Halogens—II

Compound

Usage

1. CCl₂F₂

(a) Insulating medium for high-voltage transformers

2. $(C_2F_4)_n$

(b) Insecticide

3. SF₆

(c) Anaesthetic

4. ClF₃

(d) Insulating material in cables

5. DDT

(e) Refrigerator

The Halogens—III

Molecule/ion

Stucture

1. ClF₃

(a) Linear

2. BrF₅

(b) Bent

3. IF₇

(c) Square pyramidal

4. ICl₂

(d) T-shaped

5. OF₂

(e) Pentagonal bipyramidal

The Noble Gases

Hydrolysis of XeF₄

(a) $XeF_4 + PtF_4$

2. Hydrolysis of XeF₆

(b) Clathrate

- 3. Host molecule + guest molecule
- 4. $Xe + PtF_6 + heat$
- 5. Noble gases are difficult to liquefy
- **6.** XeF₆
- 7. XeO_3F_2

- (c) Capped octahedral
- (d) $XeO_3 + O_2 + HF$
- (e) $Xe[PtF_6]$
- (f) Small dispersion forces
- (g) Positive Joule–Thomson coefficients above 40 K
- (h) Trigonal bipyramidal structure with a lone pair of electrons
- (i) XeO₃

Copper

- 1. $2Cu^{2+} + SO_3^{2-} + 2SCN^- + H_2O \rightarrow$ $2CuSCN + H_2SO_4$
- 2. $2Cu^{2+} + 4I^{-} \rightarrow 2CuI + I_2$ and $I_2 + 2S_2O_3^{2-} \rightarrow 2I^{-} + S_4O_6^{2-}$
- 3. High melting point of copper
- **4.** The first IP of Cu is higher than that of K
- 5. Reactivity of Cu
- **6.** $CuSO_4 \cdot 5H_2O$ is blue in colour
- **7.** Most of the copper(I) complexes are colourless
- 8. $CuSO_4 + Na_2S_2O_3$ (excess)
- 9. CuSO₄ + KCN (excess)
- **10.** $CuSO_4 + K_4[Fe(CN)_6]$

- (a) High enthalpy of sublimation and high ionization energy
- (b) d Electrons in copper are poorly shielded
- (c) It absorbs approximately at 550 nm in the yellow region
- (d) Cu in the +II oxidation state has an unpaired electron
- (e) $Cu_2[Fe(CN)_6]$
- (f) Low enthalpy of sublimation and low ionization energy
- (g) Cu⁺ ions have a d¹⁰ configuration
- (h) d Electrons take part in metallic bonding
- (i) Volumetric analysis of Cu
- (j) Gravimetric analysis of Cu
- (k) $Na_4[Cu_6(S_2O_3)_5]$
- (l) $K_3[Cu(CN)_4]$
- (m) $K_2[Cu(CN)_4]$
- (n) $Cu_3[Fe(CN)_5]$

Silver and Gold

- 1. AgBr is used in photography
- 2. Na₂S₂O₃ is used in photography
- 3. AuCl₃ on reacting with NH₃ forms
- 4. Cupellation process
- 5. Radius of $Ag \approx radius of Au$
- Extraction of Ag involving KCN, O₂ and active metal

- (a) Refining of Ag
- (b) Lanthanide contraction
- (c) Culminating gold
- (d) Pattinson process
- (e) Fulminating gold
- (f) As a light-sensitive material
- (g) As a fixer
- (h) As a developer
- (i) As a reducing agent
- (j) Gold sol
- (k) The ccp structures of both Ag and Au
- (l) McArthur-Forrest process
- (m) Parkes process

<u>Iron</u>

- 1. Rusting of iron
- 2. $FeSO_4 + (NH_4)_2SO_4 + H_2SO_4$
- 3. $FeCl_3 + K_4[Fe(CN)_6]$
- 4. $FeCl_2 + K_3[Fe(CN)_6]$
- 5. $FeSO_4 + AuCl_3$
- **6.** $Fe_2(SO_4)_3 + heat$
- Steel is heated at high temperature and then cooled suddenly
- 8. Steel is heated at high temperature and then cooled slowly

- (a) SO₃ only
- (b) Annealing
- (c) An electrochemical process
- (d) Quenching
- (e) Mohr salt
- (f) Prussian blue
- (g) Colloidal gold
- (h) Turnbull's blue
- (i) Fulminating gold
- (j) Ferric ammonium alum

- (k) An electrolytic process
- (l) A mixture of SO₂ and SO₃

Coordination Chemistry

- 1. Ni(CO)₄
- 2. Ni(PPh₃)₂Cl₂
- 3. $[Ni(NH_3)_4]^{2+}$
- 4. $[Zn(NH_3)_4]^{2+}$
- 5. Geometrical isomerism
- **6.** $K^{+}[Pt(C_{2}H_{4})Cl_{3}]^{-}\cdot H_{2}O$
- 7. cis-trans Isomerism
- 8. EAN for $[Ni(CN)_4^{2-}]$
- **9.** EAN for $[Pd(NH_3)_6]^{4+}$
- 10. A low-spin (spin-paired) complex
- 11. A high-spin (spin-free) complex
- **12.** A complex ion having a magnetic moment of 1.73 Bohr magnetons
- **13.** A complex ion having a magnetic moment of 2.83 Bohr magnetons
- **14.** [Ag(NH₃)₂]Cl

- (a) $[V(H_2O)_6]^{3+}$
- (b) 34
- (c) Ziese's salt
- (d) $[Co(NH_3)_4H_2O]Cl$
- (e) 54
- (f) $[Co(NH_3)_6]^{3+}$
- (g) $[Mn(CN)_6]^{4-}$
- (h) Optical as well as *cis-trans* isomerism
- (i) Ma₄b₂
- (j) A diamagnetic species
- (k) Square planar
- (l) CoF₆³⁻
- (m) Tetrahedral
- (n) A paramagnetic species
- (o) 36
- (f) 56

Analytical Chemistry—I

- Gas that turns moist lead acetate paper black
- **2.** Salt + conc. $H_2SO_4 + K_2Cr_2O_7$
- 3. Salt + H_2SO_4 + Mohr salt
- 4. Salt + Pb_3O_4 + conc. HNO_3
- 5. Salt + DMG + NH_3

- (a) Ni ²⁺
- (b) Devarda's alloy and NaOH
- (c) HCl and BaCl₂
- (d) Sodium nitroprusside
 - (e) Hydrogen sulphide

- **6.** NO₃ and Br can be distinguished from each other by
- (f) Chromyl chloride test
- 7. Na₂SO₃ and Na₂SO₄ can be distinguished from each other by
- (g) Chloride
- 8. Na₂SO₃ and Na₂S can be distinguished from each other by
- (h) Decomposition by urea followed by ring test
- 9. NaCl and Na₂SO₄ can be distinguished from each other by
- (i) Nitrate
- NaNO₂ and NaNO₃ can be distinguished from each other by
- $(j) Mn^{2+}$
- (k) Co²⁺
- (l) Zn, Co, Ni
- (m) Mg, Sn, Al

Analytical Chemistry—II

	Flame coloration	Metal
1.	Golden yellow	(a) Potassium
2.	Lilac	(b) Strontium
3.	Brick red	(c) Sodium
4.	Crimson	(d) Barium
5.	Yellowish green	(e) Calcium

Analytical Chemistry—III

	Colour of borax b	Metal	
	Hot	Cold	
1.	Green	Blue	(a) Manganese
2.	$Yellowish\ brown$	Yellow	(b) Chromium
3.	Yellow	Green	(c) Cobalt
4.	Violet	Amethyst	(d) Iron
5.	Blue	Blue	(e) Nickel
6.	Violet	Reddish brown	(f) Copper

Aluminium

- 1. By-product of the Serpek process
- 2. Alums help in purifying water
- 3. Anodized Al
- 4. Lapis lazuli, a blue rock
- 5. Removing the self-protective oxide film from Al
- 6. Ultramarine

- (a) Sodium aluminium silicate and sodium pyrosulphide
- (b) Sodium aluminium silicate
- (c) Amalgamation
- (d) Al electrically coated with Al₂O₃
- (e) Coagulation of mud
- (f) NH₃
- (g) CO₂
- (h) Flocculation of mud
- (i) Zinc cobaltate
- (j) Aluminium electrically coated with lead(IV) oxide

Tin and Lead

Reaction

- 1. $Pb \rightarrow Pb(CH_3COO)_2 \cdot 3H_2O$
- 2. $PbCl_2 \rightarrow PbCrO_4$
- 3. $Pb(NO_3)_2 \rightarrow PbCrO_4 \cdot PbO$
- **4.** $Pb(CH_3COO)_2 \rightarrow 2PbCO_3 \cdot Pb(OH)_2$ (d) Black oxide of lead
- 5. $Pb_3O_4 \rightarrow PbO$
- 6. PbO \rightarrow Pb₃O₄
- 7. $SnCl_2 \rightarrow SnO_2$
- 8. $Pb(NO_3)_2 \rightarrow PbO + Pb$
- 9. $SnCl_4 + HCl \rightarrow H_2[SnCl_6]$
- 10. Sn + Pb \rightarrow alloy

Product

- (a) White lead
- (b) Common solder
- (c) Minium
- (e) Cassiterite
- (f) Inorganic sugar
- (g) Chrome yellow
- (h) Chrome red
- (i) Litharge
- (j) A coordinately unsaturated compound
- (k) Massicot
- (m) Red lead
 - (o) A coordinately saturated compound

Zinc and Mercury

- 1. Galvanized iron pipe
- 2. Fe + Hg \rightarrow Fe_xHg_y
- 3. $ZnSO_4 + CN^- \rightarrow Zn(CN)_4^{2-} + SO_4^{2-}$
- 4. Complex halides of Hg^{II}
- $5. Zn \rightarrow Zn_2P_2O_7$
- 6. $ZnSO_4 + BaS$

- (a) Gravimetric estimation
- (b) Formation of a pigment known as lithopone
- (c) Amalgamation
- (d) Iron pipe dipped in molten nickel
- (e) Complex ion having tetrahedral structure
- (f) Nessler's reagent used to detect NH_3
- (g) No amalgam formation
- (h) Iron pipe dipped in molten zinc

Transition Elements and Lanthanoids

- **1.** A pair of metals having the same density
- 2. Wilkinson's catalyst
- 3. Adam's catalyst
- 4. A Ziegler–Natta catalyst
- 5. Fenton's reagent
- 6. Baeyer's reagent
- 7. $MnO_2 + KNO_2$
- 8. $CrO_2Cl_2 + NaOH$
- 9. $(NH_4)_2Cr_2O_7 + heat$
- 10. $Na_2Cr_2O_7 + NaOH$
- 11. Occurrence of Ce 4+
- **12.** The most common oxidation state of rare-earth elements
- 13. A radioactive rare-earth element
- **14.** An oxidizing agent used both in volumetric analysis and in preparative chemistry

- (a) Cerium(IV) solution
- (b) Lu
- (c) +III
- (d) K_2MnO_4
- (e) Na₂CrO₄
- (f) 4f¹
- (g) Pt/PtO
- (h) Rh(Cl)(PhP)₃
 - (i) $FeSO_4/H_2O_2$
 - (j) $TiCl_4/(C_2H_5)_3Al$
 - (k) KMnO₄/OH⁻
 - (l) Cr₂O₃
- (m) Ni and Co
- (n) Ag and Au
- (o) Na₂Cr₂O₇
- (p) 4f⁰
- (q) + IV
- (r) Pm

Answers

Periodic Table

1.	4	\rightarrow	f		

 $\mathbf{2.} \leftrightarrow \mathbf{h}$ $\mathbf{7.} \leftrightarrow \mathbf{k}$

 $3. \leftrightarrow i$ $8. \leftrightarrow b, c$ $4. \leftrightarrow g$ $9. \leftrightarrow d, n$ $5. \leftrightarrow j$ $10. \leftrightarrow f$

Hydrogen and Oxygen

$$1. \leftrightarrow m$$

 $6. \leftrightarrow a$

$$2. \leftrightarrow 1$$

 $3. \leftrightarrow k$

 $4. \leftrightarrow j$

 $\textbf{5.} \leftrightarrow i$

6.
$$\leftrightarrow$$
 a, q

8. ↔ b

9. \leftrightarrow f

10. ↔ d

11.
$$\leftrightarrow$$
 e

12.
$$\leftrightarrow$$
 h

Alkali Metals and Alkaline-Earth Metals—I

$$1. \leftrightarrow e$$

$$\mathbf{2.} \leftrightarrow \mathbf{d}$$

$$\textbf{3.} \leftrightarrow b$$

$$4. \leftrightarrow c$$

$$5. \leftrightarrow a$$

Alkali Metals and Alkaline-Earth Metals—II

$$\mathbf{1.} \leftrightarrow \mathbf{i}$$
$$\mathbf{6.} \leftrightarrow \mathbf{j}$$

$$\mathbf{2.} \leftrightarrow \mathbf{g}$$
$$\mathbf{7.} \leftrightarrow \mathbf{c}$$

$$3. \leftrightarrow f$$

 $8. \leftrightarrow a$

$$4. \leftrightarrow b$$

 $9. \leftrightarrow d$

$$5. \leftrightarrow h$$

 $10. \leftrightarrow e$

Boron

1.
$$\leftrightarrow$$
 d
6. \leftrightarrow c

2.
$$\leftrightarrow$$
 g, j
7. \leftrightarrow b

3.
$$\leftrightarrow$$
 h, k
8. \leftrightarrow f

Carhon

$$\textbf{4.} \leftrightarrow g$$

$$\textbf{5.} \leftrightarrow b$$

1.
$$\leftrightarrow$$
 h
6. \leftrightarrow j

$$\mathbf{2.} \leftrightarrow f$$
$$\mathbf{7.} \leftrightarrow k$$

$$3. \leftrightarrow f$$

 $8. \leftrightarrow c, 1$

$$\mathbf{9.} \leftrightarrow \mathbf{d}$$

10. ↔ e

Silicon

$$1. \leftrightarrow e$$

$$6. \leftrightarrow a$$

$$2. \leftrightarrow 1$$

 $7. \leftrightarrow c$

$$3. \leftrightarrow g$$

 $8. \leftrightarrow k$

$$\mathbf{4.} \leftrightarrow \mathbf{i}$$
 $\mathbf{9.} \leftrightarrow \mathbf{d}$

$$5. \leftrightarrow n$$

 $10. \leftrightarrow b, m$

$$2. \leftrightarrow g$$
$$7. \leftrightarrow c$$

$$3. \leftrightarrow i$$

 $8. \leftrightarrow k$

$$4. \leftrightarrow j$$

 $9. \leftrightarrow a$

$$5. \leftrightarrow b$$

$$10. \leftrightarrow d$$

6. \leftrightarrow a **Phosphorus**

$$\mathbf{1.} \leftrightarrow \mathbf{d}$$
$$\mathbf{6.} \leftrightarrow \mathbf{a}$$

$$2. \leftrightarrow g$$

 $7. \leftrightarrow b$

$$3. \leftrightarrow e$$

 $8. \leftrightarrow c, j$

$$4. \leftrightarrow h$$

$$5. \leftrightarrow f, i$$

Sulphur

1.
$$\leftrightarrow$$
 d 2. \leftrightarrow e 6. \leftrightarrow a

$$3. \leftrightarrow f$$

$$4. \leftrightarrow b$$

$$\textbf{5.} \leftrightarrow d$$

The Halogens—I

$$\mathbf{2.} \leftrightarrow \mathbf{e}$$

$$3. \leftrightarrow b$$

$$4. \leftrightarrow a, g$$

$$5. \leftrightarrow d$$

 $6. \leftrightarrow c$

The Halogens—II

$$1. \leftrightarrow e$$

$$\mathbf{2.} \leftrightarrow \mathbf{d}$$

$$3. \leftrightarrow a$$

$$4. \leftrightarrow c$$

$$5. \leftrightarrow b$$

The Halogens—III

$$1. \leftrightarrow d$$

$$2. \leftrightarrow c$$

$$3. \leftrightarrow e$$

$$4. \leftrightarrow a$$

$$5. \leftrightarrow b$$

The Noble Gases

1.
$$\leftrightarrow$$
 d
6. \leftrightarrow b

$$2. \leftrightarrow i$$
 $7. \leftrightarrow h$

$$3. \leftrightarrow b$$

$$4. \leftrightarrow e$$

$$5. \leftrightarrow f, g$$

Copper

$$\mathbf{2.} \leftrightarrow \mathrm{i}$$

$$3. \leftrightarrow h$$

$$4. \leftrightarrow b$$

$$5. \leftrightarrow a$$

6.
$$\leftrightarrow$$
 c, d

$$7. \leftrightarrow g$$

$$8. \leftrightarrow k$$

$$9. \leftrightarrow 1$$

$$\textbf{10.} \leftrightarrow e$$

Silver and Gold

$$\mathbf{1.} \leftrightarrow \mathbf{f}$$

$$\mathbf{6.} \leftrightarrow \mathbf{l}$$

$$\mathbf{2.}\leftrightarrow\mathbf{g}$$

$$3. \leftrightarrow e$$

$$\textbf{4.} \leftrightarrow a$$

$$5. \leftrightarrow b$$

Iron

1.
$$\leftrightarrow$$
 c
6. \leftrightarrow a

$$\mathbf{2.} \leftrightarrow \mathbf{e}$$

$$\mathbf{7.} \leftrightarrow \mathbf{d}$$

$$3. \leftrightarrow f$$

 $8. \leftrightarrow b$

$$\textbf{4.} \leftrightarrow h$$

$$5. \leftrightarrow g$$

Coordination Chemistry

$$\textbf{1.} \leftrightarrow m$$

$$2. \leftrightarrow k$$

$$\textbf{3.} \leftrightarrow n$$

$$4. \leftrightarrow j$$

$$5. \leftrightarrow i$$

 $10. \leftrightarrow f$

$$\mathbf{6.} \leftrightarrow \mathbf{c}$$

$$\mathbf{11.} \leftrightarrow \mathbf{l}$$

$$7. \leftrightarrow d$$

$$12. \leftrightarrow g$$

$$8. \leftrightarrow b$$

$$13. \leftrightarrow a$$

$$9. \leftrightarrow e$$

14. \leftrightarrow h

Analytical Chemistry—I

$$\mathbf{1.} \leftrightarrow \mathbf{e}$$

$$2. \leftrightarrow g$$

$$3. \leftrightarrow i$$

$$4. \leftrightarrow j$$

$$\mathbf{5.} \leftrightarrow \mathbf{a}$$

$$6. \leftrightarrow b$$

$$7. \leftrightarrow c$$

$$8. \leftrightarrow d$$

$$9. \leftrightarrow f$$

10.
$$\leftrightarrow$$
 h

Analytical Chemistry—II

$$1. \leftrightarrow c$$

$$\textbf{2.} \leftrightarrow a$$

$$3. \leftrightarrow e$$

$$4. \leftrightarrow b$$

$$\textbf{5.} \leftrightarrow d$$

Analytical Chemistry—III

$$\mathbf{1.} \leftrightarrow \mathbf{f}$$
$$\mathbf{6.} \leftrightarrow \mathbf{e}$$

$$\mathbf{2.} \leftrightarrow \mathbf{d}$$

$$3. \leftrightarrow b$$

$$\mathbf{4.} \leftrightarrow \mathbf{a}$$

$$5. \leftrightarrow c$$

Aluminium

- 1. \leftrightarrow f 2. \leftrightarrow e 3. \leftrightarrow d 4. \leftrightarrow b 5. \leftrightarrow c 6. \leftrightarrow a
- Tin and Lead
- 1. \leftrightarrow f2. \leftrightarrow g3. \leftrightarrow h4. \leftrightarrow a5. \leftrightarrow i, k6. \leftrightarrow c, m7. \leftrightarrow e8. \leftrightarrow d9. \leftrightarrow o10. \leftrightarrow b
- Zinc and Mercury
 - 1. \leftrightarrow h 2. \leftrightarrow c 3. \leftrightarrow e 4. \leftrightarrow f 5. \leftrightarrow a 6. \leftrightarrow b
- Transition Elements and Lanthanoids

Matrix-Matching-Type Questions (Mixed)

Matrix A

1. Industrial synthesis of methanol

(a) CO

2. Water gas shifting reaction at 500°C

(b) PbI₄

3. Ostwald process

(c) O_2

4. Contact process

(d) H_2

Matrix B

1. $KI + O_3 + H_2O$

(a) Acid-base and/or redox titration

2. $KH_3(C_2O_4)_2 \cdot 2H_2O$

(b) Phenolphthalein and methyl orange indicators

3. Na₂CO₃ + HCl

(c) Theory of adsorption indicator

4. $AgNO_3 + NaCl$

(d) K₂CrO₄ solution

Matrix C

Reaction

1. $CH_4 \rightarrow CO_2$

2. $PCl_5 \rightarrow H_3PO_4$

3. $HNO_3 \rightarrow N_2O_5$

4. $Ni(CO)_4 \rightarrow Ni(CN)^{2-}$

Change in hybridization of central atom

(a) $sp^3 \rightarrow dsp^2$

(b) $sp^3 \rightarrow sp$

(c) $sp^3d \rightarrow sp^3$

(d) $sp^3 \rightarrow sp, sp^2$

2-202

Matrix D

- 1. Self-reduction process (a) Purification of titanium
- 2. Carbon monoxide process (b) Copper
- 3. Decomposition of iodide (c) Iron
- 4. Face-centred-cubic lattice (d) Pyrometallurgy

Matrix E

- 1. H₂S (a) Weak dipole–dipole forces
- 2. CH₃OH (b) London dispersion forces
- 3. C_2H_6 (c) Hydrogen bonding
- 4. HNO₃ (d) Large dipole–dipole forces

Matrix F

- 1. ICl₄ (a) Tetrahedral
- 2. Al₂Cl₆ (b) Octahedral
- 3. $Fe(CN)_6^{4-}$ (c) Square planar
- 4. I_2Cl_6 (d) d^2sp^3 Hybridization

Matrix G

- 1. Two unpaired electrons (a) $Ni(en)_3^{2+}$
- **2.** No unpaired electron (b) VF_6^{3-}
- 3. Magnetic moment = 2.83 Bohr (c) $Ni(CN)_4^2$ magnetons
- 4. Square-planar geometry (d) $Fe(CN)_6^{4-}$

Matrix H

- 1. Silicon (a) Semiconductor
- 2. The charge on the anion is equal to the number of terminal oxygen atoms

- **3.** Three shared corners and ten unshared corners
- (c) SiO₄⁴⁻
- 4. A silicon atom at the centre of the tetrahedron and an oxygen atom at each corner
- (d) $Si_4O_{13}^{10-}$

Matrix I

- 1. Orthophosphoric acid
- (a) Brønsted-Lowry acid

2. B(OH)₃

(b) Nonreducing triprotic acid

3. BF₃

(c) Weak Lewis acid having a hexagonal structure

4. Borazole

(d) Dehydrated product forming Graham's salt

Matrix J

1. P₄ molecule

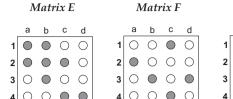
(a) One equilateral triangular face of a tetrahedral molecule showing 60° bond angles and a 90° angle between the p orbitals

2. Triphosphate ion

- (b) Strain would not be reduced by using sp³ hybrid orbitals
- Heating a stoichiometric mixture of the powdered orthophosphate salts Na₂HPO₄ and NaH₂PO₄
- (c) Eutrophication
- 4. $P_3O_{50}^{5-}$ contributes to excessive fertilization and rampant growth of algae
- (d) Synthetic detergent

Answers

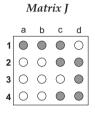
	N	latr	ix A	L		N	1atr	ix B	1		N	latr	ix C			M	latr	ix D)
	а	b	С	d		а	b	С	d		а	b	С	d		а	b	С	d
1		0	0		1		0		0	1	0		0	0	1	0	0	0	
2		\circ	\circ	\circ	2		\circ	\circ	0	2	\circ	\circ		\circ	2	\circ	\circ		
3	\circ				3			\circ	0	3	\circ	\circ	\circ		3		\circ	\circ	0
4	0	0	0	0	4	0	0	0		4	0	0	0	0	4	0	0	0	0



	N	latr	ix G		N	latr	ix H	Ī	
	а	b	С	d		а	b	С	d
1			\circ	\circ	1			0	0
2	0			\circ	2	\circ	\circ		
3		\circ		\circ	3	\circ	\circ	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	
4	0	0		0	4	0	0		

	а	b		a
1		0	\circ	
2	0	0		\circ
3	0	0		\circ
4	0	\circ		\circ

Matrix I



21

Comprehension-Type Questions

• An aqueous solution of a boron compound (A) is alkaline in the presence of phenolphthalein, and the aqueous solution of (A) in phenolphthalein can be decolorized by the addition of glycerol. When the solid compound (A) is strongly heated, two solid substances (B) and (C) are produced. When the solid product mixture of (B) and (C) is boiled with water and filtered, the filtrate contains only (C). Addition of H₂O₂ to (C) produces a peroxo compound (D). (B) is heated with a mixture of carbon and chlorine to give a volatile substance (E). (E) on heating with NH₄Cl at 140°C gives (F). (F) on reaction with NaBH₄ in ether produces (G). (G) on hydrolysis yields a gaseous mixture (H).

(b) NaBO₂

Choose the correct option. Only one option is correct.

	` '	2 0	` '	-
	(c)	$B(OH)_3$	(d)	$Na_2B_4O_7 \cdot 10H_2C$
2.	The co	ompounds (B) and (C) are res	spectiv	ely
	(a)	B(OH) ₃ and B ₂ O ₃	(b)	B ₂ O ₃ and NaBO
	(c)	NaBO ₂ and H ₃ BO ₃	(d)	B_2H_6 and B_2O_3
3.	The co	ompound (D) is used as		
	(a)	a brightener in the preparat	ion of	washing powde
	(b)	an antiseptic in medicine		

4. The compound (E) has a

1. The compound (A) is (a) B_2O_3

(a) tetrahedral structure

(d) a water softener

(b) plane triangular structure

(c) an analgesic in medicine

- (c) linear structure
- (d) trigonal bipyramidal structure

5.		ercentage of 6 27.63		rine in the mo 17.63		le of (F) is 19.75	(d)	15.43	
6.	The molecule of (G) is isoelectronic with (a) benzene, and the bond order is greater than 1.0 (b) cyclohexane, and the bond order is less than 1.5 (c) diborane, and the bond order is equal to 2.0 (d) benzene, and the bond order equals 1.0								
7.	(a)	aseous mixtu $\mathrm{NH_3}$ and $\mathrm{N_2}$ $\mathrm{NH_3}$ and $\mathrm{H_2}$	O	I) contains		$\mathrm{NH_3}$ and $\mathrm{O_2}$ $\mathrm{NH_3}$ and BF			
•	A black mineral (A) on heating in the presence of air gives a gas (B). The mineral (A) on reaction with dilute H_2SO_4 gives a gas (C) and the solution of a compound (D). On passing (C) into an aqueous solution of (B), white turbidity is obtained. The aqueous solution of (D) on reaction with $K_3[Fe(CN)_6]$ gives a blue compound (E).								
8.		nineral (A) is ZnS	(b)	FeS	(c)	FeS ₂	(d)	Fe_2O_3	
9.	_	as (B) obtaine SO ₂		SO ₃	(c)	H ₂ S	(d)	O_2	
10.	_	as (C) is CO_2	(b)	SO ₂	(c)	H_2S	(d)	N ₂ O	
11.	(a)	queous soluti FeSO ₄ FeCl ₂	on o	f (D) contains	(b)	FeSO ₄ ·(NH ₄ FeCl ₃) ₂ SO,	4	
12.	(a)	lue compoun K ₄ Fe ^{II} [Fe ^{III} (C K ₃ Fe ^{III} [Fe ^{II} (C	CN) ₆]		K ₂ Fe ^{II} [Fe ^{II} (C			
•	A green solid (A) on electrolytic oxidation in the alkaline medium gives a purple-coloured solution (B). The solution (B) oxidizes KI in the acidic medium and liberates (C), which reacts with chlorine water to give (D) and								

dic nd HCl. The potassium salt of (D) when heated gives O₂.

13. The green solid compound (A) showing sp³ hybridization is

(a) K_2MnO_4

(b) FeSO₄·7H₂O

(c) NiSO₄·6H₂O

(d) CuSO₄·5H₂O

14.	I. The purple colour of the solution (B) is due to							
		$Co(NO_3)_2$				$MnCl_2$		
	(c)	$KMnO_4$			(d)	$K_3[Co(NC)]$	$(2)_{6}$	
15.	The co	ompound (C)	libe	rated is				
	(a)	a mild redu	cing	agent				
		a strong red		~ ~				
	(c)	used in prep	oarin	g tincture of	iodir	ne in the sil	k indu	stry
	(d)	used to perf	orm	iodoform te	st			
16.		ompound (D)	_					
	(a)	HClO ₃	(b)	$HMnO_4$	(c)	HIO_3	(d)	KO_3
•	A com gentle	npound (A) is heating gives	boile (C) a	d with H ₂ SO and on strong	₄ to gi g heat	ive gypsum ing gives (I	and an)).	acid (B). (B) on
17.	The co	ompound (A)	is					
	(a)	-		a mineral	(c)	a rock	(d)	none of these
18.	The a	cid (B) is a						
	(a)	monoprotic	acid	having sp ³	hybri	dization		
	(b)	diprotic acid	d hav	ing sp ³ d hy	bridiz	zation		
	(c)	triprotic acid	d hav	ving sp ³ hyb	ridiza	ition		
	(d)	triprotic acid	d hav	ving dsp ² hy	bridiz	zation		
19.	The co	ompound (C)	sho	ws				
		sp ³ -sp ² over			(b)	sp-sp ³ ov	erlappi	ing
	(c)	sp ³ -sp overl	lappi	ing	(d)	sp^3-sp^3 or	verlapp	oing
20.	The co	ompound (C)	is					
	(a)	pyrophosph	oric	acid	(b)	orthopho	sphoric	acid
	(c)	hypophosph	noro	us acid	(d)	pyrophos	phorou	ıs acid
21.	The so	odium salt of	(D) i	is known as				
	(a)	Glauber's sa	alt		(b)	Graham's	salt	
	(c)	Rochelle sal	t		(d)	Ziese's sa	lt	
•	A white crystalline solid (A) decomposes on heating, leaving no residue. The gaseous substances evolved include water vapour and a gas (B), which rekindles a glowing splint. The solid (A) dissolves in water, yielding a colourless solution. The aqueous solution of (A) with a fresh FeSO ₄ solution							

he ch a n and concentrated H₂SO₄ produces a brown ring. The aqueous solution of (A) with alkali produces a gas (C), which responds positively to Nessler's reagent.

22. The white salt (A) shows

(a) sp^3-sp^2 overlapping

(b) sp³–sp overlapping

(c) sp³–sp³ overlapping

(d) sp³-dsp² overlapping

(a) bent structure with a high dipole moment(b) bent structure with a zero dipole moment(c) linear structure with a very low dipole moment(d) linear structure with a high dipole moment

23. The gas (B) has a

24.	. The gas (B) is neutral, having the molecular formula							
	(a)	CO_2	(b)	NO ₂	(c)	CO	(d)	N ₂ O
25.	(a) (b) (c)	JPAC name for pentaaquair pentaaquair pentaaquan pentaaquan	on(II on(I) itros	II) sulphate nitrosyl sulp oiron(II) sulp	ohate hate			
26.	In the	brown ring t	he o	xidation num	nber (of Fe is		
	(a)	_	(b)		(c)		(d)	+3
27.	(a)	rown colour i H ₂ NHgOHg NH ₂ Hg ₂ I ₃			(b)	due to the for $H_2NHg_2I_2$ $NH_2OHg_2I_3$	rmat	ion of
•	• A colourless crystalline solid (A) turns skin black. It gives two gases—(B) and (C)—and a residue (D) when (A) is heated to 950°C. One of the gases is soluble in water yielding an acidic solution (E). The residue (D) is soluble in (E). The solid (A) gives a brown precipitate with NaOH solution. The brown precipitate produces an Ag mirror with glucose. The aqueous solution of (A) gives a white precipitate, which redissolves in excess KCN solution, forming a stable complex (F). Addition of Zn to (F) gives (D).							
28.	(a)	hite solid (A) horn silver silver glance				argentite lunar causti	С	
29.	~	as (B) having dimerize				tendency to polymerize	(d)	trimerize
30.		olution (E) is a		-	-	ence of N ₃ H	(d)	HNO ₃
31.	_	as (B) has a bo 180°				ely equal to 115°	(d)	120°
32.	obtair (a)	as (C), which led by heating KClO ₃ Pb ₃ O ₄		aramagnetic	(b)	have a bond HgO all of these	orde	r equal to 2, is

33.	The brown precipitate is due to the formation of								
	(a)	Ag(OH)	(b)	Ag_2O	(c)	$AgClO_4$	(d)	$AgNO_2$	
34.	The st	tructure of the	e cor	nplex (F) is					
	(a)	sp^2	(b)	sp	(c)	sp^3	(d)	dsp ²	
35.	The ac	ddition of Zn	to (I	F) is known a	s				
	(a)	McArthur-I	Forre	st cyanide pr	oces	S			
	(b)	Pattinson pa	oces	s					
	(c)	Raschig pro	cess						
	(d)	none of thes	se						
	A white solid (A) is not completely soluble in dilute HCl/H_2SO_4 but dissolves readily in dilute HNO_3 , evolving an acidic gas (B). The gas (B) continues to burn with magnesium. (A) on heating gives a yellow solid (C) which is an amphoteric oxide. (C) when heated in air forms a red compound (D), which dissolves in concentrated HNO_3 , giving a brown residue (E).								
36.		ompound (A)	is k	nown as					
		anglesite			` /	cerrusite			
		litharge				minium			
37.		urning of Mg	witl	n (B) in the at	-	-			
		MgO only				Mg ₃ N ₂ only			
		Mg				MgO and M	lg_3N_2	2	
38.		ompound (A)	is u	sed to prepai					
	. ,	white lead	, ,		. ,	alloys			
		Zn by electr	-	S	(d)	all of these			
39.		ellow solid (C	C) is						
	` '	PbO ₂			. ,	Pb ₃ O ₄			
	()	PbO			` '	PbSO ₄			
40.		ed-coloured c	•						
	(a)	minium	(b)	litharge	(c)	fool's gold	(d)	vermilion	
41.		ompound (E)		sed to prepar	e				
	(a)	car batteries	;		(b)	alkaline bat	teries	3	
	(c)	lithium cells	3		(d)	Weston cad	miur	n cells	
•	molec		(C),	and sodium o				a paramagnetion (B) reacts with	
42.	The co	ompound (A)	imp	arts a flame	color	ation of			
	(a)	violet			(b)	brick red			
	(c)	crimson			(d)	golden yello	w		

43.	3. The compound (A) on heating produces							
	(a)	Cl_2			(b)	O_2		
	(c)	a metal oxid	le		(d)	ClO_2		
44.	The p	aramagnetic:	mole	cule (B) ha	s			
	(a)	no tendency	to d	imerize	(b)	a tendency	to dir	merize
	(c)	a bent struct	ture		(d)	a linear str	ucture	9
45.	The g	as (D) may be	e pro	duced by				
	(a)	the electroly	sis o	f acidulated	d wate	r		
	(b)	the electroly	sis o	f an aqueo	us Na ₂ s	SO ₄ solution	ı	
	(c)	the electroly	sis o	f an aqueo	us KMı	nO ₄ solution	ı	
	(d)	all of these						
•	A colourless crystalline compound (A) loses the water of crystallization on heating. (A) reacts with dilute HCl, giving a pungent-smelling gas (B) and a yellow colloidal particle (C). The gas (B) turns an acidified dichromate solution into a green solution (D). (A) decolorizes an I ₂ solution in KI to give (E). (A) gives a white precipitate (F) with AgNO ₃ and turns black on standing.							
46.	The o	xidation num			ral eler	ment in the o	compo	ound (A) are
	(a)	+2, 0, -1	(b)	+4, -2	(c)	+4, +2	(d)	+2, -2
47.	The ce	entral elemen	t in t	he molecul	e of (A) has the hy	bridiz	zation
	(a)	sp ²	(b)	sp	(c)	sp ³	(d)	dsp ²
48.	The g	as (B) is						
	(a)	$SO_3(sp^3, 10^3)$	$9\frac{1^{\circ}}{2}$		(b)	SO ₂ (sp ³ , 12	20°)	
	(c)	$SO_3(sp^2, 120)$	0°)		(d)	SO ₂ (sp ² , 12	20°)	
49.	The y	ellow particle	e (C)	is a				
		solid disper			(b)	solid dispe	rsed i	n a solid
	(c)	gas disperse	ed in	a solid	(d)	liquid disp	ersed	in a liquid
50.	The g	reen solution	(D)	contains				
		Cu ²⁺				Ni^{2+}		
	(c)	Cr 6+			(d)	Cr ³⁺		
51.	I ₂ diss	solved in I - gi	iving	; I ₃ (3 lone	pairs)	have		
	(a)	sp ³ d hybrid	izati	on with a b	ent str	ucture		
	(b)	sp ³ d hybrid	izati	on with a t	rigonal	bipyramid	struct	ture
	(c)	sp ³ d hybrid	izati	on with a li	inear st	tructure		
	(d) sp ³ d hybridization with a T-shaped structure							

52.	(a) (b) (c)	ndicator used in iodometric titr starch CuSO ₄ barium diphenylamine sulph alizarin yellow					
53.	(a)	olourless substance (E) obtaine Na ₂ S ₂ O ₃ NaI	(b)	$Na_2S_4O_6$ none of these			
54.	(a)	white precipitate (F) and the bla AgCl and Ag_2S Na ₃ [Ag(S ₂ O ₃) ₂] and Ag_2SO_4	(b)	Ag ₂ S and Na ₂ S ₄ O ₆			
55.	AgN((a)	0 1	(b)	produced by (A) with excess $Na_{4}[Ag_{6}(S_{2}O_{3})_{5}]$ None of these			
•	• A white crystalline solid (A) does not decompose on heating. When warmed with concentrated H ₂ SO ₄ , it gives a reddish brown gas (B), which when passed into water gives a colourless solution. The aqueous solution of (A) in dilute HNO ₃ /AgNO ₃ produces a pale yellow precipitate (C), which dissolves in excess Na ₂ S ₂ O ₃ to produce a soluble complex (D). The compound (A) in concentrated HCl imparts a lilac colour flame.						
56.	(a)	ompound (A) is a salt of lithium potassium		manganese calcium			
57.	(a)	as (B) may be NO_2 only $Br_2 + I_2$		$NO_2 + Br_2$ $NO + Br_2$			
58.	(a)	as (B) dissolved in excess chlor BrCl HBrO	(b)	vater produces HClO HBrO ₃			
59.	(a)	roduct (C) on fusion with Na_2O Ag_2O Ag_2CO_3	(b)	gives Ag a mixture of Ag ₂ O and Ag			

- **60.** The brown gas (B) dissolves in Na_2CO_3 solution, producing two compounds containing elements with oxidation numbers -1 and +3 respectively. The mixture on boiling with concentrated H_2SO_4 produces
 - (a) (C)
- (b) (D)
- (c) (B)
- (d) none of these

- **61.** The complex (D) so produced has
 - (a) an intense yellow colour
- (b) no colour
- (c) a light pink colour
- (d) a deep blue colour
- 62. The compound (C) is mixed with quinhydrone used in
 - (a) photography

(b) photosensitization

(c) tomography

(d) metallurgy

1. d	2. b	3. a	4. b	5. b
6. a	7. c	8. b	9. a	10. c
11. a	12. d	13. a	14. c	15. d
16. c	17. b	18. c	19. d	20. a
21. b	22. a	23. c	24. d	25. d
26. b	27. c	28. d	29. a	30. d
31. b	32. d	33. b	34. b	35. a
36. b	37. d	38. a	39. c	40. a
41. a	42. d	43. b	44. c	45. d
46. b	47. c	48. d	49. a	50. d
51. c	52. a	53. b	54. d	55. a
56. c	57. b	58. d	59. b	60. c
61. b	62. a			

Part 3

Organic Chemistry

1

Classification, Nomenclature and Hybridization

Choose the correct option. Only one option is correct.

1. The name of

$$\begin{array}{c} \mathrm{CH_3CH}(\mathrm{C_6H_5})\mathrm{CH_2CH} - \mathrm{CH_2CH_3} \\ \mathrm{OH} \end{array}$$

is

- (a) 1-ethyl-3-phenyl-1-butanol
- (b) 2-phenyl-4-hexanol
- (c) 5-phenyl-3-hexanol
- (d) 5-benzyl-3-hexanol

2. The name of



is

- (a) bicyclo [2.2.1] heptane
- (b) methylene cyclohexane
- (c) ethylene cyclopentane
- (d) none of these

3. The IUPAC name of

$$CH_2 = CH_2$$

 $CH_3CH_2 - CH_2 - CH_2CH_2CH_3$

- (a) 3-propyl-1-hexene
- (b) 3,3-dipropyl-1-propene
- (c) 4-ethenyl-heptane
- (d) none of these

4. The IUPAC name of

$$\begin{array}{cccc} \mathrm{CH_3CH_2CH-CH-CH_2-CHCH_3} \\ \mathrm{I} & \mathrm{I} & \mathrm{I} \\ \mathrm{CH_3} & \mathrm{CH_3} & \mathrm{CH_2} \\ & \mathrm{CH_3} \end{array}$$

is

- (a) 3, 4-dimethyl-6-ethylheptane
- (b) 2-ethyl-4,5-dimethylheptane
- (c) 3,4,6-trimethyloctane
- (d) 3,5,6-trimethyloctane

5. The JUPAC name of

is

- (a) 2,5-dimethyl-3-propylheptane
- (b) 3,6-dimethyl-5-propylheptane
- (c) 3-methyl-5-isopropyloctane
- (d) none of these

6. The name of the compound is

- (a) bicyclo [2.2.2] octane
- (b) bicyclo [3.2.1] octane
- (c) bicyclo [4.1.1] octane
- (d) bicyclo [4.2.0] octane

7. The IUPAC name of

$$\begin{array}{ccc} & \text{CH}_3 \\ & \text{CH}_3 & \text{CH}_2 \\ & \text{CH}_3 & \text{CH}_2 \\ \text{CH}_3 & \text{CH}_2 & \text{CH} - \text{CH}_3 \\ & \text{CH}_3 & \text{CH}_3 \end{array}$$

- (a) 2,2-dimethyl-4-ethylpentane
- (b) 3,5,5-trimethylhexane
- (c) 2,2,4-trimethylhexane
- (d) 1-tert. butyl-2-ethylpropane

8. The IUPAC name of

is

- (a) 4-sec. butyl-5-methylhexane (b) 2-methyl-3-sec. butylhexane
- (c) 2-methyl-3-isobutylhexane (d) 3-methyl-4-isopropylheptane
- 9. The correct name of the structure

$$H_{3C}$$
 $C=C$ H $C=C$ CH_{3}

is

- (a) (E), (E)-2, 4-hexadiene
- (b) (Z), (Z)-2, 4-hexadiene
- (c) (E), (Z)-3, 5-hexadiene
- (d) (Z), (E)-2, 4-hexadiene
- 10. Which of the following is a cumulated diene?
 - (a) 1,3-Pentadiene
- (b) 1,4-Pentadiene

- (c) 2,3-Pentadiene
- (d) 1,5-Hexadiene

11. The IUPAC name of

$$CH_3 - C - C = C - CH_3$$

$$CH_2CH_3$$

is

- (a) 3-methyl-4-hexyne
- (b) 4-methyl-2-hexyne
- (c) 4-ethyl-2-pentyne
- (d) 2-ethyl-3-pentyne

12. The IUPAC name of

- (a) 3,5-dimethylcyclohexene (b) 4,6-dimethylcyclohexene
- (c) 3-methyltoluene-4-ene
- (d) none of these

13. The IUPAC name of H–C≡C–CH₂CH=CH₂ is (a) 3-acetynyl-1-propene (b) 1-penten-4-yne (c) acetylene-1-propene (d) none of these 14. The IUPAC name of CH₃CHCH₂CH=CH₂ is (a) 4-hydroxypentene-1 (b) ethenylisopropanol (d) 2-hydroxy-4-pentene (c) 4-penten-2-ol 15. The IUPAC name of CH_3-CH_2 C=C CH_2-CH_3 $CH_2CH_2CH_3$ is (a) 4-ethyl-3-methyl-trans-3-heptene (b) 4-ethyl-3-methyl-cis-3-heptene (c) 5-ethyl-6-methyl-trans-5-heptene (d) 5-ethyl-6-methyl-cis-5-heptene 16. 1, 2-dimethylcyclopropane exhibits (b) position isomerism (a) geometrical isomerism (c) optical isomerism (d) nuclear isomerism 17. How many structural isomers can compounds with the molecular formula C₄H₈ have? (a) One (b) Two (d) Four (c) Three **18.** How many structures can heptane (C_7H_{16}) have? (a) Five (b) Six (c) Eight (d) Nine 19. Which of the following can have functional-group isomerism? (a) $CH_3OC_2H_5$ (b) CH₃CH₂NH₂ (c) CH₃CH₂CH=CH₂ (d) none of these 20. How many stereoisomers of cyclopropane 1, 2-dicarboxylic acid are possible? (a) One (b) Two (c) Three (d) Four

21. Which of the following statements is true about a homologous series?(a) Adjacent members of a group differ by a mass of 14.(b) Adjacent members of a group differ by one —CH₂ group.

- (c) Members of a homologous series can be prepared by the same general methods.
- (d) Members of a homologous series have the same physical and chemical properties.
- 22. The large number of organic compounds is due to
 - (a) the valency of carbon
 - (b) the small size of carbon
 - (c) a special property of carbon known as catenation
 - (d) none of these
- 23. How many isomers are possible for the alkyl group C_4H_9 —?
 - (a) Two

(b) Three

(c) Four

- (d) Five
- 24. Which of the following compounds will have only primary and tertiary carbon?
 - (a) Pentane

- (b) 2-Methylbutane
- (c) 2,3-Dimethylbutane
- (d) 2-Bromo-2-methylpropane
- 25. Which of the following compounds will have only primary and secondary carbon?
 - (a) Propane

- (b) 2,2,3-Trimethylpentane
- (c) 2-Methylpropane
- (d) n-Propylbromide
- 26. Which of the following compounds has an isopropyl group?
 - (a) 2-Methylpentane
- (b) 2,2-Dimethylpentane
- (c) 2,2,3,3-Tetramethylpentane (d) 2,2,3-Trimethylpentane
- 27. The general molecular formula of an alkyne is
 - (a) $C_n H_{2n+2}$

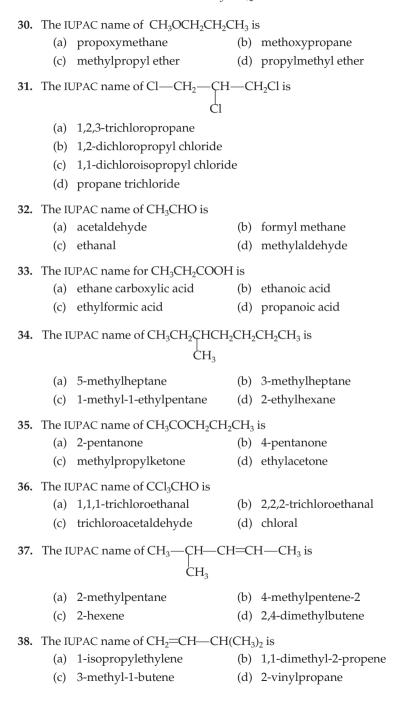
(b) C₁₁H₂₁₁

(c) $C_n H_{2n-2}$

(d) $C_n H_{2n+1} OH$

28. The IUPAC name of

- (a) 1,2-dichloropropane
- (b) 3,3-dichloropropane
- (c) 1,1-dichloropropane
- (d) dichloropropane
- **29.** The IUPAC name of $(CH_3)_2CHCH(CH_3)_2$ is
 - (a) 1,1,2,2-tetramethylethane
- (b) 1,2-di-isopropylethane
- (c) 2,3-dimethylbutane
- (d) 2,3,3-trimethylbutane



39. The IUPAC name of $CH_3CH_2C=CH_2$ is CH_3

- (a) 3-methylbutene-1
- (b) 2-methylbutene-1
- (c) vinylmethylethane
- (d) propylethene-1
- **40.** The IUPAC name of $CH_3CH_2CH = C CH_3$ is CH_2CH_3
 - (a) 2-ethyl-2-pentene
- (b) 4-ethyl-2-pentene
- (c) 3-methyl-3-hexene
- (d) 3-methyl-2-pentene
- 41. The IUPAC name of $(CH_3)_3C$ —CH= CH_2 is
 - (a) 1,1-dimethyl-3-butene
- (b) 3,3-dimethyl-1-butene

CH₂

- (c) 3,3,3-trimethyl-1-propene
- (d) 1,1,1-trimethyl-2-propene

42. The IUPAC name of the compound $H_2C = \stackrel{\downarrow}{C} - C = C - CH_3$ is

- (a) 2-methylpent-1-ene-3-yne (b) 4-methylpent-4-ene-2-yne
- (c) 2-methylpent-2-ene-3-yne
- (d) 2-methylpent-3-yne-2-ene
- **43.** The IUPAC name of C_2H_5 —C—CH— CH_3 is H_2C CH_3
 - (a) 3-methyl-2-ethylbutene-1 (b) 3-ethyl-3-methylbutene-1
 - (c) 2-ethyl-3-methylbutene-1
 - (d) ethylisopropylethene
- **44.** The IUPAC name of CH_3 —CH—CHO is
 - (a) 2-methylbutanal
- (b) butan-2-aldehyde
- (c) 2-ethylpropanal
- (d) 3-methylisobutyraldehyde
- **45.** The IUPAC name of CH_3 —CH— CH_2 — CH_2 —OH is CH_3
 - (a) pentanol

- (b) 1-pentanol
- (c) 2-methyl-4-butanol (d) 3-methyl-1-butanol
- **46.** The IUPAC name of CH₃CH₂CHCH₂OH is CH2CH3
 - (a) 2-methyl-1-pentanol
- (b) 2-ethylbutanol-1
- (c) 2-ethylpentanol-1
- (d) 3-ethylbutanol-1

47. The IUPAC name for
$$CH_3CHOHCH_2$$
— C — OH is CH_3

- (a) 2-methyl-2,4-pentanediol
- (b) 1,1-dimethyl-1,3-butanediol
- (c) 1,3,3-trimethyl-1,3-propanediol
- (d) 4-methyl-2,4-pentanediol

48. The IUPAC name for
$$CH_3$$
— CH — CH = CH — CH_2 — C — OH is CH_3

- (a) 5-carboxy-2-methylpentene
- (b) 4-isopropyl-3-butenoic acid
- (c) 5-methyl-4-hexenoic acid
- (d) none of these
- **49.** The structure of 4-methylpentene-2 is
 - (a) (CH₃)₂CH—CH=CH—CH₃
 - (b) (CH₃)₂CH—CH₂CH=CH₂
 - (c) (CH₃)₂CH—CH₂CH=CH—CH₃
 - (d) $(CH_3)_2C=CHCH_2CH_3$
- **50.** 2-methyl-2-butene is represented as

(a)
$$CH_3$$
— C = $CHCH_3$ (b) CH_3 — CH_2 — C = CH_2 CH_3

51. The IUPAC name of
$$CH_3$$
— C — Br is CH_3

- (a) tertiary butylbromide
- (b) isobutylbromide
- (c) 2-bromo-2-methylpropane
- (d) 2-methyl-2-propylbromide

52. The IUPAC name of CH₃—CH=CHCH₂Br is

- (a) 1-bromo-3-butene
- (b) 1-bromo-2-butene
- (c) 2-butene-1-bromide
- (d) 4-bromo-2-butene

53. The IUPAC name of $(CH_3)_3C$ —OH is

- (a) tert. butylalcohol
- (b) 2-methyl-2-propanol
- (c) 2-methyl-1-butanol
- (d) 2-propanol

54. The IUPAC name of CH₃COCH₂CH₂CH₃ is

- (a) methyl-*n*-propylketone
- (b) 2-pentanone

(c) 3-pentanone

(d) n-propylmethylketone

55. The IUPAC name of
$$CH_3$$
— C — C — CH_2 — CH_3 is $CH_3CH_2CH_2CH_3$

- (a) 3-ethyl-2,2,3-trimethylheptane
- (b) 2,2,3-trimethyl-3-*n*-butylpentane
- (c) 3-methyl-3-isopropylheptane
- (d) 2,2-dimethyl-3-ethyl-3-n-butylbutane

56. The IUPAC name of
$$CH_3$$
— CH — CH_2 — CH — CH_2 CH $_3$ is CH_2CH_3

- (a) 2-ethyl-4-methylhexane (b) 3,5-dimethylheptane
- (c) 5-ethyl-3-methylhexane
- (d) 2,4-diethylpentane

57. The IUPAC name of
$$CH_3CH$$
— CH — C — CH_3 is OH OH

- (a) 1,1,2-trimethyl-1,3-butanediol
- (b) 1,2-dimethyl-2,4-pentanediol
- (c) 2,3-dimethyl-2,4-pentanediol
- (d) 1,2,3,4-tetramethyl-1,3-propanediol

58. The IUPAC name of
$$CH_3C = CH - CH_3$$
 is

- (a) 4-methyl-2-pentyne
- (b) methylisopropylacetylene
- (c) 4,4-dimethyl-2-butyne
- (d) 2-methyl-4-pentyne

59. The structure of 4-methyl-2-penten-1-ol is (a) $(CH_3)_2CHCH_2$ = $CHCH_2OH$

	(b)	CH ₃ CHOH—CH=C(CH ₃) ₂		
	(c)	(CH ₃) ₂ =CHCH ₂ CH ₂ OH		
	(d)	CH ₃ CH ₂ CH=CHCH ₂ OH		
60.	Which	n of the following compounds	are n	named correctly?
	(a)	(CH ₃) ₂ CHCH ₂ CH ₂ CH ₂ CHO (5-me	ethyl-1-hexanal)
	(b)	$(CH_3)_2CHCH_2C \equiv C - COOH$	(5-n	nethyl-2-hexynoic acid)
	(c)	CH ₃ CH ₂ CH ₂ CH ₂ CH(CH ₃)CO	ОН	(2-methylhexanoic acid)
	(d)	CH ₃ CH ₂ CH=CH—COCH ₃ (3	-hex	ren-5-one)
61.	The IU	JPAC name of (CH ₃) ₃ C—CH ₂ C	H=(CH ₂ is
	(a)	2,2-dimethylpent-4-ene	(b)	2,2-dimethylhex-4-ene
	(c)	4,4-dimethylpent-1-ene	(d)	hex-1-ene
62.	In whi	ich of the following are all carb	on a	toms sp-hybridized?
	(a)	CH ₃ —CH=CH—CH ₃	(b)	CH_3 — C = C — CH_3
	(c)	HC≡C—C≡CH	(d)	CH ₃ CH ₂ —C≡CH
63.	The l	hybridization of carbon ato C—CH=CH ₂ is	ms	in the C—C single bond of
		sp^3-sp^3		sp^2-sp^3
	(c)	sp–sp ²	(d)	sp ³ –sp
64.	Which	n of the following compounds l	nave	only one type of hybrid carbon?
		CH ₂ =CH—CH=CH ₂	. ,	HC≡C—C≡CH
	(c)	CH_3 — CH_2 — CH_2 — CH_3	(d)	CH_3 — $C\equiv C$ — CH_3
65.	In the	$reaction \ CH_3CONH_2 \frac{P_2O_5}{\Delta}$	→ CH	I_3 CN, the hybridization state of
		rbon atom changes from		
		sp^2 to sp		sp^3 to sp
		sp ³ to sp ²		$sp^2 to sp^3$
66.		${f n}$ of the following have zero dip ${f H}_2$		moment? HF
		CH ₄		CHCl ₃
	. ,	•	. ,	-

67. Arrange the following resonating structures of vinyl chloride in order of decreasing stability.

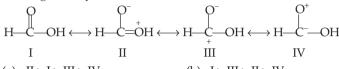
(a) I > II > III

(b) III > II > I

(c) II > I > III

(d) I > II = III

68. Arrange the following resonating structures of formic acid in order of decreasing stability.



(a) II > I > III > IV

(b) I > III > II > IV

(c) III > II > IV > I

- (d) IV > III > I > II
- **69.** Which of the following molecules show resonance?
 - (a) CO

(b) CO₂

(c) NO

(d) O_3

70. Arrange the following free radicals in order of stability.

71. Arrange the following free radicals in order of stability.

Benzyl, allyl, methyl, vinyl I II III IV

(a) IV > III > II > I

(b) I > II > III > IV

(c) II > IV > III > I

(d) III > II > IV

72. Arrange the following carbonium ions in order of decreasing stability.

+	+	+	+
$(CH_3)_3C$	$(CH_3)_2CH$	CH_3CH_2	H_3C
I	II	ĬΙΙ	Ϊ́V
(a) II > III >	I > IV	(b) IV	> III > II >

(c) I > II > III > IV

(d) I > II > III = IV

Ι

73. Which of the following compounds will produce the most stable carbonium ion?

(a)
$$CH_3$$
— CH — CH_2OH (b) CH_3 — C — OH CH_3 (c) CH_3 — CH — CH_2CH_3 (d) $CH_3CH_2CH_2CH_2OH$

- 74. The bond that undergoes heterolytic cleavage most readily is
 - (a) C—C

(b) C—O

(c) C—H

- (d) O—H
- 75. Hydrogen bonding is maximum in
 - (a) triethylamine

(b) ethyl chloride

(c) ethyl alcohol

- (d) diethyl ether
- 76. Which of the following compounds will exhibit cis-trans isomerism?
 - (a) 2-Butene

(b) 2-Butyne

(c) 1-Butene

- (d) 2-Butanol
- 77. Which of the following compounds will exhibit geometrical isomerism?

C = C CH_2 C = C CH_3

(a)
$$H$$
 $C=C$ Br
(c) H_3C $C=C$ Br
 H_3C $C=C$

- $(d) \qquad H C = C \begin{cases} Br \\ Br \end{cases}$
- 78. Which of the following compounds exhibit optical isomerism?
 - (a) $C_2H_5OC_2H_5$

- (b) CH₃OC₃H₇
- (c) CH₃CH₂CH₂CH₂OH
- (d) CH₃CHOHCH₂CH₃

The IUPAC name of this compound is

- (a) 2,2,6-trimethyl-4-(1-methylpropyl) nonane
- (b) 4,8,8-trimethyl-6-(1-methylpropyl) nonane
- (c) 3,6-dimethyl-4-(methylene tertiary butyl) nonane
- (d) 6,6-dimethyl-2-propyl-4-(1-methylpropyl) heptane

The IUPAC name of this compound is

- (a) 3-ethyl-4-chloro-1,4-pentadiene
- (b) 2-chloro-3-ethyl-1,4-pentadiene
- (c) 4-chloroethenyl-1-pentene
- (d) 3-ethenvl-4-chloro-4-pentene

The IUPAC name of this compound is

- (a) 2-ethoxy-4-methoxypentan-3-one
- (b) 2-methoxy-4-ethoxy-pentan-3-one
- (c) 2-ethoxy-4-methoxy-pentan-3-one
- (d) none of these

82. The IUPAC name of
$$CH_3$$
— CH — CH — CH_2 — CHO is CH_3 CH_3

- (a) 2,3-dimethylpentanal (b) 3,4-dimethylpentanal
- (c) 3,4,4-trimethylbutanal (d) 3-isopropylbutanal

83. The IUPAC name of
$$CH_3$$
— CH — CH_2 — $C(CH_3)_2$ is OH OH

- (a) 2-methyl-2,4-dihydroxypropane
- (b) 2,2-dimethyl-4-hydroxybutanol
- (c) 2-methyl-2,4-pentanediol
- (d) 2-hydroxy-4,4-dimethylbutanol-4

84. The IUPAC name of
$$\mathrm{BrCH_2}$$
 —CH—CO—CH $_2$ —CH $_2$ CH $_3$ is CONH_2

- (a) 2-bromomethyl-3-oxohexanamide
- (b) 1-bromo-2-amido-3-oxohexane
- (c) 1-bromo-2-amido-*n*-propylketone
- (d) 3-bromo-2-proponyl-propanamide

85. The IUPAC name of
$$CH_3$$
— CH_2 — CH — $COOC_2H_5$ is CH_2

- (a) 2-ethyl-ethylacetate
- (b) ethyl 3-methylbutanoate
- (c) ethyl 2-methylbutanoate
- (d) 2-methylbutanoic acid ethylester

86. The IUPAC name of
$$CH_3CH_2$$
—N— CH_2CH_3 is CH_3

- (a) N-methyl-N-ethylethylamine
- (b) diethylmethylamine
- (c) N-ethyl-N-methylaminoethane
- (d) methyldiethylamine

The IUPAC name of this compound is

- (a) 2-isopropylbutanal
- (b) 2-ethyl-3-methylbutanal
- (c) 3-ethyl-2-methylbutanal (d) 2-methylpentane-3-aldehyde

88. The IUPAC name of
$$C_6H_5$$
— CH — CH_2 — CCl_3 is C_6H_5

- (a) 1,1,1-trichloro-3,3-diphenylpropane
- (b) 1,1-diphenyl-3,3,3-trichloropropane
- (c) (a) as well as (b)
- (d) none of these

89. The IUPAC name of $C_6H_5CH=CH$ —COOH is

- (a) cinnamic acid
- (b) 1-phenyl-2-carboxyethene
- (c) 3-phenylprop-2-enoic acid
- (d) dihydro-3-phenylpropionic acid

90. The IUPAC name of HC \equiv CCH₂CH \equiv CH₂ is

- (a) 1-propyn-ethene
- (b) propeneacetylene
- (c) pent-4-yne-1-ene
- (d) pent-1-en-4-yne

91. The IUPAC name of
$$CH = CH - CHCH_2CH_3$$
 is CH_3

- (a) 1-cyclohexyl-3-methyl-1-pentene
- (b) 3-methyl-5-cyclohexyl-pent-1-ene
- (c) 1-cyclohexyl-3-ethyl-but-1-ene
- (d) 1-cyclohexyl-3,4-dimethyl-but-1-ene

The IUPAC name of this compound is

- (a) 3,3-dimethyl-2-pentanol
- (b) 3-methyl-3-ethyl-2-butanol
- (c) 3,3-dimethyl-3-ethyl-isopropanol
- (d) 3,3-dimethyl-3-ethyl-2-hydroxypropane

The IUPAC name of this compound is

- (a) 2-fluoro-4-chloro-2,4-diethylpentane
- (b) 3-fluoro-5-chloro-3-methyl-5-ethylhexane
- (c) 3-chloro-5-fluoro-3,5-dimethylheptane
- (d) 3,5-dimethyl-5-fluoro-3-chloroheptane
- 94. The —I effect of —NO₂, —CN, —COOH, —Cl decreases in the order

(a)
$$-NO_2 > -CN > -COOH > -Cl$$

- (b) $-Cl > -COOH > -CN > -NO_2$
- (c) $-CN > -NO_2 > -Cl > -COOH$
- (d) $-COOH > -CN > -NO_2 > -Cl$
- 95. The +I effect of $(CH_3)_3C(I)$, $(CH_3)_2CN(II)$, $CH_3C—H_2(III)$, $CH_3(IV)$ decreases in the order
 - (a) I > II > III > IV

(b) IV > III > II > I

(c) II > I > IV > III

(d) I > II > IV > III

- 96. The inductive effect of the alkyl groups on a saturated carbon chain follows the order
 - (a) $(CH_3)_3 C \longrightarrow (CH_3)_2 CH \longrightarrow CH_3 CH_2 \longrightarrow CH_3 CH_3$
 - (b) $CH_3 \longrightarrow CH_3 \longrightarrow CH_2 \longrightarrow (CH_3)_2 CH \longrightarrow (CH_3)_3 C \longrightarrow (CH_3)_3 CH \longrightarrow (CH_$
 - (c) $CH_3 CH_2 \longrightarrow CH_3 \longrightarrow (CH_3)_3 C \longrightarrow (CH_3)_2 CH \longrightarrow$
 - (d) $(CH_3)_2CH \rightarrow (CH_3)_3C \rightarrow CH_3 \rightarrow CH_3CH_2$
- 97. Give the IUPAC name of

- (a) 4-isopropyl-5-tert. butyloctane
- (b) 4-tert. butyl-5-isopropyloctane
- (c) 2-methyl-3-propyl-4-tert. butylheptane
- (d) 2, 2-dimethyl-3-propyl-4-isopropylheptane
- **98.** The IUPAC name of

$$CH_3$$
— CH_2
 $C=C$
 CH_2 — CH_3
 CH_2 — CH_3
 CH_2 — CH_3 — CH_3

- (a) 4-ethyl-3-methyl-trans-3-heptene
- (b) 4-ethyl-5-methyl-trans-4-heptene
- (c) 3-methyl-4-propyl-3-hexene
- (d) 3-propyl-4-ethyl-3-pentene

- 99. The IUPAC name of ClCH₂CH=CCH₂—OH is
 - (a) 5-chloro-3-penten-3-carbinol
 - (b) 1-chloro-3-penten-3-carbinol
 - (c) 4-chloro-2-ethyl-2-buten-1-ol
 - (d) 1-chloro-3-ethyl-2-buten-4-ol

- CH_3 **100.** The IUPAC name of CH_2 =C—CHO is
 - (a) methacrolein

- (b) methacrylaldehyde
- (c) 2-methylpropenal
- (d) propenaldehyde

101. The IUPAC name of
$$C - CH_3$$
 is

- (a) phenylethanone
- (b) methylphenylketone

(c) acetophenone

(d) phenylemethylketone

C=CH

102. The IUPAC name of HC=C-C=C-CHCH
$$_2$$
-CH $_3$ is

- (a) 5-ethyl-1,3, 6-heptatriyne (b) 3-ethyl-2,4, 5-heptatriyne
- (c) 5-ethenyl-1,3-heptatriyne (d) 3-ethenyl-4,6-heptatriyne
- 103. The IUPAC name of

$$CH_3(CH_2)_7 > C = C < H (CH_2)_7 COOH$$

is

(a) elaidic acid

- (b) trans-octadec-9-enoic acid
- (c) dihydrosteric acid
- (d) oleic acid

104. The IUPAC name of

$$CH_3(CH_2)_4$$
 $C=C < CH_2$ $C=C < (CH_2)_7COOH$

- (a) cis-cis-9,12-octadecadienoic acid
- (b) cis-trans-9,12-octadecadienoic acid
- (c) 9,10-octadecadienoic acid
- (d) 9,14-octadecadienoic acid
- 105. The IUPAC name of

- (a) 6-(1-methylbutyl)-8-(2-methylbutyl) tetradecane
 - (b) 6-(2-methylbutyl)-8-(1-methylbutyl) tetradecane
 - (c) 4-methyl-5-*n*-pentyl-7-(2-methylbutyl) tridecane
- (d) 3-methyl-5-n-hexyl-7-(1-methylbutyl) didecane

106. The IUPAC name of OHC—CH₂—CH₂—CH—CH₂—CHO is

- (a) 4, 4-di(formylmethyl) butanal
- (b) 2-(formylmethyl) butane-1,4-dicarbaldehyde
- (c) hexane-3-acetal-1.6-dial
- (d) 3-(formylmethyl) hexane-1,6-dial

107. The IUPAC name of

is

- (a) 4-carboxyheptane-1,7-dioic acid
- (b) 3-propionyl-propane-1,3-dioic acid
- (c) pentane-1,3, 5-tricarboxylic acid
- (d) 4-methionate butane-1,3-dioic acid

108. The IUPAC name of

is

- (a) 2-chlorocarbonyl ethylbenzoate
- (b) 2-carboxyethyl benzoyl chloride
- (c) ethyl-2-(chlorocarbonyl) benzoate
- (d) ethyl-1-(chlorocarbonyl) benzoate

109. The IUPAC name of C_6H_5CN is

- (a) phenyl cyanide
- (b) phenylacetonitrile
- (c) benzene cyanide
- (d) benzonitrile

110. The IUPAC name of $C_6H_5CH_2CH_2NH_2$ is

- (a) β -phenylethylamine (b) 2-phenylaminoethane
- (c) 2-phenylethanamine (d) benzyl methylamine

111. The IUPAC name of
$$CH_3$$
— NH — CH — CH_2 — CH — CH_3

$$CH_3$$

- (a) 2-(n-methylamino)-4-methylpentane
- (b) n,4-dimethylpentan-2-amine
- (c) 2-(n-methylamino)-3-isopropylpropane
- (d) 2-(n-methylamino)-1,4,4-trimethylbutane

Answers				
1. c	2. a	3. a	4. c	5. c
6. d	7. c	8. d	9. d	10. c
11. b	12. a	13. b	14. c	15. a
16. a	17. d	18. d	19. a	20. c
21. a, b, c	22. c	23. c	24. c, d	25. a, d
26. a	27. c	28. c	29. c	30. b
31. a	32. c	33. d	34. b	35. a
36. b	37. b	38. c	39. b	40. c
41. b	42. a	43. c	44. a	45. d
46. b	47. a	48. d	49. a	50. a
51. c	52. b	53. b	54. b	55. a
56. b	57. c	58. a	59. a	60. a, b, c
61. c	62. c	63. c	64. a, b, c	65. a
66. a, c	67. a	68. b	69. a, b, c, d	70. a
71. b	72. c	73. b	74. d	75. c
76. a	77. b	78. d	79. a	80. b
81. a	82. b	83. c	84. a	85. c
86. a	87. b	88. a	89. c	90. d
91. a	92. a	93. c	94. a	95. a
96. a	97. b	98. a	99. c	100. c
101. a	102. a	103. b	104. a	105. a
106. d	107. c	108. c	109. d	110. c
111. b				

Hints to More Difficult Problems

- 2 & 6 Compounds of this type are named systematically by attaching the prefix 'bicyclo' to the name of the open-chain hydrocarbon with the same total number of carbon atoms in the ring. The size of the two rings is specified by the number of carbon atoms in each of the three linkages which connect the two atoms at the ring junctions.
 - **15.** According to IUPAC nomenclature, the name is assigned by taking the longest continuous chain as it passes through the double bond.

Reaction Mechanisms

• Type 1 •

1. The nucleophilicities of R_3C^- , R_2N^- , RO^- and F^- decrease in the order

Choose the correct option. Only one option is correct.

	(a)	$R_3C^- > R_2N^- > RO^- > F^-$	(b)	$F^- > RO^- > R_2N^- > R_3C^-$
	(c)	$RO^- > F^- > R_2N^- > R_3C^-$	(d)	$R_3C^- > RO^- > F^- > R_2N^-$
2.	The nucleophilicities of RO ⁻ , HO ⁻ , RCOO ⁻ , ROH and H ₂ O are of t			
	order			
	(a)	$HO^- > RO^- > H_2O > ROH > F$	RCO(O-
	(b)	$RO^- > HO^- > RCOO^- > ROH > H_2O$		
	(c)	c) $H_2O > ROH > RCOO^- > HO^- > RO^-$		
	(d)	$ROH > H_2O > HO^- > RCOO^-$	> R0	O ⁻
3.	The o	The order of leaving group ability for halides is		
	(a)	$C1^- > Br^- > I^- > F^-$	(b)	$F^- > Br^- > I^- > Cl^-$
	(c)	$Br^- > Cl^- > F^- > I^-$	(d)	$I^- > Br^- > Cl^- > F^-$
4.	The h	hydrolysis of tert. butylchloride follows		
	(a)	first-order kinetics	(b)	second-order kinetics
	(c)	pseudo first-order kinetics	(d)	none of these
5.	The o	rder of basicity of halides is		
	(a)	$Cl^- < Br^- < I^- < F^-$	(b)	$F^- < I^- < Br^- < Cl^-$
	(c)	$I^- < Br^- < Cl^- < F^-$	(d)	$Cl^{-} < F^{-} < I^{-} < Br^{-}$
6.	The h	ydrolysis of 2-bromo-3-methyl	buta	ne yields

(b) 2-methyl-3-butanol

(d) none of these

(a) 2-methyl-2-butanol

(c) a mixture of (a) and (b)

- 7. The stability of $\mathring{1}$, $\mathring{2}$, $\mathring{3}$ and benzyl carbocations is of the order
 - (a) $\mathring{1} > \mathring{2} > \mathring{3} > \text{benzyl}$
- (b) benzyl > 3 > 2 > 1
- (c) $\stackrel{\circ}{3} > \stackrel{\circ}{2} > \stackrel{\circ}{1} > \text{benzyl}$
- (d) $\stackrel{\circ}{3} > \text{benzyl} > \stackrel{\circ}{2} > \stackrel{\circ}{1}$
- 8. The order of reactivity of alkyl halides is
 - (a) tert. alkyl halides > allyl halides > sec. alkyl halides > primary alkyl halides
 - (b) primary alkyl halides > sec. alkyl halides > tert. alkyl halides > allyl halides
 - (c) allyl halides > primary alkyl halides > sec. alkyl halides > tert. alkyl halides
 - (d) sec. alkyl halides > primary alkyl halides > tert. alkyl halides > allyl halides
- 9. Which of the following hydrolyses the fastest?
 - (a) CH₃Cl

(b) CH₃CH₂Cl

(c) CH₃OCH₂Cl

- (d) (CH₃)₂CHCl
- 10. Which of the following alkyl halides would be the most reactive in an $S_{NJ}2$ reaction?
 - (a) $C_6H_5CH_2CH_2CH_2Br$

- (d) $C_6H_5CH_2CHCH_3$
- 11. Which of the following alkyl halides would be the most reactive in an S_N 1 reaction?

 - (a) $C_6H_5CH_2CH_2CH_2Br$ (b) C_6H_5CH — CH_2Br CH_3

(c)
$$C_6H_5C$$
—Br CH_3

- (d) C₆H₅CH₂CHCH₃ | | Br
- 12. Arrange NO₂, F, COOH, Cl, OH, OR and C₆H₅ in order of electron-withdrawing effect.
 - (a) $F > NO_2 > COOH > OH > Cl > OR > C_6H_5$
 - (b) $COOH > NO_2 > F > C1 > OH > C_6H_5 > OR$
 - (c) $C_6H_5 > OR > Cl > OH > COOH > NO_2 > F$
 - (d) $NO_2 > F > COOH > C1 > OH > OR > C_6H_5$

- 13. Arrange the groups CH₃—, CH₃CH₂—, (CH₃)₂CH— and (CH₃)₃C— in order of electron-releasing effect.
 - (a) $(CH_2)_2C \longrightarrow (CH_2)_2CH \longrightarrow CH_2CH_2 \longrightarrow CH_2$
 - (b) $CH_3 \longrightarrow CH_3CH_2 \longrightarrow (CH_3)_2CH \longrightarrow (CH_3)_3C \longrightarrow$
 - (c) $CH_3CH_2 \longrightarrow CH_3 \longrightarrow (CH_3)_3C \longrightarrow (CH_3)_2CH \longrightarrow$
 - (d) $(CH_3)_2CH \rightarrow (CH_3)_3C \rightarrow CH_3 \rightarrow CH_3CH_2$
- 14. The order of the —I effect orbitals is
 - (a) $sp^3 > sp^2 > sp$

(b) $sp^2 > sp^3 > sp$

(c) $sp > sp^2 > sp^3$

- (d) $sp^3 > sp > sp^2$
- 15. The hyperconjugative effect of the group R in R-CH=CH₂, where R is CH₃-, CH₃CH₂- or (CH₃)₂CH, follows the order
 - (a) $CH_2 > CH_2CH_2 > (CH_3)_2CH -$
 - (b) $(CH_3)_2CH_->CH_3CH_2->CH_3-$
 - (c) $CH_3 > (CH_3)_2 CH > CH_3 CH_2 -$
 - (d) $(CH_2)_2CH_->CH_2->CH_2CH_2-$
- **16.** The product obtained in the reaction

$$\begin{array}{c} CH_3 \\ \vdash \\ CH_3 - C - CH - CH_3 \xrightarrow{\quad H_2SO_4 \\ \mid \quad \mid \quad \\ H \quad OH \end{array} \rightarrow$$

is

(a)
$$CH_3$$
 CH_3 (b) CH_3 — C = CH — CH_3 H

(c)
$$CH_2 = C - CH_2 - CH_3$$

- (d) none of these
- 17. The major product obtained in the reaction

- (d) none of these
- 18. Arrange the acids CH₂CH₂COOH(I), CH₂=CH—COOH(II) and HC≡C—COOH(III) in order of decrease in acidity.
 - (a) I > II > III
- (b) III > II > I
- (c) II > I > III
- (d) III > I > II
- 19. The order of stability of the carbon radicals $\mathring{3}$, $\mathring{2}$, $\mathring{1}$ and $\mathring{C}H_3$ is

 - (a) $\mathring{1} > \mathring{2} > \mathring{3} > \mathring{C}H_2$ (b) $\mathring{C}H_3 > \mathring{1} > \mathring{2} > \mathring{3}$
 - (c) $3 > 2 > 1 > CH_0$
- (d) $\dot{C}H_2 > 3 > 2 > 1$
- **20.** The order of stability of the carbocations $\mathring{1}$, $\mathring{2}$, $\mathring{3}$ and $\overset{\oplus}{C}H_3$ is
 - (a) $\overset{\oplus}{C}H_3 > \overset{\circ}{1} > \overset{\circ}{2} > \overset{\circ}{3}$
- (b) $1 > 2 > 3 > CH_3$
- (c) $\mathring{1} > \mathring{2} > \mathring{C}H_2 > \mathring{3}$
- (d) $\stackrel{\circ}{3} > \stackrel{\circ}{2} > \stackrel{\circ}{1} > \stackrel{\oplus}{CH}_{\stackrel{\circ}{a}}$
- **21.** The reaction of *cis*-2-butene with bromine gives
 - (a) meso dibromide
 - (b) racemic dibromide
 - (c) a mixture of meso and racemic dibromide
 - (d) none of these
- **22.** In the reaction

$$(CH_3)_2CH$$
- CH = CH_2 \longrightarrow HCl

the product obtained is

- (a) (CH₃)₂CCl–CH₂CH₃
- (b) (CH₃)₂CH—CH—CH₃ | | Cl
- (c) a mixture of (a) and (b)
- (d) none of these
- 23. The relative rates of addition of

- (a) I > II > III > IV > V
- (b) V > IV > III > II > I
- (c) II > III > I > IV > V
- (d) IV > III > II > V > I
- 24. The conjugation of electron-withdrawing groups

(e.g., —CHO, —C—R, —C—OR, —C
$$\equiv$$
N and —NO₂)

activates nucleophilic addition. The order of reactivity of these groups is

- 25. The order of hydrolysis of the alkyl halides (allyl, $\mathring{3}, \mathring{2}, \mathring{1}, CH_3X$) by the S_N1 path is
 - (a) $\text{allyl} > 3 > 2 > 1 > \text{CH}_3X$ (b) $1 > 2 > 3 > \text{allyl} > \text{CH}_3X$
 - (c) $\mathring{1} > \mathring{2} > \mathring{3} > CH_3X > \text{allyl}$ (d) $CH_3X > \mathring{1} > \mathring{2} > \mathring{3} > \text{allyl}$
- **26.** The rate of hydrolysis of alkyl halides 1° , 2° , 3° and CH_3X by the S_N2 path is
 - (a) $1 > 2 > 3 > CH_2X$ (b) $CH_2X > 3 > 2 > 1$
 - (c) $CH_2X > \hat{1} > \hat{2} > \hat{3}$
- (d) $3 > 2 > 1 > CH_2X$
- 27. The order of elimination of halogens in $\mathring{1}$, $\mathring{2}$ and $\mathring{3}$ alkyl halides is
 - (a) 1 > 2 > 3

(b) 3 > 1 > 2

(c) 3 > 2 > 1

- (d) 2 > 3 > 1
- 28. The major product obtained in the reaction

$$\begin{array}{c|c} CH_3 CH \\ \mid & \mid \\ CH_3 - C - C - CH_3 & \xrightarrow{-HCl} \end{array}$$

29. The major product obtained in the reaction

is

30. The elimination reaction

$$CH_3-CH_2\cdot CH-CH_3 \xrightarrow{\Delta} CH_3CH_2-CH=CH_2$$

$$\stackrel{\oplus}{}_{NMe_3} (major)$$

is governed by

- (a) the Saytzev rule
- (b) the Hofmann rule
- (c) the Saytzev as well as the Hofmann rule
- (d) neither
- **31.** The diazotization of CH₃CH₂CH₂CH₂NH₂ with NaNO₂ and dilute HCl at 0–5°C followed by the warming of the solution gives the following major product.
 - (a) 1-Butanol
 - (b) 2-Butanol
 - (c) An equimolecular mixture of these
 - (d) None of these

32. In the reaction

$$\begin{array}{c|c} CH_3CH_3 \\ & | & | \\ CH_3-C-C-C+CH_3 \end{array} \xrightarrow{H^+}$$

$$\begin{array}{c|c} CH_3CH_3 \\ & | & | \\ OHOH \end{array}$$

the product obtained is

the product obtained is (a)
$$CH_3 - C - C(CH_3)_3$$
 (b) CH_3CH_3 $CH_3 - C - C - CH_3$

(c)
$$CH_3CH_2$$
— C — CH_2 — CH_3 (d) none of these \parallel O

- 33. The reaction of cyclobutylamine with sodium nitrite and dilute HCl at 0-5°C followed by warming gives
 - (a) only cyclobutanol
 - (b) only cyclopropylcarbinol
 - (c) a mixture of cyclobutanol and cyclopropylcarbinol
 - (d) none of these

34. The reaction

is a case of

- (a) Curtius rearrangement
- (b) Hofmann rearrangement
- (c) Schmidt reaction
- (d) Beckmann rearrangement

35. Phenylallylether, C₆H₅OCH₂-CH=CH₂, in which the carbon atom next to O is ¹⁴C, on being heated to 200°C, gives

(a) OH (b) OH
$$CH_2-CH=CH_2$$
 $CH_2-CH=CH_2$ (c) $CH_2-CH=CH_2$ (d) none of these

36. 2,4-Dimethylphenylallyl ether, in which the carbon atom next to the oxygen is ¹⁴C, on being heated to 200°C, gives

(a) OH
$$CH_3$$
 $CH_2-CH=CH_2$

(b) OH
$$CH_3$$
 $CH_2-CH=\overset{*}{C}H_3$

(c) OH
$$CH_3$$
 $CH_2CH = CH_2$

- (d) none of these
- **37.** On heating phenylacetate in the presence of anhydrous aluminium chloride, we obtain
 - (a) only o-hydroxy acetophenone
 - (b) only p-hydroxy acetophenone
 - (c) a mixture of o- and p-hydroxy acetophenone
 - (d) none of these
- 38. The reaction of acetaldehyde with an excess of formaldehyde gives

(a)
$$H_2C-OH$$
 (b) CH_2OH CH_2CHO $HOH_2C-C-CHO$ CH_2OH

(c)
$$CH_2OH$$
 (d) none of these $HOH_2C-C-CH_2OH$ CH_2OH

- **39.** The condensation of acetylene with formaldehyde in the presence of Cu₂C₂ gives
 - (a) $HOH_2C-C \equiv C-CH_2OH$
- b) C≡CH
 H₂C C≡CH
- (c) $C \equiv C$ CH_2 $C \equiv C$
- (d) none of these
- **40.** The reaction of an acid chloride with diazomethane followed by treatment with water gives a carboxylic acid. The reaction occurs through the intermediate formation of
 - (a) carbene

(b) ketene

(c) benyne

- (d) a carbocation
- **41.** In the Baeyer–Villiger rearrangement, the group that migrates (in case of unsymmetrical ketones) is the one which is more electron-releasing. The aptitude of migration of the alkyl groups is of the order
 - (a) $\hat{1} > \hat{2} > \hat{3} > CH_3$
- (b) $CH_3 > \mathring{3} > \mathring{2} > \mathring{1}$
- (c) $CH_3 > \hat{1} > \hat{2} > \hat{3}$
- (d) $3 > 2 > 1 > CH_3$
- **42.** In the Baeyer–Villiger oxidation of alkyl aryl ketones, it is the more electron-releasing group that migrates. The aptitude of migration of the aryl groups is of the order
 - (a) p-chlorophenyl > p-anisyl > p-tolyl > phenyl
 - (b) phenyl > p-tolyl > p-anisyl > p-chlorophenyl
 - (c) p-anisyl > p-tolyl > p-ehlorophenyl
 - (d) p-chlorophenyl > phenyl > p-tolyl > p-anisyl
- 43. On treatment with NaOH an oxime of 2-bromo-5-nitroacetophenone did not undergo any change. However, on subjecting it to Beckmann rearrangment, N-(2-bromo-5-nitrophenyl) acetamide was obtained. On the basis of these observations, it may be concluded that the starting oxime has
 - (a) only an anti- or trans-structure
 - (b) only a syn- or cis-structure
 - (c) a mixture of syn- and anti-oximes
 - (d) none of these
- 44. The major product obtained in the reaction

NC—CH=CH
$$_2$$
 —HBr

- (b) NC-CH₂-CH₂Br
- (c) an equimolecular mixture of (a) and (b)
- (d) none of these
- **45.** In the reaction

$$(CH_3)_3CCH=CH_2 \xrightarrow{HCl}$$

the product obtained is

(a) Only
$$(CH_3)_3$$
— CH — CH_3 (b) Only $(CH_3)_2$ — C — $CH(CH_3)_2$ Cl

- (c) a mixture of (a) and (b)
- (d) none of these

46. The reaction

tion
$$CH_3CH=CH_2+HBr \xrightarrow{peroxide} CH_3CH_2CH_2Br$$
 cplained by

can be explained by

- (a) carbocation formation (b) free-radical mechanism
- (c) carboanion formation (d) none of these
- 47. The product obtained in the reaction

CH₃CH=CH₂
$$\xrightarrow{\text{HCl}}$$
 is

(a) CH₃CH - CH₃ (b) CH₃CH₂CH₂Cl

- (c) CH₃CH₂CH₂CH₂CH₂CH₃ (d) none of these
- 48. Maleic anhydride undergoes a Diels-Alder reaction with
 - (a) benzene

(b) naphthalene

(c) phenanthrene

- (d) anthracene
- 49. In the Friedel-Crafts reaction of an aromatic compound with an alkyl halide or acid halide in the presence of a catalyst, the order of reactivity of the following catalysts is
 - (a) $AlBr_3 > AlCl_3 > FeCl_3 > SbCl_5 > BCl_3 > BF_3$
 - (b) $AlCl_3 > FeCl_3 > AlBr_3 > SbCl_5 > BF_3 > BCl_3$
 - (c) $BF_3 > BCl_3 > SbCl_5 > FeCl_3 > AlCl_3 > AlBr_3$
 - (d) $FeCl_3 > AlCl_3 > AlBr_3 > SbCl_5 > BF_3 > BCl_3$

50. In the pinacol-pinacoline rearrangement

the trend of migration among the aryl groups is

- (a) p-anisyl > p-tolyl > p-ehlorophenyl
- (b) p-chlorophenyl > phenyl > p-tolyl > p-anisyl
- (c) phenyl > p-tolyl > p-anisyl > p-chlorophenyl
- (d) p-tolyl > p-anisyl > p-chlorophenyl > phenyl
- **51.** In the nitration of benzene with concentrated HNO_3 and concentrated H_2SO_4 , which of the following is the rate- determining step?

(a)
$$HO-NO_2 + H_2SO_4 \longrightarrow H_2O-NO_2 + HSO_4$$

$$\stackrel{H_2SO_4}{\longrightarrow} H_3O + NO_2 + 2HSO_4$$

(b)
$$H \times NO_2 \times$$

(c)
$$H^{NO_2} + HSO_3 \longrightarrow NO_2 + H_2SO_4$$

- (d) none of these
- 52. Arrange the following radicals in order of decreasing stability.

$$(CH_{3})_{3}C^{\bullet} \quad CH_{2} = CH - C^{\bullet} \quad CH_{3} \quad (CH_{3})_{2}CHCH_{2}^{\bullet} \quad (CH_{3})_{2}CH$$

$$I \quad II \quad III \quad IV \quad V$$

$$(a) \quad I, II, III, IV, V \quad (b) \quad V, I, III, IV, II$$

$$(c) \quad II, I, V, IV, III \quad (d) \quad II, V, I, III, V$$

- 53. Among the following, which is the strongest nucleophile for an S_N2 reaction?
 - (a) H_2O
- (b) RCO₂
- (c) OH-
- (d) RO
- 54. The weakest nucleophile in an aprotic solvent is
 - (a) I⁻
- (b) Br⁻
- (c) Cl⁻
- (d) F

- **55.** The prerequisite for an S_N 2 reaction is
 - (a) a carbocation
 - (b) a carbanion
 - (c) a penta-coordinated transition state
 - (d) none of these
- **56.** The addition of KI increases the rate of the reaction.

$$CH_3CH_2Cl + H_2O \longrightarrow CH_3CH_2OH + HCl$$

This is because I is a

- (a) a good nucleophile and a good leaving group
- (b) a poor nucleophile and a poor leaving group
- (c) a poor nucleophile and a good leaving group
- (d) none of these
- **57.** The rate-determining step in the acid catalysed dehydration of 3,3-dimethyl-2-butanol is

(a)
$$CH_3$$
 CH_3 CH_3

(b)
$$CH_3$$
 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3

(c)
$$CH_3$$

 $CH_3 - C - CH - CH_3 + H_2O \implies (CH_3)_2C = CHCH_3 + H_2O$
 CH_3

- (d) none of these
- **58.** Arrange H_2O , $HC \equiv CH$, NH_3 and CH_3CH_3 in order of acidity.
 - (a) $H_2O > HC \equiv CH > NH_3 > CH_3CH_3$
 - (b) $HC = CH > H_2O > NH_3 > CH_3CH_3$
 - (c) $CH_3CH_3 > HC = CH > NH_3 > H_2O$
 - (d) $H_2O > NH_3 > HC \equiv CH > CH_3CH_3$

59. In which of the following reactions are free-radical intermediates obtained?

(a)
$$CH_3CH=CH_2 + HBr \longrightarrow CH_3CHCH_3$$
Br

(b)
$$CH_3CH$$
— $CH_3 + H_2SO_4$ \longrightarrow CH_3CH = $CH_2 + H_2O$
OH

(c)
$$CH_2$$
= $CHCH_3 + Hg(OAc)_2 \longrightarrow CH_3$ — C — $CHCH_2OAc$

$$OH + CH_3COOH$$

(d)
$$CH_3CH=CH_2 + HBr \xrightarrow{R-O-O-R} CH_3CH_2CH_2Br$$

- 60. The Markovnikov addition of HCl to propene involves the
 - (a) initial attack of a chloride ion
 - (b) isomerization of 1-chloropropane
 - (c) formation of an n-propyl cation
 - (d) formation of an isopropyl cation
- 61. The product expected in the reaction

is

(b) HO OH

d) K MnO₄

- **62.** The reaction of 1-hexene with HBr in the presence of peroxides yields 1-bromohexane. The mechanism of the reaction involves the attack on the alkene by
 - (a) a Br⁺ ion

(b) a bromide ion

(c) a bromine atom

(d) an RO radical

- **63.** Which of the following methods of preparing alcohol proceed via anti-Markovnikov addition?
 - (a) Acid-catalysed hydration
- (b) Oxymercuration-demercuration
- (c) Hydroboration-oxidation
- (d) None of these

- **64.** The aldol condensation of (CH₃)₂CH—CH gives
 - (a) (CH₃)₂CHCH₂OH (b) CH₃ O | | | (CH₃)₂CHCH—C—CH | | OH—CH
 - (d) $O CH_3 O \\ || I || 1 \\ || CH_3|_2 CHC C CH_3 \\ || CH_3 || CH_3$ CH₃ O | | | (CH₃)₂C - C - CH
- 65. Phorone, $(CH_3)_2C=CH=C(CH_3)_2$, can be made through aldol condensation and the subsequent dehydration of
 - (a) 3 moles of acetone
 - (b) 3 moles of acetaldehyde
 - (c) 2 moles of acetone and HC-CH--C TO
 - (d) none of these
- **66.** The product of the following reaction

$$CH_3CH_2C \longrightarrow OH + CH_3^{18}OH \longrightarrow H^+$$

is

(a) O (b)
$${}^{18}O$$
 ${}^{18}O$ ${}^{18}O$

67. The final product (VI) obtained in the sequence of reactions

$$\begin{array}{c} CH_{3} \\ CH_{3}CHCOOH \xrightarrow{\quad LiAlH_{4} \quad } I \xrightarrow{\quad H_{2}O \quad } II \xrightarrow{\quad PBr_{3} \quad } III \\ \hline \frac{Mg}{Ether} IV \xrightarrow{\quad CO_{2} \quad } V \xrightarrow{\quad H_{2}O \quad } VI \end{array}$$

68. The final product (III) obtained in the sequence of reactions p-chlorotoluene $\xrightarrow{1. \text{KMnO}_4 \text{OH}^-} \text{I} \xrightarrow{\text{SOCl}_2} \text{II} \xrightarrow{\text{CH}_3\text{CH}_2\text{OH}} \text{III}$ is

69. Which of the following cannot be used for the synthesis of 2,2-dimethylpropionic acid?

(a)
$$CH_3$$
 $CH_3 - C - Br$
 $CH_3 - CH_3$
 C

(b)
$$CH_3$$
 $CH_3 - C - Br \xrightarrow{1. CN^-} CH_3 - CH_3 \xrightarrow{2. OH^-, H_2O}$

(c)
$$CH_3$$
 $CH_3 - CH_2 - CH_2OH \xrightarrow{1. \text{ KMnO}_4, \text{ OH}^-}$ $CH_3 - CH_3$

(d)
$$CH_3$$
 $CH_3 - C - CHO$ $1. \text{ KMnO}_4, OH^ CH_3 - CH_3$

70. The major product (III) obtained in the sequence of reactions

$$\begin{array}{c} CH_{3} \\ -CH_{3} \\ -CH_{2}CH_{2}COOH \\ \hline \begin{array}{c} 1 \text{ mole } Cl_{2} \\ \hline P \end{array} \end{array} I \xrightarrow{SOCl_{2}} II \xrightarrow{CH_{3}OH} III$$

is

OCH₃ (b) (CH₃)₂CHCHCH₂COCl

71. Which of the following is an example of a Hunsdieker reaction?

(a)
$$CH_2(CO_2H)_2 \xrightarrow{\Delta} CH_3CO_2H$$

(a)
$$CH_2(CO_2H)_2 \xrightarrow{\Delta} CH_3CO_2H$$

(b) $CH_3CCH_2CO_2H \xrightarrow{\Delta} CH_3CCH_3$
O

(c)
$$CH_3 - C - CH_3 \xrightarrow{KOH} CH_3CO_2^-K^+ + CHBr_3$$

(d)
$$CH_3CH_2COOH \xrightarrow{1. Ag_2O} CH_3CH_2Br$$

- 72. In nucleophilic substitutions, the relative reactivity of acyl compounds is
 - (a) acyl chloride > ester > acid anhydride > amide
 - (b) acid anhydride > acyl chloride > ester > amide
 - (c) acyl chloride > acid anhydride > ester > amide
 - (d) ester > acyl chloride > acid anhydride > amide

73. The final product III obtained in the reaction

$$\begin{array}{c|c}
CH_3 \\
\hline
Sn + HC1 \\
\hline
NO_2
\end{array}$$

$$I \xrightarrow{Br_2/H_2O} II \xrightarrow{1. HONO} III$$

is

(a) CH₃
Br

(b) CH₃

74. What will be the major product in the reaction

$$CH_3 CH_2COC_2H_5 \xrightarrow{1. \text{ NaOEt}}$$

- (a) CH₃CH₂COCH₂CH₂CO₂C₂H₅
- (b) CH₃CH₂CO₂CH₂CH₂CO₂C₂H₅
- (c) CH₃CH₂COCHCO₂C₂H

- 75. The hybridization state of a carbocation is
 - (a) sp 4

(b) sp^3

(c) sp^2

- (d) sp
- **76.** How many electrons are present in the p orbital of a methyl cation (CH_3^+) ?
 - (a) Two
- (b) Three
- (c) Four
- (d) None

77. The reaction of benzaldehyde with HCN to give benzoin takes place in the following steps.

(a)
$$C_6H_5$$
 C_6H_5 C_6H_5 C_6H_5 C_6H_5 C_6H_5

(b)
$$O_{-}^{\odot}$$
 O_{-}^{\odot} O_{-}^{O} O_{-}^{O}

(d)
$$\bigcirc$$
 OH O \bigcirc \bigcirc C₆H₅-C-C-C₆H₅ \longrightarrow C₆H₅CCHOHC₆H₅ \bigcirc CN H

Which of these is the rate-determining step?

78. In the acid-catalysed hydration of an alkene, the rate-determining step is

(a)
$$CH_2$$
 CH_3 $CH_$

(b)
$$CH_3$$
 CH_3 CH_3

(c)
$$CH_3 H$$
 $CH_3 - CH_3 + CH_3 - CH_3 - CH_3 + CH_3 - CH_3 + CH_3 - CH_3 + C$

- (d) none of these
- 79. The major product obtained in the reaction

$$CH_{3} - CH_{2} - CH_{2} \longrightarrow CH_{3}$$

$$CH_{3} - CH_{2} - CH_{2} \longrightarrow CH_{3}$$

is

- (d) a mixture of (a) and (b) in equal proportions
- **80.** What is the order of reactivity of the alkenes (CH₃)₂C=CH₂(I), CH₃CH=CH₂(II) and CH₂=CH₂(III) when subjected to acid-catalysed hydration?
 - (a) I > II > III

(b) I > III > II

(c) III > II > I

(d) II > I > III

• Type 2 •

Choose the correct options. More than one option is correct.

- **81.** Which of the following groups are *o* and *p*-directing?
 - (a) $-NH_2$

(b) —NHCOCH₃

(c) —CN

(d) $-SO_3H$

- **82.** Which of the following groups are *m*-directing?
 - (a) —CHO

(b) —OH

(c) —OCOCH₃

- (d) -COOH
- 83. Which of the following are nucleophiles?
 - (a) Water

- (b) Ammonia
- (c) Triphenylsulphide
- (d) Iodides
- 84. Which of the following are electrophiles?
 - (a) Dimethyl sulphide
- (b) Bromides
- (c) Carbon dioxide
- (d) Ammonia
- **85.** Which of the following arrangements show a correct order of nucleophilicity?
 - (a) $I^- > Br^- > Cl^- > F^-$
- (b) $RS^- > RO^-$

(c) $R_3N: > R_3P:$

- (d) $RO^- > RS^-$
- **86.** Which of the following are electrophiles?
 - (a) BF_3
- (b) Cl₂C:
- (c) NR_4^+
- (d) I⁻
- **87.** Which of the following are aprotic solvents?
 - (a) DMSO
- (b) DMF
- (c) H₂O
- (d) CH₃COOH
- 88. What is the product expected in the following reaction?

(a)

NH₂

(b) CH₃

(c)

CH₃

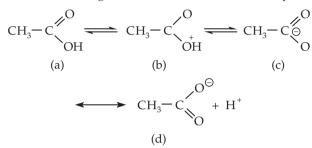
(d) none of these

89. Which of the following statements are correct for butadiene

$$^{4}_{\text{CH}_{2}} = ^{3}_{\text{CH-CH}} = ^{2}_{\text{CH}_{2}}$$

(a) The C₁−C₂ and C₃−C₄ bonds are longer than a carbon-carbon double bond.

- (b) The C_1 – C_2 and C_3 – C_4 bonds are shorter than a carbon-carbon double bond.
- (c) The C₂–C₃ bond is slightly shorter than a carbon-carbon single bond.
- (d) The C₂-C₃ bond is slightly longer than a carbon-carbon double bond.
- 90. Which of the following structures have resonance stability?



91. Which of the following groups have a +M effect (an electron-attracting mesomeric effect)?

(a)
$$-$$
NH₂

92. Which of the following groups have a –M effect (an electron-repelling mesomeric effect)?

(d)
$$-SO_3H$$

93. Br has a low reactivity in CH₂=CH—Br because

- (a) the C–Br bond has a partial double-bond character
- (b) of the +M effect of bromine
- (c) Br is electronegative
- (d) of none of these

94. Which of the following statements are correct?

- (a) Under normal conditions, 2,6-dimethylbenzoic acid cannot be esterified with ethyl alcohol and concentrated sulphuric acid.
- (b) 2,6-Dimethylphenylacetic acid can be esterified with ethyl alcohol and concentrated sulphuric acid.
- (c) The nitration of tertiary butyl benzene gives a *p*-nitro product.

- (d) 2,6-Dimethyl-N,N-dimethylaniline does not undergo azo coupling.
- **95.** Which of the following statements are correct?
 - The nucleophilic addition of HCN to CH₃CH=CHCHO results in a major addition to C=C.
 - The nucleophilic addition of HCN to CH₃CH=CHCHO results in a major addition to C=O.
 - (c) The nucleophilic addition of PhMgBr to PhCH=CH-CHO results in addition to C=0.
 - (d) The nucleophilic addition of PhMgBr to PhCH=CH-COCMe₃ results in addition to C=C.
- **96.** On treatment with strong NaOH at 340°C, p-chlorotoluene gives
 - (a) o-cresol

(b) *m*-cresol

(c) p-cresol

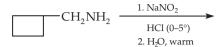
- (d) none of these
- 97. Under S_N1 conditions, the hydrolysis of neopentyl bromide,

gives

- (c) CH_3 —C=CH— CH_3 (d) none of these CH_3
- 98. Which of the following statements are correct?
 - (a) The addition of HBr to propene gives 2-bromopropane.
 - (b) The addition of HBr to propene gives 1-bromopropane.
 - (c) The addition of HCl to vinyl chloride gives ethylidene chloride.
 - (d) The addition of HCl to vinyl chloride gives ethylene chloride.
- 99. The addition of bromine to butadiene gives

 - (a) CH₂Br-CHBr-CH=CH₂ (b) CH₂Br-CH=CH-CH₂Br
 - (c) CH₃-CHBr-CHBr-CH₃ (d) CH₃-CH₂CH₂CHBr₂

100. In the reaction



the product obtained is





101. The oxidation in air of cumene (isopropyl benzene) followed by hydrolysis yields

(c)
$$O$$
 \parallel CH_3-C-CH_3

(d) none of these

102. The reaction

gives

(a)
$$H_2N$$
 NH_2

(b)
$$H_2N$$

(c)
$$NH_2$$
 H_2N

- (d) none of these
- 103. The reaction of acetone with dry HCl gas gives
 - (a) (CH₃)₂C=CHCOCH₃

- (c) (CH₃)₂C=CH—COCH=C(CH₃)₂
- (d) none of these
- **104.** The Cannizzaro reaction of benzaldehyde and formaldehyde in the presence of NaOH gives
 - (a) C₆H₅CH₂OH

(b) C₆H₅COONa

(c) HCOONa

- (d) CH₃OH
- 105. Which of the following will undergo a Cannizzaro reaction?
 - (a) (CH₃)₂CHCHO

$$(CH_3)_2N$$
 — CHO

(d) OHC—COOH

106. In the reaction

$$\begin{array}{ccc} \text{CH}_3\text{CH=CH}_2 & \xrightarrow{\text{HX}} & \text{CH}_3\text{CH}_2\text{CH}_2X \\ & \xrightarrow{\text{benzoyl}} & \text{(A)} \end{array}$$

the anti-Markovnikov product (A) cannot be obtained by using

- (a) HBr
- (b) HCl
- (c) HI
- (d) none of these

107. In the reaction

$$\begin{array}{ccc} & O & O \\ \parallel & \parallel \\ C_6H_5C-C-H & & \longrightarrow \end{array}$$

which of the following are not obtained?

(a) O (b) O
$$\parallel$$
 $C_6H_5-C-COONa$ $C_6H_5-C-CH_2OH$

108. The reaction

$$\begin{array}{c} O \\ \parallel \\ R - C - NH_2 \end{array} \xrightarrow{\qquad NaOBr(Br_2 + NaOH)} RNH_2$$

has the intermediates

(a)
$$O \\ R-C-NHBr$$
 (b) $O \\ R-C-N$:
(c) $R-N=C=O$ (d) $R-NH-C=O$

109. The products obtained in the reaction

are

- 110. Which of the following is an example of nucleophilic addition to acetone?
 - (a) Ketal formation
- (b) Reduction with hydrogen gas
- (c) Cyanohydrin formation
- (d) Bisulphite addition

- 111. Which of the following species bear a positive charge?
 - (a) H (b) H H N: N: H
 - (c) H:N:H H:O:H
- 112. Which of the following has a trigonal planar (or triangular) structure
 - (a) $: CH_3$ (b) CH_3
 - (c) BF_3 (d) $:OH_3$
- 113. Which of the following statements are true for the S_N1 reactions of alkyl halides?
 - (a) The rate of an $S_N 1$ reaction depends on the concentration of the alkyl halide.
 - (b) The S_N 1 reations of alkyl halides are favoured by polar solvents.
 - (c) The rate of an S_N1 reaction depends on the concentration of the nucleophile.
 - (d) The rate of an $S_N 1$ reaction depends on the concentration of the substrate as well as that of the nucleophile.
- **114.** In which of the following reactions is there a possibility of rearrangement?
 - (a) S_N1 reactions

- (b) $S_N 2$ reactions
- (c) E1 reactions
- (d) E2 reactions
- 115. Which of the following are examples of electrophilic addition?

(a) OH
$$CH_3CH=CH_3 + H_2O \longrightarrow CH_3-CH-CH_3$$

(b)
$$CH_3CH=CH_3 + HBr \xrightarrow{Peroxides} CH_3CHCH_3 Br$$

(c)
$$CH_3CH=CHCH_3 + Cl_2 \longrightarrow CH_3CH-CH-CH_3$$

 Cl Cl

- (d) None of these
- 116. Which of the following statements are true for the organoborane

- (a) It can be thermally rearranged.
- (b) It can be oxidized to 3-pentanol.
- (c) It can be converted to pentane.
- (d) None of these
- 117. Which of the following are intermediates in the Hofmann degradation reaction?R—N=C=O

118. The intermediates involved in the Curtius reaction

$$RCON_3 \longrightarrow RNH_2$$

are

(a)
$$O$$
 (b) $R-\ddot{N}=C=C$ $R-C-\ddot{N}$:

(c) R—C≡N

(d) none of these

119. In the reaction

$$ROH + HCl \xrightarrow{ZnCl_2} RCl + H_2O$$

the intermediates are

(a) R+

(b) Cl

(c)
$$+$$
 (d) none of these $R-O-ZnCl_2$

- 120. Which of the following statements are correct?
 - (a) RO is a stronger nucleophile than OH.
 - (b) RCO₂ is a stronger nucleophile than OH -.
 - (c) RCO₂ is a stronger nucleophile than ROH.
 - (d) RO^- is a weaker nucleophile than OH^- .

121. Which of the following are polar aprotic solvents?

- (a) Dimethylsulfoxide
- (b) Hexamethylphosphoramide

(c) Acetone

(d) N,N-Dimethylformamide

122. In the reaction

$$\begin{array}{c|c}
R \\
-C - C - OH \\
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which of the following steps are not rate-determining?

$$\begin{array}{c} \text{(b)} & \text{R} & \text{H} \\ -\text{C} - \text{C} - \text{O} - \text{H} & \text{C} - \text{C} - \text{C} \oplus & \text{+ O} - \text{H} \\ \text{H} & \text{R'} & \text{H} & \text{R'} \end{array}$$

(d) None of these

123. In the reaction

$$\begin{array}{c} \text{CH}_3 \\ \text{C} \\ \text{H} \\ \text{CH}_3 \end{array} \xrightarrow{\text{Br}_2} \begin{array}{c} \text{Br}_2 \\ \text{CCl}_4 \end{array}$$

the products obtained are

(a)
$$CH$$
 EH_3 EH_3 EH_4 EH_5 EH_5

(c)
$$\underset{\longrightarrow}{\overset{C}{C}}H_3$$
 (d) none of these $\underset{\longrightarrow}{\overset{C}{C}}H$ $\underset{\longrightarrow}{\overset{C}{C}}H$

Answers					
1. a	2. b	3. d	4. a	5. c	
6. a	7. d	8. a	9. c	10. a	
11. c	12. d	13. a	14. c	15. a	
16. b	17. b	18. b	19. c	20. d	
21. b	22. c	23. a	24. b	25. a	
26. c	27. c	28. a	29. b	30. b	
31. b	32. a	33. c	34. d	35. a	
36. a	37. c	38. c	39. a	40. b	
41. d	42. c	43. b	44. b	45. c	
46. b	47. a	48. d	49. a	50. a	
51. b	52. c	53. d	54. d	55. c	
56. a	57. a	58. a	59. d	60. d	
61. a	62. c	63. c	64. b	65. a	
66. a	67. d	68. b	69. c	70. a	
71. d	72. c	73. b	74. c	75. c	
76. d	77. c	78. a	79. c	80. a	
81. a, b	82. a, d	83. a, b, c, d	84. a, c	85. a, b	
86. a, b, c	87. a, b	88. a, b	89. a, c	90. c, d	
91. a, b	92. c, d	93. a, b	94. a, b, c, d	95. a, c, d	
96. b, c	97. a, c	98. a, c	99. a, b	100. a, b, c, d	
101. a, c	102. a, b	103. a, c	104. a, c	105. a, c, d	
106. b, c	107. a, b, d	108. a, b, c, d	109. a, b	110. a, c, d	
111. c, d	112. b, c	113. a, b	114. a, c	115. a, b, c	
116. a, b, c	117. a, b, c	118. a, b	119. a, b, c	120. a, c	
121. a, b, d	122. a, c	123. b, c			

Hints to More Difficult Problems

16. Hyperconjugation satisfactorily explains the preferential formation of the product (b).

- 17. The formation of the product (b) is due to the conjugation of the double bond with the carbonyl group, which gives resonance stability.
- **22.** The formation of the abnormal product (a) is due to the rearrangement of the secondary carbocation to the tertiary carbocation for stability.
- **29.** The unusual product (b) is obtained contrary to the Saytzev rule due to steric hindrance.
- **32.** This is a case of pinacol-pinacolone rearrangement.
- **33.** This is case of the rearrangement of the alicyclic ring system (Demjanov rearrangement).
- **35.** This is a case of Claisen rearrangement.
- **36.** End-interchange occurs twice during para migration, and so there is no rearrangement in the final product.
- **44.** Product (b) is obtained due to the strong electron-withdrawing effect of the CN group, rendering the secondary carbocation less stable than the primary carbocation. Also, the carbocation is separated from CN by two carbon atoms, and the destabilization by the inductive effect is less.
- 89. Consider the following resonance structure

$$CH_{2}=CH-CH=CH_{2} \longleftrightarrow CH_{2}-CH-CH=CH_{2}$$

$$\longleftrightarrow CH_{2}=CH-CH-CH_{2} \longleftrightarrow CH_{2}-CH=CH-CH_{2}$$

$$\longleftrightarrow CH_{2}=CH-CH-CH_{2} \longleftrightarrow CH_{2}-CH=CH-CH_{2}$$

$$\longleftrightarrow CH_{2}-CH=CH-CH_{2}$$

$$(5)$$

The structures (2) and (3) show the partial single-bond character of the C_1 – C_2 and C_3 – C_4 bonds, and (4) and (5) the partial double-bond character of the C_2 – C_3 bond. This explains the observed anomalies in the bond distances.

- **90.** Charge separation structures are less important than those in which the charge is delocalized, because there is electrostatic attraction between unlike charges.
- **93.** The low reactivity of a halogen bonded to an unsaturated carbon is due to the +M effect of the halogen. The C–Br bond in vinyl chloride has a partial double-bond character due to the +M effect of bromine, resulting in low reactivity.
- **94.** The steric effect plays a role in all these reactions.
- **95.** The formation of the products in (a) can be explained by the fact that C=O is a stronger bond than C=C, and so the preferential addition is to

- C=C. In the formation of the product in (c), steric hindrance is a major factor. However, in case of (d), the inductive effect of the Me group is a major factor, in addition to steric hindrance.
- 97. The formation of the unexpected product (a) is due to the rearrangement of the first formed $\mathring{1}$ carbocation to the $\mathring{3}$ carbocation. This also explains the formation of the product (c).
- 107. The product (c) is obtained due to an intramolecular Cannizzaro reaction.

Hydrocarbons

• *Type 1* •

1. Using which of the following reagents can one perform a simple test that can be used to differentiate between $C_6H_5C \equiv CH$ and $C_6H_5CH = CH_2$?

(b) Br₂/CCl₄

Choose the correct option. Only one option is correct.

(a) NaOH/H₂O

	(c)	Ag(NH ₃) ₂ OH	(d)	CrO ₃ /H ₂ SO ₄
2.	The ac	ddition of HCl to 1-phenylprop	ene	gives
	(a)	C ₆ H ₅ CHClCH ₂ CH ₃	(b)	C ₆ H ₅ CH ₂ CHClCH ₃
	(c)	C ₆ H ₅ CH ₂ CH ₂ CH ₂ Cl	(d)	C ₆ H ₅ CH(CH ₃)CH ₂ Cl
3.	The a	ddition of HBr in the presence	e of	a peroxide to 1-phenylpropene
	(a)	C ₆ H ₅ CHBrCH ₂ CH ₃	(b)	C ₆ H ₅ CH ₂ CHBrCH ₃
	(c)	C ₆ H ₅ CH(CH ₃)CH ₂ Br	(d)	C ₆ H ₅ CH ₂ CH ₂ CH ₂ Br
4.	By usi	ing which of the following can $PhC \equiv CH \longrightarrow PhC \equiv$		
	be ach	nieved?		
	(a)	Br ₂ /CCl ₄ , then KOH	(b)	Na, then CH ₃ CH ₂ I
	(c)	Na, then CH ₃ I	(d)	CH_2N_2
5.	Which	n conformation of ethane has th	ne lo	west potential energy?
	(a)	Eclipsed	(b)	Skewed
	(c)	Staggered	(d)	All will have equal PE
6.	The re	elative energies of the ethane co	onfo	rmations are in the order
	(a)	skewed < eclipsed < staggered	d	
	(b)	staggered < eclipsed < skewed	d	
		3-53		

- (c) skewed < staggered < eclipsed
- (d) staggered < skewed < eclipsed
- 7. The name of the compound

is

- (a) 2-methyl-5-chlorocyclohexanol
- (b) 3-chloro-6-methylcyclohexanol
- (c) 2-hydroxy-4-chlorocyclohexane
- (d) 3-hydroxy-4-methylchlorocyclohexane
- 8. The name of the compound

is

- (a) 2,5-dimethylchlorocyclohexane
- (b) 3,6-dimethylchlorocyclohexane
- (c) 3-chloro-1,4-xylene
- (d) none of these
- 9. Which cycloalkane has the lowest heat of combustion per CH₂ group?









10. The most stable conformation of 1,2-diphenylethane is

(a)
$$H$$
 (b) H C_6H_5 C_6H_5 H (c) H C_6H_5 C_6H_5 C_6H_5 C_6H_5 C_6H_5 C_6H_5 C_6H_5 C_6H_5

- **11.** The reaction of RCH₂CHX₂ with alcoholic KOH followed by treatment with NaNH₂ gives
 - (a) only RCH=CHX

Н

(b) only RC≡CH

H

(c) a mixture of (a) and (b)

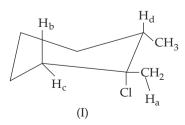
Н

- (d) none of these
- 12. A compound C_6H_{10} decolourizes bromine in carbon tetrachloride and reacts with Ag^+ in ammonia to give an insoluble salt. On treatment with excess hydrogen in the presence of an Ni catalyst, this compound gives 2-methylpentane. The structure of the compound C_6H_{10} is

(a)
$$CH_3C = CH - CH = CH_2$$
 (b) $CH_2 = C - CH = CHCH_3$ CH_3

(c)
$$CH_3CHC \equiv CCH_3$$
 (d) H $CH_3 - C - CH_2C \equiv CH_3$ CH_3

13. In the following halogen-substituted hydrocarbon, the hydrogen atom that can be eliminated most readily is



- (a) H_a
- (b) H_b
- (c) H_c
- (d) H_d

14. On catalytic hydrogenation, an organic compound $X(C_7H_{12})$ absorbs 1 mol of hydrogen and yields a compound C_7H_{14} . On ozonolysis and subsequent treatment with Zn/H_2O , the compound

$$\overset{\text{O}}{\underset{\text{II}}{\parallel}} \overset{\text{O}}{\underset{\text{CH}_3\text{C}-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2}{\overset{\text{O}}{\underset{\text{II}}{\parallel}}}$$

is obtained. The structure of X is

(a) CH
$$_3$$
—C—CH $_2$ CH $_2$ CH=CH $_2$ (b) CH $_3$ CHC=CCH $_2$ CH $_3$ CH $_3$

15. The molecules

$$CH_3$$
 $C=C=C$
 CH_3
 CH_3
 $C=C=C$
 CH_3
 CH_3
 $C=C=C$
 CH_3

are

(a) enantiomers

- (b) diastereomers
- (c) structural isomers
- (d) none of these
- **16.** Which of the following statements is true for ethane, ethene and acetylene?
 - (a) Acetylene is the weakest acid and has the longest C—H bond distance.
 - (b) Acetylene is the strongest acid and has the shortest C—H bond distance.
 - (c) Ethane is the strongest acid and has the longest C—H bond distance.
 - (d) Ethene is the strongest acid and has the shortest C—H bond distance.
- **17.** On oxidation with hot aqueous KMnO₄, a compound X gives 2-methylpropionic acid and propionic acid. Which of the following is the structure of X?

(c)
$$CH_3CHC \equiv CCH_2CH_3$$

 CH_3

- 18. On catalytic hydrogenation, a compound X (C_7H_{12}) absorbs 2 mol of hydrogen and yields 2-methylhexane. On treatment with Ag(NH₃)₂OH, X gives a precipitate which contains silver and which regenerates X on treatment with dilute HNO₃. The structure of X is
 - (a) $CH_3CH_2CH_2CH_2CH_2CH = CH$ (b) $CH_3C = CCH_2CH CH_3$ CH_3
 - (c) CH_2 =CHCH= $CHCHCH_3$ (d) CH_3 - $CHCH_2CH_2C$ =CH CH_3 CH_3
- 19. Among the following dienes, which is the most stable?
 - (a) CH₃CH=CHCH₂CH=CH₂ (b) CH₃CH=CHCH=CHCH₃
 - (c) CH_2 = $CHCH_2CH$ = CH_2 (d) CH_2 =CHCHCH=CH
 - CH₃
- 20. The main product obtained in the reaction of 2-butene with N-bromosuccinimide is
 - (a) CH₂BrCHBrCH₂CH₃
- (b) CH₃CHBrCH₂CH₃
- (c) CH₃CH=CBrCH₃
- (d) CH₃CH=CHCH₂Br
- **21.** Among the following, which has the shortest carbon-carbon single bond?
 - (a) CH₃-CH₃

- (b) CH₂=CH-CH₃
- (c) HC≡C-C≡CH
- (d) CH₂=CH-C=CH
- **22.** The product obtained in the reaction

$$CH_3$$
 $C=C$
 CH_3
 $CHCl_3$
 $CHCl_3$

(c)
$$CH_3$$
 H (d) none of these CH_3 CH_3 CI CI

- 23. Which is the most stable conformation of cyclohexane?
 - (a) Chair
- (b) Twist
- (c) Boat
- (d) Staggered
- **24.** Which of the following reagents can be used in a simple chemical test to distinguish CH₂BrCH=CH₂ from CH₃CH=CHBr?
 - (a) $Ag(NH_3)_2OH$

(b) Br_2/CCl_4

(c) Cold KMnO₄

- (d) AgNO₃/C₂H₅OH
- 25. Which alkene will undergo the following reaction?

$$\begin{array}{c} \text{Alkene} \xrightarrow{\begin{array}{c} 1.\,\,\mathrm{O_3} \\ \\ 2.\,\,\mathrm{Zn},\,\mathrm{H_2O} \end{array}} \\ \text{H-C-CH}_2 \\ \begin{array}{c} \mathrm{CH}_2 \\ \\ \mathrm{CH}_3 \end{array} \\ \text{CH}_3 \end{array} \\ + \\ \text{H-C-H}$$

(a) H
$$C=C$$
 $CH_2CH_2CH_2CH_3$

(b)
$$H$$
 $C=C$
 $CHCH_2CH_3$
 CH_3

(c) H
$$C=C$$
 H $CH_2-CH-CH_3$ CH_2

(d)
$$CH_3$$
 $C=C$ H $CH_2CH_2CH_3$

- 26. The secondary butyl group is represented as
 - (a) CH₃CH₂CH₂CH₂—

(c)
$$\mid$$
 CH₃ CH $_3$ CH $_3$ CH $_3$ CH $_3$ CH $_3$ CH $_4$ CH $_3$ CH $_4$ CH $_5$ C

27. The product obtained in the reaction

$$H_3C-C \equiv CH \xrightarrow{alc. KOH} 150^{\circ}C, 6 \text{ hr}$$

is

- (a) CH₃CH=CHOH
- (b) CH₂C=CH₂ OH

(c) $CH_2=C=CH_2$

- (d) none of these
- **28.** On ozonolysis, a compound C_7H_{12} gives acetone and carbon dioxide. The structure of the compound can be
 - (a) $CH_3CH_2C\equiv C-CH_2CH_2CH_3$ (b) CH_3 C=C=C CH_3
 - (c) $CH_3C \equiv C CH_2CH CH_3$ (d) none of these CH_3
- 29. The polymerization of butadiene according to the reaction

$$\begin{array}{c} \text{CH}_2 \\ \text{CH} \\ \text{CH} \\ \text{CH}_2 \\ \end{array} \begin{array}{c} \text{polymerization} \\ \text{CH}_2 \\ \end{array} \begin{array}{c} \text{(-H}_2\text{C} \\ \text{H} \\ \text{CH}_{\overline{2}}\text{)}_2 \\ \end{array}$$

is an example of

(a) 1,2 addition

- (b) cis-1,4 addition
- (c) trans-1,4 addition
- (d) none of these
- 30. The reaction of ethyne with bromine water gives
 - (a) cis-1,2-dibromoethene
- (b) trans-1,2-dibromoethene
- (c) 1,1,2,2-tetrabromoethane
- (d) none of these
- **31.** The order of the bond strength of C—H bonds involving sp-, sp ²- and sp ³-hybridized carbon atoms is
 - (a) $sp > sp^2 > sp^3$

(b) $sp^3 > sp^2 > sp$

(c) $sp^2 > sp^3 > sp$

(d) $sp^2 > sp > sp^3$

32. The group

$$CH_{3} - C - CH_{2} - CH_{3} - CH_{3}$$

is called the

(a) butyl group

- (b) secondary butyl group
- (c) tertiary butyl group
- (d) neopentyl group
- 33. In the oxidation of alkenes with a dilute KMnO₄ solution followed by acidification with dilute H₂SO₄ to give diol,

$$\begin{bmatrix} C \\ \parallel \\ C \end{bmatrix} + KMnO_4 \xrightarrow{OH^+} \begin{bmatrix} \\ \end{bmatrix} \xrightarrow{dil H_2SO_4} \begin{bmatrix} C - OH \\ C - OH \end{bmatrix}$$

the OH groups come from

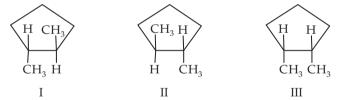
(a) NaOH

(b) H₂O

(c) KMnO₄

- (d) H₂SO₄
- 34. Which of the following hydrocarbons cannot be oxidized by KMnO₄/alkali?
 - (a) Toluene

- (b) Ethylbenzene
- (c) Isopropylbenzene
- (d) Tertiary butylbenzene
- 35. 1,2-Dimethylcyclopentane can be represented as I, II and III.



Which of these structures are enantiomers?

(a) I and II

(b) I and III

(c) II and III

- (d) I, II and III
- **36.** The final product obtained in the reaction

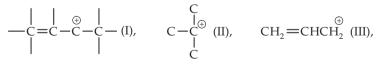
$$CH_3CH_2CH = CH_2 \xrightarrow{1 \cdot Br_2/CCl_4} \xrightarrow{2 \cdot NaNH_2, 110-160^{\circ}C}$$

is

- (a) $CH_3CH_2CH=CHBr$ (b) $CH_3CH_2C=CH_2$
- (c) CH₃CH₂C≡CH
- (d) none of these

- 37. The basicities of CH₃CH₂: ¬, CH₂=CH: ¬ and HC≡C: ¬ are of the order
 - (a) $CH_2CH_2:^->CH_2=CH:^->HC=C:^-$
 - (b) HC=C: -> CH2=CH: -> CH2CH2: -
 - (c) CH₂=CH: -> CH₂CH₂: -> HC=C:
 - (d) All the three are equally basic.
- 38. The acidities of H₂O, CH₃OH, CH₃—C=CH, NH₃, CH₂=CH₂ and CH₃CH₃ are of the order
 - (a) $H_2O > CH_2OH > CH_2C = CH > NH_2 > CH_2 = CH_2 > CH_2 = CH_2$
 - (b) $CH_3OH > H_2O > NH_3 > CH_3C = CH > CH_3 CH_3 > CH_2 = CH_2$
 - (c) $CH_3 CH_3 > CH_2 = CH_2 > NH_3 > CH_3C = CH > CH_3OH > H_2O$
 - (d) $NH_3 > CH_3C \equiv CH > CH_3 CH_3 > CH_2 = CH_2 > CH_3OH > H_2O$
- 39. The reduction of 4-octyne with H₂ in the presence of Pd/CaCO₃quinoline gives
 - (a) trans-4-octene
 - (b) cis-4-octene
 - (c) a mixture of cis- and trans-4-octene
 - (d) a completely reduced product C₈H₁₈
- 40. On halogenation, an alkane (C₅H₁₂) gives only one monohalogenated product. The alkane is
 - (a) *n*-pentane

- (b) 2-methylbutane
- (c) 2,2-dimethylpropane
- (d) cyclopentane
- 41. Which of the following free radicals is the most stable?
 - (a) Vinylic
- (b) Allylic
- (c) Ĭ
- (d) 2
- 42. The chlorination of propene at 400°C in the gaseous phase gives
 - (a) CICH₂—CH—CH₃
 Cl
- (b) CH₃—CH—CH₃
 Cl
 (d) ClCH₂—CH₂—C
 - (c) CH₂=CH-CH₂Cl
- (d) ClCH2-CH2-CH3
- **43.** The order of stability of the carbocations



$$\begin{array}{cccc} C & H & & & \\ -C & & |_{\oplus} \\ C - C & & |_{\oplus} \\ I & & H & & H \end{array} \text{ and } CH_2 \stackrel{\oplus}{=} CH \text{ (VI)}$$

is

- (a) VI > V > IV > III > II > I
- (b) I > II > III > VI > V > IV
- (c) V > VI > III > IV > I > II
- (d) I > II > III > IV > V > VI
- **44.** Arrange the compounds

CH2=CH-CH3 (I), CH2=CH-CH=CH2 (II),

 $HC \equiv C - CH = CH_2$ (III) and $HC \equiv C - C \equiv CH$ (IV)

in order of increasing carbon-carbon single-bond length.

(a) I > II > III > IV

(b) II > I > IV > III

(c) IV > III > II > I

- (d) I > II > IV > III
- **45.** In which of the following compounds is the hybridization state of the atoms in the carbon-carbon single bonds sp-sp?
 - (a) HC≡C—C≡CH
- (b) HC≡C—CH=CH₂
- (c) HC≡C—CH₃

- (d) $CH_2 = CH CH = CH_2$
- **46.** The angle strains at the CH_2 groups in cyclopropane (I), cyclobutane (II) , cyclopentane (III) and cyclohexane (IV) follow the order
 - (a) IV > III > II > I
- (b) I > II > III > IV

(c) II > I > IV > III

- (d) III > II > IV
- 47. The product obtained in the reaction

$$CF_3$$
— CH = CH_2 + HCl \longrightarrow

is

- (a) CF₃CH₂—CH₂Cl
- (b) CF₃CHClCH₃
- (c) CF₃CHCl—CH₂Cl
- (d) none of these
- **48.** The order of reactivity of F, Cl, Br and I in substitution reactions is
 - (a) F > Cl > Br > I

(b) Cl > Br > I > F

(c) I > Br > Cl > F

- (d) Cl = Br > I > F
- 49. Which of the following compounds has the highest boiling point?
 - (a) n-Octane

(b) Iso-octane

(c) *n*-Butane

- (d) 2,2,3,3-Tetramethylbutane
- 50. Which of the following conditions gives the best yield of ethyl chloride?

(a)
$$C_2H_6$$
 (excess) + Cl_2 uv light

(b)
$$C_2H_6 + Cl_2$$
 (excess) — uv light

(c)
$$C_2H_6 + Cl_2 \xrightarrow{\text{uv light}}$$

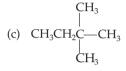
(d)
$$C_2H_6 + Cl_2 - \frac{\text{darkness}}{\text{room temp.}}$$

- 51. Which of the following compounds will have zero dipole moment?
 - (a) *cis-*1,2-Dichloroethylene
- (b) trans-1,2-Dichloroethylene

	(c)	1,1-Dichloroethylene	(d)	None of these
52.	On tr	eatment with alcoholic potas	sium	hydroxide, <i>n</i> -propyl bromide
	(a)	propane	(b)	propene
	(c)	propyne	(d)	propanol
53.	How	many types of carbon atoms ar	e pre	esent in 2,2,3-trimethylpentane?
		One	•	Two
	(c)	Three	(d)	Four
54.	In eth	ane, the H—C—C bond angle	is	
	(a)	109.5° (b) 109°		180° (d) 120°
55.	The sh	nape of the methane molecule i	is	
	(a)	trigonal planar	(b)	square planar
	(c)	tetrahedral	(d)	linear
56.	Metha	ane as well as ethane can be ob	taine	ed in one step from
	(a)	CH ₃ OH (b) C ₂ H ₅ OH	(c)	CH_3I (d) C_2H_2
57.	The re	eaction of methyl magnesium b	rom	ide with ethyl alcohol gives
		methyl alcohol		methane
	(c)	isopropyl alcohol	(d)	ethane
58.		reparation of an alkane from ar n as the	n alk	yl halide by reaction with zinc is
	(a)	Wurtz reaction	(b)	Cannizzaro reaction
	(c)	Frackland reaction	(d)	Kolbe reaction
59.		poiling point of a branched ht-chain isomer is	alka	ne as compared to that of a
	~	high	(b)	low
	(c)	equal	(d)	independent of branching
60.	The o	rder of reactivity of the hydrog	ens i	n isopentane is
	(a)	$\hat{1} > \hat{2} > \hat{3}$ (b) $\hat{1} = \hat{2} > \hat{3}$	(c)	$\mathring{3} > \mathring{2} > \mathring{1}$ (d) $\mathring{3} > \mathring{1} > \mathring{2}$
61.	The h	alogenation of alkanes is an ex	ampl	le of
	(a)	nucleophilic substitution		electrophilic substitution
	(c)	oxidation	(d)	free-radical substitution
62.	Which produ	_	give	es only one monochlorinated
	(a)	CH ₃ CH ₂ CH ₃	(b)	CH ₃ CH ₂ CH ₂ CH ₃
	(c)	$(CH_3)_4C$	(d)	$CH_3(CH_2)_3CH_3$

- **63.** A liquid hydrocarbon is converted to a mixture of gaseous hydrocarbons by
 - (a) hydrolysis

- (b) oxidation
- (c) distillation in vacuum
- (d) cracking
- 64. Which of the following compounds has the lowest boiling point?
 - (a) CH₃CH₂CH₂CH₂CH₂CH₃
- (b) CH₃CH₂CH—CH₂CH₃ CH₂



- (d) CH₃—CH—CH—CH₃ | | CH₃ CH₃
- **65.** Which of the following is the isobutyl group?
 - (a) CH₃CH₂CH₂CH₂—
- (b) CH₃—CH₂—CH—CH₃

- **66.** Which of the following compounds has three $\mathring{1}$, one $\mathring{2}$ and one $\mathring{3}$ carbon atom?
 - (a) $CH_3(CH_2)_2CH_3$
- (b) CH₃CH(CH₃)CH₃
- (c) $CH_3(CH_2)_3CH_3$
- (d) CH₃CH(CH₃)CH₂CH₃
- 67. The ease of formation of free radicals follows the order
 - (a) $3 > 2 > 1 > CH_3$
- (b) $\dot{C}H_3 > \dot{1} > \dot{2} > \dot{3}$
- (c) $\hat{1} > \hat{2} > \hat{3} > \hat{C}H_3$
- (d) $\mathring{2} > \mathring{1} > \mathring{3} > \mathring{C}H_3$
- **68.** Which of the following is correct regarding the stability of carbocations?
 - (a) 3 > 2 > 1

(b) 3 < 2 < 1

(c) $\hat{2} > \hat{1} > \hat{3}$

- (d) $\overset{\circ}{2} > \overset{\circ}{3} > \overset{\circ}{1}$
- 69. The shape of the ethylene molecule is
 - (a) linear

(b) tetrahedral

(c) pyramidal

- (d) planar
- 70. The bond angle H—C—H of alkenes is equal to
 - (a) 109°28′
- (b) 120°
- (c) 60°
- (d) 180°

71. What type of hybridization occurs in ethylene?						
	(a)	sp (b) sp^3	(c)	sp^2 (d) sp^3d		
72.		ompound having one C—C σ -bonds is	5-bor	nd, one C—C π -bond and four		
	(a)	CH_3 — CH = CH_2	(b)	CH ₂ =CH ₂		
	(c)	CH ₂ =CH—CH=CH ₂	(d)	CH ₂ =CH—CH ₃		
73.	The co	ompound having a C—C bond	dist	ance of 1.54 Å can be		
	(a)	CH ₃ —CH ₃	(b)	CH ₃ —CH ₂ —CH ₃		
	(c)	CH ₂ =CH ₂	(d)	CH≡CH		
74.	The v	inyl group is represented by				
	(a)	CH ₂ =CH—	(b)	CH ₂ =CH—CH ₂ —		
	(c)	CH_3 — CH = CH — CH_2 —	(d)	CH ₃ —CH=CH—		
75.	Amor	ng the following,				
	(a)	CH ₃ CH=CH ₂	(b)	$CH_2 = C = CH_2$		
	(c)	CH ₃ C≡CH	(d)	CH ₂ =CH—CH=CH ₂		
	has s	p- as well as sp ² -hybridized car	rbon	atoms.		
76.	The co	ompound having a conjugated	dou	ble bond is		
	(a)	butylene	(b)	propylene		
	(c)	isobutylene	(d)	butadiene		
77.	The co	ompound having only sp ² -hyb	ridiz	ed carbon atoms is		
	(a)	1-butene	(b)	2-butene		
	(c)	propene	(d)	butadiene		
78.	Which	n of the following has the lowe	st he	at of hydrogenation per mole?		
	(a)	cis-2-Butene	(b)	trans-2-Butene		
	(c)	1-Butene	(d)	1,3-Butadiene		
79.	On re	acting with alcoholic potash, 1-	-chlo	robutane gives		
	(a)	1-butene	(b)	1-butanol		
	(c)	2-butanol	(d)	2-butene		
80.		ing heated with alcoholic potas mainly	ssiun	n hydroxide, neopentyl bromide		
	(a)	2-methyl-2-butene	(b)	2-methyl-1-butene		
	(c)	2-butene	(d)	2,2-dimethyl-1-butene		
81.	-	drohalogenation can be brough				
	(a)	an aqueous solution of KOH	(b)	an alcoholic solution of KOH		
	(c)	concentrated H ₂ SO ₄	(d)	zinc dust		

82.	2. Sodium ethoxide is a specific reagent for					
	(a)	dehydrogenation	(b)	dehydration		
	(c)	dehydrohalogenation	(d)	dehalogenation		
83.		reatment of a mixture of 1-chlorolic KOH gives	ropro	opane and 2-chloropropane with		
	(a)	1-propene	(b)	2-propene		
	(c)	isopropylene	(d)	none of these		
84.	Alkar	nes and alkenes can both be ob	taine	d by		
	(a)	Wurtz reaction	(b)	Frackland reaction		
	(c)	Kolbe's electrolytic method	(d)	Williamson synthesis		
85.	at 475	reaction of 2-halogen–substitu K, the major product obtained 1-butene		butane with concentrated H ₂ SO ₄		
	(b)	2-butene				
	(c)	both 1- and 2-butene				
	(d)	the product ratio depends on the	ne ha	logen		
86.	1-But	ene exhibits				
	(a)	geometrical isomerism	(b)	optical isomerism		
	(c)	position isomerism	(d)	none of these		
87.	The n	nost common reactions of alker	nes a	re those of		
	(a)	nucleophilic substitution	(b)	electrophilic substitution		
	(c)	electrophilic addition	(d)	nucleophilic addition		
88.	The c	onversion of ethylene into etha	ne ca	an be effected by		
	(a)	nascent hydrogen	(b)	Zn + HCl		
	(c)	Raney nickel and hydrogen	(d)	Clemmensen reduction		
89.	The n	nain product of the reaction of	1-bu	tene with excess bromine is		
		1,1-dibromobutane		1,2-dibromobutane		
	(c)	2,2-dibromobutane	(d)	perbromobutane		
90.	The co	ourse of the reaction	CH	CUD CU		
	is nro	$CH_3CH=CH_2 + HBr \longrightarrow$ dicted by	СП	3СПВГСП3		
		•	(b)	the Markovnikov rule		
		the peroxide effect		Thiel's theory		
01		eaction of 1-butene with HBr in	. ,	, and the second		
71.		2-bromobutane		1-bromobutane		
	` '	1,1-dibromobutane		1,2-dibromobutane		

92.	KOH	najor produc is	et of	the reaction	on of C.	H₃CH₂C	HBrC	LH_3	with a	alcor	10l1C
	(a)	1-butene	(b)	2-butene	(c)	butane		(d)	butyr	ne-1	
93.		number of ochlorination				hat can	be	proc	duced	by	the
	(a)	one	(b)	two	(c)	three		(d)	four		
94.		ntermediate : nce of peroxi		ed during	the add	dition of	HCl	to p	oroper	ne in	the
		CH ₃ CHCH ₂				CH ₃ CH					
	(c)	CH ₃ CHCH ₃	3		(d)	CH ₃ CH	I ₂ ĊH ₂	2			
95.	anti-N	addition of l Markovnikov	beha	viour beca	ause	_					
	(a)	the iodine figure an I_2 m			med rea	idily con	nbine	wit	h each	othe	er to
	(b)	the HI bond	l is to	o strong to	o be bro	ken hon	nolyti	cally	7		
	(c)	HI is a redu	cing	agent							
	(d)	I combines	with	H to give	back HI						
96.	The p	eroxide effec	t in a	nti-Marko	vnikov	additior	invo	lves	;		
	-	the heteroly									
	(b)	the homoly	tic fis	sion of the	e double	e bond					
	(c)	a free-radic	al me	echanism							
	(d)	an ionic me	chan	ism							
97.	Orlon	is obtained f	from								
	(a)	vinyl chlori	de		(b)	vinyl cy	anid	e			
	(c)	1,3-butadie	ne		(d)	tetraflu	oroet	hyle	ne		
98.	The o	rder of stabil	ity of	the alken	es						
		$R_2C=CR_2$,	R ₂ C=	=CHR, R II	C=CH III	[₂ ,					
		RCH=CHI IV	₹ and	d RCH=C V	CH ₂						
	is										
		I > II > III >				I = II > 1					
	(c)	II > I > IV >	III >	V	(d)	V > IV	> III >	· I >	II		

99. In dehydrohalogenation, the order of reactivity of RX is

(b) $\hat{1} > \hat{2} > \hat{3}$

(d) $\mathring{3} = \mathring{2} > \mathring{1}$

(a) 3 > 2 > 1

(c) $\mathring{2} > \mathring{3} > \mathring{1}$

100.	The re	* *	in th	e presence of benzoyl peroxide
	(a)	2-chloropropane	(b)	<i>n</i> -propyl chloride
	(c)	allyl chloride	(d)	There is no reaction.
101.	R C	$C = C \xrightarrow{R} \xrightarrow{X} \xrightarrow{KOH, \text{ heat}} \xrightarrow{R}$	C=0	
	In the	above reaction, X is		
	(a)	O_3	(b)	$KMnO_4$
	(c)	HNO ₃	(d)	O_2
102.	Ethyle	ene reacts with a dilute alkalin	e KM	InO ₄ solution to give
	(a)	НСНО	(b)	oxalic acid
	(c)	glycol	(d)	ethyl alcohol
103.	Which	n of the following is obtained b	y ad	dition polymerization?
		PVC		Terylene
	(c)	Nylon	(d)	Polyamide
104.	Polyth	nene is obtained by the polyme	erizat	tion of
	` ′	butadiene	(b)	isoprene
	(c)	ethylene	(d)	styrene
105.		n is obtained from		
	(a)	difluoroethene	(b)	monofluoroethene
	(c)	tetrafluoroethene	(d)	tetrafluoroethane
106.	In the	sequence of reactions,		
		$CH_3CH_2CH_2OH \xrightarrow{PCl_5} A$	Α	$\xrightarrow{\text{alc.}} B$
	the pr	oduct B is		KOII
	(a)	propyne	(b)	propylene
	(c)	propane	(d)	propanol
107.	In the	sequence of reactions,		
		$CH_3CH_2CH_2OH \xrightarrow{PCl_5} A$	Α —	$aq. KOH \rightarrow B$
	the pr	roduct B is		
	-	propylene	(b)	propane
	(c)	propyne	(d)	propanol
108.	Ethyle	ene chlorohydrin is obtained fr	om e	ethylene by the action of
		dry chlorine gas		dry HCl gas
		dilute HCl		a solution of chlorine in water

109.	The reaction of RCH=CH $_2$ with B_2H_6 followed by oxidation with alkaline H_2O_2 gives					
	(a)	RCH(OH)CH ₂ OH	(b)	R—COCH ₃		
	(c)	RCH ₂ CH ₂ OH	(d)	RCH ₂ CHO		
110.		najor product obtained by the 4 in the presence of HgSO ₄ is	trea	tment of propyne with aqueous		
	(a)	propanal	(b)	acetone		
	(c)	propanol	(d)	propyl hydrogen sulphate		
111.	Butad	iene has				
	(a)	only sp ² -hybridized carbon at	toms			
	(b)	only sp-hybridized carbon at	oms			
	(c)	sp- as well as sp ² -hybridized	carb	on atoms		
	(d)	sp-, sp ² - and sp ³ -hybridized of	arbo	on atoms		
112.	Which	n of the following reactions wil	l yie	ld 2,2-dibromopropane?		
	(a)	$HC=CH + 2HBr \longrightarrow$	(b)	$CH_3CH = CHBr + HBr \longrightarrow$		
	(c)	$CH_3C \equiv CH + 2HBr \longrightarrow$	(d)	$CH_3CH=CH_2 + HBr \longrightarrow$		
113.	The re	eaction of propene with chlorir	ne ga	s at about 500°C gives		
	(a)	CH ₂ ClCH=CH ₂	(b)	CH ₃ CHClCH ₂ Cl		
	(c)	CH ₂ ClCHCl—CH ₂ Cl	(d)	a mixture of (a) and (b)		
114.	The m	najor product of the debromina	tion	of meso-dibromobutane is		
	(a)	<i>n</i> -butane	(b)	1-butene		
	(c)	trans-2-butene	(d)	cis-2-butene		
115.	Chlor	oprene is				
	(a)	3-chloro-2,3-butadiene	(b)	2-chloro-1,3-butadiene		
	(c)	2,3-dichlorobutadiene	(d)	none of these		
116.	A con	npound having a bond angle o	f 180	° is a/an		
	(a)	alkane	(b)	alkene		
	(c)	cycloalkane	(d)	alkyne		
117.	1-Bute	en-3-yne contains				
	(a)	six σ - and four π -bonds	(b)	five σ - and three π -bonds		
	(c)	seven σ - and three π -bonds	(d)	eight σ - and two π -bonds		
118.	How	many structures may C ₅ H ₈ rep	resei	nt?		
	(a)	Two	(b)	Four		
	(c)	Five	(d)	Six		

- 119. The conversion of 3-hexyne into trans-3-hexene can be effected by
 - (a) Na/liquid NH₃
- (b) H₂/Lindlar's catalyst
- (c) Clemmensen reduction
- (d) LiNH₂
- 120. The relative acidities of H_2O , ROH, $HC \equiv CH$ and NH_3 are of the order
 - (a) $H_2O > ROH > HC \equiv CH > NH_3$
 - (b) $HC \equiv CH > H_2O > ROH > NH_3$
 - (c) $ROH > H_2O > HC \equiv CH > NH_3$
 - (d) $ROH=H_2O > HC=CH > NH_3$
- 121. In the reaction $CH_3CH_2CCl_2CH_3 \xrightarrow{X} CH_3C = CCH_3$, the reagent X is
 - (a) KOH/C₂H₅OH
- (b) Zn

(c) HCl/H₂O

- (d) Na
- 122. The reduction of 3-hexyne with $H_2/Lindlar's$ catalyst gives predominantly
 - (a) *n*-hexane
 - (b) trans-3-hexene
 - (c) cis-3-hexene
 - (d) a mixture of cis- and trans-3-hexene
- 123. The catalyst required for the reaction

$$HC \equiv CH + dil. H_2SO_4 \xrightarrow{catalyst} CH_3CHO$$

is

- (a) HgSO₄
- (b) Pd
- (c) Pt
- (d) AlCl₃

124. In the reaction

$$CH_3C = CCH_3 \xrightarrow{(i) X} H_3 - C - C - CH_3$$

$$0 \quad O$$

Xis

(a) O_2

(b) O_3

(c) HNO₃

- (d) KMnO₄
- **125.** The decreasing order of the strength of the bases OH^- , NH_2^- , H—C \equiv C^- and CH_3 — CH_2 is
 - (a) $H C = C^- > CH_3 CH_2 > NH_2^- > OH^-$
 - (b) $CH_3 CH_2 > NH_2 > H C = C^- > OH^-$
 - (c) $NH_2^- > H C = C^- > OH^- > CH_3 CH_2$
 - (d) $OH^- > NH_2^- > H C = C^- > CH_3 CH_2$

126. In which of the following conditions does the reaction

take place?

- (a) $HC \equiv CH \xrightarrow{HOCl}$ (b) $HC \equiv CH \xrightarrow{HOCl}$ $\xrightarrow{peroxide}$ (c) $HC \equiv CH \xrightarrow{Cl_2}$ (d) $HC \equiv CH \xrightarrow{Cl_2}$ oxidizing agent

127. In which of the following conditions does the reaction

$$HC \equiv CH + CH_3OH \longrightarrow CH_3O - C \equiv CH$$

take place?

- (a) $HC \equiv CH + CH_3OH \xrightarrow{KOMe} 160-200^{\circ}C$
- (b) $HC \equiv CH + CH_3OH \xrightarrow{conc. H_2SO_4}$
- (c) $HC \equiv CH + CH_3OH \xrightarrow{anhyd. ZnCl_2}$
- (d) $HC \equiv CH + CH_3OH \xrightarrow{C!HC = CHC^1}$
- **128.** In the reaction CH \equiv CH \xrightarrow{X} Cl₂CH \longrightarrow CHCl₂, X is
 - (a) NaOCl

(b) Cl₂/CCl₄

(c) chlorine water

- (d) NaCl
- **129.** Which of the following will not react with an ammoniacal silver nitrate solution?
 - (a) CH₃C≡CH

(b) (CH₃)₂CH—C≡CH

(c) CH₃C≡CCH₃

- (d) HC≡CH
- 130. The addition of HCN to acetylene in the presence of Ba(CN)₂ as catalyst gives
 - (a) ethyl cyanide

- (b) 1,1-dicyanoethane
- (c) divinyl cyanide
- (d) vinyl cyanide
- 131. Which reagent is used to distinguish 1-butyne from 2-butyne?
 - (a) HCl

- (b) Br₂/CCl₄
- (c) Baeyer's reagent
- (d) Ammoniacal cuprous chloride
- 132. Which reagent is the most useful for distinguishing compound I from the rest of the compounds?

CH₂CH₂C≡CH CH₃C≡CCH₃ CH₃CH₂CH₂CH₃ CH₃CH=CH₂ IV

- (a) Alkaline KMnO₄
- (b) Br_2/CCl_4
- (c) Br₂/CH₃COOH
- (d) Ammoniacal AgNO₃

133.	 A compound decolourizes an alkaline KMnO₄ solution but does not react with an ammoniacal AgNO₃ solution. It is 							
	(a)	benzene	(b) butyne-	1 (c)	butyne-2	(d)	acetylene	•
134.	precij	pitate and 1	H ₈) reacts with a H. The compo	n excess				
	(a)	CH ₂ =CH-	-CH=CH—CH	H_3 (b)	$(CH_3)_2CH_3$	—C≡(CH	
	(c)	$CH_3(CH_2)_2$	C≡CH	(d)	$(CH_3)_2C=$	C=CH	2	
135.	Whic	h among the	following is th	ne most s	trained cyc	loalkar	ne?	
	(a)	Cyclopropa	ane	(b)	Cyclobuta	ne		
	(c)	Cyclopenta	ne	(d)	Cyclohexa	ine		
136.			vclopropane (I) npounds is in t			ł cyclop	oentane (II	II) to
	(a)	I > II > III		(b)	$\mathrm{I} = \mathrm{II} > \mathrm{III}$			
	(c)	I > II = III		(d)	I=III>II			
137.			the cycloalka and cyclohexa				clobutane	(II),
	(a)	II > I > III >	IV	(b)	VI > III > I	I > I		
	(c)	II = I > III >	IV	(d)	II > I = III	> IV		
138.	Whic	h among the	following has	the great	test bond a	ngle?		
	(a)	Cyclopropa	ane	(b)	Cyclobuta	ne		
	(c)	Cyclopenta	ne	(d)	Cyclohexa	ine		
139.	Whic	h of the follo	wing is the mo	st reactiv	ve cycloalka	ane?		
	(a)	Cyclopropa	ane		Cyclobuta			
	(c)	Cyclopenta	ne	(d)	Cyclohexa	ine		
140.	BrCH	I_2 — CH_2 — C	CH ₂ Br reacts v	with Na	in the pre	esence	of ethano	ol at
	100°C	to produce						
	(a)	BrCH ₂ —CI	$H=CH_2$	(b)	$CH_2 = C = C$	CH_2		
	(c)	H ₂ C	CH,	(d)	all of these	9		
		H_2C	_					
		H_2						
1.11	The		a 6 al ala access	-:6				
141.		onformation the chair fo	of cyclohexan		the boat fo	rm		
	` ,	the half-cha		` '	the twist-b		m	
	()			\ '				

(e) an equilibrium of all the forms

- 142. 3,5-Dimethylcyclopentene, on ozonolysis, yields
 - (a) only an aldehyde
- (b) only a ketone
- (c) an aldehyde and a ketone
- (d) a dialdehyde
- 143. Which of the following represents decalin?



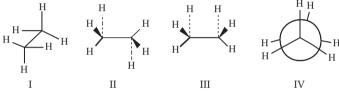


(c) H

Н



144. Which of the following represents the staggered conformation of ethane?



(a) I and II

(b) I, II and III

(c) I and III

- (d) II, III and IV
- 145. The order of reactivity of the halogens (Cl_2 , Br_2 , I_2 , F_2) towards methane is
 - (a) $F_2 > Cl_2 > Br_2 > I_2$
- (b) $F_2 > Br_2 > Cl_2 > I_2$
- (c) $F_2 > I_2 > Br_2 > Cl_2$
- (d) $I_2 > F_2 > Cl_2 > Br_2$
- **146.** The reaction of propene with chlorine at 600°C gives
 - (a) CICH₂CH=CH₂
- (b) CH₃CHCl—CH₂Cl
- (c) CH₃CHCl—CH₃
- (d) ClCH₂CH₂CH₂Cl
- 147. The product obtained in the reaction

$$CH_3$$
 $C=C$ CH_3 Br_2

is

(a) CH_3 $H \longrightarrow Br$ CH_3





148. Arrange the following in order of increasing acidity.

- (a) $HF < HCl < H_2S < H_2O < NH_3 < CH_4$
- (b) $H_2O < NH_3 < CH_4 < H_2S < HF < HCl$
- (c) $HCl < H_2S < HF < H_2O < NH_3 < CH_4$
- (d) $CH_4 < NH_3 < H_2O < HF < H_2S < HCl$
- 149. The hyperconjugative effect in the compounds

$$CH_3CH=CH_2$$
 $(CH_3)_2CH=CH_2$ $(CH_3)_3C-CH=CH_2$ III III

is of the order

(a) I > II > III

(b) III > II > I

(c) I > III > II

- (d) II > III > I
- 150. In the reaction of ethyl alcohol with an excess of concentrated H₂SO₄ at 440 K to give ethylene, the following steps are involved.
 - (a) $CH_3CH_2OH + H^+ \Longrightarrow CH_3CH_2 \stackrel{+}{\longrightarrow} CH_3$ (b) $CH_3CH_2 \stackrel{+}{O}H_2 \longrightarrow CH_3 \stackrel{+}{\longrightarrow} CH_2 + H_2O$

 - (c) $CH_3 \stackrel{+}{\longrightarrow} CH_2 \longrightarrow CH_2 = CH_2$
 - (d) $CH_3CH_2OH \xrightarrow{-H_2O} CH_2=CH_2$

Of the above, which is the fast step?

151. The stability of the alkenes

$$\begin{array}{cccc} CH_2 \!\!=\!\! CH_2 & CH_3CH \!\!=\!\! CH_2 & (CH_3)_2C \!\!=\!\! CH_2 & (CH_3)_2 \!\!=\!\! CHCH_3 \\ I & II & III & IV \end{array}$$

decreases in the order

(a) IV > III > II > I

(b) I > II > III > IV

(c) I > III > II > IV

- (d) II > III > I > IV
- 152. The carbon-carbon bond length in the compounds

$$CH_2$$
= CH_2 CH_3 — CH_3 CH = CH

follows the order

(a) III < II < IV

(b) IV < I < II < III

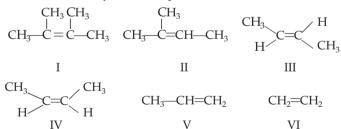
- (c) I < II < III < IV
- (d) I < IV < III < II
- **153.** What is the order of the ease of formation of the following free radicals?
 - (a) Allyl > $3 > 2 > 1 > CH_3 > vinyl$
 - (b) Vinyl > $\dot{C}H_3 > \dot{1} > \dot{2} > \dot{3} > \text{allyl}$
 - (c) $\mathring{1} > \mathring{2} > \mathring{3} > \text{allyl} > \mathring{C}H_3 > \text{vinyl}$
 - (d) $\dot{C}H_3 > \dot{1} > \dot{2} > \dot{3} > \text{allyl} > \text{vinyl}$
- **154.** A compound C₄H₈ decolourizes a KMnO₄ solution. How many structures are possible for this compound?
 - (a) 3
- (b) 4
- (c) 2
- (d) 5
- 155. Which of the following compounds will exhibit geometrical isomerism?
 - (a) Propene

- (b) Isobutylene
- (c) 1,2-Dibromoethene
- (d) 1-Butene
- **156.** The normal C=C bond length in ethylene is 1.49 Å and the normal C—C bond length in propane is 1.54 Å. What will be the bond length of the C_2 — C_3 single bond in propene (CH₃—CH₂=CH₂)?
 - (a) 1.49 Å

(b) 1.54 Å

(c) < 1.49 Å

- (d) Between 1.49 \mathring{A} and 1.54 \mathring{A}
- **157.** The relative stability of the compounds



is of the order

- (a) I > II > III > IV > V > VI
- (b) VI > V > IV > III > II > I
- (c) I > III > V > II > IV > VI
- (d) II > I > IV > III > V > VI
- **158.** An organic compound decolourizes bromine in CCl₄ and can be reduced catalytically. Also, it gives a precipitate with ammoniacal cuprous chloride. The compound can be
 - (a) CH_3 —CH=CH— CH_3
- (b) CH₃CH₂C≡CH

(c) CH₃CH=CH₂

(d) CH₃C=CH₂ | | CH₃

159.
$$CH_3$$
— CH_2 — CH — CH_3 $\xrightarrow{\overline{O}H}$ alcoholic

The product obtained in the above reaction is

- (a) CH₃CH=CHCH₃
- (b) CH₃CH₂CH=CH₂
- (c) a mixture of CH₃CH=CHCH₃ and CH₃CH₂CH=CH₂
- (d) there is no reaction
- **160.** In the formation of cyclohexane from 1-hexene, the entropy of cyclohexane is
 - (a) more than that of 1-hexene
 - (b) less than that of 1-hexene
 - (c) equal to that of 1-hexene
 - (d) more or less than that of 1-hexene depending on the conditions of the reaction

161. (A)
$$CH_3 - CH_2 - CH_2$$

Consider the reactions (A) and (B). The entropy change will be

- (a) more in reaction A than in reaction B
- (b) more in reaction B than in reaction A
- (c) equal in both reactions
- (d) zero in both reactions
- **162.** The bond dissociation energies of the following

- (c) IV > I > II > III (d) II > I > IV > III
- 163. The reaction of isobutylene with water in the presence of 10% H₂SO₄ at

(a)
$$CH_3$$
 CH_3 (b) CH_3 CH_2OH CH_3 CH_2OH

(d) none of these

- **164.** Isobutylene reacts with HBr to give *t*-butyl bromide. The intermediate formed in this reaction is
 - (a) a t-butyl cation
 - (b) an isobutyl cation
 - (c) a mixture of t- and isobutyl cations
 - (d) The addition does not take place through a cation intermediate.
- **165.** Tri-*n*-propylborane can be converted into propyl alcohol by reaction with
 - (a) hydrogen peroxide at 25–30°C
 - (b) a dilute acid at reflux temperature
 - (c) a dilute alkali at reflux temperature
 - (d) none of these
- **166.** Which of the following represents the hybrid structure of acrolein?

(a)
$$CH_2 = CH - CH = \ddot{O}$$
:

(a)
$$CH_2 = CH - CH = \ddot{O}$$
: (b) $CH_2 = CH - \ddot{C}H = \ddot{O}$:

(c)
$$\overset{\oplus}{\text{CH}}_2 = \text{CH} = \text{CH} - \ddot{\text{O}}$$
:

(c)
$$\overset{\oplus}{\text{CH}}_2 = \text{CH} = \text{CH} - \overset{\bullet}{\text{CH}} \overset{(d)}{:}$$
 $\overset{\delta \oplus}{\text{CH}}_2 = \text{CH} \overset{\delta \oplus}{=} \overset{2\delta \ominus}{\text{CH}} \overset{2\delta \ominus}{=}$

167. The rate of abstraction of hydrogen from $\mathring{3}$, $\mathring{2}$ and $\mathring{1}$ carbon atom follows the order

(a)
$$3 > 2 > 1$$

(b)
$$2 > 3 > 1$$

(c)
$$1 > 2 > 3$$

(a)
$$\mathring{3} > \mathring{2} > \mathring{1}$$
 (b) $\mathring{2} > \mathring{3} > \mathring{1}$ (c) $\mathring{1} > \mathring{2} > \mathring{3}$ (d) $3^{\circ} > 1^{\circ} > 2^{\circ}$

168. The amount of energy needed to form the radicals $\dot{C}H_3$, $\mathring{1}$, $\mathring{2}$ and $\mathring{3}$ decreases in the order

(a)
$$3 > 2 > 1 > CH_2$$

(b)
$$\dot{C}H_3 > \dot{1} > \dot{2} > \dot{3}$$

(c)
$$\mathring{1} > \mathring{2} > \mathring{3} > \mathring{C}H_3$$

(d)
$$\dot{C}H_3 > 3 > 2 > 1$$

- 169. Arrange the free radicals vinyl (H₂C=CH), allyl H₂C=CH—CH₂ and benzyl (C₆H₅—CH₂) in order of their stability.

 - (a) Benzyl > allyl > vinyl
 (b) Allyl > vinyl > benzyl
 (c) Vinyl > allyl > benzyl
 (d) Vinyl > benzyl > allyl (b) Allyl > vinyl > benzyl
- 170. The ease of abstraction of allylic, $3, 2, 1, CH_4$ and vinylic hydrogens follows the order
 - (a) allylic $> 3 > 2 > 1 > CH_4 > vinylic$
 - (b) $\mathring{3} > \mathring{2} > \mathring{1} > CH_4 > \text{allylic} > \text{vinylic}$
 - (c) $CH_4 > \mathring{3} > \mathring{1} > \mathring{2} > \text{vinylic} > \text{allylic}$
 - (d) vinylic > allylic > $\mathring{1} > \mathring{2} > \mathring{3} > CH_4$

• Type 2 •

Choose the correct options. More than one option is correct.

- 171. Which of the following compounds exhibit geometrical isomerism?
 - (a) CH₃CH=CH—COOH
- (b) Br—CH=CH—Br
- (c) C₆H₅CH=NOH
- $^{(d)}$ \bigcirc O
- 172. Which of the following statements are correct?
 - (a) 1,1-Dichloroethane exhibits geometrical isomerism.
 - (b) 1,2-Dichloroethane shows geometrical isomerism.
 - (c) 1-Butene exhibits geometrical isomerism.
 - (d) Isobutylene does not show geometrical isomerism.
- 173. Which of the following notations are correct?
 - (a) Br C=C F C C C C
- (b) Br C=C I I
- (c) H_3C C=C H (Z)
- (d) H_3C C=C $CH_2CH_2CH_3$ CH_3 CH_3
- 174. Which of the following has dipole moment?
 - (a) cis-2-butene
 - (b) trans-2-butene
 - (c) cis-1,2-dibromoethene
 - (d) trans-1,2-dibromoethene
- 175. Which of the following statements are correct about cis and trans isomers?
 - (a) *cis*-2-butene can be converted into *trans*-2-butene by irradiation.
 - (b) In general, trans isomers have zero dipole moment.

- (c) On heating, fumaric acid (a trans acid) gives an anhydride.
- (d) On heating, maleic acid (a cis acid) gives an anhydride.
- **176.** For which of the following pairs of compounds are the correct notations given?

(b)
$$H_3C$$
. $C=N$ and H_3C . $C=N$ $C=N$

(c)
$$H \subset CO_2H$$
 $C=C \subset H \subset CO_2H$ NH_2

Cis -o-aminocinnamic acid

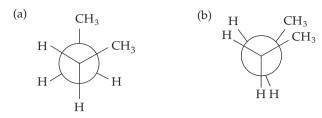
(d)
$$H_3C$$
 $CH_2CH_2CH_3$ $CICH_2$ $C=C$ CH_3 CH_3 CH_2 $C=C$ CH_3 CH

177. Which of the following statements are correct?

Trans -o-aminocinnamic acid

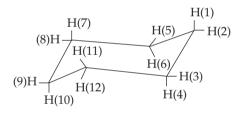
- (b) CH₃CH₂CH=CH₂ and CH₃CH₂CH=CHCH₃ are examples of position isomerism.
- (c) C₂H₅OCH₃ and CH₃CH₂CH₂OH represent functional-group isomerism.
- (d) CH₃CH₂NH₂ and CH₃NHCH₃ are examples of metamerism.

178. Which of the following represent staggered conformations of *n*-butane?



(c)
$$CH_3$$
 H H

179. In the chair conformation of cyclohexane (given below)



which of the following hydrogens (marked 1–12 in the representation above) are axial?

(a) 1

(b) 2

(c) 9

(d) 10

180. Which of the following have chiral carbon?

(a)
$$H$$
 (b) H $C_2H_5-C-CH_3$ (c) H (d) H $C_6H_5-C-CH_3$ $C_6H_5-C-CH_3$

- **181.** Which of the following representations have an R configuration?
 - (a) Br CC CI H

- (b) Br
- (c) COOH
- **182.** Which of the following compounds exhibit optical isomerism?
 - (a) CO_2H NO_2 NO_2 CO_2H
- (b) $CH_2 = C = CH_2$
- C=C=C
 β-Naphthyl β-Naphthyl
- $\begin{array}{ccc} \text{(d)} & \text{H}_2\text{C} & \text{CH}_2 \\ & & \text{H}_2\text{C} & \text{CH}_2 \end{array}$

(b)

(d)

- 183. Which of the following represent a pair of enantiomers?
 - (a) HO_2C H CO_2H
- H CO_2H HO_2C H
- (c) $HO_2C \longrightarrow CO_2H$ $H \longrightarrow H$
- H HO_2C CO_2H
- 184. 2-Bromo-3-phenylpropane can be synthesised by
 - (a) $C_6H_5CH_2CH(OH)CH_3 + PBr_3 \longrightarrow$
 - (b) $C_6H_5CH=CHCH_3 + HBr + benzoyl peroxide \longrightarrow$

(c)
$$C_6H_5CH_2CH_2CH_3 + Br_2 + light \longrightarrow$$

(d) none of these

185. Which of the following structures can exist as cis-trans isomers?

186. Which of the following reactions are expected to give

$$\begin{array}{c} CH_3 \\ | \\ CH_3 - C - CH = CH_2 \\ | \\ CH_2 \end{array}$$

in yields of more than 50%?

(a)
$$CH_3$$
 CH_3 $CH_$

(b)
$$CH_3$$
 CH_3 $CH_$

(c)
$$CH_3$$
 CH_3 $CH_$

(d) None of these

Η

187. Which of the following are correct?

(a)
$$\frac{1. O_3}{2. Zn, H_2O} OHC - CH_2CH_2CH_2 - CHO$$
(b)
$$\frac{C_6H_5C - OOH}{CH_2Cl_2} H$$

- $\begin{array}{c|c}
 & & & \\
 \hline
 \text{KMnO}_4 & & \\
 \hline
 \text{OH, heat} & & \\
 \hline
 \end{array} \quad \begin{array}{c}
 & & \\
 \hline
 \text{O-CCH}_2\text{CH}_2\text{CH}_2\text{C} \text{O}
 \end{array}$
- **188.** Which of the following reagents can be used as the basis for a simple chemical test for distinguishing 1-butene from CH₃CH₂CH₂CH₂OH?
 - (a) Br_2/CCl_4

- (b) Dilute aqueous KMnO₄
- (c) CrO₃/aqueous H₂SO₄
- (d) None of these
- 189. Addition polymerization can be brought about by
 - (a) free radicals

(b) anions

(c) cations

- (d) none of these
- 190. Which of the following can be used for the preparation of propane?

(a)
$$CH_3CH=CH_2 \xrightarrow{1. B_2H_6} 2. CH_3COOH \rightarrow$$

(b)
$$CH_3CH_2CH_2Cl \xrightarrow{1. Mg/ether}$$

(d)
$$CH_3CH_2CH_2COONa \xrightarrow{NaOH(CuO)} \Delta$$

- 191. The cis hydroxylation of alkenes can be effected by
 - (a) the addition of a 1% KMnO₄ solution
 - (b) reaction with osmium tetraoxide followed by treatment with water
 - (c) the addition of ozone to the alkene, followed by treatment of the ozonide with $\mbox{\rm H}_2\mbox{\rm O}_2$
 - (d) none of these
- **192.** Which of the following transformations are feasible?

(a)
$$CH_3CH_2CH_2CH_3 \xrightarrow{AlCl_3, \Delta} CH_3 \xrightarrow{CH_3} CH_3$$

(b)
$$CH_3$$
— CH_2 — CH_2 — $CH=CH_2$ — $AlCl_3$ — $200-300^{\circ}C$ — CH_3 — CH_2 — $CH=CH$ — CH_3

(c)
$$CH_3CH_2CH=CH_2 \xrightarrow{AICl_3} CH_2=C-CH_3$$

 CH_3

(d)
$$CH_3$$
 CH_3 $AICl_3/\triangle$

193. Which of the following olefines have Z-configurations [in the (E)–(Z) system]?

$$C = C$$
 CH_3
 $C = C$
 CH_4

$$C = C$$
 H
 $C = C$
 H

194. What are the possible intermediates in the following reaction?

$$CH_{3}CH_{2}CH = CH_{2} \xrightarrow{1. Br_{2}/CCl_{4}} CH_{3}CH_{2}C = CH_{3}CH_{2}C =$$

- (a) CH₃CH₂CH—CH₂Br (b) CH₃CH₂CH=CHBr
 Br
 (c) CH₃CH₂C=CH₂ (d) None of these

- 195. On oxidation, an unknown alkene C₈H₁₆ gives a mixture of propanoic acid (CH₃CH₂COOH) and pentanoic acid (CH₃CH₂CH₂CH₂COOH). The unknown alkene can be
 - (a) cis-3-octene

(b) trans-3-octene

(c) cis-4-octene

- (d) trans-4-octene
- **196.** The nitration of propane with concentrated HNO₃ gives
 - (a) CH₃CH₂CH₂NO₂
- (b) CH₃CHCH₃ NO₂

(d) CH₃CH₂NO₂

(d) CH₃NO₂

197. The reaction of isobutylene

with 60% H₂SO₄ at 70°C gives

(a)
$$CH_3$$
 (b) CH_3 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3

(c)
$$CH_3$$
 CH_2 (d) CH_3 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3

- 198. Which of the following are optically active?
 - (a) $H_2C=C=CH_2$

$$(d) \qquad F \qquad F \qquad Cl \qquad CO_2H$$

- 199. In which of the following compounds are all the carbon atoms in the sp³ state of hybridization?
 - (a) CH₄
- (b) C₂H₄
- (c) C_3H_8 (d) C_4H_{10}
- **200.** Which of the following statements are not correct for alkanes?
 - (a) All C—H and C—C bonds have a length of 1.112 Å and 1.54 Å respectively.

(b) All bond angles are tetrahedral, having a value of 109.5°.

201. Arrange the following in order of increase/decrease in boiling point.

202. Which of the following reactions can be used to prepare methane?

(CH₃)₂CHCH₂CH₃

(b) II > I > III

(d) III < II < I

(CH₃)₄C

(c) The C—C chain is linear and not zigzag.

(d) All alkanes exhibit isomerism.

CH₃CH₂CH₂CH₂CH₃

(a) Clemmensen reduction

(b) Wurtz reaction

(a) I > II > III

(c) III > I > II

203.

204.

205.

206.

207.

\ /					
(c)	Catalytic hydrogenation of methyl iodide				
(d)	Reduction of methyl iodide by using a zinc-copper couple				
A mixture of ethyl iodide and methyl iodide is subjected to the Wurtz reaction. The products formed are					
(a)	ethane	(b)	butane		
(c)	propane	(d)	2-methylpropane		
Metha	nne is obtained when				
(a)	sodium acetate is heated with	sod	a lime		
(b)	iodomethane is reduced				
(c)	aluminium carbide reacts wit	h wa	ter		
(d)	potassium acetate is electroly	sed			
Which	n of the following will give thre	ee m	onobromo derivatives?		
(a)	CH ₃ CH ₂ CH ₂ CH ₂ CH ₂ CH ₃	(b)	CH ₃ CH ₂ CH ₂ —CH(CH ₃)CH ₃		
(c)	CH ₃ —CH ₂ —C(CH ₃) ₂ CH ₃	(d)	CH ₃ CH(CH ₃)CH(CH ₃)CH ₃		
Which of the following compounds cannot be prepared by the Wurtz reaction?					
(a)	CH ₃ CH ₃	(b)	CH ₃ CH—CH ₃		
			CH ₃ CH—CH ₃ CH ₃		
(c)	$(CH_3)_2CHCH_3$	(d)	CH ₃ CH ₂ CH ₂ CH ₃		
Which	n of the following are cumulati	ve di	ienes?		
(a)	CH ₃ —CH=C=CH—CH ₃	(b)	CH ₂ =CH—CH=CH ₂		
(c)	CH ₂ =CH—CH ₂ CH=CH ₂	(d)	CH ₃ CH ₂ CH=C=CH—CH ₃		

208.	Alken	es undergo		
	(a)	substitution reactions	(b)	addition reactions
	(c)	ozonolysis	(d)	none of these
209.	What are the products obtained by the ozonolysis of RCH= CR_1R_2 ?			zonolysis of RCH=CR ₁ R ₂ ?
	(a)	RCHO	(b)	R_1R_2CO
	(c)	R ₂ CO	(d)	$RCH_2CH_2R_1$
210.	What	are the products obtained upo	n the	e ozonolysis of 2-pentene?
	(a)	CH ₃ CH ₂ CHO	(b)	CH ₃ CHO
	(c)	CH ₃ COCH ₃	(d)	CH ₃ COCH ₂ CH ₃
211.	Which	n of the following statements as	re co	rrect for geometrical isomers?
	(a)	The cis-isomer is more polar t	han	the trans-isomer.
	(b)	The boiling point of the cistrans-isomer.	-isoı	mer is higher than that of the
	(c)	Geometrical isomers have di chemical properties, though s		nt physical properties but their ar, are not identical.
	(d)	The stability of trans-isomers	is gr	reater than that of cis-isomers.
212.	C_4H_6	may contain		
	(a)	only single bonds	(b)	a double bond
	(c)	a triple bond	(d)	two double bonds
213.	The re	eactivities of ethane, ethylene a	nd a	cetylene are of the order
		ethane < ethylene < acetylene		
		ethane < acetylene < ethylene		
	(c)	,		
		acetylene = ethylene > ethane		
214.		n of the following contains acid		
		Ethene		Ethane Butune 1
•	(c)	Ethyne		Butyne-1
215.		n of the following will react wi		
	(a) (c)	Ethyne Butyne-2		Butyne-1 Ethane
216	. ,	•	` '	
216.		n of the following statements as		rrect: hylene to an electrophilic attack.
		•		similar reactivities towards an
	(0)	electrophilic attack.	, v v . c	minut reactivities towards are
	(c)	The reactivities of acetylene a	nd e	thylene towards an electrophilic

attack depend on the electrophilic reagent.

(d) Acetylene is less reactive than ethylene to an electrophilic attack.

217.	Which	Which of the following reactions will give an alkyne?		
	(a)	Potassium fumarate — electrolysis —		
	(b)	CH ₃ CBr ₂ CHBr ₂ —Zn/alcohol	\xrightarrow{l}	
		$CH_3CH_2CHBr_2 \xrightarrow{ alc. \ KOH }$		
	(d)	$CH_{3}CHBrCH_{2}Br \xrightarrow{ NaNH_{2} }$		
218.	That a	cetylene is a linear molecule i	s sho	wn by
		its C≡C bond distance being		
	(b)	its C—H bond distance being	g 1.08	3Å
	(c)	its H—C—C bond angle bei	ing 18	80°
		X-ray diffraction	Ü	
219.		n of the following react with C sence of diffused sunlight to p		d Br ₂ at room temperature and in ace dihalogen derivatives?
		Cyclopropane		Cyclobutane
	(c)	Cyclopentane	(d)	Cyclohexane
220.	The C	=C bond distance in an organ		=
		butene-1		butene-2
	(c)	cyclohexatriene	(d)	hexatriene
221.	The p	roduct obtained in the reaction		
		$C_6H_5CH_2$ — CH_3 — C	2, hν 73 K	\rightarrow
	is			
	(a)	C ₆ H ₅ CHCl—CH ₃	(b)	C ₆ H ₅ CH ₂ CH ₂ Cl
	(c)	$C_6H_5CCl_2$ — CH_3	(d)	C ₆ H ₅ CHClCH ₂ Cl
222.	Which	n of the following compounds	will	exhibit geometrical isomerism?
	` '	Butene-1	` '	Butene-2
	(c)	Oxime of benzaldehyde	(d)	Oxime of acetone
223.		n of the following will have di		
		Propylene	` '	1-Butene
	. ,	cis-2-Butene	` '	trans-2-Butene
224.		n of the following is a nucleop:		_
		NO _		ОН
	(c)	RCOO	(d)	RNH_2

225. Which of the following will give cis-diols?

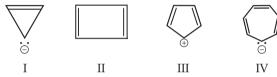
(a)
$$C = C \left(\frac{1. \text{ KMnO}_4}{2. \text{ H}_2 \text{O}} \right)$$

(b)
$$C=C \left(\frac{1.O_s O_4}{2.Na_2SO_3}\right)$$

(c)
$$\frac{1. \text{ OsO}_{4}, 25^{\circ}\text{C}}{2. \text{ Na}_{2}\text{SO}_{3}}$$

(d)
$$\frac{35\% \text{ H}_2\text{O}_2}{\text{HCO}_2\text{H},25^\circ\text{C}}$$

226. Which of the following structures



will have 4 electrons?

- (a) I
- (b) II
- (c) III
- (d) IV

227. Which of the following is a nucleophile?

- (a) $: O\overline{H}$
- (b) $: \overline{CN}$
- (c) $:\overline{C} = CR$
- (d) $: \overline{SH}$

1. c	2. a	3. b	4. c	5. c
6. d	7. a	8. a	9. d	10. b
11. b	12. d	13. d	14. c	15. a
16. b	17. c	18. d	19. b	20. d
21. c	22. c	23. a	24. d	25. c
26. c	27. c	28. b	29. c	30. b
31. a	32. d	33. c	34. d	35. a
36. c	37. a	38. a	39. b	40. c
41. b	42. c	43. d	44. a	45. a
46. b	47. a	48. a	49. a	50. a
51. b	52. b	53. d	54. a	55. c
56. c	57. b	58. c	59. b	60. c
61. d	62. c	63. d	64. c	65. c
66. d	67. a	68. a	69. d	70. b

71. c	72. b	73. c	74. a	75. b
76. d	77. d	78. d	79. a	80. a
81. b	82. c	83. a	84. c	85. b
86. c	87. c	88. c	89. b	90. b
91. b	92. b	93. b	94. c	95. a
96. c	97. b	98. a	99. a	100. a
101. b	102. c	103. a	104. c	105. c
106. b	107. d	108. d	109. c	110. b
111. a	112. c	113. a	114. c	115. b
116. d	117. c	118. d	119. a	120. a
121. a	122. c	123. a	124. b	125. b
126. a	127. a	128. b	129. c	130. d
131. d	132. d	133. c	134. b	135. a
136. a	137. b	138. d	139. a	140. c
141. e	142. d	143. c	144. a	145. a
146. a	147. a	148. d	149. a	150. c
151. a	152. b	153. a	154. b	155. c
156. d	157. a	158. b	159. a	160. b
161. b	162. c	163. a	164. a	165. a
166. d	167. a	168. b	169. a	170. a
171. a, b, c	172. b, d	173. a, c	174. a, c	175. a, b, d
176. b, c, d	177. a, b, c, d	178. a, c	179. a, d	180. a, b, d
181. a, c	182. a, c	183. a, b	184. a, b	185. a, c
186. b, c	187. a, b, c, d	188. a, b, c	189. a, b, c	190. a, b, c, d
191. a, b	192. a, b, c	193. a, c, d	194. a, b, c	195. a, b
196. a, b, c, d	197. c, d	198. b, c	199. a, b, c, d	200. c, d
201. a, d	202. c, d	203. a, b, c	204. a, b, c	205. a, c
206. b, c	207. a, d	208. a, b, c	209. a, b	210. a, b
211. a, b, c, d	212. c, d	213. a, c	214. c, d	215. a, b
216. a, b, c	217. a, b, c, d	218. c, d	219. a, b	220. a, b
221. a, c	222. b, c	223. a, b, c	224. b, c, d	225. a, b, c
226. a, b, c	227. a, b, c, d			

Hints to More Difficult Problems

6. The potential energy of the molecules is at a minimum for the staggered conformations, increases with rotation and reaches a maximum at the eclipsed conformation.

- 9. Cyclohexane is virtually strain-free. Hence the heat of combustion per CH₂ is the same as for *n*-alkanes (157.4 kcal). The increase in heat of combustion for the smaller rings is due to increase in angle strain.
- **13.** Trans elimination is easier than cis elimination. Also, the formation of a more substituted alkane is favoured.
- **15.** Enantiomers are mirror images of each other.
- **19.** Compounds in which the double bonds are in alternate positions, i.e., in case of conjugation, are stable.
- **20.** Allylic hydrogen has a low bond dissociation energy, and so NBS is used as a brominating agent in the allylic position.
- 22. The generated dichlorocarbene adds on to the double bond.
- 23. The chair form of cyclohexane is not only free of angle strain, but of torsional strain as well. It is, therefore, the most stable conformation of cyclohexane.
- **35.** The pairs which are nonsuperimposable mirror images represent enantiomers.
- **41.** Carbon radicals in suitable unsaturated systems are more stable than alkyl radicals due to the delocalization of the unpaired electron.
- **44.** An sp³-sp³ sigma bond is the longest and there is a steady decrease in the length of carbon-carbon single bonds as the hybridization state of the bonded atoms changes from sp³ to sp.
- **129.** To be able to react with ammoniacal AgNO₃, the alkyne must have an acidic (terminal) H atom.
- **171.** Alkenes with the formula baC=Cab or baC=Ced exhibit geometrical isomerism. The oximes of aldehydes also exhibit geometrical isomerism because hindered rotation is possible with a carbon-nitrogen double bond.
- 174. Trans isomers, in general, have zero dipole moment.
- **180.** A chiral carbon must have four different atoms or groups attached to it.
- **182.** The rotation of biphenyls with ortho positions substituted by a bulky group (like —COOH, —NO₂) is restricted. The molecule has no elements of symmetry, i.e., it is not superimposable on its mirror image, and therefore exhibits optical isomerism. A simple allene does not exhibit optical isomerism but allenes of the type abC=C=Cab exist in two enantiomeric forms provided a and b are bulky groups.
- **183.** The nonsuperimposable pairs are enantiomers.
- **185.** Alkenes with the formula baC=Cab or baC=Ced can exist as cis-trans isomers.

- **196.** The reaction occurs by a free-radical mechanism, and so a mixture of products is obtained.
- **197.** It is a polymerization reaction. The chain termination occurs after only one isobutylene molecule has been added. This is due to the high concentration of water; the intermediate carbonium ion loses a proton to water before it can react with another alkene molecule.
- 206. The Wurtz reaction is used to make alkanes with an even number of carbon atoms.
- **207.** In cumulative dienes, there is at least one carbon atom joined to both the neighbouring carbon atoms by double bonds, i.e., they contain the group



 \Box

Halogen Derivatives

• Type 1 •

Choose the correct option. Only one option is correct.

1. Which of the following compounds is the most likely to undergo a bimolecular nucleophilic substitution reaction with aqueous NaOH?

(c)
$$Br$$
 CH_3

- **2.** Which of the following reagents can be used to distinguish chlorobenzene from chlorocyclohexane?
 - (a) AgNO₃/C₂H₅OH
- (b) Ag(NH₃)₂OH
- (c) Na fusion; HNO₃; AgNO₃
- (d) Br₂/CCl₄
- 3. Which of the following can be used to prepare 3-bromopropene?

(a)
$$CH_3CH=CH_2 + Br_2 \xrightarrow{CCl_4} 30^{\circ}C$$

(b) $CH_3CH=CH_2 + N$ -bromosuccinimide \longrightarrow

(c)
$$CH_2 = CHCH_3 + PBr_3 \longrightarrow$$

- 4. The number of structural isomers of C₃H₅Cl₃ is
 - (a) two

(b) three

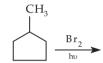
(c) four

- (d) five
- 5. Which of the following molecules has zero dipole moment?
 - (a) CCl₄

(b) CHCl₃

(c) CH₂Cl₂

- (d) CH₃Cl
- 6. The major product obtained in the reaction



is

(a) CH₂Br (b) CH_{2}

(c)

- (d) CH_{2} Br
- 7. How many monochloro derivatives does the free-radical chlorination of 2,3-dimethylbutane yield?
 - (a) One

(b) Two

(c) Three

- (d) Four
- 8. Which of the following yields one monosubstituted chloroalkane upon chlorination?
 - (a) Isobutane

(b) Cyclopentane

(c) *n*-Butane

- (d) Propane
- 9. Which of the following alkyl halides will you expect to give the highest yield of substitution products under conditions favourable to a bimolecular reaction?

 - (a) CH₃CH₂CH₂CH₂CH₂Br (b) CH₃CH₂CH₂CH—CH₃

(c)
$$CH_3CH_2$$
— CH — CH_2Br (d) CH_3CH_2 — C — CH_3 CH_3

10. What is the major product in the following reaction?

$$\begin{array}{c} CH_3 \\ CH_3 \\ CH_2 \\ CH_2 \\ CH_3 \end{array} + KOH \xrightarrow{\quad \text{ethanol} \quad \\ \text{reflux} \\ \end{array}$$

(c)
$$CH_3CH_2$$
— $C=CH_2$ (d) $CH_3CH=C$ — CH_3 CH_3

11. The major product in the reaction

$$CH_{3} \xrightarrow{CH} CH_{3} CH_{2}CH_{2}CH_{3} \xrightarrow{-} ^{+} C_{2}H_{5}OH$$

$$CH_{3} \xrightarrow{CH} CH_{2}CH_{3}$$

$$CH_{3} \xrightarrow{C} CH_{2}CH_{3}$$

is

(a)
$$(CH_3)_2C=C(CH_3)CH_2CH_3$$

(b)
$$(CH_3)_2CHC=CH_2$$

 CH_2CH_3

(d)
$$(CH_3)_2CHC(CH_3)_2$$
— C — $(CH_3)CH(CH_3)_2$
 CH_2CH_3

- 12. How many chiral stereoisomers can be drawn for CH₃CHCl—CHBrCH₃?
 - (a) Two

(b) Three

(c) Four

(d) Five

13. Which of the following represents Z-3-chloro-3-heptene?

(a)
$$CH_3$$
 $C=C$ H $CHCH_2CH_3$

(b)
$$CH_3CH_2$$
 $C=C$
 $CH_3CH_2CH_2$

(c)
$$CH_3CH_2$$
 $C=C$

$$CH_3CH_2$$

$$C=C$$

$$CH_2CH_2CH_3$$

- 14. Which of the following is true about any R-enantiomer?
 - (a) It is dextrorotatory.
 - (b) It is levorotatory.
 - (c) It is an equal mixture of + and -.
 - (d) It is the mirror image of the S-enantiomer.
- 15. Which of the following is not true about enantiomers?
 - (a) They have the same melting or boiling point.
 - (b) They have the same specific rotation.
 - (c) They have the same density.
 - (d) They have the same chemical reactivity.
- 16. 2,2-Dichloropentane can best be synthesised by

(a)
$$CH_2CH_2CH_2C \equiv CH \xrightarrow{Cl_2}$$

$$\text{(b)} \quad CH_3CH_2CH_2C \Longrightarrow CH \stackrel{H_2}{\longrightarrow} [\quad] \stackrel{Cl_2}{\longrightarrow}$$

(c)
$$CH_3CH_2CH_2C \equiv CH \xrightarrow{2HCl}$$

(d)
$$CH_3CH_2CH=CHCH_3 \xrightarrow{HCl}$$

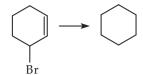
- 17. Which of the following compounds is ionic?
 - (a) BF_3

(b) CCl₄

(c) MgI₂

(d) AlCl₃

18. The transformation



can be brought about using

(a) Zn/H^+

(b) CuI

(c) Na

- (d) (CH₃)₂CuLi
- **19.** Which of the following rotamers of 1,2-dichloroethane has zero dipole moment?
 - (a) Cl H
- (b) H H CI
- (c) CIH
- (d) H H CI
- 20. In the $S_{\rm N}2$ reaction of *cis*-3-methylcyclopentyl bromide with alkali, the product formed is
 - (a) a cis alcohol
 - (b) a trans alcohol
 - (c) an equimolecular mixture of cis and trans alcohols
 - (d) there is no reaction
- 21. The reaction

$$RCH = CH - CH_2 - X \xrightarrow{y^-} R - CH - CH = CH_2$$

is

- (a) an S_N1 reaction
- (b) an $S_N 2$ reaction
- (c) an S_Ni reaction
- (d) none of these

22. The reaction

$$CH_2 = CH - CH_3 + Cl_2 \xrightarrow{400^{\circ}C} CH_2 = CH - CH_2Cl$$

proceeds through the intermediate formation of a

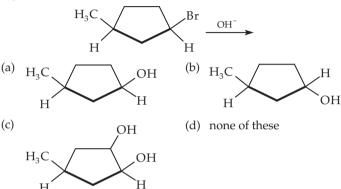
(a) free radical

(b) carbocation

(c) carbanion

(d) none of these

23. The product obtained in the reaction



- **24.** In the reaction of optically active 2-bromo-octane with sodium hydroxide to give 2-octanol, there is
 - (a) retention of configuration
 - (b) inversion of configuration
 - (c) retention and inversion of configuration in equal amounts of the product
 - (d) no reaction
- 25. Which of the following alkyl halides is unreactive in an S_N 2 reaction?
 - (a) Primary

(b) Secondary

(c) Tertiary

- (d) Methyl chloride
- **26.** The maximum yield of chloromethane (methyl chloride) by the chlorination of methane is obtained by
 - (a) using a considerable excess of methane in the reaction mixture
 - (b) using a large excess of chlorine in the reaction mixture
 - (c) using a 1:1 mixture of methane and chlorine
 - (d) carrying out the chlorination in the presence of UV light
- 27. The final product obtained in the reaction

is









28. Which of the following is the final product in the reaction between benzoyl chloride and phenyl magnesium bromide?

(a)
$$C_6H_5$$
 C_6H_5C —OMgBr

(c) C₆H₅COOH

- (d) $(C_6H_5)_2COH$
- 29. The major product obtained in the reaction of propyl benzene with chlorine in the presence of UV radiation is
 - (a) 1-chloro-1-phenylpropane
 - (b) 2-chloro-1-phenylpropane
 - (c) 3-chloro-1-phenylpropane
 - (d) none of these
- 30. Which of the following sequence of reactions will give 1-bromo-4-trichloromethylbenzene?

(a) Toluene
$$\xrightarrow{\text{Br}_2}$$
 o -bromotoluene $\xrightarrow{\text{Cl}_2}$ hv or heat \rightarrow

(b) Toluene $\xrightarrow{\text{Fe}}$ p -bromotoluene $\xrightarrow{\text{Cl}_2}$ hv or heat \rightarrow

- (c) Toluene $\xrightarrow{\text{Cl}_2}$ trichloromethyl benzene $\xrightarrow{\text{Br}_2}$ Fe
- (d) none of these
- 31. The bond dissociation energy of the C—X bond in CH₃—F, CH₃—Cl and CH₂=CHCH₂—Cl where X is F or Cl is of the order
 - (a) CH_3 — $F > CH_3$ — $Cl > CH_2$ = $CHCH_2$ —Cl
 - (b) CH₂=CHCH₂Cl > CH₃—Cl > CH₃—F
 - (c) CH₃—Cl > CH₃—F > CH₂=CHCH₂Cl
 - (d) The C—X dissociation energy in all the compounds is of the same order.

32. The products obtained in the reaction

$$CH_2=CH-CH_2Cl + RMgX \longrightarrow$$

are

- (a) CH₂=CH—CH₂MgX and RCl
- (b) CH₂=CHCH₂R and MgXCl
- (c) CH₂=CH—CH₂—CH₂CH=CH₂
- (d) none of these
- 33. The halogen derivatives of alkenes are known as
 - (a) alkyl halides

- (b) alkenyl halides
- (c) alkynyl halides
- (d) aryl halides
- **34.** In unimolecular nucleophilic substitution, alkyl halides react via the carbocation intermediate. The order of reactivity of the carbocations is
 - (a) 3 > 2 > 1

(b) $\hat{1} > \hat{2} > \hat{3}$

(c) $\stackrel{\circ}{2} > \stackrel{\circ}{1} > \stackrel{\circ}{3}$

- (d) $3^{\circ} = 1^{\circ} > 2^{\circ}$
- **35.** In bimolecular nucleophilic substitution, alkyl halides undergo hydrolysis through the formation of an intermediate product. The reactivity of the alkyl halides is in the order
 - (a) $\mathring{1} > \mathring{2} > \mathring{3}$

(b) 3 > 2 > 1

(c) $\stackrel{\circ}{2} > \stackrel{\circ}{1} > \stackrel{\circ}{3}$

- (d) 3 = 1 > 2
- **36.** In S_N2 reactions, the order of reactivity of the halides CH_3X , C_2H_5X , n- C_3H_7X , n- C_4H_9X is
 - (a) $CH_3X > C_2H_5X > n-C_3H_7X > n-C_4H_9X$
 - (b) $C_2H_5X > n-C_3H_7X > n-C_4H_9X > CH_3X$
 - (c) $C_2H_5X > n-C_3H_7X > n-C_4H_9X > CH_3X$
 - (d) $n-C_4H_9X > n-C_3H_7X > C_2H_5X > CH_3X$
- **37.** In a nucleophilic substitution reaction for a given alkyl group, the order of reactivity is
 - (a) R-I > R-Br > R-Cl > R-F (b) R-F > R-I > R-Br > R-Cl
 - (c) R-Cl>R-Br>R-I>R-F (d) R-F>R-Cl>R-Br>R-I
- **38.** The elimination of HX from an alkyl halide forms an alkene. The order of the elimination reactions is
 - (a) $\mathring{3}$ halide $> \mathring{2}$ halide $> \mathring{1}$ halide
 - (b) $\mathring{1}$ halide $> \mathring{2}$ halide $> \mathring{3}$ halide

		· ·		
	(c)	$\mathring{1}$ halide = $\mathring{2}$ halide > $\mathring{3}$ halide		
	(d)	$\overset{\circ}{2}$ halide $>\overset{\circ}{1}$ halide $>\overset{\circ}{3}$ halide		
39.	Chlor	ine is the most reactive toward	s aqı	ueous NaOH in
		methyl chloride		chlorobenzene
	(c)	vinyl chloride	(d)	benzyl chloride
40.		ine is least reactive in	(la)	athril ahlawida
	. ,	methyl chloride allyl chloride		ethyl chloride vinyl chloride
41.		-		ne by making the latter react with
		cuprous chloride		the by making the latter react with
		hydrochloric acid		
		nitrous acid followed by heat	_	_
40		chlorine in the presence of ah	-	
42.		n of the following can function HCHO + HCl		ormyl chloride in formylation? CO + HCl
	` '	HCOOCH ₃ + HCl	` /	HCONH ₂ + HCl
43.	Treati		opan	e with alcoholic KOH gives
		3-phenylpropene	(b)	1-phenylpropene
	(c)	1-phenylpropan-3-ol	(d)	1-phenylpropan-2-ol
44.		action with alcoholic KOH, 1-c		
		1-butene 1-butanol		2-butene 2-butanol
15	()		` '	
43.	-	onitrile may be prepared by he propyl alcohol with KCN		_
		propyl chloride with KCN		-
46.	Chlor	oform is slowly oxidized by air	r in t	he presence of light to form
		formyl chloride	` '	trichloroacetic acid
		formic acid		phosgene
47.		mechanism. Among the follow		ine proceeds by the free-radica, which is the chain-terminating
	(a)	$Cl_2 \longrightarrow 2Cl^{\bullet}$		
	(b)	$CH_3Cl + Cl^{\bullet} \longrightarrow {}^{\bullet}CH_2Cl$	+ HC	
	(c)	$^{\bullet}$ CH ₃ + Cl ₂ \longrightarrow CH ₃ Cl + C	Cl•	
	(d)	$Cl^{\bullet} + Cl^{\bullet} \longrightarrow Cl_2$		

(a) Chain

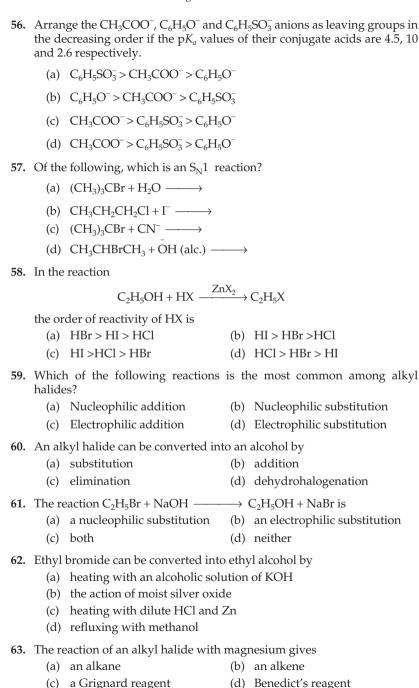
48. What is the type of isomerism exhibited by CH₃CHCl₂ and CH₂ClCH₂Cl? (b) Functional (c) Position

49. Which of the following, on being heated with alcoholic KOH, will

respond positively to the carbylamine test? (a) Chloroform and silver powder (b) Chloroform and aniline (c) Methyl chloride and aniline (d) Methyl cyanide and aniline

(d) Metamerism

50.	Carbon tetrachloride does not have a dipole moment due to (a) its regular tetrahedral structure (b) its planar structure			
	(/	ectron affinities of ca ze of the carbon and		
51.	silver nitrate.		rder of reactivity with alcoholic	
	$C_6H_5CH_2CH_2Br$ (a) II > I > III (c) III > II > I	(b)	(II) and $C_6H_5CH=CHBr$ (III) I > II > III II = I > III	
52.	silver nitrate or with	n KCN. H ₁₃ Br (II) and C ₆ H ₅ B	r (III) III > II > I	
	(c) I > III > II		I = II > III	
53.	$AgNO_3$.		ding to reactivity with alcoholic	
	Tert. Butyl chloride (a) I > II > III	(I), sec. butyl chlorid	e (II) and CCl ₄ (III)	
	(c) II > I > III	` '	II = III > I	
54.		e (III) according to re (b)	-dinitrochlorobenzene (II) and eactivity with sodium ethoxide. III $> I > II$ II = III $> I$	
55.	 55. Arrange H₂O, OH, CH₃O and CH₃COO in descending order of nucleophilicity (rate of S_N2 reactivity). 			
	(a) $H_2O > \bar{O}H > 0$	$\overline{O}CH_3 > CH_3COO^-$		
	(b) $\overline{OCH}_3 > \overline{OH} > \overline{OH}$	$> CH_3COO^- > H_2O$		
	(c) $\overline{OH} > \overline{OCH_3}$	> H ₂ O > CH ₃ COO		
	(d) $CH_3COO^- > 0$	$OCH_3 > OH > H_2O$		



64. In the reaction

$$RI + Mg \longrightarrow A \xrightarrow{H_2O} propane$$

the alkyl halide is

(a) methyl iodide

(b) ethyl iodide

(c) ethyl bromide

- (d) propyl bromide
- 65. In the reaction $CH_3MgX + CH_3OH \longrightarrow A$, the product formed (A) is
 - (a) an alcohol

(b) acetone

(c) ethane

(d) methane

66. In the reaction

chlorobenzene + Mg
$$\xrightarrow{\text{dry}}$$
 A $\xrightarrow{\text{EtOH}}$ B

the product formed (B) is

(a) ethylbenzene

(b) phenol

(c) benzene

- (d) phenylmethyl ether
- 67. The reaction of ethyl bromide with a lead-sodium alloy gives
 - (a) tetraethyl bromide
- (b) tetraethyl lead
- (c) sodium ethoxide
- (d) none of these
- 68. The Friedel-Crafts reaction of *n*-propyl bromide with benzene in the presence of anhydrous AlCl₃ gives
 - (a) *n*-propyl benzene
- (b) isopropyl benzene
- (c) 1,4-dipropyl benzene (d) 1,2-dipropyl benzene
- 69. Treatment of ammonia with excess ethyl chloride will give
 - (a) diethylamine
 - (b) ethane
 - (c) methylamine
 - (d) tetraethyl ammonium chloride
- 70. The mixture of two organic chlorine compounds, on treatment with sodium metal in ether solution, gives isobutane as one of the products. The reactants are
 - (a) methyl chloride and propyl chloride
 - (b) methyl chloride and ethyl chloride
 - (c) isopropyl chloride and ethyl chloride
 - (d) isopropyl chloride and methyl chloride
- 71. The reaction of tert. butyl bromide with sodium methoxide gives
 - (a) isobutane

(b) tert. butylmethyl ether

(c) isobutylene

(d) sodium tert, butoxide

72. In the reaction $CH_3CH_2CHBrCH_3 + (CH_3)_3COK \longrightarrow$ the main product is

	(a)	CH ₃ CH ₂ CHCH ₃	(b)	CH ₃ CH ₂ CHCH ₃
		$OC(CH_3)_3$		ОН
	(c)	CH ₃ CH ₂ CH=CH ₂	(d)	CH ₃ CH=CHCH ₃
73.	(a) (b) (c)	of the following has the high o -Dichlorobenzene m -Dichlorobenzene p -Dichlorobenzene All have the same melting po		nelting point?
74.	treatn			unsaturated hydrocarbon on anal on reaction with aqueous
	(a)	CH ₃ CHCl ₂	(b)	CH ₂ ClCH ₂ Cl
	(c)	a mixture of (a) and (b)	(d)	none of these
75.	substi Chlor 2,4-di (a)	ge the following in order tution reaction. obenzene (I), 2,4,6-trinitrochloritro-chlorobenzene (III) and I < IV < III < II II < III < IV < I	robei 4-nit (b)	
76.	The fu	usion of chlorobenzene with so	lid N	JaOH gives
		benzene		benzoic acid
	(c)	no reaction	(d)	phenol
77.	The a	ction of chloral on chlorobenze	ene g	ives
	(a)	BHC (b) DDT	(c)	gammexene (d) lindane
78.	Benzy	alcohol on reaction with PC	l ₅ giv	es
	` '	benzene	()	toluene
		benzyl chloride	` '	none of these
79.	In the		(aq.)	\longrightarrow X + Y, the compounds X
	(a)	C ₆ H ₆ and KCl	(b)	C ₆ H ₅ CH ₃ and KCl
	(c)	C ₆ H ₅ CH ₂ CN and KCl	(d)	none of these
80.	In the		C ₆ H	$_{5}CH_{2}NH_{2} + Y$, the compounds X
	(a)	HNO ₃ and HCl	(b)	NH ₃ and HCl
	(c)	HNO ₂ and H ₂ O	(d)	none of these

81.	The oxidation of benzyl chloride with Pb(NO ₃) ₂ gives			
	(a)	benzoic acid	(b)	benzene
	(c)	benzaldehyde	(d)	none of these
82.	Treati	ment of ethylidene chloride wit	h ag	ueous KOH gives
	(a)	ethylene glycol	(b)	acetaldehyde
	(c)	formaldehyde	(d)	none of these
83.	On di	stillation with bleaching powd	er, e	thyl alcohol gives
	(a)	acetone	(b)	trichloroacetone
	(c)	chloroform	(d)	acetic acid
84.	Chlor	oform is stored in bottles in the	pre	sence of
		CH ₃ COOCH ₃	. ,	C_2H_5OH
		CH₃COOH		none of these
85.	In the	reaction $2CHCl_3 + O_2 - X$	→ 20	$COCl_2 + 2HCl$, X is
		an oxidizing agent	(b)	a reducing agent
	(c)	light and air	(d)	none of these
86.		ing heated with aniline and KO	ΟH,	chloroform gives
	` '	an almond-like smell		
		a rose-like odour a smell like oil of wintergreen		
		an offensive smell		
87.	On be	ing warmed with silver powde	er, ch	nloroform gives
		C_6H_6		C_2H_4
	(c)	C_2H_2	(d)	CH ₃ Cl
88.	The re	eaction of chloroform with con-	centi	ated HNO3 gives
		CHCl ₂ NO ₂		CHCl ₂ HNO ₃
	(c)	CCl ₃ NO ₂	(d)	none of these
89.	Chlor	oprin is		
		$C_2H_5C(NO)_5SH$	(b)	CCl₃CHO
		CCl ₃ NO ₂		CCl ₃ NO ₃
90	Chlor	opicrin is used as		
<i>7</i> 0.	(a)	=	(b)	an insecticide
	` '	a hypnotic drug		all of these
91.	The f	inal product formed upon th	e hy	drolysis of CHCl ₃ by aqueous
	KOH	IS HCOOH	(b)	HCOOK
	` '	CH ₃ OH	` '	none of these
	\ /	9	` /	

92.	rne re	eaction of chioroform with ace	tone	gives
	(a)	mesitylene	(b)	ethylidene chloride
	(c)	chloretone	(d)	chloral
93.	Iodofo	orm is used as an		
	(a)	anaesthetic	(b)	analgesic
	(c)	antiseptic	(d)	antifebrin
94.	CCl ₄ i	s insoluble in H ₂ O because		
	(a)	H ₂ O is polar	(b)	CCl ₄ is nonpolar
	(c)	H ₂ O and CCl ₄ are polar	(d)	none of these
95.	Which	n of the following is known as	freor	n?
	(a)	CCl ₂ F ₂	(b)	CHCl ₃
	(c)	CH_2F_2	(d)	CF_4
96.	CCl ₄ i	s used as a fire extinguisher be	ecaus	se
	(a)	of its covalent bond		
	(b)	of its low b.p.		
	(c)	of its high m.p.		
	(d)	it gives incombustible vapou	rs	
97.	CCl ₄ i	s used in fire extinguishers un	der t	he name
	(a)	pyrene	(b)	phosphine
	(c)	phosgene	(d)	none of these
98.	The re	eactivities of CH ₃ Cl, CH ₃ CH ₂ C	H_2C	l and C ₆ H ₅ Cl are in the order
	(a)	$CH_3Cl > CH_3CH_2CH_2Cl > C_6$	H ₅ Cl	
	(b)	$CH_3CH_2CH_2Cl > CH_3Cl > C_6$	H ₅ Cl	
	(c)	$C_6H_5Cl > CH_3CH_2CH_2Cl > Cl$	H ₃ Cl	
	(d)	$CH_3Cl > C_6H_5Cl > CH_3CH_2Cl$	H ₂ Cl	
99.				cleophilic substitution reactions
		are alkyl halides. This is due to		According to a
		the formation of the less stab	ie cai	bonium ion
		resonance stabilization		
		their longer C-halogen bond		
	` ′	the inductive effect		
100.		e possible structures for the motically active?	olecu	ılar formula $C_5H_{11}Br$, how many
	_	One	(b)	Three
		Five		Four

- **101.** Of the three isomeric dihaloarenes *o-*, *m-* and *p-*, which has the highest m.p.?
 - (a) o-

(b) m-

(c) p-

- (d) They have the same m.p.
- **102.** Which of the following will be the least reactive towards nucleophilic substitution?
 - (a) CH₂Cl
- (b) Cl
- (c) CI
- (d) C_2H_5Cl
- 103. For a given alkyl group, the boiling points of alkyl halides follow the order
 - (a) RCl > RBr > RI
- (b) RI > RBr > RCl
- (c) RI > RCl > RBr
- (d) RBr > RI > RC1

104. In the reaction

$$CH_3CHCl_2 \xrightarrow{aq. KOH} X$$

X is

(a) CH₃CH₂OH

(b) (CH₃)₂CO

(c) CH₂—CH₂

- (d) CH₃CHO
- **105.** The trade name of trichloroethylene is
 - (a) Freon

(b) Westron

(c) Westrosol

- (d) DDT
- 106. Chlorobenzene can be obtained from benzene diazonium chloride by the
 - (a) Friedel–Crafts reaction
- (b) Gattermann reaction
- (c) Wurtz reaction
- (d) Fittig reaction
- 107. The main product of the reaction $(CH_3)_3CCl \xrightarrow{aq. KOH}$ is

(b)
$$CH_3$$
— C = $CHOH$ CH_3

- 108. Chlorination of toluene in the presence of light and heat followed by treatment with aqueous NaOH gives
 - (a) o-cresol

- (b) p-cresol
- (c) 2,4-dihydroxytoluene
- (d) benzoic acid
- **109.** Which of the following can give only two monochloro derivatives?
 - (a) n-Hexane

- (b) 2,4-Dimethylpentane
- (c) 2-Methylpropane
- (d) Benzene

110. In the reaction

$$CH_3CH_2I \xrightarrow{\quad alc. \; KOH \quad} X \xrightarrow{\quad Br_2 \quad} Y \xrightarrow{\quad KCN \quad} Z$$

Zis

- (a) CH₃CH₂CN
- (b) CH₂BrCH₂CN
- (c) CNCH₂CH₂CN
- (d) BrCH=CHCN

111. In the reaction

$$\begin{array}{c} CH_3CHCH_3 \xrightarrow{\quad alc. \ KOH \quad} A \xrightarrow{\quad peroxide \quad} B \xrightarrow{\quad acetone \quad} C \\ Br \end{array}$$

C is

- (a) CH₃CH₂CH₂I
- (b) CH₃—CH—CH₃

(c)
$$CH_3$$
— CH — CH_2I (d) CH_3CH = CHI

- 112. Which of the following will react according to S_N2 as well as S_N1 mechanisms?
 - (a) CH₂CH₂Cl

(b) (CH₃)₂CHCl

(c) (CH₃)₃CCl

- (d) CH₃CH₂CH₂Cl
- 113. What is the order of the ease of formation of the following carbocations?
 - (a) $3 > 2 > 1 > CH_2$
- (b) $\overset{+}{C}H_3 > \overset{\circ}{1} > \overset{\circ}{2} > \overset{\circ}{3}$
- (c) $1 > 2 > 3 > CH_2$
- (d) $\overset{+}{C}H_2 > \overset{\circ}{3} > \overset{\circ}{2} > \overset{\circ}{1}$

114. How many optically active isomers are possible for compounds with the

	molec	ular formula	$1 C_5 H_{11}$	Br?						
	(a)	Two	(b)	Гhree	(c)	Four	((d)	Five	
115.	Ā: an	ission of the d B ⁺ may be homolytic b	attribu	ited to	ound	А—В	to give	the	intern	nediate
		heterolytic								
	(c)	homolytic a	as well	as heterol	ytic bo	nd fiss	ion			
	(d)	none of the	se							
116.	Arrar	ge the follov	ving in	decreasin	g orde	er of C-l	haloger	n boı	nd leng	gth.
	(a)	$CH_3I > CH_3$	Br > C	$H_3Cl > CH$	I_3F					
	(b)	$CH_3F > CH$	$_{3}$ Br > C	$CH_3I > CH_3$	_s Cl					
	(c)	$CH_3Cl > CH$	$H_3Br >$	$CH_3I > CH$	I_3F					
	(d)	$CH_3I > CH_3$	$_{3}Cl > C$	$H_3Br > CH$	I_3F					
117.	The re	eaction CH ₃ N give	⁄IgBr +	CH ₃ C≡C	Н	\longrightarrow				
		methane				ethane				
	(c)	propane			(d)	isopro	pane			
118.		I effect of the								
	CH ₃ —	− CH₃CF II	I ₂ —	(CH ₃) ₂ C III	CH—		I ₃) ₃ C— IV			
	-	ases in the or	der	111			LV			
	(a)	I > II > III >	IV		(b)	IV > II	I > II >	Ι		
	(c)	I > IV > III	> [[(d)	II > I >	$III > \Gamma$	V		
119.	chlore	ease of deh bethane (I), 2 he order								
	(a)	I < II < III			(b)	I > II >	· III			
	(c)	II > I > III			(d)	I > III	> II			
120.	Which	n of the follo			hilic a	dditior	ı reacti	ons	?	
	(a)	CH ₃ C=C) + HC	N						
	()	CH ₃								
	(b)	CH ₃ CH=C	$H_2 + H$	$Br \longrightarrow$						
	(c)	CH ₃ CH=C	$H_2 + H$	Br <u>peroxi</u>	de →					

(d) $CH_3CONH_2 \xrightarrow{Br_2/KOH}$

- 121. How many isomers are possible for compounds having the molecular formula C₅H₁₁Br?

 (a) Five
 (b) Six
 (c) Seven
 (d) Eight

 122. The boiling points of the haloalkanes CH₃I, CH₃Br, CH₃Cl and CH₃F decrease in the order

 (a) CH₃I > CH₃Br > CH₃Cl > CH₃F
 (b) CH₃F > CH₃Cl > CH₃Br > CH₃I
 (c) CH₃Br > CH₃Cl > CH₃F > CH₃I
- (d) The boiling points of haloalkanes follow no particular order.123. The boiling points of methyl bromide (I), ethyl bromide (II), *n*-propyl bromide (III) and *n*-butyl bromide (IV) decrease in the order
 - (a) I > II > III > IV

(b) IV > III > II > I

(c) I > III > II > IV

- (d) III > IV > I > II
- **124.** The stabilities of alkyl fluorides (I), alkyl chlorides (II), alkyl bromides (III) and alkyl iodides (IV) decrease in the order
 - (a) I > II > III > IV

- (b) IV > III > II > I
- (c) I > IV > II > III
- (d) II > I > IV > III
- 125. The strengths of carbon-halogen bonds follow the order
 - (a) R F > R Cl > R Br > R I
 - (b) R-I > R-Br > R-Cl > R-F
 - (c) R F > R I > R Br > R Cl
 - (d) R-Cl > R-Br > R-I > R-F
- **126.** In the elimination reactions, the reactivities of alkyl halides follow the order
 - (a) R F > R Cl > R Br > R I (b) R Cl > R Br > R I > R F
 - (c) R-I > R-Br > R-Cl > R-F (d) R-Br > R-I > R-Cl > R-F
- **127.** The stabilites of alkenes follow the order
 - (a) $R_2C = CH_2 > R_2C = CHR > RCH = CHR > CH_2 = CH_2$
 - (b) $R_2C = CH_2 > CH_2 = CH_2 > R_2C = CHR > RCH = CHR$
 - (c) $CH_2 = CH_2 > RCH = CHR > R_2C = CHR > R_2C = CH_2$
 - (d) All alkenes are equally stable.
- 128. In the reaction

$$R$$
—Br + Cl^- dimethyl formamide R — $Cl + Br^-$

the relative rates of reaction of methyl bromide (I), ethyl bromide (II), isopropyl bromide (III) and tert. butyl bromide (IV) follow the order

(a) I > II > III > IV

(b) IV > III > II > I

(c) IV > I > II > III

(d) II > III > IV > I

129. In the reaction

$$R \longrightarrow Br + Cl^- \longrightarrow R \longrightarrow Cl + Br^-$$

the rates of reaction of ethyl bromide (I), *n*-propyl bromide (II), isobutyl bromide (III) and neopentyl bromide (IV) follow the order

(a) IV > III > II > I

(b) I > II > III > IV

(c) I > III > II > IV

- (d) III > II > IV > I
- **130.** The amounts of energy needed to form the carbocations $\overset{\scriptscriptstyle \star}{C}H_3$, $\overset{\scriptscriptstyle \star}{1}$, $\overset{\scriptscriptstyle \star}{2}$ and $\overset{\scriptscriptstyle \star}{3}$ follow the order
 - (a) $\overset{+}{C}H_3 > \overset{\circ}{1} > \overset{\circ}{2} > \overset{\circ}{3}$
- (b) $3 > 2 > 1 > CH_3$ (d) $2 > 3 > 1 > CH_3$
- (c) $\hat{1} > \hat{2} > \hat{3} > \hat{C}H_2$

• Type 2 •

Choose the correct options. More than one option is correct.

- 131. 2-phenyl-2-hexanol can be prepared by Grignard synthesis. Which of the following pair of compounds can one start with?
 - (a) $CH_3CH_2CH_2CH_2Br$ and $CH_3-C_6H_5$
 - (b) CH_3CH — CH_2Br and C— C_6H_5 CH_3 CH_3
 - (c) CH₃CH₂CH₂CH₂—CC₆H₅ and C₆H₅Br
 - (d) None of these
- 132. Which of the following reagents can be used to distinguish CH₂=CHCH₂Cl from CH₃CH₂CH₂Cl?
 - (a) Na fusion; HNO_3 , $AgNO_3$ (b) $Ag(NH_3)_2OH$
 - (c) $AgNO_3/C_2H_5OH$
- (d) Br_2/CCl_4

133. In the reaction

$$R$$
— $COOAg + Br_2 \longrightarrow RBr + CO_2 + AgBr$

the reaction proceeds through the intermediate formation of

(d) none of these

134. Alkyl iodides can be prepared by

(a)
$$RCH_2COOAg + I_2 \xrightarrow{CCl_4} RCH_2I$$

(b)
$$RCH_2Cl + NaI \xrightarrow{acetone} RCH_2I + NaCl$$

- (c) R—OH + HI \longrightarrow $RI + H_2O$
- (d) $CH_4 + I_2 \longrightarrow CH_3I$

135. Which of the following reactions are not feasible?

(a)
$$Cl$$
 $NaNH_2,NH_3$ NH_2

(b) Br
$$CH_3$$
 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3

(c) Br
$$CH_3$$
 CH_3 CH_3 CH_3 CH_3 CH_3

(d)
$$CI$$
 CH_2CH_3 $NaNH_2, NH_3$ NH_2

136. Which of the following are expected to have dipole moments?

- (a) *cis-*1,2-dichloroethene
- (b) trans-1,2-dichloroethene
- (c) cis-1,2-dibromoethene
- (d) trans-1,2-dibromoethene

- 137. Which of the following are expected to have dipole moments?
 - (a) Tetrachloroethene
- (b) Chloroform
- (c) Sulphur dioxide
- (d) Carbon dioxide
- **138.** Which of the following are Lewis bases?
 - (a) AlCl₃

(b) H+

(c) Br₂

- (d) R-O-H
- 139. The treatment of tertiary butyl chloride with 80% aqueous ethanol at 25°C gives
 - (a) $(CH_3)_3C$ —OH

- (b) (CH₃)₃C—OCH₂CH₃
- (c) $CH_2 = C < CH_3$ CH_2
- (d) none of these
- 140. Which monosubstituted product does the chlorination of isopentane [(CH₃)₂CHCH₂CH₃] with chlorine at 300°C give?

(a)
$$CICH_2$$
— C — CH_3 — CH_3 — CH_3 — C — CH_2 — CH_2 — CH_2 CI C

(c)
$$CH_3$$
 CH_3 CH_3 (d) CH_3 CH_2CH_3 CH_3

- 141. The treatment of ethyl bromide with alcoholic silver nitrite gives
 - (a) ethane

(b) ethene

(c) nitroethane

- (d) ethyl nitrite
- **142.** Acetyl chloride does not react with
 - (a) ethanol

(b) ethanal

(c) diethyl ether

- (d) propanal
- 143. Of the following, which are $S_N 2$ reactions?
 - (a) $CH_3CH_2CI + I^- \longrightarrow$
 - (b) $(CH_3)_3CBr + CN^-$ (alc.) \longrightarrow
 - (c) $CH_3CHBrCH_3 + OH^-(aq.) \longrightarrow$
 - (d) $CH_3CHBrCH_3 + OH^-$ (alc.) \longrightarrow

144. Which of the following reagents can be used to prepare an alkyl halide

	110111	ari arcorior:				
	(a)	NaCl	(b)	$HCl + ZnCl_2$		
	(c)	SOCl ₂	(d)	PCl ₅		
145.	Aryl l	nalides undergo				
	(a)	the Fittig reaction	(b)	the Ullmann reaction		
	(c)	the Grignard reaction	(d)	none of these		
146.	Which	h of the following are organom	etall	ic compounds?		
	(a)	C ₃ H ₇ MgI	(b)	C ₂ H ₅ ONa		
	(c)	$(CH_3)_3Al$	(d)	TEL		
147.	Which C ₂ H ₅ I		epict	the nucleophilic substitution of		
	(a)	$C_2H_5Br + C_2H_5SNa \longrightarrow C_2H_5Br + C_2H_5Br + C_2H_5SNa \longrightarrow C_2H_5Br + C_2H_5$	C_2H_5S	$SC_2H_5 + NaBr$		
	(b)	$C_2H_5Br + 2H \longrightarrow C_2H_6 + HBr$				
	(c)	$C_2H_5Br + AgCN \longrightarrow C_2H_5$	I ₅ NC	+ AgBr		
	(d)	$C_2H_5Br + KOH \longrightarrow C_2H_5$	OH-	+ KBr		
148.	For ar	n S_N 2 reaction, which of the fol	lowi	ng statements are true?		
	(a)	The rate of reaction is indeputed nucleophile.	pend	ent of the concentration of the		
	(b)	The nucleophile attacks the opposite to the group being of		om on the side of the molecule aced.		

(c) The reaction proceeds with simultaneous bond formation and

149. Which of the following reactions can be used for the preparation of alkyl

150. Which of the following compounds, on being warmed with iodine

(b) CH₃COCH₃

(d) CH₃OH

(a) $CH_3CH_2OH + HCl \xrightarrow{anhyd. ZnCl_2}$ (b) $CH_3CH_2OH + HCl \xrightarrow{}$

(d) $(CH_3)_2CHOH + HCl \xrightarrow{anhy.} Z_nCl.$

solution and NaOH, will give iodoform?

(c) (CH₃)₃COH + HCl ————

rupture.
(d) None of these

(a) CH₃CH₂OH

halides?

(c) OR-

158. Which of the following will give vinyl chloride?

(b) $CH_2=CH_2+Cl_2\longrightarrow CH_2$ CH_2 CH_2 C

(a) $CH = CH + HCl \xrightarrow{Hg^{2+}}$

151.	Vinyl	chloride undergoes		
	(a)	addition reactions	(b)	elimination reactions
	(c)	substitution reactions	(d)	none of these
152.		n of these statements are true f de and ethylidene chloride?	or th	e isomeric compounds ethylene
	(a)	Both react with aqueous KOF	I to g	give the same product.
	(b)	Both react with alcoholic KOI	H to	give the same product.
	(c)	They are derivatives of ethan	e.	
	(d)	They respond to Beilstein's te	st.	
153.	The re	eaction of ethyl alcohol and ble	achi	ng powder gives
	(a)	acetaldehyde	(b)	chloroform
	(c)	chloral	(d)	none of these
154.		n of the following occurs du DH and bleaching powder?	ıring	the formation of CHCl ₃ from
	(a)	Oxidation	(b)	Reduction
	(c)	Hydrolysis	(d)	Chlorination
155.	Which	n of the following statements a	re tri	ue about chloroform?
	(a)	It is used as an anaesthetic.		
	(b)	It is used as a solvent.		
	(c)	It has sp ² -hybridized carbon.		
	(d)	It has a distorted tetrahedral	shap	e.
156.	Which acetyl		can l	be used for the preparation of
	(a)	CH ₃ CHCl ₂ + alc. KOH ———	\rightarrow	
	(b)	ClCH ₂ CH ₂ Cl+ alc. KOH —	\longrightarrow	
	(c)	$ClCH_2CH_2Cl + NaNH_2$ liq.	$\frac{NH_3}{\Delta}$	\rightarrow
	(d)	None of these		
157.	The h		n be	easily replaced by nucleophiles
	(a)	CN ⁻	(b)	NO_2^-

(d) none of these

(c)
$$CH_2 = CH_2 + Cl_2 \xrightarrow{600^{\circ}C}$$

(c)
$$CH_2=CH_2+Cl_2\xrightarrow{600^{\circ}C}$$

(d) $CICH_2\longrightarrow CH_2Cl\xrightarrow{KOH}$

159. Which of the following is an S_N 2 reaction?

(a)
$$CH_3CH_2Br + KOH \longrightarrow CH_3CH_2OH + KBr$$

(b)
$$CH_3CH_2Br + CH_3CH_2ONa \longrightarrow CH_3CH_2OCH_2CH_3 + NaBr$$

(c)
$$(CH_3)_3CBr + KOH \longrightarrow (CH_3)_3COH + KBr$$

$$\begin{array}{c} CH_3 & CH_3 \\ | & | \\ (d) & CH_3-CH_2-C-Br+KOH-\longrightarrow CH_3CH_2-C-OH+KBr \\ | & | \\ CH_3 & CH_3 \end{array}$$

160. Which of the following statements are correct?

- (a) An S_N1 reaction proceeds with inversion of configuration.
- (b) An S_N2 reaction proceeds with stereochemical inversion.
- (c) An S_N2 reaction follows second-order kinetics.
- (d) The reaction of tert. butyl bromide with OH follows first-order kinetics.

	Answers				
1. d	2. a	3. b	4. d	5. a	
6. c	7. b	8. b	9. a	10. d	
11. a	12. c	13. d	14. d	15. b	
16. c	17. c	18. a	19. a	20. b	
21. a	22. a	23. b	24. b	25. c	
26. a	27. b	28. d	29. a	30. c	
31. a	32. b	33. b	34. a	35. a	
36. a	37. a	38. a	39. d	40. d	
41. c	42. b	43. b	44. a	45. c	
46. d	47. d	48. c	49. b	50. a	
51. a	52. a	53. a	54. c	55. b	
56. a	57. a	58. b	59. b	60. a	
61. a	62. b	63. c	64. d	65. d	
66. c	67. b	68. b	69. d	70. d	
71. c	72. d	73. c	74. a	75. a	
76. d	77. b	78. c	79. c	80. b	
81. c	8 2. b	83. c	84. b	85. c	

87. c	88. c	89. c	90. b
92. c	93. c	94. b	95. a
97. a	98. a	99. b	100. b
102. c	103. b	104. d	105. c
107. d	108. d	109. c	110. c
112. b	113. a	114. b	115. b
117. a	118. b	119. a	120. a
122. a	123. b	124. a	125. a
127. a	128. a	129. b	130. a
132. c, d	133. a, b, c	134. b, c	135. b, c, d
137. a, c	138. c, d	139. a, b, c	140. a, b, c, d
142. b, d	143. a, c	144. b, c, d	145. a, b, c
147. a, c, d	148. b, c	149. a, c, d	150. a, b, c
152. b, c, d	153. a, b, c	154. a, c, d	155. a, b, d
157. a, b, c	158. a, b, c, d	159. a, b	160. b, c, d
	92. c 97. a 102. c 107. d 112. b 117. a 122. a 127. a 132. c, d 137. a, c 142. b, d 147. a, c, d 152. b, c, d	92. c 93. c 97. a 98. a 102. c 103. b 107. d 108. d 112. b 113. a 117. a 118. b 122. a 123. b 127. a 128. a 132. c, d 133. a, b, c 137. a, c 138. c, d 142. b, d 143. a, c 147. a, c, d 148. b, c 152. b, c, d 153. a, b, c	92. c 93. c 94. b 97. a 98. a 99. b 102. c 103. b 104. d 107. d 108. d 109. c 112. b 113. a 114. b 117. a 118. b 119. a 122. a 123. b 124. a 127. a 128. a 129. b 132. c, d 133. a, b, c 134. b, c 137. a, c 138. c, d 139. a, b, c 142. b, d 143. a, c 144. b, c, d 147. a, c, d 148. b, c 149. a, c, d 152. b, c, d 153. a, b, c 154. a, c, d

Hints to More Difficult Problems

- Halogens activated by strong electron-withdrawing groups from o- and p-positions are displaced by a bimolecular nucleophilic substitution reaction.
- **2.** Chlorocyclohexane behaves like an aliphatic halogen-substituted hydrocarbon, and can, therefore, react with alcoholic AgNO₃.
- 3. N-Bromosuccinimide is used for allylic bromination.
- 7. 2,3-Dimethylbutane has 18 primary hydrogens (which are all equivalent) and two tertiary hydrogens (which are equivalent), and therefore, gives only two monosubstituted derivatives.
- 8. Answer this question along the same lines as Q. 7.
- 9. The rate of substitution of alkyl halides follows the order 1>2>3 for an S_N2 path. Compounds shown in options (a) and (b) are both 1 alkyl halides. The compound shown in option (b), however, is less reactive due to the steric hindrance of the methyl group.
- 11. The course of elimination in case of alkyl halides is determined by the Saytzev rule, i.e., H is eliminated preferentially from the carbon atom that is bonded to fewer hydrogen atoms, and thus the highly substituted alkene is the major product.
- **32.** A Grignard reagent can participate in $S_N 2$ reactions by furnishing R^- : as a nucleophile. Such displacements with a Grignard reagent do not proceed well with alkyl halides, but only with allyl halides.

- **34.** The transmission of the negative charge from C to a halogen (as the number of the alkyl groups increases) promotes the ionization of the halide and hence facilitates the S_N1 mechanism. So a 3 halide is the most reactive and a 1 halide the least.
- **35.** For an S_N2 attack, the enhanced inductive effect of the larger number of methyl groups in a $\ref{3}$ alkyl halide may be expected to make the carbon atom less positively polarized than in the other alkyl halides. However, steric factors are more important because an incoming $\ref{O}H$ ion will find it more difficult to attack the $\ref{3}$ carbon atom than the $\ref{1}$ carbon atom because the former is heavily substituted.
- 53. The reactivity of an alkyl halide towards alcoholic AgNO₃ depends on the ability of the alkyl halide to ionize.
- **59.** See the hint to Q. 1.
- **68.** The 1 carbonium ion isomerizes to the more stable 2 carbonium ion, which in turn reacts with benzene in the Friedel–Crafts reaction to give isopropyl benzene.
- 75. See the hint to Q. 1.
- 94. Nonpolar compounds are insoluble in water.
- **109.** See the hint to Q. 7.
- **133.** This is an example of a Hunsdiecker reaction.
- **135.** For the formation of benzyne intermediate, it is essential that an H should be available in an ortho position to the halogen.

Aromatic Hydrocarbons or Arenes

• Type 1 •

Choose the correct option/s. Only one option is correct.

1. The conversion

$$CH_2CH_2CH_3$$
 Br
 $CH_2CH_2CH_3$

can be effected using

(a) Br_2/CCl_4

(b) Br_2/H_2O

(c) Br₂/Fe

(d) Br₂/benzoyl peroxide

2. Which of the following is expected to be aromatic?









- 3. Why does 1,3-cyclohexadiene undergo dehydrogenation readily?
 - (a) It can be easily reduced.
 - (b) It has no resonance energy.
 - (c) It gains considerable stability by becoming benzene.
 - (d) It cannot undergo dehydrogenation.

- 4. How many different tetrachlorobenzenes are possible?
 - (a) Two

(b) Three

(c) Four

- (d) Five
- 5. How many isomers can tribromobenzene have?
 - (a) One

(b) Two

(c) Three

- (d) Four
- 6. The major product obtained in the reaction

is

(b) COOH

 O_2N

7. The major product obtained in the reaction

$$\begin{array}{c|c}CI\\\hline\\\hline\\CI\end{array}$$

is

(c)
$$CI$$
 (d) CI SO_3H SO_3H SO_3H

8. The major product obtained in the reaction

$$\begin{array}{c|c}
O \\
Br_2 \\
\hline
FeBr_3
\end{array}$$

is expected to be

(a)
$$Br \longrightarrow C \longrightarrow CH_2 \longrightarrow C$$

(b)
$$C - CH_2 - BI$$

$$\begin{array}{c}
\text{(c)} \\
\text{O} \\
\text{II} \\
\text{C} - \text{CH} - \text{O}
\end{array}$$

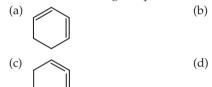
9. The final product obtained in the reaction

$$\begin{array}{c|c} & CH_3CH_2COCI \\ \hline & AlCl_3 \end{array} \longrightarrow \begin{array}{c} X & CH_3NH_2 \\ \hline & H_2, Ni \end{array}$$

is

(a)
$$\begin{array}{c} O \\ H_3C-NH - \\ \end{array} \\ \begin{array}{c} CH_2CH_2CCI \end{array}$$

10. Which of the following compounds is the most stable?



- 11. Cyclobutadiene is said to be
 - (a) aromatic (b) aliphatic (c) nonaromatic(d) none of these
- 12. The reaction

is an example of

- (a) [4+2] cycloaddition
- (b) [2+2] cycloaddition
- (c) either (4+2) or (2+2) cycloaddition
- (d) none of these
- 13. On oxidation with hot alkaline $KMnO_4$, an unknown alkene, C_7H_{12} , gives

The alkene is

(a)
$$CH_3$$
 (b) CH_3

(d) none of these

14. The final product obtained in the reaction

$$\begin{array}{c} \text{CH}_{3} \\ \hline \\ & \frac{1.\,\text{Hg(OAc)}_{2}/\text{THF-H}_{2}\text{O}/20\,\text{sec}}{2.\,\text{NaBH}_{4}(6\,\text{min})} \end{array}$$

is

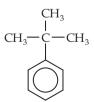
(b)

(c) CH₃

(d)

15. The major product obtained by the Friedel-Crafts reaction of *n*-butylbromide with benzene in the presence of anhydrous AlCl₃ is

(c)



(d) none of these

- 16. The sulphonation of chlorobenzene gives exclusively
 - (a) an o-product

(b) an m-product

(c) a p-product

(d) an o-, p-disubstituted product

17.		eaction of 1-methylcyclopente Alkaline H ₂ O ₂ gives	ne w	rith B ₂ H ₆ followed by treatment
	(a)	cis-2-methylcyclopentanol		
	(b)	<i>trans</i> -2-methylcyclopentanol		
		a mixture (1:1) of cis- and tra	ıns-2	-methyl cyclopentanol
		none of these		3 3 1
10	Carbo	on atoms in benzene are		
10.		sp-hybridized	(h)	sp ² -hybridized
		sp ³ -hybridized		none of these
			(u)	none of these
19.		nzene, the type of bonds are	<i>a</i> .	
		$\sin \sigma$ and nine π		$\sin \sigma$ and $\sin \pi$
	` '	nine σ and three π	(d)	twelve σ and three π
	bonds	3.		
20.	The b	enzene molecule contains		
	(a)	six sp ² -hybrid carbons	(b)	three sp ² -hybrid carbons
	(c)	six sp ³ -hybrid carbons	(d)	three sp ³ -hybrid carbons
21.	Arom	atic compounds burn with a se	ooty	flame because
	(a)		-	
	(b)	they have a ring structure		
	(c)		ercen	tage of hydrogen
	(d)	they resist the action of atmos		
22.	The o	zonolysis of benzene produces	- 3	
		glyoxal		ethylglyoxal
		dimethylglyoxal		methylglyoxal
22		, , ,		, , ,
23.				rgoes ozonolysis followed by lyoxal, monomethylglyoxal and
		hylglyoxal. It could be		, , , , , , , , , , , , , , , , , , , ,
	(a)	benzene	(b)	toluene
	(c)	o-xylene	(d)	<i>m</i> -xylene
24.	How	many monosubstituted produc	cts ca	nn be derived from benzene?
	(a)	One	(b)	Two
	(c)	Three	(d)	Four
25.	All th	e carbon and hydrogen atoms	in b	enzene are in a single plane and
	all the	e C—C bonds are of the same	lengt	h. The C—C—C bond angle is
	(a)	120°	(b)	180°
	(c)	100°	(d)	109°28′
	` '		. ,	

- **26.** Which of the following statements is true for benzene?
 - (a) The monosubstitution of benzene produces two isomeric substances.
 - (b) Benzene undergoes addition reactions because of the presence of double bonds.
 - (c) There is a cyclic delocalization of π -electrons in benzene.
 - (d) There are two types of C—C bonds in the benzene molecule.
- 27. According to the Huckel rule, an aromatic compound has
 - (a) $(4n + 2) \pi$ -electrons
- (b) $(4n + 2\pi)$ electrons
- (c) $(2n + 4) \pi$ -electrons
- (d) $(3n + 3\pi)$ electrons
- 28. The most common reactions that benzene and its derivatives undergo are
 - (a) electrophilic addition reactions
 - (b) nucleophilic addition reactions
 - (c) electrophilic substitution reactions
 - (d) nucleophilic substitution reactions
- 29. Which of the following are electrophilic substitution reactions?
 - (a) Conversion of methyl chloride to methyl alcohol
 - (b) Chlorination of benzene
 - (c) Chlorination of methane
 - (d) Formation of ethane from ethanol
- 30. The nitration of benzene by HNO₃ and H₂SO₄ is
 - (a) an electrophilic substitution (b) a nucleophilic substitution
 - (c) an electrophilic addition
- (d) a free-radical substitution

31. The reaction

$$C_6H_6 \longrightarrow C_6H_5CH_3$$

is a

- (a) Friedel-Crafts reaction
- (b) Wurtz reaction
- (c) Perkin reaction
- (d) Grignard reaction
- 32. In the nitration of benzene, the active nitrating agent is
 - (a) NO_2^-
- (b) NO_2^+
- (c) NO_3^-
- (d) HNO₃
- **33.** Which of the following represents a Friedel–Crafts reaction?

(a)
$$C_6H_6 + C_2H_5C1 \xrightarrow{AlCl_3} C_6H_5C_2H_5 + HC1$$

(b)
$$C_2H_5OH + HC1 \xrightarrow{ZnCl_2} C_6H_5Cl + H_2O$$

(c)
$$C_6H_5Cl + CH_3COCl \xrightarrow{AlCl_3} C_6H_5COOCH_3 + Cl_2$$

(d)
$$C_2H_5Br + Mg \xrightarrow{ether} C_2H_5MgBr$$

- **34.** Which of the following is the most reactive towards electrophilic nitration?
 - (a) Benzene

(b) Toluene

(c) Benzoic acid

- (d) Nitrobenzene
- 35. Which form of xylene is most easily sulphonated?
 - (a) Ortho-
 - (b) Meta-
 - (c) Para-
 - (d) All are sulphonated at the same rate
- 36. Which form of xylene has only one monobromo derivative?
 - (a) Ortho-

(b) Para-

(c) Meta-

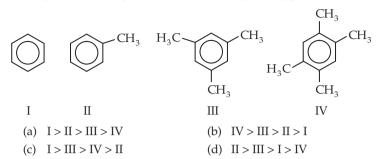
- (d) None of these
- **37.** Arrange C₆H₅CH₃ (I), C₆H₆ (II), C₆H₅COOH (III), C₆H₅NO₂ (IV) in order of decreasing reactivity to ring monobromination.
 - (a) I > III > II > IV
- (b) III > II > IV > I
- (c) I > II > III > IV
- (d) II > I > III > IV
- **38.** Which of the following is a Wurtz–Fittig reaction?

(a)
$$+ 2Na + BrCH_2CH_3 \longrightarrow CH_2CH_3 + 2NaBr$$

(c)
$$SO_3H + H_2O \xrightarrow{130-150^{\circ}C} HCI + H_2SO_4$$

$$(d) \qquad \qquad MgBr \qquad O \\ + ClCCH_2CH_3 \qquad \longrightarrow \qquad COCH_2CH_3$$

39. Arrange the following in order of decreasing boiling point.



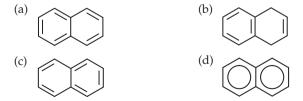
- **40.** The alkylation of benzene with *n*-propyl chloride in the presence of anhydrous AlCl₃ produces
 - (a) *n*-propyl benzene
- (b) isopropyl benzene
- (c) *o*-dipropyl benzene
- (d) a mixture of all of these
- **41.** Which of the following compounds does not undergo a Friedel–Crafts reaction?
 - (a) Benzene

(b) Toluene

(c) Nitrobenzene

- (d) Naphthalene
- 42. Toluene can be converted into benzaldehyde by oxidation with
 - (a) KMnO₄/alkali

- (b) CrO₂Cl₂
- (c) $K_2Cr_2O_7/H_2SO_4$
- (d) O_2/V_2O_5
- **43.** The heat of hydrogenation of cyclohexene is 28.6 kcal and that of cyclohexadiene is about twice as much (i.e., 55.4 kcal). What then would be the heat of hydrogenation of benzene, which has three double bonds?
 - (a) Thrice that of cyclohexene $(28.6 \times 3 \text{ kcal})$
 - (b) The same as that of cyclohexene
 - (c) The same as that of cyclohexadiene
 - (d) 49.8 kcal
- 44. Naphthalene is represented as



45. The directing power of the groups —NH $_2$, —OCH $_3$, —C $_6$ H $_5$ and —NO $_2$ follows the order

(a)
$$-NH_2 > -OCH_3 > -C_6H_5 > -NO_2$$

(b)
$$-NO_2 > -C_6H_5 > -OCH_3 > -NH_2$$

(c)
$$-OCH_3 > -NH_2 > -C_6H_5 > -NO_2$$

(d)
$$-OCH_3 > -NO_2 > -NH_2 > -C_6H_5$$

- **46.** The reactivities of $C_6H_5CH_3$ (I), $C_6H_5CH_2Cl$ (II), $C_6H_5CHCl_2$ (III) and $C_6H_5CCl_3$ (IV) toward nitration are in the order
 - (a) I > II > III > IV
- (b) II > I > III > IV
- (c) IV > III > II > I

- (d) III > I > II > IV
- **47.** Which of the following will be obtained by the bromination of ethylbenzene in the presence of light?

(a)
$$CH_2CH_3$$

(b) CH₂CH₃

(d) CH₂CH₂Br

48. A Friedel–Crafts reaction of benzene with chloroform produces

(d) all of these

- **49.** The stability of the free radicals allyl, benzyl, $\mathring{3}$, $\mathring{2}$, $\mathring{1}$ and $\overset{\bullet}{C}H_3$ is of the order
 - (a) allyl = benzyl $> \mathring{3} > \mathring{2} > \mathring{1} > \mathring{C}H_3$
 - (b) allyl > benzyl > $\mathring{3}$ > $\mathring{2}$ > $\mathring{1}$ > $\mathring{C}H_3$
 - (c) $\mathring{3} > \mathring{2} > \mathring{1} > \overset{\bullet}{C}H_3 > \text{allyl} > \text{benzyl}$
 - (d) $\mathring{3} > \mathring{2} > \mathring{1} > \mathring{C}H_3 > \text{allyl} = \text{benzyl}$

- 50. The reaction of $C_6H_5CH=CHCH_3$ with HBr produces

 - (a) $C_6H_5CHCH_2CH_3$ (b) $C_6H_5-CH_2-CH-CH_3$ R_r
 - (c) C₆H₅CH₂CH₂CH₂Br
- (d) a mixture of these
- **51.** Which is the major product formed in the following reaction?

$$C_6H_6 + (CH_3)_2CHCH_2OH \xrightarrow{H_2SO_4} ?$$

- (a) $C_6H_5CH_2CH(CH_3)_2$ (b) $C_6H_5-CH_3$ CH_3 CH_3 CH_3

$$\begin{array}{ccc} & CH_3 \\ \downarrow & \downarrow \\ (c) & C_6H_5 - C_-C_6H_5 \\ \downarrow & CH_3 \end{array}$$

- (d) A mixture of all these
- **52.** Arrange the following compounds in order of increasing dipole moment. Toluene (I), *m*-dichlorobenzene (II), *o*-dichlorobenzene (III), *p*-di-chlorobenzene (IV)
 - (a) IV < I < II < III

- (b) I < IV < II < III
- (c) IV < II < I < III

- (d) IV < I < III < II
- 53. Which of the following groups would enhance the reactivity of electrophilic aromatic substitution?

 - (a) -CN (b) -CHO (c) $-CH_3$ (d) $-NO_2$
- 54. Which of the following is the end product of the treatment of benzene with an excess of Cl_2 in the presence of I_2 ?
 - (a) Monochlorobenzene
- (b) Dichlorobenzene
- (c) Trichlorobenzene
- (d) Hexachlorocyclohexane
- 55. Benzene reacts with Cl₂ in the presence of sunlight to produce
 - (a) C_6Cl_6
- (b) C_6H_5C1
- (c) $C_6H_6Cl_6$ (d) all of these
- 56. Benzene reacts with chlorine in the presence of iron to produce
 - (a) benzene hexachloride (b) benzyl chloride

- (c) chlorobenzene
- (d) benzoyl chloride
- 57. Which of the following are produced when toluene is heated with chlorine in the presence of light and in the absence of a halogen carrier?
 - (a) $C_6H_5CH_2Cl$

(b) C₆H₅CHCl₂

(c) $C_6H_5CCl_3$

(d) Gammexene

58.	Tolue	ne reacts with chlorine in the p	rese	nce of light to produce
	(a)	o-chlorotoluene	(b)	<i>m</i> -chlorotoluene
	(c)	<i>p</i> -chlorotoluene	(d)	benzyl chloride
59.		nitration of benzene with a r ntrated H ₂ SO ₄ at 100°C produc		are of concentrated HNO ₃ and
	(a)	nitrobenzene	(b)	o-dinitrobenzene
	(c)	<i>m</i> -dinitrobenzene	(d)	<i>p</i> -dinitrobenzene
60.	The fi	nal product of the nitration of	tolue	ene is
	(a)	o-nitrotoluene	(b)	<i>m</i> -nitrotoluene
	(c)	2,4-dinitrotoluene	(d)	2,4,6-trinitrotoluene
61.	The re	eacting species in sulphonatior	is	
	(a)	SO_2	(b)	SO_3
	(c)	SO_4^{2-}	(d)	H_2SO_4
62.		e Friedel–Crafts reaction of lophilic reagent is		ene with an acid chloride, the
		RCOCl ⁺	(b)	RCO
	(c)	AlCl ₃		none of these
63.		n of the following are produce xygen is passed over the catalys		nen a mixture of benzene vapour 0 ₅ at 775 K?
	(a)	Oxalic acid	(b)	Glyoxal
	(c)	Fumaric acid	(d)	Maleic anhydride
64.	Which	n of the following is the strong	est o-	p-directing group?
	(a)	OH (b) Cl	(c)	Br (d) C_6H_5
65.	Which	n of the following can be easily	nitra	ated?
		Benzene		Phenol
	(c)	Nitrobenzene	(d)	Chlorobenzene
66.	Which	n of the following is the least re	eactiv	ve in the case of bromination?
		Phenol		Aniline
	(c)	Nitrobenzene	(d)	Anisole
67.		n of the following reacts slo tution?	wer	than benzene in electrophilic
	(a)	$C_6H_5CH_3$	(b)	$C_6H_5NO_2$
	(c)	C ₆ H ₅ OH	(d)	$C_6H_5NH_2$
68.	The n	itration of toluene is a/an		
	(a)	homolytic substitution	(b)	nucleophilic substitution
	(c)	electrophilic substitution	(d)	electrophilic addition

(a) Aniline

(a) One

(c) Aniline hydrochloride

(d) cyclopropane have a C—C—C angle of 108°?

(b) Two

(a) —NH₂>—OCH₃>—C₆H₅>—CH₃ (b) —NH₂>—CH₃>—C₆H₅>—OCH₃

69. Which of the following is the most reactive in electrophilic substitution?

70. Which among (a) cyclohexane, (b) cyclopentane, (c) cyclobutane and

71. How many geometrical isomers can in 1,2-dimethyl cyclohexane have?

72. The directing power of the following groups in arenes is in the order

(b) Nitrobenzene

(d) None of these

(d) Acetanilide

(c) Three

	(c) $-CH_3 > -NH_2 > -OCH_3 > -C_6H_5$					
	(d)	$-OCH_3 > -C_6H_5 > -CH_3 > -$	—N	H_2		
	benzo attach (a)		the (b)	(II), benzal chloride (III) and inductive effect of the group $IV < III < II < I$ $II < I < IV$		
74	()		` ′			
		ting capacity.	ın (order of decreasing electron-		
	(a)	$COOH > NO_2 > OH > Cl$	(b)	$NO_2 > COOH > Cl > OH$		
	(c)	$OH > Cl > COOH > NO_2$	(d)	$NO_2 > COOH > OH > Cl$		
	Arran capac	0 0 1	der	of decreasing electron-donating		
	(a)	$(CH_3)_3C \longrightarrow (CH_3)_2CH \longrightarrow CC$	H ₃ Cl	$H_2 \longrightarrow CH_3 \longrightarrow$		
	(b)	$CH_3 \longrightarrow CH_3CH_2 \longrightarrow (CH_3)_2CH_3$	CH—	-> (CH ₃) ₃ C—		
	(c)	$(CH_3)_3C = (CH_3)_2CH > C$	H ₃ Cl	$H_2 \longrightarrow CH_3 \longrightarrow$		
	(d)	$(CH_3)_2CH \longrightarrow (CH_3)_3C \longrightarrow CC$	H ₃ Cl	$H_2 \longrightarrow CH_3 \longrightarrow$		
		pasicities of o -toluidine (I), m e (IV) follow the order	-tolu	idine (II), p-toluidine (III) and		
		I > II > III > IV	(b)	IV > III > II > I		
	(c)	III > II > IV > I	(d)	I > IV > II > III		
		asicities of o -nitroaniline (I), n niline (IV) follow the order	n-nit	roaniline (II), <i>p</i> -nitroaniline (III)		
	(a)	I > II > III > IV	(b)	IV > III > II > I		
	(c)	I > III > II > IV	(d)	IV > II > III > I		

78. Arrange aniline (I), *N*-methylaniline (II) and *N*,*N*-dimethylaniline (III) in

79. Arrange methylamine (I), benzylamine (II) and ammonia (III) in order of

(b) III > II > I

(d) I > III > II

(b) III > I > II

(d) III > II > I

order of decreasing basicity.

(a) I > II > III(c) II > I > III

(a) II > I > III

(c) I > III > II

basicity.

80.	Benzenediazonium chloride, on coupling with aniline at 273–278 K in the presence of an acid (pH 4–5), gives				
	_			diazoaminobenzene	
	(c)	<i>p</i> -aminoazobenzene	(d)	none of these	
81.	mol ⁻¹		of b	bond in cyclohexene is 28.6 kcal penzene to cyclohexane is 50 kcal zation energy of benzene?	
	(a)	28.6 kcal mol ⁻¹	(b)	55.0 kcal mol ⁻¹	
	(c)	85.8 kcal mol ⁻¹	(d)	35.8 kcal mol ⁻¹	
82.	The regives	eaction of benzene with chlori	ine ii	n the presence of ferric chloride	
	(a)	chlorobenzene	(b)	2,4-dichlorobenzene	
	(c)	2,4,6-trichlorobenzene	(d)	hexachlorobenzene	
83.		riedel–Crafts reaction of benz rmation of	ene	with propene proceeds through	
	(a)	CH ₃ CH ₂ CH ₂	(b)	CH ₃ CHCH ₃	
	(c)	· CH ₃ CH ₂ CH ₂	(d)	· CH ₃ CHCH ₃	
84.	Arran		der	of decreasing <i>o</i> - and <i>p</i> -directing	
		—NH ₂ ,—OH,	—Cl	,—R	
	(a)	$-Cl>-R>-OH>-NH_2$			
	(b)	$-NH_2 > -OH > -R > -Cl$			
	(c)	$OH >NH_2 >R >Cl$			
	(d)	$-R > -Cl > -NH_2 > -OH$			
85.	Arran	gth.		der of decreasing <i>m</i> -directing	
		—NR ₃ , —CN, —NO	_		
	(a)	$-NR_3 > -NO_2 > -CN > -C$	00	Н	
	(b)	$-COOH > -CN > -NO_2 > -COOH $	⊸ _{NI}	\mathcal{R}_3	

(c)
$$-CN > -NO_2 > -COOH > -NR_3$$

(d)
$$-NO_2 > -CN > -NR_3 > -COOH$$

- 86. The nitration of isopropyl benzene exclusively gives a/an
 - (a) o-isomer

(b) *m*-isomer

(c) p-isomer

- (d) 2,4-disubstituted product
- 87. The reactivity of p-nitrochlorobenzene (II), 2,4-dinitrochlorobenzene (II) and 2,4,6-trinitrochlorobenzene (III) towards alkalis is of the order
 - (a) I > II > III

(b) III > II > I

(c) II > I > III

- (d) III > I > II
- 88. On being heated with calcium hydroxide, benzotrichloride gives
 - (a) CH OH OH
- (b) C OH OH

(c) CHO

- (d) COOH
- 89. *m*-Dinitrobenzene can be converted into *m*-nitroaniline by reduction with
 - (a) Raney nickel

(b) LiAlH₄

(c) $(NH_4)_2S$

(d) Na/C_2H_5OH

• *Type 2* •

Choose the correct options. More than one option is correct.

- **90.** Which of the following are true?
 - Benzene tends to undergo substitution rather than addition reactions.
 - (b) All hydrogen atoms of benzene are equivalent.
 - (c) The carbon-carbon bonds of benzene are alternatively short and long around the ring.
 - (d) There can be two o-disubstituted derivatives.
- 91. The major products of the following bromination reaction

$$CH_2CH_3 \xrightarrow{Br_2, hv}$$

are

(a)
$$Br$$
 CH_2CH_3 (b) Br CH_2CH_3 (c) CH_2CH_2Br (d) CH_2CH_3

- 92. Isopropylbenzene can be prepared by
 - (a) Benzene + $CH_3CH=CH_2 \xrightarrow{H_2SO_4}$
 - (b) Benzene + CH₃—CH—CH₃ $\xrightarrow{\text{H}_2\text{SO}_4}$ Cl
 - (c) Benzene + CH_3CH — CH_3 $AlCl_3$ Cl
 - (d) Benzene + $CH_3CH_2CH_2CI \xrightarrow{AlCl_3}$
- 93. Tert. butyl benzene can be prepared by
 - (a) Benzene + CH_2 = $C(CH_3)_2 \xrightarrow{H_2SO_4}$
 - (b) Benzene + $(CH_3)_3COH \xrightarrow{H_2SO_4}$ (c) Benzene + $(CH_3)_3C$ — $CI \xrightarrow{AlCl_3}$

 - (d) none of these
- 94. Triphenylmethane can be prepared by
 - (a) Benzene + benzal chloride $\xrightarrow{\text{AlCl}_3}$
 - (b) Benzene + chloroform $\xrightarrow{\text{AlCl}_3}$
 - (c) Benzene + benzaldehyde $\xrightarrow{ZnCl_2}$
 - (d) none of these
- 95. Which of the following characteristics does an aromatic compound exhibit?
 - (a) It should have $(4n + 2) \pi$ -electrons.
 - (b) It should be planar and conjugated.
 - It should have $4n \pi$ -electrons.
 - (d) It should possess high resonance energy.

- **96.** Which of the following do not undergo a Friedel–Crafts reaction with benzene?
 - (a) Aniline

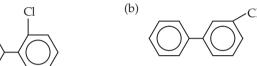
(b) Chlorobenzene

(c)
$$CH_3$$
 $C=C$ CI

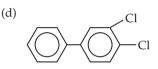
(d) Acetyl chloride

97. The reaction of biphenyl with HOCl in the presence of a strong acid gives

(a) (



(c) CI



- 98. Which of the following are obtained by the fractionation of coal tar?
 - (a) Light oil

(b) Middle oil

(c) Heavy oil

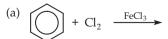
- (d) Vegetable oil
- 99. Which of the following statements are true for benzene?
 - (a) It is a flat, regular, hexagonal molecule.
 - (b) Each C—C—C angle is 109°28′.
 - (c) The C—C bond length is 1.39 Å.
 - (d) All carbon atoms are sp³-hybridized.
- 100. Which of the following meet the requirements of the Huckel rule?
 - (a) Naphthalene

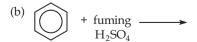
- (b) Anthracene
- (c) 1,3,5,7-Cyclooctatetraene
- (d) 1,3-Cyclobutadiene
- 101. Which of the following compounds are aromatic?
 - (a) Pyridine

(b) Pyridiazine

(c) Thiophene

- (d) Pyrrole
- 102. Which of the following are electrophilic substitution reactions?





(c) + Cl₂ - UV light

(d) All of these

- **103.** Toluene reacts with ethyl bromide in the presence of anhydrous AlCl₃ to produce
 - (a) o-ethyltoluene

(b) p-ethyltoluene

- (c) m-ethyltoluene
- (d) a mixture of all of these
- 104. Isopropylbenzene can be obtained by

(a)
$$+ CH_3 - CH = CH_2 \xrightarrow{AlCl_3}$$

(c)
$$CH_3$$
 CHCl $AlCl_3$ \rightarrow

- **105.** Benzene can be obtained by
 - (a) $C_6H_5COOH + NaOH \xrightarrow{CaO} \Delta$
 - (b) $C_6H_5OH + Zn \xrightarrow{\Delta}$
 - (c) $C_6H_5N=NCl+H_2O \longrightarrow$
 - (d) all of these
- **106.** The sulphonation of toluene with oleum at 35°C produces
 - (a) p-toluene sulphonic acid
 - (b) o-toluene sulphonic acid
 - (c) *m*-toluene sulphonic acid
 - (d) a mixture of (a), (b) and (c) in approximately equal proportions
- 107. Which of the following groups are meta-directing?
 - (a) $-NH_2$

(b) —OH

(c) —NO₂

- (d) —COOH
- 108. Which of the following groups are ortho- and para-directing?
 - (a) —OH

(b) —CHO

(c) —CN

(d) —NHCOCH₃

109. *m*-Nitrobenzoic acid can be obtained by

(a) toluene
$$\xrightarrow{[O]}$$
 A $\xrightarrow{HNO_3}$ A $\xrightarrow{H,SO_4}$

(b) toluene
$$\xrightarrow{[O]}$$
 A $\xrightarrow{\text{HNO}_3}$ B $\xrightarrow{\text{[O]}}$ KMnO₄

(c) toluene
$$\xrightarrow{\text{nitration}}$$
 A $\xrightarrow{[O]}$ KMnO₄

- (d) all these methods
- 110. Which of the following groups are electron-releasing?
 - (a) $-NH_2$

(b) —CH₃

(c) —NO₂

- (d) —CN
- 111. Which of the following groups are electron-withdrawing?
 - (a) —COOH

(b) $-C_6H_5$

(c) —NH₂

- (d) —OH
- **112.** The Friedel–Crafts reaction of benzene with *n*-butyl chloride at 0°C produces
 - (a) C_6H_5 — $CH_2CH_2CH_2CH_3$

$$\begin{array}{c} CH_3 \\ \mid \\ \text{(b)} \quad C_6H_5CH\text{---}CH_2CH_3 \end{array}$$

- (d) all of these
- **113.** Which of the following statements are correct?
 - (a) An activating group is an electron-releasing group.
 - (b) An activating group activates all positions of the benzene ring.
 - (c) The effect of any group—whether activating or deactivating—is the strongest at the ortho- and para-positions in the benzene ring.
 - (d) An activating group activates only the ortho- and para-positions in the benzene ring.
- 114. The necessary conditions for halogenation are
 - (a) low temperature
 - (b) darkness
 - (c) the presence of a halogen carrier
 - (d) none of these

- 115. The major products formed in the reaction of toluene with chlorine in the presence of ferric chloride are
 - (a) o-chlorotoluene
- (b) *m*-chlorotoluene
- (c) p-chlorotoluene
- (d) benzyl chloride

- 116. Benzene can undergo
 - (a) substitution

(b) addition

(c) elimination

- (d) oxidation
- 117. Which of the following statements about the nitration of aromatic compounds are correct?
 - (a) The rate of nitration of toluene is greater than that of benzene.
 - (b) The rate of nitration of benzene is almost the same as that of hexadeutrobenzene.
 - (c) The rate of nitration of benzene is greater than that of hexadeutrobenzene.
 - (d) Nitration is an electrophilic substitution reaction.
- 118. Which of the following reactions can be conveniently carried out?
 - (a) Benzene + $I_2 \longrightarrow$ iodobenzene
 - (b) Toluene + I_2 $\xrightarrow{HNO_3}$ o- and p-iodotoluene (c) Toluene + Cl_2 \xrightarrow{light} benzyl chloride

 - (d) Benzene + $Cl_2 \xrightarrow{hv}$ benzene hexachloride
- 119. Which of the following intermediate stages are possible in the conversion of benzaldehyde into benzoin by reaction with HCN?

(a)
$$C_6H_5$$
— C — H
OH

(b)
$$C_6H_5$$
— C_6 :

(c)
$$C_6H_5$$
— C — C_6H_5 (d) None of these OH OH

- 120. Which of the groups, if present in the benzene nucleus, are o- and *p*-directing?
 - (a) —CH₃

(b) —C≡N

(c) —NH₂

(d) —CHO

- **121.** Which of the following are electrophiles?
 - (a) R—CH₂

(b) Cl⁻

(c) C=O

- (d) NO₂
- 122. Aniline, on acetylation followed by bromination ($\rm Br_2/CH_3COOH$) and subsequent hydrolysis, gives
 - (a) o-bromoaniline
- (b) p-bromoaniline
- (c) *m*-bromoaniline
- (d) m-bromoacetanilide
- 123. Which of the following can be represented by a resonance hybrid?
 - (a) Sulphur trioxide
- (b) Benzene

(c) Toluene

- (d) Benzene sulphonic acid
- 124. Iodobenzene can be obtained by
 - (a) $C_6H_6 + I_2 \longrightarrow$

- (b) $C_6H_6 + I_2 \xrightarrow{HNO_3}$
- (c) $C_6H_5N_2^+Cl^- + KI \longrightarrow$
- (d) $C_6H_6 + HI \longrightarrow$

Answers

1. c	2. d	3. c	4. c	5. c
6. b	7. a	8. b	9. b	10. a
11. c	12. a	13. a	14. a	15. b
16. c	17. b	18. b	19. d	20. a
21. a	22. a	23. c	24. a	25. a
26. c	27. a	28. c	29. b	30. a
31. a	32. b	33. a	34. b	35. b
36. b	37. c	38. a	39. a	40. b
41. c	42. b	43. d	44. d	45. a
46. a	47. c	48. c	49. a	50. a
51. b	52. a	53. c	54. d	55. c
56. c	57. c	58. d	59. c	60. d
61. b	62. b	63. d	64. a	65. b
66. c	67. b	68. c	69. b	70. b
71. b	72. a	73. a	74. b	75. a
76. c	77. d	78. b	79. c	80. c
81. d	82. a	83. b	84. b	85. a
86. c	87. b	88. d	89. c	90. a, b
91. a, c, d	92. a, b, c, d	93. a, b, c	94. a, b, c	95. a, b, d
96. a, b, c	97. a, b, c	98. a, b, c	99. a, c	100. a, b
101. a, b, c, d	102. a, b	103. a, b	104. a, b, c	105. a, b

106. a, b	107. c, d	108. a, d	109. a, b	110. a, b
111. a, b	112. a, b	113. a, b, c	114. a, b, c	115. a, c
116. a, b, d	117. a, c, d	118. b, c, d	119. a, b, c	120. a, c
121. c, d	122. a, b	123. a, b, c, d	124. b, c	

Hints to More Difficult Problems

- **2.** An aromatic compound has $(4n + 2) \pi$ -electrons (Huckel rule).
- **6.** The COOH group is *m*-directing.
- 8. C₆H₅COCH₂C₆H₅ will behave like alkyl benzene. So the reaction (bromination) will take place preferably at the *p*-position. The attack at the *o*-position is muted by steric hindrance.
- 10. A conjugated diene is the most stable.
- 11. According to the Huckel rule, a conjugated polyene will be aromatic only if it contains (4n+2) π -electrons. If it has 4n π electrons, it will be nonaromatic.
- 12. The addition of an alkene molecule with 2 π -electrons to a conjugated diene (4 π -electrons) is called a [4 + 2] cycloaddition reaction.
- **15.** A 2 carbocation is more stable than a 1 carbocation. Therefore, wherever possible, a 1 carbocation gets isomerized or converted into a 2 carbocation.
- **24.** If all the hydrogens in an aromatic compound are equivalent, only one monosubstitution product is obtained from it.
- **31.** The Friedel–Crafts reaction of benzene with an alkyl halide gives alkyl-substituted benzene.
- **33.** See the hint to Q. 31.
- **36.** If all the available hydrogens are equivalent in a disubstituted aromatic compound, only one monosubstituted product is obtained from it.
- **100.** See the hint to Q. 2.
- **106.** The methyl group is *o* and *p*-directing.
- 116. Benzene cannot undergo an elimination reaction.
- **118.** Iodobenzene cannot be obtained by the reaction of benzene with iodine. The reaction will take place only in the presence of an oxidizing agent.

Compounds Containing Oxygen: Alcohols, Phenols, Ethers, Aldehydes, Ketones, Carboxylic Acids and Their Derivatives

• Type 1 •

Choose the correct option. Ony one option is correct.

1. The final product (IV) in the sequence of reactions

is

(a)
$$CH_3$$
— $CHOCH_2CH_2OH$ (b) CH_3 — $CHCH_2CH_2Br$ CH_3

(c)
$$CH_3-CH-CH_2CH_2OH$$
 (d) $CH_3-CHOCH_2CH_3$ CH_3

2. By which of the following procedures can ethyl *n*-propyl ether be obtained?

(c)
$$C_2H_5OH + H_2SO_4 \xrightarrow{140^{\circ}C} 3-142$$

(d)
$$C_2H_5OH + conc. H_2SO_4 \xrightarrow{180^{\circ}C} I \xrightarrow{CH_3CH_2CH_2Br}$$

3. The reaction of 1 mol each of *p*-hydroxyacetophenone and methyl magnesium iodide will give

(a)
$$CH_4 + IMgO \longrightarrow COCH_3$$

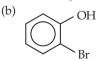
(c)
$$OMgI$$
 $H_3C - C$
 OH_2
 OH

(d)
$$MgI$$
 CH_3O $COCH_3$

- **4.** Which of the following can be used for the synthesis of 3-methyl-1-butanol?
 - (a) $n-C_3H_7MgBr + CH_3CHO$
 - (b) $(CH_3)_2CHCH_2MgBr + CH_2O$

(c)
$$\begin{array}{c} O \\ C_2H_5MgBr + CH_3-C-CH_3 \end{array}$$

- (d) $(CH_3)_3CMgBr + CH_2O$
- 5. The product obtained by the reaction of HBr with phenol is



(d) There is no reaction.

The conversion

$$\begin{array}{c} O \\ || \\ CH_3-C-CH_2CH_2CO_2CH_3 \end{array} \longrightarrow \begin{array}{c} OH \\ || \\ CH_3-CH-CH_2CH_2CH_2OH \end{array}$$
 can be effected using

- (a) LiAlH₄ and then H⁺
- (b) NaBH $_4$ and then H $^+$
- (c) H₂/Pt-C

(d) none of these

The conversion

$$\begin{array}{c} O \\ \parallel \\ CH_3 - C - CH_2CH_2CO_2CH_3 \longrightarrow \\ CH_3 - CH - CH_2CH_2CO_2CH_3 \end{array}$$
 can be effected using

- (a) LiAlH₄ and then H ⁺
- (b) NaBH $_4$ and then H $^+$
- (c) H₂/carbon

- (d) none of these
- 8. Among the following, which will react most rapidly with HBr?
 - (a) C₆H₅CH₂CH₂CH₂OH
- (b) $C_6H_5CHCH_2CH_3$ ÓН

- 9. By which of the following methods is phenol prepared on a commercial scale?
 - (a) Chlorobenzene + NaOH (350°C) and then H +
 - (b) Benzene + O₂
 - (c) Toluene + O_2 and then H^+ , Δ
 - (d) None of these
- **10.** The product X in the reaction

$$CH_3 - CH_3 + CN^{-} \xrightarrow{H^+} C_4H_7NO \xrightarrow{H_2O} X$$

is

(c)
$$CH_2 = C - COOH$$

- 11. Which of the following reactions will give benzaldehyde?
 - (a) $C_6H_5CH_2Cl \xrightarrow{OH^-}$ (b) $C_6H_5CH(OCH_3)_2 \xrightarrow{H^+}$
 - (c) $C_6H_5COOH \xrightarrow{1 \cdot LiAlH_4} (d)$ None of these
- 12. The final product (III) obtained in the reaction sequence

$$CH_{3}CH_{2}COOH \xrightarrow{\hspace{1cm}PCl_{3}} I \xrightarrow{\hspace{1cm}C_{6}H_{6}} II \xrightarrow{\hspace{1cm}NH_{2}NH_{2}} III$$

is

13. By which of the following reagents can the following conversion be effected?

$$O \longrightarrow CO_2CH_3 \longrightarrow O \longrightarrow O-CH_3$$

- (a) 2CH₃MgBr and then H₃O⁺
- (b) HOCH₂—CH₂OH, H⁺; LiAlH₄, ether; 2CH₃MgBr; H₃O⁺
- (c) HOCH₂—CH₂OH, H⁺; 2CH₃MgBr; H₃O⁺
- (d) HOCH₂—CH₂OH, H⁺; H₂, Pt; CH₃OH, H⁺
- 14. The major product obtained in the reaction

$$O \xrightarrow{HCN} \boxed{ \frac{H_2SO_4}{H_2O/\Delta}}$$

(a)
$$\bigcirc$$
 C-OH (b) \bigcirc CN

15. The conversion

$$\stackrel{\circ}{\bigsqcup} \rightarrow \stackrel{\circ}{\bigsqcup}$$

can be effected by using the reagent

(a) H_2O , H_2SO_4

- (b) O₂
- (c) O \parallel C_6H_5 —COOH
- (d) CrO_3 , H_2SO_4

16. The final product (III) obtained in the reaction

(a)
$$CH_3$$

 $CH_3 - C - CH = CHCH_2CH_2OH$
 CH_3

(b)
$$CH_3$$

 $CH_3 - C - CH = C - CH_2OH$
 $CH_3 - CH_3$

$$\begin{array}{c} \text{(d)} & \text{CH}_3 \\ \text{CH}_3 - \text{C} - \text{CH}_2 - \text{CH}_2 \text{CH}_2 \text{OH} \\ \text{CH}_3 \end{array}$$

17. The product obtained in the reaction

The product obtained in the reaction
$$CH_3CH=CHCO_2C_2H_5 \xrightarrow{NaCN} C_2H_5OH/CH_3COOH$$
(a)
$$CH_3CH_2CHCO_2C_2H_5 \qquad (b) CH_3CH=CHCH_2CN$$

$$CN$$

- (c) $CH_3CH = CHCN$
- (d) $CH_3CH CH_2CO_2C_2H_5$

18. The major product obtained in the reaction

$$(CH_3)_2C = CH - C - CH_3 \xrightarrow{1. CH_3MgBr}$$

- $(CH_{3})_{2}C = CH C CH_{3} \xrightarrow{1. CH_{3}MgBr} \xrightarrow{2. H_{3}O^{+}}$ (a) $(CH_{3})_{3}CCH_{2}COCH_{3}$ (b) $(CH_{3})_{2}C = CHC (CH_{3})_{2}OH$ (c) $(CH_{3})_{2}CH CHCOCH_{3}$ (d) $OH_{3}C$

19. Which of the following hydrogens will be the most acidic?

$$CH_2$$
— CH_2 — CH_2 — CH_3
 CH_3

20. Which of the following alcohols will dehydrate most rapidly when treated with H₂SO₄?

(c)
$$CH_3$$
 (d) CH_3 — $CH_2CH_2CH_2OH_3$
 CH_3 — C — CH — CH_3
 $OH CH_3$

- 21. The oxidation of 2-hexanol with H₂CrO₄ gives
 - (a) CH₃CO₂H

- (b) CH₃(CH₂)₂CO₂H
- O || CH₃(CH₂)₃—CCH₃
- **22.** Which of the following is the best procedure to make isopropylmethylether using the Williamson method?
 - (a) $CH_3OH + (CH_3)_2CHOH + H_2SO_4$
 - (b) $CH_3OH + (CH_3)_2CHCH_2OH + H_2SO_4$
 - (c) CH₃ONa + (CH₃)₂CHBr
 - (d) $CH_3I + (CH_3)_2CHONa$
- 23. Which is the major product obtained in the following reaction?

$$CH_{3} \xrightarrow{CH_{3} OH} CH_{3} \xrightarrow{H_{2}SO_{4}} \xrightarrow{CH_{3} CH_{3}}$$

- (a) $(CH_3)_2C=C(CH_3)_2$
- (b) $(CH_3)_3C$ —CH= CH_2
- (c) (CH₃)₂C=CHCH₃
- (d) (CH₃)₂C=CHCH₂CH₃
- 24. Among the following, which is the strongest acid?
 - (a) CHF₂—CH₂—CH₂—COOH (b) CH₃—CH₂—CF₂—COOH
- - (c) CH₂F—CHF—CH₂—COOH (d) CH₃—CF₂—CH₂—COOH
- **25.** Which of the following compounds can form a δ -lactone?
 - (a) HOOCCH₂CH₂CH₂COOH
 - (b) CH₃CH₂CH—CH₂COOH OH

 - (c) HOCH₂CH₂CH₂COOH (d) CH₃CH—CH₂CH₂COOH
- **26.** The products obtained in the reaction

$$CH_3CH_2C - CH - COH \xrightarrow{\Delta}$$

$$CH_3CH_2C - CH - COH \xrightarrow{\Delta}$$

are

- (d) none of these
- 27. The product obtained in the reaction

$$C_6H_5CO_2C_2H_5 + CH_3CO_2C_2H_5 \xrightarrow{\hspace*{1cm} 1. \text{ NaOEt} \atop \hspace*{1cm} 2. \text{ } H^+}$$

is

- (a) $C_6H_5COCH_2CO_2C_2H_5$ (b) C_6H_5CH — $CH_2CO_2C_2H_5$ OH
- (c) $C_6H_5CH_2CO_2C_2H_5$
- (d) C₆H₅CO₂CH₂CO₂C₃H₅
- 28. The final product (III) obtained in the reaction sequence

$$\begin{array}{c|c}
O & O \\
\parallel & \parallel & \parallel \\
CH_3CCH_2COC_2H_5 & \hline{1. NaOEt} & 1 & \hline{1. NaOH} & II & \hline{heat} \\
2. CH_3CH_2BT & \hline{1. NaOH} & II & \hline{-CO_2} & III
\end{array}$$

- (a)
- (b) O \parallel $C_6H_5CH_2COC_2H_5$
- (d) none of these
- 29. Which of the following compounds will not give bakelite-type resins?
 - (a) (b)

30. The major product obtained in the reaction

$$CH_3CH_2C \overline{=} CH \xrightarrow{\qquad Hg^{+},\, H_3O^+ }$$

is

- (a) CH₃CH₂CH₂CH(OH)₂ (b) CH₃CH₂CH=CHOH
- (c) CH₃CH₂CH₂CHO
- (d) CH₃CH₂C-CH₃

31. The product obtained in the reaction

$$\begin{array}{c} \text{CH}_{3} \\ \text{CH} - \text{C} - \text{CH} \\ \text{CH}_{3} \end{array} \qquad \begin{array}{c} \text{CH}_{3} \\ \text{CH}_{3} \\ \text{CH}_{3} \end{array} \qquad \begin{array}{c} \text{1.} \text{(CH}_{3})_{2} \text{CHMgBr} \\ \text{2.} \text{H}^{+} \end{array}$$

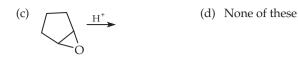
(a) OH
$$CH_3$$
 $CH - C - CH$ CH_3 CH_3 CH_3 CH_3 CH_3 CH_3

(b)
$$H_3C$$
 $CH-CH(OH)-CH < CH_3 + CH_3-CH=CH_2$

$$\begin{array}{ccc} \text{(c)} & \text{CH}_3 \\ & \text{CH}_3 \end{array} \text{C} \begin{array}{c} \text{OH} \\ \text{CH}_3 \end{array} \\ \begin{array}{ccc} \text{CH}_3 \end{array}$$

- (d) There is no reaction.
- 32. By which the following reactions can trans-cyclopentane-1,2-diol be obtained?

(a)
$$\frac{\text{KMnO}_4}{\text{dilute aqueous}}$$
 (b) $\frac{1. \text{OsO}_4}{2. \text{H}^+}$



- 33. The order of reactivity towards nucleophilic addition in CH₂O (I), CH₃CHO (II) and CH₃COCH₃ (III) is
- (a) I > II > III (b) III > II > I (c) II > I > III (d) III > I > II

34. The conversion

can be achieved by using

(a) H₂-NI

(b) LiAlH₄

(c) NaBH₄

- (d) none of these
- 35. The ease of reduction of C₆H₅COCl (I), C₆H₅CHO (II), C₆H₅COCH₃ (III) and

$$\begin{matrix} & O \\ \parallel \\ C_6H_5\text{---}C\text{---}OC_2H_5 & (IV) \end{matrix}$$

by hydrogen over a palladium catalyst follows the order

(a) I > II > III > IV

(b) IV > III > II > I

(c) II > III > I > IV

- (d) III > II > IV
- **36.** The trans-hydroxylation of cyclohexene can be done by
 - (a) dilute KMnO₄
 - (b) using OsO₄
 - (c) converting cyclohexene into an epoxide and then opening the epoxide ring
 - (d) none of these
- 37. Consider the following reaction.

Which of the following statements is correct?

- (a) The product is (R)–(-)-2-butanol.
- (b) The product is (S)–(+)-2-butanol.
- (c) The product is in racemic form.
- (d) None of these.
- 38. The relative acidities of H₂O (I), CH₃OH (II), CH₃C≡CH (III), NH₃ (IV) and CH₄ (V) follow the order
 - (a) II > I > IV > III > V
- (b) I > II > III > IV > V
- (c) III > V > IV > I > II
- (d) II > I > III > IV > V

39. In the reaction

the rate-determining step is

(a)
$$CH_3$$
 CH_3 CH_3 H CH_3 CH_3

(b)
$$CH_3 H$$
 $CH_3 - C - O - H$
 $CH_3 - CH_3 - CH_3 + H_2O$
 $CH_3 + H_2O$

(c)
$$CH_3 - C \stackrel{CH_3}{\leftarrow} + CI \stackrel{-}{\longleftarrow} CH_3 - \stackrel{CH_3}{\leftarrow} CH_3$$

(d) all steps take place with equal ease.

40. The reduction

$$\begin{array}{c} O \\ \parallel \\ - C - OCH_3 \longrightarrow HOCH_2 \longrightarrow \begin{array}{c} O \\ \parallel \\ - COCH_3 \end{array}$$

can be achieved by using

(a) NaBH₄

- (b) LiAlH₄
- (c) CuO · CuCN₂O₄
- (d) none of these

41. The product obtained in the reaction

$$\begin{array}{c} \text{O} \\ \parallel \\ \text{RCH} - \text{CH}_2\text{C} - \text{OH} \end{array} \xrightarrow{\text{heat}}$$

(b)
$$O$$
 CH_2-C-O $HC-R$ $O-C-CH_2$

- (c) RCH=CHCOOH
- (d) none of these
- **42.** Arrange phenol (I), cyclohexanol (II), 2,4,6-trinitrophenol (III) and acetic acid (IV) in order of acidity.
 - (a) III > IV > I > II

(b) I > II > III > IV

(c) III > I > II > IV

- (d) II > I > IV > III
- 43. The final product obtained in the reaction

$$H_3C$$
 OCH₃ + HBr $\xrightarrow{\Delta}$

is

(b)
$$CH_3 \longrightarrow Br$$

- (d) none of these
- 44. The final product obtained in the reaction

$$CH2O + NH3 \longrightarrow \begin{bmatrix} & & HNO3 \\ & & Ac2O \end{bmatrix}$$

(a) OH (b)
$$H_2C$$
 NH_2 H_2C CH_2 N CH_2

is

(a)

(c)
$$NO_2$$
 (d) none of these NO_2 NO_2

45. The major product obtained in the reaction

$$CH_{2}=CH-C-CH_{3} + HCN \xrightarrow{OH}$$
(a) OH (b) O
$$CH_{2}=CH-C-CH_{3} \qquad NC-CH_{2}-CH_{2}-C-CH_{3}$$
(c) OH (d) none of these
$$CH_{2}=CH-C-CH_{3} \qquad COOH$$

46. The product obtained in the reaction

47. The reaction

is known as

- (a) aldol condensation
- (b) the Cannizzaro reaction
- (c) the internal Cannizzaro reaction
- (d) benzilic acid rearrangement
- 48. The product obtained in the reaction

$$CH_3CH_2CO_2H \xrightarrow{Cl_2/P}$$

is

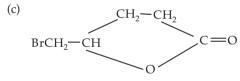
(a) CH₃CHCO₂H | | Cl

- (b) ClCH₂CH₂CO₂H
- (c) Cl $CH_3 C CO_2H$ Cl
- (d) Cl₂CHCH₂CO₂F
- 49. The final product obtained in the reaction

$$CH_2 = CHCH_2CH_2CO_2H \xrightarrow{HOBr}$$

is

(b) HOCH₂—CH₂CH₂CH₂CO₂H



- **50.** Which of the following is a primary alcohol?
 - (a) Butan-1-ol

(b) Butan-2-ol

(c) Propan-2-ol

- (d) 2-Dimethylhexane-4-ol
- **51.** How many isomers of $C_5H_{11}OH$ will be primary alcohols?
 - (a) Four

(b) Five

(c) Three

- (d) Two
- **52.** Ethyl iodide reacts with moist silver oxide to produce
 - (a) ethane

(b) propane

(c) ethyl alcohol

(d) diethyl ether

55.		eduction of R—COOH to R—	C_{n_2}	OH can be effected by
	(a)	sodium/alcohol	(b)	Zn/HCl
	(c)	LiAlH ₄	(d)	aluminium isopropoxide
54.	The al	lkaline hydrolysis of esters is k	now	n as
	(a)	hydration	(b)	esterification
	(c)	dehydration	(d)	saponification
55.	The co	onversion $CH_3CH_2CHO \longrightarrow 0$	CH ₃ C	CH ₂ CH ₂ OH can be effected by
	(a)	$NaBH_4$	(b)	Zn/HCl
	(c)	H ₂ /Ni	(d)	Na + alcohol
56.	The co	onversion CH ₂ =CH—CHO —	\rightarrow (CH ₂ =CHCH ₂ OH can be effected
	(a)	Na + alcohol	(b)	Zn + HCl
	(c)	H ₂ /Ni	(d)	LiAlH ₄
57.	The b	oiling points of isomeric alcoho	ols fo	ollow the order
	(a)	primary > secondary > tertiar	y	
	(b)	tertiary > secondary > primar	y	
	(c)	secondary > tertiary > primar	y	
	(d)	They do not follow any order		
58.	The a	cidic character of 1, 2, 3 alcohol	ls, H	₂ O and RC≡CH is of the order
	(a)	$H_2O > \mathring{1} > \mathring{2} > \mathring{3} > RC \equiv CH$		
	(b)	$RC \equiv CH > \mathring{3} > \mathring{2} > \mathring{1} > H_2O$		
	(c)	$\mathring{1} > \mathring{2} > \mathring{3} > H_2O > RC \equiv CH$		
	(d)	$\mathring{3} > \mathring{2} > \mathring{1} > H_2O > RC \equiv CH$		
59.	The o		hloro	o-compounds by reaction with
	(a)	$\mathring{1} > \mathring{2} > \mathring{3}$	(b)	$\mathring{3} > \mathring{2} > \mathring{1}$
	(c)	$\overset{\circ}{2} > \overset{\circ}{1} > \overset{\circ}{3}$	(d)	$\mathring{1} = \mathring{2} > \mathring{3}$
60.	C_2H_5	OH reacts with halogen acids K. Its reactivity with HI, HBr ar HI > HBr > HCl	nd H	the presence of ZnCl ₂ to form Cl follows the order HCl > HBr > HI
	(c)	HCl > HI > HBr	(d)	HBr=HCl > HI
61.	The d	ehydration of $\mathring{1}$, $\mathring{2}$ and $\mathring{3}$ alcohological		
	(a)	$\mathring{1} > \mathring{2} > \mathring{3}$	(b)	$\mathring{3} > \mathring{2} > \mathring{1}$
	(c)	$\overset{\circ}{2} > \overset{\circ}{1} > \overset{\circ}{3}$	(d)	There is no such order.

62. In the following reaction sequence

the alcohol is a

- (a) primary alcohol
- (b) secondary alcohol
- (c) tertiary alcohol
- (d) phenol
- **63.** The acidity of the compounds RCOOH, H₂CO₃, C₆H₅OH, ROH decreases in the order
 - (a) $RCOOH > H_2CO_3 > C_6H_5OH > ROH$
 - (b) $C_6H_5OH > RCOOH > H_2CO_3 > ROH$
 - (c) $ROH > C_6H_5OH > RCOOH > H_2CO_3$
 - (d) $H_2CO_3 > RCOOH > C_6H_5OH > ROH$
- **64.** The acidity of *p*-nitrophenol (I), *o*-nitrophenol (II), *m*-nitrophenol (III) and phenol (IV) decreases in the order
 - (a) I > II > III > IV

(b) IV > I > II > III

(c) II > III > I > IV

- (d) III > II > IV
- **65.** The acidities of phenol (I), *p*-cresol (II), *m*-cresol (III) and *o*-cresol (IV) follow the order
 - (a) I > II > III > IV
- (b) II > I > III > IV

(c) III > I > II > IV

- (d) IV > III > II > I
- **66.** Which of the following represents the Dow process for the manufacture of phenol?

(c)
$$SO_3Na$$
 + $2NaOH$ $1.625 K$ $2.H^+$

- (d) None of these
- **67.** Arrange the following in order of decreasing acidic strength. *p*-nitrophenol (I), *p*-cresol (II), *m*-cresol (III), phenol (IV)
 - (a) I > II > III > IV

(b) IV > III > II > I

- (c) I > III > II > IV
- (d) III > II > IV
- **68.** Arrange the following in order of decreasing acidic strength. 2,4,6-trinitrophenol (I), 2,4-dinitrophenol (II), *p*-nitrophenol (III), phenol (IV)

(a) I > II > III > IV

(b) I > III > II > IV

(c) IV > III > II > I

- (d) III > II > IV
- 69. Which of the following is constituent of oil of wintergreen?
 - (a) Aspirin

- (b) Salol
- (c) Methyl salicylate
- (d) None of these
- 70. In the Liebermann test for phenols, the blue or green colour produced is due to the formation of

(c)
$$O = N - OH$$

(d)
$$O = N - O Na^4$$

- 71. When ethylene glycol is heated with a mixture of concentrated HNO_3 and concentrated H_2SO_4 , it produces
 - (a) COOH

(b) $CO_2 + H_2$

(c) CH_2ONO_2 | CH_2ONO_2

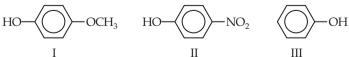
- (d) CH₂ONO₂ | CH₂OH
- 72. When glycerol is heated with an excess of HI, it produces
 - (a) allyl iodide

- (b) propene
- (c) glycerol tri-iodide
- (d) 2-iodopropane
- 73. When glycerol is heated with oxalic acid at 503 K, it produces
 - (a) formic acid

(b) allyl alcohol

(c) acrolein

- (d) glyceric acid
- 74. Arrange the following in order of decreasing acidity.



(a) I > II > III

(b) II > III > I

(c) III > II > I

(d) III > I > II

75.	Ethyl alcohol is less acidic than phenol because					
	(a)	the phenoxide ion is more resonance stabilized than phenol				
	(b)	there is more hydrogen bonding in phenol than in ethyl alcohol				
	(c)	the ethoxide ion is less resonance stabilized than ethyl alcohol				
	(d)	phenol has a higher boiling p	oint	than ethyl alcohol		
76.	Which ZnCl ₂	ich of the following is the most reactive with HCl in the presence of Cl ₂ ?				
	(a)	(CH ₃) ₃ COH	(b)	(CH ₃) ₂ CHCH ₂ OH		
	(c)	(CH ₃) ₂ CHOH	(d)	C_6H_5OH		
77.	Glyce	rol reacts with KHSO ₄ to proc	luce			
		acrolein		oxalic acid		
	(c)	formaldehyde	(d)	tartaric acid		
78.		e reaction of phenol with CHC es attacking the ring is	Cl₃ ar	nd aqueous NaOH at 345 K, the		
	(a)	CHCl ₃ (b) CHCl ₂	(c)	$COCl_2$ (d) $:CCl_2$		
79.	water	gas and excess hydrogen over		(under pressure) a mixture of ted		
	(a)	platinized asbestos	(b)	cobalt chloride		
	(c)	Zn and Cr oxides	(d)	finely divided metal		
80.	Pheno	ol $\xrightarrow{1. \text{ NaOH}}$ A $\xrightarrow{\text{H}^+/\text{H}}$	I ₂ O	$\rightarrow B \xrightarrow{Ac_2O} C$		
	In this	s reaction, the end product C is	5			
		salicylaldehyde	(b)	salicylic acid		
	(c)	phenyl acetate	(d)	aspirin		
81.	Alcoh	ols are soluble in water due to	the i	formation of		
	(a)	covalent bonds	(b)	ionic bonds		
	(c)	hydrogen bonds with water	(d)	none of these		
82.	Which of the following exhibits maximum hydrogen bonding?					
	(a)	Ethyl alcohol	(b)	Diethyl ether		
	(c)	Ethyl chloride	(d)	Triethylamine		
83.	Amor	ng the following alcohols, which	h is	the least soluble in water?		
	(a)	Ethyl alcohol	(b)	C ₄ H ₉ OH		
	(c)	$C_5H_{11}OH$	(d)	$C_{12}H_{25}OH$		
84.	-		opar	nol and tert. butanol decrease in		
	the order					

(a) ethanol > isopropanol > tert. butanol(b) tert. butanol > isopropanol > ethanol

- (c) isopropanol > tert. butanol > ethanol
- (d) tert. butanol > ethanol > isopropanol
- 85. The intermediate formed during the dehydration of alcohol is
 - (a) carbanion

(b) carbonium ion

(c) free radical

- (d) carbene
- 86. Which of the following produces the most stable carbonium ion upon dehydration?
 - (a) (CH₃)₂CHCH₂OH
- (b) (CH₃)₃COH
- (c) $CH_3(CH_2)_2CH_2OH$ (d) CH_3 —CH— CH_2 — CH_3 OH
- 87. An alcohol, on oxidation, produces a ketone with the same number of carbon atoms. When the ketone is oxidized, it yields an acid with a fewer number of carbon atoms. The alcohol could be a
 - (a) primary alcohol
- (b) secondary alcohol
- (c) tertiary alcohol
- (d) none of these
- 88. A compound with the molecular formula C₃H₈O, on vigorous oxidation, produces the acid $C_3H_6O_2$. The compound is
 - (a) a tertiary alcohol
- (b) a primary alcohol
- (c) a secondary alcohol
- (d) none of these

$$\textbf{89.} \ \ CH_3CH_2CH_2OH \xrightarrow{\quad conc. \ H_2SO_4 \\ \quad 160-180^{\circ}C} X \xrightarrow{\quad Br_2 \\ \quad KOH} Z \xrightarrow{\quad alc. \\ \quad KOH} Z$$

In this reaction sequence, Z is

- (c) CH₃—C≡CH
- (d) CH₃CH=CH₂
- 90. An alcohol, on dehydration, produces an alkene which on ozonolysis yields two molecules of acetaldehyde. The alcohol is
 - (a) CH₃CH₂CH₂OH
- (b) CH₃CH₂OH

(c)
$$CH_3$$
— CH — CH_2OH (d) CH_3CH_2CH — CH_3 CH_3

- 91. Glycerine is used in car radiators to
 - (a) facilitate evaporation
 - (b) increase the temperature of the water in the radiator
 - (c) lower the freezing point of the water in the radiator
 - (d) lower the viscosity of the water in the radiator

92.		much bromir 1.5 mol		neede 4.5 r	-		e tribromo _l 3.0 mol		from pl 6.0 mo	
93.		Ortho-nitrophenol is steam volatile, whereas para-nitrophenol is not. This is due to						is not.		
		the presence the presence	of i	ntern	olecula	r hyc	drogen bon	ding in	o-nitro	phenol
		none of thes				,	O	0	,	1
94.	(a)	phenol is he salicylaldeh <i>p</i> -chlorophe	yde	with	CCl ₄ in	(b)	aline KOH, salicylic ac There is n	cid		
95.		phenol is he phenol red salicylic acid		with	phthalio	(b)	nydride and methyl or phenolph	ange	₄ , it pro	duces
96.	In the	carbonyl gro	up, t	he ca	rbon ato					
		sp-hybridize sp ³ -hybridiz					sp ² -hybrid sp ³ d-hybr			
97.	The C	—O—C angl	e in e	ether	is					
	(a)	180°	(b)	109°	28'	(c)	110°	(d)	105°	
98.		an excess of C), the main p			ohol va	pour	is passed	over h	eated a	lumina
		ethylene				(b)	ethane			
	(c)	diethyl ethe	r			(d)	butane			
99.		diethyl ether	is h	eated	with co			it prod	uces	
		ethanol					iodoform	1:40		
	(c)	ethyl iodide					methyl iod			
100.		diethyl ether ethyl chlorid		eated	with ar		ess of PCI ₅ , diethyl etł			
		ethanoyl chl		e			perchloro			
101.		tole reacts w			oroduce	()	1	,		
		$C_6H_5I + CH_5$		•		(b)	CH ₃ CH ₂ I	+ C ₆ H ₅	ОН	
		C ₆ H ₅ CH ₂ OH			H ₂ I		CH ₃ CH ₂ I			
102		eactivity of ha								
.04.		HI > HBr> l	_	ıı acı	as will		HCl > HB			
		HBr > HI > 1					HCl>HI			

103. Diethyl ether can be distinguished from ethyl alcohol by its reaction with (b) PCl₅ (a) Na (c) 2,4-dinitrophenylhydrazine (d) none of these **104.** $CH_3CHCH_3 \xrightarrow{alc.} A \xrightarrow{peroxide} B \xrightarrow{CH_3ONa} C$ In the above reaction sequence, the final product is (a) diethyl ether (b) 1-methoxypropane (c) isopropyl alcohol (d) propylene glycol 105. Aldehydes undergo (a) electrophilic addition (b) electrophilic substitution (c) nucleophilic addition (d) nucleophilic substitution 106. The conversion CH₃CH=CHCHO → CH₃CH=CHCH₂OH can be effected with (b) 9 BBN (a) Ni/H_2 (c) Zn/Hg/HCl (d) none of these 107. Which of the following is the most reactive in nucleophilic addition reactions? (a) HCHO (b) CH₃CHO (c) CH₃COCH₃ (d) CH₃COC₂H₅ 108. Ketones are less reactive than aldehydes because (a) the C=O group is less polar in ketones (b) of the electromeric effect (c) of steric hindrance to the attacking reagent (d) of none of these 109. Aldehydes and ketones form hydrocarbons by (a) the Clemmensen reduction (b) the Cannizzaro reaction (c) the Rosenmund reduction (d) aldol condensation 110. $HC = CH \xrightarrow{HgSO_4} A \xrightarrow{LiAlH_4} B \xrightarrow{P, Br_2} C$ In this reaction sequence, C is (a) ethyl bromide (b) ethylidene bromide (c) ethylene bromide (d) 1,1-dibromoethane

111. The formation of cyanohydrins from ketones is an example of

(c) electrophilic substitution (d) nucleophilic substitution

(b) nucleophilic addition

(a) electrophilic addition

- **112.** When *m*-chlorobenzaldehyde reacts with cold, concentrated KOH at room temperature, the products formed are
 - (a) *m*-hydroxy benzaldehyde and potassium *m*-chlorobenzoate
 - (b) *m*-chlorobenzyl alcohol and *m*-hydroxybenzaldehyde
 - (c) *m*-hydroxybenzyl alcohol and *m*-chlorobenzyl alcohol
 - (d) *m*-chlorobenzyl alcohol and potassium *m*-chlorobenzoate
- 113. The reaction of acetamide with NaOBr in an alkaline medium produces
 - (a) CH_3NH_2

(b) NH₃

(c) $CH_3CH_2NH_2$

- (d) CH₃CN
- 114. Benzaldehyde can be converted into benzyl alcohol by the
 - (a) Claisen reaction
- (b) Perkin reaction
- (c) Cannizzaro reaction
- (d) Wurtz reaction
- 115. Which of the following respond positively to the iodoform test?
 - (a) 2-Pentanone

(b) 1-Pentanal

(c) 3-Pentanone

- (d) Pentanol
- 116. Arrange the following in order of decreasing acidity.

 C_6H_5OH , C_2H_5OH , HCOOH, CH_3COOH

- (a) $HCOOH > CH_3COOH > C_6H_5OH > C_2H_5OH$
- (b) $CH_3COOH > HCOOH > C_2H_5OH > C_6H_5OH$
- (c) $CH_3COOH > C_2H_5OH > HCOOH > C_6H_5OH$
- (d) $C_6H_5OH > C_2H_5OH > HCOOH > CH_3COOH$
- 117. Arrange the following in order of decreasing reactivity in nucleophilic addition reactions.

CH₃CHO, CH₃COCH₃, HCHO, C₂H₅COCH₃

- (a) $HCHO > CH_3CHO > CH_3COCH_3 > C_2H_5COCH_3$
- (b) $CH_3CHO > HCHO > C_2H_5COCH_3 > CH_3COCH_3$
- (c) CH₃CHO > HCHO > CH₃COCH₃ > C₂H₅COCH₃
- (d) $C_2H_5COCH_3 > CH_3COCH_3 > CH_3CHO > HCHO$
- **118.** The reactivities of the carbonyl compounds, HCHO, RCHO and R₂C=O, in nucleophilic addition reactions are in the following order.
 - (a) $HCHO > RCHO > R_2CO$
- (b) $RCHO > HCHO > R_2CO$
- (c) $R_2CO > RCHO > HCHO$
- (d) $HCHO > R_2CO < RCHO$
- 119. In nucleophilic addition reactions, the reactivities of the carbonyl compounds NO₂CH₂CHO (I), CICH₂CHO (II), CH₃CHO (III) and CH₃CH₂CHO (IV) follow the order
 - (a) I > II > III > IV

(b) IV > III > II > I

(c) I > III > II > IV

(d) III > II > IV

120.	. In nucleophilic addition reactions, the reactivities of the carbon compounds CH ₃ COCH ₃ (I), CH ₃ COCH ₂ NH ₂ (II) and CH ₃ COC(CH ₂ (III) follow the order					
	(a)	I>II>III>III	. ,	III > II > I		
	()	II > I > III		II > III > I		
121.		ybridization of C in formaldel	-			
	(a)	*		sp^2		
	(c)	sp^3	(d)	none of these		
122.		reatment of ethylidine chloride		=		
	(a)	CH ₂ OHCH ₂ OH		CH ₃ CHO		
	(c)	НСНО	(d)	CHOCHO		
123.	Methy	ylethylketone can be obtained	by th	ne oxidation of		
	(a)	2-butanol	(b)	2-propanol		
	(c)	1-butanol	(d)	tert. butyl alcohol		
124.	Which	n of the following yields a keto	ne o	n oxidation?		
	(a)	CH ₃ CH ₂ CH ₂ CH ₂ OH	(b)	CH ₃ CH ₂ CHOHCH ₃		
	(c)	(CH ₃) ₂ CHCH ₂ CH ₂ OH	(d)	CH ₃ C(CH ₃) ₂ CH ₂ OH		
125.	CH ₃ C	$E = CH \xrightarrow{H_2SO_4} HgSO_4$				
	The product obtained in this reaction is					
		acetaldehyde		propionic acid		
	(c)	formaldehyde	(d)	acetone		
126.	Methy	yl magnesium chloride reacts	with	acetyl chloride to produce		
	(a)	ethanol	(b)	acetone		
	(c)	methanol	(d)	ether		
127.		the product formed by the re 3MgI is hydrolysed, it yields	actio	on of HCOOC ₂ H ₅ with an excess		
		<i>n</i> -propyl alcohol	(b)	isopropyl alcohol		
		propanol		ethanol		
128.	In the	Rosenmund reduction, the ca	talys	t used is		
	(a)	Pd/BaSO ₄	(b)	Raney Ni		
	(c)	Sn/HCl	(d)	Zn/HCl		
129.	In the	reaction CH ₃ CH ₂ COCl ———————————————————————————————————	H ₂	$\xrightarrow{O_4} X$, the product is		
	(a)	propanaldehyde	(b)	acetaldehyde		
		acetic acid		acetone		

130.	The re	educing agent used in the Stepl	hen 1	reduction is
	(a)	Sn/HCl	(b)	Zn/HCl
	(c)	SnCl ₂ /HCl	(d)	Na-Hg/alcohol
131.	The o	xidation of benzene by V_2O_5 in	the	presence of air produces
	(a)	benzoic acid	(b)	benzaldehyde
	(c)	benzoic anhydride	(d)	maleic anhydride
132.	C_6H_6	+ CO + HCl $\xrightarrow{\text{anhyd.AlCl}_3}$		
		nain product obtained in this re		
	(a)	$C_6H_5CH_3$ (b) C_6H_5CHO	(c)	$C_6H_5CH_2Cl$ (d) C_6H_5COOH
133.	Aldeh	nydes and ketones undergo add	ditio	n reactions with
	(a)	phenyl hydrazine	(b)	hydrazine
	(c)	semicarbazide	(d)	hydrogen cyanide
134.		order of reactivity of CH_3COCH_3 (III) is	СНО	(I), CH ₃ CH ₂ COCH ₃ (II) and
		I > III > II	(b)	I > II > III
	(c)	II > I > III	(d)	II > II > I
135.		of an organic compound requ The compound is	ires (0.5 mol of oxygen to produce an
	(a)	an alcohol	(b)	an aldehyde
	(c)	a ketone	(d)	an ether
136.	The effects)—	→CH ₃ CH=CHCOOH can be
	(a)	alkaline KMnO ₄	(b)	acidic K ₂ Cr ₂ O ₇
	(c)	ammoniacal AgNO ₃	(d)	selenium dioxide
137.	respo		irror	cular formula C ₃ H ₆ O does not test with Tollens reagent but
	(a)	CH ₂ =CHCH ₂ OH	(b)	CH ₃ CH ₂ CHO
	(c)	CH ₂ =CHOCH ₃	(d)	CH ₃ COCH ₃
138.		aldol condensation of acetalden nediate product which is	ehyd	e involves the formation of an
	(a)	a carbocation	(b)	a carbanion
	(c)	an acetate ion	(d)	a free radical
139.	When	acetone is saturated with HCl	gas,	the final product obtained is
	(a)	diacetone alcohol	(b)	phorone
	(c)	mesityl oxide	(d)	benzene

- 140. CH₃CHO and HCHO can be differentiated using
 - (a) Fehling's solution
- (b) the Tollens reagent
- (c) the Schiff reagent
- (d) a caustic soda solution
- 141. The reaction $C_6H_5CHO + CH_3CHO \longrightarrow C_6H_5CH = CHCHO$ is called
 - (a) Claisen condensation
- (b) aldol condensation
- (c) benzoic condensation
- (d) polymerization

- **142.** Paraldehyde is a
 - (a) dimer of formaldehyde (b) trimer of acetaldehyde

 - (c) hexamer of formaldehyde (d) hexamer of acetaldehyde

$$\textbf{143.} \quad C_3H_8O \xrightarrow{\hspace{1cm} [O] \hspace{1cm}} K_2Cr_2O_7/H_2SO_4 \xrightarrow{\hspace{1cm}} C_3H_6O \xrightarrow{\hspace{1cm} I_2 \text{ alk.} \hspace{1cm}} CHI_3$$

In this reaction the first compound is

- (a) CH₃CH₂CH₂OH
- (b) CH₃CH—CH₃

(c) CH₃OCH₂CH₂

- (d) CH₃CH₂CHO
- 144. An organic compound X with the molecular formula C₅H₁₀O yields phenyl hydrazone and gives a negative response to the iodoform test and Tollens test. It produces *n*-pentane on reduction. The compound could be
 - (a) pentanal

(b) pentanone-2

(c) pentanone-3

(d) amyl alcohol

145.
$$C_6H_5CHO + CH_3COCH_3 \xrightarrow{OH^-} C_6H_5CH(OH)CH_2COCH_3$$

 $\longrightarrow C_6H_5CH=CHCOCH_3$

This reaction is known as

- (a) aldol condensation
- (b) cross aldol condensation
- (c) the Claisen-Schmidt reaction
- (d) none of these
- 146. The conversion of acetone into diacetone alcohol is carried out in the presence of
 - (a) dry HCl gas

(b) concentrated H₂SO₄

(c) $Ba(OH)_2$

(d) heat

147.
$$CH_3CH=CH_2 + CO + H_2O \xrightarrow{H_3PO_4} CH_3-CH-COOH CH_3$$

This reaction is called

- (a) the Stevens reaction
- (b) the carbonylation reaction
- (c) the Gattermannan-Koch reaction
- (d) oxidation
- **148.** Arrange the following in order of decreasing acid strength. HCOOH (I), CH₃COOH (II), CH₃COOH (III), (CH₃)₂CHCOOH (IV)
 - (a) I > II > III > IV

(b) IV > III > II > I

(c) II > I > IV > III

- (d) I > III > II > IV
- 149. The acid strengths of the following decrease in the order
 - (a) CCl₃COOH > CHCl₂COOH > CH₂ClCOOH > CH₃COOH
 - (b) CH₃COOH > CH₂CICOOH > CHCl₂CO OH > CCl₃COOH
 - (c) CCl₃COOH > CH₃COOH > CH₂CICO OH > CHCl₂COOH
 - (d) CH₂ClCOOH > CHCl₂COOH > CCl₃COOH > CH₃COOH
- **150.** The acidic strengths of chloro, bromo, iodo and fluoro acetic acids decrease in the order
 - (a) FCH₂COOH > CICH₂COOH > BrCH₂COOH > ICH₂COOH
 - (b) CICH₂COOH > BrCH₂COOH > FCH₂COOH > ICH₂COOH
 - (c) ICH₂COOH > BrCH₂COOH > ClCH₂COOH > FCH₂COOH
 - (d) BrCH₂COOH > ClCH₂COOH > FCH₂COOH > ICH₂COOH
- 151. Methanoic acid is manufactured by the reaction of carbon monoxide and
 - (a) NaOH

- (b) dilute HCl
- (c) concentrated H₂SO₄
- (d) PCl₅

152.
$$RCOOAg + Br_2 \xrightarrow{CCl_4} R - Br + AgBr + CO_2$$

This reaction is called the

- (a) Wurtz reaction
- (b) Hunsdiecker reaction
- (c) Friedel-Crafts reaction
- (d) Kolbe reaction

153.
$$CH_3CH_2COOH \xrightarrow{\text{Red P/Br}_2} CH_3CH_2CH$$
—COOH
Br

This reaction is called the

- (a) Cannizzaro reaction
- (b) Schrödinger reaction
- (c) Hell-Volhard-Zelinsky reaction
- (d) Reimer-Tiemann reaction

154. Phthalic acid $\xrightarrow{\Delta}$ A $\xrightarrow{NH_3}$ B \xrightarrow{NaOH} C $\xrightarrow{Br_2/KOH}$ D \xrightarrow{HCl} E In this reaction, the product E is

- (a) *o*-nitrobenzoic acid
- (b) salicylic acid
- (c) anthranilic acid
- (d) crotonic acid
- 155. Which of the following carboxylic acids undergoes decarboxylation easily?
 - (a) C₆H₅COCH₂COOH
- (b) C₆H₅COCOOH
- (c) C_6H_5CH —COOH (d) $C_6H_5CHCOOH$ OH NH_2
- **156.** Which of the following is the most acidic?

OH CH₃

(c)

- (d)
- 157. In the Cannizzaro reaction

$$2C_6H_5CHO \xrightarrow{OH^-} C_6H_5CH_2OH + C_6H_5COO^-$$

the slowest step is

- (a) the attack by OH on the carbonyl group
- (b) the transfer of the hydride to the carbonyl group
- (c) the exchange of protons can be slow steps
- (d) all the above
- 158. Arrange the following in order of decreasing reactivity with concentrated HCl.

(a) I > II > III > IV

(b) IV > III > II > I

(c) I > III > II > IV

(d) IV > III > I > II

159. COOCH₃ H — OH

Α

- H OH
 H OH
 COOCH
- COOH

 HO H

 COOCH₃

Which of the following statements about the compounds given above is true?

(d)

- (a) A and B are identical.
- (b) A and B are diastereomers.
- (c) A and C are enantiomers.
- (d) A and B are enantiomers.
- **160.** Among the following, which intermediate is the best hydride donor in a Cannizzaro reaction?
 - (a) H $C_6H_5-C-O^-$ OH

- (c) H O^-
- CH_3O O^-
- **161.** Mesotartaric acid and *d*-tartaric acid are
 - (a) diastereomers

(b) racemic mixtures

(c) enantiomers

- (d) position isomers
- 162. Racemic tartaric acid is optically inactive due to
 - (a) internal compensation
- (b) loss of asymmetric centre
- (c) external compensation
- (d) steric hindrance
- **163.** Which of the following para-substituted benzoic acids is the most acidic?
 - (a) Cl—C₆H₅COOH
- (b) NO_2 — C_6H_4COOH
- (c) HO—C₆H₄—COOH
- (d) CH₃—C₆H₄—COOH
- **164.** Which of the following alcohols has the highest boiling point?
 - (a) CH₃CH₂CH₂CH₂OH
- (b) CH₃C—CH₂OH

- **165.** Which of the following has the highest boiling point?
 - (a) CH₃CH₂CH₂CH₂CH₃
- (b) CH₃CH₂OCH₂CH₃
- (c) CH₃CH₂CH₂Cl
- (d) CH₃CH₂CH₂CH₂OH
- **166.** An ether reacts with H₂SO₄ to form
 - (a) an alkyl free radical
- (b) a zwitterion
- (c) an oxonium ion
- (d) an oxy anion
- 167. In the reaction $R-S-R-\frac{[O]}{KMnO_4} \times X$, the product obtained is

 (a) a sulphoxide
 (b) a sulphone
 (c) an alkane
 (d) an alkane

(c) an alkane

- (d) an alkene
- 168. Arrange the following in order of decreasing reactivity with nucleophiles.

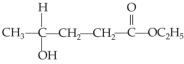
Acetone (I), ethylmethylketone (II), diethylketone (III)

(a) I > II > III

(b) III > II > I

(c) II > I > III

(d) III > I > II



This conversion is effected by using

(a) NaBH₄

(b) LiAlH₄

(c) Pd-C

- (d) Raney Ni/H₂
- 170. The conversion $CH_3CHO \longrightarrow OHC \longrightarrow CHO$ can be effected by
- (a) CrO_3 (b) SeO_2 (c) $Br_2/NaOH$ (d) $KMnO_4$
- 171. Ethylmethylketone, on heating with aluminium isopropoxide and isopropyl alcohol, gives
 - (a) CH₃CH₂CH—CHCH₂CH₃ (b) CH₃CH=CHCH₃ CH₃ CH₃
 - (c) CH₃CH(OH)CH₂CH₃ (d) CH₃CH₂CH₂CH₃

- 172. Among the following, which are the most acidic?
- (b) CH₃CH(Cl)CH₂COOH
- (c) CH₃CHClCH₂CH₂COOH (d) ClCH₂CH₂CH₂CH₂COOH
- 173. The hydrolysis of which of the following takes the longest time?
 - (a) CH₂COCl

- (b) CH₂COOCOCH₂
- (c) $CH_3C-OC_2H_5$ (d) CH_3-C-NH_7
- 174. The reactivities of acid halides (I), anhydrides (II), esters (III) and amides (IV) with nucleophilic reagents follow the order
 - (a) I > II > III > IV

(b) IV > III > II > I

(c) I > III > II > IV

- (d) III > II > I > IV
- **175.** Which of the following reactions are feasible?

(b)
$$CH_3$$
 CH_3 CH_3

- (c) Both (a) and (b)
- (d) Neither

176.
$$H_2C \xrightarrow{CH_2 \xrightarrow{(i) CH_3MgCl}} X$$

The product obtained in this reaction is

- (a) CH₃CH₂OH
- (b) (CH₃)₂CHOH
- (c) CH₃CH₂CH₂OH
- (d) HO--CH₂---CH₂---CH₂---OH

177.
$$(CH_3)_2CHCHOH \xrightarrow{acid} X$$

The major product obtained in this reaction is

- (a) $(CH_3)_2CHCH=CH_3$
- (b) (CH₃)₂C=CH—CH₂
- (c) a 1:1 mixture of (a) and (b) (d) none of these
- 178. Phenol is soluble in water because
 - (a) of weak hydrogen bonding between phenol and water molecules
 - (b) of intermolecular hydrogen bonding between phenol molecules
 - (c) it has a higher boiling point than that of water
 - (d) of none of these
- 179. The acidities of primary, secondary and tertiary alcohols follow the order
 - (a) primary > secondary > tertiary
 - (b) secondary > tertiary > primary
 - (c) tertiary > secondary > primary
 - (d) primary > tertiary > secondary
- **180.** The ease of dihydration of $\mathring{1}$, $\mathring{2}$ and $\mathring{3}$ alcohols follows the order
 - (a) $\mathring{1} > \mathring{2} > \mathring{3}$ (b) $\mathring{2} > \mathring{3} > \mathring{1}$ (c) $\mathring{3} > \mathring{2} > \mathring{1}$ (d) $\mathring{3} > \mathring{1} > \mathring{2}$

- 181. Arrange acetyl chloride (I), ethyl acetate (II), acetamide (III) and acetic anhydride (IV) in order of reactivity towards nucleophilic acyl substitution.
 - (a) I > IV > II > III

(b) I > II > III > IV

(c) III > II > IV > I

- (d) IV > III > II > I
- **182.** Which of the following is the best method for making isopropylmethyl ether?
 - (a) $CH_3I + (CH_3)_2CHOH \longrightarrow$
 - (b) $CH_3I + (CH_3)_2CHO^- \longrightarrow$
 - (c) $(CH_3)_2CHI + CH_3O^- \longrightarrow$
 - (d) (CH₃)₂CHCl + CH₃OH →
- 183. What will be the order of reactivity of the following carbonyl compounds with Grignard's reagent?

$$H > C = O$$
 $H_3 C = O$ $H_3 C > C = O$ $(CH_3)_3 C > C = O$

- (a) I > II > III > IV
- (b) IV > III > II > I
- (c) II > I > IV > III

(d) III > II > IV

184. In the bromination of acetone, the following three reactions occur.

$$CH_3COCH_3 \longrightarrow CH_3COCH_2Br(I)$$
 $CH_3COCH_2Br \longrightarrow CH_3COCHBr_2$ (II)
 $CH_3COCHBr_2 \longrightarrow CH_3COCBr_3$ (III)

Which of the following reflects the relative ease with which these reactions take place?

(a) I < II < III

(b) III < II < I

(c) I < II = III

(d) II < I < III

185. On treatment with an alkali, glyoxal gives glycolic acid:

OHC—CHO
$$\stackrel{\mathrm{OH}}{\longrightarrow}$$
 HOCH₂—COOH

To which type does this reaction belong?

- (a) Aldol condensation
- (b) Knowvenagel condensation
- (c) Cannizzaro reaction
- (d) None of these

186. Give the order of ease of the esterification of the following acids.

$$O_2N$$
 COOH

 OCH_2

IV

(a) I > II > III > IV

(b) IV > III > II > I

(c) II > I > IV > III

(d) I > II > III = IV

187. Give the order of ease of decarboxylation of the following acids.

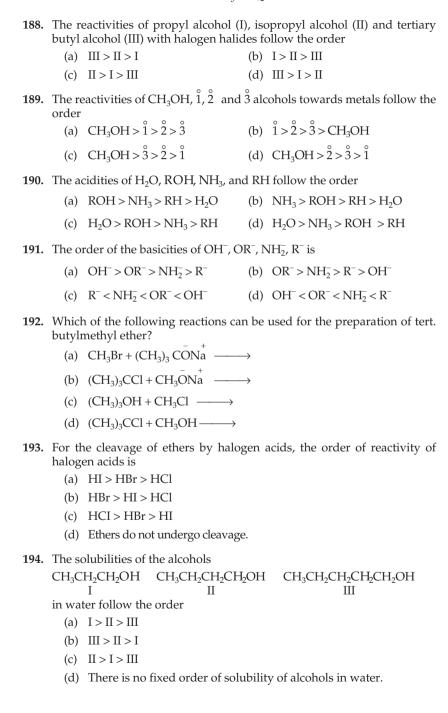
CH₃COOH CH₂=CH—CH₂COOH CH₂(COOH)₂ III III $O_2N \xrightarrow{NO_2} COOH$ NO_2 IV

(a) I > II > III > IV

(b) III > IV > II > I

(c) IV > III > II > I

(d) I > III > II > IV



196. The boiling points of isomeric $\mathring{1}$, $\mathring{2}$ and $\mathring{3}$ alcohols decrease in the order

197. Arrange CH₃OH, C₂H₅OH, (CH₃)₂CHOH and (CH₃)₃COH in order of

(b) 3 > 2 > 1

(b) 3 > 1 > 2

(d) There is no fixed order.

(d) There is no fixed order.

195. The hydrogen bonding ability of $\overset{\circ}{1}$, $\overset{\circ}{2}$ and $\overset{\circ}{3}$ alcohols is of the order

(a) $\mathring{1} > \mathring{2} > \mathring{3}$

(c) $\hat{1} > \hat{3} > \hat{2}$

(a) 3 > 2 > 1

(c) $\mathring{1} > \mathring{2} > \mathring{3}$

	decreasing acidity.					
	(a)	$CH_3OH > C_2H_5OH > (CH_3)_2CHOH > (CH_3)_3COH$				
	(b)	$(CH_3)_3COH > (CH_3)_2CHOH > C_2H_5OH > CH_3OH$				
	(c)	$C_2H_5OH > CH_3OH > (CH_3)_3C$	COH> (CH ₃) ₂ CHOH			
	(d)	There is no fixed order.				
198.	order		des of $\mathring{1}$, $\mathring{2}$ and $\mathring{3}$ alcohols follows the			
	(a)	$\hat{1} > \hat{2} > \hat{3}$ (b) $\hat{1} > \hat{3} > \hat{2}$	(c) $\mathring{3} > \mathring{2} > \mathring{1}$ (d) $\mathring{1} = \mathring{2} = \mathring{3}$			
199.	Ether	s react with concentrated H ₂ SO	O_4 to form			
	(a)	zwitterions	(b) an alkyl free radical			
	(c)	oxyanions	(d) oxonium ions			
200.	that b (a) (b) (c)	etween molecules of aldehyde weak strong equal	traction between alcohol molecules, s and ketones is attraction between molecules of			
201.	201. In ketones, the reactivites of the carbonyl group of the compounds ethylmethyl ketone (I), diethyl ketone (II) and acetone (III), decrease in the order					
	(a)	I > II > III	(b) $III > II > I$			
	(c)	I > III > II	II < I > II			
202.	02. The reactivities of the carbonyl compounds formaldehyde (I), acetaldehyde (II) and acetone (III) towards nucleophiles decrease in the order					
	(a)	I > II > III	(b) $III > II > I$			
	(c)	III > I > III	(d) III > I > II			

203.	The conversion of acetone into achieved by	2,3-dimethylbutane-2,3-diol can be
	(a) using Zn/Hg/HCl	(b) Wolff–Kishner reduction
	(c) using Mg/Hg/H ₂ O	(d) The conversion is not possible.
204.	How many structures can comp $C_4H_8O_2$ have?	oounds with the molecular formula
	(a) Five	(b) Four
	(c) Three	(d) Two
205.	Arrange the following compounds CICH ₂ CH ₂ CH ₂ COOH CH ₃ CHC	in order of decreasing acidity. ICH ₂ COOH CH ₃ CH ₂ CHCICOOH II III
	(a) $I > II > III$	(b) $III > II > I$
	(c) $I > III > II$	(d) $III > I > II$
206.	Arrange the following carboxylic a Oxalic acid Malonic acid I II (a) I > II > III	cids in order of decreasing acidity. Succinic acid III (b) III > II > I
	(c) $I > III > II$	(d) $II > III > I$
207.	Arrange OHCH ₂ COOH(I), HOCH in order of acidity.	H ₂ CH ₂ COOH (II) and CH ₃ COOH (III)
	(a) I > II > III	(b) $III > II > I$
	(c) $I > III > II$	(d) $II > III > I$
208.	The ease of hydrolysis with an alka CH_3COCl $CH_3CO-O-COCH_3$ I II is of the order	$\begin{array}{c} \text{lli in the compounds} \\ \text{CH}_3\text{COOC}_2\text{H}_5 & \text{CH}_3\text{CONH}_2 \\ \text{III} & \text{IV} \end{array}$
	(a) $I > II > III > IV$	(b) $IV > III > II > I$
	(c) $I > II > IV > III$	(d) $II > I > IV > III$
209.	RCONH ₂ (IV) with nucleophilic real	
	(a) $I > II > III > IV$	(b) $IV > III > II > I$
	(c) $I > II > IV > III$	(d) II > I > IV > III
210.	The boiling points of acetic anhydbutyric anhydride (III) follow the o	ride (I), propionic anhydride (II) and rder
	(a) $I < II < III$	(b) $III < II < I$

(d) II < III < I

(c) I < III < II

211. The reaction

$$\begin{array}{c} O \\ \parallel \\ CH_3-C-OC_2H_5+n-C_3H_7OH- \xrightarrow{\quad n-C_3H_7ONa \quad \\ \quad + C_7H_5OH \end{array}$$

is known as

(a) esterification

- (b) double decomposition
- (c) transesterification
- (d) none of these
- **212.** The reaction of acetone with CN⁻ is represented as

$$\begin{array}{c} \text{CH}_{3} \\ \text{CH}_{3} \end{array} \text{C=O} \xrightarrow{\text{Step (I)}} \begin{array}{c} \text{CH}_{3} \\ \text{CH}_{3} \end{array} \begin{array}{c} \overset{\cdot}{\text{C}} \\ \text{CH}_{3} \end{array} \begin{array}{c} \overset{\cdot}{\text{C}} \\ \text{Step (II)} \end{array} \begin{array}{c} \text{CN} \\ \text{CH}_{3} \\ \text{Step (III)} \end{array} \begin{array}{c} \text{CN} \\ \text{Step (III)} \end{array} \begin{array}{c} \text{CN} \\ \text{CH}_{3} \\ \text{C} \end{array} \begin{array}{c} \text{CN} \\ \text{CH}_{3} \\ \text{CH}_{3} \end{array} \begin{array}{c} \text{CN} \\ \text{CH}_{3} \\ \text{CH}_{3} \end{array} \begin{array}{c} \text{CN} \\ \text{CH}_{3} \\ \text{CH}_{3} \\ \text{CH}_{3} \end{array} \begin{array}{c} \text{CN} \\ \text{CH}_{3} \\ \text{CH}_{3} \\ \text{CH}_{3} \end{array} \begin{array}{c} \text{CN} \\ \text{CH}_{3} \\ \text{CH}_{4} \\ \text{CH}_{4}$$

In the above reaction, which is the fast step?

- (a) Step (I)
- (b) Step (II)
- (c) Step (III)
- (d) All the steps take place with equal ease.
- **213.** The polarization

may be attributed to the

- (a) inductive effect
- (b) electromeric effect
- (c) mesomeric effect
- (d) hyperconjugative effect
- **214.** Arrange CH₃CH₂OH (I), (CH₃)₂CHOH (II) and (CH₃)₃COH (III) in order of increasing acidity.
 - (a) I > II > III

(b) III > II > I

(c) I > III > II

- (d) II > I > III
- **215.** Arrange *p*-nitrophenol (I), *o*-nitrophenol (II), *m*-nitrophenol (III) and phenol (IV) in order of acidity.
 - (a) I > III > IV > II

(b) I > II > III > IV

- (c) IV > III > II > I
- (d) III > I > IV > II
- **216.** Arrange 2,4,6-trinitrophenol (I); 2,4-dinitrophenol (II); *o*-nitrophenol (III) and *p*-nitrophenol (IV) in order of acidity.
 - (a) I > II > III = IV
- (b) IV > III > II > I
- (c) I > II > IV > III

(d) III > II > IV

217.	Arrange phenol (I), <i>o</i> -cresol (II), <i>p</i> -cresol (III) and <i>m</i> -cresol (IV) in order of decreasing acid strength.				
	(a)	I > IV > III > II	(b)	I > II > III > IV	
	(c)	IV > III > II > I	(d)	II > I > IV > III	
218.				with methyl alcohol (I), ethyl rt. butyl alcohol (IV) follows the	
	(a)	I > II > III > IV	(b)	IV > III > II > I	
	(c)	II > I > IV > III	(d)	III > IV > I > II	
219.	The (CH ₃) order	rate of esterification of ₂ CHCOOH (III) and (CH ₃) ₃ CO		COOH (I), CH ₃ COOH (II), H (IV) with ethanol follows the	
	(a)	IV > III > II > I	(b)	I > II > III > IV	
	(c)	II > I > IV > III	(d)	III > IV > I > II	
220.	hydro (a)	lysis, the main product obtain o-hydroxybenzaldehyde	ed is (b)	<i>p</i> -hydroxybenzaldehyde <i>p</i> -hydroxybenzoic acid	
221		ene glycol, on being distilled w			
221.	-	ethylene oxide		acetaldehyde	
	(c)	1,4-dioxane	` ′	diethylene glycol	
222.	Glyce	rol, on being heated with an ex	cess	of hydriodic acid, gives	
	(a)	allyl iodide		isopropyl iodide	
	(c)	acrolein		glyceraldehyde	
223.	The o	rder of dehydration of $1, 2$ and	3° alo	cohols is	
				$\mathring{2} > \mathring{1} > \mathring{3}$ (d) $\mathring{1} > \mathring{3} > \mathring{2}$	
224.		ge formaldehyde (I), acetaldeh vity towards nucleophilic addi		(II) and acetone (III) in order of	
	(a)	II > II > II	(b)	I > II > III	
	(c)	II > I > III	(d)	II < I > III	
225.	Arran CH ₃ C	ge $[(CH_3)_3C]_2CO$ (I), $[(CH_3)_2O]_2CO$ (IV) in order of reactivity t	owa		
	(a)	I > II > III > IV	(b)	I > III > IV > II	
	(c)	IV > III > II > I	(d)	II > I > III > IV	
226.	CH ₃ C	H=CH—CHO may be reduced	d to (CH ₃ CH=CHCH ₂ OH using	
		H ₂ /Pt	` '	NaBH ₄	
	(c)	[(CH ₃) ₂ CHO] ₃ Al	(d)	Zn-Hg/HCl	

227. The reaction of benzaldehyde with excess of acetone in dilute NaOH at

228. An organic compound reduces Tollens reagent and Fehling's solution. It

(b) benzalacetone

(d) none of these

(b) C₆H₅CHO

(d) (CH₃)₃CCOCH₃

273 K gives

can be

(a) benzalacetophenone

(c) dibenzalacetone

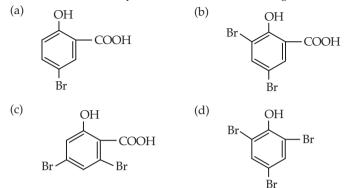
(a) CH₃CH₂CHO

(c) CH₃COCH₂CH₃

229. Benzaldehyde reacts with ammonia to form

	(a)	C_6H_5 OH NH_2	(b)	C ₆ H ₅ CH=NH		
	(c)	$(C_6H_5CH=N)_2CHC$	C_6H_5 (d)	none of these		
230.	prese (a)	nce of a catalytic amo	ount of alumin one (b)	nine in ether at 273 K in the ium trichloride gives <i>m</i> -bromoacetophenone phenacylbromide		
231.				centrated H ₂ SO ₄ at 273 K, gives		
	` '	acetic acid	` '	paraldehyde		
		metaldehyde		none of these		
232.		ige formic acid (I ylacetic acid (IV) in o		id (II), acetic acid (III) and		
		I > II > III > IV	•	IV > III > II > I		
	(c)	$\mathrm{I}>\mathrm{II}>\mathrm{IV}>\mathrm{III}$	(d)	III > IV > II > I		
233.	233. Arrange <i>p</i> -nitrobenzoic acid (I), <i>p</i> -chlorobenzoic acid (II), benzoic acid (III), <i>p</i> -toluic acid (IV) and <i>p</i> -hydroxybenzoic acid (V) in order of decreasing acidity.					
	` '	I > II > IV > V > III	()	III > I > II > IV > V		
	(c)	II > IV > I > III > V	(d)	I > II > III > IV > V		
234.	34. Arrange <i>o</i> -toluic acid (I), <i>m</i> -toluic acid (II), <i>p</i> -toluic acid (III) and benzoic acid (IV) in order of decreasing acid strength.					
	(a)	I>II>III>IV	(b)	I > II > IV > III		
	(c)	III > IV > I > II	(d)	I > IV > II > III		
235.	p-hyd			<i>m</i> -hydroxybenzoic acid (II), acid (IV) in order of decreasing		
	(a)	I > II > IV > III	(b)	I > II > III > IV		
	(c)	IV > III > II > I	(d)	II > III > I > IV		

236. The bromination of salicylic acid with bromine water gives



• *Type 2* •

Choose the correct options. More than one option is correct.

237. The synthesis of

$$\begin{array}{c} {\rm CH_3} \\ {\rm C_6H_5CH_2-C-CH_2-CH_3} \\ {\rm OH} \end{array}$$

can be achieved by

(a)
$$CH_3CH_2CCH_3 + C_6H_5CH_2MgCl \longrightarrow \begin{bmatrix} & & & \\ & & & \\ & & & \\ & & & \end{bmatrix} \xrightarrow{H_2O}$$

(b)
$$C_6H_5CH_2CCH_3 + CH_3CH_2MgBr \longrightarrow \begin{bmatrix} & & & \\ & & & \\ & & & \\ & & & \end{bmatrix}$$

(c)
$$C_6H_5CH_2CCH_2CH_3 + CH_3Mgl \longrightarrow \begin{bmatrix} \\ \\ \end{bmatrix} \xrightarrow{H_2O}$$

(d) none of these

238. Which of the following can be used for the synthesis of 2-methylbut-3-yne-2-ol?

(c)
$$O$$
 $HC \equiv CMgBr + CH_3 - C - CH_3$

(d)
$$\begin{array}{c} O \\ \parallel \\ HC \equiv CMgBr \ + \ CH_3 - CH \end{array}$$

239. Which of the following reactions will yield *p*-tert. butylphenol?

(a)
$$CH_3$$

phenol + CH_3 $C=CH_2$ $\xrightarrow{H^+}$

(b) Phenol +
$$(CH_3)_3COH \xrightarrow{H^+} AlCl_3 \rightarrow$$

(c) Phenol + $(CH_3)_3CCl \xrightarrow{}$

- (d) None of these

240. Which of the following reactions can be used to prepare

$$CH_{3} - C - C_{6}H_{5}$$

$$C_{2}H_{5}$$

(a)
$$O$$
 $CH_3 - C - C_6H_5 + C_2H_5MgBr \longrightarrow$

(b)
$$O$$
 $C_2H_5-C-C_6H_5+CH_3MgBr \longrightarrow \boxed{\frac{H_3O^+}{}}$

(d) Br
$$CH_3$$
— C_6H_5 C_7H_5 C_7H_5

241. Acetaldehyde is obtained in the reactions

(a)
$$CH_2$$
= CH - CH_2 - CH = CH_2 $\xrightarrow{1. O_3}$ $\xrightarrow{2. Zn. H.O}$

(b)
$$CH_3CH = \underbrace{\begin{array}{c} 1.O_3 \\ \hline 2.Zn, H_2O \end{array}}$$

(c)
$$HC \equiv CH + H_2O \xrightarrow{H_2SO_4} H_2SO_4$$

(d)
$$CH_3COCl + H_2 \xrightarrow{Pd-BaSO_4}$$

242. Benzophenone can be obtained by

$$C_6H_6 + C_6H_5C-CI \xrightarrow{AlCl_3}$$

(b)
$$(C_6H_5)_2CHOH \xrightarrow{\qquad CrO_3 \qquad } H_2SO_4 \xrightarrow{}$$

(c)
$$(C_6H_5)_2C = CH_2 \xrightarrow{1. O_3} \frac{1. O_3}{2. Zn, H_2O}$$

(d) none of these

243. Which of the following can be used for the synthesis of benzyl acetate?

(a)
$$C_6H_5CH_2OH + (CH_3CO)_2O \longrightarrow$$

(b)
$$C_6H_5CH_2OH + CH_3COOH \xrightarrow{H^+}$$

(c)
$$O$$
 \parallel $C_6H_5CH_2OH + CH_3CCI \xrightarrow{base}$

(d)
$$C_6H_5CH_2OH + CH_3COOH \xrightarrow{NaOH}$$

244. In which of the following reactions is benzoic acid the major product?

(a)
$$C_6H_5CH_2OH \xrightarrow{1. \text{KMnO}_4, \text{OH}^-} 2. \text{H}_3\text{O}^+$$

(b)
$$O$$
 H $C_6H_5C-CH_3 \xrightarrow{1. Cl_2, NaOH} 2. H_3O^+$

(c)
$$C_6H_5MgBr \xrightarrow{1. CO_2} 2. H_3O^+$$
(d) $C_6H_5CH_3 \xrightarrow{2. H_3O^+} 2. H_3O^+$

(d)
$$C_6H_5CH_3 \xrightarrow{1. \text{KMnO}_4, \text{OH}^-, \Delta}$$

245. Acid anhydrides can be prepared by

(a)
$$C_6H_5COOH + CH_3COCI \xrightarrow{pyridine}$$

(b)
$$C_6H_5COO^-Na^+ + C_6H_5COCl \longrightarrow$$

(c)
$$CH_3CONH_2 + CH_3CO\bar{O}Na^+ \longrightarrow$$

246. Which of the following will give cyclic products upon being heated or being treated by an acid.

- **247.** Which of the following react to give the usual products?
 - (a) Di-tert. butyl ketone + methyl magnesium bromide
 - (b) Methyl isopropyl ketone + methyl magnesium bromide
 - (c) Methyl isopropyl ketone + tert. butyl magnesium bromide
 - (d) Acetamide + methyl magnesium bromide
- **248.** The product obtained in the reaction

$$CH_3CH = CH - C - CH_3$$
 $\xrightarrow{1. C_2H_5Mgl}$ $\xrightarrow{2. H_3O^+}$

is

(a)
$$CH_3 - CH - CH_2 - C - CH_3$$

 C_2H_5

(b)
$$\begin{array}{c} OH \\ CH_3-CH=CH-C-CH_3 \\ C_2H_5 \end{array}$$

(c) OH
$$|$$
 CH $_3$ CH $_2$ - CH $_2$ - C - CH $_3$ $|$ C $_2$ H $_5$

(d) OH
$$CH_3-CH_2-C-CH_2CH_3$$
 C_2H_5

249. On treatment with a concentrated solution of zinc chloride in concentrated HCl at room temperature, an alcohol immediately gives, an oily product. The alcohol can be

250. Which of the following alcohols can be oxidized by potassium dichromate in the presence of sulphuric acid?

$$\begin{array}{c} \text{(d)} & \text{CH}_3 \\ \text{CH}_3 - \text{C} - \text{OH} \\ \text{CH}_3 \end{array}$$

251. The reaction of isopropylbenzene with oxygen in the presence of a catalytic amount of HBr followed by treatment with an acid gives phenol. The reaction proceeds through the intermediate formation of

(c)
$$C_{6}H_{5}-C_{-}O-O$$
.

- 252. Which of the following statements are correct about a carbonyl group?
 - (a) The carbonyl carbon is sp ²-hybridized
 - (b) The carbonyl carbon is sp ³-hybridized
 - (c) The three groups attached to the carbonyl carbon lie in the same plane.
 - (d) The three groups attached to the carbonyl carbon lie in different planes.
- **253.** On treatment with a clear solution of CrO_3 in dilute H_2SO_4 , an aliphatic alcohol gives a greenish opaque solution within 2–3 seconds. The alcohol can be

- **254.** The oxidation of which of the following compounds with hot alkaline KMnO₄ followed by treatment with an acid will give benzoic acid?
 - (a) $C_6H_5CH_3$

(b) C₆H₅CH=CHCH₃

(c)
$$C_6H_5C \equiv CCH_3$$

- $\begin{array}{ccc} \text{(d)} & & \text{O} \\ & & \text{II} \\ & \text{C}_6\text{H}_5\text{CCH}_2\text{CH}_3 \end{array}$
- 255. Which of the following statements are correct?
 - (a) o-Nitrophenol can be separated from p-nitrophenol because of intramolecular hydrogen bonding in o-nitrophenol.
 - (b) *o*-Nitrophenol can be separated from *p*-nitrophenol because of intermolecular hydrogen bonding in *o*-nitrophenol.

- (c) o-Hydroxybenzoic acid can be separated from p-hydroxybenzoic acid because of intramolecular hydrogen bonding in o-hydroxybenzoic acid.
- (d) *o*-Hydroxybenzoic acid can be separated from *p*-hydroxybenzoic acid because of intermolecular hydrogen bonding in *o*-hydroxybenzoic acid.
- **256.** What are the products expected in the following reaction?

$$C_{6}H_{5}-\overset{O}{C}-CH_{2}-\overset{O}{C}-CH_{3}\xrightarrow{CH_{2}N_{2}} \underbrace{}_{ether}$$
(a) $\overset{O}{O}CH_{3}\overset{O}{O}\overset{O}{O}CH_{3}$
 $C_{6}H_{5}\overset{C}{C}=CH-\overset{C}{C}-CH_{3}$
(b) $\overset{O}{O}CH_{3}\overset{C}{O}CH_{3}$
 $C_{6}H_{5}\overset{C}{C}-CH=\overset{C}{C}-CH_{3}$
(c) $\overset{O}{O}CH_{3}\overset{O}{O}\overset{O}{O}CH_{3}$
 $C_{6}H_{5}-\overset{C}{C}-CH-\overset{C}{C}-CH_{3}$
 $C_{6}H_{5}-\overset{C}{C}-CH-\overset{C}{C}-CH_{3}$

257. Under the influence of acids, acetaldehyde gives

258. The final product of the reaction

$$(CH_2)_5$$
 $C = O \xrightarrow{1. PCl_5}$ CH_2

is (a)
$$CH_2$$
 OH (b) CH_2 CH_2 OH CH_2 CH_2 CH_2 CH_2 CH_2

- (d) The starting compound is recovered.
- 259. Which of the following carboxylic acids can be decarboxylated readily by heating them to 100–150°C?
 - (a) CH₃COCH₂COOH
- (b) CCl₃COOH
- (c) CH₂=CH—CH₂COOH
 - (d) CH₃CH₂COOH
- **260.** The intermediates formed during the reaction

$$C_6H_5CH_2CO_2Ag + Br_2 \xrightarrow{\qquad dry \ CCl_4} C_6H_5CH_2Br$$

are

(a)
$$C_6H_5CH_2C$$
 (b) $C_6H_5-CH_2-C$ O

(c) C₆H₅CH₂

- (d) Br°
- **261.** The products obtained in the reaction

$$CH_{3}CH_{2}CO_{2}H \xrightarrow{\quad Cl_{2^{\prime}}\ h\nu \quad }$$

- (a) CH₃CHCO₂H
- (b) CICH2CH2CO2H
- CH_3 CH_3 CH_3 CH_3 CH_3
- (d) Cl₂CHCH₂COOH
- 262. Phosphorus pentachloride reacts with
 - (a) alcohols

(b) ketones

(c) ethers

- (d) amines
- **263.** Which of the following reactions are used in the preparation of alcohol?
 - (a) $C_2H_5Br + aq. KOH \longrightarrow$
 - (b) $(CH_3)_2C=O \xrightarrow{LiAlH_4}$

(c)
$$CH_3$$
— C — OCH_3 — $Na/EtOH$

- (d) $CH_3CH_2Cl \xrightarrow{H_2O}$
- **264.** Which of the following statements are not correct?
 - (a) All alcohols are soluble in water.
 - (b) Only the lower alcohols are soluble in water.
 - (c) All alcohols are poisonous.
 - (d) Methanol is not poisonous.
- **265.** The reaction $2CH_3CH_2OH \xrightarrow{H^+} CH_3CH_2OCH_2CH_3$ is believed to occur through the formation of

- (a) $CH_3CH_2\overset{+}{O}H_2$ (b) $CH_3\overset{+}{C}H_2$ (c) $CH_3CH_2\overset{+}{O}-CH_2CH_3$ (d) none of these
- **266.** Which of the following are correct?
 - (a) Ordinary ethyl alcohol is known as rectified spirit.
 - (b) The alcohol sold in the market for polishing, etc., is known as methylated spirit.
 - (c) Absolute alcohol is 100% ethanol.
 - (d) Power alcohol is 100% ethanol.
- **267.** Which of the following compounds are easily oxidized by K₂Cr₂O₇ and H_2SO_4 ?
 - (a) CH₃CH₂OH

(b) (CH₃)₂CHOH

(c) CH₃CHO

- (d) (CH₂)₂COH
- **268.** Which of the following exhibit hydrogen bonding?
 - (a) Chloroform

(b) Ethyl alcohol

(c) Acetic acid

- (d) Dimethyl ether
- **269.** Which of the following groups will increase the acidity of phenol?
 - (a) $-NO_2$

(b) —CN

(c) —X (halogen)

- (d) None of these
- **270.** Which of the following groups will increase the basicity of phenol?
 - (a) $-NH_2$

(b) —CH₃

(c) —NO₂

(d) None of these

271. Which of the following aldehydes undergo a Cannizzaro reaction?

	(a)	НСНО	(b) C_6H_5CHO
	(c)	CH ₃ CHO	(d) CH ₃ CH ₂ CHO
272.	Which	h of the following aldehydes	andergo aldol condensation?
		CH₃CHO	(b) C ₆ H ₅ CHO
	(c)	C ₆ H ₅ CH ₂ CHO	(d) p-ClC ₆ H ₄ CHO
273.	Which	h of the following are Canniz	zaro reactions?
	(a)	CH ₃ CHO + CH ₃ CHO ——	\longrightarrow CH ₃ CH ₂ OH + HCOOH
	(b)	$2Cl_3CCHO \longrightarrow Cl_3CCH_2$	OH + Cl ₃ CCOOH
	(c)	$C_6H_5CHO + HCHO \longrightarrow 0$	C ₆ H ₅ CH ₂ OH + HCOOH
	(d)	$C_6H_5CHO \longrightarrow C_6H_5CH_2CH_2CH_3CH_3CH_3CH_3CH_3CH_3CH_3CH_3CH_3CH_3$	$OH + C_6H_5COOH$
274.		h of the following are aldol co	
		CH ₃ CHO + CH ₃ CHO — OH	
	(b)	CH ₃ CHO + CH ₃ COCH ₃	$\xrightarrow{OH^-}$ CH ₃ CHOHCOCH ₃
	(c)	$HCHO + HCHO \xrightarrow{OH} OH$	CH₃OH + HCOONa
	(d)	$C_6H_5CHO + C_6H_5CHO - C_6H_5CHO$	$C_6H_5CH_2OH + C_6H_5COONa$
275.	Which	h of the following do not reac	t with Fehling's solution?
		Benzaldehyde	(b) Acetaldehyde
		Glucose	(d) Acetophenone
276.		h of the following do ensation?	not undergo base-catalysed aldo
		Benzaldehyde	(b) 2,2-Dimethylpropionaldehyde
	(c)	2-Methylpropionaldehyde	(d) <i>p</i> -Methylbenzaldehyde
277.	Which	h of the following statements	are correct?
	(a)	When phenol vapour is produced.	passed over Zn dust, benzene is
	(b)	The phenolic—OH group is	ortho- and para-directing.
	(c)	o-Nitrophenol has a lower b	oiling point than p -nitrophenol.
	(d)	Phenol is more acidic than o	-cresol.
278.		h of the following statements	
		Benzaldehyde reduces Fehli	ŭ
	(b)	$C_6H_5CHO + C_6H_5CHO - N$	$\xrightarrow{\text{IaOH}} C_6 H_5 \text{CH} = \text{CHC}_6 H_5 + O_2$
		is a Claisen-Schmidt reactio	n.

- (c) pK_a (formic acid) is less than pK_a (acetic acid).
- (d) o-Toluidine is more basic than aniline.
- **279.** Which of the following statements about ethers are correct?
 - (a) Peroxide is obtained in the presence of air.
 - (b) Ethers are weakly acidic.
 - (c) Ethers form oxonium salts.
 - (d) Ethers form stable complexes with Lewis acids.
- **280.** Aldehydes can be reduced to hydrocarbons by
 - (a) the Clemmensen reduction
 - (b) the Wolff-Kishner reduction
 - (c) Mg/Hg, H₂O
 - (d) the Huang-Minlon method
- **281.** Which of the following statements are correct about the C=O bond?
 - (a) It is made up of one π -bond and one σ -bond.
 - (b) It uses the sp²-hybrid orbital of carbon for its formation.
 - (c) It is planar in nature.
 - (d) It undergoes addition reactions.
- 282. CH_3 —CHO — $\overset{\mbox{o}}{\longrightarrow}$ $CH_3CH(OH)CH_2CHO$

In the aldol condensation of acetaldehyde represented above, which of the following intermediate species are obtained?

O
$$\parallel$$
 (b) $: \overset{\odot}{\operatorname{CH}}_2$ —C—H

(d)
$$CH_3$$
— C — O — CH = CH_2

- 283. Starting with CH₂=C=O, which of the following compounds can be obtained?
 - (a) CH₃COOH O \parallel C CH_3 C $COCH_2CH_3$ C CH_3CONH_2
- 284. In the context of carboxylic acid (R—COOH), which of the following statements are correct?

- (a) Hydrogen bonding is responsible for the high water solubility of simple aliphatic acids (C_1 to C_4).
- (b) Carboxylic acids ionize in aqueous solutions by transferring protons to the solvent mlecules.
- (c) Solubility decreases as chain length (R) increases.
- (d) Solubility decreases with more branching in the chain (R).
- 285. In the esterification of propanoic acid with methanol in the presence of a mineral acid, which of the following are intermediate species?

- 286. Which of the following methods are used for the conversion of carboxylic
 - (a) $RCOOH + SOCl_2 \longrightarrow$ (b) $RCOOH + PCl_5 \longrightarrow$ (c) $RCOOH + Cl_2 \longrightarrow$ (d) $RCOOH + P + Cl_2 \longrightarrow$
- **287.** In which of the following esters is the α -hydrogen acidic?
 - (a) CH₃COOC₂H₅
- (b) O₂NCH₂COOC₂H₅
- (c) NCCH₂COOC₂H₅
- (d) CH₃COCH₂COOC₂H₅
- 288. Which of the following intermediate species is/are formed in the reaction of acrylic acid with HBr to give β-bromopropionic acid

f acrylic acid with HBr to give
$$\beta$$
-bromopropionic acid

$$(CH_2 = CH - COOH \xrightarrow{HBr} BrCH_2CH_2COOH)?$$
(a)
$$CH_2 = CH - C$$

$$OH$$

$$CH_2 = CH - C$$

$$OH$$
(b)
$$CH_2 = CH - C$$

$$OH$$

$$OH$$

$$CH_2 = CH - C$$

$$OH$$

$$OH$$

289. Which of the following compounds exhibit optical activity?

(a)
$$H_2C=C=CH_2$$
 (b) $H_3C=C=CC$ CH_3 CH_3

(c)
$$COOH$$
 NO_2 (d) $COOH$ NO_2 NO_2

- **290.** Which of the following will decolourize a KMnO₄ solution?
 - (a) CH₂CH₂OH

(b) CH₃CH=CH₂

(c) (CH₃)₂CHOH

- (d) CH₃COCH₃
- **291.** Which of the following are protic solvents?
 - (a) Water

- (b) Ethanol
- (c) Dimethylformamide
- (d) Dimethylsulphoxide
- 292. The intermediate stages in the conversion

$$(CH_3)_2$$
— C — $(CH_3)_2$ — $\xrightarrow{\text{dil. H}_2SO_4}$ $CH_3COC(CH_3)_3$ are OH OH

(a)
$$(CH_3)_2$$
— C — $C(CH_3)_2$ (b) $(CH_3)_2$ — C — $C(CH_3)_2$ OH OH₂ OH

(c)
$$CH_3 - \overset{\circ}{C} - C(CH_3)_3$$

OH

(c)
$$CH_3$$
— C — $C(CH_3)_3$ (d) CH_3 — C — $C(CH_3)_3$ OH OH

- 293. Which of the following compounds will give a red precipitate on being heated with Fehling's solution?
 - (a) C₆H₅CHO

(b) CH₃CHO

(c) CH₃COCH₃

- (d) C₆H₅CH₂CHO
- 294. What types of isomerism are exhibited by hexanoic acid?
 - (a) Chain isomerism
- (b) Position isomerism
- (c) Functional group isomerism (d) Metamerism
- 295. Which of the following statements are correct?
 - (a) Carboxylic acids have higher boiling points than those of alcohols of similar molecular weight.
 - (b) Carboxylic acids have lower boiling points than those of alcohols of similar molecular weight.
 - (c) Carboxylic acids (C_1 to C_4) are soluble in water.
 - (d) The melting points of carboxylic acids increase or decrease in an irregular manner.

- **296.** Which of the following statements are correct?
 - (a) The two carbon-oxygen bond lengths in formic acid are different.
 - (b) The two carbon-oxygen bond lengths in sodium formate are equal.
 - (c) The carbon-oxygen bond length in formic acid is less than that in sodium formate.
 - (d) The carbon-oxygen bond length in formic acid is greater than that in sodium formate.
- **297.** In the context of the rearrangement of an oxime of a ketone to an amide (represented below)

$$\begin{array}{ccc}
R - C - R' \xrightarrow{PCL_5} O = C - R' \\
\parallel & \parallel & \parallel \\
N - OH & NHR
\end{array}$$

which of the following statements are correct?

- (a) It is the *trans*-hydrocarbon radical (R) with respect to the OH group that migrates.
- (b) The group that migrates never gets completely detached from the remainder of the molecule during the transformation.
- (c) The rearrangement is intermolecular.
- (d) None of these
- 298. Nitration of phenol with dilute nitric acid at 293 K gives
 - (a) o-nitrophenol

- (b) *p*-nitrophenol
- (c) *m*-nitrophenol

- (d) 2,4,6-nitrophenol
- **299.** An alcohol, on treatment with $P+I_2$ followed by the reaction of the formed product first with $AgNO_2$ and then with HNO_2 and final basicification, gives a blue colour. Which of the following alcohols can it be?
 - (a) CH₃CH₂OH

(b) (CH₃)₂CHOH

(c) (CH₃)₃C—OH

- (d) CH_3 CHOH
- **300.** Which of the following alcohols, on treatment with concentrated HCl and anhydrous ZnCl₂, will become turbid instantaneously?
 - (a) CH₃OH

(b) (CH₃)₂CHOH

(c) (CH₃)₃COH

301. Which of the following compounds will react positively to the iodoform

	(a)	(CH ₃) ₂ CHOH	(b)	CH ₃ C	CH ₂ OH	
	(c)	CH ₃ CH ₂ CH ₂ OH	(d)	CH ₃ C	CHOHCH ₂ CH ₃	
302.	The re	eaction of glycerol with HIO ₄	gives			
		formaldehyde	_	form	ic acid	
	(c)	iodic acid	(d)	oxali	c acid	
303.	Glyce	rol, on being heated with oxa	ılic aci	d at 38	33 K, gives	
		glyceryl monooxalate			, 0	
		glyceryl monoformate				
	(c)	allyl alcohol				
	(d)	formic acid				
304.	Whicl	h of the following will result:	in the	forma	tion of an ether?	
		$(CH_3)_3CONa + CH_3CH_2Br -$				
	(b)	$(CH_3)_3CBr + C_2H_5ONa$	\rightarrow			
	(c)	$C_6H_5ONa + CH_3Br \longrightarrow$				
	(d)	$C_6H_5Br + CH_3ONa \xrightarrow{r}$				
305.	Methy	ylethyl ketone can be reduced	d to <i>n-</i> l	butane	e by	
	(a)	the Meerwein-Ponndorf red	duction	n		
	(b)	the Wolff-Kishner reduction	n			
	(c)	$Mg-Hg$, H_2O				
	(d)	HI/red phosphorus at 423 I	<			
306.	On be	eing treated with HCl, aceton	e gives	5		
	(a)	mesityl oxide	(b)	phor	one	
	(c)	mesitylene	(d)	aldol		
307.	Whicl	h of the following statements	are co	rrect f	or benzoic acid?	
	(a)	Nitration gives o- and p-nitr	obenz	oic aci	d.	
	(b)	Bromination (Br ₂ /FeBr ₃) given	es m-l	oromo	benzoic acid.	
	(c)	The Friedel–Crafts reac <i>m</i> -carboxyacetophenone.	tion	with	CH ₃ COCl/AlCl ₃	gives
	(d)	The reaction with co 3-carboxybenzenesulphonic		ated	sulphuric acid	gives

Answers

1. c	2. b	3. a	4. b	5. d
6. a	7. b	8. b	9. a	10. c
11. b	12. a	13. c	14. a	15. c
16. b	17. d	18. a	19. c	20. c
21. c	22. d	23. a	24. b	25. c
26. b	27. a	28. c	29. d	30. d
31. b	32. c	33. a	34. c	35. a
36. c	37. c	38. b	39. b	40. a
41. c	42. a	43. a	44. c	45. b
46. a	47. c	48. a	49. c	50. a
51. a	52. c	53. c	54. d	55. c
56. d	57. a	58. a	59. b	60. a
61. a	62. c	63. a	64. a	65. a
66. a	67. a	68. a	69. c	70. d
71. c	72. d	73. b	74. b	75. a
76. a	77. a	78. d	79. c	80. d
81. c	82. a	83. d	84. a	85. b
86. b	87. b	88. b	89. c	90. d
91. c	92. c	93. a	94. b	95. d
96. b	97. c	98. c	99. c	100. a
101. b	102. a	103. a	104. b	105. c
106. b	107. a	108. c	109. a	110. a
111. b	112. d	113. a	114. c	115. a
116. a	117. a	118. a	119. a	120. a
121. b	122. b	123. a	124. b	125. d
126. b	127. b	128. a	129. a	130. c
131. d	132. b	133. d	134. a	135. b
136. c	137. d	138. b	139. b	140. d
141. b	142. b	143. b	144. c	145. c
146. c	147. c	148. a	149. a	150. a
151. a	152. b	153. c	154. c	155. a
156. d	157. b	158. b	159. d	160. d
161. a	162. c	163. b	164. c	165. d
166. c	167. b	168. a	169. a	170. b
171. c	172. a	173. d	174. a	175. a
176. c	177. b	178. a	179. a	180. c
181. a	182. b	183. a	184. a	185. c
186. a	187. c	188. a	189. a	190. c

191. d	192. a	193. a	194. a	195. a
196. c	197. a	198. c	199. d	200. a
201. d	202. a	203. c	204. b	205. b
206. a	207. a	208. a	209. a	210. b
211. c	212. c	213. c	214. a	215. b
216. c	217. a	218. a	219. b	220. c
221. c	222. b	223. b	224. b	225. c
226. b	227. b	228. a	229. c	230. d
231. c	232. c	233. d	234. d	235. a
236. d	237. a, b, c	238. a, c	239. a, b, c	240. a, b, c
241. b, c, d	242. a, b, c	243. a, b, c	244. a, b, c, d	245. a, b, d
246. a, b, d	247. b, d	248. a, b	249. c, d	250. a, b, c
251. a, b, c, d	252. a, c	253. a, b, c	254. a, b, c, d	255. a, c
256. a, b	257. c, d	258. b, c	259. a, b	260. a, b, c, d
261. a, b	262. a, b, c	263. a, b, c	264. a, c	265. a, b, c
266. a, b, c	267. a, b, c	268. b, c, d	269. a, b, c	270. a, b
271. a, b	272. a, c	273. b, c, d	274. a, b	275. a, d
276. a, b, d	277. a, b, c, d	278. c, d	279. a, c, d	280. a, b, d
281. a, b, c, d	282. a, b, c	283. a, b, c	284. a, b, c, d	285. a, b, c, d
286. a, b, d	287. a, b, c, d	288. a, b, c, d	289. b, c	290. a, b, c
291. a, b	292. a, b, c, d	293. b, d	294. a, b	295. a, c, d
296. a, b, c	297. a, b	298. a, b	299. b, d	300. c, d
301. a, b, d	302. a, b, c	303. a, b	304. a, c	305. b, d
306. a, b	307. b, d			

Hints to More Difficult Problems

- **35.** The more stable a carbonyl compound is, the more easily does it hydrogenate catalytically.
- **42.** The acid strengths of phenols and carboxylic acids depend on their pK_a values.
- **44.** Unlike other aliphatic aldehydes, formaldehyde reacts with ammonia to form hexamethylenetetramine, a cyclic compound. Upon nitration, this gives a trinitro-derivative—an explosive cyclonite also called RDX.
- **45.** On reaction with HCN in the presence of an alkali, α , β -unsaturated ketones do not give the expected cyanohydrins. In this case, 1,4-addition results in the formation of a β -cyanoketone.
- **46.** The addition of HCl to α , β -unsaturated aldehydes and ketones moves the halogen to the β -carbon. However, if there is a vinyl group in

conjugation with the carbonyl group, the position is opposite to that expected from Markovnikov's rule.

- 47. Simple aldehydes undergo normal Cannizzaro reactions but compounds containing two aldehyde groups undergo internal Cannizzaro reactions, i.e., one CHO is oxidized while the other CHO is reduced.
- **48.** On treatment of carboxylic acids with Cl_2/P , Cl gets attached to the α -carbon since it is more reactive.
- **49.** The first step is the formation of

Being a γ -hydroxyacid, it undergoes lactonization to give the product shown in option (c).

- **247.** The reaction of a ketone and a Grignard reagent does not yield the usual products in the following cases.
 - (a) The branching of the carbon chain near the carbonyl group prevents nucleophilic addition by the Grignard reagent due to steric hindrance.
 - (b) If the Grignard reagent has a bulky alkyl or aryl group, it fails to attack the electrophilic centre of the substrate molecule.
- **248.** Products are obtained from α , β -unsaturated aldehydes and ketones respectively through the initial 1,4- and 1,2-additions.
- **253.** On treatment with a solution of CrO₃ in dilute H₂SO₄, primary and secondary alcohols become greenish and opaque within 2–3 seconds.
- **258.** The gem dihalide obtained initially from the carbonyl compound undergoes an E2-type elimination to give allenes and acetylenes.
- **259.** Thermal decarboxylation occurs most readily when the α-carbon atom attached to the —COOH group carries a strongly electron-withdrawing group (i.e., –I substituent).
- **260.** The decarboxylation of the silver salts of carboxylic acids in the presence of bromine or chlorine is called a Hunsdiecker reaction and is useful for the synthesis of alkyl halides.
- 261. In the presence of UV light, the reaction of chlorine with CH₃CH₂CO₂H involves the free-radical chlorination at all positions along the chain. This is also true of all hydrocarbons.

Compounds Containing Nitrogen and Sulphur

• *Type 1* •

Choose the correct option. Only one option is correct.

1. Which of the following is an enamine?

- 2. What type of amine is tertiary butylamine?
 - (a) Primary
- (b) Secondary (c) Tertiary
- (d) Quaternary
- 3. Which reagent is used to distinguish between $(C_2H_5)_2NH$ and $(C_2H_5)_3N$?
 - (a) NaOH
- (b) Dilute HCl (c) C₆H₅SO₂Cl (d) KMnO₄
- 4. A molecule of which of the following will exhibit hydrogen bonding with another of the same compound?
 - (a) $CH_3 N CH_3$ (b) CH₂NH₂
 - (c) CF₃CH₃

- (d) CH₃-O-CH₃
- 5. N,N-Dimethylbenzamide cannot be made by

(a)
$$O$$
 $||$ $C_6H_5COC_2H_5 + (CH_3)_2NH \longrightarrow$

(b) O
$$\parallel$$
 $C_6H_5CCl + (CH_3)_2NH \longrightarrow$

(d) O II
$$C_6H_5CNH_2 + CH_3MgCI \longrightarrow$$

6. The final product (III) obtained in the reaction sequence

Br
$$\frac{1. \text{HONO}}{2. \text{CuCl}}$$
 I $\frac{\text{Sn/HCl}}{2. \text{H}_3 \text{PO}_2}$ III $\frac{1. \text{HONO}}{2. \text{H}_3 \text{PO}_2}$ III

is

NO₂

(d) none of these

- 7. Among the following, which is the strongest base?
 - (a) \sim NH₂

(b)
$$H_3C$$
 \longrightarrow NH_2

(c) \sim CH₂NH₂

8. Which of the following reactions can be used to synthesise *m*-bromoaniline?

(a)
$$C_6H_6 \xrightarrow{\qquad 1. \text{ HNO}_3/\text{H}_2\text{SO}_4 \qquad} \left[\qquad \right] \xrightarrow{\qquad \text{Br}_2/\text{H}_2\text{O}}$$

$$(b) \quad C_6H_6 \xrightarrow{\hspace{1cm} 1. \hspace{1cm} Br_2/FeBr_3 \hspace{1cm}} \left[\hspace{1cm} \right] \xrightarrow{\hspace{1cm} H_2 \hspace{1cm}} Pt \\$$

(c)
$$C_6H_5COOH \xrightarrow{Br} \begin{bmatrix} & 1. SOCl_2 \\ \hline & 2. NH_3 \end{bmatrix} \begin{bmatrix} & Br_2 \\ \hline & NaOH \end{bmatrix}$$

- (d) None of these
- 9. Which of the following reagents will be useful as the basis for a simple chemical test to distinguish between

$$H_3C$$
 \longrightarrow NH_2 and \bigcirc \longrightarrow CH_2NH_2 ?

- (a) $C_6H_5SO_2Cl$ and OH^- in H_2O (b) HONO, then β -naphthol
- (c) Dilute HCl

- (d) AgNO₃ in H₂O
- **10.** 3,5-Dibromotoluene can be best synthesised by
 - (a) Toluene $\xrightarrow{\text{Br}_2/\text{Fe/heat}}$

(b)
$$p ext{-Toluidine} \xrightarrow{\operatorname{Br}_2, \operatorname{H}_2\operatorname{O}} \left[\begin{array}{c} 1.\operatorname{HNO}_2 \\ \hline 2.\operatorname{H}_3\operatorname{PO}_2 \end{array} \right]$$

(c) Toluene
$$\xrightarrow{\text{fuming HNO}_3}$$
 $\xrightarrow{\text{fuming H}_2\text{SO}_4}$ $\xrightarrow{\text{In HCNO}}$ $\xrightarrow{\text{In HCNO}}$ $\xrightarrow{\text{In HCNO}}$ $\xrightarrow{\text{In HCNO}}$

- (d) 1,3-Dibromotoluene \longrightarrow CH₃Cl/AlCl₃/heat \longrightarrow
- 11. The final product (IV) obtained in the reaction sequence

$$Toluene \xrightarrow{\begin{array}{c} 1. \ KMnO_4/OH^- \\ \hline 2. \ H^+ \end{array}} I \xrightarrow{\begin{array}{c} SOCl_2 \\ \hline \end{array}} II \xrightarrow{\begin{array}{c} NH_3 \\ \hline \end{array}} III \xrightarrow{\begin{array}{c} OBr^- \\ \hline \end{array}} IV$$

is

- (a) $C_6H_5CONH_2$
- (b) p-CH₃C₆H₄NO₂

(c) $C_6H_5CH_2NH_2$

- (d) $C_6H_5NH_2$
- 12. Among the following, which is the strongest base?

(a)
$$CH_3$$
 (b) H CH_3 — $C-NH_2$ CH_3NCH_3 CH_3

- 13. The best method to synthesise m-dibromobenzene is by using the reaction
 - $Benzene \xrightarrow{Br_2/FeBr_3/heat}$
 - (b) Aniline $\xrightarrow{\text{Br}_2, \text{H}_2\text{O}} \qquad \boxed{\begin{array}{c} 1. \text{ HONO} \\ \hline 2. \text{CuBr} \end{array}}$
 - (c) Nitrobenzene $\xrightarrow{\text{fuming HNO}_3} \left[\begin{array}{c} Fe/HCl \\ \hline C_3H_5OH, \text{ heat} \end{array} \right] \xrightarrow{\text{1. HONO}} 2. \text{ CuBr}$
 - (d) Bromobenzene $\xrightarrow{\text{HNO}_3}$ $\left[\begin{array}{c} \text{Fe/HCl} \\ \text{C.H.OH. heat} \end{array}\right] \xrightarrow{\text{1. HONO}} 2. \text{CuBr}$
- 14. *m*-Fluoronitrobenzene is best synthesised by using the reaction
 - (a) Nitrobenzene $\xrightarrow{\text{fuming HNO}_3}$ $\left[\begin{array}{c} \text{NH}_3/\text{H}_2\text{S} \\ \hline \end{array} \right] \xrightarrow{\text{NH}_3/\text{H}_2\text{S}} \left[\begin{array}{c} 1. \text{ HONO} \\ \hline 2. \text{ HRE} \end{array} \right]$
 - (b) Aniline $\xrightarrow{F_2}$ heat
 - (c) Fluorobenzene $\xrightarrow{\text{HNO}_3}$ HSO. heat
 - (d) $m\text{-}C_6H_4(NH_2)_2 \xrightarrow{1. \text{ HONO}} \frac{1. \text{ HONO}}{2. \text{ CuNO}_5.3. \text{ HBF}}$
- 15. The final product obtained in the reaction

$$C_6H_5NH_2 + HONO \xrightarrow{1. HCl(0-5^{\circ}C)} 2. heat$$

is

- (a) C_6H_5Cl (b) C_6H_5OH (c) C_6H_6
- (d) none of these
- **16.** The major product obtained in the reaction

is

(a)
$$CH_3$$
 C_6H_5 OH

(b) C_6H_5OH

(c) $C_6H_5NO_2$

(d)

$$C_6H_5-N=N$$

- 17. Which of the following is a thioether?

- (c) CH₃CH₂SCH₂CH₃
- (d) None of these
- 18. The transformation $C_6H_5CH_2CH_2Br \longrightarrow C_6H_5CH_2CH_2SH$ can be effected using
 - (a) H_2S

(b) NH₂CNH₂

(c) - + CH₃SCH₂Na

(d) NaSH

19. The transformation

$$CH_3CH_2I \longrightarrow CH_3CH_2SCH_2CH_3$$

can be effected by

(a) K_2S , Δ

(c) H₂S

- (b) P_2S_5/Δ (d) NH_2CNH_2
- **20.** Which of the following statements is correct?
 - (a) Methylamine is more basic than ammonia.
 - (b) Dimethyl amine is less basic than methylamine.
 - (c) Dimethylamine is less basic than trimethylamine
 - (d) Ammonia is more basic than methylamine
- 21. Which of the following compounds is the most reactive to a nucleophilic attack?









- 22. In the reaction of (S) 2-phenylpropamide with NaOBr/H₂O to give 1-phenylethylamine,
 - (a) there is retention of configuration
 - (b) there is inversion of configuration
 - (c) a mixture of two products is obtained
 - (d) there is no reaction
- **23.** The oxidation of *m*-dinitrobenzene with alkaline potassium ferricyanide gives

(d) There is no reaction.

- **24.** Nitrobenzene can be converted into phenylhydroxylamine by reduction with
 - (a) Zn-NH₄Cl-H₂O
- (b) alkaline sodium arsenite
- (c) alkaline sodium stannite
- (d) Zn-aqueous NaOH
- 25. In pyridine, the state of hybridization of the nitrogen atom is
 - (a) sp^2

(b) sp^3

(c) sp

(d) none of these

26. The final product (III) obtained in the reaction

nal product (III) obtained in the reaction
$$CH_3CH_2NH_2 \xrightarrow{CH_3COCl} I \xrightarrow{HONO} II \xrightarrow{\Delta} III$$

is

- (a)
- (d) none of these
- 27. The product obtained in the reaction

$$C_6H_5N=C$$
: + CH_3CCH_3 + CH_3CO_2H \longrightarrow

is

- (a) $C_6H_5NH_2$
- (b) C₆H₅COOH
- (c) $C_6H_5NHC_6H_5$

28. The product obtained in the reaction

$$\begin{array}{c} CH_3 \\ | \\ HC-NO_2 \\ | \\ CH_3 \end{array} + CH_2 = CH-C \equiv N \xrightarrow{\bar{O}H} \xrightarrow{\bar{O}H}$$

is

(a)
$$N \equiv C - CH_2CH_2 - \begin{matrix} CH_3 \\ - C - NO_2 \\ CH_3 \end{matrix}$$

(b)
$$\begin{array}{c} CH_3 \\ N \equiv C - CH_2 - CH_2 - C - NH_2 \\ CH_3 \end{array}$$

(c)
$$\begin{array}{c} CH_3 \\ N \equiv C - CH_2CH_2 - C - NO_2 \\ CH_2CH_2CH_2C \equiv N \end{array}$$

(d)
$$\begin{array}{c} CH_2CH_2CH_2C \equiv N \\ | \\ N \equiv C - CH_2 - CH_2 - C - NO_2 \\ | \\ CH_2CH_2CH_2C \equiv N \end{array}$$

- 29. Which of the following species is present in a solution of glycine (H₂NCH₂COOH)?
 - (a) H_3 NC H_2 CO₂H
- (b) $\stackrel{+}{N}H_3CH_2CO_2$

(c) $NH_2CH_2CO_2$

- (d) All of these
- **30.** The oxidation of aniline with peracetic acid in the presence of acetic acid by refluxing gives
 - (a) NHOH
- (b) NO₂

(c) 0

(d) none of these

31. The final product of the reaction of ethyl bromide with H_2S in the presence of KOH,

$$CH_3CH_2Br + KOH + H_2S \xrightarrow{C_2H_5OH}$$
 heat

is

(a) CH₃CH₂SH

- (b) CH₃CH₂SCH₂CH₃
- (c) CH₃CH₂S—S—CH₂CH₃
- (d) none of these
- 32. Which of the following amines is chiral?
 - (a) CH₃CH₂CH₂NH₂
- (b) CH₃CH₂NH—CH₃
- CH₃
 CH₃
 CH₃
 CH₃
 CH₃
- $C_{2}H_{5}$ CH_{3} CH_{3} $C_{3}H_{7}$

- 33. In an aqueous solution, the order of basicity of the amines $(CH_3)_3N$, $(CH_3)_2NH$, CH_3NH_2 and NH_3 is
 - (a) $(CH_3)_3N > (CH_3)_2NH > CH_3NH_2 > NH_3$
 - (b) $(CH_3)_2NH > CH_3NH_2 > CH_3N > NH_3$
 - (c) $NH_3 > CH_3NH_2 > (CH_3)_2NH > (CH_3)_3N$
 - (d) All these amines are equally basic.
- **34.** The order of basicity of the amines $(CH_3)_3N$, $(CH_3)_2NH$, CH_3NH_2 and NH_3 in the gaseous phase is
 - (a) $(CH_3)_3N > (CH_3)_2NH > CH_3NH_2 > NH_3$
 - (b) $NH_3 > CH_3NH_2 > (CH_3)_2NH > (CH_3)_3N$
 - (c) $(CH_3)_2NH > CH_3NH_2 > (CH_3)_3N > NH_3$
 - (d) All these amines are equally basic.
- 35. The product obtained in the reduction

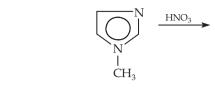
$$NO_2$$
 NO_2
 H_2S, NH_3
 NO_2

is

(a)
$$CH_3$$
 NH_2 NH_2

(d) The compound is not reduced.

36. The product obtained in the reaction



(a)
$$O_2N$$
 N
 I
 CH_3

(b)
$$N$$
 NO_2 NO_2 NO_3

(c)
$$O_2N$$
 N N N CH_3

(d) There is no reaction.

- 37. The reaction of dimethylsulphoxide with α -bromoacetophenone ($C_6H_5COCH_2Br$) gives
 - (a) C₆H₅COCH₂OH
- (b) C₆H₅COCHO
- (c) C₆H₅COCOOH
- (d) No reaction takes place.
- 38. In the conversion $C_2H_5Br \longrightarrow C_2H_5CN$, the reagent used is
 - (a) alcoholic KCN
- (b) alcoholic AgCN

(c) NH₃

- (d) none of these
- 39. The reduction of methyl cyanide with sodium and alcohol gives
 - (a) methylamine

(b) ethylamine

(c) acetic acid

- (d) methyl alcohol
- 40. An aliphatic organic compound containing C, H and N reacts with dilute HCl to produce formic acid. It is reduced to dimethylamine by Pt or Ni, and undergoes an addition reaction with chlorine and sulphur. The compound can be
 - (a) CH₃NC

(b) CH₃CN

(c) CH₃NH₂

- (d) a mixture of (a) and (b)
- 41. Ethyl isocyanide is prepared by the reaction between
 - (a) C_2H_5Br and KCN
- (b) C₂H₅Br and AgCN
- (c) C_2H_5Br and HCN
- (d) C₂H₅Br and NH₃

(a) aniline

(c) *p*-aminophenol

42.	In the		and	alcoholic AgNO ₂ , the product
	(a)	nitroethane	(b)	ethane
	(c)	ethyl nitrite	(d)	ethyl isocyanide
43.	In the	reaction between CH ₃ NC and	HgC), the product obtained is
	(a)	methyl isothiocyanate	(b)	methyl isocyanate
	(c)	methylamine	(d)	methyl cyanide
44.	Which produ		ll pro	oduce methyl nitrite as the major
	(a)	$CH_3I + AgNO_2 \longrightarrow$	(b)	$CH_3I + NaNO_2 \longrightarrow$
	(c)	both (a) and (b)	(d)	neither
45.	tempe	ganic amino compound reacterature to produce an oily nitro CH ₃ NH ₂	osoar	th aqueous nitrous acid at low nine. The compound is $CH_3CH_2NH_2$
	(c)	$(C_2H_5)_2NH$	(d)	$(C_2H_5)_3N$
46.		nitration of benzene with co , the electrophile is	ncer	ntrated HNO ₃ and concentrated
	(a)	NO_2^+	(b)	NO_2^-
	(c)	NO ₂	(d)	nitric oxide
47.	A nitroalkane reacts with nitrous acid to yield a product which is insoluble in an alkali, and turns blue on treatment with an alkali. The nitroalkane could be			
	(a)	CH ₃ CH ₂ NO ₂	(b)	(CH ₃) ₂ CHNO ₂
	(c)	(CH ₃) ₃ CNO ₂	(d)	CH ₃ CH—CH ₂ NO ₂
				CH ₃ CH—CH ₂ NO ₂ CH ₃
48.		roalkane produces a ketone lkane could be	whe	en it is boiled with HCl. The
	(a)	CH ₃ CH ₂ NO ₂	(b)	$(CH_3)_2CHNO_2$
				CH ₂
	(-)	(CII.) CNO	(4)	CII C CII NO
	(c)	(CH ₃) ₃ CNO ₂	(a)	CH ₃ CH ₃ —C—CH ₂ NO ₂ CH ₃
49.	The el		enze	ne in a strongly acidic medium

(b) azoxybenzene

(d) azobenzene

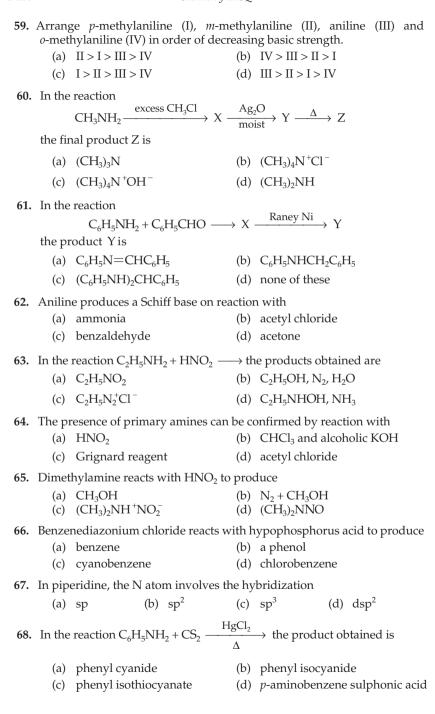
50.	The el		enze	ne in a weakly acidic medium		
	•	aniline	(b)	phenylhydroxylamine		
	(c)	<i>p</i> -aminophenol	(d)	azoxybenzene		
51.	Which	of the following will give only	y one	e monosubstituted product?		
	(a)	o-Dinitrobenzene	(b)	<i>m</i> -Dinitrobenzene		
	(c)	<i>p</i> -Dinitrobenzene	(d)	None of these		
52.	The ox	kidation of aniline with K ₂ Cr ₂ C) ₇ /H	₂ SO ₄ produces		
	(a)	benzoic acid	(b)	benzene		
	(c)	<i>p</i> -benzoquinone	(d)	<i>p</i> -nitrophenol		
53.	How ring?	many isomeric amines with th	e for	mula C ₇ H ₉ N contain a benzene		
	_	Two	(b)	Three		
	(c)	Four	(d)	Five		
54.	How 1	nany isomeric amines can hav	e the	formula C ₄ H ₁₁ N?		
	(a)	Five	(b)	Six		
	(c)	Seven	(d)	Eight		
55.	Which among the following has the highest boiling point?					
	(a)	CH ₃ CH ₂ CH ₂ NH ₂	(b)	CH ₃ CH ₂ —NH		
	$\stackrel{I}{CH_3}$					
		CH ₃ CH ₃ —N—CH ₃				
	(c)	CH_3 — N — CH_3	(d)	CH ₃ NH ₂		
56.	The N	atom in amines involves				
	(a)	sp ³ hybridization	(b)	sp ² hybridization		
	(c)	sp ² and sp ³ hybridization	(d)	none of these		
57.	Arran	ge the following in order of de	creas	sing basic strength.		
	$(C_2H_5)_2NH$, $C_2H_5NH_2$, $(C_2H_5)_3N$					
	(a)	$(C_2H_5)_2NH > C_2H_5NH_2 > (C_2H_5)_2NH_3 > (C_2H_5)_2NH_2 > (C_2H_5)_2$	$H_5)_3N$	J		
	(b)	$C_2H_5NH_2 > (C_2H_5)_2NH > (C_2H_$	$H_5)_3N$	I		
	(c)	$(C_2H_5)_3N > (C_2H_5)_2NH > C_2H$	₅ NH	2		
	(d)	$(C_2H_5)_3N > C_2H_5NH_2 > (C_2H_5)_3$) ₂ NF	I		
58.			-	d <i>p</i> -methoxyaniline (III) in order		
		reasing basic strength.	,	, , ()		

(b) I > II > III

(d) III > II > I

(a) III > I > II

(c) II > I > III



- **69.** In the reaction $CH_3CH_2NH_2 + CH_3MgBr \longrightarrow X$, the product is
 - (a) CH₃CH₃

- (b) CH₃CH₂CH₃
- (c) CH₃CH₂CH₂CH₃
- (d) CH₄
- 70. Diethylamine reacts with HNO₂ to produce
 - (a) $(C_2H_5)_2 NH^+NO_2^-$
- (b) $(C_2H_5)_2NNO$
- (c) $N_2 + C_2H_5OH$
- (d) C_2H_5OH
- 71. The product obtained in the reaction

$$CH_3CH_2CONH_2 \xrightarrow{\text{(i) bromine water}} X$$

is

- (a) CH₃CH₂COOH
- (b) CH₃CH₂CH₂NH₂

(c) CH₃CH₂NH₂

- (d) CH₃CH₂COONH₄
- **72.** Arrange *p*-toluidine (I), *N*,*N*-dimethyl-*p*-toluidine (II), *p*-nitroaniline (III) and aniline (IV) in order of decreasing basicity.
 - (a) I > IV > III > II

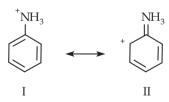
(b) I > II > III > IV

(c) II > I > IV > III

- (d) III > I > II > IV
- **73.** Arrange methylamine (I), dimethylamine (II), aniline (III) and *N*-methylaniline (IV) in order of decreasing basicity.
 - (a) I > II > III > IV

- (b) II > I > IV > III
- (c) III > II > IV

- (d) IV > III > II > I
- 74. Gabriel phthalimide synthesis is used for the preparation of
 - (a) primary aromatic amines
- (b) primary aliphatic amines
- (c) secondary amines
- (d) tertiary amines
- 75. Consider the following structure for the anilium ion.



Which of the following statements are correct regarding this structure?

- (a) It is not an acceptable canonical structure because it is nonaromatic.
- (b) It is not an acceptable canonical structure because here nitrogen has ten valence electrons.
- (c) It is an acceptable canonical structure.
- (d) It is not an acceptable canonical structure because carbonium ions are less stable than ammonium ions.

76.	Arrange phenol (I), <i>p</i> -cresol (II), <i>m</i> -nitrophenol (III) and <i>p</i> -nitropheno (IV) in order of decreasing acidity.				
		II > I > III > IV	(b)	IV > III > I > II	
	(c)	III > IV > I > II	(d)	I > IV > III > II	
77.		najor product of the reaction be NH_3 is	etwe	en p -chlorotoluene and KNH $_2$ in	
	•	o-toluidine	(b)	<i>m</i> -toluidine	
	(c)	<i>p</i> -toluidine	(d)	<i>p</i> -chloroaniline	
78.	Which	n of the following is a Sandmey	er re	eaction?	
	(a)	$C_6H_5N_2Cl \frac{\text{cuprous chloride}}{\Delta}$	→ C,	$_{6}H_{5}Cl+N_{2}$	
	(b)	$2C_6H_5Cl \xrightarrow{CuCl_2/HCl} C_6H_6$	+ Cl	2	
	(c)	$C_6H_5OH \xrightarrow{Zn dust} C_6H_6$	+ Zn	nO	
	(d)	$C_6H_5NO_2 + 6H \xrightarrow{Sn + HCl}$	C_6F	$H_5NH_2 + 2H_2O$	
79.	Which of the following compounds will produce a p -nitroso derivative on treatment with NaNO ₂ /HCl at 0–5°C?				
		Aniline		N-methylaniline	
	` '	o-Methylaniline		<i>N,N</i> -dimethylaniline	
80.		n of the following reactions wil		•	
	(a)	Benzene $\xrightarrow{\text{conc. H}_2\text{SO}_4, 80^\circ\text{C}}$		-	
	(b)	Benzenefuming sulphuric aci	d, 20	0−250°C	
	(c)	Benzenechlorosulphonic acid	$\stackrel{I}{ o}$		
	(d)	Benzene — sulphuryl chloride, p	yrid	ine →	
81.	Nitrol	benzene can be converted into	azox	ybenzene by reduction with	
	(a)	Na ₃ AsO ₃ /NaOH/H ₂ O	(b)	Zn/NaOH/CH ₃ OH	
	(c)	Sn + HCl	(d)	Zn/NH ₄ Cl/H ₂ O	
82.	Benze with	enediazonium chloride can be	conv	verted into phenol by treating it	
	(a)	H ₃ PO _{3,} H ₂ O, CuCl	(b)	H_2O , heat	
	(c)	alcohol, heat	(d)	HBF ₄ and NaNO ₂ /Cu	
83.	Arran	ge CH ₃ NH ₂ (I), CH ₃ CONH ₂ (II) aı	nd NH ₂ CH ₂ COOH (III) in order	

(b) III > II > I

(d) II > III > I

of decreasing basicity.

(a) I > III > II

(c) II > I > III

- 84. The bond angles in methane (I), ammonia (II) and trimethylamine (III) increase in the order
 - (a) I > III > II

(b) I > II > III

(c) II > I > III

- (d) III > II > I
- 85. Arrange the following compounds in order of decreasing basic strength.

 CH_3NH_2 $(CH_3)_2NH$

П

 $(CH_3)_3N$

 NH_3 IIIIV

- (a) I > II > III > IV
- (b) II > I > III > IV
- (c) IV > III > II > I

- (d) II > III > I > IV
- 86. How many structures are possible for a compound having the molecular formula C₃H₇NO₂?
 - (a) One
- (b) Two
- (c) Three (d) Four

• Type 2 •

Choose the correct options. More than one option is correct.

87. Which of the following reaction sequences will give aniline as the final product?

(a) Benzene
$$\xrightarrow{\text{HNO}_3}$$
 $\left[\quad \right] \xrightarrow{\text{1. Sn/HCl}}$ $\xrightarrow{\text{2. NaOH/H}_2O}$

(b)
$$C_6H_5CCI \xrightarrow{NH_3} \begin{bmatrix} OBr \\ (Br_2/OH) \end{bmatrix}$$

(c)
$$C_6H_5CH_2Br \xrightarrow{excess NH_3} \begin{bmatrix} 1. HONO \\ \hline 2. H_3PO_2 \end{bmatrix}$$

- (d) None of these
- 88. Which of the following reagents can be used to distinguish between *p*-O₂NC₆H₄NH₂ and C₆H₅COOH?
 - (a) $KMnO_4$
- (b) Br_2/CCl_4 (c) NaOH
- (d) HCl
- 89. Which of the following methods can be used to prepare *n*-propylamine?

(a)
$$CH_3CH_2CH_2Br \xrightarrow{NaN_3, warm} conc. H_2SO_4$$

(b)
$$CH_3CH_2CH_2CONH_2 \xrightarrow{Br_2/KOH}$$

- (d) $CH_3CH_2CH_2Br + NH_4OH \longrightarrow$
- 90. Which of the following statements are correct regarding thiol esters?
 - (a) They are not as effectively stabilized as their oxygen analogues.
 - (b) In thiol esters, the α-hydrogens are more acidic than those of ordinary esters.
 - (c) The C—S bond of a thiol ester is weaker than the C—O bond of an ordinary ester.
 - (d) $R-\bar{S}$: is a better leaving group that R-O:.
- 91. On being heated with solid potassium hydroxide, nitrobenzene gives

(b) NO₂ OH

(c) NO₂

(d) There is no reaction.

- 92. The synthesis of α -benzylethylamine using an appropriately substituted derivative of α -benzylpropionic acid occurs with retention of configuration in
 - (a) the Schmidt reaction
- (b) the Curtius rearrangement
- (c) the Lossen rearrangement
- (d) none of these
- 93. Which of the following statements are correct?
 - (a) Pyridine is a stronger base than ethylamine.
 - (b) The basic strength of trimethylamine is less than that of dimethylamine.
 - (c) Ethyl amine is a stronger base than ammonia.
 - (d) Aniline is a weaker base than ammonia.

94. Consider the structures

$$H_3C-N-CH_3$$
 H_3C
 CH_3
 O_2N
 O_2N

Which of the following statements are correct?

- (a) The base strength of II is more than that of I.
- (b) The base strength of II is less than that of I.
- (c) The base strength of IV is more than that of III.
- (d) The base strength of IV is less than that of III.
- 95. Treatment of cyclobutylmethylamine, CH_2NH_2 , with nitrous acid gives
 - (a) CH₂OH (b) CH
 (c) OH (d)
- **96.** Treatment of *n*-propylamine with nitrous acid gives
 - (a) CH₃CH=CH₂

- (b) CH₃—CH₂—CH₂OH
- (c) CH₃—CH—CH₃ OH
- (d) none of these
- 97. The reaction of cyclobutylamine with nitrous acid gives
 - (a) OH (b) O
 (c) CH₂OH (d) none of these
- 98. Which of the following statements are correct for trimethylamine?
 - (a) It has a trigonal pyramidal shape.
 - (b) The C–N–C bond angle is 108.7° .

- (c) The N atom is sp ³-hybridized.
- (d) The unshared pair of electrons on nitrogen occupies an sp² orbital.
- **99.** Which of the following compounds are likely to react with an appropriate base to generate a negative charge on the C atom adjacent to the S atom?
 - (a) C₆H₅—SCH₃

(b) CH₃—S—CH₃

(c) CH₃—SH

- (d) C_6H_5 —SH
- **100.** Which of the following statements are correct?
 - (a) Thiol esters are more susceptible to nucleophilic attack than normal esters.
 - (b) Thiol esters are less susceptible to nucleophilic attack thn normal esters.
 - (c) The α-hydrogens of thiol esters are more acidic than those of ordinary esters.
 - (d) The carbon-sulphur bond of a thiol ester is weaker than the carbon-oxygen bond of an ordinary ester.
- 101. The intermediates obtained in the reaction

$$\begin{array}{c|c}
C & & \\
R - C - C1 & \xrightarrow{\text{NaN}_3} & R - NH_2
\end{array}$$

are

(a)
$$R \longrightarrow C \longrightarrow N \longrightarrow N \longrightarrow N$$

- (b) R—N=C=O
- (c) R-CNO

- (d) none of these
- **102.** Which of the following statements are correct?
 - (a) Thiols do not form as strong intermolecular hydrogen bonds as do alcohols.
 - (b) Thiols are considerably stronger acids than the corresponding alcohols.
 - (c) Thiols are much weaker acids than the corresponding alcohols.
 - (d) Thiols form insoluble salts with heavy metals, e.g., mercury.
- 103. Which of the following reactions are feasible for the preparation of thiols?
 - (a) $C_2H_5Br + NaSH \longrightarrow C_2H_5SH + NaBr$

(b)
$$CH_3$$
 $CH_3 - C = CH_2 + H_2S \xrightarrow{H_2SO_4} CH_3 - C - CH_3$

(c) Br
$$MgBr$$
 SH

$$Mg \longrightarrow MgBr$$
 $1.S$

$$2. H^{+}$$

(d)
$$C_2H_5OH + H_2S \longrightarrow C_2H_5SH$$

104. Which of the following are feasible?

(a)
$$CH_3CH_2CH_2CN \xrightarrow{\text{LiAlH}_4} CH_3CH_2CH_2CH_2NH_2$$

(b) OH
$$(C_6H_5)_2C = N$$
 $\xrightarrow{\text{LiAlH}_4}$ $(C_6H_5)_2CHNO_2$

(c)
$$O$$
 \parallel
 $C-C+GH_3 \xrightarrow{NH_2OH} [] \xrightarrow{(CF_3CO)_2O} \triangleright NHCOCH_3$

(d)
$$C_6H_5$$
— $CH_2CO_2H + NH_3 \xrightarrow{H_2SO_4} C_6H_5CH_2NH_2$

105. In the reaction

$$CH_3CH_2CH_2NH_2 \xrightarrow{\qquad NaNO_3/dil.\ HCl} CH_3CH = CH_3CH = CH_2$$

the possible intermediates are

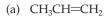
(a)
$$H$$
 (b) $CH_3CH_2CH_2-N=N-OF$ $CH_3CH_2CH_2-N-NO$ H

(c)
$$CH_3CH_2CH_2 \stackrel{+}{\longrightarrow} N$$
 (d) $CH_3CH_2\stackrel{+}{C}H_2$

106. In the reaction

$$CH_3CH_2CH_2NH_2 \xrightarrow{NaNO_2/dil. HCl} 0-5^{\circ}C$$

the products formed are





(c)
$$CH_3$$

 CH_3 — C = CH_2

(d) none of these

107. The final products (II) obtained in the reaction

$$CH_3CH_2$$
 CH_3CH_2
 $N \xrightarrow{H_2O_2} I \xrightarrow{\text{strong heating}} II$
 CH_3CH_2

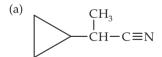
are

b) $CH_2 = CH_2$

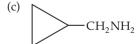
108. In the reaction

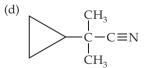
$$CH_2CN \xrightarrow{CH_3Br}$$
NaNH₂, NH₃, -80°C

the products obtained are



(b) CH₂OH





- **109.** The products obtained by the reaction of an isocyanate (R—N=C=O) with water are
 - (a) RNH₂

(c) RNHOH

(d) none of these

- 110. The reaction of CH₃(CH₂)₆Br with NaNO₂ in the presence of dimethylformamide gives
 - (a) CH₃(CH₂)₆OH
- (b) CH₂(CH₂)₆NO₂
- (c) $CH_3(CH_2)_6ONO$
- (d) There is no reaction.
- 111. Which of the following reactions are feasible?

(a)
$$(CH_3)_3CNH_2 \xrightarrow{\text{KMnO}_4} (CH_3)_3C-NO_2$$

(b)
$$ICH_2$$
— $CO_2C_2H_5 + AgNO_2 \xrightarrow{0 \circ C} O_2NCH_2CO_2C_2H_5 + AgI$

(c)
$$\sim$$
 NOH \sim CF₃CO₃H \sim NO₂
(d) \sim HONO \sim NO₂

(d)
$$\longrightarrow$$
 NO₂

112. The products obtained in the reaction

$$\begin{array}{c} CH_3 \\ I \\ -C - NO_2 \\ I \\ CH_3 \end{array} + CH_2 = O \xrightarrow{OH^{-}}$$

are

$$\begin{array}{ccc} \text{(a)} & \text{CH}_3 \\ & \text{I} \\ \text{HOCH}_2 & \text{C} & \text{NO}_2 \\ & \text{CH}_3 \end{array}$$

$$\begin{array}{c} \text{(b)} & \text{CH}_2\text{OH} \\ & \text{HOCH}_2\text{--}\text{C}\text{--NO}_2 \\ & \text{CH}_2\text{OH} \end{array}$$

- (d) none of these
- 113. The conversion $CH_3CN \longrightarrow CH_3CH_2NH_2$ can be effected by using
 - (a) Pt/H_2

(b) LiAlH₄

(c) Na/C₂H₅OH

- (d) SnCl₂/HCl
- 114. Ammonium acetate can be converted into acetamide by
 - (a) heating
 - (b) heating in the presence of P₂O₅
 - (c) using Br₂/KOH
 - (d) heating in the presence of CH₃COOH

115. Phenyl cyanide can be obtained by

(a)
$$C_6H_5CONH_2 \xrightarrow{P_2O_5, \Delta}$$

(b)
$$C_6H_5$$
— $CH=NOH$ — Ac_2O , Δ

(c)
$$C_6H_5Cl$$
 — alc. KOH \longrightarrow

(d)
$$C_6H_5NH_2 \xrightarrow{1. \text{ NaNO}_2/\text{HCl}} 2. \text{ CuCN}$$

- 116. Which of the following compounds will undergo carbylamine reactions?
 - (a) CH₃CH₂NH₂

(b) $(CH_3)_2NH$

(c) $C_6H_5NH_2$

(d) $(CH_3)_3N$

117. In the reaction

$$CH_3CONH_2 \xrightarrow{Br_2 + KOH} CH_3NH_2$$

the intermediates involved are

(a) CH₃CONHBr

(b) CH₃NHBr

(c) $CH_3N=C=O$

(d) CH₃CONBr₂

118. In the reaction

 $RCONH_2 + X \longrightarrow RNH_2$, the reagent X is

(a) PCl₅

(b) $NaOH + Br_2$

(c) soda lime

- (d) NaOBr
- **119.** Which of the following would react with nitrobenzene to produce hydrazobenzene?
 - (a) $Na_3AsO_3 + NaOH$
- (b) Zn/NaOH, CH₃OH

(c) Zn, NaOH

- (d) $NH_2NH_2 + alc. KOH$
- **120.** The conversion of *m*-dinitrobenzene into *m*-nitroaniline can be brought about with
 - (a) $(NH_4)_2S$

(b) sodium polysulphide

(c) Sn + HCl

- (d) $Zn + NH_4Cl$
- 121. Nitrobenzene can be converted into azobenzene by reduction with
 - (a) LiAlH₄/ether

- (b) Zn, NH₄Cl, Δ
- (c) Zn/NaOH, CH₃OH
- (d) Raney Ni
- 122. The nitration of aromatic organic compounds can be effected by
 - (a) a mixture of concentrated HNO₃ and concentrated H₂SO₄
 - (b) a mixture of concentrated HNO₃ and acetic anhydride

- (c) fuming nitric acid and concentrated sulphuric acid
- (d) alcoholic potassium nitrate
- 123. Which of the following represents electrophilic substitution?

(a)
$$NO_2$$
 NO_2 NO_2 NO_2 NO_2

(b)
$$NO_2 \longrightarrow NO_2$$
 aq. KOH OH

(d)
$$CH_3$$
 NO_2 NO_2 NO_2 NO_2 NO_2 NO_2

124. In the reduction of nitrobenzene into aniline, the intermediates formed are

(a)
$$N=0$$
 (b)

NHOH

- 125. Benzylamine can be prepared by
 - (a) $C_6H_5CONH_2 \xrightarrow{\text{LiAlH}_4} \text{ether}$
 - (b) $C_6H_5CH_2CONH_2 + Br_2 + KOH \longrightarrow$
 - (c) $C_6H_5CN \xrightarrow{LiAlH_4}$
 - (d) $C_6H_5CH_2NC \xrightarrow{LiAlH_4}$
- **126.** Isopropylamine can be obtained by
 - (a) $(CH_3)_2CHO + NH_2OH \longrightarrow ? \xrightarrow{LiAlH_4}$

(b)
$$(CH_3)_2CHO + NH_3 \xrightarrow{\Delta} ? \xrightarrow{H_2/Ni}$$

 CH_3
 $CHOH + NH_3 \longrightarrow$

- (d) none of these
- **127.** Which of the following statements are correct?
 - (a) In the case of primary, secondary and tertiary amines, the basic strength depends on the extent of hydrogen bonding in the protonated amines.
 - (b) The presence of groups like —OCH₃ and —CH₃ increases the basic strength of amines.
 - (c) The presence of groups like —NO₂, —CN and halogens reduces the basic strength of amines.
 - (d) The basic strength of amines depends on their concentration.
- 128. Which of the following statements are correct?
 - (a) Aniline is a weaker base than ethylamine.
 - (b) Aniline is a stronger base than *p*-methoxyaniline.
 - (c) Aniline must be acetylated before nitration with an acid mixture.
 - (d) Aniline is soluble in an ammonium hydroxide solution.
- **129.** An aliphatic nitro compound turns red with the addition of a concentrated NaOH solution, followed by the addition of an excess of an NaNO₂ solution and then dilute H₂SO₄. The colour disappears with the addition of the excess of an acid but reappears if the solution is made alkaline. The aliphatic nitro compound is
 - (a) CH₃CH₂NO₂

(b) (CH₃)₂CHNO₂

(c) (CH₃)₃CNO₂

- (d) CH₂CH₂CH₂NO₂
- **130.** *p*-Nitroaniline is obtained by
 - (a) sulphanilic acid $\xrightarrow{\text{HNO}_3}$
 - (b) benzenesulphonic acid $\xrightarrow{\text{HNO}_3/\text{H}_2\text{SO}_4}$
 - (c) aniline $\frac{1. \text{HNO}_3/\text{H}_2\text{SO}_4}{2. \text{ aq. NaOH}}$
- **131.** *p*-Nitrotoluene can be obtained by
 - (a) toluene $\xrightarrow{\text{conc. HNO}_3/\text{conc. H}_2\text{SO}_4, 30^{\circ}\text{C}}$
 - (b) toluene CH₃CO—O—NO₂

(c)
$$CH_3$$
 N_2Cl + HNO_2 Cu_2O

- (d) toluene $\frac{\text{conc. HNO}_3 + \text{conc. H}_2\text{SO}_4 > 50^{\circ}\text{C}}{\text{conc. HNO}_3 + \text{conc. H}_2\text{SO}_4 > 50^{\circ}\text{C}}$
- **132.** Which of the following amines will react with cyclohexanone to give enamine?
 - (a) CH₃NH₂

(b) (CH₃)₂NH

(c) N

- (d) N
- 133. Intermolecular hydrogen bonding is possible in the case of
 - (a) *N*-methylaniline
- (b) *N,N*-dimethylaniline

(c) methylamine

- (d) nitrobenzene
- **134.** Which of the following amines can form hydrogen bonds within themselves?
 - (a) CH₃NH₂

(b) (CH₃)₂NH

(c) (CH₃)₃N

- (d) None of these
- **135.** Which of the following reactions can be used to prepare ethyl isocyanide?
 - (a) $CH_3CH_2I + AgCN \xrightarrow{C_2H_5OH/H_2O}$
 - (b) $CH_3CH_2I + KCN \xrightarrow{alcohol} \Delta$
 - (c) $CH_3CH_2NH_2 + CHCl_3 + KOH \xrightarrow{\Delta} ACOHO$
 - (d) $CH_3CH_2Cl + KCN \xrightarrow{alcohol} \Lambda$
- 136. The conversion

$$\begin{matrix} O \\ \uparrow \\ C_6H_5NO_2 \longrightarrow C_6H_5N=N-C_6H_5 \end{matrix}$$

can be brought about by reduction with

- (a) Na₃AsO₃/NaOH
- (b) Glucose/NaOH

(c) Zn/NaOH

- (d) LiAlH₄/ether
- **137.** The electrolytic reduction of nitrobenzene in a strong acidic medium gives
 - (a) aniline

- (b) phenylhydroxylamine
- (c) *p*-aminophenol
- (d) phenylhydrazine

138. The product obtained in the reaction

$$CH_3$$
— $NO_2 + Cl_2 + NaOH$ — \longrightarrow

is

(a) ClCH₂NO₂

(b) Cl₂CHNO₂

(c) Cl₃CNO₂

(d) CH₃NH₂

Answers

1. b	2. a	3. c	4. b	5. d
6. c	7. c	8. c	9. b	10. b
11. d	12. b	13. c	14. a	15. b
16. d	17. c	18. d	19. a	20. a
21. d	22. a	23. b	24. a	25. a
26. a	27. d	28. a	29. d	30. b
31. b	32. d	33. b	34. a	35. b
36. a	37. b	38. a	39. b	40. a
41. b	42. a	43. b	44. b	45. c
46. a	47. b	48. b	49. c	50. a
51. c	52. c	53. d	54. a	55. a
56. a	57. a	58. a	59. a	60. a
61. b	62. c	63. b	64. b	65. d
66. a	67. c	68. c	69. d	70. b
71. c	72. c	73. b	74. b	75. b
76. b	77. b	78. a	79. d	80. a
81. a	82. b	83. a	84. a	85. b
86. d	87. a, b	88. c, d	89. a, b, c	90. a, b, c, d
91. a, c	92. a, b, c	93. b, c, d	94. a, c	95. a, b, c, d
96. a, b, c	97. a, c	98. a, b, c	99. a, b	100. a, c
101. a, b	102. a, b, d	103. a, b, c	104. a, b, c, d	105. a, b, c, d
106. a, b	107. a, b	108. a, d	109. a, b	110. b, c
111. a, b, c	112. a, b	113. a, b, c	114. b, d	115. a, b, d
116. a, c	117. a, c	118. b, d	119. c, d	120. a, b
121. a, c	122. a, b, c	123. a, d	124. a, b	125. a, b, c
126. a, b, c	127. a, b, c	128. a, c	129. a, d	130. a, d
131. a, c	132. b, c, d	133. a, c	134. a, b	135. a, c
136. a, b	137. b, c	138. a, b, c		

Hints to More Difficult Problems

4. Hydrogen bonding is possible only in molecules that have hydrogen atoms bound to electronegative atoms.

- 7. Aliphatic amines are generally stronger than aromatic amines.
- 22. When amides containing a chiral carbon react with NaOBr/H₂O, there is retention of configuration at the migrating atom.
- 23. The strong electron-withdrawing effect of the nitro- group coupled with the formation of the resonance-stabilized cyclohexadienylide anion constitutes the driving force for the nucleophilic attack by the hydroxide anion.
- **28.** Primary and secondary nitro-compounds undergo aldol-type and Michael additions with suitable carbonyl compounds under the influence of basic catalysts.
- **46.** A mixture of concentrated HNO₃ and concentrated H₂SO₄ is termed a nitrating mixture.

$$HO_{3}SO-H + H-O-N O H-O-N O + HSO_{4}$$
 $H-O-N O H-O-N O H-O-N O HSO_{4}$
 $H-O-N O H-O-N O HSO_{4}$

Nitronium ion reactive electrophile

- 51. The disubstituted benzene in which all the four hydrogen atoms are identical will give only one monosubstituted product.
- 53. Two types of compounds can be represented by the molecular formula C_7H_9N in case of monosubstitution.

$$C_7H_9N - C_6H_5 = -CH_2NH_2$$

One possible compound is $C_6H_5CH_2NH_2$ and the other $C_6H_5NHCH_3$. In case of disubstitution, the possible compound is

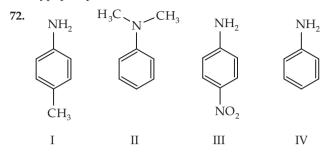
$$C_7H_9N-C_6H_4=CH_3NH_2$$
.

This is aminotoluene, which can be *o-*, *m-* or *p-*.

Therefore, in all, there can be five isomers.

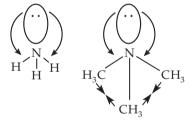
63. On reaction with nitrous acid, primary amines yield alcohols.

66. Benzenediazonium chloride undergoes deamination upon reaction with hypophosphorus acid.



In the reaction of these anilines with H^+ , electron release tends to disperse the positive charge of the resultant anilinium salt and thus stabilizes the ions relative to the amine. Electron withdrawal tends to intensify the positive charge and thereby destabilizes the ions with respect to aniline. But N,N-dimethylaniline is a stronger base than aniline. Because of the steric effect of the two methyls, the lone pair on nitrogen cannot take part in resonance with the benzene ring, and is hence more available for protonation. The correct sequence is II > I > IV > III.

84. All the three molecules are sp 3 -hybridized. Since all the bonds in CH $_4$ are equivalent, the bond angle is $109^{\circ}28'$ the normal value. However, in NH $_3$ and N(CH $_3$) $_3$, the fourth sp 3 -hybridized orbital is present as a lone pair which repels the rest of the bonds as shown below.



But this repulsion is compensated for by the steric repulsion in trimethylamine, and the H–N–H bond angle in NH_3 is less than the H_3C –N– CH_3 bond angle in triethylamine.

133. Hydrogen bonding is possible only in compounds with bonds between electronegative elements such as nitrogen, oxygen and hydrogen.

Petroleum

• Type 1 •

1. The quality of diesel oil for use in diesel engines is determined by

Choose the correct option. Only one option is correct.

(d) the amount of additives added

(a) the composition of the oil(b) the octane number(c) the cetane number

(a) isomerization(c) reforming

	C_{12} —(\mathbb{C}_{18} and a boil	ling r	ange of 300–	400°	C is known as	3	
	(a)	kerosene oil			(b)	diesel oil		
	(c)	lubricating of	oil		(d)	paraffin wax	<	
3.	Which	n of the follow	ving l	has a cetane	num	ber of 100?		
	(a)	Cetane			(b)	α-Methylna	phth	alene
	(c)	Cyclohexan	e		(d)	None of the	se	
4.		- 1				ke a 3 : 2 mixer of the diese		of cetane and is
	(a)	60	(b)	40	(c)	20	(d)	100
5.	A goo	d diesel oil s	hould	l have an oct	ane i	number of		
	(a)	30-40	(b)	> 45	(c)	10-30	(d)	5-100
6.	hydro							of aromatic essary. This is

2. During the fractionation of petroleum, the fraction with the composition

(b) alkylation

(d) distillation under vacuum

7.	The tendency of gasoline to knock decreases with change in the nature of the fuel, i.e., it depends upon whether it comprises straight-chain paraffins (I), branched-chain paraffins (II), olefins (III), aromatic hydrocarbons (IV), as follows
	(a) $I > II > III > IV$ (b) $IV > III > II > I$
	(c) $II > I > IV > III$ (d) $III > II > IV$
8.	The octane number of aviation fuel is above
	(a) 50 (b) 75
	(c) 80 (d) 100
9.	<i>n</i> -Heptane has been assigned the octane number
	(a) 0 (b) 1000
	(c) 10 (d) 50
10.	The thermal decomposition of higher hydrocarbons into lower hydrocarbons is known as
	(a) reforming (b) isomerization
	(c) cracking (d) aromatization
11.	Commercial gasoline contains mostly
	(a) straight-chain alkanes (b) cycloalkanes
	(c) branched-chain alkanes (d) aromatic hydrocarbons
12.	Arrange
	CH ₃ CH ₃
	CH ₃ CH ₃ CH ₃ CH ₃ CH ₃ CHCH ₂ CH ₃ , CH ₃ CHCH ₂ CH ₃ , CH ₃ CH ₃ CH ₃ CH ₃
	I II
	CH ₃ —CH—CH ₂ CH ₂ CH ₂ CH ₂ CH ₃ and CH ₃ CH ₂ CH ₂ CH ₂ CH ₂ CH ₂ CH ₃ CH ₃
	III IV
	in order of their octane numbers.
	(a) $I > III > II > IV$ (b) $IV > II > III > I$
	(c) $I > II > III > IV$ (d) $II > IV > I > III$
13.	Liquid hydrocarbons can be converted into a mixture of gaseous hydrocarbons by
	(a) hydrolysis
	(b) oxidation
	(c) cracking

(d) distillation under reduced pressure

14.	Petrol	eum consists mainly of		
	(a)	aromatic hydrocarbon	s (b)	aliphatic hydrocarbons
	(c)	aliphatic alcohols	(d)	none of these
15.	The fir	rst product obtained dur	ing the fract	ional distillation of petroleum is
	(a)	petroleum ether	(b)	diesel
	(c)	kerosene	(d)	none of these
16.	Which point?		tions of pe	troleum has the lowest boiling
	(a)	Diesel	(b)	Kerosene
	(c)	Gasolene	(d)	Heavy oil
17.	Paraff	in wax is		
	(a)	an ester	(b)	an alcohol
	(c)	a saturated hydrocarbo	on (d)	an unsaturated hydrocarbon
18.	Natur	al gas is a mixture of		
	(a)	CO and N ₂	(b)	CH_4 , C_2H_6 and C_3H_8
	(c)	CO and CO ₂	(d)	CO, H ₂ and CH ₄
19.	Petrol	eum ether can be used a	as	
	(a)	a fuel		
	(b)	a solvent for fats, oils a	ınd varnish	
	(c)	both		
	(d)	neither		
20.	Gasol	ine has the composition		
	(a)	$C_3 - C_5$ (b) $C_6 - C_6$	(c)	$C_8 - C_{12}$ (d) $C_{10} - C_{13}$
21.	Synth	etic petrol is produced f	rom water	gas by the
	(a)	Baeyer process	(b)	Fischer–Tropsch process
	(c)	oxo process	(d)	Bergius process
22.	The do	etection of leakage from	LPG cylino	ders is facilitated by the addition
	(a)	phenols	(b)	glycols
	(c)	thioalcohols	(d)	alcohol
23.	Which	n of the following produ		
	` '	Olefins		Aromatic hydrocarbons
	(c)	Straight-chain paraffin	s (d)	Branched-chain paraffins
24.	Which	n of the following has be	een given th	ne octane number zero?
	(a)	<i>n</i> -Octane	(b)	Iso-octane

(c) *n*-Heptane

(d) Tetraethyl lead

- 25. Tetraethyl lead is used as a
 - (a) petroleum additive
- (b) mosquito repellent

(c) painkiller

- (d) fire extinguisher
- 26. Which of the following is used as an antiknock compound?
 - (a) TEL

(b) Lead acetate

(c) Ethyl acetate

- (d) All of these
- 27. Kerosene has the composition
 - (a) $C_5 C_8$

(b) $C_8 - C_{12}$

(c) $C_{12}-C_{16}$

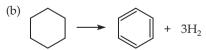
- (d) $C_{16} C_{18}$
- **28.** A sample of gasoline, on testing, behaves like a mixture of 65% iso-octane and 35% n-heptane. What is the octane number of the fuel?
 - (a) 35
- (b) 65
- (c) 30
- (d) 100
- **29.** Arrange the following in order of their knocking properties. Branched-chain paraffins (I), straight-chain paraffins (II), olefins (III) and napthenes (IV)
 - (a) I > II > III > IV
- (b) I > II > IV > III
- (c) II > I > III > IV

(d) I = II > III > IV

• *Type 2* •

Choose the correct options. More than one option is correct.

- **30.** The possible products of the thermal cracking of a straight-chain alkane $(C_{16}H_{34})$ in the presence of a catalyst are
 - (a) $C_5H_{11}CH=CH_2$
- (b) CH₂=CHCH₂CH₃
- (d) none of these
- **31.** Which of the following reactions are possible during cracking?
 - (a) $CH_3(CH_2)_{12}CH_3 \longrightarrow CH_3(CH_2)_5CH_3 + CH_3(CH_2)_4CH = CH_2$



- (c) $CH_3(CH_2)_{12}CH_3 \longrightarrow CH_3(CH_2)_4CH_3$
- (d) None of these

Petroleum 3-231

32.	Which of the following statements are correct?				
	(a)	Iso-octane has a higher octane	nur	nber than that of <i>n</i> -octane.	
		Cyclohexane has a higher octa			
	(c)	Cyclohexane has a lower octa	ne n	umber than that of <i>n</i> -hexane.	
	(d)	Iso-octane has a lower octane	num	ber than that of <i>n</i> -octane.	
33.	The th	nermal cracking of n -decane at	450-	500°C gives	
	(a)	hexane	(b)	hexene	
	(c)	pentane	(d)	pentene	
34.	Which petrol		are	obtained by the refining of	
	(a)	Toluene	(b)	Gasoline	
	(c)	Kerosene oil	(d)	Carbon tetrachloride	
35.		n of the following processes ca octane number?	ın be	e used to produce gasoline of a	
	(a)	Isomerization	(b)	Cracking	
	(c)	Alkylation	(d)	Reforming	
36.	The p	resence of oil fields below the	surf	ace of the earth can be detected	
	(a)	a survey of the gravity	(b)	determining the magnetic field	
	(c)	the seismic method	(d)	none of these	
37.	Which	n of the following statements ab	out	natural gas are true?	
	(a)	It is a mixture of gaseous hydr	rocai	rbons.	
	` '	It is used in the manufacture of	of fer	tilizers.	
	(c)	It is a mixture of CO and H_2 .			
	(d)	It is used as a fuel.			
38.	The o	ctane number of a fuel can be in	ncrea	ased by	
	(a)	isomerization	(b)	alkylation	
	(c)	reforming	(d)	fractional distillation	
39.	Which	n of the following has an octane	nur	mber of 100?	
	(a)	2,2,4-Trimethylpentane	(b)	2,2-Dimethylpentane	
	(c)	2,2,4-Triethylhexane	(d)	Iso-octane	
40.	Which	n of the following are present ir	ı coa	l tar?	
	(a)	Cresols	(b)	Alkenes	
	(c)	Cycloalkenes	(d)	Naphtha	
41.	The ca	arbonization of coal is mostly u	sefu	l for the production of	
	(a)	coke	(b)	coal-based organic chemicals	
	(c)	liquor ammonia	(d)	LPG	

- **42.** Which of the following statements are correct?
 - (a) Straight-chain alkanes have very low octane numbers.
 - (b) The branching of the chain increases the octane number of a fuel.
 - (c) Straight- and branched-chain alkanes have octane numbers higher than those of the corresponding cycloalkanes.
 - (d) Aromatic compounds have high octane numbers.

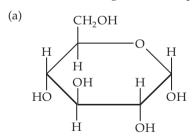
<u>Answers</u>					
1. c	2. b	3. a	4. a	5. b	
6. c	7. a	8. d	9. a	10. c	
11. d	12. c	13. c	14. b	15. a	
16. c	17. c	18. b	19. b	20. b	
21. b	22. c	23. c	24. c	25. a	
26. a	27. c	28. b	29. a	30. a, b, c	
31. a, b, c	32. a, b	33. a, b, c, d	34. b, c	35. a, b, c, d	
36. a, b, c	37. a, b, d	38. a, b, c	39. a, d	40. a, d	
41. a, b	42. a, b, d				

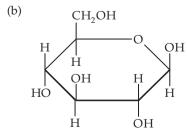
Carbohydrates

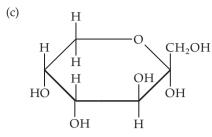
• Type 1 •

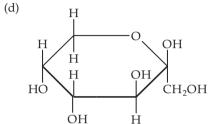
Choose the correct option. Only one option is correct.

- 1. Which of the following carbohydrates is a monosaccharide?
 - (a) Fructose
 - (b) Cellulose
 - (c) Starch
 - (d) Cane sugar
- **2.** Which of the following structures represents α -D-glucose?

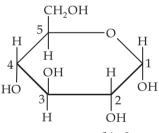








3. In α -D-glucose (structure given below), the anomeric carbon is at



(a) 1

(b) 2

(c) 4

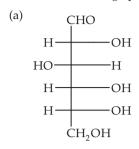
- (d) 5
- 4. On hydrolysis, which of the following carbohydrates gives only glucose?
 - (a) Sucrose

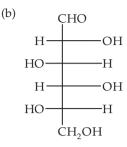
(b) Lactose

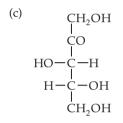
(c) Maltose

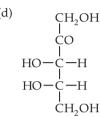
- (d) Galactose
- The presence of a primary alcoholic group (CH₂OH) in glucose is inferred by
 - (a) its oxidation by bromine water to give gluconic acid [HOOC(CHOH)₄CH₂OH], which on further oxidation with concentrated HNO₃ yields glucaric acid
 - (b) the fact that a solution of glucose in an inert solvent becomes effervescent upon the introduction of sodium
 - (c) the fact that a solution of it becomes coloured upon the addition of FeCl₃
 - (d) its reaction with the Tollens reagent

6. Which of the following open-chain structures represents D-glucose?









- 7. In the cyclic structure of glucose, the ring is
 - (a) only six-membered
- (b) only five-membered
- (c) six- as well as five-membered (d) none of these
- 8. On hydrolysis followed by heating with HI, the cyanohydrin of which of the following carbohydrates gives 2-methylhexanoic acid?
 - (a) Glucose
- (b) Fructose
- (c) Sucrose
- (d) Lactose

- 9. Grape sugar is
 - (a) glucose
- (b) fructose
- (c) maltose (d) lactose

- **10.** Which of the following is fruit sugar?

 - (a) Glucose (b) Fructose
- (c) Cane sugar (d) Starch
- 11. The urine sample of a diabetic patient contains
 - (a) sucrose
- (b) fructose
- (c) glucose
- (d) all of these
- **12.** Which of the following reduces that Tollens reagent?
 - (a) Glucose

(b) Fructose

(c) Sucrose

- (d) Starch
- 13. On oxidation with HNO₃, a carbohydrate $(C_6H_{12}O_6)$ gives a mixture of glycolic acid and tartaric acid. The carbohydrate can be
 - (a) only glucose
 - (b) only fructose
 - (c) a mixture of glucose and fructose
 - (d) none of these

- 14. An aqueous solution of a carbohydrate turns blue when a drop of iodine solution is added to it. The carbohydrate is
 - (a) glucose
- (b) fructose
- (c) lactose
- (d) starch
- 15. On oxidation with HNO₃, a carbohydrate gives a mixture of oxalic acid, tartaric acid and glucaric acid. The carbohydrate is
 - (a) glucose
- (b) fructose
- (c) cane sugar (d) starch
- 16. Glucose and fructose can be differentiated by using
 - (a) Fehling's solution
- (b) the Tollens reagent
- (c) Schiff's reagent
- (d) osazone formation
- 17. On reduction with Na-Hg, a carbohydrate gives a mixture of sorbitol and mannitol. The carbohydrate can be
 - (a) glucose

(b) fructose

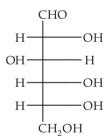
(c) cane sugar

- (d) lactose
- **18.** Which of the following is a disaccharide?
 - (a) Glucose

(b) Maltose

(c) Starch

- (d) Cellulose
- 19. A carbohydrate $(C_6H_{12}O_6)$ rotates the plane of polarization of polarized light towards the left, and the arrangement of the carbon atoms is as follows.



This carbohydrate is

(a) D(+)-glucose

(b) D(-)-glucose

(c) L(+)-glucose

(d) L(-)-glucose

• *Type 2* •

Choose the correct options. More than one option is correct.

- **20.** Which of the following are disaccharides?
 - (a) Glucose

(b) Cane sugar

(c) Maltose

(d) Starch

- **21.** Which of the following statements are correct?
 - (a) The letters D and L refer to relative configuration around the asymmetric carbon atom.
 - (b) The signs (+) and (–) refer to the direction of rotation of polarized light.
 - (c) A carbohydrate with a D configuration must belong to the (+) series.
 - (d) A carbohydrate with an L configuration must belong to the (–) series.
- 22. On hydrolysis, which of the following carbohydrates gives only glucose?
 - (a) Sucrose

(b) Lactose

(c) Maltose

- (d) Starch
- 23. The presence of a —CHO group in glucose is inferred by its,
 - (a) reaction with HCN to give a cyanohydrin
 - (b) reduction by Na-Hg to give sorbitol
 - (c) reaction with Fehling's solution
 - (d) reaction with the Tollens reagent
- **24.** Which of the following indicates that all the carbon atoms in glucose are in a straight chain?
 - (a) On being heated with HI, glucose forms *n*-hexane.
 - (b) On hydrolysis, the cyanohydrin of glucose gives *n*-heptanoic acid.
 - (c) On acetylation, glucose gives a penta-acetate.
 - (d) None of these
- 25. Which of the following statements are correct for glucose?
 - (a) It gives a positive reaction to Schiff's test for aldehydes.
 - (b) It reacts with sodium bisulphite and ammonia.
 - (c) Glucose penta-acetate does not react with hydroxylamine.
 - (d) It gives a negative reaction to Schiff's test for aldehydes.
- 26. The phenomenon of mutarotation is exhibited by
 - (a) glucose

(b) fructose

(c) lactose

- (d) maltose
- **27.** A carbohydrate gives the following compound on reaction with phenylhydrazine.

$$H-C=N.NHC_6H_5$$

 $C=N.NH.C_6H_5$
 $OH-C-H$
 $H-C-OH$
 $H-C-OH$
 $H-C-OH$
 CH_2OH

The carbohydrate can be

(a) glucose

(b) fructose

(c) lactose

- (d) maltose
- 28. Which of the following carbohydrates will give the same osazone?
 - (a) Glucose

(b) Fructose

(c) Cane sugar

- (d) Lactose
- 29. Which of the following do not undergo hydrolysis?
 - (a) Glucose

(b) Fructose

(c) Cane sugar

(d) Maltose

Answers

1. a	2. a	3. a	4. c	5. a
6. a	7. c	8. b	9. a	10. b
11. c	12. a	13. b	14. d	15. c
16. b	17. b	18. b	19. a	20. b, c
21. a, b	22. c, d	23. b, c, d	24. a, b	25 . c, d
26. a, b, c, d	27. a, b	28. a, b	29. a, b	

10

Amino Acids and Peptides

• Type 1 •

2. The proteins are hydrolysed by acids, alkalis or enzymes to give

3. Which of the following amino acids has no asymmetric carbon?

(a) a carbohydrate that gives 3 to 10 monosaccharides on hydrolysis

(c) a molecule composed of two or more α -amino acids joined by

(c) esters

(d) cycloparaffins

Choose the correct option. Only one option is correct.

(b) the phosphate ester of a nucleoside

peptide bonds

(a) amino acids (b) ethers

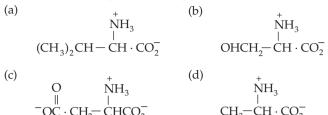
(d) an oxime

1. Peptide is

	(a)	Histidine	(b)	Glycine	(c)	α-alanine	(d)	Threonine
4.		of the essentia				corino	(4)	proling
		lysine	(D)	grychie	(C)	Sernie	(u)	promie
5.	A pep	tide bond						
	(a)	is a bond be	twee	n an oxygen	and	a carbonyl ca	ırbor	1
	(b)	is the amid- protein	e bo	nd that link	s the	amino acids	s in a	a peptide or a
	(c)	is a bond be kinds of mo			yl ca	rbon and a r	itrog	gen lipid in all
	(d)	none of thes	e					
6.	The ar	mino acids w	hich	has a nonpo	lar si	de chain is		
	(a)	lysine			(b)	serine		
	(c)	aspartic acid	d		(d)	alanine		
				3-239)			

7.	Amin	o acids are				
	(a)	acidic		(b)	basic	
	(c)	amphoteric		(d)	not dipolar ior	n
8.	Which	n of the folloing st	atements is c	orre	ct with reference	e to glycine?
	(a)	The acidic functional				
	(b)	The acidic functi	ional group is	s the	carboxylate ion	$-CO_2^-$ and the
		basic functional	group is the a	nmm	onium ion H ₃ N	<u> </u>
	(c)	The molecule is	chiral.			
	(d)	The molecules ar	re held by int	ramo	olecular hydrog	gen bonds.
9.	The to	otal number of ess	ential amino	acid	s is	
	(a)	10 (b)	20	(c)	24 (0	d) 18
10.	The h	elical structure of	protein is sta	biliz	ed bv	
		peptide bonds	1		dipeptide bon	ds
	(c)	hydrogen bonds		(d)	ionic bonds	
11.	Which	n of the following	is not found	in nı	ıcleotides?	
		Guanine		<i>a</i> .	Cytosine	
	(c)	Adenine		(d)	Tryoxine	
12.		n parts of amino in the secondary				rough hydrogen
	(a)	—COOR group		(b)	−NHCH ₃ gro	oup
	(C)	—C—NH group O)	(d)	—OH group	
13.	Rice is	s deficient in				
	(a)	alanine		(b)	glycine	
	(c)	lysine		(d)	leucine	
14.	Which	n of the following	is not a pyrin	nidiı	ne base?	
		Thymine	1,		Guanine	
	(c)	Cytosine		(d)	Uracil	

15. Which of the following structures represent for alanine?



- 16. Proteins are detected by
 - (a) Molisch's test

(b) Biuret test

(c) Benedict's test

- (d) Beilstein test
- 17. Insulin has 51 amino acids in two polypeptide chains, which are cross-linked by
 - (a) peroxide bond
- (b) disulphide bond

(c) diazo bond

- (d) two carbon-carbon double bond
- **18.** The structures obtained on acidification (H⁺) and basification (OH⁻) of alanine yields respectively.

(a)
$$CH_3-CH-COOH$$
, $CH_3-CHCOO^-$
 $| NH_3 | NH_2$

(d)
$$CH_3CH-COOH_2^+$$
, $HOCH_2-CHCOO^-$

- 19. The formation of polypeptide bond involves
 - (a) two α -amino groups and one α -carboxyl group
 - (b) one α -amino group and two α -carboxyl group
 - (c) one α -amino group of one molecule and one α -carboxyl group of another molecule
 - (d) none of these

- **20.** The pK_{a_1} and pK_{a_2} values of alanine (an amino acid) are 2.3 and 9.7 respectively. The isoelectric point (pI) of alanine is
 - (a) 3.0

(b) 7.0

(c) 8.0

- (d) 6.0
- 21. α -amino acid may be prepared by
 - (a) Williamson synthesis
- (b) Skraup synthesis
- (c) Strecker synthesis
- (d) Knorr synthesis
- 22. L-amino acids found in proteins is represented by the formula



In which of the amino acids R contains a basic group

(a) cysteine

(b) lysine

(c) aspartic acid

(d) valine

• *Type 2* •

Choose the correct options. More than one option is correct.

- **23.** Which of the following statements is correct with reference to amino acids?
 - (a) A carboxylic acid that contains an amino group
 - (b) Amino acids are the building blocks of peptides and proteins
 - (c) An α-amino acid may exists as a zwitterion under suitable condition
 - (d) none of the above
- **24.** Which of the following statements is correct with reference to isoelectric point?
 - (a) The isoelectric point is the pH at which the amino acids bear no net charge.
 - (b) It corresponds to the pH at which the concentration of the zwitterian is at a maximum.
 - (c) It is not the average of pK_{a_1} , and pK_{a_2} values
 - (d) All of these
- 25. Which of the following statements is correct?
 - (a) No enzyme can convert proteins into amino acids
 - (b) Uracil may be present in RNA but not in DNA

- (c) The left and right ends of the peptides are referred to as the N terminous (or amino terminous) and C terminous (or carboxyl terminous) respectively.
- (d) All of these.

		Answers	Answers			
1. c	2. a	3. b	4. a	5. b		
6. d	7. c	8. a	9. b	10. c		
11. d	12. c	13. c	14. b	15. d		
16. b	17. b	18. a	19. c	20. d		
21. c	22. b	23. a, b, c	24. a, b	25 . b, c		

Hints to More Difficult Problems

2. Amino acids are the basic unit of proteins

7.
$$H_2 N-CH_2-C \Longrightarrow H_3 \stackrel{+}{N} \cdot CH_2-C$$
OH

It is a neutral molecule and not an ion.

- 11. The three pyrimidine bases (cytosine, thymine and uracil) and two purine bases (adenine, guanine) are units in nucleotides.
- **12.** Proteins have $\begin{array}{cc} -C NH_2 \\ \parallel \\ O \end{array}$ linkage.
- 14. Guanine is a purine.
- **18.** Zwitterion is the hybrid of positive and negative ionic groups.

19.
$$R_1$$
 R_2
 $^+H_3N-CH-COO^-+^+H_3N-CH-COO^-$

$$R_1 O R_2$$
 $^-H_2O +^+H_3N-CH-C-NH-CH-COO^-$

20.
$$pI = \frac{pK_{a_1} + pK_{a_2}}{2} = \frac{2.3 + 9.7}{2} = 6.0$$

21.

RCHO + NH₃ + HCN
$$\longrightarrow$$
 RCHCN $\stackrel{H_3O, \text{ heat}}{\longrightarrow}$ R · CHCO $_2^-$ NH₃ $\stackrel{}{\sim}$ $\stackrel{}{\sim}$ NH₃ $\stackrel{}{\sim}$ $\stackrel{}{\sim}$ amino nitrile α -amino acid

22. —CH₂CH₂CH₂CH₂NH₂

11

Organic Polymers

• Type 1 •

Choose the correct option. Only one option is correct.

- 1. Polymer consists of large molecules
 - (a) called macromolecules which are made by linking together repeated units of small molecules, called monomer
 - (b) called capolymer
 - (c) called ε-caprolactum
 - (d) all of the above
- 2. On the basis of intermolecular forces, polymers are classified as
 - (a) rubbers
 - (b) fibres
 - (c) elastomers, fibres, thermoplastics and thermosetting
 - (d) amino acids
- **3.** Which of the following is not a natural polymer?
 - (a) Wool
- (b) Silk
- (c) Cotton
- (d) Teflon

- 4. Isoprene on polymerization, produces
 - (a) synthetic rubber
 - (b) gutta-percha
 - (c) neoprene
 - (d) cis-poly (2-methyl-1,3-butadiene)
- 5. Natural rubber is obtained from latex, which is a
 - (a) mixture of wood, plants and gums
 - (b) colloidal dispersion of rubber in water
 - (c) mixture of chloroprene and carbohydrate
 - (d) none of these

6.	Natur	al silk is		
	(a)	polyester	(b)	polyamide
	(c)	epoxide	(d)	polyurethane
7.	Natur	al rubber is a polymer, derived	d fro	m
	(a)	1,3-butadiene		isoprene
	(c)	protein	(d)	DNA
8.	Step-g	growth polymers are formed b	y	
	(a)	the reaction of a single morfunctional groups A and B.	nom	er that possesses two different
	(b)	the reaction of two differencentrated HNO ₃	erent	bifunctional monomers and
	(c)	the intermolecular reaction of	f bifu	nctional molecules
	(d)	all of these		
9.	A pol	yurethane is the product of		
	(a)	toluene-2,6-diisocyanate and blowing agent	eth	ylene glycol in presence of a
	(b)	∈-caprolactum and ethylene	glyco	ol .
	(c)	terephthalic acid and ethylen	e gly	col
	(d)	an isocyanate and an alcohol		
10.	Which	n of the following is not a copo	lyme	er?
	(a)	cross copolymer	(b)	block copolymer
	(c)	random copolymer	(d)	Graft copolymer
11.	Polyn	neric molecules are held by		
	(a)	interatomic forces	(b)	coulombic forces
	(c)	intermoleculer forces	(d)	gravitational forces
12.	The p	olymers such as polyethylene	are	
		held together by vander Waa		
		held together with the forces		•
		closely packed with coulomb	ic for	ces
	` '	none of these		
13.		ple of thermosetting plastic is/		DVIC
		Bakelite		PVC
	(c)	polyurethane		Mylar
14.		ene is a condensation polymer		
		benzoic acid terephthalic acid		acetic acid
	(()	S. I. S. DITTI GITT. GV. IU		CRATIC VIII. GIVINA

Organic Polymers 15. The fibre obtained by the condensation of hexamethylene diamine and adipic acid is (a) Dacron (b) nylon 6,6 (d) Teflon (c) Rayon 16. A raw material used in making nylon is (b) 1,3-butadiene (a) adipic acid (d) cyclohexanone (c) ethyne 17. ∈-caprolactum is the starting material for the manufacture of nylon 6 and is obtained by Beckmann rearrangement of (a) (c) (d) OH OH **18.** The repeating units of PCTFE is (a) $CF_2=CF_2$ (b) CH₂=CH₂ (c) CF_3 — CF_3 (d) FClC=CF₂ 19. The repeating units of PTFE are (a) CH≡CH (b) CF₃—CF₃ (c) CH₂=CHCN (d) $CF_2=CF_2$ 20. Glyptal is the polymer of ethylene glycol and (a) terephthalic acid (b) adipic acid (c) benzoic acid (d) picric acid 21. A polymer which is used for making ropes and carpet fibres is (b) polypropylene (a) polyacetylene

(d) PVC

(b) venyl acetate

(d) phenol and formaldehyde

(c) polyacrylonitrile

(c) neoprene

(a) methyl methacrylate

22. Hard plastic covers of telephone are made of polymer of

23.	The p	olymer which contains nitroge	n is	
	(a)	PVC	(b)	Teflon
	(c)	butyl rubber	(d)	nylon
24.	The p	roduct of addition polymeriza	tion	reaction is
	(a)	PVC	(b)	nylon
	(c)	terylene	(d)	polyamide
25.	Cellul	lose is a condensation polymer	of	
	(a)	maltose	(b)	β-glucose
	(c)	α-glucose	(d)	β-fructose
26.	Which	n of the following is a "polyam	ide"	?
	(a)	Rayon	(b)	Terylene
	(c)	nylon	(d)	Orlon
27.	Teflor	n, polystyrene and neoprene ar	e all	
		copolymers		condensation polymer
	(c)	homopolymers	(d)	monomers
28.	Teflor	1		
	(a)	$(-CF_2-CF_2-)_n$	(b)	$-(CCl_2-CCl_2)_n$
		$-(CBr_2-CBr_2)_n$		CF_2Cl_2
29.	The p	roduct of addition polymeriza	tion	reaction is
	_	PVC		nylon
	(c)	Terylene	(d)	polyamide
30.	Isopre	ene is used in making		
		petrol	(b)	nylon
	(c)	rubber	(d)	liquid fuel
31.		n of the following is an inert pastick cookware?	olyn	ner used in coating, particularly
	(a)	Teflon	(b)	Cellulose
	(c)	Bakelite	(d)	Orlon
32.	Cellul	lose trinitrate, also called "gun	cotto	on" is used in
		Cellophane paper		dyes
	(c)	explosives	(d)	making rayon
33.		lose contains glucose units j molecules are held by	oine	d by β-1,4-glycosidic linkages
		ionic bond	(b)	intramolecular hydrogen bonds
	(c)	weak vander Waals forces	(d)	all of these

- **34.** Which of the following cannot serve as a food source for human?
 - (a) Proteins
- (b) Starch
- (c) Enzymes
- (d) Cellulose
- **35.** Natural rubber is a *cis*-1,4-polyisoprene. During vulcanization, natural rubber is heated with sulphur. As a result, a reaction takes place
 - (a) that produces cross-links between the cis-polyisoprene chains
 - (b) and makes the rubber much harder
 - (c) and sulphur reacts both at the double bonds and at allylic hydrogen atoms
 - (d) all of these
- 36. Nylon-6,6 is so named because it is a polyamide, formed from a
 - (a) six-carbon dibase and a six-carbon diamine
 - (b) six-carbon diacid and a six-carbon diamine
 - (c) six-carbon diacid and a six-carbon diamide
 - (d) six-carbon ∈-caprolactum and a six-carbon diamine
- 37. Rubber is a
 - (a) conducting polymer
 - (b) oriented polymer
 - (c) elastomer
 - (d) strong commercially available fabric

• *Type 2* •

Choose the correct options. More than one option is correct.

- 38. Indicate the correct statement for chain-growth polymers.
 - (a) Chain-growth polymers are made by the addition of monomers to the end of a growing chain.
 - (b) The end of a chain is reactive because it is a radical, a cation or an anion.
 - (c) Polystyrene is the example of this class.
 - (d) None of the above.
- **39.** Chain-growth polymerization may proceed by the following mechanism
 - (a) condensation polymerization (b) cationic polymerization
 - (c) anionic polymerization
- (d) all of these
- **40.** Examples of chain-growth polymer is/are
 - (a) polystyrene

(b) nylon 6

(c) Teflon

(d) all of these

- 41. Examples of step-growth polymer is/are
 - (a) nylon 6

(b) nylon 6,6

(c) Kevlar

(d) none of these

42. Polycarbonate is

- (a) used in the manufacture of compact disc (CD)
- (b) prepared by the reaction between phosgene and bisphenol A
- (c) PVC
- (d) all of these

43. Epoxy resin is

- (a) prepared by the reaction of bisphenol A and epichlorohydrin followed by a hardener
- (b) a cross-linked polymer
- (c) an epoxy adhesive
- (d) all of these

44. Polyacetylene is a conducting polymer and

- (a) is prepared by the polymerization of acetylene using a Ziegler-Natta catalyst
- (b) the conjugated double bonds in polyacetylene causes it to conduct electricity
- (c) is used for the manufacture of electrodes for measuring pH
- (d) is not a synthetic metal

45. Crystalline polymers are

- (a) denser
- (b) harder
- (c) heavier and good conductor of heat
- (d) all of these

46. Thermoplastic polymers are those

- (a) that have ordered crystalline regions and amorphous noncrystalline regions both.
- (b) that are hard at room temperature but on heating they become soft enough to be moulded.
- (c) that are used in combs, toys, light switch plates and telephone casting.
- (d) none of these

47. Thermosetting polymers are

- (a) cross-linked chain polymers
- (b) hard enough and cannot be remelted by heating

- (c) cross-linking reduces the mobility of polymer chains and thus rendered them brittle
- (d) all of these
- **48.** Which of the following belong to the class of natural polymers?
 - (a) Proteins

(b) Cellulose

(c) Teflon

(d) All of these

	<u>Answers</u>			
1. a	2. c	3. d	4. d	5. b
6. b	7. b	8. a	9. a	10. a
11. c	12. a	13. a	14. c	15. b
16. a	17. b	18. d	19. d	20. a
21. b	22. d	23. d	24. a	25 . b
26. c	27. c	28. a	29. a	30. c
31. a	32. c	33. b	34. d	35. d
36. b	37. c	38. a, b, c	39. b, c	40 . a, c
41. a, b, c	42. a, b	43. a, b, c, d	44. a, b, c	45. a, b
46. a, b, c	47. a, b, c, d	48. a, b		

Hints to More Difficult Problems

3. Teflon is artificially made as $(...CF_2-CF_2...)_n$

14. O O O HO-CH₂CH₂·OH + HO – C
$$\stackrel{\text{H}^+}{\sim}$$
 C – OH $\stackrel{\text{H}^+}{\sim}$

Poly (ethylene terephthalate) + H₂O Terylene

- **30.** Isoprene is $CH_2 = \dot{C} CH = CH_2$
- 37. A polymer, that can stretch and then revert back to its original shape is called elastomer.

12

Practical Organic Chemistry

• Type 1 •

1. Sulphur present in an organic compound is detected by treating the

2. Which of the following nitrogenous compounds does not give blue colour in the usual Lassaigne's test for the detection of nitrogen?

(c) Aniline

(d) CH₃SH

(b) potassium ferrocyanide

(d) ammonium thiocyanate

(d) Hydrazine

Choose the correct option. Only one option is correct.

(b) Urea

'sodium extract' with

(a) Glycine

(c) $Na_2S_2O_3$

(a) potassium ferricyanide

(c) sodium nitroprusside

3.	nitrop is pre	russide is added to the sodium	ı ext	organic compound, sodium ract of the compound. If sulphur colour is obtained due to the	
	(a)	Fe(CN) ₂	(b)	$K_3[Fe(CN)_5NS]$	
	(c)	$Na_{4}[Fe(CN)_{5}NO\cdot S]$	(d)	$Na_4[Fe(CN)_6]$	
4.	 Lassaigne's test is performed to detect the presence of the elements N, S, X & P in an organic compound. In this test the organic substance is at first fused with 				
	(a)	NaCl	(b)	metallic sodium	
	(c)	metallic copper	(d)	NaOH	
5.	In La	1 1	esen	NaOH tin the organic compound, on	

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imparts green colour to the flame. This test is known as

(a) Marsh's test

(c) Gutzeit test

gives blood-red colouration with

Halogens present in organic compounds may be detected by heating the compound on a copper foil in a bunsen nonluminous flame whereby it

7. The sodium extract prepared from sulphanilic acid, contains SCN⁻. It

(b) Lassaigne's test

(d) Beilstein test

	(a)	FeCl ₃	(b)	Na_2CS_3
	(c)	FeSO ₄	(d)	a mixture of Na ₂ S and CS ₂
8.	chlori heated soluti prior	ne in the compound, the sodiud with a few drops of concenton is added to get a precipitate to addition of AgNO ₃ , is required.	im ex trate e of A red	N, S and Cl. For detection of xtract of the compound is at first d HNO ₃ and then silver nitrate AgCl. This digestion with HNO ₃ ,
		to prevent the formation NO ₂	-	
	(b)			ons to volatile HCN and H_2S , and the test by forming AgCN and
	(c)	to prevent the hydrolysis of N	laC1	N and Na ₂ S
	(d)	to form S_4N_4 which prevent the	he fo	ormation of AgCl with AgNO ₃
9.	contai blood		vith s This	ract' of an organic compound sodium nitroprusside solution, a is due to the formation of sodium thiosulphate
	(c)	ferric sulphocyanide	(d)	thiourea
10.	nitrog	a nitrogenous organic com en present in the compound is sodium nitrate	con	nd is fused with sodium, the verted into sodium nitrite
	(c)	sodamide	(d)	sodium cyanide
11.	treate appea	d with $NaNO_2$ + glacial acetic rance of	acid	ne, the sodium fusion filtrate is + CCl ₄ . Iodine is detected by the
		purple colour in the organic l		
	(b)	brown colour in the organic l	ayer	of CCl ₄
	(c)	deep blue colour in CCl ₄ laye	r	
	(d)	yellow colour in CCl ₄ layer		
12.	A mix	ture of acetone and carbon tet	rach	loride can be separated by
	(a)	fractional crystallization	(b)	fractional distillation
	(c)	steam distillation	(d)	vacuum distillation

13.	Benzoic acid can be separated from a mixture of phenol and benzoic acid by treatment with			
	(a)	NaHCO ₃ solution	(b)	NaOH solution
	(c)	Na ₂ S ₂ O ₃ solution	(d)	FeCl ₃ solution
14.	Ortho	nitrophenol can be separated f	rom	paranitrophenol by
		chromatography		solvent extraction
	(c)	steam distillation	(d)	sublimation
15.	Anthr	acene can be purified by		
		sublimation	(b)	crystallization
	(c)	distillation	(d)	filtration
16.	Rectif	ied spirit contains		
		95.6% ethanol and 4.4% meth	anol	
	(b)	100% ethanol		
	(c)	95.6% ethanol and 4.4% water	r	
	(d)	95.6% ethanol and 4.4% benze	ene	
17.	A liqu purifi		pose	es at its boiling point. It can be
	(a)	simple distillation		
	(b)	sublimation		
	(c)	distillation under reduced pre	essui	re
	(d)	all of these		
18.	Anilir	ne can be separated from pheno	ol us	ing
	(a)	NaHCO ₃	(b)	dilute HCl
	(c)	NaCl	(d)	conc. HNO ₃
19.	KOH	can be used as drying agent fo	r	
	(a)	amines	(b)	acids
	(c)	phenols	(d)	esters
20.	Which	n of the following compounds	are p	ourified by steam distillation?
	(a)	Nitrobenzene	(b)	Chlorobenzene
	(c)	Orthonitrophenol	(d)	All of these
21.	Quick	time can only be used for dry	ing	
	(a)	ethanol	(b)	phenols
	(c)	esters	(d)	carboxylic acid
22.	Silver	salt method is used for the det	term	ination of molecular weight of
	(a)	organic bases	(b)	organic acids
	(c)	aliphatic amines	(d)	esters

23.	The m	The molecular weight of aniline is determined by				
	(a)	converting it into its chloroplatinate salt and then estimating platinum obtained by ignition of the salt.				
	(b)	converting it into its acetate				
	(c)	making aniline into its tribror	no d	erivative		
	(d)	all of these				
24.		n of the following methods is usic compounds?	ısed	for the estimation of nitrogen in		
	(a)	Hypobromite method	(b)	Rast method		
	(c)	Dumas' method	(d)	Carius method		
25.		n of the following methods is a ic compounds?	ısed	for the estimation of sulphur in		
	(a)	Carius method	(b)	Victor Meyer's method		
	(c)	Kjeldahl method	(d)	Dumas' method		
26.	6:1 a			ydrogen percentages in the ratio in the ratio 3 : 4. The compound		
	(a)	CH ₂ O	(b)	CH_4O		
	(c)	C_2H_6O	(d)	CHO ₂		
27.		B mole of a carbohydrate of em drogen. The molecular formula		al formula CH ₂ O contains 1.00 g he carbohydrate is		
	(a)	$C_5H_{10}O_5$	(b)	$C_3H_4O_3$		
	(c)	$C_{12}H_{22}O_{11}$	(d)	$C_6H_{12}O_6$		
28.		n of the following aliphatic ald I solution gives an yellow resi		les on heating with concentrated precipitate?		
	(a)	CH₃CHO	(b)	CCl ₃ · CHO · H ₂ O/CCl ₃ CH(OH) ₂		
	(c)	НСНО	(d)	All of these		
29.		n of the following compound NaOH solution?	s wi	ll give chloroform on warming		
	(a)	CCl ₃ CH(OH) ₂	(b)	H · CHO		
	(c)	CH ₃ CONH ₂	(d)	C ₆ H ₅ NHCOCH ₃		
30.		n of the following organic co d with Na ₂ CO ₃ solution?	mpo	unds will not yield CO ₂ when		
	(a)	Benzoic acid	(b)	Phenol		
	(c)	Sulphanilic acid	(d)	Orthonitrophenol		
31.	Pheno	ol and carboxylic acid can be di	isting	guished from each other using		
		NaOH solution		NaCl solution		
	(c)	NaHCO ₃ solution		none of these		
	` '	5	` '			

(a) Fehling's solution

nitroprusside will produce

(a) black colouration(c) blue colouration

(c) evaporation by heating(d) alkaline KMnO₄ solution

(b) ammoniacal silver nitrate solution

32. HCHO and CH₃CHO can be distinguished from each other by the use of

33. Acetaldehyde on treatment with alkaline solution of sodium

(b) yellow colouration

(d) red colouration

34.	Aceta	ldehyde and acetone can be dis	sting	uished by
	(a)	iodoform test	(b)	nitroprusside test
	(c)	Fehling's solution test	(d)	NaHSO ₃ test
35.	Which	n of the following compounds	will	not give iodoform test?
	(a)	CH ₃ COCH ₃	(b)	$CH_3CO \cdot C_6H_5$
	(c)	CH ₃ CH ₂ COCH ₂ CH ₃	(d)	CH ₃ CH ₂ OH
36.	(a) (b) (c)	anol and ethanol can be disting iodoform reaction esterification oxidation with acidified K_2Cr acrolein test		,
37.	 Which of the following observations is correct and is useful in identifying carboxylic acids? (a) Carboxylic acids liberate CO₂ gas from NaHCO₃ solution. (b) They produce fruity smell of esters when heated with alcohol in presence of concentrated H₂SO₄. (c) Acids liberate I₂ from a mixture of KIO₃ and KI. (d) All of these 			from NaHCO ₃ solution.
38.	(a)	h of the following carboxylic ac CH ₃ CO ₂ H (COOH) ₂	(b)	will give "silver-mirror test"? $H \cdot CO_2H$ $CH_3CO \cdot CO_2H$
39.	with 2 (a)	h of the following compounds 2,4-dinitrophenylhydrazine rea $CH_3CH_2 \cdot CO_2H$ CH_3COCH_3	gent (b)	give orange-yellow precipitate? CH ₃ COOC ₂ H ₅ C ₆ H ₅ OH
40.	Which	-	mp	ounds will give foul odour of

(a) para-toluidine

- (b) Glycine
- (c) Anthranilic acid
- (d) Sulphanilic acid
- 41. An organic compound is treated with NaNO₂ and dilute HCl at 0°C and then the resulting solution is added to an alkaline solution of β -naphthol whereby a brilliant red dye is produced. This observation indicates that the compound possesses
 - (a) —NO₂ group

- (b) —CONH₂ group
- (c) aromatic—NH₂ group
- (d) aliphatic—NH₂ group
- **42.** Which of the following aromatic amines will undergo Liebermann's reaction?
 - (a) C₆H₅NHCH₃

(b) $C_6H_5N(CH_3)_2$

(c) $(C_2H_5)_3N$

- (d) C_5H_5N
- Nitrobenzene on heating with a mixture of conc. HNO₃ and conc. H₂SO₄ at 100°C produces
 - (a) *p*-dinitrobenzene
- (b) *m*-dinitrobenzene
- (c) *o*-diritrobenzene
- (d) benzene sulphonic acid

• *Type 2* •

Choose the correct options. More than one option is correct.

- **44.** In Lassaigne's test, the organic compound is at first fused with sodium metal. The sodium metal is used because
 - (a) the melting point of sodium is low so it is easily fused with organic substances
 - (b) sodium is very much effective to bring about destructive reductions of organic compounds forming ionic inorganic salts NaCN, Na₂S and NaX
 - (c) all sodium salts are soluble in water.
 - (d) none of these
- **45.** Which of the following reactions occur during the detection of nitrogen in organic substances by Lassaigne's test?
 - (a) $Na + C + N \rightarrow NaCN$
 - (b) $FeSO_4 + 6NaCN \rightarrow Na_4[Fe(CN)_6] + Na_2SO_4$
 - (c) $3Na_4[Fe(CN)_6] + 2Fe_2(SO_4)_3 \rightarrow Fe_4[Fe(CN)_6]_3 + 6Na_2SO_4$
 - (d) All of these

- **46.** Which of the following statements is correct?
 - (a) Two solid organic substances are said to be different if their mixed melting point is depressed below the melting points of both of these.
 - (b) Ethanol and water cannot be separated from each other completely by simple distillation as they form azeotropic mixture.
 - (c) Impure glycerine can be purified by ordinary distillation.
 - (d) All of these.
- **47.** Which of the following observations is correct in context to acetaldehyde?
 - (a) It usually restores pink colour rapidly to Schiff's reagent.
 - (b) It does not give silver-mirror test with ammoniacal silver nitrate solution.
 - (c) It usually reduces Fehling's solution
 - (d) All of these
- 48. Which of the following compounds will respond to iodoform test?
 - (a) CH₃CHOHCO₂H
- (b) CH₃CHOHCH₃

(c) CH₃COCO₂H

(d) None of these

1. c	2. d	3. c	4. b	5. a
6. d	7. a	8. b	9. c	10. d
11. a	12. b	13. a	14. c	15. a
16. c	17. c	18. b	19. a	20. d
21. a	22. b	23. a	24. c	25 . a
26. a	27. d	28. a	29. a	30. b
31. c	32. c	33. d	34. c	35. c
36. a	37. d	38. b	39. c	40 . a
41. c	42. a	43. b	44. a, b, c	45. a, b, c
46. a, b	47. a, c	48. a, b, c		

Hints to More Difficult Problems

- 2. It does not produce NaCN, because N_2H_4 does not contain carbon
- 7. Sulphanilic acid contains N, S and C which give NaSCN. With FeCl₃ it gives blood-red precipitate of Fe(SCN)₃
- 11. $2\text{NaI} + 2\text{NaNO}_2 + 4\text{HAc} \rightarrow I_2 + 2\text{NO} + 4\text{NaAc} + 2\text{H}_2\text{O}$

- **18.** Aniline forms $C_6H_5NH_3^+Cl^-$ (solid salt)
- 19. Amines and KOH are both bases
- 21. Rest will react with Ca(OH)₂ produced by the absorption of water by CaO.
- 26. \therefore C: H: O = 6:1:8, % C: H: O = 6/15 × 100: 1/15 × 100: 8/15 × 100 = 40: 6.67: 53.3 = 40/12: 6.67/1: 53.3/16 = 3.33: 6.67: 3.33 = 1:2:1 i.e. CH₂O
- **30.** Phenol is least acidic among all.
- 35. Does not have CH₃CO—C group and CH₃CH(OH) group.
- **40.** No odour is detected due to the non-volatility of the acidic isocyanide in the alkaline solution.
- 42. Secondary amine undergo Liebermann's reaction.

13

Assertions and Reasons

The following questions consist of an *assertion* in column 1 and a *reason* in column 2. Use the following key to choose the appropriate answer.

- (a) If the *assertion* as well as the *reason* are correct, and the *reason* is the correct explanation of the *assertion*.
- (b) If the *assertion* as well as the *reason* are correct, but the *reason* is not the correct explanation of the *assertion*.
- (c) If the *assertion* is correct, but the *reason* is incorrect.
- (d) If the *assertion* is incorrect, but the *reason* is correct.

Assertion

- The addition of HCl to alkenes in the presence of peroxides leads to the formation of an anti-Markovnikov product.
- The addition of HI to alkenes in the presence of peroxides gives an anti-Markovnikov product.
- In α, β-unsaturated compounds in which C=C and C=O groups are conjugated, there is nucleophilic addition to the C=C bond.
- In C₆H₅CH=CH—CHO, the addition of C₆H₅MgBr takes place at the C=O bond.
- **5.** Acetic acid is stronger than formic acid.

Reason

The H—Cl bond being stronger, the homolytic cleavage does not favour the production of chlorine free radicals.

The homolytic cleavage of HI does lead to the formation iodine free radicals but they combine to form iodine molecules.

The C=O is stronger than the C=C bond.

The C=O bond is stronger than the C=C bond.

In acetic acid, the electronreleasing inductive effect of the methyl group makes it difficult to break the O–H bond. NH₃ < MeNH₂ < Me₂NH > Me₃N
is the order of basicity of the
amines shown.

The basicity of amines depends on the magnitude of the +1 effect.

7. In butadiene, the C₂—C₃ single bond is slightly shorter than a carbon-carbon single bond.

The resonating structures of butadiene show that the C_2 — C_3 bond has a partial double-bond character.

8. In CH₂=CHBr, the halogen is more reactive than in CH₃CH₂Br.

Due to the +M effect of the halogen, the C—Br bond in vinyl bromide has a partial double-bond character.

The conjugation of the double bond with the carbonyl group lends resonance stability.

the major product of the dehydration of 3-hydroxy-4-methylpentanal.

2° carbocations are more stable than 1° carbocations.

10. The addition of HBr to propene gives 1-bromopropane.

Since an $S_N 1$ reaction is of the first order, its rate is independent of the concentration of OH^- .

11. In the hydrolysis of tertiary butyl chloride by the S_N1 mechanism, the rate-determining step is the ionization of tertiary butyl chloride, leading to the formation of a carbocation.

The nitro group is *m*-directing.

12. The nitration of toluene gives a mixture of *o*- and *p*-nitrotoluenes.

Normally the elimination of HX from an alkyl halide gives an alkene as per the Saytzeff rule. However, due to steric reasons a non-Saytzeff product may be obtained.

13. Contrary to the Saytzeff rule, the reaction of

$$\begin{array}{c} Me \\ He_{3}C--CH_{2}--C-Br \\ He \end{array}$$

with sodium ethoxide gives

Me
$$\mid$$
 Me₃C—CH₂—C=CH₂ as the major product.

- **14.** The diazotization of 1-aminobutane followed by warming with water gives 1-butanol as the major product.
- **15.** The Beckmann rearrangement of the oxime

$$R$$
 $C=N$
 OH

gives RCONHR'.

- **16.** The Cannizzaro reaction of C_6H_5CHO and HCHO gives C_6H_5COOH and CH_3OH .
- **17.** *p*-Dimethylaminobenzaldehyde does not undergo a Cannizzaro reaction.
- **18.** On being heated, 2,4,6-trimethylphenol allyl ether gives 3-allyl-2,4,6-trimethyl phenol.
- **19.** Benzaldehyde does not undergo aldol condensation.
- 20. On being heated with strong NaOH, C₆H₅CHO + CH₃CHO gives C₆H₅CH=CHCHO.
- 21. The addition of HBr to NCCH=CH₂ gives NC—CH₂—CH₂Br as the major product contrary to the expected NCCH(Br)CH₃.

The diazotisation of 1-aminobutane produces 1°-carbocation, which is rearranged to the more stable 2°-carbocation.

In a Beckmann rearrangement, the migrating group is always syn (i.e., cis) to the hydroxyl group.

In the crossed Cannizzaro reaction of CH_2O with aldehydes without α -hydrogen, CH_2O is oxidized and the other group is reduced.

A Cannizzaro reaction depends on the nucleophilic attack on the carbonyl carbon. So factors which reduce the positive charge on the carbonyl carbon retard the reaction. In extreme cases the reaction may not occur.

The Claisen rearragement of aryl allyl ethers give the corresponding *o*-allylphenols. If both the *o*-positions are occupied, a *p*-product is obtained. Migration to the *m*-position is not possible.

In aldol condensation it is necessary for the aldehyde to have an α -hydrogen, since the reaction involves the formation of carbanion.

In a Cannizzaro reaction, a strong alkali is used at high temperature, whereas in a Claisen reaction, a dilute alkali is used at room temperature.

Due to the strong electronwithdrawing effect of the CN group, in propene nitrile the secondary carbocation is less stable than the primary carbocation. **22.** Cyclo-octatetraene is said to be an aromatic compound.

Conjugated aromatic hydrocarbons containing $(4n + 2)\pi$ electrons are aromatic while those containing $4n\pi$ electrons are antiaromatic or nonaromatic.

23. The S_N2 reaction of *cis-*3-methylcyclopentyl bromide gives *cis-*3-methylcyclopentanol.

In S_N 2 reactions, there is an exclusive attack from behind.

24. On treatment with concentrated H₂SO₄, tert. butyl alcohol does not give ditertiary butyl ether; instead it gives 2-methyl propene.

On reaction with concentrated H_2SO_4 , tertiary alcohols do not give ethers due to the instability of ethers formed in H_2SO_4 and also due to steric crowding.

25. On being heated with *n*-butanol in the presence of sodium butoxide (*n*-C₄H₉ONa), ethylacetate gives *n*-butylacetate.

In trans esterification the alkyl group of one ester is replaced by the alkyl group of another alcohol.

26. Aliphatic as well as aromatic amines can be prepared with equal ease by making potassium phthalimide react with alkyl or aryl halides followed by hydrolysis. The halogen in aryl halides is inert compared to alkyl halides in nucleophilic displacements.

27. Aliphatic amines are weaker bases than pyridine.

In pyridine the nitrogen atom is in the sp² state and so retains a greater hold on the unshared pair of electrons than the nitrogen atom in an aliphatic amine, where it is in the sp³ state of hybridization.

28. The basic strength of *m*-nitroaniline is more than that of *p*-nitroaniline.

The nitrogen atom in a primary amine is sp ³-hybridized.

29. Guanidine is a much stronger base than other amines.

Guanidine is stabilized by resonance and a very strong base.

30. *N,N-*2,6-Tetramethylaniline has more base strength than *N,N*-dimethylaniline.

The steric inhibition of o,o-disubstituted anilines increases the base strength of amines.

 Arenediazonium salts are much more stable than their aliphatic counterparts. In arenediazonium salts there is resonance, i.e., dispersal of positive charge on the benzene ring.

- Some alcohols are poured in small quantity in the fuel tanks of automobiles, particularly in cold countries.
- **33.** Tetrachloroethene has zero dipole moment.
- **34.** The acetate ion is a weaker base than the ethoxide ion.
- Isopropyl benzene can be oxidized by KMnO₄ to give benzoic acid but tertiary butyl benzene is resistant to oxidation by KMnO₄.
- **36.** The structure

$$\begin{array}{c} CH_3 \\ H \nearrow C \nearrow Br \\ H \nearrow CH_3 \end{array}$$

has two asymmetric carbon atoms but does not show optical activity.

37. The major product obtained in the reaction

$$CH_{3} \ominus CH_{3} \ominus CH_{3} - C-O + CH_{3}CH_{2} - C-Br$$

$$CH_{3} - C-O + CH_{3}CH_{2} - C-Br$$

$$CH_{3} - CH_{3} - CH_{3}$$

$$CH_{3} - CH_{3}$$

$$CH_{2} - CH_{3}$$

$$CH_{3} - CH_{2}$$

$$CH_{3} - CH_{3}$$

(less substituted) and not

$$CH_3CH=C$$
 CH_3
 CH_3

(more substituted).

The moisture present in gasoline gets dissolved in the alcohol, and therefore, does not freeze.

The dipole moment arises from the highly polar carbon–chlorine bond.

In carboxylic acids, the carbonyl group is polarized and so the carbon of the carbonyl group bears a positive charge.

For the oxidation of the sidechain in benzene it is essential that a hydrogen atom be present in the α -position (benzylic position).

The meso compounds are optically inactive.

The dehydrohalogenation of an alkyl halide by a base (e.g. C_2H_5ONa) gives a more substituted olefine (as per Saytzeff's rule). However dehydrohalogenation with a bulky base (e.g., tertiary butoxide) gives a less substituted olefine as the major product.

38. The $S_N 2$ reaction of sodium alkylide (RC $\stackrel{-}{=}$ C: Na) with a 2° alkyl halide

$$R_2$$
 R_1 — CH_2 — CH — Br gives a mixture of RC = CH and R_1 — CH = CHR_2 instead of the expected

$$\begin{array}{c} R_1 \\ | \\ RC = C - CH = CH - R_2. \end{array}$$

39. The reaction of methyl magnesium bromide (1 mole) with 4-hydroxy-2-butanone (HOCH₂—CH₂C—CH₃) does not

give the expected

$$\begin{array}{c} CH_3 \\ | \\ HOCH_2--CH_2--C--CH_3. \\ | \\ OMgBr \end{array}$$

Instead the product obtained is $BrMgOCH_2CH_2CCH_3$.

40. The resonance structure of

- **41.** The chlorination of ethyl benzene in the presence of UV light gives 1-chloro-1-phenylethane as the major product.
- Acetaldehyde is more reactive than acetone in nucleophilic substitution.

In case of the 1° alkyl halide, the sodium alkynide acts as a nucleophile. However in case of the 2° or 3° alkyl halide, the sodium alkynide acts as a base.

Grignard reagents do not react with carbonyl compounds containing a hydroxyl group, since the OH group is much more reactive than the CO group.

Methanol cannot be represented by a resonance structure since the carbon atom has five bonds.

In ethyl benzene, the 1° radical is more stable and formed more readily than the 2° radical.

The positive charge on the carbonyl carbon in a ketone is more stable than that on the carbonyl carbon in an aldehyde. This is so because a ketone has two methyl groups and an aldehyde only one.

43. Thioanisole reacts with butyllithuim as follows.

$$SCH_3 + C_4H_9$$
 $Li^+ \rightarrow$
 $SCH_2^- Li^+ + C_4H_{10}$

However anisole does not undergo an analogous reaction.

- **44.** Phenol is less acidic than 4-methylphenol.
- **45.** *o*-Hydroxybenzaldehyde can be separated from *p*-hydroxybenzaldehyde by steam distillation. *p*-Hydroxybenzaldehyde is obtained in the steam distillate.
- **46.** On oxidation with mild oxidizing agents (e.g., atmospheric oxygen) thiols give disulphides.
- 47. The reaction of C₆H₅CO₂C₂H₅ and CH₃SO₂C₆H₅ in the presence of C₂H₅ONa gives C₆H₅COCH₂SO₂C₆H₅.
- **48.** In general, N–H…N bonds are stronger than O–H…O and F–H…F bonds.
- **49.** In cyclohexane, the boat conformation is more stable than the chair form.
- **50.** *cis-syn*. ditertiary butylethylene is stable compared to the trans form.
- **51.** The addition of bromine to an alkene gives a trans addition product.

The H atoms on C atoms that are adjacent to an alkylthio group are more acidic than those adjacent to an alkoxy group. So S atoms are easily polarized, and this can stabilize a negative charge on an adjacent atom.

The presence of an electronreleasing group in phenol makes it less acidic.

o-Hydroxyaldehydes are steam volatile due to intramolecular H bonding.

The S–H bond is stronger than the O—H bond.

In sulphones the hydrogens alpha to sulphur are acidic and undergo Claisen-type condensations with esters.

The electronegativity of nitrogen is less than that of oxygen or fluorine.

The instability of the boat form (in cyclohexane) relative to the chair form may be ascribed to relatively unfavourable interactions between the H atoms around the ring.

In *syn*. ditertiary butylethylene there is considerable steric hindrance between the bulky substituents.

The bromide ion attacks the carbon atom on the side opposite to the bridging group (viz., the bromonium ion).

52. Of CH_2O , CH_3CHO , CH_3COCH_3 and $(CH_3)_3C$ —C— $C(CH_3)_3$ the

most unreactive carbonyl compound in an addition reaction is

$$(CH_3)_3C$$
— C — $C(CH_3)_3$.

- 53. The reactivity of cyclic ketones (e.g., cyclopentanone) is much less than that of open-chain analogues (e.g., 3-hexanone) during addition reactions.
- **54.** The nitration of benzoic acid gives *o* and *p*-nitrobenzoic acid.
- 55. The nitration of 2-methoxy acetanilide gives mainly the 4-nitro derivative.
- 56. Simple aryl halides (e.g., C₆H₅Cl) are inert to nucleophilic reagents, but 2,4-dinitrochlorobenzene reacts with (CH₃)₂NH at room temperature to give

The reactivity of the carbonyl compound is greatly influenced by the bulkiness of the substituents.

In open-chain ketones, there is freedom of rotation of the groups attached to a C=O group. This causes greater steric hindrance in transition states during the addition process.

The COOH group in benzene deactivates the *o*- and *p*-positions more than the *m*-position.

If a benzene derivative contains OCH₃ and NHCOCH₃ (both being *o*- and *p*-directing), then NHCOCH₃ exerts a stronger influence than OCH₃.

The presence of strongly electron-attracting substituents (in *o-* and *p-*positions or both), activates the chloro group in 2,4-dinitrochlorobenzene.

Answers

1. d	2. d	3. a	4. b	5. d
6. c	7. a	8. d	9. a	10. d
11. a	12. d	13. a	14. d	15. c
16. d	17. a	18. d	19. a	20. d
21. a	22. d	23. d	24. a	25. a
26. d	27. d	28. b	29. a	30. a
31. a	32. a	33. b	34. b	35. a
36. a	37. d	38. a	39. a	40. d
41. c	42. a	43. a	44. d	45. d
46. c	47. a	48. d	49. d	50. d
51. a	52. a	53. d	54. d	55. a
56. a				

Part 4

Miscellaneous Questions

1

Physical Chemistry Miscellaneous Questions

• Type 1 •

Choose the correct option. Only one option is correct.

1. In the cell

$$Zn(s) | Zn^{2+}(aq, 0.10 M) | KCl(sat'd) | Zn^{2+}(aq, 1.0 M) | Zn(s),$$

- (a) E° is not zero because the concentrations of the solutions in the left and right compartments are unequal
- (b) E° is zero because the same electrode and the same type of ions are involved
- (c) E° is zero because the same electrode and the same type of ions are involved
- (d) E° is not zero because the same electrode and different concentrations of ions are involved
- 2. 25 mL of $\frac{1}{10}$ N HCl will exactly neutralize (given that the atomic weight of boron is 10.8)

 - (a) $0.2728 \text{ g of Na}_2B_4O_7\cdot 10H_2O$ (b) $0.4768 \text{ g of Na}_2B_4O_7\cdot 10H_2O$

 - (c) $0.0604 \text{ g of Na}_2B_4O_7\cdot 10H_2O$ (d) $0.8006 \text{ g of Na}_2B_4O_7\cdot 10H_2O$
- 3. The reaction between NO₂ and CO to produce NO and CO₂ is believed to occur in the following two steps.
 - Step 1:

$$2NO_2 \longrightarrow NO + NO_3$$

Step 2:

$$NO_3 + CO \longrightarrow NO + CO_2$$

The intermediate product and the rate-determining step are

- (a) NO and Step 1
- (b) NO and Step 2
- (c) NO₃ and Step 1
- (d) NO₃ and Step 2

- An orbial is
 - (a) a surface of constant probability density
 - (b) a one-electron spatial wave function
 - (c) a volume element in spherical polar coordinates
 - (d) a hydrogen atom in the ground state, the electron being confined to move within a sphere of fixed radius
- 5. Calculate the pH of a saturated 0.10-M H₂S solution (given that $K_1(H_2S) = 1 \times 10^{-7} \text{ mol } L^{-1}, K_2(H_2S^{-}) = 1 \times 10^{-14} \text{ mol } L^{-1}).$
 - (a) 7.0
- (b) 1.0
- (c) 3.0
- (d) 4.0
- 6. Which of the following equations represents Boyle's law?
 - (a) $\frac{dp}{v} = -\frac{dV}{V+n}$

- (b) $\frac{dp}{p} = -\frac{dV}{V}$
- (c) $\frac{dp}{p+V} = -\frac{dV}{V+p}$
- (d) $\frac{dp}{n} = \frac{dV}{V}$
- 7. Which of the following nuclear processes are identical?
 - (a) β^+ emission and β^- emission
 - (b) α emission and γ radiation
 - (c) β + emission and electron capture
 - (d) γ radiation and production of X-rays
- 8. Among the following, which is the most acidic?
 - (a) IO(OH)₅

(b) B(OH)₃

(c) CH₃CO₂H

- (d) HOCl
- 9. A d shell containing 5 electrons of parallel spin can exchange
 - (a) 10 electrons

(b) 6 electrons

(c) 5 electrons

- (d) 25 electrons
- **10.** Which of the following is an efficient catalytic converter?
 - A reagent which oxidizes CO and unburnt hydrocarbons to CO₂ and H_2O , and reduces NO and NO_2 to N_2 and O_2 .
 - (b) A reagent which oxidizes CO and NO to CO₂ and N₂ respectively.
 - A reagent which oxidizes CO and unburnt PbO to CO_2 and Pb O_2 , and reduces SO_2 to H_2S .
 - (d) A reagent which reduces unburnt hydrocarbons to CO, CO2 and H_2O and oxidizes N_2 to NO and NO_2 .
- 11. Which of the following is arranged in order of increasing strength of conjugate base?

 - (a) $CN^- < NH_3 < NH_2^- < OH^-$ (b) $NH_2^- < OH^- < CN^- < NH_3$
 - (c) $OH^- < NH_3 < NH_2^- < CN^-$ (d) $NH_3 < CN^- < OH^- < NH_2^-$

12. In the reaction

$$CaCO_3(s) + heat \rightleftharpoons CaO(s) + CO_2(g)$$
,

reducing the volume of the system will

- (a) shift the equilbrium to the left
- (b) shift the equilibrium to the right
- (c) reduce the particle pressure of the gas
- (d) increase the temperature of equilibrium
- **13.** Bulbs A and B contain an aqueous solution and pure water respectively. The bulbs are connected, and dry air is passed through them for some time. The air passes first through Bulb A and then through Bulb B. If the vapour pressure of water is p_1° and that of the solution is p_1 , the weight losses of the bulbs containing the solution and the solvent are directly proportional to
 - (a) p_1 and p_1°

(b) p_1 and $p_1 - p_1^{\circ}$

(c) p_1 and $p_1^{\circ} - p_1$

- (d) $p_1^{\circ} + p \text{ and } p_1^{\circ} p$
- **14.** What is the unit of *Q* (reaction quotient)?
 - (a) $(\text{mol } L^{-1})^{\Delta n}$
 - (b) $(L atm)^{\Delta n}$
 - (c) It is pressure- and temperature-dependent.
 - (d) Q does not have a unit as it is dimensionless.
- **15.** A compound AB crystallizes in the cubic-close-packed zinc blende structure. Assuming that B ions occupy the lattice points, what fraction of the tetrahedral sites is occupied by A ions?
 - (a) 25%

(b) 50%

(c) 75%

- (d) 33%
- **16.** A gas undergoes constant-temperature expansion from 250 mL to 750 mL. What is the work done by the gas if it expands against a vacuum?
 - (a) $RT \ln \frac{750}{250}$

(b) $RT \ln \frac{250}{750}$

(c) 0 J

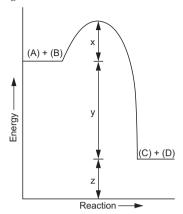
- (d) $nRT \ln \frac{750}{250}$
- 17. Which of the following bonds is resistant to oxidizing agents like Cl₂ as well as to strong nucleophiles like OH⁻?
 - (a) C—Br

(b) C—I

(c) C—F

(d) C-O

18. The enthalpy change of the reaction (A) + (B) \rightarrow (C) + (D) is y, as shown in the following figure.



The activation energy of the reaction $(C) + (D) \rightarrow (A) + (B)$ is represented by

(a)
$$x + y + z$$

(b)
$$x + y$$

(a)
$$x + y + z$$
 (b) $x + y$ (c) $x + y - z$ (d) $x - y + z$

(d)
$$x - y + 2$$

19. Determine the S²⁻ concentration in a saturated H₂S solution to which enough HCl has been added to obtain an H $^+$ concentration of 1×10^{-3} mol L $^{-1}$ (given that $K_1(\text{H}_2\text{S}) = 1.0 \times 10^{-7}$ mol L $^{-1}$ and $K_2(\text{HS}^-) = 1.0 \times 10^{-14}$ mol L $^{-1}$).

(a)
$$1 \times 10^{-14} \text{ mol L}^{-1}$$

(b)
$$1 \times 10^{-16} \text{ mol L}^{-1}$$

(c)
$$1 \times 10^{-10} \text{ mol L}^{-1}$$

(d)
$$1 \times 10^{-8} \text{ mol L}^{-1}$$

20. 40.0 mL of a 0.03-M KMnO₄ solution will completely oxidize

- (a) 30 mL of 0.15 M oxalic acid
- (b) 20 mL of a 0.20-M ferrous oxalate solution
- (c) 20 mL of a 0.10-M ferrous oxalate solution
- (d) 25 mL of a 0.20-M Mohr salt solution

21. The coefficient of expansion for an ideal gas is given by $\alpha = \frac{1}{V} \left. \frac{dV}{dT} \right|_{T}$

Using the ideal gas equation pV = nRT, we get

(a)
$$\alpha = \frac{1}{V}$$

(b)
$$\alpha = \frac{1}{V} \frac{dV}{dv}\Big|_{T}$$

(c)
$$\alpha = \frac{1}{p}$$

(d)
$$\alpha = \frac{1}{T}$$

22. A galvanic cell is set up according to the following specifications: $Zn(s) | Zn^{2+}(aq, 1 M) | KCl(sat'd) | Cu^{2+}(aq, 1 M) | Cu(s).$

Choose the correct statement.

- (a) During the overall redox reaction, electrons flow externally from the anode (Zn electrode) through the wire and voltmeter to the cathode (Cu electrode).
- (b) In the solution, the cations $(Zn^{2+}, Cu^{2+} \text{ and } K^{+})$ move towards the anode, while the anions (SO₄²⁻ and Cl⁻) move towards the cathode.
- (c) K⁺ and Cl⁻ do not move to either electrode.
- (d) The conductance of the saturated KCl solution is very low. So a KCl salt bridge is used.
- 23. The following data are given:

 - (i) $H_2(g) \longrightarrow 2H(g)$; $\Delta H^{\circ} = 436 \text{ kJ}$ (ii) $Br_2(g) \longrightarrow 2Br(g)$; $\Delta H^{\circ} = 192 \text{ kJ}$
 - (iii) $H_2(g) + Br_2(g) \longrightarrow 2HBr(g)$; $\Delta H^{\circ} = -104 \text{ kJ}$

Then ΔH° for the reaction

$$H(g) + Br(g) \longrightarrow HBr(g)$$

is

(a) -376 kI mol⁻¹

(b) $+366 \text{ kJ mol}^{-1}$

(c) -366 kJ mol⁻¹

- (d) -327 kJ mol^{-1}
- **24.** The c/a ratio for the hexagonal close packing of a sphere is
 - (a) 1.363
- (b) 1.633
- (c) 1.732
- 25. An aqueous solution of $Mg(NO_3)_2$ is electrolysed. What are the gaseous products formed?
 - (a) O_2 at the anode and N_2 at the cathode
 - (b) N_2 at the anode and H_2 at the cathode
 - (c) O_2 at the anode and H_2 at the cathode
 - (d) N₂ at the anode and Mg at the cathode
- 26. Which of the following is arranged in the increasing order of acid strength?
 - (a) $HSO_3F < H_3O^+ < HSO_4^- < NH_3$
 - (b) $H_3O^+ < HSO_3F < NH_3 < HSO_4^-$
 - (c) $NH_3 < HSO_4^- < H_3O^+ < HSO_3F$
 - (d) $HSO_4^- < NH_3 < HSO_3F < H_3O^+$
- 27. How much ethylene glycol (CH₂OH—CH₂OH) must be added to 1.0 L of water so that the solution will not freeze at -20° C? (K_f for H_2 O = 1.86 K kg mol⁻¹)
 - (a) 66 g
- (b) 667 g
- (c) 1667 g (d) 333 g

28.

$$H_2CO_3 \rightleftharpoons H^+ + HCO_3^-$$

 $HCO_3^- \rightleftharpoons H^+ + CO_3^-$
 $H_2O \rightleftharpoons H^+ + OH^-$

For the above equations, which of the following is correct on the basis of the concept of electroneutrality?

- (a) $3[H^+] = 2[HCO_3^-] + [CO_3^{2-}] + [OH^-]$
- (b) $[H^+] = [HCO_3^-] + 2[CO_3^{2-}] + [OH^-]$
- (c) $[H^+] = [HCO_3^-] + [CO_3^{2-}] + [OH^-]$
- (d) $3[H^+] = [HCO_3^-] + 2[CO_3^{2-}] + [OH^-]$
- **29.** The emf of a cell is
 - (a) dependent on the volume of the solution and the size of the
 - (b) independent of the volume of the solution and the size of the electrodes
 - (c) independent of the volume of the solution and depends on the size of the electrodes
 - (d) dependent on the volume of the solution and independent of the size of the electrodes
- **30.** A possible mechanism for the reaction

$$2NO + 2H_2 \longrightarrow N_2 + 2H_2O$$

- (i) $2NO \rightleftharpoons N_2O_2$
- (ii) $N_2O_2 + H_2 \longrightarrow N_2O + H_2O$ (slow step) (iii) $N_2O + H_2 \longrightarrow N_2 + H_2O$

The rate law for the reaction is

(a) $k[NO]^2[H_2]^{1/2}$

(b) $k[NO][H_2]^2$

(c) $k[NO]^2[H_2]$

- (d) $k[NO]^2[H_2]^2$
- 31. What is the concentration of the acetic acid which is added to 0.5 N HCOOH so that the percentage dissociation of both the acids is unchanged (given that $K_a(\text{CH}_3\text{COOH}) = 1.8 \times 10^{-5}$ mol L⁻¹ and $K_a(\text{HCOOH}) = 2.4 \times 10^{-4}$ mol L⁻¹)?
 - (a) 6.1 mol L^{-1}

(b) 0.62 mol L^{-1}

(c) 6.7 mol L^{-1}

- (d) 7.6 mol L^{-1}
- **32.** The diamond atom may be viewed as
 - (a) two fcc structures displaced from each other by a quarter of the body diagonal
 - (b) two fcc structures displaced from each other by half the body diagonal
 - (c) four fcc structures displaced from each other by a quarter of the body diagonal
 - (d) four fcc structures displaced from each other by three-fourths of the body diagonal

• *Type 2* •

Choose the correct options. More than one option is correct.

- **33.** Which of the following are correctly matched?
 - (a) Interstitial defect ↔ an extra atom in an interstitial site
 - (b) Schottky defect ↔ an atom missing from an expected site
 - (c) Frenkel defect ↔ an atom displaced to an interstitial site, creating a vacancy nearby
 - (d) Grain boundary defect ↔ a boundary between two crystals in a crystalline solid
- **34.** Which of the following statements correspond to the Hund rule?
 - (a) The state of maximum multiplicity is the lowest.
 - (b) The electrons occupy equivalent orbitals singly, as far as possible, with parallel spin.
 - (c) For a given multiplicity, the state of minimum *L* is the lowest.
 - (d) The state of maximum multiplicity is given by 2s + 1 or n + 1, where s = total spin and n = number of singly occupied orbitals.
- **35.** Which of the following statements are correct?
 - (a) In case of diffusion, molecular flux is directly proportional to molecular speed.
 - (b) In case of effusion, molecular flux is inversely proportional to molecular speed.
 - (c) For both diffusion and effusion, ΔG° tends to decrease.
 - (d) Diffusion involves the effect of collisions between molecules, whereas effusion does not.
- **36.** Which of the following statements are correct?
 - (a) All Brønsted bases are Lewis bases.
 - (b) A coordinate covalent bond is always formed in a Lewis-base reaction.
 - (c) All Lewis acids are Brønsted acids.
 - (d) All amphoteric hydroxides are insoluble in water.
- **37.** Two ideal gases have the same initial pressure, volume and temperature. They expand to the same final volume, one adiabatically and the other isothermally. Find the correct statements.
 - (a) The final pressure is greater for the isothermal process.
 - (b) The work done by the gas is greater for the isothermal process.
 - (c) The final temperature is greater for the isothermal process.
 - (d) All these statements are incorrect.

- **38.** Which of the following statements are correct?
 - (a) According to the collision theory, a reaction occurs when molecules collide with energy sufficient to break the bonds and initiate the reaction. This energy is called activation energy.
 - (b) The overall balanced equation for a reaction may be the sum of a series of simple reactions called elementary steps.
 - (c) Enzymes are catalysts occuring in nonliving systems.
 - (d) A catalyst lowers the activation energy for the forward as well as the reverse reaction.
- **39.** Which of the following statements are correct?
 - (a) Covalent character increases with decreasing cation size.
 - (b) Covalent character increases with increase in anion size.
 - (c) Cations with a non-noble-gas configuration have a greater covalent character than those of the same size with a noble-gas configuration.
 - (d) Cations with a noble-gas configuration have a greater covalent character than those of the same size with a non-noble-gas configuration.
- **40.** Which of the following reactions represent hydrolysis and not dissociation?
 - (a) $HCO_3^- + H_2O \rightleftharpoons OH^- + H_2CO_3$
 - (b) $HPO_4^{2-} + H_2O \rightleftharpoons H_2PO_4^{-} + OH^{-}$
 - (c) $H_2Gly^+ + H_2O \rightleftharpoons HGly + H_3O^+$
 - (d) $H_2PO_4^- + H_2O \rightleftharpoons HPO_4^{2-} + H_3O^+$

Answers1. b 2. b **3.** c 4. b **5**. d 6. b 7. c 8. a **9.** a 10. a 15. b **11.** d **12.** a 13. c **14.** d 16. c **17.** c 18. b 19. b **20.** c 21. d 22. a 23. c 24. b 25. c 26. c **27.** a 28. b 29. h **30.** c 31. c **32.** a 33. a, b, c **34.** a. b. d 35. a, b, c, d **36.** a, b, d **37.** a, b, c **38.** a, b, d **39.** a, b, c **40.** a, b, c

Hints to More-Difficult Problems

- **3.** Here NO₃ is an intermediate product because it is cancelled in Step 2, and so Step 1 is the rate-determining step.
- The pH of H₂S is calculated by considering the first step of dissociation of H₂S.

$$K_1 = \frac{[H^+][HS^-]}{[H_2S]} = \frac{[H^+]^2}{[H_2S]}$$
 (: [H^+] = [HS^-]).

$$\therefore 10^{-7} = \frac{[H]^{+2}}{0.1} \Rightarrow [H^+]^2 = 10^{-8} \Rightarrow pH = 4.$$

7. Positron emission: ${}^{A}_{7}X \longrightarrow {}^{A}_{7+1}Y + {}^{0}_{+1}e$

Electron capture: ${}_{Z}^{A}X + {}_{+1}^{0}e \longrightarrow {}_{Z+1}^{A}Y$

In both the processes, we obtain the identical product $_{Z+1}^{A}Y$.

- 9. ${}^{5}C_{2} = \frac{5!}{3! \ 2!} = \frac{120}{6 \times 2} = 10.$
- **10.** The conjugate acid of NH₂⁻ is weakly basic.
- 12. Apply Le Chatelier's principle.
- **13.** The loss of weight in bulb A is proportional to the vapour pressure of the solution (p_1) . The loss of weight in bulb B is proportional to the vapour pressure of the solvent $(p_1^\circ p_1)$.
- **14.** In the expression of Q, all activity terms are taken at unit pressure ($p^{\circ} = 1 \text{ bar}$) or unit concentration ($c^{\circ} = 1 \text{ mol L}^{-1}$).
- 21. We know that

$$pV = nRT$$
.

Differentiating with respect to T, we get

$$p \frac{\partial}{\partial T} \bigg]_p = nR.$$

We can work out α from this.

- 22. The process is the reverse of that in the electrolytic cell.
- **23.** Performing (iii) (i) (ii), we obtain

$$\Delta H^{\circ} = -104 \text{ kJ} - 436 \text{ kJ} - 192 \text{ kJ} = -732 \text{ kJ}$$

for the reaction 2H(g) + 2Br(g) = 2HBr(g).

 $\therefore \quad \text{for the reaction } H(g) + Br(g) = HBr,$

$$\Delta H^{\circ} = \frac{-732 \text{ kJ mol}^{-1}}{2} = -366 \text{ kJ mol}^{-1}.$$

25. Water is electrolysed.

- **30.** The slow step is the rate-determining step. So the rate of reaction is given by step (ii), i.e., then the rate law is $k[NO]^2[H_2]$.
- 31. According to the principle of isohydric solutions,

$$\frac{K_1}{V_1} = \frac{K_2}{V_2} \cdot$$

Here, V_1 is the volume of HCOOH, V_2 is the volume of CH₃COOH, K_1 is the dissociation constant of HCOOH and K_2 is the dissociation constant of CH₃COOH.

$$\begin{split} [H^+]^2 &= K_2 [HCOOH] = 2.4 \times 10^{-4} \text{ mol } L^{-1} \times 0.5 \text{ mol } L^{-1} \\ &= 1.2 \times 10^{-4} \text{ (mol } L^{-1})^2. \\ [CH_3COOH] &= \frac{[H^+]^2}{K_1} = \frac{1.2 \times 10^{-4} \text{ (mol } L^{-1})^2}{1.8 \times 10^{-5} \text{ mol } L^{-1}} = 6.7 \text{ mol } L^{-1}. \end{split}$$

40. In hydrolysis, an acid and a base are produced.

Inorganic Chemistry Miscellaneous Questions

• Type 1 •

1. Which of the following alkali metals forms a peroxide in preference to a

(a) large cations stabilize large unstable anions in CsI₃ (b) CsI₃ has a bcc structure and NaI₃ an fcc structure

(d) large cations stabilize small unstable anions in CsI₃

(c) Na

(c) CsI₃ has a low hydration energy and NaI₃ a high hydration energy

3. The temperatures at which ¹H₂O and D₂O have maximum density,

(d) Cs

Choose the correct option. Only one option is correct.

2. $CsI_3(s)$ is stable but $NaI_3(s)$ is not because

(b) Rb

superoxide?

(a) K

respectively, are

	(a)	4°C and 11.6°C	(b)	11.6°C and 4°C
	(c)	4°C and 12.5°C	(d)	12.5°C and 4°C
4.	Amor	ng the following, which is the s	trong	gest reducing agent?
	(a)	$[BH_4]^-$	(b)	$[AlH_4]^-$
	(c)	$[GaH_4]^-$	(d)	$[AlF_6]^{3-}$
5.	Which	n of the following is arranged in	n ord	der of increasing acidity?
	(a)	$N_2H_5^+ < NH_3OH^+ < NH_4^+$	(b)	$NH_3OH^+ < NH_4^+ < N_2H_5^+$
	(c)	$NH_4^+ < N_2H_5^+ < NH_3OH^+$	(d)	$NH_3OH^+ < N_2H_5^+ < NH_4^+$
6.	Fluori	ne has a lower electron affinity	tha:	n chlorine because
	(a)	there is greater electron–elect the larger Cl atom	ron	attraction in the F atom than in
	(b)	the standard reduction poten $Cl_2 \mid Cl^-$ is +1.36 V	ntial	of $F_2 \mid F^-$ is +2.87 V and that of
		1 13		

- (c) there is greater electron–electron repulsion in the F atom than in the larger Cl atom
- (d) fluorine has an oxidation number of −1, and chlorine has a maximum oxidation number of +7
- 7. The action of sodium vapour on molten KCl at 850°C gives
 - (a) Cl₂

- (b) K
- (c) solvated electrons
- (d) solvated sodium ions
- 8. In the solid compound $K[Cu(CN)_2]$,
 - (a) two CN groups are bound through C, and the third (bridging) CN group through C and N
 - (b) three CN^- groups are bound through C, and the bridging CN^- group through C and N
 - (c) one CN group is bound through C, and the other through C and N
 - (d) one CN^- group is bound through C, and two other CN^- groups through C and N
- 9. Which of the following statements is incorrect?
 - (a) LiF is much less soluble than LiCl in water.
 - (b) AgF is much more soluble than AgCl in water.
 - (c) The structures of ZnO and HgO are quite different.
 - (d) The structures of HgF₂ and HgCl₂ are identical.
- 10. Which of the following statements is correct?
 - (a) In a semiconductor, the energy gap between the filled and the empty bands is much greater than in an insulator.
 - (b) In a metal, the energy gap between the conduction band and the valence band is greater than in an insulator.
 - (c) In an insulator, the valence band and the conduction band overlap.
 - (d) In an insulator, the energy gap between the valence band and the conduction band is considerably greater than in a metal.
- 11. Among the following molecules, which has the maximum dipole moment?
 - (a) O_3
- (b) H₂O₂
- (c) N_2H_4
- (d) H₂O
- 12. The nuclear spins of protium, deuterium and helium-4 are respectively
 - (a) $\frac{1}{2}$, 0, 2

(b) $1, \frac{1}{2}, \frac{1}{2}$

(c) $\frac{1}{2}$, 1, 0

(d) $\frac{1}{2}$, 1, 2

(d) NaH(s)

13. Among the following hydrides, which is thermodynamically the most

14. Which of the following reactions is used to prepare amorphous silicon?

(c) $B_2H_6(g)$

stable at room temperature?
(a) BeH₂(s) (b) LiH(s)

(a) $Si_2H_6 \xrightarrow{400^{\circ}C}$ (P)

(b) $SiH_4 \xrightarrow{500^{\circ}C} (P)$

	(c)	SiH ₄ — electric disc	$\xrightarrow{\text{charge}}$ (P)				
	(d)	R ₃ SiCl + LiAlH ₄	\longrightarrow (P)				
15.		n of the following		lucin	g agent?		
		SO_4^{2-} (b)			-	(d) S ₂	O ₆ ²⁻
16.		n of the following					- 0
20.	(a)	•	Na	(c)		(d) C	s
17.	Which	n of the following	statements i	s cori	rect?		
	(a)	The production conversion of it which the impu	iron to steel	esse	ntially an ox	idation	process in
	(b)	The production processes.	n of iron an	d th	at of steel a	re bot	h reduction
	(c)	The production are oxidized, and is reduced to ca	d that of stee				
	(d)	The production processes.	n of iron an	d th	at of steel a	re bot	h oxidation
18.	In [Fe	$(CN)_6]^{3-}$, the d ele	ectrons occup	y the	9		
	(a)	d_{xy} , d_{yz} , d_{zx} and	d_{z^2} orbitals				
	(b)	d_{xy} , d_{yz} , and d_{zx}	orbitals				
	(c)	$d_{x^2-y^2}$ and d_{z^2} or	rbitals				
	(d)	d_{xy} , d_{yz} , d_{zx} , d_{x^2}	$-y^2$ and d_{z^2} orl	bitals			
19.	The co	omplex ion [Ni(C	$(2N)_2Br_2]^{2-}$ has	s a			
	(a)	square-planar g	eometry	(b)	tetrahedral g	geomet	ry
	(c)	square-pyramic	lal geometry	(d)	pyramidal g	eometr	y
20.		an excellent fuel	_		ecause of its		
		low specific ent			high specific		
	(c)	high bond energ	зу	(d)	low electron	affinit	y

21.	21. Among the following reactions, which gives the highest proportion of ¹ HD?			
	(a)	a) ${}^{1}\text{H}_{2} + \text{D}_{2}$ equilibrated over a platinum surface		
	(b)	The reaction between D ₂ O and	d Na	Н
	(c)	The reaction between D ₂ O and	d NI	H_3
	(d)	The electrolysis of ¹ HDO		
22.	The h	ydrolysis of Li ₃ N produces		
	(a)	HN_3	(b)	N_2H_4
	(c)	NH ₂ OH	(d)	NH_3
23.		n of the following is arrang tion in air?	ed i	n order of increasing ease of
	(a)	$NO < N_2O < NO_2$	(b)	$N_2O < NO_2 < NO$
	(c)	$NO_2 < NO < N_2O$	(d)	$NO_2 < N_2O < NO$
24.	Which	n of the following solid fluoride	es is	unstable?
	(a)	LiF	(b)	CuF
	(c)	BeF ₂	(d)	HgF_2
25.	Lithiu	ım is a highly active reducing a	igent	t because of its
	(a)	low ionization energy		
	(b)	low sublimation energy		
	(c)	strong metallic bonding		
	(d)	very high hydration energy		
26.		O_2 , the HOO angle is only 97° of in H_2O because the	comp	pared to the HOH bond angle of
	(a)	lone pair-bond pair repulsion	ı is g	reater in H ₂ O ₂ than in H ₂ O

(b) lone pair-bond pair repulsion is greater in H₂O than in H₂O₂
(c) bond pair-bond pair repulsion is greater in H₂O₂ than in H₂O
(d) bond pair-bond pair repulsion is greater in H₂O than in H₂O₂

27. Which of the following pairs give the same gaseous products upon

(b) S_4N_4 and Li_3N

(d) Na₂O₂ and KO₂

treatment with water?

(a) NaN₃ and Li₃N

(c) CH_2N_2 and $(NH_4)_2Cr_2O_7$

• Type 2 •

Choose the correct options. More than one option is correct.

- 28. Which of the following hydrated cations are colourless and diamagnetic?
 - (a) $Zn^{2+}(aq)$ (b) $Ti^{3+}(aq)$ (c) $Cu^{2+}(aq)$ (d) $Sc^{3+}(aq)$

- 29. Nitrogen reacts slowly with most compounds because of the
 - (a) high strength of the N≡N bond
 - (b) high activation energy of N₂
 - (c) low polarizability of nitrogen
 - (d) high polarizability of nitrogen
- **30.** Which of the following sets of species are isoelectronic?
 - (a) N_2 , NO^+ , CN^-
- (b) O_2^+ , NO, N₂O
- (c) O_2^{2-} , Cl_2 , $N_2H_5^+$
- (d) NO, NO₂⁺, N₂O
- **31.** Which of the following statements are correct?
 - (a) ClO₂ is an angular radical.
 - (b) The I_2O_5 molecule has an oxo-bridge between two IO_2 structures.
 - (c) HBrO₄ is more acidic but less stable than H₅IO₆.
 - (d) In XeF₇, there are six electron pairs around Xe, and one is a lone pair.
- **32.** Which of the following statements are correct?
 - (a) Xenon form a range of compounds with fluorine and oxygen.
 - (b) Helium 4 is radioactive.
 - (c) Noble gases have positive electron affinities because their valence shells are full and an incoming electron occupies an orbital of a new shell.
 - (d) Noble gases have negative electron affinities because their valence shells are full and an incoming electron occupies an orbital of a new shell.
- 33. Which of the following statements are correct?
 - (a) A metal is an array of positive ions immersed in a sea of delocalized valence electrons.
 - (b) The strength of a metallic bond is due to the cohesive force resulting from delocalized electrons.
 - (c) The valence band and the conduction band in a metal are adjacent to each other.
 - (d) An electron can travel freely through a metal since the conduction band is devoid of electrons.

- **34.** Which of the following statements are correct?
 - (a) In Al₂Cl₆, each aluminium atom is sp ³-hybridized.
 - (b) In Al₂Cl₆, each of the bridging chlorine atoms forms a normal covalent bond with one aluminium atom and a coordinate covalent bond with another.
 - (c) The reaction of $Fe_2O_3(s)$ with Al(s) is endothermic.
 - (d) Alums are represented by the general formula $M^+M^{3+}(SO_4)_2\cdot 12H_2O$.
- 35. The F-F bond is weak due to the
 - (a) strong repulsions between nonbonding electrons in the small F₂ molecule
 - (b) high standard reduction potential of fluorine
 - (c) presence of more electrons in the bonding molecular orbitals than in the antibonding molecular orbitals of F_2
 - (d) presence of more electrons in the antibonding molecular orbitals than in the bonding molecular orbitals of F₂
- **36.** Which of the following are correctly matched?
 - (a) Barium hydride ↔ covalent hydride, molecular
 - (b) Silane ↔ covalent hydride, molecular
 - (c) Palladium hydride ↔ saline
 - (d) Arsane ↔ electron-rich, molecular
- 37. Which of the following can be prepared from PCl₅?
 - (a) POCl₃

(b) H₃PO₄

(c) [PCl₄] [AlCl₄]

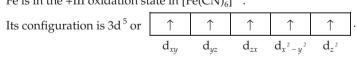
(d) $N_3P_3Cl_6$

1. c	2. a	3. a	4. b	5. c
6. c	7. b	8. a	9. d	10. d
11. b	12. c	13. b	14. c	15. b
16. a	17. a	18. b	19. a	20. b
21. b	22. d	23. d	24. b	25. d
26. a	27. d	28. a, d	29. a, b, c	30. a, c
31. a, b, c	32. a, c	33. a, b, c, d	34. a, b, d	35. a, d
36. b, d	37. a, b, c, d			

Hints to More-Difficult Problems

- The least electropositive of these metals will polarize oxygen anions the least.
- **4.** Among these anion complexes, [AlH₄] is the strongest hydride donor.
- 7. Sodium is a more powerful reducing agent than potassium.
- **8.** A portion of the spiral chain in $K[Cu(CN)_2]$ is

- 9. HgF_2 has a fluorite structure whereas mercury(II) chloride has a molecular lattice consisting of discrete linear $HgCl_2$ molecules.
- 11. H₂O₂ is more polar than the rest of the molecules shown. This can be vectorically worked out from their structures.
- **12.** (i) Nuclei with *p* and *n* even (charge and mass even) have zero spin (e.g., helium-4).
 - (ii) Nuclei with *p* and *n* odd (charge odd and mass even) have integral spin (e.g., deuterium).
 - (iii) Nuclei with odd mass have half-integral spin (e.g., hydrogen).
- 16. Polarization effect
- **18.** Fe is in the +III oxidation state in $[Fe(CN)_6]^{3-}$.



 $\rm CN^-$ is a strong ligand. So the electrons get paired and occupy the $\rm d_{\it xy\prime}$ and $\rm d_{\it zx}$ orbitals in $\rm [Fe(CN)_6]^{3-}$.

$\uparrow\downarrow$	$\uparrow\downarrow$	↑	
d_{xy}	d_{yx}	d_{zx}	

- **22.** NH_3 is produced due to the protonation of the nitride ion, a Brønsted base.
- **24.** Solid CuF disproportionates, forming Cu and CuF_2 .

- **28.** Zn ²⁺(aq) has a d ¹⁰ electronic configuration and so cannot produce d–d spectra. Thus many of its compounds are colourless and diamagnetic. Sc ³⁺(aq) has a d ⁰ configuration and cannot produce d–d spectra either. Therefore, these ions and their compounds are also colourless and diamagnetic.
- 31.



is an angular radical.

The structure of I₂O₅ is

Again,

$$HBrO_4 \longrightarrow H^+ + BrO_4^-$$
 (acidic).

 $H_5IO_6(HIO_4:2H_2O)$ is known as paraperiodic acid. It is stable because of its tetrahedral constituent IO_4^- .

34. Potash alum $[K_2SO_4\cdot Al_2(SO_4)_3\cdot 24H_2O]$, chrome alum $[K_2SO_4\cdot Cr_2(SO_4)_3\cdot 24H_2O]$ and ferric ammonium alum $[(NH_4)_2SO_4\cdot Fe_2(SO_4)_3\cdot 24H_2O]$ may be represented by the general formula $M^+M^{3+}(SO_4)_2\cdot 12H_2O$.

Structure of Al₂Cl₆

Organic Chemistry Miscellaneous Questions

• *Type 1* •

Choose the correct option. Only one option is correct.

1. With change in the hybridization of the carbon bearing the charge, the stability of a carbanion decreases in the order

(a)
$$sp < sp^2 < sp^3$$

(b)
$$sp^3 < sp^2 < sp$$

(c)
$$sp < sp^3 < sp^2$$

(d)
$$sp^2 < sp < sp^3$$

2. The order of stability of alkyl, vinyl and alkynyl carbanions is

(a)
$$RCH_2 > RCH = CH > RC = C$$
 (b) $RCH = CH > RC = C > RCH_2$

(b)
$$RCH = \overline{CH} > RC = \overline{C} > RCH_2$$

(c)
$$RC = \overline{C} > RCH = \overline{CH} > RC\overline{H}_2$$

(c)
$$RC = \overline{C} > RCH = \overline{CH} > RCH_2$$
 (d) $RCH_2 > RCH = \overline{CH} > RC = \overline{C}$

3. Which of the following alkenes is the most stable?

(a)
$$(CH_3)_2C=C(CH_3)_2$$

$$C = C$$

4. In the reaction of CH₃CHO with HCN in the presence of a base, the rate-determining step is

(a)
$$HCN + \overline{OH} \longrightarrow H_2O + \overline{CN}$$

(b)
$$CH_3-C=O + \bar{C}N \longrightarrow CH_3-C$$
 $\downarrow CN$
 $\downarrow CN$

(c)
$$CH_3 - C \downarrow O \longrightarrow CH_3 - C \downarrow CN$$

- (d) All the steps take place with equal ease.
- 5. Which of the following alkyl halides can be hydrolysed easily by an S_N2 reaction?
 - (a) CH₃CH₂CH₂CH₂Br
 - (b) CH₃CH₂CHCH₃ Br

- (d) All are hydrolysed with equal ease.
- 6. Which of the following alkyl halides will be the most reactive in an S_N1 reaction?
 - (a) CH₃CH₂CH₂CH₂Br
- (b) $CH_3CH_2CH-CH_3$ | Br
- (c) CH_3 CH_3CH_2 —C—Br CH_3
- (d) All are equally reactive.

7. The rate of the conversion

$$C_6H_5CH_2C1 \xrightarrow{KCN} C_6H_5CH_2CN$$

can be increased by carrying out the reaction

- (a) in an aqueous solution by vigorous stirring
- (b) in the presence of an acid catalyst
- (c) by stirring in an aqueous medium in the presence of a small amount of tetrabutyl ammonium chloride
- (d) by none of these methods
- 8. In the elimination reactions of appropriate substrates, the formation of the least-substituted alkane is explained by
 - (a) the Saytzeff rule
- (b) the Hofmann rule
- (c) both (a) and (b)
- (d) neither (a) nor (b)

9. The major product obtained in the reaction

$$C_6H_5$$
 — C — $CHCH_3$ + D_2O OD — COD products

is

(c) O
$$CH_3$$
 (d) none of these $C_6H_5-C-CD-CH_3$

- 10. In which of the following is the enol tautomer expected to be more predominant than the keto tautomer?
 - (a) 3-Pentanone

- (b) Propanone
- (c) Cyclohexanone
- (d) 2.4-Pentanedione
- 11. What is the major product obtained in the following reaction?

$$CH_3CH = CHCO_2C_2H_5 \xrightarrow{NaCN} products$$

12. What is the final product obtained in the reaction of

$$CH_3$$
 $C-OC_2H_5$

with an excess of Na in the presence of xylene (heating), followed by acidification with acetic acid?

(a)
$$CH_3 - C = O$$

 $CH_3 - C = O$

(c)
$$CH_3 - C = O$$

 $CH_3 - C - OH$

(d) None of these

- **13.** Which of the following aldehydes does not undergo a Cannizzaro reaction?
 - (a) C_6H_5 —CHO

- сно
- (c) (CH₃)₂—CH—CHO
- 14. In the molecular-orbital model of benzene, how many π electrons are delocalized about the ring?
 - (a) Three

(b) Four

(c) Five

- (d) Six
- 15. Cyclopentadiene is unusually acidic for a hydrocarbon. This is because
 - (a) the carbon atoms of cyclopentadiene are sp-hybridized
 - (b) cyclopentadiene is aromatic
 - (c) the removal of a proton from it yields an aromatic anion
 - (d) cyclopentadiene yields a highly stable free radical
- **16.** The compound C_7H_{12} decolorizes bromine in CCl_4 and reacts with Ag^+ in ammonia to form an insoluble salt. On reduction in the presence of Ni, it gives 3-methylhexane. The most likely structure of the compound is

(b)
$$CH_3CH_2CHCH_2C\equiv CH$$

 CH_3

(c)
$$CH_3CH=C-CH=CHCH_3$$

 CH_3

(d)
$$CH_3CH_2CHC \equiv C-CH_3$$

 CH_3

17. Which of the following alcohols will initially form the most stable carbocation on treatment with concentrated H₂SO₄?

(a)
$$CH_3$$
 (b) CH_3 CH_3 CH_3 CH_2 CH_2 CH_2 CH_2 CH_2 CH_2 CH_3 CH_2 CH_3 CH_3 CH_4 CH_5 CH_5

(c)
$$CH_3$$
 (d) CH_3 CH_3 CH_3 CH_3 CH_3 CH_4 CH_5 CH_5

18. The product (P) of the reaction

$$CH_3Li + D_2O \longrightarrow (P)$$

is

(a) CH₃OD

(b) CH₃—CH₃

(c) CH₃D

(d) CH₃—O—CH

19.

Which of the following is an enantiomer of the above structure?

(d) It does not have an enantiomer.

20. The product (P) obtained in the reaction

$$H_3C$$
 Cl
 H_2O
 $acetone$
 n - C_6H_{14}
 (P)

is

(a)
$$C_2H_5$$
 OH n - C_6H_{14}

HO C_2H_5 CH_3 n- C_6H_{14}

- (c) a mixture of (a) and (b)
- (d) none of these

- 21. In a nucleotide unit, the components of which of the following are sequentially linked?
 - (a) Monosaccharide-phosphate-heterocyclic base
 - (b) Amino acid-monosaccharide-phosphate
 - (c) Monosaccharide-amino acid-phosphate
 - (d) None of these
- 22. The transfer RNA anticodon for the messenger RNA codon G-C-A is
 - (a) G-C-U

(b) U-G-C

(c) C-G-U

- (d) G-U-C
- 23. The monomeric units of nucleic acids are
 - (a) D-ribose or 2-dioxy-D-ribose molecules
 - (b) nucleosides
 - (c) nucleotides
 - (d) all of these
- 24. The secondary structure of a protein is derived from
 - (a) peptide linkages
- (b) disulphide linkages
- (c) hydrogen-bond formation (d) none of these

• *Type 2* •

Choose the correct options. More than one option is correct.

- 25. Which of the following statements are correct about the electromeric effect?
 - (a) It involves permanent polarization.
 - (b) It operates through π bonds.
 - (c) It is a strong effect since the loose π electrons shift completely.
 - (d) None of these is correct.
- 26. The intermediate products of the reaction

$$\label{eq:me2CHCH2} \text{Me}_2\text{CH}-\text{CH}=\text{CH}_2 \xrightarrow{\text{HCl}} \text{Me}_2\text{CCl}-\text{CH}_3-\text{CH}_3+\text{Me}_2\text{CH}-\text{CH}-\text{CH}_3\\ \mid \text{Cl}$$

are

- (a) Me_2CH — $\overset{+}{C}H$ — CH_3 (b) $Me_2\overset{+}{C}$ — CH_2CH_3
- (c) Me_2CH — CH_2 — CH_2 (d) none of these

27. The products (P) obtained in the reaction

$$H_3C \xrightarrow{\text{NaNH}_2} \text{liquid NH}_3 \longrightarrow (P)$$

are

re
(a)
$$NH_2$$
 (b) H_3C NH_2

 H_2C

OH

- 28. Which of the following is an example of nucleophilic addition?
 - (a) Ketal formation
 - (b) Reduction with hydrogen
 - (c) Cyanohydrin formation
 - (d) Bisulphite addition

29. The rate of the reaction

is the same as that of

- (a) racemization
- (b) chlorination
- (c) protium–deuterium exchange at the alpha carbon
- (d) iodination with I₂ and OH⁻

30. Which of the following conversions are possible (• indicates a labelled carbon)?

(a)
$$O-\dot{C}H_2-CH=CH_2$$
 OH $CH_2-CH=\dot{C}H_2$

(b)
$$O-\dot{C}H_2-CH=CH_2$$
 OH CH_3 CH_3 CH_3 $CH_2-CH=CH_2$

(c)
$$O-\dot{C}H_2-CH=CH_2$$
 OH $\dot{C}H_2-CH=CH_2$

(d)
$$O-\dot{C}H_2-CH=CH_2$$
 OH CH_3 C

31. Which of the following aldehydes do not undergo the usual Cannizzaro reaction?

(c) O (d) CHO
$$C_6H_5$$
—C-CHO CHO

32. Which of the following reactions are possible?

(b)
$$C_{6}H_{5} \stackrel{|}{-} H_{5} \stackrel{|}{-} H_{5}$$

(d)
$$CH_3 CH_3$$
 CH_3 CH_3

$$CH_3 - C - C - CH_3 \xrightarrow{450^{\circ}C} CH_3 - C - C - CH_3$$

$$OH OH CH_3 CH_3 O (major product)$$

33. Which of the following have *trans* isomers?

- **34.** Which of the following statements are true for the S_N 2 reaction of (R)-2-bromobutane with a hydroxide ion?
 - (a) The reaction occurs with inversion of configuration.
 - (b) The reaction occurs with retention of configuration.
 - (c) Doubling the concentration of the hydroxide ion doubles the rate of the reaction.
 - (d) Doubling the concentration of the substrate doubles the rate of the reaction.
- 35. The hydrogen bonding for the base pairs of DNA are between
 - (a) amide carbonyl and —NH₂
 - (b) amide NH and cyclic amine nitrogens
 - (c) alcohols and carbonyls
 - (d) There is no hydrogen bonding.
- **36.** What are the products formed in the following reaction?

$$^{14}\text{CH}_2 = \text{CH} - \text{CH}_3 \xrightarrow{N-\text{bromosuccinimide}} \text{products}$$

(a)
$${}^{14}\text{CH}_2 = \text{CH} - \text{CH}_2\text{Br}$$
 (b) $\text{CH}_2 = \text{CH} - {}^{14}\text{CH}_2\text{Br}$

(c)
$$CH_2=CH-CH_2Br$$
 (d) None of these

37. Compare the compounds

and state which of the following statements are correct.

- (a) II is more effectively stabilized than I.
- (b) The α -hydrogens in I are more acidic than those in II.
- (c) The C—O bond in II is stronger than the C—S bond in I.
- (d) The C—O bond in II is weaker than the C—S bond in I.

1. a	2. c	3. a	4. b	5. a
6. c	7. c	8. b	9. c	10. d
11. d	12. c	13. d	14. d	15. c
16. b	17. a	18. c	19. d	20. c
21. d	22. c	23. c	24. c	25 . b, c
26. a, b	27. a, b	28. a, c, d	29. a, b, c, d	30. a, b
31. c, d	32. a, b, c	33. b, c, d	34. a, c, d	35. a, b
36. a, b	37. a, b, c			

Hints to More-Difficult Problems

- 2. The sp-hybridized carbon atom of ethyne polarizes its C—H bonds to the greatest extent because it is the most electronegative. This causes its hydrogens to be the most positive. Therefore, ethyne donates a proton to a base more readily than the other groups do.
- 5. An S_N 2 reaction proceeds through a five-membered transition state and such a transition state is destabilized by overcrowding at the central carbon atom.
- 7. Tetrabutyl ammonium chloride is a phase-transfer catalyst.
- 8. According to the Hofmann rule, in the elimination reaction of appropriate substrates, the least-substituted alkene is the major product.
- **9.** Under basic conditions, only the protons at carbon atoms α to the carbonyl group are replaced by deuterium.

15.

After the removal of one proton (H $^+$) from cyclopentadiene, a pentadienyl anion is obtained. It has a 6π electron system and is, therefore, aromatic.

- **16.** Decoloration with bromine in CCl₄ is a characteristic of unsaturated compounds. But the second reaction with Ag ⁺ to form an insoluble salt is possible only with terminal alkynes.
- **27.** This is an example of nucleophilic aromatic substitution. Reactive intermediate benzyne is produced, followed by an attack by the nucleophile NH₂ at either position.

$$H_3C$$
 NH_2
 H_3C
 NH_2
 H_3C
 NH_2
 H_3C
 NH_2

- **32.** (a) and (b) are examples of methyl/hydride shift under acidic conditions but (c) is an example of just dehydration over a support at high temperature. Under these conditions, the methyl group is not transferred.
- **33.** Geometrical isomerism is shown by compounds of the type abC=Cde.
- **36.** Both (a) and (b) are formed because *N*-bromosuccinimide is used for bromination at the allylic position, and the allylic radical formed in this reaction is resonance-stabilized.

$$^{14}\text{CH}_2$$
=CH-CH $_3$ \Longrightarrow $^{14}\text{CH}_2$ =CH-CH $_2$ \longleftrightarrow $^{14}\text{CH}_2$ -CH=CH $_2$

37. Sulphur is less electronegative than oxygen. The mesomeric effect of oxygen is greater than that of sulphur. Therefore, lone-pair electrons take part in resonance with a carbonyl group and make it less susceptible to a nucleophilic attack.

$$R-C \xrightarrow{\tilde{O}} R - C \xrightarrow{\tilde{O}} R$$

In contrast, thiol esters are not so effectively stabilized by a similar resonance contribution.

$$R-C \xrightarrow{\bar{O}} R-C \xrightarrow{\bar{O}} R-C$$

(A) requires an overlap between the 3p orbital of sulphur and a 2p orbital of carbon. Since this overlap is not large, resonance stabilization by (A) is not so effective.

In fact, (B) makes the carbonyl carbon atom more susceptible to a nucleophilic attack.

Part 5

Practice Test Papers

Practice Test Papers

In the following pages, seven model test papers are provided with the following objectives:

- (a) to assess your preparation,
- (b) to develop your skills of time management, and
- (c) to familiarize you with the actual examination-hall situation.

• Instructions •

The time for each test is 60 minutes. Each test paper contains 44–81 questions. You may attempt questions in any order within this time limit.

Each test paper contains 44-81 questions. Each question carries one mark. There is no negative marking.

Answers are provided after each test paper. Keep a separate blank sheet handy for your rough work. Do not use calculators, log tables or slide rules.

1

Practice Worksheet—1

• Type 1 •

1. B(OH)₃ is a weak acid. It can act as a strong base in the presence of

(a) The B—F bond length in BF₃ is substantially less than the sum of

(b) hydrochloric acid(d) any *trans* diol

Choose the correct option. Only one option is correct.

2. Which of the following statements is correct?

the single-bond radii of B and F.

(c) BBr₃ is a poor electron acceptor compared to BF₃.
(d) The shapes of NO₂⁺, NO₂ and NO₂⁻ are angular.

(a) sodium hydroxide

(b) NF₃ is highly reactive.

(c) glycerol

٥.	restor		of a l	blackened lea	nd pa	ainting. The g	gas also bleaches in The gas (X) is
	(a)	O_2	(b)	NH_3	(c)	O_3	(d) H ₂
4.	300 K.		e nu	mber of mole	s of	NO_2 in 100 g	and N_2O_4 is 38.3 at of the mixture. (d) 0.53
5.		oositronium on. Calculate				0	ectron bound to a
	(a)	13.6 V	(b)	6.8 V	(c)	27.2 V	(d) 3.4 V
6.	500 r		que	ous solution			ssolved to produce (Assume complete
	(a)	1.5×10^{-8}	(b)	2.5×10^{-4}	(c)	2.5×10^{-6}	(d) 2.5×10^{-3}

5-4

3. At room temperature, a blue gas (X) with a characteristic fishy odour can

- 7. The values of the solubility products at 25°C for the following salts are as given below:
 - (i) Ag_2CrO_4 : 2.5×10^{-12} (mol L⁻¹)³
 - (ii) AgCl: 1.5×10^{-10} (mol L⁻¹)²
 - (iii) $Cr(OH)_3$: 1.0×10^{-30} (mol L⁻¹)⁴
 - (iv) Mg(OH)₂: 6.0×10^{-12} (mol L⁻¹)³

Then the solubility in mol L^{-1} is maximum for

(a) Ag₂CrO₄

(b) AgCl

(c) Cr(OH)₃

- (d) $Mg(OH)_2$
- 8. Which of the following compounds is not easily hydrolysed?
 - (a) PCl₃
- (b) SnCl₄
- (c) NF_3
- (d) AsF_3
- **9.** In which of the following equilibrium reactions is the yielding of the product(s) not affected by the pressure applied?
 - (a) $CO(g) + 2H_2(g) \rightleftharpoons CH_3OH(g)$
 - (b) $C(s) + H_2O(g) \rightleftharpoons CO(g) + H_2(g)$
 - (c) $CO(g) + H_2O(g) \rightleftharpoons CO_2(g) + H_2(g)$
 - (d) $NH_4HS(s) \rightleftharpoons NH_3(g) + H_2S(g)$
- 10. Which of the following is correct for an NaCl crystal lattice?
 - (a) The centre-to-centre distance between two nearest oppositely charged ions is $\frac{1}{2}a$.
 - (b) The centres of two nearest like-charged ions are $\frac{1}{\sqrt{2}}a$ apart.
 - (c) The centres of two nearest like-charged ions are $\frac{\sqrt{3}}{2}a$ apart.
 - (d) The atomic positions of an NaCl crystal are

$$C1^{-1}: \left(\frac{1}{2} \ \frac{1}{2} \ \frac{1}{2}\right), \ \left(\frac{1}{2} \ \frac{1}{2} \ 0,\right) \left(\frac{1}{2} \ 0 \ \frac{1}{2}\right), \ \left(0 \ \frac{1}{2} \ \frac{1}{2}\right)$$

and

$$Na^{+}: \left(\frac{1}{2} \ \frac{1}{2} \ \frac{1}{2}\right), \ \left(0 \ 0 \ \frac{1}{2}\right), \ \left(0 \ \frac{1}{2} \ 0\right), \ \left(\frac{1}{2} \ 0 \ 0\right).$$

- 11. Which of the following complexes does not obey the 18-electron rule (EAN rule)?
 - (a) Ni(CO)₄

(b) $[Cr(NH_3)_6]^{3+}$

(c) $[Fe(CN)_6]^{4-}$

- (d) $[Co(NH_3)_6]^{3+}$
- **12.** The hexagonal-close-packed (hcp) and cubic-close-packed (ccp) arrangements are described respectively as
 - (a) ABABAB... and ACBABCA...
 - (b) ABBABA... and ABCABCABC...

- (c) ABABAB... and ABCABCABC...
- (d) ABCACBAB... and ACBACBABAC...
- **13.** A flask containing air at 27°C under the standard atmospheric pressure is corked. A pressure of 2.5 atm inside the flask would force the cork out. The temperature at which this will happen is
 - (a) 167 K

(b) 677 K

(c) 750 K

(d) 120 K

- 14. Which of the following complexes is a high-spin complex?
 - (a) $[Fe(CN)_6]^{4-}$

(b) $[Ni(CN)_4]^{2-}$

(c) $[Fe(C_2O_4)_3]^{3-}$

(d) $[Co(NH_3)_6]^{3+}$

- **15.** When a gas filled in a closed vessel is heated through 1°C, its pressure increases by 0.5%. The initial temperature of the gas was
 - (a) 100 K
- (b) 200 K
- (c) 300 K
- (d) 180 K
- 16. Red phosphorus is less reactive than white phosphorus because
 - (a) the P—P bond length is greater in red phosphorus
 - (b) red phosphorus exists as a polymeric structure
 - (c) red phosphorus is held by the van der Waals forces
 - (d) red phosphorus has more donor ability
- 17. Consider the tetrahedral P₄ molecule and identify the correct statement.



- (a) The P—P distance is 2.21 Å and the P—P—P angle is 120°.
- (b) The P—P—P angle indicates that it is a strained molecule having a strain energy of 6 kJ mol⁻¹.
- (c) The lengths of the six P—P bonds in this molecule are smaller than the P—P bond lengths in other molecules containing multiple phosphorus atoms.
- (d) None of these is correct.
- 18. Alumina is insoluble in water because
 - (a) it is a covalent compound
 - (b) it has a high lattice energy and a low enthalpy of hydration
 - (c) it has a low lattice energy and a high enthalpy of hydration
 - (d) Al 3+ and O 2- ions are not excessively hydrated
- 19. When hydrated MgCl₂·6H₂O is strongly heated,
 - (a) MgO is formed
- (b) anhydrous MgCl₂ is formed
- (c) Mg(OH)HCl is formed
- (d) Mg(OH)Cl is formed

20. Which of the following bases would give 1-alkene?

(a)
$$CH_3$$
 (b) CH_2CH_3 $CH_3CO^ CH_3$ (c) $CH_3CH_2O^-$ (d) $CH_3CH_2CO^ CH_3CH_2CO^-$

- **21.** Which of the following alkyl halides could be successfully used to form a Grignard reagent?
 - (a) $HOCH_2CH_2CH_2CH_2Br$ (b) CH_3 —N— $CH_2CH_2CH_2Br$ CH_3
 - (c) BrCH₂CH₂CH₂COOH (d) H₂NCH₂CH₂CH₂Br
- **22.** What product will be obtained when the *vicinal* diol shown below is heated in an acidic solution?

- **23.** Which of the following have been arranged in the order of decreasing reactivity towards electrophilic substitution?
 - (a) Ethylbenzene > anisole > benzene > chlorobenzene
 - (b) 2,4-Dinitrophenol > 1-chloro-2,4-dinitrobenzene > 2,4-dinitrotoluene
 - (c) 2,4-Dinitrophenol > 2,4-dinitrotoluene > 1-chloro-2,4-dinitrobenzene
 - (d) p-Xylene > p-cresol > toluene

• Type 2 •

Choose the correct options. More than one option is correct.

- **24.** Which of the following statements are false?
 - (a) The addition of HBr to an alkene is stereospecific.
 - (b) *meso* Compounds do not rotate polarized light.
 - (c) Diastereoisomers have the same melting point.
 - (d) The addition of HBr to an alkene is stereoselective.
- 25. Which of the following metal ions are precipitated as sulphides by passing H₂S gas in an ammoniacal medium?
 - (a) Cd^{2+}
- (b) Ni ²⁺
- (c) Pb ²⁺
- (d) Zn^{2+}
- 26. Which of the following compounds disproportionate?
 - (a) PbI₄

(b) FeI₃

(c) $Ca(ClO_3)_2$

- (d) AuCl
- 27. Which of the following statements are correct about ethanenitrile?
 - (a) The C—N σ bond is formed using sp 2 hybrids of carbon and nitrogen, just as in ethene.
 - (b) The C—N σ bond is formed using sp hybrids of carbon and nitrogen, just as in ethyne.
 - (c) The second sp orbital of the nitrogen atom is occupied by a lone pair of electrons.
 - (d) The out-of-plane 2p orbitals form the two π bonds.
- 28. Indicate the correct statements.
 - (a) An electrochemical cell is an arrangement in which a chemical reaction does electrical work or in which electrical work is used to bring about a chemical reaction.
 - (b) The galvanic cell is an electrochemical cell that produces electricity as a result of spontaneous reactions occuring inside it.
 - (c) The electrolytic cell is an electrochemical cell in which a nonspontaneous reaction is driven by an external source of current.
 - (d) Salt bridge is a concentrated electrolytic solution of $KNO_3/KCl/NH_4NO_3$ in agar jelly that completes the electrical circuit between two electrical compartments.
- 29. The structure of rock salt (NaCl)
 - (a) consists of two mutually interpenetrating and slightly expanded face-centred-cubic (fcc) arrays of ions
 - (b) consists of two interpenetrating simple cubic arrays of ions—one of the cations and the other of the anions

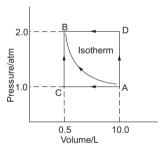
- (c) has radius ratios greater than 0.414 and less than 0.732
- (d) has a 4:4 coordination
- 30. The reaction of C₆H₅CH=CHCHO with NaBH₄ does not give
 - (a) C₆H₅CH₂CH₂CH₂OH
- (b) C₆H₅CH₂CH₂CHO
- (c) C₆H₅CH=CHCH₂OH
- (d) C₆H₅CH₂CHOHCH₃
- **31.** For the equilibrium

$$P+Q \stackrel{k_f}{===} A+B,$$

- (a) $k_f[P]_{eq}[Q]_{eq} = k_r[A]_{eq}[B]_{eq}$
- (b) $K = \frac{A_f}{A_r} \exp\left(\frac{-\Delta E}{RT}\right)$
- (c) if ΔE is positive, an increase in temperature results in an increase in equilibrium constant
- (d) $\Delta G^{\circ} = -RT \ln \frac{K}{Q}$

• Comprehension-Type Questions •

• Study the following graph, assuming perfect gas behaviour. Take $C_{V,m} = \frac{3}{2}R$ and T = 313 K.



- 32. The amounts of work done on the gas along the paths ACB and ADB are respectively
 - (a) -19.5×10^2 J and -2.9×10^4 J (b) -9.5×10^2 J and -1.9×10^4 J
 - (c) -1.9×10^4 [and -9.5×10^4] (d) -9.5×10^2 [and -9.5×10^2]
- **33.** The work done on the gas along the isotherm *AB* is
 - (a) $-4.0 \times 10^3 \text{ J}$

(b) $3.0 \times 10^3 \text{ J}$

(c) $4.0 \times 10^3 \text{ J}$

- (d) $-3.0 \times 10^3 \text{ J}$
- **34.** The value of *q* for all the three paths will be
 - (a) 939 cal (b) 1565 cal
- (c) zero
- (d) infinity

• Matching-Type Questions •

Column A

Column B

35. $\frac{\Delta_{\text{mix}}S'}{N}$

(a) Third law of thermodynamics

 $R\sum_{i=1}^{n}x_{i}\ln x_{i}$

(b) *V*

36. $\lim_{T\to 0}^{t-1} C_{p, m}$

(c) $\frac{C_V}{T}$

38. $\frac{\partial G}{\partial v}\Big|_{T}$

37. $\frac{\partial G}{\partial T}\Big|_{T}$

(d) Real gases

39. $\frac{\partial S}{\partial T}\Big|_{V}$

(e) -S

40. $\frac{\partial T}{\partial p}\Big|_{H} \neq 0$

(f) -1

41. $\Delta S \rightarrow 0$ as $T \rightarrow 0$

(g) Infinity

• Numerical Problems •

42. The element curium ($^{248}_{96}$ Cm) has a mean life of 10 13 seconds. Its primary decay modes are spontaneous fission and α -decay, the former with a probability of 8% and the latter with a probability of 92%. Each fission releases 200 MeV of energy. The masses involved in α -decay are as follows:

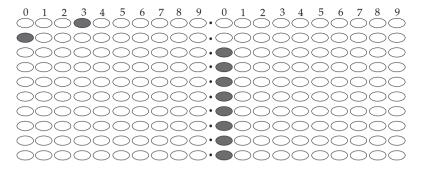
 $^{248}_{96}{\rm Cm}$ = 248.072220 u, $^{244}_{94}{\rm Pu}$ = 244.064100 u and $^{4}_{2}{\rm He}$ = 4.002603 u. Calculate the power output in microwatts from a sample of 10 20 Cm atoms (given that 1 u = 931 MeV/ c^2).

- 43. A hydrogen-like atom with the atomic number Z is in an excited state of the quantum number 2n. It can emit a photon of 204 eV maximum energy. If it makes a transition to the quantum state n, a photon of 40.8 eV energy is emitted. After obtaining the values of n, Z and the ground-state energy (in eV) of this atom, calculate the minimum energy (in eV) that can be emitted by this atom during de-excitation. The ground-state energy of a hydrogen atom is given 13.6 eV.
- 44. A KMnO₄ solution is prepared by dissolving 1.185 g KMnO₄ in water and diluting it to 500 mL. How many millilitres of this will react with the iron in 0.500 g of an ore containing 36% of Fe₂O₃ by mass? The titration reaction takes place in an acidic medium.

Answers

4	•		4	= 1
1. c	2. a	3. c	4. c	5. b
6. d	7. a	8. c	9. c	10. b
11. b	12. c	13. c	14. c	15. a
16. b	17. c	18. d	19. d	20. b
21. b	22. c	23. c	24. a, c, d	25. b, d
26. a, b, d	27. b, c, d	28. a, b, c, d	29. a, c	30. a, b, d
31. b, c				
32. b	33. d	34. c		
$\textbf{35.} \leftrightarrow f$	36. \leftrightarrow g	37. ↔ e	38. \leftrightarrow b	39. \leftrightarrow c
40. \leftrightarrow d	41. \leftrightarrow a			
42. 33.2				
0 1 2 3 0 43. 10.4				

44, 30



Hints to More-Difficult Numerical Problems

42. The reaction involved in α -decay is

$$^{248}_{96}$$
 Cm $\rightarrow ^{244}_{94}$ Pu + $^{4}_{2}$ He.

Mass defect,
$$\Delta m$$
 = mass of ${}^{248}_{96}$ Cm - mass of ${}^{244}_{94}$ Pu - mass of ${}^{4}_{2}$ He = (248.072220 - 244.064100 - 4.002603) u = 0.005517 u.

 \therefore the energy released in α -decay will be

$$E_{cr} = \Delta m \cdot c^2 = (0.005517 \times 931) \text{ MeV} = 5.136 \text{ MeV}.$$

Also, $E_{\text{fission}} = 200 \text{ MeV (given)}$.

By question, mean life,
$$\tau = 10^{13} \text{ s} = \frac{1}{\lambda}$$
.

:. disintegration constant, $\lambda = 10^{-13} \text{ s}^{-1}$.

The rate of decay at the moment when the number of nuclei is 10^{20} is given by

$$-\frac{dN}{dt} = \lambda N = (10^{-13} \text{ s}^{-1}) (10^{20}) = 10^7 \text{ Bq}.$$

In the above disintegration, 8% is by fission and 92% is by α -decay.

Therefore, the total energy released per unit time equals

$$(8\% \times 10^{7} \times 200 + 92\% \times 10^{7} \times 5.136) \text{ MeV s}^{-1}$$

= $(0.08 \times 10^{7} \times 200 + 0.92 \times 10^{7} \times 5.136) \text{ MeV s}^{-1}$

=
$$2.074 \times 10^{8}$$
 MeV s $^{-1}$ = 2.074×10^{14} eV s $^{-1}$ $\approx 2.074 \times 1.6 \times 10^{-5}$ J s $^{-1}$.

Power output = $2.074 \times 1.6 \times 10^{-5} \text{ W}$

=
$$33.2 \times 10^{-6}$$
 W = $33.2 \mu W$.

43. Energy for an H atom, $E_n = -\frac{13.6}{n^2} Z^2 \text{ eV}$.

For transition from 2n to 1,

$$204 = 13.6Z^{2} \left[\frac{1}{1^{2}} - \frac{1}{(2n)^{2}} \right]. \tag{1}$$

For transition from 2n to n,

$$40.8 = 13.6Z^{2} \left(\frac{1}{n^{2}} - \frac{1}{(2n)^{2}} \right) \cdot \dots (2)$$

From equation (2) we get 2n = Z.

... (3)

From equation (1) and (3) we get Z = 4 and n = 2.

 $\therefore 2n = 2 \times 2 = 4.$

The minimum energy for transition from 4 to 3 is

$$E = 13.6 \ 4^2 \left(\frac{1}{3^2} - \frac{1}{4^2}\right) \text{ eV} = 13.6 \times 4^2 \left(\frac{7}{9 \times 16}\right) \text{ eV} = 10.45 \text{ eV}.$$

44. Equivalent weight of KMnO₄ in acidic medium = $\frac{M}{5} = \frac{158}{5} = 31.6$ (since MnO₄⁻ + 8H ⁺ + 5e \rightarrow Mn ²⁺ + 4H₂O).

$$\therefore \text{ normality of KMnO}_4 = \frac{1.185}{500 \times 10^{-3}} \div 31.6$$
$$= \frac{2.37}{31.6} = 0.075.$$

Using milliequivalent concept,

milliequivalent of $KMnO_4$ = milliequivalent of Fe_2O_3

or
$$0.075V = \frac{0.5 \times 36\%}{80} \times 10^3 \text{ mL}$$

or
$$0.075V = \frac{0.5 \times 36}{80 \times 100} \times 10^{3} \text{ mL}$$

or
$$V = 30 \text{ mL}.$$

 \therefore volume of KMnO₄ = 30 mL.

Practice Worksheet—2

• *Type* 1 •

Cl

100	se the	correct option. Only one optic	n is correct.
1.	1.8×1	$10^{-5}~{ m s}^{-1}$ at 25°C. The observed	\rightarrow 4NO ₂ + O ₂ , the rate constant k is rate law is $r = k[N_2O_5]$. Calculate the stainer with $p_{N_2O_5} = 0.10$ atm at 25°C.
	(a)	-1.4×10^{-7} mol dm $^{-3}$ s $^{-1}$	(b) $7.2 \times 10^{-7} \text{ mol dm}^{-3} \text{ s}^{-1}$
	(c)	8.5×10^{-8} mol dm $^{-3}$ s $^{-1}$	(d) $7.2 \times 10^{-8} \text{ mol dm}^{-3} \text{ s}^{-1}$
2.	In gas	s-phase kinetics, pressures ins	tead of concentrations are sometimes
	used	* *	for $aA \rightarrow products$, one finds that

 $-\frac{1}{a}\frac{dp_A}{dt} = k_p p_{A'}^n$ where k_p is a constant and p_A is the partial pressure of A.

The value of k_P is

(a)
$$k(RT)^{1-n}$$
 (b) $k(RT)^{n-1}$ (c) $k(RT)^{a-n}$ (d) $k(RT)^{n-a}$

3. If $E_a = 30$ kcal mol⁻¹, a 10°C rise in temperature above the room temperature (298 K) causes the rate constant to be

- (a) tripled (b) quadrupled (d) halved
- (c) quintupled **4.** For reactions 1 and 2, $A_1 = 5A_2$ and $k_1 = 100k_2$ at room temperature. Then the
- difference in energy of activation $(E_{a1} E_{a2})$ is (b) 1.8 kcal mol⁻¹ (a) −1.8 kcal mol⁻¹
 - (c) -3.8 kcal mol⁻¹ (d) $-2.8 \text{ kcal mol}^{-1}$
- Ordinary hydrogen at room temperature is a mixture of
 - (a) 25% orthohydrogen + 75% parahydrogen
 - (b) 75% orthohydrogen + 25% parahydrogen

- (c) 50% orthohydrogen + 50% parahydrogen
- (d) 60% orthohydrogen + 40% parahydrogen
- 6. Which of the following solutions has the same value of pH as 0.15-M CH₃CO₂NH₄?
 - (a) 0.10-M NH₃

(b) 0.05-M CH₃CO₂NH₄

(c) 0.50-M NH₃

(d) 0.15-M Na₂CO₃

7. In the van der Waals equation of state,

$$\left(p + \frac{an^2}{V^2}\right)(V - nb) = nRT,$$

- (a) *nV* represents the total effective volume of *n* moles of the gas
- (b) a represents the magnitude of repulsive forces
- (c) the greater the value of b, the smaller is the size of the molecules
- (d) the greater the value of a, the greater will be the chance of liquefaction of gases
- 8. The ratio of the inversion temperature of a gas to its Boyle temperature (both in kelvins) is
 - (a) 2.5
- (b) 1.5
- (c) 3.5
- (d) 2.0

 $4NH_3(g) + 5O_2(g) \rightarrow 4NO(g) + 6H_2O(l); \Delta H_{298 \text{ K}}^{\circ} = -1170 \text{ kJ mol}^{-1}$ 9. $2NO(g) + O_2(g) \rightarrow 2NO_2(g); \Delta H_{298 \text{ K}}^{\circ} = -114 \text{ kJ mol}^{-1}$ $3NO_2(g) + H_2O(1) \rightarrow 2HNO_3(1) + NO(g); \Delta H_{298 \text{ K}}^{\circ} = -72 \text{ kJ mol}^{-1}$

Given the above $\Delta H_{298\text{K}}^{\circ}$ values, find $\Delta H_{298\text{K}}^{\circ}$ for the reaction $NH_3(g) + 2O_2(g) \rightarrow HNO_3(l) + H_2O(l)$.

(a) -141 kJ mol^{-1}

(b) -828 kJ mol^{-1}

(c) -414 kJ mol⁻¹

- (d) -207 kJ mol^{-1}
- 10. Which of the following constants has the same unit as that of entropy?
 - (a) Rate constant

- (b) Boltzmann constant
- (c) Equilibrium constant
- (d) Avogadro constant
- **11.** Given that the standard reduction potentials of metallic cations $E_{A^{2+}/A}^{\circ} = +0.34 \text{ V}$, $E_{B^{2+}/B}^{\circ} = -2.37 \text{ V}$ and $E_{C^{+}/C}^{\circ} = -3.05 \text{ V}$, the order of reducing power of the corresponding metals is
 - (a) A > B > C
- (b) C > A > B
- (c) B > A > C (d) C > B > A
- 12. The order of basicity of amines

 $NH_3 < (CH_3)_3N < CH_3CH_2NH_2 < CH_3NHC_2H_5$

is not governed by which of the following factors?

- (a) Inductive effect
- (b) Solvation

(c) Steric effect

- (d) Dissociation
- 13. Which of the following pairs of structures are identical?
 - (a) Borazine and cyclohexane
- (b) Diborane and ethane
- (c) Boron nitride and graphite
- (d) B(OH)₃ and Al(OH)₃

- **14.** The decreasing order of acid strength of acetic acid, trimethylacetic acid, trichloroacetic acid and nitroacetic acid is
 - (a) $(CH_3)_3CCO_2H > CH_3CO_2H > CCl_3CO_2H > O_2NCH_2CO_2H$
 - (b) $CH_3CO_2H > (CH_3)_3CCO_2H > O_2NCH_2CO_2H > CCl_3CO_2H$
 - (c) $O_2NCH_2CO_2H > (CH_3)_3CCO_2H > CH_3CO_2H > CCl_3CO_2H$
 - (d) $CCl_3CO_2H > O_2NCH_2CO_2H > CH_3CO_2H > (CH_3)_3CCO_2H$
- **15.** Which of the following sets of reactants are not used to prepare diborane?
 - (a) $Mg_3N_2 + H_3PO_4$
- (b) $BF_3 + H_2$
- (c) $(C_2H_5)_2O \cdot BF_3 + LiAlH_4$
- (d) $Na[BH_4] + I_2 + ether$
- 16. Which of the following has the highest value of dipole moment?
 - (a) CH₃OH

(b) CH₃NH₂

(c) CH₃NO₂

- (d) CH₃Cl
- 17. The buffer capacity of a solution containing a weak acid and a salt of its conjugate base is maximum when
 - (a) [salt] = [base]

(b) [salt] < [acid]

(c) [salt] > [acid]

- (d) $pH = pK_a$
- 18. Which of the following compounds is called the 'inorganic benzene'?
 - (a) $B_6N_6H_3$

(b) $B_3N_3H_6$

(c) $B_3N_3H_3$

- (d) $B_3N_6H_6$
- 19. Consider the hypothetical reversible reaction

$$\frac{1}{2}A_2(g) + \frac{3}{2}B_2(g) \rightleftharpoons AB_3(g); \Delta H = -30 \text{ kJ mol}^{-1}.$$

If the standard entropies of A_2 , B_2 and AB_3 are 80 J K⁻¹ mol⁻¹, 60 J K⁻¹ mol⁻¹ and 90 J K⁻¹ mol⁻¹ respectively, the above reaction will be at equilibrium at

- (a) 450 K
- (b) 750 K
- (c) 250 K
- (d) 1500 K
- **20.** Which of the following structures of sulphur trioxide is tetrahedral in shape?



(d) None of these

21. Indicate which of the following species is the most stable.

- (b) CH₃CHCH₂CH₃
- (c) $CH_2 = CHOH$
- (d) CH₃CH₂CH₂CH₂CH₂

22. Which of the following compounds have no chirality centres?

- (a) 2,3-dichloropentane
- (b) 2,3-dichloro-2,3-dimethylbutane
- (c) 2,3-dibromopentane

23. Which of the following pairs is not a pair of keto-enol tautomers?

(b)
$$CH_3CH_2COCO_2C_2H_5$$
 and $CH_3CH = CCO_2C_2H_5$

(c)
$$CH_3CH_2CH_2C=CH_2$$
 and $CH_3CH_2CH_2CCH_3$ OH

• *Type 2* •

Choose the correct option. More than one option is correct.

24. The chemical process in the production of metal using self-reduction of the oxide and the sulphide is applied for the ore

(a) galena

(b) chalcopyrite

(c) haematite

(d) cassiterite

25. D-Glucose and D-mannose are

(a) annomers

(b) epimers

(c) diastereomers

(d) oligosaccharides

- **26.** ClO_2 is a free radical with one unpaired electron. It has less tendency to dimerize than
 - (a) NO₂
- (b) ClO₃
- (d) NO
- (d) AlCl₃

- 27. The N₂O₃ molecule
 - (a) has an O₂N—NO structure in gas phase at low temperature
 - (b) has an O₂N—NO structure in liquid phase at low temperature
 - (c) has a very long N—N bond distance
 - (d) dissociates easily
- **28.** Which of the following pairs of molecules are those of constitutional isomers?

29. Complete the following sequences with appropriate reagents:

- (a) I is O_3/Zn , H_2O ; II is SeO_2 ; III is $OH^-/heat$
- (b) I is H₂O; II is LiAlH₄; III is O₅O₄
- (c) I is O_3/Zn , H_2O
- (d) II is LiAlH₄; III is H⁺/heat
- **30.** A solution of $Na_2S_2O_3$ is standardized iodometrically against 0.1262 g of pure KBrO₃, requiring 45 mL of the $Na_2S_2O_3$ solution. What is the strength of the $Na_2S_2O_3$ solution?
 - (a) 0.2 M

(b) 0.1 M

OH

(c) 0.05 M

(d) 0.1 N

- **31.** In the lead storage cell,
 - (a) Pb is oxidized to PbSO₄ at the anode
 - (b) PbO₂ is reduced to PbSO₄ at the cathode
 - (c) both the electrodes are immersed in the same aqueous solution of H₂SO₄
 - (d) the use of salt bridge is not necessary

• Comprehension-Type Questions •

- A colourless solid (A) on strong heating gives a brown gas (B) and a grey residue (C). On dissolution of solid (A) in NH₃, a solution of compound (D) is formed. (D) oxidizes glucose. FeSO₄ reduces the solution of (A) in water. The aqueous solution of (A) also gives a brick-red precipitate (E) with a K₂CrO₄ solution.
 - **32.** The compound (A) will be
- (a) $Pb(NO_3)_2$ (b) $AgNO_3$ (c) $Hg_2(NO_3)_2$ (d) $Bi(NO_3)_3$
- **33.** The brown gas (B) is
 - (a) NO
- (b) N_2O_3
- (c) N_2O_5
- (d) NO₂

- **34.** The grey residue (C) contains
 - (a) Hg
- (b) Bi
- (c) Ag
- (d) Pb

- **35.** The solution (D) is
 - (a) BiI₂
 - (c) $[Pb(NH_3)_2]^{2+}$

- (b) $[Ag(NH_3)_2]^+$
- (d) $[HgI_4]^{2-}$
- **36.** The brick-red precipitate (E) is
 - (a) Ag_2CrO_4

(b) $HgI_2 + Hg CrO_4$

(c) $Bi_2(CrO_4)_3$

(d) PbCrO₄

• Matching-Type Questions •

Column A

- 37. $CrO_4^{2-} \rightarrow CrO_5$
- 38. NaNO₃ \rightarrow NH₃
- 39. $Mn^{2+} \rightarrow MnO_4^{-}$
- **40.** $Al(OH)_4^- \rightarrow Al(OH)_3$
- 41. $XeF_4 \rightarrow XeO_3$

Column B

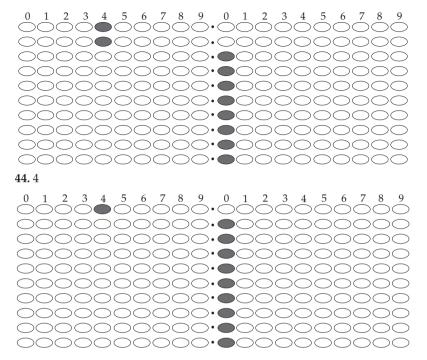
- (a) PbO_2/H^+
- (b) $H^+(aq)$
- (c) H₂O
- (d) H_2O_2/H^+
 - (e) Devarda's alloy/OH⁻

• Numerical Problems •

- **42.** The molar enthalpy of fusion of anthracene is $\Delta_{\rm fus}H_{\rm m}$ = 28.8 kJ mol ⁻¹ and its melting point is 217°C. Calculate the ideal solubility of anthracene per kilogram of benzene at 25°C.
- **43.** For the reaction $A+2B \rightarrow 3C+D$, the rate constant (k) is $2\times 10^{-3} \, L^2 \, \text{mol}^{-2} \, \text{s}^{-1}$ at 500 K. Introducing a catalyst at 1200 K, the rate increases to 2.71 times. Find the final activation energy if the initial activation energy is same as that of the reaction $2A \rightarrow P$, whose rate constant at 300 K in the presence of the catalyst is $2\times 10^{-3} \, \text{min}^{-1}$ and temperature coefficient is 2.0.
- **44.** Ice crystallizes as a hexagonal lattice. At low temperatures the lattice parameters are found to be a = 4.53 Å and c = 7.41 Å. How many H₂O molecules are there in a unit cell (given that $P_{\text{ice}} = 0.92 \text{ g cm}^{-3}$ at 0°C)?

		<u>Answers</u>		
4 1	2 1	0	4	.
1. d	2. d	3. c	4. a	5. c
6. b	7. d	8. d	9. c	10. b
11. d	12. d	13. c	14. d	15. b
16. c	17. d	18. b	19. b	20. b
21. a	22. b	23. d	24. a, b	25. b, c
26. a, b, c, d	27. a, c, d	28. b, d	29. c, d	30. b, d
31. a, b, c				
32. b	33. d	34. c	35. b	36. a
37. \leftrightarrow d	$38. \leftrightarrow \mathbf{e}$	39. \leftrightarrow a	40. \leftrightarrow b	41. \leftrightarrow c
42. 24				
0 1 2 3	4 5 6 7	8 9 0 1	2 3 4 5	6 7 8 9
0000				
0000	0000			
$\bigcirc\bigcirc\bigcirc\bigcirc\bigcirc$	0000			
0000	0000			
0000	0000			
0000	0000			

43.44



Hints to More-Difficult Numerical Problems

42. The solubility of anthracene per kilogram of benzene can be obtained from its mole fraction with the use of equation.

$$\ln x = \frac{\Delta_{\text{fus}} H_{\text{m}}}{R} \left(\frac{1}{T^{*}} - \frac{1}{T}\right)$$

$$= \frac{28.8 \times 10^{3} \text{ J mol}^{-1}}{8.314 \text{ J K}^{-1} \text{ mol}^{-1}} \left(-\frac{1}{490.15 \text{ K}} - \frac{1}{298.15 \text{ K}}\right)$$

$$= -4.55$$

$$x_{\text{B}} = e^{-4.55} = 0.0106.$$
Since $x_{\text{B}} < 1$, x_{B} (anthracene) = $\frac{n(\text{anthracene})}{n(\text{benzene})}$
or $x_{\text{B}} = \frac{\text{mol. wt of benzene}}{1000} \times \text{molality } (m)$

(the relation between mole fraction and molality)

or
$$x_{\rm B} = \frac{78}{1000} \cdot m$$
 or $0.0106 = \frac{78}{1000} \cdot m$

or $m = 0.136 \text{ mol kg}^{-1}$

:. the solubility of anthracene per kilogram of benzene is

$$(0.136 \times 178) \text{ g} \approx 24 \text{ g}.$$

43. At 500 K, $k = 2 \times 10^{-3} L^2 \text{ mol}^{-2} \text{ s}^{-1}$ (given).

$$\ln \frac{k_2}{k_1} = \ln 2.71 \approx \ln e = 1 = \frac{E_{a1} - E_{a2}}{RT}$$
 or $E_{a1} - E_{a2} = RT$ (1)

Again, temperature coefficient, $K_{t+10 \text{ K}} = 2.0 \implies \frac{K_{310 \text{ K}}}{K_{300 \text{ K}}} = 2.0$

or
$$\ln 2.0 = \frac{E_a}{R} \left(\frac{1}{300} - \frac{1}{310} \right) = \frac{E_a}{R} \cdot \frac{1}{300 \times 31}$$

or
$$2.303 \log 2.0 = \frac{E_a}{R} \times \frac{1}{9300}$$
 or $\frac{E_a}{R} = 6445$.

Here,
$$E_a = E_{a1}$$
 or $E_{a1} = 6445 R$ (2)

From equations (1) and (2),

$$E_{a2} = E_{a1} - RT = (6445 \text{ K})R - RT = R(6445 \text{ K} - T)$$

= 8.314 J K⁻¹ mol⁻¹ · (6445 – 1200) K (: here $T = 1200 \text{ K}$).

:.
$$E_a = 43.606 \text{ kJ mol}^{-1} \approx 44 \text{ kJ mol}^{-1}$$
.

44. For the hcp structure,

$$a = b \neq c$$
, $\alpha = \beta = 90^{\circ}$ and $\gamma = 120^{\circ}$.

As we know.

$$V = abc(1 - \cos^2\alpha - \cos^2\beta - \cos^2\gamma + 2\cos\alpha \cdot \cos\beta \cdot \cos\gamma)^{1/2}$$
$$= a^2c(1 - \cos^2\gamma)^{1/2} = a^2c \cdot \sin\gamma$$
$$= (4.53 \text{ Å})^2(7.41 \text{ Å}) \sin 60^\circ = 132 \times 10^{-24} \text{ cm}^3.$$

Mass of a unit cell = $V \times$ density

=
$$(132 \times 10^{-24} \text{ cm}^3)(0.92 \text{ g cm}^{-3})(6.02 \times 10^{23} \text{ u g}^{-1})$$

= 73 u.

This value is about four times the molecular weight of H_2O . Therefore, we conclude that there are 4 molecules of H_2O per unit cell.

Practice Worksheet—3

• Type 1 •

Choose the correct option. Only one option is correct.

1.	In which	of	the	following	cases	is	the	principle	of	solubility	product
	violated?									-	-

- (a) Addition of Na₂SO₄ to a BaSO₄ solution
- (b) Addition of NaBr to an AgBr solution
- (c) Addition of NaOH to an Mg(OH)₂ solution
- (d) Addition of KCN to an AgCN solution

2.	The experimental density of a solid crystal is found to be less than the	he
	calculated density. This indicates the presence of	

- (a) external impurities
- (b) a line defect
- (c) a Frenkel defect
- (d) a Schottky defect
- **3.** The radius ratio (r_+/r_-) of an ionic crystal is 0.98. The crystal will have a lattice like that of
 - (a) graphite
- (b) TiO_2
- (c) CsCl
- (d) NH₄Cl
- 4. The abnormal behaviour of colligative properties arises due to
 - (a) association of solute particles in the solvent
 - (b) lower enthalpy of dilution
 - (c) higher enthalpy of hydration
 - (d) lowering of the vapour pressure of the solution
- **5.** An electron in an excited state of the hydrogen atom is in the sixth energy level. When it comes back to the ground state,
 - (a) energy is absorbed continuously
 - (b) two spectral lines are found in the infrared region
 - (c) four spectral lines are found in the visible region
 - (d) six spectral lines are found in the ultraviolet region

the salt will be

magnetons, are respectively

(a) 0.05

(b) 0.07

6. The degree of hydrolysis of ammonium cyanide with a molarity of $0.2~\text{mol}~\text{L}^{-1}$ is 0.1. Hence, the degree of hydrolysis of a 0.4-M solution of

7. The magnetic moments of the ions $[Fe(CN)_6]^{3-}$ and $[FeF_6]^{3-}$, in Bohr

(c) 0.10

(d) 0.20

	(a)	1.73 and 2.83	(b)	1.73 and 4.90
	(c)	1.73 and 1.73	(d)	1.73 and 5.92
8.	The d	egree of dissociation of an acet	ic ac	id solution may be increased by
	(a)	adding H_2SO_4 to the solution		
		adding potassium acetate to t		
		increasing the temperature of	acet	ic acid
		diluting the solution		
9.	freezi	ng point of the solution is (give	n th	
	` '	· /	` '	-0.40° C (d) -0.20° C
10.	effect	at ordinary temperatures?		ted due to the Joule-Thomson
	()	Ammonia		Nitrogen
	` ′	Carbon dioxide		Hydrogen
11.	secon	d ionization energy?		nged in the increasing order of
	` '	N < C < Be < O		O < Be < N < C
	` ′	N < O < C < Be	(d)	Be < C < N < O
12.	The fo	ollowing cell is set up at 25°C:	C1.	
		$Ag(s) Ag^+ Cl^- Ag$	gCI(s	$) \mid Ag(s).$
		that $E_{\text{Ag}^{+}/\text{Ag}}^{\circ} = 0.799 \text{ V}$ and E_{Ag}° act of AgCl is	AgCl/	$Ag(s) + Cl^- = 0.222$ V, the solubility
	(a)	$1.8 \times 10^{10} (\text{mol L}^{-1})^2$	(b)	$1.8 \times 10^{-10} \; (\text{mol L}^{-1})^2$
	(c)	$1.8 \times 10^{-13} \text{ (mol L}^{-1})^2$	(d)	$1.8 \times 10^{-17} \text{ (mol L}^{-1})^2$
13.	Which	n of the following complexes ha	as a s	square-planar geometry?
	(a)	$[Ni(CN)_4]^{2-}$ (b) $[Ni(CO)_4]$	(c)	$[CoCl_4]^{2-}$ (d) $[Cu(Cl)_4]^{2+}$
14.	Which	n of the following is a reversible	e hal	lf-cell electrode?
	(a)	$Ag \mid AgCl(s), Cl^{-}(aq) \parallel Cl^{-}, H$	Ig ₂ C	$\operatorname{Il}_2(s) \mid \operatorname{Hg}$
	(b)	$Pt \mid MnO_4^-, Mn^{2+}, H^+ \parallel OH^-$, H ₂ ((g, 1 atm) Pt
	(c)	Pt $Cl_2(g)$, $Cl^- H^+$, $H_2(g, 1)$	ıtm)	Pt
	(d)	$Ag \mid AgCl(s) \mid Cl^{-}(aq)$		

- **15.** Which of the following statements is correct?
 - (a) BH₃ is a poor electron acceptor compared to B(CH₃)₃.
 - (b) The trend in electron affinity for group 15 (VA) is opposite to that for group 16 (VIA).
 - (c) Carbon and arsenic show a diagonal relationship.
 - (d) On being heated, lithium hydride gives Li₂ and H₂.
- 16. During the rusting of iron,
 - (a) iron becomes a positive ion (Fe ²⁺) and forms the cathode, and the noble metal which is present as an impurity in the iron serves as the anode
 - (b) iron and the noble metal present in iron form an electrochemical cell on the surface of iron
 - (c) Fe II is reduced to Fe I
 - (d) none of these happens
- **17.** Which of the following statements is valid in the context of the brown-ring test for nitrates?
 - (a) The complex forming the brown ring is [Fe(H₂O)₅NO₃]SO₄.
 - (b) The brown colour of the ring is not due to d–d transition.
 - (c) The oxidation state of iron in the brown ring is +III.
 - (d) In the iron complex which forms the brown ring, iron has three unpaired electrons.
- 18. When H_2SO_4 is added to a dark-red compound, the resultant solution is red in colour. When NaOH is added to this solution, the colour turns orange. Identify the compound.
 - (a) Pb_3O_4
- (b) HgI₂
- (c) Cu₂O
- (d) CrO_3
- 19. Which of the following reagents is used to distinguish Fe³⁺ from Fe²⁺?
 - (a) $K_3[Fe(CN)_6]$

(b) KSCN

(c) $Fe_4[Fe(CN)_6]_3$

- (d) Dimethylglyoxime
- **20.** Sodium nitroprusside reacts with sulphide ion to give a purple complex represented by the formula
 - (a) $Na_2[Fe(CN)_5NOS_2]$
- (b) $Na_2[Fe(CN)_4CNS]$
- (c) $[Fe(CN)_5NOS]^{4-}$
- (d) [Fe(CN)₄NOS] ³⁻
- **21.** The chemical process in the production of copper from chalcopyrites involves
 - (a) reduction of the oxide
 - (b) oxidation of the sulphide
 - (c) self-reduction of the oxide and the sulphide
 - (d) partial oxidation of Cu₂S and FeS

22. Oxygen is heated at constant pressure through 2°C. The percentage of the heat energy supplied, which is used in increasing internal energy, is

	(a)	60 (b)	71	(c)	85	(d)	40
23.	Which	n of the following	statements is	corr	ect?		
	(a)	A compound w		ral co	entres can	have a	maximum of
	(b)	In the most cyclohexane, bo					
	(c)	2,3-Dichloropen	tane has a ste	reois	omer that i	s a mes	o compound.
		1-Butyne is more					-
24.	The m	olecule having th	ne highest dip	ole r	noment is		
		_	CH ₃ Cl		CH ₃ Br	(d)	CH ₃ I
			T	2			
			 Type 	<u>2</u> •			
Choo	se the	correct options. N	Iore than one	opti	ion is corre	ct.	
25.	Which	n of the following	can liberate (CO_2 ?			
	(a)	Phenol		(b)	$C_6H_5SO_3H$	I	
	(c)	Cyclohexanol		(d)	2,4,6-Trin	itroph	enol
26.	Aceta	ldehyde responds	s positively to)			
	(a)	the iodoform tes	t	(b)	Benedict's	test	
	(c)	the Lucas test		(d)	the Tollen	s test	
27.	HNO	gent is added to a					
		$Na_2C_2O_4$		(b)	$(NH_4)_2S_2C$)。	
		NaBiO ₃			Pb_3O_4	0	
28.		icyanoaurate ion	. [Au(CN) ₂] -			ed by	dissolving Au
		vanide solution in				,	
	(a)	air		(b)	H_2O_2		
	(c)	I -		(d)	SCN-		
29.	Calciu	ım cyanamide is j	prepared by t	he fo	ollowing rea	action:	
		CaC ₂ (s) -	$+ N_2(g) - \frac{1100^{\circ}G}{}$: → C:	aNCN(s) +	C(s).	
	The N	ICN ²⁻ ion of CaN		. •		-(-).	
		linear					
		hont					

 $\begin{array}{ll} \text{(c)} & \text{isostructural and isoelectronic with CO}_2 \\ \text{(d)} & \text{isostructural and isoelectronic with N_2O} \\ \end{array}$

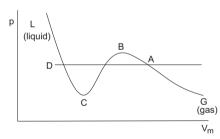
- **30.** During the electrolysis of an aqueous Zn(NO₃)₂ solution using platinum electrodes, we observe that
 - (a) Zn is deposited at the cathode
 - (b) N_2 is evolved at the anode
 - (c) H₂ and N₂ are evolved at the anode
 - (d) O₂ is evolved at the anode
- **31.** In the Langmuir adsorption isotherm, the extent of adsorption (θ) is given by

$$\theta = \frac{bp}{1 + bp} \cdot$$

On the basis of this equation, which of the following are correct statements?

- (a) When the gas is slightly adsorbed, θ will follow zero-order kinetics.
- (b) When the gas is slightly adsorbed, θ will follow first-order kinetics.
- (c) When the gas is strongly adsorbed, θ will follow first-order kinetics.
- (d) When the gas is strongly adsorbed, θ will follow zero-order kinetics.

32.



In the above $p-V_{\rm m}$ graph,

- (a) the segment AB is a metastable region corresponding to the superheated vapour
- (b) the segment CD is a region of supercooled liquid

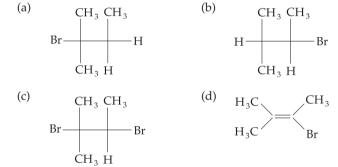
(c) at BC,
$$\frac{\partial p}{\partial V_{\rm m}}\Big]_T < 0$$

(d) at BC,
$$\frac{\partial p}{\partial V_{\rm m}}\Big|_T > 0$$

• Comprehension-Type Questions •

 Alkenes are characterized by C=C bonds. As such, they are subjected to electrophilic addition reactions. Examples of alkenes are shown below.

33. When HBr adds to the compound II above, the main product is



- **34.** What is the order of reactivity of the given compounds during the electrophilic addition of HBr?
 - (a) II > IV > I > III

(b) III > I > IV > II

(c) I > II > III > IV

- (d) IV > I > II > III
- **35.** Which of the four compounds is most likely to form an addition product with the general structure shown below?

$$H_3C$$
 CH_3 Br H (b) II

- (a) I
- (c) III

(d) IV

• Matching-Type Questions •

Column A

36.
$$CH_3CH_2C = CH \xrightarrow{\text{(i) Sia}_2BH/THF} (P)$$

(a)
$$CH_3$$
 $CH CH_2OH$ CH_3

Column B

37.
$$CH_3C = C \cdot CH_3 \xrightarrow{\text{(i) BF}_3/\text{THF}} (P)$$

b)
$$CH_2CH_3$$

 H OH
 H OH
 CH_2CH_3

$$\textbf{38.} \quad CH_3CH_2C \stackrel{HgSO_4}{=\!\!\!=\!\!\!\!=} (P)$$

(c)
$$CH_3CH_2C \equiv C \cdot CH_2OH$$

39.
$$CH_3CH_2C \equiv CH \xrightarrow{(i) CH_3MgBr/HCHO)} (P)$$

40.
$$CH_3CH_2C = CH \xrightarrow{(i) NaNH_2/CH_3CH_2Br} (P)$$

41.
$$CH_3$$
 $C=CH \xrightarrow{BF_3} (P)$

(f)
$$CH_3CH = CH \cdot CH_3$$

(g)
$$CH_3$$
 $C=C$
 CH_3

(h) CH₃CH₂CH₂CH₂CH₂CHO

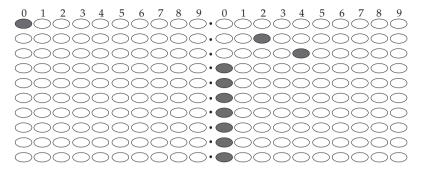
• Numerical Problems •

- **42.** Two moles of an ideal monatomic gas occupy a volume V at 57°C. The gas expands adiabatically to a volume 2V. Calculate the work done (in joules) by the gas during this process.
- 43. Two glass bulbs of equal volume are connected by a narrow tube and are filled with a gas kept at a temperature of 0°C and a pressure of 76 cmHg. One of the bulbs is then placed in melting ice and the other is placed in a water bath maintained at 62°C. Calculate the new pressure (in atm) inside the bulbs. The volume of the connecting tube is negligible.

44. Estimate the standard reduction potential (in volts) of a CuS(s) | Cu(s) electrode, given that $K_{\rm s}$ for CuS(s) is 8.5×10^{-36} (mol L⁻¹)² and $E_{\rm Cu^{2+}|Cu(s)}^{\rm out}=+0.34$ V. Finally, calculate $E^{\rm o}$ of the cell when it is coupled with an Fe²⁺ | Fe electrode, given that $E_{\rm Fe^{2+}|Fe}^{\rm e}=-0.45$ V.

<u>Answers</u>						
1. d	2. d	3. c	4. a	5. c		
6. b	7. d	8. d	9. a	10. d		
11. d	12. b	13. a	14. d	15. b		
16. b	17. d	18. d	19. b	20. c		
21. c	22. b	23. d	24. b	25. b, d		
26. a, b, d	27. c, d	28. a, b	29. a, c, d	30. a, d		
31. b, d	32. a, b, c					
33. a	34. d	35. c				
36. \leftrightarrow d	37. \leftrightarrow f	38. \leftrightarrow e	39. \leftrightarrow c	40. \leftrightarrow b		
41. \leftrightarrow a						
42. 3043						
	4 5 6 7 0000 0000 0000 0000 0000					
43. 1.1						
	† 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0					

44, 0.24



Hints to More-Difficult Numerical Problems

42.
$$\gamma = \frac{5}{3}$$
 (for monatomic gas)
 $T_1 = (273 + 57) \text{ K} = 330 \text{ K}$
 $V_1 = V$
 $V_2 = 2V$

 $T_2 = ?$

Since the gas expands adiabatically,

$$T_1 V_1^{\gamma - 1} = T_2 V_2^{\gamma - 1}$$

or
$$T_2 = T_1 \left(\frac{V_1}{V_2}\right)^{\gamma - 1} = (330 \text{ K}) \left(\frac{V}{2V}\right)^{5/3 - 1} = \frac{330}{\frac{2}{3}} \text{ K} = 208 \text{ K}.$$

Work done =
$$-\frac{nR(T_2 - T_1)}{\gamma - 1} = -\frac{2 \times 8.314(208 - 330)}{\frac{5}{3} - 1}$$
 J = $\frac{2 \times 8.314 \times 122 \times 3}{2}$ J = 3043 J.

43. Let x mol shift from the high-temperature side to the low-temperature side.

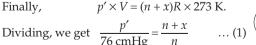
Final stage

62°C

For the left bulb, pV = nRT.

Initially, 76 cmHg \times $V = nR \times 273$ K.

 $p' \times V = (n + x)R \times 273 \text{ K}.$ Finally,



For the right bulb, pV = nRT.

Initially, 76 cmHg \times $V = nR \times 273$ K.

Finally,
$$p' \times V = (n - x)R \times (273 + 62) \text{ K}.$$

Dividing,
$$\frac{p'}{76 \text{ cmHg}} = \frac{n-x}{x} \times \frac{335 \text{ K}}{273 \text{ K}} \cdot \dots (2)$$

From (1) and (2),

$$\frac{n+x}{n} = \frac{n-x}{n} \times \frac{335 \text{ K}}{273 \text{ K}}$$
 or $n = \frac{608}{62} x$ (3)

Substituting (3) in (1), we get

$$\frac{p'}{76 \text{ cmHg}} = 1 + \frac{62}{608}$$

or $p' = \frac{670}{608} \times 76 \text{ cmHg} = 83.75 \text{ cmHg} = 84 \text{ cmHg} = 1.1 \text{ atm.}$

44. Let E° for $CuS(s) + 2e \rightleftharpoons Cu(s) + S^{2-}$ be x.

The cell reaction is

$$Cu(s) \rightarrow Cu^{2+} + 2e; E^{\circ} = 0.34 \text{ V}$$

 $\underline{CuS(s) + 2e \rightarrow Cu(s) + S^{2-}; E^{\circ} = x}$
 $\underline{CuS(s) \rightarrow Cu^{2+} + S^{2-}; E^{\circ}_{cell} = x - 0.34 \text{ V}}$

We know,
$$\log K_{\rm s} = \frac{nE_{\rm cell}^{\circ}}{0.0592}$$
 or $E_{\rm cell}^{\circ} = \frac{0.0592}{2} \log(8.5 \times 10^{-36}) \text{ V}$

or $x - 0.34 \text{ V} = 0.0296 \log(8.5 \times 10^{-36}) \text{ V}.$

$$\therefore$$
 $x = -0.69 \text{ V}.$

Finally, $E^{\circ} = E_{\text{Fe}^{2+}|\text{Fe}} - E^{\circ}_{\text{CuS}|\text{Cu}} = -0.45 \text{ V} - (-0.69 \text{ V}) = +0.24 \text{ V}.$

IIT Questions—1

Useful data

Molar gas constant, R = 0.082 L atm K⁻¹ mol⁻¹ = 8.314 J K⁻¹ mol⁻¹

Atomic numbers: Be = 4; B = 5; C = 6; N = 7; O = 8; S = 16; Xe = 54

Atomic masses: H = 1.008; O = 16.00; S = 32.07; Se = 78.96; Te = 127.6

- 1. The rate of physisorption increases with
 - (a) decrease in temperature
- (b) increase in temperature
- (c) decrease in pressure
- (d) decrease in surface area
- 2. Which of the following represents the sp², sp², sp modes of hybridization respectively from left to right?
 - (a) H₂C=CH—C≡N

(b) HC≡C—C≡CH

(c) $H_2C = C = C = CH_2$

(d) H_2C CH_2

3.
$$F \longrightarrow NO_2 \xrightarrow{DMF, \Delta} (A) \xrightarrow{0-5^{\circ}C} (B).$$
(ii) $\frac{0-5^{\circ}C}{(i) NaNO_2/HCl} \longrightarrow (B).$

In the above reaction, the product (B) is

(a)
$$H_2N$$
 \longrightarrow N CH_3 (b) O_2N CH_3

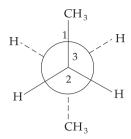
(b)
$$O_2N$$
 N N H_2N

(c)
$$H_2N \longrightarrow NH_2$$

$$-NH_2$$
 (d) O_2N NH_2

- 4. 2-Phenylpropene on acidic hydration gives
 - (a) 2-phenyl-2-propanol
- (b) 2-phenyl-1-propanol
- (c) 3-phenyl-1-propanol
- (d) 1-phenyl-2-propanol

5.



 $\overset{'}{C}$ is rotated anticlockwise 120° about the $\overset{'}{C}$ — $\overset{''}{C}$ bond. The resulting conformer is

- (a) partially eclipsed
- (b) eclipsed

(c) gauche

- (d) staggered
- 6. If the bond length of the C—O bond in carbon monoxide is 1.128 Å then what is the value of the C—O bond length in Fe(CO)₅?
 - (a) 1.15 Å

(b) 1.128 Å

(c) 1.72 Å

- (d) 1.118 Å
- 7. The species present in the solution when CO_2 is dissolved in water are
 - (a) CO_2 , H_2CO_3 , HCO_3^- , CO_3^{2-} (b) H_2CO_3 , CO_3^{2-}

(c) CO₃²⁻, HCO₃⁻

- (d) CO₂, H₂CO₃
- 8. The number of P—O—P bonds in cyclic metaphosphoric acid is
 - (a) zero

(b) two

(c) three

- (d) four
- 9. $\Delta_{\text{vap}} H_{\text{m}} = 30 \text{ kJ mol}^{-1}$ and $\Delta_{\text{vap}} S_{\text{m}} = 75 \text{ J K}^{-1} \text{ mol}^{-1}$. Find the temperature of vapour at 1 atm.
 - (a) 400 K

(b) 350 K

(c) 298 K

- (d) 250 K
- 10. 2 mol of an ideal gas expands isothermally and reversibly from 1 L to 10 L at 300 K. What is the enthalpy change?
 - (a) 4.98 kJ
- (b) 11.47 kJ
- (c) -11.47 kJ (d) 0 kJ
- 11. (A) follows the first-order reaction (A) \longrightarrow (B). The concentration of (A) changes from 0.1 mol L⁻¹ to 0.025 mol L⁻¹ in 40 min. Find the rate of reaction of (A) when the concentration of (A) is $0.01 \text{ mol } L^{-1}$.

 - (a) $3.47 \times 10^{-4} \text{ mol L}^{-1} \text{ min}^{-1}$ (b) $3.47 \times 10^{-5} \text{ mol L}^{-1} \text{ min}^{-1}$

 - (c) $1.73 \times 10^{-4} \text{ mol L}^{-1} \text{ min}^{-1}$ (d) $1.73 \times 10^{-5} \text{ mol L}^{-1} \text{ min}^{-1}$
- 12. 75.2 g of C_6H_5OH (phenol) is dissolved in a solvent of $K_f = 14 \text{ K kg mol}^{-1}$. If the depression of freezing point is 7 K then find the percentage of phenol that dimerizes.
 - (a) 85
- (b) 75
- (c) 80
- (d) 70

- 13. For the reaction $2\text{CO} + \text{O}_2 \rightarrow 2\text{CO}_2$, $\Delta H = -560 \text{ kJ}$. Two moles of CO and one mole of O_2 are taken in a container of volume 1 L. They completely form two moles of CO_2 . The gases deviate appreciably from their ideal behaviour. If the pressure in the vessel changes from 70 atm to 40 atm, find the magnitude (absolute value) of ΔU at 500 K, given that 1 L atm = 0.1 kJ.
 - (a) 583 kJ

(b) 568 kJ

(c) 563 kJ

- (d) 583 kJ
- **14.** We have taken a saturated solution of AgBr. K_s of AgBr is 12×10^{-14} (mol L⁻¹)². If 10^{-7} mol of AgNO₃ is added to 1 L of this solution, find the conductivity (specific conductance) of this solution in terms of 10^{-7} S m⁻¹, given that $\lambda_{Ag^+}^\circ = 6 \times 10^{-3}$ S m² mol⁻¹, $\lambda_{Br^-}^\circ = 8 \times 10^{-3}$ S m² mol⁻¹, $\lambda_{NO_3}^\circ = 7 \times 10^{-3}$ S m² mol⁻¹.
 - (a) 55
- (b) 65
- (c) 458
- (d) 75
- 15. The edge length of a unit cell of a metal having a molecular weight of 75 g mol⁻¹ is 5 Å which crystallizes in a cubic lattice. If the density is 2 g cm^{-3} then find the radius of the metal atom (given that $N_{\rm A} = 6 \times 10^{23} \, \text{mol}^{-1}$).
 - (a) 218.5 pm

(b) 210.5 pm

(c) 200.5 pm

- (d) 216.5 pm
- 16. Positive deviation from the ideal behaviour takes place because of the
 - (a) molecular interaction between atoms and since $\frac{pV}{nRT} > 1$
 - (b) molecular interaction between atoms and since $\frac{pV}{nRT}$ < 1
 - (c) finite size of atoms and since $\frac{pV}{nRT} > 1$
 - (d) finite size of atoms and since $\frac{pV}{nRT}$ < 1
- 17. $CH_3 \xrightarrow{H^+ -H_2O} (F) \xrightarrow{Br_2/CCl_4} C_4H_8Br_2 \xrightarrow{5 \text{ such products}} 5 \text{ such products}$

How many structures of (F) are possible?

- (a) 2
- (b) 5
- (c) 6
- (d) 3
- **18.** An enantiomerically pure acid is treated with a racemic mixture of an alcohol having one chiral carbon. The ester formed will be
 - (a) an optically active mixture
- (b) a pure enantiomer
- (c) a meso compound
- (d) a racemic mixture

19.	Lyopi	nilic sols are							
	(a)	irreversible sols							
	(b)	prepared from inorganic compounds							
	(c)	coagulated by adding electrolytes							
	(d)	self-stabilizing							
20.		n of the following resonat hoxy-1,3-butadiene is the least		structures of the compound le?					
	(a)	$\stackrel{\scriptscriptstyle{\ominus}}{\mathrm{CH}}_{2}\mathrm{CH}=\!\mathrm{CH}-\!\mathrm{CH}=\!\stackrel{\scriptscriptstyle{\oplus}}{\mathrm{O}}-\!\mathrm{CH}_{3}$							
	(b)	CH_2 = CH - CH - O - CH_3							
	(c)	[⊕] CH ₂ —CH—CH=CH—O—CI	H_3						
	(d)	CH ₂ =CH—CH—CH—O—Cl	H_3						
21.		n the extraction processes listed umn B.	d in o	column A with the metals listed					
		Column A		Column B					
	(i)	Self-reduction	(a)	Lead					
	(ii)	Carbon reduction	(b)	Silver					
	(iii)	Complex formation and	(c)	Copper					
		displacement by the metal							
	(iv)	Decomposition of the iodide	(d)	Boron					
22.		ng the following, identify the tion state.	e spe	ecies with an atom in its +VI					
	(a)	MnO_4^-	(b)	$Cr(CN)_6^{3-}$					
	(c)	NiF_6^{2-}	(d)	CrO_2Cl_2					
23.	Polyp	hosphates are used as water-so	often	ing agents because they					
		form soluble complexes with							
	(b)	precipitate anionic species							
	(c)	form soluble complexes with	catio	onic species					
	(d)	precipitate cationic species							
24.	Identi	fy the correct order of acidic st	reng	th of CO_2 , CuO , CaO and H_2O .					
	(a)	$CaO < CuO < H_2O < CO_2$	(b)	$H_2O < CuO < CaO < CO_2$					
	(c)	$CaO < H_2O < CuO < CO_2$	(d)	$H_2O < CO_2 < CaO < CuO$					
25.	Identi	fy the least stable ion amongst	the f	following.					
	(a)	Li -	(b)	Be -					
	(c)	B ⁻	(d)	C ⁻					

Consider the following equations and answer the questions below (26-28).

 $NiCl_2 + KCN$ (excess) \rightarrow (A) (a cyano complex)

 $NiCl_2 + conc. HCl (excess) \rightarrow (B) (a chloro complex)$

- 26. The IUPAC names for (A) and (B) are respectively
 - (a) Potassium tetracyanonickelate(II) and potassium tetrachloronickelate(II)
 - (b) Tetracyanopotassium nickelate(II) and teterachloropotassium nickelate(II)
 - (c) Tetracyanonickel(II) and tetrachloronickel(II)
 - (d) Potassium tetracyanonickel(II) and potassium tetrachloronickel(II)
- 27. Predict the magnetic natures of (A) and (B).
 - (a) Both are diamagnetic.
 - (b) (A) is diamagntic and (B) is paramagnetic with one unpaired electron.
 - (c) (A) is diamagnetic and (B) is paramagnetic with two unpaired electrons.
 - (d) Both are paramagnetic.
- 28. The hybridization modes of (A) and (B) are respectively
 - (a) dsp^2 and sp^3

- (b) sp^2 and sp^3
- (c) dsp^2 and dsp^2
- (d) sp^3d^3 and d^2sp^3
- **29.** At constant temperature, the equilibrium constant K_p for the decomposition reaction $N_2O_4 \Rightarrow 2NO_2$ is expressed by $K_p = \frac{4x^2p}{1-x^2}$,

where p = pressure and x = extent of decomposition. Which one of the following statements is true?

- (a) K_p increases with increase in p.
- (b) K_p increases with increase in x.
- (c) K_p increases with decrease in x.
- (d) K_p remains constant with change in p and x.
- **30.** In an electrolytic cell, flow of electrons is from
 - (a) cathode to anode in solution
 - (b) cathode to anode through external supply
 - (c) cathode to anode through internal supply
 - (d) anode to cathode through internal supply
- 31. In a first-order reaction, the concentration of the reactants decreases from 800 mol dm^{-3} to 50 mol dm^{-3} in $2 \times 10^4 \text{ s}$. The rate constant for the reaction is

(a) $2 \times 10^4 \,\mathrm{s}^{-1}$

(b) $3.45 \times 10^{-5} \text{ s}^{-1}$

(c) $1.386 \times 10^{-4} \text{ s}^{-1}$

(d) $2 \times 10^{-4} \,\mathrm{s}^{-1}$

32.

$$CH_3$$
 CH_2CH_3

The appropriate reagent required for the above transformation is

(a) Zn/Hg/HCl

(b) NH_2NH_2/OH^-

(c) H₂/Ni

(d) NaBH₄

33.

$$N_2 + 3H_2 \rightleftharpoons 2NH_3$$
.

Which is the correct statement if N_2 is added to the above system at the equilibrium condition?

- (a) The equilibrium will shift to the forward direction because, according to the second law of thermodynamics, the entropy must increase in the direction of the spontaneous reaction.
- (b) The condition for equilibrium is $G_{\rm N_2} + 3G_{\rm H_2} = 2G_{\rm NH_3}$, where G represents the Gibbs free energy per mole of the gaseous species measured at that partial pressure. The equilibrium condition is unaffected by the use of a catalyst, as it increases the rates of both the forward and backward reactions to the same extent.
- (c) The catalyst will increase the rate of the forward reaction by α and that of the backward reaction by β .
- (d) The catalyst will not alter the rate of either of the reactions.
- **34.** The reaction of propene with HOCl proceeds via the addition of
 - (a) H⁺ in the first step
- (b) Cl⁺ in the first step
- (c) OH in the first step
- (d) Cl⁺ and OH⁻ in a single step

The questions below (35–39) consist of an assertion in column A and reason in column B. Use the following key to choose the appropriate answer.

- (a) if both assertion and reason are correct and the reason is the correct explanation of the assertion
- (b) if both assertion and reason are correct but the reason is not the correct explanation of the assertion
- (c) if the assertion is correct but the reason is incorrect
- (d) if the assertion is incorrect but the reason is correct

Column A (assertion)

- 35. In any ionic solid [MX] with Schottky defects, the numbers of positive and negative ions are same.
- 36. Addition of bromine to trans-2butene yields meso-2,3-dibromobutane.
- 37. Between SiCl₄ and CCl₄, only SiCl₄ reacts with water.
- 38. Dimethyl sulphide is commonly used for the reduction of an ozonide of an alkene to get the carbonyl compounds.
- **39.** In strongly acidic solutions. aniline becomes more reactive towards electrophilic reagents.

Column B (reason)

The numbers of cation and anion vacancies are equal.

Bromine addition to an alkene is an electrophilic addition.

SiCl₄ is ionic and CCl₄ is covalent.

It reduces the ozonide, giving water-soluble dimethyl sulphoxide, and excess of it evaporates.

The amino being group completely protonated in strongly acidic solutions, the lone pair of electrons on the nitrogen atom is no longer available for resonance.

- **40.** The order of reactivity of the following alkyl halides for an S_N2 reaction is
 - (a) RF > RCl > RBr > RI
- (b) RF > RBr > RCl > RI
- (c) RCl > RBr > RF > RI
- (d) RI > RBr > RCl > RF
- 41. For the reversible reaction $N_2(g) + 3H_2(g) = 2NH_3(g)$ at 500°C, the value of K_n is 1.44×10^{-5} when partial pressure is measured in atmospheres. The corresponding value of K_c , with concentration expressed in mol L⁻¹,
 - (a) $1.44 \times 10^{-5}/(0.082 \times 500)^{-2}$
- (b) $1.44 \times 10^{-5} / (8.314 \times 773)^{-2}$
- (c) $1.44 \times 10^{-5}/(0.082 \times 773)^2$
- (d) $1.44 \times 10^{-5}/(0.082 \times 773)^{-2}$
- 42. The number of isomers of the compound with the molecular formula C₂BrClFI is
 - (a) 3

(b) 4

(c) 5

- (d) 6
- 43. The compound that will react most readily with NaOH to form methanol is

 - (a) $(CH_3)_4N^+I^-$ (b) CH_3OCH_3 (c) $(CH_3)_3S^+I^-$ (d) $(CH_3)_3CCI$
- 44. Which of the following molecular species has one or more unpaired electron(s)?
 - (a) N_2

(b) F₂

(c) O_2^-

(d) O_2^{2-}

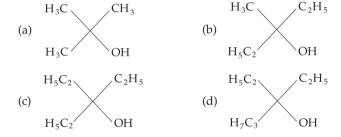
- **45.** The nodal plane in the π bond of ethene is located in
 - (a) the molecular plane
 - (b) a plane parallel to the molecular plane
 - (c) a plane perpendicular to the molecular plane which bisects the C—C σ bond at right angle
 - (d) a plane perpendicular to the molecular plane which contains the C—C $\sigma\,\text{bond}$
- 46. The set representing the correct order of first ionization potential is
 - (a) K > Na > Li

(b) Be > Mg > Ca

(c) B > C > N

- (d) Ge > Si > C
- **47.** The correct order of hybridization of the central atoms in the species NH_3 , $[PtCl_4]^{2-}$, PCl_5 , BCl_3 is
 - (a) dsp^2 , dsp^3 , sp^2 , sp^3
- (b) sp^3, dsp^2, dsp^3, sp^2
- (c) dsp^2 , sp^2 , sp^3 , dsp^3
- (d) dsp^2 , sp^3 , sp^2 , dsp^3
- 48. Ethyl ester $\xrightarrow{\text{excess}}$ (P).

The product (P) will be



49. A mixture of 0.02 mol $[Co(NH_3)_5SO_4]Br$ and 0.02 mol $[Co(NH_3)_5Br]SO_4$ is used to prepare a solution, (X), of 2 L.

1 L (X) + excess AgNO₃
$$\rightarrow$$
 (Y)
1 L (X) + excess BaCl₂ \rightarrow (Z).

The numbers of moles of (Y) and (Z) are respectively

(a) 0.01 and 0.01

(b) 0.02 and 0.01

(c) 0.01 and 0.02

(d) 0.02 and 0.02

HOOC
$$O_{2}N$$

$$CH$$

$$OH$$

$$CH$$

The product (P) will be

(a)
$$O_2N$$
 CH

(b)
$$O_2N$$
 OH C^-

53.

(d)
$$O_2N$$
 C^-

- **51.** HX is a weak acid $(K_a = 10^{-5} \text{ mol L}^{-1})$. It forms the salt NaX (0.1 M) on reaction with caustic soda. The degree of hydrolysis of NaX is
 - (a) 0.01%

(b) 0.0001%

(c) 0.1%

- (d) 0.5%
- 52. Spontaneous adsorption of a gas on a solid surface is an exothermic process because
 - (a) ΔH increases for the system
- (b) ΔS increases for the gas
- (c) ΔS decreases for the gas
- (d) ΔG increases for the gas

COOH

 $(X) + H_2SO_4 \rightarrow (Y)$ (a colourless gas with an irritating smell) $(Y) + K_2Cr_2O_7 + H_2SO_4 \rightarrow a$ green solution.

In the above equations, (X) and (Y) are respectively

(a) SO_3^{2-} and SO_2

(b) Cl and HCl

(c) S^{2-} and H_2S

- (d) CO_3^{2-} and CO_2
- 54. Which of the following reactants on reaction with concentrated NaOH followed by acidification gives the following lactone?

(a)
$$COOCH_3$$
 $COOCH_3$ (b) $COOCH_3$ (c) $COOCH_3$ $COOCH_4$ $COOCH_5$ $COOCH_7$ $COOCH_8$ $COOCH_8$

55.
$$+ Cl - CH_2CH_3 - CH_2 \xrightarrow{AlCl_3} (P) \xrightarrow{(i) O_2/\Delta} (Q) + phenol$$

- 56. Ammonia can be dried by
 - (a) concentrated H₂SO₄
- (b) P₄O₁₀

(c) CaO

- (d) anhydrous CaCl₂
- **57.** Specify the coordination geometry around and hybridization of the N and B atoms in a 1:1 complex of BF₃ and NH₃.
 - (a) N: tetrahedral, sp³; B: tetrahedral, sp³
 - (b) N: pyramidal, sp ³; B: pyramidal, sp ³
 - (c) N: pyramidal, sp ³; B: planar, sp ²
 - (d) N: pyramidal, sp³; B: tetrahedral, sp³
- **58.** The rusting of iron takes place as follows:

$$2H^+ + 2e^- + \frac{1}{2}O_2 \longrightarrow H_2O(1); E^\circ = +1.32 \text{ V}$$

$$Fe^{2+} + 2e^{-} \longrightarrow Fe(s); E^{\circ} = -0.44 \text{ V}.$$

Calculate ΔG° for the net process.

(a) -322 kJ mol^{-1}

(b) -161 kJ mol^{-1}

(c) -152 kJ mol^{-1}

- (d) -76 kI mol^{-1}
- **59.** The elevation of boiling point of a solution of 13.44 g CuCl₂ in 1 kg of water will be (given that the molecular weight of CuCl₂ is 134.4 and $K_b = 0.52$ K kg mol⁻¹)

(a) 0.16 K

occupied?

(a) NaCl

60. The ratio of the rates of diffusion of helium and methane under identical

61. In which of the following crystals are alternate tetrahedral voids

(c) 0.1 K

(c) 1

(c) CaF₂

(d) 0.2 K

(d) 0.5

(d) Na₂O

(b) 0.05 K

conditions of pressure and temperature will be (b) 2

(b) ZnS

62. Which of the following statements is false? (a) Work is a state function. (b) Temperature is a state function.

	(c)	final states are specified.	oietei	y defined when the initial and
	(d)	Work appears at the boundar	ry of	the system.
63.	Me ₂ Si	Cl ₂ on hydrolysis will produce	9	
	(a)	$Me_2Si(OH)_2$	(b)	Me ₂ Si=O
	(c)	$-[-O-Me_2Si-O-]_n$	(d)	Me ₂ SiCl(OH)
64.	with are 10 respective (a)	$10^{-16} \mathrm{M}$ sulphide ion. If the val	lues o , 10 ⁻ tate f (b)	$^+$, Fe $^{2+}$, Zn $^{2+}$ and Hg $^{2+}$ is treated of $K_{\rm s}$ for MnS, FeS, ZnS and HgS 20 (mol L $^{-1}$) 2 and 10^{-54} (mol L $^{-1}$) 2 irst? MgS ZnS
65.	In the	process of extraction of gold, roasted gold ore + CN ⁻ +		$0 \xrightarrow{O_2} (X) + OH^{-1}$ $1 \longrightarrow (Y) + Au$
	Identi	ify the complex ions (X) and (Y)	().	
	(a)	$(X) = [Au(CN)_2]^- $ and $(Y) = [Z_1]^- $	Zn(C	$N_{4}]^{2-}$
	(b)	$(X) = [Au(CN)_4]^{3-}$ and $(Y) = [$	Zn(C	$(2N)_4]^{2-}$
	(c)	$(X) = [Au(CN)_2]^- $ and $(Y) = [Z_1]^- $	Zn(C	$N)_{6}]^{4-}$
	(d)	$(X) = [Au(CN)_4]^- $ and $(Y) = [2]$	Zn(C	$N)_4]^{2-}$
66.	The core is	hemical process involved in th	ne pr	oduction of steel from haematite
	(a)	reduction		
	` '	oxidation		
		reduction followed by oxidat		
	(d)	oxidation followed by reduct	ion	

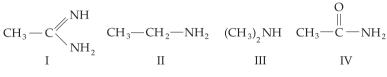
- 67. The number of radial nodes of 3s and 2p orbitals are respectively
 - (a) 2 and 0

(b) 0 and 2

(c) 1 and 2

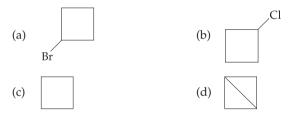
- (d) 2 and 1
- 68. Which species has the maximum number of lone pairs of electrons on the central atom?
 - (a) $[ClO_3]^-$ (b) XeF_4 (c) SF_4

- (d) I₃
- 69. Which is the correct order of basicity of the following compounds?



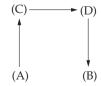
- (a) II > I > III > IV
- (b) I > III > II > IV

- (c) III > I > II > IV
- (d) I > II > III > IV
- 70. What would be the product formed when 1-bromo-3-chlorocyclobutane reacts with two equivalents of metallic sodium in ether?



- 71. In the standardization of Na₂S₂O₃ using K₂Cr₂O₇ by iodometry, the equivalent weight of K₂Cr₂O₇ is
 - (a) (molecular weight)/2 (b) (molecular weight)/6

 - (c) (molecular weight)/3 (d) same as molecular weight
- 72. The direct conversion of (A) to (B) is difficult, hence it is carried out by the following path.



Given that $\Delta S_{(A) \to (C)} = 50 \text{ e.u.}$, $\Delta S_{(C) \to (D)} = 30 \text{ e.u.}$, $\Delta S_{(B) \to (D)} = 20 \text{ e.u.}$, e.u. being the entropy unit, the value of $\Delta S_{(A) \to (B)}$ is

(a) +100 e.u.

(b) +60 e.u.

(c) -100 e.u.

(d) -60 e.u.

CH₃ CH₃ CH₃ 73. CH₂O

$$CH_3O$$
 \longrightarrow H Cl CH_3 CH_3 \longrightarrow NO_2

Which compound(s) among the following is/are given by the above compound on hydrolyis in aqueous acetone?

(I)
$$CH_3O$$
 CH_3 CH

(II)
$$CH_3O$$
 CH_3 C

(III)
$$CH_3O$$
 CH_3 CH_3

- (a) Mixture of I and II
- (b) Mixture of I and III

(c) Only III

(d) Only I

74.

$$\begin{array}{cccc} & & \oplus & & \oplus & \\ H_3 N & & & & NH_3 \\ (X) & & & & (Y) \end{array}$$

Arrange in the increasing order of acidic strength.

(a) (X) > (Z) > (Y)

(b) (Z) < (X) < (Y)

(c) (X) > (Y) > (Z)

- (c) (Z) > (X) > (Y)
- 75. A 0.004-M Na₂SO₄ solution is isotonic with a 0.01-M glucose solution. The degree of dissociation of Na₂SO₄ is
 - (a) 75%
- (b) 50%
- (c) 25%
- (d) 85%
- 76. Cyclohexene on ozonolysis followed by a reaction with Zn dust and water gives a compound (E). (E) on further treatment with aqueous KOH yields a compound (F). The compound (F) is
 - (a) CHO (b) CHO



- 77. Among the following metal carboxyls, the C—O bond order is the lowest in
 - (a) $[Mn(CO)_6]^+$

(b) [Fe(CO)₅]

(c) $[Cr(CO)_6]$

- (d) $[V(CO)_6]^{-}$
- 78. Among the following, the least stable resonance structure is

- **79.** Consider the titration of a potassium dichromate solution with an acidified Mohr salt solution using diphenylamine as the indicator. The number of moles of the Mohr salt required per mole of the dichromate is
 - (a) 3

(b) 4

(c) 5

(d) 6

80. For the process

$$H_2O(1)$$
 (1 bar, 373 K) \rightarrow $H_2O(g)$ (1 bar, 373 K),

the correct set of thermodynamic parameters is

(a) $\Delta G = 0, \Delta S > 0$

(b) $\Delta G = 0, \Delta S < 0$

(c) $\Delta G > 0$, $\Delta S = 0$

- (d) $\Delta G < 0$, $\Delta S > 0$
- 81. A solution of a metal ion when treated with KI gives a red precipitate which dissolves in excess KI to give a colourless solution. Moreover, the solution of the metal ion on treatment with a solution of cobalt(II) thiocyanate gives rise to a deep blue crystalline precipitate. The metal ion is
 - (a) Pb 2+

(b) Hg^{2+}

(c) Cu²⁺

(d) Co²⁺

Answers

1.	a 2.	a 3	. a	4. a	5. c
6.	a 7.	a 8	. C	9. a	10. d
11.	a 12.	b 13	. C	14. a	15. d
16.	a 17.	d 18	. a	19. d	20. d
21.	$(i) \leftrightarrow a$, c $(ii) \leftarrow$	\rightarrow a, c (iii) \leftrightarrow b	$(iv) \leftrightarrow d$		
22.	d 23.	c 24	. a	25. b	26. a
27.	c 28.	a 29	. d	30. c	31. c
32.	b 33.	b 34	. b	35. a	36. b
37.	c 38.	a 39	. d	40. d	41. d
42.	d 43.	c 44	. C	45. a	46. b
47.	b 48.	a 49	. a	50. a	51. a
52.	c 53.	a 54	. C	55. c	56. c
57.	a 58.	a 59	. a	60. b	61. b
62.	a 63.	c 64	. C	65. a	66. c
67.	a 68.	d 69	. b	70. d	71. b
72.	b 73.	a 74	. a	75. a	76. a
77.	d 78.	a 79	. d	80. a	81. b

IIT Questions—2

Useful data

Planck constant, $h = 6.626 \times 10^{-34}$ J s

Atomic numbers: Cr = 24, Mn = 25, Fe = 26, Co = 27, Pt = 781. For H₃PO₃ and H₃PO₄ the correct choice(s) is/are (a) H₃PO₃ is dibasic and reducing (b) H₃PO₄ is dibasic and nonreducing (c) H₃PO₄ is tribasic and reducing (d) H₃PO₃ is tribasic and nonreducing 2. When MnO₂ is fused with KOH, a coloured compound is formed. The product and its colour are respectively (a) K₂MnO₄ and purple-green (b) KMnO₄ and purple (c) Mn₂O₃ and brown (d) Mn₃O₄ and black 3. The name of the structure of silicates in which three oxygen atoms of $[SiO_4]^{4-}$ are shared is (a) pyrosilicate (b) sheet silicate (c) linear-chain silicate (d) three-dimensional silicate 4. Which is the most thermodynamically stable allotropic form of phosphorus? (a) Red phosphorus (b) White phosphorus (c) Black phosphorus (d) Yellow phosphorus 5. Which ore contains both iron and copper? (a) Cuprite (b) Chalcocite (c) Chalcopyrite (d) Malachite 6. Which of the following is not oxidized by O₃? (b) FeSO₄ (d) K_2MnO_4 (a) KI (c) KMnO₄

7.

The product on monobromination of the above compound is

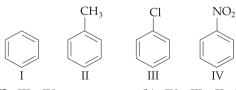
(a)
$$H_3C$$
 CH_3 (b) H_3C CH_3 Br Br

(c)
$$H_3C$$
 CH_3 H_3C H_3C CH_3 Br

- 8. (NH₄)₂Cr₂O₇ on heating gives a gas which is also given by
 - (a) heating NH₄NO₂
- (b) heating NH₄NO₃

(c) $Mg_3N_2 + H_2O$

- (d) $NaNO_3 + H_2O_2$
- **9.** Electrolytic reduction of alumina to aluminium by the Hall–Héroult process is carried out in the presence of
 - (a) NaCl
 - (b) fluorite
 - (c) cryolite, which forms a melt with a lower melting temperature
 - (d) cryolite, which forms a melt with a higher melting temperature
- **10.** Identify the correct order of reactivity in electrophilic substitution reactions of the following compounds:



(a) I > II > III > IV

(b) IV > III > II > I

(c) II > I > III > IV

(d) II > III > I > IV

11.		mide on trea aniline	tmer	nt with POCl		es benzonitrile			
	` '	chlorobenze	ne			benzylamin			
12.	The n	The methods chiefly used for the extraction of lead and tin from the ores are respectively					tin from thei	r	
		self-reduction	-	nd carbon red	luctio	on			
	(b)	self-reduction	n an	d electrolytic	c red	uction			
	(c)	carbon redu	ctior	and self-red	luctio	on			
	(d)	cyanide pro	cess	and carbon r	educ	tion			
13.		cture of benza I solution giv		nyde and for	mald	lehyde on he	ating	; with aqueou	s
	(a)	benzyl alcol	nol a	nd sodium fo	orma	te			
	(b)	sodium ben	zoate	e and methyl	alco	hol			
	(c)	sodium ben	zoate	e and sodium	ı forr	nate			
	(d)	benzyl alcol	nol a	nd methyl al	coho	1			
14.	The co	omplex ion w	hich	has no d ele	ctron	s in the centr	al m	etal atom is	
	(a)	$[MnO_4]^-$			(b)	[Co(NH ₃) ₆]	3+		
	(c)	[Fe(CN) ₆] ³⁻			(d)	$[Cr(H_2O)_6]^3$	+		
15.	Whicl	n blue liquid	is c	btained on	react	ing equimol	ar ar	nounts of tw	o
	gases	at –30°C?							
	(a)	N_2O			(b)	N_2O_3			
	(c)	N_2O_4			(d)	N_2O_5			
16.	When	PbO ₂ reacts	with	concentrated	l HN	IO3, the gas e	volve	ed is	
	(a)	NO_2	(b)	O_2	(c)	N_2	(d)	N_2O	
17.	additi		of K					tate which or ion. The cation	
	(a)	Hg^{2+}	(b)	Bi ³⁺	(c)	Pb ²⁺	(d)	Cu^+	
18.	The n	umber of noc	lal p	lanes in a p _r	orbit	al is			
		one		two		three	(d)	zero	
19.	Anhy	drous ferric c	hlori	ide is prepare	ed by	7			
	(a)	heating hyd of air	rated	d ferric chlor	ide a	t a high temp	erati	ure in a strear	n
	(b)	heating met	allic	iron in a stre	am o	of dry chlorin	e gas	;	
	(c)	reaction of f	erric	oxide with h	ydro	ochloric acid			
	(d)	reaction of r	netal	lic iron with	hydı	rochloric acid	l		

- 20. Identify the correct order of solubility of Na_2S , CuS and ZnS in the aqueous medium.
 - (a) $CuS > ZnS > Na_2S$
- (b) $ZnS > Na_2S > CuS$
- (c) $Na_2S > CuS > ZnS$
- (d) $Na_2S > ZnS > CuS$
- 21. $B(OH)_3 + NaOH \rightleftharpoons NaBO_2 + Na[B(OH)_4] + H_2O$

How can the above reaction is made to proceed in the forward direction?

- (a) Addition of cis-1,2-diol
- (b) Addition of borax
- (c) Addition of trans-1,2 diol
- (d) Addition of Na₂HPO₄

The questions below (22–26) consist of an assertion in column A and reason in column B. Use the following key to choose the appropriate answer.

- (a) if both assertion and reason are correct and the reason is the correct explanation of the assertion
- (b) if both assertion and reason are correct but the reason is not the correct explanation of the assertion
- (c) if the assertion is correct but the reason is incorrect
- (d) if the assertion is incorrect but the reason is correct

Column A (assertion)

Column B (reason)

- **22.** The first ionization energy of Be is greater than that of B.
- **23.** The heat absorbed during the isothermal expansion of an ideal gas against vacuum is zero.
- **24.** Phenol is more reactive than benzene towards electrophilic substitution reactions.
- **25.** 1-Butene on reaction with HBr in the presence of a peroxide produces 1-bromobutane.
- **26.** The pressure of a fixed amount of an ideal gas is proportional to its temperature.

The 2p orbital is lower in energy than the 2s orbital.

The volume occupied by the molecules of an ideal gas is zero.

In the case of phenol, the intermediate carbocation is more resonance-stabilized.

It involves the formation of a primary radical.

The frequency of collisions and their impact increase in proportion to the square root of temperature.

- 27. According to the MO theory,
 - (a) O₂ is paramagnetic and its bond order is greater than that of O₂
 - (b) O_2^+ is paramagnetic and its bond order is less than that of O_2^-
 - (c) O_2^+ is diamagnetic and its bond order is greater than that of O_2
 - (d) O_2^+ is diamagnetic and its bond order is less than that of O_2

Answer the questions below (28-30) using the following information.

Carbon-14 is used to determine the age of organic materials. The procedure is based on the formation of ¹⁴C by neutron capture in the upper atmosphere.

$${}^{14}_{7}N + {}^{1}_{0}n \rightarrow {}^{14}_{6}C + {}^{1}_{1}n$$

¹⁴C is absorbed by living organisms during photosynthesis. The ¹⁴C content is constant in a living organism. Once the plant or animal dies, the uptake of carbon dioxide by it ceases and the level of ¹⁴C in the dead being falls due to the decay which C ¹⁴ undergoes.

$$^{14}_{6}C \rightarrow ^{14}_{7}N + \beta^{-}$$

The half-life of 14 C is 5770 years. The decay constant (λ) can be calculated by using the formula $\lambda = \frac{0.693}{t_{1/2}}$. The comparison of the β^- activity of the dead matter with that of the carbon still in circulation enables measurement of the period of the isolation of the material from the living cycle. The method, however, ceases to be accurate over periods longer than 30,000 years. The ratio of 14 C to 12 C in living matter is $1:10^{12}$.

- **28.** Which of the following options is correct?
 - (a) In living organisms, circulation of ¹⁴C from the atmosphere is high, so the carbon content is constant in an organism.
 - (b) Carbon dating can be used to find out the age of the earth's crust and rocks.
 - (c) Radioactive absorption due to cosmic radiation is equal to the rate of radioactive decay. Hence the carbon content remains constant in living organisms.
 - (d) Carbon dating cannot be used to determine the concentration of ¹⁴C in dead beings.
- 29. What should be the age of a fossil for meaningful determination of its age?
 - (a) 6 years

(b) 6000 years

(c) 60,000 years

- (d) Any age
- **30.** A nuclear explosion has taken place leading to increase in the concentration of 14 C in nearby areas. The 14 C concentration is c_1 in nearby areas and c_2 in areas far away. If the age of the similar fossils is determined to be T_1 in nearby areas and T_2 in faraway areas then
 - (a) the age of the fossils will increase at the place where explosion has taken place and $T_1 T_2 = \frac{1}{\lambda} \ln \frac{c_1}{c_2}$
 - (b) the age of the fossils will decrease at the place where explosion has taken place and $T_1 T_2 = \frac{1}{\lambda} \ln \frac{c_1}{c_2}$
 - (c) the age of the fossils will be determined to be the same
 - (d) $\frac{T_1}{T_2} = \frac{c_1}{c_2}$

32. How many chiral compounds are possible on monochlorination of

(b) Pd/BaSO₄

(d) Pt/H_2

(b) 4

31. 2-Hexyne gives trans-2-hexene on treatment with

(a) Li/NH₃

(c) LiAlH₄

2-methylbutane? (a) 2

	(c)	6	(d)	8
33.	Which	n of the following pairs gives p	ositi	ve Tollens test?
	(a)	Glucose and sucrose	(b)	Glucose and fructose
	(c)	Hexanal and acetophenone	(d)	Fructose and sucrose
34.	Match	the following:		
		Column A		Column B
	(i)	CH ₃ —CHBr—CD ₃ on treatment with alcoholic KOH gives CH ₂ =CH—CD ₃ as the major product	(a)	E1 reaction
	(ii)	Ph—CHBr—CH ₃ reacts faster than Ph—CHBr—CD ₃	(b)	E2 reaction
	(iii)	Ph—CH ₂ —CH ₂ Br on treatment with C_2H_5OD/C_2H_5O gives Ph—CD=CH ₂ as the major product	(c)	E1cB reaction
	(iv)	Ph—CH ₂ —CH ₂ Br and Ph—CD ₂ —CH ₂ Br react at the same rate	(d)	First-order reaction
35.	$E_{X X}^{\circ}$	the electrochemical cell M = 0.33 V. From these data, one $M + X \rightarrow M^+ + X^-$ is the spont	can c	leduce that
	(b)	$M^+ + X^- \rightarrow M + X$ is the spont	taneo	ous reaction
	(c)	$E_{\text{cell}} = +0.77 \text{ V}$		
	(d)	$E_{\text{cell}} = -0.77 \text{ V}$		
36.		nole of a nonideal gas undergo (2.0 atm, 3.0 L, 95 K) -	→ (4.0	0 atm, 5.0 L, 245 K)
		a change in internal energy, (ΔH) of the process is	, Δυ	V = 30.0 L atm. The change in
		40.0 L atm		
		42.3 L atm		
		44.0 L atm		
	(d)	not defined because pressure	is no	ot constant

- 37. When one mole of a monatomic ideal gas at T K undergoes adiabatic change under a constant external pressure of 1 atm, its volume changes from 1 L to 2 L. The final temperature would be
 - (a) $\frac{T}{2^{2/3}}$ K

(b) $\left(T + \frac{2}{3} \times 0.0821\right)$ K

(c) T K

- (d) $\left(T \frac{2}{3} \times 0.0821\right)$ K
- 38. The spin quantum numbers $+\frac{1}{2}$ and $-\frac{1}{2}$ represent respectively
 - (a) the rotation of the electron in clockwise and anticlockwise directions
 - (b) the rotation of the electron in anticlockwise and clockwise directions
 - (c) the magnetic moment of the electron pointing up and down
 - (d) two quantum-mechanical spin states which have no classical analogues
- **39.** Which one of the following statements is not correct for order of reaction?
 - (a) Order can be determined experimentally.
 - (b) Order of reaction is equal to sum of the powers of concentration terms in differential rate law.
 - (c) It is not affected by the stoichiometric coefficient of the reactants.
 - (d) Order cannot be fractional.
- **40.** 0.1 mol of CH₃NH₂ ($K_b = 5 \times 10^{-4}$ mol L⁻¹) is mixed with 0.08 mol of HCl and diluted to one litre. What will be the H⁺ concentration in the solution?
 - (a) $8 \times 10^{-2} \text{ mol L}^{-1}$
- (b) $8 \times 10^{-11} \text{ mol L}^{-1}$
- (c) $1 \cdot 6 \times 10^{-11} \text{ mol L}^{-1}$
- (d) $8 \times 10^{-5} \text{ mol L}^{-1}$
- **41.** Which one of the following will most readily be dehydrated in acidic condition?

42. Consider the following equilibrium in a closed container:

$$N_2O_4(g) \rightleftharpoons 2NO_2(g)$$

At a fixed temperature, the volume of the reaction container is halved. For this change, which of the following statements holds true regarding the equilibrium constant (K_n) and degree of dissociation (α) ?

- (a) Neither K_v nor α changes.
- (b) Both K_n and α change.
- (c) K_n changes, but α does not change.
- (d) K_n does not change, but α changes.

43. 1-Propanol and 2-propanol can be best distinguished by

- (a) oxidation with alkaline KMnO₄ followed by reaction with Fehling's solution
- (b) oxidation with acidic dichromate followed by reaction with Fehling's solution
- (c) oxidation by heating with copper followed by reaction with Fehling's solution
- (d) oxidation with concentrated H₂SO₄ followed by reaction with Fehling's solution

 H_2C OCOC H_3 H_3C OCOC H_3 44. (O)

The products of the acid hydrolysis of (P) and (Q) can be distinguished by

- (a) the Lucas reagent
- (b) 2, 4-DNP
- (c) Fehling's solution
- (d) NaHSO₃

45. Complete the following reaction:

OH
$$+ C_2H_5I \xrightarrow{\text{-OC}_2H_5} \text{anhydrous} C_2H_5OH$$

- (a) C_6H_5 —O— C_2H_5
- (b) C_2H_5 —O— C_2H_5
- (c) C_6H_5 —O— C_6H_5 (d) C_6H_7I

46. A hydroxide which is acidic in water is

- (a) $Al(OH)_3$ (b) $Be(OH)_3$ (c) $La(OH)_3$ (d) $B(OH)_3$

47. Which has the maximum number of atoms?

(a) $24 \text{ g of }^{12}\text{C}$

(b) $56 \text{ g of } ^{56}\text{Fe}$

(c) 27 g of ²⁷Al

(d) $108 \text{ g of }^{108}\text{Ag}$

- **48.** Propyne and propene can be distinguished by
 - (a) concentrated H₂SO₄
- (b) Br₂ in CCl₄

(c) dilute KMnO₄

- (d) AgNO₃ in ammonia
- 49. In the presence of peroxide compounds, hydrogen chloride and hydrogen iodide do not give the anti-Markovnikov addition to alkenes because
 - (a) both are highly ionic
 - (b) one is oxidizing and the other is reducing
 - (c) one of the steps is endothermic in both the cases
 - (d) all the steps are exothermic in both the cases
- 50. If CuSO₄ decolorizes on addition of KCN,
 - (a) the product is $[Cu(CN)_4]^{2-}$
 - (b) Cu^{2+} gets reduced to form $[Cu(CN)_4]^{3-}$
 - (c) the product is $Cu(CN)_2$
 - (d) CuCN is produced
- 51. The wavelength associated with a golf ball weighing 200 g and moving at a speed of 5 m/h is of the order
- (a) 10^{-10} m (b) 10^{-20} m (c) 10^{-30} m (d) 10^{-40} m
- 52. An S_N2 reaction at an asymmetric carbon of a compound always gives
 - (a) an enantiomer of the substrate
 - (b) a product with opposite optical rotation
 - (c) a mixture of diastereoisomers
 - (d) a single stereoisomer

53.



Hydrogenation of the above compound in the presence of poisoned palladium catalyst gives

- (a) an optically active compound (b) an optically inactive compound
- (c) a racemic mixture
- (d) a diastereoisomeric mixture
- **54.** The electronic configuration of an element is $1s^2 2s^2 2p^6 3s^2 3p^6 3d^5 4s^1$. This represents its
 - (a) excited state

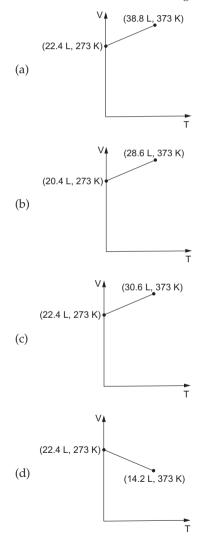
(b) ground state

(c) cationic form

- (d) anionic form
- 55. The rms velocity of hydrogen is $\sqrt{7}$ times the rms velocity of nitrogen. If *T* stands for temperature then

- (a) $T_{H_2} = T_{N_2}$ (b) $T_{H_2} > T_{N_2}$ (c) $T_{H_3} < T_{N_2}$ (d) $T_{H_3} = \sqrt{7}T_{N_2}$

- **56.** What are the common features among the species CN⁻, CO and NO⁺?
 - (a) They have the bond order 3 and are isoelectronic.
 - (b) They have the bond order 3 and are weak-field ligands.
 - (c) They are π -acceptors having the bond order 2.
 - (d) They are isoelectronic and are weak-field ligands.
- **57.** Which of the following volume–temperature (V-T) plots represents the behaviour of one mole of an ideal gas at one atmospheric pressure?



		IIT Questions—2	5-59			
58.	A substance $A_x B_y$ crystallizes in a face-centred-cubic (fcc) lattice in which the A atoms occupy each corner of the cube and the B atoms occupy the centres of each face of the cube. Identify the correct composition of the substance $A_x B_y$.					
	(a)	AB_3				
	(b)	A_4B_3				
	(c)	A_3B				
	(d)	Its composition cannot be specified.				
59.	0.0006	density of liquid water is 1.0 g cm ⁻³ and that of water vapor 6 g cm ⁻³ at 100°C and 1 atm then the volume occupied by values in 1 L of steam at that temperature and pressure is	ur is vater			

60. How many moles of electrons weigh one kilogram?

(b) 60 cm^3

(a)
$$6.023 \times 10^{23}$$
 (b) $\frac{1}{9.108} \times 10^{31}$

(c)
$$\frac{6.023}{9.108} \times 10^{54}$$
 (d) $\frac{1}{9.108 \times 6.023} \times 10^{8}$

- **61.** The IUPAC name for C_6H_5COCl is
 - (a) benzoyl chloride (b) benzene chloroketone
 - (c) benzene carbonyl chloride (d) chlorophenyl ketone

62.
$$Ag^{+} + NH_{3} \rightleftharpoons [AgNH_{3}]^{+}; k_{1} = 3.5 \times 10^{-3}$$
$$[AgNH_{3}]^{+} + NH_{3} \rightleftharpoons [Ag(NH_{3})_{2}]^{+}; k_{2} = 1.7 \times 10^{-3}$$

Then the formation constant of [Ag(NH₃)₂] ⁺ is approximately

(a) 6.0×10^{-6}

(a) 6 cm³

(b) 6.0×10^6

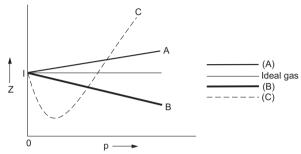
(c) 0.6 cm^3

(d) $0.06 \,\mathrm{cm}^3$

(c) 6.0×10^{-9}

- (d) none of the above
- 63. In thermodynamics, a process is called reversible when
 - the surroundings and the system change into each other
 - (b) there is no boundary between the system and the surroundings
 - the surroundings are always in equilibrium with the system
 - (d) the system changes into the surroundings spontaneously
- **64.** $CH_3NH_2 + CHCl_3 + KOH \rightarrow a$ nitrogen-containing compound $+ KCl + H_2O$. The nitrogen-containing compound is
 - (a) $CH_3 C \equiv N$ (b) CH₃—NH—CH₃
 - (c) $CH_2 \bar{N} \equiv \dot{C}$ (d) $CH_2N = \bar{C}$
- 65. Which of the following has the —O—O— linkage?
 - (a) $H_2S_2O_6$ (b) $H_2S_2O_8$
 - (c) H₂S₂O₂ (d) $H_2S_4O_6$

- **66.** When I is oxidized by MnO₄ in an alkaline medium, I converts into
 - (a) IO_3^-
- (b) I₂
- (c) IO₄
- (d) IO
- **67.** The following graph represents the variation of the compressibility factor, $Z = \frac{pV}{nRT}$, versus p, for three real gases (A), (B) and (C). Identify the incorrect statement.



- (a) For the gas (A), a = 0 and its dependence on p is linear at all values of pressure.
- (b) For the gas (B), b = 0 and its dependence on p is linear at all values of pressure.
- (c) For the gas (*C*), neither a nor b is equal to zero. By knowing the minima and the point of intersection, with Z = 1, a and b can be calculated.
- (d) At high pressure, the slope is positive for all real gases.
- **68.** The rate constant for the reaction $2N_2O_5 \rightarrow 4NO_2 + O_2$ is 3.0×10^{-5} s⁻¹. If the rate is 2.40×10^5 mol L ⁻¹ s ⁻¹ then the concentration of N_2O_5 is
 - (a) 1.4 mol L^{-1}

(b) 1.2 mol L^{-1}

(c) 0.04 mol L^{-1}

- (d) 0.8 mol L^{-1}
- **69.** Standard electrode potential data are useful for understanding the suitability of an oxidant in a redox titration. Some half-cell reactions and their standard potentials are given below:

$$\begin{split} MnO_4^-(aq) + 8H^+(aq) + 5e^- &\to Mn^{2+}(aq) + 4H_2O(l); E^\circ = +1.51 \ V \\ Cr_2O_7^{2-}(aq) + 14H^+(aq) + 6e^- &\to 2Cr^{3+}(aq) + 7H_2O(l); E^\circ = +1.38 \ V \\ Fe^{3+}(aq) + e^- &\to Fe^{2+}(aq); E^\circ = +0.77 \ V \\ Cl_2(g) + 2e^- &\to 2Cl^-(aq); E^\circ = +1.40 \ V. \end{split}$$

Now identify the only incorrect statement regarding the quantitative estimation of aqueous Fe(NO₃)₂.

- (a) MnO_4^- can be used in aqueous HCl.
- (b) $Cr_2O_7^{2-}$ can be used in aqueous HCl.
- (c) MnO₄ can be used in aqueous H₂SO₄.
- (d) $Cr_2O_7^{2-}$ can be used in aqueous H_2SO_4 .

- **70.** In a solid AB having the NaCl structure, A atoms occupy the corners of the cubic unit cell. If all the face-centred atoms along one of the axes are removed then the resultant stoichiometry of the solid is
 - (a) AB₂
- (b) A₂B
- (c) A_4B_3
- (d) A_3B_4
- 71. When diluted with H₂O and boiled, a solution gives a white precipitate. On addition of excess NH₄Cl/NH₄OH, the volume of the precipitate decreases leaving behind a white gelatinous precipitate. Identify the precipitate which dissolves in NH₄Cl/NH₄OH.
 - (a) $Zn(OH)_2$

(b) $Al(OH)_3$

(c) $Mg(OH)_2$

- (d) $Ca(OH)_2$
- **72.** When benzenesulphonic acid and p-nitrophenol are treated with NaCO₃, the gases released respectively are
 - (a) SO₂ and NO₂

(b) SO₂ and NO

(c) SO₂ and CO₂

- (d) CO₂ and CO₂
- **73.** A monatomic ideal gas undergoes a process in which the ratio of *P* to *V* at any instant is constant and equals 1. What is the molar heat capacity of the gas?
 - (a) $\frac{4R}{2}$

(b) $\frac{3R}{2}$

(c) $\frac{5R}{2}$

- (d) 0
- 74. The smallest ketone and its next homologue are reacted with NH₂OH to form oximes. Then
 - (a) two different oximes are formed
 - (b) three different oximes are formed
 - (c) two oximes are optically active
 - (d) all oximes are optically active

75.

$$H_3C$$

$$\begin{array}{c} CH_3 \\ \hline \\ CH_3 \end{array}$$

$$\begin{array}{c} hv \\ \hline \\ Cl_2 \end{array}$$
 N isomeric products

$$C_5H_{11}Cl$$
 fractional M isomeric products

What are the values of N and M?

- (a) 6 and 6
- (b) 6 and 4
- (c) 4 and 4
- (d) 3 and 3
- **76.** Consider a reaction $a(G) + b(H) \rightarrow (P)$. When the concentrations of both the reactants (G) and (H) are doubled, the rate increases by eight times. However, when the concentration of (G) is doubled keeping the concentration of (H) fixed, the rate is doubled. The overall order of the reaction is
 - (a) 0
- (b) 1
- (c) 2
- (d) 3

80. d

81. b

- Chemistry MCO 77. The number of stereoisomers obtained by the bromination of trans-2-butene is (a) 1 (b) 2 (c) 3 (d) 4 78. A positron is emitted from $^{33}_{11}$ Na. The ratio of the atomic mass and atomic number of the resulting nuclide is (c) 23:10 (a) 22:10 (b) 22:11 (d) 23:12 79. A species having a bond order different from that in CO is (a) NO (b) NO⁺ (c) CN⁻ (d) N_2 80. Among the following the paramagnetic compound is (c) N₂O (d) KO_2 (a) Na_2O_2 (b) O₃ 81. The extraction of zinc from zinc blende is achieved by (a) electrolytic reduction (b) roasting followed by reduction with carbon (c) roasting followed by reduction with another metal (d) roasting followed by self-reduction Answers 3. b 5. b 1. a, c **2.** a **4.** c **6.** c 7. b **9.** c **8.** a **10.** c **11.** b **12.** a **13.** a **14.** a 15. b **16.** b **17.** b **18.** a **19.** b **20.** d **21.** a **22.** a **23.** c **24.** a **25.** c **26.** b **27.** a **28.** c **29.** b **30.** a **32.** b **33.** b **31.** a
- **34.** (i) \leftrightarrow b (ii) \leftrightarrow b (iii) \leftrightarrow c, d (iv) \leftrightarrow a, d 35. b **36.** c **39.** d **37.** a **38.** d **40.** b **42.** d **41.** a **43.** c **44.** c **45.** b **46.** d **47.** a **48.** d **49.** c **50.** d **51.** c **52.** d **53.** b **54.** b **59.** c **55.** c **56.** a **57.** c **58.** a **61.** c **64.** d **60.** d **62.** a **63.** c 65. b **66.** a 67. b **68.** d **69.** a **70.** d **71.** a **72.** d **73.** a **74.** b **77.** a 75. b **76.** d **79.** a **78.** c

IIT Questions—3

Useful data

Mass of an electron, $m_e = 9.108 \times 10^{-31} \text{ kg}$

Avogadro constant, $N_A = 6.023 \times 10^{23} \text{ mol}^{-1}$

Molar gas constant, R = 0.082 L atm K $^{-1}$ mol $^{-1}$

Atomic numbers: Li = 3, Be = 4, B = 5, C = 6

- 1. The molecular shapes of SF_4 , CF_4 and SeF_4 are
 - (a) similar, with 2, 0 and 1 lone pairs of electrons, respectively
 - (b) similar, with 1, 1 and 1 lone pairs of electrons, respectively
 - (c) different, with 0, 1 and 2 lone pairs of electrons, respectively
 - (d) different, with 1, 0 and 2 lone pairs of electrons, respectively
- 2. (I) 1,2-Dihydroxybenzene
 - (II) 1,3-Dihydroxybenzene
 - (III) 1,4-Dihydroxybenzene
 - (IV) Hydroxybenzene

The increasing order of boiling point of the above-mentioned alcohols is

(a) I < II < III < IV

(b) I < II < IV < III

(c) IV < I < II < III

(d) IV < II < I < III

3.
$$CH_3-CH=CH_2+NOCl\rightarrow (P)$$

Identify the adduct (P).

(c) NO (d)
$$CH_2 - CH_2 - CH$$
 $CH_3 - CH_2 - CH$ NO CI

- 4. Among the following, which is the strongest base?
 - (a) C_6H_5 — NH_2

- (b) $p-(NO_2-C_6H_4-NH_2)$
- (c) m-(NO₂—C₆H₄—NH₂)
- (d) C_6H_5 — CH_2 — NH_2
- 5. Which of the following compounds will exhibit geometrical isomerism?
 - (a) 1-Phenyl-2-butene
- (b) 3-Phenyl-1-butene
- (c) 2-Phenyl-1-butene
- (d) 1,1-Diphenyl-1-propene
- 6. Consider the chemical reaction N₂(g) + 3H₂(g) → 2NH₃(g). The rate of this reaction can be expressed in terms of the time derivative of concentration of N₂(g), H₂(g) or NH₃(g). Identify the correct relationship amongst the following rate expressions.

(a) Rate =
$$-\frac{d[N_2]}{dt} = -\frac{1}{3} \cdot \frac{d[H_2]}{dt} = \frac{1}{2} \cdot \frac{d[NH_3]}{dt}$$

(b) Rate =
$$-\frac{d[N_2]}{dt}$$
 = $-3\frac{d[H_2]}{dt}$ = $2\frac{d[NH_3]}{dt}$

(c) Rate =
$$\frac{d[N_2]}{dt} = \frac{1}{3} \cdot \frac{d[H_2]}{dt} = \frac{1}{2} \cdot \frac{d[NH_3]}{dt}$$

(d) Rate =
$$-\frac{d[N_2]}{dt} = -\frac{d[H_2]}{dt} = \frac{d[NH_3]}{dt}$$

- 7. The set with correct order of acidity is
 - (a) HClO < HClO₂ < HClO₃ < HClO₄
 - (b) HClO₄ < HClO₃ < HClO₂ < HClO
 - (c) HClO < HClO₄ < HClO₃ < HClO₂
 - (d) HClO₄ < HClO₂ < HClO₃ < HClO
- 8. Among the following the molecule with the highest dipole moment is
 - (a) CH₃Cl

(b) CH₂Cl₂

(c) CHCl₃

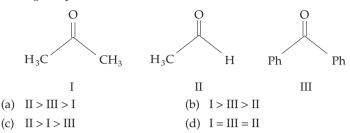
- (d) CCl₄
- 9. Which of the following are isoelectronic as well as isostructural?
 - (a) NO_3^- , CO_3^{2-}

(b) SO_3 , NO_3^-

(c) ClO_3^-, CO_3^{2-}

(d) CO_3^{2-} , SO_3

10. Find the order of reactivity of phenyl magnesium bromide with the following compounds:



• The Tollens reagent is used for the detection of aldehyde. When a solution of AgNO₃ is added to glucose with NH₄OH, gluconic acid is formed.

$$Ag^{+} + e^{-} \rightarrow Ag; E^{\circ}_{red} = +0.8 \text{ V}$$

$$C_{6}H_{12}O_{6} + H_{2}O \rightarrow C_{6}H_{12}O_{7} \text{ (gluconic acid)} + 2H^{+} + 2e^{-}; E^{\circ}_{ox} = -0.05 \text{ V}$$

$$Ag(NH_{3})_{2}^{+} + e^{-} \rightarrow Ag(s) + 2NH_{3}; E^{\circ}_{red} = +0.337 \text{ V}$$
Use $2.303 \times \frac{RT}{F} = 0.0592$ and $\frac{F}{RT} = 38.92$ at 298 K .

11. $2Ag^+ + C_6H_{12}O_6 + H_2O \rightarrow 2Ag(s) + C_6H_{12}O_2 + 2H^+$. Find ln *K* for this reaction.

(a) 66.13

(b) 58.38

(c) 28,30

- (d) 46.29
- **12.** When ammonia is added to the solution, its pH is raised to 11. Which half-cell reaction is affected by pH and by how much?
 - (a) E_{ox} will increase by a factor of 0.65 from E_{ox}° .
 - (b) E_{ox} will decrease by a factor of 0.65 from E_{ox}° .
 - (c) E_{red} will increase by a factor of 0.65 from E_{red}° .
 - (d) E_{red} will decrease by a factor of 0.65 from E_{red}° .
- **13.** Ammonia is always is added in this reaction. Which of the following must be incorrect?
 - (a) NH₃ combines with Ag ⁺ to form a complex.
 - (b) Ag(NH₃)₂⁺ is a stronger oxidizing reagent than Ag ⁺.
 - (c) In absence of NH₃, silver salt of gluconic acid is formed.
 - (d) NH₃ has affected the standard reduction potential of the glucose/gluconic acid electrode.

• RCONH₂ is converted into RNH₂ by means of the Hofmann bromamide degradation.

$$Cl \xrightarrow{O} Cl \xrightarrow{O} Cl \xrightarrow{NH_2} Cl \xrightarrow{N} -Bi$$

$$III \xrightarrow{NH_2} Cl \xrightarrow{N} Cl \xrightarrow{N} - Cl$$

$$H_2N \xrightarrow{O} Cl \xrightarrow{N} - Cl \xrightarrow{N} - Cl$$

In this reaction, RCONHBr is formed, from which this reaction has derived its name. The electron-donating group at phenyl activates the reaction. The Hofmann degradation reaction is an intramolecular reaction.

- **14.** What can bring about the conversion of I to II?
 - (a) KBr

(b) KBr + CH₃ONa

(c) KBr + KOH

- (d) $Br_2 + KOH$
- **15.** Which is the rate-determining step in the Hofmann bromamide degradation?
 - (a) Formation of I

- (b) Formation of II
- (c) Formation of III
- (d) Formation of IV

16.

$$CO-NH_2$$
 $CO-^{15}NH_2$

What are the constituent amines formed when the mixture of I and II undergoes the Hofmann bromamide degradation?

(a)
$$NH_2$$
, NH_2 , NH_2 , NH_2 , NH_2 , NH_2 , (b) NH_2 , NH_2

(c)
$$\sqrt{}$$
 $^{15}\text{NH}_2$, $\sqrt{}$ $^{15}\text{NH}_2$ (d) $\sqrt{}$ ^{15}NHD , $\sqrt{}$

17. 4-Methylbenzenesulphonic acid reacts with sodium acetate to give

18. Match the following columns: Column A

CH₃

(i) $Br^{3+} \rightarrow [BiO]^+$ (a) Heat (ii) $[AlO_2]^- \rightarrow Al(OH)_3$ (b) Hydrolysis (iii) $[SiO_4]^{4-} \rightarrow [Si_2O_7]^{4-}$ (c) Acidification (iv) $[B_4O_7]^{2-} \to B(OH)_3$ (d) Dilution by water 19. The best method to prepare cyclohexene from cyclohexanol is by using (a) concentrated HCl and ZnCl₂ (b) concentrated H₃PO₄ (c) HBr (d) concentrated HCl 20. What can convert butan-2-one to propanoic acid?

Column B

- - (a) Tollens reagent (b) Fehling's solution

(c) NaOH/I₂/H⁺

- (d) NaOH/NaI/H+
- 21. MeO CHO + (X) $\frac{CH_3COON_a}{H_3O^+}$ CH=CHCOOH

The compound (X) is

(a) CH₃COOH

(b) BrCH2-COOH

(c) $(CH_3CO)_2O$

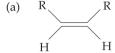
- (d) CHO-COOH
- **22.** According to Bohr's theory, E_n = total energy, K_n = kinetic energy, V_n = potential energy and r_n = radius of the nth orbit. Now match the following columns:

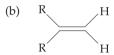
Column A

Column B

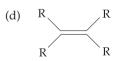
(i) V_n/K_n equals

- (a) 0
- (ii) If $r_n \propto E_n^x$ then x is
- (b) -1
- (iii) The angular momentum in the lowest orbital is
- (c) -2
- (iv) If $\frac{1}{r^n} \propto Z^y$ then y equals
- (d) 1
- **23.** Which one of the following alkenes will react fastest with H₂ under the catalytic hydrogenation condition?





(c) R R



- **24.** An aqueous solution of 6.3 g oxalic acid dihydrate is made up to 250 mL. The volume of the 0.1-N NaOH solution required to completely neutralize 10 mL of this solution is
 - (a) 40 mL

(b) 20 mL

(c) 10 mL

- (d) 4 mL
- **25.** Identify from the following list the reagents which can easily distinguish between 1-butyne and 2-butyne.
 - (a) Bromine, CCl₄

- (b) H₂, Lindlar catalyst
- (c) Dilute H₂SO₄, HgSO₄
- (d) Ammoniacal Cu₂Cl₂ solution
- **26.** The number of electrons involved in the conversion of OH to O₂ is
 - (a) one

(b) four

(c) two

(d) three

27. The pair of compounds having metals in their highest oxidation states is

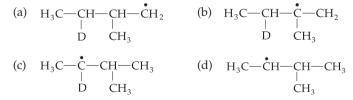
(a) MnO₂, FeCl₃ (b) $[MnO_4]^-$, CrO_2Cl_2 (d) $[NiCl_4]^{2-}$, $[CoCl_4]^{+}$ (c) $[Fe(CN)_6]^{3-}$, $Co(CN)_2$ 28. The compound having a tetrahedral geometry is (a) $[Ni(CN)_4]^{2-}$ (b) [Pd(CN)₄]²⁻ (d) [PdCl₄1²⁻ (d) $[NiCl_{4}l^{2-}]$ 29. Which hydrogen-like species has the same radius as that of the Bohr orbit of the hydrogen atom? (b) n = 2. Be ³⁺ (a) n = 2. Li²⁺ (d) $n = 3 \text{ Li}^{2+}$ (c) n = 2. He⁺ 30. The hybridization modes of atomic orbitals of nitrogen in NO₂⁺, NO₃⁻ and NH₄ are respectively (a) sp, sp 3 and sp 2 (b) sp, sp^2 and sp^3 (c) sp^2 , sp and sp^3 (d) sp^2 , sp^3 and sp31. The chemical composition of the 'slag' formed during the smelting process in the extraction of copper is (c) $CuFeS_2$ (d) $Cu_2S + FeO$ (a) $Cu_2O + FeS$ (b) $FeSiO_3$ 32. Which of the following acids has the smallest dissociation constant? (a) CH₃CHFCOOH (b) FCH2CH2COOH (c) BrCH2CH2COOH (d) CH₃CHBrCOOH 33. Consider the following reaction:

$$H_3C$$
— CH — CH — $CH_3 + \overset{\bullet}{B}r \longrightarrow (X) + HBr$
 $D \quad CH_3$

Identify the structure of the major product (X).

(b) $\sqrt{15}$

(a) $\sqrt{3}$



- 34. The spin-only magnetic moment of the compound Hg[Co(SCN)₄] is
- **35.** A sodium salt having an unknown anion when treated with MgCl₂ gives a white precipitate on boiling. The anion is

(c) $\sqrt{24}$

(d) √8

(a) SO_4^{2-} (b) HCO_3^{-} (c) CO_3^{2-} (d) NO_3^{-}

- **36.** The correct order of radius is
 - (a) N < Be < B

(b) $F^- < O^{2-} < N^{3-}$

(c) Na < Li < K

- (d) $Fe^{3+} < Fe^{2+} < Fe^{4+}$
- 37. The $\Delta H_{\rm f}^{\circ}$ for ${\rm CO_2(g)}$, ${\rm CO(g)}$ and ${\rm H_2O(g)}$ are -393.5 kJ ${\rm mol}^{-1}$, -110.5 kJ ${\rm mol}^{-1}$ and -241.8 kJ ${\rm mol}^{-1}$ respectively. The standard enthalpy change for the reaction ${\rm CO_2(g)} + {\rm H_2(g)} \rightarrow {\rm CO(g)} + {\rm H_2O(g)}$ is
 - (a) 524.1 kJ

(b) 41.2 kJ

(c) -262.5 kJ

- (d) -41.2 kJ
- **38.** Identify the set of reagents and/or reaction conditions (X) and (Y) in the following set of transformations:

$$CH_3 - CH_2 - CH_2Br \xrightarrow{(X)} (P) \xrightarrow{(Y)} CH_3 - CH - CH$$

- (a) $(X) = \text{dilute aqueous NaOH}, 20^{\circ}\text{C}; (Y) = \text{HBr/acetic acid}, 20^{\circ}\text{C}$
- (b) (X) = concentrated alcoholic NaOH, 80°C; (Y) = HBr/acetic acid, 20°C
- (c) (X) = dilute aqueous NaOH, 20° C; (Y) = $Br_2/CHCl_3$, 0° C
- (d) (X) = concentrated alcoholic NaOH, 80° C; (Y) = $Br_2/CHCl_3$, 0° C
- **39.** Among H_2O , H_2S , H_2Se and H_2Te , the one with the highest boiling point is
 - (a) H₂O because of its hydrogen bonding
 - (b) H₂Te because of its higher molecular weight
 - (c) H₂S because of its hydrogen bonding
 - (d) H₂Se because of its lower molecular weight
- **40.** A compound (A) having the molecular formula C_3H_8O is treated with acidified potassium dichromate to form a product (B) having the molecular formula C_3H_6O . (B) forms a shining silver mirror on warming with ammoniacal silver nitrate. (B) when treated with an aqueous solution of $H_2NCONHNH_2$ ·HCl and sodium acetate gives a product (C). Identify the structure of (C).
 - (a) CH₃CH₂CH=NNHCONH₂
 - (b) $CH_3 C = NNHCONH_2$ CH_3
 - (c) $CH_3-C=NCONHNH_2$ CH_3
 - (d) CH₃CH₂CH=NCONHNH₂

41. Which of the following hydrocarbons has the lowest dipole moment?

42. Which of the following has the highest nucleophilicity?

(c) CH₃CH₂C≡CH

(b) CH₃C=CCH₃

(d) CH₂=CH—C≡CH

(b) OH^{-} (c) CH_{3}^{-} (d) NH_{2}^{-}

43.	Benzo	lyl chloride is prepared from b	enzo	ic acid using
	(a)	heated Cl ₂	(b)	SO ₂ Cl ₂
	(c)	SOCl ₂	(d)	Cl ₂ and H ₂ O
44.	The co	ompressibility of a gas is less tl	าan เ	unity at stp. Therefore,
		$V_{\rm m} > 22.4 \; {\rm L \; mol}^{-1}$		$V_{\rm m}$ < 22.4 L mol ⁻¹
	(c)	$V_{\rm m} = 22.4 \; {\rm L \; mol^{-1}}$	(d)	$V_{\rm m}$ = 44.8 L mol ⁻¹
45.		n of the following processes is esium?	used	d in the extractive metallurgy of
	(a)	Fused-salt electrolysis	(b)	Self-reduction
	(c)	Aqueous-solution electrolysis	(d)	Thermite reduction
46.	treatn When black (a)	nent with dilute hydrochloric	acie d the bstar (b)	gives a white precipitate on d, which dissolves on heating rough the hot acidic solution, ance is $a(n)$ Cu^{II} salt Pb^{II} salt
47.	aqueo precip ribbon (a)	bus solution on treatment voltate. The saturated aqueous in with evolution of a colourless $(X) = CO_2$; $(Y) = Cl_2$	with solu s gas (b)	form a saturated solution. The silver nitrate gives a white tion also dissolves magnesium (Y). Identify (X) and (Y). (X) = Cl_2 ; (Y) = CO_2 (X) = H_2 ; (Y) = Cl_2
48.	If <i>I</i> is the pl	the intensity of absorbed light	and	c is the concentration of AB for B*, the rate of formation of AB*
	(a)	c (b) I	(c)	I^2 (d) $c \cdot I$
49.		orrect order of equivalent cor re dilution is	nduc	tance of LiCl, NaCl and KCl at
	(a)	LiCl > NaCl > KCl	(b)	KCl > NaCl > LiCl
	(c)	NaCl > KCl > LiCl	(d)	LiCl > KCl > NaCl

50. For a monatomic gas, kinetic energy = E. Its relation with rms velocity (u) is

(a)
$$u = \left(\frac{2E}{m}\right)^{1/2}$$
 (b) $u = \left(\frac{3E}{2m}\right)^{1/2}$ (c) $u = \left(\frac{E}{2m}\right)^{1/2}$ (d) $u = \left(\frac{E}{3m}\right)^{1/2}$

- 51. The correct order of acidic strength is
 - (a) $Cl_2O_7 > SO_2 > P_4O_{10}$
- (b) $CO_2 > N_2O_5 > SO_3$
- (c) $Na_2O > MgO > Al_2O_3$
- (d) $K_2O > CaO > MgO$
- 52. Which of the following compounds exhibits stereoisomerism?
 - (a) 2-methyl-1-butene
- (b) 3-methyl-1-butyne
- (c) 3-methylbutanoic acid
- (d) 2-methylbutanoic acid
- 53. The root-mean-square velocity of an ideal gas at constant pressure varies with its density d as
 - (a) d^{2}
- (b) *d*
- (c) √*d*
- (d) $\frac{1}{\sqrt{d}}$
- 54. The reaction $3ClO^{-}(aq) \rightarrow ClO_{3}^{-}(aq) + 2Cl^{-}(aq)$ is an example of
 - (a) oxidation reaction
- (b) reduction reaction
- (c) disproportionation reaction
- (d) decomposition reaction

55. CHO OHC $(i) \text{ NaOH/}100^{\circ}\text{C} \rightarrow \text{ different products}$ $(i) \text{ NaOH/}100^{\circ}\text{C} \rightarrow \text{ different products}$ $(ii) \text{ H}^{+}/\text{H}_{2}\text{O} \rightarrow \text{ different products}$

The major product of the above reaction is

- **56.** When two reactants (A) and (B) are mixed to give the products (C) and (D), the reaction quotient (*Q*) at the initial stages of the reaction
 - (a) is zero

(b) decreases with time

(c) is independent of time

- (d) increases with time
- 57. When temperature is increased, surface tension of water
 - (a) increases

(b) decreases

(c) remains constant

- (d) shows irregular behaviour
- 58. Rutherford's experiment, which established the nuclear model of the atom, used a beam of
 - (a) β-particles, which impinged on a metal foil and got absorbed
 - (b) γ-rays, which impinged on a metal foil and ejected electrons
 - (c) helium atoms, which impinged on a metal foil and got scattered
 - (d) helium nuclei, which impinged on a metal foil and got scattered

59. $Zn \mid Zn^{2+}(0.1 \text{ M}) \parallel Fe^{2+}(0.01 \text{ M}) \mid Fe.$

The emf of the above cell is 0.2905 V. The equilibrium constant for the cell reactions is

(a) $10^{0.32/0.0591}$

(b) 10^{0.32/0.0295}

(c) 10^{0.26/0.0295}

(d) e 0.32/0.295

- **60.** Saturated solutions of KNO_3 are used to make salt bridges because
 - (a) velocity of K + is greater than that of NO₃
 - (b) velocity of NO_3^- is greater than that of K^+
 - (c) velocities of K⁺ and NO₃⁻ are nearly the same
 - (d) KNO₃ is highly soluble in water
- **61.** During depression of freezing point in a solution, which of the following are in equilibrium?
 - (a) Liquid solvent, solid solvent
 - (b) Liquid solvent, solid solute
 - (c) Liquid solute, solid solute
 - (d) Liquid solute, solid solvent
- **62.** H_3BO_3 is a
 - (a) monobasic and weak Lewis acid
 - (b) monobasic and weak Brønsted acid
 - (c) monobasic and strong Lewis acid
 - (d) tribasic and weak Brønsted acid

63.
$$Ph - C = C - CH_{3} \xrightarrow{Hg^{2*}/H^{+}} (A).$$
In the above reaction, (A) is

(a) O

$$Ph \longrightarrow O$$

$$H_{3}C$$
(c) O
$$Ph \longrightarrow O$$

$$Ph \longrightarrow O$$

$$H_{3}C$$
(d) Ph \longrightarrow OH

- **64.** Which of the following has the most acidic hydrogen?
 - (a) 3-hexanone

- (b) 2,4-hexanedione
- (c) 2,5-hexanedione
- (d) 2,3-hexanedione
- 65. If the nitrogen atom had electronic configuration 1s⁷, it would have energy lower than that of the normal ground-state configuration 1s²2s²2p³ because the electrons would be closer to the nucleus. Yet, 1s⁷ is not observed because it violates
 - (a) the Heisenberg uncertainty principle
 - (b) the Hund rule
 - (c) the Pauli exclusion principle
 - (d) Bohr's postulate of stationary orbits
- **66.** When phenyl magnesium bromide reacts with butanol, the product would be
 - (a) benzene

(b) phenol

(c) *t*-butylbenzene

- (d) *t*-butyl phenyl ether
- 67. The two forms of D-glucopyranose obtained from the solution of D-glucose are called

 - (a) isomers (b) anomers
- (c) epimers
- (d) enantiomers
- 68. Which pair of compounds is expected to show similar colour in an aqueous medium?
 - (a) FeCl₂ and CuCl₂

(b) VOCl₂ and CuCl₂

(c) VOCl₂ and FeCl₂

- (d) FeCl₂ and MnCl₂
- 69. Which kind(s) of isomerism is/are exhibited by octahedral Co(NH₃)₄Br₂Cl?

 - (a) Geometrical and ionization (b) Geometrical and optical
 - (c) Optical and ionization (d) Geometrical only
- 70. MgSO₄ on reaction with NH₄OH and Na₂HPO₄ forms a white crystalline precipitate. What is its formula?

(a) $Mg(NH_4)PO_4$

(b) $Mg_3(PO_4)_2$

(c) MgCl₂·MgSO₄

- (d) MgSO₄
- 71. For a sparingly soluble salt $A_p B_{qr}$ the relationship of its solubility product (K_s) with its solubility (S) is
 - (a) $K_s = S^{p+q} \cdot p^p \cdot q^q$
- (b) $K_s = S^{p+q} \cdot p^q \cdot q^p$
- (c) $K_s = S^{pq} \cdot p^p \cdot q^q$
- (d) $K_s = S^{pq} \cdot (pq)^{p+q}$
- **72.** Which of the following reactions defines ΔH_f° ?
 - (a) $C(diamond) + O_2(g) \rightarrow CO_2(g)$
 - $(b) \ \frac{1}{2} \, H_2(g) + \frac{1}{2} \, F_2(g) \to HF(g)$
 - (c) $N_2(s) + 3H_2(g) \rightarrow 2NH_3(g)$
 - (d) $CO(g) + \frac{1}{2}O_2(g) \rightarrow CO_2(g)$
- 73. 23 Na is the most stable isotope of Na. Find out the process by which $^{24}_{11}$ Na can undergo radioactive decay.
 - (a) β^- emission

- (b) α emission
- (c) γ-ray emission

- (d) K-electron capture
- 74. Identify the correct order of boiling point of the following compounds:
 - (I) CH₃CH₂CH₂CH₂OH
 - (II) CH₂CH₂CH₂CHO
 - (III) CH₃CH₂CH₂COOH
 - (a) I > II > III

(b) III > I > II

(c) I > III > II

- (d) III > II > I
- **75.** The number of lone pair(s) in $XeOF_4$ is
 - (a) zero

(b) one

(c) two

(d) three

76.

O

N

$$conc. HNO_3$$
 $conc. H_2SO_4$
 $conc. H_2SO_4$

In the above reaction, the structure of the major product (X) is

(a)
$$NO_2$$

(d)
$$O_2N$$
 N N

77. The reagent(s) for the conversion

$$Br \longrightarrow H - \equiv -H$$

is/are

- (a) alcoholic KOH
- (b) alcoholic KOH followed by NaNH₂
- (c) aqueous KOH followed by NaNH₂
- (d) Zn/CH₃OH

78. The number of structural isomers of C_6H_{14} is

(a) three

(b) four

(c) five

(d) six

- 79. The percentage of p-character in the orbital forming P—P bond in P_4 is
 - (a) 25

(b) 33

(c) 50

(d) 75

- **80.** When 20 g naphthoic acid $(C_{11}H_8O_2)$ is dissolved in 50 g benzene $(k_f = 1.72 \text{ K kg mol}^{-1})$, a freezing-point depression of 2 K is observed. The van't Hoff factor (i) is
 - (a) 0.5
- (b) 1
- (c) 2
- (d) 3

- **81.** The value of $\log_{10} k$ for a reaction (A) \rightleftharpoons (B) is (given that $\Delta_{\text{fus}} H_{296 \text{ K}}^{\circ} = -54.07 \text{ kJ mol}^{-1}$, $\Delta_{\text{fus}} S_{296 \text{ K}}^{\circ} = -10 \text{ J K}^{-1} \text{ mol}^{-1}$ and $R = 8.314 \text{ J K}^{-1} \text{ mol}^{-1}$)
 - (a) 5

(b) 10

(c) 95

78. c

79. d

(d) 100

81. b

Answers

1.	d	2. c	3. a	4. d	5. a
6.	a	7. a	8. a	9. a	10. c
11.	b	12. c	13. d	14. d	15. d
16.	b	17. a			
18.	$\text{(i)} \leftrightarrow \text{b}$	$(ii) \leftrightarrow c (iii)$	\leftrightarrow a (iv) \leftrightarrow c		
19.	b	20. c	21. c		
22.	$(i) \leftrightarrow c$	$(ii) \leftrightarrow b (iii)$	\leftrightarrow a (iv) \leftrightarrow d		
23.	a	24. a	25. d	26. b	27. b
28.	d	29. b	30. b	31. b	32. c
33.	b	34. b	35. b	36. b	37. b
38.	b	39. a	40. a	41. b	42. c
43.	c	44. b	45. a	46. d	47. c
48.	d	49. b	50. a	51. a	52. d
53.	d	54. c	55. c	56. d	57. b
58.	d	59. b	60. c	61. a	62. a
63.	a	64. b	65. c	66. a	67. b
68.	b	69. a	70. a	71. a	72. b
73.	a	74. b	75. b	76. b	77. b

80. a

IIT Questions—4

- 1. 2.5 mL of a $\frac{2}{5}$ M weak monoacidic base ($K_b = 1 \times 10^{-12}$ at 25°C) is titrated with $\frac{2}{15}$ M HCl in water at 25°C. The concentration of H⁺ at equivalence point is $(K_{\rm w} = 1 \times 10^{-14} \text{ at } 25^{\circ}\text{C})$
 - (a) $3.7 \times 10^{-13} \text{ mol L}^{-1}$
- (c) $3.2 \times 10^{-2} \text{ mol L}^{-1}$
- (b) $3.2 \times 10^{-7} \text{ mol L}^{-1}$ (d) $2.7 \times 10^{-2} \text{ mol L}^{-1}$
- 2. Native silver forms a water-soluble complex with a dilute aqueous solution of NaCN in the presence of
 - (a) nitrogen

- (b) oxygen
- (c) carbon dioxide
- (d) argon
- 3. Under the same reaction conditions, the initial concentration of 1.386 mol dm⁻³ of a substance becomes half in 40 seconds and 20 seconds through first-order and zero-order kinetics, respectively. The ratio, k_1/k_0 , of the rate constants for the first order and zero order of the reactions is
 - (a) $0.5 \text{ mol}^{-1} \text{ dm}^3$

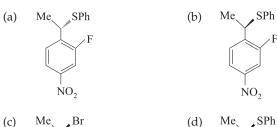
(b) 1.0 mol dm^{-3}

(c) 1.5 mol dm^{-3}

(d) 2.0 mol⁻¹ dm³

4.

The major product of the above reaction is



- 5. The aqueous solution of Na₂S₂O₃ on reaction with Cl₂ gives
 - (a) $Na_2S_4O_6$

(b) NaHSO₄

(c) NaCl

- (d) NaOH
- 6. Hyperconjugation involves overlap of the orbitals
 - (a) σ and σ

(b) σ and ρ

(c) p and p

- (d) π and π
- 7. A gas described by the van der Waals equation
 - (a) behaves similar to an ideal gas in the limit of large molar volumes
 - (b) behaves similar to an ideal gas in the limit of large pressure
 - (c) is characterised by van der Waals coefficients that are dependent on the identity of the gas but are independent of the temperature
 - (d) has the pressure that is lower than the presure exerted by the same gas behaving ideally
- 8. A solution of a colourless salt (H) on boiling with excess NaOH produces a nonflammable gas. The gas evolution ceases after some time. Upon addition of Zn dust to the same solution, the gas evolution restarts. The colourless salt(s) (H) is/are
 - (a) NH₄NO₃

- (b) NH_4NO_2 (c) NH_4Cl (d) $(NH_4)_2SO_4$

9.

Which is/are the correct statement(s) about the above compound?

- (a) The compound is optically active.
- (b) The compound possesses a centre of symmetry.
- (c) The compound possesses a plane of symmetry.
- (d) The compound possesses an axis of symmetry.

Which is/are the correct statement(s) concerning the structures (E), (F) and (G)?

- (a) (E), (F) and (G) are resonance structures.
- (b) (E), (F) and (E), (G) are tautomers.
- (c) (F) and (G) are geometrical isomers.
- (d) (F) and (G) are diastereomers.
- 11. Among the following, the surfactant that will form micelles in aqueous solution at the lowest molar concentration at ambient conditions is

 - (a) $CH_3(CH_2)_{15}N^+(CH_2)_3Br^-$ (b) $CH_3(CH_2)_{11}OSO_3^-Na^+$

 - (c) $CH_3(CH_2)_6COO^-Na^+$ (d) $CH_3(CH_2)_{11}N^+(CH_2)_3Br^-$
- **12.** The solubility product constants ($K_{\rm sp}$) of salts of types MX, MX₂ and M₃X at temperature T are 4.0×10^{-8} , 3.2×10^{-14} and 2.7×10^{-15} , respectively. The solubilities (in mol dm^{-3}) of the salts at temperature T are in the order
 - (a) $MX > MX_2 > M_3X$
- (b) $M_3X > MX_2 > MX$
- (c) $MX_2 > M_2X > MX$
- (d) $MX > M_2X > MX_2$
- 13. Electrolysis of dilute aqueous NaCl solution was carried out by passing 10 mA current. The time required to liberate 0.01 mol of H2 gas at the cathode is (given that Faraday constant = 96500 C mol^{-1})
 - (a) 9.65×10^4 s

(b) $19.3 \times 10^4 \text{ s}$

(c) 28.95×10^4 s

- (d) 38.6×10^4 s
- 14. Cellulose upon acetylation with excess acetic anhydride/H2SO4 (catalytic) gives cellulose triacetate, whose structure is

(a)
$$\begin{array}{c} AcO \\ AcO \\ H \\ OAc \\ H \\ OAc \\ H \end{array}$$
 AcO
$$\begin{array}{c} AcO \\ H \\ OAc \\ H \\ OAc \\ H \end{array}$$
 AcO
$$\begin{array}{c} AcO \\ H \\ OAc \\ H \\ OAc \\ H \end{array}$$

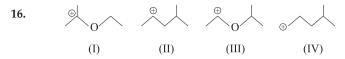
15. O O O Heat
$$\rightarrow$$
 (E) $\frac{I_2}{\text{NaOH}} \rightarrow$ (F) + (G)

(* implies ¹³C labelled carbon)

In the above reaction sequence, the correct structures of (E), (F) and (G) are respectively

(a)
$$Ph * CH_3$$
, $Ph * ONa$ and CHI_3

(c)
$$Ph$$
 CH_3 , Ph ONa and CHI_3



The correct stability order for the above set of species is

- (a) (II) > (IV) > (I) > (III)
- (b) (I) > (II) > (III) > (IV)
- (c) (II) > (I) > (IV) > (III)
- (d) (I) > (III) > (IV)
- 17. Which is a coloured compound among the following?
 - (a) CuCl

(b) $K_3[Cu(CN)_4]$

(c) CuF₂

- (d) $[Cu(CH_3CN)_4]BF_4$
- **18.** The IUPAC name for $[Ni(NH_3)_4]$ $[NiCl_4]$ is
 - (a) tetrachloronickel(II) tetraamminenickelate(II)
 - (b) tetraamminenickel(II) tetrachloronickelate(II)
 - (c) tetraamminenickel(II) tetrachloronickelate(II)
 - (d) tetrachloronickel(II) tetraamminenickelate(0)
- **19.** Both $[Ni(CO)_4]$ and $[Ni(CN)_4]^{2-}$ are diamagnetic. The hybridisations of nickel in these complexes, respectively, are
 - (a) sp^3 and sp^3

(b) sp^3 and dsp^2

(c) dsp^2 and sp^3

- (d) dsp² and dsp²
- **20.** STATEMENT-1: Pb^{4+} compounds are stronger oxidizing agents than Sn^{4+} compounds.
 - STATEMENT-2: The higher oxidation states for the group 14 elements are more stable for the heavier members of the group due to inert-pair effect.
 - (a) Statement-1 is True, Statement-2 is True; Statement-2 is a correct explanation for Statement-1.
 - (b) Statement-1 is True, Statement-2 is True; Statement-2 is not a correct explanation for Statement-1.
 - (c) Statement-1 is True, Statement-2 is False.
 - (d) Statement-1 is False, Statement-2 is True.
- **21.** STATEMENT-1: The plot of atomic number (*y*-axis) versus number of neutrons (*x*-axis) for stable nuclei shows a curvature towards the *x*-axis from the line of 45° slope as the atomic number is increased.
 - STATEMENT-2: Proton-proton electrostatic repulsions begin to overcome attractive forces involving protons and neutrons in heavier nuclides.

- (a) Statement-1 is True, Statement-2 is True; Statement-2 is a correct explanation for Statement-1.
- (b) Statement-1 is True, Statement-2 is True; Statement-2 is **not** a correct explanation for Statement-1.
- (c) Statement-1 is True, Statement-2 is False.
- (d) Statement-1 is False, Statement-2 is True.
- 22. STATEMENT-1: Bromobenzene upon reaction with Br_2/Fe gives 1,4-dibromobenzene as the major product.
 - STATEMENT-2: In bromobenzene, the inductive effect of the bromo group is more dominant than the mesomeric effect directing the incoming electrophile.
 - (a) Statement-1 is True, Statement-2 is True; Statement-2 is a correct explanation for Statement-1.
 - (b) Statement-1 is True, Statement-2 is True; Statement-2 is **not** a correct explanation for Statement-1.
 - (c) Statement-1 is True, Statement-2 is False.
 - (d) Statement-1 is False, Statement-2 is True.
- **23.** STATEMENT-1: For every chemical reaction at equilibrium, standard Gibbs energy of reaction is zero.
 - STATEMENT-2: At constant temperature and pressure, chemical reactions are spontaneous in the direction of decreasing Gibbs energy.
 - (a) Statement-1 is True, Statement-2 is True; Statement-2 is a correct explanation for Statement-1.
 - (b) Statement-1 is True, Statement-2 is True; Statement-2 is not a correct explanation for Statement-1.
 - (c) Statement-1 is True, Statement-2 is False.
 - (d) Statement-1 is False, Statement-2 is True.
- 24. STATEMENT-1: Aniline on reaction with NaNO $_2$ /HCl at 0°C followed by coupling with β -naphthol gives a dark blue precipitate.
 - STATEMENT-2: The colour of the compound formed in the reaction of aniline with NaNO $_2$ /HCl at 0°C followed by coupling with β -naphthol is due to the extended conjugation.
 - (a) Statement-1 is True, Statement-2 is True; Statement-2 is a correct explanation for Statement-1.
 - (b) Statement-1 is True, Statement-2 is True; Statement-2 is **not** a correct explanation for Statement-1.
 - (c) Statement-1 is True, Statement-2 is False.
 - (d) Statement-1 is False, Statement-2 is True.

- **25.** STATEMENT-1: The geometrical isomers of the complex $[M(NH_3)_4Cl_2]$ are optically inactive.
 - STATEMENT-2: Both geometrical isomers of the complex [M(NH₃)₄Cl₂] possess an axis of symmetry.
 - (a) Statement-1 is True, Statement-2 is True; Statement-2 is a correct explanation for Statement-1.
 - (b) Statement-1 is True, Statement-2 is True; Statement-2 is not a correct explanation for Statement-1.
 - (c) Statement-1 is True, Statement-2 is False.
 - (d) Statement-1 is False, Statement-2 is True.
- **26.** STATEMENT-1: There is a natural asymmetry between converting work to heat and converting heat to work.
 - STATEMENT-2: No process is possible in which the sole result is the absorption of heat from a reservoir and its complete conversion into work.
 - (a) Statement-1 is True, Statement-2 is True; Statement-2 is a correct explanation for Statement-1.
 - (b) Statement-1 is True, Statement-2 is True; Statement-2 is not a correct explanation for Statement-1.
 - (c) Statement-1 is True, Statement-2 is False.
 - (d) Statement-1 is False, Statement-2 is True.
- **27.** STATEMENT-1: $[Fe(H_2O)_5NO]SO_4$ is paramagnetic.

STATEMENT-2: The Fe in $[Fe(H_2O)_5NO]SO_4$ has three unpaired electrons.

- (a) Statement-1 is True, Statement-2 is True; Statement-2 is a correct explanation for Statement-1.
- (b) Statement-1 is True, Statement-2 is True; Statement-2 is **not** a correct explanation for Statement-1.
- (c) Statement-1 is True, Statement-2 is False.
- (d) Statement-1 is False, Statement-2 is True.

Answer the questions below (28-30) using the following information.

In the following reaction sequence, products (I), (J) and (L) are formed. (K) represents a reagent.

Hex-3-ynal
$$\xrightarrow{1. \text{ NaBH}_4} (I) \xrightarrow{1. \text{ Mg/ether}} (J) \xrightarrow{(K)} \text{Me} C1 \xrightarrow{Pd/\text{BaSO}_4} (L)$$

28. The structure of the product (I) is

- (a) $Me \longrightarrow Br$ (b) $Me \longrightarrow Br$ (c) $Me \longrightarrow Br$ (d) $Me \longrightarrow Br$
- **29.** The structures of the compounds (J) and (K) are respectively.
 - (a) Me COOH and SOCl₂

 (b) Me OH and SO₂Cl₂

 (c) Me COOH and SOCl₂

 (d) Me COOH and CH₃SO₂Cl

30. The structure of the product (L) is

Answer the questions below (31–33) using the following information.

There are some deposits of nitrates and phosphates in earth's crust. Nitrates are more soluble in water. Nitrates are difficult to reduce under the laboratory conditions but microbes do it easily. Ammonia forms large number of complexes with transition metal ions. Hybridization easily explains the ease of sigma donation capability of NH_3 and PH_3 . Phosphine is a flammable gas and is prepared from white phosphorous.

- **31.** Which is the correct statement among the following?
 - (a) Phosphates have no biologial significance in humans.
 - (b) Between nitrates and phosphates, phosphates are less abundant in earth's crust.
 - (c) Between nitrates and phosphates, nitrates are less abundant in earth's crust.
 - (d) Oxidation of nitrates is possible in soil.

32. Which is the correct statement among the following?

- (a) Between NH₃ and PH₃, NH₃ is a better electron donor because the lone pair of electrons occupies a spherical s orbital and is less directional.
- (b) Between NH₃ and PH₃, PH₃ is a better electron donor because the lone pair of electrons occupies an sp³ orbital and is more directional.

- (c) Between NH₃ and PH₃, NH₃ is a better electron donor because the lone pair of electrons occupies an sp³ orbital and is more directional.
- (d) Between NH₃ and PH₃, PH₃ is a better electron donor because the lone pair of electrons occupies a spherical s orbital and is less directional.
- 33. White phosphorus on reaction with NaOH gives PH_3 as one of the products. This is a
 - (a) dimerization reaction
- (b) disproportionation reaction
- (c) condensation reaction
- (d) precipitation reaction

Answer the questions below (34–36) using the following information.

Properties such as the boiling point, freezing point and vapour pressure of a pure solvent change when solute molecules are added to get a homogeneous solution. These are called colligative properties. Applications of colligative properties are very useful in our day-to-day life. One of its examples is the use of ethylene glycol and water mixture as an antifreezing liquid in the radiator of automobiles.

A solution (M) is prepared by mixing ethanol and water. The mole fraction of ethanol in the mixture is 0.9

Given: Freezing point depression constant of water $(K_f^{\text{water}}) = 1.86 \text{ K kg mol}^{-1}$

Freezing point depression constant of ethanol (K_f^{ethanol}) = 2.0 K kg mol⁻¹

Boiling point elevation constant of water (K_b^{water}) = 0.52 K kg mol⁻¹

Boiling point elevation constant of ethanol (K_h^{ethanol}) = 1.2 K kg mol⁻¹

Standard freezing point of water = 273 K

Standard freezing point of ethanol = 155.7 K

Standard boiling point of water = 373 K

Standard boiling point of ethanol = 351.5 K

Vapour pressure of pure water = 32.8 mm Hg

Vapour pressure of pure ethanol = 40 mm Hg

Molecular weight of water = 18 mol⁻¹

Molecular weight of ethanol = 46 mol⁻¹

In answering the following questions, consider the solution to be ideal dilute solutions and solutes to be nonvolatile and nondissociative.

- **34.** The freezing point of the solution (M) is
 - (a) 268.7 K

(b) 268.5 K

(c) 234.2 K

(d) 150.9 K

- **35.** The vapour pressure of the solution (M) is
 - (a) 39.3 mmHg

(b) 36.0 mmHg

(c) 29.5 mmHg

- (d) 28.8 mmHg
- **36.** Water is added to the solution (M) such that the mole fraction of water in the solution becomes 0.9. The boiling point of this solution is
 - (a) 380.4 K

(b) 376.2 K

(c) 375.5 K

(d) 354.7 K

Answer the questions below (37–39) using the following information.

A tertiary alcohol (H) upon acid-catalysed dehydration gives a product (I). Ozonolysis of (I) leads to the compounds (J) and (K). The compound J upon reaction with KOH gives benzyl alcohol and a compound (L), whereas (K) on reaction with KOH gives only (M).

$$(M):$$

$$\begin{array}{c} & & O \\ & & Ph \\ & & H \end{array}$$

37. The compound (H) is formed by the reaction of

$$(d) \quad \begin{matrix} O \\ \\ Ph \end{matrix} \quad H \qquad \qquad \begin{matrix} Me \\ \\ \\ Ph \end{matrix} \quad MgBi$$

38. The structure of the compound (I) is

(a)
$$\stackrel{\text{Ph}}{\underset{\text{H}}{\bigvee}}$$
 $\stackrel{\text{CH}_3}{\underset{\text{Ph}}{\bigvee}}$ (b) $\stackrel{\text{H}_3\text{C}}{\underset{\text{Ph}}{\bigvee}}$ $\stackrel{\text{Ph}}{\underset{\text{Ph}}{\bigvee}}$

- **39.** The structures of the compounds (J), (K) and (L) are respectively
 - (a) PhCOCH₃, PhCH₂COCH₃ and PhCH₂COO⁻K⁺
 - (b) PhCHO, PhCH₂CHO and PhCOO⁻K⁺
 - (c) PhCOCH₃, PhCH₂CHO and CH₃COO⁻K⁺
 - (d) PhCHO, PhCOCH₃ and PhCOO⁻K⁺

Answer the questions below (40–42) using the following information.

In the hexagonal system of crystals, a frequently encountered arrangement of atoms is described as a hexagonal prism. Here, the top and bottom of the cell are regular hexagons and three atoms are sandwiched between them. A space-filling model of this structure, called the *hexagonal close-packed* (hcp) structure, is constituted of a sphere on a flat surface surrounded in the same plane by six identical spheres as closely as possible. Three spheres are then placed over the first layer so that they touch each other and represent the second layer. Each one of these three spheres touches three spheres of the bottom layer. Finally,the second layer is covered with a third layer that is identical to the bottom layer in relative position. Assume radius of every sphere to be r.

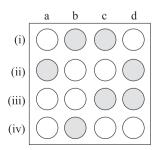


- (a) 4
- (b) 6
- (c) 12
- (d) 17

- 41. The volume of this hcp unit cell is
 - (a) $24\sqrt{2} r^3$
- (b) $16\sqrt{2} r^3$
- (c) $12\sqrt{2} r^3$
- (d) $\frac{64}{3\sqrt{3}}r^3$
- **42.** The empty space in this hcp unit cell is
 - (a) 74%
- (b) 47.6%
- (c) 32%
- (d) 26%

This section contains three questions (43–45). Each question contains statements given in two columns, which have to be matched. Statements in column A are labelled (i), (ii), (iii) and (iv), whereas statements in column B are labelled (a), (b), (c) and (d). The answers to these questions have to be appropriately bubbled, as illustrated in the following example.

If the correct matches are (i) \leftrightarrow b, (i) \leftrightarrow c, (ii) \leftrightarrow a, (ii) \leftrightarrow d, (iii) \leftrightarrow c, (iii) \leftrightarrow d and (iv) \leftrightarrow b then the correctly bubbled matrix will look like the following.



43. Column A

(i) $_{\text{H}_2\text{N}}-\text{NH}_3\text{Cl}$

- Column B
- (a) sodium fusion extract of the compound gives Prussian blue colour with FeSO₄
- (b) gives positive FeCl₃ test
- (iii) $HO \longrightarrow NH_3Cl$
- (c) gives white precipitate with AgNO₃
- (iv) O_2N \longrightarrow $NH-NH_3Br$ $\stackrel{\oplus}{NO_3}$
 - (d) reacts with aldehydes to form the corresponding hydrazone derivative

44. Column A

Column B

- (i) Orbital angular momentum of the electron in a hydrogenlike atomic orbital
- (a) Principal quantum number
- (ii) A hydrogenlike one-electron wave function obeying the Pauli principle
- (b) Azimuthal quantum number
- (iii) Shape, size and orientation of hydrogenlike atomic orbitals
- (c) Magnetic quantum number
- (iv) Probability density of electrons at the nucleus in hydrogenlike atoms
- (d) Spin quantum number

45. Column A

Column B

(i) $PbS \rightarrow PbO$

- (a) Roasting
- (ii) $CaCO_3 \rightarrow CaO$
- (b) Calcination

(iii) $ZnS \rightarrow Zn$

(c) Carbon reduction

(iv) $Cu_2S \rightarrow Cu$

(d) Self-reduction

Answers

1. d	2. b	3. a	4. a	5. b
6. b	7. a, c, d	8. a, b	9. a, d	10. b, c, d
11. c	12. d	13. b	14. a	15. c
16. d	17. c	18. c	19. b	20. c
21. a	22. c	23. d	24. d	25. b
26. b	27. a	28. d	29. c	30. c
31. c	32. c	33. b	34. d	35. b
36. b	37. b	38. a	39. d	40. b
41. a	42. d			

- 43. (ii) (iii) (iv) (iv) (iv) (iii)

IIT Questions—5

1. In the following carbocation, which is the most likely to migrate to the positively charged carbon?

$$H_{3}\overset{H}{\overset{}_{C}} - \overset{H}{\overset{}_{C}} - \overset{H}{\overset{}_{C}} - \overset{5}{\overset{}_{C}} + \overset{5}{\overset{5}} + \overset{5}{\overset{}_{C}} + \overset{5}{\overset{}_{$$

(a) CH₃ at C-4

(b) H at C-4

(c) CH₃ at C-2

- (d) H at C-2
- **2.** For a first-order reaction $A \rightarrow P$, the temperature (*T*)-dependent rate constant (*k*) was found to follow the equation $\log k = -(2000) \frac{1}{T} + 6.0$. The pre-exponential factor A and the activation energy E_a , respectively, are
 - (a) $1.0 \times 10^6 \,\mathrm{s}^{-1}$ and $9.2 \,\mathrm{kJ} \,\mathrm{mol}^{-1}$
 - (b) 6.0 s^{-1} and 16.6 kJ mol^{-1}
 - (c) $1.0 \times 10^6 \,\mathrm{s}^{-1}$ and $16.6 \,\mathrm{kJ} \,\mathrm{mol}^{-1}$
 - (d) $1.0 \times 10^6 \,\mathrm{s}^{-1}$ and $38.3 \,\mathrm{kJ} \,\mathrm{mol}^{-1}$
- 3. Given that the abundances of isotopes 54 Fe, 56 Fe and 57 Fe are 5%, 90% and 5% respectively, the atomic mass of Fe is
 - (a) 55.85

(b) 55.95

(c) 55.75

- (d) 56.05
- Among the following, the polymer in which the intermolecular force of attraction is the weakest is
 - (a) nylon

(b) polyvinyl chloride

(c) cellulose

- (d) natural rubber
- 5. The Henry's law constant for the solubility of N_2 gas in water at 298 K is 1.0×10^5 atm. The mole fraction of N_2 in air is 0.8. The number of moles of N_2 from air dissolved in 10 moles of water at 298 K and 5 atm

pressure is

(a) 4.0×10^{-4} (b) 4.0×10^{-5}

(c) 5.0×10^{-4} (d) 4.0×10^{-6}

6. I₂ is obtained from IO₃ by reduction with

(a) HCO₃

(b) H₂O₂

(c) HSO₃

(d) HNO₂

7. The spin only magnetic moment value (in Bohr magneton units) of $Cr(CO)_6$ is

(a) 0

(b) 2.84

(c) 4.90

(d) 5.92

8. The correct order of stability of the resonance structures

(II)

$$H_2C = \stackrel{+}{N} = \stackrel{-}{N}$$
 $H_2\stackrel{+}{C} - N = \stackrel{-}{N}$ $H_2\stackrel{-}{C} - \stackrel{+}{N} \equiv N$ $H_2\stackrel{-}{C} - N = \stackrel{+}{N}$

(III)

(IV)

is

(I)

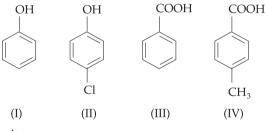
(a) (I) > (II) > (IV) > (III)

(b) (I) > (III) > (IV)

(c) (II) > (I) > (IV)

(d) (III) > (I) > (IV) > (II)

9. The correct order of acidity of



is

(a) (III) > (IV) > (II) > (I)

(b) (IV) > (III) > (I) > (II)

(c) (III) > (II) > (I) > (IV)

(d) (II) > (III) > (IV) > (I)

10. Among the electrolytes Na₂SO₄, CaCl₂, Al₂(SO₄)₃ and NH₄Cl, the most effective coagulating agent for Sb_2S_3 sol is

(a) Na_2SO_4

(b) CaCl₂

(c) $Al_2(SO_4)_3$

(d) NH₄Cl

11. The reaction of P_4 with X leads selectively to P_4O_6 . X is

(a) $dry O_2$

(b) a mixture of O_2 and N_2

(c) moist O₂

(d) O₂ in the presence of aqueous NaOH

12. Which of the following is not applicable to physical adsorptions?

(a) Adsorption on solids is reversible.

(d) Both the enthalpy and entropy of adsorption are negative.

(b) Adsorption increases with increase in temperature.

(c) Adsorption is spontaneous.

13. Among the following, the complex likely to show optical activity is (a) $trans = [Co(NH_2)_4Cl_2]$ (b) $[Cr(H_2O)_6]^{3+}$ (c) cis-[Co(NH₃)₄Cl₂] (d) $[Pt(NH_2)_2Br_2]^+$ 14. The term that corrects for the attractive forces present in a real gas in the van der Waals equation is (b) $\frac{an^2}{V^2}$ (c) $-\frac{an^2}{V^2}$ (d) -nb(a) nb 15. The IUPAC name of is (a) 4-bromo-3-cyanophenol (b) 2-bromo-5-hydroxybenzonitrile (c) 2-cyano-4-hydroxybromobenzene (d) 6-bromo-3-hydroxybenzonitrile 16. The compound(s) formed upon the combustion of sodium metal in an excess of air is(are) (b) Na₂O (a) Na_2O_2 (d) NaOH (c) NaO₂ 17. Which of the following statement(s) regarding defects in solids is(are) correct? (a) A Frenkel defect is usually favoured by a very small difference in the size of the cation and anion. (b) A Frenkel defect is a dislocation defect. (c) The trapping of an electron in the lattice leads to the formation of an F-centre. (d) Schottky defects have no effect on the physical properties of solids.

18. Which of the following compounds exhibit(s) geometrical isomerism?

(b) $[Pt(en)_2] Cl_2$

(d) $[Pt(NH_3)_2Cl_2]$

(a) $[Pt(en)Cl_2]$

(c) [Pt(en), Cl₂]Cl₂

- 19. Which of the following statements is(are) correct about the compound $H_3C(HO)HC$ —CH = CH— $CH(OH)CH_3(X)$?
 - (a) The total number of stereoisomers possible for X is 6.
 - (b) The total number of diastereomers possible for X is 3.
 - (c) If the stereochemistry about the double bond in X is trans, the number of enantiomers possible for X is 4.
 - (d) If the stereochemistry about the double bond in X is cis, the number of enantiomers possible for X is 2.
- 20. In the reaction

$$2X + B_2H_6 \rightarrow [BH_2(X)_2]^+ [BH_4]^-$$

the amine(s) X is(are)

(a) NH₃

(b) CH₃NH₂

(c) (CH₃)₂ NH

- (d) (CH₃)₃ N
- 21. Among the following, which is(are) state function(s)
 - (a) Internal energy
- (b) Irreversible expansion work
- (c) Reversible expansion work (d) Molar enthalpy
- 22. For the reduction of NO_3^- ion in an aqueous solution, E^0 is +0.96 V values of E^0 for some metal ions are given below.

$$V^{2+}(aq) + 2e^{-} \rightarrow V$$
 $E^{0} = -1.19 \text{ V}$

$$E^0 = -1.19 \text{ V}$$

$$Fe^{3+}(aq) + 3e^{-} \rightarrow Fe$$
 $E^{0} = -0.04 \text{ V}$

$$E^0 = -0.04 \text{ V}$$

$$Au^{3+}(aq) + 3e^{-} \rightarrow Au$$
 $E^{0} = +1.40 \text{ V}$

$$E^0 = +1.40 \text{ V}$$

$$Hg^{2+}(aq) + 2e^{-} \rightarrow Hg$$
 $E^{0} = +0.86 \text{ V}$

$$E^0 = +0.86 \text{ V}$$

The pair(s) of metals that is(are) oxidized by NO₃ in an aqueous solution is(are)

(a) V and Hg

(b) Hg and Fe

(c) Fe and Au

- (d) Fe and V
- 23. Which of the following are the amide derivatives of monoamino dicarboxylic acid?
 - (a) Asparagine

(b) Lysine

(c) Proline

- (d) Glutamine
- **24.** The nitrogen oxide(s) that contain(s) N N bond(s) is(are)
 - (a) N_2O

(b) N_2O_3

(c) N_2O_4

(d) N_2O_5

25. For sugars X and Y (shown below), which of the given statements is(are) correct?

- (a) X is a reducing sugar and Y is a nonreducing sugar.
- (b) X is a nonreducing sugar and Y is a reducing sugar.
- (c) The glucosidic linkages in X and Y are α and β respectively.
- (d) The glucosidic linkages in X and Y are β and α respectively.
- **26.** C_V and C_p denote the molar specific heat capacities of a gas at constant volume and constant pressure respectively. Which of the following statements is(are) correct?
 - (a) $C_p C_V$ is larger for a diatomic ideal gas than for a monoatomic ideal gas.
 - (b) $C_p + C_V$ is larger for a diatomic ideal gas than for a monoatomic ideal gas.
 - (c) C_p/C_V is larger for a diatomic ideal gas than for a monoatomic ideal gas.
 - (d) $C_p \cdot C_V$ larger for a diatomic ideal gas than for a monoatomic ideal gas.
- **27.** Match List I (reactions) with List II (reagents) and select the correct answer using the codes given below the lists.

List I (reactions)	List II (reagents)
(A) Wolff–Kishner reduction	1. NaCN
(B) Witting reaction	2. $Ph_3P = CHR$

(C) Benzoin condensation(D) Cannizzaro reaction	 3. Concentrated NaOH 4. N₂H₄, KOH 5. NaOBr 	
Codes:		_
(a) A B C D	(b) A B C D	
4 3 2 1	1 2 5 3	
(c) A B C D	(d) ABCD	
4 2 1 3	1 3 5 2	

Questions 28 through 31 contain two columns, which have to be matched. The items in **Column I** are labelled A, B, C and D, while those in **Column II** are labelled p, q, r, s and t. Any given item in **Column I** can match **ONE OR MORE** in **Column II**. The appropriate bubbles corresponding to the answers to these questions have to be darkened as illustrated in the following example.

If the correct matches are A - p and t; B - q and r; C - p and q; and D - s and t; the bubbles should be darkened as follows.

	р	q	r	s	t
Α	p	q	(r)	s	(t)
В	(a)	q	r	s	(t)
С	p	q	r	s	t
D	p	q	r	s	t

28. Match each of the reactions given in **Column I** with the corresponding product(s) given in **Column II**.

Column I	Column II
(A) $Cu + dil.HNO_3$	(p) NO
(B) $Cu + conc. HNO_3$	(q) NO ₂
(C) $Zn + dil. HNO_3$	(r) N ₂ O
(D) $Zn + conc. HNO_3$	(s) $Cu(NO_3)_2$
	(t) $Zn(NO_3)_2$

29. Match each of the compounds given in **Column I** with the reaction(s) that they can undergo, given in **Column II**.

Column I	Column II
(A) Br	(p) Nucleophilic substitution

30. Match each of the diatomic molecules in **Column I** with its property/properties in **Column II**.

Column I	Column II
(A) B_2	(p) Paramagnetic
(B) N ₂	(q) Undergoes oxidation
(C) $_{{\rm O}_{2}^{-}}$	(r) Undergoes reduction
(D) O ₂	(s) Bond order≥2
	(t) Mixing of s and p orbitals

31. Match each of the compounds in **Column I** with its characteristics reaction(s) in **Column II**.

Column I	Column II
(A) CH ₃ CH ₂ CH ₂ CN	(p) Reduction with $Pd - C/H_2$
(B) CH ₃ CH ₂ OCOCH ₃	(q) Reduction with SnCl ₂ /HCl
(C) CH ₃ —CH=CH—CH ₂ OH	(r) Development of foul smell on treatment with chloroform and alcoholic KOH
(D) CH ₃ CH ₂ CH ₂ CH ₂ NH ₂	(s) Reduction with diisobutyl- aluminium hydride (DIBAL-H)
	(t) Alkaline hydrolysis

Answer questions (32–34) using the following information.

p-Amino-*N*, *N*-dimethylaniline is added to a strongly acidic solution of X. The resulting solution is treated with a few drops of an aqueous solution of Y to yield a blue coloration due to the formation of methylene blue. Treatment of the aqueous solution of Y with the reagent potassium hexacyanoferrate(II) leads to the formation of an intense blue precipitate. The precipitate dissolves on excess addition of the reagent. Similarly, treatment of the solution of Y with the solution of potassium hexacyanoferrate(III) leads to a brown coloration due to the formation of Z.

- **32.** The compound X is
 - (a) NaNO₃

(b) NaCl

(c) Na₂SO₄

(d) Na₂S

- **33.** The compound Y is
 - (a) MgCl₂

(b) FeCl₂

(c) FeCl₃

(d) ZnCl₂

- **34.** The compound *Z* is
 - (a) $Mg_2[Fe(CN)_6]$

(b) $Fe[Fe(CN)_6]$

(c) $Fe_4[Fe(CN)_6]_3$

(d) $K_2Zn_3[Fe(CN)_6]_2$

Answer questions 35–37 using the following information.

A carbonyl compound P, which responds positively to the iodoform test, reacts with MeMgBr followed by dehydration to give an olefin Q. Ozonolysis of Q leads to the formation of a dicarbonyl compound R, which undergoes an intramolecular aldol reaction to give predominantly S.

$$P \xrightarrow[3. \text{ MeMgBr}]{1. \text{ MeMgBr}} Q \xrightarrow[2. \text{ H}^+, \text{ H}_2\text{O}]{2. \text{ Zn, H}_2\text{O}}} R \xrightarrow[2. \text{ A}]{1. \text{ OH}} S$$

35. The structure of the carbonyl compound. P is

36. The structures of the products Q and R, respectively, are

37. The structure of the product S is

The answer to each of the questions (38–45) is a single-digit integer, ranging from 0 to 9. The appropriate bubbles below the respective question numbers in the ORS have to be

darkened. For example, if the correct answers to question numbers X, Y, Z and W (say) are 6, 0, 9 and 2, respectively, the bubbles should be darkened as follows.

Χ

0 0 0 0

(1)

(2)

(3)

(4)

(5) (5) (5)

6 6 6 6

77777

(8) (8)

9 9

(3)

Z W

(3) (3)

(4)

(8)

- **38.** The dissociation constant of a substituted benzoic acid at 25°C is 1.0×10^{-4} . The pH of a 0.01 M solution of its sodium salt is
- **39.** At 400 K, the root mean square (rms) speed of a gas X (molecular weight = 40) is equal to the most probable speed of gas Y at 60 K. The molecular weight of the gas Y is
- **40.** The total number of α and β particles emitted in the nuclear reaction $^{238}_{92}U \rightarrow ^{214}_{82}Pb$ is
- **41.** The oxidation number of Mn in the product of alkaline oxidative fusion of MnO₂ is
- **42.** The coordination number of Al in the crystalline state of AlCl₃ is
- **43.** The number of water molecule(s) directly bonded to the metal centre in CuSO₄ · 5H₂O is
- 44. In a constant-volume calorimeter, 3.5 g of a gas with molecular weight 28 was burnt in excess oxygen at 298.0 K. The temperature of the calorimeter was found to increase from 298.0 K to 298.45 K due to the combustion process. Given that the heat capacity of the calorimeter is 2.5 kJ K⁻¹, the numerical value for the enthalpy of combustion of the gas in kJ mol⁻¹ is
- **45.** The total number of cyclic structures as well as stereoisomers possible for a compound with the molecular formula C_5H_{10} is

Answers

1. d	2. d	3. b	4. d	5. a
6. c	7. a	8. b	9. a	10. b
11. b	12. b	13. c	14. b	15. b
16. a, b	17. b, c	18. c, d	19. a, d	20. a, b, c
21. a, d	22. a, b, d	23. a, d	24. a, b, c	25. b, c
26. b. d	27. c			

28.

	р	q	r	S	t
Α	p	q	r	s	(t)
В	p	q	r	s	(t)
С	p	q	r	s	t
D	p	q	r	s	t

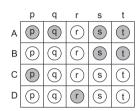
29.

	р	q	r	S	t
Α	p	q	(r)	s	t
В	p	q		S	t
С	(p)	q	r	s	t
D	p	q	(-)	(s)	t

30.

	р	q	r	s	t
Α	p	q	r	s	(t)
В	p	q	r	s	(t)
С	p	q	r	s	t
D	p	q	r	S	(t)

31.



32. d

33. c

34. b

35. b

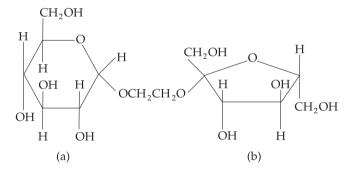
36. a

37. b

38	39	40	41	42	43	44	45
0	0	0	0	0	0	0	0
1	1	1	1	1	1	1	1
2	2	2	2	2	2	2	2
3	3	3	3	3	3	3	3
4	4	4	4	4	4	4	4
5	(5)	(5)	(5)	(5)	(5)	(5)	(5)
6	6	6	6	6	6	6	6
7	7	7	7	7	7	7	7
8	8	8	8	8	8	8	8
9	9	9	9	9	9	9	9

IIT Questions—6

1. The correct statement about the following disaccharide is



- (a) Ring (a) is pyranose with α glycosidic link
- (b) Ring (a) is furanose with α glycosidic link
- (c) Ring (b) is furanose with α glycosidic link
- (d) Ring (b) is pyranose with β glycosidic link
- The synthesis of 3-octyne is achieved by adding a bromoalkane into a mixture of sodium amide and an alkyne. The bromoalkane and alkyne respectively are
 - (a) $BrCH_2CH_2CH_2CH_3$ and $CH_3CH_2C \equiv CH$
 - (b) $BrCH_2CH_2CH_3$ and $CH_3CH_2CH_2C \equiv CH$
 - (c) $BrCH_2CH_2CH_2CH_3$ and $CH_3C \equiv CH$
 - (d) $BrCH_2CH_2CH_2CH_3$ and $CH_3CH_2C \equiv CH$
- 3. The ionization isomer of $[Cr(H_2O)_4Cl(NO_2)]Cl$ is
 - (a) $[Cr(H_2O)_4(O_2N)]Cl_2$
- (b) $[Cr(H_2O)_4Cl_2](NO_2)$
- (c) $[Cr(H_2O)_4Cl(ONO)]Cl$
- (d) $[Cr(H_2O)_4Cl_2(NO_2)] \cdot H_2O$

4. The correct structure of ethylenediaminetetraacetic acid (EDTA) is

(a)
$$HOOC-CH_2$$
 $N-CH=CH-N$ CH_2-COOH CH_2-COOH

(b)
$$N-CH_2-CH_2-N$$
 COOH

(c)
$$HOOC-CH_2$$
 $N-CH_2-CH_2-N$ CH_2-COOH CH_2-COOH

(d)
$$\begin{array}{c|c} & & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & &$$

- 5. The bond energy (in kcal mol⁻¹) of a C—C single bond is approximately
 - (a) 1

(b) 10

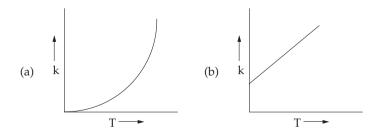
(c) 100

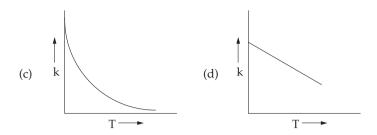
- (d) 1000
- 6. The species which by definition has ${\bf ZERO}$ standard molar enthalpy of formation at 298 K is
 - (a) $Br_2(g)$

(b) Cl₂ (g)

(c) $H_2O(g)$

- (d) CH₄ (g)
- 7. Plots showing the variation of the rate constant (*k*) with temperature (*T*) are given below. The plot that follows Arrhenius equation is





- 8. In the reaction $OCH_3 \xrightarrow{HBr}$ the products are
 - (a) Br \longrightarrow OCH₃ and H₂
 - (b) $\langle \rangle$ —Br and CH_3Br
 - (c) $\langle \rangle$ —Br and CH₃OH
 - (d) OH and CH₃Br
- 9. The species having pyramidal shape is
 - (a) SO_3
- (b) BrF₃
- (c) SiO_3^{2-}
- (d) OSF₂

10. The compounds P, Q and S

were separately subjected to nitration using $\,HNO_3/H_2SO_4$ mixture. The major product formed in each case respectively, is

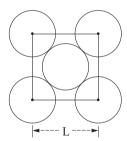
(a)
$$HO$$
 H_3C NO_2 NO_2 NO_2 NO_2

(b)
$$HO$$
 NO_2 H_3C NO_2 NO_2

(c)
$$HO$$
 NO_2 H_3C NO_2 NO_2

(d)
$$HO$$
 H_3C NO_2 NO_2 NO_2

- 11. Assuming that Hund's rule is violated, the bond order and magnetic nature of the diatomic molecule B_2 is
 - (a) 1 and diamagnetic
- (b) 0 and diamagnetic
- c) 1 and paramagnetic
- (d) 0 and paramagnetic
- **12.** The packing efficiency of the two-dimensional square unit cell shown below is



(a) 39.27%

(b) 68.02%

(c) 74.05%

(d) 78.54%

13. In the reaction
$$H_3C$$
 \longrightarrow C \longrightarrow

structure of the product T is

(c)
$$H_3C$$
 NH C

$$H_3C \longrightarrow C \longrightarrow O$$

$$NH - C$$

- 14. The complex showing a spin-only magnetic moment of 2.82 BM, is
 - (a) Ni(CO)₄

(b) [NiCl₄]²⁻

(c) Ni(PPh₃)₄

- (d) $Ni(CN)_4]^{2-}$
- **15.** Among the following, the intensive property is (properties are)
 - (a) molar conductivity
- (b) electromotive force

(c) resistance

- (d) heat capacity
- 16. The reagent(s) used for softening the temporary hardness of water is(are)
 - (a) $Ca_3(PO_4)_2$

(b) Ca(OH)₂

(c) Na₂CO₃

- (d) NaOCl
- **17.** Aqueous solutions of HNO₃, KOH, CH₃COOH, and CH₃COONa of identical concentrations are provided. The pair(s) of solutions which form a buffer upon mixing is(are)
 - (a) HNO₃ and CH₃COOH
 - (b) KOH and CH₃COONa
 - (c) HNO₃ and CH₃COONa
 - (d) CH₃COOH and CH₃COONa

- 18. In the Newman projection for 2, 2-dimethylbutane
- H₃C CH₃

- X and Y can respectively be
 - (a) H and H
- (b) H and C_2H_5
- (c) C₂H₅ and H (d) CH₃ and CH₃
- 19. In the reaction $NaOH(aq)/Br_2$ the intermediate(s) is(are)

 NaOH(aq)/ Br_2 O

 Br

 (b)

 Br

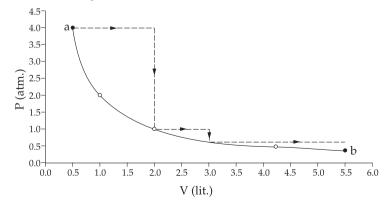
 Br

 (c)

 O

 Br

 Br
- **20.** One mole of an ideal gas is taken from **a** to **b** along two paths denoted by the solid and the dashed lines as shown in the graph below. If the work done along the solid line path is $w_{\rm s}$ and that along the dotted line path is $w_{\rm d}$, then the integer closest to the ratio $w_{\rm d}/w_{\rm s}$ is



21. Among the following, the number of elements showing only one nonzero oxidation state is

- **22.** Silver (atomic weight = 108 g mol^{-1}) has a density of 10.5 g cm^{-3} . The number of silver atoms on a surface of area 10^{-12}m^2 can be expressed in scientific notation as $y \times 10^x$. The value of x is
- 23. The total number of diprotic acids among the following is

H₃PO₄ H₂SO₄ H₃PO₃ H₂CO₃ H₂S₂O₇ H₃BO₃ H₃PO₂ H₂CrO₄ H₂SO₃

24. Total number of geometrical isomers for the complex [RhCl(CO)(PPh₃) (NH₃)] is

Paragraph Type Questions

This Section contains 2 paragraphs. Based upon the first paragraph 3 multiple choice questions and based upon the second paragraph 2 multiple choice questions have to be answered. Each of these questions has four choices (a), (b), (c) and (d) out of which ONLY ONE is correct.

Paragraph for Questions 25 to 27

Copper is the most noble of the first row transition metals and occurs in small deposits in several countries. Ores of copper include chalcanthite (CuSO $_4$ ·5H $_2$ O), atacamite [Cu $_2$ Cl(OH) $_3$], cuprite (Cu $_2$ O), copper glance (Cu $_2$ S) and malachite [CU $_2$ (OH) $_2$ CO $_3$]. However, 80% of the world copper production comes from the ore chalcopyrite (CuFeS $_2$). The extraction of copper from chalcopyrite involves partial roasting, removal of iron and self-reduction.

25.	Partial	roasting	of	chalcopyrite	produces
-----	---------	----------	----	--------------	----------

(a) Cu₂S and FeO

- (b) Cu₂O and FeO
- (c) CuS and Fe₂O₃
- (d) Cu₂O and Fe₂O₃
- **26.** Iron is removed from chalcopyrite is
 - (a) FeO

(b) FeS

(c) Fe_2O_3

- (d) FeSiO₃
- 27. In self-reduction, the reducing species is
 - (a) S

(b) O²⁻

(c) S^{2-}

(d) SO_2

Paragraph for Questions 28 to 29

The concentration of potassium ions inside a biologial cell is at least twenty times higher than the outside. The resulting potential difference across the cell is important in several processes such as transmission of nerve impulses and maintaining the ion balance. A simple model for such a concentration cell involving a metal M is

$$M(s) | M^{+}(aq; 0.05 \text{ molar}) | M^{+}(aq; 1 \text{ molar}) | M(s)$$

For the above electrolytic cell the magnitude of the cell potential $\mid E_{\rm cell} \mid = 70 \text{ mV}.$

- 28. For the above cell
 - (a) $E_{cell} < 0$; $\Delta G > 0$
- (b) $E_{cell} > 0$; $\Delta G < 0$
- (c) $E_{cell} < 0; \Delta G^{\circ} > 0$
- (d) $E_{cell} > 0$; $\Delta G^{\circ} < 0$
- **29.** If the 0.05 molar solution of M⁺ is replaced by a 0.0025 molar M⁺ solution, then the magnitude of the cell potential would be
 - (a) 35 mV

(b) 70 mV

(c) 140 mV

(d) 700 mV

Paragraph Type Questions

This Section contains **2 paragraphs**. Based upon each of the paragraph **3 multiple choice questions** have to be answered. Each of these questions has four choices (a), (b), (c) and (d) out of which **ONLY ONE** is correct.

Paragraph for Questions 30 to 32

Two aliphatic aldehydes P and Q react in the presence of aqueous K_2CO_3 to give compound R, which upon treatment with HCN provides compound S. On acidification and heating, S gives the product shown below:

30. The compounds P and Q respectively are

31. The compound R is

32. The compound S is

Paragraph for Questions 33 to 35

The hydrogen-like species Li^{2+} is in a spherically symmetric state S_1 with one radial node. Upon absorbing light the ion undergoes transition to a state S_2 . The state S_2 has one radial node and its energy is equal to the ground state energy of the hydrogen atom.

33. The State S_1 is

(a) 1s

(b) 2s

(c) 2p

(d) 3s

- **34.** Energy of the state S_1 in units of the hydrogen atom ground state energy is
 - (a) 0.75

(b) 1.50

(c) 2.25

- (d) 4.50
- 35. The orbital angular momentum quantum number of the state S_2 is
 - (a) 0

(b) 1

(c) 2

(d) 3

Integer Type Questions

This Section contains TEN questions. The answer to each question is a **single digit integer** ranging from 0 to 9. The correct digit below the question number in the ORS is to be bubbled.

36. Amongst the following, the total number of compounds whose aqueous solution turns red litmus paper blue is

KCN K₂SO₄ (NH₄)₂C₂O₄ NaCl Zn(NO₃)₂ FeCl₂ K₂CO₂ NH₄NO₂ LiCN

- 37. Based on VSEPR theory, the number of 90 degree F—Br—F angles in ${\rm BrF}_5\,{\rm is}$
- **38.** The value of *n* in the molecular formula $Be_nAl_2Si_6O_{18}$ is
- **39.** A student performs a titration with different burettes and finds titre values of 25.2 mL, 25.25 mL, and 25.0 mL. The number of significant figures in the average titre value is
- **40.** The concentration of R in the reaction $R \to P$ was measured as a function of time and the following data is obtained

[R] (molar)	1.0	0.75	0.40	0.10
t(min.)	0.0	0.05	0.12	0.18

The order of the reaction is

- **41.** The number of neutrons emitted when $^{235}_{92}$ U undergoes controlled nuclear fission to $^{142}_{54}$ Xe and $^{90}_{38}$ Sr is
- 42. The total number of basic groups in the following form of lysine is

$$\overset{\oplus}{\text{H}_3\text{N}}$$
 $-\text{CH}_2$
 $-\text{CH}_2$

- 43. The total number of cyclic isomers possible for a hydrocarbon with the molecular formula C_4H_6 is
- **44.** In the scheme given below, the total number of intramolecular aldol condensation products formed from 'Y' is

$$\frac{1. \text{ O}_3}{2. \text{ Zn, H}_2\text{O}} \quad \text{Y} \quad \frac{1. \text{ NaOH (aq)}}{2. \text{ heat}}$$

45. Amongst the following, the total number of compounds soluble in aqueous NaOH is

Matrix Type Questions

This section contains **2 questions**. Each question has four statements (a, b, c and d) given in **Column I** and five statements (p, q, r, s and t) in **Column II**. Any given statement in **Column I** can have correct matching with one or more statements given in **Column II**. For example, if for a given question, statement b matches with the statements given in q and r, then for that particular question, against statement b, darken the bubbles corresponding to q and r in the ORS.

46. Match the reactions in **Column I** with appropriate options in **Column II**.

Column I Column II

$$\begin{array}{c} & & & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & \\ & & & \\ & & & \\ & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\$$

$$\begin{array}{c} O \\ \parallel \\ C \\ C \\ C \\ -CH_3 \\ CH_3 \end{array}$$

(c)
$$CH_3 \xrightarrow{\text{1. LiAIH}_4}$$

(r) Substitution reaction

(d)
$$HS \longrightarrow Cl \xrightarrow{Base} S$$

_ _

(s) Coupling reaction

- (t) Carbocation intermediate
- **47.** All the compounds listed in **Column I** react with water. Match the result of the respective reactions with the appropriate options listed in **Column II**.

Column I

- (a) (CH₃)₂SiCl₂
- (b) XeF₄
- (c) Cl_2
- (d) VCl₅

Column II

- (p) Hydrogen halide formation
- (q) Redox reaction
- (r) Reacts with glass
- (s) polymerization
- (t) O₂ formation

Answers

1. a **2.** d **3.** b **4.** c **5.** c 6. b 7. a 8. d **9.** d **10.** c 14. b **11.** a **12.** d **13.** c 15. a, b 16. b, c **17.** d **18.** b, d **19.** a, b, c

20	21	22	23	24
0	0	0	0	0
1	1	1	1	1
2	2	2	2	2
3	3	3	3	3
4	4	4	4	4
(5)	(5)	(5)	(5)	(5)
6	6	6	6	6
7	7	7	7	7
8	8	8	8	8
9	9	9	9	9

 25. a
 26. d
 27. c
 28. b
 29. c

 30. b
 31. a
 32. d
 33. b
 34. c

 35. b

36	37	38	39	40	41	42	43	44	45
0	0	0	0	0	0	0	0	0	0
1	1	1	1	1	1	1	1	1	1
2	2	2	2	2	2	2	2	2	2
3	3	3	3	3	3	3	3	3	3
4	4	4	4	4	4	4	4	4	4
(5)	(5)	(5)	(5)	(5)	(5)	(5)	(5)	(5)	(5)
6	6	6	6	6	6	6	6	6	6
7	7	7	7	7	7	7	7	7	7
8	8	8	8	8	8	8	8	8	8
9	9	9	9	9	9	9	9	9	9

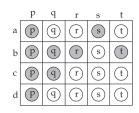
46.

	р	q	r	s	t
a	p	\bigcirc	r	\bigcirc s	t
b	p	q	r	s	t
c	p	\bigcirc	r	\bigcirc s	t
d	p	q	r	s	t

$$a \rightarrow r, s$$

 $b \rightarrow t$
 $c \rightarrow p, q$
 $d \rightarrow r$

47.



$$a \rightarrow p, s$$

 $b \rightarrow p, q, r, t$
 $c \rightarrow p, q$
 $d \rightarrow p$

IIT Questions—7

1. The freezing point (in °C) of a solution containing 0.1 g of $K_3[Fe(CN)_6]$ (mol. wt. 329) in 100 g of water ($K_f = 1.86 \text{ K kg mol}^{-1}$) is

(a)
$$-2.3 \times 10^{-2}$$

(b)
$$-5.7 \times 10^{-2}$$

(c)
$$-5.7 \times 10^{-3}$$

(d)
$$-1.2 \times 10^{-2}$$

2. Amongst the compounds given, the one that would form a brilliant colored dye on treatment with $NaNO_2$ in dil. HCl followed by addition to an alkaline solution of β -naphthol is

(a)
$$N(CH_3)_2$$

3. The major product of the following reaction is

(a) a hemiacetal

(b) an acetal

(c) an ether

- (d) an ester
- 4. The following carbohydrate is

(a) a ketohexose

(b) an aldohexose

(c) an α-furanose

- (d) an α-pyranose
- Oxidation states of the metal in the minerals haematite and magnetite, respectively, are
 - (a) II, III in haematite and III in magnetite
 - (b) II, III in haematite and II in magnetite
 - (c) II in haematite and II, III in magnetite
 - (d) III in haematite and II, III in magnetite
- **6.** Among the following complexes (K–P)

$$K_3[Fe(CN)_6]$$
 (K), $[Co(NH_3)_6]Cl_3$ (L), $Na_3[Co(oxalate)_3]$ (M),

 $[Ni(H_2O)_6]Cl_2(N), K_2[Pt(CN)_4](O) \text{ and } [Zn(H_2O)_6](NO_3)_2(P)$

the diamagnetic compounds are

(a) K, L, M, N

(b) K, M, O, P

(c) L, M, O, P

- (d) L, M, N, O
- 7. Passing H_2S gas into a mixture of Mn^{2+} , Ni^{2+} , Cu^{2+} and Hg^{2+} ions in an acidified aqueous solution precipitates
 - (a) CuS and HgS

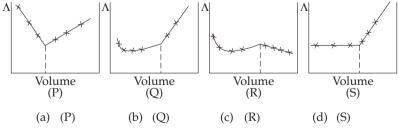
(b) MnS and CuS

(c) MnS and NiS

- (d) NiS and HgS
- 8. Consider the following cell reaction:

 $2Fe(s)+O_2(g)+4H^+(aq)\rightarrow 2Fe^{2+}(aq)+2H_2O(l)$ $E^\circ=1.67~V$ Given $[Fe^{2+}]=10^{-3}~M,$ $p_{O_2}=0.1~atm$ and pH=3 , the cell potential at $25^\circ C$ is

- (a) 1.47 V
- (b) 1.77 V
- (c) 1.87 V
- (d) 1.57 V
- 9. Dissolving 120 g of urea (mol. wt. 60) in 1000 g of water gave a solution of density 1.15 g/mL. The molarity of the solution is
 - (a) 1.78 M
- (b) 2.00 M
- (c) 2.05 M
- (d) 2.22 M
- 10. AgNO $_3$ (aq) was added to an aqueous KCl solution gradually and the conductivity of the solution was measured. The plot of conductance Λ versus the volume of AgNO $_3$ is



- 11. Among the following compounds, the most acidic is
 - (a) *p*-nitrophenol

- (b) p-hydroxybenzoic acid
- (c) *o*-hydroxybenzoic acid
- (d) p-toluic acid

12. The major product of the following reaction is

(a)
$$C$$
 N — CH_2 — Br

(b)
$$CN - CH_2CI$$

(c)
$$CN$$
 $O-CH_2-Br$

13. Extra pure N₂ can be obtained by heating

- (a) NH₃ with CuO
- (b) NH₄NO₃

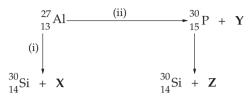
(c) $(NH_4)_2Cr_2O_7$

(d) $Ba(N_3)_2$

14. Geometrical shapes of the complexes formed by the reaction of Ni^{2+} with Cl^- , CN^- and H_2O , respectively, are

- (a) octahedral, tetrahedral and square planar
- (b) tetrahedral, square planar and octahedral
- (c) square planar, tetrahedral and octahedral
- (d) octahedral, square planar and octahedral

15. Bombardment of aluminium by α -particle leads to its artificial disintegration in two ways, (i) and (ii) as shown. Products X, Y and Z respectively are



- (a) proton, neutron, positron
- (b) neutron, positron, proton
- (c) proton, positron, neutron
- (d) positron, proton, neutron
- **16.** Amongst the given options, the compound(s) in which all the atoms are in one plane in all the possible conformations (if any), is (are)

(a)
$$H_2C$$
 $C-C$ CH_2 (b) $H-C \equiv C-C$ CH_2 CH_2 CH_2 CH_2

- 17. According to kinetic theory of gases
 - (a) collisions are always elastic
 - (b) heavier molecules transfer more momentum to the wall of the container
 - (c) only a small number of molecules have very high velocity
 - (d) between collisions, the molecules move in straight lines with constant velocities
- **18.** The correct statement(s) pertaining to the adsorption of a gas on a solid surface is (are)
 - (a) adsorption is always exothermic
 - (b) physisorption may transform into chemisorption at high temperature
 - (c) physisorption increases with increasing temperature but chemisorption decreases with increasing temperature
 - (d) chemisorption is more exothermic than physisorption, however it is very slow due to higher energy of activation
- 19. Extraction of metal from the ore cassiterite involves
 - (a) carbon reduction of an oxide ore
 - (b) self-reduction of a sulphide ore
 - (c) removal of copper impurity
 - (d) removal of iron impurity

- 20. Reduction of the metal centre in aqueous permanganate ion involves
 - (a) 3 electrons in neutral medium
 - (b) 5 electrons in neutral medium
 - (c) 3 electrons in alkaline medium
 - (d) 5 electrons in acidic medium
- 21. The correct functional group X and the reagent/reaction conditions Y in the following scheme are

- (a) $X = COOCH_2$, $Y = H_2/Ni/heat$
- (b) $X = CONH_2$, $Y = H_2/Ni/heat$
- (c) $X = CONH_2$, $Y = Br_2/NaOH$
- (d) X = CN, $Y = H_2/Ni/heat$
- 22. For the first order reaction $2N_2O_5(g) \rightarrow 4NO_2(g) + O_2(g)$
 - (a) the concentration of the reactant decreases exponentially with time
 - (b) the half-life of the reaction decreases with increasing temperature
 - (c) the half-life of the reaction depends on the initial concentration of reactant
 - (d) the reaction proceeds to 99.6% completion in eight half-life duration
- 23. The equilibrium $2Cu^{I} \rightleftharpoons Cu^{0} + Cu^{II}$ in aqueous medium at 25°C shifts towards the left in the presence of
 - (a) NO_3^- (b) CI^-
- (c) SCN^- (d) CN^-

Paragraph-Type Questions

This section contains 2 paragraphs. Based upon the first paragraph 2 multiple choice questions and based upon the second paragraph 3 multiple choice questions have to be answered. Each of these questions has four choices (a), (b), (c) and (d) out of which ONLY ONE is correct.

Paragraph for Questions 24 and 25

An acyclic hydrocarbon P, having molecular formula C₆H₁₀, gave acetone as the only organic product through the following sequence of reactions, in which Q is an intermediate organic compound.

$$P \\ (C_6H_{10}) \xrightarrow{\text{(i) dil. } H_2SO_4/HgSO_4 \\ \text{(iii) } NaBH_4/ethanol \\ \text{(iii) dil. acid}} Q \xrightarrow{\text{(i) conc. } H_2SO_4 \\ \text{(catalytic amount)}} \left\{ \begin{matrix} O \\ \text{(catalytic amount)} \\ \text{(iii) } O_3 \\ \text{(iii) } Zn/H_2O \end{matrix} \right\}$$

24. The structure of compound P is

(a)
$$CH_3CH_2CH_2CH_2-C \equiv C-H$$

(b)
$$H_3CH_2C-C \equiv C-CH_2CH_3$$

(c)
$$H_3C$$

 H_3C $+ C = C - CH_3$

(d)
$$H_3C$$
 $C-C \equiv C-H$ H_3C

25. The structure of compound Q is

Paragraph for Questions 26 to 28

When a metal rod M is dipped into an aqueous colourless concentrated solution of compound N, the solution turns light blue. Addition of aqueous NaCl to the blue solution gives a white precipitate O. Addition of aqueous NH_3 dissolves O and gives an intense blue solution.

- **26.** The metal rod M is
 - (a) Fe (b) Cu (c) Ni (d) Co
- **27.** The compound N is
- (a) $AgNO_3$ (b) $Zn(NO_3)_2$ (c) $Al(NO_3)_3$ (d) $Pb(NO_3)_2$
- 28. The final solution contains
 - $\text{(a)} \quad [Pb(NH_3)_4]^{2+} \text{ and } [CoCl_4]^{2-} \qquad \text{(b)} \quad [Al(NH_3)_4]^{3+} \text{ and } [Cu(NH_3)_4]^{2+}$
 - (c) $[Ag(NH_3)_2]^+$ and $[Cu(NH_3)_4]^{2+}$ (d) $[Ag(NH_3)_2]^+$ and $[Ni(NH_3)_6]^{2+}$

Integer-Answer-Type Questions

The answer to each question is a **single digit integer** ranging from 0 to 9. The bubble corresponding to the correct answer is to be darkened in the ORS.

29. The work function (φ) of some metals is listed below. The number of metals which will show photoelectric effect when light of 300 nm wavelength falls on the metal is

Metal	Li	Na	K	Mg	Cu	Ag	Fe	Pt	W
φ (eV)	2.4	2.3	2.2	3.7	4.8	4.3	4.7	6.3	4.75

- **30.** To an evacuated vessel with movable piston under external pressure of 1 atm., 0.1 mol of He and 1.0 mol of an unknown compound (vapour pressure 0.68 atm. at 0°C) are introduced. Considering the ideal gas behaviour, the total volume (in litre) of the gases at 0°C is close to
- **31.** Reaction of Br₂ with Na₂CO₃ in aqueous solution gives sodium bromide and sodium bromate with evolution of CO₂ gas. The number of sodium bromide molecules involved in the balanced chemical equation is
- 32. The difference in the oxidation numbers of the two types of sulphur atoms in $Na_2S_4O_6$ is
- **33.** A decapeptide (mol. wt. 796) on complete hydrolysis gives glycine (mol. wt. 75), alanine and phenylalanine. Glycine contributes 47.0% to the total weight of the hydrolysed products. The number of glycine units present in the decapeptide is
- **34.** The total number of alkenes possible by dehydrobromination of 3-bromo-3-cyclopentylhexane using alcoholic KOH is
- **35.** The maximum number of electrons that can have principal quantum number, n = 3, and spin quantum number, $m_S = -1/2$, is
- **36.** The maximum number of isomers (including stereoisomers) that are possible on monochlorination of the following compound is

$$\begin{array}{c} \operatorname{CH_3} \\ \mid \\ \operatorname{C} \\ \operatorname{CH_3CH_2} \\ \mid \\ \operatorname{CH_2CH_3} \end{array}$$

37. The total number of contributing structure showing hyperconjugation (involving C—H bonds) for the following carbocation is

- **38.** Among the following, the number of compounds that can react with PCl_5 to give $POCl_3$ is O_2 , CO_2 , SO_2 , H_2O , H_2SO_4 , P_4O_{10}
- **39.** The volume (in mL) of 0.1 M AgNO₃ required for complete precipitation of chloride ions present in 30 mL of 0.01 M solution of $[\text{Cr}(\text{H}_2\text{O})_5\text{Cl}]\text{Cl}_2$, as silver chloride is close to
- **40.** In 1 L saturated solution of AgCl $[K_{\rm sp}({\rm AgCl}) = 1.6 \times 10^{-10}]$, 0.1 mol of CuCl $[K_{\rm sp}({\rm CuCl}) = 1.0 \times 10^{-6}]$, is added. The resultant concentration of Ag⁺ in the solution is 1.6×10^{-x} . The value of "x" is
- **41.** The number of hexagonal faces that are present in a truncated octahedron is

Matrix-Matching-Type Questions

This section contains **2 questions.** Each question has four statements (a, b, c and d) given in **Column I** and five **statements** (p, q, r, s and t) in **Column II.** Any given statement in **Column I** can have correct matching with **one or more** statement(s) given in **Column II.** For example, if for a given question, statement b matches with the statements given in q and r, then for the particular question, against statement b, darken the bubbles corresponding to q and r in the ORS.

42. Match the transformations in **Column I** with appropriate options in **Column II**.

Column I

- (a) $CO_2(s) \rightarrow CO_2(g)$
- (b) $CaCO_3(s) \rightarrow CaO(s) + CO_2(g)$
- (c) $2H \bullet \rightarrow H_2(g)$
- (d) $P_{(white, solid)} \rightarrow P_{(red, solid)}$

Column II

- (p) phase transition
- (q) allotropic change
- (r) ΔH is positive
- (s) ΔS is positive
- (t) ΔS is negative
- **43.** Match the reaction in **Column I** with appropriate types of steps/reactive intermediate involved in these reactions as given in **Column II**.

Column I

Column II

(p) Nucleophilic substitution

(b)
$$(CH_2)_3Cl \xrightarrow{CH_3Mgl} \qquad (q) \ Electrophilic substitution$$

$$(CH_2)_3Cl \xrightarrow{LH_3Mgl} \qquad (r) \ Dehydration$$

$$(CH_2)_3OH \xrightarrow{L_2SO_4} \qquad (r) \ Dehydration$$

$$(CH_2)_3C(CH_3)_2 \xrightarrow{H_2SO_4} \qquad (s) \ Nucleophilic addition$$

$$(d) \qquad (d)$$

(t) Carbanion

Answers

 H_3C

CH₃

2. c **3.** c **4.** b **5.** d **1.** a **6.** c **7.** a **8.** d **9.** c **10.** d **11.** c **12.** a **13.** d **14.** b **15.** a **16.** b, c **17.** a, b, c, d **18.** a, b, d **19.** a, c, d 20. a, d 21. a, b, c, d **22.** a, b, d **23.** b, c, d **24.** d **25.** b **28.** c **26.** b **27.** a

	29	30	31	32	33	34	35	36	37	38	39	40	41
0	0	0	0	0	0	0	0	0	0	0	0	0	0
1	0	0	0	0	0	0	0	0	0	0	0	0	0
2	0	0	0	0	0	0	0	0	0	0	0	0	0
3	0	0	0	0	0	0	0	0	0	0	0	0	0
4	0	0	0	0	0	0	0	0	0	0	0	0	0
5	0	0	0	0	0	0	0	0	0	0	0	0	0
6	0	0	0	0	0	0	0	0		0	0	0	0
7	0	0	0	0	0	0	0	0	0	0	0	0	0
8	0	0	0	0	0	0	0	0	0	0	0	0	0
9	0	0	0	О	О	0		0	0	0	0	0	0