

BHU M.Sc. CHEMISTRY ENTRANCE - 2011

1. The change in entropy when two moles of a monatomic perfect gas is compressed to half its volume and simultaneously heated to twice its initial temperature is

(a) R ln 2	(b) 3 R ln 2	(c) 5 R ln 2	(d) 7 R ln 2
			(u) / K III Z

2. A heat engine operates between 100 K and 600 K. The heat discharged into the cold sink in a reversible process when 5 kJ of heat is supplied by the hot source, is

(a) 2 kJ (b) 2.5 kJ (c) 3 kJ (d) 5.5 kJ

- **3.** For which of the following processes q = 0, w = 0, $\Delta U = 0$ and $\Delta H = 0$?
 - (a) Reversible isothermal process in a perfect gas
 - (b) Reversible adiabatic process in a perfect gas
 - (c) Adiabatic expansion of a perfect gas into vacuum
 - (d) Reversible constant-volume process in a perfect gas
- **4.** The fugacity of a certain gas at 200 K and 50 bars is 25 bars. The difference of its chemical potential from that of a perfect gas in the same state is

(a) 200 R ln 25 (b) 200 R ln 2 (c) -200 R ln 2 (d) -200 R ln 25

- **5.** The entropy of mixing 1 mole of hexane with 1 mole of heptane at 298 K is 11.4 JK⁻¹. The Gibbs energy of mixing (assuming the solution to be ideal) is
 - (a) -1.72 kJ (b) -3.44 kJ (c) 1.72 kJ (d) 3.44 kJ
- 6. The number of degrees of freedom of the system $KClO_3(s) \rightleftharpoons 2KCl(s) + 3O_2(g)$ is

(a) zero	(b) one	(c) two	(d) three
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7. The expression that relates the partial molar properties of the components in a mixture is known as

(a) van't Hoff equation	(b) Gibbs-Duhem equation
(c) Duhem-Margules equation	(d) Raoult's law
For a very dilute electrolyte solution with γ	$\gamma_{_+} <$ 1, $\gamma_{_+}$ should increase with increase in
(a) solvent's density	(b) solvent's dielectric constant
(c) ionic strength	(d) cationic charge

8.



- **9.** Which of the following statements is not correct?
 - (a) ΔG_{mix} at constant temperature and pressure must be negative
 - (b) Intermolecular interactions are negligible in an ideal solution
 - (c) Solute-solute interactions are negligible in an ideally dilute solution
 - (d) Activity coefficients are never negative
- **10.** Which of the following statements is correct?

(a) If $\Delta G^{\circ} > 0$, no amount of products can be obtained when the reaction is run at constant temperature and pressure

(b) It is possible for the entropy of a closed system to decrease substantially in an irreversible process

- (c) In any closed system with P-V work only, G is always minimized at equilibrium
- (d) $\Delta(TS) = T\Delta S + S\Delta T$
- **11.** The condition for the attainment of phase-equilibrium in a closed electrochemical system is the equality of
 - (a) surface potentials (b) electric potentials
 - (c) chemical potentials (d) electrochemical potentials
- **12.** The direction of a chemical reaction at constant temperature and pressure is the direction of
 - (a) decrease of Gibbs free energy of the system
 - (b) decrease of Helmholts free energy of the system
 - (c) increase of entropy of the system
 - (d) decrease of enthalpy of the system
- **13.** The quantum yield of the photochemical decomposition of HI

HI + hv \rightarrow H + I H + HI \rightarrow H₂ + I I + I + M \rightarrow I₂ + M with respect to HI is (a) 0.5 (b) 1 (c) 1.5 (d) 2

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14. HI is absorbed strongly on gold. Assuming Langmuir isotherm to apply, the order of the reaction $2HI \rightarrow H_2 + I_2$ on gold is (b) 0.25 (c) 0.5 (d) 1 (a) zero 15. Which of the following statements is not correct for Langmuir isotherm? (a) It applies to monolayer adsorption (b) Under conditions $\theta \ll 1$ and α (Freundlich exponent) ≈ 1 , it reduces to Freundlich isotherm (c) It applies to dissociative (d) It applies to chemisorption 16. The activation energy and entropy of a bimolecular gas phase reaction at 600 K are 200 kJ mol⁻¹ and -200 JK⁻¹ mol⁻¹ respectively. The free energy of activation is (a) 70 kJ mol-1 (b) 80 kI mol⁻¹ (c) 310 kI mol⁻¹ (d) 320 kJ mol-1 17. Among the following statements which is the correct one? (a) The heat of chemisorption is always larger than that of physisorption (b) Langmuir isotherm specifically assumes the existence of active centres (c) Promoters are themselves catalysts (d) Increase in surface tension with concentration leads to negative adsorption 18. The activation energy of the gas-phase association between F_2 and IF_5 , a first-order reaction in each of the reactants, is 58.6 kJ mol⁻¹. The activation enthalpy at 340 K is (a) 53 kJ mol⁻¹ (b) 55.8 kJ mol⁻¹ (c) 58.6 kJ mol⁻¹ (d) 61.4 kJ mol⁻¹ In photochemical reaction $A \rightarrow 2B+C$, the quantum efficiency with 500 nm light is 19. 2×10^2 mol Einstein-1. After exposure of 300 m moles of A to the light, 2 m moles of B is formed. The number of photons absorbed by A is (b) 3×10^{18} (a) 1×10^{18} (c) 6×10^{18} (d) 9×10^{18} 20. The condition for which the reaction rate of an enzymolysis that follows Michaelis-Menten kinetics, is half its maximum value, is (a) $[S] \ll K_{M}$ (b) $[S] = K_{M}$ (c) $[S] = K_{M} / 2$ (d) $[S] \gg K_{M}$ 21. For the mechanism

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 $A_2 \rightleftharpoons 2A(fast)$ $A + B \rightarrow P(slow)$

the reaction order with respect to A_2 is

- (a) 0 (b) 0.5 (c) 1 (d) 2
- **22.** Which of the following relations does not hold for the activity (A) of a radioactive substance?

(a)
$$\frac{A}{A_0} = \left(\frac{1}{2}\right)^{t/t_1}$$

(b) $\frac{A}{A_0} = \exp\left(-0.693\frac{t}{t_{0.5}}\right)$
(c) $t_{0.5} / t_{0.1} = \ln 2$
(d) $\frac{A}{A_0} = 1 - 0.693\frac{t}{t_{0.5}}$ at $t \to 0$

23. A powder diffraction photograph from tungsten shows lines which indices as (110), (200), (211), (220), (310), (222), (321), (400),.... The symmetry of the unit cell is

(a) primitive (b) end-centred (c) face-centred (d) body-centred

24. Among the following halides which one forms van der Walls crystals?

(a) NaCl (b) BeCl_2 (c) HgCl_2 (d) HF

25. A form of $CaCO_3(c)$ has orthorhombic lattice with a = 5.0 Å, b = 8.0 Å, c = 5.6 Å and density = 3.0 gm cm⁻³ at room temperature. The number of Ca²⁺ ions per unit cell of the crystal is

(a) 1 (b) 2 (c) 4 (d) 8

26. The ratio of the translational partition functions of D_2 and H_2 at the same temperature and volume is

- (a) 2 (b) 1.414 (c) 2.83 (d) 4
- **27.** In which of the following systems is the energy levels separation the largest?
 - (a) an electro in a radical in a field of 0.300 T
 - (b) a ¹⁴N nucleus in 600 MHz NMR spectrometer
 - (c) a proton in the same spectrometer

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(d) a deuteron in the same spectrometer

28.	Which of the following functions is not an eigen function of $\frac{d^2}{dx^2}$?				
	(a) cos kx	(b) $\exp(-kx^2)$	(c) kx	(d) $\exp(ikx)$	
29.		molar masses of a g tation and osmotic pre		as determined by light nethods is	
	(a) 1:1:2	(b) 2:1:2	(c) 1:2:1	(d) 2:2:1	
30.		ctobacillus acidophilus tive to the initial value	0	e of 75 minutes. What is	
	(a) 4.0	(b) 2.25	(c) 2.0	(d) 1.75	
31.	If the pressure of a	gas at constant temper	ature is doubled, the v	iscosity of the gas will be	
	(a) quadrupled	(b) doubled	(c) halved	(d) unchanged	
32.	Among the followin	g molecules which one	e shows pure rotation	spectra?	
	(a) N ₂	(b) H ₂ S	(c) CO ₂	(d) CH ₄	
33.	The SI unit of radia	tion dose is			
	(a) becquerel	(b) curie	(c) rad	(d) gray	
34.	state from the grou		f frequency v. If the b	L can be excited to n = 2 od length is doubled, the	
	(a) v/4	(b) v/2	(c) 2v	(d) 4v	
35.	For a hydrogen ato angular momentum		he maximum possible	e z-component of orbital	
	(a) 2ħ	(b) 3ħ	(c) $\sqrt{12\hbar}$	(d) √ <u>6</u> ħ	
36.	Which one of the following is an acceptable approximate wave function for a state of the helium atom?				
	(a) $[1s(1)1s(2)-1]$	$s(1)1s(2)][\alpha(1)\beta(2)]$)]		
	(b) $[1s(1)1s(2)][\alpha(1)\alpha(2)+\beta(1)\beta(2)]$				

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(c)
$$[1s(1)2s(2)+2s(1)1s(2)][\alpha(1)\alpha(2)]$$

(d)
$$\left[1s(1)2s(2)+2s(1)1s(2)\right]\left[\alpha(1)\beta(2)-\beta(1)\alpha(2)\right]$$

37. Which one of the following statements concerning H_2^+ is incorrect?

(a) The non-degenerate LCAO-MOs (without spin) must be either symmetric or antisymmetric for inversion

(b) The lowest energy MO (without spin) of the molecule is anti-symmetric for inversion

(c) The ground state has a multiplicity of two

- (d) The MOs transform into AOs of the helium ion as the two nuclei are fused together
- **38.** Which one of the following is the correct formula for the lowest energy eigen function for a particle in a one-dimensional box having infinite barriers at x = -L/2 and L/2?

(a)
$$\sqrt{\frac{2}{L}}\sin\left(\frac{\pi x}{L}\right)$$
 (b) $\sqrt{\frac{2}{L}}\cos\left(\frac{\pi x}{L}\right)$ (c) $\sqrt{\frac{2}{L}}\exp\left(\frac{i\pi x}{L}\right)$ (d) $\sqrt{\frac{2}{L}}\exp\left(\frac{-i\pi x}{L}\right)$

39. Which of the following equations is used to calculate the number of theoretical plates?

(a)
$$\frac{t_{R} - t_{0}}{t_{0}}$$
 (b) $16 \left(\frac{t_{R}}{W}\right)^{2}$
(c) $\frac{k_{2}}{k_{1}}$ (d) $2 \left(t_{R_{2}} - t_{R_{1}}\right) / \left(W_{1} + W_{2}\right)$

- **40.** Which of the following techniques is based on selectively inducing radioactivity and measuring the emitted radiation?
 - (a) Isotope dilution analysis (b) Radiometric titration
 - (c) Neutron activation analysis (d) All of the above
- **41.** Which of the following techniques can be used only for volatile compounds?
 - (a) Gas chromatography (b) HPLC
 - (c) Ion chromatography (d) All of the above
- **42.** Which of the following refers to ion exchange capacity
 - (a) Nature of exchanging ions

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	(b) Nature of strong cation exchanger				
	(c) Nature of strong	anion exchanger			
(d) Total number of ion active groups per unit length of material					
43. In electrogravimetry of cations the working electrode is					
	(a) anode		(b) cathode		
	(c) both cathode and	anode	(d) neither anode no	r cathode	
44.			= =	the determination of a or for determination by	
	(a) Neutron activatio	on analysis	(b) Isotope dilution a	analysis	
	(c) Radiometric titra	tion	(d) All of the above		
45.	Which of the following	ng frequencies corresp	oonds to carbonyl stret	cch vibration in acids?	
	(a) 1625 cm ⁻¹	(b) 1715 cm ⁻¹	(c) 1745 cm ⁻¹	(d) 1800 cm ⁻¹	
46.	A particular vibration in a polyatomic molecule is IR active if during vibration, the change in				
	(a) polarizability	(b) dipole moment	(c) frequency	(d) potential energy	
47.	Moisture in a drug ca	an be determined by			
	(a) Malaprade reage	nt	(b) EDTA reagent		
	(c) Karl Fischer reag	ent	(d) chloramine-T rea	agent	
48.	Which one is more to	oxic?			
	(a) Hg	(b) $\left(CH_3\right)_2 Hg$	(c) Hg ²⁺	(d) Hg_2^{2+}	
49.	The most efficient te	chnique for the separa	tion of amino acids is		
	(a) adsorption chron	natography	(b) partition chroma	tography	
	(c) ion-exchange chr	omatography	(d) paper chromatog	raphy	
50.	Which one is not a p	ollutant?			
	(a) CO	(b) CO ₂	(c) SO ₃	(d) NO ₂	
51.	Which one is the sinl	k of CO_2 ?			
	(a) Plant	(b) Ocean	(c) Air	(d) Soil	

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52.	• Ozone layer is a protective shield against					
	(a) visible light	(b) ultraviolet light	(c) infrared rays	(d) cosmic rays		
53.	Nessler's reagent is					
	(a) KHgI ₄		(b) $K_2HgI_4 + NH_4O$	Н		
	(c) K ₂ HgI ₄		(d) $KHgI_4 + NH_4OH$	I		
54.	Which of the follow	Which of the following statements is not correct?				
	(a) In instrumental	(a) In instrumental methods of analysis visual indicators are not required				
	(b) In classical volu	(b) In classical volumetric titrations visual indications are not required				
	(c) In classical qualitative analysis removal of interfering radicals is essential					
	(d) Spectroanalytic	al techniques can be ap	plied for qualitative a	nd quantitative analysis		
55.	Visible region in the electromagnetic spectrum extends from					
	(a) 200-400 nm	(b) 800-900 nm	(c) 400-4000 cm ⁻¹	(d) 400-800 nm		
56.	Hard water can be softened in					
	(a) a calorimeter		(b) a chromatograph			
	(c) an ion exchange	er (d) an earthen ware	pot			
57.	The technique used	The technique used for the separation of components of a mixture is called				
	(a) chromatograph	У	(b) IR spectroscopy			
	(c) electornic spect	roscopy	(d) polarography			
58.	Essential constitue	Essential constituent of an amalgam is				
	(a) Ag	(b) Hg	(c) Mg	(d) Fe		
59.	Potential of which o	Potential of which of the following electrodes does not depend on pH of the solution?				
	(a) Glass electrode		(b) Hydrogen electrode			
	(c) Quinhydrone el	ectorde	(d) calomel electro	le		
60.	The unit of equivale	ent conductance is				
	(a) ohm ⁻¹ cm ⁻¹		(b) ohm ⁻¹ cm ⁻²			
	(c) ohm ⁻¹ cm ²		(d) None of the above	ve		
61.	Which quantity ren	nains unchanged on cha	inging temperature?			

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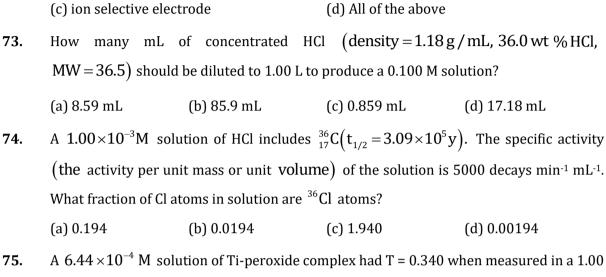
	(a) Mole fraction		(b) Molarity	
	(c) Normality		(d) None of the above	
62.	The strongest ligand	in spectro-chemical se	eries is	
	(a) F-	(b) OH-	(c) H ₂ O	(d) CO
63.	The iron is rusted, th	ien it is		
	(a) oxidized	(b) reduced	(c) evaporated	(d) decomposed
64.	Two elements canno	t be combined chemic	ally to make	
	(a) a compound		(b) another element	
	(c) a gas		(d) a liquid	
65.	Choose the law that radiations only"	states, "effective cher	nical changes are brow	ught about by absorbed
	(a) Beer's law		(b) Grotthus-Draper	law
	(c) Lambert's law		(d) law of photochemical equivalence	
66.	What quantity of lim	estone on heating will	give 56 kg of CaO?	
	(a) 10 kg	(b) 55 kg	(c) 44 kg	(d) 100 kg
67.	The unit of absorptiv	vity is		
	(a) cm ⁻¹ g ⁻¹ L	(b) cm ⁻¹ mol ⁻¹ L	(c) cm ⁻¹ g L ⁻¹	(d) No unit
68.	Oxine is a			
	(a) precipitating rea	gent	(b) redox reagent	
	(c) drug		(d) dye	
69.	Free radicals may be	detected by		
	(a) mass spectromet	ry	(b) NMR spectroscop	ру
	(c) ESR spectroscopy	/ (d) infrared spectros	зсору	
70.	Lead can be the best	analyzed colorimetric	ally using which of the following reagents?	
	(a) Dithizone	(b) EDTA	(c) DMG	(d) Acetyl acetone
71.	Which of the followi	ng is a hydride ion dor	ior?	
	(a) NAD	(b) ATP	(c) NADH	(d) Coenzyme A
72.	Na and K can be esti	mated using		

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(a) X-ray fluorescence(b) IR spectroscopy



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75. A 6.44×10^{-4} M solution of Ti-peroxide complex had T = 0.340 when measured in a 1.00 cm cell at 410 nm. What is the concentration of Ti-peroxide complex in a solution that had T = 0.516 when measured at 410 nm in a 0.500 cm cell?

(a) 7.89×10^{-2} M (b) 15.78×10^{-2} M (c) 7.89×10^{-4} M (d) 3.95×10^{-3} M

76. The partition coefficient for ethyl iodide, E, between octanol and water is

$$K_{p} = \frac{\left[E\right]_{0}}{\left[E\right]_{w}} \approx 100$$

What percent of E present in 50.00 mL of water would remain if the water was extracted with 10.00 mL of octanol?

(a) 47.6% (b) 4.76% (c) 9.52 (d) 95.2%

77. Arrange the ionization energies of the following ions in increasing order

$$\begin{array}{l} (A) \operatorname{Fe}^{2+} \to \operatorname{Fe}^{3+} \\ (B) \operatorname{Mn}^{2+} \to \operatorname{Mn}^{3+} \\ (C) \operatorname{Fe}^{+1} \to \operatorname{Fe}^{2+} \end{array}$$

$$\begin{array}{l} \text{Choices are} \\ (a) \ A < B < C \qquad (b) \ C < B < A \qquad (c) \ C < A < B \qquad (d) \ B < C < A \end{array}$$

$$(At. \operatorname{No. of} \operatorname{Fe} and \operatorname{Mn} are 26 and 25) \end{array}$$

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78. In the first row transition metal ion (2+) having four electrons in 3d orbitals (with 4s and 4p orbitals empty), the metal ion forms a hypothetical diamagnetic tetrahedral complex $[MA_4]^{2-}$ with A- anion. The electronic configuration of the M^{2+} ion in the complex is

(a)
$$[Ar] 3d_{xy}^2 3d_{xz}^2 3d_{yz}^0 3d_{z^2}^0 3d_{x^2-y^2}^0 4s^0 4p^0$$
 (b) $[Ar] 3d_{xy}^0 3d_{xz}^0 3d_{yz}^2 3d_{z^2}^2 3d_{x^2-y^2}^0 4s^0 4p^0$
(c) $[Ar] 3d_{xy}^0 3d_{xz}^0 3d_{yz}^0 3d_{z^2}^2 3d_{x^2-y^2}^2 4s^0 4p^0$ (d) $[Ar] 3d_{xy}^1 3d_{xz}^1 3d_{yz}^2 3d_{z^2}^0 3d_{x^2-y^2}^0 4s^4 p^0$

79. Out of the following which has the least tendency to form M = 0 species?

(I)Sc=0	(II)P=O	(III)Ln=0	(IV)Ac=0
(a) Sc = 0 and	P = 0		(b) Ln = 0
(c) Ac = 0			(d) $Sc = 0, P = 0 and Ac = 0$

(Ln = Lanthanides and Ac = actinides)

80. The magnetic properties of lanthanides are largely determined by the formula

(a)
$$\mu_{eff} = \sqrt{n(n+2)}$$

(b) $\mu_{eff} = \sqrt{n(n+2)}$ + some contribution of orbital contribution
(c) $\mu_{eff} = \sqrt{L(L+2)}$
(d) $\mu_{eff} = g_J \sqrt{J(J+1)}$, where $g_J = \frac{3}{2} + \frac{S(S+1) - L(L+1)}{2J(J+1)}$

- **81.** Ln^{2+} ions are largely coloured while Ln^{3+} ions are not coloured. The colour of Ln^{2+} compounds are due to transition
 - (a) $4f \rightarrow 5d$
 - (b) $4f \rightarrow 4f$
 - (c) charge transfer from liquid to metal ion
 - (d) $5d \rightarrow 5d$ transition

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- 82. Transition metals (some ions) form complexes with molecular N_2 . There are two modes of coordination of N_2 in the complexes of the type $[M(N_2)(L)_n]$. These modes are (a) end-on and (b) side-on. The stability of these complexes in these two modes is
 - (a) end-on more stable than side-on
 - (b) side-on more stable than end-on
 - (c) both are highly unstable (the relative stability depends on ligands)
 - (d) both are equally stable
 - (L = ligand)
- **83.** The structure of hyponitrous acid molecule $(H_2N_2O_2)$ is
 - (a) linear (A-B-C-D type) (b) bent $(\frown \text{ type})$ (c) square planar $(\Box \text{ type})$ (d) triangular $(\triangle \text{ type})$
- 84. We have two complexes $[Mn(H_2O)_6]^{3+}$ ion $(d^4 \text{ system})$ and $[Re(H_2O)_6]^{3+}$ ion $(d^4 \text{ system})$. Their CFSE and the pairing energy (per pair of electrons) are given below

$$\left[Mn(H_2O)_6\right]^{3+} - 10Dq = 250 \text{ kJ / mol; Pairing energy} = 300 \text{ kJ / mol}$$
$$\left[Re(H_2O)_6\right]^{3+} - 10Dq = 400 \text{ kJ / mol; Pairing energy} = 180 \text{ kJ / mol}$$

Predict from their data, the complexes are

- (a) both high spin (b) Mn-high spin and Re-low spin
- (c) both low spin (d) Mn-low spin, Re-high spin
- 85. $\left[\operatorname{NiCl}_{2}(\operatorname{PPh}_{3})_{2}\right]$ is paramagnetic with $\mu_{eff} = 2.9 \text{ BM}$. All the four ligands are monodentate. The geometry of the molecule is
 - (a) square planar (b) octahedral
 - (c) square pyramidal (d) tetrahedral

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- 86. There is no d-d transition in $\left[Mn(H_2O)_6 \right]^{2+} (d^5 \text{ system})$. This is high spin complex. The absence of any transition in the visible region is due to breaking of
 - (a) Laporte
 - (b) Parity selection rule $(g \rightarrow g \text{ and } u \rightarrow u \text{ not allowed})$
 - (c) spin selection rule (spin of the electron cannot be changed during transition)
 - (d) All the above three rules (1, 2 and 3)
- **87.** According to the IUPAC system which is the correct formula for the complex [diaquadibromodi(methylamine)] cobalt (III) nitrate?
 - (a) $\left[\text{Co}(\text{H}_2\text{O})_2 \text{Br}_2(\text{MeNH}_2)_2 \right] \text{NO}_3$ (b) $\left[\text{CoBr}_2(\text{H}_2\text{O})_2(\text{MeNH}_2)_2 \right] \text{NO}_3$
 - (c) $\left[Co(H_2O)_2(MeNH_2)_2 Br_2 \right] NO_3$ (d) $\left[Co(MeNH_2)_3(H_2O)_2 Br_2 \right] NO_3$
- **88.** $\left[Cr(H_2O)_3 \right]^{3+}$ is violet whereas $\left[Cr(NH_3)_6 \right]^{3+}$ is yellow. The wavelength absorbed by $\left[Cr(NH_3)_6 \right]^{3+}$ as compared to that by $\left[Cr(H_2O)_6 \right]^{3+}$ in their absorption spectra will be
 - (a) higher
 - (b) lower
 - (c) same
 - (d) These complexes will not show any absorption in the visible region
- **89.** Although fluorine is better oxidizing agent than oxygen, but Mn_2O_7 exists and MnF_7 does not. This is because of

(a) steric crowding and difficult to fitting seven fluoride ions around Mn centre (b) very high electron affinity of fluorine as compared to oxygen

- (c) very low lattice energy of $MnF_{\!7}\,$ compared to oxygen compound
- (d) very low bond energy of Mn F bond compared to Mn O bond
- **90.** Covalent radius of gold (125 pm) is less than that of silver (1.33 pm). This is because of
 - (a) transition metal contraction



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(b) lanthanide contraction

(c) lack of shielding of 4d orbitals compared to 5d orbitals

(d) relatively less effective shielding by 3 p and 3s orbitals in silver as compared to that by 4s, 4p and 4d

91. Using VSEPR model, the shaper of the following molecules

$$\operatorname{IF}_{5}^{-2}(A)$$
, $\operatorname{XeF}_{4}(B)$, $\operatorname{IO}_{3}^{-}(C)$, $\operatorname{BrF}_{3}(D)$

are

- (a) (A) pentagon shape, (B) square planar, (C) trigonal pyramid, (D) T-shape
- (b) (A) trigonal bipyramid, (B) tetrahedral, (C) planar, (D) planar
- (c) (A) square pyramid, (B) tetrahedral, (C) trigonal pyramid, (D) planar
- (d) (A) pentagonal pyramid, (B) tetrahedral, (C) trigonal pyramid (D) planar

Arrange the hydrides of 15-group elements $(NH_3, PH_3, AsH_3, SbH_3 < BiH_3)$ in 92. increasing contribution of p-orbital of X in the X-H bond. Choose the correct alternative from the following alternatives given below

- (a) Bi < Sb < As < P < N(b) P < N < As < Bi < Sb
- (d) Bi < Sb < N < P < As(c) N < P < As < Sb < Bi
- 93. What are A, B and C in the following reactions?
 - $(I) B_2 H_6 + N H_3 \rightarrow (A)$ $(II)B_2H_6 + H_2O \rightarrow (B)$ $(III)B_2H_6 \xrightarrow{\text{Heated at } 100^\circ C} (C)$ Here A, B and C respectively are
 - (a) borazine, B_2O_3 and B_4H_8 (b) $2BH_3$. NH₃, H₃BO₃ + H₂, B₁₀H₁₄
 - (d) BN, $H_3BO_3 + H_2$, B_4H_8 (c) BN, H_3BO_3 only, B_4H_8
- 94. Which one of the following contain (3c-2e) bonds?

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(C)BeH₂,

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 $(D)Be(NO_3)_2$

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 $(B)BeCl_2$,

 $(A) Mg(CH_3)_2$,

(a) B

95.

96.

97.

(a) A. B and C (b) A and C (c) A. C and D (d) B and C Which one of the following orders of two properties of 14- and 15-group elements is not correct? $(A)BI_3 < BBr_3 < BCl_3 < BF_3$ (Lewis acidity) $(B)Pb(CH_3)_4 < Sn(CH_3)_4 < Ge(CH_3)_4 < Sc(CH_3)_4$ (thermal stability) The correct alternative from the following ones is (b) A (c) No one is wrong (d) Both are wrong Which ones of the following compounds do not exist? $[AuXe_4], [KrF_2], [ArF_2][He-F]$ Choose the correct choice out of the following (b) $[KrF_2] and [ArF_2]$ (a) $[AuXe_4] and [He-F]$ (c) $[Ar - F_2]$ and [He - F](d) $[ArF_2][He-F], [AuXe_4]$ Alkalides and electrides are crystalline compounds of (a) alkali metals in (+1) oxidation states and electrons respectively (b) alkali metals in (-1) oxidation states and electron acting as anion (c) methyl group and electron acting as anion respectively

(d) methyl group and alkali metal group

98. Among the molecules, BiF₃, BiCl₃, BiBr₃ and BiI₃, the one which is most coloured is

(a)
$$BiF_3$$
 (b) $BiBr_3$ (c) $BiCl_3$ (d) BiI_3

- $\left[\text{Cr}\text{O}_{_{\!R}}\right]^{_{\!3^-}}$ ion is known. This ion probably is 99.
 - (a) complex of peroxide ion with Cr^{5+} ion
 - (b) polymeric complex with oxide ion acting as bridges



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(c) complex of molecular oxygen and oxide ion $(e.g. Cr^{5+} (O_2)_2 O_4)^{3-}$

(d) complex of molecular oxygen and peroxide ions

- **100.** Iodine dissolves in oleum to give bright blue colour. The blue colour is due to the formation of A, where A is
 - (a) I^- ion (b) I_3^- ion (c) I_2^+ ion (d) I_4^+ ion
- **101.** The order of increasing size of V, Nb, Ta, Db (at. no. 106) is

(a)
$$V < Nb < Ta < Db$$
 (b) $Db < Ta < V < Nb$

- (c) $V < Nb \approx Ta \approx Db$ (d) V < Db < Ta < Nb
- **102.** Given the following reaction conditions for the formations of the fluorides of Xe

$$Xe(g) + F_{2}(g) \xrightarrow{400^{\circ}C} A... (Xe in excess)$$
$$Xe(g) + F_{2}(g) \xrightarrow{600^{\circ}C} B... (Xe:F_{2} = 1:1.6)$$
$$Xe(g) + F_{2}(g) \xrightarrow{300^{\circ}C} C... (Xe:F_{2} = 1:20)$$

A, B and C in these reactions respectively are

(a)
$$XeF_4$$
, XeF_2 , XeF_6 (b) XeF_2 , XeF_4 , XeF_6

(c)
$$XeF_6$$
, XeF_4 , XeF_2 (d) XeF_2 , XeF_6 , XeF_4

- 103. In the preparation of P_4O_6 , a mixture of N_2 and oxygen is used instead of pure oxygen. The reason is
 - (a) pure oxygen will form ozone in presence of P
 - (b) pure oxygen will form P_4O_{10} despite excess of P_4 , in pure oxygen
 - (c) pure oxygen will form $\left\lceil P(O_3)_4 \right\rceil$ ozonied
 - (d) pure oxygen will not react with $P.N_2$ acts as a catalyst
- 104. Arsenic, antimony and bismuth react with concentrated HNO₃. These give respectively
 - (a) H_3AsO_4 , H_3SbO_2 , $(BiO_3)^-$ ion (b) H_3AsO_4 , Sb_2O_3 , $(BiO_3)^-$ ion (c) As_2O_3 , Sb_2O_3 , Bi_2O_3 (d) H_3AsO_4 , Sb_2O_5 and Bi^{3+} ion



- **105.** In borazine $(B_3N_3H_6)$ molecule, the number of isomers which are possible of its disubstituted borazine molecule of the formula $[B_3N_3H_4X_2]$ without changing its ring structure is
 - (a) one (b) two (c) four (d) six
- 106. Solution of alkali metals in liquid NH_3 conducts electricity. It is due to formation of
 - (a) $Na^+ + Na^-$ ion in liquid ammonia
 - (b) $Na^+ + e^- (NH_3)_{\downarrow}$ in liquid ammonia
 - (c) Na⁻, $(NH_2)^-$ and NH_4^+ ion in liquid NH_3

(d) The solution conducts like a metallic conductor with solvated electron carrying the charge

107. A compound alloy of metals P and Q has a unit cell containing P atom at the corners, while atom Q are present at the face centres. The formula of the compound should be

(a) PQ (b) PQ_2 (c) PQ_3 (d) P_3Q

108. A complex of a certain metal ion has a magnetic moment of 4.90 BM (4 unpaired electrons) and another complex of the same metal ion in the same oxidation state is diamagnetic. The complex is octahedral. The central metal could be one of the following

 $\left(Cr^{_{3+}}$, $Mn^{_{2+}}$, $Fe^{_{2+}}$, $Mn^{_{3+}}$, $Fe^{_{3+}}$, $Co^{_{2+}}
ight)$

 Cr^{3+} ion is $[Ar]4d^3$ system. The metal ion is

(a) Mn^{2+} (b) Fe^{2+} (c) Co^{2+} (d) Fe^{3+}

109. The enthalpy of hydration (ΔH) of $\operatorname{Cr}^{2+}[(d^4 \operatorname{system} \operatorname{octahedral} \operatorname{complex} (\operatorname{Cr}(H_2O)_6)^{2+}]$ is -460 kcal/mol. In the absence of CFSE, this value of ΔH is (-435) kcal/mole. The value of 10 Dq (or Δ_0) is (a) 14600 cm⁻¹ (b) 21500 cm⁻¹ (c) 9525 cm⁻¹ (d) 25252 cm⁻¹ (1 kcal = 350 cm⁻¹)

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110.	Which of the d^n ion will have the smallest CFSE if $\Delta(or 10Dq)$ is greater than pairing energy d^6 , d^7 , d^8 , d^9 , d^{10} ?				
	(a) d^6, d^{10}	(b) d ⁷ ,d ⁹	(c) d^{10}	(d) d^5 , d^8 , d^{10}	
111.	water to form its con	[*]	s. These, in turn, lowe	es \underline{A} which reacts with rs the pH in the muscles	
	(a) citric acid and H	₃ 0 ⁺	(b) lactic acid and H	₃ 0 ⁺	
	(c) enzyme and NH_4^+	(d) gluconic acid and	1 NH_4^+		
112.	Which ones of the fo	llowing are closed pac	ked structures?		
	1. AABB AABB		2. ABAC ABAC		
	3. AB AB AB		4. ABC ABC ABC		
	(a) 1, 2, 3	(b) 1, 3, 4	(c) 2, 3, 4	(d) 2, 3	
113.	Out of following acti	nides (I) Np, (II)Pu, (III)Es, (IV)Am one h	aving +2 oxidation state	
	as the only relatively	v important state is			
	(a) Np	(b) Pu	(c) Es	(d) Am	
114.	Which is not an amb	ident nucleophile?			
	(a) CN-	(b) SCN [−]	(c) NO_2^-	(d) DMSO	
115.	What is the increasing order of stability of following carbocations (give least stable first)?				
	(I) Tropylium cation	n (II) $CH_2 = CH - CH_2^+$			
	$(III)(C_6H_5)_2C^+$		$(IV)CH_3^+$		
	(a) $III < I < II < IV$	(b) IV < II < III < I	(c) $I < III < II < IV$	(d) $IV < III < II < I$	
116.	The type of rearrang	gement			
	о он он				



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1S	

	(a) Lossen	(b) Beckmann	(c) Schmidt	(d) Curtius	
117.	What is the nucleophilicity order for S_N^2 reaction?				
	$(I)C_{6}H_{5}S^{-}$		$(II)C_2H_5O^-$		
	$(III) NO_3^-$		$(II) C_2 H_5 O^-$ $(IV) CN^-$		
	(V)I ⁻				
	(a) $V > II > IV > I > I$	II	(b) $III > IV > V > II >$	·I	
	(c) $I > IV > V > II > I$	II	(d) $II > IV > V > III >$	· I	
118.	Select order of effect	iveness of Lewis acid c	atalyst in Friedel-Craf	ts reaction	
	(a) $AlCl_3 > FeCl_3 > 7$	$\operatorname{ZnCl}_2 > \operatorname{BF}_3$	(b) $AlCl_3 > BF_3 > Zn$	$Cl_2 > FeCl_3$	
	(c) $AlCl_3 > ZnCl_2 > I$	$BF_3 > FeCl_3$	(d) $AlCl_3 > FeCl_3 > FeCl_3$	$3F_3 > ZnCl_2$	
119.	One of the modern m	ethods of studying fre	e radical is		
	(a) IR spectra		(b) CIDNP		
	(c) UV spectra		(d) microwave spect	ra	
120.	For a reaction betwe	en alkyl halide and OH	I^- increase in solvent	polarity generally	
	(a) decreases the rate of $S_N 1$ reaction				
	(b) increases the rate	e of S_N^{1} reaction			
	(c) increases the rate	e of S_N^2 reaction			
	(d) does not alter the	e rate of $S_N 1$ and $S_N 2$	reaction		
121.	Ethyl acetoacetate is	prepared from ethyl a	cetate by		
	(a) Benzoin condens	ation	(b) Aldol condensatio	on	
	(c) Claisen condensa	tion	(d) Dieckmann conde	ensation	
122.	Conversion of acetop	henone to acetanilide	is best accomplished b	oy using	
	(a) Curtius	(b) Hofmann	(c) Lossen	(d) Beckmann	
123.	The order of the ease	e of the following leavi	ng groups would be		





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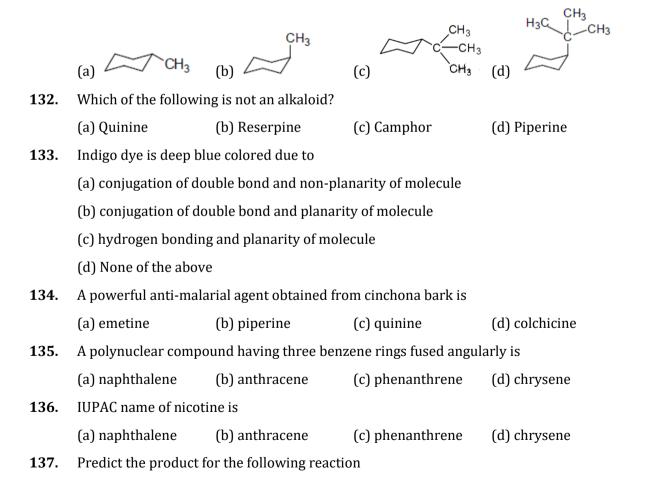
	(I) CH₃COO⁻		(II)CH₃O⁻	
	(III)CH ₃ SO ₃ ⁻		(II)CH ₃ O [−] (IV)CF ₃ SO ₃ [−]	
	(a) IV > III > I > II	(b) $I > II > III > IV$	(c) $I > III > II > IV$	(d) $I > II > IV > III$
124.	In an S_N^2 reaction t	here is		
	(a) complete racemi	zation		
	(b) mostly inversion	and little racemisatio	n	
	(c) partial racemizat	ion		
	(d) a little inversion	and mostly racemisat	ion	
125.		on atoms in pipering is		
	(a) 13	(b) 15	(c) 17	(d) 19
126.	In Beckmann rearra	ngement, the migratin	g group	
	(a) is always syn to t	he hydroxyl group		
	(b) is always anti to	the hydroxyl group		
	(c) is either anti or s	yn		
	(d) depends on the s	stereochemistry of the	molecule	
127.	In Baeyer-Villiger ox	idation, rate of reaction	on is accelerate by	
	(a) electron donatin	g groups in the ketone		
	(b) electron withdra	wing groups in the pe	racid	
	(c) Both 1 and 2 are	correct		
	(d) None of these			
128.	Reactive intermedia	te formed in the follov	ving reaction is	
	NH ₂ Br ₂ NaO	H CO ₂		
	(a) carbene	(b) nitrene	(c) carbocation	(d) carbanion
129.	The increasing orde	r of energy of various	conformations of cyclo	hexane molecule is

(a) twist boat < boat < chair < half chair

- (b) boat < twist boat < half chair < chair
- (c) chair < half chair < twist boat < boat
- (d) chair < twist boat < boat < half chair
- **130.** In cyclohexane molecule, when one chair conformation changes to other



- (a) all hydrogens originally axial become equatiorial
- (b) potential energy of chair conformation increases
- (c) angle strain reduces
- (d) None of the above
- **131.** Which of the following is least stable?



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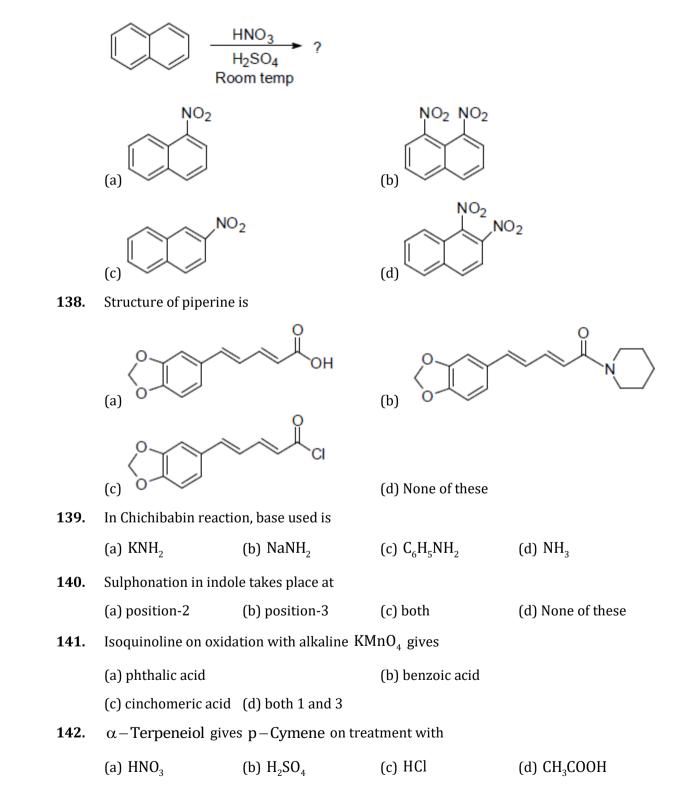
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143. The sweetest sugar amongst the following is			is	
	(a) lactose	(b) fructose	(c) glucose	(d) sucrose
144.	The bond that determines the secondary structure of protein is			
	(a) coordinate bond		(b) covalent bond	
	(c) hydrogen bond		(d) ionic bond	
145.	The main structural feature of protein is			
	(a) ester linkage	(b) ether linkage	(c) peptide linkage	(d) ionic linkage
146.	Buna-S is obtained by the polymerization of butadiene and			
	(a) chloroprene	(b) styrene	(c) acrylonitrile	(d) adipic acid
147.	 7. Terylene is a polymer of (a) adipic acid and hexamethylene diamine (b) terephthalic acid and ethylene glycol (c) phenol and formaldehyde (d) adipic acid and ethylene glycol 			
148.	3. Which is an example of thermosetting polymer?			
	(a) Polyethene		(b) Neoprene	
	(c) Polyvinyl chloride (d) Bakelite			
149.	Which of the following is chain growth polymer?			
	(a) Glyptal	(b) Nylon-66	(c) Nylon-6	(d) Polypropylene
150.	Alkaloids are found in			
	(a) seeds	(b) root	(c) bark	(d) All of the above

