DATE: 29-08-2021

TIME: 10.30 AM TO 11.50 AM

1. For the reaction

$$A(g) + B(g) \rightleftharpoons C(g) + D(g); \Delta H = -QKJ$$

The equilibrium constant cannot be disturbed by

- a) Addition of A
- b) Addition of D
- c) Increasing of pressure
- d) Increasing of temperature

Ans. c

Sol.
$$A_{(g)} + B_{(g)} \rightleftharpoons C_{(g)} + D_{(g)}; \Delta H = -Q KJ$$

Pressure has no effect on equilibrium state if $\Delta n = 0$

- 2. An organic compound 'X' on treatment with PCC in dichloromethane gives the compound Y. Compound Y' reacts with I₂ and alkali to form yellow precipitate of triiodomethane. The compound X is
 - a) CH₃CHO
- b) CH₃COCH₃
- c) CH₃CH₂OH
- d) CH₃COOH

Ans. c

Sol.

$$\begin{split} CH_3 - \overset{X}{CH_2} - OH & \xrightarrow{PCC} CH_2 CI_2 \\ & \therefore X = CH_2 CH_2 OH \end{split} \xrightarrow{Y} CHO \xrightarrow{I_2} CHI_3 + HCOONa \\ \end{split}$$

3. A compound 'A' (C_7H_8O) is insoluble in NaHCO₃ solution but dissolve in NaOH and give a characteristic colour with neutral FeCl₃ solution. When treated with Bromine water compound 'A' forms the compound B with the formula $C_7H_5OBr_3$. 'A' is

Ans. b

Sol.
$$A$$

$$OH$$

$$Br$$

$$Br$$

$$OH$$

$$Br$$

$$CH_3$$

$$CH_3$$

4. In set of reactions, identify D

$$CH_{3}COOH \xrightarrow{SOCl_{2}} A \xrightarrow{Benzene} B \xrightarrow{HCN} C \xrightarrow{H_{2}O} D$$

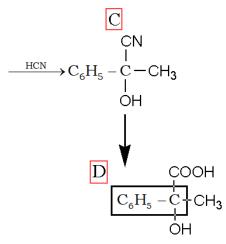
a)
$$CH_2 - CH_3$$

c)
$$CH_2 - \overset{OH}{\overset{|}{C}} - CH_3$$

d)
$$OH$$
 $COOH$ $COOH$ $COOH$ OH

Ans. d

Sol. $CH_3 - COOH \xrightarrow{SoCl_2} CH_3 \stackrel{A}{COCl} \xrightarrow{C_6H_6} COCH_3$

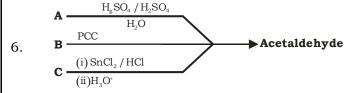


- 5. K_a values for acids H_2SO_3 , HNO_2 , CH_3COOH and HCN are respectively $1.3x10^{-2}$, $4x10^{-4}$, $1.8x10^{-5}$ and $4x10^{-10}$, which of the above acids produces stronger conjugate base in aqueous solution?
 - a) H₂SO₃
- b) HNO₂
- c) CH₃COOH
- d) HCN

Ans. d

Sol. Acidic strength ∝ Ka

The conjugate base of a weakest acid is strongest \therefore HCN $(K_a = 4 \times 10^{-10})$ (least value)



A, B and C respectively are

- a) ethanol, ethane nitrile and ethyne
- b) ethane nitrile, ethanol and ethyne
- c) ethyne, ethanol and ethane nitrile
- d) ethyne, ethane nitrile and ethanol

Ans. c

Sol.
$$A \rightarrow CH = CH \xrightarrow{\text{HgSO}_4} CH_2 = CH - OH \xrightarrow{\text{Tautamerises}} CH_3CHO$$

$$B \rightarrow CH_3CH_2OH \xrightarrow{\text{PCC}} CH_3CHO$$

$$C \rightarrow CH_3CN \xrightarrow{\text{(i)SnCl}_2/HCl} CH_3CHO$$

- 7. The reagent which can do the conversion $CH_3COOH \rightarrow CH_3 CH_2 OH$ is
 - a) LiAlH₄ / ether
- b) H₂, Pt
- c) NaBH₄
- d) Na and C₂H₅OH

Ans. a

Sol. $CH_3COOH \xrightarrow{LiAlH_4} CH_3CH_2OH$

$$8. \quad \text{CH}_3\text{CHO} \xrightarrow{\quad (i) \text{ CH}_3\text{MgBr} \\ \quad (ii) \text{ H}_3\text{O}^+} \rightarrow A \xrightarrow{\quad \text{Conc } \text{H}_2\text{SO}_4 \\ \quad \Delta \rightarrow B \xrightarrow{\quad (i) \text{ B}_2\text{H}_6 \\ \quad (ii) \text{ H}_2\text{O}, \text{ OH}^-} \rightarrow C$$

A and C are

- a) Identical
- b) Position isomers
- c) Functional
- d) Optical isomers

Ans. b

Sol.
$$\begin{array}{c} \text{CH}_{3} \\ \text{CH}_{3}\text{CHO} \xrightarrow{(i)\text{CH}_{3}\text{MgBr}} \\ \text{CH}_{3}\text{CH} - \text{OH} \xrightarrow{\text{conc.H}_{2}\text{SO}_{4}} \\ \text{A} \\ \text{CH}_{3} - \text{CH} = \text{CH}_{2} \xrightarrow{(i)\text{B}_{2}\text{H}_{6}} \\ \text{CH}_{3} - \text{CH}_{3}\text{CH}_{2}\text{CH}_{2}\text{OH} \\ \end{array}$$

- : A and C are position isomers.
- 9. Which of the following is not true for oxidation? a) addition of oxygen
 - b) addition of electronegative element
 - c) removal of hydrogen
 - d) removal of electronegative element

Ans. d

10. Which is the most suitable reagent for the following conversion?

$$\begin{array}{c} \text{O} \\ \text{CH}_3 - \text{CH} = \text{CH} - \text{CH}_2 - \overset{\mid 1}{\text{C}} - \text{CH}_3 \rightarrow \\ \text{O} \\ \text{CH}_3 - \text{CH} = \text{CH} - \text{CH}_2 - \overset{\mid 1}{\text{C}} - \text{OH} \end{array}$$

- a) Tollen's reagent
- b) Benzoyl peroxide
- c) I_2 and NaOH solution with subsequent acidification
- d) Sn and NaOH solution

Ans. c

Sol. Conceptual

- 11. $C_6H_5CH_2Cl \xrightarrow{\text{alc.NH}_3} A \xrightarrow{\text{2CH}_3Cl} B$. The product B is
 - a) N, N-Dimethyl phenyl methanamine
 - b) N, N-Dimethyl benzenamine
 - c) N-Benzyl-N-methyl methanamine
 - d) phynyl-N-N-dimethyl methanamine

Ans. a

Sol.
$$C_6H_5CH_2Cl \xrightarrow{alc.NH_3} C_6H_5CH_2NH_2 \xrightarrow{2CH_3Cl} C_6H_5CH_2N(CH_3)_2$$

- 12. The method by which aniline cannot be prepared is
 - a) Nitration of benzene followed by reduction with Sn and con. HCl
 - b) Degradation of benzamide with bromine in alkaline solution
 - c) Reduction of nitrobenzene with H_2/Pd is ethanol
 - d) Potassium salt of pthalimide treated with chlorobenzene followed by the hydrolysis with aqueous NaOH solution

Ans. d

Sol. Conceptual

- 13. Permanent hardness cannot be removed by
 - a) Using washing soda
 - b) Calgon's method
 - c) Clark's method
 - d) Ion exchange method

Ans. c

Sol. Conceptual

14. A hydrocarbon $A(C_4H_8)$ on reaction with HCl gives a compound $B(C_4H_9Cl)$ which on reaction with 1 mol of NH_3 gives compound $C(C_4H_{10}N)$. On reacting with $NaNO_2$ and HCl followed by treatment with water, compound C yields an optically active compound D. The is

Ans. b

Sol. Optically active compound is option B.

- 15. RNA and DNA are chiral molecules, their chirality is due to the presence of
 - a) D-sugar component
 - b) L-sugar component
 - c) Chiral bases
 - d) Chiral phosphate ester unit

Ans. a

Sol. Conceptual

- 16. The property of the alkaline earth metals that increases with their atomic number is
 - a) Ionisation enthalpy
 - b) Electronegativity
 - c) Solubility of their hydroxide in water
 - d) Solubility of their sulphate in water

Ans. c

Sol. Conceptual

- 17. Primary structure in a nucleic acid contains bases as GATGC ... The chain which is complementary to this chain is
 - a) G G T G A
- b) T G A A G
- c) C T A C G ...
- d) T T T A G

Ans. c

- In the detection of II group acid radical, the salt 18. containing chloride is treated with concentrated sulphuric acid, the colourless gas is liberated. The name of the gas is
 - a) Hydrogen chloride gas
 - b) Chlorine gas
 - c) Sulphur dioxide gas
 - d) Hydrogen gas

Ans. a

Sol. Conceptual

- The number of six membered and five membered rings in Buckminster Fullerence respectively is
- a) 20, 12 b) 12, 20 c) 14, 18 d) 14, 11

Ans. a

Sol. Conceptual

- 20. In chrysoberyl, a compound containing Beryllium, Aluminium and oxygen, oxide ions form cubic close packed structure. Aluminium ions occupy $\frac{1}{4}$ th of octahedral voids. The formula of the compound is
- Ans. b

Be Al O N = No. of oxide ions invoved in CCP

a) BeAlO₄ b) BeAl₂O₄ c) Be₂AlO₂ d) BeAlO₂

- **Sol.** $\frac{N}{4}: \frac{2N}{4}: N$: octahedral voids = N
 - 1: 2: 4 Tetrahedral voids = 2N
- 21. The correct statement regarding defects in solid is
 - a) Frenkel defect is a vacancy defect
 - b) Schottky defect is a dislocation defect
 - c) Trapping of an electron in the lattice leads to the formation of F-centre
 - d) Schottky defect has no effect on density

Ans. c

Sol. Frenkel defect – dislocation defect Schottky defect – decreases density F-centre - Trapping of on electrons in the lattices

- A metal crystallises in BCC lattice with unit 22. cell edge length of 300 pm and density 6.15gcm⁻³. The molar mass of the metal is
 - a) $50 \, \text{g mol}^{-1}$
- b) 60 g mol⁻¹
- c) $40 \,\mathrm{g} \,\mathrm{mol}^{-1}$
- d) 70 g mol⁻¹

Ans. a

So1. $d = \frac{ZM}{a^3N_A}$

$$M = \frac{d a^{3} N_{A}}{Z} = \frac{6.15 \times \left(300 \times 10^{-10}\right)^{3} \times 6 \times 10^{23}}{2}$$

$$\approx 50 \text{ g mol}^{-1}$$

- Henry's law constant for the solubility of N₂ 23. gas in water at 298K is 1.0×10^5 atm. The mole fraction of N_2 in air is 0.8 The number of moles of N₂ from air dissolved in 10 moles of water at 298K 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}

Ans. a

Sol. $P_{N_2} = X_{N_2} \cdot P_{total}$ $= 0.8 \times 5 = 4$ atm

$$P_{_{N_{_{2}}}}=K_{_{H}}\text{.}\,X_{_{N_{_{2}}}}$$

$$4 = 10^5 . X_N$$

$$X_{N_2} = 4 \times 10^{-5}$$

$$X_{_{N_{_{2}}}} = \frac{n_{_{N_{_{2}}}}}{n_{_{N_{_{2}}}} + n_{_{H_{_{2}}O}}} \ \left(n_{_{N_{_{2}}}} < < < < n_{_{H_{_{2}}O}}\right)$$

$$4 \times 10^{-5} = \frac{n_{N_2}}{10}$$

$$n_{N_2} = 4 \times 10^{-4}$$

- 24. A pure compound contains 2.4g of C, 1.2×10^{23} atoms of H, 0.2 moles of oxygen atoms. Its empirical formula is

 - a) C_0HO b) $C_0H_0O_0$ c) CH_0O
- d) CHO

Ans. d

Sol. 2.4g C = $\frac{2.4}{12}$ = 0.2 mol

$$1.2 \times 10^{23}$$
 atoms of $H = \frac{1.2 \times 10^{23}}{6 \times 10^{23}} = 0.2$ mol

0.2 mole of 'O' atoms

 \therefore simplest ratio = C : H : O

$$= CHO$$

- 25. Choose the correct statement
 - a) K_H value is same for a gas in any solution
 - b) Higher the K_H value more the solubility of
 - c) K_H value increases on increasing the temperature of the solution
 - d) Easily liquefiable gases usually has lesser K_H values

Ans. c

- K_H value changes with solvent nature Sol. Higher the K_H less is solubility K_H value increase with increase of T' Eerily liquefied gases have high K_H value
- 26. K_H value (K bar) of Argon Carbondioxide (II) formuldehyde (III) and methane (IV) are respectively 40.3, 167, 1.83×10⁻⁵ and 0.413 at 298 K. The increasing order of solubility of gas in liquid is
 - a) I < II < IV < III
- b) III < IV < II < I
- c) I < III < II < IV
- d) I < IV < II < III

Ans. a

Sol.
$$P_H = K_H \times$$

$$K_{\rm H} \propto \frac{1}{X(\text{solubility})}$$

- \therefore more is the K_H less is the solubility
- 27. The vapour pressure of pure liquids A and B are 450 and 700 mm of Hg at 350 K respectively. If the total vapour pressure of the mixture is 600 mm of Hg, the composition of the mixture in the solution is

a)
$$x_A = 0.4$$
, $x_B = 0.6$

b)
$$x_A = 0.6$$
, $x_B = 0.4$

c)
$$x_1 = 0.3$$
, $x_2 = 0.7$

c)
$$x_A = 0.3$$
, $x_B = 0.7$ d) $x_A = 0.7$, $x_B = 0.3$

Ans. a

Sol.
$$P_{total} = P_A^O X_A + P_B^O X_B$$

 $= P_A^O X_A + P_B^O (1 - X_A)$
 $600 = 450.X_A + 700(1 - X_A)$

$$X_A = 0.4$$

$$X_{\rm B} = 1 - 0.4 = 0.6$$

Consider the following electrodes

$$P = Zn^{2+} (0.0001M) / Zn Q = Zn^{2+} (0.1M) / Zn$$

$$R = Zn^{2+} (0.01M) / Zn$$

$$R = Zn^{2+} (0.01M) / Zn$$
 $S = Zn^{2+} (0.001M) / Zn$

 $E^{o}Zn/Zn^{2+} = -0.76V$ electrode potentials of the above electrodes in volts are in the order

a)
$$P > S > R > Q$$

b)
$$S > R > Q > P$$

c)
$$Q > R > S > P$$

d)
$$P > Q > R > S$$

Ans. c

Sol.
$$Zn^{2+}_{(aq)} + 2e^{-} \rightarrow Zn_{(s)}$$

$$E_{\mathrm{red}} = E_{\mathrm{red}}^{\Theta} - \frac{0.059}{n} log \frac{1}{\left \lceil Z n^{2^{+}} \right \rceil}$$

$$E_{\rm red} = -0.76 + \frac{0.059}{2} log \Big[Z n^{2^+} \Big]$$

$$as \Big[Zn^{^{2+}}\Big] \!\uparrow\! E_{red} \uparrow$$

The number of angular and radial nodes in 3p 29. orbital respectively are

c)
$$2,1$$

d) 2,3

Ans. b

- No. of angular nodes = l = 1 (3p) Sol. No. of radial nodes = n - l - 1 = 3-1-1=1
- 30. The resistance of 0.01 m KCl solution at 298 K is 1500 Ω . If the conductivity of 0.01 m KCl solution at 298 K is 0.1466×10^{-3} S cm⁻¹. The cell constant of the conductivity cell in cm⁻¹ is

Ans. a

Sol.
$$G^* = KR$$

$$= 0.146 \times 10^{-3} \times 1500$$
$$= 0.219$$

31. $H_{2_{(s)}} + 2AgCl_{(s)} \rightleftharpoons 2Ag_{(s)} + 2HCl_{(ag)}$

> E_{cell}^{o} at 25°C for the cell is 0.22 V. The equilibrium constant at 25°C is

a)
$$2.8 \times 10^7$$

b)
$$5.2 \times 10^8$$

c)
$$2.8 \times 10^5$$

d)
$$5.2 \times 10^4$$

Ans. a

Sol.
$$\log K_c = \frac{E_{cell}^{\Theta} \times n}{0.059} = \frac{0.22 \times 2}{0.059} = 7.45$$

$$K_c = Antilog (7.45) = 2.8 \times 10^7$$

- 32. For a reaction $A + 2B \rightarrow Products$, when concentration of B alone is increased half life remains the same. If concentration of A alone is doubled, rate remains the same. The unit of rate constant for the reaction is
 - a) S^{-1}
- b) L mol⁻¹ S⁻¹
- c) mol L⁻¹ S⁻¹
- d) atm⁻¹

Ans. a

- **Sol.** As ['B'] increase, $t_{\underline{1}}$ remains same
 - i.e. 1st order with respect to 'B'

$$rate = k[A]^0[B]^1$$

overall order = 1

 \therefore units of $k = S^{-1}$

- The third ionisation enthalpy is highest in 33.
 - a) Alkali metals
 - b) Alkaline earth metals
 - c) Chalcogens
 - d) Pnictogens

Ans. b

Sol. Conceptual

34. If the rate constant for a first order reaction is k, the time(t) required for the completion of 99% of the reaction is given by

a)
$$t = \frac{4.606}{k}$$

b)
$$t = \frac{2.303}{k}$$

c)
$$t = \frac{0.693}{k}$$

d)
$$t = \frac{6.909}{k}$$

Ans. a

- **Sol.** $t = \frac{2.303}{k} \log \frac{[R]_0}{[R]}$ $=\frac{2.303}{k}\log\left(\frac{100}{1}\right)$ $=\frac{4.606}{1}$
- The rate of a gaseous reaction is given by the 35. expression $k[A][B]^2$. If the volume of vessel is reduced to one half of the initial volume, the reaction rate as compared to original rate is
 - a) $\frac{1}{16}$ b) $\frac{1}{8}$ c) 8
- d) 16

Ans. c

Sol. rate =
$$K[A]^1[B]^2$$

$$= K \left[\frac{n}{v} \right]_{A}^{1} \left[\frac{n}{v} \right]_{B}^{2}$$

$$= K \left[\frac{n}{v} \right]^{1} \left[\frac{n}{v} \right]^{2}$$

$$= '8' K \left[\frac{n}{v} \right]^{1} \left[\frac{n}{v} \right]^{2}$$

'8'times increases

The correct IUPAC name of 36.



- a) 4-Ethyl-1-Fluoro-2-nitrobenzene
- b) 1-Ethyl-4-Fluoro-3-nitrobenzene
- c) 3-Ethyl-6-Fluoronitrobenzene
- d) 5-Ethyl-2-Fluoronitrobenzene

Ans. a

Sol. Conceptual

- 37. Higher order (>3) reactions are rare due to a) Shifting of equilibrium towards reactants
 - due to elastic collisions b) Loss of active species on collision
 - c) Low probability of simultaneous collision of all reacting species
 - d) Increase in entropy as more molecules are involved

Ans. c

Sol. Conceptual

- Arrange benzene, n-hexane and ethyne in 38. decreasing order of their acidic behaviour
 - a) Benzene > n-hexane > ethyne
 - b) n-hexane > Benzene > ethyne
 - c) ethyne > n-hexane > Benzene
 - d) ethyne > Benzene > n-hexane

Ans. d

- A colloidal solution is subjected to an electric 39. field than colloidal particles more towards anode. The amount of electrolytes BaCl₂, AlCl₃ and NaCl required to coagulate the given colloid is in the order
 - a) NaCl > BaCl₂ > AlCl₃
 - b) $BaCl_2 < AlCl_3 > NaCl$
 - c) AlCl₂ = NaCl = BaCl₂
 - d) $AlCl_3 > BaCl_2 > NaCl$

Ans. a

Sol. As ions are moving toward anode i.e. negatively charged colloid

> Coagulation value ∝ – cogulating power

- $\therefore \text{ Na}^{+1} > \text{Ba}^{+2} > \text{Al}^{+3}$
- 40. Which of the following is an incorrect statement?
 - a) Hydrogen bonding is stronger than dispersion forces
 - b) Sigma bonds are stronger than π -bonds
 - c) Ionic bonding is non-directional
 - d) σ -electrons are referred to as mobile electrons

Ans. d

Sol. Conceptual

- 41. Zeta potential is
 - Potential required bring about coagulation of a colloidal sol.
 - b) Potential required to give the particle a speed of 1 cm S^{-1}
 - c) Potential difference between fixed charged layer and the diffused layer having opposite charges
 - d) Potential energy of the colloidal particles.

Ans. c

Sol. Conceptual

- 42. Which of the following compound on heating gives N₂O?
 - a) $Pb(NO_3)_{0}$
- b) NH₄NO₃
- c) NH₄NO₂
- d) NaNO₃

Ans. b

Sol. Conceptual

Which of the following property is true for the given sequence

 $NH_3 > PH_3 > AsH_3 > SbH_3 > BiH_3$?

- a) Reducing property b) Thermal stability
- c) Bond angle
- d) Acidic character

Ans. b

Sol. Conceptual

- 44. The correct order of boiling point in the following compounds is
 - a) $HF > H_2O > NH_3$
- b) $H_2O > HF > NH_3$
- c) $NH_3 > H_2O > HF$ d) $NH_3 > HF > H_2O$

Ans. b

Sol. Conceptual

45. XeF₆ on partial hydrolysis gives a compound X, which has square pyramidal geometry 'X' is a) XeO₃ b) XeO₄ c) XeOF₄ d) XeO₂F₂

Ans. c

Sol.
$$XeF_6 + H_2O \rightarrow XeOF_4 + 2HF(XeOF_4 - square pyramidal)$$

$$XeF_6 + 2H_2O \rightarrow XeO_2F_2 + 4HF(XeO_2F_2 - See - saw)$$

- 46. A colourless, neutral, paramagnetic oxide of Nitrogen 'P' on oxidation gives reddish brown gas Q. Q on cooling gives colourless gas R. R on reaction with P gives blue solid S. Identify P, Q, R, S respectively

 - a) N_2O NO $NO_2N_2O_5$ b) N_2O NO₂ $N_2O_4N_2O_3$
 - c) NO NO₂ $N_2O_4N_2O_3$ d) NO NO $N_2O_4N_2O_5$

Ans. c

$$2NO + O_2 \rightarrow 2NO_2 \rightarrow N_2O_4 \xrightarrow{2NO} 2N_2O_3$$
P Q R S

Sol.

- 47. Which of the following does not represent property stated against it?
 - a) $CO^{+2} < Fe^{+2} < Mn^{+2}$ Ionic size
 - b) Ti < V < Mn Number of oxidation states
 - c) $Cr^{+2} < Mn^{+2} < Fe^{+2}$ Paramagnetic behaviour
 - d) Sc > Cr > Fe Density

Ans. c

- 48. Which one of the following is correct for all elements from Sc to Cu?
 - a) The lowest oxidation state shown by them is
 - b) 4S orbital is completely filled in the ground state
 - c) 3d orbital is not completely filled in the ground state
 - d) The ions in +2 oxidation states are paramagnetic

Ans. d

Sol. Conceptual

- 49. When the absolute temperature of ideal gas is doubled and pressure is halved, the volume of gas
 - a) will be half of original volume
 - b) will be 4 times the original volume
 - c) will be 2 times the original volume
 - d) will be 1/4th times the original volume

Ans. b

Sol. PV = nRT

$$\frac{P_1V_1}{T_1} = \frac{P_2V_2}{T_2}$$
 $\frac{P \times V_1}{T} = \frac{P}{2} \times \frac{V_2}{2T}$ $V_2 = 4V_1$

50. Which of the following pairs has both the ions coloured in aqueous solution? [Atomic numbers of

$$[Sc = 21, Ti = 22, Ni = 28, Cu = 29, Mn = 25]$$

- a) Sc^{3+} . Mn^{2+}
- b) Ni²⁺, Ti⁴⁺
- c) Ti³⁺, Cu⁺
- d) Mn²⁺, Ti³⁺

Ans. d

Sol. Conceptual

- 51. For the crystal field splitting in octahedral complexes,
 - a) the energy of the e_g orbitals will decrease by
 - $(3/5)\Delta_0$ and that of the t_{2g} will increase by
 - $(2/5)\Delta_0$
 - b) the energy of the e, orbitals will increase by
 - $(3\,/\,5)\Delta_0$ and that of the $\,t_{2g}\,$ will decrease by
 - $(2/5)\Delta_0$
 - c) the energy of the e_g orbitals will increase by
 - $(3/5)\Delta_0$ and that of the t_{2g} will increase by
 - $(2/5)\Delta_0$
 - d) the energy of the e, orbitals will decrease by
 - $(3/5)\Delta_0$ and that of the t_{2g} will decrease by $(2/5)\Delta_0$

Ans. b

Sol. Conceptual

- 52. Peroxide effect is observed with the addition of HBr but not with the addition of HI to unsymmetrical alkene because
 - a) H-I bond is stronger that H-Br and is not cleaved by the free radical
 - b) H-I bond is weaker than H-Br bond so that iodine free radicals combine to form iodine molecules
 - c) Bond strength of HI and HBr are same but free radicals are formed in HBr
 - d) All of these

Ans. b

Sol. Conceptual

- 53. The IUPAC name of $\left[Co(NH_3)_5(CO_3)\right]Cl$ is
 - a) Pentaamminecarbonatocobalt (III) Chloride
 - b) Carbonatopentamminecobalt (III) Chloride
 - c) Pentaamminecarbonatocobaltate (III) Chloride
 - d) Pentaammine cobalt (III) Carbonate Chloride

Ans. a

Sol. Conceptual

- 54. Homoleptic complexes among the following are
 - A) $K_3[Al(C_2O_4)_3]$,
- B) $\left[\text{CoCl}_2 \left(\text{en} \right)_2 \right]^+$
- C) $K_2 \lceil Zn(OH)_4 \rceil$
- a) A only
- b) A and B only
- c) A and C only
- d) C only

Ans. c

Sol. Conceptual

55. The correct order for wavelengths of light absorbed in the complex ions $\left[\text{CoCl}\left(\text{NH}_{3}\right)_{5}\right]^{2^{+}}, \left[\text{Co}\left(\text{NH}_{3}\right)_{6}\right]^{3^{+}} \text{ and } \left[\text{Co}\left(\text{CN}\right)_{6}\right]^{3^{-}}$

$$a) \Big[CoCl \big(NH_3\big)_5 \Big]^{2+} > \Big[Co \big(NH_3\big)_6 \Big]^{3+} > \Big[Co \big(CN\big)_6 \Big]^{3-}$$

b)
$$\left[\text{Co}\left(\text{NH}_{3}\right)_{6}\right]^{3+} > \left[\text{Co}\left(\text{CN}\right)_{6}\right]^{3-} > \left[\text{CoCl}\left(\text{NH}_{3}\right)_{5}\right]^{2+}$$

c)
$$\left[\text{Co(CN)}_{6} \right]^{3-} > \left[\text{CoCl(NH}_{3})_{5} \right]^{2+} > \left[\text{Co(CN)}_{6} \right]^{3-}$$

d)
$$\left[\text{Co(NH}_3)_6 \right]^{3+} > \left[\text{CoCl(NH}_3)_5 \right] > \left[\text{Co(CN)}_6 \right]^{3-}$$

Ans. a

Sol. Wave length of light absorbed is inversely proportional to strength of the ligand.

$$\begin{array}{c}
\text{CH}_2\text{-CH}_3\\
&\xrightarrow{\text{Br}_2}\\
\text{UV light}
\end{array}$$
A

The compound A (major product) is

c)
$$R_2$$
-CH₃ R_3 R_4 -CH₂-CH₃ R_4

Ans. b

Sol. Free radical substitution of alkane part. (Benzyl free radical) takes place.

57. Bond enthalpies of A_2 , B_2 and AB are in the ratio 2:1:2. If bond enthalpy of formation of AB is -100KJ mol $^{-1}$. The bond enthalpy of B_2

a) 100 KJ mol⁻¹

b) 50 KJ mol⁻¹

c) 200 KJ mol⁻¹

d) 150 KJ mol⁻¹

Ans. c

$$\begin{split} \textbf{Sol.} & \text{ Assume bond strength of } A_2 = 2 X \text{ , then,} \\ & B_2 = X \text{ , } AB = 2 X \\ & \Delta_r H^\Theta = \sum \Delta_{\text{diss}} H^\Theta_{\ (R)} - \sum \Delta_{\text{diss}} H^\Theta_{\ (P)} \\ & \frac{1}{2} A_2 + \frac{1}{2} B_2 \rightarrow AB \\ & -100 = \frac{1}{2} 2 X + \frac{1}{2} X - 2 \\ & X = 200 \end{split}$$

58. The order of reactivity of the compounds $C_6H_5CH_2Br$, $C_6H_5CH(C_6H_5)Br$, $C_6H_5CH(CH_3)Br$ and $C_6H_5C(CH_3)(C_6H_5)Br$ in S_N^2 reaction is

a)
$$C_6H_5 - C_-Br < C_6H_5 - C_-Br < C_6H_5 - C_-Br < C_6H_5 - C_-Br < C_6H_5 - C_-Br$$

Ans. a

Sol. Conceptual

59. The major product of the following reaction is $CH_2 = CH - CH_2 - OH \xrightarrow{HBr} Product$

b)
$$CH_2 = CH - CH_2Br$$

Ans. a

Sol. $\begin{array}{ll} \text{CH}_2 = \text{CH} - \text{CH}_2 \text{OH} & \xrightarrow{\text{HBr}} \text{CH}_3 \text{CH} \big(\text{Br} \big) - \\ & \text{CH}_2 \text{OH} & \xrightarrow{\text{HBr}} \text{CH}_3 \text{CH} \big(\text{Br} \big) - \text{CH}_2 \text{Br} \end{array}$

60.
$$\begin{array}{c} \text{CH}_3 \\ \text{CH-CH}_3 \\ \text{CH}_3\text{-C-O-OH} \\ \\ \text{H}_2\text{O} \rightarrow \text{A+B} \end{array}$$

The product 'A' gives white precipitate when treated with bromine water. The product 'B' is treated with Barium hydroxide to give the product C. The compound C is heated strongly to form product D. The product D is

to form product D. The product L

a) 4-Methylpent-3-en-2-one

b) But-2 enal

c) 3-Methylpent-3-en-2-one

d) 2-Methylbut-2-enal

Ans. a

Sol. A is phenol

B is acetone

Acetone when treated with Ba(OH)₂ Undergoes aldol condensation and followed by heating gives 4-methylpent -3-en-2-one