Solutions to IITJEE-2004 Mains Paper Chemistry

Time: 2 hours

Note: Question number 1 to 10 carries 2 marks each and 11 to 20 carries 4 marks each.

1. For the given reaction

 $A + B \longrightarrow Products$ Following data were given Initial conc. (m/L)

Initial conc. (m/L).	Initial conc. (m/L)	Initial rate [mL ⁻¹ s ⁻¹]
[A]	[B]	
0.1	0.1	0.05
0.2	0.1	0.1
0.1	0.2	0.05

- a) Write the rate equation.
- b) Calculate the rate constant.

Sol. a) Let the order w.r.t A & B are x any y respectively

$$r = K[A]^{x} [B]^{y}$$

$$0.05 = K[0.1]^{x} [0.1]^{y}$$

$$0.1 = K[0.2]^{x} [0.1]^{y}$$
or $2 = [2]^{x}$

$$x = 1$$

$$0.05 = K[0.1]^{x} [0.1]^{y}$$

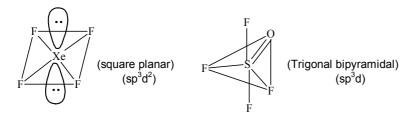
$$0.05 = K[0.1]^{x} [0.2]^{y}$$

$$1 = [2]^{y}$$

$$y = 0$$

- b) rate equation = r = K[A] [B]⁰ 0.1 = K[0.2] K = 0.5 Sec⁻¹
- 2. 100 ml of a liquid contained in an isolated container at a pressure of 1 bar. The pressure is steeply increased to 100 bar. The volume of the liquid is decreased by 1 ml at this constant pressure. Find the ΔH & ΔU .
- Sol. ΔH =0, $\Delta q_p = \Delta U$ -W W = PdV = 100×1 atmmL = 10⁻² KJ = ΔU
- 3. Draw the shape of XeF₄ and OSF₄according to VSEPR theory. Show the lone pair of electrons on the central atom

Sol.



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4. The structure of D-Glucose is as follows

-OH

-ОН

b)

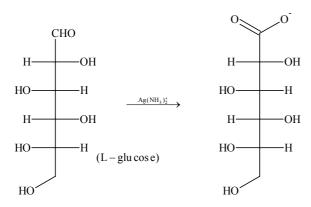
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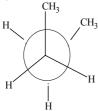
НО

- a) Draw the structure of L Glucose.
 - Give the reaction of L Glucose with Tollens reagent.

Sol.



- 5. a) Draw New mann's projection for the less stable staggered form of butane.
 - b) Relatively less stability of the staggered form is due to
 - i) Torsional strain.
 - ii) Vander Waal's strain.
 - iii) Combination of the above two.
- Sol. a



- b) Less stability is due to Vander Waal's strain
- 6. Arrange the following oxides in the increasing order of Bronsted basicity.

Sol.
$$Cl_2O_7 < SO_3 < CO_2 < B_2O_3 < BaO$$

7. AlF₃ is insoluble in anhydrous HF but when little KF is added to the compound it becomes soluble. On addition of BF₃,AlF₃ is precipitated. Write the balanced chemical equations.

Sol.
$$3KF + AlF_3 \longrightarrow K_3AlF_6$$

 $K_3AlF_6 + 3BF_3 \longrightarrow AlF_3 + 3KBF_4$

- 8. The crystal AB (rock salt structure) has molecular weight 6.023 y amu. where y is an arbitrary number in amu. If the minimum distance between cation & anion is $y^{1/3}$ nm and the observed density is $20 \, \text{Kg/m}^3$. Find the
 - a) density in Kg/m³ and
 - b) type of defect

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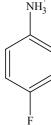
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Sol. a) Density =
$$\frac{4 \times 6.023 \times y}{6.023 \times 10^{23} \times 8 \times y \times 10^{-27}}$$
 [Since a = 2y^{1/3}]
= 5×10^3 g/m³
= 5 Kg/m³

- Since the (density) calculated < density observed, it means the defect is metal excess defect. b)
- 9. Which of the following is more acidic and why?

$$\begin{array}{c|c} NH_3^+ & NH_3^+ \\ \hline \\ F & \end{array}$$

Sol. is more acidic due to - inductive effect of fluorine NH₂



10. 7-bromo-1,3,5-cycloheptatriene exists as ionic species in aqueous solution while 5-bromo-1,3 cyclopentadiene doesn't ionise even in presence of Ag+(aq), Explain.

Sol. Aromatic Ion (highly stable)

The schrodinger wave equation for hydrogen atoms is 11.

$$\Psi_{2s} = \frac{1}{4(2\pi)^{1/2}} \left(\frac{1}{a_0}\right)^{3/2} \left(2 - \frac{r}{a_0}\right) e^{-r/a}$$

- Where a_0 is Bohr's radius. Let the radial node in 2s be at r_0 . Then find r in terms of a_0 . b) A base ball having mass 100 g moves with velocity 100 m/sec. Find out the value of wave length of
- c) $_{92}X^{234} \xrightarrow{-7\alpha \\ -6\beta} Y$. Find out atomic number, mass number of Y and identify it.

a) ψ_{2s}^2 = probability of finding electrons at any place Sol. ∴ $\Psi^2 = 0$ at node

$$\therefore \Psi^2 = 0 = \frac{1}{4} \frac{1}{\sqrt{2\pi}} \left(\frac{1}{a^0}\right)^3 \left(2 - \frac{r}{a_0}\right)^2 \times e^{-r/a_0}$$
$$\left(2 - \frac{r}{a_0}\right) = 0 \Rightarrow 2 = \frac{r}{a_0} \Rightarrow 2a_0 = r$$

b)
$$\lambda = \frac{h}{mv} = \frac{6.626 \times 10^{-34}}{100 \times 10^{-3} \times 100}$$

 $\lambda = 6.626 \times 10^{-35} \text{ m} = 6.626 \times 10^{-25} \text{ A}^{\circ}$
c) Yis $_{84}\text{Po}^{206}$

- 12. On the basis of ground state electronic configuration arrange the following molecules in increasing O-O bond length order. KO_2 O_2 $O_2[AsF_6]$

Sol.
$$O_2 = \sigma ls^2, \sigma^* ls^2, \sigma 2s^2, \sigma^* 2s^2, \sigma 2p_x^2 \begin{cases} \pi 2p_y^2 \\ \pi 2p_z^2 \end{cases} \begin{cases} \pi^* 2p_y^1 \\ \pi^* 2p_z^1 \end{cases}$$

bond order =
$$\frac{10-6}{2}$$
 = 2

$$O_{2}^{-} = \sigma_{ls^{2}}, \sigma^{*}_{ls^{2}}, \sigma_{2s^{2}}, \sigma^{*}_{2s^{2}}, \sigma_{2p_{x}^{2}} \left\{ \frac{\pi^{2}P_{y}^{2}}{\pi^{2}P_{z}^{2}} \right\} \left\{ \frac{\pi^{*}2P_{y}^{2}}{\pi^{*}2P_{z}^{1}} \right\} in \left[KO_{2} \right]$$

bond order =
$$\frac{10-7}{2} = \frac{3}{2}$$

$$O_{2}^{+} \qquad \qquad = \sigma_{ls^{2}} \,, \sigma^{*}_{\ ls^{2}} \,, \sigma_{2s^{2}} \,, \sigma^{*}_{\ 2s^{2}} \,, \sigma_{2P_{x}^{2}} \left\{ \begin{smallmatrix} \pi^{2}P_{y}^{2} \\ \pi^{2}Pz^{2} \end{smallmatrix} \right\} \left\{ \pi^{*}_{\ } 2p_{y}^{\ l} \right\}$$

in
$$[O_2(AsF_6)]$$

bond order
$$\frac{10-5}{2} = \frac{5}{2}$$

Bond length order is $O_2^+ < O_2 < O_2^-$

13.

In the following equilibrium
$$N_2O_4(g)$$
 $2NO_2(g)$

when 5 moles of each are taken, the temperature is kept at 298 K the total pressure was found to be 20 bar. Given that

$$\Delta G_f^0(N_2O_4) = 100KJ$$

$$\Delta G_f^0(NO_2) = 50KJ$$

- i) Find ΔG of the reaction
- ii) The direction of the reaction in which the equilibrium shifts
- A graph is plotted for a real gas which follows Vander Waal's equation with PV_m taken on Y axis & P on X axis. Find the intercept of the line where V_m is molar volume
- $N_2O_4(g) = 2NO_2(g)$ Sol.

Reaction quotient =
$$\frac{P_{NO_2}^2}{P_{N,O_4}} = \frac{100}{10} = 10$$
 atm

$$\Delta G^{\circ}$$
 reaction = $2\Delta G_{\rm f}^{\circ} (NO_2) - \Delta G_{\rm f}^{\circ} (N_2O_4)$

$$0 = 100 - 100$$

$$\Delta G = \Delta G^{\circ} + RT \ln k$$

$$\therefore \Delta G = RT \ln Q$$

 $= 2.303 \times .082 \times 298 \times \log 9.9 = 56.0304$ Lit atm. = Positive

ii) Therefore reaction will shift towards backward direction.

b)
$$\therefore \left(P + \frac{a}{v_m^2}\right)(v_m - b) = RT$$

$$\left(P + \frac{aP^2}{(PV)^2}\right)\left(\frac{PV}{P} - b\right) = RT$$

$$[PV)^2 P + aP^2][(PV) - b)] = P(PV)^2 RT$$

$$\Rightarrow P[(PV)^2 + aP] (PV - bP) = P(PV)^2 RT$$

$$Put P = 0$$

$$\Rightarrow (PV)^3 = (PV)^2 RT$$

$$Intercept = RT$$

14. 1.22 g C_6H_5 COOH is added into two solvent and data of ΔT_b and K_b are given as:-

i) In 100 g CH₃COCH₃

 $\Delta T_{b} = 0.17$

 $K_b = 1.7 \text{ Kg Kelvin/mol}$ In 100 g benzene, $\Delta T_b = 0.13$ and $K_b = 2.6 \text{ Kg Kelvin/mol}$ Find out the molecular weight of C_0H_5COOH in both the cases and interpret the result.

0.1 M of HA is titrated with 0.1 M NaOH, calculate the pH at end point. Given Ka(HA)= 5×10^{-6} and α << 1

In first case Sol. a)

 $\Delta T_b = K_b \times m$

$$0.17 = 1.7 \times \frac{1.22}{M \times 100 \times 10^{-3}} \Rightarrow M = 122$$

ii) In second case

$$\Delta T_b = K_b \times m$$

$$0.13 = 2.6 \times \frac{1.22}{\text{M}' \times 100 \times 10^{-3}}$$

$$M' = 244$$

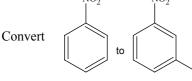
Benzoic acid dimerises in benzene

b) Since at end point molarity of salt = $=\frac{0.1}{2}$ M

: pH of salt of weak acid and strong base

$$pH = \frac{\left(pK_w + pK_a + log_c\right)}{2} = \frac{1}{2} \left[14 + 5.3010 + [-1.3010] \Rightarrow pH = 9.$$

15.



in not more than four steps. Also mention the temp and reaction

Sol.

$$\begin{array}{c|c} NO_2 & NO_2 & NO_2 \\ \hline & NO_2 & NO_2 \\$$

16.

$$\xrightarrow{\text{KCN}} (A) \xrightarrow{C_2 \text{H}_5 \text{ONa}/C_3 \text{H}_5 \text{OH}} (B) \xrightarrow{\text{H}_3 \text{O}^*} (C) \xrightarrow{\text{SOCl}_2} (D)$$

Identify A to D.

Sol.

17. A₁ & A₂ are two ores of metal M. A₁ on calcination gives black precipitate, CO₂ & water.

Calcination

Black solid +
$$CO_2$$
 + H_2O
 $A_1 \xrightarrow{dil HCI} I_2 + ppt$
 $A_2 \xrightarrow{roasting} Metal + gas$
 $K_2Cr_2O_7 + H_2SO_4$

green colour

Sol. $A_1 = Cu(OH)_2 CuCO_3$

$$A_2 = Cu_2S$$

$$Cu(OH)_2 CuCO_3 \xrightarrow{Calcination} 2CuO + CO_2 + H_2O$$
(A1) (Black Solid)

$$Cu(OH)_2CuCO_3 \xrightarrow{\text{dil}HCl} CuCl_2 + CO_2 + 3H_2O$$

$$2CuCl_2 + 4KI \rightarrow Cu_2l_2 + I_2 + 4KCl$$

$$2Cu2S + 3O2 \rightarrow 2Cu2O + 2SO2$$
(A₂)

$$Cu_2S + 2Cu_2O \rightarrow 6Cu + SO_2$$

- 18. NiCl₂ in the presence of dimethyl glyoxime (DMG) gives a complex which precipitates in the presence of NH₄OH, giving a bright red colour.
 - a) Draw its structure & show H-bonding
 - b) Give oxidation state of Ni & its hybridisation.
 - c) Predict wether it is paramagnetic or diamagnetic.

Sol.

Oxidation state of nickel is +2 and hybridization is dsp²

$$\mu s = \sqrt{n(n+2)} B.M$$

$$n = 0$$

$$\therefore \mu s = 0$$

19.

Find the equilibrium constant for the reaction
$$Cu^{+2} + In^{+2} \qquad Cu^{+} + In^{+3}$$
 Given that
$$E_{Cu^{+2}}^{\circ} /_{Cu^{+}} = 0.15V$$

$$E_{In^{+2}}^{\circ} /_{In^{+}} = -0.4V$$

$$E_{In^{+3}}^{\circ} /_{In^{+}} = -0.42 \text{ V}$$

Sol.
$$Cu^{+2} + e^{-} \longrightarrow Cu^{+}$$
 $\Delta G_{1}^{0} = -0.15 \text{ F}$
 $In^{+2} + e^{-} \longrightarrow In^{+} +$ $\Delta G_{2}^{0} = +0.4 \text{ F}$
 $In^{+} \longrightarrow In^{+3} + 2 e^{-}$ $\Delta G_{3}^{0} = -0.84 \text{ F}$
 $Cu^{+2} + In^{+2} \longrightarrow Cu^{+} + In^{+3}$ $\Delta G^{0} = -0.59 \text{ F}$

An organic compound 'P' having the molecular formula $C_5H_{10}O$ treated with dil H_2SO_4 gives two compounds, Q & R both gives positive iodoform test. The reaction of $C_5H_{10}O$ with dil H_2SO_4 gives reaction 10^{15} times faster then ethylene. Identify organic compound of Q & R. Give the reason for the extra 20. stability of P.

Sol. Highly stable carbocation

P is stabilized by resonance