FIITJEE 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	For the given reaction			
	A -	$A + B \longrightarrow Products$			
	Fo	Following data were given			
	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 a $r = K[A]^{x} [B]^{y}$ $0.05 = K[0.1]^{x} [0.1]^{y}$ $0.1 = K[0.2]^{x} [0.1]^{y}$ 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		re x any y respectively		
	b)	rate equation = $r = K[A] [B]$	$]^0$		
		0.1 = K[0.2]			
		$K = 0.5 \text{ Sec}^{-1}$			

- 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 $\Delta H \& \Delta U$.
- Sol. $\Delta H = 0$, $\Delta q_p = \Delta U W$ W = P dV $= 100 \times 1 \text{ atmmL}$ $= 10^{-2} \text{ KJ} = \Delta U$

Sol.

3. Draw the shape of XeF_4 and OSF_4 according to VSEPR theory. Show the lone pair of electrons on the central atom



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Sol.

4.



- 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.



a)



- b) Less stability is due to Vander Waal's strain
- Arrange the following oxides in the increasing order of Bronsted basicity. Cl₂O₇, BaO, SO₃, CO₂, B₂O₃
 Cl₂O₇, BaO, SO₃, CO₂, B₂O₃
- $\textbf{Sol.} \qquad Cl_2O_7 < SO_3 < CO_2 < B_2O_3 < BaO$
- 7. AlF_3 is insoluble in anhydrous HF but when little KF is added to the compound it becomes soluble. On addition of BF_3 , AlF_3 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 Kg/m^3 . Find the
 - a) density in Kg/m^3 and
 - b) type of defect

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

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





NH₃⁺

is more acidic due to - inductive effect of fluorine

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.





11. The schrodinger wave equation for hydrogen atoms is a)

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

- 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 . A base ball having mass 100 g moves with velocity 100 m/sec. Find out the value of wave length of b) base ball.
- $_{92}X^{234} \xrightarrow{-7\alpha}_{-6\beta} Y$. Find out atomic number, mass number of Y and identify it. c)

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

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

$$\left(2 - \frac{r}{a_{0}}\right) = 0 \Longrightarrow 2 = \frac{r}{a_{0}} \Longrightarrow 2a_{0} = r$$
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 $= \frac{10-6}{2} = 2$

$$O_{2}^{-} = \sigma_{ls^{2}}, \sigma^{*} ls^{2}, \sigma_{2s^{2}}, \sigma^{*} 2s^{2}, \sigma_{2P_{x}^{2}} \begin{cases} \pi 2P_{y}^{2} \\ \pi^{2} 2P_{z}^{2} \end{cases} \begin{cases} \pi^{*} 2P_{y}^{2} \\ \pi^{*} 2P_{z}^{1} \end{cases}$$
in [KO₂]
bond order $= \frac{10-7}{2} = \frac{3}{2}$

 $O_{2}^{+} = \sigma_{1s^{2}}, \sigma_{1s^{2}}^{*}, \sigma_{2s^{2}}, \sigma_{2s^{2}}^{*}, \sigma_{2P_{x}^{2}}^{2} \left\{ \pi^{2P_{y}^{2}}_{\pi^{2P_{z}^{2}}} \right\} \left\{ \pi^{*}2p_{y}^{-1} \right\}$

in $[O_2(AsF_6)]$

bond order $\frac{10-5}{2} = \frac{5}{2}$ Bond length order is $O_2^+ < O_2 < O_2^-$

a) In the following equilibrium $N_2O_4(g) = 2NO_2(g)$ 13.

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

 $\Delta G_{f}^{0}(N_{2}O_{4}) = 100 \text{KJ}$ $\Delta G_{f}^{0}(NO_{2}) = 50KJ$

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 b)

a)

i)
$$N_2O_4(g) = 2NO_2(g)$$

Reaction quotient =
$$\frac{P_{NO_2}^2}{P_{N_2O_4}} = \frac{100}{10} = 10$$
 atm
 $\Delta G^\circ \text{ reaction} = 2\Delta G_f^\circ (NO_2) - \Delta G_f^\circ (N_2O_4)$
 $0 = 100 - 100$
 $\Delta G = \Delta G^\circ + \text{RT lnk}$
 $\therefore \Delta G = \text{RT ln Q}$
 $= 2.303 \times .082 \times 298 \times \log 9.9 = 56.0304$ Lit atm. = Positive

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ii) Therefore reaction will shift towards backward direction.

b)
$$\therefore \left(P + \frac{a}{v_m^2} \right) (v_m - b) = RT$$
$$\left(P + \frac{aP^2}{(PV)^2} \right) \left(\frac{PV}{P} - b \right) = RT$$
$$[PV)^2 P + aP^2][(PV) - b)] = P(PV)^2RT$$
$$\Rightarrow P[(PV)^2 + aP] (PV - bP) = P(PV)^2RT$$
Put P = 0
$$\Rightarrow (PV)^3 = (PV)^2 RT$$
Intercept = RT

14. 1.22 g C₆H₅ COOH is added into two solvent and data of ΔT_b and K_b are given as:a) In 100 g CH₃COCH₃ i)

$$\Delta T_b = 0.17$$

- In 100 g benzene, $\Delta T_b = 0.17$ Kg Kelvin/mol Find out the molecular weight of C₆H₅COOH 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 b) << 1

 $\frac{1}{2} [14 + 5.3010 + [-1.3010] \Rightarrow pH = 9.$

Sol. a) In first case i)

ii)

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

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

ii) In second case $\Delta T_b = K_b \times m$ 1.22 $0.13 = 2.6 \times$ $M' \times 100 \times 10^{-3}$ 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 $+ pK_a + \log_c$)

 NO_2

ЮH

NH₄HS

NO₂

2 NO_2

to

NO₂

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





Convert

NO₂



16.



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17.

Sol.

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

$$A_1 \xrightarrow{\text{Calcinguon}} H_{cl}$$

$$A_2 \xrightarrow{\text{roasting}} \text{Metal} + g$$

$$\xrightarrow{\text{roasting}} \text{Metal} + \text{gas}$$

$$\downarrow K_2 \text{Cr}_2 \text{O}_7 + \text{H}_2 \text{SO}_4$$

green colour

$$\begin{aligned} \text{Sol.} \qquad & A_1 = \text{Cu}(\text{OH})_2 \text{CuCO}_3 \\ & A_2 = \text{Cu}_2 \text{S} \\ & \text{Cu}(\text{OH})_2 \text{CuCO}_3 \xrightarrow{\text{Calcination}} 2\text{CuO} + \text{CO}_2 + \text{H}_2 \text{O} \\ & \text{(Black Solid)} \end{aligned} \\ & \text{Cu}(\text{OH})_2 \text{CuCO}_3 \xrightarrow{\text{dilHCl}} \text{CuCl}_2 + \text{CO}_2 + 3\text{H}_2 \text{O} \\ & 2\text{CuCl}_2 + 4\text{KI} \rightarrow \text{Cu}_2 \text{I}_2 + \text{I}_2 + 4\text{KCl} \\ & 2\text{Cu}_2 \text{S} + 3\text{O}_2 \rightarrow 2\text{Cu}_2 \text{O} + 2\text{SO}_2 \\ & \text{(A}_2) \\ & \text{Cu}_2 \text{S} + 2\text{Cu}_2 \text{O} \rightarrow 6\text{Cu} + \text{SO}_2 \end{aligned}$$

- 18. NiCl₂ in the presence of dimethyl glyoxime (DMG) gives a complex which precipitates in the presence of NH_4OH , 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.



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19. Find the equilibrium constant for the reaction $Cu^{+2} + In^{+2} = Cu^{+} + In^{+3}$ Given that

$$E^{\circ}_{Cu^{+2}} /_{Cu^{+}} = 0.15V$$

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

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

- Sol. $Cu^{+2} + e^{-} \longrightarrow Cu^{+}$ $\Delta G_{1}^{0} = -0.15 \text{ F}$ $In^{+2} + e^{-} \longrightarrow In^{+} + \Delta G_{2}^{0} = +0.4 \text{ F}$ $In^{+} \longrightarrow In^{+3} + 2 e^{-} \qquad \Delta G_{3}^{0} = -0.84 \text{ F}$ $Cu^{+2} + In^{+2} \longrightarrow Cu^{+} + In^{+3} \quad \Delta G^{0} = -0.59 \text{ F}$ $-nFE^{\circ} = -0.59F$ $-E_{cell}^{0}F = -0.59F$ $E_{Cell}^{0} = 0.59$ $E_{cell} = E^{\circ} - \frac{0.0591}{n} \log K_{c}$ $0.59 = \frac{0.0591}{1} \log Kc$ $K_{c} = 10^{10}$
- 20. 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 stability of P.



P is stabilized by resonance

Sol.

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