Version No.			
0	0	0	0
1	1	1	1
2	2	2	2
3	3	3	3
4	4	4	4
5	5	5	5
6	6	6	6
(7)	7	7	7
8	8	8	8
9	9	9	9

CHEMISTRY HSSC–I SECTION – A (Marks 17) Time allowed: 25 Minutes

Section – A is compulsory. All parts of this section are to be answered on this page and handed over to the Centre Superintendent. Deleting/overwriting is not allowed. **Do not use lead pencil.**

Q.1 Fill the relevant bubble for each part. Each part carries one mark.

1.	Plasma is the mixture	of:				
	A. Electrons and p	orotons only	ν.	\bigcirc		
	B. Electrons and r	ositive ions	5.	Ŏ		
	C. Electrons and b	eta two nar	ticles.	ŏ		
	D Neutrons and n	rotons		ŏ		
	D. routons and p	rotons.		\bigcirc		
2	The electrode potential	of metals a	are.			
2.	$M\sigma^{2+} + 2e^{-}$	→ Mo	E°	= -2.71 v		
	$A\sigma \longrightarrow A\sigma^+$	$+ 1e^{-116}$	E	P = -0.8 v		
	Cell potential (emf) of	the cell for	$\frac{1}{1}$ med by t	these two will be:		
	$\frac{4}{4} + 351 \text{ y}$		B	-3 51 v	\bigcirc	
	$\frac{11.}{C} + 1.91 \text{ v}$	ŏ	D.	-1.91 v	X	
	C. + 1.91 V	0	D.	1.71 V	\bigcirc	
3	At constant Pressure w	hat will be	the chan	oe in temperature	when the volu	me of
5.	a gas will become twice	e of what it	the enant 1°	רקי און	when the volu	
	$A = 546^{\circ}C$		R	200°C	\cap	
	C = 546 K	X	D. D	200 C 273 K	X	
	С. Это К	\cup	D.	275 K	\cup	
4	Rate equation for a rea	ction 2A—		oduct is $Rate = K$	$[A]^2$ Unit of sr	ecific
т.	rate constant for this re	eaction is	• pro	Sauer 15 Rate IX		
	$\Delta = \text{mol}^2 \text{dm}^{-6} \text{S}^{-1}$	$\bigcap^{\text{action 13.}}$	B	$mol^{-1}dm^{3}S^{-1}$	\cap	
	$f_{\rm c}$ moldm ⁻³	X	D.	S ⁻¹	X	
	C. mordin	\bigcirc	D.	5	\cup	
5	A substance which itse	lf is not a c	atalvet h	ut increases the a	ctivity of a cata	lust is
5.	called:		alaryst 0	fut increases the a	clivity of a cala	1yst 15
	A Enzyme	\cap	R	inhibitor	\cap	
	C. Promotor	X	D. D	Doisonor	X	
	C. FIOIIIOICI	U	D.	roisonei	U	
		Page	1 of 2			

6.	Diamond is a bad conductor A. It has a tight structor C. It has no free electron	or of electri ure O	city be B. D.	ecause: It has a high densit It is transparent to	y O light O
7.	Mixture containing 0.01 m having pKb = 5 has pH of:	nole/300cm ²	³ of NF	H_4Cl and 0.1 mole/4	00cm ³ of NH ₄ OH
	A. 4.00 C. 9.88		В. D.	4.12 10.00	0
8.	5g of urea (M.wt = 60) is a solution will be:	lissolved in	a 250 ci	m ³ of its solution. C	concentration of
	A. 5 % w/w C. 0.34 M		B. D.	5 % v/w 0.34m	8
9.	The gaseous element X ex combines with two volume What is the formula of hyde A. HX_2	ists in diato e of hydrog hride of X.?	omic fo en to fo B.	orm. One volume of orm two volume of HX ₃	the element X gaseous hydride.
10.	The number of bonds in or	ne molecule	D. e of Nit	ITA trogen is:	0
	A. one σ and one π C. three σ only	0	B. D.	one σ and two π two σ and one π	000
11.	Splitting of spectral lines bA.Zeeman effectC.Photoelectric effect	by placing the object of the o	he exc B. D.	ited atom in electric Stark effect Compton effect	field is called:
12.	In the ground state of an at A. in the valence shell C. nearest to the nucle	tom, the ele	ectron i B. D.	s present: in the second shell farthest from the m	ucleus O
13.	Which one of the followinA.iceC.silicon (IV) oxide	g exists in t	the soli B. D.	id state as a giant co iodine dry ice	valent lattice?
14.	pH of 1×10^{-4} M solution A. 1.10 C. 3.52	of Phospho	ric acio B. D.	d is: 2.02 4.13	0
15.	In which substance does n A. NO C. N ₂ O ₄	itrogen exh	ibit the B. D.	e highest oxidation s N ₂ O NaNO ₂	otate?
16.	The heat of neutralization NaOH + HCl	of the given \rightarrow NaC ization of th \rightarrow FeCl	h reacting $Cl + he$ folloor $l_2 + 2I$ B.	ion is -57.3 kJ H ₂ O owing reaction? H ₂ O -114.6kJ	0
17.	C228kJ Which of these samples of) gas contair	D. ns the s	-28.6kJ same number of ato	O ms as 1g of
	hydrogen molecule? (At. N A. 22 g of CO_2 C. 20 g of Ne	$ \begin{array}{c} \text{Mass} C = 1 \\ \text{O} \vdots \\ \text{O} \vdots \\ \end{array} $	12, O B. D.	= 16, $H = 1$, Ne 8 g of CH ₄ 8 g of O ₃	= 20)
		<u> </u>			<u> </u>
		Page 2 of	2		

Federal Board HSSC-I Examination Chemistry Model Question Paper (Curriculum 2006)

MCQ'S KEY

iВ	ii A	iii C	iv B	v C	vi C	vii C	viii C	ix C	х В
xi B	xii C	xiii C	xiv C	xv C	xvii B	xvii C			

Q. No. 2(i): The bond angles of H_2O and NH_3 are not 109.5° like that of CH_4 . Although O and N atoms are SP³hybridized like C. Give reason.

Ans: Bond angle in NH₃ is 107.5°. This is due to lone pair, bond pair repulsion is greater than bond pair, bond pair repulsion. So bond angle is reduced to 107.5° although there is sp³ hybridization in N. Bond angle in H₂O is 104.5°. This is due to lone pair, lone pair repulsion is greater than lone pair, bond pair repulsion. So bond angle is reduced to 104.5° although there is sp³ hybridization in oxygen. CH4 has no lone pair. So bond angle is 109.5° and C is sp3 hybridized.

Q. No. 2(ii) As both NF₃ and BF₃ are tetra atomic molecules but have different shape and geometry. Explain according to VSEPR theory.

Ans: NF₃, N 1s², Valence Shell



Shape of NF3 is trigonal pyramidal with angle 107.5°. This is due to one lone pair and three bond pair while in BF₃

₅B 1S², 2s², 2p¹ Valence Shell

Ground state

≜♦



2s 2px 2py 2pz

Excited state



Total electron pair Lone pair Bond pair

3 0 3

Shape is trigonal due to three bond pair and no lone pair. Bond angle is 120°.

Q. No. 2(iii) Ionic Crystals are brittle in nature but metals are malleable in nature. Give reason of your answer.

Ans: It is because ionic solid consist of parallel layers in which cations and anions are present in alternate positions. Thus, when a stress is applied on crystal one layer of ions slides a little bit over the other layer. In this way like ions come in front of each other, which repel each other and thus a crystal is broken and show brittleness. When stress is applied on metal then layer slip over each other and their shape is changed. Hence they can be changed into sheets malleable or were ductile without breaking.

Q. No. 2(iv) Derive the units for general gas constant 'R' in general gas equation.

- a. When the pressure is in Nm⁻² and volume in m³.
- b. When energy is expressed in ergs.

Ans: PV= nRT

$$\frac{PV}{nT} = R$$

 $R = \frac{1.01 \times 10^5 N/m^2 \times 0.022414m^3}{1 \text{ mole } \times 273 \text{ K}}$

- R = 8.314 Nm mol⁻¹k⁻¹
- $R = 8.314 \text{ J mole}^{-1}\text{k}^{-1}$

 $I J = 10^{7} erg$

 $8.314 \text{ J} = 8.314 \times 10^7 \text{ erg mol}^{-1}\text{k}^{-1}$

 $R = 8.314 \times 10^7 \text{ erg mol}^{-1} \text{k}^{-1}$

Q. No. 2(v) Justify that the distance gaps between different orbits of an atom go on increasing from the lower to the higher orbits

Ans: For H

Z=1

lf n=1₩⇒>ell

 r_1 = 0.529 A°× n²

r₁= 0.529 A°

 $r_2 = 0.529 A^{\circ}(2)^2 = 0.529 \times 4$

 $r_2 = 2.116 \text{ A}^{\circ}$

If n=3 → M shell

 $r_3 = 0.529 (3)^2$

 $r_3 = 0.529 \times 9$

r₃ = 4.761 A°

This shows that as n is increased gap between the orbits is also increased.

Q. No. 2(vi) Describe hybridization in acetylene (C_2H_2) molecule. Also draw diagram of hybridized orbitals in this molecule.

Ans: C₂H₂

₆C 1s², 2s², 2p²

Ground State



Excited State





2px 2py 2pz non hybridization orbital's 2ps

Sp hybridization



These are four hybrid orbital's. One δ bond is formed due to sp-sp end to end overlap. Two δ bonds are formed due to sp-1s end to end overlap. One π bond is formed due to 2py-2py lateral overlap. One π bond is formed due to 2pz-2pz lateral overlap.

Q. No. 2(vii) Interpret why water and ethanol can mix easily in all proportions.

Ans: Water and ethanol are mixed in all proportion. This is due to the hydrogen bonding this between water and ethanol. Oxygen attains partial negative charge (- δ) and hydrogen attain partial positive charge ($+\delta$). Hence H-bonding is formed in between ethanol and water. Secondly both exist in liquid state



Q. No. 2(viii) Justify that Bohr's equation for the wave number can explain the spectral lines of Lyman, Balmer and Paschen series.

Ans: From Bohr's wave number equation

$$\overline{\gamma} = 1.0967 \times 10^7 \text{m}^{-1} \left[\frac{1}{n_1^2} - \frac{1}{n_2^2} \right]$$

For Lyman's series $n_1 = 1$

$$\overline{\gamma}$$
= 1.0967 × 10⁷m⁻¹ $\left[\frac{1}{1^2} - \frac{1}{n_2^2}\right]$

Where $n_2 = 2, 3, 4....$

For Bulmer's series

$$\bar{\gamma}$$
= 1.0967 × 10⁷m⁻¹ $\left[\frac{1}{2^2} - \frac{1}{n_2^2}\right]$

Where n₂ = 3, 4, 5.....

For Paschalis Series

$$\bar{\gamma}$$
= 1.0967 × 10⁷m⁻¹ $\left[\frac{1}{3^2} - \frac{1}{n_2^2}\right]$

Where $n_2 = 4, 5, 6....$

Hence Bohr's equations justify the Lyman's, Bulmer's and Paschen's series.

Q. No. 2(ix) State Dalton's law. Also write its two applications.

Ans: Sum of the partial pressure of the non-reacting gases is equal to the total pressure of the gases

 $P_t = P_1 + P_2 + P_3 + \dots$

Where P_1 , P_2 and P_3 are the partial pressure of the gases and pt is the total pressure of the gases.

Applications:

- 1. When gas is collected over vapors of water then
- $P_{gas} = P_{total} P_{H2O}$

Where P_{H2O} is called aqueous tension. The respiration in living things depends upon the difference in partial pressure. Partial pressure of O₂ outside is 159g/cm² than in lungs, where the pressure is 116 g/cm²at higher altitudes becomes 150g/cm²so pilot may have uncomfortable breathing.

Q. No. 2(x)The melting and boiling points of hydrazine (N₂H₄) are much higher than those of ethane (C₂H₄). Suggest reasons for these differences in terms of the intermolecular forces each compound possesses.

Ans: N₂H₄ has lone pair and H- bonding. So M.P and B.P of N₂H₄ is much greater while in ethane C₂H₆ there is weak London dispersion force. So B.P is very low. B.P of C₂H₆ is -88.6°C



H-bonding





Ans: Effective collisions bring about the reaction. For a collision to be effective molecules must posses the activation energy and must be properly oriented. At ordinary temperature very few molecules possess this energy of activation. All the molecules of a reactant do not possess the same energy at a particular temperature. Most of them possess average energy. A fraction of molecules has kinetic energy more than the average energy. The number of molecules having at least kinetic energy equal to Ea at temperature T in proportional to the shaded area under the Maxwell Boltzmann curve of kinetic energy. As the temp is increased the area of the shaded region increases and more molecules have kinetic energy greater than Ea. An increase in temp increase the number of reactant molecules that have enough energy for effective collision.

Q. No. 2(xii)An aqueous solution of ammonium Chloride is acidic and that of sodium acetate is basic in nature. Give reason with the help of equations.

Ans:NH₄Cl + H-OH → HCl + NH₄OH Strong acid weak base

On hydrolysis of NH4Cl, strong acid HCl is produced. Hence ammonium chloride aqueous solutions is acid PH<7

On hydrolysis of CH₃COONa, strong base NaOH is produced. Hence sodium acetate aqueous solution is basic in nature PH>7

Q. No. 2(xiii)Calculate molarity of aqueous solution of sulfuric acid from the following data.

Molar mass	Molarity	Density in g/Cm ³	
98	18	1.84	

Ans: Molarity = 18 mol/dm³ = 18 mol/ 1000cm³

Density = 1.84 cm^{-3}

$$d = \frac{m}{m}$$

m = d × v

 $m = 1.84 \times 1000$

 $m = 1840 \text{ g } H_2 \text{SO}_4$

Mass of H_2SO_4 = Mole × Molar mass

Mass of $H_2SO_4 = 18 \times 98 = 176 \text{ g} H_2SO_4$

Mass of water = $1840 - 1764 = 76 \text{ g H}_2\text{O}$

 $Molarity = \frac{Mole \times 1000}{Mass of H20}$ $Molarity = \frac{18 \times 1000}{76}$ Molarity = 236.84m

Q. No. 2(xiv)Lattice energies of LiCl and KCl are 833 kJ/mol and 690 kJ/mol, respectively. Discuss why is lattice energy of LiCl greater than KCl?

Ans: Size of Li⁺ is smaller than size of K⁺ hence force of attraction between Li⁺ and Cl⁻¹ is much greater as compare to force of attraction between K⁺ and Cl⁻¹. These force lattice energy of LiCl is Q33kJ/mol while that of KCl is 690kJ/mol

 $LiCl(s) \longrightarrow Li^+ + Q^-(g)$

KCl(s) → K⁺ + Q⁻(g)

Size of Li⁺< size of K⁺

Q. No. 2(xv)Benzene (C₆H₆) is an aromatic hydrocarbon which exists as a liquid at room temperature.

 $\Delta H_1 = +3268 \text{ kJ/mol}$

 $\Delta H_2 = -393 \times 6$

Using the following standard enthalpy changes:

Heat of formation of CO₂ = -393 KJ / mol

Heat of formation of $H_2O = -286 \text{ KJ} / \text{mol}$

Heat of combustion of $C_6H_6 = -3268 \text{ KJ} / \text{mol}$

Calculate the enthalpy change of formation of C₆H₆.

Ans:

 $C_6H_6 + \frac{15}{2}O_2 \longrightarrow 6CO_2 + 3H_2O \qquad \Delta H1 = -3268 \text{ kJ/mol}$

For formation reverse the equation

- (i) $6CO_2 + 3H_2O \longrightarrow C_6H_6 + \frac{15}{2}O_2$
- (ii) $6C + 6O_2 \longrightarrow 6CO_2$
- (iii) $3H_2 + \frac{3}{2}O_2 \longrightarrow 3H_2O\Delta H_3 = -286 \times 3$

Adding above equation

From Hisses Law

 $\Delta H_{f} = \Delta H_{1} + \Delta H_{2} + \Delta H_{3}$

 $\Delta H_{\rm f} = +3258 - 2358 - 858$

 $\Delta H_f = 52 kJ/mol$

Q. No. 2(xvi)Consider the Standard electrode potentials

 $Ag^{+}/Ag = 0.7994V$, $Fe^{3+}/Fe = 0.771V$

Write the half-cell reactions at each electrode. Also write overall reaction

Ans: $E_{cell} = E_{cahode} - E_{anode}$

 $E_{cell} = 0.7994 - 0.771$

 $E_{cell} = +0.02Q4$

At cathode Reduction

(Ag⁺ + ∉ → Ag) × 3

At Anode Oxidation

 $Fe \longrightarrow Fe^{+3} + 3e^{-1}$

3Ag⁺ + Fe → 3Ag + Fe⁺³

Overall reaction

Q. No. 2(xvii) Chemical kinetics is concerned with rates of chemical reactions and factors that affects the rates of chemical reactions. Consider the following steps of reactions:

FeCl ₃ (aq) + 2Kl (aq) Fel ₂ (aq) + 2KCl (aq) + Cl ⁻ (aq) (slo

2KI(aq) + 2Cl⁻(aq) → 2KCl(aq) +l₂(S) (fast)

a. Write the rate expression for the above reaction.

b.Give the order of reaction for the above reaction.

Ans:

- a. Rate Law expression Rate = K [*FeCl*₃][*KI*]²
- b. Order of reaction = 1 + 2

Order of reaction = 3^{rd} order reaction

Q. No. 2(xviii)What is reverse osmosis? Give its daily life applications.

Ans: If a solution in contact with pure solvent across a semi permeable membrane is subjected to an external pressure equal to osmotic pressure, it stops osmosis. If external pressure is greater than solutions osmotic pressure, it will force solvent to flow from solution to solvent. This process is called reverse osmosis.

Daily life application of reverse osmosis:

Sea water is highly hypertonic to body fluids and this is not drinkable. By reverse osmosis it is subjected to desalination i.e remove large amounts dissolve salts from sea water. The sea water is pumped under high pressure 20 atm through the semi permeable membrane, which allow water molecules to pass and stop ions.

Q. No. 2(xix)How to calculate the molecular mass of the solute by using $\Delta P/P^0 = X_2$?

Ans: From Roult's Law

$$\frac{\Delta P}{P^{\circ}} = X_2 \qquad \longrightarrow \text{Eq (1)}$$

Since $X_2 = \frac{n_2}{n_1 + n_2}$

Where n_2 is the mole of solute. Neglect the n_2 from denominator for dilute solution.

$$X_2 = \frac{n_2}{n_1}$$
$$n_1 = \frac{W_1}{M_1}$$
$$n_2 = \frac{W_2}{M_2}$$

Put in above equation

$$X_2 = \frac{w_2}{M_2} \times \frac{M_1}{W_1}$$

Put in eq 1

$$\frac{\Delta P}{P^{\circ}} = \frac{W_2 \times M_1}{M_2 \times W_1}$$

$$\mathsf{M}_2 = \frac{P^{\circ}}{\Delta P} \times \frac{W_2 M_1}{W_1}$$

Q. No. 2(xx)How to calculate standard electrode potential? Explain briefly.

Ans: Standard hydrogen electrode is used to know the electrode potential. It is made up of platinum wire, which is sealed in glass tube. It is dipped in 1MHCl solution. H₂ gas is introduced from the top. Oxidation as well as reduction potential of hydrogen is zero. So voltmeter shows the electrode potential of the required element.



Q. No. 3 (a)Derive the equation for the radius of nth orbit of hydrogen atom using Bohr's model.

Ans: Electrons revolve around the nucleus in circular path. So centrifugal force is given by



Coulomb's force of attraction between electron and proton in given by

$$\mathsf{F} = \frac{Kq_1q_2}{r^2}$$

Where $r_1 = \frac{1}{2}e$ and $q_2 = e^ K = \frac{1}{4\pi\epsilon_{\circ}}$

€ is the primitivity constant of the vacuum

€•= 8.85 × 14⁻¹²
$$C^2 N^{-1} m^{-2}$$

Put in above eq

$$F = \frac{Zee}{4\pi \in r^2}$$
$$F = \frac{Ze^2}{4\pi \in r^2} \longrightarrow Eq(2)$$

Where Z is the atomic number

These two forces are equal. So by comparing Eq(1) and Eq(2)

$$\frac{mv^2}{r} = \frac{Ze^2}{4\pi \in r^2}$$
$$V^2 = \frac{Ze^2}{4\pi \in rm} \qquad \qquad \blacksquare \mathbf{E}q(3)$$

From Bohr's postulate angular momentum is given by

$$mvr = \frac{nh}{2\pi}$$

Where n is the principal quantum number

n = 1, 2, 3.....

Shell K L M

- h = plank's constant
- $h = 6.625 \times 10^{-34}$

m is the mass of electron

v is the velocity of electron

r is the radius

$$v = \frac{nh}{2\pi mr}$$

Squaring both sides

$$v^2 = \frac{n^2 h^2}{4\pi^2 m^2 r^2} \qquad \longrightarrow \text{Eq(4)}$$

By comparing Eq(3) and Eq(4)

 $\frac{Ze^{2}}{4\pi \in rm} = \frac{n^{2}h^{2}}{4\pi^{2}m^{2}r^{2}}$ $\frac{Ze^{2}}{\in \circ} = \frac{n^{2}h^{2}}{\pi mr}$ $r = \frac{n^{2}h^{2} \in \circ}{\pi mZe^{2}}$ $r = \frac{h^{2} \in \circ}{\pi mZe^{2}} \times \frac{n^{2}}{Z} \longrightarrow Eq(5)$ Let $a^{\circ} = \frac{h^{2} \in \circ}{\pi me^{2}}$ $a^{\circ} = \frac{(6.625 \times 10^{-34})^{2}(8.85 \times 10^{-12})}{3.14 \times 9.1 \times 10^{-31} \times (1.6 \times 10^{-19})^{2}}$ $a^{\circ} = 0.529 \times 10^{-10}$ $a^{\circ} = 0.529 \text{ A}^{\circ}$ Put in Eq(5)

$$r = a^{\circ} \times \frac{n^2}{z}$$

r= 0.529 A° × $\frac{n^2}{z}$

For hydrogen $\frac{1}{2} = 1$

r= 0.529 A° × n^2

(b)Ammonia Solvay process is used to manufacture sodium carbonate. During this process ammonia is recovered by the following reaction.

2NH₄Cl + Ca(OH)₂ CaCl₂ + 2H₂O +2NH₃ When 100 g of ammonium chloride and 150 g calcium hydroxide are used then (At. Mass N=14 H=1 Cl= 35.5 Ca=40)

- i. Calculate the mass in kg of ammonia produce during chemical reaction.
- ii. Calculate the excess mass in gram of one of the reactant left unreacted.

Ans:

Mass of NH₄Cl = 100gMass of Ca(OH)2 = 150gMolar mass of NH₄Cl = 14+1×4+35.5Molar mass of Ca(OH)2 = 40+16×2+1×2Molar mass of NH₄Cl =53.5g/molMolar mass of Ca(OH)2 = 74 g/molMolar of NH₄Cl =
$$\frac{Mass in gram}{Molar mass}$$
Molar of Ca(OH)2 = 74 g/molMolar of NH₄Cl = $\frac{100}{53.5}$ Molar of Ca(OH)2 = $\frac{Mass in gram}{Molar mass}$ Molar of NH₄Cl = 1.87 mole of NH₄ClMolar of Ca(OH)2 = $\frac{150}{74}$ From equationI mole of Ca(OH)2 = 2.03 moles of Ca(OH)2Prom equation1 mole of Ca(OH)2 = 2 moles of NH31.87 moles of NH₄Cl = x $= \frac{2}{2} \times 1.87$ $= \frac{2}{2} \times 1.87$ $x = 4.06$ moles of NH3

Since NH₄Cl produces the least amount of NH₃. So NH₄Cl is the limiting reactant.

Mass of NH₃ = mole × molar mass of NH₃

Mass of $NH_3 = 1.87 \times 17$

Mass of $NH_3 = 31.97 \text{ g } NH_3$

Mass of NH₃ = $\frac{31.97}{1000}$

Mass of $NH_3 = 0.03197$ Kg of NH_3

(ii) From equation

2 moles of $NH_4Cl = 1$ mole of $Ca(OH)_2$

1.87 moles of $NH_4Cl = x$

$$\mathbf{x} = \frac{1.87 \times 1}{2}$$

x = 0.935 moles of Ca(OH)₂used

 $Ca(OH)_2 left = 2.03 - 0.935$

Ca(OH)₂ = 1.095 moles of Ca(OH)₂left

Mass of Ca(OH)₂left = mole × molar mass

Mass of Ca(OH)₂left = 1.095×74

Mass of $Ca(OH)_2$ left = Q1.03 g $Ca(OH)_2$ left

Q. No. 4Consider the following reaction:

N₂ + 3 H₂ 2NH₃

(a) Derive expression of Kc for the above reaction and calculate equilibrium concentration of N₂. The equilibrium concentration of H₂ and NH₃ are 1.0 moldm³ and 0.5 moldm⁻³ respectively. Kc of above reaction at 25°C is 1.85×10^3 .

Ans:

$$N_2 + 3H_2 \xrightarrow{} NH_3$$

At t = 0

a/v mol/dm³ b/v mol/dm³

$$\frac{a-x}{v}\frac{b-3x}{v}\frac{2x}{v}$$

K_c is written as

$$\mathsf{K}_{\mathsf{C}} = \frac{[NH_3]^2}{[N_2][H_2]^3}$$

By putting respective values.

$$K_{c} = \frac{\left[\frac{2x}{v}\right]^{2}}{\left[\frac{a-x}{v}\right]\left[\frac{b-x}{v}\right]^{3}}$$
$$K_{c} = \frac{4x^{2}}{(a-x)\frac{(b-3x)^{3}}{v^{4}}}$$

$$K_{c} = \frac{4x^{2}v^{2}}{(a-x)(b-3x)^{3}}$$

$$K_{c} = 1.85 \times 10^{3} \qquad V = 1 dm^{3}$$
Equilibrium concentration
$$[H_{2}] = 1.0 \text{ mole/dm}^{3}$$

$$[NH_{3}] = 0.5 \text{ mole/dm}^{3}$$

$$[NH_{3}] = 2x = 0.5$$

$$x = \frac{0.5}{2}$$

$$x = 0.25 \text{ mole/dm}^{3}$$

$$[N_{2}] = a - x$$

$$K_{c} = \frac{4x^{2}v^{2}}{(a-x)(b-3x)^{3}}$$

$$1.85 \times 10^{3} = \frac{4(0.25)^{2} \times 1}{(a-x) \times (1)^{3}}$$

$$(a - x) = [N_{2}] = \frac{4(0.25)^{2}}{1.85 \times 10^{3}}$$

$$[N_{2}] = 1.35 \times 10^{-4} \text{ mole/dm}^{3}$$

Equilibrium concentration of N_2

Q. No. 4 (b)Balance the following chemical equation in an acidic medium

$$Cr^{3+} + BiO_3^{1-} \longrightarrow Cr_2O_7^{2-} + Bi^{3+}$$

Ans:

$$Cr^{+3} + BiO_3^{-1} - Cr_2O_7^2 + BC^{3+}$$

Step 1 write Oxidation number

$$Cr^{+3} + [Bi^{+5}O_3^{-2} \times 3]^{-1} \longrightarrow [Cr_2^{+6} \times 2O_7^{-2} \times 7]^{2-} + BiC^{3+}$$

Steps 2 write half oxidation reaction. Balance e⁻

$$2Cr^{+3} \longrightarrow [Cr_2^{+6 \times 2}O_7^{-2 \times 7}]^{2-} + 6e^{-1}$$

Step 3 Balance oxygen by adding 7H₂O on LHS

 $2Cr^{+3} + 7H_20 \longrightarrow Cr_2O_7^{2-} + 6e^- + 14H^+Eq(1) \longrightarrow$

Steps 4 write half reduction reaction. Balance the e⁻

 $[Bi^{+5}O_3^{-2\times 3}]^{-1} + 2e^{-}Bi^{3+}$

Step 5Balance oxygen by adding 3H₂O on RHS

 $BiO_3^{-1} + 3e^- + 6H^+ \longrightarrow Bi^{3+} + 3H_20Eq(2) \longrightarrow$

Step 6 Balance the electrons by Xing Eq(2) by '2' and then add in Eq (1)

$$2Cr^{+3} + 7H_20 \longrightarrow Cr_2O_7^{2-} + 6e^- + 14H^+$$

$$2BiO_3^{-1} + 6e^- + 12H^+ \longrightarrow 2Bi^{3+} + 6H_2O$$

$$2Cr^{+3} + 7H_20 + 2BiO_3^{-1} \longrightarrow Cr_2O_7^{2-} + 14H^+ + 2Bi^{3+}$$

Balanced equation.

Q. No. 5 (a)Phosgene (COCl₂) is a toxic gas. This gas is prepared by the reaction of carbon monoxide with chlorine.

 $CO(g) + Cl_2(g) \longrightarrow COCl_2(g)$

The following data were obtained for kinetic study of this reaction.

Experiment	Initial [CO]	Initial [Cl ₂]	Initial rate (moles dm ⁻³ s ⁻¹)
1	1.000	0.100	1.29×10 ⁻²⁹
2	0.100	0.100	1.30×10 ⁻³⁰
3	0.100	1.000	1.30×10 ⁻³⁰

Use the data in the table to deduce the order of the reaction with respect to CO and Cl₂.Hence write a rate law/equation for this reaction.

Ans:

Exp	Initial conc [CO]	Initial con [<i>Cl</i> ₂]	Initial rate
1	1.0	0.1	1.29 × 10 ⁻²⁹
2	0.1	0.1	1.30 ×10 ⁻³⁰
3	0.1	1.0	1.31 ×10 ⁻³⁰

In experiment 1 and conc of Cl_2 is constant and conc of CO vary. Ratio of conc is
Exp. 1 : Exp. 2
1.0 0.1
0.1 0.1
10:1
Ratio of rate of reaction is exp 1 and 2
Exp. 1 : Exp. 2
1.29×10 ⁻²⁹ 1.30×10 ⁻³⁰
1.30×10 ⁻³⁰ 1.30×10 ⁻³⁰
10:1
Hence it is 1 st order W.r.t [<i>CO</i>]
Rate = K [<i>CO</i>] ¹
1 st order W.r.t CO
Conc of CO in experiment 2 and 3 is constant but conc of Cl_2 vary. Conc ratio is
Exp. 2 : Exp. 3
0.1 1.0
0.1 0.1
1:10
Rate ratio of Exp 2 and 3
Exp.2 : Exp. 3
1.3×10 ⁻³⁰ 1.3×10 ⁻³⁰
1.3×10 ⁻³⁰ 1.3×10 ³⁰
1:1

As the conc is increased to 10 times but rate is constant. So Cl₂ is zero order W.r.t Cl₂

Rate = $K[Cl_2]^0$

Zero order of reaction W.r.t Cl₂

Overall rate Law

Rate = $K[CO]^1[Cl_2]^0$

Overall order of reaction = 1 + 0

Overall order of reaction = 1st order

(B)Show the diamagnetic/paramagnetic nature of O_2 , $O_2^{2^+}$ and $O_2^{2^-}$ with the help of molecular orbital theory.

Ans:

O₂





2s 2px 2py 2pz



Paramagnetic behaviour:

There are two compare electron in antibonding. So oxygen shows paramagnetic behaviour i.e attract towards magnet.

 0_{2}^{-2}

 O_2^{+2}



