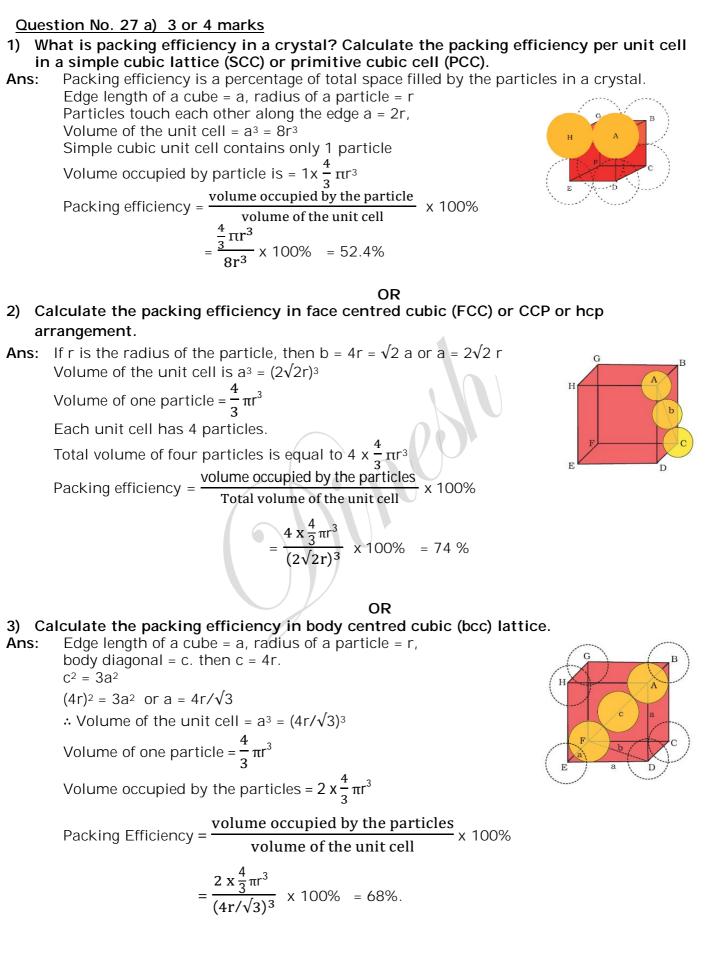
# UNIT-1 SOLID STATE



# Question No. 27 b) 2 marks

### 1) Calculate the number of particles present in per unit cell in SCC.

In SCC, the particles present at only eight corners and each particle is shared by eight Ans: other unit cells.

Therefore, the total number of particles present in a unit cell is

$$= 8 \times \frac{1}{8} = 1$$
 particle.

# OR

- 2) Calculate the number of particles present in per unit cell in FCC.
  - Ans: In FCC, the particle present at centre of each face is shared by two other unit cells where as each particle at corner is shared by eight other unit cells.

Therefore, the total number of particles present in a unit cell is

$$= 8 x \frac{1}{8} + 6 x \frac{1}{2} = 1 + 3 = 4$$
 particles.

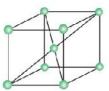
## 3) Calculate the number of particles present in per unit cell in BCC.

In BCC, the particle present at body centre is shared only unit cell where as each particle Ans: at corner is shared by eight other unit cells.

Therefore, the total number of particles present in a unit cell is

$$= 8 x \frac{1}{8} + 1 x 1 = 1 + 1 = 2 \text{ particles}$$

Ans:



### OR

4) An element having atomic mass 107.9g/mol has fcc unit cell. The edge length of unit cell is 408.6pm. Calculate the density of the unit cell. (Given  $N_A = 6.022 \times 10^{23} \text{mol}^{-1}$ ).

N<sub>A</sub> = 6.022 x 10<sup>23</sup> per mol, d = ?  
Density (d) = 
$$\frac{z.M}{N_A.a^3} = \frac{4 \times 107.9 \times 10^{-3}}{6.022 \times 10^{23} \times (408.8 \times 10^{-12})^3} = 1.05 \times 10^{-2} \text{ kgm}^{-3}$$

OR

5) Sodium metal crystallizes in a BCC structure. Its unit cell edge length is 420pm. Calculate its density. (atomic mass of sodium = 23u, NA =  $6.022 \times 10^{23}$  mol<sup>-1</sup>). Α

Ans: Given, z = 2, a = 420 pm = 288 x 
$$10^{-12}$$
 m, M = 23u = 23 x  $10^{-3}$ , N<sub>A</sub> = 6.022 x  $10^{23}$  per mol, d = ?

Density (d) = 
$$\frac{z.M}{N_A.a^3} = \frac{2 \times 23 \times 10^{-3}}{6.022 \times 10^{23} \times (420 \times 10^{-12})^3} = 3.73 \text{ kgm}^{-3}$$

OR

6) An element has a body centred cubic structure with a cell edge of 288 pm. The density of the element is 7.2 g/cm<sup>3</sup>. How many atoms are present in 208 g of the element?

z = 2,  $a = 288 \text{ pm} = 288 \text{ x} 10^{-10} \text{ cm}$ ,  $d = 7.2 \text{ g/cm}^3$ , Ans: Given,  $N_A = 6.022 \times 10^{23} \text{ per mol}$ 

i.e d = 
$$\frac{z.M}{N_A.a^3}$$

$$\therefore M = \frac{d.N_A.a^3}{z} = \frac{7.2 \times 6.022 \times 10^{23} \times (288 \times 10^{-10})^3}{2} = 51.79 \text{ g/mol}$$
79 g of element contains 6.022 x 10<sup>23</sup> atoms.

51.79 g of element contains  $6.022 \times 10^{23}$  atoms.

 $6.022 \times 10^{23} \times 208$ 208 g of the element contains = - atoms = 24.19 x 10<sup>23</sup> atoms. 51.79

Question No. 11) 2 marks 1) A cubic unit cell consists of three elements A, B & C. Atoms of A are present at the corners, atoms of B at the body centre and atoms of C at the centre of each face. What is the formula of the compound? **Ans:** No. of atoms A per unit cell = 8 x  $\frac{1}{8}$  = 1 No. of atoms B per unit cell =  $1 \times 1 = 1$ No. of atoms C per unit cell =  $6 \times \frac{1}{2} = 3$ A:B:C = 1:1:3Therefore, the formula of the compound is ABC<sub>3</sub> OR 2) A compound id formed by two elements X and Y. Atoms of the element Y (as anions) make ccp and those of the element X (as cations) occupy all the octahedral voids. What is the formula of the compound? **Ans:** Let the no. of atoms of Y involved in ccp be 'x' Therefore, no. of octahedral voids = xGiven that all the octahedral voids are occupied by atoms of X. Therefore, no. of atoms of X = xThus, ratio of X : Y = x : x = 1 : 1Hence the formula of the compound is XY. OR 3) Write any four differences between crystalline and amorphous solids. Ans: Crystalline solids Amorphous solids 1. These have regular arrangement of These have no regular arrangement of 1 constituent particles. constituent particles. These have definite shape and definite 2. These don't have definite shape and 2. definite volume. volume. 3. These have sharp melting points. 3. Melting points over a range of temperature. Anisotropic in nature. 4. Isotropic in nature. 4. These are true solids. 5. 5. These are not true solids. OR 4) Write any two differences between Schoktty defect and Frenkel defect. Ans:

| Schoktty defect |                                    | Frenkel defect |                                    |
|-----------------|------------------------------------|----------------|------------------------------------|
| 1.              | Equal number of cations and anions | 1.             | An ion occupies interstitial site. |
|                 | are missing from their lattice.    |                |                                    |
| 2.              | Decrease in density of solids.     | 2.             | No change in density of solids.    |

# Unit -2 Solutions

| Unit -2 Solutions  |  |  |  |
|--|--|--|--|
| Question No. 28) <u>3 Marks:</u>   |  |  |  |
| 1) The vapour pressure of pure benzene at certain temperature is 0.850bar. A 0.5g of   |  |  |  |
| non-volatile solute is dissolved in 39g of benzene (molar mass of benzene is 78  |  |  |  |
| g/mol). Then the vapour pressure of the solution is 0.845bar. Calculate the molar  |  |  |  |
| mass of the solute.  |  |  |  |
| Solution: Given, p <sup>o</sup> = 0.850bar, p = 0.845bar, Then, p <sup>o</sup> - p = 0.850 - 0.845 = 0.005bar                                  |  |  |  |
|  |  |  |  |
| $W_1 = 39g, \qquad M_1 = 78g/mol$  |  |  |  |
| $W_2 = 0.5g, \qquad M_2 = ?$   |  |  |  |
| $w_2M_1$ p <sup>0</sup>  |  |  |  |
| $M_2 = W_1 \times \overline{p^0 - p}$  |  |  |  |
|  |  |  |  |
| 0.5 x 78 x 0.850   |  |  |  |
| $M_2 = \frac{39 \times 0.005}{39 \times 0.005} = 170 \text{g/mol}$   |  |  |  |
| 37 X 0.000   |  |  |  |
|  |  |  |  |
| OR   |  |  |  |
| 2) The boiling point of benzene is 353.23K when 1.80g of a non-volatile, non-ionising  |  |  |  |
| solute was dissolved in 90g of benzene, the boiling point is raised to 354.11K.  |  |  |  |
|  |  |  |  |
| Calculate the molar mass of the solute. (Given $K_b$ for benzene = 2.53Kkg/mol)  |  |  |  |
| <b><u>Solution</u></b> : Given, $w_1 = 90g$ , $w_2 = 1.80g$ , $K_b = 2.53Kkg/mol$  |  |  |  |
| $\Delta T_f = T - T_0 = 354.11 - 353.23 = 0.88K \qquad M_2 = ?$  |  |  |  |
| K <sub>b</sub> x w <sub>2</sub> x 1000 2.53 x 1.80 x 1000  |  |  |  |
| $\therefore M_2 = \frac{K_b \times W_2 \times 1000}{W_1 \times 100} = \frac{2.53 \times 1.80 \times 1000}{00 \times 0.00} = 57.5 \text{g/mol}$ |  |  |  |
| $w_1 \times \Delta T_b$ 90 x 0.88  |  |  |  |
|  |  |  |  |
| OR   |  |  |  |
| 3) A 0.643g of a solute is dissolved in 50ml of benzene (density is 0.879g/ml) decreases   |  |  |  |
| the freezing point from 278.66K to 278.18K. If K <sub>f</sub> for benzene is 5.12Kkg/mol.  |  |  |  |
| Calculate the molar mass of the solute.  |  |  |  |
|  |  |  |  |
| <b>Solution:</b> Given, $w_2 = 0.643g$ , $K_f = 5.12 \text{ Kkg/mol}$  |  |  |  |
| $\Delta T_{\rm f} = T_0 - T = 278.66 - 278.18 = 0.48 \rm K$  |  |  |  |
| Volume 'v' = $50ml$ , d = $0.879g/ml$  |  |  |  |
| $w_1 = v \times d = 50 \times 0.879 = 43.95g$ $M_2 = ?$  |  |  |  |
| S.   |  |  |  |
| $\therefore M_2 = \frac{K_f x w_2 x 1000}{w_1 x \Lambda T_f} = \frac{5.12 x 0.643 x 1000}{43.95 x 0.48} = 156.1 g/mol$                         |  |  |  |
| $w_1 \times \Delta T_f$ = 43.95 x 0.48   |  |  |  |
|  |  |  |  |
| OR   |  |  |  |
|  |  |  |  |
| 4) A 200cm <sup>3</sup> of an aqueous solution of a protein contains 1.26g of protein. The osmotic   |  |  |  |
| pressure of the solution at 300K is found to be 2.57 x 10 <sup>-3</sup> bar. Calculate the molar   |  |  |  |
| mass of the protein (Ideal gas constant R = 0.083 litre bar/K/mol).  |  |  |  |
| <b>Solution:</b> Given, $V = 200 \text{ cm}^3 = 0.2 \text{ L}$ . $w = 1.26 \text{ g}$ , $T = 300 \text{ K}$                                    |  |  |  |
| $\pi = 2.57 \text{ x } 10^{-3} \text{bar},  \text{R} = 0.083 \text{ litre bar/K/mol}  \text{M} = ?$  |  |  |  |
|  |  |  |  |
| wRT  |  |  |  |
| $\pi = \frac{MV}{MV}$  |  |  |  |
|  |  |  |  |
| wRT 1.26 x 0.083 x 300 (1.000 ( 1.000 ( 1.000 )  |  |  |  |
| $M = \frac{\pi V}{\pi V} = \frac{1.20 \times 0.000 \times 0.00}{2.57 \times 10^{-3} \times 0.2} = 61.039 \text{g/mol}$                         |  |  |  |
|  |  |  |  |
|  |  |  |  |
|  |  |  |  |

### Unit - 4 Chemical kinetics

<u>Ouestion No. 30 a) 3 Marks:</u> 1) Derive an integrated rate equation for the velocity constant or rate constant of a zero order reaction. Ans: Consider a zero order reaction  $\mathbb{R} \longrightarrow \mathbb{P}$   $\mathbb{R}ate = -\frac{d[\mathbb{R}]}{dt} = k[\mathbb{R}]^0$   $-\frac{d[\mathbb{R}]}{dt} = k \times 1$   $d[\mathbb{R}] = -k dt$ Integrating both sides, we get  $[\mathbb{R}] = -kt + 1$  --- (1) where I is integration constant At, t = 0,  $[\mathbb{R}] = [\mathbb{R}]_0$ , eqn (1) becomes  $[\mathbb{R}]_0 = 1$  ---- (2) where  $[\mathbb{R}]_0$  is the initial concentration of the reactant. Substitute (2) in (1)  $[\mathbb{R}] = -kt + [\mathbb{R}]_0$ 

### OR

2) Derive an integrated rate equation for the velocity constant or rate constant of a first order reaction.

Ans: Consider a first order reaction R  $\longrightarrow$  P

Rate = 
$$-\frac{d[R]}{dt} = k[R]$$
  
 $\frac{d[R]}{[R]} = -kdt$   
Integrating the equation, we get  $\ln[R] = -kt + I$  --- (i)  
When t = 0, R =  $[R]_0$ , eqn (1) becomes  $\ln[R]_0 = I$  --- (i)  
Substitute (2) in (1)  
 $\ln[R] = -kt + \ln[R]_0$   
 $k = \frac{1}{t} \ln \frac{[R]_0}{[R]}$   
 $k = \frac{2.303}{t} \log_{10} \frac{[R]_0}{[R]}$ 

OR

3) Rate constant of a first order reaction is 0.0693 min<sup>-1</sup>. Calculate the percentage of the reactant remaining at the end of 60 minutes.

**Ans:** Given k = 0.0693 min<sup>-1</sup>, t = 60 minutes, [R]<sub>0</sub> = 100%, R = ?

$$k = \frac{2.303}{t} \quad \log_{10} \frac{[R]_0}{[R]}$$
  
0.0693 =  $\frac{2.303}{60} \quad \log_{10} \frac{100}{[R]}$   
R = 1.56%

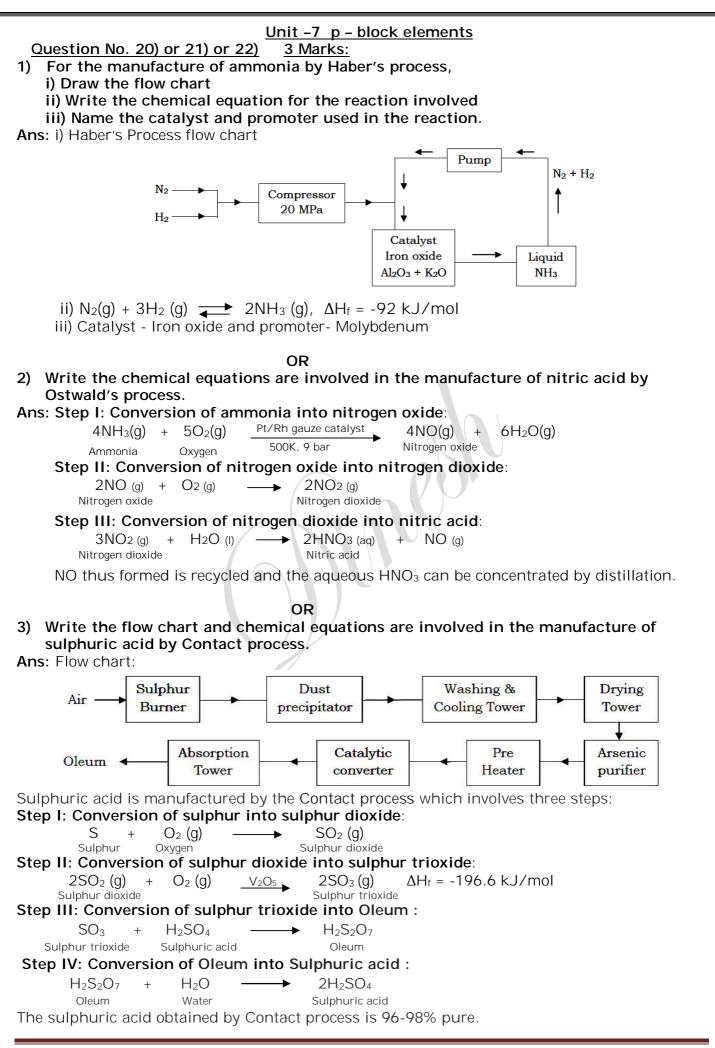
OR

4) The rate of a reaction increases by 4 times when the temperature of the reaction is raised from 340 K to 360 K. Calculate the energy of activation of the reaction. Given R = 8.314 J/K/mol.

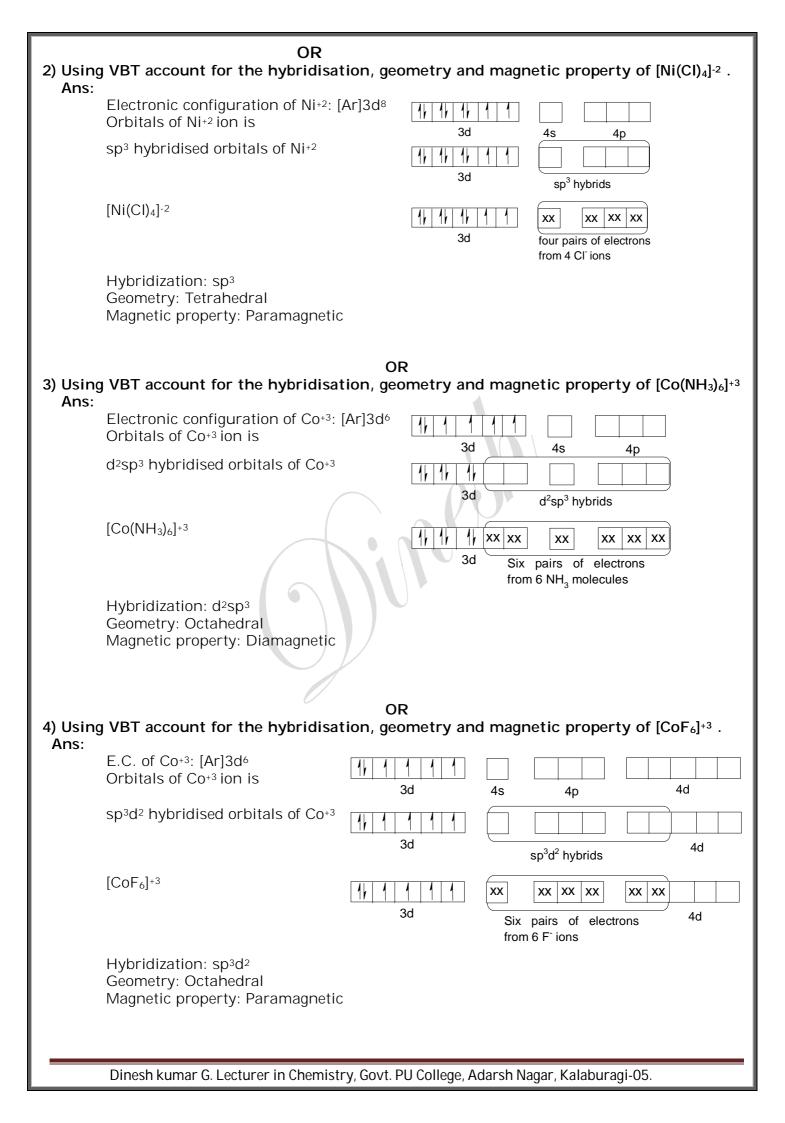
**Ans:** Given  $log(k_2/k_1) = 4$ ,  $T_1 = 340$ ,  $T_2 = 360$ , Ea = ?

$$\log \frac{k_2}{k_1} = \frac{Ea}{2.303R} \left(\frac{T_2 - T_1}{T_1 T_2}\right)$$
$$\log 4 = \frac{Ea}{2.303 \times 8.314} \left(\frac{360 - 340}{340 \times 360}\right)$$
$$\log 4 = \frac{Ea}{2.303 \times 8.314} \left(\frac{360 - 340}{340 \times 360}\right)$$

Ea = 70554J or 70.554 kJ



| OR  |  |  |  |  |
|---|--|--|--|--|
| 4) How is chlorine prepared using KMnO <sub>4</sub> ? (2 Marks)   |  |  |  |  |
| Ans: Action of conc. HCI on KMnO <sub>4</sub>   |  |  |  |  |
| $2KMnO_4 + 16HCI \longrightarrow 2KCI + 2MnCI_2 + 8H_2O + 5CI_2$  |  |  |  |  |
|   |  |  |  |  |
| Unit -8 d & f - block elements  |  |  |  |  |
| Question No. 24) <u>3 Marks:</u>  |  |  |  |  |
| 1) Write the chemical equations are involved in the manufacture potassium   |  |  |  |  |
| permanganate (KMnO <sub>4</sub> ).  |  |  |  |  |
| Ans: Step I: Conversion of manganese dioxide into potassium manganate:  |  |  |  |  |
| $2MnO_2 + 4KOH + O_2 KNO_3 2K_2MnO_4 + 2H_2O$   |  |  |  |  |
| Manganese Potassium Oxygen Potassium<br>dioxide hydroxide manganate   |  |  |  |  |
| Step II: Conversion of potassium manganate into potassium permanganate:   |  |  |  |  |
|   |  |  |  |  |
| $3K_2MnO_4 + 4H^+ \longrightarrow 2KMnO_4 + MnO_2 + 2H_2O$<br>Potassium Acid Potassium  |  |  |  |  |
| Manganate permanganate  |  |  |  |  |
|   |  |  |  |  |
| OR  |  |  |  |  |
| 2) Write the chemical equations are involved in the manufacture of potassium  |  |  |  |  |
| dichromate ( $K_2Cr_2 O_7$ ) from chromite ore.   |  |  |  |  |
| Ans. The manufacture of potassium dichromate involves the following steps.  |  |  |  |  |
| Step I: Conversion of chromite ore into sodium chromate:  |  |  |  |  |
| 4FeO.Cr <sub>2</sub> O <sub>3</sub> + 8Na <sub>2</sub> CO <sub>3</sub> + 7O <sub>2</sub> <u>900∘C</u> 8Na <sub>2</sub> CrO <sub>4</sub> + 2Fe <sub>2</sub> O <sub>3</sub> + 8CO <sub>2</sub> ↑<br>Chromite ore Sodium Oxygen Sodium Iron oxide  |  |  |  |  |
| carbonate chromate  |  |  |  |  |
| Step-II:Conversion of sodium chromate into sodium dichromate:   |  |  |  |  |
| $2Na_2CrO_4 + H_2SO_4 \longrightarrow Na_2Cr_2O_7 + Na_2SO_4 + H_2O_4$  |  |  |  |  |
| Sodium Sulpuric acid Sodium Sodium  |  |  |  |  |
| chromate dichromate sulphate  |  |  |  |  |
| The solution is concentrated and cooled, when less soluble sodium sulphate crystallized out. On filteration sodium dichromate separate in the form of filterate.<br><b>Step III:Conversion of sodium dichromate into potassium dichromate:</b><br>Na <sub>2</sub> Cr <sub>2</sub> O <sub>7</sub> + 2KCI $\longrightarrow$ K <sub>2</sub> Cr <sub>2</sub> O <sub>7</sub> + 2NaCl<br>Sodium Potassium Potassium dichromate<br>the solution mixture is cooled. The less soluble potassium dichromate crystallizes out leaving behind sodium chloride solution. |  |  |  |  |
|   |  |  |  |  |
| Unit -9 Coordination compounds  |  |  |  |  |
| Question No. 25) or 26) 3 Marks:  |  |  |  |  |
| 1) Using VBT account for the hybridisation, geometry and magnetic property of [Ni(CN) <sub>4</sub> ] <sup>-2</sup>  |  |  |  |  |
| Ans:<br>Electronic configuration of Ni <sup>+2</sup> : [Ar]3d <sup>8</sup>  |  |  |  |  |
| Orbitals of Ni+2 ion is   |  |  |  |  |
| 3d 4s 4p  |  |  |  |  |
| dsp <sup>2</sup> hybridised orbitals of Ni <sup>+2</sup>  |  |  |  |  |
|   |  |  |  |  |
| 3d dsp <sup>2</sup> hybrids 4p  |  |  |  |  |
|   |  |  |  |  |
| $[Ni(CN)_4]^{-2}$   |  |  |  |  |
|   |  |  |  |  |
| <sup>3d</sup> four pairs of electrons <sup>4p</sup><br>from 4 CN <sup>-</sup> ions  |  |  |  |  |
|   |  |  |  |  |
| Hybridization: dsp <sup>2</sup>   |  |  |  |  |
| Geometry: Square planar   |  |  |  |  |
| Magnetic property: Diamagnetic  |  |  |  |  |
|   |  |  |  |  |
| Dinesh kumar G. Lecturer in Chemistry, Govt. PU College, Adarsh Nagar, Kalaburagi-05.   |  |  |  |  |



### Unit – 10 Haloalkanes and Haloarenes

### Question No. 32 a) 3 Marks:

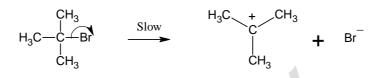
- 1) What is  $S_N 1$  mechanism? Explain  $S_N 1$  mechanism by taking hydrolysis of tertiary butyl bromide as an example.
- Ans : It is unimolecular nucleophilic substitution reaction.

When tertiary butyl bromide is heated with aqueous potassium hydroxide, tertiary butyl alcohol is formed.

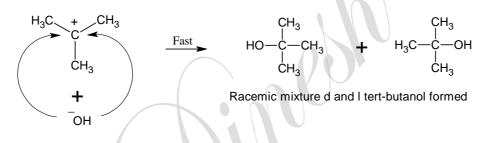


Mechanism : The mechanism involves two steps.

**Step 1 :** Formation of carbocation.



Step 2 : Attack of nucleophile (OH-).



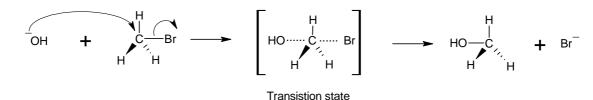
### OR

- 2) What is  $S_N 2$  mechanism? Explain  $S_N 2$  mechanism by taking an example of hydrolysis of methyl bromide.
- Ans : It is bimolecular nucleophilic substitution reaction.

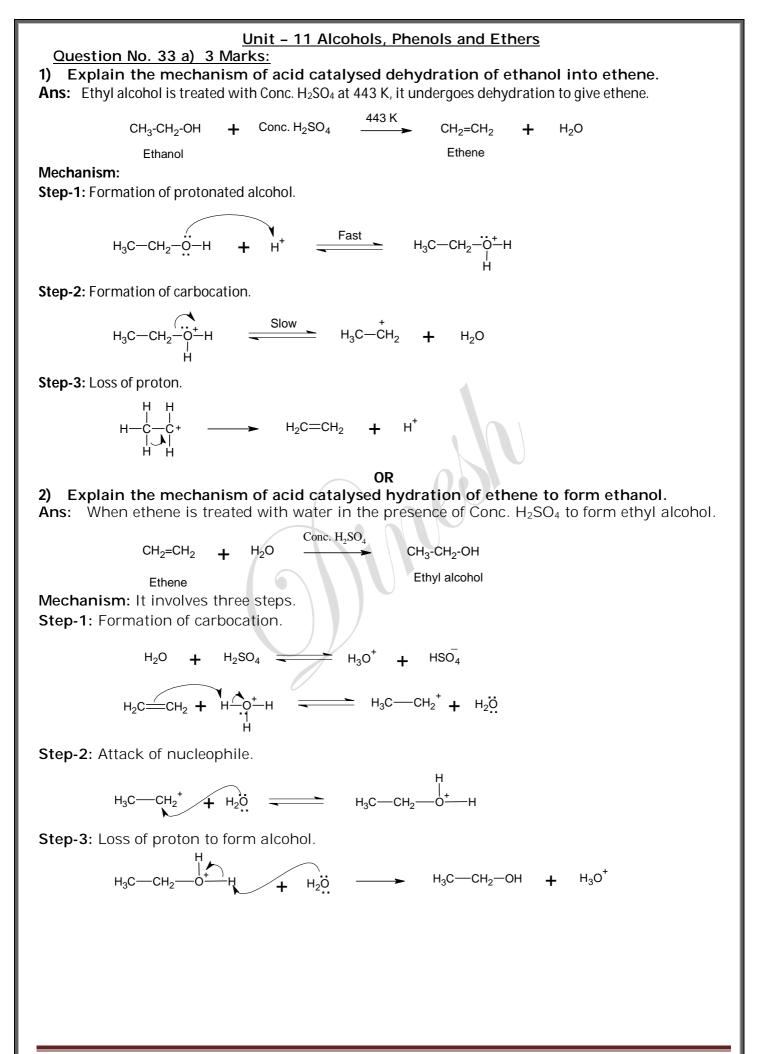
When methyl bromide is heated with aqueous potassium hydroxide, methyl alcohol is formed.

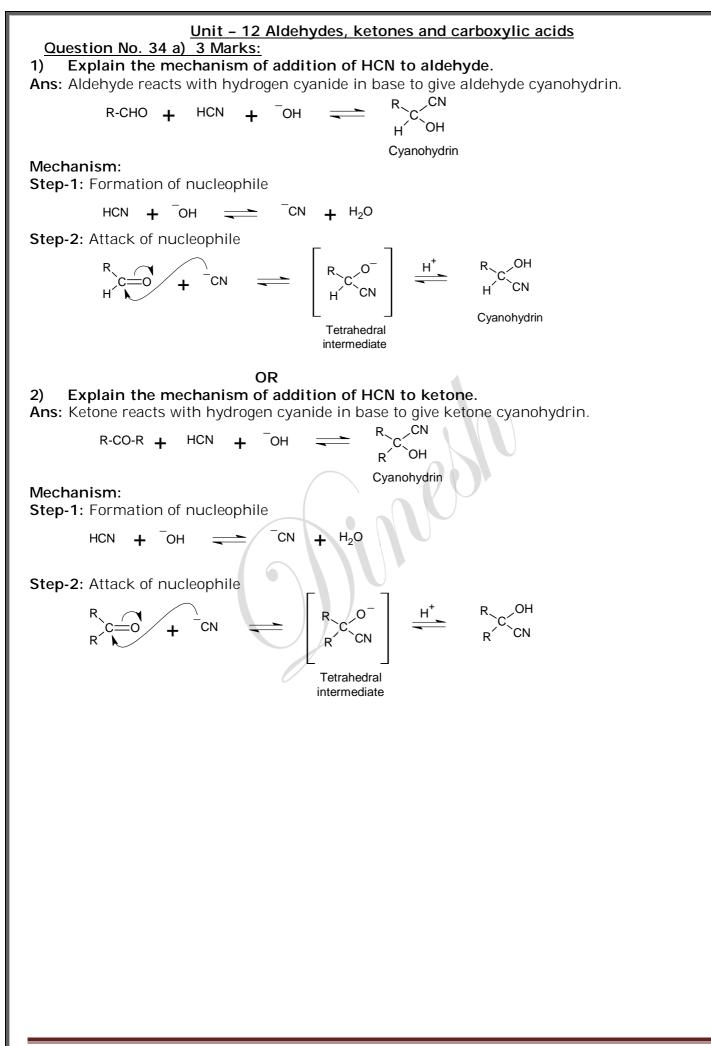
$$CH_3 - Br + KOH \xrightarrow{\Lambda} CH_3 - OH + KBr$$

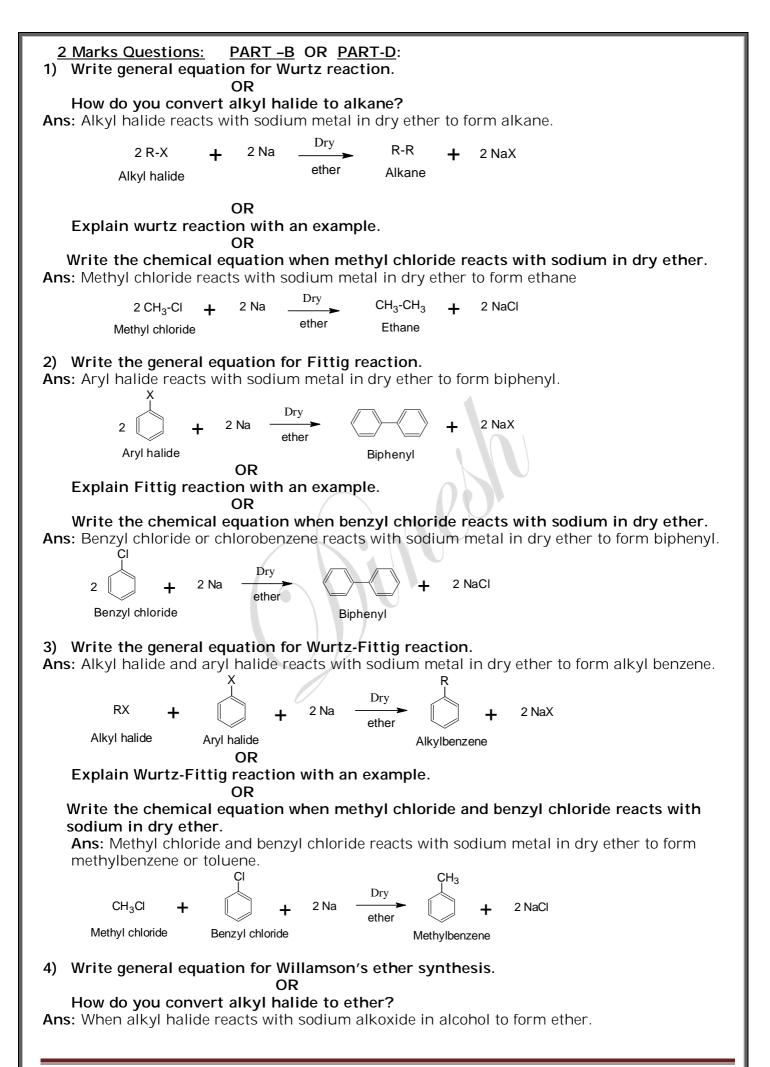
Mechanism : The mechanism involves only one step

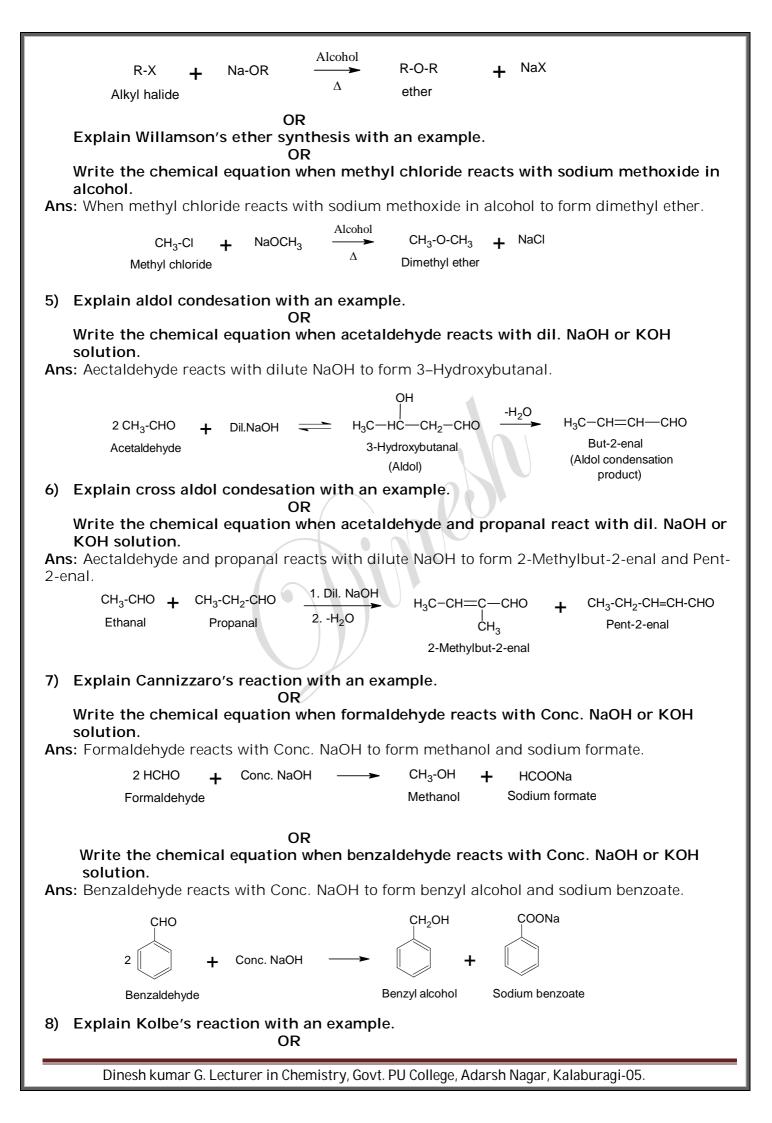


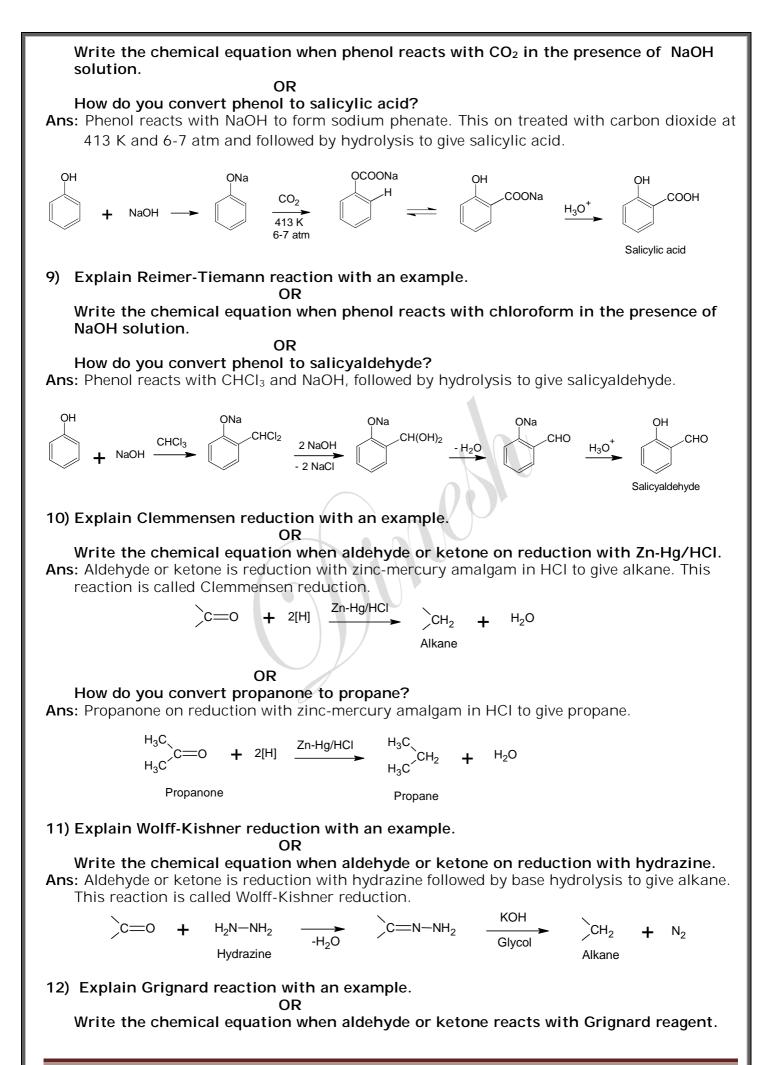
In this mechanism the nucleophile OH- attacks the carbon atom to form a new C – OH bond. Here, there exists a transition state in which C – OH bond is partially formed and C – Br bond is partially broken. Finally the C – OH bond becomes stronger and C – Br bond gets weakened, releases Br- to give end product methyl alcohol.

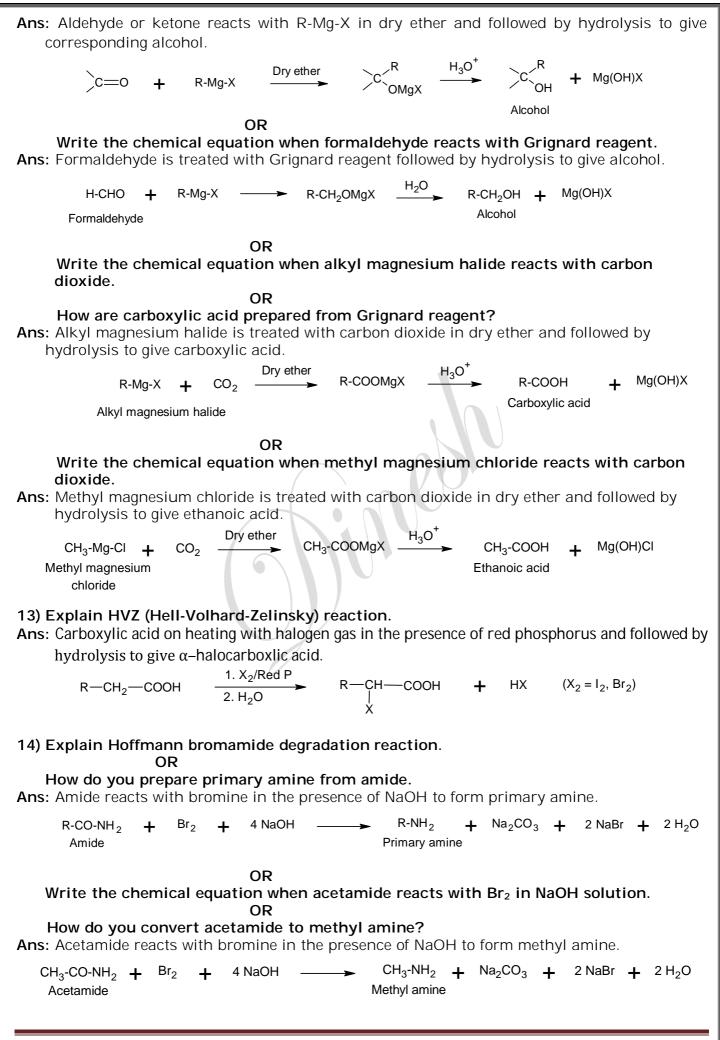












| <b>15) Explain Carbylamine reaction.</b><br><b>Ans:</b> Primary amine reacts with CHCI <sub>3</sub> & KOH to give isocyanide.  |
|--|
| R-NH <sub>2</sub> + CHCl <sub>3</sub> + 3 KOH $\xrightarrow{\Lambda}$ R-NC + 3 KCl + <sup>3</sup> H <sub>2</sub> O<br>Primary amine Isocyanide   |
| OR<br>How do you prepare methyl isocyanide from methyl amine.<br>Ans: Methyl amine reacts with CHCl <sub>3</sub> & KOH to give methyl isocyanide.  |
| $CH_3-NH_2 + CHCI_3 + 3KOH \longrightarrow CH_3-NC + 3KCI + 3H_2O$<br>Methyl amine Methyl isocyanide   |
| 16) Explain Gabriel Phthalimide synthesis with an example.   |
| OR<br>How do you prepare primary amine from Phthalimide.<br>Ans: Phthalimide is treated with alcoholic KOH to give potassium phthalimide. This on treated<br>with alloch ballede and followed by treated with any any and followed by treated with |
| with alkyl halide and followed by treated with aqueous NaOH solution to give primary amine.  |
|  |
| Phthalimide N-Alkylphthalimide   |
| aq. NaOH   |
| R-NH <sub>2</sub> +  |
| Primary amine COONa  |
|  |
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| Dinesh kumar G. Lecturer in Chemistry, Govt. PU College, Adarsh Nagar, Kalaburagi-05.  |