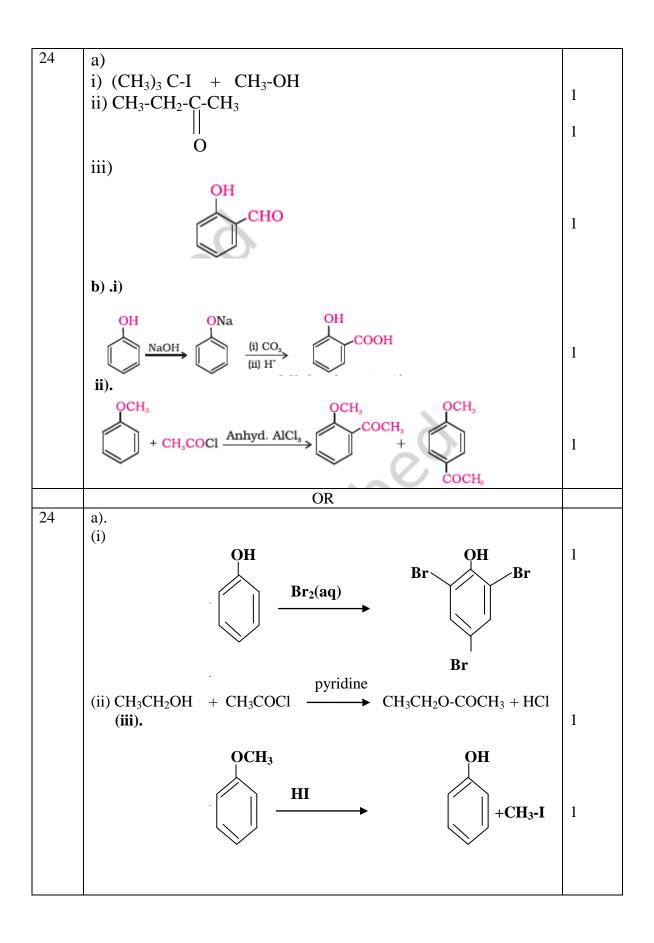
## <u>CHEMISTRY MARKING SCHEME</u> <u>FOREIGN-2016</u> <u>SET -56/2/3/F</u>

| Q.no. | Answers   | Marks |
|-------|---|-------|
| 1     | NO <sub>2</sub> gas   | 1     |
| 2     | N,N-dimethylbutanamide  | 1     |
| 3     | Like Charged particles cause repulsion/ Brownian motion/ solvation  | 1     |
| 4     | Because of some crystallization.  | 1     |
| 5     | Reaction (ii)   | 1     |
| 6     | $X = CH_3 - CO - CH_2 - CH_3 / Butan - 2 - one$   | 1     |
| Ũ     | $Y = CH_3-CH(OH)-CH_2-CH_3 / Butan-2-ol$  | 1     |
| 7     |   | -     |
| ,     | i) ii)  |       |
|       |   | 1+1   |
| 8     | i) $[Co(NH_3)_4Cl_2]Cl$   | 1     |
|       | ii) Tetraamminedichloridocobalt(III) chloride<br>When reaction is completed 99.9%, $[R]_n = [R]_0 - 0.999[R]_0$   | 1     |
| 9     |   |       |
|       | $k = \frac{2.303}{t} \log \frac{[R]_0}{[R]}$  | 1⁄2   |
|       |   |       |
|       | $=\frac{2.303}{t}\log\frac{[R]_0}{[R]_0-0.999[R]_0}=\frac{2.303}{t}\log 10^3$   |       |
|       |   | 1/2   |
|       | t = 6.909/k   |       |
|       | For half-life of the reaction   |       |
|       | $t_{1/2} = 0.693/k$   |       |
|       | $\frac{t}{t_{1/2}} = \frac{6.909}{k} \times \frac{k}{0.693} = 10$   |       |
|       | $k_{1/2} = k = 0.693$   | 1     |
|       | OR  |       |
| 9     | $R \rightarrow P$   |       |
|       | Rate = $\frac{dR}{dt} = kR$   |       |
|       |   |       |
|       | or $\frac{d R}{R} = -kdt$   | 1/2   |
|       | NEW TO AND THE REPORT OF A DESCRIPTION OF<br>A DESCRIPTION OF A DESCRIPTION O |       |
|       | Integrating this equation, we get<br>$\ln [R] = -kt + I$ (4.8)  |       |
|       | Again, I is the constant of integration and its value can be determined   |       |
|       | easily.<br>When $t = 0$ , $R = [R]_0$ , where $[R]_0$ is the initial concentration of the reactant.   |       |
|       | Therefore, equation (4.8) can be written as   |       |
|       | $\ln [R]_o = -k \times O + I$<br>$\ln [R]_o = I$  |       |
|       | Substituting the value of I in equation $(4.8)$   |       |
|       | $\ln[R] = -kt + \ln[R]_0 \tag{4.9}$   | 1⁄2   |
|       | Rearranging this equation   |       |
|       |   |       |
|       | $\ln \frac{R}{R_0} = kt$  |       |
|       | or $k = \frac{1}{t} \ln \frac{[\mathbf{R}]_0}{[\mathbf{R}]}$ $k = \frac{2.303}{t} \log \frac{[\mathbf{R}]_0}{[\mathbf{R}]}$   | 1     |
|       | · [ <sup>1</sup> ] · · · · ·  |       |

| 10 | Henry's law states that the mole fraction of gas in the solution is<br>proportional to the partial pressure of the gas over the solution.  | 1       |
|----|--|---------|
|    | Applications: solubility of $CO_2$ gas in soft drinks /solubility of air diluted with helium in blood used by sea divers or any other  | 1⁄2     |
|    | Solubility of gas in liquid decreases with increase in temperature.  | 1/2     |
| 11 | (i) Butadiene and acrylonitrile<br>$CH_2 = CH - CH = CH_2$ and $CH_2=CH-CN$  | 1/2+1/2 |
|    | (ii) Vinyl chloride  | 1/ 1/   |
|    | (iii) Chloroprene  | 1/2+1/2 |
|    | Cl   |         |
|    |  |         |
|    | $CH_2 = C - CH = CH_2$   | 1/2+1/2 |
| 12 | сн <sub>2</sub> он   | 1       |
|    |  |         |
|    | i) H OH  | 1       |
|    | ii) Peptide linkage / -CO-NH- linkage  | 1       |
|    | iii) Water soluble-Vitamin B / C   | 1/2+1/2 |
| 13 | Fat soluble- Vitamin A /D /E /K  |         |
| 15 | i) $dsp^3$ ,   | 1       |
|    | Diamagnetic, low spin  | 1/2+1/2 |
|    | ii) The energy used to split degenerate d-orbitals due to the  |         |
|    | presence of ligands in a definite geometry is called crystal   | 1       |
| 14 | field splitting energy.i)Iodine is heated with Zr or Ti to form a volatile compound which on   | 1       |
| 14 | further heating decompose to give pure Zr or Ti .  |         |
|    | or   | 1       |
|    | $Zr(impure) + 2I_2 \longrightarrow ZrI_4$<br>(volatile)  |         |
|    | $ZrI_4$ <u>1800K</u> $Zr(pure) + 2I_2$   |         |
|    | <ul> <li>ii)Cryolite lowers the m.p.of alumina mix / acts as a solvent / brings conductivity.</li> <li>(iii) Role of NaCN in the extraction of Ag is to do the leaching of silver</li> </ul> | 1       |
|    | ore in the presence of air.<br>or  |         |
|    | $4Ag(s) + 8CN^{-}(aq) + 2H_2O + O_2(g) \qquad \qquad 4OH^{-} \qquad 4[Ag(CN)_2]^{-} + 4OH^{-}$   | 1       |

| 15 | i)  |         |
|----|---|---------|
| _  | CH <sub>2</sub> Cl  |         |
|    |   |         |
|    |   |         |
|    | но  |         |
|    | ii)   |         |
|    |   |         |
|    | Br  |         |
|    | CH <sub>3</sub>   |         |
|    |   |         |
|    | iii) CH <sub>3</sub> CH <sub>2</sub> ONO  | 1 x 3=3 |
| 16 | $k = 2.303 \log p_i$  |         |
|    | $k = \frac{2.303}{t} \log \frac{p_i}{2p_i - p_t}$   | 1       |
|    | $= \frac{2.303}{300} \log \frac{0.3}{2 \times 0.3 - 0.5}$   |         |
|    | 300 2 x 0.3- 0.5  | 1       |
|    | = <u>2.303</u> log 3  |         |
|    | 300   |         |
|    | $= 2.303 \ge 0.4771$  |         |
|    | 300   |         |
|    | $= 0.0036 \text{ atm}^{-1} \text{ or } 0.004 \text{ atm}^{-1} \text{ (approx.)}$                      | 1       |
| 17 | i)Because of the resonance stabilization of the conjugate base i.e enolate                            |         |
|    | anion or diagrammatic representation.   | 11/2    |
|    | iii)Because the carboxyl group gets bonded to the catalyst anhyd.AlCl <sub>3</sub> (lewis acid).      | 11⁄2    |
|    | ( note: part ii is deleted because of printing error and mark   |         |
|    | alloted in part i and part iii )  |         |
| 17 | $OR$ $i)C_6H_5CH_3 \underline{CrO_3/(CH_3CO)_2O} C_6H_5CH(OCOCH_3)_2 \underline{H_2O}$                |         |
| 1/ | $C_6H_5CHO$   |         |
|    | ii)CH <sub>3</sub> COOH <u>Cl<sub>2</sub>/P</u> Cl-CH <sub>2</sub> -COOH                              |         |
|    |   | 1x3=3   |
|    | iii)CH <sub>3</sub> COCH <sub>3</sub> Zn(Hg)/conc.HCl CH <sub>3</sub> CH <sub>2</sub> CH <sub>3</sub> | _       |
|    | ( Or by any other correct method)   |         |

| 18 | $d = \frac{z \times M}{N_A \times a^3}$   | 1                          |
|----|---|----------------------------|
|    | $N_A x a^3$   |                            |
|    | Or  |                            |
|    |   |                            |
|    | N x a <sup>3</sup>  |                            |
|    | $d = \frac{4 \times 200 \text{ g}}{2.5 \times 10^{24} \text{ x} (400 \times 10^{-10} \text{ cm})^3}$  |                            |
|    | $2.5 \times 10^{-1} \text{ x} (400 \times 10^{-5} \text{ cm})^{\circ}$  | 1                          |
|    | $d = 5 \text{ g cm}^{-3}$   | 1                          |
|    | (or by any other correct method)  |                            |
| 19 | i) It is a process in which both adsorption and absorption can take   | 1                          |
|    | <ul><li>place simultaneously.</li><li>ii) It is the potential difference between the fixed layer and the</li></ul>  | 1                          |
|    | diffused/ double layer of opposite charges around the   | 1                          |
|    | colloidal particles.  |                            |
|    | iii) It is the temperature above which the formation of micelles takes  | 1                          |
| 20 | place.<br>$\Delta T_{\rm f} = i K_{\rm f} m$  | 1/2                        |
| 20 | $\Delta 1_{1}^{*} - \mathbf{n}_{1}^{*} \mathbf{m}$  | /2                         |
|    | For complete ionisation of $Na_2SO_4$ i=3   | 1⁄2                        |
|    |   |                            |
|    | $\Delta T_{\rm f} = T_{\rm f}^{0} - T_{\rm f} = 3 \ \text{x} \ 1.86 \ \text{K kg mol}^{-1} \text{x} \ \frac{2g}{142g \ \text{mol}^{-1}} \ \text{x} \ \frac{1000 \ \text{g kg}^{-1}}{50 \ \text{g}}$ | 1                          |
|    | $\Delta T_{\rm f} = 1.57$   | -                          |
|    |   |                            |
|    | So, $T_f = -1.57^{\circ}C$ or 271.43K<br>i)Because of higher oxidation state (+5) / high charge to size ratio /   | 1                          |
| 21 | i)Because of higher oxidation state (+5) / high charge to size ratio / high polarizing power.   |                            |
|    |   |                            |
|    | ii)Because of high interelectronic repulsion.   |                            |
|    | iii)Because of its low bond dissociation enthalpy and high hydration  | 1x3=3                      |
| 22 | enthalpy of $F^{-}$ .<br>i)A : C <sub>6</sub> H <sub>5</sub> CONH <sub>2</sub> B : C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub> C : C <sub>6</sub> H <sub>5</sub> NHCOCH <sub>3</sub>              | 11/2                       |
|    | ii)A: $C_6H_5NO_2$ B: $C_6H_5NH_2$ C: $C_6H_5NHCOCH_3$<br>ii)A: $C_6H_5NO_2$ B: $C_6H_5NH_2$ C: $C_6H_5-NC$   | $1\frac{72}{1\frac{1}{2}}$ |
| 23 | (i)Caring ,dutiful, Concerned, compassionate ( or any other two   | 1/2+1/2                    |
|    | values)   | 1                          |
|    | ii)Because higher doses may have harmful effects and act as poison<br>which cause even death.   | 1                          |
|    | iii)Tranquilizers are a class of chemical compounds used for treatment  | 1                          |
|    | of stress or even mental diseases.  |                            |
|    | ex. chlordiazepoxide, equanil, veronal, serotonin, valium (or   | 1/2+1/2                    |
|    | any other two examples)   |                            |
|    |   |                            |



| (b)(i) Warm each compound with iodine and sodium hydroxide.               | 1 |
|---|---|
| Phenol : No yellow ppt formed   |   |
| Ethanol: Yellow ppt of Iodoform are formed.                               |   |
| ii)<br>On adding lucas reagent ( $HCl/anhyd.ZnCl_2$ ) , Propan-2-ol gives | 1 |
| white turbidity after 5 minutes whereas 2-methylpropan-2-ol gives         |   |
| white turbidity immediately.  |   |
| (or any other suitable test)  |   |

| 25 | a) Given $E^{o}_{Cell} = +0.30V$ ; $F = 96500C \text{ mol}^{-1}$   |     |
|----|--|-----|
|    | n = 6 (from the given reaction)  |     |
|    | $\Delta_{\rm r} {\rm G}^{\rm O} = - n \ {\rm x} \ {\rm F} \ {\rm x} \ {\rm E}^{\rm o}_{\rm Cell}$  | 1⁄2 |
|    | $\Delta_{\rm r} {\rm G}^{\rm O} = -6 \ {\rm x} \ 96500 \ {\rm C} \ {\rm mol}^{-1} \ {\rm x} \ 0.30 {\rm V}$                                      | 1   |
|    | = - 173,700 J / mol or - 173.7 kJ / mol  |     |
|    | $\log Kc = \underline{n E^{o}_{Cell}}$   | 1/2 |
|    | 0.059  |     |
|    | $\log \mathrm{Kc} = \underline{6 \times 0.30}$   |     |
|    | 0.059<br>log Kg 20.5   | 1   |
|    | $\log \text{ Kc} = 30.5$<br>b)A  | 1   |
|    | Because $E^{\circ}$ value of A shows that on coating ,A acts as anode and Fe   | 1   |
|    | acts as a cathode and hence A oxidises in prefence to Fe and prevent   |     |
|    | corrosion / or $E_{cell}^{o}$ is positive and hence A oxidises itself to prevent   |     |
|    | corrosion of $Fe/E^{\circ}$ value is more negative.  | 1   |
|    | ( or any other correct reason) OR  |     |
| 25 | a) $\Lambda_m = \underline{\kappa}$  | 1/2 |
|    | С  |     |
|    | $= \frac{3.905 \text{ x } 10^{-5} \text{ S cm}^{-1}}{0.001 \text{ mol } \text{L}^{-1}} \text{ x } \frac{1000 \text{ cm}^{3}}{\text{L}}$          |     |
|    |  | 1   |
|    | $\Lambda_{\rm m} = 39.05  {\rm Scm}^2 {\rm mol}^{-1}$<br>$\Lambda_{\rm o} = \lambda^{\rm o}({\rm H}^+) + \lambda^{\rm o}({\rm CH}_3{\rm COO}^-)$ | 1   |
|    | $= (349.6 + 40.9) \text{ Scm}^2 \text{mol}^{-1}$   |     |
|    | $\Lambda_{\rm o} = 390.5 \ \rm Scm^2 mol^{-1}$   |     |
|    | $\alpha = \underline{\Lambda_{m}}$   | 1⁄2 |
|    | $\frac{\Lambda_0}{20.05}  \mathrm{S}  \mathrm{sm}^2 \mathrm{m}  \mathrm{s}^{1-1}$  |     |
|    | $= \frac{39.05 \text{ Scm}^2 \text{mol}^{-1}}{390.5 \text{ Scm}^2 \text{mol}^{-1}}$  |     |
|    | $\alpha = 0.1$   | 1   |
|    |  |     |
|    |  |     |

| b)Secondary battery or rechargeable battery   | 1 |
|---|---|
| $Pb(s) + PbO_{2}(s) + 2SO_{4}^{2-}(aq) + 4H^{+}(aq) \longrightarrow$<br>2PbSO <sub>4</sub> (s) + 2H <sub>2</sub> O(l) | 1 |

Т

| 26 | a)  |                             |
|----|---|-----------------------------|
|    | i)Because of higher oxidation state (+7) of Mn.   | 1                           |
|    | ii)Because it has one unpaired electron in 3d orbital in its +2 oxidation                                 | 1                           |
|    | state / or it has incompletely filled d-orbital in +2 oxidation state.                                    |                             |
|    | iii)Because of comparable energies of 5f, 6d and 7s orbitals.   | 1                           |
|    | b)  | -                           |
|    | $2MnO_2 + 4KOH + O_2 \longrightarrow 2K_2MnO_4 + 2H_2O$   |                             |
|    | $3MnO_4^{2-} + 4 H^+ \longrightarrow 2MnO_4^{-} + MnO_2 + 2H_2O$  | 1+1                         |
|    | OR  |                             |
| 26 | a)  |                             |
|    | i)Cr, because of maximum no. of unpaired electrons cause strong   |                             |
|    | metallic bonding.   | $\frac{1}{2} + \frac{1}{2}$ |
|    | ii)Mn, because it attains stable half -filled $3d^5$ configuration in +2                                  |                             |
|    | oxidation state.  | $\frac{1}{2} + \frac{1}{2}$ |
|    | iii)Zn, because of no unpaired electron in d-orbital.   |                             |
|    | b)  | $\frac{1}{2} + \frac{1}{2}$ |
|    |   |                             |
|    | $2\mathrm{Na_2CrO_4} + 2~\mathrm{H^+} \rightarrow \mathrm{Na_2Cr_2O_7} + 2~\mathrm{Na^+} + \mathrm{H_2O}$ |                             |
|    |   |                             |
|    | $Na_2Cr_2O_7 + 2 KCl \longrightarrow K_2Cr_2O_7 + 2 NaCl$   | 1.1                         |
|    |   | 1+1                         |
|    |   |                             |