## MARKING SCHEME

## SAMPLE PAPER 1

## **SECTION A**

| Q.No. | Value Point | Marks |
|-------|-------------|-------|
| 1(i)  | D           | 1     |
| (ii)  | В           |       |
|       | OR          | 1     |
|       | Α           |       |
| (iii) | Α           | 1     |
| (iv)  | С           | 1     |
| 2(i)  | В           | 1     |
| (ii)  | А           | 1     |
| (iii) | А           | 1     |
| (iv)  | Α           |       |
|       | OR          | 1     |
|       | В           |       |
| 3     | С           | 1     |
| 4     | D           |       |
|       | OR          | 1     |
|       | С           |       |
| 5     | С           | 1     |
| 6     | В           |       |
|       | OR          | 1     |
|       | В           |       |
| 7     | В           |       |
|       | OR          | 1     |
|       | D           |       |
| 8     | Α           |       |
|       | OR          | 1     |
|       | Α           |       |
| 9     | С           | 1     |
| 10    | А           | 1     |
| 11    | Α           | 1     |
| 12    | А           | 1     |
| 13    | D           | 1     |
| 14    | В           |       |
|       | OR          | 1     |
|       | В           |       |
| 15    | В           | 1     |
| 16    | Α           | 1     |

## SECTION B, C, D

| Q.No.     | VALUE POINTS   | MARKS                      |
|-----------|--|----------------------------|
| SECTION B |  |                            |
| 17        | Nitro group at ortho position withdraws the electron density from the benzene ring   | 2                          |
|           | and thus facilitates the attack of the nucleophile on haloarene.   |                            |
|           | $ \begin{array}{c} \begin{array}{c} Cl & 0 \\ \Theta \\ \Theta \\ H + \end{array} \end{array} \xrightarrow{(N \cap O)} \Theta \end{array} \xrightarrow{(N \cap O)} \left[ \begin{array}{c} Cl & O \\ \Theta \\ H \end{array} \right] \left[ \begin{array}{c} Cl \\ H \end{array} \left[ \begin{array}{c} Cl \\ $ |                            |
|           | OR   |                            |
|           | (i) $NH_2$ $N_2Cl$ $Cl$ $Cl$<br>$273 - 278 K$ $Cu_2Cl_2$   | 1                          |
|           | (ii) $CH_3CH(Br)CH_3 \xrightarrow{alc KOH} CH_3CH=CH_2 \xrightarrow{HBr, organic peroxide}$  | 1                          |
| 18        | $\Delta Tb = K_b m  \Delta Tb = 101.04-100 = 1.04 ^{\circ}C$<br>or m= 1.04 /0.52 = 2 m<br>2 m solution means 2 moles of solute in 1 kg of solvent.   | 1                          |
|           | 2 m aq solution of urea means 2 moles of urea in 1kg of water.<br>No. of moles of water = $1000/18 = 55.5$<br>Relative lowering of VP = $x_2$ (where $x_2$ is mole fraction of solute)<br>Relative lowering of VP = $n_2/n_1+n_2$ ( $n_2$ is no. of moles of solute , $n_1$ is no. of moles of solvent)<br>= $2/2+55.5 = 2/57.5 = 0.034$   | 1/2<br>1/2                 |
| 19        | (i)t <sub>2g</sub> <sup>4</sup> e <sub>g</sub> <sup>2</sup> Paramagentic<br>(ii)Dichloridobis(ethane-1,2-diamine)cobalt(III)nitrate<br>OR  | $\frac{1}{2}, \frac{1}{2}$ |
|           | (i)Square planar<br>(ii)Cu <sup>2+</sup> = $3d^9$ 1 unpaired electron so $\sqrt{1(3)} = 1.73$ BM   | 1                          |
| 20        | Reaction is a complex reaction.  | 1/2                        |
|           | Order of reaction is 1.5.  |                            |
|           | Molecularity cannot be 1.5, it has no meaning for this reaction. The reaction occurs   |                            |
|           | in steps, so it is a complex reaction.   | 1/2                        |
|           | (ii)units of k are mol <sup>-<math>1/2</math></sup> L <sup><math>1/2</math></sup> s <sup>-1</sup>  | 1                          |

|    | OR   |               |
|----|--|---------------|
|    | Ans : let the rate law expression be Rate = $k [P]^{x}[Q]^{y}$                                     |               |
|    | from the table we know that  |               |
|    | Rate $1 = 3.0 \times 10^{-4} = k (0.10)^{x} (0.10)^{y}$  |               |
|    | Rate $2 = 9.0 \times 10^{-4} = k (0.30)^{x} (0.30)^{y}$  |               |
|    | Rate $3 = 3.0 \times 10^{-4} = k (0.10)^{x} (0.30)^{y}$  |               |
|    |  |               |
|    |  |               |
|    | Rate 1/ Rate $3 = (1/3)^y$ or $1 = (1/3)^y$  |               |
|    | So $y = 0$   | 1/2           |
|    | Rate 2/ Rate $3 = (3)^x$ or $3 = (3)^x$  |               |
|    | So $x = 1$   | 1/2           |
|    | Rate = $k [P]$   | 1             |
|    |  |               |
| 21 | $k = 0.693/t_{1/2}$  |               |
|    | k = 0.693/5730 years <sup>-1</sup>   | 1/2           |
|    | $t = 2.303 \log \frac{Co}{C}$  |               |
|    | k Ct   | $1/_{2}$      |
|    | let $C_0 = 1$ $Ct = 3/10$ so $C_0/Ct = 1/(3/10) = 10/3$  | 17            |
|    | $t = \frac{2.303}{0.02} \times \frac{5}{30} \log \frac{10}{2}$                                     | 1/2           |
|    | $\begin{array}{c} 0.095 & 5 \\ t = 10042 \text{ w} (1 \ 0.4771) = 0.057 \text{ where} \end{array}$ | 1/            |
|    | t = 19042  x (1-0.4771) = 9937  years  | 72            |
| 22 |  |               |
|    | $(n_3 - cn - cn_3) \longrightarrow (n_3 - cn - cn_3)$  | 1/2           |
|    | CH <sub>3</sub> OH CH <sub>3</sub> OH <sub>2</sub>   | /2            |
|    |  |               |
|    | $CH_2 - CH - CH_2 - CH_2 - CH_2 - CH_2 - CH_2$   | $\frac{1}{2}$ |
|    |  |               |
|    | $CH_3 - QH_2 CH_3$   |               |
|    | $\bigcirc$   |               |
|    |  |               |
|    | $CH_3 - C - CH - CH_3 \xrightarrow{12 - hydride shift} CH_3 - CH_2 - CH_2$                         | 1/2           |
|    | CH <sub>3</sub> CH <sub>3</sub>  |               |
|    |  |               |
|    |  |               |
|    | Br   | • /           |
|    | $CH_3 - C - CH_2 - CH_3 \qquad Br^- \longrightarrow CH_3 - C - CH_2 - CH_3$                        | 1/2           |
|    | CH <sub>3</sub> CH <sub>3</sub>  |               |
|    | active secondar  |               |
| 23 | XeF <sub>6</sub>   | 1             |
|    | Central atom Xe has 8 valence electrons it forms 6 bonds with F and has                            |               |
|    | 1 lone pair According to VSEPR theory presence of 6 hn and 1 ln results in                         | 1             |
|    | distorted octahedral geometry  | 1             |
| 1  | Listerten commonal Beenferg  |               |

|           | F F F F F F F F F F F F F F F F F F F   |                  |
|-----------|---|------------------|
| 24.       | Racemic mixture will be given by 2 chlorobutane as it is an optically active compound.  | 1                |
|           | When 2 chlorobutane undergoes $S_N^1$ reaction, both front and rear attack are possible, resulting in a racemic mixture   | 1                |
| 25        | Let no. of Atoms of element P be x  |                  |
|           | No. of tetrahedral voids = $2x$   | 1/2              |
|           | No. Of octahedral voids = $x$   |                  |
|           | Atoms of Q = $1/3 (2x) + x = 5x/3$  | 1/2              |
|           |   | 1                |
|           | $P_3Q_5$  | 1                |
| SECTION C |   |                  |
| 26        | <ul> <li>(i)Due to large surface area and ability to show variable oxidation states</li> <li>(ii)Due to high value of third ionisation enthalpy</li> <li>(iii) Mo(VI) and W(VI) are more stable than Cr(VI).</li> <li>OR <ul> <li>(i) The general trend towards less negative <i>E</i><sup>o</sup> V values across the series is related to the general increase in the sum of the first and second ionisation enthalpies.</li> <li>(ii) The high energy to transform Cu(s) to Cu<sup>2+</sup> (aq) is not balanced by its hydration enthalpy.</li> <li>(iii) The stability of the half-filled <i>d</i> sub-shell in Mn<sup>2+</sup> and the completely filled <i>d</i><sup>10</sup> configuration in Zn<sup>2+</sup> are related to their more negative <i>E</i><sup>o</sup> V values</li> </ul> </li> </ul> | 1<br>1<br>1<br>1 |
| 27        | <ul> <li>(i) Aniline, <i>N</i>-ethylethanamine, Etanamine</li> <li>(ii)Ethanamine, ethanol, ethanoic acid</li> <li>(iii) N, N dimethylmethanamine, methanamine, N-methylmethanamine</li> </ul>  | 1<br>1<br>1      |
|           | (1) N-methyletahnamine is a secondary amine. When it reacts with benzenesulphonyl chloride, it forms N- Ethyl -N methyl sulphonamide while and  | 1                |

|     | N,N-dimethyl etahnanmine is a tertiary amine it does not react with  |                      |
|-----|--|----------------------|
|     | benzenesulphonyl chloride.<br>(ii) NO <sub>2</sub> NH <sub>2</sub> NH <sub>2</sub><br>$H_2/N_1$ $Br$ $Br$ $Br$<br>(iii)Butan-1-ol  | 1                    |
|     | Alcohol forms stronger hydrogen bonds with water than formed by amine due to<br>higher electronegativity of O in alcohol than N in amine   | 1/2<br>1/2           |
| 28  | We know that $d = zM/N_a a^3$<br>For fcc, $z=4$ therefore $d = 4 \times M / Na (3.5 \times 10^{-8})^3 \text{ g/cm}^3$<br>For bcc, $z=2$ therefore $d' = 2 \times M / Na (3.0 \times 10^{-8})^3 \text{ g/cm}^3$<br>$d/d' = 4/(3.5 \times 10^{-8})^3 / 2/(3.0 \times 10^{-8})^3 = 1.26:1$  | 1/2<br>1<br>1<br>1/2 |
| 29  | $ \begin{array}{c ccccccccccccccccccccccccccccccccccc$   | 1                    |
|     | $\begin{vmatrix} CH_2COOH & CH_3 & (CH_2)_4 - NH_2 \\   HOOC - CH - N - C - CH - N - C - CH - NH_2 \\   H & O & H & O \end{vmatrix}$   | 1                    |
|     | (ii)<br>$H_{3}N - C - COO^{-}$<br>$CH_{3}$   | 1                    |
| 30  | <ul> <li>i. Arrange the following in decreasing order of bond dissociation enthalpy<br/>Cl<sub>2</sub> &gt; Br<sub>2</sub> &gt; F<sub>2</sub> &gt; I<sub>2</sub></li> <li>ii. Bi does not form pπ-pπ bonds as its atomic orbitals are large and diffuse so<br/>effective overlapping is not possible</li> <li>iii. Due to small size of oxygen, it has greater electron electron repulsions</li> </ul> | 1                    |
|     | SECTION D  | 1                    |
| 31. | (i)<br>(a) $3Cu + 8$ HNO <sub>3</sub> (dilute) $\rightarrow 3Cu(NO_3)_2 + 2NO + 4H_2O$<br>(b)  | 1                    |





