

# ANSWER KEY

## AITS FINAL TRACK

### PART TEST-06

#### PHYSICS

##### SECTION-A

Q.1 (3)	Q.2 (4)	Q.3 (2)	Q.4 (4)	Q.5 (3)	Q.6 (1)	Q.7 (2)	Q.8 (4)	Q.9 (2)	Q.10 (3)
Q.11 (1)	Q.12 (4)	Q.13 (1)	Q.14 (2)	Q.15 (2)	Q.16 (3)	Q.17 (4)	Q.18 (2)	Q.19 (2)	Q.20 (3)
Q.21 (4)	Q.22 (3)	Q.23 (4)	Q.24 (3)	Q.25 (1)	Q.26 (3)	Q.27 (1)	Q.28 (3)	Q.29 (2)	Q.30 (2)
Q.31 (2)	Q.32 (4)	Q.33 (2)	Q.34 (4)	Q.35 (1)					

##### SECTION-B

Q.36 (3)	Q.37 (3)	Q.38 (1)	Q.39 (1)	Q.40 (2)	Q.41 (2)	Q.42 (1)	Q.43 (4)	Q.44 (3)	Q.45 (2)
Q.46 (2)	Q.47 (1)	Q.48 (2)	Q.49 (3)	Q.50 (2)					

#### CHEMISTRY

##### SECTION-A

Q.51 (1)	Q.52 (2)	Q.53 (2)	Q.54 (1)	Q.55 (1)	Q.56 (3)	Q.57 (2)	Q.58 (4)	Q.59 (1)	Q.60 (3)
Q.61 (4)	Q.62 (4)	Q.63 (3)	Q.64 (4)	Q.65 (2)	Q.66 (3)	Q.67 (1)	Q.68 (3)	Q.69 (1)	Q.70 (2)
Q.71 (1)	Q.72 (2)	Q.73 (2)	Q.74 (1)	Q.75 (3)	Q.76 (2)	Q.77 (3)	Q.78 (4)	Q.79 (4)	Q.80 (2)
Q.81 (3)	Q.82 (1)	Q.83 (1)	Q.84 (3)	Q.85 (3)					

##### SECTION-B

Q.86 (2)	Q.87 (2)	Q.88 (4)	Q.89 (4)	Q.90 (2)	Q.91 (3)	Q.92 (4)	Q.93 (2)	Q.94 (1)	Q.95 (2)
Q.96 (1)	Q.97 (1)	Q.98 (2)	Q.99 (2)	Q.100 (3)					

#### BIOLOGY-I

##### SECTION-A

Q.101 (1)	Q.102 (4)	Q.103 (1)	Q.104 (1)	Q.105 (3)	Q.106 (4)	Q.107 (3)	Q.108 (2)	Q.109 (1)	Q.110 (3)
Q.111 (2)	Q.112 (1)	Q.113 (2)	Q.114 (1)	Q.115 (2)	Q.116 (2)	Q.117 (4)	Q.118 (1)	Q.119 (3)	Q.120 (3)
Q.121 (2)	Q.122 (3)	Q.123 (3)	Q.124 (1)	Q.125 (4)	Q.126 (3)	Q.127 (4)	Q.128 (2)	Q.129 (4)	Q.130 (2)
Q.131 (3)	Q.132 (3)	Q.133 (3)	Q.134 (1)	Q.135 (3)					

##### SECTION-B

Q.136 (3)	Q.137 (4)	Q.138 (2)	Q.139 (2)	Q.140 (2)	Q.141 (4)	Q.142 (4)	Q.143 (4)	Q.144 (3)	Q.145 (4)
Q.146 (4)	Q.147 (3)	Q.148 (3)	Q.149 (2)	Q.150 (2)					

#### BIOLOGY-II

##### SECTION-A

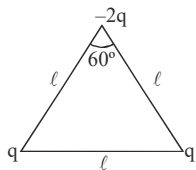
Q.151 (4)	Q.152 (4)	Q.153 (4)	Q.154 (1)	Q.155 (3)	Q.156 (1)	Q.157 (4)	Q.158 (2)	Q.159 (3)	Q.160 (4)
Q.161 (2)	Q.162 (3)	Q.163 (3)	Q.164 (1)	Q.165 (2)	Q.166 (4)	Q.167 (3)	Q.168 (3)	Q.169 (3)	Q.170 (4)
Q.171 (2)	Q.172 (1)	Q.173 (2)	Q.174 (3)	Q.175 (1)	Q.176 (4)	Q.177 (1)	Q.178 (3)	Q.179 (4)	Q.180 (3)
Q.181 (4)	Q.182 (4)	Q.183 (2)	Q.184 (4)	Q.185 (1)					

##### SECTION-B

Q.186 (2)	Q.187 (2)	Q.188 (2)	Q.189 (2)	Q.190 (3)	Q.191 (2)	Q.192 (1)	Q.193 (1)	Q.194 (3)	Q.195 (2)
Q.196 (4)	Q.197 (3)	Q.198 (2)	Q.199 (1)	Q.200 (1)					

**PHYSICS**  
**SECTION-A**

**Q.1** (3)  
According to question



It is a combination of two dipoles at  $60^\circ$

So

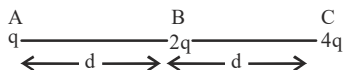
$$P_{\text{net}} = \sqrt{3}p$$

where  $p = ql$

$$\text{So } P_{\text{net}} = \sqrt{3}ql$$

**Q.2** (4)  
During charging by conduction, both the objects acquire same type of charge. Finally both rod & sphere becomes negatively charged & in order for the neutral sphere to become negative, it must gain electron.

**Q.3** (2)



$$F_A = \frac{k(q)(2q)}{d^2} + \frac{k(4q)(q)}{(2d)^2}; F_A = \frac{3kq^2}{d^2}$$

$$F_C = \frac{k(4q)(q)}{(2d)^2} + \frac{k(4q)(q)}{d^2}$$

$$F_C = \frac{9kq^2}{d^2}, \quad \frac{F_A}{F_C} = \frac{T_{AB}}{T_{BC}} = \frac{1}{3} \Rightarrow 1:3$$

**Q.4** (4)  
• When there is no net charge resides inside any closed surface then only net electric flux linked with the surface is zero.  
• Electric field due to an electric dipole is non uniform

**Q.5** (3)

$$\phi_{\text{BCGF}} = \phi_{\text{due to } q} + \phi_{\text{due to } 3q} + \phi_{\text{due to } 2q}$$

$$\phi_{\text{due to } q} = \frac{q}{24\epsilon_0}$$

$$\phi_{\text{due to } 3q} = \frac{3q}{24\epsilon_0}$$

$$\phi_{\text{due to } 2q} = 0$$

$$\phi_{\text{BCGF}} = \frac{q}{24\epsilon_0} + \frac{3q}{24\epsilon_0} + 0 = \frac{4q}{24\epsilon_0} = \frac{q}{6\epsilon_0}$$

**Q.6** (1)

$$\therefore E = \frac{KQz}{(R^2 + z^2)^{3/2}}$$

$$\therefore \frac{E_1}{E_2} = \frac{R}{(R^2 + R^2)^{3/2}} \times \frac{(R^2 + 4R^2)^{3/2}}{2R} = \frac{5\sqrt{5}}{4\sqrt{2}}$$

**Q.7** (2)

$$F = \frac{1}{4\pi\epsilon_0} \frac{q_1q_2}{r^2}$$

$$\epsilon_0 = \frac{q_1q_2}{4\pi Fr^2}$$

Unit of  $\epsilon_0$  is  $C^2/N\cdot m^2$

**Q.8** (4)  
When the Gaussian surface encloses no charge, then the electric flux through that surface is zero. Electric field over the Gaussian surface need not be zero at every point on that surface.

**Q.9** (2)

$$\text{For balance } mg = eE \Rightarrow E = \frac{mg}{e}$$

$$\text{Also } m = \frac{4}{3}\pi r^3 d = \frac{4}{3} \times \frac{22}{7} \times (10^{-7})^3 \times 1000 \text{ kg}$$

$$\Rightarrow E = \frac{\frac{4}{3} \times \frac{22}{7} \times (10^{-7})^3 \times 1000 \times 10}{1.6 \times 10^{-19}}$$

$$= 260 \text{ N/C}$$

( $g = 10 \text{ newton/kg}$ ,  $e = 1.6 \times 10^{-19} \text{ coulomb}$ )

**Q.10** (3)

$$E = \frac{Q}{A\epsilon_0\epsilon_r} \Rightarrow E \propto \frac{1}{\epsilon_r}$$

**Q.11** (1)  
Field lines of  $q_1$  passes through surface of hemisphere one time.  
Field lines of  $q_2$  passes through surface of hemisphere two time so net flux due to  $q_2$  is zero.  
Net flux due to  $q_1$  is non zero.

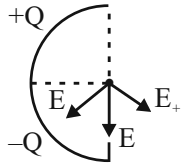
**Q.12** (4)

$$\vec{E}_P = \vec{E}_A + \vec{E}_B$$

$$= +\frac{\sigma}{2\epsilon_0} - \frac{\sigma}{2\epsilon_0} \text{ (both are in opposite direction)}$$

$$= 0$$

**Q.13** (1)



**Q.14** (2)

$$E_{\text{axis}} = \frac{2kP}{r^3}$$

$$E_{\text{eq.}} = \frac{kP}{r^3}$$

**Q.15** (2)

Maximum torque =  $pE$   
 $= 2 \times 10^{-6} \times 3 \times 10^{-2} \times 2 \times 10^5 = 12 \times 10^{-3} \text{ N-m}$

**Q.16** (3)

$$E = \frac{1}{4\pi\epsilon_0} \frac{Q}{r^2}$$

$$Q = 4\pi\epsilon_0 Er^2$$

Now, Surface charge density -

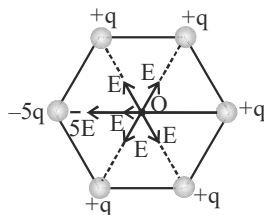
$$\sigma = \frac{Q}{4\pi R^2}$$

$$\sigma = \frac{4\pi\epsilon_0 Er^2}{4\pi R^2}$$

$$\sigma = \epsilon_0 E \left( \frac{r}{R} \right)^2$$

**Q.17** (4)

To obtained net field  $6E$  at centre  $O$ , the charge to be placed at remaining sixth corner is  $-5q$ . (see following figure)



**Q.18** (2)

$\vec{E}$  is maximum at sharp corners while entire conductor is equipotential

**Q.19** (2)

Torque  $\vec{\tau} = \vec{p} \times \vec{E} = pE \sin \theta$

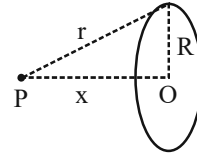
or  $4 = p \times 2 \times 10^5 \sin 30^\circ$

$$\text{or } p = \frac{4}{2 \times 10^5 \times \sin 30^\circ} = 4 \times 10^{-5} \text{ C-m}$$

Dipole moment,  $p = q\ell$

$$\therefore q = \frac{p}{\ell} = \frac{4 \times 10^{-5}}{0.02} = 2 \times 10^{-3} \text{ C} = 2 \text{ mC}$$

**Q.20** (3)



$$E = \frac{kQx}{(R^2 + x^2)^{3/2}}$$

$$r^2 = R^2 + x^2$$

$$x^2 = r^2 - R^2$$

**Q.21** (4)

Potential energy

$$U = -\vec{p} \cdot \vec{E} = -(2\hat{i} - 3\hat{j} + 4\hat{k}) \cdot (5\hat{i} + 2\hat{j} - 3\hat{k})$$

$$= -(10 - 6 - 12) = 8 \text{ joule}$$

**Q.22** (3)

$$V_A - V_B = \left( \frac{kQ}{a} + \frac{k(-Q)}{a+d} \right) - \left( \frac{k(-Q)}{a} + \frac{kQ}{a+d} \right)$$

$$= \frac{2kQd}{a(a+d)}$$

**Q.23** (4)

$$V = -\frac{dV}{dx} = -(4x)\hat{i}$$

$$V = -4(2) = -8\hat{i}$$

**Q.24** (3)

$$V' = n^{2/3} V = 8^{2/3} \times 2 \text{ volt}$$

$$= 4 \times 2 \text{ volt} = 8 \text{ volt}$$

**Q.25** (1)

$$\frac{(K_f)_Q}{(K_f)_{2Q}} = \frac{4m}{m} = \frac{4}{1}$$

$$\Rightarrow (K_f)_Q = \frac{4}{5} \left[ \frac{k(2Q)(Q)}{r} \right] = \frac{8kQ^2}{5r}$$

**Q.26** (3)

$$W_{\text{ext}} = q[V_f - V_i]$$

$$= (2\mu\text{C})\{(-5\text{V}) - (+10\text{V})\}$$

$$= -30 \mu\text{J}$$

**Q.27** (1)

$$U_{\text{system}} = \frac{kQq}{\ell} + \frac{kq^2}{\ell} + \frac{kQq}{\ell} = 0$$

$$\Rightarrow 2Qq = -q^2 \Rightarrow Q = -\frac{q}{2}$$

**Q.28** (3)

$$\Delta V_{\text{max}} = Er = 10 \times 3 = 30 \text{ volt}$$

$$\Delta V = Er \cos \theta = 25 \text{ volt}$$

**Q.29** (2)

Electrostatic field is always perpendicular to the surface of conductor.

**Q.30** (2)

$$V_B - \frac{q}{2} - 12 - \frac{q}{4} + 24 = V_A$$

$$\frac{3q}{4} = 12$$

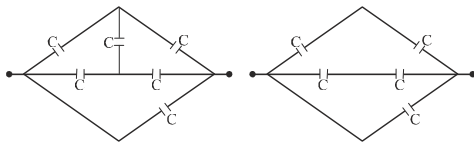
$$q = 16 \mu\text{C}$$

$$V_B - \frac{16}{2} = V_A$$

$$V_B - V_A = 8 \text{ V}$$

**Q.31** (2)

Equivalent circuit diagram

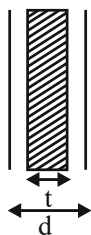


**Q.32** (4)

$$f = Q \epsilon = Q \left( \frac{Q}{2A \epsilon} \right) = \frac{Q^2}{2A \epsilon}$$

**Q.33** (2)

When a dielectric of thickness  $t$  is inserted b/w the plates



$$C = \frac{A \epsilon_0}{d - t + \frac{t}{k}}$$

For metal  $k = \infty$

$$\therefore C = \frac{A \epsilon_0}{d - t}$$

For given question  $t = \frac{d}{2}$

$$\therefore C = 2 \left( \frac{A \epsilon_0}{d} \right) \text{ (Be comes double)}$$

**Q.34** (4)

$$(V_B - V_A) \times 2\mu + (V_B - V_A) \times 3\mu = 0$$

$$(V_B - 1000) \times 2 + (V_B - 0) \times 3 = 0$$

$$2V_B - 2000 + 3V_B = 0$$

$$5V_B = 2000$$

$$V_B = 400 \text{ volt}$$

**Q.35** (1)

As the dielectric slab is pulled out, the equivalent capacity of the system decreases and hence charge supplied by battery decreases as potential of system remains constant. It means charging of battery takes place and a positive charge flows from a to b.

### SECTION-B

**Q.36** (3)

$$E = \frac{2k\lambda}{a} \sin\left(\frac{\alpha}{2}\right); \lambda = \frac{Q}{\left(\frac{\pi a}{2}\right)}; \alpha = 90^\circ$$

$$E = \frac{4k}{a} \frac{Q}{\pi a} \times \sin(45^\circ) = \frac{2\sqrt{2}kQ}{\pi a^2}$$

**Q.37** (3)

$$E_1 = \frac{KQ}{R^2}; E_2 = \frac{K(2Q)}{R^2};$$

$$E_3 = \frac{K(4Q)}{(2R)^3} \times R = \frac{KQ}{2R^2}$$

$$E_2 > E_1 > E_3$$

**Q.38** (1)

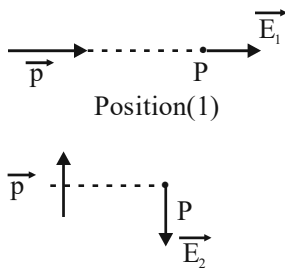
$$F = \frac{kQ^2}{R^2} \dots(1)$$

$$F_2 = \frac{k}{R^2} \left( Q - \frac{3}{4}Q \right) \left( Q + \frac{3}{4}Q \right) = \frac{7}{16} \frac{kQ^2}{R^2} \dots(2)$$

By (1) & (2)

$$F_2 = \frac{7}{16} F$$

Q.39 (1)



Q.40 (2)

$$W = \Delta K$$

$$qE \frac{l}{2} = \frac{1}{2} mv^2$$

$$v = \sqrt{\frac{qEl}{m}}$$

$$T - qE = \frac{mv^2}{l}$$

$$T = 2qE$$

Q.41 (2)

$$V_A - V_B = -\vec{E} \cdot \int_B^A d\vec{r} = -\vec{E} \cdot [\vec{r}_A - \vec{r}_B]$$

$$= \vec{E} \cdot (\vec{r}_B - \vec{r}_A) = 50\sqrt{2} \left[ \frac{\hat{i} + \hat{j}}{\sqrt{2}} \right] \cdot (-4\hat{i} - 2\hat{j})$$

$$= -300 \text{ volt}$$

Q.42 (1)

Factual.

Q.43 (4)

$$\text{Here, } V_A = V_B = V_C \\ \therefore W_1 = W_2 = 0$$

Q.44 (3)

Sol. 
$$U_i = -\frac{kq^2}{a} - \frac{kq^2}{a} + \frac{kq^2}{2a} = -\frac{3}{2} \frac{kq^2}{a}$$

$$U_f = +\frac{kq^2}{a} - \frac{kq^2}{a} - \frac{kq^2}{2a} = -\frac{1}{2} \frac{kq^2}{a}$$

$$\Delta U = U_f - U_i = \frac{kq^2}{a} = \frac{q^2}{4\pi\epsilon_0 a} = W.D$$

Q.45 (2)

$$U = -PE \cos \theta$$

Q.46 (2)

let V in potential difference of battery

$$C_{123} = 1 \mu F, q_{123} = V \mu C$$

$$q_2 = V \mu C, q_4 = CV = 4 \mu C$$

$$\frac{q_2}{q_4} = \frac{1}{4}$$

Q.47 (1)

$$\text{Energy required} = CV^2 = A \epsilon_0 E^2 d$$

$$= \frac{A \epsilon_0}{d} (\epsilon d)^2 = A \epsilon_0 \epsilon^2 d$$

$$= 25 \times 10^{-4} \times 8.85 \times 10^{-12} (600^2 \times 2.0 \times 10^{-3})$$

$$= 1.59 \times 10^{-11} = 15.9 \times 10^{-12} \text{ J}$$

Q.48 (2)

The Wheatstone bridge is in the balanced condition.

$$C_{\text{eff}} = 6 \mu F$$

$$\text{Total energy stored, } U = \frac{1}{2} C_{\text{eff}} V^2$$

$$= \frac{1}{2} \times [6 \mu] \times 2^2 = 12 \mu \text{ J}$$

Q.49 (3)

$$C_1 = \frac{\epsilon_0 A}{d} \text{ and } C_2 = \frac{K \epsilon_0 A}{2d}$$

$$\Rightarrow \frac{C_2}{C_1} = \frac{K}{2} \Rightarrow \frac{40 \times 10^{-12}}{10 \times 10^{-12}} = \frac{K}{2} \Rightarrow K = 8$$

Q.50 (2)

$$\text{Energy required} = CV^2 = \frac{A \epsilon}{d} (Ed)^2 = A \epsilon E^2 d$$

$$= A \epsilon \left( \frac{\sigma}{\epsilon} \right)^2 d$$

$$= \frac{A \sigma^2 d}{\epsilon}$$

## CHEMISTRY

### SECTION-A

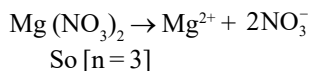
Q.51 (1)

Maximum boiling azeotrope show large negative deviation from Rault's law.

Q.52 (2)

Both statement I and statement II are correct.

**Q.53** (2)



$$\text{Degree of dissociation } \alpha = \frac{i-1}{n-1}$$

$$\therefore i = 2.74$$

$$\text{So } \alpha = \frac{2.74-1}{3-1} = \frac{1.74}{2} = 0.87$$

$$= 0.87 \times 100 = 87\%$$

**Q.54** (1)

The inter particle forces in between chloroform and acetone increases due to H-bonding and thus  $\Delta_{\text{mix}} V$  becomes negative so the resulting solution will have volume less than 70 mL.

**Q.55** (1)

Because its non volatile nature and high concentration and does not dissociate in water that will not form ions in the solution.

**Q.56** (3)

$$\frac{P^\circ - P_s}{P_s} = \frac{WM}{m \times w}$$

Given  $m = 40$   
 $w = 114 \text{ g}$   
 $M_{\text{octane}} = 114$

$$\therefore \frac{100-80}{80} = \frac{w \times 114}{40 \times 114}$$

$$[w = 10 \text{ g}]$$

**Q.57** (2)

$$\text{Concentration in ppm} = \frac{\text{mass of solute}}{\text{mass of solution}} \times 10^6$$

$$= \frac{1.04 \text{ g}}{10^5 \text{ g}} \times 10^6 = 10.4 \text{ ppm}$$

**Q.58** (4)

The entropy change of mixture of A and B is zero because the entropy of mixing for an ideal solution is positive as randomness or disorder increases.

**Q.59** (1)

$$\frac{P^\circ - P_s}{P^\circ} = \frac{w \times M}{m \times w}$$

$$= \frac{0.850 \text{ bar} - 0.845 \text{ bar}}{0.850 \text{ bar}} = \frac{0.5 \text{ g} \times 78 \text{ g/mol}}{39 \text{ g} \times M_2}$$

$$M_2 = 170 \text{ g mol}^{-1}$$

**Q.60** (3)

(1) Molality  $\therefore M$  of KI =  $39 + 127 = 166 \text{ g mol}^{-1}$

$$m = \frac{w \times 1000}{M \times W_{\text{gm}}} = \frac{20 \times 1000}{166 \times 80} = 1.5 \text{ m}$$

$$V = \frac{\text{Mass}}{\text{Density}} = \frac{100}{1.25} = 80 \text{ ml}$$

$$(2) M = \frac{w \times 1000}{M \times V(\text{ml})} = \frac{20 \times 1000}{166 \times 80 \text{ ml}} = 1.5 \text{ M}$$

Mole fraction of KI :-

$$X_B = \frac{n_B}{n_A + n_B} = \frac{\frac{20}{166}}{\frac{80}{18} + \frac{20}{166}} = \frac{0.12}{0.12 + 4.44} = 0.0263$$

**Q.61** (4)

$$P_T = P_A^\circ X_A + P_B^\circ X_B$$

$$X_A + X_B = 1$$

$$(X_B = 1 - X_A)$$

$$760 = 520 X_A + 1000(1 - X_A)$$

$$480 X_A = 240$$

$$X_A = \frac{1}{2} \text{ or } 50\%$$

**Q.62** (4)

$$i = 1 + (n-1) \times \alpha$$

$$i = 1 + (4-1) \times \frac{90}{100}$$

$$i = 3.7$$

$$\Delta T_b = i K_b m = 3.7 \times 0.52 \times 0.2 = 0.3848$$

$$\therefore T_b = T_b^\circ + 0.3848$$

$$= 373 + 0.38 = 373.38 \text{ K.}$$

**Q.63** (3)

Molarity and Normality increase with temperature % w/v also increase with temp. As they are volume related.

**Q.64** (4)

$$T_b^\circ = (\text{B.P.})_{\text{H}_2\text{O}} = 100^\circ\text{C} \quad T_b = 100.25^\circ\text{C} \text{ (for solution)}$$

$$\Delta T_b = T_b - T_b^\circ = 0.25$$

$$\Delta T_b = K_b \cdot m$$

$$0.25 = K_b \cdot \frac{1}{60 \times w} \quad \dots(1)$$

$$\Delta T_b = K_b \times \frac{3}{180 \times w} \quad \dots(2)$$

$$\frac{\text{eq}(1)}{\text{eq}(2)} = [\Delta T_b = 0.25]$$

$$T_b \text{ of glucose} = 100.25^\circ\text{C}$$

**Q.65** (2)  
 $\Delta V_{\text{mix}} > 0, \Delta S_{\text{mix}} > 0$

**Q.66** (3)  
Molarity of cane sugar =  $\frac{5\text{g}}{342 \times 100\text{mL}} \times \frac{1000}{1\text{L}}$   
= 0.146 m .....(1)

Molarity of substance X  
=  $\frac{1\text{g}}{M \times 100} \times \frac{1000}{1\text{L}} = \frac{10}{M}$  molarity .....(2)

But (1) = (2)

$\frac{10}{M} = 0.146 \quad \therefore M = \frac{10}{0.146} = 68.4$

**Q.67** (1)  
Molarity =  $\frac{M_1V_1 + M_2V_2}{V_1 + V_2} = \frac{750 \times 0.5 + 250 \times 2}{1000}$   
= 0.875 M.

**Q.68** (3)  
Raoult's law.

**Q.69** (1)  
 $\Lambda_m = \frac{K \times 1000}{M} = \frac{9.2 \times 10^{-2}}{0.02} = 4.6$

**Q.70** (2)  
For  $\text{CaCl}_2$   
 $\lambda_0 = \lambda_{\text{Ca}^{2+}}^0 + 2\lambda_{\text{Cl}^-}^0$   
=  $271.6 \text{ s cm}^2 \text{ mol}^{-1}$   
For  $\text{MgSO}_4$   
 $\lambda_0 = \lambda_{\text{Mg}^{2+}}^0 + 2\lambda_{\text{SO}_4^{2-}}^0$   
=  $266 \text{ s cm}^2 \text{ mol}^{-1}$

**Q.71** (1)  
 $K = G^* \times G$  (Where  $G^* \rightarrow$  Cell constant  
if  $[K=G]$   $G \rightarrow$  Conductance)  
 $\left[ \frac{K}{G} = 1 \right]$  So cell constant = 1

**Q.72** (2)  
Reason :- Because electrons flow takes place from anode to cathode

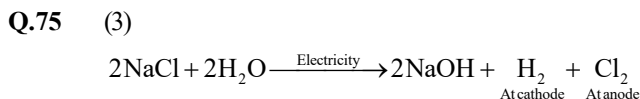
**Q.73** (2)  
Reason :- Cu is anode So it show oxidation lose  $e^-$  and cathode is Ag which gain  $e^-$  and show reduction.

**Q.74** (1)  
 $\text{Tl}^{3+} + 2e^- \rightarrow \text{Tl}^+ = nFE^\circ = 2 \times 1.26 \times F = 2.52 F$

$\text{Tl}^+ + 1e^- \rightarrow \text{Tl} = nFE^\circ = 1 \times (-0.336) \times F = -0.336 F$

$\text{Tl}^{3+} + 3e^- \rightarrow \text{Tl} \quad E^\circ = \frac{2.52F - 0.336F}{nF}$

=  $\frac{2.184}{3} = 0.728 \text{ V}$



**Q.76** (2)  
The lower the electrode potential, the stronger is the reducing agent.

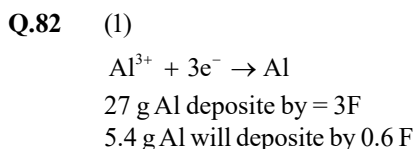
**Q.77** (3)  
 $W_{\text{Elec}} = -nF E_{\text{cell}}$   
and efficiency of cell  
=  $\frac{\Delta G}{\Delta H} \times 100 = \frac{-nF E_{\text{cell}}}{\Delta H} \times 100 = \frac{E_{\text{Elec}}}{\Delta H} \times 100$   
 $\therefore \Delta H \leq \text{Work electrical.}$

**Q.78** (4)  
 $\Lambda_m = \frac{K \times 1000}{M} = \frac{0.001 \times 1000}{0.1} = 10$

**Q.79** (4)  
All of the above.

**Q.80** (2)  
 $E_{\text{cell}}^\circ = E_{\text{Ce}^{4+}/\text{Ce}^{3+}}^\circ - E_{\text{Co}^{2+}/\text{Co}}^\circ$   
 $1.89 = E_{\text{Ce}^{4+}/\text{Ce}^{3+}}^\circ - (-0.28)$   
 $E_{\text{Ce}^{4+}/\text{Ce}^{3+}}^\circ = 1.89 - 0.28 = +1.61 \text{ V.}$

**Q.81** (3)  
As dry cell can't be recharge when it exhaust.



**Q.83** (1)  
In mercury cell cathode used is paste of HgO and carbon.

**Q.84** (3)  
On dilution the volume of the solution increases thus equivalent as well as molar conductivity increases. However the number of current carrying particles per  $\text{cm}^3$  decreases on diluting the solution.

- Q.85** (3)  
The number of faradays passed is always equal to the number of equivalents mass of produced. Just it is equivalents and not moles.  $\therefore$  2.5 F will deposit 2.5 g equivalents.

### SECTION-B

- Q.86** (2)  
HgI reacts with K.I and form  $K_2[HgI_4]$   
Due to this the no. of ions decreases thus decreasing the Vant Hoff factor  
 $[\Delta T_f = i k_f \times m]$   
Since  $i$  has decreased, it result in less depression in the freezing point causing the freezing point to raise.

- Q.87** (2)  
 $W = 15 \text{ gm}$   
 $M = 98 \text{ gm mol}^{-1}$   
 $V = \frac{\text{Mass}}{\text{Density}} = \frac{100}{1.5}$   
 $M = \frac{W \times 1000}{M \times V}$   
 $M = \frac{15 \times 1000}{98 \times \frac{100}{0.5}} = 2.29 \text{ M.}$

- Q.88** (4)  
Both solution have equal concentration and equal ions so  $\text{Ca}(\text{NO}_3)_2$  and  $\text{Na}_2\text{SO}_4$  isotonic in nature at same concentration.

- Q.89** (4)  
mole of  $\text{C}_2\text{H}_5\text{OH} = \frac{828}{46} = 18$   
mole of  $\text{H}_2\text{O} = \frac{36}{18} = 2$   
 $\therefore X_{\text{H}_2\text{O}} = \frac{n}{n+N} = \frac{2}{2+18} = 0.1$

- Q.90** (2)  
Conc. of  $\text{O}_2$  in ppm =  $\frac{\text{mass of O}_2}{\text{mass of H}_2\text{O}} = 10^6$   
[1kg  $\text{H}_2\text{O} = 10^6 \text{ mg H}_2\text{O}$ ]  
concentration in ppm =  $\frac{6 \text{ mg}}{10^6 \text{ mg}} \times 10^6 = 6 \text{ ppm.}$

- Q.91** (3)  
 $P_A = X_A \times P_A^0$        $P_B = X_B \times P_B^0$   
 $\therefore P_A^0 = \frac{P_A}{X_A}$        $P_B^0 = \frac{P_B}{X_B}$

A/c to the question  $P_A^0 > P_B^0$  thus.  $\frac{P_A}{X_A} > \frac{P_B}{X_B}$

Rearrang the relation to get

$$\frac{P_A}{P_B} > \frac{X_A}{X_B} \quad \dots(1)$$

By question YA is the mole fraction 6F A and YB is the mole fraction of B in vapour phase respectively

$$Y_A = \frac{P_A}{P_{\text{Total}}} \quad Y_B = \frac{P_B}{P_{\text{Total}}}$$

$$\left[ \frac{P_A}{P_B} = \frac{\frac{Y_A}{P_{\text{Total}}}}{\frac{Y_B}{P_{\text{Total}}}} \right] \quad \therefore \frac{P_A}{P_B} = \frac{Y_A}{Y_B} \quad \dots(2)$$

Combining the eq. (1) and eq. (2)

we get  $\left[ \frac{X_A}{X_B} < \frac{Y_A}{Y_B} \right]$

- Q.92** (4)  
Because antifreeze lowers the freezing point of water.

- Q.93** (2)
- |                    |                                |                                |
|--------------------|--------------------------------|--------------------------------|
| Ideal solution     | $\Delta H_{\text{mix}} = 0$    | $\Delta V_{\text{mix}} = 0$    |
| non ideal          | $\Delta H_{\text{mix}} \neq 0$ | $\Delta V_{\text{mix}} \neq 0$ |
| positive deviation | $\Delta H_{\text{mix}} > 0$    | $\Delta V_{\text{mix}} > 0$    |
| negative           | $\Delta H_{\text{mix}} < 0$    | $\Delta V_{\text{mix}} < 0$    |

- Q.94** (1)  
At anode  $\text{Cu} \rightarrow \text{Cu}^{2+} + 2e^-$   
At cathode  $2\text{Ag}^+ + 2e^- \rightarrow 2\text{Ag}$       where  $n = 2$   
Given  
 $E_{\text{cell}}^0 = 0.46$

$$E_{\text{cell}} = E_{\text{cell}}^0 - \frac{0.0591}{n} \log \frac{[\text{Cu}^{2+}]}{[\text{Ag}^+]^2}$$

$$E_{\text{cell}} = 0.46 - \frac{0.059}{2} \log \left[ \frac{0.01}{0.1 \times 0.1} \right] = 0.46$$

- Q.95** (2)  
An electrolytic cell

- Q.96** (1)  
 $2\text{NaCl} + 2\text{H}_2\text{O} \rightarrow 2\text{NaOH} + \text{H}_2 + \text{Cl}_2$   

Alkali	At	At
[increase basic	Cathode	Anode
nature of Solution]		

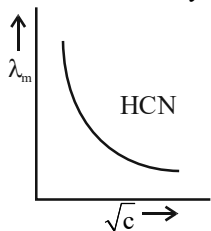
But concentration of  $\text{Na}^+$  ions not decreases.



Q.97 (1)

Q.98 (2)

For weak electrolytes



Q.99 (2)

$$\kappa = \frac{1}{77}$$

$$\Lambda_m = \frac{\kappa \times 1000}{C} = \frac{1000}{77 \times 0.1} = 130 \text{ Scm}^2\text{mol}^{-1}$$

Q.100 (3)

96500C  $\longrightarrow$   $N_A$  number of electron

$$(2 \times 60 \times 2)C \rightarrow \frac{N_A \times 240}{96500} = 0.015 \times 10^{23}$$

$$= 0.15 \times 10^{22}$$

**BIOLOGY-I  
SECTION-A**

Q.101 (1)

A typical anther is bilobed structure.

Q.102 (4)

Q.103 (1)

Egg apparatus is three celled structure.

Q.104 (1)

Spherical 25 - 50  $\mu\text{m}$

Q.105 (3)

Q.106 (4)

Q.107 (3)

Vegetative cell is layer, has irregularly shaped nucleus and stores food reserve.

Q.108 (2)

Q.109 (1)

Q.110 (3)

Q.111 (2)

Nucellus in angiosperms is diploid. MMC differentiates in the micropylar region from the cell of nucellus.

Q.112 (1)

Wheat has endospermic seeds.

Q.113 (2)

Seeds of black pepper and beet are perispermic.

Q.114 (1)

Q.115 (2)

Q.116 (2)

Q.117 (4)

The process of formation of microspores from a pollen mother cell (PMC) through meiosis is called microsporogenesis.

Q.118 (1)

Geitonogamy occurs on monoecious plant.

Q.119 (3)

Epiblast is present in embryo of monocot seeds.

Q.120 (3)

Mucilaginous covering protects the pollens from wetting in hydrophytes.

Q.121 (2)

Q.122 (3)

Q.123 (3)

Q.124 (1)

Q.125 (4)

Q.126 (3)

Q.127 (4)

Q.128 (2)

Q.129 (4)

Q.130 (2)

Q.131 (3)

Q.132 (3)

- Q.133** (3)  
Adequate moisture and temperature  $O_2$  is needed for germination
- Q.134** (1)  
Guava, orange and mango are the examples of fleshy fruits.
- Q.135** (3)
- Q.136** (3)  
Statement III is correct. Statements I and II are incorrect and can be corrected as
- Pollination by water is quite rare in flowering plants and is limited to about 30 genera, mostly monocotyledons.
  - Water is a regular mode of transport for the male gametes among lower plant groups such as algae and bryophytes.
- Q.137** (4)  
Hydrophilous flowers have mucilage covered pollen.
- Q.138** (2)
- Q.139** (2)
- Q.140** (2)
- Q.141** (4)  
Synergids are helper cells and they are not directly involved in double fertilisation.
- Q.142** (4)
- Q.143** (4)  
Asteraceae and Poaceae can form seeds without fertilisation by apomixis but when the fruit is formed without fertilisation. It is termed as parthenocarpy.
- Q.144** (3)
- Q.145** (4)
- Q.146** (4)  
In apple, fruit develops from thalamus along with ovary.
- Q.147** (3)
- Q.148** (3)
- Q.149** (2)
- Q.150** (2)  
Papaver - Multicarpellary syncarpous ovary.  
Wheat - Single ovule per ovary.
- Q.151** (4)
- Q.152** (4)
- Q.153** (4)  
Fallopian tube comprises three parts : infundibulum, ampulla and isthmus.
- Q.154** (1)
- Q.155** (3)  
1 oogonia  $\rightarrow$  1 ovum  
 $\therefore$  100 oogonia  $\rightarrow$  100 ova
- Q.156** (1)  
Both Assertion and Reason true, but Reason is not the correct explanation of Assertion.  
The correct explanation would be Fimbriae are finger-like projections of infundibulum of the oviduct which is closest to the ovary. Fimbriae collect the released from ovary and move it down into the Fallopian tube.
- Q.157** (4)  
During first trimester, major organ systems are formed.
- Q.158** (2)  
Hormones secreted by the placenta to maintain pregnancy are hCG, hPL, progesterone and estrogen.
- Q.159** (3)  
Endometrium is the innermost layer of uterus.
- Q.160** (4)  
The embryo with 8 to 16 blastomeres is called a morula.
- Q.161** (2)  
Fertilisation can only occur if the ovum and sperms are transported simultaneously to the ampullary region of fallopian tube.
- Q.162** (3)
- Q.163** (3)  
As luteal phase is fixed for 14 days, ovulation will take place on  $35-14 = 21^{\text{st}}$  day.
- Q.164** (1)
- Q.165** (2)  
Oxytocin from mother's pituitary.

- Q.166** (4)  
Interstitial cells secrete androgens.  
Leydig cells synthesize and secrete testicular hormones. Sertoli cells secrete inhibin.
- Q.167** (3)
- Q.168** (3)  
Spermatid conversion into sperm is called spermiogenesis/spermateliosis.
- Q.169** (3)
- Q.170** (4)
- Q.171** (2)
- Q.172** (1)
- Q.173** (2)
- Q.174** (3)  
The statement in option (c) is correct.  
Rest of the statements are incorrect and can be corrected as
- Size of testis is 4-5 cm in length and 2-3 cm in width.
  - The scrotum is maintained at a temperature 2-2.5°C lower than normal body temperature.
  - The earliest stages of spermatogenesis occur within the testis.
- Q.175** (1)  
Each testis has about 250 compartments called testicular lobules.
- Q.176** (4)  
In AI, the semen collected either from the husband or a healthy donor is artificially introduced either into the vagina or into the uterus (IUI - intro uterine insemination) of the female.
- Q.177** (1)  
Copper releasing and hormone releasing IUDs are considered as medicated IUDs.
- Q.178** (3)  
Lactational amenorrhea method i.e., absence of menstruation is based on the fact that ovulation and therefore the cycle do not occur during the period of intense lactation following parturition.
- Q.179** (4)  
IFT- Zygote Intra-Fallopian Transfer
- Q.180** (3)  
Amniocentesis can be used to test for the presence of certain genetic disorders such as Down's syndrome, haemophilia, sickle-cell anemia, etc.
- Q.181** (4)
- Q.182** (4)
- Q.183** (2)  
NCERT XII Pg# 63
- Q.184** (4)  
NCERT XI Pg# 64
- Q.185** (1)
- Q.186** (2)
- Q.187** (2)
- Q.188** (2)
- Q.189** (2)
- Q.190** (3)
- Q.191** (2)  
Formation of sperms is called spermatogenesis.
- Q.192** (1)  
Estrogen, LH and FSH are at high level during mid cycle.
- Q.193** (1)
- Q.194** (3)  
Near the nipple, mammary ducts expand to form mammary ampullae where some milk may be stored before going to lactiferous ducts.
- Q.195** (2)
- Q.196** (4)
- Q.197** (3)  
NCERT-XII, Pg#62
- Q.198** (2)
- Q.199** (1)
- Q.200** (1)