CBSE Test Paper-05 Class 12 Chemistry (Electrochemistry)

- 1. The standard emf of galvanic cell involving 3 moles of electrons in its redox reaction is 0.59 V. The equilibrium constant for the reaction of the cell is
 - a. 10²⁵
 - b. 10³⁰
 - c. 10¹⁵
 - d. 10²⁰
- 2. The standard reduction potential E_0 for half reactions are

$$E^0_{cell} = E^0_{cathode} - E^0_{anods}$$

The EMF of the cell reaction $Fe^{2+} + Zn = Zn^{2+} + Fe$ is--- [Given $E^0Zn^{2+}/Zn = -0.76V$;

$$E^{0}Fe^{2+}/Fe = -0.44V$$
]

- a. -1.17 V
- b. -0.32 V
- c. + 0.32 V
- d. +1.17 V
- 3. An increase in equivalent conductance of a strong electrolyte with dilution is mainly due to
 - a. increase in concentration of electrolyte
 - b. decrease in ionic mobility
 - c. increase in ionic mobility and number of ions
 - d. decrease in both i.e. number of ions and mobility of ions
- 4. Electrolytic conduction is due to the movement of:
 - a. molecules
 - b. ions
 - c. atoms
 - d. electrons
- 5. Relationship between equilibrium constant of the reaction and standard electrode potential of electrochemical cell in which that reaction takes place is

a.
$$E_{cell}^0 = rac{2.303 RT}{nF} \log K_c$$

- 6. What do you understand by corrosion?
- 7. Two metals A and B have reduction potential values -0.76 V and +0.34 V respectively. Which of these will liberate H_2 from dil H_2SO_4 ?
- 8. Define the term specific resistance and give its SI unit.
- 9. The resistance of a conductivity cell containing 0.001 M KCl solution at 298 K is $1500 \,\Omega$. What is the cell constant if conductivity of 0.001 M KCl solution at 298 K is $0.146 \times 10^{-3} S \, cm^{-1}$.
- 10. How much electricity in terms of Faraday is required to produce
 - i. 20.0 g of Ca from molten $CaCl_2$.
 - ii. 40.0 g of Al from molten Al_2O_3 .
- 11. Calculate E_{cell} for following: $2Cr(s) + 3Fe^{2+}(aq) \rightarrow 2Cr^{3+}(aq) + 3Fe(s)$ $Cr(s)|Cr^{3+}(aq)(0.1M)||Fe^{2+}(aq)(0.01M)|Fe(s)$ $E_{(Cr^{3+}/Cr)}^{\ominus} = -0.74 V$ $E_{(Fe^{2+}/Fe)}^{\ominus} = -0.44 V$
- 12. What type of a cell is the lead storage battery? Write the anode and cathode reactions and the overall reaction occurring in a lead storage battery while operating.
- 13. Write mathematical expression for Kohlrausch's law.
- 14. What is a salt bridge? What is it used for?
- 15. Calculate the standard cell potentials of galvanic cells in which the following reactions take place:
 - i. $2Cr(s) + 3Cd^{2+}(aq) \rightarrow 2Cr^{3+}(aq) + 3Cd$

ii.
$$Fe^{2+}(aq)+Ag^+(aq)
ightarrow Fe^{3+}(aq)+Ag(s)$$

Calculate the $\Delta_r G^{\Theta}$, and equilibrium constant of the reactions.

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1. b. 10^{30}

Explanation: $E^0 = \left(\frac{0.0591}{n}\right) \log k$ $0.59 = 0.0591/3 \log K$ $\log K = 3 \times 0.59/0.0591 = 30$ $K = AL 30 = 10^{30}$

2. c. + 0.32 V

Explanation: $E_{cell}^0 = E_{cathods}^0 - E_{anods}^0$ E^0 cell = E^0 Fe²⁺/Fe - E^0 Zn²⁺/Zn = -0.44-(-0.76)= +0.32V.

- a. increase in ionic mobility and number of ions
 Explanation: Equivalent conductance increases on dilution for a strong electrolyte as interionic attraction also decreases along with dilution. So ionic mobility increases which in turn increases the equivalent conductance.
- 4. c. atoms

Explanation: atoms

- 5. a. $E_{cell}^0 = \frac{2.303RT}{nF} \log K_c$ **Explanation:** $\Delta G^0 = -2.303 \text{ RT} \log \text{Kc}$; $\Delta G^0 = -n\text{FE}^0$ cell Equating, $E_{cell}^0 = \frac{2.303RT}{nF} \log Kc$
- 6. Corrosion is an electrochemical phenomenon in which metal gets decomposed in the presence of air and water and forms compounds like oxides, sulphates, carbonates, sulphides etc.
- 7. Metals having higher oxidation potential (or Lower reduction poetial) will liberate H_2 from H_2SO_4 . Thus, A will liberate H_2 from H_2SO_4 .
- 8. The specific resistance of a substance is its resistance when it is one-meter-long and its area of cross Section is one m². Its SI unit is Ωm (ohm meter).
- 9. Given,

Conductivity, $K = 0.146 \times 10^{-3} Scm^{-1}$ Resistance, R = 1500 Ω Therefore, Cell constant = $K \times R$ = $0.146 \times 10^{-3} \times 1500 = 0.219 cm^{-1}$

10. i. According to the question,

$$Ca^{2+}+2e^{-1}
ightarrow {Ca \over 40\,g}$$

Electricity required to produce 40 g of calcium = 2 F Therefore, electricity required to produce 20 g of calcium = $\frac{2 \times 20}{40}F$ = 1 F

ii. According to the question,

$$Al^{3+}+3e^{-1}
ightarrow Al_{27g}$$

Electricity required to produce 27 g of Al = 3 F

Therefore, electricity required to produce 40 g of Al = $\frac{3 \times 40}{27} F$

= 4.44 F

11. We have

$$2 ext{Cr}(s)$$
 + $3 ext{Fe}^{2+}(aq)$ $ightarrow$ $2 ext{Cr}^{3+}(aq)$ + $3 ext{Fe}(s)$
 $Cr(s)|Cr^{3+}(aq)(0.1M)||Fe^{2+}(aq)(0.01M)|Fe(s)$

Anode half reaction:

$$[Cr o Cr^{3+} + 3e^-] imes 2$$

Cathode half reaction:

$$[Fe^{2+}+2e^-
ightarrow Fe] imes 3$$
 .

Net cell reaction:

 $2Cr+3Fe^{2+}
ightarrow 2Cr^{3+}+3Fe$

Also standard emf of the cell is

$$E^{\Theta}_{cell} = E^{\Theta}_{(Fe^{2+}/Fe)} - E^{\Theta}_{(Cr^{3+}/Cr)}$$

According to Nernst equation

[Here **n=6** moles of electrons]

$$egin{split} E_{cell} &= E^{\Theta}_{cell} - rac{0.059}{n} \mathrm{log} \, rac{[Cr^{3+}]^2}{[Fe^{2+}]^3} \ E_{cell} &= E^{\Theta}_{cell} \, - \, rac{0.059}{6} \mathrm{log} iggl[rac{(0.1)^2}{(0.01)^3} iggr] \end{split}$$

- $= 0.30 \frac{0.059}{6} \log \left[\frac{10^{-2}}{10^{-6}} \right]$ = 0.30 - $\frac{0.059}{6} \log 10^4$ = 0.3 - $\frac{0.059}{6} \times 4$ = 0.30 - 0.039 = 0.261 V $\therefore E_{cell} = 0.261 V$
- 12. The lead storage battery is the most important secondary cell. The cell reaction when the battery is in use are given below:

At anode: $Pb(s) + SO_4^{2-}(aq) \to PbSO_4(s) + 2e^-$ At cathode: $PbO_2(s) + SO_4^{2-}(aq) + 4H^+(aq) + 2e^- \to 2PbSO_4(s) + 2H_2O(l)$ The overall cell reaction is: $Pb(s) + PbO_2(s) + 2H_2SO_4(aq) \to 2PbSO_4(s) + 2H_2O(l)$

13. Mathematical expression for Kohlrausch's law is Λ^∞_m or μ^∞ = Molar conductance at infinite dilution $= m\lambda^\infty_+ + n\lambda^\infty_-$

Where m and n are number of ions formed.

14. A salt bridge is a U-shaped tube containing concentrated solution of an inert electrolyte like KCl, KNO₃ etc. or solidified solution of such an electrolyte in agar-agar

and gelatine.

It is used for:

- i. To complete the electrical circuit by allowing ions to flow from one solution to the other without mixing the two solutions.
- ii. To maintain the electrical neutrality of the solution in the two half cells.

15. i.
$$E^{\Theta}_{Cr^{3+}/Cr} = 0.74V$$

$$E^{\Theta}{}_{Cd^{2+}/Cd} = 0.40V$$

The galvanic cell of the given reaction is depicted as:

$$\begin{split} &Cr_{(s)}|Cr^{3+}{}_{(aq)}||Cd^{2+}{}_{(aq)}|Cd_{(s)}\\ &\text{Now, the standard cell potential is }E^{\Theta}_{cell}=E^{\Theta}_{R}-E^{\Theta}_{L}\\ &=0.40\ \text{-}\ (\text{-}0.74)\\ &=+0.34\ \text{V}\\ &\Delta_{r}G^{\Theta}=-nFE^{\Theta}_{cell}\\ &\text{In the given equation,}\\ &\text{n}=6 \end{split}$$

$$\begin{split} & \mathsf{F} = 96487 \, \mathsf{C} \, \mathrm{mol}^{-1} \\ & E_{cell}^{\Theta} = +0.34 \, \mathsf{V} \\ & \mathsf{Then}, \, \Delta_r G^{\Theta} = -6 \times 96487 Cmol^{-1} \times 0.34V \\ & = -196833.48 \, \mathsf{CV} \, \mathsf{mol}^{-1} \\ & = -196833.48 \, \mathsf{I} \, \mathsf{mol}^{-1} \\ & = -196.86 \, \mathsf{kJ} \, \mathsf{mol}^{-1} \\ & \mathsf{Again}, \, \Delta_r G^{\Theta} = -RT \, ln \, K \\ & \Delta_r G^{\Theta} = -2.303 \, RT \, ln \, K \\ & \log \, \mathsf{K} = -\frac{\Delta_r G^{\Theta}}{2.303 RT} \\ & = \frac{196.83 \times 10^3}{2.303 \times 8.314 \times 298} \\ & = 34.496 \\ & \mathsf{Therefore}, \, \mathsf{K} = \mathsf{antilog} \, (34.496) \\ & = 3.13 \times 10^{34} \\ & \mathsf{ii}, \, E^{\Theta}_{Fe^{3+}/Fe^{2+}} = 0.77 \, V \\ & E^{\Theta}_{Ag^+/Ag} = 0.80 \, V \\ & \mathsf{The galvanic cell of the given reaction is depicted as: \\ & Fe^{2+}_{(aq)} | Fe^{3+}_{(aq)} | | Ag^+_{(aq)} | Ag_{(s)} \\ & \mathsf{Now}, \, \mathsf{the standard cell potential is} \, E_{cell}^{\Theta} = E_R^{\Theta} - E_L^{\Theta} \\ & = 0.80 \cdot 0.77 = 0.03 \, \mathsf{V} \\ & \mathsf{Here}, \, \mathsf{n} = 1. \\ & \mathsf{Then}, \Delta_r G^{\Theta} = -nFE_{cell}^{\Theta} \\ & = -1 \times 96487Cmol^{-1} \times 0.03V \\ & = -2894.61 \, \mathsf{J} \, \mathsf{mol}^{-1} \\ & \mathsf{Again}, \Delta_r G^{\Theta} = 2.303 \, RT \, In \, K \\ & \log \, K = -\frac{\Delta_r G}{2.303 RT} \\ & = \frac{-2894.61}{2.303 \times 8.314 \times 298} \\ & = 0.5073 \\ & \mathsf{Therefore}, \, \mathsf{K} = \mathsf{antilog} \, (0.5073) \\ & = 3.2 \, (\mathsf{approximately}) \end{split}$$