### **CBSE Test Paper**

## **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.  $10^{25}$
  - b.  $10^{30}$
  - c.  $10^{15}$
  - d.  $10^{20}$
- 2. The standard reduction potential  $E_{\rm o}$  for half reactions are

$$E_{cell}^0 = E_{cathode}^0 - E_{anods}^0$$

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

$$\begin{array}{l} \text{b.} \ E_{cell}^0 = \frac{2.03RT}{nF} {\rm log}\,K_c \\ \text{c.} \ E_{cell}^0 = \frac{2.230RT}{nF} {\rm log}\,K_c \\ \text{d.} \ E_{cell}^0 = \frac{2.303RT}{F} {\rm log}\,K_c \end{array}$$

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- 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  $0.146\times10^{-3}\,S\,cm^{-1}$ .
- 10. How much electricity in terms of Faraday is required to produce
  - i. 20.0 g of Ca from molten CaCl<sub>2</sub>.
  - ii. 40.0 g of Al from molten Al<sub>2</sub>O<sub>3</sub>.
- 11. Calculate  $E_{cell}$  for following:

$$egin{aligned} &2Cr(s)+3Fe^{2+}(aq) o 2Cr^{3+}(aq)+3Fe(s)\ &Cr(s)|Cr^{3+}(aq)(0.1M)||Fe^{2+}(aq)(0.01M)|Fe(s)\ &E^\ominus_{(Cr^{3+}/Cr)}\ =-0.74\ V\ &E^\ominus_{(Fe^{2+}/Fe)}\ =-0.44\ V \end{aligned}$$

- 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. 
$$2Cr(s)+3Cd^{2+}(aq)
ightarrow 2Cr^{3+}(aq)+3Cd$$

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

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

### **CBSE Test Paper-05**

# **Class 12 Chemistry (Electrochemistry)**

#### **Solutions**

1. b. 10<sup>30</sup>

Explanation: 
$$E^0 = \left(rac{0.0591}{n}
ight)\log k$$

$$0.59 = 0.0591/3 \log K$$

$$\log K = 3 \times 0.59/0.0591 = 30$$

$$K = AL 30 = 10^{30}$$

2. c. + 0.32 V

Explanation: 
$$E^0_{cell} = E^0_{cathods} - E^0_{anods}$$

$$E^{0}$$
cell =  $E^{0}$ Fe<sup>2+</sup>/Fe -  $E^{0}$ Zn<sup>2+</sup>/Zn

$$= -0.44 - (-0.76) = +0.32$$
V.

3. 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:**  $\Delta G^0 = -2.303 \text{ RT log Kc}$ ;  $\Delta G^0 = -\text{nFE}^0 \text{cell}$ 

Equating, 
$$E_{cell}^0 = rac{2.303RT}{nF} {
m log}\, Kc$$

- 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  $m^2$ . Its SI unit is  $\Omega m$  (ohm meter).
- 9. Given,

Conductivity,  $K=0.146\times 10^{-3} Scm^{-1}$ 

Resistance, R =  $1500\,\Omega$ 

Therefore, Cell constant = K imes R

$$= 0.146 \times 10^{-3} \times 1500 = 0.219 cm^{-1}$$

10. i. According to the question,

$$Ca^{2+}+rac{5}{2}e^{-1}
ightarrowrac{7}{40\,g}$$

Electricity required to produce 40 g of calcium = 2 F

Therefore, electricity required to produce 20 g of calcium =  $\frac{2\times20}{40}F$  = 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 =  $\frac{3\times40}{27}F$  = 4.44 F

$$2Cr(s) + 3Fe^{2+}(ag) \rightarrow 2Cr^{3+}(ag) + 3Fe(s)$$

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

Anode half reaction:

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

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

$$E_{cell}^{\Theta} = E_{(Fe^{2+}/Fe)}^{\Theta} - E_{(Cr^{3+}/Cr)}^{\Theta} = -0.44$$
-(-0.74) = 0.30 V

According to Nernst equation

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

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

$$= 0.30 - \frac{0.059}{6} log \left[ \frac{10^{-2}}{10^{-6}} \right]$$
 $= 0.30 - \frac{0.059}{6} log 10^{4}$ 
 $= 0.3 - \frac{0.059}{6} imes 4$ 
 $= 0.30 - 0.039 = 0.261 imes Cell = 0.261 imes V$ 

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:

$$Pb(s) + PbO_2(s) + 2H_2SO_4(aq) 
ightarrow 2PbSO_4(s) + 2H_2O(l)$$

13. Mathematical expression for Kohlrausch's law is

$$\Lambda_m^\infty$$
 or  $\mu^\infty$  = 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,  $KNO_3$  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:

$$Cr_{(s)}|Cr^{3+}{}_{(aq)}||Cd^{2+}{}_{(aq)}|Cd_{(s)}$$
  
Now, the standard cell potential is  $E^\Theta_{cell}=E^\Theta_R-E^\Theta_L$ 

$$= +0.34 \text{ V}$$

$$\Delta_r G^\Theta = -nF E^\Theta_{cell}$$

In the given equation,

$$n = 6$$

$$F = 96487 \text{ C mol}^{-1}$$

$$E_{cell}^{\Theta}$$
 = +0.34 V

Then, 
$$\Delta_r G^\Theta = -6 imes 96487 Cmol^{-1} imes 0.34 V$$

$$= -196.86 \text{ kJ mol}^{-1}$$

Again, 
$$\Delta_r G^\Theta = -RT \ln K$$

$$\Delta_r G^\Theta = -2.303\,RT\,ln\,K$$

$$\log \mathrm{K} = -rac{\Delta_r G^\Theta}{2.303 RT} = rac{196.83 \times 10^3}{2.303 \times 8.314 \times 298}$$

$$= 34.496$$

Therefore, K = antilog (34.496)

$$=3.13 \times 10^{34}$$

ii. 
$$E^{\Theta}{}_{Fe^{3+}/Fe^{2+}}=0.77\,V$$

$$E^\Theta{}_{Ag^+/Ag}=0.80\,V$$

The galvanic cell of the given reaction is depicted as:

$$Fe^{2+}{}_{(aq)}|Fe^{3+}{}_{(aq)}||Ag^{+}{}_{(aq)}|Ag_{(s)}$$

Now, the standard cell potential is  $E_{cell}^{\Theta}=E_{R}^{\Theta}-E_{L}^{\Theta}$ 

Here, n = 1.

Then,
$$\Delta_r G^\Theta = -nF E^\Theta_{cell}$$

=
$$-1 \times 96487 Cmol^{-1} \times 0.03 V$$

$$= -2.89 \text{ kJ mol}^{-1}$$

Again,
$$\Delta_r G^\Theta = 2.303\,RT\,In\,K$$

$$egin{array}{l} \log K = -rac{\Delta_r G}{2.303\,RT} \ = rac{-2894.61}{2.303 imes 8.314 imes 298} \end{array}$$

$$= 0.5073$$

Therefore, K = antilog (0.5073)

= 3.2 (approximately)