

Electrochemistry

- The equivalent conductances of sodium chloride, hydrochloric acid and sodium acetate at infinite dilution are 126.45, 426.16 and 91.0 $\text{ohm}^{-1} \text{cm}^2 \text{equiv}^{-1}$, respectively at 25 °C. Calculate the equivalent conductance of acetic acid at infinite dilution.
- 2F of electricity is passed through 20 L of a solution of aqueous solution of KCl. Calculate the pH of the solution.
- The specific conductivity of a solution containing 1.0g of anhydrous BaCl_2 in 200 cm^3 of the solution has been found to be 0.0058 S cm^{-1} . Calculate the molar conductivity of the solution. (Molecular wt. of $\text{BaCl}_2 = 208$).
- The equivalent conductivities of acetic acid at 298 K at the concentrations of 0.1 M and 0.001M are 5.20 and 49.2 $\text{S cm}^2 \text{eq}^{-1}$ respectively. Calculate the degree of dissociation of acetic acid at 0.001 M concentration.
Given that : $\Lambda^\infty(\text{H}^+)$ and $\Lambda^\infty(\text{CH}_3\text{COO}^-)$ are 349.8 and 40.9 $\text{ohm}^{-1} \text{cm}^2 \text{eq}^{-1}$ respectively.
- The amount of electricity which releases 2.0 g of gold from a gold salt is same as that which dissolves 0.967g of copper anode during the electrolysis of copper sulphate solution. What is the oxidation number of gold in the gold ion ? (At. mass of Cu = 63.5; Au = 197)
- If K_c for the reaction
$$\text{Cu}^{2+}(\text{aq}) + \text{Sn}^{2+}(\text{aq}) \longrightarrow \text{Sn}^{4+}(\text{aq}) + \text{Cu}(\text{s})$$
at 25°C is represented as 2.6×10^x then find the value of x.
(Given : $E^\circ_{\text{Cu}^{2+}|\text{Cu}} = 0.34\text{V}$; $E^\circ_{\text{Sn}^{4+}|\text{Sn}^{2+}} = 0.15\text{V}$)
- If ΔG° for the half cell $\text{MnO}_4^- | \text{MnO}_2$ in an acid solution is xF then find the value of x.
(Given : $E^\circ_{\text{MnO}_4^-|\text{Mn}^{2+}} = 1.5\text{V}$; $E^\circ_{\text{MnO}_2|\text{Mn}^{2+}} = 1.25\text{V}$)
- The standard reduction potential of a silver chloride electrode (metal-sparingly soluble salt electrode) is 0.209 V and for silver electrode is 0.80 V. If the moles of AgCl that can dissolve in 10 L of a 0.01 M NaCl solution is represented as 10^{-Z} then find the value of Z.
- Molar conductivity of aqueous solution of HA is 200 $\text{S cm}^2 \text{mol}^{-1}$, pH of this solution is 4. Calculate the value of $\text{pK}_a(\text{HA})$ at 25°C.

$$\begin{aligned}\text{Given : } \Lambda_M^\infty(\text{NaA}) &= 100 \text{ S cm}^2 \text{mol}^{-1}; \Lambda_M^\infty(\text{HCl}) \\ &= 425 \text{ S cm}^2 \text{mol}^{-1};\end{aligned}$$

$$\Lambda_M^\infty(\text{NaCl}) = 125 \text{ S cm}^2 \text{mol}^{-1}$$

- A solution containing 1M $\text{XSO}_4(\text{aq})$ and 1M $\text{YSO}_4(\text{aq})$ is electrolysed. If conc. of X^{2+} is 10^{-Z} M when deposition of Y^{2+} and X^{2+} starts simultaneously, calculate the value of Z.

$$\text{Given : } \frac{2.303RT}{F} = 0.06$$

$$E^\circ_{\text{X}^{2+}/\text{X}} = -0.12\text{V}; E^\circ_{\text{Y}^{2+}/\text{Y}} = -0.24\text{V}$$

- A solution of $\text{Ni}(\text{NO}_3)_2$ is electrolysed between platinum electrodes using 0.1 Faraday electricity. How many mole of Ni will be deposited at the cathode?
- Find the standard Gibbs energy for the given cell reaction in kJ mol^{-1} at 298 K
$$\text{Zn}(\text{s}) + \text{Cu}^{2+}(\text{aq}) \longrightarrow \text{Zn}^{2+}(\text{aq}) + \text{Cu}(\text{s}),$$
$$E^\circ = 2 \text{ V at } 298 \text{ K}$$
(Faraday's constant, $F = 96000 \text{ C mol}^{-1}$)
- All the energy released from the reaction $\text{X} \rightarrow \text{Y}$, $\Delta_r G^\circ = -193 \text{ kJ mol}^{-1}$ is used for oxidizing M^+ as $\text{M}^+ \rightarrow \text{M}^{3+} + 2\text{e}^-$, $E^\circ = -0.25 \text{ V}$
Under standard conditions, find the number of moles of M^+ oxidized when one mole of X is converted to Y
[$F = 96500 \text{ C mol}^{-1}$]
- Consider the following half-cell reactions and associated standard half-cell potentials, and determine the maximum voltage that can be obtained by combination resulting in spontaneous processes :
$$\text{AuBr}_4^-(\text{aq}) + 3\text{e}^- \longrightarrow \text{Au}(\text{s}) + 4\text{Br}^-(\text{aq}); E^\circ = -0.86 \text{ V}$$
$$\text{Eu}^{3+}(\text{aq}) + \text{e}^- \longrightarrow \text{Eu}^{2+}(\text{aq}); E^\circ = -0.43 \text{ V}$$
$$\text{Sn}^{2+}(\text{aq}) + 2\text{e}^- \longrightarrow \text{Sn}(\text{s}); E^\circ = -0.14 \text{ V}$$
$$\text{IO}^-(\text{aq}) + \text{H}_2\text{O}(\text{l}) + 2\text{e}^- \longrightarrow \text{I}^-(\text{aq}) + 2\text{OH}^-(\text{aq});$$
$$E^\circ = +0.49 \text{ V}$$
- The e.m.f. of the cell $\text{Zn} | \text{Zn}^{2+}(0.01\text{M}) || \text{Fe}^{2+}(0.001\text{M}) | \text{Fe}$ at 298 K is 0.2905 then the value of equilibrium constant for the cell reaction 10^x Find x

SOLUTIONS

1. (390.7) According to Kohlrausch's law,
 $\Lambda_{(eq)}^{\circ} \text{CH}_3\text{COONa} = \lambda_{\text{CH}_3\text{COO}^-} + \lambda_{\text{Na}^+} = 91.0 \dots (\text{i})$
 $\Lambda_{(eq)}^{\circ} \text{HCl} = \lambda_{\text{H}^+} + \lambda_{\text{Cl}^-} = 426.16 \dots (\text{ii})$
 $\Lambda_{(eq)}^{\circ} \text{NaCl} = \lambda_{\text{Na}^+} + \lambda_{\text{Cl}^-} = 126.45 \dots (\text{iii})$
 Adding equations (i) and (ii) and subtracting (iii),
 $\lambda_{\text{CH}_3\text{COO}^-} + \lambda_{\text{Na}^+} + \lambda_{\text{H}^+} + \lambda_{\text{Cl}^-} - \lambda_{\text{Na}^+} - \lambda_{\text{Cl}^-}$
 $= 91.0 + 426.16 - 126.45$
 $\lambda_{\text{CH}_3\text{COO}^-} + \lambda_{\text{H}^+} = \Lambda_{(eq)}^{\circ} \text{CH}_3\text{COOH}$
 $= 390.7 \text{ ohm}^{-1} \text{ cm}^2 \text{ equiv}^{-1}$

 2. (13) $\text{KCl} \xrightarrow[\text{at anode at cathode in solution}]{\text{Electrolysis}} \text{Cl}_2(\text{g}) + \text{H}_2(\text{g}) + \text{OH}^-$
 $1\text{F} = 1 \text{ eq of } \text{H}_2(\text{g}) = 1 \text{ eq of } \text{Cl}_2(\text{g}) \equiv 1 \text{ eq of } \text{OH}^- \text{ ions}$
 $2\text{F} = 2 \text{ eq of } \text{OH}^-$
 $[\text{OH}^-] = \frac{2 \text{ eq}}{\text{Volume in L}} = \frac{2}{20\text{L}} = 10^{-1} \text{ Nor M}$
 $\therefore \text{pOH} = -\log(10^{-1}) = 1$
 $\text{pH} = 14 - 1 = 13$

 3. (241.67) Molarity of $\text{BaCl}_2 = \frac{1 \times 1000}{208 \times 200} = 0.024 \text{ M}$
 Also, Normality of $\text{BaCl}_2 = 0.024 \times 2 = 0.048 \text{ N}$
 $(\because \text{N} = \text{M} \times \text{Valency factor})$
 Now, $\Lambda_m = \kappa \times \frac{1000}{C_M} = \frac{0.0058 \times 1000}{0.024}$
 $= 241.67 \text{ S cm}^2 \text{ mol}^{-1}$

 4. (12.5) Degree of dissociation is given by
 $\alpha = \frac{\Lambda^c}{\Lambda^\infty}$
 (i) Evaluation of $\Lambda_{\text{CH}_3\text{COOH}}^\infty$
 $\Lambda_{\text{CH}_3\text{COOH}}^\infty = \Lambda_{\text{CH}_3\text{COO}^-}^\infty + \Lambda_{\text{H}^+}^\infty$
 $= 40.9 + 349.8 = 390.7 \text{ ohm}^{-1} \text{ cm}^2 \text{ eq}^{-1}$
 (ii) Evaluation of degree of dissociation
 At $C = 0.001 \text{ M}$,
 $\alpha = \frac{\Lambda^c}{\Lambda^\infty} = \frac{49.2}{390.7} = 0.125 \text{ i.e., } 12.5\%$

 5. (3) $\frac{0.967}{63.5} \times 2 = \frac{2}{197} \times n_f$
 $n_f = 3$

 6. (6) $K_c = 10^{\frac{2(0.34 - 0.15)}{0.0591}} = 2.6 \times 10^6$

 7. (5) $4\text{H}^+ + \text{MnO}_4^- \rightleftharpoons \text{MnO}_2 + 2\text{H}_2\text{O}$
 $\Delta G^\circ = -3 \times F \times \left(\frac{1.5 \times 5 - 2 \times 1.25}{3} \right)$
 $= -5 \text{ F}; x = 5$

 8. (7) $E_{\text{Cl}^-/\text{AgCl}/\text{Ag}}^\circ = E_{\text{Ag}^+/\text{Ag}}^\circ + 1 \frac{0.0591}{1} \log K_{sp}$
 $0.209 = 0.80 + \frac{0.0591}{1} \log K_{sp}$
 $K_{sp} = 10^{-10}$; Let solubility of AgCl in 0.01 M solution is x
 $10^{-10} = x(x + 0.01)$
 $x = 10^{-8}$
 $\therefore \text{Moles of AgCl dissolved in } 10\text{L} = 10^{-8} \times 10 = 10^{-7}$

 9. (4) $\Lambda_M^\infty(\text{HA}) = \Lambda_M^\infty(\text{HCl}) + \Lambda_M^\infty(\text{NaA}) - \Lambda_M^\infty(\text{NaCl})$
 $= 425 + 100 - 125 = 400 \text{ S cm}^2 \text{ mol}^{-1}$
 $\text{pH} = 4, [\text{H}^+] = 10^{-4} = \alpha C$
 $\alpha = \frac{\Lambda_m}{\Lambda_m^\infty} = \frac{200}{400} = 0.5;$
 $K_a = \frac{(C\alpha) \cdot \alpha}{(1-\alpha)} = \frac{10^{-4}(0.5)}{(1-\alpha)} = 10^{-4}; \text{p}K_a = 4$

 10. (4) $-0.12 - \frac{0.0591}{2} \log\left(\frac{1}{X}\right) = -0.24$
 $\log \frac{1}{X} = \frac{0.12 \times 2}{0.06} = 4$
 $X = 10^{-4}$

 11. (0.05) According to the Faraday's law of electrolysis, nF of current is required for the deposition of 1 mole.
 According to the reaction,
 $\text{Ni}(\text{NO}_3)_2 \longrightarrow \text{Ni}^{2+} + 2\text{NO}_3^-$
 $2 \text{ F of current deposits} = 1 \text{ mol}$
 $\therefore 0.1 \text{ F of current deposits} = 0.05 \text{ mol}$

 12. (-384) $\Delta G^\circ = -nFE_{\text{cell}}^\circ$
 $= -2 \times (96000) \times 2 \text{ V} = -384000 \text{ J/mol}$
 $= -384 \text{ kJ/mol}$

 13. (4) $X \longrightarrow Y; \Delta G^\circ = -193 \text{ kJ mol}^{-1}$
 $M^+ \longrightarrow M^{3+} + 2e^- \quad E^\circ = -0.25 \text{V}$
 Hence ΔG° for oxidation will be
 $\Delta G^\circ = -nFE^\circ$
 $= -2 \times 96500 \times (-0.25) = 48250 \text{ J} = 48.25 \text{ kJ}$
 $48.25 \text{ kJ energy oxidises one mole } M^+$
 $\therefore 193 \text{ kJ energy oxidises } \frac{193}{48.25} \text{ mole } M^+ = 4 \text{ mole } M^+$

 14. (1.35) Maximum voltage $E^\circ = 0.49 + 0.86 = 1.35 \text{ V}$
 15. (10.85) For this cell, reaction is; $\text{Zn} + \text{Fe}^{2+} \rightarrow \text{Zn}^{2+} + \text{Fe}$
 $E = E^\circ - \frac{0.0591}{n} \log \frac{c_1}{c_2}; E^\circ = E + \frac{0.0591}{n} \log \frac{c_1}{c_2}$

$$E^{\circ} = 0.2905 + \frac{0.0591}{2} \log \frac{10^{-2}}{10^{-3}} = 0.32 \text{ V.}$$

$$E^{\circ} = \frac{0.0591}{2} \log K_{\text{eq}}$$

$$\log K_{\text{eq}} = \frac{0.32 \times 2}{0.0591} = \frac{0.32}{0.0295}$$

$$\therefore K_{\text{eq}} = 10^{\frac{0.32}{0.0295}}$$

Comparing the value of 10^x ,

$$x = \frac{0.32}{0.0295} = 10.847 \approx 10.85$$