- 1. The E^{Θ} values for $Al^{+}/Al = +0.55 \text{ V} \text{ and } Tl^{+}/Tl = -0.34 \text{ V}$ $Al^{3+}/Al = -1.66 \text{ V}$ and $T^{3+}/Tl = +1.26 \text{ V}$ 5. Identify the incorrect statement (2023)(a) Al is more electropositive than Tl (b) Tl³⁺ is a good reducing agent than Tl¹⁺ (c) Al⁺ is unstable in solution (d) Al⁺ can be easily oxidised Tl than Tl³⁺ 2. Molar conductance of an electrolyte increase with dilution according to the equation: $\Lambda_{\rm m} = \Lambda_{\rm m}^{\circ} - A\sqrt{\rm c}$ Which of the following statements are true? (a) This equation applies to both strong and weak electrolytes. (b) Value of the constant A depends upon the nature of the solvent. (c) Value of constant A is same for both 6. BaCl₂ and MgSO₄. (d) Value of constant A is same for both $BaCl_2$ and $Mg(OH)_2$. Choose the most appropriate answer from the options given below: (2023)(a) A and B only (b) A, B and C only (c) B and C only (d) B and D only 7. 3. The correct value of cell potential in volt for the reaction that occurs when the following two half cells are connected, is $Fe^{2+}(aq) + 2e^- \rightarrow Fe(s), E^\circ = -0.44 V$ $Cr_2O_7^{2-}(aq) + 14H^+ + 6e^- \rightarrow 2Cr^{3+} + 7H_2O$ $E^{\circ} = +1.33 V$ (2023)(a) +1.77 V (b) +2.65 V (c) +0.01 V (d) +0.89 V 8. The conductivity of centimolar solution of 4. KCl at 25°C is 0.0210 ohm⁻¹ cm⁻¹ and the resistance of the cell containing the solution at 25°C is 60 ohm. The value of cell constant is-(2023)(a) 3.28 cm⁻¹ (b) 1.26 cm⁻¹
- (c) 3.34 cm⁻¹
 - (d) 1.34 cm⁻¹
 - Given below are two statements: One is labelled as Assertion A and the other is labelled as Reason R: **Assertion:** In equation $\Delta_{rG} = -nFE_{cell}$, vlaue of Δ_{rG} depends on n. **Reason:** E_{cell} is an intensive property and Δ_{rG} is an extensive property. In the light of the above statements, choose the correct answer from the options given below: (2023)(a) Both A and R are true but R is NOT the correct explanation of A. (b) A is true but R is false. (c) A is false but R is true. (d) Both A and R are true R R is the correct explanation of A. Two half cell reactions are given below: $Co^{3+} + e^- \rightarrow Co^{2+}, E^{\circ}_{CO^{2+}/Co^{3+}} = -1.81 \text{ V}$ $2Al^{3+} + 6e^- \rightarrow 2Al(s), E^{\circ}_{Al/Al^{3+}} = +1.66 \text{ V}$ The standard EMF of a cell with feasible redox reaction will be: (2022)(a) −3.47 V (b) +7.09 V (c) +0.15 V (d) +3.47 V Standard electrode potential for the cell with cell reaction $Zn(s) + Cu^{2+}(aq) \rightarrow Zn^{2+}(aq) + Cu(s)$ is 1.1 V. Calculate the standard Gibbs energy change for the cell reaction. (Given: $F = 96487 \text{ C mol}^{-1}$) (2022)(a) -200.27 J mol⁻¹ (b) -200.27 kJ mol-1 (c) -212.27 kJ mol⁻¹ (d) -212.27 J mol-1 At 298 K, the standard electrode potentials of Cu^{2+}/Cu , Zn^{2+}/Zn , Fe^{2+}/Fe and Ag^{+}/Ag are 0.34 V, -0.76 V, -0.44 V and 0.80 V, respectively. On the basis of standard electrode potential, predict which of the following reaction cannot occur?
 - (a) $CuSO_4(aq) + Zn(s) \rightarrow ZnSO_4(aq) + Cu(s)$

(b) $CuSO_4(aq) + Fe(s) \rightarrow FeSO_4(aq) + Cu(s)$ (c) $FeSO_4(aq) + Zn(s) \rightarrow ZnSO_4(aq) + Fe(s)$ (d) $2CuSO_4(aq) + 2Ag(s) \rightarrow 2Cu(s) +$ $Ag_2SO_4(aq)$ 9. Given below are half cell reactions: $MnO_{4}^{-} + 8H^{+} + 5e^{-} \rightarrow Mn^{2+} + 4H_{2}O$, $E_{Mn^{2+}/MnO_{4}}^{\circ} = -1.510 \text{ V}$ $\frac{1}{2}O_2 + 2H^+ + 2e^- \longrightarrow H_2O$ $E^{\circ}_{O_2/H_2O} = +1.223 \text{ V}$ will the permanganate ion, $Mn0_4^-$ liberate O_2 from water in the presence of an acid? (2022)(a) Yes, because E_{cell}° = +0.287 V (b) No, because $E_{cell}^{\circ} = -0.287 \text{ V}$ (c) Yes, because E_{cell}° = +2.733 V (d) No, because $E_{cell}^{\circ} = -2.733 \text{ V}$ 10. Find the emf of the cell in which the following reaction takes place at 298 K $Ni(s) + 2Ag^{+}(0.001 \text{ M}) \rightarrow Ni^{2+}(0.001 \text{ M}) +$ 2Ag(s)(Given that $E_{cell}^{\circ} = 10.5 \text{ V}, \frac{2.303 \text{ RT}}{F} = 0.059 \text{ at}$ 298 K) (2022)(a) 1.0385 V (b) 1.385 V (c) 0.9615 V (d) 1.05 V 11. The molar conductance of NaCl, HCl and CH₃COONa at infinite dilution are 126.45, 426.16 and 91.0 S cm² mol⁻¹ respectively. The molar conductance of CH₃COOH at infinite dilution is. Choose the right option for your answer. (2021)(a) $390.71 S cm^2 mol^{-1}$ (b) $698.28 S cm^2 mol^{-1}$ (c) $540.48 S cm^2 mol^{-1}$ (d) $201.28 S cm^2 mol^{-1}$ 12. The molar conductivity of 0.007 M acetic acid is 20 S cm² mol⁻¹. What is the dissociation constant of acetic acid? Choose the correct option. (2021) $\Lambda_{H^+}^{\circ} = 350 \, S \, cm^2 mol^{-1}$ $\left[\Lambda_{CH_{3}COO^{-}}^{\circ} = 50 \, S \, cm^{2} mol^{-1}\right]$ (a) $2.50 \times 10^{-4} mol L^{-1}$ (b) $1.75 \times 10^{-5} mol L^{-1}$ (c) $2.50 \times 10^{-5} mol L^{-1}$ (d) $1.75 \times 10^{-4} mol L^{-1}$ 13. On electrolysis of dil sulphuric acid using Platinum (Pt) electrode, product the

obtained at anode will be:

(a) Oxygen gas (b) H_2S gas (c) SO_2 gas (d) Hydrogen gas 14. The number of Faradays (F) required to produce 20 g of calcium from molten CaCl₂ (Atomic mass of Ca = 40 g mol^{-1}) is: (2020) (a) 2 (b) 3 (c) 4 (d) 1 15. Identify the reaction from following having top position in EMF series (Std. red. Potential) according to their electrode potential at 298 K. (2020 Covid Re-NEET) (a) $Fe^{2+} + 2e^- \rightarrow Fe(s)$ (b) $Au^{3+} + 3e^- \rightarrow Au(s)$ (c) $K^+ + le^- \rightarrow K(s)$ (d) $Mq^{2+} + 2e^- \rightarrow Mq(s)$ 16. In a typical fuel cell, the reactants (R) and product (P) are (2020 Covid Re-NEET) (a) $R = H_{2(q)}, O_{2(q)}; P = H_2 O_{(l)}$ (b) $R = H_{2(g)}, O_{2(g)}, Cl_{2(g)}; P = HClO_{4(aq)}$ (c) $R = H_{2(q)}, N_{2(q)}; P = NH_{3(aq)}$ (d) $R = H_{2(q)}, O_{2(q)}; P = H_2 O_{2(l)}$ 17. For a cell involving one electron E° cell = 0.59 V at 298 K, the equilibrium constant for the cell reaction is: (2019) $\left[Given that \frac{2.303RT}{F} = 0.059V at T = 298K\right]$ (a) 1.0×10^2 (b) 1.0×10^5 (c) 1.0×10^{10} (d) 1.0×10^{30} 18. For the cell reaction $2Fe^{3+}(aq) + 2I^{-}(aq) \rightarrow 2Fe^{2+}(aq) + I_2(aq)$ $E_{cell}^{\Theta} = 0.24 V$ at 298 K. The standard Gibbs energy $(\Delta_r G^{\ominus})$ of the cell reaction is: [Given that Faraday constant F = 96500 Cmol-1] (2019)(a) -46.32 kJ mol⁻¹

- (b) -23.16 kJ mol⁻¹
- (c) 46.32 kJ mol⁻¹
- (d) 23.16 kJ mol⁻¹

(2020)

19. Consider the change in oxidation state of Bromine corresponding to different emf values as shown in the diagram below:

$$BrO_{4}^{-} \xrightarrow{1.82 \text{ V}} BrO_{3}^{-} \xrightarrow{1.5 \text{ V}} HBrO$$
$$Br^{-} \xrightarrow{1.0652 \text{ V}} Br_{2} \xrightarrow{1.595 \text{ V}}$$

24. Zinc can be coated on iron to produce Then the species undergoing galvanized iron but the reverse is not possible. It is because: disproportionation is: (2018)(2016-II) (a) BrO_3^- (a) Zinc has lower negative electrode (b) $BrO_4^$ potential than iron (c) HBrO (b) Zinc has higher negative electrode potential than iron (d) Br_2 (c) Zinc is lighter than iron 20. In the electrochemical cell (d) Zinc has lower melting point than iron $Zn|ZnSO_4(0.01M)||CuSO_4(1.0M)|Cu$, the emf of Daniel cell is E_1 . When the 25. The number of electrons delivered at the this cathode during electrolysis by a current of concentration of $ZnSO_4$ is changed to 1.0 M 1 ampere in 60 seconds is: and that of $CuSO_4$ changed to 0.01 M, the (2016-II) (charge on electron = 1.60×10^{-19} C) emf changes to E_2 . From the following, which one is the relationship between E_1 (a) 3.75×10^{20} and E₂? (Given, $\frac{RT}{F} = 0.059$) (b) 7.48 × 10²³ (c) 6×10^{23} (2017-Gujarat) (d) 6×10^{20} (a) $E_2 = 0 \neq E$ 26. If the E°cell for a given reaction has a (b) $E_1 = E_2$ (c) $E_1 < E_2$ negative value, which of the following gives (d) $E_1 > E_2$ the correct relationships for the values of ΔG° and K_{eq} ? (2016-II)21. Given that $\Lambda_m^{\circ} = 133.4, 5 \ cm^2 \ mol^{-1} \ (AgNO_3);$ (a) $\Delta G^0 < 0; K_{eq} > 1$ $\Lambda_{m}^{\circ} = 149.9 \, S \, cm^{2} \, mol^{-1} \, (KCl);$ (b) $\Delta G^0 < 0; K_{eq} < 1$ $\Lambda_{m}^{\circ} = 144.9 \, S \, cm^2 \, mol^{-1} \, (KNO_3)$ the molar (c) $\Delta G^0 > 0; K_{ea} < 1$ conductivity at infinite dilution for AgCl is: (d) $\Delta G^0 > 0; K_{eq} > 1$ (2017-Gujarat)(a) $132 S cm^2 mol^{-1}$ 27. The molar conductivity of a 0.5 mol dm⁻³ (b) 140 S cm² mol⁻¹ solution of $AgNO_3$ with electrolytic (c) $138 S cm^2 mol^{-1}$ conductivity of 5.76×10^{-3} S cm⁻¹ at 298 K (d) $134 S cm^2 mol^{-1}$ is: (2016-II) (a) 0.086 S cm² mol⁻¹ 22. The zinc/silver oxide cell is used in electric (b) 28.8 S cm² mol⁻¹ watches. The reaction is as following: (c) 2.88 S cm² mol⁻¹ $Zn^{2+} + 2e^- \rightarrow Zn ;$ $E^{\circ} = -0.760 V$ (d) 11.52 S cm² mol⁻¹ $Ag_2O + H_2O + 2e^- \rightarrow 2Ag + 2OH^-$; $E^{\circ} = 0.344 V$ 28. The pressure of H_2 required to make the potential of H₂ electrode zero in pure water If F is 96,500 C mol⁻¹, ΔG° of the cell will be: at 298 K is: (2016-I)(2017-Gujarat) (a) 10⁻⁴ atm (a) 413.21 kJ mol⁻¹ (b) 10⁻¹⁴ atm (b) 113.072 kJ mol⁻¹ (c) 10⁻¹² atm (c) -213.072 kJ mol⁻¹ (d) 10-10 atm (d) 4313.082 kJ mol⁻¹ 29. A device that converts energy of combustion 23. During the electrolysis of molten sodium of fuels like hydrogen and methane, directly chloride, the time required to produce 0.10 into electrical energy is known as: (2015) mol of chlorine gas using a current of 3 (a) Electrolytic cell (2016-II) amperes is: (b) Dynamo (a) 220 minutes (c) Ni-Cd cell (b) 330 minutes (d) Fuel cell (c) 55 minutes

(d) 110 minutes

30. The pair of compounds that can exist together is: (a) $HgCl_2, SnCl_2$

(2014)

- (b) $FeCl_2, SnCl_2$
- (c) FeCl₃, KI
- (d) $FeCl_3, SnCl_2$
- 31. Using the Gibbs energy change, ΔG⁰ = +63.3 kJ, for the following reaction, Ag₂CO₃(s) ≈ 2Ag⁺(aq) + CO₃²⁻(aq) the K_{sp} of Ag₂CO₃(s) in water at 25°C is: (2014) (R = 8.314 JK⁻¹mol⁻¹)
 (a) 8.0 × 10⁻¹²
 (b) 2.9 × 10⁻³
 (c) 7.9 × 10⁻²
 - (d) 3.2×10^{-26}
- 32. When 0.1 mol MnO_4^{2-} is oxidized, the quantity of electricity required to completely oxidise MnO_4^{2-} to MnO_4^{-} is: (2014) (a) 96500 C
 - (b) 2 × 96500 C
 - (c) 9650 C
 - (d) 96.50 C
- 33. The weight of silver (atomic weight = 108) displaced by a quantity of electricity which displaces 5600 mL of O₂ at STP will be:
 - (2014)

- (a) 10.8 g
- (b) 54.0 g
- (c) 108.0 g
- (d) 5.4 g
- 34. At 25°C, molar conductance of 0.1 molar aqueous solution of ammonium hydroxide is 9.54 ohm⁻¹ cm² mol⁻¹ and at infinite dilution its molar conductance is

238 ohm⁻¹ cm² mol⁻¹. The degree of ionization of ammonium hydroxide at the same concentration and temperature is:

(2013)

(2013)

- (a) 2.080%
- (b) 20.800%
- (c) 4.008%
- (d) 40.800%
- 35. A hydrogen gas electrode is made by dipping platinum wire in a solution of HCl of pH = 10 and by passing hydrogen gas around the platinum wire at one atm pressure. The oxidation potential of electrode would be?

0 050 V

- (a) 0.059 V(b) 0.59 V
- (c) 0.118 V
- (d) 1.18 V

- 36. A button cell used in watches functions as following $Zn(s) + Ag_2O(s) + H_2O(l) \rightleftharpoons$ $2Ag(s) + Zn^{2+}(aq) + 20H^-(aq).$ If half cell potentials are $Zn^{2+}(aq) + 2e^- \rightarrow Zn(s)$; $E^\circ = -0.76V$ $Ag_2O(s) + H_2O(l) + 2e^- \rightarrow 2Ag(s) + 20H^-(aq)$ $E^\circ = 0.34V$. The cell potential will be: (2013) (a) 1.10 V
 - (b) 0.42 V
 - (c) 0.84 V (d) 1.34 V

Answer Key	
S1. Ans. (b)	S33. Ans. (c)
S2. Ans. (d)	S34. Ans. (c)
S3. Ans. (a)	S35. Ans. (b)
S4. Ans. (b)	S36. Ans. (a)
S5. Ans. (d)	
S6. Ans. (d)	
S7. Ans. (c)	
S8. Ans. (d)	
S9. Ans. (a)	
S10. Ans. (a)	
S11. Ans. (a)	
S12. Ans. (b)	
S13. Ans. (a)	
S14. Ans. (d)	
S15. Ans. (b)	
S16. Ans. (a)	
S17. Ans. (c)	
S18. Ans. (a)	
S19. Ans. (c)	
S20. Ans. (d)	
S21. Ans. (c)	
S22. Ans. (c)	
S23. Ans. (d)	
S24. Ans. (b)	
S25. Ans. (a)	
S26. Ans. (c)	
S27. Ans. (d)	
S28. Ans. (b)	
S29. Ans. (d)	
S30. Ans. (b)	
S31. Ans. (a)	
S32. Ans. (c)	

S1. Ans. (b) Tl³ act as an oxidising agent not reducing agent. S2. Ans. (d) B and D statements are correct. S3. Ans. (a) $\mathbf{E}_{cell}^{\circ} = \mathbf{E}_{C}^{\circ} - \mathbf{E}_{A}^{\circ}$ = (1.33) - (-0.44)= +1.77 VS4. Ans. (b) Centimolar solution = 1/100 M = 0.01 MConductivity (k) = $0.0210 \text{ ohm}^{-1} \text{ cm}^{-1}$ Resistance (R) = 60 ohm $k = 1/R\left(\frac{1}{\Lambda}\right)$ $\Rightarrow 0.0210 = 1/60\left(\frac{l}{A}\right) \Rightarrow \frac{l}{A} = 1.26 \text{ cm}^{-1}$ S5. Ans. (d) S9. $\Delta_{\rm rG} = -nFE_{\rm cell}$ E_{cell} is an intensive property and Δ_{rG} is an extensive property as it depends on number of e⁻ transferred in cell reaction. S6. Ans. (d) Since, $\tilde{E_{OP}}$ of Al is more than Co^{2+} , so at anode Al will oxidise and at cathode Co3+ will reduce. $\mathbf{E}_{\text{cell}}^{\circ} = (\mathbf{E}_{\text{Cathode}}^{\circ})_{\text{RP}} - (\mathbf{E}_{\text{Anode}}^{\circ})_{\text{RP}}$ $= E_{Co^{3+}/Co^{2+}}^{\circ} - E_{Al^{3+}/Al}^{\circ}$ = (1.81) - (-1.66)= +3.47 V S7. Ans. (c) $Zn(s) + Cu^{2+}(aq) \rightarrow Zn^{2+}(aq) + Cu(s)$ S10. $E_{cell}^{\circ} = 1.1 V$ $\Delta G^{\circ} = -nFE_{cell}^{\circ}$ ∴ n = 2 $\Delta G^{\circ} = -2 \times 96487 \times 1.1$ $\Delta G^{\circ} = -212271.4 \text{ J mol}^{-1}$ $\Delta G^{\circ} = -212.27 \text{ kJ mol}^{-1}$ S8. Ans. (d) For a reaction to be spontaneous, E_{cell}° must be positive.

■ For, FeSO₄(aq) + Zn(s) \rightarrow ZnSO₄(aq) + Fe(s) $E_{cell}^{\circ} = E_{cathode}^{\circ} - E_{anode}^{\circ}$ = -0.44 V - (-0.76 V)= 0.32 V For, $2CuSO_4(aq) + 2Ag(s) \rightarrow 2Cu(s) +$ $Ag_2SO_4(aq)$ $E_{cell}^{\circ} = 0.34 \text{ V} - 0.80 \text{ V}$ = -0.46 V■ For, $CuSO_4(aq) + Zn(s) \rightarrow ZnSO_4(aq) +$ Cu(s) $E_{cell}^{\circ} = 0.34 \text{ V} - (-0.76 \text{ V})$ = 1.1 V ■ Foe, $CuSO_4(aq) + Fe(s) \rightarrow FeSO_4(aq) +$ Cu(s) $E_{cell}^{\circ} = 0.80 \text{ V} - (-0.44 \text{ V})$ = 1.24 V Ans. (a) $\blacksquare MnO_4^- + 8H^+ + 5e^- \rightarrow Mn^{2+} + 4H_2O -(i)$ $E_{MnO_{-}/Mn^{2+}}^{\circ} = -E_{Mn^{2+}/MnO_{-}}^{\circ} = 1.51 \text{ V}$ $\blacksquare H_2O \longrightarrow \frac{1}{2}O_2 + 2H^+ + 2e^-$ -(ii) $E_{0_2/H_20}^{\circ} = 1.223 V$ Using $2 \times (i) + 5 \times (ii)$, net cell reactions is $2Mn0_{4}^{-} + 6H^{+} \rightarrow 2Mn^{2+} + \frac{5}{2}O_{2} + 3H_{2}O_{3}$ $\mathbf{E}_{\text{cell}}^{\circ} = \mathbf{E}_{\text{C}}^{\circ} - \mathbf{E}_{\text{A}}^{\circ} = \mathbf{E}_{\text{MnO}_{4}^{-}/\text{Mn}^{2+}}^{\circ} - \mathbf{E}_{\text{O}_{2}/\text{H}_{2}\text{O}}^{\circ}$ = 1.51 - 1.223 = 0.287 V Since $\tilde{E}_{cell} > 0$, therefore net cell reaction is spontaneous and so $Mn0_4^-$ liberate O_2 from H₂O in presence of an acid. Ans. (a) $Ni(s) + 2Ag^{+}(0.001 \text{ M}) \rightarrow Ni^{2+}(0.001 \text{ M}) +$ 2Ag(s) \tilde{E}_{cell}° = 10.5 V $E_{cell}^{\circ} = E_{cell}^{\circ} - \frac{0.059}{n} \log \frac{[ni^{2+}]}{[Ag^{+}]^{2}}$ $\Rightarrow 10.5 - \frac{0.059}{2} \log \frac{(10^{-3})}{(10^{-3})^2}$ $\Rightarrow 10.5 - \frac{0.059}{2} \log(10)^3$ $\Rightarrow 10.5 - 0.0295 \times 3$

 $\Rightarrow 10.5 - 0.0885$

 \Rightarrow 10.4115 V S11. Ans.(a) $\Lambda_{NaCl} = \Lambda_{Na^+} + \Lambda_{Cl^-}$ $\Lambda_{HCl} = \Lambda_{H^+} + \Lambda_{Cl^-}$ $\Lambda_{CH_3COONa} = \Lambda_{Na^+} + \Lambda_{CH_3COO^-}$ Let, $\Lambda_{Na^+} = x$, $\Lambda_{Cl^-} = y$, $\Lambda_{H^+} = \Lambda_{CH_3COO^- = w}$ Given, x + y = 126.45...(i) y + z = 426.16...(ii) x + w = 91...(iii) From the above 3 equations, value of z + w = 390.71dS12. Ans.(b) $\Lambda^{\circ}_{M(CH_{3}COOH)} = \Lambda^{\circ}_{M(H^{+})} + \Lambda^{\circ}_{M(CH_{3}COO^{-})}$ $= 350 + 50 = 400 \text{ S cm}^2 \text{ mol}^{-1}$ $\alpha = \frac{\Lambda_M^C}{\Lambda_M^c}$ $a = \frac{20}{400} = 5 \times 10^{-2}$ $K_{a(CH_2COOH)} = C\alpha^2$ $= 0.007 \times (5 \times 10^{-2})^2 = 1.75 \times 10^{-5} mol L^{-1}$ S13. Ans.(a) The following reactions take place on electrolysis of dilute sulphuric acid on using pt electrodes are: At cathode: $4H^+ + 4e^-(aq) \rightarrow 2H_2(q)$ At anode: $2H_2O(l) \rightarrow O_2(g) + 4H^+(aq) + 4e^-$ Thus, the product obtained at anode is O_2 . S14. Ans.(d) 1 equivalent of any substance is deposited by 1 F of charge. We have, 20 g calcium The balance reaction will The charge on Ca in CaCl₂ Cl has -1 charge so that Ca + 2(-1) = 0Ca = 2We have to get Ca from Ca²⁺

Number of required moles = mass/molar mass Molar mass of Ca is 40 g/mol and required of Ca is 20 g Hence number of moles = 20/40= 0.5 molElectricity required to produce 1 mol of calcium = 2FThe electricity required to produce 0.5 mol of calcium = $0.5 \times 2F$ = 1FS15. Ans.(b) $Au^{3+} + 3e^- \rightarrow Au(s) E^\circ = 1.40 V$ $Fe^{2+} + 2e^- \rightarrow Fe(s) E^\circ = -0.44 V$ $Mg^{2+} + 2e^- \rightarrow Mg(s) E^\circ = -2.36 V$ $K^+ + 1e^- \rightarrow K(s) E^\circ = -2.93 V$ Au^{3+} occupies the top position in the electrochemical series. S16. Ans.(a) Cell reaction involved in hydrogenoxygen fuel cell is $2H_2(g) + O_2(g) \rightarrow 2H_2O(l)$ Thus, $R = H_2(g), O_2(g); P = H_2O(l)$ S17. Ans.(c) $E_{cell} = E^{\circ}_{cell} - \frac{0.059}{n} \log Q$...(i) At equilibrium, $Q = K_{eq}$ and $E_{cell} = 0$) $0^{\circ}E^{\circ}_{cell} - \frac{0.059}{1} log K_{eq}$ (from equation (i)) $log K_{eq} = \frac{E^{\circ}_{cell}}{0.059} = \frac{0.59}{0.059} = 10$ $K_{eq} = 10^{10} = 1 \times 10^{10}$ S18. Ans.(a) $\Delta G^{\Theta} = -nF E_{cell}^{\Theta}$ $= -2 \times 96500 \times 0.24$ $= -46320 \, Jmol^{-1} = \frac{-46320}{1000}$ $= -46.32 \, kJ/mol$ S19. Ans.(c) $\overset{+1}{HBrO} \rightarrow \overset{0}{Br_2}, E^0_{HBrO/Br_2} = 0.595\,V'$ $HBrO \rightarrow BrO_3^{+5}, E_{BrO_3/HBrO}^{\circ} = 1.5 V$ E_{cell}° for the disproportionation of *HBrO*,

$$\begin{split} E_{cell}^{*} & = E_{HB^{*}O[B^{*}_{2}} - E_{B^{*}O_{2}^{*}/HB^{*}O} \\ & = 1.595 - 1.5 \\ & = 0.095 V = +ve \\ \text{Hence, option (c) is correct answer.} \\ \text{S20. Ans.(d)} \\ & Zn[ZnSO_{4}(0.01M)]|[CuSO_{4}(1.0M)]Cu \\ & \therefore E_{1} = E_{cell}^{*} - \frac{2.303R^{*}}{2xF} \times \log\frac{(0.02)}{1} \\ \text{When concentrations are changed} \\ & \therefore E_{2} = E_{cell}^{*} - \frac{2.303R^{*}}{2xF} \times \log\frac{1}{0.01} \\ & i.e. E_{1} > E_{2} \\ \text{S21. Ans.(c)} \\ & \Lambda^{*}_{m}(AgCl) = \Lambda^{*}_{m}(AgNO_{3}) + \Lambda^{*}_{m}(KCl) - \\ & \Lambda^{*}_{m}(KNO_{3}) \\ & = (133.4 + 149 - 144.9) S cm^{2} mol^{-1} \\ & = 138.4 S cm^{2} mol^{-1} \\ \text{S22. Ans.(c)} \\ & Zn^{2^{*}} + 2e^{-} \rightarrow Zn, E^{\circ} = -0.760 V \\ & Ag_{2}O + H_{2}O + 2e^{-} \rightarrow 2Ag + 2OH^{-}, \\ & E^{\circ} = 0.344 V \\ \text{Both are reducing potential :} \\ & As, \quad E_{Ag_{2}O}^{*} 2Ag^{2} + E_{2}^{*} + 2e^{-} E^{\circ} = +0.760 V \\ & At cathode; \\ & Ag_{2}O + H_{2}O + 2e^{-} \rightarrow 2Ag + 2OH^{-}, \\ & E^{\circ} = +0.344 V \\ \text{Cell reaction:} \\ & Zn + Ag_{2}O + H_{2}O \rightarrow Zn^{2^{+}} + 2Ag + 2OH^{-} \\ & n = 2 \qquad E_{cell}^{*} = 1.104 V \\ & \ddots \ \Delta G^{\circ} = -nFE_{cell}^{*} \\ & \Delta G^{\circ} = -2 \times 96500 \times 1.10 \\ & = -213.072 J mol^{-1} \\ & = -213.072 J mol^{-1} \\ & = -213.072 kJ mol^{-1} \\ & \text{S23. Ans.(d) \\ & \text{According to Faraday's first law:} \\ & w = z.i.t \\ & z = \frac{E}{96500} (molar mass) \\ \end{aligned}$$

 $1 = \frac{35.5}{96500} \times 3 \times t \qquad \text{as x factor } 2$

min

e a higher (-ve) electrode al that is more reactive than Fe. ated on iron substances to e resistance against rusting such ess is called galvanization. But in e, that is Fe cannot be coated on corrosion will occur. In above, Zn es Fe from its salt solution.

ing to Faraday's law $\frac{\times 60}{(10^{-19})} = 3.75 \times 10^{20} \ electrons$ $-nFE^{\circ} cell \qquad E^{\circ} cell = (-ve)$ $= (+ve) \quad \Delta G > 0$ $G^{\circ} = -2.303 RT \log K_{ea}$ < 1 ntration = 0.5 mol dm⁻³, '6 × 10⁻³ S cm⁻¹ 8 K ×1000 М $\frac{10^{-3}}{2} = 11.52 \, S \, cm^2 \, mol^{-1}$ $2e^- \rightarrow H_2$ $-\frac{0.059}{2}\log\frac{P_{H_2}}{(H^+)^2}$ $\frac{0.059}{2} \log \frac{P_{H_2}}{(10^{-7})^2}$ $(10^{-7})^2 = 10^{-14} atm$.

> nnot reduce Fe^{2+} , so $FeCl_2$ and can exists together.

S31. Ans.(a)

$$\Delta G^{\circ} = -2.303 RT \log K_{sp}$$

$$63300 = -2.303 \times 8.314 \times 298 \log K_{sp}$$

$$K_{sp} \sim 8 \times 10^{-12}$$
S32. Ans.(c)

$$\stackrel{+6}{MnO_4^{-2}} \rightarrow MnO_4^{-} + e^{-}$$
0.1 mole 0.1 mole

$$Q = nF = 0.1 \times 96500 = 9650 C$$
S33. Ans.(c)

$$W_{02} = \frac{5600}{22400} \times 32 = 8g = 1 \ equivalent$$

$$= 1 \ equivalent \ of \ Ag$$

$$= 108 \ g$$

S34. Ans.(c)

According to Kohlrausch's law of limiting molar conductivity:

 $\alpha = \frac{\Lambda_m}{\Lambda_m^{\circ}} = \frac{9.54}{238} = 0.04008$ % Dissociation (\alpha) = 4.008%

S35. Ans.(b)

According to Nernst equation

$$\begin{split} E_{oxi} &= E_{red} - \frac{0.0592}{n} \log \frac{[Cation]}{[Anion]} \\ &= 0 - \frac{0.592}{2} \log \frac{[10^{-10}]}{[1]} \\ pH &= 0.59 \text{ V} \end{split}$$

 $E^{\circ}cell = E^{\circ}_{cathode} - E^{\circ}_{anode} = 0.76 - (-0.34)$ = 1.1 V