

# Chapter 17

# Electrochemistry

- In the electrochemical cell :  $\text{Zn}|\text{ZnSO}_4(0.01\text{ M})||\text{CuSO}_4(1.0\text{ M})|\text{Cu}$ , the emf of this Daniell cell is  $E_1$ . When the concentration of  $\text{ZnSO}_4$  is changed to 1.0 M and that of  $\text{CuSO}_4$  changed to 0.01 M, the emf changes to  $E_2$ . From the followings, which one is the relationship between  $E_1$  and  $E_2$ ? (Given,  $RT/F = 0.059$ )
  - $E_1 < E_2$
  - $E_1 > E_2$
  - $E_2 = 0^1 E_1$
  - $E_1 = E_2$

(NEET 2017)
- The molar conductivity of a 0.5 mol/dm<sup>3</sup> solution of  $\text{AgNO}_3$  with electrolytic conductivity of  $5.76 \times 10^{-3}\text{ S cm}^{-1}$  at 298 K is
  - 2.88 S cm<sup>2</sup>/mol
  - 11.52 S cm<sup>2</sup>/mol
  - 0.086 S cm<sup>2</sup>/mol
  - 28.8 S cm<sup>2</sup>/mol

(NEET-II 2016)
- During the electrolysis of molten sodium chloride, the time required to produce 0.10 mol of chlorine gas using a current of 3 amperes is
  - 55 minutes
  - 110 minutes
  - 220 minutes
  - 330 minutes

(NEET-II 2016)
- If the  $E^\circ_{\text{cell}}$  for a given reaction has a negative value, which of the following gives the correct relationships for the values of  $\Delta G^\circ$  and  $K_{\text{eq}}$ ?
  - $\Delta G^\circ > 0; K_{\text{eq}} < 1$
  - $\Delta G^\circ > 0; K_{\text{eq}} > 1$
  - $\Delta G^\circ < 0; K_{\text{eq}} > 1$
  - $\Delta G^\circ < 0; K_{\text{eq}} < 1$

(NEET-II 2016, 2011)
- The number of electrons delivered at the cathode during electrolysis by a current of 1 ampere in 60 seconds is (charge on electron =  $1.60 \times 10^{-19}\text{ C}$ )
  - $6 \times 10^{23}$
  - $6 \times 10^{20}$
  - $3.75 \times 10^{20}$
  - $7.48 \times 10^{23}$

(NEET-II 2016)
- Zinc can be coated on iron to produce galvanized iron but the reverse is not possible. It is because
  - zinc is lighter than iron
  - zinc has lower melting point than iron
  - zinc has lower negative electrode potential than iron
  - zinc has higher negative electrode potential than iron.

(NEET-II 2016)
- The pressure of  $\text{H}_2$  required to make the potential of  $\text{H}_2$ -electrode zero in pure water at 298 K is
  - $10^{-10}\text{ atm}$
  - $10^{-4}\text{ atm}$
  - $10^{-14}\text{ atm}$
  - $10^{-12}\text{ atm}$

(NEET-I 2016)
- A device that converts energy of combustion of fuels like hydrogen and methane, directly into electrical energy is known as
  - dynamo
  - Ni-Cd cell
  - fuel cell
  - electrolytic cell.

(2015, Cancelled)
- When 0.1 mol  $\text{MnO}_4^{2-}$  is oxidised the quantity of electricity required to completely oxidise  $\text{MnO}_4^{2-}$  to  $\text{MnO}_4^-$  is
  - 96500 C
  - $2 \times 96500\text{ C}$
  - 9650 C
  - 96.50 C

(2014)
- The weight of silver (at. wt. = 108) displaced by a quantity of electricity which displaces 5600 mL of  $\text{O}_2$  at STP will be
  - 5.4 g
  - 10.8 g
  - 54.0 g
  - 108.0 g

(2014)
- At 25°C molar conductance of 0.1 molar aqueous solution of ammonium hydroxide is  $9.54\text{ ohm}^{-1}\text{ cm}^2\text{ mol}^{-1}$  and at infinite dilution its molar conductance is  $238\text{ ohm}^{-1}\text{ cm}^2\text{ mol}^{-1}$ . The degree of ionisation of ammonium hydroxide at the same concentration and temperature is
  - 4.008%
  - 40.800%
  - 2.080%
  - 20.800%

(NEET 2013)
- A button cell used in watches function as following.
 
$$\text{Zn}_{(s)} + \text{Ag}_2\text{O}_{(s)} + \text{H}_2\text{O}_{(l)} \rightleftharpoons 2\text{Ag}_{(s)} + \text{Zn}^{2+}_{(aq)} + 2\text{OH}^{-}_{(aq)}$$

If half cell potentials are

$$\text{Zn}^{2+}_{(aq)} + 2e^- \rightarrow \text{Zn}_{(s)}; E^\circ = -0.76\text{ V}$$

$$\text{Ag}_2\text{O}_{(s)} + \text{H}_2\text{O}_{(l)} + 2e^- \rightarrow 2\text{Ag}_{(s)} + 2\text{OH}^{-}_{(aq)}; E^\circ = 0.34\text{ V}$$

The cell potential will be

  - 0.84 V
  - 1.34 V
  - 1.10 V
  - 0.42 V

(NEET 2013)

13. 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  
 (a) 0.118 V (b) 1.18 V  
 (c) 0.059 V (d) 0.59 V  
 (NEET 2013)
14. Consider the half-cell reduction reaction  
 $\text{Mn}^{2+} + 2e^- \rightarrow \text{Mn}$ ,  $E^\circ = -1.18 \text{ V}$   
 $\text{Mn}^{2+} \rightarrow \text{Mn}^{3+} + e^-$ ,  $E^\circ = -1.51 \text{ V}$   
 The  $E^\circ$  for the reaction  $3\text{Mn}^{2+} \rightarrow \text{Mn}^0 + 2\text{Mn}^{3+}$ , and possibility of the forward reaction are respectively  
 (a) -4.18 V and yes (b) +0.33 V and yes  
 (c) +2.69 V and no (d) -2.69 V and no  
 (Karnataka NEET 2013)
15. How many gram of cobalt metal will be deposited when a solution of cobalt(II) chloride is electrolyzed with a current of 10 amperes for 109 minutes (1 Faraday = 96,500 C; Atomic mass of Co = 59 u)  
 (a) 4.0 (b) 20.0  
 (c) 40.0 (d) 0.66  
 (Karnataka NEET 2013)
16. Limiting molar conductivity of  $\text{NH}_4\text{OH}$  [i.e.  $\Lambda_m^\circ(\text{NH}_4\text{OH})$ ] is equal to  
 (a)  $\Lambda_m^\circ(\text{NH}_4\text{Cl}) + \Lambda_m^\circ(\text{NaCl}) - \Lambda_m^\circ(\text{NaOH})$   
 (b)  $\Lambda_m^\circ(\text{NaOH}) + \Lambda_m^\circ(\text{NaCl}) - \Lambda_m^\circ(\text{NH}_4\text{Cl})$   
 (c)  $\Lambda_m^\circ(\text{NH}_4\text{OH}) + \Lambda_m^\circ(\text{NH}_4\text{Cl}) - \Lambda_m^\circ(\text{HCl})$   
 (d)  $\Lambda_m^\circ(\text{NH}_4\text{Cl}) + \Lambda_m^\circ(\text{NaOH}) - \Lambda_m^\circ(\text{NaCl})$  (2012)
17. Standard reduction potentials of the half reactions are given below :  
 $\text{F}_2(\text{g}) + 2e^- \rightarrow 2\text{F}^-(\text{aq})$ ;  $E^\circ = +2.85 \text{ V}$   
 $\text{Cl}_2(\text{g}) + 2e^- \rightarrow 2\text{Cl}^-(\text{aq})$ ;  $E^\circ = +1.36 \text{ V}$   
 $\text{Br}_2(\text{l}) + 2e^- \rightarrow 2\text{Br}^-(\text{aq})$ ;  $E^\circ = +1.06 \text{ V}$   
 $\text{I}_2(\text{s}) + 2e^- \rightarrow 2\text{I}^-(\text{aq})$ ;  $E^\circ = +0.53 \text{ V}$   
 The strongest oxidising and reducing agents respectively are  
 (a)  $\text{F}_2$  and  $\text{I}^-$  (b)  $\text{Br}_2$  and  $\text{Cl}^-$   
 (c)  $\text{Cl}_2$  and  $\text{Br}^-$  (d)  $\text{Cl}_2$  and  $\text{I}_2$   
 (Mains 2012)
18. Molar conductivities ( $\Lambda_m^\circ$ ) at infinite dilution of NaCl, HCl and  $\text{CH}_3\text{COONa}$  are 126.4, 425.9 and 91.0  $\text{S cm}^2 \text{ mol}^{-1}$  respectively. ( $\Lambda_m^\circ$ ) for  $\text{CH}_3\text{COOH}$  will be  
 (a) 425.5  $\text{S cm}^2 \text{ mol}^{-1}$  (b) 180.5  $\text{S cm}^2 \text{ mol}^{-1}$   
 (c) 290.8  $\text{S cm}^2 \text{ mol}^{-1}$  (d) 390.5  $\text{S cm}^2 \text{ mol}^{-1}$   
 (Mains 2012)
19. The Gibb's energy for the decomposition of  $\text{Al}_2\text{O}_3$  at  $500^\circ\text{C}$  is as follows  

$$\frac{2}{3}\text{Al}_2\text{O}_3 \rightarrow \frac{4}{3}\text{Al} + \text{O}_2$$

$$\Delta_r G = +960 \text{ kJ mol}^{-1}$$
 The potential difference needed for the electrolytic reduction of aluminium oxide ( $\text{Al}_2\text{O}_3$ ) at  $500^\circ\text{C}$  is at least  
 (a) 4.5 V (b) 3.0 V  
 (c) 2.5 V (d) 5.0 V  
 (Mains 2012)
20. Standard electrode potential of three metals X, Y and Z are -1.2 V, +0.5 V and -3.0 V respectively. The reducing power of these metals will be  
 (a)  $Y > Z > X$  (b)  $Y > X > Z$   
 (c)  $Z > X > Y$  (d)  $X > Y > Z$   
 (2011)
21. The electrode potentials for  
 $\text{Cu}^{2+}(\text{aq}) + e^- \rightarrow \text{Cu}^+(\text{aq})$  and  $\text{Cu}^+(\text{aq}) + e^- \rightarrow \text{Cu}(\text{s})$   
 are +0.15 V and +0.50 V respectively. The value of  $E^\circ_{\text{Cu}^{2+}/\text{Cu}}$  will be  
 (a) 0.500 V (b) 0.325 V  
 (c) 0.650 V (d) 0.150 V (2011)
22. Standard electrode potential for  $\text{Sn}^{4+}/\text{Sn}^{2+}$  couple is +0.15 V and that for the  $\text{Cr}^{3+}/\text{Cr}$  couple is -0.74 V. These two couples in their standard state are connected to make a cell. The cell potential will be  
 (a) +1.19 V (b) +0.89 V  
 (c) +0.18 V (d) +1.83 V (2011)
23. A solution contains  $\text{Fe}^{2+}$ ,  $\text{Fe}^{3+}$  and  $\text{I}^-$  ions. This solution was treated with iodine at  $35^\circ\text{C}$ .  $E^\circ$  for  $\text{Fe}^{3+}/\text{Fe}^{2+}$  is +0.77 V and  $E^\circ$  for  $\text{I}_2/2\text{I}^- = 0.536 \text{ V}$ . The favourable redox reaction is  
 (a)  $\text{I}_2$  will be reduced to  $\text{I}^-$   
 (b) there will be no redox reaction  
 (c)  $\text{I}^-$  will be oxidised to  $\text{I}_2$   
 (d)  $\text{Fe}^{2+}$  will be oxidised to  $\text{Fe}^{3+}$   
 (Mains 2011)
24. For the reduction of silver ions with copper metal, the standard cell potential was found to be +0.46 V at  $25^\circ\text{C}$ . The value of standard Gibb's energy,  $\Delta G^\circ$  will be  
 ( $F = 96500 \text{ C mol}^{-1}$ )  
 (a) -89.0 kJ (b) -89.0 J  
 (c) -44.5 kJ (d) -98.0 kJ (2010)

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25. An increase in equivalent conductance of a strong electrolyte with dilution is mainly due to  
 (a) increase in ionic mobility of ions  
 (b) 100% ionisation of electrolyte at normal dilution  
 (c) increase in both *i.e.*, number of ions and ionic mobility of ions  
 (d) increase in number of ions. (2010)
26. Which of the following expressions correctly represents the equivalent conductance at infinite dilution of  $\text{Al}_2(\text{SO}_4)_3$ . Given that  $\overset{\circ}{\Lambda}_{\text{Al}^{3+}}$  and  $\overset{\circ}{\Lambda}_{\text{SO}_4^{2-}}$  are the equivalent conductances at infinite dilution of the respective ions?  
 (a)  $2\overset{\circ}{\Lambda}_{\text{Al}^{3+}} + 3\overset{\circ}{\Lambda}_{\text{SO}_4^{2-}}$   
 (b)  $\overset{\circ}{\Lambda}_{\text{Al}^{3+}} + \overset{\circ}{\Lambda}_{\text{SO}_4^{2-}}$   
 (c)  $(\overset{\circ}{\Lambda}_{\text{Al}^{3+}} + \overset{\circ}{\Lambda}_{\text{SO}_4^{2-}}) \times 6$   
 (d)  $\frac{1}{3}\overset{\circ}{\Lambda}_{\text{Al}^{3+}} + \frac{1}{2}\overset{\circ}{\Lambda}_{\text{SO}_4^{2-}}$  (Mains 2010)
27. Consider the following relations for emf of an electrochemical cell  
 (i) EMF of cell = (Oxidation potential of anode) – (Reduction potential of cathode)  
 (ii) EMF of cell = (Oxidation potential of anode) + (Reduction potential of cathode)  
 (iii) EMF of cell = (Reductional potential of anode) + (Reduction potential of cathode)  
 (iv) EMF of cell = (Oxidation potential of anode) – (Oxidation potential of cathode)  
 Which of the above relations are correct?  
 (a) (iii) and (i) (b) (i) and (ii)  
 (c) (iii) and (iv) (d) (ii) and (iv)  
 (Mains 2010)
28. Given :  
 (i)  $\text{Cu}^{2+} + 2e^- \rightarrow \text{Cu}$ ,  $E^\circ = 0.337 \text{ V}$   
 (ii)  $\text{Cu}^{2+} + e^- \rightarrow \text{Cu}^+$ ,  $E^\circ = 0.153 \text{ V}$   
 Electrode potential,  $E^\circ$  for the reaction,  
 $\text{Cu}^+ + e^- \rightarrow \text{Cu}$ , will be  
 (a) 0.90 V (b) 0.30 V  
 (c) 0.38 V (d) 0.52 V (2009)
29.  $\text{Al}_2\text{O}_3$  is reduced by electrolysis at low potentials and high currents. If  $4.0 \times 10^4$  amperes of current is passed through molten  $\text{Al}_2\text{O}_3$  for 6 hours, what mass of aluminium is produced? (Assume 100% current efficiency, at. mass of Al =  $27 \text{ g mol}^{-1}$ ).  
 (a)  $8.1 \times 10^4 \text{ g}$  (b)  $2.4 \times 10^5 \text{ g}$   
 (c)  $1.3 \times 10^4 \text{ g}$  (d)  $9.0 \times 10^3 \text{ g}$  (2009)
30. The equivalent conductance of  $M/32$  solution of a weak monobasic acid is  $8.0 \text{ mho cm}^2$  and at infinite dilution is  $400 \text{ mho cm}^2$ . The dissociation constant of this acid is  
 (a)  $1.25 \times 10^{-6}$  (b)  $6.25 \times 10^{-4}$   
 (c)  $1.25 \times 10^{-4}$  (d)  $1.25 \times 10^{-5}$  (2009)
31. On the basis of the following  $E^\circ$  values, the strongest oxidizing agent is  
 $[\text{Fe}(\text{CN})_6]^{4-} \rightarrow [\text{Fe}(\text{CN})_6]^{3-} + e^-$ ;  $E^\circ = -0.35 \text{ V}$   
 $\text{Fe}^{2+} \rightarrow \text{Fe}^{3+} + e^-$ ;  $E^\circ = -0.77 \text{ V}$   
 (a)  $\text{Fe}^{3+}$  (b)  $[\text{Fe}(\text{CN})_6]^{3-}$   
 (c)  $[\text{Fe}(\text{CN})_6]^{4-}$  (d)  $\text{Fe}^{2+}$  (2008)
32. Kohlrausch's law states that at  
 (a) Infinite dilution, each ion makes definite contribution to conductance of an electrolyte whatever be the nature of the other ion of the electrolyte  
 (b) Infinite dilution, each ion makes definite contribution to equivalent conductance of an electrolyte, whatever be the nature of the other ion of the electrolyte  
 (c) Finite dilution, each ion makes definite contribution to equivalent conductance of an electrolyte, whatever be the nature of the other ion of the electrolyte  
 (d) Infinite dilution each ion makes definite contribution to equivalent conductance of an electrolyte depending on the nature of the other ion of the electrolyte. (2008)
33. Standard free energies of formation (in kJ/mol) at 298 K are  $-237.2$ ,  $-394.4$  and  $-8.2$  for  $\text{H}_2\text{O}_{(l)}$ ,  $\text{CO}_{2(g)}$  and pentane  $_{(g)}$  respectively. The value of  $E^\circ_{\text{cell}}$  for the pentane-oxygen fuel cell is  
 (a) 1.0968 V (b) 0.0968 V  
 (c) 1.968 V (d) 2.0968 V (2008)
34. The equilibrium constant of the reaction:  
 $\text{Cu}_{(s)} + 2\text{Ag}^+_{(aq)} \rightarrow \text{Cu}^{2+}_{(aq)} + 2\text{Ag}_{(s)}$ ;  
 $E^\circ = 0.46 \text{ V}$  at 298 K is  
 (a)  $2.0 \times 10^{10}$  (b)  $4.0 \times 10^{10}$   
 (c)  $4.0 \times 10^{15}$  (d)  $2.4 \times 10^{10}$  (2007)
35. The efficiency of a fuel cell is given by  
 (a)  $\Delta G/\Delta S$  (b)  $\Delta G/\Delta H$   
 (c)  $\Delta S/\Delta G$  (d)  $\Delta H/\Delta G$  (2007)

36. A hypothetical electrochemical cell is shown below.  
 $A | A^+ (xM) || B^+ (yM) | B$   
 The emf measured is +0.20 V. The cell reaction is  
 (a)  $A + B^+ \rightarrow A^+ + B$   
 (b)  $A^+ + B \rightarrow A + B^+$   
 (c)  $A^+ + e^- \rightarrow A; B^+ + e^- \rightarrow B$   
 (d) the cell reaction cannot be predicted. (2006)
37.  $E^\circ_{Fe^{2+}/Fe} = -0.441$  V and  $E^\circ_{Fe^{3+}/Fe^{2+}} = 0.771$  V, the standard EMF of the reaction  $Fe + 2Fe^{3+} \rightarrow 3Fe^{2+}$  will be  
 (a) 0.111 V (b) 0.330 V  
 (c) 1.653 V (d) 1.212 V (2006)
38. 4.5 g of aluminium (at. mass 27 amu) is deposited at cathode from  $Al^{3+}$  solution by a certain quantity of electric charge. The volume of hydrogen produced at STP from  $H^+$  ions in solution by the same quantity of electric charge will be  
 (a) 44.8 L (b) 22.4 L  
 (c) 11.2 L (d) 5.6 L (2005)
39. The mass of carbon anode consumed (giving only carbon dioxide) in the production of 270 kg of aluminium metal from bauxite by the Hall process is  
 (a) 270 kg (b) 540 kg  
 (c) 90 kg (d) 180 kg  
 (Atomic mass : Al = 27) (2003)
40. The standard e.m.f. of a galvanic cell involving cell reaction with  $n = 2$  is found to be 0.295 V at 25°C. The equilibrium constant of the reaction would be  
 (a)  $2.0 \times 10^{11}$  (b)  $4.0 \times 10^{12}$   
 (c)  $1.0 \times 10^2$  (d)  $1.0 \times 10^{10}$   
 (Given  $F = 96500$  C mol<sup>-1</sup>,  $R = 8.314$  J K<sup>-1</sup> mol<sup>-1</sup>) (2004)
41. The e.m.f. of a Daniell cell at 298 K is  $E_1$ .  

$$Zn | ZnSO_4 (0.01 M) || CuSO_4 (1.0 M) | Cu$$
  
 When the concentration of  $ZnSO_4$  is 1.0 M and that of  $CuSO_4$  is 0.01 M, the e.m.f. changed to  $E_2$ . What is the relationship between  $E_1$  and  $E_2$ ?  
 (a)  $E_1 > E_2$  (b)  $E_1 < E_2$   
 (c)  $E_1 = E_2$  (d)  $E_2 = 0 \neq E_1$  (2003)
42. On the basis of the information available from the reaction,  
 $4/3Al + O_2 \rightarrow 2/3Al_2O_3, \Delta G = -827$  kJ mol<sup>-1</sup> of  $O_2$ , the minimum e.m.f. required to carry out an electrolysis of  $Al_2O_3$  is ( $F = 96500$  C mol<sup>-1</sup>)  
 (a) 2.14 V (b) 4.28 V  
 (c) 6.42 V (d) 8.56 V (2003)
43. In electrolysis of NaCl when Pt electrode is taken then  $H_2$  is liberated at cathode while with Hg cathode it forms sodium amalgam  
 (a) Hg is more inert than Pt  
 (b) More voltage is required to reduce  $H^+$  at Hg than at Pt  
 (c) Na is dissolved in Hg while it does not dissolve in Pt  
 (d) Conc. of  $H^+$  ions is larger when Pt electrode is taken. (2002)
44. Standard electrode potentials are  $Fe^{2+}/Fe$ ,  $E^\circ = -0.44$  and  $Fe^{3+}/Fe^{2+}$ ,  $E^\circ = 0.77$  V,  $Fe^{2+}$  and Fe blocks are kept together, then  
 (a)  $Fe^{3+}$  increases  
 (b)  $Fe^{3+}$  decreases  
 (c)  $Fe^{2+}/Fe^{3+}$  remains unchanged  
 (d)  $Fe^{2+}$  decreases. (2001)
45. Equivalent conductances of  $Ba^{2+}$  and  $Cl^-$  ions are 127 and 76 ohm<sup>-1</sup> cm<sup>-1</sup> eq<sup>-1</sup> respectively. Equivalent conductance of  $BaCl_2$  at infinite dilution is  
 (a) 139.5 (b) 101.5  
 (c) 203 (d) 279 (2000)
46. For the disproportionation of copper  
 $2Cu^+ \rightarrow Cu^{2+} + Cu$ ,  $E^\circ$  is (Given  $E^\circ$  for  $Cu^{2+}/Cu$  is 0.34 V and  $E^\circ$  for  $Cu^{2+}/Cu^+$  is 0.15 V.)  
 (a) 0.49 V (b) -0.19 V  
 (c) 0.38 V (d) -0.38 V (2000)
47. The specific conductance of a 0.1 N KCl solution at 23°C is 0.012 ohm<sup>-1</sup> cm<sup>-1</sup>. The resistance of cell containing the solution at the same temperature was found to be 55 ohm. The cell constant will be  
 (a) 0.918 cm<sup>-1</sup> (b) 0.66 cm<sup>-1</sup>  
 (c) 1.142 cm<sup>-1</sup> (d) 1.12 cm<sup>-1</sup> (1999)
48. For the cell reaction,  
 $Cu^{2+} (C_1, aq) + Zn_{(s)} = Zn^{2+} (C_2, aq) + Cu_{(s)}$   
 of an electrochemical cell, the change in free energy  $\Delta G$  at a given temperature is a function of  
 (a)  $\ln (C_2)$  (b)  $\ln (C_2/C_1)$   
 (c)  $\ln (C_1)$  (d)  $\ln (C_1 + C_2)$  (1998)
49.  $E^\circ$  for the cell,  $Zn | Zn^{2+} (aq) || Cu^{2+} (aq) | Cu$  is 1.10V at 25°C, the equilibrium constant for the reaction  $Zn + Cu^{2+} (aq) \rightleftharpoons Cu + Zn^{2+} (aq)$  is the order of  
 (a)  $10^{+18}$  (b)  $10^{+17}$   
 (c)  $10^{-28}$  (d)  $10^{-37}$  (1997)

Electrochemistry

50. The molar conductances of NaCl, HCl and  $\text{CH}_3\text{COONa}$  at infinite dilution are 126.45, 426.16 and  $91 \text{ ohm}^{-1} \text{ cm}^2 \text{ mol}^{-1}$  respectively. The molar conductance of  $\text{CH}_3\text{COOH}$  at infinite dilution ( $\Lambda_m^\infty$ ) is  
 (a)  $698.28 \text{ ohm}^{-1} \text{ cm}^2 \text{ mol}^{-1}$   
 (b)  $540.48 \text{ ohm}^{-1} \text{ cm}^2 \text{ mol}^{-1}$   
 (c)  $201.28 \text{ ohm}^{-1} \text{ cm}^2 \text{ mol}^{-1}$   
 (d)  $390.71 \text{ ohm}^{-1} \text{ cm}^2 \text{ mol}^{-1}$  (1997)
51. A 5 ampere current is passed through a solution of zinc sulphate for 40 minutes. The amount of zinc deposited at the cathode is  
 (a) 0.4065 g (b) 65.04 g  
 (c) 40.65 g (d) 4.065 g (1996)
52. Reduction potential for the following half-cell reactions are  
 $\text{Zn} = \text{Zn}^{2+} + 2e^-$ ;  $E^\circ = +0.76 \text{ V}$ ;  
 $\text{Fe} = \text{Fe}^{2+} + 2e^-$ ;  $E^\circ = +0.44 \text{ V}$ .  
 The EMF for the cell reaction  
 $\text{Fe}^{2+} + \text{Zn} \rightarrow \text{Zn}^{2+} + \text{Fe}$  will be  
 (a)  $-0.32 \text{ V}$  (b)  $+1.20 \text{ V}$   
 (c)  $-1.20 \text{ V}$  (d)  $+0.32 \text{ V}$  (1996)
53. An electrochemical cell is set up as : Pt;  $\text{H}_2$  (1 atm)|HCl(0.1 M) ||  $\text{CH}_3\text{COOH}$  (0.1 M) | $\text{H}_2$  (1 atm); Pt. The e.m.f. of this cell will not be zero, because  
 (a) acids used in two compartments are different  
 (b) e.m.f. depends on molarities of acids used  
 (c) the temperature is constant  
 (d) pH of 0.1 M HCl and 0.1 M  $\text{CH}_3\text{COOH}$  is not same. (1995)
54. On heating one end of a piece of a metal, the other end becomes hot because of  
 (a) energised electrons moving to the other end  
 (b) minor perturbation in the energy of atoms  
 (c) resistance of the metal  
 (d) mobility of atoms in the metal. (1995)
55. Standard reduction potentials at  $25^\circ\text{C}$  of  $\text{Li}^+|\text{Li}$ ,  $\text{Ba}^{2+}|\text{Ba}$ ,  $\text{Na}^+|\text{Na}$  and  $\text{Mg}^{2+}|\text{Mg}$  are  $-3.05$ ,  $-2.90$ ,  $-2.71$  and  $-2.37$  volt respectively. Which one of the following is the strongest oxidising agent?  
 (a)  $\text{Ba}^{2+}$  (b)  $\text{Mg}^{2+}$   
 (c) Na (d)  $\text{Li}^+$  (1994)
56. On electrolysis of dilute sulphuric acid using platinum electrodes, the product obtained at the anode will be  
 (a) hydrogen (b) oxygen  
 (c) hydrogen sulphide (d) sulphur dioxide. (1992)

Answer Key

1. (b) 2. (b) 3. (b) 4. (a) 5. (c) 6. (d) 7. (c) 8. (c) 9. (c) 10. (d)  
 11. (a) 12. (c) 13. (d) 14. (d) 15. (b) 16. (d) 17. (a) 18. (d) 19. (c) 20. (c)  
 21. (b) 22. (b) 23. (c) 24. (a) 25. (a) 26. (b) 27. (d) 28. (d) 29. (a) 30. (d)  
 31. (a) 32. (a) 33. (a) 34. (c) 35. (b) 36. (a) 37. (d) 38. (d) 39. (c) 40. (d)  
 41. (a) 42. (a) 43. (b) 44. (b) 45. (a) 46. (c) 47. (b) 48. (b) 49. (d) 50. (d)  
 51. (d) 52. (d) 53. (d) 54. (a) 55. (b) 56. (b)