

### Numericals:

**1. A concentration cell was constructed using two iron electrodes in 0.01 M & 0.1 M Ferrous sulphate solution. Write the cell representation and calculate the emf of the cell.**

**Ans:**  $E_{\text{cell}} = 0.02955 \text{ V}$

**Given :**  $[\text{Fe}^{2+}] = 0.01 \text{ M}$

$$[\text{Fe}^{2+}] = 0.1 \text{ M}$$

$$E_{\text{cell}} = ?$$

Cell representation:  $\text{Fe} / \text{Fe}^{2+} (0.01\text{M}) // \text{Fe}^{2+} (0.1\text{M}) / \text{Fe}$

$$E_{\text{cell}} = 0.0591/n \log [0.1]/[0.01]$$

$$= 0.0591/2 (1)$$

$$= 0.02955 \text{ V}$$

**2. Represent the concentration cell formed by coupling two Cd electrodes immersed in two Cadmium sulphate solutions. Such that the concentration of  $\text{Cd}^{2+}$  ions in one electrode system is 0.05 M & that in the other system is 1.0 M, Calculate cell potential.**

**Ans:**  $E_{\text{cell}} = 0.03844 \text{ V}$

**Given :**  $[\text{Cd}^{2+}_{\text{anode}}] = 0.05 \text{ M}$ ,

$$[\text{Cd}^{2+}_{\text{cathode}}] = 1.0 \text{ M}$$

$$E_{\text{cell}} = ?$$

Cell =  $\text{Cd} / \text{Cd}^{2+} (0.05 \text{ M}) // \text{Cd}^{2+} (1.0 \text{ M}) / \text{Cd}$

$$n=2$$

$$E_{\text{cell}} = 0.0591/n \log [1.0]/[0.05]$$

$$= 0.02955 \log 20$$

$$= 0.02955 (1.3010)$$

$$= 0.03844 \text{ V}$$

**3. The spontaneous galvanic cell Tin/Tin ion (0.024 M)//Tin ion (0.064 M)/Tin develops an emf of 0.0216 V at 25° C. Calculate the valency of tin.**

**Ans:**  $n = 1$

**Given:**  $E_{\text{cell}} = 0.0216 \text{ V}$

Cell :  $\text{Tin} / \text{Tin ion} (0.024 \text{ M}) // \text{Tin ion} (0.064 \text{ M}) / \text{Tin}$

$$n = ?$$

$$E_{\text{cell}} = 0.0591/n \log [0.064 \text{ M}]/[0.024 \text{ M}]$$

$$0.0216 = 0.0591/n \log 2.6$$

$$\begin{aligned}
 n &= 0.0591/0.0216 \log 2.6 \\
 &= 2.736 (0.414) \\
 &= 1.13 \sim 1
 \end{aligned}$$

**4. In a spontaneous galvanic cell  $M/M^{n+} (0.03 \text{ N}) // M^{n+} (0.064 \text{ N})/M$  develops an emf of 0.013 V at 25° C. Calculate the valency of metal.**

**Ans:  $n = 1.5$**

**Given:  $E_{\text{cell}} = 0.013 \text{ V}$**

Cell:  $M/M^{n+} (0.03 \text{ N}) // M^{n+} (0.064 \text{ N})/M$

$n = ?$

$$E_{\text{cell}} = 0.0591/n \log [0.064 \text{ N}]/[0.03 \text{ N}]$$

$$0.013 = 0.0591/n \log 2.133$$

$$n = 0.0591/0.013 \log 2.133$$

$$= 4.546 (0.3289)$$

$$= 1.49 \sim 1.5$$

**5. Calculate the  $E_{\text{Zn}^{2+}/\text{Zn}}$  for the reaction  $\text{Zn}^{2+} + 2e \rightarrow \text{Zn}$ , Given:  $E^{\circ}_{\text{Zn}^{2+}/\text{Zn}} = -0.76 \text{ V}$ ,  $[\text{Zn}^{2+}] = 0.1 \text{ M}$**

**Ans:  $E_{\text{Zn}^{2+}/\text{Zn}} = -0.7895 \text{ V}$**

**Given:  $\text{Zn}^{2+} + 2e \rightarrow \text{Zn}$**

$$E^{\circ}_{\text{Zn}^{2+}/\text{Zn}} = -0.76 \text{ V},$$

$$[\text{Zn}^{2+}] = 0.1 \text{ M}$$

$$E_{\text{Zn}^{2+}/\text{Zn}} = ?$$

$$E_{\text{Zn}^{2+}/\text{Zn}} = E^{\circ}_{\text{Zn}^{2+}/\text{Zn}} + 0.0591/n \log [\text{Zn}^{2+}]$$

$$= -0.76 + 0.0591/2 \log [\text{Zn}^{2+}]$$

$$= -0.76 + 0.0295 \log [0.1]$$

$$= -0.76 + 0.0295 \log [-1]$$

$$= -0.7895 \text{ V}$$

**6. If two Zn electrodes were in contact with Zn ions, emf of the cell was found to be 0.065 V. If concentration of solution at cathode is 0.1 M, Calculate concentration of other solution.**

**Ans:  $[\text{Zn}^{2+}] = 6.32 \times 10^{-4} \text{ M}$**

**Given:**  $E_{\text{cell}} = 0.065 \text{ V}$ ,

$$[\text{Zn}^{2+}_{\text{cathode}}] = 0.1 \text{ M}$$

$$[\text{Zn}^{2+}_{\text{anode}}] = ?$$

Cell :  $\text{Zn}/\text{Zn}^{2+} (C_1 \text{ M}) // \text{Zn}^{2+} (0.1 \text{ M})/\text{Zn}$

$$E_{\text{cell}} = 0.0591/n \log [0.1]/[\text{Zn}^{2+}]$$

$$0.065 = 0.0591/2 \log [0.1]/[\text{Zn}^{2+}]$$

$$0.065 = 0.02955 \log [0.1]/[\text{Zn}^{2+}]$$

$$0.065/0.02955 = \log [0.1]/[\text{Zn}^{2+}]$$

$$\text{Antilog } (2.199) = \log [0.1]/[\text{Zn}^{2+}]$$

$$158.12 = \log [0.1]/[\text{Zn}^{2+}]$$

$$[\text{Zn}^{2+}] = 0.1/158.12$$

$$= 6.32 \times 10^{-4} \text{ M}$$

**7. When a Cu electrode is kept in contact with its solution at  $25^{\circ} \text{C}$ , electrode potential is found to be  $0.34 \text{ V}$ . Calculate concentration of solution. Given  $E^{\circ}_{\text{Cell}} = 0.33 \text{ V}$ .**

**Ans:**  $[\text{Cu}^{2+}] = 3.21 \text{ M}$

**Given:**  $E_{\text{Cell}} = 0.34 \text{ V}$

$$E^{\circ}_{\text{Cell}} = 0.33 \text{ V}$$

$$[\text{Cu}^{2+}] = ?$$

$$E_{\text{cell}} = E^{\circ}_{\text{Cell}} + 0.0591/n \log [\text{Cu}^{2+}]$$

$$0.345 = 0.33 + 0.0591/2 \log [\text{Cu}^{2+}]$$

$$0.345 - 0.33 = 0.0591/2 \log [\text{Cu}^{2+}]$$

$$0.015 = 0.02955 \log [\text{Cu}^{2+}]$$

$$0.015/0.02955 = \log [\text{Cu}^{2+}]$$

$$\text{Antilog } (0.5076) = \log [\text{Cu}^{2+}]$$

$$[\text{Cu}^{2+}] = 3.21 \text{ M}$$

**8.  $25 \text{ cm}^3$  of the hard water sample consumed  $10.6 \text{ cm}^3$  of  $0.01 \text{ M}$  EDTA. Calculate the total hardness.**

**Ans:**  $\text{TH} = 424 \text{ ppm}$

**Given:**  $V = 10.6 \text{ ml}$ , Molarity of EDTA =  $0.01 \text{ M}$

1000ml of 1 M EDTA= 100 g. of  $\text{CaCO}_3$

$$\begin{aligned}\text{Hence, } 10.6 \text{ ml of } 0.01 \text{ M EDTA} &= (10.6 \times 0.01 \times 100/1000) \\ &= 0.0106 \text{ g. of } \text{CaCO}_3\end{aligned}$$

$$\text{Hence, Total hardness} = [0.0106/25] 10^6 = 424 \text{ ppm}$$

**9. 25 cm<sup>3</sup> of the hard water sample consumed 12.6 cm<sup>3</sup> of 0.01 M EDTA. 25 cm<sup>3</sup> of the same water sample after boiling and filtering required 9.2 cm<sup>3</sup> of same EDTA for titration. Calculate the total hardness, temporary hardness and permanent hardness.**

**Ans: TH = 504 ppm, PH = 368 ppm and TempH = 136 ppm**

**Given:**  $V_1 = 12.6 \text{ ml}$ , Molarity of EDTA = 0.01 M

After boiling & filtering  $V_2 = 9.2 \text{ ml}$

$$\begin{aligned}12.6 \text{ ml of } 0.01 \text{ M EDTA} &= (12.6 \times 0.01 \times 100/1000) \\ &= 0.012 \text{ g. of } \text{CaCO}_3\end{aligned}$$

$$\text{Hence, Total hardness} = [0.012/25] 10^6 \text{ ppm} = 504 \text{ ppm}$$

$$\begin{aligned}9.2 \text{ ml of } 0.01 \text{ M EDTA} &= (9.2 \times 0.01 \times 100/1000) \\ &= 9.2 \times 10^{-3} \text{ g. of } \text{CaCO}_3.\end{aligned}$$

$$\text{Hence, Permanent hardness} = (9.2 \times 10^{-3}/25) 10^6 \text{ ppm} = 368 \text{ ppm}$$

$$\text{Temporary hardness} = \text{Total hardness} - \text{Permanent hardness}$$

$$= 504 - 368$$

$$= 136 \text{ ppm}$$

**10. Calculate the total & permanent hardness of the water sample. If 50 cm<sup>3</sup> of industrial hard water sample was required 20 cm<sup>3</sup> of M/50 (0.02M) EDTA salt solution. The same water sample after boiling & filtering required 13.7 cm<sup>3</sup> of the same EDTA salt solution with the same indicator.**

**Ans: TH = 800 ppm, PH = 548 ppm and TempH = 252 ppm**

**Given:**  $V_1 = 20 \text{ ml}$ , Molarity of EDTA = 0.02 M

After boiling & filtering  $V_2 = 13.7 \text{ ml}$

$$\begin{aligned}20 \text{ ml of } 0.02 \text{ M EDTA} &= (20 \times 0.02 \times 100/1000) \\ &= 0.04 \text{ g. of } \text{CaCO}_3\end{aligned}$$

$$\text{Hence, Total hardness} = [0.04/50] 10^6 = 800 \text{ ppm}$$

$$\begin{aligned}13.7 \text{ ml of } 0.02 \text{ M EDTA} &= (13.7 \times 0.02 \times 100/1000) \\ &= 0.0274 \text{ g. of } \text{CaCO}_3\end{aligned}$$

Hence, Permanent hardness =  $(0.0274/50)10^6 = 548 \text{ ppm}$

$$\begin{aligned}\text{Temporary hardness} &= \text{Total hardness} - \text{Permanent hardness} \\ &= 800 - 548 = 252 \text{ ppm}\end{aligned}$$

**11. 100 ml of water sample on titrating 0.01 M EDTA consumed 8.1 ml solution. Another 100 ml of sample after boiling for 20 minutes and filtering, the filtrate consumed 6.1 ml of same EDTA. Calculate total hardness and temporary hardness.**

**Ans: TH = 81 ppm, PH = 61 ppm and TemH = 20 ppm**

**Given:**  $V_1 = 8.1 \text{ ml}$ , Molarity of EDTA = 0.01 M

After boiling & filtering  $V_2 = 6.1 \text{ ml}$ , Water sample = 100 ml

$$\begin{aligned}8.1 \text{ ml of } 0.01 \text{ M EDTA} &= (8.1 \times 0.01 \times 100/1000) \\ &= 8.1 \times 10^{-3} \text{ g. of CaCO}_3\end{aligned}$$

$$\text{Hence, Total hardness} = [8.1 \times 10^{-3} / 100] 10^6 \text{ ppm} = 81 \text{ ppm}$$

$$\begin{aligned}6.1 \text{ ml of } 0.01 \text{ M EDTA} &= (6.1 \times 0.01 \times 100/1000) \\ &= 6.1 \times 10^{-3} \text{ g. of CaCO}_3 \text{ in } 25 \text{ ml water boiled.}\end{aligned}$$

$$\text{Hence, Permanent hardness} = (6.1 \times 10^{-3} / 100) 10^6 = 61 \text{ ppm}$$

$$\begin{aligned}\text{Temporary hardness} &= \text{Total hardness} - \text{Permanent hardness} \\ &= 81 - 61 = 20 \text{ ppm}\end{aligned}$$

**12. 20 ml of sewage sample is reacted with 25 ml of  $\text{K}_2\text{Cr}_2\text{O}_7$  solution and the unreacted  $\text{K}_2\text{Cr}_2\text{O}_7$  required 9.0 ml of 0.25 N FAS Solution. Under similar conditions in blank titration 15.0 ml of FAS is used up. Calculate the COD of the sample.**

**Ans: COD = 600 mg/dm<sup>3</sup>**

**Given :** B-A (15.0-9.0) = 6.0 ml , Normality of FAS = 0.25 N , Sewage sample = 20 ml

$$\begin{aligned}\text{Hence , normality of oxygen in waste water } N_1 &= [(15.0 - 9.0) / 20] \times 0.25 \\ &= 0.075 N_1\end{aligned}$$

$$\begin{aligned}\text{Hence, C.O.D of the sample} &= 0.075 \times 8 \times 1000 \\ &= 600 \text{ mg dm}^{-3}\end{aligned}$$

**13. Solve for COD of 25 ml of water sample if 29.5 ml and 20 ml of 0.025 N FAS was required for blank and sample titration respectively.**

**Ans: COD = 76 mg/dm<sup>3</sup>**

Given : B-A (29.5-20.0) = 9.5 ml , Normality of FAS = 0.025 N , Sewage sample = 25 ml

Hence , normality of oxygen in waste water  $N_1 = [(29.5-20.0)/25] \times 0.025$

$$= 9.5 \times 10^{-3} N_1$$

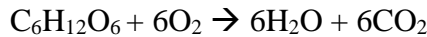
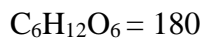
Hence, C.O.D of the sample =  $9.5 \times 10^{-3} \times 8 \times 1000$

$$= 76 \text{ mg dm}^{-3}$$

**14. Calculate the BOD value of a sewage sample containing 10.2 mg/ 200 ml of organic mater with the formula  $C_6H_{12}O_6$  (C = 12, H = 1, O = 16)**

**Ans: COD = 54.4 ppm**

**Given:** Sewage sample = 10.2 mg/ 200ml



$$\begin{array}{cc} 180 & 192 \end{array}$$

180 mg of  $C_6H_{12}O_6$  required 192 mg of  $O_2$  for Complete Oxidation

BOD of sample =  $192 \times 10.2 / 180$

$$= 10.88 \text{ mg}$$

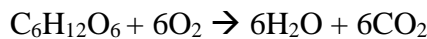
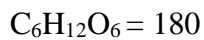
$$= 10.88 \times 1000 / 200$$

$$= 54.4 \text{ ppm}$$

**15. 1 dm<sup>3</sup> of water containing 200 mg of glucose ( $C_6H_{12}O_6$ ) completely oxidized to  $CO_2$  and  $H_2O$ . Calculate the BOD of the waste water. (Given: C = 12, H = 1, O = 16)**

**Ans: BOD = 213.3 ppm**

**Given:** 1 dm<sup>3</sup> water sample = 200 mg of glucose ( $C_6H_{12}O_6$ )



$$\begin{array}{cc} 180 & 192 \end{array}$$

180 mg of  $C_6H_{12}O_6$  required 192 mg of  $O_2$  for Complete Oxidation

BOD of sample =  $200 \times 192 / 180$

$$= 213.3 \text{ ppm}$$