

## 17.7: Quantitative Aspects of Electrolysis

Michael Faraday discovered in 1833 that there is always a simple relationship between the amount of substance produced or consumed at an electrode during electrolysis and the quantity of electrical charge *Q* which passes through the cell. For example, the half-equation

$$\mathrm{Ag}^+ + e^- \to \mathrm{Ag}$$

tells us that when 1 mol  $Ag^+$  is plated out as 1 mol Ag, 1 mol  $e^-$  must be supplied from the cathode. Since the negative charge on a single electron is known to be  $1.6022 \times 10^{-19}$  C, we can multiply by the Avogadro constant to obtain the charge per mole of electrons. This quantity is called the **Faraday constant**, symbol F:

$$F = 1.6022 \times 10^{-19} \text{ C} \times 6.0221 \times 10^{23} \text{ mol}^{-1} = 9.649 \times 10^{4} \text{ C mol}^{-1}$$

Thus in the case of Eq. (1), 96 490 C would have to pass through the cathode in order to deposit 1 mol Ag. For any electrolysis the electrical charge Q passing through an electrode is related to the amount of electrons  $n_e$  by

$$\mathrm{F} = rac{Q}{n_{e^-}}$$

Thus F serves as a conversion factor between  $n_e$  and Q.

## ✓ Example 17.7.1: Electrical Charge

Calculate the quantity of electrical charge needed to plate 1.386mol Cr from an acidic solution of  $K_2Cr_2O_7$  according to half-equation

$$H_2Cr_2O_7(aq) + 12H^+(aq) + 12e^- \rightarrow 2Cr(s) + 7H_2O(l)$$
 (17.7.1)

**Solution** 

According to Eq. 17.7.1, 12 mol  $e^-$  is required to plate 2 mol Cr, giving us a stoichiometric ratio  $S(e^-/\text{Cr})$ . Then the Faraday constant can be used to find the quantity of charge. In road-map form

$$n_{\text{Cr}} \xrightarrow{S(e^-/\text{Cr})} n_e \xrightarrow{F} Q$$

$$Q = 1.386 \text{ mol Cr} \times \frac{12 \text{ mol } e^-}{2 \text{ mol Cr}} \times \frac{9.649 \times 10^4 \text{ C}}{1 \text{ mol } e^-} = 8.024 \times 10^5 \text{ C}$$

Often the electrical current rather than the quantity of electrical charge is measured in an electrolysis experiment. Since a **coulomb** is defined as the quantity of charge which passes a fixed point in an electrical circuit when a current of one ampere flows for one second, the charge in coulombs can be calculated by multiplying the measured current (in amperes) by the time (in seconds) during which it flows:

$$Q = It$$

In this equation *I* represents current and *t* represents time. If you remember that

coulomb = 1 ampere  $\times$  1 second 1 C = 1 A s

you can adjust the time units to obtain the correct result.

## ✓ Example 17.7.2 : Mass of Hydrogen Peroxide

Hydrogen peroxide, H<sub>2</sub>O<sub>2</sub>, can be manufactured by electrolysis of cold concentrated sulfuric acid. The reaction at the anode is

$$2 H_2 SO_4 \longrightarrow H_2 S_2 O_8 + 2 H^+ + 2e^-$$
 (17.7.2)

When the resultant peroxydisulfuric acid, H<sub>2</sub>S<sub>2</sub>O<sub>8</sub>, is boiled at reduced pressure, it decomposes:

$$2 H_2 O + H_2 S_2 O_8 \longrightarrow 2 H_2 SO_4 + H_2 O_2$$
 (17.7.3)



Calculate the mass of hydrogen peroxide produced if a current of 0.893 flows for 1 h.

## **Solution**

he product of current and time gives us the quantity of electricity, Q. Knowing this we easily calculate the amount of electrons,  $n_e$ —. From half-equation 17.7.2 we can then find the amount of peroxydisulfuric acid. Equation 17.7.3 then leads to  $n_{\rm H2O2}$  and finally to  $m_{\rm H2O2}$ . The road map to describe this logic is as follows:

$$I \overset{t}{ o} Q \overset{F}{\overset{ o}{ o}} n_{e^-} \overset{S_e}{\overset{ o}{ o}} n_{\mathrm{H_2S_2O_8}} \overset{S}{ o} n_{\mathrm{H_2O_2}} \overset{M}{\overset{ o}{ o}} m_{\mathrm{H_2O_2}}$$

so that

$$\begin{split} m_{\rm H_2O_2} &= 0.893~\rm A~\times~3600~s \times \frac{1~mol~e^-}{96~490~\rm C} \times \frac{1~mol~\rm H_2S_2O_8}{2~mol~e^-} \\ &= \frac{1~mol~\rm H_2O_2}{1~mol~\rm H_2S_2O_8}~\times~\frac{34.01~\rm g~\rm H_2O_2}{1~mol~\rm H_2O_2} \end{split}$$

$$= 05666 \frac{\text{As}}{\text{C}} \times \text{g H}_2\text{O}_2 = 0.5666 \text{ g H}_2\text{O}_2$$

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