• An aqueous solution of CuSO₄ is electrolysed with a current of 1.00 A for 60 minutes. Calculate the mass of the products that are formed at the two electrodes.

If a current of 1.00 A is passed for 60 minutes, the number of electrons delivered is,

moles of
$$e^{-} = \frac{It}{F} = \frac{(1.00 \text{ A}) \times (60 \times 60 \text{ s})}{(96485 \text{ C mol}^{-1})} = 0.037 \text{ mol}$$

At the cathode, the two possible reduction reactions are:

$$Cu2+(aq) + 2e- → Cu(s) 2H2O(l) + 2e- → H2(g) + 2OH-(aq) Eo = +0.34 VEo = -0.83 V$$

The reduction of $Cu^{2+}(aq)$ is much easier (*E* is less negative) so Cu(s) is formed on the cathode.

[This potential for the reduction of water corresponds to $[H^+(aq)] = 1$ M. Using the Nernst equation gives E = -0.42 V for a pH 7 solution. An overpotential of 0.4 to 0.6 V must be added so that the reduction potential is actually ~1 V. Thus it is even more difficult to reduce water than the E° value suggests.]

Each mole of Cu(s) requires 2 moles of electrons so,

number of moles of $Cu(s) = \frac{1}{2} \times 0.037 \text{ mol} = 0.019 \text{ mol}$

The atomic mass of Cu is 63.55 g mol⁻¹ so the mass of copper deposited is,

mass of Cu(s) = number of moles \times molar mass = 0.019 mol \times 63.55 g mol⁻¹ = 1.2 g

At the anode, oxidation of H_2O occurs to produce $O_2(g)$.

 $2H_2O(l) \rightarrow O_2(g) + 4H^+(aq) + 4e^- E^\circ = -1.23 V$

As SO_4^{2-} contains S(VI), further oxidation is very difficult.

Production of each mole of $O_2(g)$ requires 4 moles of electrons so,

number of moles of $O_2(g) = \frac{1}{4} \times 0.037 \text{ mol} = 0.0093 \text{ mol}$

The molar mass of O_2 is $(2 \times 16.00) = 32.00$ g mol⁻¹ so the mass of $O_2(g)$ formed is,

mass of $O_2(g)$ = number of moles × molar mass = 0.0093 mol × 32.00 g mol⁻¹ = 0.30 g