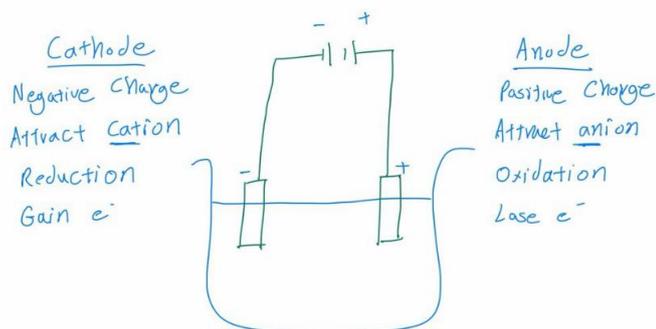


Electrolytic Cells



How to determine reactions at each electrode

Step 1: Look at the ions and molecules in the electrolyte

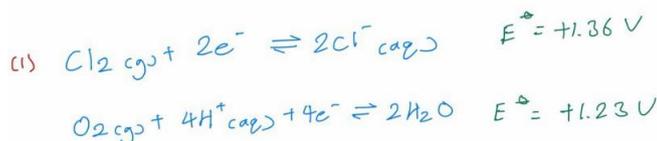
- This will require looking closely at the state of the electrolyte: molten or aqueous

Step 2: Look at the potential ions/molecules at each electrode, analyse using E^\ominus

- Species with higher E^\ominus are more readily reduced and less easily oxidised
- Species with lower E^\ominus are more readily oxidised and less easily reduced
- It's important to always think about H_2O whenever the electrolyte is in aqueous state

Step 3: Look at concentrations (usually for anions/anode)

- Usually for F^- , Cl^- , Br^- , when they are in higher concentrations, they can oxidise more favourably even when the E^\ominus says otherwise.
- For large anions like NO_3^- , a higher concentration usually will not be enough for it to be oxidised



- E.g. in NaCl (conc) the high $[Cl^-]$ will shift equilibrium \leftarrow , E^\ominus decreases, and is lesser than $E^\ominus(O_2(g)|H_2O(l))$, Cl^- becomes more easily oxidised than H_2O

Step 4: Look at nature of electrode

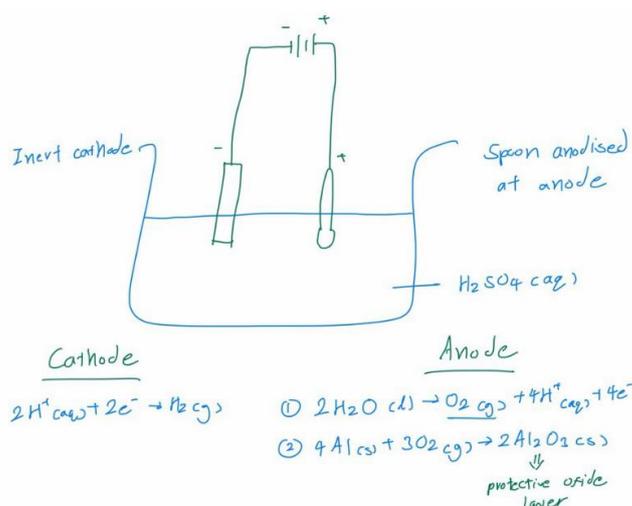
- If it is an inert electrode like Graphite or Platinum, the electrode will not be involved
- If it is an active electrode, they can participate in the reaction (usually favoured to be oxidised)
- Look at E^\ominus for oxidation of the active electrode and compare to that of ions and molecules

Calculations in Electrolysis

$$Q = It \quad \begin{matrix} \text{Quantity of Charge (C)} \\ \text{time (s)} \\ \text{Current (A)} \end{matrix}$$

$$Q = nF \quad \begin{matrix} \text{Faraday's} = 96500 \text{ C mol}^{-1} \\ \text{no. of moles of } e^- \end{matrix}$$

Anodising Aluminium



Purification of Copper

