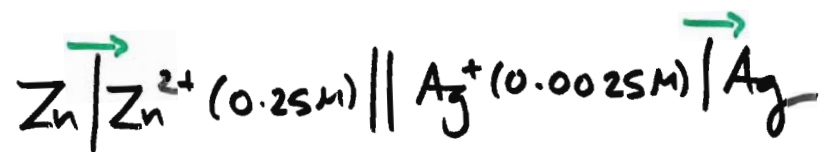
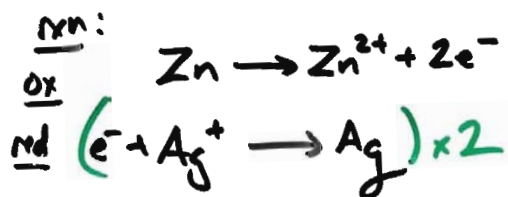


What's E_{cell}° and E_{cell} for:



$$\begin{aligned} E_{\text{cell}}^{\circ} &= E_{\text{RHS}}^{\circ} - E_{\text{LHS}}^{\circ} \\ &= +0.80\text{V} - -0.76\text{V} \\ &= +1.56\text{V} \end{aligned}$$



$$E = E^{\circ} - \frac{RT}{nF} \ln Q \quad (25^{\circ}\text{C})$$

$$= +1.56\text{V} - \frac{8.3145 \text{ J/mol}\cdot\text{K} \times 298\text{K}}{2 \times 96,500 \text{ C/mol}} \ln Q$$

$$Q = \frac{[\text{Zn}^{2+}]}{[\text{Ag}^{+}]^2} = \frac{0.25}{0.0025^2}$$

$$\begin{aligned} E &= 1.56\text{V} - \frac{8.3145 \text{ J/mol}\cdot\text{K} \times 298\text{K}}{2 \times 96,500 \text{ C/mol}} \cdot \ln\left(\frac{0.25}{0.0025^2}\right) \\ &= +1.56\text{V} - 0.0128 \frac{\text{J}}{\text{C}} \times 10.597 \end{aligned}$$

elec work = charge \times voltage.

$$1\text{J} = 1\text{C} \times 1\text{V} \quad \Rightarrow \quad 1 \frac{\text{J}}{\text{C}} = 1\text{V}$$

$$\Rightarrow \boxed{E = 1.42\text{V}}$$

Electrolysis

Voltaic/Galvanic cells ~ produce elec energy from a spontaneous redox rxn.

$$E_{\text{cell}}^{\circ} = +ve$$

$$\Delta G^{\circ} = -ve$$

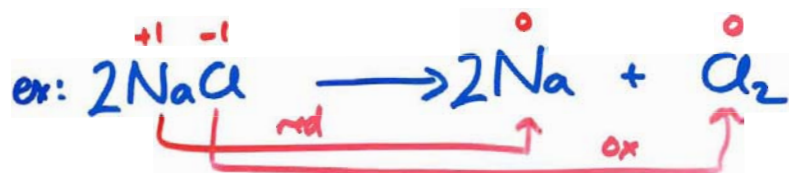
$$K \gg 1$$

Electrolytic cells ~ consume elec energy to drive an otherwise non-spont. rxn.

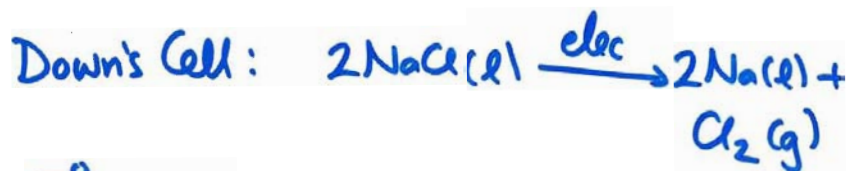
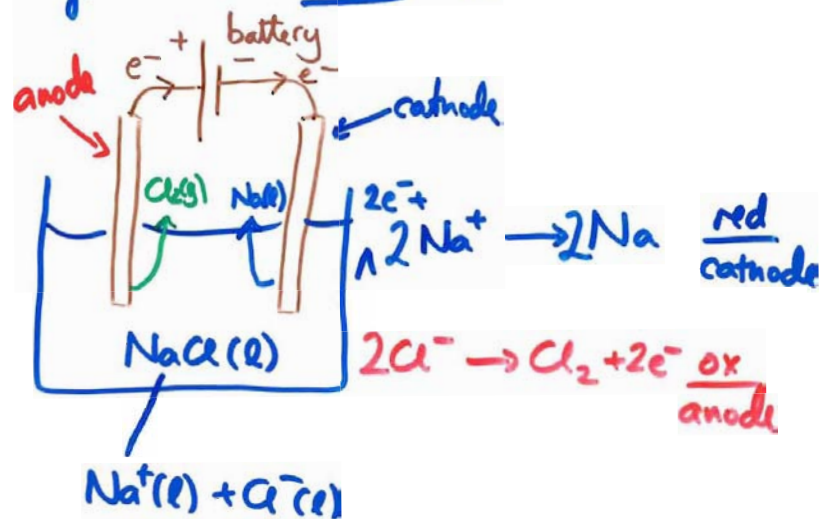
$$E_{\text{cell}}^{\circ} = -ve$$

$$\Delta G^{\circ} = +ve$$

$$K \ll 1$$



electrolysis can DRIVE this rxn!



$$E_{\text{cell}}^{\circ} \approx -4\text{V}$$

have to supply $>4\text{V}$ to drive rxn.

Quantitative aspects of electrolysis

Michael Faraday ... he noticed that the mass of products formed @ electrodes

\propto time
 \propto electrical current
 \propto molar mass of element.

$$\text{Electrical Current} = \frac{\text{Charge } Q \text{ (C)}}{\text{time } t \text{ (s)}}$$
$$I \text{ (A)}$$

$$I = \frac{Q}{t}$$

(A) (C) (s)

$$1 \text{ A} = 1 \frac{\text{C}}{\text{s}}$$

$$1 \text{ A} = 1 \frac{\text{C}}{\text{s}}, \quad 15 \text{ A} = 15 \frac{\text{C}}{\text{s}}$$

$$\text{total charge: } \boxed{Q = I \cdot t}$$

ex: What's charge when a 13A current flows for 3.0 min?

$$Q = 13 \frac{\text{C}}{\text{s}} \times 3.0 \text{ min} \times \frac{60 \text{ s}}{1 \text{ min}}$$

$$= 2300 \text{ C}$$

How many mol e⁻?

$$2300 \text{ C} \times \frac{1 \text{ mole } e^-}{96,500 \text{ C}} = 0.024 \text{ mol } e^-$$

let's say we're electrolyzing NaCl.



in general: compound



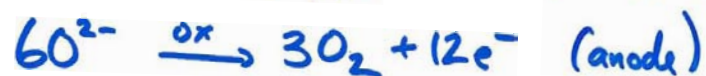
↳ elec elements

ex: What mass of Na + what mass Cl_2 would we make?

$$\frac{0.024 \text{ mole } e^-}{2 \text{ mole } e^-} \left| \frac{2 \text{ mol Na}}{1 \text{ mol Na}} \right| \frac{22.99 \text{ g Na}}{1 \text{ mol Na}} = 0.55 \text{ g Na}$$

$$\frac{0.024 \text{ mole } e^-}{2 \text{ mole } e^-} \left| \frac{1 \text{ mol } \text{Cl}_2}{1 \text{ mol } \text{Cl}_2} \right| \frac{70.90 \text{ g } \text{Cl}_2}{1 \text{ mol } \text{Cl}_2} = 0.85 \text{ g } \text{Cl}_2$$

let's try another one!



let's electrolyze Al_2O_3 for 10d @ current 12A.

Q. How many grams of Al?

Mole e^- ?

$$12 \frac{\text{C}}{\text{s}} \times 10 \text{ d} \times \frac{24 \text{ h}}{1 \text{ d}} \times \frac{60 \text{ min}}{1 \text{ h}} \times \frac{60 \text{ s}}{1 \text{ min}} =$$

$$\frac{1.0 \times 10^7 \text{ C}}{96,500 \text{ C}} = 110 \text{ mol } e^-$$

$$\frac{110 \text{ mole } e^-}{12 \text{ mol } e^-} \left| \frac{4 \text{ mol Al}}{1 \text{ mol Al}} \right| \frac{26.98 \text{ g Al}}{1 \text{ mol Al}} = 970 \text{ g Al}$$

$\approx 2 \text{ lbs.}$