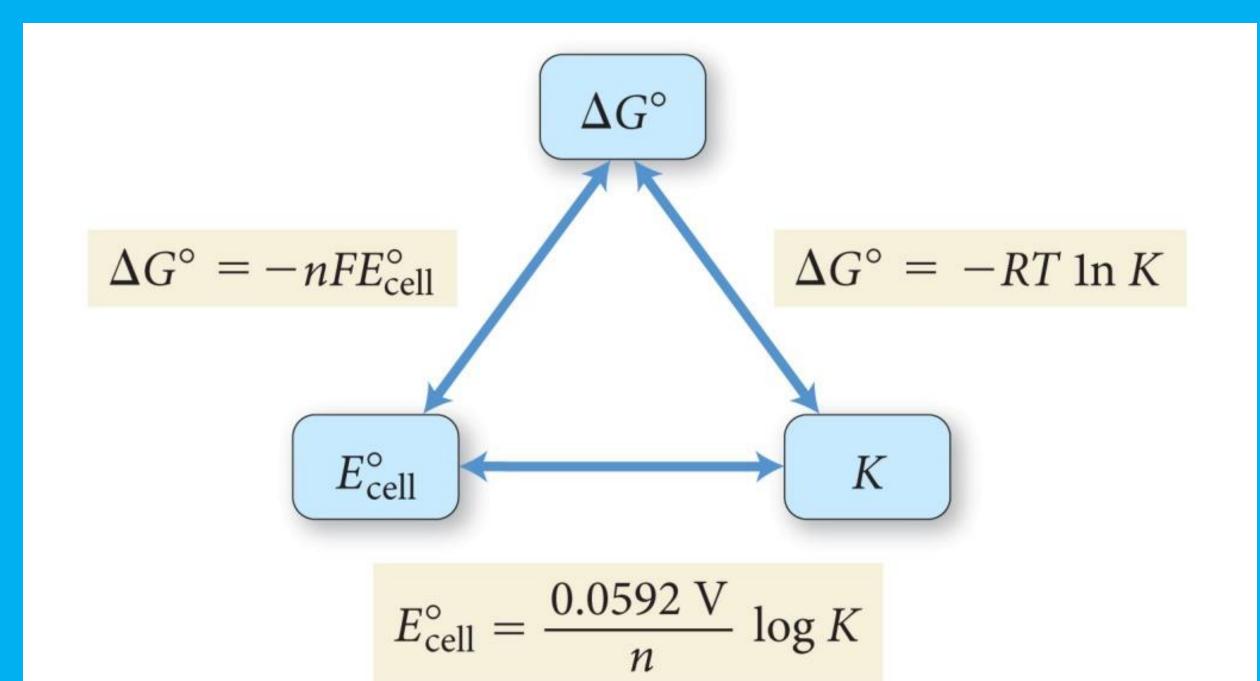
N45 - Electrochemistry

Variables Unite!

N45 - Electrochemistry

Variables Unite!

Target: I can connect Thermodynamics, Equilibrium, and Electrochemistry. The big finale of the year!



$\underline{E^{\circ}_{cell}}$, ΔG° , and K

For a spontaneous reaction

One that proceeds in the forward direction with the

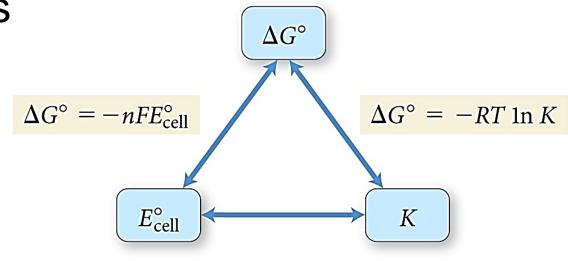
chemicals in their standard states

$$\Delta G^{\circ}$$
 < 0 (negative)

$$E^{\circ} > 0$$
 (positive)

$\Delta G^{\circ} = -RTIn(K) = -nFE^{\circ}_{cell}$

n = the number of electrons



$$E_{\text{cell}}^{\circ} = \frac{0.0592 \text{ V}}{n} \log K$$

Calculating ∆G⁰ for a Cell

$$\Delta G^0 = -nFE^\circ$$

n = moles of electrons in balanced redox equation

F = Faraday constant = 96,485 coulombs/mol e⁻

$$Zn + Cu^{2+} \rightarrow Zn^{2+} + Cu$$
 $E^{\circ} = + 1.10 \text{ V}$

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$$DG^{0} = -(2 mol e^{-})(96485 \frac{coulombs}{mol e^{-}})(1.10 \frac{Joules}{Coulomb})$$

$$\Delta G^0 = -212267 Joules = -212 kJ$$

The Nernst Equation

Standard potentials assume a concentration of 1 M. The Nernst equation allows us to calculate potential when the two cells are not 1.0 M.

$$E = E^0 - \frac{RT}{nF} \ln(Q)$$

 $R = 8.31 \text{ J/(mol \cdot K)}$

7 = Temperature in K

n = moles of electrons in balanced redox equation

F = Faraday constant = 96,485 coulombs/mol e⁻¹

Deriving the Nernst Equation

$$\Delta G = \Delta G^{\circ} + RT \ln Q$$

$$\Delta G = \Delta G^{\circ} + RT \ln Q$$

$$-nFE_{\text{cell}} = -nFE_{\text{cell}}^{\circ} + RT \ln Q$$

$$E_{\text{cell}} = E_{\text{cell}}^{\circ} - \frac{RT}{nF} \ln Q$$

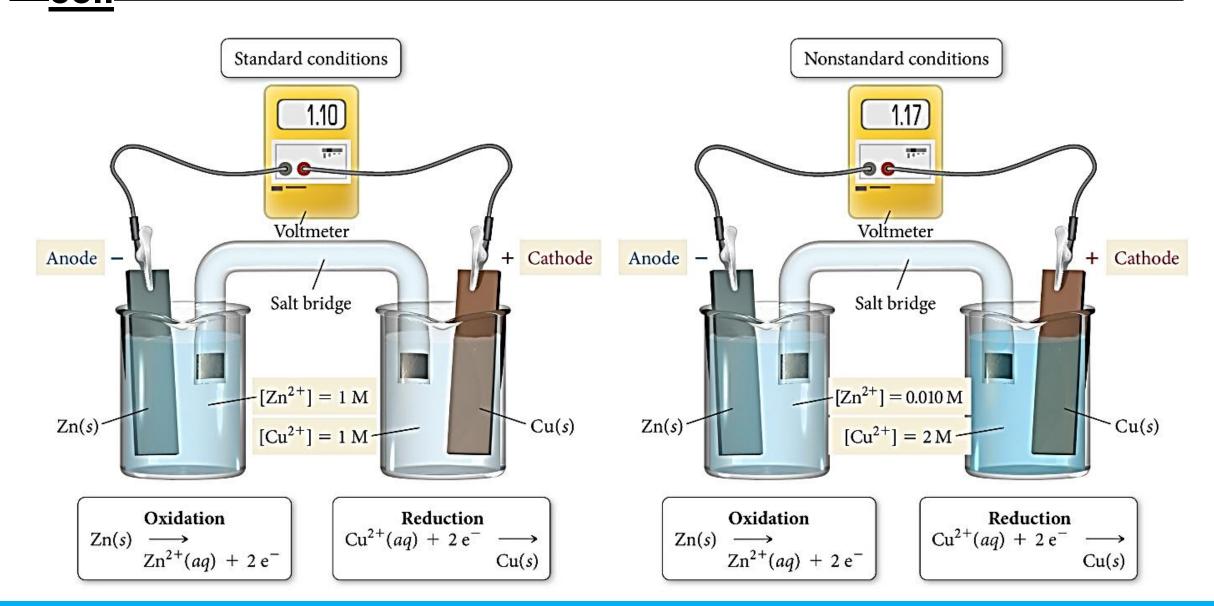
Nernst Equation Simplified

At 25 °C (298 K) the Nernst Equation is simplified this way:

$$E = E^0 - \frac{0.0592}{n} \log(Q)$$

Sometimes you may see 0.0591 on a key or online. It should be 0.0592

E_{cell} When Ion Concentrations Are Not 1 M



Equilibrium Constants and Cell Potential

At equilibrium, forward and reverse reactions occur at equal rates, therefore:

- 1. The battery is "dead" ⊗
- 2. The cell potential, E, is zero volts

Modifying the Nernst Equation (at 25 °C):

$$0 \text{ volts} = E^{\circ} - \frac{0.0592}{n} \log(K)$$

Calculating an Equilibrium Constant from a Cell Potential

$$Zn + Cu^{2+} \rightarrow Zn^{2+} + Cu$$
 $E^{\circ} = + 1.10 \text{ V}$

$$0 \ volts = 1.10 - \frac{0.0592}{2} log(K)$$

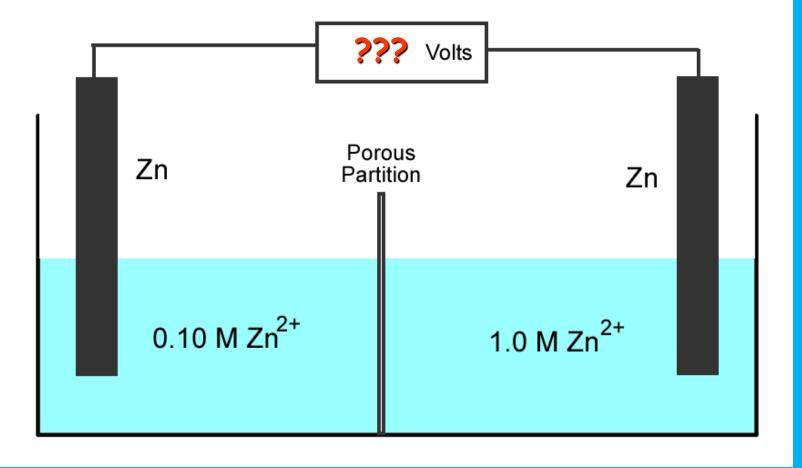
$$\frac{(1.10)(2)}{0.0592} = log(K) \qquad 37.2 = log(K)$$

$$10^{37.2} = K = 1.58 \times 10^{37}$$

Both sides of the cell have the same components but at different concentrations.

Step 1:

Determine which side undergoes oxidation, and which side undergoes reduction.

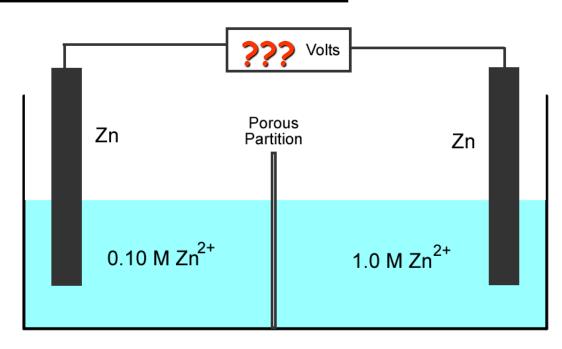


The 1.0 M Zn²⁺ must decrease in concentration, and the 0.10 M Zn²⁺ must increase in concentration

```
Zn^{2+} (1.0M) + 2e^{-} \rightarrow Zn (reduction)

Zn \rightarrow Zn^{2+} (0.10M) + 2e^{-} (oxidation)
```

 $Zn^{2+} (1.0M) \rightarrow Zn^{2+} (0.10M)$

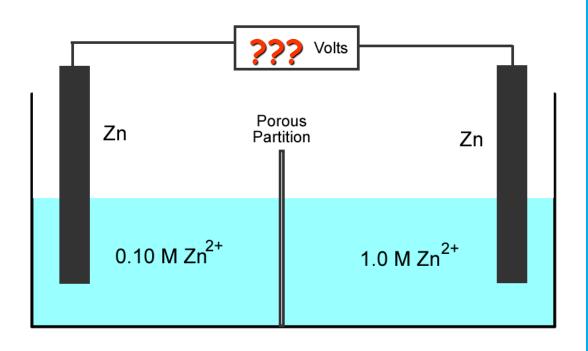


Step 2:

Calculate cell potential using the Nernst Equation (assuming 25 °C).

$$Zn^{2+}$$
 (1.0M) \rightarrow Zn^{2+} (0.10M)

$$E = E^{\circ} - \frac{0.0592}{n} log(Q)$$



$$E^{\circ} = 0$$

$$Q = \frac{[product]}{[reactant]} = \frac{[low]}{[high]}$$

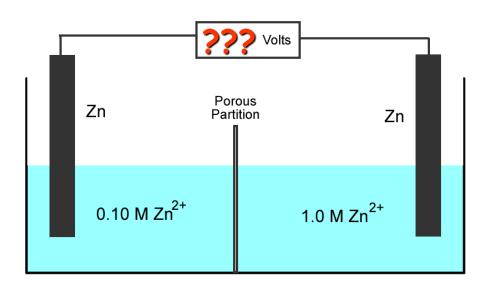
Step 2:

Calculate cell potential using the Nernst Equation (assuming 25 °C).

$$Zn^{2+} (1.0M) \rightarrow Zn^{2+} (0.10M)$$

$$E = E^{\circ} - \frac{0.0592}{n} log(Q)$$

$$E = 0 - \frac{0.0592}{2} log \left(\frac{0.10}{1.0} \right) = 0.030 volts$$



$$= 0.030 volts$$

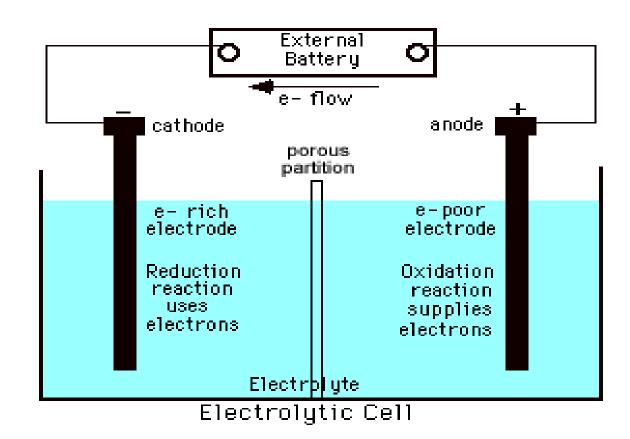
Reminder...Electrolytic Processes

Electrolytic processes are NOT spontaneous.

They have:

A negative cell potential, (-E°)

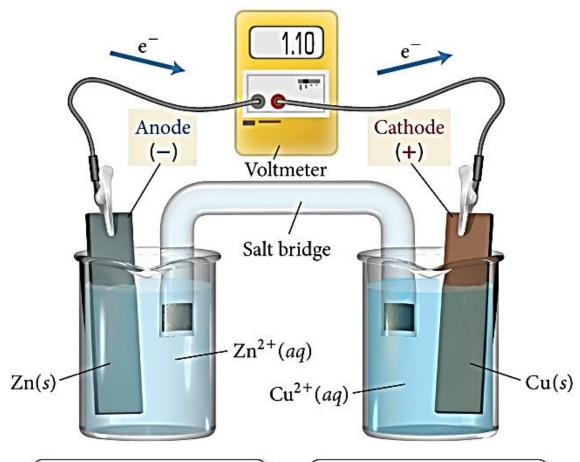
A positive free energy change, $(+\Delta G)$



Voltaic versus Electrolytic Cells

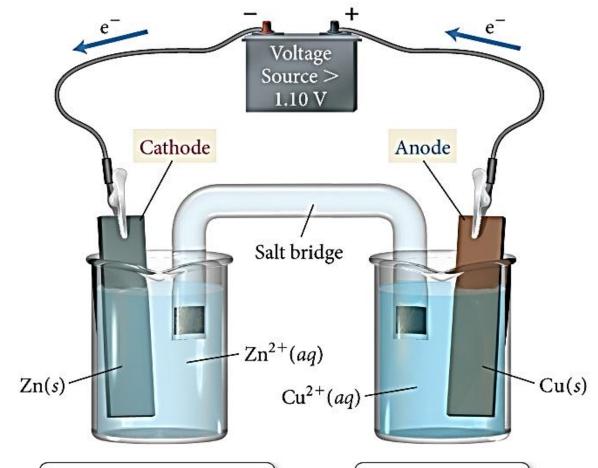
Voltaic Cell

Electrolytic Cell



$$Zn(s) \longrightarrow Zn^{2+}(aq) + 2e^{-}$$

$$Cu^{2+}(aq) + 2e^{-} \xrightarrow{Cu(s)}$$



$$Zn^{2+}(aq) + 2e^{-} \longrightarrow Zn(s)$$

$$Cu(s) \longrightarrow Cu^{2+}(aq) + 2e^{-}$$

Why Would We Want an Electrolytic Cell?

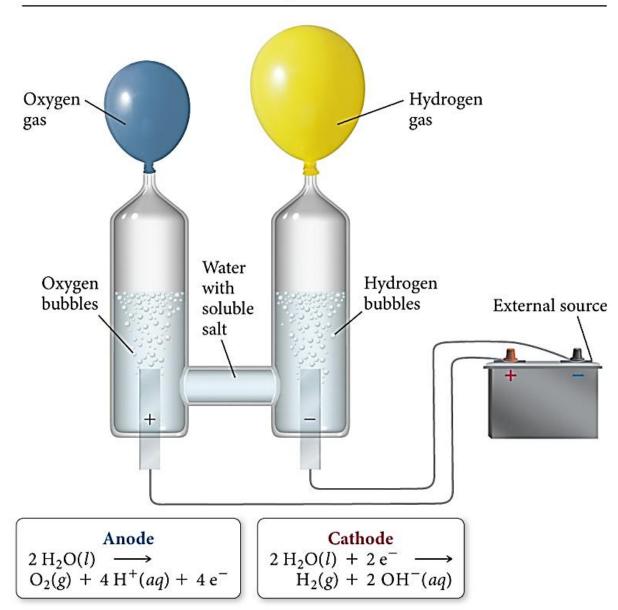
Can be useful!

Can make things happen that would not otherwise happen on their own.

Electrolysis is the process of using electrical energy to break a compound apart.

• Electrolysis is done in an electrolytic cell.

 Electrolytic cells can be used to separate elements from their compounds.



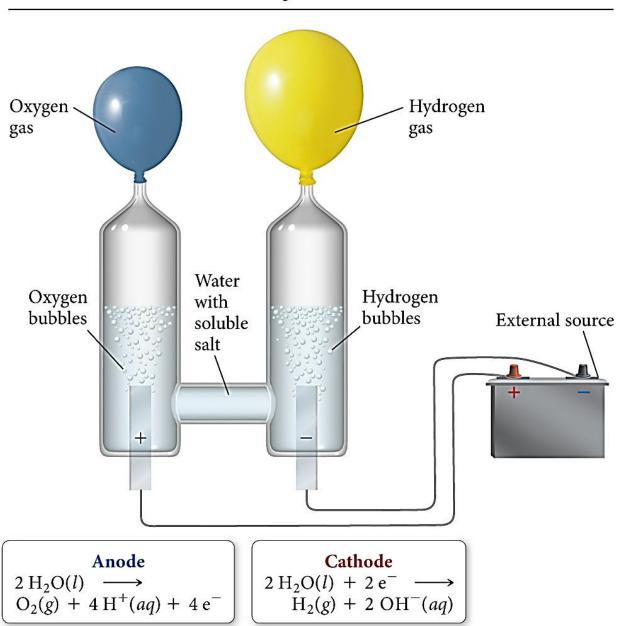
Electrolysis of Water

Anode – oxidation – loss e⁻ $2H_2O_{(I)} \rightarrow O_{2(g)} 4H^+_{(aq)} + 4e^-$

Cathode – reduction – gain e⁻ $2H_2O + 2e^- \rightarrow H_{2(g)} + 2OH_{(aq)}^-$

Memorize which equations! Everyone always forgets it, even though it is on the standard reduction sheet.

Electrolysis of Water



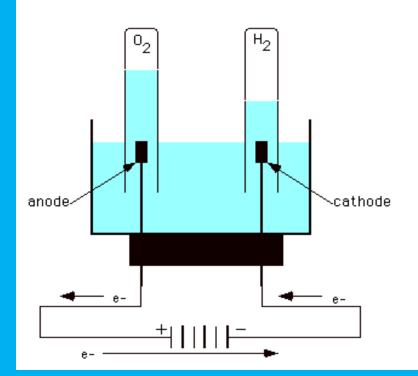
Electrolysis of Water

In acidic solution

Anode rxn:

$$2H_2O \rightarrow O_2 + 4H^+ + 4e^-$$
 -1.23 V

Cathode rxn: $4H_2O + 4e^- \rightarrow 2H_2 + 4OH^-$ -0.83 V



$$2H_2O \rightarrow 2H_2 + O_2$$

-2.06 V

E° = negative Therefore △G = positive Nonspontaneous!

That's why you need to use electricity to make it happen!

Electroplating of Silver

Anode reaction:

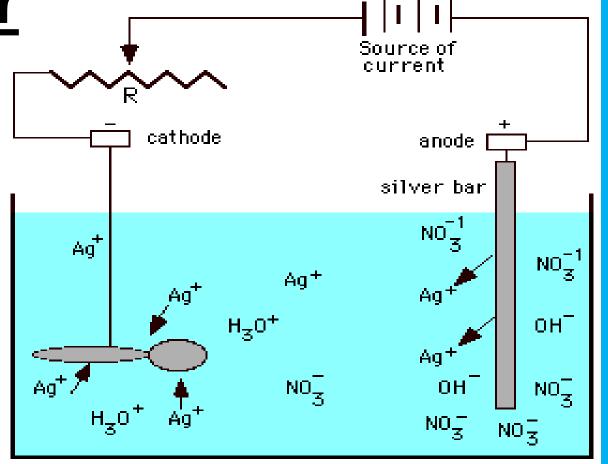
$$Ag \rightarrow Ag^+ + e^-$$

Cathode reaction:

$$Ag^+ + e^- \rightarrow Ag$$

Electroplating requirements:

- 1. Solution of the plating metal
- 2. Anode made of the plating metal
- 3. Cathode with the object to be plated
- 4. Source of current



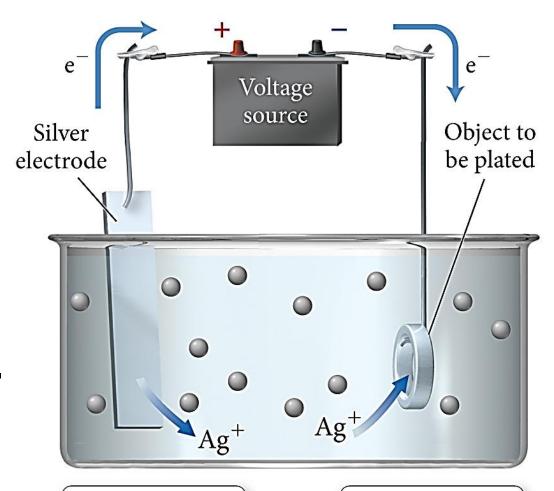
Electroplating

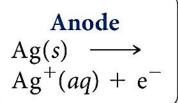
In electroplating, the work piece is the cathode.

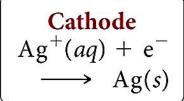
Cations are reduced at cathode, turned into neutral (s) metal, and plate on to the surface of work piece.

The anode is made of the plate metal. The anode oxidizes and replaces the metal cations in the solution. The anode will get smaller over time as it is turned into aqueous cations.

Electrolytic Cell for Silver Plating







Solving an Electroplating Problem

Q: How many seconds will it take to plate out 5.0 grams of silver from a solution of AgNO₃ using a 20.0 Ampere current?

$$Ag^+ + e^- \rightarrow Ag$$

5.0 g
$$\frac{1 \text{ mol Ag}}{107.87 \text{ g}} = \frac{1 \text{ mol e}}{1 \text{ mol Ag}} = \frac{96 485 \text{ g}}{1 \text{ mol e}} = \frac{2.2 \times 10^2 \text{ s}}{20.0 \text{ g}}$$

1 amp = 1 Coulomb/sec

96485 Coulombs = 1 mol e⁻¹

YouTube Link to Presentation

https://youtu.be/9JrlJUUGopY