

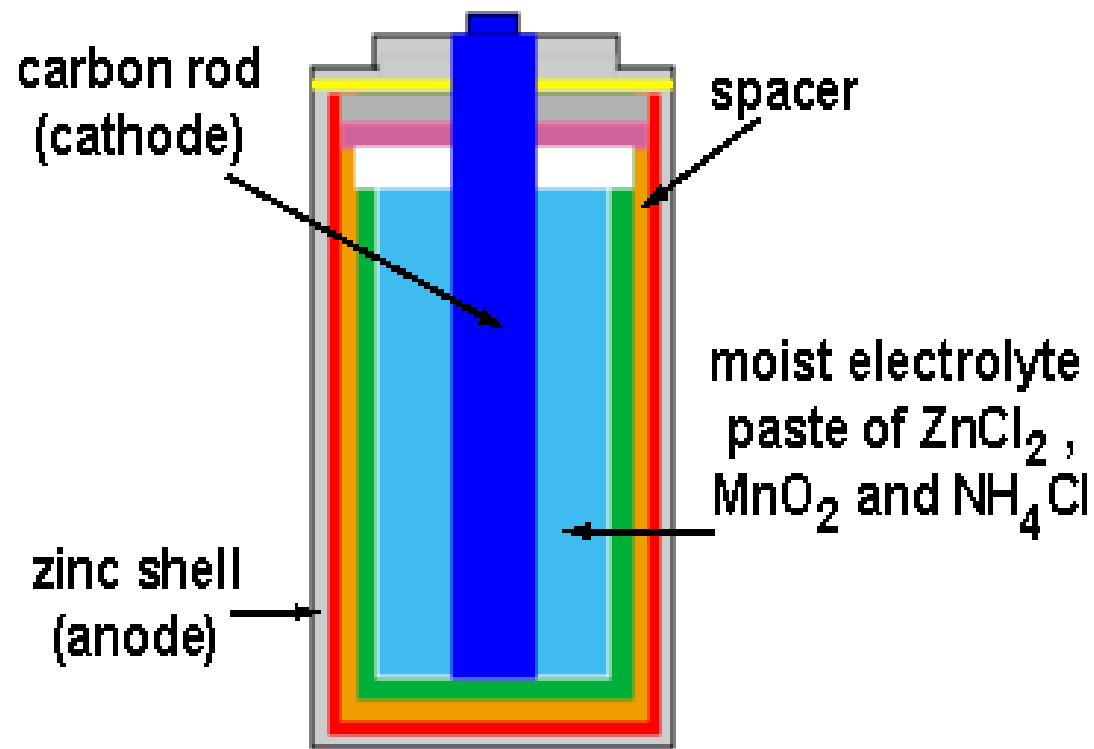
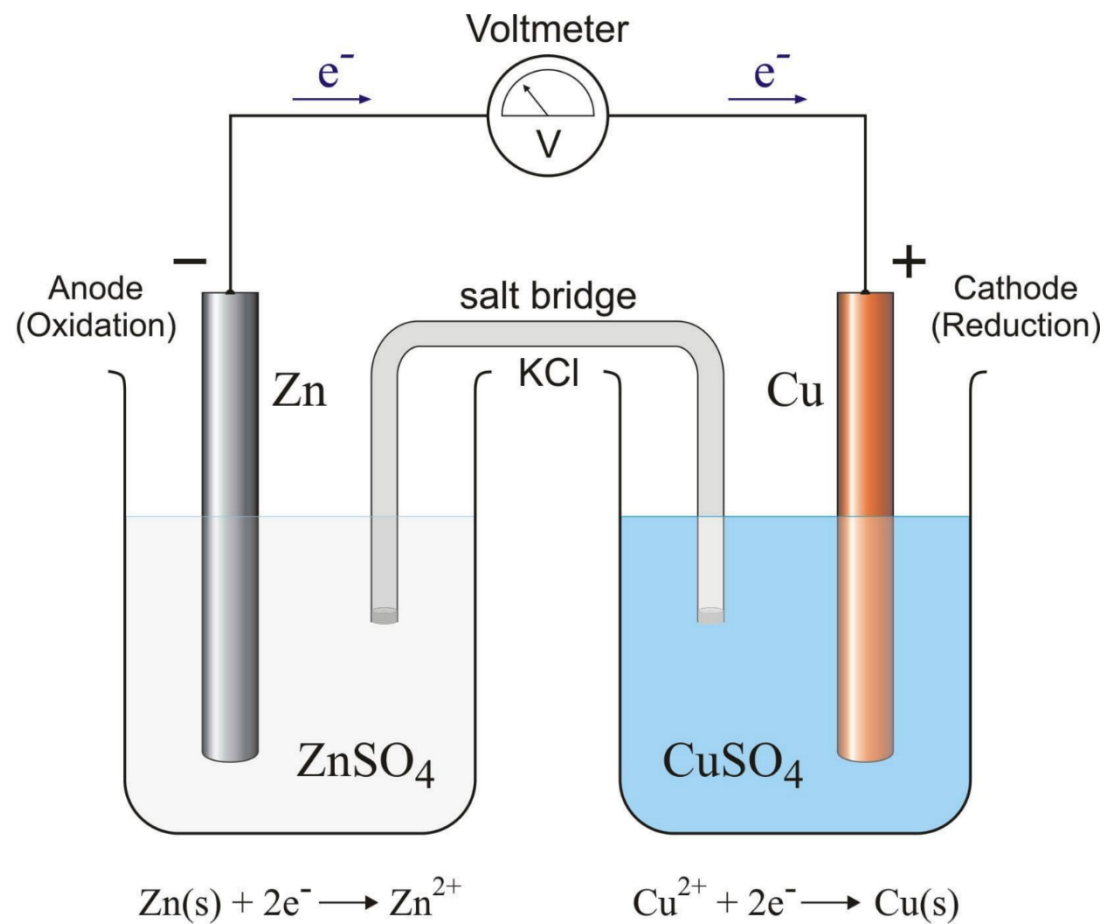
N43 - Electrochemistry

Cell Potential

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Cell Potential

Target: I can calculate the cell potential using standard reduction values.



Mnemonics

LEO goes GER

Loss of Electrons is Oxidation

Gain of Electrons is Reduction

OIL RIG

Oxidation is Loss of Electrons

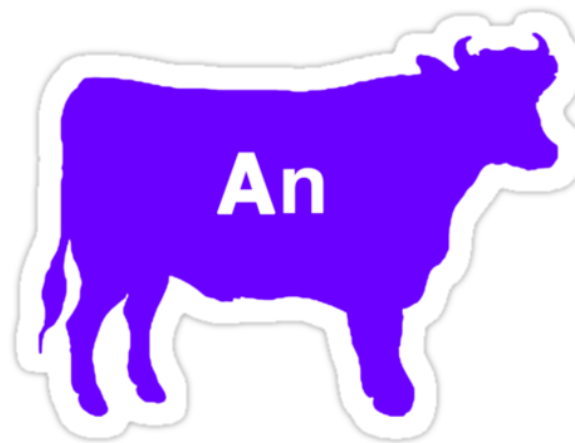
Reduction is Gain of Electrons



A Few More Electrochemistry Terms

Anode

The electrode where oxidation occurs



Anode
is
Oxidation

Cathode

The electrode where reduction occurs



Reduction
at the
Cathode

Cell Potential

Cell Potential - The difference in potential energy between the anode and the cathode in a voltaic cell

Depends on how easy one substance is reduced at the cathode and how easy the other is oxidized at the anode.

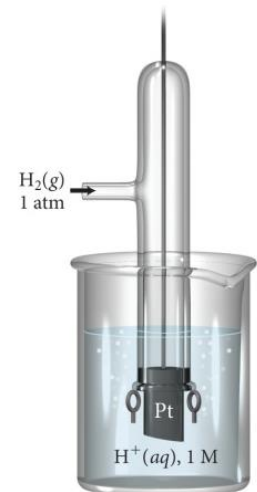
Standard emf, E°_{cell} = Cell potential @ standard conditions (25 °C, 1 atm for gases, 1 M concentration of solution)

– You add the cell potentials for each half reaction

Standard Reduction Potential

- We cannot measure the absolute tendency of a half-reaction, we can only measure it relative to another half-reaction.
- We select as a standard half-reaction the reduction of H^+ to H_2 under standard conditions, which we assign a potential difference = 0 v. (An arbitrary choice!)

Standard hydrogen electrode, SHE



Half-Cell Potentials

- SHE reduction potential is defined to be exactly 0 V.
- Standard reduction potentials compare the tendency for a particular reduction half-reaction to occur relative to the reduction of H^+ to H_2 .
 - Under standard conditions
- Half-reactions with a **stronger tendency toward oxidation** than the SHE have a **negative value for E°_{red}**
- Half-reactions with a **stronger tendency toward reduction** than the SHE have a **positive value for E°_{red}**
- For an oxidation half-reaction, **$E^\circ_{\text{oxidation}} = -E^\circ_{\text{reduction}}$**

Reduction Values

More + means more easily reduced

If you need to flip a rxn,
make sure to flip the sign on E.

If you multiply a rxn,
do **NOT** multiply E.

It is a “state function” and does
not change based on quantity!!!!

Reduction Half-Reaction	E° (V)
$F_2(g) + 2 e^- \longrightarrow 2 F^-(aq)$	2.87
$H_2O_2(aq) + 2 H^+(aq) + 2 e^- \longrightarrow 2 H_2O(l)$	1.78
$PbO_2(s) + 4 H^+(aq) + SO_4^{2-}(aq) + 2 e^- \longrightarrow PbSO_4(s) + 2 H_2O(l)$	1.69
$MnO_4^-(aq) + 4 H^+(aq) + 3 e^- \longrightarrow MnO_2(s) + 2 H_2O(l)$	1.68
$MnO_4^-(aq) + 8 H^+(aq) + 5 e^- \longrightarrow Mn^{2+}(aq) + 4 H_2O(l)$	1.51
$Au^{3+}(aq) + 3 e^- \longrightarrow Au(s)$	1.50
$PbO_2(s) + 4 H^+(aq) + 2 e^- \longrightarrow Pb^{2+}(aq) + 2 H_2O(l)$	1.46
$Cl_2(g) + 2 e^- \longrightarrow 2 Cl^-(aq)$	1.36
$Cr_2O_7^{2-}(aq) + 14 H^+(aq) + 6 e^- \longrightarrow 2 Cr^{3+}(aq) + 7 H_2O(l)$	1.33
$O_2(g) + 4 H^+(aq) + 4 e^- \longrightarrow 2 H_2O(l)$	1.23
$MnO_2(s) + 4 H^+(aq) + 2 e^- \longrightarrow Mn^{2+}(aq) + 2 H_2O(l)$	1.21
$IO_3^-(aq) + 6 H^+(aq) + 5 e^- \longrightarrow \frac{1}{2} I_2(aq) + 3 H_2O(l)$	1.20
$Br_2(l) + 2 e^- \longrightarrow 2 Br^-(aq)$	1.09
$VO_2^+(aq) + 2 H^+(aq) + e^- \longrightarrow VO^{2+}(aq) + H_2O(l)$	1.00
$NO_3^-(aq) + 4 H^+(aq) + 3 e^- \longrightarrow NO(g) + 2 H_2O(l)$	0.96
$ClO_2(g) + e^- \longrightarrow ClO_2^-(aq)$	0.95
$Ag^+(aq) + e^- \longrightarrow Ag(s)$	0.80
$Fe^{3+}(aq) + e^- \longrightarrow Fe^{2+}(aq)$	0.77
$O_2(g) + 2 H^+(aq) + 2 e^- \longrightarrow H_2O_2(aq)$	0.70
$MnO_4^-(aq) + e^- \longrightarrow MnO_4^{2-}(aq)$	0.56
$I_2(s) + 2 e^- \longrightarrow 2 I^-(aq)$	0.54
$Cu^+(aq) + e^- \longrightarrow Cu(s)$	0.52
$O_2(g) + 2 H_2O(l) + 4 e^- \longrightarrow 4 OH^-(aq)$	0.40
$Cu^{2+}(aq) + 2 e^- \longrightarrow Cu(s)$	0.34
$SO_4^{2-}(aq) + 4 H^+(aq) + 2 e^- \longrightarrow H_2SO_3(aq) + H_2O(l)$	0.20
$Cu^{2+}(aq) + e^- \longrightarrow Cu^+(aq)$	0.16
$Sn^{4+}(aq) + 2 e^- \longrightarrow Sn^{2+}(aq)$	0.15
$2 H^+(aq) + 2 e^- \longrightarrow H_2(g)$	0
$Fe^{3+}(aq) + 3 e^- \longrightarrow Fe(s)$	-0.036
$Pb^{2+}(aq) + 2 e^- \longrightarrow Pb(s)$	-0.13
$Sn^{2+}(aq) + 2 e^- \longrightarrow Sn(s)$	-0.14
$Ni^{2+}(aq) + 2 e^- \longrightarrow Ni(s)$	-0.23
$Cd^{2+}(aq) + 2 e^- \longrightarrow Cd(s)$	-0.40
$Fe^{2+}(aq) + 2 e^- \longrightarrow Fe(s)$	-0.45
$Cr^{3+}(aq) + e^- \longrightarrow Cr^{2+}(aq)$	-0.50
$Cr^{3+}(aq) + 3 e^- \longrightarrow Cr(s)$	-0.73
$Zn^{2+}(aq) + 2 e^- \longrightarrow Zn(s)$	-0.76
$2 H_2O(l) + 2 e^- \longrightarrow H_2(g) + 2 OH^-(aq)$	-0.83
$Mn^{2+}(aq) + 2 e^- \longrightarrow Mn(s)$	-1.18
$Al^{3+}(aq) + 3 e^- \longrightarrow Al(s)$	-1.66
$Mg^{2+}(aq) + 2 e^- \longrightarrow Mg(s)$	-2.37
$Na^+(aq) + e^- \longrightarrow Na(s)$	-2.71
$Ca^{2+}(aq) + 2 e^- \longrightarrow Ca(s)$	-2.76
$Ba^{2+}(aq) + 2 e^- \longrightarrow Ba(s)$	-2.90
$K^+(aq) + e^- \longrightarrow K(s)$	-2.92
$Li^+(aq) + e^- \longrightarrow Li(s)$	-3.04

Reduction Table

More + means more easily reduced

More **NIO**

More Negative Is Oxidation

More **PER**

More Positive Is Reduction

Reduction Half-Reaction	E° (V)
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Reduction Table

Example:

Which rxn is more likely to happen at the cathode and which at the anode??



Anode = oxidation = loss e⁻ = more (-) E = less (+)

Cathode = reduction = gain e⁻ = more (+) E

Calculating Cell Potentials under Standard Conditions

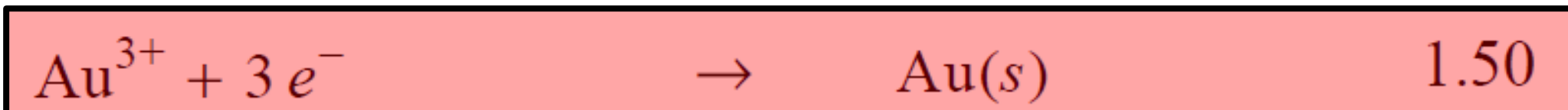
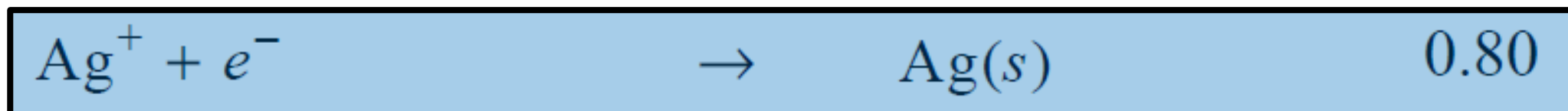
$$E^{\circ}_{\text{cell}} = E^{\circ}_{\text{oxidation}} + E^{\circ}_{\text{reduction}}$$

- When looking up values on reduction table, **flip the sign for the one that is being oxidized** because you have the opposite reaction taking place compared to what is written on the chart.
- When adding E° values for the half-cells, **do not multiply the half-cell E° values**, even if you need to multiply the half-reactions to balance the equation.

Calculating Cell Potential

Example:

What is the cell potential for a cell made with silver and gold?



Anode = oxidation = loss e⁻ = more – E (less +)

Cathode = reduction = gain e⁻ = more + E



Flipped sign for Ag half rxn b/c oxidized but did NOT multiply it by 3.

$$(+1.50) + (-0.80) = 0.70 \text{ V}$$

Sneak Peak at Spontaneity...

If $E^{\circ}\text{cell} = (+)$

then $\Delta G^{\circ} = (-)$

So it is spontaneous!

we will see why in a later lecture 😊

If $E^{\circ}\text{cell} = (-)$

then $\Delta G^{\circ} = (+)$

So it is NON-spontaneous!

YouTube Link to Presentation

<https://youtu.be/iqk4Li9toOQ>