

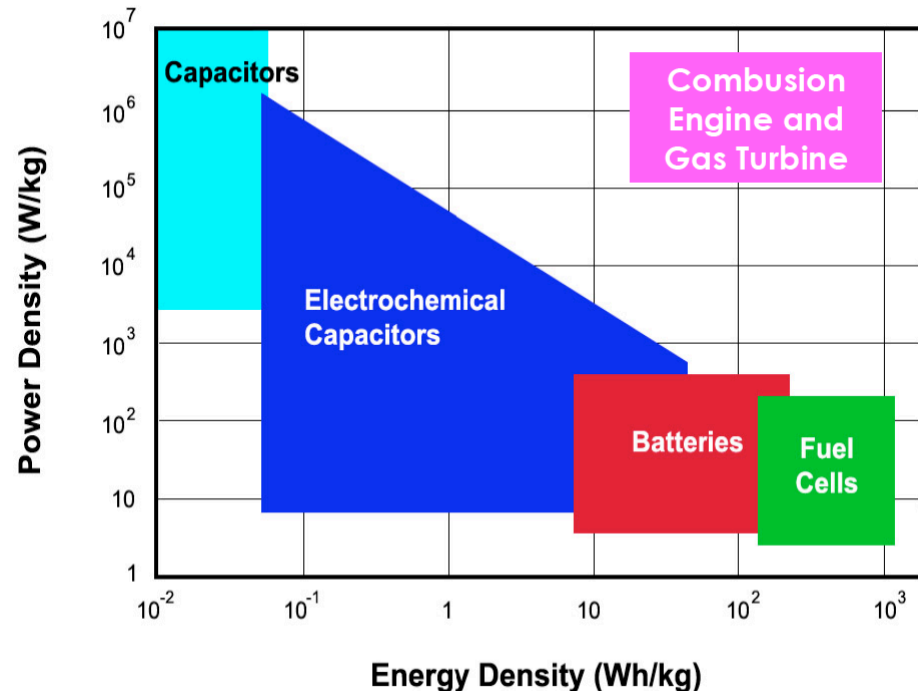
Two Major Types of Electrochemical-Based Energy Storage Devices

- **Batteries**

- ❖ Store energy in chemical reactants capable of generating charge
- ❖ High energy densities
- ❖ Many different varieties

- **Electrochemical Capacitors**

- ❖ Store energy as charge
- ❖ High power densities
- ❖ Sub-second response time



Application for LIBs and Supercapacitors

- *Mobile Electronic Devices*
- *Power-Tools*
- *Electrical Vehicles (HEVs and PEVs)*



Requirements

- **High Power for Intensity of Use**
 - ❖ More positive redox potential (cathode)
 - ❖ Fast charge transfer kinetics (large current output)
- **High Energy (High Capacity) for Length of Use**
 - ❖ More charge per weight/volume
- **Safety**
- **Cost**

Classifications of Cells and Batteries

Primary cells

- Not capable of being recharged electrically
- Good shelf life,
- high energy density at low to moderate discharge rate,
- No or little maintenance
- Ease of use

Secondary or rechargeable cells

- Can be recharged electrically
- High power density
- High discharge rate
- Flat discharge curve
- Good low temperature performance

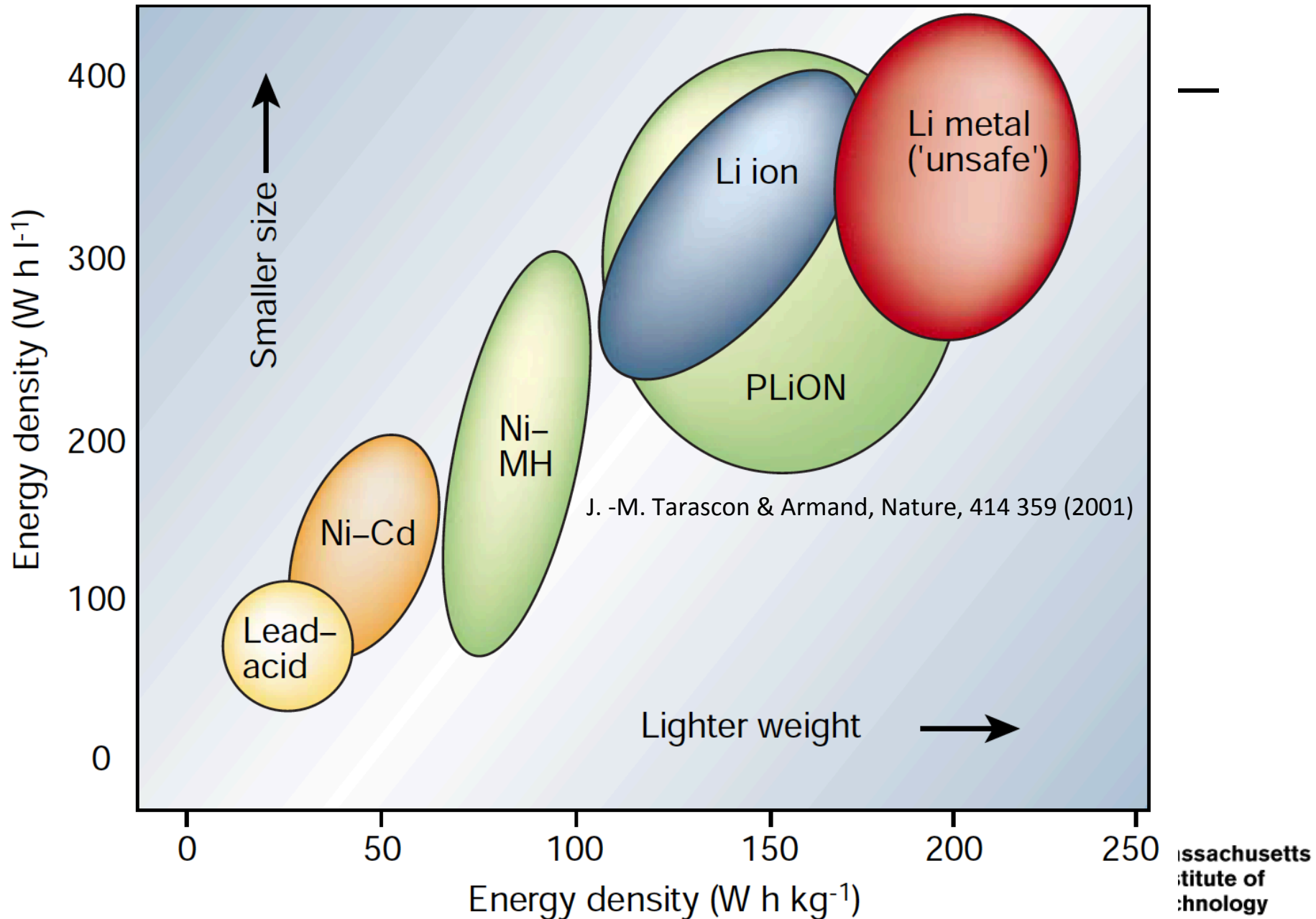
Reserve batteries

- Primary type
- long term storage

Fuel cells

- Active material are fed into the cell from an external source
- Capable of producing electrical energy as long as the active materials are fed to the electrodes

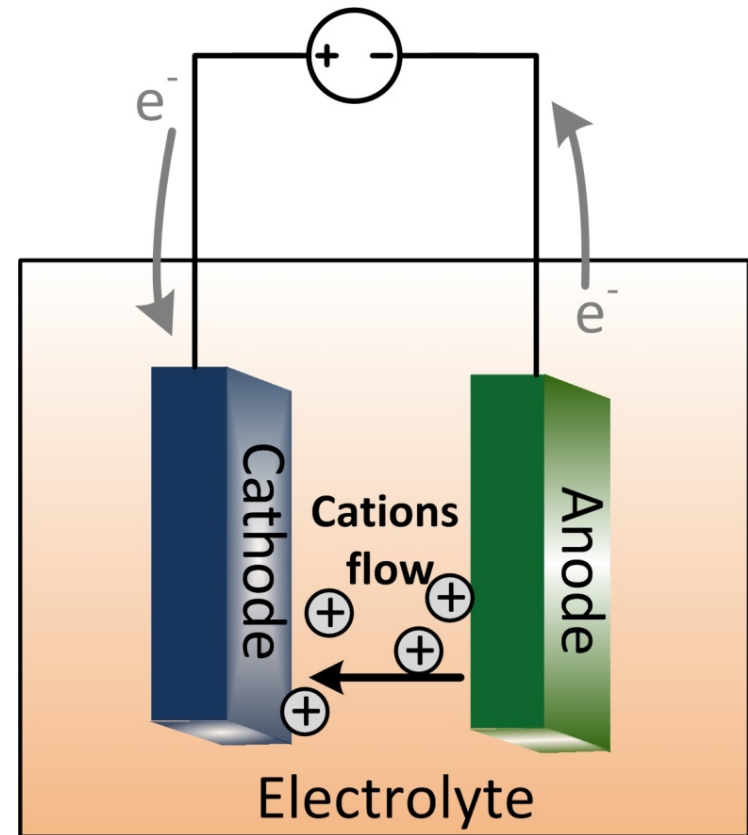
Energy Density for Secondary Batteries



Batteries: Concept and Principle

Battery is a storage device which converts chemical energy into electrical energy

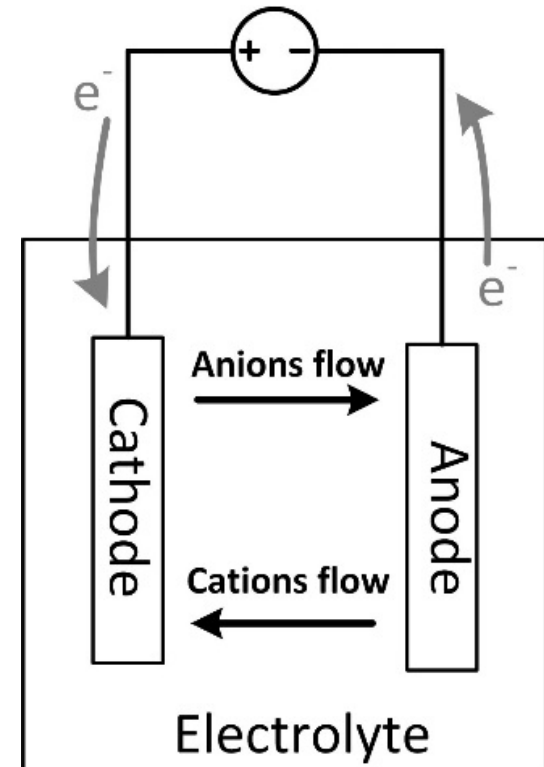
- Main components:
 - Anode or negative electrode
 - Cathode or positive electrode
 - Electrolyte – flow of ions



Operation of a cell

- **Discharge:**





- When cell is connected to an external load, electrons flow from the anode, which is oxidized, through the external load to the cathode, where the electrons are accepted and the cathode material is reduced.
- The electric circuit is completed in the electrolyte by the flow of anions (negative ions) and cations (positive ions) to the anode and cathode, respectively.



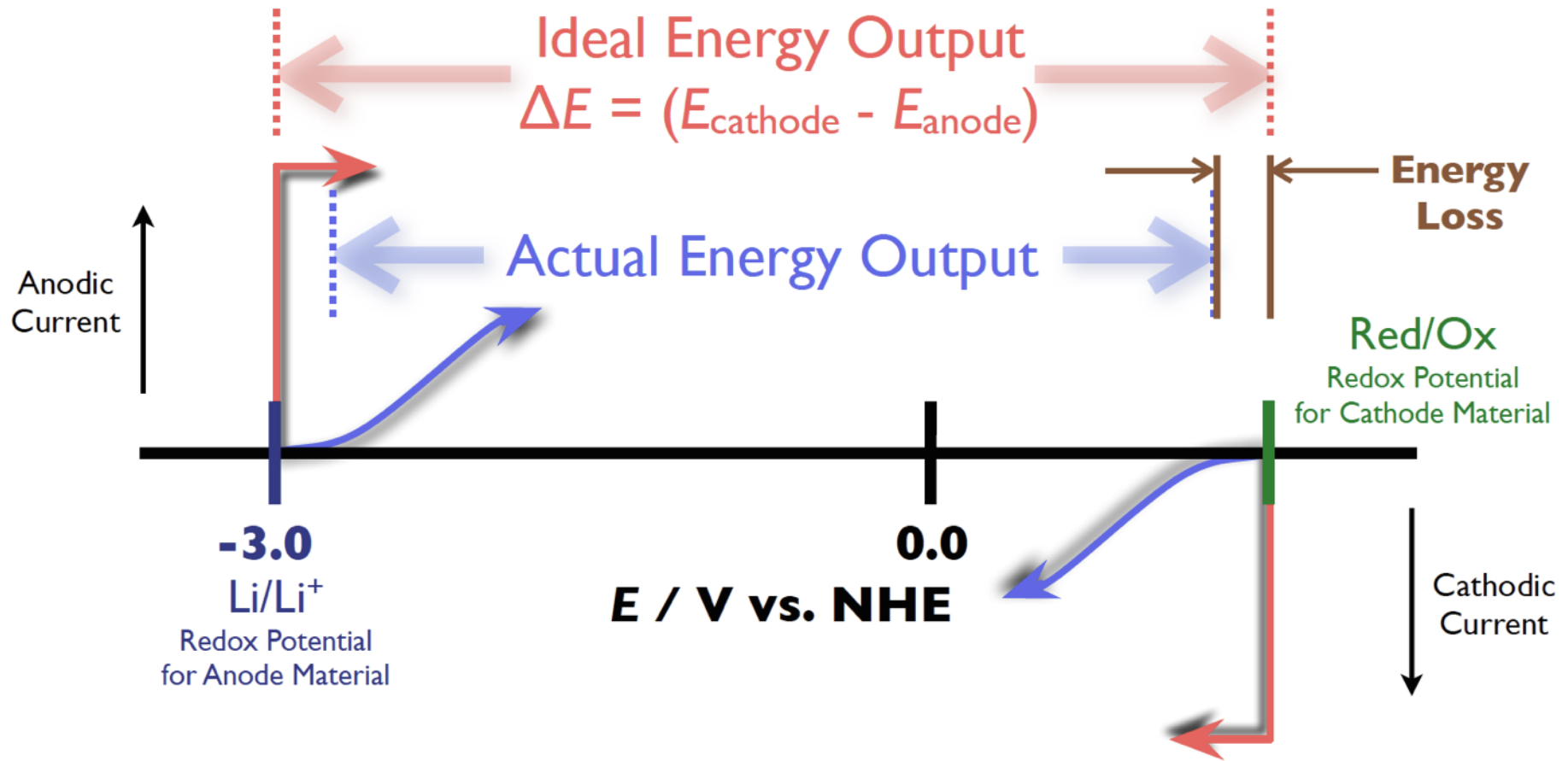
Theoretical Voltage

- The standard potential of the cell is determined by the type of the **active materials** (cathode and anode) in the cell.

TABLE 18.1 Standard Reduction Potentials at 25°C

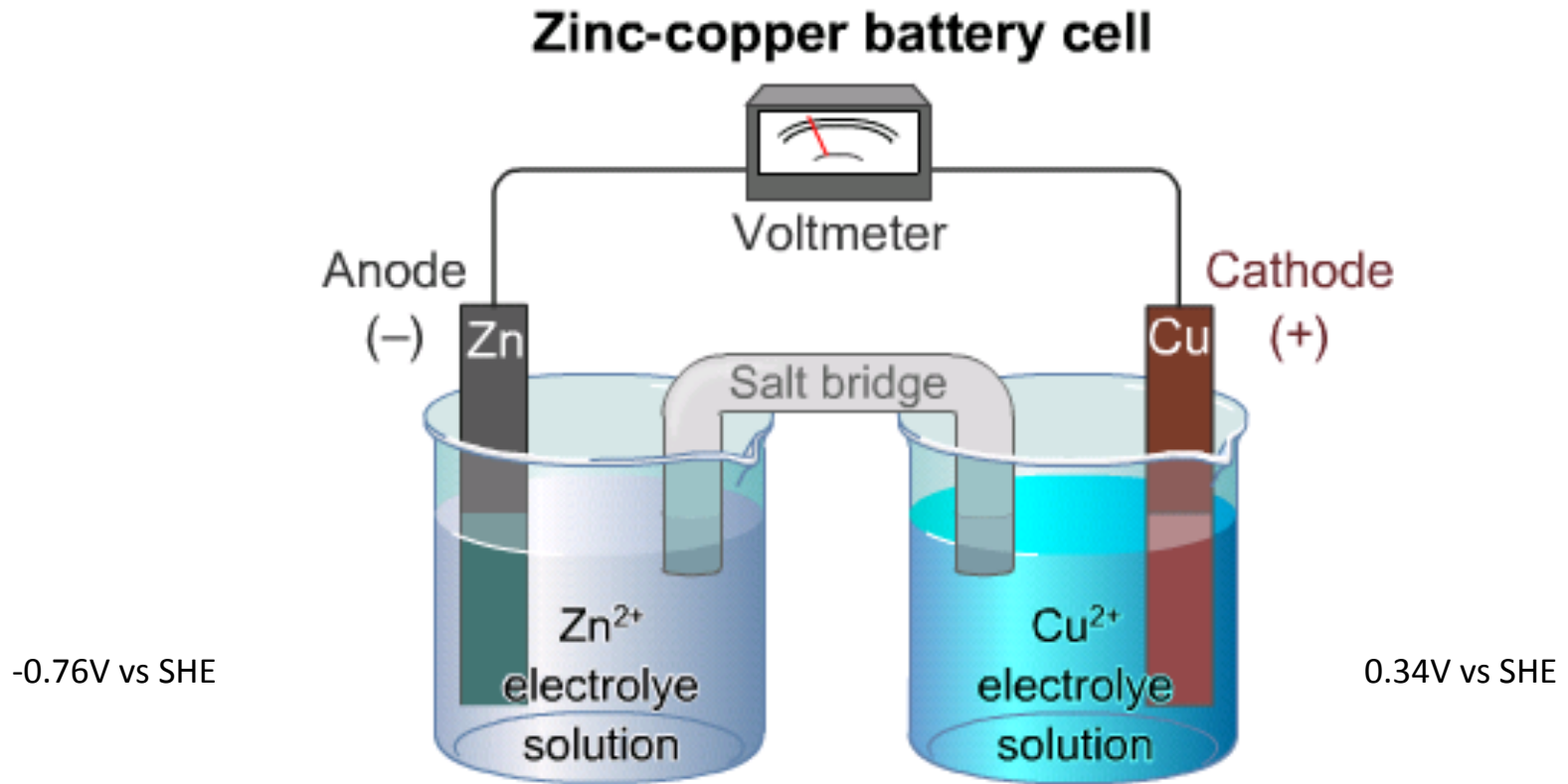
	Reduction Half-Reaction	E° (V)	
Stronger oxidizing agent 	$\text{F}_2(\text{g}) + 2 \text{e}^- \longrightarrow 2 \text{F}(\text{aq})$	2.87	Weaker reducing agent 
	$\text{H}_2\text{O}_2(\text{aq}) + 2 \text{H}^+(\text{aq}) + 2 \text{e}^- \longrightarrow 2 \text{H}_2\text{O}(\text{l})$	1.78	
	$\text{MnO}_4^-(\text{aq}) + 8 \text{H}^+(\text{aq}) + 5 \text{e}^- \longrightarrow \text{Mn}^{2+}(\text{aq}) + 4 \text{H}_2\text{O}(\text{l})$	1.51	
	$\text{Cl}_2(\text{g}) + 2 \text{e}^- \longrightarrow 2 \text{Cl}^-(\text{aq})$	1.36	
	$\text{Cr}_2\text{O}_7^{2-}(\text{aq}) + 14 \text{H}^+(\text{aq}) + 6 \text{e}^- \longrightarrow 2 \text{Cr}^{3+}(\text{aq}) + 7 \text{H}_2\text{O}(\text{l})$	1.33	
	$\text{O}_2(\text{g}) + 4 \text{H}^+(\text{aq}) + 4 \text{e}^- \longrightarrow 2 \text{H}_2\text{O}(\text{l})$	1.23	
	$\text{Br}_2(\text{l}) + 2 \text{e}^- \longrightarrow 2 \text{Br}^-(\text{aq})$	1.09	
	$\text{Ag}^+(\text{aq}) + \text{e}^- \longrightarrow \text{Ag}(\text{s})$	0.80	
	$\text{Fe}^{3+}(\text{aq}) + \text{e}^- \longrightarrow \text{Fe}^{2+}(\text{aq})$	0.77	
	$\text{O}_2(\text{g}) + 2 \text{H}^+(\text{aq}) + 2 \text{e}^- \longrightarrow \text{H}_2\text{O}_2(\text{aq})$	0.70	
	$\text{I}_2(\text{s}) + 2 \text{e}^- \longrightarrow 2 \text{I}^-(\text{aq})$	0.54	
	$\text{O}_2(\text{g}) + 2 \text{H}_2\text{O}(\text{l}) + 4 \text{e}^- \longrightarrow 4 \text{OH}^-(\text{aq})$	0.40	
	$\text{Cu}^{2+}(\text{aq}) + 2 \text{e}^- \longrightarrow \text{Cu}(\text{s})$	0.34	
	$\text{Sn}^{4+}(\text{aq}) + 2 \text{e}^- \longrightarrow \text{Sn}^{2+}(\text{aq})$	0.15	
	$2 \text{H}^+(\text{aq}) + 2 \text{e}^- \longrightarrow \text{H}_2(\text{g})$	0	
Weaker oxidizing agent 	$\text{Pb}^{2+}(\text{aq}) + 2 \text{e}^- \longrightarrow \text{Pb}(\text{s})$	-0.13	Stronger reducing agent 
	$\text{Ni}^{2+}(\text{aq}) + 2 \text{e}^- \longrightarrow \text{Ni}(\text{s})$	-0.26	
	$\text{Cd}^{2+}(\text{aq}) + 2 \text{e}^- \longrightarrow \text{Cd}(\text{s})$	-0.40	
	$\text{Fe}^{2+}(\text{aq}) + 2 \text{e}^- \longrightarrow \text{Fe}(\text{s})$	-0.45	
	$\text{Zn}^{2+}(\text{aq}) + 2 \text{e}^- \longrightarrow \text{Zn}(\text{s})$	-0.76	
	$2 \text{H}_2\text{O}(\text{l}) + 2 \text{e}^- \longrightarrow \text{H}_2(\text{g}) + 2 \text{OH}^-(\text{aq})$	-0.83	
	$\text{Al}^{3+}(\text{aq}) + 3 \text{e}^- \longrightarrow \text{Al}(\text{s})$	-1.66	
	$\text{Mg}^{2+}(\text{aq}) + 2 \text{e}^- \longrightarrow \text{Mg}(\text{s})$	-2.37	
	$\text{Na}^+(\text{aq}) + \text{e}^- \longrightarrow \text{Na}(\text{s})$	-2.71	
	$\text{Li}^+(\text{aq}) + \text{e}^- \longrightarrow \text{Li}(\text{s})$	-3.04	

Energy Density for Secondary Batteries



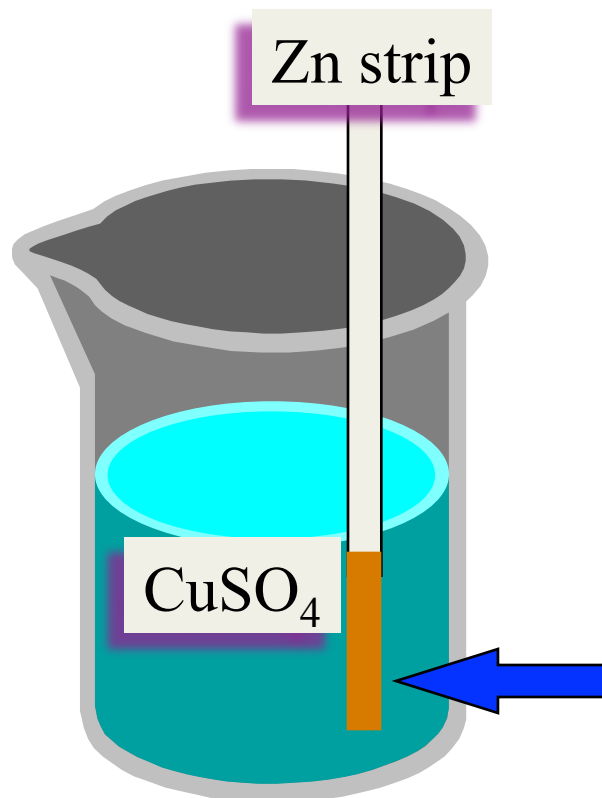
$$\text{Energy (Wh)} = I \times V \times t$$
$$\text{Power (W)} = I \times V$$

First battery



Total voltage: 1.1 V

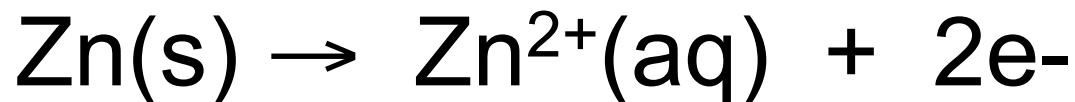
Copper Plating



With time, Cu plates out onto Zn metal strip.

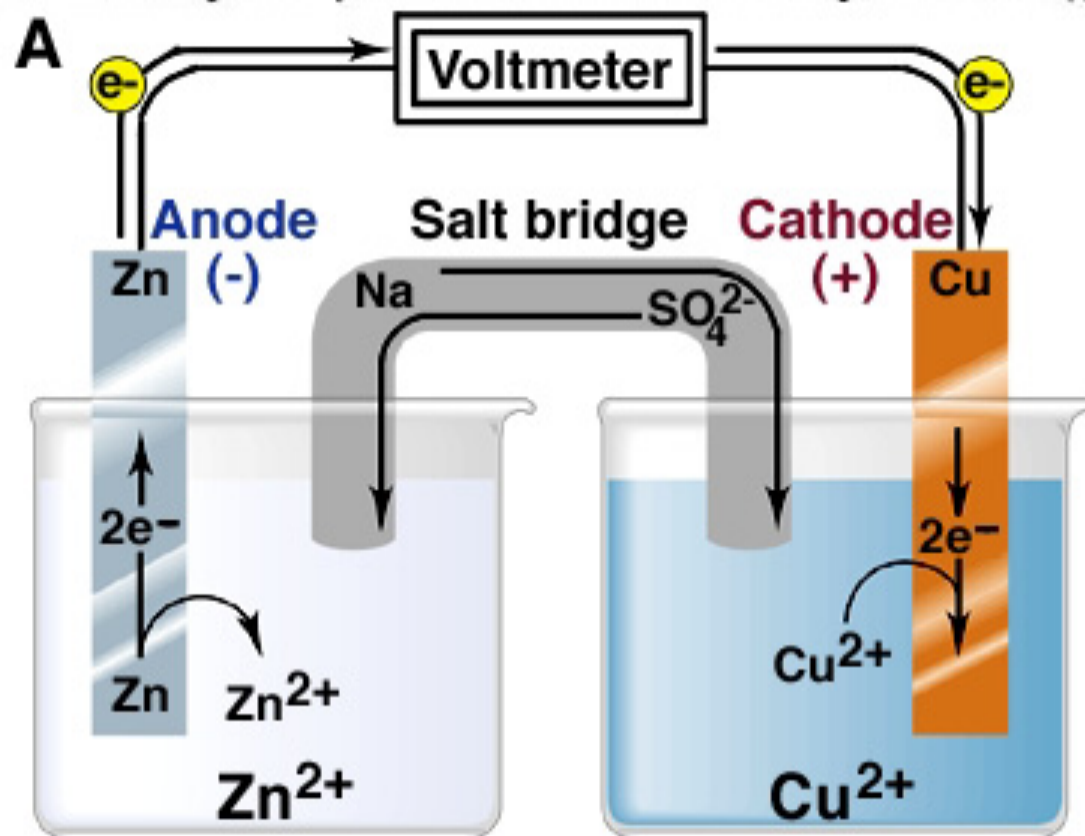
Electrons are transferred from Zn to Cu^{2+} , but there is no useful electric current.

- Zn is oxidized and is the reducing agent



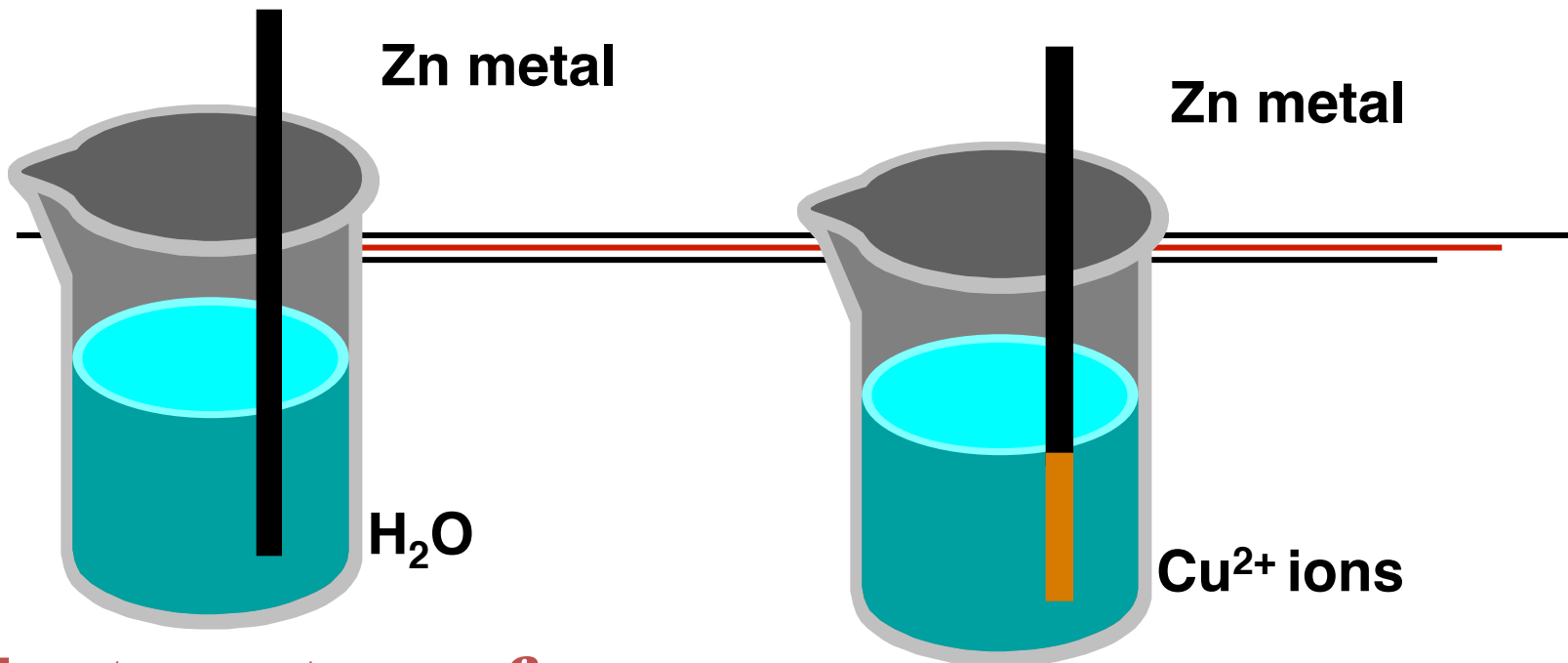
- Cu^{2+} is reduced and is the oxidizing agent



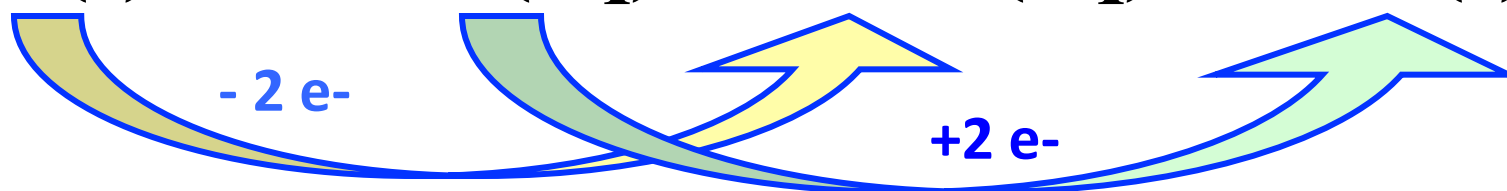


Zinc-Copper Reaction Voltaic Cell





Electron transfer



Loss of **E**lectrons =
OXIDATION (**LEO**)
Oil

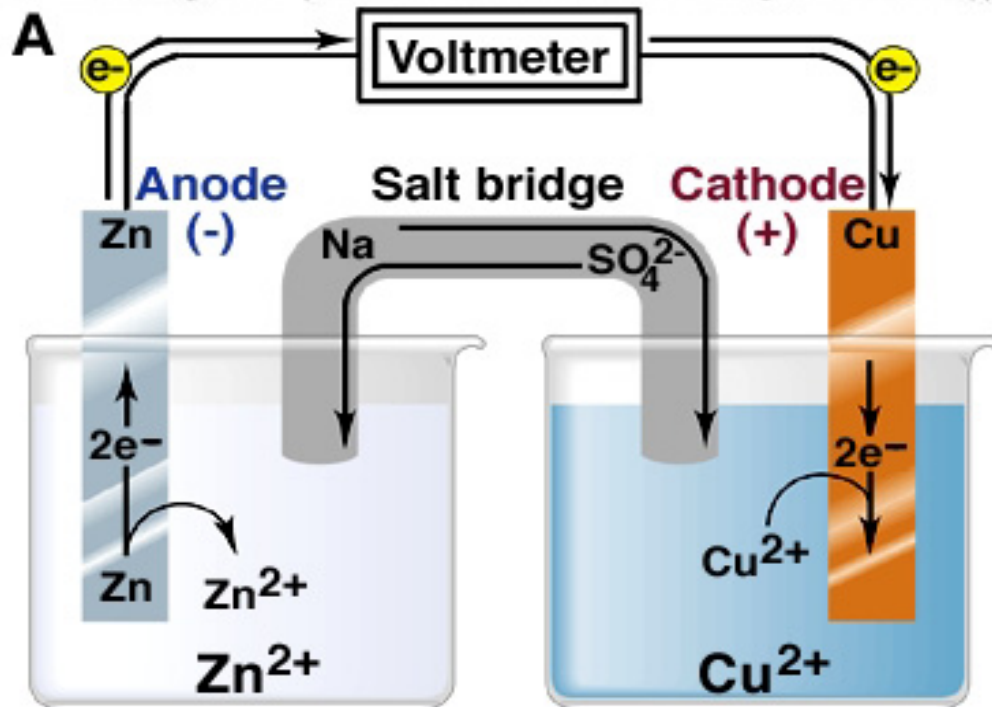
Gain of **E**lectrons =
REDUCTION (**GER**)
RiG



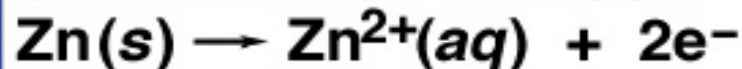
Massachusetts
Institute of
Technology

CHEMICAL CHANGE → ELECTRIC CURRENT

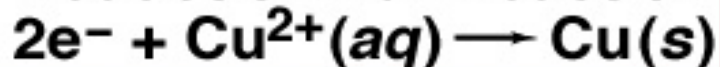
Martin S. Silberberg, *Chemistry: The Molecular Nature of Matter and Change*, 2nd Edition. Copyright © The McGraw-Hill Companies, Inc.



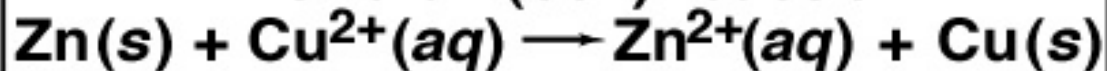
Oxidation half-reaction



Reduction half-reaction



Overall (cell) reaction



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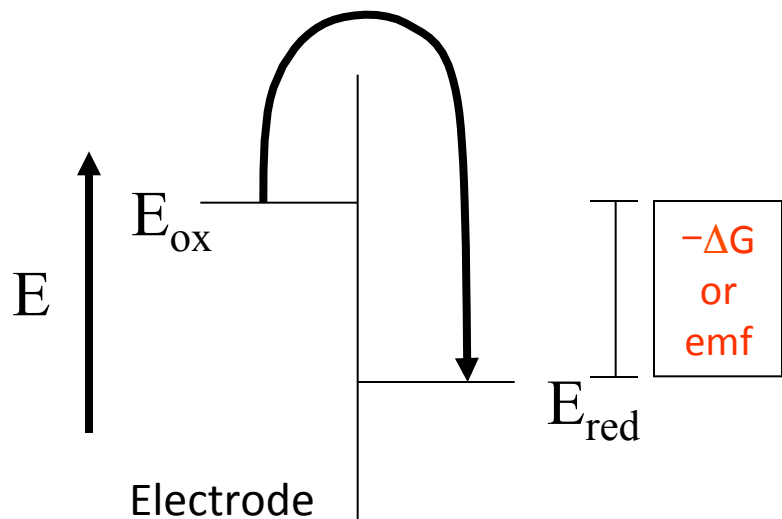
**Zinc-Copper
Reaction
Voltaic Cell**



Massachusetts
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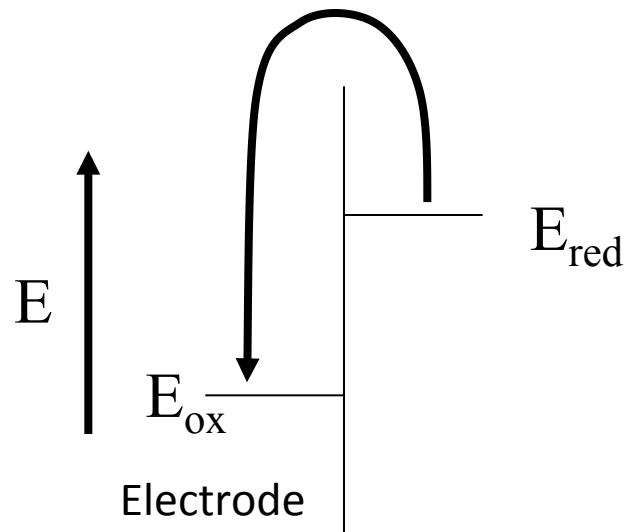
Why Electrons Transfer

Reduction



- Net flow of electrons from electrode to solute
- Electromotive force (emf) is the difference between the initial energy and reduced energy
- more cathodic
- more reducing

Oxidation



- Net flow of electrons from solute to electrode
- Positive ΔG , negative emf
- more anodic
- more oxidizing

Cell diagrams

Rather than drawing an entire cell, a type of shorthand can be used.

For our copper - zinc cell, it is:



The anode is always on the left.

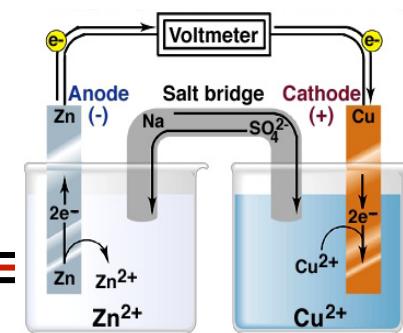
| = boundaries between phases

|| = salt bridge

Anode Left, Cathode **R**ight (**R**eduction, **R**eceiving)

Electrons flow left to right (in order of species)

Zn/Cu Electrochemical Cell

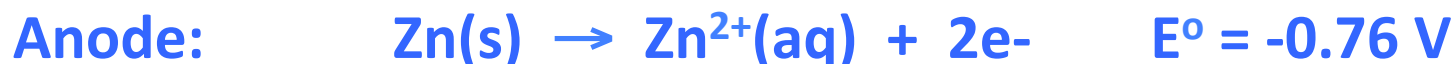


What is E° for the Zn/Cu cell (Daniel's cell) ??

$$E^\circ_{\text{cell}} = E^\circ_{\text{cathode}} - E^\circ_{\text{anode}}$$

Product gets electron
Reactant gives electron

Products - reactants



$$E_{\text{cell}} = E_{\text{cathode}} - E_{\text{anode}} = 0.34 - (-0.76) = +1.10 \text{ V}$$

E° and ΔG°

E° is related to ΔG° , the free energy change for the reaction for standard state (most stable form at 25°C and 100kPa).

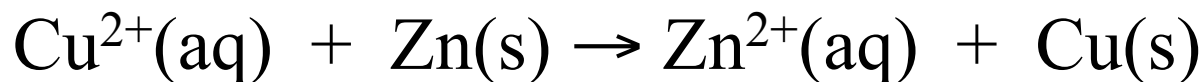
$$\Delta G^\circ = -n F E^\circ$$

F = Faraday constant
= 9.6485×10^4 C/mol

n = number of moles of e^- 's transferred.



n for Zn/Cu cell ? $n = 2$



Michael Faraday
1791-1867

Discoverer of

- electrolysis
- magnetic properties of matter
- electromagnetic induction (electric motor)

E° and ΔG°

$$\Delta G^\circ = -n F E^\circ$$

- For a **product-favored** reaction
 - Galvanic cell: Chemistry \rightarrow electric current

Reactants \rightarrow Products

$\Delta G^\circ < 0$ and so $E^\circ > 0$ (E° is positive)

- For a **reactant-favored** reaction
 - Electrolytic cell: Electric current \rightarrow chemistry

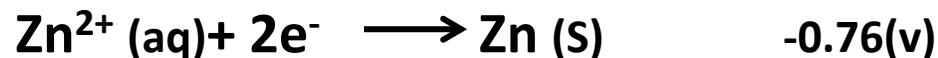
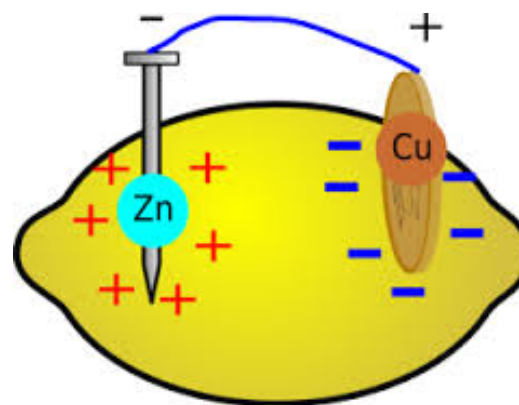
Reactants \leftarrow Products

$\Delta G^\circ > 0$ and so $E^\circ < 0$ (E° is negative)

Lemon Battery

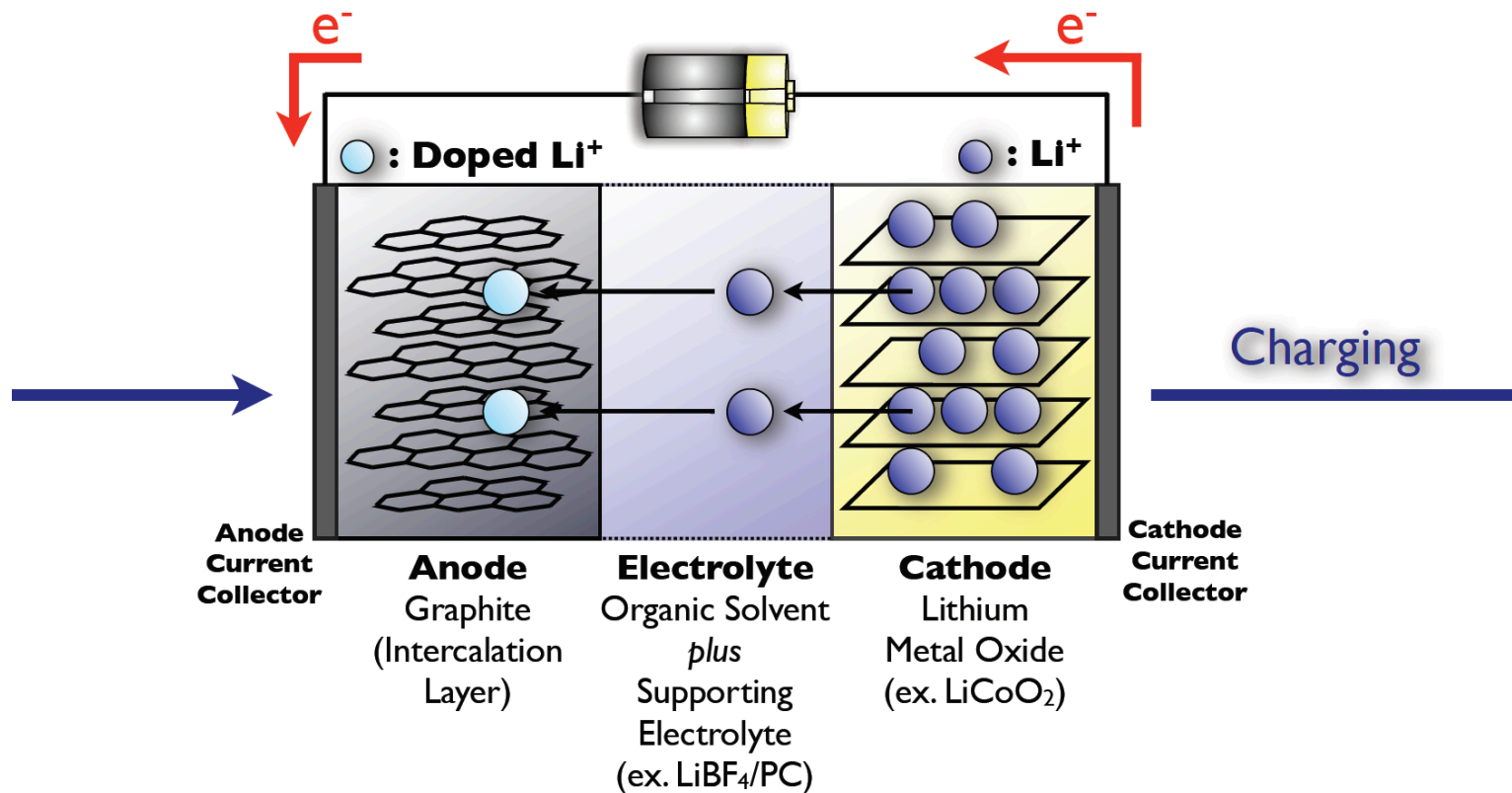
- Making a battery with a lemon, penny and galvanized nail (Zn-coated) as the electrodes

- Negative electrode?
- Positive electrode?
- Why do we need the lemon?
- What if we use 2 pennies?



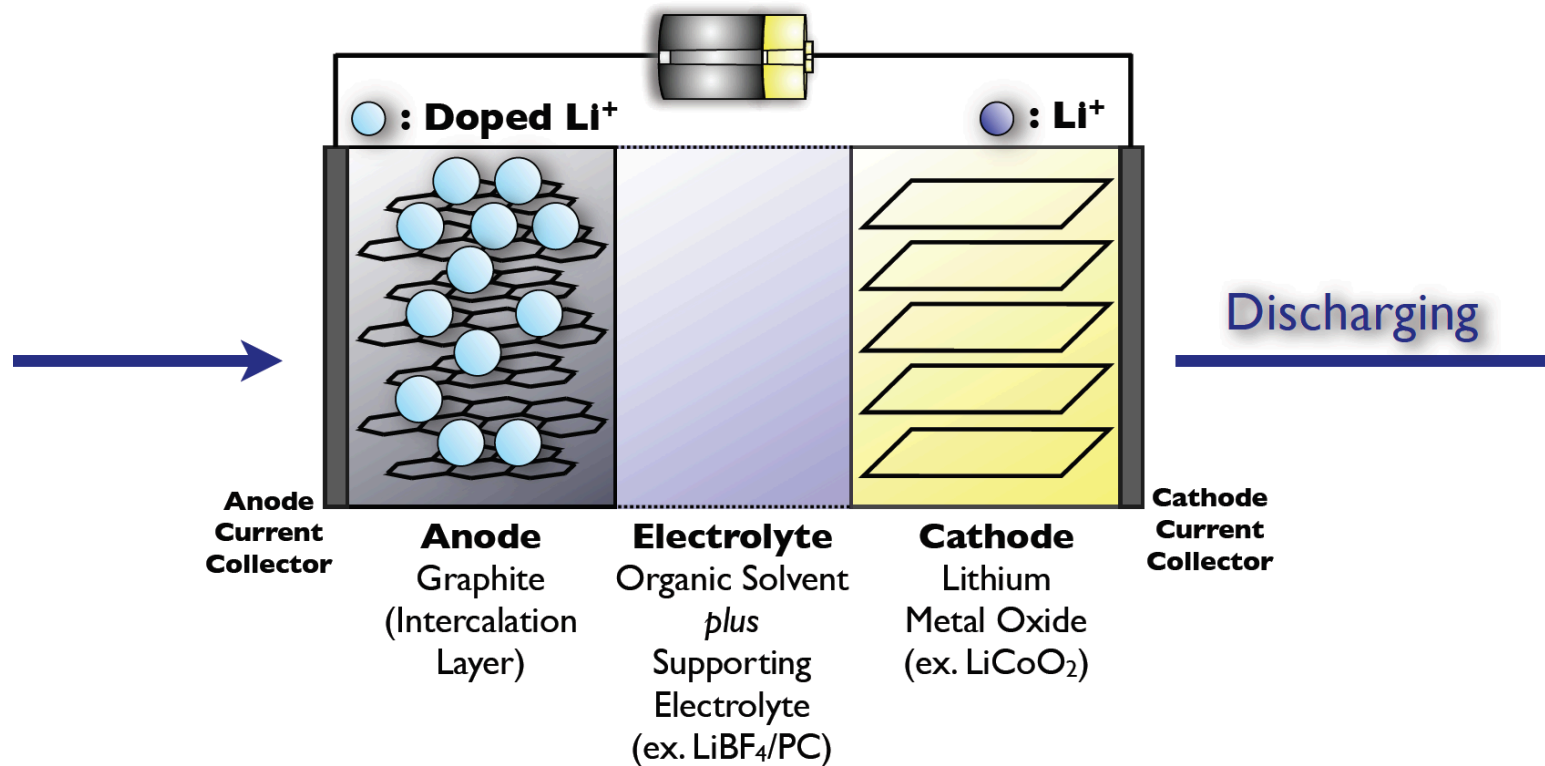
- What is the standard potential of the lemon battery?

An Intercalation-based Lithium Battery Cell



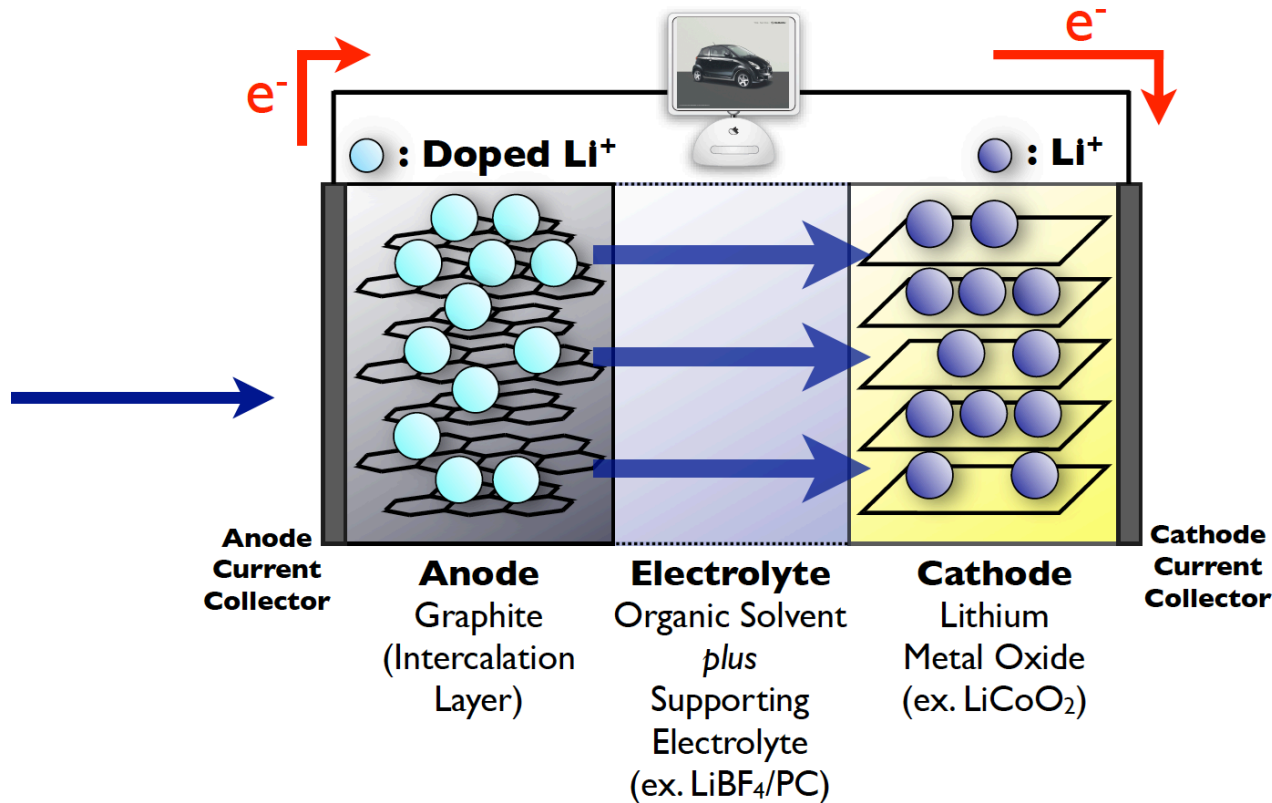
Reaction Mechanism for LIBs

Fully Charged State



Reaction Mechanism for LIBs

Fully Discharged State



What are the Batteries for Higher Energy Density LIBs?

Category	Material	Capacity (Ah/kg)	
		Theoretical	Actual
Cathode Material	LiCoO ₂	274	140
	LiMn ₂ O ₄	148	120
	LiV ₂ O ₅	142	140
	LiNiO ₂	275	200
	LiFePO ₄	170	150
Anode Material	Carbon (LiC ₆)	372	-
	Lithium	3861	-

Capacity imbalance between cathode and anode will increase further so that higher capacity cathode materials to match anode capacity will be required. Furthermore, for transportation, inexpensive and abundant materials will also be required.