

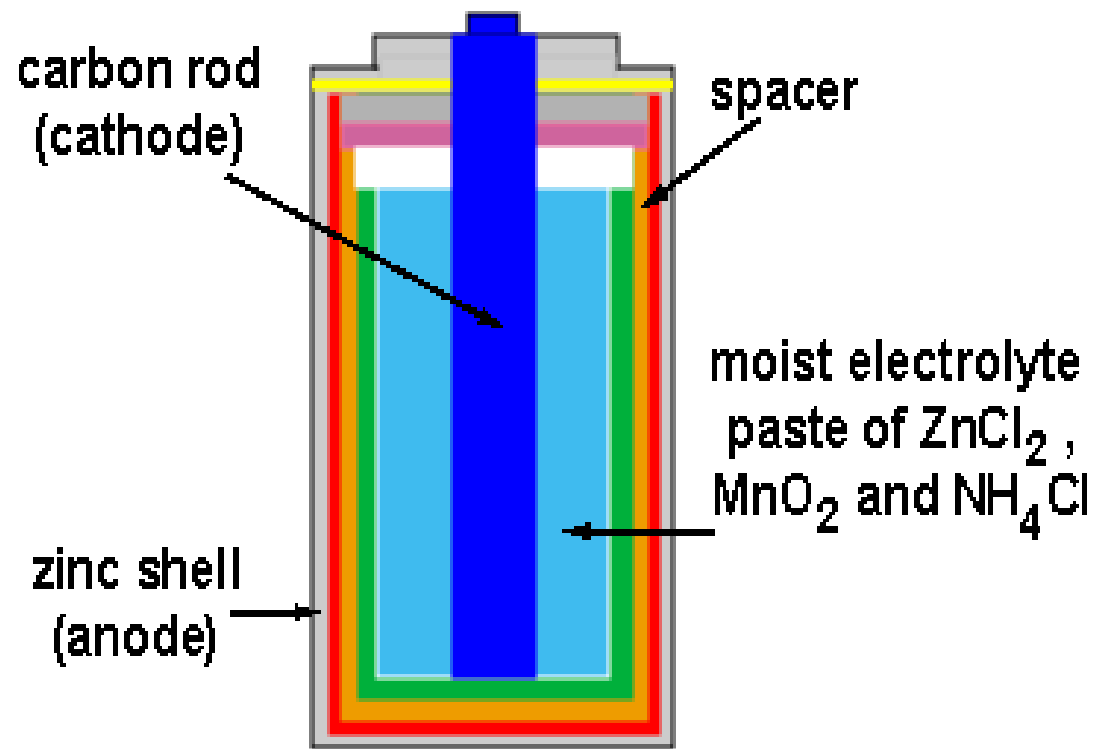
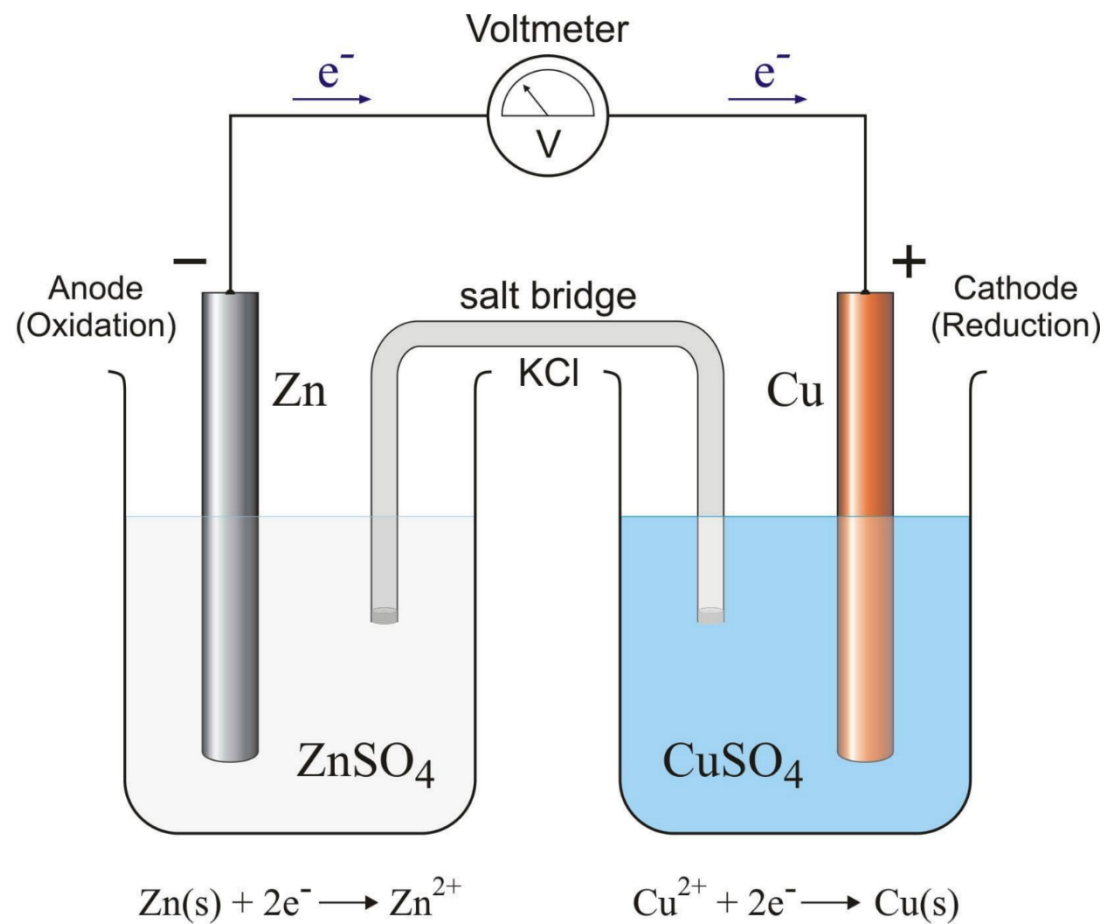
# **N44 - Electrochemistry**

## **Intro to Cells**

**Target: I can label cells and perform calculations for both Galvanic and Electrolytic cells.**

# **N44 - Electrochemistry**

## **Intro to Cells**



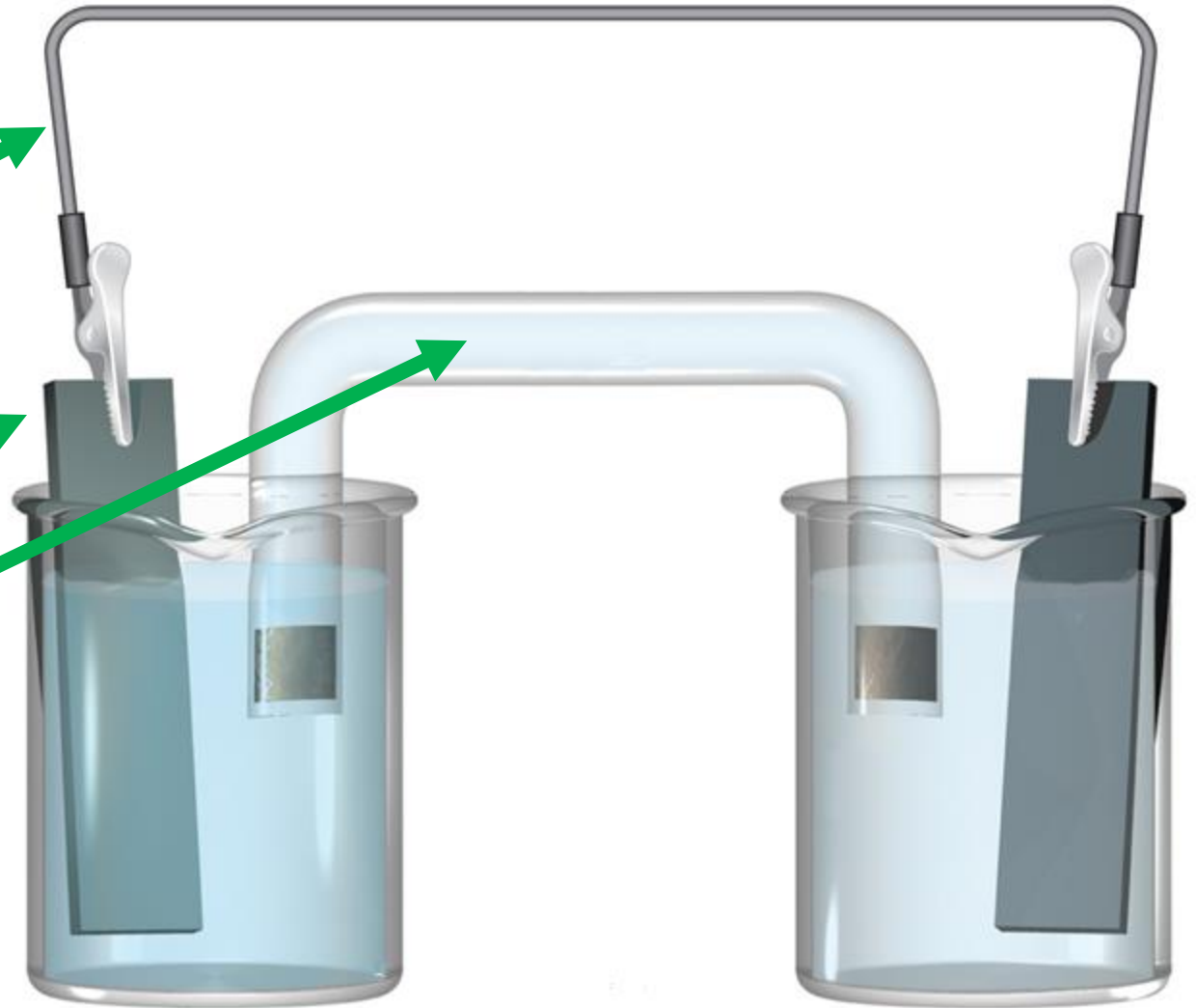
# Galvanic/Voltaic Cells

- **Thermodynamically favorable**
- **Reaction will take place spontaneously**

# Example: Mn and Ni

**Start with basics**

- Beakers
- Wire
- Electrodes
- Salt bridge



# Identifying Oxidation and Reduction

## Mn and Ni electrodes

Which will oxidize, which will reduce?

**Look at reduction tables!**



# Identifying Anode, Cathode, Electrodes

## Anode

e- are being lost.



LEO

Oxidation

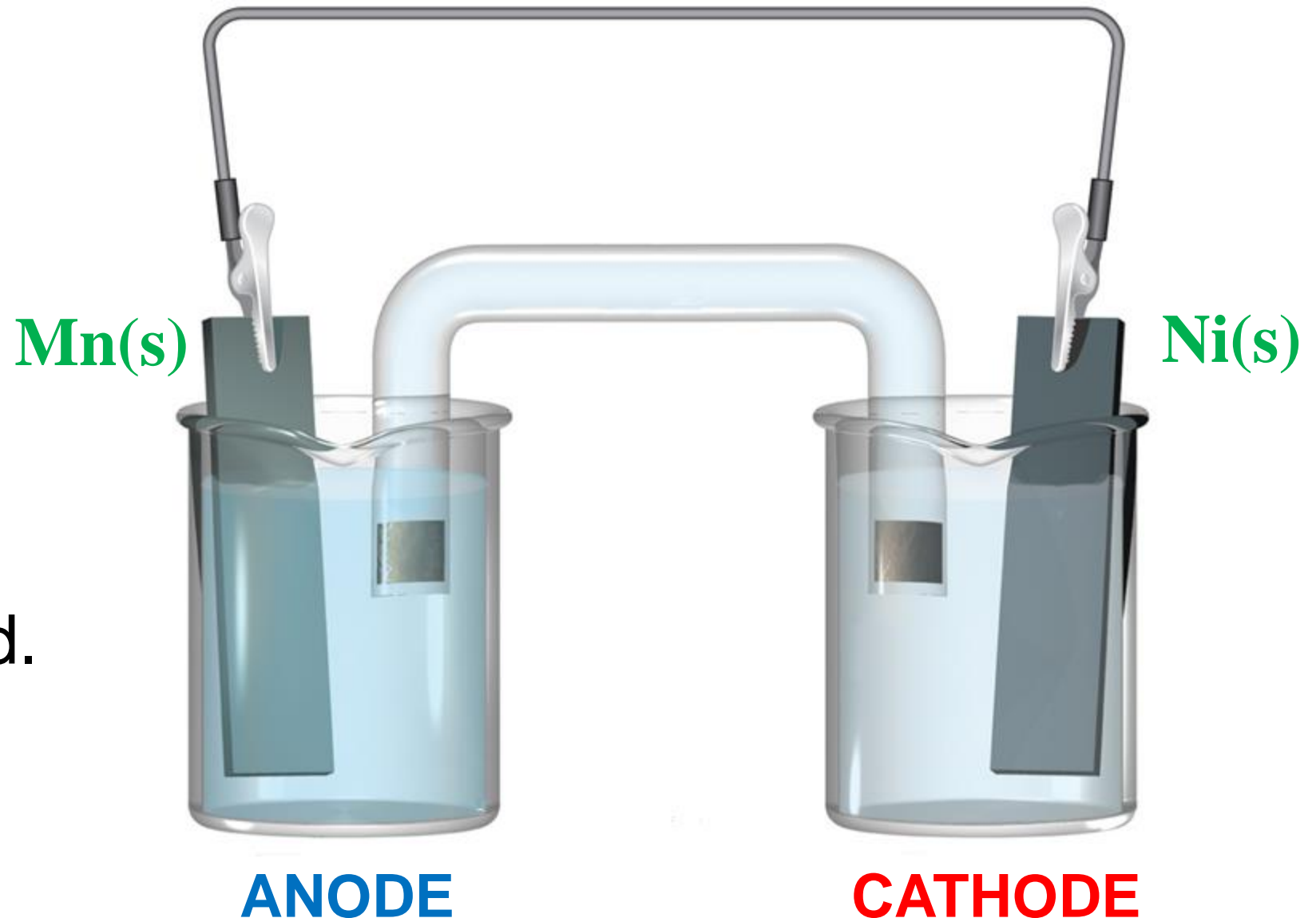
## Cathode

e- are being gained.



GER

Reduction



# Identifying Solutions

## Anode

e- are being lost.



LEO

Oxidation

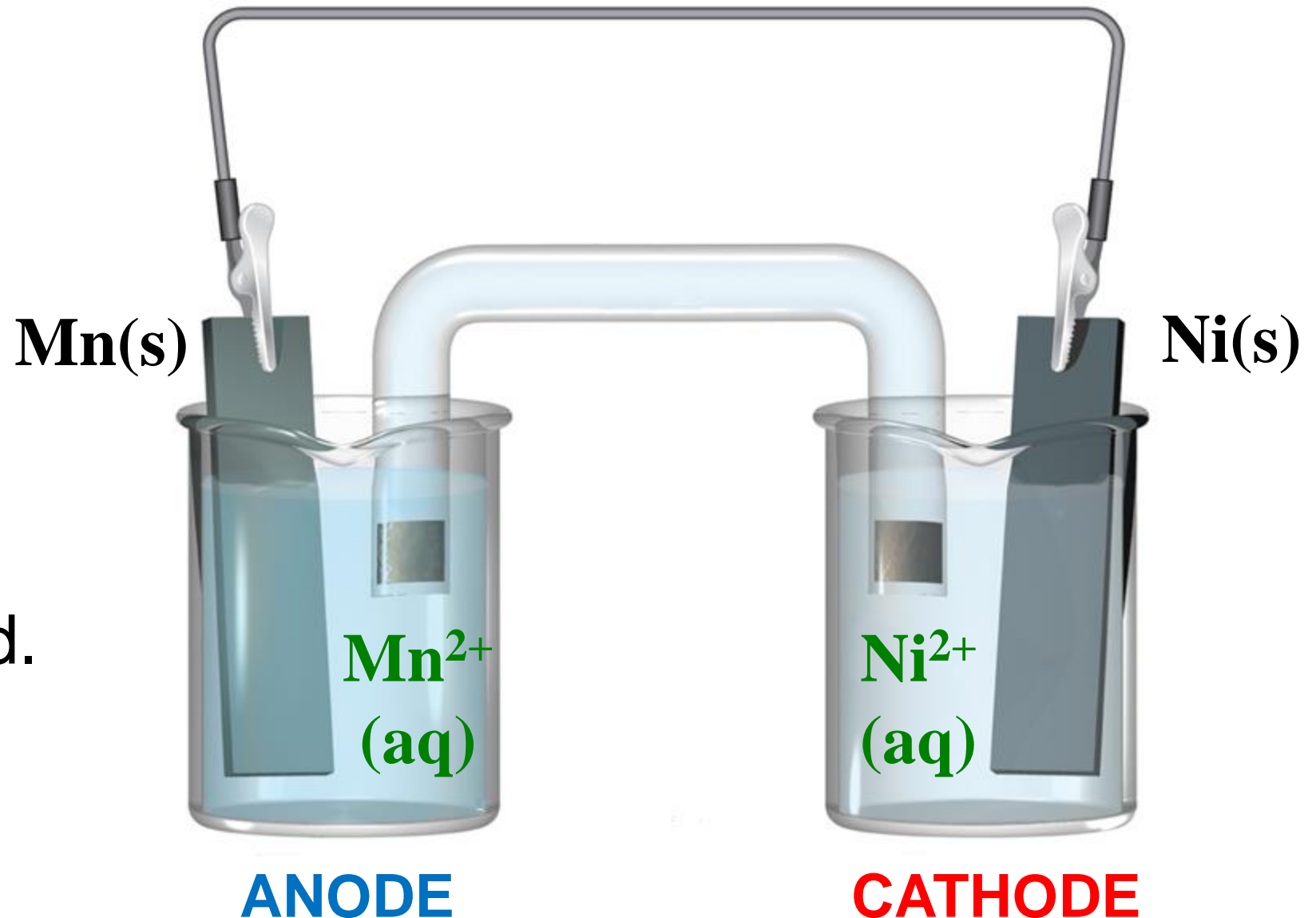
## Cathode

e- are being gained.



GER

Reduction





# Identifying Direction of Current (e<sup>-</sup>) Flow

## Anode

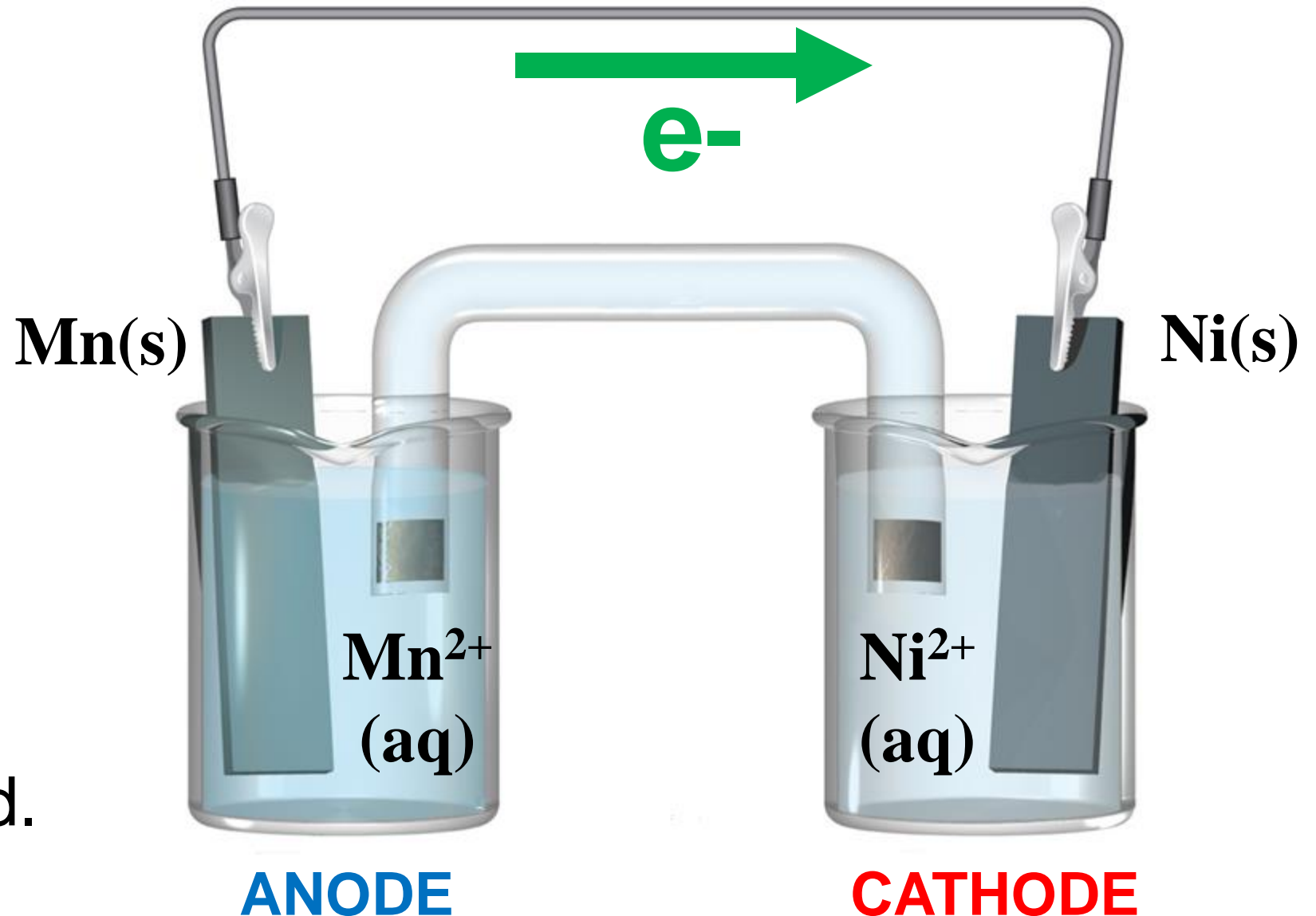
e<sup>-</sup> are being lost.



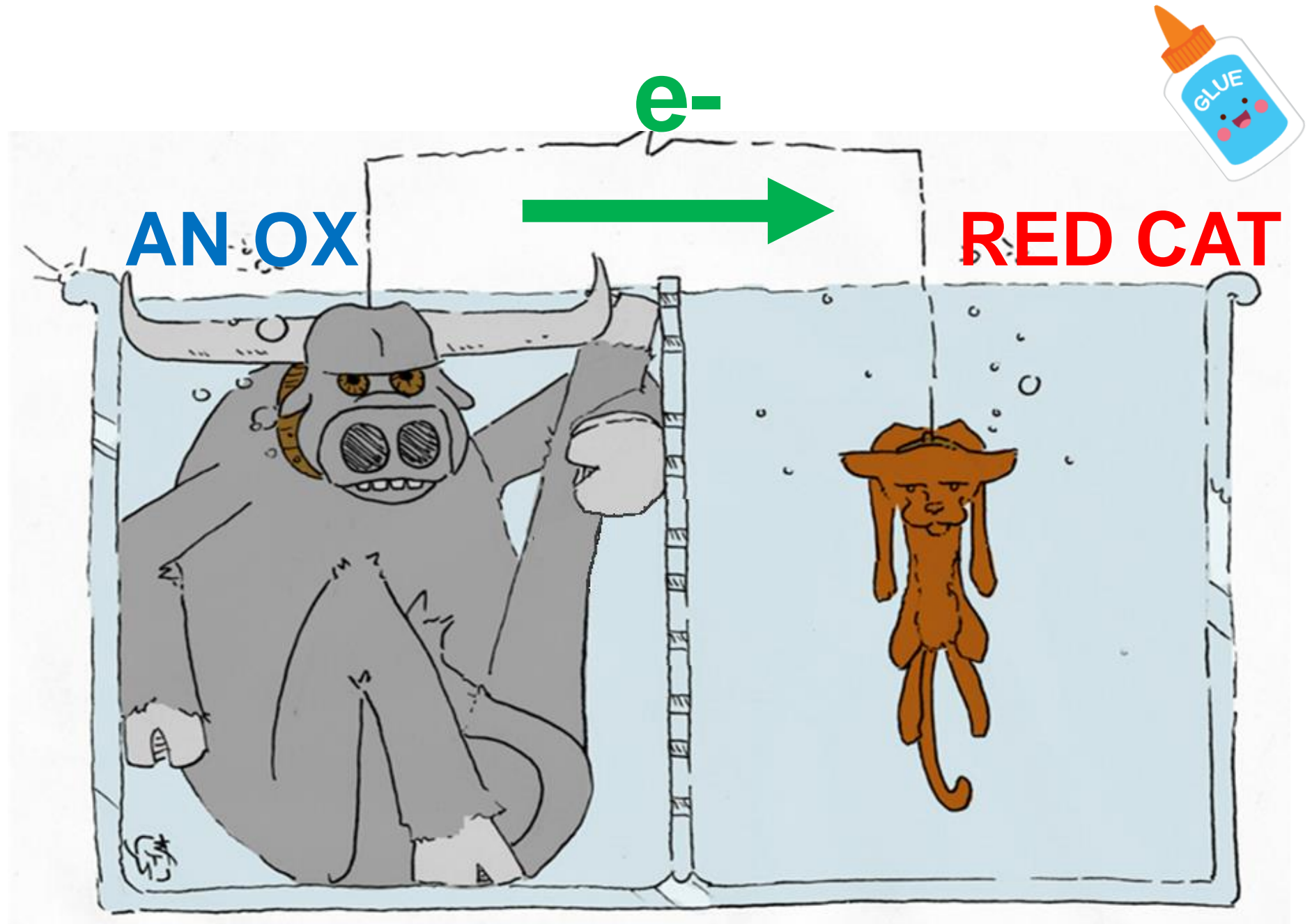
They leave anode  
and flow towards  
the Cathode.

## Cathode

e<sup>-</sup> are being gained.



a **RED**  
**CAT**  
gains,  
what  
**AN OX**  
loses



# Identifying Signs on Electrodes

## Anode

e- are being lost.

Source of e-

“e- rich”

- sign

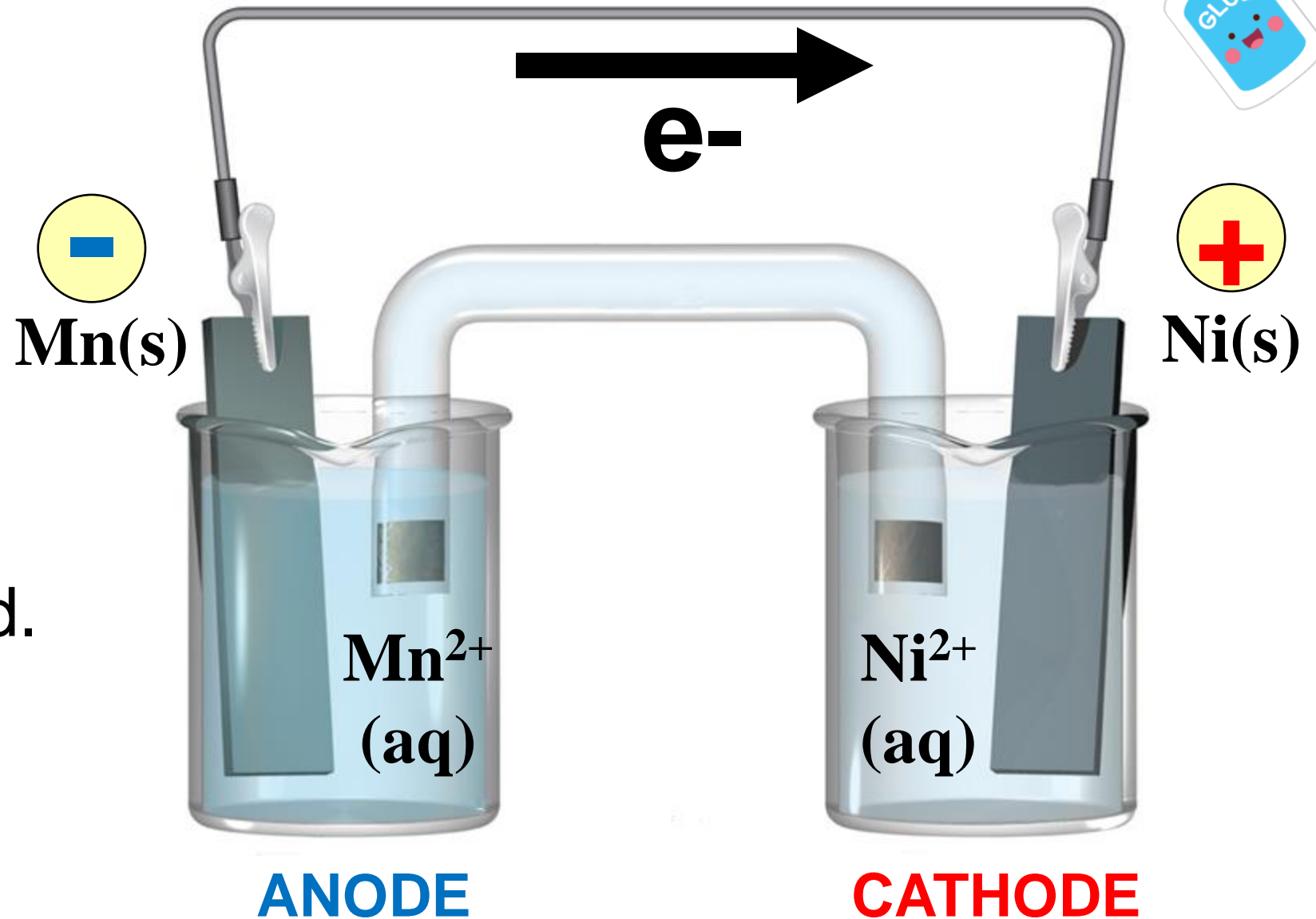
## Cathode

e- are being gained.

Drawing e-

“e- deficient”

+ sign



# Identifying $\Delta$ Mass of Electrodes

## Anode



Solid Mn turning into  $\text{Mn}^{2+}(\text{aq})$

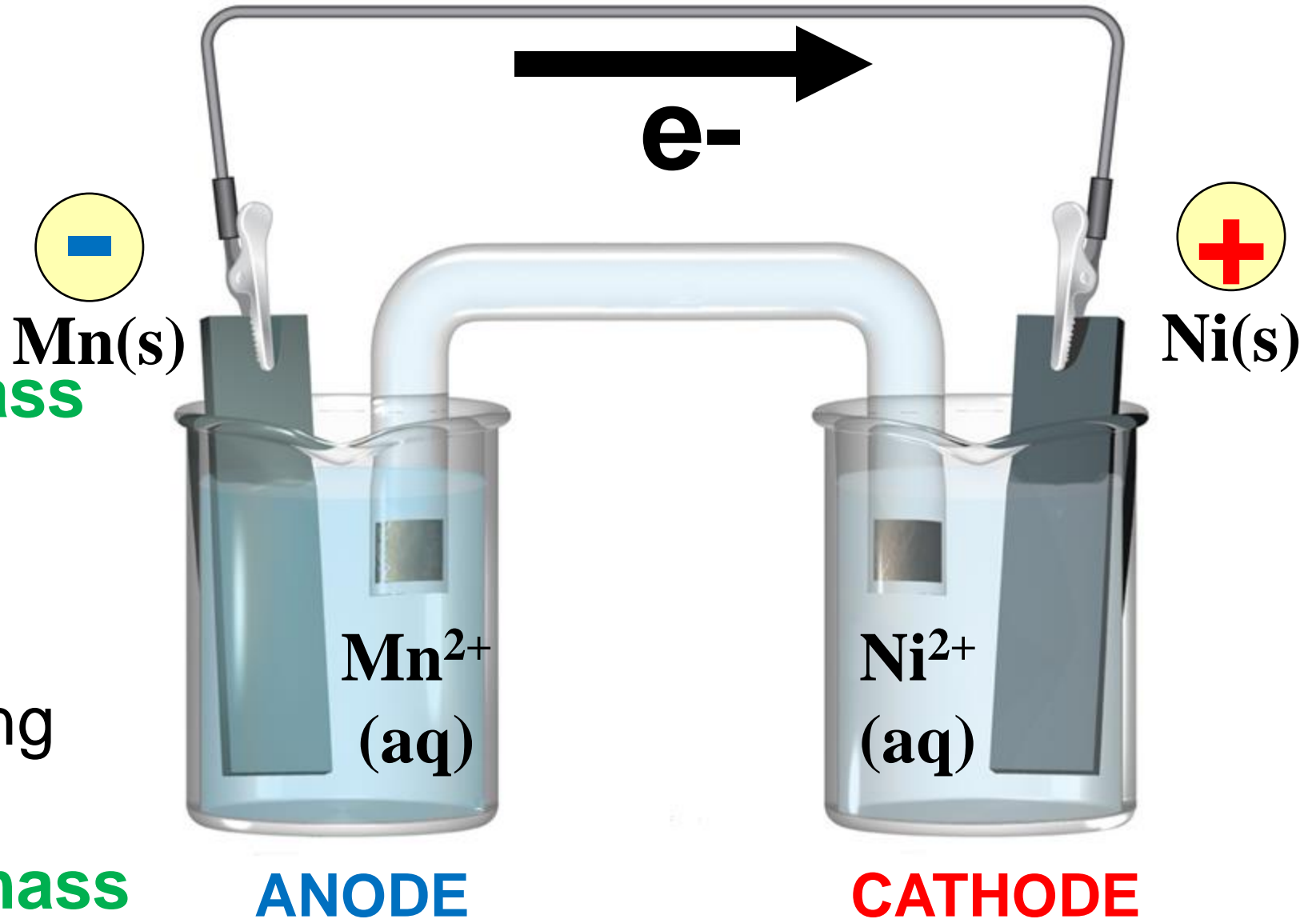
**Anode LOSES mass**

## Cathode



Aqueous  $\text{Ni}^{2+}$  turning into solid Ni metal

**Cathode GAINS mass**

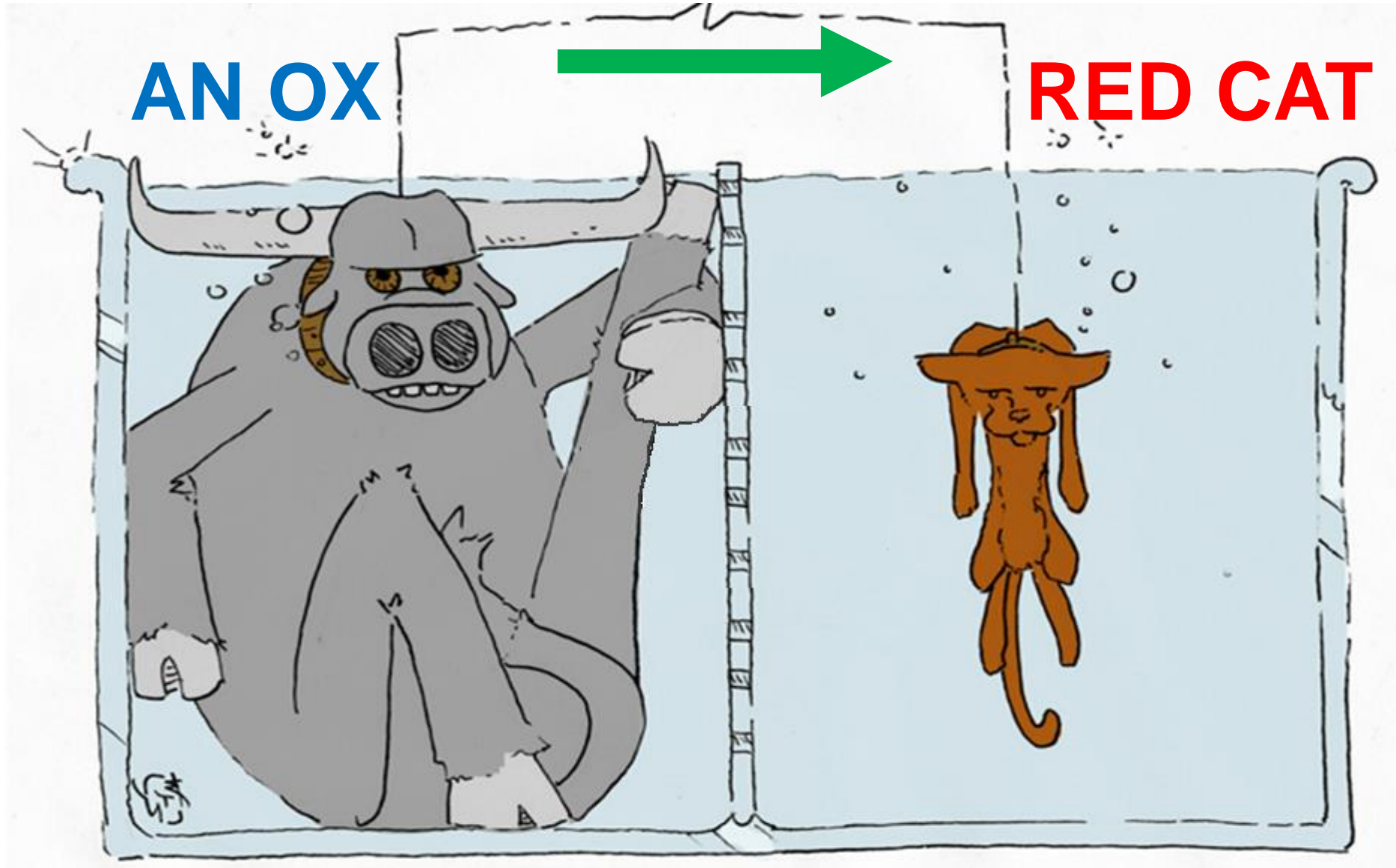


Loses Mass

Gains Mass

AN OX

RED CAT



a **RED**  
**CAT**  
gains,  
what  
**AN OX**  
loses

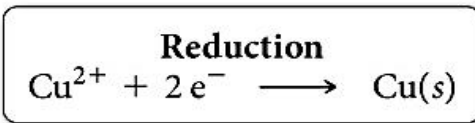
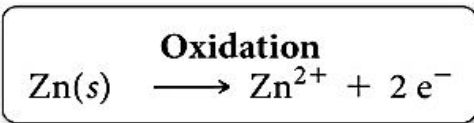
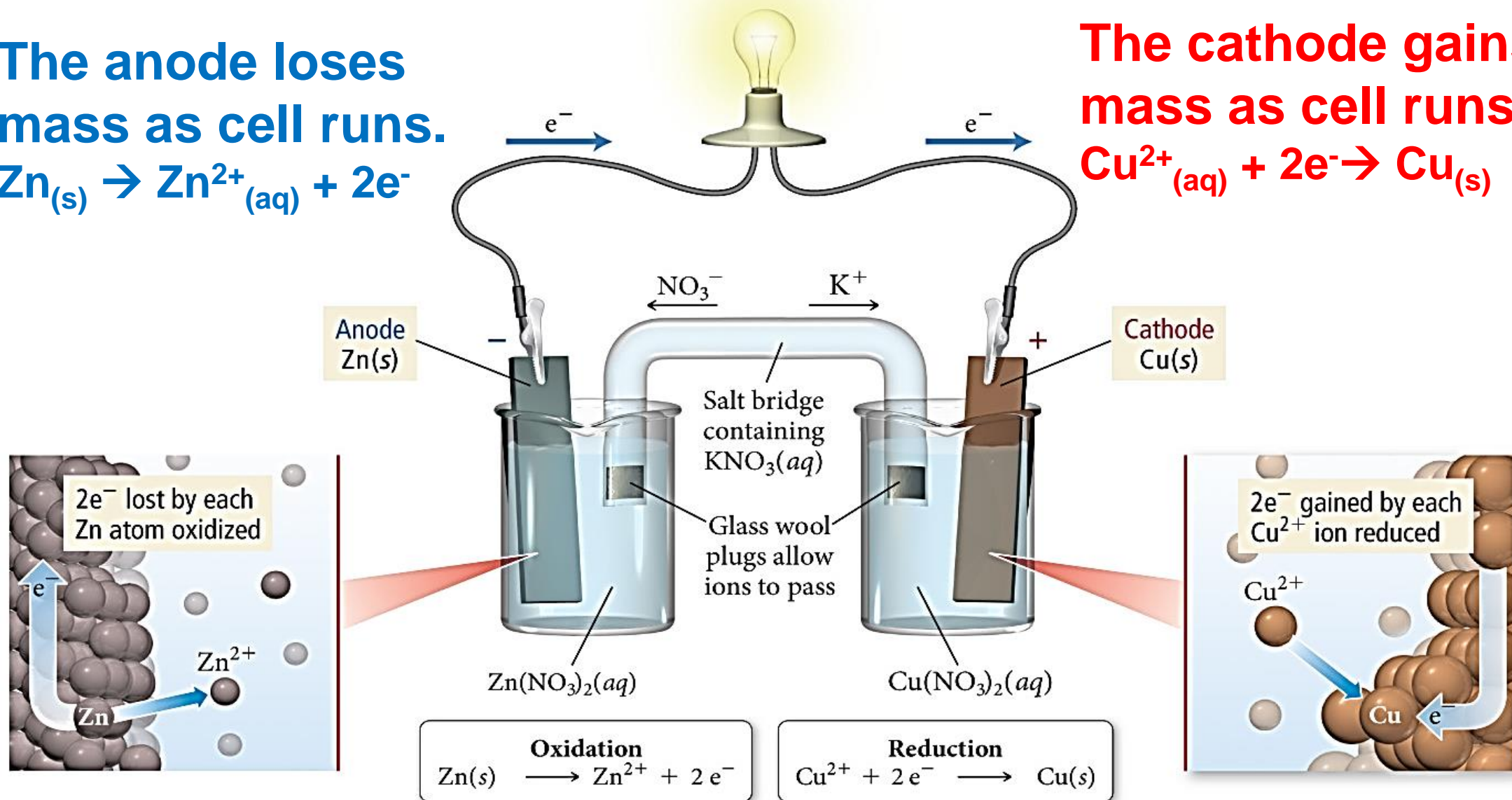
# Atomic View (different electrodes)



The anode loses mass as cell runs.



The cathode gains mass as cell runs.



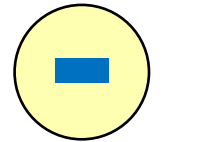
# Identifying Charge Balance – Salt Bridge

## Anode



Making more  $\text{Mn}^{2+}$

**Need anions** to  
balance new charge



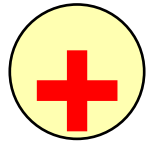
**Mn(s)**

## Cathode

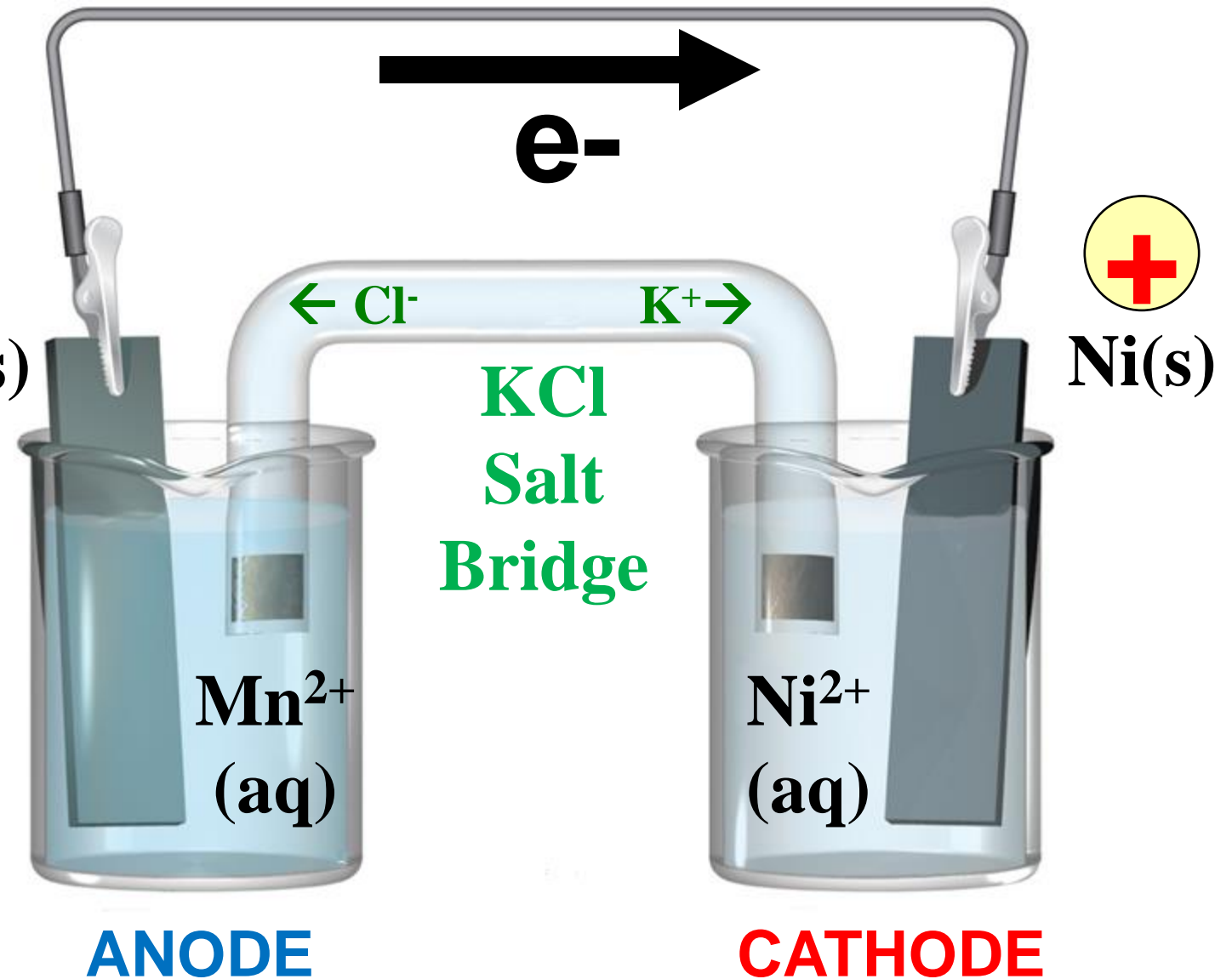


Using up  $\text{Ni}^{2+}$

**Need cations** to  
balance new charge



**Ni(s)**

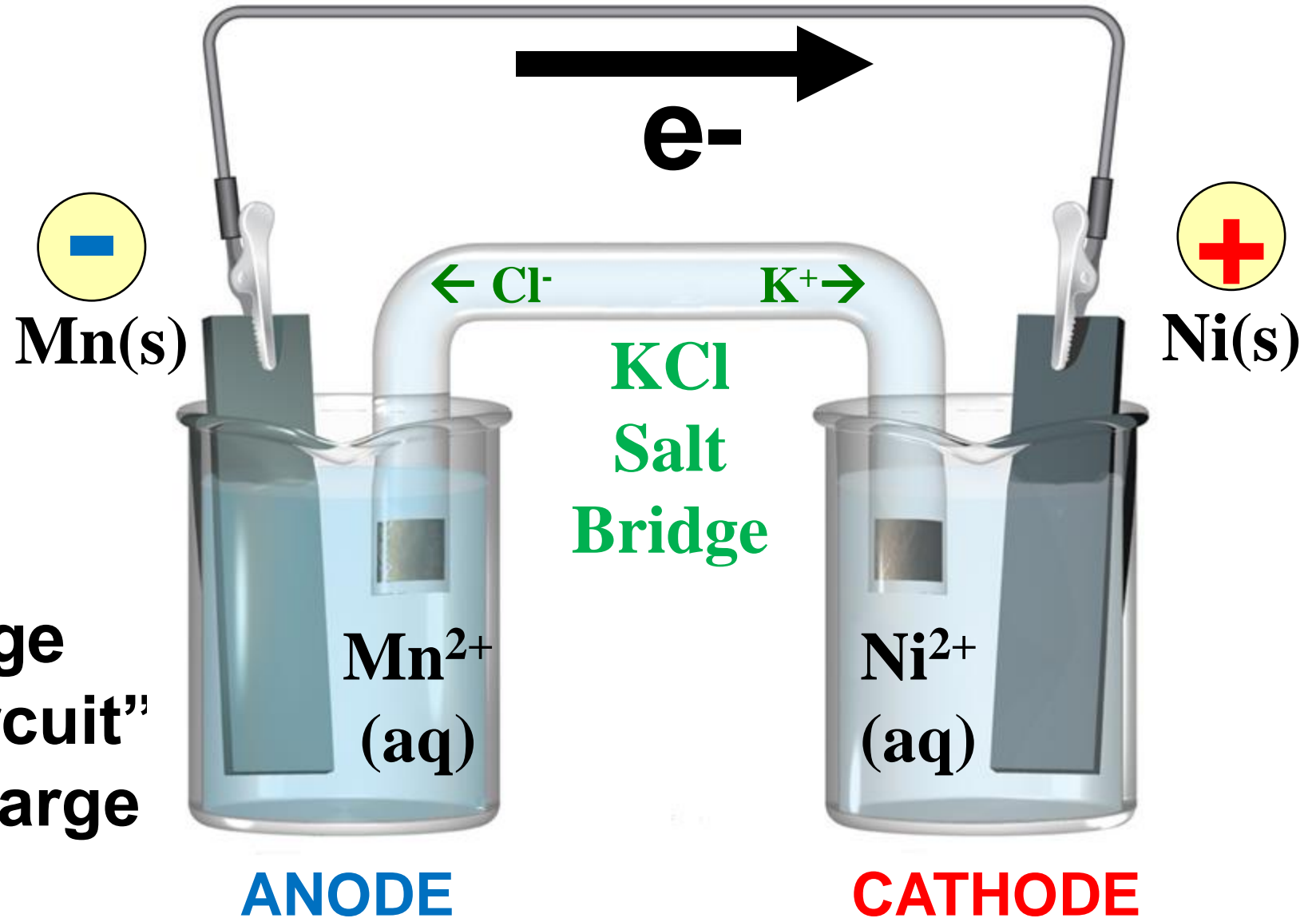


# Identifying Charge Balance – Salt Bridge

No salt bridge?  
Charge becomes  
imbalanced after  
some time...

Cell dies!!!! ☹️

Required salt bridge  
“completes the circuit”  
and “maintains charge  
balance”



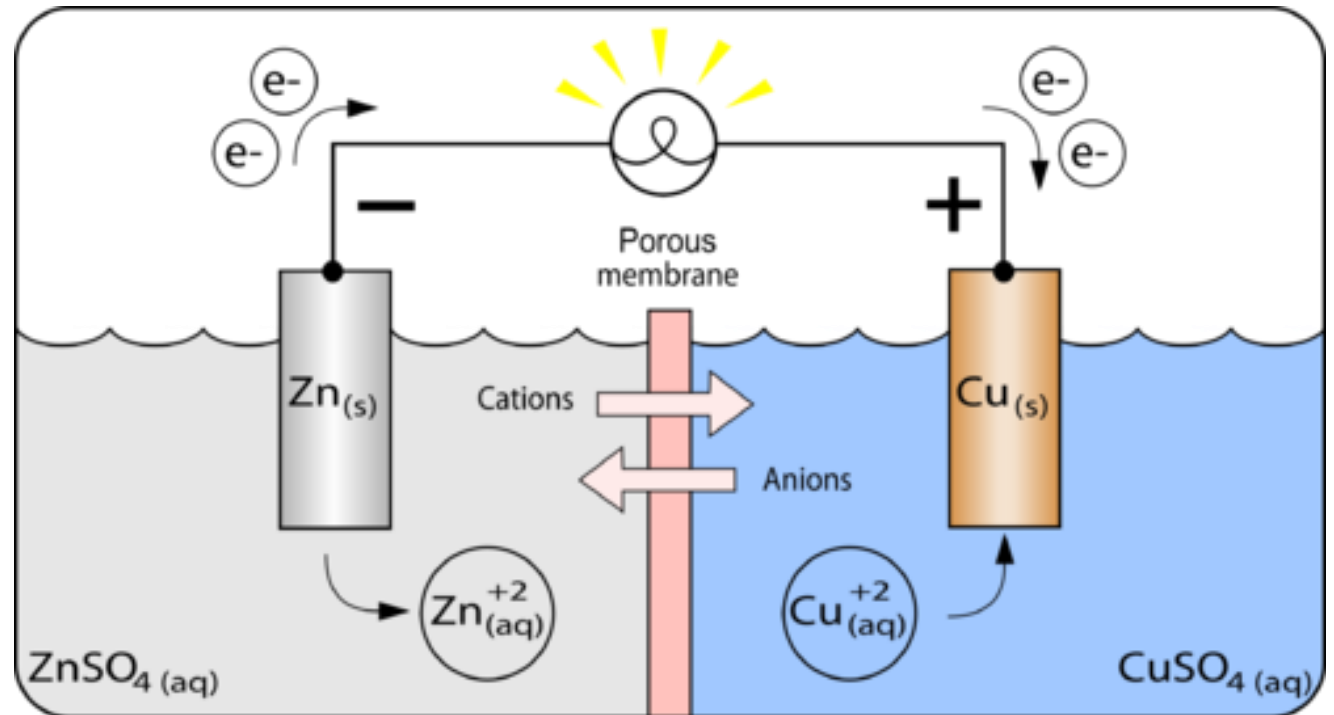


# Identifying Charge Balance – Membranes

Sometimes you have a **porous membrane partition** instead of a salt bridge

**Allows select ions to flow between sections**

**Still allows charge balance and complete circuit**



# Summary of Electrodes

## Anode

- Electrode where oxidation occurs (loss of  $e^-$ )
- Anions from salt bridge attracted to it because cations being made.
- Loses weight

## Cathode

- Electrode where reduction occurs (gain of  $e^-$ )
- Cations from salt bridge attracted to it because losing cations.
- Gains weight
  - Electrode where “plating” takes place in electroplating

# Electrolytic Cells

- **NOT** Thermodynamically favorable
- Reaction will **NOT** take place spontaneously
- Requires an outside electrical source to force it to happen!

# What would you predict for Red and Ox?

Zn and Cu



Oxidize! Lose  $\text{e}^{-}$



Reduce! Gain  $\text{e}^{-}$

**BUT!**

What if you **WANT** to do the opposite???

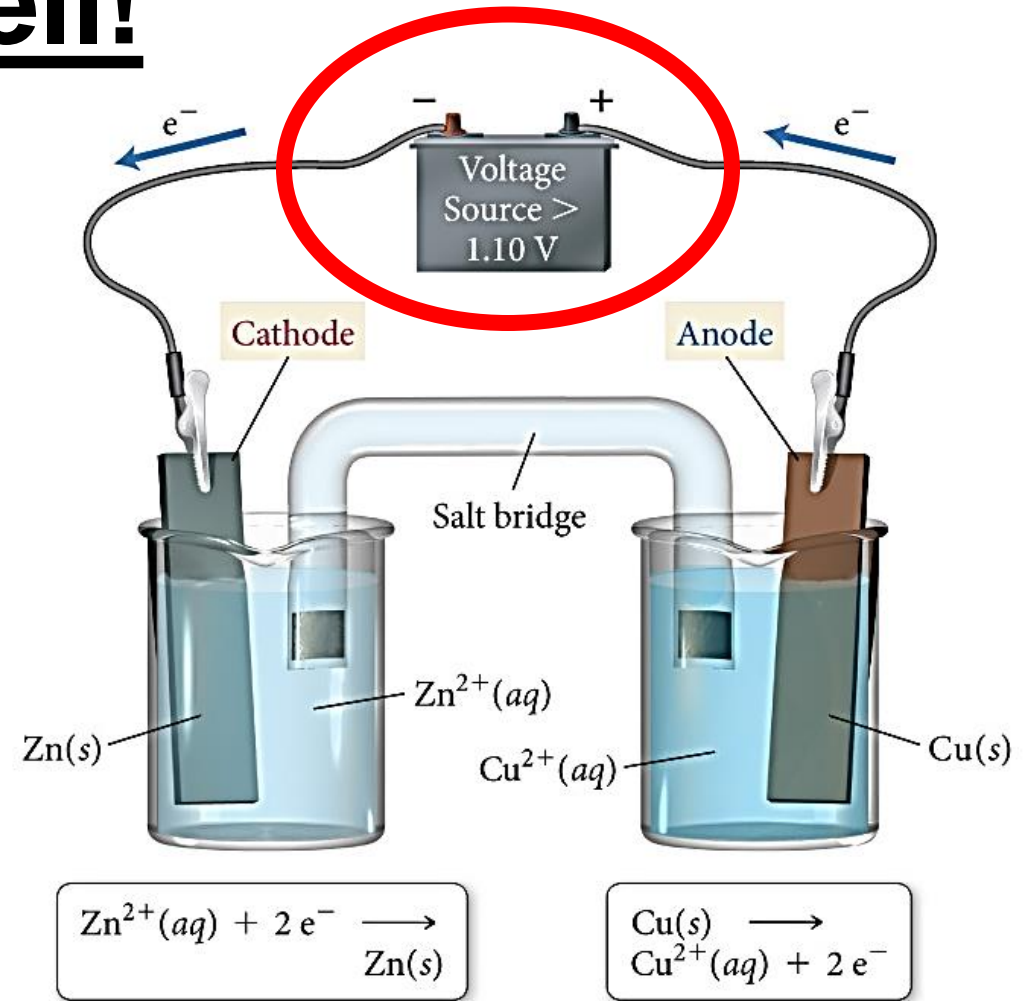
What if you want to make Zn(s) not Cu(s)???

# Make it an Electrolytic Cell!

Force it to go the other way,  
the non-thermodynamically  
favorable way!

Hook it up to an outside  
electrical source.

Will **DRIVE** the electrons the  
other direction, against their  
normal flow.



# Make it an Electrolytic Cell!

**STILL** label loss of e<sup>-</sup> as anode

**STILL** label gain of e<sup>-</sup> as cathode

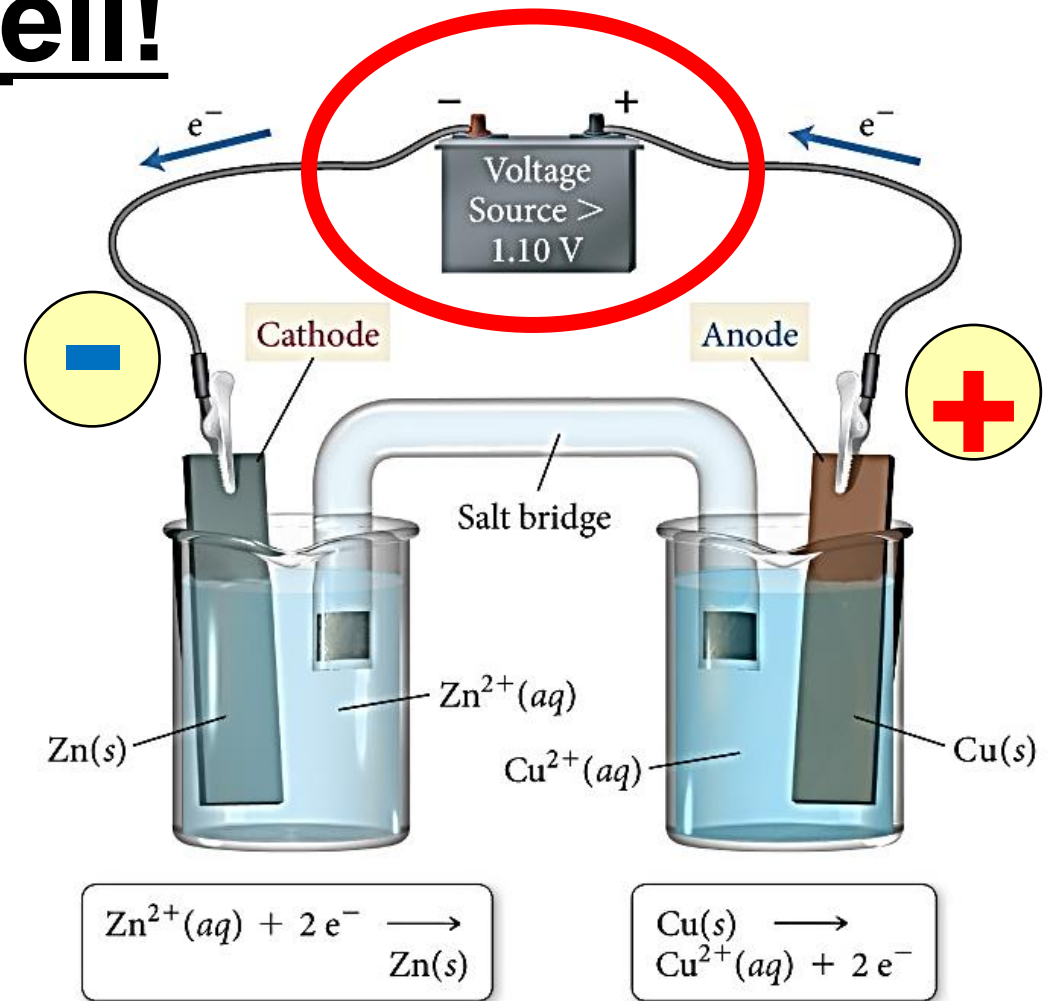
**STILL** show e<sup>-</sup> flow to cathode

**STILL** use salt bridge the same  
anions towards anode  
cations towards cathode

**BUT...**

**Anode +      Cathode –**

e<sup>-</sup> coming off anode need to go to  
the + end of the voltage source, and  
they leave – end of voltage source



# Galvanic versus Electrolytic Cells



## Galvanic

- Converts chemical energy into electrical energy.
- Positive cell potential,  $E^{\circ}_{\text{cell}} = +$
- Spontaneous,  $\Delta G = -$
- Anode = - and Cathode = +
- Electrons supplied by the chemical being oxidized.
- Electrons flow from anode to cathode.

## Electrolytic

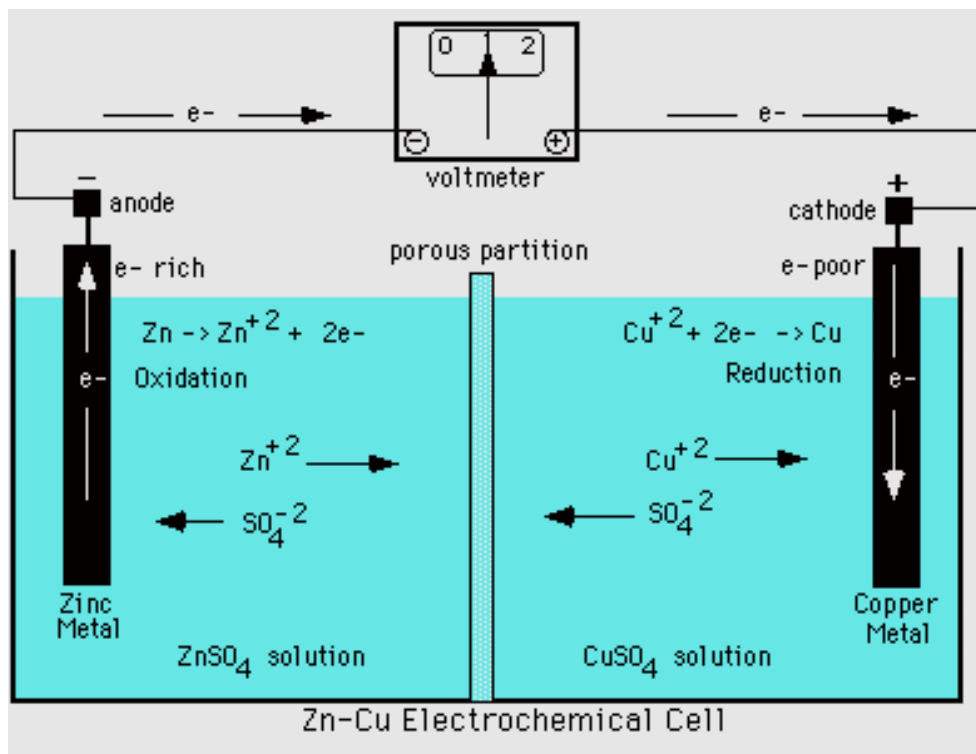
- Converts electrical energy into chemical energy
- Negative cell potential,  $E^{\circ}_{\text{cell}} = -$
- NOT spontaneous,  $\Delta G = +$
- Anode = + and Cathode = -
- Electrons supplied by an external source
- Electrons enter from the cathode and come out at the anode.

# Galvanic or Electrolytic?

Zn - Cu



From a table of reduction potentials:



Cu more positive = reduced

Zn more negative = oxidized

*flip eq and sign on E for Zn*



---

$$E = (+0.76) + (+0.34) = +1.10 \text{ V}$$

**E = + so galvanic, spontaneous,  $\Delta G = -$**



# Galvanic or Electrolytic?

A cell is undergoing this rxn:



**Careful!**

Equation is telling you that Na is going from  $\text{Na}^+ \rightarrow \text{Na}$  It is being reduced even though the table shows it would rather be oxidized!!!!

And  $2\text{Cl}^- \rightarrow \text{Cl}_2$  so it is being oxidized even though it would rather be reduced!



Told that  $\text{Na}^+$  = reduced

Told that  $\text{Cl}^-$  = oxidized

*flip eq and sign on E for Cl*



---

$$E = (-2.71) + (-1.36) = -4.07\text{V}$$

**E = - so electrolytic, NON-spontaneous,  $\Delta G = +$**

# Galvanic or Electrolytic?

A cell is undergoing this rxn:



This cell won't run by itself! You need to hook it up to an outside electrical supply!



Told that  $\text{Na}^+$  = reduced

Told that  $\text{Cl}^-$  = oxidized

*flip eq and sign on E for Cl*



---

$$E = (-2.71) + (-1.36) = -4.07\text{V}$$

**E = - so electrolytic, NON-spontaneous,  $\Delta G = +$**

# Cell (Line) Notation

Shorthand description of a voltaic cell

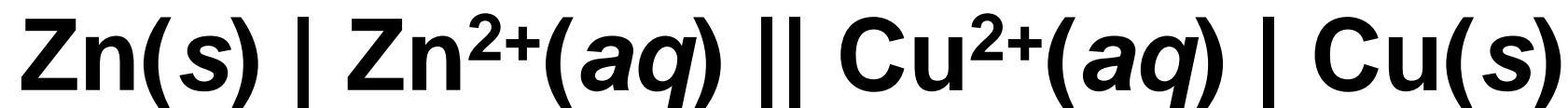
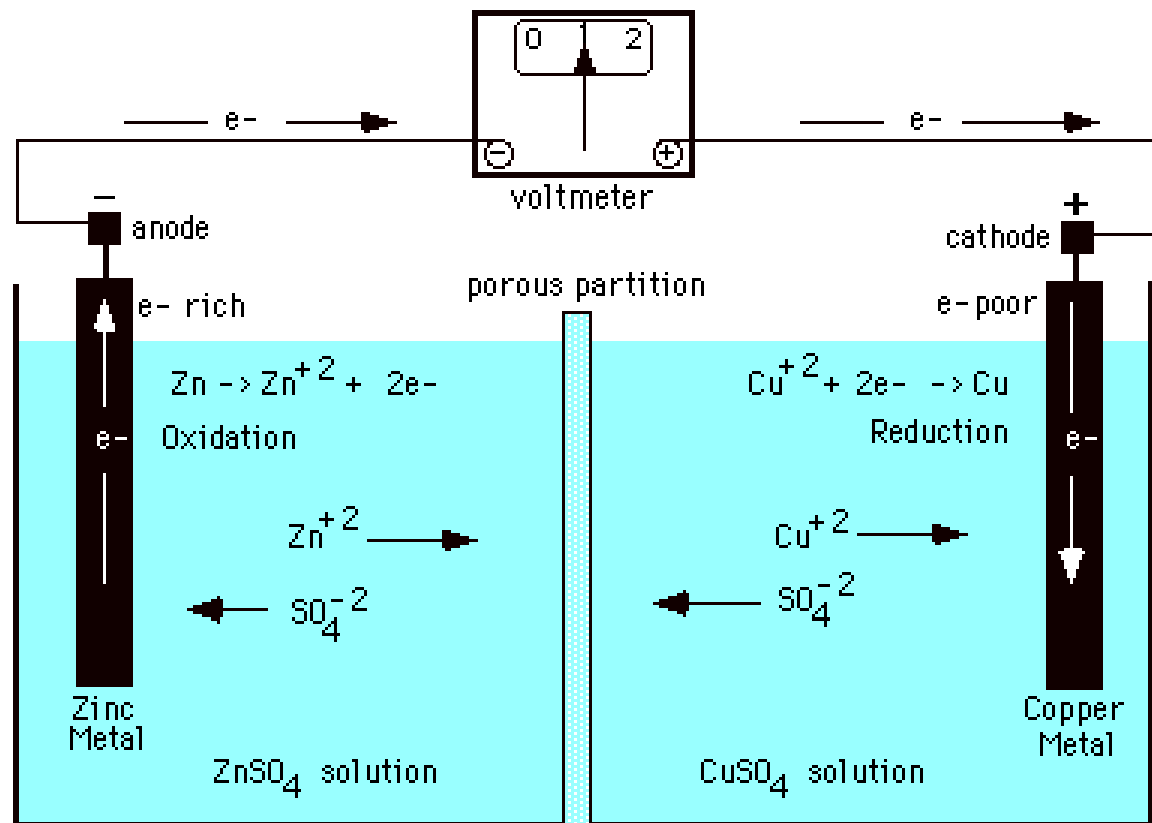
**Electrode | electrolyte || electrolyte | electrode**

*Oxidation half cell side*

*Reduction half cell side*

- Single | = phase barrier
  - If multiple electrolytes in same phase, a comma is used rather than |
  - Often use an **inert electrode**
  - Sometimes they put the concentrations in also
- Double line || = salt bridge

# Line Notation



Anode  
material

Anode  
solution

Cathode  
solution

Cathode  
material

# Requirements for Drawing/Labeling a Cell

- Determine what is reduced vs. oxidized ( $E^\circ_{\text{cell}}$  values, OR told which rxn to do)
- Beaker/container for anode and cathode AND liquid line drawn for each beaker
- Anode/cathode metal strips drawn submerged in liquid
- Label which solution/ions are in each beaker
- Label which beaker is anode and cathode
- Label anode/cathode strips with which solid metal each is
- Label anode and cathode with correct -/+ depending on if it is a galvanic cell (A-/C+) or electrolytic cell (A+/C-).
- Write the half reactions for each beaker
- Wire connecting anode/cathode strips together
- Label direction of e- flow along wire at top of drawing (anode  $\rightarrow$  cathode always)
- Salt bridge submerged in liquid on both sides
- Label direction of ion flow inside salt bridge (anions flowing to anode, cations flowing to cathode)



# Practice! Zn and Cu electrodes, ZnSO<sub>4</sub>, CuSO<sub>4</sub>

- Determine what is reduced vs. oxidized ( $E^\circ_{\text{cell}}$  values, OR told which rxn to do)
- Beaker/container for anode and cathode AND liquid line drawn for each beaker
- Anode/cathode metal strips drawn submerged in liquid
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**KCl salt bridge,  
Galvanic**



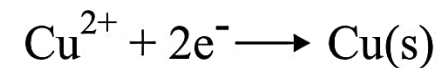
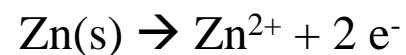
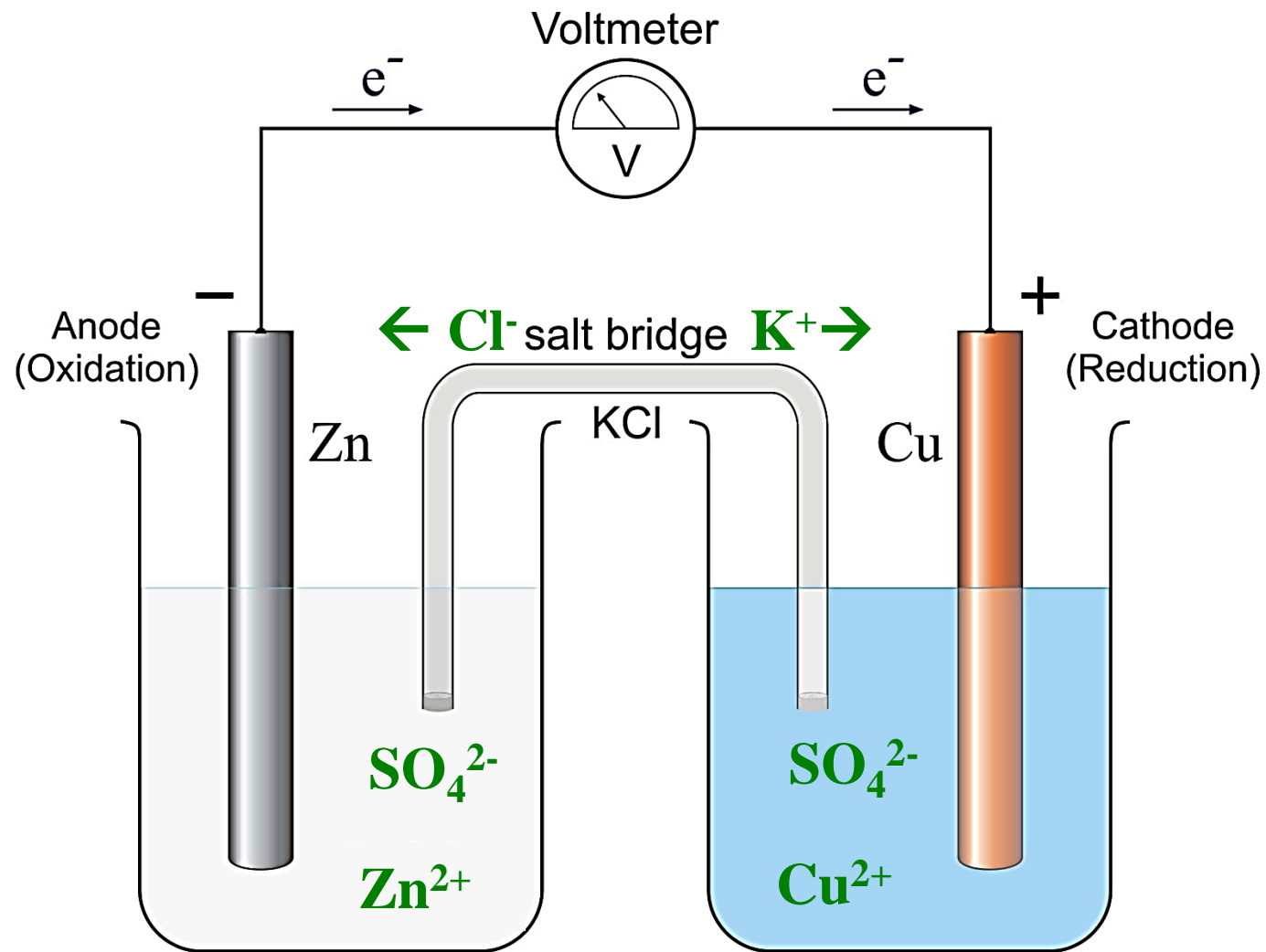
$$E^{\circ}_{\text{cell}} = -0.76$$

**Oxidized - Anode**



$$E^{\circ}_{\text{cell}} = 0.34$$

**Reduced - Cathode**



# YouTube Link to Presentation

<https://youtu.be/RRnWnKfYVDc>