### Today

Review for our Quiz!

Thermo and Electrochemistry

What happens when the conditions are not standard Nernst Equation What is the oxidation number of N in KNO<sub>3</sub>?

- A. (
- B. +
- C. -I
- D. +3
- E. +5

K is +1, O is -2

molecule is no charge

$$I(+1) + 3(-2) = -5$$

N must be +5

#### Balance this half reaction

$$HNO_2 \longrightarrow NO_3^-$$

### How many protons are in the balance 1/2 reaction?

(keeping the coeficient for  $NO_3^-$  as 1)

D. 2 on the left

$$HNO_2 \longrightarrow NO_3^-$$

$$HNO_2 + H_2O \longrightarrow NO_3^-$$

$$HNO_2 + H_2O \longrightarrow NO_3^- + 3H^+$$

$$HNO_2 + H_2O \longrightarrow NO_3^- + 3H^+ + 2e^-$$

Given following standard reduction potential, which do you think would make the best reducing agent?

$$Cl_2(g) + 2e^- \rightarrow 2Cl^- \qquad E^\circ = 1.36 V$$

$$I_2(s) + 2e^- \longrightarrow 2I^- \qquad E^\circ = 0.54 V$$

$$Br_2(I) + 2e^- \longrightarrow 2Br^- \qquad E^\circ = 1.06 V$$

A. CI- reducing agents are oxidized

B. Cl<sub>2</sub> hardest to reduce is easiest to oxidize

Lowest potential

C.  $I_2$ 

D. I Need to pick the reduced species (it can be oxidized)

E. Br<sub>2</sub>

You reduce  $H^+$  to  $H_2$  in an electrochemical cell. Your cell has a current of I Amp for 10 minutes What is the total charge that is passed through the cell?

- A. I C
- B. 10 C
- C. 600 C I A x (10 min) x (60 s min<sup>-1</sup>) = 600 C
- D. 6000 C

You reduce H<sup>+</sup> to H<sub>2</sub> in an electrochemical cell. Your cell has a current of I Amp for I0 minutes How many moles of electrons pass through the cell?



B. 600 C x F

C. IA x F

Therefore the number of moles of electrons is q/F

You reduce  $H^+$  to  $H_2$  in an electrochemical cell. The number of moles of electrons that pass through the cell is  $6.2 \times 10^{-3}$ . How many moles of  $H_2$  are formed?

$$2H^+ + 2e^- ----> H_2(g)$$

A. 
$$6.2 \times 10^{-3}$$

B. 
$$3.1 \times 10^{-3}$$

C.  $1.2 \times 10^{-2}$ 

For every mole of H<sub>2</sub> you need two moles of electrons

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# In the following standard cell, what is $E^{\circ}_{cell}$ ?

$$Zn(s) | Zn^{2+} || H^{+} | H_{2} (g)$$

$$Zn^{2+} + 2e^{-} ---> Zn(s)$$
  $E^{\circ} = -0.76 \text{ V}$   
  $2H^{+} + 2e^{-} ---> H_{2}$   $E^{\circ} = 0.0 \text{ V}$ 

anode on the left  $Zn \mid Zn^{2+}$  anode  $H^+ \mid H_2$  cathode

$$E^{\circ}_{cell} = E^{\circ}_{cathode} - E^{\circ}_{anode} = 0 - (-0.76) = +0.76$$

Voltaic Cell

In the following standard Ecell, what is the sign of the cathode?

$$Zn(s) | Zn^{2+} | | H^{+} | H_{2} (g)$$
  
 $Zn^{2+} + 2e^{-} ---> Zn(s) E^{\circ} = -0.76 V$   
 $2H^{+} + 2e^{-} ---> H_{2} E^{\circ} = 0.0 V$ 

A. 
$$+$$
  $E^{\circ}_{cell} = E^{\circ}_{cathode} - E^{\circ}_{anode} = 0 - (-0.76) = +0.76 V$ 

B.  $-$  Voltaic Cell therefore cathode +

C. neither 
$$E^{\circ}$$
cell = 0

# For a battery which of the following is correct?

A. 
$$E > 0$$
,  $\Delta G > 0$ ,  $K > I$ 

B. 
$$E > 0$$
,  $\Delta G < 0$ ,  $K > 1$ 

C. 
$$E > 0$$
,  $\Delta G < 0$ ,  $K < I$ 

D. 
$$E < 0$$
,  $\Delta G > 0$ ,  $K > I$ 

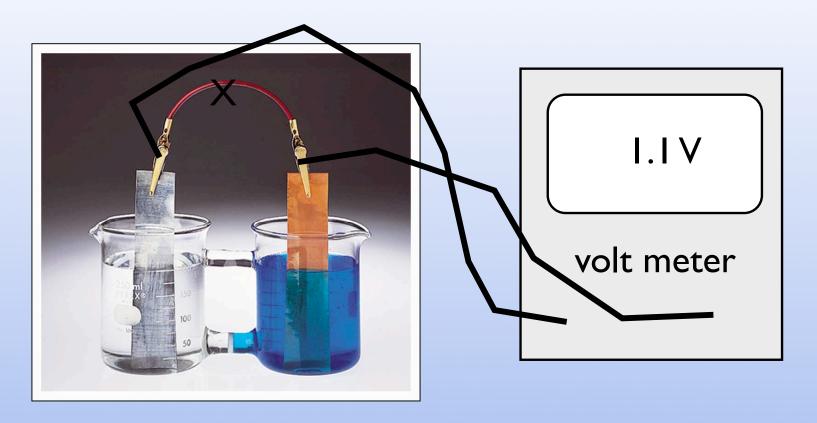
E. 
$$E < 0$$
,  $\Delta G < 0$ ,  $K > I$ 

F. 
$$E < 0$$
,  $\Delta G < 0$ ,  $K < I$ 

Battery = voltaic Spontaneous

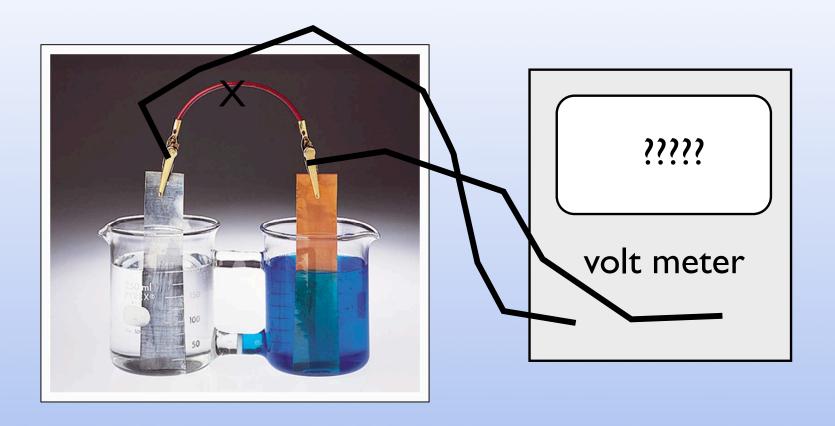
hint its not F.

#### We'll look at standard concentrations



I M Zn<sup>2+</sup> (aq) and I M Cu<sup>2+</sup> (aq) (note this is ridiculously concentrated)

#### What about other concentrations?



10<sup>-3</sup> M Zn<sup>2+</sup> (aq) and 10<sup>-1</sup> M Cu<sup>2+</sup> (aq) ???

# Relationship between E and $\Delta G$

 $\Delta G$  is energy E is electrical potential

Electric work (energy) is -charge x potential

work = 
$$-q \times E$$

$$\Delta G = work_{max}$$
  
 $\Delta G = -q \times E_{max}$ 

From now on well now the Potential we calculate are the theoretical maximum

Real world never actually that good

# Relationship between E and $\Delta G$

$$\Delta G = -q \times E$$

What is the charge q?

$$q = n \times F$$

n is number of moles of electrons F is the charge of one mole of electrons F = 96,485 C (Faraday's Constant)

$$\Delta G = - nFE$$

# Other concentrations and equilibrium Let's remember equlibrium!

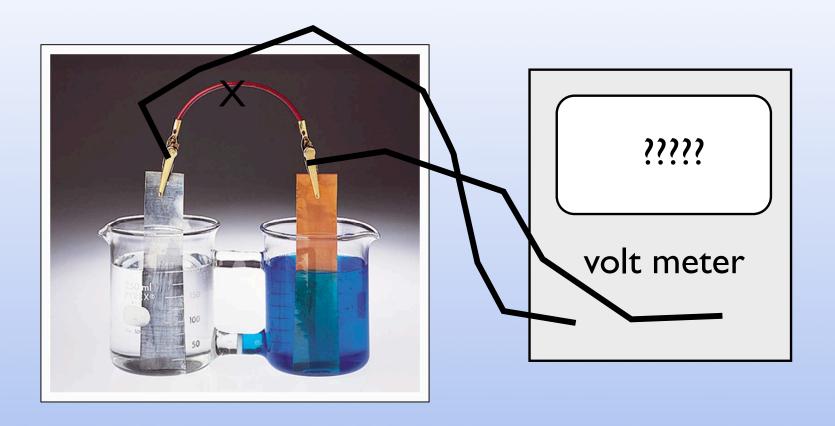
$$\Delta G = \Delta G^{\circ} + RTInQ$$
  
at equilibrium  $\Delta G = 0$   
so  $\Delta G^{\circ} = -RTInK$ 

$$-nFE = -nFE^{\circ} + RTInQ$$

$$E = E^{\circ} - \frac{RT}{nF} lnQ$$

assume 25°C
$$E = E^{\circ} - \frac{0.0591}{n} \log Q$$

#### What about other concentrations?



10<sup>-3</sup> M Zn<sup>2+</sup> (aq) and 10<sup>-1</sup> M Cu<sup>2+</sup> (aq) ???

I M Zn<sup>2+</sup> (aq) and I M Cu<sup>2+</sup> (aq) standard

$$Zn(s) + Cu^{2+}(aq) \longleftrightarrow Zn^{2+}(aq) + Cu(s)$$

$$Q = \frac{[Zn^{2+}]}{[Cu^{2+}]} = \frac{I}{I} = I$$

$$E = E^{\circ} - \frac{0.0591}{n} logQ$$

$$E = 1.10V - \frac{0.059I}{2} log(I) = 1.10V$$

$$Zn(s) + Cu^{2+}(aq) \longleftrightarrow Zn^{2+}(aq) + Cu(s)$$

$$Q = \frac{[Zn^{2+}]}{[Cu^{2+}]} = \frac{(10^{-3})}{(10^{-1})} = 10^{-2}$$

$$E = E^{\circ} - \frac{0.0591}{n} logQ$$

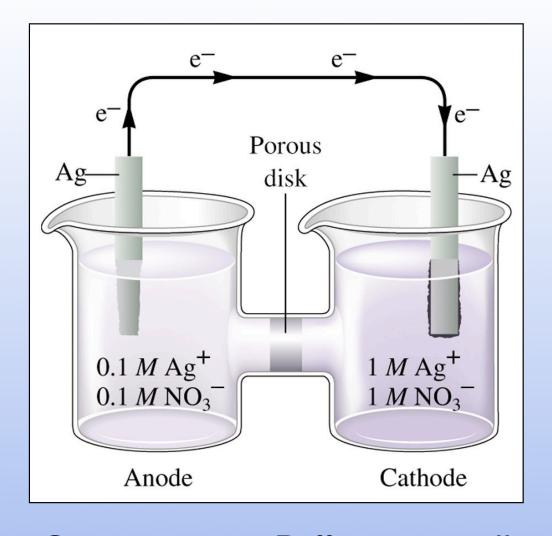
$$E = 1.10V - \frac{0.0591}{2} log(10^{-2}) = 1.16V$$

$$E = E^{\circ} - \frac{0.0591}{n} logQ$$

Current will flow until E = 0 Equilibrium

$$E^{\circ} = + \frac{0.0591}{n} \log K$$

$$logK = \frac{nE^{\circ}}{0.0591}$$



Concentration Differences will lead to potential difference

.I 
$$MAg^+$$
 (aq) and I  $MAg^+$  (aq)

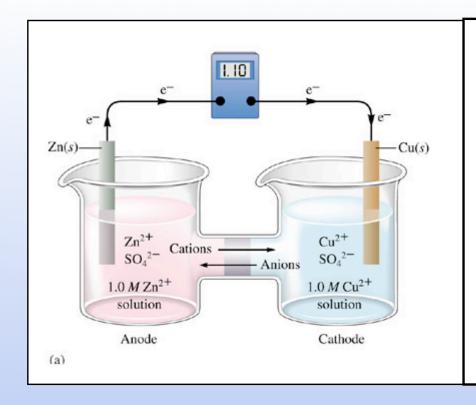
Same reaction!  $E^{\circ} = 0V$ 

$$Q = \frac{[Ag^+]_{anode}}{[Ag^+]_{cathode}} = \frac{.I}{I} = .I$$

$$E = E^{\circ} - \frac{0.0591}{n} logQ$$

$$E = 0V - \frac{0.0591}{I} log(.1) = 0.0591V$$

each factor of ten will be another 0.0591 V



If E < 0, then the reaction can be force in the non-spontaneous direction by applying a potential greater than E to the cell

$$q = I \times t$$