1. Consider the two reduction processes and their standard reduction potentials  $(E^{\circ})$ .

Cu<sup>+</sup> 
$$(aq)$$
 + 1 e<sup>-</sup>  $\rightarrow$  Cu  $(s)$   $E^{\circ}$  = +0.521 V  $(Ag^{+})(aq)$  + 1 e<sup>-</sup>  $\rightarrow$  Ag  $(s)$   $E^{\circ}$  = +0.800 V

- A) Circle  $\bigcirc$  the oxidizing agent and box  $\bigcirc$  the reducing agent. See above.
- B) Write the net ionic equation for a Galvanic/voltaic cell based on these reactions.

$$Ag^+(aq) + Cu(s) \rightleftharpoons Cu^+(aq) + Ag(s)$$

C) Determine the value of the  $E_{\text{cell}}^{\text{o}}$ 

$$E_{\text{cell}}^{\text{o}} = E_{\text{cathode}}^{\text{o}} - E_{\text{anode}}^{\text{o}} = 0.800 \text{ V} - 0.521 \text{ V} = 0.279 \text{ V}$$

D) Determine the value of the standard free energy change of the cell ( $\Delta G_{\text{cell}}^{\text{o}}$ ).

$$\Delta G_{\text{cell}}^{\text{o}} = -nFE_{\text{cell}}^{\text{o}} = -(1 \text{ mol } e^{-}) \left(96500 \frac{\text{C}}{\text{mol } e^{-}}\right) (0.279 \text{ V}) = -2.69 \times 10^{4} \text{ J}$$

E) Determine the equilibrium constant (K) for the reaction. Note: 1 J = 1 C·V

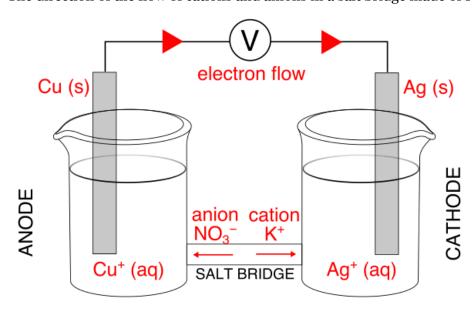
$$E_{\text{cell}} = E_{\text{cell}}^{\text{o}} - \frac{RT}{nF} \ln Q$$

$$\ln K = \frac{nF}{RT} E_{\text{cell}}^{\text{o}}$$

$$K = \exp \left\{ \frac{(1 \text{ mol } e^{-}) \left(96500 \frac{C}{\text{mol } e^{-}}\right)}{\left(8.314 \frac{J}{\text{mol } \cdot \text{K}}\right) (298.15 \text{ K})} \times 0.279 \text{ V} \right\}$$

$$K = 5.21 \times 10^{4}$$

- F) Given below is an unlabeled diagram. Label the following components in the diagram:
  - i. The solid electrodes on the anode and cathode sides.
  - ii. The ions in solutions on the anode and cathode sides.
  - iii. The direction of the flow of electrons through the voltmeter and wire.
  - iv. The direction of the flow of cations and anions in a salt bridge made of  $KNO_3$  (aq).



G) Write the cell diagram for this electrochemical cell.

2. You have constructed a Galvanic cell with the following reaction under standard conditions.

$$Zn(s) + Cu^{2+}(aq) \rightarrow Zn^{2+}(aq) + Cu(s)$$
  $E_{cell}^{o} = +1.104 \text{ V}$ 

What will the potential of the cell be when 0.50 M of Cu<sup>2+</sup> (aq) has reacted?

Assume that volume and temperature do not change.

Because the cell started under standard conditions, we know:

$$[Cu^{2+}]_i = 1.00 \text{ M}$$
  $[Zn^{2+}]_i = 1.00 \text{ M}$ 

Since 0.50 M of Cu<sup>2+</sup> is consumed after some time, we also know that 0.50 M must have been produced because the mole-mole ratio of Cu<sup>2+</sup> to Zn<sup>2+</sup> is 1:1. Therefore, the final concentrations are:

$$[Cu^{2+}]_{final} = 1.00 \text{ M} - 0.50 \text{ M} = 0.50 \text{ M}$$
  $[Zn^{2+}]_{final} = 1.00 \text{ M} + 0.50 \text{ M} = 1.50 \text{ M}$ 

Now use the Nernst equation to find the new (non-standard) cell potential:

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

$$= E_{\text{cell}}^{\text{o}} - \frac{RT}{nF} \ln \left( \frac{[\text{Zn}^{2+}]}{[\text{Cu}^{2+}]} \right)$$

$$= 1.104 \text{ V} - \frac{\left( 8.314 \frac{\text{J}}{\text{mol} \cdot \text{K}} \right) (298.15 \text{ K})}{(2 \text{ mol } e^{-}) \left( 96500 \frac{\text{C}}{\text{mol} e^{-}} \right)} \cdot \ln \left( \frac{1.50}{0.50} \right)$$

$$E_{\text{cell}} = 1.090 \text{ V}$$

3. Consider an electrochemical cell with the following cell diagram at 298.15 K.

$$Ni(s) | Ni^{2+}(1.25 M) | | Cu^{2+}(0.225 M) | Cu(s)$$

Given the following  $E^{\circ}$  values, determine whether each statement is true or false.

$${
m Cu^{2+}}\,(aq) + 2~{
m e^-} 
ightarrow {
m Cu}\,(s) \qquad E^\circ = +0.342~{
m V}$$
  ${
m Ni^{2+}}\,(aq) + 2~{
m e^-} 
ightarrow {
m Ni}\,(s) \qquad E^\circ = -0.257~{
m V}$   ${
m Ni}\,(s) + {
m Cu^{2+}}\,(aq) 
ightarrow {
m Cu}\,(s) + {
m Ni^{2+}}\,(aq) \qquad E^\circ_{\rm cell} = 0.599~{
m V}$ 

- A)  $E_{\text{cell}}$  is a smaller value than  $E_{\text{cell}}^{\text{o}}$ . <u>True</u>,  $E_{\text{cell}} = 0.578 \text{ V} < E_{\text{cell}}^{\circ}$ .
- B) The oxidation reaction takes place at the anode. <u>True</u>, oxidation always takes place at the anode.
- C) Doubling the volume of water in both half-cells will increase the cell potential. <u>False</u>, this will have no effect on the  $E_{cell}$  because the reaction quotient Q would not change.
- D) Decreasing the concentration of Ni<sup>2+</sup> will increase the cell potential. True, this will shift reaction to the right, thereby increasing the  $E_{cell}$  relative to 0.578 V.
- E) Increasing the concentration of  $Cu^{2+}$  will increase the cell potential. <u>True</u>, this will shift reaction to the right, thereby increasing the  $E_{cell}$  relative to 0.578 V.
- F) Using a Pt electrode in place of the Ni electrode will not change the cell potential. False, this will eliminate the concentration of Ni<sup>2+</sup> because Ni is the source of Ni<sup>2+</sup> ions.
- G) The mass of the Cu electrode will decrease over time.

  False, the mass of the Cu electrode (a product) increases over time.