

**Spring 2006 CH302, Practice Exam 6. Solution key answers in red.**

(the correct answer is included below. A solution key will be posted on Monday.)

1. Using the half reactions in the table of standard reduction potentials in Worksheet 11b, what is the standard emf (cell potential in V) for  $\text{Pt} | \text{Li} | \text{Li}^+(1\text{M}) || \text{F}_2 (1 \text{ atm}) | \text{F}^-(1\text{M}) | \text{Pt}$

1. +5.91 V **correct**
2. -5.91 V
3. -0.17 V
4. 0.17 V

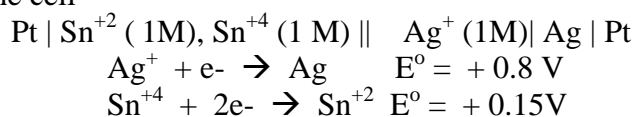
**Solution:** All compounds are standard state so simple use  $E^\circ_{\text{cell}} = E^\circ_{\text{cathode}} - E^\circ_{\text{anode}} = 2.87 - (-3.04) = 5.91 \text{ V}$ . The nearly +6V spontaneous reaction is the largest cell reaction you can make with the half reactions found in the usual table of standard reduction potentials.

2. Consider the half-reactions in the table of standard reduction potentials in Worksheet 11b. Of the species listed, the strongest reducing agent is ? and the weakest oxidizing agent is ?

1. Li,  $\text{Li}^+$  **correct**
2. Li,  $\text{F}_2$
3.  $\text{Li}^+$ ,  $\text{F}_2$
4. F,  $\text{F}_2$

**Solution:** From the table of standard reduction potentials, we want the strongest reducing agent which means we want the oxidation process that is the most positive--that is the Li oxidation in the upper right hand corner of the table (+3.04V). Next we want to find the weakest oxidizing agent which means we want the reduction half reaction compound with the most negative  $E^\circ$ ,  $\text{Li}^+$ , in the upper left hand corner (-3.04V).

3. Consider the cell



at standard conditions. Calculate the value of  $\Delta G$  for the reaction that occurs when current is drawn from this cell.

1. 125 kJ/mol
2. -125 kJ/mol **correct**
3. 62 kJ/mol
4. -62 kJ/mol

**Solution:** This is a multi-step problem.

First we need to find  $E^\circ_{\text{cell}}$ . All the compounds are standard state so we simply use  $E^\circ_{\text{cell}} = E^\circ_{\text{cathode}} - E^\circ_{\text{anode}} = 0.8\text{V} - 0.15\text{V} = 0.65 \text{ V}$  where the silver electrode is cathode and tin electrode is anode.

Also note in balancing the reaction that the least common multiple is 2 which means this is an n=2 electron process.

Finally, we find  $\Delta G = -nFE = -(2)(9.65 \times 10^4)(0.65\text{V}) = 125,000\text{J} \rightarrow -125 \text{ kJ}$ .

Note the  $E^\circ_{\text{cell}}$  is positive so free energy is negative, a spontaneous reaction.

4. An electrochemical process with a positive  $\Delta G$  value has two terminals labeled positive and negative. In this cell, electrons would flow from the ? terminal to the ? terminal through the external circuit and ? reaction occurs at the positive terminal.

1. positive; negative; a reduction
2. positive; negative; an oxidation **correct**
3. negative; positive; a reduction
4. negative; positive; an oxidation
5. positive; negative; an acid/base

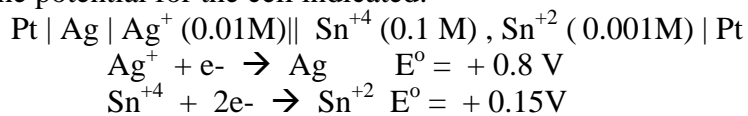
**Solution:** Because the reaction is non-spontaneous (an electrolysis cell), by definition the anode is positive and the cathode is negative. Also by definition, electrons flow from the anode to cathode so the flow is from positive electrode to negative electrode. Finally, the positive terminal is the anode in electrolysis, so oxidation is occurring there.

5. Which of the following batteries is rechargeable?

1. dry cell
2. copper/zinc
3. nickel-cadmium battery **correct**
4. copper/carrot top

**Solution.** The famous rechargeable batteries mentioned in class are the lead acid storage battery found in your car and the nickel cadmium battery which is the kind of rechargeable battery you find in stores in standard sizes like AA and D.

6. Calculate the potential for the cell indicated:



1. -0.65V
2. +0.65 V
3. -0.47 V **correct**
4. - 0.53 V
5. -0.29 V

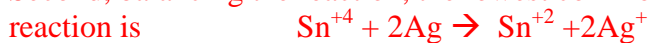
**Solution.** The potentials in this reaction are nonstandard so we use the Nernst eqn

$$E_{\text{cell}} = E^\circ_{\text{cell}} - .059/n \log Q$$

**Step by step**

First:  $E^\circ_{\text{cell}} = E^\circ_{\text{cathode}} - E^\circ_{\text{anode}} = 0.15 - 0.8 \text{ V} = -0.65\text{V}$

Second, balancing the reaction, the lowest common multiple is 2, so this is a 2 electron process. The balanced reaction is



so  $Q = (\text{Sn}^{+2})(\text{Ag}^+)^2 / (\text{Sn}^{+4})$  (we don't include solid silver because it has unit activity).

Substituting the given concentrations, and remembering to square the  $\text{Ag}^+$  we wind up with  $Q = 10^{-6}$ .

$$\text{So } E_{\text{cell}} = E^\circ_{\text{cell}} - .059/n \log Q = -.65 - .059/2 \log 10^{-6} = -0.47 \text{ V}$$

7. What weight of Au is deposited on a piece of cheap zinc chain during the electrolysis of a  $\text{Au}^{+3}$  solution when a current of 1.3 mA amps flows for 1.2 hours? (Au = 197 g/mol)

1. 3.82 mg **correct**
2. 1.27 mg
3. 11.46 mg
4. 1.44 g
5. 43.2 g

**Solution:** This is a classic stoichiometry problem where we use an initial amount of charge instead of an initial mass. Otherwise the problem is the same.

To find the charge, in Coulombs, remember that current,  $i = \text{Coulombs/second}$ .

So Coulombs =  $i$  (in Amps) \* seconds =  $1.3 \times 10^{-3} \text{A} * 1.2 \text{ hours (3600 seconds/hour)} = 5.616 \text{ Coulombs}$ .

Also note this is a three electron process because one mole of gold forms when one mole of  $\text{Au}^{+3}$  is reduced by three moles of electrons.

So:  $\text{g of Au} = 5.616 \text{ Coulombs} (1 \text{ mole } e^{-} / 9.65 \times 10^4 \text{ Coulombs}) (1 \text{ mole Au} / 3 \text{ moles } e^{-}) (197 \text{ g Au} / 1 \text{ mole Au}) = 3.82 \times 10^{-3} \text{ grams}$ .

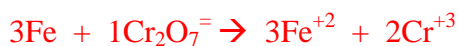
8 Using the smallest possible integer coefficients to balance the redox equation

$\text{Fe} + \text{Cr}_2\text{O}_7^{-} \rightarrow \text{Fe}^{+2} + \text{Cr}^{+3}$  in acidic solution, the coefficient for  $\text{Cr}_2\text{O}_7^{-}$  is

1. 1 **correct**
2. 2
3. 3
4. 4
5. The correct coefficient is not given.

**Solution.**

This is a six electrons process, when the least common factor 6, is multiplied through the reaction.



Note that to answer this problem you don't need to finish balancing the reaction—you only need to know the coefficient of the dichromate.