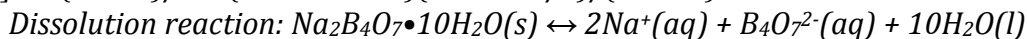


You must show your work for full credit.

Exp 9 and 10 Questions

1. A 5.0 mL aliquot of a saturated $\text{Na}_2\text{B}_4\text{O}_7 \cdot 10\text{H}_2\text{O}$ solution is titrated and gives 0.00134 moles of $\text{B}_4\text{O}_7^{2-}$ present. What is the K value for dissolution? (2 pts)

$$[\text{B}_4\text{O}_7^{2-}] = n(\text{B}_4\text{O}_7^{2-})/V = (0.00134 \text{ mol})(1000 \text{ mL/L})/(5.0 \text{ mL}) = 0.268 \text{ M}$$



ICE Table

I	0	0
C	+2x	+x
E	2x	x

Since $x = [\text{B}_4\text{O}_7^{2-}] = 0.268 \text{ M}$, $[\text{Na}^+] = 2x = 0.536 \text{ M}$ and $K = [\text{Na}^+]^2[\text{B}_4\text{O}_7^{2-}] = (0.536)^2(0.268)$

K = 0.077 (only 2 sig fig based on the volume) Can give partial credit if parts done correctly

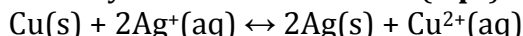
2. When $\ln K$ for dissolution of $\text{Na}_2\text{B}_4\text{O}_7 \cdot 10\text{H}_2\text{O}$ in water is plotted vs. $1/T$, a negative slope is seen. Using the equation below, explain whether dissolution of $\text{Na}_2\text{B}_4\text{O}_7 \cdot 10\text{H}_2\text{O}$ would be expected to result in the solution becoming warmer or cooler. Explain your answer. (Hint: think about how ΔH relates to temperature change from a reaction) (1 pt)

$$\ln K = \frac{-\Delta H^\circ}{RT} + \frac{\Delta S^\circ}{R}$$

Since it is reported that the $\ln K$ vs. $1/T$ plot gives a negative slope, this means $-\Delta H^\circ/R$ is negative.

Since R is positive, ΔH° must also be positive. When a reaction has a positive ΔH , the reaction takes heat, meaning that the **solution will get cooler.**

3. Given the following net equation, write the anode and cathode half reactions. Be sure to include state symbols and electrons. (2 pt)



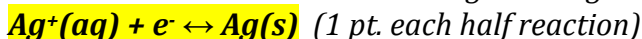
anode half reaction:

oxidation occurs at the anode. Cu is being oxidized. Anode half reaction is:



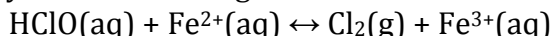
cathode half reaction:

reduction occurs at the cathode. Ag^+ is being reduced. Cathode half reaction is:



Electrochemistry problems

4. Adding H^+ and H_2O where necessary, balance the following redox reaction. To receive full credit, you need to also give two balanced half reactions. (3 pts)



reduction: Cl (from -1 to 0) balanced half-reaction: $2\text{HClO}(aq) + 2\text{H}^+(aq) + 2e^- \leftrightarrow \text{Cl}_2(g) + 2\text{H}_2\text{O}(l)$

oxidation: $\text{Fe}^{2+}(aq) \leftrightarrow \text{Fe}^{3+}(aq) + e^-$

Combine (first multiply Fe reaction by 2 so that number of e's is the same):



(one more on back) (give +1 pt for each part done correctly)

5. Given the following electrochemical reduction potential table, come up with a pair of reactions that will give the greatest potential under standard conditions and calculate the potential under standard conditions. (2 pts)

Reaction	$E^\circ(\text{V})$
$\text{Fe}^{3+}(\text{aq}) + \text{e}^- \leftrightarrow \text{Fe}^{2+}(\text{aq})$	+0.771
$\text{Cu}^{2+}(\text{aq}) + 2\text{e}^- \leftrightarrow \text{Cu}(\text{s})$	+0.337
$2\text{H}^+(\text{aq}) + 2\text{e}^- \leftrightarrow \text{H}_2(\text{g})$	0.00
$\text{Pb}^{2+}(\text{aq}) + 2\text{e}^- \leftrightarrow \text{Pb}(\text{s})$	-0.126

The greatest potential will occur when the reaction with the highest reduction potential is coupled to the reaction with the lowest reduction potential (**so this is $\text{Fe}^{3+}(\text{aq}) + \text{e}^- \leftrightarrow \text{Fe}^{2+}(\text{aq})$ for the reduction and $\text{Pb}(\text{s}) \leftrightarrow \text{Pb}^{2+}(\text{aq}) + 2\text{e}^-$ for oxidation**). $E_{\text{cell}}^\circ = 0.771 \text{ V} - (-0.126 \text{ V}) = \mathbf{0.897 \text{ V}}$ (1 pt for each part)