Understanding the Theory Behind the Ka Lab

- The purpose of the lab is to gather enough data so that we could calculate the K_A of the acid CH₃CO₂H.
- Remember for $CH_3CO_2H_{(aq)} = CH_3CO_2^{-}_{(aq)} + H^{+}_{(aq)}$ $K_A = \frac{[CH_3 CO_2^{-}][H^{+}]}{[CH_3 CO_2H]}$
- So we need to get the equilibrium [H⁺], [CH₃CO₂⁻] and [CH₃CO₂H]
- We can get the first two from the pH.
- Think of the ICE chart and you'll realize why [H⁺] & [CH₃CO₂⁻] are equal
- To get [CH₃CO₂H], we could do a titration with NaOH and see how much of NaOH is needed to neutralize the CH₃CO₂H:

$$NaOH + CH_3CO_2H \rightarrow NaCH_3CO_2 + H_2O$$

We let

 $n_1 = moles of NaOH = C_1V_1$ $n_2 = moles of CH_3CO_2H = C_2V_2$

Since the base and weak acid react in a 1:1 ratio, then $n_1 = n_2$, so:

 $\mathbf{C}_1 \mathbf{V}_1 = \mathbf{C}_2 \mathbf{V}_2$

If we measure the volume of a known concentration of NaOH needed to neutralize a known volume of CH_3CO_2H , the only unknown in the formula will be C2, the last concentration we are looking for.

Term 3 LAB EXAM STUDY GUIDE

Let's assume that an equilibrium was created by combining 0.010 L of 1.0 mole/L Fe⁺³ and 0.010 L of 1.0 mole/L SCN⁻ :

 $\begin{array}{rcl} Fe^{+3} & + & SCN^{-} & = & FeSCN^{+2} \\ Yellow-orange & colorless & DEEP RED \end{array}$

- a) Do the calculations to make sure that adding 0.0050 L of $1.0 \text{ mole/L Fe}^{+3}$ to the above will indeed increase the concentration of Fe}^{+3} and subsequently shift the equilibrium to the right.
- b) Will adding 0.40 grams of SCN⁻ be more effective in disturbing the original equilibrium? Show why or why not.

<u>Answer</u>

a) Original
$$[Fe^{+3}] = \frac{n}{V_{total}} = \frac{CV}{V_{total}} = \frac{1.0moles/L(0.010 L)}{0.010 + 0.010 L} = 0.50 \text{ moles } Fe^{+3}/L$$

Adding 0.005 L of 1.0 mole/L Fe^{+3} is adding:

 $n = CV = 1.0 \text{ mole/L} (0.005 \text{ L}) = 0.005 \text{ moles of Fe}^{+3}$ while increasing the volume by 0.005 L:

volume by 0.005 L: new [Fe⁺³] = $\frac{n_{total}}{V_{total}}$ = $\frac{new n + CV}{V_{total}}$ = $\frac{0.005 \text{ moles } + 1.0 \text{ moles}/L(0.010 \text{ L})}{0.010 + 0.010 \text{ L} + 0.005 \text{ L}}$ = 0.60 moles Fe⁺³/L

So indeed, the concentration has gone up, and it will create more effective collisions with SCN-, increasing the forward rate and driving the reaction towards more red FeSCN⁺².

b) Original $[SCN^{-}] = \frac{n}{V_{total}} = \frac{CV}{V_{total}} = \frac{1.0moles/L(0.010 L)}{0.010 + 0.010 L} = 0.50 \text{ moles SCN}^{-}/L$ Adding 0.40 grams of SCN⁻ is adding:

 $m/M = n = 0.40 \text{ grams of SCN}^{-} / (32 + 12 + 14 \text{ g/mole}) = 0.00690 \text{ moles}$ $new [SCN^{-}] = \frac{n_{total}}{V_{total}} = \frac{new n + CV}{V_{total}} = \frac{0.00690 \text{ moles} + 1.0 \text{ moles}/L(0.010 \text{ L})}{0.010 + 0.010 \text{ L}} = 0.85 \text{ moles}/L$

So the concentration goes up more dramatically it will be more effective in disturbing the equilibrium, so it should go a deeper red. (Notice that by adding a dissolving solid we don't change the volume appreciably, so we assume that it's still the same as the original created by adding the two solutions)