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## Part A. Determination of an Unknown Concentration

Q1. Record the concentration of your red solution: red solution = 12.0 ppm

Q2. (1 pt) Record the absorbance (measured at 480 nm) of your six standard solutions:

Standard	1	2	3	4	5	6
% Red	100%	80%	60%	40%	20%	0
% Blue	0	20%	40%	60%	80%	100%
Absorbance	0.224	0.179	0.135	0.089	0.045	0.0

Q3. Record the absorbance of the two unknown solutions:

Absorbance of Unknown #1 = 0.0  
0.241

Absorbance of Unknown #2 =

Q4. (1 pt) Use Excel, Google sheets or a similar program to create a graph of Absorbance vs. Concentration of Red Solution for your six *standard* solutions, where your graph must include a linear trendline as well a title and labeled axes. Add a blank page to the end of this lab report, then paste your graph onto that page, and record the equation of your trend line below in standard  $y = mx + b$  format:Equation:  $y = 0.0187x$ 

Q5. (2 pt) Calculate the % composition of red solution in each of your two unknown solutions then use the original concentration of red solution, recorded in Q1, to determine the parts-per-million concentration. Show an example of your work below:

Solution	Unknown 1	Unknown 2
Percentage Red (%)	0.0	107%
Concentration Red (ppm)	0.0	12.9

$$y = 0.0187(x)$$

y = absorbance  
x = concentration (ppm)

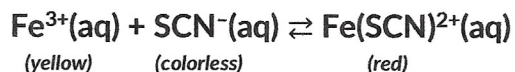
$$\therefore 0.0 = 0.0187x \rightarrow x = 0.0 \text{ ppm}$$

$$0.241 = 0.0187(x) \rightarrow x = \frac{0.241}{0.0187} = 12.888 \text{ ppm}$$

$$\frac{12.9 \text{ ppm}}{12.0 \text{ ppm}} \times 100\% = 107\%$$

## Part B. Determination of an Equilibrium Constant

The following questions refer to the reaction shown below, where, unfortunately you will not have the opportunity to conduct this portion of the experiment this summer:



- Q6. (2 pt) Five non-equilibrium standard solutions of the reaction shown above are prepared as described on the following table. Calculate the concentration of  $\text{SCN}^{-}$  in each standard once all reagents have been mixed together into a total volume of 25 mL; show at least one sample calculation in the space below:

Standard	1	2	3	4	5
Volume of 0.00150 M $\text{Fe}(\text{NO}_3)_3$	5.0 mL	5.0 mL	5.0 mL	5.0 mL	5.0 mL
Volume of 0.00150 M KSCN	5.0 mL	4.0 mL	3.0 mL	2.0 mL	1.0 mL
Volume of 0.050 M $\text{HNO}_3$	15 mL	16 mL	17 mL	18 mL	19 mL
Concentration of $\text{SCN}^{-}$ in 25 mL	$3.0 \times 10^{-4} \text{ M}$	$2.4 \times 10^{-4} \text{ M}$	$1.8 \times 10^{-4} \text{ M}$	$1.2 \times 10^{-4} \text{ M}$	$6.0 \times 10^{-5} \text{ M}$

~~$$M_{\text{KSCN}} = M_{\text{SCN}^{-}} = 0.00150 \text{ M}$$~~

$$n_{\text{KSCN}} = n_{\text{SCN}^{-}} = 0.00150 \text{ M} \times \frac{0.005 \text{ L}}{1} = 7.5 \times 10^{-6} \text{ mol}$$

\* Nitric acid is used to prevent the hydrolysis of iron (III)

$$M_{\text{SCN}^{-}} = \frac{7.5 \times 10^{-6} \text{ mol}}{0.025 \text{ L}} = 3.0 \times 10^{-4}$$

$$n_{\text{Fe}^{3+}} = 0.00150 \text{ M} \times 0.005 \text{ L} = n_{\text{SCN}^{-}}$$

- Q7. If the final concentration of  $\text{Fe}(\text{SCN})^{2+}$  is equal to the initial concentration of  $\text{SCN}^{-}$  calculated in Q6, complete the following table:

Standard	1	2	3	4	5
$[\text{Fe}(\text{SCN})^{2+}]$	$3.0 \times 10^{-4} \text{ M}$	$2.4 \times 10^{-4} \text{ M}$	$1.8 \times 10^{-4} \text{ M}$	$1.2 \times 10^{-4} \text{ M}$	$6.0 \times 10^{-5} \text{ M}$
Absorbance @ 460 nm	0.449	0.364	0.279	0.189	0.097

- Q8. (1 pt) Create a curve of Absorbance vs. Concentration of  $\text{Fe}(\text{SCN})^{2+}$  for your five standard solutions, where your graph must include a linear trendline, a proper title and labeled axes; paste this graph onto the page you added to the end of this lab report beneath your graph from Part A (created in Q4). Record the equation of your trend line in  $y = mx + b$  format:

Equation:  $y = 1465x + 0.0119$

- Q9. (2 pt) Two equilibrium mixtures of the iron / thiosulfate reaction are prepared, where  $\text{HNO}_3$  was added to bring the volume of each mixture to a total of 10.0 mL. Calculate the concentrations of  $\text{Fe}^{3+}(\text{aq})$  and  $\text{SCN}^{-}(\text{aq})$  in the 10-mL mixtures, showing at least one sample calculation of  $[\text{Fe}^{3+}]_0$  and  $[\text{SCN}^{-}]_0$  in the space below:

Mixture	A	B
Volume of 0.00150 M $\text{Fe}(\text{NO}_3)_3$	5.0 mL	5.0 mL
Volume of 0.00150 M KSCN	2.0 mL	3.0 mL
Concentration of $\text{Fe}^{3+}$ in 10 mL, $[\text{Fe}^{3+}]_0$	$7.5 \times 10^{-4} \text{ M}$	$7.5 \times 10^{-4} \text{ M}$
Concentration of $\text{SCN}^{-}$ in 10 mL, $[\text{SCN}^{-}]_0$	$3.0 \times 10^{-4} \text{ M}$	$4.5 \times 10^{-4} \text{ M}$

$$n_{\text{Fe}^{3+}} = n_{\text{Fe}(\text{NO}_3)_3} = 0.0015 \text{ M} \cdot 0.005 \text{ L} = 7.5 \times 10^{-6} \text{ mol} \rightarrow M_{\text{Fe}^{3+}} = \frac{7.5 \times 10^{-6}}{0.010 \text{ L}} = 7.5 \times 10^{-4} \text{ M}$$

$$n_{\text{SCN}^{-}} = n_{\text{KSCN}} = 0.00150 \text{ M} \cdot 0.002 \text{ L} = 3.0 \times 10^{-6} \text{ mol} \rightarrow M_{\text{SCN}^{-}} = \frac{3.0 \times 10^{-6} \text{ mol}}{0.010 \text{ L}} = 3.0 \times 10^{-4} \text{ M}$$

- Q10. (1 pt) Use your standard curve to calculate the equilibrium concentration of  $\text{Fe}(\text{SCN})^{2+}$  in both of your mixtures; show at least one sample calculation in the space below:

Mixture	A	B
Absorbance @ 460 nm	0.218	0.301
$[\text{Fe}(\text{SCN})^{2+}]_{\text{eq}}$	$1.4 \times 10^{-4} \text{ M}$	$2.0 \times 10^{-4} \text{ M}$

$$y = 1465x + 0.0119 \quad \left. \begin{array}{l} y = \text{absorbance} \\ x = \text{concentration} \end{array} \right\} 0.218 = 1465x + 0.0119 \rightarrow x = \frac{0.218 - 0.0119}{1465}$$



Q11. (3 pt) Set up ICE tables for Mixtures A and B, then determine the value of K for each mixture, as well as your average K value:

A)

	$\text{Fe}^{3+}$	$\text{SCN}^-$	$\text{Fe}(\text{SCN})^{2+}$
i	$8.9 \times 10^{-4}$	$4.4 \times 10^{-4}$	0
c	-x	-x	x
e	$7.5 \times 10^{-4}$	$3 \times 10^{-4}$	$1.4 \times 10^{-4}$

$x = 1.4 \times 10^{-4}$

$$K_A = \frac{1.4 \times 10^{-4}}{(7.5 \times 10^{-4})(3 \times 10^{-4})} = 6.2 \times 10^2$$

B)

	$\text{Fe}^{3+}$	$\text{SCN}^-$	$\text{Fe}(\text{SCN})^{2+}$
i	$9.5 \times 10^{-4}$	$6.5 \times 10^{-4}$	0
c	-x	-x	x
e	$7.5 \times 10^{-4}$	$4.5 \times 10^{-4}$	$2.0 \times 10^{-4}$

$x = 2.0 \times 10^{-4}$

$$K_B = \frac{2.0 \times 10^{-4}}{(7.5 \times 10^{-4})(4.5 \times 10^{-4})} = 5.9 \times 10^2$$

$$K_{\text{avg}} = 6.1 \times 10^2$$

Q12. (1 pt) If your classmate prepares an iron / thiosulfate mixture using different initial concentrations than you, would their K value be significantly different than yours? Explain.

No because K is a ratio of Products / reactants @ equilibrium.

no matter how much you start with, that ratio should be similar because the

Q13. (1 pt) If your classmate prepares an iron / thiosulfate mixture using the same initial concentrations as you but runs their reaction at a different temperature, could their K value be significantly different than yours? Explain.

Yes — At a higher temperature, an exothermic reaction would favor the forward reaction and K would be greater.

