**PI:** Michael Evans **Chemistry Science Education Title:** Spectrophotometric Determination of an Equilibrium Constant

**Overview**The equilibrium constant *K* for a chemical system is the ratio of product concentrations to reactant concentrations at equilibrium, each raised to the power of their respective stoichiometric coefficients. Measurement of *K* involves determination of these concentrations for systems in chemical equilibrium.

Reaction systems containing a single colored component can be studied spectrophotometrically. The relation between absorbance and concentration for the colored component is measured and used to determine its concentration in the reaction system of interest. Concentrations of the colorless components can be calculated indirectly using the balanced chemical equation and the measured concentration of the colored component.

In this video, the Beer’s law curve for Fe(SCN)2+ are determined empirically and applied to the measurement of *K* for the following reaction:



Four reaction systems with different initial concentrations of reactants are investigated to illustrate that *K* remains constant irrespective of initial concentrations.

**Principles**  
Every chemical reaction is associated with an equilibrium constant *K*, which reflects the ratio of product concentrations to reactant concentrations at chemical equilibrium. For the generic reaction *a* A + *b* B ⇄ *c* C + *d* D, the equilibrium constant is defined as



where the concentrations on the right-hand side of the equation are molarities at equilibrium. This equation is known as the equilibrium expression for the reaction. In chemical systems that are not in equilibrium, the forward and reverse reactions occur at different rates until the concentrations of reactants and products satisfy the equilibrium expression.

To measure the value of *K* for a system in chemical equilibrium, it is necessary to determine the concentrations of reactants and products either directly or indirectly. Spectrophotometric methods for the determination of *K* involve direct measurement of the concentration of one colored component and indirect measurement of the others. Visible spectroscopy of the colored component at known concentrations reveals the relation between absorbance and concentration for this component. According to Beer’s law, this relation has the form

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where  is the molar absorptivity of the component in L/mol-cm, *l* is the path length of light through the sample in cm, *c* is the molarity of the component in mol/L, and *A* is the absorbance.

The Beer’s law curve for the colored component can be applied to a reaction system in equilibrium to determine the concentration of this component from absorbance measurements. Concentrations of the remaining reactants and products can then be calculated by adjusting initial concentrations based on the measured molarity of the colored species.

The system studied here is the reaction of iron (III) cation with thiocyanate anion to form an iron (III) thiocyanate complex.



The equilibrium expression for this reaction system is



where the *eq* subscript denotes equilibrium concentrations. The iron (III) thiocyanate product is orange, but both reactants are colorless in aqueous solution. Hence, [Fe(SCN)2+]*eq* can be determined directly from absorbance measurements.



Equilibrium concentrations of the reactants can be calculated by subtracting the equilibrium concentration of the product from the initial concentrations of the reactants. An Initial-Change-Equilibrium (ICE) table illustrates how initial and equilibrium concentrations are related (**Table 1**).



Repeated trials of this experiment involving different initial concentrations of reactants should yield the same value of *K*, as the value of *K* is independent of concentration.

**Procedure**

1. Determining the Beer’s Law Curve for Fe(SCN)2+
   1. Calibrate a visible spectrophotometer using distilled water.
   2. Add 1.0 mL of 1.0 × 10–4 m Fe(NO3)3 solution to a medium test tube.
   3. To the same test tube, add 5.0 mL of 0.50 m KSCN solution.
   4. To the same test tube, add 4.0 mL of 0.10 m HNO3 solution, and mix the solution well.
   5. Use a Pasteur pipette to transfer a small quantity of the solution to a cuvette. Ensure that the liquid level is above the path of the light beam in the spectrophotometer.
   6. Place the cuvette in the spectrophotometer, so the light passes through the smooth sides.
   7. Acquire a spectrum and record the max value and the absorbance at max.
   8. Repeat steps 1.2 – 1.7 using the volumes of Fe(NO3)3, KSCN, and HNO3 solutions in **Table 2**.
   9. Plot the measured absorbance versus the concentration of Fe(SCN)2+ in each test tube and determine the line of best fit for the data. The slope of this line is the molar absorptivity  and the path length is 1 cm.
2. Measuring *K* for the Iron (III) Thiocyanate System.  
   1. Prepare four medium test tubes containing the indicated volumes of 0.0025 m Fe(NO3)3, 0.0025 m KSCN, and 0.10 m HNO3 solutions in **Table 3**.
   2. Mix each solution well and allow them to stand for at least 10 min. This resting period ensures that the solutions are at chemical equilibrium.
   3. Use a Pasteur pipette to transfer a small quantity of solution 6 to a cuvette. Ensure that the liquid level is above the path of the light beam in the spectrophotometer.
   4. Acquire a spectrum and record the max value and the absorbance at max.
   5. Repeat steps 2.3 and 2.4 for solutions 7 – 9.

**Representative Results  
Table 4** lists the absorbance and max data for solutions 1 – 5. Concentrations of Fe(SCN)2+ were determined from initial concentrations of Fe3+ under the assumption that all of the Fe3+ is converted to Fe(SCN)2+. A large excess of SCN– was used in tubes 1 – 5 to ensure that this assumption holds true.

The molarity [Fe(SCN)2+] and absorbance are plotted in **Figure 2**. The measured absorbance agrees well with Beer’s law.

**Table 5** lists measured absorbance and calculated *K* values for tubes 6 – 9. Equilibrium concentrations of Fe3+ and SCN– were calculated by subtracting the measured concentration of Fe(SCN)2+ from the initial concentrations of Fe3+ and SCN–.

The mean *K* value was 147 ± 11, illustrating that *K* is roughly constant over the range of concentrations studied.

**Applications**The equilibrium constant provides useful information about the extent to which a reaction will proceed to form products over time. Reactions with a large value of *K*, much larger than 1, will form products nearly complete given enough time (**Figure 3**). Reactions with a value of *K* less than 1 will not proceed forward to a significant degree. The equilibrium constant thus serves as a measure of the feasibility of a chemical reaction.

The equilibrium constant also provides useful thermodynamic information about the changes in free energy, enthalpy, and entropy in the course of a chemical reaction. The equilibrium constant is related to the free energy change of reaction:

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The free energy change of reaction is in turn related to the enthalpy and entropy changes of reaction:



Measurements of the temperature dependence of *K* can reveal the enthalpy change *H* and the entropy change *S* for a reaction. In addition to providing chemists with insight into patterns in molecular behavior, tables of thermodynamic data can be used to identify reactions with favorable thermodynamic properties. For example, redox reactions that release large amounts of energy (associated with negative *G* values) are attractive candidates for batteries (**Figure 4**).

**Legend**

Figure 1: Beer’s law curve for iron (III) thiocyanate.

Figure 2: Line graph of Absorbance versus Concentration for Fe(SCN)2+.

Figure 3: The equilibrium constant of this reaction is greater than 1. A significant amount of colored product forms in each case, even though the initial concentrations of reactants differ.

Figure 4: The redox reaction of copper (II) cations in aqueous solution with aluminum metal is a strongly exergonic reaction with a large value of K.

Table 1: An Initial-Change-Equilibrium (ICE) table that illustrates how initial and equilibrium concentrations are related.

Table 2: Appropriate volumes of Fe(NO3)3, KSCN, and HNO3 solutions to be placed in tubes 2 – 5.

Table 3: Appropriate volumes of 0.0025 M Fe(NO3)3, 0.0025 M KSCN, and 0.10 M HNO3 solutions.

Table 4: Absorbance versus Concentration Data for Fe(SCN)2+.

Table 5: Measured absorbance values and calculated K for the reaction of iron (III) with thiocyanate.