## Materials and Equipment

	Materials	Equipment	
	Prepared FeNCS <sup>2+</sup> Solutions	3 - 50mL Pipette	
	$2.00 \times 10^{-5} M FeNCS^{2+}$	8 - 3mL Cuvettes	
	$4.00 \times 10^{-5} M FeNCS^{2+}$	Lint-free tissue paper	
	$6.00 \times 10^{-5} M \text{ FeNCS}^{2+}$	Colorimeter	
	$8.00 \times 10^{-5} M FeNCS^{2+}$	6 15mL Test Tubes	
	$1.00 \times 10^{-4} M \text{ FeNCS}^{2+}$		
	$1.20 \times 10^{-4} M \text{ FeNCS}^{2+}$	- 1K	
	$1.40 \times 10^{-4} M \text{ FeNCS}^{2+}$	-ale.co.un	
	2.00 × 10 <sup>-3</sup> M Fe <sup>3+</sup>	otesais	
	2.00 × 10 <sup>-3</sup> M NCS 10 0 0 1		
١	$1.20 \times 10^{-4} \text{M FeNCS}^{2+}$ $1.40 \times 10^{-4} \text{M FeNCS}^{2+}$ $2.00 \times 10^{-3} \text{M Fe}^{3+}$ $2.00 \times 10^{-3} \text{M NCS}^{-}$ $1.40 \times 10^{-4} \text{M FeNCS}^{2+}$ $1.40 \times 10^{-4} \text$		
	Deionized Water		

## Method

The first step in determining the formation constant for the reaction

$$Fe^{3+}_{(aq)} + NCS^{-}_{(aq)} \leftrightarrow FeNCS^{2+}_{(aq)}$$

was to establish a relationship between concentration and absorbance for solutions of FeSCN<sup>2+</sup>. This was accomplished by establishing a Beer-Lambert calibration curve. First, a 3mL cuvette was filled with 1.40 × 10<sup>-4</sup>M FeSCN<sup>2+</sup> and placed into the colorimeter, cycling through the wavelength settings to determine which showed the strongest absorbance reading in response to FeSCN solutions. This wavelength was determined to be 470nm, as can be seen in Table 1. After the appropriate wavelength was chosen, establishing a calibration curve was a matter of filling 3mL cuvettes with the solutions specified in Table 2 and loading them into the colorimeter one at a time. The absorbance values noted in Table 2 were collected in this manner.