# Experiment 7

# Synthesis and Analysis of those Same Green Crystals what we made before Spring Break

Part 3: Spectrophotometric Determination of Iron Content

CH 204 Spring 2009

Dr. Brian Anderson

# Two weeks ago

Redox Chemistry

Oxidation — loss of electrons

Reduction — gain of electrons

Balancing redox reactions

Titration with KMnO<sub>4</sub>

## Today's lab in a nutshell

Parts 2 and 3 of the procedure in the lab manual.

- 1) Mix up a series of 5 standards by diluting from a stock solution
- 2) Measure the absorbance of each of the standards
- 3) Make a calibration curve by plotting Absorbance vs Concentration

The iron stock solution in the hood is 0.0191 grams of Fe per liter

#### Part 2 — make up the standard iron solution

Get 10 mL of the iron solution from the hood, and pipette
 5 mL into a 25 mL volumetric flask.

That's a 1 to 5 dilution of the original concentration.

- 2. Add 1 mL of hydroxylamine, NH<sub>2</sub>OH
  2 mL sodium acetate, and
  8 mL 1,10 phenanthroline
- 3. Fill the volumetric flask up to the line with deionized water using a dropper pipette, then mix it, cap it off and let it sit for 20 minutes for the reaction to occur.

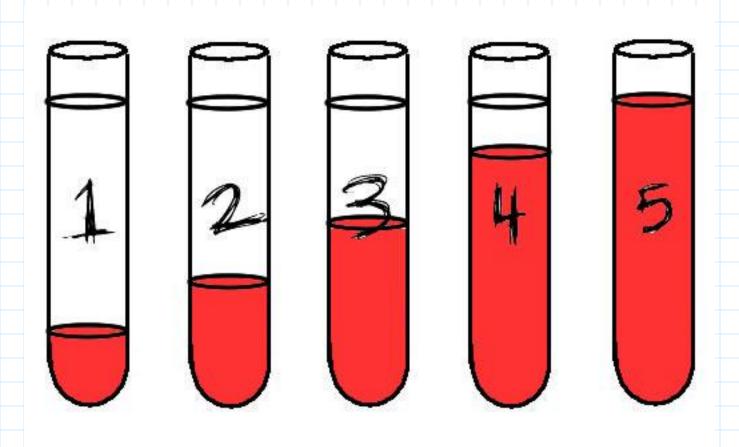
#### Part 3 — Make Individual Standards

1. Get five small test tubes and label them 1, 2, 3, 4, 5. Write directly on the glass with your marker.

Using a plastic syringe, add that many milliliters of the orange solution that you prepared in Part 2 to each test tube.

Using the plastic syringe again, fill each test tube to 5 mL total by adding 4, 3, 2, 1, and 0 mL of deionized water to test tubes 1-5 respectively.

# 5,000 words about Part 3



#### Calculating Final Concentrations

To find the actual concentration of each of the test tubes, we have to multiply by the dilution factor for each one:

Original Concentration (M) imes 1/5 imes test tube dilution factor

This dilution was in part 2 This dilution is in part 3

1: Final concentration = original conc.  $\times$  1/5  $\times$  1/5

2: Final concentration = original conc.  $\times$  1/5  $\times$  2/5

3: Final concentration = original conc.  $\times$  1/5  $\times$  3/5

4: Final concentration = original conc.  $\times$  1/5  $\times$  4/5

5: Final concentration = original conc.  $\times$  1/5  $\times$  1

# Spectrophotometry!

Spectrophotometers are the most widely used analytical instruments in the world except for the analytical balance, and they're about as easy to use as an analytical balance.

"But vat does a spectrophotometer look like?" you are wondering,

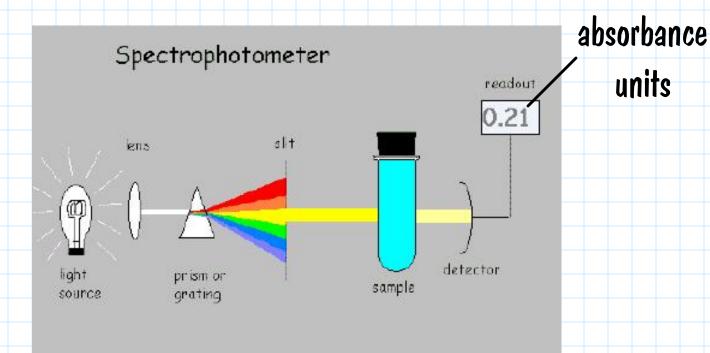
"Und how does it vork?"

I'm glad you asked!

## Looks like this



#### Works like this



Anything that is colored has color because it absorbs some wavelength (or wavelengths) of visible light.

## Using the spectrophotometer

- 1. Place a cuvette full of deionized water into the instrument.

  This is your blank. Press the button that says O ABS.
- 2. Remove the blank and put in a cuvette containing your first standard. The display will automatically read out the absorbance. Record this value.
- 3. Repeat steps 1 and 2 for each solution.
  - 2 cuvettes to a customer! Reuse the sample cuvette!

# How not to screw up this part

- 1) Rinse the cuvette twice with the sample you are about to measure before you put it in the instrument
- 2) Wipe the outside of the cuvette clean using Kim-Wipes. No fingerprints, no wetness on the outside.
- 3) No bubbles in the solution.
- 4) Fill the cuvettes at least 3/4 of the way up.

But what do these absorbance values tell us?

#### Beer's Law

Beer's Law says that absorbance depends on three factors: molar absorptivity, concentration, and path length.

$$A = \varepsilon cI$$

Sometimes written as  $A = \varepsilon bc$ 

or A = abc

## Beer's Law plots

When we plot Absorbance versus Concentration, the slope of the line is equal to  $\varepsilon$ l. In our case l=1, so the slope of the line is equal to the molar absorptivity for Fe(phen)<sub>3</sub><sup>2+</sup>.

# After you have your data

Enter the absorbance and concentration values into Excel.

Plot Absorbance (y-axis) versus concentration (x-axis). Set the y-intercept equal to zero.

You should get a straight line, and the slope of the line is your molar absorptivity,  $\varepsilon$ , in units of  $M^{-1}cm^{-1}$ . Have Excel display the equation for the line on the graph.

### Next veek

- Dissolve up some crystals
- Convert them to orange complex ion
- Measure absorbance
- Determine concentration using Beer's Law

#### Quiz Next Week!

Beer's Law

**Dilutions** 

## Also Next Week!

Turn in Post-lab 7 next week just like it was a pre-lab.

#### Quiz This Week!

After today you are 2/3 finished with the Final Exam!

## Quiz Next Week!

Covering dilutions and Beer's Law