



## EXPERIMENT 3.2

## VISIBLE LIGHT SPECTROSCOPY - COLOURIMETRY

**AIM:**

To measure how much copper is in the leach solution (copper sulphate solution)

**INTRODUCTION:**

Spectroscopy is a means of finding unknown concentrations of chemicals in aqueous solution. Visible light spectrophotometers are often referred to as colourimeters as they use "normal" coloured light to measure concentration.

Any solution which is coloured will absorb one of the wavelengths of visible light but reflect the others – thus giving it a colour. For example, copper sulphate which is a blue solution actually absorbs light in the red to orange range ( $\sim 700$  nm).

Because the red colours are absorbed, and only the other colours reflected, the copper sulphate solution looks blue.

The spectrophotometer/colourimeter simply shines a red light through the copper sulphate solution and measures how much is received the other side of the sample. The missing light is said to be absorbed by the solution, and is called the "absorbance". The more concentrated the solution the greater amount of light is absorbed (this is known as Beer's Law).

The absorbance can range between zero (the solution does not absorb any light – eg water) and one (the solution absorbs all the light). Using a colourimeter allows you to measure the absorbance of an unknown sample. By comparing the absorbance of an unknown sample to absorbances of known concentration (called standards), the concentration of the unknown solution can be worked out.

Using accurate standards (solutions of known concentrations) is an important part of colourimetry. Plotting a graph of absorbance versus concentration for the standards results in a straight line on a graph. You can use this relationship to interpolate any absorbance value within the range of your standards. Extrapolation is not considered accurate methodology with colourimetric data. If your unknown has an absorbance above the range of your standards then accurately dilute the unknown to provide a lower absorbance value that is within the range of your standards - your teacher will show you how to do this.



**MATERIALS:****For colourimetry:**

- 3 sample bottles of leach solution
- Dataquest and colourimeter
- Cuvette and tissue
- Plastic pipette x 8
- Distilled water

*Standards:*

- 3.0 g per litre Copper
- 6.0 g per litre Copper
- 9.0 g per litre Copper
- 12.0 g per litre Copper
- 15.0 g per litre Copper

**METHOD:**Using Colourimeter:

1. Turn on the Dataquest and plug the colourimeter into channel one (at the top). The Dataquest will automatically recognise the sensors as a colourimeter. The absorbance reading will appear in a band (orange) on the main screen of the Dataquest.
2. Use a permanent marker to place a dot on the top of one of the cuvette (not the cuvette lid, the actual cuvette) supplied with the colourimeter. Fill this cuvette about  $\frac{3}{4}$  full with distilled water and wipe the clear plastic sides with a dry tissue so they are as clean as possible.
3. Place this cuvette in the colourimeter so that the clear plastic sides are facing to the front and the rear of the colourimeter. Remember the direction the dot is facing – the cuvette must always be placed in the machine in this way! Close the lid on the colourimeter.
4. The distilled water has a copper concentration of zero, so we will use it to "zero" (means calibrate) the colourimeter. Press the zero button on the colourimeter.
5. Record the concentration of the solution (it is distilled water so it is 0.0 g per litre of Copper), and the absorbance reading (on the Dataquest) in your results table on the following page, even if the absorbance is zero.
6. Take out the cuvette. Pour the contents into the sink. Using the plastic pipette, fill the cuvette to approximately  $\frac{3}{4}$  full with some of the 3.2 g/L copper standard solution. Pour this out into the sink (it is a rinse solution). Again fill the cuvette to  $\frac{3}{4}$  full with some of the 3.2 g/L copper standard. Wipe the sides clean and dry with a tissue, place back in the cuvette and close the lid.
7. Again record the concentration of the solution (3.2), and the absorbance reading (on the Dataquest) in your results table.
8. Repeat steps 6 to 7 until you have recorded the absorbance of all the standards.



**Table 1**

Concentration of copper (g per litre)	Absorbance
3.0	
6.0	
9.0	
12.0	
15.0	

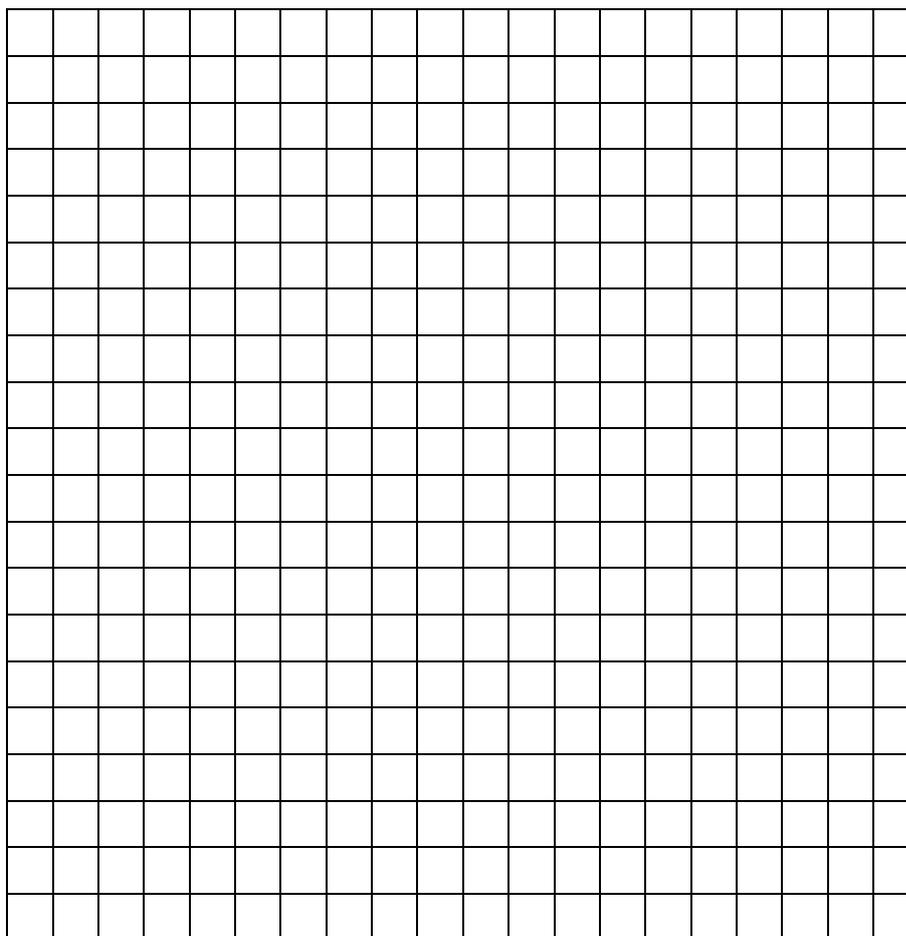
**Table 2**

Leach sample	Absorbance	Concentration of copper (g per litre)
1		
2		
3		

**Table 3**

Volume of final leach sample	=	mL
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9. Your teacher will show you how to graph your results as a scatterplot in excel. This type of graph is used all the time by scientists. As well, graph your results on the graph paper below. Mark each absorbance and concentration value with a cross on the graph and draw a straight line of best fit. Your teacher will show you how to do this. The line shows the relationship between concentration of copper in the solution and the absorbance





10. Use the graph to interpolate (fancy word for reading between the lines) the concentration of each of your leach solutions. You teacher will explain this technique. Record the concentration of copper of each leach solution in table two on the previous page.

QUESTIONS and ANALYSIS

1. Which leach solution (1, 2, 3?) had the highest concentration of copper in it. \_\_\_\_\_

2. (a) Describe the pattern you can see in the copper concentration of the three leach samples?

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(b) Is this pattern what you would have expected? Explain why or why not?

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3. You now know the concentration of leach sample three. In table three you recorded the volume of leach solution three. Use this data to calculate the total amount of copper metal that should be in Leach solution 3.

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Metal Extraction Chemistry