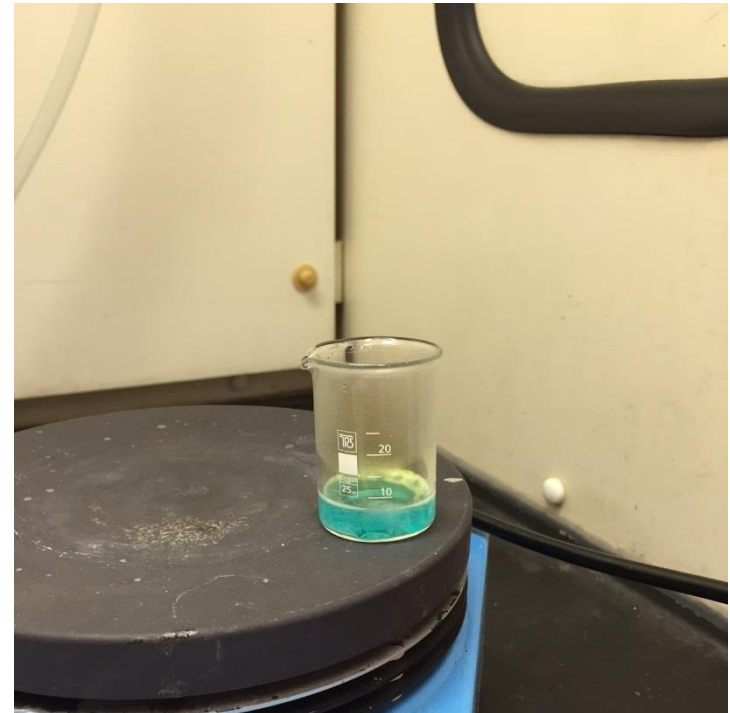
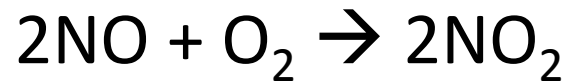
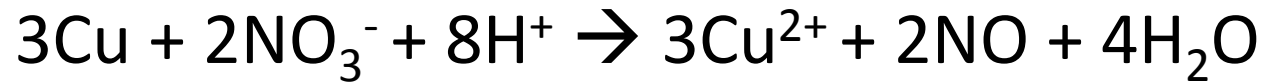
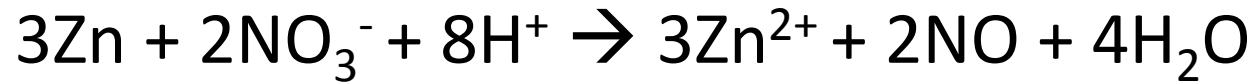


# The determination of copper in brass

## Objective

- To determine the amount of copper in a brass sample

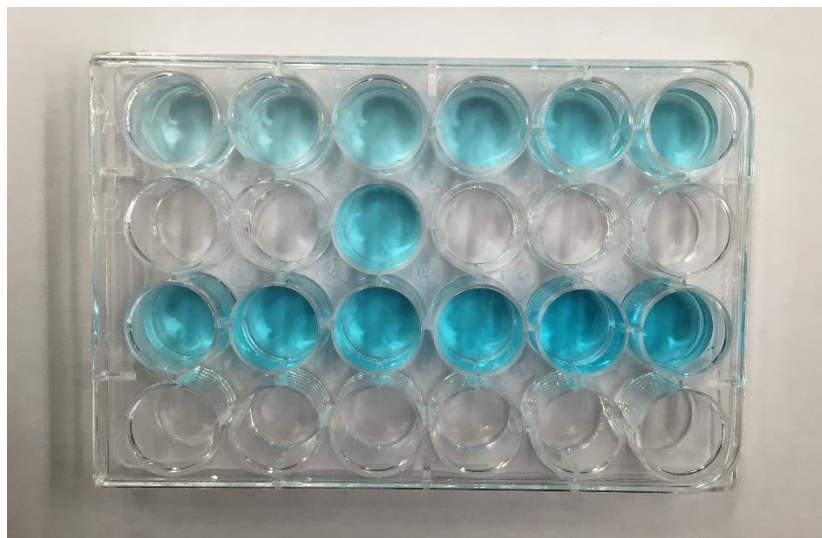
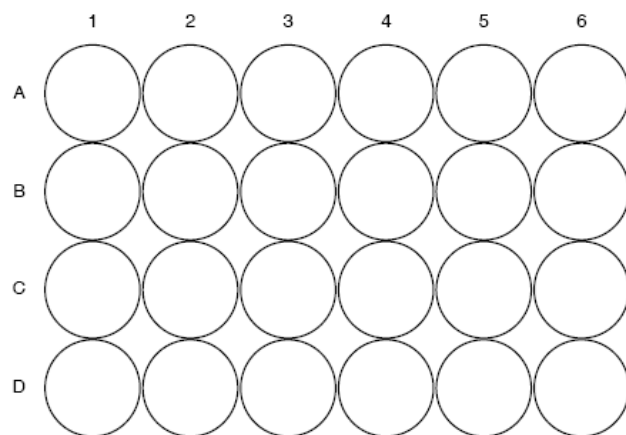
## Chemical reactions



# The determination of copper in brass

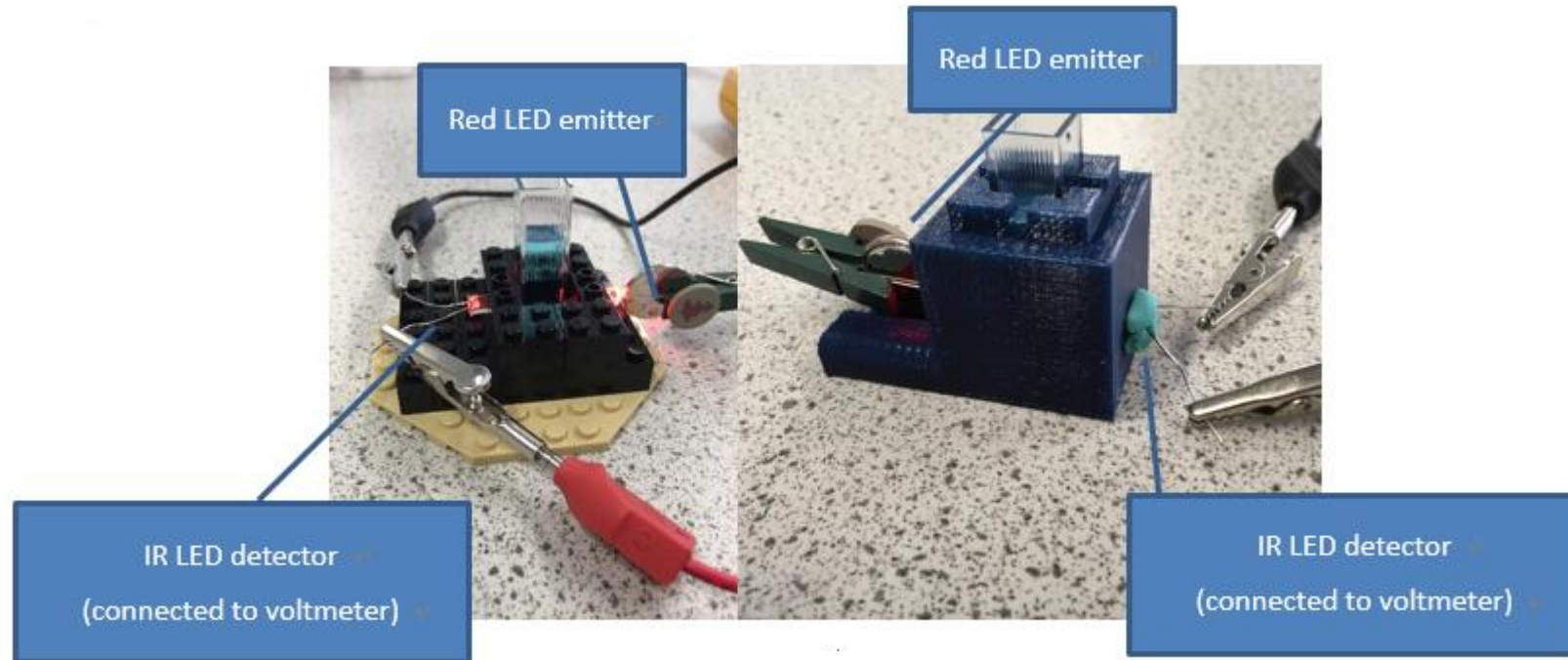
## (A) Microscale method

Well No	A1	A2	A3	A4	A5	A6	C1	C2	C3	C4	C5	C6
Drops of $0.50 \text{ mol dm}^{-3}$ copper nitrate solution	8	10	12	14	16	18	20	22	24	26	28	30
Drops of water	32	30	28	26	24	22	20	18	16	14	12	10



# The determination of copper in brass

## (B) Instrumental method



# The determination of copper in brass

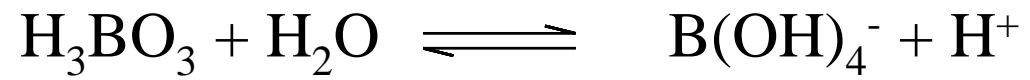
$$\text{Relative absorbance} = \log_{10}(V_0 / V_s)$$

where  $V_0$  is the voltage reading with a cuvette containing just deionised water

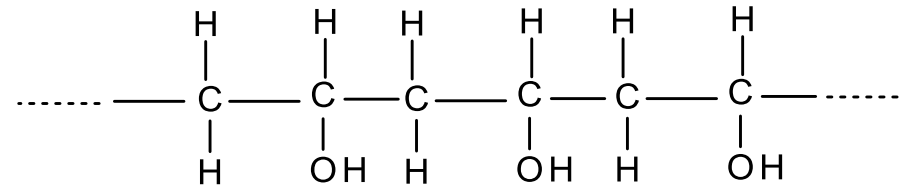
$V_s$  is the voltage reading with a sample in the cuvette

	Deionised water	0.1 M $\text{CuSO}_4(\text{aq})$	0.2 M $\text{CuSO}_4(\text{aq})$	0.3 M $\text{CuSO}_4(\text{aq})$	0.4 M $\text{CuSO}_4(\text{aq})$	0.5 M $\text{CuSO}_4(\text{aq})$	Sample solution
Voltage reading / V							
Relative absorbance							

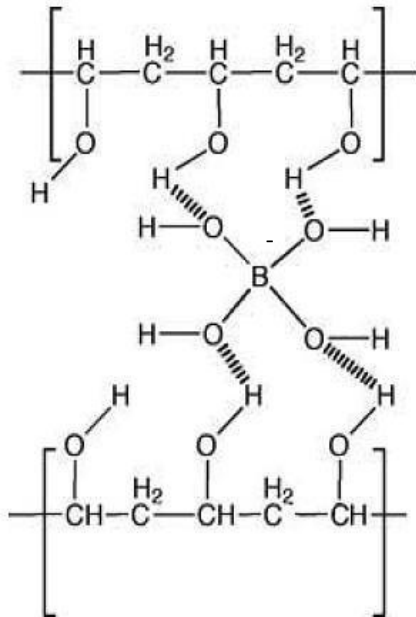
# Preparation of “Slime”



## Boric acid



PVA



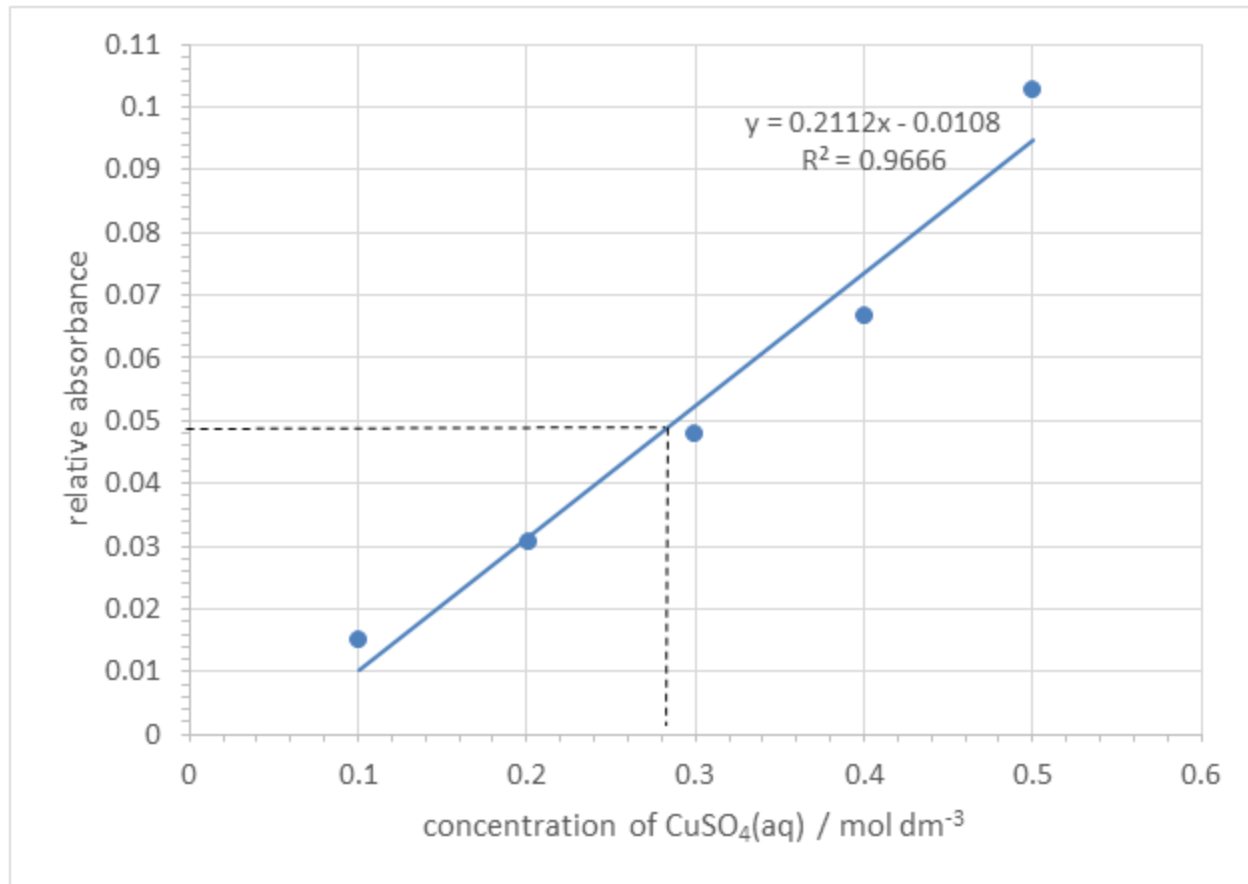
## Slime

## Teachers' notes

- The determination of copper in brass
- Preparation of “Slime”

# The determination of copper in brass

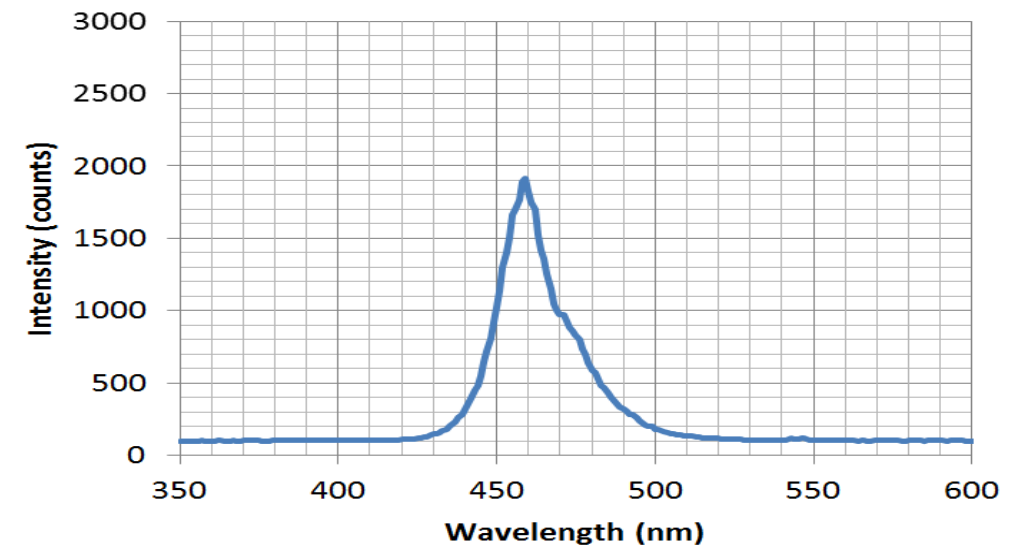
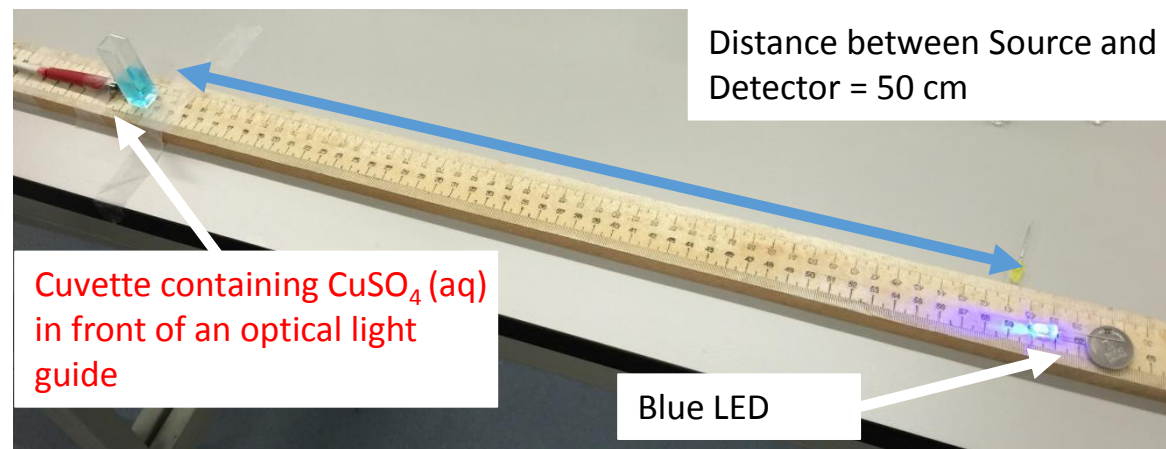
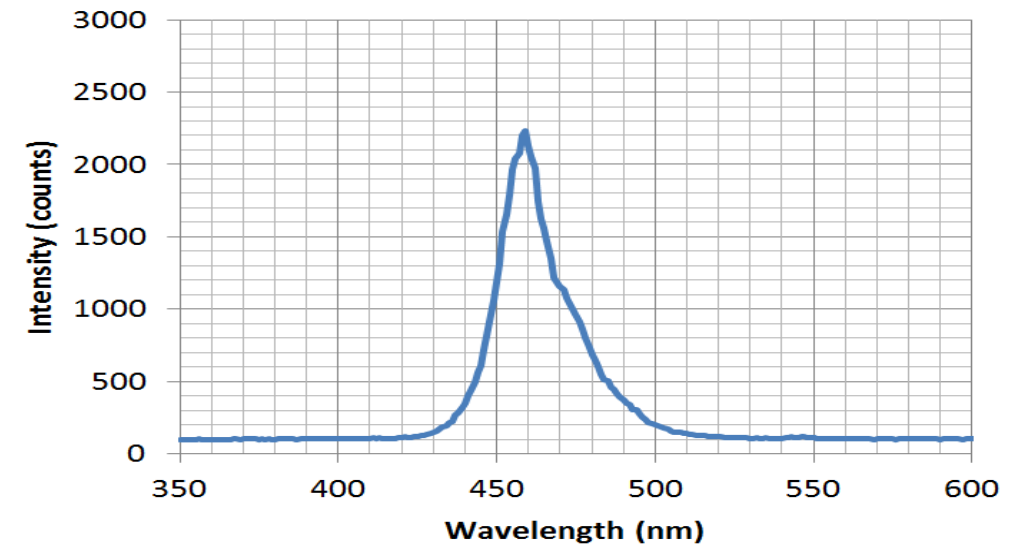
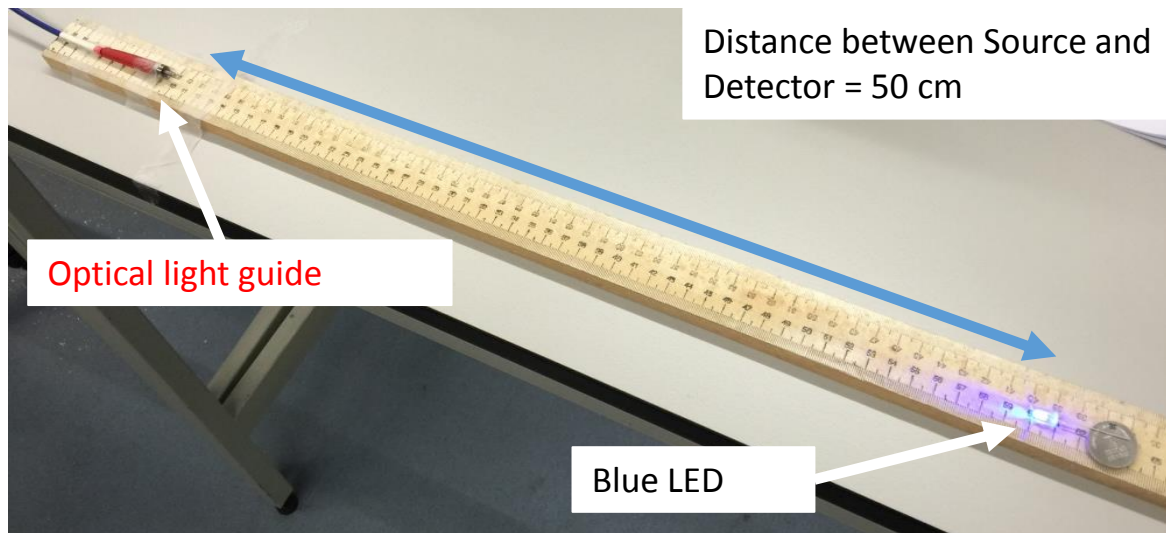
- Instrumental method



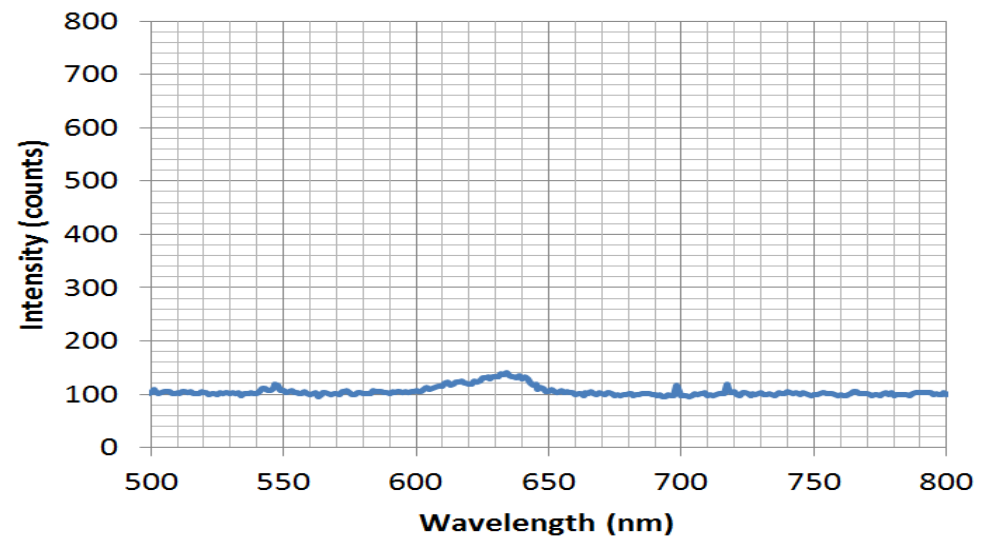
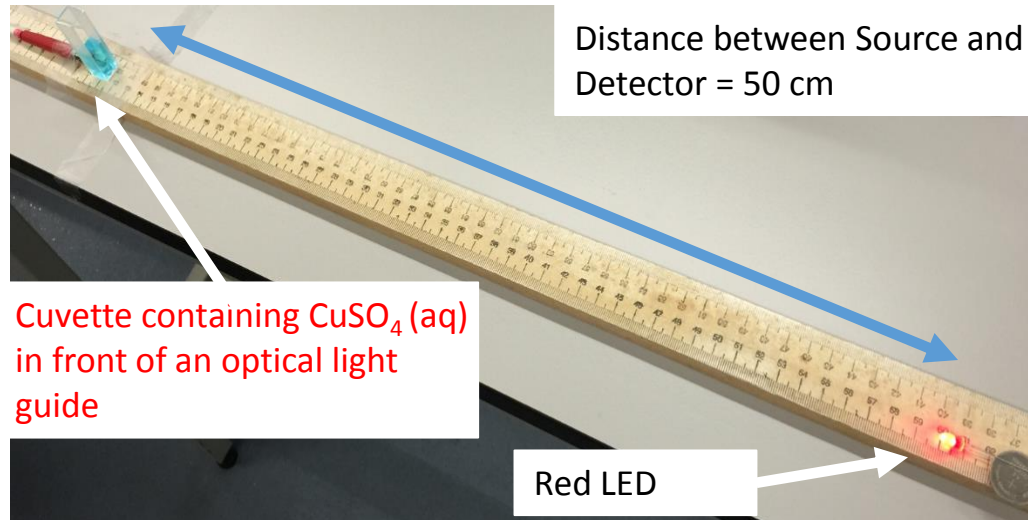
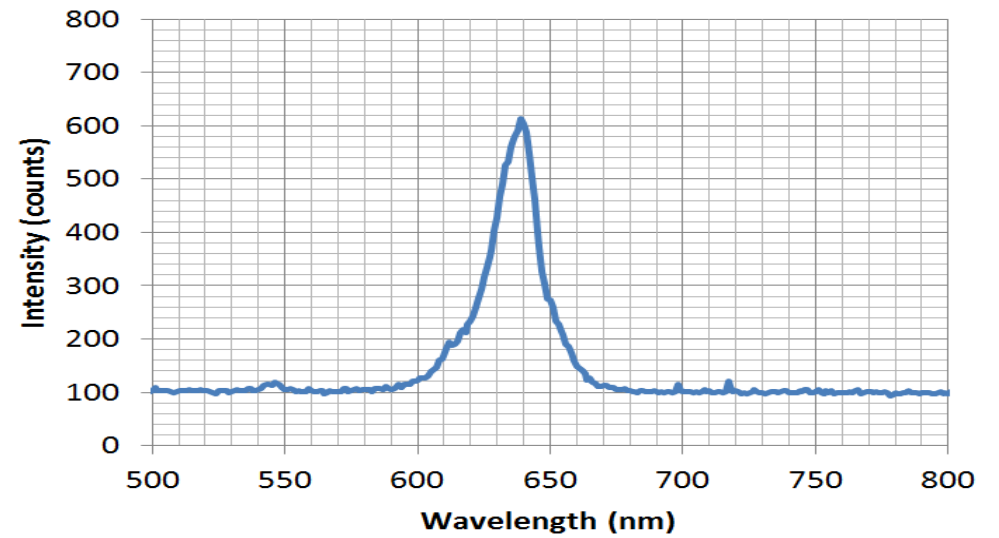
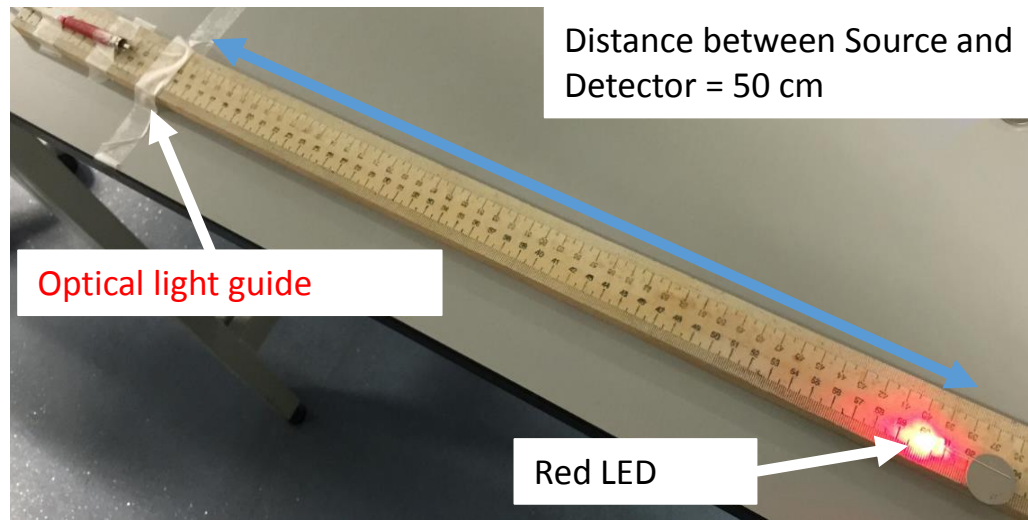
Conc. of sample  $\text{CuSO}_4(\text{aq}) = 0.281 \text{ M}$

➔ Mass of Cu in brass solution = 0.178 g

➔ % by mass of Cu in brass sample = 59.3%



- Blue LED gives peak intensity at 450 - 470 nm
- The peak intensities of the two setups are similar, i.e. no significant blue light absorption is found in  $\text{CuSO}_4$  solution



- Red LED gives peak intensity at 620 - 650 nm
- The peak intensity of the setup with cuvette is significantly lower, i.e. strong red light absorption is found in  $\text{CuSO}_4$  solution

# The LED colorimeter

- The **voltage** produced by the detector LED **may not be directly proportional to the light intensity** shining on the LED
- The Beer Lambert law **does not apply** if there is more than **one colour** change in the reaction
- The detector LED may be **saturated in very bright light**

# The Beer-Lambert Law

$$A = \varepsilon l c$$

*(A = absorbance; l = length of light; c = conc. of pigment;  
 $\varepsilon$  = constant)*

$$A \propto \log_{10} (V_o / V_s)$$

*Where*

*V<sub>o</sub> = voltage for blank solution*

*V<sub>s</sub> = voltage for sample solution*

# Preparation of “Slime”

Preparation of:

**4% PVA**

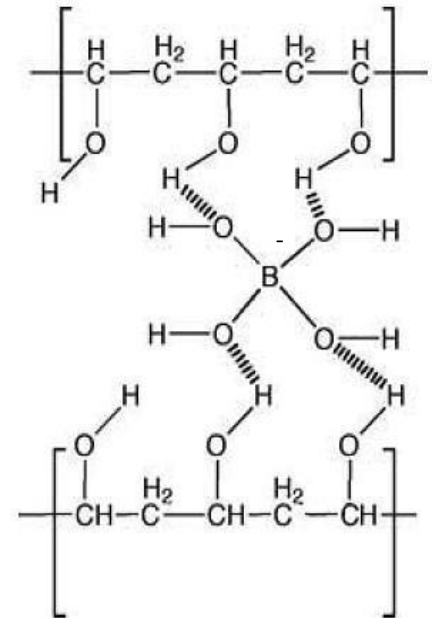
960 cm<sup>3</sup> of D. I. water + 40 g high M.W. PVA with stirring

**4% sodium tetraborate**

solid borax ( $\text{Na}_2\text{B}_4\text{O}_7 \cdot 10\text{H}_2\text{O}$ ) dissolve in D. I. water to make  
4% by weight sodium tetraborate

# Preparation of “Slime”

- The slime can adhere water-soluble ink



## H-bond Cross-linkages

Deformation of cross-linkage network through pouring, squeezing...

<https://drive.google.com/open?id=0BzCEkXNdGw3SR3lXd0RlWmNEVk0>