

# Ion Migration – Copper(II) chromate(VI)

## Introduction

We had some queries from technicians unable to get this demonstration to work. Furthermore we found that some younger technicians and chemistry staff had never even heard of it! It can be used in Standard Grade Chemistry to help show the existence of ions.

## What you will need

### Chemicals

copper(II) chromate(VI) solid (toxic)  
hydrochloric acid, 1M (irritant)  
nitric acid, 1M (corrosive)  
urea

### Equipment

filter funnel  
filter paper  
beaker, 250 cm<sup>3</sup>  
hot plate  
spatula  
stirring rod  
thermometer, stirring  
teat droppers, 2 off  
"W" tube (Griffin Education Catalogue, page 101 Cat. No EKW-224-552B, £7, [1])  
beaker, 500 cm<sup>3</sup> (used as a water bath with cold water to minimise convection of solutions)  
stand, with bosshead and clamp  
power supply, low voltage (0-12 V dc)  
wires, platinum (for electrodes)  
crocodile clips, 2 off  
leads, 4 mm connecting, 2 off  
goggles, indirect vent  
gloves, nitrile

## Preparing the solution

Heat some 1M hydrochloric acid to about 60-70°C and add excess copper(II) chromate to make a hot saturated solution. Allow to cool and filter off the excess solid.

Dissolve urea in this solution. Urea is very soluble and a large amount will be required. The density of this solution is critical, not the concentration of urea it. This should be sufficient to allow the formation of two distinct layers when added to the 1M nitric acid (see 'Setting up the apparatus'). Something approaching a saturated solution of urea in the copper chromate is ideal.

## Setting up the apparatus

Use a teat dropper to place some 1M nitric acid down the back limb into the "W" tube (Figure 2).

Now use a second dropper to carefully and slowly pour copper(II) chromate(VI) solution down the back limb into the "W" tube, making sure the tip of the dropper touches the glass. This should form a bottom layer and force the nitric acid up the side arms of the tube.



Figure 1 - Apparatus & chemicals



Figure 2 - Using dropper



Figure 3 - with electrodes

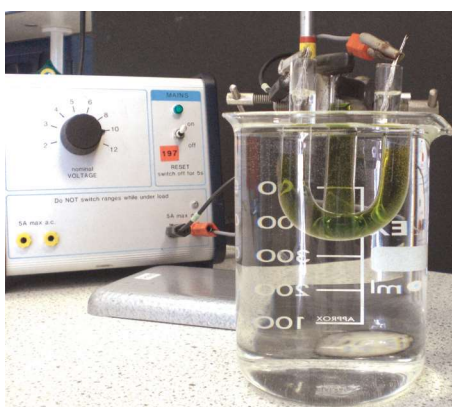


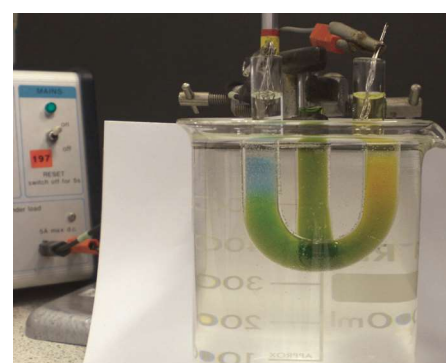
Figure 4 - Connect Pt electrodes and immerse W-tube partially in water-bath beaker  
Connect up the platinum wires (Figure 3) and immerse these in the top of the nitric acid, ensuring they are some distance from the top of the chromate solution.

Carefully clamp the "W" tube and lower the legs into the 500 cm<sup>3</sup> beaker. Fill the beaker with cold water to minimise convection currents in the W-tube.

## The Demonstration

Set the voltage to between 10-12 V dc and switch on. Bubbles should appear at the wires.

Leave it running for about an hour to see a blue colour in the acid at the negative electrode, and an orange colour in the acid at the positive electrode (Figure 6).



Figures 5 & 6 - Blue and orange colours evident at the negative and positive electrodes after about an hour.

## Hazards and Control Measures

Chemical	Main Hazard	Control Measures
Copper(II) chromate(VI)	May cause cancer and is a skin sensitiser	Avoid raising dust from compounds in the solid state and wear nitrile gloves and indirect vent goggles while preparing the solution.
Hydrochloric acid (1M)	Irritant	Wear gloves and eye protection.
Nitric acid (1M)	Corrosive	Wear gloves and indirect vent goggles.
Hot plate	Electrical: Burns	Check wires etc. & that it has been PAT tested before use; Ensure safety notice on view while hot.
Power supply	Electrical	Check power packs (wires etc) plus been PAT tested before use. Use low voltage.

1. <http://www.catalogue.fisher.co.uk/catalogue/griffinbrowse.htm>  
(Copy "EKW-224-552B" into Product Code box)