



## Making soluble salts practical - Justifying the method

Practical Method (What you did)	Justification (Explaining why you did each step)
<ol style="list-style-type: none"><li>Measure 40 cm<sup>3</sup> sulfuric acid into the 100 cm<sup>3</sup> beaker. The volume does not need to be very accurate, so you can use the graduations on the beaker.</li><li>Set up the tripod, gauze and heatproof mat. Heat the acid <b>gently</b> using the Bunsen burner until it is almost boiling. Turn off the Bunsen burner.</li><li>Use the spatula to add <b>small</b> amounts of copper (II) oxide powder. Stir with the glass rod.</li><li>Continue to add copper (II) oxide if you cannot see it when <b>stirred</b>. When the copper (II) oxide has reacted, the solution is clear blue.</li><li>Stop adding the copper (II) oxide when some of it remains after <b>stirring</b>. Allow the apparatus to cool completely.</li><li>Set up the filter funnel and paper over the conical flask. Use the clamp stand to hold the funnel.</li><li><b>Filter</b> the contents of the beaker from step 3.</li><li>When filtration is complete, pour the contents of the conical flask into the evaporating basin.</li><li>Evaporate this gently using a water bath (250 cm<sup>3</sup> beaker with boiling water) on the tripod and gauze. Stop heating once crystals start to form.</li><li>Transfer the remaining solution to the crystallising dish (evaporating dish). Leave this in a cool place for <b>at least 24 hours</b>.</li><li>Remove the crystals from the concentrated solution with a spatula. <b>Gently</b> pat the crystals dry between two pieces of filter paper. These are pure dry crystals of copper (II) sulfate.</li></ol>	<p>Why did you heat the acid?</p> <p>Why did you only add small amounts of copper oxide powder at a time?</p> <p>Why did you stir the copper oxide?</p> <p>Why did you keep adding copper oxide until no more would react?</p> <p>Why did you need to filter the solution?</p> <p>Why did you leave the solution to cool for at least 24 hours?</p>