

GCSE Chemistry

Paper 2 · Required Practicals - Cheat Sheet

AQA Chemistry (8462)

Paper 2

4 Practicals

Quick-revision reference for the Required Practicals in AQA GCSE Chemistry. For each practical: the **aim**, the **method**, the **variables**, the expected **result**, and the **exam tips** examiners reward. Learn the method and the why, not just the what.

RP5 Rates of Reaction (Cloudiness)

AIM	Investigate how the concentration of a solution affects the rate of reaction using the loss of clarity of a precipitate.
METHOD	1. Draw a black cross on paper and stand a conical flask of sodium thiosulfate on top of it. 2. Add dilute hydrochloric acid and immediately start a stopwatch. 3. Look down through the flask and stop timing when the cloudy sulfur precipitate hides the cross. 4. Repeat with different concentrations of thiosulfate, keeping the total volume and temperature the same.
VARIABLES	IV: concentration of sodium thiosulfate · DV: time for cross to disappear ($\text{rate} = 1 \div \text{time}$) · Control: volume, temperature, depth of liquid, same observer
RESULT	Higher concentration means a shorter time, so a faster rate; rate is proportional to $1 \div \text{time}$. A graph of rate against concentration gives an upward trend.
EXAM TIPS	› Sulfur dioxide gas is produced, which is toxic and an irritant, so work in a ventilated room. › Always have the same person judge when the cross vanishes to keep the end point fair. › Repeat each concentration and take a mean to improve reliability.

RP5 Rates of Reaction (Gas Volume)

AIM	Investigate how a chosen variable, such as surface area, affects the rate of reaction by measuring the volume of gas given off.
METHOD	1. Add marble chips (calcium carbonate) to dilute hydrochloric acid in a conical flask fitted with a bung and delivery tube. 2. Collect the carbon dioxide produced in a gas syringe or an inverted measuring cylinder over water. 3. Record the volume of gas every 10 seconds until the reaction stops. 4. Repeat using chips of a different size (large lumps versus small lumps or powder), keeping mass and acid the same.
VARIABLES	IV: surface area of the marble chips · DV: volume of gas collected per unit time (rate) · Control: mass of carbonate, concentration and volume of acid, temperature
RESULT	Smaller pieces have a larger surface area, giving a steeper curve and faster initial rate; all runs reach the same final gas volume when the limiting reactant is used up.
EXAM TIPS	› The steepest part of the graph is at the start when reactant concentration is highest. › A flat (horizontal) line shows the reaction has finished. › Fit the bung quickly after adding the acid so no gas escapes before measuring begins.

RP6 Chromatography

AIM	Use paper chromatography to separate a mixture of soluble dyes and calculate the R_f value of each spot.
METHOD	1. Draw a pencil baseline near the bottom of the chromatography paper and add a small spot of each ink on it. 2. Place the paper in a beaker so the solvent level sits below the baseline, then cover it. 3. Let the solvent rise up the paper, carrying the dyes, then remove it before the solvent reaches the top. 4. Mark the solvent front in pencil, leave the paper to dry, and measure the distances moved.
VARIABLES	IV: the different dyes or inks tested · DV: distance each spot travels (used for R_f) · Control: same solvent, same paper, same starting line
RESULT	Each dye moves a characteristic distance, giving $R_f = \text{distance moved by spot} \div \text{distance moved by solvent}$. R_f is always between 0 and 1, and more soluble dyes travel further.
EXAM TIPS	› Always draw the baseline in pencil, because ink would dissolve and run up the paper. › Keep the spots above the solvent so the dye is not washed straight into the beaker. › Measure to the centre of each spot when finding its distance for a fair R_f.

RP8 Identifying Ions and Water Analysis

AIM	Use chemical and flame tests to identify the cations and anions present in unknown salts and samples of water.
METHOD	1. For metal cations, dip a clean wire loop in the sample and hold it in a blue Bunsen flame to see the flame colour. 2. Add sodium hydroxide solution to a portion to test for metal hydroxide precipitates and their colours. 3. Test for carbonates by adding dilute acid and bubbling any gas through limewater. 4. Test for halides with nitric acid then silver nitrate, and for sulfates with hydrochloric acid then barium chloride.
VARIABLES	IV: the ion being tested for · DV: observation (flame colour, precipitate colour, gas produced) · Control: clean apparatus, same volumes, fresh wire loop each time
RESULT	Each ion gives a characteristic result: for example lithium burns red, sodium yellow, potassium lilac; carbonates fizz and turn limewater milky; halides give coloured silver halide precipitates; sulfates give a white barium sulfate precipitate.
EXAM TIPS	› Clean the flame test wire in acid between samples so a leftover ion does not mask the next colour. › Use nitric acid before silver nitrate, and hydrochloric acid before barium chloride, to remove ions that would give a false positive. › Flame and chemical tests struggle when a mixture of ions is present, as one strong colour can hide another.