**A Level Chemistry Required Practicals Revision Summary**

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| **Topic** | **Required practical** | **Notes** |
| 3.1.2: Amount of substance | 1 | Make up a volumetric solutionCGP Y1 p49 | Method tips to ensure **accurate** concentration:* Re-weigh bottle and calculate mass of solid by difference (some may be left in bottle)
* Wash the stirring rod into the flask / beaker after use
* Wash the beaker into the flask after transfer
* Wash the filter funnel into the flask
* Use a teat pipette to make up to the mark on flask
* Bottom of the meniscus is on the graduation mark
* Mix / shake the final solution in the flask by inverting.
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| 1 | Carry out a simple acid-base titrationCGP Y1 p50 | To ensure accuracy:* Rinse pipette & burette with solutions
* Read from bottom of meniscus
* Watch out for air bubbles
* Few drops of indicator only
* Swirl contents of flask to ensure good mixing
* Add slowly near the end point
* Rinse conical flask with distilled water near end point to ensure all solutions are in flask to react
* Use while tile to aid seeing colour change of indicator
* First rough unless shown to be concordant.
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| 3.1.4: Energetics | 2 | Measurement of an enthalpy changeCGP Y1 p106-107 | Two solutions or solid reactant added to solution.Use insulated container with lid to minimise heat loss or gain.Stir throughout to ensure even mixing.Plot data & draw 2 lines of best fit then extrapolate back to the time when the reactants were mixed to find the temperature change. |
| 3.1.5: Kinetics | 3 | Investigation of how the rate of a reaction changes with temperatureCGP Y1 p119 | The main problem with kinetics practicals is any variation in temperature will have a large effect on the rate, this can be minimised by controlling the temperature of all reactants by using a thermostatically controlled water bath. |

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| 3.2.2: Group 23.2.3: Group 7 | 4 | Carry out simple test-tube reactions to identify cations and anionsCGP Y1 p156 & p165-166 | Exact quantities not necessary as it is qualitative testing. Descriptions:* Colour
* State e.g. precipitate (ppt) or solution (aq)
* Describe change e.g. re-dissolving
* No visible change

If solution is coloured and masks colour of precipitate, decant solution and wash precipitate.Standard tests and expected results:*Carbonate ions*Test: dilute hydrochloric acid then test gas with limewater Result: goes cloudy (due to CO2)*Sulphate ions* dilute hydrochloric then barium chloride solution Result: white precipitate – BaSO4(s)*Chloride, bromide and iodide ions*Test: dilute nitric acid then silver nitrate solutionResults:chloride - white precipitate (AgCl)bromide - cream precipitate (AgBr)iodide - yellow precipitate (AgI)Further test: addition of ammonia solutionsResults:chloride - precipitate dissolves in dil. NH3bromide - precipitate dissolves in conc. NH3iodide - precipitate insoluble in dil. & conc. NH3. |
| 3.3.5: Alcohols | 5 | Distillation of a product from a reactionCGP Y1 p228-229 | To allow separation of a volatile liquid from another liquid, the product of the reaction is collected.Anti-bumping granules in the flask are a safety measure to prevent vigorous or uneven boiling through the release of large gas bubbles. They ensure only small bubbles are formed.*Distillation*http://www.micromountain.com/sci_diagrams/sci_app/sci_app_assets/distil_lab_eng.jpg*Refluxing*Don’t confuse with distillation. In this technique the volatile product condenses back into the flask so isn’t isolated and it reacts further.File:Reflux labled.svg |
| 3.3.6: Organic analysis | 6 | Tests for alcohol, aldehyde, alkene and carboxylic acid | Standard tests and expected results:*Alcohol*Test: metallic sodiumResult: effervescence (due to H2)*1O/2O alcohol*Test: Acidified potassium dichromate Result: orange to green*Aldehyde*Test: Fehling’s solutionResult: red precipitateTest: Tollens’ solutionResult: silver mirror*Alkene (unsaturation)*Test: bromine waterResult: decolourises (turns colourless due to addition of bromine to alkene)*Carboxylic acid*Test: solid sodium hydrogencarbonateResult: effervescence (due to CO2)*Haloalkane*Test: sodium hydroxide then warm and add nitric acid followed by silver nitrate solutionResults:chloride - white precipitate (AgCl)bromide - cream precipitate (AgBr)iodide - yellow precipitate (AgI) |
| 3.1.9: Rate equations | 7a | Measuring the rate of reaction by an initial rate methodCGP Y2 p58 | Time colour change – blue-black iodinePlot graph: log10 (1/t) (y) against log10 V (x) Choose points near opposite ends of the line.Measure gradient (to find rate) = y/x |
| 7b | Measuring the rate of reaction by a continuous monitoring methodCGP Y2 p42 | Reaction between magnesium and hydrochloric acid, the volume hydrogen gas produced is measured continuously at regular time intervals to monitor the rate of the reaction.Methods of measuring the volume of gas:*Using a gas syringe**Using a water trough – alternative method*Graph:Plot volume of hydrogen produced (*y*-axis) against time.Draw a line of best fit.Draw a tangent to each line at time, t = 0 sCalculate the gradient of each tangent in order to deduce the rate of each reaction. |
| 3.1.11: Electrode potential and cells | 8 | Measuring the EMF of an electrochemical cellCGP Y2 p79 | A cell produces electricity from a chemical reaction when two different metals are connected by a wire and placed in an electrolyte. emf experimentSalt bridge - allows transfer of charge between each half cell. Sandpaper - to remove any metal oxide that may have formed on the surface of metal.Electrolyte – substance dissolved in water or molten which can carry electrical charge because the liquid contains ions that are free to move.  |

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| 3.1.12: Acids and bases | 9 | Investigate how pH changes when a weak acid reacts with a strong base and when a strong acid reacts with a weak baseCGP Y2 p106-109 | Calibration of pH meter – check pH readings against buffers of known pH.Calibration graph: recorded pH reading (x axis) against the pH of the buffer solution.Record pH after each addition of alkali.Obtain correct pH reading from calibration graph.S-curve graph: corrected pH values (y) against volume (x). |
| 3.3.9: Carboxylic acids | 10a | Preparation of a pure organic solidCGP Y2 p195 | Aspirin is prepared by the acylation of salicylic acid using ethanoic anhydride as the acylating agent.Recrystallisation technique - filter under reduced pressure.http://t0.gstatic.com/images?q=tbn:ANd9GcS4rTyha2Uc9yygm7b9pgXLxLFr7Z-_A04WyYtalpgtqZ0JTWqJ |
| Test of its purityCGP Y2 p195 | The purity of an organic solid can be determined in part by measuring its melting point and comparing the value with the known melting point.A pure dry solid will melt at a precise temperature whereas an impure solid will melt over a range of temperatures which are lower than the melting point of the pure solid.Heat a small quantity of the solid in a thin-walled melting point tube using an oil bath (not water – temperature too low) or in melting point apparatus.Test first to establish approximate melting point, then repeat and heat **slowly near the melting point** until the solid collapses into a liquid without any change in temperature.The method will not work if the solid decomposes on heating.Subsequent heating of other samples can be done more slowly.If heat too fast too high a melting point will be determined.If melting point is measured at a temperature that does not match the published temperature it’s because temperature on thermometer is not the same as the sample. |
| 10b | Preparation of a pure organic liquidCGP Y2 p193-194 | Reflux to produce ester followed by distillation to isolate it.See earlier notes on the techniques and use of anti-bumping granules.Separating funnel – used to separate immiscible liquids. Image result for separating funnelSolution sodium carbonate added to react with any acid catalyst, this will stop the reaction and remove the acid. Separating funnel is stopped and inverted several time to ensure acid and carbonate react. Carbon dioxide gas is produced in the neutralisation reaction. |
| 3.2.6: Reactions of ions | 11 | Carry out simple test-tube reactions to identify transition metal ions in aqueous solutionCGP Y2 p162-163 | Learn the colour changes and states (solution or precipitate) for each reaction – see notes given. |
| 3.3.16: Chromatography | 12 | Separation of species by thin-layer chromatographyCGP Y2 p265 | Use pencil to draw base line. Place multiple small spots to produce a concentrated spot of the compounds to be used by repeated application followed by drying.Mark a reference spot (known compound).Mark a spot of unknown substance.Place in chromatography tank; ensure that the spots are above the solvent.Cover with lid to saturate the atmosphere with the solvent, the lid prevents evaporation.Allow time for the solvent to run up the chromatography plate, so it reaches just below the end of the paper.Use locating agent to find spots, e.g. ninhydrin, UV light.Compare the unknown compound with the reference compound or use Rf values.Calculate Rf value- measure distance solvent has travelled and from the baseline to solvent front- measure distance each compound has travelled from baseline to centre of each spot. |
| 3.2.5:Transition metals | Extra | Colorimetry | Sample of know concentration of a coloured solution placed in colorimeter to obtain an absorbance reading.Solution must have an intense colour:* Can intensify by reacting with another reagent
* If too intense can dilute with water.

Plot graph: absorbance (y) against concentration (x)Measure absorbance of unknown and use the graph to obtain the concentration value. |
| Extra | Redox titration | Acidified with sulphuric acid to prevent oxidation of potassium manganate(VII) to MnO2 – a brown solid.Potassium manganate(VII) **self indicating.**Read from **top of meniscus.**End point pale pink (permanent pink colour). |

**Other information**

**Graphs:**

* Use a **sensible scale** so **plotted points** cover **½ a page**
* May have **negative** numbers – make sure the **scale goes down**
* Don’t need to start at zero – use a broken scale
* **Label axes** with **measurement** & **units**
* **Plot carefully** (± one small square)
	+ Use a **pencil**!
* **Line** of **best fit** may be a **straight line** but look out for obvious **curve**
	+ Use a **long** **ruler** (no kinks or double lines!)
	+ Miss out **anomalous results**
	+ May **not** go through **origin**
	+ But **it may!**
* **Readings** from graph:
	+ Use **ruler & pencil to mark** given value and show value read form the graph

**Formula:**

* If using formula they **must be correct**

**Hazards & risks:**

* Know common **hazards** & **precautions** to minimise risks:
	+ **Flammable** or **explosive** – keep away from naked flame
	+ **Corrosive** – wear gloves/avoid contact with hands
	+ **Toxic** – wear gloves
* **Safety glasses** & **lab coats** are **assumed** they will be ignored as an answer
* **Anti-bumping granules** prevent vigorous/uneven boiling – produce small bubbles

**Errors & calculations:**

* **Repeats** used because a **single** **reading** could be **flawed** or **anomalous**
	+ **Improves reliability** of results
* **Testing** from **different batches** eliminates fact that different batches **could vary** or
**be** **contaminated**
* Calculating **equipment** errors **= margin of error / quantity measured x 100**
	+ **total errors = sum of individual equipment errors**
	+ If reactant is in excess errors not important
	+ **Smaller quantities** have **bigger errors**
		- * + To reduce errors use larger quantities or more precise measuring equipment
* Calculating **experimental** errors **= (true value – experimental value) / true value x 100**
	+ No sign needed