| Atoms, elements and | compounds |
|------------------------|-----------|

| Atom | The smallest part of an element that can exist | Have a radius of around 0.1 nanometres and have no charge (0). |
|----------|---|---|
| Element | Contains only one type of atom | Around 100 different elements each one is represented by a symbol e.g. O, Na, Br. |
| Compound | Two or more elements chemically combined | Compounds can only be separated into elements by chemical reactions. |

| I | Control muslama | 0-1-1 |
|---|-----------------|-------------------------------|
| 1 | Central nucleus | Contains protons and neutrons |
| 1 | Electron shells | Contains electrons |

| 9-5 | | |
|---------------------|--------------------|------------------|
| Name of Particle | Relative Charge | Relative Mass |
| Proton | +1 | 1 |
| Neutron | 0 | 1 |
| Electron | -1 | Very small |

number

| tronic hell | Max number of electrons |
|--------------------|----------------------------|
| 1 | 2 |
| 2 | 8 |
| 3 | 8 |
| 4 | 2 |

number of protons

Electronic

AQA Chemistry

structures

C1: Atomic Structure & The Periodic Table Rutherford's scattering experiment

A beam of alpha particles are directed at a very thin gold foil

passed right through. A few (+) alpha particles were deflected by the positive nucleus.

A tiny number of particles reflected back from the nucleus.

Most of the alpha particles

| Li ⊨ | |
|------|--|
| 3 ← | |

Mixtures

The sum of the protons and neutrons in the Mass number nucleus Atomic The number of Number of electrons =

Two or more elements or compounds

protons in the atom

not chemically combined together

Relative electrical charges of subatomic particles

Can be separated by physical processes.

| Method | Description | Example |
|----------------------------|--|---|
| Filtration | Separating an insoluble solid from a liquid | To get sand from a mixture of sand, salt and water. |
| Crystallisation | To separate a solid from a solution | To obtain pure crystals of sodium chloride from salt water. |
| Simple distillation | To separate a solvent from a solution | To get pure water from salt water. |
| Fractional distillation | Separating a mixture of liquids each with different boiling points | To separate the different compounds in crude oil. |
| Chromatography | Separating substances that move at different rates through a | To separate out the dyes in food colouring. |

Tiny solid spheres that electron, John Dalton said the Pre 1900 could not be divided solid sphere made up the different elements. JJ Thompson 's experiments 1897 A ball of positive charge 0+0+ showed that showed that an atom with negative electrons 'plum must contain small negative pudding' embedded in it charges (discovery of electrons). Ernest Rutherford's alpha particle 1909 Positively charge nucleus scattering experiment showed **(+)** • at the centre surrounded nuclear that the mass was concentrated at model negative electrons the centre of the atom. Niels Bohr proposed that electrons 1913 Electrons orbited in fixed shells; this was Bohr orbit the nucleus at supported by experimental model specific distances observations.

The development of the model of the atom

James Chadwick

Provided the evidence to show the existence of neutrons within the nucleus

Before the discovery of the

Chemical equations

atomic mass

Relative

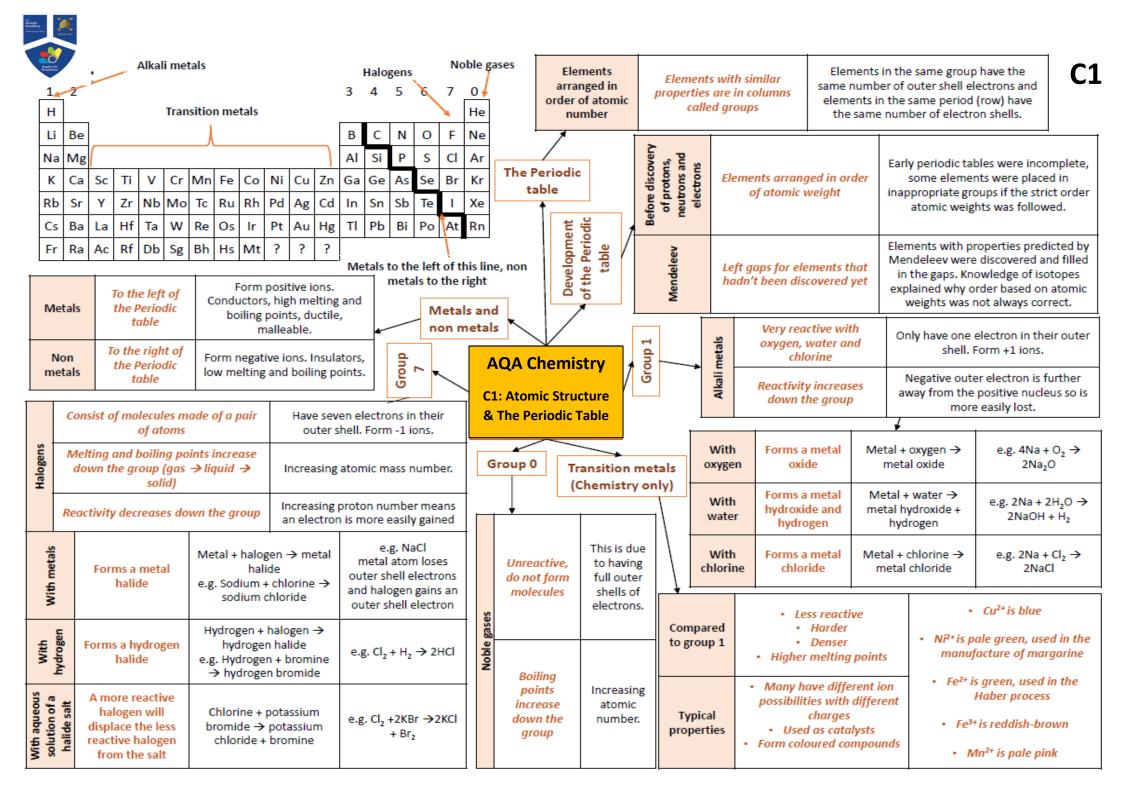
Show chemical reactions - need reactant(s) and product(s) energy always involves and energy change Law of conservation of mass states the total mass of products = the total mass of reactants.

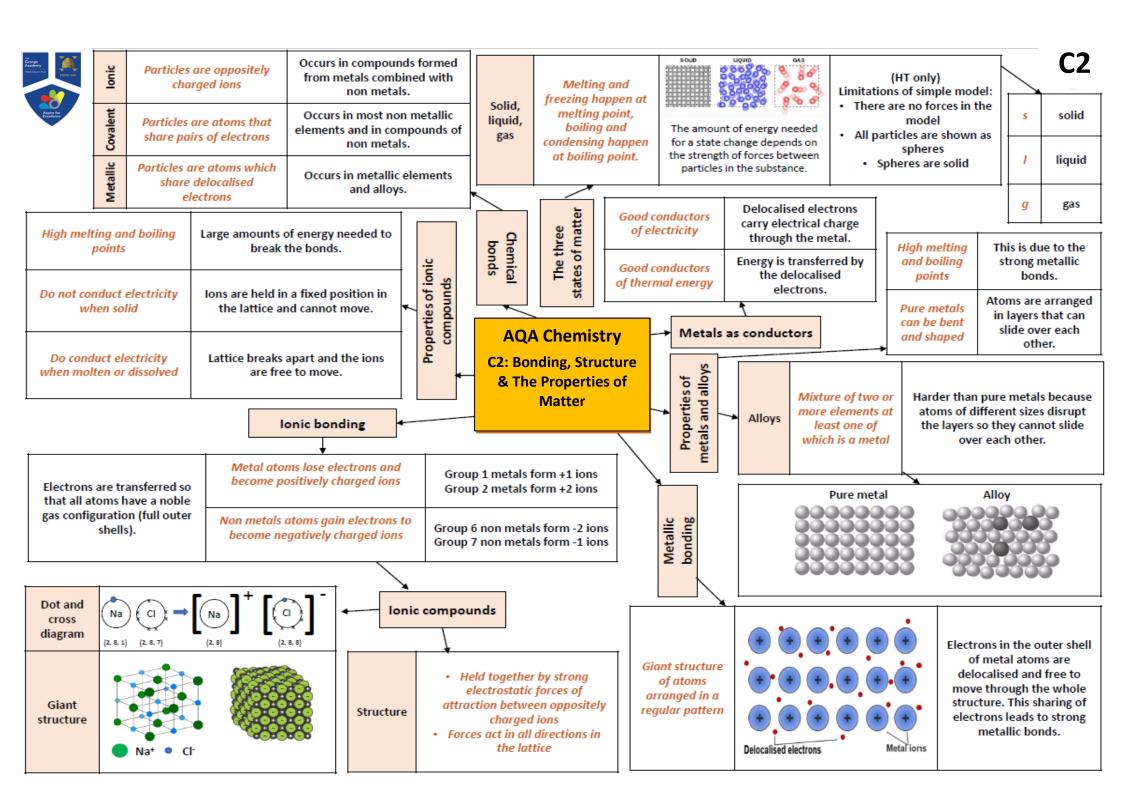
| Word equations | Uses words to show reaction reactants → products magnesium + oxygen → magnesium oxide | Does not show what is happening to the atoms or the number of atoms. |
|------------------|---|---|
| Symbol equations | Uses symbols to show reaction reactants → products 2Mg + O ₂ → 2MgO | Shows the number of atoms and molecules in the reaction, these need to be balanced. |
| | | |

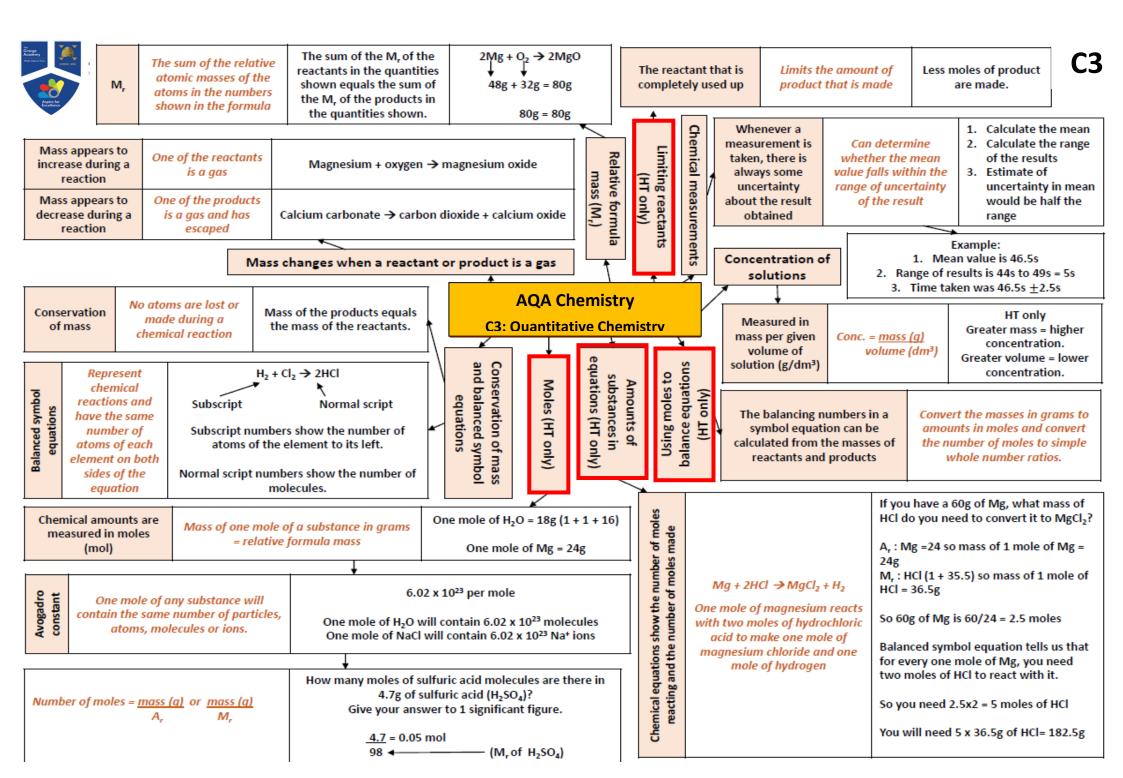
Atoms of the same element with the same number of Isotopes protons and different numbers of neutrons

35Cl (75%) and 37Cl (25%)

Relative abundance = (% isotope 1 x mass isotope 1) + (% isotope 2 x mass isotope 2) ÷ 100 e.g. (25 x 37) + (75x 35) ÷ 100 = 35.5









A measure of the amount of starting materials that end up as useful products Atom economy = Relative formula mass of desired product from equation x 100 Sum of relative formula mass of all reactants from equation

High atom economy is important or sustainable development and economic reasons

Calculate the atom economy for making hydrogen by reacting zinc with hydrochloric acid:

 M_{2} of $H_{2} = 1 + 1 = 2$ M, of Zn + 2HCl = 65 + 1 + 1 + 35.5 + 35.5 = 138

Atom economy = $\frac{2}{138} \times 100$ $= \frac{2}{130} \times 100 = 1.45\%$

This method is unlikely to be chosen as it has a low atom economy.

Concentration of a solution is the amount of solute per volume of solution

Concentration = amount (mol) volume (dm3) (mol/dm3)

What is the concentration of a solution that has 35.0g of solute in 0.5dm3 of solution?

 $35/0.5 = 70 \text{ g/dm}^3$

solutions in mol/dm3 (HT only, chemistry only)

Using concentrations of

AQA Chemistry

C3: Quantitative Chemistry

Percentage

yield

If the volumes of two solutions that react completely are known and the concentrations of one solution is known, the concentration of the other solution can be calculated.

 $2NaOH(aq) + H_2SO_4(aq) \rightarrow Na_2SO_4(aq) + 2H_2O(l)$

It takes 12.20cm3 of sulfuric acid to neutralise 24.00cm3 of sodium hydroxide solution, which has a concentration of 0.50mol/dm³.

Calculate the concentration of the sulfuric acid in mol/dm3:

0.5 mol/dm3 x (24/1000) dm3 = 0.012 mol of NaOH The equation shows that 2 mol of NaOH reacts with 1 mol of H₂SO₄, so the number of moles in 12.20cm³ of sulfuric acid is (0.012/2) = 0.006 mol of sulfuric acid

Calculate the concentration of sulfuric acid in mol/dm3 0.006 mol x (1000/12.2) dm3 = 0.49mol/dm3

HT only:

200g of calcium carbonate is heated. It decomposes to make calcium oxide and carbon dioxide. Calculate the theoretical mass of calcium oxide made.

$$CaCO_3 \rightarrow CaO + CO_2$$

M_r of $CaCO_3 = 40 + 12 + (16x3) = 100$

 M_{\bullet} of CaO = 40 + 16 = 56

100g of CaCO3 would make 56 g of CaO

So 200g would make 112g

Use of amount of substance in relation to volumes of gases (HT only, chemistry only)

Calculate the concentration of sulfuric acid in g/dm3:

 $H_2SO_4 = (2x1) + 32 + (4x16) = 98g$ $0.49 \times 98g = 48.2g/dm^3$

Yield is the amount of product obtained

It is not always possible to obtain the calculated amount of a product

The reaction may not go to completion because it is reversible.

Atom economy

Some of the product may be lost when it is separated from the reaction mixture.

Some of the reactants may react in ways different to the expected reaction.

Equal amounts of moles or gases occupy the same volume under the same conditions of temperature and pressure

The volume of one mole of any gas at room temperature and pressure (20°C and 1 atmospheric pressure) is 24 dm^3

No. of moles of gas = vol of gas (dm3) 24dm³

Percentage yield is comparing the amount of product obtained as a percentage of the maximum theoretical amount

% Yield = Mass of product made x 100 Max. theoretical mass

A piece of sodium metal is heated in chlorine gas. A maximum theoretical mass of 10g for sodium chloride was calculated, but the actual yield was only 8g.

Calculate the percentage yield.

Percentage yield = 8/10 x 100 = 80%

What is the volume of 11.6 g of butane (C4H10) gas at RTP?

 $M_r: (4 \times 12) + (10 \times 1) = 58$

11.6/58 = 0.20 mol

Volume = 0.20 x 24 = 4.8 dm³

6g of a hydrocarbon gas had a volume of 4.8 dm³. Calculate its molecular mass.

1 mole = 24 dm³, so 4.8/24 = 0.2 mol

 $M_r = 6 / 0.2 = 30$

If 6g = 0.2 mol, 1 mol equals 30 g

Oxidation Is Loss (of electrons) Reduction Is Gain (of electrons) Ionic half equations (HT only) For example: The ionic equation for the reaction Ionic half between iron and copper (II) ions is: equations show Fe + Cu²⁺ → Fe²⁺ + Cu Acids react with some metals to For what happens displacement produce salts and hydrogen. to each of the The half-equation for iron (II) is: reactions reactants during Fe → Fe2+ + 2ereactions The half-equation for copper (II) ions is: Cu²⁺ + 2e⁻ → Cu Oxidation and Acid name Salt name reduction in terms of electrons (HT ONLY) Hydrochloric Chloride acid Neutralisation of acids Sulfuric acid Sulfate and salt production Nitric acid Nitrate

sodium hydroxide + hydrochloric acid → sodium chloride + water calcium carbonate + sulfuric acid -> calcium sulfate, + carbon dioxide + water

Acids can An alkali is a soluble base e.g. metal be hvdroxide. neutralised A base is a substance that Neutralisation by alkalis neutralises an acid e.g. a soluble and bases metal hydroxide or a metal oxide.

| Metals and oxygen | Metals react with oxygen to form metal oxides | magnesium + oxygen → magnesium oxide 2Mg + O ₂ → 2MgO |
|-------------------|---|--|
| Reduction | This is when oxygen is removed from a compound during a reaction | e.g. metal oxides reacting with hydrogen, extracting low reactivity metals |
| Oxidation | This is when oxygen is gained by a compound during a reaction | e.g. metals reacting with oxygen, rusting of iron |

Reactions metal + acid → metal salt with + hydrogen acids

Reactions of acids

and metals

Reactions of

acids

AQA Chemistry

C4: Chemical Changes

Reactivity of

metals

The reactivity series

Metals form

Metal

oxides

magnesium + hydrochloric acid -> magnesium chloride + hydrogen zinc + sulfuric acid → zinc sulfate + hydrogen

Extraction using carbon

Metals less reactive than carbon can be extracted from their oxides by reduction.

For example: zinc oxide + carbon → zinc + carbon dioxide

Extraction of metals and reduction

Unreactive metals, such as gold, are found in the Earth as the metal itself. They can be mined from the ground.

| | Reactions with water | Reactions with acid |
|-----------------------|--|--|
| Group 1 metals | Reactions get more vigorous as you go down the group | Reactions get more vigorous as you go down the group |
| Group 2 metals | Do not react with water | Observable reactions include fizzing and temperature increases |
| Zinc, iron and copper | Do not react with water | Zinc and iron react slowly with acid. Copper does not react with acid. |

The reactivity series arranges

| positive ions when they react | metal is related to its tendency to form positive ions | metals in order of their reactivity (their tendency to form positive ions). |
|-------------------------------------|---|---|
| Carbon and hydrogen | Carbon and hydrogen are non-metals but are included in the reactivity series | These two non-metals are included in the reactivity series as they can be used to extract some metals from their ores, depending on their reactivity. |
| | A more reactive metal | |

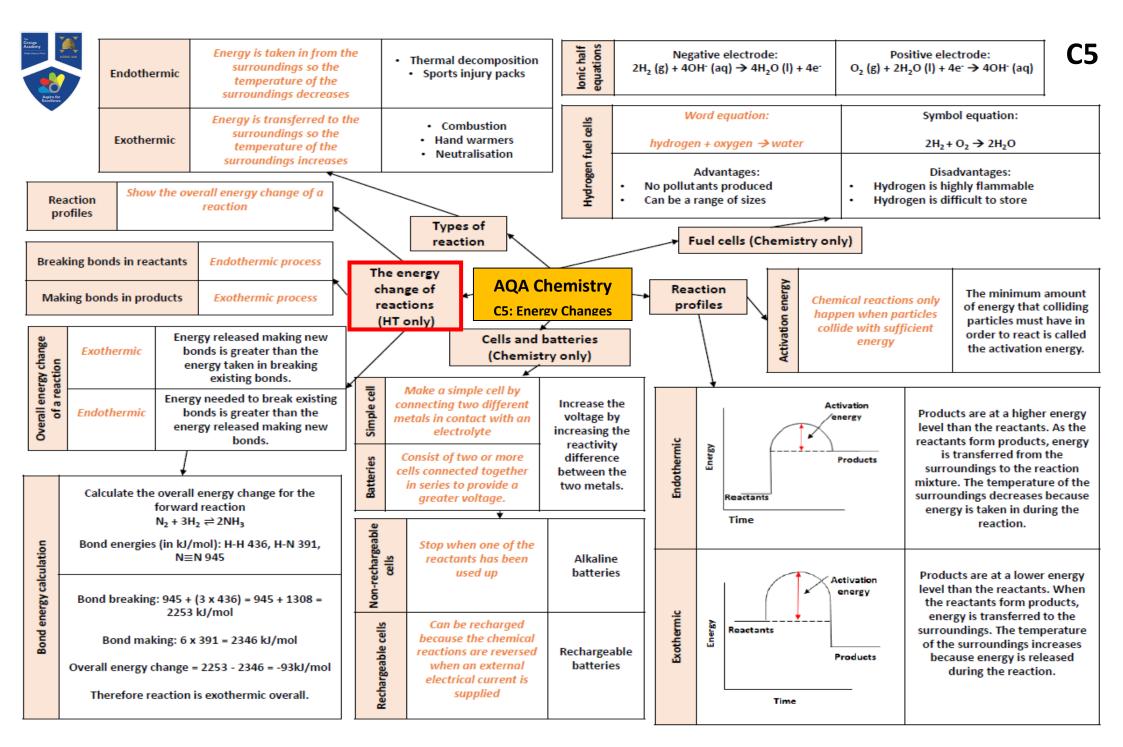
The reactivity of a

A more reactive metal Silver nitrate + Sodium chloride → can displace a less Displacement reactive metal from a Sodium nitrate + Silver chloride compound.

| potassium | most reactive | K |
|-----------|---------------|----|
| sodium | A | Na |
| calcium | | Ca |
| magnesium | | Μį |
| aluminium | | Al |
| carbon | | c |
| zinc | | Zn |
| iron | | Fe |
| tin | | Sn |
| lead | | Pb |
| hydrogen | | н |
| copper | | Cu |
| silver | | Ag |
| gold | | Au |

platinum least reactive Pt

| Change Academy Water Dear No. 2000 April 1980 April 198 | The ions discharged when solution is electrolysed u | ising inert | Process of | Splitting up | water, the | ionic compound is me | These are then able | | | C4 | | |
|--|--|---|--|--------------------------------------|----------------|--|--|-----|-------------------------|---|--|--|
| | electrodes depend on the reactivity of the element | | electrolysis | using electricity | Passing an | duct electricity and are called electrolytes. n electric current though electrolytes causes the ions to move to the electrodes. | | | B | Metals can be extracted from molten compounds using electrolysis. | | |
| At the negative electrode | Metal will be produced on if it is less reactive than Hydrogen will be produced more reactive than h | hydrogen. if the metal is | Electrode | Anode Cathode | The n | positive electrode is ca egative electrode is cal | led the cathode. | | Extracting metals using | This process is used when the metal is too reactive to be extracted by reduction with carbon. | | |
| At the positive electrode | Oxygen is formed at posit If you have a halide ion (C you will get chlorine, brom formed at that elec | ive electrode. l', l', Br') then nine or iodine | Where do the ions go? | Cations Anions | | Cations are positive ions and they move to the negative cathode. Anions are negative ions and they move to the positive anode. | | | Extracting | The process is expensive due to large amounts of energy needed to produce the electrical current. Example: aluminium is extracted in this | | |
| | Electrolysis of aqueous | solutions | (0 | Elect | trolysis | | | | | way. her tier: You can display what is happening | | |
| Strong acids | Completely ionised in a e.g. hydrochloric, nitric | | | | nemistry | Lead Bons ₽b * | Bromide@ons:Bri Molten@ead(iii) bromide | | At t | each electrode using half-equations: the cathode: Pb²+ + 2e⁻ → Pb the anode: 2Br⁻ → Br₂ + 2e⁻ | | |
| Weak acids | Only partially ionised in e.g. ethanoic acid | • | (HT ONLY) | Reactio | _ | | | | | e the pipette to add 25 cm³ of alkali to a conical flask and add a few drops of indicator. | | |
| Hydrogen ion concentration | a stronger acial the hydrogen ion | | | | 15 | Titrations (Chemistry only) | | | | rette with acid and note the starting volume. he acid from the burette to the alkali in the conical flask, swirling to mix. | | |
| Soluble salt | (e.g. metals, meta | nsoluble substanc | ces | Solub the | e precise volu | sed to work out umes of acid and that react with | appı | opr | iate c | g the acid when the end-point is reached (the blour change in the indicator happens). Note me reading. Repeat steps 1 to 3 until you get consistent readings. | | |
| Production of soluble salt | aissoives. Filter off | | then | The | | titrations involv | chemical quantities i | | | The equation shows that 2 mol of NaOH reacts with 1 mol of H ₂ SO ₄ , so the number of moles | | |
| You can use indicator of measure the alkalinity of against the | | | a pH probe t e acidity or a solution | pH scale and utralisation | | (H 2NaOH(aq) + H ₂ 2 It takes 12.20cm ³ of | n³ and in g/dm³ HT ONLY): ₂ SO ₄ (aq) → Na ₂ SO ₄ (aq) + 2H ₂ O(I) of sulfuric acid to neutra n hydroxide solution, wh | | e | in 12.20cm ³ of sulfuric acid is (0.012/2) = 0.006 mol of sulfuric acid Calculate the concentration of sulfuric acid in mol/dm ³ 0.006 mol x (1000/12.2) dm ³ = 0.49mol/dm ³ | | |
| In neutralisatio | n reactions, hydrogen hydroxide ions to | Acids | | roduce hydrogen aqueous solutio | | Calculate the concer | ation of 0.50mol/dm³ ntration of the sulfurion n g/dm³ | aci | a | Calculate the concentration of sulfuric acid in g/dm^3 $H_2SO_4 = (2x1) + 32 + (4x16) = 98g$ | | |
| produce water: | | Alkalis | | ous solutions of in hydroxide ion | | 0.5 mol/dm ³ x (24/1000) dm ³ = 0.012 mol NaOH | | | | 0.49 x 98g = 48.2g/dm ³ | | |





Quantity

Mass

Volume

Rate of

reaction

Rate of chemical reaction

Grams (g)

cm³

Unit

Grams per cm3 (g/cm3)

(mol/s)

Catalyst

How do

they work?

HT: moles per second

This can be calculated by measuring the quantity of reactant used or product formed in a given time.

Rate = quantity of reactant used time taken

| the quantity of | | time taken | 4 | , |
|------------------------------------|--|-------------------------------------|-------------------|---|
| ised or product n a given time. | Rate = <u>quant</u> | ity of product formed time taken | ing rates | |
| _ | | | ן ד | |
| | Calculating | rates of reactions | affe | |
| Volumeiom ² s | tope of tangent = $\frac{25 \text{ cm}^3}{60 \text{ s}}$ | | Factors affecting | |
| 80 80 70 | - 0.42 om² s - 1 | Rate of | <u>"</u> | |
| 50 00 00 1 | | reaction | | |
| 20 | (0) | | | |

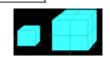
Catalysts

| Factors affecting the rate of reaction | | | | | | |
|--|--|--|--|--|--|--|
| Temperature | The higher the temperature, the quicker the rate of reaction. | | | | | |
| Concentration | The higher the concentration, the quicker the rate of reaction. | | | | | |
| Surface area | The larger the surface area of a reactant solid, the quicker the rate of reaction. | | | | | |
| Pressure (of gases) | When gases react, the higher the pressure upon them, the quicker the rate of reaction. | | | | | |

Collision theory and activation energy







| 1 | | | ACTUATION THEORY |
|---------|----------|---------------------------------------|--|
| ENERGY. | RENTANTS | ACTIVITION DATE WITH GEALINT | ACTIVITIES DESIGN HTTSO/T GENEL/SIT |
| 2 | | PROBUCTS | <u>. </u> |
| | TINE | | |

If a catalyst is used in a reaction, it is not shown in the word equation.

A catalyst changes the rate of a chemical reaction but is not used in the reaction.

These are biological Enzymes catalysts.

> Catalysts provide a different reaction pathway where reactants do not require as much energy to react when they collide.

AQA Chemistry

C6: The Rate and extent of chemical change

Reversible reactions and dynamic equilibrium

Chemical reactions can only occur when reacting Collision theory particles collide with each

other with sufficient energy. This is the minimum

amount of energy colliding particles in a reaction need in order to react.

Increasing the temperature increases the frequency of collisions and makes the collisions more energetic, therefore increasing the rate of reaction.

Increasing the concentration, pressure (gases) and surface area (solids) of reactions increases the frequency of collisions, therefore increasing the rate of reaction.

Reversible reactions

| Reversible reactions | products can react again to re-form the reactants. |
|-----------------------------------|---|
| Representing reversible reactions | A + B = C + D |
| The direction | The direction of reversible reactions can be changed by changing conditions: heat A + B |

Energy changes and reversible reactions

If one direction of a reversible reaction is exothermic, the opposite direction is endothermic. The same amount of energy is transferred in each case.

Changing conditions and equilibrium (HT)

The relative amounts of reactants and products at equilibrium depend on the conditions of the reaction.

Equilibrium in reversible reactions

Equilibrium

When a reversible reaction occurs in apparatus which prevents the escape of reactants and products, equilibrium is reached when the forward and reverse reactions occur exactly at the same rate.

For example: Hydrated copper sulfate exothermic

Anhydrous copper + Water sulfate

Le Chatelier's Principles

Activation

energy

States that when a system experiences a disturbance (change in condition), it will respond to restore a new equilibrium state.

Changing concentration

If the concentration of a reactant is increased, more products will be formed.

If the concentration of a product is decreased, more reactants will react.

Changing temperature

If the temperature of a system at equilibrium is increased:

- Exothermic reaction = products decrease
 - Endothermic reaction = products increase

Changing pressure (gaseous reactions)

For a gaseous system at equilibrium:

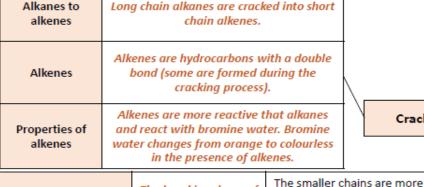
- Pressure increase = equilibrium position shifts to side of equation with smaller number of molecules.
- Pressure decrease = equilibrium position shifts to side of equation with larger number of molecules.

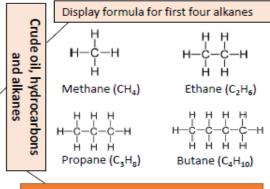
Butane

& Propane

Kerosene

| Excellence | | | | |
|-----------------------------|---|--|---|-------------|
| Crude oil | A finite resource | Consisting mainly of plankton that was buried in the mud, crude oil is the remains of ancient biomass. | | and alkanes |
| Hydrocarbons | These make up the majority of the compounds in crude oil | Most of these hydrocarbons are called alkanes. | | |
| | | For example: | 1 | |
| General formula for alkanes | C_nH_{2n+2} | C ₂ H ₆ | | |
| | | C ₆ H ₁₄ | | |
| | | | | |
| Alkanes to alkenes | | are cracked into short alkenes. | | |

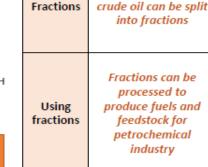




Carbon compounds as fuels

and feedstock

AQA Chemistry



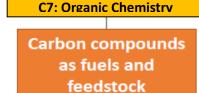
Fractional distillation and

The hydrocarbons in

| olit | number of carbon atoms in them. The process used to do this is called fractional distillation. |
|---------|---|
| e nd | We depend on many of these fuels; petrol, diesel and kerosene. |
| | Many useful materials are made by the petrochemical industry; solvents, lubricants and polymers. |

Each fraction contains

molecules with a similar



chains?

Cracking and alkenes

petrochemicals ln oil Hydrocarbon chains in crude oil come in lots of different lengths. points The boiling point of the chain depends on its length. During fractional distillation, they boil and Boiling separate at different temperatures due to this. Properties of hydrocarbons

During the complete combustion of hydrocarbons, the carbon and hydrogen in the fuels are oxidised, releasing carbon dioxide, water and energy.

| oil, |
|------|
| |
| |

150°C

r r r r

200 °C

| Cracking | Ihe breaking down of long chain hydrocarbons into smaller chains | useful. Cracking can be done by various methods including catalytic cracking and steam cracking. | | | | |
|--------------------|---|---|--|--|--|--|
| Catalytic cracking | The heavy fraction is heated until vaporised | After vaporisation, the vapour is passed over a hot catalyst forming smaller, more useful hydrocarbons. | | | | |
| Steam cracking | The heavy fraction is heated until vaporised | After vaporisation, the vapour is mixed with steam and heated to a very high temperature forming smaller, more useful hydrocarbons. | | | | |

Used to produce polymers. Alkenes They are also used as the and uses starting materials of many other chemicals, such as polymers alcohol, plastics and detergents. Without cracking, many of the Why do long hydrocarbons would be we crack wasted as there is not much long demand for these as for the

shorter chains.

Combustion

Decane → pentane + propene + ethane

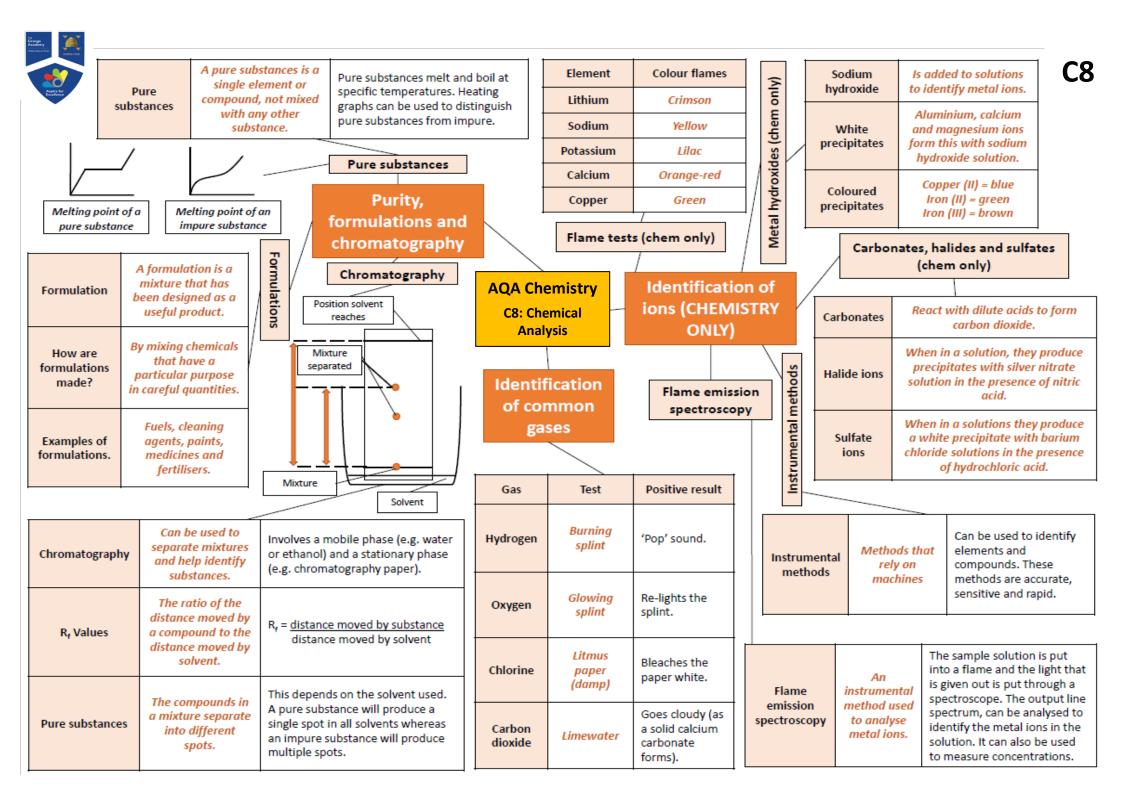
 $C_{10}H_{22} \rightarrow C_5H_{12} + C_3H_6 + C_2H_4$

Hydrocarbon chains

Complete combustion of methane: Methane + oxygen → carbon dioxide + water + energy $CH_4(g) + 2O_2(g) \rightarrow CO_2(g) + 2 H_2O(I)$

| Boiling point (temperature which liquid bo | |
|--|--|
| Viscosity (how easily it flo | As the hydrocarbon chain length increases, viscosity increases. |
| Flammability | As the hydrocarbon chain length increases, flammability decreases. |

| Grange Academy west Descen has | | | | | | | | | | | | |
|--|--|--|---|----------------------------------|-------------------------|--------------------------------------|--|---------------------------------|------|---|--|---|
| American ; | Alken | | rocarbons with a ble carbon-carbon bond. | Struct | | ctional roup | Alkenes are hydro in the functional C=C. | | 0 | organic compo | al group of an und determine eactions. | н н н н С7 |
| H H Ethene C ₃ H ₄ H C=C-C-H H Propene C ₃ H ₆ | Unsatur | rated unso | Alkenes are aturated because ey contain two r hydrogen atoms an their alkane | Structure and formula of alkenes | | kene ctions | Alkenes react with in the same way hydrocarbons, jus smoky flame d incomplete comb | as other st with a lue to | | hydrogen, v nalogens. The | o react with vater and the C=C bond allov n of other aton | |
| H - C - C - C = C | General for alke | ormula | counterparts. C"H _{2n} | ıla of alkene | | eacti | ons of alkenes | Alcohols | | Functional group | -OH For example CH₃CH₂OH | _ |
| Pentene C ₂ H ₂₀ | С—н Н | | | Š | | | AQA Chemistry | | | | | Alcohols and sodium: bubbling, hydrogen gas given off and salt formed. |
| Functional group | -COOH For example: CH₃COOH | propanoic ac | ic acid, ethanoic acid cid and butanoic acid our of the homologou series. | are | Carboxylic acids | | 7: Organic Chemistr | | | Alcohol reactions | Alcohols rea with sodium air and wate | alcohols burn in air releasing carbon |
| | Carboxylic acids and carbonates: These acids are neutralised by carbonates | | | | | thetic and natur ccurring polymer | | | | | Alcohols and water: alcohols dissolve in water to form a neutral solution. | |
| Carboxylic acid reactions | acids react with carbonates, water and alcohols. | These aci Carboxyli | rlic acids and water: ids dissolve in water ic acids and alcohols act with alcohols to f esters. | : | Addition polymerisation | Condensation polymerisat | Amino acids Amino acids have tw functional groups in | a a | Fe | ermentation | Ethanol is produced from fermentatio | produced. The conditions needed for this process include a moderate temperature |
| Strength (HT only) | Carboxylic acids are weak acids | An aqueous | ids only partially ion water. solution of a weak a | acid | lymerisati | n polymer | molecule. They reac by condensation polymerisation to produce peptides. | urally occu | | DNA | DNA gives the | eic acid is a large molecule essential for life. genetic instructions to ensure development g of living organisms and viruses. |
| | | | igh pH (but still belo | / | | isation (I | H H O | rring po | | Structure | | ecules are two polymer chains made from four omers, called nucleotides. They are in the rmation. |
| Polymers | polymers l | used to make by addition risation. | Many small mo together to form large mol | polyme | rs (very | ion (HT only) | H H O-H | lymers | | | | y occurring polymers include proteins, starch and are all important for life. |
| Displaying polymers | repeating u | polymers, the unit has the oms as the omer. | It can be displain $n \begin{bmatrix} H & H \\ C & -C \\ H & H \end{bmatrix} \underbrace{\text{polymeris}}_{\text{ethene}}$ | sation (H | this: | 11 | ndensation ymerisation | | mone | n polymerisati omers with tw oups | vo | When these types of monomers react they join ogether and usually lose small molecules, such as water. This is why they are called condensation reactions. |





| Gas | Percentage | |
|-------------------|------------|---|
| Nitrogen | ~80% | |
| Oxygen | ~20% | L |
| Argon | 0.93% | ĺ |
| Carbon dioxide | 0.04% | |

| | atmosphere | gases in the | Proportions of |
|--|------------|--------------|-----------------------|
|--|------------|--------------|-----------------------|

| Algae and plants | These produced the oxygen that is now in the atmosphere, through photosynthesis. |
|------------------|--|
| | |

First produced by algae 2.7 billion

years ago.

Reducing carbon

dioxide in the

atmosphere

Formation of

sedimentary rocks

and fossil fuels

carbon dioxide + water → glucose + oxygen 6CO, +6H,O \rightarrow C₆H₁,O₆+6O,

Over the next billion years plants evolved to gradually produce more oxygen. This gradually increased to a level that enabled animals to evolve.

| , | | | | |
|---|---|---|--|--|
| Volcano activity 1st Billion years | Billions of years ago there was intense volcanic activity | This released gases (mainly CO ₂) that formed to early atmosphere and water vapour that condensed to form the oceans. | | |
| Other gases | Released from volcanic eruptions | Nitrogen was also released, gradually building up in the atmosphere. Small proportions of ammonia and methane also produced. | | |
| Reducing carbon dioxide in the atmosphere | When the oceans formed, carbon dioxide dissolved into it | This formed carbonate precipitates, forming sediments. This reduced the levels of carbon dioxide in the atmosphere. | | |

How oxygen increased How carbon Earth's early atmosphere dioxide decreased

Composition and evolution of the atmosphere

Oxygen in the

atmosphere

AQA Chemistry

C9: Chemistry of the atmosphere

Common atmospheric

pollutants

CO₂ and methane as greenhouse gases

Global climate

change

Carbon footprints

The total amount of greenhouse gases emitted over the full life cycle of a product/event. This can be reduced by reducing emissions of carbon dioxide and methane.

Algae and plants levels in the atmosphere by absorbing it for photosynthesis. Remains of biological matter falls to the These are made out of the remains

bottom of oceans. Over millions of years layers of sediment settled on top of them of biological and the huge pressures turned them into matter, formed coal, oil, natural gas and sedimentary rocks. over millions of The sedimentary rocks contain carbon years dioxide from the biological matter.

These gradually reduced the carbon dioxide

Greenhouse gases

| Carbon dioxide, | Examples of greenhouse gases that |
|--------------------------|--|
| water vapour | maintain temperatures on Earth in |
| and methane | order to support life |
| The greenhouse effect | Radiation from the Sun enters the Earth's atmosphere and reflects off of the Earth. Some of this radiation is re-radiated back by the atmosphere to the Earth, warming up the global temperature. |

Atmospheric pollutants from fuels

| Combustion of fuels | pollutants. Most fuels may also contain some sulfur. | |
|-----------------------------|---|--|
| Gases from burning fuels | Carbon dioxide, water vapour, carbon monoxide, sulfur dioxide and oxides of nitrogen. | |
| Particulates | Solid particles and unburned hydrocarbons released when burning fuels. | |

Properties and effects of atmospheric pollutants

Toxic, colourless and odourless Carbon monoxide gas. Not easily detected, can kill. Sulfur Cause respiratory problems in dioxide and humans and acid rain which oxides of affects the environment. nitrogen Cause global dimming and health **Particulates** problems in humans.

Human activities and greenhouse gases

| Effects of climate change | Carbon dioxide | Human activities that increase carbon dioxide levels include burning fossil fuels and deforestation. | |
|--|-------------------|--|--|
| Rising sea levels | | | |
| Extreme weather events such as severe storms | Methane | Human activities that increase methane levels include raising livestock (for food) and using landfills (the decay of organic matter released methane). | |
| Change in amount and | | | |
| distribution of rainfall | Climate change | There is evidence to suggest that human activities will cause the Earth's atmospheric temperature to increase and cause climate change. | |
| Changes to distribution of wildlife species with some becoming extinct | | | |



| Corrosion | The destruction of materials by chemical reactions with substances in the environment | An example of this is iron rusting; iron reacts with oxygen from the air to form iron oxide (rust) water needs to be present for iron to rust. |
|--------------------------|---|--|
| Preventing corrosion | Coatings can be added to metals to act as a barrier | Examples of this are greasing, painting and electroplating. Aluminium has an oxide coating that protects the metal from further corrosion. |
| Sacrificial corrosion | When a more reactive metal is used to coat a less reactive metal | This means that the coating will react with the air and not the underlying metal. An example of this is zinc used to galvanise iron. |

Formulations of various These contain salts containing appropriate NPK nitrogen, fertilisers percentages of the phosphorous and potassium elements. Phosphate rock needs to be **Potassium** treated with an acid to chloride, produce a soluble salt potassium Fertiliser which is then used as a sulfate and examples fertiliser. Ammonia can be phosphate rock used to manufacture are obtained ammonium salts and nitric by mining acid.

Production and uses of NPK fertilisers

A mixture of two elements, one of which must be a metal e.g. Bronze is an alloy of copper and tin and Brass is an alloy of copper and zinc.

Gold jewellery is usually an alloy with silver, copper and zinc. The carat of the jewellery is a measure of the amount of gold in it e.g. 18 carat is 75% gold, 24 carat is 100% gold.

Alloys of iron, carbon and other metals.

High carbon steel is strong but brittle.

Low carbon steel is softer and easily shaped.

Steel containing chromium and nickel (stainless) are hard and corrosion resistant.

Aluminium alloys are low density.

Ceramics, polymers and composites

The Haber process

Polymers

Thermosetting polymers that do not melt when they are heated.

Thermosoftening polymers that melt when they are heated.

Using materials

Alloys

Gold

Steels

Corrosion and its prevention

Alloys are useful materials

AQA Chemistry

C10: Using Resources

The Haber process and the use of NPK fertilisers

| | | A mixture of materials put together for a specific purpose e.g. strength | Soda-lime glass, made by heating sand, sodium carbonate and limestone. | |
|--|------------------------|--|---|--|
| | Composite materials | | Borosilicate glass, made from sand and boron trioxide, melts at higher temperatures than soda-lime glass. | |
| | | | MDF wood (woodchips, shavings, sawdust and resin) | |
| | | | Concrete (cement, sand and gravel) | |
| | Ceramic materials | Made from clay | Made by shaping wet clay and then heating in a furnace, common examples include pottery and bricks. | |
| | Polymers | Many monomers can make polymers | These factors affect the properties of the polymer. Low density (LD) polymers and high density (HD) polymers are produced from ethene. These are formed under different conditions. | |

| Phosphate rock | | |
|--------------------|---|--|
| Treatment | Products | |
| Nitric acid | The acid is neutralised with ammonia to produce ammonium phosphate, a NPK fertiliser. | |
| Sulfuric acid | Calcium phosphate and calcium sulfate (a single superphosphate). | |
| Phosphoric acid | Calcium phosphate (a triple superphosphate). | |

| The Haber process – conditions and equilibrium | | | |
|--|--|--|--|
| Pressure | ammonia (Le Chatelier's principle). Th pressure needs to be as high as possible. | | |
| Temperature | The forward reaction is exothermic. Decreasing temperature increases ammonia production at equilibrium. The exothermic reaction that occurs releases energy to surrounding, opposing the temperature decreases. Too low though and collisions would be too infrequent to be financially viable. | | |

| The Haber process | Used to manufacture ammonia | Ammonia is used to produce fertilisers Nitrogen + hydrogen = ammonia |
|-------------------|---|--|
| Raw materials | Nitrogen from the air while hydrogen from natural gas | Both of these gases are purified before being passed over an iron catalyst. This is completed under high temperature (about 450°C) and pressure (about 200 atmospheres). |
| Catalyst | Iron | The catalyst speeds up both directions of the reaction, therefore not actually increasing the amount of valuable product. |