

WJEC Chemistry AS-Level

2.6: Halogenoalkanes

Detailed Notes

English Specification

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Elimination Reactions

When a halogenoalkane is mixed with hydroxide ions and heated to **high temperatures** under **alcoholic** conditions, **elimination** occurs. In this reaction, a hydroxide ion **nucleophile** acts as a base and accepts a proton, removing a hydrogen atom from the molecule. This results in the elimination of the **halide** too, producing a **carbon-carbon double bond**.

Mechanism - elimination

Nucleophilic Substitution

Halogenoalkanes contain **polar bonds** as the halogens are much **more electronegative** than carbon atoms. This means electron density is drawn towards the halogen, forming ∂ + and ∂ -regions.

Example:

In the reaction mechanism of nucleophilic substitution, nucleophiles attack the halogenoalkane and displace the halogen to produce compounds such as alcohols and amines.







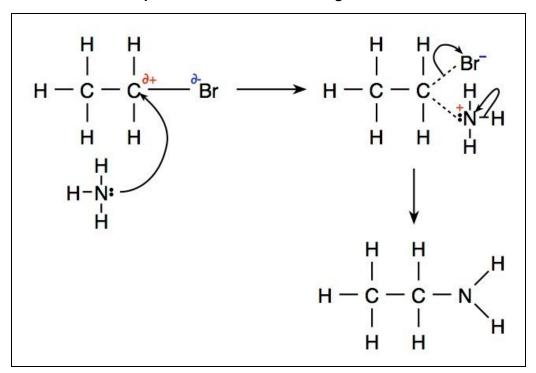




Mechanism - nucleophilic substitution of a halogenoalkane to form an alcohol

The nucleophile attacks the ∂+ carbon and the electrons are transferred to the chlorine.

Mechanism - nucleophilic substitution of a halogenoalkane to form an amine



The intermediate formed has an N⁺ atom, so electrons are transferred to it causing a hydrogen to be lost.

The greater the relative atomic mass of the halogen in the polar bond, the lower the bond enthalpy. The lower the bond enthalpy, the more easily the bond can be broken. Therefore, the higher the relative atomic mass of the halogen atom, the faster the rate of reaction for the halogenoalkane.







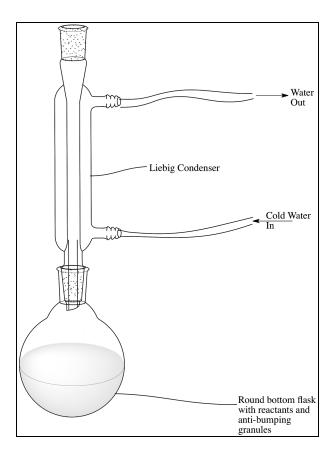




Reflux

The process of nucleophilic substitution has to be carried out under **reflux conditions** in a **closed system**. In reflux, the vapours of the reaction evaporate, condense and return to the original solution to be heated again.

There is specific **reflux equipment** that can be used to carry out this process, including a **Liebig condenser** and **anti-bumping granules** that help to disperse the heat evenly through the solution.



Testing for Halogenoalkanes

To test for halogenoalkanes, the unknown solution should be warmed with a mixture of sodium hydroxide, ethanol and water. This is a hydrolysis reaction that causes the displacement of the halide ion. After the halide ion has been displaced, dilute nitric acid is added to the solution to neutralise any excess sodium hydroxide. The type of halide ion present can then be identified specifically by using silver nitrate solution. The Ag⁺ ions from silver nitrate react with the halide to produce coloured precipitates.











The precipitates formed may not be clear to distinguish so they can be tested further using ammonia.

	CI ⁻	Br ⁻	l'
+ AgNO ₃	White precipitate (AgCl)	Cream precipitate (AgBr)	Yellow Precipitate (AgI)
+ dilute NH ₃	Precipitate dissolves	No Change	No Change
+ conc. NH ₃	Precipitate dissolves	Precipitate dissolves	No Change

Uses of Halogenoalkanes

CFCs (chlorofluorocarbons) are halogenoalkane molecules where all of the hydrogen atoms have been replaced by fluorine and chlorine atoms. Example:

They are used for refrigerants, foamed plastic production, dry cleaning solvents and aerosols due to their low flammability and toxicity.

However, once released into the atmosphere, CFCs absorb UV radiation causing them to break down to form free radicals. These free radicals are very harmful to the environment because they can catalyse ozone depletion. Example:

$$O_3 + Cl \cdot \longrightarrow \cdot OCl + O_2$$
 $\cdot OCl + O_3 \longrightarrow 2O_2 + Cl \cdot \bigcirc$
 $Overall: 2O_3 \longrightarrow 3O_2$

Ozone, O₃, is in the upper atmosphere and helps to prevent ultraviolet radiation from the sun from reaching the earth. Ultraviolet radiation causes sunburn and skin cancer so ozone is a very important layer of protection.











Ozone is formed from oxygen free radicals:

$$O_2 \longrightarrow O_4 + O_4$$
 $O_2 + O_4 \longrightarrow O_3$

CFC-free solvents are now being produced to prevent the damage to the ozone layer.







