

# **WJEC Chemistry A-Level**

OA1.2: Aromaticity

**Detailed Notes** 

**English Specification** 

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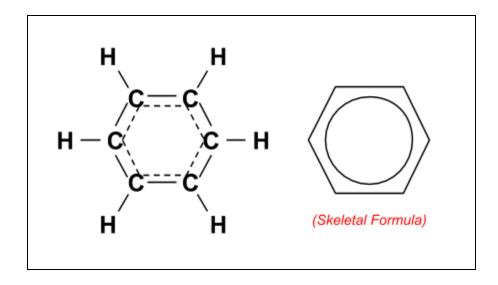
# **Aromatic Bonding**

The structure and bonding of aromatic compounds is pretty unique due to the behaviour of delocalised electrons within the compound.

## Benzene

Benzene is an example of an aromatic compound, consisting of a ring of six carbon atoms with six hydrogen atoms and a ring of delocalised electrons.

## Example:



Each carbon-carbon bond in the benzene ring has an intermediate length between that of a double and single bond.

The outer electron from the p-orbital of each carbon atom becomes delocalised to form a central ring of delocalised electrons. This ring structure makes benzene very stable compared to other molecules of a similar size.

## Arenes

Compounds that contain benzene as part of their structure are called arenes or aromatic compounds. They have high melting points due to the high stability of the delocalised ring, but low solubility as they are nonpolar molecules, meaning they generally cannot be dissolved in water.

This low polarity and high stability also means aromatic compounds resist addition reactions, as it would require the stable delocalised ring to be broken. Arenes tend to react via substitution mechanisms instead.









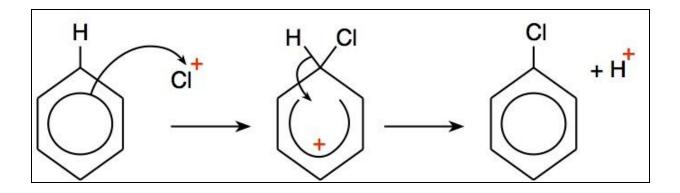
# **Electrophilic Substitution**

The delocalised ring in benzene is an area of **high electron density** making it susceptible to **attack from electrophiles**. When these species attack the electron ring, it is **partially destroyed**. It is then restored in the process of **electrophilic substitution**.

# Halogenation of Benzene

In this mechanism, an unusual and highly unstable form of halide ion with a +1 charge acts as an electrophile, attacking the electron ring. A halogen carrier catalyst is used to generate the positively charged halide ion. As a result, a halobenzene compound is formed.

#### Mechanism



## **Nitration of Benzene**

In this form of electrophilic substitution, the electrophile is the  $NO_2^+$  ion. This is a reactive intermediate, produced in the reaction of concentrated sulfuric acid  $(H_2SO_4)$  with concentrated nitric acid  $(HNO_3)$ .

Example:

$$H_2SO_4 + HNO_3 \longrightarrow H_2NO_3^+ + HSO_4^ H_2NO_3^+ \longrightarrow H_2O + NO_2^+$$

When heated with benzene the combination of these reagents lead to the substitution of the NO<sub>2</sub><sup>+</sup> electrophile onto the benzene ring, releasing a hydrogen ion. This hydrogen ion reacts with the HSO<sub>4</sub><sup>-</sup> ion from the first reaction, producing sulfuric acid. This shows how sulfuric acid is a catalyst for this reaction since it does not get used up.



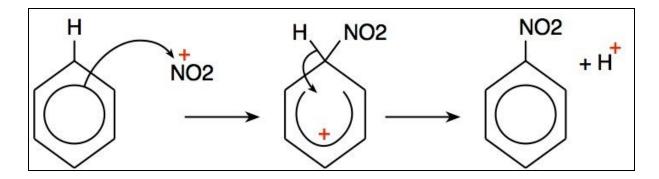








# Mechanism - electrophilic substitution



This reaction shows a mono-substitution, where a single NO<sub>2</sub><sup>+</sup> electrophile is substituted under conditions of 55°C. At temperatures greater than this, multiple substitutions can occur. It is vital that only one substitution occurs for the production of aromatic amines.

# Friedel-Crafts Alkylation

The delocalised electron ring in benzene can also act as a nucleophile, leading to an attack from acyl chlorides. This reaction is known as Friedel-Crafts alkylation.

In order for this to occur, a reactive intermediate must be produced from the **acyl chloride** and an **aluminium chloride catalyst**.

Example:

$$R\text{-COCI} + AICI_3$$
  $\longrightarrow$   $R\text{-CO}^+ + AICI_4^-$ 

This reactive intermediate is then attacked by the electron dense benzene ring.

#### Mechanism









At the end of the reaction, the H<sup>+</sup> ion removed from the ring reacts with the AlCl<sub>4</sub>- ion from the formation of the reactive intermediate to reform aluminium chloride. This shows how AlCl<sub>3</sub> is a catalyst for the reaction because it reforms at the end of the reaction.

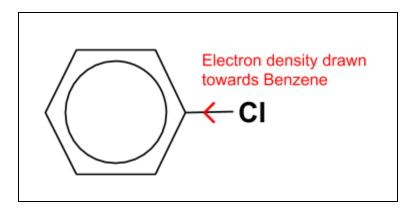
The product of this reaction is a **phenylketone**. In this case, the benzene group is called a **phenyl group**. These molecules are commonly used in the industrial production of dyes, pharmaceuticals and explosives.

# **The Negative Inductive Effect**

In organic molecules, different functional groups can **affect bond strengths** within the molecule by **changing electron density** around the bond.

Benzene rings draw electron density towards themselves, away from any substituent group such as a halogen atom. This is called the negative inductive effect. In particular, in the following example, there is some overlap between p-electrons on the chlorine and the delocalised  $\pi$ -system in the benzene ring. This gives some  $\pi$ -character to the bond, which strengthens it.

## Example:



As a result, the bond strength between Benzene and the halogen atom is **increased**, making it less likely for a reaction to occur as the molecule is **more stable**.







