

CAIE Chemistry A-level

36: Organic Synthesis

(A-level only)

Notes

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Synthetic Routes

Synthetic routes are the routes which can be used to produce a **certain product from a starting organic compound**. It is important that you understand the different methods and **conditions** required to convert compounds to other products.

Below is a table showing the typical reactions of different functional groups and how they can be identified. This table contains both reactions from AS and the A2 course.

| Homologous series | Typical reactions | Identification |
|--|---|---|
| Alkanes C-C | Combustion Electrophilic substitution/ free radical substitution with Br ₂ or Cl ₂ (forms haloalkanes) Cracking (forms short chain alkenes and alkanes) | |
| Alkenes C=C | Electrophilic addition: - Steam (forms alcohols) - Hydrogen halides (forms haloalkanes) - Halogens (forms dihaloalkanes) - Hydrogen (forms alkanes) Oxidation with H ⁺ /MnO ₄ (forms diols) Addition polymerisation (forms polymers) | React with bromine water: Decolorises in the presence of C=C. |
| Halogenoalkanes C-F/ C-Cl/ C-Br/ C-I | Combustion Nucleophilic substitution: - Hydrolysis (forms alcohols) - Reaction with ethanolic cyanide (forms nitriles) - Reaction with ammonia (forms primary amines) Elimination of hydrogen halide using ethanolic hydroxide ions (forms alkenes) | React with AgNO ₃ (aq), test precipitate with NH ₃ (aq): AgCl - white ppt soluble in dilute NH ₃ (aq) AgBr - cream ppt soluble in concentrated NH ₃ (aq) AgI - yellow ppt insoluble in NH ₃ (aq) |
| Alcohols -OH | Combustion Substitution with hydrogen halides, sulfur dichloride oxide or phosphorus(III) halides (forms haloalkanes) Ethanol and sodium (forms sodium ethoxide and hydrogen gas) Oxidation with H ⁺ /Cr ₂ O ₇ ²⁻ (forms carbonyls and carboxylic acids) Dehydration using an acid catalyst (forms alkenes) Esterification with carboxylic acids or acyl chlorides | React with H ⁺ /Cr ₂ O ₇ ²⁻ : Colour change from orange to green in the presence of primary and secondary alcohols (no change for tertiary alcohols) |











| Aldehydes | Oxidation with H ⁺ /Cr ₂ O ₇ ²⁻ (forms carboxylic acids) | React with 2,4-DNPH: A |
|------------------|--|---|
| -CHO | Reduction using NaBH ₄ or LiAlH ₄ (forms primary | yellow-orange precipitate is |
| | alcohols) | formed in the presence of a |
| | <i>Nucleophilic addition</i> with HCN (forms hydroxynitriles) | carbonyl group. |
| | | React with Tollens' reagent: |
| | | A silver mirror is produced if |
| | | an aldehyde is present. |
| | | , |
| | | React with Fehling's |
| | | reagent: The blue solution forms a brick red precipitate in |
| | | · · · |
| | | the presence of an aldehyde |
| | | React with acidified |
| | | potassium dichromate(VI): |
| | | Orange solution turns green |
| | | |
| Ketones | Reduction using NaBH ₄ or LiAlH ₄ (forms secondary | React with 2,4-DNPH: A |
| RCOR' | alcohols) | yellow-orange precipitate is |
| | <i>Nucleophilic addition</i> with HCN (forms hydroxynitriles) | formed in the presence of a |
| | (| carbonyl group. |
| Carboxylic acids | Reaction with metals, alkalis or carbonates (forms a | Test pH: pH less than 7 when |
| -COOH | salt and inorganic products) | measured using a pH probe |
| | Esterification with alcohols | React with a carbonate: |
| | Reduction with LiAlH₄ (forms alcohols) | effervescence as CO ₂ is |
| | | formed |
| | Reaction with SOCl ₂ (forms acyl chlorides, sulfur | |
| | dioxide and hydrochloric acid) | |
| | Reaction with phosphorus(V) chloride or | |
| | phosphorus(III) chloride (forms acyl chlorides) | |
| | Oxidation of methanoic acid using Fehling's or Tollens' | |
| | (forms carbon dioxide and water) | |
| | Oxidation of ethanedioic acid using acidified | |
| | potassium manganate(VII) (forms water and carbon | |
| | dioxide) | |
| Esters | Acid hydrolysis (forms a carboxylic acid and an | |
| RCOOR' | alcohol) | |
| | , | |
| | Alkali hydrolysis (forms a carboxylate salt and an alcohol) | |
| A • | , | |
| Amines | Reaction with acids (forms a salt) | |
| -NH ₂ | | |
| Nitriles | Acid hydrolysis (forms a carboxylic acid and a salt) | |
| C≣N | Alkali hydrolysis (forms a carboxylate salt and | |
| | ammonia) | |
| | | |











| Arenes | Electrophilic substitution: | |
|--------------------------------|---|--|
| -C ₆ H ₅ | - Halogen (forms chlorobenzene with Cl₂ and | |
| | bromobenzene with Br ₂) | |
| | - Nitration (forms nitrobenzene) | |
| | Friedel-Crafts acylation and alkylation | |
| | Oxidation of a side chain (forms benzoic acid) | |
| | Hydrogenation (forms cyclohexane) | |
| Phenol | Reactions with strong bases (not acidic enough to | |
| C ₆ H₅OH | react with carbonates) | |
| | Reaction with sodium (forms sodium phenoxide and hydrogen gas) | |
| | Reaction with diazonium salts (forms azo compounds) | |
| | Electrophilic substitution: | |
| | - Nitration using HNO ₃ (forms nitrophenol) | |
| | - Bromination using Br ₂ (forms bromophenol) | |
| Acyl chlorides | Hydrolysis with water (forms carboxylic acids and HCl) | |
| -COCI | Hydrolysis with sodium hydroxide (forms a carboxylate salt and water) | |
| | Esterification with alcohols or phenol | |
| | Reaction with ammonia (forms an amide and HCI) | |
| | Reactions with primary amines (forms an N-substituted amide) | |
| Amides | Acid hydrolysis (forms a carboxylic acid and | |
| -CONH₂ | ammonium ions) | |
| | Alkali hydrolysis (forms a carboxylate salt and ammonia or an amine) | |
| | Reduction using LiAlH₄ (forms a primary amine) | |

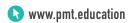
Multi-Stage Synthesis

Some organic molecules can be prepared using a **multi-stage synthesis**. Typically, this involves two stages: reactant \rightarrow intermediate \rightarrow product. It can cover more stages.

Example 1

Below is a diagram showing how ethanoic acid can be formed from chloroethane:











Example 2

2-propylamine can be formed from propene:

Analysing Synthetic Routes

When **synthesising** an organic compound, several factors are considered before deciding which synthetic route to use:

- Type of reaction addition reactions are more sustainable than substitution or elimination reactions as there are no waste products.
- Reagents renewable reagents with few safety concerns are preferred.
- By-products less harmful by-products are favoured as there would be fewer safety
 and environmental concerns. If the by-products can be used in another industry, the
 process is more sustainable.
- Conditions choose the reaction with the most energy efficient and safe conditions.







