

Organic Chemistry

Large number of compounds due to:

- 4 valence pairs
- single / double / triple bonds
- cyclic (ring) structures

Properties of hydrocarbons

- *Saturated* - all C-C bonds are single
- Insoluble in water
- Almost non-polar (similar electronegativities)
- Only dispersion forces (valence e-)
- Dispersion forces increase with length
- Branched molecules have lower density

Linear (aliphatic)

Alkanes: $C_n H_{2n+2}$

Alkenes: $C_n H_{2n}$

Alkynes: $C_n H_{2n-2}$

Naming hydrocarbons

- Branches end with *-yl*
- Indicate number of branches with di-, tri- etc.
- Longest unbranched carbon chain includes function group

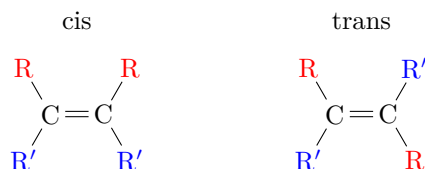
Functional groups

Alcohols	-OH	$R-OH$
Esters	-OCO-	$R-C \begin{matrix} \diagup O-R \\ \diagdown O \end{matrix}$
Aldehydes	-CHO	$R-C \begin{matrix} \diagup O \\ \diagdown H \end{matrix}$
Ketones	-CO-	$R \begin{matrix} \diagdown \\ \diagup \end{matrix} C=O$
Carboxylic acids	-COOH	$R-C \begin{matrix} \diagup O-H \\ \diagdown O \end{matrix}$
Ethers	-O-	$O \begin{matrix} \diagup R \\ \diagdown R \end{matrix}$
Amines	-NH ₂	$R-N \begin{matrix} \diagup H \\ \diagdown H \end{matrix}$
Amides	CO-NH	$R-C \begin{matrix} \diagup O \\ \diagdown N \begin{matrix} \diagup H \\ \diagdown H \end{matrix} \end{matrix}$

Isomers

- **Structural isomers** - same molecular formula, different arrangement
- **Positional isomers** - different position of functional group (affects polarity)

- **Chain isomers** - affects boiling point (more branches *implies* lower boiling point)
- **Stereoisomers** - same structural configuration, different orientation
- **Optical/chiral isomers (enantiomers)** - chiral centre, 4 groups bonded to C, non-superimposable mirror image. Racemic mixture: 50/50 of each enantiomer. Affects optical polarity of light.
- **Geometric isomers** - C=C double bond, 2 groups bonded to carbon atoms
 - **Cis** - same horizontal plane (symmetrical)
 - **Trans** - diagonal (asymmetrical)



Flash point

Lowest temperature at which the liquid gives off enough vapour to be ignited. Flammable \implies flash point $>$ 37.8 °C.

Reactions

Cracking - split molecules with heat/pressure/catalyst

Alkanes

- Relatively inert
- Non-polar solvent
- Non-soluble in H₂O
- Combusts in O₂ (forms CO₂ + H₂O)
- Reacts with halogens (**substitution** of H)

Alkenes

- More reactive than alkanes
- **Addition reactions**: C=C bond is broken (energy released)
- **Addition polymerisation**

Alcohols

- Can be formed from haloalkane substitution reaction
- **Oxidation** (combustion)
- Oxidation state \propto no. of atoms connected to C
- **Substitution** of functional group, e.g. ROH + NH₃ \longrightarrow RNH₂ + H₂O
- Primary alcohols oxidise to aldehydes then carboxylic acids
- Secondary alcohols oxidise to ketones

Carboxylic acids

- Weak acids
- **Hydrolysis**: RCOOH + H₂O \rightleftharpoons RCOO⁻ + H₂O⁺
- Rxn with amines - carboxylic acid + amine \rightarrow ammonium salt \rightarrow amide + water

Esterification

- **Condensation reactions** (esterification): RCOOH + R'OH \longrightarrow RCOOR' + H₂O
- This is reversed by **hydrolysis**: ester + water \rightarrow carboxylic acid + alcohol
- Polyesters

Summary of reactions

Combustion: $C_x H_y + z O_2 \longrightarrow x CO_2 + \frac{y}{2} H_2O$ where $z = x + \frac{y}{4}$

Substitution: e.g. halogenation/chlorination, catalysed by UV (alkanes) or compounds (alcohols etc.) which creates free radicals (unpaired e-)

Addition: C=C bond is broken, catalyst is required

Addition polymerisation: uses addition to form a polymer

Condensation polymerisation: produces H_2O

Oxidation: occurs in alcohols to form aldehydes with catalysts H^+/MnO_4^- (aq) or $H^+/Cr_2O_7^{2-}$ (aq) - see combustion equation

Cracking: heat/pressure/catalyst to break molecules