4.1.2 Alkanes

Alkanes and cycloalkanes are saturated hydrocarbons



carbon atom in saturated hydrocarbons is tetrahedral and the bond angle is 109.5°

The shape is tetrahedral as a result of the four bond pairs of electrons equally repelling.



Boiling Point

The increasing boiling points of the alkane homologous series can be explained by the increasing number of electrons in the bigger molecules causing an increase in the size of the induced dipole–dipole interactions (London forces) between molecules.

The shape of the molecule can also have an effect on the size of the induced dipole–dipole interactions (London forces). Long chain alkanes have a larger surface area of contact between molecules for London force to form than compared to spherical shaped branched alkanes and so have stronger induced dipole–dipole interactions and higher boiling points.

Reactivity

The low reactivity of alkanes with many reagents can be explained by the high bond enthalpies of the C-C and C-H bonds and the very low polarity of the -bonds present.

Hydrocarbons as fuels

Fuel : releases heat energy when burnt

Complete Combustion

In excess oxygen alkanes will burn with complete combustion

The products of <u>complete</u> combustion are CO_2 and H_2O .

 $C_8H_{18}(g)$ + 12.5 $O_2(g) \rightarrow 8CO_2(g)$ + 9 $H_2O(I)$

Incomplete Combustion

If there is a **limited amount of oxygen** then <u>incomplete</u> combustion occurs, producing CO (which is very toxic) and/or C (producing a sooty flame)

 $\begin{array}{rl} {\sf CH}_4(g) \ + \ {}^3\!/_2 \ {\sf O}_2(g) \ \to \ {\sf CO}(g) \ + \ 2 \ {\sf H}_2{\sf O}({\sf I}) \\ {\sf CH}_4(g) \ + \ {\sf O}_2(g) \ \to \ {\sf C}(s) \ + \ 2 \ {\sf H}_2{\sf O}({\sf I}) \end{array}$

Carbon monoxide is an highly toxic but odourless gas. It can cause death if it builds up in a enclosed space due to faulty heating appliances. Alkanes readily burn in the presence of oxygen. This combustion of alkanes is highly exothermic, explaining their use as **fuels**.

Incomplete combustion produces less energy per mole than complete combustion

Carbon (soot) can cause global dimming- reflection of the sun's light

CO is toxic to humans as CO can from a strong bond with haemoglobin in red blood cells. This is a stronger bond than that made with oxygen and so it prevents the oxygen attaching to the haemoglobin.

Cracking

Cracking: conversion of large hydrocarbons to smaller molecules of by breakage of C-C bonds

High Mr alkanes \rightarrow smaller Mr alkanes+ alkenes + (hydrogen)

Economic reasons for catalytic cracking

- The petroleum fractions with shorter C chains (e.g. petrol and naphtha) are in **more demand** than larger fractions.
- To make use of excess larger hydrocarbons and to supply demand for shorter ones, longer hydrocarbons are cracked.
- The products of cracking are **more valuable** than the starting materials (e.g. ethene used to make poly(ethene), branched alkanes for motor fuels, etc.)

This is a chemical process involving the splitting of strong covalent bonds so requires high temperatures.

Catalytic Cracking

Conditions: Slight pressure High Temperature (450°C) Zeolite Catalyst Turns straight chain alkanes into branched and cyclic alkanes and Aromatic hydrocarbons

Used for making motor fuels

Branched and cyclic hydrocarbons burn more cleanly and are used to give fuels a higher octane number

Substitution reactions of alkanes

Synthesis of chloroalkanes

Reaction of alkanes with bromine / chlorine in UV light

In the presence of **UV light** alkanes react with chlorine to form a mixture of products with the halogens substituting hydrogen atoms.



In general, alkanes do not react with many reagents. This is because the C-C bond and the C-H bond are relatively strong

To understand this reaction fully we must look in detail at how it proceeds step by step. This is called its **mechanism**



It proceeds via a series of steps: Step one: initiation Step two: propagation Step three: termination





Propagation steps for substituting a halogen on a 'middle' carbon

CH₃CH₂CH₃ + Br[•] → HBr + CH₃CH[•] CH₃ CH₃CH[•] CH₃ + Br₂ → CH₃CH₂BrCH₃ + Br[•]

If the question asks for the halogen to be substituted onto a middle carbon in the chain, it is important to put the free radical '**dot**' on the correct carbon in the propagation stages.

Further substitution

Excess Cl₂ present will promote further substitution and could produce CH₂Cl₂, CHCl₃ and CCl₄

These reactions could occur $CH_3Cl + Cl_2 \rightarrow CH_2Cl_2 + HCl$ $CH_2Cl_2 + Cl_2 \rightarrow CHCl_3 + HCl$ $CHCl_3 + Cl_2 \rightarrow CCl_4 + HCl$ Example propagation steps that would lead to further substitution $CH_3CI + CI \rightarrow HCI + \cdot CH_2CI$ $\cdot CH_2CI + CI_2 \rightarrow CH_2CI_2 + CI \cdot$

Overall reaction equations

You should be able to write overall reaction equations for various reactions

Example 1. Write the overall reaction equation for the formation of CCI_4 from $CH_4 + CI_2$

$$CH_4 + 4 Cl_2 \rightarrow CCl_4 + 4 HCl$$

Example 2. Write the overall reaction equation for the formation of $CFCI_3$ from $CH_3F + CI_2$

$$CH_3F + 3 Cl_2 \rightarrow CFCl_3 + 3 HCl$$

Note HCl is always the side product - never H₂