



# Hydrocarbons are the important sources of energy.

## **Objectives**

After studying this unit, you will be able to

- name hydrocarbons according to IUPAC system of nomenclature;
- recognise and write structures of isomers of alkanes, alkenes, alkynes and aromatic hydrocarbons;
- learn about various methods of preparation of hydrocarbons;
- distinguish between alkanes, alkenes, alkynes and aromatic hydrocarbons on the basis of physical and chemical properties;
- draw and differentiate between various conformations of ethane;
- appreciate the role of hydrocarbons as sources of energy and for other industrial applications;
- predict the formation of the addition products of unsymmetrical alkenes and alkynes on the basis of electronic mechanism;
- comprehend the structure of benzene, explain aromaticity and understand mechanism of electrophilic substitution reactions of benzene;
- predict the directive influence of substituents in monosubstituted benzene ring;
- learn about carcinogenicity and toxicity.

The term 'hydrocarbon' is self-explanatory which means compounds of carbon and hydrogen only. Hydrocarbons play a key role in our daily life. You must be familiar with the terms 'LPG' and 'CNG' used as fuels. LPG is the abbreviated form of liquified petroleum gas whereas CNG stands for compressed natural gas. Another term 'LNG' (liquified natural gas) is also in news these days. This is also a fuel and is obtained by liquifaction of natural gas. Petrol, diesel and kerosene oil are obtained by the fractional distillation of petroleum found under the earth's crust. Coal gas is obtained by the destructive distillation of coal. Natural gas is found in upper strata during drilling of oil wells. The gas after compression is known as compressed natural gas. LPG is used as a domestic fuel with the least pollution. Kerosene oil is also used as a domestic fuel but it causes some pollution. Automobiles need fuels like petrol, diesel and CNG. Petrol and CNG operated automobiles cause less pollution. All these fuels contain mixture of hydrocarbons, which are sources of energy. Hydrocarbons are also used for the manufacture of polymers like polythene, polypropene, polystyrene etc. Higher hydrocarbons are used as solvents for paints. They are also used as the starting materials for manufacture of many dyes and drugs. Thus, you can well understand the importance of hydrocarbons in your daily life. In this unit, you will learn more about hydrocarbons.

#### 9.1 CLASSIFICATION

Hydrocarbons are of different types. Depending upon the types of carbon-carbon bonds present, they can be classified into three main categories – (i) saturated

(ii) unsaturated and (iii) aromatic hydrocarbons. Saturated hydrocarbons contain carbon-carbon and carbon-hydrogen single bonds. If different carbon atoms are joined together to form open chain of carbon atoms with single bonds, they are termed as alkanes as you have already studied in Unit 8. On the other hand, if carbon atoms form a closed chain or a ring, they are termed as cycloalkanes. Unsaturated hydrocarbons contain carbon-carbon multiple bonds double bonds, triple bonds or both. Aromatic hydrocarbons are a special type of cyclic compounds. You can construct a large number of models of such molecules of both types (open chain and close chain) keeping in mind that carbon is tetravalent and hydrogen is monovalent. For making models of alkanes, you can use toothpicks for bonds and plasticine balls for atoms. For alkenes, alkynes and aromatic hydrocarbons, spring models can be constructed.

#### 9.2 ALKANES

As already mentioned, alkanes are saturated open chain hydrocarbons containing carbon - carbon single bonds. Methane (CH<sub>4</sub>) is the first member of this family. Methane is a gas found in coal mines and marshy places. If you replace one hydrogen atom of methane by carbon and join the required number of hydrogens to satisfy the tetravalence of the other carbon atom, what do you get? You get C<sub>2</sub>H<sub>6</sub>. This hydrocarbon with molecular formula C<sub>2</sub>H<sub>6</sub> is known as ethane. Thus you can consider C<sub>2</sub>H<sub>6</sub> as derived from CH<sub>4</sub> by replacing one hydrogen atom by -CH<sub>3</sub> group. Go on constructing alkanes by doing this theoretical exercise i.e., replacing hydrogen atom by -CH<sub>3</sub> group. The next molecules will be  $C_3H_8$ ,  $C_4H_{10}$ ...

These hydrocarbons are inert under normal conditions as they do not react with acids, bases and other reagents. Hence, they were earlier known as paraffins (*latin : parum*, little; *affinis*, affinity). Can you think

of the general formula for alkane family or **homologous series**? If we examine the formula of different alkanes we find that the general formula for alkanes is  $C_nH_{2n+2}$ . It represents any particular homologue when n is given appropriate value. Can you recall the structure of methane? According to VSEPR theory (Unit 4), methane has a tetrahedral structure (Fig. 9.1), in which carbon atom lies at the centre and the four hydrogen atoms lie at the four corners of a regular tetrahedron. All H-C-H bond angles are of 109.5.

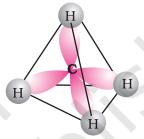


Fig. 9.1 Structure of methane

In alkanes, tetrahedra are joined together in which C-C and C-H bond lengths are 154 pm and 112 pm respectively (Unit 8). You have already read that C-C and C-H  $\sigma$  bonds are formed by head-on overlapping of  $sp^3$  hybrid orbitals of carbon and 1s orbitals of hydrogen atoms.

#### 9.2.1 Nomenclature and Isomerism

You have already read about nomenclature of different classes of organic compounds in Unit 8. Nomenclature and isomerism in alkanes can further be understood with the help of a few more examples. Common names are given in parenthesis. First three alkanes – methane, ethane and propane have only one structure but higher alkanes can have more than one structure. Let us write structures for  $C_4H_{10}$ . Four carbon atoms of  $C_4H_{10}$  can be joined either in a continuous chain or with a branched chain in the following two ways :

Butane (n- butane), (b.p. 273 K)

2-Methylpropane (isobutane) (b.p.261 K)

In how many ways, you can join five carbon atoms and twelve hydrogen atoms of  $C_5H_{12}$ ? They can be arranged in three ways as shown in structures III–V

III H H H H H H H 
$$^{1}$$
  $^{1}$   $^{2}$   $^{1}$   $^{3}$   $^{4}$   $^{5}$   $^{1}$   $^{5}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$   $^{1}$  H H H H H

Pentane (*n*-pentane) (b.p. 309 K)

2-Methylbutane (isopentane) (b.p. 301 K)

V H

$$H \leftarrow C - H \rightarrow H$$
 $H \leftarrow C - C \rightarrow C - C - H$ 
 $H \leftarrow C - C \rightarrow C - H$ 
 $H \leftarrow C - H \rightarrow H$ 

2,2-Dimethylpropane (neopentane) (b.p. 282.5 K)

Structures I and II possess same molecular formula but differ in their boiling points and other properties. Similarly structures III, IV and V possess the same molecular formula but have different properties. Structures I and II are isomers of butane, whereas structures III, IV and V are isomers of pentane. Since difference in properties is due to difference in their structures, they are known as **structural** 

**isomers**. It is also clear that structures I and III have continuous chain of carbon atoms but structures II, IV and V have a branched chain. Such structural isomers which differ in chain of carbon atoms are known as **chain isomers**. Thus, you have seen that  $C_4H_{10}$  and  $C_5H_{12}$  have two and three chain isomers respectively.

### Problem 9.1

Write structures of different chain isomers of alkanes corresponding to the molecular formula  $\rm C_6H_{14}$ . Also write their IUPAC names.

#### Solution

(i) 
$$CH_3 - CH_2 - CH_2 - CH_2 - CH_2 - CH_3$$
*n*-Hexane

(ii) 
$$CH_3 - CH - CH_2 - CH_2 - CH_3$$
  
 $CH_3$ 

$$\begin{array}{c} \text{ 2-Methylpentane} \\ \text{(iii) } \text{CH}_3 - \text{CH}_2 - \text{CH} - \text{CH}_2 - \text{CH}_3 \\ \text{CH}_3 \\ \text{ 3-Methylpentane} \end{array}$$

3-Methylpentane (iv) 
$${\rm CH_3-CH-CH-CH_3}$$
  $\stackrel{|}{\rm CH_3}$   ${\rm CH_3}$ 

2,3-Dimethylbutane

$$\begin{array}{c} \operatorname{CH_3} \\ \text{(v)} \quad \operatorname{CH_3} - \operatorname{C} - \operatorname{CH_2} - \operatorname{CH_3} \\ \text{CH_3} \end{array}$$

2,2 - Dimethylbutane

Based upon the number of carbon atoms attached to a carbon atom, the carbon atom is termed as primary (1), secondary (2), tertiary (3) or quaternary (4). Carbon atom attached to no other carbon atom as in methane or to only one carbon atom as in ethane is called primary carbon atom. Terminal carbon atoms are always primary. Carbon atom attached to two carbon atoms is known as secondary. Tertiary carbon is attached to three carbon atoms and neo or quaternary carbon is attached to four carbon atoms. Can you identify 1, 2, 3 and 4 carbon atoms in

structures I to V? If you go on constructing structures for higher alkanes, you will be getting still larger number of isomers.  $C_6H_{14}$  has got five isomers and  $C_7H_{16}$  has nine. As many as 75 isomers are possible for  $C_{10}H_{22}$ .

In structures II, IV and V, you observed that  $-CH_3$  group is attached to carbon atom numbered as 2. You will come across groups like  $-CH_3$ ,  $-C_2H_5$ ,  $-C_3H_7$  etc. attached to carbon atoms in alkanes or other classes of

compounds. These groups or substituents are known as alkyl groups as they are derived from alkanes by removal of one hydrogen atom. General formula for alkyl groups is  $C_nH_{2n+1}$  (Unit 8).

Let us recall the general rules for nomenclature already discussed in Unit 8. Nomenclature of substituted alkanes can further be understood by considering the following problem:

#### Problem 9.2

Write structures of different isomeric alkyl groups corresponding to the molecular formula  $C_5H_{11}$ . Write IUPAC names of alcohols obtained by attachment of -OH groups at different carbons of the chain.

#### **Solution**

Structures of – $C_5H_{11}$ group	Corresponding alcohols	Name of alcohol
(i) CH <sub>3</sub> - CH <sub>2</sub> - CH <sub>2</sub> - CH <sub>2</sub> - CH <sub>2</sub> -	$\mathrm{CH_3} - \mathrm{CH_2} - \mathrm{CH_2} - \mathrm{CH_2} - \mathrm{CH_2} - \mathrm{OH}$	Pentan-1-ol
(ii) CH <sub>3</sub> - CH - CH <sub>2</sub> - CH <sub>2</sub> - CH <sub>3</sub>	$\begin{array}{c} \operatorname{CH}_3 - \operatorname{CH} - \operatorname{CH}_2 - \operatorname{CH}_2 - \operatorname{CH}_3 \\ \operatorname{OH} \end{array}$	Pentan-2-ol
(iii) $CH_3 - CH_2 - CH - CH_2 - CH_3$	$\begin{array}{c} \operatorname{CH}_3 - \operatorname{CH}_2 - \operatorname{CH} - \operatorname{CH}_2 - \operatorname{CH}_3 \\ \operatorname{OH} \end{array}$	Pentan-3-ol
$\begin{array}{c} {\rm CH_3} \\ {\rm I} \\ {\rm (iv)~CH_3-CH-CH_2-CH_2-} \end{array}$	$\begin{array}{c} \operatorname{CH_3} \\   \\ \operatorname{CH_3} - \operatorname{CH} - \operatorname{CH_2} - \operatorname{CH_2} - \operatorname{OH} \end{array}$	3-Methyl- butan-1-ol
$\begin{array}{c} \text{CH}_{3} \\ \text{(v) } \text{CH}_{3} - \text{CH}_{2} - \text{CH} - \text{CH}_{2} - \end{array}$	$\begin{array}{c} \operatorname{CH_3} \\ \operatorname{CH_3} - \operatorname{CH_2} - \operatorname{CH} - \operatorname{CH_2} - \operatorname{OH} \end{array}$	2-Methyl- butan-1-ol
$\begin{array}{c} \operatorname{CH_3} \\   \\ \text{(vi) } \operatorname{CH_3} - \underset{ }{\operatorname{C}} - \operatorname{CH_2} - \operatorname{CH_3} \end{array}$	$\begin{array}{c} \operatorname{CH_3} \\ \mid \\ \operatorname{CH_3} - \operatorname{C} - \operatorname{CH_2} - \operatorname{CH_3} \\ \mid \\ \operatorname{OH} \end{array}$	2-Methyl- butan-2-ol
$\begin{array}{c} \text{CH}_3 \\ \mid \\ \text{(vii) CH}_3 - \text{C} - \text{CH}_2 - \\ \mid \\ \text{CH}_3 \end{array}$	$\begin{array}{c} \operatorname{CH_3} \\ \operatorname{I} \\ \operatorname{CH_3} - \operatorname{C} - \operatorname{CH_2OH} \\ \operatorname{CH_3} \end{array}$	2,2- Dimethyl- propan-1-ol
$\begin{array}{ccc} \mathrm{CH_3} & & \\ & \mathrm{I} & & \\ \mathrm{(viii)} \ \mathrm{CH_3} - \mathrm{CH} - \mathrm{CH} - \mathrm{CH_3} \end{array}$	$\begin{array}{ccc} \mathrm{CH_3} & \mathrm{OH} \\ & \mathrm{I} & \mathrm{I} \\ \mathrm{CH_3} - \mathrm{CH} - \mathrm{CH} - \mathrm{CH_3} \end{array}$	3-Methyl- b u t a n - 2 - o l

Table 9.1 Nomenclature of a Few Organic Compounds

Str	ucture and IUPAC Name	Remarks
(a)	$CH_3$ $CH_2$ $-CH_3$ $CH_3$ $CH_3$ $CH_3$ $CH_4$ $CH_5$ $CH_5$ $CH_5$ $CH_5$ $CH_6$ $CH_7$ $CH_8$	Lowest sum and alphabetical arrangement
(b)	$ \begin{array}{c} {\rm CH_2-CH_3} \\ {\rm ^8CH_3-^7CH_2-^6CH_2-^5CH-^4CH-^3C} \\ {\rm ^{-3}CH_2-^1CH_3} \\ {\rm ^{-1}CH_3 CH_3 CH_2-CH_3} \\ \\ {\rm ^{-1}CH_3 CH_3} \end{array} $	Lowest sum and alphabetical arrangement
	(3,3-Diethyl-5-isopropyl-4-methyloctane)	
(c)	$^{1}\text{CH}_{3}$ $^{2}\text{CH}_{2}$ $^{3}\text{CH}_{2}$ $^{4}\text{CH}_{2}$ $^{5}\text{CH}_{2}$ $^{6}\text{CH}_{2}$ $^{7}\text{CH}_{2}$ $^{8}\text{CH}_{2}$ $^{9}\text{CH}_{2}$ $^{10}\text{CH}_{3}$ $^{1}\text{H}_{3}\text{C}$ $^{-}\text{CH}$ $^{-}\text{CH}_{2}$ $^{-}\text{CH}_{3}$ $^{5}$	sec is not considered while arranging alphabetically; isopropyl is taken
(d)	$^{1}\mathrm{CH_{3}}^{-2}\mathrm{CH_{2}}^{-3}\mathrm{CH_{2}}^{-4}\mathrm{CH_{2}}^{-5}\mathrm{CH}^{-6}\mathrm{CH_{2}}^{-7}\mathrm{CH_{2}}^{-8}\mathrm{CH_{2}}^{-9}\mathrm{CH_{3}}$ $^{ }_{\mathrm{CH_{2}}}$ $^{ }_{\mathrm{CH_{3}}^{-2}\mathrm{C-CH_{3}}}$	as one word  Further numbering to the substituents of the side chain
(e)	$^{3}\mathrm{CH}_{3}$ 5-(2,2– Dimethylpropyl)nonane $^{1}\mathrm{CH}_{3} - ^{2}\mathrm{CH}_{2} - ^{3}\mathrm{CH} - ^{4}\mathrm{CH}_{2} - ^{5}\mathrm{CH} - ^{6}\mathrm{CH}_{2} - ^{7}\mathrm{CH}_{3}$ $ \qquad \qquad \mathrm{CH}_{2} - \mathrm{CH}_{3}  \mathrm{CH}_{3}$ 3–Ethyl–5–methylheptane	Alphabetical priority order

#### Problem 9.3

Write IUPAC names of the following compounds:

- (i)  $(CH_3)_3 C CH_2C(CH_3)_3$
- (ii)  $(CH_3)_2 C(C_2H_5)_2$
- (iii) tetra tert-butylmethane

### Solution

- (i) 2, 2, 4, 4-Tetramethylpentane
- (ii) 3, 3-Dimethylpentane
- (iii) 3,3-Di-*tert*-butyl -2, 2, 4, 4 tetramethylpentane

If it is important to write the correct IUPAC name for a given structure, it is equally

important to write the correct structure from the given IUPAC name. To do this, first of all, the longest chain of carbon atoms corresponding to the parent alkane is written. Then after numbering it, the substituents are attached to the correct carbon atoms and finally valence of each carbon atom is satisfied by putting the correct number of hydrogen atoms. This can be clarified by writing the structure of 3-ethyl-2, 2-dimethylpentane in the following steps:

- i) Draw the chain of five carbon atoms: C C C C C
- ii) Give number to carbon atoms:  $C^1 C^2 C^3 C^4 C^5$

iii) Attach ethyl group at carbon 3 and two methyl groups at carbon 2

$$\begin{array}{c} CH_{3} \\ I \\ C^{1}-{}^{2}C-{}^{3}C-{}^{4}C-{}^{5}C \\ I \\ CH_{3} \\ C_{2}H_{5} \end{array}$$

iv) Satisfy the valence of each carbon atom by putting requisite number of hydrogen atoms:

$$\begin{array}{c} \text{CH}_{3} \\ \text{CH}_{3} - \text{C} - \text{CH} - \text{CH}_{2} - \text{CH}_{3} \\ \text{I} & \text{I} \\ \text{CH}_{3} & \text{C}_{2}\text{H}_{5} \end{array}$$

Thus we arrive at the correct structure. If you have understood writing of structure from the given name, attempt the following problems.

#### Problem 9.4

Write structural formulas of the following compounds:

- (i) 3, 4, 4, 5–Tetramethylheptane
- (ii) 2,5-Dimethyhexane

## Solution

$$\begin{array}{ccc} \mathrm{CH_3} & \mathrm{CH_3} \\ | & | \\ \mathrm{(ii)} \ \mathrm{CH_3} - \mathrm{CH} - \mathrm{CH_2} - \mathrm{CH_2} - \mathrm{CH} - \mathrm{CH_3} \end{array}$$

## Problem 9.5

Write structures for each of the following compounds. Why are the given names incorrect? Write correct IUPAC names.

- (i) 2-Ethylpentane
- (ii) 5-Ethyl 3-methylheptane

#### Solution

(i) 
$$CH_3 - CH - CH_2 - CH_2 - CH_3$$
  
 $C_2H_5$ 

Longest chain is of six carbon atoms and not that of five. Hence, correct name is 3-Methylhexane.

Numbering is to be started from the end which gives lower number to ethyl group. Hence, correct name is 3-ethyl-5-methylheptane.

## 9.2.2 Preparation

Petroleum and natural gas are the main sources of alkanes. However, alkanes can be prepared by following methods:

## 1. From unsaturated hydrocarbons

Dihydrogen gas adds to alkenes and alkynes in the presence of finely divided catalysts like platinum, palladium or nickel to form alkanes. This process is called **hydrogenation**. These metals adsorb dihydrogen gas on their surfaces and activate the hydrogen – hydrogen bond. Platinum and palladium catalyse the reaction at room temperature but relatively higher temperature and pressure are required with nickel catalysts.

$$CH_2 = CH_2 + H_2 \xrightarrow{Pt/Pd/Ni} CH_3 - CH_3$$
  
Ethene Propane (9.1)

$$CH_3$$
– $C\equiv C$ – $H$  +  $2H$   $\xrightarrow{Pt/Pd/Ni}$   $CH_3$ – $CH_2CH_3$   
Propyne Propane (9.3)

## 2. From alkyl halides

 i) Alkyl halides (except fluorides) on reduction with zinc and dilute hydrochloric acid give alkanes.

$$\text{CH-C1+H}_{2} \xrightarrow{\quad \text{Zn,H}^{+}} \text{CH}_{4} \text{+HC1} \quad \textbf{(9.4)}$$

Chloromethane Methane

$$\begin{array}{ccc} C_2H_5-C1+H_2 & \xrightarrow{Zn,H^+} & C_2H_6+HC1 \\ Chloroethane & Ethane & (9.5) \\ CH_3CH_2CH_2C1+H_2 & \xrightarrow{Zn,H^+} & CH_3CH_2CH_3+CH1 \\ 1-Chloropropane & Propane \\ & & (9.6) \end{array}$$

ii) Alkyl halides on treatment with sodium metal in dry ethereal (free from moisture) solution give higher alkanes. This reaction is known as **Wurtz reaction** and is used for the preparation of higher alkanes containing even number of carbon atoms.

$$\begin{tabular}{ll} ${\rm CH_3Br+2Na+BrCH_3} & \xrightarrow{\rm dry\; ether} {\rm CH_3+2Na} \\ {\rm Bromomenthane} & {\rm Ethane} \\ \end{tabular}$$

$$C_2H_5br+2Na+BrC_2H_5 \xrightarrow{dry \text{ ether}} C_2H_5-C_2H$$

Bromoethane n-Butane

(9.8)

What will happen if two different alkyl halides are taken?

## 3. From carboxylic acids

i) Sodium salts of carboxylic acids on heating with soda lime (mixture of sodium hydroxide and calcium oxide) give alkanes containing one carbon atom less than the carboxylic acid. This process of elimination of carbon dioxide from a carboxylic acid is known as **decarboxylation**.

$$CH_3COO^-Na^+ + MaOH \xrightarrow{CaO} CH_4 + NaCO_3$$
  
Sodium ethanoate

#### Problem 9.6

Sodium salt of which acid will be needed for the preparation of propane? Write chemical equation for the reaction.

## Solution

Butanoic acid, 
$$\label{eq:ch3} {\rm CH_3CH_2CH_2COO^{^-}Na^{^+}} + {\rm NaOH} \xrightarrow{\rm CaO} \\ {\rm CH_3CH_2CH_3} + {\rm Na_2CO_3}$$

ii) **Kolbe's electrolytic method** An aqueous solution of sodium or potassium salt of a carboxylic acid on electrolysis gives

alkane containing even number of carbon atoms at the anode.

$$2CH_3COO^-Na^+ + 2H_2O$$

Sodium acetate

$$CH_3 - CH_3 + 2CO_2 + H_2 + 2NaOH$$
 (9.9)

The reaction is supposed to follow the following path:

i) 
$$2CH_3COO^-Na^+ \rightleftharpoons 2CH_3 - C - O^- + 2Na^+$$

## ii) At anode:

$$\begin{array}{ccc} O & O \\ 2CH_3-C-O^{-\underline{-02e^-}} 2CH_3-C-\overset{\cdot}{\Omega} : &\longrightarrow 2\dot{C}H_3+2CO_2^{\uparrow} \\ Acetate \ ion & Acetate & Methyl \ free \\ & free \ radical & radical \end{array}$$

iii) 
$$H_3C + CH_3 \longrightarrow H_3C - CH_3$$

iv) At cathode:

$$H2O+e$$
 →  $OH+\dot{H}$   
 $2\dot{H}$  →  $H_2$  ↑

Methane cannot be prepared by this method. Why?

## 9.2.3 Properties

## Physical properties

Alkanes are almost non-polar molecules because of the covalent nature of C-C and C-H bonds and due to very little difference of electronegativity between carbon and hydrogen atoms. They possess weak van der Waals forces. Due to the weak forces, the first four members,  $C_1$  to  $C_4$  are gases,  $C_5$  to  $C_{17}$ are liquids and those containing 18 carbon atoms or more are solids at 298 K. They are colourless and odourless. What do you think about solubility of alkanes in water based upon non-polar nature of alkanes? Petrol is a mixture of hydrocarbons and is used as a fuel for automobiles. Petrol and lower fractions of petroleum are also used for dry cleaning of clothes to remove grease stains. On the basis of this observation, what do you think about the nature of the greasy substance? You are correct if you say that grease (mixture of higher alkanes) is non-

polar and, hence, hydrophobic in nature. It is generally observed that in relation to solubility of substances in solvents, polar substances are soluble in polar solvents, whereas the non-polar ones in non-polar solvents *i.e.*, *like dissolves like*.

Boiling point (b.p.) of different alkanes are given in Table 9.2 from which it is clear that there is a steady increase in boiling point with increase in molecular mass. This is due to the fact that the intermolecular van der Waals forces increase with increase of the molecular size or the surface area of the molecule.

You can make an interesting observation by having a look on the boiling points of three isomeric pentanes viz., (pentane, 2-methylbutane and 2,2-dimethylpropane). It is observed (Table 9.2) that pentane having a continuous chain of five carbon atoms has the highest boiling point (309.1K) whereas 2,2 – dimethylpropane boils at 282.5K. With increase in number of branched chains, the molecule attains the shape of a sphere. This results in smaller area of contact and therefore weak intermolecular forces between spherical molecules, which are overcome at relatively lower temperatures.

## **Chemical properties**

As already mentioned, alkanes are generally inert towards acids, bases, oxidising and

reducing agents. However, they undergo the following reactions under certain conditions.

#### 1. Substitution reactions

One or more hydrogen atoms of alkanes can be replaced by halogens, nitro group and sulphonic acid group. **Halogenation** takes place either at higher temperature (573-773 K) or in the presence of diffused sunlight or ultraviolet light. Lower alkanes do not undergo nitration and sulphonation reactions. These reactions in which hydrogen atoms of alkanes are substituted are known as *substitution reactions*. As an example, chlorination of methane is given below:

## Halogenation

Table 9.2 Variation of Melting Point and Boiling Point in Alkanes

Molecular formula	Name	Molecular mass/u	b.p./(K)	m.p./(K)
CH <sub>4</sub>	Methane	16	111.0	90.5
$C_2H_6$	Ethane	30	184.4	101.0
$C_3H_8$	Propane	44	230.9	85.3
$C_4H_{10}$	Butane	58	272.4	134.6
$C_4H_{10}$	2-Methylpropane	58	261.0	114.7
$C_5H_{12}$	Pentane	72	309.1	143.3
$C_5H_{12}$	2-Methylbutane	72	300.9	113.1
$C_5H_{12}$	2,2-Dimethylpropane	72	282.5	256.4
$C_{6}H_{14}$	Hexane	86	341.9	178.5
$C_7H_{16}$	Heptane	100	371.4	182.4
$C_8H_{18}$	Octane	114	398.7	216.2
$C_9H_{20}$	Nonane	128	423.8	222.0
$C_{10}^{-1}H_{22}^{-1}$	Decane	142	447.1	243.3
$C_{20}H_{42}$	Eicosane	282	615.0	236.2

$$CH_3$$
- $CH_3$  +  $C1_2$   $\xrightarrow{hv}$   $CH_3$ - $CH_2$ C1 + HC1  
Chloroethane (9.14)

It is found that the rate of reaction of alkanes with halogens is  $F_2 > Cl_2 > Br_2 > I_2$ . Rate of replacement of hydrogens of alkanes is: 3 > 2 > 1. Fluorination is too violent to be controlled. Iodination is very slow and a reversible reaction. It can be carried out in the presence of oxidizing agents like HIO<sub>3</sub> or HNO<sub>3</sub>.

$$CH_4 + I_2 \rightleftharpoons CH_3 I + HI$$
 (9.15)

$$HIO_3 + 5HI \rightarrow 31_2 + 3H_2O$$
 (9.16)

Halogenation is supposed to proceed via free radical chain mechanism involving three steps namely initiation, propagation and termination as given below:

#### Mechanism

(i) *Initiation*: The reaction is initiated by homolysis of chlorine molecule in the presence of light or heat. The Cl–Cl bond is weaker than the C–C and C–H bond and hence, is easiest to break.

C1–C1 
$$\xrightarrow{hv}$$
  $\dot{C}H_3$  + C1  
Chlorine free radicals

**(ii) Propagation:** Chlorine free radical attacks the methane molecule and takes the reaction in the forward direction by breaking the C-H bond to generate methyl free radical with the formation of H-Cl.

(a) 
$$CH_4 + \overset{\dagger}{C}1 \xrightarrow{hv} \overset{\dagger}{C}H_3 + H-C1$$

The methyl radical thus obtained attacks the second molecule of chlorine to form  $\mathrm{CH_3}$ –  $\mathrm{Cl}$  with the liberation of another chlorine free radical by homolysis of chlorine molecule.

(b) 
$$CH_3 + C1-C1 \xrightarrow{hv} CH_3 - C1 + C1$$

The chlorine and methyl free radicals generated above repeat steps (a) and (b) respectively and thereby setup a chain of reactions. The propagation steps (a) and (b) are those which directly give principal products, but many other propagation

steps are possible and may occur. Two such steps given below explain how more highly haloginated products are formed.

$$CH_3C1 + \dot{C}1 \rightarrow \dot{C}H_2C1 + HC1$$
  
 $\dot{C}H_2C1 + C1 - C1 \rightarrow CH_2C1_2 + \dot{C}1$ 

**(iii)** *Termination:* The reaction stops after some time due to consumption of reactants and / or due to the following side reactions:

The possible chain terminating steps are:

(a) 
$$\dot{C}1 + \dot{C}1 \rightarrow C1 - C1$$

(b) 
$$H_3 \dot{C} + \dot{C}H_3 \rightarrow H_3C - CH_3$$

(c) 
$$H_3 \dot{C}1 + \dot{C}1 \rightarrow H_3C-C1$$

Though in (c), CH<sub>3</sub> – Cl, the one of the products is formed but free radicals are consumed and the chain is terminated. The above mechanism helps us to understand the reason for the formation of ethane as a byproduct during chlorination of methane.

## 2. Combustion

Alkanes on heating in the presence of air or dioxygen are completely oxidized to carbon dioxide and water with the evolution of large amount of heat.

$$CH_{4}(g) + 2O_{2}(g) \rightarrow CO_{2}(g) + 2H_{2}O(1);$$

$$\ddot{A}_{c}H^{\dot{e}} - 890\text{kJ mol}^{-1}$$

$$(9.17)$$

$$C_{4}H_{10}(g) + 13/2O_{2}(g) \rightarrow 4CO_{2}(g) + 5H_{2}O(1)$$

$$\ddot{A}_{c}H^{\dot{e}} = -2875.84\text{kJ mol}^{-1}$$

$$(9.18)$$

The general combustion equation for any alkane is :

$$C_n H_{2n+2} + \left(\frac{3n+1}{2}\right) O_2 \to nCO_2 + (n+1)H_2O$$
(9.19)

Due to the evolution of large amount of heat during combustion, alkanes are used as fuels.

During incomplete combustion of alkanes with insufficient amount of air or dioxygen, carbon black is formed which is used in the manufacture of ink, printer ink, black pigments and as filters.

$$CH_4(g) + O_2(g) \xrightarrow{\text{incomplete}} C(s) + 2H_2 O(1)$$
(9.20)

## 3. Controlled oxidation

Alkanes on heating with a regulated supply of dioxygen or air at high pressure and in the presence of suitable catalysts give a variety of oxidation products.

(i) 
$$2CH_4 + O_2 \xrightarrow{-Cu/523K/100atm} 2CH^3 OH$$
  
Methanol

(ii) 
$$CH_4 + O_2 \xrightarrow{MO_2O_3} HCHO + H_2O$$
 Methanal

(9.22)

(iii) 
$$2\text{CH}_3\text{CH}_3 + 3\text{O}_2 \xrightarrow{(\text{CH}_3\text{COO})\text{Mn}} 2\text{CH}_3\text{COOH}$$
 Ethanoic acid 
$$+ 2\text{H}_2\text{O}$$
 (9.23)

(iv) Ordinarily alkanes resist oxidation but alkanes having tertiary H atom can be oxidized to corresponding alcohols by potassium permanganate.

$$(iCH_3)_3 CH \xrightarrow{KMnO_4} (CH_3)_3 COH$$

2-Methylpropane 2-Methylpropane-2-01

(9.24)

### 4. Isomerisation

*n*-Alkanes on heating in the presence of anhydrous aluminium chloride and hydrogen chloride gas isomerise to branched chain alkanes. Major products are given below. Some minor products are also possible which you can think over. Minor products are generally not reported in organic reactions.

$$\begin{array}{c} \text{CH3(CH)2)4CH3} \xrightarrow{\text{Anhy, AICI}_3/\text{ HCI}} \\ n\text{-Hexane} \\ \text{CH}_3\text{CH-(CH}_2)_2\text{-CH}_3\text{+CH}_3\text{CH}_2\text{-CH-CH}_2\text{-CH}_3 \\ \text{CH}_3 & \text{CH}_3 \\ \text{2-Methylpen tane} & 3\text{-Methylpenatone} \\ & (9.25) \end{array}$$

## 5. Aromatization

*n*-Alkanes having six or more carbon atoms on heating to 773K at 10-20 atmospheric

pressure in the presence of oxides of vanadium, molybdenum or chromium supported over alumina get **dehydrogenated** and cyclised to benzene and its homologues. This reaction is known as *aromatization* or *reforming*.

$$\begin{array}{c|c}
CH_3 & Cr_2O_3 \text{ or } V_2O_5 \\
CH_2 & CH_3 & \text{or } Mo_2O_3 \\
CH_2 & CH_2 & 773K, \\
CH_2 & 10-20 \text{ atm}
\end{array}$$

$$\begin{array}{c}
CH_3 & Cr_2O_3 \text{ or } V_2O_5 \\
OF Mo_2O_3 & OF Mo_2O_3 \\
OF Mo_2O_3 & OF$$

Toluene (C<sub>7</sub>H<sub>8</sub>) is methyl derivative of benzene. Which alkane do you suggest for preparation of toluene?

## 6. Reaction with steam

Methane reacts with steam at 1273 K in the presence of nickel catalyst to form carbon monoxide and dihydrogen. This method is used for industrial preparation of dihydrogen gas

$$CH_4 + H_{2IO} \xrightarrow{Ni} CO + 3H_2$$
 (9.27)

### 7. Pyrolysis

Higher alkanes on heating to higher temperature decompose into lower alkanes, alkenes etc. Such a decomposition reaction into smaller fragments by the application of heat is called *pyrolysis* or *cracking*.

(0.28)

Pyrolysis of alkanes is believed to be a free radical reaction. Preparation of oil gas or petrol gas from kerosene oil or petrol involves the principle of pyrolysis. For example, dodecane, a constituent of kerosene oil on heating to 973K in the presence of platinum, palladium or nickel gives a mixture of heptane and pentene.

$$\begin{array}{cccc} C_{12}H_{26} \xrightarrow{Pt/Pd/Ni} C7H_{16} & + & C5H_{10} & Other \\ Dodecane & Heptane & Pentene \\ & & & & (9.29) \end{array}$$