

Chapter 24 Hydrocarbon

Aliphatic Hydrocarbon

Organic compounds composed of only carbon and hydrogen are called hydrocarbons. Hydrocarbons are two types

- (1) Aliphatic Hydrocarbon (Alkanes, Alkenes and Alkynes).
- (2) Aromatic Hydrocarbon (Arenes)

(1) Sources of aliphatic hydrocarbon

Mineral oil or crude oil, petroleum [Petra → rock; oleum → oil] is the dark colour oily liquid with offensive odour found at various depths in many regions below the surface of the earth. It is generally found under the rocks of earth's crust and often floats over salted water.

(2) Composition

(i) **Alkanes** : found 30 to 70% contain upto 40 carbon atom. Alkanes are mostly straight chain but some are branched chain isomers.

(ii) **Cycloalkanes** : Found 16 to 64% cycloalkanes present in petroleum are; cyclohexane, methyl cyclopentane etc. cycloalkanes rich oil is called asphaltic oil.

(iii) **Aromatic hydrocarbon** : found 8 to 15% compound present in petroleum are; Benzene, Toluene, Xylene, Naphthalene etc.

(iv) **Sulphur, nitrogen and oxygen compound** : Sulphur compound present to the extent of 6% include mercaptans [R-SH] and sulphides [R-S-R]. The unpleasant smell of petroleum is due to sulphur compounds. Nitrogenous compounds are pyridines, quinolines and pyrroles. Oxygen compounds present in petroleum are. Alcohols, Phenols and resins. Compounds like chlorophyll, haemin are also present in it.

(v) **Natural gas** : It is a mixture of Methane (80%), Ethane (13%), Propane (3%), Butane (1%), Vapours of low boiling pentanes and hexanes (0.5%) and Nitrogen (1.3%). L.P.G. Contain butanes and

pentanes and used as cooking gas. It is highly inflammable. This contain, methane, nitrogen and ethane.

(vi) **C.N.G.** : When natural gas compressed at very high pressure is called compressed natural gas (CNG). Natural gas has octane rating of 130 it consists, mainly of methane and may contain, small amount of ethane and propane.

(3) **Theories of origin of petroleum** : Theories must explain the following characteristics associated with petroleum,

Its association with brine (sodium chloride solution). The presence of nitrogen and sulphur compounds in it. The presence of chlorophyll and haemin in it. Its optically active nature. Three important theories are as follows.

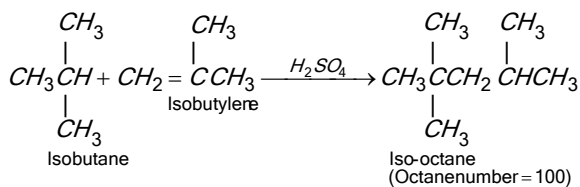
- (i) Mendeleeff's carbide theory or inorganic theory
- (ii) Engler's theory or organic theory
- (iii) Modern theory

(4) **Mining of petroleum** : Petroleum deposits occurs at varying depth at different places ranging from 500 to 15000 feet. This is brought to the surface by artificial drilling.

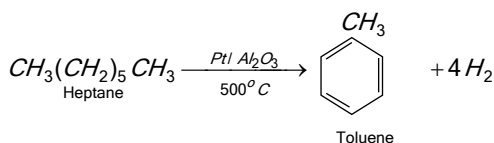
(5) **Petroleum refining** : Separation of useful fractions by fractional distillation is called petroleum refining.

Table : 24.1

Fraction	Boiling range (°C)	Approximate composition	Uses
Uncondensed gases	Upto room temperature	$C_1 - C_4$	Fuel gases; refrigerants; production of carbon black, hydrogen; synthesis of organic chemicals.
Crude naphtha on refractionation yields,	30 – 150°	$C_5 - C_{10}$	
(i) Petroleum ether	30 – 70°	$C_5 - C_6$	Solvent
(ii) Petrol or gasoline	70 – 120°	$C_6 - C_8$	Motor fuel; drycleaning; petrol gas.
(iii) Benzene derivatives	120 – 150°	$C_8 - C_{10}$	Solvent; drycleaning
Kerosene oil	150 – 250°	$C_{11} - C_{16}$	Fuel; illuminant; oil gas
Heavy oil	250 – 400°	$C_{15} - C_{18}$	As fuel for diesel engines; converted to gasoline by cracking.
Refractionation gives,			
(i) Gas oil, (ii) Fuel oil, (iii) Diesel oil			
Residual oil on fractionation by vacuum distillation gives,	Above 400°	$C_{17} - C_{40}$	
(i) Lubricating oil		$C_{17} - C_{20}$	Lubrication
(ii) Paraffin wax		$C_{20} - C_{30}$	Candles; boot polish; wax paper; etc
(iii) Vaseline		$C_{20} - C_{30}$	Toilets; ointments; lubrication.
(iv) Pitch		$C_{30} - C_{40}$	Paints, road surfacing
Petroleum coke (on redistilling tar)			As fuel.



(iii) **Aromatisation :**



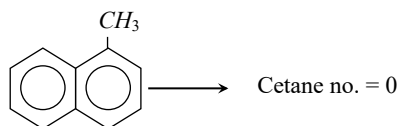
The octane no. of petrol can thus be improved.

- By increasing the proportion of branched chain or cyclic alkanes.
- By addition of aromatic hydrocarbons Benzene, Toluene and Xylene (BTX).

- By addition of methanol or ethanol.

- By addition of tetraethyl lead $(\text{C}_2\text{H}_5)_4\text{Pb}$

(5) **Cetane number :** It is used for grading the diesel oils.



The cetane number of a diesel oil is the percentage of cetane (hexadecane) by volume in a mixture of cetane and α -methyl naphthalene which has the same ignition property as the fuel oil under consideration.

(6) **Flash point :** The lowest temperature at which an oil gives sufficient vapours to form an explosive mixture with air is referred to as flash point of the oil.

The flash point in India is fixed at 44°C , in France it is fixed at 35°C , and in England at 22.8°C . The flash point of an oil is usually determined by means of "Abel's apparatus".

Chemists have prepared some hydrocarbons with octane number even less than zero (e.g., *n*-nonane has octane number - 45) as well as hydrocarbon with octane number greater than 100 (e.g., 2, 2, 3 trimethyl-butane. has octane number of 124).

(7) **Petrochemicals :** All such chemicals which are derived from petroleum or natural gas called petrochemicals. Some chemicals which are obtained from petroleum are summarised in table :

Table : 24.2

	nitroglycerine, dodecylbenzene, cumene, bakelite.
Hexane	Benzene, DDT, gammexane.
Heptane	Toluene
Cycloalkanes	Benzene, toluene, xylenes, adipic acid.
Benzene	Ethyl benzene, styrene, phenol, BHC (insecticide), adipic acid, nylon, cyclohexane, ABS detergents.
Toluene	Benzoic acid, TNT benzaldehyde, saccharin, chloramine-T, benzyl chloride, benzal chloride.

Alkanes [Paraffines]

"Alkanes are saturated hydrocarbon containing only carbon-carbon single bond in their molecules."

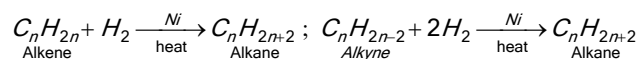
Alkanes are less reactive so called paraffins; because under normal conditions alkanes do not react with acids, bases, oxidising agents and reducing agent.

General formula : $\text{C}_n\text{H}_{2n+2}$

Examples are $\text{CH}_4, \text{C}_2\text{H}_6, \text{C}_3\text{H}_8$,

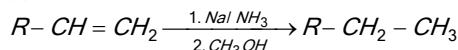
(1) **General Methods of preparation**

(i) **By catalytic hydrogenation of alkenes and alkynes** (Sabatier and Senderen's reaction)

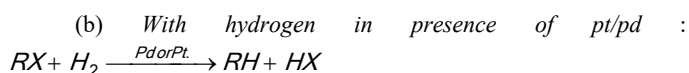
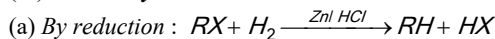


□ Methane is not prepared by this method

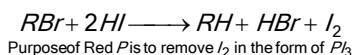
(ii) **Birch reduction :**



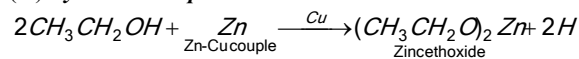
(iii) **From alkyl halide**



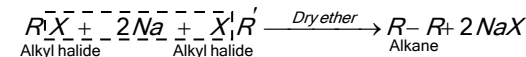
(c) *With HI in presence of Red phosphorus* :



(iv) **By Zn-Cu couple :**

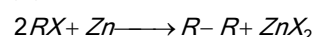


(v) **Wurtz reaction :**

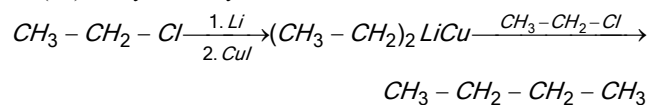


□ *R-Br* or *RI* preferred in this reaction. The net result in this reaction is the formation of even no. of carbon atoms in molecules.

(vi) **Frankland's reaction :**



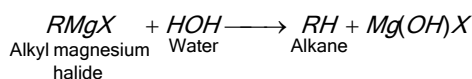
(vii) **Corey-house synthesis**



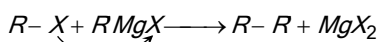
□ Reaction is suitable for odd number of Alkanes.

(viii) **From Grignard reagent**

(a) *By action of acidic 'H'* :

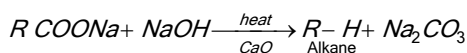


(b) By reaction with alkyl halide :



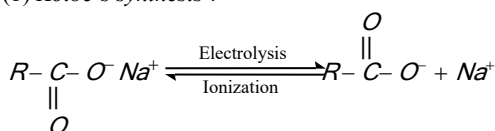
(ix) **From carboxylic acids**

(a) **Laboratory method** [Decarboxylation reaction or Duma reaction]

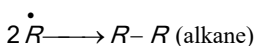
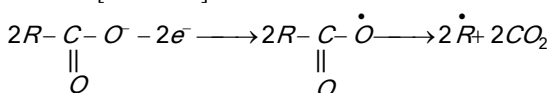


□ NaOH and CaO is in the ratio of 3 : 1. (Sodalime)

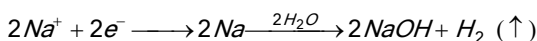
(b) **Kolbe's synthesis** :



At anode [Oxidation] :

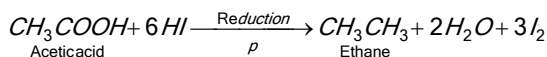


At cathode [Reduction] :

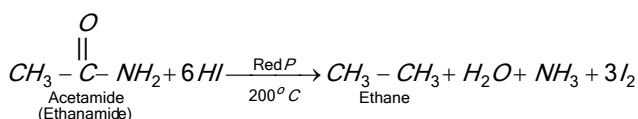
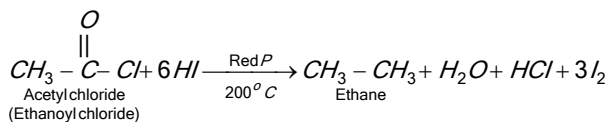
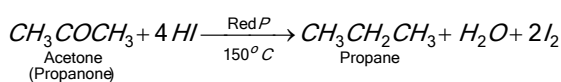
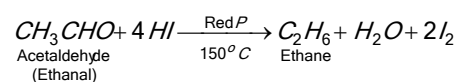
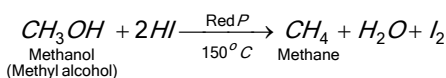


□ Both ionic and free radical mechanism are involved in this reaction.

(c) **Reduction of carboxylic acid** :

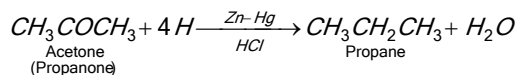
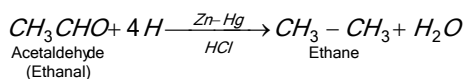


(x) **By reduction of alcohols, aldehyde, ketones or acid derivatives**



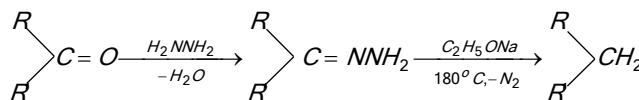
□ Aldehyde and ketones when reduced with amalgamated zinc and conc. HCl also yield alkanes.

Clemmenson reduction :



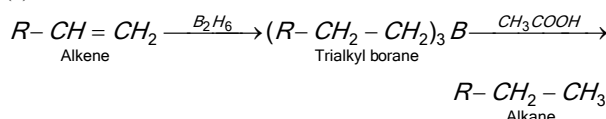
□ Aldehydes and ketones (>C=O) can be reduced to hydrocarbon in presence of excess of hydrazine and sodium alkoxide on heating.

Wolff-kishner reduction :

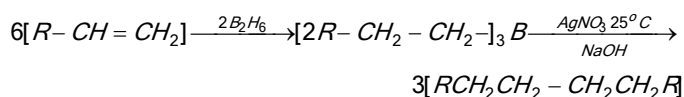


(xi) **Hydroboration of alkenes**

(a) **On treatment with acetic acid**



(b) **Coupling of alkyl boranes by means of silver nitrate**



(2) **Physical Properties**

(i) **Physical state** : Alkanes are colourless, odourless and tasteless.

Alkanes State

C₁ - C₄ Gaseous state

C₅ - C₁₇ Liquid state [Except neo pentane which is gas]

C₁₈ and above Solid like waxes

(ii) **Density** : Alkanes are lighter than water.

(iii) **Solubility** : Insoluble in water, soluble in organic solvents,

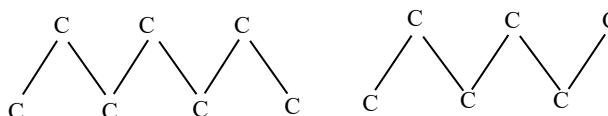
$$\text{solubility} \propto \frac{1}{\text{Molecular mass}}$$

(iv) **Boiling points and Melting points** : Melting points and

$$\text{boiling points} \propto \frac{1}{\text{No. of branches}}$$

Alkane :	C ₃ H ₈	C ₄ H ₁₀	C ₅ H ₁₂	C ₆ H ₁₄	C ₇ H ₁₆	C ₈ H ₁₈
M.P.(K)	85.9	138	143.3	179	182.5	216.2
:						

□ Melting points of even > Odd no. of carbon atoms, this is because, the alkanes with even number of carbon atoms have more symmetrical structure and result in closer packing in the crystal structure as compared to alkanes with odd number of carbon atoms.



(3) **Chemical properties**

(i) **Substitution reactions of Alkanes**

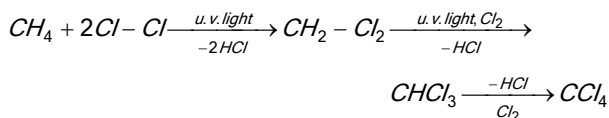
(a) **Halogenation** : R-H + X-X → R-X + HX

The reactivity of halogen is : F₂ > Cl₂ > Br₂ > I₂

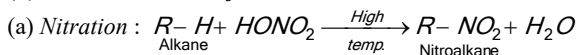
Fluorine can react in dark. Cl_2, Br_2 require light energy. I_2 does not show any reaction at room temperature, but on heating it shows iodination.

Iodination of methane is done in presence of oxidising agent such as HNO_3 / HIO_3 / HgO which neutralises HI .

Chlorination of methane :



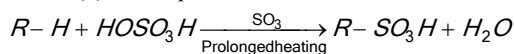
(ii) **Reaction based on free radical mechanism**



Nitrating mixture : (i) (Con. HNO_3 + Con. H_2SO_4) at $250^\circ C$

(ii) (HNO_3 vapour at $400^\circ - 500^\circ C$).

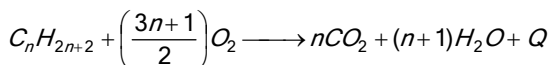
(b) **Sulphonation** : Free radical mechanism



Lower alkanes particularly methane, ethane, do not give this reaction.

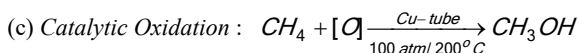
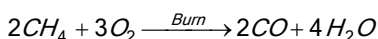
(iii) **Oxidation**

(a) **Complete Oxidation or combustion :**



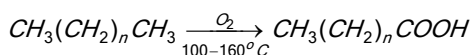
This is exothermic reaction.

(b) **Incomplete combustion or oxidation**

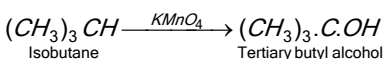


This is the industrial method for the manufacture of methyl alcohol.

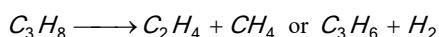
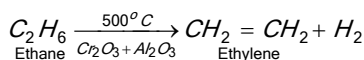
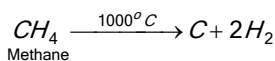
Higher alkanes are oxidised to fatty acids in presence of manganese stearate.



(d) **Chemical oxidation :**

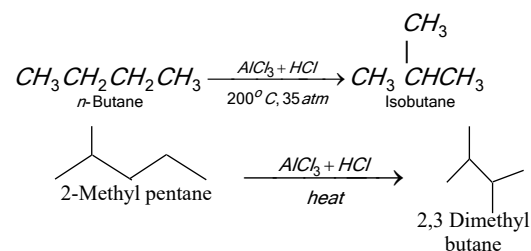


(iv) **Thermal decomposition or cracking or pyrolysis or fragmentation**

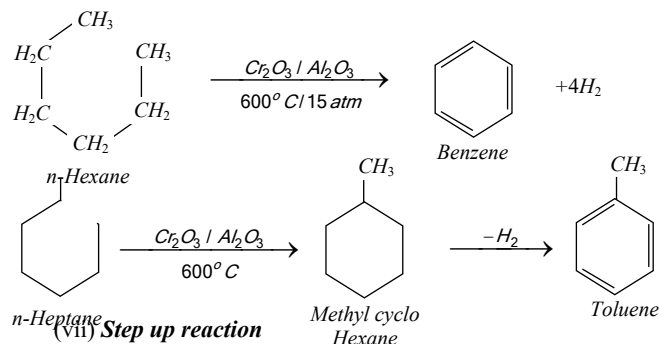


This reaction is of great importance to petroleum industry.

(v) **Isomerisation :**

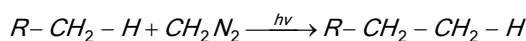


(vi) **Aromatisation :**

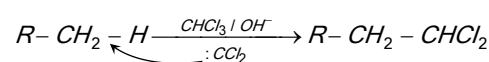


(vii) **Step up reaction**

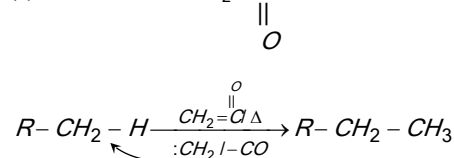
(a) **Reaction with CH_2N_2 (Diazo methane) :**



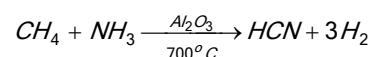
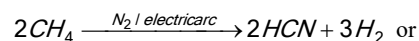
(b) **Reaction with $CHCl_3$ / $NaOH$:**



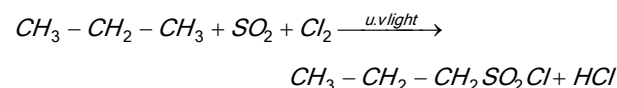
(c) **Reaction with $CH_2=C$:**



(viii) **HCN formation :**

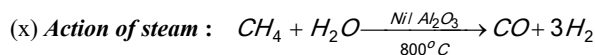


(ix) **Chloro sulphonation/Reaction with $SO_2 + Cl_2$**



This reaction is known as Reed's reaction.

This is used in the commercial formation of detergent.

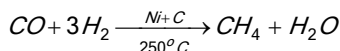


Individual members of alkanes

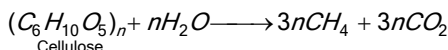
(1) **Methane** : Known as *marsh gas*.

(i) **Industrial method of preparation** : Methane gas is obtained on a large scale from natural gas by liquefaction. It can also be obtained by the application of following methods,

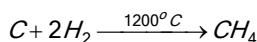
(a) *From carbon monoxide* : A mixture of carbon monoxide and hydrogen is passed over a catalyst containing nickel and carbon at 250°C when methane is formed.



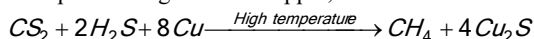
(b) *Bacterial decomposition of cellulose material present in sewage water* : This method is being used in England for production of methane.



(c) *Synthesis* : \square By striking an electric arc between carbon electrodes in an atmosphere of hydrogen at 1200°C , methane is formed.



By passing a mixture of hydrogen sulphide and carbon disulphide vapour through red hot copper, methane is formed.



(ii) Physical properties

(a) It is a colourless, odourless, tasteless and non-poisonous gas.

(b) It is lighter than air. Its density at NTP is 0.71 g/L.

(c) It is slightly soluble in water but is fairly soluble in ether, alcohol and acetone.

(d) Its melting point is -182.5°C and boiling point is -161.5°C .

(iii) Uses

(a) In the manufacture of compounds like methyl alcohol, formaldehyde, methyl chloride, chloroform, carbon tetrachloride, etc.

(b) In the manufacture of hydrogen, used for making ammonia.

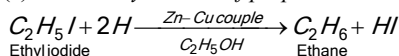
(c) In the preparation of carbon black which is used for making printing ink, black paints and as a filler in rubber vulcanisation.

(d) As a fuel and illuminant.

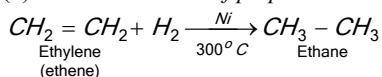
(2) Ethane

(i) Methods of preparation

(a) *Laboratory method of preparation* :



(b) *Industrial method of preparation* :



(ii) Physical properties

(a) It is a colourless, odourless, tasteless and non-poisonous gas.

(b) It is very slightly soluble in water but fairly soluble in alcohol, acetone, ether, etc.

(c) Its density at NTP is 1.34 g/L

(d) It boils at -89°C . Its melting point is -172°C .

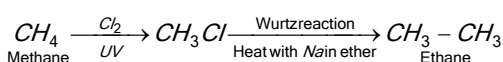
(iii) Uses

(a) As a fuel. (b) For making hexachloroethane which is an artificial camphor.

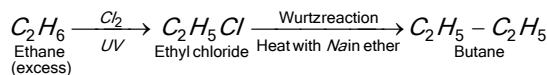
(3) Interconversion of Alkanes

Ascent of alkane series,

(i) Methane to ethane :

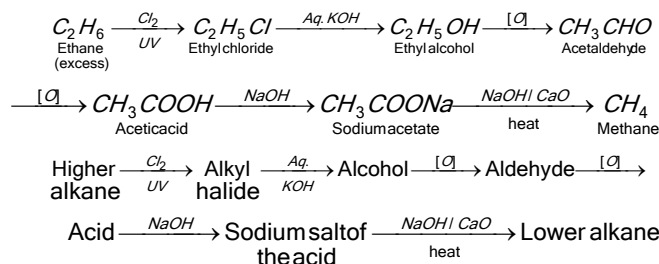


(ii) Butane from ethane :



Descent of alkane series : Use of decarboxylation reaction is made. It is a multistep conversion.

Ethane to methane

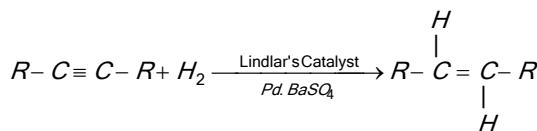


Alkenes

These are the acyclic hydrocarbon in which carbon-carbon contain double bond. These are also known as olefins, because lower alkene react with halogens to form oily substances. General formula is C_nH_{2n} . Examples, C_2H_4 , C_3H_6 , C_4H_8 .

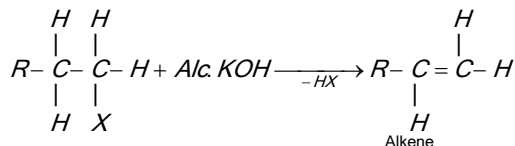
(1) Preparation methods

(i) From Alkynes :



\square Poison's catalyst such as BaSO_4 , CaCO_3 are used to stop the reaction after the formation of alkene.

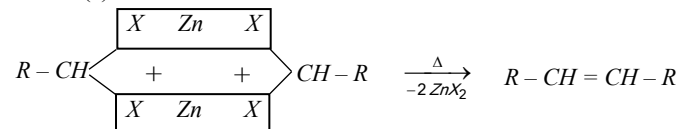
(ii) From mono halides :



\square If we use alc. NaOH in place of KOH then trans product is formed in majority because of its stability. According to saytzeff rule.

(iii) From dihalides

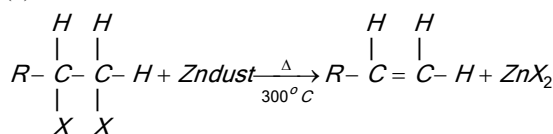
(a) From Gem dihalides



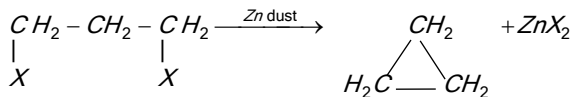
\square If we take two different types of gemdihalides then we get three different types of alkenes.

\square Above reaction is used in the formation of symmetrical alkenes only.

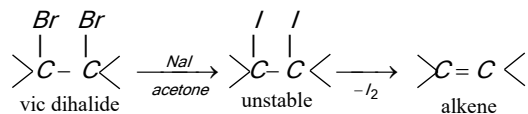
(b) From vicinal dihalides :



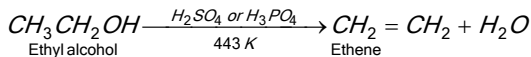
\square Alkene is not formed from 1, 3 dihalides. Cycloalkanes are formed by dehalogenation of it.



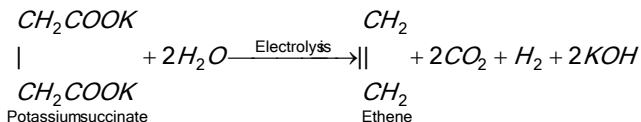
(iv) **By action of NaI on vicinal dihalide :**



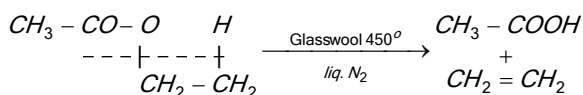
(v) **From alcohols** [Laboratory method] :



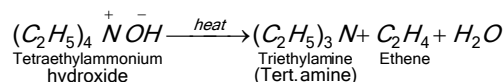
(vi) **Kolbe's reaction :**



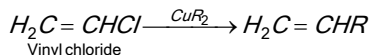
(vii) **From esters** [Pyrolysis of ester] :



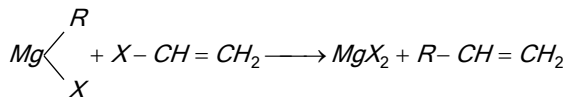
(viii) **Pyrolysis of quaternary ammonium compounds :**



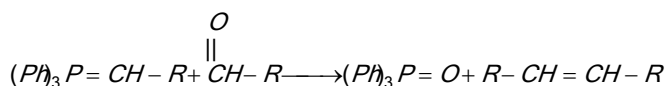
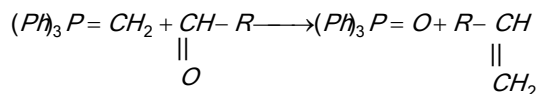
(ix) **Action of copper alkyl on vinyl chloride :**



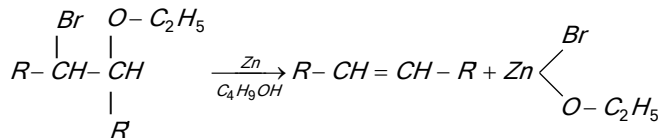
(x) **By Grignard reagents :**



(xi) **The Wittig reaction :**



(xii) **From β bromo ether** [Boord synthesis]

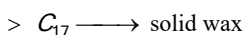
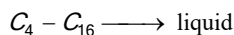
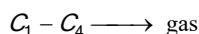


(2) **Physical Properties**

(i) Alkenes are colourless and odourless.

(ii) These are insoluble in water and soluble in organic solvents.

(iii) Physical state



(iv) **B.P.** and **M.P.** decreases with increasing branches in alkene.

(v) The melting points of cis isomers are lower than trans isomers because cis isomer is less symmetrical than trans. Thus trans packs more tightly in the crystal lattice and hence has a higher melting point.

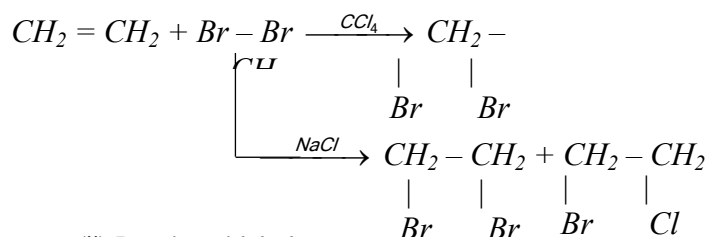
(vi) The boiling points of cis isomers are higher than trans isomers because cis-alkenes has greater polarity (Dipole moment) than trans one.

(vii) These are lighter than water.

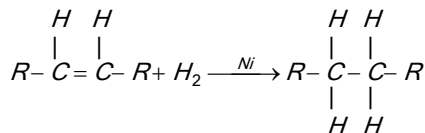
(viii) **Dipole moment** : Alkenes are weakly polar. The, π-electron's of the double bond. Can be easily polarized. Therefore, their dipole moments are higher than those of alkanes.

(3) **Chemical properties**

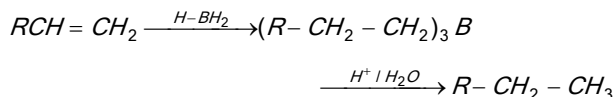
(i) **Francis experiment** : According to Francis electrophile first attacks on olefinic bond.



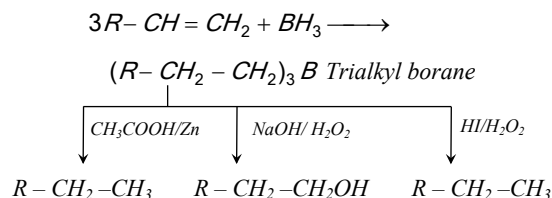
(ii) **Reaction with hydrogen :**



(iii) **Reduction of alkene via hydroboration** : Alkene can be converted into alkane by protolysis

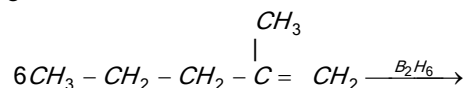


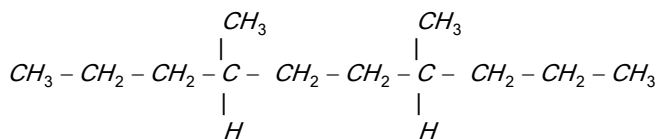
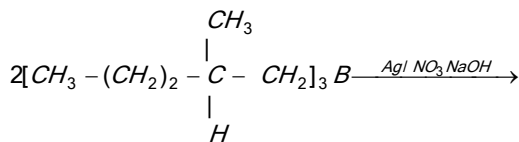
Hydroboration : Alkene give addition reaction with diborane which called hydroboration. In this reaction formed trialkylborane, Which is very important and used for synthesis of different organic compound



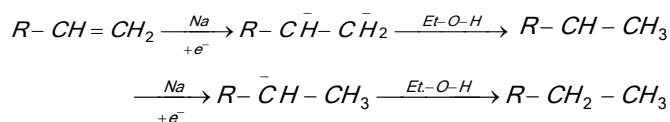
The overall result of the above reaction appears to be antimarkownikoff's addition of water to a double bond.

(iv) **By treatment with AgNO₃ + NaOH** : This reaction gives coupling

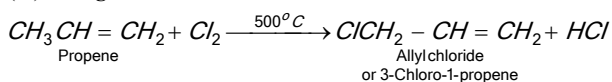




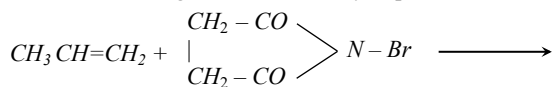
(v) **Birch reduction** : This reaction is believed to proceed via anionic free radical mechanism.



(vi) **Halogenation**

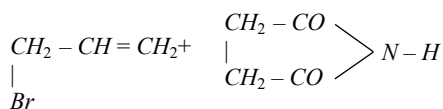


□ If NBS [N-bromo succinimide] is a reagent used for the specific purpose of brominating alkenes at the allylic position.



Propene

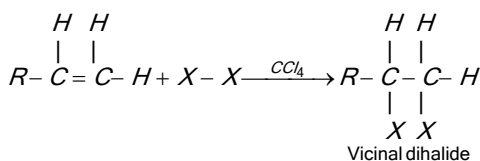
NBS



Allyl bromide

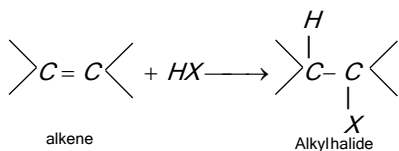
Succinimide

□ In presence of polar medium alkene form vicinal dihalide with halogen.

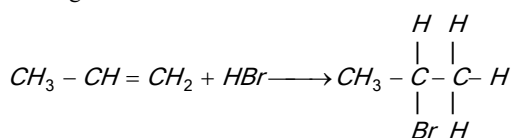


Reactivity of halogen is $\text{F}_2 > \text{Cl}_2 > \text{Br}_2 > \text{I}_2$

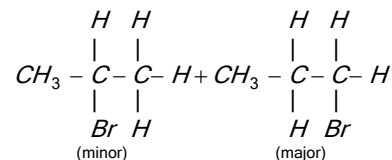
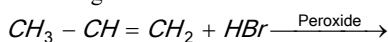
(vii) **Reaction with HX** [Hydrohalogenation]



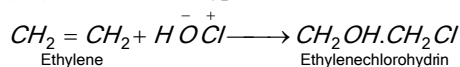
According to markownikoff's rule and kharasch effect.



According to Anti Markownikoff rule (Based on F.R.M.)

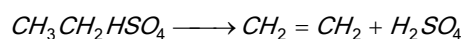
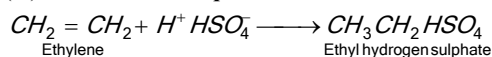


(viii) **Reaction with hypohalous acids** :



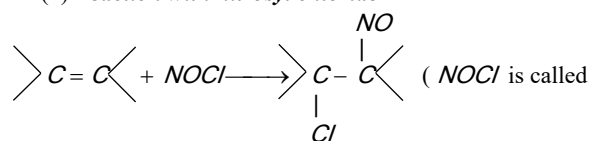
□ In case of unsymmetrical alkenes markownikoff rule is followed.

(ix) **Reaction with sulphuric acid** :



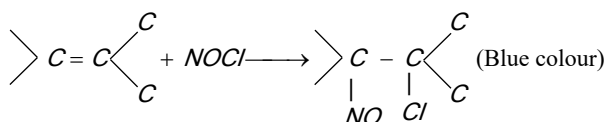
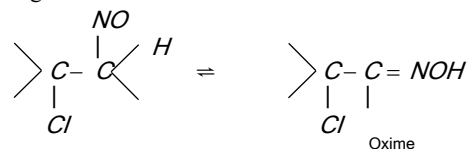
□ This reaction is used in the separation of alkene from a gaseous mixture of alkanes and alkenes.

(x) **Reaction with nitrosyl chloride**



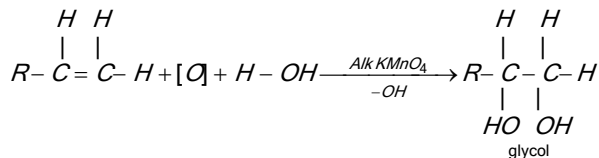
Tillden reagent)

□ If hydrogen is attached to the carbon atom of product, the product changes to more stable oxime.

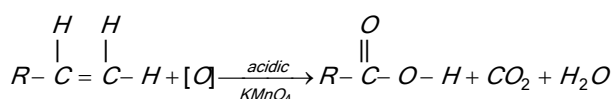


(xi) **Oxidation** : With alkaline KMnO_4 [Bayer's reagent] :

This reaction is used as a test of unsaturation.

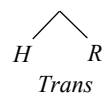
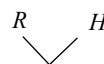
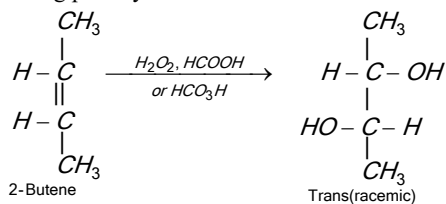


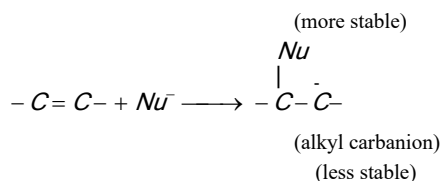
With acidic KMnO_4 :



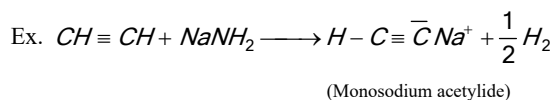
(xii) **Hydroxylation**

(a) Using per oxy acid :

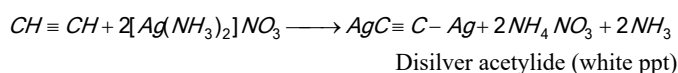
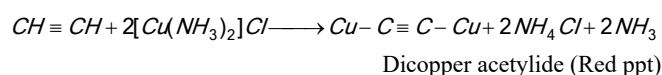




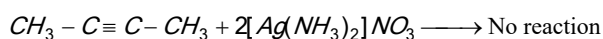
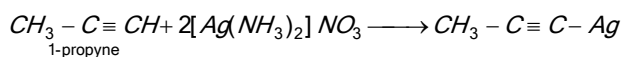
(i) **Acidity of alkynes** : Acetylene and other terminal alkynes (1-alkynes) are weakly acidic in character



The acetylenic hydrogen of alkynes can be replaced by copper (I) and silver (I) ions. They react with ammoniacal solutions of cuprous chloride and silver nitrate to form the corresponding copper and silver alkynides.



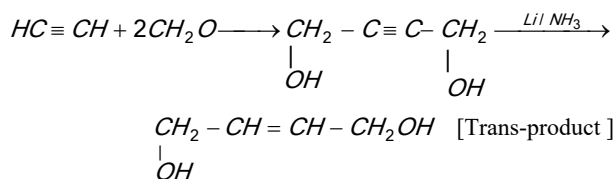
This reaction can be used to distinguish between 2-alkynes and 1-alkynes. 1-alkynes will give this test while 2-alkynes, will not give this test.



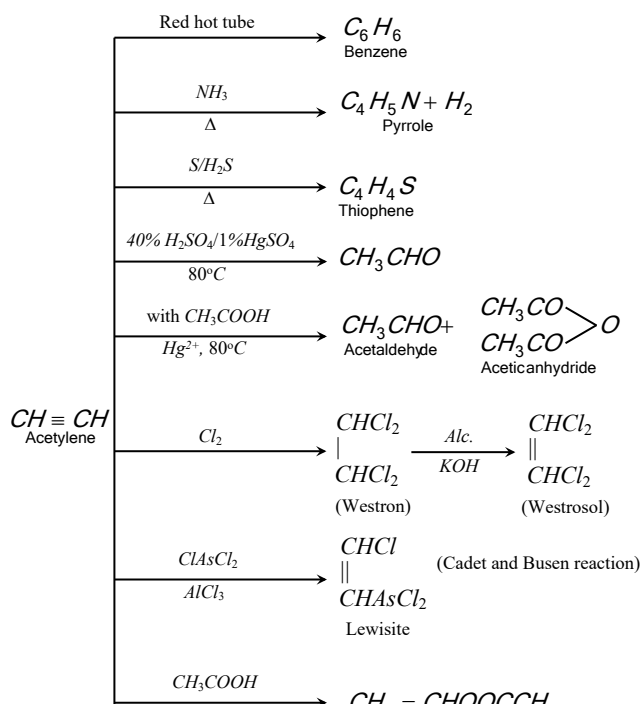
Explanation for the acidic character : It explained by *sp*

hybridisation. We know that an electron in *s*-orbital is more tightly held than in a *p*-orbital. In *sp* hybridisation *s*-character is more (50%) as compared to *sp²* (33%) or *sp³* (25%), due to large *s*-character the carbon atom is quite electronegative.

(ii) **Reaction with formaldehyde**



(4) Chemical properties of acetylene

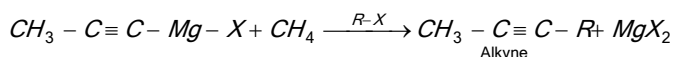
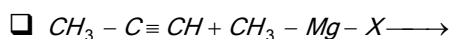
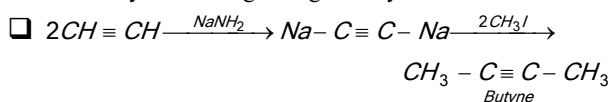


□ In reaction with gem dihalide, Alc. KOH is not used for elimination in 2nd step.

□ In reaction with vicinal dihalide, if the reactant is 2-butylene chloride then product is 2-butyne as major product.

Preparation of higher alkynes (by metal acetylide)

□ Acetylene gives salt with $NaNH_2$ or $AgNO_3$ (ammonical) which react with alkyl halide to give higher alkyne.



(2) Physical properties

(i) Acetylene is a colourless gas. It has a garlic odour. The odour is due to presence of impurities of phosphorous and hydrogen sulphide. However, pure acetylene has pleasant odour.

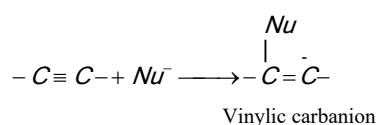
(ii) It is insoluble in water but highly soluble in acetone and alcohol. Acetylene is transported under high pressure in acetone soaked on porous material packed in steel cylinders.

(iii) Its boiling point is $-84^\circ C$.

(iv) It is lighter than air. It is somewhat poisonous in nature.

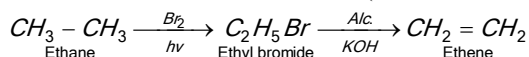
(v) It burns with luminous flame and forms explosive mixture with air.

(3) **Chemical reactivity of alkynes** : $C \equiv C$ is less reactive than the carbon-carbon double bond towards electrophilic addition reaction. This is because in alkyne carbon has more *s*-character so more strongly will be the attraction for π electrons. Alkyne also undergo nucleophilic addition with electron rich reagents. Ex. Addition of water, cyanide, carboxylic acid, alcohols. Nucleophilic addition can be explained on the basis that alkynes form vinylic carbanion which is more stable than alkyl carbanion formed by alkene

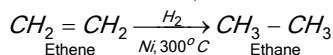


(6) Interconversion

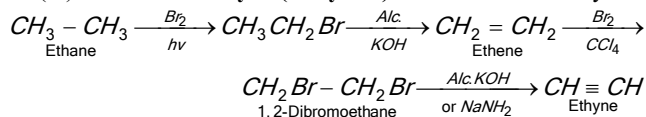
(i) Conversion of ethane into ethene : (Alkane into alkene)



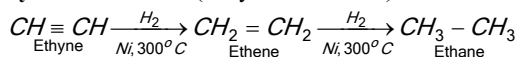
(ii) Ethene into ethane : (Alkene into alkane)



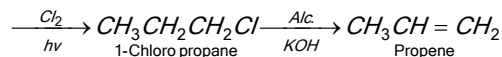
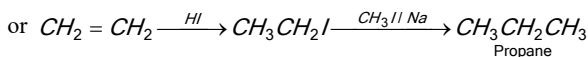
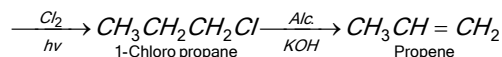
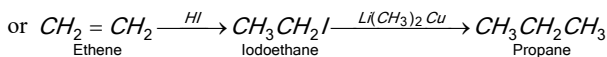
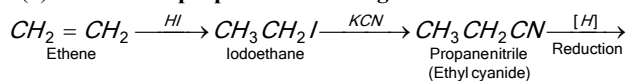
(iii) Ethane into ethyne (acetylene) : i.e., alkane into alkyne



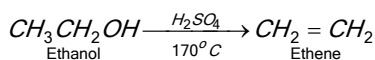
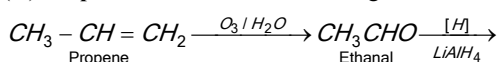
(iv) Ethyne into ethane : (Alkyne into alkane)



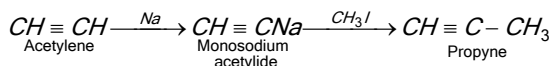
(v) Ethene into propene : Ascending in alkene series



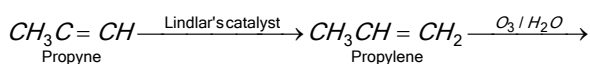
(vi) Propene into ethene : Descending an alkene series



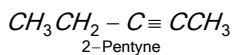
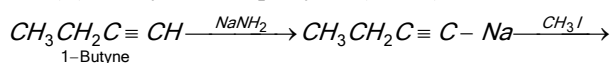
(vii) Acetylene into propyne (methyl acetylene) : (Ascent)



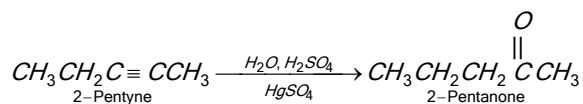
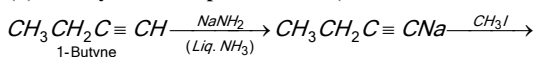
(viii) Propyne into acetylene : (Descent)



(ix) 1-Butyne into 2-pentyne : (Ascent)

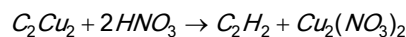
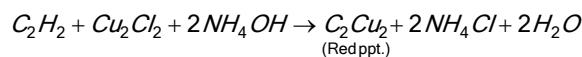


(x) 1-Butyne into 2-pentanone : (Not more than three steps)

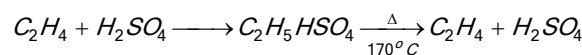


Separation of alkane, alkene and alkyne

The gaseous mixture is passed through ammonical cuprous chloride solution. The alkyne (acetylene) reacts with Cu_2Cl_2 and forms a red precipitate. It is filtered. The alkyne or acetylene is recovered by decomposition of the precipitate with an acid.



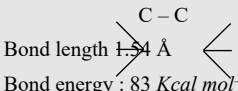
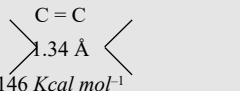
The remaining gaseous mixture is passed through concentrated H_2SO_4 . Alkene is absorbed. The Hydrogen sulphate derivatives is heated at 170°C to regenerate ethene.



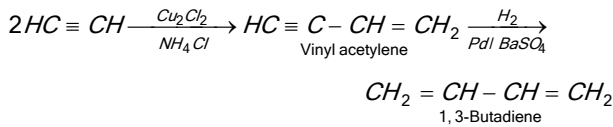
The methane or ethane is left behind unreacted.

Distinction between alkanes, Alkenes and Alkynes

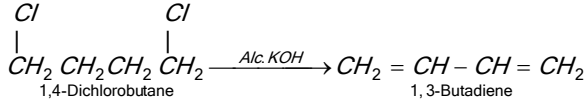
Table : 24.3

Property	Alkane (Ethane)	Alkene (Ethene)	Alkyne (Ethyne)
Molecular formula	$C_nH_{2n+2}(C_2H_6)$	$C_nH_{2n}(C_2H_4)$	$C_nH_{2n-2}(C_2H_2)$
Nature	Saturated	Unsaturated	Unsaturated
	Single bond between carbon atoms. Each carbon atom is sp^3 -hybridized Bond length 1.54 \AA Bond energy : 83 Kcal mol^{-1} 	Double bond between two carbon atoms. Both carbon atoms are sp^2 -hybridized Bond length 1.34 \AA $146 \text{ Kcal mol}^{-1}$ 	Triple bond between two carbon atoms both carbon atoms are sp -hybridized $-C \equiv C-$ 1.20 \AA $200 \text{ Kcal mol}^{-1}$
Burning	Burns with nonluminous flame $C_2H_6 + 7/2O_2 \rightarrow 2CO_2 + 3H_2O$	Burns with luminous flame $C_2H_4 + 3O_2 \rightarrow 2CO_2 + 2H_2O$	Burns with smoky flame $C_2H_2 + 5/2O_2 \rightarrow 2CO_2 + H_2O$
Reaction with H_2	–	Forms alkane $C_nH_{2n} + H_2 \xrightarrow[300^\circ C]{Ni} C_nH_{2n+2}$ Alkane $C_2H_4 + H_2 \rightarrow C_2H_6$	Forms alkene and alkane $C_nH_{2n} + H_2 \xrightarrow[300^\circ C]{Ni} C_nH_{2n+2}$ Alkane $C_nH_{2n-2} + H_2 \xrightarrow[300^\circ C]{Ni} C_nH_{2n}$ Alkene
Reaction with conc. H_2SO_4 and hydrolysis	–	Addition $C_2H_4 + H_2SO_4 \rightarrow C_2H_5HSO_4$ $\xrightarrow{H_2O} C_2H_5OH$ Alcohol	Addition $C_2H_2 \rightarrow CH_3CH(HSO_4)_2 \xrightarrow{H_2O} CH_3CHO$ Aldehyde
Br_2/CCl_4	–	Decolourises Dibromo derivative, $C_2H_4 + Br_2 \rightarrow C_2H_4Br_2$	Decolourises Tetrabromo derivative, $C_2H_2Br_4$
Baeyer's reagent (Alk. $KMnO_4$)	–	Decolourises Glycol is formed $CH_2 \quad CH_2OH$ $\parallel + H_2O + O \rightarrow $ $CH_2 \quad CH_2OH$	Decolourises Oxalic acid is formed $CH \quad COOH$ $\parallel + 4O \rightarrow $ $CH \quad COOH$
Ammonical Cu_2Cl_2	–	–	Red precipitate $CH \quad CCu$ $\parallel + Cu_2Cl_2 + 2NH_4OH \rightarrow \parallel$ $CH \quad CCu$ (Red) $+ 2NH_4Cl + 2H_2O$
Ammonical silver nitrate	–	–	White precipitate $CH \quad C-Ag$ $\parallel + 2AgNO_3 + 2NH_4OH \rightarrow \parallel$ $CH \quad C-Ag$ $+ 2NH_4Cl + 2H_2O$

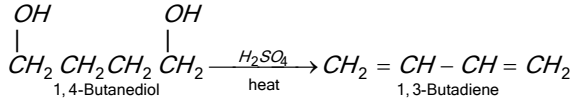
(i) **From acetylene :**



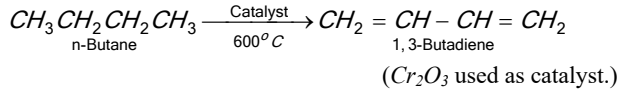
(ii) **From 1, 4-dichlorobutane :**



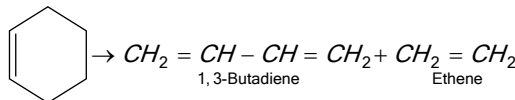
(iii) **From 1,4-butanediol :**



(iv) **From butane :**



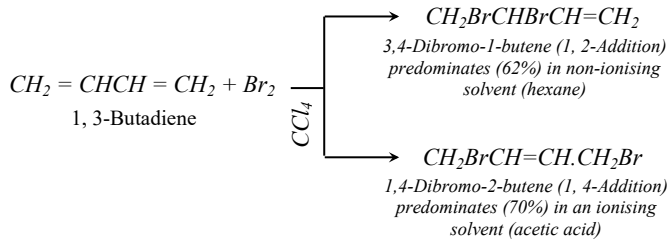
(v) **From cyclohexene :**



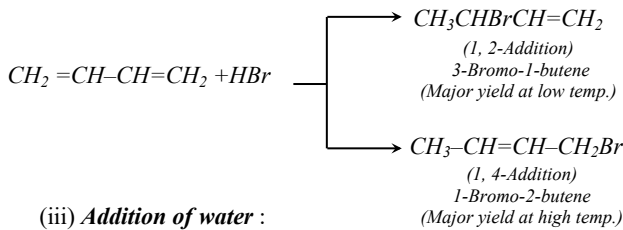
(2) **Physical property :** 1,3-butadiene is a gas.

(3) **Chemical properties**

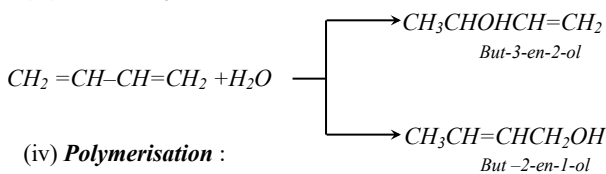
(i) **Addition of halogens :**



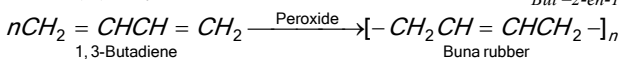
(ii) **Addition of halogen acids :**



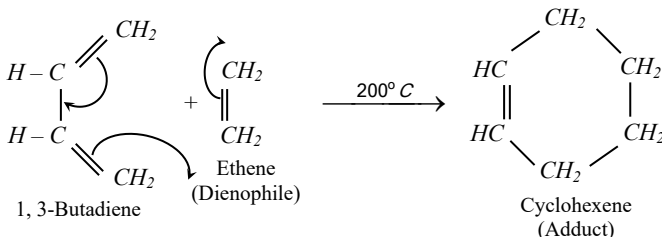
(iii) **Addition of water :**



(iv) **Polymerisation :**

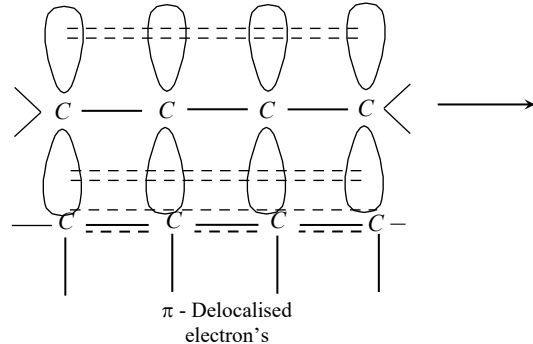


Diels-alder reaction :

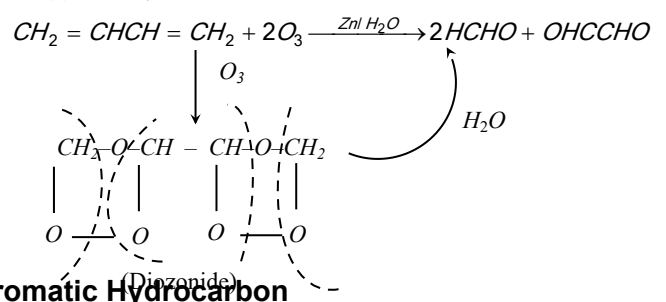


Stability of conjugated dienes : It is explained on the basis of delocalisation of electron cloud between carbon atoms.

The four π electrons of 1, 3-butadiene are delocalised over all the four atoms. This delocalisation of the π electrons makes the molecule more stable.



(v) **Ozonolysis :**



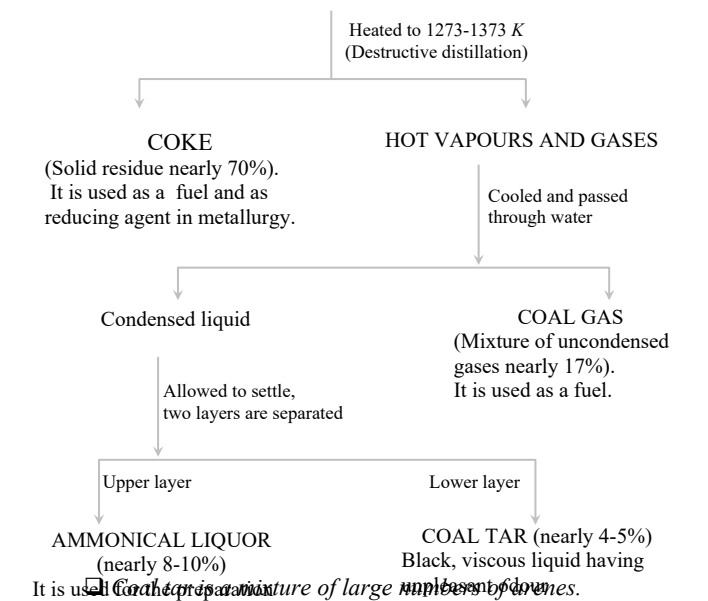
Aromatic Hydrocarbon

(1) **Source of Arenes**

Source of arenes is coal. It contains benzene, xylene, naphthalene etc. Arenes are obtained by destructive distillation of coal.

(2) **Distillation of coal**

COA



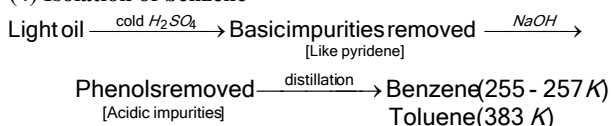
(3) **Distillation of coal tar :** Arenes are isolated by fractional distillation of coal tar,

Table : 24.4

Name of the fraction	Temperature range (K)	Main constituents
Light oil (or crude oil) fraction	Upto 443	Benzene, toluene, xylene
Middle oil fraction (Carbolic oil)	443-503	Phenol, naphthalene, pyridine
Heavy oil fraction (Creosote oil)	503-543	Naphthalene, naphthol and cresol
Green oil (Anthracene oil)	543-633	Anthracene, phenanthrene
Pitch (left as residue)	Non-volatile	Carbon

□ The residue left after fractional distillation of coal-tar is called pitch.

(4) Isolation of benzene



General characteristics of arenes

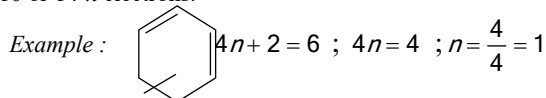
(1) All arenes have general formula $[C_nH_{2n-6}]_y$. Where y is number of benzene rings and n is not less than 6.

(2) Arenes are cyclic and planar. They undergo substitution rather than addition reactions.

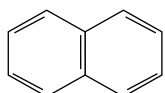
(3) **Aromaticity or aromatic character** : The characteristic behaviour of aromatic compounds is called aromaticity. Aromaticity is due to extensive delocalisation of π -electrons in planar ring system. Huckel (1931) explained aromaticity on the basis of following rule.

Huckel rule : For aromaticity the molecule must be planar, cyclic system having delocalised $(4n+2)\pi$ electrons where n is an integer equal to 0, 1, 2, 3,-----.

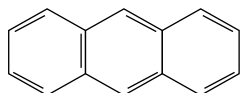
Thus, the aromatic compounds have delocalised electron cloud of 2,6,10 or 14 π electrons.



Benzene 6π electrons
 $n=1$

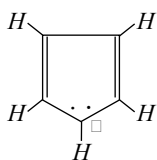


Naphthalene 10π electrons
 $n=2$

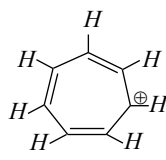


Anthracene 14π electrons
 $n=3$

Similarly cyclopentadienyl anion or tropylium ion are also aromatic because of containing 6π electrons ($n=1$).



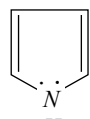
Cyclopentadienyl anion 6π electrons ($n=1$)



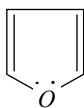
Tropylium ion 6π electrons ($n=1$)



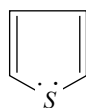
Cyclopropenyl anion 2π electrons ($n=0$)



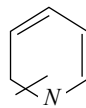
Pyrrole



Furan



Thiophene

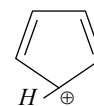


Pyridine

Molecules do not satisfy huckel rule are not aromatic.



Cyclopentadiene
 4π electrons



Cyclopentadienyl cation
 4π electrons



Cyclooctatetraene
 8π electrons



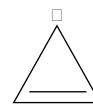
Cyclopropenyl anion
 4π electrons

(4) **Antiaromaticity** : Planar cyclic conjugated species, less stable than the corresponding acyclic unsaturated species are called antiaromatic. Molecular orbital calculations have shown that such compounds have $4n\pi$ electrons. In fact such cyclic compounds which have $4n\pi$ electrons are called antiaromatic compounds and this characteristic is called antiaromaticity.

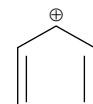
Example : 1,3-Cyclobutadiene, It is extremely unstable antiaromatic compound because it has $4n\pi$ electrons ($n=1$) and it is less stable than 1,3 butadiene by about 83.6 KJ mol^{-1} .



Thus, cyclobutadiene shows two equivalent contributing structures and it has $n=1$.



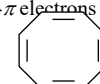
Cyclopropenyl anion
 4π electrons



Cyclopentadienyl
 4π electrons



Cycloheptatrienyl anion
 8π electrons



Cyclooctatetraene
 8π electrons

Benzene (C_6H_6)

Benzene is the first member of arenes. It was first discovered by Faraday (1825) from whale oil. Mitscherlich (1833) obtained it by distilling benzoic acid with lime. Hofmann (1845) obtained it from coal tar, which is still a commercial source of benzene.

(1) **Structure of benzene** : Benzene has a special structure, which is although unsaturated even then it generally behave as a saturated compound.

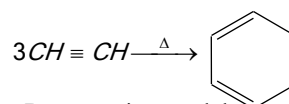
(i) **Kekule's structure** : According to Kekule, in benzene 6-carbon atoms placed at corner of hexagon and bonded with hydrogen and double bond present at alternate position.

(a) **Evidence in favour of Kekule's structure**

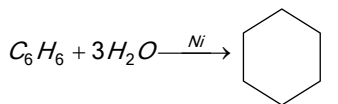
- Benzene combines with 3 molecules of hydrogen or three molecules of chlorine. It also combines with 3 molecules of ozone to form triozonide. These reactions confirm the presence of three double bonds.

- Studies on magnetic rotation and molecular refraction show the presence of three double bonds and a conjugated system.

- The synthesis of benzene from three molecule of acetylene also favour's Kekule's structure.



- Benzene gives cyclohexane by reduction with hydrogen.



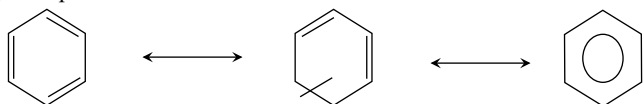
Cyclohexane

(b) *Objections against Kekule's formula*

- Unusual stability of benzene.
- According to Kekule, two ortho disubstituted products are possible. But in practice only one ortho disubstituted product is known.
- Heat of hydrogenation of benzene is 49.8 kcal/mole, whereas theoretical value of heat of hydrogenation of benzene is 85.8 kcal/mole. It means resonance energy is 36 kcal/mole.

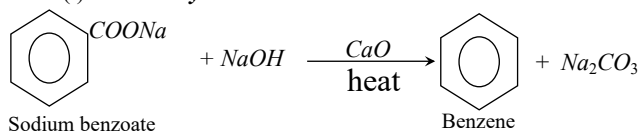
• C-C bond length in benzene are equal, (although it contains 3 double bonds and 3 single bonds) and are 1.39 Å.

Kekule explained this objection by proposing that double bonds in benzene ring were continuously oscillating between two adjacent positions.



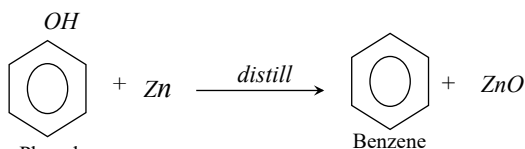
(2) **Methods of preparation of benzene**

(i) **Laboratory method :**

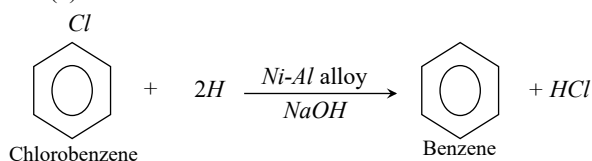


(ii) **From benzene derivatives**

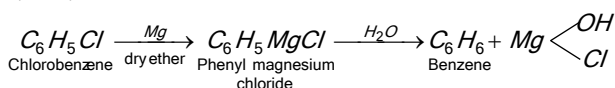
(a) **From phenol :**



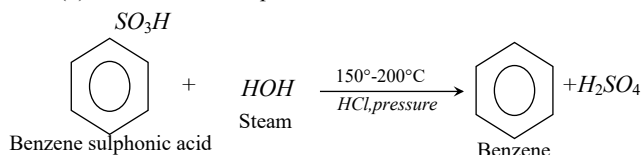
(b) **From chlorobenzene :**



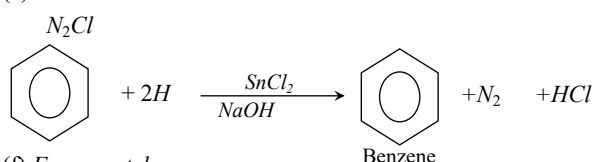
(c) **By first preparing grignard reagent of chlorobenzene and then hydrolysed**



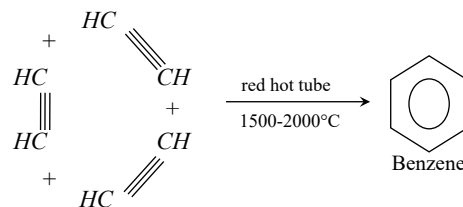
(d) **From benzene sulphonic acid :**



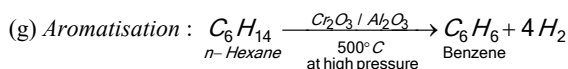
(e) **From benzene diazonium chloride :**



(f) **From acetylene :**



Three molecules of acetylene
□ Cyclic polymerisation takes place in this reaction.



(3) **Properties of benzene**

(i) **Physical properties**

(a) Benzene is a colourless, mobile and volatile liquid. It's boiling point is 80°C and freezing point is 5.5°C. It has characteristic odour.

(b) It is highly inflammable and burns with sooty flame.

(c) It is lighter than water. It's specific gravity at 20°C is 0.8788.

(d) It is immiscible with water but miscible with organic solvents such as alcohol and ether.

(e) Benzene itself is a good solvent. Fats, resins, rubber, etc. dissolve in it.

(f) It is a non-polar compound and its dipole moment is zero.

(g) It is an extremely poisonous substance. Inhalation of vapours or absorption through skin has a toxic effect.

(ii) **Chemical properties :** Due to the presence of π electron clouds above and below the plane benzene ring, the ring serves as a source of electrons and is easily attacked by electrophiles (Electron loving reagents). Hence electrophilic substitution reaction are the characteristic reactions of aromatic compounds.

Substitution reactions in benzene are preferred rather than addition are due to the fact that in the former reactions resonance stabilised benzene ring system is retained while the addition reactions lead to the destruction of benzene ring. Principal reactions of benzene can be studied under three heads,

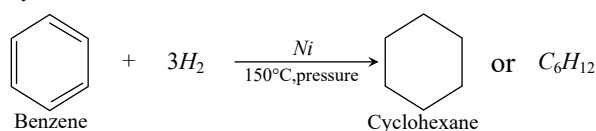
(a) Addition reactions

(b) Substitution reactions

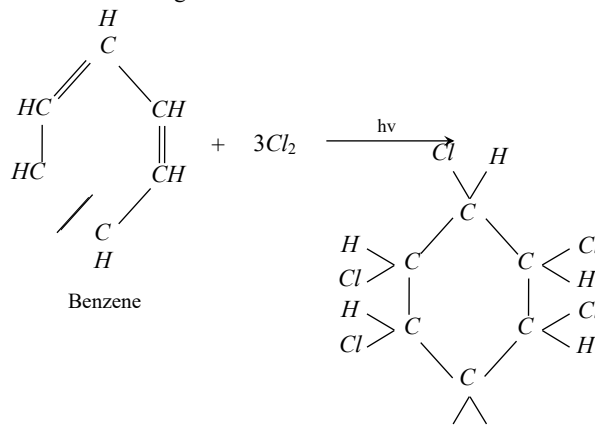
(c) Oxidation reactions

(a) **Addition reactions :** In which benzene behaves like unsaturated hydrocarbon.

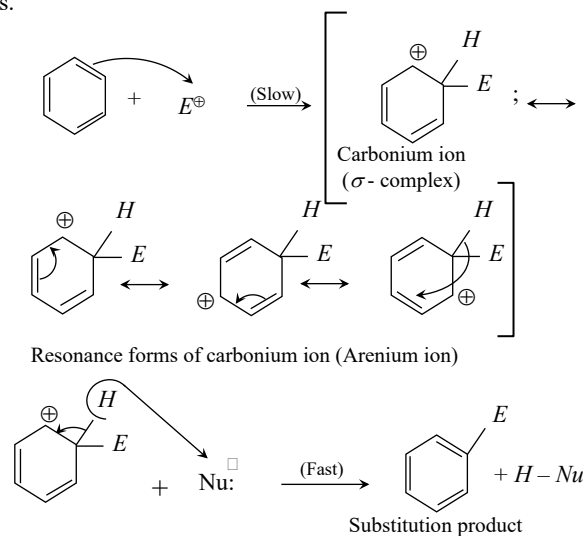
Addition of hydrogen : Benzene reacts with hydrogen in the presence of nickel (or platinum) as catalyst at 150°C under pressure to form cyclohexane.



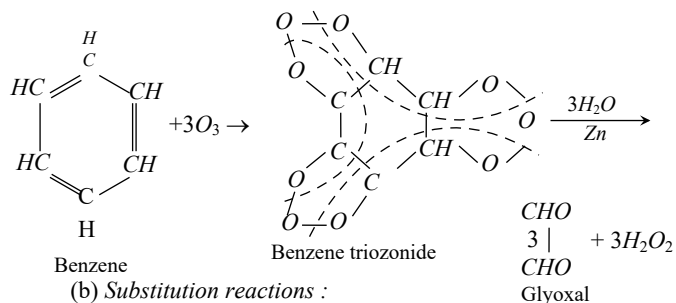
Addition of halogen :



• Electrophilic substitution reaction : Benzene undergoes this reaction because it is an electron rich system due to delocalized π -electrons.



Addition of ozone :

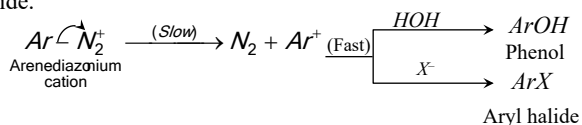


Benzenes

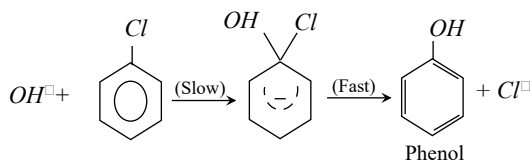
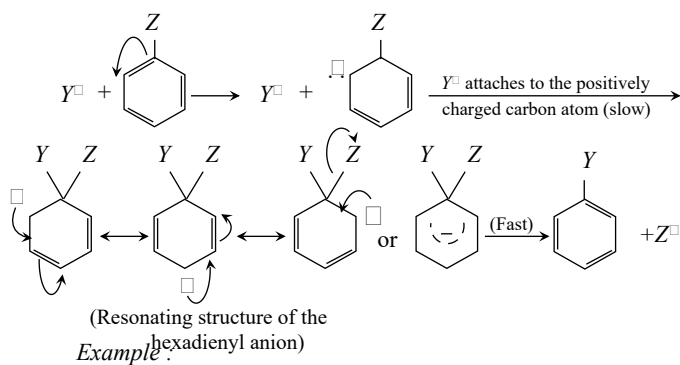
(b) Substitution reactions :

Nucleophilic substitution :

Unimolecular : Mostly uncommon in aromatic substitution, there is only one example which obtain in benzene diazonium dichloride.



• Bimolecular :



• Elimination-addition mechanism (Benzyne mechanism)

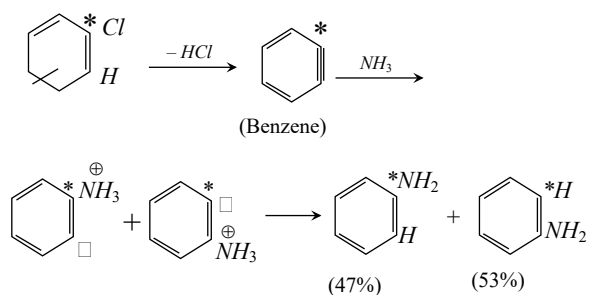
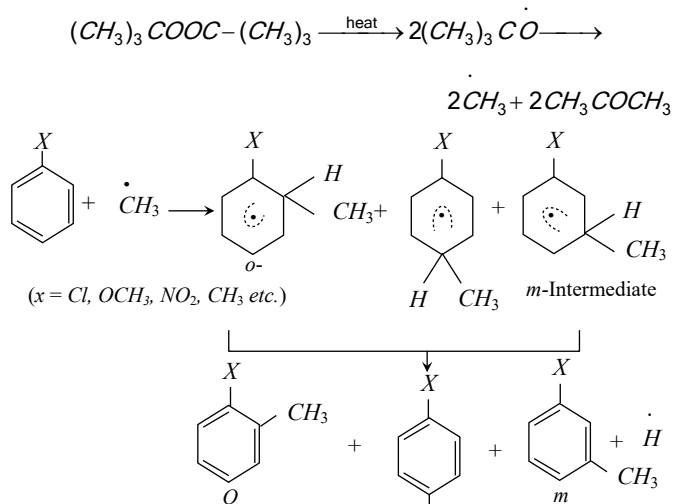


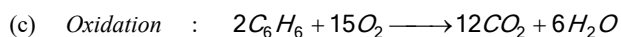
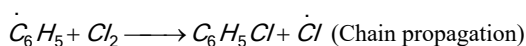
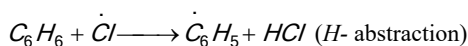
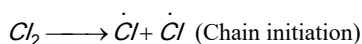
Table : 24.5

Electrophile (E^{\oplus})	Name	Source	Name of substitution reaction
Cl^+	Chloronium	$Cl_2 + AlCl_3$ or $FeCl_3$	Chlorination
Br^+	Bromonium	$Br_2 + AlBr_3$ or $FeBr_3$	Bromination
NO_2^+	Nitronium	$HNO_3 + H_2SO_4$	Nitration
SO_3	Sulphur trioxide	Conc. H_2SO_4 , Fuming sulphuric acid	Sulphonation
R^+	Alkyl carbonium	$RX + AlX_3$ ($X = Cl$ or Br), $ROH + H^+$	Friedel-Craft's (Alkylation)
$R-\overset{+}{C}=O$	Acyl carbonium	$RCOCl + AlCl_3$	Friedel-Craft's (Acylation)

• Free radical aromatic substitution : The aromatic substitution reactions which follow free radical mechanisms are very few and have limited synthetic value. But some typical example of these reactions are:

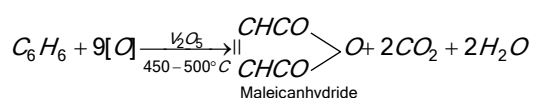


The mechanism of chlorination of benzene at high temperature is similar to that of the free radical aliphatic substitution



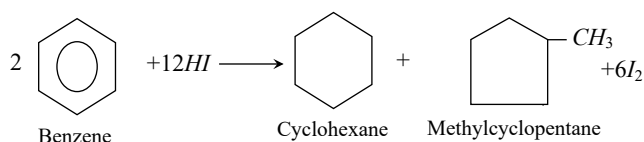
$$\Delta H = 6530 \text{ kJ/mole}$$

When vapours of benzene and air are passed over vanadium pentoxide at 450 – 500°C, maleic anhydride is obtained.



□ Strong oxidising agents converts benzene slowly into CO₂ and water on heating.

(d) Reduction :



(iii) **Uses** : (a) In dry cleaning (b) As a motor fuel when mixed with petrol. (c) As a solvent. (d) In the manufacture of gammexane (As insecticide). (e) In the preparation of nitrobenzene, chlorobenzene, benzene sulphonic acid, aniline, styrene, etc. Many of these are employed for making dyes, drugs, plastics, insecticides, etc.

Directive effect in substituted benzene derivatives

(1) **Directive effect in mono substituted benzene derivatives**

: The substituent already present on the benzene ring directs the

incoming substituent to occupy ortho (2 or 6), meta (3 or 5) or para (4) position. This direction depends on the nature of the first substituent and is called **directive or the orientation effect**.

The substituent already present can increase or decrease the rate of further substitution, *i.e.*, it either activates or deactivates the benzene ring towards further substitution. These effects are called **activity effects**.

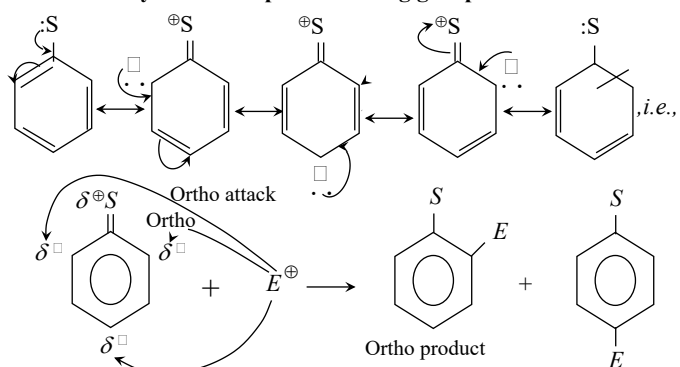
There are two types of substituents which produce directive effect are,

(i) Those which direct the incoming group to ortho- and para-positions simultaneously (Neglecting meta all together).

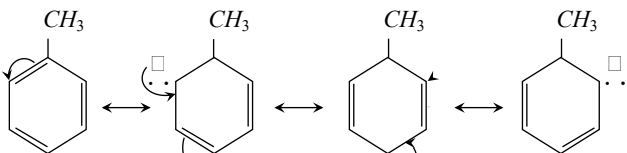
(ii) Those which direct the incoming group to meta-position only (Neglecting ortho- and para-positions all together).

Ortho-para directors	<i>Meta directors</i>
Strongly activating $-\ddot{N}H_2, -\ddot{N}HR, -\ddot{N}R_2, -\ddot{O}H, -\ddot{O}^-$	Moderately deactivating $-C \equiv N, -SO_3H, -COOH, -COOR, -CHO, COR$
Moderately activating $-\ddot{N}HCOCH_3, -\ddot{N}HCO R, -\ddot{O}CH_3, -\ddot{O}R$	Strongly deactivating $-NO_2, -NR_3^{\oplus}, -CF_3, -CCl_3$
Weakly activating $-CH_3, -C_2H_5, -R, -C_6H_5$ Weakly deactivating $-\ddot{F}, -\ddot{Cl}, -\ddot{Br}, -\ddot{I},$	

Theory of ortho - para directing group



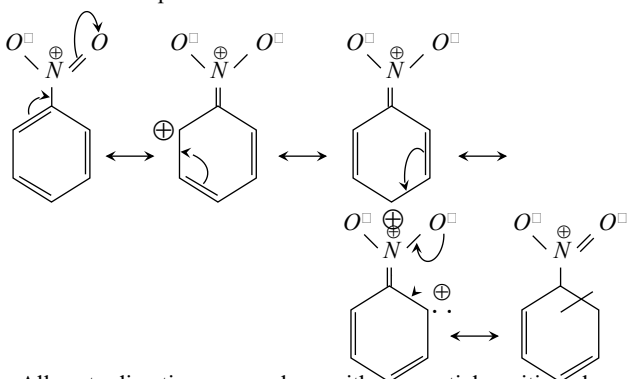
The above mechanism is followed when S is $-OH, -NH_2, -Cl, -Br, -I, -OR, -NR_2, -NHCOR$ etc.



In methyl or alkyl group, the $+I$ effect of the methyl group or alkyl group initiates the resonance effect.

Thus, methyl or alkyl group directs all electrophiles to ortho and para positions.

Theory of meta directing group : The substituent, S withdraws electrons from ortho and para positions. Thus, *m*-position becomes a point of relatively high electron density and further substitution by electrophile occurs at meta position. For example, $-NO_2$ group is a meta directing (Electron withdrawing). Its mechanism can be explained as :

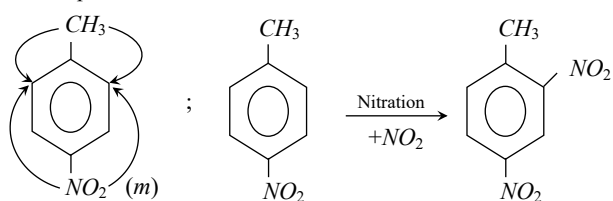


All meta-directing groups have either a partial positive charge or a full positive charge on the atom directly attached to the ring.

(2) Directive effect in disubstituted benzene

(i) If the directive effects of two substituents reinforce, then a single product is formed.

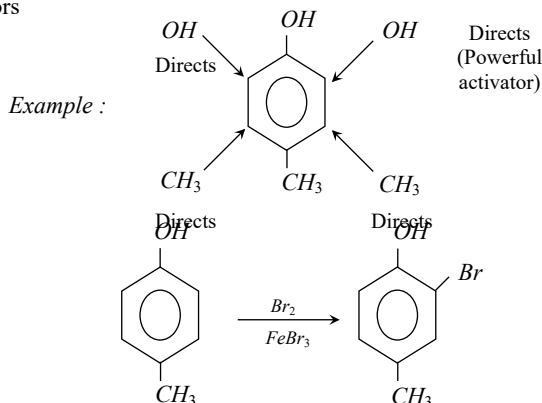
Example :



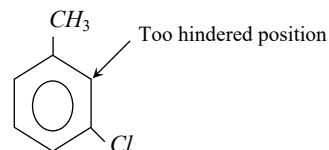
Thus, both (CH_3, NO_2) direct further substitution to the same position (Ortho with respect to CH_3).

(ii) If the directing effect of two groups oppose each other strongly activating groups win over deactivating or weakly activating group. The sequence of directing power is

$-NH_2 > -OH > -OCH_3 > -NHCOR > -C_6H_5 > CH_3 >$ meta directors



(iii) There is normally little substitution when the two groups are meta to each other. Aromatic rings with three adjacent substituents are generally prepared by same other routes.

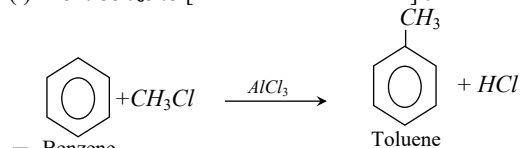


Toluene, methyl benzene or phenyl methane

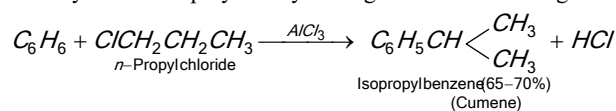
Toluene is the simplest homologue of benzene. It was first obtained by dry distillation of tolu balsam and hence named toluene. It is commercially known as toluol.

(1) Methods of preparation

(i) **From benzene** [Friedel-craft's reaction] :

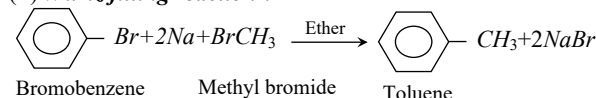


Alkyl halide employed may undergo an isomeric change

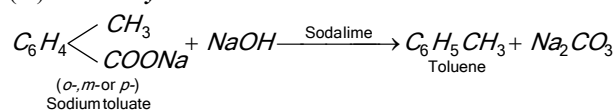


Catalysts can be used in place of anhydrous $AlCl_3$ are, $AlCl_3 > SbCl_3 > SnCl_4 > BF_3 > ZnCl_2 > HgCl_2$

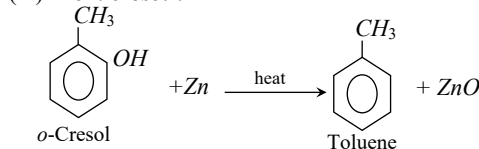
(ii) **Wurtz fitting reaction :**



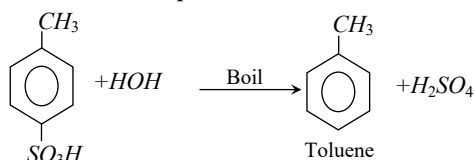
(iii) **Decarboxylation :**



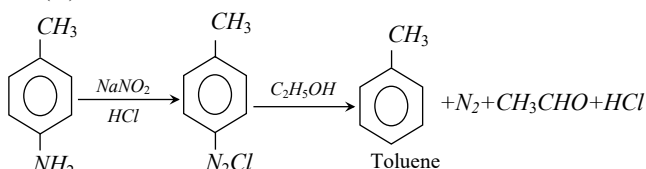
(iv) **From cresol :**



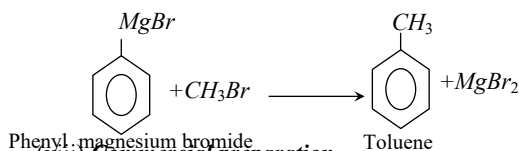
(v) **From toluene sulphonic acid :**



(vi) **From toluenamine :**



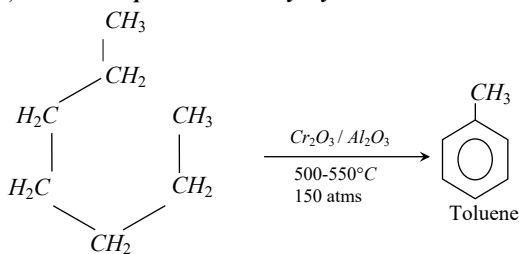
(vii) **From Grignard reagent :**



(viii) **Commercial preparation**

From coal tar : The main source of commercial production of toluene is the light oil fraction of coal-tar. The light oil fraction is washed with conc. H_2SO_4 to remove the bases, then with $NaOH$ to remove acidic substances and finally with water. It is subjected to fractional distillation. The vapours collected between $80-110^\circ C$ is 90% benzol which contains 70-80% benzene and 14-24% toluene. 90% benzol is again distilled and the portion distilling between $108-110^\circ C$ is collected as toluene.

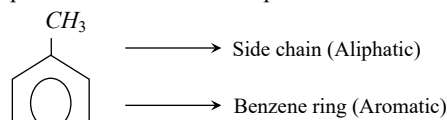
(ix) **From n-heptane and methyl cyclohexane**



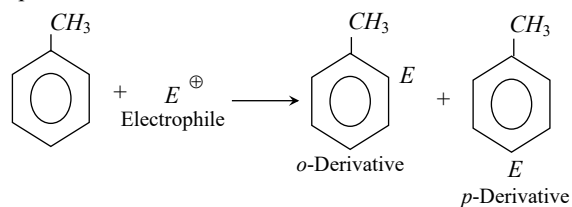
(2) **Physical properties**

- (i) It is a colourless mobile liquid having characteristic aromatic odour.
- (ii) It is lighter than water (*sp. gr.* 0.867 at $20^\circ C$).
- (iii) It is insoluble in water but miscible with alcohol and ether in all proportions.
- (iv) Its vapours are inflammable. It boils at $110^\circ C$ and freezes at $-96^\circ C$.
- (v) It is a good solvent for many organic compounds.
- (vi) It is a weak polar compound having dipole moment 0.4D.

(3) **Chemical properties :** Toluene shows the behaviour of both an aliphatic and an aromatic compound.



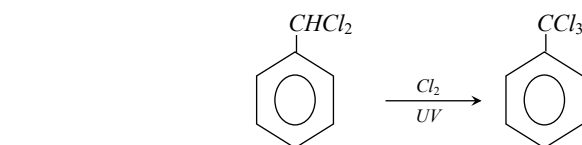
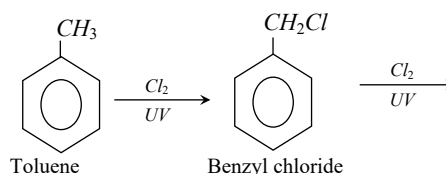
(i) **Electrophilic substitution reactions :** Aromatic character (More reactive than benzene) due to electron releasing nature of methyl group.



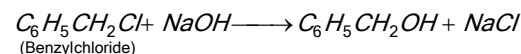
□ E^+ may be Cl^+ , NO_2^+ , SO_3^+ , H^+ etc.

(ii) **Reactions of side chain**

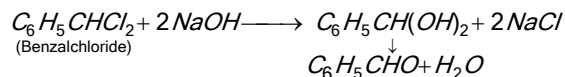
(a) **Side chain halogenation :**



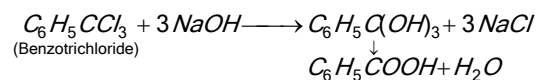
□ Benzyl chloride on hydrolysis with aqueous caustic soda forms benzyl alcohol.



□ Benzal chloride on hydrolysis forms benzaldehyde.

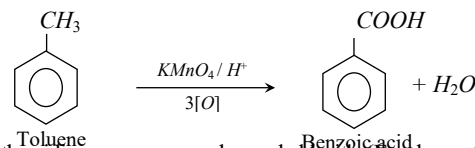


□ Benzo trichloride on hydrolysis forms benzoic acid.

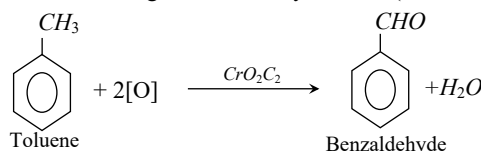


(b) **Oxidation :**

• With hot acidic $KMnO_4$:

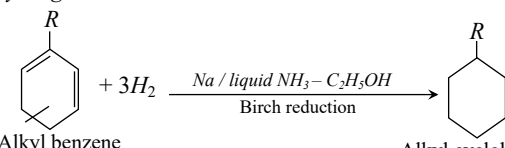


• With acidic manganese or chromyl chloride (Etard's reaction) :



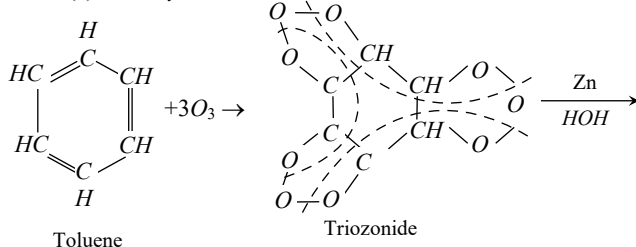
□ All alkyl benzenes on oxidation with hot acidic $KMnO_4$ or $Na_2Cr_2O_7$ form benzoic acid. The length of the side chain does not matter.

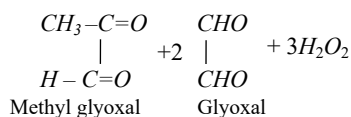
(c) **Hydrogenation :**



(d) **Combustion :** $C_6H_5CH_3 + 9O_2 \longrightarrow 7CO_2 + 4H_2O$

(e) **Ozonolysis :**





(4) Uses

(i) In the manufacture of benzyl chloride, benzal chloride, benzyl alcohol, benzaldehyde, benzoic acid, saccharin, etc.

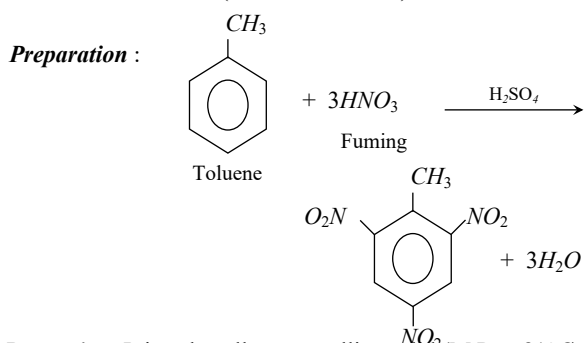
(ii) In the manufacture of trinitrotoluene (TNT), a highly explosive substance.

(iii) As an industrial solvent and in drycleaning.

(iv) As a petrol substitute.

(v) In the manufacture of certain dyes and drugs.

T.N.T. (Tri-nitro toluene)



Properties : It is pale yellow crystalline solid (M.P. = 81°C).

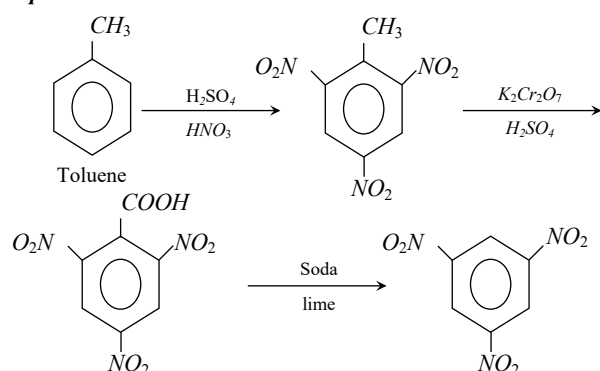
Uses : • It is used as an explosive in shells, bombs and torpedoes under the name **trotyl**.

• When mixed with 80% ammonium nitrate it forms the explosive **amatol**.

• TNT is also used as a mixture of aluminium nitrate, alumina and charcoal under the name **ammonal**.

T.N.B. (Tri-nitro benzene)

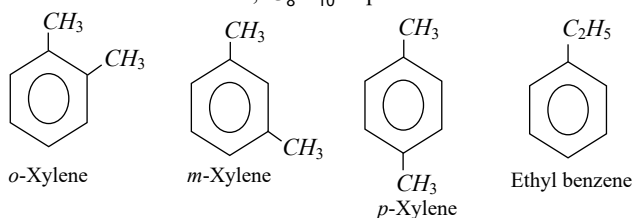
Preparation :



Properties and uses: It is colourless solid (M.P. = 122°C). It is more explosive than T.N.T. and used for making explosive.

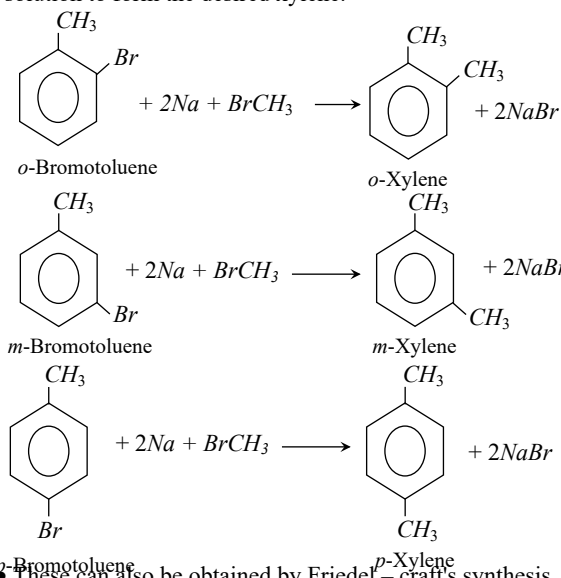
Xylenes (Dimethyl benzene) $\text{C}_6\text{H}_4(\text{CH}_3)_2$

The molecular formula, C_8H_{10} represents four isomers.



These are produced along with benzene, toluene and ethylbenzene when aromatisation of $\text{C}_6 - \text{C}_8$ fraction of petroleum naphtha is done. The xylenes are isolated from the resulting mixture (BTX) by fractional distillation.

These can be prepared by Wurtz – Fittig reaction. A mixture of bromotoluene and methylbromide is treated with sodium in dry ethereal solution to form the desired xylene.



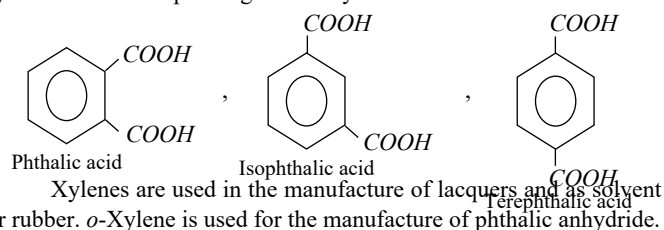
• These can also be obtained by Friedel-Craft's synthesis,

• *m*-Xylene can be obtained from mesitylene.

Xylenes are colourless liquids having characteristic odour. The boiling points of three isomers are,

o-Xylene=144°C; *m*-Xylene=139°C; *p*-Xylene=138°C.

Xylenes undergo electrophilic substitution reactions in the same manner as toluene. Upon oxidation with KMnO_4 or $\text{K}_2\text{Cr}_2\text{O}_7$, Xylenes form corresponding dicarboxylic acids.

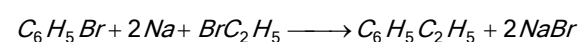


Xylenes are used in the manufacture of lacquers and as solvent for rubber. *o*-Xylene is used for the manufacture of phthalic anhydride.

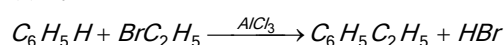
Ethyl benzene ($\text{C}_6\text{H}_5\text{C}_2\text{H}_5$)

It can be prepared by the following reactions,

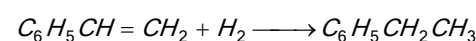
(1) By Wurtz-Fittig reaction :



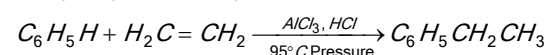
(2) By Friedel-Craft's reaction :



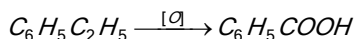
(3) By catalytic reduction of styrene :



(4) By alkyl benzene synthesis :



It undergoes electrophilic substitution reactions in the same way as toluene. When oxidised with dil. HNO_3 or alkaline $KMnO_4$ or chromic acid it forms benzoic acid.

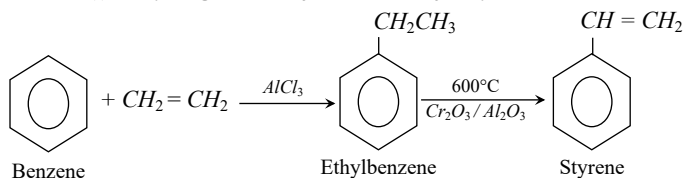


Styrene ($C_6H_5CH=CH_2$)

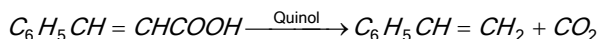
It is present in storax balsam and coal-tar in traces.

(1) Preparation

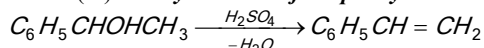
(i) Dehydrogenation of side chain of ethylbenzene :



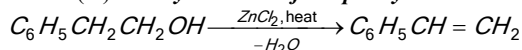
(ii) **Decarboxylation of cinnamic acid** : This is the laboratory preparation and involves heating of cinnamic acid with a small amount of quinol.



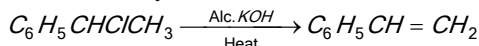
(iii) Dehydration of 1-phenyl ethanol with H_2SO_4 :



(iv) Dehydration of 2-phenyl ethanol with $ZnCl_2$:

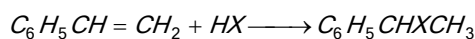
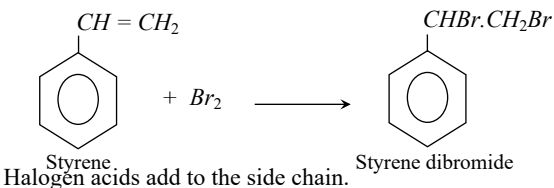
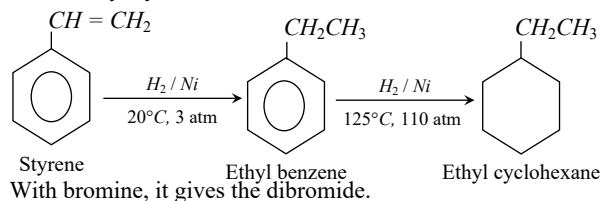


(v) **Dehydrohalogenation of 1-phenyl-1-chloro ethane** : On heating with alcoholic potassium hydroxide, a molecule of hydrogen chloride is eliminated by the chloro derivative.

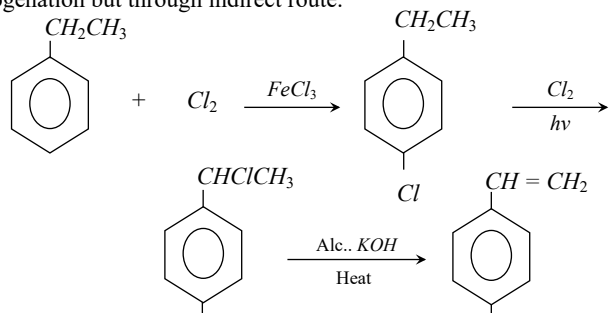


(2) **Properties** : It is a colourless liquid, boiling point $145^\circ C$. On keeping, it gradually changes into a solid polymer called **metastyrene**. The polymerisation is rapid in sunlight or when treated with sodium. It shows properties of benzene ring (Electrophilic substitution) and unsaturated side chain (Electrophilic addition). However, the side chain double bond is more susceptible to electrophilic attack as compared to benzene ring.

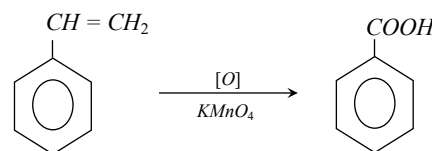
At lower temperature and pressure, it reacts with hydrogen to produce ethylbenzene and at higher temperature and pressure, it is converted into ethyl cyclohexane.



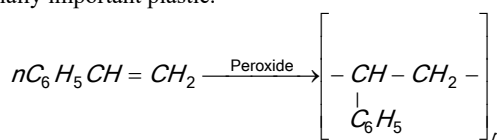
Preparation of ring substituted styrenes is not done by direct halogenation but through indirect route.



When oxidised under drastic conditions, the side chain is completely oxidised to a carboxyl group.



In presence of peroxides, styrene undergoes free radical polymerisation resulting in the formation of polystyrene – an industrially important plastic.



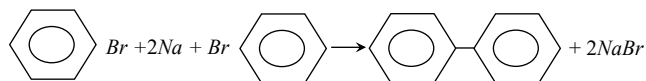
Co-polymers of styrene with butadiene and other substances are also important since many of them are industrially useful products such as **SBR** (A rubber substitute).

Bi-phenyl ($C_6H_5 - C_6H_5$)

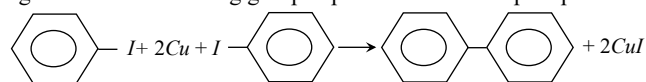
It occurs in coal-tar. It is the simplest example of an aromatic hydrocarbon in which two benzene rings are directly linked to each other.

(1) Methods of formation

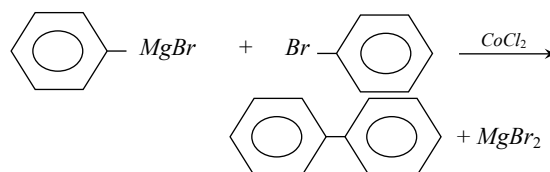
(i) **Fittig reaction** : It consists heating of an ethereal solution of bromobenzene with metallic sodium.



(ii) **Ullmann biaryl synthesis** : Iodobenzene, on heating with copper in a sealed tube, forms biphenyl. The reaction is facilitated if a strong electron withdrawing group is present in ortho or para position.

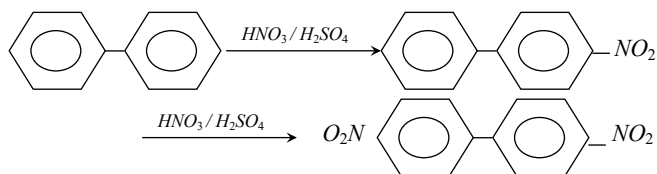


(iii) **Grignard reaction** : Phenyl magnesium bromide reacts with bromo benzene in presence of $CoCl_2$.



(2) **Properties** : It is a colourless solid, melting point $71^\circ C$. It undergoes usual electrophilic substitution reactions. Since aryl groups are electron withdrawing, they should have deactivating and *m*-orientating effect. But, it has been experimentally shown that presence of one benzene ring activates the other for electrophilic substitution and directs the incoming group to *o*- and *p*- positions. It has been shown that monosubstitution in the bi-phenyl results in the formation of para isomer as the major product.

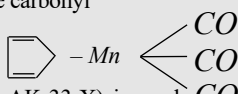
Another special feature of the biphenyl is the behaviour towards second substitution in a monosubstituted biphenyl. The second substituent invariably enters the unsubstituted ring in the ortho and para position no matter what is the nature of substituent already present.



Tips & Tricks

✍ Octane number may be less than zero (e.g., *n*-Nonane has an octane number-45) and higher than 100 (e.g., Triptane or 2, 3, 3-Trimethylbutane has an octane number of 124).

✍ To avoid lead pollution, a new compound cyclopentadienyl manganese carbonyl

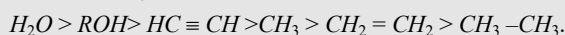


(called as AK-33-X) is used as antiknock now a days in developed countries (unleaded petrol).

✍ Acetylene has a garlic odour when impure due to impurities of phosphine and hydrogen sulphide.

✍ Fluorination is a violent reaction and can be controlled by diluting fluorine with nitrogen.

✍ The relative acidic character of water, alcohols acetylene, ammonia, ethylene and ethane follows the order :



Obviously, the basic character of their conjugate bases follows the reverse order, i.e.,



✍ *Wilkinson's catalyst* : (Triphenylphosphine) rhodium, $(\text{PPh}_3)_3\text{RhCl}$ is called wilkinson's catalyst. It reduces alkenes and alkynes while other common functional groups such as $\text{C}=\text{O}$, NO_2 and $\text{C} \equiv \text{N}$ remain unaffected,

✍ The order of reactivity of primary (1°), secondary (2°) and tertiary (3°) hydrogens in alkanes follows the sequence : $3^\circ > 2^\circ > 1^\circ$.