

Hydrocarbons

FAST TRACK: QUICK REVISION

Hydrocarbons are the organic compounds containing carbon and hydrogen only. Depending upon the types of carbon-carbon bonds present, hydrocarbons can be classified into three categories- (i) Saturated (ii) Unsaturated (iii) Aromatic hydrocarbons.

Saturated hydrocarbons contain carbon-carbon muiltiple bonds—double bonds, triple bonds or both.

ALKANES: Saturated open chain hydrocarbons containing carbon-carbon single bonds. These are inert under normal conditions i.e. do not react with acids, bases and other reagents. Alkanes exhibit Chain isomerism, Position isomerism and conformational isomerism.

General methods of preparation of alkanes:

1. From Unsaturated hydrocarbons : By hydrogenation in the presence of platinium, palladium or nickel as catalyst.

General Chemical Equation:

$$R-CH=CH_2+H_2 \xrightarrow{Ni} R-CH_2-CH_3$$

[Where R is H or Alkyl group]

e.g.
$$CH_2 = CH_2 + H_2 \xrightarrow{Ni} CH_3 - CH_3$$

Ethane
$$CH_3 - C \equiv CH + H_2 \xrightarrow{Ni} CH_3 - CH = CH_2 + H_2 \xrightarrow{Ni} CH_3 - CH_2 - CH_3$$
Propene Propane

2. From alkyl halides: on reduction with Zinc and dilute hydrochloric acid

$$\begin{array}{ccc} \text{CH}_3-\text{Cl} + \text{H}_2 & \xrightarrow{Zn, \, \text{H}^+} & \text{CH}_4 + \text{HCl} \\ \text{Chloromethane} & \text{Methane} \\ \text{C}_2 \, \text{H}_5-\text{Cl} + \text{H}_2 & \xrightarrow{Zn, \, \text{H}^+} & \text{C}_2 \text{H}_6 + \text{HCl} \\ \text{Chloroethane} & \text{Ethane} \end{array}$$

$$\begin{array}{ccc} \text{CH}_3\text{CH}_2\text{CH}_2\text{Cl} + \text{H}_2 & \xrightarrow{\text{Zn, H}^+} & \text{CH}_3\text{CH}_2\text{CH}_3 + \text{HCl} \\ \text{1-Chloropropane} & & \text{Propane} \end{array}$$

3. From alkyl halides by Wurtz reaction : Reaction of alkyl halide with sodium in dry ether, useful only for the preparation of symmetrical alkanes.

$$CH_{3}CH_{2}Br + 2Na + Br CH_{2}CH_{3} \xrightarrow{Dry Ether} CH_{3}CH_{2}CH_{2}CH_{3}$$
Bromoethane
$$Butane$$

$$\begin{array}{c} \text{CH}_3\text{CH}_2\text{CH}_2\text{Br} + 2\text{Na} + \text{Br}\text{CH}_2\text{CH}_2\text{CH}_3 \xrightarrow{\text{Dry Ether}} \text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3 \\ \text{1-Bromopropane} \end{array}$$

$$\begin{array}{c} \text{CH}_3\text{CH}_2\text{ Br} + 2\text{Na} + \text{Br CH}_2\text{CH}_3 & \xrightarrow{\text{Dry Ether}} & \text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_3 \\ \text{1-Bromopropane} & \text{1-Bromoethane} & \text{n-Pentane (desired)} \end{array}$$

4. From Carboxylic acids : By decarboxylation with soda lime

5. By Kolbe's electrolytic method : Electrolysis of an aqueous solution sodium or potassium salt of carboxylic acid. Alkane containing even number of carbon atoms is formed at anode.

$$2CH_3COO Na + 2H_2O \xrightarrow{Electrolysis} CH_3 - CH_3 + 2CO_2 + H_2 + 2NaOH$$

At Anode : (Oxidation)

$$2CH_{3} - \overset{O}{C} - O^{-} \xrightarrow{-2e^{-}} 2CH_{3} - \overset{O}{C} - O^{-} \longrightarrow 2CH_{3}^{\bullet} + 2CO_{2}$$

$$2CH_{3}^{\bullet} \longrightarrow CH_{3} - CH_{3}$$

At Cathode: (Reduction)

$$2H_2O \longrightarrow 2OH^- + 2H^+$$

$$2H^+ + 2e^- \longrightarrow H_2$$

Physical Properties of alkanes:

- 1. Boiling point of alkanes decreases on branching due to decrease in surface area of molecule with branching which decreases magnitude of van der Waal's forces of attraction.
- 2. Alkanes being non-polar in nature are soluble in non-polar solvents.

Chemical properties of Alkanes:

• Alkanes undergo substitution reactions. *e.g.*, Halogenation, Nitration, Sulphonation.

Halogenation: For example Chlorination of methane

Rate of reaction of alkanes with halogens is $F_2 > Cl_2 > Br_2 > I_2$. Rate of replacement of hydrogen in alkanes is $3^{\circ} > 2^{\circ} > 1^{\circ}$. Fluorination is too violent to be controlled. Iodination is reversible and it is therefore carried out in the presence of oxidising agent like HNO₃.

Mechanism of halogenation: Free radical mechanism

$$Cl - Cl \xrightarrow{h\upsilon} Cl \bullet + \bullet Cl \qquad \text{Initiation}$$

$$Cl \bullet + CH_4 \longrightarrow H - Cl + \bullet CH_3 \qquad \text{Propagation}$$

$$\bullet CH_3 + Cl - Cl \longrightarrow CH_3 Cl + \bullet Cl \qquad \text{Propagation}$$

$$Cl \bullet + \bullet CH_3 \longrightarrow CH_3 Cl \qquad \text{Termination}$$

Combustion: Complete combustion gives carbon dioxide and water.

$$CH_4 + 2O_2 \longrightarrow CO_2 + 2H_2O$$

Isomerisation:

$$CH_3 CH_2CH_2CH_3 \xrightarrow{AlCl_3 + HCl} CH_3 CHCH_3$$

$$CH_3 CH_2CH_2CH_3 \xrightarrow{200^{\circ} \text{ C}, 35 \text{ atm}} CH_3 CHCH_3$$
Isobutane

Aromatisation:

(vi) **Reaction with steam**

$$CH_4 + H_2O \xrightarrow{Ni} CO + 3H_2$$

Pyrolysis: Decomposition of higher alkanes to lower alkanes on heating.

$$C_6H_{14}$$
 773K $C_6H_{12} + H_2$ $C_4H_8 + C_2H_6$ $C_3H_6 + C_2H_4 + CH_4$

Conformations:

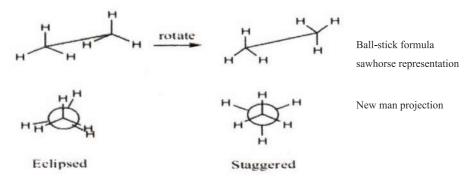
Different spatial arrangement of atoms arising due to rotation around C-C single bond.

Conformation of ethane, CH3CH3

Two conformational isomers or conformers.

Eclipsed form = all hydrogen atoms nearest to each other.

Staggered form = all hydrogen atoms are farthest apart.



Stability of eclipsed conformation is least while staggered conformation is most stable. The energy difference between two extreme forms is 12.5kJmol⁻¹. Due to this small energy difference the two forms are easily inter-convertible at ordinary temperature and cannot be separated and isolated.

ALKENES

These are unsaturated non-cyclic hydrocarbons which have sp^2 -hybridisation with 120° bond angle.

Alkenes are also called olefins [oil.forming] which indicates their high reactive nature.

Alkenes have general formula $C_n H_{2n}$, where $n = 2, 3, 4, \dots$ C_2H_4 (ethene), C_3H_6 (propene), etc.

Methods of Preparation of Alkenes

(i) From alkynes

$$R - C = C - R' + H_2 \xrightarrow{Pd/C} \xrightarrow{R} C - C \xrightarrow{R}$$
cis-alkene

(ii) From alkyl halide [Dehydrohalogenational]

(iii) From vicinal dihalides

(iv) From alcohols by acidic dehydrogenation

• Chemical Properties of Alkenes:

1. Addition of Halogens:

1, 2-Dichloropropane

2. Addition of hydrogen halides HCl, HBr, Hl: Add up to alkenes to form alkyl halides as per their reactivity order in HI>HBr>HCl.

Addition reaction of HBr to unsymmetrical alkenes (Markownikov's rule) According to Markownikovs' rule, the negative part of the addendum (adding molecule) gets attached to that carbon atom which possesses lesser number of hydrogen atoms.

Anti Markownikov addition or peroxide effect or Kharasch effect in the presence of organic peroxide, addition of only HBr molecule on unsymmetrical alkene takes place contrary to the Markownikov's rule. Peroxide effect is not observed in case of HF, HCl and HI.

$$CH_3 - CH = CH_2 + HBr \xrightarrow{(C_6H_5CO)_2O_2} CH_3CH_2CH_2Br$$
1-bromopropane

3. Addition of sulphuric acid

4. Addition of water

5. Oxidation : Alkenes decolourise cold dilute aqueous solution potassium permanganate (Baeyer's reagent). It is used as a test for unsaturation.

Acidic KMnO₄ or acidic K₂Cr₂O₇ oxidises alkenes to ketones and/or acids depending upon the nature of alkene and the experimental conditions.

$$(CH_3)_2C=CH_2 \xrightarrow{KMnO_4/H^+} (CH_3)_2CO + CO_2 + H_2O$$
2-Methyl propene Propane-2-one
$$CH_3-CH=CH-CH_3 \xrightarrow{KMnO_4/H^+} 2CH_3COOH$$
But-2-ene Ethanoic acid

6. Ozonolysis: Reaction of ozone with alkene to form ozonide which on subsequent reductive cleavage with zinc dust and water give carbonyl compounds (aldehydes & ketones).

$$H_3C$$
 $C=CH_2 + O_3 \xrightarrow{Zn/H_2O} C=O + HCHO + H_2O_2$
 H_3C
 $Zn + H_2O_2 \longrightarrow ZnO + H_2O$

7. Polymerization

$$n(\text{CH}_2 = \text{CH}_2) \xrightarrow{\text{High temperature/pressure} \atop \text{catalyst}} -[\text{CH}_2 - \text{CH}_2 -]_n \\ \text{polythene}$$

$$n(\text{CH}_3 - \text{CH} = \text{CH}_2) \xrightarrow{\text{High temperature/pressure} \atop \text{catalyst}} \left[-\text{CH} - \text{CH}_2 - \right]_n$$

ALKYNES

These are unsaturated hydrocarbons with general formula C_nH_{2n-2} e.g., C_2H_2 (ethyne), C_3H_4 (propyne).

Alkynes also exhibit electrophilic addition reaction but less reactive than alkenes because the dissociation of π -electron cloud requires more energy.

H–C≡C–H contins 3σ and 2π -bonds and bond length is 120 pm. In acetylene. H–C–C bond angle is 180°.

Methods of Preparation of Alkynes

1. From calcium carbide

$$CaCO_3 \longrightarrow CaO + CO_2$$

 $CaO + C \longrightarrow CaC_2 + CO$
 $CaC_2 + 2H_2O \longrightarrow Ca(OH)_2 + C_2H_2$

2. From vicinal dihalides

3. From tetrahalides

$$Br_2CH$$
— $CHBr_2 + 2Zn$ $\xrightarrow{CH_3 OH}$ H — $C \equiv C$ — $H + 2ZnBr_2$

Physical Properties of Alkynes:

- 1. The first two members are gases next eight members $(C_5 C_{12})$ are liquids and higher members are solids.
- 2. They are all colourless and odourless with the exception of acetylene which has slightly garlic odour due to the presence of PH₃ and H₂S as impurities.
- 3. Alkynes are insoluble in water but soluble in organic solvents like ethers, carbon tetrachloride and benzene.
- 4. Melting point, boiling point and density increase with increase in molar mass.

Chemical properties of Alkynes

Alkynes show electrophilic as well as nucleophilic addition reactions.

(i) Acidic character of alkyne

$$HC \equiv CH + Na \longrightarrow HC \equiv \overline{C}Na + \frac{1}{2}H_{2}$$

$$monosodium$$

$$acetylides$$

$$H - C \equiv CNa^{+} + Na \longrightarrow Na^{+}C \equiv \overline{C}Na^{+} + \frac{1}{2}H_{2}$$

$$disodium acetylide$$

$$CH_{3} - C \equiv C - H + NaNH_{2} \longrightarrow CH_{3} - C \equiv \overline{C}Na^{+} + NH_{3}$$

$$sodium propynide$$

These reactions are not shown by alkenes, alkanes and non-terminal alkynes, hence used for distinction between alkane, alkene and alkyne.

Acetylenic hydrogens are acidic in nature due to 50% s-character in sp-hybridised orbitals. Acidity of alkynes is lesser than water.

Acidic behaviour order

(i)
$$HC \equiv CH > CH_2 = CH_2 > CH_3 - CH_3$$

 sp sp^2 sp^3

(ii)
$$HC \equiv CH > CH_3 - C \equiv CH >> CH_3 - C \equiv C - CH_3$$

(ii) Electrophilic addition reactions

$$-C \equiv C - + HZ \xrightarrow{H^+} -C \equiv C - + : Z^- \longrightarrow -C \equiv C -$$
vinyl cation

The addition product formed depends upon the stability of vinylic cation. Addition on unsymmetrical alkynes takes place according to Markovnikov's rule.

$$CH_3 - C = C - H + H_2 \xrightarrow{Pt/Pd/Ni} [CH_3 - CH = CH_2] \xrightarrow{H_2} CH_3CH_2CH_3$$
propene

Addition of halogens

$$\begin{split} HC &\equiv CH + Cl \longrightarrow \begin{bmatrix} HCCl = CHCl \end{bmatrix} \\ 1, 2\text{-Dichloropropene} \\ Cl_2 \downarrow \\ Cl & Cl \\ | & | \\ HC - CH \\ | & | \\ Cl & Cl \\ 1, 1, 2, 2\text{-Tetrachloroethane} \\ or westron \\ \downarrow & -HCl \\ CH = CCl_2 \\ | & Cl \\ Cl \\ \end{split}$$

westrosol (1, 1, 2-Trichloroethene)

Addition of hydrogen halides

Addition for water

CH₃—C
$$\equiv$$
CH + HOH $\xrightarrow{\text{Hg}^{2+}/\text{H}^+}$ CH₃—C \equiv CH₂ \rightleftharpoons CH₃—C—CH₃ propyne O—H

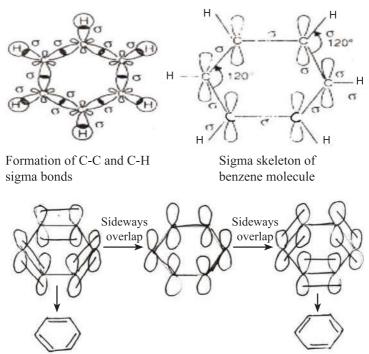
(iii) Cyclic polymerisation of ethyne

AROMATIC HYDROCARBONS

These hydrocarbon are also known as arenes. The parent member of the family aromatic hydrocarbons is benzene.

Aromatic compounds containing benzene ring are known as benzenoids.

Structure of benzene : Hexagonal ring of carbon atoms with alternate single and double bonds. Each carbon atom is sp^2 hybridised. Planar ring, bond angle 120° . All C-C bond lengths are equal due to complete delocalisation of π electrons.



HUCKEL'S RULE

- Huckel's rule, (based on calculations): a planar cyclic molecule with alternating double and single bonds has aromatic stability if it has $(4n + 2)\pi$ electrons (n is 0, 1, 2, 3, 4)
- For n = 1 : 4n+2 = 6; benzene is stable and electrons are delocalized.



Three double bonds;

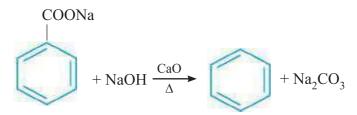
Benzene

 $\sin \pi$ electrons

METHODS OF PREPARATION

1. Cyclic polymerisation of ethyne

2. Decarboxylation of aromatic acids



3. Reduction of phenol

$$\begin{array}{c}
\text{OH} \\
\hline
\end{array}$$

$$\begin{array}{c}
\text{Zn} \\
\end{array}$$

$$+ \text{ZnC}$$

Physical Properties of Benzene:

- (i) Aromatic hydrocarbons are non-polar molecules and are usually colourless liquids or solids with a characteristic aroma.
- (ii) Aromatic hydrocarbons are immiscible with water but readily miscible with organic solvents.
- (iii) Aromatic compounds burn with sooty flame.

Chemical Reactions of Benzene:

- (i) Benzene gives electrophile substitution reactions.
- (ii) According to experimental evidences, electrophile substitution reaction involve following three steps:
 - Generation of electrophilie
 - Formation of carbocation intermediate.
 - Removal of proton from the carbocation intermediate.

(i) Nitration

(ii) Halogenation

(iii) Sulphonation

(iv) Friedel-Craft's alkylation reaction

(v) Friedel-Crafts acylation reaction

$$\begin{array}{c|c} & & & & & & & & & \\ & & & & & & & & \\ \hline & + \text{CH}_3\text{COCl} & & & & & & \\ & \text{or} & & & & & \\ & \text{(CH}_3\text{CO)}_2\text{O} & & & & \\ & & & & & & \\ \end{array}$$

Benzene also undergoes addition reactions e.g.

(i)
$$+3H_2 \xrightarrow{Ni}$$
 cyclohexane

Cl

Cl

Cl

Cl

Cl

Cl

Cl

benzene bexachloride or 666 (BHC or Gammexane or lindane)

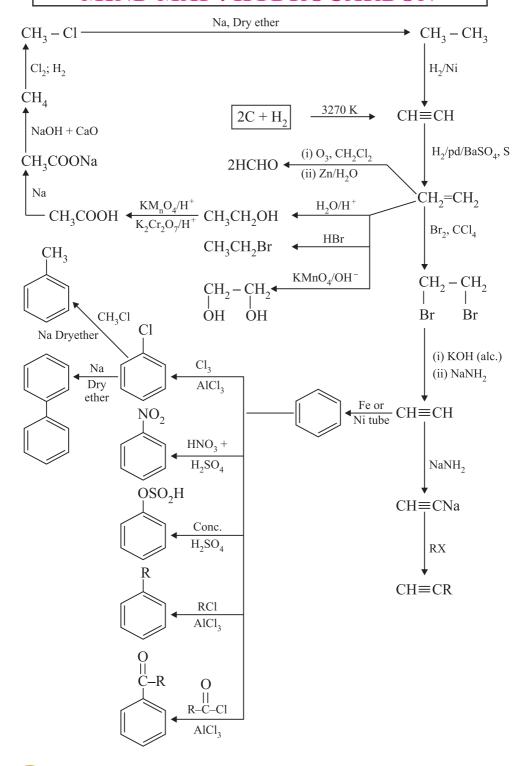
COMBUSTION

$$2C_6H_6 + 15O_2 \longrightarrow 12CO_2 + 6H_2O$$

Directing influence of substituents in monosubstituted benzene

- (i) Ortho and para directing groups: Ring activating groups e.g., NH_2 , $-CH_3$, $-C_2H_5$, $-OCH_3$ etc. (+R effect)
- (ii) **Meta directing groups :** Ring deactivating groups e.g. NO₂, CN, CHO, COOH, SO₃H (-R effect).

MIND MAP: HYDROCARBON



CASE BASED STUDY - QUESTIONS

PASSAGE-I

Presence of double bond in an alkene is tested by reacting with either water or with dilute solution of Bayer's regent. The position of double bond is located with the help of ozonalysis. It is done by joining together the product of ozonalysis which are carbonyl compounds at their carbonyl carbon atoms by double bond. One mole of ozone is used in the ozonalysis reaction per mole of double bond in a particular alkene.

- 1. An alkene upon ozonalysis will give CH₃CH₂ CHO and CH₃COCH₃. Give IUPAC name and structure of the given alkene.
- 2. Complete the given chemical equation.

$$CH_3 - CH = CH - CH_3 + O_3 \xrightarrow{R_2O} A$$

- 3. On ozonolysis one molecule of a hydrocarbon produces two molecules of ethanal and one molecule of ethanedial. Give the structure on ozonolysis of hydrocarbon.
- 4. An alkene give 5-keto-2-methyl hexanal. Give IUPAC name of the given alkene.

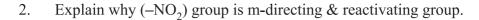
5. (a)
$$A \xrightarrow{O_3} CH_3CHO$$
 ?
(b) $A \xrightarrow{O_3} CH_3CHO + CH_3-C-CH_2-CH_3$

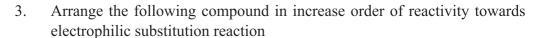
PASSAGE-II

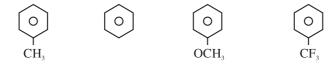
Benzene responds to electrophilic substitution reactions. However some reagents are needed in these reactions in order to generate the attaching electrohiles. The directive influence in disubstitution taking place in the ring depends upon the nature of the group already present. An activating group in creases the electron density at the ortho and para positions. The deactivating group decreases the same at these positions thereby healing the metal position comparatively a point of high electron density.

1. Complete the following chemical equation

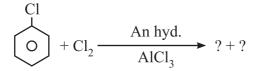




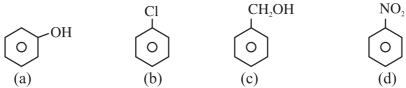




4. Write the product in the following reaction



5. Which of the following is most reactive to electrophilic attack.



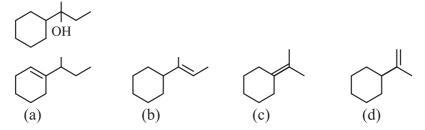
MULTIPLE CHOICE QUESTIONS (MCQ)

- 1. Which of the following has zero dipole moment?
 - (a.) cis-But-2-ene
- (b) trans-But-2-ene

(c) But-1-ene

- (d) 2-Methylprop-1-ene
- 2. Bond length of (I) ethane, (II) ethene, (III) Acetylene, (IV) Benzene follows the order:
 - (a) I > II > III > IV
- $(b) \quad I > II > IV > III$
- $(c) \quad I > IV > II > III$
- (d) III > IV > II > I
- 3. The methyl group in benzene ring is:
 - (a) Ortho directing
- (b) Ortho and meta directing
- (c) Para directing
- (d) Ortho and para directing

4. Which of th following is not the product of dehydration of



5. In the hydrocarbon:

$$6 5 4 3 2 1$$

 $H_3C - CH = CH - CH_2 - C \equiv CH$

The state of hybridization of carbon 1, 3 and 5 are in the order

(a) sp, sp 2 , sp 3

(b) sp^3 , sp^2 , sp

(c) sp^2 , sp, sp^3

(d) sp, sp 3 , sp 2

6. Action of acetylene on dilute H₂SO₄ gives:

(a) Acetic acid

- (b) Acetone
- (c) Acetaldehyde
- (d) Ethyl alcohol

7. Which of the following compounds will exhibit cis-trans (geometrical) isomerism?

(a) Butanol

(b) 2-Butyne

(c) 2-Butenol

(d) 2-Butene

8. Basic strength of:

 H_3C CH, H_2C = CH and $HC \equiv C$ is in the order of

(a) I > III > II

(b) I > II > III

(c) II > I > III

(d) III > II > I

9. Reaction of hydrogen bromide with propene in the absence of a peroxide is a/an

- (a) free radical reaction
- (b) nucleophilic substitution
- (c) electrophilic addition
- (d) nucleophilic substitution

10. Among the following compounds, the one which is most reactive towards electrophilic nitration is:

- (a) Benzoic acid
- (b) Nitrobenzene

(c) Toluene

(d) Benzene

ANSWERS: 1. b 2. c 3. d 4. a 5. d 6. c 7. d 8. c 9. c 10. c

FILL IN THE BLANKS

1.	The addition of HBr to an unsymmetrical alkene takes place in accordan				
	with rule.				
2.	Benzene favours	_ substitution react	ion.		
3.	The Dipole moment of Benzene is				
4.	The nitro group in the benzene r	nucleus is	_directing. It		
	the reactivity of the benzene ring.				
5.	Melting point and boiling point increase as the molar masses				
ANS	WERS : 1. Markownikov's	2. electrophilic	3. Zero		
	4. Meta, decreases	5. Increase			

TRUE AND FALSE TYPE QUESTIONS

- 1. Alkanes mainly undergo substitution reactions using the free-radical mechanism.
- 2. The decreasing order of boiling points among the isomeric pentanes is neo > iso > n.
- 3. The acidic character of three types of hydrocarbons follows the order alkanes > alkenes > alkynes.
- 4. The peroxide effect is observed only in addition of HBr, and not with HCl and HI.
- 5. Wurtz reaction is suitable for the preparation of both symmetrical and unsymmetrical alkanes.
- 6. For a compound to be aromatic it must have $(4n + 2)\pi$ electrons.
- 7. Benzene has planar structure.
- 8. The benzene molecule has two different carbon-carbon bond lengths, corresponding to alternate single and double bonds.
- 9. In Friedel-Crafts reaction, AlCl₃ is an electrophile.
- 10. An electron-donating substituent in benzene ring gives a meta product.

ANSWERS: 1. T 2. F 3. F 4. T 5. F 6. T 7. T 8. F 9. F 10. F

MATCH THE COLUMNS

Match the statements (a,b,c,d) in column I with the statements (i, ii, iii, iv) in column II.

1.	Column I	Column II	Column III	
	a. CH_4	i. sp^2	e. Ozonalysis	
	b. $CH_2 = CH_2$	ii. sp^3	f. Oxidising agent	
	c. $CH \equiv CH$	iii. sp³d	g. Saturated nature	
	d. PCl ₅	iv. sp	h. Un Saturated natur	e

2. Column I

Column II

- a. Alkanes
- i. Saturated nature
- b. Alkenes
- ii. Ozonolysis
- c. Alkynes
- iii. Geometrical isomerism
- d. Arenes
- iv. Aromatic character

ANSWERS: 1. a. \rightarrow ii. - g.

- $b. \rightarrow i.-h.-e.$
- $c. \rightarrow i.-h.-e.$
- $d. \rightarrow iii.-f.$

2. $a \rightarrow i$.

- $b. \rightarrow ii., iii.$
- $c. \rightarrow ii.$
- $d. \rightarrow i.$, iv.

ASSERTION-REASON TYPE QUESTIONS

Type 1. The questions given below consist of Assertion(A) and Reason (R). Use the following key to select correct answer.

- (a) If both assertion and reason are correct and reason is correct explanation for assertion.
- (b) If both assertion and reason are correct but reason is not correct explanation for assertion.
- (c) If assertion is correct but a reason is incorrect.
- (d) If assertion and reason both are incorrect.
- 1. **Assertion:** The IUPAC name of $CH_3CH = CH C \equiv CH$ is pent-3-en-1-yne and not pent-2-en-4-yne.

Reason: While deciding the locants of double and triple Bonds, lowest sum rule is always followed.

- 2. **Assertion:** Tropylium cation
 - **Reason:** The only property which decides the aromatic character is its planar nature.
- 3. **Assertion:** Friedel-craft reaction between benzene and acetic anhydride in the presence of anhydrous AlCl₃ yields acetophenone and not polysubstituted products.

Reason: Acetophenone formed poisons the catalyst preventing further reaction.

4. **Assertion:** But-1-ene on reacting with HBr in the presence of peroxide, products 1-bromobutane.

Reason: It involves the formation of a primary free radical.

5. **Assertion:** Cyclopentadienyl anion is aromatic in nature.

Reason: Cyclopentaclienyl anion has six π -electrons.

6. **Assertion:** Benzene reacts with chlorine in the form of light to form BHC.

Reason: BHC is also called gammexane or 666.

is aromatic in character.

- 7. **Assertion:** All the hydrogen atoms in $CH_2 = C = CH_2$ lie in one plane. **Reason:** All the carbon atoms in it are sp² hybridised.
- 8. **Assertion:** Propene reacts with HBr in the presence of benzoyl peroxide to yield 2-bromopropane.

Reason: In the presence of peroxide, the addition of HBr to propene follows ionic mechanism.

9. **Assertion:** Benzene does not decolourise bromine water.

Reason: Benzene is stabilised by resonance due to delocalisation of π electrons.

10. **Assertion:** Acidity of C-H bond decreases in the order:

$$HC \equiv CH > H_2C = CH_2 > H_3C - CH_3$$

Reason: Greater the percentage s-character, more is the acidity of C—H bond.

ANSWERS: 1.a 2.c 3.c 4.c 5.a 6.b 7.c 8.d 9.a 10.a

ONE WORD TYPE QUESTIONS

- 1. What is the state of hybridisation of Carbon atoms in alkanes?
- 2. What is the number of bonds in But-3-en-1-yne?
- 3. Name the product formed when Propyne is treated with aqueous H₂SO₄ in the presence of dil.HgSO₄.
- 4. Name the product formed when Benzene reacts with CH₃Cl in the presence of anhydrous aluminium chloride.
- 5. —COOH is ortho, para directing or Meta directing group?

1-MARK QUESTIONS

- 1. Write the reaction involved in Kolbe's electrolytic method to prepare ethane.
- 2. Define term decarboxylation.
- 3. Why dry ether and not water is used as a solvent in the preparation of alkane by Wurtz reaction?
- 4. Sodium salt of which carboxylic acid will be needed for the preparation of propane by decarboxylation method?
- 5. Complete the following reaction:

$$CH_3Cl + Na$$
 dry ether

- 6. Amongst the following which one has the maximum boiling point? *n*-Pentane, iso-pentane, neo-pentane.
- 7. Write IUPAC name of CH₃CH=CHCH₂CH=CCH₂CH=CH₂

 |
 CH₂CH₃
- 8. Draw the cis and trans isomers of CHCl=CHCl.
- 9. What happens when 2-Bromobutane is being treated with alc. KOH?
- 10. Name the reagents used to carry out the following conversions:

$$CH_3 - CH = CH_2 \rightarrow CH_3CH(OH)CH_2OH$$

11. Complete the following reaction:

$$CH_3 - CH = CH_2 + HBr$$
 Organic peroxide

- 12. An alkene A on ozonolysis gives a mixture of ethanol and pentan-3-one. Write IUPAC name of element.
- 13. When alkyne is treated with bromine water then what will be the colour of the product?
- 14. Why alkynes do not exhibit geometrical isomerism?
- 15. Complete the following reaction:

(i)
$$CH_3C \equiv CH \xrightarrow{H_2O, Hg^{2+}/H^+}$$
?

(ii)
$$CaC_2 + 2H_2O \longrightarrow +$$

- 16. How will you convert ethyne to benzene?
- 17. Write IUPAC name of $C_6H_5-CH_2-CH_2-CH=CH_2$.
- 18. Why is benzene extraordinarily stable although it contains three double bonds?
- 19. Write chemical reaction to exemplify Friedel-Crafts alkylation of benzene.
- 20. Why j is not aromatic?
- 21. C-C bond length in benzene is intermediate between C-C and C=C. Why?
- 22. Starting from benzene, how would you synthesize m- Bromonitrobenzene.
- 23. Give one example each of o, p-directing group and m-directing group.
- 24. Complete the reaction:

$$C_6H_6 + CH_3COC1$$
 Anhy. AlCl₃

2-MARKS QUESTIONS

- 1. What effect does branching have on the boiling point of an alkane and why?
- 2. An alkene A contains three C–C eight C–H and one C–C (π) bond upon ozonlysis A gives two moles of an aldehyde of molar mass 44u. Write the IUPAC name of A.
- 3. Write the structures of all the alkenes which upon hydrogenation give 2-methyl butane.
- 4. Wurtz reaction cannot be used for the preparation of unsymmetrical alkanes? Give reason.
- 5. How can ethene be prepared from (i) Ethanol (ii) Ethyl bromide?
- 6. Melting point of cis-But-2-ene is lower than that of trans-But-2-ene. Give reason.
- 7. Draw the structures of cis and trans Hex-2-ene.
- 8. Explain with the help of equation : Ozonolysis of propene.
- 9. What do you understand by peroxide effect (Kharasch effect)?
- 10. What factor determines the stability of alkene?
- 11. Out of benzene, *m*-dinitro benzene and toluene, which will undergo nitration most easily and why.
- 12. Complete the reaction:

(i)
$$+ HBr \rightarrow ?$$

(ii) $+ O_3 \xrightarrow{CH_3COOH} ? + ?$

13. An alkene on treatment with HBr in presence of peroxide can generate two types of free radicals $CH_3 - C \cdot - CH_2$ -Br and $CH_3 - C(CH_3) - CH_2 \cdot$ CH_3 Br

Predict the final product of the reaction and give reason.

(Hint: Stability of free radicals)

- 14. What happens when But-2-ene reacts with acidified potassium permanganate solution?
- 15. You are provided with But-2-yne, how will you convert it into:
 - (i) cis-But-2-ene
 - (ii) trans-But-2-ene

- An alkene C₄H₈ reacts with HBr both in the presence and absence of peroxide 16. to give the same product. Identify the alkene.
- 17. Arrange ethane, ethene and ethyne in the order of increasing acidity.
- 18. Identify A and B in the following reaction:

A
$$\xrightarrow{\text{Na}}$$
 CH=CH $\xrightarrow{\text{Red hot iron tube}}$ B

Write the structures of the products A and B of the following reactions:

(i)
$$HC \equiv CH \xrightarrow{Na} A \xrightarrow{CH_3Br} B$$

(ii)
$$BrH_2C-CH_2Br \xrightarrow{Alc.KOH} A \xrightarrow{NaNH_2} B$$

3-MARKS QUESTIONS

- 1. Write the structures and name of products obtained in the reaction of sodium with a mixture of 1-Iodo-2-methylpropane and 2-Iodopropane.
- State Markownikov's rule. Using this rule, write the reaction of propene 2. with (i) HCl & (ii) H₂O.
- Complete the following reactions: 3.

(i)
$$CH_3CH_2Br \xrightarrow{Alc.KOH}$$

(ii)
$$CH_3CH = CH_2 + O_3 \xrightarrow{Zn/H_2O}$$

(i)
$$CH_3CH_2Br$$
 $\xrightarrow{Alc.KOH}$ (ii) $CH_3CH = CH_2 + O_3$ $\xrightarrow{Zn/H_2O}$ $\xrightarrow{Dil.KMnO_4}$

- (i) Write the structure of 3, 4-Dimethylhept-3-ene. 4.
 - (ii) Name the compounds obtained by ozonolysis of 3-Methylpent-2-ene.
- Complete the following reactions: 5.

(i)
$$CH \equiv CH \xrightarrow{NaNH_2, CH_3Br}$$
 ?

(i)
$$CH \equiv CH \xrightarrow{NaNH_2, CH_3Br}$$
?
(ii) $CH \equiv CH \xrightarrow{H_2O, HgSO_4/H_2SO_4}$?

(iii)
$$CH_3C = CH + H_2 \xrightarrow{Pt} ? \xrightarrow{H_2} ?$$

- Write the mechanism of nitration of benzene. 6.
- 7. Arrange in the order of decreasing relative reactivity with an electrophile and explain:

Toluene, p-Nitrotoluene, 1, 4-Dinitrobenzene

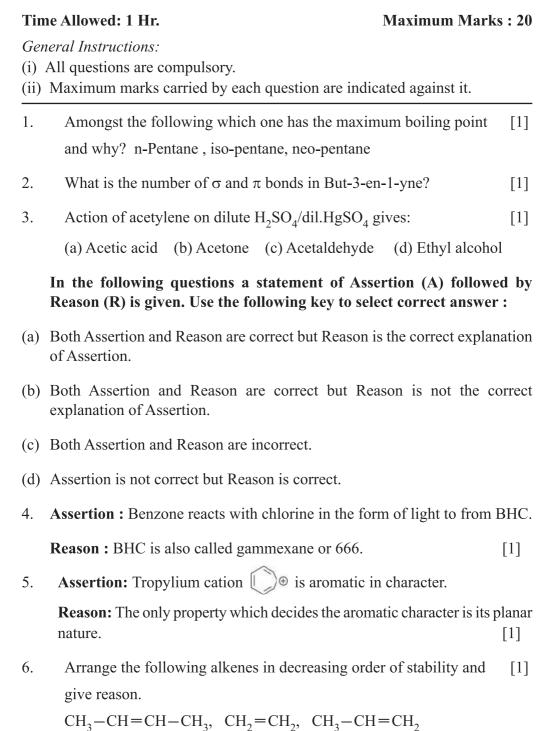
What is meant by delocalization of π electrons? How does it affect stability 8. of benzene?

- 9. What are the conditions for a compound/species to be aromatic according to Huckel's rule?
- 10. How will you convert benzene into
 - (i) Acetophenone
 - (ii) m- Chloronitrobenzene?

5-MARKS QUESTIONS

- 1. Define isomerism. Write all the structural isomers of hexane(C_6H_{14}) and arrange them in increasing order of boiling points.
- 2. Write short note on (i) Wurtz reaction (ii) Kolbe's electrolysis (iii) Ozonolysis
- 3. An alkene 'A' of molecular mass '28u' on treatment with bromine gives a product 'B'. The Compound 'B' on further dehalogenation with zinc gives back 'A'. Give the structures of 'A' and 'B' and also the sequence of reactions.
- 4. An organic compound 'A' with formula C_4H_9Br on treatment with KOH(alc.) gave two isomeric compounds 'B' and 'C' with formula C_4H_8 . Ozonolysis of 'B' gave only one product CH3CHO while 'C' gave two different products. Identify A, B and C.
- 5. How will you convert Ethyne into (i) 1, 1, 2, 2-Tetrachloroethane (ii) Ethene (iii) Ethanal (iv) Benzene (v) Sodium ethynide

UNIT TEST-I



- 7. (i) Give a chemical test to distinguish between ethyne and ethene. [2]
 - (ii) Melting point of cis-But-2-ene is lower than that of trans-But-2-ene. Give reason.
- 8. Complete the following reactions: [3]
 - (i) CH₃CH₂Br Alc.KOH
 - (ii) $CH_3CH = CH_2 + O_3 \xrightarrow{Zn/H_2O}$
 - (iii) $CH_2 = CH_2 + H_2O + [O]$ Dil.KMnO₄
- 9. (i) What are the conditions for a compound/species to be aromatic [3] according to Huckel's rule?
 - (ii) How will you convert Benzene to acetophenone?
- 10. (i) An alkene 'A' of molecular mass '28u' on treatment with [5] bromine gives a product 'B'. The Compound 'B' on further dehalogenation with zinc gives back 'A'. Give the structures of 'A' and 'B' and also the sequence of reactions.
 - (ii) Why is benzene extraordinarily stable although it contains three double bonds?
 - (iii) How can we convert ethyne into benzene?

UNIT TEST-II

Time Allowed: 1 Hr. Maximum Marks: 20 General Instructions: (i) All questions are compulsory. (ii) Maximum marks carried by each question are indicated against it. 1. The methyl group in Benzene ling is [1] Ortho directing Ortho and meta directing (a) (b) Ortho and para directing (c) Para directing (d) Action of acetylene on dilute H₂SO₄ gives : 2. [1] Acetic acid Acetone (a) (b) Acetaldehyde (d) Ethyl alcohol (c) What is the other name for Geometrical isomerism? 3. [1] In the following questions a statement of Assertion (A) followed by Reason (R) is given. Use the following key to select correct answer: (a) Both Assertion and Reason are correct but Reason is the correct explanation of Assertion. (b) Both Assertion and Reason are correct but Reason is not the correct explanation of Assertion. (c) Both Assertion and Reason are incorrect. (d) Assertion is not correct but Reason is correct. **Assertion:** Benzene does not decolourise bromine water. 4. **Reason:** Benzene is stabilised by resonance due to delocalisation of π

5. **Assertion:** Acidity of C-H bond decreases in the order:

$$HC \equiv CH > H_2C = CH_2 > H_3C - CH_3$$

Reason: Greater the percentage s-character, more is the acidity of C—H bond.

Reason: Greater the percentage s-character, more is tHow will you convert methyl bromide to ethane?

[2]

7. Explain with the help of equation ozonolysis of propene.

[2]

8. Write the mechanism of nitration of benzene.

[3]

9. How will you convert benzene into

[3]

- (i) Acetophenone
- (ii) Toluene

electrons.

10. Write short note on the following:

[5]

- (i) Wurtz reaction.
- (ii) Kolbe's electrolysis
- (iii) Ozonolysis
