**21** 

# **Hydrocarbons**

## • Classification of Hydrocarbons

Simple organic compounds having only carbon and hydrogen are called hydrocarbons. Their is classification as follows:

Hydro carbon	Type	Charac-	General	Bond	Example
		teristic	formula		
Alkane	acyclic	saturated	$C_nH_{2n+2}$	C – C	$\mathrm{CH_3} - \mathrm{CH_2} - \mathrm{CH_3}$ propane
	cyclic	saturated	$C_nH_{2n}$	C - C $C - C$	CH <sub>2</sub> cyclo propane
					$H_2C$ $CH_2$
alkene	acyclic	unsaturated	$C_nH_{2n}$	C = C	$CH_3 - CH = CH_2$ propene
	cyclic	unsaturated	$C_nH_{2n-2}$	C = C	CH <sub>2</sub> cyclo propene
					нс
alkyne	acyclic	unsaturated	$C_nH_{2n-2}$	$C \equiv C$	$CH_3 - C \equiv CH \text{ propyne}$
arene	cyclic	specific	$C_nH_{2n-6m}$	alternate	
		unsaturation		C - C and	Benzene
				C = C	

## • Classification of carbon atoms in hydrocarbon

- (i) Carbon connected with one other carbon is called primary (1°) carbon.
- (ii) Carbon connected with two other carbon is called secondary (2°) carbon.
- (iii) Carbon connected with three other carbon is called tertiary (3°) carbon.
- (iv) Carbon connected with four other carbon is called quaternary (4°) carbon.
- 1. Which of the following is a saturated hydrocarbon?
  - (A) Propene
- (B) Benzene
- (C) Cyclohexane
- (D) Acetylene

- 2. Which of the following is isomer of cyclo alkene?
  - (A) cyclo alkane
- (B) alkyne
- (C) alkene
- (D) arene

- **3.** What is the general formula of cyclic alkene?
  - (A)  $C_{n}H_{2n+2}$
- (B)  $C_{n}H_{2n-2}$
- (C)  $C_n H_{2n}$
- (D)  $C_n H_{2n+1}$
- **4.** Which of the following substance contains tertiary carbon?
  - (A) Propane
- (B) n butane
- (C) 2 Methyl propane(D) Methane

- 5. Which of the following compound is only acyclic?
  - (A) alkane
- (B) alkene
- (C) alkyne
- (D) arene

- **6.** Which hydro carbon has 4° carbon?
  - (A) isobutane
- (B) n-hexane
- (C) neopentane
- (D) n-butane
- 7. In which of the following hydro carbon compound all four types of carbon (1°, 2°, 3°, 4°) are present ?
  - (A) 2, 3-dimethyl butane

(B) 2, 2 dimethyl butane

(C) neo pentane

- (D) 2, 2, 3 trimethyl pentane
- **8.** How many tertiary carbon atoms are present in 2, 3 dimethyl butane?
  - (A) 1
- (B) 2

- (C) 3
- (D) 4
- 9. Number of 1°, 2°, and 3° hydrogen atoms present in neopentane are ..... respectively.
  - (A) 0, 12, 6
- (B) 12, 2, 0
- (C) 6, 2, 0
- (D) 12, 1, 1

- 10. ..... is example of alkyne.
  - (A)  $C_5H_8$
- (B)  $C_8 H_{10}$
- $(C) C_{9}H_{18}$
- (D)  $C_7H_1$

Answers : 1. (C), 2. (B), 3. (B), 4. (C), 5. (C), 6. (C), 7. (D), 8. (B), 9. (B), 10. (A)

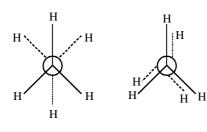
- General information, Nomenclature and isomerism in alkane
  - General Information:
    - $\rightarrow$  sp<sup>3</sup> hybridization and tetrahedral shape.
    - → C C and C H bond length 154 and 112 pm.respectively
    - $\rightarrow$  H C H bond angle 109° 28'
    - → Nomenclature: Refer Unit: 20
    - $\rightarrow$  Isomerism :

In alkane chain isomerism and conformational isomerism is observed. Out of which chain isomerism is discussed in unit 20.

Conformational isomerism: In alkane C - C single bond can rotate along with its bond axis. As a result hydrogen or atoms connected with both carbon can arranged at different distance and various spatial arrangement become possible. These are known as conformer isomers or rotamers. Out of these two isomers staggerd and eclipsed are important other isomers are in between two out of which skew or gauche is also important onc.

# Two conformer of ethane are as follows:

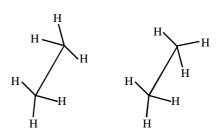
Newmann projection



Staggered

**Eclipsed** 

Sawhorse projection



Staggered

**Eclipsed** 

- In staggered structure hydrogen of both carbon are arranged far from each other hence repulsion is less so it is more stable. (dihedral angle 180°)
- In eclipsed structure hydrogen of both carbon are arranged closer (in diff plane) from each other hence, repulsion is more so stability is less (dihedral angle 0°)
- Out of staggered and eclipsed, staggered is more stable.
- Stability order for conformers of butane is as follows: Staggered (anti) > Skew or Gauche > Partially eclipsed > Fully eclipsed
- In cyclohexane chair and boat conformers are observed out of which chair conformer is more stable.
- In alkane compound C H bond length and, H C H bond angle is ..... respectively. 11.
  - (A) 112 pm 120°
- (B) 112 pm, 109° 281
- (C) 154 pm, 109° 28<sup>1</sup> (D) 135 pm, 180°
- 12. Which of the following is the structure of isobutyl group?
  - (A)  $CH_3 CH_2 CH_2 CH_2 -$
- (B) CH<sub>3</sub> CH CH<sub>2</sub> CH<sub>3</sub>
- (C)  $CH_3 CH CH_2 CH_3$
- (D)  $(CH_3)_3 C -$
- 13. Which sentence is incorrect with reference to isopentane?
  - (A) It has three methyl group.
- (B) It has only one CH<sub>2</sub> group.

(C) It has one - CH group .

- (D) It has one quaternary carbon
- 14. Which of the following compound has isopropyl group?
  - (A) 3, 3-dimethyl pentane

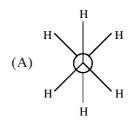
(B) 2, 2, 3, 3-tetramethyl pentane

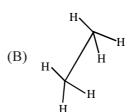
(C) 2-methyl pentane

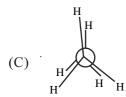
- (D) 2, 2, 3-tri methyl pentane
- Acidic saturated hydrocarbon has molor mass 72 gm/mol then how many isomers are possible for 15.
  - (A) 2
- (B) 4
- (C) 5

(D) 3

- C H bond length is least in ..... 16.
  - (A)  $C_2H_2$
- (B)  $C_2H_4$
- $(C) C_2H_6$
- (D)  $C_2H_2Br_2$
- 17. Which of the following is staggered conformer of ethane?







(D) None of the three.

- 18. Out of four conformer of n-butane which one is most stable?
  - (A) eclipsed
- (B) partially eclipsed
- (C) staggered
- (D) gauch

- 19. Which structure of cyclo hexa triene is most stable?
  - (A) chair
- (B) boat
- (C) half chair
- (D) plannar
- **20.** In which of the following cyclic structure maximum strain is observed?
  - (A) Cyclohexane
- (B) Cyclo pentane
- (C) Cyclo butane
- (D) Cyclo propane

Answers: 11. (B), 12. (B), 13. (D), 14. (C), 15. (D), 16. (C), 17. (A), 18. (C), 19. (D), 20. (D)

# • Preparation and properties of alkane

## • Preparation:

By hydrogenation of alkene and alkyne

$$R - CH = CH_2 + H_2 \xrightarrow{Pt/Pd} R - CH_2 - CH_3$$

$$R - C \equiv CH + 2H_2 \xrightarrow{Pt/Pd} R - CH_2 - CH_3 \qquad (R = alkyl group)$$

• Reduction of alkyl halide

$$R - X + H_2 \xrightarrow{Zn/HCl} R - H + HX$$

• By decarboxylation of sodium salt of carboxylic acid

$$R - COONa \xrightarrow{\text{NaOH} + CaO} R - H + Na_2CO_3$$

• Wurtz reaction

$$2R - X + 2Na \xrightarrow{\text{ether}} R - R + 2 NaX$$

• Grignard reaction

$$R - X + Mg \xrightarrow{\text{other}} R - Mg - X \xrightarrow{HX} R - H + Mg X_2$$

Frankland reaction

$$2R - I + Zn \xrightarrow{\text{ether}} R - R + ZnI_2$$

- By reduction of carbocyclic acid, aldehyde ketone and alcohol alkane is formed.
- Correy-house reaction

$$R - X + R_2^{'}CuLi \longrightarrow R - R' + RCu + LiX$$
 (R and R' same or different)

## • Kolbe's electrolytic method

## • Physical properties

- ullet Alkane having  $C_1$  to  $C_4$  are gaseous,  $C_5$  to  $C_{17}$  are liquid  $C_{18}$  or more than  $C_{18}$  are solids.
- In alkane series or number of carbon increases boiling point increases.
- As alkane is nonpolar insoluble in water but in non polar solvent like benzene it is soluble.
- In alkane boiling point decreases from  $1^{\circ} \rightarrow 2^{\circ} \rightarrow 3^{\circ}$ .

## • Chemical properties

• Halogenation: Reactivity order  $F_2 > Cl_2 > Br_2 > I_2$ .

$$CH_4 + Cl_2 \xrightarrow{hv} CH_3 - Cl + HCl$$

• Reaction with steam : 
$$CH_{4(g)} + H_2O_{(g)} \xrightarrow{Ni} CO_{(g)} + H_{2(g)}$$
 water gas

#### • Arromatization :

$$CH_3 - (CH_2)_4 - CH_3 \xrightarrow{Mo_2O_3 \text{ or } V_2O_5 \text{ or } Cr_2O_3} + 4H_2$$

n-hexane benzene

#### • Isomerization:

$$\begin{array}{c} \text{CH}_3 - \text{CH}_2 - \text{CH}_2 - \text{CH}_3 & \xrightarrow{\text{[AlCl}_3]} & \text{CH}_3 - \text{CH}_4 - \text{CH}_3 & \text{....} & \text{isobutane} \\ & & & \text{n-butane} & & & \text{CH}_3 \end{array}$$

## • Racking:

• Nitration : 
$$CH_4 + HNO_3 \longrightarrow CH_3NO_2 + H_2O$$

Methane Nitro methane

• Sulphonation: 
$$C_6H_{14} + H_2SO_4 \xrightarrow{675 \text{ K}} C_6H_{13} - SO_3H + H_2O$$
  
hexane hexane sulphonic acid

• Oxidation: 
$$CH_{4(g)} + 2O_{2(g)} \xrightarrow{\Delta} CO_{2(g)} + 2H_2O_{(g)} + heat$$

21.	Conversion of but-1-en	ne into but-2-ene is done	by 1	reacting it with		
	(A) Pd / $H_2$	(B) Zn / HCl	(C)	Sn / HCl	(D) Zn – Hg	
22.	By which method prop	oane can be obtained for	m pr	ropene ?		
	(A) Wurtz reaction		(B)	Grignard reaction		
	(C) Catalytic hydrogen	ation	(D)	Frank land reaction	1	
23.	In preparation of propa	ane from propyne hybrid	izatio	on of carbon change	es from	
	(A) sp <sup>2</sup> to sp	(B) $sp^2$ to $sp^3$	(C)	sp to sp <sup>3</sup>	(D) sp <sup>3</sup> to sp	
24.	By which reaction but	ane can be prepared from	n eth	nyl chloride in one	step?	
	(A) Wurtz reaction	(B) Frankland reaction	(C)	Hydrogenation	(D) Grignard reaction	
25.	Preparation of methano	e can be done by				
	(A) hydrogenation	(B) water reaction	(C)	decarboxcylation	(D) all the given	
26.	By which method alke	ne having same no of ca	arbor	n can be prepared fi	rom halo alkane?	
	(A) Wurtz reaction	(B) Grignard reaction	(C)	reduction	(D) both (B) and (C)	
27.	Kolbe's electrolysis of	sodium acetate gives				
	(A) methane	(B) ethane	(C)	ethene	(D) acetic acid	
28.	has highest boiling	g point.				
	(A) n-hexane		(B)	n-pentane		
	(C) 2-methyl butane		(D)	2, 2-dimethyl propa	nne	
29.	Boiling point of neope	ntane is more than				
	(A) iso pentane	(B) n-pentane	(C)	butane	(D) n-hexane	
30.	Benzene obtained from	n n-hexane by react	ion.			
	(A) isomerism	(B) Craking	(C)	rearrangement	(D) cydization	
31.	Kerosene is a mixture	of				
	(A) alkanes	(B) aromatic compound	ds	(C) alcohols	(D) CO + $H_2$	
32.	Which compound give	s methane by reaction w	ith v	vater?		
	(A) $Al_4C_3$	(B) CaC <sub>2</sub>	(C)	VC	(D) SiC	
33.	For following conversi	on CH <sub>3</sub> CH <sub>2</sub> COOH —	→ CI	H <sub>3</sub> CH <sub>2</sub> CH <sub>3</sub> which re	eagnet is appropriate?	
	(A) LiAlH <sub>4</sub>	(B) Sodalime	(C)	Red P / HI	(D) Zn / HCl	
34.	Halogenation of alkane	e is example of which of	the	following ?		
	(A) Electrophilic substi	tuion	(B)	Nucleophilic substit	cution	
	(C) Free radical substit	tution	(D)	addition reaction		
35.	Which parafin is solid	at room temperature ?				
	(A) $C_3H_8$	(B) $C_8H_{18}$	(C)	$C_4H_{10}$	(D) C <sub>20</sub> H <sub>42</sub>	
Ansv		(C), <b>23.</b> (C), <b>24.</b> (A), A			B), <b>28.</b> (A), <b>29.</b> (C),	
	<b>30.</b> (D), <b>31.</b> (A), <b>32.</b> (A), <b>33.</b> (C), <b>34.</b> (C), <b>35.</b> (D)					

## • General information, Nomenclature and isomerism of alkene

- sp² hybridization and trigonal plannar shape
- C C and C H bond length 134 and 110 pm respectively
- H C H bond angle  $116^{\circ}6'$  and C C H bond angle  $121^{\circ}7'$
- Nomenclature: Refer Unit: 20
- **Isomerism**: Alkene has position isomerism, chain isomerism and geometrical isomerism position isomerism is discussed in unit 20.
- Geometrical isomerism: In some alkene compounds though their molecular and structural formula are same but atoms or groups arranged differently in space in different direction. Which is called geometrical isomerism. Geometrical isomers shown by cis and trans.
- This isomerism is observed in planner molecule having C = C.
- Due to restricted rotation around C = C geometrical isomerism arises.
- Two isomers of but-2-ene are as follows:

$$CH_3$$
  $C = C$   $CH_3$   $CH_3$   $C = C$   $CH_3$   $C = C$   $CH_3$ 

cis but-2-ene

trans but-2-ene

- Geometrical isomerism is observed in compounds like ab C = C ab, ab C = C ad ab C = C de but it is not observed in the compounds like ab C = C bb or aa C = Cab type of compounds.
- cis isomer is polar while trans isomer is non-polar.
- cis isomer has greater solubilly in water than trans isomer.
- **36.** In ethene C = C and C H bond length is ..... respectively.
  - (A) 134, 112
- (B) 154, 112
- (C) 134, 110
- (D) 139, 110

- 37. Shape of ethene molecule is ......
  - (A) linear
- (B) tetrahedral
- (C) octahedral
- (D) trigonal plannar
- 38. IUPAC name of  $(CH_3)_2$ . CH CH = CH CH = CH is ......

$$CH - CH_3$$
 $C_2H_5$ 

- (A) 2, 7-dimethyl nona 3, 5-diene
- (B) 2, 7-dimethyl octa-3, 5-diene
- (C) 2-ethyl 7-methyl octa-3, 5-diene
- (D) 7-methyl non-3-ene

39.	IUPAC	name	of /	\ <u></u>	is	

- (A) 4-ethyl 3-methyl pent-4-en-1-yne
- (B) 2-ethyl 4-methyl pent-1-en- 4-yne
- (C) 4-ethyl 3-Methyl pent-1-yne-4-ene
- (D) 2-ethyl 3-methyl pent-4-yne-1-ene
- 40. Which compound shows geometrical isomerism?
  - (A) butane
- (B) but-1-ene
- (C) but–2–ene
- (D) but-2-yne

- 41. Number of possible alkene isomers of C<sub>4</sub>H<sub>8</sub> is ......
  - (A) 2
- (B) 3
- (C) 4

- (D) 5
- 42. In which of the following compound carbon has sp and sp<sup>2</sup> hybridization.
  - (A) propene

(B) propyne

(C) propadiene

- (D) none
- 43. Hybridization of carbon atoms buta 1, 3-diene is ..... type.
  - (A) sp,  $sp^2$ ,  $sp^3$
- (B)  $sp^2$ ,  $sp^3$
- (C) sp, sp<sup>2</sup>
- (D) Võík sp<sup>2</sup>

- 44. How many alkene isomers of C<sub>5</sub>H<sub>10</sub> are possible ?
  - (A) 7
- (B) 5

- (C) 4
- (D) 6

- Number of isomers of  $C_2H_2Br_2$  is ..... . **45.**

(A) 1 (B) 2 (C) 3 (D) 0 **Answers : 36.** (C), **37.** (D), **38.** (A), **39.** (B), **40.** (C), **41.** (C), **42.** (C), **43.** (D), **44.** (D), **45.** (C)

- Preparation and properties of alkene
  - (I) Preparation:
  - Hydrogenation of alkyne:  $R C \equiv CH \xrightarrow{H_2} R CH = CH_2$

Lindler's catalyst

• Dehydrohalogenation of alkylholide ( $\beta$  – elimination reaction) :

$$^{\beta}\text{CH}_3 - {^{\alpha}\text{CH}_2} - \text{Cl} + \text{KOH} \xrightarrow{\text{Ethanol}} \text{CH}_2 = \text{CH}_2 + \text{KCl} + \text{H}_2\text{O}$$

• From vicinal dihalide:

$$Br - CH_2 - CH_2 - Br + Zn \xrightarrow{\Delta} CH_2 = CH_2 + Zn Br_2$$

• Dehydration of alcohol:

$$CH_3 CH_2 OH \xrightarrow{conc H_2SO_4} CH_2 = CH_2 + H_2O$$

• Kolbe's electrolysis method :

## (II) Physical properties:

- Boiling point of alkyne is more than corresponding alkane due to polar nature.
- Boilling point of cis isomer is more than trans isomer.
- Melting point of trans alkene is more than cis isomer.
- In soluble in water but soluble in organic solvent.
- In isomers boiling point decreases from  $1^{\circ} \rightarrow 2^{\circ} \rightarrow 3^{\circ}$ .

## (III) Chemical Properties:

Due to  $\pi$ -bond alkene shows following additional reactions :

• Hydrogenation : 
$$CH_2 = CH_2 + H_2 \xrightarrow{Pt/Pd} CH_3 - CH_3$$

• Halogenation : 
$$CH_2 = CH_2 + X_2 \longrightarrow X - CH_2 - CH_2 - X$$
 where,  $X = CI$ , Br, I

• Hydrohalogenation:

(a) Symmetric alkene 
$$CH_2 = CH_2 + HX \longrightarrow CH_3 - CH_2 - X$$
 (X = Cl, Br, I)

(b) Asymmetric alkene : 
$$CH_3 - CH = CH_2 + HX \longrightarrow CH_3 - CH - CH_3$$
 2-halo propane

**Note:** Hydrohalogenation of asymmetric alkene follows Markovnikoff rule: As per this rule negative part of reagent. Combine with ethylenic carbon having less number of hydrogen.

(c) Reaction of asymmetric alkene with HBr in presence of peroxide catalyst follows anti Markovnikoff rule. In this reaction negative part of reagent  $(\overline{X})$  combines with ethylenic carbon having more number of hydrogen.

eg., 
$$CH_3 - CH = CH_2 + HBr \xrightarrow{benzoyl peroxide} CH_3 - CH_2 - CH_2 - Br$$

$$1 - Bromo propane$$

• This reaction do not take place with HCl or HI.

• Hydration : 
$$CH_2 = CH_2 + H_2O \xrightarrow{\text{dil } H_2SO_4} CH_3 - CH_2 - OH$$

$$CH_3 - CH = CH_2 + H_2O \xrightarrow{\text{dil } H_2SO_4} CH_3 - CH_3 - CH_3 \text{ (Markovnikoff)}$$

• Oxidation : (a) 
$$CH_2 = CH_2$$
  $\xrightarrow{Cold}$   $CH_2 - CH_2$   $CH_2 - CH_2$   $CH_3 - CH_4$   $CH_4 - CH_5$   $CH_5 - CH_6$   $CH_6 - CH_6$ 

ethylene glycol

During this reaction pink colour of KMnO<sub>4</sub> disappear it is known as Bayer's test for unsaturation.

(b) 
$$CH_2 = CH_2 \xrightarrow{\text{Hot}} 2 \text{ HCOOH}$$

Formic acid

• Ozonolysis:

$$R - CH = CH_2 + O_3 \longrightarrow R - CH \longrightarrow CH_2 \xrightarrow{C} R - CHO + HCHO$$
Ozonide

- Which major product is obtained on reaction of chloro ethene with alcoholic KOH? 46.
  - (A) ehtane
- (C) ethanol
- (D) ethyne

47. 
$$CH_3 - CH_2 - CH_2 - CH_3 \xrightarrow{alcoholic} major product.$$

- (A) butane
- (C) but-2-ene
- (D) butyne

- 48. Lindlar's catalyst is mixture of what?
  - (A)  $Ni + H_2$
- (B) Pt / H<sub>2</sub>
- (C) Pd + Pt
- (D) Pd + charlcol
- 49. Which substance can form alkene by elimination reaction?
  - (A) haloalkane
- (B) dihaloalkane
- (C) alcohol
- (D) all
- Which compound on heating with Zn gives but-2-ene? **50.** 
  - (A) 2, 3-dibromobutane

(B) 1, 2-dibromobutane

(C) but-2-yne

- (D) None
- 51. ..... is the gas which removes colour of basic KMnO<sub>4</sub> produced by reaction between ethyl iodide and alcoholic KOH.
  - (A) C<sub>2</sub>H<sub>6</sub>
- (B)  $C_2H_4$
- $(C) C_2H_2$
- (D) CH<sub>4</sub>
- 52. Preparation of alkene from vicinal dihalide known as ......
  - (A) decarboxylation
- (B) dehydrohelogenation(C) dehalogenation
- (D) dehydrogenation

53. Product of 
$$CH_3 - CH_3 - CH_3 \xrightarrow{H_2SO_4} is.$$

- (B) but -1 ene
- (C) propanol
- (D) 2-methyl propene

**54.** 
$$CH_3 - CH_2 - CH = CH_2 + HBr \longrightarrow \dots$$

$$(A) CH3 - CH2 - CH2 - CH3$$

(B) 
$$CH_3 - CH_2 - CH_2 - CH_2 - Br$$

(C) 
$$CH_3 - CH_2 - CH - CH_3$$
  
Br
$$(D) CH_3 - CH_2 - C \equiv CH$$

(D) 
$$CH_3 - CH_2 - C \equiv CH_3$$

55.	$CH_{a} =$	CH -	C1 + HC1	$\longrightarrow$	
			0		

- (A)  $CH_3 CH_2 CI$  (B)  $CH_3 CHCl_2$  (C)  $CH_2 CH_2$  (D)  $CH \equiv CH$  CI CI
- ..... product is obtained on reaction between 2-methyl prop-1-ene and HBr in presence of benzoyl **56.** peroxide.
  - (A) CH<sub>3</sub> CH CH<sub>2</sub>Br CH<sub>3</sub>,

(B)  $CH_3 - C - CH_3$ 

(C) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>Br

- (D)  $CH_3 CH_2 CH CH_3$
- 57. For the reaction of which of the following compounds Markonikoff rule is useful?
  - (A)  $C_2H_4 + HC1$

(B)  $C_3H_6 + Br_2$ 

(C)  $C_3H_6 + HBr$ 

(D)  $C_2H_4 + I_2$ 

**58.** 
$$R - CH_2 - CH = CH_2 + IC1 \longrightarrow$$

- (A)  $R CH_2 CH CH_2 I$  (B)  $R CH_2 CH CH_2 CI$  (C)  $R CH_2 CH = CH_2$  (D)  $R CH = CH CH_2 I$

- 59. In presence of peroxide alkene do not give anti Markovnikoff reaction with HCl or HI because .....
  - (A) both are highly ionic.
  - (B) one is oxidising agent other is reducing agent.
  - (C) In both case one step is endothermic.
  - (D) In both case all the steps are endothermic.
- **60.** Major product of reaction between 3-phenyl propene with HBr is ......
  - (A)  $C_6H_5CH_2CH(Br)CH_3$

(B) C<sub>6</sub>H<sub>5</sub>CH(Br)CH<sub>2</sub>CH<sub>3</sub>

(C) C<sub>6</sub>H<sub>5</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>Br

- (D)  $C_6H_5CH(Br)CH = CH_7$
- Which of the following compound do not give reaction with HBr according to Markovnikoff **61.** rule.
  - (A) Propene
- (B) but–1–ene
- (C) but-2-ene
- (D) pent-2-ene

- isobutene + HBr  $(C_6H_5CO)_2O_2 \longrightarrow \cdots$ . 62.
  - (A) 3° butyl bromide

(B) isobutyl bromide

(C) 3° butyl alcohol

- (D) isobutyl alcohol
- **63.** Butene from butane can be formed by reaction with .....
  - (A) Zn HC1
- (B) Sn HC1
- (C) Zn Hg
- (D) Pd / H<sub>2</sub>

	(A) Reaction takes	place by more stable car	bonium ion.		
	(B) Reaction takes	place by more stable fre	e radical.		
	(C) Reaction takes	place by more stable car	bonium ion.		
	(D) none of the abo	ove.			
65.	Identify B in the fo	llowing reaction sequence	e :		
	$CH_3 - CH = CH -$	$CH_3 \xrightarrow{O_3} A \xrightarrow{Zn} B$			
	(A) 2CH <sub>3</sub> CHO	(B) 2CH <sub>3</sub> COCH <sub>3</sub>	(C) 2CH <sub>3</sub> COOH	(D) $CH_3CH_2CH_2CH_3$	
Ans	Answers: 46. (B), 47. (C), 48. (D), 49. (D), 50. (A), 51. (B), 52. (C), 53. (D), 54. (C), 55. (B), 56. (A), 57. (C), 58. (A), 59. (C), 60. (B), 61. (C), 62. (B), 63. (D), 64. (A), 65. (A)				
• Ge	eneral Information, N	Nomenclature and isome	erism of alkyne		
	• General information	n : sp hybridization and l	inear shape.		
	• $C \equiv C$ and $C - H$	bond length 120 and 106	pm respectively.		
	• Bond angle 180°				
	• Nomenclature: Re	fer Unit: 20			
	• Isomerism: In alky	vne position, chain and ri	ing chain isomerism is	observed.	
66.	Compound with whi	ich of the following bour	nd is most reactive ?		
	(A) C – C	(B) $C = C$	(C) $C \equiv C$	(D) All	
67.	In which of the foll	owing compound all aton	ns are linearly bonded	?	
	(A) Propane	(B) Propene	(C) Propyne	(D) All	
68.	Hybridization of sec	cond carbon in buta 1, 2-	-diene is		
	(A) sp	(B) $sp^2$	(C) $sp^3$	(D) dsp <sup>2</sup>	
69.	Triple bond between	n two carbon in alkyne i	s formed by		
	(A) 1 sigma 2 pi	(B) 1 pi 2 sigma	(C) 3 sigma	(D) 3 pi	
70.	IUPAC name of (C	$H_3)_3 C - C \equiv C - C (C$	CH <sub>3</sub> ) <sub>3</sub> is		
	(A) 3, 3, 4, 4-tetra	methyl Hex-3-yne	(B) 2, 2, 5, 5-tetra	n methyl Hex-3-yne	
	(C) 2, 2, 5, 5-tetra	methyl Hex-4-yne	(D) di(trimethyl)–2	-yne	
71.	How many alkyne i	somers are possible for c	compound having mole	cular formula C <sub>5</sub> H <sub>8</sub> ?	
	(A) 2	(B) 4	(C) 3	(D) 5	
72.	Which of the follow	ving is isomer of propyne	?		
	(A) Cyclo propyne	(B) Cyclo propene	(C) Propene	(D) Propane	
Ans	swers : 66. (C), 67	(. (C), <b>68.</b> (A), <b>69.</b> (A),	<b>70.</b> (B), <b>71.</b> (C), <b>72.</b>	(B)	
		50:	1 ———		

Propene on reaction with HI gives isopropyl iodide not n-propyl iodide because ......

64.

## • Preparation and properties of alkyne

#### • Preparation:

Hydrolysis of Calcium Carbide:

$$CaC_2 + 2H_2O \longrightarrow Ca(OH)_2 + CH \equiv CH$$

Dehydrohalogenation of vicinal dihalide:

Dehalogenation of tetrahalide:

$$R - C Br_2 - CH Br_2 + 2Zn \xrightarrow{\Delta} R - C \equiv CH + 2ZnBr_2$$

Kolbe's electrolysis method:

Higher alkyne from acetylene:

$$CH \equiv CH \xrightarrow{\text{Na/Liq. NH}_3} HC \equiv C^- - \text{Na}^+ \xrightarrow{R - X} R - C \equiv CH + \text{NaX}$$

#### • Physical properties

- Alkyne is weakly polar.
- Insoluble in water but soluble in non polar solvent.
- Compared to corresponding alkane and alkene, alkynes have higher boiling point and melting point.
- Ethyne and terminal alkyne shows acidic character. They are weak acids.

## • Chemical properties

Like alkene alkyne also gives addition reactions. :

• Hydrogenation:

• CH = CH + H<sub>2</sub> 
$$\xrightarrow{Pd/C}$$
 CH<sub>2</sub> = CH<sub>2</sub>  $\xrightarrow{H_2}$  CH<sub>3</sub> - CH<sub>3</sub>

Lindlar's catalyst

H<sub>2</sub>, Pd / C

R

C = C

H

cis - alkene

R

C = C

H

rtrans-alkene

## Halogenation:

$$CH \equiv CH + Cl_2 \rightarrow CHCl = CHCl \xrightarrow{Cl_2} CHCl_2 - CHCl_2$$

## **Hydrohalogenation:**

$$CH \equiv CH + HBr \rightarrow CH_2 = CHBr \xrightarrow{HBr} CH_3 - CHBr_2$$
 (Markovnikoff)

## **Hydration:**

#### **Polymerization:**

$$3CH \equiv CH \xrightarrow{\text{Red hot Fe tube}}$$

#### Oxidation:

(a) 
$$CH_3 - C \equiv CH \xrightarrow{\text{Cold}} CH_3COOH + CO_2$$
  
terminal alkyne

non terminal alkyne

(b) 
$$CH_3 - C \equiv CH \xrightarrow{Hot} CH_3COOH + CO_2$$

terminal alkyne

$$CH_3 - C \equiv C - CH_3 \xrightarrow{\text{Hot}} CH_3COOH + CH_3COOH$$

non-terminal alkyne

#### Ozonolysis:

$$CH \equiv CH + O_3 \longrightarrow \begin{matrix} CH \longrightarrow CH \\ \hline \\ O \longrightarrow O \end{matrix} \qquad \begin{matrix} CH \longrightarrow CH \\ \hline \\ O \longrightarrow O \end{matrix} \qquad \begin{matrix} CH \longrightarrow CH \\ \hline \\ O \longrightarrow O \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \\ O \longrightarrow O \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \\ O \longrightarrow O \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \\ O \longrightarrow O \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \\ O \longrightarrow O \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \\ O \longrightarrow O \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \\ O \longrightarrow O \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \\ O \longrightarrow O \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \\ O \longrightarrow O \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \\ O \longrightarrow O 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H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \end{matrix} \qquad \begin{matrix} CD / H_2O \\ \hline \end{matrix} \qquad \end{matrix} \qquad \end{matrix}$$

#### Acidic nature:

(i) Acetylene and terminal alkynes reacts with strong base like sodamide due to acidic nature.

$$CH \equiv CH + NaNH_2 \longrightarrow HC \equiv C^- \cdot Na^+ + NH_3$$

Sodium acetylide

(ii) By decomposing Grignard reagent gives alkane

$$CH \equiv CH + R - Mg - X \longrightarrow R - H + HC \equiv C - MgX$$

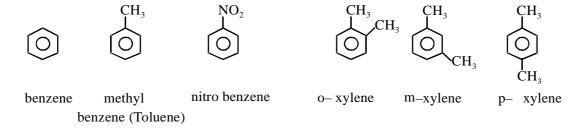
73.	Colourless gas obtain	ned by reaction between	metal carbide and wat	er is
	(A) Methane	(B) ethane	(C) acetylene	(D) ethylene
74.	Identify X and Y in t	the following reaction:		
	$CaC_{2(S)} + H_2O_{(1)}$ -	$\rightarrow X \xrightarrow{\text{dil } H_2SO_4} \rightarrow$	Y	
	(A) X-ethylene Y-ac	etaldehyde	(B) X-acetylene Y-	ethanol
	(C) X-ethane Y-etha	anol	(D) X-acetylene Y-	ethenol
75.	1, 2-di bromo ehtane	e on reaction with alcoh	olic KOH gives	
	(A) ethane	(B) acetylene	(C) ethylene	(D) methane
76.	Product obtain by re-	action between haloalke	ne and sodamide is	
	(A) alkane	(B) alkene	(C) alkyne	(D) holo alkane
77.	Which of the followi	ng compound has acidic	hydrogen ?	
	(A) ethene	(B) ethyne	(C) propyne	(D) both (B) and (C)
78.	By which of the follo	owing reaction propyne	is formed ?	
	(A) CH <sub>3</sub> Br with acet	tylene	(B) CH <sub>3</sub> Br with sod	ium acetylide
	(C) CH <sub>3</sub> I with sodius	m acetate	(D) Methane with e	thene
<b>79.</b>	-	ne by hot alkaline KMn	O <sub>4</sub> gives	
	(A) CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> CO		(B) CH <sub>3</sub> CH <sub>2</sub> COOH	
	(C) CH <sub>3</sub> CH <sub>2</sub> COOH -		(D) CH <sub>3</sub> CH <sub>2</sub> COOH	+ НСООН
80.		eaction between acetyler		
	(A) Butane	(B) Ethane	(C) but-1-ene	(D) but-2-ene
81.	Final product of redu	active ozonolysis of but-	-2-yne is	
	(A) glyoxal	(B) 2CH <sub>3</sub> COOH	(C) but 2, 3 – dione	(D) butanal
82.	Final product of C <sub>2</sub> H	I <sub>2</sub> with HCl is		
	(A) $CH = CH - CI$	(B) $CH_3 - CHCl_2$	II	(D) None
	**	r. C1	CH – Cl	
83.	$CH \equiv CH + HCI - \frac{H}{C}$	$\xrightarrow{\text{IgCl}_2}$ product.		
	(A) Methyl chloride	(B) Dichloro ethane		(D) Ethyl chloride
84.		is obtained by reactio	n of propyne with dilu	ite H <sub>2</sub> SO <sub>4</sub> in presence of
	HgSO <sub>4</sub> .		(D) many l hydrogon	aulmhata
	<ul><li>(A) propanal</li><li>(C) propanal</li></ul>		<ul><li>(B) propyl hydrogen</li><li>(D) acetone</li></ul>	Surpriate
85.		s are given by alkyne co		
	(A) substitution	(B) addition	(C) polymerization	(D) all
Ans		(C), <b>75.</b> (B), <b>76.</b> (C), (C), <b>84.</b> (D), <b>85.</b> (D)	, 77. (D), 78. (B), 79.	. (C), <b>80.</b> (B), <b>81.</b> (C),

## • Arene (aromatic hydro carbon) General information, Nomencluture and structure of benzene

- Compounds having benzene ring are called aromatic compounds.
- Aromatic compounds having only carbon and hydrogen are called aromatic hydrocarbon or arene. Their first member is benzene.
- General formula of arene is  $C_n H_{2n-6m}$  where m = number of rings
- Aromatic compounds having benzene ring are called benzanoids. eg. biphenyl, napthalene, anthracene, phenanthrene, napthacene
- Aromatic compounds which do not have benezene ring are called non-benzaoids. eg. pyrol, furan, thiofin, pyridine.

#### • Nomencluture

• By displacing hydrogen of benzene with other group substituted benzene can be obtained.



## • Structure of Benzene

- Molecular formula of benzene is  $C_6H_6$ . It has hexagonal cyclic structure with alternate C C and C = C.
- Resonating structure of benzene as per kekule is as follows:

$$\bigcirc \longleftrightarrow \bigcirc \equiv \bigcirc$$

- Mono substituted benzene has only one isomer while disubstituted has three isomers (or the meta and para)
- Benzene has less chemical reactivity and greater stability is called aromatic character.
- Benzene follows  $(4n + 2) \pi e^{-}$  rule of Huckel hence it is aromatic.
- (A) Napthalene (B) Aniline (C) Pyridine (D) Napthacene **89.** Reaction of benzene with 3 moles Cl<sub>2</sub> in presence of sunlight gives ...... product.
- (A)  $C_6H_3Cl_3$  (B)  $C_6Cl_6$  (C)  $C_6H_6Cl_6$  (D)  $C_6H_5Cl$  **90.** Chemical name of pesticide gamaxene is ......
- (A) DDT (B) BHC (C) chloral (D) hexa chlorethane

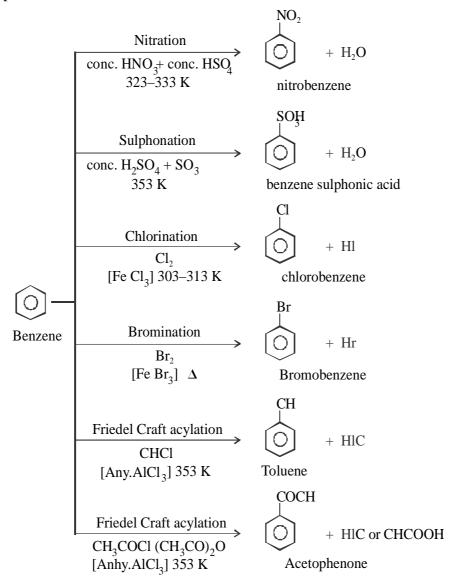
91.	Carbon-carbon bond le	ingth in benzene is	piii.		
	(A) 154 and 134	(B) only 134	(C) only 154	(D) 139	
92.	Which product is obta	ined by ozonolysis of be	enzene followed by hydo	orlysis of product ?	
	(A) Benzenetriozonide	(B) acetaldehyde	(C) glyoxal	(D) benzone	
93.	According to Huckel's	s rule how many $\pi e^-s$ a	re present in phenanthre	ene ?	
	(A) 6	(B) 10	(C) 12	(D) 14	
94.	Benzene shows ty	ype of reaction.			
	(A) substitution	(B) addition	(C) oxidation	(D) all the given	
95.	How many $\sigma$ and $\pi$ be	onds are present in bipl	nenyl respectively?		
	(A) 22 and 6	(B) 23 and 6	(C) 13 and 5	(D) 12 and 6	
96.	Which of the followin	g sentence is correct for	or benzene ?		
	(A) Due to unsaturation benzen gives addition reaction easily.				
	(B) In benzen 3 types of C-H bond.				
	(C) In benzene cyclic	delocalized $\pi$ bonds an	re present.		
	(D) In benzene $6\pi$ ele	ectrons are localized.			
97.	Number of $\sigma$ and $\pi$ b	onds in phenol are	respectively.		
	(A) 13, 2	(B) 12, 3	(C) 13, 3	(D) 13, 4	
98.	What is the hybridizat	ion of all six carbon in	benzene ?		
	(A) sp	(B) $sp^2$	(C) $sp^3$	(D) sp <sup>2</sup> and sp <sup>3</sup>	
99.	Why benzene has high	er stability and less rea	ectivity?		
	(A) sp³ hybridization of	of six carbon atom	(B) cyclic structure of	f six carbon atom	
	(C) high resonance en	nergy	(D) unsaturated natur	e of benzene ring	
100.	Which compound do r	ot follow Huckel's rule	?		
	(A) Benzene	(B) Cyclohexane	(C) Napthalene	(D) Pyrol	
,		(D), <b>88.</b> (C), <b>89.</b> (C), <b>97.</b> (C), <b>98.</b> (B), <b>99.</b> (	90. (B), 91. (D), 92. (C), 100. (B)	(C), <b>93.</b> (D), <b>94.</b> (D),	
• Pre	eparation of Benzene				
	<ul> <li>By cyclic polymerizat</li> </ul>	•			
	$3CH \equiv CH - \frac{\text{red hot}}{87}$	$\frac{\text{Fe tube}}{3 \text{ K}} \longrightarrow $			
•	By decarboxylation or	f sodium benzoate			
	COONa L sodalii	me			
	NaOH +	$\longrightarrow$ $  \cap   + N_2 $	2CO <sub>3</sub>		
	• By reduction of phenomenature $\Delta$	ol			

$$\begin{array}{c|cccc}
OH & & & \\
\hline
O & & \Delta
\end{array}$$

$$\begin{array}{c}
Zn & & \\
\hline
\end{array}$$

$$\begin{array}{c}
+ & ZnO
\end{array}$$

## • Electrophillic substitution reaction of benzene



## • Directive effect of inductive group

- Due to effect of first group connected with benzene, at which position second group is connected is determined. Hence 1st group present in mono substituted benzene is called inductive group and this effect is called directive effect.
  - Directive effect are at two types:
    - (a) o and p directive group (e donor group)

(b) meta directive group (e attracting group)

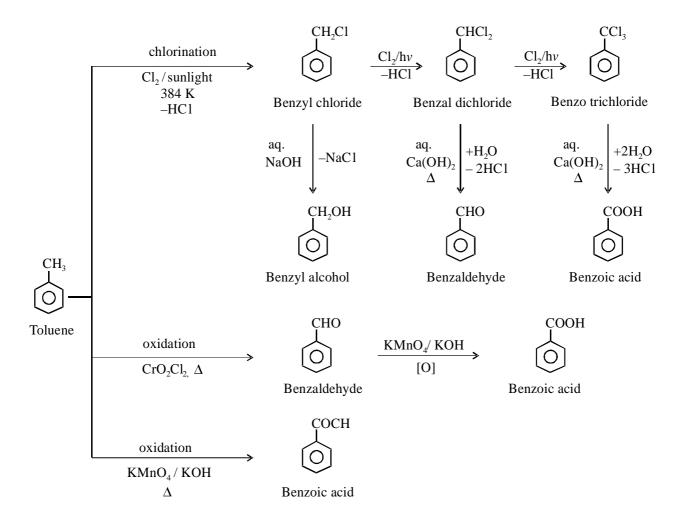
$$-\mathrm{NO}_2,\ -\mathrm{SO}_3\mathrm{H},\ -\mathrm{COOH},\ -\mathrm{COOR},\ -\mathrm{CN},\ -\mathrm{CHO},\ -\mathrm{COR},\ -\mathrm{CCl}_3$$

	(A) +CH <sub>3</sub>	(B) CH <sub>3</sub> <sup>-</sup>	(C) +SO <sub>3</sub> H	(D) CH <sub>3</sub> <sup>+</sup> CO
106.		ý.	(Temperature 323 K	3
100.	•	intration of benzene is .		)
	(A) nitro benzene		(B) nitroso benzene	
	(C) O-dinitro benzene		(D) m-dinitrobenzene	
	Product obtained by	reaction of benzene w	rith concentrated HNO <sub>3</sub>	and H <sub>2</sub> SO <sub>4</sub> of 353 to
107.	363 K temperature is		3	2 4
107.			(B) nitro benzene	2 4
107.	363 K temperature is	c acid		- ,
<ul><li>107.</li><li>108.</li></ul>	363 K temperature is (A) benzene sulphonic (C) m-dinitro benzen	c acid	<ul><li>(B) nitro benzene</li><li>(D) (o and p) dinitro</li></ul>	- ,
	363 K temperature is (A) benzene sulphonic (C) m-dinitro benzen Which of the following	c acid	<ul><li>(B) nitro benzene</li><li>(D) (o and p) dinitro</li></ul>	benzene
	363 K temperature is  (A) benzene sulphonic  (C) m-dinitro benzen  Which of the followir reaction ?  (A) benzene		(B) nitro benzene (D) (o and p) dinitro factive towards electroph	benzene ilic aromatic substitution
108.	363 K temperature is  (A) benzene sulphonic  (C) m-dinitro benzen  Which of the followir reaction ?  (A) benzene	c acid e ng compound is most re (B) toluene	(B) nitro benzene (D) (o and p) dinitro factive towards electroph	benzene ilic aromatic substitution
108.	363 K temperature is  (A) benzene sulphonic  (C) m-dinitro benzene  Which of the following reaction?  (A) benzene  In which of the follow  (A) toluene	c acid e ng compound is most re  (B) toluene ving compound second s	(B) nitro benzene (D) (o and p) dinitro factive towards electroph (C) benzoic acid figure acid figure (C) phenol	benzene tilic aromatic substitution (D) nitro benzene

# Reaction of phenyl group in toluene:

- Toluene also gives electrophillic substitution reaction like benzene for eg. nitration, sulphonation, chlorination, bromination, alkylation, acylation etc.
- - CH<sub>3</sub> group in toluene is ortho-para directive group hence second substitution takes place at ortho and para position. (Study the reactions from Textbook.)

# • Reaction of methyl group in toluene :



**Conversion :** Process to convert one organic compound into another is called organic conversion. eg, Benzene to benzoic acid.

$$\begin{array}{c|c}
\hline
& CH_3Cl \\
\hline
& [Anhy.AlCl_3]
\end{array}$$

$$\begin{array}{c}
CH_3 \\
\hline
& \Delta [O]
\end{array}$$

$$\begin{array}{c}
COOH \\
\hline
& \Delta [O]
\end{array}$$
Benzoic acid

- 111. Product obtained by complete nitration of toluene is ......
  - (A) o-nitro toluene

(B) p-nitro toluene

(C) 2, 4-dinitro toluene

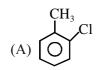
- (D) 2, 4, 6-trinitro toluene
- 112. ..... is obtained by reaction of toluene with Cl<sub>2</sub> in presence of FeCl<sub>3</sub>.
  - (A) m-chloro benzene

(B) benzyl chloride

(C) o and p chloro toluene

(D) chloro benzene

113. ..... is obtained by chlorination of toluene in presence of sunlight.



(B) CH<sub>2</sub>C

(C) CH<sub>3</sub>



- 114. In Friedel Craft reaction ..... catalyst is used.
  - (A) anhy.ZnCl<sub>2</sub>
- (B) anhy.AlCl<sub>2</sub>
- (C) FeCl<sub>2</sub>
- (D) CH<sub>3</sub>Cl
- 115. What is the function of anhy AlCl<sub>3</sub> in Friedel Craft reaction?
  - (A) To absorb water.

- (B) To produce free radical.
- (C) To produce electrophilic ion.
- (D) To produce nucleophilic ion.
- 116. Methylation of toluene in presence of HF/BF<sub>3</sub> gives ......
  - (A) o-xylene
- (B) p-xylene
- (C) o and p xylene
- (D) m-xylene
- 117.  $\overbrace{\bigcirc}^{\text{CH}_3} \xrightarrow{\text{KMnO}_4/\text{KOH}} X \xrightarrow{\text{sodalime}} Y, X \text{ and } Y \text{ are } \dots ...$

(A) 
$$X = \bigcirc$$

$$Y = \bigcirc$$

(B) 
$$X = \bigcirc$$

$$Y = \bigcirc$$
SO<sub>3</sub>H

(C) 
$$X = \bigcirc$$

(D) 
$$X = \bigcirc$$

$$Y = \bigcirc$$

- 118.  $X \xrightarrow{Zn} Benzene \xrightarrow{CH_3COCl} Y identify X and Y.$ 
  - (A) X-benzoic acid, Y-toluene
- (B) X-phenol, Y-toluene
- (C) X-phenol, Y-aceto phenone
- (D) X-toluene, Y-aceto phenone
- 119. In the reaction  $\bigcirc$   $\xrightarrow[\text{anhy AlCl}_3]{C_2H_5Cl}$  P  $\xrightarrow[903 \text{ K}]{}$  Q final product Q is .......
  - (A) Benzene
- (B) m-Xylene
- (C) ethyl benzene
- (D) styrene

**120.** Identify A and B in the following reaction :

$$CaC_{2(s)} + H_2O_{(1)} \longrightarrow A \xrightarrow{H_2SO_4} B$$

- (A) A-ethylene B-acetaldehyde
- (B) A-acetylene B-propanol

(C) A-ethane B-ethanol

(D) A-acetylene B-acetaldehyde

Instru		: For questi		correct sentence T and	d for incorrect sence F then
121.	(i)	Each 'carbon	in benzene is sp² hybr	ridized.	
	(ii)	In benzene C	–C bondlength is 154	and 134 para alternativel	y.
	(iii)	In benzene 63	π electrons are loalized	l.	
	(iv)	In benzene 6	carbon and 6 hydroge	n are identical.	
	(A)	TFFT	(B) FTFT	(C) TTFF	(D) TTTF
122.	(i)	At normal ter	mperature benzene give	es addition reaction with	$H_2$ , $Cl_2$ and $O_3$ .
	(ii)	Benzene give	s nucleophilic substituti	ion reaction easily.	
	(iii)	Reactivity of	benzene is more than	alkene.	
	(A)	TFT	(B) TTT	(C) TTF	(D) FFF
123.	(i)	Benzene is p	lannar.		
	(ii)	In benzene cy	yelic shaped molecular	orbital of $6\pi$ electron.	
	(iii)	Due to reson	ance reactivity of benz	zene increases.	
	(iv)	Benzene is in	soluble in water.		
	(A)	TTFT	(B) TTFF	(C) FTTF	(D) TTFF
124.	(i)	-NHCH <sub>3</sub> is n	neta directing group.		
	(ii)	Chlorination	of benzoic acid gives C	-chloro benzoic acid.	
	(iii)	TNT is used	as explosive.		
	(iv)	Ozonolysis of	f benzene gives addition	n reaction.	
	(A)	TFFT	(B) TTFT	(C) FFTT	(D) FFTF
125.	(i)	Chlorobenzen	e gives Friedel Craft r	eaction.	

(ii) -OH group is electron donor group.

(iii) Nitration of benzene done by nucleophile <sup>+</sup>NO<sub>2</sub>.

(iv) Benzene burns with sooty flame.

(B) TFFF (A) TTFT

Instruction: Read the paragraph carefully and answer the questions below it.

Paragraph: The compound or ions which follows Huckel's rule are called aromatic compounds. Main important point of this law are as follows:

(C) FTTF

(D) TFFT

(i) Compound or ion must be plannar.

- (ii) It has delocalized  $\pi$  electron coloured.
- (iii)  $\pi$  electron cloud should cover all the carbon atom of cyclic system.
- (iv) Total number of  $\pi$  e<sup>-</sup> should be 4n + 2 where n = 0, 1, 2, ....

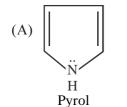
126. Which of the following compound is not aromatic?

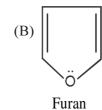


127. Which of the following system is aromatic?



**128.** Which of the following compound is aromatic?







(D) all

Thiofene

• In question 129 to 131 column-I and column-II aer given match them and select correct option :

129.	Column-I	Column-II
	(1) alkane	$(p) C_n H_{2n-2}$
	(2) alkene	$(q) C_n H_{2n}$
	(3) alkyne	$(r) C_n H_{2n+2}$
	(A) arene	(c) C H

(A) 
$$(1)$$
– $(r)$ ,  $(2)$ – $(q)$ ,  $(3)$ – $(p)$ ,  $(4)$ – $(s)$ 

(B) 
$$(1)$$
– $(p)$ ,  $(2)$ – $(q)$ ,  $(3)$ – $(r)$ ,  $(4)$ – $(s)$ 

(C) 
$$(1)$$
– $(s)$ ,  $(2)$ – $(r)$ ,  $(3)$ – $(q)$ ,  $(4)$ – $(r)$ 

(D) 
$$(1)$$
– $(r)$ ,  $(2)$ – $(s)$ ,  $(3)$ – $(p)$ ,  $(4)$ – $(q)$ 

130.	Column-I (Reaction)	Column-II (Hydro carbon product)
	(a) decarboxylation of sodium acetate	(p) ethyne
	(b) Wurtz reaction	(q) 2-methyl propane
	(c) Correy-House reaction	(r) n-butane
	(d) dehydro halogenation	(s) methane

(A) 
$$(a)-(r)$$
,  $(b)-(p)$ ,  $(c)-(q)$ ,  $(d)-(s)$ 

(C) 
$$(a)-(q)$$
,  $(b)-(s)$ ,  $(c)-(r)$ ,  $(d)-(p)$ 

131.	Column-I (Reaction)	Column-II (Reagent)
	(a) $CH_3COCH_3 \rightarrow CH_3CH_2CH_3$	(p) Sodalime
	(a) $CH_3COCH_3 \rightarrow CH_3CH_2CH_3$ (b) $CH_3CH_2Br \rightarrow CH_2 = CH_2$	(q) Zn powder
	(c) $C_6H_5COOH \rightarrow C_6H_6$	(r) alcoholic KOH
	$(d) C_6 H_5 OH \rightarrow C_6 H_6$	(s) HI / Red P

(A) 
$$(a)-(p)$$
,  $(b)-(s)$ ,  $(c)-(q)$ ,  $(d)-(r)$ 

- In the following questions two sentences are given out of which one is assertion (A) and other is reason (R). Study the sentences given below properly and select the open given below:
  - (A) Assertion (A) and reason (R) both are correct and reason (R) is correct explanation of assertion (A).
  - (B) Assertion (A) and reason (R) both are correct but reason (R) is not correct explanation of assertion (A).
  - (C) Assertion (A) is correct while reason (R) is incorrect.
  - (D) Assertion (A) is incorrect while reason (R) is correct.

132. Assertion (A): Stability of benzene is less than alkene

**Reason** (R): Resonance energy of benzene indicates it's greater stability and less chemical reactivity.

133. Assertion (A) : cis but-2-ene is polar while trans but-2-ene is non polar.

**Reason** (R): In trans isomer both methyl groups are on opposite sides hence net dipole moment is zero.

134. Assertion (A): Though benzene has double bond it does not undergo polymerization.

**Reason** (R): In normal condition benzene does not behave as alkane.

135. Assertion (A): Compared to n-pentane boiling point of neo pentane is more.

**Reason** (R): neo-pentane has quaternary carbon.

**136.** Assertion (A): Dehydration of butan-2-ol mainly gives but-2-ene.

**Reason** (R): Dehydration takes place mainly through carbocation intermediate.

137. Assertion (A): Benzene easily gives electrophilic substitution reaction.

**Reason** (R): Benzene is unsaturated hydro carbon.

**138. Assertion (A)** : By addition reaction of propene with HCl in presence of peroxide mainly gives 2-chloro propane.

**Reason** (R): This reaction takes place by free radical intermediate.

139. Assertion (A): Cyclopenta dienyl anion is more stable than allyl anion.

**Reason** (R): Cyclopenta dienyl anion is aromatic.

140. Assertion (A): Reaction of acetylene with sodamide gives sodium acetylide and ammonia.

**Reason** (R): sp hybridized carbon of atoms acetylene are highly electro negative.

Answers: 111. (D), 112. (C), 113. (B), 114. (B), 115. (C), 116. (D), 117. (C), 118. (C),

119. (D), 120. (D), 121. (A), 122. (D), 123. (A), 124. (C), 125. (A), 126. (B),

127. (B), 128. (D), 129. (A), 130. (B), 131. (C), 132. (D), 133. (D), 134. (C),

135. (B), 136. (D), 137. (B), 138. (D), 139. (A), 140. (A).