

Alkane

Which represents an alkane 1.

[CPMT 1976]

- (a) $C_5 H_8$
- (b) $C_8 H_6$
- (c) C_9H_{10}
- (d) $C_7 H_{16}$

The decreasing order of boiling points is[BHU 1999] 2.

- (a) *n*-Pentane > iso-Pentane > neo-Pentane
- (b) iso-Pentane > n-Pentane > neo-Pentane
- (c) neo-Pentane > iso-Pentane > n-Pentane
- (d) *n*-Pentane > *n*eo-Pentane > iso-Pentane

To prepare a pure sample of n-hexane using 3. sodium metal as one reactant, the other reactant [BHU 1999] will be

- (a) *n*-propyl bromide
- (b) Ethyl bromide and *n*-butyl bromide
- (c) Ethyl chloride and n-butyl chloride
- (d) Methyl bromide and n -pentyl chloride

In the preparation of Grignard reagent from 4. haloalkane, the metal [RPET 1999]

- (a) Mq
- (b) Zn

- (c) Li
- (d) K

5. Sodium acetate can be converted to ethane by

[Pune CET 1998]

- (a) Heating with LiAlH 4
- (b) Electrolysing its aqueous solution
- (c) Heating with sodalime
- (d) Heating with calcium acetate

6. Which of the following compounds is used in antiknock compositions to prevent the deposition of oxides of lead on spark plug, combustion chamber and exhaust pipe

[KCET 1998]

- (a) Glycerol
- (b) Glycol
- (c) 1, 2-dibromoethane (d) Benzene

Which of petroleum corresponds to kerosene oil 7.

[DCE 1999]

- (a) $C_{15} C_{18}$
- (b) $C_{10} C_{12}$
- (c) $C_5 C_9$
- (d) $C_1 C_9$

In the reaction $CH_3 - Br + 2Na + Br - CH_3 \rightarrow$, the 8.

[Pb. CET 1999; CPMT 1983. 86; KCET 1992; MP PMT 1994; BHU 1998; MP PMT 2002; MP PET 1986]

- (a) Wurtz reaction
- (b) Aldol condensation
- (c) Perkin's reaction
- (d) Levit reaction

Iodoethane reacts with sodium in the presence of 9. dry ether. The product is [AFMC 1997; KCET 1998]

- (a) Pentane
- (b) Propane
- (c) Butene
- (d) Butane

Which of the following is oxidised by KMnO₄

- (a) Methane
- (b) Pentane
- (c) Isobutane
- (d) Neopentane

Which of the following has maximum stability[AIIMS 2001

- (a) $\stackrel{+}{CH}_3$ (b) $CH_3 \stackrel{|}{C} H$ CH_3 (c) $CH_3 \stackrel{|}{C} CH_3$ (d) $CH_3 \stackrel{|}{C} CH_3$

- [DPMT 2000]

12. The most volatile compound is (a) 2, 2-dimethyl propane (b) 2-methyl butane

- (c) Isobutane (d) *n*-pentane
- In Wurtz reaction, the reagent used is [EAMCET 1998] 13.
 - (a) Na
- (b) Na/liquid NH₃
- (c) Na/dry ether
- (d) Na/dry alcohol

Which of the following has highest octane number [MP PMT 2000]

- (a) *n*-hexane
- (b) *n*-heptane
- (c) *n*-pentane pentane
- (d) 2, 2, 4-trimethyl

What is freon-12

16.

[RPET 1999]

- (a) Pesticide
- (b) Refrigerant (d) Lubricant
- (c) Solvent
- The petrol having octane number 80 has[MP PET 2000]
- (a) 20% normal heptane + 80% iso-octane
- (b) 80% normal heptane + 20% iso-octane
- (c) 20% normal heptane + 80% normal octane
- (d) 80% normal heptane + 20% normal octane

Which of the following reactions will not give 17. propane

[DPMT 2005]

- (a) $CH_3CH_2CH_2Cl \frac{Mg/\text{ether}}{H_2O}$
- (b) $CH_3COCl \frac{CH_3MgX}{H_2O}$
- (c) $CH_3CH = CH_2 \xrightarrow{B_2H_6} CH_3COOH$
- (d) $CH_3CH CH_3 \xrightarrow{P/HI}$

The shape of methane molecule is[MP PET 1997, 2001] (b) Trigonal planar

- (a) Linear
- (c) Square planar Which of the following shows only one
- (d) Tetrahedral

brominated compound (a) Butene-2

- [CPMT 1996] (b) 2, 2-dimethylpropane
- (c) Butyne-1
- (d) Butanol-3

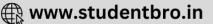
20. Kerosene is used as fuel because it is [CPMT 1996] (a) Less volatile

- (b) More volatile
- (c) Cheap
- (d) Abundantly available

 $CH_3 - CH_2 - CH_2 - CH_3 \xrightarrow{AlCl_3} \text{Product in}$ above reaction is

[RPMT 2003]





- (a) $CH_3 CH CH_2 CH_3$ Rr
- (b) $CH_3 CH CH_3$ CH_3
- (c) $CH_2 CH_2 CH_2$ Rr CH_3
- (d) All of these
- Which of the following statements is not true for 22. ethane

[AIIMS 1996]

- (a) It can be chlorinated with chlorine
- (b) It can be catalytically hydrogenated
- (c) When oxidised produces CO_2 and H_2O
- (d) It is a homologue of iso-butane
- Petroleum refining is [AIIMS 1996; KCET 2004] 23.
- (a) Distillation of petroleum to get different fractions
 - (b) Obtaining aromatic compounds from aliphatic compounds present in petroleum
- (c) Cracking of petroleum to get gaseous hydrocarbons
 - (d) Purification of petroleum
- The chemical added to leaded petrol to prevent the deposition of lead in the combustion chamber

[Kerala (Med.) 2003]

- (a) Iso-octane
- (b) Ethylene dibromide
- (c) Tetraethyl lead
- (d) Mercaptan
- (e) n-Heptane
- In the commercial gasolines, the type of 25. hydrocarbons which are more desirable is[CBSE PMT 1997; AFMC 1997]
 - (a) Branched hydrocarbon
 - (b) Straight-chain hydrocarbon
 - (c) Linear unsaturated hydrocarbon
 - (d) Toluene
- Which of the following is not formed by the 26. reaction of Cl_2 on CH_4 in sunlight [AIIMS 1987]
 - (a) CHCl₃
- (b) CH_3Cl
- (c) CH_3CH_3
- (d) $CH_3CH_2CH_3$
- Which of the following has the highest boiling point

[DPMT 1986]

- (a) Neopentane
- (b) *n*-butane
- (c) *n*-heptane
- (d) Isobutane
- Which gives CH₄ when treated with water 28.

[CPMT 1974, 79; NCERT 1976; IIT-JEE 1990]

- (a) Silicon carbide
- (b) Calcium carbide
- (c) Aluminium carbide
- (d) Iron carbide
- Which of the following does not react with PCl₅

[CPMT 1973]

- (a) CH_3OH
- (b) CH₃COOH

- (c) CH₃CHO
- (d) C_2H_6
- Which of the following compounds is insoluble 30. even in hot concentrated H_2SO_4 [IIT-JEE 1983]
 - (a) Ethylene
- (b) Benzene
- (c) Hexane
- (d) Aniline
- A reaction between methyl magnesium bromide and ethyl alcohol gives[CPMT 1979; MNR 1986; UPSEAT 1999
 - (a) Methane
- (b) Ethane
- (c) Propane
- (d) Butane
- Methane and ethane both can be obtained in 32. single step from

[CPMT 1974; MP PET 1995; AFMC 1998, 2000; BHU 2005]

- (a) CH_3I
- (b) C_2H_5I
- (c) CH_3OH
- (d) C_2H_5OH
- 33. Paraffin wax is
- [MP PMT 1986; CPMT 1993]
- (a) Ester
- (b) Alcohol
- (c) Unsaturated hydrocarbon
- (d) Saturated hydrocarbon
- The number of possible enantiomeric pairs that can be produced during monochlorination of 2methylbutane is

[IIT-JEE 1997]

(a) 2

(b) 3

(c) 4

- (d) 1
- Petroleum consists mainly of [CPMT 1985, 94; KCET 1991]
 - (a) Aliphatic hydrocarbons (b)Aromatic hydrocarbons
 - (c) Aliphatic alcohols (d) None of these
- Petroleum ether can be used as
 - (a) Solvent for fat, oil, varnish and rubber

- (c) Both (a) and (b)
- (d) None of these
- Which of the following are produced from coaltar

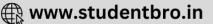
[MNR 1987; UPSEAT 2002]

- (a) Synthetic dyes
- (b) Drugs
- (c) Perfumes
- (d) All the three
- 38. In alkanes, the bond angle is [MP PMT 1989; BHU 1996]
 - (a) 109.5°
- **(b)** 109 °
- (c) 120°
- (d) 180°
- In the preparation of alkanes; a concentrated aqueous solution of sodium or potassium salts of saturated carboxylic acid are subjected to [CPMT 1985; MP
 - (a) Hydrolysis
- (b) Oxidation
- (c) Hydrogenation
- (d) Electrolysis
- Halogenation of alkanes is an example of 40.

[MP PET 1993; KCET 1998]

- (a) Electrophilic substitution
- (b) Nucleophilic substitution
- (c) Free-radical substitution
- (d) Oxidation
- Propionic acid is subjected to reduction with hydroiodic acid in the presence of a little P, the product formed is





In catalytic reduction of hydrocarbons which [JIPMER 1997] 53. (a) Ethane (b) Propane catalyst is mostly used [DCE 2001] (c) Butane (d) None of these (a) Pt /Ni (b) Pd When ethyl iodide and propyl iodide react with (d) Misch Metal (c) SiO_2 *Na* in the presence of ether, they form [BHU 1997] 54. Ethylene reacts with bromine to form[MP PET 2001] (a) One alkane (b) Two alkanes (a) $Br - CH_2 - CH_3$ (b) $CH_3 - CBr_3$ (c) Four alkanes (d) Three alkanes (c) $Br - CH_2 - CH_2Br$ (d) CHBr₂ The alkane that yields two isomeric monobromo derivatives is 55. [EAMCET 1997] obtained from crude petroleum oil by (b) Ethane (a) Neopentane (c) Methane (d) Propane [MP PMT 1999] **44.** Kerosene is a mixture of [CPMT 1979; AFMC 1992] (a) Fractional distillation (b) Vacuum distillation (b) Aromatic compounds (a) Alkanes (c) Steam distillation (d) Pyrolysis (c) Alcohols (d) Aliphatic acids 56. Which of the following does not give alkane 45. When petroleum is heated the vapours contain [MP PMT 1999] mainly (a) Reaction of CH_3I with Na in ether [CPMT 1981] (a) Kerosene (b) Petroleum ether (b) Reaction of sodium acetate with sodalime (c) Diesel (d) Machine oil (c) Electrolysis of concentrated sodium acetate solution **46.** Iso-octane is mixed to the petrol [NCERT 1972] (d) Reaction of ethyl chloride with alco. KOH (a) To precipitate inorganic substances **57.** *LPG* is a mixture of [MP PMT 1999; KCET 2005] (b) To prevent freezing of petrol (c) To increase boiling point of petrol (a) $C_6H_{12} + C_6H_6$ (b) $C_4H_{10} + C_3H_8$ (d) As an antiknock (c) $C_2H_4 + C_2H_2$ (d) $C_2H_4 + CH_4$ Tetraethyl lead is used as[NCERT 1976, 79; DPMT 1984; 58. Carbon black, which is used in making printer's CPMT 1989, 91; BHU 1995] ink, is obtained by decomposition of [MP PET 1993] (a) Fire extinguisher (b) Pain reliever (a) Acetylene (b) Benzene (c) Petroleum additive (d) Mosquito repellent (c) Carbon tetrachloride (d) Methane 48. Cyclohexane, a hydrocarbon floats on water The addition of tetraethyl lead to petrol[MP PET 1993] because (a) Lowers its octane number [NCERT 1976] (b) Raises its octane number (a) It is immiscible with water (c) May raise or lower the octane number (b) Its density is low as compared to water (d) Has no effect on octane number (c) It is non-polar substance 60. Which of the following compound has maximum (d) It is immiscible and lighter than water boiling point [IIT-JEE 1982; MP PMT 1986; 49. Natural gas contains mainly MADT Bihar 1995; Pb. PMT 1999] [MNR 1990; UPSEAT 1999, 2000, 01, 02; BCECE 2005] (a) *n*-hexane (b) *n*-pentane (a) Methane (b) *n*-butane (c) 2, 2-dimethyl propane (d) 2-methyl butane (d) Mixture of octane (c) *n*-octane Knocking sound occurs in engine when fuel[CPMT 1981] **50.** Which compound is not inflammable [MP PET 2001] (a) Ignites slowly (a) CCl_{4} (b) C_2H_5OH (b) Ignites rapidly (c) CH₄ (d) $C_6 H_6$ (c) Contains water Propane is obtained from propene, by which of (d) Is mixed with machine oil 51. the following methods [CBSE PMT 2001; AFMC 2001] Petroleum is mainly a mixture of (b) Dehydrogenation (a) Wurtz reaction [CPMT 1984; Pb. PMT 1999] (c) Frankland reaction (d) Catalytic (a) Alkanes (b) Cyclohexane hydrogenation (c) Benzenoid hydrocarbons (d) Alkenes The organic compound used as antiknock agent in Which of the following has maximum boiling petroleum is [DCE 1999; CPMT 2000; Pb. CET 2000; point MP PET 2001] [IIT-JEE 1986; MP PMT 1986; CPMT 1989] (a) $(C_2H_5)_4 Pb$ (b) *TNT* (a) iso-octane (b) *n*-octane (c) CH_3MgBr (d) $(C_2H_5)_2Hg$ (c) 2, 2, 3, 3-tetramethyl butane

- (d) *n*-butane
- **64.** Aqueous solution of the following compound on electrolysis gives ethane

[NCERT 1983; MP PET 1985; CPMT 1975, 79]

- (a) Acetic acid
- (b) Acetamide
- (c) Potassium acetate
- (d) Ethyl acetate
- **65.** Which of the following does not decolourise bromine solution in carbon disulphide[MP PET 1986]
 - (a) Acetylene
- (b) Propene
- (c) Ethane
- (d) Propyne
- **66.** Anhydrous sodium acetate on heating with sodalime gives

[CPMT 1972, 84; Pb. CET 2001, 2003]

- (a) Acetic acid
- (b) Methane
- (c) Calcium acetate
- (d) Ethane
- **67.** Water gas is
- [CPMT 1993, 2004; Pb. PMT 2004]
- (a) $CO + CO_2$
- (b) $CO + N_2$
- (c) $CO + H_2$
- (d) $CO + N_2 + H_2$
- **68.** A sample of gasoline contains 81% *iso*-octane and 19% *n*-heptane. Its octane number will be[MP PMT 1995]
 - (a) 19
- (b) 81
- (c) 100
- (d) 62
- **69.** The natural petroleum contains [MP PMT 1995]
 - (a) Saturated hydrocarbons
 - (b) Cyclic saturated hydrocarbons
 - (c) Compounds of sulphur
 - (d) All of these
- 70. The preparation of ethane by electrolysis of aqueous solution of potassium acetate is called as [MP PNI PSS]
 - (a) Wurtz reaction
 - (b) Sabatier-Senderen's reaction
 - (c) Kolbe's synthesis
 - (d) Grignard reaction
- **71.** Action of hydrogen chloride on $CH_3 C = CH_2$ CH_3

and on $CH \equiv CH$ will predominantly give the compounds, respectively

(a)
$$CH_3 - CH = CH_2Cl$$
 and $CH_2Cl - CH_2Cl$

$$CH_3$$

(b)
$$CH_3 - CCl = CH_3$$
 and $CH_3 - CHCl_2$

$$CH_3$$

(c)
$$CH_3 - CH = CH_2Cl$$
 and $CH_3 - CHCl_2$
 CH_3

(d)
$$CH_3 - CH = CH_3$$
 and $CH_2Cl - CH_2Cl$
 CH_3

- **72.** As the number of carbon atoms in a chain increases the boiling point of alkanes [AFMC 1989]
 - (a) Increases
 - (b) Decreases
 - (c) Remains same

- (d) May increase or decrease
- 73. In the fractional distillation of crude petroleum
 [Roorkee 1989]
 - (a) Petrol condenses at the bottom of the column
 - (b) The gases condense at the top of the column
 - (c) High boiling constituents condense at the bottom of the column
 - (d) High boiling constituents condense at the top of the column
- **74.** Which of the following is not an endothermic reaction

[J & K 2005]

- (a) Dehydrogenation
- (b) Ethane to ethene
- (c) Combustion of propane
- (d) Change of chlorine molecule into chlorine atoms.
- **75.** Gasoline is the name of **[Roorkee 1989]**
 - (a) Crude oil
 - (b) The gaseous constituents of petroleum
 - (c) The mixture of uncondensed gases produced in the distillation of crude oil
 - (d) The mixture of the residue and gas oil obtained in the distillation of crude oil
- **76.** In the process of cracking [Roorkee 1989]
 - (a) Organic compounds decompose into their constituent elements
 - (b) Hydrocarbons decompose into carbon and
 - (c) High molecular weight organic compounds decompose to give low molecular weight
- organic compounds

 (d) Hydrocarbons yield alkyl radicals and hydrogen
- 77. Octane number has o value for

[Roorkee 1989; MP PET 1999, 2002; MP PMT 2001; KCET 2002]

- (a) iso-octane
- (b) *n*-hexane
- (c) *n*-heptane
- (d) iso-heptane
- **78.** Dry distillation of sodium propanoate with sodalime gives

[CPMT 1996]

- (a) Propane
- (b) Propene
- (c) Ethane
- (d) Ethene
- **79.** What is the chief product obtained when *n*-butane is treated with bromine in the presence of light at 130 ° *C*

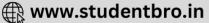
[IIT-JEE 1995]

(a)
$$CH_3 - CH_2 - CH - Br$$
 CH_3

(b)
$$CH_3 - CH - CH_2 - Br$$

$$CH_3$$





(c)
$$CH_3 - C - Br$$
 CH_3

(d)
$$CH_3 - CH_2 - CH_2 - CH_2 - Br$$

- **80.** A mixture of propene and methane is obtained by the cracking of
 - (a) 1-butene
- (b) 2-butene
- (c) *n*-butane
- (d) Isobutane
- Which of the following fractions of petroleum refining contains kerosene? (Boiling ranges in ° *C* are given below)
 - (a) 40 80
- (b) 80 200
- (c) 200 300
- (d) Above 300
- Which of the following statements is incorrect? The members of the homologous series of alkanes

[NCERT 1974]

- (a) Are all straight chain compounds
- (b) Have the general formula $C_n H_{2n+2}$
- (c) Have similar chemical properties
- (d) Show a regular gradation of properties
- 83. On mixing tetraethyl lead to gasoline available at petrol pumps [CPMT 1981]
 - (a) Calorific value of the fuel increases
 - (b) Odour diminishes
 - (c) Less smoke is obtained on combustion
 - (d) Antiknock property of fuel increases
- A liquid hydrocarbon can be converted to gaseous hydrocarbon by [CPMT 1980; MP PMT 2001]
 - (a) Cracking
 - (b) Hydrolysis
 - (c) Oxidation
 - (d) Distillation under reduced pressure
- 85. The tetrahedral nature of carbon was first given by

[MP PMT 1994]

- (a) Kekule
- (b) Le Bell and Van't

Hoff

- (c) Pauling
- (d) Armstrong and Bayer
- **86.** Formation of alkane by the action of Zn on alkyl halide is called [DPMT 1984; MHCET 2004]
 - (a) Frankland's reaction (b) Wurtz reaction
 - (c) Cannizzaro reaction (d) Kolbe's reaction
- Which of the following compounds will form a 87. hydrocarbon on reaction with Grignard reagent[CPMT 9788, §3]mple of 2, 3-dibromo-3-methylpentane is
 - (a) CH_3CH_2OH
- (b) CH₃CHO
- (c) CH₃COCH₃
- (d) $CH_3CO_2CH_2$
- **88.** Name the hydrocarbon that is a liquid at STP
 - (a) Ethane
- (b) Propane
- (c) n-butane
- (d) n-pentane
- 89. Which statement is not true concerning alkanes

[MP PET 2003]

- (a) Large number alkanes are soluble in water
- (b) All alkanes have a lower density than water
- (c) At room temperature some alkanes liquids, some solids and some gases
- (d) All alkanes burn
- 90. Fischer Tropsch process for the is used manufacture of

[DCE 1999; MP PET 2003]

- (a) Synthetic petrol plastics
- (c) Ethanol
- (d) Benzene

(b) Thermosetting

91. Which one of the following compounds cannot be prepared by Wurtz reaction[Kurukshetra CEE 2002;

MP PMT 2002; MP PET 2003]

- (a) CH_4
- (b) C_2H_6
- (c) C_3H_8
- (d) $C_4 H_{10}$
- A fuel contains 25 % n-heptane and 75 % isooctane. Its octane number is[MP PMT 1993; MP PET 1994]
 - (a) 50
- (b) 75
- (c) 100
- (d) 25
- Sodium ethoxide is a specific reagent for [CPMT 1985] 93.
 - (a) Dehydration
 - (b) Dehydrogenation
 - (c) Dehydrohalogenation
 - (d) Dehalogenation
- Which of the following has highest percentage of 94. hydrogen

[CPMT 1975; 79]

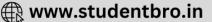
- (a) CH_4
- (b) C_2H_4
- (c) C_6H_6
- (d) C_2H_2
- What is the molecular formula of the alkane, the 5.6 *litre* of which weight 11 q at STP[MP PMT 2003]
 - (a) C_6H_{14}
- (b) $C_4 H_{10}$
- (c) C_3H_8
- (d) C_2H_6
- The reference compound 'iso-octane' which is 96. used in determining the octane number of gasoline has the structure

(a)
$$CH_3 - CH(CH_3) - CH(CH_3) - CH(CH_3) - CH_3$$

- (b) $CH_3 C(CH_3)_2 CH_2 CH(CH_3) CH_3$
- (c) $CH_3 C(CH_3)_2 CH(CH_3) CH_2 CH_3$
- (d) $CH_3 C(CH_3)_2 C(CH_3)_2 CH_3$
- heated with zinc dust. The resulting product is isolated and heated with HI in the presence of phosphorus. Indicate which is the structure that represent the final organic product formed in the reaction [CBSE PMT 1991]
- (a) $CH_3 CH_2 CH CH_2 CH_3$ CH_{2}







$$CH_3$$

- (b) $CH_2 = CH CH CH_2 CH_3$
- (c) $CH_3 CHI CH CH_2 CH_3$ CH_3
- (d) $CH_2 = CH C(I) CH_2 CH_3$
- The order of appearance of the following with rising temperature during the refining of crude oil

[MNR 1993; UPSEAT 2002]

- (a) Kerosene oil, gasoline, diesel
- (b) Diesel, gasoline, kerosene oil
- (c) Gasoline, diesel, kerosene oil
- (d) Gasoline, kerosene oil, diesel
- When sodium propionate is heated with soda lime, the main product is
 - (a) Ethane
- (b) Methane
- (c) Propane
- (d) Butane
- 100. Gasoline is a mixture of alkanes with the number of carbon atoms [CPMT 1983, 84; BVP 2003]
 - (a) $C_3 C_5$
- (b) $C_5 C_6$
- (c) $C_6 C_8$
- (d) $C_7 C_9$
- 101. The final product of complete oxidation of hydrocarbons is

[CPMT 1981]

- (a) Acid
- (b) Aldehyde
- (c) $H_2O + CO_2$
- (d) Dihydric alcohol
- 102. Which of the following will have least hindered rotation about carbon-carbon bond[IIT-JEE 1987; CPMT 1989, the fact that
 - (a) Ethane
- (b) Ethylene
- (c) Acetylene
- (d) Hexachloroethane
- 103. Which of the following represents the most oxidized form of hydrocarbon [MP PMT/PET 1988]
 - (a) *CO*,
- (b) RCHO
- (c) RCOOH
- (d) RCOOOH
- 104. Name the reaction

$$C_{10}H_{22} \xrightarrow{-900\,K} C_4H_8 + C_6H_{14}$$

[MP PET 1995; MP PMT 1997]

- (a) Alkylation
- (b) Cracking
- (c) Pyrolysis
- (d) Fractionation
- 105. How many types of carbon atoms are present in 2, 2, 3-trimethylpentane
 - (a) One
- (b) Two
- (c) Three
- (d) Four
- 106. Which one gives only one monosubstitution product on chlorination [AIEEE 2003]
 - (a) *n*-pentane
- (b) Neopentane
- (c) Isopentane
- (d) n-butane

107. CH_4 is formed when

[AFMC 1987]

- (a) Sodium acetate is heated with sodalime
- (b) Iodomethane is reduced
- (c) Aluminium carbide reacts with water
- (d) All of these
- 108. A mixture of methane, ethylene and acetylene gases is passed through a Wolf's bottle containing ammoniacal cuprous chloride. The gas coming out is [NCERT 1976]
 - (a) Methane
 - (b) Acetylene
 - (c) A mixture of methane and ethylene
 - (d) The original mixture
- 109. At room temperature solid paraffin is[RPET/PMT 1999]
 - (a) C_3H_8
- (b) $C_8 H_{18}$
- (c) $C_4 H_{10}$
- (d) $C_{20}H_{42}$
- 110. Which one of the following compounds does not give addition reactions [MADT Bihar 1981]
 - (a) Aldehydes
- (b) Alkanes
- (c) Alkenes
- (d) Alkynes
- (e) Ketones
- (f) All of these
- 111. The most important method of preparation of hydrocarbons of lower carbon number is[CBSE PMT 1989]
- (a) Pyrolysis of higher carbon number hydrocarbons
 - (b) Electrolysis of salts of fatty acids
 - (c) Sabatier and Senderen's reaction
 - (d) Direct synthesis
- 112. The inorganic origin of petroleum is indicated by
 - (a) Its constituents can be separated by fractional distillation
 - (b) Carbon and hydrocarbon can combine by absorption of solar energy give hydrocarbons
 - (c) Petroleum contains traces of chlorophyll
- (d) Oil fields are located with the help of seismograph
- 113. Which of the following is a gemdihalide[CPMT 1976, 88]
 - (a) CH₃.CHBr.CHBr.CH₃ (b) CH₂Br.CH₂Br
 - (c) CHBr = CHBr
- (d) CH_3CHBr_2
- 114. Which one of the following contain isopropyl group

[BHU 2005]

- (a) 2,2,3,3-tetramethylpentane
- (b) 2-methylpentane
- (c) 2,2,3-trimethylpentane
- (d) 3,3-dimethylpentane
- 115. Natural gas is a mixture of [MP PMT 1986]
 - (a) $CO + CO_2$
- (b) $CO + N_2$







1112 Hydrocarbon (c) $CO + H_2 + CH_4$ (d) $CH_4 + C_2H_6 + C_3H_8$ 116. By Wurtz reaction, a mixture of methyliodide and ethyliodide gives (a) Butane (b) Ethane (c) Propane (d) A mixture of the above three

117. Product obtained by nitration of propane is[RPMT 2003]

(a) Nitropropane

(b) Nitromethane

(c) Nitroethane

(d) All of these

118. Isomerism in saturated hydrocarbons is due to

(a) Change in the valence of carbon

(b) Change in the ratio of elements in compounds

(c) Formation of branches in the chain of C atoms

(d) Formation of double bond

119. Photochemical chlorination of alkane is initiated by a process of [DPMT 1985; NCERT 1978]

(a) Pyrolysis

(b) Substitution

(c) Homolysis

(d) Peroxidation

120. Which of the following is not linked with methane

(a) Marsh gas

(b) Natural gas

(c) Producer gas

(d) Coal gas

121. Which of the following has highest octane number

[MP PET 1996]

[BHU 2003]

(a) n-hexane

(b) *n*-heptane

(c) Iso-octane

(d) *n*-heptane and iso-octane mixed in ratio 50:

50

122. A mixture of ethyl iodide and n-propyl iodide is subjected to Wurtz reaction. The hydrocarbon that will not be formed is

[IIT-JEE (Screening) 1990]

(a) *n*-butane

(b) *n*-propane

(c) *n*-pentane

(d) *n*-hexane

123. Most of the hydrocarbons from petroleum are obtained by

[CPMT 1974, 80]

(a) Fractional distillation (b)Fractional crystallization

(c) Vaporization

(d) Polymerization

124. Which is the best antiknock compound or Which one of the following substances is used as an antiknock compound

[CPMT 1974, 81, 99, 2000; RPMT 2002; CBSE PMT 1996; KCET (Med.) 2000 MP PET 1985, 87, 97, 2001;

MP PMT 1994, 96; AIIMS 2000]

(a) Lead tetrachloride

(b) Lead acetate

(c) Zinc ethyl

(d) Tetraethyl lead (TEL)

125. In the dichlorination reaction of propane, mixture of products are obtained. How many isomers, the mixture contains [Orissa JEE 2003]

(a) 2

(b) 3

(c) 4

(d) 5

126. Which of the following cycloalkane gives open chain compound, when reacts with bromine [Orissa JEE 20

(a) Cyclopropane

(b) Cyclopentane

(c) Cyclohexane

(d) Cyclo-octane

127. Grignard reagent is not prepared in aqueous medium but prepared; in ether medium because the reagent

[KCET 2002]

(a) Reacts with water

(b) Is insoluble in water

(c) Is highly reactive in ether

(d) Becomes inactive in water

128. A sample of petrol is a mixture of 30% *n*-heptane and 70% iso-octane. The sample has octane number

[MP PET 1985]

(a) 30

(b) 70

(c) 15

(d) 35

129. For the reduction of ketones to hydrocarbon, the appropriate agent is [DPMT 2002]

(a) HI

(b) Zn - Hg/HCl

(c) Red phosphorous

(d) H_2SO_4

130. Heating of alkanes with fuming sulphuric acid or oleum at high temperature, which forms sulphonic acid, is called

[MH CET 1999]

(a) Nitration

(b) Halogenation

(c) Sulphonation

(d) Oxidation

131. Propane is obtained from propene by which method?

> [CPMT 1997; CBSE PMT 2001; AFMC 2001; MH CET 2001]

(a) Catalyst hydrogenation (b) Wurtz reaction

(c) Dehydrogenation

(d) Frankland reaction

132. B.P. of branched chain alkanes as compared to straight chain alkanes are[MP PMT 1987; AIIMS 1999]

(a) Lower

(b) Equal

(c) Higher

(d) Independent of the chain

133. Daily use candles (paraffin wax) contain[CPMT 1996]

(a) Higher saturated hydrocarbon

(b) Lower saturated hydrocarbon

(c) Higher unsaturated hydrocarbon

(d) Lower unsaturated hydrocarbon

134. The reaction $CH_4 + Cl_2 \xrightarrow{\text{uv light}} CH_3Cl + HCl$ is an

example of

[CBSE PMT 1999, 2002]

(a) Addition reactions (b) Substitution reaction

(c) Elimination reaction (d) Rearrangement reaction





135.	Normal butane convert	into isobutane by[RPMT 2002]		(a) Wurtz reaction	(b) Kolbe's reac	tion
	(a) LiAlH ₄	(b) AlCl ₃		(c) Reduction of alkyl h	alide (d)Hydroger	nation of alkene
	(c) <i>NaBH</i> ₄	•	147.	Which hydrocarbon wi	ll be most stable	
126	•				[MP PET	2000, 03]
130.	Alcoholic solution of K			(a) Methane	(b) Ethane	
	(a) Dehydration	CPMT 1982, 86; IIT-JEE 1990] (b) Dehydrogenation		(c) Propane	(d) Butane	
	(c) Dehydrohalogenati		148.	C-H bond length is grea		
125	-	reacting with water gives			1989; MNR 1990; A	MU 2002]
137.	Aluminium carolice on	[NCERT 1981; MP PET 1985]		(a) C_2H_2	(b) C_2H_4	
	(a) Methane	(b) Ethane		(c) C_2H_6	(d) $C_2H_2Br_2$	
	(c) Ethene	(d) Ethyne	149.	Which one of the follo	wing compounds	does not
128		on bond distance is found		form an ozonide	[EAM	CET 1997]
150.	in	on bond distance is round		(a) Ethene	(b) Propyne	
	[MP PMT 1987; I	IT-JEE 1981; Bihar MEE 1995]		(c) Propene	(d) Propane	_
	(a) Ethyne	(b) Ethene	150.	Which type of hybridisa		-
	(c) Ethane	(d) Benzene		[CBSE PMT 1991; Bi	(b) sp^2	MER 1997]
139.	Which of the following	ng reaction is expected to		-	•	
	readily give a hydrocar	bon product in good yields[CBs	SE PM'	r (f9)97) ²	(d) sp^3d	
	(a) RCOOK — Oxidation Elect	roly is	151.	Silver acetylide when h (a) C_2H_2	eated with HCl gi (b) H_2	ves
	(b) $RCOOAg \xrightarrow{I_2}$			= =	(d) None of thes	20
	(c) $CH_3 - CH_3 \xrightarrow{Cl_2}$		4	(c) C_2H_4		
	nv		152.	When sodium reacts we the following hydrocard		which of
	(d) $(CH_3)_2 CCl - \frac{C_2 H_5 OH}{C_2 H_5 OH}$	\rightarrow		the following nyurocart	[NCERT 1984; I	3HU 1982]
140.	Out of the following f	ractions of petroleum, the		(a) Methane	(b) Ethane	
		boiling point is or Which of		(c) Butane	(d) Ethene	
		ned at lowest temperature			DPMT 1983; CBSE I	PMT 1989]
	=	n of petroleum[MP PMT 1993; M	MP PEI	(c) Covalent solid	(b) Ionic solid(d) Not possible	
	(a) Kerosene(c) Gasoline	(b) Diesel oil (d) Heavy oil	154.	The shape of ethane is	_	: CEE 1995]
141.		r used by miners works on	-54.	(a) Triangular		1999]
141.	the principle of	[AMU 1984]		(c) Linear	(d) None of thes	se
	= =	ites of diffusion of gases	155.	CH_3MgI will give meth	rkee 1995]	
	(b) Avogadro's hypothe			(a) C_2H_5OH	(b) $CH_3 - CH_2 -$	NH_2
	(c) Gay-Lussac's law of	f gaseous volumes		(c) $CH_3 - CO - CH_3$	(d) All of these	
	(d) Berzelius hypothes	is	156.	Propane-1-ol can be pre	epared from prope	ene by its
142.	Methane can be prepar	ed by [DCE 2001]		reaction with	_	PMT 2003]
	(a) Wurtz's reaction	(b) Decarboxylation		(a) CH_3COOH	(b) H_3BO_3	
	(c) Hydrogenation reac	ction (d) All of these		(c) $B_2H_6/NaOH, H_2O_2$		
143.	The most strained cycle	oalkane is [IIT-JEE 1981]	157.	The process in which		
	(a) Cyclopropane	(b) Cyclobutane		broken down into controlled pyrolysis, is	•	bons by
	(c) Cyclopentane	(d) Cyclohexane		controlled pyrolysis, is		PMT 2002]
144.		vith chlorine in dark[Pb. PMT 20	000]	(a) Hydrolysis	(b) Cracking	-
	(a) C_2H_4	(b) C_2H_2		(c) Oxidation	(d) Reduction	
	(c) <i>CH</i> ₄	(d) CH ₃ CHO	158.	Successive alkanes diffe	· .	PMT 2002]
145.	Main constituent of ma	rsh gas is		(a) CH_2	(b) > CH	
		o; MP PMT 1994; AFMC 1997]		(c) $-CH_3$	(d) C_2H_4	
	(a) C_2H_2	(b) <i>CH</i> ₄	159.	General formula of alka		
	(c) H_2S	(d) CO		(a) $C_n H_{2n+2}$	ET 1979; Manipal N	иее 1995]
146	_	method can be used for the		(a) $C_n H_{2n+2}$ (c) $C_n H_{2n}$	(b) $C_n H_{2n-1}$ (d) $C_n H_{2n+1}$	
-70.	preparation of methan			$C_n \Pi_{2n}$	$(\mathbf{u}) \cup_{n} \mathbf{n}_{2n+1}$	

- **160.** Methane and ethane both can be prepared in one step by which of the following compound [BHU 2004]
 - (a) C_2H_4
- (b) CH_3O
- (c) CH_3Br
- (d) CH_3CH_2OH
- **161.** Photochemical chlorination of alkane is initiated by a process of **[Kerala PMT 2004]**
 - (a) Pyrolysis
- (b) Substitution
- (c) Cracking
- (d) Peroxidation
- (e) Homolysis
- **162.** A petroleum fraction having boiling range 70- $200^{\circ}C$ and containing 6-10 carbon atoms per molecule is called

[UPSEAT 2004]

- (a) Natural gas
- (b) Gas oil
- (c) Gasoline
- (d) Kerosene
- 163. Producer gas is a mixture of

[Pb. CET 2002; UPSEAT 2004]

- (a) CO and N_2
- (b) CO_2 and H_2
- (c) N_2 and O_2
- (d) CH_4 and N_2
- 164. The highest boiling point is expected for [DEC. 2003]
 - (a) n butane
 - (b) iso-octane
 - (c) n octane
 - (d) 2,2,3,3-tetramethyl butane
- **165.** Which of the following is a good conductor of heat of electricity **[Pb. CET 2003]**
 - (a) Diamond
- (b) Graphite
- (c) Anthracite
- (d) Charcoal
- **166.** Which one of the following has the minimum boiling point

[AIEEE 2004]

- (a) 1-Butene
- (b) 1-Butyne
- (c) n-Butane
- (d) Isobutane
- **167.** Octane number can be changed by [AFMC 2004]
 - (a) Isomerisation
- (b) Alkylation
- (c) Cyclisation
- (d) All of these
- **168.** Gasoline has composition
- [AFMC 2004]
- (a) $C_8 C_{12}$
- (b) $C_2 C_5$
- (c) $C_6 C_{11}$
- (d) None of these
- **169.** The complete combustion of CH_4 gives [BHU 2004]
 - (a) $CO + H_2$
- (b) $CO + N_2$
- (c) $CO_2 + H_2O$
- (d) $CO + N_2O$
- 170. Which of the following has highest knocking

[UPSEAT 2004]

- (a) Olefins
- (b) Branched chain olefins
- (c) Straight chain olefins
- (d) Aromatic hydrocarbons
- 171. Which one of the following compounds gives Where methane on treatment with water[Kerala PMT 2004; MH CET 2004]
 - (a) Al_4C_3
- (b) *CaC*₂
- (c) VC
- (d) SiC
- (e) $B_{A}C$

- 172. Pick out the alkane which differs from the other members of the group. [KCET 2004]
 - (a) 2,2-dimethyl propane
 - (b) Pentane
 - (c) 2-methyl butane
 - (d) 2,2-dimethyl butane
- **173.** 2-Methylbutane on reacting with bromine in the presence of sunlight gives mainly **[AIEEE 2005]**
 - (a) 1-bromo-2-methylbutane
 - (b) 2-bromo-2-methylbutane
 - (c) 2-bromo-3-methylbutane
 - (d) 1-bromo-3-methylbutane
- 174. Of the five isomeric hexanes, the isomer which can give two monochlorinated compounds is[AIEEE 2005]
 - (a) *n*-hexane
 - (b) 2, 3-dimethylbutane
 - (c) 2, 2-dimethylbutane
 - (d) 2-methylpentane
- 175. The product obtained on reaction of C_2H_5Cl with hydrogen over palladium carbon is **[AFMC 2005]**
 - (a) C_3H_8
- (b) C_4H_{10}
- (c) C_2H_6
- (d) C_2H_4

Alkene

1. Addition of bromine to 1, 3-butadiene gives

[CPMT 1987, 93]

- (a) 1, 2 addition product only
- (b) 1, 4 addition product only
- (c) Both 1, 2 and 1, 4 addition products
- (d) No reaction
- **2.** When ethylene bromide is treated with Zn, we get

[RPMT 1997]

- (a) Alkane
- (b) Alkene
- (c) Alkyne
- (d) All
- 3. Ethene when treated with Br_2 in the presence of CCl_4 which compound is formed

[RPMT 1997; DCE 2001; KCET (Med.) 1999]

- (a) 1, 2-dibromoethane
- (b) 1-bromo-2-chloroethane
- (c) Both (a) and (b)
- (d) 1, 1, 1-tribromoethane
- **4.** In a reaction

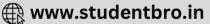
$$CH_2 = CH_2 \xrightarrow{\text{Hy pochlorous}} M \xrightarrow{R} |$$

$$CH_2 = OH$$

$$CH_2 = OH$$

Where M = molecule; R = reagent M and R are2004] [CBSE PMT 1997; CPMT 2001]

- (a) CH_3CH_2Cl and NaOH
- (b) $CH_2Cl CH_2OH$ and aq. $NaHCO_3$
- (c) CH_3CH_2OH and HCl



- (d) $CH_2 = CH_2$ and heat
- Alkenes usually show which type of reaction 5.

[AIIMS 1999; MADT Bihar 1980]

- (a) Addition
- (b) Substitution
- (c) Elimination
- (d) Superposition
- 6. The propene reacts with HBr to form

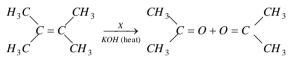
[AIIMS 1999; RPET 1999]

- (a) Ethane
- (b) Hexane
- (c) 1-bromo-propane
- (d) 2-bromo propane
- Ethylene may be obtained by dehydration of 7. which of the following with concentrated H_2SO_4 at 160 - 170° C

[RPET 1999]

- (a) C_2H_5OH
- (b) CH₃OH
- (c) $CH_3CH_2CH_2OH$
- (d) $(CH_3)_2 CHCH_2 OH$

8.



X in the above reaction is

- (a) HNO_3
- (c) O_3
- (d) $KMnO_4$
- The disappearance of the characteristic purple Q. colour of KMnO4 in its reaction with an alkene is the test for unsaturation. It is known as

[CPMT 1989, 94; CBSE PMT 1990]

- (a) Markownikoff's test (b) Baeyer's test
- (c) Wurtz's test
- (d) Grignard test
- A gas formed by the action of alcoholic KOH on 10. ethyl iodide, decolourises alkaline KMnO_4 . The gas

[KCET 2003]

- (a) C_2H_6
- (b) CH₄
- (c) C_2H_2
- (d) C_2H_4
- $CH_3 CH_2 Cl \xrightarrow{alc.KOH} A$, the product is 11.

[CPMT 2003]

- (a) CH_3CH_2OK
- (b) CH₃CHO
- (c) $CH_3CH_2OCH_2CH_3$
- (d) $CH_2 = CH_2$
- The final product formed when ethyl bromide is 12. treated with excess of alcoholic KOH is [MP PET 1999]
 - (a) Ethylene
- (b) Ethane
- (c) Ethyne
- (d) Vinyl bromide
- Which of the following hydrocarbons cannot be 13. obtained by Sabatier and Senderen's reaction
 - (a) CH_{4}
- (b) C_2H_6
- (c) C_3H_8
- (d) All
- When 3, 3-dimethyl-2-butanol is heated with H_2SO_4 the major product obtained is [CBSE PMT 1995]

- (a) cis and trans isomers of 2, 3-dimethyl-2butene
 - (b) 3, 3-dimethyl-1-butene
 - (c) 2, 3-dimethyl-2-butene
 - (d) 2, 3-dimethyl-1-butene
- The intermediate during the addition of HCl to 15. propene in the presence of peroxide is[IIT-JEE 1997]
 - (a) $CH_3 CHCH_2 Cl$ (b) $CH_3 CHCH_3$
 - (c) $CH_3CH_2CH_2$
 - (d) $CH_3CH_3CH_3$
- $CH_2 = CH_2 \xrightarrow{KMnO_4} X$. Product 'X' in reaction is

[RPMT 2003]

- (a) Ethylene glycol
- (b) Glucose
- (c) Ethanol
- (d) All of these
- Which of the following compounds represents acrylonitrile

[JIPMER 1997]

- (a) Vinyl cyanide
- (b) Cyanoethene

[CPMT 1985-23]ne nitrile

- (d) All of them
- When acetylene reacts with arsenic trichloride in the presence of anhydrous aluminium chloride, it produces

[AFMC 1999]

- (a) Lewisite
- (b) β -chlorovinyl dichloroarisine
- (c) Nitrobenzene
- (d) Both (a) and (b)
- Ozonolysis of which one of the following will give 19. two molecules of acetaldehyde

[Bihar MEE 1997; MP PET 2000]

- (a) 1-butene
- (b) 2-butene
- (c) 1-pentene
- (d) 2-pentene
- (e) None of these
- In which of the following, addition of HBr does not take place against Markownikoff's rule or Anti-Markownikoff addition of *HBr* observed for

[IIT-JEE 1985; CBSE PMT 1994; MADT Bihar 1995; MP PMT 1999; AMU 2002]

- (a) Propene
- (b) But-1-ene
- (c) But-2-ene
- (d) Pent-2-ene
- Which one of the following characteristics apply to both ethene and ethyne
 - (a) Explode when mixed with chlorine
- (b) Decolourise Baeyer's reagent giving brown precipitate
 - (c) Rapidly absorbed by cold conc. H_2SO_4
- (d) Form white precipitate with silver nitrate solution
- 22. Which of the following has highest knocking
 - (a) Aromatic hydrocarbons
 - (b) Olefins





- (c) Branched chain paraffins
- (d) Straight chain paraffins
- Dilute aqueous KMnO_4 , at room temperature 23. reacts with R - CH = CH - R to give [Roorkee 1992]
 - (a) R CHO
- (b) R COOH
- (c) RCHOH CHOHR
- (d) $CO_2 + H_2O$
- Aqueous sulphuric acid reacts with 2-methyl-1-24. butene to give predominantly
 - (a) Isobutyl hydrogen sulphate
 - (b) 2-methyl-2-butanol
 - (c) 2-methyl-1-butanol
 - (d) Secondary butyl hydrogen sulphate
- How can ethene be produced from ethanol [BHU 1996] 25.
- (a) By dehydrohalogenation
 - (b) By dehydrogenation
 - (c) By dehydration with conc. H_2SO_4 at $170^{\circ}C$
 - (d) By reduction with hydrogen iodide
- 26. Baeyer's reagent is used in the laboratory for [CBSE PMT 1991, 92; AIIMS 1998; AFMC 1999]
 - (a) Detection of double bonds
 - (b) Detection of glucose
 - (c) Reduction
 - (d) Oxidation
- Isopropyl alcohol is obtained by reacting which of 27. the following alkenes with conc. H_2SO_4 and H_2O

[MP PMT 1999]

- (a) Ethylene
- (b) Propylene
- (c) 2-methyl propene
- (d) Isoprene
- 28. Which of the following compound is produced when $CH_2 = CH - (CH_2)_2 COOH$ reacts with HBr in presence of peroxides [AIIMS 2000]
 - (a) $CH_3CH(CH_2)_5COOH$
 - (b) $BrCH_2CH_2(CH_2)_5COOH$
 - (c) $CH_3CH_2CH_2(CH_2)_5COOH$
 - (d) CH₃CH₂BrCH₂CH₂COOH
- One mole of each of the following alkenes is 29. catalytically hydrogenated. The quantity of heat evolved will be the lowest in the case of
 - (a) 1-butene
- (b) Trans-2-butene
- (c) Cis-2-butene
- (d) 1, 3-butadiene
- **30.** Which of the following is not used to distinguish ethene from ethane

[KCET (Med.) 2001; UPSEAT 2002; CBSE PMT 2002]

- (a) Iodine in CCl₄
- (b) Bromine in CCl₄
- (c) Alkaline KMnO₄
- (d) Ammonical Cu_2Cl_2
- A hydrocarbon X adds on one mole of hydrogen to 31. give another hydrocarbon and decolourised bromine water. X reacts with $KMnO_4$ in presence of acid to give two moles of the same carboxylic acid. The structure of *X* is

[JIPMER 2001]

- (a) $CH_2 = CH CH_2CH_2CH_3$
- (b) $CH_3CH_2CH_2 CH = CHCH_3$

- (c) $CH_3CH_2CH = CHCH_2CH_3$
- (d) $CH_3CH = CHCH_2CH_2CH_3$
- When 2-bromobutane reacts with alcoholic KOH, 32. the reaction is called [KCET (Med.) 2001]
 - (a) Halogenation
- (b) Hydrogenation
- (c) Chlorination
- (d) Dehydro-

- halogenation [Roorkee 1992]
 1, 3-butadiene reacts with ethylene to form[BHU 2001]
 - (a) Benzene
- (b) Cyclohexane
- (c) Cyclohexene
- (d) 2, 3 dimethyl butane
- Ethylene reacts with ozone gas to form the compound [UPSEAT 2001]
 - (a) HCHO
- (b) C_2H_5OH
- (c) $O \stackrel{CH_2-O}{\underset{CH_2-O}{\leftarrow}}$
- (d) CH₃CHO
- Oils are converted into fats by [Kerala (Med.) 2002] 35.
 - (a) Hydration
- (b) Decarboxylation
- (c) Hydrogentation
- (d) Dehydrogenation
- (e) Hydrogenolysis
- Which process converts olefins into parafins 36.

[MP PET 2002]

- (a) Halogenation
- (b) Dehydration
- (c) Hydrogenation
- (d) Hydrolysis
- Of the following the formula which represents a saturated cyclic compound is
 - (a) C_3H_6
- (b) C_3H_8
- (c) C_8H_{10}
- (d) $C_8 H_{12}$
- 38. In a reaction, if half of the double bond is broken and two new bonds are formed, this is a case of

[AMU 1983; NCERT 1978; CPMT 1983]

- (a) Elimination
- (b) Addition
- (c) Displacement
- (d) Rearrangement
- Which of the following are formed on addition reaction of DCI with 3-methyl-1-butene[Roorkee 2000]

 - (a) $CH_2DCHClCH(CH_3)_2$ (b) $CH_2DCH_2CCl(CH_3)_2$
 - [Roorkee 2000] CH₃CDClCH(CH₃)₂
- (d) $ClCH_2CHDCH(CH_3)_2$
- Major product of the following reaction is

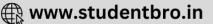
$$CH_3 - \overset{|}{C} - CH_2 - CH_3 + alco.KOH \rightarrow \quad \text{[MP PMT 1986]}$$

- (a) Butene-1
- (b) Butene-2
- (c) Butane
- (d) Butyne-1
- Cyclopentene on treatment with alkaline KMnO₄ gives

[CPMT 1987]

- (a) Cyclopentanol
- (b) trans 1, 2-cyclopentanediol
- (c) cis 1, 2-cyclopentanediol
- (d) 1: 1 mixture of cis and trans 1, 2cyclopentanediol
- **42.** Which of the following is the most stable alkene





[AIIMS 1998; KCET (Med.) 2000; CPMT 2003] (a) Ethyl alcohol (b) Acetaldehyde (a) $R_2C = CR_2$ (b) RCH = CHR(c) Acetic acid (d) Ethylene glycol (c) $RCH_2 = CH_2R$ (d) $CH_2 = CH_2$ A gas decolourised $KMnO_4$ solution but gives no 53. precipitate with ammoniacal cuprous chloride is **43.** Ethene gives with acidic $KMnO_A$ solution [MP PMT 1997] or Which of the following gases does not give a (b) Ethylene oxide (a) Ethylene glycol precipitate with ammoniacal solution of silver (c) Formaldehyde (d) Acetaldehyde nitrate but decolourizes KMnO₄ (neutral or In paraffins, with the increasing molecular 44. slightly alkaline) [CPMT 1974] weight, it is found that [NCERT 1974, 77; CPMT 1974, 77, 78; (a) Freezing point decreases MP PMT 1996; MP PET 1996, 99] (b) Boiling point decreases (a) Ethane (b) Methane (c) Boiling point increases (c) Ethene (d) Acetylene (d) Vapour pressure decreases A hydrocarbon reacts with hypochlorous acid to When alcoholic solution of ethylene dibromide is give 1-chloro-2-hydroxyethane. The hydrocarbon heated with granulated zinc, the compound formed is [CPMT 1990] [CBSE PMT 1989] (a) Ethylene (b) Ethyne (b) Methane (a) Ethylene (c) Cyclobutane (d) Butane (c) Ethane (d) Acetylene A gas formed by the action of alcoholic KOH on ethyl iodide, decolorises alkaline $KMnO_4$ solution. When ethene is heated at 400°C under high 55. pressure, the product is/are (a) Carbon and H_2 (b) Polyethylene [CPMT 1974, 91; MP PET 1985; IIT-JEE 1982] (a) CH_{4} (b) C_2H_6 (c) Acetylene and H_2 (d) None of these (d) C_2H_2 (c) C_2H_4 Which decolorize aqueous bromine and gives Markownikoff's rule white fumes of *HCl* on reaction with *PCl*₅ [Pb. PMT 1999] provides guidance of 47. addition of HBr on (a) $CH_3COCH_2CH = CH_2$ [MNR 1994] (b) $CH_3CH_2CH_2CH_3CH_3$ (a) $CH_2 = CH_2$ (b) $CH_3 - CH_2 - CH_3$ (c) $CH_3CH = CHCH_2CH_2OH$ (c) $CH_3CH = CHCH_3$ (d) $CH_2 = CHBr$ (d) CH₃OCH₂CH₂CH₂CH₂OH Ethyl bromide gives ethylene when reacted with 48. [CPMT 1982, 93; RPET 2000; Pb. PMT 2001] During debromination of meso-dibromobutane, 57. the major compound formed is (a) Ethyl alcohol (b) Dilute H_2SO_4 [IIT-JEE 1997] (a) *n*-butane (b) 1-butane (c) Aqueous KOH (d) Alcoholic KOH (c) cis-2-butene (d) trans-2-butene Ethylene is prepared by the dehydration of 49. What product is formed when 1-chlorobutane [CPMT 1974, 79; DPMT 1985; BHU 1989] react with alcoholic KOH [RPMT 2002] (a) Ethyl alcohol (b) Methyl alcohol (a) 1-butene (b) 2-butene (c) Acetic acid (d) Oxalic acid (c) 1-butanol (d) 2-butanol Which reactions are most common in alkenes 50. The olefin which on ozonolysis gives CH₃CH₂CHO (a) Electrophilic substitution reactions [Roorkee 1992] and CH₃CHO is (b) Nucleophilic substitution reactions (a) 1-butene (b) 2-butene (c) Electrophilic addition reactions (c) 1-pentene (d) 2-pentene (d) Nucleophilic addition reactions Bond length between carbon-carbon in ethylene 60. mixture of 1-chloropropane 51. molecule is chloropropane when treated with alcoholic KOH [MP PET 1997] gives [NCERT 1990] (a) 1.54 \mathring{A} (b) 1.35 Å(a) 1-propene (b) 2-propene (c) 1.19 Å (d) 2.4 Å(d) All the three (c) Isopropylene The compound having both sp and sp^2 hybridised The compound formed by passing ethylene gas 52. into cold alkaline solution of KMnO4 is carbon atom is [IIT-JEE 1981]

(a) Propene

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(c) Propadiene

(b) Propyne

(d) None of these

[NCERT 1974, 81; CPMT 1979, 86, 88;

MP PET 1985, 95; AFMC 1998]

- 62. The halogen which is most reactive in the halogenation of alkenes under sunlight is [IIT-JEE 1981]
 - (a) Chlorine
- (b) Bromine
- (c) Iodine
- (d) All equal
- When ethene reacts with bromine, it forms

[AFMC 2000; KCET 2001]

- (a) Chloroethane
- (b) Ethylene dibromide
- (c) 1 bromopropane
- (d) 1,2-dichloroethene
- **64.** Paraffins are soluble in
- [NCERT 1978]
- (a) Distilled water
- (b) Benzene
- (c) Methanol
- (d) Sea water
- 65. Addition of HCl to propene in presence of peroxides gives

[BHU 1981, 98]

- (a) 1-Chloropropane
- (b) 2-Chloropropane
- (c) 3-Chloropropane
- (d) Propene dichloride
- The name of the product obtained by the addition of HI to propene in presence of peroxide catalyst is [KCET 2000]
 - (a) Isopropyl iodide
- (b) 2-Iodopropene
- (c) 2-Iodopropane
- (d) 1-Iodopropane
- **67.** In the reaction $C_2H_5CH = CH_2 + H X \rightarrow \text{Product.}$ What is the product [BHU 2002]
 - (a) $C_2H_5 CH_3$
 - (b) $C_2H_5CH_2 CH_2X$
 - (c) $C_2H_5 CHX CH_3$
 - (d) $CH_3 CH_2X CH = CH_2$
- 68. Alkene can be prepared from alkyl halide by the following reagent $R - X + Nu^- \rightarrow \text{Alkene} + NuH \text{ [RPET 2000]}$
 - (a) Alc. KOH + heat
- (b) Aq. KOH + cold water
- (c) NaOH
- (d) LiOH
- **69.** 2-chlorobutane is heated with alcoholic *NaOH*, the product formed in larger amount is [RPET 1999; AMU 2000]
 - (a) 1-Butene
- (b) 1-Butyne
- (c) 2-Butene
- (d) All of these
- 70. Ethylene has high b.p. and high vapour pressure at 100 ° C and does not dissolve in water. Hence ethylene is separated by this method
 - (a) Simple distillation
- (b) Vacuum distillation
- (c) Vapour distillation (d) Alkali treatment
- 71. Addition of bromine to 1, 3-butadiene gives[AMU 1999] 79.
 - (a) 1, 4-addition product only
 - (b) 1, 2-addition product only
 - (c) Both 1, 2-and 1, 4 addition product
 - (d) None of these
- In the presence of peroxide, hydrogen chloride and hydrogen iodide do not give anti-Markovnikoff's addition to alkenes because[IIT-JEE Screening 2001]
 - (a) Both are highly ionic
 - (b) One is oxidising and the other is reducing
 - (c) One of the steps is endothermic in both the
 - (d) All the steps are exothermic in both the cases

- The compound most likely to decolourize a solution of potassium permanganate is[NCERT 1978]
- (a) CH_3CH_3

73.



(c) $CH_3CH = CHCH_2CH_3$

$$CH_3$$

- (d) $CH_3 C CH_3$
- Ethylene is converted to X on passing through a mixture of an acidified aqueous solution of palladium chloride and cupric chloride. Which of the following reagents readily take part in addition reaction with X[UPSEAT 2003]
 - (a) Br_2
- (b) HBr
- (c) HCl
- (d) HCN
- Addition of HCl does not obey antimarkownikoff's rule because

[UPSEAT 2003]

- (a) It is a strong acid (b) It is a gas
- (c) Its bond energy is high (d)Its bond energy is less
- Correct statement about1, 3-dibutene[UPSEAT 2003]
- (a) Conjugated double bonds are present
 - (b) Reacts with HBr
 - (c) Forms polymer
 - (d) All of these
 - At low temperatures, the slow addition of molecular bromine to $CH_2 = CH - CH_2 - C \equiv CH$ gives

[Roorkee Qualifying 1998]

- (a) $CH_2 = CH CH_2 CBr = CHBr$
- (b) $BrCH_2 CHBr CH_2 C \equiv CH$
- (c) $CH_2 = CH CH_2 CH_2 CBr_3$
- (d) $CH_3 CBr_2 CH_2 C \equiv CH$
- PCl_5 reacts with propanone, to give [Pb. PMT 2001] [UPSEAT 1999]
 - (a) vic-dichloride
- (b) Propanal
- (c) propane-chloride
- (d) gem-dichloride

The compounds that will give an isomer of 2; 2dimethyl propane on catalytic hydrogenation are [AMU 19

- (1) $CH_3CH = C CH_3$
- (2) $CH_3CH = CHCH_3$

 CH_3 CH_3

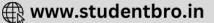
 CH_3

(3) $CH_3C = CHCH_2CH_3$ (4) $CH_3C = C - CH_3$

- (a) 1 and 4
- (b) 2 and 4
- (c) 1 and 3
- (d) 1 and 2
- **80.** Alkene $R-C-H=CH_2$ reacts readily with B_2H_6 and the product on oxidation with alkaline hydrogen peroxides produces







- (a) $R CH_2 CHO$
- (b) $R CH_2 CH_2 OH$
- (c) $R-C-CH_3$ $\stackrel{\scriptscriptstyle{\sqcap}}{o}$
- (d) $R CH CH_3$ OH OH
- Bayer's reagent is used for detection of [RPMT 2002]
 - (a) Amines
- (b) Glucose
- (c) Unsaturated bond (d) Alcohol
- Which of the following is(are) example(s) of nucleophilic addition reaction in case of acetylene
 - (a) Addition of water
- (b) Addition of HCN
- (c) Addition of AsCl₃
- (d) All
- Structural formula for lewisite is

CHCl

CHCl 2

(a) |

(b) |

CHAsCl 3

CHAsCl 3

CHCl

- (d) None of these (c) | CHAsCl 2
- 84. Propene when heated with chlorine at about 500°C forms

[MP PET 1997]

- (a) $CH_2Cl.CH = CH_2$
- (b) CH₃.CHCl.CH₂Cl
- (c) CH₂Cl.CHCl.CH₂Cl
- (d) All the three
- 85. PVC is obtained from vinyl chloride by a reaction called
 - (a) Addition
- (b) Isomerization
- (c) Polymerization
- (d) Substitution
- **86.** Reaction of Br_2 on ethylene in presence of NaCl
 - (a) $BrCH_2 CH_2Br$
- (b) $ClCH_2 CH_2Br$
- (c) Both (a) and (b)
- (d) None of these
- $CH_3 CH = CH_2 + HBr \longrightarrow \dots$, the product 87. formed is[AIIMS 1983; CPMT 1997; RPMT 1999, 2003]
 - (a) $CH_3 CH_2 CH_2 Br$ (b) $CH_3 CHBr CH_3$
 - (c) $BrCH_2 CH = CH_2$
- (d) $CH_2 = C = CH_2$
- The product of reaction between propene and HBr in the presence of a peroxide is
 - (a) $CH_3 CH_2 CH_2Br$ (b) $CH_3 CHBr CH_3$
 - (c) $CH_3 CH_2Br$
- (d) $CH_3 CH = CHBr$
- Ozonolysis of 2-methyl butene-2 yields
 - (a) Only aldehyde
 - (b) Only ketone
 - (c) Both aldehyde and ketone
 - (d) None of these
- The final product formed by the ozonolysis of 90. compound $RCH = CR_2$ is [NCERT 1978]
 - (a) RCHO
- (b) R_2CO
- (c) Both (a) and (b)
- (d) None of these
- Which one is an unsaturated compound [BIT 1990]

- (a) $C_6 H_{14}$
- (b) C_4H_8
- (c) C_3H_7OH
- (d) CH_3OH
- Ethyl alcohol on heating with conc. H_2SO_4 gives

[EAMCET 1979; MP PMT 1996]

- (a) $CH_3COOC_2H_5$
- (b) C_2H_6
- (c) C_2H_4
- (d) C_2H_2
- Monohalides on reacting with alcoholic KOH give [MP PET 1982, 86; DPMT 1981; CPMT 1979, 83]
 - (a) Alkanes
- (b) Alkenes
- (c) Alkynes
- (d) Aromatic

hydrocarbons

- **94.** Ethylene is a member of.... series [BHU 1979]
 - (a) Alkyne
- (b) Olefin
- (c) Paraffin
- (d) Amine
- In a double bond between two carbon atoms of ethene, there are [NCERT 1981]
 - (a) Two sigma bonds perpendicular to each other
 - (b) One sigma and one pi bond
 - (c) Two pi bonds perpendicular to each other
 - (d) Two pi bonds at an angle of 60°
- 96. The formation of alkene from alkyl halide is an example of

[CPMT 1983; AMU 1982; Pb. CET 1986]

- (a) Addition
- (b) Elimination
- (c) Substitution
- (d) (a) and (c)
- 97. In the following reaction

$$CH_3 - CH_2 - CH_2 - CH_3 \xrightarrow{H_2SO_4}$$
 [AIIMS 1983]

- (a) $CH_3CH = CHCH_3$ predominates
- (b) $CH_2 = CHCH_2CH_3$ predominates
- (c) Both are formed in equal amounts
- (d) The amount of production depends on the nature of catalyst
- 98. The compound B formed in the following sequences of reactions is

$$CH_3CH_2CH_2OH \xrightarrow{PCl_3} A \xrightarrow{Alco.KOH} B$$

[NCERT 1981]

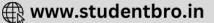
- (a) Propyne
- (b) Propene
- (c) Propanol
- (d) Propane
- n-propyl bromide on treatment with ethanolic 99. potassium hydroxide produces[IIT-JEE 1987; MP PMT 1997]
 - (a) Propane
- (b) Propene
- (c) Propyne
- (d) Propanol
- 100. The dehydrohalogenation of neopentyl bromide with alcoholic KOH mainly gives

[IIT-JEE (Screening) 1990; MP PET 1993]

- (a) 2-methyl-1-butene (b) 2-methyl-2-butene
- (c) 2, 2-dimethyl-1-butene (d)
- 2-butene
- 101. Which is the most reactive hydrocarbon in the following







[JIPMER 2002]

- (a) Ethane
- (b) Ethyne
- (c) Ethene
- (d) Methane
- **102.** Shape of ethylene molecule is
 - e is [MP PET 1993]
 - (a) Tetrahedral
- (b) Pyramidal
- (c) Planar
- (d) Linear
- **103.** Electrophilic addition on a carbon-carbon double bond involves the intermediate formation of a more stable carbocation. This statement is called
 - (a) Saytzeff's rule
- (b) Baeyer's effect
- (c) Markownikoff's rule (d) None of these
- **104.** $CH_2 = CHCl$ reacts with HCl to form[CPMT 1985, 93]
 - (a) $CH_2Cl CH_2Cl$
- (b) $CH_3 CHCl_2$
- (c) $CH_2 = CHCl.HCl$
- (d) None of these
- **105.** Deviation from Markownikoff's rule occurs in presence of
 - (a) Zinc
- (b) Peroxides
- (c) Hg Zn / HCl
- (d) All of these
- 106. Presence of peroxides affects the addition of [BHU 1987] $_{13}$. How many gm of bromine will react with 21 gm
 - (a) HBr
- (b) HCl
- (c) HI
- (d) All of these
- **107.** Catalyst used in dimerisation of acetylene to prepare chloroprene is **[BHU 1984]**
 - (a) $HgSO_4 + H_2SO_4$
- (b) Cu_2Cl_2
- (c) $Cu_2Cl_2 + NH_4Cl$
- (d) $Cu_2Cl_2 + NH_4OH$
- 108. Chloroprene is
 - (a) 2-chloro-1, 3-butadiene
 - (b) 3-chloro-2, 3-butadiene
 - (c) 2, 3-dichlorobutadiene
 - (d) None of these
- **109.** Chloroprene is used in making [MP PET 1985]
 - (a) Synthetic rubber
- (b) Plastic
- (c) Petrol
- (d) All of these
- **110.** When isobutyl magnesium bromide in dry ether is treated with absolute ethyl alcohol, the products formed are

[IIT-JEE 1995]

- (a) $CH_3 CH CH_2OH$ and CH_3CH_2MgBr CH_3
- (b) $CH_3 CH CH_2 CH_2 CH_3$ and Mg(OH)Br CH_3
- (c) $CH_3 CH CH_3$ and $CH_3 CH_2OMgBr$ CH_3
- (d) $CH_3 CH CH_3$, $CH_2 = CH_2$ and Mg(OH)Br CH_3
- 111. The predominant product formed, when 3-methyl-2-pentene reacts with *HOCl*, is [IIT-JEE 1995]

(a)
$$CH_3 - CH_2 - C - CH - CH_3$$

Cl Cl

(b)
$$CH_3 - CH_2 - C - CH - CH_3$$

 CH_3

ЭН

(c)
$$CH_3 - CH_2 - C - CH - CH_3$$

 $CH_3 Cl$

 $CH_3 OH$

(d)
$$CH_3 - C - CH - CH_3$$

$$CH_3$$

112. Which of the following occurs easily in ethylene

[MNR 1987; NCERT 1979]

- (a) Addition
- (b) Substitution
- (c) Elimination
- (d) Rearrangement
- How many gm of bromine will react with 21 gm C_3H_6

[MP PET 1985]

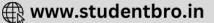
- (a) 80
- (b) 160
- (c) 240
- (d) 320
- 114. Conjugate double bond is present in [MP PMT 1987]
 - (a) Propylene
- (b) Butadiene
- (c) Isobutylene
- (d) Butylene
- 115. On passing vapours of an organic liquid over finely divided Cu at $573\,K$ the product was an alkene. This reaction is
 - (a) Catalytic oxidation of primary alcohol
- (b) Catalytic dehydrogenation of secondary alcohol
 - (c) Catalytic dehydrogenation of tertiary alcohol
 - (d) Catalytic dehydration of tertiary alcohol
- 116. The total number of $\operatorname{sigma} \sigma$ and $\operatorname{pi}(\pi)$ bonds in an ethylene molecule are
 - (a) 4σ , 2π
- (b) 4σ , 1π
- (c) 5σ , 2π
- (d) 5σ , 1π
- 117. Cyclic hydrocarbon molecule A has all the carbon and hydrogens in a single plane. All the carbon-carbon bonds are of same length and less that 1.54 \mathring{A} but more than 1.34 \mathring{A} . C-C-C bond angle will be [CBSE PMT 1989]
 - (a) 120°
- **(b)** 180°
- (c) 100°
- (d) 109°28′
- 118. General formula of alkenes is

[CPMT 1975, MNR 1987; NCERT 1987; MP PMT 1994]

- (a) $C_n H_{2n}$
- (b) $C_n H_{2n-2}$
- (c) $C_n H_{2n+2}$
- (d) $C_n H_{2n-1}$
- **119.** The product of following reaction is







$$CH_{3}$$

$$CH_{3} - \overset{|}{C} - CH = CH_{2} - \underbrace{\overset{(i) Hg(OAc)_{2}; H_{2}O}{(ii) NaBH_{4}}}$$

$$CH_{3}$$

[MP PMT 1986; MP PET 1997]

(a)
$$CH_3 - CH_2$$

$$CH_3 - CH_3 - CH_2$$

$$CH_3 OH$$

$$CH_3$$

(b)
$$CH_3 - \overset{\downarrow}{C} - CH_2 - CH_2OH$$

 CH_3

$$OH \ CH_3$$
 (c) $CH_3 - C - CH - CH_3$

(d)
$$HOCH_2 - CH_2 - CH_2 - CH_2$$

- 120. Which one of the following organic compounds decolourizes an alkaline KMnO₄ solution[CPMT 1987, 93]
 - (a) CS_2
- (b) C_3H_6
- (c) C_3H_8
- (d) CH₃OH
- 121. Decolourization of alkaline $KMnO_4$ is used as a test for

[AMU 1983]

- (a) Aromatic hydrocarbons
- (b) Olefinic hydrocarbons
- (c) Acetylenic hydrocarbons
- (d) Cycloalkanes
- 122. The reaction

$$CH_2 = CH_2 + H_2 \xrightarrow[250-300^{\circ}C]{Ni} CH_3 - CH_3$$
 is called

[MP PMT 1996; CBSE PMT 2001; MH CET 2001; BHU 2002]

- (a) Wurtz's reaction
- (b) Kolbe's reaction
- (c) Sabatier and Senderen's reaction
- (d) Carbylamine reaction
- 123. The alkene which on ozonolysis yields acetone is [MP PMT 1986, 2000]
 - (a) $CH_2 = CH_2$
 - (b) $CH_3 CH = CH_3$
 - (c) $(CH_3)_2 C = C(CH_3)_2$
 - (d) $CH_3 CH = CH CH_3$
- **124.** $CH_3CH = CHCHO$ oxidized to $CH_3CH = CHCOOH$ using
 - (a) Alkaline potassium permanganate
 - (b) Acidified potassium permanganate
 - (c) Selenium dioxide
 - (d) Osmium tetroxide

- 125. The order of increasing reactivity towards HCl of the following compounds will be
 - (1) $CH_2 = CH_2$
 - (2) $(CH_3)_2 C = CH_2$
 - (3) $CH_3CH = CHCH_3$

[MP PET 1994]

- (a) 1 < 2 < 3
- (b) 1 < 3 < 2
- (c) 3 < 2 < 1
- (d) 2 < 1 < 3
- 126. The reagent which is used to distinguish between propene and propyne is [MP PET 1994; IIT-JEE (Screening): AIIMS 2000; Pb. PMT 2002; BHU 2003]
 - (a) Bromine
- (b) Alkaline KMnO
- (c) Ammoniacal AgNO 3 (d) Ozone
- 127. Which one of the following reactions would be the best for the formation of 2-bromobutane[MP PET 1994]
 - (1) $CH_3CH = CHCH_2CH_3 \xrightarrow{HBr}$
 - (2) $CH_3CH_2CH = CH_2 \xrightarrow{HBr}$
 - (3) $CH_3CH = CHCH_3 \xrightarrow{Br_2}$
 - (4) $CH_3CH_2CH = CH_2 \xrightarrow{HBr}$
 - (a) 1

(b) 2

(c) 3

128. If HCl is added over $CH_{2} = C$

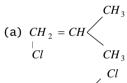
 CH_3

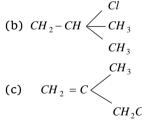
 CH_3

then

what is formed

[CPMT 1996]





- (d) None of these
- 129. Position of double bond in an organic compound is determined by [DCE 2001; RPMT 2002]
 - (a) Ozonolysis
- (b) Oxidation
- (c) Reduction
- (d) Hydrogenation
- 130. A gas decolourises Bayer's reagent but does not react with Tollen's reagent, this gas is [MP PMT 2001]
 - (a) Ethene
- (b) Ethyne
- (c) Ethane
- (d) Methane
- 131. Formation of 2-butene from 2-bromobutane is according to
 - [NCERT 1978] (a) Markowikoff's
- (b) Bayer
- (c) Saytzeff
- (d) Wurtz
- 132. An alkene on ozonolysis gave acetaldehyde the alkene is







- (a) Ethylene
- (b) Propene
- (c) 1-butene
- (d) 2-butene
- 133. Indicate the organic structure for the product expected when 2-methyl propene is heated with acetyl chloride in presence of anhydrous zinc chloride [CBSE PMT 1989]
 - (a) $CH_3 \overset{|}{C} CH_2 CO CH_3$
 - (b) $CH_3 C CH_2 CO CH_3$
 - (c) $CH_3 C O C \stackrel{Me}{\longleftarrow} Me$ $O CH_3 C O C \stackrel{Me}{\longleftarrow} Me$ $O CH_3$
 - (d) $CH_3 C C = CH_2$
- **134.** The reaction

$$CH_{3} \xrightarrow{CH_{3}} CH_{3} \xrightarrow{H_{2}SO_{4}} CH_{3} - \overset{|}{C} = CH_{2}$$

$$OH$$

is the example of

[AMU 1983]

- (a) Sulphonation
- (b) Dehydration
- (c) Alkylation
- (d) Decomposition
- 135. Olefins can be hydrogenated by
- [AIIMS 1991]
- (a) Zinc and HCl (c) Raney Ni and H_2
- (b) Nascent hydrogen
- (d) Lithium hydride in

ether

- **136.** Electrolysis of cold concentrated aqueous solution of potassium succinate yields
 - (a) Ethane
- (b) Ethyne
- (c) Ethene
- (d) Ethane-1, 2-diol
- 137. A hydrocarbon containing 2 carbon atoms gives Sabatier and Senderen's reaction but does not give precipitate with ammoniacal silver nitrate solution. The hydrocarbon in the question is [MADT Bihar 1983]
 - (a) Ethane
- (b) Acetylene
- (c) Ethylene
- (d) None of these
- **138.** The reaction

$$CH_2 = CH - CH_3 + HBr \rightarrow CH_3CHBr - CH_3$$
 is

[CBSE PMT 1996]

- (a) Nucleophilic addition
- (b)Electrophilic addition
- (c) Electrophilic substitution
- (d)
- 139. What is the product of the reaction of 1, 3butadiene with Br_2
 - (a) 1,4 -dibromobutene (b) 1,2 -dibromobutene
 - (c) 3,4-dibromobutene (d) 2,3-dibromo-2butene
- 140. An alkene given two moles of HCHO, one mole of CO2 and one mole of CH3COCHO on ozonolysis. What is its structure [Orissa JEE 2003]

(a) $CH_2 = C = CH - CH_2 - CH_3$

- (b) $CH_2 = CH CH CH = CH_2$
- (c) $CH_2 = C = C CH_3$ CH_3

 CH_{2}

- (d) $CH_2 = C = C CH = CH_2$
- **141.** In the reaction

$$CH_3CH = CH_2 + H_2O + [O] \xrightarrow{KMnO_4}$$

$$CH_3 - CH - CH_2 \xrightarrow{[O]} X + HCOOH$$

$$OH OH$$

X is

- (a) CH_3CH_2COOH
- (b) CH₃COOH
- (c) CH_3CH_2CHO
- (d) CH_3CH_2OH
- 142. Which of the following alkenes gives only acetic and on oxidation with potassium permanganate solution

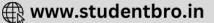
[MP PET 2003]

- (a) Ethylene
- (b) 1-Butene
- (c) Propene
- (d) 2-Butene
- 143. Butene-1 may be converted to butane by reaction with

[AIEEE 2003]

- (a) Zn-HCl
- (b) Sn-HCl
- (c) Zn-Hq
- (d) Pd/H_2
- 144. The major product formed when propene reacts [CPMith 613; inpropense of peroxides is [NCERT 1980; CBSE PM
 - (a) *n*-propyl bromide
- (b) Isopropyl bromide
- (c) *n*-propyl alcohol
- (d) 1, 3-dibromopropane
- 145. Ethyl hydrogen sulphate is obtained by the reaction of H_2SO_4 on [CPMT 1985]
- (b) Ethane
- (c) Ethyl chloride
- (d) Ethanol
- 146. Ethylene reacts with ozone to give [DPMT 1981]
 - (a) Formaldehyde
- (b) Ethyl alcohol
- (c) Ozonide
- (d) Acetaldehyde
- 147. Which of the following aliphatic compounds will discharge red colour of bromine
 - (a) C_2H_4 Free radical addition
- (b) C_3H_6
- (c) $C_4 H_8$
- (d) All of these
- 1480 Gsda J Hazioon can be done on
 - (a) $CH_3 CH = CH_2$
- (b) $CH_2 = CH_2$
- (c) $CH \equiv CH$
- (d) None of these
- 149. Addition of HI on the double bond of propene yields isopropyl iodide and not n-propyl iodide as the major product. This is because the addition proceeds through





[CPMT 1988]

- (a) A more stable carbonium ion
- (b) A more stable carbanion
- (c) A more stable free radical
- (d) None of the above being a concerted reaction
- 150. When butene-1 is mixed with excess of bromine, the expected reaction product is [CPMT 1974; BHU 1980]
 - (a) 1, 2-dibromobutane (b) 1, 1-dibromobutane
 - (c) 2, 2-dibromobutane (d) Perbromobutane
- **151.** A compound 'X' on ozonolysis forms two molecules of HCHO. Compound 'X' is[AIIMS 1987; CPMT 1993]phenyl propene is
 - (a) C_2H_4
- (b) C_2H_2
- (c) C_2H_6
- (d) $C_6 H_6$
- 152. For the reaction

$$CH_3 - CH = CH_2 + HOCl \rightarrow A$$
 the product A is

[Orissa JEE 2002]

- (a) $CH_3 CHCl CH_2OH$
- (b) $CH_3 CH CH_2 Cl$
- (c) $CH_3 CH_2 CH_2 COCl$

(d)
$$CH_3 - C - CH_3$$

$$OH$$

- **153.** $(CH_3)_2 C = CH \xrightarrow{\text{Cataly st}} \text{Optical isomers} [BHU 2003]$
 - (a) 2

- (b) 4
- (c) Zero
- (d) 3
- **154.** Isobutene $+HBr \xrightarrow{\text{Peroxide}}$ product is [BHU 2003]
 - (a) Tertiary butyl bromide (b) Isobutyl bromide
- - (c) Tertiary butyl alcohol
- (d) Isobutyl alcohol
- 155. Which of the following represents the given mode of hybridisation $sp^2 - sp^2 - sp - sp$ from left to right

[IIT-JEE (Screening) 2003]

- (a) $H_2C = CH C \equiv CH$
- (b) $HC \equiv C C \equiv CH$
- (c) $H_2C = C = C = CH_2$ (d) CH_2
- 156. "The negative part of addenda adds on to the carbon atom linked with least number of hydrogen atoms". This statement is called [DPMT 1982; AIIMSMASSDAFINIO 1984].lle
 - (a) Thiele's principle
- (b) Bayer's strain theory
- (c) Markownikoff's rule (d) Peroxide effect
- 157. The product obtained, heating ethanol with conc. H_2SO_4 at $165^{\circ} - 170^{\circ}$, is [MP PMT 2003]
 - (a) $(C_2H_5)_2SO_4$
- (b) $CH_2 = CH_2$
- (c) CH₃COOH
- (d) $C_2H_5HSO_4$
- **158.** Which of the following is the most stable
 - (a) 1-butene
- (b) 2-butene

- (c) 1-pentene
- (d) 2-pentene
- 159. Which doesn't follow Markownikoff's rule

[JEE Orissa 2004; MP PMT 2004; BCECE 2005]

- (a) $CH_3 CH = CH_2$
- (b) $CH_3CH = CHCH_3$
- (c) $CH_3 CH CH = CH_2$ CH_3
- (d) $CH_3 CH_2 CH = CH_2$
- 160. The product of acid catalyzed hydration of 2-[IIT JEE (Screening) 2004]
 - (a) 3-phenyl-2-propanol (b) 1-phenyl-2-propanol
 - (c) 2-phenyl-2-propanol (d) 2-phenyl-1-propanol
- **161.** A reagent used to test for unsaturation of allkene is

[BHU 2004]

- (a) conc. H_2SO_4
- (b) Ammonical Cu_2Cl_2
- (c) Ammonical AgNO 3
- (d) Solution
- of

 CCl_A

- 162. Propylene on hydrolysis with sulphuric acid forms [MH CET-2003]
 - (a) *n*-propyl alcohol
- (b) Isopropyl alcohol
- (c) Ethyl alcohol
- (d) Butyl alcohol
- 163. An alkene, on ozonolysis gives formaldehyde and acetaldehyde. The alkene is:
 - (a) Ethene
- (b) Propene
- (c) Butene-1
- (d) Butene-2
- cold alkaline KMnO₄ **164.** In the reaction, $H_2C = CH_2$ \rightarrow (A):

Product A is:

[Pb. CET 2000]

- (a) Ethylene glycol
- (b) Acetic acid
- (c) Ethane
- (d) Butyric acid
- 165. Using anhydrous AlCl3 as catalyst, which one of the following reaction produces ethylbenzene (PhEt)

[CBSE PMT 2004]

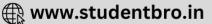
- (a) $H_2C = CH_2 + C_6H_6$
- (b) $H_3C CH_3 + C_6H_6$
- (c) $H_3C CH_2OH + C_6H_6$
- (d) $CH_3 CH = CH_2 + C_6H_6$
- of these does

[Orissa JEE 2005]

- (a) 2-butene
- (b) 1-butene
- (c) 2-pentene
- (d) 2-hexene
- 167. Reaction of HBr with propene in the presence of peroxide gives [CBSE PMT 2004]
 - (a) Allyl bromide
- (b) *n*-propyl bromide
- (c) Isopropyl bromide
- (d) 3-bromo propane
- **168.** Which of the following react with $KMnO_4$ but does not react with AgNO 3? [BCECE 2005]







- (a) C_2H_6
- (b) CH_4
- (c) C_2H_4
- (d) C_2H_2
- **169.** 3-Phenylpropene on reaction with *HBr* gives (as a major product) **[AIIMS 2005]**
 - (a) $C_6H_5CH_2CH(Br)CH_3$
 - (b) $C_6H_5CH(Br)CH_2CH_3$
 - (c) $C_6H_5CH_2CH_2CH_2Br$
 - (d) $C_6H_5CH(Br)CH = CH_2$
- 170. Reaction of one molecule of HBr with one molecule of 1,3-butadiene at $40^{\circ}C$ gives predominantly [AIEEE 2005]
 - (a) 3-bromobutene under kinetically controlled conditions
 - (b) 1-bromo-2-butene under thermodyanamically controlled conditions
 - (c) 3-bromobutene under thermodynamically controlled conditions
 - (d) 1-bromo-2-butene under kinetically controlled conditions
- 171. The only alcohol that can be prepared by the indirect hydration of alkene is [AFMC 2005]
 - (a) Ethyl alcohol
- (b) Propyl alcohol
- (c) Isobutyl alcohol
- (d) Methyl alcohol

 CH_3

- 172. The reaction of HBr with $CH_3 C = CH_2$ in the presence of peroxide will give [BHU 2005]
 - (a) CH_3CBrCH_3
- (b) CH₃CH₂CH₂CH₂Br

 CH_3 CH_3

- CH_3
- (c) CH₃CHCH₂Br
- (d) CH₃CH₂CHCH₃
- 173. A gas decolourised by KMnO_4 solution but gives no precipitate with ammoniacal cuprous chloride is

[KCET 2005]

- (a) Ethane
- (b) Methane
- (c) Ethene
- (d) Acetylene
- **174.** Cyclohexene on reaction with OsO_4 followed by reaction with $NaHSO_3$ gives [Orissa JEE 2005]
 - (a) cis-diol
- (b) trans-diol
- (c) epoxy
- (d) alcohol

Alkyne

1. Which of the following gases is used for welding

[CPMT 1996]

- (a) Methane
- (b) Ethane
- (c) Acetylene
- (d) Ethene
- 2. A metallic carbide on treatment with water gives a colourless gas which burns readily in air and which gives a precipitate with ammoniacal silver nitrate solution. Gas evolved is

[NCERT 1975; CPMT 1977; MP PET 2002]

- (a) Methane
- (b) Ethane
- (c) Acetylene
- (d) Ethylene
- 3. 1-butyne reacts with cold alkaline KMnO_4 to produce

[AIIMS 1997]

- (a) CH_3CH_2COOH
- (b) $CH_3CH_2CH_2COOH$
- (c) $CH_3CH_2COOH + CO_2$
- (d) $CH_3CH_2COOH + HCOOH$
- Identify the product D in the following series of reaction

$$CH_3COOH \xrightarrow{LiAlH_4} A \xrightarrow{H^+} B \xrightarrow{Br_2} C \xrightarrow{alc.} D$$

[CBSE PMT 1998]

- (a) Methane
- (b) Alcohol
- (c) Acetylene
- (d) Benzaldehyde
- 5. The correct order towards bond length is

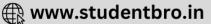
[RPMT 1997]

- (a) $C C < C = C < C \equiv C$ (b) $C \equiv C < C = C < C C$
- (c) $C = C < C \equiv C < C C$ (d) $C = C < C C < C \equiv C$
- **6.** In the molecule $CH \equiv C CH = CH_2$, the hybridisation of C C bond is
 - (a) $sp^2 sp$
- (b) $sp^3 sp^3$
- (c) $sp^2 sp^2$
- (d) $sp^3 sp$
- 7. The product formed when acetylene is passed through red hot tube is [BHU 1989; RPMT 2003]
 - (a) Benzene
- (b) Cyclohexane
- (c) Neoprene
- (d) Ethane
- **8.** Acetylenic hydrogens are acidic because

[CBSE PMT 1989; Pb. PMT 1999]

- (a) Sigma electron density of C-H bond in acetylene is nearer to carbon, which has 50% s-character
- (b) Acetylene has only one hydrogen on each carbon
 - (c) Acetylene contains least number of hydrogens among the possible hydrocarbons having two carbons
 - (d) Acetylene belongs to the class of alkynes with molecular formula $C_n H_{2n-2}$
- Which is the most suitable reagent among the following to distinguish compound (iii) from rest of the compounds
 - (i) $CH_3 C \equiv C CH_3$
 - (ii) $CH_3 CH_2 CH_2 CH_3$





$$CH_3 - CH_2 - C \equiv CH$$

(iv)
$$CH_3 - CH = CH_2$$

[CBSE PMT 1989]

- (a) Bromine in carbon tetrachloride
- (b) Bromine in acetic acid
- (c) Alkaline KMnO₄
- (d) Ammoniacal silver nitrate reagent
- A hydrocarbon of formula C_6H_{10} absorbs only one molecule of H_2 upon catalytic hydrogenation. Upon ozonolysis, the hydrocarbon yields

$$H \cap H \cap H \cap H \cap C = C - CH_2 - CH_2$$

The hydrocarbon is

[MP PMT 1986]

- (a) Cyclohexane
- (b) Benzene
- (c) Cyclohexene
- (d) Cyclobutane
- Poisonous gas 'Lewissite' is obtained by the 11. reaction of

[MP PMT 2003]

- (a) $CH \equiv CH$ and $AsCl_3$
- (b) $CH_2 = CH_2$ and $AsCl_3$
- (c) $CH \equiv CH$ and S_2Cl_2
- (d) $CH_2 = CH_2$ and NOCl
- Products of following reaction 12. $CH_3C \equiv C CH_2CH_3 - \frac{(2) Hydrolysis}{(2) Hydrolysis}$ \rightarrow are

[CBSE PMT 2005]

- (a) $CH_3CHO + CH_3CH_2CHO$
- (b) $CH_3COOH + CH_3CH_2CHO$
- (c) $CH_3COOH + HOOCCH_3CH_3$
- (d) $CH_3COOH + CO_2$
- By coaltar distillation, which is not obtained 13.

[SCRA 1990; MP PMT 1986]

- (a) Light oil
- (b) Middle oil
- (c) Heavy oil
- (d) Mobil oil
- Hydrocarbon containing following bond is most 14. reactive

[AIIMS 1987]

- (a) $C \equiv C$
- (b) C = C
- (c) C C
- (d) All of these
- The shapes of methane, ethene and ethyne 15. molecules are, respectively
 - (a) Tetrahedral, planar and linear
 - (b) Tetrahedral, linear and planar
 - (c) Pyramidal, planar and linear
 - (d) Tetrahedral, pyramidal and planar
- 16. To synthesize the unsymmetrical alkyne $CH_3 - C \equiv C - CH_2 - CH_3$ the reagents needed would be

- iodoethane, iodomethane (a) Ethene, and potassium hydroxide
- (b) Acetaldehyde, 1-bromopropane and conc. H_2SO_4
 - (c) 1, 2-dichloroethane, 1-propanol and alcoholic potassium hydroxide
 - (d) Ethyne, iodomethane, iodoethane and sodamide
- When propyne is treated with dilute H_2SO_4 and 17. HgSO₄, the major product is [Kurukshetra CEE 2002]
 - (a) Propanal
 - (b) Propanol
 - (c) Propyl hydrogen sulphate
 - (d) Propanone
- 18. Which of the following will be the final product when C_2H_2 reacts with HCl

[DPMT 1984; AFMC 1982; Bihar MEE 1982]

- CH(a) || CHCl
- CHCl
- (d) None of these
- What is the end product of the following sequences operations $\xrightarrow{\text{dil. } H_2SO_4} B -$

[CPMT 1978; MP PMT 1996]

- (a) Methyl alcohol
- (b) Acetaldehyde
- (c) C_2H_5OH
- (d) C_2H_4
- **20.** $R CH_2 CCl_2 R \xrightarrow{\text{Reagent}} R C \equiv C R$

The reagent is [CBSE PMT 1989; MP PET 1995]

- (a) Na
- (b) HCl and H_2O

- (c) KOH in C_2H_5OH
- (d) Zn
- Acetylene can be prepared from
 - (a) Potassium fumarate (b) Calcium carbide

 - (c) Ethylene bromide
- (d) All of these
- Acetylene is obtained by the electrolysis of [BHU 1986]
 - (a) Sodium succinate
- (b) Potassium fumarate
- (c) Both (a) and (b)
- (d) None of these
- The compound C_3H_4 has a triple bond, which is indicated by its reaction with
 - (a) Bromine water
- (b) Bayer's reagent
- (c) Fehling solution nitrate
- (d) Ammonical silver
- $CH \equiv CH \xrightarrow{H_2O/Hg^{2+}} X \xrightarrow{LiAlH_4} Y \xrightarrow{P_4/Br_2} Z$ Here Z

[JIPMER 2002]

[CPMT 1988]

- (a) Ethylene bromide
- (b) Ethanol
- (c) Ethyl bromide
- (d) Ethylidene bromide
- $CH \equiv CH \xrightarrow{Ni(CN)_2} X$. Here X in the reaction 25.







[JIPMER 2002]

- (a) Benzene
- (b) Ethane
- (c) Cycloctatetraene
- (d) Cyclohexane
- A salt producing hydrocarbon among these compounds is

[KCET (Engg.) 2002]

- (a) Ethane
- (b) Methane
- (c) Ethene
- (d) Ethyne
- An unknown compound A has a molecular formula C_4H_6 . When A is treated with an excess of Br_2 a new substance B with formula $C_4H_6Br_4$ is formed. A forms a white precipitate with ammoniacal silver nitrate solution. A may be

[MP PET/PMT 1998]

- (a) Butyne-1
- (b) Butyne-2
- (c) Butene-1
- (d) Butene-2
- 28. Which of the following reacts with sodium with the elimination of hydrogen
 - (a) CH_A
- (b) C_2H_6
- (c) C_2H_4
- (d) C_2H_2
- 29. Acetylene gives

[CPMT 1985]

- (a) White precipitate with AgNO 3 and red precipitate with Cu_2Cl_2
- (b) White precipitate with Cu_2Cl_2 and red precipitate with AgNO 3
- (c) White precipitate with both the reagents
- (d) Red precipitate with both the reagents
- The bond length between sp^3 hybridised carbon atom and other carbon atom is minimum in

[CBSE PMT 1996; Pb. PMT 1999]

- (a) Propane
- (b) Butane
- (c) Propene
- (d) Propyne
- 31. The C - H bond length is minimum in the bond formed by
 - (a) *sp s* overlapping (as in alkynes)
 - (b) $sp^2 s$ overlapping (as in alkenes)
 - (c) $sp^3 s$ overlapping (as in alkanes)
 - (d) None of these
- **32.** Which of the *C C* bond is strongest
 - (a) Formed by $sp^3 sp^3$ hybridised carbon atoms (as in alkanes)
 - (b) Formed by $sp^2 sp^2$ hybridised carbon atoms (as in alkenes)
 - (c) Formed by sp sp hybridised carbon atoms (as in alkynes)
 - (d) All are equal
- Which of the following pairs has the same bond
- (a) Ethane and ethylene (b) Ethylene and acetylene
 - (c) Ethylene and benzene (d)Acetylene and benzene

The product(s) obtained via oxymercuration 34. $(HgSO_4 + H_2SO_4)$ of 1-butyne would be[IIT-JEE 1999]

- (a) $CH_3 CH_2 C CH_3$
- (b) $CH_3 CH_2 CH_2 CHO$
- (c) $CH_3 CH_2 CHO + HCHO$
- (d) $CH_3CH_2COOH + HCOOH$
- **35.** A compound is treated with NaNH, to give sodium salt. Identify the compound [AFMC 1998]
 - (a) C_2H_2
- (b) $C_6 H_6$
- (c) C_2H_6
- (d) C_2H_4
- A gas decolourises bromine in CCl4 and forms a precipitate with ammoniacal silver nitrate. The [EAMCET 1998]
 - (a) C_2H_2
- (b) C_2H_4

[BHU 1983]

- (d) CH_{Λ}
- Among the following compounds which have more than one type of hybridisation for carbon atom
 - (i) $CH_3CH_2CH_2CH_3$
 - (ii) $CH_3 CH = CH CH_3$
 - (iii)
 - $CH_2 = CH C \equiv CH$
 - (iv) $H C \equiv C H$
- (b) (ii)
- (a) (ii) and (iii) (c) (iii) and (iv)
- (d) (iv)
- **38.** The homologue of ethyne is
- [EAMCET 1998]

[EAMCET 1998]

- (a) C_2H_4
- (b) C_2H_6
- (c) C_3H_8
- (d) C_3H_4
- When acetylene reacts with HCl in the presence of $HgCl_2$, the product is [MNR 1985; MP PET 1996; UPSEAT 200
 - (a) Methyl chloride
- (b) Dichloroethane
- (c) Vinyl chloride
- (d) Ethylidine chloride
- When propyne reacts with aqueous H_2SO_4 in the presence of $HgSO_4$, the major product is

[IIT-JEE 1983; AFMC 1991; KCET 1993]

- (a) Propanal
- (b) Propyl
- hydrogen

- sulphate
 - (c) Acetone
- (d) Propanol
- Propyne on polymerisation gives [CPMT 1999, 2002]
 - (a) Mesitylene
- (b) Benzene
- (c) Ethyl benzene
- (d) Propyl benzene
- When treated with ammoniacal cuprous chloride, which one among the following forms copper derivative

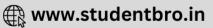
[CBSE PMT 1989; MP PMT 1993]

- (a) C_2H_6
- (b) C_2H_4
- (c) C_2H_2
- (d) C_6H_6





43.	Which of the following catalyst is used in polymerisation of $CH \equiv CH$ to C_6H_6 [CPMT 1				(c) Metal silver (d) Silver mirror					
		0 0	[CPMT 1999]	53.		can be prepared by the				
	(a) $AlCl_3$	(b) $HgSO_4$			reaction of <i>HCl</i> and	(h)				
	(c) $NbCl_3$	(d) HCl			(a) C_2H_4	(b) C_2H_2				
44.	KMnO ₄ will oxidise ace	tylene to	[CPMT 1999]		(c) C_2H_5	(d) All of these				
	(a) Ethylene glycol(c) Oxalic acid	(b) Ethyl ald (d) Acetic ad		54.		g order of reagent is chosen ene from C_2H_2 [RPET 2000]				
45.	Ethyne on reaction w	ith dil. H_2SO	$_4$ and $Hg(II)$		(a) $CuCl/NH_4Cl$ and $H_2/Pd(BaSO_4)$					
	gives				(b) $NH_4Cl/CuCl$ and I	$H_2/Pd(BaSO_4)$				
	(a) Ethanol				(c) $H_2/Pd(BaSO_4)$ and					
	(b) Ethanal				(d) $H_2/Pd(BaSO_4)$ and $NH_4Cl/CuCl$					
	(c) Methoxymethane	shata								
46	(d) Ethyl hydrogen sulp		dictinguich	55.	= -	r of[RPET 1999; Bihar MEE 19				
46.	Which of the following ethylene and acetylene	ng is used it [MP PET 2000:	KCET 2000: IIPN	/JER 20	(a) Methane	(b) Ethane				
		CPMT 1977	; NCERT 1973]			(d) Ethyne				
	(a) Alkaline KMnO ₄	3,,	, 5,01	56.	СН	acid in presence of Hg^{2+} to				
	(b) Bromine water				give	[D1111 acc-1				
	(c) Ammoniacal cuprou	ıs chloride			CH_3	[BHU 2005] CH(CH ₃ COO) ₂				
	(d) Conc. H_2SO_4				(a) \downarrow $CH(CH_3COO)_2$	(b) $+$ $CH(CH_3COO)_2$ $CH(CH_3COO)_2$				
47.	The distinguishing test	_	_		3	$CH(CH_3COO)_2$				
	acidic hydrogen is	[.	JIPMER 2000]		(c) $\stackrel{CH_3}{\downarrow}$	(d) None of these				
	(a) $Ag(NH_3)_2^+$	(b) Br_2 in CC	Cl_4		$CH_2(CH_3COO)$					
_	·	(d) AlCl ₃		57•	electric discharge thr	ed industrially by passing ough graphite electrodes in				
48.	If acetylene is passed through an electric arc in the atmosphere of nitrogen, the compound formed is				the atmosphere of	[CPMT 1985]				
					(a) Air	(b) N_2				
	[1		[RPMT 1999]		(c) H_2	(d) CO_2				
	(a) HCN	(b) Pyrrole		- 8	-	assed into dilute sulphuric				
	(c) Pyrazole	(d) Pyridine		50.	-	ions, the product formed is				
49.	Ozonolysis of acetylene	gives	[RPMT 1999]		•	e 1995; BHU 1998; KCET 1999;				
	(a) Glycol	(b) Glyoxal,	formic acid			; DCE 1999; DPMT 1999, 2002;				
	(c) Formaldehyde	(d) None				, 82, 83, 90; MP PMT 1994, 97;				
50.	The bond length between	een the hybri	dised carbon			; AIIMS 2002; CBSE PMT 1999;				
	atom and other carbon	atom is minim	ium in [Pb. PMT	2000]	(a) Acetone	ET (Med.) 1999, JIPMER 1999] (b) Acetic acid				
	(a) Butane	(b) Propyne			(c) Acetaldehyde	(d) Formaldehyde				
	(c) Propene	(d) Propane		59.	Which of the following	· · · · · · · · · · · · · · · · · · ·				
51.	The reaction of propen		-			; CPMT 1986; Bihar MEE 1997;				
	the addition of [IIT-JEE (Screening) 2001]					RPET 1999; AFMC 1999]				
	 (a) H⁺ in the first step (b) Cl⁺ in the first step 				(a) Ethyne	(b) Ethene				
					(c) Ethane	(d) Benzene				
	(c) OH^- in the first step			60.	Ayrenes on oxidation v	with acidic <i>KMnO</i> ₄ gives				
	(d) Cl^+ and OH^- in a single step				(a) Terphthalic acid	[JIPMER 2000] (b) Phthalic acid				
52.	Acetylene reacts with a	-	O_3 forming		(c) Isophthalic acid	(d) All of these				
	[MH CET 1999; C	=	2		(s) === promise word	(, 32 0000				
	(a) Silver acetylene	(b) Silver ac								
		(-,								



The structure of the product(Z) in the reactions 61. given

$$HC \equiv CH \xrightarrow{NaNH_2, CH_3COCH_3} X \xrightarrow{Hg^{2+}, H_3O^+} Z$$
 is

[Roorkee 2000]

- (a) $CH_3 CH_2 CH_2 CH_2OH$
- (b) $CH_3 C CH CH_2OH$ O
- (c) $CH_3 C C CH_3$ O OH CH_3
- (d) $CH_3 CHOH CH CHO$
- Carbon-carbon bond length is minimum in 62.

[CBSE PMT 1988, 91; MNR 1984; CPMT 1989; RPMT 1997; Pb. PMT 2001]

- (a) Ethane
- (b) Ethene
- (c) Ethyne
- (d) Benzene
- (e) Ethanol
- Triple bond of ethyne is made of or Cylindrical 63. shape of an alkyne is due to

[EAMCET 1978; NCERT 1979; CBSE PMT 1997; Manipal MEE 1995; Bihar MEE 1996]

- (a) Three σ bonds
- (b) Three π bonds
- (c) Two σ and one π bond
- (d) Two π and one σ bond
- An organic compound has a triple bond and not double bond. It can be tested by [MP PMT 2000, 03]
 - (a) Bromine water
 - (b) Bayer's reagent
 - (c) Fehling solution
 - (d) Ammonical silver nitrate
- Which of these will not react with acetylene 65.

[AIEEE 2002; DCE 2002]

- (a) NaOH
- (b) Ammonical AgNO 3
- (c) Na
- (d) HCl
- 66. What is the product when acetylene reacts with hypochlorous acid [RPMT 2002; AIEEE 2002]
 - (a) CH₃COCl
- (b) ClCH2CHO
- (c) Cl₂CHCHO
- (d) CICHCOOH
- The alkene C_6H_{10} producing $OHC (CH_2)_4 CHO$ on ozonolysis is [Roorkee 1999]
 - (a) Hexene-1
- (b) Hexene-3
- (c) Cyclohexene
- (d) 1-
- methylcyclohexene-1

- The number of moles of proton which can be easily given by butyne-1(1 mole) is [MP PMT 2000]
 - (a) 1

(b) 2

(c) 3

- (d) 6
- Which will undergo reaction with ammoniacal

[DPMT 1996]

(a)
$$\frac{CH_3}{CH_3}$$
 $CH - CH_2 - CH = CH - CH_3$

- (b) $CH_3 CH = CH C \equiv CH$
- (c) $CH_3 CH_2 CH = CH CH_2 CH_3$
- (d) $CH_2 = CH CH_2 CH_3$
- (e) None
- 70. Acetylene gas when passed through the 20% H_2SO_4 at $80^{\circ}C$ gives acetaldehyde. The catalyst required for this conversion is
 - (a) Anhydrous AlCl₃
- (b) $HgSO_4$
- (c) Pd
- (d) Pt
- Which of the following reactions will yield 2, 2-71. dibromopropane [MNR 1993; UPSEAT 2001]
 - (a) $HC \equiv CH + 2HBr \rightarrow$
 - (b) $CH_3C \equiv CH + 2HBr \rightarrow$
 - (c) $CH_3CH = CH_2 + HBr \rightarrow$
 - (d) $CH_3CH = CHBr + HBr \rightarrow$
- Which of the following does not give white 72. precipitate with ammoniacal AgNO₃
 - (a) $CH \equiv CH$
- (b) $CH_3 C \equiv CH$
- (c) $CH_3 C \equiv C CH_3$ (d) $CH_2 C \equiv CH$
- $||| \xrightarrow{O_3 / NaOH} X \xrightarrow{Z_n / CH_3COOH} Y 'Y' \text{ is[AIIMS 1988]}$
 - (a) $\begin{vmatrix} CH_2OH \\ CH_2OH \end{vmatrix}$
- (b) CH₃CH₂OH

- (c) CH₃COOH
- (d) CH_3OH
- Which is represented by the formula $C_n H_{2n-2}$ 74.

[CPMT 1975, 76; EAMCET 1979; MP PET 2003

- (a) Alkane
- (b) Alkyne
- (c) Alkene
- (d) None of these
- What is the major product of the following reaction $CH_3C \equiv C - CH_2 - CH_3 \xrightarrow{1 \text{ mole of } Cl_2}$

[Kerala (Med.) 2003]

(a)
$$CI > C = C < CI > CH_2CH_3$$

(b)
$$CH_3 - CH_2 - CH_2 - CH_2 + CH_3$$



- (c) $Cl > C = C < \frac{CH_2CH_3}{Cl}$
- (d) $CH_3 \overset{\mid}{C} \overset{\mid}{C} CH_2CH_3$
- A compound C_5H_8 which give white ppt. with ammonical AgNO 3. A give (CH 3)2 CHCOOH with hot alcoholic *KOH* then compound is **[RPMT 2002]**
 - (a) $CH_3CH_2 CH_2 CH = CH_2$
 - (b) $CH_3 CH_2 C \equiv CH$
 - (c) $(CH_3)_2 CH C \equiv CH$
 - (d) $CH_2 = CH CH_2 CH = CH_2$
- 77. 1, 2-dibromoethane when heated with alcoholic potash gives [Kerala PMT 2004]
 - (a) Ethane
- (b) Acetylene
- (c) Ethylene
- (d) Methane
- (e) None of these
- 78. Which of the following is not a member of homologous series [RPMT 2002]
 - (a) Ethene
- (b) 1-butene
- (c) 2-butene
- (d) 2-butyne
- 79. The compound formed as a result of potassium permanganate oxidation of ethylbenzene is

[MP PET/PMT 1998]

- (a) Benzoic acid
- (b) Benzyl alcohol
- (c) Benzophenone
- (d) Acetophenone
- What is the product when 2-butyne is treated with liquid NH_3 in presence of lithium[Orissa JEE 2003]
 - (a) *n*-butane
- (b) cis-2-butene
- (c) trans-2-butene
- (d) 1-butene
- Distinction in pentene-1 and pentyne-1 is done by [CPMT 1996]
 - (a) $[Ag(NH_3)_2]^+$
- (b) Conc. H_2SO_4
- (c) HCl
- (d) Br_2
- 82. A mixture of ethane, ethene and ethyne is passed through ammoniacal $AgNO_3$ solution. The gases which remain unreacted are
 - (a) Ethane and ethene
- (b) Ethane and ethyne
- (c) Ethene and ethyne
- (d) Ethane only
- 83. In its reaction with silver nitrate acetylene shows
 - [MP PET 1999]
 - (a) Oxidising property
- (b) Reducing property
- (c) Basic property
- (d) Acidic property
- **84.** Simplest alkyne is represented by [CPMT 1974]
 - (a) CH
- (b) CH₂
- (c) C_2H_2
- (d) C_2H_4
- **85.** Which of the following bonds is most acidic
 - (a) = C H
- (b) -C-H
- (c) $\equiv C H$
- (d) All are equally acidic

- 86. The hybridisation in methane, ethene and Ethyne [CPMT 2003] respectively is
 - (a) sp^3 , sp^2 and sp
- (b) sp^{3} , sp, sp^{2}
- (c) sp^2 , sp^3 and sp
- (d) sp^3 , sp^2 , sp
- Number of acidic hydrogen atoms in butyne-1 are **[MP PET 1986]**
 - (a) 2

(b) 3

(c) 1

- (d) 4
- 88. Which of the following shows linear structure[CPMT 2003]
 - (a) Ethane
- (b) Ethene
- (c) Acetylene
- (d) CCl_{\perp}
- 89. Calcium carbide on reacting with water gives

[CBSE PMT 1991; MP PMT 1993, 94; RPMT 2002; J & K 20051

- (a) Methane
- (b) Ethane
- (c) Ethene (d) Acetylene
- **90.** Addition of *HCN* to ethyne in presence of Ba(CN), as catalyst gives [AFMC 1991]
 - (a) 1, 1-dicyano ethane
- (b) Ethyl cyanide
- (c) Vinyl cyanide
- (d) Divinyl cyanide
- Which compound will react with an aqueous 91. solution of $Ag(NH_3)^+_2OH^-_3$
 - (a) $CH_2 = CH_2$
- (b) $CH_3 CH_3$
- (c) $CH_3CH_2C \equiv CH$
- (d) $CH_3 C \equiv C CH_3$
- **92.** Which of the following give H_2 gas with Na[RPMT 2002]
 - (a) CH_4
- (b) C_2H_6
- (c) C_2H_4
- (d) C_2H_2
- $CH_3 C \equiv CH \xrightarrow{O_3}$ Product .Product above Zn/H_2O_2

reaction is

[RPMT 2003]

- (a) CH₃COOH
- (b) HCOOH
- (c) Both (a) and (b)
- (d) $CH_3CHO + HCHO$
- The number of π bonds in the product formed by passing acetylene through dilute sulphuric acid containing mercuric sulphate is [EAMCET 1997]
 - (a) Zero [CPMT 1990] (c) Two
- (b) One
- (d) Three
- Which of the following is weakly acidic
 - (a) $CH_2 = CH_2$
- (b) C_6H_6
- (c) $CH_3 C \equiv CH$
- (d) $CH_3 C \equiv C CH_3$
- Which of the following reactions is shown by alkynes

[AMU 1984; RPMT 2000]

- (a) Addition
- (b) Substitution
- (c) Polymerization
- (d) All of these
- 97. Shortest C-C bond length is present in [BVP 2004]
 - (a) $CH_3 CH_2 CH_3$
- (b) $CH_3CH_2CH_2CH_3$







- (c) $CH_2 = CH CH = CH_2$ (d) $CH \equiv C C \equiv CH$
- 98. Acetylene can be obtained by the reaction[MH CET 2004]
 - (a) $HCOOK \xrightarrow{\text{electroly sis}}$
 - (b) $CHI_3 + 6Ag + CHI_3 \xrightarrow{\Delta}$
 - (c) $CH_3CH_2OH \xrightarrow{\text{Conc. } H_2SO_4}$
 - (d) $Be_2C + H_2O \rightarrow$
- **99.** Which of the following used for the conversion of 2-hexyne into trans-2-hexane[IIT JEE (Screening) 2004]
 - (a) $H_2/Pd/BaSO_4$
- (b) H_2 , PtO_2
- (c) NaBH ₄
- (d) $Li NH_3 / C_2H_5OH$
- **100.** In which of the following, the bond length between hybridized carbon atom and other carbon atom is minimum

[MH CET 2003]

- (a) Propyne
- (b) Propene
- (c) Butane
- (d) Propane
- **101.** What happens when a mixture of acetylene an hydrogen is passed over heated Lindlar's catalyst

[Kerala PMT 2004; AIIMS 1987]

- (a) Ethane and water are formed
- (b) Ethylene is formed
- (c) Acetylene and ethane are formed
- (d) None of these
- **102.** In acetylene molecule, the two carbon atoms are linked by

[KCET 2004]

- (a) One sigma bond and two pi bonds
- (b) Two sigma bonds and one pi bond
- (c) Three sigma bonds
- (d) Three pi bonds
- **103.** Which reacts with ammoniacal *AgNO* 3

[Orissa JEE 2005]

- (a) Propyne
- (b) 2-butyne
- (c) 1,3-butadiene
- (d) Pentene
- **104.** $CH \equiv CH \xrightarrow{HgSO_4} \xrightarrow{CH_3MgBr} \xrightarrow{P/Br_2} \xrightarrow{P/Br_2}$

[DPMT 2005]

- (a) $CH_3CH(Br)CH_3$
- (b) $CH_3CH_2CH_2Br$
- (c) $CH_2 = CH Br$
- (d) $BrCH = CH CH_3$
- **105.** Carbide, which react with water to give propyne is

[Kerala CET 2005]

- (a) CaC_2
- (b) SiC
- (c) Mg_2C_3
- (d) Al_4C_3
- (e) Be_2C

Aromatic hydrocarbon

The function of anhydrous $AlCl_3$ in the Friedel-Craft's reaction is to

[MNR 1986, 1995; Roorkee 1999; BHU 2001; CPMT 2002; MPPET 2001]

- (a) Absorb water
- (b) Absorb HCl
- (c) To produce electrophile (d)To produce nucleophile
- Benzene reacts with CH_3COCl in the presence of
 - $AlCl_3$ to give

[DPMT 1983; CBSE PMT 1991]

- (a) C_6H_5Cl
- (b) C_6H_5COCl
- (c) $C_6H_5CH_3$
- (d) $C_6H_5COCH_3$
- Acylation process is preferred than direct alkylation because (by the Friedel-Craft's reaction)
 - (a) In alkylation, a poisonous gas is evolved
 - (b) In alkylation, large amount of heat is evolved
 - (c) In alkylation, polyalkylated product is formed
 - (d) Alkylation is very costly
- 4. Benzene cannot undergo
 - (a) Substitution
- (b) Addition
- (c) Elimination
- (d) Oxidation
- 5. Coaltar is main source of
- [DPMT 1984]
- (a) Aromatic compounds
- (b) Aliphatic compounds
- (c) Cycloalkanes
- (d) Heterocyclic compounds
- **6.** Which of the following is not formed by the ozonolysis of *o*-xylene
 - (a) Glyoxal
- (b) Ethyl glyoxal
- (c) Dimethyl glyoxal
- (d) Methyl glyoxal
- 7. The number of σ and π bonds in a molecule of benzene is

[MP PMT/PET 1988; BHU 1995; CPMT 1997]

- (a) 6σ and 9π
- (b) 9σ and 3π
- (c) 12σ and 3π
- (d) 6σ and 6π
- **8.** The ratio of σ and π bonds in benzene is

[CPMT 1991; BHU 1995]

- (a) 2
 - 5 ((
- (c) 6
- (b) 4 (d) 8
- Carbon atoms in benzene molecule is inclined at an angle of [BHU 1985]

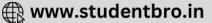
[Bite 1905]

- (a) 120°
- **(b)** 180 °
- (c) 109°28′
- (d) 60°
- 10. When benzene is treated with excess of Cl_2 in the presence of I_2 , the end product is
 - (a) Monochlorobenzene (b) Trichlorobenzene
 - (c) Hexachlorobenzene (d) Benzene

hexachloride

 Chemical name of the insecticide gammexene is [CPMT 1981; MP PET 1995; MP PMT 1996;





				Hydrocarbon 1131
	(a) DDT	CBSE PMT 1999; MP PET 1999] (b) Benzene	20.	Nitration of benzene by nitric acid and sulphuric acid is
hexa	chloride			[MNR 1989; CPMT 1990; BCECE 2005]
	(c) Chloral	(d) Hexachloroethane		(a) Electrophilic substitution (b)
12.		ned from benzene when it		(c) Nucleophilic substitution(d)Free radical substit
	reacts with	light (in the change of a	21.	Necessary conditions for halogenation are[CPMT 19
_	=	llight (in the absence of a		(a) Cold and dark
catal				(b) Presence of halogen carrier
	-	llight (in the absence of a		(c) Both (a) and (b)
atal	=			(d) None
	(c) CH_3Cl in the prese	ence of anhydrous AlCl ₃	22.	$C_6H_6 + CH_3Cl \xrightarrow{\text{anhy drous}} C_6H_5CH_3 + HCl$
	(d) $COCl_2$ in the prese	ence of anhydrous AlCl ₃		
3.	Point out the wrong s	statement in relation to the		is an example of [NCERT 1979; CPMT 1974, 85, 90; Bihar CEE 1995; BHU 1979, 2001; MP PET 1995;
	structure of benzene			MP PMT 1995; KCET 1993; EAMCET 1998; AIIMS 1998;
		nonosubstitution product		CBSE PMT 2000; AFMC 2000; JIPMER 2000]
		ance in benzene is uniformly		(a) Friedel-Craft's reaction (b) Kolbe's synthesis
	1.397 Å			(c) Wurtz reaction (d) Grignard reaction
		e hybrid of a number of	23.	The reaction of benzene with chlorine in the
ano	nical forms	lind male and an authority		presence of iron gives [MP PET 1993]
_		lised π - molecular orbitals		(a) Benzene hexachloride (b) Chlorobenzene
4.	•	sents an example of Friedel-		(c) Benzyl chloride (d) Benzoyl chloride
	Craft's reaction	[MNR 1993; CPMT 1996]	24.	Benzene was discovered by [NCERT 1981]
	(a) $C_6H_6 + C_2H_5Cl - Al$			(a) Ramsay (b) Dalton
	(b) $C_2H_5OH + HCl - Z_{10}$	$C_2H_5Cl + H_2O$		(c) Faraday (d) Priestley
	(c) $C_c H_c Cl + CH_c COCl$	$\xrightarrow{AlCl_3} C_6H_5COCH_3 + Cl_2$	25.	The correct structure of benzene was proposed by
		0 0 0 2		[CPMT 1972]
	(d) $C_2H_5Br + Mg - Ether$	- *		(a) Faraday (b) Davy
5.		arbonium ion among the		(c) Kekule (d) Wohler
	following is	[VVDVCTD ATVCC]	26.	1 1 5
	+	[JIPMER 2002; AFMC 2002]		[CPMT 1982, 83, 89]
	(a) $C_6H_5^+CHC_6H_5$	(b) $C_6H_5CH_2$		(a) Dewar (b) Ladenberg
	(c) $CH_3^+CH_2$		D	(c) Kekule (d) Armstrong and
_		* * = =	Baey	
16.	The reaction of toluen	e with chlorine in presence		The bond order of individual carbon-carbon bonds [IIT-JEE 1981; MP PET 2000]
		predominantly [IIT-JEE 1986;		(a) One (b) Two
	=	(b) <i>m</i> -chlorotoluene		(c) Between one and two(d) One and two,
hlor	(c) Benzyl chloride otoluenes	(d) <i>o</i> - and <i>p</i> -	alter	rnately
11101 7 •		when toluene is heated in		Six carbon atoms of benzene are of
/•		absence of halogen carrier is	_••	(a) One type (b) Two types
	(a) Benzotrichloride			(c) Three types (d) Six types
	(c) Chlorobenzene	(b) Gammexene(d) None of these	29.	On heating a mixture of sodium benzoate and
8.		or electrophilic species in	_5.	sodalime, the following is obtained
σ.	<u>o</u>	is or In the nitration of		[CPMT 1990; AIIMS 1996; MP PET 1999; AFMC 1999]
		trated HNO_3 and H_2SO_4 the		(a) Toluene (b) Phenol
	attack on ring is made			(c) Benzene (d) Benzoic acid
	_	PET 1996, 2000; Pb. PMT 1998;	30.	Benzene on treatment with a mixture of conc.
		[U 2001; BVP 2004; DCE 2003]	-	HNO_3 and conc. H_2SO_4 at $100 ^{o}C$ gives
	(a) NO_2^-	(b) NO_{2}^{+}		(a) Nitrobenzene (b) <i>m</i> -dinitrobenzene
				(c) p-dinitrobenzene (d) o-dinitrobenzene
	(c) NO_3^-	(d) NO_2	31.	-
9.	-	he following reactions takes place when		What is the end product which is obtained on the nitration of toluene [MP PMT/PET 1988]
	a mixture of concentrated HNO_3 and H_2SO_4			- ,
	reacts on benzene at 3	50 K	[0	(a) o-nitrotoluene (b) p-nitrotoluene (c) 2, 4-dinitrotoluene (d) 2, 4, 6-
	(a) Sulphonation	(b) Nitration		(c) 2, 4-dinitrotoluene (d) 2, 4, 6-itrotoluene
	(c) Hydrogenation	(d) Dehydration	¢1 1111	ici ocoraciic





	1132 Hydrocarbor			
32.	Which of the following	processes is reversible		(c) Toluene (d) Phenol
	(a) Halogenation(c) Nitration	(b) Sulphonation(d) None	45.	Select the true statement about benzene from amongst the following [CBSE PMT 1992]
33.	The attacking (ele	ctrophilic) species in		(a) Because of unsaturation benzene easily
	=	e is[RPMT 1997; CPMT 1999, 2 0	002]	undergoes addition reactions
	(a) SO_2	(b) SO_3		(b) There are two types of <i>C</i> - <i>C</i> bonds in benzene
	(c) SO_4^{2-}	(d) HSO_3^-		molecule (c) There is a cyclic delocalisation of π electrons in
34.	Which xylene is most ea	sily sulphonated		benzene
J 1	(a) Ortho	(b) Para		(d) Monosubstitution of benzene group gives
	(c) Meta	(d) All at the same rate		three isomeric substances
35.	Toluene on oxidation	with dilute HNO_3 and	46.	Anhydrous AlCl ₃ is used in the Friedel-Craft's
	alkaline KMnO ₄ gives	[DPMT 1981]		reaction because it is [CBSE PMT 1991]
	(a) Benzaldehyde	(b) Phenol		(a) Electron rich
	(c) Nitrotoluene	(d) Benzoic acid		(b) Soluble in ether
36.	• •	with air when passed over		(c) Insoluble to chloride and aluminium ions
	V_2O_5 catalyst at 775 K g	<u>-</u>		(d) Electron deficient
	[AFMC 1991	; CPMT 2001; MP PMT 2003]	47.	(i) Chlorobenzene and (ii) benzene hexachloride
	(a) Glyoxal	(b) Oxalic acid		are obtained from benzene by the reaction of chlorine, in the presence of
	-	(d) Fumaric acid		(a) (i) Direct sunlight and (ii) anhydrous <i>AlCl</i> ₃
37.		ns of benzene (aromatic		-
	hydrocarbon) and its de			(b) (i) Sodium hydroxide and (ii) sulphuric acid
		9; AFMC 1997; BHU 1996, 98]		(c) (i) Ultraviolet light and (ii) anhydrous FeCl ₃
	(a) Electrophilic addition			(d) (i) Anhydrous AlCl ₃ and (ii) direct sunlight
	(b) Electrophilic substit		48.	In Friedel Craft's alkylation, besides $AlCl_3$ the
	(c) Nucleophilic additio(d) Nucleophilic substit			other reactants are [AFMC 1997; CBSE PMT 1999]
38.	Which is most readily n			(a) $C_6H_6 + CH_3Cl$ (b) $C_6H_6 + CH_4$
50.	(a) Benzene	(b) Phenol		(c) $C_6H_6 + NH_3$ (d) $C_6H_6 + CH_3COCl$
	(c) Aniline	(d) Nitrobenzene	49.	Nitration of benzene is a [RPMT 1999]
39.	o, p-directing groups ar		10	(a) Electrophilic displacement
		(b) Deactivating groups		(b) Electrophilic addition
	(c) Neutral groups	(d) None of these		(c) Nucleophilic addition
40.	Which among the follow	wing is the strongest o, p-		(d) Nucleophilic displacement
	directing group	[CBSE PMT 1989]	50.	Benzene shows [RPMT 1999]
	(a) OH	(b) <i>Cl</i>		(a) Substitution (b) Addition
	(c) $C_6 H_5$	(d) Br		(c) Oxidation (d) All of these
41.		s most reactive towards	51.	Benzene can be obtained in the reaction
	electrophilic nitration is			[RPET 2000; Bihar MEE 1997]
		MS 1998; MP PET/PMT 1998]		(a) Ethene + 1, 3-butadiene
	(a) Toluene	(b) Benzene		(b) Trimerisation of ethyne
42	(c) Benzoic acid	(d) Nitrobenzene		(c) Reduction of <i>PhCHO</i>
42.	most readily sulphonate	the compound that can be		(d) All of these
		ADT Bihar 1995; KCET 2005]	52.	Thiophene and benzene are separated by [RPET 2000]
	(a) Benzene	(b) Nitrobenzene		(a) Sulphonation of thiophene
	(c) Toluene	(d) Chlorobenzene		(b) Sulphonation of benzene
43.	Which of the following	g would be least reactive		(c) Nitration of thiophene
	towards bromine	[NCERT 1981]		(d) Nitration of benzene
	(a) Nitrobenzene	(b) Phenol	53.	Which of the following is a hydrocarbon[AFMC 1992]
	(c) Anisole	(d) Chlorobenzene		(a) Urea (b) Benzene
44.		g, the compound that is		(c) Ammonium cyanate (d) Phenol
	nitrated with difficulty		54.	Aromatic compounds burn with sooty flame
	(a) Benzene	(b) Nitrobenzene		because



[BIT 1991]

- (a) They have a ring structure of carbon atoms
- (b) They have a relatively high percentage of hydrogen
- (c) They have a relatively high percentage of carbon
 - (d) They resist reaction with oxygen of air
- Among the following compound which one is 55. planar in shape [MP PMT 2000]
 - (a) Methane
- (b) Acetylene
- (c) Benzene
- (d) Isobutane
- Among the following statements on the nitration of aromatic compounds, the false one is[IIT-JEE 1997]
 - (a) The rate of nitration of benzene is almost the same as that of hexadeuterobenzene
 - (b) The rate of nitration of toluene is greater than that of benzene
 - (c) The rate of nitration of benzene is greater than that of hexadeuterobenzene
- (d) Nitration is an electrophilic substitution reaction
- Methyl group attached to benzene can be oxidised 57. to carboxyl group by reacting with
 - (a) Fe_2O_3
- (b) $AgNO_3$
- (c) $KMnO_A$
- (d) Cr_2O_3
- How is O_2N^{-1}

is widely used[MP PET 2002]

 NO_{2}

 CH_3

- (a) Insecticide
- (b) Drug
- (c) Explosive
- (d) Dye
- The compound 'A' when treated with HNO3 (in presence of H_2SO_4) gives compound 'B' which is then reduced with Sn and HCl to aniline. The compound 'A' is [MP PET 2002]
 - (a) Toulene
- (b) Benzene
- (c) Ethane
- (d) Acetamide
- Which is formed when benzene is heated with 60. chlorine in the presence of sunlight

[CPMT 2000; KCET (Med.) 2000; MP PMT 1993; MP PET 2002 AIIMS 1999]

- (a) $C_6H_5CCl_3$
- (b) $C_6H_5CHCl_2$
- (c) $C_6H_5CH_2Cl_2$
- (d) $C_6H_6Cl_6$
- The compound used as an explosive is 61.

[Kerala (Engg.) 2002; MP PET 2002; MP PMT 1993]

- (a) 2,4, 6-tribromoaniline (b)1,3, 5-trinitrobenzene
- (c) 2,4, 6-trichlorotoluene (d)1,3, 5-trichlorobenzene
- (e) 2,4, 6-trinitrotoluene
- **62.** Adding of Cl_2 to benzene in the presence of $AlCl_3$ is an example of [Bihar MEE 1996]
 - (a) Addition
- (b) Halogenation
- (c) Substitution
- (d) Elimination

- (e) None of these
- 63. What happens when naphthalene balls are put inside kerosene [Kerala (Med.) 2002]
 - (a) Precipitates heating
- (b) Dissolves
 - upon
- (c) Dissolves easily
- (e) None of these Three fused benzene rings are found in

[Kerala (Engg.) 2002]

- (a) Naphthalene
- (b) Anthracene
- (c) Phenanthroline
- (d) Triphenyl methane

(d) Does not dissolve

Product obtained after nitration of nitrobenzene

[RPMT 1997]

(a) TNT

64.

- (b) 1, 3-dinitrobenzene
- (c) Picric acid
- (d) 1, 4-dinitrobenzene
- After ozonolysis of benzene (not hydrolysis), the product is

[RPMT 1997; CPMT 1997]

- (a) Benzene triozonide (b) Glyoxal
- (c) Ethanediol
- (d) All of them
- Which acid will not form hydrocarbon [CPMT 1997]
 - (a) Cinnamic acid
- (b) Isothallic acid
- (c) Salicylic acid
- (d) Picric acid
- **68.** Catalytic dehydrogenation of *n*-haptane in presence of Cr_2O_3/Al_2O_3 at 750 K gives[Roorkee 1999]
 - (a) iso-heptane
- (b) 1-heptene
- (c) toluene
- (d) 2,
- dimethylpentene-1
- $C_6H_6 \xrightarrow{HNO_3} X \xrightarrow{Cl_2} Y$. In the above sequence Y $FeCl_2$

[AIIMS 1999]

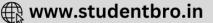
3-

- (a) 1-nitrochloro benzene
- (b)3-nitrochlorobenzene
- (c) 4-nitrochlorobenzene
- (d)1,2-nitrochlorobenzene Which of the following has lowest knocking
- property (a) Olefins (b) Straight
- paraffins

- - (c) Aromatic hydrocarbons (d)Branched chain paraffins In which of the following, the bond length
- between carbon and carbon atom is equal[CPMT 1997]
 - (a) 2-butene
- (b) Benzene
- (c) 1-butene
- (d) 1-propyne
- Benzene is prepared in laboratory from which one of the following compounds [MP PMT 1996]
 - (a) $C_6N_5N_2Cl$
- (b) C_6H_5OH
- (c) C_6H_5COONa
- (d) $C_6H_5SO_3H$
- Which of the following is not used in Friedelcrafts reaction

[KCET 2000]

- (a) Phenyl acetanilide
- (b) Bromobenzene
- (c) Benzene
- (d) Chlorobenzene
- In chlorination of benzene, the reactive species is 74. [MP PET 2000]
 - (a) Cl^+
- (b) Cl⁻



76. Benzene molecule is MP PET 2001; Pb. PMT 2004] (a) Tetrahedral (b) Planar (c) Pyramidal (d) Trigonal 77. Pyridine is less basic than triethylamine because [AIIMS 2005] (a) Pyridine has aromatic character (b) Nitrogen in pyridine is sy² hybridized (c) Pyridine is a cyclic system (d) In pyridine, lone pair of nitrogen is delocalized 78. Electrophile in the case of chlorination of benzene in the presence of FeCl₃ is (a) Cl² (b) Cl² (c) Cl (d) FeCl₃ (a) FeCl₃ (a) FeCl₃ (a) FeCl₃ (a) FeCl₃ (a) FeCl₃ (b) Cl² (c) Cl (d) FeCl₃ (a) FeCl₃ (a) FeCl₃ (a) FeCl₃ (a) FeCl₃ (b) FeCl₃ (a) FeCl₃ (a) FeCl₃ (b) FeCl₃ (b) FeCl₃ (c) Fellowing will undergo meta substitution on monochlorination (b) M-position (c) Pyposition (d) Both o- and p-positions 80. Nitration of toluene takes place at [NCERT 1990] (a) -position (b) M-position (c) Pyposition (d) Both o- and p-positions 81. Which of the following is not o, p—directing group (a) -position (b) M-position (c) Pyposition (d) FeCl₃ (d)	34 пуштосать	11,	iyul ocal boli				
75. Which of following having delocalised electron (BCECE 2005) (a) Benzene (b) Cyclohexane (c) C C H ₄ (d) C ₂ H ₆ (d) C ₂ H ₆ (d) C ₂ H ₆ (a) Tetrahedral (b) Planar (c) Pyramidal (d) Trigonal (d) Trigonal (e) Pyridine is less basic than triethylamine because (a) Pyridine is a cyclic system (d) In pyridine is sys p² hybridized (c) Pyridine is a cyclic system (d) In pyridine, lone pair of nitrogen is the presence of PcC ₄ , is (a) C' (b) C' (c) Cl (d) PcC ₄ , (d) PcC ₄ (d) PcC ₇ (d) PcC ₇ (e) Cl (d) PcC ₁ (d) PcC ₁ (e) Cl (d) PcC		(c) <i>Cl</i> .	(d)) Cl ₂ -		(c) C_2H_5I /anhy. $AlCl_3$	(d) CH ₃ COCl / AlCl ₃
(a) Benzene (b) Cyclohexane (c) CH ₄ (d) C ₂ H ₆ (d) C ₃ H ₆ (d) C ₄ H ₇ (d) Anthracene oil (d) Trigonal (e) Pyridine is less basic than triethylamine because (a) Pyridine is a sorbit character (b) Nitrogen in pyridine is sorbit of (c) Pyridine is a cyclic system (d) In pyridine, lone pair of nitrogen is the presence of FcC ₄ , is (d) C ₇ (d) FcC ₁ , is (d) C ₇ (d) FcC ₁ , is (e) C ₁ (d) FcC ₁ (d) FcC ₁ (d) FcC ₁ (e) C ₁ (d) C ₁ (d) C ₁ (d) C ₁ (e) C ₂ (d) FcC ₁ (e) C ₁ (d) FcC ₁ (e) C ₂ (d) FcC ₁ (e) C ₁ (d) FcC ₁ (e) C ₂ (d) FcC ₁ (e) C ₂ (d) FcC ₁ (e) C ₁ (d) FcC ₂ (e) C ₂ (d) FcC ₁ (e) C ₂ (d) FcC ₁ (e) C ₃ (d) C ₁ (d) C ₁ (e) C ₄ (e	of following h	-		delocalised electron	88.		
(c) CH ₄ (d) C ₂ H ₆ (e) Benzene molecule is [MF PET 2001; Pb. PMT 2004] (a) Tetrahedral (b) Planar (c) Pyramidal (d) Trigonal (d) Trigonal (d) Pyridine is less basic than triethylamine because [LIIMS 2005] (a) Pyridine has aromatic character (b) Nitrogen in pyridine is sy ² hybridized (c) Pyridine is a cyclic system (d) In pyridine, lone pair of nitrogen is delocalized 78. Electrophile in the case of chlorination of benzene in the presence of FeCt ₄ is (a) Ct (c) Ct (d) FeCt ₃ 79. Which one of the following will undergo metas substitution on monochlorination [LIIMS 1991] (a) Ethoxy ethane (b) Chlorobenzene (c) Ethyl benzoate (d) Phenol 80. Nitration of foluene takes place at [NoERT 1990] (a) -position (b) m-position (a) -NIT ₂ (b) -OII (c) Ct (d) FeCt ₃ 81. Which of the following is not o, p—directing group (a) -position (d) Both o- and p-position (d) Both o- and p-position (e) p-yosition (e) p-yosition (d) Both o- and p-position (e) p-yosition (d) Both o- and p-position (e) p-yosition (e) p-yosition (d) Both o- and p-position (e) p-yosition (e) p-	zene	(a) Be	e (b)			(a) Light oil	(b) Heavy oil
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79. Which one of the following will undergo meta substitution on monochlorination [AIIMS 1991] (a) Ethoxy ethane (b) Chlorobenzene (c) Ethyl benzoate (d) Phenol (c) Ethyl benzoate (d) Phenol (d) Soda lime and zinc dust (c) Zinc dust and sodaium hydroxide (d) Soda lime and copper (order of reactivity of C_2H_6 , C_2H_4 and C_2H_2 in the solution of the following is not o, p—directing group (c) C_1 (halogens) (d) C_2 (d) C_1 (d) C_2 (e) C_2 (d) C_2 (d) C_2 (d) C_2 (e) C_2 (e) C_2 (d) C_2 (d) All are equally reactive (e) C_1 (e) C_2 (d) C_2 (d) C_2 (d) C_2 (e) C_2 (e) C_2 (e) C_2 (f) C_2 (d) All are equally reactive (e) C_1 (e) C_2 (d) C_2 (d) C_2 (d) C_2 (e) C_2 (e) C_2 (e) C_2 (e) C_2 (f) C_2 (f) C_2 (d) All are equally reactive (e) C_1 (e) C_2 (f)		(c) Cl	(d)) FeCl ₃			or with 1. A and 1 are
substitution on monochlorination [AIIMS 1991] (a) Ethoxy ethane (b) Chlorobenzene (c) Ethyl benzoate (d) Phenol (c) Ethyl benzoate (d) Phenol (d) Phenol (d) Phenol (e) Order of reactivity of C_2H_6 , C_2H_4 and C_2H_2 (d) Soda lime and copper (d) Soda lime and copp	one of the f	Which	of the followir	ng will undergo meta		1 copout . cry	[KCET 2004]
(c) Ethyl benzoate (d) Phenol 80. Nitration of toluene takes place at [NCERT 1990] (a) o -position (b) m -position (c) p -position (d) Both o - and p -positions 81. Which of the following is not o , p -directing group (a) a - M + a 2 (b) a - M + a 2 (c) a 4 a 4 a 4 a 4 a 4 a 5 a 5 a 6 a 7 a 8. Benzene can react with (a) B 7 a 8. Benzene can react with (b) B 8. The compound 'A' having formula C_8H_{10} (a) $B^{-1}V_{10}$ (b) $C^{-1}V_{10}$ (c) $C^{-1}V_{10}$ (d) $C^{-1}V_{10}$ (e) $C^{-1}V_{10}$ (d) $C^{-1}V_{10}$ (e) $C^{-1}V_{10}$ (d) $C^{-1}V_{10}$ (e) $C^{-1}V_{10}$ (d) $C^{-1}V_{10}$ (e) $C^{-1}V_{10}$ (e) $C^{-1}V_{10}$ (d) $C^{-1}V_{10}$ (e) $C^{-1}V_{10}$ (e) $C^{-1}V_{10}$ (e) $C^{-1}V_{10}$ (for $C^{-1}V_{10}$ (h) $C^{-1}V_{10}$	ition on mond	substit	n on monochlorin	nation [AIIMS 1991]		(a) Zinc dust and soda	
80. Nitration of toluene takes place at [NCERT 1990] (a) o -position (b) m -position (c) p -positions (d) Both o - and p -positions 81. Which of the following is not o , p -directing group (a) $-NH_2$ (b) $-OH$ (c) $-X$ (halogens) (d) $-CHO$ 82. Benzene can react with (a) Br_2 water (b) HNO_3 (c) H_2O (d) CH_3OH 83. The compound 'A' having formula C_8H_{10} (a) C_8H_{10} (a) C_9H_{10} (b) C_9H_{10} (c) C_9H_{10} (d) C_9H_{10} (a) C_9H_{10} (a) C_9H_{10} (b) C_9H_{10} (c) C_9H_{10} (d) C_9H_{10} (e) C_9H_{10} (e) C_9H_{10} (e) C_9H_{10} (e) C_9H_{10} (e) C_9H_{10} (e) C_9H_{10} (f) C_9H_{10} (h) C_9H_{10} (e) C_9H_{10} (e) C_9H_{10} (f) C_9H_{10} (h) C_9H_{10} (g) C_9H_{10} (h) C_9H_{10}	oxy ethane	(a) Eth	ethane (b)) Chlorobenzene		(b) Soda lime and zinc	dust
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(c) p -position (d) Both o- and p -positions 81. Which of the following is not o , p -directing group (a) $-NH_2$ (b) $-OH$ (c) $-X$ (halogens) (d) $-CHO$ 82. Benzene can react with [UPSEAT 2003] (a) Br_2 water (b) HNO_3 (c) H_2O (d) CH_3OH 83. The compound 'A' having formula C_8H_{10} (aromatic) which gives 1 mononitro substitute and 3 nitrosubstitute compound is (a) m -Xylene (b) p -Xylene (c) O -Xylene (d) E -thylene (c) E -thylene (b) E -thylene (c) E -thylene (d) E -thylene (e) E -thylene (c) E -thylene (c) E -thylene (c) E -thylene (d) E -thylene (e) E -thylene (c) E -thylene (c) E -thylene (c) E -thylene (d) E -thylene (e) E -thylene (e) E -thylene (following the most basic contains (a) E -thylene (b) E -thylene (b) E -thylene (c) E -thylene (a) E -thylene (b) E -thylene (b) E -thylene (c) E -thylene (d) E -thylene (e) E -thylene (a) E -thylene (b) E -thylene (b) E -thylene (c) E -thylene (b) E -thylene (c) E -thylene (d) E -thylene (e) E -thylene (hand) (following the most basic contains (a) E -thylene (b) E -thylene (b) E -thylene (b) E -thylene (c) E -thylene (b) E -thylene (c) E -thylene (d) E -thylene (e) E -thylene (following the most basic contains (a) E -thylene (b) E -thylene (b) E -thylene (b) E -thylene (c) E -thylene (b) E -thylene (c) E -thylene (d) E -thylene (e) E -thylene (following the most basic contains (a) E -thylene (b) E -thylene (b) E -thylene (b) E -thylene (c) E -thylene (d) E -thylene (e) E -thylene (following the following the most basic contains (a) E -thylene (b) E -thylene (b) E -thylene (b) E -thylene (c) E -thylene (a) E -thylene (b) E -thylene (b) E -thylene (c) E -thylene (d) E -thylene (e) E -thylene (following the following the following the following the following the following the following t	on of toluene t	. Nitrati	f toluene takes p	lace at [NCERT 1990]		(d) Soda lime and copp	er
positions 81. Which of the following is not o , p —directing group (a) $-NH_2$ (b) $-OH$ (c) $-X$ (halogens) (d) $-CHO$ (c) $-X$ (halogens) (d) $-CHO$ (e) $-X$ (halogens) (d) $-X$ (halogens) (e) $-X$ (halogens) (d) $-X$ (e) $-X$ (halogens) (f) $-X$ (halogens) (having formula $-X$ (gives formula $-X$ (halogens) (having formula $-X$ (halogens) (hal	osition	(a) o-r	ion (b)) <i>m</i> -position	92.	Order of reactivity of C	C_2H_6, C_2H_4 and C_2H_2 is
81. Which of the following is not o , p —directing group (a) $-NH_2$ (b) $-OH$ (c) $-X$ (halogens) (d) $-CHO$ 82. Benzene can react with [UPSEAT 2003] (a) Br_2 water (b) Br_2 (c) Br_2 water (b) Br_2 (a) Br_2 water (b) Br_2 (c) Br_2 water (b) Br_2 (a) Br_2 water (b) Br_2 (c) Br_2 (a) Br_2 water (b) Br_2 (c) Br_2 (a) Br_2 water (b) Br_2 (c) Br_2 (a) Br_2 water (b) Br_2 (a) Br_2 (b) Br_2 (c) Br_2 (c) Br_2 (d) Br_2 (e) Br_2 (e) Br_2 (e) Br_2 (e) Br_2 (f) Br_2 (e) Br_2 (e) Br_2 (f) Br_2 (f) Br_2 (h) Br_2 (c) Br_2 (c) Br_2 (d) Br_2 (e) Br_2 (e) Br_2 (e) Br_2 (f) $Br_$	osition		ion (d)) Both o- and p-			[MH CET 2004]
81. Which of the following is not 0, p —directing group (a) $-NH_2$ (b) $-OH$ reactive (c) $-X$ (halogens) (d) $-CHO$ 82. Benzene can react with [UPSEAT 2003] (a) Br_2 water (b) HNO_3 [AFMI (c) H_2O (d) CH_3OH (a) Kolbe's reaction (b) Williamson's synthesis (aromatic) which gives 1 mononitro substitute and 3 nitrosubstitute compound is (a) m -Xylene (b) p -Xylene (c) o -Xylene (d) Ethyl benzene (c) o -Xylene (d) Ethyl benzene (e) Benzoic acid (d) Toluene (c) Benzoic acid (d) Toluene (c) Animal (d) Charcoal (d) Charcoal (e) Final (c) Animal (d) Charcoal (d) Enzero (e) Naphthalene (d) Xylene (d) Xylene (e) Naphthalene (d) Xylene (d) Xylene (e) Naphthalene (d) Xylene (d) Senzoic acid (d) Xylene (e) Senzaldehyde (d) Benzoic acid (d) Benzoic acid (d) Senzoic acid (d) Xylene (e) Senzaldehyde (d) Benzoic acid (d) Benzoic acid (d) Senzoic acid (d) Xylene (e) Senzaldehyde (d) Benzoic acid (d) Benzoic acid						(a) $C_2H_6 > C_2H_4 > C_2H$, (b) $C_2H_2 > C_2H_6 > C_2H_4$
(a) $2M_{12}$ (b) $2M_{12}$ (c) $-X$ (halogens) (d) $-CHO$ reactive 82. Benzene can react with (a) Br_2 water (b) HNO_3 (a) Br_2 water (b) HNO_3 (a) Kolbe's reaction (b) Williamson's synthesis (c) H_2O (d) CH_3OH (a) Kolbe's reaction (b) Williamson's synthesis (c) Wurtz reaction (d) Sandmeyer reactive and 3 nitrosubstitute compound is (a) m -Xylene (b) p -Xylene (c) o -Xylene (d) Ethyl benzene 84. Catalytic hydrogenation of benzene gives[AIIMS 1996] (a) Xylene (b) Cyclohexane (c) Benzoic acid (d) Toluene 85. Benzene is obtained from [CPMT 1996] (a) Coaltar (b) Plant (c) Animal (d) Charcoal 86. The 'middle oil' fraction of coaltar distillation contains [MP PET 2001] (a) Benzene (b) Anthracene (c) Naphthalene (d) Xylene (d) Xylene (d) Senzaldehyde (d) Benzoic acid			_				
82. Benzene can react with (a) Br ₂ water (b) HNO ₃ (c) H ₂ O (d) CH ₃ OH 83. The compound 'A' having formula C ₈ H ₁₀ (aromatic) which gives 1 mononitro substitute and 3 nitrosubstitute compound is (a) m-Xylene (b) p-Xylene (c) o-Xylene (d) Ethyl benzene 84. Catalytic hydrogenation of benzene gives[AIIMS 1996] (a) Xylene (b) Cyclohexane (c) Benzoic acid (d) Toluene 85. Benzene is obtained from [CPMT 1996] (a) Coaltar (b) Plant (c) Animal (d) Charcoal 86. The 'middle oil' fraction of coaltar distillation contains [MP PET 2001] (a) Benzene (b) Anthracene (c) Naphthalene (d) Xylene (d) Xylene (d) Xylene (e) Naphthalene (d) Xylene (d) Sandmeyer reaction (d) Folicient (c) Mixture of both (d) Heptylene (a) Benzene (b) Aniline (c) Acetanilide (d) p-nitroaniline (d) Benzene (e) Benzaldehyde (d) Benzoic acid (d) Benzoic acid	I_2	(a) −Λ	(b)) <i>–OH</i>	reac		6 (a) im are equally
alkene (a) Br_2 water (b) HNO_3 (c) H_2O (d) CH_3OH 83. The compound 'A' having formula C_8H_{10} (a) m -xylene (b) m -xylene (c) m -xylene (d) m -xylene (d) m -xylene (d) m -xylene (e) m -xylene (f) m -xylene (h) m -xylene	(halogens)	(c) $-X$	ılogens) (d)) <i>-CHO</i>			ng vield both alkane and
(a) Br_2 water (b) HNO_3 (c) H_2O (d) CH_3OH (a) Kolbe's reaction (b) Williamson's S3. The compound 'A' having formula C_8H_{10} (aromatic) which gives 1 mononitro substitute and 3 nitrosubstitute compound is (a) m -Xylene (b) p -Xylene (c) o -Xylene (d) Ethyl benzene (b) p -Xylene (c) o -Xylene (d) Ethyl benzene (e) o -Xylene (b) Cyclohexane (c) Benzoic acid (d) Toluene (e) Benzene is obtained from [CPMT 1996] (a) Coaltar (b) Plant (c) Animal (d) Charcoal (d) Charcoal (e) Animal (d) Charcoal (e) Naphthalene (d) Xylene (e) Maphthalene (d) Xylene (c) Benzaldehyde (d) Benzoic acid (d) Benzoic acid (d) Representation (b) Williamson's synthesis (c) Wurtz reaction (d) Sandmeyer reaction	e can react wi	. Benzer	in react with	[UPSEAT 2003]	93.		ing yield both dikane and
83. The compound 'A' having formula C_8H_{10} (a) Kolbe's reaction (b) Williamson's synthesis (cromatic) which gives 1 mononitro substitute and 3 nitrosubstitute compound is (a) m -Xylene (b) p -Xylene (c) o -Xylene (d) Ethyl benzene (e) o -Xylene (d) Ethyl benzene (e) o -Xylene (b) Cyclohexane (c) Benzoic acid (d) Toluene (e) Plant (c) Animal (d) Charcoal (d) Charcoal (e) Men toluene is treated with o -Milline (c) Animal (d) Charcoal (d) Enzoic acid (d) Toluene (e) Naphthalene (d) Xylene (b) Anthracene (c) Naphthalene (d) Xylene (d) Enzoic acid (d) Xylene (e) Anthracene (c) Renzaldehyde (d) Benzoic acid (d) Sandmeyer reasynthesis synthesis (c) Wurtz reaction (d) Sandmeyer reasynthesis synthesis (c) Wurtz reaction (d) Sandmeyer reasynthesis (c) Wurtz reaction (d) Animperson (d) Possible (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3 + Cr_2O_3$) catalyst at 773 K gives [DCI (Al $_2O_3$	water	(a) <i>Br</i> ₂	ter (b)) HNO_3			[AFMC 2004]
(aromatic) which gives 1 mononitro substitute and 3 nitrosubstitute compound is (a) m-Xylene (b) p-Xylene (c) o-Xylene (d) Ethyl benzene 84. Catalytic hydrogenation of benzene gives[AIIMS 1996] (a) Xylene (b) Cyclohexane (c) Benzoic acid (d) Toluene 85. Benzene is obtained from (d) Coaltar (e) Animal (f) Charcoal 86. The 'middle oil' fraction of coaltar distillation contains [MP PET 2001] (a) Benzene (b) Cyclohexane (c) Acetanilide (d) Charcoal (a) Benzene (b) Anthracene (c) Benzoic acid (d) Charcoal (a) Benzene (b) Anthracene (c) Benzaldehyde (d) Benzoic acid (d) Coaltar (d) Charcoal (a) Benzene (b) Chlorobenzene (c) Benzaldehyde (d) Benzoic acid)	(c) H_2	(d)) <i>CH</i> ₃ <i>OH</i>		(a) Kolbe's reaction	
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(a) m -Xylene (b) p -Xylene (c) o -Xylene (d) Ethyl benzene (a) Benzene (b) Toluene (a) Xylene (b) Cyclohexane (c) Benzoic acid (d) Toluene (e) Plant (c) Animal (d) Charcoal (d) Charcoal (e) Animal (d) Charcoal (e) Animal (d) Charcoal (e) Animal (d) Charcoal (e) Animal (e) Anthracene (formal matrix m					94.	Aromatisation of n-h	neptane by passing over
(c) o-Xylene (d) Ethyl benzene 84. Catalytic hydrogenation of benzene gives[AIIMS 1996] (a) Xylene (b) Cyclohexane (c) Benzoic acid (d) Toluene 85. Benzene is obtained from [CPMT 1996] (a) Coaltar (b) Plant (c) Animal (d) Charcoal 86. The 'middle oil' fraction of coaltar distillation contains [MP PET 2001] (a) Benzene (b) Anthracene (c) Naphthalene (d) Xylene (a) Benzene (b) Toluene (c) Mixture of both (d) Heptylene (a) Benzene (b) Amongst the following the most basic contise (a) Benzylamine (b) Aniline (c) Acetanilide (d) p-nitroaniline (d) Poundant (=			$(Al_2O_3 + Cr_2O_3)$ catalyst	at 773 K gives [DCE 2004]
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(c) Benzoic acid (d) Toluene is 85. Benzene is obtained from [CPMT 1996] (a) Coaltar (b) Plant (c) Animal (d) Charcoal 86. The 'middle oil' fraction of coaltar distillation contains [MP PET 2001] (a) Benzene (b) Anthracene (c) Naphthalene (d) Xylene is [Senzene is obtained from [CPMT 1996] (a) Benzylamine (b) Aniline (c) Acetanilide (d) p-nitroaniline (c) Acetanilide (d) p-nitroaniline (d) Produced (d) Benzene (e) Co Benzaldehyde (d) Benzoic acid	c hydrogenat	. Cataly	ydrogenation of t	oenzene gives[AIIMS 1996]		(c) Mixture of both	(d) Heptylene
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(a) Coaltar (b) Plant (c) Animal (d) Charcoal (c) Acetanilide (d) p-nitroaniline (d) Per 2001] (a) Benzylamine (b) Aniline (d) p-nitroaniline (e) Acetanilide (d) p-nitroaniline (f) Per 2001] (a) Benzene (b) Anthracene (a) Benzene (b) Chlorobenzene (c) Naphthalene (d) Xylene (c) Benzaldehyde (d) Benzoic acid	zoic acid	(c) Be	c acid (d)) Toluene		is	
(c) Animal (d) Charcoal (c) Acetanilide (d) p-nitroaniline 86. The 'middle oil' fraction of coaltar distillation contains [MP PET 2001] (a) Benzene (b) Anthracene (c) Naphthalene (d) Xylene (c) Acetanilide (d) p-nitroaniline (d) p-nitroan	e is obtained f	. Benzer	obtained from	[CPMT 1996]			[AIEEE 2005]
86. The 'middle oil' fraction of coaltar distillation contains [MP PET 2001] (a) Benzene (b) Anthracene (c) Naphthalene (d) Chartoal 96. When toluene is treated with KMnO ₄ , we produced [AFMO (a) Benzene (b) Chlorobenzene (c) Benzaldehyde (d) Benzoic acid	ltar	(a) Co	· (b)) Plant		=	
contains produced [MP PET 2001] [AFMo (a) Benzene (b) Anthracene (a) Benzene (b) Chlorobenzene (c) Naphthalene (d) Xylene (c) Benzaldehyde (d) Benzoic acid	mal	(c) An	(d)) Charcoal		• •	-
[MP PET 2001] [AFMO (a) Benzene (b) Anthracene (a) Benzene (b) Chlorobenzene (c) Naphthalene (d) Xylene (c) Benzaldehyde (d) Benzoic acid	iddle oil' fra	. The 'r	le oil' fraction	of coaltar distillation	96.		ted with $KMnO_4$, what is
(a) Benzene(b) Anthracene(a) Benzene(b) Chlorobenzene(c) Naphthalene(d) Xylene(c) Benzaldehyde(d) Benzoic acid	s	contai				produced	
(c) Naphthalene (d) Xylene (c) Benzaldehyde (d) Benzoic acid						() D	[AFMC 2005]
87. Lindane can be obtained by reaction of benzene 97. In presence of light & heat toluene chloring				-			
react with across NaOU to give	can be obta		in be obtained b	y reaction of benzene	97.		
with react with aqueous NaOH to give		with		[DCD cccc]		react with aqueous Nat	OH to give [Kerala CET 2005]
[Cl Janhy AlC	(a) C	Janhy AICI (b)			(a) n-Cresol	[Nerala CE1 2005]
(a) CH_3Cl /anhy. $AlCl_3$ (b) Cl_2 /sunlight (a) o-Cresol	₃ Cι / aiiiiy. AlCl	(a) CE	I annly. $AiCl_3$ (D)	j ci ₂ /sumignt		(a) 0 CIC301	

2004]

- (b) p-Cresol
- (c) Mixture of o- Cresol & p-Cresol
- (d) Benzoic acid
- (e) 1, 3, 5 trihydroxy toluene



- In the case homologous series of alkanes, which 1. one of the following statements is incorrect [JIPMER 2000]
 - (a) The members of the series are isomers of each other
 - (b) The members of the series have similar chemical properties
 - (c) The members of the series have the general formula $C_n H_{2n+2}$, where n is an integer
 - (d) The difference between any two successive members of the series corresponds to 14 unit of relative atomic mass
- How many primary, secondary, tertiary and 2. quaternary carbons are present in the following hydrocarbon

$$CH_3 - CH(CH_3) - C(CH_3)_2 - CH_2 - CH(CH_3) - CH_2 - CH_3$$

	Primary	Secondar	Tertiary	Quaterna
		у		ry
(a	6	2	2	1
)				
(b	2	6	3	0
)				
(c	2	4	3	2
)				
(d	2	2	4	3
)				

The octane number of a sample of petrol is 40. It 3. means that its knocking property is equal to the mixture of

[MP PMT 2003]

- (a) 40% *n*-heptane + 60% *iso*-octane
- (b) 40% petrol + 60% iso-octane
- (c) 60% *n*-heptane + 40% *iso*-octane
- (d) 60% petrol + 40% iso-octane
- Formation of 2-butene as major product by 4. dehydration of 2-butanol is according to [MP PMT 1995]
 - (a) Markownikoff rule (b) Saytzeff rule
 - (c) Peroxide effect
- (d) Anti-Markownikoff

rule

5.
$$CH_3C \equiv CCH_3 \xrightarrow{(ii)H_2O/Z_1} CH_3 - C - C - CH_3$$

X in the above reaction is [CPMT 1985; MP PET 1997; Roorkee Qualifying 1998; DPMT 2001]

- (a) HNO_3
- (b) O_2
- (c) O_3
- (d) $KMnO_A$
- 6. Which of the following is Friedel-Craft's reaction
 - (a) $C_6H_6 + FeCl_3 + Cl_2 \rightarrow C_6H_5Cl$
 - (b) $C_6H_5CHO + CH_3CHO + KOH \rightarrow C_6H_5CH = CH CHO$

(c)
$$C_6H_6 + CH_3COCl + AlCl_3 \rightarrow C_6H_5 - C - CH_3$$

(d) $C_6H_5OH + CHCl_3 + KOH \longrightarrow$ Salicylald ehyde

Condition for maximum yield of C_2H_5Cl is

[IIT-JEE 1986]

- (a) C_2H_6 (excess) $+Cl_2 \xrightarrow{UV \text{ Light}}$
- (b) $C_2H_6 + Cl_2 \xrightarrow{\text{Dark}} \text{Room temp.}$
- (c) $C_2H_6 + Cl_2$ (excess) $\xrightarrow{UV \text{ Light}}$
- (d) $C_2H_6 + Cl_2 \xrightarrow{UV \text{Light}}$
- When ethyl alcohol is heated with red phosphorus and HI, then which of the following is formed

[Kurukshetra CEE 1998]

- (a) C_2H_6
- (b) CH_{4}
- (c) C_3H_8
- (d) C_2H_4
- In the Fischer-Tropsch synthesis of petrol.... and 9. are used as the raw materials [KCET 1998]
 - (a) H_2 ; CO
- (b) $CH_4; H_2$
- (c) CH_4 ; CH_3OH
- (d) CH₃OH;CO
- Which one of the following reactions is most suitable for the preparation of n-propyl benzene[MP PET/]
 - (a) Friedel-Craft's reaction (b) Wurtz reaction
 - (c) Wurtz-Fittig reaction(d) Grignard reaction Propane cannot be prepared from which reaction
- 11. [DCE 2003]

(a)
$$CH_3 - CH = CH_2 \xrightarrow{B_2H_6} OH^-$$

- (b) $CH_3CH_2CH_2I \xrightarrow{HI}_P$
- (c) $CH_3CH_2CH_2Cl \xrightarrow{Na}$
- (d) None of these
- 12. The reaction

$$CH_3CH = CH_2 \xrightarrow{(CO+H_2)} CH_3 - CH - CH_3$$
 is

known as

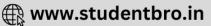
- (a) Wurtz reaction
- (b) Koch reaction
- (c) Clemmensen reduction
- Kolbe's reaction (d)

The compound $CH_3 - C = CH - CH_3$ on reaction with NaIO4 in the presence of KMnO4 gives[CBSE PMT 2003

- (a) $CH_3CHO + CO_2$
- (b) CH₃COCH₃



13.



- (c) $CH_3COCH_3 + CH_3COOH$
- (d) $CH_3COCH_3 + CH_3CHO$
- In the reaction: 14.

 $HC \equiv CH + 2AgNO_3 \xrightarrow{NH_4OH} X + 2NH_4NO_3 + 2H_2O$

- (a) Ag_2C
- (b) Ag_2C_2
- (c) AgC
- (d) AgOH
- Naphthalene is a/an 15.

[AFMC 2004]

- (a) Ionic solid
- (d) Covalent solid (d) Molecular solid
- (c) Metallic solid
- Which of the following is not aromatic [Pb. CET 2000]
 - (a) Benzene
- (b) Naphthalene
- (c) Pyridine
- (d) 1,3,5 heptatriene
- Acetylene reacts with HCN in the presence of 17. Ba(CN), to yield
 - [UP SEAT 2004]
 - (a) 1, 1-dicyanoethane
- (b) 1, 2-dicyanoethane
- (c) Vinyl cyanide
- (d) None of these
- 18. Write the products of the addition reaction $C = C + XY \rightarrow$ [Kerala (Med.) 2002]
 - (a) >C-C <

- Formation of polyethylene from calcium carbide 19. takes place as follows

 $CaC_2 + 2H_2O \rightarrow Ca(OH)_2 + C_2H_2$

 $C_2H_2 + H_2 \rightarrow C_2H_4$

 $n(C_2H_4) \rightarrow (-CH_2 - CH_2 -)_n$

The amount of polyethylene obtained from 64.1 kg CaC_2 is

[AIIMS 1997]

- (a) 7 kg
- (b) 14 kg
- (c) 21 kg
- (d) 28 kg
- Nitrobenzene can be prepared from benzene by using a mixture of conc. HNO_3 and conc. H_2SO_4 . In the nitrating mixture, HNO₃ acts as a[IIT-JEE 1997]
 - (a) Base
- (b) Acid
- (c) Reducing agent
- (d) Catalyst
- A group which deactivates the benzene ring 21. towards electrophilic substitution but which directs the incoming group principally to the oand p-positions is

[Pb. PMT 1998]

- (a) $-NH_2$
- (b) -Cl
- (c) $-NO_2$
- (d) $-C_2H_5$
- Which order is correct for the decreasing reactivity to ring monobromination of the following compounds

- $C_6H_5CH_3$, C_6H_5COOH , C_6H_6 $C_6H_5NO_2$
- (a) I > II > III > IV
- (b) I > III > II > IV
- (c) II > III > IV > I
- (d) III > I > II > IV
- Benzene is obtained by
- [DPMT 2002]
- (a) Substitution of three acetylene molecules
- (b) Addition of three C_2H_2 molecules
- (c) Polymerisation of three C_2H_2 molecules
- (d) Condensation of three C_2H_2 molecules
- Toluene can be oxidised to benzoic acid by[AIIMS 1999]
 - (a) $KMNO_{A}$
- (b) $K_2Cr_2O_7$
- (c) H_2SO_4
- (d) Both (a) and (b)
- $CaC_2 + H_2O \rightarrow A \xrightarrow{H_2SO_4/HgSO_4} B$. Identify A and B 25. in the given reaction [CPMT 2000; BVP 2004]
 - (a) C_2H_2 and CH_3CHO (b) CH_4 and HCOOH

 - (c) C_2H_4 and CH_3COOH (d) C_2H_2 and CH_3COOH
- **26.** The compound *X* on reaction with $HgSO_4 + H_2SO_4$ gives Y which on oxidation gives acetic acid. X is [MP PMT
 - (a) C_2H_2
- (b) C_2H_4
- (c) C_3H_4
- (d) C_4H_6
- What is formed when calcium carbide react with 27. heavy water [CPMT 1999]
 - (a) C_2D_2
- (b) *CaD*₂
- (c) CaD_2O
- (d) CD_2
- The addition of *HBr* is easiest with [MP PMT 2000]
 - (a) $ClCH_2 = CHCl$
- (b) ClCH = CHCl
- (c) $CH_3 CH = CH_2$
- (d) $(CH_3)_2 C = CH_2$
- Identify the species X in the reaction :

Propene + $O(\text{conc. acidic } KMnO_4) \rightarrow X + \text{Formic acid}$

- (a) Acetone
- (b) Acetaldehyde
- (c) Isopropanol
- (d) Acetic acid
- In benzene 1, 3 position is called 30.
 - (a) Meta
- (b) Para
- (c) Ortho
- (d) Odd position
- Which of the following is formed as a result of biological oxidation of benzene in the body of the

[Manipal MEE 1995]

- (a) Acrylic acid
- (b) Cinnamic acid
- (c) Maleic acid
- (d) Gluconic acid
- When acetylene is reacted with HBr, we get 32.

[CPMT 1979; JIPMER 2002]

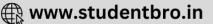
- (a) Methyl bromide
- (b) Ethyl bromide
- (c) Ethylene bromide
- (d) Ethylidene bromide
- The only o, p-directing group which is 33. deactivating in nature is

Which kind of isomerism will butene-2 show

- (a) $-NH_2$
- (b) -OH
- (c) -X (halogens)
- (d) -R (alkyl groups)
- (a) Geometrical
- (b) Optical







- (c) Position (d) None of these
- In presence of light toluene on reaction with 35. chlorine gives

[RPET 1999]

- (a) Benzoyl chloride
- (b) Ortho chlorotoluene
- (c) Para chloro toluene (d) Benzyl chloride
- If ethylene, carbon monoxide and water is heated at high temperature, which of the following is formed

[AIIMS 2000]

- (a) $C_4 H_8 O_2$
- (b) C_2H_5COOH
- (c) CH₃COOH
- (d) $CH_2 = CH COOH$
- Compound C_6H_{12} is an

[AMU 1983]

- (a) Aliphatic saturated compound
 - (b) Alicyclic compound
 - (c) Aromatic compound
 - (d) Heterocyclic compound
- **38.** Identify Z in the following series

$$CH_2 = CH_2 \xrightarrow{HBr} X \xrightarrow{\text{Hydrolysis}} Y \xrightarrow{Na_2CO_3} Z$$

[AIIMS 1983; RPMT 1999]

- (a) C_2H_5I
- (b) C_2H_5OH
- (c) CHI_3
- (d) CH₃CHO
- *n*-pentane and iso pentane can be distinguished by

- (a) Br_2
- (b) O_3
- (c) conc. H_2SO_A
- (d) $KMnO_A$
- $CH \equiv CH + HBr \rightarrow X$, product *X* is 40. [Pb. CET 2003]
 - (a) Ethylene bromide
- (b) Vinyl bromide
- (c) Bromo ethane
- (d) Ethyledine bromide



- Assertion: 1-Butene on reaction with HBr in 1.
 - the presence of a peroxide produces 1-bromo-butane.

involves Tt the Reason free radical

mechanism.

[IIT-JEE (Screening) 2000]

Assertion: Addition of Br_2 to 1-butene gives 2.

two optical isomers.

Reason product contains one

asymmetric carbon. [IIT 1998]

Assertion: Cyclobutane is less stable than 3.

cyclopentane

Presence of bent bonds causes "loss Reason

> of orbital overlap". [AIIMS 1996]

Assertion: Pyrrole is an aromatic heterocyclic

compound.

It has a cyclic, delocalised 6π Reason

electrons.

[AIIMS 1995]

5. Assertion: CH_4 does not react with Cl_2 in

dark.

Reason Chlorination of CH, takes place in

sunlight.

[AIIMS 2001]

Assertion: Alkyl benzene is not prepared by

Friedel-Crafts alkylation

benzene.

Alkyl halides are less reactive than Reason

acyl halides. [AIIMS 2003]

Assertion: 2-Bromobutane on reaction with 7. sodium ethoxide in ethanol gives 1-

butene as a major product. [AIIMS 2004]

1-Butene is more stable than 2-Reason

butene.

8. Assertion: Styrene on reaction with HBr gives

2-bromo-2- phenyl-ethane.

Benzyl radical is more stable than Reason

alkyl radical. [AIIMS 2004]

9. Assertion: Melting point of *n*-butane is higher

than propane.

Reason It is called oscillation effect.

Assertion: Iodination of alkanes is reversible. 10.

> Reason Iodination is carried out

presence of iodic acid.

Assertion: Isobutane on oxidation with KMnO4 11.

gives tert-butyl alcohol.

Oxidising agents have no effect on Reason

alkanes.

Halogenation of 12. Assertion: alkanes is

catalysed by tetraethyl lead.

Reason Halogenation proceeds through free

radical mechanism.

13. Assertion: Neopentane forms

monosubstituted compound.

Reason Neopentane has high bond energy.

Freezing point of neopentane is 14. Assertion:

more than *n*-pentane.

Increase in Van der Waals forces Reason

increases freezing point.

Assertion: Knocking lowers the efficiency of the engine.

> Fuel minimum with

property is preferred.

The presence of Ag^+ enhances the

solubility of alkenes in water.

Alkenes are weakly polar in nature. Reason Assertion: 2-Butanol on heating with H_2SO_4

gives 1-butene and 2-butene.

Reason Dehydration of 2-butanol follows

saytzeff rule.

Reason

Assertion:

16.



knocking

18. Assertion: Ethene on treating with Br_2 in

presence of NaCl forms CH_2ClCH_2Br and CH_2Br-CH_2-Br .

Reason : This addition involves the

formation of free radicals.

19. Assertion: Straight chain alkanes have very

low octane number.

Reason : Quality of gasoline is measured in

terms of octane number.

20. Assertion: Corey-House reaction can be used

to prepare both symmetrical and

unsymmetrical alkanes.

Reason : The reaction involves the

interaction between lithium dialkyl copper with an alkyl halide both of which may contain even or odd

number of carbon atoms.

21. Assertion : All the hydrogen atoms in $CH_2 = C = CH_2$ lie in one plane.

Reason : All the carbon atoms in it are sp^2

hybridized.

22. Assertion: Propene reacts with *HBr* in

presence of benzoyl peroxide to

yield 2-bromopropane.

Reason : In presence of peroxide, the

addition of HBr to propene follows

ionic mechanism.

23. Assertion: Acetylene reacts with sodamide to

evolve H_2 gas.

Reason : Acetylene is a weaker acid than

ammonia.

24. Assertion: Aryl halides are less reactive

towards substitution of halogen

atom.

Reason : Halogens are o, p – directing in

nature.

25. Assertion: Benzene is a solvent for the Friedel

Craft's alkylation of bromobenzene.

Reason : Friedel Craft's reaction is used to

introduced on alkyl or acyl group in

benzene nucleus.

26. Assertion: Benzene removes a butter stain

from a table cloth.

Reason : Butter has an affinity towards

benzene.

27. Assertion: Nitration of toluene is easier than

benzene.

Reason : The methyl group in toluene is

electron-releasing.

28. Assertion: Benzene forms benzene sulphonic

acid with fuming H_2SO_4 at high

temperature.

Reason : The attacking species is SO_3 .

29. Assertion: Activating groups are electron

donors.

Reason : Nitroso group is activating group.

30. Assertion: Benzene reacts with CH_3COCl to

give chlorobenzene.

Reason : Chlorination is an electrophilic

substitution reaction.

31. Assertion: Conjugated polyenes containing odd

number of carbon atoms is known

as annulenes.

Reason : General formula of annulenes is

 $(CH = CH)_n$ where n = 2,3,4 etc.

32. Assertion: Tropylium cation is aromatic in

nature

+

Reason : The only property that determines

its aromatic behaviour is its planar

structure.

33. Assertion: [10] Annulene is not aromatic

though it contains Huckel number

of π -electrons.

Reason : Steric interaction between internal

hydrogens makes it non-planar.

34. Assertion: Rates of nitration of benzene and

hexadeuterobenzene are different.

Reason : C-H bond is stronger than

C-Dbond

35. Assertion: Cyclolpentadienyl anion is much

more stable than allyl anion.

Reason : Cyclopentadienyl anion is aromatic

in character.

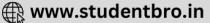


Alkane

1	d	2	а	3	а	4	а	5	b
6	С	7	b	8	а	9	d	10	С
11	d	12	а	13	С	14	d	15	b
16	а	17	b	18	d	19	b	20	b
21	b	22	b	23	а	24	b	25	а
26	d	27	С	28	С	29	d	30	С
31	а	32	а	33	d	34	а	35	а
36	а	37	d	38	а	39	d	40	С
41	b	42	d	43	d	44	а	45	b
46	d	47	С	48	d	49	а	50	С
51	d	52	а	53	а	54	С	55	а
56	d	57	b	58	d	59	b	60	а







61	b	62	а	63	b	64	С	65	С
66	b	67	С	68	b	69	d	70	С
71	b	72	а	73	С	74	С	75	b
76	С	77	С	78	С	79	а	80	С
81	С	82	а	83	d	84	а	85	b
86	а	87	а	88	d	89	а	90	а
91	а	92	b	93	С	94	а	95	С
96	b	97	а	98	d	99	а	100	d
101	С	102	а	103	а	104	b	105	d
106	b	107	d	108	С	109	d	110	b
111	а	112	b	113	d	114	b	115	d
116	d	117	d	118	С	119	С	120	С
121	С	122	b	123	а	124	d	125	С
126	а	127	а	128	b	129	b	130	С
131	а	132	а	133	а	134	b	135	b
136	С	137	а	138	С	139	а	140	С
141	а	142	b	143	а	144	С	145	b
146	С	147	d	148	С	149	d	150	b
151	а	152	С	153	а	154	b	155	ab
156	С	157	b	158	а	159	а	160	С
161	е	162	С	163	а	164	С	165	b
166	d	167	d	168	С	169	С	170	С
171	а	172	d	173	b	174	b	175	С

Alkene	è
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1	С	2	b	3	а	4	b	5	а
6	d	7	а	8	d	9	b	10	d
11	d	12	а	13	a	14	С	15	b
16	a	17	d	18	d	19	b	20	С
21	b	22	d	23	С	24	b	25	С
26	a	27	b	28	d	29	d	30	d
31	С	32	d	33	С	34	а	35	С
36	С	37	а	38	b	39	ac	40	b
41	С	42	а	43	С	44	С	45	а
46	С	47	d	48	d	49	а	50	С
51	a	52	d	53	С	54	а	55	b
56	С	57	d	58	a	59	d	60	b
61	С	62	а	63	b	64	b	65	b
66	d	67	С	68	a	69	С	70	С
71	С	72	С	73	С	74	d	75	С
76	d	77	а	78	d	79	С	80	b
81	С	82	d	83	С	84	а	85	С
86	С	87	b	88	а	89	С	90	С
91	b	92	С	93	b	94	b	95	b

96	b	97	а	98	b	99	b	100	b
101	b	102	С	103	С	104	b	105	b
106	а	107	С	108	а	109	а	110	С
111	С	112	а	113	а	114	b	115	d
116	d	117	а	118	а	119	а	120	b
121	b	122	С	123	С	124	b	125	С
126	С	127	b	128	b	129	а	130	а
131	С	132	b,d	133	а	134	b	135	С
136	С	137	С	138	b	139	а	140	d
141	b	142	С	143	d	144	а	145	d
146	а	147	d	148	abc	149	а	150	а
151	а	152	b	153	С	154	b	155	а
156	С	157	b	158	d	159	b	160	С
161	d	162	b	163	b	164	а	165	а
166	а	167	b	168	С	169	b	170	b
171	а	172	С	173	С	174	а		

Alkyne

1	С	2	С	3	С	4	С	5	b
6	а	7	а	8	а	9	d	10	С
11	а	12	С	13	d	14	а	15	а
16	d	17	d	18	b	19	С	20	С
21	d	22	b	23	d	24	С	25	С
26	d	27	а	28	d	29	а	30	d
31	а	32	С	33	С	34	а	35	а
36	а	37	С	38	d	39	С	40	С
41	а	42	С	43	а	44	С	45	b
46	С	47	а	48	а	49	b	50	b
51	d	52	а	53	b	54	а	55	d
56	а	57	С	58	С	59	а	60	d
61	С	62	С	63	d	64	d	65	a
66	С	67	С	68	а	69	b	70	b
71	b	72	С	73	а	74	b	75	d
76	С	77	b	78	d	79	а	80	С
81	а	82	а	83	d	84	С	85	С
86	а	87	С	88	С	89	d	90	С
91	С	92	d	93	С	94	b	95	d
96	d	97	d	98	b	99	d	100	а
101	b	102	а	103	а	104	а	105	С

Aromatic hydrocarbon

1	С	2	d	3	С	4	С	5	а
6	b	7	С	8	b	9	а	10	С
11	b	12	b	13	а	14	а	15	а



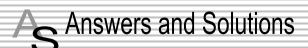
16	d	17	а	18	b	19	b	20	a
21	С	22	а	23	b	24	С	25	С
26	d	27	С	28	а	29	С	30	b
31	d	32	b	33	b	34	С	35	d
36	С	37	b	38	b	39	а	40	а
41	а	42	С	43	а	44	b	45	С
46	d	47	d	48	а	49	а	50	d
51	b	52	b	53	b	54	С	55	С
56	С	57	С	58	С	59	b	60	d
61	е	62	С	63	С	64	b	65	b
66	а	67	d	68	С	69	b	70	С
71	b	72	С	73	а	74	а	75	а
76	b	77	d	78	а	79	С	80	d
81	d	82	b	83	b	84	b	85	а
86	С	87	b	88	а	89	d	90	а
91	b	92	С	93	а	94	b	95	a
96	d	97	d						

Critical Thinking Questions

1	а	2	а	3	С	4	b	5	С
6	С	7	а	8	а	9	а	10	С
11	а	12	b	13	d	14	b	15	d
16	d	17	С	18	а	19	d	20	a
21	b	22	b	23	С	24	b	25	a
26	а	27	а	28	d	29	d	30	a
31	b	32	d	33	С	34	а	35	d
36	b	37	b	38	С	39	d	40	b

Assertion and Reason

1	а	2	а	3	С	4	а	5	b
6	b	7	d	8	b	9	b	10	b
11	b	12	е	13	С	14	b	15	b
16	b	17	а	18	С	19	b	20	а
21	d	22	d	23	С	24	b	25	е
26	b	27	а	28	е	29	С	30	е
31	е	32	С	33	а	34	b	35	а



Alkane

1. (d) C_7H_{16} (C_nH_{2n+2})

3. (a) According to wurtz reaction.

$$2CH_3CH_2CH_2Br + 2Na \xrightarrow{\text{ether}}$$

$$CH_3(CH_2)_4 CH_3 + 2NaBr$$

5. (b) $2CH_3COONa + 2H_2O \xrightarrow{\text{Electolysis}} \rightarrow$ Sodium acetate

$$CH_3 - CH_3 + 2CO_2 + 2NaOH + H_2$$

6. (c) $Pb(C_2H_5)_4 \xrightarrow{\text{heat}} Pb + 4CH_3CH_2$

$$CH_2 - CH_2 + Pb \longrightarrow CH_2 = CH_2 + PbBr_2$$
| Ethene Lead bromide

As leaded gasoline burns, lead metal gets deposited in the engine which is removed by adding ethylene dibromide. The lead bromide is volatile and is carried off with the exhaust gases from the engine

9. (d)
$$C_2H_5I + 2Na + IC_2H_5 \xrightarrow{\text{Dry}} C_2H_5 - C_2H_5 + 2NaI$$

10. (c)
$$(CH_3)_3 CH \xrightarrow{KMnO_4} (CH_3)_3 C - OH$$
 tertiary butyl alcohol

13. (c)
$$RCl + 2Na + RCl \xrightarrow{Dry} 2NaCl + R - R$$
Alkane





- **14.** (d) *iso*-octane *i.e.* 2,2,4-trimethyl pentane has highest octane number.
- **17.** (b) With calculated amount of Grignard reagent, acetyl chloride forms ketones.

$$CH_3COCl + XMgCH_3 \rightarrow CH_3COCH_3 + Mg < \frac{Cl}{X}$$

- 18. (d) CH_4 is tetrahedral H H H SP^3 hybridization
- **22.** (b) Ethane is a saturated compound it can not be catalytically hydrogenated.
- **25.** (a) Branched hydrocarbons are more desirable because they are more volatile.
- **26.** (d) $CH_4 \xrightarrow{Cl_2} CH_3Cl \xrightarrow{Cl_2} CH_2Cl_2 \xrightarrow{Cl_2} CH_2Cl_3 \xrightarrow{Cl_2} CCl_4$. Since this reaction takes place by free radical mechanism. Hence, there is a possibility of formation of ethane.

$$CH_3 + CH_3 \rightarrow CH_3 - CH_3$$
Methyl free radicals Ethane

- **28.** (c) $Al_4C_3 + 6H_2O \rightarrow 3CH_4 + 2Al_2O_3$ Aluminium
 Carbide

 Methane
- 31. (a) $C_2H_5O[H+CH_3] Mg Br \rightarrow CH_4 + Mg$
- 32. (a) $CH_3I + 2H \xrightarrow{Zn/HCl} CH_4 + HI$ $CH_3I + 2Na + ICH_3 \xrightarrow{Dry} CH_3 - CH_3 + 2NaI$
- **36.** (a) Solvent for fat, oil, varnish and rubber
- **37.** (d) Synthetic dyes, drugs, perfumes all are made from coal tar.
- **38.** (a) In alkanes, hybridization is sp^3 . Hence bond angle is $109^o.5'$.
- **39.** (d) $2CH_3COONa + 2H_2O \xrightarrow{\text{Electrolysis}}$

$$CH_3 - CH_3 + 2CO_2 + 2NaOH + H_2$$

41. (b) $CH_3 - CH_2 - COOH + 6HI \xrightarrow{\text{Red } P}$ Propanoic acid

$$CH_3 - CH_2 - CH_3 + 2H_2O + 3I_2$$
Propane

- **42.** (d) $C_2H_5I + 2Na + C_3H_7I \xrightarrow{\text{Dry}} C_2H_5 C_3H_7 + 2NaI$ $C_2H_5I + 2Na + C_2H_5I \xrightarrow{\text{Dry}} C_2H_5 - C_2H_5 + 2NaI$ $C_3H_7I + 2Na + C_3H_7I \xrightarrow{\text{Dry}} C_3H_7 - C_3H_7 + 2NaI$ $C_3H_7I + 2Na + C_3H_7I \xrightarrow{\text{Ether}} C_3H_7 - C_3H_7 + 2NaI$ $C_3H_7I + 2Na + C_3H_7I \xrightarrow{\text{Ether}} C_3H_7 - C_3H_7 + 2NaI$ $C_3H_7I + 2NaI \xrightarrow{\text{Hexane}} C_3H_7 - C_3H_7 + 2NaI$
- **48.** (d) Cyclohexane, is immiscible and lighter than water. Hence, floats on the surface of water.

- **49.** (a) Methane is the main component of natural gas.
- **53.** (a) *Pt./Ni* is used in catalytic reduction of hydrocarbon.
- **55.** (a) Fractional distillation is used because the difference between the boiling point of different component is less.
- **56.** (d) $CH_3 CH_2 Cl + KOH \rightarrow CH_2 = CH_2 + KCl + H_2O$

In presence of alc. *KOH* dehydrohalogenation occur and alkene is formed.

- **57.** (b) Liquefied petroleum gas is a mixture of ethane, propane and butane. The main component is butane.
- **58.** (d) $CH_4 + O_2 \xrightarrow{\Delta} C + 2H_2O$ $\xrightarrow{\text{Limited supply of air}} C + 2H_2O$

It contains 98-99% carbon. It is used in making black ink, paints and shoe polishes.

- **59.** (b) Tetraethyl lead is anti-knocking agent it increases the octane no. of the fuel.
- **60.** (a) *n*-hexane because it is linear therefore strong Vander Waal force.
- **61.** (b) Knocking Sudden and irregular burning of the fuel mixture causing jerks against the piston and gives rise to violent sound. This is known as knocking.
- **63.** (b) *n*-octane
 - Boiling point depends on molecular mass.
 Greater the molecular mass higher will be the boiling point.
 - Boiling point also depends on the structure. If two compounds have same molecular mass then straight chain or linear compound has higher boiling point.
- **64.** (c) $2CH_3COOK + 2H_2O \xrightarrow{\text{Electrolysis}}$

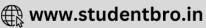
$$CH_3-CH_3+2CO_2+2KOH+H_2\\ \text{Anode} \qquad \text{Cathode}$$

- **65.** (c) Ethane does not decolourise bromine solution because it is a saturated compound.
- **66.** (b) $CH_3COONa + NaOH \xrightarrow{CaO} CH_4 + Na_2CO_3$ Sodium acetate Methane
- **68.** (b) Octane number is related to the percentage of *iso*-octane since *iso*-octane is 81% hence octane number is 81%.
- **70.** (c) $2CH_3COOK + 2H_2O \xrightarrow{\text{Electrolysis}}$

$$\underbrace{2CO_2 + CH_3 - CH_3 + 2KOH + H_2}_{\text{Anode}}$$
 Cathode

71. (b) $CH_3 - C = CH_2 + HCl \rightarrow CH_3 - C - CH_3$ $CH_3 - C + HCl \rightarrow CH_3 - C + CH_3$

$$CH = CH + HCl \rightarrow CH_2 = CH - Cl \xrightarrow{HCl} CH_3 - CHCl_2$$



- **72.** (a) Boiling point of alkanes increases with the number of carbon atoms because surface area increases which increases the Vander Waal forces.
- **74.** (c) The enthalpy of combustion *i.e.*, ΔH is always negative. It means combustion is an exothermic reaction.
- **78.** (c) $CH_3CH_2COONa + NaOH \xrightarrow{CaO} C_2H_6 + Na_2CO_3$
- **79.** (a) $CH_3 CH_2 CH_2 CH_3 + Br_2 \xrightarrow{\text{Light}}$ $CH_3 CH CH_2 CH_3 + CH_3 CH_2 CH_2 CH_2 Br$ Br2-Bromo butane
 (Minor)

2-Bromobutane is the main product because 2^o carbonium ion is more stable than 1^o .

80. (c) $CH_3CH_2CH_2CH_3 \xrightarrow{\text{Cracking}}$

(Main product)

$$CH_4 + CH_3 - CH = CH_2$$

- 83. (d) Anti-knocking properties of fuel increases.
- **84.** (a) $C_6H_{14} \xrightarrow{\Delta} C_4H_{10} + C_2H_2$ Hexane
 Liquid
 Butane
 Ethene
 Gas
- 87. (a) Compounds having active hydrogen $(ROH, H_2O, R-NH_2)$ can form alkane when treated with Grignard's reagent $CH_3CH_2OH + CH_3MgBr \rightarrow CH_4 + Mg < \frac{Br}{OCH_2CH_3}$
- **91.** (a) It is not possible to prepare CH_4 by wurtz reaction.
- 92. (b) Octane number is the percentage by volume of iso-octane in the mixture of iso-octane and *n*-heptane which has the same antiknocking properties as the fuel under examination.
 Given fuel (25% *n*-heptane +75% iso-octane)
 Hence, octane number = 75 (because iso octane is 75%)
- **93.** (c) $CH_3 CH_2 CH_2 Br \xrightarrow{C_2H_5ONa}$ Dehydro halogenation

$$CH_3 - CH = CH_2 + HBr$$

- 94. (a) % of hydrogen = $\frac{\text{Mass of hydrogen}}{\text{Mass of compound}} \times 100$ $CH_4 = \frac{4}{16} \times 100 = 25\%$.
- **95.** (c) Molecular mass can be obtained by the victor mayer process

Molecular mass =
$$\frac{\text{Weight}}{Vml.} \times 22400$$

= $\frac{11}{5600} \times 22400 = 44$

97. (a)
$$CH_3 - CH - C - CH_2 - CH_3 + Zn \rightarrow Br Br$$
2,3-dibromo -3-methyl pentane

$$CH_{3}$$

$$CH_{3} - CH = C - CH_{2} - CH_{3} + ZnBr_{2}$$

$$HI + \bigvee_{\text{Red P (Reduction)}} CH_{3} - CH_{2} - CH_{3} - CH_{2} - CH_{3}$$

$$CH_{3}$$

$$CH_{3}$$

$$3 - \text{Methyl pentane}$$

- 98. (d) Gasoline, kerosene oil, diesel
- 99. (a) $CH_3 CH_2 COONa \xrightarrow{\text{Soda lime}} CH_3 CH_3$
- **100.** (d) Gasoline or petrol composition $C_7 C_{12}$.
- **101.** (c) $CH_4 + 2O_2 \rightarrow CO_2 + 2H_2O$ Methane $C_2H_4 + 3O_2 \rightarrow 2CO_2 + 2H_2O$ Ethene

All hydrocarbons saturated or unsaturated on complete combustion always produce CO_2 and H_2O .

- 102. (a) Free rotation around carbon-carbon bond takes place easily in alkanes. Now ethane and hexachloroethane both are alkanes. But in hexachloroethane bulky chlorine atom hinders the rotation. Therefore least hindered rotation takes place in ethane.
- 103. (a) Hydrocarbons on complete oxidation produce ${\it CO}_2$ and water

$$CH_3 - CH_3 + 3\frac{1}{2}O_2 \rightarrow 2CO_2 + 3H_2O$$

104. (b) $C_{10}H_{22} \xrightarrow{900 \text{ K}} C_{4}H_{8} + C_{6}H_{14}$ Decane Hexane

106. (b)
$$CH_3 - C - CH_3$$

Replaceable hydrogen atoms are present only on 4 primary carbon atoms. Hence, it gives only are monochloro Substituted product.

107. (d) $CH_3COONa + NaOH \xrightarrow{CaO} CH_4 + Na_2CO_3$ Sod. acetate $CH_3I + 2H \xrightarrow{Reduction} CH_4 + HI$

Iodomethan e Reauction
$$Al_4C_3 + 6H_2O \rightarrow 3CH_4 + 2Al_2O_3$$
 Aluminium carbide

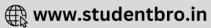
108. (c) Acetylene reacts with ammonical cuprous chloride to form red ppt. of copper acetylide while methane and ethylene do not react (since they do not have acidic hydrogen) They come out from the bottle

$$CH \equiv CH + 2[Cu(NH_3)_2]OH \rightarrow$$

$$Cu - C \equiv C - Cu + 4NH_3 + 2H_2O$$

 CH_4 + Ammonical $Cu_2Cl_2 \rightarrow No$ reaction





 C_2H_4 +Ammonical $Cu_2Cl_2 \rightarrow No$ reaction

- **110.** (b) Alkanes do not give addition reactions because multiple bond is absent.
- 111. (a) $C_6H_{14} \xrightarrow{-Pyrolysis} C_2H_4 + C_4H_{10}$ Hexane Δ Ethene Butane
- 113. (d) In gemdihalide both the halogen atoms are present on the same carbon atom while in vicdihalide both the halogen atoms are present on adjacent carbon atoms.

$$\begin{array}{cccc} CH_3-CHBr_2 & CH_2-CH_2 \\ \text{Gemdihalide} & & | & | \\ Br & Br \\ & & vic\text{-dihalide} \end{array}$$

$$H_{3}C-H_{2}C-H_{2}C-HC-CH_{3}\\ \text{isopropy1 group 2-methy lpentane}\\ CH_{3}CH_{3} & CH_{3}\\ H_{3}C-H_{2}C-HC-C-CH_{3} & H_{3}C-H_{2}C-C-CH_{2}-CH_{3}\\ CH_{3} & CH_{3}\\ 2,2,3\text{-trimethy lpentane} & 3,3\text{-dimethy lpentane}\\ \end{array}$$

- **116.** (d) All-butane, Ethane and Propane are possible in this reaction.
- **118.** (c) Formation of branches in the chain of C atoms C-C-C-C straight chain

Branched chain

- 119. (c) Chlorination of alkane in photochomical reaction which takes place by free radical mechanism. Free radicals are formed by homolytic bond fission or homolysis.
- 120. (c) Marsh gas, Natural gas and coal gas contains CH_4 but producer gas is a mixture of CO and N_2

$$\begin{array}{ccc}
2C & + \underbrace{O_2 + 4N_2}_{Air} & \rightarrow \underbrace{2CO + 4N_2}_{Producer\ gas}
\end{array}$$

- **123.** (a) Fractional distillation is based on the difference in the boiling point of different components.
- 124. (d) Tetraethyl lead (TEL) is an anti-knocking compounds when mixed with petrol tend to improve the octane no. and therefore, decreases the knocking in the cylinder of the combustion engine.
- **128.** (b) Petrol sample 30% *n*-heptane + 70% *iso*-octane since *iso*-octane is 70%. Hence, octane no. is 70.
- **131.** (a) $CH_3 CH = CH_2 + H_2 \xrightarrow{Ni} CH_3 CH_2 CH_3$ Propane
- **132.** (a) As the number of branches increases, surface area decreases, due to which Vander Waal

forces of attraction decreases. Hence, boiling point also decreases.

- 135. (b) $CH_{3}CH_{2}CH_{2}CH_{3} \xrightarrow{\text{Anhyd.} AlCl_{3}} CH_{3} CH_{3} CH_{3}$ $CH_{3} \xrightarrow{\text{iso butane}} CH_{3} \xrightarrow{\text{iso butane}}$
- **136.** (c) $CH_3 CH_2 Br + KOH \xrightarrow{\text{Dehydrohal ogenation}}$ $CH_2 = CH_2 + KBr + H_2O$

In alcoholic *KOH* alkoxide ions (RO^-) are present which is a strong base. They abstract proton from β -carbon of alkyl halide and favours elimination reaction

$$\begin{array}{c} ROH + KOH \rightarrow ROK + H_2O \\ \text{Alcohol} \end{array} \rightarrow \begin{array}{c} ROK + H_2O \\ \text{Potassium alkoxide} \end{array}$$

$$ROK \rightarrow RO^- + K^+$$
Alkoxideion

$$RO^- + H - CH_2 - CH_2 - Br \rightarrow ROH + CH_2 = CH_2 + Br$$

- **137.** (a) $Al_4C_3 + 6H_2O \rightarrow 3CH_4 + 2Al_2O_3$ Methane
- **138.** (c) In C_2H_6 , C-C bond length is 1.54Å.
- 139. (a) $R COOK + 2H_2O \xrightarrow{\text{Electrolysis}} R R + CO_2 + 2KOH + H_2$ Alkane
- **142.** (b) $CH_3COONa + NaOH \xrightarrow{CaO} CH_4 + Na_2CO_3$
- 143. (a)
 60
 90
 Cyclopropan
 Cyclopentane
 Cyclobutane

Cyclopropane with a bond angle of 60° is very strained and hence very reactive.

- 145. (b) Marsh gas mainly consists of methane.
- **146.** (c) $CH_3 Cl + 2H \xrightarrow{Zn/HCl} CH_4 + HCl$
- **149.** (d) Propane is a saturated compound. Ozonide is formed only by alkenes or alkynes

$$CH_2 = CH_2 + O_3 \rightarrow \begin{array}{c|c} CH_2 & CH_2 \\ \hline CH_2 & O & O \\ \hline \\ \hline CH_2 & CH_2 \\ \hline \\ \hline \\ CH_2 & CH_2 \\ \hline \\ \hline \\ CH_2 & CH_2 \\ \hline \\ CH_2 & CH_$$

$$CH \equiv CH + O_3 \rightarrow \begin{array}{c} O \\ CH - CH \\ O - O \\ Ethyne ozonide \end{array}$$

150. (b)
$$H - \overset{sp^2}{C} = \overset{sp^2}{C} - H$$
 sp^2 -hybridisation (trigonal H H

planar).

151. (a)
$$Ag - C \equiv C - Ag + 2HCl \rightarrow CH \equiv CH + 2AgCl$$

152. (c) Wurtz reaction
$$C_2H_5 - I + 2Na + I - C_2H_5 \xrightarrow{\text{Dry}} C_2H_5 - C_2H_5 + 2NaI$$
Rutane

154. (b) All the
$$C-C$$
 bond are single bonds. Hence sp^3 -hybridization and tetrahedral structure.

155. (a.b)
$$CH_3MgI + CH_3 - CH_2 - NH_2 \rightarrow$$

$$CH_4 + CH_3CH_2NHMgI$$

$$C\!H_3MgI + C_2H_5O\!H \rightarrow C\!H_4 + C_2H_5O\!MgI$$

Alkyl group of Grignard's reagent is involved in the formation of alkane.

159. (a) General formula of alkane $C_n H_{2n+2}$ (n = no. of atoms).

160. (c)
$$CH_3Br + H_2 \xrightarrow{LiAlH_4} CH_4$$
 (methane)
$$\xrightarrow{Na} CH_3 - CH_3$$
 (Ethane)

$$\xrightarrow{Na}$$
 $CH_3 - CH_3$ (Ethane)

161. (e) Photochemical chlorination of alkane take place by free radical mechanism which are possible by Homolysis of C - C bond

$$\begin{aligned} &Cl_2 \overset{hv}{\longrightarrow} Cl^{\bullet} + Cl^{\bullet} \\ &CH_3 - CH_3 + Cl^{\bullet} \to CH_3Cl^{\bullet} + \dot{C}H_3 \end{aligned}$$

163. (a) Producer gas –
$$CO$$
 and N_2

- 164. (c) Among alkanes, boiling point increase with increasing molecular weight. For isomeric alkanes straight chain alkanes have higher boiling point than the branched alkanes.
- 165. (b) Graphite is a good conductor of heat of electricity.
- 166. (d) Among the isomeric alkanes, the normal isomer has a higher boiling point than the branched chain isomer. The greater the branching of the chain, the lower is the boiling point. The n-alkane have larger surface area in comparison to branched chain isomer (as the shape approaches that of a sphere in the branched chain isomers). Thus, intermolecular forces are weaker in branched chain isomers, there fore they have lower point in comparision to straight chain isomers.
- 167. (d) The octane numbers of Fuel can be improved by increasing the percentage of branched alkanes, alkenes and hydrocarbon. Thus octane number can be by isomerisation (reforming), alkylation and aromatisation (cyclisation) etc.
- 168. (c) The approximate composition of gasoline is $C_6 - C_{11}$ at boiling point 70-200°C and is used in motor fuel, dry cleaning, petrol gas etc.

169. (c)
$$CH_4 + O_2 \rightarrow CO_2 + 2H_2O$$

170. (c) Straight chain olefins has highest knocking.

171. (a)
$$Al_4C_3 + 12H_2O \rightarrow 3CH_4 + 4Al(OH)_3$$

 $CaC_2 + 2H_2O \rightarrow C_2H_2 + Ca(OH)_2$

172. (d) Except 2,2 dimethyl butane rest compound contain 5 carbon i.e., pantane while 2,2 dimethyl butane contain 6 carbon i.e., Hexane

$$CH_3 - CH_2 - C - CH_3$$

$$CH_3 - CH_2 - C - CH_3$$

$$CH_3$$

173. (b)
$$H_3C - CH - CH_2 - CH_3 + Br_2 \longrightarrow CH_3$$

 $CH_3 CH_3$

$$Br$$
 $|$
 $H_3C - C - CH_2 - CH_3$
 $|$
 CH_3

Moior

174. (b)
$$H_3C - C - C - CH_3 \xrightarrow{Cl_2} H H$$

175. (c)
$$C_2H_5Cl + H_2 \xrightarrow{Pd/C} C_2H_6 + HCl$$

This reaction is used for the preparation of pure alkanes.

Alkene

2. (b)
$$CH_2 - CH_2 + Zn \rightarrow CH_2 = CH_2 + Br_2$$

| Alkene
 Br Br

4. (b)
$$CH_2 = CH_2 \xrightarrow{HOCl} CH_2 - CH_2 \xrightarrow{aq \ NaHCO_3} CH_2 - OH$$

$$CH_2 - OH$$

$$CH_2 - OH$$





- (a) Alkenes are unsaturated hydrocarbon having 5. double bond so generally gives addition
- 6. (d) According to markownikoff's rule.

7.

(a)
$$CH_{3} - CH_{2} - OH \xrightarrow{Conc. H_{2}SO_{4}} CH_{2} = CH_{2} + H_{2}O$$

8. (d)
$$CH_3$$
 $C = C$ CH_3 CH_3

- (d) $C_2H_5I + alc.KOH \rightarrow C_2H_4 + KI + H_2O$ 10.
- (a) $CH_3 CH_2 Br + KOH \rightarrow CH_2 = CH_2 + KBr + H_2O$ 12.
- (a) Methane can not be obtained by Sabatier and 13. Sendern's reaction because in this the product obtained contain minimum two carbon atoms.

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

$$CH = CH + 2H_2 \xrightarrow{Ni} CH_3 - CH_3$$

14. (c)
$$CH_3 - CH - C - CH_3 \xrightarrow{H_2SO_4} OH CH_3$$

$$CH_{3}$$

$$CH_{3} - \overset{+}{C}H - \overset{-}{C} - CH_{3} + OH^{-} \xrightarrow{\text{Methyl shift}} CH_{3}$$

$$CH_{3}$$

$$2^{\circ} \text{ carbonium ion}$$

20. (c)
$$CH_3 - CH = CH - CH_3 + HBr \xrightarrow{\text{Peroxide}}$$

$$CH_3 - CH_2 - CH - CH_3$$

$$Br$$
2-Bromobutan e

Anti-markownikoff's rule is not applicable to symmetrical alkenes.

21. (b)
$$CH_2 = CH_2 + KMnO_4 \xrightarrow{[O]} CH_2 - CH_2 + MnO_2$$
alk. $OH OH$
Colour less

$$CH = CH + KMnO_4 \xrightarrow{[O]} COOH + MnO_2$$

$$COOH \\ Oxalic acid$$
Brown

- (d) Octane number increases in the order Straight chain alkanes < Branched chain alkanes < Olefins < Cyclo alkanes < Aromatic Since, straight chain alkane has minimum
 - octane number. Hence, it produces maximum knocking.

23. (c)
$$R-CH = CH - R \xrightarrow{\text{dil. aqueous } KMnO_4} R - CH - CH - R$$
room temp.
 $OH OH$
(Alcohol)

$$R - CH = CH - R \xrightarrow{\text{Conc. } KMnO_4} R - COOH + R - COOH$$

24. (b)
$$CH_2 = C - CH_2 - CH_3 + H_2O \xrightarrow{H_2SO_4}$$

$$CH_2 = C - CH_2 - CH_3 + H_2O \xrightarrow{Markowniko ffs rule}$$

$$CH_3$$

$$CH_3$$

$$CH_3$$

$$CH_3$$

$$CH_3$$

$$CH_2 = C - CH_2 - CH_3 + H_2O \xrightarrow{Markowniko ffs rule}$$

$$CH_3 \\ CH_3 - C - CH_2 - CH_3 \\ OH \\ 2 - Methyl-2-butanol$$

27. (b)
$$CH_3 - CH = CH_2 + H_2O \xrightarrow{\text{Conc. } H_2SO_4} CH_3 - CH - CH_3$$

Analowniko ffs rule

OH

 $CH_3 - CH - CH_2 - CH_3 \xrightarrow{alc.KOH} CH_3CH = CHCH_3$

The reaction is dehydrohalogenation.

34. (a)
$$CH_2 = CH_2 + O_3 \longrightarrow CH_2 CH_2 \longrightarrow 2HCHO$$

$$O \longrightarrow O$$

(c) Oil are unsaturated esters which are converted into fats by saturating it by catalytic hydrogenation.

36. (c)
$$>C = C < \xrightarrow{\text{Hydrogenation}} > C - C < \xrightarrow{H}$$

37. (a)
$$CH_2$$

$$CH_2 - CH_2$$
Cyclo Propans

32. (d)

38. (b) e.g.
$$CH_2 = CH_2 + Br_2 \rightarrow CH_2 - CH_2$$
 $| | | Br Br$

Half of the double bond is broken. It means π bond is broken while sigma bond is retained also two new C-Br bonds are formed.

40. (b)
$$CH_3 - CH - CH_2 - CH_3 + KOH \rightarrow \text{(alc)}$$

$$CH_3 - CH = CH - CH_3 + KBr + H_2O$$
Buttener-2

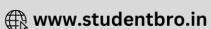
cis 1, 2-cyclopentanediol

(a) We know that greater the number of alkyl 42. groups attached to double bonded carbon atoms, more stable is the alkene. Therefore most stable is $R_2C = CR_2$

$$\begin{array}{ccc}
R & & & R \\
R & & & C = C \\
R & & & R
\end{array}$$







43. (c)
$$CH_2 = CH_2 + 2[O] \xrightarrow{KMnO_4} HCHO + HCHO$$
 Formaldehy de

- **44.** (c) Paraffins are non-polar compounds. The intermolecular forces are weak Vander Waal's forces. As the molecular mass increases Vander Waal's forces increases. Hence boiling point increases.
- **45.** (a) $CH_2 CH_2 + Zn \rightarrow ZnBr_2 + CH_2 = CH_2$ Br Br
- **46.** (c) $CH_3CH_2I + KOH(alc) \rightarrow CH_2 = CH_2 + KI + H_2O$ $CH_2 = CH_2 + KMnO_4 \xrightarrow{H_2O} CH_2 CH_2 + MnO_2$ Alk.Pink OH OHColourless

47. (d)
$$CH_2 = CH - Br \xrightarrow{HBr} CH_3 - CH < \frac{Br}{Br}$$

According to Markownikoff's rule H atom or positive part goes to that carbon atom which is more hydrogenated.

48. (d)
$$CH_3 - CH_2 - Br + KOH \rightarrow CH_2 = CH_2 + KBr + H_2O$$

49. (a)
$$CH_3 - CH_2 - OH \xrightarrow{\text{Conc. } H_2SO_4} CH_2 = CH_2 + H_2O$$

50. (c) Electrophillic addition reactions are shown by alkenes or alkynes in these reactions attacking species is electrophiles

$$CH_3 - CH = CH_2 + H^+ \xrightarrow{\text{Slow}} CH_3 - \overset{+}{CH} - CH_3$$

$$2^{\sigma} \text{ carbonium ion}$$

$$CH_3 - CH - CH_3 + Br^- \xrightarrow{\text{Fast}} CH_3 - CH - CH_3$$

$$Br$$
2-Bromo propane

51. (a)
$$CH_3CH_2CH_2Cl + KOH \rightarrow$$
 $CH_3 - CH = CH_2 + KCl + H_2O$ $CH_3 - CH - CH_3 + KOH \rightarrow CH_3 - CH = CH_2 + KCl + H_2O$

53. (c) Ethane and Methane does not decolourise $KMnO_4$ and does not react with ammonical cuprous chloride. Acetylene decolourise $KMnO_4$ solution and also gives red ppt. with ammonical cuprous chloride. On the other hand ethene decolourize $KMnO_4$ solution but does not react with ammonical cuprous chloride.

54. (a)
$$CH_2 = CH_2 + HOCl \xrightarrow{-} CH_2 - CH_2$$
 $OH Cl$
1-Chloro-2-hydroxy ethane

55. (b)
$$n(CH_2 = CH_2) \xrightarrow{400^{\circ} C} (-CH_2 - CH_2 -)_n$$
(ethylene) (polyethylene)

Trans-2-butene is more stable than its *cis* isomer.

58. (a)
$$CICH_2 - CH_2 - CH_2 - CH_3 \xrightarrow{alc. KOH}$$

$$CH_2 = CH - CH_2 - CH_3$$
1-butene

59. (d)
$$CH_3CH_2CH = CHCH_3 \xrightarrow{O_3} CH_3CH_2CHO + CHOCH_3$$
Ethanal

61. (c)
$$CH_3 = C = CH_2$$

$$sp^2 \qquad sp \qquad sp^2$$
Propadiene

Propadiene has both sp and sp^2 -hybridized carbon atoms.

62. (a)
$$C_2H_4 + Cl_2 \xrightarrow{CCl_4} Cl - CH_2 - CH_2 - Cl$$
 Ethelene Chloride

64. (b) Paraffins or alkanes are non-polar compounds. Hence soluble in benzene.

65. (b)
$$CH_3 - CH = CH_2 + HCl \xrightarrow{Peroxide} CH_3 - CH - CH_3$$

Peroxide rule is applicable only to HBr.

66. (d)
$$CH_3 - CH = CH_2 + HI \xrightarrow{\text{Peroxide}} CH_3 - CH_2 - CH_2I$$

69. (c)
$$CH_3 - CH - CH - CH_2 \xrightarrow{alc. NaOH} -HCl$$

$$H \quad Cl \quad H$$

$$CH_3CH = CH - CH_3$$

73. (c) $CH_3 - CH = CH - CH_2 - CH_3$ it decolourizes KMnO_4 solution because double bond is present.

77. (a)
$$CH_2 = CH - CH_2 - C \equiv CH + Br_2 \rightarrow CH_2 = CH - CH_2 - C = CH_2 - C =$$

78. (d)
$$CH_3COCH_3 + PCl_5 \rightarrow CH_3 - CCl_2 - CH_3 + POCl_3$$

80. (b)
$$3R - CH = CH_2 + \frac{1}{2}B_2H_6 \xrightarrow{\text{Dry}} (R - CH_2CH_2)_3 B$$

$$OH^-_{H_2O} \downarrow H_2O_2$$

$$3R - CH_2CH_2OH$$
Alcohol

83. (c)
$$CH$$
 Cl $Anhydrous AlCl_3$ $CH + AsCl_2$ Arsenic trichloriæ $CHCl$

Lewisite is more poisonous than mustard gas and was used in world war -II.







84. (a)
$$CH_3 - CH = CH_2 + Cl_2 \xrightarrow{500^{\circ} C}$$
 $CH_2 - CH = CH_2 + HCl$

This reaction is called allylic halogenation reaction because halogenation occurs at the allylic position of an alkene

85. (c)
$$n(CH_2 = CH - Cl) \xrightarrow{\text{Polymerization}} (-CH_2 - CH - -)_n$$

$$Cl$$
(PVC)

87. (b)
$$CH_3 - CH = CH_2 + HBr \xrightarrow{1-\text{bromo } -2-\text{chloroethane}}$$

$$CH_3 - CH - CH_3$$

$$Br$$
2-Bromopropa ne

88. (a)
$$CH_3 - CH = CH_2 + HBr \xrightarrow{Peroxide}$$

$$CH_3 - CH_2 - CH_2 - Br$$
89. (c) $CH_3 - C = CH - CH_3 \xrightarrow{(1)O_2 \atop (2)Z_1/H_2O}$

$$CH_3$$

69. (c)
$$CH_3 - C = CH - CH_3 \xrightarrow{(2)Zn/H_2O}$$

$$CH_3$$

$$CH_3COCH_3 + CHOCH_3$$

(a) $CH_3 - CH = CH_2 + HBr \xrightarrow{\text{Peroxide}}$

90. (c)
$$R-CH = C-R \xrightarrow{(1)O_3} R-CHO + R_2CO$$
 R

R

(c) $R-CH = C-R \xrightarrow{(1)O_3} R-CHO + R_2CO$
Ketone

92. (c)
$$CH_3 - CH_2 - OH \xrightarrow{\text{Conc.}} CH_2 = CH_2 + H_2O$$

93. (b)
$$CH_3 - CH_2 - Cl \xrightarrow{\text{alc. KOH}} CH_2 = CH_2 + KCl + H_2O$$
Ethyl Chloride

(b) Olefin because double bond is present. 94.

95. (b) In
$$CH_2 = CH_2$$
 double bond consist of one σ and one π bond

96. (b)
$$R - CH_2 - CH_2 - X \xrightarrow{\text{Elimination}}$$
 alc. KOH

$$R - CH = CH_2 + HX$$
Alkene Hallogen

97. (a)
$$CH_3 - CH_2 - CH_2 - CH_3 \xrightarrow{H_2SO_4}$$

$$CH_3 - CH = CH - CH_3$$
More symmetrica l(major product)
$$CH_3 - CH_2 - CH_2 - CH_3 \xrightarrow{H_2SO_4}$$

$$CH_2 = CH - CH_2 - CH_3$$
Less symmetrica l or

It is based on Saytzeff's rule. According to this more symmetrical or more alkylated alkene predominates.

98. (b)
$$CH_3 - CH_2 - CH_2 - OH \xrightarrow{PCl_3}$$

$$CH_3 - CH_2 - CH_2 - Cl \xrightarrow{\text{Alc.}KOH} CH_3 - CH = CH_2$$
(a) (b)
propylchloride Propene

99. (b)
$$CH_3 - CH_2 - CH_2 - Br + KOH \xrightarrow{C_2H_5OH}$$
 n -Propyl bromide

$$CH_3 - CH = CH_2 + KBr + H_2O$$
Propene

100. (b)
$$CH_3 - \overset{|}{C} - CH_2 - Br + KOH \longrightarrow CH_3$$

$$CH_3$$

$$CH_3 - \overset{|}{C} = CH - CH_3 + KBr + H_2O$$

In this reaction 1° carbonium ion is formed which rearranges to form 3° carbonium ion from which base obstruct proton. Hence 2methyl-2-butene is formed as a main product.

$$CH_{3} \xrightarrow{C} CH_{2} \xrightarrow{Methyl \, shift} CH_{3} \xrightarrow{C} CH_{2} - CH_{2}$$

$$CH_{3} \xrightarrow{C} CH_{3}$$

$$CH_{3} \xrightarrow{C} CH_{3}$$

$$CH_{3} \xrightarrow{C} CH_{2} - CH_{3}$$

$$CH_{3} \xrightarrow{C} CH_{3}$$

$$CH_{3} \xrightarrow{C} CH_{3}$$

$$CH_{3} \xrightarrow{C} CH_{3} \xrightarrow{C} CH_{3}$$

Elimination of proton from
$$\beta$$

$$CH_3 \quad \text{carbon which is less} \quad | \quad \text{hydrogenated} \quad \rightarrow$$

$$CH_3 - C = CH - CH_3$$
2-Methyl-2-Butene

101. (b) Alkyne > Alkene > Alkane

102. (c)
$$H \subset C = C \subset H$$

Ethylene sp^2 -hybridization; Shape = Planar.

104. (b)
$$CH_2 = CH - Cl \xrightarrow{HCl} CH_3 - CH < \stackrel{Cl}{Cl}$$

According to Markownikoff's rule H atom of the reagent goes to that carbon atom which is more hydrogenated.

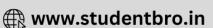
105. (b)
$$CH_3 - CH = CH_2 + HBr \xrightarrow{\text{Markownikoffs rule}} \rightarrow CH_3 - CH - CH_3 + HBr \xrightarrow{\text{Presence of peroxid e}} \rightarrow Br$$
2-Bromopropa ne

$$CH_3 - CH_2 - CH_2 - Br$$
1-Bromopropa ne

106. (a) Peroxide rule is applicable only to *HBr* and not for HCl, HF and HI.

unsymmetri cal (minor pro duct)





109. (a)
$$n \left(CH_2 = C - CH = CH_2 \right) \xrightarrow{\text{Polymerization}} Cl$$
Chloroproene

$$\begin{pmatrix} -CH_2 - C = CH - CH_2 - \\ Cl \\ Neoprene \end{pmatrix}_n$$
 120. (b) C_3H_6 is an alkene theref alkaline $KMnO_4$ solution.

123. (c) $CH_3 - C = C - CH_3 \xrightarrow{(1)O_3} \xrightarrow{(2)Zn/H_2O}$

110. (c)
$$CH_3 - CH_3 - CH_2 - Mg - Br + C_2H_5OH \longrightarrow CH_3$$
 $CH_3 - CH_3 - CH_3 + C_2H_5OMgBr$

$$CH_3 - CH - CH_3 + C_2H_5OMgBr$$

Ethoxy magnesium

 CH_3

Isobutane

111. (c)
$$CH_3 - CH_2 - C = CH - CH_3 + HOCl \rightarrow CH_3$$

$$CH_3 - CH_2 - CH_2 - CH_3$$

$$CH_3 - CH_2 - CH_3$$

Addition takes place according to

Markownikoff's rule in which Cl^+ goes to that carbon atom which is more hydrogenated.

112. (a) In case of ethene double bond is present. Hence, addition reactions occur easily.

113. (a)
$$CH_3 - CH = CH_2 + Br_2 \rightarrow CH_3 - CH_2 - CH_2$$
Propane
1 mole
1 mole
42 gms
1,2-dibromo propane

: 42 gms of propene reacts with 160 gms of bromine.

$$\therefore$$
 21gms of propene $\frac{160}{42} \times 21 = 80 \text{ gms}$.

114. (b) Butadiene $CH_2 = CH - CH = CH_2$

A single bond separated by two double bonds is known as conjugated double bond.

115. (d)
$$CH_3 - C - OH \xrightarrow{Cu} CH_3 - C + H_2O$$

$$CH_3 - C - OH \xrightarrow{300^{\circ}C} CH_3 - C + H_2O$$

$$CH_3 - C + H_2O$$
Isobutene

117. (a) Cyclic hydrocarbon in which all the carbon atoms are present in the same plane is benzene. In this C-C bond length is 1.39Å which is more than 1.34A° but less than 1.54Å. angle is 120° with sp^2 Hence bond hybridization.

119. (a)
$$CH_3 - CH_3 - CH_4 = CH_2 - \frac{DH_3(OAc)_2; H_2O}{ii) NaBH_4} \rightarrow CH_3$$

$$CH_3 - CH_3 - CH - CH_3$$

$$CH_3 - CH - CH_3$$

$$CH_3 OH$$

120. (b) C_3H_6 is an alkene therefore decolourizes

123. (c)
$$CH_3 - C = C - CH_3 \xrightarrow{(1)O_3} \xrightarrow{(2)Zn/H_2O} CH_2 CH_2$$

$$CH_3 - CO + OC - CH_3$$
 $CH_3 CH_3$
Acetone

124. (b)
$$CH_3 - CH = CH - CHO \xrightarrow{Acidic} KMnO_4$$

$$CH_3 - CH = CH - COOH$$

127. (b)
$$CH_3 - CH_2 - CH = CH_2 \xrightarrow{HBr}$$
 $CH_3 - CH_2 - CH - CH_3$

$$Br$$
2-Bromo butane

$$CH_3-CH_2-CH_2-CH_3$$

$$CH_3-CH_3$$

$$CH_3$$

$$C$$

129. (a) Ozonolysis is useful in locating the position of a double bond in an alkene. The double bond is obtained by joining the carbon atoms of the two carbonyl compounds. For example Let the product of ozonolysis be two molecules of ethanal.

$$\begin{array}{ccc} H & H \\ | & | \\ CH_3 - C = O + O = C - CH_3 \rightarrow CH_3 - CH = CH - CH_3 \\ \hline & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ \end{array}$$

131. (c)
$$CH_3 - CH - CH_2 - CH_3 \rightarrow CH_3 - CH = CH - CH_3 + HBr$$

$$Br$$
2-Bromo butane

134. (b)
$$CH_3 - C - CH_3 \xrightarrow[]{H_2SO_4} CH_3 - C = CH_2 + H_2O$$
OH
2Methyl-2-hydroxy propane Isobutene

2Methyl-2-hydroxy propane
135. (c)
$$CH_2 = CH_2 + H_2 \xrightarrow{Ni} CH_3 - CH_3$$

Ethene CH_2COOK

136. (c)
$$\stackrel{CH_2COOK}{\vdash} + 2H_2O \xrightarrow{\text{Electrolysis}} CH_2COOK + 2H_2O \xrightarrow{\text{Electrolysis}} CH_2 + 2CO_2 + 2KOH + H_2$$
 $\stackrel{\parallel}{\vdash} CH_2 \qquad Cathode$

137. (c)
$$CH_2 = CH_2 + H_2 \frac{Ni}{300^{\circ}C} CH_3 - CH_3$$

Ethylene does not give precipite

Ethylene does not give precipitate with ammonical silver nitrate solution because it does not have acidic hydrogen.





139. (a)

143. (d) Pd acts as catalyst.

144. (a)
$$CH - CH = CH_2 + HBr \xrightarrow{\text{peroxide}}$$

$$CH_3 - CH_2 - CH_2 - Br$$

$$n - \text{propyl bromide}$$

In presence of peroxide addition takes place according to Anti Markownikoff's rule and hydrogen atom goes to that carbon atom which is less hydrogenated.

145. (d)
$$CH_3 - CH_2 - OH + H_2SO_4 \xrightarrow{110^o C}$$
 Ethanol Conc.

CH₃CH₂HSO₄ Ethylhydrogen sulphate

146. (a)
$$CH_2 = CH_2 \xrightarrow{1)O_3} HCHO + HCHO$$
 Formaldehy de

147. (d) C_2H_4, C_3H_6 and C_4H_8 all an alkene. Therefore they discharge the red colour of bromine.

149. (a)
$$HI \rightarrow H^+ + I^-$$

$$CH_3 - CH = CH_2 + H^+ \rightarrow$$

$$CH_3 - CH_2 - \overset{+}{C}H_2 + CH_3 - \overset{+}{C}H - CH_3$$
 (Major) (Major) (More stable)
$$\overset{1^{\circ}}{\text{Carbonium ion}} (\text{Less stable})$$
 (More stable)

$$CH_3 - CH - CH_3 + I^- \rightarrow CH_3 - CH - CH_3$$

$$2^o \text{ Carbonium ion } I$$
Isopropyliodide (Major product)

150. (a)
$$CH_3 - CH_2 - CH = CH_2 + Br_2 \rightarrow CH_3 - CH_2 - CH - CH_2$$

$$\begin{vmatrix} & & & & & & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & & \\ & & & & & & & & & \\ & & & & & & & & & \\ & & & & & & & & & \\ & & & & & & & & \\ & & & & & & & & \\ & & & & & & & \\ & & & & & & & \\ & & & & & & & \\ & & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & \\ & & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & \\ & & & \\ & & \\ & & & \\ & & & \\ & & \\ & & & \\ & \\ & & \\ & & \\ & & \\ & \\ & \\ & \\ & \\$$

151. (a)
$$CH_2 = CH_2 \xrightarrow{(1)O_3} HCHO + HCHO$$

153. (c)
$$(CH_3)_2 - C = CH \xrightarrow{\text{Catalytic} \\ \text{hydro genation}} (CH_3)_2 - CH - CH_2$$

154. (b)
$$CH_3 - C = CH_2 + HBr \xrightarrow{\text{Peroxide}} CH_3 - \overset{H}{C} - CH_2 - Br$$

$$CH_3 \xrightarrow{C} CH_3$$

$$2 - \text{isobutylbromide}$$

155. (a)
$$CH_2 = CH - C = CH$$

156. (c)
$$CH_3 - CH = CH_2 + H^+Br^- \rightarrow CH_3 - CH - CH_3$$

Br 2-Bromopropa ne

$$C_2H_5OH \xrightarrow{\text{Conc. } H_2SO_4} CH_2 = CH_2$$

158. (d) $CH_3 - CH = CH - CH_2 - CH_3$ will be the most stable because greater the number of alkyl groups attached to double bonded carbon atoms, more stable is the alkene.

159. (b) Markownikoff's rule can not be applied for symmetrical alkene.

160. (c)
$$CH_3 - C = CH_2 \xrightarrow{H_3O^+} CH_3 - C - CH_3$$

161. (d) Solution of bromine in carbon tetrachloride is used to test for unsaturation of alkene. Red colour of bromine disappears due to the formation of colourless dibromo ethane $(C_2H_4Br_2)$.

$$CH_2 = CH - CH_3 + H_2O \xrightarrow{H_2So_4} \rightarrow$$
Propylene
$$OH$$

$$|$$

$$CH_3 - CH - CH_3$$
Isopropyl alcohol

Thus in this reaction isopropyl alcohol is formed.

163. (b) Propene gives formaldehyde and acetaldehyde on ozonolysis.

$$CH_{2} = CH.CH_{3} \xrightarrow{O_{3}} CH_{2} - CH.CH_{3} - \underbrace{\frac{Z_{1}/H_{2}O}{H_{2}O_{2}}} HCHO + CH_{3}CHO$$

164. (a) When ethylene is treated with cold alkaline *KMnO*4 , ethylene glycol is formed.

KMnO4, ethylette grycol is formed.

$$\frac{H}{H} > C = C < \frac{H}{H} \xrightarrow{\text{KMnO4}} \frac{H}{\text{Cold alkaline}} + \frac{C - C}{H} < \frac{H}{OH \ OH}$$
Ethylene glycol

165. (a)
$$C_6H_6 + H_2C = CH_2 \xrightarrow{AlCl_3.Hl} C_6H_5CH_2CH_3$$

166. (a) Markownikoff as well as anti-Markownikoff's rule is valid only for unsymmetrical alkenes.

167. (b) The formation of *n*-propyl bromide in presence of peroxide can be explained as follows.

Step-1: Peroxide undergo fission to give free radicals $R-O-O-R \rightarrow 2-R-O$

Step-2: HBr combines with free radical to form bromine free radical $R - \dot{O} + HBr \rightarrow R - OH + B\dot{r}$

Step-3 : $B\dot{r}$ attacks the double bond of the alkene to form a more stable free radical

$$CH_{3} - \dot{C}H - CH_{2}Br$$

$$(more stable)$$

$$CH_{3}CH = CH_{2} + Br$$



$$Br$$
 $|$
 $CH_3 - CH - CH_2$
(less stable)

Step-4 : More stable free radical attacks the ${\it HBr}$

$$CH_3$$
 – CH – CH_2 – Br + HBr \rightarrow $CH_3CH_2CH_2Br$ + Br n-propyl bromide

Step-5: $B\dot{r} + B\dot{r} \rightarrow Br_2$

168. (c)
$$CH_2 = CH_2 \xrightarrow{KMnO_4} CH_2 - CH_2$$

$$OH OH$$

$$AgNO_3$$
No

169. (b) According to Markownikoff's rule, the negative part of the unsymmetrical reagent adds to less hydrogenated (more substituted) carbon atom of the double bond.

$$C_6H_5CH = CH - CH_3 + HBr \rightarrow C_6H_5CHCH_2CH_3$$

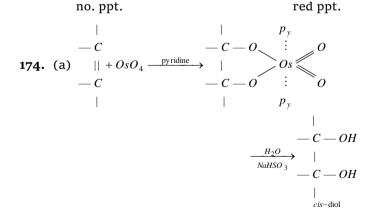
171. (a)
$$CH_2 CH_3 CH_3 CH_3 CH_3 CH_2 CH_2 CH_2 HSO_4 CH_2OH CH_2OH$$

Except ethyl alcohol, no other primary alcohol can be prepared by this method as the addition of H_2SO_4 follows Markownikoff's rule. Generally secondary and tertiary alcohols are obtained.

172. (c)
$$CH_3 - C = CH_2 \xrightarrow{HBr / peroxide} CH_3 - CHCH_2 - Br$$

$$CH_3 \qquad CH_3 \qquad CH_3$$

173. (c) Decolourisation of KMnO_4 Ethene Acetylene $\mathit{Cu}_2\mathit{Cl}_2$, $\mathit{NH}_4\mathit{OH}$ $\mathit{Cu}_2\mathit{Cl}_2$,



Alkyne

1. (c) $2CH \equiv CH + 5O_2 \rightarrow 4CO_2 + 2H_2O \Delta H = -1300 \ KJ$ The combustion of acetylene is highly exothermic and the heat produced during the combustion can be used for welding purposes in the form of oxy acetylene flame.

2. (c)
$$CaC_2 + 2H_2O \rightarrow Ca(OH)_2 + C_2H_2$$

 $CH = CH + 2[Ag(NH_3)_2]OH \rightarrow$
 $Ag - C = C - Ag + 4NH_3 + 2H_2O$
Whitepart

3. (c)
$$CH_3 - CH_2 - C \equiv CH \xrightarrow{\text{Cold}} CH_3CH_2COOH + CO_2$$

4. (c)
$$CH_3 - COOH \xrightarrow{LiAlH_4} CH_3 - CH_2 - OH \xrightarrow{H^+} 443 K$$

$$CH_2 = CH_2 \xrightarrow{Br_2} CH_2 - CH_2 \xrightarrow{alc.} KOH$$

$$Br Br$$

$$CH \equiv CH + 2KBr + 2H_2O$$

Acetylene

5. (b)
$$C \equiv C < C = C < C - C$$

1.20 A^o 1.34 A^o 1.54 A^o

6. (a)
$$HC = C - CH = CH_2$$

7. (a)
$$3 CH \equiv CH \xrightarrow{\text{Re } d \text{ hot}} \text{Fe tube}$$

10. (c)
$$H_2$$

Cyclohexene Cyclohexane

$$CH_2$$

$$CH_2$$

Cyclohexene CH2

$$CH_2$$

Cyclohexene CH2

11. (a)
$$HC \equiv CH + AsCl_3 \longrightarrow ClHC = CH AsCl_2$$
2-chlorovinyl dichloroarsine (Lewsite)

(c)
$$CH_3 - C \equiv C - CH_2 - CH_3 \xrightarrow{O_3}$$

$$CH_3 - C \downarrow C - CH_2 - CH_3 \downarrow C - CH_2 - CH_3 \downarrow C - CH_3 - COOH + CH_3 CH_2 COOH$$

14. (a)
$$-C \equiv C - is$$
 most reactive because *sp*-hybridization.

16. (d)
$$CH = CH \xrightarrow{NaNH_2} CH = C^-Na^+ \xrightarrow{CH_3I} CH = C - CH_3$$

$$C_2H_5 - C = C - CH_3 \xleftarrow{C_2H_5I} Na^+C^- = C - CH_3$$
 Pent-2-yne

17. (d)
$$CH_3 - C = CH + H_2O \xrightarrow{H_2SO_4/H_8SO_4}$$



$$CH_3 - C - CH_3$$

$$O$$
Propanone
$$CH_3 - C - CH_3$$

$$O$$
Prop-1-ene 2-ol

18. (b)
$$CH = CH + HCl \rightarrow CH_2 = CH - Cl \xrightarrow{HCl}$$

$$CH_3 - CH < \frac{Cl}{Cl}$$

1-di-chloroethane

19. (c)
$$CaC_2 \xrightarrow{H_2O} HC \equiv CH \xrightarrow{Dil.H_2SO_4} CH_3CHO$$
Acetaldelyde

 $\xrightarrow{Ni} CH_3CH_2 - OH$ Ethylalcohol

20. (c)
$$R - CH_2 - CCl_2 - R \xrightarrow{KOH + C_2H_5OH}$$

$$R - C \equiv C - R + 2HCl$$

This reaction is an example of dehydrohalogenation Hence, alcoholic *KOH* is used as a reagent.

CH - COOK

21. (d) |||
$$+2H_2O \xrightarrow{\text{Electrolysis}}$$
 $CH - COOK$

cii coon

$$CH$$
||| + 2 CO_2 + 2 KOH + H_2
Cathode

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

 $CH_2 - CH_2 + 2KOH \rightarrow CH \equiv CH + 2KBr + 2H_2O$
 $\mid \qquad \mid$
 $Br \qquad Br$

22. (b)
$$CH - COOK + 2H_2O \xrightarrow{Electrolysis} CH + 2CO_2$$

$$||| CH - COOK CH$$

 $+2KOH + H_2$

$$CH_2 - COONa + 2H_2O \xrightarrow{Electrolysis} CH_2COONa$$

$$\begin{array}{l} CH_2 + 2\,CO_2 + 2\,NaOH + H_2 \\ \parallel \\ CH_2 \end{array}$$

23. (d)
$$CH_3 - C \equiv C - H + AgNO_3 \rightarrow CH_3 - C \equiv C - Ag$$
Propyne Annonical Siver salt of Propyne

24. (c)
$$CH = CH \xrightarrow{H_2O/Hg^{2+}} H_3C - C = H \xrightarrow{LiAlH_4} CH_3 - CH_2 - OH \xrightarrow{P_4/Br_2} C_2H_5Br$$

25. (c)
$$4 CH = CH \xrightarrow{Ni(CN)_2} Cyclo$$

26. (d) Ethyne have acidic hydrogen to form salt.

27. (a)

$$CH_{3} - CH_{2} - C \equiv CH + 2Br_{2} \rightarrow CH_{3} - CH_{2} - \begin{bmatrix} Br & Br \\ CH_{3} - CH_{2} - C \\ C - C - H_{3} \end{bmatrix}$$

$$Br = Br$$

$$Br = Br$$

Since the molecule takes 2 moles of Br_2 . Therefore it is alkyne. Also it gives white ppt with Tollen's reagent therefore acidic H is present. Hence it is 1-Butyne.

28. (d)
$$2CH = CH + 2Na \rightarrow 2CH = C^{-}Na^{+} + H_{2}$$

29. (a)
$$CH_3 = CH + AgNO_3 \rightarrow Ag - C = C - Ag$$
 Whiteppt.

$$CH_3 \equiv CH + CuCl_2 \rightarrow Cu - C \equiv C - Cu$$
(Ammonica) $\rightarrow Cu - C \equiv C - Cu$

32. (c)
$$sp - sp > sp^2 - sp^2 > sp^3 - sp^3$$

Order of bond strength

33. (c) Ethylene and Benzene

$$120^{\circ} 120^{\circ}$$

$$sp^2 sp^2$$

34. (a)
$$CH_3 - CH_2 - C \equiv CH \xrightarrow{H_2SO_4, 60^{\circ}C} CH_3 - CH_2 - C - CH_3$$

35. (a)
$$CH \equiv CH + NaNH_2 \rightarrow CH \equiv C^-Na^+ + NH_3$$

36. (a)
$$CH = CH + 2Br_2 \xrightarrow{CCI_4} CH - CH = CH + NH_4OH + AgNO_3 \rightarrow Ag - C = C - Ag$$

Disilver acety lide white ppt.

38. (d) C_2H_2 and C_3H_4 are homologue because they differ by $-CH_2$ group. Both have triple bond in their molecule.

39. (c)
$$CH = CH + HCl \xrightarrow{HgCl_2} CH_2 = CH - Cl$$
Vinylchloride

41. (a)
$$3CH_3 - C \equiv CH \longrightarrow \begin{array}{c|c} CH & CH_3 \\ \hline C & C \\ \hline C & C \\ \hline CH & CH \\ \hline CH & CH \\ \hline CH_3 \\ \hline CH_3 \\ \hline \end{array}$$

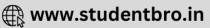
Mesitylene or 1,3,5-trimethyl benzene

(I) hydroxide

42. (c)
$$2NH_4OH + Cu_2Cl_2 \rightarrow 2CuOH + 2NH_4Cl$$

$$NH_4OH + CuOH \rightarrow [Cu(NH_3)_2]OH$$
 Diamnine copper





$$2[Cu(NH_3)_2]OH + HC \equiv CH \rightarrow$$

$$Cu - C \equiv C - Cu + 4NH_3 + 2H_2O$$
 copper acetylide Red ppt.

43. (a)
$$3CH \equiv CH \xrightarrow{\text{Hot } Cu \text{ tube}} AlCl_3$$

44. (c)
$$CH \xrightarrow{KMnO_4} COOH$$
 $CH \xrightarrow{CH} COOH$
Acetylene Oxalic acid

45. (b)
$$CH = CH + H_2O \xrightarrow{40\% H_2SO_4} CH_3 - CHO$$

- **46.** (c) Acetylene reacts with ammonical cuprous chloride to give brown *ppt* where as ethylene does not give this reaction.
- **47.** (a) $Ag(NH_3)_2^+$ gives white ppt with terminal alkynes.
- **50.** (b) Bond length decreases with increase in number of bonds.

55. (d)
$$3HC \equiv CH \xrightarrow{\text{Red}}$$
Benzen

56. (a) Reaction of acetic acid with acetylene is catalysed by Hg^{2+} salts.

$$HC \equiv CH \xrightarrow{CH_3COOH} CH_2 = CHOOCCH_3$$

$$\xrightarrow{CH_3COOH} CH_3 - CH_3COCCH_3)_2$$
ethylidene diacetate

Hg(OOCCH₃)₂ ethylidene diacetate
57. (c)
$$2C + H_2 \xrightarrow{Electricarc} CH \equiv CH$$

58. (c)
$$CH = CH \xrightarrow{40\% H_2SO_4} CH_2 = CH - OH$$
Vinylalcohol

$$\frac{\text{Rearrangem ent}}{\text{Acetaldehyde}} \rightarrow CH_3 - CHO$$

- **59.** (a) $2CH = CH + 2Na \rightarrow 2CH = C^-Na^+ + H_2$ Reaction with sodium metal shows that hydrogen is acidic.
- **63.** (d) In $CH \equiv CH$ triple bond consists of one σ and two π bonds.
- **65.** (a) Nucleophiles and alkalies normally do not react with acetylene. Thus *NaOH* does not react.

66. (c)
$$HC \equiv CH + 2HOCl \rightarrow [CHCl_2 - CH(OH)_2]$$

$$-H_2O$$

$$CHCl_2 - CHO$$
Dichloroacetal dehyde

69. (b)
$$CH_3 - CH = CH - C \equiv C - H$$
. Acidic hydrogen (*H* atom attached to triple bond) is present therefore it gives reaction with ammoniacal $AgNO_3$.

70. (b)
$$CH = CH + H_2O \xrightarrow{40\% H_2SO_4} CH_2 = CHOH \\ \downarrow \\ CH_3CHO$$

71. (b)
$$CH_3C \equiv CH + 2HBr \longrightarrow CH_3 - C - CH_3$$

$$Br$$

$$Br$$

$$CH_3C \equiv CH + 2HBr \longrightarrow CH_3 - C - CH_3$$

72. (c) $CH_3 - C \equiv C - CH_3$, 2-butyne does not have acidic hydrogen. Hence, does not give white ppt. with ammonical $AgNO_3$ solution.

73. (a)
$$(CH \equiv CH) \xrightarrow{O_3} CH - CH \xrightarrow{Hydrolysis} O \longrightarrow O$$

$$\begin{array}{c} CHO-CHO \xrightarrow{Zn} & CH_2-OH \\ \text{Glyoxal} & CH_3COOH & CH_2-OH \end{array}$$

77. (b)
$$| +2KOH \text{ (alcoholic)} \xrightarrow{\Delta} CH \equiv CH + 2KBr$$
 CH_2Br

 $+2H_{2}O$

- **78.** (d) All are alkenes but 2-butyne is not.
- **80.** (c) Reduction of alkynes with liquid NH_3/Li gives trans alkenes.
- **82.** (a) Ethyne reacts with ammonical $AgNO_3$ to give white ppt of silver acetylide while ethane and ethene do not react because acidic hydrogen is absent.
- **83.** (d) Acidic property because H atoms are replaced by Silver metal atoms.
- **87.** (c) The hydrogen atom which is attached to triple bond is acidic.
- **88.** (c) In acetylene both carbons are *sp*-hybridised so it has linear structure.

89. (d)
$$CaC_2 + 2H_2O \rightarrow CH \equiv CH + Ca(OH)_2$$
Acetylene

90. (c)
$$CH \equiv CH + HCN \xrightarrow{Ba(CN)_2} CH_2 = CH - C \equiv N$$
Vinyley anide

91. (c) Because $CH_3CH_2 - C \equiv CH$ has one acidic hydrogen.

92. (d)
$$CH = CH + 2Na \xrightarrow{Liq.NH_3} C.Na = C.Na + H_2$$

94. (b)
$$CH = CH + H_2O \xrightarrow{\text{dil.} H_2SO_4} CH_2 = C - OH \rightarrow CH_3 - C-$$

- **95.** (d) $CH_3 C \equiv C CH_3$ has not acidic character.
- **96.** (d) Addition $CH = CH + 3H_2 \xrightarrow{Ni} CH_3 CH_3$ Substitution –

$$CH \equiv CH + Na \longrightarrow CH \equiv C^{-}Na^{+} + \frac{1}{2}H_{2}$$

Polymerization -

$$3CH \equiv CH \xrightarrow{\text{hot } Cu \text{ tube}} C_6H_6$$
Polymerization Benzene





(d) Bond length decrease with increase in Bond 97. order. Hence triple bonded carbon has minimum bond length.

C - C bond length = 1.54 Å

C = C bond length = 1.33 Å

 $C \equiv C$ bond length = 1.22 Å

(b) Acetylene can be obtained by the reaction of 98. silver and chloroform (or iodo form)

 $2CHI_3 + 6Ag \xrightarrow{\Delta} C_2H_2 + 6AgI$

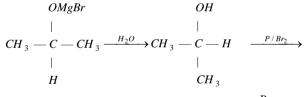
- (d) The partial reduction of alkynes by active 99. metal in liquid ammonia takes place through trans vinylic anion which ultimately produces trans alkene.
- **100.** (a) We know that C C bond length = 1.54 A, C = C bond length =1.34A and $C \equiv C$ bond length = 1.20A. Since propyne has triple bond; there fore it has minimum bond length.

101. (b) $CH \equiv CH + H_2 \xrightarrow{\text{Lindlar}} CH_2 = CH_2$ Cataly st Acety lene

- **102.** (a) $HC\sigma \equiv CH$ one sigma and two π bond
- 103. (a) Propyne reacts with ammoniacal $AgNO_3$ due to presence of acidic hydrogen atom. $CH_3C \equiv CH + AgNO_3 + NH_4OH \rightarrow$

 $CH_3C \equiv CAg + NH_4NO_3 + H_2O$

104. (a) $CH \equiv CH \xrightarrow{HgSO_4} CH_3 CHO \xrightarrow{CH_3 MgBr}$



105. (c) $Mg_2C_3 + 4H_2O \rightarrow CH_3C \equiv CH + 2Mg(OH)_2$

Aromatic Hydrocarbon

- $CO CH_3$ (d) Acetophenone
- (a) Coal tar is a main source of aromatic 5. hydrocarbons like benzene, naphthalene, anthracene, phenol etc

 $C - C\pi$ bonds = 3 $C - C\sigma$ bonds = 6 7. (c)

 $C - H\sigma$ bonds = 6

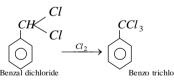
Hence 12σ and 3π bonds

(b) Ratio = $\frac{\sigma \text{ bonds}}{\pi \text{ bonds}} = \frac{12}{3} = 4$

(BHC)

- (a) $C_6H_6 + C_2H_5Cl \xrightarrow{AlCl_3} C_6H_5 C_2H_5 + HCl$
- (a) $C_6H_5CHC_6H_5$ is the most stable carbonium ion due to resonance.

o- and p-chlorotoluene



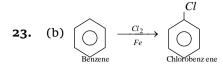
(b) $HO - NO_2 + H^+ HSO_4^- \rightarrow H_2O + NO_2^+ + HSO_4^-$

22. (a) $C_6H_6 + CH_3Cl \xrightarrow{\text{anhyd.}} C_6H_5CH_3 + HCl$, it is a

Friedel-craft's reaction.







$$CH - C$$

$$CH - C$$

$$CH - C$$

$$O$$
Maleic anhydride

29. (c)
$$+ NaOH$$
 CaO $+ Na_2CO_3$ Sodium benzoate Benzene

$$\begin{array}{c} NO_2 \\ \\ \bullet \\ \bullet \\ \bullet \\ \end{array}$$

31. (d)
$$\begin{array}{c} & \xrightarrow{>55^{\circ}C} \\ & & & \\ Nitrobenzene \end{array}$$

$$+3HNO_{3} \xrightarrow{Conc.H_{2}SO_{4}} \xrightarrow{NO_{2}} \\ & & \\ NO_{2} \xrightarrow{CH_{3}} NO_{2} \\ & & \\ NO_{2} \xrightarrow{NO_{2}} \\ & & \\ 2, 4, 6- trinitro \end{array}$$

 NO_2

32. (b)
$$+H_2SO_4 \rightarrow +H_2O_4$$

34. (c)
$$CH_3$$
 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3

m-xylene o-xylene p-xylene

m-xylene is most easily sulphonated because O and P both positions are free with respect to methyl group.

35. (d)
$$COOH$$
 A

Toluene Benzoic acid

36. (c)
$$+ \frac{9}{2}O_2 \xrightarrow{V_2O_5} | H - COOH \xrightarrow{-H_2O}$$
Maleic acid

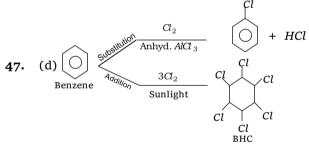
38. (b) Phenol is most easily nitrated.

41. (a)
$$\longleftrightarrow$$
 \longleftrightarrow \longleftrightarrow \longleftrightarrow \longleftrightarrow \longleftrightarrow

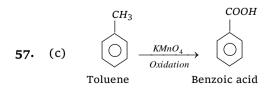
43. (a) Nitrobenzene is least reactive towards bromine because of presence of $-NO_2$ group decreases electron density at o and p positions and hence attack of electrophile on the benzene nucleus is difficult because + ve charge on o and p position repel the incoming electrophile.

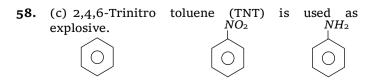
46. (d) $AlCl_3$ is an electron deficient compound. It generates electrophile in the reaction

$$CH_3Cl + AlCl_3 \rightarrow CH_3^+ + AlCl_4^-$$
Electrophile

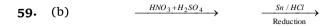


- **51.** (b) Benzene can be obtained by polymerisation of acetylene.
- 53. (b) Benzene C_6H_6 is made up of hydrogen and carbon only.
- **54.** (c) They have a relatively high percentage of carbon
- **55.** (c) All 6 carbons of Benzene are sp^2 -hybridised so it is planar.









61. (e) TNT is used as an explosive.

62. (c)
$$+Cl_2 \xrightarrow{AlCl_3}$$
 + $HCl_3 \xrightarrow{\text{Substitution}}$

65. (b)
$$+ HNO_3 \xrightarrow{H_2SO_4 conc.} NO_2$$

$$NO_2 \xrightarrow{NO_2}$$

$$NO_2 \xrightarrow{NO_2}$$
1,3-dinitrobenzene or m -

68. (c) *n*-heptane
$$\frac{Cr_2O_3 / Al_2O_3}{750 \text{ K}}$$

69. (b)
$$HNO_3$$
 H_2SO_4 O_2 O_2 O_2 O_3 O_4 O_2 O_3 O_4 O_5 O_5 O_6 O_7 O_8 $O_$

3-nitrochloro

- 71. (b) In benzene due to resonance all the carbon-carbon bond lengths are equal 1.39Å which is between C C(1.54 Å) and C = C(1.34 Å)
- 72. (c) $C_6H_5COONa + NaOH \xrightarrow{CaO} C_6H_6 + Na_2CO_3$ Sodium benzoate Benzene
- 73. (a) Friedel-Craft's reaction involves the introduction of an alkyl or acyl group into benzene ring in the presence of a catalyst. The presence of an electron withdrawing group in the ring hinders the reaction. Therefore phenyl acetanilide is not used.
- 74. (a) Halogenation is initiated by free redical.
- **76.** (b) In benzene all the six carbon atoms are present in the same plane. All the carbon atoms are sp^2 hybridized. Hence, it is a planar molecule.
- 77. (d) Basicity of amines is due to availability of an unshared pair (lone pair) of electrons on nitrogen. This lone pair of electrons is available for the formation of a new bond with a proton or Lewis acid.

 Pyridine is less basic than triethylamine because lone pair of nitrogen in pyridine is delocalised.

78. (a) In chlorination electrophile is Cl^+ (chloronium ions).

79. (c)
$$\bigcup_{\text{Ethyl benzoate}}^{COOC_2H_5}$$

-COOH group is meta directing group hence, ethyl benzoate undergo meta substitution.

$$COOC_2H_5$$
 $COOC_2H_5$ $+ Cl_2 \rightarrow Cl$ Ethyl benzoate $- chloro ethyl$

80. (d)2
$$O$$
 + 2HNO₃ C Conc. H_2SO_4 O + O + O + O + O 2

o- and P - nitro toluene

81. (d)
$$H-C=O$$
 $H-C-O^ H-C-O^ H-C-O^-$

Presence of -CHO decrease electron density at o to p positions. Hence, attack of an electrophile occurs at m-position. therefore it is meta directing group.

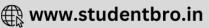
84. (b)
$$\longrightarrow$$
 $+3H_2 \xrightarrow{Ni}$ Cyclohexane

87. (b)
$$\begin{array}{c} Cl & Cl \\ +3Cl_2 & \xrightarrow{hv} & Cl & Cl \\ & & Cl & Cl \\ & & BHC \end{array}$$

- **88.** (a) Benzene on fractional distillation gives light oil [It is lighter than water and hence called as light oil]
- **89.** (d) Because Ammonical $AgNO_3$ reduce C = CH bond of yne.
- 90. (a) Bond length $\propto \frac{1}{\text{Bond order}}$ Bond order = $\frac{\text{Bonding} \text{antibondin g electron}}{2}$

Bond order is highest for C_2H_6 so it has minimum bond strength.





91. (b)
$$C_6H_5COOH + NaOH \rightarrow C_6H_5COONa + H_2O$$

$$\xrightarrow{NaOH} C_6H_6 + Na_2CO_3$$

$$C_6H_5OH + Z_0 \xrightarrow{\text{distill}} C_6H_6 + Z_0O$$

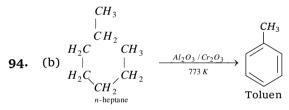
- (c) Unsaturated hydrocarbons are more reactive than saturated hydrocarbons. Among ethyne (C_2H_2) and ethene (C_2H_4) the later is more reactive as $C \equiv C$ triple bond is quite strong bond and therefore ethyne generally require catalysts (like Hg^{2+} etc) in its reactions.
- (a) Kolbe's methods Electrolysis 93. concentrated aqueous solution of either sodium or potassium salts of saturated mono carboxylic acids yields higher alkanes at

$$CH_3COONa = CH_3COO^- + Na +$$

Anode:
$$2CH_3COO^- \xrightarrow{-2e^-} CH_3 - CH_3 + 2CO_2$$

Cathode:
$$2Na^+ + 2e^- \rightarrow 2Na$$

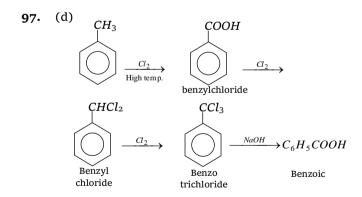
$$2Na + 2H_2O \rightarrow 2NaOH + H_2$$



(a) $-NH_2$ group is not linked with benzene ring. 95.

96. (d)
$$CH_3$$
 $COOH$

$$\underbrace{KMnO_4}$$
benzoic



Critical Thinking Question

1. (a) The difference between any two successive members of the homologous series $-CH_2$ i.e., the molecular weight of every two adjacent members differ by 14. $(CH_2 = 12 + 2 = 14)$

$$CH_{3}$$
(a)
$$CH_{3} - CH - C^{4^{o}} - CH_{2} - CH_{2} - CH_{2} - CH_{3}$$

$$CH_{3} - CH_{3} - CH_{3} - CH_{3}$$

$$CH_{3} - CH_{3} - CH_{3}$$

$$CH_{3} - CH_{3} - CH_{3}$$

$$1^o \Rightarrow \text{Primary } 6, 2^o \Rightarrow \text{Secondary } 2$$

$$3^o \Rightarrow$$
 Tertiary 2, $4^o \Rightarrow$ Quanternar y 1

- (c) Octane number of fuel is the percentage of 3. iso- octane in mixture.
- (b) $CH_3 CH CH_2 CH_3 \rightarrow CH_3 CH = CH CH_3$ 2-Butanol

$$+H_2O$$

According to this rule H atom goes from that β - carbon which is less hydrogenated.

5. (c)
$$CH_3 - C \equiv C - CH_3 \xrightarrow{\text{(ii)} O_3} CH_3 - C - C - CH_3$$

 OOO

6. (c) Friedel-craft's reaction

$$\begin{array}{c} CH_3COCl + C_6H_6 \xrightarrow{\hspace*{1cm} \text{anhy drous } AlCl_3} \to CH_3COC_6H_5 + HCl \\ \text{Acety klhloride} \quad \text{Benzene} \qquad \qquad \text{Acetophenone} \end{array}$$

7. (a)
$$C_2H_6(\text{excess}) + Cl_2 \xrightarrow{U.V.\text{Light}} C_2H_5Cl + HCl \xrightarrow{\text{Ethylchloride} \text{(Maj or product)}}$$

- (a) $CH_3CH_2 OH + 2HI \xrightarrow{Red P} CH_3 CH_3 + H_2O + I_2$ Ethane 8.
- (a) Fischer-Tropsch process -9.

Fischer-Tropsch process –
$$\underbrace{CO + H_2}_{\text{Water gas}} + \underbrace{H_2}_{\text{Excess}} \xrightarrow{\text{heat}} \underbrace{\text{Mixture of hydrocarbo ns} + H_2O}_{\text{(Petrol)}}$$

10. (c)
$$CH_3 - CH_2 - CH_2 - Br + \bigcirc + 2Na \xrightarrow{Dry} + 2Na \xrightarrow{Propyl bromide}$$

$$2NaBr + CH_3 - CH_2 - CH_2$$

(a) Hydroboration of alkenes followed hydrolysis in basic medium yield alcohols and not the alkanes.

$$R-CH=CH_2 \xrightarrow{B_2H_6} (R-CH_2-CH_2)_3-B$$

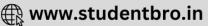
$$\xrightarrow{OH^-} R-CH_2-CH_2OH$$

(b) Koch reaction: (Carboxylation of Alkene)

$$CH_3 - CH = CH_2 \xrightarrow{\text{Water gas}(CO + H_2)} CH_3 - CH - CH_3$$
Isobuty ric acid

$$CH_3$$

13. (d)
$$CH_3 - C = CH - CH_3 \xrightarrow{NaIO_4} CH_3 COCH_3$$



+CH₃CHO

14. (b)
$$HC \equiv CH + 2AgNO_3 \xrightarrow{NH_4OH}$$

$$Ag - C \equiv C - Ag + 2NH_4NO_3 + 2H_2O$$

- (d) Naphthalene is a molecular solid. If the 15. crystals contains only individuals atoms; as in solid argon or krypton or if they are composed of non polar molecules as in naphthalene, the only attraction between the molecules are the London forces.
- (d) A compound is said to be aromatic if it is 16. planar and there is complete delocalization of π electrons, which is only possible if it is a conjugated cyclic system and number of electrons used in delocalization is (4n+2). 1, 3, 5 heptatriene is not an aromatic compound complete delocalization electrons, is not possible in it.



17. (c)
$$CH \equiv CH + HCN \xrightarrow{Ba(CN)_2} CH_2 = CHCN$$

No Vinylcyanide

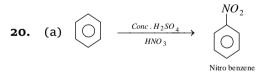
18. (a) Addition reaction means addition on double

19. (d)
$$CaC_2 + 2H_2O \rightarrow Ca(OH)_2 + C_2H_2$$

$$\begin{array}{c} C_2H_2+H_2 \rightarrow C_2H_4 \\ {}_{28\,g} \end{array}$$

64g of CaC_2 gives 28g of ethylene

 \therefore 64kg of CaC_2 will give 28kg of polyethylene

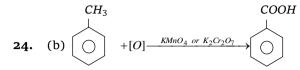


(b) Electron accepting groups which make the 21. substitution difficult are known deactivating groups, the group or substituent already present on the ring also decides the position of incoming group.

> ortho and para directing groups are as follow $-CH_3, C_2H_5(-R), -NH_2, -OH$, halogens, (Cl, Br, I)

23. (c) Polymerisation

$$3CH \equiv CH \xrightarrow[Cu,500^{o}C]{\text{heat}}$$



(a) Wohler reaction: 25.

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

$$C_2H_2 \xrightarrow{dil.H_2SO_4/HgSO_4} [CH_2 = CHOH] \rightarrow CH_3CHO$$
Acetylene Acetaldehyde

26.

(a)
$$CH = CH + H_2O \xrightarrow{H_3SO_4} CH_2 = CH - OH \xrightarrow{KMnO_4} CH_3CHO \xrightarrow{Oxidation} CH_3COOH$$

27. (a)
$$CaC_2 + 2D_2O \longrightarrow C_2D_2 + Ca(OD)_2$$

- (d) It is a unsymmetrical olefin. In such cases addition of H-Xis governed 'Markownikoff's rule'
- (d) $CH_3 CH = CH_2 \xrightarrow{\text{Conc.}} CH_3COOH + HCOOH$

(b) Biological oxidation of benzene gives cinnamic 31. acid.

32. (d)
$$CH \equiv CH \xrightarrow{HBr} CH_2 = CH - Br \xrightarrow{HBr}$$

$$CH_3CH < \frac{Br}{Br}$$

(c) Electron accepting groups which make the substitution difficult are known deactivating groups.

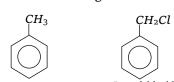
o. p. directing groups are as follow:

$$-CH_3, C_2H_5(-R), -NH_2, -OH$$
, halogens (Cl, Br, I) .

34. (a) 2-butene shows geometrical isomerism.

$$\begin{array}{ccc} H-C-CH_3 & H-C-CH_3 \\ & || & || \\ H-C-CH_3 & CH_3-C-H \\ \text{cis-Butene -2} & \text{trans-Butene -2} \end{array}$$

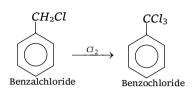
(d) Sidechain chlorination and bromination is 35. favoured by high temperature, light and in absence of halogen carrier.











36. (b)
$$C_2H_4 + CO + H_2O \xrightarrow{\text{Ligh temp.}} C_2H_5 - COOH$$
Ethylene Propionic acid

37. (b) Cyclohexane C_6H_{12} is alicyclic compound.

38. (c)
$$CH_2 = CH_2 \xrightarrow{HBr} CH_3 - CH_2 \xrightarrow{\text{Hydroly sis}} Br$$

$$CH_3 - CH_2 \xrightarrow{Na_2CO_3} CHI_3 \text{ Yellowppt (Iodoform)}$$

39. (d) On oxidation, with $KMnO_4$, they give different alcohols

$$\begin{array}{c} CH_{3}CH_{2}CH_{2}CH_{2}CH_{3} \xrightarrow{KMnO_{4}} \\ {}_{n \text{ pentane}} \end{array} \longrightarrow \\ CH_{3}CH_{2}CH_{2}CH_{2}CHOH_{2}$$

$$CH_{3}CH_{2} - CH - CH_{3} \xrightarrow{KMnO_{4}} CH_{3}CH_{2} - C - CH_{3}$$

$$CH_{3} \xrightarrow{CH_{3}} CH_{3} \xrightarrow{3^{\circ} \text{alcohol}}$$

40. (b)
$$CH \equiv CH + HBr \rightarrow CH_2 = CHBr$$
Vinyl bromide

Assertion & Reason

 (a) This reaction is followed by against of Markownikoff rule

$$\begin{array}{c|cccc} H & H & H & H \\ & & & & & \\ H - C - C - C - C & C & + HBr & \xrightarrow{\text{Peroxide}} \\ & & & & \\ & & & & \\ H & H & & H & \\ \end{array}$$

In this reaction anti Markownikoff's addition is explained on the basis of the fact that in the presence of peroxide the addition takes place Via a free radical mechanism rather than the ionic mechanism.

2. (a)
$$CH_3 - CH_2 - CH = CH_2 + Br_2 \rightarrow$$

$$CH_2 - CH_2 - C^{\bullet} - CH_2Br$$

$$Br$$
1.2 dibromobut ane

4. (a) Pyrrole is a heterocyclic compound having five membered ring. It has 6π electrons. As 4π electrons in the ring and 2π electrons donated by nitrogen atom present in the ring.

5. (b) The assertion that chlorination of ${\it CH}_4$ does not take place in dark is correct because it is a free radical reaction and free radicals are obtained in presence of sun light.

6. (b) The alkyl benzene is not prepared by Friedel Craft's alkylation of benzene because the monoalkyl product formed undergo alkylation to produce polyalkylated benzene.

The reason that alkyl halides are less reactive than acyl halides is also correct but this is not the correct explanation of assertion.

7. (d) 2-bromobutane on reaction with sodium ethoxide in ethanol gives 2-butene as a major product.

$$EtOH + CH = CH + Br^{-}$$

$$CH_{3}$$

This is according to saytzeff's rule i.e. the Predominant product is the most substituted alkene, i.e. are carrying the largest number of alkyl substituents of hydrogen is eliminated preferentially from the carbon atom joined to the least number of hydrogen atoms.

2-butene is more stable than 1-butene due to presence of large number of hyperconjugating structures in 2-butene.

8. (b) *HBr* adds to the double bond of the styrene forming 1-bromo-1-phenylethane.

$$\xrightarrow{HBr} \xrightarrow{Br}$$
Styren

This is an example of elimination-addition reaction.

Note: Here given assertion is wrong.

9. (b) Alkanes with odd carbon atoms have their and carbon atom on the same side of the molecule





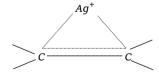
and in even carbon atom alkane, the end carbon atom on opposite side. Thus alkanes with even carbon atoms are packed closely in crystal lattice to permit greater intermolecular attractions and hence higher melting point.

10. (b) Iodination is reversible since formed HI is a strong reducing agent and reduces the alkyl iodide back to alkane.

$$CH_4 + I_2 = CH_3I + HI$$

11. (b) Normally oxidising agents have no effect on alkanes. However, alkanes containing a tertiary hydrogen atom can be oxidised by oxidising agents such as KMnO_4 to the corresponding alcohols. For example,

- **12.** (e) Halogenation of alkanes is catalysed by radical initiators like benzoyl peroxide.
- 13. (c) Neopentane has four identical methyl groups attached on 4° carbon atom and thus only one product is possible during monosubstitution.
- **14.** (b) Neopentane is more symmetric than *n*-pentane and it is expected to show more stable crystal lattice than that of *n*-pentane and thus have high melting point.
- **15.** (b) Knocking involves the production of metallic sound due to irregular burning of the fuel. This irregular burning lowers the efficiency of the engine.
- **16.** (b) Ag^+ coordinates with the alkene by $p\pi d\pi$ bonding giving an ion and the solubility increases.



17. (a) In the dehydration of secondary and tertiary alcohols. When there is a possibility of formation of two isomers. The hydrogen atom is preferentially eliminated from the adjacent carbon atom with the fewer number of hydrogen atoms. This is Saytzeff's rule.

$$CH_{3} - CH_{2} - \begin{matrix} CH_{3} \\ CH - CH_{3} \end{matrix} \xrightarrow{60\% H_{2}SO_{4}} \\ CH_{3} \\ CH_{3} \\ CH_{3}CH_{2}CH = CH_{2} + CH_{3}CH = CHCH_{3} \\ \\ 1-\text{butene (20\%)} \\ 2-\text{butene (80\%)} \end{matrix}$$

18. (c) The addition of Br_2 , follows ionic mechanism

$$H_2C = CH_2 \xrightarrow[NaCl \to Na^+ + Cl^-]{\delta^+ \quad \delta^- \atop NaCl \to Na^+ + Cl^-}} H_2 \stackrel{+}{C} = \stackrel{-}{C} H_2 \xrightarrow{\quad +Br \atop +Br}$$

$$H_2 \stackrel{+}{C} = \stackrel{-}{C} H_2 Br$$

$$H_2 \stackrel{+}{C} = \stackrel{-}{C} H_2 Br$$

$$1, 2$$

$$+Cl^- \longrightarrow Cl \ CH_2 - CH_2 \ Br$$
1-bromo-2-chloro

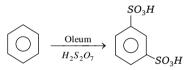
- 19. (b) Straight chain alkanes have very low octane number because as the length of the chain increases, octane number further decreases.
- **21.** (d) The two hydrogen atoms on first carbon and the two H-atoms on the third carbon atom lie in perpendicular planes. The central carbon atom is sp-hybridized while terminal carbon atoms are sp^2 -hybridized.
- **22.** (d) Propene reacts with *HBr* in presence of benzoyl peroxide to give 1-bromopropane. In presence of peroxide, the addition of *HBr* to propane occurs by free radical mechanism.
- **24.** (b) Arylhalide shows resonance stabilization and thus resistant to substitution reaction.
- **25.** (e) During alkylation of bromobenzene, if benzene is used as solvent, alkylation of benzene will take place because benzene is more reactive for S_E reactions than bromo benzene, benzene is not used as a solvent for this reaction.



- **26.** (b) Benzene is a polar solvent Butter is composed of organic compounds of low polarity. So it gets dissolved in benzene.
- 27. (a) CH_3 In toluene , the $-CH_3$ group is electron

Pushing. It is an activating group. By its+I effect, $-CH_3$ group activates the benzene ring at ortho and para position relative to it. Due to increased activity towards electrophilitic substitution toluene can be more easily nitrated than benzene.

28. (e) With fuming H_2SO_4 or oleum(conc. $H_2SO_4 + SO_3$) at high temperature benzene forms m-benzene disulphonic acid.



m-Benzene disulphonic

- **29.** (c) Like halogens, the nitroso group (-N=O) is also deactivating but o, p-directing. It is deactivating because O is more electronegative than N and hence No group as whole withdraws electrons from the benzene ring.
- **31.** (e) Completely conjugated polyenes containing even number of carbon atoms is known as annulenes.
- **32.** (c) $(4n+2)\pi$ electrons and planar structure are the essential conditions for aromaticity.
- **33.** (a)



Due to severe non-bonded interactions between the internal hydrogen (as shown in figure), the ring assumes non-planar geometry.