KD EDUCATION ACADEMY [9582701166] street no. 21 A-1 block Bengali colony sant nagar burari delhi -110084

Time: 5 Hour

STD 11 Science Chemistry kd 90+ Ch-9 hydrocarbon

Total Marks: 270

* Choose The Right Answer From The Given Options.[1 Marks Each]

[78]

1. In which of the following, all atoms are coplanar?

(A)



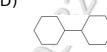
(B)



(C)



(D)



- 2. The bond order of individual carbon-carbon bonds in benzene is:
 - (A) One

(B) Two

(C) Between one and two

- (D) One and two, alternately
- 3. Which of the following compounds will exhibit geometrical isomerism?
 - (A) 2-Phenyl-1-butene.

(B) 1, 1-Diphenyl propane.

(C) 1-Phenyl-2-butene.

- (D) 3-phenyle butane.
- 4. What are A and B in following reaction?

 \mathbf{H}

$$ext{H}_2 ext{C} - ext{C} - ext{H} + ext{KOH} \xrightarrow{ ext{Alcohol}} ext{A} \xrightarrow{ ext{NaNH}_2} ext{E}$$

	Di	
	A	В
(a)	$ m H_2C=CH_2$	$\mathrm{CH} \equiv \mathrm{CH}$
(b)	$\mathrm{CH}_2 = \mathrm{CHBr}$	$\mathrm{CH} \equiv \mathrm{CH}$
(c)	$\mathrm{CH}_2 = \mathrm{CHBr}$	$\mathrm{CH}_2 = \mathrm{CH}_2$
(d)	$\mathrm{CH}_2 = \mathrm{CH}_2$	$\mathrm{CH} \equiv \mathrm{CBr}$

- 5. 2.84gms of methyl iodide was completely converted into CH₃Mgl and the product was decomposed by the excess of ethanol. The volume of the gaseous hydrocarbon produced at NTP will be_____.
 - (A) 22.4lit.
- (B) 2240lit.
- (C) 0.448lit.
- (D) 224lit.

- 6. The geometry of ethyne molecule is:
 - (A) Linear.

(B) Trigonal planar.

(C) Tetrahedral.

(D) Bent.

7.	IUPAC name of the fir	st member of al	kyne is						
	(A) Ethene	(B) Ethyne	1	(C) Ethan	e	(D) Methane			
8.	An optically active hydrocarbon (X) on catalytic hydrogenation gives an optically inactive compound (Y), C_6H_{14} . The hydrocarbon (X) is-								
	(A) 3-methyl-1-penter	ne	(B) 3-me	thyl-2-penter	ne				
	(C) 2-ethyl-1-butene			(D) 3-me	thylcyclopen	tene			
9. Calcium heptane dioate on distillation produces:									
	(A) Cyclo hexanal		(B) Cyclohexene						
	(C) Cyclohexanone			(D) Cyclohexanoic acid					
10.	Cycloalkanes are iso	meric with:							
	(A) Alkanes	(B) Alkenes		(C) Alkyne	es	(D) Arenes			
11.	The percentage of iso room temperature re		_			-			
	(A) 81%	(B) 56%	((C) 15%		(D) 22%			
12.	The addition of HBr t	o 1-butene gives	s a mixture	of produc	cts A, B and C				
	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	H ₂ —CH ₂ —Br							
	The mixture consists	of:							
	(A) A and B as major	and C as minor	products.						
	(B) B as major, A and	C as minor prod	ducts.						
	(C) B as minor, A and	C as major prod	ducts.						
	(D) A and B as minor	and C as major	products.	/					
13.	How many chain ison	ners can be forn	ned by but	ane?					
	(A) One	(B) Three	V	(C) Two		(D) Four			
14.	4. Which of the following is not a cumulated diene?								
(A) Hexa-1, 2-diene (B) Hexa-2, 3-die									
	(C) Penta-2, 3-diene)	(D) Penta	a-1, $3-diene$				
15. The correct IUPAC name of the following alkane is: H,C—CH2—CH2—CH2—CH2—CH3									
	$ \begin{array}{ccc} $								
	(A) 3, 6 – Diethyl – 2 –	(4)			opropyl – 3 –	-			
	(C) 3 – Ethyl – 5 – isop	ropyloctane.		(D) 3 – Is	opropyl – 6 –	ethyloctane.			
16.	A direct iodination of		possible be	ecause:					
	(A) lodine is an oxidis								
	(B) Resulting C_6H_5I is	reduced to C ₆ H	₆ by HI.						
	(C) HI is unstable.								
	(D) The ring gets dea								
17.	Which of the followin	_	trical isom						
	(A) 2-methylpropene.			(B) But-1		_			
	(C) But-2-ene.			(D) 2, 3-d	dibromo-but-	2-ene.			
18.									

	Examples of addition out?	reactions include all but	one of the following. Wh	ich is the odd one				
	(A) Reaction of HBr w	ith but-2-ene.	(B) Combustion of propene.					
	(C) Reaction of Cl ₂ wi	th propene.	(D) Polymerization of e	ethene.				
19.	CH ₃ -C \equiv C MgBr can be prepared by the reaction of							
	(A) CH ₃ −C≡C−Br wit	h MgBr ₂	(B) CH ₃ −C≡CH with M	gB ₂				
	(C) CH ₂ −C≡CH with H	〈Br and Mg metal	(D) CH ₃ −C≡CH with C	H ₃ MgBr				
20.	On heating sodium p	ropionate with sodalime	the hydrocarbon obtaine	d is:				
	(A) Butane	(B) Ethane	(C) Propane	(D) Ethene				
21.	In an alkane if the va	lue of $n = 19$, it is:						
	(A) Solid		(B) Liquid					
	(C) Gaseous		(D) None of the above					
22.	Arrange the following	g carbanions in order of t	heir decreasing stability.					
	A. H							
	$ m CH_3 - C -$	$ m CH_2Br$						
	CH.	•						
	B. $CH_3 - CH_2$							
		$ ho - \mathrm{CH}_2 - \mathrm{Br}$	(C) D > C > A	(D) A > C > D				
		(B) $C > B > A$	(C) $B > C > A$	(D) $A > C > B$				
23.		g gives propyne on hydro		(D) 1 - C				
	(A) Al ₄ C ₃	(B) M _g 2C ₃	(C) B ₄ C	(D) La ₄ C ₃				
24.		the alkene H ₂ C=CH–CH=						
	(, ,, 1, 5 5 4 4 4 4 1 1 1		(B) 1, 3-butanediene					
	(C) 1, 3-dibutene		(D) 1, 3-dibutadiene					
25.	5. The compound formed as a result of oxidation of propyl benzene by hot alkaline $KMnO_4$ is:							
	(A) Benzoic acid.		(B) 2-phenyl ethanoic acid.					
	(C) 3-phenyl ethanoic	(D) Acetophenone.	phenone.					
26.		with Na to give $CH_3CH_2CH_3$		is:				
	(A) CH ₃ CH ₂ OH		(B) CH ₃ CH ₂ -Cl					
	(C) $CH_3 - CH_3$		(D) $CH_3 - CH_2 - CH_2 - CH_3 - CH_$	_				
27.	The degree of unsatu phenol is:	ration of the compound	formed by the partial hyc	lrogenation of				
	(A) 3	(B) 4	(C) 2	(D) 1				
28.	In the reaction,							
	+ H ₂ SO ₄ - (Fuming sulphuric acid)	$\xrightarrow{\Delta}$ Y,						
	(A) C ₆ H ₅ SO ₃ H	(B) C ₆ H ₅ OH	(C) $C_6H_5SO_2$	(D) C ₆ H ₅ CHO				
29.								

Secondary butyl chloride will undergo alkaline hydrolysis in the polar solvent by the mechanism:

(A) S_N2

(B) S_N1

(C) S_N1 and S_N2

- (D) None of the above
- 30. Four structures are given in options (i) to (iv). Examine them and select the aromatic structures.

(A)



(B)



(C)



(D)



- 31. Which of the following is Aromatic Hydrocarbon?
 - (A) C_2H_2
- (B) C_3H_8
- (C) C₅H₁₂
- (D) C_6H_6
- 32. Which of the following will not show geometrical isomerism?

(A)

(B)

$$C = C$$

(C)

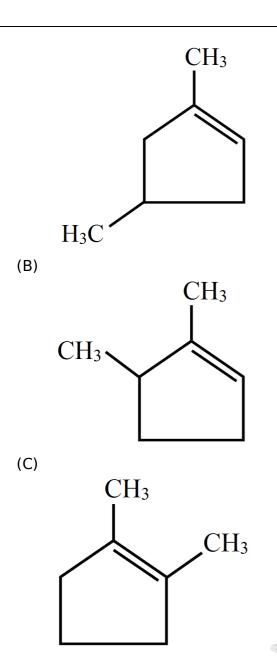
$$H_3C$$
 $C = C$
 C_2H_5
 $C = C$
 CH_3

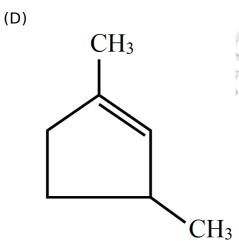
(D)

$$CH_3$$
 $C = C$ CH_3 CH_5

33.	Consider the nitration of benzene using cor $KHSO_4$ is added to mixture, the rate of nitrations of the same of the contract of the same of the contract of t		c. HNO ₃ . If large amount of			
	(A) Unchanged. (B) Doubled.	(C) Faster.	(D) Slower.			
34.	How many isomers are possible in C ₄ H ₈ ?					
	(A) 3 (B) 2	(C) 4	(D) 5			
35.	Which of the following deactivates the benzand is o/ p directing?	zene ring towards e	electrophilic substitution			
	(A) $-\mathrm{Cl}^\ominus$	(B) -OCH ₃				
		(C) -CHO				
	(D) All of these.					
36.	The IUPAC name of $(CH_3)_3C-CH=CH_2$ is:					
	(A) 1, 1, 1-trimethyl-2-propene	(B) 3, 3, 3-trir	methyl-2-propene			
	(C) 2, 2-dimethyl -3-butene	(D) 3, 3-Dime	(D) 3, 3-Dimethyl-1-butene			
37.	Ethylene is a/an:					
	(A) Saturated hydrocarbon	(B) Alkene				
	(C) Paraffin	(D) Aromatic	compound			
38.	Which of the following alkenes on ozonolys	is give a mixture o	f ketones only?			
	(a) CH_3 — CH = CH — CH_3					
	(b) CH_3 — C — CH = CH_2 CH_3					
	Cn_3					
	(c) \subset \subset CH_3 CH_3	3				
	(d) $(CH_3)_2 C = C CH_3$ CH_3					
39.	The IUPAC name of CH ₃					
	is					
	CH ₃					
	(A) Dimethylbenzene.	(B) Methyltol	uene.			
	(C) 1, 3-dimethylbenzene.	(D) 1, 4-dime	thylbenzene.			
40.	Which one of the following compounds give	es methane on trea	atment with water?			
	(A) AI_4C_3 (B) CaC_2	(C) VC	(D) SiC			
41.	Propene is allowed to react with $B_2\ D_6$ and final product obtained is:	the product is trea	ted with acetic acid. The			
	(A) 1-deuteriopropane	(B) 2-deuterio	opropane			
	(C) 1-deuteriopropene	(D) 2-deuteri	opropene			
42.	Which type of isomerism is shown by But-1-	yne and cyclobute	ne?			

	(A) Ring chain isomerism	(B) Chain isomerism
	(C) Position isomerism	(D) Metamerism
43.	Which of the following is properly matched?	
	(A) Kerosene - $C_5 - C_{10}$	(B) Diesel oil - $C_5 - C_6$
	(C) Petrol - C ₇ -C ₉	(D) All are correct
44.	When cyclohexane is poured on water, it floats	s because cyclohexane is:
	(A) In boat form	(B) In chair form
	(C) In crown form	(D) Less dense than water.
45.	What happens when calcium carbide is treated	d with water?
	(A) Ethane is formed.	(B) Methane and ethane are formed.
	(C) Ethyne is formed.	(D) Ethene and ethyne are formed.
46.	Compound A (molecular formula C_5H_{10}) gives the structure of compound A.	only one monochlorinated product. Write
	(A) 1-pentene	(B) 2-pentene
	(C) 1-methyl cyclobutane	(D) Cyclopentane
47.	For an electrophilic substitution reaction, the p benzene ring	resence of a halogen atom in the
	(A) Deactivates the ring by inductive effect.	
	(B) Deactivates the ring by resonance.	
	(C) Increases the charge density at ortho and presonance.	para position relative to meta position by
	(D) Directs the incoming electrophile to meta p to ortho and para position.	position by increasing the charge density relative
48.	The C-C bond length of the following molecule	es is in the order:
	(A) $CH_2 > C_2H_4 > C_6H_6 < C_2H_2$	(B) $C_2H_2 < C_2H_4 < C_6H_6 < C_2H_6$
	(C) $C_2H_6 < C_2H_2 < C_6H_6 < C_2H_4$	(D) $C_2H_4 < C_2H_6 < C_2H_2 < C_6H_6$
49.	Arrange the halogens F_2 , Cl_2 , Br_2 , l_2 , in order c	of their increasing reactivity with alkanes:
	(A) $I_2 < Br_2 < CI_2 < F_2$.	(B) $Br_2 < Cl_2 < F_2 < l_2$.
	(C) $F_2 < Cl_2 < Br_2 < l_2$.	(D) $Br_2 < I_2 < CI_2 < F_2$.
50.	Which of the following organic materials dama	age DNA of our body?
	(A) Tobacco.	(B) Coal.
	(C) Petroleum.	(D) All of these.
51.	Which compound would give 5-keto 2-methyl has reaction with $\frac{Zn}{H_2O}$?	nexanal upon ozonolysis followed by
	(A)	





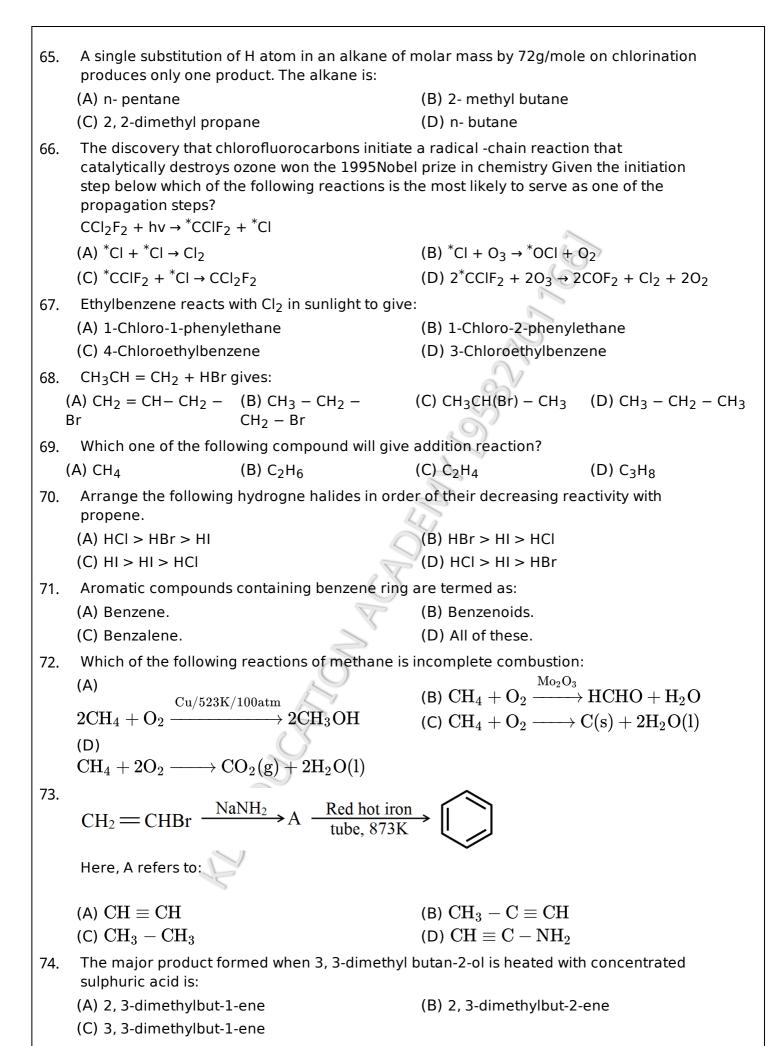
- 52. Conformations of ethane in which hydrogen atoms attached to two carbon atoms are closest to each other are known as:
 - (A) Eclipsed.

(B) Skew boat.

(C) Staggered.

- (D) Partially eclipsed.
- 53. Nitration and chlorination of benzene are:
 - (A) Nucleophilic and electrophilic substitution respectively.

	(B) Electrophilic and nucleophilic substitu	-								
	(C) Electrophilic substitution in both the reactions.									
	(D) Nucleophilic substitution in both the	eactions.								
54.	4. Alkynes on reduction with sodium in liquid ammonia forms:									
	(A) Cis-alkene.	(B) Alkane.								
	(C) Trans-alkene.	(D) Both (a)	and (c).							
55.	The reaction of Toluene with Cl ₂ in prese	ence of FeCl ₃ predon	ninanly gives:							
	(A) Benzyle Chloride.	(B) o- & p-ch	lorotoluene.							
	(C) m-Chlorotoluene.	(D) Benzoyl	chloride.							
56.	Which content(s) of middle oil separate of	on cooling?								
	(A) Naphthalene (B) Phenol	(C) Benzene	(D) Pyridine							
57.	The energy barrier to free rotation abou	t carbon carbon bon	d of ethane is about:							
	(A) $30kJ \text{ mol}^{-1}$	(B) 3kJ mol	ì							
	(C) $3kcal mol^{-1}$	(D) 40kJ mo	_1							
58.	2-methyl propan-2-ol is obtained by the conc. H ₂ SO ₄ . The X is:	reaction of X with wa	ater in the presence of							
	(A) 1-methylpropene.	(B) 2, 2-dim	ethylhexane.							
	(C) 2-methylpropene.	(D) 2-methy	lbutane.							
59.	Name the IUPAC nomenclature of CH ₃ -0	CH ₂ -CH ₂ -CH=CH ₂ :								
	(A) 1-Pentene	(B) 2-Proper	ne							
	(C) 3-Propene	(D) None of	these							
50.	Which of the following can be used as th crafts reaction:	e halide (electrophil	e provider) for Friedel							
	(A) Chloro benzene.	(B) Bromo b	enzene.							
	(C) Chloro ethane.	(D) Isopropy	ıl chloride.							
51.	The hydrocarbon obtained by treating so	odium ethanoate wit	h soda lime is							
	(A) Ethane (B) Methane	(C) Propane	(D) Butane							
52.	Ethylene is prepared by:									
	(A) Dehalogenation of chloroform.									
	(B) Pyrolysis of ethane at 450°C									
	(C) Dehydration of methanol with Al ₂ O ₃ /	350°C								
	(D) Methyl chloride on reduction									
53.	The highest boiling point is expected for	:								
	(A) Isooctane	(B) n-Octane	è							
	(C) 2, 2, 3, 3-Tetramethyl butane	(D) n-Butan	е							
54.	In the reaction, ${ m Br}$	A is:								
	$ ext{CH}_3 - ext{CH}_2 \xrightarrow{ ext{alc. KOH}} A$	L ,								
	(A) $CH_2 = CH_2$	(B) CH ₃ CH ₃								
	(C) $\mathrm{CH} \equiv \mathrm{CH}$	(D) Both (a)	and (b)							



(D) Cis and trans-isomers of 2, 3-dimethylbut-1-ene

75. The IUPAC name of given compound is: $CH_3-C\equiv C-C(CH_3)_2-CH_3$

(A) 4, 4-dimethyl-2-heptene-5-yne

(B) 4, 4-dimethyl-2-heptene-6-yne

(C) 4, 4-dimethyl-5-heptene-2-yne

(D) None of these

76. Phenyl magnesium bromide reacts with methanol to give:

(A) Benzene.

(B) Toluene.

(C) Phenol.

(D) Arisole.

77. Monochlorination of ethylbenzene (PhCH₂CH₃) with Cl₂ under heat produces _____

(A) PhCH2CH2CI

(B) PhCHClCH₃

(C) Both (a) and (b) in equal amounts

(D) None of (b) and less of (a)

78. Among the following, which hydrocarbon is not produced by Wurtz reaction?

(A) Methane.

(B) Ethane.

(C) Propane.

(D) All given options can be prepared.

* Answer The Following Questions In One Sentence.[1 Marks Each]

[20]

79. Write IUPAC names of the following compound:

80. Write IUPAC names of the products obtained by the ozonolysis of the following compound:

Pent-2-ene.

81. Write IUPAC names of the products obtained by the ozonolysis of the following compound:

3,4-Dimethylhept-3-ene.

- 82. Why is addition reaction of bromine to benzene difficult?
- 83. Give the structural formula of 4-tert, butyl-4-ethyl-2, 2, 5-tetra methyl hexane.
- 84. Bring out the following conversion ethane to ethene.
- 85. Convert ethyne to propyne.
- 86. Why eclipsed and staggered forms of ethane cannot be isolated at room temperature?
- 87. Name the reagent used in the following changes

- 88. In which of the following pair both angle strain and torsional strain is present?
 - a. Cyclopropane, Cyclobutane
 - b. Cyclopropane, Cyclohexane

- Cyclobutane, Cyclohexane c.
- d. Cyclohexane, Cyclopentane
- 89. Arrange the three isomers of pentane in increasing order of their boiling points.
- 90. Identify the structure of A and B;

$$\operatorname{CH}_3 - \operatorname{CH}_2\operatorname{COOH} \xrightarrow{\operatorname{KOH}} \operatorname{A} \xrightarrow{\operatorname{Kolbe's electrolysis}} \operatorname{E}$$
 $\operatorname{CH}_3 - \operatorname{CH}_2\operatorname{COOH} \xrightarrow{\operatorname{KOH}} \operatorname{CH}_3\operatorname{CH}_2\operatorname{COO}^-\operatorname{K}^+$
 A
 $\operatorname{Kolbe's electrolysis} \longrightarrow \operatorname{CH}_3\operatorname{CH}_2\operatorname{CH}_2\operatorname{CH}_3$

- Draw the Newman's projection formula of the staggered form of 1, 2-dichloroethane. 91.
- 92. Convert methane to ethane.
- Write an equation of the reaction of propyne with water in the presence of H₂SO₄ and 93. HgSO₄. Show the intermediate.
- Why is dipole moment of trans-1, 2-dichloroethene zero? 94.
- 95. Convert 1-bromopropane to 2-bromopropane.
- Why do alkenes prefer to undergo electrophilic addition reaction while arenes prefer 96. electrophilic substitution reactions? Explain.
- 97. Alkynes on reduction with sodium in liquid ammonia form trans alkenes. Will the butene thus formed on reduction of 2-butyne show the geometrical isomerism?
- 98. Give the structure of the alkene (C_4H_8) which adds on HBr in the presence and in absence of peroxide to give the same product, C₄H₉Br.

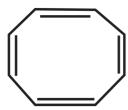
Given Section consists of questions of 2 marks each.

[28]

99. Explain why the following system are not aromatic?



100. Explain why the following system are not aromatic?



101. Complete the following:

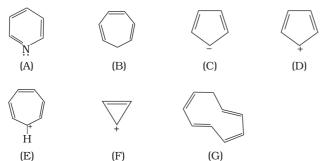
(i)
$$H_3C$$
 COONa Sodalime Heat

An alkyl bromide (X) react with sodium metal dissolved in anhydrous ether to form 4, 5-102. diethyl octane. Identify 'X'.



- i. Planar ring containing conjugated π bonds.
- ii. Complete delocalisation of the $\pi-$ electrons in ring system i.e. each atom in the ring has unhybridised p-orbital, and
- iii. Presence of $(4n + 2)\pi$ electrons in the ring where n is an integer (n = 0, 1, 2,) [Huckel rule].

Using this information classify the following compounds as aromatic/ nonaromatic.



104. Complete the following reactions:

i.
$$(CH_3)_2C=CH-CH_3\xrightarrow{KMnO_4/KOH}$$
 ii. $(CH_3)_2C=CH-CH_3\xrightarrow{cold\ KMnO_4/KOH}$

105. Complete:

i. Cl $\xrightarrow{\text{Na}}$ Ether ii.

Cl + CH_3 - Cl Na Ether

106. Explain why the following systems are not aromatic?

$$\left\langle \begin{array}{c} \\ \end{array} \right\rangle = CH_2$$

- 107. What happens when 1, 2-Dibromo ethane reacts with excess of alc. KOH?
- 108. Complete the following reactions.
 - $\begin{array}{ll} \text{i.} & Iso\text{-propyl bromide} \xrightarrow{Alc. \ KOH} A \xrightarrow{HBr} B \\ \\ \text{ii.} & n\text{-propyl alcohol} \xrightarrow{Conc. \ H_2SO_4} A \xrightarrow{O_2, \ Ag} B \end{array}$

109. What will be the product obtained as a result of the following reaction and why?

$$+ H_3C-CH_2-CH_2Cl$$
 \longrightarrow

- 110. What happens when:
 - i. Bromoethane is treated with zinc and hydrochloric acid?
 - ii. Hydrogen is passed into 2-bromopropane in the presence of palladium?
- 111. Write the structures and names of products obtained in the reactions of sodium with a mixture of 1-iodo-2-methylpropane and 2-iodopropane.
- 112. Convert Acetylene to Propylene.

* Given Section consists of questions of 3 marks each.

[66]

113. 2 Arrange the following set of compounds in order of their decreasing relative reactivity with an electrophile, E⁺.

Toluene, $p-H_3C-C_6H_4-NO_2$, $p-O_2N-C_6H_4-NO_2$.

- 114. For the following compound, write structural formulas and IUPAC names for all possible isomers having the number of double or triple bond as indicated: C_5H_8 (one triple bond).
- 115. How will you convert benzene into: p-nitrobromobenzene.
- 116. For the following compound, write structural formulas and IUPAC names for all possible isomers having the number of double or triple bond as indicated: C_4H_8 (one double bond).
- 117. Suggest the name of a Lewis acid other than anhydrous aluminium chloride which can be used during ethylation of benzene.
- 118. 2 Arrange the following set of compounds in order of their decreasing relative reactivity with an electrophile, E⁺.

Chlorobenzene, 2,4-dinitrochlorobenzene, p-nitrochlorobenzene.

119. Give the structures of A and B.

$$\begin{array}{c} C+D \xleftarrow{\text{Hot KMnO}_4} & A \\ \text{(Isomers of formula C_3H_6O)} & A \\ \end{array} \xrightarrow{(i) \ O_3} B+C \\ \text{(Isomers of formula C_3H_6O)} \\ \end{array}$$

120. Complete the following reactions:

i.
$$CH_3-CH=CH_2+HBr\xrightarrow{\mathrm{Peroxide}}$$
ii. $CH_3-Cl+Na\xrightarrow{\mathrm{Dry\;ether}}$
iii. $CH_3-CH-CH_2-CH_3\xrightarrow{\mathrm{alc.\;KOH}}$

121. Complete the reactions:

i.
$$\text{CH}_3\text{CH}_2\text{CH}_3 + \text{HNO}_3 \xrightarrow{\text{Vapour phase}}$$
ii. $\text{CH}_4 + \text{O}_2 \xrightarrow{\text{Copper tube}} \xrightarrow{723-773\text{K}}$
iii. $\text{C}_2\text{H}_5 \xrightarrow{\text{COONa}} + \text{NaOH} \xrightarrow{\text{CaO}}$

122. Pent-1-yne
$$\xrightarrow{\text{(i) NaNH}_2/\text{NH}_3}$$
 A $\xrightarrow{\text{H}_2}$ B $\xrightarrow{\text{Br}_2}$ 'C $\xrightarrow{\text{Lindlar's catalyst}}$

Identify A, B and C compounds and give their reactions.

- 123. Convert:
 - i. Ethylene to Nitrobenzene.
 - ii. Write short note on Markovnikov's Rule.
- 124. Which of the following compounds are aromatic according to Huckel's rule?

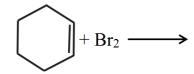
$$(A) \qquad (B) \qquad (C) \qquad H_1C \longrightarrow CH_2$$

$$(D) \qquad (E) \qquad (F) \qquad (F)$$

- 125. For the following reactions, complete and Identify the type of reactions:
 - i. $CH_3 CH_2 CH_2Br + KOH(alc.) \longrightarrow$

ii.
$$CH_3 - CH - CH_3 + KOH(aq) \longrightarrow$$

iii.

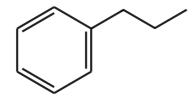


- 126. An alkyl halide (X) of formula $C_3H_{13}Cl$ on treatment with alcoholic KOH or potassium tert-butoxide gives two isomeric alkenes Yand $Z(C_6H_{12})$. Both alkenes on hydrogenation give 2, 3-dimethylbutane. Predict the structure of X, Y and Z.
- 127. Give the reactions involved in the preparation of propane from the following:
 - i. $CH_3 CH = CH_2$
 - ii. $CH_3 CH_2 CH_2Cl$
 - iii. $CH_3CH_2CH_2COO^-Na^+$
- 128. An alkyl halide 'A' of formula $C_6H_{13}Cl$ on treatment with alcoholic KOH gives two isomeric alkenes 'B' and 'C' with molecular formula C_6H_{12} . Both alkenes on hydrogenation give 2, 3-dimethyl butane. Predict the structure of 'A', 'B' and 'C'.
- 129. Arrange 2, 2-dimethylbutane, 3-methylpentane and n-hexane in increasing order of their boiling point.
- 130. a. Name a compound that will be required to obtain butane using Kolbe's electrolytic method.
 - b. Why does benzene show electrophilic substitution easily?
 - c. Complete the following:

$$\mathrm{CH_3} - \mathrm{C} = \mathrm{CH_2} + \mathrm{H_2O} \overset{\mathrm{H^+}}{\longrightarrow} \ \mathrm{CH_3}$$

- 131. Write chemical reactions of:
 - i. Wurtz-Fittig reaction.
 - ii. Kolbe's electrolytic method.
 - iii. B-Elimination reaction.
- 132. Write IUPAC names of the following:

a.



b.

c.

$$Cl$$
 3
 1
 H

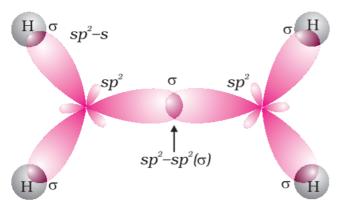
- 133. An alkane has a molecular mass of 72. Give all the possible structural isomers along with their IUPAC names.
- 134. i. How will you convert benzene into:
 - a. Acetophenone?
 - b. m-nitrochlorobenzene?
 - ii. Write the structures of products obtained by ozonolysis of pent-2-ene.

* Case study based questions

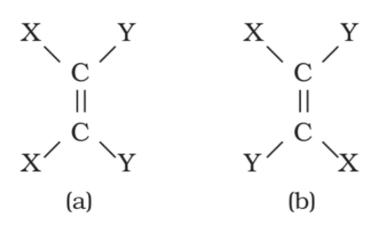
[8]

135. Read the passage given below and answer the following questions from 1 to 5. Alkenes are unsaturated hydrocarbons containing at least one double bond. What should be the general formula of alkenes? If there is one double bond between two carbon atoms in alkenes, they must possess two hydrogen atoms less than alkanes. Hence, general formula for alkenes is C_nH_{2n} . Alkenes are also known as olefins (oil forming) since the first member, ethylene or ethene (C_2H_4) was found to form an oily liquid on reaction with chlorine.

Structure of Double Bond Carbon-carbon double bond in alkenes consists of one strong sigma (σ) bond (bond enthalpy about 397 kJ mol⁻¹) due to head-on overlapping of sp² hybridised orbitals and one weak pi π bond (bond enthalpy about 284 kJ mol⁻¹) obtained by lateral or sideways overlapping of the two 2p orbitals of the two carbon atoms. The double bond is shorter in bond length (134 pm) than the C-C single bond (154 pm). You have already read that the pi (π) bond is a weaker bond due to poor sideways overlapping between the two 2p orbitals. Thus, the presence of the pi (π) bond makes alkenes behave as sources of loosely held mobile electrons. Therefore, alkenes are easily attacked by reagents or compounds which are in search of electrons. Such reagents are called electrophilic reagents. The presence of weaker (π) -bond makes alkenes unstable molecules in comparison to alkanes and thus, alkenes can be changed into single bond compounds by combining with the electrophilic reagents. Strength of the double bond (bond enthalpy, 681 kJ mol⁻¹) is greater than that of a carbon-carbon single bond in ethane (bond enthalpy, 348 kJ mol⁻¹). Orbital diagrams of ethene molecule are shown in Figure.



Geometrical isomerism: Doubly bonded Carbon atoms have to satisfy the remaining two Valences by joining with two atoms or groups. If the two atoms or groups attached to each Carbon atom are different, they can be Represented by YX C = C XY like structure. YX C = C XY can be represented in space in the Following two ways:



In (a), the two identical atoms i.e., both the X or both the Y lie on the same side of the Double bond but in (b) the two X or two Y lie Across the double bond or on the opposite Sides of the double bond. This results in Different geometry of (a) and (b) i.e. disposition Of atoms or groups in space in the two Arrangements is different. Therefore, they are Stereoisomers. They would have the same Geometry if atoms or groups around C = Cbond Can be rotated but rotation around C = C bond Is not free. It is restricted. For understanding This concept, take two pieces of strong Cardboards and join them with the help of two Nails. Hold one cardboard in your one hand And try to rotate the other. Can you really rotate The other cardboard? The answer is no. The Rotation is restricted. This illustrates that the Restricted rotation of atoms or groups around The doubly bonded carbon atoms gives rise to Different geometries of such compounds. The Stereoisomers of this type are called Geometrical isomers. The isomer of the type (a), in which two identical atoms or groups lie On the same side of the double bond is called Cis isomer and the other isomer of the type (b), in which identical atoms or groups lie on The opposite sides of the double bond is called Trans isomer. Thus cis and trans isomers Have the same structure but have different Configuration (arrangement of atoms or groups In space). Due to different arrangement of Atoms or groups in space, these isomers differ In their properties like melting point, boiling Point, dipole moment, solubility etc. Geometrical or cis-trans isomers of but-2-ene Are represented below:

Cis form of alkene is found to be more polar Than the trans form. For example, dipole Moment of cis - but - 2-ene is 0.33 Debye, Whereas, dipole moment of the trans form Is almost zero or it can be said that trans - but - 2 -ene is non-polar. This can be understood by drawing geometries of the two forms as given below from which it is clear that in the trans - but - 2 -ene, the two methyl groups are in opposite directions, Therefore, dipole moments of C - CH_3 bonds cancel, thus making the trans form non-polar.

In the case of solids, it is observed that the trans isomer has higher melting point than the cis form. Geometrical or cis-trans isomerism is also shown by alkenes of the types XYC = CXZ and XYC = CZW

Preparation – From alkynes: Alkynes on partial reduction with calculated amount of dihydrogen in the presence of palladised charcoal partially deactivated with poisons like sulphur

compounds or quinoline give alkenes. Partially deactivated palladised charcoal is known as Lindlar's catalyst. Alkenes thus obtained are having cis geometry. However, alkynes on reduction with sodium in liquid ammonia form trans alkenes.

ii)
$$RC \equiv CR^1 + H_2 \xrightarrow{Na/liquid NH_3} C = C$$
Alkyne
$$H$$

$$trans-Alkene$$

iii)
$$CH \equiv CH + H_2 \xrightarrow{Pd/C} CH_2 = CH_2$$
 (13.32)
Ethyne Ethene

iv)
$$CH_3-C\equiv CH+H_2 \xrightarrow{Pd/C} CH_3-CH=CH_2$$

Propyne Propene

From alkyl halides: Alkyl halides (R-X) on heating with alcoholic potash (potassium hydroxide dissolved in alcohol, say, ethanol) eliminate one molecule of halogen acid to form alkenes. This reaction is known as dehydrohalogenation i.e., removal of halogen acid. This is example of $\beta-$ elimination reaction, since hydrogen atom is eliminated from the β carbon atom (carbon atom next to the carbon to which halogen is attached).

$$H = H$$

$$H = C$$

$$H =$$

Nature of halogen atom and the alkyl group determine rate of the reaction. It is observed that for halogens, the rate is: iodine > bromine > chlorine, while for alkyl groups it is: tert > secondary > primary.

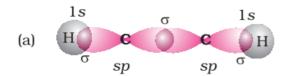
Physical properties Alkenes as a class resemble alkanes in physical properties, except in types of isomerism and difference in polar nature. The first three members are gases, the next fourteen are liquids and the higher ones are solids. Ethene is a colourless gas with a faint sweet smell. All other alkenes are colourless and odourless, insoluble in water but fairly soluble in non- polar solvents like benzene, petroleum ether. They show a regular increase in boiling point with increase in size i.e., every -CH₂ group added increases boiling point by 20–30 K. Like alkanes, straight chain alkenes have higher boiling point than isomeric branched chain compounds.

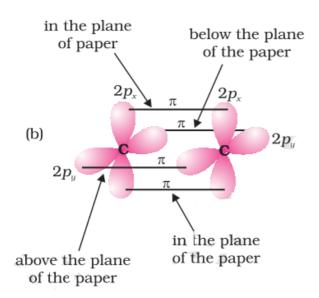
- i. The first three members of alkenes are ...?
 - a. Gases
 - b. Liquids

- c. Solids
- d. None of above
- ii. General formula for alkenes is?
 - a. C_nH_{2n+1}
 - b. C_nH_{2n}
 - c. C_nH_{2n-1}
 - d. C_nH_{2n+2}
- iii. The colour of ethene gas is ...?
 - a. Red
 - b. White
 - c. Pale Green
 - d. None of above
- iv. The bond length of carbon carbon double bond is ... pm 3
 - a. 154
 - b. 143
 - c. 134
 - d. 120
- v. Alkenes are also knows as ...?
 - a. Olefines
 - b. Paraffines
 - c. Oleofines
 - d. Paracetofines
- 136. Read the passage given below and answer the following questions from 1 to 5.

Alkynes– Like alkenes, alkynes are also unsaturated hydrocarbons. They contain at least one triple bond between two carbon atoms. The number of hydrogen atoms is still less in alkynes as compared to alkenes or alkanes. Their general formula is C_nH_{2n-2} . The first stable member of alkyne series is ethyne which is popularly known as acetylene. Acetylene is used for arc welding purposes in the form of oxyacetylene flame obtained by mixing acetylene with oxygen gas. Alkynes are starting materials for a large number of organic compounds. Hence, it is interesting to study this class of organic compounds.

Structure of Triple Bond Ethyne is the simplest molecule of alkyne series. Structure of ethyne is shown in Figure. Each carbon atom of ethyne has two sp hybridised orbitals. Carbon-carbon sigma (σ) bond is obtained by the head-on overlapping of the two sp hybridised orbitals of the two carbon atoms. The remaining sp hybridised orbital of each carbon atom undergoes overlapping along the internuclear axis with the 1s orbital of each of the two hydrogen atoms forming two C-H sigma bonds. H-C-C bond angle is of 180° .





Each carbon has two unhybridised p orbitals which are perpendicular to each other as well as to the plane of the C - C sigma bond. The 2p orbitals of one carbon atom are parallel to the 2p orbitals of the other carbon atom, which undergo lateral or sideways overlapping to form two pi (π) bonds between two carbon atoms. Thus ethyne molecule consists of one C - C σ bond, two C - H σ bonds and two C - C π bonds. The strength of C π C bond (bond enthalpy 823 kJ mol⁻¹) is more than those of C = C bond (bond enthalpy 681 kJ mol⁻¹) and C - C bond (bond enthalpy 348 kJ mol⁻¹). The C π C bond length is shorter (120 pm) than those of C = C (133 pm) and C - C (154 pm). Electron cloud between two carbon atoms is cylindrically symmetrical about the internuclear axis. Thus, ethyne is a linear molecule.

Preparation – From calcium carbide: On industrial scale, ethyne is prepared by treating calcium carbide with water. Calcium carbide is prepared by heating quick lime with coke. Quick lime can be obtained by heating limestone as shown in the following reactions:

$$\operatorname{CaO} + \operatorname{3C} \longrightarrow \operatorname{CaC}_2 + \operatorname{CO}$$

$$\operatorname{Calcium}$$

$$\operatorname{carbide}$$

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

From vicinal dihalides: Vicinal dihalides on treatment with alcoholic potassium hydroxide undergo dehydrohalogenation. One molecule of hydrogen halide is eliminated to form alkenyl halide which on treatment with sodamide gives alkyne.

Aromatic hydrocarbon- These hydrocarbons are also known as 'arenes'. Since most of them possess pleasant odour, the class of compounds was named as 'aromatic compounds'. Most of such compounds were found to contain benzene ring. Benzene ring is highly unsaturated but in a majority of reactions of aromatic compounds, the unsaturation of benzene ring is retained. However, there are examples of aromatic hydrocarbons which do not contain a benzene ring but instead contain other highly unsaturated ring. Aromatic compounds containing benzene ring are known as benzenoids and those not containing a benzene ring are known as non-benzenoids. Some examples of arenes are given below:

$$+ CH_{3}Cl \xrightarrow{Anhyd. AlCl_{3}} + HCl$$

$$+ CH_{3}Cl \xrightarrow{Anhyd. AlCl_{3}} + HCl$$

$$+ C_{2}H_{5}Cl \xrightarrow{Anhyd. AlCl_{3}} + HCl$$

Friedel-Crafts alkylation reaction: When benzene is treated with an alkyl halide in the presence of anhydrous aluminium chloride, alkylbenene is formed.

Friedel-Crafts acylation reaction: The reaction of benzene with an acyl halide or acid anhydride in the presence of Lewis acids (AlCl₃) yields acyl benzene.

$$+ CH_{3}COC1 \xrightarrow{Anhyd. AlCl_{3}} + HCl$$

$$+ CH_{3}COC1 \xrightarrow{Anhyd. AlCl_{3}} + HCl$$

$$+ (CH_{3}CO)_{2}O \xrightarrow{Anhyd. AlCl_{3}} + CH_{3}COOH$$

If excess of electrophilic reagent is used, further substitution reaction may take place in which other hydrogen atoms of benzene ring may also be successively replaced by the electrophile. For example, benzene on treatment with excess of chlorine in the presence of anhydrous $AICI_3$ can be chlorinated to hexachlorobenzene (C_6CI_6)

$$+ 6Cl_{2} \xrightarrow{Anhyd. AlCl_{3}} Cl Cl + 6HCl$$

$$+ 6Cl_{2} \xrightarrow{Anhyd. AlCl_{3}} Cl Cl + 6HCl$$

$$+ 6$$

- i. The general formula of Alkynes is ...?
 - a. C_nH_{2n-2}
 - b. C_nH_{2n-2}
 - c. C_nH_{2n-2}
 - d. C_nH_{2n-2}
- ii. Calcium carbide is prepared by heating quick lime with ...?
 - a. Backing soda
 - b. Coke
 - c. Carbide
 - d. Salt
- iii. Vicinal dihalides on treatment with alcoholic potassium hydroxide undergo ...?
 - a. Dehydrogenation.
 - b. Hydrohalogenation.
 - c. Dehydrohalogenation
 - d. Dehalogenation.
- iv. The bond enthalpy of $C \equiv C$ is ...?
 - a. 523 kJ mol
 - b. 623 kJ mol
 - c. 723 kJ mol
 - d. 823 kJ mol
- v. The $C \equiv C$ bond length is ...?

- a. 1200 pm
- b. 620 pm
- c. 240 pm
- d. 120 pm

* Given Section consists of questions of 5 marks each.

[70]

- 137. 8 Arrange benzene, n-hexane and ethyne in decreasing order of acidic behaviour. Also give reason for this behaviour.
- 138. What are the necessary conditions for any system to be aromatic?
- 139. Write chemical equations for combustion reaction of the following hydrocarbons:
 - i. Butane.
 - ii. Pentene.
 - iii. Hexyne.
 - iv. Toluene.
- 140. Out of benzene, m-dinitrobenzene and toluene which will undergo nitration most easily and why?
- 141. Why is benzene extra ordinarily stable though it contains three double bonds?
- 142. 4 In the alkane $H_3C-CH_2-C(CH_3)_2-CH_2-CH(CH_3)_2$, identify 1°,2°,3° carbon atoms and give the number of H atoms bonded to each one of these.
- 143. Write down the products of ozonolysis of 1,2-dimethylbenzene (o-xylene). How does the result support Kekulé structure for benzene?
- 144. Why is Wurtz reaction not preferred for the preparation of alkanes containing odd number of carbon atoms? Illustrate your answer by taking one example.
- 145. Draw the cis and trans structures of hex-2-ene. Which isomer will have higher b.p. and why?
- 146. 6 An alkene 'A' contains three C-C, eight C-H σ bonds and one C-C π bond. 'A' on ozonolysis gives two moles of an aldehyde of molar mass 44 u. Write IUPAC name of 'A'.
- 147. How do you account for the formation of ethane during chlorination of methane?
- 148. An alkene 'A' on ozonolysis gives a mixture of ethanal and pentan-3-one. Write structure and IUPAC name of 'A'.
- 149. Propanal and pentan-3-one are the ozonolysis products of an alkene? What is the structural formula of the alkene?
- 150. Addition of HBr to propene yields 2-bromopropane, while in the presence of benzoyl peroxide, the same reaction yields 1-bromopropane. Explain and give mechanism.

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		4		chan	ge you	life					