ORGANIC CHEMISTRY - SOME BASIC PRINCIPLES AND TECHNIQUES

CHEMISTRY





d) All of the above





- c) 2-methylpent-2-en-4-yne
- d) 4-methylpent-1-en-4-yne
- 40. Which of the following is elimination reaction





$$\begin{array}{c} CH_{3} \leftarrow \downarrow \leftarrow \downarrow \\ CH_{3} \leftarrow \downarrow \leftarrow \downarrow \\ COOH \\ \hline CH_{3} \leftarrow \downarrow \\ COOH \\ \hline CH_{3} \leftarrow \downarrow \\ COOH \\ \hline CH_{2} \leftarrow \downarrow \\ COOH \\ \hline CH_{2} \leftarrow \downarrow \\ CH_{2} \leftarrow \\ CH_{2} \leftarrow \\ CH_{2} \leftarrow \downarrow \\ CH_{2} \leftarrow \\$$

c) 3 a) 2 b) 4

59. "The negative part of the addendum adds on the carbon atom joined to the least number of hydrogen atoms." This statement is called : a) Markownikoff's rule b) Peroxide effect c) Baeyer's strain theory d) Thiele's theory 60. The total number of isomeric carbocations possible for the formula $C_4H_9^+$ is : a) 3 b) 4 c) 2 d) 5 61. The correct order for homolytic bond dissociation energies. (ΔH in kcal/mol) for CH₄(A), C₂H₆(B) and CH₃Br(C), under identical experimental conditions a) C>B>A b) B > C > Ac) C>A>B d) A > B > C62. The sodium extract of an organic compound on treatment with FeSO₄ solution, FeCl₃ and HCl gives a red solution. The organic compound contains a) Both nitrogen and sulphur b) Nitrogen only c) Sulphur only d) Halogen 63. *d*-tartaric acid and *l*-tartaric acid are : a) Structureal isomers b) Diastereoisomers c) Tautomers d) Enantiomers 64. Which of the following is a pair of functional isomers? a) CH₃COCH₃, CH₃CHO b) C₂H₅CO₂H, CH₃CO₂CH₃ c) $C_2H_5CO_2H$, $CHCO_2C_2H_5$ d) CH₃CO₂H, CH₃CHO 65. Which of the following is an optically active compound? a) Lactic acid b) Chloro acetic acid c) Meso-tartaric acid d) Acetic acid 66. Give the correct IUPAC name for CH_3 CH₃. CH₂OCH. CH₂. CH₂. CH₂Cl a) 2-ethoxy-5-chloropentane b) l-chloro-4-ethoxy-4-methylbutane d) Ethyl-1-chloropentylether c) 1-chloro-4-ethoxypentane 67. The IUPAC name of the compound, $\begin{array}{c} CH_2 & -- CH & -- CH_2 \\ I & I & I \\ OH & OH & OH \end{array}$ a) 1,2,3-trihydrosypropane b) 3-hydroxypentane-1,5-diol c) 1,2,3-hydroxypropane d) Propane-1,2,3-triol 68. Bond energy with the increase in number of lone pairs on the bonded atoms. a) Decreases c) Does not change d) None of these b) Increases 69. A liquid decomposes at its normal boiling point. It can be purified by a) Sublimation b) Steam distillation c) Vacuum distillation d) Fractional distillation 70. On monochlorination of 2-methyl butane, the number of chiral compounds formed are : a) 2 b) 4 d) 8 c) 6 71. Stability of which intermediate is not governed by hyperconjugation? a) Carbon cation b) Carbon anion c) Carbon free radical d) None of these 72. The ammonia evolved from the treatment of 0.30g of an organic compound for the estimation of nitrogen was passed in 100mL of 0.1M sulphuric acid. The excess of acid required 20mL of 0.5 M sodium hydroxide solution for complete neutralisation. The organic compound is a) Acetamide b) Benzamide d) Thiourea c) Urea 73. Conversion of CH_4 to CH_3Cl is an example of which of the following reaction? a) Electrophilic substitution b) Free radical addition

	c) Nucleophilic substitution		d) Free radical substitution	
74.	Number of possible	isomers of glucose are :		
	a) 10	b) 14	c) 16	d) 20
75.	The reaction			
	$CH_3CH_2CHCH_3 \xrightarrow{NaNH}$	$\stackrel{_2}{\rightarrow}$ Butene-1 and butane -2 (majo	or)	
	Br			
	The correct stateme	nt (s) are		
	a) 2-butene is Saytze	eff product	b) 1-butene is Hofmanı	n (s) product
	c) The elimination r	eaction follows Saytzeff rule	d) All of the above	
76.	Consider the followi	ng carbanions		
	\frown			
	(II) $O_2 N \rightarrow \overline{C}$	H ₂		\sim
			~	
			~	*
	Correct order of stat	oility is		
	a) I>II>III	b) III>II>I	c) II>III>I	d) I>III>II
77.	The stability of 2,3-d	limethyl but-2-ene is more than	1 2-butene. This can be ex	plained in terms of :
	a) Resonance	b) Hyperconjugation	c) Electromeric effect	d) Inductive effect
78.	Protin solvent is			
-0	a) Diethyl ether	b) <i>n</i> -hexane	c) Acetone	d) Ethanol
79.	. Addition of Br ₂ on <i>trans</i> -butene-2 gives :			
	a) A racemic mixture	e of 2,3-dibromobutane		
	b) Meso form of 2,3-	dibromobutane		
	c) Dextro form of 2,3	3-dibromobutane		
00	d) Laevo form of 2,3	-dibromobutane	and an of no other with all	atuan hili a uga gan dia
80.		NO ₂	order of reaction with ele	ectrophilic reagand is
	I II	III		
	a) II>III>I	b) III <i<ii< th=""><th>c) I>II>III</th><th>d) I=II>III</th></i<ii<>	c) I>II>III	d) I=II>III
81.	During AgNO ₃ test for	or detection of halogens, sodium	n extract is boiled with fe	w drops of conc. HNO ₃ to
	decompose			
	a) NaCN	b) Na ₂ S	c) Both (a) and (b)	d) None of these
82.	Which is true about	following?		
	CH ₃	CH ₃		
C	$C_2H_5 - C_H - C_3H_7$	$CH_3 - C - CH - C_2H_5$		
	(I)	O (II)		
	н	(11)		
		$C_2H_5 - CH - C_2H_5$		
	H − Ċ [♥]	L CH ₂		
	I H			
	(III)	(1V)		
	a) Only III is a chiral compound			

b) Only II and IV are chiral compounds c) All four are chiral compounds d) Only I and II are chiral compounds 83. How many chiral compounds are possible on monochlorination of 2-methyl butane? b) 4 a) 2 c) 6 d) 8 84. How many isomers of $C_5H_{11}OH$ will be primary alcohols? a) 5 b) 4 c) 3 d) 2 85. The epoxide ring consists of which of the following? a) Three membered ring with two carbon and one oxygen b) Four membered ring with three carbon and one oxygen c) Five membered ring with four carbon and one oxygen. d) Six membered ring with five carbon and one oxygen. 86. The reaction which is not the example of nucleophilic substitution among the following is a) $CH_3C - Br + CH_3OH \rightarrow CH_3C - OCH_3 + HBr$ b) $CH_3C - Cl + aq. KOH \rightarrow CH_3C - OH + KCl$ c) CI + alc. KOH \rightarrow + KCl + H₂O d) H = KOH \rightarrow COH + HO 87. Consider the following reaction >C=O + H₂NOH \rightarrow >C=NOH +H₂O Is an example of a) Substitution b) Elimination c) Addition d) Addition elimination 88. An important chemical method to resolve a racemic mixture makes use of the formation of : c) Racemers a) *meso* compound b) Enantiomer d) diastereoisomers 89. Red colour complex ion formed on adding FeCl₃ to sodium extract when N and S both are present in organic compound is d) $[Fe(CN)_6]^{3-1}$ a) $[Fe(CN)_6]^{4-}$ b) $[Fe(CNS)]^{2+}$ c) $[Fe(CNS)_2]^+$ 90. (I) $CH_3CH_2Br \xrightarrow{LAH} C_2H_6and$ (II) $(CH_3)_3 CBr \xrightarrow{LAH}$ alkene, The reason for this is a) (I)S_N2(II) E₁ mechanism b) (I) S_N1, (II) E₂ mechanism c) (I)S_N1, (II) E_1 mechanism d) (I) S_N2,(II) E₂ mechanism 91. How many σ and π -bonds are there in the molecule of tetracyanoethylene? a) 9σ and 9π b) 5σ and 9π c) 9σ and 7π d) 5σ and 8π 92. Which of the following complex formation indicates presence of sulphur in the organic compound when sodium nitroprusside is added to sodium extract of the compound? a) $Fe_4[Fe(CN)_6]_3$ b) $Na_2[Fe(NO)(CN)_5]$ c) $Fe_4(CNS)_3$ d) Na_4 [Fe(CN)₅NOS] 93. Who pointed out the concept hyperconjugation? a) Nathan and Baker b) Mullikan c) Kekule d) Kolbe 94. Alkyl halide can be converted into alkene by a) Nucleophilic substitution reaction b) Elimination reaction c) Both nucleophilic substitution and elimination reaction d) Rearrangement

95. The order of reactivities of the following alkyl halides for a $S_N 2$ reaction is :

- a) RF > RCl > RBr > RI
- b) RF > RBr > RCl > RI
- c) RCl > RBr > RF > RI d) RI > RBr > RCl > RF

96. The optically active alkane with lowest molecular weight is :

	j i i j i i i i i		II	
	а) СН ₃ СН ₂ С СН	b) CH ₃ CH ₂ —CH ₃	-CH ₃ c) CH ₃ $- \overset{n}{\overset{l}{_{C_2H_5}}}$	d) CH ₃ CH ₂ ·CH ₂ CH ₃
9	7. Which type of isomeris	sm is most common am	ong ethers?	
	a) Metamerism	b) Functional	c) Chain	d) Position
98	8. With a change in hybri	disation of the carbon l	bearing the charge, the stab	ility of a carbanion increase in the
	order			\frown
	a) sp < sp ² < sp ³	b) sp $< sp^3 < sp^2$	c) $sp^3 < sp^2 < sp$	d) $sp^2 < sp < sp^3$
99	9. A molecule is R_3C — H	I. If H is replaced by Z	$R(R_3C-Z)$ and on doing so	b electron density on R_3 – C part
	increases, then Z is :			
	a) Electron attracting g	group		
	b) Electron withdrawin	ng group		
	c) Electron repelling g	roup		
	d) Either of the above			
10	00. Which of the following	compounds are not ar	ranged on order of decreasi	ng reactivity towards
	electrophilic substituti	on?	4	\mathbf{N}
	a) Fluorobenzene > ch	lorobenzene > bromo	benzene	
	b) Phenol> <i>n</i> -propyl b	enzene> benzoic acid>	>	×
	c) Chlorotoluene >par	a-nitrotoluene>2-chlo	oro-4-nitro toluene	
	d) Benzoic acid> phen	ol>n-propyl benzene		
1	01. A mixture of camphor a	and benzoic acid can be	e separated by	
	a) Sublimation		b) Extraction with a	a solvent
	c) Chemical method		d) Fractional crysta	llisation
1	02. Resonance in benzene	is accompanied by dele	ocalisation of π -electrons. E	ach π -electron is attached with :
	a) 4 carbon	b) 2 carbon	c) 3 carbon	d) 6 carbon
10	03. Grignard reagent adds	to		
	a) >c==0	b) $-C \equiv N$	c) >c==s	d) All of these
10	04. Resonance energy is m	ore for		
	a) C ₆ H ₆		b) Cylohexene	
	c) Cycloheptene		d) Cyclohexa -1,2,3	-triene
1	05. The reaction			
	H + I	ICN		
	is an example of			
	a) Electrophilic additio	on	b) Electrophilic sub	ostitution
C	c) Nucleophilic substit	ution	d) Nucleophilic add	lition
1	06. Which of the following	is not chiral?		
	a) 3-bromopentane			
	b) 2-hydroxy propanoi	c acid		
	c) 2-butanol			
	d) 2,3-dibromopentane	e		
10	07. The structures $(CH_3)_3$	CBr and $CH_3[CH_2]_3$ Br	represent	
	a) Chain isomerism		b) Position isomeri	sm
	c) Chain as well as pos	ition isomerism	d) Functional isome	erism









159. Anti-Markownikoff addition of HBr is not observed in :









189. The number of optical isomers of $CH_3CH(OH)CH(OH)CHO$ is

a) Zero b) 2 c) 3 d) 4 190. The ratio of σ -to π -bonds in benzene is: a) 2 b) 4 c) 6 d) 8 191. In a $S_N 2$ substitution reaction of the type $R - Br + Cl^{-} \xrightarrow{DMF} R - Cl + Br^{+}$ Which one of the following has the highest relative rate? CH_3 —CH— CH_2Br + Cla) ĊH₃ $C - CH_2Br$ b) CH₃c) CH₃CH₂Br d) $CH_3 - CH_2 - CH_2Br$ 192. Hyperconjugation is a) $\sigma - \pi$ delocalisation b) No bond resonance c) $\sigma - \pi$ odd electron d) All of these 193. Which one of the following reactions is a condensation reaction? a) HCHO \rightarrow para-formaldehyde b) CH₃CHO \rightarrow para-aldehyde c) $CH_3COCH_3 \rightarrow$ mesityl oxide d) $CH_2 = CH_2 \rightarrow polyethylene$ 194. Which group has the maximum-Inductive effect? c) -COOH a) $-N0_2$ b) – CN d) – F 195. The correct IUPAC name of the following compound is $CH_3CH_2CH - C = C - CHCH_2CH_2CH_2CH_3$ Т CH_3 C_2H_5 C_2H_5 b) 6-butyl-5-ethyl-3-methyl oct-4-ene a) 5, 6-dimethyl-8-methyl dec-6-ene c) 5, 6-diethyl-3-methyl dec-4-ene d) 2, 4, 5-triethyl non-3-ene 196. Which is incorrect about enantiomorphs? a) They rotate the plane of polarized light in different directions b) They have mostly identical physical properties c) They have same configuration d) They have different biological properties 197. Which one is the seniormost functional group in the nomenclature of an organic compound if it possesses more than one functional group? a) —CHO b) —COOH c) — OH d) CO 198. The absolute configuration of the following CH₃ CI , are C_2H_5 are a) 2S, 3R b) 2S, 3S d) 2R, 3R c) 2R, 3S 199. Which step is chain termination step in the following mechanism?

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(i) $Cl_2 \xrightarrow{hv} Cl^{\bullet} + Cl^{\bullet}$ (ii) $Cl^{\bullet} + CH_4 \longrightarrow CH_3 + HCl$ (iii) $\overset{\bullet}{CH}_3 + Cl_2 \longrightarrow CH_3Cl + Cl^{\bullet}$ (iv) $Cl^{\bullet} + CH_{3} \longrightarrow CH_{3}CH_{3$ a) (i) b) (ii) c) (iii) d) (iv) 200. The reaction intermediate produced, by homolytic cleavage of a bond is called a) Carbene b) Carbocation c) Carbanion d) Free redical 201. Fractional distillation is useful in distillation of d) All of these a) Petroleum b) Coal-tar c) Crude alcohol 202. Which of the following species is paramagnetic? b) A free radical d) All of these a) A carbocation c) A carbanion ion 203. Sulphur trioxide is : a) An electrophile b) A nucleophile c) A homolytic reagent d) A base 204. In Kjeldahl's method, ammonia from 5g of food neutralizes 30 cm^3 of 0.1 N acid. The percentage of nitrogen in the food is a) 0.84 b) 8.4 c) 16.8 d) 1.68 205. The number of isomeric alkanes having the molecular formula $C_5 H_{12}$ is b) Five d) Thirty two a) Three c) Nine 206. Select the organic compound which was prepared for the first time in laboratory from its elements : c) C_2H_5OH a) Urea b) CH₃COOH d) None of these 207. Which of the following compounds can exist in optically active form? c) 3-pentanol a) 1-butanol b) 2-butanol d) 4-heptanol 208. The compound in which carbon uses only its sp^3 hybrid orbitals for bond formation is a) $(CH_3)_3COH$ b) HCOOH c) CH₃CHO d) $(H_2N)_2CO$ 209. How many types of functional group can be present in an amine with the formula C_3H_9N ? a) 1 b) 2 c) 3 d) 4 210. Select the most reactive cycloalkane b) Cyclobutane a) Cyclopropane c) Cyclopentane d) Cyclohexane 211. The -I effect is shown by : a) –COOH c) $-CH_3CH_2$ b) - CH₃ d) $-CHR_2$ 212. The stability of carbanions in the following; (1) $-RC = \vec{C}$ (2)(3) R_2C (4) $R_3C - \breve{C}H_2$ is in the order of : a) (2) > (3) > (4) > (1)b) (4) > (2) > (3) > (1)c) (1) > (3) > (2) > (4)d) (1) > (2) > (3) > (4)213. Glyoxal is a) $CH_2OH - CH_2OH$ b) CHO $- CH_2OH$ c) COOH – CO – COOH d) CHO - CHO 214. IUPAC name of acraldehyde is b) Propenyl aldehyde a) But-3-en-1-al c) But-2-ene-1-al d) Prop-2-en-1-al

215. The IUPAC name of $CH_3 - C \equiv CH(CH_3)_2$ is a) 4-methyl-2-pentyne b) 4, 4-dimethyl-2-butyne c) methyl isopropyl acetylene d) 2-methyl-4-pentyne 216. What information is provided by reaction mechanism? a) The bonds broken and formed b) The reaction intermediates c) The relative rates of discrete steps, especially the slowest one d) All of the above 217. The enolic from of acetone contains a) 8σ bonds, 2π -bonds and 1 lone pair b) 9σ bonds, 1π -bonds and 2 lone pairs c) 9σ bonds, 2π -bonds and 1 lone pair d) 10σ bonds, 1π -bonds and 1 lone pair 218. Which of the following acids has the smallest dissociation constant? d) CH₃CHBrCOOH c) BrCH₂CH₂COOH a) CH₃CHFCOOH b) FCH₂CH₂COOH 219. IUPAC name of, CH₃CH(OH)CH₂CH₂COOH is : a) 4-hydroxypentanoic acid b) 1-carboxy-3-butanoic acid c) 1-carboxy-4-butanol d) 4-carboxy-2-butanol 220. The number of isomers for the compound with the molecular formula C₂BrClFI is a) 3 b) 4 c) 5 d) 6 221. Among the following the strongest nucleophile is c) CH₃NH₂ a) C_2H_5SH b) CH₃COO⁻ d) NCCH $_{2}^{-}$ 222. Which of the following statements is correct? b) +I group stabilises a carbanion a) +I group stabilises a carbocation c) -I group stabilises a carbocation d) -I group stabilises a free radical 223. Which of the following species does not exert a resonance effect? b) C₆H₅NH₃ c) C_6H_5OH a) $C_6H_5NH_2$ d) C_6H_5Cl 224. The number of different amines corresponding to the formula C₃H₉N is : a) 2 b) 3 c) 4 d) 5 225. Lactic acid molecule has a) One chiral carbon atom b) Two chiral carbon atoms c) No chiral carbon atom d) asymmetric molecule 226. The arrangement of atoms that characterises a particular stereoisomer is called : b) Configuration a) Geometry of isomer d) None of these c) Conformers 227. An alkane forms isomers if the number of carbon atoms is : a) ≥ 1 b) ≥ 2 c) ≥ 3 d) ≥ 4 228. Which of the following statements is correct? a) Desmotropism is another name for tautomerism b) Allyl carbocation is less stable than isopropyl carbocation c) –*I* effect is exhibited by– $^{+}_{NH_3}$ d) The formula CH_2Cl_2 is non-polar 229. The IUPAC name of $CH_3COCH(CH_3)_2$ is a) Isopropylmethyl ketone b) 2-methyl-3-butanone c) 4-methylisopropyl ketone d) 3-methyl-2-butanone 230. Qualitative test of halogens in an organic compound is made by a) Fleming's test b) Beilstein test c) Bayer's test d) Fehling's test 231. Which one of the following will most readily be dehydrated in acidic conditions?





2			
3 4			
\int_{5} is :			
6			
a) 3-(1-ethyl propy	l) hex-1-ene		
b) 4-Ethyl-3-propyl	hex-1-ene		
c) 3-Ethyl-4-etheny	'l heptane		\frown
d) 3-Ethyl-4-propyl	hex-5-ene		
255. IUPAC name of (CH	$_{3})_{2}N - C_{2}H_{5}$ is :		
a) Dimethyl ethyl ar	mine		
b) Dimethylaminon	iethane		
c) Dimethylaminoe	thane		
d) <i>N</i> , <i>N</i> -dimethyleth	ianamine		
256. Among the followin	g compounds, the most acidic	is	\sim
a) <i>p</i> -nitrophenol		b) <i>p</i> -hydroxybenzoic act	id
c) <i>o</i> -hydroxybenzo	ic acid	d) <i>p</i> -toluic acid	
257. Electrophiles are :			
a) Lewis bases	b) Lewis acids	c) Amphoteric	d) None of these
258. On monochlorination	on of <i>n</i> -pentane, the number o	f isomers formed is :	
a) 4	b) 3	c) 2	d) 1
259. Cyclohexane is :			
a) Aliphatic compo	und		
 b) Alicyclic compou 	nd	XY	
c) Aromatic compo	und	Y	
d) Heterocyclic com	ipound	Y	
260. Which of the follow	ing is a primary halide?		
a) Isopropyl iodide		b) Secondary butyl iodio	le
c) Tertiary butyl br	omide	d) Neo hexyl chloride	
261. The percentage of '	<i>s</i> ' character of the hybrid orb	oital of carbon in ethane, etl	nane and ethyne respectively
are :			
a) 25, 33, 50	b) 20, 50, 33	c) 25, 50, 75	d) 33, 66, 99
262. Which is a chiral mo	olecule?		
a) CH ₃ Cl	b) CH ₂ Cl ₂	c) CHBr ₃	d) CHClBrl
263. The stability of a ca	rbonium ion depends upon		
a) The bond angle of	if the attached group		
b) The substrate wi	th which it reacts		
c) The inductive eff	ect and hyper-conjugative effe	ect of the attached group	
d) None of the abov	e		
264. The IUPAC name of	the compound,		
CH ₃ -(CH ₂) ₄ -CH	$-\dot{C}$ - CH_2 - CH_3 is:		
▼ CH	(CH ₂) ₂ CH ₂		
a) 2.4 dimathul 2 m			
a) $3,4$ -almethyl- $3-n$	propymonane		
a) 6.7 dimethyl 7 m	nyuetane		
d) 6.7 dimethyl 7 a	propymonane thyldocano		
265 Bromination of all	unyinecane		
200. DI UIIIII ati UII OI dika	h) Carbocations	c) Carbonas	d) Free radicals
aj Jai Daillulis	uj carbucacions	cj cai belles	uj rice i duicais

266. The isomeric *cis*-2-butene and *trans*-2-butene can be distinguished on the basis of : a) Their physical nature b) Their reduction products c) The products they give on ozonolysis d) The products they give on addition to bromine 267. Lassaigne's test is not used for the detection of a) Carbon d) Sulphur b) Halogens c) Nitrogen 268. Consider the following carbocations, (I) $C_6H_5CH_2$ (II) $C_6H_5CH_2CH_2$ (III) $C_6H_5CHCH_3$ (IV) $C_6H_5C(CH_3)_2$ d) IV<III<I<II a) II<I<III<IV b) II<III<IV c) III<I<IIV 269. The simplest formula of a compound containing 50% of element *X* (at. wt 10) and 50% of element *Y* (at. wt. 20) is a) *XY* b) XY_2 c) X_2Y $X_{2}Y_{2}$ 270. *n*-pentane, iso-pentane, and *neo* – pentane are examples for isomers of the type b) Optical a) Geometrical c) Chain d) Positional 271. Homolytic fission of C— C bond in ethane gives an intermediate in which carbon ishybridized. b) sp^2 a) sp^3 d) sp^2d c) sp 272. Pick out the correct statement from the following and choose the correct answer from the codes given below I. Hexa-1, 5-diene is a conjugated diene II. Prop-1, 2-diene is conjugated diene III. Hexa-1, 3-diene is a conjugated diene IV. Buta-1, 3-diene is an isolated diene V. Prop-1, 2-diene is a cumulative diene b) II, III c) IV, V d) II, V a) I,II 273. The IUPAC name of CH₂ $CH_3 - CH - CH_2 - C - CH_3$ Т OH OH a) 1, 1-dimethyl-1, 3-butanediol b) 2-methyl-2, 4-pentanediol c) 4-methyl-2, 4-pentanediol d) l, 3, 3-trimethyl-1, 3-propane diol 274. Which among the following is the correct IUPAC name of isoamylene? a) 1-pentene b) 2-methyl-2-butene c) 3-methyl-1-butene d) 2-mythyl-1-butene 275. Which of the following compounds exhibits geometrical isomerism? a) C_2H_5Br b) $(CH)_2(COOH)_2$ c) CH₃CHO d) $(CH_2)_2(COOH)_2$ 276. Which one of the following compounds, is most acidic? OH a) $Cl - CH_2 - CH_2 - OH$ b) CH_3 NO₂ 277. An organic compound has carbon and hydrogen percentage in the ratio 6:1 and carbon and oxygen percentages in the ratio 3:4. The compound has the empirical formula d) CH_2O a) C_2H_6O b) CHO_2 c) CH₄O 278. Among the following the one which does not exhibit functional group isomerism is :

c) C_4H_{10}

a) $C_2 H_6 O$

b) $C_{3}H_{8}O$

d) $C_4 H_{10} O$

279. How many structural for	rmulae are possible for C ₅ H	1 ₁₁ Cl?	
a) 6	b) 8	c) 10	d) 12
280. Which one of the follow	ing is a secondary alcohol?		
a) 2-methyl-1-propanol	b) 2-methyl-2-propanol	c) 2-butanol	d) l-butanol
281. Among the following an	tions (a) $\overline{C}H_3(b) \overline{N}H_2, (c)O$	H^{-} , $(d)F^{-}$ the order of basi	city is :
a) $a > b > c > d$	b) $b > a > c > d$	c) $c > b > a > d$	d) $c > a > b > d$
282. Electromeric effect is			
a) Permanent effect	b) Temporary effect	c) Resonance effect	d) Inductive effect
283. In the following groups,	-OAC -OMe-OSO2Me-OS	50 ₂ CF ₃ the order of leaving	group ability is :
(1)			
284 The angle of rotation of	nlane of polarized light den	ends upon ·	
a) The nature of the light	t beam		
b) The number of the m	olecules		
c) The number of asymm	netric carbon atoms in the	molecule of the substance	
d) All of the above			\sim
285. Which of the following s	hows geometrical isomeris	m?	X
a) C_2H_5Br	b) (CH ₂)(COOH) ₂	c) (CH) ₂ (COOH) ₂	d) $C_2 H_6$
286. Which of the following c	annot undergo nucleophilio	substitution under ordina	ry conditions?
a) Chlorobenzene	b) <i>Tert</i> –butylchloride	c) Isopropyl chloride	d) None of these
287. The C— C bond length o	f the following molecules is	in the order.	
a) $C_2H_6 > C_2H_4 > C_6$	$H_6 > C_2 H_2$		
b) $C_2H_2 < C_2H_4 < C_6H$	$_{6} < C_{2}H_{6}$		
c) $C_6H_6 > C_2H_2 > C_2$	$H_6 > C_2 H_4$		
d) $C_2H_4 > C_2H_6 > C_2$	$H_2 > C_6 H_6$	X) ^x	
288. Isomerism among comp	ounds due to the migration	of a proton is known as :	
a) Geometrical	b) Optical	c) Tautomerism	d) Position
289. Removal of hydrogen at	om is easier when it is attac	ched to :	
a) 1° carbon	b) 2° carbon	c) 3° carbon	d) Same in all
290. The order of stability of	carbanions is :		
a) $CH_3^- > 1^\circ > 2^\circ > 3^\circ$	b) $3^{\circ} > 2^{\circ} > 1^{\circ} > CH_3^-$	c) $3^{\circ} > 1^{\circ} > 2^{\circ} > CH_{3}^{-}$	d) $2^{\circ} > 3^{\circ} > 1^{\circ} > CH_{3}^{-}$
291. Glycerine contains			
a) 1° carbon	$\mathbf{X}\mathbf{Y}$	b) 2° carbon	
c) 3° carbon		d) Both 1° and 2° carbon	L
292. Which of the following p	bairs of carbon skeletons in	an example of isomerism?	
Ĭ			
a) $C - \dot{C} - C - C$ and C	C-C-C		
	C		
b) C-C-C and C-C	—C		
\mathbf{O}	С		
c) $C = C = C$ and C	C - C - C		
	L C		
d) ^C			
C - C - C - C - C and C	-cc		
	l C		

293. In cannizzaro reaction given below

$$2PhCHO \xrightarrow{OH} PhCH_2OH + PhCO_2^{\Theta}$$

The slowest step is

- a) The attack of :: OH^{\ominus} at the carboxyl group
- b) The transfer of hydride to the carbonyl group
- c) The abstraction of proton from the carboxylic group
- d) The deprotonation of PhCH₂OH
- 294. In the following the most stable conformation of n-butane is :





317. Chlorobenzene is 0, *p*-directing in electrophilic substituting reaction. The directing influence is explained by

a) +M of Ph b) +I of Cl c) +M of Cl d) -I of Ph

a)
$$H_{1,C}(W_{1,C}) = H_{1,C}(W_{1,C}) = H_{1,C}($$

a)
$$\begin{array}{l} H_{2}C = CH = CH = CH \\ h_{3} \\ h_{4} = C \\ h_{5} \\ h_{5} \\ h_{5} \\ h_{5} \\ h_{6} \\ h_{5} \\ h_{6} \\ h_{7} \\$$

$$CH_2 = CH - CH(CH_2CH_3)C = CH_2$$

|
Br

a) 4-bromo-3-ethyl-1,4-pentadiene

- b) 2-bromo-3-ethyl-1,4pentadiene
- c) 2-bromo-3-ethyl-1-5-pentadiene
- d) None of the above
- 354. The product of reaction,

$$CH_3$$

 CH_3 — CH_2 — C — CH — CH_3 + HOCl—

product is :

a)
$$CH_{3}CH_{2}$$
 CH_{3} $CH_{3}CH_{2}$ CH_{3} $CH_{3}CH_{2}$ CH_{3} CH_{3}
b) CH_{3} CH_{2} CH_{2} CH_{3} CH_{3} CH_{3} CH_{3} CH_{3} CH_{2} CH_{3} CH_{3} CH_{3} CH_{2} CH_{3} CH_{3} CH_{3} CH_{2} $CH_$

ĊH₃

355. The most unlikely representation of resonance structures of *p*-nitrophenoxide ion is :



359. Which of the following is an example of substitution reaction?

$$a) \oint_{C_{H_{3}}} \underbrace{\operatorname{Me}_{C_{4}} \operatorname{Le}_{U}}_{U_{1}} \oint_{C_{H_{3}}} \oint_{C_{4}} \underbrace{\operatorname{D}}_{U_{4}} \oint_{C_{4}} \underbrace{\operatorname{D}}_{U_{4}} \oint_{U_{4}} \underbrace{\operatorname{D}}_{U_{4}} \bigoplus_{U_{4}} \underbrace{\operatorname{D}}_{U_{4}} \bigoplus_{U_{4}} \underbrace{\operatorname{D}}_{U_{4}} \bigoplus_{U_{4}} \underbrace{\operatorname{D}}_{U_{4}} \bigoplus_{U_{4}} \underbrace{\operatorname{D}}_{U_{4}} \bigoplus_{U_{4}} \underbrace{\operatorname{D}}_{U_{4}} \underbrace{\operatorname{D}}_{U$$





394. The increasing order of +ve *I*-effect shown by H, CH_3 , C_2H_5 and C_3H_7 is :



c)	Their melting point		d) Their solubility in achiral reagents	
406. Th	e number of chiral cent	res in (+) -glucose		
a)	4	b) 3	c) 2	d) 1
407. Hv	vdrogen cvanide and hv	drogen isocvanide are :	-)	
a)	Tautomers			
b)	Positional isomers			
c)	Metamers			
d)	Chain isomers			
408. W	hich of the following hy	drocarbons is most unsatu	rated?	
a)	C ₂ H ₄	b) C_2H_2	c) C ₂ H ₆	d) $CH_3CH = CH_2$
409. So	metimes the behaviour	of a compound is explaine	d by assuming that it exist	s in a world between two or
m	ore different possible st	ructures. This phenomeno	n is called :	
a)	Isomerism	b) Resonance	c) Mutarotation	d) Allotropism
410. Ho	w many primary amine	es are possible with the for	mula $C_4 H_{11} N$?	
a)	1	b) 2	c) 3	d) 4
411. W	hich one of the followin	g pairs represents stereois	omerism?	
a)	Geometrical isomerism	, position isomerism	Ć	
b)	Geometrical isomerism	, conformational isomerism	n	
c)	Optical isomerism, geor	metrical isomerism		
d)	Optical isomerism, met	amerism		
412. Th	e large number of organ	nic compounds is due to:		
a)	Catenation property of	carbon		
b)	Covalent bond formation	on		
c)	Isomerism		G, Y	
d)	polymerization	A	N [°]	
413. Th	ie IUPAC name of CH_3 -	$-C - CH - CH_3$ is		
			·	
		0 CH ₃		
a)	2-methyl-3-butanone		b) 3-methyl-butan-2-one	
c)	3-methyl butanone		d) None of these	
414. Fo	rmic acid is a stronger a	acid than acetic acid. This c	an be explained using	
a)	+Meffect	b) –/ effect	c) $+I$ effect	d) $-M$ effect
415. Th	e energy of C— C triple	bond in acetylene in kcal i	S :	
a)	140	b) 192	c) 60	d) 100
416. In	which of the following i	molecules, the resonance e	ffect is not present?	
a)		h H_{a}	с) ОН	
uj				
417. W	hich of the following rep	presents the given mode of	hybridization $sp^2 - sp^2 - p^2$	sp – sp from left to right?
a)	$CH_2 = CH - C \equiv CH$	b) HC \equiv C $-$ C \equiv N	c) $CH_2 = C - C = CH_2$	d) CH
418 Th	e Lassaigne's extract is	hoiled with dil HNO ₂ hefor	re testing for halogens beca	
2110. II.	AGCN is soluble in HNC		h) Silver halides are solut	has e
	Na ₂ S and NaCN are dec	romposed by HNO ₂	d) $Ag_{a}S$ is soluble in HNO	2
419. Th	e tautomeric form which	ch is less stable is called :		3
a)	Anion form	b) Cation form	c) Labile form	d) All of these
420. Th	e effect involving the o	complete transfer of a sha	red pair of electrons to or	ne of the atoms joined by a
m	multiple bond at the requirement of attacking reagent is called :			
a)	Inductive effect	b) Mesomeric effect	c) Electromeric effect	d) None of these
421. W	hich of the following aci	ds does not exhibit optical	isomerism?	,
a)	Lactic acid	b) Tartaric acid	c) Maleic acid	d) α-amino acids
422. Many organic compounds are prepared by using PCl₅ because : a) OH group of alcohol is easily replaced by chlorine atom b) Chlorines are added to the unsaturated compounds c) It removes water from organic compounds d) Phosphorus atoms are entered in the alcohol 423. Which of the following conformations of cyclohexane is chiral? a) Twist boat b) Rigid c) Chair d) Boat 424. Which type of isomerism is shown by propanal and propanone? c) Tautomerism a) Functional group b) Metamerism d) Chain isomerism 425. Identify the product in the given reaction: $CH_3 - CH = CH_2 + NOCI \rightarrow Product$ a) $CH_3CHCLCH_2$. NO b) CH₃CH(NO). CH₂Cl c) $CH_3CH_2CH(Cl)(NO)$ d) $CH_2(NO)$. CH_2 . CH_2Cl 426. A straight chain hydrocarbon has the molecular formula C_8H_{10} . The hybridization for the carbon atoms from one end of the chain to the other are respectively sp^3 , sp^2 , sp^2 , sp^3 , sp^2 , sp^2 , sp^2 , sp and sp. The structural formula of the hydrocarbon would be : a) $CH_3 - C \equiv C - CH_2 - CH = CH - CH = CH_2$ b) $CH_3 - CH_2 - CH = CH - CH = CH - CH \equiv CH$ c) $CH_3 - CH = CH - CH_2 - C \equiv C - CH = CH_2$ d) $CH_3 - CH = CH - CH_2 - CH = CH - C \equiv CH$ 427. Reaction, a) Electrophilic substitution b) Nucleophilic substitution c) Electrophilic addition d) Nucleophilic addition 428. The total number of acylic iosmers including the stereoisomers (geometrical and optical), with the molecular formula C₄H₇Cl is a) 12 b) 11 c) 10 d) 9 429. The best method to separate the mixture of *ortho* -and-para nitrophenol (1:1) is **b)** Colour spectrum c) Distillation a) Vaporisation d) Crystallisation 430. Which of the following does not show electrometric effect? a) Alkenes b) Ethers c) Aldehyde d) Ketones 431. Shifting of electrons of a multiple bond under the influence of a reagent is called : a) *I*-effect b) E-effect c) *M*-effect d) *T*-effect 432. 0.4 g of a silver salt of a monobasic orgainc acid gave 0.26 g pure silver on ignition. the molecular weight of the acid is (atomic weight of silver=108) a) 58 b) 37 c) 89 d) 105 433. The S_N2 mechanism for, $R - X + KOH(aq) \rightarrow R - OH + KX$ follows with : a) 100% inversion b) 50% inversion c) 40% inversion d) 30% inversion 434. IUPAC name of, is : a) 4,5-dimethyl oct-4-ene b) 3,4-dimethyl oct-5-ene c) 4,5-dimethyl oct-5-ene d) None of the above 435. The reaction,

 \cap •X + Nu⁻ – is fastest when X is a) OCOR b) OC_2H_5 c) NH₂ d) Cl 436. Which pair represents chain isomers? a) CH₃CHCl₂ and ClCH₂CH₂Cl b) Propyl alcohol and isopropyl alcohol c) 2-methylbutane and neopentane d) Diethylether and dipropylether 437. The empirical formula of an acid is CH_2O_2 , the probable molecular formula of the acid may be d) CH_2O_2 b) $C_3 H_6 O_4$ c) $C_2H_2O_4$ a) $C_2H_4O_2$ 438. The number of valence electrons in the excited carbon atom is : a) Two in s and two in p-orbitals b) 4 single *p*-orbitals c) One in s and three in p-orbitals d) None of the above 439. A hydrocarbon contains 10.5 g carbon and 1 g hydrogen. Its 2.4 g has 1 L volume at 1 atm and 127°C. Hydrocarbon is a) C_6H_7 b) C_6H_8 d) C_6H_6 c) $C_5 H_6$ 440. The number of stereoisomers possible for a compound of the molecular formula $CH_3 - CH = CH -$ CH(OH) - Me is a) 3 b) 2 d) 6 441. The structural formula of methyl aminomethane is : a) $(CH_3)_2 CHNH_2$ b) $(CH_3)_3N$ c) $(CH_3)_2NH$ d) CH₃NH₂ 442. A mixture of oil and water is separated b a) Filtration b) Fractional distillation c) Sublimation d) Using separating funnel 443. The stability of $CH_3 - CH = CH - CH_3, CH_3$ $C - CH_3$ CH₃ CH₃ (II) (I) $CH_3 - C = CH_2$ $CH_2 - C = CH - CH_2$ L CH₂ CH₃ (III) (IV)In the increasing order is a) III<I<IV<II d) II<III<IV<I b) I<II<III<IV c) IV<III<II<I 444. The number of optically active isomers of tartaric acid are a) 1 b) 3 c) 4 d) 2 445. The nodal plane in the π -bond of ethene is located in a) The molecular plane b) A plane parallel to the molecular plane c) A plane perpendicular to the molecular plane which bisects the carbon-carbon sigma bond at right angle

d) A plane perpendicular to the molecular plane which contains the carbon-carbon sigma bond 446. Which of the following has the highest degree of coordination bond?



a) Tautomers b) Metamers	c) Functional isomers	d) Geometrical isomers
460. A racemic mixture is a mixture of :		
a) <i>meso</i> and its isomers		
b) <i>d</i> -and its <i>l</i> -isomers in equal proportions		
c) <i>d</i> -and its <i>l</i> -isomers in different proportions		
d) <i>meso</i> and <i>d</i> -isomers		
461. A mixture of iodine and sodium chloride can b	e easily separated by	
a) Fractional distillation	b) Steam distillation	
c) Chromatography	d) Sublimation	
462. The property by virtue of which a compound c	an turn the plane of polarization	n of light is known as :
a) Photolysis b) Phosphorescence	e c) Optical activity	d) polarization
463. Correct order of stability is		
a) HC $\equiv \overline{C} > CH_2 = \overline{C}H > CH_3 - \overline{C}H_2$	b) $CH_3 - \overline{C}H_2 > CH_2 = 0$	$\overline{C}H > CH \equiv \overline{C}$
c) $CH_3 - \overline{C}H_2 > CH \equiv CH \cong CH_2 = \overline{C}H$	d) All are equally stable	
464. In the estimation of sulphur in an organic com	pound, fuming nitric acid is used	l to convert sulphur into
a) SO_2 b) H_2S	c) H_2SO_3	d) H_2SO_4
465. The IUPAC name of compound	Ċ	
0		
		-
$CH_2 - C - OH$		
CH2-COOH		
a) 1,2,3-tricarboxy-2,1-propane		
b) 3-carboxy-3-hydroxy-1,5-pentanedioic acid		
c) 3-hydroxy-3-carboxy-1,5-pentanedioic acid		
d) None of the above	Y	
466. Which of the following will be chiral?	$\mathbf{\mathcal{I}}$	
a) CH_3CHCl_2 b) $CH_3CHBrCl$	c) CD_2Cl_2	d) CH ₂ ClBr
^{467.} In the dehydration reaction $CH_3CONH_2 \xrightarrow{P_2O_5} CI$	$H_3C \equiv N$, the hybridization state	e of carbon change from
a) $lsp^3to sp^2$ b) $lsp to sp$	c) $lsp^2 to sp$	d) lsp to sp^3
468. The correct acidity order of the following is		
он он соон соон		
(I) (II) (III) (IV)		
a) $(III) > (IV) > (II) > (I)$ b) $(IV) > (III) > (I) > (IV) > (III) > (IV) > $	(II) c) (III)>(II)>(I)>(IV)	(II)<(VI)<(III)<(II)
469. Which of the following is not a nucleophile?		
a) BF_2 b) CN^-	c) OH-	d) NH ₂
470 Least stable conformer of cyclohexane is	0,011	
a) Chair b) Boat	c) Twist hoat	d) Planar hexagon
471 The compound isomeric with acetone is :	ej i wist bout	uj i lanar nexagon
a) Pronionaldehyde b) Pronionic acid	c) Ethoxy ethane	d) None of these
472 Which of the chloride is less reactive towards l	hvdrolvsis?	aj none or mese
a) Vinyl chlorida b) Allyl chlorida	c) Ethvl chlorida	d) t -butyl chloride
473 Chycerol is an alcohol which can be classified a	cj Ediyi chidride	
a) Tribudric b) Monobudric	s c) Dibudric	d) Hovahudria
a) ITHINGTON D MONONYUPIC		uj nexaliyullu
474 . Addition of Br_2 on $CH_2 = CH_2$ in presence of N	aci(uq.) gives :	

a) $CH_2Br. CH_2Br$ b) $CH_2Br. CH_2Cl$ c) CH₂Br. CH₂OH d) All of these 475. The electromeric effect in organic compounds is a : a) Temporary effect b) Permanent effect c) Temporary-permanent effect d) None of the above 476. The function of boiling the sodium extract with conc. HNO₃ before testing for halogen is a) To make the solution acidic b) To make the solution clear c) To convert Fe²⁺ to Fe³⁺ d) To destroy CN⁻ and S²⁻ ions 477. Copper wire test of halogens is known as a) Liebig's test b) Lassaigne's test c) Fusion test d) Beilstein's test 478. Which of the following is singlet carbine? c) CH₃ C HCH₃ a) $(CH_3)_3C^+$ b) C₂H₅ C–H 479. Which of the following will be easily nitrated? CH_3 NO_2 c) CH_3NO_2 a) 480. Optical isomerism is shown by a) Propanol-2 b) Butanol-2 c) Ethanol d) Methanol 481. Williamson's synthesis involves b) Nucleophilic addition a) S_N1 mechanism d) S_E mechanism c) S_N2 mechanism 482. Free radicals can undergo : a) Disproportionation to two species b) Rearrangement to a more stable free radical c) Decomposition to give another free radical d) All of the above are correct 483. During addition of bromine on ethene, the first species formed is Br^+ c) ⁺_{CH₂CH₂Br} b) $C_2H_4OH^+$ d) $C_2H_5^+$ a) CH₂-CH₂ 484. Metamers of ethyl propionate are a) C_4H_9COOH and $HCOOC_4H_9$ b) C₄H₉COOH and CH₃COOC₃H₇ c) CH₃COOCH₃ and CH₃COOC₃H₇ d) CH₃COOC₃H₇ and C₃H₇COOCH₃ 485. Which statement is correct about the hybridization of carbon atoms in, a) C_1 and C_4 are sp^2 -hybridized b) C_2 and C_3 are sp^2 -hybridized c) All are *sp*-hybridized d) All are sp^2 -hybridized 486. Which one is not is IUPAC system? -CH—CH—CH₃ (3-methyl-2-butanol) $\begin{bmatrix} I \\ OH \end{bmatrix}$ a)

 CH_3 CH-CH-CH₂CH₃ CH₃CH₂CH₂-ĊH₂CH₃ (3-methyl-4-ethyl heptane) -C-CH-CH₃ || | CH₂CH₃ c) (2-ethyl-3-methyl-but-1-ene) d) $CH_3 - C \equiv C - CH(CH_3)_2$ (4-methyl-2-pentyne) 487. The compound which exhibits optical isomerism is : d) CH₃CCl₂CH₂CH a) CH₃CHOHCH₃ b) $(CH_3)_2 CHCH_2 CH_3$ c) CH₃CHClCH₂CH₃ 488. Which of the following applies in the reaction $CH_3CHBrCH_2CH_3 \xrightarrow{Alco.KOH}$? $(I)CH_3CH = CHCH_3(Major product)$ $(II)CH_2 = CHCH_2CH_3$ (Minoe product) a) Hofmann's rule b) Savtzeff's rule c) Kharasch effect d) Markownikoff's rule 489. Homologous compounds have : a) Same chemical properties b) Same molecular weight c) Same physical properties d) Same m.p. and b.p. 490. How many chiral compound are possible on mono chlorination of 2-methyl butane? b) 4 d) 8 a) 2 491. Which of the following may exist in enantiomorphs? СН₃—СН—СООН | CH₃ a) b) $CH_3 = CHCH_2CH_2CH_3$ CH-CH₃ -CH₂—CH—CH₃ NH_2 492. How many isomers are possible for the alkane C_4H_{10} ? a) 3 b) 5 c) 2 d) 4 493. Which of the following IUPAC names is correct? a) 2-methyl-3-ethylpentane b) 2-ethyl-3-methylpentane c) 3-ethyl-2-methylpentane d) 3-methyl-2-ethylpentane 494. Ethyl acetoacetate shows, which type of isomerism? a) Chain b) Optical c) Metamerism d) Tautomerism 495. Enol content is highest in a) Acetone b) Acetophenone c) Acetic acid d) Acetyl acetone 496. The maximum number of stereoisomers possible for 3-hydroxy-2-methyl butanoic acid is : b) 2 c) 3 d) 4 a) 1 497. Which of the following will exhibit *cis* – *trans* isomerism? d) $CBr_2 = CH_2$ a) $CH_2Br - CH_2Br$ b) $CBr_3 - CH_3$ c) CHBr = CHBr498. Which of the following is most reactive towards electrophilic nitration? b) Benzene c) Benzoic acid d) Nitrobenzene a) Toluene 499. The number of structural isomers possible for an organic compound with molecular formula C_5H_{12} is : a) 5 b) 3 c) 4 d) 2



$$\begin{array}{c} CH_{3} & CH_{3$$

a) Isomerism b) Resonance c) Tautomerism d) Inductive effect 534. Nitration of benzene is a) Electrophilic substitution b) Electrophilic addition c) Nucleophilic substitution d) Nucleophilic addition 535. The following reaction is described as a) S_E 2 b) S_N 2 c) S_N 1 d) S_N 0 536. Nitrogen containing organic compound when fused with sodium forms b) Sodium cyanide a) Sodium azide c) Sodamide d) Sodium cyanate 537. Which of the following is strongest nucleophile? c) CN d) C₂H₅O: a) Br⁻ b) :OH 538. In Kjeldahl's method for the estimation of nitrogen, the formula used to a) % of N = $\frac{1.4 Vw}{N}$ b) % of N = $\frac{1.4 VN}{W}$ c) % of N = $\frac{V N V}{1.8}$ 1.4 wN d) % of N = 539. The most satisfactory method to separate sugars is to use a) Fractional crystallisation b) Chromatography c) Benedict's reagent d) Carius method 540. The IUPAC name of an unsymmetrical ether with the molecular formula $C_4 H_{10} O$ a) Ethoxy propane b) Methoxy ethane c) Ethoxy ethane d) Methoxy propane 541. S_N 1 reaction on optically active substrates mainly gives : a) Retention in configuration b) Inversion in configuration c) Racemic product d) No product 542. The structures that do not actually exist are called a) Tautomers b) Conformational isomers c) Canonical structures d) Optical isomers 543. The correct IUPAC name of, a) Isopropyl benzene b) Cumene c) Phenyl isopropane d) 2-phenyl propane 544. When SCN⁻ is added to an aqueous solution containing $Fe(NO_3)_3$, the complex ion produced is a) $[Fe(OH_2)_2(SCN)]^{2+}$ b) $[Fe(OH_2)_5(SCN)]^{2+}$ c) $[Fe(OH_2)_8(SCN)]^{2+}$ d) $[Fe(OH_2)(SCN)]^{6+}$ 545. Which of the following is the most stable carbocation? b) $\stackrel{+}{RC}$ H₂ c) $R_{2}C^{+}H$ d) $R_{3}C^{+}$ 46. Which one of the following compound is most acidic? OH a) $Cl - CH_2 - CH_2 - OH$ b) 547. The number of carbon atoms present in neopentane are : a) Four 1° carbon, one 4° carbon b) two 1° carbon, two 2° carbon c) one 1° carbon, three 4° carbon d) None of the above is correct



c) $CH_3^+ > R_2CH^+ > RCH_2^+ > R_3C^+$ d) $CH_3^+ > RCH_2^+ > R_2CH^+ > R_3C^+$ 560. How many delocalized π -electrons are there in the compounds HC-HC a) 8 b) 2 d) 6 c) 4 561. What will be the compound if two valencies of carbonyl group are satisfied by two alkyl groups? a) Aldehvde b) Ketone c) Acid d) Acidic anhydride 562. The most stable carbocation is : CH_3 a) $\begin{bmatrix} C_6H_5 & C_6H_5 \\ C_6H_5 \end{bmatrix}$ b) $\begin{bmatrix} C_6H_5 & C_6H_5 \\ C_{H_3} \end{bmatrix}$ c) $\begin{bmatrix} CH_3 \\ CH_3 \end{bmatrix}$ 563. Which of the following belongs to – *I* group? d) $-C(CH_3)_3$ c) $-CH_2CH_3$ a) $-C_6H_5$ b) $-CH_3$ 564. IUPAC name of following compound, $CH_3 - CH_2 - CH_3$ is: a) 2-cyclohexylbutane b) 2-phenylbutane 3-cyclohexylbutane d) 3-phenylbutane 565. Which is most commonly used to dry organic liquids? a) Lithium b) Sodium c) Potassium d) Rubidium 566. In Lassaigne's solution, pink/violet colouration is produced when sodium nitroprusside solution is added. It indicates the presence of b) Nitrogen a) Sulphur c) Chlorine d) None of these 567. A carbonium ion is formed when a covalent bond between two atoms in an organic compound undergoes : b) Heterolysis a) Homolysis 4 c) Cracking d) Pyrolysis 568. Racemic mixture is formed by mixing two a) Isomeric compounds b) Chiral compounds c) *meso* compounds d) Enantiomers with chiral carbon 569. In a solution, solvent can be separated from solute by one of the following process a) Decantation b) Filtration c) Distillation d) Sedimentation 570. Buta-1,3-diene and But-2-yne are : a) Position isomers b) Functional isomers d) Tautomers c) Chain isomers 571. $CH_3 - CHCl - CH_2 - CH_3$ has a chiral centre. Which one of the following represent its *R*-configuration? b) $Cl - CH_3$ c) $H = \begin{bmatrix} CH_3 \\ I \\ C \\ C \\ C_2H_5 \end{bmatrix}$ a) $H - C - CH_3$ d) $H_3C - C - Cl$ 572. Which of the following statements (s) is (are) not true? a) Carbanions and carbonium ions, usually exist in ion pairs or else solvated Acidity increases and basicity decreases in going from left to right across a row of Periodic Table b) $CH_4 < NH_3 < H_2O < HF(acidity)$ $CH_3^- > NH_2^- > OH^- > F^-$ (basicity)

c) RCOOH like RCOR reacts with H₂NOH to give an oxime

Decreasing order of ionizing power of solvents is $CF_3COOH > HCOOH > H_2O > CH_3COOH > CH_3OH > C_2H_5OH > (CH_3)_2SO > CH_3CN$ 573. The intermediate during the addition of HCl to propene in the presence of peroxide is : a) CH₃ CHCH₂Cl c) CH_3CH_2 $\dot{C}H_2$ b) $CH_3 \stackrel{+}{C}HCH_3$ d) $CH_3CH_2 \stackrel{+}{C}H_2$ 574. Which of the following represents neo -pentyl alcohol? a) $CH_3CH(CH_3)CH_2CH_2OHb)$ (CH_3)₃C – CH_2OH c) $CH_3(CH_2)_3OH$ d) $CH_3CH_2CH(OH)C_2H_5$ 575. 2-methyl-2-butene will be represented as : $CH_3 - CH - CH_2 - CH_3$ a) СH₃-СH-СH=СH₂ | СH₃ b) $CH_3 - CH_2 - C = CH_2$ c) d) CH₃—C=CH—CH₃ II CH₃ 576. The most abundant organic compound in the world is : b) Chlorophyll c) Alkaloids d) Cellulose a) CH₄ 577. The chain initiating species in free radical chlorination of methane is : c) CH₃ radical a) Cl free radical b) HCl d) Methylene radical 578. Which of the following belongs to +I group? c) -COOH b) $-0CH_3$ a) –0H d) $-CH_3$ 579. Different structures generated due to rotation about, C – C axis, of an organic molecule, are examples of a) Geometrical isomerism b) Conformational isomerism c) Optical isomerism d) Structural isomerism 580. Which of the following molecules is expected to rotate the plane of plane-polarised light? COOH сно NH_2 H_2N d) ^{H₂N} н c) a) Ph ĊH₂OH 581. Chromatography was discovered by a) Kekule b) Pauling c) Rutherford d) Tswett 582. Sodium nitroprusside when added to an alkaline solution of sulphide ions produces a colouration a) Red b) brown c) Blue d) Purple 583. $(CH_3)_4$ N is neither an electrophile, nor a nucleophile because it : a) Does not have electron pair for donation as well as cannot attract electron pair b) Neither has electron pair available for donation nor can accommodate electron since all shells of N are fully occupied c) Can act as Lewis acid and base d) None of the above 584. Isopentane can form four isomeric mono bromo derivatives. How many of them are optically active? d) None of these a) 1 b) 2 c) 3 585. Which one of the following does not show resonance? a) Carbon dioxide b) Benzene c) Nitromethane d) Propane 586. Select the organic compounds aliphatic in nature but burn with smoky flame :

a) CCl ₄ 587. Which of the foll	b) CHCl ₃ owing is an electrophilic reagent?	c) C ₆ H ₅ CH ₂ OH	d) Both (a) and (b)
a) <i>R</i> O ⁻	b) BF ₃	c) NH ₃	d) R Ö H
588. The molecular fo	ormula of diphenyl methane is C ₁₃	₃ H ₁₂ .	
How many struc	tural isomers are possible when o	one of the hydrogen is rej	placed by a chorine atom?
589 A mixture of iror	u fillings and sulphur cannot be se	cj o pharated by	
a) Heating	i mings and surphur cannot be se	h) Magnet	
c) Shaking with	CS ₂	d) Washing in a curre	ent of water
590. Isomers of propi	onic acid are		
a) HCOOC ₂ H ₅ an	d CH ₂ COOCH ₂	b) HCOOC ₂ H ₅ and C ₂	Н-СООН
c) CH ₃ COOCH ₃ a	Ind C ₃ H ₇ OH	d) C_3H_7OH and CH_3C	COCH ₃
591. Reactions involv	ing heterolytic fission are said to	proceed via :	
a) Ionic mechani	ism b) Polar mechanism	c) Both (a) and (b)	d) None of these
592. Which of the foll	owing orders is true regarding th	e acidic nature of COOH?	
a) Formic acid >	acetic acid> propanoic acid	b) Formic acid > ace	tic acid< propanoic acid
c) Formic acid <	acetic acid> propanoic acid	d) Formic acid > ace	tic acid< propanoic acid
593. Which behaves b	ooth as a nucleophile as well as an	electrophile?	
a) CH ₃ OH	b) CH ₃ NH ₂	c) CH ₃ CN	d) CH ₃ Cl
594. Alkaline hydroly	sis of an ester (A) gives alcohol a	nd salt	
	011-		
CH ₃ −Ċ−O−CH·	$- CH_2CH_3 \xrightarrow{OH} Alcohol + salt$	V	
CH ₃			
The correct state	ement about the reaction is :		
a) In alcohol con	figuration about chiral carbon ato	om is retained	
b) In alcohol con	figuration about chiral carbon ato	om is inverted	
c) Alcohol loses	optical activity		
d) All statement	are incorrect		
595. In which case the	e carbon-carbon bond length is sa	ime?	
a) 2-butene	b) Benzene	c) 1-butene	d) 1-propyne
596. Incorrect statem	ent is		
a) Aniline can be	purified by steam distillation		
b) Belistein test	is not given by fluorine	hur	
d) Lassaigen's te	st is used in the qualitative detect	tion of elements in organ	ic compounds
597 The increasing o	rder of stability of the following f	ree radicals are	le compounds
($CH_{3})_{2}$ CH < ($(CH_3)_3 C < (C_6H_5)_2 CH $ $< (C_6H_5)_3 C$		
b) $(C_6H_5)_3 \stackrel{\bullet}{C} < (C_6H_5)_3 \stackrel{\bullet}{C}$	$C_6H_5)_2$ CH < (CH ₃) ₃ C < (CH ₃) ₂ CH		
c) (C ₆ H ₅) ₂ CH <	$(C_6H_5)_3 \overset{\bullet}{C} < (CH_3)_3 \overset{\bullet}{C} < (CH_3)_2 \overset{\bullet}{C} < (CH_3)_2 \overset{\bullet}{C}H$		

d)
$$(CH_3)_2 CH < (CH_3)_3 C < (C_6H_5)_3 C < (C_6H_5)_2 CH$$

598. Which one of the following explain, why propene undergo electrophilic addition with HBr, but not with HCN?

a) Br⁻is better nucleophile than CN⁻

b) HBr being better source of proton as it is stronger acid than HCN

c) HCN attacks preferentially via lone pair of nitrogen

d) The C – Br bond being stronger is formed easily as compared to C – CN bond

599. The structural formula of 2,2,3-trimethyl hexane is :

CH₃ CH₃
a) CH₃
$$\xrightarrow{-C}_{-C}_{-CH}_{-CH_2}_{-CH_2}_{-CH_3}$$

b) CH₃ $\xrightarrow{-C}_{-C}_{-CH_2}_{-CH_2}_{-CH_2}_{-CH_3}$
c) CH₃ $\xrightarrow{-C}_{-C}_{-CH_3}$

d)
$$CH_3 - CH - CH_2 - CH_2 - CH_3 - CH_3 - CH_3 - CH_2 - CH_2 - CH_3 -$$

600. The IUPAC name of the compound,

a) 2-iodo-3-chloro-4-pentanoic acid

b) 4-oxo-3-chloro-2-iodo pentanoic acid

- c) 4-carboxy-4,3-chloro-2-butanone
- d) 3-chloro-2-iodo-4-oxo-pentanoic acid

601. Select the correct statement about the detection of sulphur in organic compounds

- a) Sulphur present in organic compound on fusion with sodium is converts to $Na_2S_2O_3$
- b) FeCl₃ gives purple colour when added to the sodium fusion extract
- c) Sodium nitroprusside is used to detect the presence of sulphur
- d) All of the above

602. The name of the compound



- c) Show geometrical isomerism
- d) Are unstable molecules
- 615. Who is called the 'Father of Chemistry'?a) Faradayb) Priestley
- a) Faraday
 b) Priestley
 c) Rutherford
 d) Lavoisier
 616. With a change in hybridization of the carbon bearing the charge, the stability of a carbanion increase in the order
 - a) $sp < sp^3 < sp^2$ b) $sp < sp^2 < sp^3$
- 617. The correct IUPAC name of the acid



- a) Z-3-ethyl-4-methyl hex-3-en-1-oic acid
- c) Z-3, 4-diethylpent-3-en-1-oic acid
- 618. IUPAC name of,

$$CH_{3} - C = C - CH - CH = CH_{2} is:$$

$$CH_{3} - C = CH - CH = CH_{2} is:$$

$$CH_{2} - CH_{2} - CH_{2}$$

a) 2-ethyl-3-methyl-hexa-l-en-4-yne

b) 5-ethyl-4-methyl-hexa-2-yn-5-ene

- c) 3-methylene-4-methylhepta-5-yne
- d) 5-methylene-5-ethyl-4-methylhepta-2-yne
- 619. The total number of alkenes possible by dehydromination of 3-bromo-3-cyclopentylhexane using alcoholic KOH is

a) 1	b) 3	c) 5	d) 7
620. The Cl—C—Cl bon	d angle in dichloro methane	e will be :	
a) > 109°28′	b) < 109°28′	c) 109°28′	d) 120°
621. In the following rea	action sequence, the chain i	nitiation steps is :	

a)
$$Cl \rightarrow Cl \rightarrow Cl + Cl$$

b)
$$CH_4 + Cl \longrightarrow CH_3 + HCl$$

c)
$$\dot{C}H_3 + Cl_2 \longrightarrow CH_3Cl + \dot{C}l$$

d)
$$\overset{\bullet}{CH_3+} \overset{\bullet}{Cl} \longrightarrow CH_3Cl$$

622. Amongst the following which of the above are true for $S_N 2$ reaction?

(i) The rate of reaction is independent of the concentration of the nucleophile.

(ii) The nucleophile attacks the carbon atom on the side of the molecule opposite to the group being displaced.

(iii) the reaction proceeds with simultaneous bond formation and bond rupture.

a) (i), (ii) b) (i), (iii) c) (i), (iii) d) (ii), (iii) 623. Predict the nature of principal product in the reaction, BrCH₂CH₂CH₂CH₂CH₂Br + KOH (alc.) \rightarrow Product : a) 1,3-butadiene b) Cyclobutane c) BrCH₂CH₂CH = CH₂ d) None of these 624. The most stable carbanion is

a) CH_3^{Θ} b) RCH_2^{Θ} c) R_3C^{Θ} d) CH_2CHO

625. Crown ethers are named as X-crown-Y. In the following crown ether, X and Y are respectively

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b) *Z*-3-ethyl-4-methyl hexanoic acidd) *E*-3-ethyl-4-methylhex-4-en-1-oic acid

b) $sp < sp^2 < sp^3$ c) $sp^2 < sp < sp^3$ d) $sp^3 < sp^2 < sp$





benzyl allyl methyl vinyl Ι Π III IV a) IV>III>II>I b) I>II>II>IV c) II>IV>III>I d) III>II>IV 650. The prefix name of —SH group in IUPAC system is : b) Thiol c) Sulphide d) None of these a) Mercapto 651. The correct name for CH₃COCH₂OH is : a) 2-keto propanol b) 1-hydroxy propan-2-one c) Propan-2-one-1-ol d) 3-hydroxy propan-2-one 652. The maximum number of alkyl groups in $\rm C_8H_{18}$ is : d) 2 a) 6 b) 5 c) 4 653. The chlorination of methane to give CCl₄ is an example of d) Chain reaction a) Addition b) Elimination c) Substitution 654. The number of isomers for the aromatic compound of the formula $C_7 H_8 O$ is : a) 2 b) 3 c) 4 d) 5 655. The IUPAC name of the compound having the formula $CH \equiv C - CH = CH_2$ is : a) 1-butene-3-yne b) 3-butene-1-yne c) 1-butyn-3-ene d) But-1-yne-3-ene $656. H_3C - C = CH - CH - CH_3$ Cl CH_3 b) 4-chloro-2-methyl-3-pentene a) 2-chloro-4-methyl-2-pentene d) 2-chloro-4, 4-dimethyl-2-butene c) 4-methyl-2-chloro-2-pentene 657. Give the IUPAC name of the compound a) 1,1,3-trimethylcyclohex-2-ene b) 1,3,3-trimethylcyclohex-1-ene c) 1,1,5-trimethylcyclohex-5-ene d) 2,6,6-trimethylcyclohex-1-ene 658. C - H bond energy is about101 kcal/mol for methane, ethane and other alkanes but is only77 kcal/mol for C – H bond of CH₃ in toluene. This is because a) Of inductive effect due to $-CH_3$ in toluene b) Of the presence of benzene ring in toluene c) Of resonance among the structures of benzyl radical in toluene d) Aromaticity of toluene 659. The reaction $(CH_3)_3C - Br \xrightarrow{H_2O} (CH_3)_3COH$ is a) Elimination b) Substitution c) Free radical d) Addition 660. The number of structural and configurational isomers of a bromo compound, C₅H₉Br, formed by the addition of HBr to 2-pentyne respectively are : a) 1 and 2 b) 2 and 4 c) 4 and 2 d) 2 and 1 661. Fractional distillation is a process by which the separation of different fractions from mixture of solution is carried by making use of the following property of the fractions d) Solubility a) Freezing point b) Boiling point c) Melting point 662. The maximum number of carbon atoms arranged linearly in the molecule, $CH_3 - C \equiv C - CH = CH_2$ is a) 5 b) 4 c) 2 d) 3 663. α -D-glucose and β -D-glucose have a specific rotation of +112° and +19° respectively. In aqueous solution





c)
$$HC \equiv C - C - H$$

d) CI
 $HC \equiv C - C - H$

684. Which of the following statements is incorrect?

- a) The rate of reaction increases with increase in water concentration in the hydrolysis of tertiary butyl bromide in methanol and water
- The relative nucleophilicity in protic solvent is

b) $CN^- > I^- > \overline{O}H > Br^- > CI^- > F^- > H_2O$

- c) In $S_N 2$ reactions, the order of reactivity of alkyl halides is in the order methyl>primary>secondary> tertiary
- d) $S_N 2$ reaction involves carbonium ions

685. Arrange in order of increasing acidic strength.



b) Z < X > Y

d) Z > X > Y

d) HBr

686. For the purification, isolation and separation of organic compounds, the latest technique followed is b) Steam distillation a) Chromatography

c) X>

c) HF

c) Fractional crystallisation d) Sublimation

b) H_2S

- 687. Arrange *p*-toluidine (I) N,N-dimethyl-*p* toluidine (II) *p* nitroaniline (III) and aniline (IV) in order of decreasing basicity
 - b) I>II>II>IIV a) I>IV>III>II c) II>I>IV>III d) III>I>II>IV
- 688. The reagent showing addition on alkene against the Markownikoff's rule of :
- a) Br_2 689. Carbocation can undergo :

a) Loss of a proton

- b) Addition to multiple bond
- c) Combination with anions
- d) All of the above
- 690. Lactic acid is :
 - a) Propionic acid
 - b) β-hydroxypropanoic acid
 - c) α-hydroxypropanoic acid
 - d) None of the above

691. Of the following compounds which will have a zero dipole moment?

- a) 1,1-dichloroethylene
 - b) trans-1,2-dichloroethylene
 - c) cis-1,2-dichloroethylene
 - d) None of the above
- 692. The IUPAC name of













722. Lactic acid in which a methyl group, a hydroxyl grou	up, a carboxylic acid group	and a hydrogen atom are
attached to a central carbon atom shows optical iso	merism due to the molecu	lar geometry at the :
a) Carbon atom of the methyl group		
b) Carbon atom of the carboxylic acid group		
c) Central carbon atom		
d) Oxygen of the hydroxyl group		
723. Which of the following process is suitable for the pu	irification of aniline?	
a) Simple distillation	b) Fractional distillation	
c) Fractional crystallisation	d) Steam distillation	
724. Maleic and fumaric acids are :		
a) Tautomers b) Geometrical isomers	c) Chain isomers	d) Functional isomers
725. $CH_3Br + Nu \rightarrow CH_3 - Nu + Br$ The decreasing	order of the rate of the ab	ove reaction with
nucleophiles (Nu) A to D is:		
$[NU = (A)PNO , (B)ACO , (C)HO , (D)CH_3O]$		
a) $D > C > A > B$ b) $D > C > B > A$ 726 Which are in least reactive in a nuclear hile substitu	CJ A > B > C > D	a) B > D > C > A
2 26. Which one is least reactive in a nucleophile substitution of the substitution 2	$r_{\rm a}$ $r_{\rm cu}$ $ r_{\rm cu}$	
a) $\Box \Pi_3 \Box \Pi_2 \Box$ D) $\Box \Pi_2 = \Box \Pi \Box \Pi_2 \Box$	$C_{1} CH_{2} = CHCI$	$(UH_3)_3UU$
dibromoethene begauge :	to yield $DICH_2CH_2OCH_3$ if	ir addition to 1,2-
a) The intermediate carbocation may react with Pr		
b) The methyl alcohol solvolates the bromine	01 CH30H	
c) The reaction follows Markownikoff's rule		
d) This is a free radical mechanism		
728 Number of tertiary carbon atoms in tertiary butyl a	lcohol is :	
a) 1 b) 2	c) Zero	d) 4
729. Which step is chain propagation step in the followit	ng mechanism?	
	0	
(i) $Cl_2 \longrightarrow Cl^2 + Cl^2$		
(ii) $Cl^{\bullet} + CH_4 \longrightarrow CH_3 + HCl$		
$(iii) Cl^{\bullet} + Cl^{\bullet} \longrightarrow Cl_2$		
$(iv) \stackrel{\bullet}{C}H_2 + Cl \stackrel{\bullet}{\longrightarrow} CH_2 Cl$		
		-) (:)
$\frac{1}{2} \frac{1}{2} \frac{1}$	cj (III)	u) (IV)
730. The IOPAC name of the compound $CH_3 - N \equiv C$ is:		
a) Ethane nitrile b) Methane isonitrile	c) Ethane isonitrile	d) None of these
/31.10 PAC name of CIL CL $(CPr) = CIL CL is$		
$CH_3CH_2C(BF) = CH - CHS$	h) 1 chlara 2 hrama hu	tono 1
a) 2-bromo-1-chioro butene-1 c) 2-chloro 2-bromo butene 2	d) None of the above	tene-1
C) 5-CHIOTO-Z-DI OHIO DUTEHE-Z 722 Which of the following undergoes nucleonbilic subs	uj None of the above	achanicm?
2) Repart chlorida b) Icopropul chlorida	$_{\rm c}$ Chlorobonzono	d) Ethyl chlorido
722 The sigma hand energy of C—H hand in C. H. is:	cj chiorobenzene	d) Ediyi chioride
2) 90 kcal b) 140 kcal	c) 200 kcal	d) 60 kcal
734 The general formula (H. O. could be for open cha	cj 200 Kcal	u) oo kcar
a) Diketones b) Carbovylic acids	c) Diols	d) Dialdebydes
735 The correct sequence of steps involved in the mech	anism of Cannizzaro's read	tion is
a) Nucleophilic attack transfer of H ⁻ and transfer of	fH ⁺	
b) Transfer of H^- transfer of H^+ and nucleophilic a	ottack	
c) Transfer if H^+ , nucleonhilic attack and transfer of H^-		
d) Electrophilic attack by OH^- transfer of H^+ and the	ransfer of H ⁻	
736. Examine the following statements regarding $S_{\rm M}$ 2 re	eaction	

(1) The rate of reaction is independents of concentration of nucleophile

(2) The nucleophile attacks the carbon atom on the side of molecule opposite to the group being displaced(3) The reaction proceeds with simultaneous bond formation and rupture

Which of the above written statements is correct? b) 1, 3 d) 2, 3 a) 1, 2 c) 1, 2, 3 737. Propanol and propanone are a) Functional isomers b) Enantiomers c) Chain isomers d) Structural isomers 738. Diastereomers can be separated by : c) Electrophoresis a) Fractional distillation b) Simple distillation d) All of these 739. Angle strain in cyclopropane is d) -5°16' a) 24°44' b) 9°44' c) 44' 740. The function of AlCl₃ in Friedel-Craft's reaction is a) To absorb HCl b) To absorb water c) To produce nucleophiled) To produce electrophile 741. In Kjeldahl's method of estimation of nitrogen, CuSO₄ acts as d) Hydrolysis agent a) Oxidising agent b) Reducing agent c) Catalytic agent 742. A mixture of acetone and methanol can be separated by a) Steam distillation b) Vaccum distillation c) Fractional distillation d) None of these 743. The IUPAC name of, $\begin{array}{c} CH_3 - CH - CH_2 - CH - CHO \quad is: \\ I & I \\ OH & CH_3 \end{array}$ a) 4-hydroxy-1-methylpentanal b) 4-hydroxy-2-methylpentanal c) 3-hydroxy-2-methylpentanal d) 3-hydroxy-3-methylpentanal 744. The oxygen atom in phenol a) Exhibits only inductive effect b) Exhibits only resonance effect c) Has more dominating resonance effect than inductive effect d) Has more dominating inductive effect than the resonance effect 745. 2-methylpent-3-enoic acid shows : a) Optical isomerism b) Geometrical isomerism c) Both (a) and (b) d) None of these 746. In the reaction, $ROH - R'COOH \rightarrow R' - C - OR + H_2O$ water is formed by the combination of : a) Hydroxyl of acid with alcoholic hydroxyl hydrogen b) Hydroxyl of alcohol with carboxylic hydrogen c) Both the above changes d) None of the above 747. Pyridine is : a) An aromatic compound and a primary base b) A heterocyclic amino compound and a tertiary base c) An aromatic amino compound and forms salts

d) A cyano derivative of benzene and secondary base



d) _H / ^{Cl}		
H ₃ C ⁷ Cl		
759. When thiourea is heated with metallic sodium, the c	compound which can't be fo	ormed is
a) NaCNS b) NaCN	c) Na ₂ SO ₄	d) Na ₂ S
760. An unknown compound <i>A</i> has a molecular formula	C_4H_6 . When A is treated wi	th excess of Br_2 a new
substance B with formula $C_4H_6Br_4$ is formed. A form	ns a white ppt. with ammo	niacal silver nitrate solution.
A may be :		
a) But-1-yne b) But-2-yne	c) But-1-ene	d) But-2-ene
/61. The facemisation of optically active compounds is a	riven by :	d) Element of a montation
a) Entropy D) Enthalpy	C II Clara classified and t	a) Element of Symmetry
762. A cyclic steleoisoniel having the molecular formula	$C_4 \Pi_7 CI are classified and t$	abulated. Find out the
Competical Optical		
a) 6 2	h) 4 2	
$r = \frac{1}{2}$	d) 4 0	\circ
763 The correct name for the following hydrocarbon is		
a) Tricycle [4.1.0]heptane	b) Bicyclo [5.2.1] heptan	2
c) Bicyclo [4.1.0] heptane	d) Bicyclo [4.1.0] hexane	
764. Which of the following is the most stable radical?		
a) CH_{2}^{\bullet} b) RCH_{2}^{\bullet}	c) R ₂ CH [•]	d) $R_2 C^{\bullet}$
	0) 12000	uj rize
765. The number of 4° carbon atoms in $2,2,4,4$ -tetrametr	iyipentane :	
a) 1 DJ 2	° CJ 3	a) 4
r_{0} Delegalization of σ electrons	b) Displacement of a ala	strong
c) Delocalisation of π -electrons	d) Displacement of π -elec	ctrons
767 Compounds whose molecules are superimposable	e on their mirror images	even though they contain
asymmetric carbon atoms or chiral centres are know	wn as ·	even though they contain
a) Enantiomers b) Racemers	c) Mesomers	d) Conformers
768. Percentage of hydrogen is maximum in .	0) 11000111010	
a) C_2H_4 b) CH_4	c) C ₂ H ₂	d) $C_6 H_6$
769. Which of the following has most acidic hydrogen?	, , , , , , , , , , , , , , , , , , , ,	5 6 6
a) 3-hexanone b) 2, 4-hexanedione	c) 2, 4-hexanedione	d) 2, 3-hexanedione
770. IIIPAC name of CH ·N·CH.		
C ₂ H ₅		
a) $N_{\rm N}$ - dimethylethanamine		
b) <i>N</i> -methyl <i>N</i> -ethylmethanamine		
c) Dimethyl-ethylamine		
d) None of the above		
771. Ease of abstraction of hydrogen is greater when atta	ached to :	
a) 1° carbon b) 2° carbon	c) 3° carbon	d) <i>neo-</i> carbon
772. <i>neo</i> -Heptyl alcohol is correctly represented as :	-	-



	a) Aq. NaHCO ₃	b) H ₂ O	c) Aq. HCl	d) Aq. NaOH
781. The name formic acid was given for HCOOH because it was prepared from :				
	a) Acetum	b) Ant	c) Wood	d) Oxalis plant
782. 2, 3-dimethyl hexane contains tertiary secondary andprimary carbon atmos, respectively				
	a) 2, 2, 1	b) 2, 4, 3	c) 4, 3, 2	d) 3, 2, 4

d) None of these

d) A nucleophile

783. Which one of the following is the correct formula for dichlorodiphenyltrichloro ethane?



b) RS^{\ominus} is less basic but more nucleophilic than RO^{\ominus}



a) 6 b) 8	c) 4	d) 2
804. What is the number of possible optical isomers i	n glucose?	
a) 3 b) 4	c) 12	d) 16
805. In which reaction addition takes place according	to Markownikoff's rule?	
a) $CH_3CH = CHCH_3 + Br \rightarrow$	b) $CH_2 = CH_2 + HBr \rightarrow$	
c) $CH_3CH = CH_2 + HBr \rightarrow$	d) $CH_3CH = CH_2 + Br_2$	\rightarrow
806. Presence of halogen in organic compounds can b	e detected using	
a) Leibig's test b) Duma's test	c) Kjeldahl test	d) Beilstein's test
807. The bond energy for catenation next to carbon is	5 :	
a) N b) S	c) Si	d) P
808. The hydrolysis of alkyl halides by aqueous NaOH	I is best termed as :	
a) Electrophilic substitution reaction		
b) Electrophilic addition reaction		
c) Nucleophilic addition reaction		
d) Nucleophilic substitution reaction		
809. Which of the following compounds exhibit stere	oisomerism?	
a) 3-methyl butyne –1	b) 2-methyl butene –1	
c) 2-methyl butanoic acid	d) 3-methyl butanoic ad	cid
810. The + I.E.(inductive effect) is shown by :		
a) CH ₃ b) —OH	c) F	d) — C_6H_5
811. In paper chromatography		
a) Mobile phase is liquid and stability phase is so	olid	
b) Mobile phase is solid and stationary phase is l	iquid	
c) Both phases are liquids	C.XY	
d) Both phases are solids		
812. Which one of the following is not found in alkene	es?	
a) Chain isomerism		
b) Geometrical isomerism		
c) Metamerism		
d) Position isomerism		
813. Select the correct statement :		
a) The prefixes are written before the name of c	ompound	
b) The suffixes are written after the name of com	npound	
c) The IUPAC name of a compound is always wri	itten as one word	
d) All of the above		
814. A compound contains 2 dissimilar asymmetric ca	arbon atoms. The number of	optically active isomers is :
a) 2 b) 3	c) 4	d) 5
815. The inductive effect		
a) Implies the atom's ability to cause bond polar	ization	
b) Increases with increase of distance		
c) Implies the transfer of lone pair of electrons f	rom more electronegative at	om to the lesser
electronegative atom in a molecule		
d) Implies the transfer of lone of electrons from	lesser electronegative atom t	to the more electronegative
atom in a molecule		
816. IUPAC name of the compound, ClCH ₂ CH ₂ COOH i	S :	
a) 3-chloropropanoic acid		
b) 2-chloropropanoic acid		
c) 2-chloroethanoic acid		
d) Chlorosuccinic acid		
817. Which one is a nucleophilic substitution reaction	n among the following?	
a) $CH_3CHO + HCN \rightarrow CH_3CH(OH)CN$	-	

b)
$$CH_{3}-CH=CH_{1}+H_{1}O_{1}H_{2}^{-1}CH_{2}-CH_{3}^{-1}O_{1}^{-1}CH_{3}^{-1}O_{1}$$


a) Partially eclipsed	b) Eclipsed	c) gauche	d) Staggered
	~) = F =	-) 8	

839. Amongst the following compounds, the optically active alkane having lowest molecular mass is

 CH_3 a) $CH_3 - CH_2 - CH_2 - CH_3$ b) $\dot{CH}_3 - CH_2 - \dot{CH} - CH_3$ c) CH₃—c d) $CH_3 - CH_2 - C \equiv CH$ 840. How many chiral isomers can be drawn from 2-bromo, 3-chloro butane? d) 5 a) 2 b) 3 c) 4 841. Glycerol can be separated from spent-lye in soap industry by a) Steam distillation b) Fractional distillation c) Distillation under reduced pressure d) Ordinary distillation 842. The IUPAC name of $H_3C - CH - C_3H_7$ I OC₃H₇ a) 4-propoxy pentane b) Pentyl-propyl ether c) 2-propoxy pentane d) 2-pentoxy propane 843. Correct gradation of basic charactor b) $CH_3NH_2 > NH_3 > NF_3$ a) $NH_3CH_3NH_2 > NF_3$ c) $NF_3 > CH_3NH_2 > NH_3$ d) $CH_3NH_2 > NF_3 > NH_3$ 844. An organic compound contains 49.3% carbon, 6.84% hydrogen and its vapour density is 73. Molecular formula of compound is a) $C_6 H_9 O_3$ b) C₄H₁₀O₂ c) $C_3H_5O_2$ d) $C_3 H_{10} O_2$ 845. Vital force theory of the origin of organic compounds was discarded by : a) Kolbe's synthesis b) Haber's synthesis c) Wöhler's synthesis d) Berthelot's synthesis 846. In the presence of peroxide, hydrogen chloride and hydrogen iodide do not give anti-Markownikoff addition to alkenes because : a) Both are highly ionic b) One is oxidizing and the other is reducing c) One of the steps are exothermic in both the cases d) All the steps are exothermic in both the cases 847. Which of the following does not have a resonance structure? a) Benzene b) Benzaldehyde c) Acetaldehyde d) Benzylamine 848. Which of the following is the correct order of stability of different conformations of butane? a) Staggered > Gauche > Partially eclipsed > Fully eclipsed b) Gauche > Staggered > Partially eclipsed > Fully eclipsed c) Staggered > Fully eclipsed > Partially eclipsed > Gauche d) None of the above 849. Glucose and fructose are : c) Functional isomers d) Optical isomers a) Chain isomers b) Position isomers 850. The enol form of acetone after treatment with D_2O gives : b) O CH₃-C=CD₂

 $(I)CH_3CH_2CH_2CH_3 \qquad (II)CH_3CH = CH - CH_3 \\ (III)CH_2 = CH - CH = CH_2 \quad (IV)H - C \equiv C - H$





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886. During the fusion of an organic compound with sodium metal, nitrogen of the compound is converted into
a) NaNO₂
b) NaNH₂
c) NaCN
d) NaNC
887. The structure representing a heterocyclic compound is :



895. Lactic acid, CH₃CH(OH)COOH molecule shows :

a) Geometrical isomerism		
b) Metamerism		
c) Optical isomerism		
d) Tautomerism		
896. <i>n</i> -pentane and neopentane are :		
a) Functional isomers b) Geometrical isomers	c) Chain isomers	d) Position isomers
897. The IUPAC name of acryldehyde is		
a) Prop-2-en-1-al b) Propenylaldehyde	c) But-2-en-1-al	d) Propenal
898. Due to presence of an unpaired electron, free radica	ls are	· · ·
a) Cations b) Anions	c) Chemically inactive	d) Chemically reactive
899. 2-methylpent-3-ene is a chiral because it has :		
a) A centre of symmetry		
b) A plane of symmetry		
c) Symmetry at C ₂ carbon		
d) Both centre and a plane of symmetry		
900. Which of the following molecules contain asymmetr	ric carbon atom?	
a) CH ₃ CHClCOOH b) CH ₃ CH ₂ COOH	c) ClCH ₃ . CH ₂ COOH	d) Cl ₂ CHCOOH
901. Cyclobutane and butene-1 are :		
a) Chain isomers b) Position isomers	c) Ring-chain isomers	d) Metamers
902. Which of the following is not true for carbanions?		2
a) The carbon carrying the charge has eight valence	electrons	
b) They are formed by heterolytic fission		
c) They are paramagnetic		
d) The carbon carrying the charge is sp^3 hydridised		
903. Which of the following structures permits <i>cis-trans</i>	isomerism?	
a) $X_2C = CY_2$ b) $XYC = CZ_2$	c) $X_2C = CXY$	d) $XYC = CXY$
904. Which one of the following compound will show opt	tical iosmerism?	- ,
a) $(CH_2)_2 - CH - CH_2 - CH_2$	b) CH ₂ – CHOH – CH ₂	
c) $CH_3 - CHCl - CH_2 - CH_3$	d) $CH_3 - CCl_2 - CH_2 - C$	H₃
905. The Kolbe's electrolysis proceeds <i>via</i>		3
a) Nucleophilic substitution mechanism	b) Electrophilic addition	mechanism
c) Free radical mechanism	d) Electrophilic substitut	ion reaction
906. Which of the following statements is not correct?	.)	
a) Primary carbocation are more stable than second	larv ones	
b) Secondary free radicals are more stable than prin	nary free radicals	
c) Tertiary free radicals are more stable than second	dary ones	
d) Tertiary carbonium ions are more stable than pri	mary ones	
907. Adsorbent is made of in TLC.		
a) Silica gel b) Alumina	c) Both (a) and (b)	d) None of these
908 Amongst the following the total number of compou	nds soluble in aqueous Na()H is
, the cotal number of compou	nae soluble in aqueous nat	
CUT.		
\checkmark		





921. Which of the following Fischer's projection formula is identical to D-glyceraldehyde?

Predominant product is b) $R_3N - CH_2 - CH_2 - Br$ Ð d) No reaction c) _{CH2}==CH---NR₃ Br 932. *n*- pentane and 2-methyl butane are a pair of a) Enantiomers b) Stereoisomers c) Diastereomers d) Constitutional isomers 933. IUPAC name of compound, is : a) 3-ethyl-4,4-dimethyl heptane b) 1,1-diethyl-2,2-dimethyl pentane c) 4,4-dimethyl-5,5-diethyl pentane d) 5,5-diethyl-4, 4-dimethyl pentane 934. IUPAC name of, $(C_2H_5)_2$ CHCH₂OH is : a) 2-ethylbutanol-1 b) 2-methylpentanol-1 c) 2-ethylpentanol-1 d) 3-ethylbutanol-1 935. Amongst the following the compound that can most readily get sulphonated is b) Toluene c) Nitrobenzene a) Benzene d) Chlorobenzene 936. In E2 elimination, some compounds follow Hofmann's rule which means: a) The double bond goes to the most substituted carbon b) The compound is resistant to elimination c) No double bond is formed d) The double bond goes mainly towards the least substituted carbon 937. How many asymmetric carbon atoms are present in (i) 1, 2-dimethylcyclohexane (ii) 3-methylcyclopentane and (iii) 3-methylcyclohexene? b) One, one, one a) Two, one , one c) Two, none, two d) Two, none One 938. Which of the following is a chiral compound? a) Hexane b) *n*-butane c) Methane d) 2,3,4,trimethyl hexane 939. How many structures can compound with molecular formula C₇H₉N have? a) 5 b) 4 c) 3 d) 2 940. The maximum number of possible optical isomers in 1-bromo-2-methyl cyclobutane is b) 2 a) 4 c) 8 d) 16 941. Which of the following types of reaction occurs when a substituent has got a double bond with evently distributed π electron cloud? a) Electrophilic addition b) Nucleophilic addition c) Any of the (a) and (b) d) None of the above 942. Vinyl alcohol and acetaldehyde are : a) Geometrical isomers b) Keto-enol tautomers c) Chain isomers d) None of these 943. 0.25 g of an organic compound on Kjeldahl's analysis gave enough ammonia to just neutralise 10cm³ of $0.5M H_2SO_4$. The percentage of nitrogen in the compound is



+ HBr _ Is an example of a) Nucleophilic substitution b) Electrophilic addition c) Elimination reaction d) Nucleophilic addition 957. Acetone and propen-2-ol are : a) Positional isomers c) Geometrical isomers d) Chain isomers b) Leto-enol tautomers 958. The number of stereoisomers obtained by bromination of trans - 2 -butene is? b) 2 d) 4 a) 1 c) 3 959. The compound which forms one monochloro product when treated with chlorine is : d) None of these a) *n*-pentane b) Isopentane c) neo-pentane 960. Reactivity towards nucleophilic addition reaction of (I)HCHO (II)CH₃CHO (III)CH₃COCH₃is a) II>III>I b) III>II>I c) I>II>III d) I>II<III 961. Maleic acid and fumaric acid are a) Position isomers b) Geometric isomers c) Enantimoers d) Functional isomers 962. The ease of nitration of the following three hydrocarbons follows the order CH_3 NO_2 I Π Ш c) III>II>I a) II=III≈I b) II>III>I d) I=III>II 963. Which represents the condensed formula for pentanes? a) $CH_3(CH_2)_3CH_3$ b) $(CH_3)_3CCH_3$ c) $(CH_3)_2CHCH_2CH_3$ d) All of these 964. Which of the substance is purified by sublimation? a) Benzoic acid b) Camphor c) Naphthalene d) All of these 965. The halogen compound which most readily undergoes nucleophilic substitutions is a) $CH_2 = CHCl$ b) $CH_3CH = CHCl$ c) $CH_2 = CHC(CI) = CH_2$ d) $CH_2 = CHCH_2Cl$ 966. Which of the following order is correct regarding the acidity of carboxylic acids? a) $Cl_3CCOOH > Cl_2CHCOOH > ClCH_2COOH$ b) $Cl_3CCOOH > Cl_2CHCOOH < ClCH_2COOH$ c) $Cl_3CCOOH < Cl_2CHCOOH > ClCH_2COOH$ d) $Cl_3CCOOH < Cl_2CHCOOH < ClCH_2COOH$ 967. An S_N2 reaction at an asymmetric carbon of a compound always gives a) A mixture of diastereomers b) A single stereoisomer c) An enantiomer of the substrate d) A product with opposite optical rotation 968. The IUPAC name of the compound, $CH_3CH = CHC \equiv CH$ is : a) Pent-4-yn-2-ene b) Pent-3-en-1-yne c) Pent-2-en-4-yne d) Pent-1-yn-3-ene 969. Reaction of methyl bromide with aqueous sodium hydroxide involves a) Racemisation b) S_N1 mechanism c) Retention of configuration d) $S_N 2$ mechanism 970. An organic compound X(mol. formula $C_6H_5O_2N$) has six carbons in a ring system, three double bonds and also a nitro group as substituent. X is : a) Homocyclic but not aromatic b) Aromatic but not homocyclic c) Homocyclic and aromatic

d) heterocyclic

uj neterocyclic		
971. The compounds CH_3NH_2 and CH_3CH_2 . NH_2 are :		
a) Isomers b) Isobars	c) Homologous	d) Allotropes
972. The following compound will undergo electrophili	c substitution more readily	than benzene
a) Nitrobenzene b) Benzoic acid	c) Benzaldehyde	d) Phenol
973. Which of the following elements can't be detected	by direct tests?	
a) N b) O	c) S	d) Br
974. IUPAC name of ,		
H_3C — CH — CH_2 — CH — CH_2CI is:		· · ·
c) 2 chloromethyl 4 methyl hevenal		
a) 2-chloromethyl-4-methyl-nexanal		
b) 1-chloro 4-ethyl 2 heyerol		
c) 1-chloro-4-methyl-2-nexanal		
075 Desition isomerican is shown have		
975. Position isomerism is snown by :		N I
a) 0 -nitroinenoi and p -nitrophenoi	C	× ×
b) Dimetnyi etner and etnahol		
c) Pentan-2-one and pentan-3-one		
a) Acetaidenyde and acetone		
976. Formulae of phenyl carbinol and chloral are respec	ctively:	
a) L_6H_5 . LH_2LH_2 OH and LHL_2LHO		
b) $L_6H_5CH_2OH$ and CLl_3CHO		
c) L_6H_5 OH and LH_2 CI. CHO	$\mathcal{L}^{\mathbf{Y}}$	
d) C_6H_5CHO and $CHCl_2CHO$	V.	
977. How many primary carbon atoms are there in the	compound,	
CH_3 CH_3		
CH_3 — CH_2 — CH_2 — CH_3 ?		
a) 6 b) 2	c) 4	d) 3
978. IUPAC name of,		
OHC—CH $=$ CH—CH $=$ CH $=$ CH $_2$ is:		
CH ₂ CH ₂ CH ₂ CH ₂ CH ₃		
a) 4-butyl-2,5-hexadien-1-al		
b) 5-vinvloct-3-en-1-al		
c) 5-vinyloct-5-en-8-al		
d) 3-butyl-1,4-hexadien-6-al		
979. The molecular formula of a saturated compound is	$C_2H_4Br_2$.	
This formula permits the existence of :		
a) Functional isomers b) Optical isomers	c) Positional isomers	d) <i>cis – trans</i> isomers
980. Which of the following solvents are aprotic?	,	,
$(A)NH_3$ $(B)SO_2$		
$(C)CH_{2}CN$ $(D)CH_{2}CO_{2}H$		
a) A,B,C b) A,C,D	c) B,C	d) A.C
981. The reaction of sodium ethoxide with iodoethane t	o from diethyl is termed as	, .
a) Electrophilic substitution	b) Nuclephilic substituti	ion
c) Electrophilic addition	d) Radical substitution	
982. The sodium extract of an organic compound on aci	dification with acetic acid a	and addition of lead acetate
solution gives a black precipitate. The organic com	pound contains	
0		



is influenced by the hyper conjugation effect of group *R*. If *R* sequentially is

I.
$$CH_3 - H_2 - H_3C - CH_2 - H_3C - CH_3$$

III. $CH_3 - CH_3$
III. $CH_3 - CH_3$
 $H_3C - C - H_3$
IV. CH_3

the increasing order of speed of the above reaction is

- a) IV, III, II, I
- b) I, II, III, IV
- c) I, IV, III, II
- d) III, II, I, IV

Will be in order

995. The compound, whose stereo-chemical formula is written below, exhibits *x* geometrical isomers and y optical isomers





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a) 4 b) 3 c) 2 d) 1 101 How many π -electrons are there in following? HC — CH 1. CH HC d) 8 a) 2 c) 6 101 2. **IUPAC** name of is a) Dimethyl amine b) 2-amino propane c) Isopropylamine d) 2-propanamine 101 An organic compound having molecular mass 60 is found to contain C=20%, H=6.67% and N=46.67% while rest is oxygen. On heating it gives NH₃ along with a solid residue. The solid residue gives violet 3. colour with alkaline copper sulphate solution. The compound is a) CH₃CH₂CONH₂ b) $(NH_2)_2CO$ d) CH₃NCO c) CH_3CONH_2 101 How many chiral carbon atoms are present in 2, 3, 4- trichloropentane? 4. d) 3 a) 4 b) 1 c) 2 101 Which one of the following compounds is most polar? 5. c) CH_2Cl_2 a) CH_2I_2 d) CH₂Br₂ b) CH_2F_2 101 Geometrical isomerism is not shown by 6. a) 1, 1-dichloro-1-pentene b) 1,2-dichloro-1-pentene c) 1, 3-dichloro-2-pentene d) 1, 4-dichloro-2-pentene 101 The change in optical rotation with time of freshly prepared solution of sugar is known as: 7. a) Specific rotation) b) Inversion c) Rotatory motion d) Mutarotation 101 Which of the following does not show stereo isomerism? 8. CH₃ a) CH_3 b) Η CH₃ COOH c) CH

d) COOH Н-- OH COOH 101 One of the following compounds exhibit geometrical iosmerism 9. a) CH₃CH₂CH₂CH₃ b) $CH_3 - HC(CH_3) - H(C)CH_3 - CH_3$ c) $CH_3 - HC(CH_3) - CH_3$ d) $CH_3CH = CH - CH_3$ 102 Which one of the following shows functional isomerism? 0. b) C_3H_6 c) C_2H_5OH d) CH_2Cl_2 a) C_2H_4 R 102 1. The chirality of the compound H₃C a) R b) S c) Z 102 In the compound 2. **Ring II** Ring I electrophilic substitution occurs at a) ortho/para position at ring I b) *meta* position at ring I d) meta position at ring II c) ortho/para position at ring II 102 In the reaction of phenol with chloroform and aqueous solution of NaOH at 70°C, the electrophile attacking 3. the ring is b) CHCl₂ c) : CCl_2 a) $CHCl_3$ d) $COCl_2$ 102 The production of an optically active compound from a symmetric molecule without resolution in termed as : 4. a) Walden inversion b) Partial racemisation c) Asymmetric synthesis d) Partial resolution 102 An organic compound on heating with CuO produces CO₂ but no water. The organic compound may be 5. b) Chloroform d) Ethyl iodide a) Carbon tetrachloride c) Methane 102 Which of the following statement is not applicable to Beilstein test? 6. a) Green or bluish green flame is due to the formation of volatile cupric halides b) It does not tell us to which halogen is present in the organic compound c) It is very sensitive test can be easily performed d) It is a sure test for the presence of halogen 102 Essential oils can be isolated by 7. a) Crystallization b) Steam distillation c) Sublimation d) Distillation 102 Mesomeric effect involves delocalisation of

8.

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102 9.	a) Pi-electrons The IUPAC name of the co	b) Sigma electrons ompound,	c) Protons	d) None of these
	is :			
	a) 1.2-dimethyl-2-butenc	ป		
	b) 3-methylpent-3-en-2-	ol		
	c) 3,4-dimethyl-2-buten-	4-ol		
	d) 2,3-dimethyl-3-penter	ıol		\frown
103	Which of the following sp	oecies is paramagnetic in na	ature?	
0.				
100	a) Carbonium ion	b) Free radical	c) Carbene	d) Nitrene
103	Isobutyl chloride is :			
1.	a) $(H_{-}($	b) (ርዝ ₋)-ርዝርዝ-ር]	c) ርዝ ₂ ርዝ2ርዝርነርዝ2	d) $(CH_{2})_{2}C = Cl$
103	How many isomers will (h_{c} H _c have?		
2.		3316 114701	A	X
	a) 1	b) 2	c) Zero	d) 4
103	Which one of the following	ng compounds is capable of	f existing in a <i>meso</i> form?	
3.				
	a) 3, 3-dibromopentane		b) 4-bromo-2-pentanol	
400	c) 3-bromo-2-pentanol		d) 2, 3-dibromopentane	
103	Geometrical isomerism is	s caused :		
4.	a) Purportricted rotation	around $C = C$ hand	G.XY	
	b) By the presence of one	a = 0 asymmetric carbon atom	N [°]	
	c) Due to different group	s attached to the same fund	rtional group	
	d) By swing of hydrogen	atom between two polyval	ent atoms	
103	Lassaigne's test is used for	or the detection of		
5.				
	a) Carbon only		b) Hydrogen only	
	c) Oxygen only		d) Nitrogen, sulphur and	halogens
103	Which of the following is	arranged according to the	nature indicated?	
6.	a) Electrophile			
	h) Flectronhile			
	c) Electrophile – $CH_{2}OH$	N_{-}^{-} Nucleonhile – NO_{+}^{+} Br ⁺	F	
	d) Electrophile – Br^+ N_{-}^-	Nucleonhile – CH_0OH		
103	The number of possible a	lkvnes with molecular forr	nula C- Ho is	
7.				
	a) 3	b) 4	c) 5	d) 6
103	Example of geometrical is	somerism is		
8.	×			
	a) 2-butanol	b) 2-butene	c) Butanal	d) 2-butyne
103	Which of the following is	an example of elimination	reaction?	
9.	a) Chlorination of mathe	no		
	a) Unior mation of ethance			
	c) Nitration of henzene	,ı		
	d) Hydroxylation of ethyl	lene		
104	The order of stability of t	he following carbanion is		

0. \ominus RCH₂ III IV Π I a) I>II>II>IV b) I>III>II>IV c) IV>III>II>I d) III>IV>I>II 104 Which nomenclature in IUPAC is not correct? 1. a) Pentyne-3 b) Pentyne-2 c) Hexyne-3 d) None of these 104 The C—C bond angle in cyclopropane is : 2. b) 120° c) 109°28' a) 60° d) 180° 104 Absolute alcohol cannot be obtained by simple fractional distillation because 3. a) Pure C_2H_5OH is unstable b) C₂H₅OH forms hydrogen bonds with water c) Boiling point of C_2H_5OH is very close to that of water d) Constant boiling azeotropic mixture is formed with water 104 Racemic compound has 4. b) 1:1 mixture of enantiomer and diastereomer a) Equimolar mixture of enantiomers d) 1:2 mixture of enantiomers c) 1:1 mixture of diastereomers 104 In case of a homologous series each member differs from the preceding or the succeeding member by : 5. b) a CH₃ group c) Two hydrogen atoms d) Four hydrogen atoms a) a CH₂ group 104 Vaccum distillation is used to purify liquids which 6. a) Are highly volatile b) Are explosive in nature c) Decompose below their boiling points d) Have high boiling point 104 The number of isomeric structures for C_2H_7N would be : 7. a) 4 c) 2 d) 1 b) 3 104 The IUPAC name of $CH_3 - CH_2$ 8. b) 2-methylbutanal c) Butanal-1 d) Pentanal-1 a) Propanal-1 104 Which of the following is the weakest base? 9. a) Ethyl amine b) Ammonia c) Dimethyl amine d) Methyl amine 105 The structure which has positive charge on the oxygen atom : 0. b) CH₃—Ö,—CH₃ с) СН₃—**ё:** d) н—**ё**—**ё**—н

ORGANIC CHEMISTRY - SOME BASIC PRINCIPLES AND TECHNIQUES

CHEMISTRY

														—
					:	ANS	W	ER K	EY :					
1)	d	2)	а	3)	b	4)	С	165)	а	166)	b	167) c	168)	а
5)	С	6)	d	7)	b	8)	С	169)	d	170)	d	171) d	172)	d
9)	b	10)	а	11)	с	12)	С	173)	b	174)	С	175) c	176)	b
13)	b	14)	b	15)	а	16)	d	177)	b	178)	d	179) d	180)	a
17)	d	18)	b	19)	С	20)	b	181)	b	182)	b	183) a	184)	d
21)	а	22)	С	23)	b	24)	а	185)	а	186)	a	187) c	188)	С
25)	b	26)	b	27)	b	28)	а	189)	d	190)	b	191) c	192)	d
29)	а	30)	d	31)	d	32)	С	193)	С	194)	а	1 95) c	196)	С
33)	С	34)	С	35)	d	36)	d	197)	b	198)	b	199) d	200)	d
37)	b	38)	b	39)	а	40)	С	201)	d	202)	b	203) a	204)	а
41)	b	42)	а	43)	а	44)	b	205)	а	206)	b	207) b	208)	а
45)	b	46)	b	47)	b	48)	d	209)	с	210)	а	211) a	212)	d
49)	d	50)	С	51)	d	52)	С	213)	d	214)	d	215) a	216)	d
53)	d	54)	С	55)	d	56)	С	217)	b	218)	С	219) a	220)	d
57)	d	58)	d	59)	а	60)	b	221)	a	222)	а	223) b	224)	С
61)	b	62)	а	63)	d	64)	b	225)	а	226)	b	227) d	228)	а
65)	а	66)	а	67)	d	68)	С	229)	d	230)	b	231) a	232)	С
69)	С	70)	b	71)	b	72)	С	233)	а	234)	С	235) b	236)	С
73)	d	74)	С	75)	d	76)	С	237)	С	238)	b	239) c	240)	С
77)	b	78)	d	79)	b	80)	С	241)	С	242)	d	243) b	244)	d
81)	С	82)	d	83)	b	84)	b	245)	d	246)	d	247) d	248)	С
85)	а	86)	С	87)	d	88)	d	249)	d	250)	b	251) a	252)	d
89)	b	90)	a 🖌	91)	а	92)	d	253)	С	254)	b	255) d	256)	С
93)	а	94)	b	95)	d	96)	С	257)	b	258)	b	259) b	260)	d
97)	а	98)	С	99)	С	100)	d	261)	а	262)	d	263) c	264)	b
101)	С	102)	d	103)	d	104)	a	265)	d	266)	d	267) a	268)	а
105)	d	106)	a	107)	С	108)	b	269)	С	270)	С	271) a	272)	С
109)	b	110)	С	111)	С	112)	b	273)	b	274)	С	275) b	276)	b
113)	а	114)	С	115)	d	116)	b	277)	d	278)	С	279) b	280)	С
117)	d	118)	С	119)	b	120)	С	281)	а	282)	b	283) b	284)	d
121)	a	122)	b	123)	а	124)	d	285)	С	286)	а	287) b	288)	С
125)	d	126)	b	127)	d	128)	d	289)	С	290)	а	291) d	292)	а
129)	d	130)	С	131)	d	132)	b	293)	b	294)	b	295) a	296)	а
133)	d	134)	С	135)	а	136)	b	297)	С	298)	а	299) b	300)	b
137)	а	138)	d	139)	С	140)	С	301)	а	302)	d	303) a	304)	а
141)	а	142)	d	143)	b	144)	b	305)	b	306)	b	307) b	308)	С
145)	С	146)	С	147)	a	148)	С	309)	а	310)	а	311) c	312)	С
149)	d	150)	С	151)	С	152)	С	313)	С	314)	b	315) d	316)	a
153)	b	154)	d	155)	d	156)	С	317)	С	318)	b	319) b	320)	b
157)	а	158)	С	159)	С	160)	С	321)	а	322)	а	323) b	324)	С
161)	а	162)	С	163)	d	164)	d	325)	С	326)	d	327) b	328)	d

329)	d	330)	а	331)	d	332)	a	533)	b	534)	а	535)	b	536)	b
333)	С	334)	d	335)	b	336)	С	537)	d	538)	b	539)	а	540)	d
337)	d	338)	b	339)	b	340)	d	541)	с	542)	С	543)	d	544)	b
341)	b	342)	С	343)	d	344)	С	545)	d	546)	b	547)	а	548)	а
345)	b	346)	С	347)	a	348)	b	549)	b	550)	а	551)	с	552)	d
349)	d	350)	а	351)	b	352)	d	553)	d	554)	d	555)	а	556)	d
353)	b	354)	b	355)	С	356)	а	557)	b	558)	а	559)	b	560)	d
357)	b	358)	С	359)	a	360)	b	561)	b	562)	а	563)	a	564)	b
361)	а	362)	С	363)	a	364)	а	565)	b	566)	а	567)	b	568)	b
365)	b	366)	b	367)	d	368)	b	569)	с	570)	b	571)	b	572)	с
369)	С	370)	С	371)	a	372)	С	573)	b	574)	b	575)	d	576)	d
373)	b	374)	b	375)	b	376)	d	577)	а	578)	d	579)	b	580)	а
377)	b	378)	С	379)	d	380)	С	581)	d	582)	d	583)	b	584)	b
381)	d	382)	а	383)	С	384)	С	585)	d	586)	d	587)	b	588)	b
385)	С	386)	а	387)	b	388)	b	589)	а	590)	a	591)	C	592)	а
389)	b	390)	b	391)	d	392)	а	593)	С	594)	a	595)	b	596)	С
393)	d	394)	а	395)	b	396)	d	597)	а	598)	b	599)	с	600)	d
397)	с	398)	С	399)	a	400)	b	601)	с	602)	a	603)	с	604)	а
401)	d	402)	b	403)	С	404)	С	605)	а	606)	d	607)	a	608)	С
405)	b	406)	а	407)	a	408)	b	609)	b	610)	d	611)	с	612)	d
409)	b	410)	d	411)	С	412)	а	613)	c	614)	а	615)	d	616)	d
413)	b	414)	b	415)	b	416)	b	617)	c —	618)	а	619)	с	620)	С
417)	а	418)	С	419)	С	420)	С	621)	a	622)	d	623)	а	624)	d
421)	С	422)	а	423)	С	424)	a	625)	b	626)	b	627)	с	628)	а
425)	а	426)	d	427)	d	428)	а	629)	d	630)	b	631)	С	632)	С
429)	С	430)	b	431)	b	432)	a	633)	d	634)	С	635)	d	636)	b
433)	а	434)	а	435)	d	436)	С	637)	d	638)	d	639)	d	640)	а
437)	d	438)	С	439)	a	440)	С	641)	b	642)	а	643)	d	644)	а
441)	С	442)	d	443)	a	444)	d	645)	а	646)	b	647)	С	648)	а
445)	С	446)	d	447)	с	448)	d	649)	b	650)	а	651)	b	652)	a
449)	С	450)	С	451)	d	452)	С	653)	d	654)	d	655)	а	656)	а
453)	d	454)	b	455)	d	456)	b	657)	b	658)	С	659)	b	660)	b
457)	b	458)	C	459)	С	460)	b	661)	b	662)	С	663)	С	664)	С
461)	d	462)	С	463)	а	464)	d	665)	b	666)	а	667)	b	668)	С
465)	b	466)	b	467)	С	468)	а	669)	d	670)	а	671)	С	672)	а
469)	а	470)	d	471)	а	472)	а	673)	b	674)	С	675)	С	676)	b
473)	a	474)	ď	475)	a	476)	d	677)	b	678)	b	679)	b	680)	b
477)	d	478)	b	479)	a	480)	b	681)	а	682)	а	683)	b	684)	d
481)	c	482)	d	483)	a	484)	d	685)	а	686)	а	687)	С	688)	d
485)	с	486)	b	487)	С	488)	b	689)	d	690)	С	691)	b	692)	d
489)	a	490)	а	491)	d	492)	С	693)	а	694)	С	695)	С	696)	а
493)	С	494)	d	495)	d	496)	d	697)	d	698)	а	699)	d	700)	С
497)	С	498)	а	499)	b	500)	d	701)	b	702)	а	703)	b	704)	b
501)	а	502)	d	503)	d	504)	b	705)	b	706)	С	707)	а	708)	d
505)	b	506)	b	507)	С	508)	а	709)	b	710)	С	711)	b	712)	а
509)	а	510)	С	511)	a	512)	С	713)	d	714)	С	715)	b	716)	С
513)	b	514)	b	515)	a	516)	d	717)	b	718)	а	719)	a	720)	b
517)	С	518)	d	519)	а	520)	d	721)	b	722)	С	723)	d	724)	b
521)	d	522)	b	523)	b	524)	С	725)	b	726)	С	727)	a	728)	а
525)	С	526)	а	527)	d	528)	d	729)	b	730)	b	731)	a	732)	а
529)	а	530)	С	531)	b	532)	a	733)	а	734)	b	735)	а	736)	d

737)	a	738)	а	739)	а	740)	d	897)	а	898)	d	899)	С	900)	a
741)	С	742)	С	743)	b	744)	С	901)	С	902)	С	903)	d	904)	С
745)	С	746)	а	747)	b	748)	a	905)	С	906)	а	907)	С	908)	d
749)	С	750)	а	751)	d	752)	а	909)	С	910)	d	911)	d	912)	а
753)	b	754)	d	755)	с	756)	b	913)	d	914)	а	915)	b	916)	d
757)	a	758)	b	759)	с	760)	a	917)	d	918)	b	919)	b	920)	С
761)	a	762)	а	763)	с	764)	d	921)	b	922)	b	923)	С	924)	d
765)	b	766)	b	767)	с	768)	b	925)	С	926)	a	927)	a	928)	b
769)	b	770)	а	771)	с	772)	С	929)	a	930)	a	931)	b	932)	d
773)	b	774)	а	775)	b	776)	С	933)	a	934)	a	935)	b	936)	d
777)	d	778)	а	779)	с	780)	С	937)	a	938)	d	939)	a	940)	а
781)	b	782)	а	783)	а	784)	С	941)	a	942)	b	943)	b	944)	d
785)	С	786)	с	787)	d	788)	а	945)	b	946)	b	947)	b	948)	а
789)	b	790)	d	791)	а	792)	b	949)	С	950)	a	951)	C	952)	d
793)	С	794)	а	795)	а	796)	а	953)	С	954)	с	955)	a	956)	b
797)	d	798)	d	799)	а	800)	С	957)	b	958)	a	959)	С	960)	С
801)	С	802)	b	803)	d	804)	d	961)	b	962)	b	963)	d	964)	d
805)	С	806)	d	807)	С	808)	d	965)	d	966)	a	967)	b	968)	b
809)	С	810)	а	811)	С	812)	С	969)	d	970)	С	971)	С	972)	d
813)	d	814)	с	815)	а	816)	а	973)	b	974)	a	975)	a	976)	b
817)	d	818)	а	819)	d	820)	b	977)	a	978)	a	979)	С	980)	a
821)	b	822)	b	823)	d	824)	d	981)	b	982)	С	983)	С	984)	С
825)	b	826)	d	827)	С	828)	С	985)	a	986)	d	987)	b	988)	b
829)	b	830)	d	831)	С	832)	d	989)	d	990)	а	991)	b	992)	d
833)	b	834)	С	835)	b	836)	b	993)	a	994)	а	995)	b	996)	b
837)	d	838)	С	839)	С	840)	С	997)	С	998)	С	999)	b	1000)	С
841)	С	842)	С	843)	b	844)	а	1001)	a	1002)	d	1003)	b	1004)	а
845)	С	846)	С	847)	с	848)	a	1005)	С	1006)	d	1007)	а	1008)	b
849)	С	850)	а	851)	b	852)	b	1009)	a	1010)	С	1011)	С	1012)	d
853)	С	854)	С	855)	b	856)	d	1013)	b	1014)	С	1015)	b	1016)	а
857)	С	858)	а	859)	d	860)	b	1017)	b	1018)	С	1019)	d	1020)	С
861)	b	862)	d	863)	d	864)	С	1021)	a	1022)	С	1023)	С	1024)	С
865)	b	866)	d	867)	b	868)	а	1025)	a	1026)	d	1027)	b	1028)	a
869)	d	870)	d	871)	b	872)	С	1029)	b	1030)	b	1031)	b	1032)	b
873)	а	874)	b	875)	d	876)	а	1033)	С	1034)	a	1035)	d	1036)	а
877)	d	878)	C	879)	d	880)	С	1037)	а	1038)	b	1039)	b	1040)	d
881)	С	882)	C	883)	а	884)	d	1041)	a	1042)	a	1043)	d	1044)	а
885)	d	886)	С	887)	С	888)	а	1045)	a	1046)	С	1047)	С	1048)	а
889)	b	890)	b	891)	b	892)	С	1049)	b	1050)	a				
893)	d	894)	С	895)	С	896)	С								
								I							
CN															
7															
~															

ORGANIC CHEMISTRY - SOME BASIC PRINCIPLES AND TECHNIQUES

CHEMISTRY

	: HINTS AND	SO	LUTIONS :
1	(d)		nitroprusside
	$C_{\rm H_{2}}$		(nurnle
n	$c_{6}n_{5}$ — c_{11} = $c_{11}c_{00}$ is chinaline actu.		colour)
Ζ		0	
-	Draw position and chain isomers.	9	
3	(b)		-do -
	Strain = $\frac{1}{2}$ [Normal valence angel-valence angel]	10	(a)
	1		Cyclopropyl methyl carbocations are more stable
	$=\frac{1}{2} [109^{\circ}28' - 60^{\circ}] = 24^{\circ}44'.$		than benzyl carbocations due to conjugation
6	(d) 2		between bent orbitals of cyclopropyl group.
0	2 2 dimethyl bytane is 6 carbon bydrocarbon	11	(c)
			The shape of π electron cloud in acetylene in
	(L_6H_{14})		cylindrical
	CH ₃	12	
			Acidified active fusion outpatt on addition of
	$\mathrm{CH}_3 - \mathrm{C} - \mathrm{CH}_2 - \mathrm{CH}_3$	\sim	
			ferric chloride solution gives blood red
	CH ₃		colouration, which confirms the presence of N and
	Rest all are the chain isomers of pentane (C_5H_{12}) .		S.
	CH ₃		$3NaCNS + aq. FeCl_3 \rightarrow Fe(CNS)_3 + 3NaCl$
			red
	$H_{2}C - C - CH_{2}$		(ferric thiocyanide)
		13	(b)
			СІ ОН
	CII3		
			\downarrow \downarrow
	(<i>neo</i> -pentane)		
	CH ₃		
			chlorobenzene phenol
	$H_3C - CH - CH_2 - CH_2$		In this process one group is replaced by other
	2- methyl butane		In this process one group is replaced by other,
	(iso –pentane)		nence, it is a substitution process and both the
	CH ₃ CH ₂ CH ₂ CH ₂ CH ₃		leaving and attacking groups are nucleophilic,
	<i>n</i> -pentane		therefore it is an example of nucleophilic
7	(b)		substitution reaction.
	Follow IUPAC rules.	14	(b)
8	(c)		(i) $Fe(CNS)_3$ is red in colour and is formed when
-	Detection of sulphur in sodium extract is done by		both N and S are present in organic compound
	lead acetate and sodium nitronrusside		(ii) $Na_2S + Na_2[Fe(CN)_5NO] \rightarrow$
	Na ₂ S + (CH ₂ COO). Ph \rightarrow PhS + 2CH COON ₂		[NaFe(C) ₅ NOS]
	$\frac{1}{1020} + \frac{1}{10000000000000000000000000000000000$		from organic sodium violet colour
	IEAU ACETALE DIACK PPL		compound nitroprusside
	$\operatorname{INa}_2 \circ + \operatorname{INa}_2[\operatorname{re}(\operatorname{LN})_5\operatorname{NU}] \to \operatorname{Na}_4[\operatorname{re}(\operatorname{LN})_5\operatorname{NUS}]$	15	(a)
	Sodium nitroprusside sodium thio	10	Bond $C = H C = C C =$
		I	

N C – O		
Bond energy (in kcal mol ^{-1}) 97	83	73
86		

16 **(d)**

1° carbon is attached to one carbon atom.

2° carbon is attached to two carbon atms.

3° carbon is attached to three carbon atoms.

The hydrogen attached to 2° carbon atom are 2°.



 \therefore It has one 2° carbon atom and two 2° hydrogen atoms.

17 **(d)**

-do-18 **(b)**

(b) Non-staggered means eclipsed form.

19 (c)

Carbanion is electron rich species. Stability of carbanion increases with increase in s —character of hybrid orbitals of carbon bearing the charge.

:. $sp^3 < sp^2 < sp$ (25%s-character) (33%s-character) (50%scharacter)

20 **(b)**



It is example of addition reaction

21 **(a)**

It is a fact.

22 **(c)**

It is the latest modified definition of organic chemistry.

23 **(b)**

In the lassaigne test, if organic compound consists of both N and S, then a red colour is obtained on adding aqueous

 $FeCl_3$ To sodium extract. $NH_2 - C - NH_2$ contains both N and S hence, it will give red colour in Lassaigne test NaCNS + $FeCl_3 \rightarrow [Fe(SCN)]Cl_2 + NaCl$ Blood red colour

24 **(a)**

If nitrogen is present in organic compound then sodium extract contains NaCN.

 $Na + C + N \xrightarrow{Fuse} NaCN$ $FeSO_4 + 6NaCN \rightarrow Na_4[Fe(CN)_6] + Na_2SO_4$ (A)A changes to Prussian blue $Fe_4[Fe(CN)_6]_3$ on reaction with FeCl₃. 4FeCl₃ + 3Na₄[Fe(CN)₆] \rightarrow Fe₄[Fe(CN)₆]₃ + 12NaCl 25 (b) Follow IUPAC rules. 27 (b) It is the definition of asymmetric synthesis. 28 (a) (i) has sp^3 ; (ii) has sp^2 , sp^3 ; (iii) has sp^2 , sp; (iv) 31 (d) The number of isomeric alkenes with molecular formula $C_6 H_{12}$ are 13. (1) $CH_2 = CH - CH_2 - CH_2 - CH_2 - CH_3$ (2) $CH_3 - CH = CH - CH_2 - CH_2 - CH_3$ (cis and trans) (3) $CH_3 - CH_2 - CH = CH - CH_2 - CH_3$ (cis and trans) CH₃ $(4)CH_2 = CH - CH - CH_2 - CH_3$ CH₃ $(5)CH_2 = C - CH_2 - CH_2 - CH_3$ CH₃ $(6)CH_2 = CH - CH_2 - CH - CH_3$ CH₃ $(7)CH_3 - CH = CH - CH - CH_3$ (cis and trans) CH_3 (8) $CH_3 - C = CH - CH_2 - CH_3$

CH₃

I

$$(9)CH_3 - CH = C - CH_2 - CH_3$$

(cis and trans)

32 **(c)**

It is 3-methyl butan-2-ol.

33 (c)

This statement is not true now.

36 (d)

Heterolysis involves the bond fission in a manner when either of the two atoms involved in bond fission retains the shared pair of electron, producing +ve and -ve ions, *e*.g.

34 (c)

$$\rightarrow$$
C-Cl \rightarrow \rightarrow C⁺+Cl⁻;Cl is more electronegative.

37 **(b)**

Keto and enol forms are inter convertable. The enol content will be maximum when enol form is stabilised by hydrogen bonding.

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

 $CH_{3} - C = CH - CH_{3} - CH_{3} - CH_{4} - C$

In acetyl acetone, the enol form is stabilised by Hbonding, hence it has more enol content than other.

38 **(b)**

Compounds having bivalent functional group (like C=O, -O-, -S - etc) with atleast 4 carbon atoms (in case of ether and thioether) or atleast 5 carbon atoms (in case of ketones) exhibit metamerism. Hence, $C_2H_5 - S - C_2H_5$ will show metamerism.

39 (a) Follow IUPAC ru

Follow IUPAC rules. 40 (c)

$$(CH_3)_3 + N(CH_3)_3 + CH_3 - C - CH_2CH_3 \xrightarrow{Alc.KOH} CH_3CH = CHCH_3 + H$$

is an example of elimination reaction

41 **(b)**

The dehydrohalogenation in presence of OH⁻is correctly represented by





OH⁻ In this mechanism the base OH⁻ removes a

proton from the β carbon.

43 **(a)**

Covalent bonds are cleaved in homolytic way in presence of UV light. It results in formation of free radical.

$$CI \xrightarrow{\bullet\bullet} CI \xrightarrow{UV} CI + CI$$

45 **(b)**

One asymmetric carbon atom is present.

46 **(b)**

Both alkene and cyclo alkane have general formula C_nH_{2n} .

IUPAC name of the above compound is 3-carboxyhexane-l, 6-dioic acid.

48 **(d)**

 $Na_2S + Na_2[Fe(CN)_5NO] \rightarrow Na_4[Fe(CN)_5NOS]$ Solution introprusside sodium thio nitro Solution prusside (purple colure)

49 **(d)**

50

Greater the difference in electronegativity of bonded atoms easier will be heterolytic cleavage (c)

$$HO - CH_2 - CH_2 - F$$



Gauche conformation is comparatively more stable due to hydrogen linkage in between F and H (at O-atom), hence order is *Eclipse*, *Anti* (staggered), *Gauche*.

51 **(d)**

Phosphorous is estimated as $Mg_2P_2O_7$ P $\xrightarrow{HNO_3}_{\Lambda}$ H_3PO_4

$$\begin{split} &H_3 PO_4 + Mg^{2+} + NH_4 OH \rightarrow MgNH_4 PO_4 \\ &2MgNH_4 PO_4 \rightarrow Mg_2 P_2 O_7 + H_2 O + 2NH_3 \\ &\% \text{ of } P = \frac{62 \times \text{wtof } Mg_2 P_2 O_7 \times 100}{222 \times \text{w}} \end{split}$$

n = 2 and $a = 2^n = 2^2 = 4$.

54 **(c)**

Among carbonyl compounds, reactivity decrease with increase in alkyl groups as alkyl groups (+*I* effects) decrease positive character on Catom. Thus, the correct order of reactivity is HCHO > CH_3 CHO > C_6H_5 CHO

55 **(d)**

Primary and secondary alkyl halides gives $S_N 2$ reaction

56 **(c)**

Follow IUPAC rules.

57 **(d)**

2-butanol is optically active as it contain as it contain chiral carbon atom.

CH

$$CH_{3} - CH_{2} - C - OH$$

H
58 (d)

$$a = 2^n; n = 3.$$

59 (a)

It is Markownikoff's rule.

61 **(b)**

The order of homolytic bond dissociation energies of CH_4 , C_2H_6 and CH_3Br is as

$$CH_4 > C_2H_6 > CH_3Br$$

 $\Delta H \text{ (kcal/mol)}$ 105 100 70

63 **(d)**

Each d form has its l form and the pair is known as enantiomer.

64 **(b)**

Carboxylic acid and esters show functional group isomerism. When two compounds have same molecular formula but different functional groups, then functional isomerism arises.

e.g.,

C₂H₅COOH and CH₃COOCH₃

65 **(a)**

One asymmetric carbon atom is present in a lactic acid molecule. Hence, it is an optically active compound.

$$H = \begin{bmatrix} CH_3 \\ Asymmetric carbon atom \\ COOH \\ COOH \end{bmatrix}$$

66 **(**

According to IUPAC system, ether are named as alkoxy alkanes. The larger alkyl group froms the parent chain while lower alkyl group is taken ethereal oxygen and forms a part of alkoxy group.

$$CH_{3} - CH_{2} - O - CH_{2} - CH_{2}$$

Follow IUPAC rules.

68 **(c)**

It is a fact.

69 **(c)**

A liquid, which decomposes at its normal boiling point can be purified by vacuum distillation.

71 **(b)**

Stability of carbanion is not governed by hyperconjugation. Its stability depends on the +I or -I group

72 **(c)**

Let unreacted $0.1M(= 0.2N)H_2SO_4 = V'mL$ $\therefore 20 \text{ mL of } 0.5M \text{ NaOH}$ $=V'mL \text{ of } 02 \text{ N } H_2SO_4$ $\therefore 20 \times 0.5 = V' \times 0.2$ $\therefore V' = 50mL$ Used $H_2SO_4 = 100 - 50 = 50mL$ $\% \text{Nitrogen} = \frac{1.4 \text{ NV}}{W}$ where, N=normality of H_2SO_4 $V=\text{volume of } H_2SO_4 \text{ used}$ $\therefore \% \text{ nitrogen} = \frac{1.4 \times 0.2 \times 50}{0.30}$ % of nitrogen in

(a)
$$CH_3CONH_2 = \frac{14 \times 100}{59} = 23.73\%$$

(b) $C_6H_5CONH_2 = \frac{14 \times 100}{122} = 11.48\%$
(c) $NH_2CONH_2 = \frac{28 \times 100}{60} = 46.67\%$
(d) $NH_2CSNH_2 = \frac{28 \times 100}{76} = 36.84\%$

= 46.67%

Therefore, the organic compound is urea.

73 (d)

 $CH_4 + Cl_2 \xrightarrow{hv} CH_3Cl$ This is an example of free radical substitution reaction

$$CI \longrightarrow CI^{\bullet} + CI^{\bullet}$$

$$CH_{4} + CI^{\bullet} \longrightarrow CH_{3}^{\bullet} + HCI$$

$$CH_{3}^{\bullet} + CI - CI \longrightarrow CH_{3}CI + CI^{\bullet}$$

$$CI^{\bullet} + CI^{\bullet} \longrightarrow Cl_{2}$$

74 (c)

> Glucose has four dissimilar asymmetric carbon atoms; $a = 2^4$.

75 (d)

The elimination takes place according to Saytzeff rule. The most substituted alkene (butane-2) is called Saytzeff product whereas less substituted alkene (butane-1) is called Hofmann product

76 (c)

 $-NO_2$ group shows -M effect while CH_3O - group shows +M effect. (-M effect stabilizes an anion) Hence, the order of stability is



77

78

The former possesses 12α -H atom whereas, later possesses six α -H atom. More is the no. of α -H atom, more is delocalisation and more is stability. (d)

 H_2O , ROH, R – COOH etc are protic solvents because they are polar in nature and contain a hydrogen directly bonded to oxygen while other are aprotic solvents as they do not have a hydrogen bonded directly to oxygen. They are especially favourable for $S_N 1$ reactions. While aprotic solvents cannot have hydrogen bond to the nucleophile because they does not have

hydrogen bonded to nitrogen or oxygen. They are favourable for $S_N 2$ reactions.

79 **(b)**

Follow mechanism of addition reactions.

80 (c)

 $\cap \cap \sqcup$

Methoxy group is electron releasing group it increases electron density of benzene nucleus

- NO₂ group is electron withdrawing group, it decreases the electron density of benzene nucleus. Thus, the order of reaction with electrophilic regent is

$$NaCN + HNO_3 \xrightarrow{\Delta} HCN \uparrow + NaNO_2$$

$$Na_2S + HNO_3 \rightarrow H_2S \uparrow + 2NaNO_3$$

81

The central carbon in I and II is asymmetric.



Thus, out of four isomers only two have chiral carbon. Each have two isomers.

85 (a)

$$| \sum_{CH_2}^{CH_2} > 0$$

The structural formula of epoxide is CH_2 It consists three membered ring with two carbon and one oxygen.

86 (c)

87

$$\begin{array}{l} \mathrm{CH}_3 - \mathrm{CH}_2 - \mathrm{CH}_2 - \mathrm{Cl} + \mathrm{alc.} \, \mathrm{KOH} \longrightarrow \mathrm{CH}_3 - \mathrm{CH} \\ &= \mathrm{CH}_2 \end{array}$$

It is an example of elimination reaction

(d)

$$>C=O+H_2NOH \rightarrow >C=NOH+H_2O$$

Both addition and elimination takes place simultaneously. Thus, the reaction is addition elimination

88 (d)

Resolution of racemic mixture involves the

formation of distereoisomers.

89 **(b)**

When N and S both are present in the organic compound, then a red colour complex ion of $[Fe(CNS)]^{2+}$ is formed on adding $FeCl_3$ to sodium extract

 $NaCNS + FeCl_3 \rightarrow [Fe(SCN)]Cl_2 + NaCl$ blood red colour

90 **(a)**

$$CH_3CH_2Br \xrightarrow{H-H} CH_3 - CH_3 + HBr$$

 S_{N^2}

$$CH_{3}$$

$$|$$

$$CH_{3} - C - Br \xrightarrow{LAH} CH_{3} - C = CH_{2} + HBr$$

$$|$$

$$(E_{2}) \qquad |$$

$$CH_{3} \qquad CH_{3}$$

91 **(a)**

Count σ and π -bonds.

92 **(d)**

The formation of sodium thionitroprusside (blue) shows the presence of sulphur.

 $Na_2S + Na_2[Fe(NO)(CN)_5]$ $\rightarrow Na_4[Fe(CN)_5NOS]$

Sodium sodium nitroprusside sodium thionitroprusside purple extract

93 **(a)**

It is a fact.

94 **(b)**

 $R - CH_2 - CH_2X + KOH(alc.) \rightarrow R - CH$ $= CH_2 + KX + H_2O$

Alkyl halid undergo β -elimination to form alkene.

95 **(d)**

The rate of reaction follows the order : RI > RBr > RCl > RF; whether it obeys S_N1 or S_N2 mechanism due to steric hindrance of alkyl group.

(c) Only this is optically active due to central carbon being asymmetric.

97 **(a)**

96

Ethers show metamerism. Metamerism arises when a polyvalent functional group

$$(e.g., -0, >c=0 \text{ etc.})$$
 is attached to different alkyl groups but the molecular formula

remains same e.g., $C_2H_5 - O - C_2 H_5$ and $CH_3 - O - C_3H_7$ Carbanion is electron rich species. Stability of carbanion increases with increase in s —character of hybrid orbitals of carbon bearing the charge.

: $sp^3 < sp^2 < sp$ (25%s-character) (33%s-character) (50%scharacter)

99 **(c)**

The Z repells electrons and thus, electron density increases on R_3C part.

100 **(d)**

∵ -COOH group is a deactivating group
 ∴ Benzoic acid is less reactive towards
 electrophilic substitution.
 So, benzoic acid> phenol>*n*-propyl benzene is
 not arranged correctly.

101 **(c)**

Aqueous solution of $NaHCO_3$ can be used to separate benzoic acid from its mixture with camphor. Benzoic acid form water soluble sodium benzoate with $NaHCO_3$.

102 (d)

Each π -electron is delocalised over six carbon atoms in ring.

103 **(d)**

Grignard reagent reacts with $>C=0, -C \equiv N, > C = S$ as follows

$$>C=O + RMg X \longrightarrow C OH + Mg < OH X$$

$$-C = N + RMg X \longrightarrow -C = O + Mg \bigvee_{X}^{R}$$
$$>C = S + RMg X \longrightarrow C - SH + Mg \bigvee_{X}^{OH}$$

104 **(a)**

Benzene is the most stable and we know that resonance energy is a direct measure of the stability of a molecule

105 (d) O C H HCN O CN CN

It is an example of nucleophilic addition reaction. 107 (c) CH₃ $H_3C - C - Br$ and $CH_3CH_2CH_2CH_2Br$ are chain as well as position isomers.

108 **(b)**

Detection of phosphorus in the organic compound can be done by its conversion into phosphate. The ammonium molybdate is used to identify phosphate ion

 $Na_3PO_4 + 3HNO_3 \rightarrow H_3PO_4 3NaNO_3$

$$H_3PO_4 + 12(NH_4)_2MoO_4 + 21HNO_3 -$$

ammonium modybdate

$$(NH_4)_3PO_4.12MoO_3 + 21NH_4NO_3 + 12H_2O$$

yellow ppt.

109 **(b)**

Homolytic bond fission is one in which each entity involved in bond formation retains its electron involved in shared pair of electron to form free radicals.

110 **(c)**

 CH_3CN has sp^3 and sp-hybridised carbon atom.

111 **(c)**

 CH_3OCH_3 , C_2H_5OH Methoxy methaneEthanol (ether)(alcohol)functional group (C_2H_6O) (C_2H_6O) molecular formulaIn methoxy methane and ethanol both molecularformula is same but functional groups aredifferent, so they are functional isomers.

112 **(b)**

The main conditions for exhibiting geometrical isomerism are

(i) Presence of double bond.

(ii) Presence of different groups on same double bonded carbon.

(iii) Presence of at least one similar group on adjacent double bonded carbon atoms.

 $C_3H_6(H_3C \longrightarrow CH \implies CH_2)$ does not exhibit geometric isomerism due to presence of same group on double bonded carbon atom (C_1).

113 **(a)**

Follow mechanism of addition reaction.

114 **(c)**

Vicinal or alkylene dihalides.

115 **(d)**

The electrophile involved in the sulphonation of

benzene is SO_3 .

$$2\mathrm{H}_2\mathrm{SO}_4 \rightarrow \mathrm{SO}_3 + \mathrm{H}_3\mathrm{O}^+ + \mathrm{HSO}_4^-$$

116 **(b)**

Number of meso structures in compound having odd number of chiral carbon atoms and

symmetrical molecule
$$=2^{\left(\frac{n}{2}-\frac{1}{2}\right)}$$

Given, $n = 5$

 $\therefore \text{ Number of } meso \text{ forms} = 2^{\frac{5}{2}-\frac{1}{2}} = 2^2 = 4$

117 **(d)**

In Kjeldahl's method, the nitrogen is quantitatively converted into ammonia by heating with NaOH

$$C + H + N \xrightarrow{\Delta} (NH_4)SO_4 + CO_2 + H_2O$$

(from organic compound) $(NH_4)_2SO_4 + 2NaOH \rightarrow Na_2SO_4 + 2NH_3 + 2H_2O$ In Dumas method nitrogen present in organic compound is quantitatively converted into N₂.

118 **(c)**

It is a fact. 119 **(b)**

9 (b)
(CH₃)₃ C
$$\rightarrow$$
 OH $\xrightarrow{+H^+}_{-H_2O}$ (CH₃)₃C
3° carbocation
(more stable)
CH₂=CH \rightarrow CH₂ \rightarrow CH₂OH $\xrightarrow{+H^+}_{-H_2O}$
CH₂=CH \rightarrow CH₂ \rightarrow CH₂

Increasing order of stability of carbocation. 1°carbocation <2° carbocation<3°carbocation

120 **(c)**

Both geometrical and optical isomerism are included in stereoisomerism.

121 **(a)**

Phenol reacts with chloform and NaOH to give o-hydroxy benzaldehyde or salicylaldehyde. In this reaction dichlorocarbene (: CCl_2) electrophile is generated. This reaction is called Reimer-Tiemann reaction.

$$OH^{-} + CHCl_{3} \longrightarrow HOH + CCl_{3}^{-}$$

$$unstable$$

$$:CCl_{3}^{-} \longrightarrow Cl^{-} + CCl_{2}^{-}$$

dichlorocarbene

122 **(b)**

Homolytic fission is favoured by sunlight. In it, each bonded atom takes away its shared electrons and thus free radicals are produced.

 $CI \longrightarrow CI^{\bullet} + CI^{\bullet}$

124 (d)







The closed ring cycloalkanes beyond five carbon atoms has puckered ring structure maintaining tetrahedral nature or stainless rings, e.g., cyclo hexane has chair and boat form.

These are characteristics of $S_N 1$ mechanism.

-I power of groups in decreasing order with respect to the reference H $NO_{2} > CHO > COOR > F > Cl > Br > l > OH$

$$> OR > NH_2$$

Asymmetry is present in all states.

132 (b)

The structure of cyclopropane, cyclobutane and cyclo hexane are as



Hence, the common group in cyclopropane, cyclobutane and cyclohexane is $>CH_2$ group.

Draw all possible isomers.

+I effect is shown by - CH₃ while - I effect is shown by -Br, -Cl and $-NO_2$.

It is a strong reducing agent.

% of C=
$$\frac{12}{44} \times \frac{0.535}{0.765} \times 100 = 19.07$$

% of H = $\frac{2}{18} \times \frac{0.138}{0.765} \times 100 = 2.004$
Ratio of % of C:H=19:2 (approx.)

Benzene has planar structure.

127 (d)

 $CH_3CHOHC_2H_5$ is optically active because it has chiral C*-atom



Unsaturation (double bond) is given priority over halogen, then lowest set of locants. So, the correct IUPAC name is 3-bromo-1-chlorocyclohexene.

140 (c)

Follow conformation; The conformers for n-butane are two gauche, two eclipsed and one anti.

142 (d)

Select longest possible carbon atom chain, number it and name compound according to IUPAC, rules.



3, 3-diethyl-4-methyl-5-(1'-methyl ethyl)-octane

143 **(b)**

Benzal is C_6H_5CH group.

144 **(b)**

A 2° carbon is one of which two valencies are attached to carbon atom.

145 **(c)**

Eclipsed conformation of butane contain angle and steric strain both. Follow conformation.

146 **(c)**

Weight of organic compound =29.5mg $NH_3 + HCl \rightarrow Na_4Cl$ $HCl + NaOH \rightarrow NaCl + H_2O$ (remaining) $15 \times 0.1 M$ =1.5 mmol Total millimole of HCl=2 Millimole used by $NH_3 = 2 - 1.5 = 0.5$ Weight of $NH_3=0.5 \times 17mg=8.5mg$ Weight of nitrogen $=\frac{14}{17} \times 8.5mg = 7mg$ % Nitrogen $=\frac{7}{29.5} \times 1100 = 23.7\%$

147 (a)

Fluoro group causes negative inductive effect increasing ionisation, thus 0.1M difluoroacetic acid has highest electrical conductivity.



148 (c)

 $-NO_2$ group shows -M effect white CH_3O -group shows +M effect (-M effect stabilises an anion)

149 **(d)**

When the groups with higher priority (*i. e.*, with high atomic number) are present on same side of double bond, then the configuration is Z but when present on opposite side of double bond, the configuration is E.



150 (c)

Nucleophiles are electron rich species and can donate lone pair of electron to carbocation or any +ve centre.

152 **(c)**

 C_4H_6 can have five cyclic isomers.



153 (b) -do 154 (d) Urea shows tautomerism as

 $\begin{array}{ccc} 0 & OH \\ \parallel & \parallel \\ NH_2 - C - NH_2 \leftrightarrows NH = C - NH_2 \\ \text{keto form} & \text{enolic form} \end{array}$

155 (d) The conditions given are for C_6H_6 .

156 **(c)**

Count π -bonds. Delocalisation is not possible.

Hence, correct IUPAC name is bicyclo [2,2,0] hexane.

172 (d)

Chlorinolysis involves substitution reactions by chlorine.

173 (b)

 $a = 2^{n-1}$; where *n* is no. of asymmetric carbon; when molecule possesses symmetry.

1

3-chlorobutanal \therefore The order is priority is -CHO175 (c)

% C =
$$\frac{12}{44} \times \frac{12.517}{4.0} \times 100 = 85.7$$

% H = $\frac{2}{18} \times \frac{5.143}{4.0} \times 100 = 14.3$
The mole ratio of C to H is $\frac{85.7}{12} : \frac{14.3}{1}$
= 7.14: 14.3 = 1: 2 = CH₂

176 **(b)**

3-chloro-2, 3-dimethyl pentane contains all the four 1°,2°, 3° and 4° carbon atoms.



3-chloro-2-3-dimethyl pentane

178 (d)

hybridized

The carbon, four valencies of which are satisfied by four different groups, is termed as chiral carbon atom. The structures of the given compounds are as



Hence, succinic acid does not contain any chiral carbon atom.

180 (a)

In homolysis, the covalent bond is broken in such a way that each resulting species known as free radical.

$$CH_3CH_2$$
— $CI \xrightarrow{Homolytic} CH_3CH_2 + CI$
fission

181 **(b)**

p-orbitals are at 90° to each other.

182 **(b)**

 $S_{\rm N}2$ reactions are greatly controlled by steric factor.

 $\begin{array}{c} R-CH_2-X R_2 CH-X R_3 C-X \\ 1^\circ \qquad 2^\circ \qquad 3^\circ \end{array}$

 $S_N 2$ reactivity decreases as bulkyness of alkyl group increases.

183 (a)

$$\begin{array}{c|c}
 & \mathsf{OH} \\
 & 5 & 4 & 3 \\
 & \mathsf{CH}_3 & \mathsf{CH}_2 & \mathsf{CH}_2 & \mathsf{CH}_2 & \mathsf{CH}_2 \\
 & & & & \\
 & & & & \\
 & & & \mathsf{CH}_3
\end{array}$$

3-methyl pentane 3-ol

Hydroxy is used when – OH group is written in prefix. So, choice (b) and (c) are wrong.

184 (d)

Due to -R effect of – CHO group, oxygen carries – δ charge while the terminal carbon carries + δ , *ie*,

$$^{+\delta}$$
 CH=CH-CH=O

185 **(a)**



Its IUPAC name is 3,5-dimethylcyclohexene. 186 (a)

Lactic acid obtained in the given reaction is an optically active compound due to the presence of chiral C-atom. It exists as *d* and *l* forms whose ratio 1:1.



187 (c)

The mirror-image isomerism is a class of stereoisomerism and are included in optical isomerism.

188 **(c)**

 CH_3^+ acts as an intermediate in the given reaction (Friedel Craft's alkylation). It is an example of electrophilic aromatic substitution. In this reaction CH_3^+ is electrophile.

189 **(d)**

isomers = 2^n

$$= 2^2 = 4$$

190 **(b)**

 C_6H_6 has 12σ and 3π -bonds.

191 **(c)**

For $S_N 2$ mechanism, there should be least steric hinderance.

193 **(c)**

Rest all are polymerisation.

194 **(a)**

The increasing order of inductive effect is: $-F < -COOH < -CN < -NO_2$.

197 **(b)**

-COOH is on top in preference table.

200 **(d)**

In homolytic cleavage, covalent bond is cleaved in such a way that each atom takes its shared electrons with itself and free radicals are formed.

$$A \stackrel{\text{Homolytic fission}}{\longrightarrow} A^{\bullet} + B^{\bullet}$$

free radicals

201 (d)

Fractional distillation is used for the separation of

crude petroleum into various fractions like coal-— COOH is electron withdrawing group. tar, crude alcohol and petroleum 213 (d) 202 **(b)** CHO Free radical has unpaired electron. is called glyoxal 203 (a) CHO SO_3 can accept lone pair of electron in *d*-subshell. 214 (d) 204 (a) IUPAC name of acraldehyde From Kjeldahl's method, $(CH_2 = CH. CHO)$ is: Percentage of nitrogen $=\frac{1.4\times N\times V}{W}=\frac{1.4\times 0.1\times 30}{5}$ prop-2-ene-1-al = 0.84%216 (d) 205 (a) These are characteristics known from mechanism The isomers alkanes having the molecular of reaction. formula $C_5 H_{12}$ are as 217 (b) (ii) (i) *n*-pentane iso-pentane enolic form of acetone (2-methyl butane) 9σ bonds, 1π bond, 2 lone pairs 218 (c) (iii) BrCH₂CH₂COOH is the weakest acid and have lowest dissociation constant because. I.E. of Br is neo-pentane (2,2-dimethyl propane) lesser than F and is far away from – COOH group. 206 **(b)** 219 (a) Just after few years when Wöhler prepared urea Follow IUPAC rules. from KCNO and $(NH_4)_2SO_4$, Kolbe prepared acetic 220 (d) acid in laboratory from its element and gave final Molecular formula C_2Br ClFl six isomers are blow to Vital force theory. possible. 207 (b) Br Br Due to the presence of asymmetric carbon atom, e.g., CI CH₃CH₂•CHOHCH Br 208 (a) sp^2 Br CH OH C CH_3 sp^3 221 (a) Nucleophilies are those substances which can All bonds are σ - bonds hence C uses only its sp^3 donate a pair of electrons. They can be neutral or hybrid orbitals. In all other compounds there is negatively charged. The nucleophilic power one C = O double bond, therefore, this carbon is depends on the tendency of species to donate the sp^2 - hybridized electrons. Due to the presence of +I effect it 210 (a) increases. Hence, higher the +I effect, higher the Follow Baeyer's strain theory for stability of nucleophilic power. The +I effect of ethyl is cycloalkane. greater than +I effect of methyl group 211 (a)
0



 C_6H_6 has more canonical forms.

247 **(d)**

Chemical methods are based upon the distinguishing chemical properties of one class of organic compounds from the other. for example camphor and benzoic acid

248 **(c)**

Fractional crystalliation is used to purify organic solids which dissolve in a particular solvent. But their rate of solubility is different

249 **(d)**

NH₄CNO is functional isomer of urea. 250 **(b)**

$\stackrel{\Theta}{\text{Carbanion}} ^{(\text{CH}_3)}$

Here, the carbon atom carries a negative charge with lone pair of electrons and has eight electrons in outermost orbit and complete its octet.

H Reactions in which carbanions are formed as intermediate are said to proceed by a "Carbanion mechanism".

Carbanion is sp^3 hybridised, three sp^3 hybrid orbitals form covalent bonds with three atoms while the fourth sp^3 hybrid orbital has a nonbonding pair of electrons. It is pyramidal in shape as similar to NH₃.

251 **(a)**

Ether group(-0)has propyl and isopropyl group on its two sides.

252 **(d)**

Follow carbenes.

$$CH_3 \longrightarrow CH \longrightarrow CH \longrightarrow CH_3$$

There are two chiral C-atoms (*)

Thus, optical isomerism is possible.

254 **(b)**

Follow IUPAC rules.

255 **(d)**

-do -

256 **(c)**

A monosubstituted benzoic acid is stronger than a monosubstituted phenol as former being a carboxylic acid. Among the given substituted benzoic acid, ortho – hydroxy acid is strongest acid although - OH causes electron donation by resonance effect which tends to decreases acid strength. It is due to very high stabilisation of conjugate base by intramolecular H-bond which outweight the electron donating resonance effect of – OH.



The overall order of acid-strength of given four acids is *ortho*-hydroxybenzoic acid ($pK_a =$ 2.98) > Toluic acid pk_a = 4.37) > p –hydroxybenzoic acid $(pk_a = 4.58) > p$ –nitrophenol $(pk_a = 7.15)$. 257 (b)

Electrophiles are electron pair acceptors.

258 **(b)**

CH₃CHClCH₂CH₂CH₃; CH₃CH₂CHClCH₂CH₃; CH₂ClCH₂CH₂CH₂CH₃.

259 (b)

Benzene and all its derivatives along with heteroaromatics possess aromatic nature. Rest all possess aliphatic nature.

260 (d)

Neohexyl chloride is a primary halide as in it, Clatom is attached to a primary carbon.

CH₃

$$CH_3 - C - CH_2 - CH_2CI$$

$$|$$

$$CH_3$$

261 (a)

Ethane, ethene and ethyne have sp^3 , sp^2 and sp-270 (c) hybridization respectively.

262 (d)

Carbon is asymmetric as all its valencies are attached to different groups.

264 **(b)**

Follow IUPAC rules.

265 (d)

Bromination of alkanes in the presence of sunlight involves the formation of free radical, e.g.,

$$CH_4 \xrightarrow{Br_2} CH_3Br$$

Mechanism

$$Br \longrightarrow Br + Br$$

Propagation

$$CH_4 + Br \longrightarrow CH_3^{\bullet} + HBr$$

$$CH_3^{\bullet} + Br \longrightarrow Br \longrightarrow CH_3Br + Br$$
Termination
$$Br + Br \longrightarrow Br_2$$

266 (d)

The two butene give different products on addition of Br2-cis butene gives racemic mixture whereas trans butene gives meso form of 2,3,3dibromo butene.

267 (a)

Lassaigne's test is used for the detection of halogens, nitrogen and sulphur.

268 (a)

Resonance and inductive effect decide stability of carbocations.



∴Correct order of stability is II<I<III<IV

269 (c)

Atom	At mass (a)	% (b)	$\frac{b}{a}$	Ratio
X	10	50	$\frac{50}{10} = 5$	2
Y	20	50	$\frac{50}{20} = 2.5$	1

Hence, empirical formula = $X_2 Y$

The compounds which differ in the nature of carbon chain are called chain isomers, e.g.,

$$CH_3 - CH_2 - CH_2 - CH_2 - CH_3$$

n-pentane

$$CH_3$$

$$|$$

$$CH_3 - CH - CH_2 - CH_3$$

iso-pentane

$$CH_3$$

$$|$$

$$H_3C - C - CH_3$$

$$|$$

$$CH_3$$

neo-pentane

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272 **(c)**

In conjugated diene alternate single and double bonds are present while in cumulative diene, double bonds are present at adjacent positions.

I.
$$H_2C = CHCH_2 CH_2 CH_2 CH_2 CH_2 CH_2$$

hexa-1, 5-diene

 \Rightarrow It is an isolated diene.

II.
$$H_2C = C = CH_2$$

prop-1, 2-diene

 \Rightarrow It is a cumulative diene.

$$\lim_{n \to \infty} \frac{1}{2} = 2 + 3 + 5 + 6$$
$$\lim_{n \to \infty} \frac{1}{2} = 2 + 2 + 3 + 6$$
$$\lim_{n \to \infty} \frac{1}{2} + 2 + 3 + 6$$

hexa-1, 3-diene

 \Rightarrow It is a conjugated diene.

IV.
$$H_2C \xrightarrow{2} CH \xrightarrow{3} CH \xrightarrow{4} CH_2$$

buta-1, 3-diene

 \Rightarrow It is a conjugated diene.

V. Prop -1, 2-diene is a cumulative diene. Hence, statement 3 and 5 are correct.

276 **(b)**

Phenols are much more acidic than alcohol due to the stabilisation of phenoxide ion resonance.



phenol phenoxide ion

Phenoxide ion is stabilised due to following resonating structures.



$$R \longrightarrow R \longrightarrow R \longrightarrow R^{+} H^{+}$$

alcohol (Not stabilised due to
absence of resonance)

ortho nitrophenol is most acidic because in – NO_2 electron attracting group is attached to ortho position which helps in stabilising the negative charge on the oxygen of phenoxide ion. Hence, due to this reason acidic character of phenol is increased, while on attachment of – CH_3 group (electron donating group) acidic strength of phenol is decreased in cresol due to the destabilisation of phenoxide ion.

277 (d)
C: H: 0 = 6: 1: 8

$$= \frac{6}{15} \times 100: \frac{1}{15} \times 100: \frac{8}{15} \times 100$$
40: 6.67: 53.3

$$= \frac{40}{12}: \frac{6.67}{1}: \frac{53.3}{16}$$
= 1: 2: 1 ie, CH₂O
278 (c)

Alkanes never show functional isomerism, metamerism, tautomerism and geometrical isomerism.

279 **(b)**

Draw all possible chain and position isomers.

2-butanol has following structure OH

$$R \rightarrow C \longrightarrow OH \longrightarrow 3^{\circ}alcohols$$

281 **(a)**

Stronger is acid, weaker is its conjugate base or weaker is nucleophilicity.

The acidic order $HF > H_2O > NH_3 > CH_4$.

282 **(b)**

Electromeric effect occurs only in the presence of attacking reagent. It operates in the molecules having multiple bonds. Since, it exists only on the demand of attacking reagent, it is a temporary effect. *e.g.*,



283 **(b)**

Leaving group ability depends upon basicity of group.

284 (d)

Read optical activity.

285 (c)

The compounds must fulfill two conditions to show geometrical isomerism.

(i) The compound should have at least one C=C.

(ii) The two groups attached to same carbon must be different.

Out of given choices only (c) fulfill both conditions and shows geometrical isomerism.

H - C - COOH

II

H - C - COOH

(c)

2-butene-1,4-dioic acid

286 (a)

C – Cl bond is aryl chloride is stable due to delocalisation of electron by resonance. Also C – Cl bond possess a double bond character like vinyl chloride, hence S_N reactions are not possible in chlorobenzene under ordinary conditions.



287 **(b)**

C - C, C = C and $C \equiv C$ bond length are 1.54 Å, 1.34Å and 1.20Å respectively. In benzene C = C is 1.40Å.

288 **(c)**

Definition of tautomerism.

289 **(c)**

The reactivity order is $3^{\circ}H > 2^{\circ}H > 1^{\circ}H$.

The +ve inductive effect of CH_3 group on carbanions intensifies negative charge on C^- centre and thus, 3° carbanion is more reactive.

291 **(d)**

Glycerine contains ^{α}CH₂ − OH ← 1° alcohol | ^{β}CH − OH ← 2°alcohol

 $^{\alpha}CH_2 - OH \leftarrow 1^{\circ} alcohol$

292 **(a)**

In rest all carbon chain is same.

293 **(b)**

In Cannizzaro reaction the transfer of H^- to another carbonyl group is difficult and slowest step. (rate determining step or key step)



294 **(b)**

Anti conformation is the most stable form of *n*-butane (Bulky groups far apart).

 $R - X \longrightarrow R^+ + X^- \xrightarrow{\mathrm{OH}^-} R - \mathrm{OH}.$

It has altogether different groups.

It is an ether and the name of ether is given as alkoxy alkane. So, its name is methoxy benzene. 300 **(b)**

butane-2-3-diol

Where *C*^{*}=asymmetric C atom

It is a symmetrical molecule, so the number of

optically active stereomers= 2^{n-1}

(*n*=number of asymmetric C atom)

$$= 2^{2-1}$$

 $=2^{-1}=2$

302 **(d)**

Wöhler prepared urea from KCNO and (NH₄)₂SO₄

304 **(a)**

Electron donors having lone pair of electrons are nucleophile.

(i) BF_3 is not nucleophile because it does not have lone pair of electrons. It is infact Lewis acid because it accepts pair of electron

(ii)NH₃, CN^- and OH^- all have lone pair of electrons, so they are nucleophiles.

305 **(b)**

Due to asymmetric carbon atom.

306 **(b)**

Aryl carbonium ions are more stable than alkyl carbonium ions. The order of stability of carbocation is

Triphenyl methyl >Diphenyl methyl>

 $Benzyl > Allyl > 3^{\circ} > 2^{\circ} > 1^{\circ} > methyl carbocation.$

308 **(c)**

Follow IUPAC rules.

310 **(a)**

The dispersal of the charge stabilises the carbocation. More the number of alkyl groups; the greater the dispersal of positive charge and therefore, more the stability of carbocation, thus $C_2H_5^+ > CH_3^+, O - CH_3$ is also an electron donating group, thus it will increase the stability of carbocation, hence, the correct order of stability is C>B>A.

311 (c)

Chromatography method is used to separate sugars.

312 **(c)**

Only urea does not sublime while naphthalene, camphor and benzoic acid do

313 **(c)**

Esters are named by prefixing the name of the alkyl or aryl group (of OR' part) before the name of the parent acid and changing the suffix ic acid to ate. Hence, the structure of phenyl ehtanoate is

314 **(b)**

Inductive effect of groups is measured with respect to H.

315 **(d)**

Methyl halides are methylating agents.

316 **(a)**

Nucleophiles are the species which have excess of electrons. Among the given species, the lone pair of nitrogen of pyrrole is involved in delocalisation of the ring, thus, are not available for donation. In aniline, the lone pair is involved in conjugation with the π -electrons of the ring while in pyridine, these are relatively free for donation. Thus, nitrogen of pyridine is most nucleophilic.



(phenyl and $- \text{COCH}_3$ both are electron withdrawing groups, thus decreases the nuleophilicity of nitrogen).

317 **(c)**

Chlorobenzene is o, p directing in electrophilic substitution reaction. The directing influence is explained by +M of Cl atom



+ M effect of chlorine atom

318 **(b)**

Free radicals are electrons deficient compounds. Alkyl groups are electron donor groups and they increase the stability of free radicals.

 \therefore More the number of alkyl groups, more will be stability of free radicals.

 \div 3°>2°>1° is the correct order of stability of free

radicals.

319 **(b)**

This give rise to net resultant of four C - Cl vectors equal to zero.

320 **(b)**

IUPAC name of compound.



Straight chain which contains large number of side chains taken as parent chain and counting starts from that side where the side chain is nearest.

3-methyl-5(1-methyl ethyl) octane.

321 **(a)**

The isomerism which arises due to rotation about a C-C is called conformational isomerism and the isomers are called conformational isomers or rotational isomers or conformers.

322 (a)

 $CH_3CH = CH_2CHOHCH_3$ has one asymmetric carbon.

323 **(b)**

(i) In nucleophilic substitution reaction more powerful nucleophile replaces weaker nucleophile.

(ii) In rearrangement reaction atoms replace their position within molecule.

(iii) In elimination reaction small molecules (e. g. , H_2O , NH_3) are lost.

$$R - CH_2CH_2Cl + KOH (alc.) \xrightarrow{\Delta} RCH$$
$$= CH_2 + KCl + H_2O$$

 \therefore KCl and H₂Omolecules are lost during reaction.

 \therefore It is an elimination reaction.

324 **(c)**

CH₃COCH₃ is simplest ketone.

$$CH_3 - C - CH_3$$

OH

is the formula of tertiary butyl alcohol as in it – OH group is attached to tertiary carbon.

327 **(b)**

The neopentane:

CH₃

 $CH_3 - C - CH_3$

IUPAC name:2,2-dimethyl propane

328 **(d)**

Triple bond possesses maximum bond energy.

329 (d)

If molecule having asymmetric carbon atom and is not superimoposable on its mirror image then it is chiral while if it is superimposable on its mirror image, it is achiral.

330 **(a)**

(a) When optically active acid reacts with racemic mixture of an alcohol, it forms two types of isomeric esters. In each, the configuration of the chiral centre of acid will remain the same. So, the mixture will be optically active.

331 **(d)**

Due to H^- shift from C_2 to C_3 . Driving force is conjugation from oxygen. Also bulky gps hinders in hydride shift.

332 **(a)**

 C_6H_6 has six delocalized π -electrons.

333 (c)

Due to same molecular formula.

334 (d)

Cis – *trans* isomers generally contain double bonded carbon atoms.

335 **(b)**

Due to the presence of lone pair on N atom.

336 **(c)**

A molecule having doubly bonded carbon atoms shows geometrical isomerism only if both the doubly bonded carbon have altogether different group, *i. e.*, $_{ba}C \equiv C_{ab}$ or $_{ab}C = C_{ac}$ or $_{dc}C = C_{ab}$.

337 **(d)** I>III>II>IV

 $\rightarrow +M$ effect,

hyperconjugation (6Hs) and +*I* effect (2 Me-groups)

H effect, hyperconjugation (3Hs) and +I effect (1-Me group)

338 **(b)**



known as hyperconjugation.



362 **(c)**

 sp^3 -hybridization with one position occupied by lone pair like NH₃.

364 (a)

2-hexyne gives trans-2-hexene on treatment Li/NH₃

$$CH_{3} - CH_{2} - CH_{2} - C \equiv C - CH_{3} + H_{2} \frac{LL/NH_{3}}{H}$$

$$H$$

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

$$H$$

$$H$$

$$trans-2-hexene$$

365 **(b)**

Lassaigne's test is given by those nitrogenous compounds in which carbon is also present along with nitrogen.

In NH₂. NH₂. HCl, carbon is absent, so it does not given Lassaigne's test.

366 **(b)**

$$CH_{3}CH_{2}-\overset{O}{C}_{1}-\overset{O}{C}_{2}| \xrightarrow{Heterolytic fission} CH_{3}CH_{2}-\overset{O}{C}^{+}+\overset{O}{C}I_{3}$$

 $C I_3$ is most stable carbanion because of -I effect of I which disperses negative charge on \overline{C} . Center



The above compound is an ether and its name is

written as alkoxy alkane. Oxy is attached with the lower group. Hence, the IUPAC name of above compound is 2-ethoxy propane.

368 **(b)**

In thin layer chromatography the relative adsorption of each component of the mixture is expressed in terms of retention factor (R_f) distance moved by the spot centre from

$$R_f = \frac{\text{base line}}{\text{distance moved by the solvent from}}$$
the base line

 $(CH_3)_3CBr + H_2O \rightarrow (CH_3)_3C - OH + HBr$ Br is subsituted by $-OH^-$ (nucleophile) S_N1 (unimolecular nuclerophilic substitution reaction)

370 **(c)**

 $C_6H_5CHC_6H_5$ is the most stable since the +ve charge can be delocalized on both phenyl rings 371 (a)

2-methyl propane-2-ol is *tert*-butyl alcohol. CH₃

$$H_3C - C - CH_3$$

Since, in the above structures, position of Cl is different, these are position isomers, which is a type of structural isomerism.

374 **(b)**

More directionally concentrated orbitals show more overlapping. Also more closer are shells to the nucleus more is overlapping.

375 **(b)**

Removal of H from ketone gives resonance stabilized carbanion.

376 **(d)**

 $CH_3CH_2X \rightarrow CH_2 = CH_2(sp^3 \text{ to } sp^2);$ $CH_2 = CHX \rightarrow CH \equiv CH(sp^2 \text{ to } sp);$

$$CH_2XCH_2CH_2X \rightarrow \Delta$$
 (No change).

377 **(b)**

Follow IUPAC rules.

378 **(c)**

It is a fact.

379 **(d)**

 $S_N 1$ (Unimolecular nucleophilic substitution reactions) Rate∝ (substrate) Rate determining step in the formation of carbocation depends on the stability of carbocation formed. The stability of carbocations follow the order $(C_6H_5)_2 C^+(CH_3) > (C_6H_5)_2CH > (CH_3)_2CH > CH_3CH_2^+$ \therefore Order of S_N 1 reacticity is $(C_6H_5)_2C(CH_3)Br > (C_6H_5)_2CHBr > (CH_3)_3CBr$ $> (CH_3)_2 CHBr > C_2 H_5 Br$ *i.e.*, *iii>ii>iv>v*

380 (c)

 $CH_3CH_2CH_3 \xrightarrow{\text{Hetrolysis}} \overline{CH}_3 + CH_3 \overset{+}{CH}_2, CH_3 \overset{+}{CH}_2 \text{ is more stable than } CH_3 \xrightarrow{--} \overline{CH}_2$

due to dispersal of +ve charge on ethylium ion on account of +ve inductive effect. Thus,

propane will not give $\stackrel{+}{C}H_3$ and CH_3 $\stackrel{-}{C}H_2$.

381 (d)

Delocalised electrons are present in benzene, 1, 3butadiene and 1,3,5-hexatriene

382 (a)

A characteristic of *dextrorotatory*.

383 (c)

It is a fact.

384 (c)

If a liquid decomposes at or below its boiling point, it is purified by vacuum distillation, impure glycerine is purified by this method

386 (a)

Distillation process is not used for purification of solid impurities. It is used for the purification of liquids which boils without decomposition and contains non-volatile impurities.

387 (b)

 sp^3 , sp^2 and sp

– orbitals are at 109°28′, 120° and 180°.

388 (b)

 \bigcirc 2°H is more reactive than 1°.

389 (b)

Dehydration of alcohol involves the loss of two atoms or groups from the adjacent carbon atoms, hence it is an example of β -elimination reaction.

B CH₃CH₂OH ethanol

391 (d)

(i) Eantiomers are pair of optical isomers which are related as non-superimposable mirror images of each other.

(ii) Diasteriomers are pair of optical isomers which cannot be related as non-superimposable mirror images of each other.

: The only correct statement about given structures is that (*A*) and (*B*) are enantiomers.

392 (a)

Two positive charges present at the adjacent place, elevates the energy, thus lowers the stability most.

393 (d)

Due to H-bonding.

394 (a)

Follow inductive effect.

395 (b)

Naphthalene and benzoic acid cannot be separated by the sublimation method because the naphthalene and benzoic acid both are sublimes on heating. They are separated by hot water in which benzoic acid dissolves but naphthalene does not

396 (d)

All are used as dehydrohalogenating agent.

$$C_{2}H_{5}X \xrightarrow{(i) \text{ KOH alc.}} C_{2}H_{4} + HX.$$

or (ii) C₂H₅ONa
or (iii) NaNH₂

397 (c)

In the given electrophile

Group is the same. So, only *X* affects their activity, *i. e.*, we have to discuss activity due to

(a)
$$- OCH_3$$
 (b) $- Cl$

(c) N < MeMe (d) $-S - CH_3$

Since, amines are less actives, therefore, electrophile (c) will be least active.

398 **(c)**

The positions of Cl are different.

399 **(a)**

Higher the stability of carbocation, faster is the reaction because $S_N 1$ reactions involve the formation of carbocation intermediate.



reduction are used to convert CO to CH_2 .

The later is not suitable as it will also attack – OH group of ring.

401 (d)

In the following carbocation; H/CH_3 that is most likely to migrate to the positively charged carbon is



(A resonance stabilised carbocation)

402 **(b)**

Hyperconjugation arises due to the partial overlap of a $sp^3 - s$ (a C-H bond) with the empty p –orbital of an adjacent positively charged carbon atom.



Hyperconjugation in ethyl cation

403 (c)

In the triphenyl methyl carbonium ion the π electrons of all the three benzene rings are delocalised with the vacant *p*-orbital of central carbon atom. So, it is resonance stabilised. It is the most stable of all the carbonium ions given

CH₃ is stabilised by hyperconjugattion, a

second order resonance.

404 (c)

During nitration of benzene the attacking electrophile is NO_2^+ . It is formed as follows by reaction between HNO₃ and H₂SO₄.

(I)
$$HNO_3 + H_2SO_4 \longrightarrow H \longrightarrow O + HSO_4$$

(II) $H \longrightarrow O + NO_2 \longrightarrow H_2O + NO_2^+$
(II) $H \longrightarrow H + H_2O + NO_2^+$

405 (b)

The R and S enantiomers of an optically active compound differ in their optical rotation of plane polarised light.

406 (a)

н-

A carbon atom which is attached by four different group is called chiral centre of asymmetric carbon atom. (+)-glucose has four chiral centres.



(ii)
$$CH_3 - CH - CH_2 - NH_2$$

 CH_3
(iii) $CH_3 - C - CH_3$

$$|$$

$$NH_{2}$$

$$(iv) CH_{3} - CH_{2} - CH - NH_{2}$$

$$|$$

$$CH_{2}$$

411 (c)

Stereoisomerism is of two types-optical and geometrical.

412 (a)

Catenation is the tendency to unite atoms of an element to form a long carbon chain or ring.

414 (b)

Electron withdrawing group has – I effect while electron donating group has +I effect. In CH_3COOH , the alkyl group $(-CH_3)$ due to its greater +I effect increases the electron density on oxygen atom of the O-H bond. Due to this the release of H⁺ ion in acetic acid will be more difficult as compared to formic acid.

415 (b)

It is a fact.

416 **(b)**

If positive charge is present on nitrogen then positive charge will not be in conjugation to the ring because in this case nitrogen will become pentavalent

$$\overset{\textcircled{\baselineskip}{\baselineskip} \mathsf{NH}_3 \qquad \qquad \mathsf{NH}_3 \overset{\baselineskip}{\baselineskip} \mathsf{Pentavalent} \\ & & & & \\ & & & & & \\ & & & & \\ &$$

418 (c)

HNO₃ is added to decompose Na₂S and NaCN if present

 $Na_2S + 2HNO_3 \rightarrow 2NaNO_3 + H_2S \uparrow$ $NaCN + HNO_3 \rightarrow NaNO_3 + HCN \uparrow$

419 (c)

It is a fact.

420 (c)

The definition of electromeric effect.

СН•СООН Maleic acid is

It has no asymmetric carbon, however it shows geometrical isomerism.

422 (a)

Sbstituion of – OH group from a substrate can be easily made by PCl₅.

423 (c)



Chair form is unsymmetrical due to absence of any element of symmetry.

424 (a)

When two compounds have similar molecular formula but differ in the functional group then the isomerism is called functional group isomerism *i.e.*,

0|| CH₃CH₂CHO and CH₃. CCH₃

426 (d)

Count σ -and π -bonds on each carbon and report hybridization.

427 (d)

Carbonyl compounds show nucleophilic addition.

428 (a)

SMA

Twelve in all

(a) Six geometrical isomers



distillation because p —nitrophenol has higher boiling point than o-nitrophenol due to Hbonding.

430 **(b)**

Electromeric effect implies complete transfer of π electrons in presence of a reagent. Since, simple ethers do not contain a multiple bond, therefore, they do not show electromeric effect

431 **(b)**

It is definition of electromeric effect.

432 **(a)**

Mass of silver salt=0.4 g Mass of silver =0.26 g Eq. mass of silver salt/Eq. mass of Ag= $\frac{\text{wt.of silver salt}}{\text{wt.of silver}}$ Eq. mass of silver salt = $\frac{108 \times 0.4}{0.26}$ = 166

Eq. mass of side side side $-\frac{1}{0.26} = 1$ Eq. mass of acid = 166 - 108 = 58

Eq. mass of actu = 100 - 108

433 **(a)**

The $S_N 2$ mechanism always involves 100% inversion since nucleophile attacks from back side of leaving group.

434 **(a)**

Follow IUPAC rules.

435 (d)

 $\begin{array}{c} 0 \\ \parallel \\ R - C - X + Nu^{-} \rightarrow R - C - Nu^{-} + X^{-} \\ \end{array}$

Best leaving group (poorest nucleophile) is Cl^{\oplus} , thus fastest reaction is with Cl.

436 **(c)**

Note that propyl (propan-l-ol) and isopropyl alcohol (propan-2-ol) are position isomers.

437 (d)

Empirical formula of acid = CH_2O_2 We know that molecular formula = n (empirical formula)

If n = 1 molecular formula = $(CH_2O_2)_1 = CH_2O_2$ If n = 2 molecular formula = $(CH_2O_2)_2 = C_2H_4O_4$

Thus, the probable molecular formula =
$$(CH_2O_2)_3 = C_3H_6C_2$$

438 **(c)**

Excited state of carbon is $2s^1 2p^3$.

439 **(a)**

C = 10.5 g =
$$\frac{10.5}{12}$$
 mol = 0.87 mol
H = 1 g = $\frac{1}{1}$ mol = 1 mol
∴ (C_{0.87}H₁)₇ = C_{6.09}H₇ ≈ C₆H₇

 $pV = nRT = \frac{w}{m}RT$ $1 \times 1 = \frac{2.4}{m} \times 0.082 \times 400$ m = 79Hence, the hydrocarbon is C_6H_7 440 (c) There are four stereoisomers cis - Rcis - Strans – R trans – S 441 (c) IUPAC name is N-methyl methanamine. 442 (d) Oil and water are immiscible liquids thus are separated by using separating funnel 443 (a) Can be solved on the basis of hyperconjugative structures 444 (d) Total number of optical isomers= $(2)^n$ (where *n*=number of asymmetric carbon atom). $=(2)^{2}=4$ Out of these four optical isomers two are meso structures which are optically in active. . Only two structures d and l- tartaric acid are optically active. соон COOH HO-Ċ—ОН H-·ОН HO--C-ĊOOH ĊOOH *l*-tartaric acid *d*-tartaric acid 446 (d) Three coordinate bond on O atom. 447 (c) Furan is heterocyclic and aromatic due to Huckel's rule of aromaticity, *i.e.*, 6π -electrons. 448 (d) In mixture of o-nitrophenol, p-nitrophenol, onitrophenol is steam volatile due to intra molecular hydrogen bonding whereas p-

nitrophenol is less volatile due to inter molecular hydrogen bonding.



449 (c)

The total number of cyclic isomers are six as shown below.



451 (d)

Electromeric effect involves complete transfer of π -electron pair to more electronegative atom on the need of attacking reagent.

452 (c)

Cannizzaro reaction involves oxidation as well as reduction of aldehydes having lack of $\alpha - H$ atom. The mechanism of this reaction is as





(ii) Transfer of hydride ion





C: H: Br = $\frac{2.27}{12}$: $\frac{5.69}{1}$: $\frac{65.04}{80}$ = 2.43: 5.69: 0.813 = 3: 7: 1

or empirical formula = C_3H_7Br

454 **(b)**

The terminal e' of ene is retained if suffix name starts with consonant.



IUPAC name=3, 3-dimethyl-1-butene. 456 (b)

CaO is added to NaOH to retard activity of NaOH, otherwise decarboxylation of acids will occur more violently.

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

457 **(b)**

$$\begin{array}{ccc} CH_3 & CH_3 \\ | & | \\ CH_3 - C - Cl + OH^- \rightarrow HO - C - CH_3 + Cl^- \\ | & | \\ CH_3 & CH_3 \end{array}$$

Rate
$$\propto$$
 [*t*-butyl chloride]

Tertiary butyl carbocation is first formed which is more stable

458 **(c)**

In heterolysis, the covalent bond is broken in such a way that one species (less electronegative) is deprived if its own electron, while the other species gain both the electrons

$$CH_3CH_2CH_3 \longrightarrow \overset{\Theta}{C}H_3 + \overset{\Theta}{C}_2H_5$$

459 **(c)**

Both have different functional groups, i.e., -CN and - NC.

460 **(b)**

A mixture of 50-50% of d and its l form is called racemic mixture.

461 (d)

Sublimation is the process employed for those solids which convert directly into vapours on heating without converting into liquid phase 462 (c)

It is the definition of optical activity.

463 **(a)**

Stability of alkyl carbanion \propto

465 466 468 469	I = I = I = I = I = I = I = I = I = I =	470 471 472 473	example BeCl ₂ , BH ₃ , ZnCl ₂ , AlCl ₃ (d) Planar hexagon conformer has considerable angle strain due to the fact that its bonds are not 109.5°. It also has torsional strain. Due to presence of these strains planar hexagon conformer of cyclohexane is least stable. (a) Both have same molecular formula. (a) Due to resonance partial double bond character is created on vinyl chloride. So, chlorine atom is not replaced easily $CH_2 = CH = CI = CH = CI$ esonance hybrid (a) Glycerol is a trihydric alcohol. It is $CH_2OH = CH = CH$ $CHOH = CH_2OH$. It contains three hydroxyl group. It is present in nature in oils and fats as its carboxylic esters.
469	Thus, the overall order is: (a) III>IV>II>I. (a) All neutral covalent compound in which central atom has incomplete octet are electrophile. For		
474	(d) Once the carbocation is formed as an intermediate attach it in addition of Br ⁻ . $CH_2 \xrightarrow{+\delta} CH_2 + Br \xrightarrow{+\delta} CH_2 \xrightarrow{+} CH_2 \xrightarrow{OH^-} CH_2 \xrightarrow{OH^-} P$	e, the roduc	nucleophile Cl ⁻ and OH ⁻ present in solution also
475	(a) It is arised on the need of attacking reagent, <i>e</i> .g.,		attracted towards electron rich position in benzene ring. Hence, electron donating groups

example BeCl₂, BH₃, ZnCl₂, AlCl₃

$$C = C \left\langle \underbrace{A.R}_{A.R} \right\rangle^{+}_{C} - \overline{C} \left\langle \right\rangle$$

1

478 (b)

An organic reaction intermediate, neutral species having divalent carbon atom with six valence electron out of which two are present in same orbital with opposite spin is called singlet carbene.

479 (a)

Nitration or aromatic compounds takes place by an electrophile. The electrophile will be more

will be easily nitrated.

Toluene will be most easily nitrated among these compounds due to presence of electron donating group $(i.e., CH_3)$.

Nitrobenzene will be most slowly nitrated due to the presence of electron withdrawing group $(i.e., NO_2).$

CH₃NO₂ will be formed by free radical substitution of CH₄

480 **(b)**

Optical isomerism is shown by an asymmetric

carbon atom which has a carbon atom attached to four different atoms or groups.

H
$$H_3C - CH_2 - C - CH_3$$

|
OH
butanol-2

481 **(c)**

When sodium or potassium alkoxide is heated with an alkyl halide to give ether, this reaction is known as Williamson's synthesis.

 $RONa + R'X \rightarrow R - O - R' + NaX$ This is an example of nucleophilic substitution

and follow $S_N 2$ mechanism.

482 (d)

These are characteristics of free radicals.

483 (a)

Addition of Br_2 on ethane follow electrophilic addition

$$CH_2 = CH_2 + Br_2 \longrightarrow CH_2 - CH_2$$

Intermediate is cyclic bromonium ion

484 (d)

Metamers of ethyl propionate are as $CH_3COOC_3H_7$, $C_3H_7COOCH_3$

- 485 **(c)**
- Each carbon has two σ -and two π -bonds. 486 **(b)**

It should be 4-ethyl-3-methyl heptane.

487 (c)

Due to asymmetric carbon atom in it.

488 **(b)**

$$\overset{\beta}{\underset{\text{CH}_3\text{CH}-\text{CH}_2\text{CH}_3}{\overset{\beta}{\underset{\text{CH}_3\text{CH}-\text{CH}_2\text{CH}_3}}} \xrightarrow{\beta} \overset{\beta}{\underset{\text{CH}_3\text{CH}=\text{CH}-\text{CH}_3}{\overset{\text{major}}{\underset{\text{minor}}{\overset{\beta}{\underset{\text{CH}_2\text{CH}_2\text{CH}_3}}}} \xrightarrow{\text{CH}_3\text{CH}=\text{CH}-\text{CH}_3$$

This reaction is governed by Saytzeff's rule. According to this rule the elimination of β hydrogen atom take place from the carbon having the lesser number of H-atoms or in other words a stable alkene is formed. (More substituted alkene is more stable)

489 (a)

Due to same functional group.

490 (a)

On chlorination of 2-methyl butane

496 **(d)**

CH₃-CH-CH-COOH has two asymmetric carbon atoms and OH CH₃



The amount of enolic form is highest in acetyl due to the stabilisation of enolic form by hydrogen

bonding $CH_{3} \xrightarrow{O} CH_{2} \xrightarrow{C} CH_{3} \xrightarrow{O} CH_{3}$

molecule has no symmetry. Thus, number of optical isomers $= 2^n = 2^2 = 4$.

497 **(c)**

Due to restricted rotation about double bond, the alkene shows geometrical isomerism because the relative position of atoms or groups attached to the carbon atoms of the double bond get fixed. If same groups or atoms attached with double bond bearing carbon, then alkene doesn't show geometrical isomerism.



498 **(a)**

Due to +I effect of CH_3 in toulene, it is more reactive than bezene. Due to electron withdrawing nature of – COOH group in benzoic acid and – NO₂ group in nitrobenzene, both benzoic acid and nitrobenzene are less reactive than benzene.

499 **(b)**

These are isopentane, neopentane and *n*-pentane. 500 **(d)**

(a) Ethylene dichloride and ethylidine chloride both react with alc. KOH to produce ethyne. $H_3C - CHCl_2 + 2KOH(alc.) \rightarrow$

$$IC \equiv CH + 2KCl + 2H_2O | ^5$$

 $ClCH_2 - CH_2Cl + 2KOH alc. \rightarrow$

 $HC \equiv CH + 2KCl + 2H_2O$ (b) They are position isomers because they have

same molecular formula but different position of chlorine atom.

H Cl Cl
H
$$H_3C - C - Cl$$
 and $H - C - C - H$
Cl H H

ethylidene dichloride ethylene dichloride (c) : Their molecular formula is same \therefore They have same percentage of chlorine. (d) They give different product on hydrolysis $CH_3CHCl_2 + 2KOH(aq) \rightarrow$ ethyledene chloride

$$\begin{array}{c} 0 \\ \parallel \\ \mathrm{CH}_3 - \mathrm{CH}(\mathrm{OH})_2 \xrightarrow{-\mathrm{H}_2 \mathrm{O}} \mathrm{CH}_3 - \mathrm{CH} \\ \mathrm{Unstable} \\ \mathrm{CH}_2 \mathrm{Cl} - \mathrm{CH}_2 \mathrm{Cl} + 2\mathrm{KOH} \ (\mathrm{aq}) \longrightarrow \mathrm{CH}_2 - \mathrm{CH}_2 \end{array}$$

ethylene dichloride



501 (a)

First the longest continuous chain of carbon atoms is selected. Now numbered the chain from the side containing senior functional group (*i. e.*, the group placed above in the seniority table).

$$H_{3}C \xrightarrow{C} C \xrightarrow{C} C H_{2} \xrightarrow{C} C H_{2} \xrightarrow{C} C H \xrightarrow{C} C H_{3}$$

502 **(d)**

$$CH_3 - CH - CH - CH_1$$

$$2$$
-methyl butan-2-ol

н

503 (d)
$$H_{3}C - C - CH_{3}$$

t- butyl iodide

504 **(b)**

Number of hybrid orbitals = number of σ bonds+number of *lps*

In
$$\operatorname{CH}_3 \begin{pmatrix} H \\ H \\ H \end{pmatrix}$$
; σ -bonds = 3
 $lps = 3$

∴Number of hybrid orbitals =3+0=3Hence, hybridisation is sp^2 and geometry is planar.

506 **(b)**

Follow the concept of hyperconjugation.

507 **(c)**

On the basis of stability of carbocation formed.

508 **(a)**

φ is phenyl group, the question can be solved on the basis of number of conjugative structures509 (a)

 S_N^2 order : methyl > 1° > 2° > 3°.

Activating groups like – OCH_3 , –OH etc activates the benzene ring towards electrophilic substitution while deactivating groups like– NO_2 , –COOH etc. deactivates the benzene ring towards electrophilic substitution. Thus, order of reaction towards electrophile (of the given compounds) is as I>II>III.

511 (a)

Electron deficient species or electron acceptor is electrophile. For example

 CH_3 , CH_2 , CX_2

513 **(b)**

As the min mol wt. must have at least one S-atom so

$$S \% = \frac{\text{wt. of one } S - \text{atom}}{\text{min. mol. wt}} \times 100$$
$$4 = \frac{32}{\text{min mol. wt}} \times 100$$
Min mol wt. $\frac{32 \times 100}{4} = 800$

514 **(b)**

A carbonium ion is sp^2 -hybridized.

515 (a)

It is a fact.

517 (c)

Ortho and para directing groups facilitate the ring for electrophilic substitution reaction. – NH_2 group increase electron density in ring, hece activite it is to electrophilic substitutipon.

518 (d)

n-pentanol, 2-pentanol, 3-pentanol, 2methylbutanol, 2-methylbutan-2-ol, 3methylbutanol, 2, 2-dimethypropanol, and 3methylbutan-2-ol (8 isomers)

519 (a)

Substances which sublimes on heating is usually purified by sublimation. Hence, naphthalene is purified by sublimation.

520 **(d)**

There are total 6α – H to sp^2 carbon and they all can participate in hyperconjugation.



533 (b)

Resonance in benzene gives rise to identical C— C bond lengths.

534 (a)

During nitration benzene ring is attacked by NO_2^+ and hydrogen of benzene ring is replaced by NO_2 group.

 \therefore Nitration of benzene is electrophilic substitution because NO_2^+ is an electrophile.

535 **(b)**

 $\begin{array}{c} CH_{3}(CH_{2})_{5} \\ H_{3}C \end{array} \xrightarrow[H]{} C -Br \xrightarrow[optical]{} OH^{-} \\ H_{3}C \xrightarrow[H]{} OH^{-} \\ H_{3}C \xrightarrow[optical]{} HO - C \xrightarrow[H]{} (CH_{2})_{5}CH_{3} \\ H_{3}C \xrightarrow[h]{} CH_{3} \\ H_{3}C \xrightarrow[h]{} OH^{-} \\ OH^{-}$

In this reaction inversion takes place. Hence, it is an example of $S_N 2$ reaction. In this mechanism the attack of OH⁻ ions take place from the back side while the Br⁻ ion leaves from the front side

536 **(b)**

Organic compound containing nitrogen is fused with a small piece of sodium metal to form NaCN. Na + C + N \rightarrow NaCN

from organic compound

537 (d)

The order of nucleophilicity depends upon the nature of alkyl group '*R*' on which a nucleophile to attack as well as on nature of solvent. However, if these are same, then weaker is acid, stronger is base, *i.e.*, stronger is nucleophilicity. This acidic character is.

 $HI > HBr > HCl > HCN > H_2O > EtOH$

540 **(d)**

Only two unsymmetrical ether is possible from the formula $C_4H_{10}O$ viz

 $CH_3CH_2CH_2 - 0 - CH_3$

methoxy propane

H₃C

2-methoxy propane

541 (c)

 S_N 1 mechanism gives rise to 50% inversion as it involves front seat as well as back seat substitution. This leads to racemic products.

542 (c)

Canonical structures proposed in resonance are not the real structure of compound. The compound showing resonance has a definite structure which can however be not drawn on paper.

543 **(d)**

All the names are correct. Options (d) is IUPAC name.

544 **(b)**

On adding SCN⁻ to an aqueous solution of $Fe(NO_3)_3$, a blood red colour, due to formation of $[Fe(H_2O)_5(SCN]^{2+}$ complex is obtained. This test is used for the detection of Fe^{3+} ion. $SCN^- + Fe(NO_3)_3 + 5H_2O \rightarrow$

 $[Fe(OH_2)_5(SCN)]^{2+} + 3NO_3^{-}$

Blood red colour

545 (d)

In case of alkyl carbocations as the number of *R* group decreases stability decreases. Thus, the correct order of stability of carbocation is

$$R_3C^+ > R_2\overset{+}{C}H > R\overset{+}{C}H_2 > \overset{+}{C}H_3$$

546 **(b)**

Ortho nitrophenol is the most acidic because electron withdraeing group increases acidic character due to -I effect of NO₂

o - nitrophenol

548 (a)

549 **(b)**

 $C_6H_5C \equiv N$ and $C_6H_5N \equiv C$ are functional isomers. 551 (c)

Second ring is in conjugation with lone pair of oxygen



% of N
$$\frac{28}{22400} \times \frac{\text{volume of N}_2 \text{at NTP}}{\text{wt. of compound}} \times 100$$

= $\frac{28}{22400} \times \frac{224}{1.18} \times 100$
= $\frac{28}{1.18} = 23.728$
(d)

In (II) and (IV) lone pair is involved in resonance. 554 **(d)**

A primary carbon is one which is joined to 1 carbon atom. A secondary carbon atom is joined to two carbon atoms and the tertiary carbon is attached to three carbon atoms. A quaternary 570 (b) carbon has all its four valencies attached to carbon atoms.

555 (a)

The compounds which contain active methylene group at the adjacent position of carboxyl group show tautormerism.

This compound does not contain active methylene group, hence does not exhibit tautomerism. Moreover, this compound is highly stable due to extensive cross-conjugation.

557 (b)

CH₃CH₂OH and CH₃OCH₃ have different functional groups.

(ie., -OH in a alcohol and -O-in ether), hence they are the example of functional isomerism.

558 (a)

Two pairs of *cis* and *trans* forms.

559 **(b)**

Stability of alkyl carbocations can be explained by inductive effect and hyperconjugation. According to these two effect the stability order is

$$R \stackrel{\textcircled{\begin{subarray}{c}} \bullet \\ \mathsf{C} \\ \mathsf{C} \\ \mathsf{C} \\ \mathsf{R} \\ \mathsf{R}$$

560 (d)

In the given compound four π - electrons of double bond and 1 lone pair on N atom leads to delocalization of six electrons

562 (a)

Due to resonance, extra stability in 3° carbocation.

564 **(b)**

Follow IUPAC rules.

565 **(b)**

Na reacts with water less violently than K and Rb.

566 (a)

 $Na_2S + Na_2[Fe(CN)_5NO] \rightarrow Na_4[Fe(CN)_5NOS]$ Sod. extract pink/violet colour

567 **(b)**

Heterolytic bond fission produces +ve and -ve ions.

568 (b)

Racemic mixture is formed by mixing two chiral compounds.

569 (c)

Distillation is applied if organic liquid is stable at its boiling point and contains a non-volatile impurity

Two double bonds are treated as different functional group with a triple bond.

571 (b)

Follow priority rule.

572 (c)

Aldehydes and ketones combine with a variety of compounds of the $Z - NH_2$ to form oxime

$$R_2 CO + NH_2 OH \longrightarrow \begin{bmatrix} R_2 C < OH \\ NHOH \end{bmatrix}$$

$$\frac{-H_2O}{R_2C} = NOH$$
oxime

573 (b)

Addition of HCl is not peroxide effect and it occurs via electrophilic addition.

 $(CH_3)_3C$ CH_2OH is neo -pentyl alcohol. 575 (d)

-do 576 (d)

A lot of plant kingdom is made up of cellulose. 579 (b)

The different arrangement of atoms in space that results from the carbon-carbon single bond free rotation by 360° are called conformations or conformational isomers and this phenomenon is called conformational isomerism.

580 (a)

The molecule, which is optically active, has chiral centre, is expected to rotate the plane of polarised light.







Two chiral centres, but plane of symmetry within $molecule \Rightarrow optically inactive$

583 (b)

Note the fact.

585 (d)

Alkanes do not show resonance.

586 (d) and a Br⁻ ion Both CHCl₃ and CCl₄ burn with smoky flame $HBr \rightarrow H^+ + Br^-$ Thus, H⁺attack the π bond of propene to form although both are aliphatic. C₆H₅CH₂OH, aromatic but burns with non smoky flame. These are carbonium ion as exceptions. $\begin{array}{ccc} CH_{3}-CH & H^{+} \\ \parallel \\ CH_{2} \end{array} \xrightarrow{} CH_{3}-CH_{-}CH_{3} \\ 2^{o} \text{ (more stable)} \\ -CH_{3}-CH_{2}-CH_{2} \end{array}$ 587 **(b)** BF_3 is an electron deficient compound. 590 (a) Isomers of propionic acid are as 0 0 $CH_3 - \stackrel{+}{CH} + Br^- \longrightarrow CH_3 - CH_ | \qquad | \qquad | \qquad | \qquad CH_3 \qquad CH_3$ CH₃ $CH_3 - C - OCH_3 H - C - OC_2H_5$ (major product methyl ethanoate ethyl formate 591 (c) 599 (c) Follow IUPAC rules. Heterolytic bond fission give rise to formation of 600 (d) ions. Report prefixes in alphabetic order. 592 (a) 601 (c) This can be judged by comparing the stabilities of Sulphur is present in the sodium extract in the carboxylate ions formed. The most stable form of sodium sulphide (Na₂S) carboxylate ion is formed by strongest acid FeCl₃ gives blood red colour with sodium extract 593 (c) contain N and S CH_3NH_2 and CH_3OH are nucleophiles, $CH_3 - Cl$ is 602 (a) an electrophile. But $CH_3 - C = N^{\circ}$ is a The name of the compound (2Z,4Z)-2, 4-hexadiene. nucleophile due to the presence of a lone pair of 603 (c) electrons on N and is an electrophile due to the It is a fact. presence of a partial positive charge on C 604 (a) 594 (a) The number of stereoisomers=2' No bond around chiral carbon is broken and so (Here, n =chiral carbon atom) configuration will be retained. Thus, number of stereoisomers= 2^3 =8 595 (b) 605 (a) Due to resonance in benzene. The structural formula of the compound 5-nitro-596 (c) 3-methoxy-3-methyl hexanoyl chloride is as Kjeldahl's method is used for estimation of nitrogen 597 (a) $\begin{array}{c|c} 6 & 5 \\ CH_3 & -CH & -CH_2 & -CH_2 & -CH_2 \\ \hline \end{array} \begin{array}{c} 6 & 5 \\ CH_2 & -CH_2 & -CH_2 \\ \hline \end{array} \begin{array}{c} CH_2 & -CH_2 \\ \end{array} \begin{array}{c} CH_2 & -CH_2 \\ \end{array} \end{array} \begin{array}{c} CH_2 & -CH_2 \\ \end{array} \begin{array}{c} CH_2 \\ \end{array} \begin{array}{c} CH_2 \\ \end{array} \begin{array}{c} CH_2 & -CH_2 \\ \end{array} \end{array} \begin{array}{c} CH_2 \\ \end{array} \end{array}$ \end{array} Free radicals stability Free radicals stability $C_6H_5 > C_6H_5$ — CH **ÓCH**₃ 606 (d) Presence of methyl group on NH₃ molecule Highly stable by delocalisation increases the tendency of N atom to lose electron pair. However, tertiary, amines are less basic due >H₃C $\stackrel{\bullet}{\underset{CH_3}{\longrightarrow}}$ CH₃ > H₃CC $\stackrel{\bullet}{\underset{CH_3}{\longrightarrow}}$ CH₃ to steric hindrance. 607 (a) $CHCl = CHCH_2CH_2CH = CH_2;$ 9-hyperconjugative $CH_2 = CClCH_2CH_2CH = CH_2;$ hydrogens and +I effect $CH_2 = CHCHCH_2CH = CH_2$. 598 (b) HBr being better source of proton. It gives a H⁺ Ċl

608 **(c)**

2° carbocation is more stable.

609 **(b)**

In such cases where migrating group is cycloalkyl group, ring expansion may occur.





Chiral molecules should not contain any kind of symmetry.

It is honour to Lavoisier.

Stability of carbanions increase with increasing in *s*-character of hybrid orbitals of carbon bearing charge therefore, the order is $sp^3 < sp^2 < sp$



E-3-ethyl-4-methyl hex-3-en-1-oic acid. [The configuration of this compound is *E* because bulkier groups are present at opposite of the double bond.]

618 **(a)**

-do -

619 **(c)**

The substrate has three different types of B - H, therefore, first, three structural isomers of alkenes are expected as



The last two alkenes II and III are also a capable of showing geometrical isomerism hence two geometrical isomers for each of them will be counted giving a total of five isomers.

620 **(c)**

 CH_2Cl_2 has sp^3 -hybridization and tetrahedral nature.

621 (a)

Chain initiation step involves formation of free 631 (c) radicals only.

622 **(d)**

Follow characteristics of $S_N 2$ mechanism.

623 **(a)**

Follow elimination of HBr from two ends.

624 **(d)**

 $^{\mbox{$\Theta$}}\mbox{CH}_2\mbox{CHO}$ is the most stable carbanion since it is stabilize by resonance

$$0 \qquad 0^{-}$$

$$|| \qquad |$$

$$CH_2 - C - H \leftrightarrow CH_2 = C - H$$

625 **(b)**

X-crown-Y, 18-crown-6

First number X is the total number of 'C' and 'O' atoms in the ring and second number Y is the number of oxygen atom in ring.

626 **(b)**

When the nucleophilic site is the same, nucleophilicity parallels basicity. It means more basic the nucleophile, stronger is the nucleophile.



(vi)
$$H_3C \longrightarrow O \longrightarrow CH \begin{pmatrix} CH_3 \\ CH_3 \end{pmatrix}$$

methyl isopropyl ether
 $O \begin{pmatrix} CH_3 \\ CH_3 \longrightarrow CH_2 \longrightarrow CH_3 \end{pmatrix}$
(vii) methyl arcmed ether

methyl propyl ether

Hence, three isomeric ethers are possible. 632 (c)

> Liebig's method is used to estimate carbon and hydrogen.

C and H
$$\xrightarrow{[0]} CO_2 + H_2O$$

%C = $\frac{12}{44} \times \frac{\text{weight of } CO_2}{\text{weight of compound}} \times 100$
%H = $\frac{2}{18} \times \frac{\text{weight of } H_2O}{\text{weight of compound}} \times 100$

633 (d)

There are six isomers possible for the compounds having molecular formula C₄H₈O, which are as follows



carbonium ion, the π -electrons of all the three benzene rings are delocalised with the vacant *p*orbital of central carbon atom. So, it is resonance stabilised. Therefore, it is most stable of the given carbonium ions.

More the number of resonatic structures more will be the stability.

635 (d)

Chlorine atoms are strongly electro negative (show negative inductive effect i.e., -I effect). They deactivate the ring towards electrophilic reaction.

The increasing order of substituent-E towards electrophilic substitution is

 $-\mathrm{CCl}_3 < -\mathrm{CHCl}_2 < -\mathrm{CH}_2\mathrm{Cl} < -\mathrm{CH}_3$

636 **(b)**

Fructose has three chiral carbon atoms, hence the number of optical isomerism $= 2^3 = 8$ CH₂OH

ĊHOH

CH₂OH

637 (d)

O has two unpaired electrons in ground state.

638 (d)

An organic ion with a pair of available electrons and a negative charge on the central carbon atom is called a carbanion.

Electron attracting group (----CN , >c==-o) increases stability and electron releasing group $(-CH_3 etc)$ decreases the stability of carbanion. In $(CH_3)_3C^-$, three – CH_3 groups (electron releasing group) are present, so it least stable.

639 (d)

The stability order of conformation of cyclohexane is chair>twist boat>boat>half chair. Hence, half chair is less stable due to torsional and angle strain.

640 (a)

It is a fact.

642 (a)

Formation of 2° carbocation, i.e.,

	\sim	$+\mathcal{S}$	-8		
CH ₂ CH=	'⊨CH-	<u>H-</u>	<u>-Н</u>	CH.	HCH.
Chigon	OII_2	2		Chige	meng

643 **(d)**

In CH_3CH_2OH , there is intermolecular H-bonding, while it is absent in isomeric ether CH_3OCH_3

- 1. Larger heat is required to vaporise CH_3CH_2OH as compared to CH_3, OCH_3 , thus (a) is incorrect.
- CH₃CH₂OH is less volatile than CH₃OCH₃, thus vapour pressures are different, thus
 (b) is incorrect.
- 3. Boiling point of $CH_3CH_2OH > CH_3OCH_3$, thus (c) is incorrect.

Density = $\frac{\text{mass}}{\text{volume}}$, due to ideal behaviour at a given temperature and pressure volume and molar mass are same.

Hence, they have same vapour density.

644 **(a)**

Follow IUPAC rules.

646 **(b)**

% of chlorine =
$$\frac{35.5}{143.5} \times \frac{\text{mass of AgCl}}{\text{mass of the compound}} \times 100$$

= $\frac{35.5}{143.5} \times \frac{0.287}{0.099} \times 100$
= 71.71 %

647 **(c)**

Due to presence of delocalised π -electrons in the aromatic compounds, the electron density is maximum inside the ring. Therefore, aromatic compounds undergo electrophilic substitution reaction and resistance to addition reactions.

648 **(a)**

64

The acidic nature is
$$H_2 O > C_2 H_2 > C_2 H_4 > C_2 H_6$$
.
49 **(b)**



methyl CH_3 , vinyl $CH_2 = CH$ given options can be solved on the basis of conjugative and hyperconjugative structures

650 (a)

Follow IUPAC rules.

652 **(a)**

 $(CH_3)_3C - C(CH_3)_3$ has maximum number of alkyl groups in it.

Halogenation of methane is chain reaction, propagate through free radical

655 **(a)**

Follow IUPAC nomenclature.

658 **(c)**

 \because C – H bond in toluene has partial double bond character due to resonance.

 \div C – H bond in toluene has less energy as

compared to others.

659 **(b)**

The reaction $(CH_3)_3C - Br \xrightarrow{H_2O} (CH_3)_3COH$ is an example of substitution reaction

660 **(b)**

Draw the isomers.

661 **(b)**

Distillation is used to purify liquids and based on difference in their boiling points. When the boiling points of liquids are very closed to each other, then fractional distillation is used.

662 **(c)**

The C-atoms attached to the triple bond lie in a straight line while the carbon of the CH_2 group is inclined at an angle of 120°. Therefore, only 2 carbon atoms are linearly arranged

663 **(c)**

The conversion of $\alpha\mbox{-glucose}$ to $\beta\mbox{-glucose}$ is called mutarotation.

664 **(c)**

Positively charged species in which central atom has incomplete octet is called electrophile, H^+, X^+, R^+ are electrophile

665 **(b)**

An organic reaction intermediate, neutral species having divalent carbon atom with six valence electron out of which two are present in same orbital with opposite spin is called singlet carbene

666 **(a)**

The structure of D-glyceraldehyde is as CHO

н–|-----он

I CH₂OH

The priority of groups is decided by the following rules

(i) Atom having higher atomic number gets higher priority

653 **(d)**

(ii) If the priority cannot be decided by rule 1 then the next atoms are considered for priority assignment.

(ii) Where there is a = bond or=bond both atoms are considered to be duplicated or triplicated

 $(----c^{-})^{\circ}$ has higher priority than – CH_2OH). Hence, the correct order of priority of groups in D-glyceraldehyde is as:

OH(1), CHO(2), CH₂OH (3) and H(4)

668 **(c)**

Nucleophile always attacks on electron deficient site. Presence of electron withdrawing groups such as NO_2 , CHO etc decreases the electron density on benzene nucleus, hence such groups activate the ring towards nucleophilic attack. While presence of electron releasing groups such as R or OR increases the electron density, thus deactivates the nucleus towards nucleophilic attack.

 $\rm NO_2$ group activates the ring more than Cl towards nucleophilic attack,

Hence reacts readily with nucleophile.

NO₂ 669 (d)

CI

The increasing order is :

$$-CH_3 < CH_3 - CH_2 - < (CH_3)_2 CH - < (CH_3)_3 C -$$

670 (a)

The treatment with FeCl_3 yield ferric ferrocyanide which has bulish green or prussian blue colouration

 $3Na_4[Fe(CN)_6] + 4FeCl_3 \rightarrow Fe_4[Fe(CN)_6]_3$ 671 (c)

Each π -bond contributes two π electrons and the two electrons of the lone pair are present in a *p*-orbital. Therefore, total number of π electrons is six

672 **(a)**

Allyl carbocations are more stable than the alkyl carbocations due to the resonance stabilization

673 **(b)**

The middle carbon has 2σ -and 2π -bonds.

675 **(c)**

Di-chloro acetic acid due to presence of two electron with drawing chloro groups (-I showing group) is more acidic than acetic acid $(+I \text{ showing} - \text{CH}_3\text{group})$.

677 **(b)**

Na₃PO₄ + 3HNO₃ → H₃PO₄ + 3NaNO₃ sod. extract H₃PO₄ + 12(NH₄)₂MoO₄ + 21HNO₃ → (NH₄)₃PO₄. 12MoO₃ + 21NH₄NO₃ + 12H₂O yellow

ppt.

678 **(b)** 1,2,3; 1,2,4 and 1,3,5-trimethyl benzene.

679 **(b)**

Hydride ion is formed when hydrogen accept a proton, so it has a tendency to donate electron. Since, hydride ion (H^-) has a tendency to donate electron, it functions as nucleophile.

The structure of isomers from $C_2H_2Br_2$ are $CH_2 = CBr_2$;

$$CH_2 = CBr;$$

 I
 H
 Br
 $C = C < Br$
 H
 Br

cis-isomer II

$$H c = c < H H$$

trans-isomer III

681 **(a)**

$$CH_{3}MgX + CH_{3}C$$

$$\equiv C - H \rightarrow CH_{4} + CH_{3}C$$

$$\equiv C MgX.$$

682 **(a)**

The correct decreasing order of priority for the functional groups of organic compounds in the IUPAC system is

$$\label{eq:coord} \begin{split} -\text{COOH} &> S\text{O}_3\text{H} > -\text{COOR} > \text{COCl} > -\text{CONH}_2 \\ &> -\text{CN} > -\text{CH} = 0 \end{split}$$

683 **(b)**

Optical isomerism is shown by compounds which have one or more chiral carbon atoms.

HC
$$\stackrel{H}{=} C \stackrel{H}{\stackrel{}{=} C} C \stackrel{H}{\stackrel{}{=} CH_3}$$

 \therefore It has asymmetric or chiral carbon atom.

∴ It shows optical isomerism.

684 **(d)**

 $S_N 2\ reaction$ does not involve ion formation, these infact involve formation of transition state

685 **(a)**

 pK_a value of carboxylic group is less than pK_a of

 $^{+}_{\mathrm{NH}_{3}}$ in amino acid and $^{--}_{\mathrm{NH}_{3}}$ (Z) will have

comparatively less pK_a than $-NH_3^+$ (Y) due to -I effect of carboxylic group. We know that acidic strength in inversely proportional to pK_a . Hence, correct order of acidic strength is

$$--- \operatorname{CO}_{2} \operatorname{H}(X) > ---- \operatorname{NH}_{3}(Z) > ---- \operatorname{NH}_{3}(Y)$$

686 **(a)**

Chromatography is a modern technique used for the separation of mixtures into its components, purification of compounds and also to test the purity of compounds.

688 **(d)**

Kharasch effect involves addition of HBr.

689 **(d)**

These are characteristics of carbocations.

691 **(b)**

Symmetrical *trans*-form has non-polar nature. 694 **(c)**

The nucleophilic addition reaction is the characteristic addition of carbonyl compounds. Reactivity order of carbonyl compounds is in the order.

$$H \rightarrow C = 0 > H_3C \rightarrow C = 0 > H_3C \rightarrow C = 0$$

This is due to increase in the intensity of charge on carbon of carbonyl group due to +I effect of alkyl groups.

695 (c)

Chlorobenzene has only one deactivating group, . *e.*, -Cl. In 2, 4-dinitrochlorobenzene three deactivating group, *i. e.*, two – NO₂ and one – Cl are present and *p* –nitrochlorobenzene two deactivation groups, *i. e.*, one NO₂ and one Cl is present. So, the order of reactivity is A>C>B.

Acid has —COOH group whereas, ester has — COOR group.

697 **(d)**

The given reaction can be represented as $NaOH \rightarrow Na^+ + OH^-$



Since in this reaction, a nucleophile replaces the other group, it is a example of nucleophilic substitution reaction.

The mechanism shows that the rate depends on the concentration of both alkyl halide and nucleophile. So, it is an example of $S_N 2$ (nucleophilic substitution of II order) reaction.

698 (a)

Follow IUPAC rules.

699 **(d)**

Follow IUPAC rules.

702 (a)

Carbonyl compounds undergoes nucleophilic addition reaction.

$$X \xrightarrow{\mathbf{C}}_{\mathbf{S}^+}^{\mathbf{S}^-} [X-\text{shows negative inductive effect}]$$

If group or atom attached with carbonyl carbon shows negative inductive effect, then it decreases electron density on carbonyl carbon and facilitate the attack of nucleophile, hence reactivity of carbonyl compound increases. The aromatic aldehydes and ketones are less reactive than their aliphatic analogues due to +R effect of benzene ring. The increasing order of the nucleophilic addition reaction in the following compounds will be.

 $\label{eq:CH3} CH_3 CHO > CH_3 COCH_3 > PhCOCH_3 > PhCOPh$ 703 (b)

Chiral compounds which have one chiral centre. All four atoms or groups attached to carbon are

696 **(a)**

different.

704 **(b)**

Due to the presence of methyl group positive inductive effect increases and the stability of carbocation also increases. The stability order of carbocation is

Tertiary > Secondary > Primary

705 **(b)**

According to IUPAC system ethers are named as alkoxy alkanes. The larger alkyl group forms the parent chain while lower alkyl group is taken with the ethereal oxygen and forms a part of alkoxy group.

CH₃—O
$$\xrightarrow{2}_{CH}$$
 CH₂ $\xrightarrow{3}_{CH_2}$ CH₃
|
1CH₃
2-methoxybutane

706 **(c)**

 H_3^+ O cannot accept electron pair.

707 (a)

Epoxide is ambident substrate for nucleophilic substitution reactions. In protonated epoxide carbon-2 and carbon-3 both aquire some positive charge due to the highly electronegative atom

$$MeO^- > MeCH_2O^- >$$

708 **(d)**

Beilstein test is used to detect halogens in organic compounds.

 $> Me_2 CHO^- > Me_3 CO^-$

709 **(b)**

Free radical chain reaction is initiated by UV light. It proceeds in three main steps like initiation, propagation and termination. It gives major products derived from most stable free radical

711 **(b)**

Follow IUPAC rules.

713 **(d)**

Elimination reactions involves removal of a molecule (HBr here) from a substrate.

and



0

Ш



Fumaric acid

OH



$$\begin{array}{l} \mathrm{CH}_{2}\mathrm{CH}_{2}-\mathrm{CH}_{2}-\mathrm{C}-\mathrm{CH}_{3}\rightleftharpoons\mathrm{CH}_{3}\mathrm{CH}_{2}\mathrm{CH}_{2}-\mathrm{C}\\ &=\mathrm{CH}_{2} \end{array}$$

ketoform

enol form

715 **(b)**

The species which are electron deficient and accept a pair of electron are called electrophile. Hence, SO₃ is a electrophile as it contains an electron deficient centre. While H_2O , NH_3 and R - O - R are nucleophiles.

In the above reaction more stable carbocation is generated hence, the compound dehydrated very easily

719 (a)

In case of kjeldahl's method the percentage of $\rm N_2$ is then calculated from the amount of $\rm NH_3$

721 **(b)**

Such dehydrohalogenation follow E_2 mechanism. The driving force of such reaction is the stability of alkene produced. Since, tertiary alkyl halide can give more substituted alkene, it reacts fastest followed by secondary and primary *i.e.*, $3^\circ > 2^\circ > 1^\circ$.

722 **(c)**

Central carbon atom is chiral carbon.

723 **(d)**

Those organic compounds, which are volatile in steam are purified by steam distillation. Since, aniline is a steam volatile compound, hence it is purified by steam distillation. 725 **(b)**

 $C_6H_5O^-$ possess less nucleophilicity due to stabilized nature of phenoxide ion. CH_3OH is weaker acid than CH_3COOH and thus CH_3O^- is stronger base.

Acidic order : $CH_3COOH > H_2O > CH_3OH$

726 **(c)**

Vinyl chloride is least reactive for $S_{\ensuremath{N}}$ reaction due to resonance

$$CH_2 = CH - CI \leftrightarrow CH_2 - CH = CI$$

729 **(b)**

The chain propagation step involves the use of free radical and regeneration of another free radical.

730 **(b)**

CH₃NC is methaneisonitrile.

732 (a)

Benzyl carbonium is more stable due to resonance and thus, benzyl chloride is more reactive.

733 (a)

It is a fact.

734 **(b)**

 $C_n H_{2n} O_2$ is general formula for open chain acid and ester.

 $n = 3 \quad C_3 H_6 O_2$ 0 $\|$ Acid CH₃CH₂ - C - O - H

U II _ 4

Easter $CH_3 - C - 0 - CH_3$

735 **(a)**

The Cannizzaro reaction is as HCHO + HCHO $\xrightarrow{\text{KOH(conc.)}}$ CH₃OH

 $CHO + HCHO \xrightarrow{\text{Reflection}} CH_3OH + HCOOK^+$ methyl alcohol acetic acid

The mechanism of Cannizzaro reaction is as **Step I** Attack of nucleophile OH⁻ to the carbonyl carbon



Step II The transfer of hydride ion from anion (I) to second molecule of aldehyde and finally rapid transfer of proton takes place.



 $(NH_4)_2 SO_4.$

742 (c)

Acetone and methanol have nearly equal boiling point. thus, they are separated by fractional distillation

743 (b)

Follow IUPAC rules.

744 (c)

The oxygen atom in phenol has more dominating 745 **(c)**

The acid exist in *cis* and *trans* forms :



Also it has asymmetric carbon atom $CH_3CH = HCOOH$.

746 (a)

Follow the mechanism of esterification.

747 (b)

Pyridine is a heterocyclic compound having six

membered ring formed with C and N-atoms.

748 (a)

When – OH group of lactic acid is replaced by H, then chiral carbon is lost.

$$CH_{3} - C - COOH \rightarrow CH_{3} - C - COOI$$

Η

lactic acid

∴Its optical activity is lost.

750 **(a)**

752 (a)

Ozonolysis of the compound may be given as : $CH_3 H$



753 **(b)**

2-aminopentane and 3-aminopentane; Position is different.

755 **(c)**

resonance effect than inductive effect. Increase in charge separation decreases the stability of a resonating structure

Stability of resonating structure in decreasing order will be $I > II \equiv IV > III$

2-butene exhibit rotamers. Rotamers are the isomers formed by restricted rotation.751 (d)

It contains lone pair electron on N atom.

$$=\frac{2}{18} \times \frac{0.9}{0.5} \times 100 = 20\%$$

 $\times 100$

% of H = $\frac{2}{18}$

weight of H₂O weight of organic compound o – and p-directing groups facilitate S_E reactions whereas m-directing groups deactivate benzene ring for S_E reactions.

757 **(a)**

(+) and (-) tartaric acid does not possess any element of symmetry.

758 **(b)**

A molecule having doubly bonded carbon atoms shows geometrical isomerism only if both the doubly bonded carbon have altogether different group, *i. e.*, $_{ba}C \equiv C_{ab}$ or $_{ab}C = C_{ac}$ or $_{dc}C = C_{ab}$.

759 **(c)**

The chemical formula of thiourea is NH_2CSNH_2 so here Na_2S , NaCN and NaCNS will be formed but not Na_2SO_4

760 **(a)**

A white precipitate with am. AgNO₃ confirms the presence of terminal alkyne.

761 **(a)**

Racemisation involves change in entropy, *i.e.*, change in arrangement of groups position leading to a change in entropy of disorderness.

762 **(a)**

The acyclic stereoisomers of $\rm C_4H_7Cl$ are



Number of optical isomers $=2^n = 2^1 = 2$ Hence, total number of geometrical isomers=6 Total number of optical isomers =2.

763 **(c)**



bicyclo (4, 1, 0) heptane

This compound contains 7 carbon atoms, so the corresponding alkane is heptane. Two bridges contain 4 and 1 carbon atom respectively and one bridge does not contain any carbon atom. So the name of the compound is bicyclo (4,1,0) heptane.

764 **(d)**

Stability of alkyl free radicals can be explained by hyperconjugation and number of resonating structure due to the hyperconjugation. The decreasing order of stability of alkyl free radical is as follows

 3° free radical > 2° free radical > 1° free radical > $\overset{\bullet}{C}H_3$

766 **(b)**

Inductive effect involves only displacement (and not delocalisation) of σ –electrons.

767 (c)

Meso forms are optically inactive as they are superimposable to their mirror images.

768 **(b)**

 CH_4 has highest ratio of H to C

769 **(b)**

active methylene group



When methylene group $(-CH_2)$ is attached with two electron withdrawing groups (like, -CHO, > C = O, -COOH, -CN, -X, etc), its acidity will increase due to -I effect of the electron withdrawing groups.

770 (a)

Follow IUPAC rules.

771 **(c)**

The reactivity order for H atom is $3^{\circ} > 2^{\circ} > 1^{\circ}$; Neocarbon does not have H atom.

773 **(b)**

— do—

774 **(a)**

Organic compound which are volatile in steam can be purified by steam distillation. It is based on the fact that vaporisation of organic liquid takes place at lower temperature than its boiling point

775 **(b)**

Follow IUPAC rules.

777 **(d)**

 $CH_3\bar{O}$ is nucleophile;

 $CH_3OH+ Na \longrightarrow CH_3O Na + (1/2)H_2$

778 **(a)**

Inductive effect is the permanent effect on σ –electrons. It involve the electron displacement along the chain of saturated carbon atoms due to the presence of a polar covalent bond at one end of the chain.

779 **(c)**

Homologous differ by a group — CH_2 and cannot be isomer.

780 **(c)**

The reagent selected should be such that only one of components to be separated, reacts with it. Aniline+aq. HCl \rightarrow salt, which is water soluble Nitrobenzene +aq. HCl \rightarrow no reaction

 \therefore aq. HCl is used to separate aniline and nitrobenzene.

781 **(b)**

Formic acid was obtained from ant (fromica in greek). This is trivial name for HCOOH.

782 **(a)**

The structure of 2, 3-dimethyl hexane is



So, the number of tertiary carbon atoms=2 The number of secondary carbon atoms=2 The number of primary carbon atoms=4

783 **(a)**

Follow IUPAC rules.

786 (c)

CH₃⁺ has planar structure. 787 (**d**)

These are characteristics of carbanion.

788 **(a)**

Follow Saytzeff rule for elimination. 3-halopentane will give only pentene-2.

789 **(b)**

Atom Ratio	Atomic	Percentage	$\frac{b}{a} = x$	
	Mass (a)	(b)		
С	12	10.06	$\frac{10.06}{12}$	1
Н	1	0.84	$\frac{0.84}{1}$	1
Cl	35.5	89.10	$\frac{89.10}{35.5}$	3

Empirical formula = $CHCl_3$

Empirical formula mass = 12 + 1 + 106.5 =119.5 \approx 120

Molecular mass = $2 \times V.D = 2 \times 60 = 120$ molar mass

 $n = \frac{120}{\text{empirical formula mass}}$ $= \frac{120}{120} = 1$

Molecular formula = $(CHCl_3)_1 = CHCl_3$

790 **(d)**

During nucleophilic substitution weaker nucleophile is replaced by stronger nucleophile. The compound having C-Cl bond which can be most easily broken will be most reactive towards nuclophilic substitution reaction. In vinyl chloride $CH_2 = CH - Cl$ and chlorobenzene C_6H_5Cl the C - Cl bond has partial double bond character due to resonance. ∴ They do not give nucleophilic substitution reaction easily

Benzyl chloride, give nucleophilic substitution easily because they carbocation formed is stabilised due to resonance.

$$\xrightarrow{\text{OH}^{-}} \text{HOCH}_2 \xrightarrow{} \text{CH} \xrightarrow{} \text{CH}_2$$
allyl alcohol

791 (a)

Enantiomers are non-superimposable mirror images, *e.g.* lactic acid



Diastereomers are non-superimposable and are not the mirror images of each other. Moreover, *meso* form has plane of symmetry.

792 **(b)**

Nucleophilic strength increases down a column of the Periodic Table (in solvents that can have hydrogen bonds, such as water, alcohols, thio alcohols).

Nucleophilic strength $RO^- < RS^-$

Base strength $RO^{\ominus} > RS^{-}$

Thus, RO^{\ominus} is more nucleophilic but less basic than RO^{-}

795 (a)

We know that there are seven isomers in $C_4H_{10}O$. Out of these seven isomers, four are of alcohol and three are of ether.

796 **(a)**

Tertiary halide always favours $S_N 1$ mechanism (as they give comparatively stabler carbocation) white primary halide favours $S_N 2$ mechanism.

797 **(d)**

Electron donors are bases. Since, electron density is highest at



798 **(d)**

Follow IUPAC rules.

800 (c)

To be optically active, compound or structure should posses a chiral or asymmetric carbon atom. 1-chloropentane is not chiral.

801 **(c)**

Stearic hinderance in tertiary halides give rise to less reactivity for $S_N 2$.

802 **(b)**

Addition of Br_2 gives altogether different products units *cis* and *trans* butene-2.

803 **(d)**



It has one chiral centre (two enantiomer) and two geometrical isomers

cis–d, trans–d, cis– and trans–l.

804 **(d)**

Glucose contains four chiral carbon atoms hence number of possible optical isomers are $2^4 = 16$.

805 **(c)**

Markownikoff's rule is for addition of unsymmetrical additive on unsymmetrical alkene.

806 **(d)**

Presence of halogen in organic compound can be detected by Beilstein's test.

807 **(c)**

The bond energy of catenation order is C > Si > S > P.

808 **(d)**

$$\begin{array}{l} R - X \xrightarrow{\text{NaOH}} R - \text{OH} + \text{Na}X \\ R - X \xrightarrow{\text{OH}^{-}} R - \text{OH} + X^{-}. \end{array}$$
 This is nucleophilic substitution.

809 **(c)**

2-methyl butanoic acid exhibits stereo isomerism. $$\rm CH_3$$

It shows optical isomerism because it contains asymmetric carbon atom.

810 **(a)**

$-CH_3$ is electron repelling group.		CH ₂ B	Sr 	СН	
812 (c)		This is	ne albronna	e acetylene	action
metamerism is found in molecules having	826	(d)	a dellydroll	alogenation re	
813 (d)	020	Stereo	oisomerism i	s of two types	geometrical and
There are HIPAC rules		ontica].	s of two types,	geometrical and
	827	(c)			
$a = 2^{n}$: where n is no. of dissimilar asymmetric		Follow	v IUPAC rule	s.	
carbon atoms and a is no. of optically active	828	(c)			
isomers.		Compo	ounds having	g asymmetric (2-atom is
816 (a)		optical	lly active, e. g	g.,	
Follow IUPAC rules.			ŀ	┥ ┃ ▲	
817 (d)			*		•
Nucleophile $(-NH_3)$ replaces other nucleophile		Н ₃ С—	—H ₂ С—С	;СНО 	
(-Br) in the reaction.			r		
818 (a)		The C-	ztom whose	four valencies	are satisfied by
819 (d)		four d	ifferent grou	ns is asymmet	ric C-atom.
Formation of ethylene from acetylene is an	829	(b)	increne grou		
example of addition reaction		Chlori	ne of vinyl cl	hloride ($CH_2 =$	CHCl) is non-
CH CH ₂		reactiv	ve (less react	tive) towards r	ucleophile in
$H_{1} \rightarrow H_{2} \rightarrow H_{3}$		nucleo	ophilic substi	itution reactior	n because it
CH CH_{2}		shows	the followin	ng resonating s	tructure due to
Ethyne ethene	K,	<i>+M</i> ef	fect of – Cl at	com.	
820 (b)		Си —		Б- СН₂—СН=	δ+ =Cl
Sodium hydrogen sulphite adds to aldehydes and		CH ₂ —	I	II	
ketones to form crystalline bisulphite addition		In stru	icture II, Cl-a	tom have posi	tive charge and
products. The product is water soluble and can be		partial	l double bon	d character wi	th C of vinyl
converted back to the original carbonyl		group,	, so it is more	e tightly attract	ed towards the
compound by treating it with dilute mineral acid		nucleu	is and it does	s not get replac	ed by
or alkall. Therefore, these are useful for		nucleo	pphile in S_N^-	reaction.	
acetaldehydes	830	(d)	h :	6 .] .]	L'
821 (b)	021	Follow	mechanism	l ol debromina	uon.
Zn dust is used for dehalogenation,	051	(C) Atom	A b c c c b c	D	b
CH V CH V Zn dust CH - CH		Atom	Atomic	Percentage	$\frac{-}{a} = x$
$CH_2A.CH_2A$ alc. CH_2 CH_2 .		Ratio	Maga (g)	(b)	
823 (d)		C	$\max(u)$	(b)	40 2.22
Resonance in a molecule is arised due to		L	12	40	$\frac{1}{12} = 3.33$
delocalisation of π -electrons.		1			6 66
824 (d)		Н	1	6.66	$\frac{0.00}{1} = 6.66$
CH ₃ -CH ₂ -CH=CH		2			
sp^3 sp^3 sp^2 sp		0	16	53.34	$\frac{53.34}{16} = 3.33$
Electronegativity of different hybrid and unhybrid		1			
orbitals in decreasing order is as follows		Hence	, empirical fo	ormula = CH_2C)
$s > sp > sp^2 > sp^3 > p$	832	(d)			_
825 (b)		Grigna	ard reagents	s can act as	electrophile and
Δ	0.00	nucleo	ophile.		
$ + 2KOH \longrightarrow + 2KBr + 2H_2O$	833	(0)			

Both these carbon atoms have 3σ -and 1π -bond. Recall hybridized orbitals never from π -bonds.

834 **(c)**

 $S_N 1$ mechanism involves the formation of carbocation intermediate. Hence, the species which gives the most stable carbocation readily undergoes $S_N 1$ mechanism. *t*-butyl bromide gives the most stable carbocation, *i. e.*, 3° carbocation, so it readily undergoes $S_N 1$ reaction.

835 (b)

Follow IUPAC rules.

836 **(b)**

In the Lassaigne's test, a blue colour is obtained if the organic compound contains nitrogen. The blue colour is due to ferri-ferro cyanide *i.e.*, $Fe_4[Fe(CN)_6]_3$.

837 (d)

According to Cahn-Ingold-Prelog sequence rules, the priority of groups is decided by the atomic number of their atoms. When the atom (which is directly attached to the asymmetric carbon atom) of a group has higher atomic number, then the group gets higher priority. Groups which atoms of comparable atomic number having double or triple bond, have high priority than those have single bond.

Hence, the order of priority of group is $-OH > -COOH > -CHO > -CH_2OH$

838 (c)



 $\dot{C}H_3$ Here, when C_2 is rotated anticlockwise 120° about $C_2 - C_3$ bond the resulting conformer is *Gauche* conformer.



839 (c)





carbon, thus optically active.



2-bromo 3-chloro butane

∴ Number of asymmetric carbon atoms=2

: Number of chiral isomers = $2^n = 2^2 = 4$

841 (c)

Glycerol can be separated from spent lye in soap industry by the distillation under reduced pressure because it decomposes near its boiling point

843 **(b)**

In gas phase tertiary amines are more basic than secondary amines which are more basic than ammonia

-I group present on central atom decreases electron density, hence decreases basicity $CH_3NH_2 > NH_3 > NF_3$

844 **(a)**

Atom	At mass (a)	% (b)	$\frac{b}{a}$
Ratio			-
С	12	49.3	$\frac{49.3}{12} = 4.10$
2			
Н	1	6.84	$\frac{6.84}{1} = 6.84$
3			1
0	16	43.86	$\frac{43.86}{16} = 2.74$
1			10
Hence	, empirical fo	rmula = (C	$L_2H_3O)$
Molec	ular mass $= 2$	\times VD = 2 $>$	× 73
= 146			
n —	molar ma	ss	$\frac{146}{2} \sim 3$

 $n = \frac{1}{\text{empirical formula mass}} = \frac{1}{43} \approx 3$

So, formula= $(C_2H_3O)_3 \approx C_6H_9O_3$

845 **(c)**

Wöhler prepared urea from inorganic compounds and rejected the vital force theory that organic compounds can only be synthesised from living organisms.

846 (c)

Follow mechanism of addition of HCl and HI in presence of peroxide. One of the chain
propagation step is endothermic in both cases.

847 **(c)**

All aromatic compounds are resonance hybrid.

- 848 **(a)** It is the stability order for various conformers.
- 849 **(c)**

Glucose has aldehyde group and fructose keto group. The general formula for both is $C_6H_{12}O_6$.

- 851 **(b)** Follow conformation.
- 852 **(b)**

In o-, m-, p – derivatives vectors are at 60°, 120° and 180°. Thus, *para* has zero dipole moment. Also *ortho* form has more dipole moment than *meta* form.

853 **(c)**

The staggered form has lower energy than eclipsed form because of repulsive interaction between the H-atoms attached to two carbon atoms are minimum due to maximum distance between them.

854 **(c)**

Victor Mayer's method is applicable only for the determination of molecular mass of volatile substance

856 (d)

Hexane is non-polar molecule.

857 **(c)**

Nucleophilies may be neutral or negatively charged, whereas substrate undergoing nucleophilic substitution may be neutral or positively charged

 $C_2H_5 - I + OH^- \rightarrow C_2H_5OH + I^-$

858 **(a)**

Nucleophilicity increases on going down in the group of the Periodic Table

 $I^{\Theta} > Br^{\Theta} > Cl^{\Theta} > F^{\Theta}$

859 **(d)**

Free radicals have unpaired electrons, but are neutrals and are reactive.

$$CH_3 + CH_3 \longrightarrow CH_3 \longrightarrow CH_3$$

861 **(b)**

Follow mechanism of Kharasch effect.

862 **(d)**

—do—

864 **(c)**

Reaction of NaOH with dinitrofluorobenzene represents nucleophilic aromatic substitution reaction because – NO₂ group is deactivating

group. They make benzene nucleus electron deficient and facilitate the nucleophile to attack the ring.



Cyano group has the highest priority therefore, parent name must be benzonitrile. Br occurs at 2position, and hydroxyl at 3-position, hence the IUPAC name is 2-bromo-5-hydroxy benzonitrile.

866 **(d)**

Ethers show metamerism.

867 **(b)**

Due to resonance; the carbonyl group of benzoic acid is coplanar with the ring. If the electron withdrawing substituent (i.e., -I showing) is present at *ortho* position, it prevents the coplanarity and thus, the resonance. Hence, makes the acid more stronger.

Thus, among the given acids, *ortho* hydroxy benzene acid is the most acidic.

868 **(a)**

Diamond (sp^3) , Graphite (sp^2) , Acetylene (sp).

869 **(d)**

CH₃CHClCOOH contains asymmetric carbon atom.

870 (d)

Statement (c) is wrong.

871 **(b)**

The –ve inductive effect of –CHO group play role to give anti Markownikoff's addition.

$$CH_2 \longrightarrow CH \rightarrow -CHO \rightarrow CH_2 - \overline{C}H - CHO.$$

The structure of 1-chloro-2-nitroethene is as O_2N_1 , CI



In this compound E-Z isomerism is possible because it is highly substituted alkene. The E-Zsystem of nomenclature is developed by Cahn, Inglod and Prelog.

873 **(a)**

 $CH_3C = N$ is known as acetonitrile or methyl cyanide.

874 **(b)**

Ketone undergoes nucleophilic addition reaction because nucleophilic end of reagent attack first followed by electrophilic end of reagent.



875 **(d)**

Halogen containing compounds (C_6H_5Cl) When placed in a flame, the presence of halogen is revealed by a green to blue flame.

876 **(a)**

Two similar asymmetric carbon atoms; $\therefore a =$

 2^{n-1} . Also meso form $= 2^{\frac{n}{2}-1}$.

Total = a + m.

877 **(d)**

In C_6H_5 ring there are three $\pi\text{-bonds}$ and one $\pi\text{-}$

C = 0

bond is present in OH group. Therefore, in all there are four π - bonds in C_6H_5COOH . In $CH_3CH_2COCH_3$ there is only one π bond in C = 0 group, in $CH_2 = CH - CH =$ CH_2 there are two π -bonds while in $HC \equiv C CH = CH_2$ there are three π - bonds

878 (c)

Order of bond length σ bond $(sp^3) > \sigma$ bond $(sp^2) > \sigma$ bond (sp)

879 **(d)**

It is a reason for the given fact.

880 **(c)**

The octet of all atoms are complete in structures *a* and *b*. The molecule in which all the atoms have completed octet is more stable than atom which have incomplete octet. Larger the number of resonating structures, larger will be the stability, thus structures *a* and *b* are stable. In structure (d), the electron deficient of positive charged carbon is duly compensated by one pair

charged carbon is duly compensated by one pair electrons of adjacent oxygen atoms while such neighbour group support is not available in structure (c). Hence, structure (c) is least stable in comparison to structure (d).

883 (a)

CH₃CH₂Cl; CH₃CHCl₂; CH₂ClCH₂Cl; CH₃CCl₃; CH₂ CHCl₂CCl₃; CCl₃CCl₃

884 **(d)**

(1) and (3) are enantiomeric forms to each other . 885 **(d)**

Methoxy group, due to +I effect, increase electron density on OH- group, thus making it less acidic. Thus, *o*-methoxy phenol and acetylene are less than phenol.

p-nitrophenol is more acidic than phenol.

886 **(c)**

When organic compound is fused with sodium metal, nitrogen of the compound is converted into sodium cyanide as

$$Na + C + N \rightarrow NaCN$$

887 **(c)**

It is structure of furan, a heterocyclic compound. 888 **(a)**

Diazonium salts are highly reactive. In Sandmeyer reaction diazo group is replaced by chlorine or bromine in presence of CuCl or CuBr.(Substitution reaction)

$$\overset{\textcircled{}}{C_6H_5N_2Cl} \overset{\textcircled{}}{\underbrace{CuCl}} \blacktriangleright C_6H_5Cl + N_2$$

 X^{-} is replaced by OH⁻.

890 **(b)**

There are four structural isomers are possible for $\rm C_4H_9Cl$

(a)
$$CH_{3}CH_{2}CH_{2}CH_{2}CH_{2}CI$$

(b) $CH_{3} - CH_{2} - CH - CH_{3}$
(c) $CH_{3} - CH - CH_{2}CI$
(c) $CH_{3} - CH - CH_{2}CI$
(d) $CH_{3} - C - CH_{3}$
(d) $CH_{3} - C - CH_{3}$
(CH₃)

891 **(b)**

A carbanion or carboanion has –ve charge on it. 894 **(c)**

The case with which a nucleophile attacks the carbonyl groups depends upon the electron-deficiency, *i. e*, magnitude of the positive charge on the carbonyl carbon. Since, an alkyl groups has electron-donating inductive effect.

(+*I* effect), therefore, greater the number of alkyl groups attached to the carbonyl groups greater is

the electron-density on the carbonyl carbon and
hence, lower is its reactivity towards nucleophilic9addition reactions.9

$$R \rightarrow C = 0 > R \rightarrow C = 0$$

896 (c)

n-pentane and isopentane or 2-methylbutane are chain isomers since both have different hydrocarbon chain.

897 (a)

 $CH_2 = CH - CHO$

898 **(d)**

Free radicals have unpaired electrons but are neutrals and are reactive.

 $CH_3 + CH_3 \rightarrow CH_3 - CH_3$

900 **(a)**

The second carbon is asymmetric.

901 **(c)**

Both have different mode of linkage, *i.e.*, chain and ring.

902 **(c)**

Carbanions contain even number of valence electrons and thus, show diamagnetic behaviour.

903 **(d)**

Molecules with two similar groups attached on either of the doubly bonded carbon do not show geometrical isomerism.

904 **(c)**

Compound $CH_3 - CHCl - CH_2 - CH_3$ shows optical isomerism due to the presence of chiral carbon atom.

H ______chiral carbon atom

905 (c)

The Kolbe's electrolysis proceeds *via* free radical mechanism. For example, when sodium propionate is electrolysed, n —butane, ethane, ethylene are obtained. The propionate ion discharge at the anode to form free radicals. $C_2H_5COO^- \rightarrow C_2H_5COO^* + e^ C_2H_5COO^* \rightarrow C_2H_5^* + CO_2$ $2C_2H_5^* \rightarrow C_4H_{10}$

$$C_2H_5^{\bullet} + C_2H_5^{\bullet} \rightarrow C_2H_4 + C_2H_6$$

In TLC, adsorbent is made of silica gel or alumina

gel 908 **(d)**



Lead unreached 0.1 M (=0.2 N)H₂SO₄ = V mL 20 Ml of 0.5 NaOH=V mL of 0.2 N H₂SO₄ 20 × 0.5 = V × 0.2 $V = \frac{20 \times 0.5}{0.2} = 50 \text{ mL}$ Used H₂SO₄ = 100 - 50 = 50 mL % of N = $\frac{1.4 \text{ NV}}{w} = \frac{1.4 \times 0.2 \times 50}{0.30}$ = 46.67 % % of nitrogen in (a) CH₃CONH₂ = $\frac{14 \times 100}{59}$ = 23.73 % (b) C₆H₅CONH₂ = $\frac{14 \times 100}{121}$ = 11.57 % (c) NH₂CONH₂ = $\frac{28 \times 100}{60}$ = 46.67 % (d) NH₂CSNH₂ = $\frac{28 \times 100}{76}$ = 36.84 %

910 **(d)**

Removal of H from alkane, alkene and alkyne gives alkyl, alkenyl, alkynyl groups respectively.

911 **(d)**

There are two symmetrical hexenes as given in (a) and (b).



The solution of D(+)-2-chloro-2-phenyl ethane in toluene racemises slowly in the presence of $SbCl_5$ due to the formation in carbocation

914 **(a)**

Tautomerism it is functional isomerism in which

the isomers are readily interchangeable and maintain a dynamic equilibrium with each other.



915 **(b)**

If acid is weak, its conjugate base (nucleophile) is strong and *vice versa*.

(B)
$$CH_3O^-$$
 is a conjugate base of CH_3OH (II)

(C) CN⁻ is a conjugate base of HCN (III)

(D)H₃ H_3C SO_3^- is a conjugate base

Of H₃C SO₃ H(IV)

0

Acidic nature of IV>I>III>II and nucleophilicity of B>C>A>D.

916 **(d)**

 $S_N 2$ reaction proceeds with inversion and a transition state is formed which does not carry any charge

918 **(b)**

The most stable one is that in which the positive and negative charges reside on the most electropositive and most electronegative atoms of the species respectively. Like

÷с: сн₂−сн=с−сн 919 **(b)**

Organic compound $\xrightarrow{[0]} CO_2 + H_2O$

% of
$$C = \frac{12}{44} \times \frac{11.6}{5.6} \times 100 = 85.7\%$$

% of $H = \frac{2}{18} \times \frac{7.2}{5.6} \times 100 = 14.28\%$

Ele	Percen	Relative no.	Simplest ratio
me	tage	of atoms	
nt			
С	85.7	85.7/12=7.1	7.14/7.14=1
		4	
Н	14.28	14.28/1=14.	14.28/7.14=
		28	2

Hence, empirical formula of compound of $= CH_2$ \therefore Molecular formula of compound $=C_4H_8$ 920 **(c)**

0

Stronger is acid, weaker is its conjugate base. Acidic nature : $CH \equiv CH > CH_2 = CH_2 >$ $CH_3 - CH_3$ Conjugate basic : $CH \equiv C^- < CH_2 = CH^- <$ $CH_3 - CH_2^-$ nature. 921 (b) $H \xrightarrow[-]{} CHO$ $H \xrightarrow[-]{} OH$ CH_2OH

D-glyceraldehyde has the above formula. So, the Fischer's projection formula which is identical to

922 **(b)**

Percentage of *N* in an organic Compound = $\frac{1.4 \times N \times V}{W}$

$$=\frac{1.4 \times 1 \times 30}{1.2} = 35$$

923 (c)

Dissociation of proton from ddd is very difficult due to -I effect of Cl⁻and N⁺ while in C₆H₅OH due to the reasonance stabilization of phenoxide ion proton eliminates easily. Similarly due to Hbonding in C₆H₅CH₂OH it can be eliminate easily and in CH₃C \equiv CH the proton is acidic in nature hence, it can be dissociated

% of S =
$$\frac{32}{233} \times \frac{\text{wt. of BaSO}_4}{\text{wt. of organic compound}} \times 100$$

= $\frac{32}{233} \times \frac{1.158}{0.53} \times 100$
= 30%

925 (c)

Tautomerism is a dynamic isomerism because two forms (keto and enol) of substance cannot be separated they are in dynamic equilibrium with each other.

926 **(a)**

Enantiomers are non-superimposable mirror images. *e.g.*, lactic acid



927 (a)

Vinyl carbocations are more stable than primary carbocation but less stable than secondary carbocation.

928 **(b)**

Duma's method involve the determination of nitrogen content in the organic compound in the form of $N_{\rm 2}$

 $N_2O + Cu → N_2 + Cu$ d% of N = $\frac{28}{22400} \times \frac{\text{volume of } N_2 \text{at NTP}}{\text{weight of compound}} \times 100$

930 **(a)**

The correct order of reactivity is

RI > RBr > RCl > RF

It is due to fact, that weaker the base, better it will be the leaving group.

Hence, I^- is the best leaving group.

931 **(b)**

 $R_3 N \longrightarrow CH \longrightarrow CH_2 \xrightarrow{HBr}$

Due to R_3^{\bigoplus} (*e*⁻withdrawing tendency) carbocation will appear farther to that (terminal). Hence, product is $R_3N - CH_2 - CH_2Br$.

932 (d)

n-pentane and 2-methyl butane are constitutional isomers or chain isomers or skeletal isomers.

934 (a)

Follow IUPAC rules.

935 **(b)**

In toluene $(-CH_3)$ group is present which has +I effect and increases electron density on *ortho* and *para* position. While in sulphonation $-SO_3$ acts as electrophile. Therefore, it $(-SO_3H)$ attacks on *ortho* and *para* position readily.

936 **(d)**

Follow elimination rules.

937 (a)



944 **(d)**

It is the definition of distereoisomers.

945 **(b)**

The central carbon is attached to four different substituents, hence it is chiral, therefore optically active.

946 **(b)**

 $HNO_2 + H_2SO_4 \rightarrow NO^+ + HSO_4^- + H_2O$ Nitrosonium ion

947 **(b)**

-Cl is an electron withdrawing (*i.e.*, -I showing) group. It withdraws electrons when attached to the carboxylic acid and decreases the electron density on the oxygen atom. This will facilitate the release of H⁺ by making O – H bond more polar and thus – Cl increases the acidity of acetic acid when attached at, α position because of – *I* effect. 948 (a)

 $-NH_2$ has +R effect, it donates electrons to the benzene ring. As a result, the lone pair of electron on the N-atom gets delocalized over the benzene ring. As a result, the lone pair of electron on the Natom gets delocalized over the benzene ring and thus it is less readily available for protonation. Hence, aniline is a weaker base than cyclohexylamine.



951 (c)

Acetophenone oxime can show geometrical isomerism.



952 (d)

Ethers show functional isomerism with molecular formula $C_n H_{2n+2}O$. For example, $C_2 H_6 O$ CH₃OCH₃, CH₃CH₂OH dimethyl ether, ethyl alcohol

953 **(c)**

The alkyl halide which produce stable carbonium ion undergo $S_N 1$ reaction. Tertiary alkyl halide is most stable. Thus, decreasing order of tendency of alkyl halides undergoes $S_N 1$ mechanism.

t-alkyl halide>*sec*-alkyl halide> *primary*-alkyl halide

954 **(c)**

Vinyl chloride($CH_2 = CHCl$) undergoes addition and elimination reactions. Substitutions reaction is shown by compounds having single bonds only.

955 **(a)**

Fisher projections are for illustration of optical isomers.

956 **(b)**

A hydrogen halide contain a highly polar H —

X bond can easily lose $\stackrel{\frown}{H}$ to the pi bond of an alkene. The result of the attack of H^{\oplus} is an intermediate carbocation, which quickly undergoes reaction with the negative halide ion (X^{-}) to yield an alkyl halide



960 **(c)**

Aldehydes and ketones readily undergo nucleophilic addition reaction. The order of reactivity, is as the +I effect of alkyl group increases

$$\underset{H}{\overset{H}{\rightarrow}}c=0>\underset{H}{\overset{CH_{3}}{\rightarrow}}c=0>\underset{CH_{3}}{\overset{CH_{3}}{\rightarrow}}c=0$$

961 (b)

The structures of maleic and fumaric acids are given below

$$H - C - COOH$$

∥ HOOC − C − H

(fumaric acid)

molecular

H – C – COOH (maleic acid) molecular formula ($C_4H_4O_4$) formula($C_4H_4O_4$)

The structures of fumaric and maleic acid suggest that they are geometrical isomers because they have same molecular formula but different spatial arrangement of atoms around a double bond.

962 **(b)**

Stability order of arenium ion II>III>I

963 **(d)**

In condensed structure formula, paranthesis is used for identical group of atoms.

964 **(d)**

Compound which sublime on heating can be purified by sublimation method.

Benzoic acid, camphor and naphthalene sublime on heating hence, they are purified by sublimation method.

965 **(d)**

 $CH_2 = CH. CH_2Cl$ compound undergoes nucleophilic substitution most readily.

966 **(a)**

As the – *I* group increases at the α -carbon, acidity increases

967 **(b)**

In $S_N 2$ reactions, the nucleophile attaches itself from the direction opposite to that of the nucleophile already present in the second step, the previous nucleophile is removed and a single stereoisomer is obtained

968 **(b)**

Follow IUPAC rules.



Since, the reaction rate depends upon the concentration of both reactant and nucleophile, it is a $S_N 2$ reaction. It involves inversion of configuration.

970 (c)

The given compound is nitrobenzene.

971 **(c)**

Both differs by a $-CH_2$ group.

972 **(d)**

During electrophilic substitution electrophile attacks the double bond of benzene ring. The aromatic compounds having electron donating groups undergo electrophilic substitution more easily due to favouable effect of electron donating group.



(i) NO₂, COOH and CHO groups are electron withdrawing groups so, they decrease the reactivity of organic compounds.

 (ii) - OHgroup is electron donating group, so it increases the electron density in benzene ring ad increases the rate of reaction



973 **(b)**

Oxygen cannot be detected by direct test because oxygen is present in atmosphere and all tests are carried in atmosphere of oxygen

974 **(a)**

Follow IUPAC rules.

975 **(a)**

o-, m-, p- isomers are position isomers.

976 **(b)**

Carbinol is trivial name for HCH_2OH . Thus, $C_6H_5CH_2OH$ is phenyl carbinol and chloral is CCl_3CHO .

977 **(a)**

A primary carbon is one which is attached with one carbon atom.

978 **(a)**

Follow IUPAC rules.

979 **(c)**

1,1-dibromoethane and 1,2-dibromoethane.

980 (a)

A solvent molecule lacking a polar X-H bond is called aprotic solvent. NH_3 , SO_2 and CH_3CN are aprotic solvent while CH_3COOH is protic solvent.

981 **(b)**

When sodium ethoxide reacts with iodoethane, diethyl ether is obtained (**Williamson's synthesis**) The mechanism of this reaction is as follows $C_2H_5ONa \rightleftharpoons C_2H_5O^- + Na^+$

$$C_2H_5O^- + C_2H_5 - I \xrightarrow{\text{Slow}} [C_2H_5 - O^- - C_2H_5 - I]$$

transition state

<u>Fast</u> $C_2H_5OC_2H_5 + I^-$ Since, the reaction involves substitution of a group by a nucleophile, it is an example of nucleophilic substitution reaction.

982 (c)

The organic compounds is fused with Na metal, Na_2S is formed which is tested as

It reacts with lead acetate and forms black ppt. of PbS

 $Na_2S + (CH_3COO)_2Pb \rightarrow PbS + 2CH_3COONa$ black ppt

983 (c)

0 Ш $CH_3 - C - NHBr$ N-bromoethanamide 984 (c) Ε 100 $\frac{108}{108} = \frac{1}{60}$ Eq. wt of the silver salt $E = 108 \times$ = 180 \therefore Eq. wt. of the acid = E 108 + 1 = 73985 (a) 2 1 $C_2H_5 - C - CH_2OH$ 3 || CH₂ 2-ethylprop-2-en-1-ol 986 (d) The structure of butane-2, 3-idol is as H_3C ÓН ÔН : Optical isomers in compounds have similar asymmetric carbon atom, which are even in number = $2^n - 1$ Here, n = 2: Total number of optically active stereoisomers

$$= 2^{2} - 1 = 3$$
987 (b)
CH₄ \rightarrow CH₃⁺ + H⁻;
CH₃⁺ is methyl carbonium.
988 (b)
(A)CH₃ - CH - CH₂CH₃
i
Cl
One asymmetric carbon atom, forms *d*, and
l-optical and isomers.
(b) Two asymmetric carbon atoms, forms *d*, *l* and
meso forms

$$= CH_{3}$$
H + Cl
Plane of
symmetry + cl
Cl
Plane of
symmetry - Cl
CH₃ + CH - CH₂CH₃
Meso due to internal compensation
(c)CH₃ + CH - CH₂CH₃
Cl
Two asymmetric carbon atoms but does not have
symmetry. Hence, *meso* form is not formed.
(c)CH₃ + CH - COOH
One asymmetric carbon atom, *meso* form is not
formed.
989 (d)
More the number of hyperconjugated structures,
more will be electronegative chlorine atom.
(a) CH₃ - Cl
3-hyperconjugated structures
(b) CH₃ - CH₂ - Cl
2-hyperconjugated structures
(c) CH₃ - CH₂ - Cl
(c) CH₃ - CH₂ - Cl

(d)
$$CH_3 - CH_2 - C - Cl$$

CH₃ 8-hyperconjugated structures ∵ 8-hyperconjugated structures are possible for

 CH_3

(d)

 $\div \mbox{Chlorine}$ in this is most electronegative.

990 **(a)**

Follow resonance characteristics.

991 **(b)**

Diastereoisomers are a pair of optical isomers which cannot be related as non-superimposable mirror images of each other.

994 **(a)**

The rate of reaction is influenced by the hyperconjugation effect of group R. it depends on the electron donating power of alkyl group (R). The electron releasing power of R group depends on the number of hydrogen present on α carbon. The increasing order of speed with R group in the reaction is

 $H_{3}C \xrightarrow{CH_{3}} H_{3}C \xrightarrow{H_{3}C} CH \xrightarrow{H_{3}C} CH \xrightarrow{CH_{3}} CH_{2} \xrightarrow{CH_{3}} CH_{3}$

i.e., IV<III<II<I

995 **(b)**

The given compound form two geometrical isomers and two optical isomers.

996 **(b)**

Geometrical isomerism is shown by >C=C< only when identical groups are not present on the double bonded carbon atoms.

999 **(b)**

 $RX + I^- \rightarrow R - I + X^-$ This reaction is an example of nucleophilic substitution.

100 (c)

0

1

Its IUPAC name is bicyclo [1,1,0] butane.

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Weak base \rightarrow strong conjugate acid
```

 $CH \equiv C^{-} \rightarrow CH \equiv CH$ weakest base (strongest acid among the given)

$$CH_3 - CH_2^- \rightarrow CH_3CH_3$$

(strongest base) (weakest acid)

- 100 **(d)**
- 2 With the increasing basicity of the added base, the rates of the elimination reactions have been found to increase. Thus, *RO⁻* is most reactive

100 **(b)**

3 Follow IUPAC rules.

100 (a)

Number of isomers in hexane are five as follows $CH_3 - CH_2 - CH_2 - CH_2 - CH_2 - CH_3$ $CH_2 - CH - CH_2 - CH_2 - CH_3$ CH_3 $CH_3 - CH_2 - CH - CH_2 - CH_3$ CH_3 CH₃ CH_3 $CH_3 - C - CH_3$ $CH_3 - C - CH_2 - CH_3$ CH_2 CH3 100 (c) (a) it is Diels Alder's reaction (cyclo addition) 5 =O + NaHSO₃ -(b) It is nucleophilic addition reaction $C - OH + \frac{Dry HCl/Anhy. ZnCl_2}{Lucas reagent} CH_3$ CH₃ (c) It is nucleophilic substitution reaction Br $(CH_3)_2C = CH_2 + BrCl \rightarrow (CH_3)_2 - C - CH_2$ (d) It is electrophilic addition reaction 100 (d) (d) is with maximum conjugative structure 6 among them 100 (a) 7 Based on heat of hydrogenation. 101 (c) Butyne-2 is $CH_3 - C \equiv C - CH_3$; Two carbon of 0 corner are sp^3 -hybridized. 101 (c) 1 Four π -electrons of double bond and 1 lone pair on N atom leads to delocalisation of six electrons. 101 (d) 2 NH₂ $CH_3 - CH - CH_3$ 2-propanamine

101 **(b)**

3	Elem Perce		Percentage	Simple	
	ent	ntage	at.wt.	ratio	

С	20.0	$\frac{20.0}{12} = 1.66$	$\frac{1.66}{1.66} = 1$
Н	6.67	$\frac{6.67}{1} = 6.67$	$\frac{6.67}{1.66} = 4$
N	46.67	$\frac{46.67}{14} = 3.33$	$\frac{3.33}{1.66} = 2$
0	26.66	$\frac{26.66}{16} = 1.66$	$\frac{1.66}{1.66} = 1$

Empirical formula = CH_4N_2O

Empirical formula weight

 $= 12 + (4 \times 1) + (2 \times 14) + 16 = 60$ mol. formula weight

 $\therefore n = \frac{1}{\text{emp. formula weight}}$ $=\frac{60}{60}=1$

: Molecular formula = CH_4N_2O Given compound gives biuret test. Thus, given compound is urea $(NH_2)_2CO$.

101 (c)





Hence, two chiral carbon atoms are present in 2, 3, 4-trichloropentane.

101 **(b)**

The electronegativity of F is maximum and thus, 5 C— F bond is more polar.

101 (a)

1, 1-dichloro-1-pentene does not exhibit 6 geometrical isomerism.

101 **(b)**

solution 102 (a) The 7 hydrolysis 🔺 of sugar of 5 leads to (*dextrorotatory*) formation laevorotatory mixture due to formation of (*dextrorotatory*) glucose and fructose (laevorotatory).

101 (c)

First two and fourth will show stereoisomerism. 8 101 (d)

Geometrical isomers of $CH_3CH = CH - CH_3$ are



102 (c)

0 Ethyl alcohol shows functional isomerism with dimetyl ether.

 C_2H_5OH $CH_3 - O - CH_3$ alcohol ether

- 102 (a)
- Highest to lowest priority $(Br > Cl > CH_3)$ is 1 clockwise than R.

102 (c) 2



Electrophilic substitution reaction takes place in compounds in which π –electrons are highly delocalised. The electrophile attacks the region of high electron density, therefore, electrophilic substitution occurs at ortho/para position at ring II.

102 (c)

3

When phenol reacts with chlrtoform and aqueous NaOH solution, it give salicyladehyde.



102 (c)

4

It is definition of asymmetric synthesis.

Since, the compound on heating with CuO produced CO_2 , it contains carbon. Again, it does not produce water, hence it does not contain hydrogen. So, the organic compound is carbon tetrachloride (CCl_4).

102 (d)

6

8

If organic compound gives blue or green colour at the tip of red hot copper wire, this indicates the presence of halogens in the compound. However, compounds like urea, thiourea etc also give blue or green colour in this test even in the absence of halogens, thus, it is not a sure test for halogens

102 (a)

Mesomeric effect involves complete transfer of π or lone pair of electrons to the adjacent atom or covalent bond. Hence, it involves delocalisation of pi (π) electrons.

103	(b)				3-methyl but-1-yne
0	Species	Valence electrons	Magnetic	104	(d)
	behavior			0	I can have maximum 3 hyperconjugative
	Carbonium ion	6	Diamagnetic		structures. II has maximum 5 hyperconjugative
	Free radical	7			structure, III has 2 conjugative structure while IV
	Paramagnetic				has 1 conjugative structure
	Carbene	6	Diamagnetic	104	(a)
	Nitrane	6	Diamagnetic	1	$CH_3C \equiv CCH_2CH_3$; It is always pentyne-2 and
103	(b)				never pentyne-3.
1	It should conta	in $(CH_3)_2 CH - gro$	up to be named	104	(d)
	as iso.			3	The components of an azeotropic mixture are
103	(b)				separated by special method, <i>ie</i> , fractional
2	One propene ar	nd one cyclopropan	e.		distillation. The simple fraction which distils at
103	(c)				337.8 K is a ternary azeotrope consisting of all
3	Those compour	nds which contain t	wo or more		water. Some alcohol and benzene
	asymmetric car	bon atoms but are o	optically	104	(a)
	inactive due to	presence of plane o	f symmetry, are	4	Hence, an equimolar mixture of the enantiomers
	called <i>meso</i> con	npounds. M <i>eso</i> con	npounds are		(dextro or laevo forms) is called racemic
	optically inactiv	ve due to internal co	ompensation.		mixture. It is represented as dl -form or \pm form
	Out of the giver	n compounds only 2	, 4-		and is optically inactive due to external
	dibromopentan	e have a plane of sy	vmmetry, so it		compensation. Separation of racemic mixture into
	is a <i>meso</i> comp	ound.			<i>d</i> - and <i>l</i> -forms is called resolution.
	C	CH ₃		104	(a)
				5	Two successive homologous differ in their
	н——с	C——Br	4	X	formula by CH ₂ or have a difference of 14 units in
				\bigcirc	their molecular weight.
	н——с	с——н	5	104	(c)
				6	If a liquid decomposes at or below its boiling
	H——Č	C——Br			point, it is purified by vaccum distillation
				104	(c)
	0	CH ₃		7	$CH_3CH_2NH_2$ and CH_3NHCH_3 .
	2, 4-dibromop	entane (<i>meso</i> comp	ound)	104	(a)
103	(a)	$ \lambda $		8	IUPAC name of CH_3CH_2CHO is propan-1-al.
4	It is a fact.	CXY		104	(b)
103	(a)			9	Alkyl group (an electron releasing (+ <i>I group</i>)
6	Electrophiles an	re the species havin	g a tendency to		increases electron density at N-atom, hence, basic
	accept a pair of	electron, $e.g.$, NO ₂	, Br+ etc.		nature is increased. In ammonia, no alkyl group is
	Nucleophiles ar	e the species havin	g a tendency to		present, so it is least basic.
	donate a pair of	f electron. <i>e</i> . <i>g</i> ., CH ₃	$OH.N_3^-$		
103	(a)				
7	C_5H_8 has three	possible alkynes. Tl	hese are		
\mathbf{C}	$CH_3CH_2 - CH_2$	$C \equiv CH, CH_3CH_2C$	$\equiv C - CH_3$		
	pent-1-yne	pent-	2-yne		
	CH ₃				
	I				
	$CH_3 - CH - C$	$\equiv CH$			