AMINES

CHEMISTRY

Single Correct Answer Type

- 1. During diazotization of benzenamine with sodium nitrite and hydrochloric acid, the excess of hydrochloric acid is used primarily to
 - a) Check the hydrolysis of ϕ OH

- b) Ensure a stoichiometric amount of nitrous acid
- c) Check the concentration of free aniline
- d) Neutralize any base formed during reaction
- 2. Hofmann's bromamide reaction is to convert
 - a) Acid to alcohol
- b) Alcohol to acid
- c) Amide to amine
- d) Amine to amide
- 3. Examine the following two structures for the anilinium ion and choose the correct statement from the ones given below

- a) II is not acceptable as canonical structure because carbonium ions are less stable than ammonium ions
- b) II is not an acceptable canonical structure because it is non-aromatic
- c) II is not an acceptable canonical structure because in it N has 10 valence electrons
- d) II is an acceptable as canonical structure
- 4. Choose the amide which on reduction with $LiAIH_4$ yields a secondary amine
 - a) Ethanamide

b) N-methylethanamide

c) N, N-dimethylethanamide

- d) Phenylmethanamide
- 5. When methyl cyanide is hydrolysed in presence of alkali, the product is:
 - a) Acetamide
- b) Methane
- c) $CO_2 + H_2O$
- d) Acetic acid

6. In the following reactions, reactants A, B and C are:

$$Cl_2H_5NH_2 + A \rightarrow C_2H_5N = CH - C_6H_5 + H_2O$$

Urea +B \rightarrow H₂N - NHCONH₂ + NH₃

 $CH_2H_5NH_2 + C \rightarrow C_2H_5Cl + H_2O + N_2$

a) CH₃CHO, NH₂ - NH₂ and PCl₅

b) C₆H₅CHO, NH₂ - NH₂and SOCl₂

c) C₆H₅CHO, NH₂ – NH₂and NOCl

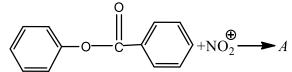
- d) CH₃CHO, NH₂ NH₂and PCl₃
- 7. Toluene is nitrated and the resulting product is reduced with tin and hydrochloric acid. The product so obtained is diazotised and then heated with cuprous bromide. The reaction mixture so formed contains.
 - a) Mixture of o-and p-bromotoluenes
- b) Mixture of o-and p-dibromobenzenes
- c) Mixture of o-and p-bromoanilines
- d) Mixture of o-and m-bromotoluenes
- 8. C=0 compounds reacts with NH₃ or amines followed by H₂/Ni. The reaction is called
 - a) Mendius reaction

b) Hofmann bromamide

c) Reductive amination

- d) Gabriel's phthalimide
- 9. A compound which on reaction with aqueous nitrous acid gives an oily nitrosoamine is:
 - a) Methylamine
- b) Ethylamine
- c) Diethylamine
- d) Triethylamine

10.



The product A is

b)
$$NO_2$$

d)
$$\sim$$
 O \sim NO₂

- 11. The active species produced in Hofmann's bromamide reaction is:
 - a) Br⁻
 - b) Br₂
 - c) OBr
 - d) OBr₂
- 12. $C_5H_{13}N$ reacts with HNO_2 to give an optically active alcohol. The compound is
 - a) Pentan-1-amine

b) Pentan-2-amine

c) N, N-dimethylpropan-2-amine

- d) N-methylbutan-2-amine
- 13. Reduction of alkyl nitriles, produces

a) Secondary amine

b) Primary amine

c) Tertiary amine

d) amide

14. Which one of the following compound is most basic?

$$(A) \qquad (B) \qquad (C)$$

a) (A)

b) (B)

c) (C)

- d) All are equally basic
- 15. Alkyl halide (*RX*) on treatment with KCN followed by reduction leads to formation of:
 - a) RNH-

- b) RCH₂NH₂
- c) $RH + NH_3$
- d) $RCH_3 + N_2$
- 16. A gaseous carbon compound is soluable in dilute HCI. The solution on treating with ${\rm NaNO_2}$ gives off nitrogen leaving behind a solution which smells of wood spirit. The carbon compound is
 - a) HCHO
- b) CO

- c) $C_2H_5NH_2$
- d) CH₃NH₂
- 17. Benzaldehyde condenses with N, N-diamethylaniline in presence of anhydrous ZnCl₂ to give
 - a) Azo dye
- b) Malachite green
- c) Michler's ketone
- d) Buffer yellow

- 18. Which of the following statements are correct?
 - a) Aniline is a stronger base ethyl amine
 - b) Aniline is a stronger base than *p*-methoxyaniline
 - c) Aniline must be acetylated before nitration with an acid mixture
 - d) Aniline is soluble in an ammonium hydroxide solution
- 19. CHCl₃ and KOH on heating with a compound from a bad smelling product, compound is
 - a) C₂H₅CN
- b) C_2H_5NC
- c) C_2H_5OH
- d) $C_2H_5NH_2$
- 20. On heating urea, a gas evolves along with formation of biuret. Identify the gas.
 - a) CO

b) NH₃

c) CO_2

d) H_2

The alkene formed as a major product in the above elimination reaction is

- a) Ma
- b) $CH_2 = CH_2$
- c) Me

- 22. $CH_3NH_2 + CHCI_3 + KOH \rightarrow nitrogen containing compound + KCI + H_2O$. Nitrogen containing compound is
 - a) $CH_3 C \equiv N$
- b) $CH_3 NH CH_3$
- c) $CH_3 \longrightarrow \stackrel{-}{N} \longrightarrow \stackrel{+}{C}$ d) $CH_3 \stackrel{+}{N} \longrightarrow C$

- 23. A secondary amine is:
 - a) A compound with two —NH₂ groups
 - b) A compound with 2 carbon atoms and a —NH₂ group
 - c) A compound with a —NH₂group on the carbon atom in number 2 position
 - d) A compound in which 2 of the hydrogens of NH₃ have been replaced by alkyl or aryl groups

24.

$$\mathsf{F} - \bigvee \mathsf{NO}_2 \xrightarrow{(\mathsf{CH}_3)_2 \mathsf{NH}} A \longrightarrow$$

a)
$$H_2N$$
 N CH_3 CH_3

- 25. The name urea given by:
 - a) Wöhler
- b) Berzelius
- c) Roulle
- d) Lemery

26. In the reaction

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

$$X \longrightarrow NO_2$$

$$NO_2$$

Xis

a) SiC

- b) H_2SO_4
- c) KMnO₄
- d) Fe/HCl
- 27. Which of the following enzymes can hydrolyse urea into CO₂ and NH₃?
 - a) Amylase
- b) Urease
- c) Lipase
- d) Zymase

 $C_6H_5NH_2 \xrightarrow{H_2SO_4} H_2NC_6H_4(SO_3H)$ (para)

The true statement about the product is

- a) It does not exist as Zwitter ion
- b) -NH2 displays a powerful basic character
- c) It does not act as inner salt
- d) -SO₃ dimminishes the basic character of -NH₂
- 29. Aniline on treatment with NaNO₂ in HCI at 0°C followed by treatment with alkaline β –naphthol gives
 - a) A violet solution

b) A red solution

c) A green solution

- d) A blue precipitate
- 30. Which of the test is used for detection of secondary amines?
 - a) Liebermann's nitroso test

b) Lucas test

c) Tollen's test

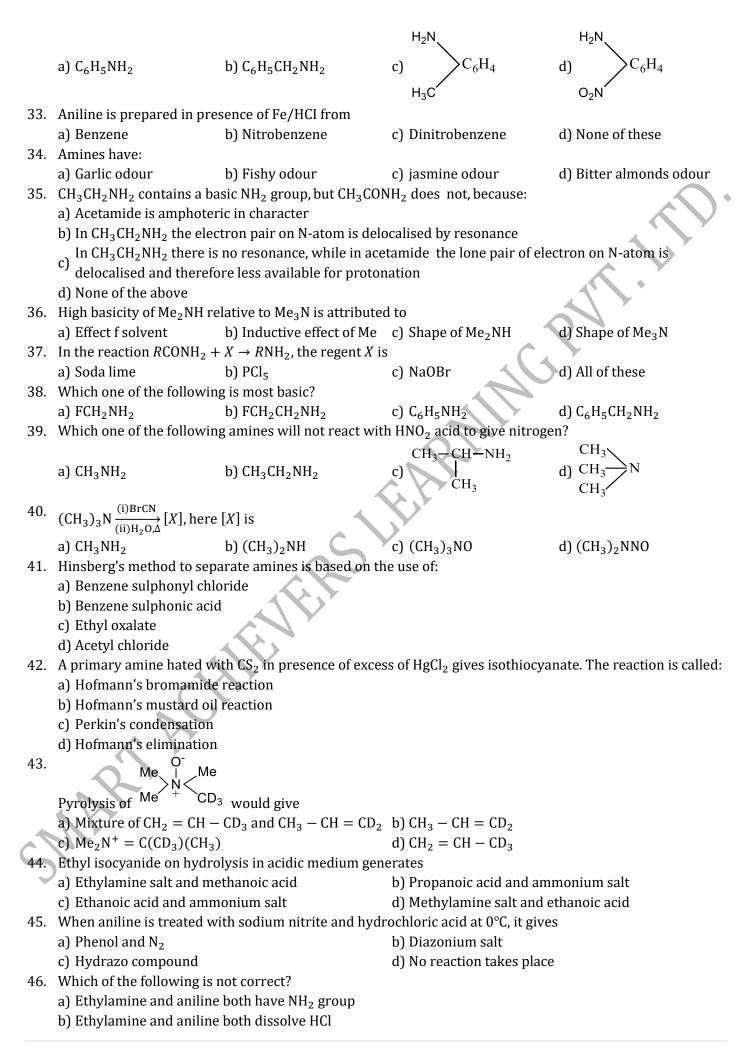
- d) Carbylamine reaction
- 31. Gas evolved during the reaction of sodium metal on ethyl amine is:

b) C_2H_2

c) H₂

d) CO_2

32. Which will not go for diazotization?



- c) Ethylamine and aniline both react with CHCl₃ and KOH to form unpleasant smell d) Ethylamine and aniline both react with NaNO₂ + HCl to give hydroxyl compounds in cold 47. Amine is not formed in the reaction
 - (A) Hydrolysis of RCN
 - (B) Reduction of RCH = NOH
 - (C) Hydrolysis of RNC
 - (D) Hydrolysis of RCONH₂

The correct answer is

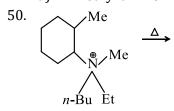
- a) A, B, D
- b) A, D

c) B, C

- d) A, B, C
- 48. During coupling reaction of benzene diazonium chloride and aniline, the pH of reaction medium should be approximately
 - a) 1-2

- b) 9-10
- c) 4-5

- 49. The amine which will not liberate nitrogen on reaction with nitrous acid is
 - a) Trimethyl amine
- b) Ethyl amine
- c) Sec-butyl amine
- d) t-butyl amine



The alkane formed as a major product in the given elimination reaction is:

- b) $CH_2 = CH_2$

- 51. Carbylamine reaction is given by aliphatic
 - a) Primary amine
 - c) Tertiary amine

- b) Secondary amine
- d) Quaternary ammonium salt
- 52. Nitrobenzene is reduced by Zn and alcoholic potash mixture to get
 - a) $C_6H_5 NH_2$

b) $C_6H_5 - NH - NH - C_6H_5$

c) $C_6H_5 - N - N - C_6H_5$

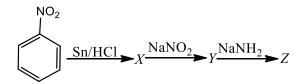
- d) $C_6H_5 NH CO C_6H_5$
- 53. The decreasing order of basic characters of the three amines and ammonia is
 - a) $NH_3 > CH_3NH_2 > C_2H_5NH_2 > C_6H_5NH_2$
- b) $C_2H_5NH_2 > CH_3NH_2 > NH_3 > C_6H_5NH_2$
- c) $C_6H_5NH_2 > C_2H_5NH_2 > CH_3NH_2 > NH_3$
- d) $CH_3NH_2 > C_2H_5NH_2 > C_6H_5NH_2 > NH_3$
- 54. Which of the following is strongest base?
 - a) $C_6H_5NH_2$
- b) $p NO_2 C_6H_4NH_2$
- c) $m NO_2 C_6H_4NH_2$ d) $C_6H_5CH_2CH_2$

- 55. Benzyl amine cannot be prepared by
 - a) $C_6H_5CONH_2 \xrightarrow{\text{LiAIH}_4}$

b) $C_6H_5CH_2CONH_2 + Br_2 + KOH \rightarrow$

- d) $C_6H_5CH_2NC \xrightarrow{LiAlH_4}$
- 56. Urea when heated a white residue is formed. Its alkaline solution when treated with few drops of CuSO₄ solution gives:
 - a) Red colour
- b) Violet colour
- c) Green colour
- d) Yellow colour
- 57. An organic compound 'A' having molecular formula C_2H_3N on reduction gave another compound B, upon treatment with nitrous acid 'B' gave ethyl alcohol. On warming with chloroform and alcoholic KOH, it formed an offensive smelling compound 'C'. The compound 'C' is
 - a) CH₃CH₂NH₂
- b) $CH_3CH_2N \Longrightarrow C$ c) $CH_3C \equiv N$
- d) CH₃CH₂. OH

58. What is 'Z'in the following reaction?



- a) Benzoic acid
- b) Cyanobenzoic acid
- c) Benzamide
- d) Aniline
- 59. Amino group is *ortho*/ *para*-directing for aromatic electrophilic substitution. On nitration of aniline, a good amount of *m*-nitroaniline is obtained. This is due to
 - a) In nitration mixture, ortho, para-activity of NH2 group is completely lost
 - b) -NH₂ because NH₃⁺, which is m-directing
 - c) $-NH_2$ becomes $-NH^+SO_4^-$, which is *m*-directing
 - d) $-NH_2$ becomes $-NH^-NO_2^+$, which is *m*-directing
- 60. Carbonyl chloride reacts with ammonia to form:
 - a) CO_2

- b) NH₂CONH₂
- c) CH₃COONH₄
- d) CH₃CONH₂

- 61. The action of nitrous acid on a primary amine gives:
 - a) Nitroalkane
- b) Alkyl nitrite
- c) Alcohol
- d) Secondary amine

- 62. The reduction of CH₃CN to CH₃CH₂NH₂ is called:
 - a) Rosenmund's reduction
 - b) Clemmensen's reduction
 - c) Mendius reduction
 - d) Hofmann's reduction
- 63. Aniline is reacted with Br₂ water and the resulting product is treated with an aqueous solution of sodium nitrite in the presence of diluteHCl. The compound so formed is converted into tetrafluoroborate which is subsequently heated dry. The end product is
 - a) p-bromofluorobenzene

b) *p*-bromoaniline

c) 2, 4, 6- tribromofluoro benzene

d) 1, 3, 5- tribromobenzene

64. The reaction,

$$RCOOH \xrightarrow{\text{NaN}_3/\text{conc.H}_2SO_4} RNH_2 + N_2 + CO_2$$

is known as

- a) Curtius reaction
- b) Lossen reaction
- c) Schmidt reaction
- d) Hofmann reaction
- 65. Which of the following compounds on treatment first with NaNO₂/HCI and then coupled with phenol produces *p*-hydroxyazobenzene?
- a) Nitrobenzene
- b) Azobenzene
- c) Phenol
- d) Aniline

- 66. The structural formula of methyl amino methane is:
 - a) (CH₃)₂CHNH₂
- b) (CH₃)₃N
- c) $(CH_3)_2NH$
- d) CH₃NH₂
- 67. An organic compound (C_3H_9N) (A), when treated with nitrous acid, gave an alcohol and N_2 gas was evolved. (A) on warming with $CHCl_3$ and caustic potash gave (C) which on reduction gave isopropylmethylamine. Predict the structure of (A).

a)
$$CH_3$$
 $CH-NH_2$

c)
$$CH_3 - N - C$$

- d) CH₃CH₂CH₂— NH₂
- 68. Urea when heated slowly, product formed is:
 - a) N₂
 - b) CO₂
 - c) biuret
 - d) Ammonium carbamate

- 69. Which of the following statements is not correct?

 a) Primary amines show intermolecular hydrogen bonding
 - b) Secondary amines show intermolecular hydrogen bonding
 - 2) Tartiana anima ah animbana ah animbana ah animbana ah animbana
 - c) Tertiary amines show intermolecular hydrogen bonding
 - d) Amines have lower boiling points as compared to those of alcohols and carboxylic acids of comparable molar masses
- 70. Compare boiling point of isomeric alkyl amines.
 - a) $1^{\circ} > 2^{\circ} > 3^{\circ}$
- b) $1^{\circ} > 2^{\circ} < 3^{\circ}$
- c) $1^{\circ} < 2^{\circ} < 3^{\circ}$
- d) $1^{\circ} < 2^{\circ} > 3^{\circ}$

- 71. Hofmann's hypobromite reaction affords a method of:
 - a) Preparing a tertiary amine
 - b) Preparing a mixture of amines
 - c) Stepping down a series
 - d) Stepping up a series
- 72. A colourless, odourless and non-combustible gas is liberated when ethylamine reacts with:
 - a) NaOH
- b) CH₃COCl
- c) $NaNO_2 + HCl$
- d) H₂SO₄

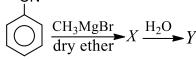
- 73. Reaction of benzaldehyde with methylamine gives
 - a) C₆H₅COOH

b) $C_6H_5N = NCl$

c) $C_6H_5 - CH = N - CH_3$

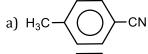
d) $C_6H_5NH_2$

74. Çİ



Identify Y

- a) Benzophenone
- b) Acetophenone
- c) Benzoic acid
- d) phenol
- 75. What is the proper sequence of reagent in the Hofmann's degradation reaction?
 - a) Br_2 , KOH, H_2O
- b) KOH, Br₂, H₂O
- c) H₂O, KOH, Br₂
- d) KOH, H₂O, Br₂
- 76. The reaction of chloroform with alcoholic KOH and *p*-toluidine form



b)
$$H_3C$$
 \longrightarrow N_2CI

d)
$$H_3C$$

- 77. Ethyl isocyanide on hydrolysis in acidic medium generated
 - a) Ethyl amine salt and methanoic acid
- b) Propanoic acid and ammonium salt
- c) Ethanoic acid and ammonium salt
- d) Methyl amine salt and ethanoic acid
- 78. When methyl iodide is treated with ammonia, the product obtained is:
 - a) Methylamine
- b) Dimethylamine
- c) Trimethylamine
- d) All of these

- 79. Aliphatic amines are soluble in water because:
 - a) They are basic
 - b) They are amino compounds
 - c) They are lighter than water
 - d) Of formation of hydrogen bonds with water
- 80. An organic amino compound reacts with aqueous nitrous acid at low temperature to produce an oily nitrosoamine. The compound is
 - a) CH₃NH₂
- b) CH₃CH₂NH₂
- c) CH₃CH₂NHCH₂CH₃
- d) $(CH_3CH_2)_3N$

- 81. Allyl isocyanide containsand.....bonds.
 - a) 9σ , 3π
- b) 9σ , 9π
- c) 3σ, 4π
- d) 5σ , 7π

- 82. Mendius method of preparation of amines consists of:
 - a) Catalytic reduction of alkyl cyanides
 - b) Reduction of amide with LiAlH₄
 - c) Reduction of nitroparaffin with Sn + HCl

	d) Reduction of oximes with Na + C ₂ H ₅ OH		
83.	The compound having the molecular formula C ₃ H ₉ I	N represent :	
	a) Trimethylamine b) <i>n</i> -propylamine	c) Isopropylamine	d) All of these
84.	From the following compounds which does not read	ct with C ₆ H ₅ SO ₂ CI?	
	a) C ₂ H ₅ .NH ₂	b) CH ₃ . NH ₂	
	c) (CH ₃) ₂ NH	d) $(C_2H_5)_3N$	
85.	Identify <i>A</i> and <i>B</i> in the reaction given below.	7 (2 3/3	
	Wandaraharia Danada ambatian		
	Ethane nitrile $\xrightarrow[+2H_2O]{\text{Hydrolysis}} A \xrightarrow[]{\text{Decarboxylation}} B$		
	$^{+2} ext{H}_2 ext{O}$ $^{\Delta}$ $^{-} ext{CO}_2$		
	a) Acetic acid, methanol	b) Acetone, methane	
	c) Ethanoic acid, ethane	d) Ethanoic acid, methan	e
86.	The compound formed when malonic ester reacts w	-	A . Y
	a) Cinnamic acid b) Butyric acid	c) Barbituric acid	d) Crotonic acid
87.	Decreasing order of basicity of the three isomers of	,	
	a) $p-CH_3OC_6H_4NH_2 > o-CH_3OC_6H_4NH_2 > m-CC$	•	
	b) p -CH ₃ OC ₆ H ₄ NH ₂ > m - CH ₃ OC ₆ H ₄ NH ₂ > o - C		
	c) $o - CH_3OC_6H_4NH_2 > p - CH_3OC_6H_4NH_2 > m - CH_3OC_6H_4NH$	$H_3OC_6H_4NH_2$	
	d) o -CH ₃ OC ₆ H ₄ NH ₂ > m - CH ₃ OC ₆ H ₄ NH ₂ > p - C		
88.	Nitrogen of nitrobenzene at 125°C with mixed acid		
	a) <i>meta</i> -dinitrobenzene	b) <i>ortho</i> -dinitrobenzene	
	c) <i>para</i> -dinitrobenzene	d) 1, 3, 5-trinitrobenzene	
89.	The value of K_b is highest in case of:		
	a) <i>p</i> -methoxy aniline b) <i>p</i> -chloroaniline	c) <i>p</i> -nitroaniline	d) p-methylaniline
90.	Benzene diazonium chloride on reaction with phene	ol in weakly basic medium g	gives
	a) Diphenyl ether b) <i>p</i> -hydroxy azobenzen	e c) Chlorobenzene	d) Benzene
91.	$R - N = C + HgO \longrightarrow A + Hg_2O$; What is A?	7	
	a) RNH ₂ b) RCONH ₂	c) R—NCO	d) RCOOH
92.	Amine oxide, when heated froms alkene. The reaction	on is known as	
	a) Curtius	b) Cope elimination	
	c) Mannich reaction	d) Hofmann elimination	
93.	Identify the product in the following sequence $3,4$,	5-tribromoaniline	
	(i)Diazotization		
	(ii)H ₃ PO ₂	1) 4 2 2	
	a) 3, 4, 5-tribromobenzene	b) 1, 2, 3-tribromobenzer	
0.4	c) 2, 4, 6-tribromobenzene	d) 3, 4, 5-tribromonitrob	enzene
94.	Identify the product Z in the following reaction scho	eme	
	$C_6H_5NH_2 \xrightarrow{Ac_2O} X \xrightarrow{Br_2/CCl_4} Y \xrightarrow{HOH} Z$		
	a) <i>p</i> -bromoaniline	b) p-bromoacetophenone	
	c) p-bromoacetanilide	d) o-bromoacetophenone	
95.	In the following reaction, $X \xrightarrow{\text{Bromination}} Y \xrightarrow{\text{NaNO}_2} Z \xrightarrow{\text{Boil}} Z \xrightarrow{\text{C}_2 \text{H}}$	$\stackrel{\text{ling}}{\longrightarrow}$ tribromo benzene. X is	
		c) Phenol	
06	a) Benzoic acid b) Salicylic acid The compound, N-ethyl-N-methylpropanamine forr	•	d) Aniline
70.	show optical activity. This is due to	iis iioii- superiiiiposable iiii	iroi illiage but does not
	a) Absence of a chiral N-atom	b) Presence of a chiral N-	atom
	c) Presence of lone pair on N-atom	d) Rapid flipping of one fi	
97	Which of the following statement about primary am	, , , , ,	om mito another
<i>,,</i> ,	a) Alkylamines are stronger base than arylamines		
	b) Alkylamines react with nitrous acid to produce a	alcohols	
	5, 1, rainines react with the ous acid to produce a		

- c) Arylamines react with nitrous acid to produce phenols
- d) Alkylamines are stronger bases than ammonia.
- 98. How may primary amines are possible for the formula C₄H₁₁N?
 - a) 1

b) 2

c) 3

- d) 4
- 99. What is decreasing order of basicity of p-, s-, t-ethyl amines and NH_3 ?
 - a) $NH_3 > C_2H_5NH_2 > (C_2H_5)_2NH > (C_2H_5)_3N$
 - b) $(C_2H_5)_3N > (C_2H_5)_2NH > C_2H_5NH_2 > NH_3$
 - $(C_2H_5)_2NH > C_2H_5NH_2 > NH_3 > (C_2H_5)_3N/'$
 - d) $(C_2H_5)_2NH > (C_2H_5)_3N > C_2H_5NH_2 > NH_3$
- 100. In the reaction

$$CH_3CN + 2H \xrightarrow{HCI}_{SnCl_2} X \xrightarrow{Boiling H_2O} Y$$
,

The term Y is,

- a) Acetone
- b) Ethanamine
- c) Acetaldehyde
- d) Dimethyl amine

- 101. Which is not the property of ethanenitrile (CH₃CN)?
 - a) Undergoes acidic hydrolysis to give carboxylic acid
 - b) Undergoes alkaline hydrolysis to give salt of carboxylic acid
 - c) It tautomerises to give methyl isocyanide
 - d) It gives carbylamines reaction with chloroform
- 102. Acetoneoxime on catalytic hydrogenation gives:
 - a) 1-propanamine
- b) Isopropylamine
- c) Ethyl methyl amine
- d) CH4 and ethanamine

103. The product of Hofmann elimination of

$$\leftarrow$$
 CH₂N(CH₃)₂OH is

a)
$$\langle _ \rangle$$
 $- CH_2N(CH_3)_2$

c)
$$\langle \bigcirc \rangle$$
 = CH₂ + (CH₃)₂NOH

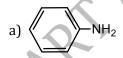
b)
$$\langle \bigcirc \rangle$$
 NH₂ + (CH₃)₂N

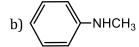
d)
$$\left\langle \right\rangle + (CH_3)_4 \stackrel{+}{\text{NOH}}$$

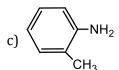
- 104. Hofmann's rearrangement during the conversion of an amide to amine involves....... rearrangemet.
 - a) Intermolecular
- b) Intramolecular.
- c) Both (a) and (b)
- d) None of these

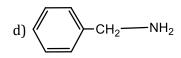
- 105. Aniline reacts with ... to yield ... as the final product.
 - a) Bromine, 2-bromoaniline

- b) Bromine, 2, 4, 6-tribromoaniline
- c) Chloroform/KOH, phenyl cyanide
- d) Acetyl chloride, benzanilide
- 106. Which of the following is the strongest base?









- 107. Which of the following can be used to distinguish acetamide and urea?
 - a) Fehling's solution
- b) Biuret test
- c) Hofmann's reaction
- d) NaOH solution

- 108. Which of the following amines is optically active?
 - a) CH₃NH₂
 - b) CH₃NHCH₃

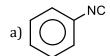
- d) Sec. butylamine
- 109. Which one of the following is not the correct reaction of aryl diazonium salts?

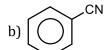
a)
$$C_6H_5N_2^+CI^- + Cu_2CI_2 \rightarrow C_6H_5CI$$

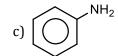
b)
$$C_6H_5N_2^+CI^- + HBF_4 \xrightarrow{Heat} C_6H_5F$$

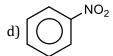
c) $C_6H_5N_2^+CI^- + H_3PO_2 \rightarrow C_6H_5PO_4$ d) $C_6H_5N_2^+CI^- + SnCI_2/HCI \rightarrow C_6H_5NHNH_2$ 110. Hinsberg's reagent is a) C₆H₅COCI b) CH₃COCI c) C₆H₅CH₂CI d) C₆H₅SO₂CI 111. Which one of the following compound when heated with KOH and primary amines gives carbylamine test? b) CH₃Cl d) CH₃NC a) CHCl₃ c) CCl₄ 112. Ethyl amine on acetylation gives a) N-ethyl acetamide b) Acetamide c) Methyl acetamide d) None of these 113. The oxidation of aniline with per acetic acid in the presence of acetic acid by refluxing gives d) None of these 114. Aniline reacts with acetaldehyde to form d) None of these a) Schiff's base b) Carbylamine c) Immine 115. Aniline gives a precipitate with bromine. The colour of precipitate is d) White b) Black c) Blue 116. a) Chain b) Functional c) Position d) All of these 117. A compound of molecular formula C₃H₉N when reacts with benzene sulphonyl chloride gives a product soluble in dilute NaOH solution. The compound should be d) All of these b) $(CH_3)_2CH - NH_2$ a) $(CH_3)_3N$ 118. Which one does not liberate NH₃ when undergoes hydrolysis? a) Acetanilide b) Acetonitrile c) Acetamide d) Phenyl isocyanide 119. n-butylamine (I), diethylamine (II) and N, N-dimethylethylamine (III) have the same molar mass. The increasing order of their boiling point is a) III < II < Ib) I < II < III c) II < III < Id) II < I < III120. Correct order of basic nature of $CH_3NH_2(A)$, $CH_3CN(B)$ and $CH_3N = CHCH_3(C)$ is a) A > B > Cb) B > C > Ac) A > C > Bd) C > A > B121. Nitroparaffins on reduction give: a) Amides b) Alkylamines c) Ammonium salts d) Acetanilides 122. Which of the following is not characteristic of amines? a) They smell like ammonia b) They are inflammable in air c) They show the property of hydrogen bonding d) They are amphoteric in nature 123. On heating benzyl amine with chloroform and ethanolic KOH, product obtained is a) Benzyl alcohol b) Benzaldehyde c) Benzonitrile d) Benzyl isocyanide 124. Benzyl amine reacts with nitrous acid to give a) Azobenzene b) Benzene c) Benzyl alcohol d) Phenol 125. Which of the following statements is not correct? a) Alkyl isocyasnides have bad odours while alkyl cyanides have pleasant odours b) Alkyl cyanides are not as poisonous as KCN

c) Alkyl cyanides have lower boiling points than the corresponding alkyl isocyanides d) Acetonitrile is soluble in water but methylcarbylamine is not										
126. When NaNO ₂ and dilute HCl were added to an am		s was evoloued and an ionic								
compound is formed. The amine is:	ine at 0 c, a colouriess gas	was evoloved and an ionic								
a) An primary amine										
b) An aromatic primary amine										
c) Any amine										
d) None of the above										
127. Which of the following reactions is given by only pri	mary amines?									
a) Reaction with HONO	b) Reaction with chlorofo	orm and alcoholic KOH								
c) Reaction with acetyl chloride	d) Reaction with Grignar									
128. In hypobromite reaction of amide, carbonyl carbon	atom is lost as:									
a) CO b) CO ₂	c) CO_3^{2-}	d) None of these								
129. Correct order of basicity of $\phi NH_2[A]$, \bigcirc $\rightarrow NH_2[B]$,										
OCH ₃		V								
$\langle \bigcirc \rangle$ -NH ₂ [C] H ₃ CO- $\langle \bigcirc \rangle$ -NH ₂ [D] is	. C 4									
OCH ₃ a) $A > B > C > D$ b) $D > C > A > B$	c) $B > D > C > A$	d) $D > A > B > C$								
	C) B > D > C > A	u) D > A > B > C								
130. The IUPAC name of CH_3 – CH – CH_2 – CH – CH_3 is:										
$\overset{1}{\text{CH}_{3}}$ $\overset{1}{\text{CN}}$										
a) 2-cyano-3-methyl hexane										
b) 2-dimethyl-4-cyanopentane										
c) 2,4-dimethyl pentane nitrile										
d) 2-cyano-3-methyl hexane										
131. Choose the incorrect statement.										
a) Primary amines show intermolecular hydrogen b	onds.									
b) Tert-butylamine is primary amine.										
c) Tertiary amines do not show intermolecular hydr	rogen bonds.									
d) Isopropylamine is a secondary amine.										
132. N_2 gas is liberated when [HCl + NaNO ₂] reacts with	the following compounds									
(A)CH ₃ CH ₂ NH ₂ (B) Urea										
$(C)CH_3CONH_2$ $(D)C_6H_5NH_2$										
The answer is										
a) A, B, C b) B, C, D	c) A, C, D	d) A, B, D								
133. Urea on heating with ethanol gives:	> xx	D.M. C.I								
a) Urethane b) Urea alcohol	c) Ureides	d) None of these								
134. Treatment of nitrobenzene with acetyl chloride in th		AICI ₃ gives								
a) 2-nitroacetophenone	b) 3-nitroacetophenone									
c) 4-nitroacetophenone	d) None of these	a								
135. The correct sequence of reactions to convert p-nitro	= =									
a) Reduction, diazotization and hydrolysisc) Hydrolysis, reduction and diazotization	b) Hydrolysis, diazotizatid) Diazotization, reduction									
136. The reduction of which of the following compound	=									
a) Alkyl nitrile	b) Carbylamine	iiiie:								
c) Primary amine	d) Secondary nitro comp	ound								
137. The compound with foul odour among the following	_	vuiiu								
137. The compound with rour odour among the following	, 10									









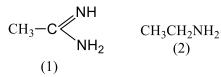
- 138. Reduction of nitrobenzene in the presence of Zn/NH₄Cl gives
 - a) Azobenzene

b) Hydrazobenzene

c) N-phenyl hydroxylamine

- d) Aniline
- 139. Name of method use to separate primary, secondary and tertiary amines is
 - a) Hofmann method
- b) Lucas method
- c) Victor Meyer method
- d) Kolbe method

140. The correct order of basicities of the following compound is



$$(CH_3)_2NH$$
 (3)

$$CH_3 - C - NH_2$$

$$(4)$$

- a) 2>1>3>4
- b) 1>3>2>4
- c) 3>1>2>4
- d) 1>2>3>4

- 141. Dye test can be used to distinguish
 - a) Ethyl amine and acetamide

b) Ethyl amine and aniline

c) Urea and acetamide

- d) Methyl amine and ethyl amine
- 142. In the reaction of (S) 2-phenylpropamide with NaBr/H₂O to give 1-phenylethylamine
 - a) There is retention of configuration
- b) There is inversion of configuration
- c) A mixture of two products is obtained
- d) There is no reaction
- $143. RNH_2$ reacts with $C_6H_5SO_2Cl$ in aqueous KOH to give a clear solution. On acidification a precipitate is obtained which is due to the formation of

b)
$$R - N^{-}SO_{2}C_{6}H_{5}K^{+}$$

c) $C_6H_5SO_2NH_2$

d) $R - NH - SO_2 - C_6H_5$

144. The reaction,

$$CH_3CN + 4H \xrightarrow{Na/C_2H_5OH} CH_3CH_2NH_2$$
 is called:

- a) Hofmann's bromamide reaction
- b) Mendius reaction
- c) Sabatier reaction
- d) None of the above

F—
$$O_2$$

DMF (CH₃)₂ NH
$$A = \frac{\text{(i) NaNO}_2/\text{HCl, 0-5}^{\circ}\text{C}}{\text{(ii) H.Ps}} B$$

In the above sequence *B* is

a)
$$H_2N$$
— O — N
 CH_3
 CH_3

$$C)$$
 O_2N-CH_3 CH_3 CH_3

d)
$$O_2N - \bigcirc -NH_2$$

146. A compound A when reacted with PCl_5 and then with ammonia gave B. B when treated with bromine and caustic potash produced C. C on treatment with $NaNO_2$ and HCl at $0^{\circ}C$ and then boiling produced

orthocresol. Compound *A* is:

- a) o-toluic acid
- b) o-chlorotoluene
- c) o-bromotoluene
- d) m-toluic acid

147.
$$C_2H_5NH_2 \xrightarrow{HNO_2} A \xrightarrow{PCI_3} B \xrightarrow{NH_3} C$$
.

Recognize the compound C from the following

- a) Propanenitrile
- b) Methylamine
- c) Ethylamine
- d) Acetamide

148. The conversion

$$C_6H_5NO_2 \longrightarrow C_6H_5N = N - C_6H_5$$

Can be brought about by reduction with

- a) Na₃AsO₃/NaOH
- b) Glucose/HCl
- c) Zn/NaOH
- d) LiAlH₄/ether

- 149. Benzoyl chloride does not react with:
 - a) Primary or secondary amines
 - b) Aliphatic compounds
 - c) Aromatic compounds
 - d) Carboxylic acids
- 150. Which compound will liberate CO₂ from NaHCO₃ solution?

- c) $(CH_3)_4N^+OH^-$
- d) CH₃N⁺H₃CL⁻

151.
$$C_6H_5NH_2 \xrightarrow{NaNO_2} X \xrightarrow{Cu_2(CN)_2} Y \xrightarrow{H_2O/H^+} Z$$

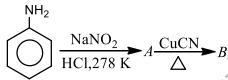
Z is identified as

- a) $C_6H_5 NH CH_3$
- b) $C_6H_5 COOH$
- $-CH_2 NH_2$
- d) $C_6H_5 CH_2COOH$

- 152. Ketones and 1° amines react to form:
 - a) Amides

d) Anils

153. In the chemical reactions,



Compounds A and B respectively are

- a) Fluorobenzene and phenol
- c) Nitrobenzene and chlorobenzene
- b) Benzene diazonium chloride and benzonitrile
- d) Phenol and bromobenzene

- 154. Dehydration of an amide gives:
 - a) Cyanide
- b) Amine
- c) Isocyanide
- d) Fatty acid

155. Given the following sequence of reactions,

$$\operatorname{CH_3CH_2I} \xrightarrow{\operatorname{NaCN}} A \xrightarrow[\operatorname{Partial}\\ \operatorname{hydrolysis}]{\operatorname{OH}} B \xrightarrow{\operatorname{Br_2/NaOH}} C$$

The major product C' is

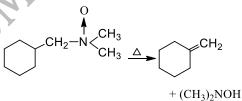
$$CH_3$$
. $CH_2C - NHBr$

 CH_3 . $CH_2C - NBr_2$

a) CH₃CH₂NH₂

Ш

c) CH_3 . $CH_2 - COONH_4$



This reaction is called

- a) Cope reaction
- b) Ritter reaction
- c) Schmidt reaction
- d) Gabriel reaction
- 157. Which one of the following compounds forms a quaternary salt on reacting with excess methyl iodide?
 - a) $C_2H_5OCH_3$
- b) $(CH_3)_2CHOC_2H_5$
- c) $C_6H_5NH_2$
- d) $C_6H_5NO_2$

158. Which of the following reactions can produce aniline as main product?

a) $C_6H_5NO_2 + Zn/KOH$

b) $C_6H_5NO_2 + Zn/NH_4Cl$

c) $C_6H_5NO_2 + LiAlH_4$

d) $C_6H_5NO_2 + Zn/HCl$

159. Reaction of aniline with benzaldehyde is

- a) Substitution
- b) Addition
- c) Condensation
- d) Polymerisation

160. The product *D* in the following sequence of reactions is,

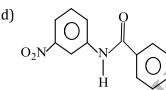
$$CH_3COOH \xrightarrow{NH_3} A \xrightarrow{Heat} B \xrightarrow{P_2O_5} C \xrightarrow{Na+C_2H_5OH} D:$$

- a) Ester
- c) Acid

d) Alcohol

161. In the following reaction, the product *X* is:

$$\begin{array}{c|c} & & & \\ & & & \\$$



162. Indicate the correct statement.

- a) C₂H₅N⁺H₃OH⁻ is acidic
- b) C₂H₅NH₂ is less basic than NH₃
- c) C₂H₅NH₂ is a stronger base than NH₃
- d) C₂H₅NH₂ forms salts with bases

163. The compound will react most readily with NaOH to form methanol is

- a) $(CH_3)_4N^+I^-$
- b) CH₃OCH₃
- c) $(CH_3)_3S^+I^-$
- d) $(CH_3)_3CI$

164. Increasing order of basicity of $CH_3CH_2CH_2NH_2$ $H_2C = CHCH_2NH_2$ and $HC \equiv CCH_2NH_2$ is

- $CH_3CH_2CH_2NH_2 < HC \equiv CCH_2NH_2 < H_2C =$ CHCH₂NH₂
- $_{\mathrm{bl}}$ $^{\mathrm{CH_3CH_2CH_2NH_2}} < \mathrm{H_2C} = \mathrm{CHCH_2NH_2} < \mathrm{CH}$

 $\equiv CCH_2NH_2$

HC \equiv CCH₂NH₂ < H₂C = CHCH₂NH₂ < CH₃CH₂CH₂NH₂

 $_{\rm d}$ CH \equiv CCH₂NH₂ < CH₃CH₂CH₂NH₂ < H₂C = CHCH₂NH₂

165. Reduction of aniline with acetyl chloride in presence of NaOH produce

- a) Aniline hydrochloride b) Acetanilide
- c) p-choloroaniline
- d) A red dye

166. The molecular formula C₃H₉N cannot represent

- a) 1°amine
- b) 2°amine
- c) 3°amine
- d) Quaternary salt

167. (A) $C_2H_5NH_2 \xrightarrow{(ii)AgNO_2} [W]$

$$(B)(CH_3)_2CHNH_2 \xrightarrow[(ii)AgNO_2]{(ii)AgNO_2} [X]$$

$$(C)(CH_3)_3 CNH_2 \xrightarrow{(i)NOCl}_{(ii)AgNO_2} [Y]$$

(D)CH₃CH(NH₂)C₂H₅
$$\xrightarrow{\text{(i)NOCl}}_{\text{(ii)AgNO}_2}$$
 [Z]

Which product will not show tautomerism?

168. Carcinogens are the products of the reaction between:

a)
$$R_2NH + HNO_2$$

b)
$$R_3N + HNO_2$$

c)
$$RNH_2 + HNO_2$$

169. Acetonitriles on hydrolysis produce which of the following?

a) Amine

170. Primary, secondary and tertiary nitroalkanes can be identified by the action of:

a)
$$HNO_2 + NaOH(aq.)$$

b)
$$CHCl_3 + NaOH(aq.)$$

c)
$$CHCl_3 + KOH(alc.)$$

171. Methyl cyanide gives on hydrolysis

a) Methyl amine

b) Acetic acid

c) Formic acid

d) Ethyl amine

172. The hydrochlorides of amines form double salt with:

a) PtCl₄

173. General formula of an amine is:

a)
$$C_n H_{2n+1} N$$

b)
$$C_n H_{2n+2} N$$

c)
$$C_nH_{2n+3}N$$

d)
$$C_n H_{2n} N$$

174. Aniline when diazotized in cold and then treated with dimethyl aniline gives a coloured product. Its structure would be

a)
$$CH_3NH$$
 $N=N$ $N+CH_3$

c)
$$N(CH_3)_2N$$
 $N=N$

d)
$$(CH_3)_2N$$
 $N=N$ NH_2

175. NH₂

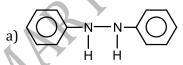
$$\frac{\text{(i) NaNO}_2/\text{HCl}}{\text{(ii) CuCN/H}_3\text{O}^+} A; A \text{ is}$$







176. The structure of the compound formed, when nitrobenzene is reduced by lithium aluminium hydride $(LiAIH_4)$ is



c)
$$\langle \bigcirc \rangle$$
-N=N- $\langle \bigcirc \rangle$

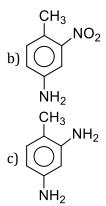
$$d) \bigcirc^{\mathsf{NH}_2}$$

177. Aniline and ethylamine resembles in:

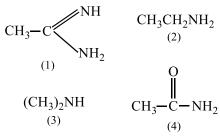
- a) Solubility
- b) Action with HNO₂
- c) Action of Grignard reagent
- d) Coupling reaction

178. Reaction of cyclohexanone with dimethylamine in the presence of catalytic amount of an acid forms a

compound of water during the reaction is continuously removed. The compound formed is generally known as a) An amine b) An imine c) An enamine d) A Schiff's base 179. Comparing basic strength of NH₃, CH₃NH₂ and C₆H₅NH₂ it may be concluded that a) Basic strength remains unaffected b) Basic strength of alkyl amines is lowest c) Basic strength of aryl amines is lowest d) Basic strength of NH₃ is highest 180. The product obtained when methylamine is treated with nitrous acid is: a) CH₂OH b) CH₃—ONO c) CH₃OCH₃ d) Both (b) and (c) 181. Hofmann bromamide reaction is used to prepare a) 1° amine b) 2° amine c) 3° amine d) All of these 182. Tertiary amine is obtained in the reaction a) Aniline $\xrightarrow{CH_3I} \xrightarrow{CH_3I}$ c) Nitrobenzene $\xrightarrow{Sn/HCl}$ b) Aniline $\stackrel{CH_3I}{\longrightarrow}$ d) None of these 183. Which of the following on reduction with LiAlH₄ gives a secondary amine? a) CH₃NC b) CH₃CONH₂ c) CH₃CN 184. Which of the following is most basic in aqueous medium? a) CH₃CH₂CH₂CH₂NH₂ b) $CH_3 - CH_2 - CH_2 = NH$ CH₃-N-CH₃ 185. The product *A* and *B* in the reaction are: $C_2H_5NH_2 + CHCl_3 + 3KOH \rightarrow A + B + 3H_2O$ a) $C_2H_5NC + 3KCl$ b) $C_2H_5CN + 3KCl$ c) $C_2H_5CONH_2 + 3KCl$ d) $C_2H_5NC + K_2CO_3$ 186. p – amine and s – amine are distinguished by: a) Br₂/KOH b) HClO d) NH₃ 187. Which one of the following compounds will dissolve in an alkali solution after it has undergone reaction with Hinsberg reagent? a) CH₃NH₂ b) $(CH_3)_3N$ c) $(C_2H_5)_2NH$ d) C₆H₅NHC₆H₅ 188. The reaction of chloroform with alcoholic KOH and p-toluidine from 189. Which one of the following functional groups undergoes hydrolysis with alkali to yield an acid group? b) -CHO a) -CN c) $-COCH_3$ d) -Br190. Ethylamine reacts with nitrosyl chloride (NOCl)to form: a) Ethyl chloride b) Ethyl alcohol c) Ethyl nitrite d) Nitroethane 191. The product obtained in the reduction ÇH₃ NO_2 H_2S , NH_3 is



- d) The compound is not reduced
- 192. The correct order of basic nature of the following compounds is:



- a) 2 > 1 > 3 > 4
- b) 1 > 3 > 2 > 4
- c) 3 > 1 > 2 > 4
- d) 1 > 2 > 3 > 4
- 193. In reduction of nitrobenzene, which of the following is the intermediate?

c)
$$\phi N = N - \phi$$

d)
$$\uparrow$$
 $\phi N = N -$

- 194. Nitration of aniline also gives *m*-nitro aniline in strong acidic medium because

 - a) In electrophilic substitution reaction amino group by $\frac{1}{m}$ Inspite of substituents nitro group always goes to $\frac{1}{m}$ is meta directive
 - c) In strong acidic medium aniline aniline present as d) None of the above
 - anilinium ion
- 195. Gabriel's sunthesis is used frequently for the preparation of which of the following?
 - a) Primary amines
- b) Primary alcohols
- c) Tertiary amines
- d) Tertiary alcohols

'X' is identified as

- c) NHOH
- d) None of these

- - In the above sequence A and C are
 - a) Methanal, methyl amine

b) Acetone, ethaneamine

c) Ethanal, diamethyl amine

d) Acetaldehyde, ethyl amine

198. In the following reaction, X is

$$X \xrightarrow{\text{Bromination}} Y \xrightarrow{\text{NaNO}_2} Z \xrightarrow{\text{Boiling}} \text{Tribromo benzene}$$

- a) Benzoic acid
- b) Salicyclic acid
- c) Phenol
- d) Aniline

199. Which of the following is not a nitro-derivative?

- a) $C_6H_5NO_2$
- b) CH₃CH₂ONO
- c) CH_3CH O D $C_6H_5(OH)NO_2$ CH_3
- 200. Decreasing order of basic nature in aqueous solutions
 - a) $C_6H_5NH_2 > NH_3 > CH_3NH_2 > (CH_3)_2NH$
- b) $NH_3 > C_6H_5NH_2 > CH_3NH_2 > (CH_3)_2NH$ d) $CH_3NH_2 > (CH_3)_2NH > NH_3 > C_6H_5NH_2$
- c) $(CH_3)_2NH > CH_3NH_2 > NH_3 > C_6H_5NH_2$
- 201. The IUPAC name of, CH₃-CH₂-CH-NH₂ is:
 - a) 1-methyl-amino propane
 - b) 2-aminobutane
 - c) 2-methyl-2-aminopropane
 - d) None of the above
- 202. When aqueous solution of benzene diazonium chloride is boiled, the product formed is
 - a) C₆H₅CH₂OH
- b) $C_6H_6 + N_2$
- c) C_6H_5COOH
- d) C_6H_5OH
- 203. Methylethylpropylamine forms non-superimposable mirror images but it does not show optical activity because:
 - a) Of rapid flipping
 - b) Amines are basic in nature
 - c) Nitrogen has a lone pair of electron
 - d) Of absences of asymmetric nitrogen

204.
$$Y \stackrel{\text{Reduction}}{=} [R_2 C = NH] \stackrel{\text{H}_3 O^+}{=} X$$

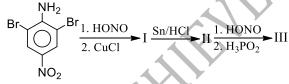
In the above sequence of reaction X, Y, Z are respectively

a) Aldehyde, ketone, NH₃

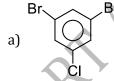
b) Ketone, 1° amine, KMnO₄

c) Ketone, 2° amine, KMnO₄

- d) Ketimine, 1° amine, H₂SO₅
- 205. The final product (III) obtained in the reaction sequence



is



b) b)

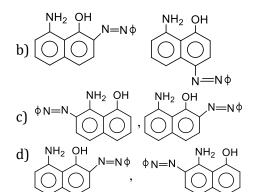
- c) Br CI Br
- d) None of these
- 206. Which of the following compound reacts with chloroform and a base to form phenyl isocyanide?
 - a) Phenol
- b) Aniline
- c) Benzene
- d) Nitrobenzene

Ph = 5-7

[X] and [Y]are
$$_{\text{NH}_2}$$
 OH $_{\text{NH}_2}$ OH

NH₂ OH

a)
$$\phi N=N$$
 $\downarrow NH_2 OH$ $\downarrow NH_2 OH$ $\downarrow NH_2 OH$ $\downarrow NH_2 OH$



208. In the reaction,

$$RNH_2 \xrightarrow{HNO_2} A + B + C \uparrow; C$$
 is

a) NH_a

b) N₂

c) 0_2

d) CO₂

209. The correct set of the products obtained in the following reactions is

$$(A)RCN \xrightarrow{Reduction}$$

(B)
$$RCN \xrightarrow{(I)CH_3MgBr}$$
 (II) H_2O

 $(C)RNC \xrightarrow{Hydrolysis}$

		HNO_2
(D)	RNH_2	\longrightarrow

	A	D	C	U
a)	2°amine	Methyl ketone	1°amine	Alcohol
c)	2°amine	Methyl ketone	2°amine	Acid

b)	1°amine	Methyl ketone	1°amine	Alcohol
d)	2°amine	Methyl ketone	2°amine	aldehyde

210. Which of the following is hydrolysed to give secondary amine?

a) Alkyl



c) Nitroparaffins

d) Acidamide

211. $\text{CH}_3\text{CH}_2\text{Br} \xrightarrow{aq.\text{KOH}} A \xrightarrow{\text{KMnO}_4/\text{H}^+} B \xrightarrow{\text{NH}_3} C \xrightarrow{\text{Br}_2} D, 'D' \text{ is}$

a) CH₃Br

- b) CH2CONH2
- c) CH₃NH₂
- d) CHBr₃

212. The product [A] formed in the reaction;

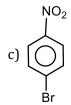
$$2C_5H_5 - CN \xrightarrow{Na} [A]$$
 is:

213.
$$\underbrace{ \frac{\text{HNO}_3}{\text{H}_2\text{SO}_4}} A \underbrace{ \frac{\text{Br}_2}{\text{FeBr}_3}} B$$

The compound *B* is







d)
$$Br$$
 Bl

214.
$$\left[\bigwedge \right] \xrightarrow{\text{I}^-} \underset{\text{H}_2\text{O}}{\text{Ag}_2\text{O}} X \xrightarrow{400 \text{ K}}$$

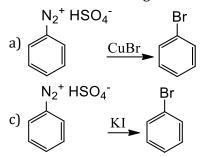
The products of above sequence of reactions are

a) $CH_2 = CH_2$ and $(CH_3)_2CHCN$

b) CH₃CH₂CN and C₂H₅NH₂

c) $CH_2 = CH_2$ and $(CH_3)_3N$

- d) $(CH_3)_2$ C = CH_2 and NH_3
- 215. Which of the following reactions is an example of Sandmeyer reaction?



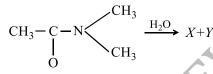


216. Isopropyl amine with excess of acetyl chloride will give

a)
$$(CH_3CO)_2N - CH - (CH_3)_2$$

$$(CH_3)_2CH - N - COCH_3$$
b) | H
$$CH_3CH_2CH_2 - N - COCH_3$$
d) |

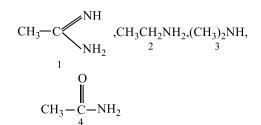
- c) (CH₃)₂CHN(COCH₃)₂
- 217. *X* and *y* in the given reaction are:



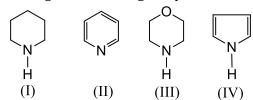
- a) $CH_3COOH + (CH_3)_2NH_1$
- b) $CH_3CONH_2 + CH_3OH$
- c) $CH_3CHO + (CH_3)_2NH$
- d) $CH_3COCH_3 + CH_3NH_2$
- 218. Primary nitroalkanes on hydrolysis give:
 - a) $RCOOH + NH_2OH$
- b) RCOOH
- c) NH₂OH
- d) RCOR
- 219. Aliphatic nitriles are prepared by the treatment of alkyl halides with
 - a) Sodium cyanide
- b) Sodium isocyanide
- c) Sodium isocyanate
- d) Cyanamide

- 220. Urea is not used:
 - a) As fertilizer
 - b) In manufacture of plastic
 - c) In preparation of medicines
 - d) In purification of water
- 221. When aniline is heated with glacial acetic acid in presence of anhydrous ZnCl₂, the product is:
 - a) Acetamide
- b) Acetanilide
- c) Phenyl acetamide
- d) Chlorobenzene
- 222. Benzene diazonium chloride on treatment with hypo phosphorous acid and water in presence of Cu⁺ catalyst produce
 - a) Benzene
- b) Toluene
- c) Aniline
- d) Chlorobenzene
- 223. Which of the following cannot be used for following conversion?
 - $CH_3CN \rightarrow CH_3CH_2NH_2$

	a) Nitrobenzene	b) LiAlH ₄ ce formed by the action of a b) Phenyl isocyanide	c) Phenyl cyanide	d) SnCl ₂ /HCl chloroform and aniline is d) Phenyl isocyanate
	a) 1°	I_5SO_2Cl and the product is s b) 2°	soluble in alkali, amine is: c) 3°	d) All of these
226.	Which of the following re-	action will not occur?		
	a) $\phi N_2^+ + CuBr \xrightarrow{HBr} \phi - F$	3r	b) $\phi N_2 + \phi OH \longrightarrow \phi -$	$-N=N-\langle \bigcirc \rangle$ OH
	c) $\phi N_2^+ \xrightarrow[HOH]{HOH} \phi H$		d) $\phi N_2^+ + I^- \rightarrow \phi - I$	
227.	The end product in the be			
	$C_2H_5NH_2 \xrightarrow{HNO_2} A \xrightarrow{PCl_5} B \xrightarrow{N}$	$\stackrel{n_3}{\to} \mathcal{C}$		
	a) Ethyl cynide	, ,	c) Methyl amine	d) Acetamide
228.	Among the amines $(A)C_6$ I $(CH_3)_2$ NH, $(D)(CH_3)_3$ N, t		4	21
	a) $A < B < D < C$. (4	
	b) $D < C < B < A$			
	c) $A < B < C < D$			
	d) $B < C < D < A$			
229.	= = = = = = = = = = = = = = = = = = =	reaction with HNO ₂ gives a		DAI CM
220	a) 11.2 L of N ₂	b) 5.6 L of N ₂	c) 22.4 L of N ₂	d) 1 L of N ₂
230.		with chloroform and ethan	A 1/1	
224	a) Benzyl alcohol	b) Benzaldehyde	c) Benzonitrile	d) Benzyl isocyanide
231.	Which nitro compound w		CH CH NO	15
222	a) C ₆ H ₅ NO ₂	b) (CH ₃) ₃ CNO ₂	c) CH ₃ CH ₂ NO ₂	d) <i>o</i> -nitrotoluene
Z3Z.	Benzamide can be conver		a) IZCN	4) D O
222	a) H ₃ 0 ⁺	b) 0H ⁻ /H ₂ 0	c) KCN	d) P_2O_5
233.	Choose the incorrect state	ement condary and tertiary amine	s the basic strength denon	ds on the extent on the
	extent of hydrogen bor	nding in the protonated am	ines	
		s like – OCH_3 and – CH_3 incr		
		s like – NO_2 , –CN and halog		gth of amines
	d) The basic strength of a	mines depends on their cor	ncentration	
234.	Nitrosoamines (R_2 N – N secondary amines. The re	= 0) are soluable in water. action is called	On heating them with cond	CH ₂ SO ₄ , they give
	a) Perkin's reaction		b) Fries reaction	
	c) Liebermann nitroso re	action	d) Etard reaction	
235.		not used for nitration of org		
	a) A mixture of concentra H ₂ SO ₄	ted HNO ₃ and concentrated	d b) A mixture of concentra anhydride	ted HNO ₄ and acetic
	c) Fuming nitric acid and	concentrated sulphuric	d) Alcoholic potassium nit	trate
	acid			
236.	m-fluoronitrobenzene is l	best synthesized by using tl	ne reaction	
	a) Nitrobenzene $\frac{\text{Fuming HI}}{\text{H}_2\text{SO}_4,\text{he}}$	$\underset{\text{at}}{\overset{\text{NO}_3}{\longrightarrow}} \left[\right] \xrightarrow{\underset{\text{NH}_3/\text{H}_2S}{\text{NH}_3/\text{H}_2S}} \left[\right] \xrightarrow{\text{1.HONO}}$	b) Aniline $\xrightarrow{F_2}$ heat	
	c) Fluorobenzene $\frac{HNO_3}{H_2SO_4,he}$	\rightarrow	d) m -C ₆ H ₄ (NH ₂) ₂ $\frac{1.\text{HO}}{2.\text{CuNO}_2}$	NO
227			_	3.HBF ₄
۷3/.	The correct order of basic	cities of the following comp	ounus is:	



- a) 2 > 1 > 3 > 4
- b) 1 > 3 > 2 > 4
- c) 3 > 1 > 2 > 4
- d) 1 > 2 > 3 > 4
- 238. Arrange the following compounds in increasing order of basic strength



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

239. In the reaction

$$R - C \equiv N + 4(H) \xrightarrow{X} RCH_2NH_2$$

X can be

- a) LiAIH₄
- b) H₂SO₄
- c) Ni

- d) 2KBr
- 240. p-chloro aniline and anilinium hydrogen chloride can be distinguished by
 - a) Sandmeyer reaction

b) Carbylamines reaction

c) Hinsberg's reaction

d) AgNO₃

241. In the chemical reaction,

$$CH_3CH_2NH_2 + CHCI_3 + 3KOH \rightarrow (A) + (B) + 3H_2O$$

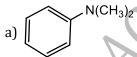
The compounds (A) and (B) are respectively

a) CH₃CH₂CONH₂ and 3KCI

b) C₂H₅NC and K₂CO₃

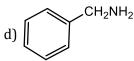
c) C₂H₅NC and 3KCI

- d) C₂H₅CN and 3KCI
- 242. The reagent that reacts with nitromethane to form methyl hydroxylamine is
 - a) Zn/HCI
- b) Zn/NH₄CI
- c) Zn/NaOH
- d) Sn/HCI
- 243. The compound which on reaction with cold HNO₂ gives only nitrosoamine is:
 - a) CH₃NH₂
- b) (CH₃)₂NH
- c) $(CH_3)_3N$
- d) $(C_2H_5)_3N$
- 244. Amongst the compound given, the one that would form a brilliant coloured dye on treatment with $NaNO_2$ in dil. HCI followed by addition to an alkaline solution of β —naphthol is



b) (N

c) H₃C



- 245. Primary, secondary and tertiary amines may be separated by using:
 - a) Ethanoyl chloride
- b) Diethyl oxalate
- c) Thionyl chloride
- d) None of these

246. Consider the following reaction,

$$C_6H_5NO_2 \xrightarrow{Sn/HCI} X \xrightarrow{C_6H_5COCI} Y + HCI$$

What is Y?

- a) Acetanilide
- b) Benzanilide
- c) Azobenzene
- d) Hydrazobenzene
- 247. Nitration of aniline in strongly acidic medium, result in the formation of *m*-nitroaniline also. This is because
 - a) Amino group is *meta* orienting during electrophilic substitution reaction.
 - b) Nitro group goes always to the *meta* position irrespective of the substituents.
 - c) Nitration of aniline is a nucleophilic substitution reaction in strongly acidic medium.
 - d) In strongly acidic conditions aniline is present as anilinium ion.
- 248. The major product of the following reaction is

a)
$$N - H_2C - Br$$

c)
$$N$$
 C
 $O-CH_2$
 Br

b)
$$C$$
 N CH_2CI

d)
$$N - H_2C - CH_2CI$$

- 249. Which of the following compounds is soluble in benzene but almost insoluble in water?
 - a) C₂H₅OH
- b) CH₃CO₂H
- c) CH₃CHO
- d) $C_6H_5NO_2$

250. H₂NH₂C

$$\frac{\text{HNO}_2}{\text{Product is}}$$
?



251. $CHCI_3 + C_6H_5NH_2 + 3NaOH \rightarrow A + 3B + 3C$

In the above reaction, the product 'A' is

- a) Chlorobenzene
- b) Phenyl isocyanide
- c) Phenyl cyanide
- d) Phenyl chloride

- 252. Alkyl cyanides undergo Stephen reduction to produce
 - a) Aldehyde
- b) Secondary amine
- c) Primary amine
- d) amide

- 253. Which of the following is not a nitro derivative?
 - a) C₆H₅NO₂
- b) CH₃CH₂ONO
- c) CH₃-CH-N CH₃
- d) $C_6H_4(OH)NO_2$

254.

 $\begin{array}{c|c} CH_2CH_2NH_2HCI\\ \text{In the reaction,} \end{array}$

 $\stackrel{\Delta}{\longrightarrow}$ Product:

CH2CH2NH2HCl

The product is:

b)
$$\begin{bmatrix} NH_2 \\ NH_3 \end{bmatrix}$$

b) N of CN group

d)
$$\langle \bigcirc \rangle$$
 $N(CH_3)_3$

- 255. In alkyl cyanide alkyl group attached with
 - a) C of CN group
 - c) Either C or N of CN group
- 256. The diamide of carbonic acid is:
 - a) Acetamide
- b) Formamide
- c) Benzamide
- d) Urea

- 257. A positive carbylamines test is given by
 - a) N, N-dimethylaniline

b) 2,4-dimethylaniline

d) Both C and N of CN group

c) N-methy-o -methylaniline

- d) N-methylbenzylamine
- 258. Which of the following amines can be directly oxidized to the corresponding nitro compound by potassium permanganate?



$$\begin{array}{c|c} \operatorname{CH}_2 - \operatorname{CH} - \operatorname{CH}_3 \\ \text{b)} & | \\ \operatorname{NH}_2 \end{array}$$

c)
$$(CH_3)_2NH$$

d)
$$(CH_3)_3C - NH_2$$

259. Arrange the following CH_3NH_2 (I); CH_3NH (II); $C_6H_5NH_2$ (III); $(CH_3)_3N$ (IV) in increasing order of basic nature in aqueous medium:

a)
$$II < I < IV < III$$

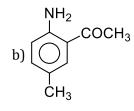
b)
$$III < IV < I < II$$

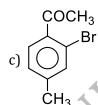
c)
$$I < II < III < IV$$

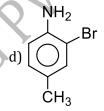
d)
$$II < III < I < IV$$



The final product 'C' in the above reaction is







261. Identify X in the series,





$$\frac{\text{HNO}_3}{\text{H}_2\text{SO}_4} \text{Intermediate} \xrightarrow{\text{H}_2\text{O}} X$$

a)
$$\bigcirc$$
 $\stackrel{NH_2}{\bigcirc}$

$$NH_2$$

$$O_2N$$
 NH_2
 NO_2

262. In the reaction sequence

The product 'C' is

- a) Benzonitrile
- b) Benzaldehyde
- c) Benzoic acid
- d) Benzyl amine

263. Nitroalkane is acidic only towards:

- a) Na₂CO₃
- b) NaOH
- c) Alcohol
- d) Liquid NH₃

264. Urea reacts with hydrazine to form:

- a) Nitrogen
- b) Phenyl hydrazine
- c) Semicarbazide
- d) Urethane

265. Phenyl cyanide cannot be obtained by

a)
$$C_6H_5CONH_2 \xrightarrow{P_2O_5,\Delta}$$

b)
$$C_c H_r - CH = NOH \xrightarrow{Ac_2 O, \Delta}$$

c)
$$C_6H_5Cl \xrightarrow{alc.KOH}$$

b)
$$C_6H_5 - CH = NOH \xrightarrow{Ac_2O,\Delta}$$

d) $C_6H_5NH_2 \xrightarrow{1.NaNO_2/HCl}$

266. Substitution of one alkyl group by replacing hydrogen of primary amines:

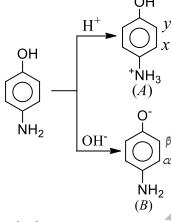
- a) Increases the base strength
- b) Decreases the base strength
- c) Remains the same
- d) None of the above

267. Acetanilide is prepared by the reaction of acetyl chloride on:

- a) Acetamide
- b) Aniline
- c) Acetaldehyde
- d) Benzene

- 268. Aqueous solution of urea is:
 - a) Acidic
- b) Alkaline
- c) Almost neutral
- d) Amphoteric

269. Consider *p*-aminophenol



Which positions are activated for coupling reaction in acidic and basic media respectively?

a) x in A and β in B

a) A diazonium salt

- b) x in A and α in B
- c) y in A and α in B
- d) y in A and β in B

270. The general formula of quaternary ammonium compound is:

- a) R—NH₂
- b) R_3N

- c) $[R_4N]^+X^-$
- d) NH_4X

271. Reaction of nitrous acid on 1° aliphatic amines in cold will give:

- b) An alcohol
- c) A nitrile
- d) A dye

272. In pyridine, the state of hybridization of the nitrogen atom is

a) sp^2

b) sp^3

d) None of these

273. Which of the following will give a primary amine on hydrolysis?

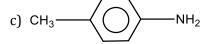
- b) Alkyl cyanide
- c) Oxime
- d) Alkyl isocyanate

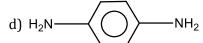
274. Which of the following compounds will form alcohol on treatment with NaNO₂, HCI/H₂O at 0° C?

 \sim a) (CH₃)₂CHNH₂

a) Nitroparaffin

b) $C_6H_5NH_2$





275. Which of the following statements is correct?

- a) Aniline is stronger base than ammonia
- b) Methylamine is a stronger base than aniline and ammonia
- c) Aniline is stronger than ammonia, but weaker base than methylamine

d) Methylamine is strong	d) Methylamine is stronger than aniline, but weaker base than ammonia									
276. Benzenediazonium chlor	ride on reaction with pheno	l in weakly basic medium g	ives							
a) Diphenyl ether		b) <i>p</i> -hydroxyazobenzene								
c) Chlorobenzene		d) benzene								
277. Which of the following m	nethods neither means for the	he synthesis nor for separa	tion of amines?							
a) Hinsberg's method	b) Hofmann's method	c) Wurtz reaction	d) Curtius method							
278. Which substance when b		•	· , ·····							
a) Ethylamine	b) Aniline	c) Acetamide	d) Acetoxime							
279. Acetonitrile on reduction	•	,								
a) Propanamine	b) Methanamine	c) Ethanamine	d) None of these							
280. When ethanol is mixed w	•	•								
a) C ₂ H ₅ NH ₂	b) C ₂ H ₄	c) C ₂ H ₅ OC ₂ H ₅	d) CH ₃ OCH ₃							
281. The molecular formula o		-) -2323	.,,,							
a) C ₆ H ₅ CN	b) C ₆ H ₅ NC	c) C ₆ H ₅ CNO	d) C ₆ H ₅ NCO							
282. Which of the following a										
a) CH ₃ NH ₂	b) (CH ₃) ₂ NH	c) $(CH_3)_3N$	d) None of these							
283. The correct order of the										
a) Methylamine < anilin		netnyr annine, anninoma ant	diffiffic 15.							
b) Aniline < ammonia <										
c) Aniline < methylamin	•									
d) Ammonia < aniline <										
284. Diazotisation can be carr		NO and dilute HCl at ice co	ld tamparatura on:							
a) Aromatic secondary a	-	NO ₂ and unute fici at ice co.	iu temperature on.							
b) Aromatic primary am										
c) Aromatic nitro compo	ulia									
d) Aromatic amine	hadia than NII hut around	ia aminag aya hagia tha	NII							
285. Aliphatic amines are										
a) More, less	b) Less, more	c) Both (a) and (b)	d) None of these							
286. Aniline is weaker base th		9.1.1. 6								
	of N-atom is not freely avai	llable for coordination with	a proton due to resonance							
than in ethylamine										
b) Its b. p. is higher than	•	* = 1								
•	ifficient concentration of OI									
-	while ethylamine is solubl									
287. The basic character of m										
a) 3°>2°>1°>NH ₃	5	c) $1^{\circ}>2^{\circ}>3^{\circ}>NH_3$	d) None of these							
288. Isopropylamine $\xrightarrow{\text{KMnO}_4} X$	$Y \xrightarrow{H_3O^+} Y$. In the above seque	ence X and Y are respectivel	y							
a) Acetaldimine, ethanal		b) Ethanal, ketimine	•							
c) Ketimine, acetone		d) Acetone, propan-2-ol								
289. Which of the following co	ompound does not undergo		tion?							
a) Phenol	b) Primary amine	c) Secondary amine	d) Tertiary amine							
290. Production of amines by	-	-	a) rereary among							
a) Frankland reaction		b) Hofmann's ammonolys	sis							
c) Hofmann's mustrard	oil reaction	d) Hofmann's bromamide								
291. Which of the following is		a, mondani o oromanina								
a) CH ₃ CONH ₂	b) NH ₂ CONH ₂	c) CH ₂ (NH ₂)CONH ₂	d) CO(OH)NH ₂							
292. Aniline reacts with conc.	·	o, onz(miz)oomiz	as do (onsimiz							
	1111177 117 EIVL									

b)
$$NH_2$$
 NO_2 and NO_2

$$d) \overbrace{\hspace{1cm}}^{\hspace{1cm} NO_2}$$

293. The correct structure of ethylenediamine-tetra acetic acid (EDTA) is:

294. Hydrazobenzene $\xrightarrow{\text{NaIO}_3}$ $(X) \xrightarrow{\text{CH}_3\text{CO}_3\text{H}} (Y)$

Both *X* and *Y* on reduction with Sn/HCl give *Z*. Which of the following does not represent *X*, *Y* or *Z*?

- a) Azobenzene
- b) Phenol

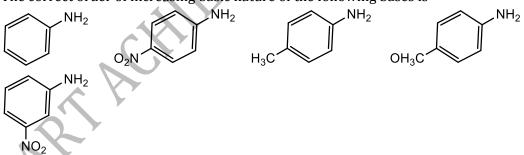
CH₂COOH

- c) Aniline
- d) Azoxybenzene

- 295. The pri., sec. and ter. amines can be distinguished by:
 - a) Hinsberg's reagent
- b) Grignard reagent
- c) Fehling's solution
- d) Tollen's reagent

- 296. Final product of hydrolysed alkyl cyanide is
 - a) RCOOH
- b) RCONH₂
- R C = NHc) | OH
- d) R C = NH

297. The correct order of increasing basic nature of the following bases is



a) II < V < I < III < IV b) V < II < I < III < IV c) II < V < I < IV < III d) V < II < I < IV < III 298. Which of the following is the strongest base?

a)
$$\langle \bigcirc \rangle$$
—NH₂

d)
$$\langle \bigcirc \rangle$$
 — CH_2 — NH_2

299. The basicity of compounds I, II, III and IV

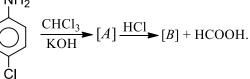
 CH_3NH_2 , $(CH_3)_2NH$, $(CH_3)_3N$, $C_6H_5CH_2NH_2$

- varies in the order
- III
- II IV
- a) I > II > III > IV
- b) II > I > III > IV
- c) III > I > II > IV
- III < II < I > II

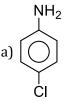
300. Which one of the following does not have sp^2 hybri	dised carbon?	
a) Acetone b) Acetic acid	c) Acetonitrile	d) Acetamide
301. The basic character of amines can be explained:		
a) In terms of Lewis and Arrhenius concept		
b) Only in terms of Lowry Bronsted concept		
c) It terms of Lewis and Lowry Bronsted concept		
d) Only in Lewis concept		
302. In the compound given below,		
H_3 N H_3 (Z)		
СООН		\ \'
(X)		A Y
the correct order of acidic nature of the positions (, , , , , ,	
a) $Z > X > Y$ b) $X > Y > Z$	c) $X > Z > Y$	d) Y > X > Z
303. KCN reacts readily to give a cyanide		
a) Ethyl alcohol b) Ethyl bromide	c) Bromobenzene	d) chlorobenzene
304. A colourless organic compound gave brisk efferves)" -
a) Glucose b) Oxalic acid	c) Urea	d) Benzoic acid
305. Which of the following reactions can be used to pre		
a) $CH_3CH_2I + NaCN \xrightarrow{C_2H_5OH/H_2O}$	b) $CH_3CH_2I + KCN \frac{Alcoh}{\Lambda}$	ol →
	d) None of the above	
c) $CH_3CH_2NH_2 + CHCL_3 + KOH \xrightarrow{Alcohol}$	u) None of the above	
306. Diethyl oxalate is used for distinguishing primary, s	secondary and tertiary	
a) Alcohols	b) Amines	
c) Alkyl halides	d) Hydrogens in hydroc	arbons
307. Identify <i>X</i> in the sequence,	Y	
$X \xrightarrow{\text{HNO}_2} \text{C}_3\text{H}_8\text{O} \xrightarrow{\text{H}_2\text{SO}_4} \text{C}_3\text{H}_6\text{O}_2$:		
a) CH ₃ —NH—CH ₂ —CH ₃		
b) CH ₃ —CH ₂ —CH ₂ —NH ₂		
c) (CH ₃) ₃ N		
d) None of the above		
$308. \text{ CH}_3\text{CONH}_2 + \text{NaOH} \rightarrow \text{CH}_3\text{COONa} + A$		C 11
Urea is obtained if product if product 'A' in the above		
a) Ethyl carbonate b) Ethyl urethane	c) Phosgene	d) All of these
309. Which of the following is involved in Sandmeyer's r	eaction?	
a) Ferrous salt		
b) Diazonium salt		
c) Ammonium salt		
d) Cupraammonium salt		
310. <i>R</i> Mg <i>X</i> on reacting with cyanogen chloride gives:		
a) R—NC b) R—Cl	c) R—CN	d) None of these
311. Methyl amine reacts with methyl iodide. For compl	etion of reaction, how mar	ny moles of methyl iodide are
required?		
a) 1 b) 2	c) 3	d) 4
312. Aniline on oxidation with $Na_2Cr_2O_7$ and H_2SO_4 give		
a) Benzoic acid b) <i>m</i> -amino benzoic acid	c) Schiff's base	d) <i>p</i> -bezoquinone
313. Among the following the weakest base is		
a) C ₆ H ₅ CH ₂ NH ₂ b) C ₆ H ₅ CH ₂ NHCH ₃	c) $O_2NCH_2NH_2$	d) CH ₃ NHCHO
314. Why do 2° and 3° amines fail to undergo the carbyla	amines test?	

- a) They combine with chloroform to give a stable compound
- b) They react with alcoholic KOH
- c) They nitrogen atom of the amine group does not have the required number of hydrogen atoms
- d) All the given reasons are correct
- 315. The compound that will react most readily with NaOH to form methanol is
 - a) $(CH_3)_4N^+I^-$
- b) CH₃OCH₃
- c) $(CH_3)S^+I^-$
- d) $(CH_3)_3CCl$
- 316. Alkanamide, which on Hofmann's reaction gives 1-phenylethylamine, is:
 - a) 2-phenylpropanamide
 - b) 3-phenylpropanamide
 - c) 2-phenylethanamide
 - d) N-phenylethanamide
- 317. Which of the following compounds is expected to be most basic?
 - a) Aniline
- b) Ethylamine
- c) Hydroxylamine
- d) Methylamine

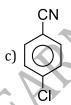
318. NH₂

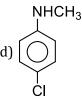


What is [B]?









- 319. Aniline is not the major product in one of the following reactions. Identify that reaction.
 - a) $C_6H_5OH + NH_3 \xrightarrow[300^{\circ}C]{ZnCI_2}$
 - C₆H₅Ci + NH₃ $\frac{200^{\circ}\text{C}}{\text{Cu}_{2}\text{O}}$ High pressure
- b) $C_6H_5NO_2 + Zn$ powder $\xrightarrow{Alcoholic KOH}$

d)
$$C_6H_5NO_2 + 6(H) \xrightarrow{\text{Fe+H}_2O} HCI$$

- 320. In the reaction between CH₃NC and HgO, the product obtained is
 - a) Methyl isothiocyanate

b) Methyl isocyanate

c) Methyl amine

d) Methyl cyanide

321. Complete the following reaction

$$RNH_2 + H_2SO_4 \rightarrow$$

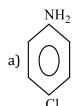
- a) $[RNH_3]^+HSO_4^-$
- **b)** $[RNH_3]_2^+SO_4^{2-}$
- c) RNH_2 . H_2SO_4
- d) No reaction
- 322. Which one of the following is the strongest base in aqueous solution?
 - a) Trimethylamine
- b) Aniline
- c) Dimethylamine
- d) Methylamine

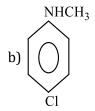
- 323. Nitrobenzene on reduction with Al-Hg and water gives:
 - a) Azobenzene
 - b) Aniline
 - c) Azoxy benzene
 - d) phenylhydroxylamine
- 324. Gabriels phthalimide reaction is used to prepare:
 - a) p –amine
- b) s amine
- c) t amine
- d) All of these

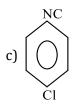
325. Identify *X* in the reaction,

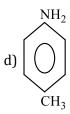


 $\xrightarrow{\text{CHCl}_3, \text{KOH}} \text{Intermediate } \xrightarrow{\text{HCl}} X:$

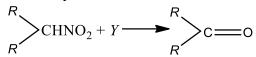








326. Secondary nitroalkanes can be converted into ketones by using Y. Identify Yfrom the following



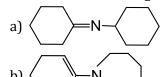
- a) Aqueous HCl
- b) Aqueous NaOH
- c) KMnO₄
- d) CO

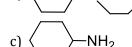
- 327. The strongest base among the following is
 - a) $C_6H_5NH_2$
- b) $(C_6H_5)_2NH$
- c) NH₃

d) $(C_2H_5)_2NH$

- 328. Alkyl nitrite on reduction with Sn/HCl gives:
 - a) Alcohol
- b) Hydroxylamine
- c) Both (a) and (b)
- d) hydrazine

329. Which of the following is an enamine?





$$d$$
) $N=C=N-$

- 330. The number of π -bonds in the formula given below, NC—CH=CH—CN are:
 - a) 5

b) 4

c) 3

d) 2

- 331. Which of the following is most basic in nature?
 - a) NH.

- b) CH₃NH₂
- c) $(CH_3)_2NH$
- d) $C_6H_5N(CH_3)_2$

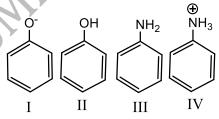
- 332. Diazomethane reacts with carboxylic acids to produce:
 - a) Ester

- b) Alcohol
- c) Amine
- d) Imines

- 333. Which compound is known as alkyl carbylamines?
 - a) R. CN
- b) R. NC
- c) Ar. CN
- d) Ar. NC
- 334. n-propylamine yields a volatile compound X on warming with alc. alkali and chloroform. X has an offensive odour. The structure of X is
- a) CH₃CH₂CH₂CN
- b) (CH₃)₂CHCN
- c) CH₃CH₂CH₂NC
- d) (CH₃)₂CHNC
- 335. The reaction of CHCI₃ and alcoholic KOH with *p*-toluidine gives

c)
$$H_3C$$
 NC

336. Coupling of diazonium salts of following takes place in the order



- a) IV < II < III < I
- b) IV > III < II < I
- c) II < IV < I < III
- d) I < II < III < IV

- 337. Tertiary nitroalkane cannot tautomerise because
 - a) Their tautomeric forms are highly unstable
 - c) They do not have labile H-atom

- b) They do not contain any multiple bond
- d) They are not basic in nature

338.	In aqueous solutions, th	e basic strength of amines d	ecreases in the order				
	a) $CH_3NH_2 > (CH_3)_2NH_3$	$I_2 > (CH_3)_3 N$	b) $(CH_3)_2NH > (CH_3)_3N$	$> CH_3NH_2$			
	c) $(CH_3)_3N > (CH_3)_2NH$	$H > CH_3NH_2$	d) $(CH_3)_2NH_2 > CH_3NH_2 > (CH_3)_3N$				
339.	Dichlorocarbene does n	ot form as an intermediate i	n this reaction				
	a) $phenol + CHCl_3 + 4K$	ОН	b) Ethyl amine +CHCl ₃ + KOH				
	c) Phenol+CCl ₄ + 4KOH	I	d) CHCl ₃ + KOH				
340.	Which of the following is	s not a nitroderivative?					
			ρ				
	a) C ₆ H ₅ NO ₂	b) CH ₃ CH ₂ ONO	c) $\begin{array}{c} \text{CH}_3 \text{ CH} - \text{N} \\ \\ \text{CH}_3 \end{array}$	d) C ₆ H ₄ (OH)NO ₂			
341.	Urea reacts with HNO ₃ t	o give:					
	a) Urea nitrite	b) Urea nitrate	c) H_2CO_3	d) None of these			
342.	Which of the following r	eagents will convert nitrom	ethane into methylamine?	4			
	a) Zn/HCl	b) Zn/NaOH	c) Zn/C ₂ H ₅ OH	d) Ni/H ₂			
343.	NH_2			0 1			
	(i) (CH ₂ CO) ₂ O (ii)	HOSO ₂ C1					
	(i) (CH ₃ CO) ₂ O (ii) (iii) RNH ₂ (iv) dil	[X];	4 4				
	~	Here A IS	W 10 0 0 (1)	NILLI			
	NHCOCH ₃	NH ₂	NHCOCH3	NH ₂ 			
	a) ()	b) [()	c) (d) [()			
	Y H) 00 NUID	CO KILID	Y			
	SO₃H	ŚO₂NH <i>R</i>	ŚO₂NH <i>R</i>	ŚO₃H			
344.	In which reaction, nitre	ne is not the intermediate?	V , y				
	a) Schmidt		b) Curtius				
	c) Hofmann bromamide		d) Gabriel's phthalimide				
345.	CH ₃ CONH ₂ . Br ₂ and KOI	H give CH ₃ NH ₂ as the produc	ct. The intermediates of the	reaction are			
	O						
	(A) CH ₃ —C—NHBr	$B) CH_3 - N = C = 0$					
	(C)CH ₃ NHBr	(D) CH_3CONBr_2					
	The correct answer is	(D) Cli3CONDI2					
	a) A, B						
346	aj n, D	h) / (c) C D	4) B D			
340.	The HIDAC name of the	b) A, C	c) C, D	d) B, D			
		b) A, C compound having formula,	c) C, D	d) B, D			
	$O = C - CH - CH_2$ is:		c) C, D	d) B, D			
			c) C, D	d) B, D			
	$O = C - CH - CH_2$ is:	compound having formula,	c) C, D	d) B, D			
	$ \begin{array}{c c} O = C - CH - CH_2 \text{ is:} \\ & & \\ OH \text{ NH}_2 \text{ OH} \end{array} $	compound having formula,	c) C, D	d) B, D			
	$O = C - CH - CH_2$ is: $\begin{vmatrix} & & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & \\ & & & \\ & & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & \\ & & & \\ & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\$	compound having formula, pionic acid c acid	c) C, D	d) B, D			
	O=C-CH-CH ₂ is:	compound having formula, pionic acid c acid anoic acid	c) C, D	d) B, D			
347.	O=C-CH-CH ₂ is:	compound having formula, pionic acid c acid anoic acid					
347.	O=C-CH-CH ₂ is:	compound having formula, pionic acid c acid anoic acid ropanoic acid ound reacts with aqueous ni					
347.	O=C-CH-CH ₂ is:	compound having formula, pionic acid c acid anoic acid ropanoic acid ound reacts with aqueous ni					
347.	O=C-CH-CH ₂ is:	compound having formula, pionic acid c acid anoic acid ropanoic acid ound reacts with aqueous ni	trous acid at low temperat b) CH3CH2NH2				
5	O=C-CH-CH ₂ is:	compound having formula, pionic acid anoic acid ropanoic acid bund reacts with aqueous ni pound is	trous acid at low temperat				
5	O=C-CH-CH ₂ is:	compound having formula, pionic acid anoic acid ropanoic acid bound reacts with aqueous ni bound is	trous acid at low temperat b) ${\rm CH_3CH_2NH_2}$ d) ${\rm CH_3CH_2-NHCH_2CH_3}$	ure to produce an oily			
348.	O=C-CH-CH ₂ is:	compound having formula, pionic acid c acid anoic acid ropanoic acid bund reacts with aqueous ni bound is by: b) Carbohydrates	trous acid at low temperat b) CH3CH2NH2				
348.	O=C-CH-CH ₂ is:	compound having formula, pionic acid c acid anoic acid ropanoic acid ound reacts with aqueous ni pound is by: b) Carbohydrates mpounds, the most basic is	trous acid at low temperat b) $CH_3CH_2NH_2$ d) $CH_3CH_2 - NHCH_2CH_3$ c) Polypeptides	ure to produce an oily d) Urea			
348. 349.	O=C-CH-CH ₂ is:	compound having formula, pionic acid c acid anoic acid ropanoic acid ound reacts with aqueous ni pound is by: b) Carbohydrates mpounds, the most basic is b) Acetanilide	trous acid at low temperat b) ${\rm CH_3CH_2NH_2}$ d) ${\rm CH_3CH_2-NHCH_2CH_3}$	ure to produce an oily			

351. When (NH ₄) ₂ SO ₄ +	KCNO are heated, we get:		
a) Nitrogen			
b) Carbon dioxide			
c) Biuret			
d) Ammonium carbo	nate		
352. Grignard reagent and	d acetyl chloride does not	react with:	
a) RNH ₂	b) R ₂ NH	c) R_3N	d) None of these
353. Acetaldoxime reacts	s with P_2O_5 to give:		
a) CH ₃ CN	b) C ₂ H ₅ CNO	c) C ₂ H ₅ CN	d) All of these
354. 2,4,6-tribromo anilir	ne is a product of:		\mathcal{L}
a) Electrophilic addi	-		
	stitution on C ₆ H ₅ NH ₂		
c) Nucleophilic addi			A . Y
d) Nucleophilic subs	-		41
355. Choose the incorrect	° ° -		
CH ₃	, comparion (c)		
-	CHNHCH.	CH CH CH NH	COUL
$H_3C-C-NH_2 < a$	CII3—INFICII3	b) CH ₃ CH ₂ CH ₂ NH ₂	
CH ₃		(basicity in aque	eous medium)
(basicity in aque	eous medium)		
CTT OIL NILL	CIT NILIOLI OLI	3 N - 1 1 1 1 1 1	21/2
CH_3 - CH - NH_2 <	CH ₃ -NHCH ₂ CH ₃	2 $N(1)$	>N(3)
c) CH ₃		d) IN	
(basicity in the	gaseous state)		(basicity in aqueou
356. Identify the major pr	roduct of the reaction	medium)	
550. Identity the major pr	oduct of the reaction		
$\left[\begin{array}{c} + \end{array}\right] \qquad \stackrel{\text{OH}}{\longrightarrow}$	C	Y	
N CH ₃			
CH ₃ CH ₃		7	
	h)	C) L CH3	d) L. LCH3
a) $N \subset CH_3$	b) N CH ₂	o) N	d) N —CH ₃
CH ₃ CH ₃	H ₃ C´CH ₃	H ₃ C´\CH ₃	H ₃ C´ `CH ₃
357. Which of the following	ng compounds gives a sec	ondary amine on reduction	n?
a) Nitromethane	b) Nitrobenzene	c) Methyl isocyani	de d) Methyl cyanide
358. Which of the following		s the basic for a simple ch	emical test to distinguish
between?		•	<u> </u>
	1		
$H_3C - \langle O \rangle - NH_2$	and $\langle \bigcirc \rangle$ — CH_2NH_2		
a) C ₆ H ₅ SO ₂ Cl and OI	H ⁻ in H ₋ O	b) HONO, then β-n	anhthol
c) Dilute HCl	1 1111120		αριταίοι
		d) AgNO ₃ in H ₂ O	
359. Amine may contain:	_	_	4) All = 641 :
a) —NH ₂ gp	b) NH gp.	c) \rightarrow N gp.	d) All of these

of the excess of an acid but reappears if the solution is made alkaline. The aliphatic nitro compound is a) $CH_3CH_2NO_2$ b) $(CH_3)_2CHNO_2$ c) $(CH_3)_3CNO_2$ d) All of these 362. In the following reaction sequence predict the compound X and Y.

361. An aliphatic nitro compound turns red with the addition of a concentrated NaOH solution, followed by the addition of an excess of an $NaNO_2$ solution and then dilute H_2SO_4 . The colour disappears with the addition

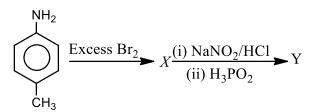
c) Tetraethyl hydrazine

360. Diethylamine on oxidation with $KMnO_4$ gives:

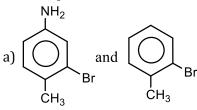
b) Propanone

a) Ethanal

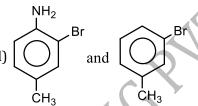
d) None of these



The compound X and Y are



c)
$$Br$$
 Br Br Br Br Br Br



- 363. Primary amine (RNH₂) reacts with nitrous acid to give
 - a) $RNH_3^+NO_2$
- b) ROH

c) ROR

- d) None of these
- 364. Carbylamine reaction tubes are not thrown into sink, to avoid bad odour, but are treated with conc. HCl to give:
 - a) $RCOOH + NH_3$
- b) RNH₂
- c) $RNH_2 + HCOOH$
- d) $RCOOH + N_2$
- 365. The compound obtained by heating a mixture of 1° amine and chloroform with ethanolic potassium hydroxide is
 - a) An alkyl isocyanide

b) An alkyl isothiocyanate

c) An amide

- d) An amide and nitro compound
- 366. The best method to synthesise *m*-dibromobenzene is by using the reaction
 - a) Benzene $\frac{Br_2/FeBr_3/heat}{}$

b) Aniline $\xrightarrow{Br_2,H_2O}$ [] $\xrightarrow{1.HONO}$

Nitrobenzene

- C) $\frac{\text{Fuming HNO}_3}{\text{H}_2\text{SO}_4\Delta}$ [] $\frac{\text{Fe/HCl}}{\text{C}_2\text{H}_5\text{OH,heat}}$
- Bromobenzene d) $\xrightarrow{\text{HNO}_3}$ [] $\xrightarrow{\text{Fe/HCl}}$ [] $\xrightarrow{\text{1.HONO}}$ 2.CuBr

367. The main product in the reaction,

HCONHR
$$\xrightarrow{\text{POCl}_3}$$
 is:

a) RCN

b) RNC

- c) RCNO
- d) RNCO

368. The type of isomerism shown by C₆H₅CN

and C_6H_5NC is:

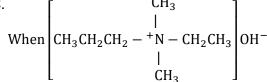
- a) Position
- b) Functional
- c) Enantiomerism
- d) Tautomerism

- 369. Which among the following has the highest boiling point?
 - a) CH₃CH₂CH₂NH₂
- CH₃CH₂-NH CH₃
- c) CH_3 CH_3 -N $-CH_3$
- d) CH₃NH₂

- 370. Benzaldehyde reacts with methyl amine to give:
 - a) $C_6H_5NH_2$
- b) C₆H₅CH₂NH₂
- c) $C_6H_5CH = NCH_3$
- d) C₆H₅CONH₂

- 371.
 - $\xrightarrow{\text{NH}_2\text{OH}} X \xrightarrow{\text{LiAlH}_4} Y. \text{ In the above sequence } Y \text{ is}$
 - a) Tertiary amine
- b) Secondary amine
- c) Primary amine
- d) 2-nitropropane
- 372. Ethylamine undergoes oxidation in the presence of KMnO₄ followed by hydrolysis to form:
 - a) An acid
- b) An alcohol
- c) An aldehyde
- d) a N-oxide

373.



Is heated, then

a) Propene is the major product

- b) Ethane and C₃H₇N(CH₃)₂ are the only product
- c) Ethane and propene are obtained while ethane as d) Equimolar amounts of ethane and propene are the major product obtained
- 374. Diethyl carbonate on heating with ammonia gives:
 - a) $C_2H_5NH_2$
 - b) $(C_2H_5)_3N$
 - c) $(C_2H_5)_2NH$
 - d) Urea
- 375. In which case formation of butane nitrile is possible?
 - a) $C_3H_7Br + KCN$
- b) $C_4H_9Br + KCN$
- c) $C_3H_7OH + KCN$
- d) $C_4H_9OH + KCN$

- 376. Ethyl amine reacts with nitrous acid to form
 - a) C₂H₅OH
- b) C_2H_5OH , N_2 , H_2O
- c) $C_2H_5N_2^+CI^-$
- d) C₂H₅NHOH, NH₃
- 377. Which of the following compounds will undergo carbylamine reactions?
 - a) $(CH_3CH_2)_2NH$
- b) (CH₃)₂NH
- c) $C_6H_5NH_2$
- d) $(CH_3)_3N$
- 378. Aniline first reacts with acetyl chloride producing compound 'A'. 'A' reacts with nitric acid/sulphuric acid mixture and produces compound 'B', which hydrolyses to compound 'C'. What is the identify of 'C'?
 - a) Acetanilide
- b) *p*-nitroacetanilide
- c) p-nitroaniline
- d) Aniline

AMINES

CHEMISTRY

: ANSWER KEY:															
1)	С	2)	С	3)	С	4)		177)	С		С	179)	С	180)	d
5)	d	6)	c	7)	a	8)	С	181)	a	400	a	183)	a	184)	d
9)	c	10)	c	11)	c	12)	d	185)	a	404	С	187)	a	188)	d
13)	b	14)	d	15)	b	16)	d	189)	a	400	a	191)	b	192)	b
17)	b	18)	b	19)	b	20)	b	193)	a		С	195)	a	196)	b
21)	b	22)	b	23)	d	24)	a	197)	d	-	d	199)	b	200)	c
25)	c	26)	d	27)	b	28)	d	201)	b	202)	d	203)	a	204)	b
29)	d	30)	a	31)	c	32)	b	205)	b	206)	b	207)	c	208)	b
33)	b	34)	b	35)	c	36)	a	209)	b	210)	b	211)	c	212)	b
37)	c	38)	d	39)	d	40)	b	213)	a	214)	С	215)	a	216)	c
41)	a	42)	b	43)	a	44)	a	217)	a	218)	a 🗸	219)	a	220)	d
45)	b	46)	d	47)	b	48)	c	221)	b	222)	a	223)	d	224)	b
49)	a	50)	d	51)	a	52)	b	225)	a	226)	b	227)	b	228)	a
53)	b	54)	b	55)	d	56)	b	229)	C,	230)	d	231)	c	232)	d
57)	b	58)	d	59)	a	60)	b	233)	d	234)	С	235)	c	236)	a
61)	C	62)	C	63)	c	64)	c	237)	b	238)	b	239)	a	240)	d
65)	d	66)	C	67)	a	68)	c	241)	c	242)	b	243)	b	244)	C
69)	c	70)	a	71)	C	72)	C	245)	b	246)	b	247)	b	248)	a
73)	c	74)	b	75)	a	76)	d	249)	d	250)	d	251)	b	252)	a
77)	a	78)	d	79)	d	80)	c	253)	b	,	d	255)	a	256)	d
81)	a	82)	a	83)	d	84)	b	257)	b	,	С	259)	b	260)	d
85)	d	86)	c	87)	C	88)	a	,	b	•	d	263)	b	264)	C
89)	a	90)	b	91)	C	92)	b	265)	C	,	a	267)	b	268)	C
93)	b	94)	a	95)	d	96)	b	269)	b	,	С	271)	b	272)	a
97)	C	98)	d		d	100)	C	273)	d	,	a	275)	b	276)	a
101)	d	102)	b	103)	c	104)	b	,	C	,	С	279)	C	280)	a
105)	b	106)	d	107)	b	108)	d	281)	a	,	a	283)	b	284)	b
109)	C	110)	d	111)	a	112)	a	,	a	,	a	287)	a	288)	С
113)	b	114)	a	115)	d	116)		289)	d	-	b	291)	b	292)	С
117)	b	118)	d	119)	a	120)		293)	c	•	b L	295)	a	296)	a
121)		122)	ď	123)	d h	124)		297)	a	•	d L	299)	b b	300)	c
125) 129)	c d	126) 130)	b	127) 131)	b d	128) 132)		301) 305)	c	•	b b	303) 307)	b b	304) 308)	C
133)		134)	c	131) 135)		136)		309)	c b			311)	b	312)	d d
137)	a a	134)	c c	139)	a a	140)		313)	d	-	c b	315)	c a	316)	a
141)		142)	a	143)	d	144)		317)	b		a	319)	b	320)	b
145)	a	146)	a	147)	C	148)		321)	d		c C	323)	d	324)	a
149)	d	150)	d	151)	b	152)		325)	a		a	327)	d	328)	c
153)	b	154)	a	155)	a	156)		329)	a		a	331)	c	332)	a
157)	c	158)	d	159)	С	160)		333)	b		c	335)	c	336)	a
161)	b	162)	c	163)	a	164)		337)	c	-	d	339)	c	340)	b
165)	d	166)	d	167)	С	168)		341)	b		a	343)	b	344)	d
169)	b	170)	a	171)	b	172)		345)	a	-	d	347)	d	348)	b
173)	c	174)	c	175)	b	176)		349)	d		a	351)	c	352)	c
-,	-	- 7	-	-,		-,	-	1 7		- ,		-,	-	,	-

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b 369)
353)
            354)
                  b
                        355)
                             d
                                    356)
                                                 a
                                                         370)
                                                             C
                                                                     371)
                                                                                 372) c
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                                                                          С
357)
            358)
                        359)
                              d
                                    360)
                                          c 373) c
                                                        374) d
                                                                     375) a
                                                                                 376) b
     C
                  b
                        363)
                                    364)
                                          c 377) c
                                                         378) c
361)
            362)
                  d
                              b
                        367)
365)
            366)
                                    368)
                                          b
```

SMART ACHIEVERS LEARNING BUT. LITT

AMINES

CHEMISTRY

: HINTS AND SOLUTIONS :

2 **(c)**

Hofmann's bromamide reaction is used to convert amide to amine.

 $RCONH_2 + Br + 4KOH$

$$\rightarrow RNH_2 + K_2CO_3 + 2KBr + 2H_2O$$
amine

amide

4 **(d)**

Secondary amides such as N-methylethanamide on reduction with ${\rm LiAIH_4}$ give secondary amines.

 $CH_3CONHCH_3 + [H] \xrightarrow{LiAIH_4} CH_3CH_2NHCH_3 + H_2O$ N-methylethanamide 2°amine

5 **(d)**

$$CH_3CN \xrightarrow{HOH} CH_3COOH$$

6 **(c)**

$$H$$

$$|$$

$$C_2H_5NH_2 + O = C - C_6H_5 \rightarrow C_2H_5N$$

$$= HCC_6H_5 + H_2O$$

$$Benzaldehyde$$

(A)

 $NH_2CO(NH_2 + H)NH\cdot NH_2 \longrightarrow NH_2CONHNH_2 + NH_3$ urea hydrazine

$$C_2H_5NH_2 + NOCl \rightarrow C_2H_5Cl + H_2O + N_2$$
(C)

11 (c)

The reaction is believed to follow the mechanism.

$$R \longrightarrow CONH_2 + OBr^- \longrightarrow RCONHBr + OH^-$$

$$RCONHBr + OH^- \longrightarrow RCONBr + H_2O$$

$$RCONBr \longrightarrow R - C = O + Br^- R - C = O \longrightarrow R - \ddot{N} = C = O$$

$$\vdots$$

$$\vdots$$

$$N$$

$$\vdots$$

$R-N=C=O+2OH^- \longrightarrow RNH_2 + CO_3^{2-}$

12 (d)

 HNO_2 reacts to give an alcohol means the compound is primary amine.

C₅H₁₃N means C₅H₁₁NH₂(primary amine)

Optically active alcohol means C_5H_{11} segment contain a chiral carbon.

$$\begin{array}{cccc} & & & H \\ 5 & 4 & 3 & 2 \mid \\ \text{CH}_3 - \text{CH}_2 - \text{CH}_2 - \text{C} - \text{NH}_2 \\ & & \mid \\ & & 1 \text{ CH}_3 \end{array}$$

Pentan-2-amine

13 **(b)**

$$R - CN + 4[H] \xrightarrow{\text{LiAIH}_4} R - CH_2NH_2$$

Alkyl nitriles primary amine

14 (d)

Electron donors are bases. In the given choices structure which does not involve resonance will have electron easily available for donation, hence most basic.

- ∴ Only in choice (b) electrons are not in conjugation with double bond of adjacent atom.
- ∴ Compound in choice (b) is most basic.
- 15 **(b**)

$$RBr + KCN \rightarrow RCN \xrightarrow{Reduction} RCH_2NH_2$$

16 **(d)**

It is methyl amine which, being basic dissolves in dilute HCI. It with ${\rm NaNO_2}$ evolves nitrogen gas leaving behind methyl alcohol which has smell of wood-spirit.

$$\begin{aligned} \mathsf{CH_3NH_2} &\overset{\mathsf{HCI}}{\longrightarrow} \mathsf{CH_3NH_2}. \mathsf{HCI} \\ \mathsf{CH_3NH_2} &+ \mathsf{HNO_2} &\overset{\mathsf{NaNO_2/HCI}}{\longrightarrow} \mathsf{CH_3OH} + \mathsf{N_2} \uparrow + \mathsf{H_2O} \\ && \mathsf{methyl} \ \mathsf{alcohol} \end{aligned}$$

17 **(b)**

Benzaldehyde condenses with N, N-diamethyl aniline in presence of anhydrous ${\rm ZnCl_2}$ to give malachite green

$$\begin{array}{c} \text{H-} & \text{CH}_3 \\ \text{C}_{6}\text{H}_5\text{CH=O} & + \\ \text{H-} & \text{CH}_3 & \frac{\text{Anhy.}}{\text{ZnCl}_2} \\ \text{CH}_3 & -\text{H}_2\text{O} \end{array}$$

19 **(b)**

This is carbylamine reaction which is used to distinguish 1° amines from other amines. The reaction is given by 1° amines only.

$$C_2H_5NH_2 + CHCI_3 + 3KOH$$

 $\rightarrow C_2H_5N \xrightarrow{=} C + 3KCI + 3H_2O$

$$RNH_2 + CHCI_3 + 3KOH$$

 $\rightarrow R - N \xrightarrow{=} C + 3KCI + 3H_2O$
1°amine chloroform isocyanide (bad smelling)

20 **(b)** $NH_2CONH_2 + HNHCONH_2 \rightarrow NH_2CONHCONH_2 + NH_3$

There are four β - hydrogens, in this quaternary ammonium salt.

On heating quaternary ammonium salt gives Hofmann elimination (abstraction of most acidic hydrogen which is β^1).

Hence, major product is $CH_2 = CH_2$. (Least substituted alkene).

$$CH_3NH_2 + CHCI_3 + 3KOH$$

 $\rightarrow CH_3NC + 3KCI + 3H_2O$

 CH_3NC or $CH_3 - N^+ \equiv C^-$ methyl isocyanide or methyl carbylamine.

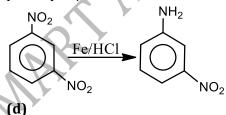
This reaction is an example of carbylamine reaction and it is used for the distinction of *p*-amines from *s*- and *t*-amines or identification of *p*-amino group.

25 **(c)**

Roulle first isolated urea (in 1773) from urine and named it as urea.

26 **(d)**

Reduction of NO₂ group to NH₂group is taking place by Fe/HCI.



$$\begin{array}{c|c} \mathsf{NH_2} & \mathsf{NH_2}.\mathsf{H_2SO_4} & \mathsf{NHSO_3H} \\ \hline \\ & & & \\ \hline \end{array}$$

Sulphanilic acid exists as a dipolar ion which has acidic and basic groups in the same molecule. Such ions are called Zwitter ions or inner salts

30 **(a**

For detection of secondary amines Liebermann's nitroso test is used.

31 (c) $C_2H_5NH_2 + Na \rightarrow C_2H_5NHNa + \frac{1}{2}H_2$

32 **(b**)

Only 1° aromatic amine (primary aromatic amine) from diazonium salts at low temperature (0° – 5°C). A reaction in which – NH₂ group is converted into diazo group ($-N^+ \equiv N$) is called diazotization. Diazotized salts are stable in cold aqueous solution.

$$C_6H_5NH_2+HC1 \xrightarrow{0^{\circ}-5^{\circ}C} C_6H_5NH2C1^{-}$$
 $C_6H_5NH_2+HC1 \xrightarrow{0^{\circ}-5^{\circ}C} HNO_2 +NaC1$
 $C_6H_5NC1^{\circ}+HNO_2 \xrightarrow{0^{\circ}-5^{\circ}C} C_6H_5N_2C1^{\circ}+2H_2O$
 $C_6H_5NH_2$
 $C_6H_5NH_2$
 C_6H_4
 $C_6H_5NH_2$
 C_6H_4
 $C_6H_5NH_2$
 C_6H_4

Amines, so undergo diazotization but $C_6H_5CH_2NH_2$ (aliphatic amine) will not undergo diazotisation.

33 **(b)**

Aniline is prepared by the reduction of nitrobenzene in acidic medium.

$$+6H$$
 Fe/HCl $+2H_2O$ aniline

34 **(b)**

Amines possess fishy smell.

36 **(a)**

Electrons donors are bases. Greater the stabilisation of cation formed by loss of electron more will be basicity of amine.

2° amine is more basic than 3° amine because 2° amine is stabilized by hydrogen bonding with solvent molecule.

$$RCONH_2 + NaOBr$$

 $\rightarrow RNH_2 + NaBr + Na_2CO_3$
 $+ 2H_2O$

38 **(d)**

Benzyl amine $(C_6H_5CH_2NH_2)$ is more basic than aniline $(C_6H_5NH_2)$ because N-atom of aniline is delocalized over the benzene ring. However in benzyl amine the lone pair of electrons on the N-atom is not conjugated with the benzene ring and therefore it is not delocalized. Hence, the lone pair of electrons on the N-atom in benzyl amine is more readily available for protonation than that on the N-atom of aniline. Thus, the benzyl amine is a stronger base than aniline.

39 **(d)**

Tertiary amines react as, $(CH_3)_3N + HNO_2 \rightarrow (CH_3)_3NHNO_2$

41 **(a)** Follow text.

44 (a)

$$C_2H_5NC + H_2O \xrightarrow{H^+} HCOOH + C_2H_5NH_2$$
formic acid
 $C_2H_5NH_2 + H^+ \rightarrow C_2H_2NH_3^+$
salt

45 **(b)**

It gives diazonium salt.

$$NH_2$$
 $+ NaNO_2 + 2HCl$
 $+ 2H_2O + NaCl$
benzene diazonium
chloride

It is known as diazotization reaction.

46 (d)

Aniline undergoes diazotisation.

47 (b)

$$RCN \xrightarrow{\text{Hydrolysis}} RCOOH + \text{NH}_3$$

$$RCH = \text{NOH} \xrightarrow{\text{Reduction}} RCH_2\text{NH}_2 + \text{H}_2\text{O}$$

$$RCN + 2H_2O \xrightarrow{\Delta} R\text{NH}_2 + \text{HCOOH}$$

$$RCONH_2 \xrightarrow{\text{Hydrolysis}} RCOOH + \text{NH}_3$$

49 (a)

Trimethyl amine is a tertiary amine. It dissolve in

cold nitrous acid to form salts which decompose on warming to nitrosoamine and alcohol. It will not liberate nitrogen.

 $(CH_3)_3N + HNO_2 \rightarrow [(CH_3)_3NH]^+NO_2^$ trimethyl ammonium nitrite

51 **(a)**

Carbylamine reaction is given by aliphatic and aromatic primary amine hence, it can be used for the distinguish of primary amine with secondary and tertiary amine. In this reaction, a primary amine reacts with chloroform and alcoholic KOH to give poisonous substance isocyanide.

$$RNH_2 + CHCI_3 + 3KOH(alc.) \xrightarrow{\Delta} RNC + 3KCI + 3H_2O$$

Primary amine

alkyl isocyanide

52 **(b)**

Nitrobenzene is reduced by Zn and alcoholic KOH into hydrazobenzene.

$$2C_6H_5NO_2 + 10H \xrightarrow{Zn+alc.KOH} C_6H_5 - NH - NH - C_6H_5 + H_2O$$
hydrazobenzene

53 **(b)**

Electron withdrawing groups (*e.g.*, benzyl) because the basicity of amines. Electron donating groups (*e.g.*, alkyl) increase the acidity of amines. \therefore The correct order of basicity of amines is $C_2H_5NH_2 > CH_3NH_2 > NH_3 > C_6H_5NH_2$

54 **(b**

Aliphatic amines (in which amino group is attached with alkyl group) are more basic than aromatic amines (in which amino group is bonded directly with benzene nucleus). Hence, $C_6H_5CH_2NH_2 \ (benzyl \ amine), being \ an \ aliphatic \ amine, is the most basic among the given the compounds.$

55 **(d)**

(a)
$$C_6H_5CONH_2 \xrightarrow{\text{LiAlH}_4} C_6H_5CH_2NH_2$$

Benzylamine

(b) $C_6H_5CH_2CONH_2 \xrightarrow{\text{Br}_2/KOH} C_6H_5CH_2NH_2$

Benzylamine

(c) $C_6H_5CN \xrightarrow{\text{LiAlH}_4} C_6H_5CH_2NH_2$

Benzylamine

(d) $C_6H_5CH_2NC \xrightarrow{\text{LiAlH}_4} C_6H_5CH_2NHCH_3$

56 **(b)**

Biuret formed gives violet colour with $CuSO_4$ in alkaline medium.

57 **(b)**

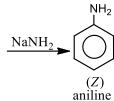
$$CH_3 - C \equiv N \xrightarrow{\text{Reduction}} CH_3CH_2NH_2$$

$$\xrightarrow{\text{HONO}} CH_3CH_2OH$$
methyl cyanide ethanamine ethanol
1. (B)

$$CH_3CH_2NH_2 \xrightarrow{CHCI_3,KOH} CH_3CH_2N \cong C$$
2. Ethyl isocyanide

(C)

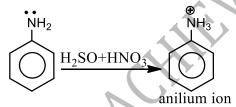
58 **(d)** $\begin{array}{c}
NO_2 \\
N_2CI \\
N_3CI \\
N_4CI \\
N_2CI



 $\therefore Z$ is aniline

59 **(a)**

On direct nitration of aniline, lone pair of electrons present at nitrogen atom will accept proton from the nitrating mixture to give anilium ion which is *meta* directing.



61 **(c)** R—NH₂ + HNO₂ \rightarrow ROH + N₂ + H₂O; But note that CH₃NH₂ gives CH₃ONO or CH₃OCH₃ on treating with HNO₂.

62 **(c)**The conversion of —CN to —CH₂NH₂ by catalytic reduction is called Mendius reaction.

64 (c) Schmidt reaction

0 ||
$$R - C - OH \xrightarrow{\text{NaN}_3}_{\text{H}_2\text{SO}_4} R - \text{NH}_2 + \text{N}_2 + \text{CO}_2$$

66 **(c)**

Methylaminomethane is trivial name of N-methyl methanamine (CH₃)₂NH.

67 **(a**

$$C_3H_9N(A) \xrightarrow{HNO_2} Alcohol + N_2$$

 $\Rightarrow A \text{ is } 1^\circ \text{ amine, } i.e.,$

 C_3H_9N is $C_3H_7NH_2$

$$C_3H_9N \xrightarrow{KOH+CHCl_3} Carbylamine \xrightarrow{Reduction}$$

$$CH_3$$
 $CH-NH-CH_3 \Rightarrow Alkyl \text{ part is } CH_3$ $CH-Hence,$ CH_3 $CH-$

$$C_3H_9N$$
 is CH_3 $CH-NH_2$

68 **(c)**

$$NH_2CONH_2 + HNHCONH_2 \xrightarrow{\Delta} NH_2CONHCONH_2$$
Biuret

69 **(c**)

In tertiary amines, no H-atom is attached directly to the more electronegative N-atom. Hence, it has no tendency to form H-bond

70 **(a)**

The order of boiling points of the isomeric amines is as fallows:

Primary amines > secondary amines > tertiary amines

$$(1^{\circ} > 2^{\circ} > 3^{\circ})$$

71 **(c)**

$$R$$
— $CONH_2 + Br_2 + KOH \rightarrow R$ — NH_2

72 **(c**

$$C_2H_5NH_2 \xrightarrow{HNO_2} C_2H_5OH + N_2 \uparrow + H_2O$$

76 **(d)**

$$NH_2$$
 +CHCl₃+ alc.KOH \triangle CH₃

p-toluidine (a carbylamine reaction)

77 (a)

Ethyl isocyanide on hydrolysis in acidic medium gives methanoic acid and ethyl amline salt

$$C_2H_5NC + H_2O \xrightarrow{H+} HCOOH + C_2H_5NH_2$$

methanoic acid

$$C_2H_5NH_2 + H^+ \rightarrow C_2H_5NH_3^+$$

Ethylamine salt

78 **(d)**

$$\begin{array}{c} CH_3I + NH_3 \longrightarrow CH_3NH_2 \stackrel{CH_3I}{\longrightarrow} (CH_3)_2NH \\ \stackrel{CH_3I}{\longrightarrow} (CH_3)_3N \stackrel{CH_3I}{\longrightarrow} (CH_3)_4N^+I^- \end{array}$$

80 **(c**)

Secondary amine on reaction with aq. $\rm HNO_2$ at low temperature produces yellow oily nitrosoamines. $\rm CH_3CH_2NHCH_2CH_3$ is secondary amine.

82 (a)

$$R \longrightarrow CN \xrightarrow{\text{Reduction}} RCH_2CH_2$$

83 (d)

 C_3H_9N may have the structures as: $(CH_3)_3N$, $CH_3CH_2CH_2NH_2$, $(CH_3)_2CHNH_2$

84 **(b)**

Tertiary amines, due to lack of H-atom, attached directly with N, does not react with benzene sulphonyl chloride ($C_6H_5SO_2CI$), i.e., Hinsberg's reagent. (C_2H_5)₃N is a tertiary amine, so does not react with $C_6H_5SO_2CI$.

85 (d)

$$\begin{array}{c} \text{CH}_3\text{CN} \xrightarrow[A=2]{\text{H.OH}} \text{CH}_3\text{COOH} \xrightarrow[A=2]{\text{Decarboxylation}} \text{CH}_4 \\ \xrightarrow[A=2]{\text{CH}_2\text{O}} \text{-NH}_3 \\ \end{array}$$

Ethane ethanoic acid

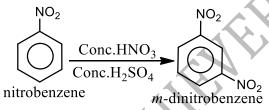
methane

Nitrile

88 (a)

Nitrobenzene on nitration gives *m*-dinitro benzene as – NO₂ group is meta-directing.

(A)



In this reaction the attacking reagent is NO_2^+ .

89 (a

Presence of—OCH₃ gp. on aniline makes it more basic than the presence of — NO₂, —Cl or — CH₃ gp.

90 **(b)**

Benzene diazonium chloride reacts with phenol in weakly basic medium gives *p*-hydroxy azobenzene.

chloride

p-hydroxy azobenzene

91 **(c)**

R—NC+2HgO $\rightarrow R$ NCO+Hg₂O

99 (d)

The abnormal trend of 3° amines is explained in terms of steric effect. Note basic order of amines on the basis of pK_b reported in Finar

$$(CH_3)_2NH > CH_3NH_2 > (CH_3)_3N > NH_3$$

3.23 3.32 4.2

4.73

 pK_b

$$(C_2H_5)_2NH > (C_2H_5)_3N > C_2H_5NH_2 > NH_3$$

3.07 3.13 3.37

4.73

 pK_b

100 (c)

SnCI₂ + 2HCI(g)
$$\rightarrow$$
 SnCI₄ + 2[H]
CH₃CN + 2[H] + HCI \rightarrow CH₃CH
= NH. HCI $\stackrel{\text{H}_2\text{O}}{\text{boil}}$ CH₃CHO + NH₄CI

 $(X) \qquad \qquad (Y)$

(X) is acetaldimine hydrochloride and (Y) is acetaldehyde.

101 **(d)**

Carbylamine reaction is given by aliphatic and aromatic primary amine.

CH₃CN does not give carbylamine reaction with chloroform because it is not an amine.

CH₃CN undergoes acidic hydrolysis to give carboxylic acid.

$$\begin{array}{c} \text{CH}_3\text{CN} \xrightarrow{\text{H}_2\text{O}/\text{H}^+} \text{CH}_3\text{CONH}_2 \xrightarrow{\text{H}_2\text{O}/\text{H}^+} \text{CH}_3\text{COOH} \\ + \text{NH}_3 \end{array}$$

acetic acid

 ${
m CH_3CN}$ undergoes alkaline hydrolysis to give salt of carboxylic acis.

$$\mathsf{CH_3CN} \xrightarrow[\mathsf{H_2O}]{\mathsf{NaOH}} \mathsf{CH_3CONH_2} \xrightarrow[\mathsf{H_2O}]{\mathsf{NaOH}} \mathsf{CH_3COONa} + \mathsf{NH_3}$$

Sodium acetate

CH₃CN tautomerises to give methyl isocyanide.

$$CH_3$$
— $C \equiv N \longrightarrow C \equiv N \longrightarrow CH_3$

105 **(b)**

Aniline reacts with Br₂to give 2, 4, 6-tribromoaniline not bromoaniline as

$$H_2$$
 $+ 3Br_2$
 Br
 $+ 3HBr$

2,4,6-tribromoaniline

106 (d)

 ${
m CH_3}$ — (an electron releasing (+I)group) increases electron density at N-atom hence, basic nature is increased.

$$I^{-}$$
 NH_{2}
 I^{-}
 NH
 CH_{3}
 I^{-}
 NH_{2}
 CH_{3}

 C_6H_5 decreases electron density at N-atom thus basic nature is decreased. (Lone-pair on N in aniline compounds is delocalised along with π -electrons in benzene).

Thus, (d) is the strongest base.

107 **(b)**

Urea gives biuret test. Biuret formed gives violet colour with CuSO₄ in alkaline medium.

110 (d)

or C₆H₅SO₂CI is called Hinsberg

reagent.

It is used for the distinction of 1°, 2°, 3° amine.

111 (a)

$$R - NH_2 + CHCl_3 + 3KOH \rightarrow 3KCl + 3H_2O$$

+ RNC
Offensive

112 (a)

Ethyl amine, on acetylation, gives N-ethyl acetamide.

$$\mathsf{C_2H_5NH_2} \xrightarrow{\mathsf{CH_3COCI}} \mathsf{C_2H_5NHCOCH_3}$$

N-ethyl acetamide

113 **(b)**

Aniline on oxidation with per acetic acid, CH₃CO₃H gives nitrobenzene

114 (a)

Aniline or any 1° amine reacts with aldehyde to form Schiff's base or anils.

$$NH_2$$

$$+ CH_3CHO \xrightarrow{-H_2O} N \equiv CHCH_3$$
aniline acetadehyde schiff's base or anil

115 (d)

Aniline reacts with bromine and give white ppt. of 2, 4, 6 tribromoaniline.

116 **(b)**

$$R - N$$
 is nitroalkane having $-N$ of f. gp.

R—0— N=0 is alkylnitrite having —0—N=0 f. gp. ; f. gps are different.

118 (d)

Isocyanide on hydrolysis forms primary amine not ammonia.

$$C_6H_5NC + H_2O \xrightarrow{H^+} C_6H_5NH_2 + HCOOH$$

119 (a)

Intermolecular H-bonding is more in primary amines than in secondary amines as there are two H-atom available for H-bonding. Tertiary amines do not have intermolecular H-bonding due to absence of H-atom available for H-bonding. Therefore, the order of boiling points of the given amines is as fallows

$$nC_4H_9NH_2 > (C_2H_5)_2NH > C_2H_5N(CH_3)_2$$

b.p. 350.8 K b.p. 329.3 K b.p. 310.5 K

121 **(b)**

$$R$$
— $NO_2 \xrightarrow{Sn/HCl} R$ — NH_2

123 (d)

It is carbylamine reaction,

benzyl amine
$$CH_2NH_2$$
 CH_2NH_2
 CH_2NC
 CH

$$C_6H_5NH_2 \xrightarrow{Diazotisation} C_6H_5N_2Cl + H_2O$$

127 **(b)**

In carbylamines reaction, when a primary amine reacts with chloroform in presence of alc. KOH, it gives iso-cyanide which has abonxious odour.

This reaction is given by primary amine

$$RNH_2 + CHCl_3 + 3KOH \xrightarrow{\Delta} RNC + 3KCl + 3H_2O$$

Primary (alc) alkyl
Amine *iso-cyanide*

128 **(c)**

K₂CO₃ is formed in Hofmann's degradation reactio

129 **(d)**

Electron withdrawing groups result in decreased basicity while electron releasing groups increases the basicity. Thus, the order of basic character is D > A > B > C

131 **(d)**

Isopropyl amine is a primary amine because one hydrogen atom of ammonia is replaced by isopropyl group.

$$\begin{array}{c} \rm NH_2 \\ | \\ \rm CH_3 - CH - CH_3 \\ 1^{\circ} \ amine \end{array}$$

132 (a)

$$CH_3CH_2NH_2 \xrightarrow{NaNO_2+HCl} CH_3 CH_2 OH + N_2 + H_2Cl$$

Ethylamine

$$NH_2CONH_2 \xrightarrow{NaNO_2+HCl} 2N_2 + H_2O + CO_2 + NaCl$$

Urea

$$CH_3CONH_2 \xrightarrow{NaNO_2 + HCl} CH_3COOH + N_2 + H_2O$$
+ NaCl

Acetamide

$$C_6H_5NH_2 \xrightarrow{NaNO_2+HCl} C_6H_5N_2^+Cl^-H_2O + NaCl$$
Aniline

133 (a)

$$NH_2CONH_2 + HOC_2H_5 \xrightarrow{\Delta} H_2NCOOC_2H_5 + NH_3$$
Urethane

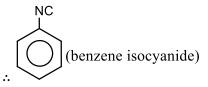
136 (b)

Carbylamine (or isocyanides) give secondary amine on reduction.

$$R - N \stackrel{\supseteq}{=} C \xrightarrow{\text{Ni/H}_2} R - \text{NH} - \text{CH}_3$$
 carbylamine secondary amine

137 (a)

Isocyanides (carbylamines) are foul odour compounds.



As foul odour

138 (c)

Reduction of nitrobenzene by Zn and NH₄CI gives N-phenyl hydroxylamine.

$$+4[H]$$
 $2n/NH_4Cl$ $+H_2O$

N-phenyl hydroxylamine

139 (a)

3. In Hofmann method, a mixture of primary, secondary and tertiary amines is treated with diethyloxalate, when primary amine forms solid oxamide, secondary amine forms a liquid oxamic ester whereas tertiary amine remains unaffected.

$$(COOC_2H_5)_2 + 2HNHR \rightarrow (CONHR)_2 + 2C_2H_5OH$$

Diethyl oxalate 1° amine solid

$$(COOC_2H_5)_2 + HNR_2 \rightarrow |$$

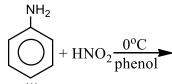
$$COOC_2H_5 + C_2H_5OH$$

2°amine liquid

$$(COOC_2H_5)_2 + NR_3 \rightarrow No reaction$$

141 **(b)**

 $C_2H_5NH_2 + HNO_2 \rightarrow C_2H_5OH + N_2 + H_2O$ ethyl amine ethyl alcohol (1°aliphatic amine)



aniline

1° aromatic amine

∴Dye test is used to distinguish between

1°aliphatic and

1° aromatic amine

142 (a)

Hofmann bromamide degradation takes place with complete retention of stereochemical configuration in the migrating alkyl group

143 (d)

$$C_6H_5SO_2Cl + RNH_2 \rightarrow RNHSO_2C_6H_5 \xrightarrow{KOH} R - NKSO_2C_6H_5$$

Benzene sulphyonyl N-alkyl benzene soluble in KOH

chloride sulphonamide

144 **(b)**

The conversion of —CN to — CH_2NH_2 by catalytic 162 (c) reduction is called Mendius reaction.

147 (c)

$$\mathsf{C_2H_5NH_2} \xrightarrow[-N_2]{\mathsf{HNO}_2} \mathsf{C_2H_5OH} \xrightarrow{\mathsf{PCI}_3} \mathsf{C_2H_5CI} \xrightarrow{\mathsf{NH}_3} \mathsf{C_2H_5NH_2}$$

ethyl amine ethanol ethyl chloride ethyl amine

4.

(B)

(C)

148 (a)

Azoxybenzene is the main product when reducing agent used is Na₃AsO₃/NaOH

149 (d)

C₆H₅COCl has no reaction with carboxylic acids.

150 (d)

CH₃NH₃Cl is acidic in nature.

153 **(b)**



Formation of *A* is by diazotization and formation of B from A is by S_N reaction.

154 (a)

$$RCONH_2 \xrightarrow{P_2O_5} RCN + H_2CONH_2 \xrightarrow{P_2O_5$$

155 (a)

$$\begin{array}{c} \text{CH}_3\text{CH}_2\text{I} \xrightarrow{\text{NaCN}} \text{CH}_3\text{CH}_2\text{CN} \xrightarrow{\text{OH}^-} \text{CH}_3\text{CH}_2\text{CONH}_2 \\ \xrightarrow{\text{Br}_2/\text{NaOH}} \text{CH}_3\text{CH}_2\text{NH}_2 \end{array}$$

Amines react with alkyl halide (excess) to give quaternary ammonium salt.

 $C_6H_5NH_2 + 3CH_3I \rightarrow C_6H_5N^+(CH_3)_3I^-$

159 (c)

Reaction of aniline with benzaldehyde is condensation reaction.

> Η I

$$\begin{aligned} \mathsf{C}_6\mathsf{H}_5 - \mathsf{C} &= \mathsf{O} + \mathsf{H}_2\mathsf{N}\mathsf{C}_6\mathsf{H}_5 \xrightarrow{\mathsf{Condensation}} \mathsf{C}_6\mathsf{H}_5\mathsf{CH} \\ &= \mathsf{N}\mathsf{C}_6\mathsf{H}_5 + \mathsf{H}_2\mathsf{O} \end{aligned}$$

Benzaldehyde aniline benzylidene aniline

161 **(b)**

O
$$\parallel$$
 $-\ddot{N}H$ and $-C-gp$. are o-and p-directing (activating) and m-directing gps.(deactivating)

respectively. In such case o-(minor) and p isomers (major) predominates.

C₂H₅NH₂ is stronger base than NH₃. The presence of alkyl group on N-atom intensifies -ve charge on N-atom and thus, electron pair is donated more readily.

$$R \longrightarrow NH_2$$
 $H-NH_2$ (+ve *IE* of alkyl gp.)

163 (a)

$$(CH_3)_4N^+I^- + NaOH \rightarrow (CH_3)_4N^+OH^- + Nal$$

 $(CH_3)_4N^+OH^- \stackrel{\Delta}{\rightarrow} (CH_3)_3N + CH_3OH$
methanol

164 (c)

Electron deficient group decreases the electron density of N-atom, thus, makes its lone pair less available for donation

165 (d)

$$\begin{array}{c} {\sf C_6H_5NH_2+CI-COCH_3} \stackrel{{\sf NaOH}}{\longrightarrow} {\sf C_6H_5NHCOCH_3} \\ & + {\sf HCI} \end{array}$$

aniline acetyl chloride acetanilide

166 **(d)**

C₃H₉N represent following structures CH₃CH₂CH₂NH₂

Propanamine

(1°amine)

$$\mathrm{CH_3CH_2} - \mathrm{N} - \mathrm{CH_3}$$

N-methyl ethanamine

(2°amine)

$$CH_3$$

 $|$
 $CH_3 - N - CH_3$
 N,N -dimethyl methanamine
 $(3^\circ amine)$

167 (c)

Presence of α -H atom is the main condition for exhibiting tautomerism.

The reactant taken in reaction (C) does not

contain any α -H atom, thus the product (Y)will also show the absence of α -H atom, Hence, Y will show tautomerism

168 (a)

 R_2 NH + HNO₂ \rightarrow R_2 N—N=0 + H₂O Nitrosoamines are carcinogens.

169 **(b)**

Acetonitriles on hydrolysis produce carboxylic acids with the evolution of ammonia.

$$\begin{array}{ccc} & & & & & & & & & & \\ & & & & & & & || & & & || \\ \text{CH}_3 - \text{C} & \equiv \text{N} & \xrightarrow{\text{H}_2\text{O}} \text{CH}_3 - \text{C} - \text{NH}_2 & \xrightarrow{\text{H}_2\text{O}} \text{CH}_3 - \text{C} \\ & & & & - \text{OH} + \text{NH}_3 \end{array}$$

acetamide

171 **(b)**

Acetonitrile

Methyl cyanide gives acetic acid on hydrolysis.

$$CH_3CN \xrightarrow{H_2O/H^+} CH_3 - C - NH_2 \xrightarrow{H_2O/H^+} CH_3COOH + NH_3$$

172 (c)

 $2RNH_2 + 2HCl + PtCl_4 \rightarrow (RNH_3)_2PtCl_6; RNH_3AuCl_4$

173 (c)

General formula for any amine is $C_nH_{2n+3}N$; also note that for primary amine, it is C_nH_{2n+1} NH_2 ; for secondary amine, it is $C_nH_{2n+2}NH$ and for tertiary amine, it is $C_nH_{2n+3}N$.

174 (c)

Aniline on diazotization in cold (at 0° to 5°C) gives benzene diazonium chloride.

$$NH_2 + NaNH_2 + 2 HCI \frac{0-5^{\circ}C}{Diazotization}$$
 $N=N-CI + NaCI + 2H_2O$

benzene diazonium chloride

This benzene diazonium chloride on coupling reaction with dimethyl aniline gives a coloured product *i.e.*, p(N, N dimethyl) amino azobenzene (azodye)

$$N=N.Cl+H \longrightarrow N(CH_3)_2 \xrightarrow{NaOH} N=N-N-N(CH_3)_2 \longrightarrow N(CH_3)_2 \longrightarrow N(CH_$$

176 (c)

Nitrobenzene on reduction with lithium aluminium hydride (LiAIH₄) gives azobenzene.

$$2 \underbrace{ \frac{\text{LiAlH}_4}{\text{Direction}}}_{\text{Nitrobenzene}} \underbrace{ \frac{\text{LiAlH}_4}{\text{azobenzene}}}_{\text{azobenzene}}$$

177 (c)

Both gives alkane (*RH*) with Grignard reagents *RMgX* due to the presence of acidic hydrogen (N—H)

179 (c)

Presence of methyl group increase the electron density on nitrogen. So, increases the basicity. Aniline is weaker base than the primary aliphatic amines and this may be explained by resonance. The lone pair of N is involved in resonance, thus not available for donation. That's why basic strength of aryl amines (aniline) is lowest.

180 (d)

acetic acid

All primary (aliphatic) amines give alcohol with H which forms

$$CH_3$$
— 0 — N = 0 and CH_3 0 CH_3 .

181 **(a)**

Hofmann bromamide reaction is used to prepare 1° amine form primary amides. In this method, amides are treated with bromine in presence of KOH

$$R - CONH_2 + 4KOH + Br_2 \xrightarrow{\Delta} RNH_2 + K_2CO_3 + 2KBr + 2H_2O$$

184 **(d)**

The amines are basic in nature due to presence of lone pair of electron on nitrogen. The 2°amines are basic among 1°, 2° and 3°amines because of steric effect and hydration effect

185 (a)

This is carbylamine reaction.

186 **(c)**

p-amine forms alcohol; *s*-amine forms only nitrosoamine.

187 (a)

$$\begin{aligned} \mathsf{CH_3NH_2} + \mathsf{C_6H_5SO_2CI} \\ &\to \mathsf{C_6H_5SO_2NHCH_3} \\ &\xrightarrow[(\mathsf{alkali})]{\mathsf{NaOH}} \mathsf{C_6H_5SO_2N(Na)CH_3} \end{aligned}$$

1° amine hinsberg's N-methyl benzene soluble sodium salt

Reagent sulphonamide

188 (d)

$$NH_2$$
 $+ CHCl_3 + alc 3KOH$
 CH_3
 $- CH_3$
 $-$

it is a example of carbylamines reaction

189 (a)

Cyanides are hydrolysed either by alkali or acid to give carboxylic acid.

$$R - CN + 2H_2O \xrightarrow{\text{NaOH}} R - COOH + NH_3$$

190 (a)

$$C_2H_5NH_2 + NOCl \rightarrow C_2H_5Cl + N_2 + H_2O$$

191 **(b)**

By using H_2S , NH_3 as reagent, selective reduction takes place

$$NO_2$$
 NO_2
 NO_2
 NO_2
 NO_2
 NO_2

192 **(b)**

 2° amines are more basic than 1° amines due to +ve IE of — CH_3 gp. In amide the resonance give rise to less availability to electron pair for coordination and thus less basic. The negative

charge developed on N in CH_3-C NH

NH

NH2

due to resonance makes it more basic.

Nitration of aniline also gives m-nitro aniline in strong acidic medium because in strong acidic condition protonation of – NH_2 group gives anilinum ion (+ NH_3) which is deactivating in nature and of m-directive nature

195 (a)

Grabriel's synthesis: Phthalimide is reacted with KOH to form potassium phthalimide. The potassium salt is treated with an alkyl halide. The product N-alkyl phthalimide is put to hydrolyse with hydrochloric acid, then primary amine is formed.

196 **(b)**

$$C_6H_5NO_2 \xrightarrow{Sn/HCl} C_6H_5NH_2$$

Nitrobenzene Aniline Nitrobenzene in reduction with Sn and HCI produce aniline. Hence, 'X' is identified as – NH₂ group.

197 (d)

 $\begin{array}{c} \text{CH}_3\text{CHO} \\ \xrightarrow{\text{H}_2\text{NOH}} \text{CH}_3\text{CH} = \text{NOH} \xrightarrow{\text{Reduction}} \text{CH}_3\text{CH}_2\text{NH}_2 \xrightarrow{\text{NOCI}} \text{CH}_2\text{NH}_2 \xrightarrow{\text{H}_2\text{O}} \text{CH}_2\text{NH}_2 \\ & \text{Acetaldehyde} \end{array}$

chloride

(A) (B)

199 **(b)**

 $CH_3CH_2 - 0 - N = 0$ is a nitrite derivative, hence it is not a nitro derivative.

200 **(c)**

Basic nature of an amine depends upon availability of lone pair on nitrogen atom. If lone pair is easily available the compound would be more basic.

Dut to +I effect of methyl group, methyl amine is more basic than ammonia and dimethyl amine is more basic than methyl amine. While aniline is a weaker base than ammonia due to delocalization of lone pair of nitrogen atom at different position.

202 **(d)**

When aqueous solution of benzene diazonium chloride boiled, it gives phenol.

$$N_2$$
⁺Cl⁻ OH
benzene
diazonium
chloride

206 **(b)**

$$C_2H_5NH_2 + CHCI_3 + 3KOH$$

 \rightarrow C₆H₅NC + 3KCI + 3H₂O

Aniline chloroform phenyl isocyanide Thus in this reaction phenyl isocyanide is produced. This is called carbylamine reaction.

208 **(b)**

$$RNH_2 \xrightarrow{HNO_2} ROH + H_2O + N_2 \uparrow$$

5. (B) (C)

209 (h)

$$(A)RCN \xrightarrow{\text{Reduction}} RCH_2NH_2$$

$$1^{\circ} \text{amine}$$

$$RCN + CH_{3}MgBr \longrightarrow R - C = N - MgBr \xrightarrow{H. OH} Br$$

$$CH_{3} \longrightarrow C = O + NH_{3}$$

$$CH_{3} \longrightarrow C \longrightarrow C$$

$$CH_{3} \longrightarrow C \longrightarrow C$$

$$CH_{4} \longrightarrow C$$

(C)
$$RNC \xrightarrow{Hydrolysis} H - COOH + RNH_2$$
1°amine

(D)
$$RNH_2 \xrightarrow{HNO_2} ROH + N_2 + H_2O$$

211 (c)

$$\begin{array}{c} \operatorname{CH_3CH_2Br} \xrightarrow{\operatorname{aq.KOH}} \operatorname{CH_3CH_2OH} \xrightarrow{\operatorname{KMnO_4/H}^+} \operatorname{CH_3COOH} \\ \xrightarrow{\operatorname{NH_3}} \operatorname{CH_3COONH_4} \xrightarrow{\Delta} \operatorname{CH_3CONH_2} \end{array}$$

6. Acetic acid acetamide

(B)

(C)

 $\xrightarrow{\text{Br}_2/\text{alkali}} \text{CH}_3\text{NH}_2$

Methyl amine (D)

212 **(b)**

$$C \equiv N \xrightarrow{\text{Na}} C$$

$$\text{Ether}$$

$$NH$$

$$CN$$

This is thorpe nitrile condensation involving only α-H-atom of nitrile due to strong attraction nature of CN gp.

213 (a)

 $-NO_2$ is a *meta* directing group. As it is also a deactivating group, so no chance of introduction of second - Br atom.

216 **(c)**

$$(CH_3)_2CH.NH_2 + 2CH_3COCI \xrightarrow{-HCI} (CH_3)_2CH$$

- $N(COCH_3)_2$

iso-propyl amine

(ter-amine)

(pri amine)

218 (a)

$$RCH_2NO_2 \xrightarrow{HOH} RCOOH + NH_2OH$$

219 (a)

 $R - X + \text{NaCN} \rightarrow R - \text{CN} + \text{Na} X$ Alkyl halide aliphatic cyanide

220 **(d)**

It is not used in purification of water.

223 **(d)**

The mixture of SnCl₂ + HCl, reduces, -CN group partially

224 **(b)**

Action of alcoholic caustic potash on chloroform and aniline forms a bad smelling compound phenyl isocyanide.

 $C_6H_5NH_2 + CHCI_3 + 3KOH(alc.)$

$$\rightarrow$$
 C₆H₅NC + 3KCI + 3H₂O

chloroform aniline

isocyanide

This reaction is called carbylamine reaction and it is actually the test of primary amines.

225 (a)

$$RNH_2 + C_6H_5SO_2Cl \rightarrow C_6H_5SO_2NHR$$
(Sulphonamide soluble in alkali)

226 (b)

$$\phi N_2^+ + \phi OH \rightarrow \phi N_2 \phi OH$$
or $\langle OH \rangle$
coupling product

227 **(b)**

$$\begin{array}{c} C_2H_5NH_2 \\ \xrightarrow{HNO_2} C_2H_5OH \xrightarrow{PCl_5} C_2H_5Cl \xrightarrow{NH_3} C_2H_5NH_2 \\ \text{Ethyl amine} \qquad \text{ethyl alcohol} \qquad \text{ethyl chloride} \\ \text{ethyl amine} \\ \end{array}$$

(*A*) (B)

228 (a)

The order of basicity among the following amines

$$(CH_3)_2NH > (CH_3)_3N > CH_3NH_2 > C_6H_5NH_2$$

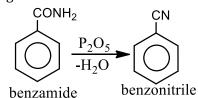
230 (d)

$$C_6H_5CH_2NH_2 + CHCI_3 + 3KOH$$

 $\rightarrow C_6H_5CH_2NC + 3KCI + 2H_2O$

232 (d)

With P₂O₅benzamide loses a water molecule and gives benzonitrile.



Here, P₂O₅ is a dehydrating agent.

233 (d)

Concentration does not affect the basis strength of amines

234 **(c)**

Nitrosoamine on heating with $conc.H_2SO_4$ gives secondary amine. This reaction is called Liebermann nitroso reaction.

$$R_2 N - N = 0 + H_2 O \xrightarrow{Conc.H_2SO_4} R_2 NH + HNO_2$$

nitrosoamine secondary amine

237 **(b)**

Negative charge developed on N in the resonance hybrid makes it more basic as it loses electron pair readily. Also 2° amine is more basic than 1° amine due to +ve IE of alkyl group. In amide the lone pair remains less available due to delocalisation in resonance.

239 (a)

$$R - C \equiv N + 4[H] \xrightarrow{\text{LiAIH}_4} RCH_2 NH_2$$

240 **(d)**

Anilinium hydrogen chloride ($C_6H_5NH_2$. HCI) gives white precipitate. With AgNO $_3$ solution as it gives CI $^-$ ion, p-chloro aniline has CI atom attached directly to the nucleus, hence cannot be ionised.

241 **(c)**

$$CCH3CH2NH2 + CHCI3 + 3KOH$$

$$\rightarrow CH3CH2NC + 3KCI + 3H2O$$
7. (B)

This reaction is known as carbylamine reaction

242 **(b)**

Nitromethane forms methyl hydroxylamine on reduction in netural medium with Zn/NH $_4$ CI. $\text{CH}_3 \text{NO}_2 + 4 \text{[H]} \xrightarrow{\text{Zn/NH}_4 \text{CI}} \text{CH}_3 \text{NHOH} + \text{H}_2 \text{O}$

Nitromethane

N-methyl hydroxyl

amine

243 **(b)**

A characteristic reaction of secondary amines. $(CH_3)_2NH \xrightarrow{HNO_2} (CH_3)_2 N-N=0$ (nitrosoamine).

244 (c)

As we know, benzenediazonium salt forms brilliant coloured dye with β -naphthol, the compound under consideration must be p-toludine (c) as it is a primary aromatic amine.

Primary aromatic amine, on treatment with ${\rm NaNO_2}$ in dil. HCI forms the corresponding diazonium chloride salt.

$$NH_2$$
 N_2
 ## 245 **(b)**

Hofmann's method is used to separate primary, secondary and tertiary amines. The compound used is diethyl oxalate for this purpose.

246 **(b)**

$$C_6H_5NO_2 \xrightarrow{Sn/HCI} C_6H_5NH_2$$

$$\xrightarrow{C_6H_5COCI} C_6H_5CONHC_6H_5$$

Nitrobenzene aniline benzanilide

247 **(b)**

Nitro group goes always to meta position, in aromatic compounds, irrespective to the substituents.

248 (a)

It is the first step of Gabriel's phthalimide synthesis. The hydrogen bonded to nitrogen is sufficiently acidic due to two α -carbonyls.

$$N \longrightarrow H + OH \longrightarrow H_2O + \bigcirc$$

$$N \longrightarrow N$$

The conjugate base forms above act as nucleophile in the subsequent step of reaction. As shown above, the nucleophile exist in three resonating form, one may think of oxygen being the donor atom in the nucleophilic attack. However, nitrogen act as donor as it is better donor than oxygen.

$$\begin{array}{c|c}
\hline
S_{N^2} \\
\hline
\end{array}$$
N- CH₂
Br

Bromine is not substituted in the above reaction as it is in resonance with benzene ring giving partial double bo0nd character to C – Br bond, hence difficult to break.

$$H_2CCI$$
 H_2CCI H_2CCI

249 (d)

Nitrobenzene is insoluble in water but soluble in benzene alcohol etc.

251 **(b)**

CHCI₃ + C₆H₅NH₂ + 3NaOH

$$\rightarrow$$
 C₆H₅NC + 3NaCI + 3H₂O
phenylisocyanide 267 **(b)**

252 (a)

An alkyl cyanide is dissolved in ether or better in ethyl formate or ethyl acetate, and reduced with SnCI₂ and HCI and then steam distilled. The whole process is called Stephen reaction. In this process alkyl cyanide is reduced to aldehyde.

$$CH_3 - C \equiv NH = 2[H] + HCI \xrightarrow{SnCI_2 + HCI}$$

There is no analogous method for the preparation 273 (d) of ketones.

253 (b)

 $CH_3CH_2 - O - N = O$ is a nitrite derivative, hence, it is not a nitro derivative

255 (a)

$$R - C \equiv N$$

: Alkyl group is attached to carbon.

Only primary amines give positive carbylamine test

258 (c)

Tertiary amine can be directly oxidised to corresponding nitro compound by KMnO₄.

259 **(b)**

Due ΙE to +ve in alkylamines and resonance in C₆H₅NH₂.

$$\begin{array}{c|c}
 & \text{NHCOCH}_3 \\
\hline
 & \text{HNO}_3 \\
\hline
 & \text{H}_2\text{SO}_4
\end{array}$$

$$\begin{array}{c|c}
 & \text{NHCOCH}_3 \\
\hline
 & \text{NO}_2
\end{array}$$

$$\begin{array}{c}
 & \text{NHCOCH}_3 \\
\hline
 & \text{NO}_2
\end{array}$$

$$\begin{array}{c}
 \text{NH}_2 \\
 \text{NO}_2 \\
 \text{NO}_2
\end{array}$$

 $(-NHCOCH_3 \text{ is } o\text{- and } p\text{-directing})$

263 **(b)**

$$C_2H_5NO_2 \xrightarrow{\text{NaOH}} CH_3COONa + NH_2OH$$

264 (c)

$$NH_2CONH_2 + NH_2 \cdot NH_2 \rightarrow NH_2CONHNH_2 + NH_3$$

$$C_6H_5NH_2 + CH_3COCl \rightarrow C_6H_5NHCOCH_3$$

268 (c)

The aqueous solution of urea is neutral in nature but urea behaves as a weak monoacid-base and forms salts with strong acids, e.g.,

$$\mathrm{NH_2CONH_2} + \mathrm{HNO_3} \longrightarrow \mathrm{NH_2CONH_2} \cdot \mathrm{HNO_3}$$
 Urea nitrate

270 (c)

Quaternary ammonium compounds, e.g., $[R_4N]^+X^-$, tetra alkyl amm. halide or $[R_4N]^+OH^-$, tetra alkyl amm. Hydroxide.

271 **(b)**

$$R - NH_2 \xrightarrow{HNO_2} R - OH$$

$$R$$
—N=C=0 $\xrightarrow{\text{HOH}} R$ NH₂ + H₂CO₃

274 (a)

Aliphatic primary amines on treatment with NaNo₂/HCI gives alcohols with evolution of nitrogen gas.

$$(CH_3)_2CHNH_2 + HONO \xrightarrow{NaNO_2/HCI} (CH_3)_2CHOH + N_2 + H_2O$$

275 **(b)**

+ ve IE in alkyamines and resonance in $C_6H_5NH_2$.

278 (c)

 $RCONH_2 + NaOH \rightarrow RCOONa + NH_3$

279 (c)

$$CH_3CN \xrightarrow{\text{Reduction}} CH_3CH_2NH_2$$

acetonitrile ethanamine

280 (a)

$$C_2H_5OH + NH_3 \rightarrow C_2H_5NH_2 + H_2O$$

281 (a)

Molecular formula of benzonitrile is C₆H₅CN.



phenyl cyanide or benzonitrile

282 (a)

Primary amines have tendency of forming H-bonds

283 **(b)**

Due to +ve IE in alkylamines and resonance in $C_6H_5NH_2$.

284 **(b)**

Only these aromatic primary amines undergo diazotisation in which $-\mathrm{NH}_2$ is attached to nucleus.

285 (a)

Due to +ve IE in alkylamines and resonance in $C_6H_5NH_2$.

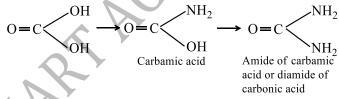
287 (a)

In vapour phase the basic character of methylamines is $3^{\circ} > 2^{\circ} > 1^{\circ} > NH_3$.

This is due to less acidic character in conjugate

291 **(b)**

Urea is monoamide of carbamic acid or diamide of carbonic acid,



293 (c)

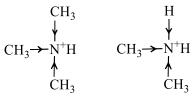
EDTA is ethylenediamine-tetra acetic acid.

295 (a)

Hinsberg reagent $C_6H_5CO_2Cl$ reacts with primary amines and gives alkali soluble benzene sulphonamide; with secondary amine it gives alkali insoluble benzene sulphonamide, with tertiary amines it does not react.

$$\begin{array}{ccc} {\rm C_6H_5SO_2Cl} + R{\rm NH_2} \longrightarrow & \\ R{\rm NHSO_2C_6H_5} & \stackrel{{\rm KOH}}{\longrightarrow} & R{\rm NKSO_2C_6H_5} \\ & & {\rm Soluble\ in\ KOH} \end{array}$$

acid of amines because of +ve $\it IE$ of methyl group which disperses +ve charge on N-atom more effectively in 3° ion.



+ve charge on N is more dispersed and thus, more stable.

In case of solution phase steric hindrance or solvation effect play role to give the basic order $2^{\circ} > 1^{\circ} > 3^{\circ} > NH_3$.

289 **(d)**

Compounds having active hydrogen such as, phenols, alcohols, primary or secondary amines and amides show Schotten-Baumann reaction. But tertiary amines do not have active hydrogen, hence, do not undergo Schotten-Baumann reaction.

$$R$$
—OH + NaOH $\frac{-H_2O}{-NaCl}$ OCOR

296 **(a)**

$$R-C \equiv N \xrightarrow{\begin{subarray}{c} \begin{subarray}{c} \begin{subarra$$

297 **(a)**

 $-0\mathrm{CH_3}$ is strongest electron releasing group (+M effect) which opposes most the dispersion of lone pair of electron of nitrogen into the ring. Thus, $-0\mathrm{CH_3}$ being at *para* position imparts hifhest basicity. $\mathrm{NO_2}$ being at *meta* position

stabilises the electron pair of nitrogen only by -I effect. While NO_2 being present at *para* position due to -M effect and -I effect stabilizes the lone pair of electron of nitrogen, most and impart least basicity.

$$O_2N$$
 NH_2
 NH_2
 NH_2
 NH_2

$$\begin{array}{c|c} & & & \\ &$$

298 (d)

Benzyl amine is the strongest base among the given compounds because lone of \ddot{N} are not taking part in conjugation whereas in other compounds lone pairs are taking part in conjugation

$$\bigcirc$$
 CH₂- $\stackrel{\bullet}{N}$ H₂

299 **(b)**

Basicity of amines increases with increasing +I effect of alkyl group.3°amine has greater +I effect than 2° and 1°amines but less basic than these, due to steric hindrance of bulky groups.

Moreover, benzyl amine is a weaker base than aliphatic amines. Hence, the following compounds has the order of basicity.

$$CH_3NH_2(1^\circ), (CH_3)_2NH(2^\circ), (CH_3)_3N(3^\circ), C_6H_5CH$$
 $II \qquad III \qquad IV$
 $II > I > III > V$

300 (c)

8. 0

$$|| CH3 - C2 - CH3 sp3 sp2 sp2$$

(b) 0

$$CH_3 - C_2 - OH$$

$$sp^3 sp^2$$

(c)

$$CH_3 - C - \equiv N$$

$$sp^3 \quad sp$$

(d) 0

$$\begin{array}{c} || \\ \mathrm{CH_3} - \mathrm{C_2} - \mathrm{NH_2} \\ sp^3 & sp^2 \end{array}$$

Acetonitrile does not contain sp^2 hybridised carbon.

301 (c)

Availability of lone pair on N-atom (Lewis concept),

 $RNH_2 + H_2O \rightarrow RNH_3^+ + OH^-$ (Bronsted concept proton acceptor).

302 **(b)**

The order for acidic nature depends upon the ease to lose H⁺ ion. —COOH is resonance stabilized and thus lose H⁺ at the earliest. Also NH₃⁺ near to —COOH releases H⁺ more easily due to electron withdrawing nature of —COOH than NH₃⁺ far away from —COOH.

303 **(b)**

Ethyl bromide reacts with KCN to give ethyl cyanide.

 $C_2H_5Br + KCN(alc.) \rightarrow C_2H_5CN + KBr$ Ethyl bromide ethyl cyanide

305 **(c)**

Carbylamine reaction is used to prepare *iso* cyanides.

306 **(b)**

Dimethyl oxalate is used for distinguishing primary, secondary and tertiary amines (Hofmann's method).

9. The primary (1°) amine forms the corresponding substituted oxamide which is usually a crystalline solid.

10. The secondary amine forms a diethyl oxamic ester which is generally a liquid.

$$COOC_2H_5$$
 $CONR_2$

$$| +NHR_2 \rightarrow | +C_2H_5OH$$

$$COOC_2H_5 \qquad COOC_2H_5$$

11. The tertiary amine under these conditions does not react at all since it does not

contain a replaceable hydrogen atom.

307 **(b)**

 C_3H_8O is alcohol and $C_3H_6O_2$ is acid. Thus, C_3H_9N i

308 (d)

$$CH_3CONH_2 + NaOH \rightarrow CH_3COONa + NH_3$$
(A)

$$(C_2H_5)_2CO_3 + 2NH_3 \rightarrow NH_2CONH_2 + 2C_2H_5OH$$

ethyl carbonate urea

$$NH_2COOC_2H_5 + NH_3 \rightarrow NH_2CONH_2 + 2C_2H_5OH$$

$$COCl_2 + 2NH_3 \rightarrow NH_2CONH_2 + 2HCl$$

phosgene

311 (c)

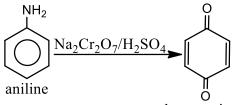
$$CH_3 - NH_2 + 3CH_3I \rightarrow (CH_3)_4N^+I^-$$

methyl amine quaternary

therefore, 3 moles of $\mathrm{CH_{3}I}$ (methyl iodide) are required for reaction with methyl amine.

312 (d)

Aniline on oxidation with Na₂Cr₂O₇and H₂SO₄ gives *p*-benzoquinone.



p-benzoquinone (quinone)

313 (d)

Presence of electron withdrawing group like -CN, -CHO, $-\text{NO}_2$ etc decreases the electron density over N- atom of amines, and thus decreases their basic character as lone pair is less available for donation

314 **(b)**

Secondary and tertiary amines fail to undergo the carbylamine test because they react with alcoholic KOH.

317 **(b)**

Due to +ve IE of alkyl gp., N-atom of amines acquires partial -ve charge and thus, electron pair is easily donated.

319 (b)

Aniline is not obtained as a major product by the reaction.

$$C_6H_5NO_2 + Zn powder \xrightarrow{\text{Alcoholic KOH}} H_1$$

Nitrobenezene

321 (d)

Bases react with acid to form salt.

- ∴ Amines are basic in nature.
- ∴It forms salt on reaction with H_2SO_4 $2RNH_2 + H_2SO_4 \rightarrow [RNH_3]_2^+SO_4^{2-}$

322 **(c)**

In aqueous solution, basicity order dimethyl amine > methyl amine > trimethyl amine > aniline

326 **(a)**

Secondary nitroalkanes can be converted into ketones by using aqueous HCI.

$$2 \xrightarrow{R} CHNO_2 + HC1 \longrightarrow 2 \xrightarrow{R} C = O + N_2O + H_2C$$

327 (d)

 $(C_2H_5)_2$ NH (2°amine) is strongest base. Basic nature of amines due to presence of lone pair of electron on nitrogen atom which is available for the bond formation with Lewis acid. Due to the +I effect 2° amine is better base than 1° amine and NH₃. In case of aromatic amines the lone pair on nitrogen atom involved in resonance, therefore, not available for bond formation, so aromatic amines are less basic.

328 (c)

$$R = 0 - N = 0 \xrightarrow{4[H]} R = 0H + NH_2OH$$

330 (a)

N≡C—CH=CH—C≡N; It has five π -bonds.

331 (c)

The basic character of amines depends upon the capacity nitrogen atom to donate an electron pair. More the ability of nitrogen atom to donate electron pair, more will be the basic character. In presence of electron releasing groups (+I showing group, $-CH_3$) the basic character of amines increases due to more availability of electrons on nitrogen atom.

Tertiary amines are least basic due to steric hindrance caused by three bulky alkyl groups.

∴The order of basic character is

$$NH(CH_3)_2 > CH_3NH_2 > N(CH_3)_3 > NH_3$$

332 (a)
$$CH_2N_2 + CH_3COOH \rightarrow CH_3COOCH_3 + N_2$$
Ester

333 **(b)**

The name of isocyanides is carbylamine, and when it is attached with an alkyl group, the compound is called alkyl carbylamine, *i.e.*,RNC.

334 **(c)**

Primary amines react with alc alkali and chloroform to give an offensive odour compound *i.e.*, isocyanide. This reaction is called carbylamine reaction.

$${\rm CH_3CH_2CH_2NH_2+CHCI_3+3KOH} \
ightarrow {\rm CH_3CH_2CH_2NC+3KCI+3H_2O} \ n$$
-propyl amine (alco.) propyl isocyanide

335 **(c)**

Aliphatic and aromatic primary amines on warming with CHCI_3 and alcoholic KOH form isocyanide or carbylamine which has very unpleasant smell. This reaction is known as carbylamine reaction.

Since *p*-toluidine contains an aromatic primary amine group, it undergoes similar reaction and give 4-methyl phenyl isocyanide.

NC

$$+CHCl_3+3(alc)KOH$$

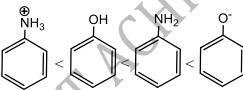
Warm

 $+3KCl+3H_2O$
 $+3C$
 p -toluidine

 $+3KCl+3H_2O$
 $+3C$
 $+$

336 (a)

Coupling of diazonium salts takes place in the following order as



338 (d)

Basicity of amines depends upon the availability of lone pair of electrons of nitrogen for donation. Electron releasing group increases the electron density over nitrogen, thus increases the basic character. 3° methyl amine although contains three electron realeasing groups but is least basic because of steric hindrance. Hence, the order of basic character is

$$(CH_3)_2NH > CH_3NH_2 > (CH_3)_3N$$

341 **(b)**

 $NH_2CONH_2 + HNO_3 \rightarrow NH_2CONH_2 \cdot HNO_3$

344 (d)

Nitrene is not the intermediate of Gabriel's phthalimide reaction

347 (d)

Secondary amines give oily nitrosoamine with nitrous acid

$$(CH_3CH_2)_2NH + HNO_2$$

$$\rightarrow$$
 (CH₃CH₂)₂N - NO + H₂O

Secondary amine nitrous acid oily nitrosoamine

348 **(b)**

Biuret test is given by compounds having — CONH₂gp.

349 **(d)**

Benzyl amine is most basic because positive inductive effect (+1) increases due to presence of methylene group.

350 (a)

Due to sp^3 -hybridisation and lone pair of electron (like NH₃)

351 **(c)**

$$NH_4CNO \xrightarrow{\Delta} NH_2CONH_2 \xrightarrow{\Delta} NH_2CONHCONH_2$$
Urea Biuret

352 **(c)**

Tertiary amines do not have replaceable H-atom.

353 (a)

$$CH_3CH = NOH \xrightarrow{P_2O_5} CH_3CN + H_2O$$

354 **(b)**

H-atom of C_6H_6 ring is replaced by S_E reactions

356 **(b**)

Less substituted alkene is the main product

357 (c)

$$CH_3NC \xrightarrow{Reduction} CH_3NHCH_3$$

360 (c)

$$R_2$$
NH $\stackrel{[0]}{\rightarrow}$ R_2 N—N R_2

361 (a)

Primary nitro alkanes when treated with nitrous acid from nitrolic acid which when dissolve in alkali gives, salt of nitrolic acid which is red in colour. In excess of acid, the salt dissociated, thus colour disappears while in excess of alkali the red colour reappears

363 **(b)**

1° aliphatic amines on reduction with HNO₂ form

(red colour)

alcohol and evolve N₂.

$$RNH_2 + HONO \xrightarrow{NaNO_2 - HCI} ROH + N_2 + H_2$$

365 (a)

Primary amines on heating with chloroform and ethanolic KOH, give alkyl isocyanide. This reaction is called carbylamine reaction.

$$R - NH_2 + CHCI_3 + 3KOH \xrightarrow{\Delta} R - NC + 3KCI + 3H_2O$$

1°amine

alcohol alkyl isocyanide

367 **(b)**

$$HCONHR \xrightarrow{POCl_3} RNC \times H_2O$$

368 **(b)**

—CN and —NC are different functional groups.

369 (a)

As the molecular mass increase, boiling point increases. In case of isometric amines, however, as the number of H-atoms attached directly to Natom decreases, boiling point decreases because tendency to form H-bonds decreases. Hence, the order of boiling points of given amines is $CH_3NH_2 < (CH_3)_3 N < CH_3CH_2N(NH_3)$

$$CH_3NH_2 < (CH_3)_3 N < CH_3CH_2N(NH_3)$$

 $< CH_3CH_2CH_2NH_2$

370 **(c)**

$$C_6H_5CH = O + H_2NCH_3 \rightarrow C_6H_5CH = NCH_3$$

372 **(c)**

$$C_2H_5NH_2 \xrightarrow{[0]} CH_3CH = NH \xrightarrow{HOH} CH_3CHO + NH_3$$

In Hofmann elimination reaction, less substitudes

alkene is the main product

$$\begin{bmatrix} \mathsf{CH_3} \\ \mathsf{CH_3}\mathsf{CH_2}\mathsf{CH_2} - {}^{\mathsf{+}}\mathsf{N} - \mathsf{CH_2}\mathsf{CH_3} \\ \mathsf{CH_3} \end{bmatrix} \mathsf{OH}^{\mathsf{-}} \stackrel{\triangle}{\longrightarrow}$$

$$CH_3CH_2CH_2$$
-N(CH_3)₂ + CH_3CH - CH_2 + CH_2 - CH_2
(minor) major

375 (a)

When propyl bromide is reacted with KCN, butanenitrile is formed.

$$CH_3CH_2CH_2Br + KCN \rightarrow CH_3CH_2CH_2C \equiv N + KBr$$

propyl bromide butane nitrile

376 **(b)**

Ethyl amine reacts with nitrous acid to give ethyl alcohol and nitrogen.

$$C_2H_5NH_2 + HNO_2 \rightarrow C_2H_5OH + N_2 + H_2O$$

377 (c)

Carbylamine reaction is given by only primary amines (both aliphatic and aromatic). In this reation a primary amine reacts with chloroform in basic medium, to form a very bad smelling compound, called carbylamines $C_6H_5NH_2 + CHCl_3KOH \rightarrow C_6H_5NC + KCl + H_2O$