## 13.AMINES

## **Single Correct Answer Type**

- 1. Acetonitrile on reduction gives
  - a) Propanamine
- b) Methanamine
- c) Ethanamine
- d) None of these
- 2. 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
- 3. The reduction of which of the following compound would yield secondary amine?
  - a) Alkyl nitrile

b) Carbylamine

c) Primary amine

- d) Secondary nitro compound
- 4. 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
- 5. From the following compounds which does not react with C<sub>6</sub>H<sub>5</sub>SO<sub>2</sub>CI?
  - a)  $C_2H_5$ .  $NH_2$

b) CH<sub>3</sub>. NH<sub>2</sub>

c)  $(CH_3)_2NH$ 

- d)  $(C_2H_5)_3N$
- 6. 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
- 7. Which of the following reaction will not occur?

a) 
$$\phi N_2^+ + CuBr \xrightarrow{HBr} \phi - Br$$

b) 
$$\phi N_2 + \phi OH \longrightarrow \phi - N = N - \langle O \rangle - OH$$

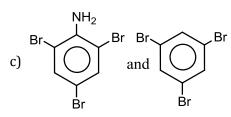
c) 
$$\Phi N_2^+ \xrightarrow{H_3PO_2} \Phi H$$

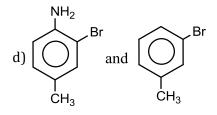
d) 
$$\phi N_2^+ + I^- \rightarrow \phi - I$$

- 8. 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
- 9. In the following reaction sequence predict the compound *X* and *Y*.

The compound *X* and *Y* are

a) 
$$\bigcap_{CH_3}^{NH_2}$$
 and  $\bigcap_{CH_3}^{Br}$   $\bigcap_{CH_3}^{Br}$ 





- 10. The type of isomerism shown by C<sub>6</sub>H<sub>5</sub>CN and C<sub>6</sub>H<sub>5</sub>NC is:
  - a) Position
- b) Functional
- c) Enantiomerism
- d) Tautomerism

11. Identify *A* and *B* in the reaction given below.

Ethane nitrile 
$$\xrightarrow[+2H_2O]{\text{Hydrolysis}} A \xrightarrow[-NH_3]{\text{Decarboxylation}} A \xrightarrow[-CO_2]{\text{Decarboxylation}} B$$

a) Acetic acid, methanol

b) Acetone, methane

c) Ethanoic acid, ethane

- d) Ethanoic acid, methane
- 12. When ethanol is mixed with ammonia and passed over catalyst, the compound formed is:
  - a)  $C_2H_5NH_2$
- b)  $C_2H_4$

- c)  $C_2H_5OC_2H_5$
- d) CH<sub>3</sub>OCH<sub>3</sub>

- 13. 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)  $C_6H_5CH_2CH_2$

14. In the reaction

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

X can be

- a) LiAIH<sub>4</sub>
- b) H<sub>2</sub>SO<sub>4</sub>
- c) Ni

- 15. Increasing order of basicity of  $CH_3CH_2CH_2NH_2$   $H_2C = CHCH_2NH_2$  and  $HC \equiv CCH_2NH_2$  is
  - $CH_3CH_2CH_2NH_2 < HC \equiv CCH_2NH_2 < H_2C =$ a) CHCH NII CHCH<sub>2</sub>NH<sub>2</sub>
- $_{\text{bl}}$  CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub> < H<sub>2</sub>C = CHCH<sub>2</sub>NH<sub>2</sub> < CH
- $HC \equiv CCH_2NH_2 < H_2C = CHCH_2NH_2$ < CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>
- $\begin{aligned} \mathsf{CH} &\equiv \mathsf{CCH_2NH_2} < C\mathsf{H_3CH_2CH_2NH_2} < \mathsf{H_2C} \\ &= \mathsf{CHCH_2NH_2} \end{aligned}$

16. In the compound given below,

$$H_3$$
 $(Y)$ 
 $(Y)$ 
 $(X)$ 
 $(Y)$ 
 $(X)$ 

the correct order of acidic nature of the positions (X), (Y) and (Z) is:

- a) Z > X > Y
- b) X > Y > Z
- c) X > Z > Y
- d) Y > X > Z

- 17. Which of the following is carbamide?
  - a) CH<sub>3</sub>CONH<sub>2</sub>
- b) NH<sub>2</sub>CONH<sub>2</sub>
- c) CH<sub>2</sub>(NH<sub>2</sub>)CONH<sub>2</sub>
- d) CO(OH)NH<sub>2</sub>
- 18. Aniline is not the major product in one of the following reactions. Identify that reaction.

- b)  $C_6H_5NO_2 + Zn$  powder  $\xrightarrow{Alcoholic KOH}$
- a)  $C_6H_5OH + NH_3 \xrightarrow[300^{\circ}C]{200^{\circ}C}$ c)  $C_6H_5Ci + NH_3 \xrightarrow[Cu_2O]{200^{\circ}C}$  High pressure
- d)  $C_6H_5NO_2 + 6(H) \xrightarrow{Fe+H_2O}_{HCI}$
- 19. An organic compound 'A' having molecular formula C<sub>2</sub>H<sub>3</sub>N 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) CH<sub>3</sub>CH<sub>2</sub>NH<sub>2</sub>
- b) CH<sub>3</sub>CH<sub>2</sub>N  $\rightleftharpoons$  C
- c)  $CH_3C \equiv N$
- d) CH<sub>3</sub>CH<sub>2</sub>. OH

20. The IUPAC name of the compound having formula,

$$O = C - CH - CH_2$$
 is  $C - CH - CH_2$  is  $C - CH - CH_2$  OH

- a) 3-aminohydroxy propionic acid b) 2-amino-propan-3-oic acid c) Amino hydroxy propanoic acid d) 2-amino-3-hydroxy propanoic acid Methyl amine reacts with methyl iodic required?
- 21. Methyl amine reacts with methyl iodide. For completion of reaction, how many moles of methyl iodide are required?
  - a) 1

b) 2

c) 3

d) 4

- 22. The IUPAC name of  $CH_3$  CH—  $CH_2$  CH—  $CH_3$  is:  $CH_3$  CN
  - a) 2-cyano-3-methyl hexane
  - b) 2-dimethyl-4-cyanopentane
  - c) 2,4-dimethyl pentane nitrile
  - d) 2-cyano-3-methyl hexane
- 23. Identify *X* in the series,



 $\xrightarrow{\text{HNO}_3} \text{Intermediate} \xrightarrow{\text{H}_2\text{O}} X$ :



 $NH_2$ 



NHCOCH<sub>3</sub>



ŇO



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

Recognize the compound C from the following

- a) Propanenitrile
- b) Methylamine
- c) Ethylamine
- d) Acetamide
- 25. Dichlorocarbene does not form as an intermediate in this reaction
  - a)  $phenol + CHCl_3 + 4KOH$

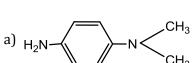
b) Ethyl amine +CHCl<sub>3</sub> + KOH

c) Phenol+CCl<sub>4</sub> + 4KOH

- d)  $CHCl_3 + KOH$
- 26. Which of the following methods neither means for the synthesis nor for separation of amines?
  - a) Hinsberg's method
- b) Hofmann's method
- c) Wurtz reaction
- d) Curtius method

- 27. Reaction of aniline with benzaldehyde is
  - a) Substitution
- b) Addition
- c) Condensation
- d) Polymerisation
- 28. On heating benzyl amine with chloroform and ethanolic KOH, product obtained is

	a) Benzyl alcohol	b) Benzaldehyde	c) Benzonitrile	d) Benzyl isocyanide							
29.	Choose the amide which	on reduction with $\it LiAIH_4$ y	vields a secondary amine								
	a) Ethanamide		b) N-methylethanamide								
	c) N, N-dimethylethanam	iide	d) Phenylmethanamide								
30.	Hofmann bromamide rea	ction is used to prepare									
	a) 1° amine	b) 2° amine	c) 3° amine	d) All of these							
31.	CN	•	•								
	$\frac{\text{CH}_3\text{MgBr}}{\text{dry ether}} X \xrightarrow{\text{H}_2}$	<u>O</u> <b>Y</b>									
	Identify Y										
	a) Benzophenone	b) Acetophenone	c) Benzoic acid	d) phenol							
32.	Which of the following compound does not undergoes Schotten-Baumann reaction?										
	a) Phenol	b) Primary amine									
33.	Arrange the following CI	$H_3NH_2$ (I); $CH_3NH$ (II);	$C_6H_5NH_2$ (III); $(CH_3)_3N$	(IV) in increasing order of							
	basic nature in aqueous medium:										
	<del>-</del>		c) $I < II < III < IV$	d) $II < III < I < IV$							
34.	The reaction,	,									
	$RCOOH \xrightarrow{\text{NaN}_3/\text{conc.H}_2SO_4} R$	NII I N I CO									
		$NH_2 + N_2 + CO_2$									
	is known as	h) I	a) Calauri de ura reti au	d) II - 6							
25	a) Curtius reaction			d) Hofmann reaction							
35.			st with NaNO <sub>2</sub> /HCI and the	n coupled with phenoi							
	produces <i>p</i> -hydroxyazob		) DII	15 A - 11:							
26	a) Nitrobenzene	b) Azobenzene	,	d) Aniline							
36.	<del>-</del>	<del>-</del>	is acid gives an oily nitroso								
0.7	a) Methylamine	b) Ethylamine	, ,	•							
37.			ol in weakly basic medium g								
	a) Diphenyl ether	b) <i>p</i> -hydroxy azobenzene	e c) Chlorobenzene	d) Benzene							
38.											
	$F \sim NO_2 - CH_2$	$A \longrightarrow A$									
	\DI	MF,∠∆									
				N. I							
				CH <sub>3</sub>							



$$O_1 O_2 N$$
 $O_2 N$ 
 $O_2 N$ 
 $O_2 N$ 

d) 
$$O_2N$$
  $NH_2$ 

- 39. 1 mole of ethyl amine on reaction with HNO<sub>2</sub> gives at NTP
  - a) 11.2 L of N<sub>2</sub>
- b) 5.6 L of N<sub>2</sub>
- c) 22.4 L of N<sub>2</sub>
- d) 1 L of N<sub>2</sub>

- 40. Dehydration of an amide gives:
  - a) Cyanide
- b) Amine
- c) Isocyanide
- d) Fatty acid
- 41. In aqueous solutions, the basic strength of amines decreases in the order
  - a)  $CH_3NH_2 > (CH_3)_2NH_2 > (CH_3)_3N$
- b)  $(CH_3)_2NH > (CH_3)_3N > CH_3NH_2$
- c)  $(CH_3)_3N > (CH_3)_2NH > CH_3NH_2$
- d)  $(CH_3)_2NH_2 > CH_3NH_2 > (CH_3)_3N$
- 42. Methyl cyanide gives on hydrolysis
  - a) Methyl amine
- b) Acetic acid
- c) Formic acid
- d) Ethyl amine

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

	a) 1°amine	b) 2°amine	c) 3°amine	d) Quaternary salt						
45.	2,4,6-tribromo aniline is a product of:									
	a) Electrophilic addition on C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub>									
	b) Electrophilic substitution on C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub>									
	c) Nucleophilic addition									
	d) Nucleophilic substitution on $C_6H_5NH_2$									
46	The value of $K_b$ is highes									
10.		b) <i>p</i> -chloroaniline	c) <i>p</i> -nitroaniline	d) p-methylaniline						
47.			c) p-introamme	$u_j p$ -methylannine						
	( ) = /		c) (CH <sub>3</sub> ) <sub>3</sub> NO	4) (CH ) NNO						
10	· · ·	- · · · - · · · · · · · · · · · · · · ·		u) (CH3)2NNO						
40.	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) $C_2H_5NH_2 > CH_3NH_2 > NH_3 > C_6H_5NH_2$									
40	c) $C_6H_5NH_2 > C_2H_5NH_2 > CH_3NH_2 > NH_3$ d) $CH_3NH_2 > C_2H_5NH_2 > C_6H_5NH_2 > NH_3$									
49.	<del>-</del>	perate CO <sub>2</sub> from NaHCO <sub>3</sub> so		12 022 22 22						
	a) CH <sub>3</sub> CONH <sub>2</sub>	b) CH <sub>3</sub> NH <sub>2</sub>	c) $(CH_3)_4N^+OH^-$	d) CH <sub>3</sub> N <sup>+</sup> H <sub>3</sub> CL <sup>-</sup>						
50.	Acetaldoxime reacts wit									
	a) CH <sub>3</sub> CN	b) C <sub>2</sub> H <sub>5</sub> CNO	c) C <sub>2</sub> H <sub>5</sub> CN	d) All of these						
51.	Diethyl oxalate is used for distinguishing primary, secondary and tertiary									
	a) Alcohols b) Amines									
	c) Alkyl halides		d) Hydrogens in hydroca	rbons						
52.	Nitration of aniline in strongly acidic medium, result in the formation of									
	<i>m</i> -nitroaniline also. This is because									
	a) Amino group is <i>meta</i> orienting during electrophilic substitution reaction.									
	b) Nitro group goes always to the <i>meta</i> 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.									
53.	On heating benzyl amine with chloroform and ethanolic KOH, product obtained is									
	a) Benzyl alcohol	b) Benzaldehyde	c) Benzonitrile	d) Benzyl isocyanide						
54.	, ,	•	•							
<i>J</i> 1.	An organic amino compound reacts with aqueous nitrous acid at low temperature to produce an oily nitroso amine. The compound is									
	a) CH <sub>3</sub> NH <sub>2</sub>	odila is	b) CH <sub>3</sub> CH <sub>2</sub> NH <sub>2</sub>							
	c) $(CH_3CH_2)_3N$		d) $CH_3CH_2 - NHCH_2CH_3$							
55.		( = 0) are columble in wate		as U CO thou give						
JJ.	Nitrosoamines ( $R_2N - N = 0$ ) are soluable in water. On heating them with conc $H_2SO_4$ , they give									
	secondary amines. The reaction is called									
	a) Perkin's reaction		b) Fries reaction							
	c) Liebermann nitroso re		d) Etard reaction							
56.			rgoes hydrolysis with alkali							
	a) -CN	b) -CHO	c) -COCH <sub>3</sub>	d) –Br						
57.	<del>-</del>	oducts obtained in the follo	wing reactions is							
	$(A)RCN \xrightarrow{\text{Reduction}} (B) RCN \xrightarrow{(I)CH_3MgBr}$									
	(II)H <sub>2</sub> O									
	$(C)RNC \xrightarrow{Hydrolysis} (D)RNH_2 \xrightarrow{HNO_2}$									
	A B	C D								
	a) 2°amine Methyl ketone	e 1°amine Alcohol	b) 1°amine   Methyl ketone	1°amine Alcohol						
	c) 2°amine Methyl ketone		d) 2°amine   Methyl ketone	2°amine aldehyde						
58.	Which of the following is	not a nitroderivative?								

 $44. \ \ \, The \, molecular \, formula \, C_3H_9N \, cannot \, represent$ 

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- a)  $C_6H_5NO_2$
- b) CH<sub>3</sub>CH<sub>2</sub>ONO
- d)  $C_6H_4(OH)NO_2$

- 59.  $CHCI_3 + C_6H_5NH_2 + 3NaOH \rightarrow A + 3B + 3C$ In the above reaction, the product 'A' is
  - a) Chlorobenzene
- b) Phenyl isocyanide
- c) Phenyl cyanide
- d) Phenyl chloride

- 60. Carbonyl chloride reacts with ammonia to form:
  - a)  $CO_2$

- b) NH<sub>2</sub>CONH<sub>2</sub>
- c) CH<sub>3</sub>COONH<sub>4</sub>
- d) CH<sub>3</sub>CONH<sub>2</sub>

- 61. 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
- 62. The major product of the following reaction is

a) 
$$N - H_2C - Br$$

$$d) \bigcirc N \longrightarrow H_2C \longrightarrow CH_2CI$$

- 63. Diazotisation can be carried out by the action of NaNO<sub>2</sub> and dilute HCl at ice cold temperature on:
  - a) Aromatic secondary amine
  - b) Aromatic primary amine
  - c) Aromatic nitro compound
  - d) Aromatic amine
- 64. The molecular formula of benzonitrile is
  - a) C<sub>6</sub>H<sub>5</sub>CN
- b)  $C_6H_5NC$
- c) C<sub>6</sub>H<sub>5</sub>CNO
- d) C<sub>6</sub>H<sub>5</sub>NCO
- 65. Which of the following reactions is given by only primary amines?
  - a) Reaction with HONO

b) Reaction with chloroform and alcoholic KOH

c) Reaction with acetyl chloride

d) Reaction with Grignard reagent

- 66. The diamide of carbonic acid is:
  - a) Acetamide
- b) Formamide
- c) Benzamide
- d) Urea

- 67. Urea reacts with hydrazine to form:
- a) Nitrogen
- b) Phenyl hydrazine
- c) Semicarbazide
- d) Urethane

- 68. Final product of hydrolysed alkyl cyanide is
  - a) RCOOH
- b) RCONH<sub>2</sub>
- R C = NHc)

OH

69.	=	can be converted into keton	es by using $Y$ . Identify $Y$ from	om the following								
	R CHNO <sub>2</sub> + $Y$	R										
	$\sim$ CHNO <sub>2</sub> + $\gamma$	<b>&gt;</b> c==0										
	/ \	N	a) VMO	4) CO								
70	a) Aqueous HCl	b) Aqueous NaOH	c) KMnO <sub>4</sub>	d) CO								
70.		npounds, the most basic is	a) it ili	J) D								
71	a) Aniline	b) Acetanilide	c) <i>p</i> -nitroaniline	d) Benzyl amine								
71.	<u>-</u>	group by replacing hydroge	en or primary annines:									
<ul><li>a) Increases the base strength</li><li>b) Decreases the base strength</li></ul>												
		rength										
	c) Remains the same d) None of the above											
72		xidation in the presence of l	KMnO.followed by hydroly	rsis to form:								
, 2.	a) An acid	b) An alcohol	c) An aldehyde	d) a N-oxide								
73.	-	ompounds will undergo car	-	aj a iv oxiac								
, 0.	a) (CH <sub>3</sub> CH <sub>2</sub> ) <sub>2</sub> NH	b) (CH <sub>3</sub> ) <sub>2</sub> NH	c) $C_6H_5NH_2$	d) (CH <sub>3</sub> ) <sub>3</sub> N								
74.		forms non-superimposable										
	because:											
	a) Of rapid flipping											
	b) Amines are basic in na	nture										
	c) Nitrogen has a lone pa											
	d) Of absences of asymm											
75.	Urea on heating with eth	<del>-</del>										
	a) Urethane	b) Urea alcohol	c) Ureides	d) None of these								
76.	Aliphatic amines are solu	ıble in water because:										
	a) They are basic											
	b) They are amino compo	ounds										
	c) They are lighter than v	water										
	d) Of formation of hydrog	gen bonds with water										
77.	A positive carbylamines	test is given by										
	a) N, N-dimethylaniline		b) 2,4-dimethylaniline									
	c) N-methy-o -methylani		d) N-methylbenzylamine									
78.	Which of the following an	mines form maximum hydr		ves?								
	a) CH <sub>3</sub> NH <sub>2</sub>	b) $(CH_3)_2NH$	c) $(CH_3)_3N$	d) None of these								
79.	<u> </u>	ne with dimethylamine in th	=									
	<del>-</del>	ng the reaction is continuou	isly removed. The compour	nd formed is generally								
	known as											
	a) An amine	b) An imine	c) An enamine	d) A Schiff's base								
80.		ride on treatment with hypo	o phosphorous acid and wa	iter in presence of Cu <sup>+</sup>								
	catalyst produce	1.) m. 1	) A 11:	1) (1) 1								
01	a) Benzene	b) Toluene	c) Aniline	d) Chlorobenzene								
81.		ng does not have $sp^2$ hybrid		J) A + : J -								
വ	a) Acetone	b) Acetic acid	c) Acetonitrile	d) Acetamide								
82.	The product of Hofmann											
	$\leftarrow$ CH <sub>2</sub> N(CH <sub>3</sub> ) <sub>2</sub> O	oH is										
	a) $\langle -CH_2N(CH_3)_2 \rangle$		b) $\langle \bigcirc \rangle$ NH <sub>2</sub> + (CH <sub>3</sub> )	) <sub>2</sub> N								
	c) CH <sub>2</sub> + (CH <sub>3</sub>	) <sub>2</sub> NOH	d) $\left\langle \begin{array}{c} \\ \\ \\ \end{array} \right\rangle + (CH_3)_4 \stackrel{+}{\text{NOH}}$									

- 83. The best method to synthesise *m*-dibromobenzene is by using the reaction
  - a) Benzene  $\xrightarrow{Br_2/FeBr_3/heat}$

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

Nitrobenzene

- C)  $\xrightarrow{\text{Fuming HNO}_3}$  []  $\xrightarrow{\text{Fe/HCl}}$  []  $\xrightarrow{\text{1.HONO}}$  2.CuBr
- $\frac{\text{d)} \xrightarrow{\text{HNO}_3}}{\text{H}_2\text{SO}_4} \left[ \right] \xrightarrow{\text{Fe/HCl}} \left[ \right] \xrightarrow{\text{1.HONO}} 2.\text{CuBr}$
- 84. Aniline is reacted with Br<sub>2</sub> 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
- 85. 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 stronger than aniline, but weaker base than ammonia
- 86. RNH<sub>2</sub> reacts with C<sub>6</sub>H<sub>5</sub>SO<sub>2</sub>Cl 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) 
$$R - NH - SO_2 - C_6H_5$$

- 87. When  $NaNO_2$  and dilute HCl were added to an amine at  $0^{\circ}$ C, a colourless gas was evoloved and an ionic compound is formed. The amine is:
  - a) An primary amine
  - b) An aromatic primary amine
  - c) Any amine
  - d) None of the above
- 88. Choose the incorrect comparision(s)

b)  $CH_3CH_2CH_2NH_2 > (CH_3)_3N$  (basicity in aqueous medium)

(basicity in aqueous medium)

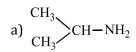
d)  $\frac{3 N}{2 N}$  N(1) > N(3)(basicity aqueous medium)

- 89. Grignard reagent and acetyl chloride does not react with:
  - a)  $RNH_2$
- b)  $R_2$ NH
- c)  $R_3N$

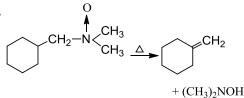
- d) None of these
- 90. Which of the following can be used to distinguish acetamide and urea?
  - a) Fehling's solution
- b) Biuret test
- c) Hofmann's reaction
- d) NaOH solution

- 91. Among the amines  $(A)C_6H_5NH_2$ ,  $(B)CH_3NH_2$ , (C) $(CH_3)_2NH$ ,  $(D)(CH_3)_3N$ , the order of basicity is
  - a) A < B < D < C
  - b) D < C < B < A
  - c) A < B < C < D
  - d) B < C < D < A
- 92. Choose the incorrect statement
  - a) In the case primary, secondary and tertiary amines, the basic strength depends on the extent on the extent of hydrogen bonding in the protonated amines

	1.) ml	lel ocu l cu :		<i>c</i> .
	<del>-</del>	groups like – $OCH_3$ and – $CH_3$ in	_	
	<del>-</del>	groups like – $NO_2$ , –CN and hal	=	ength of amines
02	,	th of amines depends on their o		ture to anoduce on eiler
93.	nitrosoamine. The	ompound reacts with aqueous	mitrous acid at low tempera	ture to produce an ony
	a) CH <sub>3</sub> NH <sub>2</sub>	b) CH <sub>3</sub> CH <sub>2</sub> NH <sub>2</sub>	c) CH <sub>3</sub> CH <sub>2</sub> NHCH <sub>2</sub> CH <sub>3</sub>	4) (CH CH ) N
94.		with cyanogen chloride gives:	c) GH3GH2NHGH2GH3	u) (61136112)31V
<i>)</i> 1.	a) R—NC	b) R—Cl	c) R—CN	d) None of these
95.	•	hydrolysis in acidic medium g	•	a) None of those
	= =	and methanoic acid		mmonium salt
	c) Ethanoic acid an		d) Methyl amine salt and	
96.	=	ring will give a primary amine o		
	a) Nitroparaffin	b) Alkyl cyanide	c) Oxime	d) Alkyl isocyanate
97.	Urea when heated	a white residue is formed. Its	alkaline solution when trea	ted with few drops of CuSO <sub>4</sub>
	solution gives:			
	a) Red colour	b) Violet colour	c) Green colour	d) Yellow colour
98.	Which one of the fo	llowing is most basic?		
	a) FCH <sub>2</sub> NH <sub>2</sub>	b) FCH <sub>2</sub> CH <sub>2</sub> NH <sub>2</sub>	c) $C_6H_5NH_2$	d) C <sub>6</sub> H <sub>5</sub> CH <sub>2</sub> NH <sub>2</sub>
99.	The basicity of com	pounds I, II, III and IV		
	$CH_3NH_2$ , $(CH_3)_2NH$	$(CH_3)_3N, C_6H_5CH_2NH_2$		
	I II	III IV		
	varies in the order			
	a) $I > II > III > IV$	VI $<$ III $>$ IV	c) $III > I > II > IV$	d) $IV > I > II > III$
100	. A gaseous carbon c	ompound is soluable in dilute I	HCI. The solution on treating	g with NaNO <sub>2</sub> gives off
		hind a solution which smells of	f wood spirit. The carbon co	<del>-</del>
	а) НСНО	b) CO	c) $C_2H_5NH_2$	d) CH <sub>3</sub> NH <sub>2</sub>
101	. The correct order o	of basic nature of the following	compounds is:	
	NH	CH <sub>3</sub> CH <sub>2</sub> NH <sub>2</sub>		
	CH <sub>3</sub> -C	(2)		
	$^{NH_2}$			
	( )			
	$(CH_3)_2NH$	$CH_3$ - $C$ - $NH_2$		
	(3)	(4)		
	a) 2 > 1 > 3 > 4			
	b) $1 > 3 > 2 > 4$			
	c) $3 > 1 > 2 > 4$			
	d) $1 > 2 > 3 > 4$			
102	,	of amines can be explained:		
		s and Arrhenius concept		
	•	Lowry Bronsted concept		
		and Lowry Bronsted concept		
	d) Only in Lewis co			
103	· •	the reaction of sodium metal o	n ethyl amine is:	
	a) N <sub>2</sub>	b) C <sub>2</sub> H <sub>2</sub>	c) H <sub>2</sub>	d) CO <sub>2</sub>
104	, <u>-</u>	de is hydrolysed in presence of	, <u>-</u>	
	a) Acetamide	b) Methane	c) $CO_2 + H_2O$	d) Acetic acid
105	. An organic compo	und $(C_3H_9N)$ (A), when treat	ed with nitrous acid, gave	an alcohol and N2 gas was
	evolved. (A) on	warming with CHCl <sub>3</sub> and c	caustic potash gave (C)	which on reduction gave
	isopropylmethylam	nine. Predict the structure of ( $A$	).	



106.



This reaction is called

- a) Cope reaction
- b) Ritter reaction
- c) Schmidt reaction
- d) Gabriel reaction
- 107. Which of the following regents will be useful as the basic for a simple chemical test to distinguish between?

$$H_3C-\bigcirc -NH_2$$
 and  $\bigcirc -CH_2NH_2$ 

a) C<sub>6</sub>H<sub>5</sub>SO<sub>2</sub>Cl and OH<sup>-</sup> in H<sub>2</sub>O

b) HONO, then β-naphthol

c) Dilute HCl

- d) AgNO<sub>3</sub> in H<sub>2</sub>O
- 108. Alkanamide, which on Hofmann's reaction gives 1-phenylethylamine, is:
  - a) 2-phenylpropanamide
  - b) 3-phenylpropanamide
  - c) 2-phenylethanamide
  - d) N-phenylethanamide
- 109. Reduction of nitrobenzene in the presence of Zn/NH<sub>4</sub>Cl gives
  - a) Azobenzene

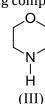
b) Hydrazobenzene

c) N-phenyl hydroxylamine

- d) Aniline
- 110. 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>IV
- 111. Which of the following reactions can be used to prepare ethyl isocyanide?
  - a)  $CH_3CH_2I + NaCN \xrightarrow{C_2H_5OH/H_2O}$

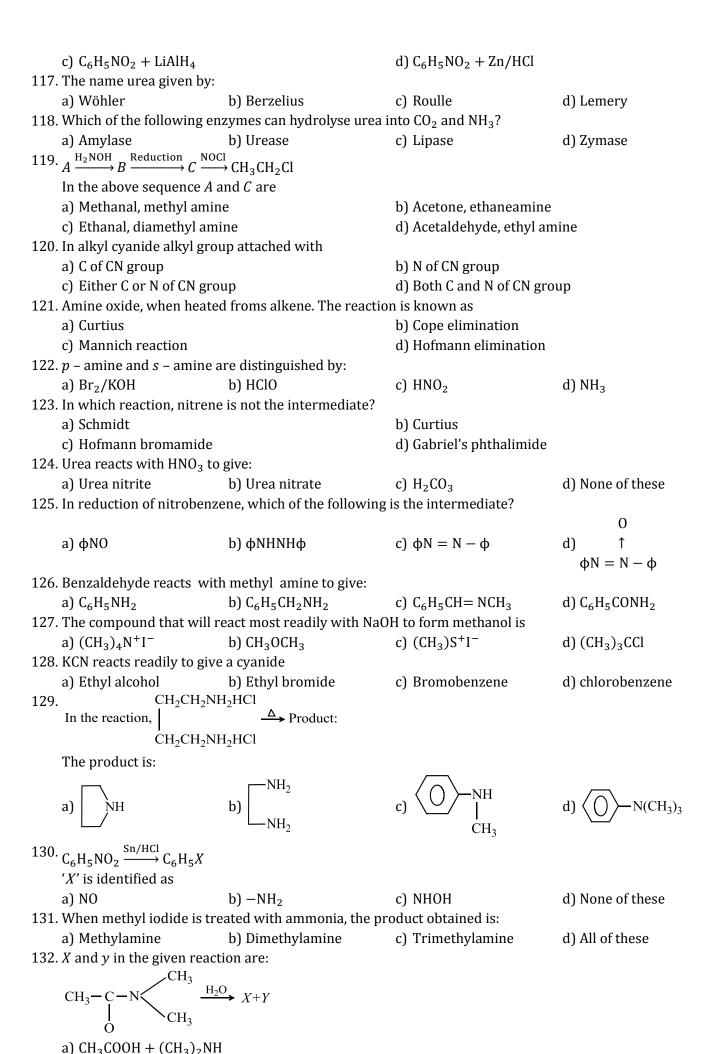
- b)  $CH_3CH_2I + KCN \xrightarrow{Alcohol}$
- c)  $CH_3CH_2NH_2 + CHCL_3 + KOH \xrightarrow{Alcohol}$
- d) None of the above
- 112. An amine reacts with C<sub>6</sub>H<sub>5</sub>SO<sub>2</sub>Cl and the product is soluble in alkali, amine is:
  - a) 1

b) 2°

c) 3°

- d) All of these
- 113. 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) RNH<sub>2</sub>
- c)  $RNH_2 + HCOOH$
- d)  $RCOOH + N_2$
- 114. The product obtained when methylamine is treated with nitrous acid is:
  - a) CH<sub>3</sub>OH
- b) CH<sub>3</sub>—ONO
- c) CH<sub>3</sub>OCH<sub>3</sub>
- d) Both (b) and (c)
- 115. Correct order of basic nature of  $CH_3NH_2(A)$ ,  $CH_3CN(B)$  and  $CH_3N = CHCH_3(C)$  is
  - a) A > B > C
- b) B > C > A
- c) A > C > B
- d) C > A > B
- 116. Which of the following reactions can produce aniline as main product?
  - a)  $C_6H_5NO_2 + Zn/KOH$

b)  $C_6H_5NO_2 + Zn/NH_4Cl$ 



- b)  $CH_3CONH_2 + CH_3OH$
- c)  $CH_3CHO + (CH_3)_2NH$
- d)  $CH_3COCH_3 + CH_3NH_2$
- 133. The correct sequence of reactions to convert p-nitrophenol into quinol involves
  - a) Reduction, diazotization and hydrolysis
- b) Hydrolysis, diazotization and reduction
- c) Hydrolysis, reduction and diazotization
- d) Diazotization, reduction and hydrolysis
- 134. Reaction of benzaldehyde with methylamine gives
  - a)  $C_6H_5COOH$

b)  $C_6H_5N = NCl$ 

c)  $C_6H_5 - CH = N - CH_3$ 

- d)  $C_6H_5NH_2$
- 135. The compound formed when malonic ester reacts with urea is:
  - a) Cinnamic acid
- b) Butyric acid
- c) Barbituric acid
- d) Crotonic acid
- 136. A colourless organic compound gave brisk effervescence with a mixture of NaNO<sub>2</sub> and dil. HCl. It could be:
  - a) Glucose
- b) Oxalic acid
- c) Urea

- d) Benzoic acid
- 137. A colourless, odourless and non-combustible gas is liberated when ethylamine reacts with:
  - a) NaOH
- b) CH<sub>3</sub>COCl
- c)  $NaNO_2 + HCl$
- d)  $H_2SO_4$
- 138. The compound obtained by heating a mixture of  $1^{\circ}$  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
- 139.  $CH_3CONH_2 + NaOH \rightarrow CH_3COONa + A$

Urea is obtained if product if product 'A' in the above reaction reacts with the following compound

- a) Ethyl carbonate
- b) Ethyl urethane
- c) Phosgene
- d) All of these
- 140. Which of the following amines can be directly oxidized to the corresponding nitro compound by potassium permanganate?

$$CH_2 - CH - CH_3$$

- a) CH<sub>3</sub>NH<sub>2</sub>
- $\begin{array}{c} \mathrm{CH_2} \mathrm{CH} \mathrm{CH_3} \\ | \\ \mathrm{NH_2} \end{array}$ b)
- c)  $(CH_3)_2NH$
- d)  $(CH_3)_3C NH_2$
- 141. Which one of the following compound when heated with KOH and primary amines gives carbylamine test?
  - a) CHCl<sub>3</sub>
- b) CH<sub>3</sub>Cl
- c) CCl<sub>4</sub>

d) CH<sub>3</sub>NC

- 142. The hydrochlorides of amines form double salt with:
  - a) PtCl<sub>4</sub>

- b) AuCl<sub>3</sub>
- c) Both (a) and (b)
- d) None of these

143. 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  $\rightarrow$  H<sub>2</sub>N - NHCONH<sub>2</sub> + NH<sub>3</sub>

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

a) CH<sub>3</sub>CHO, NH<sub>2</sub> - NH<sub>2</sub>and PCl<sub>5</sub>

- b) C<sub>6</sub>H<sub>5</sub>CHO, NH<sub>2</sub> NH<sub>2</sub>and SOCl<sub>2</sub>

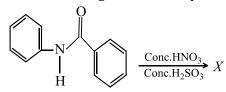
c) C<sub>6</sub>H<sub>5</sub>CHO, NH<sub>2</sub> – NH<sub>2</sub>and NOCl

d) CH<sub>3</sub>CHO, NH<sub>2</sub> - NH<sub>2</sub>and PCl<sub>3</sub>

- 144. Nitroparaffins on reduction give:
  - a) Amides
- b) Alkylamines
- c) Ammonium salts
- d) Acetanilides

- 145. Aniline is prepared in presence of Fe/HCI from
  - a) Benzene
- b) Nitrobenzene
- c) Dinitrobenzene
- d) None of these

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



a) 
$$NO_2$$
 $NO_2$ 
 $NO_2$ 
 $NO_2$ 
 $NO_2$ 
 $NO_2$ 
 $NO_2$ 
 $NO_2$ 
 $NO_2$ 
 $NO_2$ 

147. 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

148. 
$$C_6H_5NH_2 \xrightarrow{H_2SO_4} H_2NC_6H_4(SO_3H_2)$$
(para)

The true statement about the product is

- a) It does not exist as Zwitter ion
- b) -NH2displays a powerful basic character
- c) It does not act as inner salt
- d) -SO<sub>3</sub> dimminishes the basic character of -NH<sub>2</sub>

$$Ac_{2}O \rightarrow A \xrightarrow{Br_{2}} B\xrightarrow{H_{2}O} B\xrightarrow{H_{2}O}$$

$$CH_{3}$$

The final product C in the above reaction is

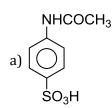
$$\begin{array}{c}
\mathsf{NH}_2\\\mathsf{H}_3\\\mathsf{CH}_3
\end{array}$$

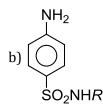
150. Which of the following compound reacts with chloroform and a base to form phenyl isocyanide?

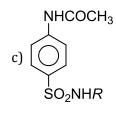
here Xis

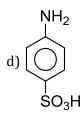
- a) Phenol
- b) Aniline
- c) Benzene
- d) Nitrobenzene

 $(i) (CH_3CO)_2O (ii) HOSO_2CI \longrightarrow [X];$   $(iii) RNH_2 (iv) dil. HCl, \triangle$ 









- 152. Amine is not formed in the reaction
  - (A) Hydrolysis of RCN
  - (B) Reduction of RCH = NOH
  - (C) Hydrolysis of RNC
  - (D) Hydrolysis of RCONH<sub>2</sub>

The correct answer is

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

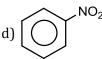
c) B, C

d) A, B, C

153. Aniline reacts with conc. HNO<sub>3</sub> to give

a) 
$$H_2N$$
— $NH_2$ 

b) 
$$NH_2$$
  $NO_2$  and  $NO_2$   $NO_2$ 



- 154. 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
- 155. 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)  $C_6H_5N_2^+CI^- + SnCI_2/HCI \rightarrow C_6H_5NHNH_2$

- 156. The geometry of ethylamine is:
  - a) Pyramidal
- b) Tetrahedral
- c) Triangular
- d) Square planar

157. 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

158. 
$$\left[ \bigcap_{H_2O} \right] \xrightarrow{I^-} Ag_2O \xrightarrow{X} \xrightarrow{400 \text{ K}}$$

The products of above sequence of reactions are

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

b) CH<sub>3</sub>CH<sub>2</sub>CN and C<sub>2</sub>H<sub>5</sub>NH<sub>2</sub>

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

- d)  $(CH_3)_2 C = CH_2$  and  $NH_3$
- 159. Which of the following is hydrolysed to give secondary amine?
  - a) Alkyl
- b) HCON CH<sub>3</sub>
- c) Nitroparaffins
- d) Acidamide

- 160. Carbylamine reaction is given by aliphatic
  - a) Primary amine

b) Secondary amine

c) Tertiary amine

- d) Quaternary ammonium salt
- 161. Ethyl amine reacts with nitrous acid to form
  - a)  $C_2H_5OH$
- b) C<sub>2</sub>H<sub>5</sub>OH, N<sub>2</sub>, H<sub>2</sub>O
- c)  $C_2H_5N_2^+CI^-$
- d) C<sub>2</sub>H<sub>5</sub>NHOH, NH<sub>3</sub>

- 162. 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

- 163. 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 164. Which of the following reagents will convert nitromethane into methylamine? a) Zn/HCl b) Zn/NaOH c)  $Zn/C_2H_5OH$ d) Ni/H<sub>2</sub> 165. Tertiary nitroalkane cannot tautomerise because a) Their tautomeric forms are highly unstable b) They do not contain any multiple bond c) They do not have labile H-atom d) They are not basic in nature NH<sub>2</sub> OH 166. [X] and [Y] are NH<sub>2</sub> OH 167. In the reaction  $NO_2$ Xis a) SiC b) H<sub>2</sub>SO<sub>4</sub> c) KMnO<sub>4</sub> d) Fe/HCl caustic potash produced C. C on treatment with NaNO2 and HCl at 0°C and then boiling produced orthocresol. Compound A is: a) o-toluic acid c) o-bromotoluene d) m-toluic acid b) o-chlorotoluene 169. The correct order of basicities of the following compound is
- 168. A compound A when reacted with PCl<sub>5</sub> and then with ammonia gave B. B when treated with bromine and

c) 3>1>2>4

CH<sub>3</sub>-C
$$\stackrel{\text{NH}}{\stackrel{\text{NH}_2}}{\stackrel{\text{NH}_2}{\stackrel{\text{NH}_2}{\stackrel{\text{NH}_2}{\stackrel{\text{NH}_2}{\stackrel{\text{NH}_2}}{\stackrel{\text{NH}_2}{\stackrel{\text{NH}_2}}{\stackrel{\text{NH}_2}{\stackrel{\text{NH}_2}}}}\stackrel{\text{NH}_2}}{\stackrel{\text{NH}_2}}{\stackrel{\text{NH}_2}}}\stackrel{\text{NH}_2}}{\stackrel{\text{NH}_2}}{\stackrel{\text{NH}_2}}}\stackrel{\text{NH}_2}}{\stackrel{\text{NH}_2}}{\stackrel{\text{NH}_2}}}\stackrel{\text{NH}_2}}{\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}{\stackrel{\text{NH}_2}}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}\stackrel{\text{NH}_2}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}\stackrel{\text{NH}_2}}\stackrel{\text{NH}_2}\stackrel{\text{NH}_2}\stackrel{\text{N$$

(CH<sub>3</sub>)<sub>2</sub>NH 
$$CH_3$$
– $C$ – $NH_2$ 
(3)

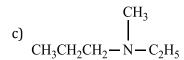
170. Which of the following amines is optically active?

b) 1>3>2>4

- a) CH<sub>3</sub>NH<sub>2</sub>
- b) CH<sub>3</sub>NHCH<sub>3</sub>

a) 2>1>3>4

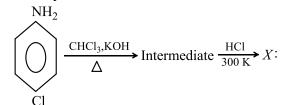
d) 1>2>3>4

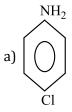


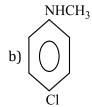
- d) Sec. butylamine
- 171. CH<sub>3</sub>CH<sub>2</sub>NH<sub>2</sub> contains a basic NH<sub>2</sub> group, but CH<sub>3</sub>CONH<sub>2</sub> does not, because:
  - a) Acetamide is amphoteric in character
  - b) In CH<sub>3</sub>CH<sub>2</sub>NH<sub>2</sub> the electron pair on N-atom is delocalised by resonance
  - c) In  $CH_3CH_2NH_2$  there is no resonance, while in acetamide the lone pair of electron on N-atom is delocalised and therefore less available for protonation

  - d) None of the above
- 172. Benzaldehyde condenses with N, N-diamethylaniline in presence of anhydrous ZnCl<sub>2</sub> to give
  - a) Azo dye
- b) Malachite green
- c) Michler's ketone
- d) Buffer yellow

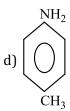
173. Identify *X* in the reaction,





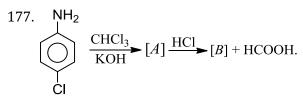






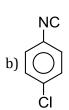
- 174. Aliphatic nitriles are prepared by the treatment of alkyl halides with
  - a) Sodium cyanide
- b) Sodium isocyanide
- c) Sodium isocyanate
- d) Cyanamide

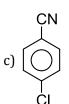
- 175. Diethylamine on oxidation with KMnO<sub>4</sub> gives:
  - a) Ethanal
- b) Propanone
- c) Tetraethyl hydrazine
- d) None of these
- 176. Which one of the following amines will not react with HNO<sub>2</sub> acid to give nitrogen?
  - a) CH<sub>3</sub>NH<sub>2</sub>
- b) CH<sub>3</sub>CH<sub>2</sub>NH<sub>2</sub>
- c) CH<sub>3</sub>-CH-NH<sub>2</sub> CH<sub>3</sub>

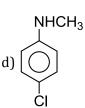


What is [B]?









- $178. C_5H_{13}N$  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
- 179. Which of the following is not a nitro-derivative?

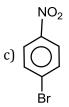
	// <sup>0</sup>							
a) C <sub>6</sub> H <sub>5</sub> NO <sub>2</sub>	b) CH <sub>3</sub> CH <sub>2</sub> ONO	c) CH <sub>3</sub> CH—N	d) C <sub>6</sub> H <sub>5</sub> (OH)NO <sub>2</sub>					
180. Hinsberg's reagent	is	v						
a) C <sub>6</sub> H <sub>5</sub> COCI	b) CH <sub>3</sub> COCI	c) C <sub>6</sub> H <sub>5</sub> CH <sub>2</sub> CI	d) C <sub>6</sub> H <sub>5</sub> SO <sub>2</sub> CI					
181. Among the followin		-7 -0 3 - 2 -	- 7 - 0 - 3 2 -					
a) C <sub>6</sub> H <sub>5</sub> CH <sub>2</sub> NH <sub>2</sub>	b) C <sub>6</sub> H <sub>5</sub> CH <sub>2</sub> NHCH <sub>3</sub>	c) O <sub>2</sub> NCH <sub>2</sub> NH <sub>2</sub>	d) CH <sub>3</sub> NHCHO					
182. A secondary amine		-	- J - 3					
<del>-</del>	h two —NH <sub>2</sub> groups							
	h 2 carbon atoms and a $-NH_2$	group						
	h a $-NH_2$ group on the carbor		on					
	which 2 of the hydrogens of NF	<del>-</del>						
	s with nitrous acid to give		J J J I					
a) Azobenzene	b) Benzene	c) Benzyl alcohol	d) Phenol					
184. Urea is not used:	,	., . ,						
a) As fertilizer								
b) In manufacture o	of plastic							
c) In preparation of	_							
d) In purification of								
	ing statements is not correct?							
	show intermolecular hydrogen	ı bonding						
	es show intermolecular hydrog	=						
= = =	show intermolecular hydrogen	<del>-</del>						
		=	carboxylic acids of comparable					
molar masses	or some bornes as compared		on sony no delue of comparable					
	ula of methyl amino methane	is:						
a) $(CH_3)_2CHNH_2$	b) (CH <sub>3</sub> ) <sub>3</sub> N	c) (CH <sub>3</sub> ) <sub>2</sub> NH	d) CH <sub>3</sub> NH <sub>2</sub>					
187. Aniline and ethylar		0) (0113)21111	a) 31131112					
a) Solubility								
b) Action with HNO	2							
c) Action of Grignar	=							
d) Coupling reaction	_							
,	sequence of reagent in the Ho	fmann's degradation reac	tion?					
a) Br <sub>2</sub> , KOH, H <sub>2</sub> O	b) KOH, Br <sub>2</sub> , H <sub>2</sub> O	c) H <sub>2</sub> O, KOH, Br <sub>2</sub>	d) KOH, H <sub>2</sub> O, Br <sub>2</sub>					
	and tertiary nitroalkanes can	·						
	(q.) b) CHCl <sub>3</sub> + NaOH( $(aq.)$ )	=						
	and tertiary amines may be s		d wone of these					
a) Ethanoyl chlorid		c) Thionyl chloride	d) None of these					
•	produced in Hofmann's broma	•	u) None of these					
a) Br <sup>-</sup>	noduced in Hollianii 3 brollia	iniae reaction is.						
b) Br <sub>2</sub>								
c) OBr <sup>-</sup>								
d) OBr <sub>2</sub>								
· -	amnound turns rad with the a	ddition of a concentrated	NaOH solution, followed by the					
<del>=</del>	=		_					
	_		ur disappears with the addition					
	icid but reappears if the soluti		<del>-</del>					
a) CH <sub>3</sub> CH <sub>2</sub> NO <sub>2</sub>	b) (CH <sub>3</sub> ) <sub>2</sub> CHNO <sub>2</sub>	c) $(CH_3)_3CNO_2$	d) All of these					
193. Isopropylamine KM	$\xrightarrow{\text{nO}_4} X \xrightarrow{\text{H}_3\text{O}^+} Y$ . In the above sec	quence $X$ and $Y$ are respec	tively					

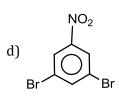
	a) Acetaldimine, ethanal	1	_	Ethanal, ketimine			
104	c) Ketimine, acetone	nuonauad her	a)	Acetone, propan-2-ol			
194.	Benzyl amine cannot be	prepared by					
	a) $C_6H_5CONH_2 \xrightarrow{\text{LiAlH}_4}$		b)	$C_6H_5CH_2CONH_2 + Br_2$	+ KOH →		
	c) $C_6H_5CN \xrightarrow{LiAlH_4}$		d)	$C_6H_5CH_2NC \xrightarrow{LiAlH_4}$			
195	When $(NH_4)_2SO_4 + KCN$	NO are heated we get:	,	061150112110			
175.	a) Nitrogen	to are neated, we get.					
	b) Carbon dioxide						
	c) Biuret						
	d) Ammonium carbonat	e					
196.	=	s most basic in aqueous med	lium	n?			
	a) CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> NH <sub>2</sub>	•	b)	$CH_2 - CH_2 - CH_2 - NH$	2		
	-			CH <sub>3</sub>			
	CH <sub>3</sub> c) CH <sub>3</sub> -N-CH <sub>3</sub>		d)	CH <sub>3</sub> CH <sub>3</sub> -CH <sub>2</sub> -NH-CH <sub>3</sub>			
197	9	ith carboxylic acids to produc		3 2 0			
177.	a) Ester	b) Alcohol		Amine	d) Imines		
198.	•	eparate amines is based on th	-		a) mines		
	a) Benzene sulphonyl ch	-					
	b) Benzene sulphonic ac						
	c) Ethyl oxalate						
	d) Acetyl chloride						
199.	CH <sub>3</sub> CONH <sub>2</sub> . Br <sub>2</sub> and KOI	H give CH <sub>3</sub> NH <sub>2</sub> as the produc	ct. T	the intermediates of the	reaction are		
	(A) CH <sub>3</sub> -C-NHBr						
	(A) $CH_3$ —C—NПБІ	$B) CH_3 - N = C = 0$					
	(C)CH <sub>3</sub> NHBr	(D) CH <sub>3</sub> CONBr <sub>2</sub>					
	The correct answer is	13.4.6					
200	a) A, B	b) A, C	c)	C, D	d) B, D		
200.	Which will not go for dia	azotization?		⊔ N	⊔N		
				H <sub>2</sub> N	H <sub>2</sub> N		
	a) $C_6H_5NH_2$	b) C <sub>6</sub> H <sub>5</sub> CH <sub>2</sub> NH <sub>2</sub>	c)	$ ho C_6 H_4$	d) $C_6H_4$		
				H <sub>3</sub> C	$O_2N$		
201.	Amine may contain:			-	_		
	a) —NH <sub>2</sub> gp	b) >NH gp.	c)	$\rightarrow$ N gp.	d) All of these		
202.		gives $m$ -nitro aniline in stro		,			
		itution reaction amino group	_		nitro group always goes to		
	a) is <i>meta</i> directive	0 1	b)	<i>m</i> -position	0 1 7 0		
	c) In strong acidic media	um aniline aniline present as	s d)	•			
	anilinium ion	•					
203.	In the reaction RCONH <sub>2</sub>	$+ X \rightarrow RNH_2$ , the regent X is	S				
	a) Soda lime	b) PCl <sub>5</sub>	c)	NaOBr	d) All of these		
204.	The bad smelling substa	nce formed by the action of	alco	holic caustic potash on o	chloroform and aniline is		
	a) Nitrobenzene	b) Phenyl isocyanide	c)	Phenyl cyanide	d) Phenyl isocyanate		
205.	How may primary amin	es are possible for the formu	ıla C	$L_4H_{11}N$ ?			
	a) 1	b) 2	c)	3	d) 4		
206.	$\frac{\text{HNO}_3}{\text{H}_2\text{SO}_4} A \frac{\text{Br}_2}{\text{FeBr}}$	$\stackrel{\circ}{\longrightarrow} B$					

The compound B is









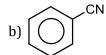
- 207.  $R N = C + HgO \longrightarrow A + Hg_2O$ ; What is A?
  - a)  $RNH_2$
- b) RCONH<sub>2</sub>
- c) R-NCO
- d) RCOOH
- 208. Which of the following on reduction with LiAlH<sub>4</sub> gives a secondary amine?
  - a) CH<sub>3</sub>NC
- b) CH<sub>3</sub>CONH<sub>2</sub>
- c) CH<sub>3</sub>CN
- d) CH<sub>3</sub>NO<sub>2</sub>
- 209. Aniline on treatment with NaNO<sub>2</sub> in HCI at 0°C followed by treatment with alkaline  $\beta$  –naphthol gives
  - a) A violet solution

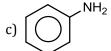
b) A red solution

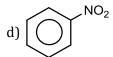
c) A green solution

- d) A blue precipitate
- 210. The compound with foul odour among the following is









- 211. 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

212.  $F \longrightarrow NO_2$ DMF  $\downarrow (CH_3)_2 NH$   $A \xrightarrow{\text{(i) NaNO}_2/HCl, 0-5°C} B$ 

In the above sequence *B* is

$$a)$$
  $H_2N O$   $N< CH_3$   $CH_3$ 

b) 
$$H_2N-\langle \bigcirc \rangle -NH_2$$

$$C_1$$
  $O_2N$   $O$ 

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

213. The product *A* and *B* in the reaction are:

$$C_2H_5NH_2 + CHCl_3 + 3KOH \rightarrow A + B + 3H_2O$$

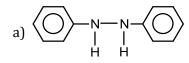
- a)  $C_2H_5NC + 3KCl$
- b)  $C_2H_5CN + 3KCl$
- c)  $C_2H_5CONH_2 + 3KCl$
- d)  $C_2H_5NC + K_2CO_3$
- 214. When aqueous solution of benzene diazonium chloride is boiled, the product formed is
  - a)  $C_6H_5CH_2OH$
- b)  $C_6H_6 + N_2$
- c)  $C_6H_5COOH$
- d)  $C_6H_5OH$

215. Correct order of basicity of  $\phi$  NH<sub>2</sub> [A],  $\langle OCH_2 \rangle$  OCH<sub>2</sub>

$$NH_2[C]$$
  $H_3CO$   $NH_2[D]$  is DCH<sub>3</sub>

- a) A > B > C > D
- b) D > C > A > B
- c) B > D > C > A
- d) D > A > B > C

- 216. Tertiary amine is obtained in the reaction
  - a) Aniline  $\xrightarrow{CH_3I} \xrightarrow{CH_3I}$
- b) Aniline  $\stackrel{CH_3I}{\longrightarrow}$
- c) Nitrobenzene  $\xrightarrow{Sn/HCl}$
- d) None of these
- 217. The structure of the compound formed, when nitrobenzene is reduced by lithium aluminium hydride  $(\text{LiAIH}_4)$  is





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



- 218. Why do 2° and 3° amines fail to undergo the carbylamines 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
- 219. Nitrogen of nitrobenzene at 125°C with mixed acids gives
  - a) meta-dinitrobenzene

b) *ortho*-dinitrobenzene

c) para-dinitrobenzene

d) 1, 3, 5-trinitrobenzene

220. Complete the following reaction

 $RNH_2 + H_2SO_4 \rightarrow$ 

- a)  $[RNH_3]^+HSO_4^-$
- b)  $[RNH_3]_2^+SO_4^{2-}$
- c) RNH<sub>2</sub>. H<sub>2</sub>SO<sub>4</sub>
- d) No reaction

- 221. Ethylamine reacts with nitrosyl chloride (NOCl)to form:
  - a) Ethyl chloride
- b) Ethyl alcohol
- c) Ethyl nitrite
- d) Nitroethane
- 222. Reduction of aniline with acetyl chloride in presence of NaOH produce
  - a) Aniline hydrochloride b) Acetanilide
- c) *p*-choloroaniline
- d) A red dye
- 223. 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) -NH<sub>2</sub> because NH<sub>3</sub><sup>+</sup>, 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
- 224. p-chloro aniline and anilinium hydrogen chloride can be distinguished by
  - a) Sandmeyer reaction

b) Carbylamines reaction

c) Hinsberg's reaction

- d) AgNO<sub>3</sub>
- 225. Nitrobenzene on reduction with Al-Hg and water gives:
  - a) Azobenzene
  - b) Aniline
  - c) Azoxy benzene
  - d) phenylhydroxylamine
- 226. Which of the following is most basic in nature?

- b) CH<sub>3</sub>NH<sub>2</sub>
- c)  $(CH_3)_2NH$
- d)  $C_6H_5N(CH_3)_2$
- 227. The reaction of CHCI<sub>3</sub> and alcoholic KOH with *p*-toluidine gives

228. The IUPAC name of , $CH_3-CH_2-CH-NH_2$  is :



- a) 1-methyl-amino propane
- b) 2-aminobutane
- c) 2-methyl-2-aminopropane
- d) None of the above

229. a) Chain b) Functional c) Position d) All of these 230. Which of the following is not used for nitration of organic compounds? A mixture of concentrated HNO<sub>3</sub> and concentrated by A mixture of concentrated HNO<sub>4</sub> and acetic c) Fuming nitric acid and concentrated sulphuric d) Alcoholic potassium nitrate 231. *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) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>CN b)  $(CH_3)_2CHCN$ c) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>NC d)  $(CH_3)_2CHNC$ 232. *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 < Ic) II < III < Ib) I < II < IIId) II < I < III233. Aniline reacts with acetaldehyde to form a) Schiff's base b) Carbylamine c) Immine d) None of these 234. Coupling of diazonium salts of following takes place in the order  $NH_3$  $NH_2$ IV II Ш a) IV < II < III < Ib) IV > III < II < Ic) II < IV < I < IIId) I < II < III < IV235. The general formula of quaternary ammonium compound is: b)  $R_3N$ d)  $NH_4X$ a) R—NH<sub>2</sub> c)  $[R_4N]^+X^-$ 236. Which of the following statement about primary amines is false? a) Alkylamines are stronger base than arylamines b) Alkylamines react with nitrous acid to produce alcohols c) Arylamines react with nitrous acid to produce phenols d) Alkylamines are stronger bases than ammonia. 237. The end product in the below reaction is  $C_2H_5NH_2 \xrightarrow{HNO_2} A \xrightarrow{PCl_5} B \xrightarrow{NH_3} C$ a) Ethyl cynide b) Ethyl amine c) Methyl amine d) Acetamide 238. In hypobromite reaction of amide, carbonyl carbon atom is lost as: c)  $CO_3^{2-}$ d) None of these a) CO b)  $CO_2$ 239. Identify the product Z in the following reaction scheme  $\mathsf{C_6H_5NH_2} \xrightarrow{\mathsf{Ac_2O}} X \xrightarrow{\mathsf{Br_2/CCl_4}} Y \xrightarrow{\mathsf{HOH}} Z$ a) p-bromoaniline b) p-bromoacetophenone d) o-bromoacetophenone c) p-bromoacetanilide

240. Which of the following is not correct?

- a) Ethylamine and aniline both have NH<sub>2</sub> group
- b) Ethylamine and aniline both dissolve HCl
- c) Ethylamine and aniline both react with CHCl<sub>3</sub> and KOH to form unpleasant smell
- d) Ethylamine and aniline both react with  $NaNO_2 + HCl$  to give hydroxyl compounds in cold
- 241. The reduction of CH<sub>3</sub>CN to CH<sub>3</sub>CH<sub>2</sub>NH<sub>2</sub> is called:

- a) Rosenmund's reduction
- b) Clemmensen's reduction
- c) Mendius reduction
- d) Hofmann's reduction
- 242. Phenyl cyanide cannot be obtained by

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

b) 
$$C_6H_5 - CH = NOH \frac{Ac_2O_5}{C}$$

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

b) 
$$C_6H_5 - CH = NOH \xrightarrow{Ac_2O,\Delta}$$
  
d)  $C_6H_5NH_2 \xrightarrow{1.NaNO_2/HCI}$ 

- 243. Which nitro compound will show tautomerism?
  - a)  $C_6H_5NO_2$
- b)  $(CH_3)_3CNO_2$
- c) CH<sub>3</sub>CH<sub>2</sub>NO<sub>2</sub>
- d) o-nitrotoluene
- 244. Alkyl halide (RX) on treatment with KCN followed by reduction leads to formation of:
  - a)  $RNH_2$
- b) RCH<sub>2</sub>NH<sub>2</sub>
- c)  $RH + NH_3$
- d)  $RCH_3 + N_2$
- 245. Aniline gives a precipitate with bromine. The colour of precipitate is
  - a) Red

- b) Black
- c) Blue

- d) White
- 246. The reagent that reacts with nitromethane to form methyl hydroxylamine is
  - a) Zn/HCI
- b) Zn/NH<sub>4</sub>CI
- c) Zn/NaOH
- d) Sn/HCI
- 247. Identify the product in the following sequence 3, 4, 5-tribromoaniline

$$\xrightarrow{\text{(i)Diazotization}} ?$$

a) 3, 4, 5-tribromobenzene

b) 1, 2, 3-tribromobenzene

c) 2, 4, 6-tribromobenzene

- d) 3, 4, 5-tribromonitrobenzene
- 248. Production of amines by ammonia and alkyl halides is called
  - a) Frankland reaction

- b) Hofmann's ammonolysis
- c) Hofmann's mustrard oil reaction
- d) Hofmann's bromamide reaction
- m-fluoronitrobenzene is best synthesized by the specific problem of the synthesized by the specific problem is best synthesized by the specific problem i 249. *m*-fluoronitrobenzene is best synthesized by using the reaction

- 250. Allyl isocyanide contains ......and.....bonds.
  - a)  $9\sigma$ ,  $3\pi$
- c) 3σ, 4π
- d)  $5\sigma$ ,  $7\pi$

251.

a) 
$$9\sigma$$
,  $3\pi$  b)  $9\sigma$ ,  $9\pi$ 

$$\begin{array}{c} CH_{3} \\ | \\ CH_{3}CH_{2}CH_{2} - {}^{+}N - CH_{2}CH_{3} \\ | \\ CH_{3} \end{array} \right] OH^{-}$$

Is heated, then

a) Propene is the major product

- b) Ethane and  $C_3H_7N(CH_3)_2$  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
- 252. In the following reaction,  $X \xrightarrow{\text{Bromination}} Y \xrightarrow{\text{NaNO}_2} Z \xrightarrow{\text{Boiling}} \text{tribromo benzene. } X \text{ is}$ 
  - a) Benzoic acid
- b) Salicylic acid
- c) Phenol
- d) Aniline
- 253. Reaction of nitrous acid on 1° aliphatic amines in cold will give:
  - a) A diazonium salt
- b) An alcohol
- c) A nitrile
- d) A dye

- 254. Benzoyl chloride does not react with:
  - a) Primary or secondary amines
  - b) Aliphatic compounds
  - c) Aromatic compounds
  - d) Carboxylic acids
- 255. The compound, N-ethyl-N-methylpropanamine forms non-superimposable mirror image but does not show optical activity. This is due to

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 from into another
- 256. > C=O compounds reacts with NH $_3$  or amines followed by H $_2$ /Ni. The reaction is called

b) Hofmann bromamide

a) Mendius reaction c) Reductive amination

- d) Gabriel's phthalimide
- 257. Decreasing order of basic nature in aqueous solutions
  - a)  $C_6H_5NH_2 > NH_3 > CH_3NH_2 > (CH_3)_2NH$
- b)  $NH_3 > C_6H_5NH_2 > CH_3NH_2 > (CH_3)_2NH$
- c)  $(CH_3)_2NH > CH_3NH_2 > NH_3 > C_6H_5NH_2$
- d)  $CH_3NH_2 > (CH_3)_2NH > NH_3 > C_6H_5NH_2$
- 258. Comparing basic strength of NH<sub>3</sub>, CH<sub>3</sub>NH<sub>2</sub> and C<sub>6</sub>H<sub>5</sub>NH<sub>2</sub> 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 NH<sub>3</sub> is highest
- 259. Decreasing order of basicity of the three isomers of methoxyaniline is

a) 
$$p$$
-CH<sub>3</sub>OC<sub>6</sub>H<sub>4</sub>NH<sub>2</sub> >  $o$  - CH<sub>3</sub>OC<sub>6</sub>H<sub>4</sub>NH<sub>2</sub> >  $m$  - CH<sub>3</sub>OC<sub>6</sub>H<sub>4</sub>NH<sub>2</sub>

b) 
$$p$$
-CH<sub>3</sub>OC<sub>6</sub>H<sub>4</sub>NH<sub>2</sub> >  $m$  - CH<sub>3</sub>OC<sub>6</sub>H<sub>4</sub>NH<sub>2</sub> >  $o$  - CH<sub>3</sub>OC<sub>6</sub>H<sub>4</sub>NH<sub>2</sub>

c) 
$$o-\text{CH}_3\text{OC}_6\text{H}_4\text{NH}_2 > p-\text{CH}_3\text{OC}_6\text{H}_4\text{NH}_2 > p-\text{CH}_3\text{OC}_6\text{H}_4\text{NH}_2$$

d) 
$$o$$
-CH<sub>3</sub>OC<sub>6</sub>H<sub>4</sub>NH<sub>2</sub> >  $m$  - CH<sub>3</sub>OC<sub>6</sub>H<sub>4</sub>NH<sub>2</sub> >  $p$  - CH<sub>3</sub>OC<sub>6</sub>H<sub>4</sub>NH<sub>2</sub>

- 260. Which one of the following is the strongest base in aqueous solution?
- a) Trimethylamine
  - b) Aniline
- c) Dimethylamine
- d) Methylamine

- 261. Which compound is known as alkyl carbylamines?
  - a) R. CN
- b) R. NC

- c) Ar. CN
- d) Ar. NC

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

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

- c) Acid

d) Alcohol

263. 
$$C_6H_5NH_2 \xrightarrow{NaNO_2} X \xrightarrow{Cu_2(CN)_2} Y \xrightarrow{H_2O/H^+} Z$$

Z is identified as

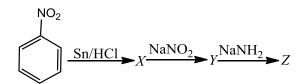
- a)  $C_6H_5 NH CH_3$
- b)  $C_6H_5 COOH$
- c)  $C_6H_5 CH_2 NH_2$  d)  $C_6H_5 CH_2COOH$

264. The product [*A*] formed in the reaction;

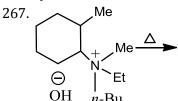
$$2C_5H_5$$
 –  $CN \xrightarrow{Na} [A]$  is:

- 265. 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

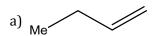
266. What is 'Z'in the following reaction?



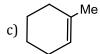
- a) Benzoic acid
- b) Cyanobenzoic acid
- c) Benzamide
- d) Aniline



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



b)  $CH_2 = CH_2$ 





268. Nitroalkane is acidic only towards:

- a) Na<sub>2</sub>CO<sub>3</sub>
- b) NaOH
- c) Alcohol
- d) Liquid NH<sub>3</sub>

269. 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)  $C_6H_5 - NH - CO - C_6H_5$ 

270. CH<sub>3</sub>NH<sub>2</sub> + CHCl<sub>3</sub> + KOH →nitrogen containing compound +KCI + H<sub>2</sub>O. Nitrogen containing compound is

- a)  $CH_3 C \equiv N$
- b)  $CH_3 NH CH_3$  c)  $CH_3 N \stackrel{-}{=} \stackrel{+}{C}$  d)  $CH_3 \stackrel{+}{N} \stackrel{-}{=} C$

271. 
$$Y \xrightarrow{\text{Reduction}} [R_2 C = NH] \xrightarrow{\text{H}_3 O^+} X$$

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

a) Aldehyde, ketone, NH<sub>3</sub>

b) Ketone, 1° amine, KMnO<sub>4</sub>

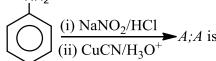
c) Ketone, 2° amine, KMnO<sub>4</sub>

d) Ketimine, 1° amine, H<sub>2</sub>SO<sub>5</sub>

272. Aniline is weaker base than ethylamine because:

- a) Lone pair of electrons of N-atom is not freely available for coordination with a proton due to resonance than in ethylamine
- b) Its b. p. is higher than that of ethylamine
- c) It does not produce sufficient concentration of OH<sup>-</sup> ions in solution
- d) It is insoluble in water while ethylamine is soluble in water

 $NH_2$ 273.







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

- a)  $(CH_3)_4N^+I^-$
- b) CH<sub>3</sub>OCH<sub>3</sub>
- c)  $(CH_3)_3S^+I^-$
- d)  $(CH_3)_3CI$

275. N<sub>2</sub> gas is liberated when [HCl + NaNO<sub>2</sub>] reacts with the following compounds

- $(A)CH_3CH_2NH_2$ 
  - (B) Urea
  - (C)CH<sub>3</sub>CONH<sub>2</sub>
- $(D)C_6H_5NH_2$

The answer is

- a) A, B, C
- b) B, C, D
- c) A, C, D
- d) A, B, D

276. 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

- 277. (A) $C_2H_5NH_2 \xrightarrow{(ii)AgNO_2} [W]$ 
  - $(B)(CH<sub>3</sub>)<sub>2</sub>CHNH<sub>2</sub> \xrightarrow{(i)NOCl}_{(ii)AgNO<sub>2</sub>} [X]$
  - $(C)(CH_3)_3 CNH_2 \xrightarrow[(ii)AgNO_2]{(ii)AgNO_2} [Y]$
  - (D)CH<sub>3</sub>CH(NH<sub>2</sub>)C<sub>2</sub>H<sub>5</sub> $\xrightarrow{\text{(i)NOCl}}_{\text{(ii)AgNO}_2}$ [Z]

Which product will not show tautomerism?

c) Y

d)Z

a) W b) X

- 278. Name of method use to separate primary, secondary and tertiary amines is

- a) Hofmann method b) Lucas method c) Victor Meyer method 279. Alkyl cyanides undergo Stephen reduction to produce
  - a) Aldehyde
- b) Secondary amine
- c) Primary amine
- d) amide
- 280. A compound of molecular formula C<sub>3</sub>H<sub>9</sub>N when reacts with benzene sulphonyl chloride gives a product soluble in dilute NaOH solution. The compound should be
  - a)  $(CH_3)_3N$
- b)  $(CH_3)_2CH NH_2$
- c)  $\sim$  NH
- d) All of these

d) Kolbe method

- 281. Benzenediazonium chloride on reaction with phenol in weakly basic medium gives
  - a) Diphenyl ether

b) p-hydroxyazobenzene

c) Chlorobenzene

d) benzene

282. The conversion

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

Can be brought about by reduction with

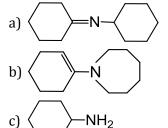
- a) Na<sub>3</sub>AsO<sub>3</sub>/NaOH
- b) Glucose/HCl
- c) Zn/NaOH
- d) LiAlH<sub>4</sub>/ether

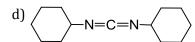
283. The correct order of basicities of the following compounds is:

- a) 2 > 1 > 3 > 4
- b) 1 > 3 > 2 > 4
- c) 3 > 1 > 2 > 4 d) 1 > 2 > 3 > 4
- 284. Aniline on oxidation with Na<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> and H<sub>2</sub>SO<sub>4</sub> gives
  - a) Benzoic acid
- b) *m*-amino benzoic acid c) Schiff's base
- d) p-bezoquinone

- 285. Which among the following has the highest boiling point?
  - a) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>
- $CH_3CH_2$ -NH
- CH<sub>3</sub> c) CH<sub>3</sub>—N—CH<sub>2</sub>
- d) CH<sub>3</sub>NH<sub>2</sub>

286. Which of the following is an enamine?





- 287. When aniline is heated with glacial acetic acid in presence of anhydrous ZnCl<sub>2</sub>, the product is:
  - a) Acetamide
- b) Acetanilide
- c) Phenyl acetamide
- d) Chlorobenzene

- 288. Acetonitriles on hydrolysis produce which of the following?
  - a) Amine
- b) Acid

- c) Amides
- d) Carbonyl compounds
- 289. The number of  $\pi$ -bonds in the formula given below, NC—CH=CH—CN are:
  - a) 5

b) 4

c) 3

d) 2

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

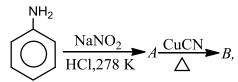
$$e \xrightarrow{\text{NaIO}_3} (X) \xrightarrow{\text{CH}_3 \text{CO}_3 \text{H}} (Y)$$

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
- c) Aniline
- d) Azoxybenzene
- 291. Examine the following two structures for the anilinium ion and choose the correct statement from the ones given below

$$^{\dagger}NH_3$$
  $^{\dagger}NH_3$   $^{\dagger}$   $^{\dagger}$   $^{\dagger}$   $^{\dagger}$   $^{\dagger}$   $^{\dagger}$   $^{\dagger}$   $^{\dagger}$   $^{\dagger}$ 

- 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
- 292. 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

293. In the chemical reaction,

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

The compounds (A) and (B) are respectively

- a) CH<sub>3</sub>CH<sub>2</sub>CONH<sub>2</sub> and 3KCI
- c) C<sub>2</sub>H<sub>5</sub>NC and 3KCI

- b) C<sub>2</sub>H<sub>5</sub>NC and K<sub>2</sub>CO<sub>3</sub>
- d) C<sub>2</sub>H<sub>5</sub>CN and 3KCI
- 294. The product obtained in the reduction

$$O_2$$
 $O_2$ 
 $O_3$ 
 $O_2$ 
 $O_3$ 
 $O_4$ 
 $O_2$ 
 $O_4$ 
 $O_4$ 
 $O_5$ 
 $O_4$ 
 $O_5$ 
 $O_5$ 
 $O_6$ 
 $O_7$ 
 $O_8$ 
 $O_8$ 

- d) The compound is not reduced
- 295. In the reaction between  $\text{CH}_3\text{NC}$  and HgO, the product obtained is
  - a) Methyl isothiocyanate

b) Methyl isocyanate

c) Methyl amine

d) Methyl cyanide

296. The compound which on reaction with cold  $HNO_2$  gives only nitrosoamine is:

a) CH<sub>3</sub>NH<sub>2</sub>

- b)  $(CH_3)_2NH$
- c)  $(CH_3)_3N$

d)  $(C_2H_5)_3N$ 

- 297. Which of the following is involved in Sandmeyer's reaction?
  - a) Ferrous salt
  - b) Diazonium salt
  - c) Ammonium salt
  - d) Cupraammonium salt
- 298. During diazotization of benzenamine with sodium nitrite and hydrochloric acid, the excess of hydrochloric acid is used primarily to
  - a) Check the hydrolysis of  $\phi$  OH

- b) Ensure a stoichiometric amount of nitrous acid
- c) Check the concentration of free aniline
- d) Neutralize any base formed during reaction
- 299. The correct structure of ethylenediamine-tetra acetic acid (EDTA) is:

- 300. 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
- 301. Identify *X* in the sequence,

$$X \xrightarrow{\text{HNO}_2} \text{C}_3\text{H}_8\text{O} \xrightarrow{\text{H}_2\text{SO}_4} \text{C}_3\text{H}_6\text{O}_2$$
:

- c)  $(CH_3)_3N$
- d) None of the above
- 302. Reduction of alkyl nitriles, produces

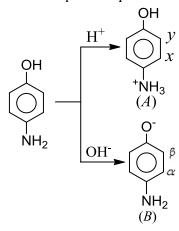
- a) Secondary amine
- b) Primary amine
- c) Tertiary amine
- d) amide

- 303. General formula of an amine is:
  - a)  $C_nH_{2n+1}N$
- b)  $C_n H_{2n+2} N$
- c)  $C_nH_{2n+3}N$
- d)  $C_n H_{2n} N$

- 304. Biuret test is not given by:
  - a) Proteins
- b) Carbohydrates
- c) Polypeptides
- d) Urea

- 305. Which of the following is not a nitro derivative?
  - a) C<sub>6</sub>H<sub>5</sub>NO<sub>2</sub>
- b) CH<sub>3</sub>CH<sub>2</sub>ONO
- $_{\text{c})} \stackrel{\text{CH}_3-\text{CH}-\text{N}}{\stackrel{\text{C}}{\text{CH}_3}} \circ$
- d)  $C_6H_4(OH)NO_2$

306. Consider *p*-aminophenol



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

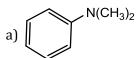
- a) x in A and  $\beta$  in B
- b) x in A and  $\alpha$  in B
- c) y in A and  $\alpha$  in B
- d) y in A and  $\beta$  in B

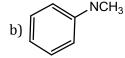
307. Diethyl carbonate on heating with ammonia gives:

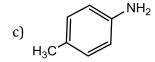
- a)  $C_2H_5NH_2$
- b)  $(C_2H_5)_3N$
- c)  $(C_2H_5)_2NH$
- d) Urea

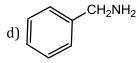
308. A primary amine hated with CS<sub>2</sub> in presence of excess of HgCl<sub>2</sub> gives isothiocyanate. The reaction is called:

- a) Hofmann's bromamide reaction
- b) Hofmann's mustard oil reaction
- c) Perkin's condensation
- d) Hofmann's elimination
- 309. 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  $\beta$  —naphthol is



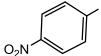


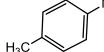


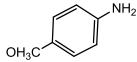


310. The correct order of increasing basic nature of the following bases is



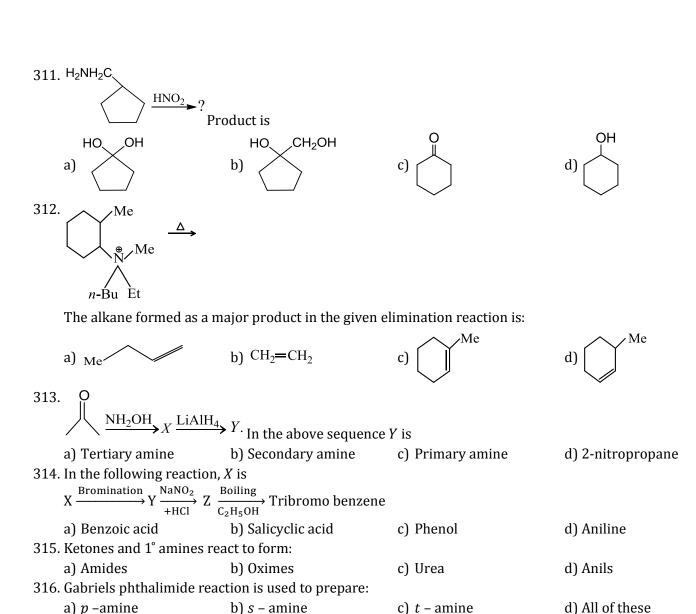








- a) II < V < I < III < IV
- b) V < II < I < III < IV
- c) II < V < I < IV < III
- d) V < II < I < IV < III



c)  $RNH_2 + HNO_2$ 

c)  $(C_2H_5)_2NH$ 

c) ROR

b) Diazonium salt

317. Carcinogens are the products of the reaction between:

318. Mendius method of preparation of amines consists of:

322. Primary amine (RNH<sub>2</sub>) reacts with nitrous acid to give

a) Catalytic reduction of alkyl cyanides b) Reduction of amide with LiAlH<sub>4</sub>

c) Reduction of nitroparaffin with Sn + HCl d) Reduction of oximes with Na +  $C_2H_5OH$ 

a)  $R_2$ NH + HNO<sub>2</sub>

with Hinsberg reagent?

c) Hydrazo compound

a) N-ethyl acetamide

321. Ethyl amine on acetylation gives

a) CH<sub>3</sub>NH<sub>2</sub>

a) Phenol and N<sub>2</sub>

a)  $RNH_3^+NO_2$ 

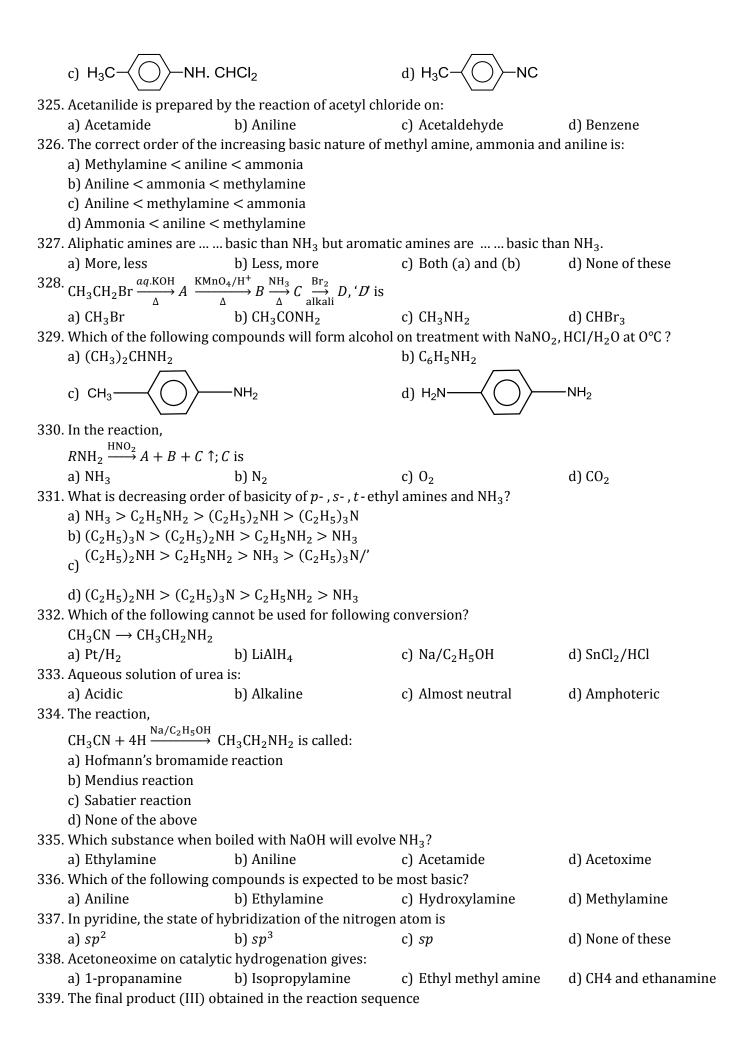
b)  $R_3$ N + HNO<sub>2</sub>

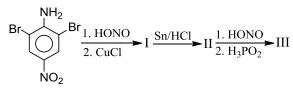
b)  $(CH_3)_3N$ 

b) Acetamide

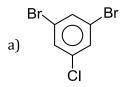
b) ROH

b) NH<sub>3</sub> 324. The reaction of chloroform with alcoholic KOH and p-toluidine from d) None of these

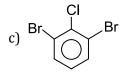




is



b) Br B



d) None of these

340. The main product in the reaction,

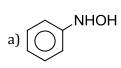
HCONHR 
$$\xrightarrow{\text{POCl}_3}$$
 is:

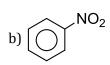
a) RCN

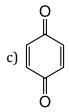
b) RNC

- c) RCNO
- d) RNCO

- 341. Ethyl isocyanide on hydrolysis in acidic medium generates
  - a) Ethylamine salt and methanoic acid
- b) Propanoic acid and ammonium salt
- c) Ethanoic acid and ammonium salt
- d) Methylamine salt and ethanoic acid
- 342. The oxidation of aniline with per acetic acid in the presence of acetic acid by refluxing gives







d) None of these

- 343. The compound having the molecular formula C<sub>3</sub>H<sub>9</sub>N represent :
  - a) Trimethylamine
- b) *n*-propylamine
- c) Isopropylamine
- d) All of these

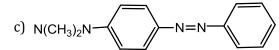
- 344. Amines have:
  - a) Garlic odour
- b) Fishy odour
- c) jasmine odour
- d) Bitter almonds odour
- 345. Which of the following compounds gives a secondary amine on reduction?
  - a) Nitromethane
- b) Nitrobenzene
- c) Methyl isocyanide
- d) Methyl cyanide

- 346. Indicate the correct statement.
  - a)  $C_2H_5N^+H_3OH^-$  is acidic
  - b) C<sub>2</sub>H<sub>5</sub>NH<sub>2</sub> is less basic than NH<sub>3</sub>
  - c) C<sub>2</sub>H<sub>5</sub>NH<sub>2</sub> is a stronger base than NH<sub>3</sub>
  - d) C<sub>2</sub>H<sub>5</sub>NH<sub>2</sub> forms salts with bases
- 347. Choose the incorrect statement.
  - a) Primary amines show intermolecular hydrogen bonds.
  - b) Tert-butylamine is primary amine.
  - c) Tertiary amines do not show intermolecular hydrogen bonds.
  - d) Isopropylamine is a secondary amine.
- 348. 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
- 349. The strongest base among the following is
  - a)  $C_6H_5NH_2$
- b)  $(C_6H_5)_2NH$
- c) NH<sub>3</sub>

- d)  $(C_2H_5)_2NH$
- 350. Aniline when diazotized in cold and then treated with dimethyl aniline gives a coloured product. Its structure would be

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

b) 
$$CH_3$$
  $N=N$   $NH_2$ 



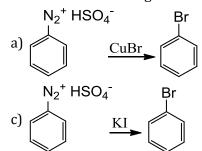
d) 
$$(CH_3)_2N$$
  $N=N$   $NH_2$ 

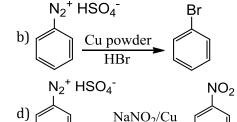
- 351. Treatment of nitrobenzene with acetyl chloride in the presence of anhydrous AlCl<sub>3</sub> gives
  - a) 2-nitroacetophenone

b) 3-nitroacetophenone

c) 4-nitroacetophenone

- d) None of these
- 352. Urea when heated slowly, product formed is:
  - a)  $N_2$
  - b) CO<sub>2</sub>
  - c) biuret
  - d) Ammonium carbamate
- 353. Which of the following reactions is an example of Sandmeyer reaction?





354. Given the following sequence of reactions,

$$\operatorname{CH_3CH_2I} \xrightarrow{\operatorname{NaCN}} A \xrightarrow[\operatorname{Partial}]{\operatorname{OH}^-} B \xrightarrow[\operatorname{hydrolysis}]{\operatorname{Br}_2/\operatorname{NaOH}} C$$

The major product 'C' is

a) CH<sub>3</sub>CH<sub>2</sub>NH<sub>2</sub>

 $CH_3$ .  $CH_2C - NHBr$ b) Ш 0

c) CH<sub>3</sub>. CH<sub>2</sub> - COONH<sub>4</sub>

CH<sub>3</sub>. CH<sub>2</sub>C - NBr<sub>2</sub> d) Ш 0

- 355. CHCl<sub>3</sub> and KOH on heating with a compound from a bad smelling product, compound is
  - a) C<sub>2</sub>H<sub>5</sub>CN
- b) C<sub>2</sub>H<sub>5</sub>NC
- c)  $C_2H_5OH$
- d)  $C_2H_5NH_2$

- 356. Benzamide can be converted into benzonitrile with
  - a) H<sub>2</sub>0+

- b)  $0H^{-}/H_{2}O$
- c) KCN

d)  $P_2O_5$ 

- 357. 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)  $1^{\circ} < 2^{\circ} > 3^{\circ}$

- 358. Hofmann's bromamide reaction is to convert
  - a) Acid to alcohol
- b) Alcohol to acid
- c) Amide to amine
- d) Amine to amide
- 359. During coupling reaction of benzene diazonium chloride and aniline, the pH of reaction medium should be approximately
  - a) 1-2

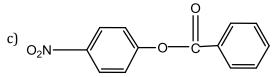
- b) 9-10
- c) 4-5

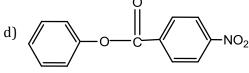
d) 7 - 8

360.

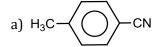
The product *A* is

a) 
$$O - C - C$$





361. The reaction of chloroform with alcoholic KOH and p-toluidine form



b) 
$$H_3C$$
  $\longrightarrow$   $N_2CI$ 

c) 
$$H_3C$$
 NHCHC $I_2$ 

d) 
$$H_3C$$
  $\longrightarrow$   $NC$ 

362.

Pyrolysis of Me´ + `CD<sub>3</sub> would give

- a) Mixture of  $CH_2 = CH CD_3$  and  $CH_3 CH = CD_2$  b)  $CH_3 CH = CD_2$
- c)  $Me_2N^+ = C(CD_3)(CH_3)$

d)  $CH_2 = CH - CD_3$ 

363. Which of the following compounds is soluble in benzene but almost insoluble in water?

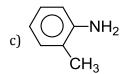
- a) C<sub>2</sub>H<sub>5</sub>OH
- b) CH<sub>3</sub>CO<sub>2</sub>H
- c) CH<sub>3</sub>CHO
- d) C<sub>6</sub>H<sub>5</sub>NO<sub>2</sub>

364. The action of nitrous acid on a primary amine gives:

- a) Nitroalkane
- b) Alkyl nitrite
- c) Alcohol
- d) Secondary amine

365. Which of the following is the strongest base?

a) 
$$\langle \bigcirc \rangle$$
  $\sim$   $NH_2$ 



d) 
$$\langle CH_2 - NH_2 \rangle$$

366. Which is not the property of ethanenitrile (CH<sub>3</sub>CN)?

- 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

367. The basic character of methylamines in vapour phase is:

- a)  $3^{\circ}>2^{\circ}>1^{\circ}>NH_{3}$
- b)  $2^{\circ}>3^{\circ}>1^{\circ}>NH_{3}$
- c)  $1^{\circ}>2^{\circ}>3^{\circ}>NH_{3}$
- d) None of these

368. 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)  $C_4H_9OH + KCN$

369. Isopropyl amine with excess of acetyl chloride will give

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

$$(CH_3)_2CH - N - COCH_3$$

DJ

$$CH_3CH_2CH_2 - N - COCH_3$$

c)  $(CH_3)_2CHN(COCH_3)_2$ 

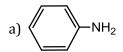
370. In the reaction of (S) 2-phenylpropamide with NaBr/H<sub>2</sub>O to give 1-phenylethylamine

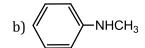
- a) There is retention of configuration
- b) There is inversion of configuration
- c) A mixture of two products is obtained
- d) There is no reaction

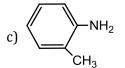
371. High basicity of Me<sub>2</sub>NH relative to Me<sub>3</sub>N is attributed to

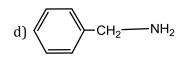
- a) Effect f solvent
- b) Inductive effect of Me
- c) Shape of Me<sub>2</sub>NH
- d) Shape of Me<sub>3</sub>N

372. Which of the following is the strongest base?

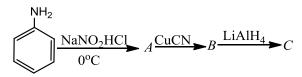








373. In the reaction sequence



The product 'C' is

- a) Benzonitrile
- b) Benzaldehyde
- c) Benzoic acid
- d) Benzyl amine
- 374. Which one does not liberate NH<sub>3</sub> when undergoes hydrolysis?
  - a) Acetanilide
- b) Acetonitrile
- c) Acetamide
- d) Phenyl isocyanide

375. Identify the major product of the reaction

- 376. Primary nitroalkanes on hydrolysis give:
  - a)  $RCOOH + NH_2OH$
- b) RCOOH
- c) NH<sub>2</sub>OH
- d) RCOR

377. In the reaction

$$CH_3CN + 2H \xrightarrow{HCI}_{SnCI_2} X \xrightarrow{Boiling H_2O} Y$$
,

The term Y is,

- a) Acetone
- b) Ethanamine
- c) Acetaldehyde
- d) Dimethyl amine
- 378. Which one of the following compounds forms a quaternary salt on reacting with excess methyl iodide?
  - a) C<sub>2</sub>H<sub>5</sub>OCH<sub>3</sub>
- b) (CH<sub>3</sub>)<sub>2</sub>CHOC<sub>2</sub>H<sub>5</sub>
- c)  $C_6H_5NH_2$
- d)  $C_6H_5NO_2$

						: ANSW	V	ER K	EY					
1)	С	2)	b	3)	b			189)	a	190)	b	191)	С	192) a
5)	b	6)	c	7)	b	-	d	-	c	194)	d	195)	c	196) d
9)	d	10)	b	11)	d	40)	a	197)	a	198)	a	199)	a	200) b
13)	b	14)	a	15)	c		b		d	202)	c	203)	c	204) b
17)	b	18)	b	19)	b		d		d	206)	a	207)	c	208) a
21)	c	22)	c	23)	b	2.0	С	209)	d	210)	a	211)	c	212) a
25)	c	26)	c	27)	c		d	213)	a	214)	d	215)	d	216) a
29)	d	30)	a	31)	b	32)	d	217)	c	218)	b	219)	a	220) d
33)	b	34)	c	35)	d	36)	c	221)	a	222)	d	223)	a	224) d
37)	b	38)	a	39)	c	40)	a	225)	d	226)	c	227)	c	228) b
41)	d	42)	b	43)	c	44)	d	229)	b	230)	c	231)	c	232) a
45)	b	46)	a	47)	b	<b>48)</b>	b	233)	a	234)	a	235)	c	236) c
49)	d	50)	a	51)	b	<b>52)</b>	b	237)	b	238)	C	239)	a	<b>240)</b> d
53)	d	54)	d	55)	c	<b>56</b> ) a	a	241)	C	242)	c	243)	c	244) b
57)	b	58)	b	59)	b	<b>60)</b>	b	245)	d	246)	b	247)	b	248) b
61)	b	62)	a	63)	b	<b>64)</b>	a	249)	a	250)	a	251)	C	252) d
65)	b	66)	d	67)	C	<b>68</b> ) :	a	253)	b	254)	d	255)	b	256) c
69)	a	70)	d	71)	a	72)	c	257)	C	258)	C	259)	C	260) c
73)	c	74)	a	75)	a	76)	d	261)	b	262)	b	263)	b	264) b
77)	b	78)	a	79)	c	80)	a	265)	a	266)	d	267)	b	268) b
81)	C	82)	C	83)	c	,	C	,	b	270)	b	271)	b	272) a
85)	b	86)	d	87)	b	,	d	,	b	274)	a	275)	a	276) b
89)	C	90)	b	91)	a	•		277)	C	278)	a	279)	a	280) b
93)	C	94)	C	95)	a	•		_	a	282)	a	283)	b	284) d
97)	b	98)	d	99)	b	,		,	a	286)	a	287)	b	288) b
101)	b	102)	С	103)	c	,	d	,	a	290)	b	291)	C	292) b
105)	a	106)	a	107)	b	,	a	,	C	294)	b	295)	b	296) b
109)	С	110)	b	111)	С	,	a	_	b	298)	c	299)	C	300) c
113)	c	114)	d	115)	C	=		301)	b	302)	b	303)	C	304) b
117)	C	118)	b	119)	d	-		305)	b	306)	b	307)	d	308) b
121)	b	122)	c	123)	d	-		309) 313)	c	310)	a d	311)	d	312) d
125) 129)	a d	126) 130)	c b	127) 131)	a d	-		317)	c	314) 318)	d	315) 319)	d	316) a 320) b
133)	a	134)	C	131)	u C	-		317) 321)	a a	322)	a b	323)	a b	320) b
137)	a C	134)	a	139)	d	,		321) 325)	a b	326)	b	323) 327)	a	324) u
141)	a	142)	C	143)	c	-		329)	a	330)	b	331)	d	332) d
145)	b	146)	b	147)	d	-		333)	c	334)	b	335)	c	336) b
149)	d	150)	b	151)	b	-		337)	a	338)	b	339)	b	340) b
153)	c	154)	a	151) 155)	c	-		341)	a	342)	b	343)	d	344) b
157)	b	154)	c	159)	b	_		345)	c	346)	С	347)	d	348) a
161)	b	162)	b	163)	a	-		349)	d	350)	c	351)	c	352) c
165)	c	166)	c	167)	d	-		353)	a	354)	a	355)	b	356) d
169)	b	170)	d	171)	c	-		357)	a	358)	c	359)	С	360) c
173)	a	174)	a	175)	c	-		361)	d	362)	a	363)	d	364) c
177)	a	178)	d	179)	b	-		365)	d	366)	d	367)	a	368) a
181)	d	182)	d	183)	c	-		369)	С	370)	a	371)	a	372) d
_	c	•	c	187)		-		373)	d	374)	d	-	b	376) a
,	_			-3.7		===,								

# : HINTS AND SOLUTIONS :

# 1 **(c)**

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

acetonitrile ethanamine

# 3 **(b**

Carbylamine (or isocyanides) give secondary amine on reduction.

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

## 4 (a)

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

# 5 **(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. ( $C_2H_5$ )<sub>3</sub>N is a tertiary amine, so does not react with  $C_6H_5SO_2CI$ .

#### 7 **(b**)

$$\phi N_2^+ + \phi OH \rightarrow \phi N_2 \phi OH$$
or  $\langle O \rangle - N = N - \langle O \rangle - OH$ 

coupling product

## 10 **(b)**

—CN and —NC are different functional groups.

#### 11 (d)

$$\begin{array}{c} \text{CH}_3\text{CN} \xrightarrow[\text{-PM}]{\text{H.OH}} \xrightarrow{\text{CH}_3\text{COOH}} \xrightarrow[\text{-CO}_2]{\text{Decarboxylation}} \text{CH}_4 \\ \xrightarrow[\text{-PM}_3]{\text{CH}_3\text{COOH}} \xrightarrow[\text{-CO}_2]{\text{Decarboxylation}} \xrightarrow{\text{CH}_4} \\ \xrightarrow[\text{-PM}_3]{\text{CH}_3\text{COOH}} \xrightarrow[\text{-CO}_2]{\text{Decarboxylation}} \\ \xrightarrow[\text{-PM}_3]{\text{CH}_3\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{Decarboxylation}} \\ \xrightarrow[\text{-PM}_3]{\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{COOH}} \\ \xrightarrow[\text{-PM}_3]{\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{COOH}} \\ \xrightarrow[\text{-PM}_3]{\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{COOH}} \\ \xrightarrow[\text{-PM}_3]{\text{COOH}} \xrightarrow[\text{-PM}_3]{\text{$$

Ethane

ethanoic acid

methane

# Nitrile

(A)

12 **(a)** 

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

# 13 **(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.

# 14 **(a)**

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

# 15 **(c)**

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

# 16 **(b)**

The order for acidic nature depends upon the ease to lose H<sup>+</sup> ion. —COOH is resonance stabilized and thus lose H<sup>+</sup> at the earliest. Also NH<sub>3</sub><sup>+</sup> near to —COOH releases H<sup>+</sup> more easily due to electron withdrawing nature of —COOH than NH<sub>3</sub><sup>+</sup> far away from —COOH.

## 17 **(b)**

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

# 18 **(b)**

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

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

Nitrobenezene

$$\begin{array}{c} \text{CH}_3 - \text{C} \equiv \text{N} \xrightarrow{\substack{\text{Reduction} \\ \text{HONO} \\ \end{array}}} \text{CH}_3 \text{CH}_2 \text{NH}_2 \\ \xrightarrow{\substack{\text{HONO} \\ \text{methyl cyanide} \\ \end{array}}} \text{CH}_3 \text{CH}_2 \text{OH}$$

methyl cyanide ethanamin 1. (B)

 $\begin{array}{c} \operatorname{CH_3CH_2NH_2} \xrightarrow{\operatorname{CHCI_3,KOH}} \operatorname{CH_3CH_2N} \stackrel{\cong}{=} \operatorname{C} \\ 2. & \text{Ethyl isocyanide} \end{array}$ 

(C)

# 21 **(c)**

$$CH_3 - NH_2 + 3CH_3I \rightarrow (CH_3)_4N^+I^-$$
  
methyl amine quaternary  
ammonium salt

therefore, 3 moles of  $\mathrm{CH_3I}$  (methyl iodide) are required for reaction with methyl amine.

# 23 **(b)**

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

$$\begin{array}{c|c}
 & \text{NHCOCH}_3 \\
\hline
 & \text{NHCOCH}_3
\end{array}$$

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

$$\xrightarrow{\text{HOH}} \begin{array}{c} NH_2 \\ NO_2 \\ NO_2 \end{array} + \begin{array}{c} NH_2 \\ NO_2 \end{array}$$

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

# 24 **(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

3.

(B)

(C)

#### 27 **(c)**

Reaction of aniline with benzaldehyde is condensation reaction.

H
$$C_6H_5 - C = O + H_2NC_6H_5 \xrightarrow{Condensation} C_6H_5CH$$

$$= NC_6H_5 + H_2O$$

Benzaldehyde aniline benzylidene aniline

#### 28 **(d)**

$$C_6H_5CH_2NH_2 + CHCI_3 + 3KOH$$
  
 $\rightarrow C_6H_5CH_2NC + 3KCI + 2H_2O$ 

Secondary amides such as N-methylethanamide on reduction with LiAIH<sub>4</sub> give secondary amines.

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

# 30 **(a)**

ethanol

Hofmann bromamide reaction is used to prepare  $1^{\circ}$  amine form primary amides. In this method, amides are treated with bromine in presence of KOH

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

# 32 **(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 +  $NaOH$   $-H_2O$  OCOR

33 **(b)** 

Due to +ve IE

in alkylamines and resonance in  $C_6H_5NH_2$ .

# 34 **(c**)

Schmidt reaction

$$0$$

$$||$$

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

## 7 **(b)**

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

Weakly basic 
$$N = N - OH + HCI$$
  $p$ -hydroxy azobenzene

40 **(a)** 

$$RCONH_2 \xrightarrow{P_2O_5} RCN + H_2O$$

# 41 (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

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

42 **(b)** 

Methyl cyanide gives acetic acid on hydrolysis.

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

43 **(c**)

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

44 (d)

C<sub>3</sub>H<sub>9</sub>N represent following structures CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>

Propanamine

(1°amine)

H I

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

N-methyl ethanamine

(2°amine)

CH<sub>3</sub>

 $CH_3 - N - CH_3$ 

N,N-dimethyl methanamine

(3°amine)

45 **(b**)

H-atom of  $C_6H_6$  ring is replaced by  $S_E$  reactions

46 (a)

Presence of—OCH<sub>3</sub> gp. on aniline makes it more basic than the presence of — NO<sub>2</sub>, —Cl or — CH<sub>3</sub> gp.

48 **(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  $C_2H_5NH_2 > CH_3NH_2 > NH_3 > C_6H_5NH_2$ 

49 **(d)** 

CH<sub>3</sub>NH<sub>3</sub>Cl is acidic in nature.

50 **(a**)

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

51 **(b)** 

Dimethyl oxalate is used for distinguishing primary, secondary and tertiary amines

(Hofmann's method).

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

$$COOC_2H_5$$
  $H-NHR$   $CONHR$   
 $| + \rightarrow | +2C_2H_5OH$   
 $COOC_2H_5$   $H-NHR$   $CONHR$ 

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

COOC<sub>2</sub>H<sub>5</sub> CONR<sub>2</sub> 
$$| +NHR2 \rightarrow | +C2H5OH$$
 COOC<sub>2</sub>H<sub>5</sub> COOC<sub>2</sub>H<sub>5</sub>

6. The tertiary amine under these conditions does not react at all since it does not contain a replaceable hydrogen atom.

52 **(b)** 

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

53 **(d)** 

It is carbylamine reaction,

benzyl amine 
$$CH_2NH_2$$
  $CHCl_3$   $CH_2NC$   $CH_2$ 

54 **(d)** 

Secondary amines give oily nitrosoamine with nitrous acid

$$(CH_3CH_2)_2NH + HNO_2$$
  
 $\rightarrow (CH_3CH_2)_2N - NO + H_2O$ 

Secondary amine nitrous acid oily nitrosoamine

55 (c

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

$$R_2$$
N - N = O + H<sub>2</sub>O  $\xrightarrow{\text{Conc.H}_2\text{SO}_4} R_2$ NH + HNO<sub>2</sub> nitrosoamine secondary amine

56 **(a)** 

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

$$R - \text{CN} + 2\text{H}_2\text{O} \xrightarrow{\text{NaOH}} R - \text{COOH} + \text{NH}_3$$
57 **(b)**

$$(A)R\text{CN} \xrightarrow{\text{Reduction}} R\text{CH}_2\text{NH}_2$$

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

$$R\text{CN} + \text{CH}_3\text{MgBr} \xrightarrow{} R - \text{C} = \text{N-MgBr} \xrightarrow{\text{H. OH}} \text{CH}_3$$

$$C\text{H}_3 \xrightarrow{} \text{OH}$$

(B) 
$$R$$
  $C=O + NH$   $CH_3$  methyl ketone

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

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

59 **(b)**

$$CHCI_3 + C_6H_5NH_2 + 3NaOH$$

$$\rightarrow C_6H_5NC + 3NaCI + 3H_2O$$
phenylisocyanide

62 **(a)** It is the first step of Gabriel's phthalimide synthesis. The hydrogen bonded to nitrogen is sufficiently acidic due to two  $\alpha$ -carbonyls.

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.

$$S_{N^2}$$
 $N^- + CH_2$ 
 $N^- + CH_2$ 

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.

$$H_2CCI$$
  $H_2CCI$   $H_2CCI$ 

63 **(b)**Only these aromatic primary amines undergo diazotisation in which —NH<sub>2</sub> is attached to

nucleus.

Molecular formula of benzonitrile is  ${\rm C_6H_5CN}$ .  ${\rm CN}$ 

69

(a)

phenyl cyanide or benzonitrile

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

- 67 (c)  $NH_2CONH_2 + NH_2 \cdot NH_2 \rightarrow NH_2CONHNH_2 + NH_3$ 68 (a)
  - $R-C \equiv N \xrightarrow{\substack{\text{hydrolysis} \\ \text{hydrolysis}}} R \text{CONH}_2 \xrightarrow{\substack{\text{hydrolysis} \\ \text{alkyl cyanide}}} R\text{COOH}$  Alkyl cyanide alkyl amide carboxylic acid
  - Secondary nitroalkanes can be converted into ketones by using aqueous HCI. R  $CHNO_2+HCI$  R  $C=O+N_2O+H_2O$

70 **(d)** 

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

72 **(c)** 

 ${\rm C_2H_5NH_2} \mathop \to \limits^{\rm [O]} {\rm CH_3CH} \mathop = \limits^{\rm HOH} {\mathop \to \limits^{\rm HOH}} {\rm CH_3CHO} + {\rm NH_3}$ 

73 **(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  $C_6H_5NH_2+CHCl_3KOH\rightarrow C_6H_5NC+KCl+H_2O$ 

75 **(a)** 

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

77 **(b)** 

Only primary amines give positive carbylamine test

78 **(a)** 

Primary amines have tendency of forming H-bonds

- 81 **(c)** 
  - 7. 0

 $CH_3 - C_2 - CH_3$   $sp^3 \quad sp^2 \quad sp^2$ 

(b) 0

 $CH_3 - C_2 - OH$   $sp^3 sp^2$ 

(c)

 $CH_3 - C - \equiv N$  $sp^3 \quad sp$ 

(d) (

 $CH_3 - C_2 - NH_2$   $sp^3 sp^2$ 

Acetonitrile does not contain  $sp^2$  hybridised carbon.

85 **(b)** 

Due to

+ ve *IE* in alkyamines and resonance in C<sub>6</sub>H<sub>5</sub>NH<sub>2</sub>.

86 **(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

87 **(b)** 

 $C_6H_5NH_2 \xrightarrow{Diazotisation} C_6H_5N_2Cl + H_2O$ 

89 **(c**)

Tertiary amines do not have replaceable H-atom.

90 **(b)** 

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

91 **(a**)

The order of basicity among the following amines is

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

92 **(d**)

Concentration does not affect the basis strength of amines

93 **(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.

95 **(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

96 **(d)** 

R—N=C=0  $\xrightarrow{\text{HOH}} R$ NH<sub>2</sub> + H<sub>2</sub>CO<sub>3</sub>

97 **(b** 

Biuret formed gives violet colour with  $\text{CuSO}_4$  in alkaline medium.

98 **(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.

# 99 **(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.

$$CH_3NH_2(1^\circ), (CH_3)_2NH(2^\circ), (CH_3)_3N(3^\circ), C_6H_5CH_1$$
 $II \qquad III \qquad IV$ 
 $II > I > III > V$ 

## 100 **(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.

$$\begin{array}{c} \overset{\text{HCI}}{\text{CH}_3\text{NH}_2} \overset{\text{HCI}}{\longrightarrow} \text{CH}_3\text{NH}_2. \\ \text{HCI} \\ \text{CH}_3\text{NH}_2 + \text{HNO}_2 \overset{\text{NaNO}_2/\text{HCI}}{\longrightarrow} \text{CH}_3\text{OH} + \text{N}_2 \uparrow + \text{H}_2\text{O} \\ \\ \text{methyl alcohol} \end{array}$$

# 101 **(b)**

 $2^{\circ}$  amines are more basic than  $1^{\circ}$  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 
$$CH_3-C$$

$$NH_2$$

due to resonance makes it more basic.

#### 102 (c)

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

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

## 103 (c)

$$C_2H_5NH_2 + Na \rightarrow C_2H_5NHNa + \frac{1}{2}H_2$$

#### 104 (d)

$$CH_3CN \xrightarrow{HOH} CH_3COOH$$

# 105 (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  $C_3H_7NH_2$ 
 $C_3H_9N \xrightarrow{KOH+CHCl_3} Carbylamine_{(R-NC)} \xrightarrow{Reduction}$ 

$$\begin{array}{c} \text{CH}_3 \\ \text{CH-NH-CH}_3 \Longrightarrow \text{Alkyl part is} \\ \text{CH}_3 \\ \text{Isopropyl methyl amine} \end{array} \\ \begin{array}{c} \text{CH}_3 \\ \text{CH}_3 \end{array}$$

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

# 109 (c)

Reduction of nitrobenzene by Zn and NH<sub>4</sub>CI gives N-phenyl hydroxylamine.

$$+4[H]$$
  $Zn/NH_4Cl$   $+H_2O$ 

N-phenyl hydroxylamine

# 111 (c)

Carbylamine reaction is used to prepare *iso* cyanides.

# 112 **(a)**

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

# 114 **(d)**

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

$$CH_3$$
— $O$ — $N$ = $O$  and  $CH_3OCH_3$ .

#### 117 **(c)**

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

# 119 **(d)**

CH<sub>3</sub>CHO

$$\xrightarrow{\text{H}_2\text{NOH}} \text{CH}_3\text{CH} = \text{NOH} \xrightarrow{\text{Reduction}} \text{CH}_3\text{CH}_2\text{NH}_2 \xrightarrow[-\text{H}_2\text{O}]{\text{NOCl}} \text{CH}_3\text{CH}_2$$

Acetaldehyde ethyl amine ethyl chloride

$$(A) (B)$$

# 120 **(a)**

$$R - C \equiv N$$

 $\therefore$  Alkyl group is attached to carbon.

#### 122 **(c)**

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

# 123 **(d)**

Nitrene is not the intermediate of Gabriel's phthalimide reaction

#### 124 **(b)**

$$NH_2CONH_2 + HNO_3 \rightarrow NH_2CONH_2 \cdot HNO_3$$

#### 126 (c)

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

#### 128 **(b)**

Ethyl bromide reacts with KCN to give ethyl cyanide.

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

130 **(b)** 

$$\mathsf{C_6H_5NO_2} \xrightarrow[[H]]{\mathsf{Sn/HCl}} \mathsf{C_6H_5NH_2}$$

Nitrobenzene Aniline Nitrobenzene in reduction with Sn and HCI produce aniline. Hence, 'X' is identified as –  $NH_2$  group.

131 **(d)** 

$$\begin{array}{c} \text{CH}_3\text{I} + \text{NH}_3 \longrightarrow \text{CH}_3\text{NH}_2 \stackrel{\text{CH}_3\text{I}}{\longrightarrow} (\text{CH}_3)_2\text{NH} \\ \stackrel{\text{CH}_3\text{I}}{\longrightarrow} (\text{CH}_3)_3\text{N} \stackrel{\text{CH}_3\text{I}}{\longrightarrow} (\text{CH}_3)_4\text{N}^+\text{I}^- \end{array}$$

137 (c)

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

138 (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

139 **(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  $NH_2COOC_2H_5 + NH_3 \rightarrow NH_2CONH_2 + 2C_2H_5OH$ Ethyl urethane urea  $COCl_2 + 2NH_3 \rightarrow NH_2CONH_2 + 2HCl$ phosgene urea

140 **(c)** 

Tertiary amine can be directly oxidised to corresponding nitro compound by KMnO<sub>4</sub>.

141 (a)

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

142 (c)

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

143 (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  
(B)

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

144 **(b)** 

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

145 **(b)** 

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

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

146 **(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.

147 (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.

148 (d)

$$\begin{array}{c|c} NH_2 & NH_2 .H_2SO_4 & NHSO_3H \\ \hline \\ \hline \\ H_2SO_4 & \hline \\ \hline \\ \hline \\ -H_2O & \hline \\ \end{array}$$

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

150 **(b)** 

$$C_2H_5NH_2 + CHCI_3 + 3KOH$$
  
 $\rightarrow C_6H_5NC + 3KCI + 3H_2O$ 

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

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

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

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

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

#### 154 (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.

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

## 156 (a)

Due to  $sp^3$ -hybridisation and lone pair of electron (like NH<sub>3</sub>)

# 157 **(b)**

$$\begin{array}{c} C_6H_5NO_2 \xrightarrow{Sn/HCI} C_6H_5NH_2 \\ \xrightarrow{C_6H_5COCI} C_6H_5CONHC_6H_5 \end{array}$$

Nitrobenzene aniline benzanilide

# 160 (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

# 161 **(b)**

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

$$\mathrm{C_2H_5NH_2} + \mathrm{HNO_2} \rightarrow \mathrm{C_2H_5OH} + \mathrm{N_2} + \mathrm{H_2O}$$

# 162 **(b)**

Aniline reacts with Br<sub>2</sub>to give 2, 4, 6-tribromoaniline not bromoaniline as

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

2,4,6-tribromoaniline

163 (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

# 167 **(d)**

Reduction of NO<sub>2</sub> group to NH<sub>2</sub>group is taking place by Fe/HCI.

# 172 **(b)**

Benzaldehyde condenses with N, N-diamethyl aniline in presence of anhydrous ZnCl<sub>2</sub> to give malachite green

$$\begin{array}{c} \text{H-} & \text{CH}_3 \\ \text{CH}_3 & \text{CH}_3 \\ \text{CH}_3 & \text{CH}_2 \\ \text{CH}_3 & \text{-H}_2 \text{O} \end{array}$$

#### 174 (a)

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

# 175 (c)

$$R_2$$
NH  $\stackrel{[O]}{\longrightarrow} R_2$ N—N $R_2$ 

#### 176 (d

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

## 178 (d)

HNO<sub>2</sub> reacts to give an alcohol means the compound is primary amine.

C<sub>5</sub>H<sub>13</sub>N means C<sub>5</sub>H<sub>11</sub>NH<sub>2</sub>(primary amine)

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

Pentan-2-amine

# 179 **(b)**

 $CH_3CH_2 - O - N = O$  is a nitrite derivative, hence

it is not a nitro derivative.

180 (d)

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

reagent.

It is used for the distinction of  $1^{\circ}$ ,  $2^{\circ}$ ,  $3^{\circ}$  amine.

181 **(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

184 (d)

It is not used in purification of water.

185 **(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

186 **(c)** 

Methylaminomethane is trivial name of Nmethyl methanamine (CH<sub>3</sub>)<sub>2</sub>NH.

187 **(c)** 

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

190 **(b)** 

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

191 (c)

192 (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

$$H_3CH_2NO_2$$
  $\xrightarrow{HONO}$   $CH_3C=NOH$   $\xrightarrow{NO_2}$   $CH_3C=NOH$   $\xrightarrow{acid}$   $NO_2$   $CH_3-C=NO^ Na^+$  sodium salt (red colour)

194 (d)

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

Benzylamine

(b) 
$$C_6H_5CH_2CONH_2 \xrightarrow{Br_2/KOH} C_6H_5CH_2NH_2$$
  
Benzylamine

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

(d) 
$$C_6H_5CH_2NC \xrightarrow{\text{LiAlH}_4} C_6H_5CH_2NHCH_3$$
  
2°amine

195 (c)

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

196 (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

197 (a)  $CH_2N_2 + CH_3COOH \longrightarrow CH_3COOCH_3 + N_2_{Ester}$ 

198 **(a)** 

Follow text.

200 **(b)** 

Only 1° aromatic amine (primary aromatic amine) from diazonium salts at low temperature (0° – 5°C). A reaction in which – NH<sub>2</sub> group is converted into diazo group ( $-N^+ \equiv N$ ) is called diazotization. Diazotized salts are stable in cold aqueous solution.

$$C_{6}H_{5}NH_{2}+HC1 \xrightarrow{0^{\circ}-5^{\circ}C} C_{6}H_{5}NH2C1^{-}$$
 $C_{6}H_{5}NH_{2}+HC1 \xrightarrow{0^{\circ}-5^{\circ}C} HNO_{2} +NaC1$ 
 $C_{6}H_{5}NC1^{-}+HNO_{2} \xrightarrow{0^{\circ}-5^{\circ}C} C_{6}H_{5}N_{2}C1^{-}+2H_{2}O$ 
 $C_{6}H_{5}NH_{2}$ 
 $C_{6}H_{4}$ 
 $C_{6}H_{5}NH_{2}$ 
 $C_{6}H_{4}$ 

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

202 **(c)**

$$NH_{2} \longrightarrow NH_{2} \longrightarrow NH_{2} \longrightarrow NH_{2} \longrightarrow NO_{2}$$
aniline
$$NO_{2} \longrightarrow MO_{2} \longrightarrow NO_{2}$$

$$-nitroaniline \longrightarrow NO_{2} \longrightarrow NO_{2}$$

(51%)

Nitration of aniline also gives *m*-nitro aniline in strong acidic medium because in strong acidic condition protonation of - NH<sub>2</sub> group gives anilinum ion (+NH<sub>3</sub>) which is deactivating in nature and of *m*-directive nature

$$RCONH_2 + NaOBr$$
  
 $\rightarrow RNH_2 + NaBr + Na_2CO_3$   
 $+ 2H_2O$   
 $'X'$ 

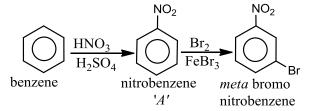
### 204 **(b)**

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

$$C_6H_5NH_2 + CHCI_3 + 3KOH(alc.)$$
  
 $\rightarrow C_6H_5NC + 3KCI + 3H_2O$   
aniline chloroform phenyl  
isocyanide

This reaction is called carbylamine reaction and it |219|is actually the test of primary amines.

#### 206 (a)



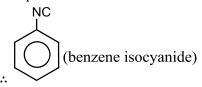
 $-NO_2$  is a *meta* directing group. As it is also a deactivating group, so no chance of introduction of second - Br atom.

## 207 **(c)**

$$R$$
—NC+2HgO $\rightarrow R$ NCO+Hg<sub>2</sub>O

# 210 (a)

Isocyanides (carbylamines) are foul odour compounds.



As foul odour

## 213 (a)

This is carbylamine reaction.

# 214 (d)

When aqueous solution of benzene diazonium

chloride boiled, it gives phenol.

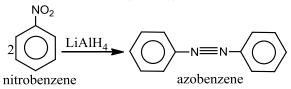
$$N_2$$
+Cl-OH
benzene
diazonium
chloride

## 215 (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

# 217 (c)

Nitrobenzene on reduction with lithium aluminium hydride (LiAIH<sub>4</sub>) gives azobenzene.



## 218 **(b)**

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

Nitrobenzene on nitration gives *m*-dinitro benzene as – NO<sub>2</sub> group is meta-directing.

In this reaction the attacking reagent is  $NO_2^+$ .

#### 220 (d)

Bases react with acid to form salt.

- : Amines are basic in nature.
- ∴It forms salt on reaction with H<sub>2</sub>SO<sub>4</sub>  $2RNH_2 + H_2SO_4 \rightarrow [RNH_3]_2^+SO_4^{2-}$

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

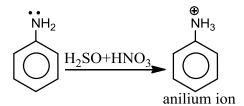
222 **(d)** 

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

acetvl chloride aniline acetanilide

#### 223 (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.



# 224 (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.

# 226 **(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  $NH(CH_3)_2 > CH_3NH_2 > N(CH_3)_3 > NH_3$ 

# 227 **(c)**

Aliphatic and aromatic primary amines on warming with  $\mathrm{CHCI}_3$  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.

$$NH_2$$
 +CHCl<sub>3</sub>+3(alc)KOH Warm + 3KCl+3H<sub>2</sub>O  $H_3$ C  $P$ -toluidine 4-methyl phenyl isocyanide

#### 229 **(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.

# 231 **(c)**

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

$$CH_3CH_2CH_2NH_2 + CHCI_3 + 3KOH$$
  
 $\rightarrow CH_3CH_2CH_2NC + 3KCI + 3H_2O$   
 $n$ -propyl amine (alco.) propyl isocyanide

# 232 **(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

# 233 **(a)**

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

$$NH_2$$

$$+ CH_3CHO - H_2O$$

$$+ CH_3CHO - Schiff's base or anil$$

#### 234 (a)

Coupling of diazonium salts takes place in the following order as

# 235 (c)

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

## 237 **(b)**

$$C_2H_5NH_2$$
 $\xrightarrow{HNO_2}$   $C_2H_5OH \xrightarrow{PCl_5}$   $C_2H_5Cl \xrightarrow{NH_3}$   $C_2H_5NH_2$ 
Ethyl amine ethyl alcohol ethyl chloride ethyl amine

(A) (B)

# 238 **(c)**

 $\rm K_2CO_3$  is formed in Hofmann's degradation reactio

# 240 **(d)**

Aniline undergoes diazotisation.

# 241 **(c)**

The conversion of —CN to —CH<sub>2</sub>NH<sub>2</sub> by catalytic reduction is called Mendius reaction.

$$RBr + KCN \longrightarrow RCN \xrightarrow{\text{Reduction}} RCH_2NH_2$$

# 245 (d)

Aniline reacts with bromine and give white ppt. of

2, 4, 6 tribromoaniline.

$$+3 \text{ Br}_2$$
 $+3 \text{ HBr}$ 
white ppt.

246 **(b)** 

Nitromethane forms methyl hydroxylamine on reduction in netural medium with Zn/NH<sub>4</sub>CI.

$$CH_3NO_2 + 4[H] \xrightarrow{Zn/NH_4CI} CH_3NHOH + H_2O$$

Nitromethane

N-methyl hydroxyl

amine

## 251 (c)

In Hofmann elimination reaction, less substitudes alkene is the main product

$$\begin{bmatrix} \mathsf{CH_3} \\ \mathsf{CH_3CH_2CH_2} \overset{+}{\mathsf{N}} - \mathsf{CH_2CH_3} \\ \mathsf{CH_3} \end{bmatrix} \mathsf{OH}^{-} \xrightarrow{\Delta} \\ \mathsf{CH_3CH_2CH_2} - \mathsf{N}(\mathsf{CH_3})_2 + \mathsf{CH_3CH} = \mathsf{CH_2} + \mathsf{CH_2} = \mathsf{CH_2} \\ (minor) \quad major \\ \end{bmatrix}$$

253 **(b)** 

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

254 (d)

C<sub>6</sub>H<sub>5</sub>COCl has no reaction with carboxylic acids.

257 **(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.

258 (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.

260 **(c)** 

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

2° 1° 3°

261 **(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.

264 **(b)** 

$$-C \equiv N \xrightarrow{Na} -C \xrightarrow{Na} NH$$

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

265 **(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.

266 **(d)** 

$$\begin{array}{c|c}
 & NO_2 \\
\hline
 & NH_2 \\
\hline
 & NaNO_2 \\
\hline
 & HCl, 0^{\circ}C
\end{array}$$

$$\begin{array}{c}
 & (Y) \\
 & \text{diazonium salt} \\
\hline
 & NaNH_2
\end{array}$$

 $\therefore Z$  is aniline

267 **(b)** 

OH 
$$_{\text{H}}^{\text{OH}}$$
  $_{\text{H}}^{\text{Me}}$   $_{\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_3}^{\text{Me}}$   $_{\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_3}^{\text{OH}}$   $_{\text{CH}_2}^{\text{OH}}$   $_{\text{$ 

There are four  $\beta$ - hydrogens, in this quaternary ammonium salt.

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

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

268 **(b)** 

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

## 269 **(b)**

Nitrobenzene is reduced by Zn and alcoholic KOH into hydrazobenzene.

$$\begin{aligned} 2 C_6 H_5 NO_2 + 10 H & \xrightarrow{Zn+alc.KOH} C_6 H_5 - NH - NH \\ & - C_6 H_5 + H_2 O \\ & \text{hydrazobenzene} \end{aligned}$$

$$CH_3NH_2 + CHCI_3 + 3KOH$$
  
 $\rightarrow CH_3NC + 3KCI + 3H_2O$ 

 $CH_3NC$  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.

# 274 (a)

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

# 275 (a)

$$CH_3CH_2NH_2 \xrightarrow{NaNO_2+HCl} CH_3 CH_2 OH + N_2 + H_2O$$
  
+ NaCl

Ethylamine

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

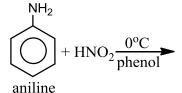
$$CH_3CONH_2 \xrightarrow{NaNO_2+HCl} CH_3COOH + N_2 + H_2O$$
+ NaCl

Acetamide

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

#### 276 **(b)**

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



1°aromatic amine

∴Dye test is used to distinguish between

1°aliphatic and

1° aromatic amine

#### 277 (c)

Presence of  $\alpha$ -H atom is the main condition for exhibiting tautomerism.

The reactant taken in reaction (C) does not contain any  $\alpha$ -H atom, thus the product (Y)will

also show the absence of  $\alpha$ -H atom, Hence, Y will show tautomerism

# 278 (a)

8. 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

$$(\mathsf{COOC}_2\mathsf{H}_5)_2 + \mathsf{HN}R_2 \to |$$

$$COOC_2H_5 + C_2H_5OH$$

2°amine liquid

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

3° amino

# 279 **(a)**

An alkyl cyanide is dissolved in ether or better in ethyl formate or ethyl acetate, and reduced with  $SnCI_2$  and HCI and then steam distilled. The whole process is called Stephen reaction. In this process alkyl cyanide is reduced to aldehyde.

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

$$CH_{3} \longrightarrow CH = \begin{vmatrix} O \\ H_{2} \\ - NH \cdot HCl \end{vmatrix} \xrightarrow{H_{2}O} CH_{3}CHO + NH_{4}Cl$$

There is no analogous method for the preparation of ketones.

#### 282 (a)

Azoxybenzene is the main product when reducing agent used is Na<sub>3</sub>AsO<sub>3</sub>/NaOH

# 283 **(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.

# 284 **(d)**

Aniline on oxidation with  $Na_2Cr_2O_7$  and  $H_2SO_4$  gives *p*-benzoquinone.

$$NH_2$$
 $Na_2Cr_2O_7/H_2SO_4$ 
 $p$ -benzoquinone (quinone)

#### 285 (a)

As the molecular mass increase, boiling point increases. In case of isometric amines, however, as the number of H-atoms attached directly to N-atom decreases, boiling point decreases because tendency to form H-bonds decreases. Hence, the order of boiling points of given amines is  $CH_3NH_2 < (CH_3)_3 N < CH_3CH_2N(NH_3)$   $< CH_3CH_2CH_2NH_2$ 

# 288 **(b)**

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

$$0 \qquad 0$$

$$|| \qquad ||$$

$$CH_3 - C \equiv N \xrightarrow{H_2 0} CH_3 - C - NH_2 \xrightarrow{H_2 0} CH_3 - C$$

$$- OH + NH_2$$

Acetonitrile

acetamide

acetic acid

# 289 (a)

 $N \equiv C - CH = CH - C \equiv N$ ; It has five  $\pi$ -bonds.

#### 292 **(b)**

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

$$CCH_3CH_2NH_2 + CHCI_3 + 3KOH$$
  
 $\rightarrow CH_3CH_2NC + 3KCI + 3H_2O$   
9. (B)

This reaction is known as carbylamine reaction

#### 294 **(b)**

By using  $\mathrm{H}_2\mathrm{S}$ ,  $\mathrm{NH}_3$  as reagent, selective reduction takes place

$$NO_2$$
 $H_2S, NH_3$ 
 $NO_2$ 
 $NH_2$ 
 $NH_2$ 

# 296 **(b)**

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

299 (c)

EDTA is ethylenediamine-tetra acetic acid.

300 (c)

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

301 **(b)** 

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

302 **(b)** 

$$R - \text{CN} + 4[\text{H}] \xrightarrow{\text{LiAIH}_4} R - \text{CH}_2\text{NH}_2$$
  
Alkyl nitriles primary amine

# 303 **(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  $C_nH_{2n+3}N$ .

# 304 **(b)**

Biuret test is given by compounds having — CONH<sub>2</sub>gp.

305 **(b)** 

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

309 (c)

As we know, benzenediazonium salt forms brilliant coloured dye with  $\beta$ -naphthol, the compound under consideration must be p-toludine (c) as it is a primary aromatic amine. Primary aromatic amine, on treatment with NaNO $_2$  in dil. HCI forms the corresponding diazonium chloride salt.

$$NH_2$$
 $N_2$ 
 $N_2$ 

## 310 (a)

 $-OCH_3$  is strongest electron releasing group (+M effect) which opposes most the dispersion of lone pair of electron of nitrogen into the ring. Thus,  $-OCH_3$  being at *para* position imparts hifhest basicity.  $NO_2$  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.

$$\mathsf{O_2N} \overset{\mathsf{NH_2}}{\longleftarrow} \mathsf{NH_2} \overset{\mathsf{NH_2}}{\longleftarrow} \mathsf{NH_2}$$

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

317 (a)

 $R_2$ NH + HNO<sub>2</sub>  $\rightarrow$   $R_2$ N—N=0 + H<sub>2</sub>O Nitrosoamines are carcinogens.

318 **(a)** 

$$R$$
—  $CN \xrightarrow{Reduction} RCH_2CH_2$ 

319 (a)

$$CH_3NH_2 + C_6H_5SO_2CI$$

$$\rightarrow C_6H_5SO_2NHCH_3$$

$$\xrightarrow[(alkali)]{NaOH} C_6H_5SO_2N(Na)CH_3$$

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

Reagent sulphonamide

320 **(b)** 

It gives diazonium salt.

It is known as diazotization reaction.

321 (a)

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

$$C_2H_5NH_2 \xrightarrow{CH_3COCI} C_2H_5NHCOCH_3$$

N-ethyl acetamide

322 **(b)** 

 $1^{\circ} aliphatic amines on reduction with <math display="inline">\text{HNO}_2$  form alcohol and evolve  $\text{N}_2.$ 

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

323 **(b)** 

 $NH_2CONH_2 + HNHCONH_2 \rightarrow NH_2CONHCONH_2 + NH_3$ 

324 **(d)** 

it is a example of carbylamines reaction

325 **(b)** 

$$C_6H_5NH_2 + CH_3COCl \rightarrow C_6H_5NHCOCH_3$$

326 **(b)** 

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

327 **(a)** 

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

328 **(c)** 

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

10.

Acetic acid

acetamide

(*B*)

(C)

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

Methyl amine (*D*)

329 (a)

Aliphatic primary amines on treatment with NaNo<sub>2</sub>/HCI gives alcohols with evolution of nitrogen gas.

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

330 **(b)** 

$$RNH_2 \xrightarrow{HNO_2} ROH + H_2O + N_2 \uparrow$$
11. (B) (C)

331 (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

 $pK_b$ 

332 **(d)** 

The mixture of  $SnCl_2 + HCl$ , reduces, -CN group partially

333 **(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

334 **(b)** 

The conversion of —CN to —CH<sub>2</sub>NH<sub>2</sub> by catalytic reduction is called Mendius reaction.

335 (c)  $RCONH_2 + NaOH \rightarrow RCOONa + NH_3$ 

336 **(b)** 

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

340 **(b)** 

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

341 (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

342 **(b)** 

Aniline on oxidation with per acetic acid, CH<sub>3</sub>CO<sub>3</sub>H gives nitrobenzene

343 (d)

 $C_3H_9N$  may have the structures as:  $(CH_3)_3N_{,(a)}CH_3CH_2CH_2NH_2, (CH_3)_2CHNH_2$ 

344 **(b**)

Amines possess fishy smell.

345 (c)

$$CH_3NC \xrightarrow{Reduction} CH_3NHCH_3$$

346 (c)

 $\rm C_2H_5NH_2$  is stronger base than  $\rm NH_3$ . 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.)

347 (d)

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

$$$\operatorname{NH}_2$$$
  $|$   $$\operatorname{CH}_3-\operatorname{CH}-\operatorname{CH}_3$$   $1^\circ$  amine

349 (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 NH $_3$ . 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.

350 (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)

$$N=N.CI+H- N(CH_3)_2 \xrightarrow{NaOH} N=N-(CH_3)_2 -HCI$$

352 (c)

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

354 (a)

$$\begin{array}{c} \operatorname{CH_3CH_2I} \xrightarrow{\operatorname{NaCN}} \operatorname{CH_3CH_2CN} \xrightarrow{\operatorname{OH}^-} \operatorname{CH_3CH_2CONH_2} \\ \xrightarrow{\operatorname{Br_2/NaOH}} \operatorname{CH_3CH_2NH_2} \end{array}$$

355 **(b)** 

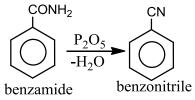
This is carbylamine reaction which is used to distinguish  $1^{\circ}$  amines from other amines. The reaction is given by  $1^{\circ}$  amines only.

$$C_2H_5NH_2 + CHCI_3 + 3KOH$$
  
 $\rightarrow C_2H_5N \stackrel{=}{\rightarrow} C + 3KCI + 3H_2O$ 

$$RNH_2 + CHCI_3 + 3KOH$$
  
 $\rightarrow R - N \xrightarrow{=} C + 3KCI + 3H_2O$   
1° amine chloroform isocyanide (bad smelling)

356 (d)

With P<sub>2</sub>O<sub>5</sub> benzamide loses a water molecule and gives benzonitrile.



Here, P<sub>2</sub>O<sub>5</sub> is a dehydrating agent.

357 (a)

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

Primary amines > secondary amines > tertiary amines

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

358 (c)

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

$$RCONH_2 + Br + 4KOH$$
  
 $\rightarrow RNH_2 + K_2CO_3 + 2KBr + 2H_2O$   
amide amine

361 (d)

$$\leftarrow$$
 +CHCl<sub>3</sub>+ alc.KOH  $\leftarrow$   $\leftarrow$  CH<sub>3</sub>

*p*-toluidine (a carbylamine reaction)

363 **(d)** 

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

364 (c)

R—NH<sub>2</sub> + HNO<sub>2</sub>—ROH + N<sub>2</sub> + H<sub>2</sub>O; But note that CH<sub>3</sub>NH<sub>2</sub> gives CH<sub>3</sub>ONO or CH<sub>3</sub>OCH<sub>3</sub> on treating with HNO<sub>2</sub>.

365 (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

$$\langle \bigcirc \rangle$$
-CH<sub>2</sub>- $\mathring{N}$ H<sub>2</sub>

366 **(d)** 

Carbylamine reaction is given by aliphatic and aromatic primary amine.

CH<sub>3</sub>CN does not give carbylamine reaction with chloroform because it is not an amine.

CH<sub>3</sub>CN undergoes acidic hydrolysis to give carboxylic acid.

$$CH_3CN \xrightarrow{H_2O/H^+} CH_3CONH_2 \xrightarrow{H_2O/H^+} CH_3COOH + NH_3$$

acetic acid

CH<sub>3</sub>CN undergoes alkaline hydrolysis to give salt of carboxylic acis.

$$\text{CH}_{3}\text{CN} \xrightarrow[\text{H}_{2}\text{O}]{\text{NaOH}} \text{CH}_{3}\text{CONH}_{2} \xrightarrow[\text{H}_{2}\text{O}]{\text{NaOH}} \text{CH}_{3}\text{COONa} + \text{NH}_{3}$$

Sodium acetate

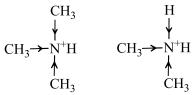
CH<sub>3</sub>CN tautomerises to give methyl isocyanide.

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

367 (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 acid of amines because of +ve IE of methyl group which disperses +ve charge on N-atom more effectively in  $3^{\circ}$  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  $2^{\circ} > 1^{\circ} > 3^{\circ} > NH_3$ .

368 (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

369 (c)

$$(CH_3)_2CH.NH_2 + 2CH_3COCI \xrightarrow{-HCI} (CH_3)_2CH$$
  
-  $N(COCH_3)_2$ 

*iso*-propyl amine (*ter*-amine) (*pri* amine)

370 (a)

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

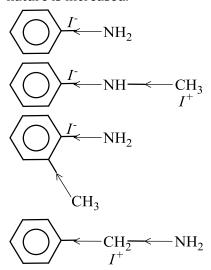
# 371 (a)

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

 $2^{\circ}$  amine is more basic than  $3^{\circ}$  amine because  $2^{\circ}$  amine is stabilized by hydrogen bonding with solvent molecule.

# 372 **(d)**

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



 ${\rm C_6H_5}$  decreases electron density at N-atom thus basic nature is decreased. (Lone-pair on N in aniline compounds is delocalised along with  $\pi$ -electrons in benzene).

Thus, (d) is the strongest base.

# 374 **(d)**

Isocyanide on hydrolysis forms primary amine

not ammonia.

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

375 **(b)** 

Less substituted alkene is the main product

376 **(a)** 

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

377 **(c)** 

$$SnCI_2 + 2HCI(g) \rightarrow SnCI_4 + 2[H]$$
  
 $CH_3CN + 2[H] + HCI \rightarrow CH_3CH$   
 $= NH. HCI \xrightarrow{H_2O} CH_3CHO + NH_4CI$ 

$$(X)$$
  $(Y)$ 

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

# 378 (c)

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

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

