

it is a example of carbylamines reaction

(a)

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

$$R - CN + 2H_2ONaOHR - COOH + NH_3$$

10 **(a)**

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

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(b)

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



12 **(b)**

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2 amines are more basic than 1 amines due to +ve IE of -CH₃ gp. In amide the resonance give rise to less availability to electron pair for coordination and thus less basic. The negative



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

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(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.

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(b)

$$C_6H_5NO_2$$
 Sn/HClC₆H₅NH₂

Nitrobenzene Aniline

Nitrobenzene in reduction with Sn and HCI produce aniline. Hence, 'X' is identified as -N H₂ group.

17 **(d)**

 $CH_{3}CHOH_{2}NOHCH_{3}CH = NOHReductionCH_{3}CH_{2}NH_{2}NOClCH_{3}CH_{2}Cl$

Acetaldehyde	ethyl amine	ethyl chloride

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

19 **(b)**

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

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(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.

ANSWER-KEY										
Q .	1	2	3	4	5	6	7	8	9	10
A.	Α	Α	Α	D	Α	С	Α	D	Α	A
Q .	11	12	13	14	15	16	17	18	19	20
A.	В	В	Α	С	Α	В	D	D	В	С