DPP

DAILY PRACTICE PROBLEMS

Class: XIIth Date:

Solutions

Subject : CHEMISTRY

DPP No.: 2

Topic:- Amines

1 **(b)**

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

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

2 **(c)**

 $C_2H_5NH_2$ is stronger base than NH_3 . The presence of alkyl group on N-atom intensifies –ve charge on N-atom and thus, electron pair is donated more readily.

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

H-NH₂

3 **(a)**

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

$$(CH_3)_4 N^+ OH^- \Delta (CH_3)_3 N + CH_3 OH$$

methanol

4 **(c)**

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

5 **(d)**

$${\sf C_6H_5NH_2+CI-COCH_3\underline{NaOHC_6H_5NHCOCH_3+HCI}}$$

aniline acetyl chloride acetanilide

6 **(d)**

C₃H₉N represent following structures

CH₃CH₂CH₂NH₂

Propanamine

(1°amine)

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 $CH_3CH_2 - N - CH_3$

N-methyl ethanamine

(2°amine)

 CH_3

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 $CH_3 - N - CH_3$

N,N-dimethyl methanamine

(3°amine)

7 **(c)**

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

The reactant taken in reaction (C) does not contain any α -H atom, thus the product (Y) will also show the absence of α -H atom, Hence, Y will show tautomerism

8 **(a)**

 $R_2NH + HNO_2 \rightarrow R_2N - N = O + H_2O$ Nitrosoamines are carcinogens.

9 **(b**)

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

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 $CH_3 - C \equiv NH_2OCH_3 - C - NH_2H_2OCH_3 - C - OH + NH_3$

Acetonitrile acetamide acetic acid

11 **(b)**

Methyl cyanide gives acetic acid on hydrolysis.

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 $CH_3CNH_2O/H^+CH_3 - C - NH_2H_2O/H^+CH_3COOH + NH_3$

12 **(c)**

 $2RNH_2 + 2HCl + PtCl_4 \rightarrow (RNH_3)_2 PtCl_6; RNH_3 AuCl_4$

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

14 (c)

Aniline on diazotization in cold (at 0° to 5°C) gives 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)

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

16 **(c)**

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

$$2 \underbrace{ \frac{\text{LiAlH}_4}{\text{nitrobenzene}}} \underbrace{ \frac{\text{LiAlH}_4}{\text{azobenzene}}}$$

17 **(c)**

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

19 **(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.

20 **(d)**

All primary (aliphatic) amines give alcohol with HNO_2 except CH_3NH_2 which forms $CH_3-O-N=0$ and CH_3OCH_3 .



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

