# Pharmaceutical Organic Chemistry-1

**Chapter-8: Amines** 

# Structure and Classification of Amines

 Amines are compounds that derived from ammonia by replacement of one, two, or three hydrogens by alkyl or aryl groups.

Aliphatic amines contain only alkyl groups bonded directly to the nitrogen atom.

$$CH_3$$
— $N$ — $H$   $CH_3$ — $N$ — $CH_3$   $CH_3$ — $N$ — $CH_2$ — $CH_2$ — $CH_3$ 

Aliphatic amines

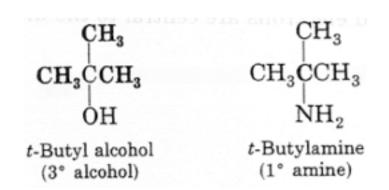
 Aromatic amines are those in which one or more aryl groups are bonded directly to nitrogen.

# Classification and Structure of Amir

The relation between ammonia and amines is illustrated by the following structures:

H—N—H R—N—H R—N—R R—N—R H H H R R

- Amines are classified as primary, secondary, or tertiary, depending on whether one, two, or three organic groups are attached to the nitrogen.
- NOTE:

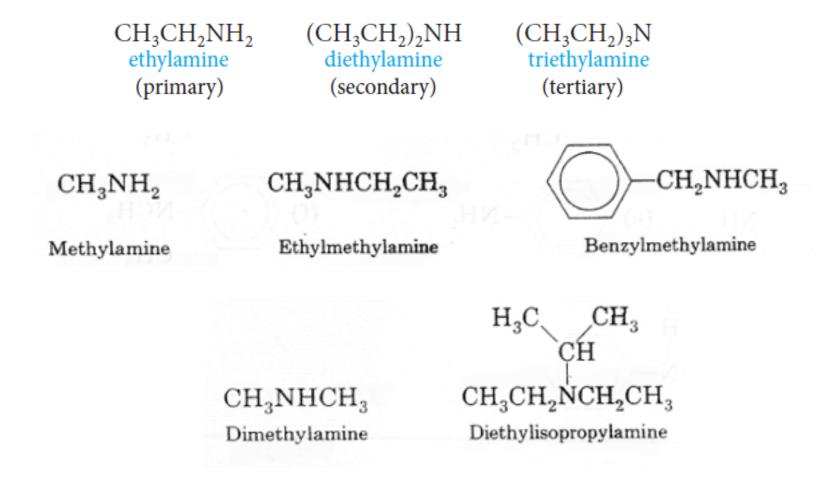


- ?t-butyl alcohol is a tertiary alcohol (because three carbons are attached to the carbinol carbon).
- r-butyl amine is a primary amine (because only one carbon is attached directly to the nitrogen atom).

# **Nomenclature of Amines**

C o m m o n

Marrines are named by specifying the alkyl groups attached to the nitrogen and adding the suffix —amine (Alkylamine).



# Nomenclature of Amines

# I U P A C System

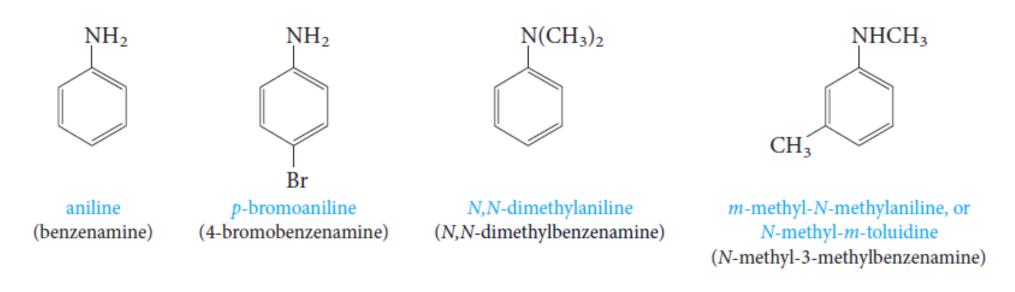
Amines can be named as alkanamines.

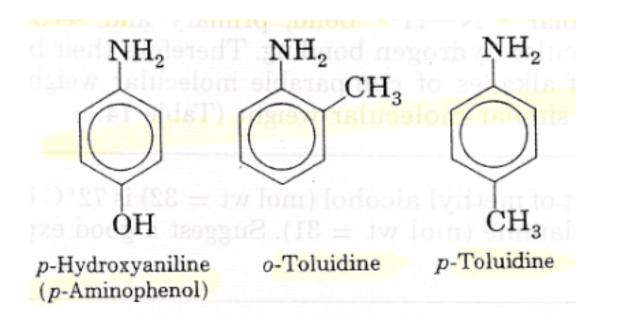
# Nomenclature of Amines

#### I U P A C

Syst then other functional groups are present, the amino group, -NH<sub>2</sub>, is named as a substituent.

- Aromatic amines are named as derivatives of aniline.
- o In the IUPAC system, aniline is called benzenamine.





# **Physical Properties of Amines**

# Boiling

- Point thylamine and ethylamine are gases, but primary amines with three or more carbons are liquids.
- Primary amines boil well above alkanes with comparable molecular weights, but below comparable alcohols.

Intermolecular N-H···N hydrogen bonds are important and raise the boiling points of primary and secondary amines but are not as strong as the O-H···O bonds of alcohols.

The reason for this is that nitrogen is not as electronegative as evugen

alkane	CH <sub>3</sub> CH <sub>3</sub> (30) bp -88.6°C	CH <sub>3</sub> CH <sub>2</sub> CH <sub>3</sub> (44) bp -42.1°C
amine	$CH_3NH_2$ (31) bp $-6.3^{\circ}C$	$CH_3CH_2NH_2$ (45) bp +16.6°C
alcohol	CH <sub>3</sub> OH (32) bp +65.0°C	CH <sub>3</sub> CH <sub>2</sub> OH (46) bp +78.5°C

# Physical Properties of Amine

# Boiling

Potertiary amines are also polar compounds, but because hydrogen is not bonded to nitrogen, these amines are incapable of intermolecular hydrogen bonding.

Their boiling points are Lower than primary and secondary amines of identical molecular weights and Higher than those of alkanes of similar molecular weight.

# Solubility in

- Watthree classes of amines can form hydrogen bonds with the -OH group of water (that is, O-H···N).
  - Primary and secondary amines can also form hydrogen bonds with the oxygen atom in water: N-H···O.
  - Amines with up to six carbons show appreciable solubility in water.

# The Basicity of Amines

- The unshared pair of electrons on the nitrogen atom dominates the chemistry of amines.
- Because of this electron pair, amines are both basic and nucleophilic.
- Aqueous solutions of amines are basic because of the following equilibrium:

$$N: +H - \ddot{O}H \Longrightarrow N^+ - H + - \ddot{O}H$$
amine ammonium hydroxide ion ion

Flectron-donating groups increase the basicity of amines.

?Electron-withdrawing groups decrease their hasicity

One electron-donating group

Two electron-donating groups

# The Basicity of Amine

- Aromatic amines are much weaker than aliphatic amines or ammonia.
  - Example: aniline is less basic than cyclohexylamine.

The reason is the resonance delocalization of the unshared electron pair that is possible in aniline, but not in cyclohexylamine:

# **Preparation of Amines**

#### 1) Alkylation of Ammonia

Ammonia reacts with alkyl halides to give amines via a two-step process.
 The first step is a nucleophilic substitution reaction. The free amine can then be obtained from its salt by treatment with a strong base

$$H_3N: +R-X \longrightarrow R-NH_3X^ R-NH_3X^ RNH_2 + H_2O + Na^+X^ RNH_3$$
 $RNH_2 + H_3O + Na^+X^ RNH_3$ 
 $RNH_3$ 
 $RNH_3$ 

Primary, secondary, and tertiary amines can be similarly alkylated.

$$\begin{array}{c} \overset{+}{\text{RNH}_2} + \overset{+}{\text{R}} - \overset{+}{\text{X}} \longrightarrow \overset{+}{\text{R}_2 \text{NH}_2} \overset{+}{\text{X}} - \overset{\text{NaOH}}{\text{Maine}} \\ & \overset{+}{\text{R}_2 \text{NH}_2} + \overset{+}{\text{R}} - \overset{+}{\text{X}} \longrightarrow \overset{+}{\text{R}_3 \text{N}_1} + \overset{+}{\text{R}} - \overset{+}{\text{X}} \longrightarrow \overset{+}{\text{R}_3 \text{N}_1} + \overset{+}{\text{R}} - \overset{+}{\text{X}} \longrightarrow \overset{+}{\text{R}_3 \text{N}_1} + \overset{+}{\text{R}} - \overset{+}{\text{X}} \longrightarrow \overset{+}{\text{R}_4 \text{N}} + \overset{+}{\text{X}} - \overset{+}{\text{NaOH}} \\ & \overset{+}{\text{Secondary}} & \overset{+}{\text{amine}} & \overset{+}{\text{R}_3 \text{N}_1} + \overset{+}{\text{R}} - \overset{+}{\text{X}} \longrightarrow \overset{+}{\text{R}_4 \text{N}} + \overset{+}{\text{X}} - \overset{+}{\text{NaOH}} \\ & \overset{+}{\text{Secondary}} & \overset{+}{\text{amine}} & \overset{+}{\text{A}} & \overset{+}{\text{NaOH}} + \overset{+}{\text{R}_3 \text{N}_1} + \overset{+}{\text{R}} - \overset{+}{\text{X}} \longrightarrow \overset{+}{\text{R}_4 \text{N}} + \overset{+}{\text{X}} - \overset{+}{\text{NaOH}} \\ & \overset{+}{\text{Secondary}} & \overset{+}{\text{amine}} & \overset{+}{\text{MaOH}} & \overset{+}{\text{R}_3 \text{N}_1} + \overset{+}{\text{R}} - \overset{+}{\text{X}} \longrightarrow \overset{+}{\text{R}_4 \text{N}} + \overset{+}{\text{X}} - \overset{+}{\text{NaOH}} \\ & \overset{+}{\text{Secondary}} & \overset{+}{\text{amine}} & \overset{+}{\text{A}} & \overset{+}{\text{NaOH}} + \overset{+}{\text{NaOH}} \\ & \overset{+}{\text{Secondary}} & \overset{+}{\text{A}} & \overset{+}{\text{A}} & \overset{+}{\text{NaOH}} + \overset{+}{\text{NaOH}} + \overset{+}{\text{NaOH}} \\ & \overset{+}{\text{Secondary}} & \overset{+}{\text{A}} & \overset{+}{\text{A}} & \overset{+}{\text{NaOH}} + \overset{+}{\text{NaOH}} + \overset{+}{\text{NaOH}} \\ & \overset{+}{\text{Secondary}} & \overset{+}{\text{A}} & \overset{+}{\text{NaOH}} & \overset{+}{\text{A}} & \overset{+}{\text{NaOH}} + \overset{+}{\text{NaOH}} \\ & \overset{+}{\text{Secondary}} & \overset{+}{\text{A}} & \overset{+}{\text{NaOH}} & \overset{+}{\text{A}} & \overset{+}{\text{NaOH}} & \overset{+}{\text{NaOH}} \\ & \overset{+}{\text{Secondary}} & \overset{+}{\text{A}} & \overset{$$

# **Preparation of Amines**

### 2) Reduction of Nitro Groups

 The best route to aromatic primary amines is by reduction of the corresponding nitro compounds.

The nitro group is easily reduced, either catalytically with hydrogen or by chemical reducing agents.

# **Preparation of Amines**

#### 3) Reduction of

#### **Nitriles**

Reduction of nitriles (cyanides) gives primary amines.

$$R-C \equiv N \xrightarrow{\text{LiAlH}_4} RCH_2NH_2$$

# 4) Reduction of

Amides can be reduced to amines with lithium aluminum hydride.

$$R - C - N \xrightarrow{R'} \xrightarrow{\text{LiAlH}_4} RCH_2N \xrightarrow{R'} R'$$

$$R'' \xrightarrow{\text{R'}} RCH_2N \xrightarrow{R'} R'$$

$$R'' \xrightarrow{\text{R'}} RCH_2N \xrightarrow{\text{R'}} R'$$

$$R'' \xrightarrow{\text{R'}} RCH_2N \xrightarrow{\text{R'}} R'$$

#### **Reactions of Amines**

#### 1) Reactions with Acids: Salt

Formites General with strong acids to form alkylammonium salts.

#### 2) Acylation of Amines: Amides

Formation discondary amines react with acyl halides to form amides.

#### Reactions of Amines

#### 3) Imines

Formationnines, R-NH<sub>2</sub> or ArNH<sub>2</sub>, undergo nucleophilic addition with aldehydes or ketones in an acidic buffer to give substituted imines.