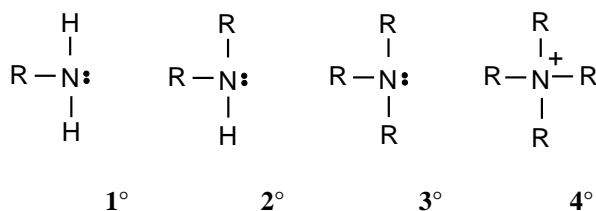


AMINES

Structure Contain the NH_2 group.

Classification **primary** (1°) amines
secondary (2°) amines
tertiary (3°) amines
quarternary (4°) **ammonium** salts



Aliphatic methylamine, ethylamine, dimethylamine

Aromatic NH_2 group is **attached directly** to the benzene ring (phenylamine)

Nomenclature Named after the groups surrounding the nitrogen + *amine*

Aliphatic amines

$\text{C}_2\text{H}_5\text{NH}_2$ ethylamine
 $(\text{CH}_3)_2\text{NH}$ dimethylamine
 $(\text{CH}_3)_3\text{N}$ trimethylamine

Aromatic amines

$\text{C}_6\text{H}_5\text{NH}_2$ phenylamine (*aniline*)

Q.1 Draw structures for all amines of molecular formula $\text{C}_4\text{H}_{11}\text{N}$.
 Classify them as primary, secondary or tertiary amines.

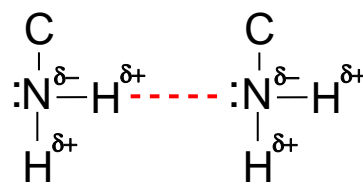
Properties The presence of the lone pair in 1° , 2° and 3° amines makes them ...

- **Lewis bases** - they can be lone pair donors
- **Brønsted-Lowry bases** - can be proton acceptors
- **Nucleophiles** - provide a lone pair to attack a positive (electron deficient) centre

Physical properties

Boiling point • Boiling points increase with molecular mass.

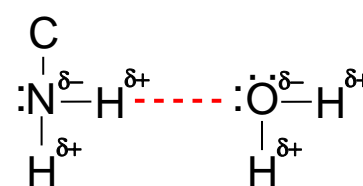
- Amines have higher boiling points than corresponding alkanes because of intermolecular hydrogen bonding.
- Quarternary ammonium salts are **ionic** - they exist as crystalline salts.



intermolecular hydrogen bonding in amines

Solubility • Soluble in organic solvents.

- Lower mass compounds are soluble in water due to hydrogen bonding with the solvent.
- Solubility decreases as molecules get heavier.



hydrogen bonding between amines and water

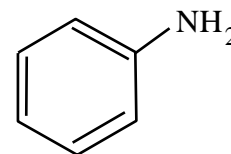
Basic properties

Bases The **lone pair** on nitrogen makes amines **basic**. $\text{RNH}_2 + \text{H}^+ \rightarrow \text{RNH}_3^+$

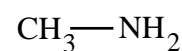
Strength

- depends on the availability of the lone pair and thus its ability to pick up protons
- the greater the electron density on the N, the better its ability to pick up protons
- this is affected by the groups attached to the nitrogen.

- electron withdrawing substituents (e.g. benzene rings) decrease basicity as the electron density on N is lowered.



- electron releasing substituents (e.g. CH_3 groups) increase basicity as the electron density is increased



draw arrows to show the electron density movement

Measurement • the strength of a weak base is depicted by its K_b or $\text{p}K_b$ value

pK and *pK_b*

values

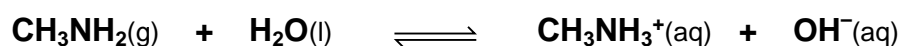
- the larger the K_b value the stronger the base
- the smaller the pK_b value the stronger the base.
- the pK_a value can also be used - ($pK_a + pK_b = 14$)
- the smaller the pK_b , the larger the pK_a .

<i>Compound</i>	<i>Formula</i>	<i>pK_b</i>	<i>Comments</i>
ammonia	NH ₃	4.76	
methylamine	CH ₃ NH ₂	3.36	methyl group is electron releasing
phenylamine	C ₆ H ₅ NH ₂	9.38	electrons delocalised into the ring

strongest base **methylamine > ammonia > phenylamine** *weakest base*

Reactions

- Amines which dissolve in water **produce weak alkaline solutions**

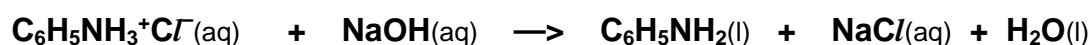


- Amines **react with acids to produce salts.**



This reaction allows one to dissolve an amine in water as its salt.

Addition of aqueous sodium hydroxide liberates the free base from its salt

**Nucleophilic Character**

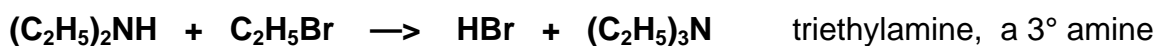
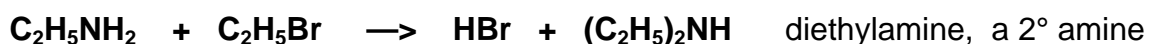
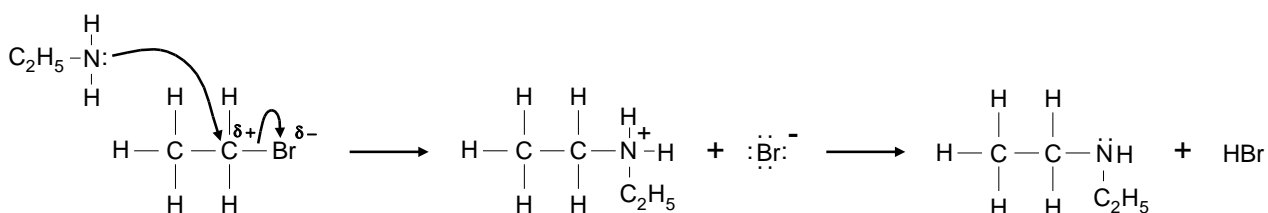
Due to their **lone pair**, amines react as **nucleophiles** with

- haloalkanes *forming substituted amines* **nucleophilic substitution**
- acyl chlorides *forming N-substituted amides* **addition-elimination**

Haloalkanes Amines can be prepared from haloalkanes (see below and previous notes).

<i>Reagent</i>	Aqueous, alcoholic ammonia
<i>Conditions</i>	Reflux in aqueous, alcoholic solution under pressure
<i>Product</i>	Amine (or its salt due to a reaction with the acid produced)
<i>Nucleophile</i>	Ammonia (NH ₃)
<i>Equation</i>	$\text{C}_2\text{H}_5\text{Br} + \text{NH}_3(\text{aq / alc}) \longrightarrow \text{C}_2\text{H}_5\text{NH}_2 + \text{HBr} \quad (\text{or } \text{C}_2\text{H}_5\text{NH}_3^+\text{Br}^-)$

Problem The amine produced is also a nucleophile and can attack another molecule of haloalkane to produce a secondary amine. This in turn can react further producing a tertiary amine and, eventually an ionic quarternary amine.



Uses Quarternary ammonium salts with long chain alkyl groups

eg $[\text{CH}_3(\text{CH}_2)_{17}]_2\text{N}^+(\text{CH}_3)_2\text{Cl}^-$ are used as cationic surfactants in fabric softening.

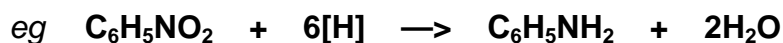
Prepared from

haloalkanes Nucleophilic substitution using ammonia ... see above

nitriles **Reduction** of nitriles using $\text{Li}^+\text{AlH}_4^-$ in **dry ether**



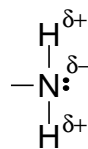
nitro compounds **Reduction** by refluxing with **tin and conc. hydrochloric acid**



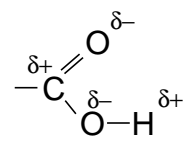
α - AMINO ACIDS

Structure Amino acids contain 2 functional groups

- **amine** NH_2
- **carboxyl** COOH

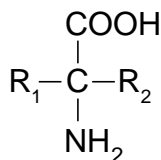


Amine

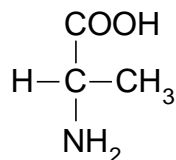


Carboxyl

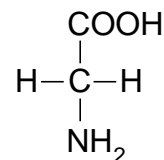
They all have a similar structure - the identity of R_1 and R_2 vary



general structure



2-aminopropanoic acid (Alanine)



2-aminoethanoic acid (Glycine)

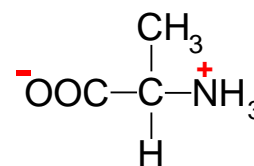
Optical Isomerism

Amino acids can exist as **optical isomers** if they have **different R_1 and R_2** groups

- optical isomers exist when a molecule contains an **asymmetric carbon** atom
- asymmetric carbon atoms have **four different atoms or groups** attached
- **two isomers** are formed
- one **rotates plane polarised light** to the left, one rotates it to the right
- **no optical isomerism with glycine** - two H's are attached to the C atom

Zwitterions

- a zwitterion is a **dipolar ion**
- it has a **plus and a minus charge** in its structure
- a proton from the COOH group moves to NH_2
- amino acids exist as **zwitterions** at a certain pH
- the pH value is called the **isoelectric point**
- produces increased inter-molecular forces
- melting and boiling points are higher

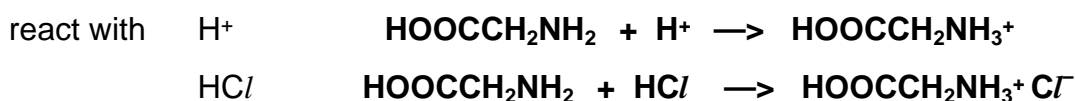


a zwitterion

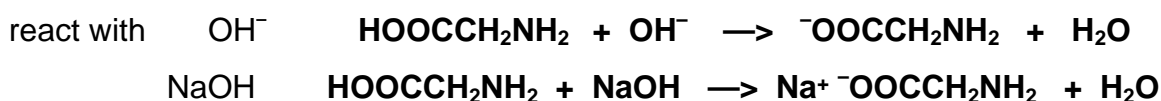
Acid/base properties

- amino acids possess acidic and basic properties due to their functional groups
- they will form salts when treated with acids or alkalis.

Basic properties:



Acidic properties:

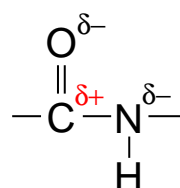


Q.2 Describe the arrangement of bonds in the amino acid H_2NCH_2COOH around...
 the N atom in the NH_2
 the C atom in the $COOH$
 the C atom in the CH_2

What change, if any, takes place to the arrangement around the N if the amino acid is treated with dilute acid?

PEPTIDES

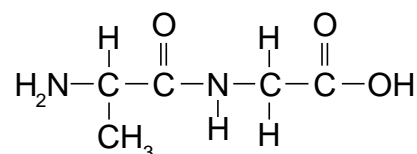
Formation • α -amino acids can join up together to form peptides via an **amide** or **peptide link**



the peptide link

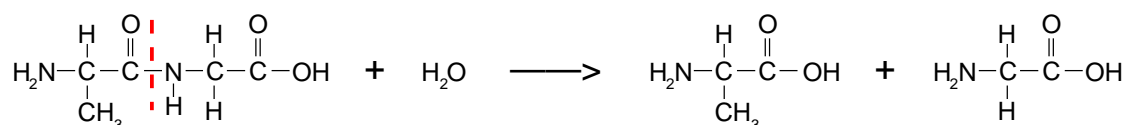
Structure Sequences of amino acids joined together by peptide links

- 2 amino acids joined **dipeptide**
- 3 amino acids joined **tripeptide**
- many amino acids joined **polypeptide**



a dipeptide

Hydrolysis Peptides can be broken down into their constituent amino acids by **hydrolysis**



- attack takes place at the slightly positive C of the $C=O$
- the C-N bond next to the $C=O$ is broken
- **hydrolysis with just water is not feasible**
- **hydrolysis in alkaline/acid conditions is quicker**
- hydrolysis in acid/alkaline conditions (e.g. NaOH) will produce salts

with HCl	NH_2	will become	$NH_3^+Cl^-$
H^+	NH_2	will become	NH_3^+
NaOH	$COOH$	will become	$COO^- Na^+$
OH^-	$COOH$	will become	COO^-

Q.3 Draw structural isomers for the compounds produced when

- $H_2NCH_2CONHCH(CH_3)COOH$ is hydrolysed by water
- $H_2NCH_2CONHC(CH_3)_2COOH$ is hydrolysed in **acidic** solution
- $H_2NCH_2CONHCH(CH_3)COOH$ is hydrolysed in **alkaline** solution

Q.4 Write out possible sequences for the **original** peptide if the hydrolysis products are

- 1 mole of amino acid **A**, 1 mole of amino acid **B** and 1 mole of amino acid **C**
- 1 mole of amino acid **A**, 2 moles of amino acid **B** and 1 mole of amino acid **C**
- 1 mole of amino acid **A**, 1 mole of **B**, 1 mole of **C**, 1 mole of **D** and 1 mole of **E**

Proteins

- **polypeptides with high molecular masses**
- chains can be lined up with each other
- the **C=O** and **N-H** bonds are **polar** due to a difference in electronegativity
- **hydrogen bonding** exists between chains

dotted lines -----
represent hydrogen bonding

