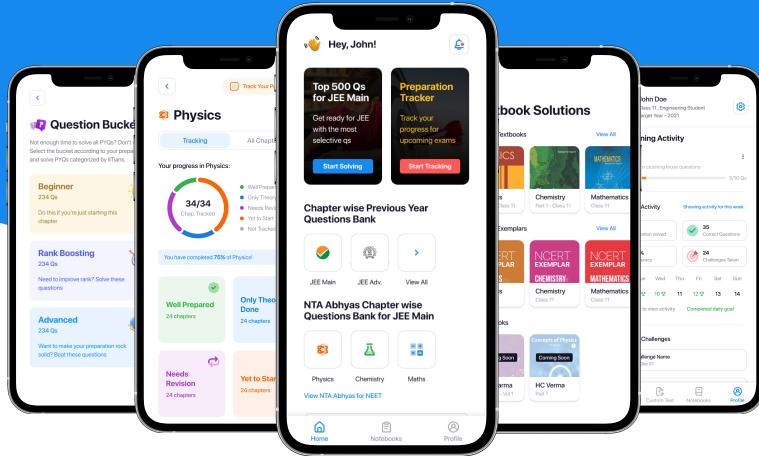




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Biomolecules

The hydrates of carbon was originally defined as carbohydrates, with general formula of $C_n(H_2O)_y$. But this definition has found several limitations today.

- (i) Carbon do not forms hydrates
- (ii) There are number of organic molecules having $C_n(H_2O)_y$ formula but are not carbohydrates.
 - e.g. (a) Formaldehyde ($HCHO$) : CH_2O
 - (b) Acetic acid CH_3COOH : $C_2H_4O_2$ etc.
- (iii) A number of carbohydrates do not have $C_n(H_2O)$ formula.
 - e.g. (i) Rhamnose - $C_6H_{12}O_5$
 - (ii) Deoxyribose - $C_5H_{10}O_4$

MODERN DEFINITION :

Polyhydroxy aldehyde or Ketone or substances which gives these on hydrolysis is called carbohydrates. They have at least one chiral carbon in general

CLASSIFICATION :

TYPE : 1

Carbohydrates		
(1) Monosaccharides	(2) Oligosaccharides	(3) Polysaccharides
Simplest carbohydrates which can't be hydrolysed into simpler molecules Gen. formula = $(CH_2O)_n$ $n = 3$ to 7.	* oligo = few Carbohydrates which on hydrolysis gives 2 to 10 molecules of carbohydrates	Carbohydrates which on hydrolysis gives number of carbohydrates. e.g. $(C_6H_{10}O_5)_n$ $n = 100$ to 3000

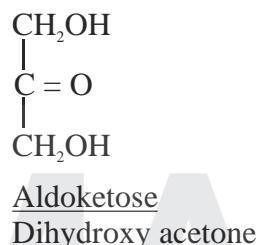
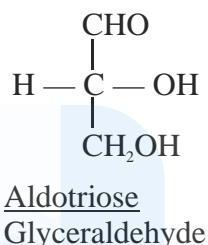
TYPE : 2

Carbohydrates	
(i) Sugar	(ii) Non-Sugar
All the Mono and Oligosaccharides have sweet taste and these are collectively called sugars. They are crystalline in nature.	Polysaccharides are tasteless and are called non-sugar. They are amorphous in nature.

TYPE : 3

Carbohydrates	
(i) Reducing	(ii) Non-Reducing
<p>All those carbohydrates which has aldehydic or ketonic group in their hemiacetal and hemiketal structures have the ability to reduce Tollen's Reagent or Fehling solution. They are called Reducing sugar.</p> <p>* All Monosaccharides whether aldose or ketose are reducing sugars.</p>	<p>The carbohydrates which can not reduce Tollen's reagent or Fehling solutions are called as Non-reducing.</p> <p>* All Polysaccharides are non reducing. e.g. Starch, Cellulose, dextrins, Glycogens etc.</p>

Simplest Triose : The simplest monosaccharides are triose.

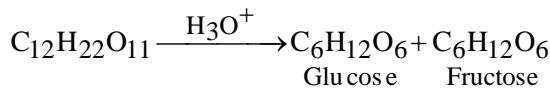


GLUCOSE

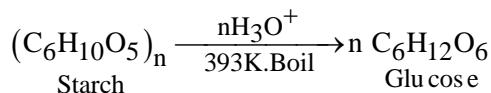
- * They are also called as Grape sugar or Dextrose. It is present in sweet fruits.
- Molecular formula — $\text{C}_6\text{H}_{12}\text{O}_6$
It is aldohexose sugar

PREPARATION OF GLUCOSE :

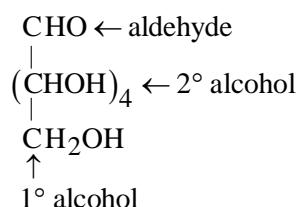
(i) From Sucrose (Cane - Sugar) : By acidic hydrolysis.



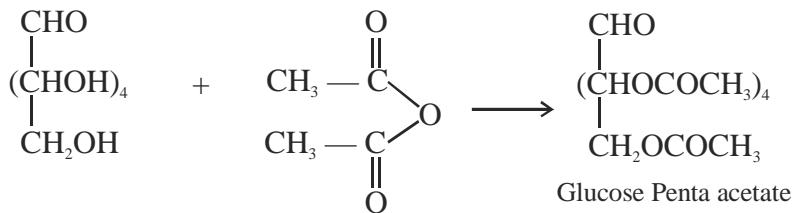
(ii) From Starch : Commercial method.



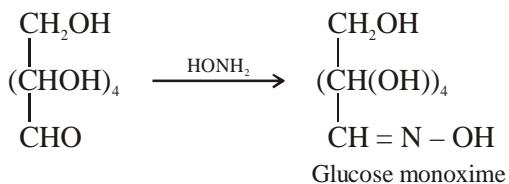
CHEMICAL REACTION OF GLUCOSE :



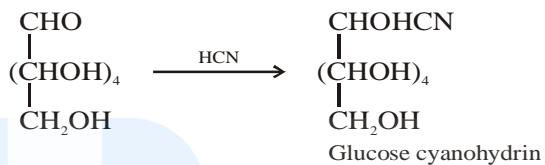
1. Reaction with Acetic anhydride : Acetylation



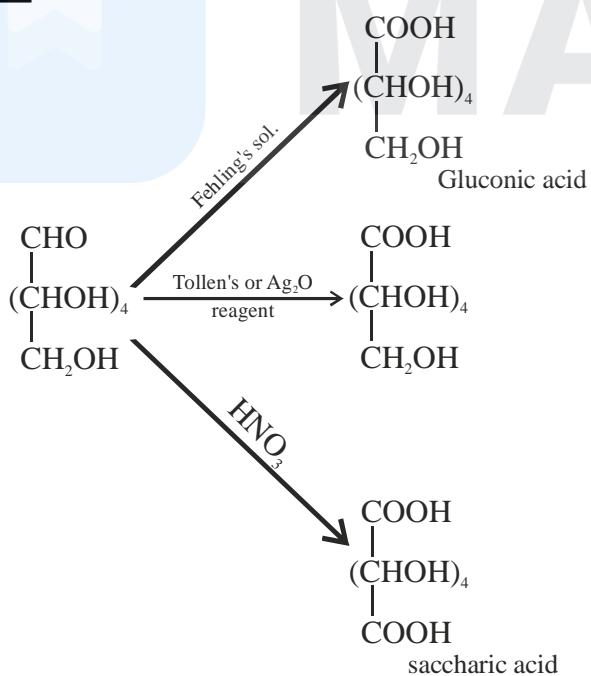
2. Reaction with hydrosyamines :



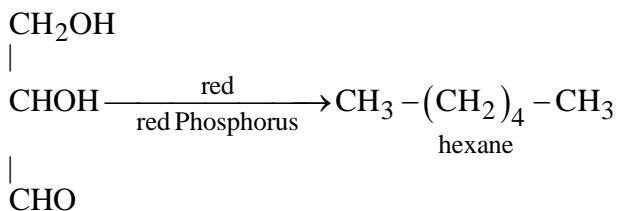
3. Reaction with HCN :



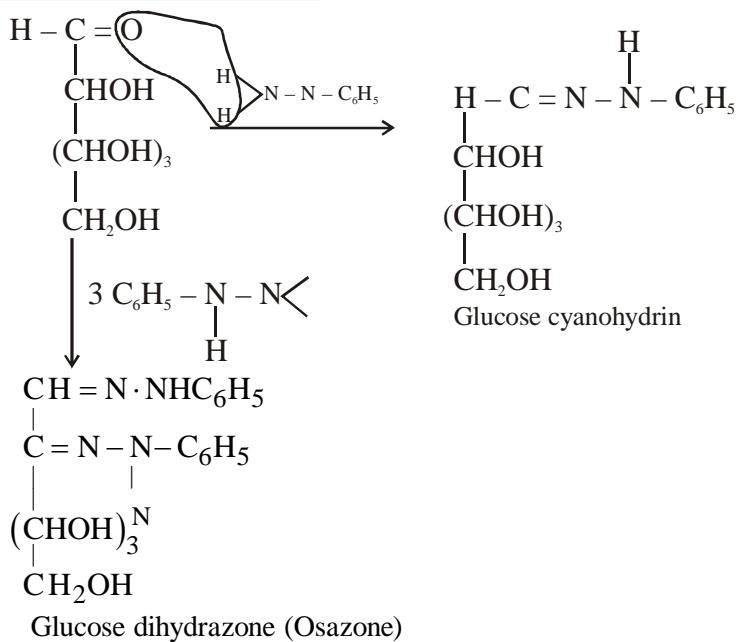
4. Oxidation :



5. Reduction :



6. Reaction with Phenyl hydrazine :



7. Action of Alkali : Lobry de Bruyn-van Ekenstein reaction

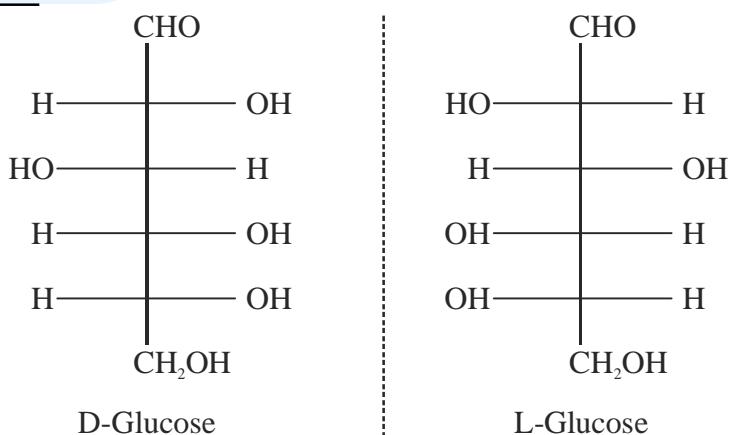


On adding concentrated solution of NaOH this occurs.

⇒ It is probably on account of this isomerisation, that fructose reduces Fehling's solution and Tollen's Reagent in alkaline medium.

STRUCTURE OF GLUCOSE

Open-chain strs :



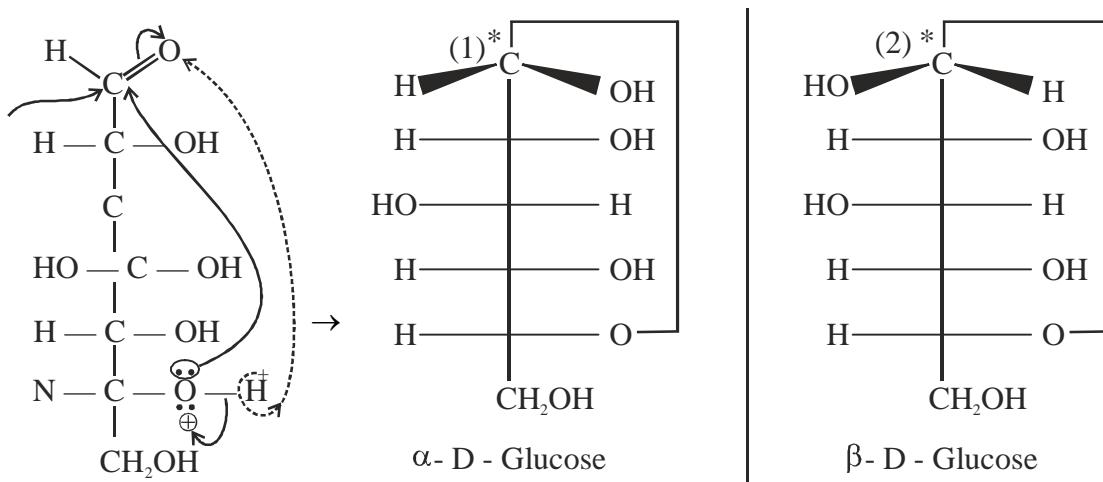
Limitation of open-chain strs :

- (i) Glucose doesn't undergo characteristic reaction of aldehydes such as
 - (a) Glucose does not react with NaHSO₃ (sodium bisulphite)
 - (b) Glucose does not respond to Schiff's test.
- (ii) Glucose doesn't react with G.R.

Cyclic structures of Glucose :

α – D and β – D Glucose : (Hemiacetal structures)

- In α -D-Glucose OH group is towards right while in β -D-Glucose the OH group is towards left at C₁.



ANOMERS :

“The stereo isomers which differs in configuration at “C₁” is called anomers”.

The “C₁” is also called as anomeric carbon or Glycosidic carbon.

Note : ‘ α ’ and ‘ β - D’ Glucose are not enantiomers, since the configuration at other carbon remains same.

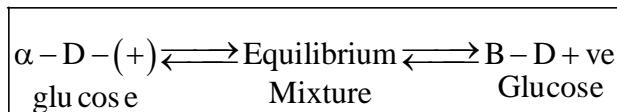
Mutarotation :

When Glucose is crystallized from a concentrated solution at 300 K, α form of D-Glucose is isolated. But, from a hot saturated solution (aqueous) at a temperature in excess of 100°C, the β -Glucose is obtained.

Term	α	β
M.pt :	146°C	150°C
Sp. rotation :	+111°	+19.2°

If either of the two forms is dissolved in water and allowed to stand, the specific rotation of the solution slowly changes and reaches a constant value of +52.5°.

“The spontaneous change in specific rotation of an optically active compound is called mutarotation”.



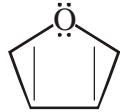
Haworth structures of Glucose : The molecules of glucose and fructose can exist in any of the two cyclic structures.

(a) Pyranose structure
Six membered ring
Derived from pyran

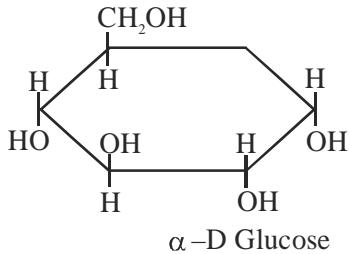
(b) Furanose structure
5 Membered Ring
Derived from furan



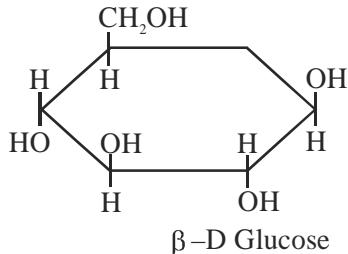
Pyran



Furan



α -D Glucose



β -D Glucose

OLIGOSACCHARIDES

These are carbohydrates which gives 2 to 10 molecules of monosaccharides.

* In some standard text (particularly in Biology) $n = 2$ to 7 has been frequently used.

Example

- (1) Disaccharides : Sucrose, Maltose, Lactose
(2) Trisaccharides : Raffinose
(3) Tetrasaccharides : Stachyrose

Disaccharides :

Gives two molecules of same or different monosaccharides.

General formula :

Formula : C₁₂ H₂₂ O₁₁

Examples and its monomers :

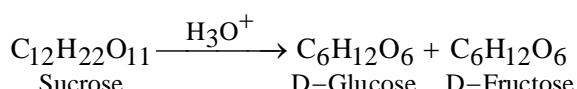
- (1) Sucrose $\xrightarrow[\text{H}_3\text{O}^+]{\text{Invertase}}$ Glucose + Fructose

(2) Maltose $\xrightarrow[\text{Maltase}]{\text{H}_3\text{O}^+}$ Glucose + Glucose

(3) Lactose $\xrightarrow[\text{Lactase}]{\text{H}_3\text{O}^+}$ Glucose + Galactose

- The disaccharides can be reducing or non reducing. If carbonyl group is free, sugar is reducing.

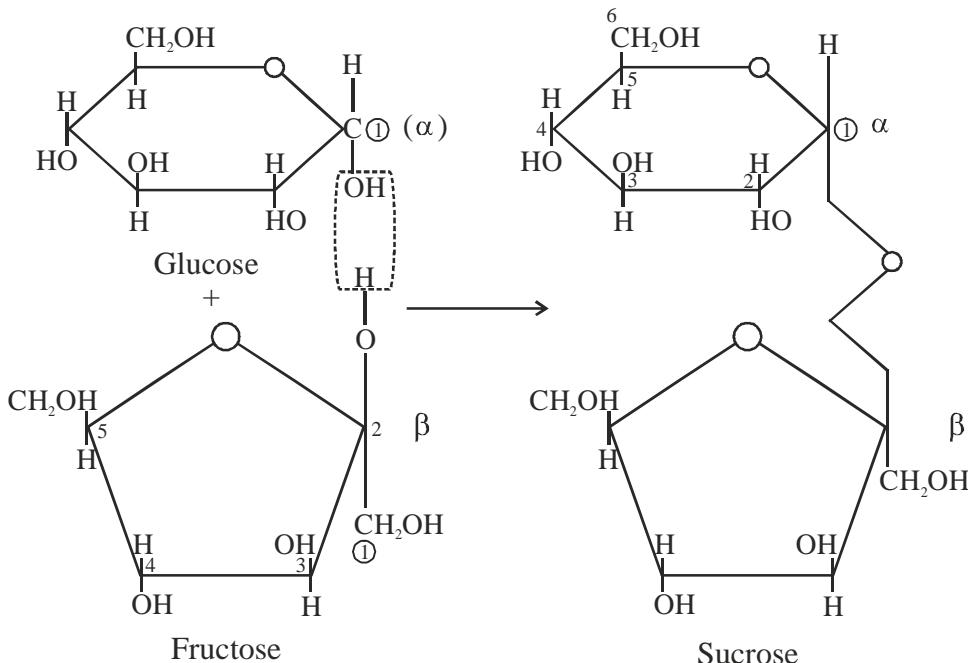
Sucrose or cane sugar or Table Sugar :



Specific rotation + 66.5° +52.5° -92.4°

Since the laevo-rotation of fructose is more than the dextro-rotation of glucose, the resulting solution will be laevo-rotatory. This overall process is called inversion of sugar.

HAWORTH STRUCTURE OF SUCROSE :



- Determination of ring size by spectral methods has revealed that, in sucrose glucose is in its pyranose form and fructose it is furanose form.
- There is thus, α,β glucosidic bond between glucose and fructose as monomers.

POLYSACCHARIDES :

Polysaccharides are formed when large number of monosaccharide joins together with simultaneous elimination of water molecules.

Some common polysaccharides are :

- (i) Cellulose (ii) Starch (iii) Glycogen (iv) Dextrin

STARCH :

(i) It is also called as Amylum.

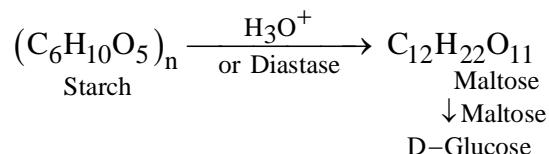
General formula - $(C_6H_{10}O_5)_n$

where, $n \approx 100$ to 300

It is the chief food storage material in plants.

(ii) Starch is white amorphous powder which is sparingly soluble in water.

(iii) Hydrolysis of starch :



(iv) Starch does not reduce Fehling solution or Tollen's reagent and does not form osazone. This clearly suggests that all hemiacetal hydroxy group of glucose unit at C_1 is involved in glycosidic linkage.

(v) Starch is a mixture of two poly saccharides Amylose and Amylopectins.

Natural starch has

Amylose : 10 to 20%

Amylopectin : 80 to 90%

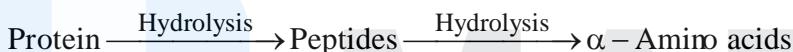
CELLULOSE :

- (i) It is the major constituent material of plant cell wall. In general, wood contains 50% cellulose and cotton contains 90-95% cellulose.
- (ii) It is a colourless, amorphous solid which can be decomposed by heating.
- (iii) It is almost linear and its individual monomeric units are joined through multiple H-bond.
- (iv) Cellulose does not reduce Fehling solution, Tollen's reagent and does not form osazone.
- (v) Large population of cellulolytic bacteria is present in the stomach of ruminant mammals. These bacteria cause decomposition of cellulose in stomach cellulose is digested as glucose in ruminant mammals.
- (vi) Human does not have any system to digest cellulose.
- (vii) Structure of Cellulose : Cellulose is a straight chain polymer of D-glucose which are joined by β -Glucosidic bond b/w C₁ of one glucose and C₄ of other glucose.

PROTEINS

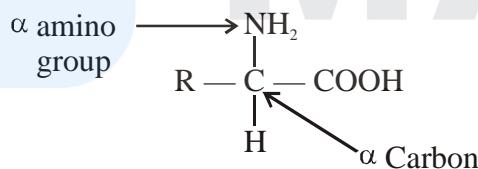
Proteins are vital chemical substance essential for the growth and maintenance of life. Chemically proteins are condensation polymers in which the monomeric unit is α amino acids. All proteins contain the elements like carbon, hydrogen, oxygen, nitrogen and sulphur in major.

HYDROLYSIS OF PROTEINS



α Amino acids :

α amino acids are the building block of proteins.



The total of 20 amino acids has been isolated by hydrolysis of various proteins.

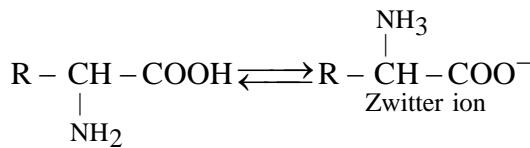
Classification of α Amino acids :

Amino acids	
Essential	Non-essential
The amino acids which can't be synthesised in human body is called essential amino acids * They are 10.	The amino acids which can be synthesised in human body is called non-essential amino acids. * They are also 10.

α - Amino acids		
(a) Neutral	(b) Alkaline	(c) Acidic
Has one NH_2 group and one COOH group.	Two NH_2 group and one COOH group	Has two COOH group and one NH_2 group

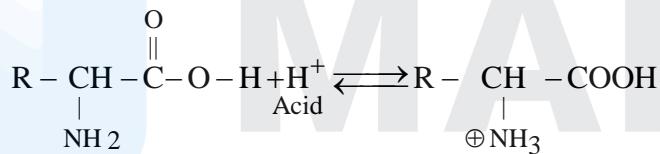
Zwitterion ion structure :

α - amino acid largely exists as dipolar ion.



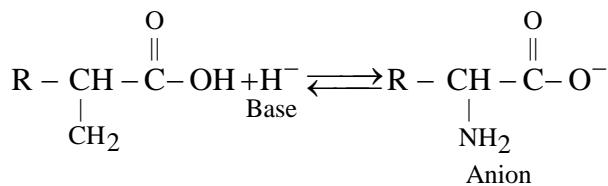
Electrical behaviour of Amino acids :

- (a) In acidic solution : α amino acid exist as cations and thus will migrat towards the cathode under electrical field.



The cation will migrat towards cathode.

- (b) In alkaline solution : α Amino acid exist as



The anion will migrat towards +ve electrode i.e. Anode.

Iso electric point :

The pH at which amino acid has no net migration towards any of the electrode under influence of electric field is called isoelectric point.

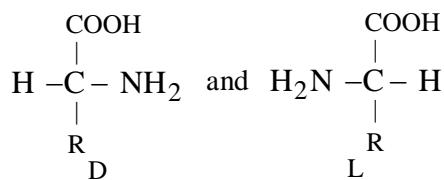
Each amino acid has a characteristic isoelectric point.

Amino acid	Iso electric point (pH)
(1) Neutral	Slightly less than 7 (≈ 6.1)
(2) Acidic	3.2 to 3.5
(3) Alkaline	7.6 to 10.8

At isoelectric point amino acids have least solubility.

D, L Nomenclature of Amino acids :

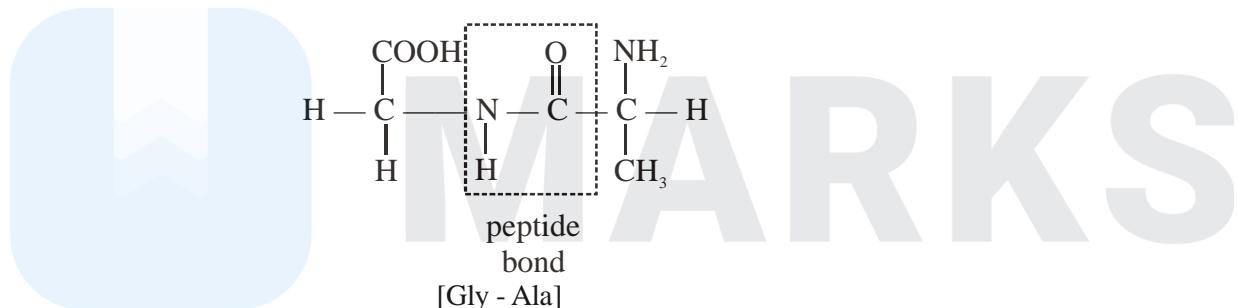
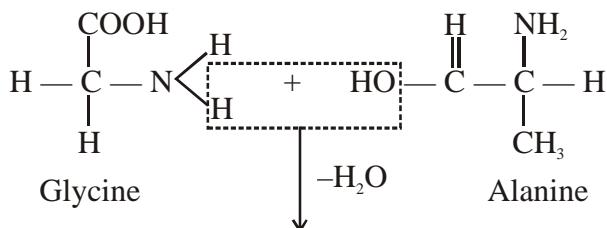
All amino acids except glycine is chiral and has two stereo isomeric forms.



All naturally occurring amino acid belongs to L series.

Peptide bond :

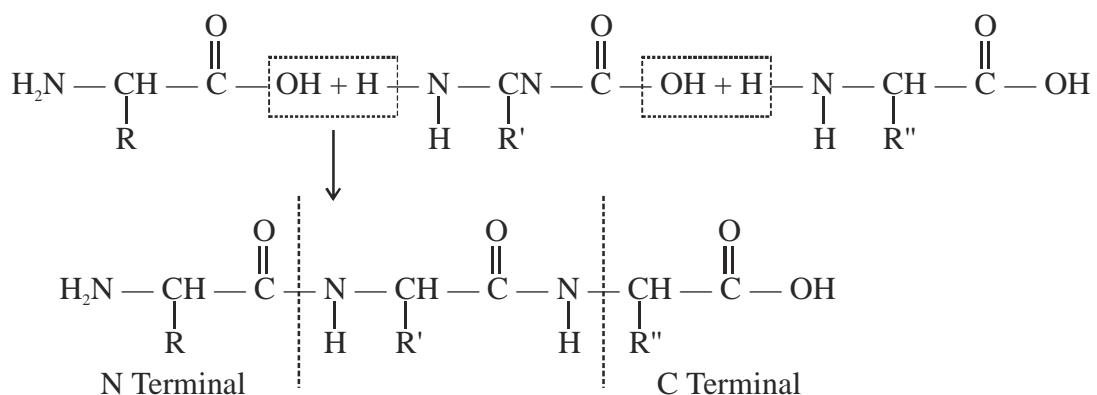
Peptides are organic amides formed by the condensation of amino group of one α amino acid and carboxylic acid of other amino acid, by simultaneous elimination of water.



* Thus $\overset{\text{O}}{\parallel}\text{C}-\text{N}-$ linkage is called the peptide bond.

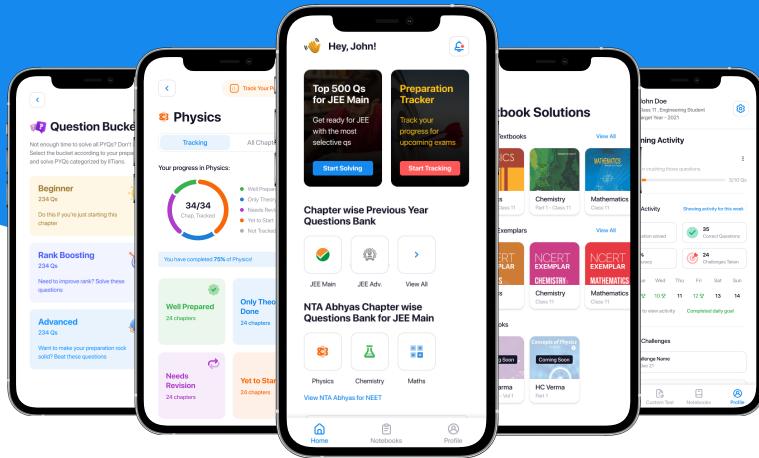
Poly peptides :

If large number of α amino acids are joined together by peptide bonds, the polyamide is formed. Such polyamides are called polypeptides.





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