Bíomolecules



MONOSACCHARIDES Chemical Reactions of Glucose:

$$C_{12}H_{22}O_{11} + H_{20} \xrightarrow{H^+} Glucose + Fructose$$

 $C_6H_{12}O_6 \quad C_6H_{12}O_6$

1. Reaction with acetyl chloride (CH_3CO_{Cl}):

When glucose reacts with acetyl chloride in the presence of <u>ZnCl2</u>, it forms glucose pentaacetate and HCl. This reaction demonstrates the presence of hydroxyl groups in glucose.

CHOOCCH3) + HCl. CH2ODCCH3

Treatment with PCl5:

Glucose reacts with phosphorus pentachloride to form pentachloroglucose (CHCl)_& along with POCl₃ and HCl. This further confirms the presence of hydroxyl groups.



(CHCL) & along with PUCL3 and HCL. Inis further confirms the presence of hydroxyl groups.



Reduction reaction: When glucose is treated with sodium amalgam and water $(Na-Hg/H_2O)$, it forms sorbitol $(CH_2OH)(CHOH)_4CH_2OH)$. This reaction shows that glucose can be reduced at its aldehyde group.

Oxídatíon reactíons:

CH2OH (CHOH) COOH o With bromine water: Forms gluconic acid

• With nitric acid: Produces saccharic acid with additional oxidation

O The formation of these acids proves the presence of both aldehyde and primary alcohol groups



Formation of glucosazone: Glucose reacts with excess phenylhydrazine (C6H5NHNH2) to form glucosazone, which has characteristic yellow needleshaped crystals. This is a crucial identification test.

$$CH = N NHC_{6}H_{5}$$

$$C = N NHC_{6}H_{5}$$

$$(CHOH)_{3}$$

$$CH_{2}OH$$

Structural Insights: The straight-chain structure of glucose explains most properties but fails to explain three important observations:

- No reaction with 2,4-DNP and NaHSO3
- No reaction of its pentaacetate with NH2OH
- \bullet Existence of α and β forms



These observations are better explained by glucose's cyclic structure, which exists in equilibrium with the open chain form. The cyclic structure can take two forms:
Furanose form: Five-membered ring containing oxygen





• Pyranose form: Six-membered ring containing oxygen



Mutarotation: A particularly important phenomenon is mutarotation, where α -D-(+)-glucose converts to β -D-(+)-glucose in aqueous solution, reaching an equilibrium. The specific rotation changes from +112° to +19° at equilibrium (+ 52.7°).

Anomers:

stereoisomers that differ in configuration only around C-1 (the anomeric or glycosidic carbon). This explains the existence of α and β forms of glucose



Sucrose, C₁₂H₂₂O₁₁ (Cane sugar, non-reducing) :





• Lactose (C₁₂H₂₂O₁₁) (Milk sugar, reducing) :



Polysaccharides:

Starch is one of the most significant polysaccharides, serving as the primary energy source for humans.

key characterístics and behaviour: The structure of starch consists of two main components: anylose and anyl<u>opectin</u>.

Amylose forms a straight chain with α -1,4-glycosidic bonds between glucose units. **Amylopectin**, however, is more complex, having both α -1,4-glycosidic bonds in the main chain and α -1,6-glycosidic bonds at branching points. When we eat starchy foods, our body begins breaking them down immediately.

In our saliva, the enzyme amylase initiates this process:

 $Starch \rightarrow Dextrín \rightarrow Maltose \rightarrow Glucose$

 $(\mathcal{C}_{6} \vdash_{10} \mathcal{O}_{5})_{n} \rightarrow (\mathcal{C}_{6} \vdash_{10} \mathcal{O}_{5})_{x} \rightarrow \mathcal{C}_{12} \vdash_{22} \mathcal{O}_{11} \rightarrow \mathcal{C}_{6} \vdash_{12} \mathcal{O}_{6}$

A distinctive property of starch is its reaction with iodine, producing a characteristic blue color. This serves as a key identification test. Importantly, starch doesn't reduce Fehling's or Tollens' reagents because its reducing groups are involved in glycosidic bonds.

Cellulose:

Another crucial polysaccharide, differs significantly from starch. It forms the structural framework of plant cell walls and is remarkably abundant - making up 90-95% of cotton and 45-50% of wood.

Its structure involves β -D-glucose units connected by β -1,4-glycosidic linkages between C-1 of one glucose unit and C-4 of the next.



Glycogen, often called "animal starch," serves as the storage form of glucose in animals. It's primarily found in the liver, muscles, and brain, with a structure similar to amylopectin but more extensively branched.

Proteins and Amino Acids:

Proteíns are complex bíomolecules con	nposed of amino acids linked through pe f	stide
bonds (-CONH-).	COOHYHAN COHN -	
understanding amino acids is crucil	al for comprehending protein structure au	rd
function:		
Amíno acíds share a common structu	ure but díffer ín their síde chains. They c	an
be classified into several categories:	Except alucine	
1. Non-polar síde chaíns:	- uper ne	
• Símple ones líke glycíne (no síde cl	hain) and alanine (methyl group)	
	NH2	
HOOC-CH2-NH2	CH3 CH	
	COOH	
Gtly cine.	Alarine	
1		

• Branched ones like valine, leucine, and isoleucine (all essential)

• Special cases like proline, which forms a unique ring structure



2) Polar but neutral side chains:

- Hydroxyl-containing (serine, threonine)
- Sulfur-containing (cysteine, methionine)
- Aromatic (tryptophan, tyrosine)
- Amide-containing (asparagine, glutamine) (_X)

Amíno acíds with charged síde chains:

- Basic (lysine, arginine, histidine)
- Acídic (aspartic acíd, glutamic acíd)

Amphoterísm:

A fascinating property of amino acids is their amphoteric nature - they can act as both acids and bases. This leads to the formation of **zwitterions**, where the amino acid carries both positive and negative charges. At a specific pH called the isoelectric point, the amino acid exists as a neutral zwitterion, showing minimum solubility.

Proteins can be classified based on their shape and structure:

- 1. Fíbrous proteíns: These have parallel polypeptide chains held together by hydrogen and disulfide bonds, making them insoluble in water. Examples include keratin and myosin.
- 2. Globular proteíns: These form three-dímensíonal spherical structures and are typically soluble in water. Insulin and albumins are prime examples.

Proteín Structure and Its Levels:



Proteín Structure and Its Levels:



The structure of proteins can be understood through four hierarchical levels, each building upon the previous one:

Primary Structure: This is the most basic level - imagine it as the raw sequence of amino acids linked together by peptide bonds. Each peptide bond forms between the carboxyl group (-COOH) of one amino acid and the amino group (-NH₂) of another, creating a backbone with side chains (R_1 , R_2 , R_3 , etc.) hanging off like branches from a tree.



Secondary Structure: This level shows how the protein chain begins to fold into regular patterns:

 α-Helíx: Pícture a spíral staírcase, where the proteín chain coils around itself líke a spring. Hydrogen bonds between every fourth amino acid stabilize this structure, making proteíns líke keratín in hair incredibly strong yet flexible.



1. β -Pleated Sheet: Imagine a folded paper fan - the protein chain folds back and forth upon itself, creating a flat sheet structure. These sheets are particularly common in silk proteins, giving them their unique strength.



Tertiary Structure: Think of this as the final <u>3D shape</u> of a single protein chain. Various forces come into play here - hydrophobic interactions push some parts inward, while polar regions stay on the surface. This level gives proteins like hemoglobin their specific functional shape.

Quaternary Structure: This is like a protein team - multiple folded protein chains (called subunits) come together to form one functional unit. Hemoglobin, with its four subunits, is a classic example.

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Enzymes and Their Properties:

Enzymes are nature's catalysts, and their behavior follows several key principles: 1. Mechanism of Action:

The process follows a clear sequence:

- Enzyme (E) meets substrate (S) \rightarrow Forms ES complex
- ES complex transforms \rightarrow EI (intermediate) complex \checkmark
- El complex changes \rightarrow EP (product) complex
- Finally, EP breaks apart \rightarrow Free enzyme + Product

2. Key Properties:

- High Efficiency: They can catalyze millions of reactions per second
- Specificity: Like a lock and key, each enzyme works with specific substrates
- Temperature Sensitive: Most work best around body temperature (37°C)
- pH Dependent: Each has an optimal pH range (often around 7)

Vítamíns

These are organic compounds which cannot be produced by the body and must be supplied in small amounts in diet for normal health, growth and maintenance of body. Multiple deficiencies caused by lack of more than one vítamín are common ín human beings. This condition of vitamín deficiency is known as avitaminosis. * The vitamins which are soluble in water are called water soluble vitamins. e.g. vit. B, vit. C, etc. * The vitamins which are soluble i.n. fate are called fat fat Boluble. * The vitamins which are soluble in fats are called fat soluble vítamíns. e.g. vít. A, D, E, K.

"ase"

Nucleic acids are polynucleotides present in the nuclei of all living cells in the form of nucleoproteins. They are the fundamental molecules of heredity and contain genetic information essential for all living organisms. Basic Components (High Priority Topic)

- Each nucleotide consists of three essential components:
- Pentose Sugar (5-carbon sugar)
- Nítrogenous Base
- Phosphate Group (H3PO4)



	Feature	DNA (X.)	RNA (X)		
<u></u>	Sugar	2-deoxy-D-ríbose	D-ríbose	/	\nearrow
	Pyrimidine bases	Cytosíne, Thymíne	uracíl, Cytosíne		
	Puríne bases	Adeníne, Guaníne	Adeníne, Guaníne		
	Structure	Double-stranded helíx	Síngle-stranded		Х
	Location	Nucleus (mainly)	Nucleus + Cytoplasm		$(\)$
	Replication	Self-replicating 🗸	Non-self-replicating	-/	$\langle \rangle$
	Síze	Larger	Smaller		Х
	Stabílíty	More stable	Less stable		/

Chargaff's Rules (Important for Both Exams) First Rule: A = T, G = C (in DNA) Second Rule: Purine = Pyrimidine Base Pairing: A=T (2 H-bonds) $G\equiv C$ (3 H-bonds) In RNA: A=U

Bíological Functions (Essential for NEET)

Replication

- Semíconservative process
- Produces identical DNA copies
- Enzyme-catalyzed

Proteín Synthesis

- Transcription: DNA \rightarrow RNA
- Translation: $RNA \rightarrow Protein$
- Genetic code: Triplet codon system

Mutations (Important for NEET)

- Definition: Changes in nitrogenous base sequence
- Effects:
- Altered amíno acíd sequence
- Modífied proteín structure
- Possíble genetic disorders

Important Practice Problems JEE-Level Problem:

3. Calculate the number of hydrogen bonds in a DNA segment containing 30% adenine. Total number of nucleotides = 100.

$$A = T = 30 f.$$

 $G1 = C = 20 f$

$$No. of H-bonds = (A-T paulis x 2) + = (30 \times 2) + (G1-C pauli x 3) = (20.)$$

NEET-Level Problem:

1. If a DNA segment has 20% cytosine, predict the percentage of other bases.

$$C = G_1 = 20 \gamma$$
.
 $A = T = 30 \gamma$. = 100 γ

Hormones:

These chemical messengers can be classified into three main types: 1. Steroid Hormones:

- Examples: testosterone, estrogen, cortísone
- Fat-soluble molecules that easily cross cell membranes
- Control development and metabolism
- 2. Peptide Hormones:
- Examples: insulin, oxytocin
- Water-soluble proteins that work through cell surface receptors
- Control various metabolic processes
- з. Amíne Hormones:
- Examples: thyroxine, epinephrine
- Derived from amino acids
- Control metabolism and stress responses

JEE Maín 2023: Q:

A polypeptide has 8 amino acids. Calculate the total number of peptide bonds present.

"No. of Peptide bonds is always one less
than
$$AA$$
." $B-1=7$ Peptide bond.

NEET 2022: Q: Which of the following statements about enzymes is incorrect? a) They are highly specific b) They possess both active and allosteric sites c) They are predominantly proteins de they can be used indefinitely • JEE Advanced 2023:

Q: Match the following:

	Column A	Column B	
Ļ	finsulín p.	Peptíde hormone	A
ł	Thyroxíne	q. Steroíd hormone	C
Č	Estrogen	r. Amíne hormone	B

NEET 2023: Q: Which statement about nucleic acids is correct?

- a) RNA contains thymine > Uracid. b) DNA is single-stranded > double Storanded.
- c) In DNA, adenine pairs with thymine through two hydrogen bonds
- d) RNA contains uracil and ribose sugar

JEE Main 2022: Q: Calculate the pH of isoelectric point of glycine if $pKa_1 = 2.34$ and $pKa_2 = 9.60$

$$pI = \left(\frac{P_{ka_1} + P_{ka_2}}{2}\right)$$

= $\frac{2 \cdot 39 + 9 \cdot 60}{2} = \frac{11 \cdot 94}{2} = 5.97$

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