

BIOMOLECULES

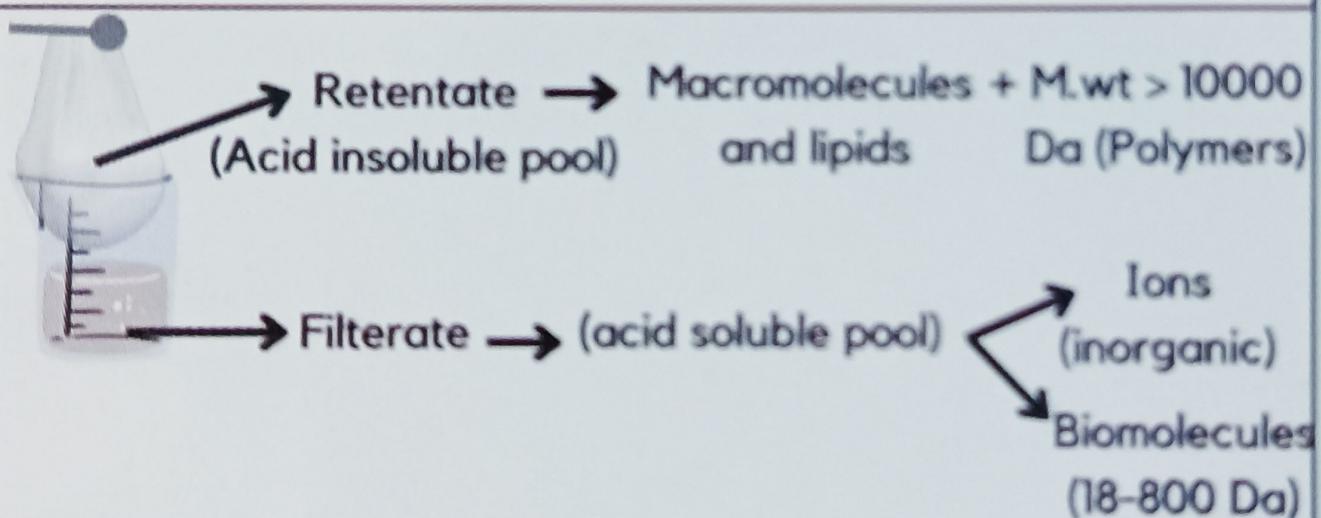
C-compounds obtained from living tissue - Biomolecules

Living tissue + Cl_3CCOOH trichloro acetic acid

grind in mortar

Obtained a thick slurry

Strain the slurry through a cheesecloth



To analyse inorganic elements

Living tissue $\xrightarrow{\text{dry}}$ dry weight $\xrightarrow{\text{burn}}$ carbon compounds \rightarrow 'ash' \rightarrow inorganic elements

oxidised

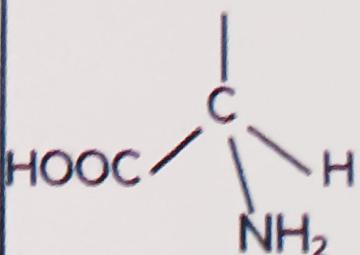
Contains inorganic elements (Ca, Mg, etc)

Usually Found

Na^+ , K^+ , Ca^{+2} , Mg^{+2} , H_2O , NaCl , CaCO_3 , PO_4^{3-} , SO_4^{2-}

α-Amino Acids - Substituted Methanes

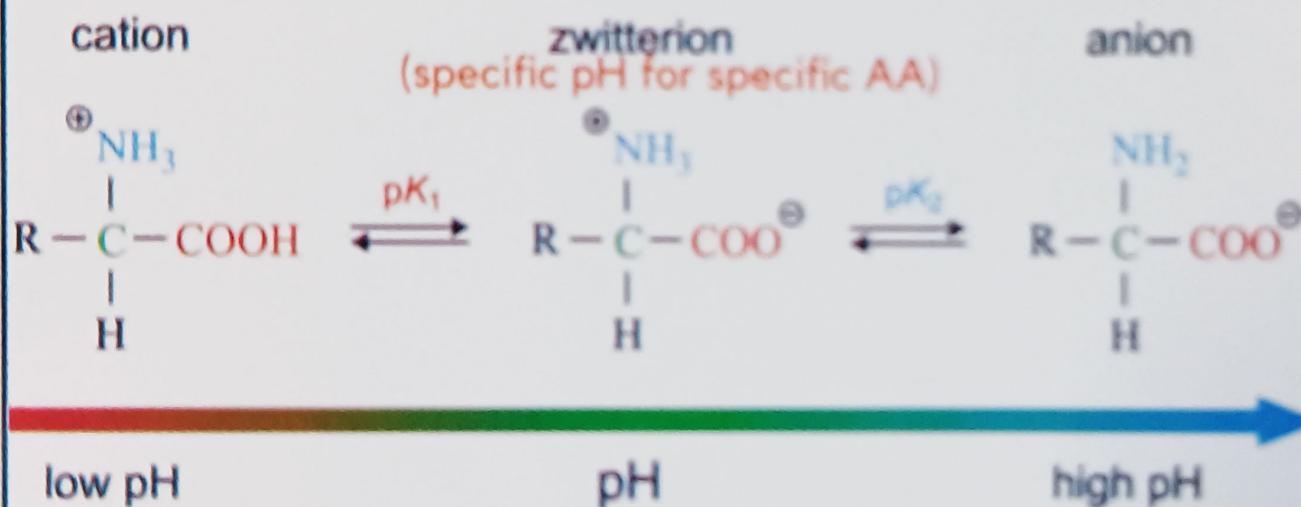
variable group or (R) → 20 types of Amino acid



R group	H	CH ₃	CH ₂ OH
Amino acid	Glycine	Aniline	Serine

Acidic AA	Basic AA	Aromatic AA
Glutamic acid	Histidine, arginine, Lysine	Tyrosine, tryptophan, Phenylalanine

Amino acids are ionisable (due to -NH₂ & -COOH)



FATTY ACIDS- water insoluble ; (R-COOH) form

- **Chain lengths**

- Palmitic acid = 16-C → H₃C-(CH₂)₁₄-COOH

- Arachidonic acid=20-C

- **Fatty Acids**
 - Unsaturated (with atleast 1 double bond)
 - Saturated (without multiple bonds)

Lipids (Esters of Fatty acids)

Monoglycerides (1FA)

Diglycerides (2FA)

Triglycerides (3FA)

Fats

Oils

High MP (Solid in winters)

Low MP (Liquid in winters)

Phospholipids : E.g. Lecithin (cell membranes)

Lipids that have phosphorus/phosphorylated organic compound

NUCLEIC ACID

Polynucleotides. → DNA/RNA (genetic material)

Nucleotide = Nucleoside (ester bond) Phosphate group

Nitrogenous Base + Monosaccharide

Purines
Learn by
CTU

Adenine
Guanine
Cytosine
Uracil
Thymine

Pyrimidine

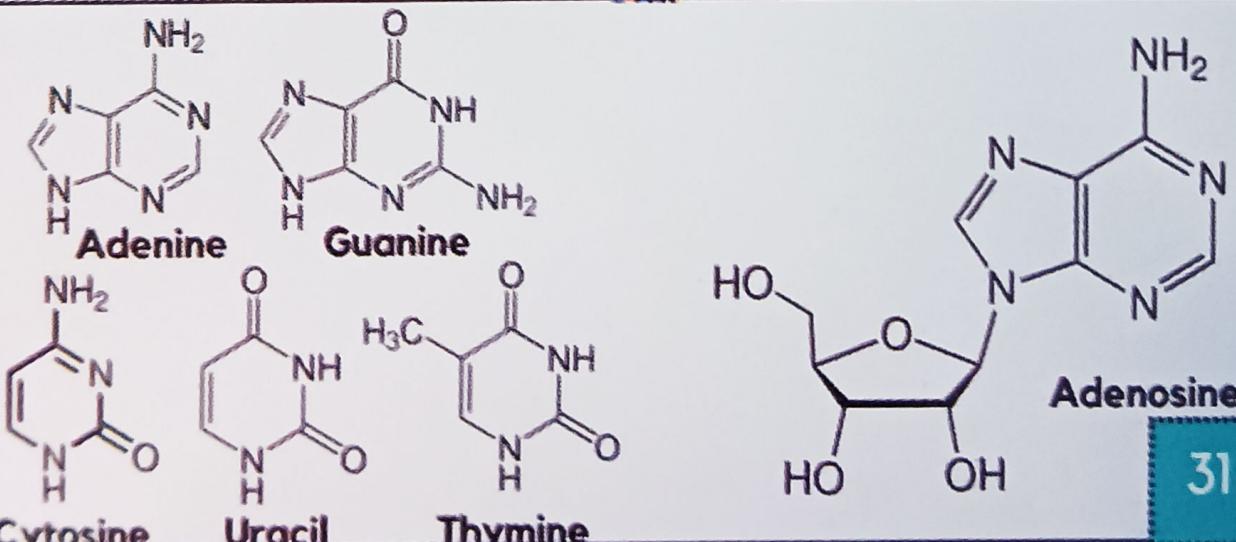


Uracil → RNA

Thymine → DNA

Linkage (b/w 2 nucleotides)

Phosphodiester linkage



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WATSON-CRICK MODEL

- Describes structure of B-DNA
- Antiparallel polynucleotide chains (move in opposite direction)
- Sugar-phosphate backbone
- A-T (2H-bonds) and G-C (3H-bonds)
[$\therefore A+G=T+C$]
- Each base pair turns strand to 36° (1 full turn = 10 base pairs)
- Pitch = 34 \AA° (per base pair - 3.4 \AA°)



METABOLITES

1° Metabolites	2° Metabolites
<ul style="list-style-type: none">• Identifiable functions (physiological processes) e.g.- Amino acids, nucleic acids, Carbohydrates etc.	<ul style="list-style-type: none">• Not involved in 1° metabolism• No direct function in growth & development• Useful for human welfare & also have ecological importance e.g.-Flavonoids, essential oils etc.

Pigments-Carotenoid, Anthocyanin

Alkaloids-Morphine, Codeine

Terpenoids-Monoterpenes, Diterpenes

Essential oils-Lemon grass oil

Toxin-Abrin, Ricin

Lectins-Concanavalin A

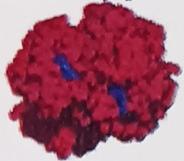
Drugs-Vinblastin, curcumin

Polymeric substance-Rubber, gums, cellulose

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PROTEINS

Amino acids linked by peptide bonds (heteropolymer)
(20 type of AA's)



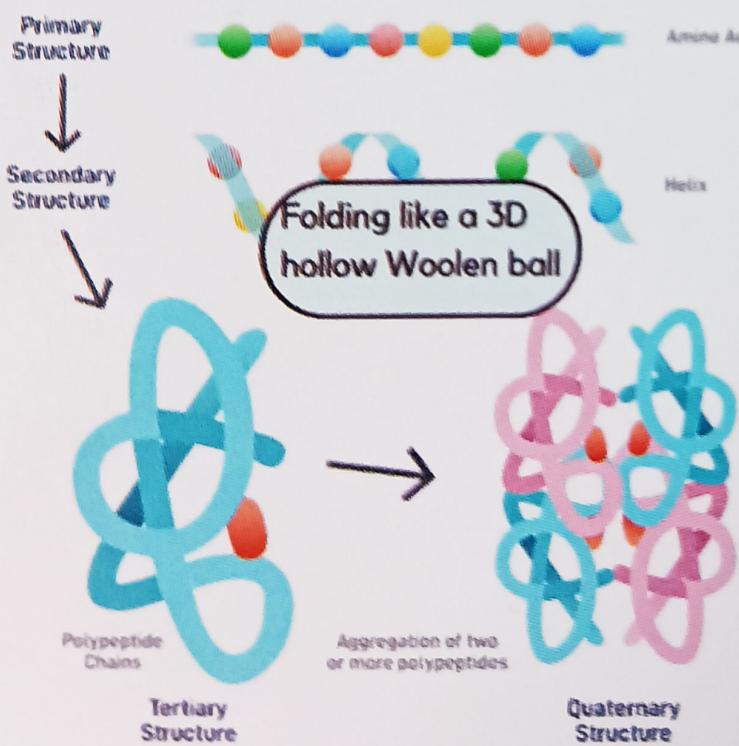
COOH of one amino acid +
NH₂ group of other
aminoacid (dehydration)

- Collagen- most abundant animal protein
- RuBisCO (Ribulose biphosphate Carboxylase Oxygenase) most abundant protein of Biosphere

TYPES OF AMINO ACIDS

ESSENTIAL	NON ESSENTIAL
<ul style="list-style-type: none">• Important for health• Supplied via diet	<ul style="list-style-type: none">• Produced by body itself• Not supplied via diet

PROTEIN STRUCTURE



Amino Acids attached as line/chain
Left : N-Terminus(1st AA)
Right : C-Terminus(Last AA)

Right Handed Helix formed



More than one polypeptide chain arranged with respect to each other

e.g - Haemoglobin is made of 4 subunits (2α + 2β)

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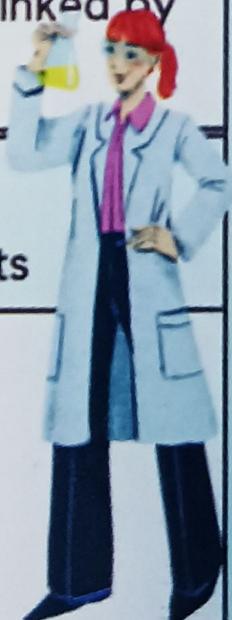
PROTEINS AND THEIR FUNCTIONS

- 1. Collagen
- 2. Trypsin
- 3. Insulin
- 4. Antibody
- 5. Receptor
- 6. GLUT-4

- Intercellular ground substance
- Enzyme
- Hormone
- Fight Infectious Agents
- Sensory Receptors (Smell, taste etc.)
- Enables glucose transport in cells

Polysaccharide

Long chain of carbohydrates / monosaccharides (linked by glycosidic bond) (formed by dehydration)



Homopolysaccharide- same monomer units

Heteropolysaccharides - different monomer units

- 1. Cellulose
- 2. Insulin
- 3. Glycogen
- 4. Starch
- 5. Chitin

- Homopolymer of Glucose
- Homopolymer of Fructose
- Homopolymer of Glucose
- Amylose and Amylopectin
- N- Acetyl Glucosamine

- Glycogen & Starch give Red & Blue colour with I_2 respectively.
- Plant cell wall is made of cellulose
- Exoskeleton of arthropods is made of chitin



METABOLISM

- Biomolecules tend to turnover (dynamic transformation to other biomolecules) through chemical reactions [Process - Metabolism]
- Such chemical reactions are linked forming a metabolic pathway.
- Flow of metabolites → dynamic state of body constituents.
- Types of metabolic pathways

Catabolic

- Degradation
- Complex to simple
- Energy released.

Glucose → lactic acid

Glucose → Ethanol

Glucose → Pyruvic acid

Anabolic

- Biosynthesis
- Simple to complex
- Energy used

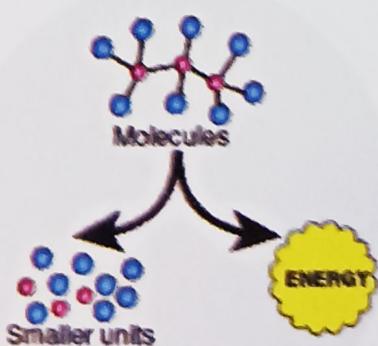
Acetic acid → cholesterol

Amino acid → Proteins.

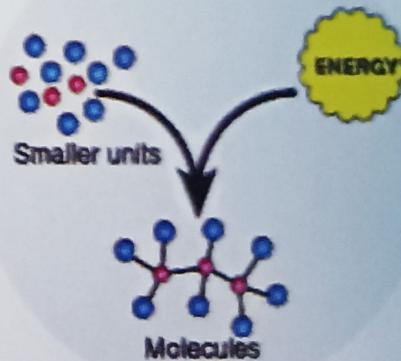
*Living state is a non-equilibrium steady-state to be able to perform work.

(Related to conc. of biomolecules) Blood conc. of glucose :
4.2-6.1 mmol/L

Catabolic



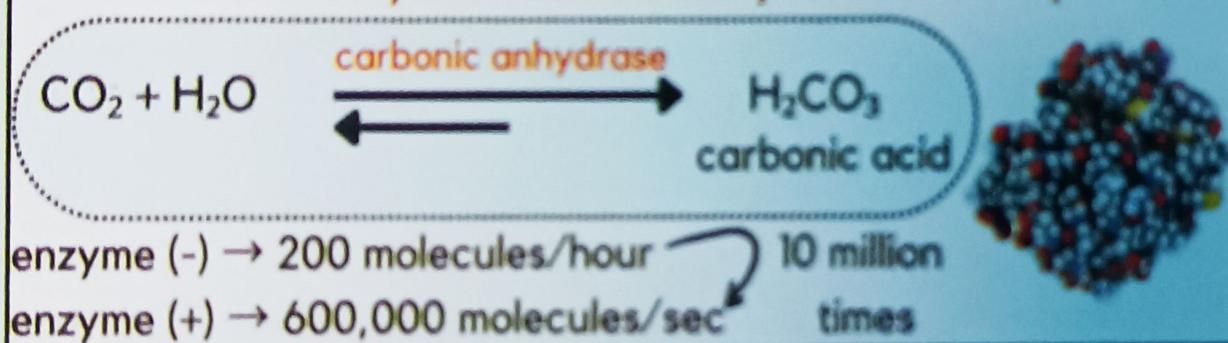
Anabolic



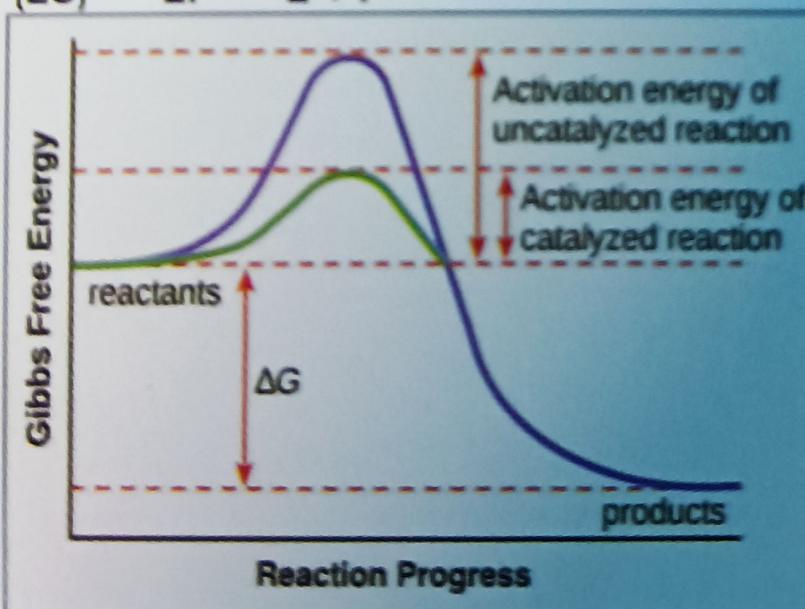
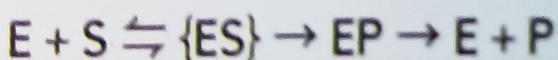
ENZYMES

- Proteins with catalytic action (fastening metabolic processes)
- Some nucleic acid are also enzymes-ribozymes
- Tertiary structure, remain unchanged at the end of reaction.
- Active site-specific sites to fit the particular substrate
- Enzymes denature at high temp. (may have specific optimal conditions to work) Rate = $\delta P / \delta T$

Rate usually doubles with every 10° rise in Temp



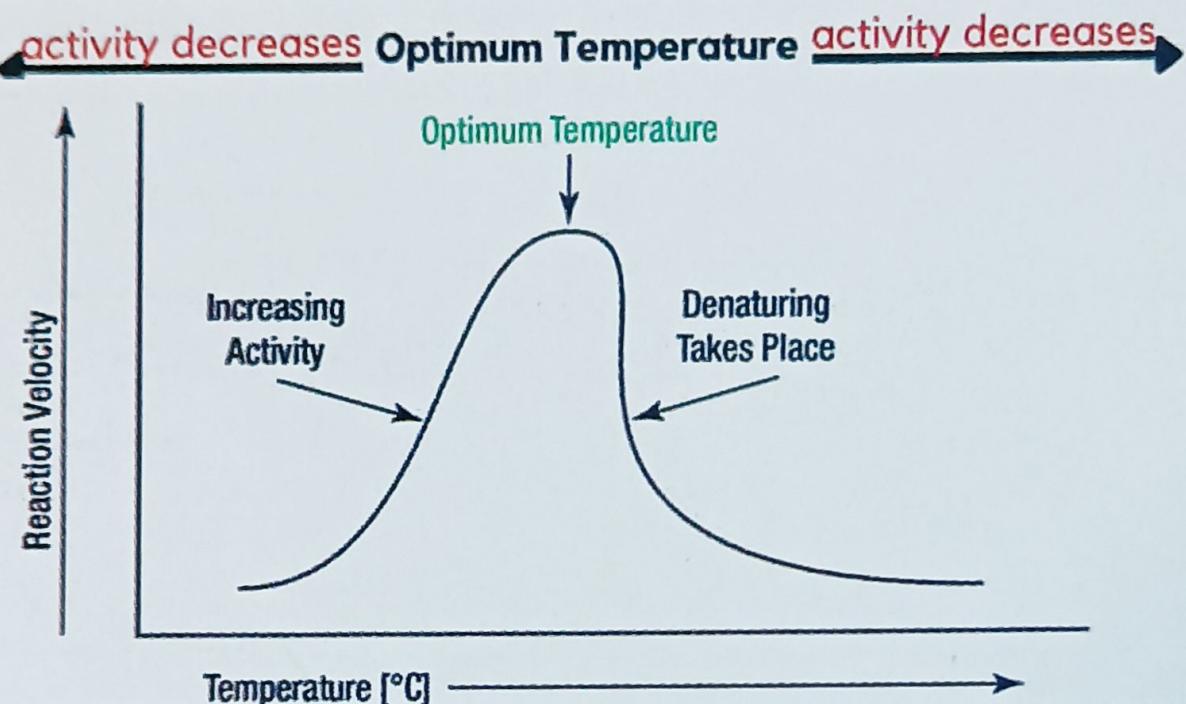
Action substrate 'S' binds to enzyme 'E' forming an obligatory 'ES' complex (transient phenomenon); making & breaking of bonds convert substrate to product, later releasing the product & unchanged enzyme



Factors Affecting Enzyme Activity

1. Temperature & pH

Optimum temp, and pH → (best activity)



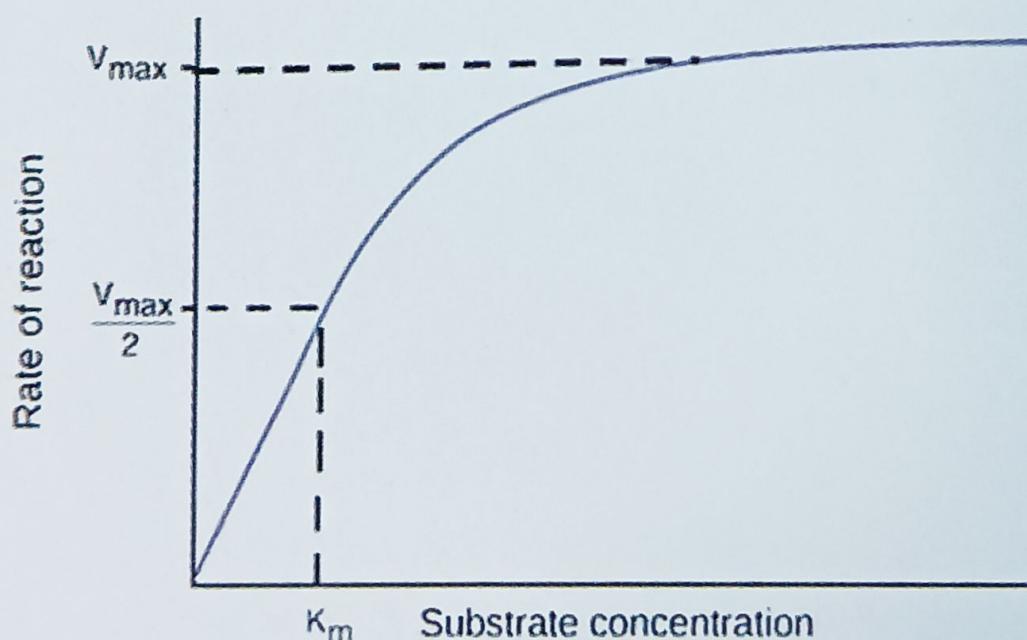
2. Substrate Concentration

On increasing enzyme inc. till saturation

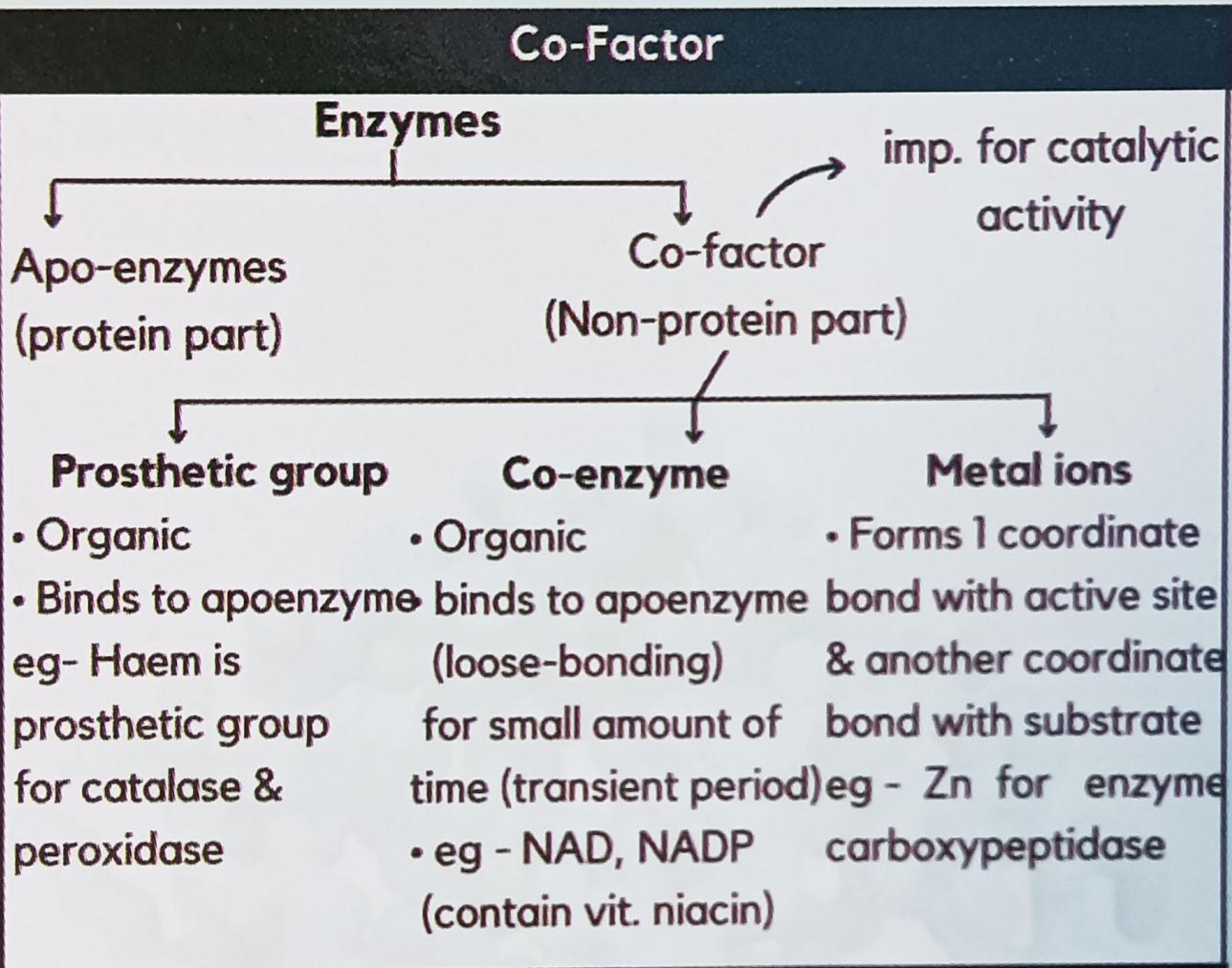
substrate \rightarrow reaction \rightarrow is attained \rightarrow
conc speed (V_{max})

No
free enzyme

↓
constant velocity



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Classification of Enzymes

Classification of enzymes - Basis-function (6-classes)

1. Oxidoreductase/dehydrogenase - catalyses simultaneous oxidation & reduction
 $S \text{ reduced} + S' \text{ oxidised} \rightarrow \text{oxidised} + S' \text{ reduced}$
2. Transferase-catalyses group transfer
 $S-G + S' \rightarrow S + S'-G$ ($\therefore S \text{ & } S'$ are substrates)
3. Hydrolases - catalyses hydrolysis of ester, ether, peptide etc.
4. Lyases - Catalyses group removal, forming double bonds

$$\begin{array}{c} X \quad Y \\ | \quad | \\ C-C \end{array} \rightarrow X-Y + C=C$$
5. Isomerases - catalyses interconversion of different isomers (optical, geometrical, etc.)
6. Ligases - catalyses linking of 2 component (forming C-O, C-S, C-N etc bonds)