

Quaternary: Containing more than 1 A.A chain.

## 29/10 ENZYMES

- Enzymes are mostly proteins (with exception of ribosomes) which are made up of RNA.
- All are globular proteins
- Catalysts
- Catalyzed reaction is reversible
- Efficient
- Highly specific.

Enzymes are biological catalyst which enhance the rate of biochemical reaction from  $10^6$  to  $10^{16}$  times when compared to uncatalyzed reaction.

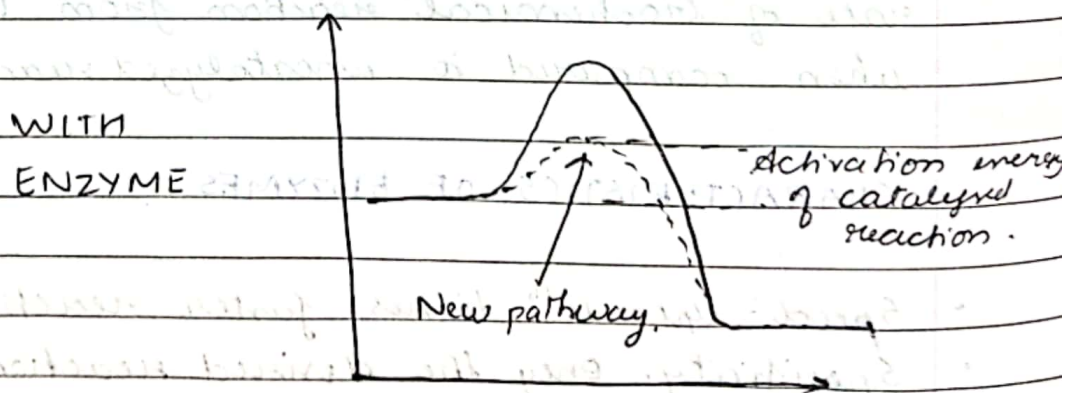
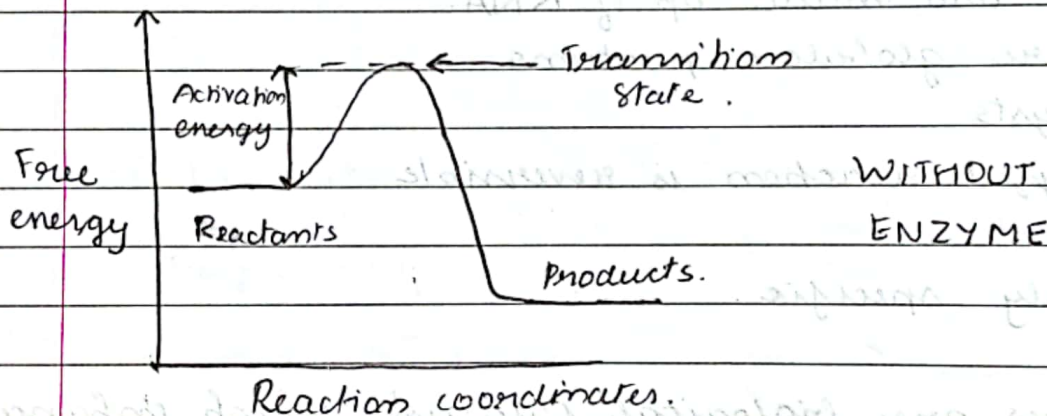
### CHARACTERISTICS OF ENZYMES

- Speed: Up to  $10^{16}$  times faster reaction rate.
- Specificity: Only the desired reaction occurs. Hence, high quality products, fewer by-products & purification process becomes much easier.
- Permit reactions ~~at~~ under mild conditions.

- Enzyme carries out its activity without being consumed in the reaction. Hence a small amount of ~~the~~ enzyme can act on large amount of substrate.
- Enzymes can be classified based on the chemical reaction it catalyses eg. oxidoreductase, isomerase, hydrolases etc.

## CHEMICAL REACTIONS

- Chemical reactions need an initial input energy i.e. Activation energy.
- During this part of the reaction the molecules are said to be in a transition state.





## ENZYME STRUCTURE

- Enzymes are proteins
- They have a globular shape.
- A complex 3-D structure.

## THE ACTIVE SITE

- One part of an enzyme, the active site, is particularly important
- The shape and the chemical environment inside the active site permits a chemical reaction to proceed more easily.

## COFACTORS

- An additional non-protein molecule that is needed by some enzymes to help the reaction.
- Tightly bound co-factors are called prosthetic groups
- Cofactors that are bound and released easily are called coenzymes.
- Many vitamins are called coenzymes.

## THE CATALYTIC SITE OF ENZYME

- Usually the size of the enzyme is bigger than that of the substrate
- Enzyme binds to substrate at least in three different regions called as catalytic sites.
- Different models were proposed regarding how enzyme binds to substrate.

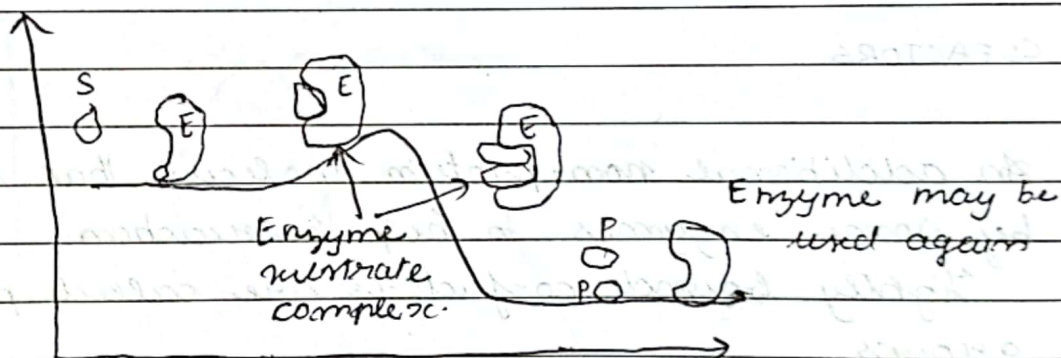
### A. The 'Lock & Key' or Template Model.

- The model was proposed by Emil Fischer.
- The enzyme is considered to be rigid.
- The catalytic site is presumed to be pre-shaped to fit the substrate.

### B. The "Induced Fit" Model.

- The model was proposed by Koshland.
- The enzyme is considered to be flexible.
- The substrate induces a conformational change in the enzyme.

### THE LOCK AND KEY HYPOTHESIS.



### WHAT HAPPENS IN AN ENZYME CATALYZED REACTION?

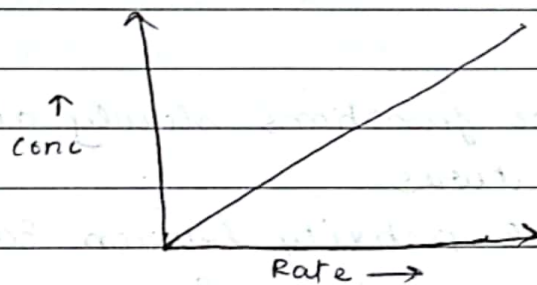
1.  $[E]$  binds to  $[S]$  to form  $[ES]$  complex (rate  $k_1$ )
2.  $[ES]$  can release substrate (rate  $k_3$ ) or convert  $[S]$  to  $[P]$  (rate  $k_2$ )
3. In steady state, the production and consumption of the transition state proceed at the same rate. So the concentration of transition state keeps a constant.
4. At equilibrium  $[S]$  and  $[P]$  remain constant.



## FACTORS AFFECTING ENZYMES ACTION

1. Concentration of Enzyme.
2. Substrate concentration.
3. Temperature.
4. pH.
5. Effect of product concentration.
6. Effect of activators.
7. Effect of light and radiation.

### Concentration of enzyme:

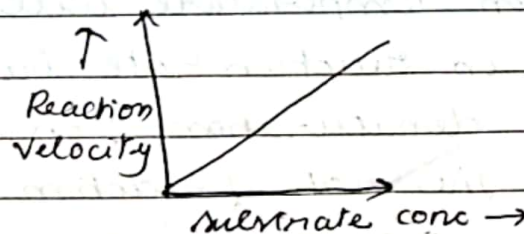


### The substrate:

They are the reactants activated by the enzyme.  
The Enzymes are specific to their substrates.

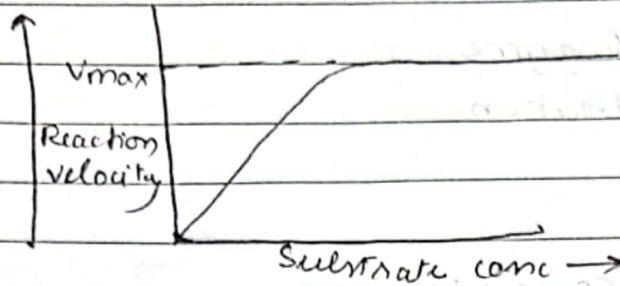
The specificity is determined by the active site.

Substrate conc: Non-enzymic reactions.



Increase in velocity  $\propto$  substrate conc.

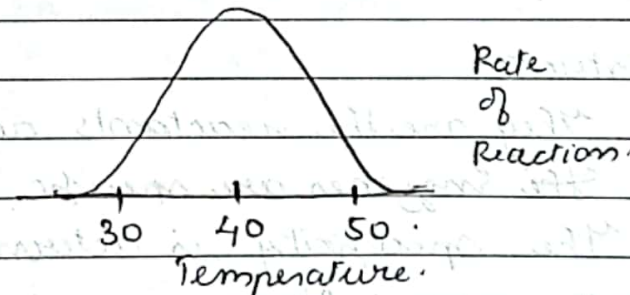
- Substrate conc: Enzymic reactions.



- Faster reaction but it reaches a saturation point when all ~~enz~~ enzyme molecules are occupied.
- If you ~~alter~~ alter the concentration of the enzyme then  $V_{max}$  will change too.

- Temperature:

- Enzyme functions slowly at sub-freezing temperatures
- Optimal-activity between  $30-40^{\circ}\text{C}$ .
- Denatures above  $45^{\circ}\text{C}$ .



- Increase in temperature will lead to an increase in reaction rate, but it also can lead to denaturation of enzyme.
- Usually the rate of reaction will double for every  $10^{\circ}\text{C}$  rise in temperature, thus speeding up the process.



## • The effect of temperature

- For most enzymes the optimum temperature is about  $30^{\circ}\text{C}$ .
- Many are a lot lower, cold water fish die at  $30^{\circ}\text{C}$  because their enzymes denature.
- A few bacteria have enzymes that can withstand temperatures up to  $100^{\circ}\text{C}$ .
- Most enzymes are however fully denatured at  $70^{\circ}\text{C}$ .
- The optimum temperature for an enzyme controlled reaction will be a balance between the  $Q_{10}$  and denaturation

## • pH:

- Extremes generally inactivates enzymes.
- pH optimum
- Maximum pH activity of most enzymes between pH 4.5 and 8.0.
- Narrow pH range.
- Exceptions.
  - Pepsin: Optimum pH is 1.8.
  - Trypsin: Optimum pH is 9.8.

## • The effect of pH.

- Extreme pH levels will produce denaturation.
- The structure of enzyme is changed.
- The active site is distorted and the substrate molecule will no longer fit in it.
- At pH values slightly different from the enzyme's optimum value, small changes in

the charges of the enzyme and its substrate molecules will occur.

- This change in ionisation will affect the binding of the substrate with the active site.

### Inhibitors.

- Inhibitors are chemicals that reduce the rate of enzymic reactions.
  - They are usually specific and they work at low concentrations.
  - They block the enzyme but they do not usually destroy it.
  - Many drugs and poisons are inhibitors of enzymes in the nervous system.
- The effect of enzyme inhibition.
    - Irreversible inhibitors: Combine with the functional groups of the amino acids in the active site, irreversibly.
    - Reversible inhibitors: These can be washed out of the solution of enzyme by dialysis.
- Effect of Product Concentration.
    - The accumulation of products generally decreases the enzyme velocity.
    - For certain enzymes, the product combine with the site of enzyme and form a loose complex, and thus inhibit enzyme activity (Feed back mechanism).



## • Effect of activators.

Some of the enzymes requires certain inorganic metallic cations like  $Mg^{2+}$ ,  $Mn^{2+}$ ,  $Ca^{2+}$ ,  $Co^{2+}$ ,  $Cu^{2+}$ ,  $Na^{2+}$ ,  $K^{+}$  etc. for their optimum activity.

- Metal activated enzymes: Enzyme is tightly held by the enzyme. Eg ATPase ( $Mg^{2+}$  &  $Ca^{2+}$ )
- Metalloenzymes: Enzymes hold the metals tightly. Eg: Alcohol dehydrogenase, carbon anhydrase.

## • Effect of light and radiation.

Exposure of enzymes to ultraviolet, gamma and/or x-rays may inactivate certain enzymes due to the formation of peroxides.

## SOURCES OF ENZYMES

- Biologically active enzymes are extracted from any living organisms.
- Selection of source of enzymes influences:
  - Type of extraction & purification process
  - Stability of the enzyme and,
  - Cost of the enzymes.