



**Enzymes** are <u>biological catalysts</u> made up of <u>proteins</u> used to speed up rate of chemical reactions. They <u>lower activation energy</u> by providing an alternative pathway and <u>remains chemically unchanged</u> at the end of the chemical reaction.

#### Characteristics of enzymes:

- three dimensional shape held by weak hydrogen bonds
- <u>specific</u> in nature
- affected by <u>temperature and pH</u>
- can catalyse <u>reversible</u> reactions



Human salivary amylase

#### Types of enzymes

- Hydrolases are hydrolytic enzymes that use water to cleave chemical bonds, usually dividing a large molecule into two smaller molecules. Examples of common hydrolases include esterases, proteases, glycosidases, nucleosidases, and lipases.
- **Synthases** are enzymes that catalase the synthesis of a substance without the use of a high-energy source such as cleavage of a phosphate bond in ATP.
- **Synthetases** are enzymes that catalyses the linking together of two molecules especially by using the energy derived from the concurrent splitting off of a pyrophosphate group from a triphosphate (as ATP).





**Topic: Enzymes** 



**KEY BIOLOGICAL PROCESS: How Enzymes Work** Substrates Product Active site Enzyme-substrate complex Enzyme Enzyme Enzymes have a complex An enzyme and its substrate(s) bind As a result, a chemical reaction three-dimensional surface to which tightly together, forming an occurs within the active site, forming enzyme-substrate complex. The binding the product. The product then particular reactants (called substrates of brings key atoms near each other and diffuses away, freeing the enzyme to that enzyme) fit, like a hand in a glove. stresses key covalent bonds. work again.

## LO: Explain enzyme action in terms of the 'lock and key' hypothesis

- Enzyme is the <u>"lock"</u>
- Substrate is the <u>"key"</u>
- Specific substrate fits into <u>active site of the enzyme</u>. The <u>three dimensional shape</u> of the substrate is <u>complementary</u> to the shape of the <u>active site</u> of the <u>enzyme</u>
- Formation of <u>enzyme-substrate complex</u>
- Reaction takes place then the enzyme and substrate separates.
- Enzymes remain <u>chemically unchanged</u> at the <u>end of the chemical reaction</u>.







## The induced fit model



#### LO: Explain the mode of action of enzymes in terms of an active site, enzymesubstrate complex, lowering of activation energy and enzyme specificity

- Enzyme binds to the substrate at the <u>active site</u> on the <u>enzyme</u> molecule.
- Results in the formation of an <u>enzyme-substrate complex.</u>
- Enzymes <u>reduce</u> the <u>activation energy</u> needed for the reaction to occur, so the reaction occurs more readily. <u>Activation energy</u> is the energy needed to start a chemical reaction.
- Enzyme specificity due to the <u>three dimensional shape of the enzyme held by</u> weak hydrogen bonds. The <u>active site of the enzyme is complementary to the</u> <u>three dimensional shape of the substrate</u>.



LO: Investigate and explain the effects of temperature and pH on the rate of enzyme catalysed reactions

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#### Low temperature

- Kinetic energy of particles is low, enzyme is inactive.
- Enzyme substrate complex formation is slow and rate of reaction is low.

#### Increasing temperature to optimum temperature

- Kinetic energy of particles increases, <u>increasing the frequency of collision of the</u> <u>substrate and enzyme molecules</u>.
- Rate of formation of enzyme substrate complex <u>increases</u> and rate of reaction increases.
- Note: Rate of reaction usually doubles for every 10°C increase.



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## **Optimum temperature**

- Enzyme is most active.
- Rate of formation of enzyme substrate complex is at its <u>maximum</u> and rate of reaction is the <u>highest</u>.

## Beyond optimum temperature

- <u>Weak hydrogen bonds</u> in three dimensional enzyme structure is broken.
- <u>Three-dimensional shape</u> of enzyme is lost and <u>shape of active site is altered.</u>
- <u>Substrate can no longer bind to active site of enzyme and no enzyme-substrate</u> <u>complexes are formed.</u>
- Enzyme is <u>denatured</u> and rate of reaction <u>decreases to zero</u>.



# Below optimum pH

- <u>Weak hydrogen bonds</u> in three dimensional enzyme structure is broken.
- <u>Three-dimensional shape</u> of enzyme is lost and <u>shape of active site is altered.</u>
- <u>Substrate can no longer bind to active site of enzyme and no enzyme-substrate</u> <u>complexes are formed.</u>
- Enzyme is <u>denatured</u> and rate of reaction <u>decreases to zero</u>.

# Optimum pH

- Enzyme is most active.
- Rate of formation of enzyme substrate complex is at its <u>maximum</u> and rate of reaction is the <u>highest</u>.

# Above optimum pH

- <u>Weak hydrogen bonds</u> in three dimensional enzyme structure is broken.
- <u>Three-dimensional shape</u> of enzyme is lost and <u>shape of active site is altered.</u>
- <u>Substrate can no longer bind to active site of enzyme and no enzyme-substrate</u> <u>complexes are formed.</u>
- Enzyme is <u>denatured</u> and rate of reaction <u>decreases to zero</u>.



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(a) Spontaneous refolding of ribonuclease following denaturation. Anfinsen's experiment showed that all the information needed for the proper folding of a ribonuclease polypeptide into its native three-dimensional conformation is present in its amino acid sequence.

