

# Enzymes (HL)

Where can we find ENZYMES?

	Intracellular	Extracellular
Location	Occur within a cell (free or attached)	Occur outside a cell (free or attached)
Example	Glycolysis in cytoplasm & Krebs cycle in mitochondria	Chemical digestion in stomach 

## METABOLIC ( BIOCHEMICAL ) PATHWAYS

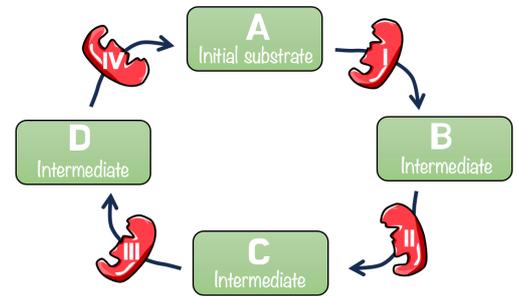
### 1. Linear metabolic pathway

Ex. glycolysis



### 2. Cyclic metabolic pathway

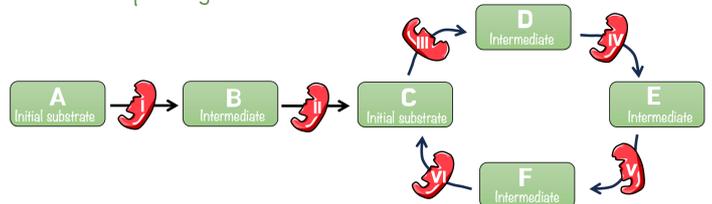
Ex. Krebs cycle (citric acid cycle), Calvin cycle



### KEY CONCEPTS

- Different pathways have **DIFFERENT** lengths.
- Each reaction requires its own **UNIQUE ENZYME** since enzymes are **SPECIFIC**.
- Cyclical pathway begins and ends with the **SAME** substance.
- ☀ The final product of one pathway could be the substrate of another pathway.

☀ In this example, C is both the final product in a linear metabolic pathway and the initial substrate in a cyclical metabolic pathway.

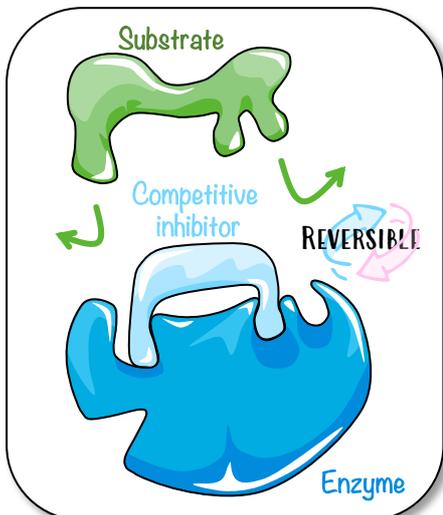


**INHIBITOR** – Molecule that binds to an enzyme & decreases its activity.

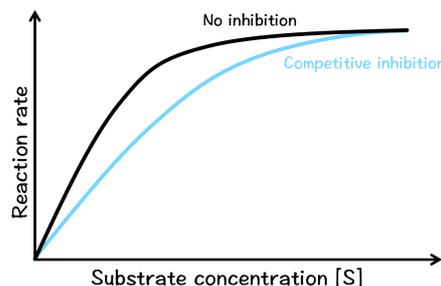
### NOTE!

- There are **3** types of inhibition:
- Competitive inhibition
  - Non-competitive inhibition
  - Mechanism-based inhibition

### Competitive INHIBITION



The competitive inhibitor **COMPETES DIRECTLY** with the usual substrate for the active site – it has a **SIMILAR STRUCTURE** to that normal substrate and therefore can **BLOCK THE ACTIVE SITE**.



It causes a decrease in rate of the chemical reaction

It is affected by substrate concentration and can be overcome by increasing the substrate concentration

Example – **STATINS** inhibit production of cholesterol 

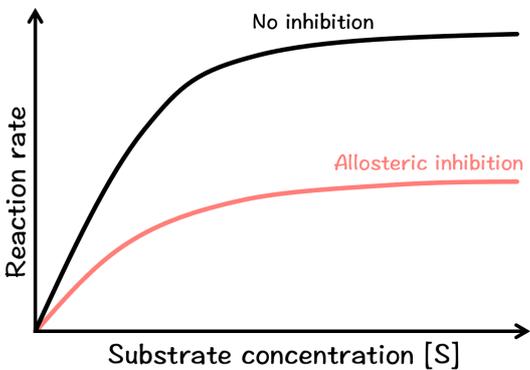
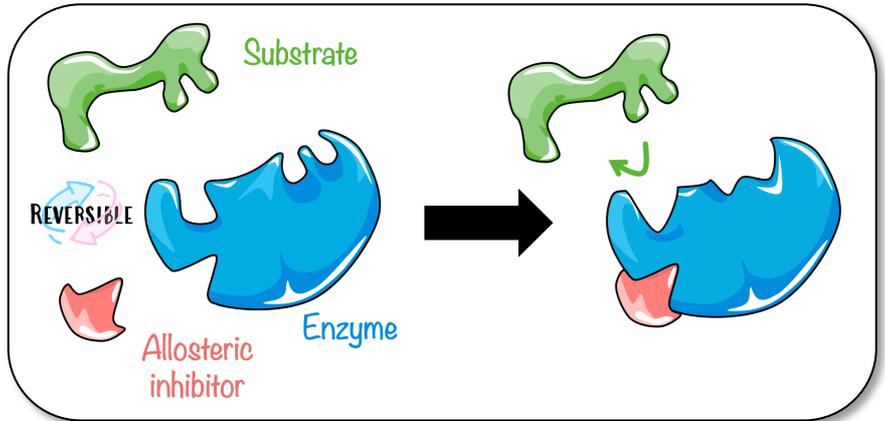


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## Non-competitive INHIBITION (allosteric)

The non-competitive inhibitor Instead binds to **ALLOSTERIC SITE** (AKA allosteric inhibition) - either on free-enzyme or enzyme-substrate complex.

It causes a change in the shape (CONFORMATIONAL CHANGE) of an enzymes active site, making it non-functional.



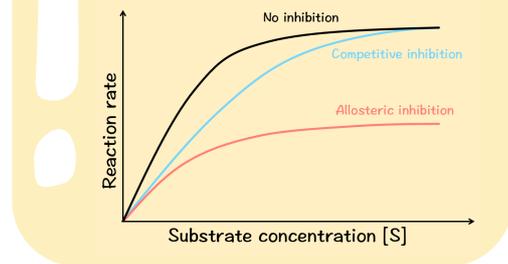
It causes a decrease in rate of the chemical reaction.  
Lower maximum rate

They are not influenced by the concentration of the substrate

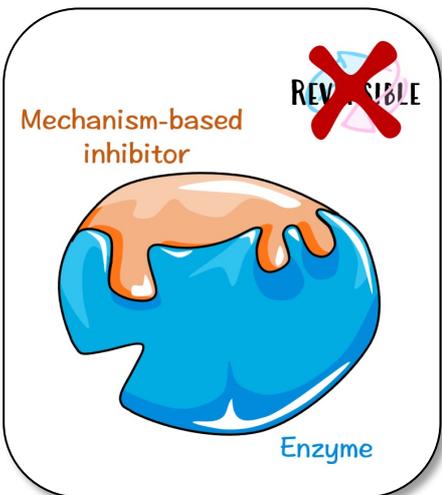
Rate can increase if enzyme concentration is increased

### NOTE!

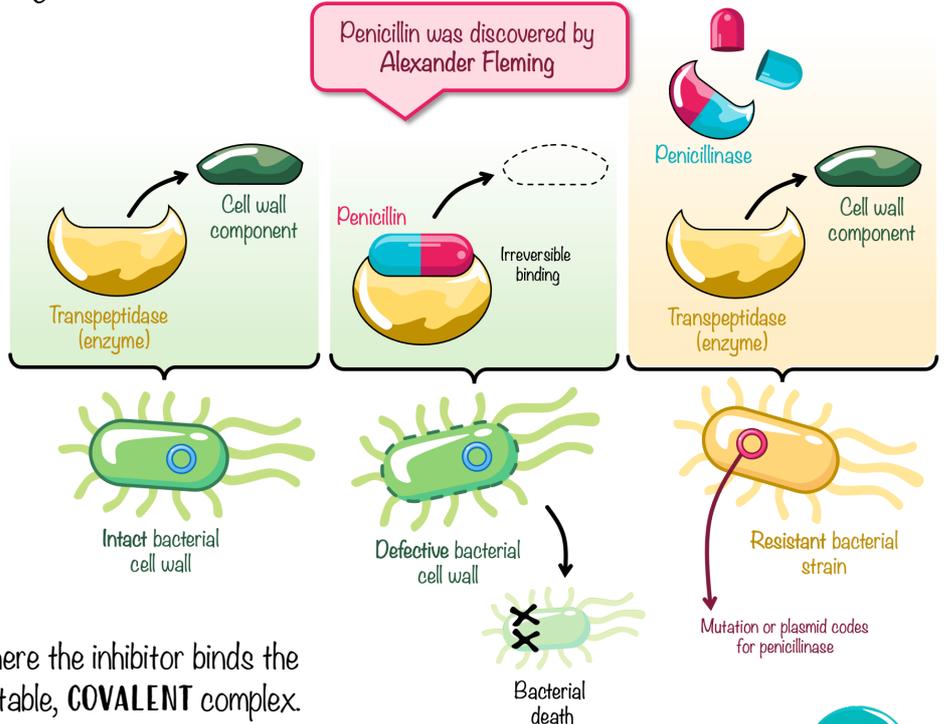
Make sure you can recognize the graphs for competitive and non-competitive inhibition in comparison to no inhibition.



## Mechanism-Based INHIBITION



Penicillin was discovered by Alexander Fleming



An irreversible form of inhibition where the inhibitor binds the enzymes' active site and forms a stable, **COVALENT** complex.

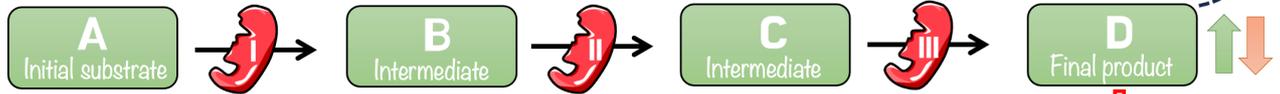
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## FEEDBACK INHIBITION (End product inhibition)

Prevents the cell from wasting **CHEMICAL RESOURCES AND ENERGY** by making more of substance than it doesn't need. A form of negative feedback.



Concept: Linear metabolic pathway



Typically act on the allosteric site

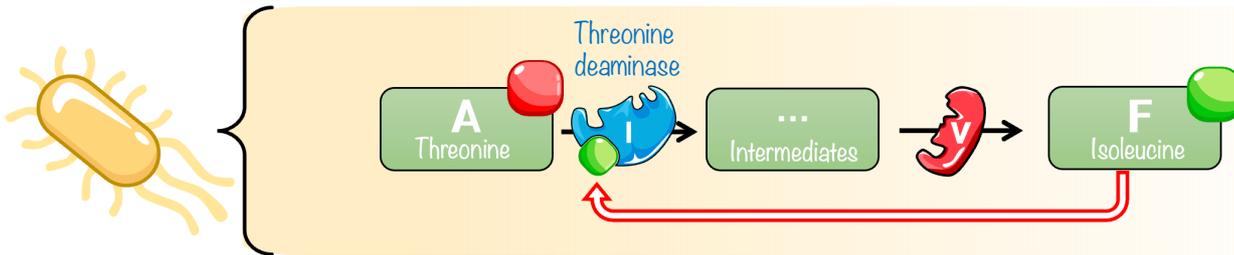
The end-product inhibits the action of the enzyme in the first step of the pathway.

↑ When present in high concentrations, the end-product binds with the allosteric site of the first enzyme, thus bringing about inhibition.

↓ As the existing end-product is used up by the cell, the first enzyme is reactivated.

This is done so that a build up of intermediate substrates in the cell does not occur. Energy is also conserved since the following reactions in the pathway would not occur.

Real example:



## ENZYMES AND COFACTORS

Many enzyme need an inorganic ion of a non-protein organic molecule to work properly. These are called **COFACTORS**.

A common example of a cofactor are vitamins.

### BIG BRAIN TIP!

#### ORGANIC MATTER -

Contain C-H bonds. Includes the 4 macromolecules.

The **4** macromolecules

CARBOHYDRATES

NUCLEIC ACIDS

LIPIDS

PROTEINS

#### INORGANIC MATTER -

Do not contain C-H bonds but may contain carbon, includes rocks, minerals etc.

