



## Commentary

# Enzyme inhibition and its types

Peter Alexanderio\*

Department of Biochemistry and Molecular Biology, Dakar Bourguiba University of Dakar, Daka, Senegal

\*Corresponding Author's Email: [peteralexrio@gmail.com](mailto:peteralexrio@gmail.com)

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## DESCRIPTION

Enzymes are essential for the majority, of present life's functions. Enzymes catalyse reactions by lowering the activation energy required for them to take place. Enzymes, on the other hand, must be strictly managed to ensure that product levels do not escalate to unacceptably high levels. Enzymes catalyse nearly every cellular function. Modifiers are inorganic and organic compounds that modify the catalytic activity of specific enzymes. Activators (Positive modifiers) are compounds that boost enzyme activity, while inhibitors are substances that decrease enzyme activity (Negative modifiers). Enzyme inhibitors are compounds that turn enzymes into inactive substances, slowing down the pace of enzyme-catalyzed reactions. Enzyme inhibition is the term for this type of process.

### Types of inhibition environmental impacts

Inhibitors, both reversible and irreversible, attach to an enzyme and decrease its activity. One way to do this is to bind to an enzyme for an extended period of time. These inhibitors are known as irreversible inhibitors. Other substances, on the other hand, can temporarily bind to an enzyme. These are said to as reversible. Competitive inhibitors bind to an active site, while reversible inhibitors bind to another site on the enzyme (non-competitive inhibitors).

**Competitive inhibitors:** The inhibitor can mix with the free enzyme in such a way that it competes with the usual substrate for binding at the active site. Analogue of the substrate. It's also known as inhibition. An enzyme-inhibitor complex [EI] is formed similarly to the enzyme-substrate complex [ES]. By increasing the substrate concentration, the rate of inhibition can be delayed. The concentration level of substrate and inhibitor define the degree of inhibition.

**Uncompetitive Inhibitors:** The inhibitor does not interact with the free enzyme or its regular substrate; however, it does interact with the enzyme-substrate complex. Here, an

inactive enzyme-substrate-inhibitor complex [ESI] forms, preventing the usual product from undergoing further processing. When the substrate concentration is increased, the degree of inhibition may increase.

**Noncompetitive inhibitors:** The inhibitor can bind to either the free enzyme or the enzyme-substrate complex, preventing both from working. Inhibitors bind to a place other than the active site of the enzyme. Inhibitors frequently distort the enzyme, preventing it from forming the [ES] complex at its normal pace and, if created, from decomposing at its normal rate to give products. [ESI] and [EI], two inactive complexes, are produced. Increasing the substrate concentration does not reverse the degree of inhibition.

**Mixed inhibition:** In multi-substrate reactions, this sort of inhibition is widespread. It's the result of a mix of competitive and non-competitive inhibition. The mixed inhibitor has the ability to bind to both the active and allosteric sites.  $V_{max}$  drops as  $K_m$  rises in the kinetics of the reaction. Because the inhibitor binds to the allosteric site non-competitively and distorts the enzyme, the  $V_{max}$  lowers. Similarly,  $K_m$  rises as the inhibitor competes with the substrate for the active site. Increased substrate concentration will not remove this form of inhibition.

### Function of enzyme inhibitor

- Enzyme inhibitors are used to learn about the structure of an enzyme's active site and the amino acid residues that make up the active site.
- They're utilised to figure out how a metabolic pathway is regulated or controlled.
- They can be utilized to create new drugs.
- They're crucial for resolving metabolic issues.
- They're utilised to create herbicides, pesticides, and disease killers.