

## Chapter 5 - Chemical Kinetics

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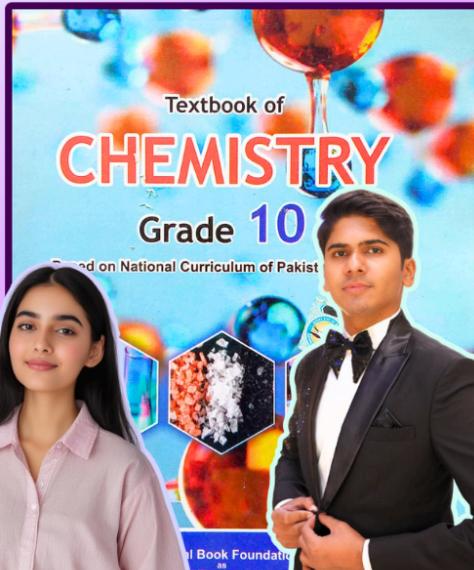
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### Chemical Kinetics

Chemical Kinetics is the study of:

- The **rates** (speeds) of chemical reactions.
- The **mechanisms** (steps) by which reactions occur.
- The **factors** that influence these rates.

Reaction rates vary greatly:

- **Slow:** Fermentation (weeks), Digestion.
- **Fast:** Acid-base neutralization (microseconds).

- **Moderate:** Muscle contraction, nerve impulses, photography.

**Importance:** Understanding kinetics is crucial for industry to make chemical processes **cost-effective**.

## 5.1. Rates of Reactions

The rate of a reaction tells us how quickly **reactants are consumed** or **products are formed** over time. It is defined as the **change in concentration** of a reactant or product per unit time.

**Mathematical Formula:**

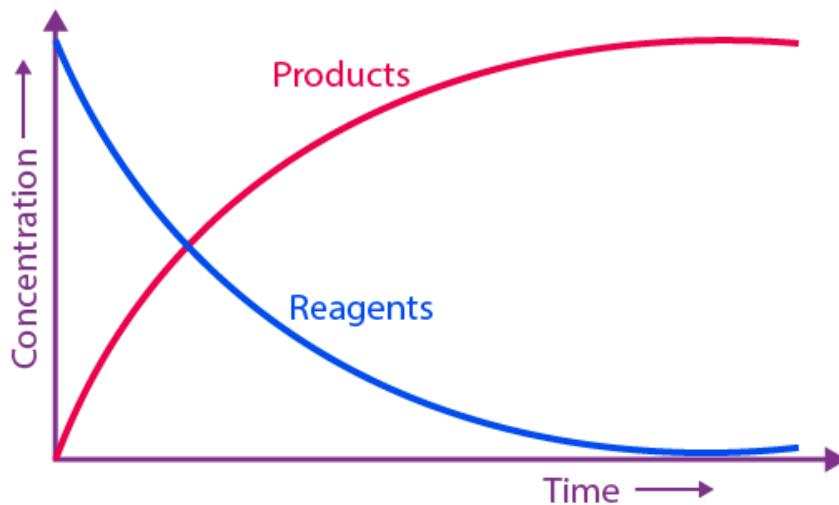
$$\text{Rate} = \frac{\text{Change in concentration of a substance}}{\text{Time taken for change}}$$

**Common Units:** moles per cubic decimeter per second ( $\text{mol dm}^{-3} \text{ s}^{-1}$ ).

### Graphical Representation (Concentration vs. Time)

A typical graph shows two curves:

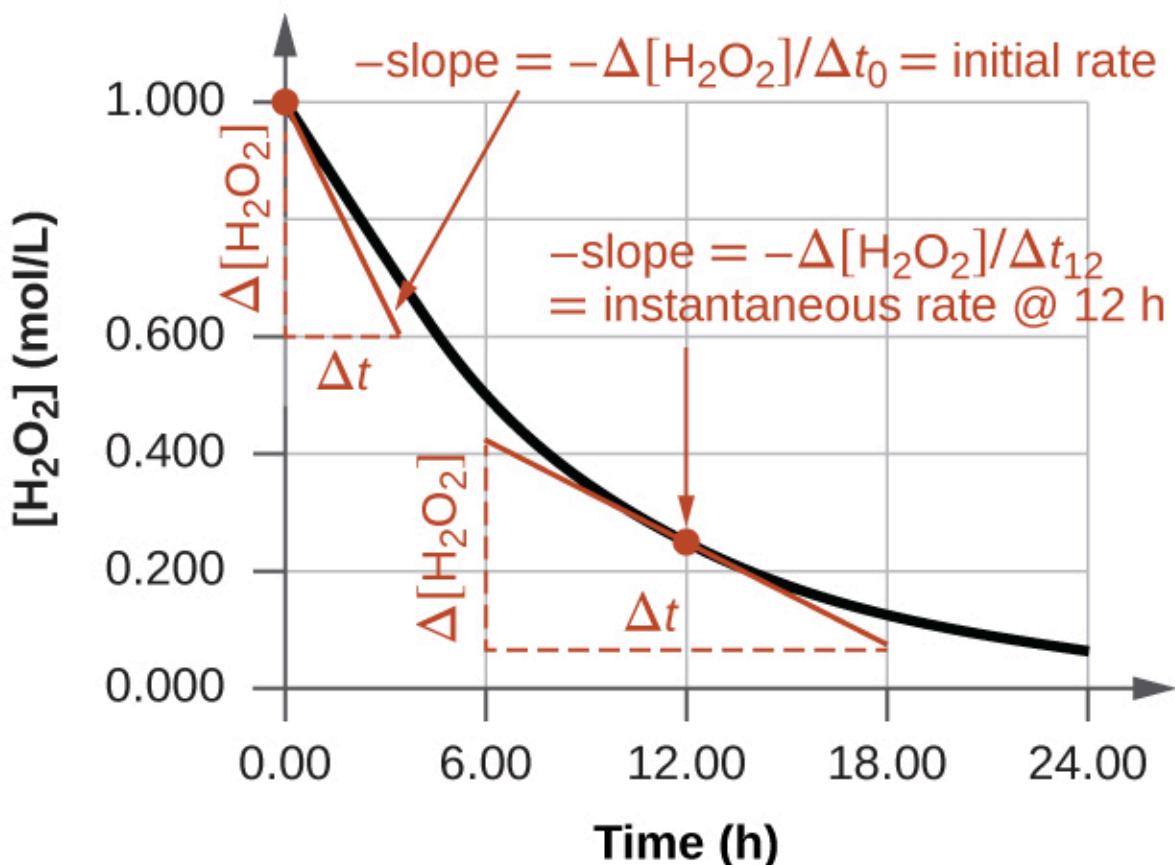
- **Reactant Curve:** Starts high and decreases over time.
- **Product Curve:** Starts at zero and increases over time.



**Observations from the Graph:**

### 1. Initial Stage (Steep Slope):

At the start, the curves are steep. This means the reaction is **fast** because there are many reactant particles available to collide.



### 2. As Time Progresses (Slope Decreases):

The curves become less steep. The reaction **slows down** because the concentration of reactants decreases, leading to fewer successful collisions.

### 3. Final Stage (Flat Curve):

The curves become horizontal (flat). The reaction has **stopped** because the reactants are fully used up, or the system has reached **equilibrium**.

**Conclusion:** The rate of a reaction is **not constant**; it is highest at the start and decreases over time.

## Average Rate of Reaction

This gives the overall speed of the reaction over a specific time interval.

**Formula:**

$$\text{Average rate} = \frac{\text{Total change in concentration}}{\text{Total time taken}}$$

## Expressing Reaction Rate Mathematically

For a simple reaction:  $\text{A} \rightarrow \text{B}$

- The rate can be expressed as the **disappearance of reactant A**:

$$\text{Rate} = -\frac{d[\text{A}]}{dt}$$

(The negative sign shows concentration is decreasing)

- Or as the **appearance of product B**:

$$\text{Rate} = +\frac{d[\text{B}]}{dt}$$

(The positive sign shows concentration is increasing)

Where  $d[\text{A}]$  and  $d[\text{B}]$  are the small changes in concentration, and  $dt$  is the small change in time.

## Interpreting Reaction Rate Data

### Example Reaction: $\text{A} + \text{B} \rightarrow \text{C}$

The rate of this reaction can be followed by measuring the **concentration of the product (C)** at regular time intervals.

### Data Table & Analysis:

Time(min)	Concentration of C ( $\text{mol dm}^{-3}$ )	Interpretation
0.0	0.00	Reaction has not yet started.
20	15	Product C is forming <b>rapidly</b> . The rate is high.
40	21	The increase in [C] is slower, indicating the <b>reaction is slowing down</b> .
60	23	The reaction continues to slow. Very little increase in [C].
80	25	No further change in concentration.
100	25	The reaction has <b>stopped</b> ; likely because a reactant has been used up.

The data visually demonstrates that the **reaction rate decreases over time** as reactants are consumed.

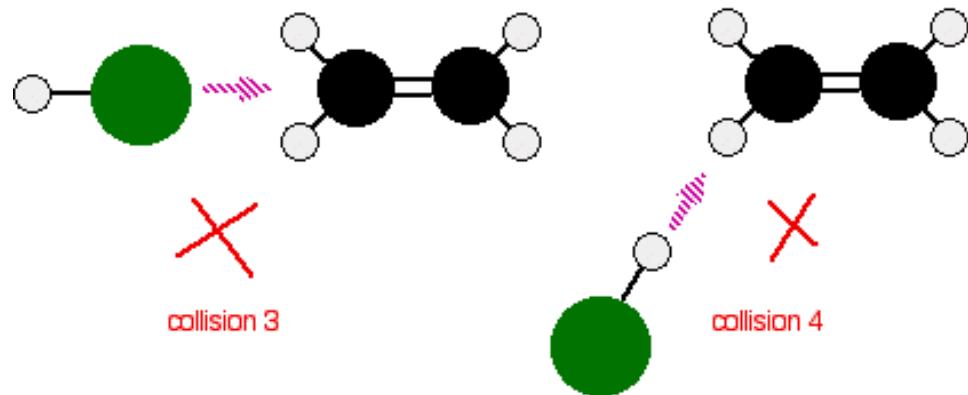
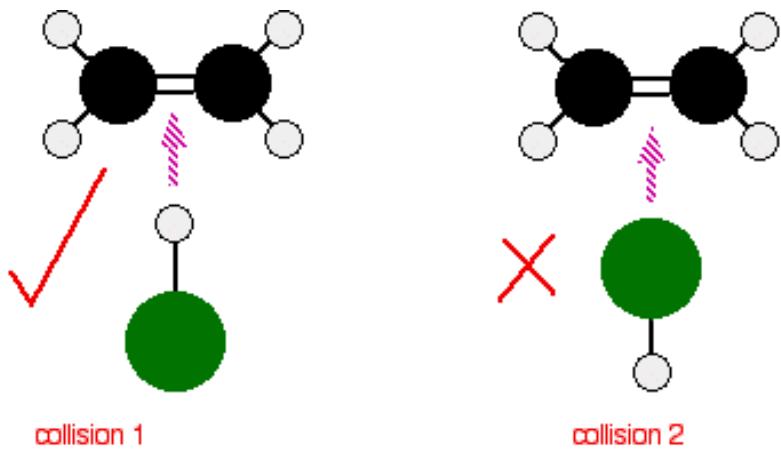
## 5.2. Collision Theory and Activation Energy

Collision theory explains the requirements for a chemical reaction to occur at the molecular level.

### Two Conditions for a Successful Reaction

For a collision between particles to result in a reaction, two criteria must be met:

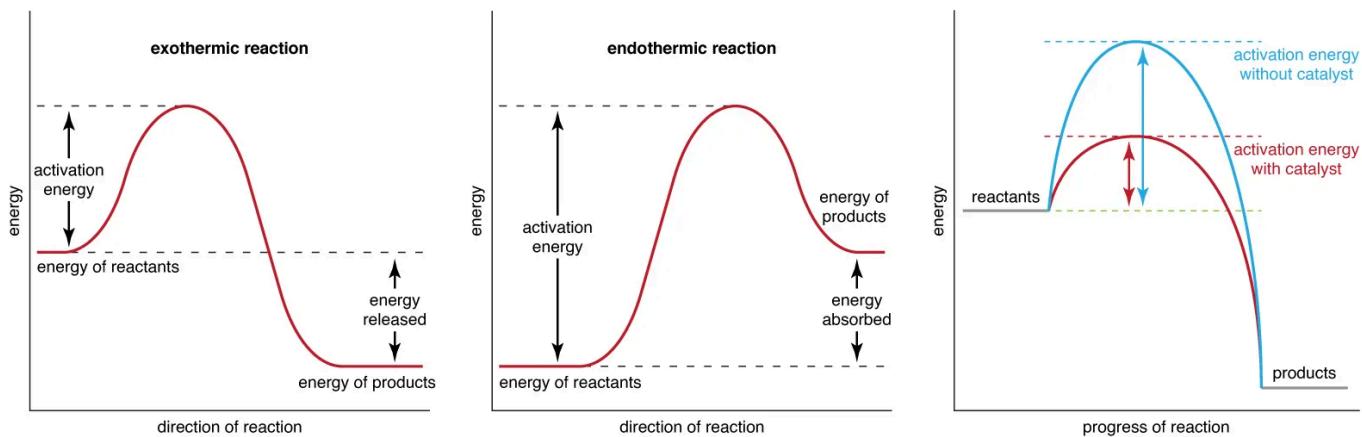
- **Sufficient Energy:** Colliding particles must possess enough kinetic energy to overcome the repulsion between their electrons and break existing chemical bonds.
- **Correct Orientation:** The particles must collide in a specific spatial alignment that allows the atoms to rearrange and form new bonds.



### Activation Energy ( $E_a$ ):

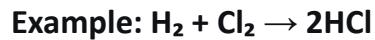
This is the **minimum amount of energy** required for a collision to be effective and lead to a reaction.

- **High  $E_a$ :** Fewer particles have this energy, so the reaction is slower.
- **Low  $E_a$ :** More particles have this energy, so the reaction is faster.

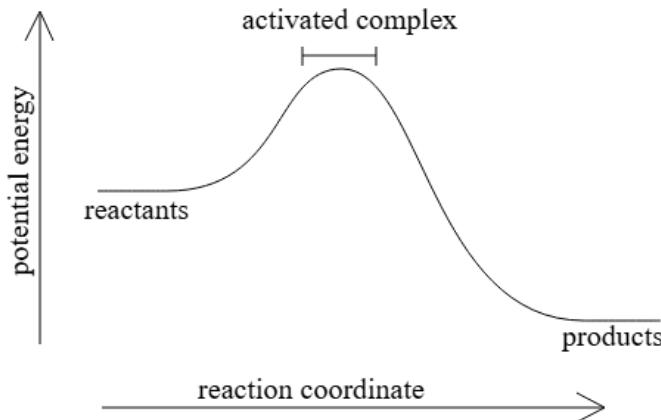


### Activated Complex (Transition State):

- During an effective collision, particles form a temporary, high-energy, and unstable species called the **activated complex**.
- It is a transitional structure where old bonds are breaking and new bonds are beginning to form.
- It exists for a very short time before breaking apart to form the final products.

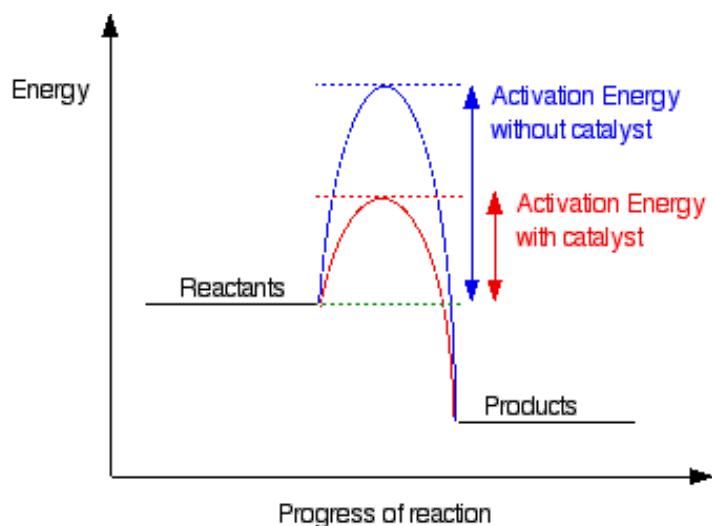


1. **Reactants:**  $\text{H}_2$  and  $\text{Cl}_2$  molecules collide effectively.
2. **Activated Complex:** A high-energy arrangement (e.g., H--Cl--Cl) forms momentarily.
3. **Products:** The complex breaks down to form two stable  $\text{HCl}$  molecules



### 5.3. Catalysts and their Role in Reaction Kinetics

A catalyst is a substance that speeds up a chemical reaction without being consumed or used up in the process. It provides an alternative pathway for the reaction that has a lower activation energy.



#### How Does a Catalyst Work? (The "Hill" Analogy)

- **Without a Catalyst:** Reacting particles must overcome a high energy "hill" (the original activation energy). Few particles have enough energy, so the reaction is slow.
- **With a Catalyst:** The catalyst provides a different, **lower hill** (lower activation energy). This allows **more particles** to have the required energy to react, leading to a faster reaction.

#### Properties of Catalysts

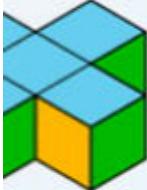
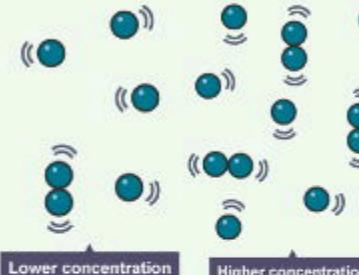
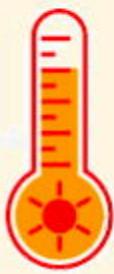
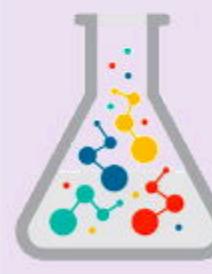
- **Not Consumed:** A catalyst remains **unchanged in mass and composition** at the end of the reaction.
- **Does Not Initiate Reactions:** A catalyst **cannot** make a non-spontaneous reaction happen. It only speeds up a reaction that is already feasible.
- **No Overall Energy Change:** A catalyst does not alter the **total energy change ( $\Delta H$ )** of the reaction between reactants and products.

### 5.3.1. Physical Parameters that Affect the Rate of Reaction

Several measurable physical changes indicate that a reaction is occurring and can be used to track its rate.

#### 1. Change in Mass

As reactants are used up, their **mass decreases**. As products are formed, their **mass increases**. Monitoring mass change over time is a direct way to measure the reaction rate.

What factors affect the rate of reaction?	
<b>Surface Area</b>  <p>The <b>larger</b> the surface area of your reactants, the <b>faster</b> the rate of reaction ➤ More surface for a reaction so <b>more reactant collisions</b>.</p>	<b>Concentration</b>  <p>The <b>higher</b> concentration of reactants, the <b>faster</b> the rate of reaction ➤ There are <b>more particles</b> in the same volume so <b>more collisions</b> between reactants.</p>
<b>Temperature</b>  <p>The <b>higher</b> the temperature of your reactants, the <b>faster</b> the rate of reaction ➤ Particles have <b>more energy</b> so move <b>faster</b>, increasing the <b>rate of collisions</b>.</p>	<b>Catalyst</b>  <p>A catalyst <b>speeds up</b> the reaction but is <b>NOT</b> used up ➤ Catalysts increase the <b>number</b> of successful collisions between reacting particles by <b>lowering activation energy</b>, thus reactions take place at <b>lower temperatures</b>.</p>

#### 2. Formation of a Gas

In an **open system**, if a gas is produced and escapes, the **mass of the reaction mixture will decrease**. In a **closed system**, the gas cannot escape. As more gas is produced, the **number of gas particles increases**, leading to an **increase in pressure** (if volume is constant). This pressure increase can be measured to track the rate.

### 3. Temperature

**Increasing the temperature** increases the reaction rate. At higher temperatures, particles have **more kinetic energy**. They move faster, leading to more frequent collisions. A greater proportion of particles have energy equal to or greater than the activation energy, leading to more successful collision

#### 5.3.2. Factors Affecting Rate of Reactions

**Several** factors influence the reaction rate by affecting the frequency and **effectiveness** of collisions between particles.

##### 1. Concentration of Reactants

Higher concentration leads to a faster reaction rate. More reactant particles in a given volume increase the frequency of collisions.

###### Examples:

- Stronger (higher  $\text{H}^+$  concentration) in acid rain damages marble faster.
- Two antacid tablets neutralize acid faster than one because more reacting particles are present.
- Doubling the concentration of  $\text{H}_2$  or  $\text{Cl}_2$  gas doubles the reaction rate.

##### 2. Surface Area (for Solids)

A larger surface area leads to a **faster reaction rate**. Breaking a solid into smaller pieces or powder exposes more particles to the other reactant, increasing the **number of sites where collisions can occur**.

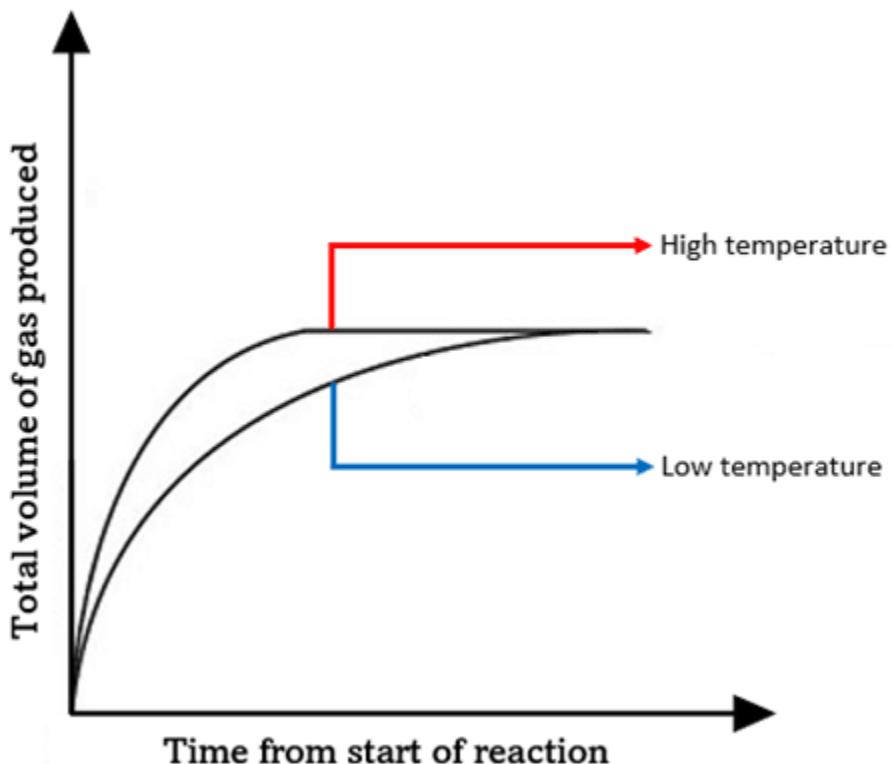
###### Examples:

- Zinc powder reacts with dilute HCl faster than a zinc lump.
- Powdered aluminum reacts quickly with NaOH, while aluminum foil reacts slowly.

### 3. Temperature

Higher temperature leads to a **much faster reaction rate**.

- **More Frequent Collisions:** Particles move faster, leading to more collisions per second.
- **More Energetic Collisions:** A greater proportion of particles have kinetic energy equal to or greater than the **activation energy ( $E_a$ )**.



#### 4. Pressure (for Gaseous Reactions)

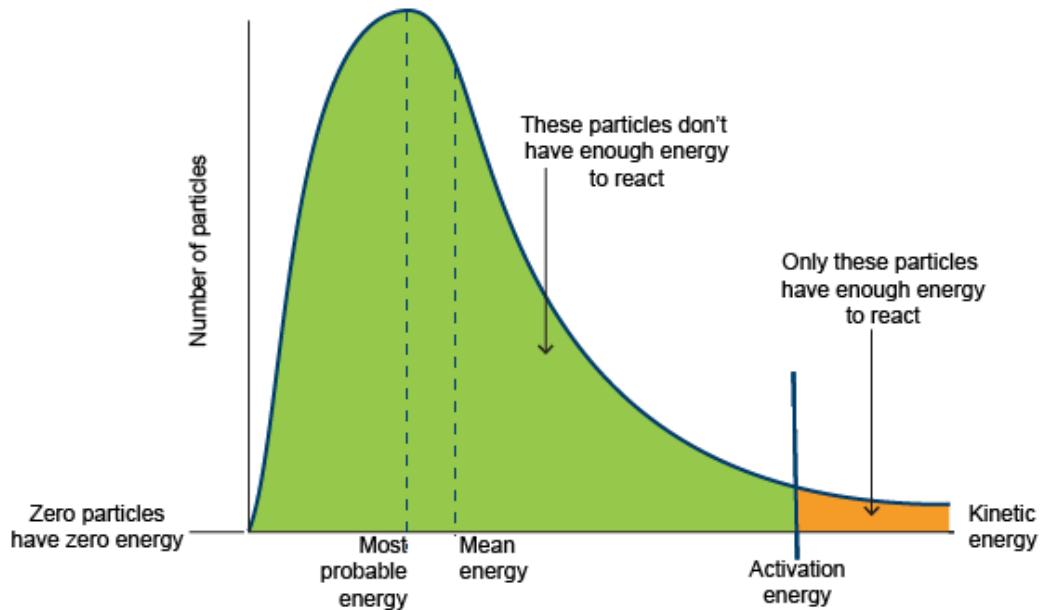
Higher pressure leads to a **faster reaction rate**. Increasing the pressure of a gas is equivalent to increasing its **concentration** in a given volume. This forces gas particles closer together, resulting in more frequent collisions.

**Example:** Doubling the partial pressure of  $H_2$  or  $Cl_2$  in their reaction mixture doubles the reaction rate.

#### Maxwell-Boltzmann Energy Distribution

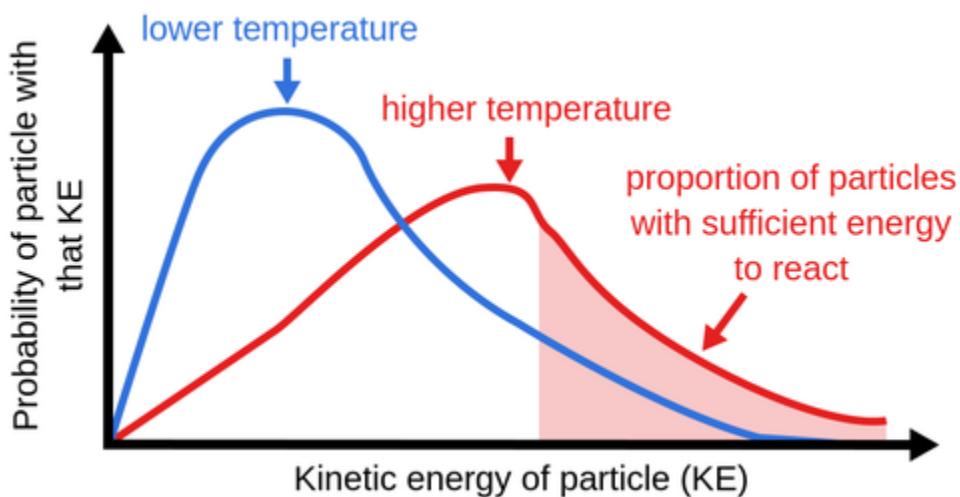
This concept explains *why* temperature has such a significant effect on reaction rate. The Maxwell-Boltzmann curve shows the distribution of kinetic energies among particles in a

gas at a specific temperature. No particles have zero energy. Only a **small fraction** of particles have energy greater than or equal to the activation energy ( $E_a$ ) at a given temperature ( $T_1$ ).



#### Effect of Increasing Temperature (to $T_2$ ):

The entire curve shifts to the right and flattens, meaning the **average particle energy increases**. Crucially, the area under the curve beyond the activation energy ( $E_a$ ) **significantly increases**. This means a **much larger proportion of particles** now possess the required energy to react upon collision.



**Conclusion:** An increase in temperature doesn't just cause more collisions; it dramatically increases the number of **effective collisions** (those with energy  $\geq E_a$ ), which is the primary reason for the faster reaction rate.

#### 5.3.4. Enzymes

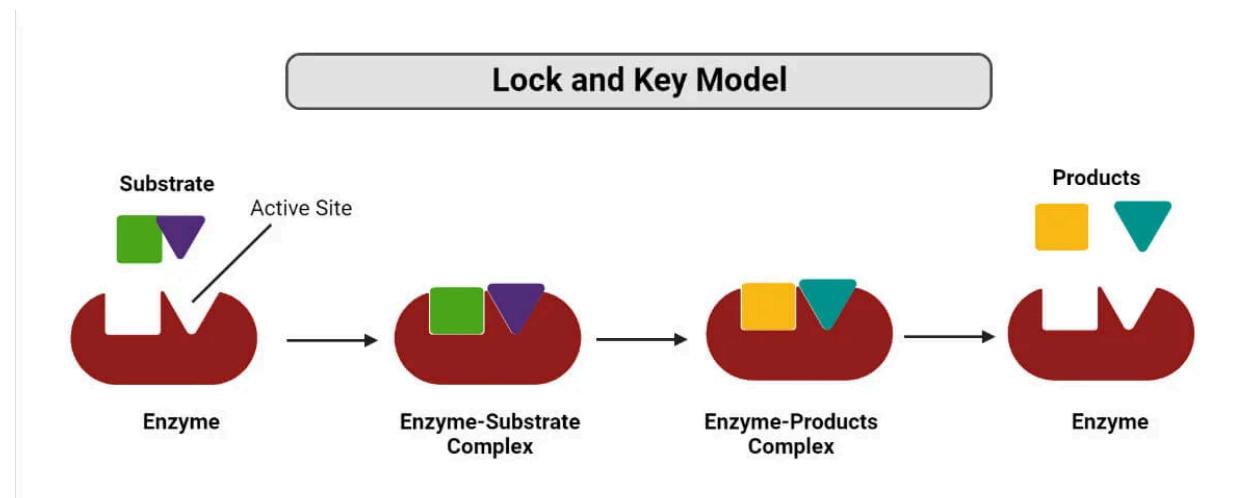
Enzymes are **biochemical catalysts** that regulate the vast majority of chemical reactions within living organisms. **Nature:** They are specialized **proteins**.

**Properties:**

- **Not Consumed:** Like catalysts, they are not used up in the reactions they facilitate.
- **Highly Specific:** Each enzyme catalyzes **only one specific reaction**.
- **Extremely Efficient:** They can speed up reactions by a factor as high as **10<sup>20</sup>**.
- **Location:** Found inside cells and in extracellular fluids (e.g., saliva, gastric juice).

**How They Work (Lock and Key Model):**

Enzymes have a specific three-dimensional shape. They bind to reactant molecules (substrates) and hold them in the **precise orientation** required for a successful collision. This precise binding dramatically **lowers the activation energy** and increases the reaction rate.

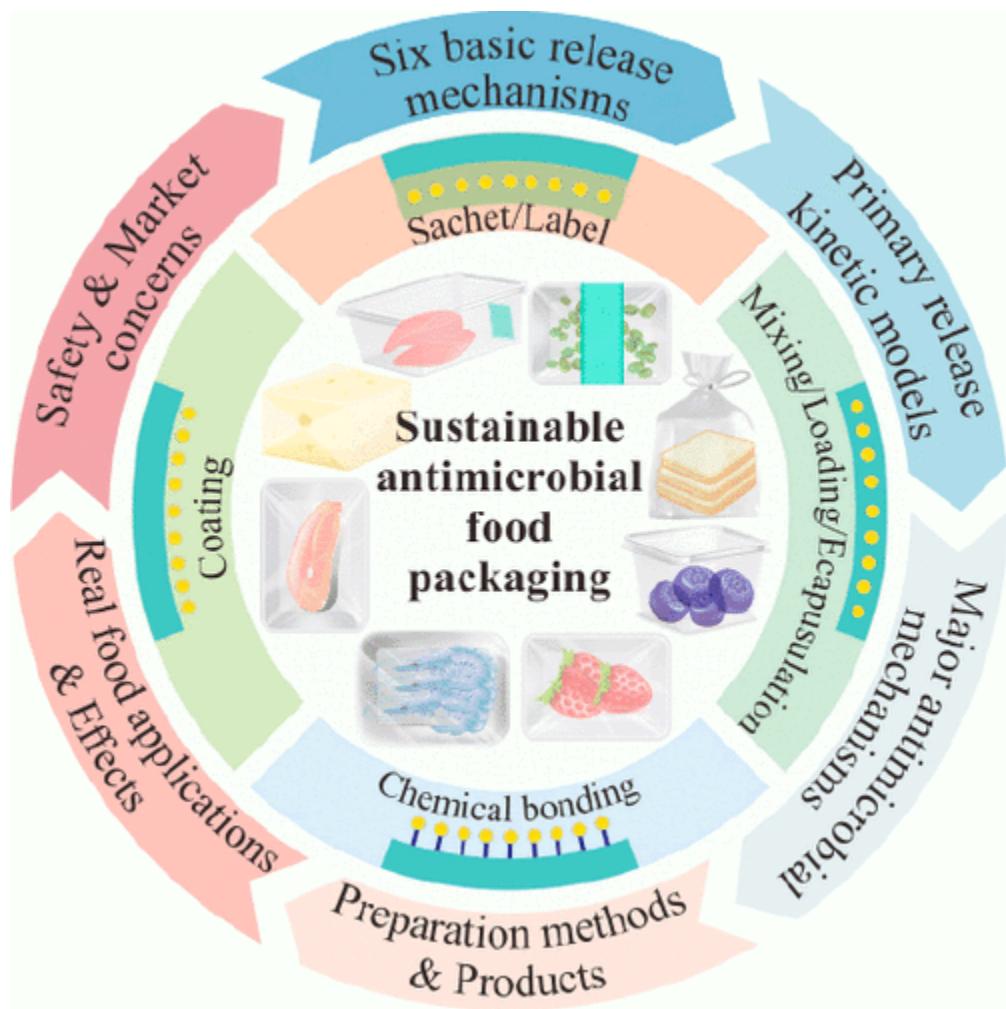


## 5.4. Role of Chemical Kinetics in the Food Industry

Chemical kinetics is applied in the food industry to control quality, minimize waste, and extend shelf life. Key applications include:

**Optimizing Harvest and Transport:** Determining the best time to harvest and transport produce to ensure it arrives with optimal taste, texture, and nutritional value.

**Minimizing Losses:** Estimating harvest time so products reach the market at peak quality, reducing losses from over-ripening during transit.



**Understanding Degradation:** Identifying factors (like oxidation) that cause food spoilage during transportation and storage.

**Improving Storage Methods:** Developing storage and transportation conditions (e.g., controlled atmospheres) that preserve nutritional content.

**Shelf Life:** Using kinetics to find methods (like refrigeration or modified packaging) that **slow down degradation reactions** by controlling temperature and humidity.



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