# N11 - KINETICS

# Collision Theory and More

# N11 - KINETICS

Target: I can use Collision Theory to describe how factors like temperature, concentration, and catalysts affect the speed of a reaction.

#### Rate Laws

#### **Reaction Mechanism**

The series of elementary steps by which a chemical reaction occurs.

#### **Collision Model**

**Key Idea:** Molecules must collide to react.

However, only a small fraction of collisions produces a reaction. Why?



### **Collision Model**

Collisions must have <u>sufficient energy</u> to produce the reaction (must equal or exceed the activation energy).

Colliding particles must be <u>correctly oriented</u> to one another in order to produce a reaction.

#### **Molecularity**

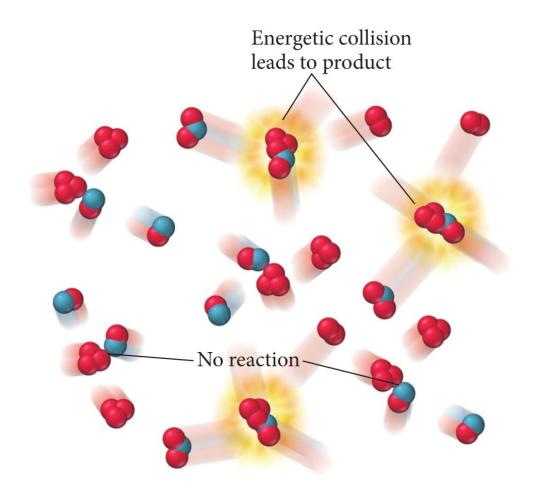
The # of species that must collide to produce the rxn indicated by that step

- Unimolecular step a reaction involving one molecule
- Bimolecular step reaction involving the collisions of two species
- Termolecular step reaction involving the collisions of three species

Elementary Step	Molecularity	Rate Law
$A \rightarrow products$	Unimolecular	Rate = k[A]
$A + A \rightarrow products$ (2A $\rightarrow products$ )	Bimolecular	Rate = k[A] <sup>2</sup>
$A + B \rightarrow products$		Rate = k[A][B]
$A + A + B \rightarrow products$ (2A + B $\rightarrow products$ )	Termolecular	Rate = $k[A]^2[B]$
$A + B + C \rightarrow products$		Rate = $k[A][B][C]$

### **Effective Collisions- Kinetic Energy Factor**

For a collision to lead to overcoming the energy barrier, the reacting molecules must have sufficient kinetic energy so that when they collide the activated complex can form.



#### **Effective Collisions - Orientation Factor**

$$\operatorname{NOCl}(g) + \operatorname{NOCl}(g) \longrightarrow 2 \operatorname{NO}(g) + \operatorname{Cl}_2(g)$$
 $+ \longrightarrow + \longrightarrow + \longrightarrow$ 
Ineffective collision

Effective collision

### <u>Molecular Interpretation of Factors</u> <u>Affecting the Rate – Reactant Nature</u>

- Reactions generally occur faster in solution than in pure substances.
  - Mixing gives more particle contact.
  - Particles are separated, allowing more effective collisions per second.
  - Forming some solutions breaks bonds that need to be broken.

# Molecular Interpretation of Factors Affecting the Rate – Reactant Nature

## Some materials undergo similar reactions at different rates either because they have a

- 1) Higher initial potential energy and are therefore closer in energy to the activated complex, or
- 2) Because their reaction has a lower activation energy.

CH<sub>4</sub> + Cl<sub>2</sub> → CH<sub>3</sub>Cl + HCl is about 12 times faster than

 $CD_4 + CI_2 \rightarrow CD_3CI + DCI$  because the C—H bond is weaker and less stable than the C—D bond.

 $CH_4 + X_2 \rightarrow CH_3X + HX$  occurs about 100x faster with  $F_2$  than  $CI_2$  because the  $A_E$  for  $F_2$  is 5 kJ/mol, but for  $CI_2$  is 17 kJ/mol.

# Molecular Interpretation of Factors Affecting the Rate – Concentration

- Reaction rate generally increases as the concentration or partial pressure of reactant molecules increases.
  - Except for zero order reactions
- More molecules leads to more molecules with sufficient kinetic energy for effective collision.
  - Distribution the same, just bigger curve

# Molecular Interpretation of Factors Affecting the Rate – Temperature

- Increasing the temperature raises the average kinetic energy of the reactant molecules.
- There is a minimum amount of kinetic energy needed for the collision to be converted into enough potential energy to form the activated complex.
- Increasing the temperature increases the number of molecules with sufficient kinetic energy to overcome the activation energy.

#### **Effect of Temperature on Rate**

- Changing the temperature changes the rate constant of the rate law.
- Svante Arrhenius investigated this relationship and showed the following:  $k = Ae^{\left(\frac{-Ea}{RT}\right)}$

*T* = temperature in kelvins.

R = gas constant in energy units, 8.314 J/(mol - K).

**A** = called the frequency factor, the rate the reactant energy approaches the Ea

 $E_a$  = activation energy, the extra energy needed to start the molecules reacting.

### The Arrhenius Equation Rearranged

$$\ln(k) = -\frac{E_a}{R} \left(\frac{1}{T}\right) + \ln(A)$$

- Simplifies solving for E<sub>a</sub>
- $-E_a/R$  is the slope when graphing ln(k) vs. (1/T)
- In(A) is the y-intercept
- Graphing In(k) vs (1/T) and taking line of best fit can quickly yield a slope

### <u>Arrhenius Equation – Exponential Factor</u>

- The exponential factor in the Arrhenius equation is a number between 0 and 1.
- $k = Ae^{\left(\frac{-Ea}{RT}\right)}$
- Represents the fraction of reactant molecules with sufficient energy so they can make it over the energy barrier.
  - The higher the energy barrier (larger E<sub>A</sub>), the fewer molecules that have sufficient energy to overcome it.
- That extra energy comes from converting the kinetic energy of motion to potential energy in the molecule when the molecules collide.
  - Increasing the temp increases the average kinetic energy of the molecules.
  - Therefore, increasing the temp will increase the number of molecules with sufficient energy to overcome the energy barrier.
  - Therefore, increasing the temperature will increase the reaction rate.

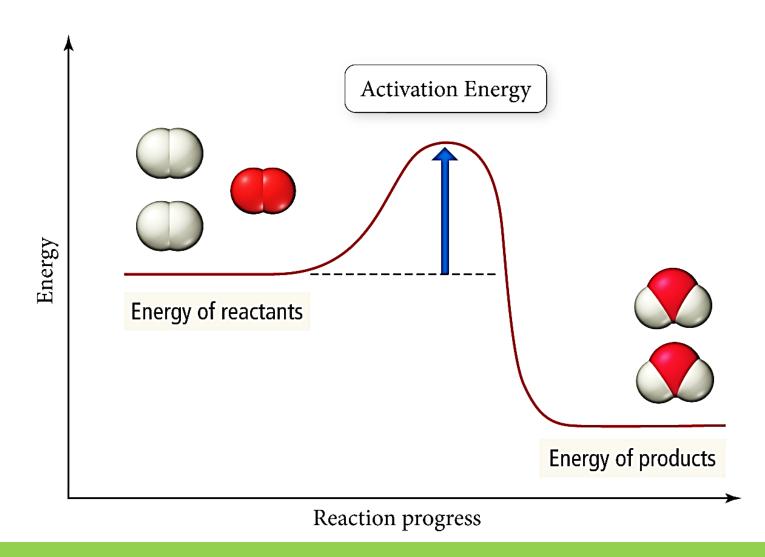
#### **Arrhenius Equation – Two Point Format**

If you only have two (T, k) data points, the following form of the Arrhenius equation can be used:

$$ln\left(\frac{k_2}{k_1}\right) = \frac{E_a}{R} \left(\frac{1}{T_1} - \frac{1}{T_2}\right)$$

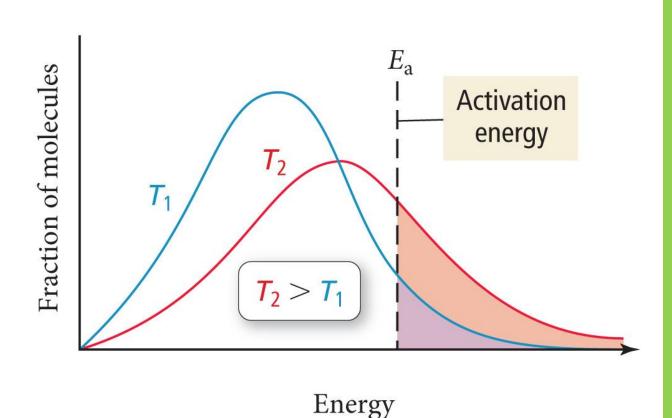
#### **Activation Energy**

$$2 H_2(g) + O_2(g) \rightleftharpoons 2 H_2O(g)$$



#### **Thermal Energy Distribution**

As temperature increases, the fraction of molecules with enough energy to surmount the activation energy barrier also increases.



### **Catalysts**

Catalyst - A substance that speeds up a reaction without being consumed during the reaction.

Enzyme - A large molecule (usually a protein) that catalyzes biological reactions.

Homogeneous catalyst - Present in the same phase as the reacting molecules.

Heterogeneous catalyst - Present in a different phase than the reacting molecules.

### **Catalysts**

- Catalysts work by providing an alternative mechanism for the reaction with a lower activation energy.
- Catalysts are consumed in an early mechanism step, and then re-made in a later step.

#### **Mechanism without catalyst:**

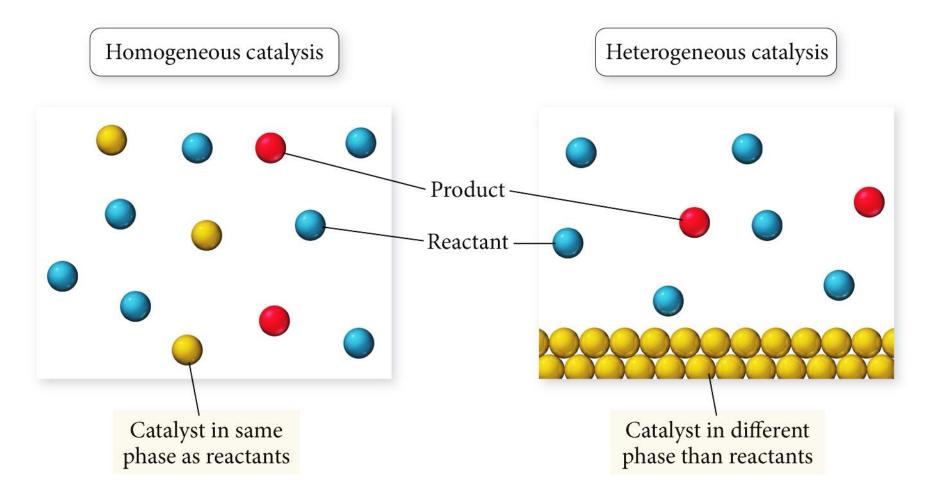
$$O_{3(g)} + O_{(g)} \rightarrow 2 O_{2(g)}$$
 Very Slow

#### **Mechanism with catalyst:**

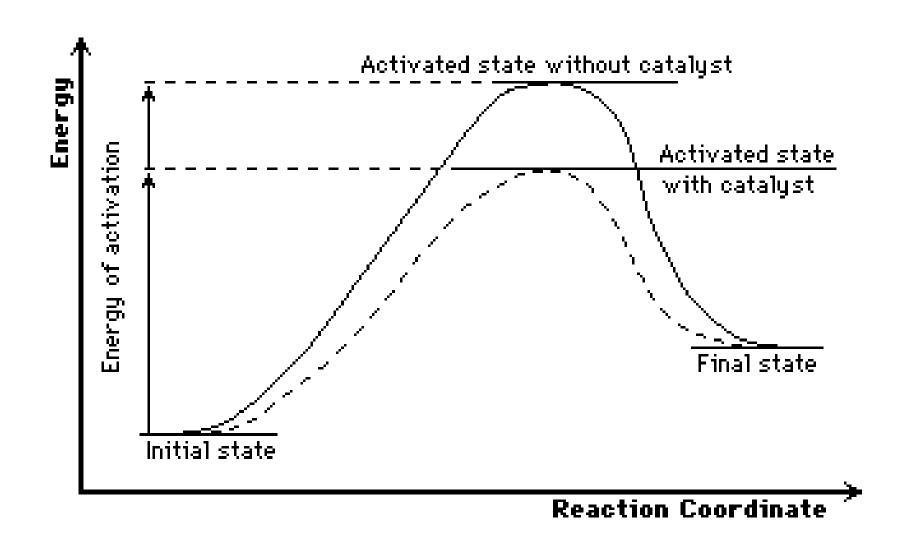
$$CI_{(g)} + O_{3(g)} \Leftrightarrow O_{2(g)} + CIO_{(g)}$$
 Fast

$$ClO_{(g)} + O_{(g)} \rightarrow O_{2(g)} + Cl_{(g)}$$
 Slow

### **Types of Catalysts**



## **Lowering of E<sub>A</sub> by a Catalyst**

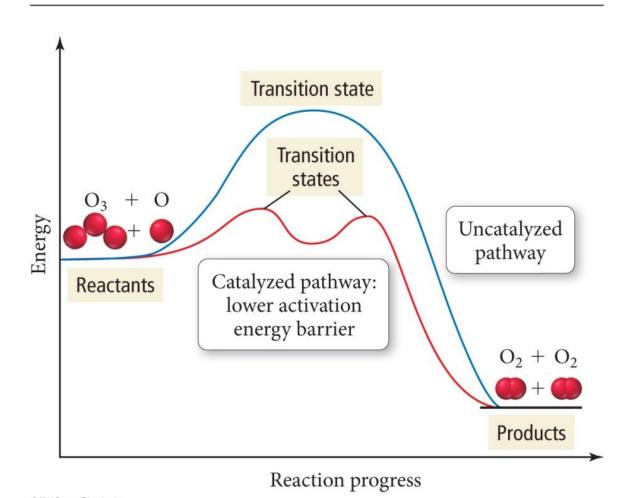


#### **Energy Profile of a Catalyzed Reaction**

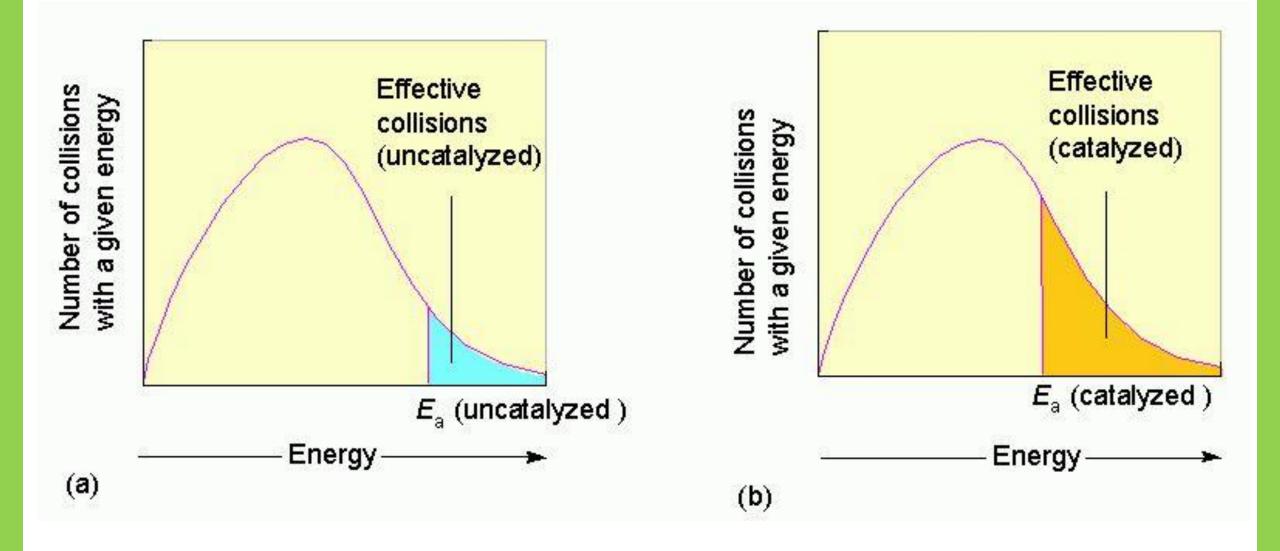


Polar stratospheric clouds contain ice crystals that catalyze reactions that release Chlorine from atmospheric chemicals.

Energy Diagram for Catalyzed and Uncatalyzed Pathways



#### Catalysts Increase the # of Effective Collisions



# Molecular Interpretation of Factors Affecting the Rate – Catalysts

Give reactant molecules a different path to follow w/ lower E<sub>A</sub>.

- Homogeneous catalysts react with one of the reactant molecules to form a more stable activated complex with a lower activation energy.
- Heterogeneous catalysts hold one reactant molecule in proper orientation for reaction to occur when the collision takes place.
  - Sometimes they also help to start breaking bonds.

#### **Step #1:**

Adsorption and activation of the reactants.

Carbon monoxide and nitrogen monoxide adsorbed on a platinum surface **Platinum** 

#### **Step #2:**

Migration of the adsorbed reactants on the surface.

Carbon monoxide and nitrogen monoxide arranged prior to reacting



#### **Step #3:**

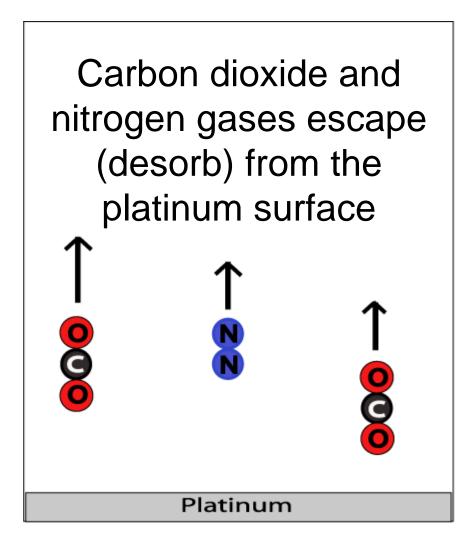
Reaction of the adsorbed substances.

Carbon dioxide and nitrogen form from previous molecules



#### **Step #4:**

Escape, or desorption, of the products.



# YouTube Link to Presentation https://youtu.be/Ogol25FK9\_M