

# Reaction Kinetics

**A measure of the rate at  
which a reaction occurs**



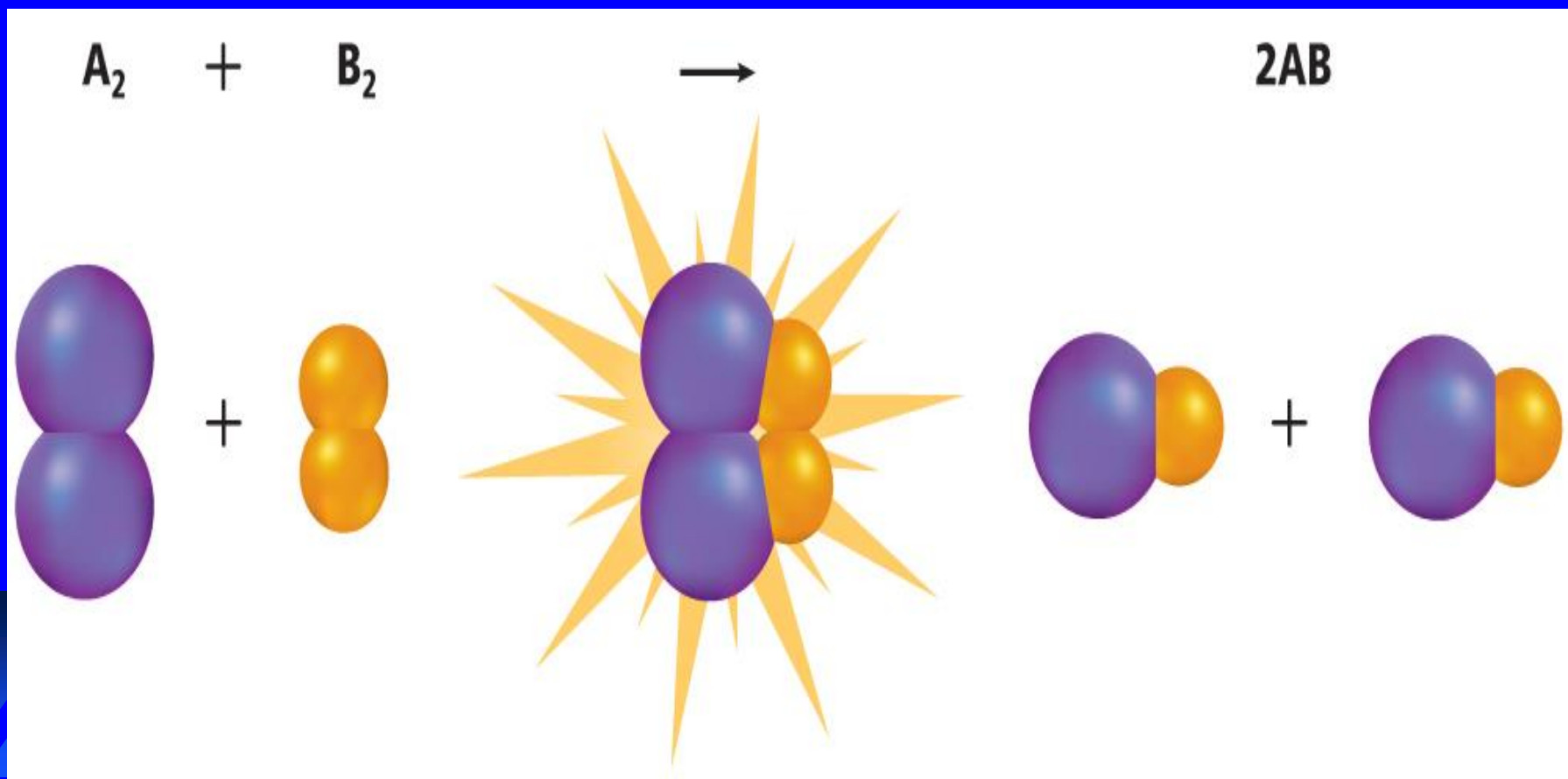
# Kinetic Energy

- Related to molecular speed
- $KE = \frac{1}{2}mv^2$ 
  - higher velocity = higher KE

# Reaction Rate

- The rate of reaction tells us the **speed** at which a reaction takes place

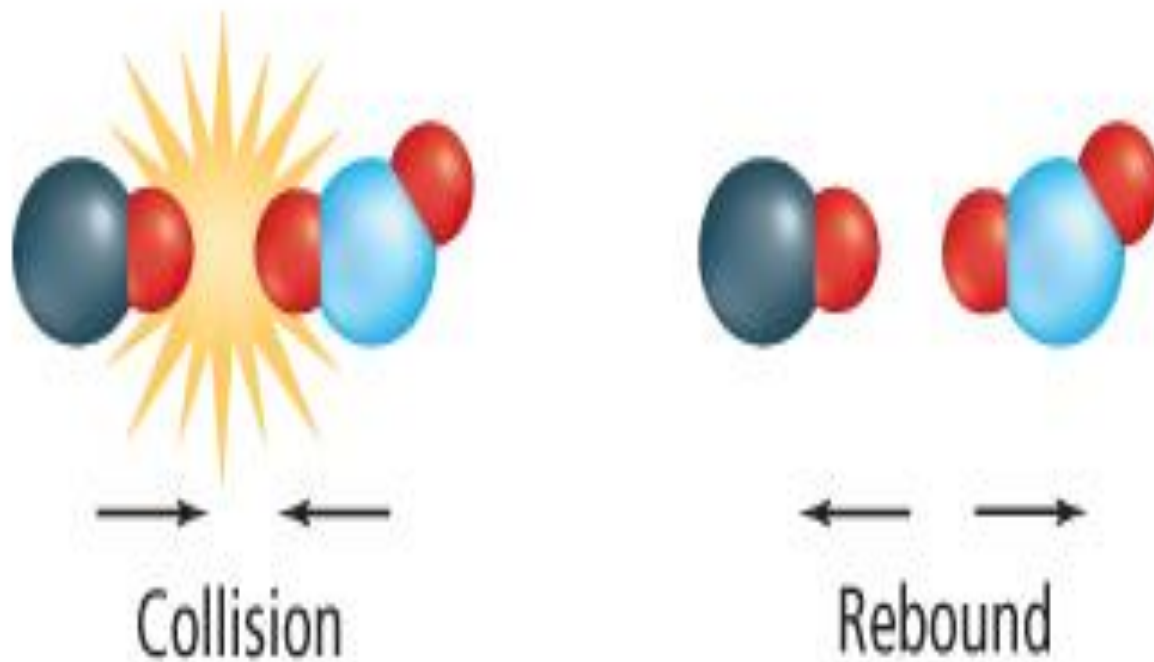
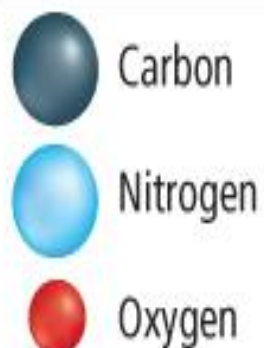
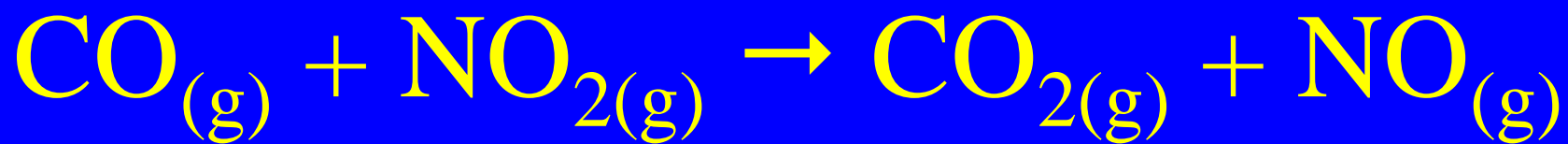
**Collision Theory** states that atoms, ions, and molecules must collide in order to react.



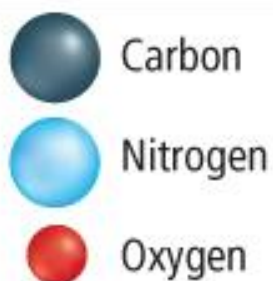
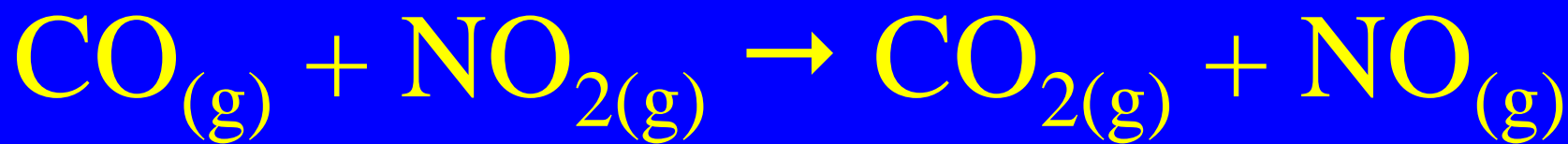
# Collision Theory

**Not every collision between atoms or molecules results in a reaction**

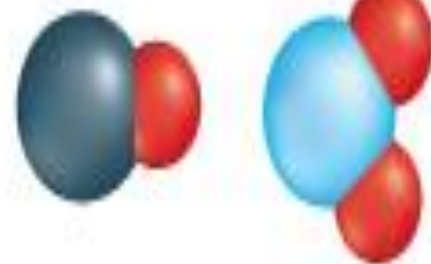
- 1. Orientation is crucial**  
ex: darts, train cars



**Incorrect orientation**

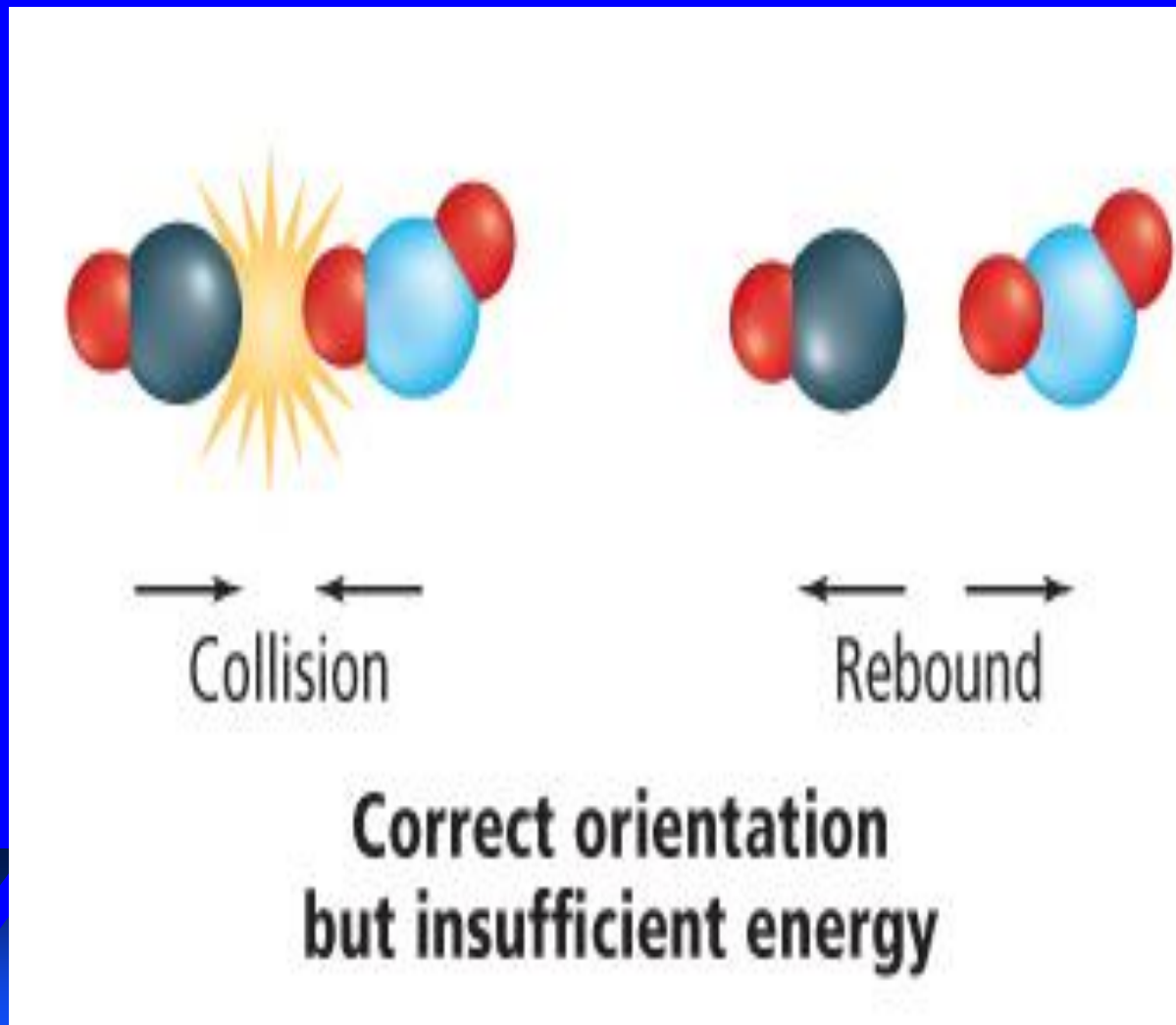
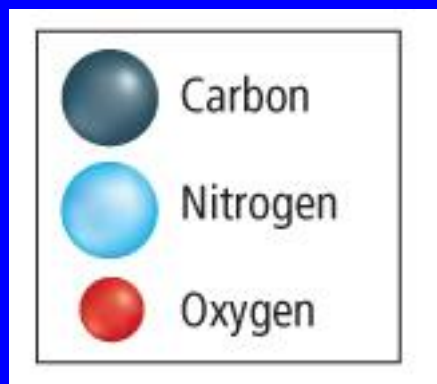
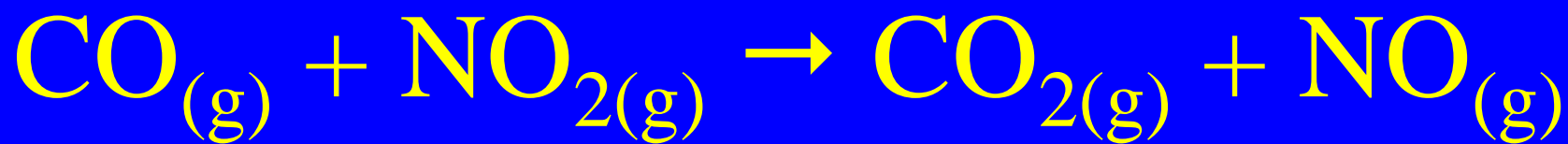


Collision

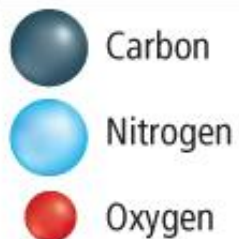
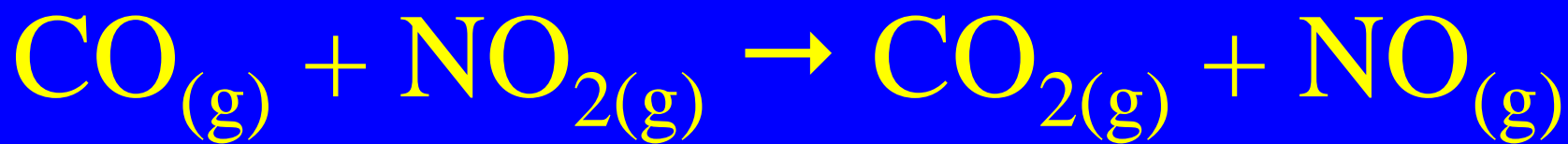


Rebound

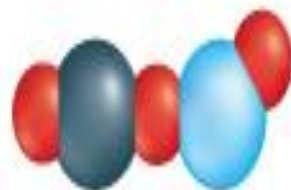
**Incorrect orientation**



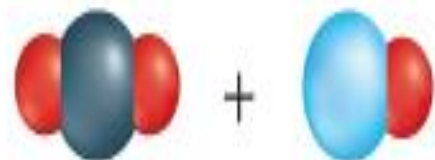




→ ←  
Collision



Activated complex



Products

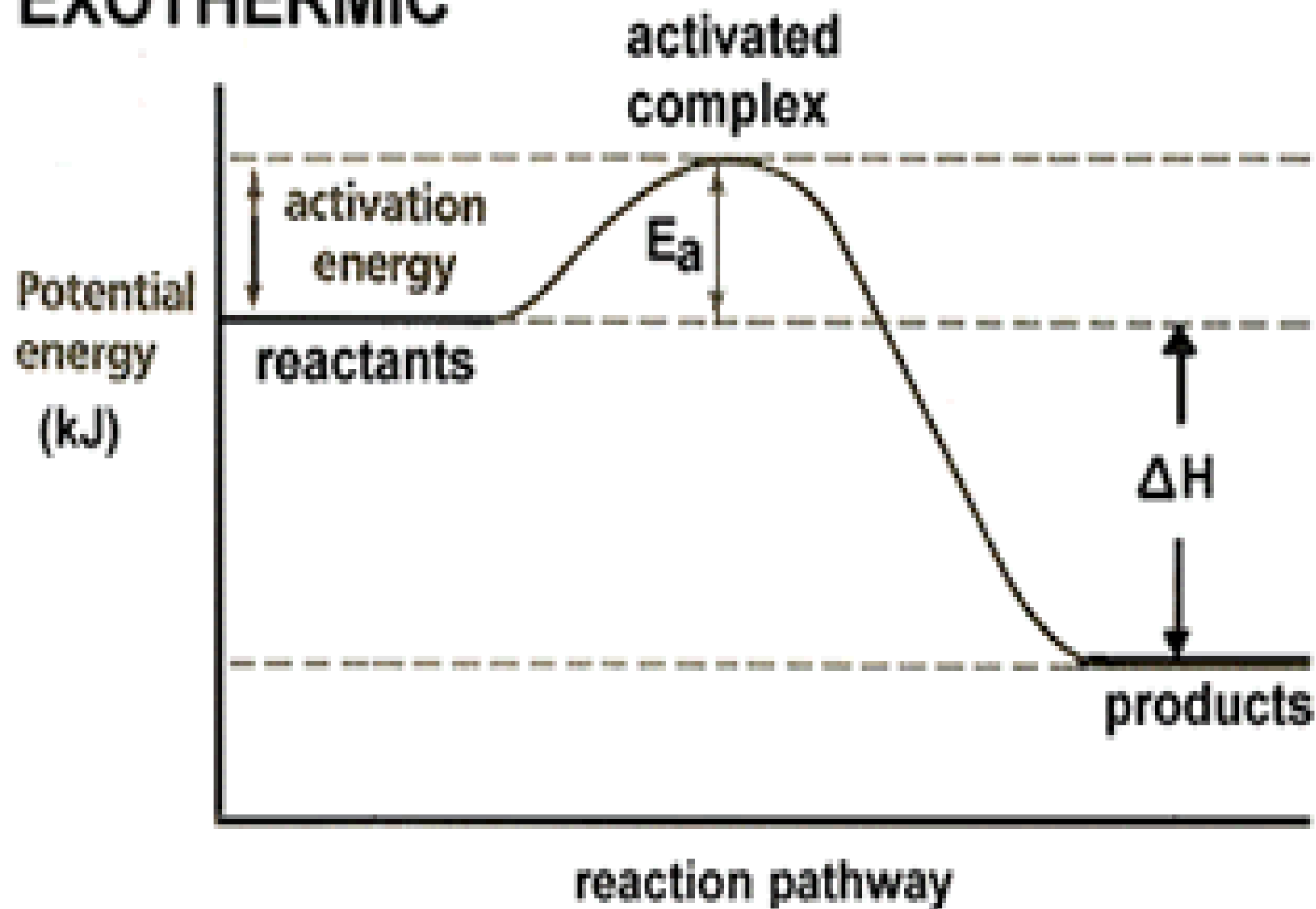
**Correct orientation**

# Collision Theory

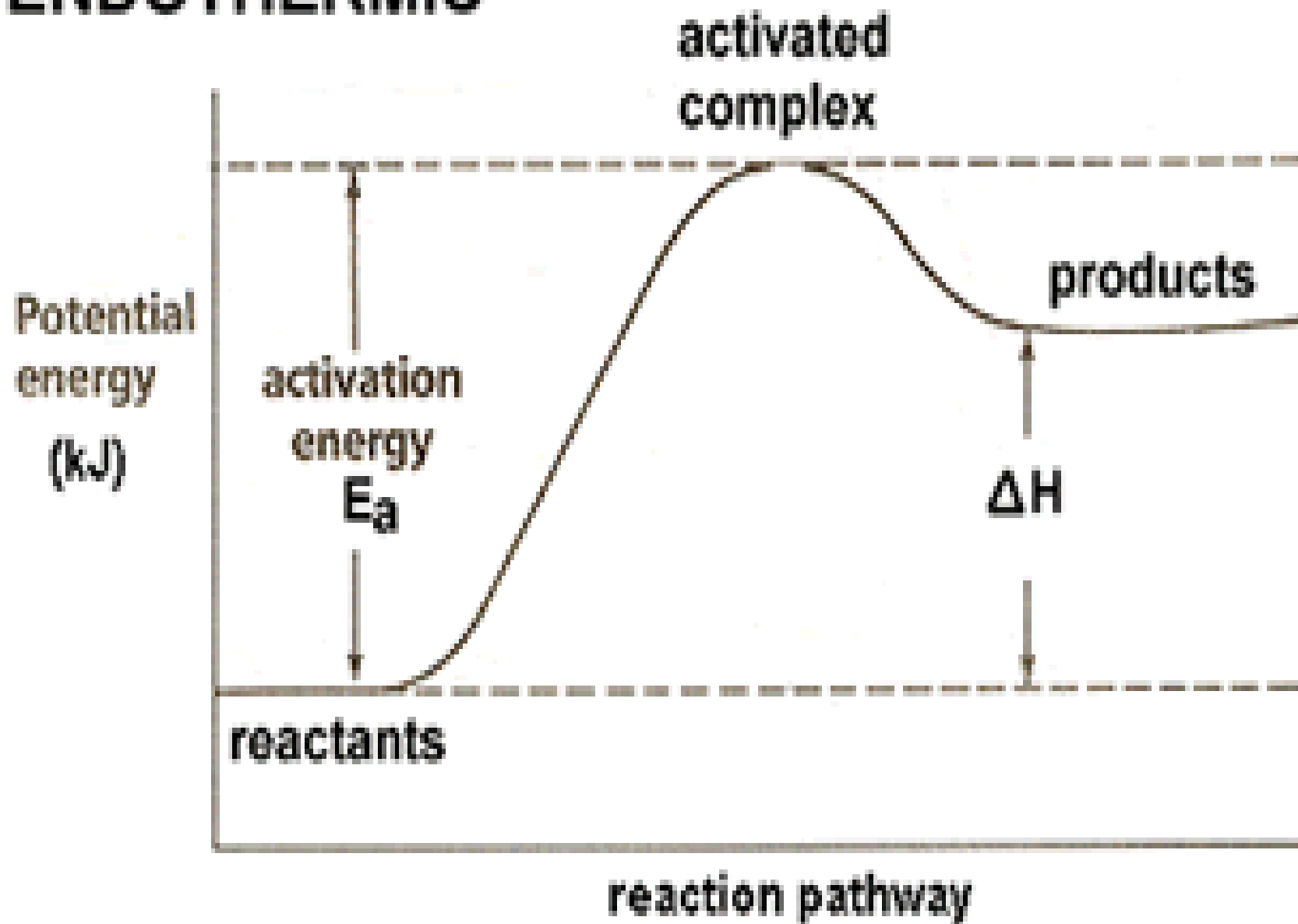
**Not every collision between atoms or molecules results in a reaction**

2. They must collide with sufficient energy
  - the “**Activation energy**” ( $E_a$ )

# EXOTHERMIC



# ENDOTHERMIC



# Collision Theory

Any change that increases the number of collisions should increase reaction rate

# Factors in reaction rates...



# 1. The Nature of the reactants

- what are their “reactivities” / tendencies toward bond formation?
- $\text{Cs} + \text{H}_2\text{O} \rightarrow \text{CsOH} + \text{H}_2$ 
  - instantaneous
  - $\text{Cs} \Rightarrow$  very reactive (low IE, EN)
- $\text{Fe} + \text{H}_2\text{O} \text{-----} > \text{Fe}_2\text{O}_3 + \text{H}_2$ 
  - very slow

## 2. The ability of the reactants to meet (collide)

- reactions usually occur in liquid, gas or aqueous phase
- Related to **surface area** -  
higher surface area means faster reaction in heterogeneous reactions
  - reaction only occurs at phase interface
  - ex: log vs firewood vs kindling vs sawdust



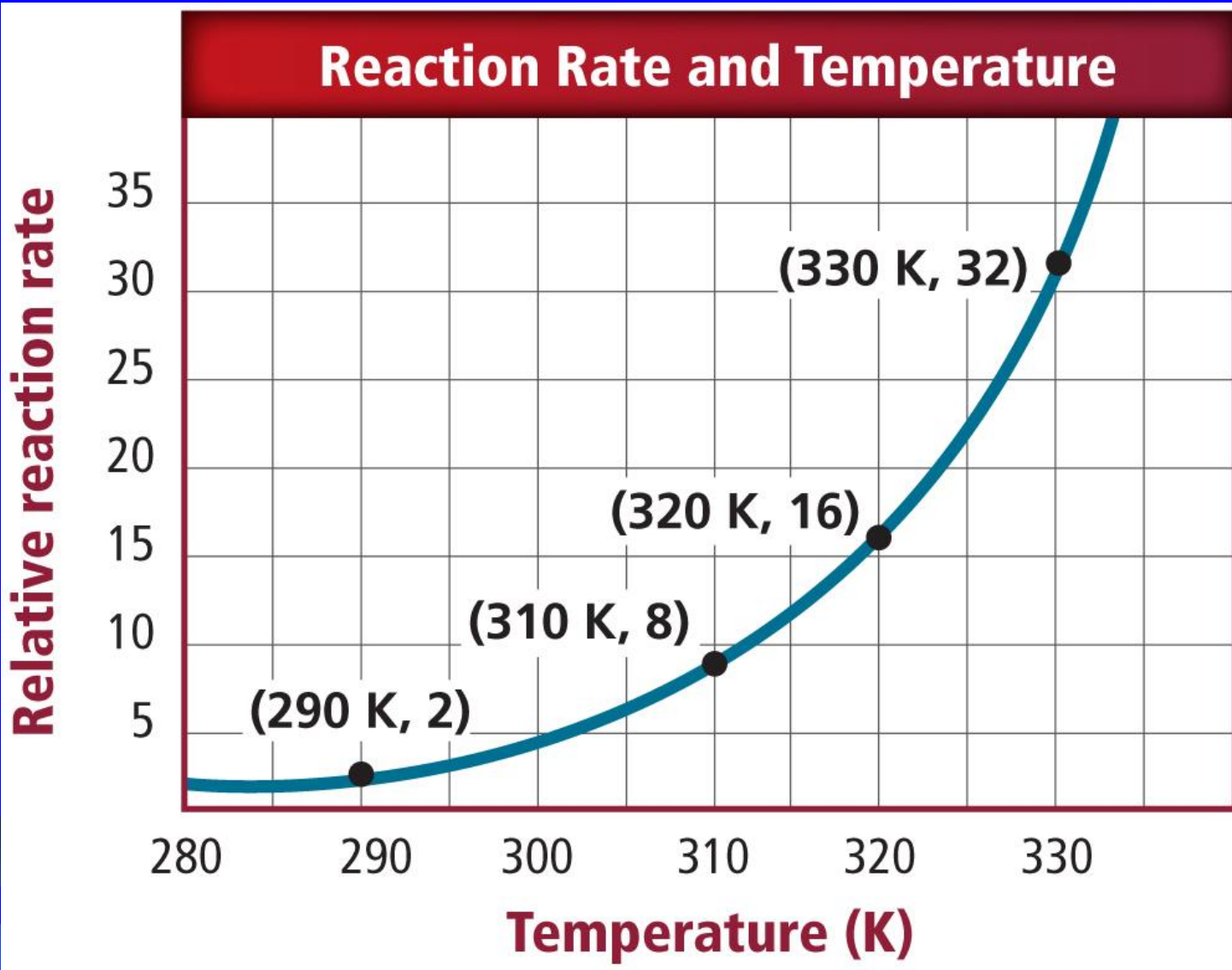
### 3. Concentration of the reactants

- often listed as molarity (mol/L)
- written as [square brackets]
  - ex:  $[\text{HCl}]$  = “the M of the HCl”
- more reactants = more collisions
- more collisions = increased reaction rate

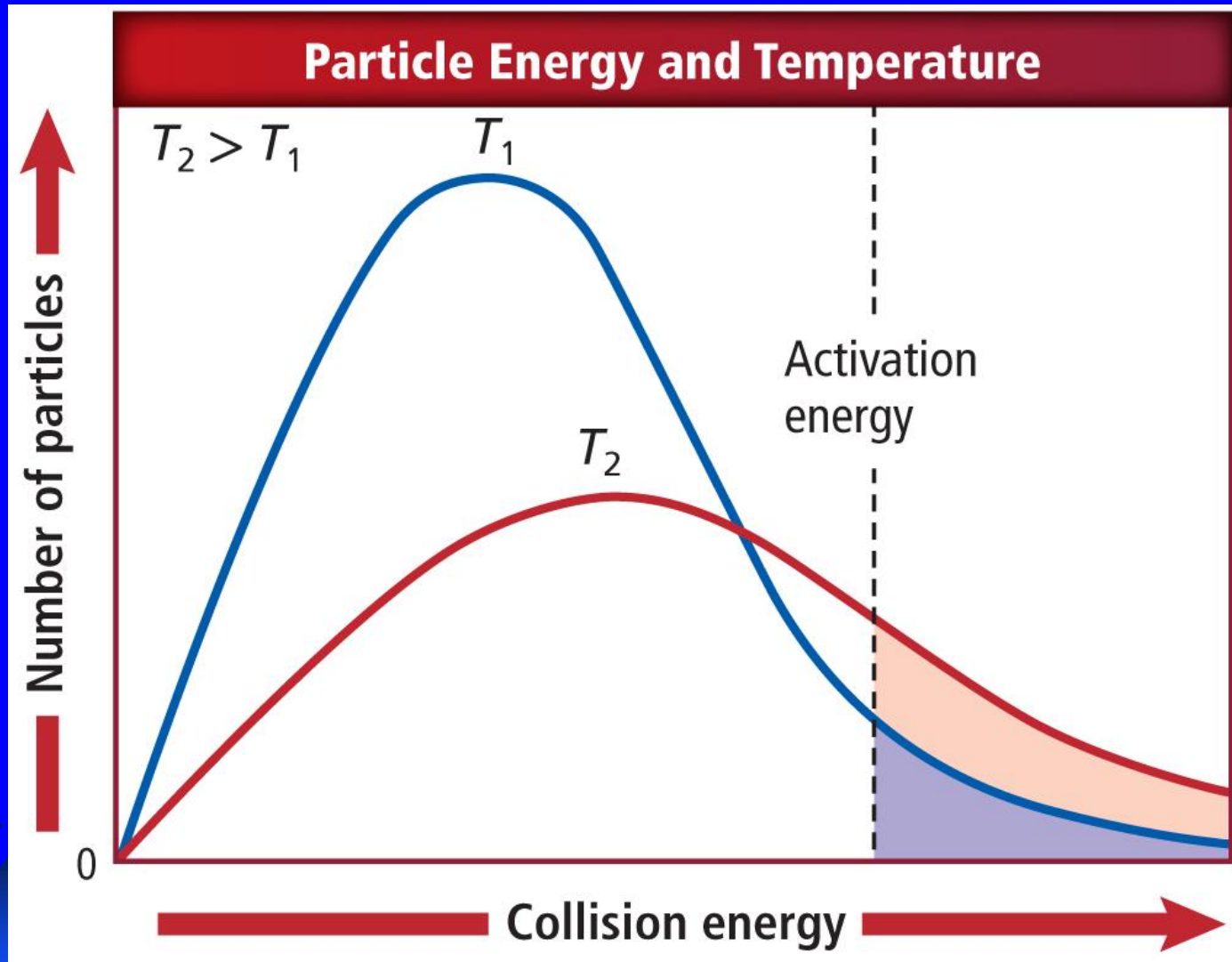
# 4. Temperature

- T is a measure of the average KE
- higher T means higher KE
- higher KE = molecules moving faster
- faster moving molecules means more collisions
- more collisions = faster rate


# 4. Temperature



# 4. Temperature

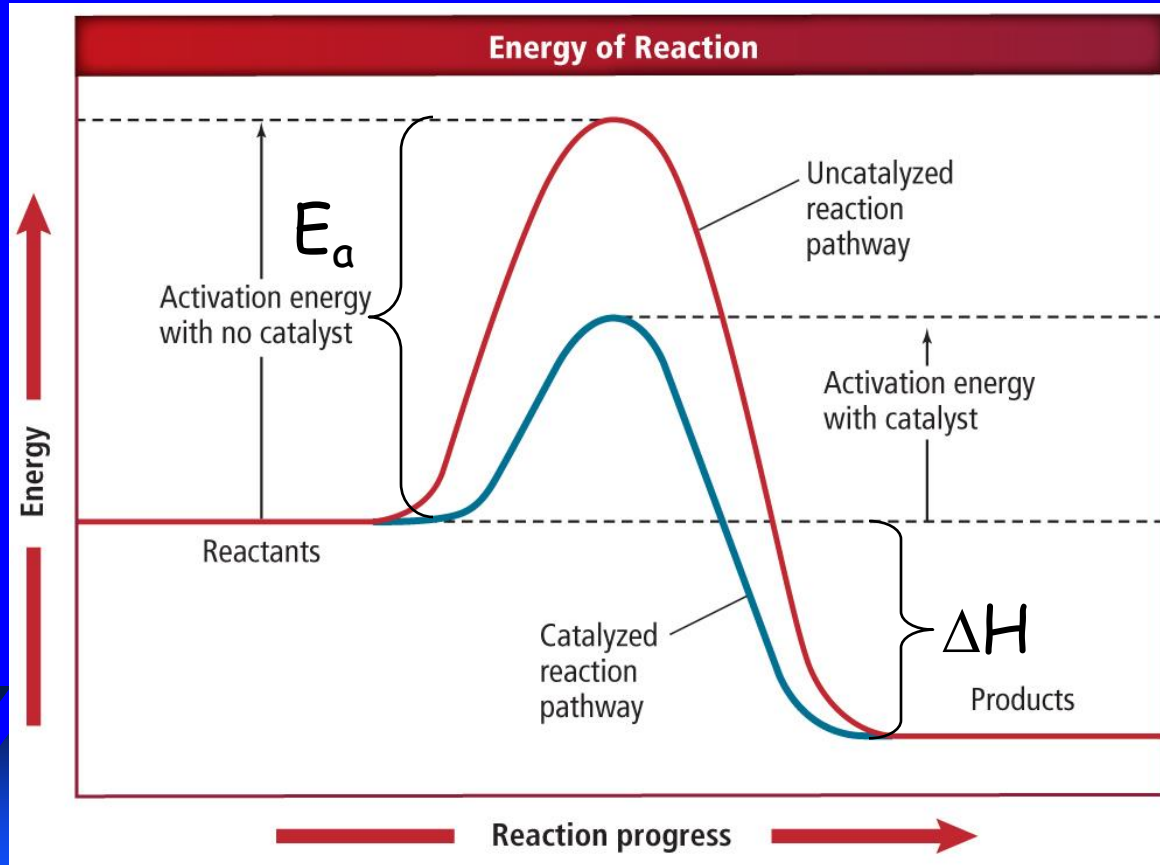


# 5. Catalysts

- increase the reaction rate without being consumed in the process
  - lower the energy barrier needed to overcome in order to react ( $E_a$ )
- 

# 5. Catalysts

- Lower activation energy means more collisions between particles have sufficient energy to react.



# 5. Catalysts

- A **heterogeneous catalyst** exists in a physical state different than that of the reaction it catalyzes.
- A **homogeneous catalyst** exists in the same physical state as the reaction it catalyzes.
- A **reforming catalyst** is consumed in the reaction, but then re-made
- An **enzyme** is a biologically active catalyst
- An **inhibitor** slows down a reaction by increasing the activation energy

***Next:  
a look inside  
the  
numbers....***






# Reaction Kinetics

A measure of the rate at which a  
reaction occurs



# To Review....Five factors that change reaction rate

- 1. The nature of the reactants**
  - 2. the ability of the reactants to collide**
  - 3. concentration of reactants**
  - 4. temperature of the system**
  - 5. catalysts**
- 

# Measuring Rates

- A rate is something per unit of time
  - interest rate = \$ earned / time
  - speed = distance traveled/ time
  - pay = dollars / hour

# Measuring Reaction Rates

- as the reaction proceeds, the reactants are “used up”
  - [reactants] goes down
  - [products] goes up
- Reaction rate is measured as :  
 $\Delta \text{ concentration} \div \Delta \text{ time}$

◆ *rate unit:*  $\frac{\Delta \text{concentration}}{\Delta \text{time}} = \frac{M}{s} = \frac{\text{mol/L}}{s} =$   
 $M s^{-1} = \text{mol } L^{-1} s^{-1}$

# Coefficients and rate

- The coefficients of a chemical equation give us some insight as to the relative rates of consumption of reactants and production of products, but not “absolute” amounts.

Imagine a machine where you put in 1 dime, 2 nickels, and 5 pennies in a slot on the top, and a quarter drops out the other side

$$1 D + 2 N + 5 P = 1 Q$$

If you had a pile of 100 dimes, a pile of 100 nickels, and a pile of 100 pennies, are the piles being used up at the same rate?

$$1D + 2N + 5P = 1Q$$

How does the “rate of disappearance” of pennies compare to the “rate of appearance” of quarters?

If the nickels are disappearing at a rate of 3 per second, how fast are the pennies disappearing?

$$3 \text{ N s}^{-1} \times 5 \text{ P} / 2 \text{ N} = 7.5 \text{ P s}^{-1}$$

consider the reaction



- it takes 2 A's for every single B that reacts, so A is “disappearing” twice as fast as B is



consider the reaction



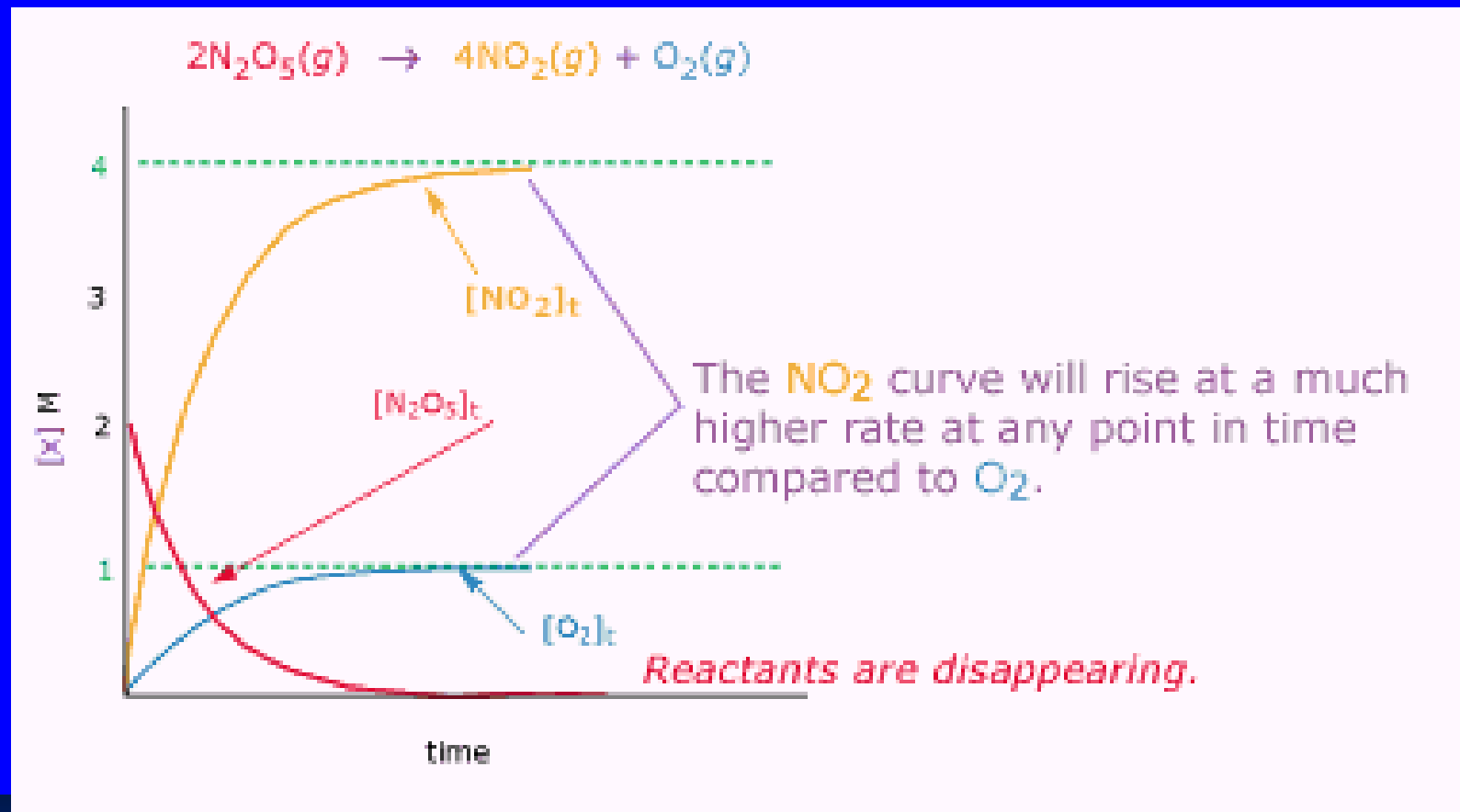
- there are 2 D's produced for every B and every 2 A's that are "consumed", so D is appearing at the same rate A is disappearing and twice as fast as B is disappearing

consider the reaction



- C is being produced 3 times faster than B is being used up, and  $3/2$  times faster than A is being used up

# Graphing the rate of $\Delta M$ for



A Rate Law is an equation that describes how a change in concentration affects the reaction rate.

- for the reaction:  $A + B \rightarrow \text{products}$
- The rate law would be:

$$\text{rate} = k[A]^m[B]^n$$

$$\text{rate} = k[A]^m[B]^n$$

- **k** = the rate constant → depends on the **temperature**, different for each reaction
- **m** = the “*order of reaction*” with respect to A
- **n** = the “*order of reaction*” with respect to B

$$\text{rate} = k[A]^m[B]^n$$

- **m** and **n** have to be experimentally determined; they are not the same as the reaction coefficients except in a “one step mechanism” (pretty rare).

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