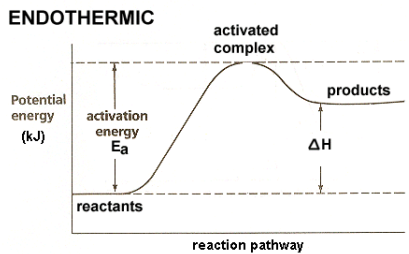
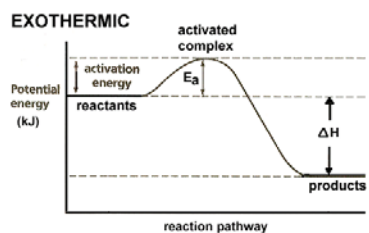


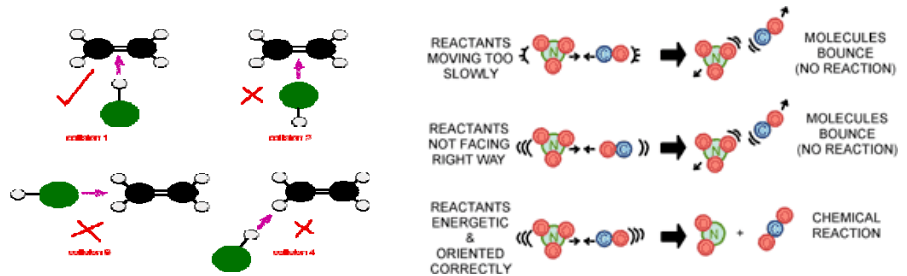
## Kinetics/Equilibrium/Thermochemistry



1

## Kinetics

- study of the rates of chemical \_\_\_\_\_
- Collision theory: in order for a reaction to occur, particles must \_\_\_\_\_ with proper \_\_\_\_\_ and \_\_\_\_\_



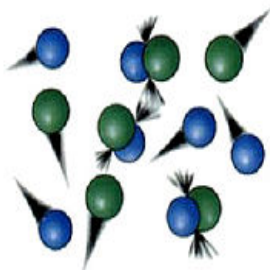
## Factors affecting rate of reaction

- Type of reactant (covalent vs. ionic)
  - Covalent
    - \_\_\_\_\_ to break bond
    - \_\_\_\_\_ energy required
    - \_\_\_\_\_ bonds to break
    - Reacts \_\_\_\_\_
  - Ionic
    - \_\_\_\_\_ to break bond
    - \_\_\_\_\_ energy needed
    - \_\_\_\_\_ bonds to break
    - Reacts \_\_\_\_\_

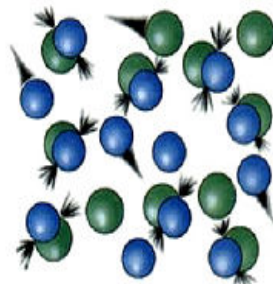
3

## Factors affecting rate of reaction

- Concentration: \_\_\_ in amount of reactants will \_\_\_ rate of rxn (more \_\_\_\_\_ occurring when more is present)



Low concentration = Few collisions



High concentration = More collisions

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## Factors affecting rate of reaction

- Surface Area: \_\_\_ in surface area will \_\_\_ rate of rxn
  - Ex. clump of sugar; inner portions are \_\_\_ making \_\_\_ with water (the other reactant)
  - Rxn is \_\_\_ than if material was crushed

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## Factors affecting rate of reaction

- Pressure: Specifically affects gases
  - \_\_\_ in pressure will \_\_\_ amount of collisions
  - In a closed system a volume \_\_\_ will lead to a pressure \_\_\_ and rxn rate \_\_\_
- Temperature: \_\_\_ in temp will \_\_\_ KE of particles which will result in more energy and more effective collisions
  - As collisions \_\_\_, rxn rate also \_\_\_
  - [Temperature and Movement animation](#)

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## Factors affecting rate of reaction

- Catalyst: Provides an \_\_\_\_\_ for a reaction
  - \_\_\_ rate of rxn
  - Causes activation (\_\_\_\_\_) energy to be \_\_\_\_\_, so rxn may start \_\_\_\_\_
  - remains \_\_\_\_\_ when rxn is complete

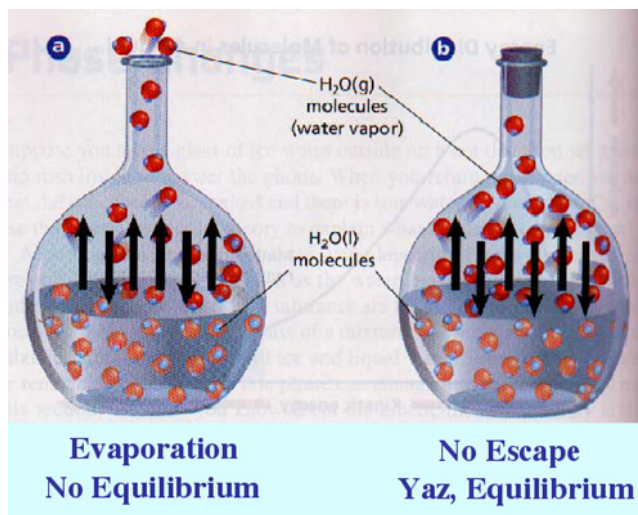
7

## Equilibrium

- when both forward and reverse reactions occur at the \_\_\_\_\_ (in a closed system)
- Ex.  $\text{H}_2\text{O}(\text{l}) \rightleftharpoons \text{H}_2\text{O}(\text{g})$  (represented by double arrow)
- \_\_\_\_\_ of reactants and products do \_\_\_\_\_ have to be equal, but there of \_\_\_\_\_ or \_\_\_\_\_ up are equal
  - Ex. rate of water \_\_\_\_\_ is the same is water vapor \_\_\_\_\_ back to liquid
  - Boiling water  $\rightarrow$  steam rises and condenses back into the pot

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# Equilibrium



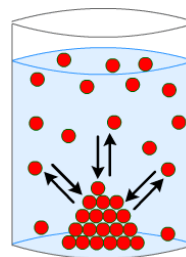
9

## Physical Equilibrium

- Changing of \_\_\_\_\_ occurs
- Phase equilibrium: between solid/liquid or melting/freezing
  - Water will \_\_\_\_\_ and \_\_\_\_\_ at the same rate at 0°C ( $\text{H}_2\text{O(s)} \rightleftharpoons \text{H}_2\text{O(l)}$ )
- Solution Equilibrium: solids in liquids
  - When \_\_\_\_\_ = \_\_\_\_\_ (at a constant temperature)

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## Physical Equilibrium



– Ex. sugar in a saturated tea solution → some may \_\_\_\_\_, but other molecules will \_\_\_\_\_  
\_\_\_\_\_ ( $C_{12}H_{22}O_{11}(s) \rightleftharpoons C_{12}H_{22}O_{11}(aq)$ )

- Can also occur with a gas in a liquid
  - $CO_2$  in soda ( $CO_2(g) \rightleftharpoons CO_2(aq)$ )
  - If temperature \_\_\_\_, more gas will \_\_\_\_\_  
then will be converted/dissolved to aqueous form

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## Chemical Equilibrium

- When reactants are 1st mixed, only the \_\_\_\_\_ reaction will occur
- When reactant amount \_\_\_\_ and product \_\_\_\_ the \_\_\_\_\_ reaction will be favored
- When forward = reverse, \_\_\_\_\_ is achieved

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## Le Chatelier's Principle

- Systems will \_\_\_\_\_ due to changes in \_\_\_\_\_, \_\_\_\_\_ or \_\_\_\_\_ so as to relieve stress on the system
  - May shift forward (to the \_\_\_\_\_)
  - May shift reverse (to the \_\_\_\_\_)

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## Le Chatelier's Principle

- **Changes in Temperature:** ex.  $\text{N}_2 + 3\text{H}_2 \rightleftharpoons 2\text{NH}_3 + \text{Heat}$ 
  - If Heat \_\_\_\_\_, \_\_\_\_\_ rxn favored
    - Easier to \_\_\_\_\_ larger molecules into smaller ones with the addition of heat
    - Typically takes place with \_\_\_\_\_ reactions (heat \_\_\_\_\_)
  - If Heat \_\_\_\_\_, \_\_\_\_\_ rxn favored
    - Easier for smaller reactants to \_\_\_\_\_ and form a \_\_\_\_\_ product
    - Typically takes place with \_\_\_\_\_ reactions (heat \_\_\_\_\_)

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## Le Chatelier's Principle

- **Changes in Concentration:**

- \_\_\_\_\_ of a \_\_\_\_\_ will shift equilibrium to the \_\_\_\_\_
  - More \_\_\_\_\_ is formed
  - \_\_\_\_\_ rxn favored over \_\_\_\_\_
  - \_\_\_\_\_ will eventually be met
- \_\_\_\_\_ of a reactant will shift equilibrium to the \_\_\_\_\_
  - Less product can be formed
  - \_\_\_\_\_ rxn favored over \_\_\_\_\_
  - \_\_\_\_\_
  - Equilibrium is eventually met

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## Concentration Shift

- Ex.  $4\text{NH}_3(g) + 5\text{O}_2(g) \rightleftharpoons 4\text{NO}(g) + 6\text{H}_2\text{O}(g) + \text{Heat}$
- If \_\_\_\_\_(g) is removed from the \_\_\_\_\_, NO(g) and H<sub>2</sub>O(g) would be impossible to \_\_\_\_\_

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## Le Chatelier's Principle

- **Changes in Pressure:** (typically with gases)
  - When only 1 gas present on both sides of a reaction;  $\text{CO}_2(g) \rightleftharpoons \text{CO}_2(aq)$ 
    - Pressure \_\_\_ will favor the \_\_\_\_\_ rxn
    - More  $\text{CO}_2(g)$  going \_\_\_\_\_ solution  $\rightarrow$  becoming  $\text{CO}_2(aq)$
    - Pressure \_\_\_ will favor the \_\_\_\_\_ rxn
    - $\text{CO}_2(g)$  will come \_\_\_\_\_ solution (when you open a soda can)

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When # of gas molecules/moles is different on both sides of the reaction

- ex.  $\text{N}_2(g) + 3\text{H}_2(g) \rightleftharpoons 2\text{NH}_3(g)$ 
  - \_\_\_ moles of reactants  $\rightleftharpoons$  \_\_\_ moles of products
- An \_\_\_ in pressure will \_\_\_ concentration of reactants and products on \_\_\_\_\_ sides of equation
  - Molecules move closer together so reactants \_\_\_\_\_ to make \_\_\_\_\_ product
  - Will favor the \_\_\_\_\_ rxn towards the side with the \_\_\_\_\_ gas moles (\_\_\_\_\_(g) above)
- A \_\_\_ in pressure will favor the \_\_\_\_\_ direction.
  - \_\_\_\_\_ product made as molecules are farther apart and will \_\_\_\_\_ as successful as under high pressure

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## Le Chatlier's Priciple: Pressure continued

- When # of gas molecules/moles is the \_\_\_\_\_ on both sides, a pressure change will have \_\_\_\_\_ affect on the system reaction rate
  - $\text{H}_2(g) + \text{Cl}_2(g) \rightleftharpoons 2\text{HCl}(g)$
  - \_\_\_\_\_ molecules  $\rightleftharpoons$  \_\_\_\_\_ molecules in above example

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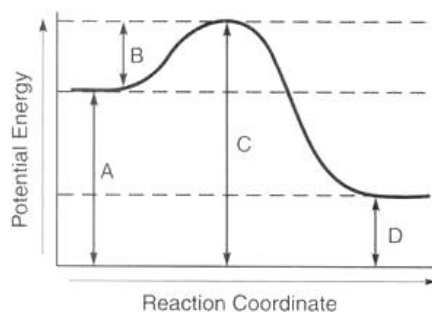
## Le Chatelier's Principle

- **Catalyst**: Introduction of a catalyst will change the rate of reaction in \_\_\_\_\_ directions
  - may cause equilibrium to be met \_\_\_\_\_
  - has no affect on \_\_\_\_\_

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## Potential Energy Diagrams (PED's)

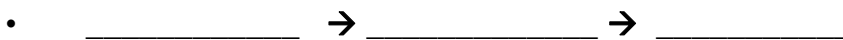
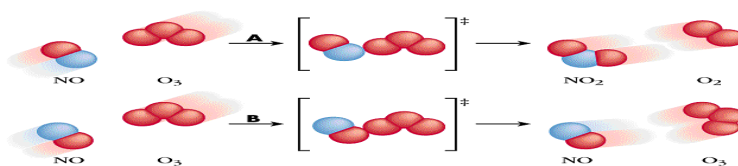
- Used to illustrate the Potential Energy changes that occur during a chemical reaction
- A= Potential Energy of \_\_\_\_\_
- B= \_\_\_\_\_
- C= Potential Energy of \_\_\_\_\_
- D= Potential Energy of \_\_\_\_\_



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## Activated Complex

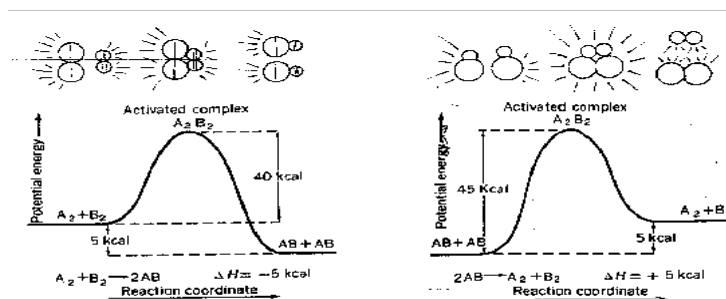
- For a reaction to occur, reactants need an effective \_\_\_\_\_ with proper \_\_\_\_\_ and \_\_\_\_\_
- As they collide/bond, their \_\_\_\_\_ energy is \_\_\_\_\_ to stored \_\_\_\_\_ energy



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# Activated Complex

- Occurs when colliding particles form a \_\_\_\_\_ intermediate product that may break apart and \_\_\_\_\_ or make a new \_\_\_\_\_

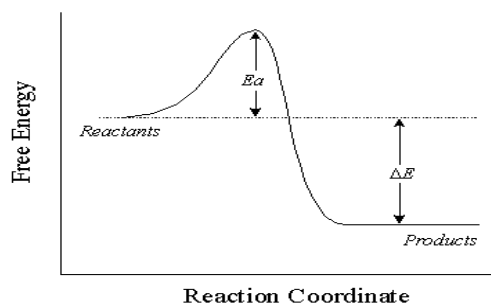


On the left is a potential energy diagram for the changes occurring in a chemical system during an exothermic reaction. On the right is the same type of diagram for a system undergoing an endothermic reaction.

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# Activation Energy

- To form the activated complex, the reactants must collide with enough \_\_\_\_\_
  - Measured as the distance between \_\_\_\_\_ and the \_\_\_\_\_ (sometimes listed as \_\_\_\_\_)



- May be measured in \_\_\_\_\_

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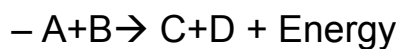
## Heat of Reaction: ( $\Delta H$ )

- = Difference between PE of \_\_\_\_\_ and PE of \_\_\_\_\_
- $\Delta H$  is the quantity of heat \_\_\_\_\_ or \_\_\_\_\_ in a chemical Rxn
- $\Delta H = H_{\text{_____}} - H_{\text{_____}}$ 
  - ( ) values indicate a \_\_\_\_\_ of heat and therefore an \_\_\_\_\_ reaction
  - See Table \_\_\_\_ for common reactions

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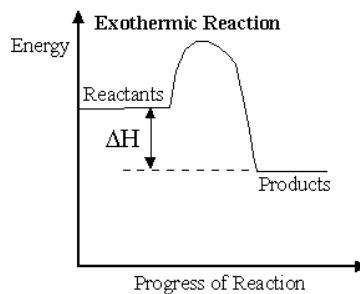
## Exothermic Reactions

- when PE of product is \_\_\_\_\_ than PE of reactant



– Heat given \_\_\_\_\_

–  $\Delta H$  is \_\_\_\_\_



- The \_\_\_\_\_ the (-) value for  $\Delta H$ , the more \_\_\_\_\_ the material

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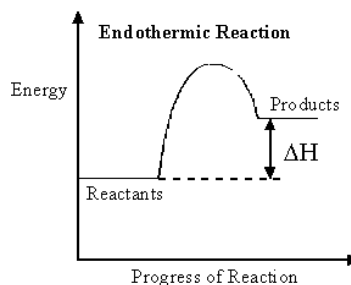
# Endothermic Reactions

- when PE of reactant is \_\_\_\_\_ than PE of product



– Heat \_\_\_\_\_

–  $\Delta H$  is \_\_\_\_\_



- If a rxn is Exothermic in the \_\_\_\_\_ direction, it will be endothermic in the \_\_\_\_\_ (and vice versa)

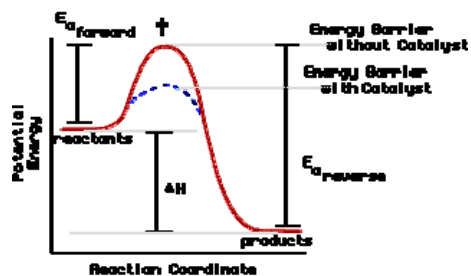
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# Catalysts

- cause a reaction to reach equilibrium \_\_\_\_\_ as they \_\_\_\_\_

– Reactant and product values remain \_\_\_\_\_

– Catalyst remains \_\_\_\_\_ and can be \_\_\_\_\_



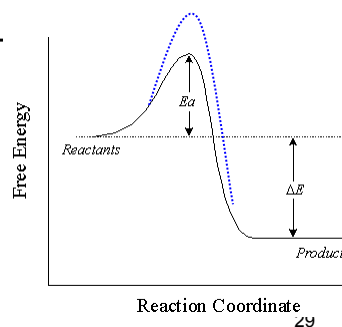
‡ Transition State

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# Inhibitor

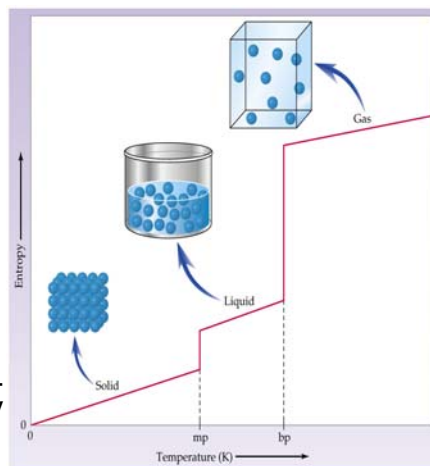
- causes a Rxn to \_\_\_\_\_ (opposite of \_\_\_\_\_) or take \_\_\_\_\_ to reach equilibrium
  - Reactant and product values remain \_\_\_\_\_
  - Activation energy \_\_\_\_\_

– Activation Energy with Inhibitor



# Entropy

- going to a state of \_\_\_\_\_
  - \_\_\_\_\_ or chaos/lack of regularity
  - Systems will tend to go from \_\_\_\_\_ (great order) to \_\_\_\_\_ (great disorder)
    - Due to particle \_\_\_\_\_
  - Ex. \_\_\_\_\_ → \_\_\_\_\_ → \_\_\_\_\_ would be the trend in entropy for molecular arrangement



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# Enthalpy

- Tendency to change to a lower \_\_\_\_\_
  - Associated with Heat of Reaction ( $\Delta H$ )
  - Exothermic Rxn's  $\rightarrow$  \_\_\_\_\_ energy, thus products will have less PE than the reactants
  - Systems tend to move towards \_\_\_\_\_ energy (\_\_\_\_\_) and greater disorder (\_\_\_\_\_)

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