

Unit 7 – Kinetics and Thermodynamics

17-1 Thermochemistry

Thermochemistry - The study of the changes in heat energy that accompany chemical reactions and physical changes

I. Heat and Temperature

A. Calorimeter

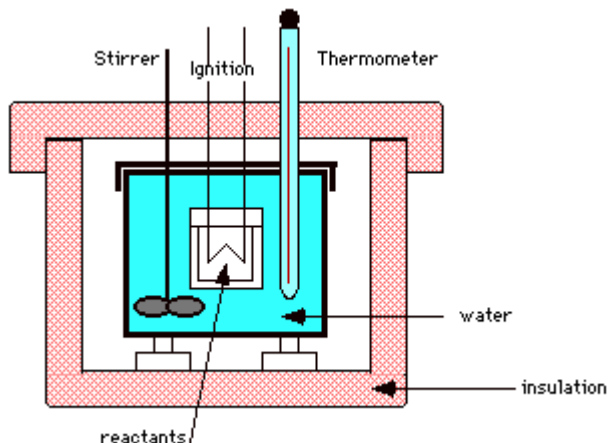
1. Heat given off or absorbed is determined from the temperature of a known mass of water

B. Temperature

1. A measure of the average kinetic energy of the particles in a sample of matter

C. Heat (or Heat Energy)

1. The sum total of the kinetic energies of the particles in a sample of matter
 - a. Heat flows spontaneously from matter at higher temperature to matter at lower temperature
2. Heat is usually measured in joules, J



$$1 \text{ J} = \text{N} \cdot \text{m} = \frac{\text{kg} \cdot \text{m}^2}{\text{s}^2}$$

- a. Heat cannot be measured directly, but is measured by temperature changes when heat is transferred

II. Heat Capacity and Specific Heat

A. Specific Heat

1. The amount of heat required to raise the temperature of one gram of substance by one Celsius degree (1°C) or one Kelvin (1 K)

B. Calculating Specific Heat

1. Specific Heat (c_p)
2. $q = \text{heat}$ $m = \text{mass}$ $\Delta T = \text{change in temperature (in Celsius or Kelvin)}$

$$c_p = \frac{q}{m \cdot \Delta T}$$

$$q = c_p \cdot m \cdot \Delta T$$

12.3 Latent Heat of Phase Change

I. Boiling and Condensation

A. Molar Heat of Vaporization

1. The amount of heat energy required to vaporize one mole of a liquid at its boiling point
2. The amount of heat energy released when one mole of a vapor condenses to a liquid at its condensation point
3. Strong attractive forces between particles result in high molar heat of vaporization

II. Freezing and Melting

A. Molar Heat of Fusion

1. The amount of heat energy required to melt one mole of solid at its melting point
2. The amount of heat energy released when one mole of a liquid freezes to a solid at its freezing point

Problem: The molar heat of fusion of water is 6.009 kJ/mol. How much energy is needed to convert 60 grams of ice at 0°C to liquid water at 0°C?

$$\frac{60 \text{ g } H_2O}{18.02 \text{ g } H_2O} \left| \frac{1 \text{ mol } H_2O}{1 \text{ mol } H_2O} \right| \frac{6.009 \text{ kJ}}{1 \text{ mol}} = 20.00 \text{ kJ}$$

III. Heat of Reaction

A. Enthalpy (H)

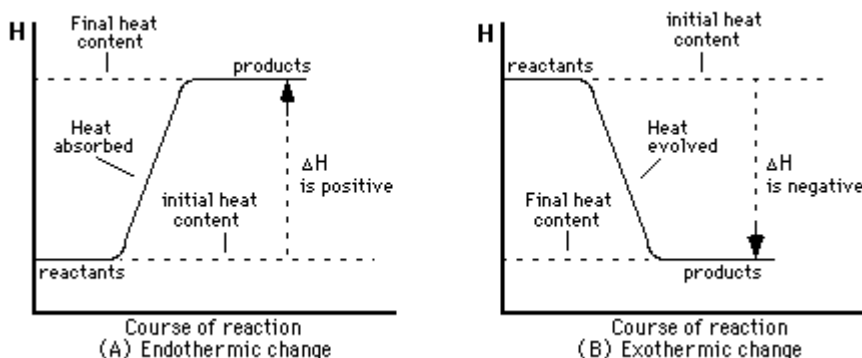
1. Heat content of a substance under constant pressure
2. Enthalpy cannot be measured directly (it is NOT the same as the temperature)

B. Enthalpy Change (ΔH)

1. The amount of heat absorbed or lost by a system during a process at constant temperature

$$\Delta H = H_{\text{products}} - H_{\text{reactants}}$$

- a. ΔH is positive for an endothermic rxn
 - (1) Heat content of products is greater than the heat content of the reactants
- b. ΔH is negative for an exothermic rxn
 - (1) Heat content of the reactants is greater than the heat content of the products



C. Writing Thermochemical Equations

1. Fraction coefficients may be used because coefficients represent mole quantities, not atoms or molecules
2. Use appropriate state/phase symbols (g) (l) (s)
3. ΔH is proportional to the number of moles
4. ΔH is usually not influenced significantly by the temperature of the system

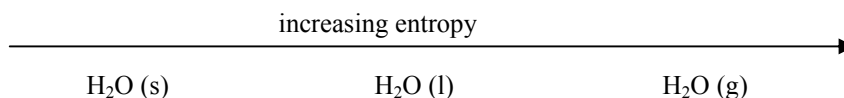
17-2 Driving Forces of Reactions

A. Enthalpy and Reaction Tendency

1. Tendency for processes to occur that lead to the lowest possible energy state
2. Most spontaneous reactions have a negative value for ΔH

B. Entropy (S)

1. Entropy is a measure of the degree of randomness of the particles, such as molecules in a system



C. Entropy and Reaction Tendency

1. There is a tendency for the disorder of the system to increase

17-3 The Reaction Process

I. Reaction Mechanisms

A. Reaction Mechanism

1. The step-by-step sequence of reactions by which the overall chemical change occurs
2. Reactions proceed by a series of simple steps called the reaction pathway

B. Intermediates

1. Species that appear in some steps but not in the net equation

C. Homogeneous Reaction

1. A reaction whose reactants and products exist in the same phase

II. The Two Principals of Collision Theory

- A. Particles must collide while favorably oriented
- B. Particles must collide with enough energy to disrupt the bonds of the molecules

III. Activation Energy and the Activated Complex

A. Activated Complex

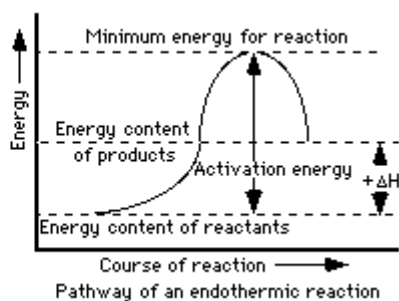
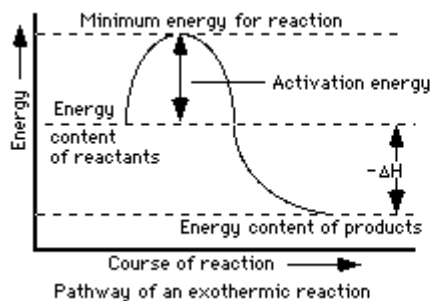
1. A transitional structure resulting from an effective collision that persists while old bonds are breaking and new bonds are forming

B. Activation Energy

1. The minimum energy required to transform reactants into the activated complex

C. Sources of Activation Energy

1. Flame, spark, high temperature, radiation
2. Once an exothermic rxn is activated (reaches the top) the energy released by the rxn sustains the rxn by activating other molecules



17-4 Reaction Rate

I. Rate Influencing Factors

A. Nature of the Reactants

1. Rate depends on the particular bonds involved

B. Amount of Surface

1. The more surface area of a solid exposed, the faster the rxn

C. Temperature

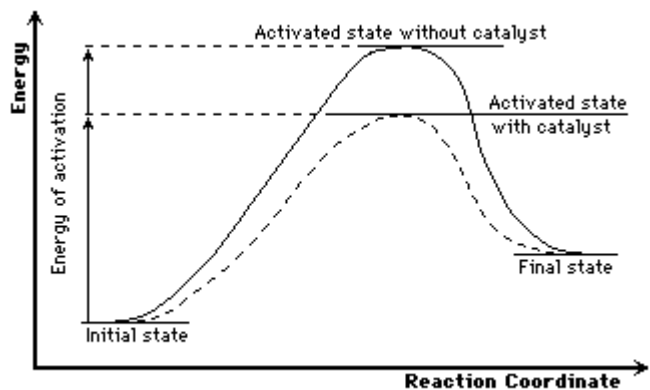
1. Rates of rxns are roughly doubled for every 10 °C increase in temperature
 - a. increase in collision energy
 - b. increase in collision frequency

D. Effect of Concentration

1. Increasing concentrations of liquid or gaseous solutions usually increases rates of rxns
2. Actual effect of concentration change must be determined experimentally for each rxn

E. Presence of Catalysts

1. A catalyst is a substance of substances that increase the rate of a chemical rxn without itself being changed
2. Catalysts provide an alternate reaction mechanism or pathway with a lower activation energy



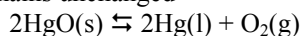
18.1 – 18.2 Equilibrium and Le Chatelier's Principle

A. Reversible Reactions

1. A chemical reaction in which the products can react to re-form the reactants

B. Chemical Equilibrium

1. When the rate of the forward reaction equals the rate of the reverse reaction and the concentration of products and reactants remains unchanged



2. Arrows going both directions indicates equilibrium in a chemical equation

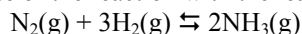
C. Le Chatelier's Principle

1. When a closed system at equilibrium is disturbed by application of a stress, it attains a new equilibrium position that minimizes the stress

D. Equilibrium and System Stresses – Predicting the direction of shift

1. Pressure

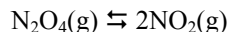
- a. An increase in pressure will cause the reaction to shift in the direction that results in fewer collisions – toward the side of the reaction with the least number of moles of gas molecules



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An increase in pressure causes the equilibrium position to shift to the right, where there are only two moles of gas. The left side has four moles of gas

- b. Decreases in pressure cause the reaction to shift in the direction that increases collisions; toward the side with more moles of gas



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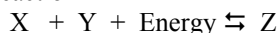
A decrease in pressure causes the equilibrium position to shift to the right, because there are more moles of gas (2 moles) than on the left (1 mole)

- c. Pressure has no effect on equilibrium if there are no gas phase reactants or products

2. Temperature

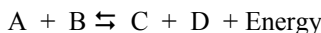
- a. Increasing temperature causes the reaction to shift in a way so as to use up the added energy (away from the side that has energy)

Endothermic reaction



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Exothermic reaction



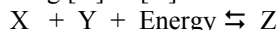
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- b. Decreasing temperature causes the reaction to shift in a way so as to replace the missing energy (toward the side that has energy)

3. Concentration

- a. Increasing the concentration of a reactant or product causes the reaction to shift in such a way as to try to use up the added substance (away from that substance)

Result of increasing [X] or [Y]

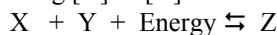


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The reaction shifts away from [X] and [Y] and produces more [Z]

- b. Decreasing the concentration of a reactant or product causes the reaction to shift in such a way as to try to replace the missing substance (toward that substance)

Result of decreasing [X] or [Y]



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The reaction shifts toward [X] and [Y] and produces more [X] and [Y]