Skills to Develop

- To predict the effects of stresses on a system at equilibrium.

Chemists use various strategies to increase the yield of the desired products of reactions. When synthesizing an ester, for example, how can a chemist control the reaction conditions to obtain the maximum amount of the desired product? Only three types of stresses can change the composition of an equilibrium mixture:

1. a change in the concentrations (or partial pressures) of the components by adding or removing reactants or products,
2. a change in the total pressure or volume, and
3. a change in the temperature of the system.

In this section, we explore how changes in reaction conditions can affect the equilibrium composition of a system. We will explore each of these possibilities in turn.

Changes in Concentration

If we add a small volume of carbon tetrachloride (CCl₄) solvent to a flask containing crystals of iodine, we obtain a saturated solution of I₂ in CCl₄, along with undissolved crystals:

\[ I_{2(s)} \rightleftharpoons I_{2(soln)} \tag{15.5.1} \]

The system reaches equilibrium, with \( K = [I_2] \). If we add more CCl₄, thereby diluting the solution, \( Q \) is now less than \( K \). Le Châtelier’s principle tells us that the system will react to relieve the stress—but how? Adding solvent stressed the system by decreasing the concentration of dissolved I₂. Hence more crystals will dissolve, thereby increasing the concentration of dissolved I₂ until the system again reaches equilibrium if enough solid I₂ is available (Figure 15.5.1). By adding solvent, we drove the reaction shown in Equation 15.5.1 to the right as written.
Figure 15.5.1: The Concentration of Dissolved $I_2$ as a Function of Time Following the Addition of More Solvent to a Saturated Solution in Contact with Excess Solid $I_2$. The concentration of $I_2$ decreases initially due to dilution but returns to its original value as long as solid $I_2$ is present.

We encounter a more complex system in the reaction of hydrogen and nitrogen to form ammonia:

$$[N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)] \tag{15.5.2}$$

The $K_p$ for this reaction is $2.14 \times 10^{-2}$ at about 540 K. Under one set of equilibrium conditions, the partial pressure of ammonia is $P_{NH_3} = 0.454$ atm, that of hydrogen is $P_{H_2} = 2.319$ atm, and that of nitrogen is $P_{N_2} = 0.773$ atm. If an additional 1 atm of hydrogen is added to the reactor to give $P_{H_2} = 3.319$ atm, how will the system respond? Because the stress is an increase in $P_{H_2}$, the system must respond in some way that decreases the partial pressure of hydrogen to counteract the stress. The reaction will therefore proceed to the right as written, consuming $H_2$ and $N_2$ and forming additional $NH_3$. Initially, the partial pressures of $H_2$ and $N_2$ will decrease, and the partial pressure of $NH_3$ will increase until the system eventually reaches a new equilibrium composition, which will have a net increase in $P_{H_2}$.

We can confirm that this is indeed what will happen by evaluating $Q_p$ under the new conditions and comparing its value with $K_p$. The equations used to evaluate $K_p$ and $Q_p$ have the same form: substituting the values after adding hydrogen into the expression for $Q_p$ results in the following:

$$Q_p = \frac{(P_{NH_3})^2}{(P_{N_2})(P_{H_2})^3} = \frac{(0.454)^2}{(0.773)(2.319+1.00)^3} = 7.29 \times 10^{-3}$$

Thus $Q_p < K_p$, which tells us that the ratio of products to reactants is less than at equilibrium. To reach equilibrium, the reaction must proceed to the right as written: the partial pressures of the products will increase, and the partial pressures of the reactants will decrease. $Q_p$ will thereby increase until it equals $K_p$, and the system will once again be at equilibrium. Changes in the partial pressures of the various substances in the reaction mixture (Equation 15.5.2) as a function of time are shown in Figure 15.5.2.
The added hydrogen is consumed by reacting with nitrogen to produce more ammonia, allowing the system to reach a new equilibrium composition.

We can force a reaction to go essentially to completion, regardless of the magnitude of \(K\), by continually removing one of the products from the reaction mixture. Consider, for example, the methanation reaction, in which hydrogen reacts with carbon monoxide to form methane and water:

\[
\text{CO} + 3\text{H}_2 \rightleftharpoons \text{CH}_4 + \text{H}_2\text{O}
\]

This reaction is used for the industrial production of methane, whereas the reverse reaction is used for the production of \(\text{H}_2\) (Example 14). The expression for \(Q\) has the following form:

\[
Q = \frac{[\text{CH}_4][\text{H}_2\text{O}]}{[\text{CO}][\text{H}_2]^3}
\]

Regardless of the magnitude of \(K\), if either \([\text{H}_2\text{O}]\) or \([\text{CH}_4]\) can be removed from the reaction mixture so that \([\text{H}_2\text{O}]\) or \([\text{CH}_4]\) is approximately zero, then \(Q \approx 0\). In other words, when product is removed, the system is stressed \((Q \ll K)\), and more product will form to counter the stress. Because water \((\text{bp} = 100^\circ\text{C})\) is much less volatile than methane, hydrogen, or carbon monoxide \((\text{all of which have boiling points below } -100^\circ\text{C})\), passing the gaseous reaction mixture through a cold coil will cause the water vapor to condense to a liquid that can be drawn off. Continuing to remove water from the system forces the reaction to the right as the system attempts to equilibrate, thus enriching the reaction mixture in methane. This technique, referred to as driving a reaction to completion, can be used to force a reaction to completion even if \(K\) is relatively small. For example, esters are usually synthesized by removing water via a "condensation reaction" as shown here:
Example 15.5.1

For each equilibrium system, predict the effect of the indicated stress on the specified quantity.

a. \(2\text{SO}_2(g) + \text{O}_2(g) \rightleftharpoons 2\text{SO}_3(g)\):
   1. the effect of removing \(\text{O}_2\) on \(P_\text{SO}_2\);
   2. the effect of removing \(\text{O}_2\) on \(P_\text{SO}_3\)

b. \(\text{CaCO}_3(s) \rightleftharpoons \text{CaO}_(s) + \text{CO}_2(g)\):
   1. the effect of removing \(\text{CO}_2\) on the amount of \(\text{CaCO}_3\);
   2. the effect of adding \(\text{CaCO}_3\) on \(P_\text{CO}_2\)

Given: balanced chemical equations and changes

Asked for: effects of indicated stresses

Strategy:

Use \(Q\) and \(K\) to predict the effect of the stress on each reaction.

Solution:

1. Removing \(\text{O}_2\) will decrease \(P_\text{O}_2\), thereby decreasing the denominator in the reaction quotient and making \(Q_p > K_p\). The reaction will proceed to the left as written, increasing the partial pressures of \(\text{SO}_2\) and \(\text{O}_2\) until \(Q_p\) once again equals \(K_p\).

2. Removing \(\text{O}_2\) will decrease \(P_\text{O}_2\) and thus increase \(Q_p\), so the reaction will proceed to the left. The partial pressure of \(\text{SO}_3\) will decrease.

\(K_p\) and \(Q_p\) are both equal to \(P_\text{CO}_2\).

1. Removing \(\text{CO}_2\) from the system causes more \(\text{CaCO}_3\) to react to produce \(\text{CO}_2\), which increases \(P_\text{CO}_2\) to the partial pressure required by \(K_p\).

2. Adding (or removing) solid \(\text{CaCO}_3\) has no effect on \(P_\text{CO}_2\) because it does not appear in the expression for \(K_p\) (or \(Q_p\)).

Exercise 15.5.1

For each equilibrium system, predict the effect that the indicated stress will have on the specified quantity.

a. \(\text{H}_2(g) + \text{CO}_2(g) \rightleftharpoons \text{H}_2\text{O}(g) + \text{CO}(g)\):
   1. the effect of adding \(\text{CO}\) on \(\text{H}_2\);
   2. the effect of adding \(\text{CO}_2\) on \(\text{H}_2\)

b. \(\text{CuO}(s) + \text{CO}(g) \rightleftharpoons \text{Cu}(s) + \text{CO}_2(g)\):
   1. the effect of adding \(\text{CO}\) on the amount of \(\text{Cu}\);
2. the effect of adding \(\text{CO}_2\) on \([\text{CO}])\)

Answer:

- (1) \([\text{H}_2]\) increases; (2) \([\text{H}_2]\) decreases.
- (1) the amount of \(\text{Cu}\) increases; (2) \([\text{CO}])\) increases.

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**Changes in Total Pressure or Volume**

Because liquids are relatively incompressible, changing the pressure above a liquid solution has little effect on the concentrations of dissolved substances. Consequently, changes in external pressure have very little effect on equilibrium systems that contain only solids or liquids. In contrast, because gases are highly compressible, their concentrations vary dramatically with pressure. From the ideal gas law, \((PV = nRT)\), the concentration \((\text{mol/L})\) of a gas is related to its pressure as follows:

\[
[C = \frac{\text{n}}{V} = \frac{P}{RT} \tag{15.5.5}]
\]

Hence the concentration of any gaseous reactant or product is directly proportional to the applied pressure \((P)\) and inversely proportional to the total volume \((V)\). Consequently, the equilibrium compositions of systems that contain gaseous substances are quite sensitive to changes in pressure, volume, and temperature.

These principles can be illustrated using the reversible dissociation of gaseous \(\text{N}_2\text{O}_4\) to gaseous \(\text{NO}_2\)

\[
\text{N}_2\text{O}_4 \rightleftharpoons 2 \text{NO}_2 \]

The syringe shown in Figure 15.5.3 initially contains an equilibrium mixture of colorless \(\text{N}_2\text{O}_4\) and red-brown \(\text{NO}_2\). Decreasing the volume by 50% causes the mixture to become darker because all concentrations have doubled. Decreasing the volume also constitutes a stress, however, as we can see by examining the effect of a change in volume on \(Q\). At equilibrium,

\[
[Q = K = \frac{[\text{NO}_2]^2}{[\text{N}_2\text{O}_4]} \tag{15.5.6a}]\]

If the volume is decreased by half, the concentrations of the substances in the mixture are doubled, so the new reaction quotient is now

\[
[Q = \frac{([\text{NO}_2]_i)^2}{[\text{N}_2\text{O}_4]_i} = \frac{4([\text{NO}_2]_i)^2}{2[\text{N}_2\text{O}_4]_i} = 2K \tag{15.5.6b}]\]

Because \([Q]\) is now greater than \([K]\), the system is no longer at equilibrium. The stress can be relieved if the reaction proceeds to the left, consuming 2 mol of \(\text{NO}_2\) for every 1 mol of \(\text{N}_2\text{O}_4\) produced. This will decrease the concentration of \(\text{NO}_2\) and increase the concentration of \(\text{N}_2\text{O}_4\), causing \([Q]\) to decrease until it once again equals \([K]\). Thus, as shown in Figure 15.5.3, the intensity of the brown color due to \(\text{NO}_2\) decreases with time following the change in volume.
Figure 15.5.3 The Effect of Changing the Volume (and Thus the Pressure) of an Equilibrium Mixture of \(N_2O_4\) and \(NO_2\) at Constant Temperature. (a) The syringe with a total volume of 15 mL contains an equilibrium mixture of \(N_2O_4\) and \(NO_2\); the red-brown color is proportional to the \(NO_2\) concentration. (b) If the volume is rapidly decreased by a factor of 2 to 7.5 mL, the initial effect is to double the concentrations of all species present, including \(NO_2\). Hence the color becomes more intense. (c) With time, the system adjusts its composition in response to the stress as predicted by Le Châtelier's principle, forming colorless \(N_2O_4\) at the expense of red-brown \(NO_2\), which decreases the intensity of the color of the mixture.

Note

Increasing the pressure of a system (or decreasing the volume) favors the side of the reaction that has fewer gaseous molecules and vice versa.

In general, if a balanced chemical equation contains different numbers of gaseous reactant and product molecules, the equilibrium will be sensitive to changes in volume or pressure. Increasing the pressure on a system (or decreasing the volume) will favor the side of the reaction that has fewer gaseous molecules and vice versa.

Example 15.5.2

For each equilibrium system, write the reaction quotient for the system if the pressure is decreased by a factor of 2 (i.e., if the volume is doubled) at constant temperature and then predict the direction of the reaction.

a. \(N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)\)

b. \(C_2H_2(g) + C_2H_6(g) \rightleftharpoons 2C_2H_4(g)\)

c. \(2 NO(g) \rightleftharpoons 2NO_2(g) + O_2(g)\)
**Given:** balanced chemical equations

**Asked for:** direction of reaction if pressure is halved

**Strategy:**

Use Le Châtelier’s principle to predict the effect of the stress.

**Solution:**

Two moles of gaseous products are formed from 4 mol of gaseous reactants. Decreasing the pressure will cause the reaction to shift to the left because that side contains the larger number of moles of gas. Thus the pressure increases, counteracting the stress. \(K\) for this reaction is \(\frac{[NH_3]^2}{[N_2][H_2]^3}\). When the pressure is decreased by a factor of 2, the concentrations are halved, which means that the new reaction quotient is as follows:

\[
Q = \frac{1/4[NH_3]^2}{1/16[N_2][H_2]^3} = 4K
\]

Two moles of gaseous products form from 2 mol of gaseous reactants. Decreasing the pressure will have no effect on the equilibrium composition because both sides of the balanced chemical equation have the same number of moles of gas. Here \(K = [C_2H_4]^2/[C_2H_2][C_2H_6]\). The new reaction quotient is as follows:

\[
Q = \frac{1/4[C_2H_4]^2}{1/4[C_2H_2][C_2H_6]} = K
\]

Three moles of gaseous products are formed from 2 mol of gaseous reactants. Decreasing the pressure will favor the side that contains more moles of gas, so the reaction will shift toward the products to increase the pressure. For this reaction \(K = [NO]^2[O_2]/[NO_2]^2\). Under the new reaction conditions the reaction quotient is as follows:

\[
Q = \frac{1/8[NO]^2[O_2]}{1/4[NO_2]^2} = 1/2K
\]

**Exercise 15.5.2**

For each equilibrium system, write a new reaction quotient for the system if the pressure is increased by a factor of 2 (i.e., if the volume is halved) at constant temperature and then predict the direction in which the reaction will shift.

a. \(H_2O_{(g)}+CO_{(g)} \rightleftharpoons H_{2(g)}+CO_{2(g)}\)

b. \(H_{2(g)}+C_2H_{4(g)} \rightleftharpoons C_2H_{6(g)}\)

c. \(2SO_{2(g)}+O_{2(g)} \rightleftharpoons 2SO_{3(g)}\)

**Answer:**

a. \(Q = K\); no effect

b. \(Q = 1/2 K\); to the right

c. \(Q = 1/2 K\); to the right
Changes in Temperature

In all the cases we have considered so far, the magnitude of the equilibrium constant, $K$ or $K_p$, was constant. Changes in temperature can, however, change the value of the equilibrium constant without immediately affecting the reaction quotient ($Q \neq K$). In this case, the system is no longer at equilibrium; the composition of the system will change until $Q$ equals $K$ at the new temperature.

To predict how an equilibrium system will respond to a change in temperature, we must know something about the enthalpy change of the reaction ($\Delta H_{rxn}$). As you learned previously, heat is released to the surroundings in an exothermic reaction ($\Delta H_{rxn} < 0$), and heat is absorbed from the surroundings in an endothermic reaction ($\Delta H_{rxn} > 0$). We can express these changes in the following way (although not technically correct since heat is not a species):

- Exothermic ($\Delta H < 0$): $\text{reactants} \rightleftharpoons \text{products} + \text{heat}$
- Endothermic ($\Delta H > 0$): $\text{reactants} + \text{heat} \rightleftharpoons \text{products}$

Thus heat can be thought of as a product in an exothermic reaction and as a reactant in an endothermic reaction. Increasing the temperature of a system corresponds to adding heat. Le Châtelier's principle predicts that an exothermic reaction will shift to the left (toward the reactants) if the temperature of the system is increased (heat is added). Conversely, an endothermic reaction will shift to the right (toward the products) if the temperature of the system is increased. If a reaction is thermochemically neutral ($\Delta H_{rxn} = 0$), then a change in temperature will not affect the equilibrium composition.

We can examine the effects of temperature on the dissociation of $N_2O_4$ to $NO_2$, for which $\Delta H = +58$ kJ/mol. This reaction can be written as follows:

$$58\;kJ+N_2O_4(g) \rightleftharpoons 2\;NO_2(g)$$

Increasing the temperature (adding heat to the system) is a stress that will drive the reaction to the right, as illustrated in Figure 15.5.4. Thus increasing the temperature increases the ratio of $NO_2$ to $N_2O_4$ at equilibrium, which increases $K$. 
The effect of temperature on the equilibrium between gaseous \(N_2O_4\) and \(NO_2\). The value of \(K\) changes with temperature of the same 99.9% pure \(NO_2/N_2O_4\) sealed in an ampoule. From left to right -196°C, 0°C, 23°C, 35°C, 50°C. In the endothermic reaction, the equilibrium position can be shifted by changing the temperature. When heat is added and the temperature increases, the reaction shifts to the right and the flask turns reddish brown due to an increase in \(NO_2\). When heat is removed and the temperature decreases, the reaction shifts to the left and flask turns colorless due to an increase in \(N_2O_4\). This demonstrates Le Chatelier’s Principle because the equilibrium shifts in the direction that consumes energy. Figure used with permission from Wikipedia.

The effect of increasing the temperature on a system at equilibrium can be summarized as follows: increasing the temperature increases the magnitude of the equilibrium constant for an endothermic reaction, decreases the equilibrium constant for an exothermic reaction, and has no effect on the equilibrium constant for a thermally neutral reaction. Table 15.5.1 shows the temperature dependence of the equilibrium constants for the synthesis of ammonia from hydrogen and nitrogen, which is an exothermic reaction with \(\Delta H^\circ = -91.8 \text{ kJ/mol}\). The values of both \(K\) and \(K_p\) decrease dramatically with increasing temperature, as predicted for an exothermic reaction.

**Table 15.5.1: Temperature Dependence of K and Kp for**

\[N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)\]

<table>
<thead>
<tr>
<th>Temperature (K)</th>
<th>(K)</th>
<th>(K_p)</th>
</tr>
</thead>
<tbody>
<tr>
<td>298</td>
<td>(3.3 \times 10^8)</td>
<td>(5.6 \times 10^5)</td>
</tr>
<tr>
<td>400</td>
<td>(3.9 \times 10^4)</td>
<td>(3.6 \times 10^1)</td>
</tr>
<tr>
<td>450</td>
<td>(2.6 \times 10^3)</td>
<td>(1.9)</td>
</tr>
<tr>
<td>500</td>
<td>(1.7 \times 10^2)</td>
<td>(1.0 \times 10^{-1})</td>
</tr>
<tr>
<td>550</td>
<td>(2.6 \times 10^1)</td>
<td>(1.3 \times 10^{-2})</td>
</tr>
<tr>
<td>600</td>
<td>(4.1)</td>
<td>(1.7 \times 10^{-3})</td>
</tr>
</tbody>
</table>
Increasing the temperature causes endothermic reactions to favor products and exothermic reactions to favor reactants.

Example 15.5.3

For each equilibrium reaction, predict the effect of decreasing the temperature:

a. \( \text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \rightleftharpoons 2\text{NH}_3(\text{g}) \); \( \Delta H_{\text{rxn}} = -91.8 \text{ kJ/mol} \)

b. \( \text{CaCO}_3(\text{s}) \rightleftharpoons \text{CaO}(\text{s}) + \text{CO}_2(\text{g}) \); \( \Delta H_{\text{rxn}} = 178 \text{ kJ/mol} \)

**Given:** balanced chemical equations and values of \( \Delta H_{\text{rxn}} \)

**Asked for:** effects of decreasing temperature

**Strategy:**

Use Le Châtelier’s principle to predict the effect of decreasing the temperature on each reaction.

**Solution:**

The formation of \( \text{NH}_3 \) is exothermic, so we can view heat as one of the products:

\[
\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \rightleftharpoons 2\text{NH}_3(\text{g}) + 91.8 \text{ kJ}
\]

If the temperature of the mixture is decreased, heat (one of the products) is being removed from the system, which causes the equilibrium to shift to the right. Hence the formation of ammonia is favored at lower temperatures.

The decomposition of calcium carbonate is endothermic, so heat can be viewed as one of the reactants:

\[
\text{CaCO}_3(\text{s}) + 178 \text{ kJ} \rightleftharpoons \text{CaO}(\text{s}) + \text{CO}_2(\text{g})
\]

If the temperature of the mixture is decreased, heat (one of the reactants) is being removed from the system, which causes the equilibrium to shift to the left. Hence the thermal decomposition of calcium carbonate is less favored at lower temperatures.

Exercise 15.5.3

For each equilibrium system, predict the effect of increasing the temperature on the reaction mixture:

- \( 2\text{SO}_2(\text{g}) + \text{O}_2(\text{g}) \rightleftharpoons 2\text{SO}_3(\text{g}) \) with \( \Delta H_{\text{rxn}} = -198 \text{ kJ/mol} \)
- \( \text{N}_2(\text{g}) + \text{O}_2(\text{g}) \rightleftharpoons 2\text{NO}(\text{g}) \) with \( \Delta H_{\text{rxn}} = +181 \text{ kJ/mol} \)

**Answer:**

- Reaction shifts to the left.
- Reaction shifts to the right.
Summary

Three types of stresses can alter the composition of an equilibrium system: adding or removing reactants or products, changing the total pressure or volume, and changing the temperature of the system. A reaction with an unfavorable equilibrium constant can be driven to completion by continually removing one of the products of the reaction. Equilibria that contain different numbers of gaseous reactant and product molecules are sensitive to changes in volume or pressure; higher pressures favor the side with fewer gaseous molecules. Removing heat from an exothermic reaction favors the formation of products, whereas removing heat from an endothermic reaction favors the formation of reactants.

Key Takeaway

- Equilibria are affected by changes in concentration, total pressure or volume, and temperature.

Conceptual Problems

1. If an equilibrium reaction is endothermic in the forward direction, what is the expected change in the concentration of each component of the system if the temperature of the reaction is increased? If the temperature is decreased?

2. Write the equilibrium equation for the following system: \(4\text{NH}_3(g) + 5\text{O}_2(g) \rightleftharpoons 4\text{NO}_2(g) + 6\text{H}_2\text{O}(g)\). Would you expect the equilibrium to shift toward the products or reactants with an increase in pressure? Why?

3. The reaction rate approximately doubles for every 10°C rise in temperature. What happens to K?

4. The formation of \(\text{A}_2\text{B}_2(g)\) via the equilibrium reaction \(2\text{AB}(g) \rightleftharpoons \text{A}_2\text{B}_2(g)\) is exothermic. What happens to the ratio \(k_f/k_r\) if the temperature is increased? If both temperature and pressure are increased?

5. In each system, predict the effect that the indicated change will have on the specified quantity at equilibrium:
   a. \(\text{H}_2(g) + \text{I}_2(g) \rightleftharpoons 2\text{HI}(g)\): If \(\text{H}_2\) were removed; what is the effect on \(P_{\text{I}_2}\)?
   b. \(2\text{NOBr}(g) \rightleftharpoons 2\text{NO}(g) + \text{Br}_2(g)\): If \(\text{Br}_2\) were removed; what is the effect on \(P_{\text{NOBr}}\)?
   c. \(2\text{NaHCO}_3(s) \rightleftharpoons \text{Na}_2\text{CO}_3(g) + \text{CO}_2(g) + \text{H}_2\text{O}(g)\): If \(\text{CO}_2\) were removed; what is the effect on \(P_{(\text{NaHCO}_3)}\)?

6. What effect will the indicated change have on the specified quantity at equilibrium?
   a. \(\text{NH}_4\text{Cl}(s) \rightleftharpoons \text{NH}_3(g) + \text{HCl}(g)\): \(\text{NH}_4\text{Cl}\) is increased; what is the effect on \(P_{(\text{HCl})}\)?
   b. \(2\text{H}_2\text{O}(g) \rightleftharpoons 2\text{H}_2(g) + \text{O}_2(g)\): \(\text{O}_2\) is added; what is the effect on \(P_{(\text{H}_2)}\)?
   c. \(\text{PCl}_3(g) + \text{Cl}_2(g) \rightleftharpoons \text{PCl}_5(g)\): \(\text{Cl}_2\) is removed; what is the effect on \(P_{(\text{PCl}_5)}\)?

Numerical Problems

For each equilibrium reaction, describe how Q and K change when the pressure is increased, the temperature is increased, the volume of the system is increased, and the concentration(s) of the reactant(s) is increased.
\[
\begin{align*}
\text{(1) } & \quad A_{(g)} \rightleftharpoons B_{(g)}: \quad \Delta H = -20.6 \text{ kJ/mol} \\
\text{(2) } & \quad 2A_{(g)} \rightleftharpoons B_{(g)}: \quad \Delta H = 0.3 \text{ kJ/mol} \\
\text{(3) } & \quad A_{(g)} + B_{(g)} \rightleftharpoons 2C_{(g)}: \quad \Delta H = 46 \text{ kJ/mol}
\end{align*}
\]

For each equilibrium reaction, describe how \(Q\) and \(K\) change when the pressure is decreased, the temperature is increased, the volume of the system is decreased, and the concentration(s) of the reactant(s) is increased.

\[
\begin{align*}
\text{(4) } & \quad 2A_{(g)} \rightleftharpoons B_{(g)}: \quad \Delta H = -80 \text{ kJ/mol} \\
\text{(5) } & \quad A_{(g)} \rightleftharpoons 2B_{(g)}: \quad \Delta H = 0.3 \text{ kJ/mol} \\
\text{(6) } & \quad 2A_{(g)} \rightleftharpoons 2B_{(g)} + C_{(g)}: \quad \Delta H = 46 \text{ kJ/mol}
\end{align*}
\]

Le Châtelier's principle states that a system will change its composition to counteract stress. For the system \(CO(g) + Cl_2(g) \rightleftharpoons COCl_2(g)\), write the equilibrium constant expression \(K_p\). What changes in the values of \(Q\) and \(K\) would you anticipate when (a) the volume is doubled, (b) the pressure is increased by a factor of 2, and (c) \(COCl_2\) is removed from the system?

For the equilibrium system \(3O_2(g) \rightleftharpoons 2O_3(g), \Delta H^\circ = 284 \text{ kJ}\), write the equilibrium constant expression \(K_p\). What happens to the values of \(Q\) and \(K\) if the reaction temperature is increased? What happens to these values if both the temperature and pressure are increased?

Carbon and oxygen react to form \(CO_2\) gas via \(C(s) + O_2(g) \rightleftharpoons CO_2(g)\), for which \(K = 1.2 \times 10^{69}\). Would you expect \(K\) to increase or decrease if the volume of the system were tripled? Why?

The reaction \(COCl_2(g) \rightleftharpoons CO(g) + Cl_2(g)\) has \(K = 2.2 \times 10^{-10}\) at 100°C. Starting with an initial \(PCOCl_2\) of 1.0 atm, you determine the following values of \(PCO\) at three successive time intervals: \(6.32 \times 10^{-6}\) atm, \(1.78 \times 10^{-6}\) atm, and \(1.02 \times 10^{-5}\) atm. Based on these data, in which direction will the reaction proceed after each measurement? If chlorine gas is added to the system, what will be the effect on \(Q\)?

The following table lists experimentally determined partial pressures at three temperatures for the reaction \(Br_2(g) \rightleftharpoons 2Br(g)\).

<table>
<thead>
<tr>
<th>T (K)</th>
<th>1123</th>
<th>1173</th>
<th>1273</th>
</tr>
</thead>
<tbody>
<tr>
<td>(P_{Br_2}) (atm)</td>
<td>3.000</td>
<td>0.3333</td>
<td>6.755 \times 10^{-2}</td>
</tr>
<tr>
<td>(P_{Br}) (atm)</td>
<td>(3.477 \times 10^{-2})</td>
<td>(2.159 \times 10^{-2})</td>
<td>(2.191 \times 10^{-2})</td>
</tr>
</tbody>
</table>

Is this an endothermic or an exothermic reaction? Explain your reasoning.

The dissociation of water vapor proceeds according to the following reaction: \(H_2O(g) \rightleftharpoons 1/2O_2(g) + H_2(g)\). At 1300 K, there is 0.0027% dissociation, whereas at 2155 K, the dissociation is 1.18%. Calculate \(K\) and \(K_p\). Is this an endothermic reaction or an exothermic reaction? How do the magnitudes of the two Equilibria compare? Would increasing the pressure improve the yield of \(H_2\) gas at either temperature? (Hint: assume that the system initially contains 1.00 mol of \(H_2O\) in a 1.00 L container.)
When 1.33 mol of CO\(_2\) and 1.33 mol of H\(_2\) are mixed in a 0.750 L container and heated to 395°C, they react according to the following equation: CO\(_2\)(g)+H\(_2\)(g) \rightleftharpoons CO(g)+H\(_2\)O(g). If K = 0.802, what are the equilibrium concentrations of each component of the equilibrium mixture? What happens to K if H\(_2\)O is removed during the course of the reaction?

The equilibrium reaction H\(_2\)(g)+Br\(_2\)(g) \rightleftharpoons 2HBr(g) has K\(_p\) = 2.2 \times 10^9 at 298 K. If you begin with 2.0 mol of Br\(_2\) and 2.0 mol of H\(_2\) in a 5.0 L container, what is the partial pressure of HBr at equilibrium? What is the partial pressure of H\(_2\) at equilibrium? If H\(_2\) is removed from the system, what is the effect on the partial pressure of Br\(_2\)?

Iron(II) oxide reacts with carbon monoxide according to the following equation: \(\text{FeO}_{(s)} + \text{CO}_{(g)} \rightleftharpoons \text{Fe}_{(s)}+\text{CO}_{2(g)}\). At 800°C, K = 0.34; at 1000°C, K = 0.40.

A 20.0 L container is charged with 800.0 g of \(\text{CO}_2\), 1436 g of \(\text{FeO}\), and 1120 g of iron. What are the equilibrium concentrations of all components of the mixture at each temperature?

What are the partial pressures of the gases at each temperature?

If CO were removed, what would be the effect on \(P_{\text{(CO}_2)}\) at each temperature?

The equilibrium constant K for the reaction \(\text{C}_{(s)} + \text{CO}_2(g) \rightleftharpoons 2\text{CO}_{(g)}\) is 1.9 at 1000 K and 0.133 at 298 K.

If excess C is allowed to react with 25.0 g of \(\text{CO}_2\) in a 3.00 L flask, how many grams of CO are produced at each temperature?

What are the partial pressures of each gas at 298 K? at 1000 K?

Would you expect K to increase or decrease if the pressure were increased at constant temperature and volume?

Data for the oxidation of methane, \(\text{CH}_4(g)+2\text{O}_2(g) \rightleftharpoons \text{CO}_2(g)+2\text{H}_2\text{O}_{(g)}\), in a closed 5.0 L vessel are listed in the following table. Fill in the blanks and determine the missing values of Q and K (indicated by ?) as the reaction is driven to completion.

<table>
<thead>
<tr>
<th>(\text{CH}_4)</th>
<th>(\text{O}_2)</th>
<th>(\text{CO}_2)</th>
<th>(\text{H}_2\text{O})</th>
<th>(Q)</th>
<th>(K)</th>
</tr>
</thead>
<tbody>
<tr>
<td>initial (moles)</td>
<td>0.45</td>
<td>0.90</td>
<td>0</td>
<td>0</td>
<td>?</td>
</tr>
<tr>
<td>at equilibrium</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>1.29</td>
</tr>
<tr>
<td>add 0.50 mol of methane</td>
<td>0.95</td>
<td></td>
<td></td>
<td>?</td>
<td></td>
</tr>
<tr>
<td>new equilibrium</td>
<td></td>
<td></td>
<td></td>
<td>?</td>
<td></td>
</tr>
</tbody>
</table>
### CH₄ | O₂ | CO₂ | H₂O | Q | K
---|---|---|---|---|---
**initial**<br>moles | 0.45 | 0.90 | 0 | 0 | 1.29
**at equilibrium**<br> | 0.215 | 0.43 | 0.235 | 0.47 | K | 1.29
**add 0.50 mol of methane**<br> | 0.715 | 0.43 | 0.235 | 0.47 | 0.39 | 1.29
**new equilibrium**<br> | 0.665 | 0.33 | 0.285 | 0.57 | K | 1.29
**remove water**<br> | 0.665 | 0.33 | 0.285 | 0 | 0 | 1.29
**new equilibrium**<br> | 0.57 | 0.14 | 0.38 | 0.19 | K | 1.29

None of the changes would affect K; (a) Q doubles; (b) Q is halved; Q decreases.

K would not change; it does not depend on volume.

\([\text{CO}] = [\text{H}_2\text{O}] = 0.839 \text{ M}, [\text{CO}_2] = [\text{H}_2] = 0.930 \text{ M}; \text{no effect on K}\)

At 800°C, \([\text{CO}] = 0.678 \text{ M}, [\text{CO}_2] = 0.231 \text{ M}; \text{at 1000°C, [CO]} = 0.645 \text{ M}, [\text{CO}_2] = 0.264 \text{ M}\)

At 800°C, \(\text{PCO} = 59.7 \text{ atm}, \text{PCO}_2 = 20.3 \text{ atm}; \text{at 1000°C, PCO} = 67.4 \text{ atm}, \text{PCO}_2 = 27.6 \text{ atm}\).

Removing CO would cause the reaction to shift to the right, causing \(\text{PCO}_2\) to decrease.