

# 3

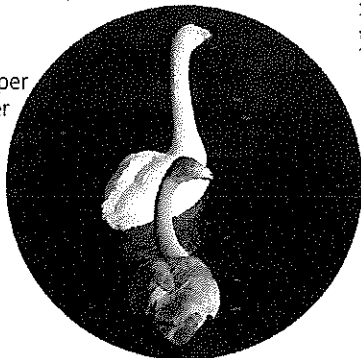
## Water and Life

### KEY CONCEPTS

- 3.1 Polar covalent bonds in water molecules result in hydrogen bonding
- 3.2 Four emergent properties of water contribute to Earth's suitability for life
- 3.3 Acidic and basic conditions affect living organisms

**AP** **BIG IDEAS:** Water figures into any theory about the origin of life, as it provides a medium in which the building blocks of life can come together and get organized into more complex molecules and form simple cells (**Big Idea 1**). The chemical and physical properties of water support living things in both their internal and external environments (**Big Ideas 2 & 4**). These complex properties relate directly to the molecular structure of this simple molecule.

► A young whooper swan paddles after its parent.



▲ **Figure 3.1** How does the habitat of a whooper swan depend on the chemistry of water?

### The Molecule That Supports All of Life

Life on Earth began in water and evolved there for 3 billion years before spreading onto land. Water is the substance that makes possible life as we know it here on Earth. All organisms familiar to us are made mostly of water and live in an environment dominated by water. Water is the biological medium here on Earth, and possibly on other planets as well.

Three-quarters of Earth's surface is covered by water. Although most of this water is in liquid form, water is also present on Earth as a solid (ice) and a gas (water vapor). Water is the only common substance to exist in the natural environment in all three physical states of matter. Furthermore, the solid state of water floats on the liquid, a rare property emerging from the chemistry of the water molecule. All three states of water can be seen in **Figure 3.1**, which shows water vapor rising from hot springs that feed into a partially frozen lake in Hokkaido, Japan. The lake is a migratory stop for the elegant whooper swan (*Cygnus cygnus*). The growing young require a watery habitat because their legs can't support their body weight on land for long periods of time.

In this chapter, you will learn how the structure of a water molecule allows it to interact with other molecules, including other water molecules. This ability leads to water's unique emergent properties that help make Earth suitable for life.

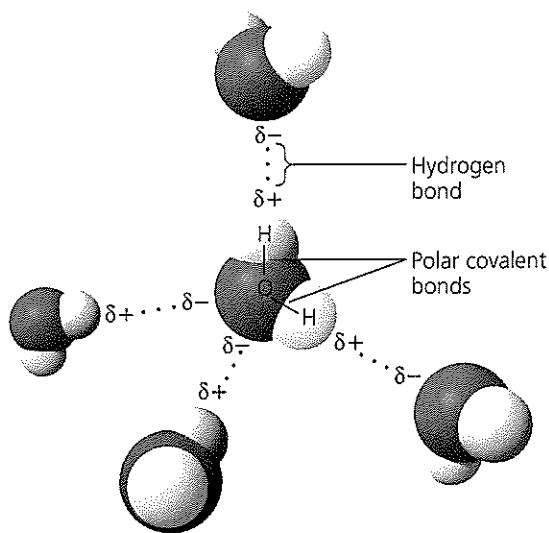
## CONCEPT 3.1

### Polar covalent bonds in water molecules result in hydrogen bonding

Water is so familiar to us that it is easy to overlook its many extraordinary qualities. Following the theme of emergent properties, we can trace water's unique behavior to the structure and interactions of its molecules.

Studied on its own, the water molecule is deceptively simple. It is shaped like a wide V, with its two hydrogen atoms joined to the oxygen atom by single covalent bonds. Oxygen is more electronegative than hydrogen, so the electrons of the covalent bonds spend more time closer to oxygen than to hydrogen; these are **polar covalent bonds** (see Figure 2.11). This unequal sharing of electrons and water's V-like shape make it a **polar molecule**, meaning that its overall charge is unevenly distributed. In water, the oxygen region of the molecule has a partial negative charge ( $\delta^-$ ), and each hydrogen has a partial positive charge ( $\delta^+$ ).

The properties of water arise from attractions between oppositely charged atoms of different water molecules: The slightly positive hydrogen of one molecule is attracted to the slightly negative oxygen of a nearby molecule. The two molecules are thus held together by a hydrogen bond (Figure 3.2). When water is in its liquid form, its hydrogen bonds are very fragile, each only about 1/20 as strong as a covalent bond. The hydrogen bonds form, break, and re-form with great

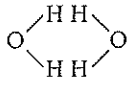


▲ **Figure 3.2** Hydrogen bonds between water molecules. The charged regions in a water molecule are due to its polar covalent bonds. Oppositely charged regions of neighboring water molecules are attracted to each other, forming hydrogen bonds. Each molecule can hydrogen-bond to multiple partners, and these associations are constantly changing.

**DRAW IT** Draw partial charges on the water molecule at the far left, and draw two more water molecules hydrogen-bonded to it.

frequency. Each lasts only a few trillionths of a second, but the molecules are constantly forming new hydrogen bonds with a succession of partners. Therefore, at any instant, most of the water molecules are hydrogen-bonded to their neighbors. The extraordinary properties of water emerge from this hydrogen bonding, which organizes water molecules into a higher level of structural order.

#### CONCEPT CHECK 3.1

1. **MAKE CONNECTIONS** What is electronegativity, and how does it affect interactions between water molecules? (Review Figure 2.11.)
2. Why is it unlikely that two neighboring water molecules would be arranged like this?  

3. **WHAT IF?** What would be the effect on the properties of the water molecule if oxygen and hydrogen had equal electronegativity?

For suggested answers, see Appendix A.

## CONCEPT 3.2

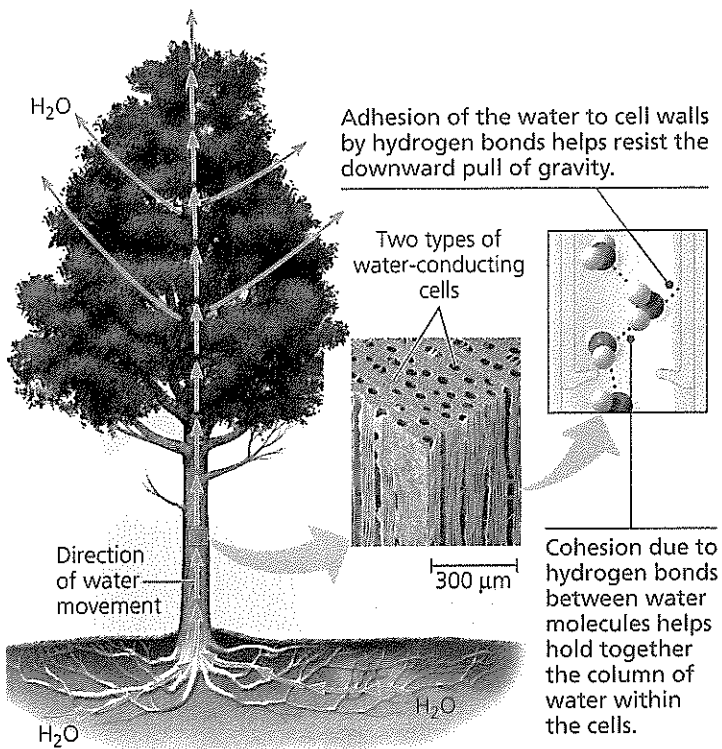
### Four emergent properties of water contribute to Earth's suitability for life

We will examine four emergent properties of water that contribute to Earth's suitability as an environment for life: cohesive behavior, ability to moderate temperature, expansion upon freezing, and versatility as a solvent.

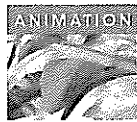
#### Cohesion of Water Molecules

Water molecules stay close to each other as a result of hydrogen bonding. Although the arrangement of molecules in a sample of liquid water is constantly changing, at any given moment many of the molecules are linked by multiple hydrogen bonds. These linkages make water more structured than most other liquids. Collectively, the hydrogen bonds hold the substance together, a phenomenon called **cohesion**.

Cohesion due to hydrogen bonding contributes to the transport of water and dissolved nutrients against gravity in plants. Water from the roots reaches the leaves through a network of water-conducting cells (Figure 3.3). As water evaporates from a leaf, hydrogen bonds cause water molecules leaving the veins to tug on molecules farther down, and the upward pull is transmitted through the water-conducting cells all the way to the roots. **Adhesion**, the clinging of one substance to another, also plays a role. Adhesion of



▲ **Figure 3.3 Water transport in plants.** Evaporation from leaves pulls water upward from the roots through water-conducting cells. Because of the properties of cohesion and adhesion, the tallest trees can transport water more than 100 m upward—approximately one-quarter the height of the Empire State Building in New York City.

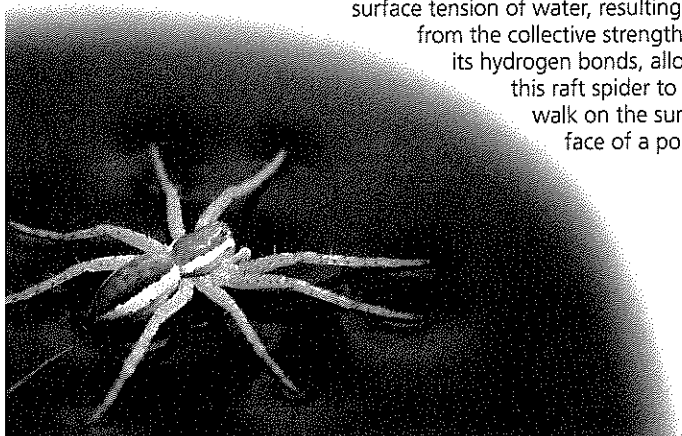


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water by hydrogen bonds to the molecules of cell walls helps counter the downward pull of gravity (see Figure 3.3).

Related to cohesion is **surface tension**, a measure of how difficult it is to stretch or break the surface of a liquid. At the interface between water and air is an ordered arrangement of water molecules, hydrogen-bonded to one another and to the water below. This gives water an unusually high surface tension, making it behave as though it were coated with an invisible film. You can observe the surface tension of water by slightly overfilling a drinking glass; the water will stand above the rim. The spider in **Figure 3.4** takes advantage of the surface tension of water to walk across a pond without breaking the surface.

▼ **Figure 3.4 Walking on water.** The high surface tension of water, resulting from the collective strength of its hydrogen bonds, allows this raft spider to walk on the surface of a pond.



## Moderation of Temperature by Water

Water moderates air temperature by absorbing heat from air that is warmer and releasing the stored heat to air that is cooler. Water is effective as a heat bank because it can absorb or release a relatively large amount of heat with only a slight change in its own temperature. To understand this capability of water, let's first look at temperature and heat.

### Temperature and Heat

Anything that moves has **kinetic energy**, the energy of motion. Atoms and molecules have kinetic energy because they are always moving, although not necessarily in any particular direction. The faster a molecule moves, the greater its kinetic energy. The kinetic energy associated with the random movement of atoms or molecules is called **thermal energy**. Thermal energy is related to temperature, but they are not the same thing. **Temperature** is a measure of energy that represents the *average* kinetic energy of the molecules in a body of matter, regardless of volume, whereas the *total* thermal energy depends in part on the matter's volume. When water is heated in a coffeemaker, the average speed of the molecules increases, and the thermometer records this as a rise in temperature of the liquid. The total amount of thermal energy also increases in this case. Note, however, that although the pot of coffee has a much higher temperature than, say, the water in a swimming pool, the swimming pool contains more thermal energy because of its much greater volume.

Whenever two objects of different temperature are brought together, thermal energy passes from the warmer to the cooler object until the two are the same temperature. Molecules in the cooler object speed up at the expense of the thermal energy of the warmer object. An ice cube cools a drink not by adding coldness to the liquid, but by absorbing thermal energy from the liquid as the ice itself melts. Thermal energy in transfer from one body of matter to another is defined as **heat**.

One convenient unit of heat used in this book is the **calorie (cal)**. A calorie is the amount of heat it takes to raise the temperature of 1 g of water by 1°C. Conversely, a calorie is also the amount of heat that 1 g of water releases when it cools by 1°C. A **kilocalorie (kcal)**, 1,000 cal, is the quantity of heat required to raise the temperature of 1 kilogram (kg) of water by 1°C. (The "calories" on food packages are actually kilocalories.) Another energy unit used in this book is the **joule (J)**. One joule equals 0.239 cal; one calorie equals 4.184 J.

### Water's High Specific Heat

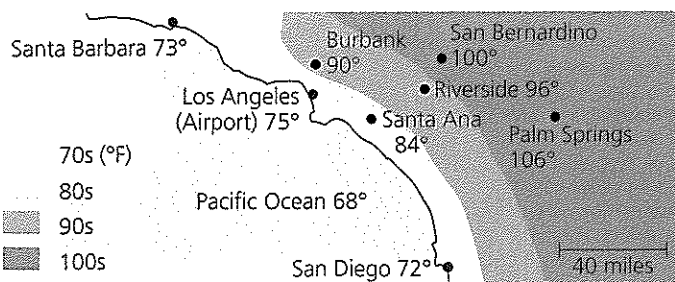
The ability of water to stabilize temperature stems from its relatively high specific heat. The **specific heat** of a substance is defined as the amount of heat that must be absorbed or lost for 1 g of that substance to change its temperature by 1°C. We

already know water's specific heat because we have defined a calorie as the amount of heat that causes 1 g of water to change its temperature by 1°C. Therefore, the specific heat of water is 1 calorie per gram and per degree Celsius, abbreviated as 1 cal/g · °C. Compared with most other substances, water has an unusually high specific heat. For example, ethyl alcohol, the type of alcohol in alcoholic beverages, has a specific heat of 0.6 cal/g · °C; that is, only 0.6 cal is required to raise the temperature of 1 g of ethyl alcohol by 1°C.

Because of the high specific heat of water relative to other materials, water will change its temperature less than other liquids when it absorbs or loses a given amount of heat. The reason you can burn your fingers by touching the side of an iron pot on the stove when the water in the pot is still lukewarm is that the specific heat of water is ten times greater than that of iron. In other words, the same amount of heat will raise the temperature of 1 g of the iron much faster than it will raise the temperature of 1 g of the water. Specific heat can be thought of as a measure of how well a substance resists changing its temperature when it absorbs or releases heat. Water resists changing its temperature; when it does change its temperature, it absorbs or loses a relatively large quantity of heat for each degree of change.

We can trace water's high specific heat, like many of its other properties, to hydrogen bonding. Heat must be absorbed in order to break hydrogen bonds; by the same token, heat is released when hydrogen bonds form. A calorie of heat causes a relatively small change in the temperature of water because much of the heat is used to disrupt hydrogen bonds before the water molecules can begin moving faster. And when the temperature of water drops slightly, many additional hydrogen bonds form, releasing a considerable amount of energy in the form of heat.

What is the relevance of water's high specific heat to life on Earth? A large body of water can absorb and store a huge amount of heat from the sun in the daytime and during summer while warming up only a few degrees. At night and during winter, the gradually cooling water can warm the air. This capability of water serves to moderate air temperatures in coastal areas (Figure 3.5). The high specific heat of water also



▲ **Figure 3.5** Temperatures for the Pacific Ocean and Southern California on an August day.

**INTERPRET THE DATA** Explain the pattern of temperatures shown in this diagram.

tends to stabilize ocean temperatures, creating a favorable environment for marine life. Thus, because of its high specific heat, the water that covers most of Earth keeps temperature fluctuations on land and in water within limits that permit life. Also, because organisms are made primarily of water, they are better able to resist changes in their own temperature than if they were made of a liquid with a lower specific heat.

### Evaporative Cooling

Molecules of any liquid stay close together because they are attracted to one another. Molecules moving fast enough to overcome these attractions can depart the liquid and enter the air as a gas (vapor). This transformation from a liquid to a gas is called vaporization, or evaporation. Recall that the speed of molecular movement varies and that temperature is the *average* kinetic energy of molecules. Even at low temperatures, the speediest molecules can escape into the air. Some evaporation occurs at any temperature; a glass of water at room temperature, for example, will eventually evaporate completely. If a liquid is heated, the average kinetic energy of molecules increases and the liquid evaporates more rapidly.

**Heat of vaporization** is the quantity of heat a liquid must absorb for 1 g of it to be converted from the liquid to the gaseous state. For the same reason that water has a high specific heat, it also has a high heat of vaporization relative to most other liquids. To evaporate 1 g of water at 25°C, about 580 cal of heat is needed—nearly double the amount needed to vaporize a gram of alcohol or ammonia. Water's high heat of vaporization is another emergent property resulting from the strength of its hydrogen bonds, which must be broken before the molecules can exit from the liquid in the form of water vapor (see Figure 3.1).

The high amount of energy required to vaporize water has a wide range of effects. On a global scale, for example, it helps moderate Earth's climate. A considerable amount of solar heat absorbed by tropical seas is consumed during the evaporation of surface water. Then, as moist tropical air circulates poleward, it releases heat as it condenses and forms rain. On an organismal level, water's high heat of vaporization accounts for the severity of steam burns. These burns are caused by the heat energy released when steam condenses into liquid on the skin.

As a liquid evaporates, the surface of the liquid that remains behind cools down (its temperature decreases). This **evaporative cooling** occurs because the "hottest" molecules, those with the greatest kinetic energy, are the most likely to leave as gas. It is as if the hundred fastest runners at a college transferred to another school; the average speed of the remaining students would decline.

Evaporative cooling of water contributes to the stability of temperature in lakes and ponds and also provides a mechanism that prevents terrestrial organisms from overheating. For example, evaporation of water from the leaves of a plant

helps keep the tissues in the leaves from becoming too warm in the sunlight. Evaporation of sweat from human skin dissipates body heat and helps prevent overheating on a hot day or when excess heat is generated by strenuous activity. High humidity on a hot day increases discomfort because the high concentration of water vapor in the air inhibits the evaporation of sweat from the body.

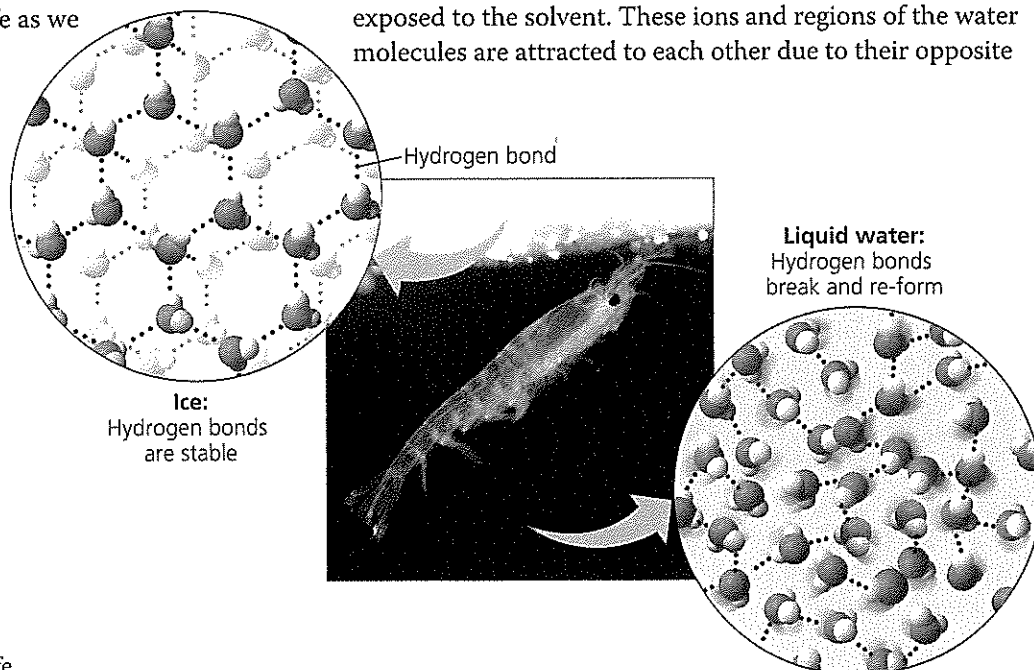
## Floating of Ice on Liquid Water

Water is one of the few substances that are less dense as a solid than as a liquid. In other words, ice floats on liquid water. While other materials contract and become denser when they solidify, water expands. The cause of this exotic behavior is, once again, hydrogen bonding. At temperatures above 4°C, water behaves like other liquids, expanding as it warms and contracting as it cools. As the temperature falls from 4°C to 0°C, water begins to freeze because more and more of its molecules are moving too slowly to break hydrogen bonds. At 0°C, the molecules become locked into a crystalline lattice, each water molecule hydrogen-bonded to four partners (**Figure 3.6**). The hydrogen bonds keep the molecules at “arm’s length,” far enough apart to make ice about 10% less dense (10% fewer molecules for the same volume) than liquid water at 4°C. When ice absorbs enough heat for its temperature to rise above 0°C, hydrogen bonds between molecules are disrupted. As the crystal collapses, the ice melts, and molecules are free to slip closer together. Water reaches its greatest density at 4°C and then begins to expand as the molecules move faster. Even in liquid water, many of the molecules are connected by hydrogen bonds, though only transiently: The hydrogen bonds are constantly breaking and re-forming.

The ability of ice to float due to its lower density is an important factor in the suitability of the environment for life. If ice sank, then eventually all ponds, lakes, and even oceans would freeze solid, making life as we know it impossible on Earth. During

► **Figure 3.6 Ice: crystalline structure and floating barrier.** In ice, each molecule is hydrogen-bonded to four neighbors in a three-dimensional crystal. Because the crystal is spacious, ice has fewer molecules than an equal volume of liquid water. In other words, ice is less dense than liquid water. Floating ice becomes a barrier that insulates the liquid water below from the colder air. The marine organism shown here is a type of shrimp called krill; it was photographed beneath floating ice in the Southern Ocean near Antarctica.

**WHAT IF?** If water did not form hydrogen bonds, what would happen to the shrimp’s habitat, shown here?



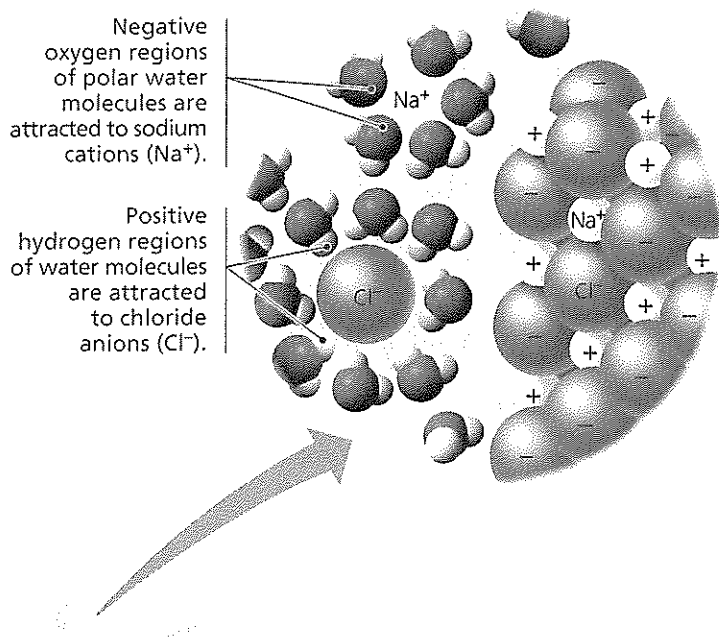
summer, only the upper few inches of the ocean would thaw. Instead, when a deep body of water cools, the floating ice insulates the liquid water below, preventing it from freezing and allowing life to exist under the frozen surface, as shown in the photo in **Figure 3.6**. Besides insulating the water below, ice also provides a solid habitat for some animals, such as polar bears and seals.

Many scientists are worried that these bodies of ice are at risk of disappearing. Global warming, which is caused by carbon dioxide and other “greenhouse” gases in the atmosphere, is having a profound effect on icy environments around the globe. In the Arctic, the average air temperature has risen 1.4°C just since 1961. This temperature increase has affected the seasonal balance between Arctic sea ice and liquid water, causing ice to form later in the year, to melt earlier, and to cover a smaller area. The rate at which glaciers and Arctic sea ice are disappearing is posing an extreme challenge to animals that depend on ice for their survival.

## Water: The Solvent of Life

A sugar cube placed in a glass of water will dissolve with a little stirring. The glass will then contain a uniform mixture of sugar and water; the concentration of dissolved sugar will be the same everywhere in the mixture. A liquid that is a completely homogeneous mixture of two or more substances is called a **solution**. The dissolving agent of a solution is the **solvent**, and the substance that is dissolved is the **solute**. In this case, water is the solvent and sugar is the solute. An **aqueous solution** is one in which the solute is dissolved in water; water is the solvent.

Water is a very versatile solvent, a quality we can trace to the polarity of the water molecule. Suppose, for example, that a spoonful of table salt, the ionic compound sodium chloride (NaCl), is placed in water (**Figure 3.7**). At the surface of each grain, or crystal, of salt, the sodium and chloride ions are exposed to the solvent. These ions and regions of the water molecules are attracted to each other due to their opposite



▲ **Figure 3.7 Table salt dissolving in water.** A sphere of water molecules, called a hydration shell, surrounds each solute ion.

**WHAT IF?** What would happen if you heated this solution for a long time?

charges. The oxygen regions of the water molecules are negatively charged and are attracted to sodium cations. The hydrogen regions are positively charged and are attracted to chloride anions. As a result, water molecules surround the individual sodium and chloride ions, separating and shielding them from one another. The sphere of water molecules around each dissolved ion is called a **hydration shell**. Working inward from the surface of each salt crystal, water eventually dissolves all the ions. The result is a solution of two solutes, sodium cations and chloride anions, homogeneously mixed with water, the solvent. Other ionic compounds also dissolve in water. Seawater, for instance, contains a great variety of dissolved ions, as do living cells.

A compound does not need to be ionic to dissolve in water; many compounds made up of nonionic polar molecules, such as the sugar in the sugar cube mentioned earlier, are also water-soluble. Such compounds dissolve when water molecules surround each of the solute molecules, forming hydrogen bonds with them. Even molecules as large as proteins can dissolve in water if they have ionic and polar regions on their surface (**Figure 3.8**). Many different kinds of polar

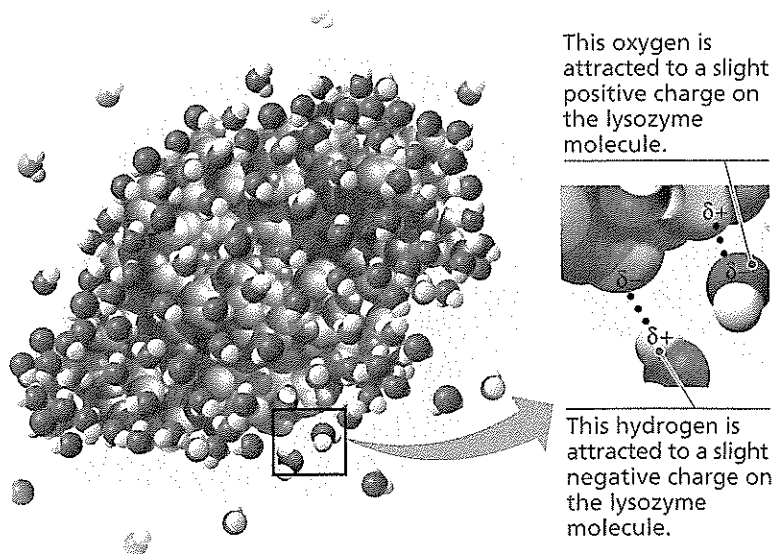
► **Figure 3.8 A water-soluble protein.** Human lysozyme is a protein found in tears and saliva that has antibacterial action. This model shows the lysozyme molecule (purple) in an aqueous environment. Ionic and polar regions on the protein's surface attract water molecules.

compounds are dissolved (along with ions) in the water of such biological fluids as blood, the sap of plants, and the liquid within all cells. Water is the solvent of life.

### Hydrophilic and Hydrophobic Substances

Any substance that has an affinity for water is said to be **hydrophilic** (from the Greek *hydro*, water, and *philos*, loving). In some cases, substances can be hydrophilic without actually dissolving. For example, some molecules in cells are so large that they do not dissolve. Another example of a hydrophilic substance that does not dissolve is cotton, a plant product. Cotton consists of giant molecules of cellulose, a compound with numerous regions of partial positive and partial negative charges that can form hydrogen bonds with water. Water adheres to the cellulose fibers. Thus, a cotton towel does a great job of drying the body, yet it does not dissolve in the washing machine. Cellulose is also present in the walls of water-conducting cells in a plant; you read earlier how the adhesion of water to these hydrophilic walls helps water move up the plant against gravity.

There are, of course, substances that do not have an affinity for water. Substances that are nonionic and nonpolar (or otherwise cannot form hydrogen bonds) actually seem to repel water; these substances are said to be **hydrophobic** (from the Greek *phobos*, fearing). An example from the kitchen is vegetable oil, which, as you know, does not mix stably with water-based substances such as vinegar. The hydrophobic behavior of the oil molecules results from a prevalence of relatively nonpolar covalent bonds, in this case bonds between carbon and hydrogen, which share electrons almost equally. Hydrophobic molecules related to oils are major ingredients of cell membranes. (Imagine what would happen to a cell if its membrane dissolved!)



## Solute Concentration in Aqueous Solutions

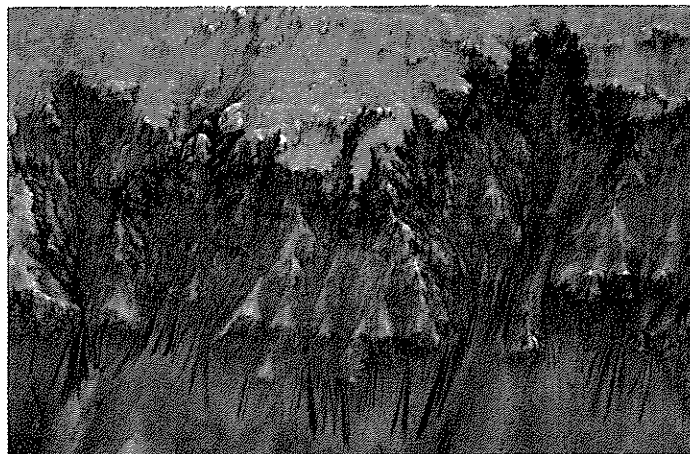
Most of the chemical reactions in organisms involve solutes dissolved in water. To understand such reactions, we must know how many atoms and molecules are involved and calculate the concentration of solutes in an aqueous solution (the number of solute molecules in a volume of solution).

When carrying out experiments, we use mass to calculate the number of molecules. We must first calculate the **molecular mass**, which is the sum of the masses of all the atoms in a molecule. As an example, let's calculate the molecular mass of table sugar (sucrose),  $C_{12}H_{22}O_{11}$ . In round numbers of daltons, the mass of a carbon atom is 12, the mass of a hydrogen atom is 1, and the mass of an oxygen atom is 16. Thus, sucrose has a molecular mass of  $(12 \times 12) + (22 \times 1) + (11 \times 16) = 342$  daltons. Because we can't weigh out small numbers of molecules, we usually measure substances in units called moles. Just as a dozen always means 12 objects, a **mole (mol)** represents an exact number of objects:  $6.02 \times 10^{23}$ , which is called Avogadro's number. Because of the way in which Avogadro's number and the unit *dalton* were originally defined, there are  $6.02 \times 10^{23}$  daltons in 1 g. Once we determine the molecular mass of a molecule such as sucrose, we can use the same number (342), but with the unit *gram*, to represent the mass of  $6.02 \times 10^{23}$  molecules of sucrose, or 1 mol of sucrose (this is sometimes called the *molar mass*). To obtain 1 mol of sucrose in the lab, therefore, we weigh out 342 g.

The practical advantage of measuring a quantity of chemicals in moles is that a mole of one substance has exactly the same number of molecules as a mole of any other substance. If the molecular mass of substance A is 342 daltons and that of substance B is 10 daltons, then 342 g of A will have the same number of molecules as 10 g of B. A mole of ethyl alcohol ( $C_2H_6O$ ) also contains  $6.02 \times 10^{23}$  molecules, but its mass is only 46 g because the mass of a molecule of ethyl alcohol is less than that of a molecule of sucrose. Measuring in moles makes it convenient for scientists working in the laboratory to combine substances in fixed ratios of molecules.

How would we make a liter (L) of solution consisting of 1 mol of sucrose dissolved in water? We would measure out 342 g of sucrose and then gradually add water, while stirring, until the sugar was completely dissolved. We would then add enough water to bring the total volume of the solution up to 1 L. At that point, we would have a 1-molar (1 M) solution of sucrose. **Molarity**—the number of moles of solute per liter of solution—is the unit of concentration most often used by biologists for aqueous solutions.

Water's capacity as a versatile solvent complements the other properties discussed in this chapter. Since these remarkable properties allow water to support life on Earth so well, scientists who seek life elsewhere in the universe look for water as a sign that a planet might sustain life.



▲ **Figure 3.9 Evidence for subsurface liquid water on Mars.** The dark streaks running down the lower portion of the photo are proposed to be streams of subsurface flowing water because they appear only during the warm season. The gullies in the middle of the photo could have been formed by flowing water.

## Possible Evolution of Life on Other Planets

**EVOLUTION** Biologists who look for life elsewhere in the universe (known as *astrobiologists*) have concentrated their search on planets that might have water. More than 800 planets have been found outside our solar system, and there is evidence for the presence of water vapor on a few of them. In our own solar system, Mars has been a focus of study. Like Earth, Mars has an ice cap at both poles. Images from spacecraft sent to Mars show that ice is present just under the surface of Mars and enough water vapor exists in its atmosphere for frost to form. **Figure 3.9** shows streaks that form along steep slopes during the Mars spring and summer, features that vanish during the winter. Some scientists have proposed that these are seasonal streams of flowing water occurring when subsurface ice melts during the warm season, while others think they are the result of  $CO_2$  rather than water. Drilling below the surface may be the next step in the search for signs of life on Mars. If any life-forms or fossils are found, their study will shed light on the process of evolution from an entirely new perspective.

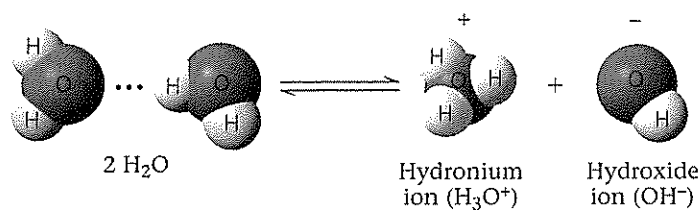
### CONCEPT CHECK 3.2

1. Describe how properties of water contribute to the upward movement of water in a tree.
2. Explain the saying "It's not the heat; it's the humidity."
3. How can the freezing of water crack boulders?
4. **WHAT IF?** A water strider (which can walk on water) has legs that are coated with a hydrophobic substance. What might be the benefit? What would happen if the substance were hydrophilic?
5. **INTERPRET THE DATA** The concentration of the appetite-regulating hormone ghrelin is about  $1.3 \times 10^{-10}$  M in the blood of a fasting person. How many molecules of ghrelin are in 1 L of blood?

For suggested answers, see Appendix A.

## Acidic and basic conditions affect living organisms

Occasionally, a hydrogen atom participating in a hydrogen bond between two water molecules shifts from one molecule to the other. When this happens, the hydrogen atom leaves its electron behind, and what is actually transferred is a **hydrogen ion** ( $H^+$ ), a single proton with a charge of  $1+$ . The water molecule that lost a proton is now a **hydroxide ion** ( $OH^-$ ), which has a charge of  $1-$ . The proton binds to the other water molecule, making that molecule a **hydronium ion** ( $H_3O^+$ ). We can picture the chemical reaction as follows:



By convention,  $H^+$  (the hydrogen ion) is used to represent  $H_3O^+$  (the hydronium ion), and we follow that practice in this book. Keep in mind, though, that  $H^+$  does not exist on its own in an aqueous solution. It is always associated with a water molecule in the form of  $H_3O^+$ .

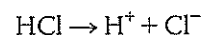
As indicated by the double arrows, this is a reversible reaction that reaches a state of dynamic equilibrium when water molecules dissociate at the same rate that they are being reformed from  $H^+$  and  $OH^-$ . At this equilibrium point, the concentration of water molecules greatly exceeds the concentrations of  $H^+$  and  $OH^-$ . In pure water, only one water molecule in every 554 million is dissociated; the concentration of each ion in pure water is  $10^{-7} M$  (at  $25^\circ C$ ). This means there is only one ten-millionth of a mole of hydrogen ions per liter of pure water and an equal number of hydroxide ions. (Even so, this is a huge number—over 60,000 *trillion*—of each ion.)

Although the dissociation of water is reversible and statistically rare, it is exceedingly important in the chemistry of life.  $H^+$  and  $OH^-$  are very reactive. Changes in their concentrations can drastically affect a cell's proteins and other complex molecules. As we have seen, the concentrations of  $H^+$  and  $OH^-$  are equal in pure water, but adding certain kinds of solutes, called acids and bases, disrupts this balance. Biologists use something called the pH scale to describe how acidic or basic (the opposite of acidic) a solution is. In the remainder of this chapter, you will learn about acids, bases, and pH and why changes in pH can adversely affect organisms.

### Acids and Bases

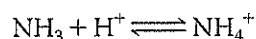
What would cause an aqueous solution to have an imbalance in  $H^+$  and  $OH^-$  concentrations? When acids dissolve in

water, they donate additional  $H^+$  to the solution. An **acid** is a substance that increases the hydrogen ion concentration of a solution. For example, when hydrochloric acid (HCl) is added to water, hydrogen ions dissociate from chloride ions:

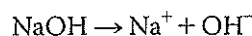


This source of  $H^+$  (dissociation of water is the other source) results in an acidic solution—one having more  $H^+$  than  $OH^-$ .

A substance that reduces the hydrogen ion concentration of a solution is called a **base**. Some bases reduce the  $H^+$  concentration directly by accepting hydrogen ions. Ammonia ( $NH_3$ ), for instance, acts as a base when the unshared electron pair in nitrogen's valence shell attracts a hydrogen ion from the solution, resulting in an ammonium ion ( $NH_4^+$ ):



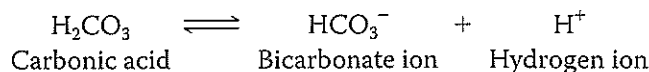
Other bases reduce the  $H^+$  concentration indirectly by dissociating to form hydroxide ions, which combine with hydrogen ions and form water. One such base is sodium hydroxide (NaOH), which in water dissociates into its ions:



In either case, the base reduces the  $H^+$  concentration. Solutions with a higher concentration of  $OH^-$  than  $H^+$  are known as basic solutions. A solution in which the  $H^+$  and  $OH^-$  concentrations are equal is said to be neutral.

Notice that single arrows were used in the reactions for HCl and NaOH. These compounds dissociate completely when mixed with water, so hydrochloric acid is called a strong acid and sodium hydroxide a strong base. In contrast, ammonia is a weak base. The double arrows in the reaction for ammonia indicate that the binding and release of hydrogen ions are reversible reactions, although at equilibrium there will be a fixed ratio of  $NH_4^+$  to  $NH_3$ .

Weak acids are acids that reversibly release and accept back hydrogen ions. An example is carbonic acid:



Here the equilibrium so favors the reaction in the left direction that when carbonic acid is added to pure water, only 1% of the molecules are dissociated at any particular time. Still, that is enough to shift the balance of  $H^+$  and  $OH^-$  from neutrality.

### The pH Scale

In any aqueous solution at  $25^\circ C$ , the *product* of the  $H^+$  and  $OH^-$  concentrations is constant at  $10^{-14}$ . This can be written

$$[H^+][OH^-] = 10^{-14}$$

In such an equation, brackets indicate molar concentration. In a neutral solution at  $25^\circ C$  (close to room temperature),  $[H^+] = 10^{-7}$  and  $[OH^-] = 10^{-7}$ , so in this case,  $10^{-14}$  is the



product of  $10^{-7}$  and  $10^{-7}$ . If enough acid is added to a solution to increase  $[H^+]$  to  $10^{-5} M$ , then  $[OH^-]$  will decline by an equivalent factor to  $10^{-9} M$  (note that  $10^{-5} \times 10^{-9} = 10^{-14}$ ). This constant relationship expresses the behavior of acids and bases in an aqueous solution. An acid not only adds hydrogen ions to a solution, but also removes hydroxide ions because of the tendency for  $H^+$  to combine with  $OH^-$ , forming water. A base has the opposite effect, increasing  $OH^-$  concentration but also reducing  $H^+$  concentration by the formation of water. If enough of a base is added to raise the  $OH^-$  concentration to  $10^{-4} M$ , it will cause the  $H^+$  concentration to drop to  $10^{-10} M$ . Whenever we know the concentration of either  $H^+$  or  $OH^-$  in an aqueous solution, we can deduce the concentration of the other ion.

Because the  $H^+$  and  $OH^-$  concentrations of solutions can vary by a factor of 100 trillion or more, scientists have developed a way to express this variation more conveniently than in moles per liter. The pH scale (Figure 3.10) compresses the range of  $H^+$  and  $OH^-$  concentrations by employing

logarithms. The **pH** of a solution is defined as the negative logarithm (base 10) of the hydrogen ion concentration:

$$\text{pH} = -\log [H^+]$$

For a neutral aqueous solution,  $[H^+]$  is  $10^{-7} M$ , giving us

$$-\log 10^{-7} = -(-7) = 7$$

Notice that pH *declines* as  $H^+$  concentration *increases*.

Notice, too, that although the pH scale is based on  $H^+$  concentration, it also implies  $OH^-$  concentration. A solution of pH 10 has a hydrogen ion concentration of  $10^{-10} M$  and a hydroxide ion concentration of  $10^{-4} M$ .

The pH of a neutral aqueous solution at 25°C is 7, the midpoint of the pH scale. A pH value less than 7 denotes an acidic solution; the lower the number, the more acidic the solution. The pH for basic solutions is above 7. Most biological fluids, such as blood and saliva, are within the range of pH 6–8. There are a few exceptions, however, including the strongly acidic digestive juice of the human stomach, which has a pH of about 2.

Remember that each pH unit represents a tenfold difference in  $H^+$  and  $OH^-$  concentrations. It is this mathematical feature that makes the pH scale so compact. A solution of pH 3 is not twice as acidic as a solution of pH 6, but a thousand times ( $10 \times 10 \times 10$ ) more acidic. When the pH of a solution changes slightly, the actual concentrations of  $H^+$  and  $OH^-$  in the solution change substantially.

## Buffers

The internal pH of most living cells is close to 7. Even a slight change in pH can be harmful, because the chemical processes of the cell are very sensitive to the concentrations of hydrogen and hydroxide ions. The pH of human blood is very close to 7.4, which is slightly basic. A person cannot survive for more than a few minutes if the blood pH drops

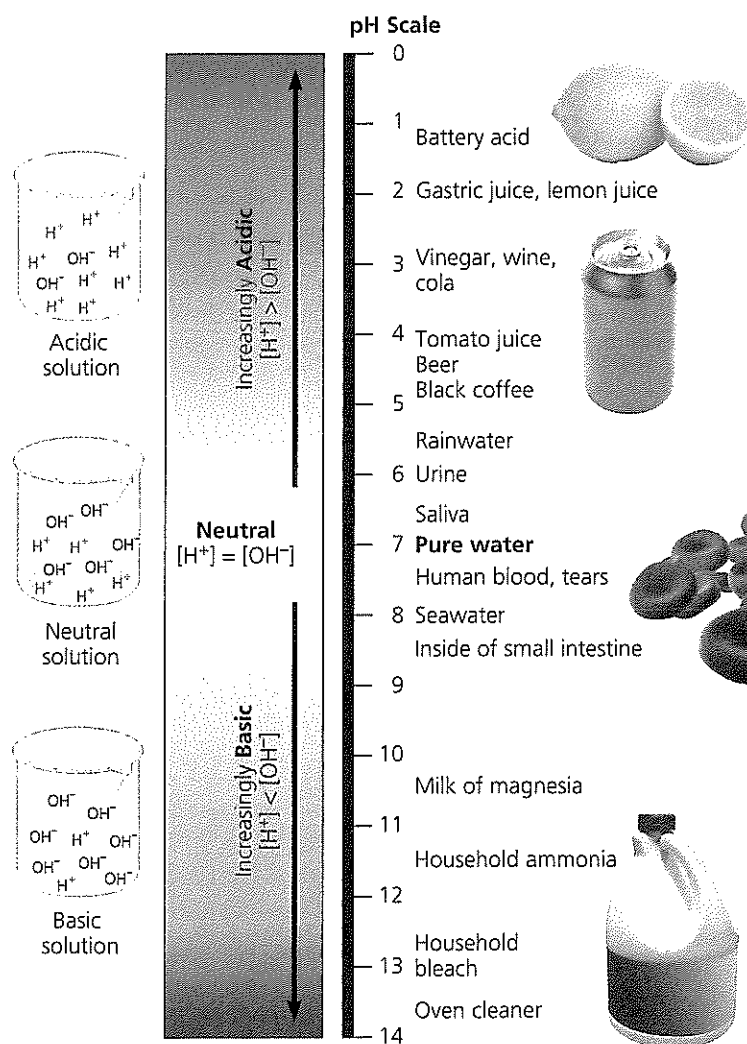
to 7 or rises to 7.8, and a chemical system exists in the

blood that maintains a stable pH. If 0.01 mol of a strong acid is added to a liter of pure water, the pH drops from 7.0 to 2.0. If the same amount of acid is added to a liter of blood, however, the pH decrease is only from 7.4 to 7.3. Why does the ad-

dition of acid have so much less of an effect on the pH of blood than it does on the pH of water?

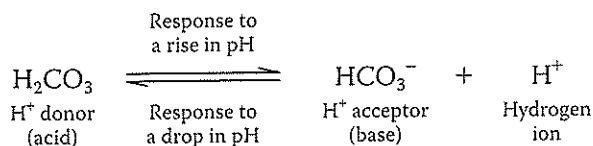
The presence of substances called buffers allows biological fluids to maintain a relatively constant pH despite the addition of acids or bases. A **buffer** is a substance that minimizes changes in the concentrations of  $H^+$  and  $OH^-$  in a solution. It does so by accepting hydrogen ions from the solution when they are in excess and donating hydrogen ions to the solution when they have been depleted. Most buffer solutions contain a weak acid and its corresponding base, which combine reversibly with hydrogen ions.

Several buffers contribute to pH stability in human blood and many other biological solutions. One of these is



▲ **Figure 3.10** The pH scale and pH values of some aqueous solutions.

carbonic acid ( $\text{H}_2\text{CO}_3$ ), which is formed when  $\text{CO}_2$  reacts with water in blood plasma. As mentioned earlier, carbonic acid dissociates to yield a bicarbonate ion ( $\text{HCO}_3^-$ ) and a hydrogen ion ( $\text{H}^+$ ):



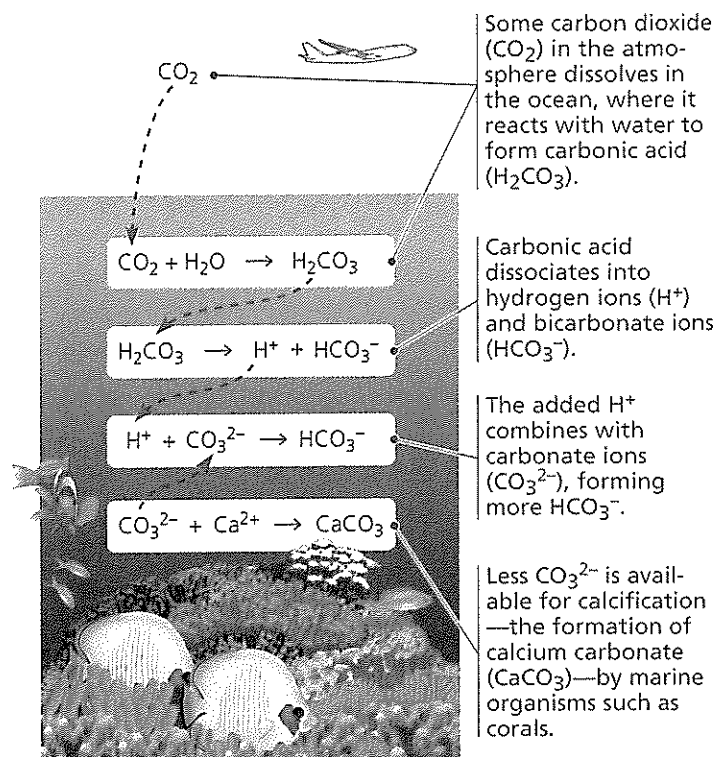
The chemical equilibrium between carbonic acid and bicarbonate acts as a pH regulator, the reaction shifting left or right as other processes in the solution add or remove hydrogen ions. If the  $\text{H}^+$  concentration in blood begins to fall (that is, if pH rises), the reaction proceeds to the right and more carbonic acid dissociates, replenishing hydrogen ions. But when the  $\text{H}^+$  concentration in blood begins to rise (when pH drops), the reaction proceeds to the left, with  $\text{HCO}_3^-$  (the base) removing the hydrogen ions from the solution and forming  $\text{H}_2\text{CO}_3$ . Thus, the carbonic acid–bicarbonate buffering system consists of an acid and a base in equilibrium with each other. Most other buffers are also acid–base pairs.

## Acidification: A Threat to Water Quality

Among the many threats to water quality posed by human activities is the burning of fossil fuels, which releases gaseous compounds into the atmosphere. When certain of these compounds react with water, the water becomes more acidic, altering the delicate balance of conditions for life on Earth. Carbon dioxide is the main product of fossil fuel combustion. About 25% of human-generated  $\text{CO}_2$  is absorbed by the oceans. In spite of the huge volume of water in the oceans, scientists worry that the absorption of so much  $\text{CO}_2$  will harm marine ecosystems.

Recent data have shown that such fears are well founded. When  $\text{CO}_2$  dissolves in seawater, it reacts with water to form carbonic acid, which lowers ocean pH, a process known as **ocean acidification**. Based on measurements of  $\text{CO}_2$  levels in air bubbles trapped in ice over thousands of years, scientists calculate that the pH of the oceans is 0.1 pH unit lower now than at any time in the past 420,000 years. Recent studies predict that it will drop another 0.3–0.5 pH unit by the end of this century.

As seawater acidifies, the extra hydrogen ions combine with carbonate ions ( $\text{CO}_3^{2-}$ ) to form bicarbonate ions ( $\text{HCO}_3^-$ ), thereby reducing the carbonate ion concentration (**Figure 3.11**). Scientists predict that ocean acidification will cause the carbonate ion concentration to decrease by 40% by the year 2100. This is of great concern because carbonate ions are required for calcification, the production of calcium carbonate ( $\text{CaCO}_3$ ) by many marine organisms, including reef-building corals and animals that build shells. The **Scientific Skills Exercise** allows you to work with data



▲ **Figure 3.11** Atmospheric  $\text{CO}_2$  from human activities and its fate in the ocean.

from an experiment examining the effect of carbonate ion concentration on coral reefs. Coral reefs are sensitive ecosystems that act as havens for a great diversity of marine life. The disappearance of coral reef ecosystems would be a tragic loss of biological diversity.

If there is any reason for optimism about the future quality of water resources on our planet, it is that we have made progress in learning about the delicate chemical balances in oceans, lakes, and rivers. Continued progress can come only from the actions of informed individuals, like yourselves, who are concerned about environmental quality. This requires understanding the crucial role that water plays in the suitability of the environment for continued life on Earth.

### CONCEPT CHECK 3.3

1. Compared with a basic solution at pH 9, the same volume of an acidic solution at pH 4 has \_\_\_\_\_ times as many hydrogen ions ( $\text{H}^+$ ).
2.  $\text{HCl}$  is a strong acid that dissociates in water:  $\text{HCl} \rightarrow \text{H}^+ + \text{Cl}^-$ . What is the pH of 0.01 M  $\text{HCl}$ ?
3. Acetic acid ( $\text{CH}_3\text{COOH}$ ) can be a buffer, similar to carbonic acid. Write the dissociation reaction, identifying the acid, base,  $\text{H}^+$  acceptor, and  $\text{H}^+$  donor.
4. **WHAT IF?** Given a liter of pure water and a liter solution of acetic acid, what would happen to the pH if you added 0.01 mol of a strong acid to each? Use the reaction equation from question 3 to explain the result.

For suggested answers, see Appendix A.

## SCIENTIFIC SKILLS EXERCISE

AP<sup>®</sup> SPs 1.4, 2.3, 6.4, 7.1

### Interpreting a Scatter Plot with a Regression Line

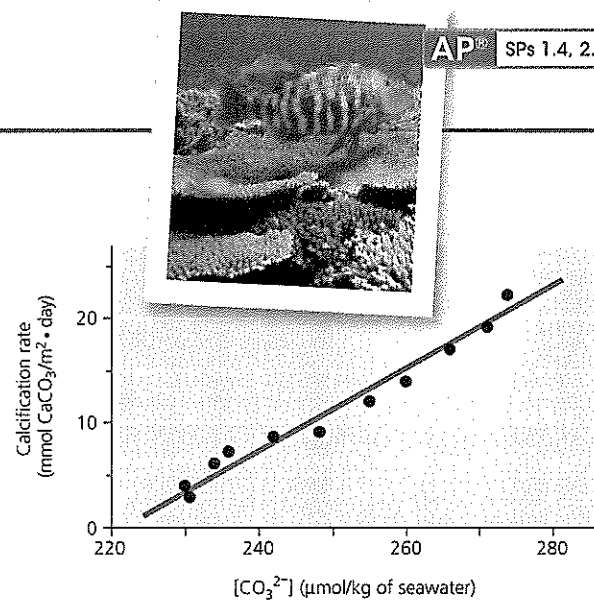
**How Does the Carbonate Ion Concentration of Seawater Affect the Calcification Rate of a Coral Reef?** Scientists predict that acidification of the ocean due to higher levels of atmospheric  $\text{CO}_2$  will lower the concentration of dissolved carbonate ions, which living corals use to build calcium carbonate reef structures. In this exercise, you will analyze data from a controlled experiment that examined the effect of carbonate ion concentration ( $[\text{CO}_3^{2-}]$ ) on calcium carbonate deposition, a process called calcification.

**How the Experiment Was Done** The Biosphere 2 aquarium in Arizona contains a large coral reef system that behaves like a natural reef. For several years, a group of researchers measured the rate of calcification by the reef organisms and examined how the calcification rate changed with differing amounts of dissolved carbonate ions in the seawater.

**Data from the Experiment** The black data points in the graph form a scatter plot. The red line, known as a linear regression line, is the best-fitting straight line for these points.

#### Interpret the Data

- When presented with a graph of experimental data, the first step in analysis is to determine what each axis represents. (a) In words, explain what is being shown on the x-axis. Be sure to include the units. (b) What is being shown on the y-axis (including units)? (c) Which variable is the independent variable—the variable that was *manipulated* by the researchers? (d) Which variable is the dependent variable—the variable that responded to or depended on the treatment, which was *measured* by the researchers? (For additional information about graphs, see the Scientific Skills Review in Appendix F and in the Study Area in MasteringBiology.)
- Based on the data shown in the graph, describe in words the relationship between carbonate ion concentration and calcification rate.
- (a) If the seawater carbonate ion concentration is  $270 \mu\text{mol/kg}$ , what is the approximate rate of calcification, and approximately how many days would it take 1 square meter of reef to accumulate 30 mmol of



- (a) Referring to the equations in Figure 3.11, determine which step of the process is measured in this experiment. (b) Are the results of this experiment consistent with the hypothesis that increased atmospheric  $[\text{CO}_2]$  will slow the growth of coral reefs? Why or why not?

**MB** A version of this Scientific Skills Exercise can be assigned in MasteringBiology.

**Data from** C. Langdon et al., Effect of calcium carbonate saturation state on the calcification rate of an experimental coral reef, *Global Biogeochemical Cycles* 14:639–654 (2000).

## 3 Chapter Review

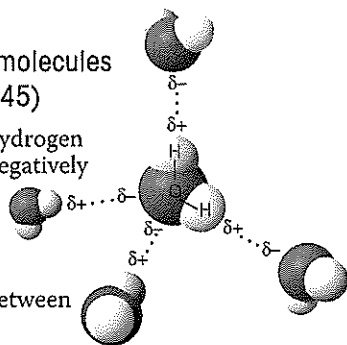
AP<sup>®</sup> How do the polar covalent bonds that form between oxygen and hydrogen atoms in a water molecule contribute to all the different emergent properties of water upon which living systems depend? (**Big Idea 2**)

### SUMMARY OF KEY CONCEPTS

#### CONCEPT 3.1

Polar covalent bonds in water molecules result in hydrogen bonding (p. 45)

- Water is a **polar molecule**. A hydrogen bond forms when the slightly negatively charged oxygen of one water molecule is attracted to the slightly positively charged hydrogen of a nearby water molecule. Hydrogen bonding between water molecules is the basis for water's properties.



**DRAW IT** Label a hydrogen bond and a polar covalent bond in this figure. Is a hydrogen bond a covalent bond? Explain.

#### CONCEPT 3.2

Four emergent properties of water contribute to Earth's suitability for life (pp. 45–50)

- Hydrogen bonding keeps water molecules close to each other, and this **cohesion** helps pull water upward in the microscopic water-conducting cells of plants. Hydrogen bonding is also responsible for water's **surface tension**.
- Water has a high **specific heat**: Heat is absorbed when hydrogen bonds break and is released when hydrogen bonds form. This helps keep temperatures relatively steady, within limits that permit life. **Evaporative cooling** is based on water's high **heat of vaporization**. The evaporative loss of the most energetic water molecules cools a surface.
- Ice floats because it is less dense than liquid water. This property allows life to exist under the frozen surfaces of lakes and polar seas.
- Water is an unusually versatile **solvent** because its polar molecules are attracted to ions and polar substances that can form

hydrogen bonds. **Hydrophilic** substances have an affinity for water; **hydrophobic** substances do not. **Molarity**, the number of moles of **solute** per liter of **solution**, is used as a measure of solute concentration in solutions. A **mole** is a certain number of molecules of a substance. The mass of a mole of a substance in grams is the same as the **molecular mass** in daltons.

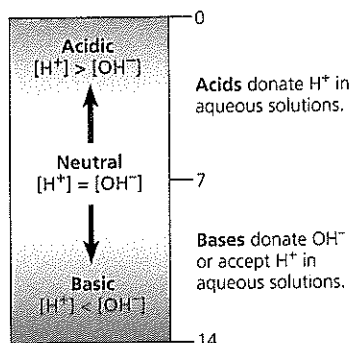
- The emergent properties of water support life on Earth and may contribute to the potential for life to have evolved on other planets.

2 Describe how different types of solutes dissolve in water. Explain what a solution is.

### CONCEPT 3.3

#### Acidic and basic conditions affect living organisms (pp. 51–54)

- A water molecule can transfer an  $H^+$  to another water molecule to form  $H_3O^+$  (represented simply by  $H^+$ ) and  $OH^-$ .
- The concentration of  $H^+$  is expressed as **pH**;  $pH = -\log [H^+]$ . A **buffer** consists of an acid-base pair that combines reversibly with hydrogen ions, allowing it to resist pH changes.
- The burning of fossil fuels increases the amount of  $CO_2$  in the atmosphere. Some  $CO_2$  dissolves in the oceans, causing **ocean acidification**, which has potentially grave consequences for coral reefs.



2 Explain how increasing amounts of  $CO_2$  dissolving in the ocean leads to ocean acidification. How does this change in pH affect carbonate ion concentration and the rate of calcification?

### TEST YOUR UNDERSTANDING

#### LEVEL 1: KNOWLEDGE/COMPREHENSION

- Which of the following is a hydrophobic material?
 

(A) paper	(C) wax
(B) table salt	(D) sugar
- We can be sure that a mole of table sugar and a mole of vitamin C are equal in their
 

(A) mass.	(C) number of atoms.
(B) volume.	(D) number of molecules.
- Measurements show that the pH of a particular lake is 4.0. What is the hydrogen ion concentration of the lake?
 

(A) $4.0 M$	(C) $10^{-4} M$
(B) $10^{-10} M$	(D) $10^4 M$
- What is the *hydroxide* ion concentration of the lake described in question 3?
 

(A) $10^{-10} M$	(C) $10^{-7} M$
(B) $10^{-4} M$	(D) $10.0 M$

#### LEVEL 2: APPLICATION/ANALYSIS

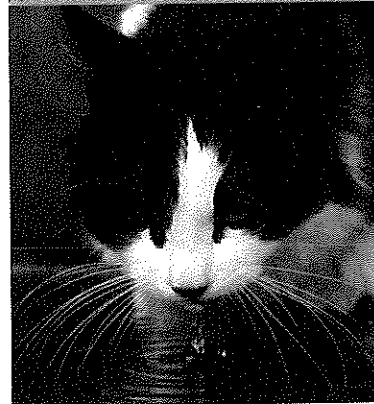
- A slice of pizza has 500 kcal. If we could burn the pizza and use all the heat to warm a 50-L container of cold water, what would be the approximate increase in the temperature of the water? (Note: A liter of cold water weighs about 1 kg.)
 

(A) $50^\circ C$	(C) $100^\circ C$
(B) $5^\circ C$	(D) $10^\circ C$

- DRAW IT** Draw the hydration shells that form around a potassium ion and a chloride ion when potassium chloride (KCl) dissolves in water. Label the positive, negative, and partial charges on the atoms.

#### LEVEL 3: SYNTHESIS/EVALUATION AP<sup>®</sup>

- SCIENTIFIC INQUIRY/Science Practice 7**  
In agricultural areas, farmers pay close attention to the weather forecast. Right before a predicted overnight freeze, farmers spray water on crops to protect the plants. Use the properties of water to **explain** how this method works. Be sure to mention why hydrogen bonds are responsible for this phenomenon.
- CONNECT TO BIG IDEA 1**  
This chapter explains how the emergent properties of water contribute to the suitability of the environment for life. Until fairly recently, scientists assumed that other physical requirements for life included a moderate range of temperature, pH, atmospheric pressure, and salinity, as well as low levels of toxic chemicals. That view has changed with the discovery of organisms known as extremophiles, which have been found flourishing in hot, acidic sulfur springs, around hydrothermal vents deep in the ocean, and in soils with high levels of toxic metals. **Explain** why astrobiologists would be interested in studying extremophiles. **Speculate** as to what the existence of life in such extreme environments says about the possibility of life on other planets.
- SCIENTIFIC INQUIRY/Science Practice 4**  
**Design** a controlled experiment to test the hypothesis that water acidification caused by acidic rain would inhibit the growth of *Elodea*, a freshwater plant (see Figure 2.17).
- CONNECT TO BIG IDEA 4**  
Several emergent properties of water contribute to the suitability of the environment for life. In a short essay (100–150 words), **describe** how the ability of water to function as a versatile solvent arises from the structure of water molecules.
- SYNTHESIZE YOUR KNOWLEDGE**



**SCIENTIFIC INQUIRY/ Science Practice 6**  
How do cats drink? While dogs form their tongues into spoons and scoop water into their mouths, scientists using high-speed video have shown that cats use a different technique. Four times a second, the cat touches the tip of its tongue to the water and draws a column

of water up into its mouth (as seen in the photo), which then shuts before gravity can pull the water back down. **Describe** how the properties of water allow cats to drink in this fashion, including how water's molecular structure contributes to the process.

For selected answers, see Appendix A.

### MasteringBiology<sup>®</sup>

**Students** Go to **MasteringBiology** for assignments, the eText, and the Study Area with practice tests, animations, and activities.

**Instructors** Go to **MasteringBiology** for automatically graded tutorials and questions that you can assign to your students, plus Instructor Resources.

# 4

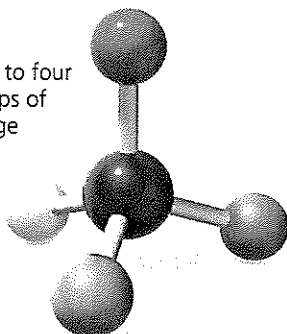
## Carbon and the Molecular Diversity of Life

### KEY CONCEPTS

- 4.1 Organic chemistry is the study of carbon compounds
- 4.2 Carbon atoms can form diverse molecules by bonding to four other atoms
- 4.3 A few chemical groups are key to molecular function

**AP** **BIG IDEAS:** Carbon is one of the key elements around which living matter is organized and core biological processes have evolved (Big Idea 1), plays a vital role in the capture, use, and storage of the energy needed to support life (Big Idea 2), and it serves as the backbone around which a great diversity of biomolecules are synthesized and then cycle through the environment (Big Idea 4).

► Carbon can bond to four other atoms or groups of atoms, making a large variety of molecules possible.



▲ **Figure 4.1** What properties make carbon the basis of all life?

### Carbon: The Backbone of Life

Living organisms, such as the plants and the Qinling golden snub-nosed monkeys shown in **Figure 4.1**, are made up of chemicals based mostly on the element carbon. Carbon enters the biosphere through the action of plants and other photosynthetic organisms. Plants use solar energy to transform atmospheric  $\text{CO}_2$  into the molecules of life, which are then taken in by plant-eating animals.

Of all the chemical elements, carbon is unparalleled in its ability to form molecules that are large, complex, and varied, making possible the diversity of organisms that have evolved on Earth. Proteins, DNA, carbohydrates, and other molecules that distinguish living matter from inanimate material are all composed of carbon atoms bonded to one another and to atoms of other elements. Hydrogen (H), oxygen (O), nitrogen (N), sulfur (S), and phosphorus (P) are other common ingredients of these compounds, but it is the element carbon (C) that accounts for the enormous variety of biological molecules.

Large biological molecules, such as proteins, are the main focus of Chapter 5. In this chapter, we investigate the properties of smaller molecules. We will use these small molecules to illustrate concepts of molecular architecture that will help explain why carbon is so important to life, at the same time highlighting the theme that emergent properties arise from the organization of matter in living organisms.

## CONCEPT 4.1

### Organic chemistry is the study of carbon compounds

For historical reasons, compounds containing carbon are said to be organic, and their study is called **organic chemistry**. By the early 1800s, chemists had learned to make simple compounds in the laboratory by combining elements under the right conditions. Artificial synthesis of the complex molecules extracted from living matter seemed impossible, however. Organic compounds were thought to arise only in living organisms, which were believed to contain a life force beyond the jurisdiction of physical and chemical laws.

Chemists began to chip away at this notion when they learned to synthesize organic compounds in the laboratory. In 1828, Friedrich Wöhler, a German chemist, tried to make an “inorganic” salt, ammonium cyanate, by mixing solutions of ammonium ions ( $\text{NH}_4^+$ ) and cyanate ions ( $\text{CNO}^-$ ). Wöhler was astonished to find that instead he had made urea, an organic compound present in the urine of animals.

The next few decades saw laboratory synthesis of increasingly complex organic compounds, supporting the view that physical and chemical laws govern the processes of life. Organic chemistry was redefined as the study of carbon compounds, regardless of origin. Organic compounds range from simple molecules, such as methane ( $\text{CH}_4$ ), to colossal ones, such as proteins, with thousands of atoms.

### Organic Molecules and the Origin of Life on Earth

**EVOLUTION** In 1953, Stanley Miller, a graduate student of Harold Urey’s at the University of Chicago, helped bring the abiotic (nonliving) synthesis of organic compounds into the context of evolution. Study **Figure 4.2** to learn about his classic experiment. From his results, Miller concluded that complex organic molecules could arise spontaneously under conditions thought at that time to have existed on the early Earth. You can work with the data from a related experiment in the **Scientific Skills Exercise**. These experiments support the idea that abiotic synthesis of organic compounds, perhaps near volcanoes, could have been an early stage in the origin of life (see Chapter 25).

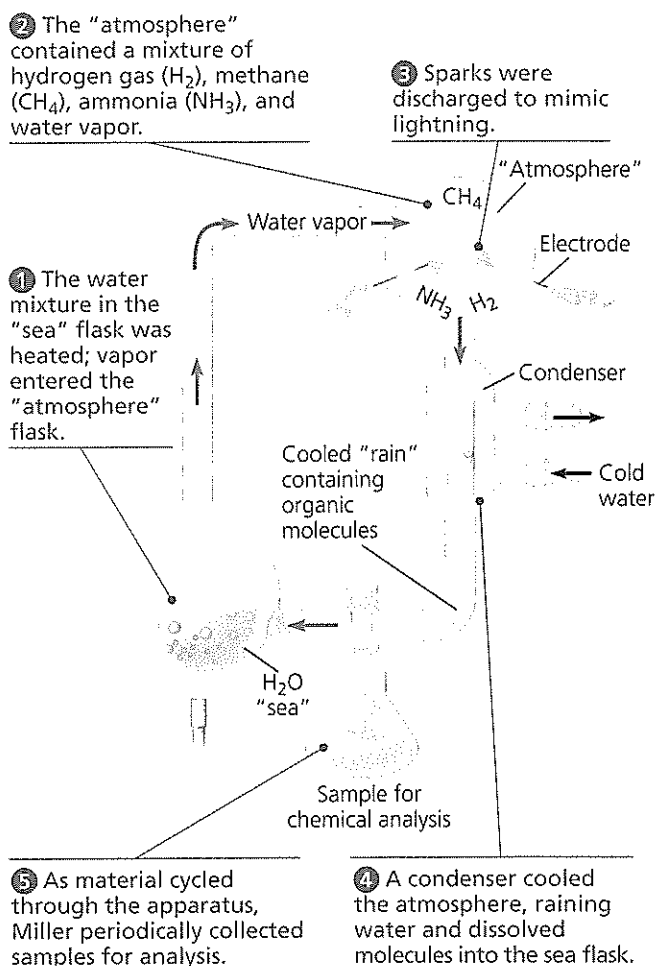
The overall percentages of the major elements of life—C, H, O, N, S, and P—are quite uniform from one organism to another, reflecting the common evolutionary origin of all life. Because of carbon’s ability to form four bonds, however, this limited assortment of atomic building blocks can be used to build an inexhaustible variety of organic molecules. Different species of organisms, and different individuals within a species, are distinguished by variations in the types

## ▼ Figure 4.2

## Inquiry

### Can organic molecules form under conditions estimated to simulate those on the early Earth?

**Experiment** In 1953, Stanley Miller set up a closed system to mimic conditions thought at that time to have existed on the early Earth. A flask of water simulated the primeval sea. The water was heated so that some vaporized and moved into a second, higher flask containing the “atmosphere”—a mixture of gases. Sparks were discharged in the synthetic atmosphere to mimic lightning.



**Results** Miller identified a variety of organic molecules that are common in organisms. These included simple compounds, such as formaldehyde ( $\text{CH}_2\text{O}$ ) and hydrogen cyanide ( $\text{HCN}$ ), and more complex molecules, such as amino acids and long chains of carbon and hydrogen known as hydrocarbons.

**Conclusion** Organic molecules, a first step in the origin of life, may have been synthesized abiotically on the early Earth. Although new evidence indicates that the early Earth’s atmosphere was different from the “atmosphere” used by Miller in this experiment, recent experiments using the revised list of chemicals also produced organic molecules. (We will explore this hypothesis in more detail in Chapter 25.)

*Source:* S. L. Miller, A production of amino acids under possible primitive Earth conditions, *Science* 117:528–529 (1953).

**WHAT IF?** If Miller had increased the concentration of  $\text{NH}_3$  in his experiment, how might the relative amounts of the products  $\text{HCN}$  and  $\text{CH}_2\text{O}$  have differed?

## Working with Moles and Molar Ratios

**Could the First Biological Molecules Have Formed Near Volcanoes on Early Earth?** In 2007, Jeffrey Bada, a former graduate student of Stanley Miller's, discovered some vials of samples that had never been analyzed from an experiment performed by Miller in 1958. In this experiment, Miller used hydrogen sulfide gas ( $\text{H}_2\text{S}$ ) as one of the gases in the reactant mixture. Since  $\text{H}_2\text{S}$  is released by volcanoes, the  $\text{H}_2\text{S}$  experiment was designed to mimic conditions near volcanoes on early Earth. In 2011, Bada and colleagues published the results of their analysis of these "lost" samples. In this exercise, you will make calculations using the molar ratios of reactants and products from the  $\text{H}_2\text{S}$  experiment.

**How the Experiment Was Done** According to his laboratory notebook, Miller used the same apparatus as in his original experiment (see Figure 4.2), but the mixture of gaseous reactants included methane ( $\text{CH}_4$ ), carbon dioxide ( $\text{CO}_2$ ), hydrogen sulfide ( $\text{H}_2\text{S}$ ), and ammonia ( $\text{NH}_3$ ). After three days of simulated volcanic activity, he collected samples of the liquid, partially purified the chemicals, and sealed the samples in sterile vials. In 2011, Bada's research team used modern analytical methods to analyze the products in the vials for the presence of amino acids, the building blocks of proteins.

**Data from the Experiment** The table below shows 4 of the 23 amino acids detected in the samples from Miller's 1958  $\text{H}_2\text{S}$  experiment.

Product Compound	Molecular Formula	Molar Ratio (Relative to Glycine)
Glycine	$\text{C}_2\text{H}_5\text{NO}_2$	1.0
Serine	$\text{C}_3\text{H}_7\text{NO}_3$	$3.0 \times 10^{-2}$
Methionine	$\text{C}_5\text{H}_{11}\text{NO}_2\text{S}$	$1.8 \times 10^{-3}$
Alanine	$\text{C}_3\text{H}_7\text{NO}_2$	1.1

## Interpret the Data

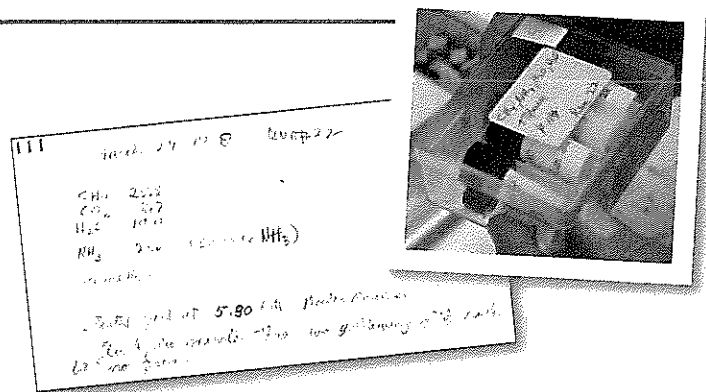
1. A *mole* is the number of grams of a substance that equals its molecular (or atomic) mass in daltons. There are  $6.02 \times 10^{23}$  molecules (or atoms) in 1.0 mole (Avogadro's number; see Concept 3.2). The data table shows the "molar ratios" of some of the products from the Miller  $\text{H}_2\text{S}$  experiment. In a molar ratio, each unitless value is expressed relative to a standard for that experiment. Here, the standard is the number of moles of the amino acid glycine, which is set to a value of 1.0. For instance, serine has a molar ratio of  $3.0 \times 10^{-2}$ , meaning that for every mole of glycine, there is  $3.0 \times 10^{-2}$  mole of serine. (a) Give the molar ratio of methionine to glycine and explain what it means. (b) How many molecules of glycine are present in 1.0 mole? (c) For every 1.0 mole of glycine in the sample, how many molecules of methionine are present? (Recall that to multiply two

of organic molecules they make. In a sense, the great diversity of living organisms we see on the planet (and in fossil remains) is made possible by the unique chemical versatility of the element carbon.

## CONCEPT CHECK 4.1

1. Why was Wöhler astonished to find he had made urea?
2. **WHAT IF?** Miller carried out a control experiment without the electrical discharge and found no organic compounds. What might explain this result?

For suggested answers, see Appendix A.



▲ Some of Stanley Miller's notes from his 1958 hydrogen sulfide ( $\text{H}_2\text{S}$ ) experiment along with his original vials.

- numbers with exponents, you add their exponents; to divide them, you subtract the exponent in the denominator from that in the numerator.)
2. (a) Which amino acid is present in higher amounts than glycine? (b) How many more molecules of that amino acid are present than the number of molecules in 1.0 mole of glycine?
  3. The synthesis of products is limited by the amount of reactants. (a) If one mole each of  $\text{CH}_4$ ,  $\text{NH}_3$ ,  $\text{H}_2\text{S}$ , and  $\text{CO}_2$  is added to 1 liter of water (= 55.5 moles of  $\text{H}_2\text{O}$ ) in a flask, how many moles of hydrogen, carbon, oxygen, nitrogen, and sulfur are in the flask? (b) Looking at the molecular formula in the table, how many moles of each element would be needed to make 1.0 mole of glycine? (c) What is the maximum number of moles of glycine that could be made in that flask, with the specified ingredients, if no other molecules were made? Explain. (d) If serine or methionine were made individually, which element(s) would be used up first for each? How much of each product could be made?
  4. The earlier published experiment carried out by Miller did not include  $\text{H}_2\text{S}$  in the reactants (see Figure 4.2). Which of the compounds shown in the data table can be made in the  $\text{H}_2\text{S}$  experiment but could not be made in the earlier experiment?

**MB** A version of this Scientific Skills Exercise can be assigned in MasteringBiology.

**Data from** E. T. Parker et al., Primordial synthesis of amines and amino acids in a 1958 Miller  $\text{H}_2\text{S}$ -rich spark discharge experiment, *Proceedings of the National Academy of Sciences USA* 108:5526-5531 (2011). [www.pnas.org/cgi/doi/10.1073/pnas.1019191108](http://www.pnas.org/cgi/doi/10.1073/pnas.1019191108).

## CONCEPT 4.2

## Carbon atoms can form diverse molecules by bonding to four other atoms

The key to an atom's chemical characteristics is its electron configuration. This configuration determines the kinds and number of bonds an atom will form with other atoms. Recall that it is the valence electrons, those in the outermost shell, that are available to form bonds with other atoms.

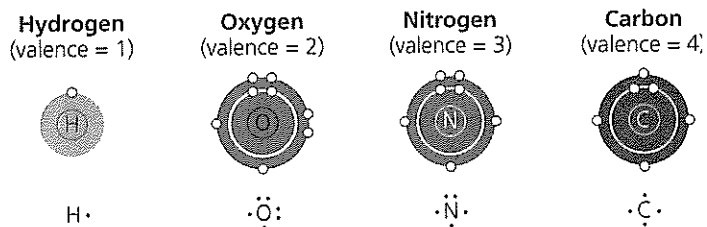
Molecule and Molecular Shape	Molecular Formula	Structural Formula	Ball-and-Stick Model (molecular shape in pink)	Space-Filling Model
(a) <b>Methane.</b> When a carbon atom has four single bonds to other atoms, the molecule is tetrahedral.	CH <sub>4</sub>	<pre>       H         H — C — H               H           </pre>		
(b) <b>Ethane.</b> A molecule may have more than one tetrahedral group of single-bonded atoms. (Ethane consists of two such groups.)	C <sub>2</sub> H <sub>6</sub>	<pre>       H   H             H — C — C — H                   H   H           </pre>		
(c) <b>Ethene (ethylene).</b> When two carbon atoms are joined by a double bond, all atoms attached to those carbons are in the same plane, and the molecule is flat.	C <sub>2</sub> H <sub>4</sub>	<pre>       H   H        \ /         C = C        / \       H   H           </pre>		

▲ **Figure 4.3** The shapes of three simple organic molecules.

## The Formation of Bonds with Carbon

Carbon has 6 electrons, with 2 in the first electron shell and 4 in the second shell; thus, it has 4 valence electrons in a shell that can hold up to 8 electrons. A carbon atom usually completes its valence shell by sharing its 4 electrons with other atoms so that 8 electrons are present. Each pair of shared electrons constitutes a covalent bond (see Figure 2.10d). In organic molecules, carbon usually forms single or double covalent bonds. Each carbon atom acts as an intersection point from which a molecule can branch off in as many as four directions. This enables carbon to form large, complex molecules.

When a carbon atom forms four single covalent bonds, the arrangement of its four hybrid orbitals causes the bonds to angle toward the corners of an imaginary tetrahedron. The bond angles in methane (CH<sub>4</sub>) are 109.5° (**Figure 4.3a**), and they are roughly the same in any group of atoms where carbon has four single bonds. For example, ethane (C<sub>2</sub>H<sub>6</sub>) is shaped like two overlapping tetrahedrons (**Figure 4.3b**). In molecules with more carbons, every grouping of a carbon bonded to four other atoms has a tetrahedral shape. But when two carbon atoms are joined by a double bond, as in ethene (C<sub>2</sub>H<sub>4</sub>), the bonds from both carbons are all in the same plane, so the atoms joined to those carbons are in the same plane as well (**Figure 4.3c**). We find it convenient to write molecules as structural formulas, as if the molecules being represented are two-dimensional, but keep in mind that molecules are three-dimensional and that the shape of a molecule is central to its function.



▲ **Figure 4.4** Valences of the major elements of organic molecules. Valence is the number of covalent bonds an atom can form. It is generally equal to the number of electrons required to complete the valence (outermost) shell (see Figure 2.7). All the electrons are shown for each atom in the electron distribution diagrams (top). Only the valence shell electrons are shown in the Lewis dot structures (bottom). Note that carbon can form four bonds.

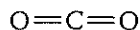
**MAKE CONNECTIONS** Draw the Lewis dot structures for sodium, phosphorus, sulfur, and chlorine. (Refer to Figure 2.7.)

The electron configuration of carbon gives it covalent compatibility with many different elements. **Figure 4.4** shows the valences of carbon and its most frequent bonding partners—hydrogen, oxygen, and nitrogen. These are the four major atomic components of organic molecules. These valences are the basis for the rules of covalent bonding in organic chemistry—the building code for the architecture of organic molecules.

How do the rules of covalent bonding apply to carbon atoms with partners other than hydrogen? We'll look at two examples, the simple molecules carbon dioxide and urea.

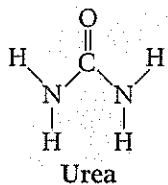


In the carbon dioxide molecule ( $\text{CO}_2$ ), a single carbon atom is joined to two atoms of oxygen by double covalent bonds. The structural formula for  $\text{CO}_2$  is shown here:



Each line in a structural formula represents a pair of shared electrons. Thus, the two double bonds in  $\text{CO}_2$  have the same number of shared electrons as four single bonds. The arrangement completes the valence shells of all atoms in the molecule. Because  $\text{CO}_2$  is a very simple molecule and lacks hydrogen, it is often considered inorganic, even though it contains carbon. Whether we call  $\text{CO}_2$  organic or inorganic, however, it is clearly important to the living world as the source of carbon for all organic molecules in organisms.

Urea,  $\text{CO}(\text{NH}_2)_2$ , is the organic compound found in urine that Wöhler synthesized in the early 1800s. Again, each atom has the required number of covalent bonds. In this case, one carbon atom participates in both single and double bonds.



Urea and carbon dioxide are molecules with only one carbon atom. But as Figure 4.3 shows, a carbon atom can also use one or more valence electrons to form covalent bonds to other carbon atoms, each of which can also form four bonds. Thus, the atoms can be linked into chains of seemingly infinite variety.

## Molecular Diversity Arising from Variation in Carbon Skeletons

Carbon chains form the skeletons of most organic molecules. The skeletons vary in length and may be straight, branched, or arranged in closed rings (Figure 4.5). Some carbon skeletons have double bonds, which vary in number and location. Such variation in carbon skeletons is one important source of the molecular complexity and diversity that characterize living matter. In addition, atoms of other elements can be bonded to the skeletons at available sites.

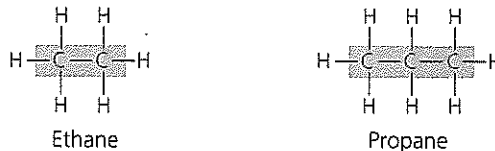
### Hydrocarbons

All of the molecules that are shown in Figures 4.3 and 4.5 are **hydrocarbons**, organic molecules consisting of only carbon and hydrogen. Atoms of hydrogen are attached to the carbon skeleton wherever electrons are available for covalent bonding. Hydrocarbons are the major components of petroleum, which is called a fossil fuel because it consists of the partially decomposed remains of organisms that lived millions of years ago.

Although hydrocarbons are not prevalent in most living organisms, many of a cell's organic molecules have regions consisting of only carbon and hydrogen. For example, the

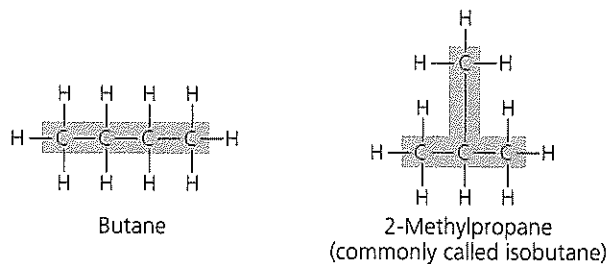
▼ Figure 4.5 Four ways that carbon skeletons can vary.

#### (a) Length



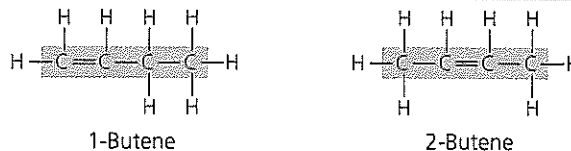
Carbon skeletons vary in length.

#### (b) Branching



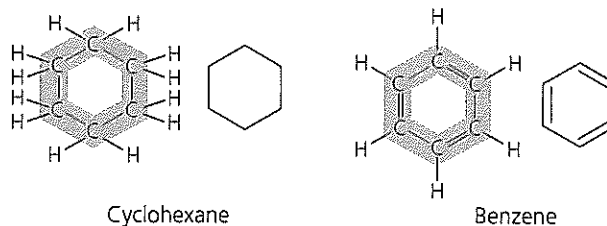
Skeletons may be unbranched or branched.

#### (c) Double bond position



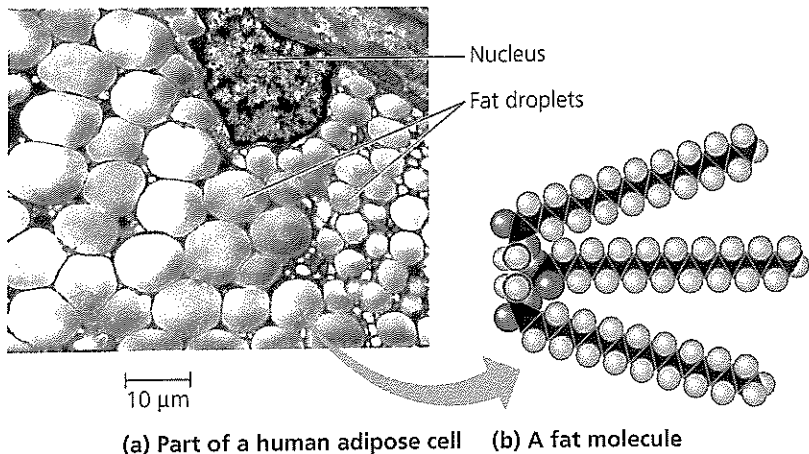
The skeleton may have double bonds, which can vary in location.

#### (d) Presence of rings



Some carbon skeletons are arranged in rings. In the abbreviated structural formula for each compound (at the right), each corner represents a carbon and its attached hydrogens.

molecules known as fats have long hydrocarbon tails attached to a nonhydrocarbon component (Figure 4.6). Neither petroleum nor fat dissolves in water; both are hydrophobic compounds because the great majority of their bonds are relatively nonpolar carbon-to-hydrogen linkages. Another characteristic of hydrocarbons is that they can undergo reactions that release a relatively large amount of energy. The gasoline that fuels a car consists of hydrocarbons, and the hydrocarbon tails of fats serve as stored fuel for plant embryos (seeds) and animals.



**▲ Figure 4.6** The role of hydrocarbons in fats. (a) Mammalian adipose cells stockpile fat molecules as a fuel reserve. This colorized micrograph shows part of a human adipose cell with many fat droplets, each containing a large number of fat molecules. (b) A fat molecule consists of a small, nonhydrocarbon component joined to three hydrocarbon tails that account for the hydrophobic behavior of fats. The tails can be broken down to provide energy. (Black = carbon; gray = hydrogen; red = oxygen.)

**MAKE CONNECTIONS** How do the tails account for the hydrophobic nature of fats? (See Concept 3.2.)

## Isomers

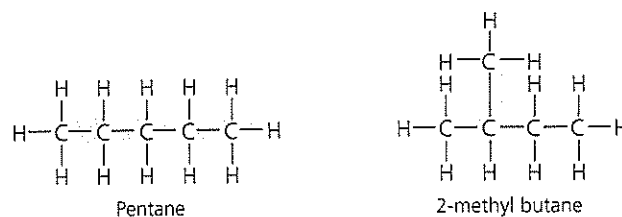
Variation in the architecture of organic molecules can be seen in **isomers**, compounds that have the same numbers of atoms of the same elements but different structures and hence different properties. We will examine three types of isomers: structural isomers, *cis-trans* isomers, and enantiomers.

**Structural isomers** differ in the covalent arrangements of their atoms. Compare, for example, the two five-carbon compounds in **Figure 4.7a**. Both have the molecular formula  $C_5H_{12}$ , but they differ in the covalent arrangement of their carbon skeletons. The skeleton is straight in one compound but branched in the other. The number of possible isomers increases tremendously as carbon skeletons increase in size. There are only three forms of  $C_5H_{12}$  (two of which are shown in Figure 4.7a), but there are 18 variations of  $C_8H_{18}$  and 366,319 possible structural isomers of  $C_{20}H_{42}$ . Structural isomers may also differ in the location of double bonds.

In *cis-trans* isomers (formerly called *geometric isomers*), carbons have covalent bonds to the same atoms, but these atoms differ in their spatial arrangements due to the inflexibility of double bonds. Single bonds allow the atoms they join to rotate freely about the bond axis without changing the compound. In contrast, double bonds do not permit such rotation. If a double bond joins two carbon atoms, and each C also has two different atoms (or groups of atoms) attached to it, then two distinct *cis-trans* isomers are possible. Consider a simple molecule with two double-bonded carbons, each of which has an H and an X attached to it (**Figure 4.7b**). The arrangement with both Xs on the same side of the double bond is called a *cis isomer*, and that with the Xs on opposite sides

**▼ Figure 4.7** Three types of isomers, compounds with the same molecular formula but different structures.

### (a) Structural isomers



Structural isomers differ in covalent partners, as shown in this example of two isomers of  $C_5H_{12}$ .

### (b) *Cis-trans* isomers

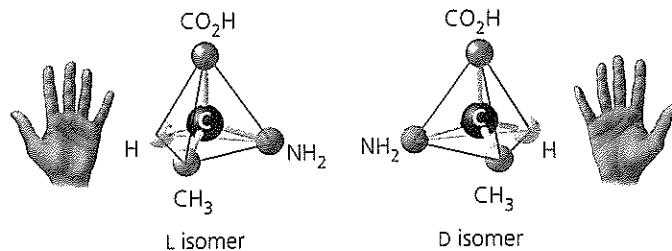


*cis* isomer: The two Xs are on the same side.

*trans* isomer: The two Xs are on opposite sides.

*Cis-trans* isomers differ in arrangement about a double bond. In these diagrams, X represents an atom or group of atoms attached to a double-bonded carbon.

### (c) Enantiomers







Enantiomers differ in spatial arrangement around an asymmetric carbon, resulting in molecules that are mirror images, like left and right hands. The two isomers here are designated the L and D isomers from the Latin for "left" and "right" (*levo* and *dextro*). Enantiomers cannot be superimposed on each other.

**DRAW IT** There are three structural isomers of  $C_5H_{12}$ ; draw the one not shown in (a).

called a *trans isomer*. The subtle difference in shape between such isomers can dramatically affect the biological activities of organic molecules. For example, the biochemistry of vision involves a light-induced change of retinal, a chemical compound in the eye, from the *cis* isomer to the *trans* isomer (see Figure 50.17). Another example involves *trans* fats, which are discussed in Chapter 5.

**Enantiomers** are isomers that are mirror images of each other and that differ in shape due to the presence of an *asymmetric carbon*, one that is attached to four different atoms or groups of atoms. (See the middle carbon in

Drug	Effects	Effective Enantiomer	Ineffective Enantiomer
Ibuprofen	Reduces inflammation and pain	 S-Ibuprofen	 R-Ibuprofen
Albuterol	Relaxes bronchial (airway) muscles, improving airflow in asthma patients	 R-Albuterol	 S-Albuterol

▲ **Figure 4.8** The pharmacological importance of enantiomers. Ibuprofen and albuterol are drugs whose enantiomers have different effects. (*S* and *R* are used here to distinguish between enantiomers.) Ibuprofen is commonly sold as a mixture of the two enantiomers; the *S* enantiomer is 100 times more effective than the *R* form. Albuterol is synthesized and sold only as the *R* form of the drug; the *S* form counteracts the active *R* form.

the ball-and-stick models shown in **Figure 4.7c**.) The four groups can be arranged in space around the asymmetric carbon in two different ways that are mirror images. Enantiomers are, in a way, left-handed and right-handed versions of the molecule. Just as your right hand won't fit into a left-handed glove, a "right-handed" molecule won't fit into the same space as the "left-handed" version. Usually, only one isomer is biologically active because only that form can bind to specific molecules in an organism.

The concept of enantiomers is important in the pharmaceutical industry because the two enantiomers of a drug may not be equally effective, as is the case for both ibuprofen and the asthma medication albuterol (**Figure 4.8**). Methamphetamine also occurs in two enantiomers that have very different effects. One enantiomer is the highly addictive stimulant drug known as "crank," sold illegally in the street drug trade. The other has a much weaker effect and is the active ingredient in an over-the-counter vapor inhaler for treatment of nasal congestion. The differing effects of enantiomers in the body demonstrate that organisms are sensitive to even the most subtle variations in molecular architecture. Once again, we see that molecules have emergent properties that depend on the specific arrangement of their atoms.

#### CONCEPT CHECK 4.2

- DRAW IT** (a) Draw a structural formula for  $C_2H_4$ . (b) Draw the *trans* isomer of  $C_2H_2Cl_2$ .
- Which molecules in Figure 4.5 are isomers? For each pair, identify the type of isomer.
- How are gasoline and fat chemically similar?
- Can propane ( $C_3H_8$ ) form isomers? Explain.

For suggested answers, see Appendix A.

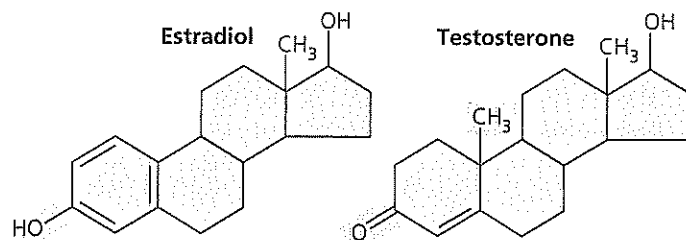
### CONCEPT 4.3

## A few chemical groups are key to molecular function

The properties of an organic molecule depend not only on the arrangement of its carbon skeleton but also on the chemical groups attached to that skeleton. We can think of hydrocarbons, the simplest organic molecules, as the underlying framework for more complex organic molecules. A number of chemical groups can replace one or more hydrogens of the hydrocarbon. These groups may participate in chemical reactions or may contribute to function indirectly by their effects on molecular shape; they help give each molecule its unique properties.

## The Chemical Groups Most Important in the Processes of Life

Consider the differences between estradiol (a type of estrogen) and testosterone. These compounds are female and male sex hormones, respectively, in humans and other vertebrates. Both are steroids, organic molecules with a common carbon skeleton in the form of four fused rings. They differ only in the chemical groups attached to the rings (shown here in abbreviated form); the distinctions in molecular architecture are shaded in blue:

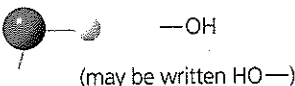
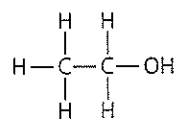
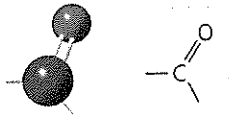
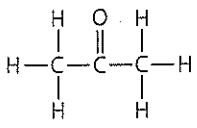
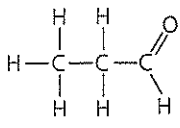
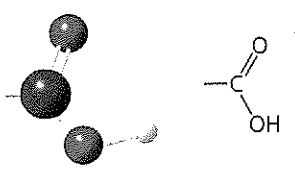
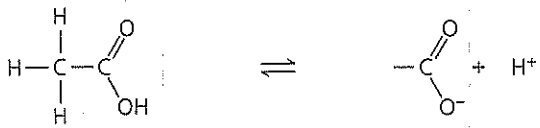
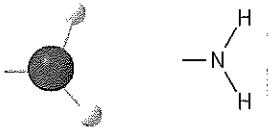
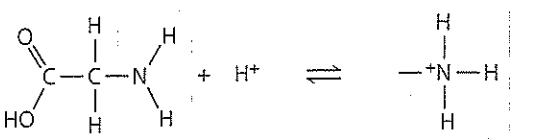
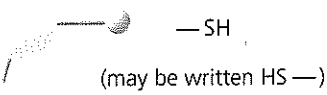
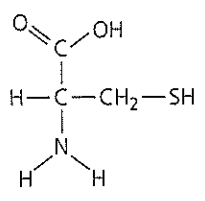
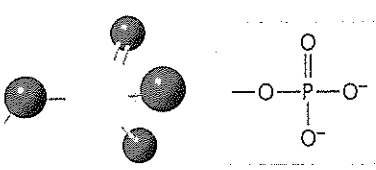
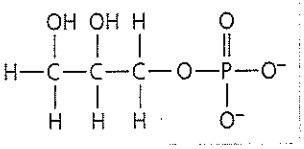
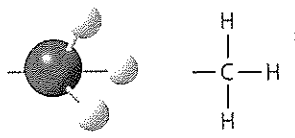
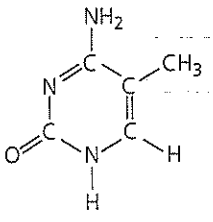


The different actions of these two molecules on many targets throughout the body are the basis of gender, producing the contrasting features of male and female vertebrates. In this case, the chemical groups are important because they affect molecular shape, contributing to function.

In other cases, chemical groups are directly involved in chemical reactions; such groups are known as **functional groups**. Each has certain properties, such as shape and charge, that cause it to participate in chemical reactions in a characteristic way.

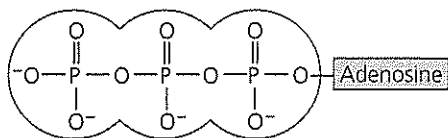
The seven chemical groups most important in biological processes are the hydroxyl, carbonyl, carboxyl, amino, sulfhydryl, phosphate, and methyl groups. The first six groups can be chemically reactive; of these, all except the sulfhydryl group are also hydrophilic and thus increase the solubility of organic compounds in water. The methyl group is not reactive, but instead often serves as a recognizable tag on biological molecules. Study **Figure 4.9** to become familiar with these biologically important chemical groups.

▼ Figure 4.9 Some biologically important chemical groups.

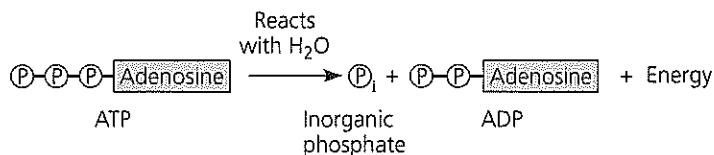
Chemical Group	Group Properties and Compound Name	Examples
<b>Hydroxyl group</b> ( $\text{—OH}$ ) 	<p>Is polar due to electronegative oxygen. Forms hydrogen bonds with water, helping dissolve compounds such as sugars.</p> <p>Compound name: Alcohol (specific name usually ends in <i>-ol</i>)</p>	 <p><b>Ethanol</b>, the alcohol present in alcoholic beverages</p>
<b>Carbonyl group</b> ( $\text{>C=O}$ ) 	<p>Sugars with ketone groups are called ketoses; those with aldehydes are called aldoses.</p> <p>Compound name: Ketone (carbonyl group is within a carbon skeleton) or aldehyde (carbonyl group is at the end of a carbon skeleton)</p>	 <p><b>Acetone</b>, the simplest ketone</p>  <p><b>Propanal</b>, an aldehyde</p>
<b>Carboxyl group</b> ( $\text{—COOH}$ ) 	<p>Acts as an acid (can donate <math>\text{H}^+</math>) because the covalent bond between oxygen and hydrogen is so polar.</p> <p>Compound name: Carboxylic acid, or organic acid</p>	 <p><b>Acetic acid</b>, which gives vinegar its sour taste</p> <p>Ionized form of <math>\text{—COOH}</math> (carboxylate ion), found in cells</p>
<b>Amino group</b> ( $\text{—NH}_2$ ) 	<p>Acts as a base; can pick up an <math>\text{H}^+</math> from the surrounding solution (water, in living organisms).</p> <p>Compound name: Amine</p>	 <p><b>Glycine</b>, an amino acid (note its carboxyl group)</p> <p>Ionized form of <math>\text{—NH}_2</math>, found in cells</p>
<b>Sulfhydryl group</b> ( $\text{—SH}$ ) 	<p>Two <math>\text{—SH}</math> groups can react, forming a "cross-link" that helps stabilize protein structure. Hair protein cross-links maintain the straightness or curliness of hair; in hair salons, permanent treatments break cross-links, then re-form them while the hair is in the desired shape.</p> <p>Compound name: Thiol</p>	 <p><b>Cysteine</b>, a sulfur-containing amino acid</p>
<b>Phosphate group</b> ( $\text{—OPO}_3^{2-}$ ) 	<p>Contributes negative charge (1<math>^-</math> when positioned inside a chain of phosphates; 2<math>^-</math> when at the end). When attached, confers on a molecule the ability to react with water, releasing energy.</p> <p>Compound name: Organic phosphate</p>	 <p><b>Glycerol phosphate</b>, which takes part in many important chemical reactions in cells</p>
<b>Methyl group</b> ( $\text{—CH}_3$ ) 	<p>Affects the expression of genes when on DNA or on proteins bound to DNA. Affects the shape and function of male and female sex hormones.</p> <p>Compound name: Methylated compound</p>	 <p><b>5-Methyl cytosine</b>, a component of DNA that has been modified by addition of a methyl group</p>

## ATP: An Important Source of Energy for Cellular Processes

The “Phosphate group” row in Figure 4.9 shows a simple example of an organic phosphate molecule. A more complicated organic phosphate, **adenosine triphosphate**, or **ATP**, is worth mentioning here because its function in the cell is so important. ATP consists of an organic molecule called adenosine attached to a string of three phosphate groups:



When three phosphates are present in series, as in ATP, one phosphate may be split off as a result of a reaction with water. This inorganic phosphate ion,  $\text{HOPO}_3^{2-}$ , is often abbreviated  $\text{P}_i$  in this book, and a phosphate group in an organic molecule is often written as  $\text{P}$ . Having lost one phosphate, ATP becomes adenosine *diphosphate*, or **ADP**. Although ATP is sometimes said to store energy, it is more accurate to think of it as storing the potential to react with water. This reaction releases energy that can be used by the cell. You will learn about this in more detail in Chapter 8.



### CONCEPT CHECK 4.3

1. What does the term *amino acid* signify about the structure of such a molecule?
2. What chemical change occurs to ATP when it reacts with water and releases energy?
3. **WHAT IF?** Suppose you had an organic molecule such as cysteine (see Figure 4.9, sulfhydryl group example), and you chemically removed the  $-\text{NH}_2$  group and replaced it with  $-\text{COOH}$ . Draw the structural formula for this molecule and speculate about its chemical properties. Is the central carbon asymmetric before the change? After?

For suggested answers, see Appendix A.

## The Chemical Elements of Life: A Review

Living matter, as you have learned, consists mainly of carbon, oxygen, hydrogen, and nitrogen, with smaller amounts of sulfur and phosphorus. These elements all form strong covalent bonds, an essential characteristic in the architecture of complex organic molecules. Of all these elements, carbon is the virtuoso of the covalent bond. The versatility of carbon makes possible the great diversity of organic molecules, each with particular properties that emerge from the unique arrangement of its carbon skeleton and the chemical groups appended to that skeleton. This variation at the molecular level provides the foundation for the rich biological diversity found on our planet.

# 4 Chapter Review

**AP** Of all the elements in the periodic table, why is carbon central to all forms of life? (**Big Idea 2**)

### SUMMARY OF KEY CONCEPTS

#### CONCEPT 4.1

Organic chemistry is the study of carbon compounds (pp. 57–58)

- Organic compounds, once thought to arise only within living organisms, were finally synthesized in the laboratory.
- Living matter is made mostly of carbon, oxygen, hydrogen, and nitrogen. Biological diversity results from carbon's ability to form a huge number of molecules with particular shapes and properties.

**?** How did Stanley Miller's experiments support the idea that, even at life's origins, physical and chemical laws govern the processes of life?

#### CONCEPT 4.2

Carbon atoms can form diverse molecules by bonding to four other atoms (pp. 58–62)

- Carbon, with a valence of 4, can bond to various other atoms, including O, H, and N. Carbon can also bond to other carbon

atoms, forming the carbon skeletons of organic compounds. These skeletons vary in length and shape and have bonding sites for atoms of other elements.

- **Hydrocarbons** consist of carbon and hydrogen.
- **Isomers** are compounds that have the same molecular formula but different structures and therefore different properties. Three types of isomers are **structural isomers**, **cis-trans isomers**, and **enantiomers**.

**?** Refer back to Figure 4.9. What type of isomers are acetone and propanal? How many asymmetric carbons are present in acetic acid, glycine, and glycerol phosphate? Can these three molecules exist as forms that are enantiomers?

#### CONCEPT 4.3

A few chemical groups are key to molecular function (pp. 62–64)

- Chemical groups attached to the carbon skeletons of organic molecules participate in chemical reactions (**functional groups**) or contribute to function by affecting molecular shape (see Figure 4.9).
- **ATP (adenosine triphosphate)** consists of adenosine attached to three phosphate groups. ATP can react with water, forming

