

## 17

TEMPERATURE  
AND HEAT

## LEARNING GOALS

**By studying this chapter, you will learn:**

- The meaning of thermal equilibrium, and what thermometers really measure.
- How different types of thermometers function.
- The physics behind the absolute, or Kelvin, temperature scale.
- How the dimensions of an object change as a result of a temperature change.
- The meaning of heat, and how it differs from temperature.
- How to do calculations that involve heat flow, temperature changes, and changes of phase.
- How heat is transferred by conduction, convection, and radiation.

**?** At a steelworks, molten iron is heated to  $1500^{\circ}$  Celsius to remove impurities. Is it accurate to say that the molten iron contains heat?



**W**hether it's a sweltering summer day or a frozen midwinter night, your body needs to be kept at a nearly constant temperature. It has effective temperature-control mechanisms, but sometimes it needs help. On a hot day you wear less clothing to improve heat transfer from your body to the air and for better cooling by evaporation of perspiration. You drink cold beverages and may sit near a fan or in an air-conditioned room. On a cold day you wear more clothes or stay indoors where it's warm. When you're outside, you keep active and drink hot liquids to stay warm. The concepts in this chapter will help you understand the basic physics of keeping warm or cool.

The terms "temperature" and "heat" are often used interchangeably in everyday language. In physics, however, these two terms have very different meanings. In this chapter we'll define temperature in terms of how it's measured and see how temperature changes affect the dimensions of objects. We'll see that heat refers to energy transfer caused by temperature differences and learn how to calculate and control such energy transfers.

Our emphasis in this chapter is on the concepts of temperature and heat as they relate to *macroscopic* objects such as cylinders of gas, ice cubes, and the human body. In Chapter 18 we'll look at these same concepts from a *microscopic* viewpoint in terms of the behavior of individual atoms and molecules. These two chapters lay the groundwork for the subject of **thermodynamics**, the study of energy transformations involving heat, mechanical work, and other aspects of energy and how these transformations relate to the properties of matter. Thermodynamics forms an indispensable part of the foundation of physics, chemistry, and the life sciences, and its applications turn up in such places as car engines, refrigerators, biochemical processes, and the structure of stars. We'll explore the key ideas of thermodynamics in Chapters 19 and 20.

## 17.1 Temperature and Thermal Equilibrium

The concept of **temperature** is rooted in qualitative ideas of “hot” and “cold” based on our sense of touch. A body that feels hot usually has a higher temperature than a similar body that feels cold. That’s pretty vague, and the senses can be deceived. But many properties of matter that we can *measure* depend on temperature. The length of a metal rod, steam pressure in a boiler, the ability of a wire to conduct an electric current, and the color of a very hot glowing object—all these depend on temperature.

Temperature is also related to the kinetic energies of the molecules of a material. In general this relationship is fairly complex, so it’s not a good place to start in *defining* temperature. In Chapter 18 we will look at the relationship between temperature and the energy of molecular motion for an ideal gas. It is important to understand, however, that temperature and heat can be defined independently of any detailed molecular picture. In this section we’ll develop a *macroscopic* definition of temperature.

To use temperature as a measure of hotness or coldness, we need to construct a temperature scale. To do this, we can use any measurable property of a system that varies with its “hotness” or “coldness.” Figure 17.1a shows a familiar system that is used to measure temperature. When the system becomes hotter, the colored liquid (usually mercury or ethanol) expands and rises in the tube, and the value of  $L$  increases. Another simple system is a quantity of gas in a constant-volume container (Fig. 17.1b). The pressure  $p$ , measured by the gauge, increases or decreases as the gas becomes hotter or colder. A third example is the electrical resistance  $R$  of a conducting wire, which also varies when the wire becomes hotter or colder. Each of these properties gives us a number ( $L$ ,  $p$ , or  $R$ ) that varies with hotness and coldness, so each property can be used to make a **thermometer**.

To measure the temperature of a body, you place the thermometer in contact with the body. If you want to know the temperature of a cup of hot coffee, you stick the thermometer in the coffee; as the two interact, the thermometer becomes hotter and the coffee cools off a little. After the thermometer settles down to a steady value, you read the temperature. The system has reached an *equilibrium* condition, in which the interaction between the thermometer and the coffee causes no further change in the system. We call this a state of **thermal equilibrium**.

If two systems are separated by an insulating material or **insulator** such as wood, plastic foam, or fiberglass, they influence each other more slowly. Camping coolers are made with insulating materials to delay the ice and cold food inside from warming up and attaining thermal equilibrium with the hot summer air outside. An *ideal insulator* is a material that permits no interaction at all between the two systems. It prevents the systems from attaining thermal equilibrium if they aren’t in thermal equilibrium at the start. An ideal insulator is just that, an idealization; real insulators, like those in camping coolers, aren’t ideal, so the contents of the cooler will warm up eventually.

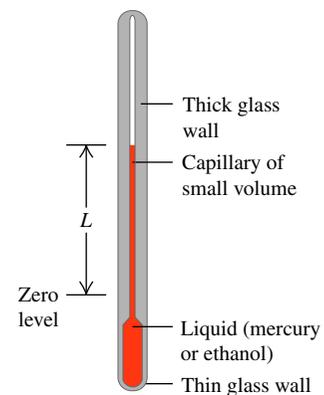
### The Zeroth Law of Thermodynamics

We can discover an important property of thermal equilibrium by considering three systems,  $A$ ,  $B$ , and  $C$ , that initially are not in thermal equilibrium (Fig. 17.2). We surround them with an ideal insulating box so that they cannot interact with anything except each other. We separate systems  $A$  and  $B$  with an ideal insulating wall (the green slab in Fig. 17.2a), but we let system  $C$  interact with both systems  $A$  and  $B$ . This interaction is shown in the figure by a yellow slab representing a thermal **conductor**, a material that *permits* thermal interactions through it. We wait until thermal equilibrium is attained; then  $A$  and  $B$  are each in thermal equilibrium with  $C$ . But are they in thermal equilibrium *with each other*?

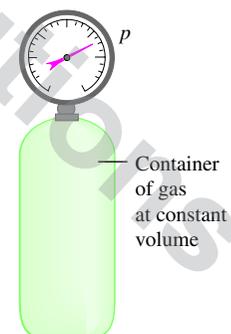
To find out, we separate system  $C$  from systems  $A$  and  $B$  with an ideal insulating wall (Fig. 17.2b), and then we replace the insulating wall between  $A$  and  $B$  with a

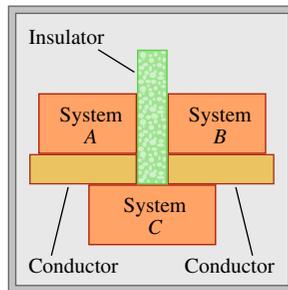
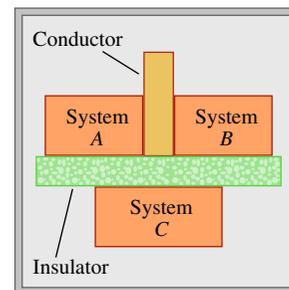
**17.1** Two devices for measuring temperature.

(a) Changes in temperature cause the liquid’s volume to change.



(b) Changes in temperature cause the pressure of the gas to change.



**17.2** The zeroth law of thermodynamics.**(a)** If systems *A* and *B* are each in thermal equilibrium with system *C* ...**(b)** ... then systems *A* and *B* are in thermal equilibrium with each other.

conducting wall that lets *A* and *B* interact. What happens? Experiment shows that *nothing* happens; there are no additional changes to *A* or *B*. We conclude

**If *C* is initially in thermal equilibrium with both *A* and *B*, then *A* and *B* are also in thermal equilibrium with each other. This result is called the zeroth law of thermodynamics.**

(The importance of this law was recognized only after the first, second, and third laws of thermodynamics had been named. Since it is fundamental to all of them, the name “zeroth” seemed appropriate.)

Now suppose system *C* is a thermometer, such as the tube-and-liquid system of Fig. 17.1a. In Fig. 17.2a the thermometer *C* is in contact with both *A* and *B*. In thermal equilibrium, when the thermometer reading reaches a stable value, the thermometer measures the temperature of both *A* and *B*; hence *A* and *B* both have the *same* temperature. Experiment shows that thermal equilibrium isn’t affected by adding or removing insulators, so the reading of thermometer *C* wouldn’t change if it were in contact only with *A* or only with *B*. We conclude

**Two systems are in thermal equilibrium if and only if they have the same temperature.**

This is what makes a thermometer useful; a thermometer actually measures *its own* temperature, but when a thermometer is in thermal equilibrium with another body, the temperatures must be equal. When the temperatures of two systems are different, they *cannot* be in thermal equilibrium.

**Test Your Understanding of Section 17.1** You put a thermometer in a pot of hot water and record the reading. What temperature have you recorded? (i) the temperature of the water; (ii) the temperature of the thermometer; (iii) an equal average of the temperatures of the water and thermometer; (iv) a weighted average of the temperatures of the water and thermometer, with more emphasis on the temperature of the water; (v) a weighted average of the water and thermometer, with more emphasis on the temperature of the thermometer. **MP**

## 17.2 Thermometers and Temperature Scales

To make the liquid-in-tube device shown in Fig. 17.1a into a useful thermometer, we need to mark a scale on the tube wall with numbers on it. These numbers are arbitrary, and historically many different schemes have been used. Suppose we label the thermometer’s liquid level at the freezing temperature of pure water “zero” and the level at the boiling temperature “100,” and divide the distance between these two points into 100 equal intervals called *degrees*. The result is the **Celsius temperature scale** (formerly called the *centigrade* scale in English-

speaking countries). The Celsius temperature for a state colder than freezing water is a negative number. The Celsius scale is used, both in everyday life and in science and industry, almost everywhere in the world.

Another common type of thermometer uses a *bimetallic strip*, made by bonding strips of two different metals together (Fig. 17.3a). When the temperature of the composite strip increases, one metal expands more than the other and the strip bends (Fig. 17.3b). This strip is usually formed into a spiral, with the outer end anchored to the thermometer case and the inner end attached to a pointer (Fig. 17.3c). The pointer rotates in response to temperature changes.

In a *resistance thermometer* the changing electrical resistance of a coil of fine wire, a carbon cylinder, or a germanium crystal is measured. Because resistance can be measured very precisely, resistance thermometers are usually more precise than most other types.

Some thermometers work by detecting the amount of infrared radiation emitted by an object. (We'll see in Section 17.7 that *all* objects emit electromagnetic radiation, including infrared, as a consequence of their temperature.) A modern example is a *temporal artery thermometer* (Fig. 17.4). A nurse runs this over a patient's forehead in the vicinity of the temporal artery, and an infrared sensor in the thermometer measures the radiation from the skin. Tests show that this device gives more accurate values of body temperature than do oral or ear thermometers.

In the **Fahrenheit temperature scale**, still used in everyday life in the United States, the freezing temperature of water is 32°F (thirty-two degrees Fahrenheit) and the boiling temperature is 212°F, both at standard atmospheric pressure. There are 180 degrees between freezing and boiling, compared to 100 on the Celsius scale, so one Fahrenheit degree represents only  $\frac{100}{180}$ , or  $\frac{5}{9}$ , as great a temperature change as one Celsius degree.

To convert temperatures from Celsius to Fahrenheit, note that a Celsius temperature  $T_C$  is the number of Celsius degrees above freezing; the number of Fahrenheit degrees above freezing is  $\frac{9}{5}$  of this. But freezing on the Fahrenheit scale is at 32°F, so to obtain the actual Fahrenheit temperature  $T_F$ , multiply the Celsius value by  $\frac{9}{5}$  and then add 32°. Symbolically,

$$T_F = \frac{9}{5}T_C + 32^\circ \quad (17.1)$$

To convert Fahrenheit to Celsius, solve this equation for  $T_C$ :

$$T_C = \frac{5}{9}(T_F - 32^\circ) \quad (17.2)$$

In words, subtract 32° to get the number of Fahrenheit degrees above freezing, and then multiply by  $\frac{5}{9}$  to obtain the number of Celsius degrees above freezing—that is, the Celsius temperature.

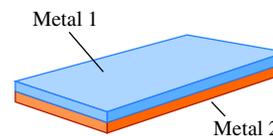
We don't recommend memorizing Eqs. (17.1) and (17.2). Instead, try to understand the reasoning that led to them so that you can derive them on the spot when you need them, checking your reasoning with the relationship  $100^\circ\text{C} = 212^\circ\text{F}$ .

It is useful to distinguish between an actual temperature and a temperature *interval* (a difference or change in temperature). An actual temperature of 20° is stated as 20°C (twenty degrees Celsius), and a temperature *interval* of 10° is 10 C° (ten Celsius degrees). A beaker of water heated from 20°C to 30°C undergoes a temperature change of 10 C°.

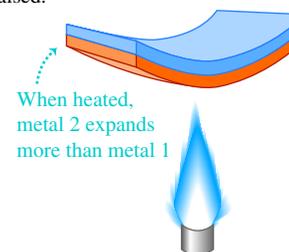
**Test Your Understanding of Section 17.2** Which of the following types of thermometers have to be in thermal equilibrium with the object being measured in order to give accurate readings? (i) a bimetallic strip; (ii) a resistance thermometer; (iii) a temporal artery thermometer; (iv) both (i) and (ii); (v) all of (i), (ii), and (iii).

**17.3** Use of a bimetallic strip as a thermometer.

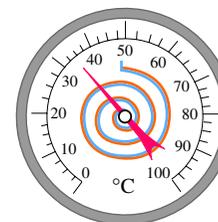
(a) A bimetallic strip



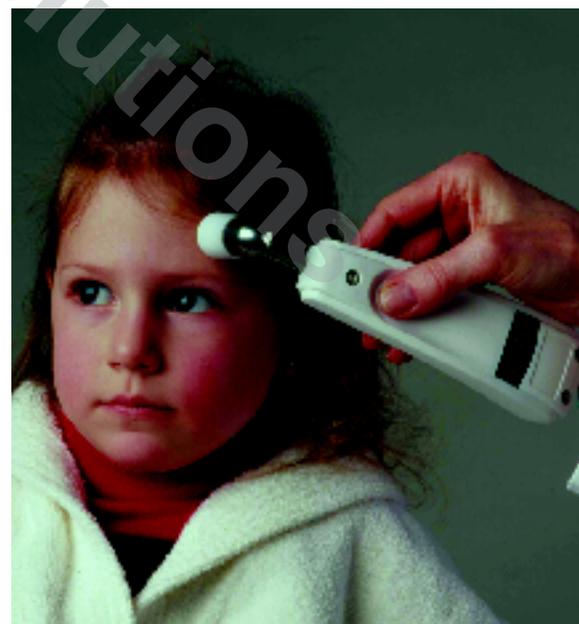
(b) The strip bends when its temperature is raised.



(c) A bimetallic strip used in a thermometer



**17.4** A temporal artery thermometer measures infrared radiation from the skin that overlies one of the important arteries in the head. Although the thermometer cover touches the skin, the infrared detector inside the cover does not.



### 17.3 Gas Thermometers and the Kelvin Scale

When we calibrate two thermometers, such as a liquid-in-tube system and a resistance thermometer, so that they agree at  $0^{\circ}\text{C}$  and  $100^{\circ}\text{C}$ , they may not agree exactly at intermediate temperatures. Any temperature scale defined in this way always depends somewhat on the specific properties of the material used. Ideally, we would like to define a temperature scale that *doesn't* depend on the properties of a particular material. To establish a truly material-independent scale, we first need to develop some principles of thermodynamics. We'll return to this fundamental problem in Chapter 20. Here we'll discuss a thermometer that comes close to the ideal, the *gas thermometer*.

The principle of a gas thermometer is that the pressure of a gas at constant volume increases with temperature. A quantity of gas is placed in a constant-volume container (Fig. 17.5a), and its pressure is measured by one of the devices described in Section 14.2. To calibrate a constant-volume gas thermometer, we measure the pressure at two temperatures, say  $0^{\circ}\text{C}$  and  $100^{\circ}\text{C}$ , plot these points on a graph, and draw a straight line between them. Then we can read from the graph the temperature corresponding to any other pressure. Figure 17.5b shows the results of three such experiments, each using a different type and quantity of gas.

By extrapolating this graph, we see that there is a hypothetical temperature,  $-273.15^{\circ}\text{C}$ , at which the absolute pressure of the gas would become zero. We might expect that this temperature would be different for different gases, but it turns out to be the *same* for many different gases (at least in the limit of very low gas density). We can't actually observe this zero-pressure condition. Gases liquefy and solidify at very low temperatures, and the proportionality of pressure to temperature no longer holds.

We use this extrapolated zero-pressure temperature as the basis for a temperature scale with its zero at this temperature. This is the **Kelvin temperature scale**, named for the British physicist Lord Kelvin (1824–1907). The units are the same size as those on the Celsius scale, but the zero is shifted so that  $0\text{ K} = -273.15^{\circ}\text{C}$  and  $273.15\text{ K} = 0^{\circ}\text{C}$ ; that is,

$$T_{\text{K}} = T_{\text{C}} + 273.15 \quad (17.3)$$

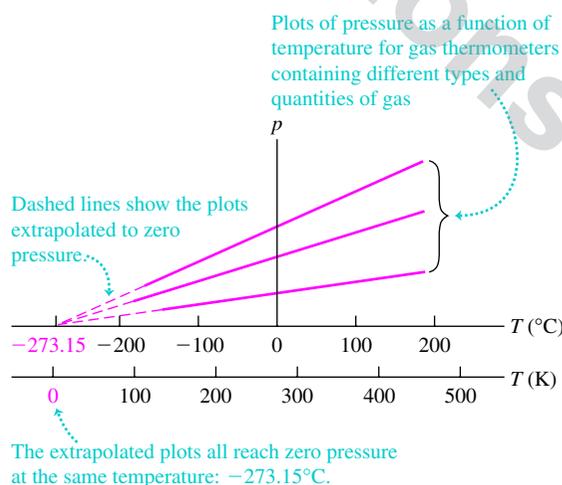
This scale is shown in Fig. 17.5b. A common room temperature,  $20^{\circ}\text{C}$  ( $= 68^{\circ}\text{F}$ ), is  $20 + 273.15$ , or about  $293\text{ K}$ .

**17.5** (a) Using a constant-volume gas thermometer to measure temperature.  
(b) The greater the amount of gas in the thermometer, the higher the graph of pressure  $p$  versus temperature  $T$ .

(a) A constant-volume gas thermometer

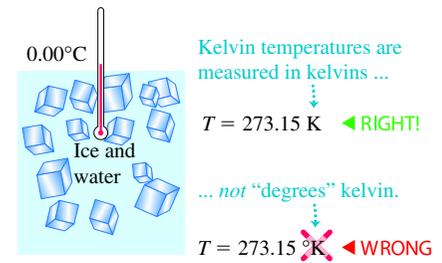


(b) Graphs of pressure versus temperature at constant volume for three different types and quantities of gas



**CAUTION Never say “degrees kelvin”** In SI nomenclature, “degree” is not used with the Kelvin scale; the temperature mentioned above is read “293 kelvins,” not “degrees kelvin” (Fig. 17.6). We capitalize Kelvin when it refers to the temperature scale; however, the *unit* of temperature is the *kelvin*, which is not capitalized (but is nonetheless abbreviated as a capital K). ■

**17.6** Correct and incorrect uses of the Kelvin scale.



### Example 17.1 Body temperature

You place a small piece of melting ice in your mouth. Eventually, the water all converts from ice at  $T_1 = 32.00^\circ\text{F}$  to body temperature,  $T_2 = 98.60^\circ\text{F}$ . Express these temperatures as  $^\circ\text{C}$  and K, and find  $\Delta T = T_2 - T_1$  in both cases.

#### SOLUTION

**IDENTIFY:** Our target variables are temperatures  $T_1$  and  $T_2$  expressed in Celsius degrees and in kelvins, as well as the difference between these two temperatures.

**SET UP:** We convert Fahrenheit to Celsius temperatures using Eq. (17.2), and Celsius to Kelvin temperatures using Eq. (17.3).

**EXECUTE:** First we find the Celsius temperatures. We know that  $T_1 = 32.00^\circ\text{F} = 0.00^\circ\text{C}$ , and  $98.60^\circ\text{F}$  is  $98.60 - 32.00 =$

$66.60^\circ\text{F}$  above freezing; we multiply this by  $(5^\circ\text{C}/9^\circ\text{F})$  to find  $37.00^\circ\text{C}$  above freezing, or  $T_2 = 37.00^\circ\text{C}$ .

To get the Kelvin temperatures, we just add 273.15 to each Celsius temperature:  $T_1 = 273.15 \text{ K}$  and  $T_2 = 310.15 \text{ K}$ . “Normal” body temperature is  $37.0^\circ\text{C}$ , but if your doctor says that your temperature is 310 K, don’t be alarmed.

The temperature *difference*  $\Delta T = T_2 - T_1$  is  $37.00^\circ\text{C} = 37.00 \text{ K}$ .

**EVALUATE:** The Celsius and Kelvin scales have different zero points but the same size degrees. Therefore any temperature difference is the *same* on the Celsius and Kelvin scales but not the same on the Fahrenheit scale.

## The Kelvin Scale and Absolute Temperature

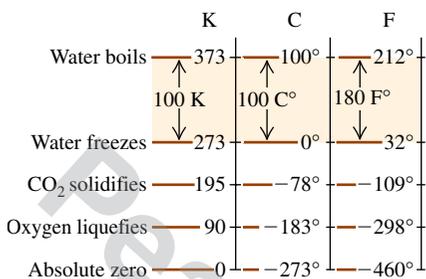
The Celsius scale has two fixed points, the normal freezing and boiling temperatures of water. But we can define the Kelvin scale using a gas thermometer with only a single reference temperature. We define the ratio of any two temperatures  $T_1$  and  $T_2$  on the Kelvin scale as the ratio of the corresponding gas-thermometer pressures  $p_1$  and  $p_2$ :

$$\frac{T_2}{T_1} = \frac{p_2}{p_1} \quad (\text{constant-volume gas thermometer, } T \text{ in kelvins}) \quad (17.4)$$

The pressure  $p$  is directly proportional to the Kelvin temperature, as shown in Fig. 17.5b. To complete the definition of  $T$ , we need only specify the Kelvin temperature of a single specific state. For reasons of precision and reproducibility, the state chosen is the *triple point* of water. This is the unique combination of temperature and pressure at which solid water (ice), liquid water, and water vapor can all coexist. It occurs at a temperature of  $0.01^\circ\text{C}$  and a water-vapor pressure of 610 Pa (about 0.006 atm). (This is the pressure of the *water*; it has nothing to do directly with the gas pressure in the *thermometer*.) The triple-point temperature  $T_{\text{triple}}$  of water is *defined* to have the value  $T_{\text{triple}} = 273.16 \text{ K}$ , corresponding to  $0.01^\circ\text{C}$ . From Eq. (17.4), if  $p_{\text{triple}}$  is the pressure in a gas thermometer at temperature  $T_{\text{triple}}$  and  $p$  is the pressure at some other temperature  $T$ , then  $T$  is given on the Kelvin scale by

$$T = T_{\text{triple}} \frac{p}{p_{\text{triple}}} = (273.16 \text{ K}) \frac{p}{p_{\text{triple}}} \quad (17.5)$$

**17.7** Relationships among Kelvin (K), Celsius (C), and Fahrenheit (F) temperature scales. Temperatures have been rounded off to the nearest degree.



Low-pressure gas thermometers using various gases are found to agree very closely, but they are large, bulky, and very slow to come to thermal equilibrium. They are used principally to establish high-precision standards and to calibrate other thermometers.

Figure 17.7 shows the relationships among the three temperature scales we have discussed. The Kelvin scale is called an **absolute temperature scale**, and its zero point ( $T = 0 \text{ K} = -273.15^\circ\text{C}$ , the temperature at which  $p = 0$  in Eq. (17.5)) is called **absolute zero**. At absolute zero a system of molecules (such as a quantity of a gas, a liquid, or a solid) has its *minimum* possible total energy (kinetic plus potential); because of quantum effects, however, it is *not* correct to say that all molecular motion ceases at absolute zero. To define more completely what we mean by absolute zero, we need to use the thermodynamic principles developed in the next several chapters. We will return to this concept in Chapter 20.

**Test Your Understanding of Section 17.3** Rank the following temperatures from highest to lowest: (i)  $0.00^\circ\text{C}$ ; (ii)  $0.00^\circ\text{F}$ ; (iii)  $260.00 \text{ K}$ ; (iv)  $77.00 \text{ K}$ ; (v)  $-180.00^\circ\text{C}$ .

## 17.4 Thermal Expansion

Most materials expand when their temperatures increase. Rising temperatures make the liquid expand in a liquid-in-tube thermometer (Fig. 17.1a) and bend bimetallic strips (Fig. 17.3b). The decks of bridges need special joints and supports to allow for expansion. A completely filled and tightly capped bottle of water cracks when it is heated, but you can loosen a metal jar lid by running hot water over it. These are all examples of *thermal expansion*.

### Linear Expansion

Suppose a rod of material has a length  $L_0$  at some initial temperature  $T_0$ . When the temperature changes by  $\Delta T$ , the length changes by  $\Delta L$ . Experiments show that if  $\Delta T$  is not too large (say, less than  $100 \text{ C}^\circ$  or so),  $\Delta L$  is *directly proportional* to  $\Delta T$  (Fig. 17.8a). If two rods made of the same material have the same temperature change, but one is twice as long as the other, then the *change* in its length is also twice as great. Therefore  $\Delta L$  must also be proportional to  $L_0$  (Fig. 17.8b). Introducing a proportionality constant  $\alpha$  (which is different for different materials), we may express these relationships in an equation:

$$\Delta L = \alpha L_0 \Delta T \quad (\text{linear thermal expansion}) \quad (17.6)$$

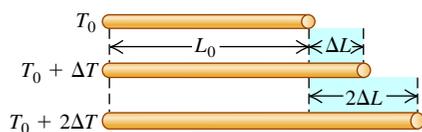
If a body has length  $L_0$  at temperature  $T_0$ , then its length  $L$  at a temperature  $T = T_0 + \Delta T$  is

$$L = L_0 + \Delta L = L_0 + \alpha L_0 \Delta T = L_0(1 + \alpha \Delta T) \quad (17.7)$$

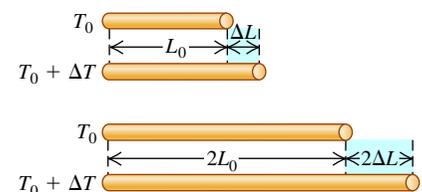
The constant  $\alpha$ , which describes the thermal expansion properties of a particular material, is called the **coefficient of linear expansion**. The units of  $\alpha$  are  $\text{K}^{-1}$  or  $(\text{C}^\circ)^{-1}$ . (Remember that a temperature *interval* is the same in the Kelvin and Celsius scales.) For many materials, every linear dimension changes according to Eq. (17.6) or (17.7). Thus  $L$  could be the thickness of a rod, the side length of a square sheet, or the diameter of a hole. Some materials, such as wood or single crystals, expand differently in different directions. We won't consider this complication.

**17.8** How the length of a rod changes with a change in temperature. (Length changes are exaggerated for clarity.)

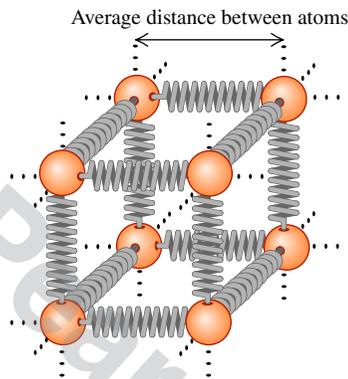
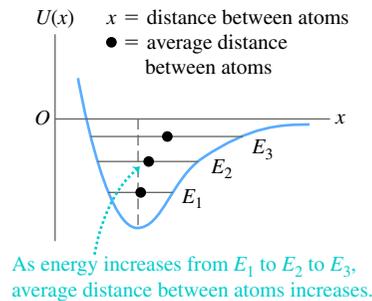
(a) For moderate temperature changes,  $\Delta L$  is directly proportional to  $\Delta T$ .



(b)  $\Delta L$  is also directly proportional to  $L_0$ .



(a) A model of the forces between neighboring atoms in a solid

(b) A graph of the “spring” potential energy  $U(x)$ 

We can understand thermal expansion qualitatively on a molecular basis. Picture the interatomic forces in a solid as springs, as in Fig. 17.9. (We explored the analogy between spring forces and interatomic forces in Section 13.4.) Each atom vibrates about its equilibrium position. When the temperature increases, the energy and amplitude of the vibration also increase. The interatomic spring forces are not symmetrical about the equilibrium position; they usually behave like a spring that is easier to stretch than to compress. As a result, when the amplitude of vibration increases, the *average* distance between atoms also increases. As the atoms get farther apart, every dimension increases.

**CAUTION Heating an object with a hole** If a solid object has a hole in it, what happens to the size of the hole when the temperature of the object increases? A common misconception is that if the object expands, the hole will shrink because material expands into the hole. But the truth of the matter is that if the object expands, the hole will expand too (Fig. 17.10); as we stated above, *every* linear dimension of an object changes in the same way when the temperature changes. If you’re not convinced, think of the atoms in Fig. 17.9a as outlining a cubical hole. When the object expands, the atoms move apart and the hole increases in size. The only situation in which a “hole” will fill in due to thermal expansion is when two separate objects expand and close the gap between them (Fig. 17.11). ■

The direct proportionality expressed by Eq. (17.6) is not exact; it is *approximately* correct only for sufficiently small temperature changes. For a given material,  $\alpha$  varies somewhat with the initial temperature  $T_0$  and the size of the temperature interval. We’ll ignore this complication here, however. Average values of  $\alpha$  for several materials are listed in Table 17.1 on page 578. Within the precision of these values we don’t need to worry whether  $T_0$  is  $0^\circ\text{C}$  or  $20^\circ\text{C}$  or some other temperature. Note that typical values of  $\alpha$  are very small; even for a temperature change of  $100^\circ\text{C}$ , the fractional length change  $\Delta L/L_0$  is only of the order of  $1/1000$  for the metals in the table.

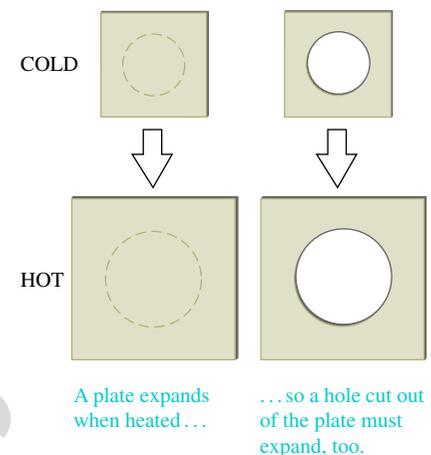
## Volume Expansion

Increasing temperature usually causes increases in *volume* for both solid and liquid materials. Just as with linear expansion, experiments show that if the temperature change  $\Delta T$  is not too great (less than  $100^\circ\text{C}$  or so), the increase in volume  $\Delta V$  is approximately proportional to both the temperature change  $\Delta T$  and the initial volume  $V_0$ :

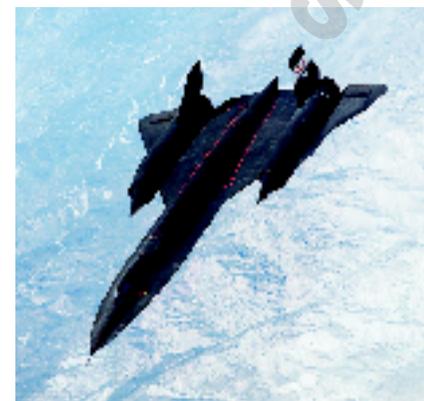
$$\Delta V = \beta V_0 \Delta T \quad (\text{volume thermal expansion}) \quad (17.8)$$

**17.9** (a) We can model atoms in a solid as being held together by “springs” that are easier to stretch than to compress. (b) A graph of the “spring” potential energy  $U(x)$  versus distance  $x$  between neighboring atoms is *not* symmetrical (compare Fig. 13.20b). As the energy increases and the atoms oscillate with greater amplitude, the *average* distance increases.

**17.10** When an object undergoes thermal expansion, any holes in the object expand as well. (The expansion is exaggerated.)



**17.11** When this SR-71 aircraft is sitting on the ground, its wing panels fit together so loosely that fuel leaks out of the wings onto the ground. But once it is in flight at over three times the speed of sound, air friction heats the panels so much that they expand to make a perfect fit. (In-flight refueling makes up for the lost fuel.)



**Table 17.1** Coefficients of Linear Expansion

| Material                  | $\alpha$ [ $\text{K}^{-1}$ or $(\text{C}^\circ)^{-1}$ ] |
|---------------------------|---|
| Aluminum                  | $2.4 \times 10^{-5}$                                    |
| Brass                     | $2.0 \times 10^{-5}$                                    |
| Copper                    | $1.7 \times 10^{-5}$                                    |
| Glass                     | $0.4\text{--}0.9 \times 10^{-5}$                        |
| Invar (nickel-iron alloy) | $0.09 \times 10^{-5}$                                   |
| Quartz (fused)            | $0.04 \times 10^{-5}$                                   |
| Steel                     | $1.2 \times 10^{-5}$                                    |

The constant  $\beta$  characterizes the volume expansion properties of a particular material; it is called the **coefficient of volume expansion**. The units of  $\beta$  are  $\text{K}^{-1}$  or  $(\text{C}^\circ)^{-1}$ . As with linear expansion,  $\beta$  varies somewhat with temperature, and Eq. (17.8) is an approximate relationship that is valid only for small temperature changes. For many substances,  $\beta$  decreases at low temperatures. Several values of  $\beta$  in the neighborhood of room temperature are listed in Table 17.2. Note that the values for liquids are generally much larger than those for solids.

For solid materials there is a simple relationship between the volume expansion coefficient  $\beta$  and the linear expansion coefficient  $\alpha$ . To derive this relationship, we consider a cube of material with side length  $L$  and volume  $V = L^3$ . At the initial temperature the values are  $L_0$  and  $V_0$ . When the temperature increases by  $dT$ , the side length increases by  $dL$  and the volume increases by an amount  $dV$  given by

$$dV = \frac{dV}{dL} dL = 3L^2 dL$$

Now we replace  $L$  and  $V$  by the initial values  $L_0$  and  $V_0$ . From Eq. (17.6),  $dL$  is

$$dL = \alpha L_0 dT$$

Since  $V_0 = L_0^3$ , this means that  $dV$  can also be expressed as

$$dV = 3L_0^2 \alpha L_0 dT = 3\alpha V_0 dT$$

This is consistent with the infinitesimal form of Eq. (17.8),  $dV = \beta V_0 dT$ , only if

$$\beta = 3\alpha \quad (17.9)$$

You should check this relationship for some of the materials listed in Tables 17.1 and 17.2.

**Table 17.2** Coefficients of Volume Expansion

| Solids         | $\beta$ [ $\text{K}^{-1}$ or $(\text{C}^\circ)^{-1}$ ] | Liquids          | $\beta$ [ $\text{K}^{-1}$ or $(\text{C}^\circ)^{-1}$ ] |
|----------------|--|------------------|--|
| Aluminum       | $7.2 \times 10^{-5}$                                   | Ethanol          | $75 \times 10^{-5}$                                    |
| Brass          | $6.0 \times 10^{-5}$                                   | Carbon disulfide | $115 \times 10^{-5}$                                   |
| Copper         | $5.1 \times 10^{-5}$                                   | Glycerin         | $49 \times 10^{-5}$                                    |
| Glass          | $1.2\text{--}2.7 \times 10^{-5}$                       | Mercury          | $18 \times 10^{-5}$                                    |
| Invar          | $0.27 \times 10^{-5}$                                  |                  |  |
| Quartz (fused) | $0.12 \times 10^{-5}$                                  |                  |  |
| Steel          | $3.6 \times 10^{-5}$                                   |                  |  |

### Problem-Solving Strategy 17.1 Thermal Expansion

**IDENTIFY** the relevant concepts: Decide whether the problem involves changes in length (linear thermal expansion) or in volume (volume thermal expansion).

**SET UP** the problem using the following steps:

1. Choose Eq. (17.6) for linear expansion and Eq. (17.8) for volume expansion.
2. Identify which quantities in Eq. (17.6) or (17.8) are known and which are the unknown target variables.

**EXECUTE** the solution as follows:

1. Solve for the target variables. Often you will be given two temperatures and asked to compute  $\Delta T$ . Or you may be given an

initial temperature  $T_0$  and asked to find a final temperature corresponding to a given length or volume change. In this case, plan to find  $\Delta T$  first; then the final temperature is  $T_0 + \Delta T$ .

2. Unit consistency is crucial, as always.  $L_0$  and  $\Delta L$  (or  $V_0$  and  $\Delta V$ ) must have the same units, and if you use a value of  $\alpha$  or  $\beta$  in  $\text{K}^{-1}$  or  $(\text{C}^\circ)^{-1}$ , then  $\Delta T$  must be in kelvins or Celsius degrees ( $\text{C}^\circ$ ). But you can use K and  $\text{C}^\circ$  interchangeably.

**EVALUATE** your answer: Check whether your results make sense. Remember that the sizes of holes in a material expand with temperature just the same way as any other linear dimension, and the volume of a hole (such as the volume of a container) expands the same way as the corresponding solid shape.



**Example 17.2 Length change due to temperature change I**

A surveyor uses a steel measuring tape that is exactly 50.000 m long at a temperature of 20°C. What is its length on a hot summer day when the temperature is 35°C?

**SOLUTION**

**IDENTIFY:** This problem concerns linear expansion. We are given the initial length and initial temperature of the tape, and our target variable is the tape's length at the final temperature.

**SET UP:** We use Eq. (17.6) to find the change  $\Delta L$  in the tape's length. We are given  $L = 50.000$  m,  $T_0 = 20^\circ\text{C}$ , and  $T = 35^\circ\text{C}$ , and the value of  $\alpha$  is found from Table 17.1. The target variable is the new length  $L = L_0 + \Delta L$ .

**EXECUTE:** The temperature change is  $\Delta T = T - T_0 = 15^\circ\text{C}$ , so from Eq. (17.6) the change in length  $\Delta L$  and the final length  $L = L_0 + \Delta L$  are

$$\begin{aligned}\Delta L &= \alpha L_0 \Delta T = (1.2 \times 10^{-5} \text{ K}^{-1})(50 \text{ m})(15 \text{ K}) \\ &= 9.0 \times 10^{-3} \text{ m} = 9.0 \text{ mm} \\ L &= L_0 + \Delta L = 50.000 \text{ m} + 0.009 \text{ m} = 50.009 \text{ m}\end{aligned}$$

Thus the length at 35°C is 50.009 m.

**EVALUATE:** Note that  $L_0$  is given to five significant figures but that we need only two of them to compute  $\Delta L$ . Note also that  $\Delta L$  is proportional to the initial length  $L_0$ : A 5.0-m tape would expand by 0.90 mm, and a 0.50-m (50-cm) tape would expand by a mere 0.090 mm.

This example shows that metals expand very little under moderate temperature changes. Even a metal baking pan in a 200°C (392°F) oven is only slightly larger than it is at room temperature.

**Example 17.3 Length change due to temperature change II**

In Example 17.2 the surveyor uses the measuring tape to measure a distance when the temperature is 35°C; the value that she reads off the tape is 35.794 m. What is the actual distance? Assume that the tape is calibrated for use at 20°C.

**SOLUTION**

**IDENTIFY:** As we saw in Example 17.2, at 35°C the tape has expanded slightly. The distance between two successive meter marks is slightly more than 1 meter, so the scale underestimates the actual distance.

**SET UP:** The actual distance (our target variable) is *larger* than the distance read off the tape by a factor equal to the ratio of the tape's length  $L$  at 35°C to its length  $L_0$  at 20°C.

**EXECUTE:** The ratio  $L/L_0$  is  $(50.009 \text{ m})/(50.000 \text{ m})$ , so the true distance is

$$\frac{50.009 \text{ m}}{50.000 \text{ m}}(35.794 \text{ m}) = 35.800 \text{ m}$$

**EVALUATE:** Although the difference of 0.008 m = 8 mm between the scale reading and the actual distance seems small, it can be important in precision work.

**Example 17.4 Volume change due to temperature change**

A glass flask with volume 200 cm<sup>3</sup> is filled to the brim with mercury at 20°C. How much mercury overflows when the temperature of the system is raised to 100°C? The coefficient of linear expansion of the glass is  $0.40 \times 10^{-5} \text{ K}^{-1}$ .

**SOLUTION**

**IDENTIFY:** This problem involves the volume expansion of the glass and of the mercury. The amount of overflow depends on the *difference* between the volume changes for these two materials.

**SET UP:** The amount of overflow is equal to the difference between the values of  $\Delta V$  for mercury and for glass, both given by Eq. (17.8). For the mercury to overflow, its coefficient of volume expansion  $\beta$  must be larger than that for glass. The value for mercury is  $\beta_{\text{mercury}} = 18 \times 10^{-5} \text{ K}^{-1}$  from Table 17.2, and we find the value of  $\beta$  for this type of glass from Eq. (17.9),  $\beta = 3\alpha$ .

**EXECUTE:** The coefficient of volume expansion for the glass is

$$\beta_{\text{glass}} = 3\alpha_{\text{glass}} = 3(0.40 \times 10^{-5} \text{ K}^{-1}) = 1.2 \times 10^{-5} \text{ K}^{-1}$$

*Continued*

The increase in volume of the glass flask is

$$\begin{aligned}\Delta V_{\text{glass}} &= \beta_{\text{glass}} V_0 \Delta T \\ &= (1.2 \times 10^{-5} \text{ K}^{-1})(200 \text{ cm}^3)(100^\circ\text{C} - 20^\circ\text{C}) \\ &= 0.19 \text{ cm}^3\end{aligned}$$

The increase in volume of the mercury is

$$\begin{aligned}\Delta V_{\text{mercury}} &= \beta_{\text{mercury}} V_0 \Delta T \\ &= (18 \times 10^{-5} \text{ K}^{-1})(200 \text{ cm}^3)(100^\circ\text{C} - 20^\circ\text{C}) \\ &= 2.9 \text{ cm}^3\end{aligned}$$

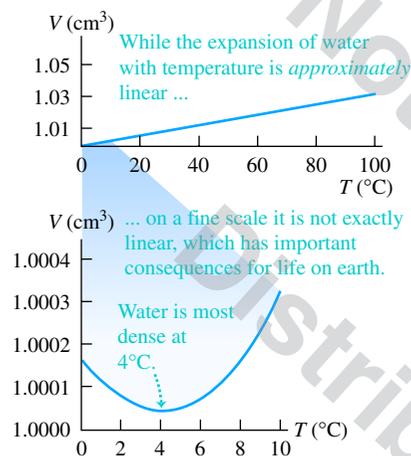
The volume of mercury that overflows is

$$\Delta V_{\text{mercury}} - \Delta V_{\text{glass}} = 2.9 \text{ cm}^3 - 0.19 \text{ cm}^3 = 2.7 \text{ cm}^3$$

**EVALUATE:** This is basically how a mercury-in-glass thermometer works, except that instead of letting the mercury overflow and run all over the place, the thermometer has it rise inside a sealed tube as  $T$  increases.

As Tables 17.1 and 17.2 show, glass has smaller coefficients of expansion  $\alpha$  and  $\beta$  than do most metals. This is why you can use hot water to loosen a metal lid on a glass jar; the metal expands more than the glass does.

**17.12** The volume of 1 gram of water in the temperature range from  $0^\circ\text{C}$  to  $10^\circ\text{C}$ . By  $100^\circ\text{C}$  the volume has increased to  $1.034 \text{ cm}^3$ . If the coefficient of volume expansion were constant, the curve would be a straight line.



**17.13** The interlocking teeth of an expansion joint on a bridge. These joints are needed to accommodate changes in length that result from thermal expansion.



## Thermal Expansion of Water

Water, in the temperature range from  $0^\circ\text{C}$  to  $4^\circ\text{C}$ , *decreases* in volume with increasing temperature. In this range its coefficient of volume expansion is *negative*. Above  $4^\circ\text{C}$ , water expands when heated (Fig. 17.12). Hence water has its greatest density at  $4^\circ\text{C}$ . Water also expands when it freezes, which is why ice humps up in the middle of the compartments in an ice cube tray. By contrast, most materials contract when they freeze.

This anomalous behavior of water has an important effect on plant and animal life in lakes. A lake cools from the surface down; above  $4^\circ\text{C}$ , the cooled water at the surface flows to the bottom because of its greater density. But when the surface temperature drops below  $4^\circ\text{C}$ , the water near the surface is less dense than the warmer water below. Hence the downward flow ceases, and the water near the surface remains colder than that at the bottom. As the surface freezes, the ice floats because it is less dense than water. The water at the bottom remains at  $4^\circ\text{C}$  until nearly the entire lake is frozen. If water behaved like most substances, contracting continuously on cooling and freezing, lakes would freeze from the bottom up. Circulation due to density differences would continuously carry warmer water to the surface for efficient cooling, and lakes would freeze solid much more easily. This would destroy all plant and animal life that cannot withstand freezing. If water did not have this special property, the evolution of life would have taken a very different course.

## Thermal Stress

If we clamp the ends of a rod rigidly to prevent expansion or contraction and then change the temperature, tensile or compressive stresses called **thermal stresses** develop. The rod would like to expand or contract, but the clamps won't let it. The resulting stresses may become large enough to strain the rod irreversibly or even break it. (You may want to review the discussion of stress and strain in Section 11.4).

Engineers must account for thermal stress when designing structures. Concrete highways and bridge decks usually have gaps between sections, filled with a flexible material or bridged by interlocking teeth (Fig. 17.13), to permit expansion and contraction of the concrete. Long steam pipes have expansion joints or U-shaped sections to prevent buckling or stretching with temperature changes. If one end of a steel bridge is rigidly fastened to its abutment, the other end usually rests on rollers.

To calculate the thermal stress in a clamped rod, we compute the amount the rod *would* expand (or contract) if not held and then find the stress needed to com-

press (or stretch) it back to its original length. Suppose that a rod with length  $L_0$  and cross-sectional area  $A$  is held at constant length while the temperature is reduced (negative  $\Delta T$ ), causing a tensile stress. The fractional change in length if the rod were free to contract would be

$$\left(\frac{\Delta L}{L_0}\right)_{\text{thermal}} = \alpha \Delta T \quad (17.10)$$

Both  $\Delta L$  and  $\Delta T$  are negative. The tension must increase by an amount  $F$  that is just enough to produce an equal and opposite fractional change in length  $(\Delta L/L_0)_{\text{tension}}$ . From the definition of Young's modulus, Eq. (11.10),

$$Y = \frac{F/A}{\Delta L/L_0} \quad \text{so} \quad \left(\frac{\Delta L}{L_0}\right)_{\text{tension}} = \frac{F}{AY} \quad (17.11)$$

If the length is to be constant, the *total* fractional change in length must be zero. From Eqs. (17.10) and (17.11), this means that

$$\left(\frac{\Delta L}{L_0}\right)_{\text{thermal}} + \left(\frac{\Delta L}{L_0}\right)_{\text{tension}} = \alpha \Delta T + \frac{F}{AY} = 0$$

Solving for the tensile stress  $F/A$  required to keep the rod's length constant, we find

$$\frac{F}{A} = -Y\alpha \Delta T \quad (\text{thermal stress}) \quad (17.12)$$

For a decrease in temperature,  $\Delta T$  is negative, so  $F$  and  $F/A$  are positive; this means that a *tensile* force and stress are needed to maintain the length. If  $\Delta T$  is positive,  $F$  and  $F/A$  are negative, and the required force and stress are *compressive*.

If there are temperature differences within a body, nonuniform expansion or contraction will result and thermal stresses can be induced. You can break a glass bowl by pouring very hot water into it; the thermal stress between the hot and cold parts of the bowl exceeds the breaking stress of the glass, causing cracks. The same phenomenon makes ice cubes crack when dropped into warm water. Heat-resistant glasses such as Pyrex™ have exceptionally low expansion coefficients and high strength.

### Example 17.5 Thermal stress

An aluminum cylinder 10 cm long, with a cross-sectional area of  $20 \text{ cm}^2$ , is to be used as a spacer between two steel walls. At  $17.2^\circ\text{C}$  it just slips in between the walls. When it warms to  $22.3^\circ\text{C}$ , calculate the stress in the cylinder and the total force it exerts on each wall, assuming that the walls are perfectly rigid and a constant distance apart.

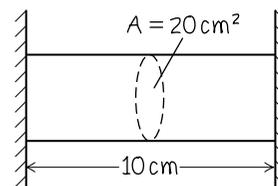
#### SOLUTION

**IDENTIFY:** Our target variables are the thermal stress in the cylinder and the associated force it exerts on each of the walls that holds it in place.

**SET UP:** Figure 17.14 shows our sketch of the situation. We use Eq. (17.12) to relate the stress to the temperature change. The

relevant values of Young's modulus  $Y$  and the coefficient of linear expansion  $\alpha$  are those for aluminum, the material of which the cylinder is made; we find these values from Tables 11.1 and 17.1, respectively.

**17.14** Our sketch for this problem.



*Continued*

**EXECUTE:** For aluminum,  $Y = 7.0 \times 10^{10}$  Pa and  $\alpha = 2.4 \times 10^{-5} \text{ K}^{-1}$ . The temperature change is  $\Delta T = 22.3^\circ\text{C} - 17.2^\circ\text{C} = 5.1 \text{ C}^\circ = 5.1 \text{ K}$ . The stress is  $F/A$ ; from Eq. (17.12),

$$\begin{aligned} \frac{F}{A} &= -Y\alpha\Delta T = -(0.70 \times 10^{11} \text{ Pa})(2.4 \times 10^{-5} \text{ K}^{-1})(5.1 \text{ K}) \\ &= -8.6 \times 10^6 \text{ Pa (or } -1200 \text{ lb/in.}^2\text{)} \end{aligned}$$

The negative sign indicates that compressive rather than tensile stress is needed to keep the cylinder's length constant. This stress

is independent of the length and cross-sectional area of the cylinder. The total force  $F$  is the cross-sectional area times the stress:

$$\begin{aligned} F &= A\left(\frac{F}{A}\right) = (20 \times 10^{-4} \text{ m}^2)(-8.6 \times 10^6 \text{ Pa}) \\ &= -1.7 \times 10^4 \text{ N} \end{aligned}$$

or nearly 2 tons. The negative sign indicates compression.

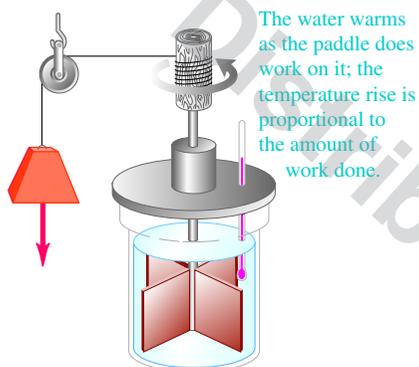
**EVALUATE:** The stress on the cylinder and the force it exerts on each wall are immense. This points out the importance of accounting for such thermal stresses in engineering.

**Test Your Understanding of Section 17.4** In the bimetallic strip shown in Fig. 17.3a, metal 1 is copper. Which of the following materials could be used for metal 2? (There may be more than one correct answer). (i) steel; (ii) brass; (iii) aluminum.

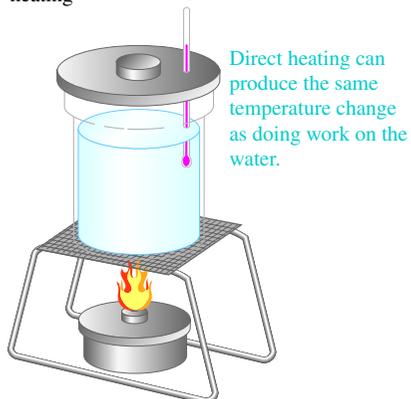
## 17.5 Quantity of Heat

**17.15** The same temperature change of the same system may be accomplished by (a) doing work on it or (b) adding heat to it.

(a) Raising the temperature of water by doing work on it



(b) Raising the temperature of water by direct heating



When you put a cold spoon into a cup of hot coffee, the spoon warms up and the coffee cools down as they approach thermal equilibrium. The interaction that causes these temperature changes is fundamentally a transfer of *energy* from one substance to another. Energy transfer that takes place solely because of a temperature difference is called *heat flow* or *heat transfer*, and energy transferred in this way is called **heat**.

An understanding of the relationship between heat and other forms of energy emerged gradually during the 18th and 19th centuries. Sir James Joule (1818–1889) studied how water can be warmed by vigorous stirring with a paddle wheel (Fig. 17.15a). The paddle wheel adds energy to the water by doing work on it, and Joule found that *the temperature rise is directly proportional to the amount of work done*. The same temperature change can also be caused by putting the water in contact with some hotter body (Fig. 17.15b); hence this interaction must also involve an energy exchange. We will explore the relationship between heat and mechanical energy in greater detail in Chapters 19 and 20.

**CAUTION Temperature vs. heat** It is absolutely essential for you to keep clearly in mind the distinction between *temperature* and *heat*. Temperature depends on the physical state of a material and is a quantitative description of its hotness or coldness. In physics the term “heat” always refers to energy in transit from one body or system to another because of a temperature difference, never to the amount of energy contained within a particular system. We can change the temperature of a body by adding heat to it or taking heat away, or by adding or subtracting energy in other ways, such as mechanical work (Fig. 17.15a). If we cut a body in half, each half has the same temperature as the whole; but to raise the temperature of each half by a given interval, we add *half* as much heat as for the whole. ■

We can define a *unit* of quantity of heat based on temperature changes of some specific material. The **calorie** (abbreviated cal) is defined as *the amount of heat required to raise the temperature of 1 gram of water from 14.5°C to 15.5°C*. The kilocalorie (kcal), equal to 1000 cal, is also used; a food-value calorie is actually a kilocalorie (Fig. 17.16 on the next page). A corresponding unit of heat using Fahrenheit degrees and British units is the **British thermal unit**, or Btu. One Btu

is the quantity of heat required to raise the temperature of 1 pound (weight) of water 1 F° from 63°F to 64°F.

Because heat is energy in transit, there must be a definite relationship between these units and the familiar mechanical energy units such as the joule. Experiments similar in concept to Joule's have shown that

$$1 \text{ cal} = 4.186 \text{ J}$$

$$1 \text{ kcal} = 1000 \text{ cal} = 4186 \text{ J}$$

$$1 \text{ Btu} = 778 \text{ ft} \cdot \text{lb} = 252 \text{ cal} = 1055 \text{ J}$$

The calorie is not a fundamental SI unit. The International Committee on Weights and Measures recommends using the joule as the basic unit of energy in all forms, including heat. We will follow that recommendation in this book.

### Specific Heat

We use the symbol  $Q$  for quantity of heat. When it is associated with an infinitesimal temperature change  $dT$ , we call it  $dQ$ . The quantity of heat  $Q$  required to increase the temperature of a mass  $m$  of a certain material from  $T_1$  to  $T_2$  is found to be approximately proportional to the temperature change  $\Delta T = T_2 - T_1$ . It is also proportional to the mass  $m$  of material. When you're heating water to make tea, you need twice as much heat for two cups as for one if the temperature change is the same. The quantity of heat needed also depends on the nature of the material; raising the temperature of 1 kilogram of water by 1 C° requires 4190 J of heat, but only 910 J is needed to raise the temperature of 1 kilogram of aluminum by 1 C°.

Putting all these relationships together, we have

$$Q = mc \Delta T \quad (\text{heat required for temperature change } \Delta T \text{ of mass } m) \quad (17.13)$$

where  $c$  is a quantity, different for different materials, called the **specific heat** of the material. For an infinitesimal temperature change  $dT$  and corresponding quantity of heat  $dQ$ ,

$$dQ = mc dT \quad (17.14)$$

$$c = \frac{1}{m} \frac{dQ}{dT} \quad (\text{specific heat}) \quad (17.15)$$

In Eqs. (17.13), (17.14), and (17.15),  $Q$  (or  $dQ$ ) and  $\Delta T$  (or  $dT$ ) can be either positive or negative. When they are positive, heat enters the body and its temperature increases; when they are negative, heat leaves the body and its temperature decreases.

**CAUTION The definition of heat** Remember that  $dQ$  does not represent a change in the amount of heat *contained* in a body; this is a meaningless concept. Heat is always energy *in transit* as a result of a temperature difference. There is no such thing as “the amount of heat in a body.” ■

The specific heat of water is approximately

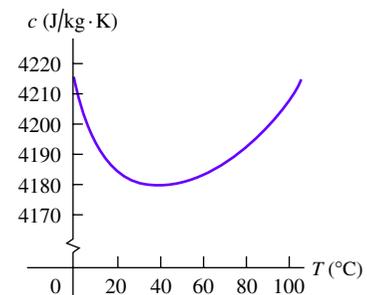
$$4190 \text{ J/kg} \cdot \text{K} \quad 1 \text{ cal/g} \cdot \text{C}^\circ \quad \text{or} \quad 1 \text{ Btu/lb} \cdot \text{F}^\circ$$

The specific heat of a material always depends somewhat on the initial temperature and the temperature interval. Figure 17.17 shows this dependence for water. In the problems and examples in this chapter we will usually ignore this small variation.

**17.16** The motto “Komm in Schwung mit Zucker” on these German sugar packets can be translated as “Sugar gives you momentum.” In fact, sugar gives you *energy*: According to the label, each packet has an energy content of 22 kilocalories (22 food-value calories) or 92 kilojoules. (We discussed the difference between energy and momentum in Section 8.1.)



**17.17** Specific heat of water as a function of temperature. The value of  $c$  varies by less than 1% between 0°C and 100°C.



**Example 17.6 Feed a cold, starve a fever**

During a bout with the flu an 80-kg man ran a fever of  $39.0^{\circ}\text{C}$  ( $102.2^{\circ}\text{F}$ ) instead of the normal body temperature of  $37.0^{\circ}\text{C}$  ( $98.6^{\circ}\text{F}$ ). Assuming that the human body is mostly water, how much heat is required to raise his temperature by that amount?

**SOLUTION**

**IDENTIFY:** This problem uses the relationship among heat (the target variable), mass, specific heat, and temperature change.

**SET UP:** We are given the values of  $m = 80\text{ kg}$ ,  $c = 4190\text{ J/kg}\cdot\text{K}$  (for water), and  $\Delta T = 39.0^{\circ}\text{C} - 37.0^{\circ}\text{C} = 2.0\text{ C}^{\circ} = 2.0\text{ K}$ . We use Eq. (17.13) to determine the required heat.

**EXECUTE:** From Eq. (17.13),

$$Q = mc\Delta T = (80\text{ kg})(4190\text{ J/kg}\cdot\text{K})(2.0\text{ K}) = 6.7 \times 10^5\text{ J}$$

**EVALUATE:** This corresponds to 160 kcal, or 160 food-value calories. In fact, the specific heat of the human body is more nearly equal to  $3480\text{ J/kg}\cdot\text{K}$ , about 83% that of water. The difference is due to the presence of protein, fat, and minerals, which have lower specific heats. With this value of  $c$ , the required heat is  $5.6 \times 10^5\text{ J} = 133\text{ kcal}$ . Either result shows us that were it not for the body's temperature-regulating systems, taking in energy in the form of food would produce measurable changes in body temperature. (In the case of a person with the flu, the elevated temperature results from the body's extra activity in fighting infection.)

**Example 17.7 Overheating electronics**

You are designing an electronic circuit element made of 23 mg of silicon. The electric current through it adds energy at the rate of  $7.4\text{ mW} = 7.4 \times 10^{-3}\text{ J/s}$ . If your design doesn't allow any heat transfer out of the element, at what rate does its temperature increase? The specific heat of silicon is  $705\text{ J/kg}\cdot\text{K}$ .

**SOLUTION**

**IDENTIFY:** The energy added to the circuit element gives rise to a temperature increase, just as if heat were flowing into the element at a rate of  $7.4 \times 10^{-3}\text{ J/s}$ . Our target variable is the *rate* of temperature change.

**SET UP:** From Eq. (17.13), the temperature change  $\Delta T$  in kelvins is proportional to the heat transferred in joules, so the rate of temperature change in  $\text{K/s}$  is proportional to the rate of heat transfer in  $\text{J/s}$ .

**EXECUTE:** In 1 second,  $Q = (7.4 \times 10^{-3}\text{ J/s})(1\text{ s}) = 7.4 \times 10^{-3}\text{ J}$ . From Eq. (17.13),  $Q = mc\Delta T$ , the temperature change in 1 second is

$$\Delta T = \frac{Q}{mc} = \frac{7.4 \times 10^{-3}\text{ J}}{(23 \times 10^{-6}\text{ kg})(705\text{ J/kg}\cdot\text{K})} = 0.46\text{ K}$$

Alternatively, we can divide both sides of Eq. (17.14) by  $dt$  and rearrange:

$$\begin{aligned} \frac{dT}{dt} &= \frac{dQ/dt}{mc} \\ &= \frac{7.4 \times 10^{-3}\text{ J/s}}{(23 \times 10^{-6}\text{ kg})(705\text{ J/kg}\cdot\text{K})} = 0.46\text{ K/s} \end{aligned}$$

**EVALUATE:** At this rate of temperature rise (27 K every minute), the circuit element would soon self-destruct. Heat transfer is an important design consideration in electronic circuit elements.

**Molar Heat Capacity**

Sometimes it's more convenient to describe a quantity of substance in terms of the number of *moles*  $n$  rather than the *mass*  $m$  of material. Recall from your study of chemistry that a mole of any pure substance always contains the same number of molecules. (We will discuss this point in more detail in Chapter 18.) The *molar mass* of any substance, denoted by  $M$ , is the mass per mole. (The quantity  $M$  is sometimes called *molecular weight*, but *molar mass* is preferable; the quantity depends on the mass of a molecule, not its weight.) For example, the molar mass of water is  $18.0\text{ g/mol} = 18.0 \times 10^{-3}\text{ kg/mol}$ ; 1 mole of water has a mass of  $18.0\text{ g} = 0.0180\text{ kg}$ . The total mass  $m$  of material is equal to the mass per mole  $M$  times the number of moles  $n$ :

$$m = nM \quad (17.16)$$

Replacing the mass  $m$  in Eq. (17.13) by the product  $nM$ , we find

$$Q = nMc\Delta T \quad (17.17)$$

The product  $Mc$  is called the **molar heat capacity** (or *molar specific heat*) and is denoted by  $C$  (capitalized). With this notation we rewrite Eq. (17.17) as

$$Q = nC\Delta T \quad (\text{heat required for temperature change of } n \text{ moles}) \quad (17.18)$$

Comparing to Eq. (17.15), we can express the molar heat capacity  $C$  (heat per mole per temperature change) in terms of the specific heat  $c$  (heat per mass per temperature change) and the molar mass  $M$  (mass per mole):

$$C = \frac{1}{n} \frac{dQ}{dT} = Mc \quad (\text{molar heat capacity}) \quad (17.19)$$

For example, the molar heat capacity of water is

$$C = Mc = (0.0180 \text{ kg/mol})(4190 \text{ J/kg} \cdot \text{K}) = 75.4 \text{ J/mol} \cdot \text{K}$$

Values of specific heat and molar heat capacity for several substances are given in Table 17.3. Note the remarkably large specific heat for water (Fig. 17.18).

**CAUTION The meaning of “heat capacity”** The term “heat capacity” is unfortunate because it gives the erroneous impression that a body *contains* a certain amount of heat. Remember, heat is energy in transit to or from a body, not the energy residing in the body. ■

Precise measurements of specific heats and molar heat capacities require great experimental skill. Usually, a measured quantity of energy is supplied by an electric current in a heater wire wound around the specimen. The temperature change  $\Delta T$  is measured with a resistance thermometer or thermocouple embedded in the specimen. This sounds simple, but great care is needed to avoid or compensate for unwanted heat transfer between the sample and its surroundings. Measurements for solid materials are usually made at constant atmospheric pressure; the corresponding values are called the *specific heat* and *molar heat capacity at constant pressure*, denoted by  $c_p$  and  $C_p$ . For a gas it is usually easier to keep the substance in a container with constant *volume*; the corresponding values are called the *specific heat* and *molar heat capacity at constant volume*, denoted by  $c_v$  and  $C_v$ . For a given substance,  $C_v$  and  $C_p$  are different. If the system can expand while heat is added, there is additional energy exchange through the performance of *work* by the system on its surroundings. If the volume is constant, the system does no work. For gases the difference between  $C_p$  and  $C_v$  is substantial. We will study heat capacities of gases in detail in Section 19.7.

The last column of Table 17.3 shows something interesting. The molar heat capacities for most elemental solids are about the same: about  $25 \text{ J/mol} \cdot \text{K}$ . This correlation, named the *rule of Dulong and Petit* (for its discoverers), forms the basis for a very important idea. The number of atoms in 1 mole is the same for all

**17.18** Water has a much higher specific heat than the glass or metals used to make cookware. This helps explain why it takes several minutes to boil water on a stove, even though the pot or kettle reaches a high temperature very quickly.



**Table 17.3** Approximate Specific Heats and Molar Heat Capacities (Constant Pressure)

| Substance                   | Specific Heat, $c$<br>(J/kg · K) | Molar Mass, $M$<br>(kg/mol) | Molar Heat Capacity, $C$<br>(J/mol · K) |
|-----------------------------|----------------------------------|-----------------------------|---|
| Aluminum                    | 910                              | 0.0270                      | 24.6                                    |
| Beryllium                   | 1970                             | 0.00901                     | 17.7                                    |
| Copper                      | 390                              | 0.0635                      | 24.8                                    |
| Ethanol                     | 2428                             | 0.0461                      | 111.9                                   |
| Ethylene glycol             | 2386                             | 0.0620                      | 148.0                                   |
| Ice (near 0°C)              | 2100                             | 0.0180                      | 37.8                                    |
| Iron                        | 470                              | 0.0559                      | 26.3                                    |
| Lead                        | 130                              | 0.207                       | 26.9                                    |
| Marble (CaCO <sub>3</sub> ) | 879                              | 0.100                       | 87.9                                    |
| Mercury                     | 138                              | 0.201                       | 27.7                                    |
| Salt (NaCl)                 | 879                              | 0.0585                      | 51.4                                    |
| Silver                      | 234                              | 0.108                       | 25.3                                    |
| Water (liquid)              | 4190                             | 0.0180                      | 75.4                                    |

elemental substances. This means that on a *per atom* basis, about the same amount of heat is required to raise the temperature of each of these elements by a given amount, even though the *masses* of the atoms are very different. The heat required for a given temperature increase depends only on *how many* atoms the sample contains, not on the mass of an individual atom. We will see the reason the rule of Dulong and Petit works so well when we study the molecular basis of heat capacities in greater detail in Chapter 18.

**Test Your Understanding of Section 17.5** You wish to raise the temperature of each of the following samples from 20°C to 21°C. Rank these in order of the amount of heat needed to do this, from highest to lowest. (i) 1 kilogram of mercury; (ii) 1 kilogram of ethanol; (iii) 1 mole of mercury; (iv) 1 mole of ethanol.



## 17.6 Calorimetry and Phase Changes

Calorimetry means “measuring heat.” We have discussed the energy transfer (heat) involved in temperature changes. Heat is also involved in *phase changes*, such as the melting of ice or boiling of water. Once we understand these additional heat relationships, we can analyze a variety of problems involving quantity of heat.

### Phase Changes

We use the term **phase** to describe a specific state of matter, such as a solid, liquid, or gas. The compound H<sub>2</sub>O exists in the *solid phase* as ice, in the *liquid phase* as water, and in the *gaseous phase* as steam. (These are also referred to as **states of matter**: the solid state, the liquid state, and the gaseous state.) A transition from one phase to another is called a **phase change** or *phase transition*. For any given pressure a phase change takes place at a definite temperature, usually accompanied by absorption or emission of heat and a change of volume and density.

A familiar example of a phase change is the melting of ice. When we add heat to ice at 0°C and normal atmospheric pressure, the temperature of the ice *does not* increase. Instead, some of it melts to form liquid water. If we add the heat slowly, to maintain the system very close to thermal equilibrium, the temperature remains at 0°C until all the ice is melted (Fig. 17.19). The effect of adding heat to this system is not to raise its temperature but to change its *phase* from solid to liquid.

To change 1 kg of ice at 0°C to 1 kg of liquid water at 0°C and normal atmospheric pressure requires  $3.34 \times 10^5$  J of heat. The heat required per unit mass is called the **heat of fusion** (or sometimes *latent heat of fusion*), denoted by  $L_f$ . For water at normal atmospheric pressure the heat of fusion is

$$L_f = 3.34 \times 10^5 \text{ J/kg} = 79.6 \text{ cal/g} = 143 \text{ Btu/lb}$$

More generally, to melt a mass  $m$  of material that has a heat of fusion  $L_f$  requires a quantity of heat  $Q$  given by

$$Q = mL_f$$

This process is *reversible*. To freeze liquid water to ice at 0°C, we have to *remove* heat; the magnitude is the same, but in this case,  $Q$  is negative because heat is removed rather than added. To cover both possibilities and to include other kinds of phase changes, we write

$$Q = \pm mL \quad (\text{heat transfer in a phase change}) \quad (17.20)$$

The plus sign (heat entering) is used when the material melts; the minus sign (heat leaving) is used when it freezes. The heat of fusion is different for different materials, and it also varies somewhat with pressure.

**17.19** The surrounding air is at room temperature, but this ice–water mixture remains at 0°C until all of the ice has melted and the phase change is complete.



For any given material at any given pressure, the freezing temperature is the same as the melting temperature. At this unique temperature the liquid and solid phases (liquid water and ice, for example) can coexist in a condition called **phase equilibrium**.

We can go through this whole story again for *boiling* or *evaporation*, a phase transition between liquid and gaseous phases. The corresponding heat (per unit mass) is called the **heat of vaporization**  $L_v$ . At normal atmospheric pressure the heat of vaporization  $L_v$  for water is

$$L_v = 2.256 \times 10^6 \text{ J/kg} = 539 \text{ cal/g} = 970 \text{ Btu/lb}$$

That is, it takes  $2.256 \times 10^6 \text{ J}$  to change 1 kg of liquid water at  $100^\circ\text{C}$  to 1 kg of water vapor at  $100^\circ\text{C}$ . By comparison, to raise the temperature of 1 kg of water from  $0^\circ\text{C}$  to  $100^\circ\text{C}$  requires  $Q = mc \Delta T = (1.00 \text{ kg})(4190 \text{ J/kg} \cdot \text{C}^\circ) \times (100 \text{ C}^\circ) = 4.19 \times 10^5 \text{ J}$ , less than one-fifth as much heat as is required for vaporization at  $100^\circ\text{C}$ . This agrees with everyday kitchen experience; a pot of water may reach boiling temperature in a few minutes, but it takes a much longer time to completely evaporate all the water away.

Like melting, boiling is a reversible transition. When heat is removed from a gas at the boiling temperature, the gas returns to the liquid phase, or *condenses*, giving up to its surroundings the same quantity of heat (heat of vaporization) that was needed to vaporize it. At a given pressure the boiling and condensation temperatures are always the same; at this temperature the liquid and gaseous phases can coexist in phase equilibrium.

Both  $L_v$  and the boiling temperature of a material depend on pressure. Water boils at a lower temperature (about  $95^\circ\text{C}$ ) in Denver than in Pittsburgh because Denver is at higher elevation and the average atmospheric pressure is lower. The heat of vaporization is somewhat greater at this lower pressure, about  $2.27 \times 10^6 \text{ J/kg}$ .

Table 17.4 lists heats of fusion and vaporization for several materials and their melting and boiling temperatures at normal atmospheric pressure. Very few *elements* have melting temperatures in the vicinity of ordinary room temperatures; one of the few is the metal gallium, shown in Fig. 17.20.

**17.20** The metal gallium, shown here melting in a person's hand, is one of the few elements that melt in the vicinity of room temperature. Its melting temperature is  $29.8^\circ\text{C}$ , and its heat of fusion is  $8.04 \times 10^4 \text{ J/kg}$ .



**Table 17.4** Heats of Fusion and Vaporization

| Substance | Normal Melting Point |                  | Heat of Fusion, $L_f$ (J/kg) | Normal Boiling Point |                  | Heat of Vaporization, $L_v$ (J/kg) |
|-----------|----------------------|------------------|------------------------------|----------------------|------------------|------------------------------------|
|           | K                    | $^\circ\text{C}$ |                              | K                    | $^\circ\text{C}$ |                                    |
| Helium    | *                    | *                | *                            | 4.216                | -268.93          | $20.9 \times 10^3$                 |
| Hydrogen  | 13.84                | -259.31          | $58.6 \times 10^3$           | 20.26                | -252.89          | $452 \times 10^3$                  |
| Nitrogen  | 63.18                | -209.97          | $25.5 \times 10^3$           | 77.34                | -195.8           | $201 \times 10^3$                  |
| Oxygen    | 54.36                | -218.79          | $13.8 \times 10^3$           | 90.18                | -183.0           | $213 \times 10^3$                  |
| Ethanol   | 159                  | -114             | $104.2 \times 10^3$          | 351                  | 78               | $854 \times 10^3$                  |
| Mercury   | 234                  | -39              | $11.8 \times 10^3$           | 630                  | 357              | $272 \times 10^3$                  |
| Water     | 273.15               | 0.00             | $334 \times 10^3$            | 373.15               | 100.00           | $2256 \times 10^3$                 |
| Sulfur    | 392                  | 119              | $38.1 \times 10^3$           | 717.75               | 444.60           | $326 \times 10^3$                  |
| Lead      | 600.5                | 327.3            | $24.5 \times 10^3$           | 2023                 | 1750             | $871 \times 10^3$                  |
| Antimony  | 903.65               | 630.50           | $165 \times 10^3$            | 1713                 | 1440             | $561 \times 10^3$                  |
| Silver    | 1233.95              | 960.80           | $88.3 \times 10^3$           | 2466                 | 2193             | $2336 \times 10^3$                 |
| Gold      | 1336.15              | 1063.00          | $64.5 \times 10^3$           | 2933                 | 2660             | $1578 \times 10^3$                 |
| Copper    | 1356                 | 1083             | $134 \times 10^3$            | 1460                 | 1187             | $5069 \times 10^3$                 |

\*A pressure in excess of 25 atmospheres is required to make helium solidify. At 1 atmosphere pressure, helium remains a liquid down to absolute zero.

**17.21** Graph of temperature versus time for a specimen of water initially in the solid phase (ice). Heat is added to the specimen at a constant rate. The temperature remains constant during each change of phase, provided that the pressure remains constant.

**Phase of water changes.** During these periods, temperature stays constant and the phase change proceeds as heat is added:  $Q = +mL$ .

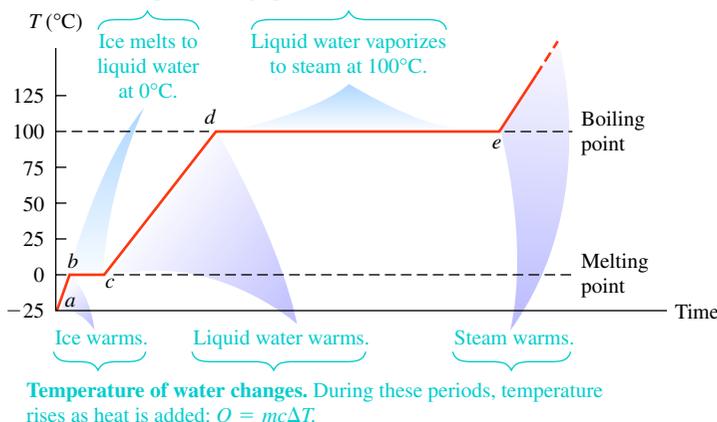


Figure 17.21 shows how the temperature varies when we add heat continuously to a specimen of ice with an initial temperature below  $0^\circ\text{C}$  (point  $a$ ). The temperature rises until we reach the melting point (point  $b$ ). As more heat is added, the temperature remains constant until all the ice has melted (point  $c$ ). Then the temperature rises again until the boiling temperature is reached (point  $d$ ). At that point the temperature again is constant until all the water is transformed into the vapor phase (point  $e$ ). If the rate of heat input is constant, the line for the solid phase (ice) has a steeper slope than does the line for the liquid phase (water). Do you see why? (See Table 17.3.)

A substance can sometimes change directly from the solid to the gaseous phase. This process is called *sublimation*, and the solid is said to *sublime*. The corresponding heat is called the *heat of sublimation*,  $L_s$ . Liquid carbon dioxide cannot exist at a pressure lower than about  $5 \times 10^5$  Pa (about 5 atm), and “dry ice” (solid carbon dioxide) sublimates at atmospheric pressure. Sublimation of water from frozen food causes freezer burn. The reverse process, a phase change from gas to solid, occurs when frost forms on cold bodies such as refrigerator cooling coils.

Very pure water can be cooled several degrees below the freezing temperature without freezing; the resulting unstable state is described as *supercooled*. When a small ice crystal is dropped in or the water is agitated, it crystallizes within a second or less. Supercooled water *vapor* condenses quickly into fog droplets when a disturbance, such as dust particles or ionizing radiation, is introduced. This principle is used in “seeding” clouds, which often contain supercooled water vapor, to cause condensation and rain.

A liquid can sometimes be *superheated* above its normal boiling temperature. Any small disturbance such as agitation causes local boiling with bubble formation.

Steam heating systems for buildings use a boiling–condensing process to transfer heat from the furnace to the radiators. Each kilogram of water that is turned to steam in the boiler absorbs over  $2 \times 10^6$  J (the heat of vaporization  $L_v$  of water) from the boiler and gives it up when it condenses in the radiators. Boiling–condensing processes are also used in refrigerators, air conditioners, and heat pumps. We will discuss these systems in Chapter 20.

The temperature-control mechanisms of many warm-blooded animals make use of heat of vaporization, removing heat from the body by using it to evaporate water from the tongue (panting) or from the skin (sweating). Evaporative cooling enables humans to maintain normal body temperature in hot, dry desert climates where the air temperature may reach  $55^\circ\text{C}$  (about  $130^\circ\text{F}$ ). The skin temperature may be as much as  $30^\circ\text{C}$  cooler than the surrounding air. Under these conditions a normal person may perspire several liters per day, and this lost water must be replaced. Old-time desert rats (such as one of the authors) state that in the desert, any canteen that holds less than a gallon should be viewed as a toy! Evaporative

cooling also explains why you feel cold when you first step out of a swimming pool (Fig. 17.22).

Evaporative cooling is also used to cool buildings in hot, dry climates and to condense and recirculate “used” steam in coal-fired or nuclear-powered electric-generating plants. That’s what goes on in the large, tapered concrete towers that you see at such plants.

Chemical reactions such as combustion are analogous to phase changes in that they involve definite quantities of heat. Complete combustion of 1 gram of gasoline produces about 46,000 J or about 11,000 cal, so the **heat of combustion**  $L_c$  of gasoline is

$$L_c = 46,000 \text{ J/g} = 4.6 \times 10^7 \text{ J/kg}$$

Energy values of foods are defined similarly. When we say that a gram of peanut butter “contains 6 calories,” we mean that 6 kcal of heat (6,000 cal or 25,000 J) is released when the carbon and hydrogen atoms in the peanut butter react with oxygen (with the help of enzymes) and are completely converted to  $\text{CO}_2$  and  $\text{H}_2\text{O}$ . Not all of this energy is directly useful for mechanical work. We will study the *efficiency* of energy utilization in Chapter 20.

### Heat Calculations

Let’s look at some examples of calorimetry calculations (calculations with heat). The basic principle is very simple: When heat flow occurs between two bodies that are isolated from their surroundings, the amount of heat lost by one body must equal the amount gained by the other. Heat is energy in transit, so this principle is really just conservation of energy. Calorimetry, dealing entirely with one conserved quantity, is in many ways the simplest of all physical theories!

**17.22** The water may be warm and it may be a hot day, but these children will feel cold when they first step out of the swimming pool. That’s because as water evaporates from their skin, it removes the heat of vaporization from their bodies. To stay warm, they will need to dry off immediately.



#### Problem-Solving Strategy 17.2 Calorimetry Problems



**IDENTIFY** *the relevant concepts:* When heat flow occurs between two bodies that are isolated from their surroundings, the amount of heat lost by one body must equal the amount gained by the other body.

**SET UP** *the problem* using the following steps:

1. Identify which objects exchange heat. To avoid confusion with algebraic signs, take each quantity of heat *added* to a body as *positive* and each quantity *leaving* a body as *negative*. When two or more bodies interact, the *algebraic sum* of the quantities of heat transferred to all the bodies must be zero.
2. Each object will undergo a temperature change with no phase change, a phase change at constant temperature, or both. Use Eq. (17.13) to describe temperature changes and Eq. (17.20) to describe phase changes.
3. Consult Table 17.3 for values of the specific heat or molar heat capacity and Table 17.4 for heats of fusion or vaporization.
4. Be certain to identify which quantities are known and which are the unknown target variables.

**EXECUTE** *the solution* as follows:

1. Solve Eq. (17.13) and/or Eq. (17.20) for the target variables. Often you will need to find an unknown temperature. Represent it by an algebraic symbol such as  $T$ . Then if a body has an initial temperature of  $20^\circ\text{C}$  and an unknown final temperature  $T$ , the temperature change for the body is  $\Delta T = T_{\text{final}} - T_{\text{initial}} = T - 20^\circ\text{C}$  (not  $20^\circ\text{C} - T$ ).
2. In problems where a phase change takes place, as when ice melts, you may not know in advance whether *all* the material undergoes a phase change or only part of it. You can always assume one or the other, and if the resulting calculation gives an absurd result (such as a final temperature higher or lower than *any* of the initial temperatures), you know the initial assumption was wrong. Back up and try again!

**EVALUATE** *your answer:* A common error is to use the wrong algebraic sign for either a  $Q$  or  $\Delta T$  term. Double check your calculations, and make sure that the final results are physically sensible.

#### Example 17.8 A temperature change with no phase change

A geologist working in the field drinks her morning coffee out of an aluminum cup. The cup has a mass of 0.120 kg and is initially at  $20.0^\circ\text{C}$  when she pours in 0.300 kg of coffee initially at  $70.0^\circ\text{C}$ . What is the final temperature after the coffee and the cup attain thermal equilibrium? (Assume that coffee has the same specific heat as water and that there is no heat exchange with the surroundings.)

#### SOLUTION

**IDENTIFY:** The two objects we must consider are the cup and the coffee, and the target variable is their common final temperature.

**SET UP:** No phase changes occur in this situation, so the only equation we need is Eq. (17.13).

*Continued*

**EXECUTE:** By using Table 17.3, the (negative) heat gained by the coffee is

$$\begin{aligned} Q_{\text{coffee}} &= m_{\text{coffee}} c_{\text{water}} \Delta T_{\text{coffee}} \\ &= (0.300 \text{ kg})(4190 \text{ J/kg} \cdot \text{K})(T - 70.0^\circ\text{C}) \end{aligned}$$

The (positive) heat gained by the aluminum cup is

$$\begin{aligned} Q_{\text{aluminum}} &= m_{\text{aluminum}} c_{\text{aluminum}} \Delta T_{\text{aluminum}} \\ &= (0.120 \text{ kg})(910 \text{ J/kg} \cdot \text{K})(T - 20.0^\circ\text{C}) \end{aligned}$$

We equate the sum of these two quantities of heat to zero, obtaining an algebraic equation for  $T$ :

$$\begin{aligned} Q_{\text{coffee}} + Q_{\text{aluminum}} &= 0 \quad \text{or} \\ (0.300 \text{ kg})(4190 \text{ J/kg} \cdot \text{K})(T - 70.0^\circ\text{C}) \\ &+ (0.120 \text{ kg})(910 \text{ J/kg} \cdot \text{K})(T - 20.0^\circ\text{C}) = 0 \end{aligned}$$

Solution of this equation gives  $T = 66.0^\circ\text{C}$ .

**EVALUATE:** The final temperature is much closer to the initial temperature of the coffee than to that of the cup; water has a much higher specific heat than aluminum, and we have more than twice as much mass of water. We can also find the quantities of heat by substituting the value  $T = 66.0^\circ\text{C}$  back into the original equations. We find that  $Q_{\text{coffee}} = -5.0 \times 10^3 \text{ J}$  and  $Q_{\text{aluminum}} = +5.0 \times 10^3 \text{ J}$ ;  $Q_{\text{coffee}}$  is negative, which means that the coffee loses heat.

### Example 17.9 Changes in both temperature and phase

A physics student wants to cool 0.25 kg of Diet Omni-Cola (mostly water), initially at  $25^\circ\text{C}$ , by adding ice initially at  $-20^\circ\text{C}$ . How much ice should she add so that the final temperature will be  $0^\circ\text{C}$  with all the ice melted if the heat capacity of the container may be neglected?

#### SOLUTION

**IDENTIFY:** The ice and the Omni-Cola are the objects that exchange heat. The Omni-Cola undergoes a temperature change only, while the ice undergoes both a temperature change and a phase change from solid to liquid. The target variable is the mass of ice,  $m_{\text{ice}}$ .

**SET UP:** We use Eq. (17.13) to find the amount of heat involved in warming the ice to  $0^\circ\text{C}$  and cooling the Omni-Cola to  $0^\circ\text{C}$ . In addition, we'll need Eq. (17.20) to calculate the heat required to melt the ice at  $0^\circ\text{C}$ .

**EXECUTE:** The Omni-Cola loses heat, so the heat added to it is negative:

$$\begin{aligned} Q_{\text{Omni}} &= m_{\text{Omni}} c_{\text{water}} \Delta T_{\text{Omni}} \\ &= (0.25 \text{ kg})(4190 \text{ J/kg} \cdot \text{K})(0^\circ\text{C} - 25^\circ\text{C}) \\ &= -26,000 \text{ J} \end{aligned}$$

From Table 17.3, the specific heat of ice (not the same as for liquid water) is  $2.1 \times 10^3 \text{ J/kg} \cdot \text{K}$ . Let the mass of ice be  $m_{\text{ice}}$ ; then the heat  $Q_1$  needed to warm it from  $-20^\circ\text{C}$  to  $0^\circ\text{C}$  is

$$\begin{aligned} Q_1 &= m_{\text{ice}} c_{\text{ice}} \Delta T_{\text{ice}} \\ &= m_{\text{ice}}(2.1 \times 10^3 \text{ J/kg} \cdot \text{K})[0^\circ\text{C} - (-20^\circ\text{C})] \\ &= m_{\text{ice}}(4.2 \times 10^4 \text{ J/kg}) \end{aligned}$$

From Eq. (17.20) the additional heat  $Q_2$  needed to melt this mass of ice is the mass times the heat of fusion. Using Table 17.4, we find

$$\begin{aligned} Q_2 &= m_{\text{ice}} L_f \\ &= m_{\text{ice}}(3.34 \times 10^5 \text{ J/kg}) \end{aligned}$$

The sum of these three quantities must equal zero:

$$\begin{aligned} Q_{\text{Omni}} + Q_1 + Q_2 &= -26,000 \text{ J} + m_{\text{ice}}(42,000 \text{ J/kg}) \\ &+ m_{\text{ice}}(334,000 \text{ J/kg}) = 0 \end{aligned}$$

Solving this for  $m_{\text{ice}}$ , we get  $m_{\text{ice}} = 0.069 \text{ kg} = 69 \text{ g}$ .

**EVALUATE:** This mass of ice corresponds to three or four medium-size ice cubes, which seems reasonable for the quantity of Omni-Cola in this problem.

### Example 17.10 What's cooking?

A heavy copper pot of mass 2.0 kg (including the copper lid) is at a temperature of  $150^\circ\text{C}$ . You pour 0.10 kg of water at  $25^\circ\text{C}$  into the pot, then quickly close the lid of the pot so that no steam can escape. Find the final temperature of the pot and its contents, and determine the phase (liquid or gas) of the water. Assume that no heat is lost to the surroundings.

#### SOLUTION

**IDENTIFY:** The two objects that exchange heat are the water and the pot. Note that there are three conceivable outcomes in this situation. One, none of the water boils, and the final temperature is less than  $100^\circ\text{C}$ ; two, a portion of the water boils, giving a mixture of water and steam at  $100^\circ\text{C}$ ; or three, all the water boils, giving 0.10 kg of steam at a temperature of  $100^\circ\text{C}$  or greater.

**SET UP:** We again use Eq. (17.13) for the heat transferred in a temperature change and Eq. (17.20) for the heat transferred in a phase change.

**EXECUTE:** The simplest case to calculate is the first possibility. Let the common final temperature of the liquid water and the copper pot be  $T$ . Since we are assuming that no phase changes take place, the sum of the quantities of heat added to the two materials is

$$\begin{aligned} Q_{\text{water}} + Q_{\text{copper}} &= m_{\text{water}} c_{\text{water}}(T - 25^\circ\text{C}) \\ &+ m_{\text{copper}} c_{\text{copper}}(T - 150^\circ\text{C}) \\ &= (0.10 \text{ kg})(4190 \text{ J/kg} \cdot \text{K})(T - 25^\circ\text{C}) \\ &+ (2.0 \text{ kg})(390 \text{ J/kg} \cdot \text{K})(T - 150^\circ\text{C}) \\ &= 0 \end{aligned}$$

Solving this for  $T$ , we find  $T = 106^\circ\text{C}$ . But this is above the boiling point of water, which contradicts our assumption that none of the water boils! So this assumption can't be correct; at least some of the water undergoes a phase change.

If we try the second possibility, in which the final temperature is  $100^\circ\text{C}$ , we have to find the fraction of water that changes to the gaseous phase. Let this fraction be  $x$ ; the (positive) amount of heat needed to vaporize this water is  $(xm_{\text{water}})L_v$ . Setting the final temperature  $T$  equal to  $100^\circ\text{C}$ , we have

$$\begin{aligned} Q_{\text{water}} &= m_{\text{water}}c_{\text{water}}(100^\circ\text{C} - 25^\circ\text{C}) + xm_{\text{water}}L_v \\ &= (0.10 \text{ kg})(4190 \text{ J/kg} \cdot \text{K})(75 \text{ K}) \\ &\quad + x(0.10 \text{ kg})(2.256 \times 10^6 \text{ J/kg}) \\ &= 3.14 \times 10^4 \text{ J} + x(2.256 \times 10^5 \text{ J}) \\ Q_{\text{copper}} &= m_{\text{copper}}c_{\text{copper}}(100^\circ\text{C} - 150^\circ\text{C}) \\ &= (2.0 \text{ kg})(390 \text{ J/kg} \cdot \text{K})(-50 \text{ K}) = -3.90 \times 10^4 \text{ J} \end{aligned}$$

Now require that the sum of all the quantities of heat be zero:

$$\begin{aligned} Q_{\text{water}} + Q_{\text{copper}} &= 3.14 \times 10^4 \text{ J} + x(2.256 \times 10^5 \text{ J}) \\ &\quad - 3.90 \times 10^4 \text{ J} = 0 \\ x &= \frac{3.90 \times 10^4 \text{ J} - 3.14 \times 10^4 \text{ J}}{2.256 \times 10^5 \text{ J}} = 0.034 \end{aligned}$$

This makes sense, and we conclude that the final temperature of the water and copper is  $100^\circ\text{C}$ . Of the original 0.10 kg of water,  $0.034(0.10 \text{ kg}) = 0.0034 \text{ kg} = 3.4 \text{ g}$  has been converted to steam at  $100^\circ\text{C}$ .

**EVALUATE:** Had  $x$  turned out to be greater than 1, we would have again had a contradiction (the fraction of water that vaporized can't be greater than 1). In this case the third possibility would have been the correct description, all the water would have vaporized, and the final temperature would have been greater than  $100^\circ\text{C}$ . Can you show that this would have been the case if we had originally poured less than 15 g of  $25^\circ\text{C}$  water into the pot?

### Example 17.11 Combustion, temperature change, and phase change

In a particular gasoline camp stove, 30% of the energy released in burning the fuel actually goes to heating the water in the pot on the stove. If we heat 1.00 L (1.00 kg) of water from  $20^\circ\text{C}$  to  $100^\circ\text{C}$  and boil 0.25 kg of it away, how much gasoline do we burn in the process?

#### SOLUTION

**IDENTIFY:** In this problem all of the water undergoes a temperature change and part of the water also undergoes a phase change from liquid to gas. This requires a certain amount of heat, which we use to determine the amount of gasoline that must be burned (the target variable).

**SET UP:** We use Eqs. (17.13) and (17.20) as well as the idea of heat of combustion.

**EXECUTE:** The heat required to raise the temperature of the water from  $20^\circ\text{C}$  to  $100^\circ\text{C}$  is

$$\begin{aligned} Q_1 &= mc \Delta T = (1.00 \text{ kg})(4190 \text{ J/kg} \cdot \text{K})(80 \text{ K}) \\ &= 3.35 \times 10^5 \text{ J} \end{aligned}$$

To boil 0.25 kg of water at  $100^\circ\text{C}$  requires

$$Q_2 = mL_v = (0.25 \text{ kg})(2.256 \times 10^6 \text{ J/kg}) = 5.64 \times 10^5 \text{ J}$$

The total energy needed is the sum of these, or  $8.99 \times 10^5 \text{ J}$ . This is only 0.30 of the total heat of combustion, so that energy is  $(8.99 \times 10^5 \text{ J})/0.30 = 3.00 \times 10^6 \text{ J}$ . As we mentioned earlier, 1 gram of gasoline releases 46,000 J, so the mass of gasoline required is

$$\frac{3.00 \times 10^6 \text{ J}}{46,000 \text{ J/g}} = 65 \text{ g}$$

or a volume of about 0.09 L of gasoline.

**EVALUATE:** This result is a testament to the tremendous amount of energy that can be released by burning even a small quantity of gasoline. Note that most of the heat delivered was used to boil away 0.25 L of water. Can you show that another 123 g of gasoline would be required to boil away the remaining water?

**Test Your Understanding of Section 17.6** You take a block of ice at  $0^\circ\text{C}$  and add heat to it at a steady rate. It takes a time  $t$  to completely convert the block of ice to steam at  $100^\circ\text{C}$ . What do you have at time  $t/2$ ? (i) all ice at  $0^\circ\text{C}$ ; (ii) a mixture of ice and water at  $0^\circ\text{C}$ ; (iii) water at a temperature between  $0^\circ\text{C}$  and  $100^\circ\text{C}$ ; (iv) a mixture of water and steam at  $100^\circ\text{C}$ .

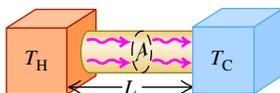


## 17.7 Mechanisms of Heat Transfer

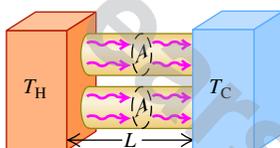
We have talked about *conductors* and *insulators*, materials that permit or prevent heat transfer between bodies. Now let's look in more detail at *rates* of energy transfer. In the kitchen you use a metal or glass pot for good heat transfer from the stove to whatever you're cooking, but your refrigerator is insulated with a material that *prevents* heat from flowing into the food inside the refrigerator. How do we describe the difference between these two materials?

**17.23** Steady-state heat flow due to conduction in a uniform rod.

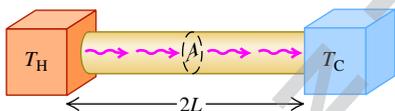
(a) Heat current  $H$



(b) Doubling the cross-sectional area of the conductor doubles the heat current ( $H$  is proportional to  $A$ ).



(c) Doubling the length of the conductor halves the heat current ( $H$  is inversely proportional to  $L$ ).



**Table 17.5** Thermal Conductivities

| Substance                             | $k$ (W/m · K) |
|---------------------------------------|---------------|
| <i>Metals</i>                         |               |
| Aluminum                              | 205.0         |
| Brass                                 | 109.0         |
| Copper                                | 385.0         |
| Lead                                  | 34.7          |
| Mercury                               | 8.3           |
| Silver                                | 406.0         |
| Steel                                 | 50.2          |
| <i>Solids (representative values)</i> |               |
| Brick, insulating                     | 0.15          |
| Brick, red                            | 0.6           |
| Concrete                              | 0.8           |
| Cork                                  | 0.04          |
| Felt                                  | 0.04          |
| Fiberglass                            | 0.04          |
| Glass                                 | 0.8           |
| Ice                                   | 1.6           |
| Rock wool                             | 0.04          |
| Styrofoam                             | 0.01          |
| Wood                                  | 0.12–0.04     |
| <i>Gases</i>                          |               |
| Air                                   | 0.024         |
| Argon                                 | 0.016         |
| Helium                                | 0.14          |
| Hydrogen                              | 0.14          |
| Oxygen                                | 0.023         |

The three mechanisms of heat transfer are conduction, convection, and radiation. *Conduction* occurs within a body or between two bodies in contact. *Convection* depends on motion of mass from one region of space to another. *Radiation* is heat transfer by electromagnetic radiation, such as sunshine, with no need for matter to be present in the space between bodies.

**Conduction**

If you hold one end of a copper rod and place the other end in a flame, the end you are holding gets hotter and hotter, even though it is not in direct contact with the flame. Heat reaches the cooler end by **conduction** through the material. On the atomic level, the atoms in the hotter regions have more kinetic energy, on the average, than their cooler neighbors. They jostle their neighbors, giving them some of their energy. The neighbors jostle *their* neighbors, and so on through the material. The atoms themselves do not move from one region of material to another, but their energy does.

Most metals also use another, more effective mechanism to conduct heat. Within the metal, some electrons can leave their parent atoms and wander through the crystal lattice. These “free” electrons can rapidly carry energy from the hotter to the cooler regions of the metal, so metals are generally good conductors of heat. A metal rod at 20°C feels colder than a piece of wood at 20°C because heat can flow more easily from your hand into the metal. The presence of “free” electrons also causes most metals to be good electrical conductors.

Heat transfer occurs only between regions that are at different temperatures, and the direction of heat flow is always from higher to lower temperature. Figure 17.23a shows a rod of conducting material with cross-sectional area  $A$  and length  $L$ . The left end of the rod is kept at a temperature  $T_H$  and the right end at a lower temperature  $T_C$ , so heat flows from left to right. The sides of the rod are covered by an ideal insulator, so no heat transfer occurs at the sides.

When a quantity of heat  $dQ$  is transferred through the rod in a time  $dt$ , the rate of heat flow is  $dQ/dt$ . We call this rate the **heat current**, denoted by  $H$ . That is,  $H = dQ/dt$ . Experiments show that the heat current is proportional to the cross-sectional area  $A$  of the rod (Fig. 17.23b) and to the temperature difference ( $T_H - T_C$ ) and is inversely proportional to the rod length  $L$  (Fig. 17.23c). Introducing a proportionality constant  $k$  called the **thermal conductivity** of the material, we have

$$H = \frac{dQ}{dt} = kA \frac{T_H - T_C}{L} \quad (\text{heat current in conduction}) \quad (17.21)$$

The quantity  $(T_H - T_C)/L$  is the temperature difference *per unit length*; it is called the magnitude of the **temperature gradient**. The numerical value of  $k$  depends on the material of the rod. Materials with large  $k$  are good conductors of heat; materials with small  $k$  are poor conductors or insulators. Equation (17.21) also gives the heat current through a slab or through *any* homogeneous body with uniform cross section  $A$  perpendicular to the direction of flow;  $L$  is the length of the heat-flow path.

The units of heat current  $H$  are units of energy per time, or power; the SI unit of heat current is the watt ( $1 \text{ W} = 1 \text{ J/s}$ ). We can find the units of  $k$  by solving Eq. (17.21) for  $k$ ; you can show that the SI units are  $\text{W/m} \cdot \text{K}$ . Some numerical values of  $k$  are given in Table 17.5.

The thermal conductivity of “dead” (that is, nonmoving) air is very small. A wool sweater keeps you warm because it traps air between the fibers. In fact,

many insulating materials such as Styrofoam and fiberglass are mostly dead air. Figure 17.24 shows a ceramic material with very unusual thermal properties, including very small conductivity.

If the temperature varies in a nonuniform way along the length of the conducting rod, we introduce a coordinate  $x$  along the length and generalize the temperature gradient to be  $dT/dx$ . The corresponding generalization of Eq. (17.21) is

$$H = \frac{dQ}{dt} = -kA \frac{dT}{dx} \quad (17.22)$$

The negative sign shows that heat always flows in the direction of *decreasing* temperature.

For thermal insulation in buildings, engineers use the concept of **thermal resistance**, denoted by  $R$ . The thermal resistance  $R$  of a slab of material with area  $A$  is defined so that the heat current  $H$  through the slab is

$$H = \frac{A(T_H - T_C)}{R} \quad (17.23)$$

where  $T_H$  and  $T_C$  are the temperatures on the two sides of the slab. Comparing this with Eq. (17.21), we see that  $R$  is given by

$$R = \frac{L}{k} \quad (17.24)$$

where  $L$  is the thickness of the slab. The SI unit of  $R$  is  $1 \text{ m}^2 \cdot \text{K}/\text{W}$ . In the units used for commercial insulating materials in the United States,  $H$  is expressed in Btu/h,  $A$  is in  $\text{ft}^2$ , and  $T_H - T_C$  in  $^\circ\text{F}$ . (1 Btu/h = 0.293 W.) The units of  $R$  are then  $\text{ft}^2 \cdot ^\circ\text{F} \cdot \text{h}/\text{Btu}$ , though values of  $R$  are usually quoted without units; a 6-inch-thick layer of fiberglass has an  $R$  value of 19 (that is,  $R = 19 \text{ ft}^2 \cdot ^\circ\text{F} \cdot \text{h}/\text{Btu}$ ), a 2-inch-thick slab of polyurethane foam has an  $R$  value of 12, and so on. Doubling the thickness doubles the  $R$  value. Common practice in new construction in severe northern climates is to specify  $R$  values of around 30 for exterior walls and ceilings. When the insulating material is in layers, such as a plastered wall, fiberglass insulation, and wood exterior siding, the  $R$  values are additive. Do you see why? (See Problem 17.110.)

**17.24** This protective tile, developed for use in the space shuttle, has extraordinary thermal properties. The extremely small thermal conductivity and small heat capacity of the material make it possible to hold the tile by its edges, even though its temperature is high enough to emit the light for this photograph.



### Problem-Solving Strategy 17.3 Heat Conduction



**IDENTIFY** *the relevant concepts:* The concept of heat conduction comes into play whenever two objects at different temperature are placed in contact.

**SET UP** *the problem* using the following steps:

1. Identify the direction of heat flow in the problem (from hot to cold). In Eq. (17.21),  $L$  is always measured along this direction, and  $A$  is always an area perpendicular to this direction. Often when a box or other container has an irregular shape but uniform wall thickness, you can approximate it as a flat slab with the same thickness and total wall area.
2. Identify the target variable.

**EXECUTE** *the solution* as follows:

1. If heat flows through a single object, use Eq. (17.21) to solve for the target variable.
2. In some problems the heat flows through two different materials in succession. The temperature at the interface between the two

materials is then intermediate between  $T_H$  and  $T_C$ ; represent it by a symbol such as  $T$ . The temperature differences for the two materials are then  $(T_H - T)$  and  $(T - T_C)$ . In steady-state heat flow, the same heat has to pass through both materials in succession, so the heat current  $H$  must be *the same* in both materials.

3. If there are two *parallel* heat-flow paths, so that some heat flows through each, then the total  $H$  is the sum of the quantities  $H_1$  and  $H_2$  for the separate paths. An example is heat flow from inside to outside a house, both through the glass in a window and through the surrounding frame. In this case the temperature difference is the same for the two paths, but  $L$ ,  $A$ , and  $k$  may be different for the two paths.
4. As always, it is essential to use a consistent set of units. If you use a value of  $k$  expressed in  $\text{W}/\text{m} \cdot \text{K}$ , don't use distances in centimeters, heat in calories, or  $T$  in degrees Fahrenheit!

**EVALUATE** *your answer:* As always, ask yourself whether the results are physically reasonable.

**Example 17.12 Conduction through a picnic cooler**

A Styrofoam box used to keep drinks cold at a picnic (Fig. 17.25a) has total wall area (including the lid) of  $0.80 \text{ m}^2$  and wall thickness  $2.0 \text{ cm}$ . It is filled with ice, water, and cans of Omni-Cola at  $0^\circ\text{C}$ . What is the rate of heat flow into the box if the temperature of the outside wall is  $30^\circ\text{C}$ ? How much ice melts in one day?

**SOLUTION**

**IDENTIFY:** The first target variable is the heat current  $H$ . The second is the amount of ice melted, which depends on the heat current (heat per unit time), the elapsed time, and the heat of fusion.

**SET UP:** We use Eq. (17.21) to describe the heat current and Eq. (17.20),  $Q = mL_f$ , to determine the mass  $m$  of ice that melts due to the heat flow.

**EXECUTE:** We assume that the total heat flow is approximately the same as it would be through a flat slab of area  $0.80 \text{ m}^2$  and thickness  $2.0 \text{ cm} = 0.020 \text{ m}$  (Fig. 17.25b). We find  $k$  from Table 17.5. From Eq. (17.21) the heat current (rate of heat flow) is

$$H = kA \frac{T_H - T_C}{L} = (0.010 \text{ W/m} \cdot \text{K})(0.80 \text{ m}^2) \frac{30^\circ\text{C} - 0^\circ\text{C}}{0.020 \text{ m}} = 12 \text{ W} = 12 \text{ J/s}$$

The total heat flow  $Q$  in one day (86,400 s) is

$$Q = Ht = (12 \text{ J/s})(86,400 \text{ s}) = 1.04 \times 10^6 \text{ J}$$

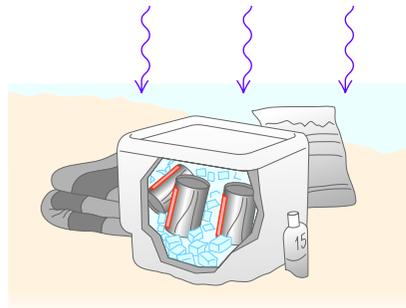
The heat of fusion of ice is  $3.34 \times 10^5 \text{ J/kg}$ , so the quantity of ice melted by this quantity of heat is

$$m = \frac{Q}{L_f} = \frac{1.04 \times 10^6 \text{ J}}{3.34 \times 10^5 \text{ J/kg}} = 3.1 \text{ kg}$$

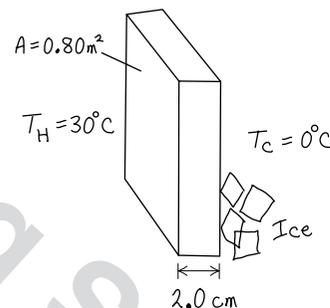
**EVALUATE:** The low heat current is a result of the low thermal conductivity of Styrofoam. A substantial amount of heat flows in 24 hours, but a relatively small amount of ice melts because the heat of fusion is high.

**17.25 Conduction of heat across the walls of a Styrofoam cooler.**

(a) A cooler at the beach



(b) Our sketch for this problem



**Example 17.13 Conduction through two bars I**

A steel bar  $10.0 \text{ cm}$  long is welded end to end to a copper bar  $20.0 \text{ cm}$  long. Both bars are insulated perfectly on their sides. Each bar has a square cross section,  $2.00 \text{ cm}$  on a side. The free end of the steel bar is maintained at  $100^\circ\text{C}$  by placing it in contact with steam, and the free end of the copper bar is maintained at  $0^\circ\text{C}$  by placing it in contact with ice. Find the temperature at the junction of the two bars and the total rate of heat flow.

**SOLUTION**

**IDENTIFY:** In this problem there is heat flow through two bars of different composition. As we discussed in Problem-Solving Strategy 17.3, the heat currents in the two end-to-end bars must be the same.

**SET UP:** Figure 17.26 shows the situation. We write Eq. (17.21) twice, once for each bar, and set the heat currents  $H_{\text{steel}}$  and  $H_{\text{copper}}$  equal to each other. Both expressions for the heat current involve the temperature  $T$  at the junction, which is one of our target variables.

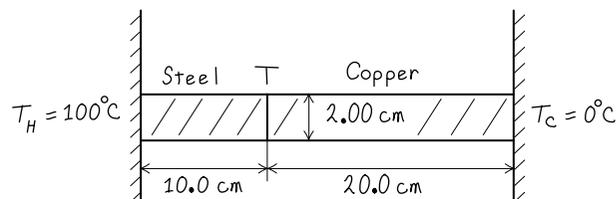
**EXECUTE:** Setting the two heat currents equal, we have

$$H_{\text{steel}} = \frac{k_{\text{steel}}A(100^\circ\text{C} - T)}{L_{\text{steel}}} = H_{\text{copper}} = \frac{k_{\text{copper}}A(T - 0^\circ\text{C})}{L_{\text{copper}}}$$

The areas  $A$  are equal and may be divided out. Substituting  $L_{\text{steel}} = 0.100 \text{ m}$ ,  $L_{\text{copper}} = 0.200 \text{ m}$ , and numerical values of  $k$  from Table 17.5, we find

$$\frac{(50.2 \text{ W/m} \cdot \text{K})(100^\circ\text{C} - T)}{0.100 \text{ m}} = \frac{(385 \text{ W/m} \cdot \text{K})(T - 0^\circ\text{C})}{0.200 \text{ m}}$$

**17.26 Our sketch for this problem.**



Rearranging and solving for  $T$ , we find

$$T = 20.7^\circ\text{C}$$

We can find the total heat current by substituting this value for  $T$  back into either of the above expressions:

$$\begin{aligned} H_{\text{steel}} &= \frac{(50.2 \text{ W/m} \cdot \text{K})(0.0200 \text{ m})^2(100^\circ\text{C} - 20.7^\circ\text{C})}{0.100 \text{ m}} \\ &= 15.9 \text{ W} \end{aligned}$$

or

$$H_{\text{copper}} = \frac{(385 \text{ W/m} \cdot \text{K})(0.0200 \text{ m})^2(20.7^\circ\text{C})}{0.200 \text{ m}} = 15.9 \text{ W}$$

**EVALUATE:** Even though the steel bar is shorter, the temperature drop across it is much greater than across the copper bar (from  $100^\circ\text{C}$  to  $20.7^\circ\text{C}$  in the steel versus from  $20.7^\circ\text{C}$  to  $0^\circ\text{C}$  in the copper). This difference arises because steel is a much poorer conductor than copper.

### Example 17.14 Conduction through two bars II

In Example 17.13, suppose the two bars are separated. One end of each bar is maintained at  $100^\circ\text{C}$  and the other end of each bar is maintained at  $0^\circ\text{C}$ . What is the *total* rate of heat flow in the two bars?

#### SOLUTION

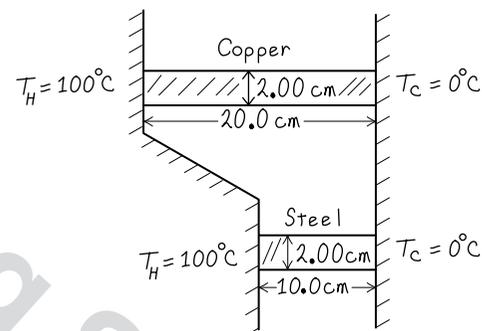
**IDENTIFY:** In this case the bars are side by side rather than end to end. The total heat current is now the *sum* of the currents in the two bars.

**SET UP:** Figure 17.27 shows the situation. For each bar,  $T_H - T_C = 100^\circ\text{C} - 0^\circ\text{C} = 100 \text{ K}$ .

**EXECUTE:** We write the heat currents for the two rods individually, and then add them to get the total heat current:

$$\begin{aligned} H &= H_{\text{steel}} + H_{\text{copper}} = \frac{k_{\text{steel}}A(T_H - T_C)}{L_{\text{steel}}} + \frac{k_{\text{copper}}A(T_H - T_C)}{L_{\text{copper}}} \\ &= \frac{(50.2 \text{ W/m} \cdot \text{K})(0.0200 \text{ m})^2(100 \text{ K})}{0.100 \text{ m}} \\ &\quad + \frac{(385 \text{ W/m} \cdot \text{K})(0.0200 \text{ m})^2(100 \text{ K})}{0.200 \text{ m}} \\ &= 20.1 \text{ W} + 77.0 \text{ W} = 97.1 \text{ W} \end{aligned}$$

**17.27** Our sketch for this problem.



## Convection

**Convection** is the transfer of heat by mass motion of a fluid from one region of space to another. Familiar examples include hot-air and hot-water home heating systems, the cooling system of an automobile engine, and the flow of blood in the body. If the fluid is circulated by a blower or pump, the process is called *forced convection*; if the flow is caused by differences in density due to thermal expansion, such as hot air rising, the process is called *natural convection* or *free convection* (Fig. 17.28).

Free convection in the atmosphere plays a dominant role in determining the daily weather, and convection in the oceans is an important global heat-transfer mechanism. On a smaller scale, soaring hawks and glider pilots make use of thermal updrafts from the warm earth. The most important mechanism for heat transfer within the human body (needed to maintain nearly constant temperature in various environments) is *forced* convection of blood, with the heart serving as the pump.

Convective heat transfer is a very complex process, and there is no simple equation to describe it. Here are a few experimental facts:

1. The heat current due to convection is directly proportional to the surface area. This is the reason for the large surface areas of radiators and cooling fins.
2. The viscosity of fluids slows natural convection near a stationary surface, giving a surface film that on a vertical surface typically has about the same insulating value as 1.3 cm of plywood ( $R$  value = 0.7). Forced convection

**17.28** A heating element in the tip of this submerged tube warms the surrounding water, producing a complex pattern of free convection.



decreases the thickness of this film, increasing the rate of heat transfer. This is the reason for the “wind-chill factor”; you get cold faster in a cold wind than in still air with the same temperature.

- The heat current due to convection is found to be approximately proportional to the  $\frac{5}{4}$  power of the temperature difference between the surface and the main body of fluid.

## Radiation

**Radiation** is the transfer of heat by electromagnetic waves such as visible light, infrared, and ultraviolet radiation. Everyone has felt the warmth of the sun’s radiation and the intense heat from a charcoal grill or the glowing coals in a fireplace. Most of the heat from these very hot bodies reaches you not by conduction or convection in the intervening air but by *radiation*. This heat transfer would occur even if there were nothing but vacuum between you and the source of heat.

Every body, even at ordinary temperatures, emits energy in the form of electromagnetic radiation. At ordinary temperatures, say  $20^\circ\text{C}$ , nearly all the energy is carried by infrared waves with wavelengths much longer than those of visible light (see Figs. 17.4 and 17.29). As the temperature rises, the wavelengths shift to shorter values. At  $800^\circ\text{C}$  a body emits enough visible radiation to be self-luminous and appears “red-hot,” although even at this temperature most of the energy is carried by infrared waves. At  $3000^\circ\text{C}$ , the temperature of an incandescent lamp filament, the radiation contains enough visible light that the body appears “white-hot.”

The rate of energy radiation from a surface is proportional to the surface area  $A$ . The rate increases very rapidly with temperature, depending on the fourth power of the absolute (Kelvin) temperature. The rate also depends on the nature of the surface; this dependence is described by a quantity  $e$  called the **emissivity**. A dimensionless number between 0 and 1, it represents the ratio of the rate of radiation from a particular surface to the rate of radiation from an equal area of an ideal radiating surface at the same temperature. Emissivity also depends somewhat on temperature. Thus the heat current  $H = dQ/dt$  due to radiation from a surface area  $A$  with emissivity  $e$  at absolute temperature  $T$  can be expressed as

$$H = Ae\sigma T^4 \quad (\text{heat current in radiation}) \quad (17.25)$$

where  $\sigma$  is a fundamental physical constant called the **Stefan–Boltzmann constant**. This relationship is called the **Stefan–Boltzmann law** in honor of its late-19th-century discoverers. The current best numerical value of  $\sigma$  is

$$\sigma = 5.670400(40) \times 10^{-8} \text{ W/m}^2 \cdot \text{K}^4$$

We invite you to check unit consistency in Eq. (17.25). Emissivity ( $e$ ) is often larger for dark surfaces than for light ones. The emissivity of a smooth copper surface is about 0.3, but  $e$  for a dull black surface can be close to unity.

**17.29** This false-color infrared photograph reveals radiation emitted by various parts of the man’s body. The strongest emission (colored red) comes from the warmest areas, while there is very little emission from the bottle of cold beverage.



### Example 17.15 Heat transfer by radiation

A thin square steel plate, 10 cm on a side, is heated in a blacksmith’s forge to a temperature of  $800^\circ\text{C}$ . If the emissivity is 0.60, what is the total rate of radiation of energy?

#### SOLUTION

**IDENTIFY:** The target variable is  $H$ , the rate of emission of energy.

**SET UP:** We use Eq. (17.25) to calculate  $H$  from the given values.

**EXECUTE:** The total surface area, including both sides, is  $2(0.10 \text{ m})^2 = 0.020 \text{ m}^2$ . We must convert the temperature to the Kelvin scale;  $800^\circ\text{C} = 1073 \text{ K}$ . Then Eq. (17.25) gives

$$\begin{aligned} H &= Ae\sigma T^4 \\ &= (0.020 \text{ m}^2)(0.60)(5.67 \times 10^{-8} \text{ W/m}^2 \cdot \text{K}^4)(1073 \text{ K})^4 \\ &= 900 \text{ W} \end{aligned}$$

**EVALUATE:** A blacksmith standing nearby will easily feel heat being radiated from this plate.

## Radiation and Absorption

While a body at absolute temperature  $T$  is radiating, its surroundings at temperature  $T_s$  are also radiating, and the body *absorbs* some of this radiation. If it is in thermal equilibrium with its surroundings,  $T = T_s$  and the rates of radiation and absorption must be equal. For this to be true, the rate of absorption must be given in general by  $H = Ae\sigma T_s^4$ . Then the *net* rate of radiation from a body at temperature  $T$  with surroundings at temperature  $T_s$  is

$$H_{\text{net}} = Ae\sigma T^4 - Ae\sigma T_s^4 = Ae\sigma(T^4 - T_s^4) \quad (17.26)$$

In this equation a positive value of  $H$  means a net heat flow *out of* the body. Equation (17.26) shows that for radiation, as for conduction and convection, the heat current depends on the temperature *difference* between two bodies.

### Example 17.16 Radiation from the human body

If the total surface area of the human body is  $1.20 \text{ m}^2$  and the surface temperature is  $30^\circ\text{C} = 303 \text{ K}$ , find the total rate of radiation of energy from the body. If the surroundings are at a temperature of  $20^\circ\text{C}$ , what is the *net* rate of heat loss from the body by radiation? The emissivity of the body is very close to unity, irrespective of skin pigmentation.

#### SOLUTION

**IDENTIFY:** We must take into account both the radiation that the body emits and the radiation that the body absorbs from its surroundings.

**SET UP:** The rate of radiation of energy from the body is given by Eq. (17.25), and the net rate of heat loss is given by Eq. (17.26).

**EXECUTE:** Taking  $e = 1$  in Eq. (17.25), we find that the body radiates at a rate

$$\begin{aligned} H &= Ae\sigma T^4 \\ &= (1.20 \text{ m}^2)(1)(5.67 \times 10^{-8} \text{ W/m}^2 \cdot \text{K}^4)(303 \text{ K})^4 \\ &= 574 \text{ W} \end{aligned}$$

This loss is partly offset by absorption of radiation, which depends on the temperature of the surroundings. The *net* rate of radiative energy transfer is given by Eq. (17.26):

$$\begin{aligned} H_{\text{net}} &= Ae\sigma(T^4 - T_s^4) \\ &= (1.20 \text{ m}^2)(1)(5.67 \times 10^{-8} \text{ W/m}^2 \cdot \text{K}^4) \\ &\quad \times [(303 \text{ K})^4 - (293 \text{ K})^4] = 72 \text{ W} \end{aligned}$$

**EVALUATE:** The value of  $H_{\text{net}}$  is positive because the body is losing heat to its colder surroundings.

## Applications of Radiation

Heat transfer by radiation is important in some surprising places. A premature baby in an incubator can be cooled dangerously by radiation if the walls of the incubator happen to be cold, even when the *air* in the incubator is warm. Some incubators regulate the air temperature by measuring the baby's skin temperature.

A body that is a good absorber must also be a good emitter. An ideal radiator, with an emissivity of unity, is also an ideal absorber, absorbing *all* of the radiation that strikes it. Such an ideal surface is called an ideal black body or simply a **blackbody**. Conversely, an ideal *reflector*, which absorbs *no* radiation at all, is also a very ineffective radiator.

This is the reason for the silver coatings on vacuum ("Thermos") bottles, invented by Sir James Dewar (1842–1923). A vacuum bottle has double glass walls. The air is pumped out of the spaces between the walls; this eliminates nearly all heat transfer by conduction and convection. The silver coating on the walls reflects most of the radiation from the contents back into the container, and the wall itself is a very poor emitter. Thus a vacuum bottle can keep coffee or soup hot for several hours. The Dewar flask, used to store very cold liquefied gases, is exactly the same in principle.

**Test Your Understanding of Section 17.7** A room has one wall made of concrete, one wall made of copper, and one wall made of steel. All of the walls are the same size and at the same temperature of  $20^\circ\text{C}$ . Which wall feels coldest to the touch? (i) the concrete wall; (ii) the copper wall; (iii) the steel wall; (iv) all three walls feel equally cold to the touch.

# CHAPTER 17 SUMMARY

**Temperature and temperature scales:** A thermometer measures temperature. Two bodies in thermal equilibrium must have the same temperature. A conducting material between two bodies permits them to interact and come to thermal equilibrium; an insulating material impedes this interaction.

The Celsius and Fahrenheit temperature scales are based on the freezing temperature ( $0^{\circ}\text{C} = 32^{\circ}\text{F}$ ) and boiling temperature ( $100^{\circ}\text{C} = 212^{\circ}\text{F}$ ). One Celsius degree equals  $\frac{9}{5}$  Fahrenheit degrees. (See Example 17.1.)

The Kelvin scale has its zero at the extrapolated zero-pressure temperature for a gas thermometer,  $-273.15^{\circ}\text{C} = 0\text{ K}$ . In the gas-thermometer scale, the ratio of two temperatures  $T_1$  and  $T_2$  is defined to be equal to the ratio of the two corresponding gas-thermometer pressures  $p_1$  and  $p_2$ . The triple-point temperature of water ( $0.01^{\circ}\text{C}$ ) is defined to be 273.16 K.

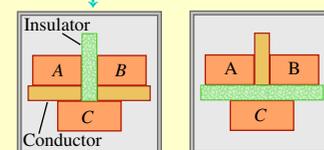
$$T_{\text{F}} = \frac{9}{5}T_{\text{C}} + 32^{\circ} \quad (17.1)$$

$$T_{\text{C}} = \frac{5}{9}(T_{\text{F}} - 32^{\circ}) \quad (17.2)$$

$$T_{\text{K}} = T_{\text{C}} + 273.15 \quad (17.3)$$

$$\frac{T_2}{T_1} = \frac{p_2}{p_1} \quad (17.4)$$

If systems A and B are each in thermal equilibrium with system C ...



... then systems A and B are in thermal equilibrium with each other.

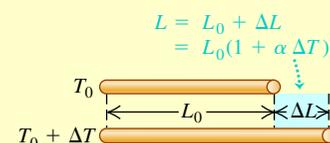
**Thermal expansion and thermal stress:** A temperature change  $\Delta T$  causes a change in any linear dimension  $L_0$  of a solid body. The change  $\Delta L$  is approximately proportional to  $L_0$  and  $\Delta T$ . Similarly, a temperature change causes a change  $\Delta V$  in the volume  $V_0$  of any solid or liquid material that is approximately proportional to  $V_0$  and  $\Delta T$ . The quantities  $\alpha$  and  $\beta$  are the coefficients of linear expansion and volume expansion, respectively. For solids,  $\beta = 3\alpha$ . (See Examples 17.2–17.4.)

When a material is cooled or heated and held so it cannot contract or expand, it is under a tensile stress  $F/A$ . (See Example 17.5.)

$$\Delta L = \alpha L_0 \Delta T \quad (17.6)$$

$$\Delta V = \beta V_0 \Delta T \quad (17.8)$$

$$\frac{F}{A} = -Y\alpha \Delta T \quad (17.12)$$



**Heat, phase changes, and calorimetry:** Heat is energy in transit from one body to another as a result of a temperature difference. The quantity of heat  $Q$  required to raise the temperature of a quantity of material by a small amount  $\Delta T$  is proportional to  $\Delta T$ . This proportionality can be expressed either in terms of the mass  $m$  and specific heat capacity  $c$  or in terms of the number of moles  $n$  and the molar heat capacity  $C = Mc$ . Here  $M$  is the molar mass and  $m = nM$ . (See Examples 17.6 and 17.7.)

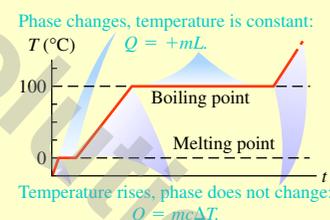
To change a mass  $m$  of a material to a different phase at the same temperature (such as liquid to solid or liquid to vapor) requires the addition or subtraction of a quantity of heat. The amount of heat is equal to the product of  $m$  and  $L$ , the heat of fusion, vaporization, or sublimation.

When heat is added to a body, the corresponding  $Q$  is positive; when it is removed,  $Q$  is negative. The basic principle of calorimetry comes from conservation of energy. In an isolated system whose parts interact by heat exchange, the algebraic sum of the  $Q$ 's for all parts of the system must be zero. (See Examples 17.8–17.11.)

$$Q = mc \Delta T \quad (17.13)$$

$$Q = nC \Delta T \quad (17.18)$$

$$Q = \pm mL \quad (17.20)$$



**Conduction, convection, and radiation:** Conduction is the transfer of energy of molecular motion within bulk materials without bulk motion of the materials. The heat current  $H$  or conduction depends on the area  $A$  through which the heat flows, the length  $L$  of the heat-flow path, the temperature difference ( $T_H - T_C$ ), and the thermal conductivity  $k$  of the material. (See Examples 17.12–17.14.)

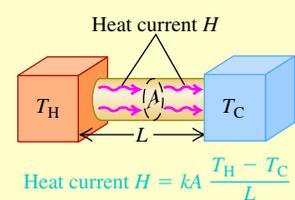
Convection is a complex heat-transfer process that involves mass motion from one region to another. It depends on surface area, orientation, and the temperature difference between a body and its surroundings.

Radiation is energy transfer through electromagnetic radiation. The radiation heat current  $H$  depends on the surface area  $A$ , the emissivity  $e$  of the surface (a pure number between 0 and 1), and the Kelvin temperature  $T$ . It involves a fundamental constant  $\sigma$  called the Stefan–Boltzmann constant. When a body at temperature  $T$  is surrounded by material at temperature  $T_s$ , the *net* heat current  $H_{\text{net}}$  from the body to its surroundings depends on both  $T$  and  $T_s$ . (See Examples 17.15 and 17.16.)

$$H = \frac{dQ}{dt} = kA \frac{T_H - T_C}{L} \quad (17.21)$$

$$H = Ae\sigma T^4 \quad (17.25)$$

$$H_{\text{net}} = Ae\sigma(T^4 - T_s^4) \quad (17.26)$$



## Key Terms

thermodynamics, 570

temperature, 571

thermometer, 571

thermal equilibrium, 571

insulator, 571

conductor, 571

zeroth law of thermodynamics, 572

Celsius temperature scale, 572

Fahrenheit temperature scale, 573

Kelvin temperature scale, 574

absolute temperature scale, 576

absolute zero, 576

coefficient of linear expansion, 576

coefficient of volume expansion, 578

thermal stress, 580

heat, 582

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convection, 595

radiation, 596

emissivity, 596

Stefan–Boltzmann constant, 596

Stefan–Boltzmann law, 596

blackbody, 597

## Answer to Chapter Opening Question

No. By “heat” we mean energy that is in transit from one body to another as a result of temperature difference between the bodies. Bodies do not *contain* heat.

## Answers to Test Your Understanding Questions

**17.1 Answer: (ii)** A liquid-in-tube thermometer actually measures its own temperature. If the thermometer stays in the hot water long enough, it will come to thermal equilibrium with the water and its temperature will be the same as that of the water.

**17.2 Answer: (iv)** Both a bimetallic strip and a resistance thermometer measure their own temperature. For this to be equal to the temperature of the object being measured, the thermometer and object must be in contact and in thermal equilibrium. A temporal artery thermometer detects the infrared radiation from a person’s skin, so there is no need for the detector and skin to be at the same temperature.

?

**17.3 Answer: (i), (iii), (ii), (v), (iv)** To compare these temperatures, convert them all to the Kelvin scale. For (i), the Kelvin temperature is  $T_K = T_C + 273.15 = 0.00 + 273.15 = 273.15$  K; for (ii),  $T_C = \frac{5}{9}(T_F - 32^\circ) = \frac{5}{9}(0.00^\circ - 32^\circ) = -17.78^\circ\text{C}$  and  $T_K = T_C + 273.15 = -17.78 + 273.15 = 255.37$  K; for (iii),  $T_K = 260.00$  K; for (iv),  $T_K = 77.00$  K; and for (v),  $T_K = T_C + 273.15 = -180.00 + 273.15 = 93.15$  K.

**17.4 Answer: (ii) and (iii)** Metal 2 must expand more than metal 1 when heated and so must have a larger coefficient of linear expansion  $\alpha$ . From Table 17.1, brass and aluminum have larger values of  $\alpha$  than copper, but steel does not.

**17.5 Answer: (ii), (i), (iv), (iii)** For (i) and (ii), the relevant quantity is the specific heat  $c$  of the substance, which is the amount of heat required to raise the temperature of 1 *kilogram* of that substance by 1 K (1  $^\circ\text{C}$ ). From Table 17.3, these values are (i) 138 J for mercury and (ii) 2428 J for ethanol. For (iii) and (iv) we need the molar heat capacity  $C$ , which is the amount of heat required to raise the temperature of 1 *mole* of that substance by 1  $^\circ\text{C}$ . Again from Table 17.3, these values are (iii) 27.7 J for mercury and (iv) 111.9 J

for ethanol. (The ratio of molar heat capacities is different from the ratio of the specific heats because a mole of mercury and a mole of ethanol have different masses.)

**17.6 Answer: (iv)** In time  $t$  the system goes from point  $b$  to point  $e$  in Fig. 17.21. According to this figure, at time  $t/2$  (halfway along the horizontal axis from  $b$  to  $e$ ), the system is at  $100^\circ\text{C}$  and is still boiling; that is, it is a mixture of liquid and gas. This says that most of the heat added goes into boiling the water.

**17.7 Answer: (ii)** When you touch one of the walls, heat flows from your hand to the lower-temperature wall. The more rapidly heat flows from your hand, the colder you will feel. Equation (17.21) shows that the rate of heat flow is proportional to the thermal conductivity  $k$ . From Table 17.5, copper has a much higher thermal conductivity ( $385.0\text{ W/m}\cdot\text{K}$ ) than steel ( $50.2\text{ W/m}\cdot\text{K}$ ) or concrete ( $0.8\text{ W/m}\cdot\text{K}$ ), and so the copper wall feels the coldest to the touch.

## PROBLEMS

For instructor-assigned homework, go to [www.masteringphysics.com](http://www.masteringphysics.com)



### Discussion Questions

**Q17.1.** Explain why it would not make sense to use a full-size glass thermometer to measure the temperature of a thimbleful of hot water.

**Q17.2.** If you heat the air inside a rigid, sealed container until its Kelvin temperature doubles, the air pressure in the container will also double. Is the same thing true if you double the Celsius temperature of the air in the container? Explain.

**Q17.3.** Many automobile engines have cast-iron cylinders and aluminum pistons. What kinds of problems could occur if the engine gets too hot? (The coefficient of volume expansion of cast iron is approximately the same as that of steel.)

**Q17.4.** Why do frozen water pipes burst? Would a mercury thermometer break if the temperature went below the freezing temperature of mercury? Why or why not?

**Q17.5.** Two bodies made of the same material have the same external dimensions and appearance, but one is solid and the other is hollow. When their temperature is increased, is the overall volume expansion the same or different? Why?

**Q17.6.** The inside of an oven is at a temperature of  $200^\circ\text{C}$  ( $392^\circ\text{F}$ ). You can put your hand in the oven without injury as long as you don't touch anything. But since the air inside the oven is also at  $200^\circ\text{C}$ , why isn't your hand burned just the same?

**Q17.7.** A newspaper article about the weather states that "the temperature of a body measures how much heat the body contains." Is this description correct? Why or why not?

**Q17.8.** To raise the temperature of an object, must you add heat to it? If you add heat to an object, must you raise its temperature? Explain.

**Q17.9.** A student asserts that a suitable unit for specific heat capacity is  $1\text{ m}^2/\text{s}^2\cdot\text{C}^\circ$ . Is she correct? Why or why not?

**Q17.10.** In some household air conditioners used in dry climates, air is cooled by blowing it through a water-soaked filter, evaporating some of the water. How does this cool the air? Would such a system work well in a high-humidity climate? Why or why not?

**Q17.11.** The units of specific heat capacity  $c$  are  $\text{J}/\text{kg}\cdot\text{K}$ , but the units of heat of fusion  $L_f$  or heat of vaporization  $L_v$  are simply  $\text{J}/\text{kg}$ . Why do the units of  $L_f$  and  $L_v$  not include a factor of  $(\text{K})^{-1}$  to account for a temperature change?

**Q17.12.** Why is a hot, humid day in the tropics generally more uncomfortable for human beings than a hot, dry day in the desert?

**Q17.13.** A piece of aluminum foil used to wrap a potato for baking in a hot oven can usually be handled safely within a few seconds after the potato is removed from the oven. The same is not true of the potato, however! Give two reasons for this difference.

**Q17.14.** Desert travelers sometimes keep water in a canvas bag. Some water seeps through the bag and evaporates. How does this cool the water inside the bag?

**Q17.15.** When you first step out of the shower, you feel cold. But as soon as you are dry you feel warmer, even though the room temperature does not change. Why?

**Q17.16.** The climate of regions adjacent to large bodies of water (like the Pacific and Atlantic coasts) usually features a narrower range of temperature than the climate of regions far from large bodies of water (like the prairies). Why?

**Q17.17.** When water is placed in ice-cube trays in a freezer, why doesn't the water freeze all at once when the temperature has reached  $0^\circ\text{C}$ ? In fact, the water freezes first in a layer adjacent to the sides of the tray. Why?

**Q17.18.** Before giving you an injection, a physician swabs your arm with isopropyl alcohol at room temperature. Why does this make your arm feel cold? (*Hint:* The reason is *not* the fear of the injection! The boiling point of isopropyl alcohol is  $82.4^\circ\text{C}$ .)

**Q17.19.** A cold block of metal feels colder than a block of wood at the same temperature. Why? A hot block of metal feels hotter than a block of wood at the same temperature. Again, why? Is there any temperature at which the two blocks feel equally hot or cold? What temperature is this?

**Q17.20.** A person pours a cup of hot coffee, intending to drink it five minutes later. To keep the coffee as hot as possible, should she put cream in it now, or wait until just before she drinks it? Explain.

**Q17.21.** When a freshly baked apple pie has just been removed from the oven, the crust and filling are both at the same temperature. Yet if you sample the pie, the filling will burn your tongue but the crust will not. Why is there a difference? (*Hint:* The filling is moist while the crust is dry.)

**Q17.22.** Old-time kitchen lore suggests that things cook better (evenly and without burning) in heavy cast-iron pots. What desirable characteristics do such pots have?

**Q17.23.** In coastal regions in the winter, the temperature over the land is generally colder than the temperature over the nearby ocean; in the summer, the reverse is usually true. Explain. (*Hint:* The specific heat of soil is only 0.2–0.8 times as great as that of water.)

**Q17.24.** It is well known that a potato bakes faster if a large nail is stuck through it. Why? Does an aluminum nail work better than a steel one? Why or why not? (*Note:* Don't try this in a microwave oven!) There is also a gadget on the market to hasten the roasting of meat; it consists of a hollow metal tube containing a wick and some water. This is claimed to work much better than a solid metal rod. How does it work?

**Q17.25.** Glider pilots in the Midwest know that thermal updrafts are likely to occur in the vicinity of freshly plowed fields. Why?

**Q17.26.** Some folks claim that ice cubes freeze faster if the trays are filled with hot water, because hot water cools off faster than cold water. What do you think?

**Q17.27.** We're lucky that the earth isn't in thermal equilibrium with the sun (which has a surface temperature of 5800 K). But why aren't the two bodies in thermal equilibrium?

**Q17.28.** When energy shortages occur, magazine articles sometimes urge us to keep our homes at a constant temperature day and night to conserve fuel. They argue that when we turn down the heat at night, the walls, ceilings, and other areas cool off and must be reheated in the morning. So if we keep the temperature constant, these parts of the house will not cool off and will not have to be reheated. Does this argument make sense? Would we really save energy by following this advice?

## Exercises

### Section 17.2 Thermometers and Temperature Scales

**17.1.** Convert the following Celsius temperatures to Fahrenheit: (a)  $-62.8^{\circ}\text{C}$ , the lowest temperature ever recorded in North America (February 3, 1947, Snag, Yukon); (b)  $56.7^{\circ}\text{C}$ , the highest temperature ever recorded in the United States (July 10, 1913, Death Valley, California); (c)  $31.1^{\circ}\text{C}$ , the world's highest average annual temperature (Lugh Ferrandi, Somalia).

**17.2.** Find the Celsius temperatures corresponding to (a) a winter night in Seattle ( $41.0^{\circ}\text{F}$ ); (b) a hot summer day in Palm Springs ( $107.0^{\circ}\text{F}$ ); (c) a cold winter day in northern Manitoba ( $-18.0^{\circ}\text{F}$ ).

**17.3.** While vacationing in Italy, you see on local TV one summer morning that the temperature will rise from the current  $18^{\circ}\text{C}$  to a high of  $39^{\circ}\text{C}$ . What is the corresponding increase in the Fahrenheit temperature?

**17.4.** Two beakers of water, *A* and *B*, initially are at the same temperature. The temperature of the water in beaker *A* is increased  $10^{\circ}\text{F}$ , and the temperature of the water in beaker *B* is increased  $10\text{ K}$ . After these temperature changes, which beaker of water has the higher temperature? Explain.

**17.5.** You put a bottle of soft drink in a refrigerator and leave it until its temperature has dropped  $10.0\text{ K}$ . What is its temperature change in (a)  $^{\circ}\text{F}$  and (b)  $^{\circ}\text{C}$ ?

**17.6.** (a) On January 22, 1943, the temperature in Spearfish, South Dakota, rose from  $-4.0^{\circ}\text{F}$  to  $45.0^{\circ}\text{F}$  in just 2 minutes. What was the temperature change in Celsius degrees? (b) The temperature in Browning, Montana, was  $44.0^{\circ}\text{F}$  on January 23, 1916. The next day the temperature plummeted to  $-56^{\circ}\text{C}$ . What was the temperature change in Celsius degrees?

**17.7.** (a) You feel sick and are told that you have a temperature of  $40.2^{\circ}\text{C}$ . What is your temperature in  $^{\circ}\text{F}$ ? Should you be concerned? (b) The morning weather report in Sydney gives a current temperature of  $12^{\circ}\text{C}$ . What is this temperature in  $^{\circ}\text{F}$ ?

### Section 17.3 Gas Thermometers and the Kelvin Scale

**17.8.** (a) Calculate the one temperature at which Fahrenheit and Celsius thermometers agree with each other. (b) Calculate the one temperature at which Fahrenheit and Kelvin thermometers agree with each other.

**17.9.** Convert the following record-setting temperatures to the Kelvin scale: (a) the lowest temperature recorded in the 48 contiguous states ( $-70.0^{\circ}\text{F}$  at Rogers Pass, Montana, on January 20, 1954); (b) Australia's highest temperature ( $127.0^{\circ}\text{F}$  at Cloncurry, Queensland, on January 16, 1889); (c) the lowest temperature recorded in the northern hemisphere ( $-90.0^{\circ}\text{F}$  at Verkhoyansk, Siberia, in 1892).

**17.10.** Convert the following Kelvin temperatures to the Celsius and Fahrenheit scales: (a) the midday temperature at the surface of

the moon ( $400\text{ K}$ ); (b) the temperature at the tops of the clouds in the atmosphere of Saturn ( $95\text{ K}$ ); (c) the temperature at the center of the sun ( $1.55 \times 10^7\text{ K}$ ).

**17.11.** Liquid nitrogen is a relatively inexpensive material that is often used to perform entertaining low-temperature physics demonstrations. Nitrogen gas liquefies at a temperature of  $-346^{\circ}\text{F}$ . Convert this temperature to (a)  $^{\circ}\text{C}$  and (b)  $\text{K}$ .

**17.12.** A gas thermometer registers an absolute pressure corresponding to 325 mm of mercury when in contact with water at the triple point. What pressure does it read when in contact with water at the normal boiling point?

**17.13.** The pressure of a gas at the triple point of water is 1.35 atm. If its volume remains unchanged, what will its pressure be at the temperature at which  $\text{CO}_2$  solidifies?

**17.14.** Like the Kelvin scale, the *Rankine scale* is an absolute temperature scale: Absolute zero is zero degrees Rankine ( $0^{\circ}\text{R}$ ). However, the units of this scale are the same size as those of the Fahrenheit scale rather than the Celsius scale. What is the numerical value of the triple-point temperature of water on the Rankine scale?

**17.15. A Constant-Volume Gas Thermometer.** An experimenter using a gas thermometer found the pressure at the triple point of water ( $0.01^{\circ}\text{C}$ ) to be  $4.80 \times 10^4\text{ Pa}$  and the pressure at the normal boiling point ( $100^{\circ}\text{C}$ ) to be  $6.50 \times 10^4\text{ Pa}$ . (a) Assuming that the pressure varies linearly with temperature, use these two data points to find the Celsius temperature at which the gas pressure would be zero (that is, find the Celsius temperature of absolute zero). (b) Does the gas in this thermometer obey Eq. (17.4) precisely? If that equation were precisely obeyed and the pressure at  $100^{\circ}\text{C}$  were  $6.50 \times 10^4\text{ Pa}$ , what pressure would the experimenter have measured at  $0.01^{\circ}\text{C}$ ? (As we will learn in Section 18.1, Eq. (17.4) is accurate only for gases at very low density.)

### Section 17.4 Thermal Expansion

**17.16.** The tallest building in the world, according to some architectural standards, is the Taipei 101 in Taiwan, at a height of 1671 feet. Assume that this height was measured on a cool spring day when the temperature was  $15.5^{\circ}\text{C}$ . You could use the building as a sort of giant thermometer on a hot summer day by carefully measuring its height. Suppose you do this and discover that the Taipei 101 is 0.471 foot taller than its official height. What is the temperature, assuming that the building is in thermal equilibrium with the air and that its entire frame is made of steel?

**17.17.** The Humber Bridge in England has the world's longest single span, 1410 m. Calculate the change in length of the steel deck of the span when the temperature increases from  $-5.0^{\circ}\text{C}$  to  $18.0^{\circ}\text{C}$ .

**17.18. Ensuring a Tight Fit.** Aluminum rivets used in airplane construction are made slightly larger than the rivet holes and cooled by "dry ice" (solid  $\text{CO}_2$ ) before being driven. If the diameter of a hole is 4.500 mm, what should be the diameter of a rivet at  $23.0^{\circ}\text{C}$ , if its diameter is to equal that of the hole when the rivet is cooled to  $-78.0^{\circ}\text{C}$ , the temperature of dry ice? Assume that the expansion coefficient remains constant at the value given in Table 17.1.

**17.19.** A U.S. penny has a diameter of 1.9000 cm at  $20.0^{\circ}\text{C}$ . The coin is made of a metal alloy (mostly zinc) for which the coefficient of linear expansion is  $2.6 \times 10^{-5}\text{ K}^{-1}$ . What would its diameter be on a hot day in Death Valley ( $48.0^{\circ}\text{C}$ )? On a cold night in the mountains of Greenland ( $-53^{\circ}\text{C}$ )?

**17.20.** A geodesic dome constructed with an aluminum framework is a nearly perfect hemisphere; its diameter measures 55.0 m on a winter day at a temperature of  $-15^{\circ}\text{C}$ . How much more interior

space does the dome have in the summer, when the temperature is  $35^{\circ}\text{C}$ ?

**17.21.** A metal rod is 40.125 cm long at  $20.0^{\circ}\text{C}$  and 40.148 cm long at  $45.0^{\circ}\text{C}$ . Calculate the average coefficient of linear expansion of the rod for this temperature range.

**17.22.** A copper cylinder is initially at  $20.0^{\circ}\text{C}$ . At what temperature will its volume be 0.150% larger than it is at  $20.0^{\circ}\text{C}$ ?

**17.23.** The density of water is  $999.73\text{ kg/m}^3$  at a temperature of  $10^{\circ}\text{C}$  and  $958.38\text{ kg/m}^3$  at a temperature of  $100^{\circ}\text{C}$ . Calculate the average coefficient of volume expansion for water in that range of temperature.

**17.24.** A steel tank is completely filled with  $2.80\text{ m}^3$  of ethanol when both the tank and the ethanol are at a temperature of  $32.0^{\circ}\text{C}$ . When the tank and its contents have cooled to  $18.0^{\circ}\text{C}$ , what additional volume of ethanol can be put into the tank?

**17.25.** A glass flask whose volume is  $1000.00\text{ cm}^3$  at  $0.0^{\circ}\text{C}$  is completely filled with mercury at this temperature. When flask and mercury are warmed to  $55.0^{\circ}\text{C}$ ,  $8.95\text{ cm}^3$  of mercury overflow. If the coefficient of volume expansion of mercury is  $18.0 \times 10^{-5}\text{ K}^{-1}$ , compute the coefficient of volume expansion of the glass.

**17.26.** (a) If an area measured on the surface of a solid body is  $A_0$  at some initial temperature and then changes by  $\Delta A$  when the temperature changes by  $\Delta T$ , show that

$$\Delta A = (2\alpha) A_0 \Delta T$$

where  $\alpha$  is the coefficient of linear expansion. (b) A circular sheet of aluminum is 55.0 cm in diameter at  $15.0^{\circ}\text{C}$ . By how much does the area of one side of the sheet change when the temperature increases to  $27.5^{\circ}\text{C}$ ?

**17.27.** A machinist bores a hole of diameter 1.35 cm in a steel plate at a temperature of  $25.0^{\circ}\text{C}$ . What is the cross-sectional area of the hole (a) at  $25.0^{\circ}\text{C}$  and (b) when the temperature of the plate is increased to  $175^{\circ}\text{C}$ ? Assume that the coefficient of linear expansion remains constant over this temperature range. (*Hint:* See Exercise 17.26.)

**17.28.** As a new mechanical engineer for Engines Inc., you have been assigned to design brass pistons to slide inside steel cylinders. The engines in which these pistons will be used will operate between  $20.0^{\circ}\text{C}$  and  $150.0^{\circ}\text{C}$ . Assume that the coefficients of expansion are constant over this temperature range. (a) If the piston just fits inside the chamber at  $20.0^{\circ}\text{C}$ , will the engines be able to run at higher temperatures? Explain. (b) If the cylindrical pistons are 25.000 cm in diameter at  $20.0^{\circ}\text{C}$ , what should be the minimum diameter of the cylinders at that temperature so the pistons will operate at  $150.0^{\circ}\text{C}$ ?

**17.29.** The outer diameter of a glass jar and the inner diameter of its iron lid are both 725 mm at room temperature ( $20.0^{\circ}\text{C}$ ). What will be the size of the mismatch between the lid and the jar if the lid is briefly held under hot water until its temperature rises to  $50.0^{\circ}\text{C}$ , without changing the temperature of the glass?

**17.30.** A brass rod is 185 cm long and 1.60 cm in diameter. What force must be applied to each end of the rod to prevent it from contracting when it is cooled from  $120.0^{\circ}\text{C}$  to  $10.0^{\circ}\text{C}$ ?

**17.31.** (a) A wire that is 1.50 m long at  $20.0^{\circ}\text{C}$  is found to increase in length by 1.90 cm when warmed to  $420.0^{\circ}\text{C}$ . Compute its average coefficient of linear expansion for this temperature range. (b) The wire is stretched just taut (zero tension) at  $420.0^{\circ}\text{C}$ . Find the stress in the wire if it is cooled to  $20.0^{\circ}\text{C}$  without being allowed to contract. Young's modulus for the wire is  $2.0 \times 10^{11}\text{ Pa}$ .

**17.32.** Steel train rails are laid in 12.0-m-long segments placed end to end. The rails are laid on a winter day when their temperature is

$-2.0^{\circ}\text{C}$ . (a) How much space must be left between adjacent rails if they are just to touch on a summer day when their temperature is  $33.0^{\circ}\text{C}$ ? (b) If the rails are originally laid in contact, what is the stress in them on a summer day when their temperature is  $33.0^{\circ}\text{C}$ ?

### Section 17.5 Quantity of Heat

**17.33.** An aluminum tea kettle with mass 1.50 kg and containing 1.80 kg of water is placed on a stove. If no heat is lost to the surroundings, how much heat must be added to raise the temperature from  $20.0^{\circ}\text{C}$  to  $85.0^{\circ}\text{C}$ ?

**17.34.** In an effort to stay awake for an all-night study session, a student makes a cup of coffee by first placing a 200-W electric immersion heater in 0.320 kg of water. (a) How much heat must be added to the water to raise its temperature from  $20.0^{\circ}\text{C}$  to  $80.0^{\circ}\text{C}$ ? (b) How much time is required? Assume that all of the heater's power goes into heating the water.

**17.35.** You are given a sample of metal and asked to determine its specific heat. You weigh the sample and find that its weight is 28.4 N. You carefully add  $1.25 \times 10^4\text{ J}$  of heat energy to the sample and find that its temperature rises  $18.0^{\circ}\text{C}$ . What is the sample's specific heat?

**17.36. Heat Loss During Breathing.** In very cold weather a significant mechanism for heat loss by the human body is energy expended in warming the air taken into the lungs with each breath. (a) On a cold winter day when the temperature is  $-20^{\circ}\text{C}$ , what amount of heat is needed to warm to body temperature ( $37^{\circ}\text{C}$ ) the 0.50 L of air exchanged with each breath? Assume that the specific heat of air is  $1020\text{ J/kg}\cdot\text{K}$  and that 1.0 L of air has mass  $1.3 \times 10^{-3}\text{ kg}$ . (b) How much heat is lost per hour if the respiration rate is 20 breaths per minute?

**17.37.** While running, a 70-kg student generates thermal energy at a rate of 1200 W. To maintain a constant body temperature of  $37^{\circ}\text{C}$ , this energy must be removed by perspiration or other mechanisms. If these mechanisms failed and the heat could not flow out of the student's body, for what amount of time could a student run before irreversible body damage occurred? (*Note:* Protein structures in the body are irreversibly damaged if body temperature rises to  $44^{\circ}\text{C}$  or higher. The specific heat of a typical human body is  $3480\text{ J/kg}\cdot\text{K}$ , slightly less than that of water. The difference is due to the presence of protein, fat, and minerals, which have lower specific heats.)

**17.38.** While painting the top of an antenna 225 m in height, a worker accidentally lets a 1.00-L water bottle fall from his lunchbox. The bottle lands in some bushes at ground level and does not break. If a quantity of heat equal to the magnitude of the change in mechanical energy of the water goes into the water, what is its increase in temperature?

**17.39.** A crate of fruit with mass 35.0 kg and specific heat  $3650\text{ J/kg}\cdot\text{K}$  slides down a ramp inclined at  $36.9^{\circ}\text{C}$  below the horizontal. The ramp is 8.00 m long. (a) If the crate was at rest at the top of the incline and has a speed of 2.50 m/s at the bottom, how much work was done on the crate by friction? (b) If an amount of heat equal to the magnitude of the work done by friction goes into the crate of fruit and the fruit reaches a uniform final temperature, what is its temperature change?

**17.40.** A 25,000-kg subway train initially traveling at 15.5 m/s slows to a stop in a station and then stays there long enough for its brakes to cool. The station's dimensions are 65.0 m long by 20.0 m wide by 12.0 m high. Assuming all the work done by the brakes in stopping the train is transferred as heat uniformly to all the air in the station, by how much does the air temperature in the station

rise? Take the density of the air to be  $1.20 \text{ kg/m}^3$  and its specific heat to be  $1020 \text{ J/kg} \cdot \text{K}$ .

**17.41.** A nail driven into a board increases in temperature. If we assume that 60% of the kinetic energy delivered by a 1.80-kg hammer with a speed of  $7.80 \text{ m/s}$  is transformed into heat that flows into the nail and does not flow out, what is the temperature increase of an 8.00-g aluminum nail after it is struck ten times?

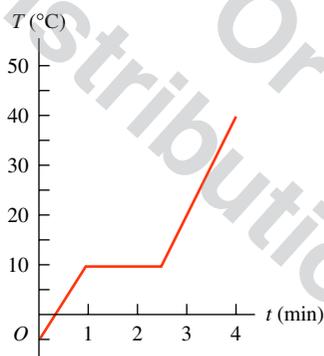
**17.42.** A technician measures the specific heat of an unidentified liquid by immersing an electrical resistor in it. Electrical energy is converted to heat transferred to the liquid for 120 s at a constant rate of 65.0 W. The mass of the liquid is 0.780 kg, and its temperature increases from  $18.55^\circ\text{C}$  to  $22.54^\circ\text{C}$ . (a) Find the average specific heat of the liquid in this temperature range. Assume that negligible heat is transferred to the container that holds the liquid and that no heat is lost to the surroundings. (b) Suppose that in this experiment heat transfer from the liquid to the container or surroundings cannot be ignored. Is the result calculated in part (a) an *overestimate* or an *underestimate* of the average specific heat? Explain.

**17.43.** You add 8950 J of heat to 3.00 mol of iron. (a) What is the temperature increase of the iron? (b) If this same amount of heat is added to 3.00 kg of iron, what is the iron's temperature increase? (c) Explain the difference in your results for parts (a) and (b).

### Section 17.6 Calorimetry and Phase Changes

**17.44.** As a physicist, you put heat into a 500.0-g solid sample at the rate of  $10.0 \text{ kJ/min}$ , while recording its temperature as a function of time. You plot your data and obtain the graph shown in Fig. 17.30. (a) What is the latent heat of fusion for this solid? (b) What are the specific heats of the liquid and solid states of the material?

**Figure 17.30** Exercise 17.44.



**17.45.** A 500.0-g chunk of an unknown metal, which has been in boiling water for several minutes, is quickly dropped into an insulating Styrofoam beaker containing 1.00 kg of water at room temperature ( $20.0^\circ\text{C}$ ). After waiting and gently stirring for 5.00 minutes, you observe that the water's temperature has reached a constant value of  $22.0^\circ\text{C}$ . (a) Assuming that the Styrofoam absorbs a negligibly small amount of heat and that no heat was lost to the surroundings, what is the specific heat of the metal? (b) Which is more useful for storing thermal energy: this metal or an equal weight of water? Explain. (c) What if the heat absorbed by the Styrofoam actually is not negligible. How would the specific heat you calculated in part (a) be in error? Would it be too large, too small, or still correct? Explain.

**17.46.** Before going in for his annual physical, a 70.0-kg man whose body temperature is  $37.0^\circ\text{C}$  consumes an entire 0.355-L can of a soft drink (mostly water) at  $12.0^\circ\text{C}$ . (a) What will his body temperature be after equilibrium is attained? Ignore any heating by the man's metabolism. The specific heat of the man's body is  $3480 \text{ J/kg} \cdot \text{K}$ . (b) Is the change in his body temperature great enough to be measured by a medical thermometer?

**17.47.** In the situation described in Exercise 17.46, the man's metabolism will eventually return the temperature of his body (and of the soft drink that he consumed) to  $37.0^\circ\text{C}$ . If his body releases energy at a rate of  $7.00 \times 10^3 \text{ kJ/day}$  (the *basal metabolic rate*, or BMR), how long does this take? Assume that all of the released energy goes into raising the temperature.

**17.48.** An ice-cube tray of negligible mass contains 0.350 kg of water at  $18.0^\circ\text{C}$ . How much heat must be removed to cool the water to  $0.00^\circ\text{C}$  and freeze it? Express your answer in joules, calories, and Btu.

**17.49.** How much heat is required to convert 12.0 g of ice at  $-10.0^\circ\text{C}$  to steam at  $100.0^\circ\text{C}$ ? Express your answer in joules, calories, and Btu.

**17.50.** An open container holds 0.550 kg of ice at  $-15.0^\circ\text{C}$ . The mass of the container can be ignored. Heat is supplied to the container at the constant rate of  $800.0 \text{ J/min}$  for 500.0 min. (a) After how many minutes does the ice *start* to melt? (b) After how many minutes, from the time when the heating is first started, does the temperature begin to rise above  $0.0^\circ\text{C}$ ? (c) Plot a curve showing the temperature as a function of the elapsed time.

**17.51.** The capacity of commercial air conditioners is sometimes expressed in "tons," the number of tons of ice (1 ton = 2000 lb) that can be frozen from water at  $0^\circ\text{C}$  in 24 h by the unit. Express the capacity of a 2-ton air conditioner in Btu/h and in watts.

**17.52. Steam Burns Versus Water Burns.** What is the amount of heat input to your skin when it receives the heat released (a) by 25.0 g of steam initially at  $100.0^\circ\text{C}$ , when it is cooled to skin temperature ( $34.0^\circ\text{C}$ )? (b) By 25.0 g of water initially at  $100.0^\circ\text{C}$ , when it is cooled to  $34.0^\circ\text{C}$ ? (c) What does this tell you about the relative severity of steam and hot water burns?

**17.53.** What must the initial speed of a lead bullet be at a temperature of  $25.0^\circ\text{C}$  so that the heat developed when it is brought to rest will be just sufficient to melt it? Assume that all the initial mechanical energy of the bullet is converted to heat and that no heat flows from the bullet to its surroundings. (Typical rifles have muzzle speeds that exceed the speed of sound in air, which is  $347 \text{ m/s}$  at  $25.0^\circ\text{C}$ .)

**17.54.** Evaporation of sweat is an important mechanism for temperature control in some warm-blooded animals. (a) What mass of water must evaporate from the skin of a 70.0-kg man to cool his body  $1.00^\circ\text{C}$ ? The heat of vaporization of water at body temperature ( $37^\circ\text{C}$ ) is  $2.42 \times 10^6 \text{ J/kg} \cdot \text{K}$ . The specific heat of a typical human body is  $3480 \text{ J/kg} \cdot \text{K}$  (see Exercise 17.37). (b) What volume of water must the man drink to replenish the evaporated water? Compare to the volume of a soft-drink can ( $355 \text{ cm}^3$ ).

**17.55. "The Ship of the Desert."** Camels require very little water because they are able to tolerate relatively large changes in their body temperature. While humans keep their body temperatures constant to within one or two Celsius degrees, a dehydrated camel permits its body temperature to drop to  $34.0^\circ\text{C}$  overnight and rise to  $40.0^\circ\text{C}$  during the day. To see how effective this mechanism is for saving water, calculate how many liters of water a 400 kg camel would have to drink if it attempted to keep its body temperature at a constant  $34.0^\circ\text{C}$  by evaporation of sweat during

the day (12 hours) instead of letting it rise to  $40.0^\circ\text{C}$ . (Note: The specific heat of a camel or other mammal is about the same as that of a typical human,  $3480\text{ J/kg}\cdot\text{K}$ . The heat of vaporization of water at  $34^\circ\text{C}$  is  $2.42 \times 10^6\text{ J/kg}$ .)

**17.56.** An asteroid with a diameter of 10 km and a mass of  $2.60 \times 10^{15}\text{ kg}$  impacts the earth at a speed of 32.0 km/s, landing in the Pacific Ocean. If 1.00% of the asteroid's kinetic energy goes to boiling the ocean water (assume an initial water temperature of  $10.0^\circ\text{C}$ ), what mass of water will be boiled away by the collision? (For comparison, the mass of water contained in Lake Superior is about  $2 \times 10^{15}\text{ kg}$ .)

**17.57.** A refrigerator door is opened and room-temperature air ( $20.0^\circ\text{C}$ ) fills the  $1.50\text{-m}^3$  compartment. A 10.0-kg turkey, also at room temperature, is placed in the refrigerator and the door is closed. The density of air is  $1.20\text{ kg/m}^3$  and its specific heat is  $1020\text{ J/kg}\cdot\text{K}$ . Assume the specific heat of a turkey, like that of a human, is  $3480\text{ J/kg}\cdot\text{K}$ . How much heat must the refrigerator remove from its compartment to bring the air and the turkey to thermal equilibrium at a temperature of  $5.00^\circ\text{C}$ ? Assume no heat exchange with the surrounding environment.

**17.58.** A laboratory technician drops a 0.0850-kg sample of unknown material, at a temperature of  $100.0^\circ\text{C}$ , into a calorimeter. The calorimeter can, initially at  $19.0^\circ\text{C}$ , is made of 0.150 kg of copper and contains 0.200 kg of water. The final temperature of the calorimeter can and contents is  $26.1^\circ\text{C}$ . Compute the specific heat capacity of the sample.

**17.59.** An insulated beaker with negligible mass contains 0.250 kg of water at a temperature of  $75.0^\circ\text{C}$ . How many kilograms of ice at a temperature of  $-20.0^\circ\text{C}$  must be dropped into the water to make the final temperature of the system  $30.0^\circ\text{C}$ ?

**17.60.** A glass vial containing a 16.0-g sample of an enzyme is cooled in an ice bath. The bath contains water and 0.120 kg of ice. The sample has specific heat  $2250\text{ J/kg}\cdot\text{K}$ ; the glass vial has mass 6.00 g and specific heat  $2800\text{ J/kg}\cdot\text{K}$ . How much ice melts in cooling the enzyme sample from room temperature ( $19.5^\circ\text{C}$ ) to the temperature of the ice bath?

**17.61.** A 4.00-kg silver ingot is taken from a furnace, where its temperature is  $750.0^\circ\text{C}$ , and placed on a large block of ice at  $0.0^\circ\text{C}$ . Assuming that all the heat given up by the silver is used to melt the ice, how much ice is melted?

**17.62.** A copper calorimeter can with mass 0.100 kg contains 0.160 kg of water and 0.0180 kg of ice in thermal equilibrium at atmospheric pressure. If 0.750 kg of lead at a temperature of  $255^\circ\text{C}$  is dropped into the calorimeter can, what is the final temperature? Assume that no heat is lost to the surroundings.

**17.63.** A vessel whose walls are thermally insulated contains 2.40 kg of water and 0.450 kg of ice, all at a temperature of  $0.0^\circ\text{C}$ . The outlet of a tube leading from a boiler in which water is boiling at atmospheric pressure is inserted into the water. How many grams of steam must condense inside the vessel (also at atmospheric pressure) to raise the temperature of the system to  $28.0^\circ\text{C}$ ? You can ignore the heat transferred to the container.

### Section 17.7 Mechanisms of Heat Transfer

**17.64.** Use Eq. (17.21) to show that the SI units of thermal conductivity are  $\text{W/m}\cdot\text{K}$ .

**17.65.** Suppose that the rod in Fig. 17.23a is made of copper, is 45.0 cm long, and has a cross-sectional area of  $1.25\text{ cm}^2$ . Let  $T_H = 100.0^\circ\text{C}$  and  $T_C = 0.0^\circ\text{C}$ . (a) What is the final steady-state temperature gradient along the rod? (b) What is the heat current in the rod in the final steady state? (c) What is the final steady-state temperature at a point in the rod 12.0 cm from its left end?

**17.66.** One end of an insulated metal rod is maintained at  $100.0^\circ\text{C}$ , and the other end is maintained at  $0.00^\circ\text{C}$  by an ice–water mixture. The rod is 60.0 cm long and has a cross-sectional area of  $1.25\text{ cm}^2$ . The heat conducted by the rod melts 8.50 g of ice in 10.0 min. Find the thermal conductivity  $k$  of the metal.

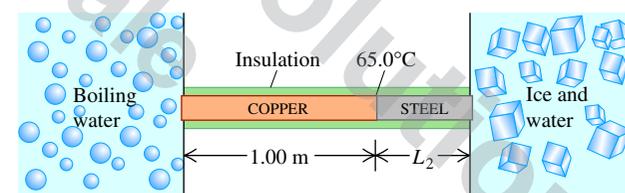
**17.67.** A carpenter builds an exterior house wall with a layer of wood 3.0 cm thick on the outside and a layer of Styrofoam insulation 2.2 cm thick on the inside wall surface. The wood has  $k = 0.080\text{ W/m}\cdot\text{K}$ , and the Styrofoam has  $k = 0.010\text{ W/m}\cdot\text{K}$ . The interior surface temperature is  $19.0^\circ\text{C}$ , and the exterior surface temperature is  $-10.0^\circ\text{C}$ . (a) What is the temperature at the plane where the wood meets the Styrofoam? (b) What is the rate of heat flow per square meter through this wall?

**17.68.** An electric kitchen range has a total wall area of  $1.40\text{ m}^2$  and is insulated with a layer of fiberglass 4.00 cm thick. The inside surface of the fiberglass has a temperature of  $175^\circ\text{C}$ , and its outside surface is at  $35.0^\circ\text{C}$ . The fiberglass has a thermal conductivity of  $0.040\text{ W/m}\cdot\text{K}$ . (a) What is the heat current through the insulation, assuming it may be treated as a flat slab with an area of  $1.40\text{ m}^2$ ? (b) What electric-power input to the heating element is required to maintain this temperature?

**17.69.** The ceiling of a room has an area of  $125\text{ ft}^2$ . The ceiling is insulated to an  $R$  value of 30 (in units of  $\text{ft}^2\cdot\text{F}^\circ\cdot\text{h/Btu}$ ). The surface in the room is maintained at  $69^\circ\text{F}$ , and the surface in the attic has a temperature of  $35^\circ\text{F}$ . What is the heat flow through the ceiling into the attic in 5.0 h? Express your answer in Btu and in joules.

**17.70.** A long rod, insulated to prevent heat loss along its sides, is in perfect thermal contact with boiling water (at atmospheric pressure) at one end and with an ice–water mixture at the other (Fig. 17.31). The rod consists of a 1.00-m section of copper (one end in boiling water) joined end to end to a length  $L_2$  of steel (one end in the ice–water mixture). Both sections of the rod have cross-sectional areas of  $4.00\text{ cm}^2$ . The temperature of the copper–steel junction is  $65.0^\circ\text{C}$  after a steady state has been set up. (a) How much heat per second flows from the boiling water to the ice–water mixture? (b) What is the length  $L_2$  of the steel section?

Figure 17.31 Exercise 17.70.



**17.71.** A pot with a steel bottom 8.50 mm thick rests on a hot stove. The area of the bottom of the pot is  $0.150\text{ m}^2$ . The water inside the pot is at  $100.0^\circ\text{C}$ , and 0.390 kg are evaporated every 3.00 min. Find the temperature of the lower surface of the pot, which is in contact with the stove.

**17.72.** You are asked to design a cylindrical steel rod 50.0 cm long, with a circular cross section, that will conduct  $150.0\text{ J/s}$  from a furnace at  $400.0^\circ\text{C}$  to a container of boiling water under 1 atmosphere. What must the rod's diameter be?

**17.73.** A picture window has dimensions of  $1.40\text{ m} \times 2.50\text{ m}$  and is made of glass 5.20 mm thick. On a winter day, the outside temperature is  $-20.0^\circ\text{C}$ , while the inside temperature is a comfortable  $19.5^\circ\text{C}$ . (a) At what rate is heat being lost through the window by conduction? (b) At what rate would heat be lost through the window if you covered it with a 0.750-mm-thick layer of paper (thermal conductivity 0.0500)?

**17.74.** What is the rate of energy radiation per unit area of a black-body at a temperature of (a) 273 K and (b) 2730 K?

**17.75.** What is the net rate of heat loss by radiation in Example 17.16 (Section 17.7) if the temperature of the surroundings is 5.0°C?

**17.76.** The emissivity of tungsten is 0.350. A tungsten sphere with radius 1.50 cm is suspended within a large evacuated enclosure whose walls are at 290.0 K. What power input is required to maintain the sphere at a temperature of 3000.0 K if heat conduction along the supports is neglected?

**17.77. Size of a Light-Bulb Filament.** The operating temperature of a tungsten filament in an incandescent light bulb is 2450 K, and its emissivity is 0.350. Find the surface area of the filament of a 150-W bulb if all the electrical energy consumed by the bulb is radiated by the filament as electromagnetic waves. (Only a fraction of the radiation appears as visible light.)

**17.78. The Sizes of Stars.** The hot glowing surfaces of stars emit energy in the form of electromagnetic radiation. It is a good approximation to assume  $e = 1$  for these surfaces. Find the radii of the following stars (assumed to be spherical): (a) Rigel, the bright blue star in the constellation Orion, which radiates energy at a rate of  $2.7 \times 10^{32}$  W and has surface temperature 11,000 K; (b) Procyon B (visible only using a telescope), which radiates energy at a rate of  $2.1 \times 10^{23}$  W and has surface temperature 10,000 K. (c) Compare your answers to the radius of the earth, the radius of the sun, and the distance between the earth and the sun. (Rigel is an example of a *supergiant* star, and Procyon B is an example of a *white dwarf* star.)

## Problems

**17.79.** You propose a new temperature scale with temperatures given in °M. You define 0.0°M to be the normal melting point of mercury and 100.0° to be the normal boiling point of mercury. (a) What is the normal boiling point of water in °M? (b) A temperature change of 10.0 M° corresponds to how many C°?

**17.80.** Suppose that a steel hoop could be constructed to fit just around the earth's equator at a temperature of 20.0°C. What would be the thickness of space between the hoop and the earth if the temperature of the hoop were increased by 0.500 C°?

**17.81.** At an absolute temperature  $T_0$ , a cube has sides of length  $L_0$  and has density  $\rho_0$ . The cube is made of a material with coefficient of volume expansion  $\beta$ . (a) Show that if the temperature increases to  $T_0 + \Delta T$ , the density of the cube becomes approximately

$$\rho \approx \rho_0(1 - \beta\Delta T)$$

(Hint: Use the expression  $(1 + x)^n \approx 1 + nx$ , valid for  $|x| \ll 1$ .) Explain why this approximate result is valid only if  $\Delta T$  is much less than  $1/\beta$ , and explain why you would expect this to be the case in most situations. (b) A copper cube has sides of length 1.25 cm at 20.0°C. Find the change in its volume and density when its temperature is increased to 70.0°C.

**17.82.** A 250-kg weight is hanging from the ceiling by a thin copper wire. In its fundamental mode, this wire vibrates at the frequency of concert A (440 Hz). You then increase the temperature of the wire by 40 C°. (a) By how much will the fundamental frequency change? Will it increase or decrease? (b) By what percentage will the speed of a wave on the wire change? (c) By what percentage will the wavelength of the fundamental standing wave change? Will it increase or decrease?

**17.83.** You are making pesto for your pasta and have a cylindrical measuring cup 10.0 cm high made of ordinary glass

[ $\beta = 2.7 \times 10^{-5} (\text{C}^\circ)^{-1}$ ] that is filled with olive oil [ $\beta = 6.8 \times 10^{-4} (\text{C}^\circ)^{-1}$ ] to a height of 1.00 mm below the top of the cup. Initially, the cup and oil are at room temperature (22.0°C). You get a phone call and forget about the olive oil, which you inadvertently leave on the hot stove. The cup and oil heat up slowly and have a common temperature. At what temperature will the olive oil start to spill out of the cup?

**17.84.** Use Fig. 17.12 to find the approximate coefficient of volume expansion of water at 2.0°C and at 8.0°C.

**17.85.** A Foucault pendulum consists of a brass sphere with a diameter of 35.0 cm suspended from a steel cable 10.5 m long (both measurements made at 20.0°C). Due to a design oversight, the swinging sphere clears the floor by a distance of only 2.00 mm when the temperature is 20.0°C. At what temperature will the sphere begin to brush the floor?

**17.86.** You pour 108 cm<sup>3</sup> of ethanol, at a temperature of  $-10.0^\circ\text{C}$ , into a graduated cylinder initially at 20.0°C, filling it to the very top. The cylinder is made of glass with a specific heat of 840 J/kg · K and a coefficient of volume expansion of  $1.2 \times 10^{-5} \text{K}^{-1}$ ; its mass is 0.110 kg. The mass of the ethanol is 0.0873 kg. (a) What will be the final temperature of the ethanol, once thermal equilibrium is reached? (b) How much ethanol will overflow the cylinder before thermal equilibrium is reached?

**17.87.** A metal rod that is 30.0 cm long expands by 0.0650 cm when its temperature is raised from 0.0°C to 100.0°C. A rod of a different metal and of the same length expands by 0.0350 cm for the same rise in temperature. A third rod, also 30.0 cm long, is made up of pieces of each of the above metals placed end to end and expands 0.0580 cm between 0.0°C and 100.0°C. Find the length of each portion of the composite rod.

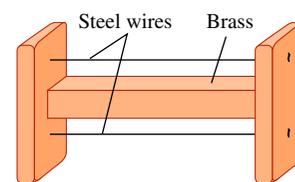
**17.88.** On a cool (4.0°C) Saturday morning, a pilot fills the fuel tanks of her Pitts S-2C (a two-seat aerobatic airplane) to their full capacity of 106.0 L. Before flying on Sunday morning, when the temperature is again 4.0°C, she checks the fuel level and finds only 103.4 L of gasoline in the tanks. She realizes that it was hot on Saturday afternoon, and that thermal expansion of the gasoline caused the missing fuel to empty out of the tank's vent. (a) What was the maximum temperature (in °C) reached by the fuel and the tank on Saturday afternoon? The coefficient of volume expansion of gasoline is  $9.5 \times 10^{-4} \text{K}^{-1}$ , and the tank is made of aluminum. (b) In order to have the maximum amount of fuel available for flight, when should the pilot have filled the fuel tanks?

**17.89.** (a) Equation (17.12) gives the stress required to keep the length of a rod constant as its temperature changes. Show that if the length is permitted to change by an amount  $\Delta L$  when its temperature changes by  $\Delta T$ , the stress is equal to

$$\frac{F}{A} = Y \left( \frac{\Delta L}{L_0} - \alpha \Delta T \right)$$

where  $F$  is the tension on the rod,  $L_0$  is the original length of the rod,  $A$  its cross-sectional area,  $\alpha$  its coefficient of linear expansion, and  $Y$  its Young's modulus. (b) A heavy brass bar has projections at its ends, as in Fig. 17.32. **Figure 17.32** Problem 17.89.

Two fine steel wires, fastened between the projections, are just taut (zero tension) when the whole system is at 20°C. What is the tensile stress in the steel wires when the temperature of the system is raised to 140°C? Make any simplifying assumptions you think are justified, but state what they are.



**17.90.** A steel rod 0.350 m long and an aluminum rod 0.250 m long, both with the same diameter, are placed end to end between rigid supports with no initial stress in the rods. The temperature of the rods is now raised by  $60.0^\circ\text{C}$ . What is the stress in each rod? (*Hint:* The length of the combined rod remains the same, but the lengths of the individual rods do not. See Problem 17.89.)

**17.91.** A steel ring with a 2.5000-in. inside diameter at  $20.0^\circ\text{C}$  is to be warmed and slipped over a brass shaft with a 2.5020-in. outside diameter at  $20.0^\circ\text{C}$ . (a) To what temperature should the ring be warmed? (b) If the ring and the shaft together are cooled by some means such as liquid air, at what temperature will the ring just slip off the shaft?

**17.92. Bulk Stress Due to a Temperature Increase.** (a) Prove that, if an object under pressure has its temperature raised but is not allowed to expand, the increase in pressure is

$$\Delta p = B\beta\Delta T$$

where the bulk modulus  $B$  and the average coefficient of volume expansion  $\beta$  are both assumed positive and constant. (b) What pressure is necessary to prevent a steel block from expanding when its temperature is increased from  $20.0^\circ\text{C}$  to  $35.0^\circ\text{C}$ ?

**17.93.** A liquid is enclosed in a metal cylinder that is provided with a piston of the same metal. The system is originally at a pressure of 1.00 atm ( $1.013 \times 10^5 \text{ Pa}$ ) and at a temperature of  $30.0^\circ\text{C}$ . The piston is forced down until the pressure on the liquid is increased by 50.0 atm, and then clamped in this position. Find the new temperature at which the pressure of the liquid is again 1.00 atm. Assume that the cylinder is sufficiently strong so that its volume is not altered by changes in pressure, but only by changes in temperature. Use the result derived in Problem 17.92. (*Hint:* See Section 11.4.)

Compressibility of liquid:  $k = 8.50 \times 10^{-10} \text{ Pa}^{-1}$

Coefficient of volume expansion of liquid:  $\beta = 4.80 \times 10^{-4} \text{ K}^{-1}$

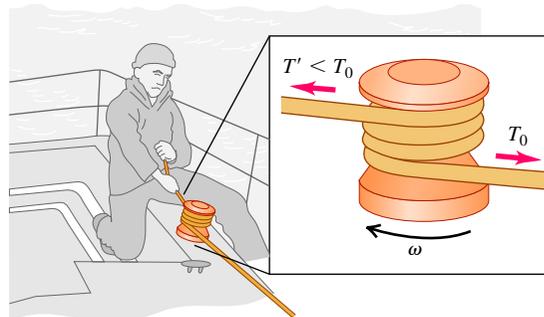
Coefficient of volume expansion of metal:  $\beta = 3.90 \times 10^{-5} \text{ K}^{-1}$

**17.94.** You cool a 100.0-g slug of red-hot iron (temperature  $745^\circ\text{C}$ ) by dropping it into an insulated cup of negligible mass containing 75.0 g of water at  $20.0^\circ\text{C}$ . Assuming no heat exchange with the surroundings, a) what is the final temperature of the water and b) what is the final mass of the iron and the remaining water?

**17.95. Spacecraft Reentry.** A spacecraft made of aluminum circles the earth at a speed of 7700 m/s. (a) Find the ratio of its kinetic energy to the energy required to raise its temperature from  $0^\circ\text{C}$  to  $600^\circ\text{C}$ . (The melting point of aluminum is  $660^\circ\text{C}$ . Assume a constant specific heat of  $910 \text{ J/kg} \cdot \text{K}$ .) (b) Discuss the bearing of your answer on the problem of the reentry of a manned space vehicle into the earth's atmosphere.

**17.96.** A capstan is a rotating drum or cylinder over which a rope or cord slides in order to provide a great amplification of the rope's tension while keeping both ends free (Fig. 17.33). Since the added tension in the rope is due to friction, the capstan generates thermal energy. (a) If the difference in tension between the two ends of the rope is 520.0 N and the capstan has a diameter of 10.0 cm and turns once in 0.900 s, find the rate at which thermal energy is generated. Why does the number of turns not matter? (b) If the capstan is made of iron and has mass 6.00 kg, at what rate does its temperature rise? Assume that the temperature in the capstan is uniform and that all the thermal energy generated flows into it.

**Figure 17.33** Problem 17.96.



**17.97. Debye's  $T^3$  Law.** At very low temperatures the molar heat capacity of rock salt varies with temperature according to Debye's  $T^3$  law:

$$C = k \frac{T^3}{\Theta^3}$$

where  $k = 1940 \text{ J/mol} \cdot \text{K}$  and  $\Theta = 281 \text{ K}$ . (a) How much heat is required to raise the temperature of 1.50 mol of rock salt from 10.0 K to 40.0 K? (*Hint:* Use Eq. (17.18) in the form  $dQ = nC dT$  and integrate.) (b) What is the average molar heat capacity in this range? (c) What is the true molar heat capacity at 40.0 K?

**17.98.** A person of mass 70.0 kg is sitting in the bathtub. The bathtub is 190.0 cm by 80.0 cm; before the person got in, the water was 10.0 cm deep. The water is at a temperature of  $37.0^\circ\text{C}$ . Suppose that the water were to cool down spontaneously to form ice at  $0.0^\circ\text{C}$ , and that all the energy released was used to launch the hapless bather vertically into the air. How high would the bather go? (As you will see in Chapter 20, this event is allowed by energy conservation but is prohibited by the second law of thermodynamics.)

**17.99. Hot Air in a Physics Lecture.** (a) A typical student listening attentively to a physics lecture has a heat output of 100 W. How much heat energy does a class of 90 physics students release into a lecture hall over the course of a 50-min lecture? (b) Assume that all the heat energy in part (a) is transferred to the  $3200 \text{ m}^3$  of air in the room. The air has specific heat capacity  $1020 \text{ J/kg} \cdot \text{K}$  and density  $1.20 \text{ kg/m}^3$ . If none of the heat escapes and the air conditioning system is off, how much will the temperature of the air in the room rise during the 50-min lecture? (c) If the class is taking an exam, the heat output per student rises to 280 W. What is the temperature rise during 50 min in this case?

**17.100.** The molar heat capacity of a certain substance varies with temperature according to the empirical equation

$$C = 29.5 \text{ J/mol} \cdot \text{K} + (8.20 \times 10^{-3} \text{ J/mol} \cdot \text{K}^2) T$$

How much heat is necessary to change the temperature of 3.00 mol of this substance from  $27^\circ\text{C}$  to  $227^\circ\text{C}$ ? (*Hint:* Use Eq. (17.18) in the form  $dQ = nC dT$  and integrate.)

**17.101.** For your cabin in the wilderness, you decide to build a primitive refrigerator out of Styrofoam, planning to keep the interior cool with a block of ice that has an initial mass of 24.0 kg. The box has dimensions of  $0.500 \text{ m} \times 0.800 \text{ m} \times 0.500 \text{ m}$ . Water from melting ice collects in the bottom of the box. Suppose the ice block is at  $0.00^\circ\text{C}$  and the outside temperature is  $21.0^\circ\text{C}$ . If the top of the empty box is never opened and you want the interior of the

box to remain at  $5.00^\circ\text{C}$  for exactly one week, until all the ice melts, what must be the thickness of the Styrofoam?

**17.102. Hot Water Versus Steam Heating.** In a household hot-water heating system, water is delivered to the radiators at  $70.0^\circ\text{C}$  ( $158.0^\circ\text{F}$ ) and leaves at  $28.0^\circ\text{C}$  ( $82.4^\circ\text{F}$ ). The system is to be replaced by a steam system in which steam at atmospheric pressure condenses in the radiators and the condensed steam leaves the radiators at  $35.0^\circ\text{C}$  ( $95.0^\circ\text{F}$ ). How many kilograms of steam will supply the same heat as was supplied by 1.00 kg of hot water in the first system?

**17.103.** A copper calorimeter can with mass 0.446 kg contains 0.0950 kg of ice. The system is initially at  $0.0^\circ\text{C}$ . (a) If 0.0350 kg of steam at  $100.0^\circ\text{C}$  and 1.00 atm pressure is added to the can, what is the final temperature of the calorimeter can and its contents? (b) At the final temperature, how many kilograms are there of ice, how many of liquid water, and how many of steam?

**17.104.** A Styrofoam bucket of negligible mass contains 1.75 kg of water and 0.450 kg of ice. More ice, from a refrigerator at  $-15.0^\circ\text{C}$ , is added to the mixture in the bucket, and when thermal equilibrium has been reached, the total mass of ice in the bucket is 0.778 kg. Assuming no heat exchange with the surroundings, what mass of ice was added?

**17.105.** In a container of negligible mass, 0.0400 kg of steam at  $100^\circ\text{C}$  and atmospheric pressure is added to 0.200 kg of water at  $50.0^\circ\text{C}$ . (a) If no heat is lost to the surroundings, what is the final temperature of the system? (b) At the final temperature, how many kilograms are there of steam and how many of liquid water?

**17.106.** A tube leads from a 0.150-kg calorimeter to a flask in which water is boiling under atmospheric pressure. The calorimeter has specific heat capacity  $420\text{ J/kg}\cdot\text{K}$ , and it originally contains 0.340 kg of water at  $15.0^\circ\text{C}$ . Steam is allowed to condense in the calorimeter at atmospheric pressure until the temperature of the calorimeter and contents reaches  $71.0^\circ\text{C}$ , at which point the total mass of the calorimeter and its contents is found to be 0.525 kg. Compute the heat of vaporization of water from these data.

**17.107.** A worker pours 1.250 kg of molten lead at a temperature of  $365.0^\circ\text{C}$  into 0.5000 kg of water at a temperature of  $75.00^\circ\text{C}$  in an insulated bucket of negligible mass. Assuming no heat loss to the surroundings, calculate the mass of lead and water remaining in the bucket when the materials have reached thermal equilibrium.

**17.108.** One experimental method of measuring an insulating material's thermal conductivity is to construct a box of the material and measure the power input to an electric heater inside the box that maintains the interior at a measured temperature above the outside surface. Suppose that in such an apparatus a power input of 180 W is required to keep the interior surface of the box  $65.0^\circ\text{C}$  (about  $120^\circ\text{F}$ ) above the temperature of the outer surface. The total area of the box is  $2.18\text{ m}^2$ , and the wall thickness is 3.90 cm. Find the thermal conductivity of the material in SI units.

**17.109. Effect of a Window in a Door.** A carpenter builds a solid wood door with dimensions  $2.00\text{ m} \times 0.95\text{ m} \times 5.0\text{ cm}$ . Its thermal conductivity is  $k = 0.120\text{ W/m}\cdot\text{K}$ . The air films on the inner and outer surfaces of the door have the same combined thermal resistance as an additional 1.8-cm thickness of solid wood. The inside air temperature is  $20.0^\circ\text{C}$ , and the outside air temperature is  $-8.0^\circ\text{C}$ . (a) What is the rate of heat flow through the door? (b) By what factor is the heat flow increased if a window 0.500 m on a side is inserted in the door? The glass is 0.450 cm thick, and the glass has a thermal conductivity of  $0.80\text{ W/m}\cdot\text{K}$ . The air films on the two sides of the glass have a total thermal resistance that is the same as an additional 12.0 cm of glass.

**17.110.** A wood ceiling with thermal resistance  $R_1$  is covered with a layer of insulation with thermal resistance  $R_2$ . Prove that the effective thermal resistance of the combination is  $R = R_1 + R_2$ .

**17.111.** Compute the ratio of the rate of heat loss through a single-pane window with area  $0.15\text{ m}^2$  to that for a double-pane window with the same area. The glass of a single pane is 4.2 mm thick, and the air space between the two panes of the double-pane window is 7.0 mm thick. The glass has thermal conductivity  $0.80\text{ W/m}\cdot\text{K}$ . The air films on the room and outdoor surfaces of either window have a combined thermal resistance of  $0.15\text{ m}^2\cdot\text{K/W}$ .

**17.112.** Rods of copper, brass, and steel are welded together to form a Y-shaped figure. The cross-sectional area of each rod is  $2.00\text{ cm}^2$ . The free end of the copper rod is maintained at  $100.0^\circ\text{C}$ , and the free ends of the brass and steel rods at  $0.0^\circ\text{C}$ . Assume there is no heat loss from the surfaces of the rods. The lengths of the rods are: copper, 13.0 cm; brass, 18.0 cm; steel, 24.0 cm. (a) What is the temperature of the junction point? (b) What is the heat current in each of the three rods?

**17.113. Time Needed for a Lake to Freeze Over.** (a) When the air temperature is below  $0^\circ\text{C}$ , the water at the surface of a lake freezes to form an ice sheet. Why doesn't freezing occur throughout the entire volume of the lake? (b) Show that the thickness of the ice sheet formed on the surface of a lake is proportional to the square root of the time if the heat of fusion of the water freezing on the underside of the ice sheet is conducted through the sheet. (c) Assuming that the upper surface of the ice sheet is at  $-10^\circ\text{C}$  and the bottom surface is at  $0^\circ\text{C}$ , calculate the time it will take to form an ice sheet 25 cm thick. (d) If the lake in part (c) is uniformly 40 m deep, how long would it take to freeze all the water in the lake? Is this likely to occur?

**17.114.** A rod is initially at a uniform temperature of  $0^\circ\text{C}$  throughout. One end is kept at  $0^\circ\text{C}$ , and the other is brought into contact with a steam bath at  $100^\circ\text{C}$ . The surface of the rod is insulated so that heat can flow only lengthwise along the rod. The cross-sectional area of the rod is  $2.50\text{ cm}^2$ , its length is 120 cm, its thermal conductivity is  $380\text{ W/m}\cdot\text{K}$ , its density is  $1.00 \times 10^4\text{ kg/m}^3$ , and its specific heat is  $520\text{ J/kg}\cdot\text{K}$ . Consider a short cylindrical element of the rod 1.00 cm in length. (a) If the temperature gradient at the cooler end of this element is  $140^\circ\text{C/m}$ , how many joules of heat energy flow across this end per second? (b) If the average temperature of the element is increasing at the rate of  $0.250^\circ\text{C/s}$ , what is the temperature gradient at the other end of the element?

**17.115.** A rustic cabin has a floor area of  $3.50\text{ m} \times 3.00\text{ m}$ . Its walls, which are 2.50 m tall, are made of wood (thermal conductivity  $0.0600\text{ W/m}\cdot\text{K}$ ) 1.80 cm thick and are further insulated with 1.50 cm of a synthetic material. When the outside temperature is  $2.00^\circ\text{C}$ , it is found necessary to heat the room at a rate of 1.25 kW to maintain its temperature at  $19.0^\circ\text{C}$ . Calculate the thermal conductivity of the insulating material. Neglect the heat lost through the ceiling and floor. Assume the inner and outer surfaces of the wall have the same temperature as the air inside and outside the cabin.

**17.116.** The rate at which radiant energy from the sun reaches the earth's upper atmosphere is about  $1.50\text{ kW/m}^2$ . The distance from the earth to the sun is  $1.50 \times 10^{11}\text{ m}$ , and the radius of the sun is  $6.96 \times 10^8\text{ m}$ . (a) What is the rate of radiation of energy per unit area from the sun's surface? (b) If the sun radiates as an ideal blackbody, what is the temperature of its surface?

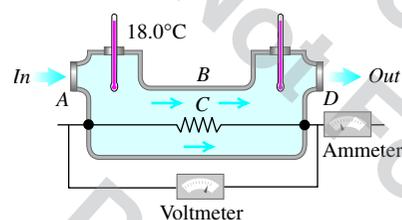
**17.117. A Thermos for Liquid Helium.** A physicist uses a cylindrical metal can 0.250 m high and 0.090 m in diameter to store liquid helium at  $4.22\text{ K}$ ; at that temperature the heat of vaporization of helium is  $2.09 \times 10^4\text{ J/kg}$ . Completely surrounding the metal

can are walls maintained at the temperature of liquid nitrogen, 77.3 K, with vacuum between the can and the surrounding walls. How much helium is lost per hour? The emissivity of the metal can is 0.200. The only heat transfer between the metal can and the surrounding walls is by radiation.

**17.118. Thermal Expansion of an Ideal Gas.** (a) The pressure  $p$ , volume  $V$ , number of moles  $n$ , and Kelvin temperature  $T$  of an ideal gas are related by the equation  $pV = nRT$ , where  $R$  is a constant. Prove that the coefficient of volume expansion for an ideal gas is equal to the reciprocal of the Kelvin temperature if the expansion occurs at constant pressure. (b) Compare the coefficients of volume expansion of copper and air at a temperature of 20°C. Assume that air may be treated as an ideal gas and that the pressure remains constant.

**17.119.** An engineer is developing an electric water heater to provide a continuous supply of hot water. One trial design is shown in Fig. 17.34. Water is flowing at the rate of 0.500 kg/min, the inlet thermometer registers 18.0°C, the voltmeter reads 120 V, and the ammeter reads 15.0 A [corresponding to a power input of  $(120 \text{ V}) \times (15.0 \text{ A}) = 1800 \text{ W}$ ]. (a) When a steady state is finally reached, what is the reading of the outlet thermometer? (b) Why is it unnecessary to take into account the heat capacity  $mc$  of the apparatus itself?

**Figure 17.34** Problem 17.119.



**17.120. Food Intake of a Hamster.** The energy output of an animal engaged in an activity is called the basal metabolic rate (BMR) and is a measure of the conversion of food energy into other forms of energy. A simple calorimeter to measure the BMR consists of an insulated box with a thermometer to measure the temperature of the air. The air has density  $1.20 \text{ kg/m}^3$  and specific heat  $1020 \text{ J/kg} \cdot \text{K}$ . A 50.0-g hamster is placed in a calorimeter that contains  $0.0500 \text{ m}^3$  of air at room temperature. (a) When the hamster is running in a wheel, the temperature of the air in the calorimeter rises  $1.60 \text{ C}^\circ$  per hour. How much heat does the running hamster generate in an hour? Assume that all this heat goes into the air in the calorimeter. You can ignore the heat that goes into the walls of the box and into the thermometer, and assume that no heat is lost to the surroundings. (b) Assuming that the hamster converts seed into heat with an efficiency of 10% and that hamster seed has a food energy value of  $24 \text{ J/g}$ , how many grams of seed must the hamster eat per hour to supply this energy?

**17.121.** The icecaps of Greenland and Antarctica contain about 1.75% of the total water (by mass) on the earth's surface; the oceans contain about 97.5%, and the other 0.75% is mainly groundwater. Suppose the icecaps, currently at an average temperature of about  $-30^\circ\text{C}$ , somehow slid into the ocean and melted. What would be the resulting temperature decrease of the ocean? Assume that the average temperature of ocean water is currently  $5.00^\circ\text{C}$ .

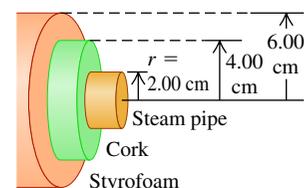
### Challenge Problems

**17.122.** (a) A spherical shell has inner and outer radii  $a$  and  $b$ , respectively, and the temperatures at the inner and outer surfaces

are  $T_2$  and  $T_1$ . The thermal conductivity of the material of which the shell is made is  $k$ . Derive an equation for the total heat current through the shell. (b) Derive an equation for the temperature variation within the shell in part (a); that is, calculate  $T$  as a function of  $r$ , the distance from the center of the shell. (c) A hollow cylinder has length  $L$ , inner radius  $a$ , and outer radius  $b$ , and the temperatures at the inner and outer surfaces are  $T_2$  and  $T_1$ . (The cylinder could represent an insulated hot-water pipe, for example.) The thermal conductivity of the material of which the cylinder is made is  $k$ . Derive an equation for the total heat current through the walls of the cylinder. (d) For the cylinder of part (c), derive an equation for the temperature variation inside the cylinder walls. (e) For the spherical shell of part (a) and the hollow cylinder of part (c), show that the equation for the total heat current in each case reduces to Eq. (17.21) for linear heat flow when the shell or cylinder is very thin.

**17.123.** A steam pipe with a radius of 2.00 cm, carrying steam at  $140^\circ\text{C}$ , is surrounded by a cylindrical jacket with inner and outer radii 2.00 cm and 4.00 cm and made of a type of cork with thermal conductivity  $4.00 \times 10^{-2} \text{ W/m} \cdot \text{K}$ . This in turn is surrounded by a cylindrical jacket made of a brand of Styrofoam with thermal conductivity  $1.00 \times 10^{-2} \text{ W/m} \cdot \text{K}$  and having inner and outer radii 4.00 cm and 6.00 cm (Fig. 17.35). The outer surface of the Styrofoam is in contact with air at  $15^\circ\text{C}$ . Assume that this outer surface has a temperature of  $15^\circ\text{C}$ .

**Figure 17.35** Challenge Problem 17.123.



(a) What is the temperature at a radius of 4.00 cm, where the two insulating layers meet? (b) What is the total rate of transfer of heat out of a 2.00-m length of pipe? (Hint: Use the expression derived in part (c) of Challenge Problem 17.122.)

**17.124.** Suppose that both ends of the rod in Fig. 17.23 are kept at a temperature of  $0^\circ\text{C}$ , and that the initial temperature distribution along the rod is given by  $T = (100^\circ\text{C}) \sin \pi x/L$ , where  $x$  is measured from the left end of the rod. Let the rod be copper, with length  $L = 0.100 \text{ m}$  and cross-sectional area  $1.00 \text{ cm}^2$ . (a) Show the initial temperature distribution in a diagram. (b) What is the final temperature distribution after a very long time has elapsed? (c) Sketch curves that you think would represent the temperature distribution at intermediate times. (d) What is the initial temperature gradient at the ends of the rod? (e) What is the initial heat current from the ends of the rod into the bodies making contact with its ends? (f) What is the initial heat current at the center of the rod? Explain. What is the heat current at this point at any later time? (g) What is the value of the *thermal diffusivity*  $k/\rho c$  for copper, and in what unit is it expressed? (Here  $k$  is the thermal conductivity,  $\rho = 8.9 \times 10^3 \text{ kg/m}^3$  is the density, and  $c$  is the specific heat.) (h) What is the initial time rate of change of temperature at the center of the rod? (i) How much time would be required for the temperature of the rod to reach its final temperature if the temperature continued to decrease at this rate? (This time is called the *relaxation time* of the rod.) (j) From the graphs in part (c), would you expect the magnitude of the rate of temperature change at the midpoint to remain constant, increase, or decrease as a function of time? (k) What is the initial rate of change of temperature at a point in the rod 2.5 cm from its left end?

**17.125. Temperature Change in a Clock.** A pendulum clock is designed to tick off one second on each side-to-side swing of the pendulum (two ticks per complete period). (a) Will a pendulum clock gain time in hot weather and lose it in cold, or the reverse? Explain your reasoning. (b) A particular pendulum clock keeps cor-

rect time at  $20.0^{\circ}\text{C}$ . The pendulum shaft is steel, and its mass can be ignored compared with that of the bob. What is the fractional change in the length of the shaft when it is cooled to  $10.0^{\circ}\text{C}$ ? (c) How many seconds per day will the clock gain or lose at  $10.0^{\circ}\text{C}$ ? (d) How closely must the temperature be controlled if the clock is not to gain or lose more than 1.00 s a day? Does the answer depend on the period of the pendulum?

**17.126.** One end of a solid cylindrical copper rod 0.200 m long is maintained at a temperature of  $20.00\text{ K}$ . The other end is blackened and exposed to thermal radiation from surrounding walls at  $500.0\text{ K}$ . The sides of the rod are insulated, so no energy is lost or gained except at the ends of the rod. When equilibrium is reached, what is the temperature of the blackened end? (*Hint:* Since copper is a very good conductor of heat at low temperature, with  $k = 1670\text{ W/m}\cdot\text{K}$  at  $20\text{ K}$ , the temperature of the blackened end is only slightly higher than  $20.00\text{ K}$ .)

**17.127. A Walk in the Sun.** Consider a poor lost soul walking at  $5\text{ km/h}$  on a hot day in the desert, wearing only a bathing suit. This person's skin temperature tends to rise due to four mechanisms:

(i) energy is generated by metabolic reactions in the body at a rate of  $280\text{ W}$ , and almost all of this energy is converted to heat that flows to the skin; (ii) heat is delivered to the skin by convection from the outside air at a rate equal to  $k'A_{\text{skin}}(T_{\text{air}} - T_{\text{skin}})$ , where  $k'$  is  $54\text{ J/h}\cdot\text{C}^{\circ}\cdot\text{m}^2$ , the exposed skin area  $A_{\text{skin}}$  is  $1.5\text{ m}^2$ , the air temperature  $T_{\text{air}}$  is  $47^{\circ}\text{C}$ , and the skin temperature  $T_{\text{skin}}$  is  $36^{\circ}\text{C}$ ; (iii) the skin absorbs radiant energy from the sun at a rate of  $1400\text{ W/m}^2$ ; (iv) the skin absorbs radiant energy from the environment, which has temperature  $47^{\circ}\text{C}$ . (a) Calculate the net rate (in watts) at which the person's skin is heated by all four of these mechanisms. Assume that the emissivity of the skin is  $e = 1$  and that the skin temperature is initially  $36^{\circ}\text{C}$ . Which mechanism is the most important? (b) At what rate (in L/h) must perspiration evaporate from this person's skin to maintain a constant skin temperature? (The heat of vaporization of water at  $36^{\circ}\text{C}$  is  $2.42 \times 10^6\text{ J/kg}$ .) (c) Suppose instead the person is protected by light-colored clothing ( $e \approx 0$ ) so that the exposed skin area is only  $0.45\text{ m}^2$ . What rate of perspiration is required now? Discuss the usefulness of the traditional clothing worn by desert peoples.

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