## FUNDAMENTALS OF THERMAL RADIATION

S o far, we have considered the conduction and convection modes of heat transfer, which are related to the nature of the materials involved and the presence of fluid motion, among other things. We now turn our attention to the third mechanism of heat transfer: *radiation*, which is characteristically different from the other two.

We start this chapter with a discussion of *electromagnetic waves* and the *electromagnetic spectrum*, with particular emphasis on *thermal radiation*. Then we introduce the idealized *blackbody*, *blackbody radiation*, and *blackbody radiation*, together with the *Stefan–Boltzmann law*, *Planck's law*, and *Wien's displacement law*.

Radiation is emitted by every point on a plane surface in all directions into the hemisphere above the surface. The quantity that describes the magnitude of radiation emitted or incident in a specified direction in space is the *radiation intensity*. Various radiation fluxes such as *emissive power, irradiation,* and *radiosity* are expressed in terms of intensity. This is followed by a discussion of radiative properties of materials such as *emissivity, absorptivity, reflectivity,* and *transmissivity* and their dependence on wavelength, direction, and temperature.

The *greenhouse effect* is presented as an example of the consequences of the wavelength dependence of radiation properties. The last section is devoted to the discussions of *atmospheric* and *solar radiation* because of their importance.

# CHAPTER

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### HEAT TRANSFER



#### FIGURE 11-1

A hot object in a vacuum chamber loses heat by radiation only.



### FIGURE 11-2

Unlike conduction and convection, heat transfer by radiation can occur between two bodies, even when they are separated by a medium colder than both of them.

### 11–1 • INTRODUCTION

Consider a hot object that is suspended in an evacuated chamber whose walls are at room temperature (Fig. 11–1). The hot object will eventually cool down and reach thermal equilibrium with its surroundings. That is, it will lose heat until its temperature reaches the temperature of the walls of the chamber. Heat transfer between the object and the chamber could not have taken place by conduction or convection, because these two mechanisms cannot occur in a vacuum. Therefore, heat transfer must have occurred through another mechanism that involves the emission of the internal energy of the object. This mechanism is *radiation*.

Radiation differs from the other two heat transfer mechanisms in that it does not require the presence of a material medium to take place. In fact, energy transfer by radiation is fastest (at the speed of light) and it suffers no attenuation in a *vacuum*. Also, radiation transfer occurs in solids as well as liquids and gases. In most practical applications, all three modes of heat transfer occur concurrently at varying degrees. But heat transfer through an evacuated space can occur only by radiation. For example, the energy of the sun reaches the earth by radiation.

You will recall that heat transfer by conduction or convection takes place in the direction of decreasing temperature; that is, from a high-temperature medium to a lower-temperature one. It is interesting that radiation heat transfer can occur between two bodies separated by a medium colder than both bodies (Fig. 11–2). For example, solar radiation reaches the surface of the earth after passing through cold air layers at high altitudes. Also, the radiation-absorbing surfaces inside a greenhouse reach high temperatures even when its plastic or glass cover remains relatively cool.

The theoretical foundation of radiation was established in 1864 by physicist James Clerk Maxwell, who postulated that accelerated charges or changing electric currents give rise to electric and magnetic fields. These rapidly moving fields are called **electromagnetic waves** or **electromagnetic radiation**, and they represent the energy emitted by matter as a result of the changes in the electronic configurations of the atoms or molecules. In 1887, Heinrich Hertz experimentally demonstrated the existence of such waves. Electromagnetic waves transport energy just like other waves, and all electromagnetic waves travel at the *speed of light* in a vacuum, which is  $C_0 = 2.9979 \times 10^8$  m/s. Electromagnetic waves are characterized by their *frequency*  $\nu$  or *wavelength*  $\lambda$ . These two properties in a medium are related by

 $\lambda = \frac{c}{\nu} \tag{11-1}$ 

where *c* is the speed of propagation of a wave in that medium. The speed of propagation in a medium is related to the speed of light in a vacuum by  $c = c_0/n$ , where *n* is the *index of refraction* of that medium. The refractive index is essentially unity for air and most gases, about 1.5 for glass, and about 1.33 for water. The commonly used unit of wavelength is the *micrometer* (µm) or micron, where 1 µm =  $10^{-6}$  m. Unlike the wavelength and the speed of propagation, the frequency of an electromagnetic wave depends only on the source and is independent of the medium through which the wave travels. The *frequency* (the number of oscillations per second) of an electromagnetic wave can range

from less than a million Hz to a septillion Hz or higher, depending on the source. Note from Eq. 11-1 that the wavelength and the frequency of electromagnetic radiation are inversely proportional.

It has proven useful to view electromagnetic radiation as the propagation of a collection of discrete packets of energy called **photons** or **quanta**, as proposed by Max Planck in 1900 in conjunction with his *quantum theory*. In this view, each photon of frequency  $\nu$  is considered to have an energy of

$$e = hv = \frac{hc}{\lambda} \tag{11-2}$$

where  $h = 6.6256 \times 10^{-34}$  J · s is *Planck's constant*. Note from the second part of Eq. 11-2 that the energy of a photon is inversely proportional to its wavelength. Therefore, shorter-wavelength radiation possesses larger photon energies. It is no wonder that we try to avoid very-short-wavelength radiation such as gamma rays and X-rays since they are highly destructive.

### 11–2 • THERMAL RADIATION

Although all electromagnetic waves have the same general features, waves of different wavelength differ significantly in their behavior. The electromagnetic radiation encountered in practice covers a wide range of wavelengths, varying from less than  $10^{-10}$  µm for cosmic rays to more than  $10^{10}$  µm for electrical power waves. The **electromagnetic spectrum** also includes gamma rays, X-rays, ultraviolet radiation, visible light, infrared radiation, thermal radiation, microwaves, and radio waves, as shown in Figure 11–3.

Different types of electromagnetic radiation are produced through various mechanisms. For example, *gamma rays* are produced by nuclear reactions, *X-rays* by the bombardment of metals with high-energy electrons, *microwaves* by special types of electron tubes such as klystrons and magnetrons, and *radio waves* by the excitation of some crystals or by the flow of alternating current through electric conductors.

The short-wavelength gamma rays and X-rays are primarily of concern to nuclear engineers, while the long-wavelength microwaves and radio waves are of concern to electrical engineers. The type of electromagnetic radiation that is pertinent to heat transfer is the **thermal radiation** emitted as a result of energy transitions of molecules, atoms, and electrons of a substance. Temperature is a measure of the strength of these activities at the microscopic level, and the rate of thermal radiation emission increases with increasing temperature. Thermal radiation is continuously emitted by all matter whose temperature is above absolute zero. That is, everything around us such as walls, furniture, and our friends constantly emits (and absorbs) radiation (Fig. 11–4). Thermal radiation is also defined as the portion of the electromagnetic spectrum that extends from about 0.1 to 100  $\mu$ m, since the radiation emitted by bodies due to their temperature falls almost entirely into this wavelength range. Thus, thermal radiation includes the entire visible and infrared (IR) radiation as well as a portion of the ultraviolet (UV) radiation.

What we call **light** is simply the *visible* portion of the electromagnetic spectrum that lies between 0.40 and 0.76  $\mu$ m. Light is characteristically no different than other electromagnetic radiation, except that it happens to trigger the









FIGURE 11–4 Everything around us constantly emits thermal radiation.

#### TABLE 11-1

### The wavelength ranges of different colors

Color	Wavelength band
Violet	0.40–0.44 μm
Blue	0.44–0.49 μm
Green	0.49–0.54 μm
Yellow	0.54–0.60 μm
Orange	0.60–0.67 μm
Red	0.63–0.76 μm



#### FIGURE 11–5

Food is heated or cooked in a microwave oven by absorbing the electromagnetic radiation energy generated by the magnetron of the oven. sensation of seeing in the human eye. Light, or the visible spectrum, consists of narrow bands of color from violet (0.40–0.44  $\mu$ m) to red (0.63–0.76  $\mu$ m), as shown in Table 11–1.

A body that emits some radiation in the visible range is called a light source. The sun is obviously our primary light source. The electromagnetic radiation emitted by the sun is known as **solar radiation**, and nearly all of it falls into the wavelength band  $0.3-3 \mu m$ . Almost *half* of solar radiation is light (i.e., it falls into the visible range), with the remaining being ultraviolet and infrared.

The radiation emitted by bodies at room temperature falls into the **infrared** region of the spectrum, which extends from 0.76 to 100  $\mu$ m. Bodies start emitting noticeable visible radiation at temperatures above 800 K. The tungsten filament of a lightbulb must be heated to temperatures above 2000 K before it can emit any significant amount of radiation in the visible range.

The **ultraviolet** radiation includes the low-wavelength end of the thermal radiation spectrum and lies between the wavelengths 0.01 and 0.40 µm. Ultraviolet rays are to be avoided since they can kill microorganisms and cause serious damage to humans and other living organisms. About 12 percent of so*lar radiation is in the ultraviolet range*, and it would be devastating if it were to reach the surface of the earth. Fortunately, the ozone (O<sub>3</sub>) layer in the atmosphere acts as a protective blanket and absorbs most of this ultraviolet radiation. The ultraviolet rays that remain in sunlight are still sufficient to cause serious sunburns to sun worshippers, and prolonged exposure to direct sunlight is the leading cause of skin cancer, which can be lethal. Recent discoveries of "holes" in the ozone layer have prompted the international community to ban the use of ozone-destroying chemicals such as the refrigerant Freon-12 in order to save the earth. Ultraviolet radiation is also produced artificially in fluorescent lamps for use in medicine as a bacteria killer and in tanning parlors as an artificial tanner. The connection between skin cancer and ultraviolet rays has caused dermatologists to issue strong warnings against its use for tanning.

Microwave ovens utilize electromagnetic radiation in the **microwave** region of the spectrum generated by microwave tubes called *magnetrons*. Microwaves in the range of  $10^2$ – $10^5$  µm are very suitable for use in cooking since they are *reflected* by metals, *transmitted* by glass and plastics, and *absorbed* by food (especially water) molecules. Thus, the electric energy converted to radiation in a microwave oven eventually becomes part of the internal energy of the food. The fast and efficient cooking of microwave ovens has made them as one of the essential appliances in modern kitchens (Fig. 11–5).

Radars and cordless telephones also use electromagnetic radiation in the microwave region. The wavelength of the electromagnetic waves used in radio and TV broadcasting usually ranges between 1 and 1000 m in the **radio wave** region of the spectrum.

In heat transfer studies, we are interested in the energy emitted by bodies because of their temperature only. Therefore, we will limit our consideration to *thermal radiation*, which we will simply call *radiation*. The relations developed below are restricted to thermal radiation only and may not be applicable to other forms of electromagnetic radiation.

The electrons, atoms, and molecules of all solids, liquids, and gases above absolute zero temperature are constantly in motion, and thus radiation is constantly emitted, as well as being absorbed or transmitted throughout the entire volume of matter. That is, radiation is a **volumetric phenomenon**. However, for opaque (nontransparent) solids such as metals, wood, and rocks, radiation is considered to be a **surface phenomenon**, since the radiation emitted by the interior regions can never reach the surface, and the radiation incident on such bodies is usually absorbed within a few microns from the surface (Fig. 11–6). Note that the radiation characteristics of surfaces can be changed completely by applying thin layers of coatings on them.

### 11–3 • BLACKBODY RADIATION

A body at a temperature above absolute zero emits radiation in all directions over a wide range of wavelengths. The amount of radiation energy emitted from a surface at a given wavelength depends on the material of the body and the condition of its surface as well as the surface temperature. Therefore, different bodies may emit different amounts of radiation per unit surface area, even when they are at the same temperature. Thus, it is natural to be curious about the *maximum* amount of radiation that can be emitted by a surface at a given temperature. Satisfying this curiosity requires the definition of an idealized body, called a *blackbody*, to serve as a standard against which the radiative properties of real surfaces may be compared.

A **blackbody** is defined as *a perfect emitter and absorber of radiation*. At a specified temperature and wavelength, no surface can emit more energy than a blackbody. A blackbody absorbs *all* incident radiation, regardless of wavelength and direction. Also, a blackbody emits radiation energy uniformly in all directions per unit area normal to direction of emission. (Fig. 11–7). That is, a blackbody is a *diffuse* emitter. The term *diffuse* means "independent of direction."

The radiation energy emitted by a blackbody per unit time and per unit surface area was determined experimentally by Joseph Stefan in 1879 and expressed as

$$E_b(T) = \sigma T^4$$
 (W/m<sup>2</sup>) (11-3)

where  $\sigma = 5.67 \times 10^{-8}$  W/m<sup>2</sup> · K<sup>4</sup> is the *Stefan–Boltzmann constant* and *T* is the absolute temperature of the surface in K. This relation was theoretically verified in 1884 by Ludwig Boltzmann. Equation 11-3 is known as the **Stefan–Boltzmann law** and  $E_b$  is called the **blackbody emissive power**. Note that the emission of thermal radiation is proportional to the *fourth power* of the absolute temperature.

Although a blackbody would appear *black* to the eye, a distinction should be made between the idealized blackbody and an ordinary black surface. Any surface that absorbs light (the visible portion of radiation) would appear black to the eye, and a surface that reflects it completely would appear white. Considering that visible radiation occupies a very narrow band of the spectrum from 0.4 to 0.76  $\mu$ m, we cannot make any judgments about the blackness of a surface on the basis of visual observations. For example, snow and white paint reflect light and thus appear white. But they are essentially black for infrared radiation since they strongly absorb long-wavelength radiation. Surfaces coated with lampblack paint approach idealized blackbody behavior.

Another type of body that closely resembles a blackbody is a *large cavity with a small opening*, as shown in Figure 11–8. Radiation coming in through the opening of area *A* will undergo multiple reflections, and thus it will have several chances to be absorbed by the interior surfaces of the cavity before any



### FIGURE 11-6

Radiation in opaque solids is considered a surface phenomenon since the radiation emitted only by the molecules at the surface can escape the solid.



#### FIGURE 11–7

A blackbody is said to be a *diffuse* emitter since it emits radiation energy uniformly in all directions.



FIGURE 11-8

A large isothermal cavity at temperature T with a small opening of area A closely resembles a blackbody of surface area A at the same temperature. part of it can possibly escape. Also, if the surface of the cavity is isothermal at temperature T, the radiation emitted by the interior surfaces will stream through the opening after undergoing multiple reflections, and thus it will have a diffuse nature. Therefore, the cavity will act as a perfect absorber and perfect emitter, and the opening will resemble a blackbody of surface area A at temperature T, regardless of the actual radiative properties of the cavity.

The Stefan–Boltzmann law in Eq. 11-3 gives the *total* blackbody emissive power  $E_b$ , which is the sum of the radiation emitted over all wavelengths. Sometimes we need to know the **spectral blackbody emissive power**, which is the amount of radiation energy emitted by a blackbody at an absolute temperature T per unit time, per unit surface area, and per unit wavelength about the wavelength  $\lambda$ . For example, we are more interested in the amount of radiation an incandescent light bulb emits in the visible wavelength spectrum than we are in the total amount emitted.

The relation for the spectral blackbody emissive power  $E_{b\lambda}$  was developed by Max Planck in 1901 in conjunction with his famous quantum theory. This relation is known as **Planck's law** and is expressed as

$$E_{b\lambda}(\lambda, T) = \frac{C_1}{\lambda^5 [\exp\left(C_2/\lambda T\right) - 1]} \qquad (W/m^2 \cdot \mu m) \tag{11-4}$$

where

$$C_1 = 2\pi h c_0^2 = 3.742 \times 10^8 \,\mathrm{W} \cdot \mu \mathrm{m}^4 / \mathrm{m}^2$$
  
$$C_2 = h c_0 / k = 1.439 \times 10^4 \,\mu \mathrm{m} \cdot \mathrm{K}$$

Also, *T* is the absolute temperature of the surface,  $\lambda$  is the wavelength of the radiation emitted, and  $k = 1.38065 \times 10^{-23}$  J/K is *Boltzmann's constant*. This relation is valid for a surface in a *vacuum* or a *gas*. For other mediums, it needs to be modified by replacing  $C_1$  by  $C_1/n^2$ , where *n* is the index of refraction of the medium. Note that the term *spectral* indicates dependence on wavelength.

The variation of the spectral blackbody emissive power with wavelength is plotted in Figure 11–9 for selected temperatures. Several observations can be made from this figure:

- **1.** The emitted radiation is a continuous function of *wavelength*. At any specified temperature, it increases with wavelength, reaches a peak, and then decreases with increasing wavelength.
- **2.** At any wavelength, the amount of emitted radiation *increases* with increasing temperature.
- **3.** As temperature increases, the curves shift to the left to the shorterwavelength region. Consequently, a larger fraction of the radiation is emitted at *shorter wavelengths* at higher temperatures.
- 4. The radiation emitted by the *sun*, which is considered to be a blackbody at 5780 K (or roughly at 5800 K), reaches its peak in the visible region of the spectrum. Therefore, the sun is in tune with our eyes. On the other hand, surfaces at  $T \le 800$  K emit almost entirely in the infrared region and thus are not visible to the eye unless they reflect light coming from other sources.

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#### FIGURE 11–9

The variation of the blackbody emissive power with wavelength for several temperatures.

As the temperature increases, the peak of the curve in Figure 11–9 shifts toward shorter wavelengths. The wavelength at which the peak occurs for a specified temperature is given by **Wien's displacement law** as

$$(\lambda T)_{\text{max power}} = 2897.8 \ \mu\text{m} \cdot \text{K} \tag{11-5}$$

This relation was originally developed by Willy Wien in 1894 using classical thermodynamics, but it can also be obtained by differentiating Eq. 11-4 with respect to  $\lambda$  while holding *T* constant and setting the result equal to zero. A plot of Wien's displacement law, which is the locus of the peaks of the radiation emission curves, is also given in Figure 11–9.

The peak of the solar radiation, for example, occurs at  $\lambda = 2897.8/5780 = 0.50 \,\mu\text{m}$ , which is near the middle of the visible range. The peak of the radiation emitted by a surface at room temperature (T = 298 K) occurs at 9.72  $\mu$ m, which is well into the infrared region of the spectrum.

An electrical resistance heater starts radiating heat soon after it is plugged in, and we can feel the emitted radiation energy by holding our hands facing the heater. But this radiation is entirely in the infrared region and thus cannot



### FIGURE 11–10

A surface that reflects red while absorbing the remaining parts of the incident light appears red to the eye.



### FIGURE 11–11

On an  $E_{b\lambda}$ - $\lambda$  chart, the area under a curve for a given temperature represents the total radiation energy emitted by a blackbody at that temperature. be sensed by our eyes. The heater would appear dull red when its temperature reaches about 1000 K, since it will start emitting a detectable amount (about  $1 \text{ W/m}^2 \cdot \mu m$ ) of visible red radiation at that temperature. As the temperature rises even more, the heater appears bright red and is said to be *red hot*. When the temperature reaches about 1500 K, the heater emits enough radiation in the entire visible range of the spectrum to appear almost *white* to the eye, and it is called *white hot*.

Although it cannot be sensed directly by the human eye, infrared radiation can be detected by infrared cameras, which transmit the information to microprocessors to display visual images of objects at night. *Rattlesnakes* can sense the infrared radiation or the "body heat" coming off warm-blooded animals, and thus they can see at night without using any instruments. Similarly, honeybees are sensitive to ultraviolet radiation. A surface that reflects all of the light appears *white*, while a surface that absorbs all of the light incident on it appears black. (Then how do we see a black surface?)

It should be clear from this discussion that the color of an object is not due to emission, which is primarily in the infrared region, unless the surface temperature of the object exceeds about 1000 K. Instead, the color of a surface depends on the absorption and reflection characteristics of the surface and is due to selective absorption and reflection of the incident visible radiation coming from a light source such as the sun or an incandescent lightbulb. A piece of clothing containing a pigment that reflects red while absorbing the remaining parts of the incident light appears "red" to the eye (Fig. 11–10). Leaves appear "green" because their cells contain the pigment chlorophyll, which strongly reflects green while absorbing other colors.

It is left as an exercise to show that integration of the *spectral* blackbody emissive power  $E_{b\lambda}$  over the entire wavelength spectrum gives the *total* blackbody emissive power  $E_b$ :

$$E_b(T) = \int_0^\infty E_{b\lambda}(\lambda, T) \, d\lambda = \sigma T^4 \qquad (W/m^2) \tag{11-6}$$

Thus, we obtained the Stefan–Boltzmann law (Eq. 11-3) by integrating Planck's law (Eq. 11-4) over all wavelengths. Note that on an  $E_{b\lambda}$ – $\lambda$  chart,  $E_{b\lambda}$  corresponds to any value on the curve, whereas  $E_b$  corresponds to the area under the entire curve for a specified temperature (Fig. 11–11). Also, the term *total* means "integrated over all wavelengths."

### **EXAMPLE 11-1** Radiation Emission from a Black Ball

Consider a 20-cm-diameter spherical ball at 800 K suspended in air as shown in Figure 11–12. Assuming the ball closely approximates a blackbody, determine (*a*) the total blackbody emissive power, (*b*) the total amount of radiation emitted by the ball in 5 min, and (*c*) the spectral blackbody emissive power at a wavelength of 3  $\mu$ m.

**SOLUTION** An isothermal sphere is suspended in air. The total blackbody emissive power, the total radiation emitted in 5 minutes, and the spectral blackbody emissive power at 3 mm are to be determined. *Assumptions* The ball behaves as a blackbody.

*Analysis* (a) The total blackbody emissive power is determined from the Stefan–Boltzmann law to be

$$E_b = \sigma T^4 = (5.67 \times 10^{-8} \,\text{W/m}^2 \cdot \text{K}^4)(800 \,\text{K})^4 = 23.2 \times 10^3 \,\text{W/m}^2 = 23.2 \,\text{kW/m}^2$$

That is, the ball emits 23.2 kJ of energy in the form of electromagnetic radiation per second per  $m^2$  of the surface area of the ball.

(*b*) The total amount of radiation energy emitted from the entire ball in 5 min is determined by multiplying the blackbody emissive power obtained above by the total surface area of the ball and the given time interval:

$$A_{s} = \pi D^{2} = \pi (0.2 \text{ m})^{2} = 0.1257 \text{ m}^{2}$$
$$\Delta t = (5 \text{ min}) \left( \frac{60 \text{ s}}{1 \text{ min}} \right) = 300 \text{ s}$$
$$Q_{\text{rad}} = E_{b} A_{s} \Delta t = (23.2 \text{ kW/m}^{2})(0.1257 \text{ m}^{2})(300 \text{ s}) \left( \frac{1 \text{ kJ}}{1000 \text{ W} \cdot \text{ s}} \right)$$
$$= 876 \text{ kJ}$$

That is, the ball loses 876 kJ of its internal energy in the form of electromagnetic waves to the surroundings in 5 min, which is enough energy to raise the temperature of 1 kg of water by 50°C. Note that the surface temperature of the ball cannot remain constant at 800 K unless there is an equal amount of energy flow to the surface from the surroundings or from the interior regions of the ball through some mechanisms such as chemical or nuclear reactions.

(c) The spectral blackbody emissive power at a wavelength of 3  $\mu m$  is determined from Planck's distribution law to be

$$E_{b\lambda} = \frac{C_1}{\lambda^5 \left[ \exp\left(\frac{C_2}{\lambda T}\right) - 1 \right]} = \frac{3.743 \times 10^8 \,\mathrm{W} \cdot \mu \mathrm{m}^4 / \mathrm{m}^2}{(3 \,\mu \mathrm{m})^5 \left[ \exp\left(\frac{1.4387 \times 10^4 \,\mu \mathrm{m} \cdot \mathrm{K}}{(3 \,\mu \mathrm{m})(800 \,\mathrm{K})}\right) - 1 \right]}$$
  
= 3848 W/m<sup>2</sup> · \mu m

The Stefan–Boltzmann law  $E_b(T) = \sigma T^4$  gives the *total* radiation emitted by a blackbody at all wavelengths from  $\lambda = 0$  to  $\lambda = \infty$ . But we are often interested in the amount of radiation emitted over *some wavelength band*. For example, an incandescent lightbulb is judged on the basis of the radiation it emits in the visible range rather than the radiation it emits at all wavelengths.

The radiation energy emitted by a blackbody per unit area over a wavelength band from  $\lambda = 0$  to  $\lambda$  is determined from (Fig. 11–13)

$$E_{b,0-\lambda}(T) = \int_0^{\lambda} E_{b\lambda}(\lambda, T) \, d\lambda \qquad (W/m^2) \tag{11-7}$$

It looks like we can determine  $E_{b, 0-\lambda}$  by substituting the  $E_{b\lambda}$  relation from Eq. 11-4 and performing this integration. But it turns out that this integration does not have a simple closed-form solution, and performing a numerical integration each time we need a value of  $E_{b, 0-\lambda}$  is not practical. Therefore, we define a dimensionless quantity  $f_{\lambda}$  called the **blackbody radiation function** as







FIGURE 11–12 The spherical ball considered in Example 11–1.

TABLE 11-2							
Blackbody radiation functions $f_{\lambda}$							
λΤ,		λ <i>Τ</i> ,					
μm · K	$f_{\lambda}$	μm・K	$f_{\lambda}$				
200	0.000000	6200	0.754140				
400	0.000000	6400	0.769234				
600	0.000000	6600	0.783199				
800	0.000016	6800	0.796129				
1000	0.000321	7000	0.808109				
1200	0.002134	7200	0.819217				
1400	0.007790	7400	0.829527				
1600	0.019718	7600	0.839102				
1800	0.039341	7800	0.848005				
2000	0.066728	8000	0.856288				
2200	0.100888	8500	0.874608				
2400	0.140256	9000	0.890029				
2600	0.183120	9500	0.903085				
2800	0.227897	10,000	0.914199				
3000	0.273232	10,500	0.923710				
3200	0.318102	11,000	0.931890				
3400	0.361735	11,500	0.939959				
3600	0.403607	12,000	0.945098				
3800	0.443382	13,000	0.955139				
4000	0.480877	14,000	0.962898				
4200	0.516014	15,000	0.969981				
4400	0.548796	16,000	0.973814				
4600	0.579280	18,000	0.980860				
4800	0.607559	20,000	0.985602				
5000	0.633747	25,000	0.992215				
5200	0.658970	30,000	0.995340				
5400	0.680360	40,000	0.997967				
5600	0.701046	50,000	0.998953				
5800	0.720158	75,000	0.999713				
6000	0.737818	100,000	0.999905				



$$f_{\lambda}(T) = \frac{\int_{0}^{\lambda} E_{b\lambda}(\lambda, T) \, d\lambda}{\sigma T^4}$$
(11-8)

The function  $f_{\lambda}$  represents the fraction of radiation emitted from a blackbody at temperature T in the wavelength band from  $\lambda = 0$  to  $\lambda$ . The values of  $f_{\lambda}$  are listed in Table 11–2 as a function of  $\lambda T$ , where  $\lambda$  is in  $\mu$ m and T is in K.

The fraction of radiation energy emitted by a blackbody at temperature *T* over a finite wavelength band from  $\lambda = \lambda_1$  to  $\lambda = \lambda_2$  is determined from (Fig. 11–14)

$$f_{\lambda_1 - \lambda_2}(T) = f_{\lambda_2}(T) - f_{\lambda_1}(T)$$
 (11-9)

where  $f_{\lambda_1}(T)$  and  $f_{\lambda_2}(T)$  are blackbody radiation functions corresponding to  $\lambda_1 T$  and  $\lambda_2 T$ , respectively.

### **FIGURE 11–14**

Graphical representation of the fraction of radiation emitted in the wavelength band from  $\lambda_1$  to  $\lambda_2$ .

### **EXAMPLE 11–2** Emission of Radiation from a Lightbulb

The temperature of the filament of an incandescent lightbulb is 2500 K. Assuming the filament to be a blackbody, determine the fraction of the radiant energy emitted by the filament that falls in the visible range. Also, determine the wavelength at which the emission of radiation from the filament peaks.

**SOLUTION** The temperature of the filament of an incandescent lightbulb is given. The fraction of visible radiation emitted by the filament and the wavelength at which the emission peaks are to be determined.

Assumptions The filament behaves as a blackbody.

**Analysis** The visible range of the electromagnetic spectrum extends from  $\lambda_1 = 0.4 \ \mu m$  to  $\lambda_2 = 0.76 \ \mu m$ . Noting that  $T = 2500 \ K$ , the blackbody radiation functions corresponding to  $\lambda_1 T$  and  $\lambda_2 T$  are determined from Table 11–2 to be

$$\lambda_1 T = (0.40 \ \mu\text{m})(2500 \ \text{K}) = 1000 \ \mu\text{m} \cdot \text{K} \longrightarrow f_{\lambda_1} = 0.000321$$
  
 $\lambda_2 T = (0.76 \ \mu\text{m})(2500 \ \text{K}) = 1900 \ \mu\text{m} \cdot \text{K} \longrightarrow f_{\lambda_2} = 0.053035$ 

That is, 0.03 percent of the radiation is emitted at wavelengths less than 0.4  $\mu$ m and 5.3 percent at wavelengths less than 0.76  $\mu$ m. Then the fraction of radiation emitted between these two wavelengths is (Fig. 11–15)

$$f_{\lambda_1 - \lambda_2} = f_{\lambda_2} - f_{\lambda_1} = 0.053035 - 0.000321 = 0.0527135$$

Therefore, only about 5 percent of the radiation emitted by the filament of the lightbulb falls in the visible range. The remaining 95 percent of the radiation appears in the infrared region in the form of radiant heat or "invisible light," as it used to be called. This is certainly not a very efficient way of converting electrical energy to light and explains why fluorescent tubes are a wiser choice for lighting.

The wavelength at which the emission of radiation from the filament peaks is easily determined from Wien's displacement law to be

$$(\lambda T)_{\text{max power}} = 2897.8 \ \mu\text{m} \cdot \text{K} \quad \rightarrow \quad \lambda_{\text{max power}} = \frac{2897.8 \ \mu\text{m} \cdot \text{K}}{2500 \ \text{K}} = 1.16 \ \mu\text{m}$$

*Discussion* Note that the radiation emitted from the filament peaks in the infrared region.



Graphical representation of the fraction of radiation emitted in the visible range in Example 11–2.

### 11–4 • RADIATION INTENSITY

Radiation is emitted by all parts of a plane surface in all directions into the hemisphere above the surface, and the directional distribution of emitted (or incident) radiation is usually not uniform. Therefore, we need a quantity that describes the magnitude of radiation emitted (or incident) in a specified direction in space. This quantity is *radiation intensity*, denoted by *I*. Before we can describe a directional quantity, we need to specify direction in space. The direction of radiation passing through a point is best described in spherical coordinates in terms of the zenith angle  $\theta$  and the azimuth angle  $\phi$ , as shown in

### HEAT TRANSFER



FIGURE 11–16

Radiation intensity is used to describe the variation of radiation energy with direction.



A slice of pizza of plain angle  $\boldsymbol{\alpha}$ 



A slice of watermelon of solid angle  $\omega$  FIGURE 11–17

Describing the size of a slice of pizza by a plain angle, and the size of a watermelon slice by a solid angle. Figure 11–16. Radiation intensity is used to describe how the emitted radiation varies with the zenith and azimuth angles.

If all surfaces emitted radiation uniformly in all directions, the *emissive power* would be sufficient to quantify radiation, and we would not need to deal with intensity. The radiation emitted by a blackbody per unit normal area is the same in all directions, and thus there is no directional dependence. But this is not the case for real surfaces. Before we define intensity, we need to quantify the size of an opening in space.

### **Solid Angle**

Let us try to quantify the size of a slice of pizza. One way of doing that is to specify the arc length of the outer edge of the slice, and to form the slice by connecting the endpoints of the arc to the center. A more general approach is to specify the angle of the slice at the center, as shown in Figure 11–17. An angle of 90° (or  $\pi/2$  radians), for example, always represents a quarter pizza, no matter what the radius is. For a circle of unit radius, the length of an arc is equivalent in magnitude to the *plane angle* it subtends (both are  $2\pi$  for a complete circle of radius r = 1).

Now consider a watermelon, and let us attempt to quantify the size of a slice. Again we can do it by specifying the outer surface area of the slice (the green part), or by working with angles for generality. Connecting all points at the edges of the slice to the center in this case will form a three-dimensional body (like a cone whose tip is at the center), and thus the angle at the center in this case is properly called the **solid angle**. The solid angle is denoted by  $\omega$ , and its unit is the *steradian* (sr). In analogy to plane angle, we can say that *the area of a surface on a sphere of unit radius is equivalent in magnitude to the solid angle it subtends* (both are  $4\pi$  for a sphere of radius r = 1).

This can be shown easily by considering a differential surface area on a sphere  $dS = r^2 \sin \theta d\theta d\phi$ , as shown in Figure 11–18, and integrating it from  $\theta = 0$  to  $\theta = \pi$ , and from  $\phi = 0$  to  $\phi = 2\pi$ . We get

$$S = \int_{\text{sphere}} dS = \int_{\phi=0}^{2\pi} \int_{\theta=0}^{\pi} r^2 \sin \theta d\theta \phi = 2\pi r^2 \int_{\theta=0}^{\pi} \sin \theta d\theta = 4\pi r^2$$
(11-10)

which is the formula for the area of a sphere. For r = 1 it reduces to  $S = 4\pi$ , and thus the solid angle associated with a sphere is  $\omega = 4\pi$  sr. For a hemisphere, which is more relevant to radiation emitted or received by a surface, it is  $\omega = 2\pi$  sr.

The differential solid angle  $d\omega$  subtended by a differential area dS on a sphere of radius *r* can be expressed as

$$d\omega = \frac{dS}{r^2} = \sin \theta d\theta d\phi \tag{11-11}$$

Note that the area dS is normal to the direction of viewing since dS is viewed from the center of the sphere. In general, the differential solid angle  $d\omega$  subtended by a differential surface area dA when viewed from a point at a distance r from dA is expressed as

$$d\omega = \frac{dA_n}{r^2} = \frac{dA\cos\alpha}{r^2}$$
(11-12)

### CHAPTER 11



FIGURE 11–18

The emission of radiation from a differential surface element into the surrounding hemispherical space through a differential solid angle.

where  $\alpha$  is the angle between the normal of the surface and the direction of viewing, and thus  $dA_n = dA \cos \alpha$  is the normal (or projected) area to the direction of viewing.

Small surfaces viewed from relatively large distances can approximately be treated as differential areas in solid angle calculations. For example, the solid angle subtended by a 5 cm<sup>2</sup> plane surface when viewed from a point O at a distance of 80 cm along the normal of the surface is

$$\omega \simeq \frac{A_n}{r^2} = \frac{5 \text{ cm}^2}{(80 \text{ cm})^2} = 7.81 \times 10^{-4} \text{ sr}$$

If the surface is tilted so that the normal of the surface makes an angle of  $\alpha = 60^{\circ}$  with the line connecting point *O* to the center of the surface, the projected area would be  $dA_n = dA \cos \alpha = (5 \text{ cm}^2)\cos 60^{\circ} = 2.5 \text{ cm}^2$ , and the solid angle in this case would be half of the value just determined.

### Intensity of Emitted Radiation

Consider the emission of radiation by a differential area element dA of a surface, as shown in Figure 11–18. Radiation is emitted in all directions into the hemispherical space, and the radiation streaming though the surface area dS is proportional to the solid angle  $d\omega$  subtended by dS. It is also proportional to the radiating area dA as seen by an observer on dS, which varies from a maximum of dA when dS is at the top directly above dA ( $\theta = 0^{\circ}$ ) to a minimum of zero when dS is at the bottom ( $\theta = 90^{\circ}$ ). Therefore, the effective area of dA for emission in the direction of  $\theta$  is the projection of dA on a plane normal to  $\theta$ , which is  $dA \cos \theta$ . Radiation intensity in a given direction is based on a unit area normal to that direction to provide a common basis for the comparison of radiation emitted in different directions.

The **radiation intensity** for emitted radiation  $I_e(\theta, \phi)$  is defined as *the rate* at which radiation energy  $d\dot{Q}_e$  is emitted in the  $(\theta, \phi)$  direction per unit area normal to this direction and per unit solid angle about this direction. That is,

$$I_e(\theta, \phi) = \frac{dQ_e}{dA\cos\theta \cdot d\omega} = \frac{dQ_e}{dA\cos\theta\sin\theta d\theta d\phi} \qquad (W/m^2 \cdot sr) \qquad (11-13)$$





Radiation intensity is based on projected area, and thus the calculation of radiation emission from a surface involves the projection of the surface.



**FIGURE 11–20** Radiation incident on a surface in the direction  $(\theta, \phi)$ .

The *radiation flux* for emitted radiation is the **emissive power** E (the rate at which radiation energy is emitted per unit area of the emitting surface), which can be expressed in differential form as

$$dE = \frac{d\dot{Q}_e}{dA} = I_e(\theta, \phi) \cos \theta \sin \theta d\theta d\phi$$
(11-14)

Noting that the hemisphere above the surface will intercept all the radiation rays emitted by the surface, the emissive power from the surface into the hemisphere surrounding it can be determined by integration as

$$E = \int_{\text{hemisphere}} dE = \int_{\phi=0}^{2\pi} \int_{\theta=0}^{\pi/2} I_e(\theta, \phi) \cos \theta \sin \theta d\theta d\phi \qquad (W/m^2)$$
(11-15)

The intensity of radiation emitted by a surface, in general, varies with direction (especially with the zenith angle  $\theta$ ). But many surfaces in practice can be approximated as being diffuse. For a *diffusely emitting* surface, the intensity of the emitted radiation is independent of direction and thus  $I_e = \text{constant}$ . Noting that  $\int_{\phi=0}^{2\pi} \int_{\theta=0}^{\pi/2} \cos \theta \sin \theta d\theta d\phi = \pi$ , the emissive power relation in Eq. 11-15 reduces in this case to

Diffusely emitting surface:  $E = \pi I_e$  (W/m<sup>2</sup>) (11-16)

Note that the factor in Eq. 11-16 is  $\pi$ . You might have expected it to be  $2\pi$  since intensity is radiation energy per unit solid angle, and the solid angle associated with a hemisphere is  $2\pi$ . The reason for the factor being  $\pi$  is that the emissive power is based on the *actual* surface area whereas the intensity is based on the *projected* area (and thus the factor  $\cos \theta$  that accompanies it), as shown in Figure 11–19.

For a *blackbody*, which is a diffuse emitter, Eq. 11-16 can be expressed as

Blackbody: 
$$E_b = \pi I_b$$
 (11-17)

where  $E_b = \sigma T^4$  is the blackbody emissive power. Therefore, the intensity of the radiation emitted by a blackbody at absolute temperature *T* is

Blackbody: 
$$I_b(T) = \frac{E_b(T)}{\pi} = \frac{\sigma T^4}{\pi}$$
 (W/m<sup>2</sup> · sr) (11-18)

### **Incident Radiation**

All surfaces emit radiation, but they also receive radiation emitted or reflected by other surfaces. The intensity of incident radiation  $I_i(\theta, \phi)$  is defined as *the rate at which radiation energy dG is incident from the*  $(\theta, \phi)$  *direction per unit area of the receiving surface normal to this direction and per unit solid angle about this direction* (Fig. 11–20). Here  $\theta$  is the angle between the direction of incident radiation and the normal of the surface.

The radiation flux incident on a surface from *all directions* is called **irradiation** *G*, and is expressed as

$$G = \int_{\phi=0}^{2\pi} dG = \int_{\phi=0}^{2\pi} \int_{\theta=0}^{\pi/2} I_i(\theta, \phi) \cos \theta \sin \theta d\theta d\phi \qquad (W/m^2)$$
(11-19)

Therefore irradiation represents the rate at which radiation energy is incident on a surface per unit area of the surface. When the incident radiation is diffuse and thus  $I_i = \text{constant}$ , Eq. 11-19 reduces to

Diffusely incident radiation:

on:  $G = \pi I_i$ 

(11-20)

Again note that irradiation is based on the *actual* surface area (and thus the factor  $\cos \theta$ ), whereas the intensity of incident radiation is based on the *projected* area.

 $(W/m^2)$ 

### **Radiosity**

Surfaces emit radiation as well as reflecting it, and thus the radiation leaving a surface consists of emitted and reflected components, as shown in Figure 11–21. The calculation of radiation heat transfer between surfaces involves the *total* radiation energy streaming away from a surface, with no regard for its origin. Thus, we need to define a quantity that represents *the rate at which radiation energy leaves a unit area of a surface in all directions*. This quantity is called the **radiosity** *J*, and is expressed as

$$J = \int_{\phi=0}^{2\pi} \int_{\theta=0}^{\pi/2} I_{e+r}(\theta, \phi) \cos \theta \sin \theta d\theta d\phi \qquad (W/m^2)$$
(11-21)

where  $I_{e+r}$  is the sum of the emitted and reflected intensities. For a surface that is both a diffuse emitter and a diffuse reflector,  $I_{e+r} = \text{constant}$ , and the radiosity relation reduces to

Diffuse emitter and reflector:  $J = \pi I_{e+r}$  (W/m<sup>2</sup>) (11-22)

For a blackbody, radiosity J is equivalent to the emissive power  $E_b$  since a blackbody absorbs the entire radiation incident on it and there is no reflected component in radiosity.

### **Spectral Quantities**

So far we considered *total* radiation quantities (quantities integrated over all wavelengths), and made no reference to wavelength dependence. This lumped approach is adequate for many radiation problems encountered in practice. But sometimes it is necessary to consider the variation of radiation with wavelength as well as direction, and to express quantities at a certain wavelength  $\lambda$  or per unit wavelength interval about  $\lambda$ . Such quantities are referred to as *spectral* quantities to draw attention to wavelength dependence. The modifier "spectral" is used to indicate "at a given wavelength."

The spectral radiation intensity  $I_{\lambda}(\lambda, \theta, \phi)$ , for example, is simply the total radiation intensity  $I(\theta, \phi)$  per unit wavelength interval about  $\lambda$ . The **spectral intensity** for emitted radiation  $I_{\lambda, e}(\lambda, \theta, \phi)$  can be defined as the rate at which radiation energy  $d\dot{Q}_e$  is emitted at the wavelength  $\lambda$  in the  $(\theta, \phi)$  direction per unit area normal to this direction, per unit solid angle about this direction, and it can be expressed as

$$I_{\lambda, e}(\lambda, \theta, \phi) = \frac{d\dot{Q}_e}{dA\cos\theta \cdot d\omega \cdot d\lambda} \qquad (W/m^2 \cdot \text{sr} \cdot \mu m)$$
(11-23)



The three kinds of radiation flux (in W/m<sup>2</sup>): emissive power, irradiation, and radiosity. HEAT TRANSFER



Integration of a "spectral" quantity for all wavelengths gives the "total" quantity.

Then the spectral emissive power becomes

$$E_{\lambda} = \int_{\Phi=0}^{2\pi} \int_{\theta=0}^{\pi/2} I_{\lambda,e}(\lambda,\theta,\phi) \cos\theta \sin\theta d\theta d\phi \qquad (W/m^2)$$
(11-24)

Similar relations can be obtained for spectral irradiation  $G_{\lambda}$ , and spectral radiosity  $J_{\lambda}$  by replacing  $I_{\lambda, e}$  in this equation by  $I_{\lambda, i}$  and  $I_{\lambda, e+r}$  respectively.

When the variation of spectral radiation intensity  $I_{\lambda}$  with wavelength  $\lambda$  is known, the total radiation intensity *I* for emitted, incident, and emitted + reflected radiation can be determined by integration over the entire wavelength spectrum as (Fig. 11–22)

$$I_e = \int_0^\infty I_{\lambda, e} d\lambda, \qquad I_i = \int_0^\infty I_{\lambda, i} d\lambda, \qquad \text{and} \qquad I_{e+r} = \int_0^\infty I_{\lambda, e+r} d\lambda \qquad (11-25)$$

These intensities can then be used in Eqs. 11-15, 11-19, and 11-21 to determine the emissive power *E*, irradiation *G*, and radiosity *J*, respectively.

Similarly, when the variations of spectral radiation fluxes  $E_{\lambda}$ ,  $G_{\lambda}$ , and  $J_{\lambda}$  with wavelength  $\lambda$  are known, the total radiation fluxes can be determined by integration over the entire wavelength spectrum as

$$E = \int_0^\infty E_\lambda d\lambda, \qquad G = \int_0^\infty G_\lambda d\lambda, \text{ and } J = \int_0^\infty J_\lambda d\lambda$$
 (11-26)

When the surfaces and the incident radiation are *diffuse*, the spectral radiation fluxes are related to spectral intensities as

$$E_{\lambda} = \pi I_{\lambda,e}, \qquad G_{\lambda} = \pi I_{\lambda,i}, \qquad \text{and} \qquad J_{\lambda} = \pi I_{\lambda,e+r}$$
(11-27)

Note that the relations for spectral and total radiation quantities are of the same form.

The spectral intensity of radiation emitted by a blackbody at an absolute temperature *T* at a wavelength  $\lambda$  has been determined by Max Planck, and is expressed as

$$I_{b\lambda}(\lambda, T) = \frac{2hc_0^2}{\lambda^5 [\exp(hc_0/\lambda kT) - 1]} \qquad (W/m^2 \cdot \mathrm{sr} \cdot \mu m) \tag{11-28}$$

where  $h = 6.6256 \times 10^{-34} \text{ J} \cdot \text{s}$  is the Planck constant,  $k = 1.38065 \times 10^{-23}$  J/K is the Boltzmann constant, and  $c_0 = 2.9979 \times 10^8$  m/s is the speed of light in a vacuum. Then the spectral blackbody emissive power is, from Eq. 11-27,

$$E_{b\lambda}(\lambda, T) = \pi I_{b\lambda}(\lambda, T)$$
(11-29)

A simplified relation for  $E_{b\lambda}$  is given by Eq. 11-4.

#### **EXAMPLE 11–3** Radiation Incident on a Small Surface

A small surface of area  $A_1 = 3 \text{ cm}^2$  emits radiation as a blackbody at  $T_1 = 600$  K. Part of the radiation emitted by  $A_1$  strikes another small surface of area  $A_2 = 5 \text{ cm}^2$  oriented as shown in Figure 11–23. Determine the solid angle subtended by  $A_2$  when viewed from  $A_1$ , and the rate at which radiation emitted by  $A_1$  that strikes  $A_2$ .



FIGURE 11–23 Schematic for Example 11–3. **SOLUTION** A surface is subjected to radiation emitted by another surface. The solid angle subtended and the rate at which emitted radiation is received are to be determined.

**Assumptions** 1 Surface  $A_1$  emits diffusely as a blackbody. **2** Both  $A_1$  and  $A_2$  can be approximated as differential surfaces since both are very small compared to the square of the distance between them.

**Analysis** Approximating both  $A_1$  and  $A_2$  as differential surfaces, the solid angle subtended by  $A_2$  when viewed from  $A_1$  can be determined from Eq. 11-12 to be

$$\omega_{2-1} \cong \frac{A_{n,2}}{r^2} = \frac{A_2 \cos \theta_2}{r^2} = \frac{(5 \text{ cm}^2) \cos 40^\circ}{(75 \text{ cm})^2} = 6.81 \times 10^{-4} \text{ sr}$$

since the normal of  $A_2$  makes 40° with the direction of viewing. Note that solid angle subtended by  $A_2$  would be maximum if  $A_2$  were positioned normal to the direction of viewing. Also, the point of viewing on  $A_1$  is taken to be a point in the middle, but it can be any point since  $A_1$  is assumed to be very small.

The radiation emitted by  $A_1$  that strikes  $A_2$  is equivalent to the radiation emitted by  $A_1$  through the solid angle  $\omega_{2-1}$ . The intensity of the radiation emitted by  $A_1$  is

$$I_1 = \frac{E_h(T_1)}{\pi} = \frac{\sigma T_1^4}{\pi} = \frac{(5.67 \times 10^{-8} \text{ W/m}^2 \cdot \text{K}^4)(600 \text{ K})^4}{\pi} = 2339 \text{ W/m}^2 \cdot \text{sr}$$

This value of intensity is the same in all directions since a blackbody is a diffuse emitter. Intensity represents the rate of radiation emission per unit area normal to the direction of emission per unit solid angle. Therefore, the rate of radiation energy emitted by  $A_1$  in the direction of  $\theta_1$  through the solid angle  $\omega_{2-1}$  is determined by multiplying  $I_1$  by the area of  $A_1$  normal to  $\theta_1$  and the solid angle  $\omega_{2-1}$ . That is,

 $\dot{Q}_{1-2} = I_1(A_1 \cos \theta_1)\omega_{2-1}$ = (2339 W/m<sup>2</sup> · sr)(3 × 10<sup>-4</sup> cos 55° m<sup>2</sup>)(6.81 × 10<sup>-4</sup> sr) = **2.74 × 10<sup>-4</sup> W** 

Therefore, the radiation emitted from surface  $A_1$  will strike surface  $A_2$  at a rate of 2.74  $\times$  10<sup>-4</sup> W.

**Discussion** The total rate of radiation emission from surface  $A_1$  is  $\dot{Q}_e = A_1 \sigma T_1^4 = 2.204$  W. Therefore, the fraction of emitted radiation that strikes  $A_2$  is 2.74 ×  $10^{-4}/2.204 = 0.00012$  (or 0.012 percent). Noting that the solid angle associated with a hemisphere is  $2\pi$ , the fraction of the solid angle subtended by  $A_2$  is 6.81 ×  $10^{-4}/(2\pi) = 0.000108$  (or 0.0108 percent), which is 0.9 times the fraction of emitted radiation. Therefore, the fraction of the solid angle a surface occupies does not represent the fraction of radiation energy the surface will receive even when the intensity of emitted radiation is constant. This is because radiation energy emitted by a surface in a given direction is proportional to the *projected area* of the surface in that direction, and reduces from a maximum at  $\theta = 0^\circ$  (the direction normal to surface) to zero at  $\theta = 90^\circ$  (the direction parallel to surface).

### **11–5 • RADIATIVE PROPERTIES**

Most materials encountered in practice, such as metals, wood, and bricks, are *opaque* to thermal radiation, and radiation is considered to be a *surface* 

*phenomenon* for such materials. That is, thermal radiation is emitted or absorbed within the first few microns of the surface, and thus we speak of radiative properties of *surfaces* for opaque materials.

Some other materials, such as glass and water, allow visible radiation to penetrate to considerable depths before any significant absorption takes place. Radiation through such *semitransparent* materials obviously cannot be considered to be a surface phenomenon since the entire volume of the material interacts with radiation. On the other hand, both glass and water are practically opaque to infrared radiation. Therefore, materials can exhibit different behavior at different wavelengths, and the dependence on wavelength is an important consideration in the study of radiative properties such as emissivity, absorptivity, reflectivity, and transmissivity of materials.

In the preceding section, we defined a *blackbody* as a perfect emitter and absorber of radiation and said that no body can emit more radiation than a blackbody at the same temperature. Therefore, a blackbody can serve as a convenient *reference* in describing the emission and absorption characteristics of real surfaces.

### **Emissivity**

The **emissivity** of a surface represents *the ratio of the radiation emitted by the surface at a given temperature to the radiation emitted by a blackbody at the same temperature*. The emissivity of a surface is denoted by  $\varepsilon$ , and it varies between zero and one,  $0 \le \varepsilon \le 1$ . Emissivity is a measure of how closely a surface approximates a blackbody, for which  $\varepsilon = 1$ .

The emissivity of a real surface is not a constant. Rather, it varies with the *temperature* of the surface as well as the *wavelength* and the *direction* of the emitted radiation. Therefore, different emissivities can be defined for a surface, depending on the effects considered. The most elemental emissivity of a surface at a given temperature is the **spectral directional emissivity**, which is defined as the ratio of the intensity of radiation emitted by the surface at a specified wavelength in a specified direction to the intensity of radiation emitted by a blackbody at the same temperature at the same wavelength. That is,

$$\varepsilon_{\lambda,\,\theta}(\lambda,\,\theta,\,\phi,\,T) = \frac{I_{\lambda,\,e}(\lambda,\,\theta,\,\phi,\,T)}{I_{b\lambda}(\lambda,\,T)}$$
(11-30)

where the subscripts  $\lambda$  and  $\theta$  are used to designate *spectral* and *directional* quantities, respectively. Note that blackbody radiation intensity is independent of direction, and thus it has no functional dependence on  $\theta$  and  $\phi$ .

The **total directional emissivity** is defined in a like manner by using total intensities (intensities integrated over all wavelengths) as

$$\varepsilon_{\theta}(\theta, \phi, T) = \frac{I_e(\theta, \phi, T)}{I_b(T)}$$
(11-31)

In practice, it is usually more convenient to work with radiation properties averaged over all directions, called *hemispherical properties*. Noting that the integral of the rate of radiation energy emitted at a specified wavelength per unit surface area over the entire hemisphere is *spectral emissive power*, the **spectral hemispherical emissivity** can be expressed as

$$\varepsilon_{\lambda}(\lambda, T) = \frac{E_{\lambda}(\lambda, T)}{E_{b\lambda}(\lambda, T)}$$
(11-32)

Note that the emissivity of a surface at a given wavelength can be different at different temperatures since the spectral distribution of emitted radiation (and thus the amount of radiation emitted at a given wavelength) changes with temperature.

Finally, the total hemispherical emissivity is defined in terms of the radiation energy emitted over all wavelengths in all directions as

$$\varepsilon(T) = \frac{E(T)}{E_b(T)}$$
(11-33)

Therefore, the total hemispherical emissivity (or simply the "average emissivity") of a surface at a given temperature represents the ratio of the total radiation energy emitted by the surface to the radiation emitted by a blackbody of the same surface area at the same temperature.

Noting from Eqs. 11-26 and 11-32 that  $E = \int_0^\infty E_\lambda d\lambda$  and  $E_\lambda(\lambda, T) =$  $\varepsilon_{\lambda}(\lambda, T)E_{h\lambda}(\lambda, T)$ , and the total hemispherical emissivity can also be expressed as

$$\varepsilon(T) = \frac{E(T)}{E_b(T)} = \frac{\int_0^\infty \varepsilon_\lambda(\lambda, T) E_{b\lambda}(\lambda, T) d\lambda}{\sigma T^4}$$
(11-34)

since  $E_b(T) = \sigma T^4$ . To perform this integration, we need to know the variation of spectral emissivity with wavelength at the specified temperature. The integrand is usually a complicated function, and the integration has to be performed numerically. However, the integration can be performed quite easily by dividing the spectrum into a sufficient number of wavelength bands and assuming the emissivity to remain constant over each band; that is, by expressing the function  $\varepsilon_{\lambda}(\lambda, T)$  as a step function. This simplification offers great convenience for little sacrifice of accuracy, since it allows us to transform the integration into a summation in terms of blackbody emission functions.

As an example, consider the emissivity function plotted in Figure 11–24. It seems like this function can be approximated reasonably well by a step function of the form

$$\varepsilon_{\lambda} = \begin{cases} \varepsilon_{1} = \text{constant}, & 0 \leq \lambda < \lambda_{1} \\ \varepsilon_{2} = \text{constant}, & \lambda_{1} \leq \lambda < \lambda_{2} \\ \varepsilon_{3} = \text{constant}, & \lambda_{2} \leq \lambda < \infty \end{cases}$$
(11-35)

Then the average emissivity can be determined from Eq. 11-34 by breaking the integral into three parts and utilizing the definition of the blackbody radiation function as

$$\varepsilon(T) = \frac{\varepsilon_1 \int_0^{\lambda_1} E_{b\lambda} d\lambda}{E_b} + \frac{\varepsilon_2 \int_{\lambda_1}^{\lambda_2} E_{b\lambda} d\lambda}{E_b} + \frac{\varepsilon_3 \int_{\lambda_2}^{\infty} E_{b\lambda} d\lambda}{E_b}$$
$$= \varepsilon_1 f_{0-\lambda_1}(T) + \varepsilon_2 f_{\lambda_1-\lambda_2}(T) + \varepsilon_3 f_{\lambda_2-\infty}(T)$$
(11)

Radiation is a complex phenomenon as it is, and the consideration of the wavelength and direction dependence of properties, assuming sufficient data



-36)

ε

Approximating the actual variation of emissivity with wavelength by a step function.

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```
Real surface:

\epsilon_{\theta} \neq \text{constant}

\epsilon_{\lambda} \neq \text{constant}

Diffuse surface:

\epsilon_{\theta} = \text{constant}

Gray surface:

\epsilon_{\lambda} = \text{constant}

Diffuse, gray surface:

\epsilon = \epsilon_{\lambda} = \epsilon_{\theta} = \text{constant}
```

#### **FIGURE 11–25**

The effect of diffuse and gray approximations on the emissivity of a surface.



### FIGURE 11–26

Typical variations of emissivity with direction for electrical conductors and nonconductors.

exist, makes it even more complicated. Therefore, the *gray* and *diffuse* approximations are often utilized in radiation calculations. A surface is said to be *diffuse* if its properties are *independent of direction*, and *gray* if its properties are *independent of wavelength*. Therefore, the emissivity of a gray, diffuse surface is simply the total hemispherical emissivity of that surface because of independence of direction and wavelength (Fig. 11–25).

A few comments about the validity of the diffuse approximation are in order. Although real surfaces do not emit radiation in a perfectly diffuse manner as a blackbody does, they often come close. The variation of emissivity with direction for both electrical conductors and nonconductors is given in Figure 11–26. Here  $\theta$  is the angle measured from the normal of the surface, and thus  $\theta = 0$  for radiation emitted in a direction normal to the surface. Note that  $\varepsilon_{\theta}$  remains nearly constant for about  $\theta < 40^{\circ}$  for conductors such as metals and for  $\theta < 70^{\circ}$  for nonconductors such as plastics. Therefore, the directional emissivity of a surface in the normal direction is representative of the hemispherical emissivity of the surface. In radiation analysis, it is common practice to assume the surfaces to be diffuse emitters with an emissivity equal to the value in the normal ( $\theta = 0$ ) direction.

The effect of the gray approximation on emissivity and emissive power of a real surface is illustrated in Figure 11–27. Note that the radiation emission from a real surface, in general, differs from the Planck distribution, and the emission curve may have several peaks and valleys. A gray surface should emit as much radiation as the real surface it represents at the same temperature. Therefore, the areas under the emission curves of the real and gray surfaces must be equal.

The emissivities of common materials are listed in Table A–18 in the appendix, and the variation of emissivity with wavelength and temperature is illustrated in Figure 11–28. Typical ranges of emissivity of various materials are given in Figure 11–29. Note that metals generally have low emissivities, as low as 0.02 for polished surfaces, and nonmetals such as ceramics and organic materials have high ones. The emissivity of metals increases with temperature. Also, oxidation causes significant increases in the emissivity of metals. Heavily oxidized metals can have emissivities comparable to those of nonmetals.



### FIGURE 11-27

Comparison of the emissivity (*a*) and emissive power (*b*) of a real surface with those of a gray surface and a blackbody at the same temperature.

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The variation of normal emissivity with (a) wavelength and (b) temperature for various materials.

Care should be exercised in the use and interpretation of radiation property data reported in the literature, since the properties strongly depend on the surface conditions such as oxidation, roughness, type of finish, and cleanliness. Consequently, there is considerable discrepancy and uncertainty in the reported values. This uncertainty is largely due to the difficulty in characterizing and describing the surface conditions precisely.

#### **EXAMPLE 11–4** Emissivity of a Surface and Emissive Power

The spectral emissivity function of an opaque surface at 800 K is approximated as (Fig. 11–30)

$$\varepsilon_{\lambda} = \begin{cases} \varepsilon_1 = 0.3, & 0 \le \lambda < 3 \ \mu m \\ \varepsilon_2 = 0.8, & 3 \ \mu m \le \lambda < 7 \ \mu m \\ \varepsilon_3 = 0.1, & 7 \ \mu m \le \lambda < \infty \end{cases}$$

Determine the average emissivity of the surface and its emissive power.

**SOLUTION** The variation of emissivity of a surface at a specified temperature with wavelength is given. The average emissivity of the surface and its emissive power are to be determined.

*Analysis* The variation of the emissivity of the surface with wavelength is given as a step function. Therefore, the average emissivity of the surface can be determined from Eq. 11-34 by breaking the integral into three parts,

$$\varepsilon(T) = \frac{\varepsilon_1 \int_0^{\lambda_1} E_{b\lambda} d\lambda}{\sigma T^4} + \frac{\varepsilon_2 \int_{\lambda_1}^{\lambda_2} E_{b\lambda} d\lambda}{\sigma T^4} + \frac{\varepsilon_3 \int_{\lambda_2}^{\infty} E_{b\lambda} d\lambda}{\sigma T^4}$$
$$= \varepsilon_1 f_{0-\lambda_1}(T) + \varepsilon_2 f_{\lambda_1 - \lambda_2}(T) + \varepsilon_3 f_{\lambda_2 - \infty}(T)$$
$$= \varepsilon_1 f_{\lambda_1} + \varepsilon_2 (f_{\lambda_2} - f_{\lambda_1}) + \varepsilon_3 (1 - f_{\lambda_2})$$



Typical ranges of emissivity for various materials.



The spectral emissivity of the surface considered in Example 11–4.

where  $f_{\lambda_1}$  and  $f_{\lambda_2}$  are blackbody radiation functions corresponding to  $\lambda_1 T$  and  $\lambda_2 T$ . These functions are determined from Table 11–2 to be

 $\lambda_1 T = (3 \ \mu m)(800 \ K) = 2400 \ \mu m \cdot K \rightarrow f_{\lambda_1} = 0.140256$  $\lambda_2 T = (7 \ \mu m)(800 \ K) = 5600 \ \mu m \cdot K \rightarrow f_{\lambda_2} = 0.701046$ 

Note that  $f_{0-\lambda_1} = f_{\lambda_1} - f_0 = f_{\lambda_1}$ , since  $f_0 = 0$ , and  $f_{\lambda_2-\infty} = f_{\infty} - f_{\lambda_2} = 1 - f_{\lambda_2}$ , since  $f_{\infty} = 1$ . Substituting,

$$\varepsilon = 0.3 \times 0.140256 + 0.8(0.701046 - 0.140256) + 0.1(1 - 0.701046)$$
  
= 0.521

That is, the surface will emit as much radiation energy at 800 K as a gray surface having a constant emissivity of  $\varepsilon = 0.521$ . The emissive power of the surface is

 $E = \varepsilon \sigma T^4 = 0.521(5.67 \times 10^{-8} \text{ W/m}^2 \cdot \text{K}^4)(800 \text{ K})^4 = 12,100 \text{ W/m}^2$ 

**Discussion** Note that the surface emits 12.1 kJ of radiation energy per second per  $m^2$  area of the surface.

### Absorptivity, Reflectivity, and Transmissivity

Everything around us constantly emits radiation, and the emissivity represents the emission characteristics of those bodies. This means that every body, including our own, is constantly bombarded by radiation coming from all directions over a range of wavelengths. Recall that radiation flux *incident on a surface* is called **irradiation** and is denoted by *G*.

When radiation strikes a surface, part of it is absorbed, part of it is reflected, and the remaining part, if any, is transmitted, as illustrated in Figure 11–31. *The fraction of irradiation absorbed by the surface* is called the **absorptivity**  $\alpha$ , *the fraction reflected by the surface* is called the **reflectivity**  $\rho$ , and *the fraction transmitted* is called the **transmissivity**  $\tau$ . That is,

Absorptivity:	$\alpha = \frac{\text{Absorbed radiation}}{\text{Incident radiation}} = \frac{G}{G}$	$\frac{abs}{7}$ ,	$0 \le \alpha \le 1$	(11-37)
		J		

Reflectivity: 
$$\rho = \frac{\text{Reflected radiation}}{\text{Incident radiation}} = \frac{G_{\text{ref}}}{G}, \qquad 0 \le \rho \le 1$$
 (11-38)

Transmissivity: 
$$\tau = \frac{\text{Transmitted radiation}}{\text{Incident radiation}} = \frac{G_{\text{tr}}}{G}, \quad 0 \le \tau \le 1$$
 (11-39)

where G is the radiation energy incident on the surface, and  $G_{abs}$ ,  $G_{ref}$ , and  $G_{tr}$  are the absorbed, reflected, and transmitted portions of it, respectively. The first law of thermodynamics requires that the sum of the absorbed, reflected, and transmitted radiation energy be equal to the incident radiation. That is,

$$G_{\rm abs} + G_{\rm ref} + G_{\rm tr} = G$$
 (11-40)

Dividing each term of this relation by G yields

$$\alpha + \rho + \tau = 1 \tag{11-41}$$

For opaque surfaces,  $\tau = 0$ , and thus



#### FIGURE 11-31

The absorption, reflection, and transmission of incident radiation by a semitransparent material.

 $\alpha + \rho = 1$ 

(11-42)

This is an important property relation since it enables us to determine both the absorptivity and reflectivity of an opaque surface by measuring either of these properties.

These definitions are for *total hemispherical* properties, since *G* represents the radiation flux incident on the surface from all directions over the hemispherical space and over all wavelengths. Thus,  $\alpha$ ,  $\rho$ , and  $\tau$  are the *average* properties of a medium for all directions and all wavelengths. However, like emissivity, these properties can also be defined for a specific wavelength and/or direction. For example, the **spectral directional absorptivity** and **spectral directional reflectivity** of a surface are defined, respectively, as the absorbed and reflected fractions of the intensity of radiation incident at a specified wavelength in a specified direction as

$$\alpha_{\lambda,\,\theta}(\lambda,\,\theta,\,\phi) = \frac{I_{\lambda,\,abs}(\lambda,\,\theta,\,\phi)}{I_{\lambda,\,i}(\lambda,\,\theta,\,\phi)} \quad \text{and} \quad \rho_{\lambda,\,\theta}(\lambda,\,\theta,\,\phi) = \frac{I_{\lambda,\,ref}(\lambda,\,\theta,\,\phi)}{I_{\lambda,\,i}(\lambda,\,\theta,\,\phi)} \quad (11-43)$$

Likewise, the **spectral hemispherical absorptivity** and **spectral hemispherical reflectivity** of a surface are defined as

$$\alpha_{\lambda}(\lambda) = \frac{G_{\lambda, \text{ abs}}(\lambda)}{G_{\lambda}(\lambda)} \quad \text{and} \quad \rho_{\lambda}(\lambda) = \frac{G_{\lambda, \text{ ref}}(\lambda)}{G_{\lambda}(\lambda)}$$
(11-44)

where  $G_{\lambda}$  is the spectral irradiation (in W/m<sup>2</sup> ·  $\mu$ m) incident on the surface, and  $G_{\lambda, abs}$  and  $G_{\lambda, ref}$  are the reflected and absorbed portions of it, respectively.

Similar quantities can be defined for the transmissivity of semitransparent materials. For example, the **spectral hemispherical transmissivity** of a medium can be expressed as

$$\tau_{\lambda}(\lambda) = \frac{G_{\lambda, \text{tr}}(\lambda)}{G_{\lambda}(\lambda)}$$
(11-45)

The average absorptivity, reflectivity, and transmissivity of a surface can also be defined in terms of their spectral counterparts as

$$\alpha = \frac{\int_0^\infty \alpha_\lambda G_\lambda d\lambda}{\int_0^\infty G_\lambda d\lambda}, \qquad \rho = \frac{\int_0^\infty \rho_\lambda G_\lambda d\lambda}{\int_0^\infty G_\lambda d\lambda}, \qquad \tau = \frac{\int_0^\infty \tau_\lambda G_\lambda d\lambda}{\int_0^\infty G_\lambda d\lambda}$$
(11-46)

The reflectivity differs somewhat from the other properties in that it is *bidirectional* in nature. That is, the value of the reflectivity of a surface depends not only on the direction of the incident radiation but also the direction of reflection. Therefore, the reflected rays of a radiation beam incident on a real surface in a specified direction will form an irregular shape, as shown in Figure 11–32. Such detailed reflectivity data do not exist for most surfaces, and even if they did, they would be of little value in radiation calculations since this would usually add more complication to the analysis than it is worth.

In practice, for simplicity, surfaces are assumed to reflect in a perfectly *specular* or *diffuse* manner. In **specular** (or *mirrorlike*) **reflection**, *the angle of reflection equals the angle of incidence of the radiation beam*. In **diffuse reflection**, *radiation is reflected equally in all directions*, as shown in Figure









### FIGURE 11–33

Variation of absorptivity with the temperature of the source of irradiation for various common materials at room temperature.



**FIGURE 11–34** 

The absorptivity of a material may be quite different for radiation originating from sources at different temperatures. 11–32. Reflection from smooth and polished surfaces approximates specular reflection, whereas reflection from rough surfaces approximates diffuse reflection. In radiation analysis, smoothness is defined relative to wavelength. A surface is said to be *smooth* if the height of the surface roughness is much smaller than the wavelength of the incident radiation.

Unlike emissivity, the absorptivity of a material is practically independent of surface temperature. However, the absorptivity depends strongly on the temperature of the source at which the incident radiation is originating. This is also evident from Figure 11–33, which shows the absorptivities of various materials at room temperature as functions of the temperature of the radiation source. For example, the absorptivity of the concrete roof of a house is about 0.6 for solar radiation (source temperature: 5780 K) and 0.9 for radiation originating from the surrounding trees and buildings (source temperature: 300 K), as illustrated in Figure 11–34.

Notice that the absorptivity of aluminum increases with the source temperature, a characteristic for metals, and the absorptivity of electric nonconductors, in general, decreases with temperature. This decrease is most pronounced for surfaces that appear white to the eye. For example, the absorptivity of a white painted surface is low for solar radiation, but it is rather high for infrared radiation.

### **Kirchhoff's Law**

Consider a small body of surface area  $A_s$ , emissivity  $\varepsilon$ , and absorptivity  $\alpha$  at temperature *T* contained in a large isothermal enclosure at the same temperature, as shown in Figure 11–35. Recall that a large isothermal enclosure forms a blackbody cavity regardless of the radiative properties of the enclosure surface, and the body in the enclosure is too small to interfere with the blackbody nature of the cavity. Therefore, the radiation incident on any part of the surface of the small body is equal to the radiation emitted by a blackbody at temperature *T*. That is,  $G = E_b(T) = \sigma T^4$ , and the radiation absorbed by the small body per unit of its surface area is

$$G_{abs} = \alpha G = \alpha \sigma T^4$$

(

The radiation emitted by the small body is

$$E_{\rm emit} = \varepsilon \sigma T^4$$

Considering that the small body is in thermal equilibrium with the enclosure, the net rate of heat transfer to the body must be zero. Therefore, the radiation emitted by the body must be equal to the radiation absorbed by it:

$$A_s \varepsilon \sigma T^4 = A_s \alpha \sigma T^4$$

Thus, we conclude that

$$\varepsilon(T) = \alpha(T) \tag{11-47}$$

That is, the total hemispherical emissivity of a surface at temperature T is equal to its total hemispherical absorptivity for radiation coming from a blackbody at the same temperature. This relation, which greatly simplifies the radiation analysis, was first developed by Gustav Kirchhoff in 1860 and is now called **Kirchhoff's law.** Note that this relation is derived under the condition that the surface temperature is equal to the temperature of the source of irradiation, and the reader is cautioned against using it when considerable difference (more than a few hundred degrees) exists between the surface temperature and the temperature of the source of irradiation.

The derivation above can also be repeated for radiation at a specified wavelength to obtain the *spectral* form of Kirchhoff's law:

$$\varepsilon_{\lambda}(T) = \alpha_{\lambda}(T)$$
 (11-48)

This relation is valid when the irradiation or the emitted radiation is independent of direction. The form of Kirchhoff's law that involves no restrictions is the *spectral directional* form expressed as  $\varepsilon_{\lambda, \theta}(T) = \alpha_{\lambda, \theta}(T)$ . That is, the emissivity of a surface at a specified wavelength, direction, and temperature is always equal to its absorptivity at the same wavelength, direction, and temperature.

It is very tempting to use Kirchhoff's law in radiation analysis since the relation  $\varepsilon = \alpha$  together with  $\rho = 1 - \alpha$  enables us to determine all three properties of an opaque surface from a knowledge of only *one* property. Although Eq. 11-47 gives acceptable results in most cases, in practice, care should be exercised when there is considerable difference between the surface temperature and the temperature of the source of incident radiation.

### The Greenhouse Effect

You have probably noticed that when you leave your car under direct sunlight on a sunny day, the interior of the car gets much warmer than the air outside, and you may have wondered why the car acts like a *heat trap*. The answer lies in the spectral transmissivity curve of the glass, which resembles an inverted U, as shown in Figure 11-36. We observe from this figure that glass at thicknesses encountered in practice transmits over 90 percent of radiation in the visible range and is practically opaque (nontransparent) to radiation in the longer-wavelength infrared regions of the electromagnetic spectrum (roughly  $\lambda > 3 \mu m$ ). Therefore, glass has a transparent window in the wavelength range 0.3  $\mu$ m  $< \lambda < 3 \mu$ m in which over 90 percent of solar radiation is emitted. On the other hand, the entire radiation emitted by surfaces at room temperature falls in the infrared region. Consequently, glass allows the solar radiation to enter but does not allow the infrared radiation from the interior surfaces to escape. This causes a rise in the interior temperature as a result of the energy build-up in the car. This heating effect, which is due to the nongray characteristic of glass (or clear plastics), is known as the greenhouse effect, since it is utilized primarily in greenhouses (Fig. 11–37).

The greenhouse effect is also experienced on a larger scale on earth. The surface of the earth, which warms up during the day as a result of the absorption of solar energy, cools down at night by radiating its energy into deep space as infrared radiation. The combustion gases such as  $CO_2$  and water vapor in the atmosphere transmit the bulk of the solar radiation but absorb the infrared radiation emitted by the surface of the earth. Thus, there is concern that the energy trapped on earth will eventually cause global warming and thus drastic changes in weather patterns.

In *humid* places such as coastal areas, there is not a large change between the daytime and nighttime temperatures, because the humidity acts as a barrier

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FIGURE 11–35

The small body contained in a large isothermal enclosure used in the development of Kirchhoff's law.



#### FIGURE 11–36

The spectral transmissivity of low-iron glass at room temperature for different thicknesses.



### FIGURE 11–37

A greenhouse traps energy by allowing the solar radiation to come in but not allowing the infrared radiation to go out.





Solar radiation reaching the earth's atmosphere and the total solar irradiance.



#### FIGURE 11–39

The total solar energy passing through concentric spheres remains constant, but the energy falling per unit area decreases with increasing radius. on the path of the infrared radiation coming from the earth, and thus slows down the cooling process at night. In areas with clear skies such as deserts, there is a large swing between the daytime and nighttime temperatures because of the absence of such barriers for infrared radiation.

### 11–6 • ATMOSPHERIC AND SOLAR RADIATION

The sun is our primary source of energy. The energy coming off the sun, called *solar energy*, reaches us in the form of electromagnetic waves after experiencing considerable interactions with the atmosphere. The radiation energy emitted or reflected by the constituents of the atmosphere form the *atmospheric radiation*. Below we give an overview of the solar and atmospheric radiation because of their importance and relevance to daily life. Also, our familiarity with solar energy makes it an effective tool in developing a better understanding for some of the new concepts introduced earlier. Detailed treatment of this exciting subject can be found in numerous books devoted to this topic.

The *sun* is a nearly spherical body that has a diameter of  $D \approx 1.39 \times 10^9$  m and a mass of  $m \approx 2 \times 10^{30}$  kg and is located at a mean distance of  $L = 1.50 \times 10^{11}$  m from the earth. It emits radiation energy continuously at a rate of  $E_{\rm sun} \approx 3.8 \times 10^{26}$  W. Less than a billionth of this energy (about  $1.7 \times 10^{17}$  W) strikes the earth, which is sufficient to keep the earth warm and to maintain life through the photosynthesis process. The energy of the sun is due to the continuous *fusion* reaction during which two hydrogen atoms fuse to form one atom of helium. Therefore, the sun is essentially a *nuclear reactor*, with temperatures as high as 40,000,000 K in its core region. The temperature drops to about 5800 K in the outer region of the sun, called the convective zone, as a result of the dissipation of this energy by radiation.

The solar energy reaching the earth's atmosphere is called the **total solar** irradiance  $G_s$ , whose value is

$$G_s = 1373 \text{ W/m}^2$$
 (11-49)

The total solar irradiance (also called the **solar constant**) represents *the rate at which solar energy is incident on a surface normal to the sun's rays at the outer edge of the atmosphere when the earth is at its mean distance from the sun* (Fig. 11–38).

The value of the total solar irradiance can be used to estimate the effective surface temperature of the sun from the requirement that

$$(4\pi L^2)G_s = (4\pi r^2)\,\sigma T_{\rm sun}^4\tag{11-50}$$

where *L* is the mean distance between the sun's center and the earth and *r* is the radius of the sun. The left-hand side of this equation represents the total solar energy passing through a spherical surface whose radius is the mean earth–sun distance, and the right-hand side represents the total energy that leaves the sun's outer surface. The conservation of energy principle requires that these two quantities be equal to each other, since the solar energy experiences no attenuation (or enhancement) on its way through the vacuum (Fig. 11–39). The **effective surface temperature** of the sun is determined from Eq. 11-50 to be  $T_{sun} = 5780$  K. That is, the sun can be treated as a

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blackbody at a temperature of 5780 K. This is also confirmed by the measurements of the spectral distribution of the solar radiation just outside the atmosphere plotted in Figure 11–40, which shows only small deviations from the idealized blackbody behavior.

The spectral distribution of solar radiation on the ground plotted in Figure 11–40 shows that the solar radiation undergoes considerable attenuation as it passes through the atmosphere as a result of absorption and scattering. About 99 percent of the atmosphere is contained within a distance of 30 km from the earth's surface. The several dips on the spectral distribution of radiation on the earth's surface are due to *absorption* by the gases  $O_2$ ,  $O_3$  (ozone),  $H_2O$ , and  $CO_2$ . Absorption by *oxygen* occurs in a narrow band about  $\lambda = 0.76 \ \mu\text{m}$ . The *ozone* absorbs *ultraviolet* radiation at wavelengths below 0.3  $\mu$ m almost completely, and radiation in the range 0.3–0.4  $\mu$ m considerably. Thus, the ozone layer in the upper regions of the atmosphere protects biological systems on earth from harmful ultraviolet radiation. In turn, we must protect the ozone layer from the destructive chemicals commonly used as refrigerants, cleaning agents, and propellants in aerosol cans. The use of these chemicals is now banned in many countries. The ozone gas also absorbs some radiation in the visible range. Absorption in the infrared region is dominated by water vapor and carbon dioxide. The dust particles and other pollutants in the atmosphere also absorb radiation at various wavelengths.

As a result of these absorptions, the solar energy reaching the *earth's sur-face* is weakened considerably, to about 950 W/m<sup>2</sup> on a clear day and much less on cloudy or smoggy days. Also, practically all of the solar radiation reaching the earth's surface falls in the wavelength band from 0.3 to 2.5  $\mu$ m.

Another mechanism that attenuates solar radiation as it passes through the atmosphere is scattering or reflection by air molecules and the many other kinds of particles such as dust, smog, and water droplets suspended in the atmosphere. Scattering is mainly governed by the size of the particle relative to the wavelength of radiation. The oxygen and nitrogen molecules primarily scatter radiation at very short wavelengths, comparable to the size of the molecules themselves. Therefore, radiation at wavelengths corresponding to violet and blue colors is scattered the most. This molecular scattering in all directions is what gives the sky its bluish color. The same phenomenon is responsible for red sunrises and sunsets. Early in the morning and late in the afternoon, the sun's rays pass through a greater thickness of the atmosphere than they do at midday, when the sun is at the top. Therefore, the violet and blue colors of the light encounter a greater number of molecules by the time they reach the earth's surface, and thus a greater fraction of them are scattered (Fig. 11–41). Consequently, the light that reaches the earth's surface consists primarily of colors corresponding to longer wavelengths such as red, orange, and yellow. The clouds appear in reddish-orange color during sunrise and sunset because the light they reflect is reddish-orange at those times. For the same reason, a red traffic light is visible from a longer distance than is a green light under the same circumstances.

The solar energy incident on a surface on earth is considered to consist of *direct* and *diffuse* parts. The part of solar radiation that reaches the earth's surface without being scattered or absorbed by the atmosphere is called **direct** solar radiation  $G_D$ . The scattered radiation is assumed to reach the earth's surface uniformly from all directions and is called **diffuse solar radiation**  $G_d$ .



#### FIGURE 11-40

Spectral distribution of solar radiation just outside the atmosphere, at the surface of the earth on a typical day, and comparison with blackbody radiation at 5780 K.



### FIGURE 11-41

Air molecules scatter blue light much more than they do red light. At sunset, the light travels through a thicker layer of atmosphere, which removes much of the blue from the natural light, allowing the red to dominate.

### HEAT TRANSFER



FIGURE 11–42 The direct and diffuse radiation incident on a horizontal surface at the earth's surface.



### FIGURE 11-43

Radiation interactions of a surface exposed to solar and atmospheric radiation.

Then the *total solar energy* incident on the unit area of a *horizontal surface* on the ground is (Fig. 11–42)

$$G_{\text{solar}} = G_D \cos \theta + G_d \qquad (W/m^2) \tag{11-51}$$

where  $\theta$  is the angle of incidence of direct solar radiation (the angle that the sun's rays make with the normal of the surface). The diffuse radiation varies from about 10 percent of the total radiation on a clear day to nearly 100 percent on a totally cloudy day.

The gas molecules and the suspended particles in the atmosphere *emit* radiation as well as absorbing it. The atmospheric emission is primarily due to the CO<sub>2</sub> and H<sub>2</sub>O molecules and is concentrated in the regions from 5 to 8  $\mu$ m and above 13  $\mu$ m. Although this emission is far from resembling the distribution of radiation from a blackbody, it is found convenient in radiation calculations to treat the atmosphere as a blackbody at some lower fictitious temperature that emits an equivalent amount of radiation energy. This fictitious temperature is called the **effective sky temperature**  $T_{sky}$ . Then the radiation emission from the atmosphere to the earth's surface is expressed as

$$G_{\rm sky} = \sigma T_{\rm sky}^4$$
 (W/m<sup>2</sup>) (11-52)

The value of  $T_{sky}$  depends on the atmospheric conditions. It ranges from about 230 K for cold, clear-sky conditions to about 285 K for warm, cloudy-sky conditions.

Note that the effective sky temperature does not deviate much from the room temperature. Thus, in the light of Kirchhoff's law, we can take the absorptivity of a surface to be equal to its emissivity at room temperature,  $\alpha = \varepsilon$ . Then the sky radiation absorbed by a surface can be expressed as

$$E_{\rm sky, \, absorbed} = \alpha G_{\rm sky} = \alpha \sigma T_{\rm sky}^4 = \varepsilon \sigma T_{\rm sky}^4$$
 (W/m<sup>2</sup>) (11-53)

The net rate of radiation heat transfer to a surface exposed to solar and atmospheric radiation is determined from an energy balance (Fig. 11–43):

$$\dot{q}_{\text{net, rad}} = \sum E_{\text{absorbed}} - \sum E_{\text{emitted}}$$

$$= E_{\text{solar, absorbed}} + E_{\text{sky, absorbed}} - E_{\text{emitted}}$$

$$= \alpha_s G_{\text{solar}} + \varepsilon \sigma T_{\text{sky}}^4 - \varepsilon \sigma T_s^4$$

$$= \alpha_s G_{\text{solar}} + \varepsilon \sigma (T_{\text{sky}}^4 - T_s^4) \qquad (W/m^2) \qquad (11-54)$$

where  $T_s$  is the temperature of the surface in K and  $\varepsilon$  is its emissivity at room temperature. A positive result for  $\dot{q}_{\text{net, rad}}$  indicates a radiation heat gain by the surface and a negative result indicates a heat loss.

The absorption and emission of radiation by the *elementary gases* such as  $H_2$ ,  $O_2$ , and  $N_2$  at moderate temperatures are negligible, and a medium filled with these gases can be treated as a *vacuum* in radiation analysis. The absorption and emission of gases with *larger molecules* such as  $H_2O$  and  $CO_2$ , however, can be *significant* and may need to be considered when considerable amounts of such gases are present in a medium. For example, a 1-m-thick layer of water vapor at 1 atm pressure and 100°C emits more than 50 percent of the energy that a blackbody would emit at the same temperature.

In solar energy applications, the spectral distribution of incident solar radiation is very different than the spectral distribution of emitted radiation by the surfaces, since the former is concentrated in the short-wavelength region and the latter in the infrared region. Therefore, the radiation properties of surfaces will be quite different for the incident and emitted radiation, and the surfaces cannot be assumed to be gray. Instead, the surfaces are assumed to have two sets of properties: one for solar radiation and another for infrared radiation at room temperature. Table 11–3 lists the *emissivity*  $\varepsilon$  and the *solar absorptivity*  $\alpha_s$  of the surfaces of some common materials. Surfaces that are intended to *collect solar energy*, such as the absorber surfaces of solar collectors, are desired to have high  $\alpha_s$  but low  $\varepsilon$  values to maximize the absorption of solar radiation and to minimize the emission of radiation. Surfaces that are intended to *remain cool* under the sun, such as the outer surfaces of fuel tanks and refrigerator trucks, are desired to have just the opposite properties. Surfaces are often given the desired properties by coating them with thin layers of *selective* materials. A surface can be kept cool, for example, by simply painting it white.

We close this section by pointing out that what we call *renewable energy* is usually nothing more than the manifestation of solar energy in different forms. Such energy sources include wind energy, hydroelectric power, ocean thermal energy, ocean wave energy, and wood. For example, no hydroelectric power plant can generate electricity year after year unless the water evaporates by absorbing solar energy and comes back as a rainfall to replenish the water source (Fig. 11–44). Although solar energy is sufficient to meet the entire energy needs of the world, currently it is not economical to do so because of the low concentration of solar energy on earth and the high capital cost of harnessing it.

### **EXAMPLE 11–5** Selective Absorber and Reflective Surfaces

Consider a surface exposed to solar radiation. At a given time, the direct and diffuse components of solar radiation are  $G_D = 400$  and  $G_d = 300$  W/m<sup>2</sup>, and the direct radiation makes a 20° angle with the normal of the surface. The surface temperature is observed to be 320 K at that time. Assuming an effective sky temperature of 260 K, determine the net rate of radiation heat transfer for these cases (Fig. 11–45):

(a)  $\alpha_s = 0.9$  and  $\varepsilon = 0.9$  (gray absorber surface)

(b)  $\alpha_s = 0.1$  and  $\varepsilon = 0.1$  (gray reflector surface)

(c)  $\alpha_s = 0.9$  and  $\varepsilon = 0.1$  (selective absorber surface)

(d)  $\alpha_s = 0.1$  and  $\varepsilon = 0.9$  (selective reflector surface)

**SOLUTION** A surface is exposed to solar and sky radiation. The net rate of radiation heat transfer is to be determined for four different combinations of emissivities and solar absorptivities.

Analysis The total solar energy incident on the surface is

$$G_{\text{solar}} = G_D \cos \theta + G_d$$
  
= (400 W/m<sup>2</sup>) cos 20° + (300 W/m<sup>2</sup>)  
= 676 W/m<sup>2</sup>

Then the net rate of radiation heat transfer for each of the four cases is determined from:

$$\dot{q}_{\text{net, rad}} = \alpha_s G_{\text{solar}} + \varepsilon \sigma (T_{\text{sky}}^4 - T_s^4)$$

#### CHAPTER 11

### TABLE 11-3

Comparison of the solar absorptivity  $\alpha_s$  of some surfaces with their emissivity  $\epsilon$  at room temperature

Surface	αs	З
Aluminum		
Polished	0.09	0.03
Anodized	0.14	0.84
Foil	0.15	0.05
Copper		
Polished	0.18	0.03
Tarnished	0.65	0.75
Stainless steel		
Polished	0.37	0.60
Dull	0.50	0.21
Plated metals		
Black nickel oxide	0.92	0.08
Black chrome	0.87	0.09
Concrete	0.60	0.88
White marble	0.46	0.95
Red brick	0.63	0.93
Asphalt	0.90	0.90
Black paint	0.97	0.97
White paint	0.14	0.93
Snow	0.28	0.97
Human skin	0.00	0.07
(caucasian)	0.62	0.97



FIGURE 11–44 The cycle that water undergoes in a hydroelectric power plant.



### FIGURE 11–45

Graphical representation of the spectral emissivities of the four surfaces considered in Example 11–5.

(a)  $\alpha_s = 0.9$  and  $\varepsilon = 0.9$  (gray absorber surface):

 $\dot{q}_{\text{net, rad}} = 0.9(676 \text{ W/m}^2) + 0.9(5.67 \times 10^{-8} \text{ W/m}^2 \cdot \text{K}^4)[(260 \text{ K})^4 - (320 \text{ K})^4]$ = 307 W/m<sup>2</sup>

(b)  $\alpha_s = 0.1$  and  $\varepsilon = 0.1$  (gray reflector surface):

 $\dot{q}_{\text{net, rad}} = 0.1(676 \text{ W/m}^2) + 0.1(5.67 \times 10^{-8} \text{ W/m}^2 \cdot \text{K}^4)[(260 \text{ K})^4 - (320 \text{ K})^4]$ = 34 W/m<sup>2</sup>

(c)  $\alpha_s = 0.9$  and  $\varepsilon = 0.1$  (selective absorber surface):

 $\dot{q}_{\text{net, rad}} = 0.9(676 \text{ W/m}^2) + 0.1(5.67 \times 10^{-8} \text{ W/m}^2 \cdot \text{K}^4)[(260 \text{ K})^4 - (320 \text{ K})^4]$ = 575 W/m<sup>2</sup>

(d)  $\alpha_s = 0.1$  and  $\varepsilon = 0.9$  (selective reflector surface):

 $\dot{q}_{\text{net, rad}} = 0.1(676 \text{ W/m}^2) + 0.9(5.67 \times 10^{-8} \text{ W/m}^2 \cdot \text{K}^4)[(260 \text{ K})^4 - (320 \text{ K})^4]$ = -234 W/m<sup>2</sup>

**Discussion** Note that the surface of an ordinary gray material of high absorptivity gains heat at a rate of  $307 \text{ W/m}^2$ . The amount of heat gain increases to  $575 \text{ W/m}^2$  when the surface is coated with a selective material that has the same absorptivity for solar radiation but a low emissivity for infrared radiation. Also note that the surface of an ordinary gray material of high reflectivity still gains heat at a rate of  $34 \text{ W/m}^2$ . When the surface is coated with a selective material that has the same reflectivity for solar radiation but a high emissivity for infrared radiation, the surface loses  $234 \text{ W/m}^2$  instead. Therefore, the temperature of the surface will decrease when a selective reflector surface is used.

### **TOPIC OF SPECIAL INTEREST\***

### Solar Heat Gain Through Windows

The sun is the primary heat source of the earth, and the solar irradiance on a surface normal to the sun's rays beyond the earth's atmosphere at the mean earth–sun distance of 149.5 million km is called the total solar irradiance or solar constant. The accepted value of the solar constant is 1373 W/m<sup>2</sup> (435.4 Btu/h  $\cdot$  ft<sup>2</sup>), but its value changes by 3.5 percent from a maximum of 1418 W/m<sup>2</sup> on January 3 when the earth is closest to the sun, to a minimum of 1325 W/m<sup>2</sup> on July 4 when the earth is farthest away from the sun. The spectral distribution of solar radiation beyond the earth's atmosphere resembles the energy emitted by a blackbody at 5780°C, with about 9 percent of the energy contained in the ultraviolet region (0.4 to 0.7  $\mu$ m), and the remaining 52 percent in the near-infrared region (0.7 to 3.5  $\mu$ m). The peak radiation occurs at a wavelength of about 0.48  $\mu$ m, which corresponds to the green color portion of the visible spectrum. Obviously a

<sup>\*</sup>This section can be skipped without a loss in continuity.