

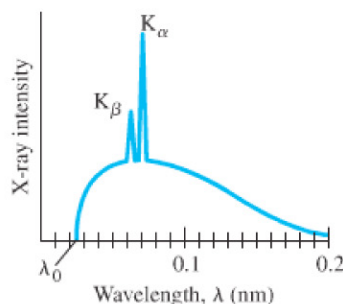
**CAUTION**  
Subshells are not always filled in "order"

The presence of the **transition elements** in the center of the table, as well as the lanthanides (rare earths) and actinides below, is a result of incomplete inner shells. For the lowest  $Z$  elements, the subshells are filled in a simple order: first  $1s$ , then  $2s$ , followed by  $2p$ ,  $3s$ , and  $3p$ . You might expect that  $3d$  ( $n = 3$ ,  $l = 2$ ) would be filled next, but it isn't. Instead, the  $4s$  level actually has a slightly lower energy than the  $3d$  (due to electrons interacting with each other), so it fills first (K and Ca). Only then does the  $3d$  shell start to fill up, beginning with Sc, as can be seen in Table 28–3. (The  $4s$  and  $3d$  levels are close, so some elements have only one  $4s$  electron, such as Cr.) Most of the chemical properties of these transition elements are governed by the relatively loosely held  $4s$  electrons, and hence they usually have valences of  $+1$  or  $+2$ . A similar effect is responsible for the *lanthanides* and *actinides*, which are shown at the bottom of the periodic table for convenience. All have very similar chemical properties, which are determined by their two outer  $6s$  or  $7s$  electrons, whereas the different numbers of electrons in the unfilled inner shells have little effect.

## \* 28–9 X-Ray Spectra and Atomic Number

The line spectra of atoms in the visible, UV, and IR regions of the EM spectrum are mainly due to transitions between states of the outer electrons. Much of the charge of the nucleus is shielded from these electrons by the negative charge on the inner electrons. But the innermost electrons in the  $n = 1$  shell "see" the full charge of the nucleus. Since the energy of a level is proportional to  $Z^2$  (see Eq. 27–15), for an atom with  $Z = 50$ , we would expect wavelengths about  $50^2 = 2500$  times shorter than those found in the Lyman series of hydrogen (around 100 nm), or  $10^{-2}$  to  $10^{-1}$  nm. Such short wavelengths lie in the X-ray region of the spectrum.

**FIGURE 28–11** Spectrum of X-rays emitted from a molybdenum target in an X-ray tube operated at 50 kV.



X-rays are produced when electrons accelerated by a high voltage strike the metal target inside the X-ray tube (Section 25–11). If we look at the spectrum of wavelengths emitted by an X-ray tube, we see that the spectrum consists of two parts: a continuous spectrum with a cutoff at some  $\lambda_0$  which depends only on the voltage across the tube, and a series of peaks superimposed. A typical example is shown in Fig. 28–11. The smooth curve and the cutoff wavelength  $\lambda_0$  move to the left as the voltage across the tube increases. The sharp lines or peaks (labeled  $K_\alpha$  and  $K_\beta$  in Fig. 28–11), however, remain at the same wavelength when the voltage is changed, although they are located at different wavelengths when different target materials are used. This observation suggests that the peaks are characteristic of the material used. Indeed, we can explain them by imagining that the electrons accelerated by the high voltage of the tube can reach sufficient energies that when they collide with the atoms of the target, they can knock out one of the very tightly held inner electrons. Then we explain these **characteristic X-rays** (the peaks in Fig. 28–11) as photons emitted when an electron in an upper state drops down to fill the vacated lower state. The K lines result from transitions *into* the K shell ( $n = 1$ ). The  $K_\alpha$  line consists of photons emitted in a transition that originates from the  $n = 2$  (L) shell and drops to the  $n = 1$  (K) shell, whereas the  $K_\beta$  line reflects a transition from the  $n = 3$  (M) shell down to the K shell. An L line is due to a transition into the L shell, and so on.

Characteristic X-rays