

## ATOMIC STRUCTURE

### Line Spectra

Although objects at high temperature emit a continuous spectrum of electromagnetic radiation, a different kind of spectrum is observed when pure samples of individual elements are heated. For example, when a high-voltage electrical discharge is passed through a sample of hydrogen gas at low pressure, the resulting individual isolated hydrogen atoms caused by the dissociation of  $\text{H}_2$  emit a red light. Unlike blackbody radiation, the color of the light emitted by the hydrogen atoms does not depend greatly on the temperature of the gas in the tube. When the emitted light is passed through a prism, only a few narrow lines, called a line spectrum, rather than a continuous range of colors. The light emitted by hydrogen atoms is red because, of its four characteristic lines, the most intense line in its spectrum is in the red portion of the visible spectrum, at 656 nm. With sodium, however, we observe a yellow color because the most intense lines in its spectrum are in the yellow portion of the spectrum, at about 589 nm.



**Figure 2.3.1 The Emission of Light by Hydrogen Atoms** (a) A sample of excited hydrogen atoms emits a characteristic red light. (b) When the light emitted by a sample of excited hydrogen atoms is split into its component wavelengths by a prism, four characteristic violet, blue, green, and red emission lines can be observed, the most intense of which is at 656 nm.

Such *emission spectra* were observed for many other elements in the late 19th century, which presented a major challenge because classical physics was unable to explain them. Part of the explanation is provided by Planck's equation (as seen above): the observation of only a few values of  $\lambda$  (or  $\nu$ ) in the line spectrum meant that only a few values of  $E$  were possible. Thus *the energy levels of a hydrogen atom had to be quantized*; in other words, only states that had certain values

of energy were possible, or *allowed*. If a hydrogen atom could have *any* value of energy, then a continuous spectrum would have been observed, similar to blackbody radiation.

In 1885, a Swiss mathematics teacher, Johann Balmer (1825–1898), showed that the frequencies of the lines observed in the visible region of the spectrum of hydrogen fit a simple equation that can be expressed as follows:

$$\nu = \text{constant} \left( \frac{1}{2^2} - \frac{1}{n^2} \right)$$

where  $n = 3, 4, 5, 6$ . As a result, these lines are known as the *Balmer series*. The Swedish physicist Johannes Rydberg (1854–1919) subsequently restated and expanded Balmer's result in the *Rydberg equation*:

$$\frac{1}{\lambda} = \mathfrak{R} \left( \frac{1}{n_1^2} - \frac{1}{n_2^2} \right)$$

where  $n_1$  and  $n_2$  are positive integers,  $n_2 > n_1$ , and  $\mathfrak{R}$  the *Rydberg constant*, has a value of  $1.09737 \times 10^7 \text{ m}^{-1}$ .

Like Balmer's equation, Rydberg's simple equation described the wavelengths of the visible lines in the emission spectrum of hydrogen (with  $n_1 = 2$ ,  $n_2 = 3, 4, 5, \dots$ ). More important, Rydberg's equation also described the wavelengths of other series of lines that would be observed in the emission spectrum of hydrogen: one in the ultraviolet ( $n_1 = 1$ ,  $n_2 = 2, 3, 4, \dots$ ) and one in the infrared ( $n_1 = 3$ ,  $n_2 = 4, 5, 6$ ). Unfortunately, scientists had not yet developed any theoretical justification for an equation of this form.

### The Bohr Model

In 1913, a Danish physicist, Niels Bohr (1885–1962; Nobel Prize in Physics, 1922), proposed a theoretical model for the hydrogen atom that explained its emission spectrum. Bohr's model required only one assumption: *The electron moves around the nucleus in circular orbits that can have only certain allowed radii.* As discussed earlier, Rutherford's earlier model of the atom had also assumed that electrons moved in circular orbits around the nucleus and that the atom was held together by the electrostatic attraction between the positively charged nucleus and the negatively charged electron. Although we now know that the assumption of circular orbits was

incorrect, Bohr's insight was to propose that *the electron could occupy only certain regions of space*.

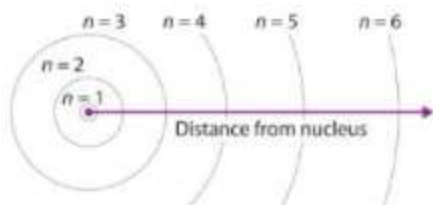
Using classical physics, Bohr showed that the energy of an electron in a particular orbit is given by

$$E_n = \frac{-\mathfrak{R}hc}{n^2}$$

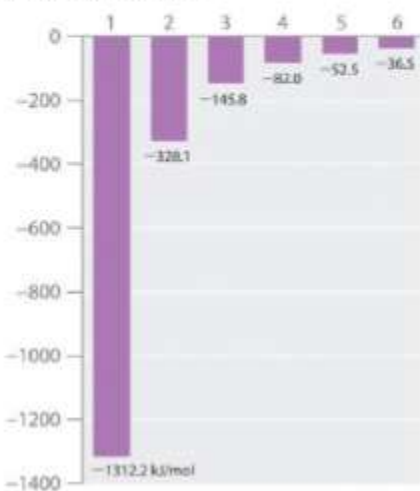
where  $\mathfrak{R}$  is the Rydberg constant,  $h$  is Planck's constant,  $c$  is the speed of light, and  $n$  is a positive integer corresponding to the number assigned to the orbit, with  $n = 1$  corresponding to the orbit closest to the nucleus. In this model  $n = \infty$  corresponds to the level where the energy holding the electron and the nucleus together is zero. In that level, the electron is unbound from the nucleus and the atom has been separated into a negatively charged (the electron) and a positively charged (the nucleus) ion. In this state the radius of the orbit is also infinite. The atom has been ionized.

As  $n$  decreases, the energy holding the electron and the nucleus together becomes increasingly negative, the radius of the orbit shrinks and more energy is needed to ionize the atom. The orbit with  $n = 1$  is the lowest lying and most tightly bound. The negative sign in the Bohr equation above indicates that the electron-nucleus pair is more tightly bound when they are near each other than when they are far apart. Because a hydrogen atom with its one electron in this orbit has the lowest possible energy, this is the ground state, the most stable arrangement for a hydrogen atom. As  $n$  increases, the radius of the orbit increases; the electron is farther from the proton, which results in a less stable arrangement with higher potential energy. A hydrogen atom with an electron in an orbit with  $n > 1$  is therefore in an excited state: its energy is higher than the energy of the ground state. When an atom in an excited state undergoes a transition to the ground state in a process called *decay*, it loses energy by emitting a photon whose energy corresponds to the difference in energy between the two states. Figure below shows **the Bohr**

**Model of the Hydrogen Atom:** (a) The distance of the orbit from the nucleus increases with increasing  $n$ . (b) The energy of the orbit becomes increasingly less negative with increasing  $n$ .

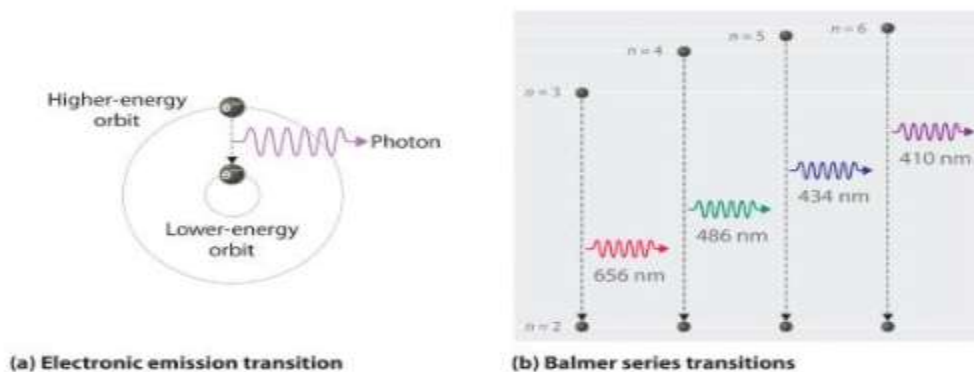


(a) Hydrogen orbits (Bohr model). Orbits are not drawn to scale.



(b) Hydrogen orbit energy (kJ/mol) vs.  $n$

The Emission of Light by a Hydrogen Atom in an Excited State is shown below: (a) Light is emitted when the electron undergoes a transition from an orbit with a higher value of  $n$  (at a higher energy) to an orbit with a lower value of  $n$  (at lower energy). (b) The Balmer series of emission lines is due to transitions from orbits with  $n \geq 3$  to the orbit with  $n = 2$ . The differences in energy between these levels corresponds to light in the visible portion of the electromagnetic spectrum.



So the difference in energy ( $\Delta E$ ) between any two orbits or energy levels is given by  $\Delta E = E_{n_1} - E_{n_2}$  where  $n_1$  is the final orbit and  $n_2$  the initial orbit. Substituting from Bohr's equation for each energy value gives

$$\Delta E = \frac{hc}{\lambda} = -Rhc \left( \frac{1}{n_2^2} - \frac{1}{n_1^2} \right)$$

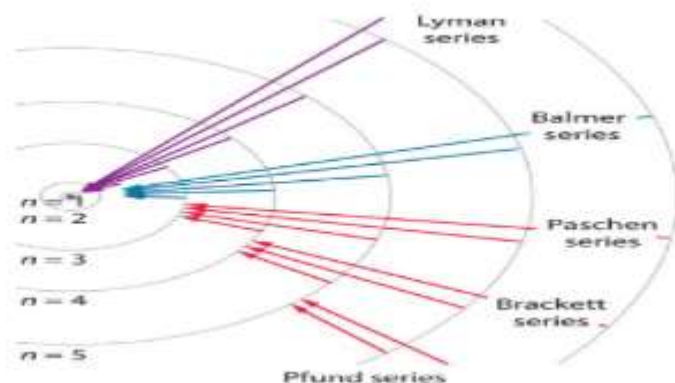
Canceling  $hc$  on both sides gives

$$\frac{1}{\lambda} = -R \left( \frac{1}{n_2^2} - \frac{1}{n_1^2} \right)$$

Except for the negative sign, this is the same equation that Rydberg obtained experimentally.

We can now understand the physical basis for the Balmer series of lines in the emission spectrum of hydrogen. In the spectrum of hydrogen as discussed earlier, the lines in this series correspond to transitions from higher-energy orbits ( $n > 2$ ) to the *second* orbit ( $n = 2$ ). Thus the hydrogen atoms in the sample have absorbed energy from the electrical discharge and decayed from a higher-energy excited state ( $n > 2$ ) to a lower-energy state ( $n = 2$ ) by emitting a photon of electromagnetic radiation whose energy corresponds exactly to the *difference* in energy between the two states. The  $n = 3$  to  $n = 2$  transition gives rise to the line at 656 nm (red), the  $n = 4$  to  $n = 2$  transition to the line at 486 nm (green), the  $n = 5$  to  $n = 2$  transition to the line at 434

nm (blue), and the  $n = 6$  to  $n = 2$  transition to the line at 410 nm (violet). Because a sample of hydrogen contains a large number of atoms, the intensity of the various lines in a line spectrum depends on the number of atoms in each excited state. At the temperature in the gas discharge tube, more atoms are in the  $n = 3$  than the  $n \geq 4$  levels. Consequently, the  $n = 3$  to  $n = 2$  transition is the most intense line, producing the characteristic red color of a hydrogen discharge. Other families of lines are produced by transitions from excited states with  $n > 1$  to the orbit with  $n = 1$  or to orbits with  $n \geq 3$ . These transitions are shown schematically in the Figure below.



**Figure above shows Electron Transitions Responsible for the Various Series of Lines Observed in the Emission Spectrum of Hydrogen.** *The Lyman series of lines is due to transitions from higher-energy orbits to the lowest-energy orbit ( $n = 1$ ); these transitions release a great deal of energy, corresponding to radiation in the ultraviolet portion of the electromagnetic spectrum. The Paschen, Brackett, and Pfund series of lines are due to transitions from higher-energy orbits to orbits with  $n = 3$ ,  $4$ , and  $5$ , respectively; these transitions release substantially less energy, corresponding to infrared radiation. (Orbits are not drawn to scale.)*

### Application of Electron Transitions

In contemporary applications, electron transitions are used in timekeeping that needs to be exact. Telecommunications systems, such as cell phones, depend on timing signals that are accurate to within a millionth of a second per day, as are the devices that control the US power grid. Global positioning system (GPS) signals must be accurate to within a billionth of a second per day, which is equivalent to gaining or losing no more than one second in 1,400,000 years.

Quantifying time requires finding an event with an interval that repeats on a regular basis. To achieve the accuracy required for modern purposes, physicists have turned to the atom. The current standard used to calibrate clocks is the cesium atom. Supercooled cesium atoms are placed in a vacuum chamber and bombarded with microwaves whose frequencies are carefully controlled. When the frequency is exactly right, the atoms absorb enough energy to undergo an electronic transition to a higher-energy state. Decay to a lower-energy state emits radiation. The microwave frequency is continually adjusted, serving as the clock's pendulum. In 1967, the second was defined as the duration of 9,192,631,770 oscillations of the resonant frequency of a cesium atom, called the *cesium clock*. Research is currently under way to develop the next generation of atomic clocks that promise to be even more accurate. Such devices would allow scientists to monitor vanishingly faint electromagnetic signals produced by nerve pathways in the brain and geologists to measure variations in gravitational fields, which cause fluctuations in time, that would aid in the discovery of oil or minerals.

### Exercise

1. The so-called Lyman series of lines in the emission spectrum of hydrogen corresponds to transitions from various excited states to the  $n = 1$  orbit. Calculate the wavelength of the lowest-energy line in the Lyman series to three significant figures. In what region of the electromagnetic spectrum does it occur?

**Given:** lowest-energy orbit in the Lyman series

**Asked for:** wavelength of the lowest-energy Lyman line and corresponding region of the spectrum

**Strategy:**

**A** Substitute the appropriate values into

$$\frac{1}{\lambda} = -\mathfrak{R} \left( \frac{1}{n_2^2} - \frac{1}{n_1^2} \right)$$

(the Rydberg equation) and solve for  $\lambda$ .

**B** Use suitable colour chart to locate the region of the electromagnetic spectrum corresponding to the calculated wavelength.

**Answer:**  $\lambda = 1.215 \times 10^{-7} \text{ m} = 122 \text{ nm}$

This emission line is called Lyman alpha. It is the strongest atomic emission line from the sun and drives the chemistry of the upper atmosphere of all the planets producing ions by stripping electrons from atoms and molecules. It is completely absorbed by oxygen in the upper stratosphere, dissociating  $\text{O}_2$  molecules to O atoms which react with other  $\text{O}_2$  molecules to form stratospheric ozone

**B** This wavelength is in the ultraviolet region of the spectrum.

2. The Pfund series of lines in the emission spectrum of hydrogen corresponds to transitions from higher excited states to the  $n = 5$  orbit. Calculate the wavelength of the *second* line in the Pfund series to three significant figures. In which region of the spectrum does it lie?

**Answer:**  $4.65 \times 10^3 \text{ nm}$ ; infrared

Bohr's model of the hydrogen atom gave an exact explanation for its observed emission spectrum. The following are his key contributions to our understanding of atomic structure:

- Electrons can occupy only certain regions of space, called *orbits*.
- Orbits closer to the nucleus are lower in energy.
- Electrons can move from one orbit to another by absorbing or emitting energy, giving rise to characteristic spectra.

Unfortunately, Bohr could not explain *why* the electron should be restricted to particular orbits. Also, despite a great deal of tinkering, such as assuming that orbits could be ellipses rather than circles, his model could not quantitatively explain the emission spectra of any element other than



hydrogen. In fact, Bohr's model worked only for species that contained just one electron: H,  $\text{He}^+$ ,  $\text{Li}^{2+}$ , and so forth. Scientists needed a fundamental change in their way of thinking about the electronic structure of atoms to advance beyond the Bohr model.

### Applications of Emission and Absorption Spectra

If white light is passed through a sample of hydrogen, hydrogen atoms absorb energy as an electron is excited to higher energy levels (orbits with  $n \geq 2$ ). If the light that emerges is passed through a prism, it forms a continuous spectrum with *black* lines (corresponding to no light passing through the sample) at 656, 468, 434, and 410 nm. These wavelengths correspond to the  $n = 2$  to  $n = 3$ ,  $n = 2$  to  $n = 4$ ,  $n = 2$  to  $n = 5$ , and  $n = 2$  to  $n = 6$  transitions. Any given element therefore has both a characteristic emission spectrum and a characteristic absorption spectrum, which are essentially complementary images.

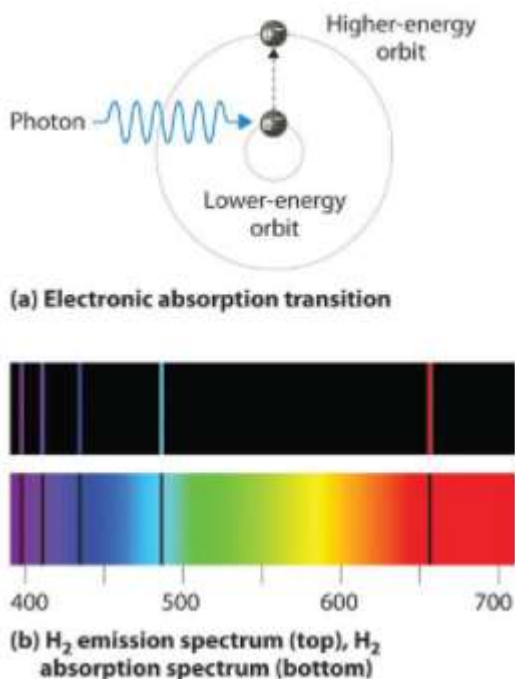
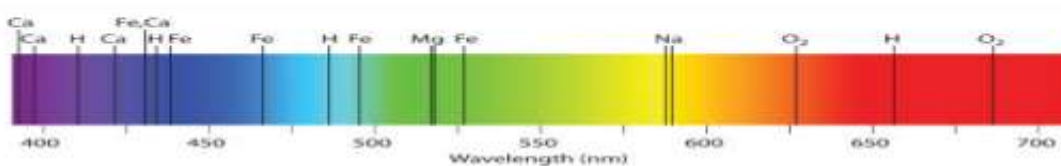


Figure: **Absorption and Emission Spectra** (a) When a hydrogen atom absorbs a photon of light, an electron is excited to an orbit that has a higher energy and larger value of  $n$ . (b) Images of the emission and absorption spectra of hydrogen are shown here.

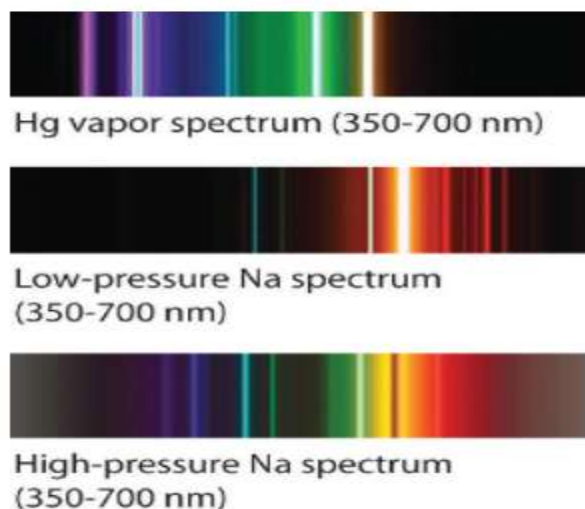
Emission and absorption spectra form the basis of *spectroscopy*, which uses spectra to provide information about the structure and the composition of a substance or an object. In particular, astronomers use emission and absorption spectra to determine the composition of stars and interstellar matter. As an example, consider the spectrum of sunlight shown in the figure below. Because the sun is very hot, the light it emits is in the form of a continuous emission spectrum. Superimposed on it, however, is a series of dark lines due primarily to the absorption of specific frequencies of light by cooler atoms in the outer atmosphere of the sun. By comparing these lines with the spectra of elements measured on Earth, we now know that the sun contains large amounts of hydrogen, iron, and carbon, along with smaller amounts of other elements. During the solar eclipse of 1868, the French astronomer Pierre Janssen (1824–1907) observed a set of lines that did not match those of any known element. He suggested that they were due to the presence of a new element, which he named *helium*, from the Greek *helios*, meaning “sun.” Helium was finally discovered in uranium ores on Earth in 1895. Alpha particles are helium nuclei. Alpha particles emitted by the radioactive uranium, pick up electrons from the rocks to form helium atoms.



**Figure: The Visible Spectrum of Sunlight.** *The characteristic dark lines are mostly due to the absorption of light by elements that are present in the cooler outer part of the sun’s atmosphere; specific elements are indicated by the labels. The lines at 628 and 687 nm, however, are due to the absorption of light by oxygen molecules in Earth’s atmosphere.*

The familiar red color of “neon” signs used in advertising is due to the emission spectrum of neon. Similarly, the blue and yellow colors of certain street lights are caused, respectively, by mercury and sodium discharges. In all these cases, an electrical discharge excites neutral atoms to a higher energy state, and light is emitted when the atoms decay to the ground state. In the case of

mercury, most of the emission lines are below 450 nm, which produces a blue light. In the case of sodium, the most intense emission lines are at 589 nm, which produces an intense yellow light.



**Figure: The emission spectra of sodium and mercury** *Many street lights use bulbs that contain sodium or mercury vapor. Due to the very different emission spectra of these elements, they emit light of different colors. The lines in the sodium lamp are broadened by collisions. The dark line in the center of the high pressure sodium lamp where the low pressure lamp is strongest is caused by absorption of light in the cooler outer part of the lamp.*

### **The Chemistry of Fireworks: An Applications of Emission Spectra**

The colors of fireworks are also due to atomic emission spectra. As shown in part (a) in the figure below, a typical shell used in a fireworks display contains gunpowder to propel the shell into the air and a fuse to initiate a variety of reactions that produce heat and small explosions. Thermal energy excites the atoms to higher energy states; as they decay to lower energy states, the atoms emit light that gives the familiar colors. When oxidant/reductant mixtures are ignited, a flash of white or yellow light is produced along with a loud bang. Achieving the colors requires adding a small amount of a substance that has an emission spectrum in the desired portion of the visible spectrum.



**Figure:** The Chemistry of Fireworks *(a) In the “multibreak” shell used for fireworks, the chambers contain mixtures of fuels and oxidizers plus compounds for special effects (“stars”) connected by time-delay fuses so that the chambers explode in stages. (b) The finale of a fireworks display usually consists of many shells fired simultaneously to give a dazzling multicolor display. The labels indicate the substances that are responsible for the colors of some of the fireworks shown.*

For example, sodium is used for yellow because of its 589 nm emission lines. The intense yellow color of sodium would mask most other colors, so potassium and ammonium salts, rather than sodium salts, are usually used as oxidants to produce other colors, which explains the preponderance of such salts. Strontium salts, which are also used in highway flares, emit red light, whereas barium gives a green color. Blue is one of the most difficult colors to achieve. Copper(II) salts emit a pale blue light, but copper is dangerous to use because it forms highly unstable explosive compounds with anions such as chlorate. As you might guess, preparing fireworks with the desired properties is a complex, challenging, and potentially hazardous process.

## Lasers

Most light emitted by atoms is *polychromatic*—containing more than one wavelength. In contrast, *lasers* (from *light amplification by stimulated emission of radiation*) emit *monochromatic* light—a single wavelength only. Lasers have many applications in fiber-optic telecommunications, the reading and recording of compact discs (CDs) and digital video

discs (DVDs), metal cutting, semiconductor fabrication, supermarket checkout scanners and laser pointers. Laser beams are generated by the same general phenomenon that gives rise to emission spectra, with one difference: only a single excited state is produced, which in principle results in only a single frequency of emitted light. In practice, however, inexpensive commercial lasers actually emit light with a very narrow range of wavelengths.

### Exercise 1

1. Is the spectrum of the light emitted by isolated atoms of an element discrete or continuous? How do these spectra differ from those obtained by heating a bulk sample of a solid element? Explain your answers.
2. Explain why each element has a characteristic emission and absorption spectra. If spectral emissions had been found to be continuous rather than discrete, what would have been the implications for Bohr's model of the atom?
3. Explain the differences between a ground state and an excited state. Describe what happens in the spectrum of a species when an electron moves from a ground state to an excited state. What happens in the spectrum when the electron falls from an excited state to a ground state?
4. What phenomenon causes a neon sign to have a characteristic color? If the emission spectrum of an element is constant, why do some neon signs have more than one color?
5. How is light from a laser different from the light emitted by a light source such as a light bulb? Describe how a laser produces light.

### Exercise 2

1. Using a Bohr model and the transition from  $n = 2$  to  $n = 3$  in an atom with a single electron, describe the mathematical relationship between an emission spectrum and an absorption spectrum. What is the energy of this transition? What does the sign of the energy value represent in this case? What range of light is associated with this transition?
2. If a hydrogen atom is excited from an  $n = 1$  state to an  $n = 3$  state, how much energy does this correspond to? Is this an absorption or an emission? What is the wavelength of the

- photon involved in this process? To what region of the electromagnetic spectrum does this correspond?
3. The hydrogen atom emits a photon with a 486 nm wavelength, corresponding to an electron decaying from the  $n = 4$  level to which level? What is the color of the emission?
  4. An electron in a hydrogen atom can decay from the  $n = 3$  level to  $n = 2$  level. What is the color of the emitted light? What is the energy of this transition?
  5. Calculate the wavelength and energy of the photon that gives rise to the third line in order of increasing energy in the Lyman series in the emission spectrum of hydrogen. In what region of the spectrum does this wavelength occur? Describe qualitatively what the absorption spectrum looks like.
  6. The wavelength of one of the lines in the Lyman series of hydrogen is 121 nm. In what region of the spectrum does this occur? To which electronic transition does this correspond?
  7. The emission spectrum of helium is shown. Estimate what change in energy ( $\Delta E$ ) gives rise to each line?
  8. Removing an electron from solid potassium requires 222 kJ/mol. Would you expect to observe a photoelectric effect for potassium using a photon of blue light ( $\lambda = 485$  nm)? What is the longest wavelength of energy capable of ejecting an electron from potassium? What is the corresponding color of light of this wavelength?
  9. The binding energy of an electron is the energy needed to remove an electron from its lowest energy state. According to Bohr's postulates, calculate the binding energy of an electron in a hydrogen atom. There are  $6.02 \times 10^{23}$  atoms in 1g of hydrogen atoms. What wavelength in nanometers is required to remove such an electron from one hydrogen atom?
  10. As a radio astronomer, you have observed spectral lines for hydrogen corresponding to a state with  $n = 320$ , and you would like to produce these lines in the laboratory. Is this feasible? Why or why not?

#### Answers

1. 656 nm; red light
- 2.

3.  $n = 2$ , blue-green light
- 4.
5. 97.2 nm,  $2.04 \times 10^{-18}$  J/photon, ultraviolet light, absorption spectrum is a single dark line at a wavelength of 97.2 nm
- 6.
7. Violet: 390 nm, 307 kJ/mol photons; Blue-purple: 440 nm, 272 kJ/mol photons; Blue-green: 500 nm, 239 kJ/mol photons; Orange: 580 nm, 206 kJ/mol photons; Red: 650 nm, 184 kJ/mol photons
- 8.
9. 1313 kJ/mol,  $\lambda \leq 91.1$  nm
- 10.