

## 4.3: Electron Configurations- How Electrons Occupy Orbitals

### Learning Objectives

- Derive the predicted ground-state electron configurations of atoms
- Identify and explain exceptions to predicted electron configurations for atoms and ions
- Relate electron configurations to element classifications in the periodic table

Having introduced the basics of atomic structure and quantum mechanics, we can use our understanding of quantum numbers to determine how atomic orbitals relate to one another. This allows us to determine which orbitals are occupied by electrons in each atom. The specific arrangement of electrons in orbitals of an atom determines many of the chemical properties of that atom.

### Orbital Energies and Atomic Structure

The energy of atomic orbitals increases as the principal quantum number,  $n$ , increases. In any atom with two or more electrons, the repulsion between the electrons makes energies of subshells with different values of  $l$  differ so that the energy of the orbitals increases within a shell in the order  $s < p < d < f$ . Figure 4.3.1 depicts how these two trends in increasing energy relate. The 1s orbital at the bottom of the diagram is the orbital with electrons of lowest energy. The energy increases as we move up to the 2s and then 2p, 3s, and 3p orbitals, showing that the increasing  $n$  value has more influence on energy than the increasing  $l$  value for small atoms. However, this pattern does not hold for larger atoms. The 3d orbital is higher in energy than the 4s orbital. Such overlaps continue to occur frequently as we move up the chart.

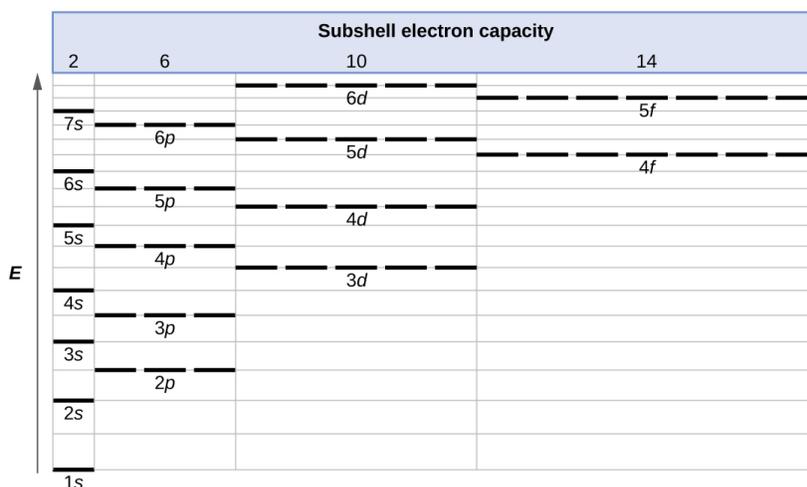


Figure 4.3.1: Generalized energy-level diagram for atomic orbitals in an atom with two or more electrons (not to scale).

Electrons in successive atoms on the periodic table tend to fill low-energy orbitals first. Thus, many students find it confusing that, for example, the 5p orbitals fill immediately after the 4d, and immediately before the 6s. The filling order is based on observed experimental results, and has been confirmed by theoretical calculations. As the principal quantum number,  $n$ , increases, the size of the orbital increases and the electrons spend more time farther from the nucleus. Thus, the attraction to the nucleus is weaker and the energy associated with the orbital is higher (less stabilized). But this is not the only effect we have to take into account. Within each shell, as the value of  $l$  increases, the electrons are less penetrating (meaning there is less electron density found close to the nucleus), in the order  $s > p > d > f$ . Electrons that are closer to the nucleus slightly repel electrons that are farther out, offsetting the more dominant electron–nucleus attractions slightly (recall that all electrons have  $-1$  charges, but nuclei have  $+Z$  charges). This phenomenon is called shielding and will be discussed in more detail in the next section. Electrons in orbitals that experience more shielding are less stabilized and thus higher in energy. For small orbitals (1s through 3p), the increase in energy due to  $n$  is more significant than the increase due to  $l$ ; however, for larger orbitals the two trends are comparable and cannot be simply predicted. We will discuss methods for remembering the observed order.

The arrangement of electrons in the orbitals of an atom is called the electron configuration of the atom. We describe an electron configuration with a symbol that contains three pieces of information ( Figure 4.3.2):

1. The number of the principal quantum shell,  $n$ ,

2. The letter that designates the orbital type (the subshell,  $l$ ), and
3. A superscript number that designates the number of electrons in that particular subshell.

For example, the notation  $2p^4$  (read "two-p-four") indicates four electrons in a  $p$  subshell ( $l = 1$ ) with a principal quantum number ( $n$ ) of 2. The notation  $3d^8$  (read "three-d-eight") indicates eight electrons in the  $d$  subshell (i.e.,  $l = 2$ ) of the principal shell for which  $n = 3$ .

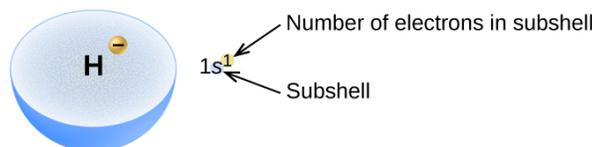


Figure 4.3.2: The diagram of an electron configuration specifies the subshell ( $n$  and  $l$  value, with letter symbol) and superscript number of electrons.

## The Aufbau Principle

To determine the electron configuration for any particular atom, we can "build" the structures in the order of atomic numbers. Beginning with hydrogen, and continuing across the periods of the periodic table, we add one proton at a time to the nucleus and one electron to the proper subshell until we have described the electron configurations of all the elements. This procedure is called the Aufbau principle, from the German word *Aufbau* ("to build up"). Each added electron occupies the subshell of lowest energy available (in the order shown in Figure 4.3.3), subject to the limitations imposed by the allowed quantum numbers according to the Pauli exclusion principle. Electrons enter higher-energy subshells only after lower-energy subshells have been filled to capacity. Figure 4.3.3 illustrates the traditional way to remember the filling order for atomic orbitals.

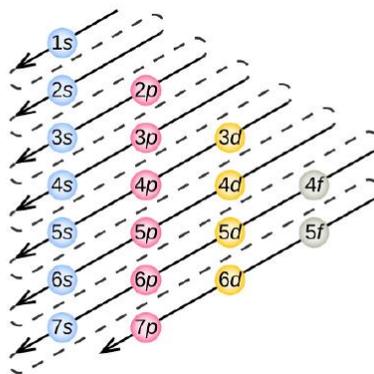


Figure 4.3.3: The arrow leads through each subshell in the appropriate filling order for electron configurations. This chart is straightforward to construct. Simply make a column for all the  $s$  orbitals with each  $n$  shell on a separate row. Repeat for  $p$ ,  $d$ , and  $f$ . Be sure to only include orbitals allowed by the quantum numbers (no  $1p$  or  $2d$ , and so forth). Finally, draw diagonal lines from top to bottom as shown.

Since the arrangement of the periodic table is based on the electron configurations, Figure 4.3.4 provides an alternative method for determining the electron configuration. The filling order simply begins at hydrogen and includes each subshell as you proceed in increasing  $Z$  order. For example, after filling the  $3p$  block up to Ar, we see the orbital will be  $4s$  (K, Ca), followed by the  $3d$  orbitals.

**Electron Configuration Table**

Period	Group																	18	
1	1	H 1																	He 1
		1s																	1s
2	1	Li 1	2													10	18		
		2s	2													2p	18		
3	1	Na 1	Mg 2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18
		3s	2	3	4	5	6	7	8	9	10	11	12	3p	4	5	6	18	
4	1	K 1	Ca 2	Sc 1	Ti 2	V 3	Cr 4	Mn 5	Fe 6	Co 7	Ni 8	Cu 9	Zn 10	Ga 1	Ge 2	As 3	Se 4	Br 5	Kr 6
		4s	2	3d										4p					
5	1	Rb 1	Sr 2	Y 1	Zr 2	Nb 3	Mo 4	Tc 5	Ru 6	Rh 7	Pd 8	Ag 9	Cd 10	In 1	Sn 2	Sb 3	Te 4	I 5	Xe 6
		5s	2	4d										5p					
6	1	Cs 1	Ba 2	La *1	Hf 2	Ta 3	W 4	Re 5	Os 6	Ir 7	Pt 8	Au 9	Hg 10	Tl 1	Pb 2	Bi 3	Po 4	At 5	Rn 6
		6s	2	5d										6p					
7	1	Fr 1	Ra 2	Ac **1	Rf 2	Db 3	Sg 4	Bh 5	Hs 6	Mt 7	Ds 8	Rg 9	Cn 10	Uut	Fl	Uup	Lv	Uus	Uuo
		7s	2	6d															
		* Ce 1 Pr 2 Nd 3 Pm 4 Sm 5 Eu 6 Gd 7 Tb 8 Dy 9 Ho 10 Er 11 Tm 12 Yb 13 Lu 14																	
		** Th 1 Pa 2 U 3 Np 4 Pu 5 Am 6 Cm 7 Bk 8 Cf 9 Es 10 Fm 11 Md 12 No 13 Lr 14																	

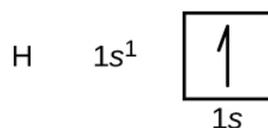
Name → **H** ← Electrons

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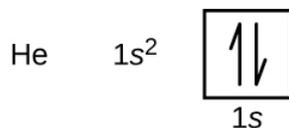
**1s** ← Subshell

Figure 4.3.4: This periodic table shows the electron configuration for each subshell. By “building up” from hydrogen, this table can be used to determine the electron configuration for any atom on the periodic table.

We will now construct the ground-state electron configuration and orbital diagram for a selection of atoms in the first and second periods of the periodic table. Orbital diagrams are pictorial representations of the electron configuration, showing the individual orbitals and the pairing arrangement of electrons. We start with a single hydrogen atom (atomic number 1), which consists of one proton and one electron. Referring to either Figure 4.3.3 or 4.3.4, we would expect to find the electron in the 1s orbital. By convention, the  $m_s = +\frac{1}{2}$  value is usually filled first. The electron configuration and the orbital diagram are:

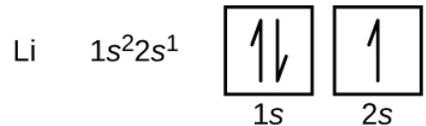


Following hydrogen is the noble gas helium, which has an atomic number of 2. The helium atom contains two protons and two electrons. The first electron has the same four quantum numbers as the hydrogen atom electron ( $n = 1, l = 0, m_l = 0, m_s = +\frac{1}{2}$ ). The second electron also goes into the 1s orbital and fills that orbital. The second electron has the same  $n, l,$  and  $m_l$  quantum numbers, but must have the opposite spin quantum number,  $m_s = -\frac{1}{2}$ . This is in accord with the Pauli exclusion principle: No two electrons in the same atom can have the same set of four quantum numbers. For orbital diagrams, this means two arrows go in each box (representing two electrons in each orbital) and the arrows must point in opposite directions (representing paired spins). The electron configuration and orbital diagram of helium are:

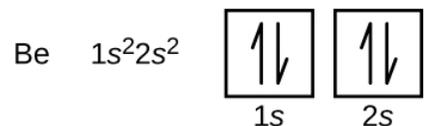


The  $n = 1$  shell is completely filled in a helium atom.

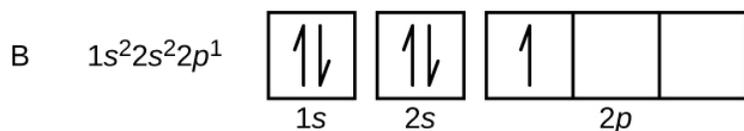
The next atom is the alkali metal lithium with an atomic number of 3. The first two electrons in lithium fill the 1s orbital and have the same sets of four quantum numbers as the two electrons in helium. The remaining electron must occupy the orbital of next lowest energy, the 2s orbital (Figure 4.3.3 or 4.3.4). Thus, the electron configuration and orbital diagram of lithium are:



An atom of the alkaline earth metal beryllium, with an atomic number of 4, contains four protons in the nucleus and four electrons surrounding the nucleus. The fourth electron fills the remaining space in the 2s orbital.



An atom of boron (atomic number 5) contains five electrons. The  $n = 1$  shell is filled with two electrons and three electrons will occupy the  $n = 2$  shell. Because any  $s$  subshell can contain only two electrons, the fifth electron must occupy the next energy level, which will be a  $2p$  orbital. There are three degenerate  $2p$  orbitals ( $m_l = -1, 0, +1$ ) and the electron can occupy any one of these  $p$  orbitals. When drawing orbital diagrams, we include empty boxes to depict any empty orbitals in the same subshell that we are filling.



Carbon (atomic number 6) has six electrons. Four of them fill the 1s and 2s orbitals. The remaining two electrons occupy the  $2p$  subshell. We now have a choice of filling one of the  $2p$  orbitals and pairing the electrons or of leaving the electrons unpaired in two different, but degenerate,  $p$  orbitals. The orbitals are filled as described by Hund's rule: the lowest-energy configuration for an atom with electrons within a set of degenerate orbitals is that having the maximum number of unpaired electrons. Thus, the two electrons in the carbon  $2p$  orbitals have identical  $n$ ,  $l$ , and  $m_s$  quantum numbers and differ in their  $m_l$  quantum number (in accord with the Pauli exclusion principle). The electron configuration and orbital diagram for carbon are:



Nitrogen (atomic number 7) fills the 1s and 2s subshells and has one electron in each of the three  $2p$  orbitals, in accordance with Hund's rule. These three electrons have unpaired spins. Oxygen (atomic number 8) has a pair of electrons in any one of the  $2p$  orbitals (the electrons have opposite spins) and a single electron in each of the other two. Fluorine (atomic number 9) has only one  $2p$  orbital containing an unpaired electron. All of the electrons in the noble gas neon (atomic number 10) are paired, and all of the orbitals in the  $n = 1$  and the  $n = 2$  shells are filled. The electron configurations and orbital diagrams of these four elements are:

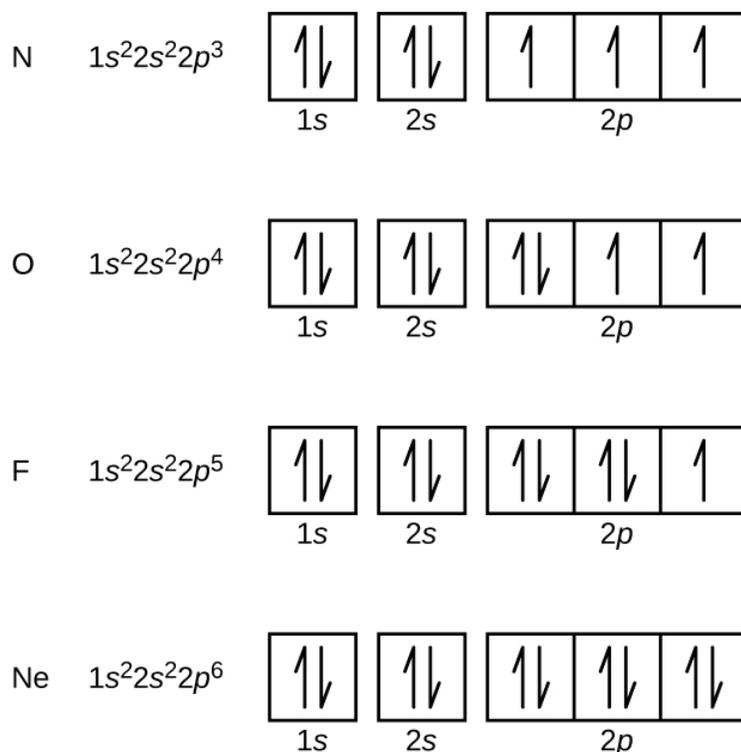


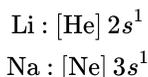
Figure 4.3.5: Since the core electron shells correspond to noble gas electron configurations, we can abbreviate electron configurations by writing the noble gas that matches the core electron configuration, along with the valence electrons in a condensed format. For our sodium example, the symbol [Ne] represents core electrons, ( $1s^2 2s^2 2p^6$ ) and our abbreviated or condensed configuration is [Ne] $3s^1$ .

The alkali metal sodium (atomic number 11) has one more electron than the neon atom. This electron must go into the lowest-energy subshell available, the 3s orbital, giving a  $1s^2 2s^2 2p^6 3s^1$  configuration. The electrons occupying the outermost shell orbital(s) (highest value of  $n$ ) are called valence electrons, and those occupying the inner shell orbitals are called core electrons ( Figure \PageIndex5\PageIndex5). Since the core electron shells correspond to noble gas electron configurations, we can abbreviate electron configurations by writing the noble gas that matches the core electron configuration, along with the valence electrons in a condensed format. For our sodium example, the symbol [Ne] represents core electrons, ( $1s^2 2s^2 2p^6$ ) and our abbreviated or condensed configuration is [Ne] $3s^1$ .



Figure 4.3.5: A core-abbreviated electron configuration (right) replaces the core electrons with the noble gas symbol whose configuration matches the core electron configuration of the other element.

Similarly, the abbreviated configuration of lithium can be represented as [He] $2s^1$ , where [He] represents the configuration of the helium atom, which is identical to that of the filled inner shell of lithium. Writing the configurations in this way emphasizes the similarity of the configurations of lithium and sodium. Both atoms, which are in the alkali metal family, have only one electron in a valence s subshell outside a filled set of inner shells.



The alkaline earth metal magnesium (atomic number 12), with its 12 electrons in a [Ne] $3s^2$  configuration, is analogous to its family member beryllium, [He] $2s^2$ . Both atoms have a filled s subshell outside their filled inner shells. Aluminum (atomic number 13), with 13 electrons and the electron configuration [Ne] $3s^2 3p^1$ , is analogous to its family member boron, [He] $2s^2 2p^1$ .

The electron configurations of silicon (14 electrons), phosphorus (15 electrons), sulfur (16 electrons), chlorine (17 electrons), and argon (18 electrons) are analogous in the electron configurations of their outer shells to their corresponding family members carbon, nitrogen, oxygen, fluorine, and neon, respectively, except that the principal quantum number of the outer shell of the heavier elements has increased by one to  $n = 3$ . Figure 4.3.6 shows the lowest energy, or ground-state, electron configuration for these elements as well as that for atoms of each of the known elements.

**Electron Configuration Table**

Period	Group																	
	1											13	14	15	16	17	18	
1	1 H 1s <sup>1</sup>																2 He 1s <sup>2</sup>	
2	3 Li 2s <sup>1</sup>	4 Be 2s <sup>2</sup>											5 B 2s <sup>2</sup> 2p <sup>1</sup>	6 C 2s <sup>2</sup> 2p <sup>2</sup>	7 N 2s <sup>2</sup> 2p <sup>3</sup>	8 O 2s <sup>2</sup> 2p <sup>4</sup>	9 F 2s <sup>2</sup> 2p <sup>5</sup>	10 Ne 2s <sup>2</sup> 2p <sup>6</sup>
3	11 Na 3s <sup>1</sup>	12 Mg 3s <sup>2</sup>	3	4	5	6	7	8	9	10	11	12	13 Al 3s <sup>2</sup> 3p <sup>1</sup>	14 Si 3s <sup>2</sup> 3p <sup>2</sup>	15 P 3s <sup>2</sup> 3p <sup>3</sup>	16 S 3s <sup>2</sup> 3p <sup>4</sup>	17 Cl 3s <sup>2</sup> 3p <sup>5</sup>	18 Ar 3s <sup>2</sup> 3p <sup>6</sup>
4	19 K 4s <sup>1</sup>	20 Ca 4s <sup>2</sup>	21 Sc 4s <sup>2</sup> 3d <sup>1</sup>	22 Ti 4s <sup>2</sup> 3d <sup>2</sup>	23 V 4s <sup>2</sup> 3d <sup>3</sup>	24 Cr 4s <sup>1</sup> 3d <sup>5</sup>	25 Mn 4s <sup>2</sup> 3d <sup>5</sup>	26 Fe 4s <sup>2</sup> 3d <sup>6</sup>	27 Co 4s <sup>2</sup> 3d <sup>7</sup>	28 Ni 4s <sup>2</sup> 3d <sup>8</sup>	29 Cu 4s <sup>1</sup> 3d <sup>10</sup>	30 Zn 4s <sup>2</sup> 3d <sup>10</sup>	31 Ga 4s <sup>2</sup> 3d <sup>10</sup> 4p <sup>1</sup>	32 Ge 4s <sup>2</sup> 3d <sup>10</sup> 4p <sup>2</sup>	33 As 4s <sup>2</sup> 3d <sup>10</sup> 4p <sup>3</sup>	34 Se 4s <sup>2</sup> 3d <sup>10</sup> 4p <sup>4</sup>	35 Br 4s <sup>2</sup> 3d <sup>10</sup> 4p <sup>5</sup>	36 Kr 4s <sup>2</sup> 3d <sup>10</sup> 4p <sup>6</sup>
5	37 Rb 5s <sup>1</sup>	38 Sr 5s <sup>2</sup>	39 Y 5s <sup>2</sup> 4d <sup>1</sup>	40 Zr 5s <sup>2</sup> 4d <sup>2</sup>	41 Nb 5s <sup>1</sup> 4d <sup>4</sup>	42 Mo 5s <sup>1</sup> 4d <sup>5</sup>	43 Tc 5s <sup>1</sup> 4d <sup>5</sup>	44 Ru 5s <sup>1</sup> 4d <sup>7</sup>	45 Rh 5s <sup>1</sup> 4d <sup>8</sup>	46 Pd 4d <sup>10</sup>	47 Ag 5s <sup>1</sup> 4d <sup>10</sup>	48 Cd 5s <sup>2</sup> 4d <sup>10</sup>	49 In 5s <sup>2</sup> 4d <sup>10</sup> 5p <sup>1</sup>	50 Sn 5s <sup>2</sup> 4d <sup>10</sup> 5p <sup>2</sup>	51 Sb 5s <sup>2</sup> 4d <sup>10</sup> 5p <sup>3</sup>	52 Te 5s <sup>2</sup> 4d <sup>10</sup> 5p <sup>4</sup>	53 I 5s <sup>2</sup> 4d <sup>10</sup> 5p <sup>5</sup>	54 Xe 5s <sup>2</sup> 4d <sup>10</sup> 5p <sup>6</sup>
6	55 Cs 6s <sup>1</sup>	56 Ba 6s <sup>2</sup>	57 La 6s <sup>2</sup> 5d <sup>1</sup>	72 Hf 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>2</sup>	73 Ta 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>3</sup>	74 W 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>4</sup>	75 Re 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>5</sup>	76 Os 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>6</sup>	77 Ir 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>7</sup>	78 Pt 6s <sup>1</sup> 4f <sup>14</sup> 5d <sup>9</sup>	79 Au 6s <sup>1</sup> 4f <sup>14</sup> 5d <sup>10</sup>	80 Hg 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>10</sup>	81 Tl 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>10</sup> 6p <sup>1</sup>	82 Pb 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>10</sup> 6p <sup>2</sup>	83 Bi 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>10</sup> 6p <sup>3</sup>	84 Po 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>10</sup> 6p <sup>4</sup>	85 At 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>10</sup> 6p <sup>5</sup>	86 Rn 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>10</sup> 6p <sup>6</sup>
7	87 Fr 7s <sup>1</sup>	88 Ra 7s <sup>2</sup>	89 Ac 7s <sup>2</sup> 6d <sup>1</sup>	104 Rf 7s <sup>2</sup> 5f <sup>14</sup> 6d <sup>2</sup>	105 Db 7s <sup>2</sup> 5f <sup>14</sup> 6d <sup>3</sup>	106 Sg 7s <sup>2</sup> 5f <sup>14</sup> 6d <sup>4</sup>	107 Bh 7s <sup>2</sup> 5f <sup>14</sup> 6d <sup>5</sup>	108 Hs 7s <sup>2</sup> 5f <sup>14</sup> 6d <sup>6</sup>	109 Mt 7s <sup>2</sup> 5f <sup>14</sup> 6d <sup>7</sup>	110 Ds 7s <sup>2</sup> 5f <sup>14</sup> 6d <sup>8</sup>	111 Rg 7s <sup>2</sup> 5f <sup>14</sup> 6d <sup>9</sup>	112 Cn 7s <sup>2</sup> 5f <sup>14</sup> 6d <sup>10</sup>	113 Uut	114 Fl	115 Uup	116 Lv	117 Uus	118 Uuo
			* 58 Ce 6s <sup>2</sup> 4f <sup>2</sup>	59 Pr 6s <sup>2</sup> 4f <sup>3</sup>	60 Nd 6s <sup>2</sup> 4f <sup>4</sup>	61 Pm 6s <sup>2</sup> 4f <sup>5</sup>	62 Sm 6s <sup>2</sup> 4f <sup>6</sup>	63 Eu 6s <sup>2</sup> 4f <sup>7</sup>	64 Gd 6s <sup>2</sup> 4f <sup>7</sup> 5d <sup>1</sup>	65 Tb 6s <sup>2</sup> 4f <sup>9</sup>	66 Dy 6s <sup>2</sup> 4f <sup>10</sup>	67 Ho 6s <sup>2</sup> 4f <sup>11</sup>	68 Er 6s <sup>2</sup> 4f <sup>12</sup>	69 Tm 6s <sup>2</sup> 4f <sup>13</sup>	70 Yb 6s <sup>2</sup> 4f <sup>14</sup>	71 Lu 6s <sup>2</sup> 4f <sup>14</sup> 5d <sup>1</sup>		
			** 90 Th 7s <sup>2</sup> 6d <sup>2</sup>	91 Pa 7s <sup>2</sup> 5f <sup>2</sup> 6d <sup>1</sup>	92 U 7s <sup>2</sup> 5f <sup>3</sup> 6d <sup>1</sup>	93 Np 7s <sup>2</sup> 5f <sup>4</sup> 6d <sup>1</sup>	94 Pu 7s <sup>2</sup> 5f <sup>6</sup>	95 Am 7s <sup>2</sup> 5f <sup>7</sup>	96 Cm 7s <sup>2</sup> 5f <sup>7</sup> 6d <sup>1</sup>	97 Bk 7s <sup>2</sup> 5f <sup>9</sup> 6d <sup>1</sup>	98 Cf 7s <sup>2</sup> 5f <sup>10</sup>	99 Es 7s <sup>2</sup> 5f <sup>11</sup>	100 Fm 7s <sup>2</sup> 5f <sup>12</sup>	101 Md 7s <sup>2</sup> 5f <sup>13</sup>	102 No 7s <sup>2</sup> 5f <sup>14</sup>	103 Lr 7s <sup>2</sup> 5f <sup>14</sup> 6d <sup>1</sup>		

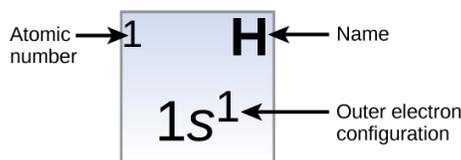


Figure 4.3.6: This version of the periodic table shows the outer-shell electron configuration of each element. Note that down each group, the configuration is often similar.

When we come to the next element in the periodic table, the alkali metal potassium (atomic number 19), we might expect that we would begin to add electrons to the  $3d$  subshell. However, all available chemical and physical evidence indicates that potassium is like lithium and sodium, and that the next electron is not added to the  $3d$  level but is, instead, added to the  $4s$  level (Figure 4.3.3 or 4.3.4). As discussed previously, the  $3d$  orbital with no radial nodes is higher in energy because it is less penetrating and more shielded from the nucleus than the  $4s$ , which has three radial nodes. Thus, potassium has an electron configuration of  $[\text{Ar}]4s^1$ . Hence, potassium corresponds to Li and Na in its valence shell configuration. The next electron is added to complete the  $4s$  subshell and calcium has an electron configuration of  $[\text{Ar}]4s^2$ . This gives calcium an outer-shell electron configuration corresponding to that of beryllium and magnesium.

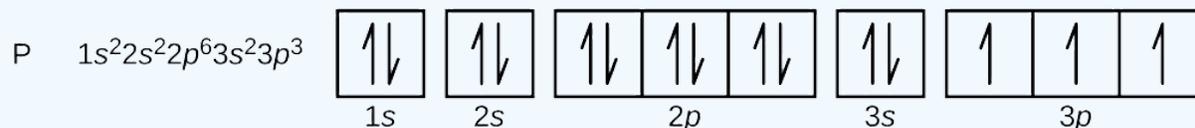
Beginning with the transition metal scandium (atomic number 21), additional electrons are added successively to the  $3d$  subshell. This subshell is filled to its capacity with 10 electrons (remember that for  $l = 2$  [ $d$  orbitals], there are  $2l + 1 = 5$  values of  $m_l$ , meaning that there are five  $d$  orbitals that have a combined capacity of 10 electrons). The  $4p$  subshell fills next. Note that for three series of elements, scandium (Sc) through copper (Cu), yttrium (Y) through silver (Ag), and lutetium (Lu) through gold (Au), a total of 10  $d$  electrons are successively added to the  $(n - 1)$  shell next to the  $n$  shell to bring that  $(n - 1)$  shell from 8 to 18 electrons. For two series, lanthanum (La) through lutetium (Lu) and actinium (Ac) through lawrencium (Lr), 14  $f$  electrons ( $l = 3$ ,  $2l + 1 = 7$   $m_l$  values; thus, seven orbitals with a combined capacity of 14 electrons) are successively added to the  $(n - 2)$  shell to bring that shell from 18 electrons to a total of 32 electrons.

### ✓ Example 4.3.1: Quantum Numbers and Electron Configurations

What is the electron configuration and orbital diagram for a phosphorus atom? What are the four quantum numbers for the last electron added?

#### Solution

The atomic number of phosphorus is 15. Thus, a phosphorus atom contains 15 electrons. The order of filling of the energy levels is  $1s$ ,  $2s$ ,  $2p$ ,  $3s$ ,  $3p$ ,  $4s$ , . . . The 15 electrons of the phosphorus atom will fill up to the  $3p$  orbital, which will contain three electrons:



The last electron added is a  $3p$  electron. Therefore,  $n = 3$  and, for a  $p$ -type orbital,  $l = 1$ . The  $m_l$  value could be  $-1$ ,  $0$ , or  $+1$ . The three  $p$  orbitals are degenerate, so any of these  $m_l$  values is correct. For unpaired electrons, convention assigns the value of  $+\frac{1}{2}$  for the spin quantum number; thus,  $m_s = +\frac{1}{2}$ .

### ? Exercise 4.3.1

Identify the atoms from the electron configurations given:

- $[\text{Ar}]4s^2 3d^5$
- $[\text{Kr}]5s^2 4d^{10} 5p^6$

#### Answer a

Mn

#### Answer b

Xe

The periodic table can be a powerful tool in predicting the electron configuration of an element. However, we do find exceptions to the order of filling of orbitals that are shown in Figure 4.3.3 or 4.3.4. For instance, the electron configurations of the transition metals chromium (Cr; atomic number 24) and copper (Cu; atomic number 29), among others, are not those we would expect. In general, such exceptions involve subshells with very similar energy, and small effects can lead to changes in the order of filling.

In the case of Cr and Cu, we find that half-filled and completely filled subshells apparently represent conditions of preferred stability. This stability is such that an electron shifts from the  $4s$  into the  $3d$  orbital to gain the extra stability of a half-filled  $3d$  subshell (in Cr) or a filled  $3d$  subshell (in Cu). Other exceptions also occur. For example, niobium (Nb, atomic number 41) is predicted to have the electron configuration  $[\text{Kr}]5s^2 4d^3$ . Experimentally, we observe that its ground-state electron configuration is actually  $[\text{Kr}]5s^1 4d^4$ . We can rationalize this observation by saying that the electron–electron repulsions experienced by pairing the electrons in the  $5s$  orbital are larger than the gap in energy between the  $5s$  and  $4d$  orbitals. There is no simple method to predict the exceptions for atoms where the magnitude of the repulsions between electrons is greater than the small differences in energy between subshells.

## Electron Configurations and the Periodic Table

As described earlier, the periodic table arranges atoms based on increasing atomic number so that elements with the same chemical properties recur periodically. When their electron configurations are added to the table (Figure 4.3.6), we also see a periodic recurrence of similar electron configurations in the outer shells of these elements. Because they are in the outer shells of an atom, valence electrons play the most important role in chemical reactions. The outer electrons have the highest energy of the electrons in

an atom and are more easily lost or shared than the core electrons. Valence electrons are also the determining factor in some physical properties of the elements.

Elements in any one group (or column) have the same number of valence electrons; the alkali metals lithium and sodium each have only one valence electron, the alkaline earth metals beryllium and magnesium each have two, and the halogens fluorine and chlorine each have seven valence electrons. The similarity in chemical properties among elements of the same group occurs because they have the same number of valence electrons. It is the loss, gain, or sharing of valence electrons that defines how elements react.

It is important to remember that the periodic table was developed on the basis of the chemical behavior of the elements, well before any idea of their atomic structure was available. Now we can understand why the periodic table has the arrangement it has—the arrangement puts elements whose atoms have the same number of valence electrons in the same group. This arrangement is emphasized in Figure 4.3.6, which shows in periodic-table form the electron configuration of the last subshell to be filled by the Aufbau principle. The colored sections of Figure 4.3.6 show the three categories of elements classified by the orbitals being filled: main group, transition, and inner transition elements. These classifications determine which orbitals are counted in the valence shell, or highest energy level orbitals of an atom.

- Main group elements** (sometimes called **representative elements**) are those in which the last electron added enters an  $s$  or a  $p$  orbital in the outermost shell, shown in blue and red in Figure 4.3.6. This category includes all the nonmetallic elements, as well as many metals and the intermediate semimetallic elements. The valence electrons for main group elements are those with the highest  $n$  level. For example, gallium (Ga, atomic number 31) has the electron configuration  $[\text{Ar}]4s^23d^{10}4p^1$ , which contains three valence electrons (underlined). The completely filled  $d$  orbitals count as core, not valence, electrons.
- Transition elements or transition metals.** These are metallic elements in which the last electron added enters a  $d$  orbital. The valence electrons (those added after the last noble gas configuration) in these elements include the  $ns$  and  $(n - 1)d$  electrons. The official IUPAC definition of transition elements specifies those with partially filled  $d$  orbitals. Thus, the elements with completely filled orbitals (Zn, Cd, Hg, as well as Cu, Ag, and Au in Figure 4.3.6) are not technically transition elements. However, the term is frequently used to refer to the entire  $d$  block (colored yellow in Figure 4.3.6), and we will adopt this usage in this textbook.
- Inner transition elements** are metallic elements in which the last electron added occupies an  $f$  orbital. They are shown in green in Figure 4.3.6. The valence shells of the inner transition elements consist of the  $(n - 2)f$ , the  $(n - 1)d$ , and the  $ns$  subshells. There are two inner transition series:
  - The lanthanide series: lanthanide (La) through lutetium (Lu)
  - The actinide series: actinide (Ac) through lawrencium (Lr)

Lanthanum and actinium, because of their similarities to the other members of the series, are included and used to name the series, even though they are transition metals with no  $f$  electrons.

## Electron Configurations of Ions

We have seen that ions are formed when atoms gain or lose electrons. A cation (positively charged ion) forms when one or more electrons are removed from a parent atom. For main group elements, the electrons that were added last are the first electrons removed. For transition metals and inner transition metals, however, electrons in the  $s$  orbital are easier to remove than the  $d$  or  $f$  electrons, and so the highest  $ns$  electrons are lost, and then the  $(n - 1)d$  or  $(n - 2)f$  electrons are removed. An anion (negatively charged ion) forms when one or more electrons are added to a parent atom. The added electrons fill in the order predicted by the Aufbau principle.

### ✓ Example 4.3.2: Predicting Electron Configurations of Ions

What is the electron configuration and orbital diagram of:

- $\text{Na}^+$
- $\text{P}^{3-}$
- $\text{Al}^{2+}$
- $\text{Fe}^{2+}$
- $\text{Sm}^{3+}$

**Solution**

First, write out the electron configuration for each parent atom. We have chosen to show the full, unabbreviated configurations to provide more practice for students who want it, but listing the core-abbreviated electron configurations is also acceptable.

Next, determine whether an electron is gained or lost. Remember electrons are negatively charged, so ions with a positive charge have *lost* an electron. For main group elements, the last orbital gains or loses the electron. For transition metals, the last *s* orbital loses an electron before the *d* orbitals.

- Na:  $1s^2 2s^2 2p^6 3s^1$ . Sodium cation loses one electron, so  $\text{Na}^+$ :  $1s^2 2s^2 2p^6 3s^1 = \text{Na}^+$ :  $1s^2 2s^2 2p^6$ .
- P:  $1s^2 2s^2 2p^6 3s^2 3p^3$ . Phosphorus trianion gains three electrons, so  $\text{P}^{3-}$ :  $1s^2 2s^2 2p^6 3s^2 3p^6$ .
- Al:  $1s^2 2s^2 2p^6 3s^2 3p^1$ . Aluminum dication loses two electrons  $\text{Al}^{2+}$ :  $1s^2 2s^2 2p^6 3s^2 3p^1 = \text{Al}^{2+}$ :  $1s^2 2s^2 2p^6 3s^1$ .
- Fe:  $1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^6$ . Iron(II) loses two electrons and, since it is a transition metal, they are removed from the 4s orbital  $\text{Fe}^{2+}$ :  $1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^6 = 1s^2 2s^2 2p^6 3s^2 3p^6 3d^6$ .
- Sm:  $1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^{10} 4p^6 5s^2 4d^{10} 5p^6 6s^2 4f^6$ . Samarium trication loses three electrons. The first two will be lost from the 6s orbital, and the final one is removed from the 4f orbital.  $\text{Sm}^{3+}$ :  $1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^{10} 4p^6 5s^2 4d^{10} 5p^6 6s^2 4f^6 = 1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^{10} 4p^6 5s^2 4d^{10} 5p^6 4f^5$ .

### ? Exercise 4.3.2

- Which ion with a +2 charge has the electron configuration  $1s^2 2s^2 2p^6 3s^2 3p^6 3d^{10} 4s^2 4p^6 4d^5$ ?
- Which ion with a +3 charge has this configuration?

#### Answer a

Tc<sup>2+</sup>

#### Answer b

Ru<sup>3+</sup>

## Summary

The relative energy of the subshells determine the order in which atomic orbitals are filled (1s, 2s, 2p, 3s, 3p, 4s, 3d, 4p, and so on). Electron configurations and orbital diagrams can be determined by applying the Pauli exclusion principle (no two electrons can have the same set of four quantum numbers) and Hund's rule (whenever possible, electrons retain unpaired spins in degenerate orbitals).

Electrons in the outermost orbitals, called valence electrons, are responsible for most of the chemical behavior of elements. In the periodic table, elements with analogous valence electron configurations usually occur within the same group. There are some exceptions to the predicted filling order, particularly when half-filled or completely filled orbitals can be formed. The periodic table can be divided into three categories based on the orbital in which the last electron to be added is placed: main group elements (*s* and *p* orbitals), transition elements (*d* orbitals), and inner transition elements (*f* orbitals).

## Glossary

### Aufbau principle

procedure in which the electron configuration of the elements is determined by “building” them in order of atomic numbers, adding one proton to the nucleus and one electron to the proper subshell at a time

### core electron

electron in an atom that occupies the orbitals of the inner shells

### electron configuration

electronic structure of an atom in its ground state given as a listing of the orbitals occupied by the electrons

### Hund's rule

every orbital in a subshell is singly occupied with one electron before any one orbital is doubly occupied, and all electrons in singly occupied orbitals have the same spin

**orbital diagram**

pictorial representation of the electron configuration showing each orbital as a box and each electron as an arrow

**valence electrons**

electrons in the outermost or valence shell (highest value of  $n$ ) of a ground-state atom; determine how an element reacts

**valence shell**

outermost shell of electrons in a ground-state atom; for main group elements, the orbitals with the highest  $n$  level ( $s$  and  $p$  subshells) are in the valence shell, while for transition metals, the highest energy  $s$  and  $d$  subshells make up the valence shell and for inner transition elements, the highest  $s$ ,  $d$ , and  $f$  subshells are included

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