Chapter 8

Periodic Properties of the Elements

8.1 Outline

- History of the periodic table
- Effective nuclear charge
- Sizes of atoms and ions
- Trends in ionization energies
- Trends in electron affinities

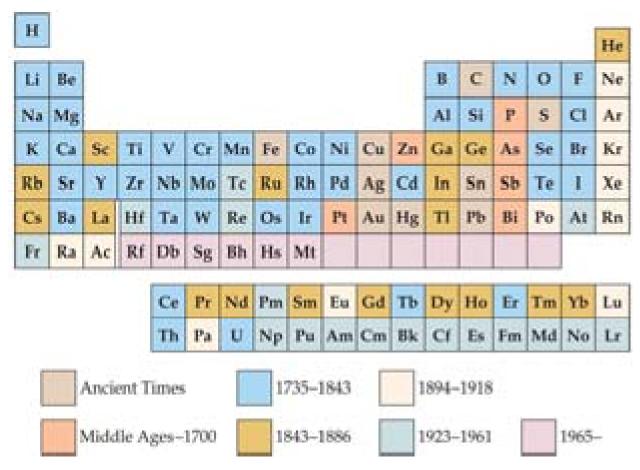


Figure 8.1: Discovery Dates of the Elements

8.2 Development of the Periodic Table

- Mendeleev's insistence that elements with similar properties be listed in the same group lead him to leave several blanks in the periodic table.
- For example, Mendeleev predicted some properties of now what is called Germanium based on the fact that it is in the same group as Silicon. Silicon was discovered almost 100 years before that of Germanium!
- Once germanium was discovered, its observed properties matched exceptionally well with Mendeleev's predictions (see the table on the next slide).

Table 8.1: Comparison of the Properties of Eka-Silicon ("under" silicon) Predicted by Mendeleev with the Observed Properties of Germanium

Property		Observed Properties of
	tions for Eka-Silicon (made in 1871)	Germanium (discovered in 1886)
Atomic weight	72	72.59
Density (g/cm^3)	5.5	5.35
Specific heat $(J/g \times K)$	0.305	0.309
Melting point (°C)	High	947
Color	Dark gray	Grayish white
Formula of oxide	XO_2	GeO_2
Density of oxide (g/cm^3)	4.7	4.70
Formula of chloride	XCI ₄	$GeCl_4$
Boiling point of chloride	A little under 100	84
(°C)		

Periodic law 1860–1870's (Mendeleev and Meyer) – A periodic repetition of physical and chemical properties occurs when the elements are arranged in order of increasing atomic $weight \ [number]$

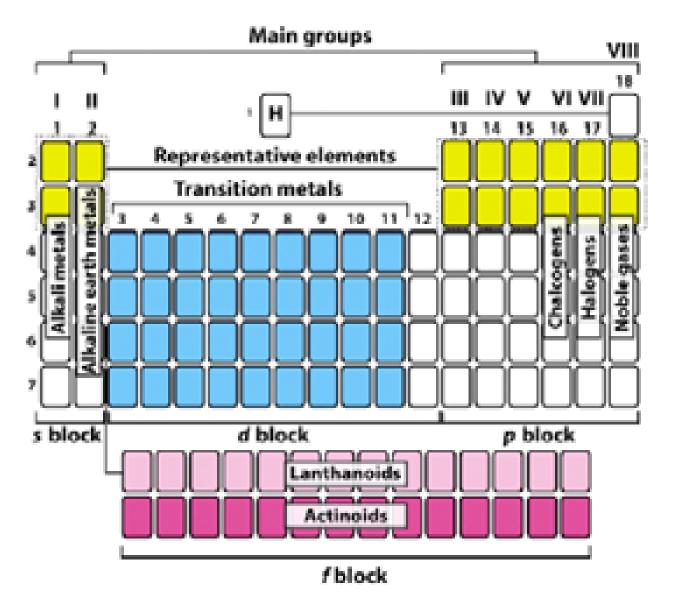


Figure 8.2

8.3 Ordering by Atomic Weight

Inconsistencies in ordering by atomic weight:

- $\bullet\,$ Co (58.93 amu; Z=27) and Ni (58.69 amu; Z=28)
- $\bullet\,$ Ar (39.95 amu; Z=18) and K (39.10 amu; Z=19)
- $\bullet\,$ Te (127.60 amu; Z=52) and I (126.90 amu; Z=53)

However, all of the above are correctly ordered by atomic number, Z (i.e., the number of protons).

8.4 Development of Periodic Table

- Elements in the same group generally have similar chemical properties.
- However, physical properties are not necessarily similar.
- For example, even though Oxygen and Sulfur are in the same group (6A), Oxygen is a colorless gas, while Sulfur is a yellow solid under normal conditions.

8.5 But why do elements in the same group have similar properties?

- Underlying valence electronic structure is the same (almost) for elements of a group:
 - e.g., alkaline earth metals (Group 2) metals are all

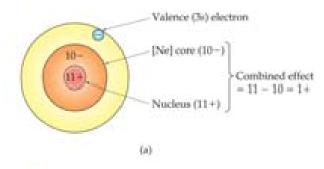
[Noble Gas] ns²

- halogens (Group 17) are all

[Noble Gas] ns²np⁵

- So, broadly speaking, elements in the same group should have similar chemistry
- But why do properties like atomic size, ionization energies and electron affinity vary from group to group?.

8.6 Effective Nuclear Charge



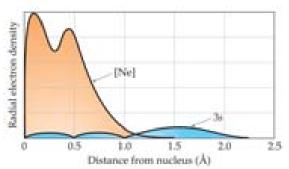


Figure 8.3: Effective Nuclear Charge

- In a many-electron atom, electrons are both attracted to the nucleus and repelled by other electrons.
- The nuclear charge that an electron experiences depends on both factors.
- The effective nuclear charge, Z_{eff} , is found this way:

$$Z_{eff} = Z - \sigma \tag{8.1}$$

where Z is the atomic number and σ is the screening constant, usually close to the number of inner/core electrons.

- Z_{eff} is always less than Z (atomic number) due to shielding!
- Electrons in the same shell don't screen one another very well.

8.7 What is the Size of an Atom?

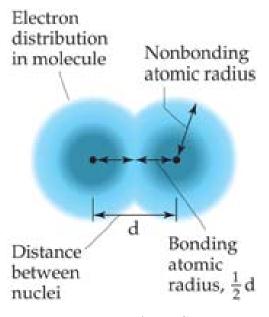


Figure 8.4: Atom Size

• The bonding atomic radius is defined as $\frac{1}{2}$ of the distance between covalently bonded nuclei.

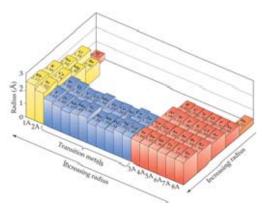


Figure 8.5: Sizes of Atoms

- Bonding atomic radii tend to:
 - (A) decrease from left to right across a row (due to increasing Z_{eff}).
 - (B) increase from top to bottom of a column (due to increasing value of n).

8.8 Estimating Bond Lengths

- Atomic radii allows one to estimate bond lengths.
- \bullet Example, estimate the carbon-chlorine bond length in carbon tetrachloride, CCl₄. Atomic radius of C: 0.77 Å Atomic radius of Cl: 0.99 Å Intenuclear bond length = 0.77 Å + 0.99 Å = 1.76 Å
- The experimentally determined carbon-chlorine bond length in carbon tetrachlroide is actually 1.77 Å.

8.9 Radii of Cations and Anions

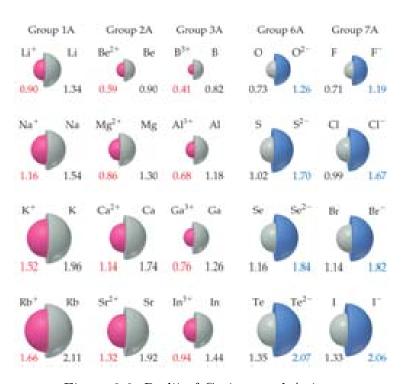


Figure 8.6: Radii of Cations and Anions

- Cations are smaller than parent neutral atoms (e.g., Li⁺ is smaller than Li.) Why?
- Increased Z_{eff} upon electron removal.
- Anions are larger than parent neutral atoms (e.g., I⁻ is larger than I.) Why?
- Increased electron-electron repulsions upon electron addition, causing electrons to spread out more in space.
- Ions with the same charge ion size increases going down a group due to increasing values of *n*.
- For example, a Sr^{2+} ion (radius = 1.32 Å) is larger than a Ca^{2+} ion (radius = 1.14 Å).

8.10 Sizes of Isoelectronic Ions

- In an isoelectronic series, ions have the same number of electrons.
- Ionic size decreases with an increasing nuclear charge (Z_{eff}) .



Figure 8.7: Sizes of Isoelectronic Ions

8.11 Ionization Energy

• Ionization energy – energy required to remove an electron from a gaseuos species.

- The first ionization energy is the energy required to remove the first electron.
- The second ionization energy is the energy required to remove the second electron, etc.

$$\text{Mg} \overset{\mathit{IE}_1 = 738 \frac{\mathit{kJ}}{\mathit{mol}}}{\longrightarrow} \text{Mg}^+ \overset{\mathit{IE}_2 = 1451 \frac{\mathit{kJ}}{\mathit{mol}}}{\longrightarrow} \text{Mg}^{2+}$$

- It requires more energy to remove each successive electron.
- When all valence electrons have been removed, the ionization energy takes a large leap.
- Ionization energies for all elements can be found at: https://www.webelements.com

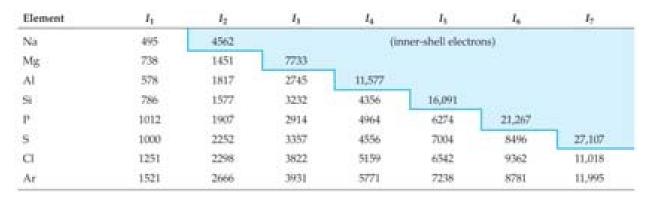


Figure 8.8: Ionization Energy

8.12 Trends in First Ionization Energies

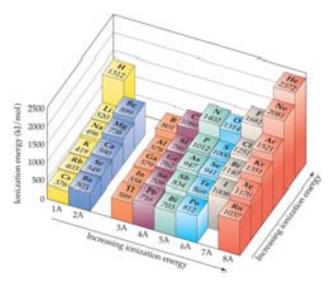


Figure 8.9: Periodic Table Ionization Energies

- As one goes down a group, less energy is required to remove the first electron.
 - For atoms in the same group, Z_{eff} is essentially the same, but the valence electrons are farther from than nucleus (i.e., increasing value of n).

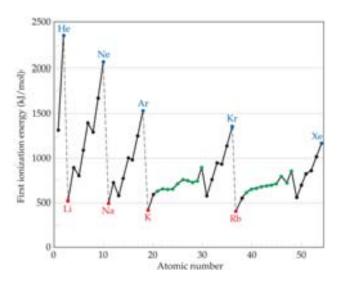


Figure 8.10: Trends in Ionization Energy

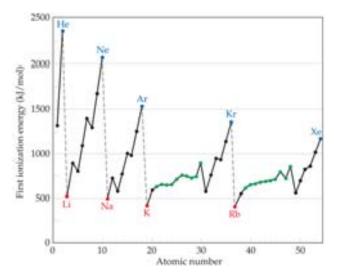


Figure 8.11: Discontinuities in Trends in Ionization Energy

- Generally, as one goes across a row/period, it becomes more difficult to remove an electron.
 - As you go from left to right $\Rightarrow Z_{eff}$ increases!
- However, there are two apparent discontinuities in this trend.
- The first occurs between Groups IIA (2) and IIIA (13).
- In this case, the electron is removed from a *p*-orbital rather than an *s*-orbital.
 - The electron removed is further from the nucleus.
 - There is also a small amount of repulsion by the s electrons.
- The second occurs between Groups VA (15) and VIA (16).
 - The electron removed comes from a doubly occupied orbital.
 - Repulsion from the other electron in the orbital aids in the removal.

8.13 Electron Affinity

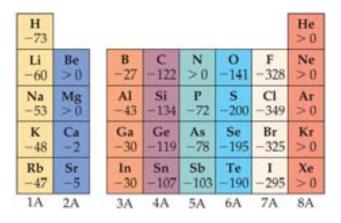
Electron affinity is the energy change accompanying the addition of an electron to a gaseous atom:

$$\mathsf{CL}(\mathsf{g}) + \mathsf{e}^- \longrightarrow \mathsf{CI}^-(\mathsf{g}) \qquad E_a = -349 \frac{\mathrm{kJ}}{\mathrm{mol}}$$

Energy is typically released when an electron is added to a gaseous atom. The process is said to be exothermic, so the energy has a negative sign associated with it.

The electron affinity of lithium is a negative value, whereas the electron affinity of Beryllium is a positive value. Use electron configuration to account for this observation.

8.14 Trends in Electron Affinity



• In general, electron affinities becomes more exothermic/more negative as you go from left to right across a period.

Figure 8.12: Trends in Electron Affinity

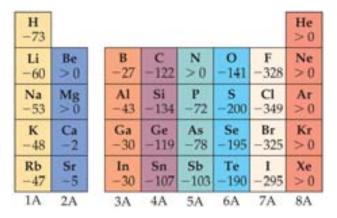


Figure 8.13: Discontinuities in Trends in Electron Affinity

• In general, electron affinities becomes more exothermic/more negative as you go from left to right across a period.