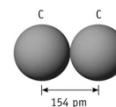


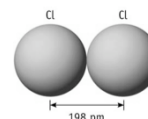
Periodic Properties and Trends

Chapter 8

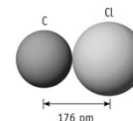
Atomic Radius (Size)



- The atomic radius of an element can be defined as half the distance between the nuclei of a diatomic molecule of the element.



- For example, the atomic radius of carbon is 77 picometers ($7.7 \times 10^{-11} \text{ m}$).



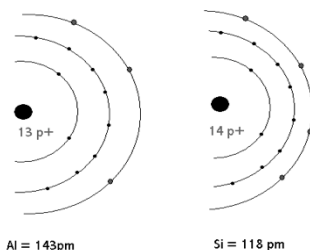
We might expect the atomic radii of elements to increase as we move from LEFT to RIGHT across the periodic table. However, experimentally, this is not observed. **Why not?**

Effective nuclear charge (Z_{eff}) and Shielding

- $Z_{\text{effective}}$ is the average shielded charge felt by an electron.
- Shielding** in an atom is the “blocking” or canceling of positive charge by core electrons.

- As we move from **left** \rightarrow **right** on the periodic table, shielding of the nucleus by inner (core) electrons remains constant. (Shielding by other outer electrons is minimal.)
- However, as the number of protons increases, the electrons in the valence (outer) shell experience a greater attraction for the nucleus – a higher $Z_{\text{effective}}$.

- Because of the greater pull on the outer electrons as we move across the periodic table, atomic radius generally **DECREASES** from **left** \rightarrow **right** within a period.



Atomic Radii (pm)

As we move from **top** \rightarrow **bottom** on the periodic table, we would expect the atomic radius to **INCREASE**, and it does, because the energy level of the outer e⁻s increases.

Ionic Radius

- Reported Ionic radii are determined from the measured distances between nuclei in ionic compounds.
- **Cations** are always *smaller* than the atom they are derived from. In many cases, the outer energy level of the cation is one energy level lower than that of the atom.
- **Anions** are always *larger* than the atom they are derived from, primarily due to repulsive forces of electrons in the outer energy level.

Ionic Radii
(pm)

- The **L→R** trend for ionic radius is **DECREASING** for the same type of ion – cation or anion.
- The **T→B** trend is **INCREASING**.

Examples: Atomic and Ionic Radii

- Put the following species into order of **DECREASING** particle size:

1. P P³⁻ S S²⁻

2. Rb Rb⁺ Sr I I⁻

3. B C Al

Ionization Energy

- **First Ionization Energy** – the energy required to remove the highest energy electron from an atom.
- **Second Ionization Energy** – the energy required to remove the second highest energy electron (from the (+1) ion).
- **Third, Fourth, Fifth, etc.**
- Ionization energies are positive, as they refer to the energy going into the atom (system) to remove an electron.

1st
Ionization
Energy

- **L→R** trend: 1st ionization energy **INCREASES**, because the pull on the outermost electron increases.
- **T→B** trend: 1st ionization energy **DECREASES**, because the outermost electron is at a higher energy level.

2nd Ionization Energy and higher...

A "jump" between subsequent ionization energies is observed when a **STABLE** electron configuration is reached.

Beryllium 1st ionization:

Beryllium 2nd ionization:

Element	First	Second	Third	Fourth	Fifth	Sixth	Seventh
H	1312						
He	2372	5250					
Li	520	7298	11,815				
Be	899	1757	14,848	21,006			
B	801	2427	3660	25,025	32,826		
C	1086	2353	4620	6222	37,829	47,276	
N	1402	2857	4578	7475	9445	53,265	64,358
O	1314	3388	5300	7469	10,989	13,326	71,333
F	1681	3374	6020	8407	11,022	15,164	17,867
Ne	2081	3952	6122	9370	12,177	15,238	19,998

Examples – Ionization energy

- Put the following in order of increasing 1st ionization energy:
Li Be Na
- Where would magnesium fit in?
- Which of the following will have the largest jump in ionization energy between its 4th and 5th ionization energies?
Na Al Si S

Electron Affinity

- The energy released (negative value) when an atom of an element in the gas phase takes on an electron to become a negative ion.
- L→R trend:** Generally increasing, because the attraction for the outermost electrons increases from left to right.
- T→B trend:** No reliable trend. Generally decreasing, but trend is not highly predictable from top to bottom.

Table of Electron Affinities

Period	IA	IIIA	IVA	VA	VIA	VIIA
1	H -73					
2	Li -60	B -27	C -122	N 0	O -141	F -328
3	Na -53	Al -44	Si -134	P -72	S -200	Cl -349
4	K -48	Ga -30	Ge -120	As -77	Se -195	Br -325
5	Rb -47	In -30	Sn -121	Sb -101	Te -190	I -295
6	Cs -45	Tl -30	Pb -110	Bi -110	Po -180	At -270

*Atoms of the alkaline earth metals (Group IIA) and the noble gases (Group VIIIA) do not form stable negative ions.

Electronegativity

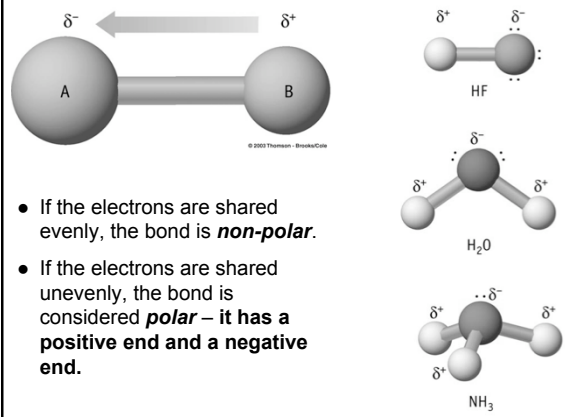
- A measure of the relative attraction that a nucleus has on electrons in a chemical bond - the larger the number, the greater the attraction.
- L→R trend: INCREASES**
- T→B trend: DECREASES**
- If two atoms are in a bond, the difference in their electronegativities will determine if the electrons are shared evenly or unevenly.

Electronegativity and Bond Polarity

- The greater the electronegativity difference (ΔEN) between two atoms in a bond, the more polar the bond.
- If the difference is very great, then one atom essentially pulls the electrons away from the other making an ionic bond.
- There is a continuum of non-polar to polar to ionic compounds, with no clear breaks.
- However, we will define somewhat arbitrary cut-off points:

$\Delta EN \leq 0.4$	Non-polar covalent
$0.5 \leq \Delta EN \leq 1.6$	Polar covalent
$\Delta EN \geq 1.7$	Ionic

The cutoff for "ionic" is the point at which a bond has 50% ionic character.



Summary of Periodic Properties

<i>Property</i>	<i>L→R</i>	<i>T→B</i>
$Z_{\text{effective}}$		
Shielding		
Electron Energy Level		
Atomic Radius		
Ionic Radius		
Ionization Energy		
Electron Affinity		
Electronegativity		