- Giving the four parameters will uniquely identify an electron around an atom. No two electrons in the same atom can share all four. These parameters are called QUANTUM NUMBERS.

) PRINCIPAL QUANTUM NUMBER (n):

- "energy level", "shell"

- Represents two things:

* The distance of the electron from the nucleus.

* Energy. "n" is one factor that contributes to the energy of the electron.

$$n = 1, 2, 3, 4, ...$$
 (integers)

.) ANGULAR MOMENTUM QUANTUM NUMBER: $\, \ell$

- "subshell"
- Represents the SHAPE of the region of space where the electron is found.

- (Bohr assumed CIRCULAR orbits for electrons ... but there are more possibilities.)

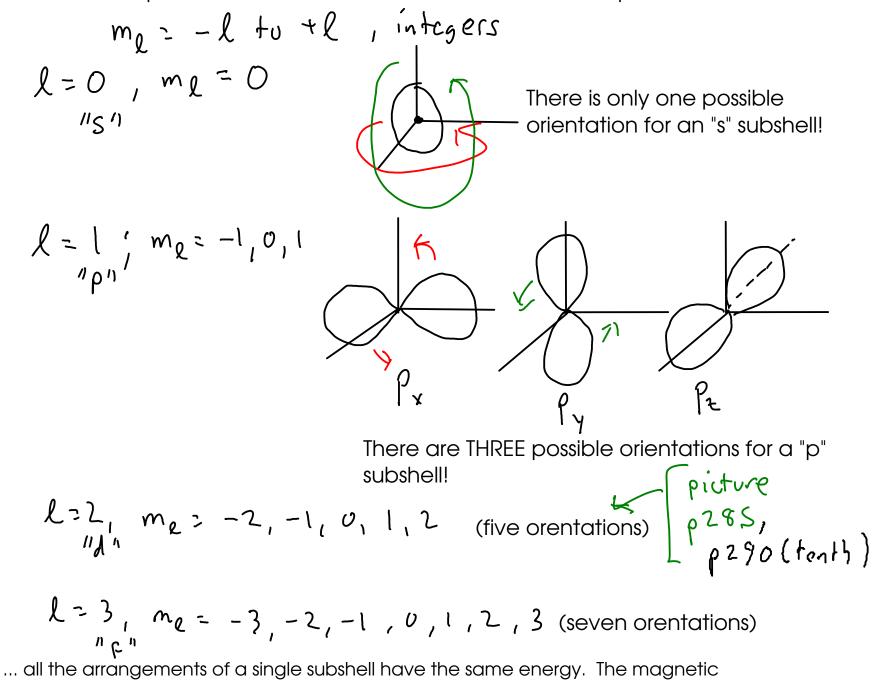
-"I" also contributes ENERGY. Higher values for "I" mean the electron has higher energy.

For

(3) MAGNETIC QUANTUM NUMBER M_{Q}

172

- Represents the ORIENTATION of a subshell in 3D space.



quantum number DOESN'T contribute to the energy of an electron.

(4) (MAGNETIC) SPIN QUANTUM NUMBER: m_{s}

$$M_{S} = \frac{1}{2} \circ \frac{R}{2} \frac{1}{2}$$
 "spin down" or "spin up"

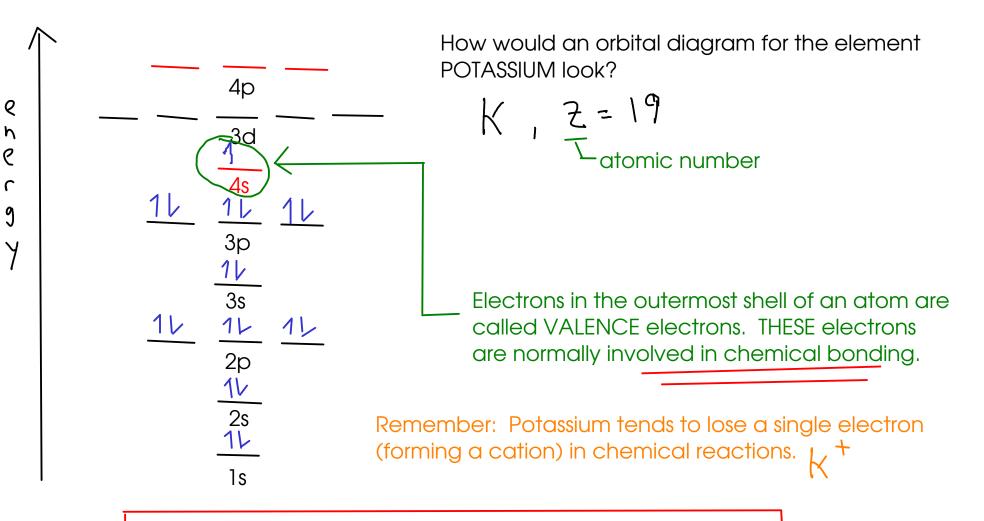
- An ORBITAL (region with fixed "n", "I" and "ml" values) can hold TWO electrons.

ORBITAL DIAGRAM

173

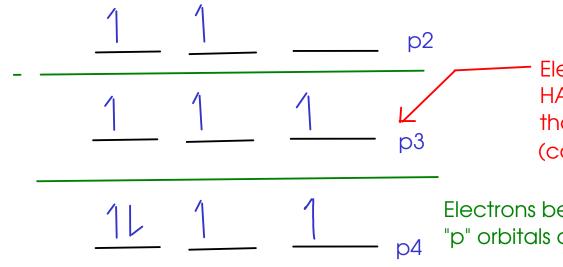
- A graphical representation of the quantum number "map" of electrons around an atom.

~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	4p Each blank represents an ORBITAL, and can hold two electrons.
	3d
	<ul> <li>The 4s subshell is lower energy than the 3d subshells remember</li> <li>4s that both n and "I" contribute to energy!</li> </ul>
9 7	3p Orbitals fill in order: The lower energy orbitals fill first.
	3s X Where there's more than one orbital with the same energy, electrons don't pair up until each one has
	2p one electron in it!
	2s
	1s ] n= \ she    "s" subshell ("1"=0)



A note on chemical bonding and electron arrangement: - Filled and half-filled subshells seem to be preferred by atoms. Hund's Rule

- When you have two or more orbitals with equivalent energy, electrons will go into each equivalent orbital BEFORE pairing. Pairing costs a bit of energy - less than going to a higher-energy orbital, but more than going to another equivalent orbital.



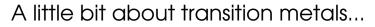
Electron configurations with filled subshells OR HALF-FILLED SUBSHELLS are more stable than other configurations. (can explain some transition metal chemistry)

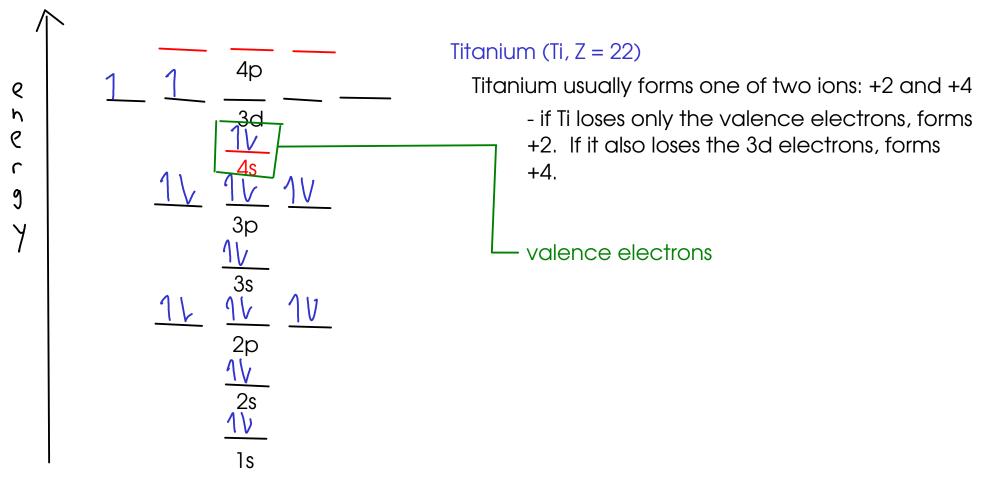
Electrons begin to pair only AFTER all equivalent "p" orbitals are full.

Experimental evidence for Hund's rule:

"Paramagnetism" - attraction of an atom to a magnetic field

- Spinning electrons are magnetic, but OPPOSITE spins cancel each other out.
- * Atoms with unpaired electrons are paramagnetic, while atoms containing only paired electrons are not.

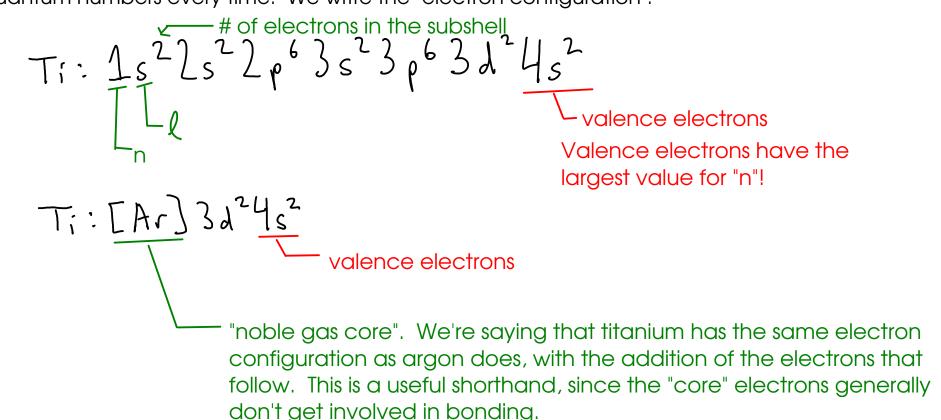


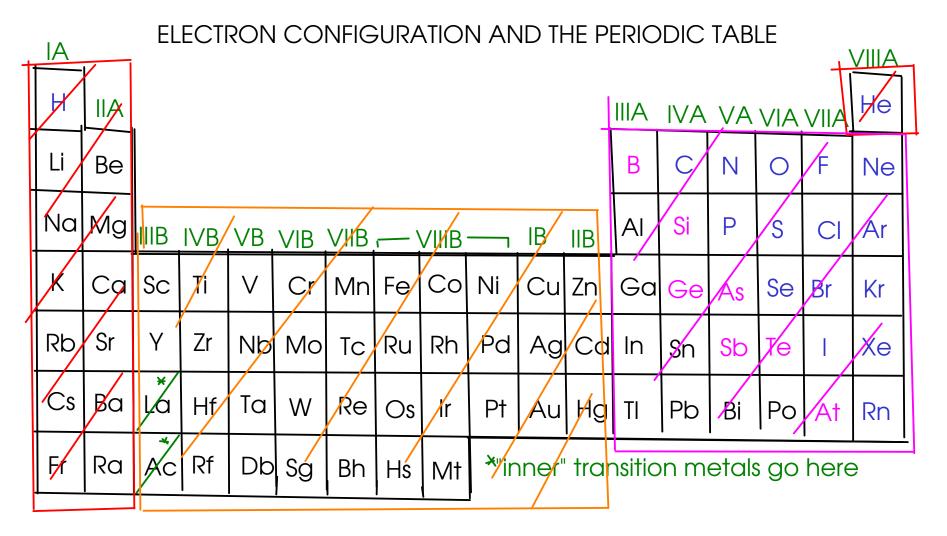


- Most transition metals have TWO valence electrons (in an "s" subshell), and the other ions they form come from electron loss in "d" subshells.

#### ELECTRON CONFIGURATION (SHORT FORM)

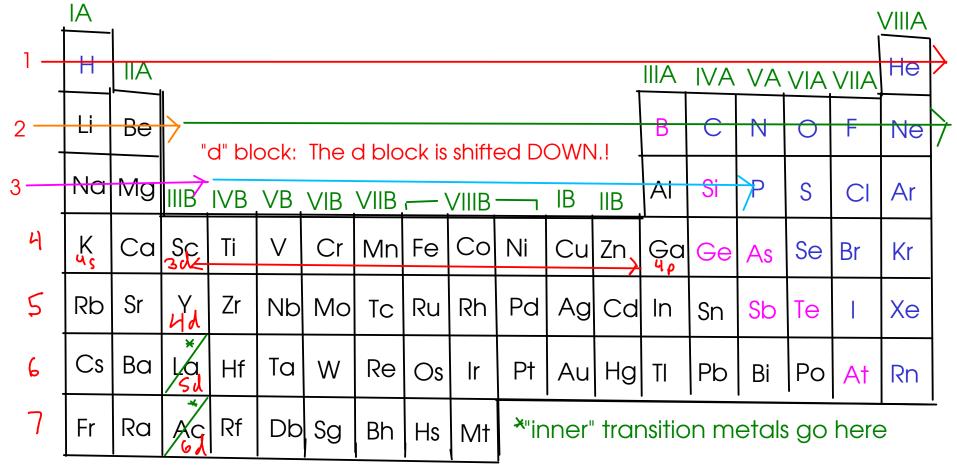
- We can represent the electron configuration without drawing a diagram or writing down pages of quantum numbers every time. We write the "electron configuration".





- "s" block: last electron in these atoms is in an "s" orbital!
- "p" block: last electron in these atoms is in a "p" orbital!
- "d" block: last electron in these atoms is in a "d" orbital

- To write an electron configuration using the periodic table, start at hydrogen, and count up the electrons until you reach your element!



Example: Phosphorus (P):  $\frac{1}{2}^{2}$   $\frac{2}{5}^{2}$   $\frac{2}{5}^{2}$   $\frac{2}{5}^{3}$ 

Noble gas core notation for P:  $[Ne]3s^23p^3$ 

EXAMPLES: $F \left[ s^{2} 2 s^{2} 2 \rho^{S} \right]$	Remember - valence electrons are ALL of the electrons in the outermost SHELL (n)! More that one subshell (I) may be included in the valence electrons	
s 1s ² 2s ² 2p ⁶ 3s ²	when the +2 ion forms, while the 4s AND 3d electrons are lost to form the +4!	
CI $ s^{2}2s^{2}2\rho^{6}3s^{2}3r^{5}$ CNe] $3s^{2}3\rho^{5}$ Ti $ s^{2}2s^{2}2\rho^{6}3s^{2}3r^{5}$	You can order the subshells in numeric order OR in filling order $p^{6}3d^{2}4s^{2}$ or $1s^{2}2s^{2}2p^{6}3s^{2}3p^{6}4s^{2}3d^{2}$	
Se $ s^2 2s^2 2p^6 3s^2 3p^6 3a^{10} 4s^2 4p^4$ $\int Ar [Ar] 3a^{10} 4s^2 4p^4$ Noble gas core notation. Use the previous noble gas on the table, then add the electrons that it doesn't have to the end.		
Kr [Ar] 3a104s24p6 $Ce:[$	Sample f-block element $xe ] 6c^2 Sd^1 4f^1$	

### PERIODIC TRENDS

- Some properties of elements can be related to their positions on the periodic table.

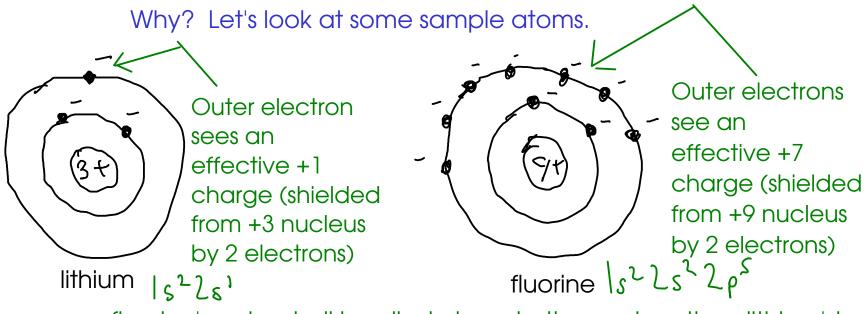
ATOMIC RADIUS

- The distance between the nucleus of the atoms and the outermost shell of the electron cloud.

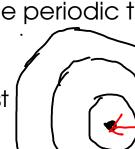
- Relates to the size of the atom.
- As you go DOWN A GROUP (  $\sqrt{}$  ), the atomic radius INCREASES.

- Why? As you go down a period, you are ADDING SHELLS!

- As you go ACROSS A PERIOD ( $\longrightarrow$ ), the atomic radius DECREASES



... so fluorine's outer shell is pulled closer to the nucleus than lithium's!



182

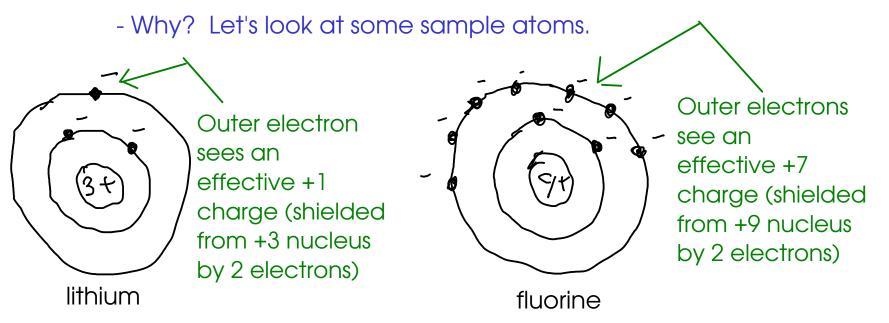
- The amount of energy required to remove a single electron from the outer shell of an atom.

- Relates to reactivity for metals. The easier it is to remove an electron, the more reactive the metal.

- As you go DOWN A GROUP (  $\int$  ), the ionization energy DECREASES.

- Why? As you go down a period, you are ADDING SHELLS. Since the outer electrons are farther from the nucleus and charge attraction lessens with distance, this makes electrons easier to remove as the atoms get bigger!

- As you go ACROSS A PERIOD (  $\longrightarrow$  , the ionization energy INCREASES.



... since fluorine's outer electrons are held on by a larger effective charge, they are more difficult to remove than lithium's.