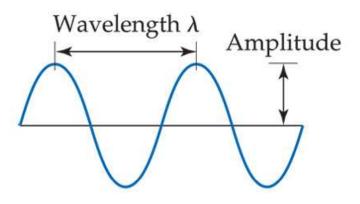
Chemistry, The Central Science, 10th edition Theodore L. Brown; H. Eugene LeMay, Jr.; and Bruce E. Bursten

Chapter 6 Electronic Structure of Atoms



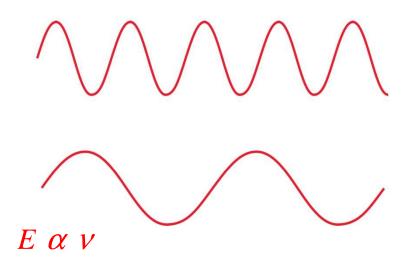
Waves



- To understand the electronic structure of atoms, one must understand the nature of electromagnetic radiation.
- The distance between corresponding points on adjacent waves is the wavelength (λ).



Waves



where α means 'is prapotionate to ' Therefore

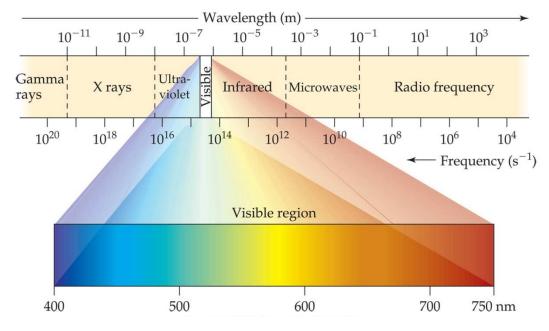
 $v\lambda = \text{constant}$

this constant is the speed of light

- The number of waves passing a given point per unit of time is the frequency (ν).
- For waves traveling at the same velocity, the longer the wavelength, the smaller the frequency.



Electromagnetic Radiation



- All electromagnetic radiation travels at the same velocity: the speed of light (*c*), 3.00 $\times 10^8$ m/s.
- Therefore,

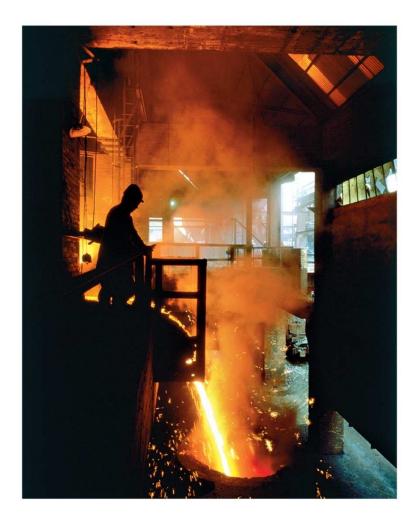
$$C = \lambda V$$



• Isaac Newton believed that light consists of beam of particles.

- Thomas Young shoes that light had the ability to diffract which is a wave property.
- Einstein postulated that light had both wave and particulate properties.





- When solids are heated they emit radiation.
- Red hot objects give out red light and at higher temperatures white light is emitted.
- Max Planck explained this phenomenon by making an assumption that in light energy comes in packets called quanta.



• He concluded that energy is proportional to frequency:

 $E \alpha v$

E = hv

where *h* is Planck's constant, = 6.63×10^{-34} J-s.



• The units for planks constant:

E = hv

$$h = \frac{E}{v} = \frac{J}{/s} = J.s$$



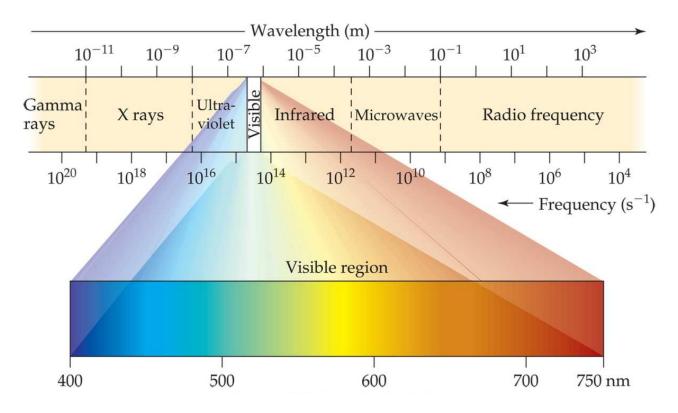
 Therefore, if one knows the wavelength of light, one can calculate the energy in one photon, or packet, of that light:

$$c = \lambda v$$

Gives us the frequency E = hvGives us the energy







You can calculate the energy of the photon of every kind of wave as long as you know that

> h = 6.63×10^{-34} J-s.and c= 3.0×10^8 m/s



- On Monday we were calculating the energy in photon from the visible light that had the wavelength of 400nm.
- We know that $\lambda v = c$

So $v = c / \lambda = \frac{3 \times 10^8 \text{m/s}}{400 \text{nm} \times \frac{1 \text{m}}{10^9 \text{nm}}}$ =7.5 x 10¹⁴ /s $E = h v = 6.63 \times 10^{-34} \text{ J-s x } 7.5 \times 10^{14} \text{ /s}$ = 4.9725x10⁻¹⁹ J = 5 x10⁻¹⁹ J

- The Si unit for frequency is Hz or Hertz.
- It is the same as /s.



 Max Plank in 1900 postulated that energy can either be released or absorbed by atoms in discreet chunks This is related to the frequency by the equation

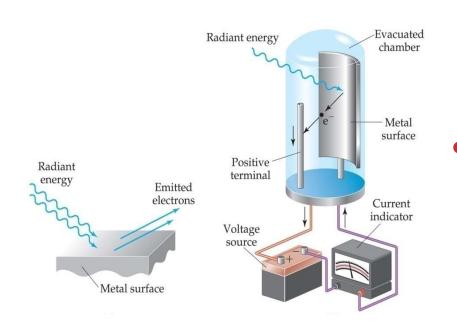
$$E = hv$$



- He also said that the atom can absorb energy in multiples of hv only.
- The smallest value of hv is called a quanta of energy



Photoelectric effect



- Einstein used Planck's theory to explain photoelectric effect.
- He said that the radiant energy striking the metal surface is behaving like a stream of tiny energy packets.



These packets are photons and behave like tiny particles

Einstein also deduced that each photon must have the energy equivalent to the Planck's constant times the frequency of that light.

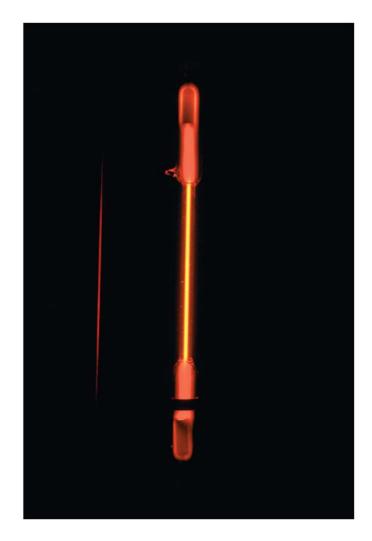
E = hv



Under the right conditions, a photon can strike the metal surface and be absorbed. Then it transfers it's energy to an electron in the metal. Once enough energy is absorbed, the electron is emitted from the metal

• The minimum energy to release the electron is different for different metals

Structure of Atoms

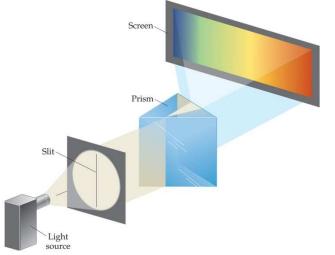


Another mystery involved the emission spectra observed from energy emitted by atoms and molecules.



Before we go any further
Monochromatic light:
The radiation has one single wavelength.
Spectrum:

when the radiation has many wavelengths.

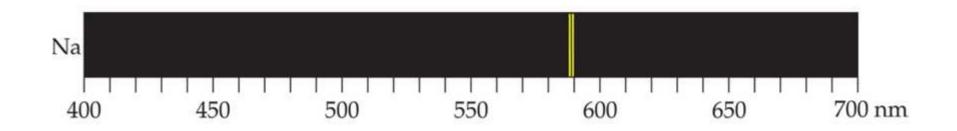




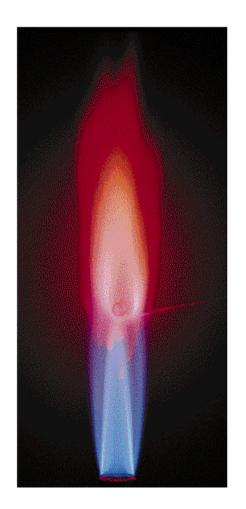
• When high voltage is applied to a tube containing a gas at reduced pressure, the gas emits different colors of light



Na vapor emits a light which when passed through a prism gives a single line at 590 nm wavelength in the yellow range.



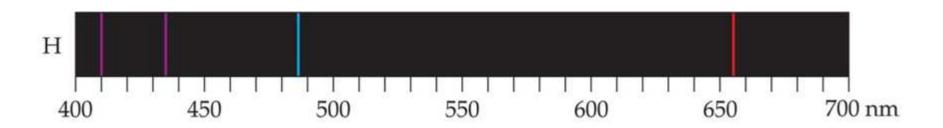








• For Hydrogen the spectrum has four distinct lines

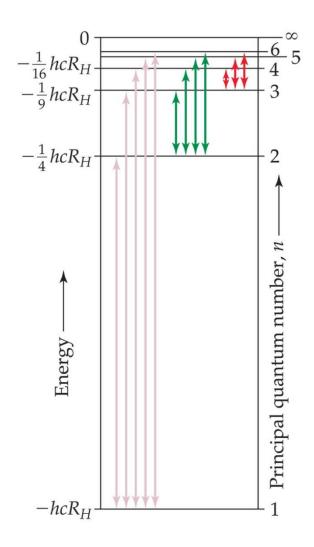




Bohr's Model:

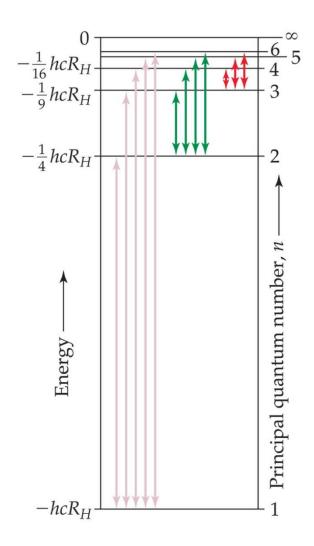
According to classical physics, electrically charged particles that are moving in a circular orbit should continuously be losing energy and collapse into the nucleus but this does not seem to happen. Bohr adapted Planck's idea that energies are quantized.





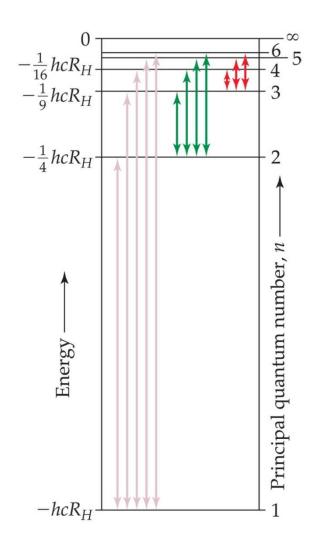
- Niels Bohr adopted Planck's assumption and explained these phenomena in this way:
 - Electrons in an atom can only occupy certain orbits (corresponding to certain energies).





- Niels Bohr adopted Planck's assumption and explained these phenomena in this way:
 - Electrons in permitted orbits have specific, "allowed" energies; these energies will not be radiated from the atom.





- Niels Bohr adopted
 Planck's assumption and
 explained these
 phenomena in this way:
 - Energy is only absorbed or emitted in such a way as to move an electron from one "allowed" energy state to another; the energy is defined by

$$E = hv$$



Bohr calculated the energy corresponding to each each orbit Uhere h = planck's Constant $E = <math>(-he R_H) \begin{pmatrix} 1 \\ n^2 \end{pmatrix} \qquad C = Speed of Light$ $R_H = Rydburg's Constant$ And the product of all thethree is 2.18 × 15¹⁸ J



N = principal quantum number and can have the values



Bohr accurred that electors could jump from one energy state to another A radient energy is cmitted when electron jumps from a sigher to a lower energy state $\Delta E = E_f - E_i = E_{photon} = h \nu$



$$\Delta E = h = \frac{h c}{\lambda} = (-2.18 \times 10^{18} \text{J}) (\frac{1}{n_f^2} - \frac{1}{h_1^2})$$
So if $n_i = 3$ and $n_f = 4$

$$\Delta E = (-2.18 \times 10^{18} \text{J}) (\frac{1}{1^2} - \frac{1}{3^2}) = -2.18 \times 10^{18} \text{J} (\frac{1}{-1})$$

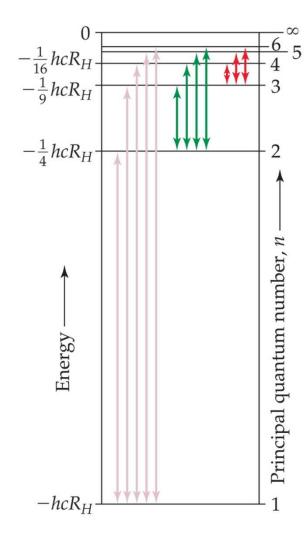
$$= -2.18 \times 10^{18} \text{J} (\frac{8}{-9})$$

$$= -1.94 \times 10^{-18} \text{J}$$



$$\lambda = \frac{\zeta}{V} = \frac{hC}{\Delta E} = \frac{(L \cdot LS \times 10^{-34} \text{ J} \text{ s})(3 \times 10^{-18} \text{ J})}{1 \cdot 94 \times 10^{-18} \text{ J}}$$
$$= 1 \cdot 93 \times 10^{-7} \text{ m}.$$





The energy absorbed or emitted from the process of electron promotion or demotion can be calculated by the equation:

$$E=-hcR_{H}\left(\frac{1}{n_{f}^{2}}-\frac{1}{n_{i}^{2}}\right)$$

where R_H is the Rydberg constant, 2.18 × 10⁻¹⁸ J, and n_i and n_f are the initial and final energy levels of the electron. Electronic Structure

of Atoms

imitations of Bohr's model

• It applied only to hydrogen atom



The Wave Nature of Matter

- Louis de Broglie posited that if light can have material properties, matter should exhibit wave properties.
- He demonstrated that the relationship between mass and wavelength was

$$\lambda = \frac{h}{mv}$$



The Uncertainty Principle

• Heisenberg showed that the more precisely the momentum of a particle is known, the less precisely is its position known:

$$(\Delta x) (\Delta mv) \geq \frac{h}{4\pi}$$

 In many cases, our uncertainty of the whereabouts of an electron is greater than the size of the atom itself!



Nora Tobin has completed her orientation and Student Employment paperwork and is now ready to tutor Chem 115 and 116. Students can register for tutoring in the Academic Support Programs Office (CC-1-1300).

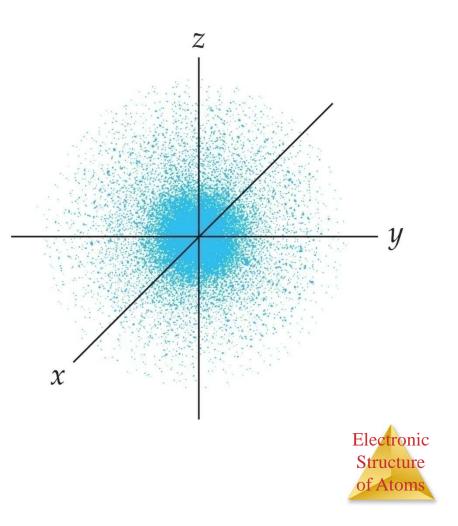


- With DeBrogie's correlation of electron and wave and Heisenberg's uncertainty principal the wave nature of electron was accepted
- Now the location of electrons is described in probability and the energy of an electron can be calculated.



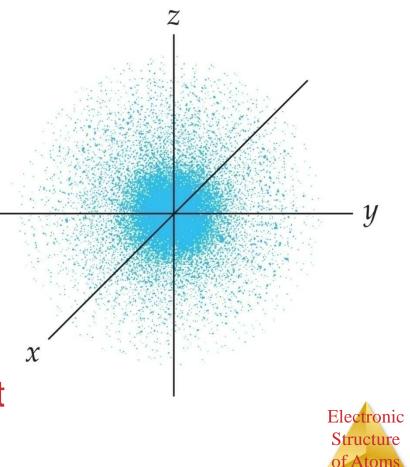
Quantum Mechanics

- Erwin Schrödinger developed a mathematical treatment into which both the wave and particle nature of matter could be incorporated.
- It is known as quantum mechanics.



Quantum Mechanics

- The wave function is designated with a lower case Greek *psi* (ψ).
- The square of the wave equation, \u03c8², gives a probability density map of where an electron has a certain statistical likelihood of being at any given instant in time.



Quantum Numbers

- Solving the wave equation gives a set of wave functions, or orbitals, and their corresponding energies.
- Each orbital describes a spatial distribution of electron density.
- An orbital is described by a set of three quantum numbers.



- Remember now the term being used will be orbital, not orbit the term used by Bohr
- Bore had only the quantum number n for orbit.
- The quantum mechanics uses four quantum numbers.



Principal Quantum Number, n

- The principal quantum number, *n*, describes the energy level on which the orbital resides.
- The values of *n* are integers 1, 2,3 and so forth



Azimuthal Quantum Number, /

- This quantum number defines the shape of the orbital.
- Allowed values of *I* are integers ranging from 0 to *n* 1.
- We use letter designations to communicate the different values of / and, therefore, the shapes and types of orbitals.



Azimuthal Quantum Number, /

Value of <i>I</i>	0	1	2	3
Type of orbital	S	p	d	f



Magnetic Quantum Number, m₁

- Describes the three-dimensional orientation of the orbital.
- Values are integers ranging from -/ to /: $-l \le m_l \le l.$
- Therefore, on any given energy level, there can be up to 1 *s* orbital, 3 *p* orbitals, 5 *d* orbitals, 7 *f* orbitals, etc.



• n=1, l=0, m=0

• n=2, I=0 m=0

- n=2 I = 1 m = -1
- n=2 I = 1 m = 0
- n=2 I = 1 m = +1



$$n=3 \qquad l=0 \qquad m=0 \qquad s$$

$$\begin{cases} n=3 \qquad l=1 \qquad m=-1 \\ n=3 \qquad l=1 \qquad m=0 \\ n=3 \qquad l=1 \qquad m=+1 \end{cases} p$$

$$\begin{cases} n=3 \qquad l=2 \qquad m=-2 \\ n=3 \qquad l=2 \qquad m=-1 \\ n=3 \qquad l=2 \qquad m=0 \\ n=3 \qquad l=2 \qquad m=+1 \\ n=3 \qquad l=2 \qquad m=+2 \end{cases} d$$



Magnetic Quantum Number, m₁

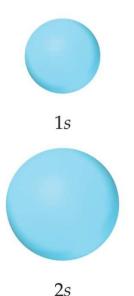
- Orbitals with the same value of *n* form a shell.
- Different orbital types within a shell are subshells. \bullet
- Magnetic quantum numbers represent orbitals.

n	Possible Values of <i>l</i>	Subshell Designation	Possible Values of m_l	Number of Orbitals in Subshell	Total Number of Orbitals in Shell	
1	0	1 <i>s</i>	0	1	1	
2	0	2 <i>s</i>	0	1		
	1	2p	1,0,-1	3	4	
3	0	3 <i>s</i>	0	1		
	1	Зр	1,0,-1	3		
	2	3d	2, 1, 0, -1, -2	5	9	
4	0	4s	0	1		
	1	4p	1,0,-1	3		
	2	4d	2, 1, 0, -1, -2	5		
	3	4f	3, 2, 1, 0, -1, -2, -3	7	16	Electroni
		-500				Structure

TT (1) 1

of Atom

s Orbitals

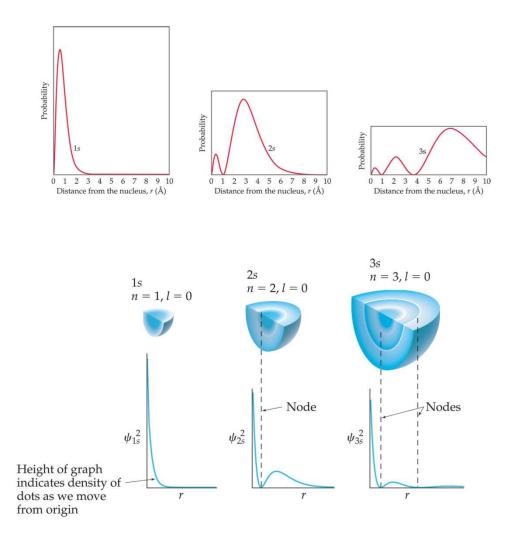




- Value of l = 0.
- Spherical in shape.
- Radius of sphere increases with increasing value of *n*.



s Orbitals

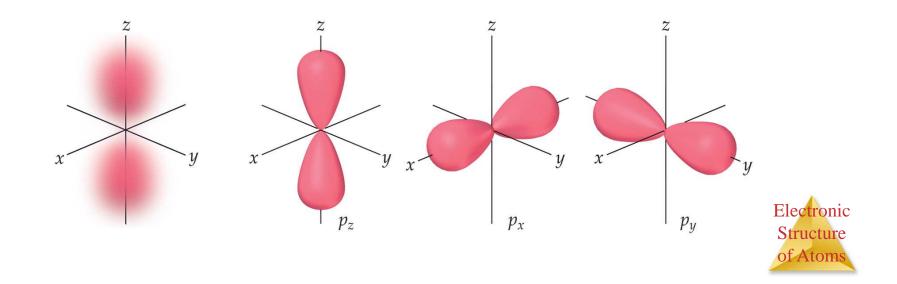


Observing a graph of probabilities of finding an electron versus distance from the nucleus, we see that *s* orbitals possess *n*–1 nodes, or regions where there is 0 probability of finding an electron.



p Orbitals

- Value of *I* = 1.
- Have two lobes with a node between them.

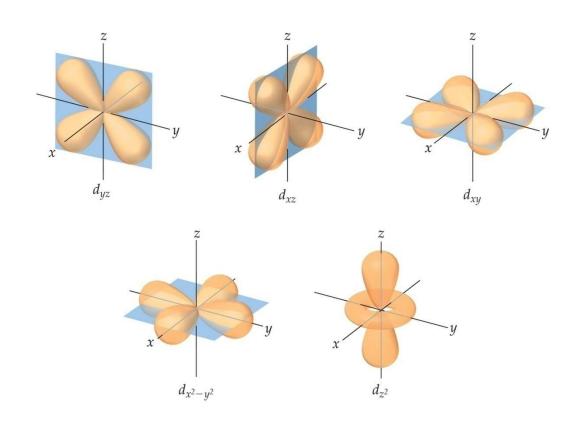


http://www.d.umn.edu/~pkiprof/ChemWebV2/AOs/ao2.html

and the second s



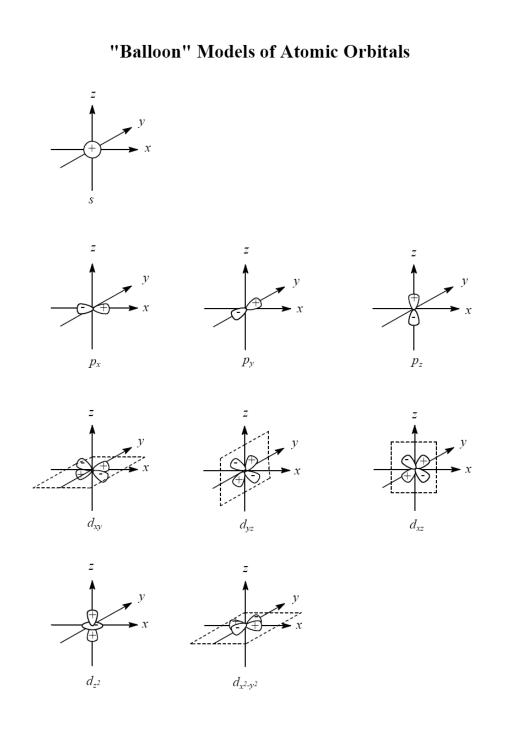
d Orbitals



• Value of / is 2.

• Four of the five orbitals have 4 lobes; the other resembles a *p* orbital with a doughnut around the center.





From Dr Carter's website.

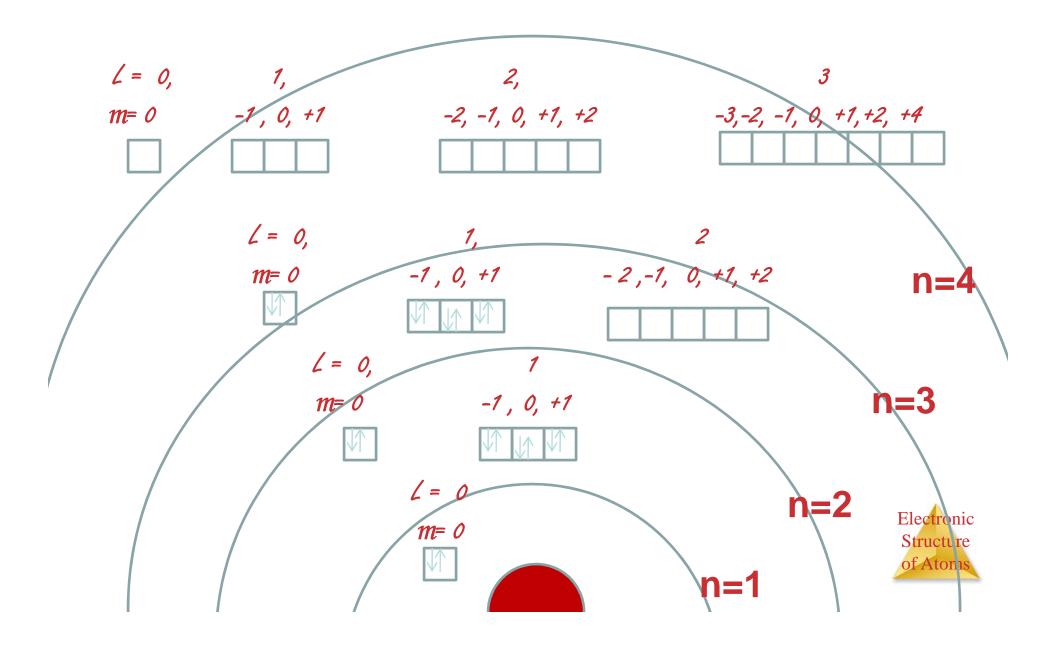


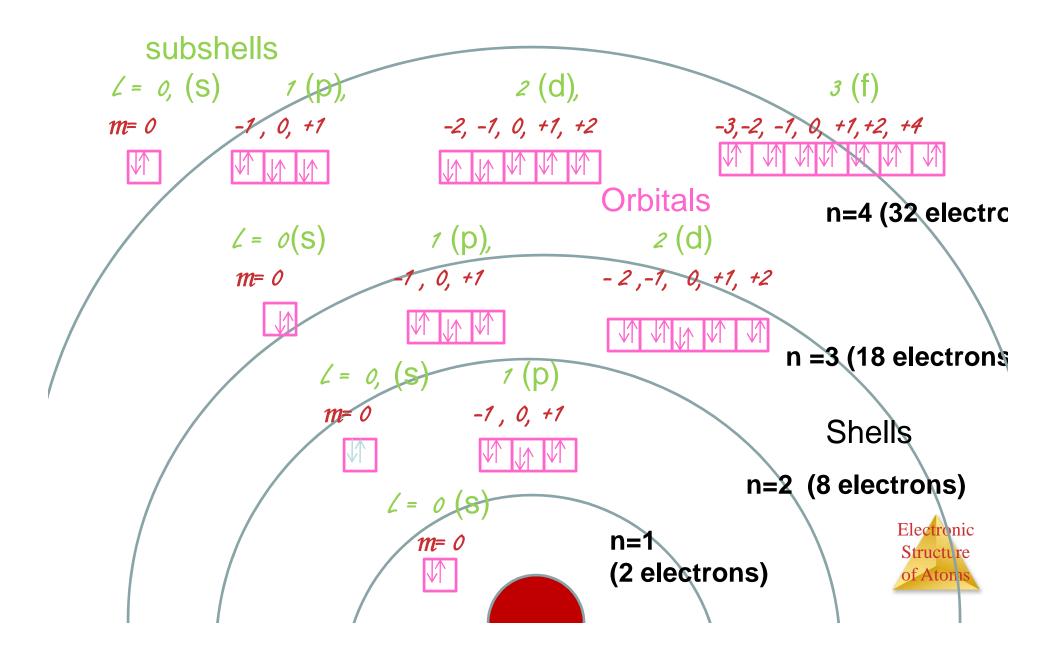
Shells n is a shell **Subshells** s, p d and f are sub shells **Orbitals** s has 1, p has 3, d has 5 and f has 7 orbitals.



- The number of orbitals in a each shell is n²
- n=1 orbital 1²= 1
- n=2 orbital 2^2 = 4 1s and 3p
- n=3 orbital 3^2 = 9 1s, 3p and 5d
- n=4 orbital 4²= 16 1s, 3p, 5d and 7f

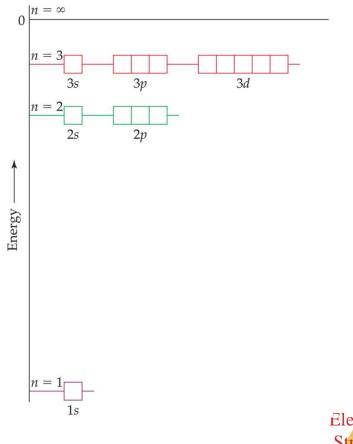






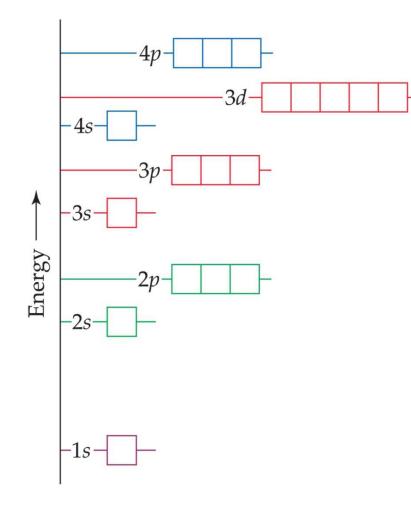
Energies of Orbitals

- For a one-electron hydrogen atom, orbitals on the same energy level have the same energy.
- That is, they are degenerate.



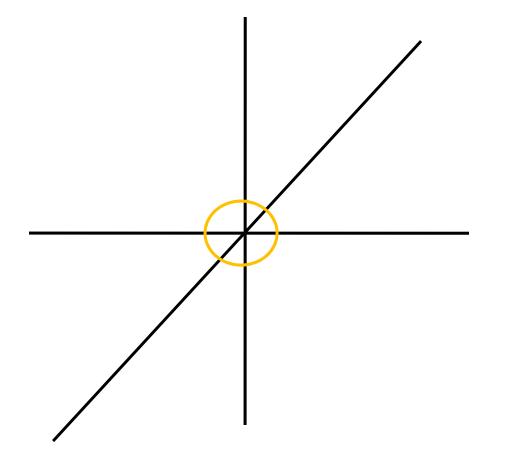


Energies of Orbitals

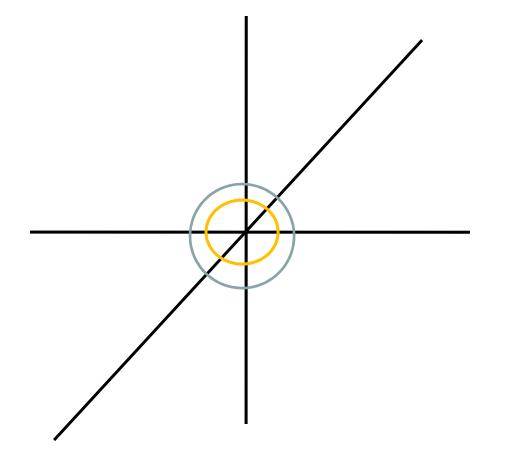


- As the number of electrons increases, though, so does the repulsion between them.
- Therefore, in manyelectron atoms, orbitals on the same energy level are no longer degenerate. Electronic

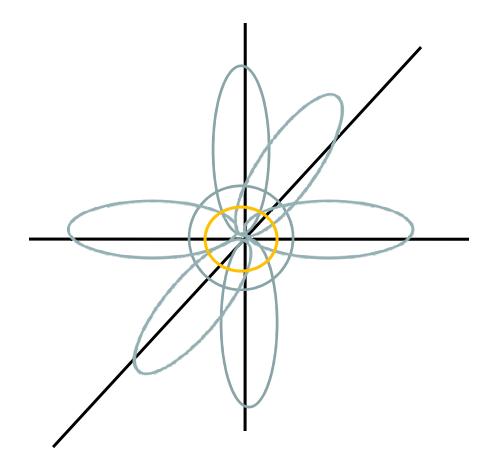
Structure of Atoms



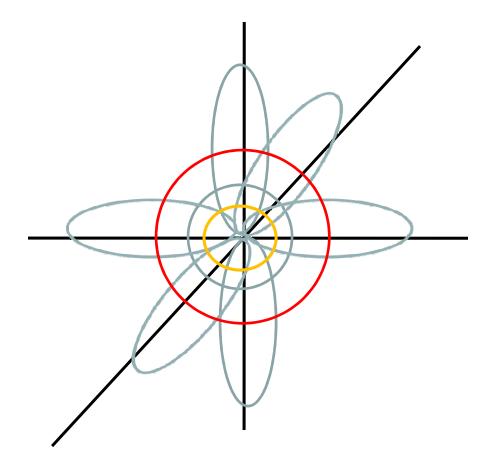




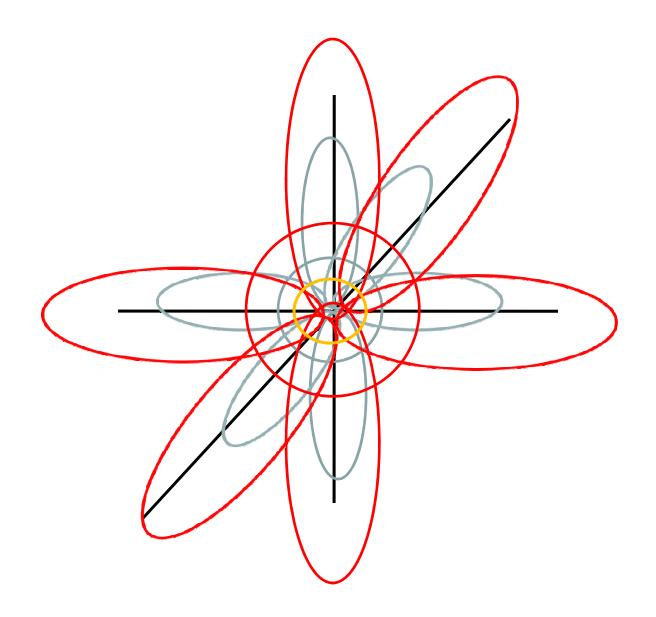




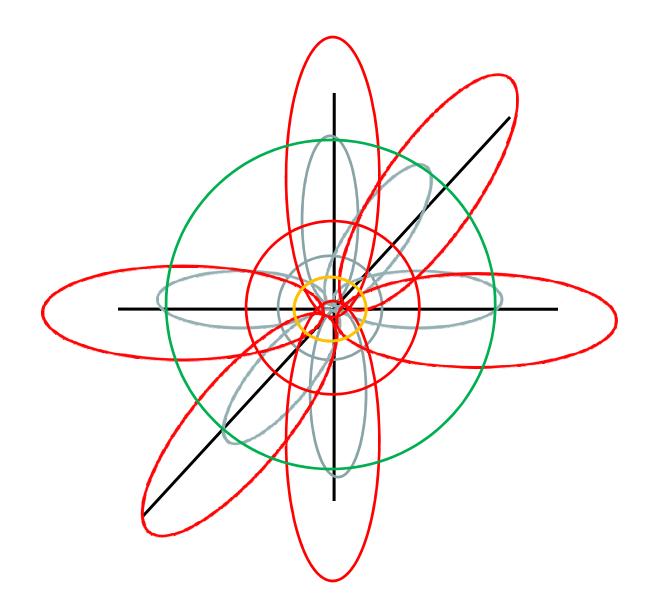




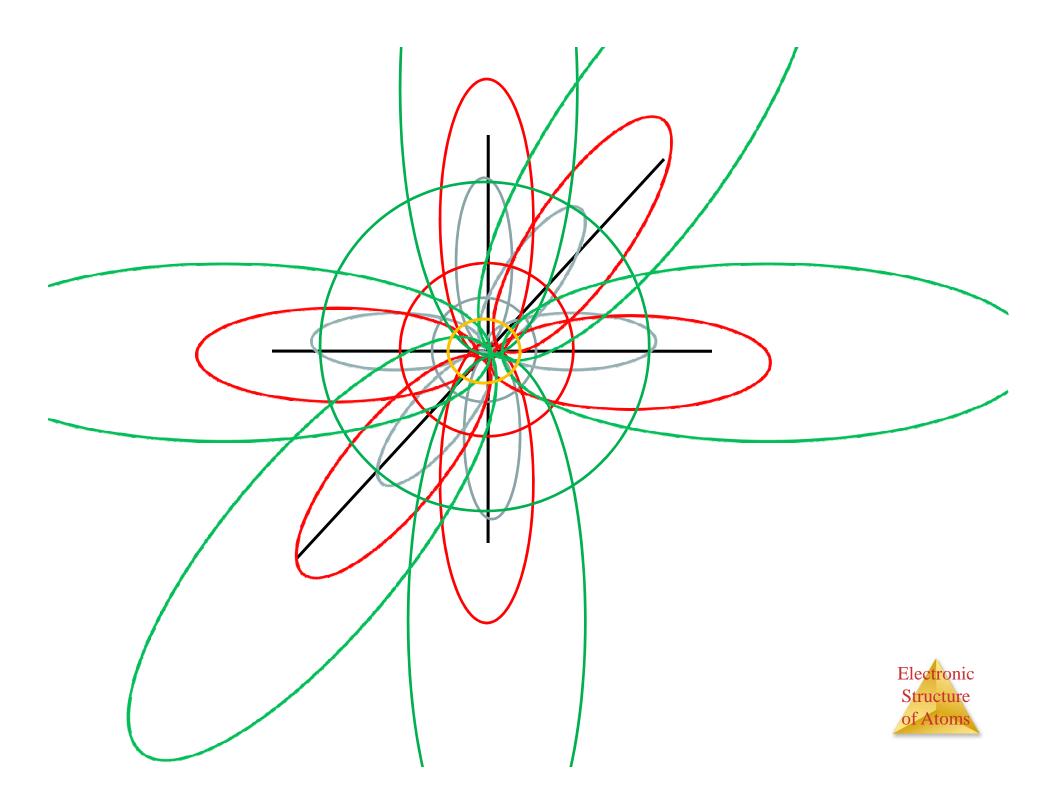






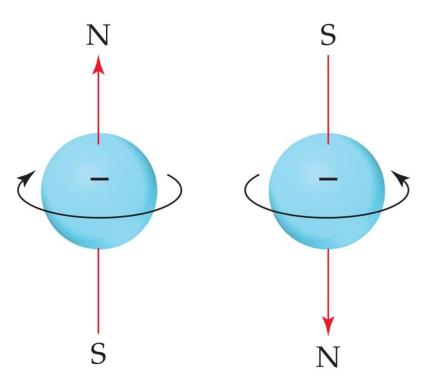






Spin Quantum Number, m_s

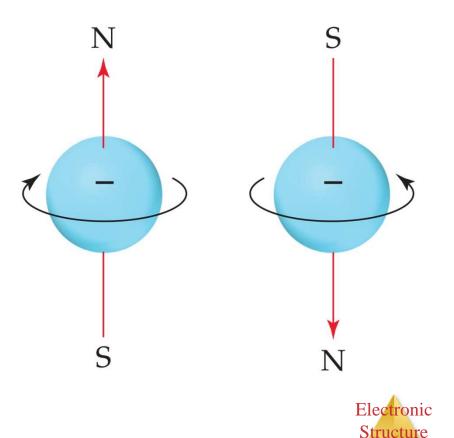
- In the 1920s, it was discovered that two electrons in the same orbital do not have exactly the same energy.
- The "spin" of an electron describes its magnetic field, which affects its energy.



Electronic Structure of Atoms

Spin Quantum Number, m_s

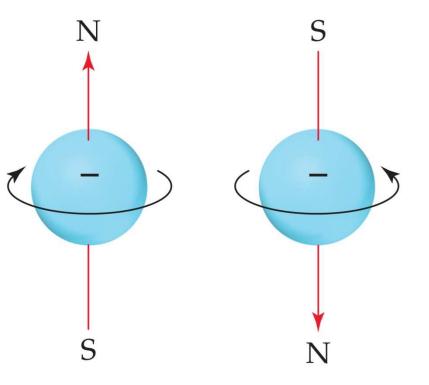
- This led to a fourth quantum number, the spin quantum number, m_s.
- The spin quantum number has only 2 allowed values: +1/2 and -1/2.



of Atoms

Pauli Exclusion Principle

- No two electrons in the same atom can have exactly the same energy.
- For example, no two electrons in the same atom can have identical sets of quantum numbers.



Electronic Structure of Atoms

Electron Configurations

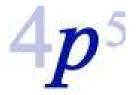


- Consist of
 - Number denoting the energy level





Electron Configurations



- Distribution of all electrons in an atom
- Consist of
 - Number denoting the energy level
 - Letter denoting the type of orbital



Electron Configurations

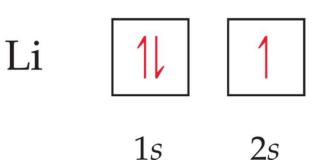


- Distribution of all electrons in an atom.
- Consist of
 - Number denoting the energy level.
 - Letter denoting the type of orbital.
 - Superscript denoting the number of electrons in those orbitals.



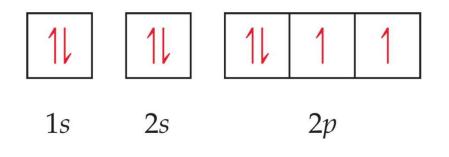
Orbital Diagrams

- Each box represents one orbital.
- Half-arrows represent the electrons.
- The direction of the arrow represents the spin of the electron.





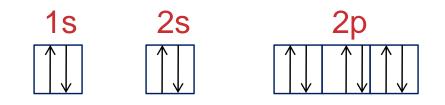
Hund's Rule



"For degenerate orbitals, the lowest energy is attained when the number of electrons with the same spin is maximized."



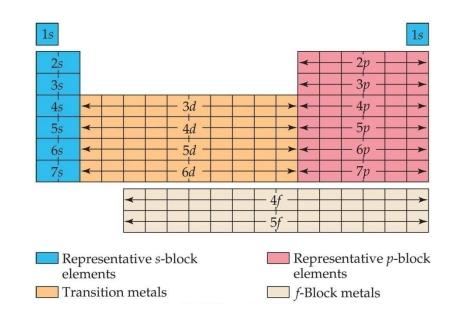
The sequence of filling of electrons in atoms will be:





Periodic Table

- We fill orbitals in increasing order of energy.
- Different blocks on the periodic table, then correspond to different types of orbitals.

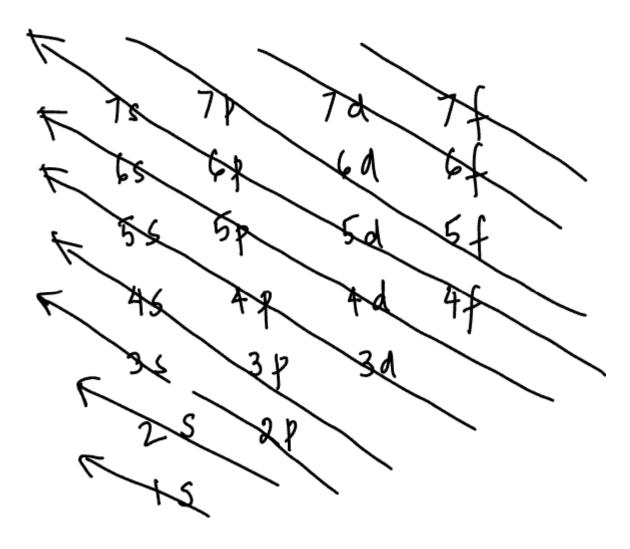




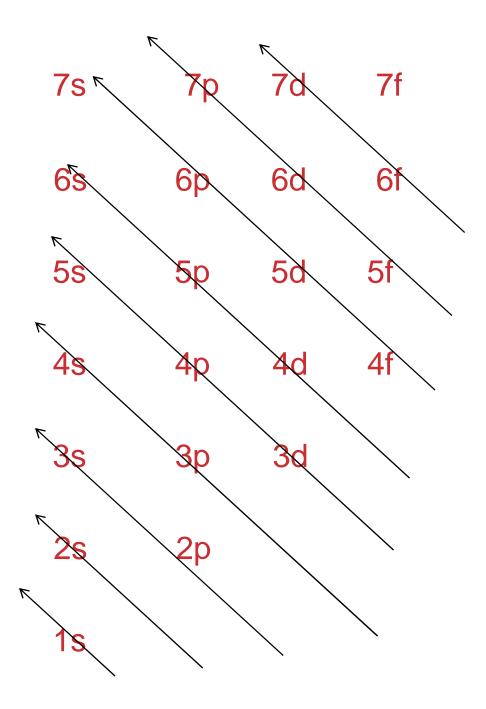
Aufbau Principle

- This is the building up principle of the electrons.
- Aufbau in German means building up.











						7s - 69	TR 6p	78 68	7f 61												
	1A 1	1				55	50	50	57									8A 18			
Core	$\begin{array}{c} 1\\ \mathbf{H}\\ 1s^1 \end{array}$	2A 2				45 *35	49 30	xd 3d	4f				3A 13	4A 14	5A 15	6A 16	7A 17	2 He 1s ²	1s	1s	
[He]	3 Li $2s^1$	4 Be 2s ²			ħ	26	2p						$ \begin{array}{c} 5\\ \mathbf{B}\\ 2s^22p^1 \end{array} $	$\begin{pmatrix} 6\\ C\\ 2s^2 2p^2 \end{pmatrix}$	$ \begin{array}{c} 7 \\ \mathbf{N} \\ 2s^2 2p^3 \end{array} $		9 F 2s ² 2p ⁵	$10 \\ Ne \\ 2s^2 2p^6$	2s2p	2sp	
[Ne]	11 Na 3s ¹	12 Mg 3s ²	3B 3	4B 4	5B 5	6B 6	7B 7	8	8B 9	10	1B 11	2B 12	$13 \\ A1 \\ 3s^2 3p^1$	$14 \\ Si \\ 3s^2 3p^2$	$15 \\ P \\ 3s^2 3p^3$			$18 \\ Ar \\ 3s^2 3p^6$	3s3p	3spd	
[Ar]	$19 \\ \mathbf{K} \\ 4s^1$	20 Ca 4s ²	21 Sc $3d^14s^2$	22 Ti 3d ² 4s ²	$\begin{array}{c} 23 \\ \mathbf{V} \\ 3d^3 4s^2 \end{array}$	$\begin{array}{c} 24 \\ \mathbf{Cr} \\ 3d^54s^1 \end{array}$	25 Mn $3d^54s^2$	26 Fe ^{3d⁶4s²}	27 Co $3d^{7}4s^{2}$	28 Ni 3d ⁸ 4s ²	$29 \\ Cu \\ 3d^{10}4s^1$	30 Zn $3d^{10}4s^2$	$31 \\ Ga \\ 3d^{10}4s^2 \\ 4p^1$	$32 \\ Ge \\ 3d^{10}4s^2 \\ 4p^2 \end{cases}$	33 As $3d^{10}4s^2$ $4p^3$	$34 \\ Se \\ 3d^{10}4s^2 \\ 4p^4$	$35 \\ Br \\ 3d^{10}4s^2 \\ 4p^5$	$36 \\ Kr \\ 3d^{10}4s^2 \\ 4p^6$	4s3d4p	4spdf	
[Kr]	37 Rb 5s ¹	38 Sr 5s ²	39 Y 4d ¹ 5s ²	$\begin{array}{c} 40\\ \mathbf{Zr}\\ 4d^25s^2\end{array}$	$\begin{array}{c} 41\\ \mathbf{Nb}\\ 4d^35s^2 \end{array}$	42 Mo $4d^{5}5s^{1}$	$43 \\ Tc \\ 4d^{5}5s^{2}$	$44 \\ \mathbf{Ru} \\ 4d^75s^1$	$45 \\ \mathbf{Rh} \\ 4d^{8}5s^{1}$	$\begin{array}{c} 46\\ \mathbf{Pd}\\ 4d^{10}\end{array}$	47 Ag $4d^{10}5s^1$	$48 \\ Cd \\ 4d^{10}5s^2$	49 In	50 Sn	$51 \\ 5b \\ 4d^{10}5s^2 \\ 5p^3$	52	$53 \\ I \\ 4d^{10}5s^2 \\ 5p^5$	54 34 $4d^{10}5s^2$ $5p^6$	5a4d4p	5spdf	
[Xe]	55 Cs _{6s¹}	56 Ba _{6s²}	71 Lu $4f^{14}5d^{1}$ $6s^{2}$	${{{\rm Hf}}\atop{{\rm 4}f^{14}5d^2}\atop{6s^2}}$	$73 \\ Ta \\ 4f^{14}5d^3 \\ 6s^2$	$\begin{matrix} 74 \\ \mathbf{W} \\ 4f^{14}5d^4 \\ 6s^2 \end{matrix}$	$75 \\ Re \\ 4f^{14}5d^5 \\ 6s^2$	$76 \\ Os \\ 4f^{14}5d^6 \\ 6s^2$	$77 \\ Ir \\ 4f^{14}5d^7 \\ 6s^2$	$78 \\ Pt \\ 4f^{14}5d^9 \\ 6s^1$	$79 \\ Au \\ 4f^{14}5d^{10} \\ 6s^{1}$	$80 \\ Hg \\ 4f^{14}5d^{10} \\ 6s^2$	$81 \\ f^{14} 5d^{10} \\ 6s^2 6p^1$	$82 \\ Pb \\ 4f^{14}5d^{10} \\ 6s^26p^2$	$83 \\ Bi \\ 4f^{14}5d^{10} \\ 6s^26p^3$	$84 \\ Po \\ 4f^{14}5d^{10} \\ 6s^26p^4$	$85 \\ At \\ 4f^{14}5d^{10} \\ 6s^26p^5$	$86 \\ \mathbf{Rn} \\ 4f^{14}5d^{10} \\ 6s^26p^6$	6s4f5d6p	6spdf	
[Rn]	87 Fr _{7s¹}	88 Ra 7s ²	103 Lr 5f ¹⁴ 6d ¹ 7s ²	$ \begin{array}{r} 104 \\ \mathbf{Rf} \\ 5f^{14}6d^2 \\ 7s^2 \end{array} $	$ \begin{array}{r} 105 \\ \mathbf{Db} \\ 5f^{14}6d^3 \\ 7s^2 \end{array} $	$ \begin{array}{r} 106 \\ Sg \\ 5f^{14}6d^4 \\ 7s^2 \end{array} $	$107 \\ Bh \\ 5f^{14}6d^5 \\ 7s^2$	$108 \\ Hs \\ 5f^{14}6d^6 \\ 7s^2$	$109 \\ Mt \\ 5f^{14}6d^7 \\ 7s^2$	110	111	112	113	114	115	116			7s5f6d	7spdf	
[Xe]	Lanth series	nanide		57 La $5d^{1}6s^{2}$	$58 \\ Ce \\ 4f^{1}5d^{1} \\ 6s^{2}$	$59 \\ \mathbf{Pr} \\ 4f^3 6s^2$	$\begin{array}{c} 60\\ \mathbf{Nd}\\ 4f^46s^2 \end{array}$	$61 \\ \mathbf{Pm} \\ 4f^56s^2$	$62 \\ Sm \\ 4f^{6}6s^{2}$	$63 \\ Eu \\ 4f^76s^2$	$64 \\ Gd \\ 4f^{7}5d^{1} \\ 6s^{2}$	$65 \\ Tb \\ 4f^{9}6s^{2}$	$ \begin{array}{c} 66 \\ Dy \\ 4f^{10}6s^2 \end{array} $	$67 \\ Ho \\ 4f^{11}6s^2$	$68 \\ Er \\ 4f^{12}6s^2$	69 Tm 4f ¹³ 6s ²	$70 \\ Yb \\ 4f^{14}6s^2$				
[Rn]	Actin	ide se	ries	89 Ac 6d ¹ 7s ²	90 Th $6d^27s^2$	91 Pa $5f^{2}6d^{1}$ $7s^{2}$	92 U $5f^{3}6d^{1}$ $7s^{2}$	93 Np $5f^{4}6d^{1}$ $7s^{2}$	94 Pu 5f ⁶ 7s ²	95 Am 5f ⁷ 7s ²	$\frac{65^{\circ}}{96}$ $\frac{65^{\circ}}{5f^{\circ}6d^{1}}$ $\frac{75^{\circ}}{7s^{\circ}}$	97 Bk 5f ⁹ 7s ²	98 Cf 5f ¹⁰ 7s ²	99 Es 5f ¹¹ 7s ²	$100 \ Fm \ 5f^{12}7s^2$	101 Md 5f ¹³ 7s ²	102 No 5f ¹⁴ 7s ²				
				N	/letals		7	lloids		Noni	netals		1							Electro Struct of Ato	ure

- Electronic configuration
- Condensed electronic configuration
- Orbital Diagram





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GROUP	ELEMENT	ATOMIC NUMBER	VALENCE-SHELL ELECTRON CONFIGURATION
1A	Li (lithium)	3	$2s^1$
2A			
7A			
8A			
OA			

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	•	

GROUP	ELEMENT	ATOMIC NUMBER	VALENCE-SHELL ELECTRON CONFIGURATION
1A	Li (lithium)	3	$2s^1$
	Na (sodium)	11	$3s^{1}$
2A			
7A			
8A			

GROUP	ELEMENT	ATOMIC NUMBER	VALENCE-SHELL ELECTRON CONFIGURATION
1A	Li (lithium)	3	$2s^1$
	Na (sodium)	11	$3s^1$
	K (potassium)	19	$4s^1$
2A			
7A			
8A			

GROUP	ELEMENT	ATOMIC NUMBER	VALENCE-SHELL ELECTRON CONFIGURATION
1A	Li (lithium)	3	$2s^1$
	Na (sodium)	11	$3s^1$
	K (potassium)	19	$4s^1$
	Rb (rubidium)	37	$5s^1$
2A 7A 8A			

GROUP	ELEMENT	ATOMIC NUMBER	VALENCE-SHELL ELECTRON CONFIGURATION
1A	Li (lithium)	3	$2s^1$
	Na (sodium)	11	$3s^1$
	K (potassium)	19	$4s^1$
	Rb (rubidium)	37	$5s^1$
	Cs (cesium)	55	$6s^1$
2A 7A			
8A			

GROUP	ELEMENT	ATOMIC NUMBER	VALENCE-SHELL ELECTRON CONFIGURATION
1A	Li (lithium)	3	$2s^1$
	Na (sodium)	11	$3s^1$
	K (potassium)	19	$4s^1$
	Rb (rubidium)	37	$5s^1$
	Cs (cesium)	55	$6s^1$
2A	Be (beryllium)	4	$2s^2$
7A			
8A			

GROUP	ELEMENT	ATOMIC NUMBER	VALENCE-SHELL ELECTRON CONFIGURATION
1A	Li (lithium)	3	$2s^1$
	Na (sodium)	11	$3s^1$
	K (potassium)	19	$4s^1$
	Rb (rubidium)	37	$5s^{1}$
	Cs (cesium)	55	$6s^1$
2A	Be (beryllium)	4	$2s^2$
	Mg (magnesium)	12	$3s^{2}$
7A 8A			

GROUP	ELEMENT	ATOMIC NUMBER	VALENCE-SHELL ELECTRON CONFIGURATION
1A	Li (lithium)	3	$2s^1$
	Na (sodium)	11	$3s^1$
	K (potassium)	19	$4s^1$
	Rb (rubidium)	37	$5s^{1}$
	Cs (cesium)	55	$6s^1$
2A	Be (beryllium)	4	$2s^2$
	Mg (magnesium)	12	$3s^2$
	Ca (calcium)	20	$4s^2$
7A 8A			

conic ture

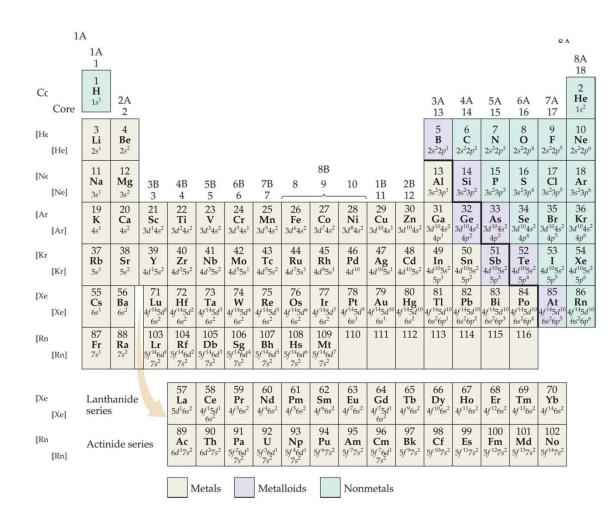
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GROUP	ELEMENT	ATOMIC NUMBER	VALENCE-SHELL ELECTRON CONFIGURATION
1A	Li (lithium)	3	$2s^1$
	Na (sodium)	11	$3s^1$
	K (potassium)	19	$4s^1$
	Rb (rubidium)	37	$5s^1$
	Cs (cesium)	55	$6s^1$
2A	Be (beryllium)	4	$2s^2$
	Mg (magnesium)	12	$3s^2$
	Ca (calcium)	20	$4s^2$
	Sr (strontium)	38	$5s^2$
	Ba (barium)	56	$6s^2$
4 A	hav	ve 4 electrons	
5A		5 electrons	
6A		6 electrons	
7 A		7 electrons	
8A		8 electrons	
	in the	eir outer most sh	nell

GROUP	ELEMENT	ATOMIC NUMBER	VALENCE-SHELL ELECTRON CONFIGURATION
1A	Li (lithium)	3	$2s^1$
	Na (sodium)	11	$3s^1$
	K (potassium)	19	$4s^1$
	Rb (rubidium)	37	$5s^{1}$
	Cs (cesium)	55	$6s^1$
2A	Be (beryllium)	4	$2s^2$
	Mg (magnesium)	12	$3s^2$
	Ca (calcium)	20	$4s^2$
	Sr (strontium)	38	$5s^{2}$
	Ba (barium)	56	$6s^2$
7A	F (fluorine)	9	$2s^2 2p^5$
	Cl (chlorine)	17	$3s^2 3p^5$
	Br (bromine)	35	$4s^2 4p^5$
	I (iodine)	53	$5s^2 5p^5$
8A	He (helium)	2	$1s^2$
	Ne (neon)	10	$2s^2 2p^6$
	Ar (argon)	18	$3s^2 3p^6$
	Kr (krypton)	36	$4s^2 4p^6$
	Xe (xenon)	54	$5s^2 5p^6$

conic ture oms

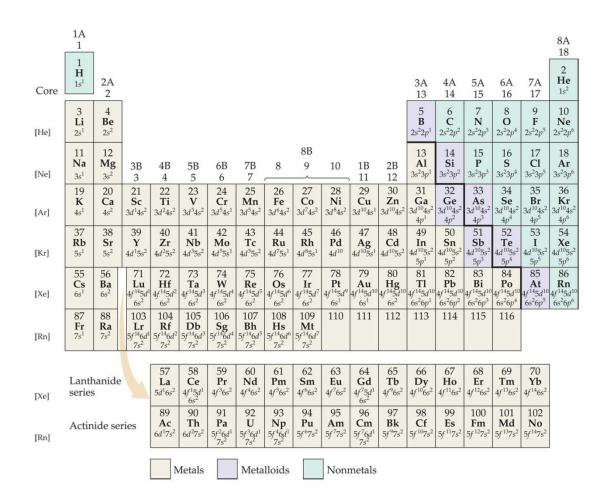
Some Anomalies



Some irregularities occur when there are enough electrons to halffill *s* and *d* orbitals on a given row.



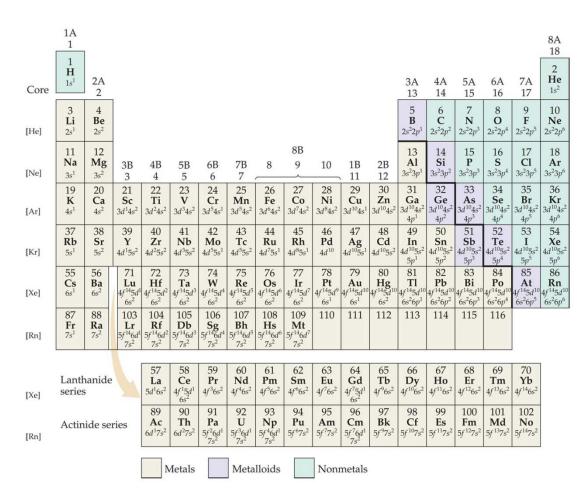
Some Anomalies



For instance, the electron configuration for copper is [Ar] $4s^1 3d^{10}$ rather than the expected [Ar] $4s^2 3d^9$.



Some Anomalies



- This occurs because the 4s and 3d orbitals are very close in energy.
- Mo, Pd and Ag also show these anomalies.





- Ag [Kr] 4d¹⁰ 5s¹
- Pd [Kr] 4d¹⁰
- Mo [Kr] 4d⁵ 5s¹

Actual

Not [Kr] 4d⁴ 5s² [Kr] 4d⁸ 5s² [Kr] 4d⁹ 5s²

• There are some anomalies in the f block elements too.

• These anomalies though interesting do not have great chemical significance.

