

Chapter Seven

The Electronic Structure of Atoms

<http://clipart-library.com>

Wave Theory

Wave

- Repeating disturbance spreading out from a defined origin
- Characterized by wavelength, frequency and amplitude

Wavelength (λ)

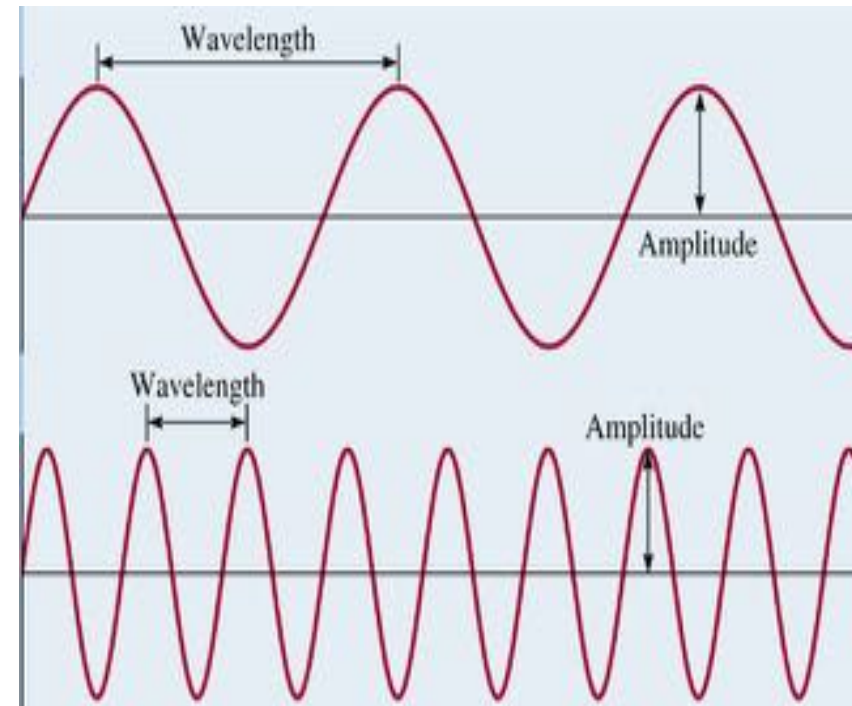
- Distance between identical pts
- Units some form of meters

Frequency (ν)

- Number of waves that pass through a point in 1 second
- Units of cycles/sec or Hz

Amplitude

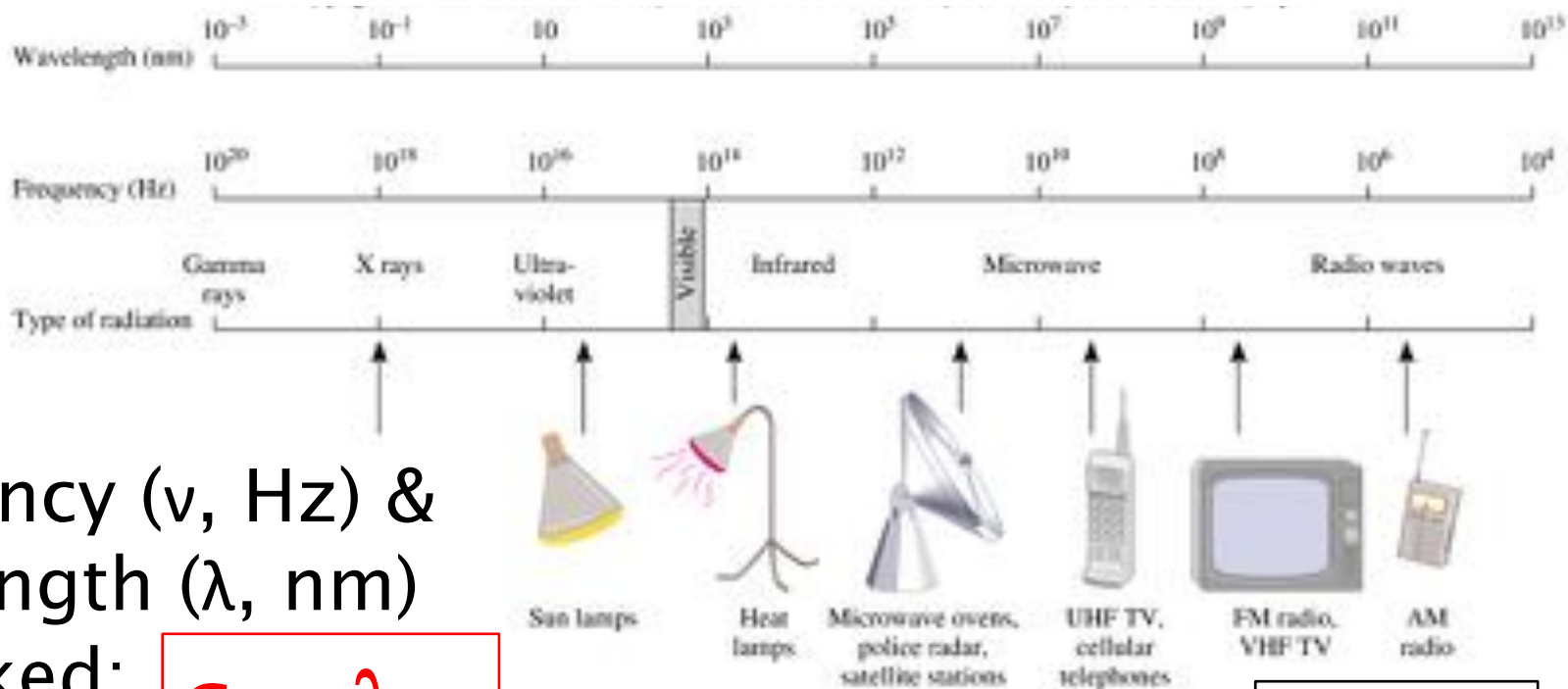
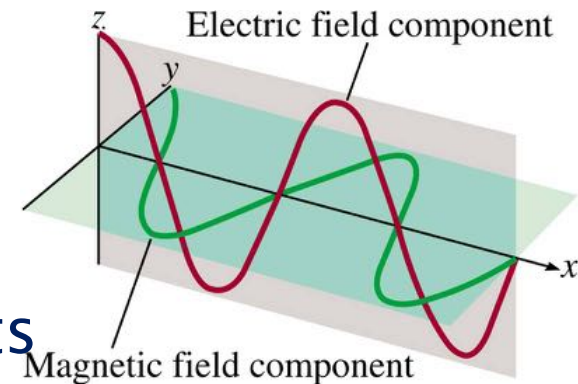
- Height of wave from center point
- Intensity of wave



Electromagnetic Radiation

Electromagnetic Radiation

- Emission/transmission of energy
- In form of waves
- Has electrical & magnetic components
- Travels at the speed of light ($c = 3.00 \times 10^8 \text{ m/s}$)



Frequency (ν , Hz) & wavelength (λ , nm) are linked:

$$c = \lambda \nu$$

$$\text{Hz} = \text{s}^{-1}$$

Using the relationship $c = \lambda\nu$:

What is the wavelength of an FM-radiowave with a 94.9 MHz frequency?

A: 3.16 m

Max Planck's Quantum Theory

Studied energy emitted by objects

- Amount of energy emitted was directly related to wavelength at which energy was emitted

Theory: Energy must be in discrete amounts.

- Amounts were defined by λ (& ν – they are related!)

$$E = h\nu = hc/\lambda$$

- Can have multiples of these discrete amounts

$$E = h\nu, E = 2h\nu, E = 3h\nu \dots$$

- **h = Planck's constant = $6.626 \times 10^{-34} \text{ J s}$**

Called the smallest amount of energy a **Quantum**.

Didn't know why, but math worked over entire spectrum

Einstein and the Photoelectric Effect

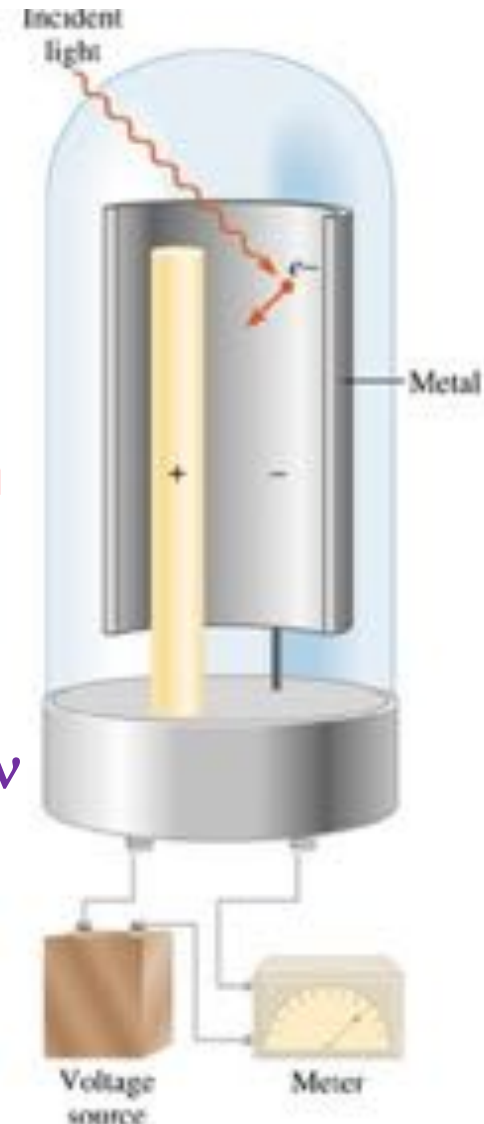
Experiment to prove why $E = h\nu$

- Full spectrum of light hits metal surface
- Energy transferred to electrons in metal
- Electrons break free and escape to anode
- Flow of electrons recorded with voltmeter
- **Light energy must be at or above a certain frequency to dislodge electrons**

Conclusions:

- **Light energy has wave properties: $E = h\nu$**
&
- **Light energy has particle properties**

Particles of light were later called
“photons”



Using $E = h\nu$ ($h = 6.626 \times 10^{-34} \text{ Js}$)

What is the energy of a radiowave with a frequency of 94.9 MHz? A: $6.29 \times 10^{-26} \text{ J}$

What wavelength has an energy of $1.00 \times 10^{-20} \text{ J}$?

A: $1.99 \times 10^{-5} \text{ m}$
Or $19.9 \text{ } \mu\text{m}$

Using $E = h\nu$ ($h = 6.626 \times 10^{-34} \text{ Js}$)

What is the energy per photon and per mole of photons of violet light, with a wavelength of 415 nm?

A: $4.79 \times 10^{-19} \text{ J/photon}$
A: $2.88 \times 10^5 \text{ J/mol}$

Continuous vs. Line Spectra

Continuous spectrum:

- Occurs when all visible light is present: white light



Line Spectrum

- Occurs when light is produced through an element
- Pattern of lines is characteristic of the element
- Can be used for identification of elements



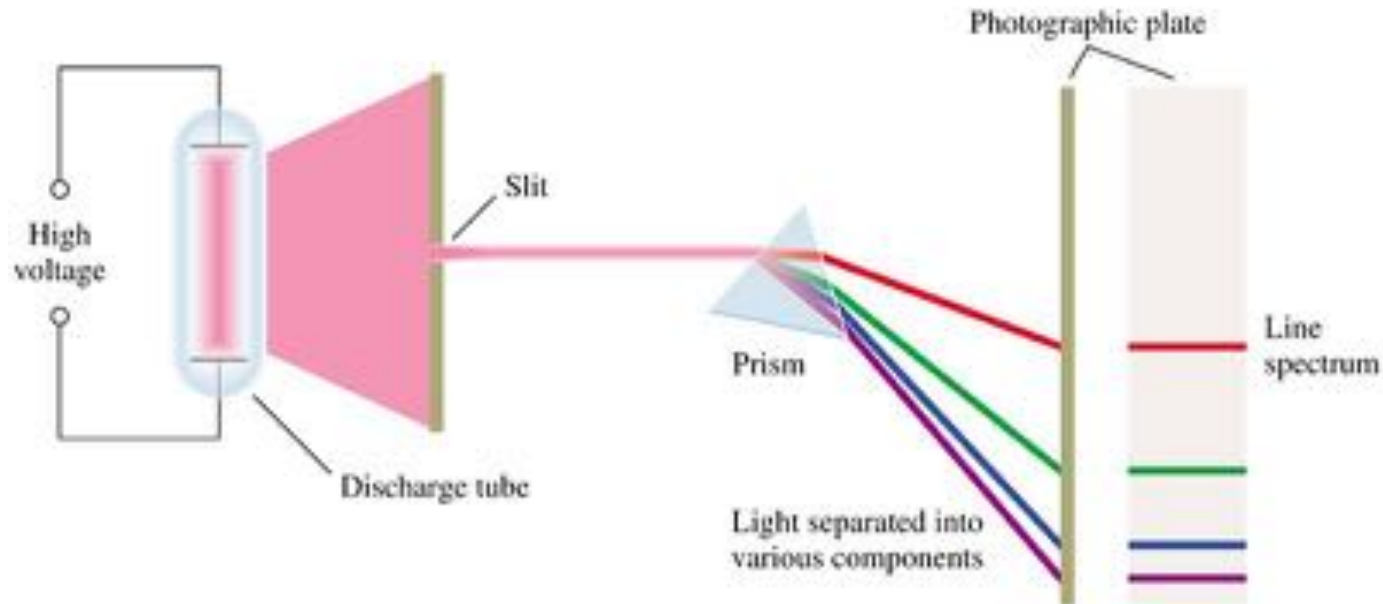
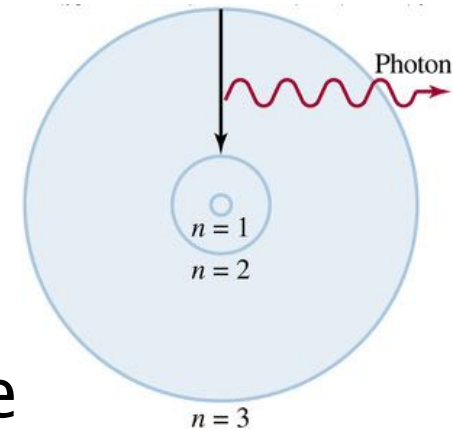
Bohr's Theory of the Hydrogen Atom

10

Emission Spectra: Pattern of radiation that is emitted when photons are removed from a substance.

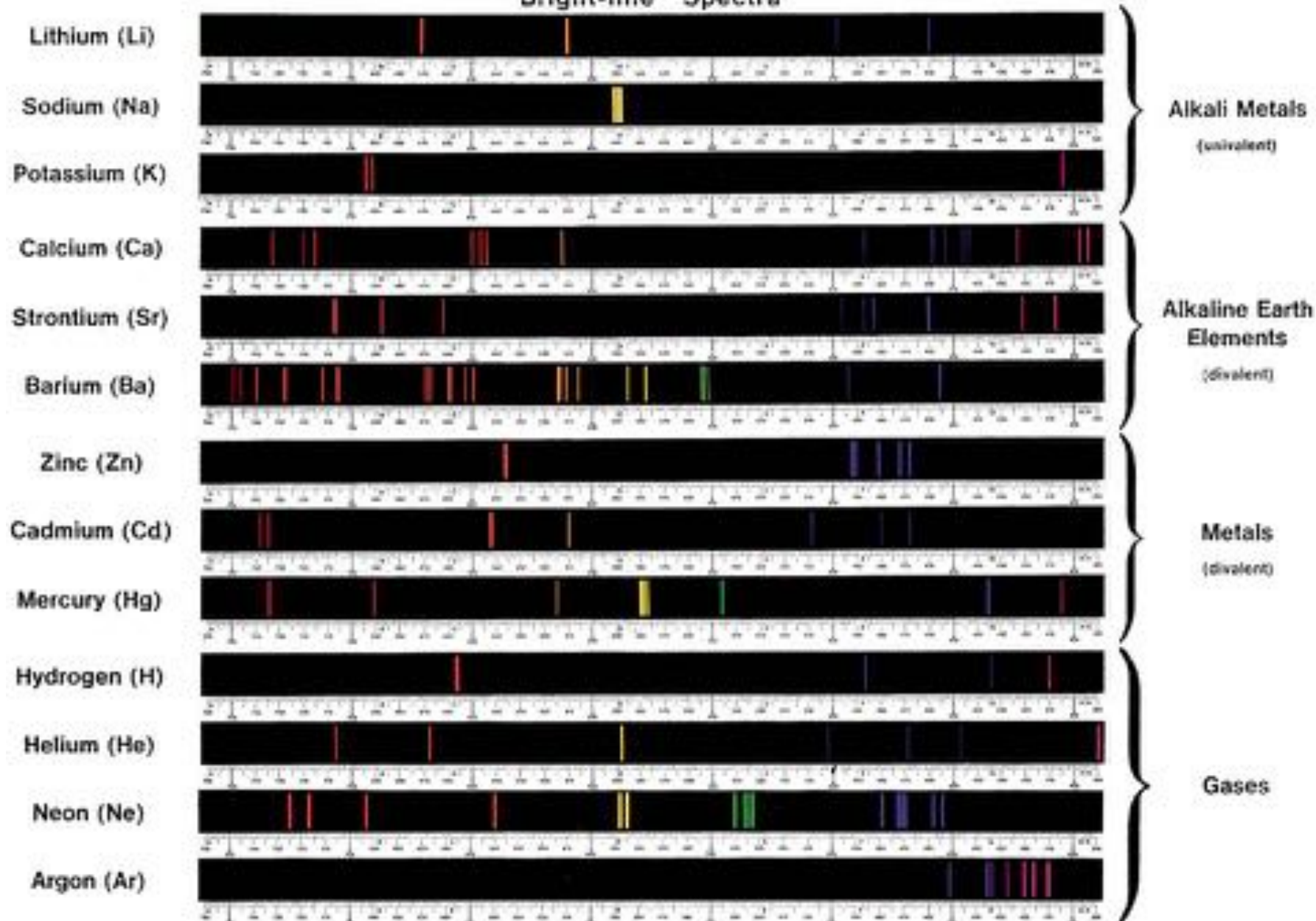
Procedure

- Add energy to a substance
- Photons are emitted as a beam of light
- Separate wavelengths through a prism
- Record pattern on a photographic plate



Elemental Line Spectra

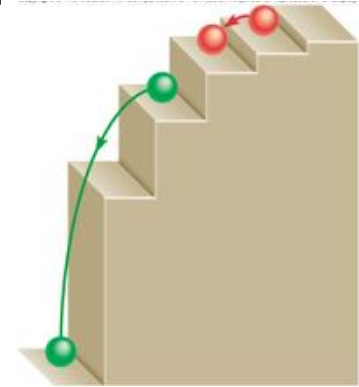
Bright-line Spectra



Bohr's Hydrogen Atom

Niels Bohr (1913): Electron energy (E_n) was quantized

- Only certain specified values allowed
- Stable levels called energy levels
- Photon absorbed/released when electron moves from 1 level to another



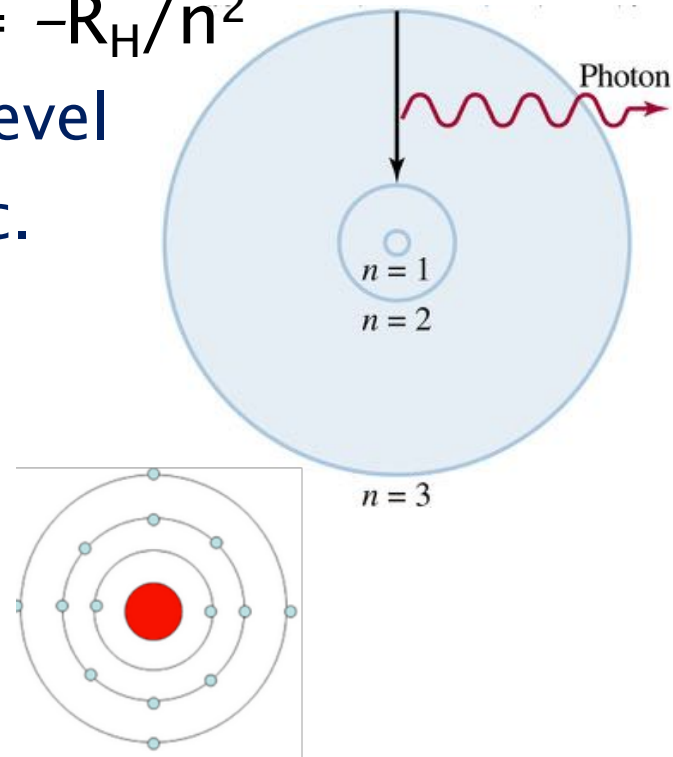
The energy of each stable orbit: $E_n = -R_H/n^2$

- n is the quantum number of the level
- n is always an integer, 1,2,3,...etc.

Proportionality constant R_H

- Rydberg constant
- **$R_H = 2.18 \times 10^{-18} \text{ J}$**

Leads to orbit description of atoms



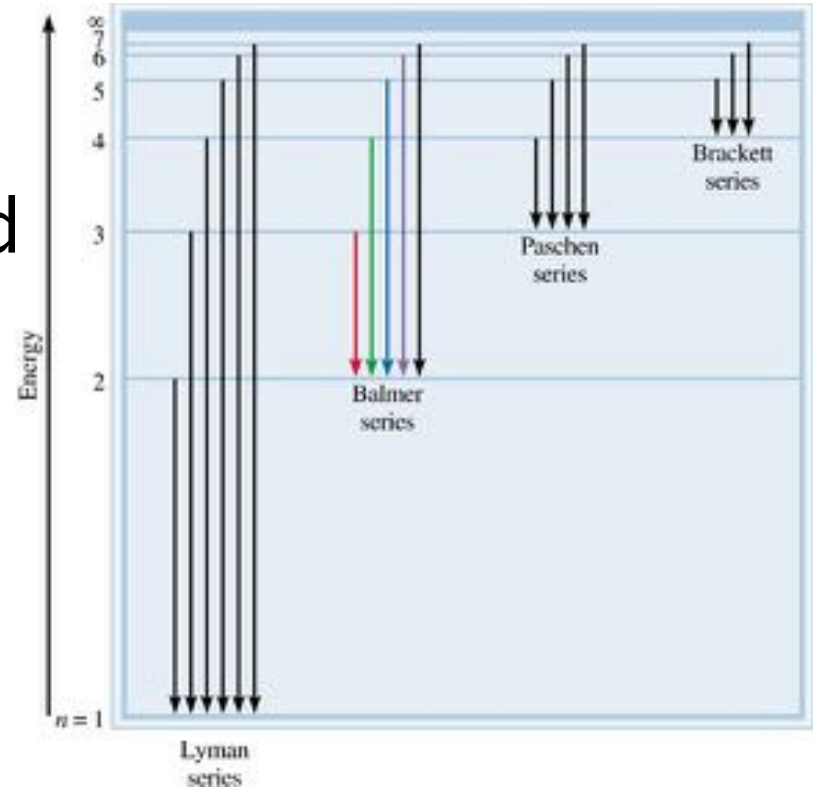
Energy Level Calculations

All calculations done by comparing energy levels

- Electron moves between levels
- $E = -R_H (1/n_f^2 - 1/n_i^2)$

Energy emitted or absorbed

- High to low level:
 - energy released (-)
- Low to high level:
 - energy absorbed (+)



Ground state: An e^- 's lowest possible energy level

Excited state: All other levels

Calculate the wavelength of the electron shift from¹⁴
 $n = 4$ to $n = 2$. Is light emitted or absorbed?

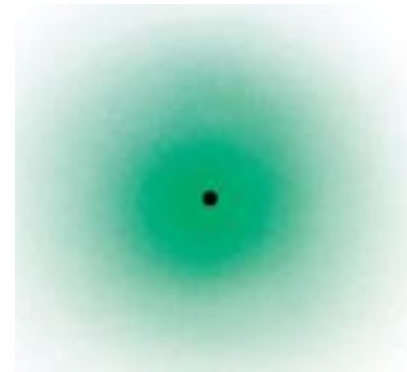
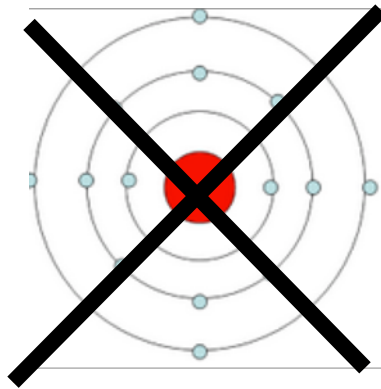
$$E = -R_H \left(\frac{1}{n_f^2} - \frac{1}{n_i^2} \right)$$

$$R_H = 2.18 \times 10^{-18} \text{ J}$$

A: $\lambda = 486 \text{ nm}$ Visible blue green light is emitted (neg E value)

Modern View of the Atom: Quantum Mechanics – a very brief intro

- (Nucleus in center, protons & neutrons in nucleus)
- Electrons outside nucleus
 - located in “cloud” surrounding the nucleus
 - likely location based on probability functions
 - quantum numbers used to describe probable location



$$i\hbar \frac{\partial \Psi}{\partial t} = -\frac{\hbar^2}{2m} \nabla^2 \Psi + [V_1(x) + iV_2(x)] \Psi$$

Quantum Numbers and Atomic Orbitals

Atomic orbital

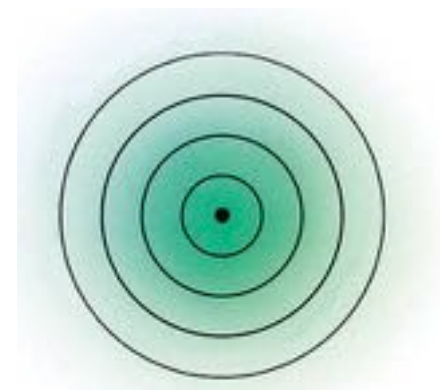
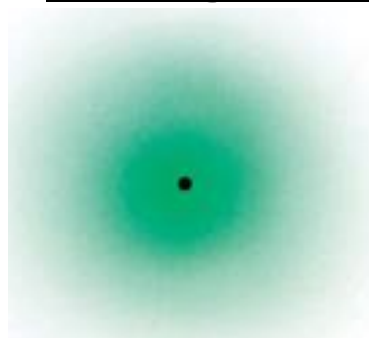
- A region in space with a high probability of finding an electron.
- Identified by 4 quantum numbers.

4 Quantum Numbers (think of it as a dorm address)

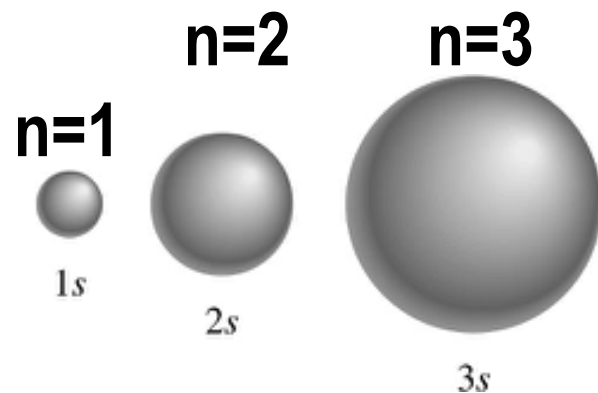
- | | |
|--|----------|
| 1. Principal quantum number (n): | Building |
| 2. Angular momentum quantum number (l) | Floor |
| 3. Magnetic quantum number (m_l) | Room # |
| 4. Electron spin quantum number (m_s) | Bed |

The Principal Quantum Number (n)

- Restricted to the positive integers: 1, 2, 3, 4, 5, 6, 7
- The shell or energy level of the orbital



- Indicates the size of the orbital
 - max distance e^- can travel from nucleus
- Integers correspond to row numbers in Periodic Table
 - row an element is in tells you the highest energy level in the ground state



The Angular Momentum Quantum Number (ℓ) ¹⁸

- Indicates orbital shape

- Designation: s, p, d or f

level	0	1	2	3
Name	s	p	d	f

- Designates the **subshell**

- Values range from 0 to $n-1$
- 0-6 theoretically, but realistically 0-3
- Give rise to “Blocks” in periodic table

Energy Level (n)	Math	Allowed ℓ values	Orbitals
1	$1-1 = 0$	0	s only
2	$2-1 = 1$	0, 1	s & p
3	$3-1 = 2$	0, 1, 2	s, p, & d

The Magnetic Quantum Number (m_ℓ):

Determines the orientation in space of the orbitals

- “orientation” refers to proximity to axes (x, y, z)
- Integers from $-\ell$ to $+\ell$

Determines the number of orbitals in a subshell

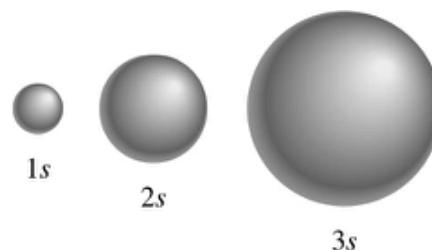
- The number of possible values for $m_\ell = 2\ell + 1$

Orbital	ℓ value	Allowed m_ℓ values	Number of Orbitals per Energy Level
s	0	0	1
p	1	-1, 0, 1	3
d	2	-2, -1, 0, 1, 2	5
f	3	-3, -2, -1, 0, 1, 2, 3	7

Orbital Shapes = ℓ quantum number

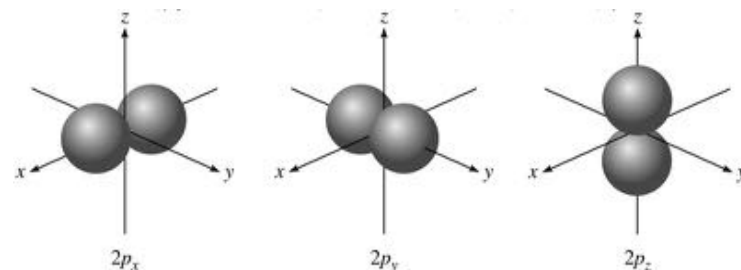
$\ell = 0$: s orbitals

- Spherical
- **One** per energy level



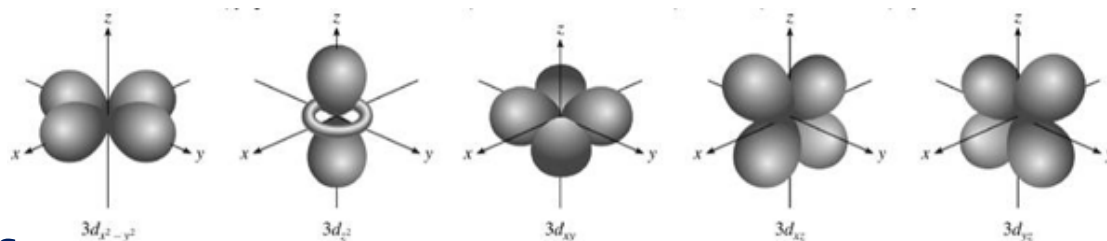
$\ell = 1$: p orbitals

- 2 teardrops joined at center
- **Three** per energy level



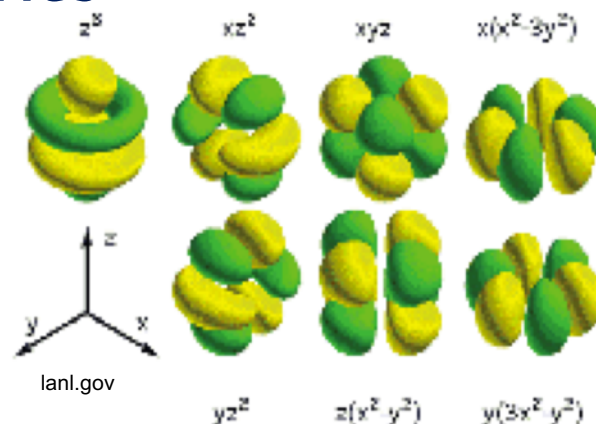
$\ell = 2$: d orbitals

- Most are like two p orbitals along different axes
- **Five** per energy level



$\ell = 3$: f orbitals.

- Complicated shapes
- **Seven** per energy level



Orbitals with same n & ℓ values are “degenerate”

degenerate = same energy

(Note: In some cases there are slight energy differences)

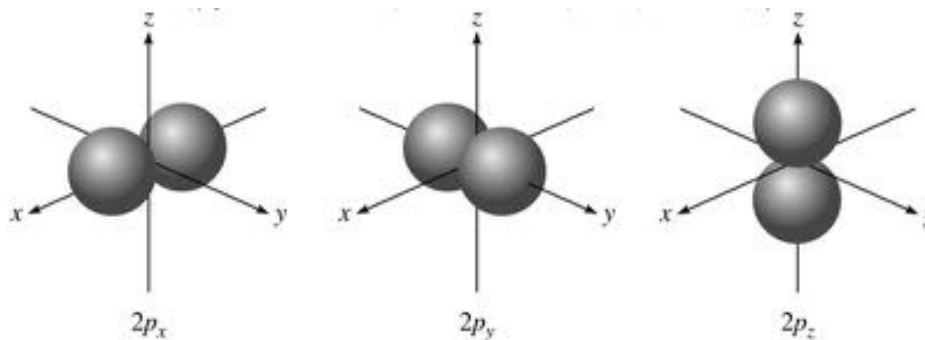
Possible quantum numbers for an electron in a 3p orbital:

$$n = 3$$

ℓ can be 0 to $3-1$ (0, 1, 2) BUT if it is a p orbital $\ell = 1$

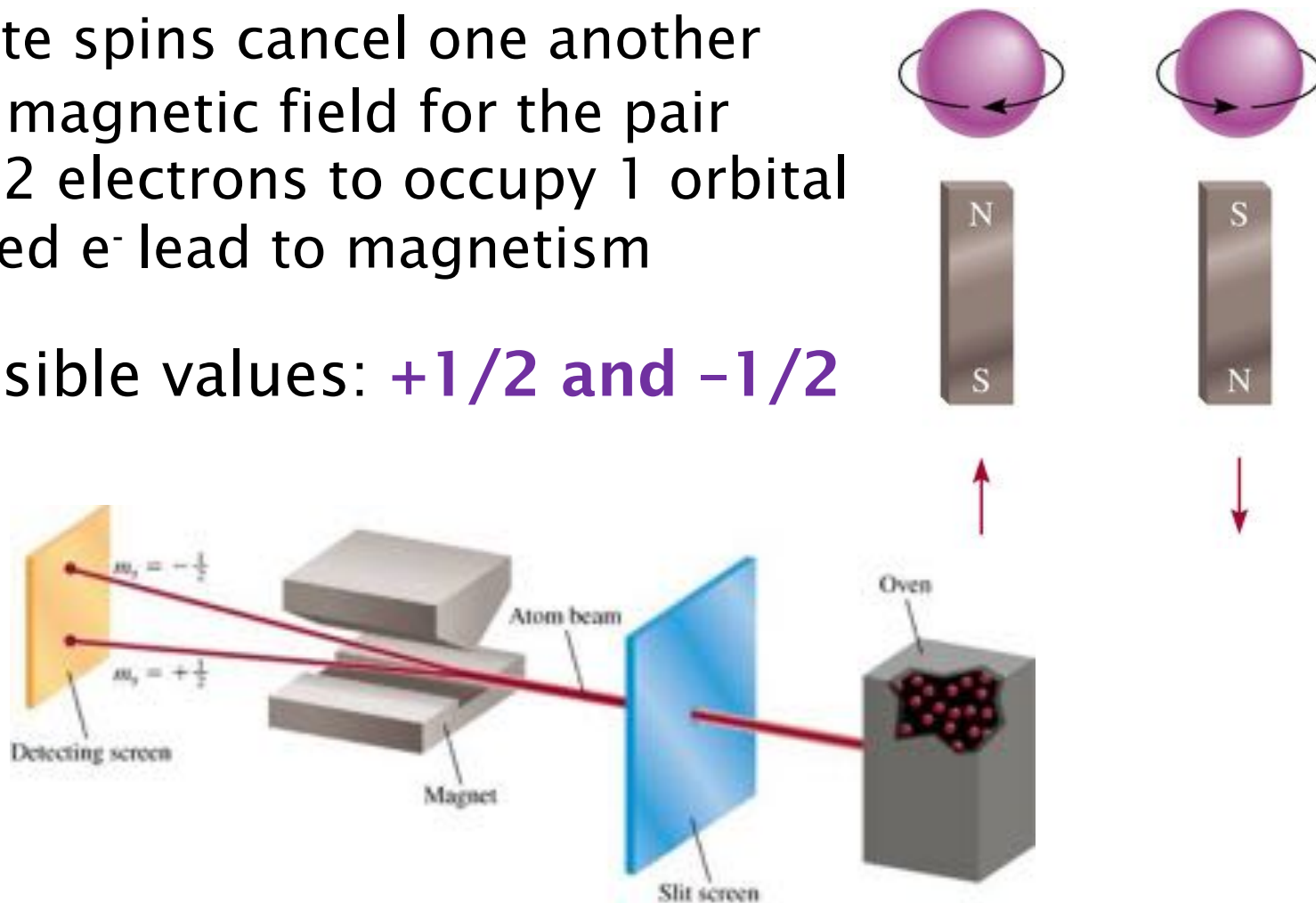
m_ℓ can be $+\ell$ to $-\ell = -1, 0, +1$

Since the 3p orbitals are degenerate, any of the three m_ℓ values could be correct



Electron Spin Quantum Number (m_s)

- A magnetic field is induced by the moving electric charge of an electron as it spins
 - Opposite spins cancel one another
 - No net magnetic field for the pair
 - Allows 2 electrons to occupy 1 orbital
 - Unpaired e^- lead to magnetism
- Two possible values: **$+1/2$ and $-1/2$**



Quantum Numbers Summary

TABLE 7.2

Relation Between Quantum Numbers and Atomic Orbitals

n	ℓ	m_ℓ	Number of Orbitals	Atomic Orbital Designations
1	0	0	1	$1s$
2	0	0	1	$2s$
	1	-1, 0, 1	3	$2p_x, 2p_y, 2p_z$
3	0	0	1	$3s$
	1	-1, 0, 1	3	$3p_x, 3p_y, 3p_z$
	2	-2, -1, 0, 1, 2	5	$3d_{xy}, 3d_{yz}, 3d_{xz},$ $3d_{x^2-y^2}, 3d_{z^2}$
⋮	⋮	⋮	⋮	⋮
⋮	⋮	⋮	⋮	⋮
⋮	⋮	⋮	⋮	⋮

A possible set of quantum numbers for the last electron added to complete an atom of selenium would be:

n :

l :

m_l :

m_s :

Electron Configuration:

Finding a home for each electron

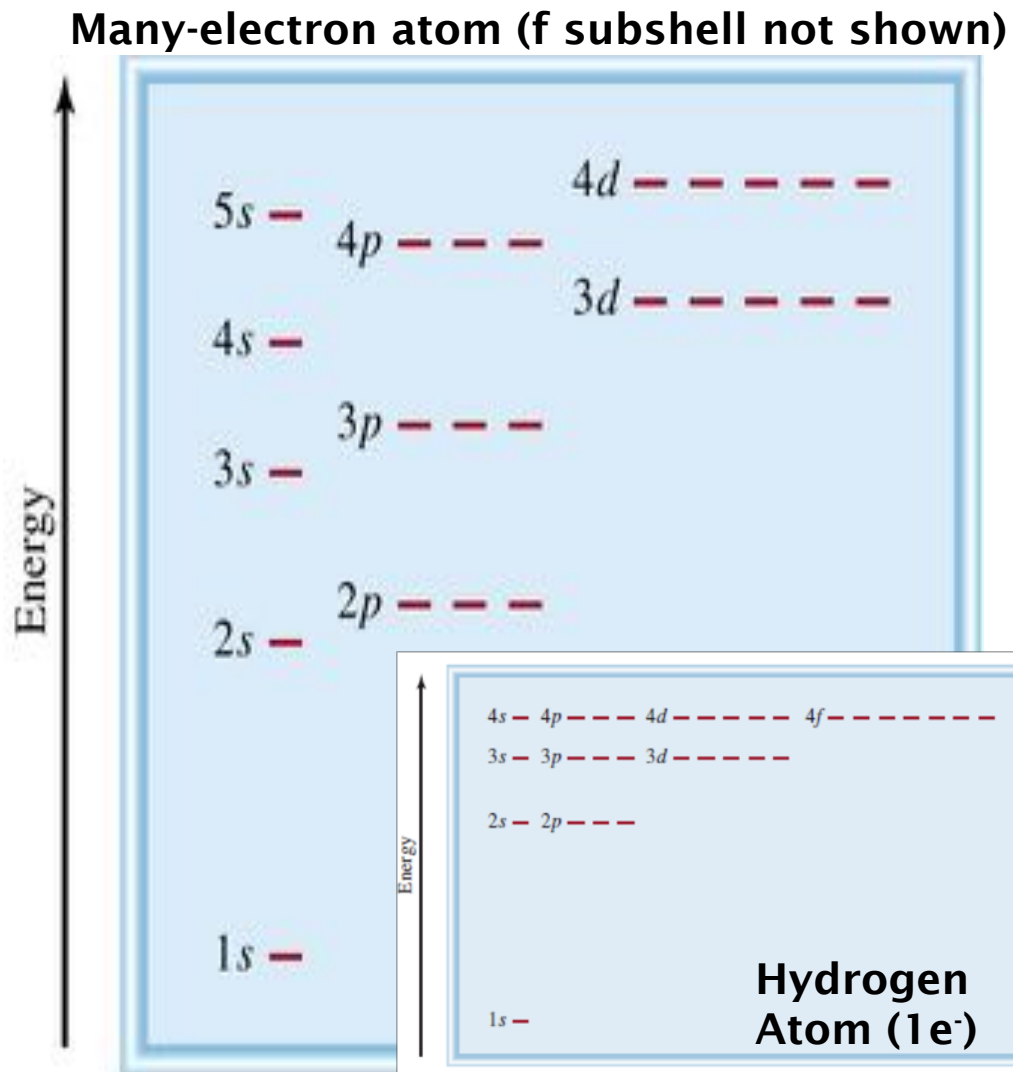
The energy of an electron is defined by both n & ℓ

- Principle shells (size)

- $n = 1, 2, 3, 4$ or 5

- Subshells (shape)

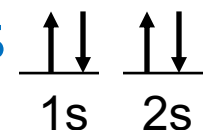
- $\ell = 0, 1, 2$, or 3
- n determines number of subshells
- s, p, d, f orbitals
- Shielding impacts relative energies



Rules & Principles Governing e⁻ Configurations

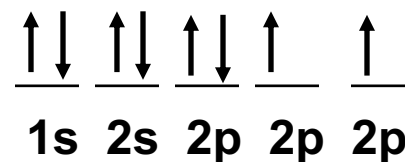
Pauli Exclusion Principle:

- No 2 e⁻ in an atom can have the same set of 4 quantum #s
 - If in the same orbital, e⁻ must have opposite spins



Hund's rule:

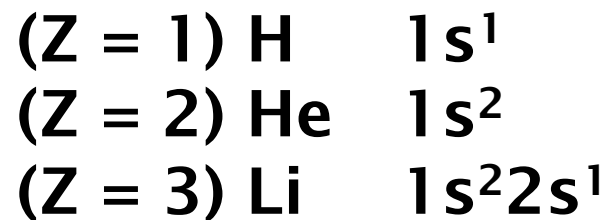
- Electrons in the same subshell occupy degenerate orbitals singly, before pairing
 - Degenerate = same energy



Ex: Oxygen, O Z = 8

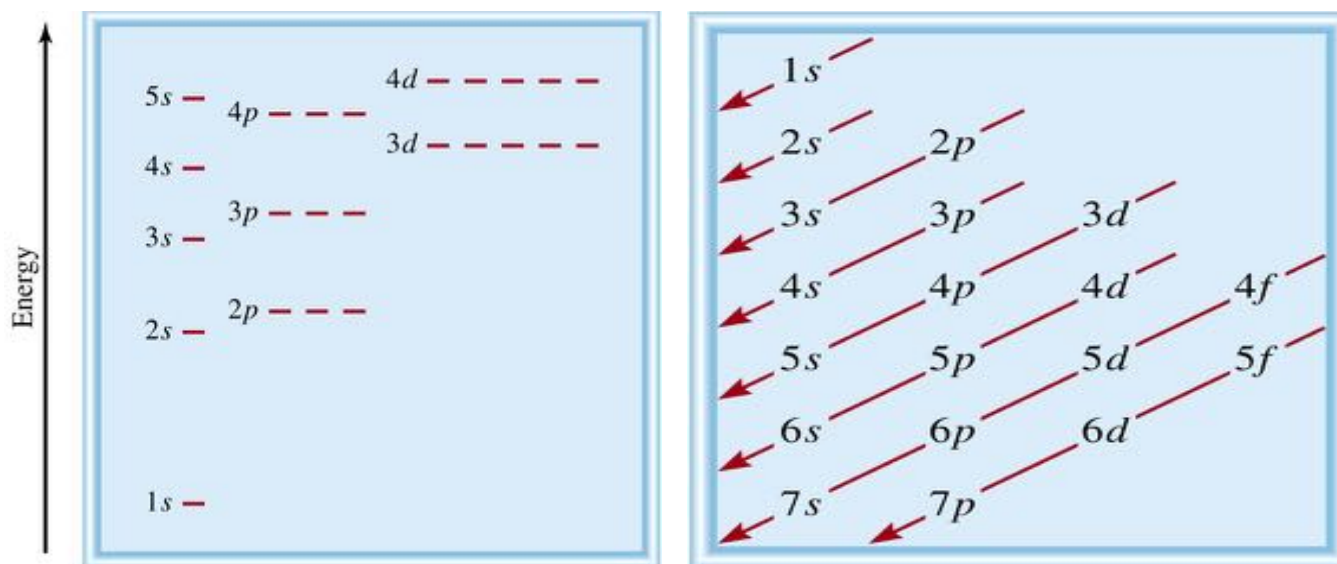
The Aufbau Principle:

- In general, each successive electron added to an atom occupies the lowest energy orbital available
 - There are some exceptions



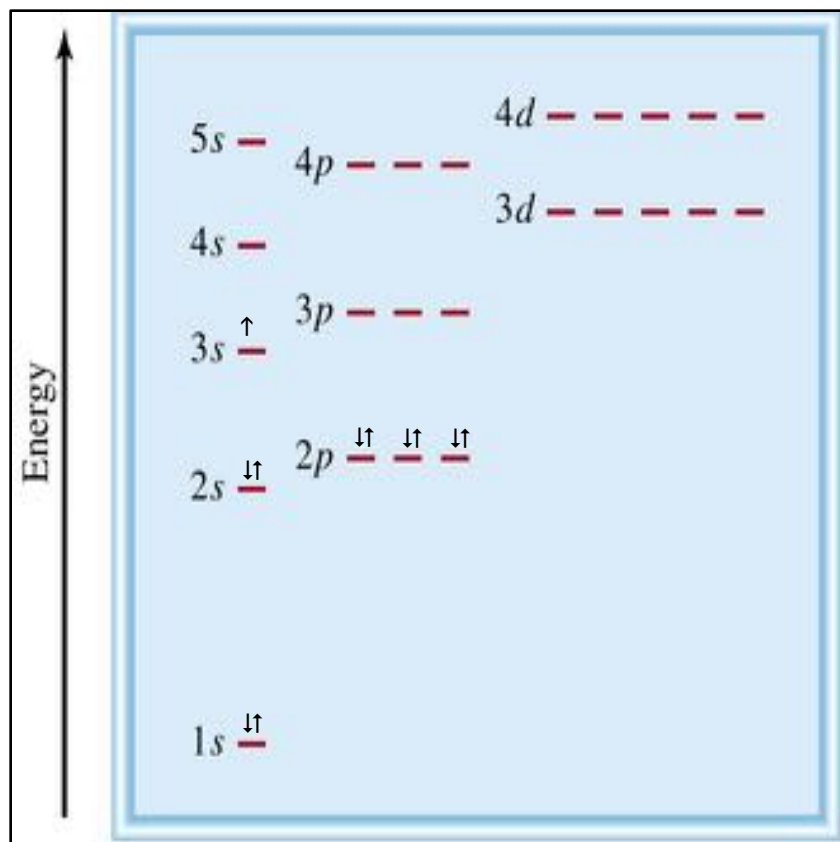
Orbital Filling in Multi-electron Atoms

- Fill low to high energy
 - 2 electrons per orbital
 - Shielding impacts the energy of orbitals
 - Use chart to account for overlap of n values
- $1s < 2s < 2p < 3s < 3p < 4s < 3d < 4p < 5s < 4d < 5p < 6s$
- Format: spdf or orbital notation
 - Ends when a home is found for each electron



Electron Configuration con't

- Defines the orbital (“home”) for each electron
- # electrons = atomic number (Z) of atom (if neutral)
 - Max 2 electrons per orbital



Orbital Diagrams

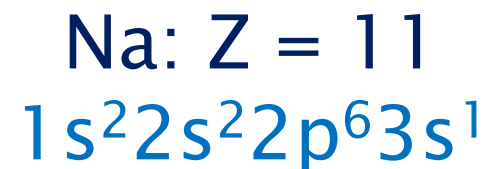
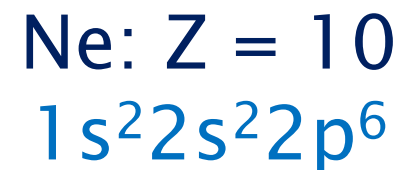
- Energy increases from bottom to top
 - Higher energy levels at top
- Boxes or lines represent orbitals
 - # lines at one level = # degenerate orbitals
- Arrows ($\uparrow\downarrow$) represent e^-
- 2 e^- allowed per orbital
 - one arrow up & one arrow down to show the different spins

Formats for Electron Configurations

spdf Notation

- Front number = energy level
- Letter = type of orbital (s, p, d, or f)
 - Degenerate orbitals are combined together
- Superscript = # electrons in that type of orbital
 - Degenerate orbitals are combined, so the superscript can be more than 2 if it is a p, d, or f orbital (p max 6, d max 10, f max 14)

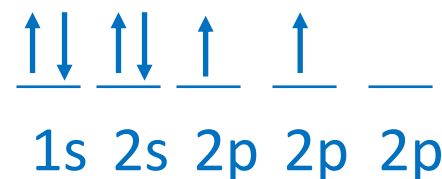
Examples:



Orbital Notation

- Number = energy level
- Letter = type of orbital
 - Degenerate orbitals are NOT combined
- Arrows = electrons
 - Put one e^- in each degenerate orbital before pairing
 - If 2 e^- in one orbital, one arrow must be up, the other down

Example:



Writing Electron Configurations

Sulfur:

Vanadium:

Writing Electron Configurations for Ions

Remove Electrons from Highest Energy Level First

Magnesium ion (Mg^{2+}):

Fluorine ion (F^-):

Manganese (II) ion (Mn^{2+}):

Noble Gas Configuration

- Abbreviation of Electron Configuration (ex: Na: [Ne]3s¹)
- Noble gas symbol replaces the portion of the e⁻ config. that is identical to the e⁻ config. of the noble gas.
- Always use the largest noble gas that is smaller than the element
- Can use for either spdf or orbital notation
- ex: Arsenic
regular configuration: 1s²2s²2p⁶3s²3p⁶4s²3d¹⁰4p³
- ex: Strontium
regular configuration: 1s²2s²2p⁶3s²3p⁶4s²3d¹⁰4p⁶5s²

Exceptions To The Aufbau Principle

Half filled & filled subshells provide additional stability

Cr and Cu ½ fill/fill their 3d shell before the 4s shell.

Elements in same columns as Cr & Cu behave in same way.

Similar behavior seen in p block.

Cr & Cu are the only exceptions you need to know.

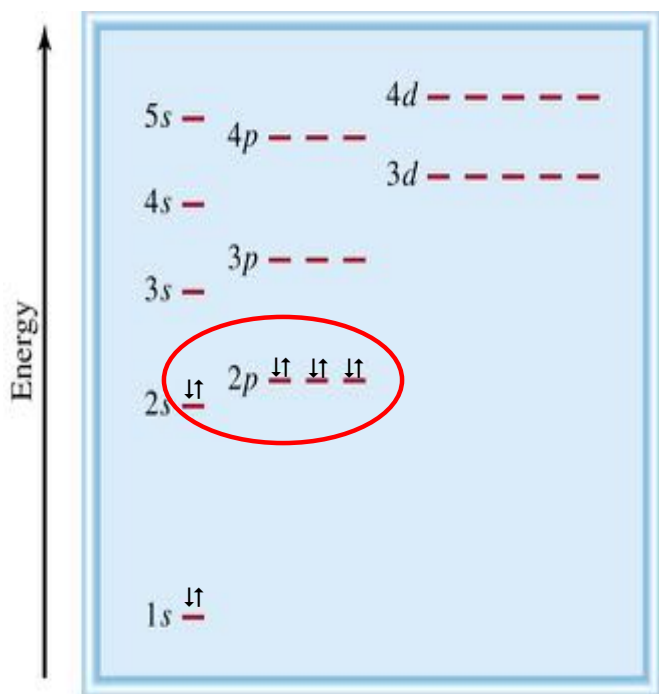
		3d	4s	
Sc	[Ar]			[Ar]3d ¹ 4s ²
Ti	[Ar]			[Ar]3d ² 4s ²
V	[Ar]			[Ar]3d ³ 4s ²
Cr	[Ar]			[Ar]3d ⁵ 4s ¹
Mn	[Ar]			[Ar]3d ⁵ 4s ²
Fe	[Ar]			[Ar]3d ⁶ 4s ²
Co	[Ar]			[Ar]3d ⁷ 4s ²
Ni	[Ar]			[Ar]3d ⁸ 4s ²
Cu	[Ar]			[Ar]3d ¹⁰ 4s ¹
Zn	[Ar]			[Ar]3d ¹⁰ 4s ²

Cr: 3d ½ full
& 4s ½ full

Cu: 3d full
& 4s ½ full

Magnetism in Multi-electron Atoms

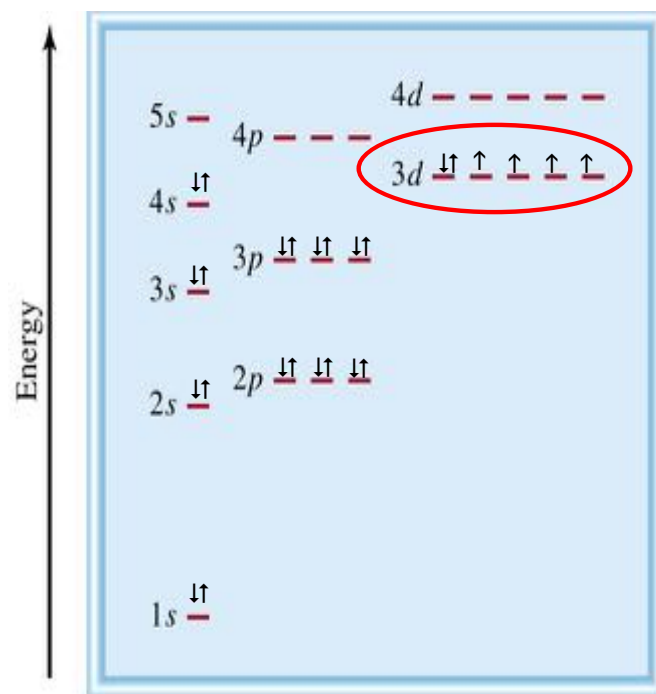
- $+1/2$ & $-1/2$ spins will cancel if electrons paired
- No magnetic properties without spin present
 - # unpaired electrons proportional to magnetic properties



Diamagnetic

All electrons paired

Ne: $1s^2 2s^2 2p^6$



Paramagnetic

At least 1 unpaired electron

Fe: $1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^6$

Quantum Numbers, Electron Configuration, & the Periodic Table

- **Principle quantum number, n**
 - Row number of periodic table, values of 1-7
- **Angular momentum quantum number, ℓ**
 - Specific area of periodic table, spdf “blocks”
- **Can follow the periodic table to fill e^- configuration**
- **Can use location on Periodic Table to determine where e^- configuration will end**

1s							1s
2s						2p	
3s						3p	
4s	3d					4p	
5s	4d					5p	
6s	5d					6p	
7s	6d					7p	
				4f			
				5f			

Electronic Configurations and the Periodic Table

Add 1 electron for each square (ie element) in the Periodic Table

Electrons in the outermost energy level are the valence electrons.

1 1A	2 2A											13 3A	14 4A	15 5A	16 6A	17 7A	18 8A
1 H 1s ¹												3 B 2s ² 2p ¹	6 C 2s ² 2p ²	7 N 2s ² 2p ³	8 O 2s ² 2p ⁴	9 F 2s ² 2p ⁵	10 Ne 2s ² 2p ⁶
2 Li 2s ¹	4 Be 2s ²											13 Al 3s ² 3p ¹	14 Si 3s ² 3p ²	15 P 3s ² 3p ³	16 S 3s ² 3p ⁴	17 Cl 3s ² 3p ⁵	18 Ar 3s ² 3p ⁶
3 Na 3s ¹	12 Mg 3s ²	3 3B	4 4B	5 5B	6 6B	7 7B	8 8B	9 9B	10 10B	11 11B	12 12B	13 Al 3s ² 3p ¹	14 Si 3s ² 3p ²	15 P 3s ² 3p ³	16 S 3s ² 3p ⁴	17 Cl 3s ² 3p ⁵	18 Ar 3s ² 3p ⁶
4 K 4s ¹	20 Ca 4s ²	21 Sc 4s ² 3d ¹	22 Ti 4s ² 3d ²	23 V 4s ² 3d ³	24 Cr 4s ¹ 3d ⁵	25 Mn 4s ² 3d ⁵	26 Fe 4s ² 3d ⁶	27 Co 4s ² 3d ⁷	28 Ni 4s ² 3d ⁸	29 Cu 4s ¹ 3d ¹⁰	30 Zn 4s ² 3d ¹⁰	31 Ga 4s ² 3d ¹⁰ 4p ¹	32 Ge 4s ² 3d ¹⁰ 4p ²	33 As 4s ² 3d ¹⁰ 4p ³	34 Se 4s ² 3d ¹⁰ 4p ⁴	35 Br 4s ² 3d ¹⁰ 4p ⁵	36 Kr 4s ² 3d ¹⁰ 4p ⁶
5 Rb 5s ¹	38 Sr 5s ²	39 Y 5s ² 4d ¹	40 Zr 5s ² 4d ²	41 Nb 5s ¹ 4d ⁵	42 Mo 5s ² 4d ⁵	43 Tc 5s ² 4d ⁵	44 Ru 5s ¹ 4d ⁷	45 Rh 5s ¹ 4d ⁸	46 Pd 4d ¹⁰	47 Ag 5s ¹ 4d ¹⁰	48 Cd 5s ² 4d ¹⁰	49 In 5s ² 4d ¹⁰ 5p ¹	50 Sn 5s ² 4d ¹⁰ 5p ²	51 Sb 5s ² 4d ¹⁰ 5p ³	52 Te 5s ² 4d ¹⁰ 5p ⁴	53 I 5s ² 4d ¹⁰ 5p ⁵	54 Xe 5s ² 4d ¹⁰ 5p ⁶
6 Cs 6s ¹	56 Ba 6s ²	57 La 6s ² 5d ¹	72 Hf 6s ² 5d ²	73 Ta 6s ² 5d ³	74 W 6s ² 5d ⁴	75 Re 6s ² 5d ⁵	76 Os 6s ² 5d ⁶	77 Ir 6s ² 5d ⁷	78 Pt 6s ¹ 5d ⁹	79 Au 6s ¹ 5d ¹⁰	80 Hg 6s ² 5d ¹⁰	81 Tl 6s ² 5d ¹⁰ 6p ¹	82 Pb 6s ² 5d ¹⁰ 6p ²	83 Bi 6s ² 5d ¹⁰ 6p ³	84 Po 6s ² 5d ¹⁰ 6p ⁴	85 At 6s ² 5d ¹⁰ 6p ⁵	86 Rn 6s ² 5d ¹⁰ 6p ⁶
7 Fr 7s ¹	88 Ra 7s ²	89 Ac 7s ² 6d ¹	104 Rf 7s ² 6d ²	105 Db 7s ² 6d ³	106 Sg 7s ² 6d ⁴	107 Bh 7s ² 6d ⁵	108 Hs 7s ² 6d ⁶	109 Mt 7s ² 6d ⁷	110 Ds 7s ² 6d ⁸	111 Rg 7s ² 6d ⁹	112 Cn 7s ² 6d ¹⁰	(113)	(114)	(115)	(116)	(117)	(118)

90 Th 6d ² 7s ²	91 Pa 6d ¹ 7s ²	92 U 6d ¹ 7s ²	93 Np 6d ¹ 7s ²	94 Pu 6d ¹ 7s ²	95 Am 6d ¹ 7s ²	96 Cm 6d ¹ 7s ²	97 Bk 6d ¹ 7s ²	98 Cf 6d ¹ 7s ²	99 Es 6d ¹ 7s ²	100 Fm 6d ¹ 7s ²	101 Md 6d ¹ 7s ²	102 No 6d ¹ 7s ²	103 Lr 6d ¹ 7s ²
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