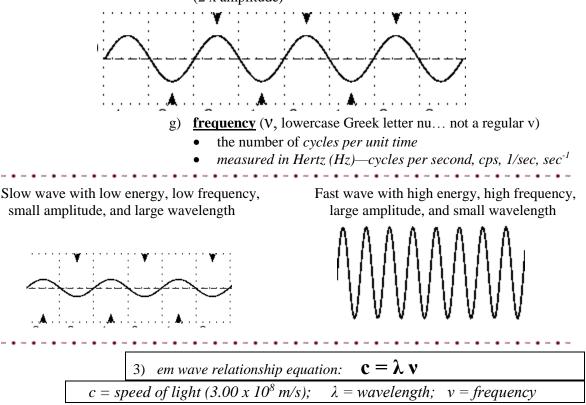
Ch. 4 Notes - ELECTRONS IN ATOMS

NOTE: Vocabulary terms are in **boldface and underlined**. Supporting details are in *italics*.

- I. Light and Quantized Energy
 - A. The Wave Nature of Light
 - 1) the wave nature of light: light acts as a moving wave
 - 2) wave anatomy and characteristics
 - a) **<u>origin</u>**, zero line, base line—flat line running horizontally, cutting the wave in two; line of zero movement
 - b) <u>crest (</u>"peak")
 - c) trough ("valley")
 - d) <u>amplitude</u>—vertical distance from the origin to the crest, or from the origin to the trough
 - e) wavelength (λ)—horizontal distance between two equivalent points on a wave, such as between two crests or two troughs
 - f) <u>wave height</u>—vertical distance from crest level to trough level; (2 x amplitude)



EXAMPLE 1) A wave has a frequency of 6.6×10^6 Hz. What is its wavelength?

$$c = \lambda v$$
 $\lambda = \underline{c} = 3.00 \text{ x } 10^8 \text{ m/s} = 45 \text{ m}$
 $v = 6.6 \text{ x } 10^6 \text{ 1/s}$

EXAMPLE 2) A wave is estimated to have a wavelength of 4.47×10^{-4} m. What is its frequency?

$$c = \lambda v$$
 $v = \underline{c}$ $= \underline{3.00 \times 10^8 \text{ m/s}}_{\lambda} = \underline{6.71 \times 10^{11} \text{ Hz}} (1/\text{s} = \text{Hz})$
 $\lambda = 4.47 \times 10^{-4} \text{ m}$

B. The Particle Nature of Light

the particle nature of light: light acts as a stream of moving bundles of light energy called <u>photons</u>
<u>wave-particle duality of nature</u>—waves can act as particles, AND particles can act as waves
<u>quantum</u>– a bundle of energy; the minimum amount that can be gained or lost by an atom
<u>photoelectric effect</u>— electrons (photoelectrons) being ejected from a metal when the metal is exposed to high-frequency light
<u>E = energy</u> of a wave, usually measured in Joules (J)

6) energy of a quantum or photon: **E** = **h v** *E* = energy; *h* = Planck's constant (6.626 x 10⁻³⁴ Js); *v* = frequency

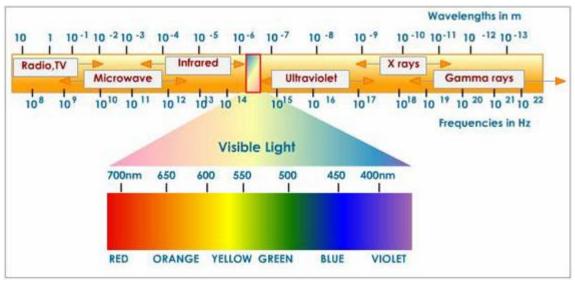
EXAMPLE 3) A photon in a wave has a frequency of 6.6×10^6 Hz. What is its energy?

E = h v $E = (6.6 \text{ x } 10^6 \text{ 1/s}) (6.626 \text{ x } 10^{-34} \text{ Js}) = 4.4 \text{ x } 10^{-27} \text{ Js}$

EXAMPLE 4) A quantum has a measured energy of 1.333×10^{-20} J. What is its frequency?

$$E = h v \qquad v = \underline{E}_{h} = \underline{1.333 \times 10^{-20} J}_{6.626 \times 10^{-34} Js} = \underline{2.012 \times 10^{13} Hz} (1/s = Hz)$$

- II. Spectra
 - A. Electromagnetic (em) spectrum—broad radiation spectrum



(source: Redmcolsoqo)

(low energy) **RADIO WAVES** ... **MICROWAVES** ... **INFRARED (IR) RADIATION** ... <u>VISIBLE LIGHT</u> "ROYGBIV": red, orange, yellow, green, blue, indigo, violet ... <u>ULTRAVIOLET (uv)</u> ... <u>X-RAYS</u> ... <u>GAMMA RAYS</u> ... <u>COSMIC RAYS</u> (high energy)

B. atomic emission spectrum

1) array of colors from a heated element, separated by a prism

2) set of frequencies of em waves emitted by atoms of an element

III. Electrons and Light

- A. evidence for energy levels: explanation of atomic spectra
 - 1) **ground state**—electron(s) at their lowest possible energy level
 - 2) <u>excited state</u>
 - a) *electron(s) at higher energy than normal*
 - b) *e- absorb energy and jump to higher levels where there is room*
- B. Niels Bohr (1885 1962); Bohr model proposed in 1913
 - 1) **<u>quantum number</u>** = n ("ring" number 1-7)
 - 2) nucleus as the center, composed of p^+ and n^0
 - 3) *e- orbit the nucleus; similar to planetary motion*
 - 4) e⁻ in an orbit have a fixed energy level
 - 5) lowest energy levels are closest to the nucleus
 - 6) **<u>quantum</u>** (see section I B)—*a bundle of energy* needed to make an electron "jump" to a higher level, which is a **<u>quantum leap</u>**
- C. <u>spectroscopy</u>—the study of substances exposed to continuous energy
 - 1) when exposed to intense energy, atoms absorb energy and become excited
 - 2) when atoms are in their excited state, their electrons jump to higher energy levels
 - 3) when the electrons eventually return to their normal (ground) state, energy is given off (emitted)

D. quantum mechanical model of the atom

- 1) electrons as waves
- 2) Louis deBroglie (1892-1987)
 - a) all moving particles have wave characteristics
 - b) particle em wave relationship equation: $\lambda = \underline{h}$

mv

 λ = wavelength; h = Planck's constant (6.626 x 10⁻³⁴ Js); m = mass; v = velocity (speed with magnitude and direction)

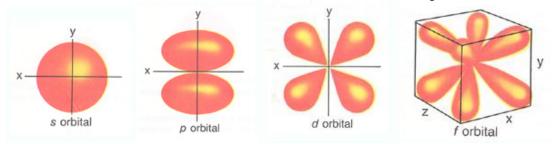
mv = momentum

- 3) **Heisenberg Uncertainty Principle**—*it is not possible to know both the exact position and velocity of an object simultaneously*
 - a) if you measure the position, you have to stop it (technically), so you cannot find the velocity or momentum
 - b) if you measure the velocity, it is moving, so you cannot pinpoint one position to measure
- 4) Erwin Schrödinger (1887-1961); quantum mechanical model 1926
 - a) based on probability of e⁻ location, not exact path
 - b) e cloud model; "boundary surface diagram"
 - c) equation: $\frac{\rho^2 \psi}{\rho x^2} + \frac{\rho^2 \psi}{\rho y^2} + \frac{\rho^2 \psi}{\rho z^2} + \frac{8\pi^2 m}{h^2} (E-V)\psi = 0$
 - d) e^{-} are mathematically treated as a wave
 - e) wave function—amplitude (ψ)

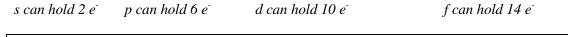
- f) solved the equation for a series of wave functions (definite energy states of an e⁻ and the probability of finding it there)
- IV. Electron configurations—Distribution of Electrons in Energy Levels
 - A. <u>electron configuration</u>—specific e⁻ arrangement in orbitals
 - B. <u>atomic orbital</u>—area of high probability (~95%) of finding electrons
 - C. principal quantum number = n
 - electron energy level or "shell" number
 n = 1, 2, 3, 4, 5, 6, 7
 - D. energy <u>sublevels</u>—divisions within a level
 - 1) many similar energy states grouped together in a level
 - 2) different shapes: spherical, dumbbell, cloverleaf
 - 3) sublevels: s, p, d, f
 - (s p d f stand for sharp, principal, diffuse, fundamental)
 - 4) maximum number of e^{-} in a principal energy level = $2n^{2}$

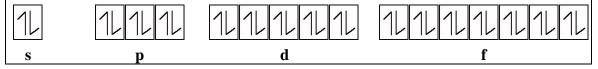
<u>n</u>	<u>SUBLEVELS</u>
n = 1	1s
n = 2	2s, 2p
n = 3	3s, 3p, 3d
n = 4	4s, 4p, 4d, 4f
n = 5	5s, 5p, 5d, 5f
n = 6	6s, 6p, 6d
n = 7	7s, 7p
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SHAPES OF ORBITALS (boomeria.org)



E. electron capacity of the sublevels





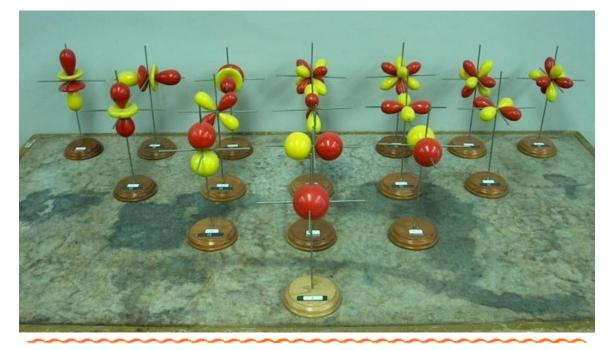
1) orbital = one "box;" can also be a line or a small circle

- 2) electrons = "arrows" (can be drawn as full \uparrow or half | arrows)
- 3) <u>antiparallel spin</u>—electrons in an orbital spin in opposite directions, so as not to repel one another (this is shown as arrows pointing in opposite directions)

4) rules

- <u>Aufbau Principle</u>—e⁻ enter low-energy orbitals first
- <u>**Pauli Exclusion Principle**</u>—no more than two e⁻ per orbital; the emust have compatible spins (shown by ↑ and ↓)
- <u>Hund's Rule</u>—degenerate orbitals (orbitals of equal energy) have one e⁻ filling each orbital before pairs can exist
- 5) sample electron configuration: $1s^2 2s^2 2p^6 3s^2 3p^2 = read "1 s 2, 2 s 2, 2 p 6, 3 s 2, 3 p 2"$ total e⁻ = 14 (Si)
- 6) the sum of the superscripts equals the atomic number of the element
- 7) *superscripts are NOT exponents* (nothing is being squared, etc.)
- 8) *** normal valence configurations will be <u>s</u> OR <u>s and p</u> ***
- 9) ways to do electron configurations: Aufbau (diagram or periodic table location) <u>Aufbau diagram</u> = orbital-filling diagram, orbital notation
- 10) Examples with the Aufbau diagram (see last page of notes for Aufbau diagram)
- EXAMPLE 5) Write out the electron configuration for nitrogen.
- EXAMPLE 6) Write the electron configuration for cadmium.

ANSWERS: E 5) $1s^2 2s^2 2p^3$ E 6) $1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^{10} 4p^6 5s^2 4d^{10}$



V. The Periodic Table and Atomic Structure

A. electron configurations according to location

- 1) Group IA (1) and IIA (2) = end in s
- 2) Group IIIA-VIIIA (13-18) = end in p
- 3) B Groups (Groups 3-12 transition metals) = end in d
- 4) Inner transition metals = end in f
- 5) Noble gases configurations end in $s^2 p^6$
- 6) Pseudo-Noble gas configurations are exceptions $= s^2 p^6 d^{10}$

B. Complete vs. valence electron configurations

- 1) complete—lists all orbitals (superscripts match the atomic number)
- 2) *valence—only lists the outer shell* (matches the dot diagram)

DOTS: "right, left, up, down, top all the way around, counterclockwise."

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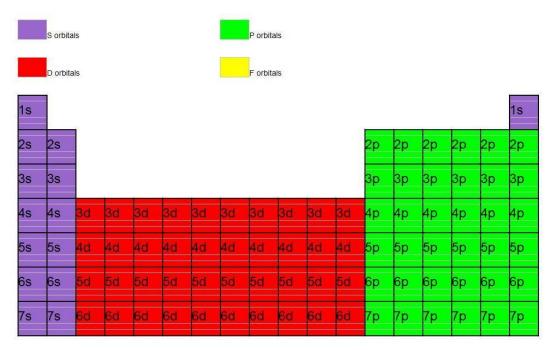
*** There are many ways to draw dots in Lewis structures, but this is the order we will use. ***

- C. Why do normal valence configurations show <u>s</u> OR <u>s and p</u> and not normally include d or f?
 - 1) *d* orbitals are "step-downs" they are one level below the valence
 - 2) *f orbitals are "two step-downs" they are two levels below the valence*

GROUP #	VALENCE #	SAMPLE VALENCE CONFIG.	# DOTS		
IA (1)	1	3s ¹	1		
IIA (2)	2	$3s^2$	2		
IIIA (13)	3	3s ² 3p ¹	3		
IVA (14)	4	$3s^23p^2$	4		
VA (15)	5	$3s^23p^3$	5		
VIA (16)	6	$3s^23p^4$	6		
VIIA (17)	7	$3s^23p^5$	7		
VIIIA (18)	8	$3s^23p^6$	8		

ELECTRON CONFIGURATIONS AND THE PERIODIC TABLE (scientifictutor.org)

(for black-and-white copies, arbitrary colors used are s = purple, p = green, d = red, f = yellow)



| 4f |
|----|----|----|----|----|----|----|----|----|----|----|----|----|----|
| 5f |

*** There is a discrepancy between the $5d^{1}/6d^{1}$ and $4f^{1}/5f^{1}$ positions

D. Noble Gas (Condensed, Abbreviated) Electron Configurations

1) use the previous Noble Gas as the starting point in brackets, then finish the configuration

2) Example 7: Indium #49 complete: Noble Gas (condensed): valence:

 $\begin{array}{l} 1s^2\,2s^2\,2p^6\,3s^2\,3p^6\,4s^2\,3d^{10}\,4p^6\,\underline{5s^2}\,4d^{10}\,\underline{5p^1}\\ [Kr]\,5s^2\,4d^{10}\,5p^1\\ 5s^2\,5p^1\end{array}$

- 3) Example 8: Titanium #22 complete: $1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^2$ condensed: [Ar] $4s^2 3d^2$ valence: $4s^2$
- E. *Exceptions* to electron configurations
 - 1) to fill a sublevel = stable
 - 2) to half-fill a sublevel = stable
 - 3) new unpredicted configurations are observed Cu, expected: $1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^9$ Cu, observed: $1s^2 2s^2 2p^6 3s^2 3p^6 4s^1 3d^{10}$

ELECTRON CONFIGURATIONS EXCEPTIONS AND THE PERIODIC TABLE (from hackaday)

