Measurement and Units

 SI – similar to (but not exactly the same as) the metric system

Measurement and Units

Physical Quantity	Name of unit	Symbol of Unit
Length	Meter	m
Mass	Kilogram	kg
Time	Second	S
Temperature	Kelvin	Κ
Amount of	Mole	mol
substance		
Electric current	Ampere	A
Luminous intensity	Candela	cd

Prefix	Multiple
Tera (T)	1012
Giga (G)	10 ⁹
Mega (M)	106
Kilo (k)	10 ³
Centi (c)	10-2
Milli (m)	10-3
Micro (u or μ)	10-6
Nano (n)	10 ⁻⁹
Pico (p)	10-12
Femto (f)	10 ⁻¹⁵

More on measurement

- Precision how "close" experimental values are to each other (consistency)
- Accuracy how "close" experimental values are to a "true" or "accepted" value
- "closeness" can be measured by a variety of statistical techniques – mean, median, mode, standard deviation, etc.

Significant figures

- We live in the real world, not in theory!
- Aid in reporting experimentally measured quantities
 - Any instrument used for measurement will have a specified precision (+/-)
 - -We are allowed to report <u>all known</u> digits and <u>one</u> *unknown* digit

Significant figures

- Any non-zero digit is significant (Ex. 1234)
- Zeros sandwiched between digits are significant (Ex. 1023)
- Zeros to the left of a decimal are NOT significant (Ex. 0.123)
- Zeros to the left of the first non-zero digit are NOT significant (Ex. 0.0000123)
- Zeros to the right of the last non-zero digit are significant (Ex. 0.123000)
- If there is no decimal point, zeros are NOT significant (Ex. 100 vs 100.)

Calculations involving significant figures

- "A chain is only as strong as its weakest link"
- Addition and Subtraction use the number with the least number of significant figures AFTER the decimal (or least number if there is no decimal)
- Multiplication and Division use the number with the least number of TOTAL significant figures
- Propagation of error round only at the last step of a multi-step calculation (but keep track of how many sig figs there should be at each point)

Dimensional Analysis and Unit Conversion

- Can be used as a problem-solving tool
- It is always a good idea to include units, not just numbers!
- Ex. How many seconds are in one year?

 $1 year \left(\frac{365 \ days}{1 \ year}\right) \left(\frac{24 \ hours}{1 \ day}\right) \left(\frac{60 \ minutes}{1 \ hour}\right) \left(\frac{60 \ seconds}{1 \ minute}\right)$

Chemical Reactions (Equations)

- Note: In this course the phases for each chemical reaction are omitted
- Example
 - $-2C_2H_6(g) + 7O_2(g) \rightarrow 4CO_2(g) + 6H_2O(I)$ will be written as
 - $-2C_2H_6 + 7O_2 \rightarrow 4CO_2 + 6H_2O$

Neutralizations

- Reaction of an acid with a base
 Acid + Base → Salt + Water
- Overall/Complete formula/Molecular reaction:
 HCl(aq) + NaOH(aq) → NaCl(aq) + H₂O(l)
- However, we should really show this reaction as it would "look" in solution

Neutralizations

- lonic equation:
 - $\begin{array}{l} -\mathsf{H}^{\scriptscriptstyle +}(\mathsf{aq}) + \mathsf{CI}^{\scriptscriptstyle -}(\mathsf{aq}) + \mathsf{Na}^{\scriptscriptstyle +}(\mathsf{aq}) + \mathsf{OH}^{\scriptscriptstyle -}(\mathsf{aq}) \xrightarrow{} & \mathsf{Na}^{\scriptscriptstyle +}(\mathsf{aq}) + \mathsf{CI}^{\scriptscriptstyle -}(\mathsf{aq}) + \\ \mathsf{H}_2\mathsf{O}(\mathsf{I}) \end{array}$
 - (water only dissociates about every 1 in 10⁷ molecules)
- Net ionic equation:
 - $-H^{+}(aq) + OH^{-}(aq) \rightarrow H_{2}O(I)$
 - *—Spectator ions* identity is irrelevant, however they are necessary for charge neutrality

Naming compounds

- Usually put metal first, then nonmetal (go from left to right from the periodic table)
- Exceptions N, H, O
- Name = firstelement secondelement(-ide)
- Prefixes
 - -Ionic substances generally have no prefixes
 - –Covalent substances prefixes are always used for the 2nd element (even if it only has one of them) but are only used for the 1st element if >1

Number	Prefix
1	Mono
2	Di
3	Tri
4	Tetra
5	Penta
6	Неха
7	Hepta
8	Octa
9	Nona
10	Deca

Ionic Compounds

- lons atoms that have gained or lost electrons (have + or – charge)
 - Can have very different properties than their corresponding elements
- Cations + charge (lost electrons)
 —Usually originate from metals
- Anions - charge (gained electrons)
 —Usually originate from nonmetals
- Ions can also be *polyatomic* (composed of more than one atom)

Determining the charge for an ion

 For Groups IA-VIIIA the "usual" charge of an ion is based on its position

-+1, +2, +3, +/-4, -3, -2, -1, 0

- For Group B (transition metals), use the Stock system
 - -Roman numerals represent charges

-Ex. Fe(II) ion = Fe^{2+}

Periodic Table of the Elements

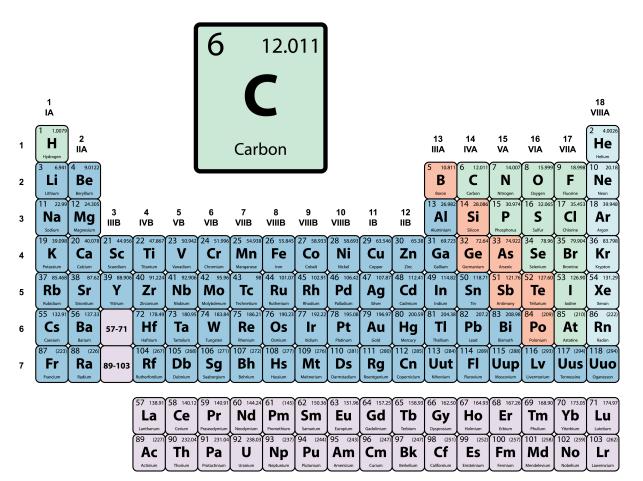


Table E Selected Polyatomic Ions

$\mathrm{H_{3}O^{+}}$	hydronium	CrO ₄ ²⁻	chromate
${{{\rm Hg}_{2}}^{2+}}$	dimercury (I)	Cr2072-	dichromate
$\mathrm{NH_4}^+$	ammonium	MnO ₄ ⁻	permanganate
C ₂ H ₃ O ₂ -	} acetate	NO2 ⁻	nitrite
CH ₃ COO	205	NO ₃ -	nitrate
CN-	cyanide	O ₂ ²⁻	peroxide
CO32-	carbonate	OH-	hydroxide
HCO3-	hydrogen carbonate	PO4 ³⁻	phosphate
C2042-	oxalate	SCN-	thiocyanate
ClO-	hypochlorite	SO3 ²⁻	sulfite
ClO ₂ -	chlorite	SO42-	sulfate
ClO ₃ ⁻	chlorate	HSO ₄ -	hydrogen sulfate
ClO ₄ -	perchlorate	S2032-	thiosulfate

Oxyanions

• Contain a varying number of oxygen atoms as part of a polyatomic ion

Oxyanion	Name
CIO ⁻	Hypochlorite
ClO₂ ⁻	Chlorite
ClO₃ ⁻	Chlorate
CIO₄ ⁻	Perchlorate

Formula Unit

- Strictly speaking, this term should be used to describe ionic compounds
- It represents the smallest collection of ions that combine to form something neutral
- Ex. NaCl, Al₂O₃
- In naming formula units, prefixes are NOT used.

Hydrates

- Chemicals that contain H₂O in their formula
- The water molecules are actually associated with the cations/anions in a well-defined way
- A prefix must be used to indicate the number of H₂O molecules
- Anhydrous (dry) no H₂O present
- Ex. $CuSO_4 vs. CuSO_4 \cdot 5H_2O$

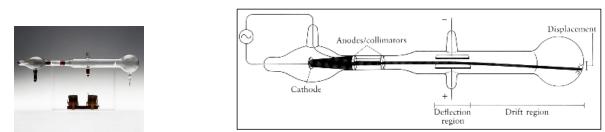


What's an atom made out of?

- All atoms are comprised of subatomic particles, which are fundamental.
- All subatomic particles are created equal
 - They are exactly the same, even if they are present in different atoms
- Three are important for chemistry
 - Proton
 - Neutron
 - Electron

J. J. Thomson (1897)

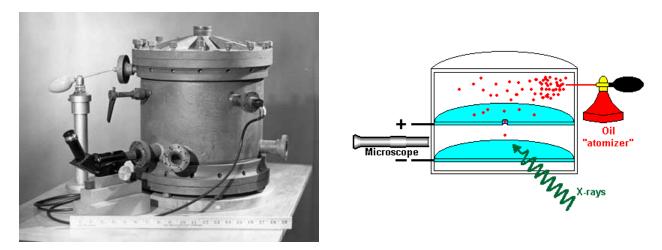
- Discovery of the electron
- (-) charged particles were produced, and they behaved exactly the same, regardless of the metal that was used.
- Was able to calculate the m/z ratio, -5.69X10⁻¹² kg/C, but wasn't able to get individual values for the mass or charge.



http://www.makingthemodernworld.org.uk/icons_of_invention/science/1880-1939/IC.026/ http://dbhs.wvusd.k12.ca.us/webdocs/AtomicStructure/Disc-of-Electron-Images.html

Robert Millikan (1909)

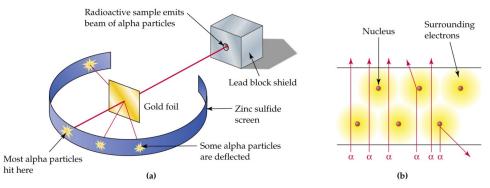
- Measured the velocity of a falling oil droplet in the presence/absence of a magnetic field
- Determined the charge on an electron (-1.602X10⁻¹⁹ C)
 - Mass of electron = $9.11X10^{-31}$ kg



http://www.juliantrubin.com/bigten/millikanoildrop.html

Ernest Rutherford (1911)

- α particle = $\frac{4}{2}He^{2+}$
- Most particle went straight through, but some were deflected
- Most of the atom is empty space, but all the (+) charge is concentrated in the center (nucleus)



http://wps.prenhall.com/wps/media/objects/602/616516/Media_Assets/Chapter02/Text_Imag es/FG02_05.JPG

Some definitions

- Isotope same # of protons, but different # of neutrons
- Atomic Number (Z) # of protons
- Mass Number (A, M) # of nucleons (protons and neutrons)
- Atomic Mass weighted average of all mass numbers (weighted by fractional abundance)

$$A.M.=\sum_{i}f_{i}M_{i}$$

The atomic mass unit (amu)

- One amu = 1/12 the mass of one atom of C-12 (by definition)
- This is the basic unit of mass for chemists, though it isn't an SI unit

Example

• Calculate the atomic weight of carbon.

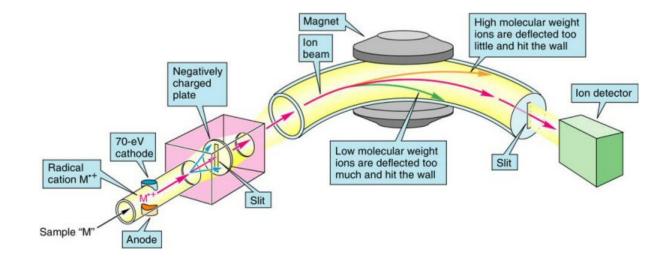
Solution

- Carbon exists in three isotopic forms: ¹²C, ¹³C and ¹⁴C.
- The relative abundances of these isotopes are approximately 98.8%, 1.1% and 0.1%, respectively (this can be determined by mass spectrometry)
- Therefore the atomic weight would be = 12(0.988) + 13(0.011) + 14(.001) = 12.011 amu

Gas chromatography – Mass spectrometry (GC-MS)

- Usually requires ionization
- Form charged species with an unpaired electron (radical)
- Fragmentation pattern
 - Based on broken chemical bonds
 - Each piece (fragment) has a characteristic m/z ratio
- Molecular jigsaw puzzle

Diagram of a mass spectrometer



http://chemistry.umeche.maine.edu/CHY251/Ch13-Overhead4.html

GC-MS instruments



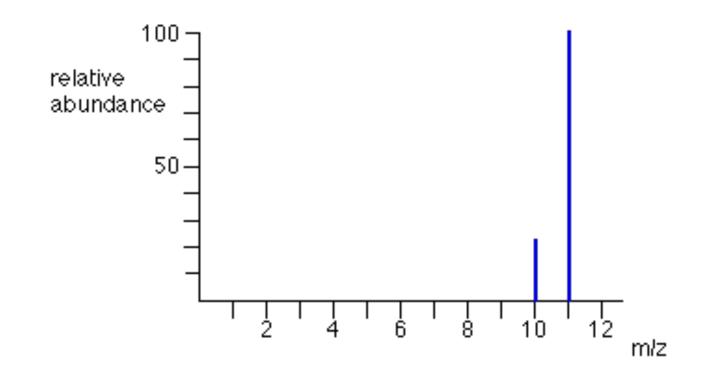
http://www.cooper.edu/~newmark/CH251/gcms.html

Common isotopic ratios

Element	lsotopes	Abundance (%)
Hydrogen	¹ H, ² H, ³ H	99.985, 0.015, (0)
Carbon	¹² C, ¹³ C, ¹⁴ C	98.90, 1.10, (0)
Nitrogen	¹⁴ N, ¹⁵ N	99.63, 0.37
Oxygen	¹⁶ 0, ¹⁷ 0, ¹⁸ 0	99.762, 0.038, 0.200
Chlorine	³⁵ Cl, ³⁷ Cl	75.77, 24.23
Bromine	⁷⁹ Br, ⁸¹ Br	50.69, 49.31

 Ratios can tell you which atoms you have present (by comparing relative intensities)

Example: Boron

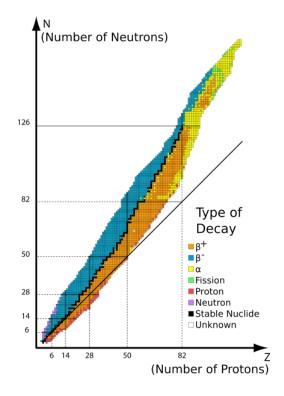


http://www.chemguide.co.uk/analysis/masspec/elements.html

Average atomic masses listed by IUPAC are based on a study of experimental results. Bromine has two isotopes ⁷⁹Br and ⁸¹Br, whose masses (78.9183 and 80.9163 amu) and abundances (50.69% and 49.31%) were determined in earlier experiments. Calculate the average atomic mass of bromine based on these experiments.

Nuclear Stability

- Nuclei can be predicted to be stable or unstable "radioactive" based on the number of nucleons (protons and neutrons).
- Generally if Z>84 (Po) the nuclide will undergo radioactive decay. All elements where Z>92 are "artificial" in the sense that they are not naturally occurring.
- For "small" nuclei, stable configurations are achieved when (A-Z)/Z is 1.
- For "large" nuclei, (A-Z)/Z is >1 (1.2-1.4)
- "Magic numbers" exist where nuclei are exceptionally stable: 2, 8, 20, 28, 50, 82, 126.



Radioactive Decay

- In order to achieve stability, radioactive nuclei will typically try to change their (A-Z)/Z ratio so they can fall in the band of stability. (α and β decays)
- It is also possible to become more stable yet keep the mass of the nucleus the same (γ decay)
- Other possibilities are *fission* (splitting of a heavy nuclide into smaller nuclides) and *fusion* (joining lighter nuclides into a heavier nuclide)

$\alpha \text{ decay}$

- Loss of a helium nucleus
 Results in ejection of positive particles
- Typically occurs with heavier nuclei
- Example $^{238}_{92}U \rightarrow ^{4}_{2}He + ^{234}_{90}Th$

β decay

- Common for medium-sized nuclides
- β^{-} decay loss of an electron $__{1e}^{0}$
 - Example ${}^{14}_{6}C \rightarrow {}^{0}_{-1}e + {}^{14}_{7}N$
 - Net conversion of a neutron into a proton [(A-Z)/Z too high] $\frac{1}{0}n \rightarrow -\frac{1}{1}e + \frac{1}{1}H$
- β^+ decay (positron emission) loss of an positron $\frac{1}{1}e$
 - Example $22_{11}^{22}Na \rightarrow 0_{1}^{0}e + 22_{10}^{22}Ne$
 - A positron is the *antiparticle* of an electron ${}^{0}_{1}e + {}^{0}_{-1}e \rightarrow 2{}^{0}_{0}y$
- <u>Electron capture</u> gain of an electron
 - Example $73_{33}As + 0_{-1}e \rightarrow 73_{32}Ge$
 - Net conversion of a proton into a neutron [(A-Z)/Z too low]

γ decay

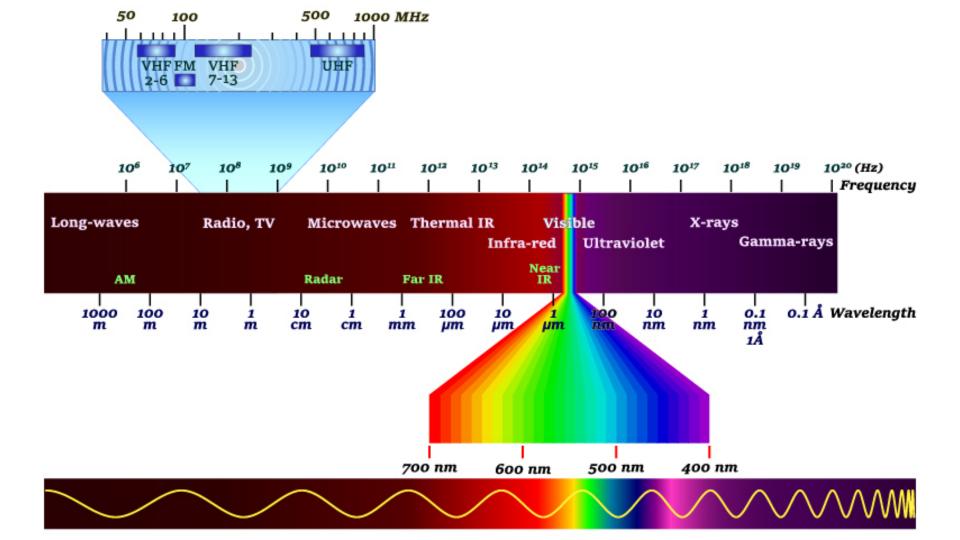
- Loss of a high energy photon
- No change in atomic or mass number
 - Example ${}^{99,m}_{43}Tc \rightarrow {}^{0}_{0}\gamma + {}^{99}_{43}Tc$
- We can think of the nucleons as being found in various energy levels, just like electrons

Nuclear fission

- Artificial transmutation process that releases a tremendous amount of energy $\frac{1}{0}n + \frac{235}{92}U \rightarrow \frac{141}{56}Ba + \frac{92}{36}Kr + 3\frac{1}{0}n$
- Typically initiated by a "magic bullet", commonly a neutron:
- Notice that for every one neutron that is used, three neutrons are produced. Each of these neutrons can then be used for another fission reaction, and so the reaction leads to an unstable (supercritical) situation since the number of particles grows exponentially. This is known as a *chain reaction*.

Light and spectroscopy

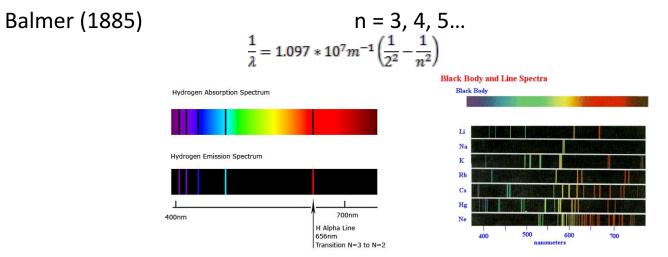
- EM Spectrum has waves of varying frequencies and wavelengths
- $E = hv = hc/\lambda$
- Spectroscopy deals with the interaction of matter with light



Atomic line spectra

- A cuvette filled with a sample is then exposed to a beam of light. Since light of all possible wavelengths are incident, it was believed that all possible wavelengths should be emitted, so the spectrum should be a rainbow (continuous emission)
- However, something else was observed...

٠



http://www.faculty.virginia.edu/consciousness/new_page_6.htm http://www.astronomyknowhow.com/hydrogen-alpha.htm

A simple, yet revolutionary idea

• Planck proposed that energy is quantized:

E=hv

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

"Old" Quantum Mechanics

- Niels Bohr (1913)
 - Assumed that the angular momentum (not the energy!) of the electron in a hydrogen atom is quantized
 - Used a combination of classical physics and this new interpretation for energy to derive "orbits", or energy levels (very similar to a planetary model)

$$E_n = -\frac{me^4}{8\varepsilon_o h^2} \frac{1}{n^2} = -\frac{B}{n^2}$$

 This was based on well-understood fundamental constants in physics (and Planck's constant)

A theoretical explanation of atomic line spectra

- Photons of light are emitted when electrons go from a higher to lower energy level (opposite is true for absorption)
- Because the energy levels are fixed, only certain wavelengths of light will be observed

The good, the bad and the ugly

- The good
 - Bohr was able to come up with a theoretical model for the energy levels in the hydrogen atom which accounted for the experimentally observed line spectra (Balmer series)
- The bad
 - It only worked for hydrogen!!! (and other one-electron systems)
- The ugly
 - The necessary mathematics get very difficult very quickly
 - Multi-electron systems often don't have closed form solutions

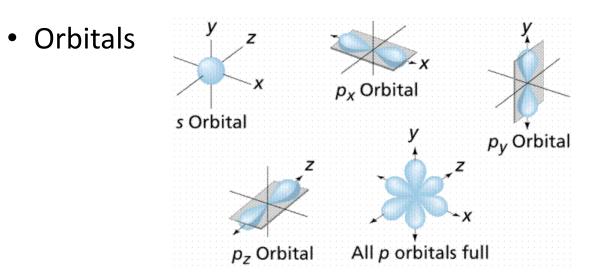
Quantum numbers

- Principal quantum number (n)
 - n = 1, 2, 3...
 - Same as Bohr's energy levels
 - Indicates what "shell" the electron is in
- Angular momentum quantum number (I)
 - $-l \le n-1$
 - − Ex. I=0 → s orbital, I=1 → p orbital, I=2 → d orbital, I=3 → f orbital
 - Determines the shape of the orbital, or "subshell"

Quantum numbers

- Magnetic quantum number (m_l)
 - $\mid m_l \mid \leq 1$
 - Determines the spatial *orientation* and degeneracy of the orbital
 - Ex. if l=1 (p orbital) then $m_l = -1$, 0, 1. These are usually called p_x , p_y , and p_z (directions do not directly correspond to these numbers). We can also see why there are three p orbitals, since there are three allowed values for m_l .

Subshell (orbital) shapes



 Nodes are possible – regions of zero probability of finding the electron

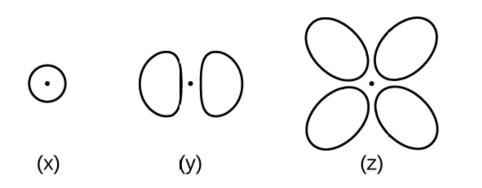
http://www.emc.maricopa.edu/faculty/farabee/biobk/biobookchem

Quantum numbers

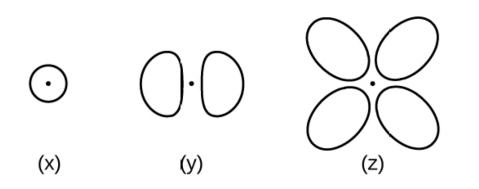
- Spin quantum number (m_s)
 - Unrelated to the other three quantum numbers
 - Unrelated to spatial coordinates
 - Each electron has an "intrinsic" spin coordinate
 - There is no classical analog, but it behaves similar to angular momentum
 - $-m_s = +/- \frac{1}{2}$ (half-integer)

- "All electronic wave functions must be antisymmetric under the interchange of any two electrons"
- It is impossible for two electrons in the same orbital to have the same spin

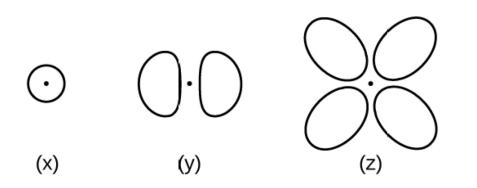
 No two electrons can have identical quantum numbers (in the same atom)



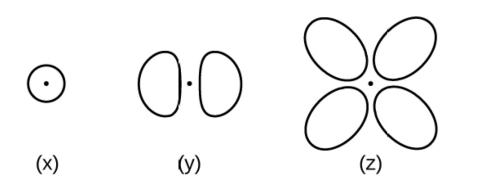
(a) What is the maximum number of electronscontained in an orbital of type (x)? Of type (y)? Oftype (z)?



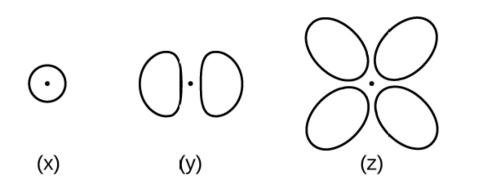
(b) How many orbitals of type (x) are found in a shell with n = 2? How many of type (y)? How many of type (z)?



(c) Write a set of quantum numbers for an electron in an orbital of type (x) in a shell with n = 4. Of an orbital of type (y) in a shell with n = 2. Of an orbital of type (z) in a shell with n = 3.

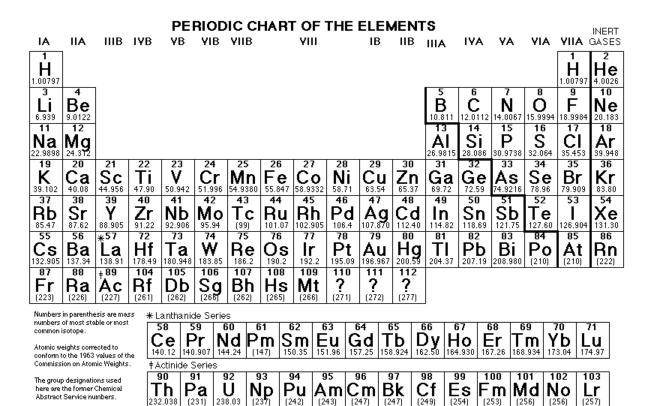


(d) What is the smallest possible *n* value for an orbital of typ



(e) What are the possible *I* and *mI* values for an orbital of type (x)? Of type (y)? Of type (z)?

The periodic table



Main-group elements ("the A-list")

- The valence (outer shell) consists only of s and p orbital electrons
- Group number = # of electrons in the valence shell (using the older Roman numeral system)
- Period number = principal quantum number (n)
- s block alkali metals and alkali earth metals
- p block metals, metalloids and nonmetals (including halogens and noble gases)

Transition metals ("the B-team")

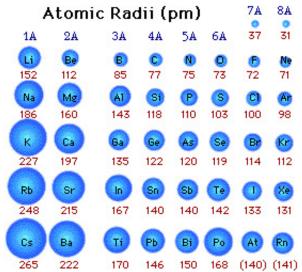
- Contain d and f orbitals
- d block transition metals
- f block rare earth (lanthanide/actinide)
- These are considered "inner shell" electrons
- The highest energy electrons are actually in a shell with a smaller value of n that that of the outermost shell (valence shell)
 - d block (n-1)
 - -f block (n-2)

Periodic Trends

- Patterns that emerge in chemical and physical properties when elements are arranged in the periodic table
- Can usually be explained by the number of valence electrons, the number of core electrons, and the number of protons (nuclear charge)

Atomic Radius

- Generally atomic radius decreases across a period and increases down a group
 - The trend only works for main group elements



http://edtech2.boisestate.edu/kilnerr/502/jigsaw.html

Atomic Radius

- Group increase in the number of principal energy levels (greater average distance that the electron is from the nucleus)
- Period increase in effective nuclear charge (the net charge the valence electrons "feel")

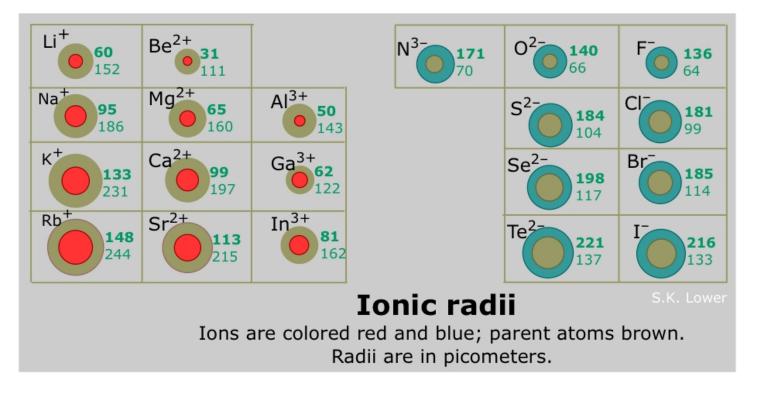
Effective Nuclear Charge (Z_{eff})

- Z_{eff} is meant to incorporate the shielding effect of core (inner) electrons
 - Valence electrons can penetrate inner shells (ex 3d and 4s)
 - Core electrons are not all equally effective in shielding valence electrons
 - Valence electrons can shield each other, though the effect is weak
- Transition metals in the same period have almost the same radius since $\rm Z_{eff}$ is the same

Ionic Radius

- Defined in a similar fashion to atomic radius (distance between two ions in a formula unit)
- Metals tend to lose valence electrons, so their highest occupied principal energy level decrease by one
 - Ionic radii for metals are smaller than those of the corresponding atomic radii
- Nonmetals tend to gain valence electrons, so their highest occupied principal energy level remains the same, but there is increased repulsion among the electrons in that level
 - Ionic radii for nonmetals are larger than those of the corresponding atomic radii

In pictures



http://chemwiki.ucdavis.edu/Wikitexts/UC_Davis/UCD_Chem_124A%3A_Kauzlarich/ChemWiki _Module_Topics/Periodic_Trends_in_Ionic_Radii

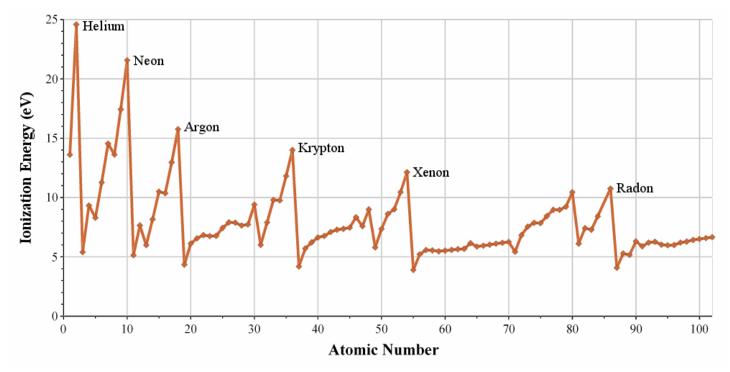
Ionization Energy

- Defined as the energy required to *remove* an electron from the ground state, in the gas phase
 - $A(g) \rightarrow A^+(g) + e^-$
- This can be repeated successively (1st, 2nd, 3rd, etc.)
 - It gets progressively harder to remove electrons since the species is already charged
 - Large jumps occur for a given element as you break up an octet (going from valence electrons to core electrons)

Ionization Energy

- Generally decreases as you go down a group
 - Outermost electrons are (on average) further away from the nucleus, so there is a greater shielding effect
- Generally increases as you go across a period
 - Elements have a greater tendency to gain electrons (rather than lose)
- Minor effects can be due to
 - what subshell the electron is in (s>p>d>f because of energy)
 - Paired vs. unpaired electrons (unpaired>paired because of repulsions)

In pictures



http://websites.pdesas.org/jvogus/2010/5/18/44324/page.aspx

Electron Affinity

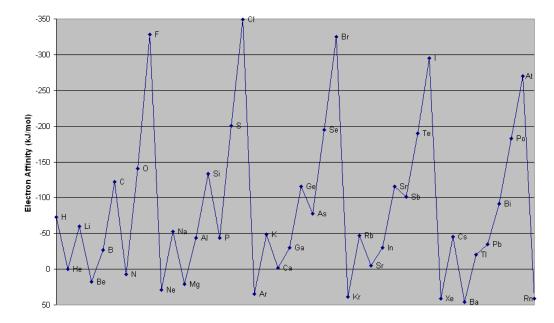
Defined as the energy required to *add* an electron to the ground state, in the gas phase

 $-A(g) + e^{-} \rightarrow A^{-}(g)$

- This has the same general trend as ionization energy, although it is less clear-cut
 - Complications due to repulsions between the incoming electron and the atomic electrons

In pictures





http://www.angelo.edu/faculty/kboudrea/periodic/trends_electron_affinity.htm

Electronegativity

• "Tendency" of an element to gain electrons

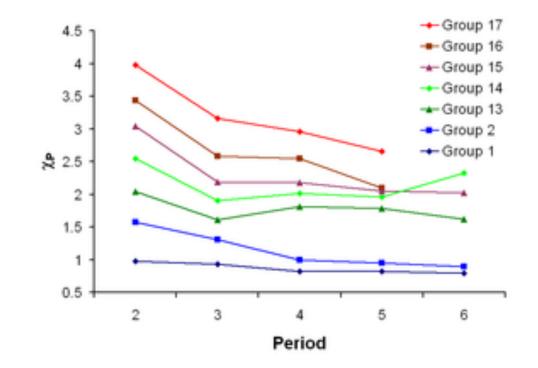
$$\chi = \frac{I.E. - E.A.}{2}$$

• Pauling scale:

$$\chi_B = \chi_A + 0.102 \{ (A - B) - [(A - A)(B - B)]^{1/2} \}^{1/2}$$

- (i-j) = bond-dissociation energy between i and j
- F is arbitrarily given the maximum value of 4.0
- Also follows the same general trend as ionization energy and electron affinity

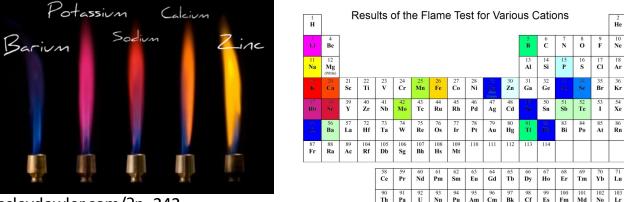
In pictures



http://en.wikipedia.org/wiki/Electronegativity

Chemical Properties of Elements

- Flame test
 - Based on characteristic absorbance of light energy
 - Wavelength emitted will be related to the energy gap between electronic levels
 - Used to identify various metals



http://wesleydowler.com/?p=242

http://alchemist.edublogs.org/2008/11/17/which-ion-causes-the-color/

Redox reactions (an introduction)

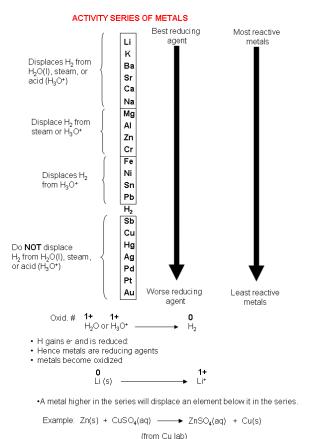
- Redox reactions involve a simultaneous *reduction* and *ox*idation.
- Reduction gain of electrons
 - oxidation number is decreased
- Oxidation loss of electrons
 - Oxidation number is increased
- Disproportionation redox reaction where the same species is both oxidized and reduced.

 $- \text{Ex. } 2\text{H}_2\text{O}_2 \rightarrow 2\text{H}_2\text{O} + \text{O}_2$

Agents

- Reducing agent causes a reduction
 - Gets oxidized
 - Usually metal
- Oxidizing agent causes an oxidation
 - Gets reduced
 - Usually nonmetal

Activity series



"Noble" metals (Cu, Ag, Hg, Au)
 – can't produce H₂

http://employees.csbsju.edu/hjakubowski/cla sses/ch123/summer_chem/ch123OLSGMM04 05.htm

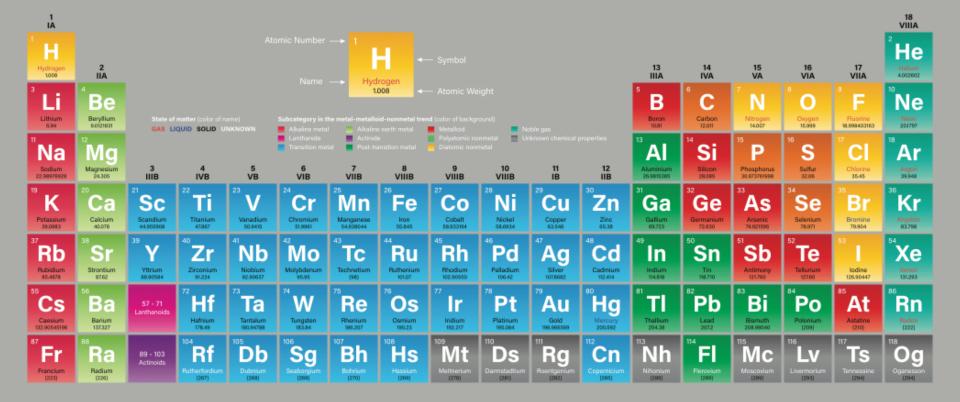
Chemical Properties of Elements

- Reduction
 - Reducing agents have a tendency to lose electrons
 - This property can be correlated with ionization energy, electronegativity and electron affinity
- Metals can react with sources of H⁺ (acids, or even water if they are active enough) to generate ions and hydrogen gas
- Mg + 2H⁺ \rightarrow Mg²⁺ + H₂
- Ca + 2H₂O \rightarrow Ca²⁺ + 2OH⁻ + H₂

Chemical Properties of Elements

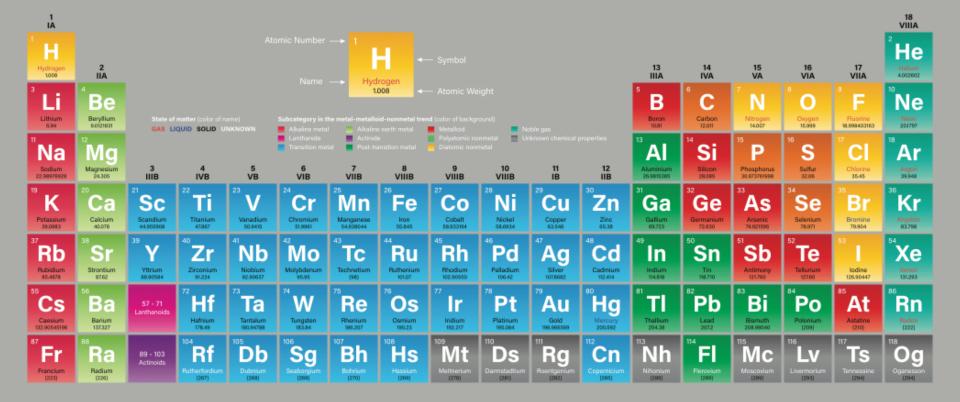
- Oxidation
 - Oxidizing agents have a tendency to gain electrons
 - This property can be correlated with ionization energy, electronegativity, and electron affinity
- Cl₂ + 2l⁻ → 2Cl⁻ + l₂ will occur since Cl atoms have a higher (more negative) electron affinity than I atoms (-349 kJ/mol vs. -295 kJ/mol)
- $I_2 + CI^- \rightarrow 2I^- + CI_2$ will NOT occur

List the following ions in order of increasing radius: Li⁺, Mg²⁺, Br⁻, Te²⁻.



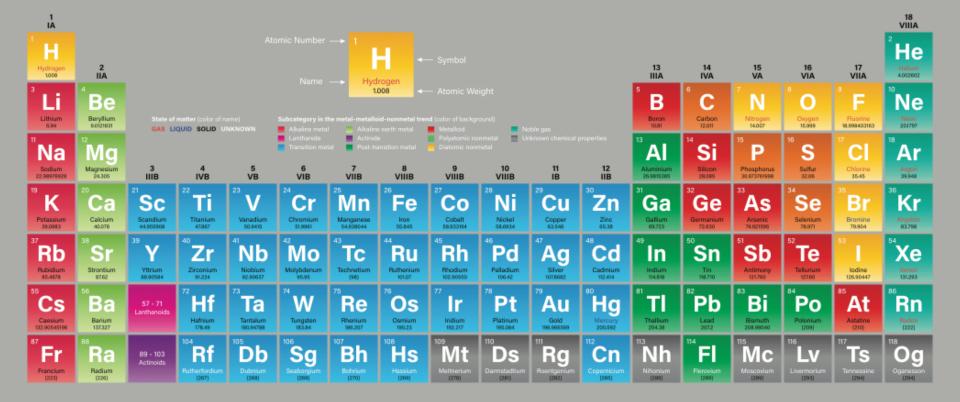


Write the Lewis structure for SeCl₃⁺.





Explain why the H₂O molecule is bent, whereas the BeH₂ molecule is linear.





Reactions of alkali metals

- With halogens (F_2, Cl_2, Br_2, I_2) : - M + X₂ \rightarrow MX
- With hydrogen: - $M + H_2 \rightarrow MH$
- With (excess) oxygen:
 - − Li + O_2 → Li₂O (plus some Li₂O₂)
 - − Na + $O_2 \rightarrow Na_2O_2$ (plus some Na_2O)
 - $-M + O_2 \rightarrow MO_2$ (M = K, Rb, Cs)
- With water:
 - $-M + H_2O \rightarrow MOH + H_2$

Reactions of alkaline earth metals

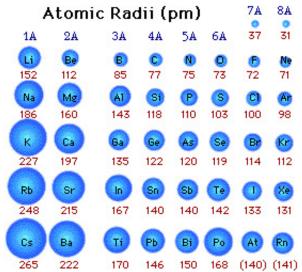
- With halogens (F_2, Cl_2, Br_2, I_2) :
 - $-M + X_2 \rightarrow MX_2$
- With nitrogen:
 - $-M + N_2 \rightarrow M_3N_2$
- With oxygen:
 - $-M + O_2 \rightarrow MO$
- With water:
 - $-Mg + H_2O(g) \rightarrow MgO + H_2$
 - $-M + H_2O \rightarrow M(OH)_2 + H_2 (M \neq Mg)$

Periodic Trends

- Patterns that emerge in chemical and physical properties when elements are arranged in the periodic table
- Can usually be explained by the number of valence electrons, the number of core electrons, and the number of protons (nuclear charge)

Atomic Radius

- Generally atomic radius decreases across a period and increases down a group
 - The trend only works for main group elements



http://edtech2.boisestate.edu/kilnerr/502/jigsaw.html

Ionization Energy

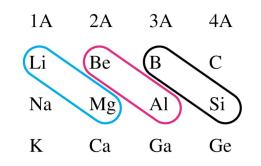
- Generally decreases as you go down a group
 - Outermost electrons are (on average) further away from the nucleus, so there is a greater shielding effect
- Generally increases as you go across a period
 - Elements have a greater tendency to gain electrons (rather than lose)
- Minor effects can be due to
 - what subshell the electron is in (s>p>d>f because of energy)
 - Paired vs. unpaired electrons (unpaired>paired because of repulsions)

Overview - Periodic Trends in Group 13

- B is a nonmetal/metalloid forms covalent bonds but displays electrical properties of semiconductors (diagonal relationship with Si)
- Al is a metal/metalloid –forms covalent bonds but can also lose valence electrons to form ions (Al³⁺)
- Ga forms Ga³⁺ ions to achieve stable configuration ([Ar]3d¹⁰)
- In and TI tend to form +1 ions because they lose the valence p electron but NOT the valence s electrons (inert pair)

Diagonal relationships

- Often the 1st member of a group has properties that are different from the other members of the group, but are similar to those of the 2nd member of the adjacent group
 - Relatively high charge density



- Example: Li
- Li₂CO₃, LiF, LiOH and Li₃PO₄ are much less soluble than the corresponding salts of the other alkali metals
 - Li_2CO_3 and LiOH form Li_2O
- Li + N₂ → Li₃N (other alkali metals don't react)
- Li + O₂ → Li₂O (other alkali metals form peroxides or superoxides)