### CHEMISTRY 20 -- UNIT 3: CHEMICAL BONDING

#### **INTRODUCTION**

- A. *Periodicity*: \_\_\_\_\_\_ that can be obtained from the periodic table (Chapter 12)
  - 1. Groups & Period Characteristics
    - a. 3 MAIN GROUPS:
    - b. 5 GROUP NAMES:
    - c. PERIODS:
  - 2. Reactivity on the periodic table
    - \* Most reactive metal: \_\_\_\_\_; Most reactive nonmetal: \_\_\_\_\_
    - \* Halogens react vigorously with \_\_\_\_\_ metals because they transfer electrons
    - \* Alkali metals and Alkaline metals react with \_\_\_\_\_ to form \_\_\_\_\_ coats
    - \* Alkali metals also react violently with \_\_\_\_\_
    - \* Nobel gases are very \_\_\_\_\_ because their orbitals are full
  - 3. State: solid, liquid or gas at room temperature
- B. Electrons in Atoms (Chapter 11)

History:

- 1. J. J. Thomson's Billiard Ball Model:
- 2. E. Rutherford's Raisin Bun Model:
- 3. Bohr's Energy level Model
  - a. <u>Energy levels</u>:
  - b. Energy level diagrams for atoms
  - a. Energy level diagrams for ions
- 4. *Quantum mechanics* Orbital Model

	Ionic	Molecular
State		
Color		
Solubility		
Conductivity		
Bond		
Ions Present		
Components		
Melting point		
Naming Rules	1.	1.
	2.	2. Common: water, ozone, peroxide, ammonia, methane, ethane, methanol, ethanol, sucrose, glucose
Formula Rules	1. 2. 3. 4. 5.	1. 2.

## C. Properties of Ionic and molecular compounds

- D. What is a chemical *Bond*
- A chemical bond is \_\_\_\_\_
- Atoms try to achieve the electron structure (not proton structure) or the nearest \_\_\_\_\_
- When a chemical bond is formed, the atoms form a more stable energy level. They will have a net \_\_\_\_\_\_ of energy to obtain this. Forming a chemical bond is \_\_\_\_\_\_

ie)  $H + H = ---> H_2 + 435 \text{ kJ}$ 

\*

- When a chemical bond is broken, the atoms will have a net \_\_\_\_\_\_ of energy. Breaking a chemical bond is \_\_\_\_\_\_
- ie)  $H_2 + 435 \text{ kJ} ----> H + H$
- GRAPH: Potential Energy vs distance for a Hydrogen bond

\* There are two main types of bonding forces

a. Intermolecular forces: \_\_\_\_\_

ie)

The amount of force depends upon many factors like size, polarity, number and type of electrons etc...

b) Intramolecular forces:

ie)

\* Approximate Bond Force strength..

Strongest						> Weak	est
	>	>	>	>	>	>	

In the flow diagram, place INTRA below the bonding force that is intramolecular and INTER below the force that is intermolecular.

#### VALENCE ORBITALS

A. Bohr said that electrons exist in energy \_\_\_\_\_\_ outside the nucleus. These \_\_\_\_\_\_ are regions around the nucleus where you would most likely find the electrons. B. <u>AUFBAU'S PRINCIPLE:</u>

Valence electrons

(free electrons) Group # = # of valence electrons.

Valence orbital \_\_\_\_\_

<u>Valence orbital</u> (<u>Valere</u> in Latin means to be strong.)

C. The rules of valence electrons are as follows

# 1. PAUL'S EXCLUSIO<u>N PRINCIPLE</u>:

2.

3. *HUND'S RULE*: \_\_\_\_\_\_

lone pairs.	
bonding electron	
4. <u>OCTET RULE</u> :	
Exceptions:	
EXAMPLE: sulphur atom	
6e <sup>-</sup> 2e,2e,1e,1e	Valence orbital with 2 lone pairs & 2 bonding electrons
8e <sup>-</sup> 2e,2e,2e,2e	
2e <sup>-</sup> 2e	
16p <sup>+</sup>	
S	

## D. Electron Configurations (optional)

1. There are principle energy levels with sublevels. Each sublevel has a different cloud shape and can hold a different number of electrons.

s orbital	holds 2 e	spherical
p orbital	holds 6 e	dumbbell shaped
d orbital	holds 10 e	complex
f orbital	holds 14 e	complex

The principle energy levels fill the sublevels as follows:

1s <sup>2</sup>			
2s <sub>2</sub>	$2p^6$		
$3s^2$	3p <sup>6</sup>	3d <sup>10</sup> (period 4 atoms)	
$4s^2$	$4p^6$	4d <sup>10</sup> (period 5 atoms)	4f <sup>14</sup> (period 6)
$5s^2$	5p <sup>6</sup>	5d <sup>10</sup> (period 6 atoms)	5f <sup>14</sup> (period 7)
$6s^2$	бр <sup>6</sup>	6d <sup>10</sup> (period 7 atoms)	
$7s^2$	7p <sup>6</sup> (future elements)		

Electrons wish to find the lowest possible energy state

The s orbitals fill the alkali metals and alkaline earths

The p orbitals are the last 6 elements on the right of each row. The very last is the Noble gas.

The "d" orbitals form the transition metal part of the table

The "f" orbitals form the rare earths (lanthanide and actinide series)

#### LEWIS DIAGRAM OR ELECTRON DOT DIAGRAM FOR ELEMENTS & IONS

Gilbert Lewis (1875 - 1946) developed a scheme for drawing particles with valence electrons shown as dots: <u>Lewis Dot Diagram</u>. (see page 300 for chart of beginning elements in the periodic table)

- A. <u>Electron dot</u>

C. Exam	ples			
Element or Ion	Number of valence electrons	Electron Dot Diagram	Number of bonding electrons	Number of lone pairs
calcium (Ca <sup>+</sup> )				
calcium (Ca)				
germanium (Ge)				
potassium (K)				
sulphur (S)				
nitrogen (N)				
aluminum (Al)				
neon (Ne)				
flouride (F <sup>-</sup> )				
fluorine (F)				
helium (He)				

D. Incorrect Lewis diagrams:

1. C 2. P 3. B 4. Na 5. P 6. S

E. Lewis Diagrams for Molecular Compounds

\* Sum the valence electrons for all the atoms in the molecule. Only this number of valence electrons can be used.

\* Determine if necessary which atom can form the most number of bonds. & This is the

	<ul> <li>* Valence electrons obey the rule (except hydrogen)</li> <li>EXAMPLES:</li> <li>1) Cl<sub>2</sub></li> </ul>
	2) CH <sub>4</sub>
F.	Lewis Diagrams for multiple bonds (F & G) above * Same rules as above * covalent bonds have two pairs of electrons shared between atoms * covalent bonds have three pairs of electrons shared between atoms 1) O <sub>2</sub>
	2) CO <sub>2</sub>
	3) N <sub>2</sub>
G.	Lewis Diagrams for Ionic Compounds * Electrons are from metal to nonmetal * The electrons around the metal are * Charged species are enclosed in EXAMPLES: 1) NaCl
	2) $MgS_2$
H.	<ul> <li>Lewis Diagrams for Complex Ions (OPTIONAL)</li> <li>* the valence electrons for all the atoms in the molecule. Add electrons for negative charges and subtract for positive charges.</li> <li>* Arrange the electrons around the atom so the sum is achieved. Use double and triple bonds if necessary.</li> <li>* Charged species are enclosed in</li> <li>EXAMPLES:</li> <li>1) Cyanide</li> </ul>
	2) Sulphate
	3) Ammonium

#### **BONDS**

## I. Ionic Bonding

A. Review

\* Ions:

Metallic ions:	(electronegativity)
Non-metallic ions:	( electro)
Complex ions:	

\* <u>Ionic bond:</u>\_\_\_\_\_

\* Ionic bonds produce a \_\_\_\_\_\_ with a repeating pattern.

\* Ionic compounds characteristics (Look at beginning of your notes)

B. Electron dot diagrams to show ionic compounds (Review)

Na + Cl  $\rightarrow [Na]^+[Cl]^-(brackets show that they are not sharing)$ 

C. Oxidation - Reduction (LEO goes GER; OIL RIG) (Optional)

1. <u>Oxidation</u>: \_\_\_\_\_\_\_of electrons (metal) Na ----> Na<sup>+</sup> + 1e<sup>-</sup>(half reaction)

The lower the electronegativity, the easier it is oxidized

2. <u>*Reduction*</u>: \_\_\_\_\_\_\_\_ of electrons (non-metal) Cl + 1e<sup>-</sup> ----> Cl<sup>-</sup>

. Stopping Corrosion (optional)

Method	Examples	EXPLANATION
Plating	* w/gold or silver or tin	* Covers the reactive metal
<u>Cathodic protection</u>	*blocks of Zn or Mg in an Fe/Be ship.	Attach a more reactive metal that corrode/oxidize easier. This saves the other metal
Oxide coating	chromium(III) oxide, aluminum oxide, zinc oxide, tin(IV) oxide	<i>Galvanization</i> : Covers the steel with ZnO
Creating alloys	stainless steel	
Painting or covering with plastic		

D. Ionic models (optional diagrams)

# II. <u>Metallic bonding</u>:

PROPERTIES:		 
*		
*		
* Diagram		

ie) Al, Fe, Cu, Zn, Ca, Na, Ag, Pb, Hg

III. <u>Network covalent bonding</u>:\_\_\_\_\_\_

PROPERTIES:
\*\_\_\_\_\_\*\_\_\_\_\_

\* Diagram

ie) diamonds (C<sub>n</sub>), Carborundum (SiC),

## III. Covalent bonds:

A. Introduction

1. Summary:

\* occur between \_\_\_\_\_\_ atoms and result in the mutual \_\_\_\_\_ of e-

\* unpaired electrons between two atoms that collide, are shared until they

have an octet (except Hydrogen) & have the same electron structure as a

Noble gas. ie) Cl + Cl ----- Cl Cl

\* Covalent bonds may be single, double or triple

# 2. <u>Bonding Capacity</u>: The maximum number of bonds a molecular atom can form.

Covalent bonds are represented by a dash. (- single, = double,  $\pm$  triple)

Атом	# OF VALENCE Electrons	# OF BONDING Electrons	BONDING CAPACITY & TYPES OF BONDS
hydrogen (H)			
carbon (C)			
nitrogen (N)			
oxygen (O)			
fluorine(halogen)(F)			
neon(noble) (Ne)			

3. Draw the *structural diagram* by replacing the shared pairs with a line (bond) &

omitting the lone pairs. The *molecular formula* is written with only the symbols.

$\mathcal{O}$ 1	<i>u</i>		5
<u>Examples</u>	<u>Molecular</u>	<u>Lewis</u>	<u>Structural</u>
a. Diatomic:			

b. One Lone pair:

c. Double bond

d. Triple bond:

e. Cyclic carbon compounds:

4. Problems with the lewis diagram model
<u>Coordinate Covalent Bond</u>:

a.	CO; C <u>≤</u> O; C O
b.	$\mathrm{NH_4}^+$
c.	$H_3O^+$

# Resonance:

<u>Hybrid</u>:

ie) ozone

Exceptions to the octet rule (pg. 326)

a. NO<sub>2</sub>

b. P and S may sometimes expand the octet to ten or twelve electrons

#### SUMMARY

\* Lewis diagrams tell us: 1) electrons involved in bonding 2) how many bonds

\* Lewis diagrams do not tell us: 1) what an atom looks like 2) where electrons are

\* Bonds are formed by \_\_\_\_\_ electrons

\* \_\_\_\_\_ electrons are involved in each bond, called a \_\_\_\_\_\_ bond

\* Diagrams and structural formula's usually satisfy the \_\_\_\_\_ rule

Stereochemistry:	
* <u>VSEPR Theory</u> : . ASSUMPTIONS:	
* all molecules have a 3-D shape	
1	r of bonding electrons and the central atom
must be known to determine the sha	
. RULES	pe.
. SHAPES & POLARITY	
1) Tetrahedral	e.g.) CH <sub>4</sub>
2) <u>Pyramidal</u>	e.g.) NH <sub>3</sub>
3) <u>V-shape</u>	e.g.) H <sub>2</sub> O
4) <u>Trigonal Planar</u>	e.g.) C <sub>2</sub> H <sub>4</sub>
5) <u>Linear</u> : *	e.g.) HCl
*	e.g.)Cl <sub>2</sub>
*	e.g.) C <sub>2</sub> H <sub>2</sub>
*	

\_

6) Combinations: e.g.) CH<sub>3</sub>OH(tetrahedral around C & V-shaped around O)

## C. Electronegativity:

(electron attractivity or an atoms desire for an octet)

- \* Principles of electronegativity developed by Linus Pauling (on periodic table)
  - 1. As you go left and down the periodic table(metals), e- are held more \_\_\_\_\_\_.
  - Therefore the element's reactivity to lose e- increases & electronegativity \_\_\_\_\_.
    - (Least electronegative: Fr,Ce = 0.7)
  - 2. As you go right and up the periodic table(non-metals), e- are held more \_\_\_\_\_.

Therefore the element's reactivity to gain increases & electronegativity \_\_\_\_\_.

\_\_\_\_\_ (Most electronegative: F = 4.0)

3. NOBLE GAS EXCEPTION: The noble gases hold their electrons most tightly,

but have \_\_\_\_\_\_ and are hardly reactive because they have an \_\_\_\_\_.

- Bond type based on electronegativity
  - 1. If the electronegative difference between two bonded atoms is equal or greater than 1.7 the bond is \_\_\_\_\_.
  - 2. If the electronegativity difference between two bonded atoms is less than 1.7 than the bond is polar \_\_\_\_\_\_.
  - 3. If the electronegativity difference between two bonded atoms is O than the bond is nonpolar covalent or metallic
- Polarity based on electronegativity

# Polar covalent bond:

e.g.) HCl-->\_\_\_\_\_

\*

Affected by: 1) electronegativity of the atoms; 2) bond dipoles & 3) shape *Polar molecule*:

electric poles:			 	 
bond dipole: _				
Step 1)			 	
Step 2)				 _
	•.1 1	1 /	 	(11)

POLAR: Elements with larger electronegativities become negative regions (delta ) & the elements with smaller electronegativities become positive regions (delta ).

NON POLAR: If the electronegativities are equal on the ends, than they cancel each other out. Step 3) \_\_\_\_\_

Non-polar covalent bond: \_\_\_\_\_\_.

e.g.) Cl<sub>2</sub> --> \_\_\_\_\_ CO<sub>2</sub> --> \_\_\_\_\_

CONCLUSION: Polarity is based on

D. Saturated vs. Unsaturated fats (optional)

	<b>EXAMPLES</b>	<b>EFFECT ON BODY</b>	<b>BONDS</b>
SATURATED			
POLYUNSATURATED			

#### **INTERMOLECULAR BONDING AND FORCES** Intermolecular bonding:

\*

\*

\*

- The size of the molecules and the molecular polarity determine the strength (weak)
- electron & nuclear attractions between molecules cause intermolecular bonding
- intermolecular forces affect \_\_\_\_\_\_ (Like substances (polar) dissolve like (polar) substances.)
- There are three main types:

  - form *temporary* dipoles. These are the \_\_\_\_\_\_ intermolecular forces.
- LD hold molecules together when they are in solid and liquid states
- \* LD are weaker than covalent bonds because electrons are farther from nuclei
  - If two molecules have the same # of electrons they are called ie) Ar = 18e & HCl = 18e BUT HCl has higher boiling point due to other forces.

\_\_\_\_\_ are an indirect measure of these forces.  $Br_2$  is a liquid &  $Cl_2$  is not because Br<sub>2</sub> has stronger dispersion forces. 2. \_\_\_\_\_: when electrostatic attraction between two polar molecules ie)

3. \_\_\_\_\_: special, relatively strong dipole-dipole forces between molecules containing O-H, F-H, & N-H bonds.

- These bonds form because of the \_\_\_\_\_\_ between hydrogen and \_\_\_\_\_ \* \* The small size of hydrogen atom means
  - and this positive pole exerts a strong attraction to the negative pole of a nearby molecule.
  - V shaped molecules with hydrogen bonds, like water, have stronger hydrogen bonds than other molecules, like ethanol because \_\_\_\_\_

ie) H & F ---- H & F (Hydrogen bonds are represented by dotted lines because \_\_\_\_

ie) water

ie) ethanol

\* PHYSICAL PROPERTIES Affected by Intermolecular bonds

#### 1.\_\_\_\_\_

ie) H<sub>2</sub>O is a liquid at room temperature and boils at 100 C because of extra strong hydrogen bonds
HCl (18 e) bp = -83.7 C
HBr (36 e) bp = -67.0 C
HI (54 e) bp = -35.4 C

2. \_\_\_\_

ie) water ---> ice - patterns form & it becomes less dense due to hydrogen bonds

### SUMMARY

\* Strength of all the bonds from strongest to weakest

network covalent & ionic & metallic(vary) & dispersion & hydrogen & dipole-dipole \* Limitations of bonding model

- 1. Structures of common gases such as NO2 & O3 cannot be explained easily
- 2. Special properties of gases like oxygen are difficult to explain
- 3. Formation of complex ions like nitrate, nitrite, carbonate, phosphate and sulphate are difficult to explain
- 4. Graphite and diamond are different structural forms of carbon. Reasons for the variety are difficult to explain
- \* Summary of two bond types

INTERMOLECULAR BONDS	INTRAMOLECULAR BONDS
Types: Van der Waals(London dispersion & dipole-dipole), Hydrogen	Types: Ionic, covalent, metallic, network covalent
Between molecules	Within molecules
Small energy change when broken	Large energy change when broken
Creates physical properties (melting point, boiling point, surface tension, solubility)	Creates chemical properties
Distance between charges is larger	Distance between charges is smaller
Polarity affects strength	Electronegativity affects type & structure
Generally weaker bonds	Generally stronger bonds

\* Electronegativity difference

Electronegativity difference	Type of bond	Description
> or = 1.7	Ionic	Transfer of electrons between metal & nonmetal
< 1.7	Polar covalent	Electrons shared between unlike nonmetal atoms
0	Nonpolar covalent	Electrons shared between like nonmetal atoms
0	Metallic	Electrons move freely between metal ions