

This review is intended to help review the majority of topics we have discussed in class. It is not 100% inclusive of all material that may be on the exam. Questions may be asked about topics that are not listed on this review sheet.

### Chapter 1 – Matter

Matter, properties of matter; physical vs. chemical changes, states: solids, liquids, gases  
Pure substances and mixtures; types of mixtures – heterogeneous vs. homogeneous,  
Types of pure substances: elements, compounds; Atoms and molecules  
Chemical symbols of the elements: know symbols and names of selected elements on periodic table  
Chemical formulas (with subscripts)

### Chapter 2 – Measurements

Metric system: meter, liter, gram  
Exact and inexact numbers, significant numbers, calculations involving significant figures  
Scientific notation (only significant figures are written)  
Dimensional analysis (unit factor method): include unit labels with numbers to help set up calculations  
Know by memory: 454g = 1pound, 3.78 liters = 1 gallon, 2.54 cm = 1 inch  
Density = mass/ volume (g/cm<sup>3</sup>); be able to calculate density given mass and volume.  
Temperature scales: °F, °C, K: convert from one to another, understand the relationships between scales  
Heat energy content is measured in calories; 1 calorie is the amount of energy to raise 1 g H<sub>2</sub>O by 1°C.

### Chapter 3 – Atomic Structure

Structure of the atom: nucleus, subatomic particles, protons, neutrons, electrons  
Atomic number and mass number; each atom has unique numbers that define it  
Isotopes: atoms with the same number of protons and electrons, but different numbers of neutrons  
Atomic mass: average atomic mass of an element, depends on the natural abundances of elements  
Metals vs. Nonmetals: Metals are located to the left of the staircase line on the periodic table, nonmetals to the right. Metals conduct heat and electricity, exhibit luster, and are malleable.  
Electrons are located on quantized energy levels around the nucleus called “orbitals” and have very little mass.  
We will use the Bohr model of the atom:  
    circular orbits having 8 electrons in the outermost orbital (except H, He that have only 2)  
    we will not discuss electron “shells,” “subshells,” or “configurations” (such as 1s<sup>2</sup>2s<sup>2</sup>2p<sup>6</sup>3s<sup>2</sup>... etc)  
Be able to draw Lewis dot structures with the appropriate numbers of valence electrons  
Understand the relationships between valence electrons and the representative groups in the periodic table  
Distinguish between groups and periods on the periodic table; identify the following elements on the table:  
    noble gases, representative elements, transition elements, alkali metals and alkaline earth metals  
Nuclear chemistry - fission: parent and daughter nuclei, general concept of E=mc<sup>2</sup>  
Nuclear stability and radioactivity: excessive neutrons contribute to instability  
Radioactive emissions: alpha ( $\frac{4}{2}\alpha$ ), beta ( ${}_{-1}^0\beta$ ), gamma ( $\gamma$ )  
    Describe each type of emission, its mass, velocity, penetration depth, damage to tissues, etc.  
    Be able to balance nuclear equations (e.g.,  ${}_{15}^{32}\text{P} \rightarrow \frac{4}{2}\alpha + \underline{\quad}$ )  
Half life: Understand the concept of half life for radioisotopes  
    Calculate how much of an isotope is remaining after a given number of half lives, or  
    calculate the half life, given the amount remaining after a measured period of time.

### Chapter 4 – Chemical Bonds

Understand the octet rule and the related role of valence electrons in determining bonding  
Ionic compounds: charged attraction between metals and nonmetals, determine ionic formulas & names  
Molecular compounds: covalent bonds between two or more nonmetal atoms, bonding & non-bonding pairs  
Covalent bonds can be single, double or triple bonds; predict bonding using valence electrons and octet rule  
Polarity of a covalent bond is determined by the electronegativity of the atoms involved in the bond.  
Predict molecular geometry (shape) of molecules from bonding and non-bonding pairs around central atom.  
Assign names to ionic and covalently-bonded compounds; e.g., NaCl (ionic) or CO<sub>2</sub> (covalent)  
Memorize 7 polyatomic ions and their charges: OH<sup>-</sup>, CN<sup>-</sup>, CO<sub>3</sub><sup>2-</sup>, SO<sub>4</sub><sup>2-</sup>, NO<sub>3</sub><sup>-</sup>, PO<sub>4</sub><sup>3-</sup>, NH<sub>4</sub><sup>+</sup>.  
Be able to write compound formulas involving polyatomic ions, using correct parentheses and subscripts.

### Chapter 5 – Chemical Calculations

Calculate formula masses and molecular weights using the periodic table.  
Understand the concept of the mole; know Avogadro’s number and what it means (6.023x10<sup>23</sup>)  
Using the mass of a mole, calculate the number of molecules or individual atoms in a given mass.  
Write and understand chemical formulas with subscripts and parentheses.  
Write and balance chemical equations.  
Calculate the grams and/or moles of reactants and products in chemical equations. (g → mole → mole → g)

**Chapter 6 – Gases, Liquids, and Solids**

Matter is in motion; motion of molecules increases with heat.

Absolute zero is the temperature at which all atomic and molecular motion ceases. (0K, or -273°C).

Combined gas law,  $P_1V_1/T_1 = P_2V_2/T_2$ , with calculation (Terms cancel from both sides when they are constant.)

Ideal Gas Law:  $PV=nRT$   $P$ =pressure(atm),  $V$ =volume(liters),  $n$ =moles,  $R=0.082$  (L·atm/K·mole),  $T$  (Kelvins).

Dalton's Law of Partial Pressures:  $P_t = p_1 + p_2 + p_3...$

Vapor pressure: evaporation, boiling, explain why boiling point of water changes with altitude

Intermolecular forces help determine boiling points: Define hydrogen bonding.

**Chapter 7 – Solutions**

Solute, solvent, alloys, liquids, solids, gasses as solutes. Explain the phrase, "Like dissolves Like."

Solution formation: dissolution of ionic compounds to form solvated ions.

Solubility rules for ionic compounds: e.g., which is more soluble,  $\text{CaCO}_3$  or  $\text{Na}_2\text{CO}_3$ ?

Solute concentration units: %(w/w), %(w/v), %(v/v), molarity ( $M$ =moles/L), normality ( $N$ =equivalents/L)

Dilution calculations:  $C_1V_1=C_2V_2$  ← Use this equation... given 3 values, calculate the 4<sup>th</sup>.

Colligative Properties: freezing (melting) point depression (e.g. antifreeze).

Osmosis and Osmotic Pressure: solvent (water) passes through a membrane into liquids with less water.

Osmotic pressure draws water into plants, through the digestive tract, etc.

Osmotic pressure is determined by TOTAL dissolved particles: 1mole NaCl yields 2 total moles of ions (both  $\text{Na}^+$  &  $\text{Cl}^-$ )

Define and know the meanings of the terms: Isotonic, hypertonic and hypotonic solutions.

Dialysis

**Chapter 8 – Chemical Reactions**

Combination reactions  $X+Y \rightarrow XY$ ; Decomposition reactions  $XY \rightarrow X+Y$ ; Single & Double displacement reactions

Oxidation and reduction reactions ("Redox" reactions): Oxidation = loss of electrons; Reduction = gain of electrons

(Definition on page #205 is not correct!); Examples: Oxidation:  $\text{Cu} \rightarrow \text{Cu}^{2+} + 2e^-$  Reduction:  $\text{Ag}^+ + 1e^- \rightarrow \text{Ag}$

Formation of positively-charged metal ions from their elements is oxidation.

Reactants must collide with each other to react. Reaction rates depend on frequency, orientation, & energy of collisions.

Rates of chemical reactions depend on reactant concentrations, temperatures, and presence of catalysts.

Chemical Equilibrium: LeChatelier's principle: equilibrium shifts to relieve stress... temperature, reactants, pressure.

**Chapter 9 – Acids, Bases, and Salts**

Definition of acids and bases: acids donate  $\text{H}^+$ , bases increase  $\text{OH}^-$  ions in water

Names (formal & common) and formulas of common acids: HCl,  $\text{HNO}_3$ ,  $\text{H}_2\text{CO}_3$ ,  $\text{H}_2\text{SO}_4$ ,  $\text{H}_3\text{PO}_4$ .

Identify the economically important acids and bases and their uses.

Names and formulas of common bases: NaOH, KOH,  $\text{Ca}(\text{OH})_2$ ,  $\text{Mg}(\text{OH})_2$ ,  $\text{NH}_4\text{OH}$

Reactions of acids with metals (e.g., copper); know their reaction products (metal salt plus  $\text{H}_2$  gas)

Identify elements oxidized and reduced during a reaction with acid and metals

Neutralization reactions between acids and bases: acid + base  $\rightarrow$  salt + water; give examples, write eqns.

Self-ionization of water:  $\text{H}_2\text{O} \leftrightarrow \text{H}^+ + \text{OH}^-$

pH definition:  $-\log[\text{hydrogen ion concentration in moles per liter}]$ ; calculate pH using calculator

Convert pH to  $\text{H}^+$  concentration

Understand the pH scale: powers of 10, acid vs. basic pH

Buffers resist pH changes when acids or bases are added to the solution

Bicarbonate buffer system:  $\text{CO}_2 + \text{H}_2\text{O} \leftrightarrow \text{H}_2\text{CO}_3 \leftrightarrow \text{H}^+ + \text{HCO}_3^- \leftrightarrow \text{H}^+ + \text{CO}_3^{2-}$

**Chapter 10 – Saturated Hydrocarbons**

Carbon atoms bond to other carbons and hydrogen atoms to form hydrocarbons; valence of carbon, 4 bonds

Alkanes are hydrocarbons with single bonds between carbon atoms; non-cyclic:  $\text{C}_n\text{H}_{2n+2}$ , cyclic alkanes:  $\text{C}_n\text{H}_{2n}$

Structural formulas and structural isomerism; different conformations are due to rotation of single bonds

Names of alkanes:  $\text{C}_1$ - $\text{C}_{10}$ ; methane, ethane, propane, ..., decane

Cyclic alkanes can exhibit *cis*- and *trans*- isomerism

Assign proper IUPAC names of branched and/or cyclic alkanes

Hydrocarbons come from petroleum; distillation, cracking and reformation give us a wide variety of products

Physical properties of hydrocarbons: gases, liquids, solubilities

Chemical reactions of alkanes: combustion, halogenations (substitution of F, Cl, Br, or I for H)

Key reactions: Alkane +  $\text{O}_2 \rightarrow \text{CO}_2 + \text{H}_2\text{O}$ , and  $\text{R-H} + \text{X}_2 \rightarrow \text{R-X} + \text{H-X}$  (R=alkane, X=halogen such as Cl, Br)

**Chapter 11 – Unsaturated Hydrocarbons**

Unsaturated hydrocarbons contain one or more C=C or C≡C bonds

General formula for non-cyclic alkenes: C<sub>n</sub>H<sub>2n</sub>, alkynes: C<sub>n</sub>H<sub>2n-2</sub>

Names of alkenes: C<sub>2</sub>-C<sub>10</sub>; ethene, propene, ..., decene

Alkenes can exhibit *cis*- and *trans*- isomers

Names of alkynes: C<sub>2</sub>-C<sub>10</sub>; ethyne, propyne, ..., decyne

Assign IUPAC names for branched and/or cyclic alkenes and alkynes

Chemical reactions of alkenes:

Addition reactions: H<sub>2</sub>, Cl<sub>2</sub>, Br<sub>2</sub>, HCl, HBr, H<sub>2</sub>O all add to double bonds; (Markovnikov's rule)

Polymerization of alkenes: small alkenes combine to form long chains of polymers

Know polymerization reactions and recognize names of important polymers in Table 11.2

Aromatic hydrocarbons: Rings with delocalized multiple double bonds; most common is benzene, C<sub>6</sub>H<sub>6</sub>

Name substituted benzene rings with proper numbering systems; also *ortho*-, *meta*-, and *para*- prefixes

Know the structures of toluene, xylenes, naphthalene, anthracene

Phenyl groups are benzene rings attached to carbon atoms as substituents on longer chains.

Reactions of benzene: alkylation and halogenation

Know reactions of alkenes and alkynes (pp. 304-305)

**Chapter 12 – Single-bond Derivatives**

Halogenated hydrocarbons: R-X (where X is F, Cl, Br, I)

Names start with number- and fluoro-, chloro-, bromo-, or iodo-: e.g., 2-chlorobutane

Common names for some solvents: e.g., carbon tetrachloride: CCl<sub>4</sub> or ethyl bromide

Alcohols & Phenols: R-OH

IUPAC (one-word) names: change alkane name by dropping “-e” and adding “-ol.”: ethanol, methanol

Common names for common solvents: ethyl alcohol, methyl alcohol, isopropyl alcohol. (pg312)

Ethanol concentrations are often expressed using the “proof” scale, where 100% = “200 Proof”

Polyhydroxy alcohols: 1,2-ethanediol (ethylene glycol): HO-C-C-OH, propylene glycol, glycerin (p313)

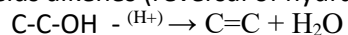
Phenols names: IUPAC: phenol, 4-methylphenol; common name: phenyl alcohol

Properties of alcohols: boiling points go up with number of carbons and alcohol groups (p316-7)

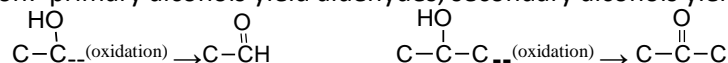
Alcohols form strong hydrogen bonds to themselves and water

Reactions of alcohols:

dehydration yields alkenes (reversal of hydration)



oxidation: primary alcohols yield aldehydes, secondary alcohols yield ketones



Ethers: R-O-R

Names: ethyl methyl ether: C-O-C-C, dimethyl ether: C-O-C (p322)

Properties: flammable, anesthetic activities

Thiols: R-SH (*sulfhydryl group*)

Recognize thiol functional group(s)

Thiols have strong, disagreeable odors; e.g., skunk odor, added to natural gas for safety

Oxidation of thiols yields disulfide bonds: C-C-SH + HS-C-C → C-C-S-S-C-C decreases odor;

(dilute H<sub>2</sub>O<sub>2</sub> is an excellent oxidizing agent)

Amines: R-NH<sub>2</sub> (R-NH-R, or R-N-R<sub>2</sub>)

Names: ethyl amine: C-C-NH<sub>2</sub>, ethylmethylamine: C-C-NH-C, *N,N*-dimethylpropanamine:  $\text{H}_3\text{C}-\text{CH}_2-\text{CH}_2-\overset{\text{CH}_3}{\underset{|}{\text{N}}}-\text{CH}_3$

Dopamine, epinephrine, and ephedrine are all biologically active amines (p330)

Low molecular weight amines have a “fishy” smell

Amines can be protonated with acid to form amine salts:



Heterocyclic amines have ring systems where N has been substituted for at least one C in the ring.

Caffeine, nicotine morphine, codeine, heroin are all examples of heterocyclic amines (p334)

Review key reactions and equations on pg 336

**Chapter 13 – Carbon-oxygen Double Bonds**

Carbonyl group: C=O

Aldehydes: -CHO; names end in “-al”:



methanal (formaldehyde), ethanol (acetaldehyde), propanal, benzaldehyde, etc.

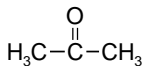
usually have peculiar odors (almonds, cinnamon, vanilla)

reactions: reduction to alcohols: C=O + H<sub>2</sub> → CH-OH

oxidation to acids: C=O → -COOH

during oxidation, aldehydes can reduce metal ions: Cu<sup>2+</sup> to Cu<sup>1+</sup> (Benedict’s test)or Ag<sup>1+</sup> to Ag<sup>0</sup> (Tollen’s test - mirror forms)

Ketones: names end in “-one”:



2-propanone (acetone), 2-butanone (ethylmethylketone), 3-pentanone (diethylketone), etc.

Reactions: reduction to alcohols C=O + H<sub>2</sub> → CH-OH

oxidation does not occur!

Excellent solvents used industrially e.g., finger-nail polish remover, varnish remover, etc.

Carboxylic acids: -COOH, names end in “-ic acid”:

Formic acid (from ants), acetic acid (vinegar), propanoic acid, butanoic acid, benzoic acid, etc.

Acids release protons in aqueous solutions: -COOH → -COO<sup>-</sup> + H<sup>+</sup>

Some acids have more than one -COOH group: succinic acid(2), glutaric acid(2), citric acid(3)

Three common OTC pain relievers are carboxylic acids: Ibuprofen and Naproxen and aspirin

Carboxylate salts form when acids are neutralized with base: -COOH + NaOH → -COO<sup>-</sup> Na<sup>+</sup> + H<sub>2</sub>O

Carboxylate salts are principle ingredient in soaps

Small molecular weight carboxylic acids are preservatives: benzoic acid, sorbic acid, propionic acid

Carboxylic acid reactions (condensations):

with alcohols to form esters: R-COOH + HO-R' → R-COO-R' + H<sub>2</sub>Owith amines to form amides: R-COOH + H<sub>2</sub>N-R' → R-CONH-R' + H<sub>2</sub>O

Esters: -COOR, names end in “-ate.”

Methylformate, methylacetate, propylbutanolate, ethylbenzoate, etc.

Recognize the acid and alcohol building blocks for esters

Esters have characteristic flavors and odors (“fruit-like” smells)

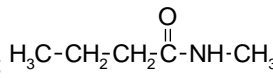
Aspirin is an ester of acetic acid and the alcohol group on salicylic acid, “acetylsalicylic acid”

Esters are easily hydrolyzed, yielding the original acids and alcohols.

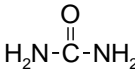
Amides: -CONH-R, names end in “-amide.”

Formamide, acetamide, benzamide, etc.

IUPAC names include “N-”: N-methylpropanamide:



Urea, diamide, is an important amide biologically:



The pain reliever, acetaminophen, is an amide of acetic acid (pg371)

Reactions of amides: hydrolysis yields carboxylic acids and amines

Condensation Polymers: polyesters and polyamides

Polyesters are synthesized from diacids and dialcohols: PET: Dacron, Mylar, soft-drink bottles, etc.

Polyamides are synthesized from diacids and diamines: nylons, Kevlar, etc.

**Chapter 14 - Carbohydrates**General: C<sub>n</sub>H<sub>2n</sub>O<sub>n</sub>; Subclasses: pentoses(C<sub>5</sub>), hexoses(C<sub>6</sub>); aldoses (aldehydes) and ketoses (ketones)

Handedness (“D” &amp; “L”) forms of monosaccharides; rotation of plane-polarized light right or left

Important monosaccharides:

Glucose (dextrose, blood sugar) is a hexose (C<sub>6</sub>) and an aldose, (an aldohexose), is a reducing sugar, and rotates light to the *right* in aqueous solution; it is an important nutrient in the blood stream.Fructose (levulose) is a ketohexose (hexose and ketose); it is the sweetest tasting sugar; it rotates light to the left, and is an important sweetener for soft drinks and other foods;Ribose is an aldopentose (aldose and pentose); 5-carbon sugar, important building block in DNA and RNA.

Monosaccharides exist primarily as cyclic forms in solution

Most monosaccharides are reducing sugars; that is, they oxidize easily and concomitantly reduce other chemicals such as Cu<sup>2+</sup> ions (Benedict’s test).

**Disaccharides:**

Maltose: glucose-glucose	reducing sugar; common in foods (malt flavoring, beer)
Lactose: galactose-glucose	reducing sugar; “milk sugar”
Sucrose: glucose-fructose	non-reducing sugar; “table sugar,” “cane and beet sugar”

**Polysaccharides:**

Cellulose: polyglucose, linked with $\beta(1\rightarrow4)$ bonds, not digestible, structural component of plants
Starch: polyglucose, linked with $\alpha(1\rightarrow4)$ bonds, complex carbohydrate, good glucose source
Glycogen: branched polyglucose, linked with $\alpha(1\rightarrow4)$ bonds; very soluble, “animal starch”
Chitin: linear polymer with $\beta(1\rightarrow4)$ bonds, contains acetylamino groups on each glucose monomer

**Chapter 15 – Lipids**

Lipids are fats and oils that are extractable into non-polar organic solvents such as hexane

Fatty acids are the principle components of fats and oils; fatty acids have 12-20 carbon reduced atoms (contain mostly hydrogen atoms);

Saturated and unsaturated fatty acids: (*cis*- double bonds in fatty acid chains)

Essential fatty acids: linoleic and linolenic acid, omega-3 family of fatty acids,

Triacylglycerols are tri-esters of 3 fatty acids and glycerol

Unsaturated nature of fatty acids lowers the melting point and as such, “oils” are liquids at room temp.

Reactions: hydrolysis / saponification: yields free fatty acids and glycerol; components of soaps  
 Hydrogenation: adding hydrogen across C=C double bonds to create more saturated FAs  
 Oxidation: oxygen reacts with C=C double bonds to create small aldehydes and acids.

Phospholipids: Tri-esters of glycerol with 2 fatty acids and one phosphoric acid group. (“lecithin”)

Principle components of biological membranes

Sphingolipids: fatty-acid linked to sphingosine via an amide linkage; present in nerve tissues.

Steroids: Lipids that contain a characteristic fused 4-ring system (pg 423)

Cholesterol – the most abundant human steroid; alcohol; building block for hormones

Bile salts: emulsifying agents to help solubilize dietary lipids; stored in the gall bladder

Steroid hormones: estrogens (estradiol), androgens (testosterone), progestins

Cell Membranes are composed of lipids; phospholipids are most common lipids; membranes also contain significant amounts of protein and some carbohydrates and cholesterol.

**Chapter 16 – Proteins**

Amino acids are the building blocks for proteins

AA properties are determined by their side-chains: e.g., polar, non-polar, aromatic, acids, bases, etc.

Two AAs contain sulfur; cystine, forms disulfide linkages that are easily oxidized and reduced.

AAs are classified as either “essential” or “non-essential”

Phenylalanine can cause developmental problems for PKU babies

Tryptophan is a precursor for serotonin

AAs are linked together with amide (peptide) bonds to form long polymers called proteins

Peptide bonds may be hydrolyzed in acid, yielding free amino acids and smaller peptide chains

There are four levels of protein structure:

Primary: linear sequence of AAs

Secondary: alpha-helix and beta-sheet

Tertiary: Overall 3-D shape of protein

Quaternary: Interactions between separate subunits of proteins

Enzymes are proteins that act as catalysts

Enzymes are unique catalysts that are highly specific and are regulated by conditions in the cells

Enzymes react with “substrates” and help speed up their conversion to “products”

Enzyme turnover numbers describe how fast substrates can be converted to products.

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