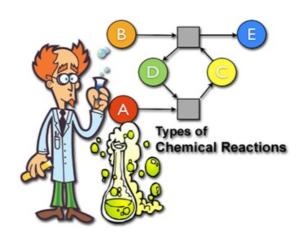
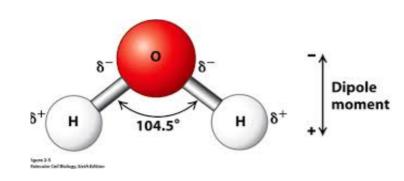
Ch 4 Reactions and Solution Chemistry:

- Solutions concentrations and types
- Chemical Reactions
 - Precipitate Rxns
 - Acid/Base Rxns
 - Other types of Rxns



H₂O - The Common Solvent:



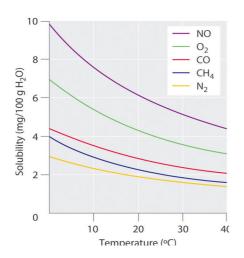
- Type of bonds?
- Types of compounds it can dissolve?
- How does it dissolve or dissociate?

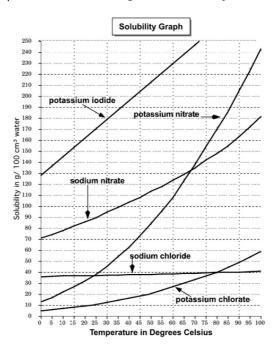
Solubility:

- Quantitative amount (measured in X g solute/100 g $\rm H_2O$) of solute that will dissolve in solvent
 - Solute:
 - Solvent:

Trends in solubility:

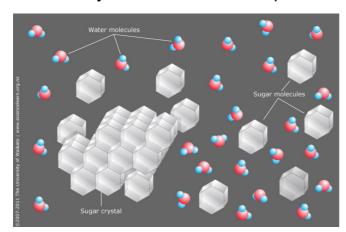
- Solids:
 - Generally an \int in temp = \int in amt. of solute soluble in 100 g H₂O
- Gases:
 - Opposite is true: ↑ temp = ↓ in amount of gas solubility in solvent

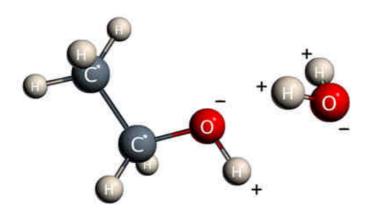




H₂O also dissolves non-ionic substances:

- Breaks them into molecules by attraction to waters S- and S+ sides
 - this applies to molecular substances like sugar and alcohols
- Does not Dissociate them into ions or atoms
 - Only breaks a mass clump of molecules into individual molecules





Strong vs Weak Electrolytes:

Strong electrolyte - 100% of compound ionizes when placed in water

Ex: Strong acids, strong bases, soluble salts

Weak electrolyte - a small degree of ionization occurs when placed in water

Ex: Weak acids, weak bases, insoluble salts

 \Longrightarrow

Non-electrolyte - no dissociation into ions occurs, but compound dissolves into individual molecules

Ex: Sugars, alcohols

Writing dissociation equations for compounds:

- Decide if compound is an acid, base, or salt
- Decide if it is strong or weak (this tells you what arrow to use)
- Dissociate the compound, balancing each side. Show the ions as charged
- If it is a polyprotic acid, be sure to dissociate the H+ in steps

D	iss	ociati	on r	evie	w/p	racti	ce:
	.00	Joiati	0	0110	*** P	uoti	.

 $Ca(OH)_2$

 $Mg(NO_3)_2$

Oxalic acid

Determining the concentration of solutions:

Molarity =
$$\frac{\text{moles solute}}{\text{Liters of solution}}$$

% concentration =
$$g$$
 solute * 100 g total solution



molality =
$$\frac{\text{moles solute}}{\text{kg solvent}}$$

Dilution equation: $M_1V_1=M_2V_2$

1 = initial conditions

2 = final conditions

Difference in V_1 to V_2 is due to ?

Moles of solute stays CONSTANT between change of volumes!



Practice with concentration math:

#1: Determine Molarity if 11.5 g of NaOH is used to make a 1.50 L solution.

#2: If a solution of NaCl has a density of 1.25 g/mL and a % by mass concentration of 16.0%, Determine its Molarity.

#3: You need to make a 0.10 M solution of H_2SO_4 . How much stock solution should you use to make 2.00 L of you dilute solution if the stock solution is 16 M.

#4: Typical blood serum is 0.14 M NaCl. What volume of blood contains 1.0 mg of NaCl.

More concentration practice:

Introduction of ICE box method

Ex 1: Calculate the molarity of each ion present in solution if you dissolve 8.00 g of $Fe(CIO_4)_3$ into 1.50 L of total solution.

Ex: Calculate the moles of Na^{+1} ions present in a solution of 1.25 M Na_2SO_4 solution.

Reactions that involve solutions:

- Metathesis (double replacement)
 - liquid, gas or precipitate = products possible
- Acid-Base (neutralization)
 - salt and H₂O are your products
- Reduction oxidation (redox)
 - involves a transfer of e- and reduction of 1 compound, oxidation of other

Metathesis reactions:

- pretend cations switch places, thus showing possible products
- think of solubility rules and ID states of products (s) for insoluble or slightly soluble compoiunds
- ppt is indicated with]
- other possible products are:

NH₃ and H₂O (due to NH₄OH being formed) CO₂ and H₂O (due to H₂CO₃ being formed)





Predict the following: Draw the ions/compounds in beakers as the reaction progresses.

Ex: Potassium nitrate + Barium chloride

Ex: Sodium sulfate + lead (II) nitrate

Ex: Potassium hydroxide + Iron (III) nitrate

Net ionic equations:

Use of solubility rules and understanding of strong vs weak electrolytes to show what is happening in a reaction

Compounds that totally ionize (------):

Compounds that have little/partial ionization (→):

Rules of follow in net ionic equations:

- 1.) Only dissociate the compounds that totally ionize, leave others together
- 2.) Show charges of ions and elements
 - -what is the charge on an element?
- 3.) Balance all ions/compounds in the equation
- 4.) Determine the net ionic equation and spectator ions after completing the overall ionic equation

Practice with net ionic equations:

Ex 3: Na +
$$H_2O$$
 -----> NaO $H_{(aq)}$ + H_2

Ex 4: Potassium hydroxide solution is poured into acetic acid

Stoichiometry of solutions:

- Focus on who the product is
- Need moles, use Molarity
- Use stoichiometry rules we know may need to use ICE box if asked about concentrations of ions also

Ex: Find the mass of NaCl needed to ppt out all Ag+ ion from AgNO₃ with a molarity of 0.100 M and a volume of 1.50 Liters

Ex 2: Aqueous solutions of sodium sulfate and lead (II) nitrate are mixed. Calculate the mass of product formed from 1.25 L of 0.0500 M lead (II) nitrate and 2.00 L of 0.0250 M sodium sulfate.

Acid - Base Reactions:

- Arrhenius Theory:

Acids - produce H+ ions

Bases - produce OH- ions

- Bronsted - Lowry Theory:

Acids - H+ donor

Bases - H+ acceptor

Neutralization reaction - acid + base ---> salt + H₂O

Practice stoichiometry of acid-base reactions:

Steps for stoichiometry of acid base reactions:

- 1.) Write the reaction, balance, and determine the net ionic equation Special note about net ionic equations of acid/base: If a weak acid is reacting with a strong base, a 100% reaction occurs even though the acid was weak
- 2.) Stoichiometry math
- 3.) Convert grams, volume, molarity if needed

Ex: What volume of 0.100 M HCl is needed to neutralize 25.0 mL if 0.350 M Ca(OH)₂?

Ex 2: In an experiment, 28.0 mL of 0.250 M HNO₃ and 53.0 mL of 0.320 M KOH are mixed. What concentration of H+ and OH- ions are in excess after the reaction?

Titrations of acids and bases:

Titration- delivering a measurable volume of a known concentration solution to an unknown concentration solution

Equivalence point - pt in titration when mole of titrant has 100% reacted with analyte

Endpoint - pt in titration when indicators color changes

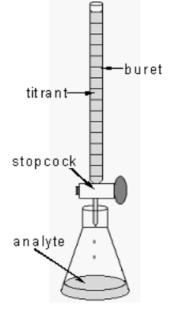
-(this ideally should coincide with the equivalence point - if you chose the correct indicator)

Titrant - solution in buret

- measure amount you use at end

Analyte - solution in flask

- measured the amount at the beginning



Steps to follow as you do a titration and its math:

- 1.) Know the balanced reaction of titrant + analyte
- 2.) Pick proper indicator
- 3.) Careful, slow.... stop as soon as that 1 single drop changes the color permanently
- 4.) Know the exact moles or volume of analyte you used

Ex: 1.3009 g of KHP (KHC₈H₄O₄) is dissolved in approximately 50.0 mL of H₂O. The titration used 41.20 mL of the titrant (NaOH). Determine the M of NaOH.

Ex: A chemist analyzed effluent water from an industrial process known to contain CCl_4 and benzoic acid $(HC_7H_5O_2)$, a weak acid that has one acid proton per molecule. A sample of this effluent weighing 0.3518 g was shaken with water, and the resulting aqueous solution required 10.59 mL of 0.1546 M NaOH to neutralize. Calculate the mass percent of $HC_7H_5O_2$ in the original sample.

Oxidation-Reduction reactions:

- -Reactions where 1 or more e- are transferred
- -Covalent and ionic compounds can undergo Redox reaction
- How to know if REDOX rxn occurred = if the oxidation #s of atoms changed

How to determine oxidation # of atoms:

- 1.) An unbonded atom = 0
- 2.) Monatomic ions = its normal charge due to its group
- 3.) If atoms are in a polyatomic ion or compound:
 - a.) Fluorine present = it has priority = it is -1
 - b.) Oxygen is normally -2
 - c.) Hydrogen = +1
 - d.) The summed charges of the atoms needs to = total charge of ion or compound

Practice:

Find the oxidation #s on each of the atoms:

 CO_2 SF_6 NO_3^{-1}

Fractional oxidation charges may occur... that is ok (WE are the ones that assign e- a home)

ex: Fe₃O₄

Details about Redox reactions:

Electrons are transferred to:

- 1.) form ions
- 2.) rearrange atoms

Oxidation = increase in oxidation state (become more +)

Reduction = decrease in oxidation state (become more -)

OIL RIG

Oxidizing agent (OA) - electron acceptor (gets reduced)

Reducing agent (RA) - electron donor (gets oxidized)

Examples: ID who is oxidized, who is reduced, and ID OA and RA

$$CH_4 + 2O_2 ---> CO_2 + 2H_2O$$

The following reactions take place as chemists take galena (PbS), a lead conaining ore, to its elemental state.

Balancing Redox Reactions:

(FOR EASIER REDOX REACTIONS)

Steps:

- 1.) Assign oxidation #s to all atoms
- 2.) e- lost must = e- gained

 Show the e- lost and gained with tie lines
- 3.) Use coefficients to = e- lost/gained
- 4.) Add states (s, I, g, aq)
- 5.) Balance the rest of the equation

Example:

$$Cu_{(s)}$$
 + $Ag^{+}_{(aq)}$ ----> $Ag_{(s)}$ + $Cu^{+2}_{(aq)}$

$$H^+ + Cl^- + Sn + NO_3^{-1} ----> SnCl^{-2}_6 + NO_2 + H_2O$$

PbO +
$$NH_3$$
 ----> N_2 + H_2O + Pb

Gravimetric analysis:

- Use of precipitation reaction to measure product made
- Through stoichiometry determines amount of a certain ion in solution

Example:

An ore sample is to be analyzed for sulfur. As part of the procedure, the ore is dissolved, and the sulfur is converted to sulfate ion (SO_4^{-2}) . Barium nitrate is added which causes the sulfate to precipitate out. The original sample had a mass of 3.187 g. The dried product is 2.005 g. What is the percent of original Sulfur in the ore?

Example: The thallium (present in Tl_2SO_4) in a 9.486 g pesticide sample was precipitated as thallium (I) iodide. Calculate the mass percent of Tl_2SO_4 in the sample if 0.1824 g of thallium (I) iodide was recovered.

Determination of Concentration of ions in solution:

- A) Determine the amount of moles of each reactant ion (Molarity --> moles and ICEBOX)
- B) Use stoichiometry to calculate Limiting and excess reactant
- C) Complete LR to ER stoich to find the moles of ER needed
 - Then subtract from given amount of ER ion to find moles of ER left over
- D) After you know the moles of each ion, use molarity formula to find [] Conc.

Ex: Determine the molarity of Fe⁺³ ions and SO₄⁻² ions in a solution prepared by dissolving 48.05 g of Iron (III) sulfate in enough water to make a total volume solution of 800. mL.

Ex: You react 35.0 mL of 2.00 M Sodium carbonate solution with 25.0 mL of 1.50M Calcium chloride solution. Determine the mass of product expected. Then find the concentration of all ions in the filtrate solution.

REDOX practice:

Determine the oxidation #s, who is reduced, who is oxidized, OA and RA for the following problems. Then balance and ID # e- transferred and if they are gained or lost.

Ex:

$$H_2$$
 + O_2 ---> H_2O

$$Zn_{(s)}$$
 + Cu^{+2} ---> Zn^{+2} + $Cu_{(s)}$

$$2MnO_{4^{-}(aq)} \quad + \quad 16H^{+}_{(aq)} \quad + \quad 5C_{2}O_{4}^{2^{-}}_{(aq)} \quad ---> \\ \\ 2Mn^{2+}_{(aq)} \quad + \quad 10CO_{2} \quad + \quad 8H_{2}O$$

Single Replacement Reaction Review:

Element + Compound (aq) ---> Element + Compound (aq)

Cation replaces cation Anion replaces anion

<u>Activity series -</u> stronger element can replace weaker element

Ex: Ca + $Fe(NO_3)_3$

Ex: Chlorine gas is bubbled into a solution of hydrobromic acid

Ex: Copper is added to a solution of zinc chloride.