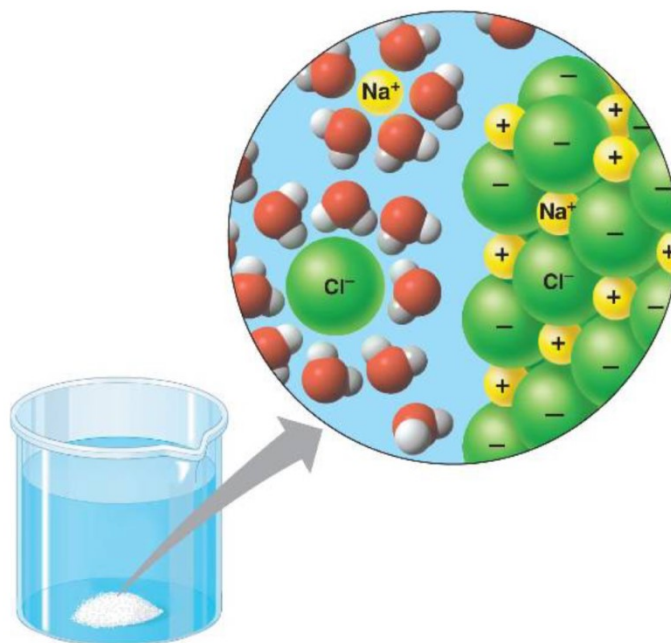


# Note Packet

## UNIT 7: SOLUTIONS



### Unit Vocabulary:

Alloy	heterogeneous	saturated
Aqueous	homogeneous	solubility
boiling point	insoluble	soluble
boiling point elevation	miscible	solution
colligative properties	mixture	solute
colloid	molarity (M)	solvent
concentrated	parts per million (ppm)	supersaturated
concentration	percent by mass	suspension
dilute	percent by volume	Tyndall Effect
freezing point depression	precipitate	unsaturated

**Unit Objectives** (Upon the Completion of this unit you will be able to):

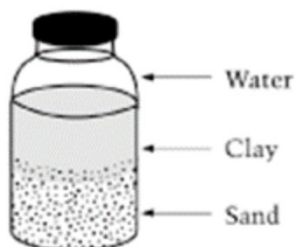
- ◆ Differentiate between heterogeneous and homogeneous mixtures
- ◆ Identify the various types of heterogeneous mixtures and their properties
  - ▶ colloid
  - ▶ suspension
- ◆ Identify the various types of homogeneous mixtures and their properties
  - ▶ alloy
  - ▶ solution
- ◆ Define solubility and understand the factors that contribute to solubility
  - ▶ nature of solute and solvent (like dissolves like)
  - ▶ temperature
  - ▶ Use Table F to determine if precipitate is formed in a chemical reaction
- ◆ Distinguish between saturated, unsaturated, or supersaturated solutions
- ◆ Read the solubility curve (Table G) to determine if a solution is (1) saturated, (2) unsaturated, or (3) supersaturated
- ◆ Differentiate between dilute and concentrated solutions
- ◆ Calculate various concentrations of a solution using the following:
  - ▶ Molarity (M)
  - ▶ Percent by Mass
  - ▶ Percent by Volume
  - ▶ Parts per Million (ppm)
- ◆ Prepare a solution of known concentration
- ◆ Explain a solute's effect on a solution (colligative properties)
  - ▶ Freezing Point Depression
  - ▶ Boiling Point Elevation

**A mixture is:** a combination of 2+ pure substances (elements and/or compounds) that can be separated by **PHYSICAL** means

**Mixtures** can be categorized into one of two groups:

**1. HETEROGENEOUS MIXTURE:** mixture of 2 or more pure substances in **DIFFERENT PHYSICAL STATES**; **VARYING COMPOSITION** (**not** uniform throughout)

➤ **Suspension:** heterogeneous mixture that **clearly** separates



- eventually **SEPARATES** upon standing
- solid does not visibly dissolve in the liquid (or break down)
- separate by **FILTRATION** (using a filter)
- relatively **LARGE** particle size
- Examples: **SAND IN WATER, ITALIAN DRESSING**

➤ **Colloid:** heterogeneous mixture that does **NOT CLEARLY SEPARATE**



- **CLOUDY** (*appear* homogeneous, but are **NOT**)
- do not settle (naturally; on their own) through time
- can be separated by a **CENTRIFUGE** (forced gravity)
- **MEDIUM** to **SMALL** particle size
- properties in between a suspension and a solution
- Examples: **BLOOD, MILK**

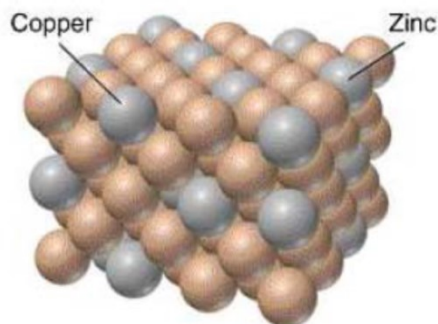


**2. HOMOGENEOUS MIXTURE (AKA Solution):** mixture of 2 or more pure substances in the **SAME PHYSICAL STATE**; **CONSTANT COMPOSITION** (uniform throughout)

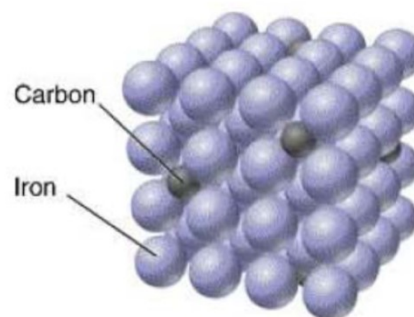


- a. looks like **CLEAR, COLORED** or **COLORLESS** water
- b. **EXTREMELY SMALL** particle size (you can't see them)
- c. may exist in any phase (liquid phase most common)
- d. Examples: **SALTWATER, KOOL-AID, CLUB SODA**

➤ **Alloy:** solution where at least one of the materials is a **METAL** (although most times it is a mixture of 2+ metals)



**A** Brass, a substitutional alloy



**B** Carbon steel, an interstitial alloy

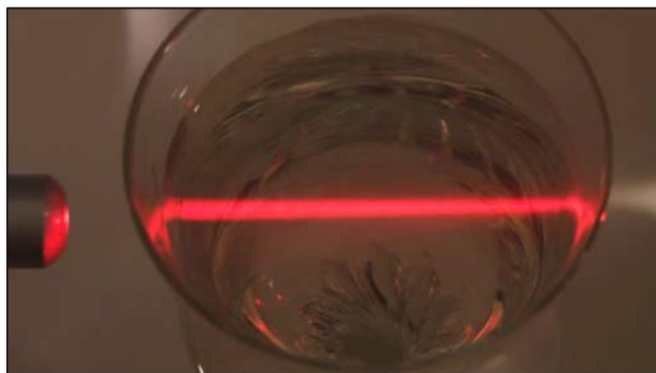
- Metals arrange in an alternating pattern with **SEA OF MOBILE VALENCE ELECTRONS** gluing them together
- Examples: **BRASS** (zinc + copper), **STEEL** (iron + carbon)

**Now, we will focus on SOLUTIONS only...**

**Characteristics  
of  
SOLUTIONS**

1. **HOMOGENOUS** mixtures
2. **CLEAR** (as opposed to cloudy) and do not disperse light (no Tyndall observed)
3. Can have **COLOR** or be **COLORLESS**
4. **DO NOT SETTLE** upon standing (after any amount of time)
5. **CANNOT** be separated by filtration

**TYNDALL EFFECT:** **NOT** seen in solutions; seen when a mixture makes a beam of light **VISIBLE**



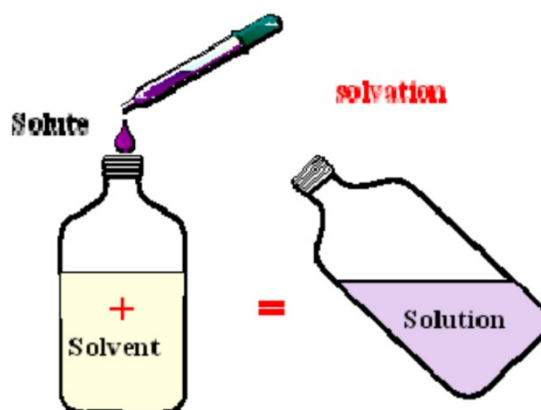
**(NOT a solution!)**

**\*\*Solutions always assume the phase of the solvent!**

Phase of Solute	Phase of Solvent	Example
Solid	Solid	Metallic alloy
Solid	Liquid	Table salt & water
Liquid	Liquid	Ethyl alcohol & water
Gas	Liquid	CO <sub>2</sub> & water (club soda)
Gas	Gas	Air

## SOLUTIONS ARE MADE UP OF 2 PARTS:

- ✓ **SOLUTE**: substance or substances that are **DISSOLVED** in a solution (example: the salt in saltwater); solute is usually a **SOLID** and is the **LESSER** quantity of the two (versus the solvent)
- ✓ **SOLVENT**: the substance that is **DISSOLVING** the solute (example: the water in salt water); solvent is usually a **LIQUID** and is the **GREATER** quantity of the two (versus the solute)

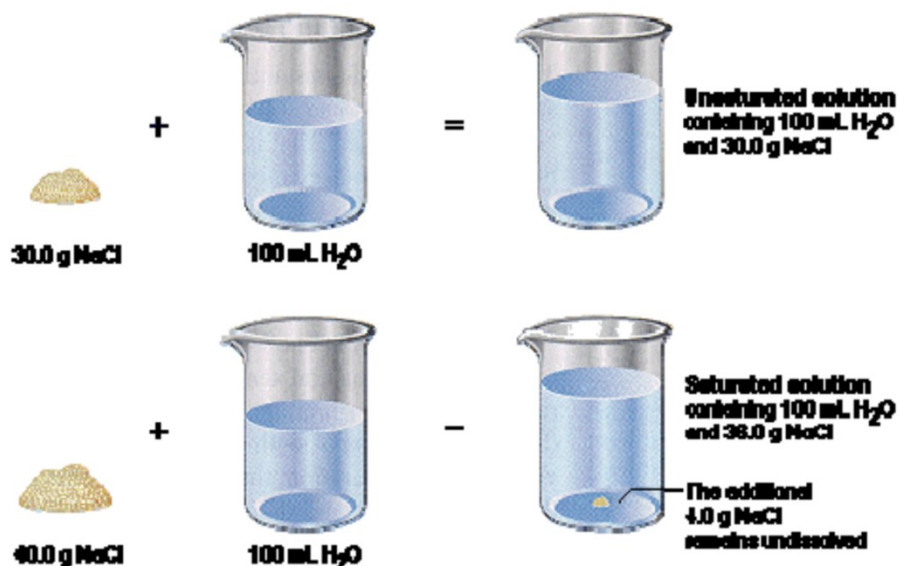


**AQUEOUS** = a solution in which **WATER** is the solvent (aq)

Example: NaCl(aq) is otherwise known as **SALT WATER**, is actually **Na<sup>+</sup>(aq) + Cl<sup>-</sup>(aq)**

## DIFFERENT TYPES OF SOLUTIONS:

- ✓ **UNSATURATED:** a soln in which **MORE SOLUTE** could be dissolved in a given volume of solvent
- ✓ **SATURATED:** a soln containing the **MAXIMUM AMT.** of solute that will dissolve in a given volume of solvent
  - Saturated = solution **EQUILIBRIUM**

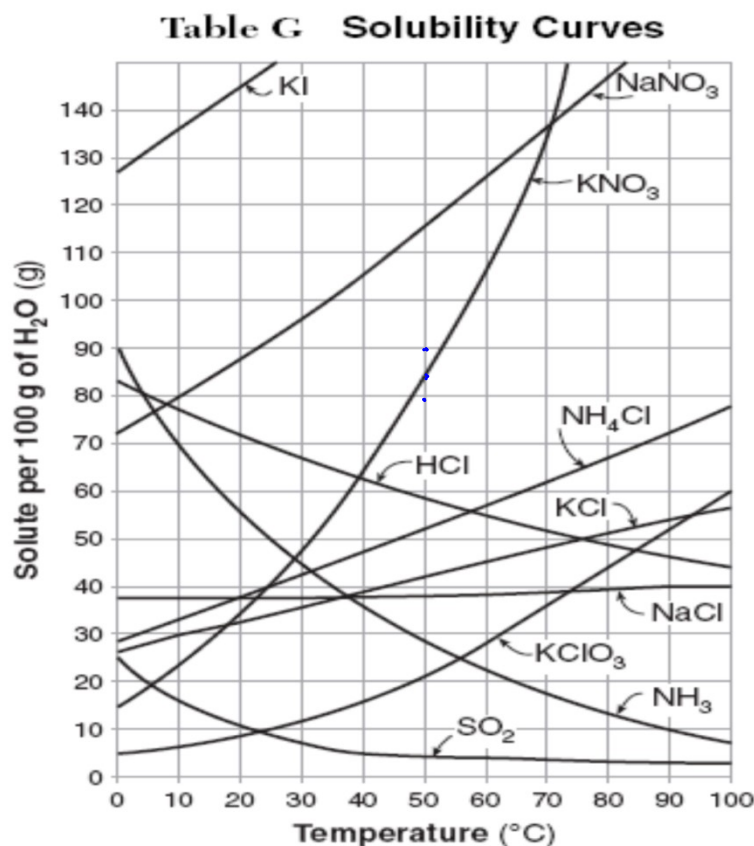


- ✓ **SUPERSATURATED:** a soln that contains **MORE SOLUTE** than would normally dissolve in a given volume of solvent; usually requires an **INCREASE** in **TEMPERATURE** or **PRESSURE** *initially* (**VERY UNSTABLE**)
  - *Very rare! Allow me to demonstrate...*

### Situation:

- If you have an UNSATURATED solution and you add crystals, what will you observe? **CRYSTALS DISSOLVE/DISAPPEAR**
  - The rate of dissolving is **GREATER** than the rate of precipitation
- If you have a SUPERSATURATED solution and you add crystals, what will you observe? **ALL CRYSTALS PRECIPITATE OUT**
  - The rate of dissolving is **LESS** than the rate of precipitation
- If you have a SATURATED solution and you add crystals, what will you observe? **\*CRYSTALS SINK TO BOTTOM**
  - \*The rate of dissolving is **EQUAL** to the rate of precipitation

## SOLUBILITY CURVES - USING TABLE G (of the Reference Tables):



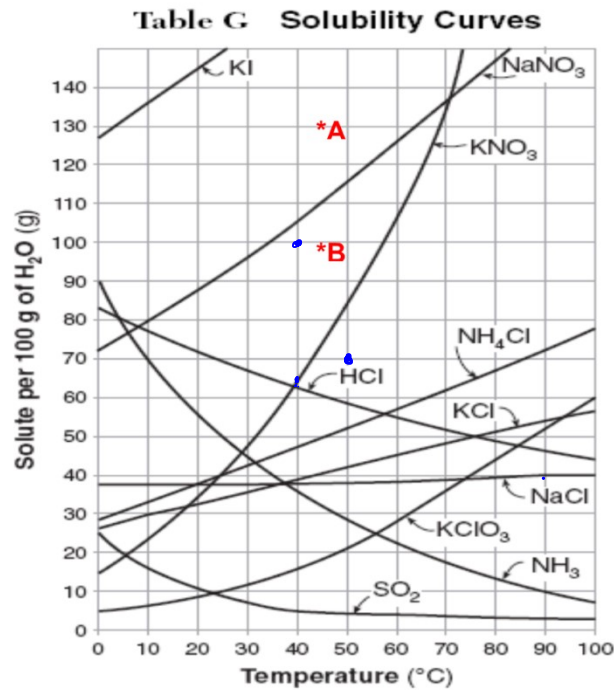
**HOW TO USE TABLE G:** Table G shows the number of grams of a substance that can be dissolved in **100 GRAMS OF WATER** at temperatures between 1°C and 100°C. Each curve represents the maximum amount of a substance that can be dissolved at a given temperature. All lines that show an increase in solubility as temperature increases represent **SOLIDS**. The lines for **GASES** slope downward.

**\*\*\*If the intersection of GRAMS OF SOLUTE vs. TEMPERATURE falls:**

- **ON THE CURVE**, solution is **SATURATED**
- **UNDER THE CURVE**, solution is **UNSATURATED**
- **ABOVE THE CURVE**, solution is **SUPERSATURATED**



Use Table G to answer the following questions:



- How many grams of NaCl will dissolve in 100 g of H<sub>2</sub>O at 90°C? 40g
- How many grams of KCl will dissolve in 50g of H<sub>2</sub>O at 30°C? 100g 38g ≈ 18g ± 1g
- How many grams of KCl will dissolve in 200g of H<sub>2</sub>O at 30°C? 276g 11g
- Name two substances on Table G that are solids.  
KI + NaCl (slope up)
- Name two substances on Table G that are gases.  
HCl + NH<sub>3</sub> (slopes down)
- Is a solution that is made up of 100 grams of H<sub>2</sub>O and 70 grams of KNO<sub>3</sub> at 50°C saturated, unsaturated, or supersaturated? Why?  
unsaturated
- At 40°C, if 100 grams of KNO<sub>3</sub> are added to 100 grams of water, how many grams of solute will fall to the bottom of the solution (and won't dissolve)?  
100g - 65g = 35g will fall out of solution
- Which is more concentrated on Table G, point A or point B? Why?  
point A is more concentrated

## SOLUBILITY FACTORS (Why do some things dissolve and others do not?)

### Factor 1 – Nature of Solute & Solvent

	<i>Acetone</i> Nonpolar Solvent	<i>Water</i> Polar Solvent
<i>Nail polish</i>	<b>SOLUBLE</b> <u>          </u>	<b>INSOLUBLE</b> <u>          </u>
<i>Sugar</i>	<b>INSOLUBLE</b> <u>          </u>	<b>SOLUBLE</b> <u>          </u>
<i>Salt</i>	<b>INSOLUBLE</b>	<b>SOLUBLE</b>

### Factor 2 – Temperature

- For solids, as **temperature ↑**, **solubility ↑**

**Remember:** Table G is there to help you!  
Gases slope ↓ solids slope ↑.

- For MOST gases, as **temperature ↑**, **solubility ↓**  
Example: heat soda → carbonation is LOST!  
(warm)

### Factor 3 – Pressure

- For solid or liquid solutes → **LITTLE TO NO EFFECT**
- For gases dissolved in liquids → As pressure increases, **VOLUME ↓, SOLUBILITY ↑**

Recall: We can change the volume of gases!

Reason: Gas is forced into less space per liquid, so it DISSOLVES more!

Ex: can of soda



When packaged, soda is pressurized (so  $\text{CO}_2$  dissolves in the liquid). When you open the can, pressure is let out (pressure decreases in the can). So the gas particles are released and you get CARBONATION!

CartoonChurch.com



I AM FINDING IT DIFFICULT  
TO CONCENTRATE BUT I  
AM NOT SURE WHY



## SOLUBILITY - TABLE F:

- ❖ Tells you what substances are soluble or insoluble
- ❖ Is used to predict whether or not a precipitate will form when two ionic solutions are mixed and undergo a double replacement reaction

**Precipitate** = **INSOLUBLE** product that falls to the bottom of a mixture/solution (following a double replacement rxn.)

- ❖ A precipitate will form (or a visible reaction will occur) if one or both of the products listed is insoluble (this is a precipitate!)

Ions That Form Soluble Compounds	(insoluble) Exceptions	Ions That Form Insoluble Compounds	(soluble) Exceptions
Group 1 ions (Li <sup>+</sup> , Na <sup>+</sup> , etc.)		carbonate (CO <sub>3</sub> <sup>2-</sup> )	when combined with Group 1 ions or ammonium (NH <sub>4</sub> <sup>+</sup> )
ammonium (NH <sub>4</sub> <sup>+</sup> )		chromate (CrO <sub>4</sub> <sup>2-</sup> )	when combined with Group 1 ions, Ca <sup>2+</sup> , Mg <sup>2+</sup> , or ammonium (NH <sub>4</sub> <sup>+</sup> )
nitrate (NO <sub>3</sub> <sup>-</sup> )		phosphate (PO <sub>4</sub> <sup>3-</sup> )	when combined with Group 1 ions or ammonium (NH <sub>4</sub> <sup>+</sup> )
acetate (C <sub>2</sub> H <sub>3</sub> O <sub>2</sub> <sup>-</sup> or CH <sub>3</sub> COO <sup>-</sup> )		sulfide (S <sup>2-</sup> )	when combined with Group 1 ions or ammonium (NH <sub>4</sub> <sup>+</sup> )
hydrogen carbonate (HCO <sub>3</sub> <sup>-</sup> )		hydroxide (OH <sup>-</sup> )	when combined with Group 1 ions, Ca <sup>2+</sup> , Ba <sup>2+</sup> , Sr <sup>2+</sup> , or ammonium (NH <sub>4</sub> <sup>+</sup> )
chlorate (ClO <sub>3</sub> <sup>-</sup> )			
perchlorate (ClO <sub>4</sub> <sup>-</sup> )			
halides (Cl <sup>-</sup> , Br <sup>-</sup> , I <sup>-</sup> )	when combined with Ag <sup>+</sup> , Pb <sup>2+</sup> , and Hg <sub>2</sub> <sup>2+</sup>		
sulfates (SO <sub>4</sub> <sup>2-</sup> )	when combined with Ag <sup>+</sup> , Ca <sup>2+</sup> , Sr <sup>2+</sup> , Ba <sup>2+</sup> , and Pb <sup>2+</sup>		

COLUMN 1

COLUMN 2

COLUMN 3

COLUMN 4

## HOW TO USE TABLE F:

- ❖ Find one of the ions in your formula in either **Column 1** or **Column 3**
- ❖ Check if there are any exceptions to that ion (glance over to Column 2/Column 4)
- ❖ If there are no exceptions, scroll up Column 1 or 3 (whichever you are in) to find if your compound is soluble or insoluble
- ❖ If there is an exception (that ion listed in column 2 or 4 as an exception will be what your ion is bonded to), then your answer is the heading to Column 2 or 4.

**PRACTICE:** Using Table F, determine whether the following are **SOLUBLE** or **INSOLUBLE**

1) **KNO<sub>3</sub>** Soluble  
 2) **CaCO<sub>3</sub>** Insoluble

3) **MgSO<sub>4</sub>** Soluble  
 4) **NH<sub>4</sub>OH** Soluble

**\*SOLUBILITY:** a measure of how much SOLUTE will dissolve in a certain amt. of SOLVENT @ a given temp.

- **Soluble:** will dissolve; AKA miscible  
*\*The more soluble something is, the more concentrated it can be in soln*
- **Insoluble:** will NOT dissolve
- **Concentration:** RATIO of the amt. of solute per amount of solvent (SOLUTE/SOLVENT)

**DILUTE:** soln that has a relatively SMALL amount of SOLUTE dissolved into a relatively LARGE amount of SOLVENT (WEAK solution)  
(verb: to WEAKEN the CONCENTRATION of a soln)

$$\text{Dilute} = \frac{\text{solute}}{\text{SOLVENT}}$$

**CONCENTRATED:** Solution that has a relatively LARGE amount of SOLUTE dissolved into a relatively SMALL amount of SOLVENT (STRONG solution)

$$\text{Concentrated} = \frac{\text{SOLUTE}}{\text{solvent}}$$

## Most vs. Least Concentrated (Using Table F)

(STRONG) (WEAK)

- **CONCENTRATION** is directly related to amt. of solute **DISSOLVED**

➤ REGENTS QUESTION:

- Most concentrated = **SOLUBLE** compounds
- Least concentrated = **INSOLUBLE** compounds

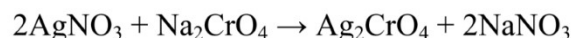
*\*If a compound doesn't dissolve, it doesn't add to the concentration of the soln*

## REACTIONS AND TABLE F:

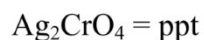
Click here to watch vodcast: <https://www.youtube.com/watch?v=tB4O09KnFVo&feature=youtu.be>

**EXAMPLE:** Silver nitrate and sodium chromate solutions are mixed together. Will a precipitate form? If so, what is the name of the precipitate?

**STEP 1** → Write the chemical or word equation for the double replacement reaction

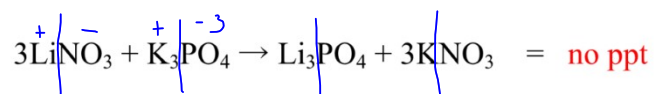


**STEP 2** → Check the solubilities for both products (on Table F). If there are any that are insoluble according to Table F, you have found your precipitate (ppt)!

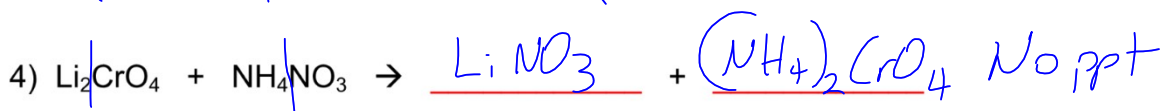


### NOW YOU TRY SOME:

1) Will a precipitate form when lithium nitrate is mixed with potassium phosphate?



For the following reactions, determine the products formed and then circle any precipitate formed as a product:



## Calculating [Concentration] - all these formulas can be found on Table T!

Use the following methods/formulas to calculate the concentrations of the solutions below.  
SHOW YOUR WORK.

<p><b>Molarity (M)</b></p> <p>Formula:</p> $\text{molarity} = \frac{\text{moles of solute}}{\text{liters of solution}}$ <p><b>Example:</b> What is the molarity of a solution that contains 4.0 moles of NaOH in 0.50 L solution?</p> $\text{molarity} = \frac{4.0 \text{ mol}}{0.5 \text{ L}} = 8.0 \text{ M}$	<p><b>Parts per Million (ppm)</b></p> <p>Formula:</p> $\text{PPM} = \frac{\text{mass of solute}}{\text{mass of whole}} \times 1,000,000$ <p><b>Example:</b> Approximately 0.0043 g of oxygen can be dissolved in 100. mL of = 100g H<sub>2</sub>O water at 20°C. Express this in terms of part per million.</p> $\text{PPM} = \frac{0.0043 \text{ g}}{100 \text{ g}} \times 1,000,000 = 43 \text{ ppm}$
<p><b>Percent by Mass</b></p> <p>Formula: <math>\% \text{ Comp} = \frac{\text{mass of part}}{\text{mass of whole}} \times 100</math></p> <p><b>Example:</b> What is the percent by mass of sodium hydroxide if 2.50 g of NaOH is added to 50.00 g of H<sub>2</sub>O?</p> $\% \text{ Comp} = \frac{2.50}{50.0} \times 100 = 4.76\%$	<p><b>Percent by Volume</b></p> <p>Formula: <math>\% \text{ Comp} = \frac{\text{mass of part}}{\text{mass of whole}} \times 100</math></p> <p><b>Example:</b> What is the percent by volume of alcohol if 50.0 mL of ethanol is diluted with water to form a total volume of 300. mL?</p> $\% \text{ Comp} = \frac{50 \text{ mL}}{300 \text{ mL}} \times 100 = 16.7\%$

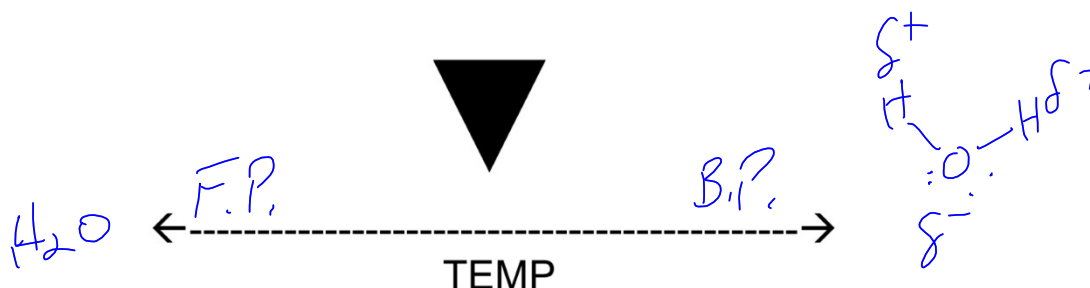
## COLLIGATIVE PROPERTIES FOR ELECTROLYTES

(Based on **NUMBER OF PARTICLES**)

Electrolyte = A substance that produces **IONS** when dissolved in a solution. Because the ions are **FREE TO MOVE AROUND (MOBILE)** in the solution, the solution is able to **CONDUCT** electricity (salts).

\*Same **concept** as mobile electrons in metals, but not the same thing. Don't get confused on the Regents!!!

When a solute is dissolved in a solvent, solvent molecules surround the particles of the solute. This causes the boiling point and freezing point of the solution to change in a very specific and predictable way.



Boiling Point Elevation = b.p. **INCREASES** when solute is added

Ex: Adding salt to water allows you to boil pasta at 102-103°C (cooks it faster)

Freezing Point Depression = f.p. **DECREASES** when solute is added

Ex: Putting salt on roads causes ice to melt because it drops f.p. below 0°C.



**IONIC VS MOLECULAR SOLUTES:**

\* Why do we salt the roads in the winter rather than sugar them?!?!?

	MOLECULAR: C <sub>12</sub> H <sub>22</sub> O <sub>11</sub> (sucrose – covalent!)	IONIC: Salt (NaCl)
Reaction produced when solute dissolved in water	$\text{C}_{12}\text{H}_{22}\text{O}_{11}(\text{s}) \xrightarrow{\text{H}_2\text{O}} \text{C}_{12}\text{H}_{22}\text{O}_{11}(\text{aq})$ <p style="text-align: center;">1 mol                  1 mol</p>	$\text{NaCl}(\text{s}) \xrightarrow{\text{H}_2\text{O}} \text{Na}^+(\text{aq}) + \text{Cl}^-(\text{aq})$ <p style="text-align: center;">1 mol                  2 mol</p>
Number of moles of product produced	1 mol → 1 mol	1 mol → 2 mol
General Rule	The solute that dissolves to form a greater number of products (ions) will have a greater f.p. depression (f.p. will ↓ by more) and a greater boiling point elevation (b.p. will ↑ by more)	
Example:	$\text{CaCl}_2(\text{s}) \xrightarrow{\text{H}_2\text{O}} \text{Ca}^{2+}(\text{aq}) + 2\text{Cl}^-(\text{aq})$	
# ions produced:	1 mol → 3 mol	

\*CaCl<sub>2</sub> is even better than NaCl, so why don't we use it to salt our roads?

**Because IT COSTS MORE MONEY!**

Freezing point depressions (laboratory conditions):

NaCl = -29.4°C (-21°F)

CaCl<sub>2</sub> = -51°C (-60°F)

\*\*NaCl becomes ineffective as a melting agent at around -9°C (15°F). CaCl<sub>2</sub> works effectively at temperatures as low as -29°C (-20°F).