# Unit 5 – Liquids, Solids, and Solutions

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# **Objectives:**

## **Intermolecular Forces (IMFs)**

- Describe hydrogen bonding, London Dispersion Forces, and Dipole-Dipole Forces
- Identify the type of IMF based on the substance
- Relate the IMF to the physical properties of the substance

## **Properties of Water**

- Describe the structure of a water molecule
- Explain how the physical properties of water are determined by the structure of water

## **Solutions**

- Distinguish solutions from suspensions and colloids
- Identify and explain physical and chemical factors that affect solubility
- Define concentration terms of molarity, molality, and percent by volume.
- Perform calculations using molarity (M) and molality (m)

## **Ions in Aqueous Solutions and Colligative Properties**

- Describe the dissociation of ionic compounds and the ionization of some molecular compounds when they dissolve in water
- Define electrolyte
- Distinguish between strong and weak electrolytes
- Write a net ionic equation using the solubility rules
- Perform calculations with different solution concentrations such as molarity, mass percent, molality, and mole fraction.
- Discuss effects of temperature, pressure, and structure on solubility.

- Use a solubility chart to determine if a solution is unsaturated, saturated, or supersaturated.
- Define colligative properties and use to calculate boiling point elevation, freezing point depression, and vapor pressure lowering
- Use colligative properties to determine the molar mass of a solute

## Vocabulary

- Capillary action
- Surface tension
- Soluble
- Insoluble
- Precipitate
- Precipitation
- Aqueous
- Solubility
- Solution
- Electrolyte
- Nonelectrolyte
- Solute
- Solvent
- Solvation
- Effervescence
- Henry's Law
- Hydration
- Immiscible
- Miscible
- Saturated
- Unsaturated

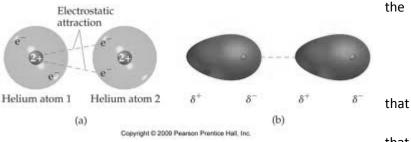
- Supersaturated
- Solution equilibrium
- Concentration
- Molality
- Molarity
- Dissociation
- Ionization
- Net ionic equation
- Spectator ion
- Strong electrolyte
- Weak electrolyte
- Colligative property
- Boiling point elevation
- Freezing point depression
- Molal boiling point constant (k<sub>b</sub>)
- Molal freezing point constant (k<sub>f</sub>)
- Vapor pressure lowering
- Hydrogen Bonding
- London Dispersion Forces
- Dipole-Dipole Forces
- Van der Waals Forces

## Lesson 5.1: Don't Flip your Lid and Intermolecular Forces

**Comparing Intermolecular Forces** 

The forces that hold one molecule to another molecule are referred to as *intermolecular forces (IMFs)*. IntrAmolecular forces are the bonds (ionic or covalent) that hold atoms together within a molecule. These forces arise from unequal distributions of the electrons in the molecule and the electrostatic attraction between oppositely charged portions of molecules.

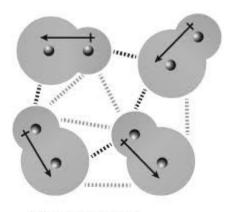
Van der Waals forces are a function of number of electrons in a given molecular and how tightly those electrons are held. The term Van der Waals force is a generic umbrella term used to describe IMFs. Let us assume the molecule involved is nonpolar. A good example would be O<sub>2</sub>. Pretend



that

the molecule is alone in the universe. If that were the case, the electrons in the molecule would be perfectly symmetrical. However, the molecule is not really alone. It is surrounded by other molecules that are constantly colliding with it. When these collisions occur, the electron cloud around the molecule is distorted. This produces a momentary induced dipole within the molecule. The amount of distortion of the electron

could is referred to as polariziability. Since the molecule now has a positive side and a negative side, it can be attracted to the other molecules. This attractive force is called a London dispersion force. Since all molecules have electrons, all molecules have London forces. These forces range from  $5 - 40 \, \text{kJ/mol}$ .

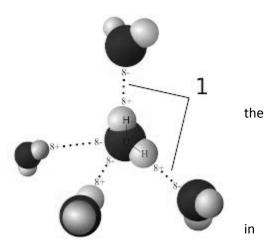


attraction .....repulsion

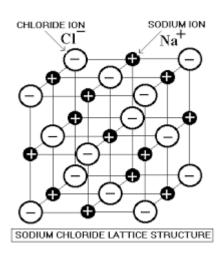
Some molecules are naturally polar; therefore, in addition to dispersion forces, they can also have a permanent dipole which attracts other polar molecules (either induced or permanent). This is called a dipole-dipole force.

A third type of intermolecular force is *hydrogen bonding*. When hydrogen is covalently bonded to a small electronegative atom like nitrogen, oxygen or fluorine, electron cloud on the hydrogen is very distorted and

hydrogen is very distorted and pulled toward the electronegative atom. Since hydrogen has no inner core electrons, the positive nuclear charge is somewhat exposed. This sets up the potential for a reasonably strong attraction between this hydrogen and an electronegative atom



another molecule. Hydrogen bonds are significant in determining such factors as the high boiling point of water, solubility of acetone in water, and the shape and structure of proteins and DNA.



lonic bonds are formed between charged particles. Ionic compounds do not form molecules, but rather are help together in a crystal by electrostatic attraction. These electrostatic attractions are strong and are omnidirectional.

## **PURPOSE**

In this laboratory activity you will determine the relative melting points of four substances, paraffin, NaCl (table salt),  $C_6H_{12}O_6$  (glucose) and iodine ( $I_2$ ). Paraffin is a medium size nonpolar molecule, NaCl is an ionic compound. Glucose is a medium size polar molecule and Iodine is a large nonpolar molecule.

## **MATERIALS**

Bunsen burner Ring stand Tin Can Lid Solid samples of the following: sodium chloride (NaCl), glucose ( $C_6H_{12}O_6$ ), paraffin, iodine ( $I_2$ ).

SAFETY ALERT!!! Wear your goggles. Do not inhale the iodine vapors. Allow the can lid to cool before dispersal

## **PROCEDURE**

- 1. In the space marked HYPOTHESIS on your student answer page, write a statement predicting in which order the salt, paraffin, iodine and glucose will melt.
- 2. Set up your ring stand. Place the can lid on the ring stand
- 3. Transfer a small amount of each of the four samples into one of the four areas in a groove of the outer side of the can lid. You only need a very small sample just enough to be able to see the compound. The teacher will transfer the crystal of iodine.
- 4. Turn the hotplate on and observe. You will need to record the relative order of melting. As soon as three of the compounds melt, turn off the hot plate.
- 5. Allow the can lid to cool before using tongs to dispose of it in a trash can or as your teacher directs.
- 6. Answer the CONCLUSIONS questions on your student answer page.

## **HYPOTHESIS:**

OBSERVATIONS: Record the relative order of melting for the four compounds.

## **ANALYSIS:**

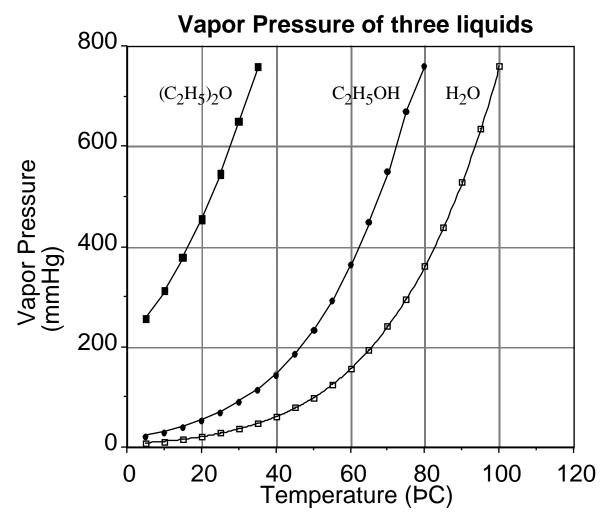
From what you know about intermolecular forces, explain the relative order of the melting points.

## **CONCLUSIONS QUESTIONS:**

- 1. Glucose, C<sub>6</sub>H<sub>2</sub>O<sub>6</sub>, and sucrose, C<sub>12</sub>O<sub>22</sub>O<sub>11</sub>, are both sugars with similar polarities. Which do you think would have the higher melting point? Justify your answer.
- 2. Hydrogen sulfide is a gas at room temperature, while water is a liquid, yet hydrogen sulfide has more electrons that water. Explain this anomaly.
- 3. Consider the halogens at room temperature and 1 atmosphere of pressure. Why are fluorine and chlorine gases at room temperature, while bromine is a liquid and iodine is a solid?

There are three basic intermolecular forces that hold covalent molecules together as either solids or liquids. To change the phase of a substance, the IMFs must be overcome. The amount of energy required for the phase change is dependent on the substance and the strength (type) of the IMF. Typically substances that contain hydrogen bonding will require more energy for the phase change to occur. Substances with LDFs will typically require the least amount of energy.

Vapor pressure is the pressure due to particles of a substance in the vapor phase above its liquid in a closed container at a given temperature. The weaker the forces holding the liquid together, the higher the vapor pressure of the liquid will be.

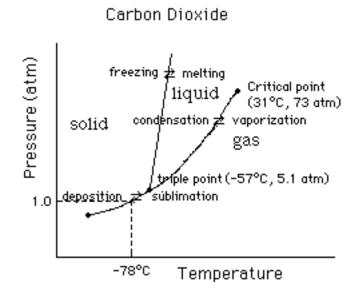


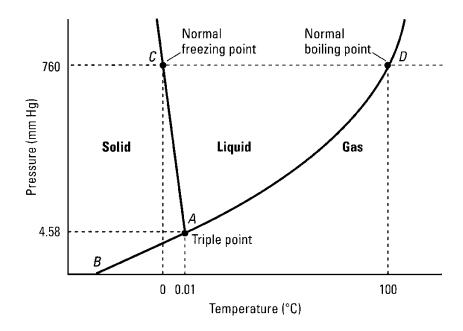
Boiling point is the temperature at which the vapor pressure of a liquid equals the atmospheric pressure. The normal boiling point is the temperature at which the vapor pressure of the liquid equals 1 atmosphere.

The molar heat of vaporization ( $\Delta H_{vap}$ ) is the amount of heat required to change one mole of a pure liquid into one mole of a pure gas. The molar heat of fusion ( $\Delta H_{fus}$ ) is the amount of heat required to change one mole of a pure substance from a solid to a liquid. What generalization can be made between IMFs and boiling temperature?

## **Phase Diagrams**

Phase diagrams are a way of graphically representing the relationship of the three states of a pure substance. Below is an example of a phase diagram for carbon dioxide.



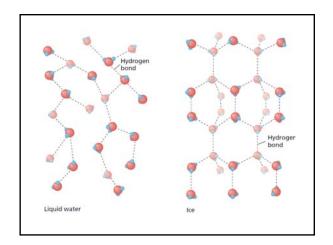


Above is the phase diagram for water. One very important point to notice is the negative slope of the solid liquid interface. This shows that the density of ice is actually less than the density of water - a very unusual property of water that makes much of life possible.

The phase of substance is not only dependent on the IMF present but also on the physical conditions, temperature and pressure.

# Solutions- Day 1 \*\*These homework patches really reduce your cravings. This one is 'Chemistry,' but you can get them in any subject.\*\*

# A group of water molecules will orient at the molecular scale because of electrical forces. This effect is especially great at low temperatures. The water molecules move less rapidly, and they are able to be drawn closer together by dipole-dipole attractions. The volume of the water decreases because the molecules pull together.



# Surface Tension • A water molecule forms a drop because of surface tension, which is the force needed to overcome intermolecular forces and break through the surface of a liquid or spread the liquid out. • The higher the surface tension of a liquid is, the more resistant the liquid is to having its surface broken. • The net inward force makes the surface of the drop contract

# **Capillary Action**

- the rising of liquids in narrow tubes, also known as capillarity
- Capillarity results from the competition between the IMFs between the molecules of liquid and the attractive forces between the liquid and the tube that contains it.

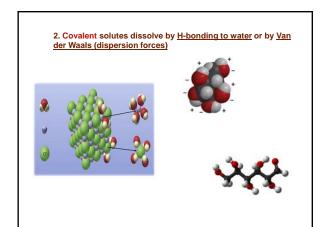
## Like Dissolves Like

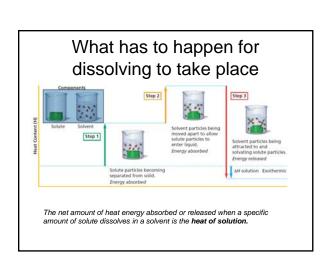
- Polar dissolves polar
- Non polar dissolves non polar

## **Definitions**

- A solution is a homogeneous mixture
- A solute is dissolved in a solvent.
  - solute is the substance being dissolved
  - solvent is the liquid in which the solute is dissolved
  - an aqueous solution has water as solvent

# Dissolution of Solid Solute (Solvation) What are the driving forces which cause solutes to dissolve to form solutions? 1. lonic solutes dissolve by dissociation into their ions. Video of dissociation: http://www.youtube.com/watch?v=EBfGcTAJF4o





## Solution and Concentration

- 3 ways of expressing concentration
- -Molarity(M): = moles solute / Liter solution
- -Mass percent = (mass solute / mass of solution) x 100%
- -Molality\* (m) = moles solute / kg solvent
  - \* Note that <u>molality</u> is the only concentration unit in which denominator contains only <u>solvent</u> information rather than solution.
- Chemists never apply the terms strong and weak to solution concentrations always use concentrated or dilute.

## % Concentration

• % (m/m) = 
$$\frac{mass\ solute}{mass\ solution}$$
 x100

• % (m/v) = 
$$\frac{mass\ solute}{volume\ solution}$$
 x100

• % 
$$(V/V) = \frac{volume \ solute}{volume \ solution} x 100$$

## **Example Problems**

- 1) How many grams of beryllium chloride are needed to make 125 mL of a 0.0500 M beryllium chloride solution?
- 2) How many grams of beryllium chloride would you need to add to 125 mL of water to make a 0.0500 molal solution?
- 3) What will the volume (in L) of a 0.500 M solution be if it contains 25.0 grams of coloium bydrovide?
- 4) How many grams of ammonia are present in 5.00 L of a 0.0500 M solution?
- 5) If 3.50 g of CoCl<sub>2</sub> is dissolved in 100. mL solvent, what is concentration of the solution in % mass (m/m)? (Assuming the <u>density</u> of the solvent is 1.00 g/mL)
- 6) Explain how to make at least 1.00 liter of a 1.25 molal ammonium hydroxide solution
- 7) Explain how to make 1.00 liters of a 1.25 molar ammonium hydroxide solution.

## Dilution

- -When a solution is diluted, solvent is added to lower its concentration.
- -The amount of solute remains constant before and after the dilution:
- -Therefore moles BEFORE = moles AFTER  $M_1V_1 = M_2V_2$

M= concentration in molarity

## Example Problems

v= volume

- 1) If 25.0 mL of water is added to 125 mL of a 0.150 M NaOH solution, what will the molarity of the diluted solution be?
- 2) Given 345 mL of a 1.50 M NaCl solution. If the solution is boiled so its volume decreases 95.0 mL, what will the molarity of the solution be?
- 3) How much water would have to be added to 500. L of a 2.40 M KCl solution to make it a 1.00 M solution?

# Solutions Day 2



## How to describe a solution

- A <u>saturated</u> solution contains the maximum amount of solute per amount of the solution under the given conditions - no more solute is able to dissolve.
  - A saturated solution represents an equilibrium: the rate of dissolving is equal to the rate of crystallization. The salt (solute) continues to dissolve, but crystallizes at the same rate so that there "appears" to be nothing happening.
- A <u>unsaturated</u> solution contains less dissolved solute dissolved than the maximum that could be dissolved.
- A <u>supersaturated</u> solution more dissolved solute than the usual maximum amount and are unstable.

## Factors Affecting Solubility Liquids and Gases

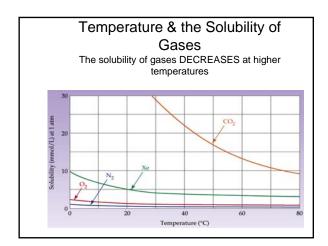
- 1. Nature of Solute / Solvent.  $\underline{\text{Like dissolves like}} \ (IMF) \ need \ to \ have \ polar \ with \ polar \ or \ non \ polar \ with \ non \ polar$
- 2. Temperature -
- i) Solids/Liquids- Solubility increases with Temperature (usually)
   Increase K.E. increases motion and collision between solute / solvent.
- ii) gas Solubility decreases with Temperature
- Increase K.E. result in gas escaping to atmosphere.
- 3. Pressure Factor -
- i) Solids/Liquids Very little effect

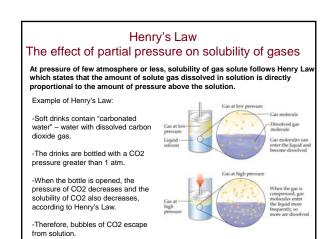
Solids and Liquids are already close together, extra pressure will not increase solubility.

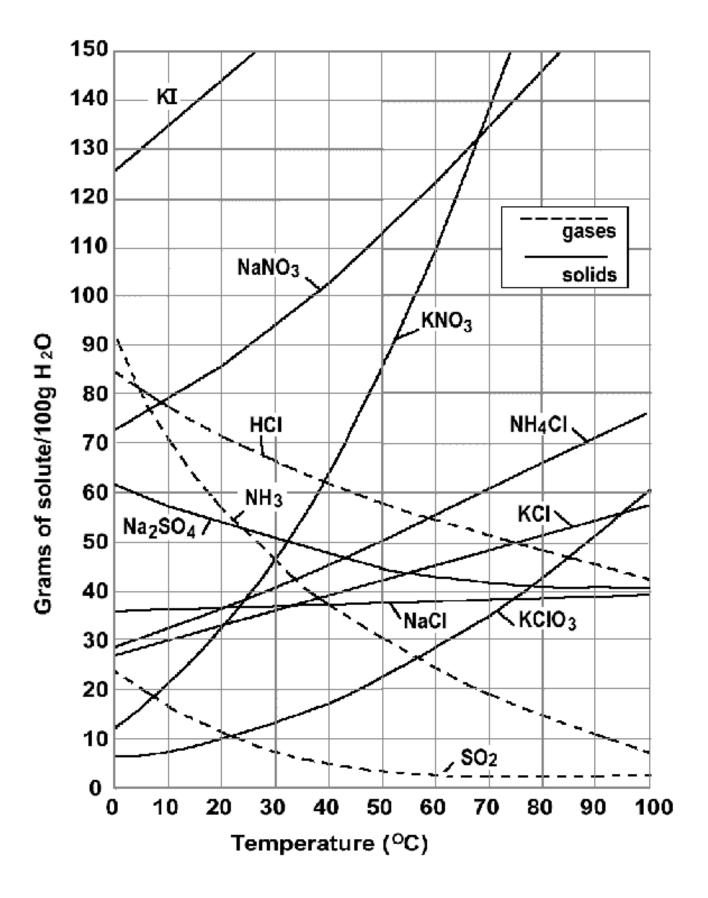
ii) gas - Solubility increases with Pressure.

Increase pressure squeezes gas solute into solvent.

### Solubilities of Ionic Solids vs Temperature Solubilities of several 90 ionic solid as a function of 100 g H<sub>2</sub>O) temperature. MOST salts have greater solubility in hot water. 60 Heat of solution of most ionic solids is positive endothermic. (g of salt i 50 40 A few salts have NaC negative heat of 30 solution, (exothermic 20 process) and they become less soluble 10 with increasing Ce2(SO4) temperature. 10 20 30 40 50 60 70 80 90 100







**Solubility Curves** Based on the solubility below, decide whether each of the following is A: unsaturated, B: saturated, C: supersaturated, or whether D: not enough information is given. \* assume it's dissolved \* 1) 50 g KCl in 100 g of water at 4) 50 g KNO<sub>3</sub> in 25 g of water at 7) 25 g NaCl in 100 g of water. 90°C. 60°C. 2) 50 g KCl in 100 g of water at 5) 65 g  $KNO_3$  in 50 g of water at 60°C. \_\_\_\_ 8) 40 g of KCl in 100 g of water at 20°C. \_\_\_\_ 3) 50 g KNO<sub>3</sub> in 100 g of water 6) 25 g KNO<sub>3</sub> in 100 g of water. at 60°C. 9) How many grams of KCl can dissolve in 100.0 g of water at 65°C? \_\_\_\_\_ 10) What temperature would be required to get 85 g of KNO<sub>3</sub> to dissolve in 100.0 g of water? SHOW ALL WORK FOR THE FOLLOWING KNO<sub>3</sub> 180 170 11) How many grams of KNO<sub>3</sub> can be dissolved in 50.0g of 160 water at 50.0°C? \_\_\_\_\_ 150 140 130 12) What mass of KCl can be dissolved in 200.0 g of water at 15.0°C? \_\_\_\_\_ 13) How much KNO<sub>3</sub> can be dissolved in 14.3 g of water at 69.0°C? \_\_\_\_\_ **KCI** 14) How many grams of water will it take to dissolve 28.0 g NaCl NaCl at 60.0°C? \_\_\_\_\_ 20 15) How much water is needed to dissolve 46.6 g of KNO3 at 10 20 30 40 50 60 70 80 90 100 52°C? \_\_\_\_\_ temperature (C) 19) What temperature is required to make a 63.0% 16) What temperature would be required to get 51.0 g of KCl to dissolve in 156 g of water? KNO<sub>3</sub> solution? \_\_\_\_\_

17) What is the % KCl in a solution that is saturated at 61°C? \_\_\_\_\_

18) What temperature is required to make a 50.0%  $KNO_3$  solution?

20) Based on what you've learned in class about soda & fish, do gases behave the same as or different than solids when it comes to solubility & temperature? What do you think a solubility graph of gases would look like?

# Molarity

Determine the concentration (molarity) for each of the	e solutions:
a) 3.0 mol sugar dissolved in 2.0 L of solution	b) 0.40 mol NaCl dis. in 10.0 L of soln
c) $0.030 \text{ mol KNO}_3$ dis. in $50.0 \text{ mL of soln.}$	d) 350 g KNO $_3$ dis. in 5.0 L of soln
e) 6.45 g of Na₂SO₄ dis in 250 mL of soln	f) 465 mg KF of dis. in 0.054 L of soln
2. How many moles of sugar are needed to make 2.5 L of 1.4 M sugar solution? Ans:	6. What volume of 0.25 M sugar solution can be made using 4.0 moles sugar? Ans:
3. How many moles of NaBr are needed to make 150 mL of 3.0 M NaBr solution? Ans:	7. How many mL of 2.50 M Na <sub>3</sub> PO <sub>4</sub> solution can be made using 1.8 g of Na <sub>3</sub> PO <sub>4</sub> ? Ans:
4. How many grams of NaNO₂ are needed to make 3.5 L of 0.50 M NaNO2 solution? Ans:	8. 65.0 mL of $K_3PO_4$ solution are evaporated, and 1.54 g of solid $K_3PO_4$ are recovered. What was the molarity of the original solution? Ans:
5. How many grams of K <sub>2</sub> CO <sub>3</sub> are needed to make 300.0 mL of 1.25 M K <sub>2</sub> CO <sub>3</sub> solution? Ans:	

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n	ш	ΙIŤ	10	ns

1. Determine the concentrations for each of the	e following mixtures:
a) equal volumes of 3.0 M KCl & water:	b) equal volumes of 3.0M KCl & 7.0 M KCl:
c) one vol. of 8.0 M KCl & one vol. water:	d) one vol. of 6.0 M KCl & two vol's water:
e) one vol. water & two vol's of 6.0M KCl:	f) one vol. of 5.0 M KCl & 4 vol's of water:
g) one vol. of 2.5 M KCl & 9 vol's water:	h) one vol. of 2.5 M KCl & 99 vol's water:
	te, one usually mixes the can of concentrate with three cans of (what fraction?) its original concentration.
3. Use the dilution equation to determine the	concentrations of the following mixtures
a) 45 L of 3.6 M KCl & 71 L of water:	b) 215 mL of 2.8 M KCl & 47 mL water:
Ans:	Ans:
c) 45 mL of 3.6 M KCl & 71 mL of 6.2 M KCl:	
	Ans:
d) 83 mL of 2.0 M KCl & 25 mL of water:	e) 38 mL of 6.0 M KCl dil. to a tot vol of 100 mL:
Ans:	Ans:
4. To what total volume must 26.0 mL of 4.80 N a) 2.10 M	M KCl be diluted to reduce its concentration to b) 0.480 M
Ans:	

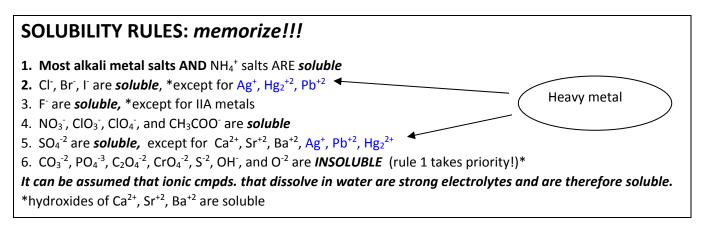
(Warning: one of the questions on this public what volume of water must be added t			
a) 1.2 M	b) . Ans:	0.26 M	Ans:
6. What volume of 2.5 M KCl must be ac	 dded to 37 mL of 6.0	M KCl to make the total co	
a) 1.5 M	b).	4.2 M	
	Ans:		Ans:
7. What volume of 2.5 M KCl must be a	uded to 37 ML of wa	ter to make the total conce	entration 1.8MF
8. You mix 32 mL of 4.5 M KCl, 56 mL of be 1.7 M. How much water must have be		e water, and the total conce	
9. Sketch a volumetric flask and explain some 1.500 M NaNO₃ solution. (You have equipment you need) <b>How much 2.000</b>	ve available some 2.0	000 M NaNO₃ solution and	
10. You need to make up some 5.0 M K	Cl solution but all yo	u have is 125 mL of 3.0 M I	CCI.

Explain what to do to make up the 5.0 M solution. How much 5.0 M KCl will you get? Show calculations: (hint-

calculate how much water to evaporate)

## **Electrolytes, Precipitates and Balancing Net Ionic Equations**

<u>Electrolytes:</u> substances that form ions in aqueous solutions (dissolved in water). Positive and negative ions carry current (conduct electricity) in an aqueous solution. Usually these are soluble ionic salts, strong acids (completely dissociate), and strong bases (completely dissociate). Substances that will not dissociate in solution are insoluble and nonelectrolytes (cannot conduct electricity). The following table is a guide to whether a solution is soluble or not.



Identify the following compounds as either electrolytes or nonelectrolytes. Also list the ions they will form in solution if applicable. Assume all are aqueous.

- 1) LiCl
- 2) (NH<sub>4</sub>)<sub>2</sub>S
- 3) Ammonium Chlorate
- 4) Ca(OH)<sub>2</sub>
- 5) Mg(OH)<sub>2</sub>
- 6) Silver Chloride

<u>Precipitates:</u> An insoluble solid that emerges from an aqueous solution. The emergence of the insoluble solid from solution is called precipitation.

Precipitates can form when two soluble salts react in solution to form one or more insoluble products. The insoluble product separates from the liquid and is called a precipitate.

Precipitates can also form when the temperature of a solution is lowered. The lower temperature lower temperature reduces the solubility of the salt resulting in the formation of a precipitate.

<u>Spectator lons:</u> lons in a solution that do not participate in a chemical reaction.

Balancing Net Ionic Equations: typically these are double replacement/displacement reaction types. Net ionic equations are used to show which chemical species are actively reacting and eliminates spectator ions. The reactions we will be considering will take place in  $H_2O(I)$ .

Below is an example of how these are done.

EXAMPLE:  $KCl_{(aq)} + Pb(NO_3)_{2(aq)} \rightarrow$ 

- Take <u>only one</u> of the first cation(s) and match it with <u>one</u> of the second anion(s). (Write the cation first)
  - b. Take <u>only one</u> of the second cation(s) and match it with <u>one</u> of the first anion(s). (Write the cation first)

$$KCl_{(aq)} + Pb(NO_3)_{2(aq)} \rightarrow KNO_3 + PbCl$$

2. Correct the formulas of the products based on the charges of the ions. (Note the subscript change on CI)

$$KCl_{(aq)} + Pb(NO_3)_{2(aq)} \rightarrow KNO_3 + PbCl_2$$

3. Balance the equation

$$\underline{2} \text{ KCl}_{(aq)} + \text{Pb}(\text{NO}_3)_{2(aq)} \rightarrow \underline{2} \text{ KNO}_3 + \text{PbCl}_2$$

4. Consult the solubility rules and assign the correct state symbol. This should agree with any observations concerning the formation of a precipitate which gets the symbol (s). If water is formed, water is a molecule; it does not ionize to any significant extent. It is given (I). Note the state variables on the product side. Why does KNO<sub>3</sub> get (aq) and PbCl<sub>2</sub> get (s).

2 KCl (aq) + Pb(NO<sub>3</sub>)<sub>2</sub> (aq) 
$$\rightarrow$$
 2 KNO<sub>3</sub> (aq) + PbCl<sub>2</sub> (s)

5. Write the Total Ionic Equation. All compounds that are aqueous (aq) break up into individual cations and anions

$$2 K^{+}(aq) + 2Cl^{-}(aq) + Pb^{2+}(aq) + 2NO_{3}^{-}(aq) \rightarrow 2 K^{+}(aq) + 2NO_{3}^{-}(aq) + PbCl_{2}(s)$$

6. Eliminate spectator ions. Spectator ions are in the same form on each side of the equation arrow.

$$2 K^{+}_{(aq)} + 2CI^{-}_{(aq)} + Pb^{2+}_{(aq)} + 2NO_{3}^{-}_{(aq)} \rightarrow 2 K^{+}_{(aq)} + 2NO_{3}^{-}_{(aq)} + PbCI_{2(s)}$$

7. Write the Net Ionic Equation. The convention is to write the cation first followed by the anion on the "reactants" side. Don't forget that chemical equations are written using the lowest common coefficients (including net ionic equations). If all ions cancel each other out then it is NR.

$$Pb^{2+}_{(aq)} + 2Cl^{-}_{(aq)} \rightarrow +PbCl_{2(s)}$$

Class work Net Ionic Equations. Write and Balance the net ionic equation for the following reactions.

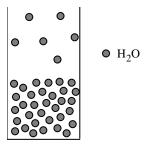
- 1..  $K_3PO_4$  (aq) +  $AI(NO_3)_3$  (aq)  $\rightarrow$
- 2. Bel<sub>2</sub> (aq) + Cu<sub>2</sub>SO<sub>4</sub> (aq)  $\rightarrow$
- 3. Ni(NO<sub>3</sub>)<sub>3</sub> (aq) + KBr (aq)  $\rightarrow$
- 4. cobalt(III) bromide + potassium sulfide →
- 5. barium nitrate + ammonium phosphate →

## **Colligative Properties**

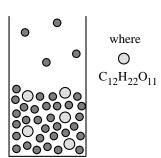
Colligative properties are properties that depend on the <u>number of molecules or ions</u> of solute present, and not on what the particles are (as long as they are not volatile). Properties of solutions that are colligative properties,

- 1) Vapor Pressure lowering
- 2) Boiling Point elevation
- 3) Freezing Point depression

The following diagram shows why a solute will affect the colligative properties



Before addition of the solute: The vapor above a pure liquid is shown on the right. The equilibrium vapor pressure is a result of the presence of molecules in the vapor phase above the molecules in the liquid phase. The figure below shows the effect on the vapor pressure of the solution upon addition of a nonvolatile solute.



After addition of the solute: The addition of a nonvolatile solute decreases the vapor pressure due to the liquid in the vapor phase. This occurs because the presence of solute particles on or near the surface decreases the ability of the solvent particles to escape into the vapor phase. Because proportionally fewer solvent particles are on the surface, the vapor pressure drops.

The above leads to Raoults law:

 $P_{\text{solution}} = \chi_{\text{solvent}} \cdot P_{\text{solvent}}$ 

Where;

P<sub>solution</sub> is the vapor pressure of the solvent above the solution at the particular temperature

$$\chi_{solvent}$$
 is the mole fraction of the solvent in the solution  $\chi_{solvent} = \frac{mol_{solvent}}{mol_{solvent} + mol_{solute}}$ 

P<sub>solvent</sub> is the vapor pressure of the pure solvent at the particular temperature.

The other colligative properties are freezing point depression and boiling point elevation.

$$\Delta T_f = i k_f m$$
 or  $\Delta T_b = i k_b m$ 

where:

 $\Delta T_f$  or  $\Delta T_h$  is the change in freezing point or the change in boiling point.

k<sub>f</sub> or k<sub>b</sub> are the freezing point and boiling point constants (solvent dependent)

$$m$$
 is the molality of the particles in the solution  $\left(\frac{\text{mol solute}}{\text{kg solvent}}\right)$ 

i the Van't Hoff factor and is the number of ions of solute in solution

example: if the substance is a nonelectrolyte - i = 1

$$NaCl - i = 2$$

## **Practice Problems**

1. Calculate the expected vapor pressure of at 25°C for a solution prepared by dissolving 97.4 grams of common table sugar (MM 342 g/mol) in 453 mL of water. Vapor pressure of water at 25°C is 23.8 mmHg (Ans 23.49 mmHg).

2. Glucose is often used as a form of nourishment to hospital patients through intravenous feeding. Calculate the new boiling point and freezing point for a 0.70 m solution of glucose,  $C_6H_{12}O_6$ .

3.	Calcium chloride is a common drying agent and is very exothermic when dissolved in water. If 47.50 grams of calcium chloride is dissolved in 395 mL of water, calculate the boiling point and freezing point of the solution that is formed.
4.	Moth balls are often composed of pure naphthalene, $C_{10}H_8$ . The normal freezing point of naphthalene is 80.2°C. When 5.25 grams of an unknown nonvolatile substance was mixed with 35.00 grams of naphthalene, the freezing point was found to be 78.8°C. Calculate the molecular weight of the unknown substance. The $k_f$ for naphthalene is $6.9^{\circ}C/m$ .
5.	Which solution freezes at a lower temperature, 0.20 <i>m</i> sodium sulfate or 0.20 <i>m</i> magnesium sulfate? Explain.

# **Unsaturated, Saturated, Supersaturated Lab**

Follow the procedure below, and record all observations. Ignore the letters in ( ).

1) Obtain a clean, unscratched test tube. Add 2.0 mL of water (A). Then add 4.00 g of NaC2H3O2 ("sodium

acetate," which we will abbreviate SA from here on), but don't shake yet (B). Use a perm. pen to mark the top of the undissolved SA level on the side of test tube. Stopper and mix for 3 sec, tilt to get any undissolve crystals off the sides, and note any changes (C).
Obs:
2) Re-mark the top of the undissolved solute level. Mix for another 5 sec and observe the changes, including feeling the test tube (D).
Obs:
3) Repeat step #2 until no more change occurs (E).
Obs:
4) REMOVE STOPPER! Heat test tube (by placing in hot water) for 90 sec (while waiting, weigh out another 0.10 g of SA for step #5). Remove test tube from heat, stopper & mix for 10 sec (F).
Obs:
5) While still hot, add the 0.10 additional grams of SA, stopper & mix for 10 sec (G)
Obs:
6) Add an additional 4.00 g of SA, stopper & mix for 10 sec (H).
Obs:
7) <b>REMOVE STOPPER!</b> Heat for 2 min (while waiting, weigh out 1.00 g of SA for step #8), then remove from heat, stopper and mix for 10 sec (I). Obs:
8) Add the 1.00 g of SA & mix (J). Obs:
9) Reheat until all crystals have dissolved (K), stopper and mix, and then cool in cold water for 50-60 sec (L), (In recrystallization occurs during cooling, reheat to redissolve it, then re-cool it.)
* Then add 1 crystal SA & observe (M). Obs:
10) (bonus) Reheat until all crystals have dissolved and then an additional 30 sec (N), make sure your test tuber rim is ULTRA-clean, and cool in water for 50-60 sec (O). Place a crystal or two on a clean petri dish lid. Then, carefully, drop-by-drop, pour your solution out onto the crystal. Observe what happens (P). Advice Don't allow the growing pillar to come too close to the mouth of the test tube.  Obs:

## **QUESTIONS:**

each particular moment, the test tube con	ntained a solution that was unsat, sat. or sup	ersat.
Briefly justify your answers. The first one	is done for you.	
A unsat _it's pure waterno solute	l	
В	J	
C	К	
D	L	
E	M	
F	N	
G	0	
н	P	
cases.	determine whether it was unsaturated, satu lo and what you would expect to see for eac	
supersaturated:		
3. A solution has some undissolved crystals si	itting on the bottom. Could it be	
unsaturated? Y / N Explain:		
saturated? Y / N Explain:		
supersat.? Y / N Explain:		

1. Consider each of the points throughout the procedure indicated by the letters (A-P) and decide whether at

4. Use the solubility curves in the packet to explain **precisely**, step-by-step, how you would go about making a **supersaturated** KNO<sub>3</sub> solution. State precisely how many grams of water, how many grams of KNO3 and what temperatures you would use.

## **Preparing Solutions Lab:**

**Background:** In chemistry, spectrophotometry is the quantitative measurement of the reflection or transmission properties of a material as a function of wavelength. Spectrophotometry involves the use of a spectrophotometer. A spectrophotometer is a photometer that can measure intensity in absorbance and transmittance. In lab you will prepare solutions of Copper (II) Sulfate Pentahydrate and measure their absorbance at a wavelength of 635 nm.

## Fill in the data table below from the board:

Concentration (M)	Absorbance at 635 nm
0.019 M	
0.038 M	
0.075 M	
0.150 M	

0.075 M				
0.150 M				
1) Use the data table above to generate a linea concentration is x and absorbance is y.	r regression equation in the format y=mx +b where			
2) Use the molarity equation, a 50.0 mL volumetric flask, distilled water, and a balance to prepare 50.0 mL of a 0.130 M CuSO <sub>4</sub> solution. Describe below how to make this solution.				
	to prepare a 50.0 mL of a 0.060 M solution using your etric flask. Describe how to make this solution below.			
4) Use a 50.0 mL volumetric flask, distilled water solution. Describe below how to make this solution.	er, and a balance to prepare 50.0 mL of a 0.030 M CuSO <sub>4</sub>			

5)	5) Use your linear regression equation to predict the absorbance of your prepared solutions.				
6)	Use a spectrophotom nm. Record this data		absorbance of each of you	ur prepared solutions at 635	
Co	ncentration (M)	Predicted Absorbance	Actual Absorbance	Percent Error	
UN	KNOWN SAMPLE ABSO	ORBANCE			
PRI	EDICTED MOLARITY OF	UNKNOWN			
Coi	nclusion Questions:				
1)	1) Molarity is considered a temperature dependent concentration unit while molality is considered a temperature independent concentration unit. Why is this true?				
2)	What mass (in grams	) of calcium hydroxide is ne	eded to make 500.0 mL of	a 2.40 M solution?	
3)	To what volume shou solution is 1.50 M?	ıld 200. mL of an 8.00 M Na	aOH solution be diluted so t	that the final molarity of the	