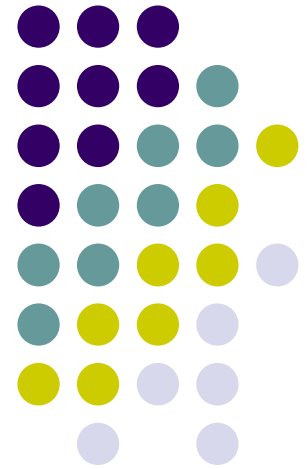


# Colligative Properties of Solutions

**Yanovska Anna Olexandrivna**

PhD physical chemistry

Lecturer of General chemistry  
department





● In our life we can see decreasing of freezing point by adding salt on the road. Due to this road is free of snow and ice.



# IONIC COMPOUNDS

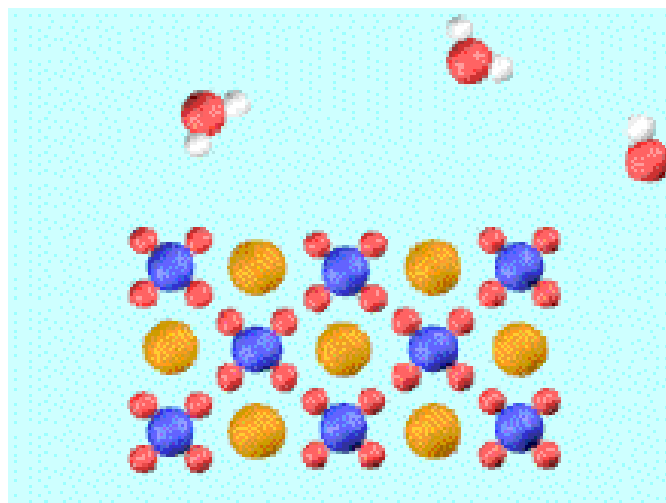
## *Compounds in Aqueous Solution*



Many reactions involve ionic compounds, especially reactions in water — **aqueous solutions**.

**$\text{KMnO}_4$  in water**

**$\text{K}^+(\text{aq}) + \text{MnO}_4^-(\text{aq})$**



# Aqueous Solutions



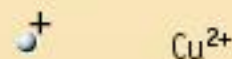
If compounds completely dissociate into ions they are called

**ELECTROLYTES**

HCl,  $\text{MgCl}_2$ , and NaCl are **strong electrolytes**.

Degree of dissociation  $\alpha$  for such electrolytes is 1 or nearly 100%.

Strong Electrolyte



A strong electrolyte conducts electricity.  $\text{CuCl}_2$  is completely dissociated into  $\text{Cu}^{2+}$  and  $\text{Cl}^-$  ions.

# Electrolytes in the Body



- Carry messages to and from the brain as electrical signals
- Maintain cellular function with the correct concentrations electrolytes



Make your own

**50-70 g sugar**  
**One liter of warm water**

**Pinch of salt**  
**200ml of sugar free fruit squash**  
**Mix, cool and drink**



# Aqueous Solutions

Some compounds dissolve in water but do not conduct electricity. They are called **nonelectrolytes**.

**Examples include:**

**sugar**

**ethanol**

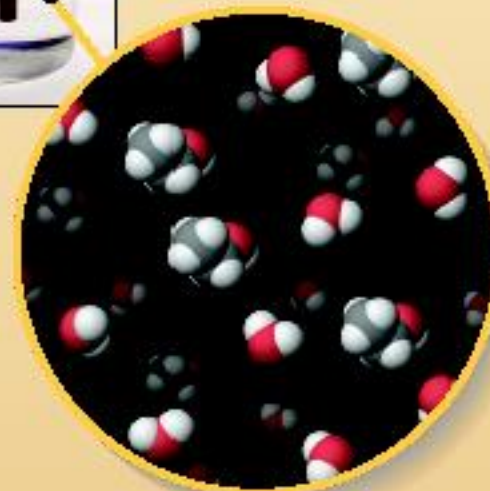
**ethylene glycol**



Nonelectrolyte



Ethanol



A nonelectrolyte does not conduct electricity because no ions are present in solution.

# Constant of dissociation for weak electrolytes



electrolytes

$$K = \frac{[Kt^{y+}]^x \cdot [An^{x-}]^y}{[Kt_xAn_y]}$$

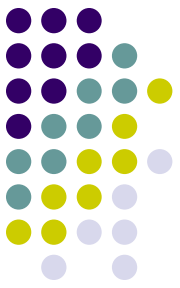
$$[Kt^+] = [An^-] = \alpha C,$$

$$[KtAn] = C - \alpha C = C(1 - \alpha).$$

$$K_d = \frac{[Kt^+][An^-]}{[KtAn]} = \frac{\alpha C \cdot \alpha C}{C(1 - \alpha)} = \frac{\alpha^2 C}{1 - \alpha}$$

$$K = \alpha^2 C,$$

$$\alpha = \sqrt{K / C}.$$



# Colligative properties

- Properties of ideal solutions don't depend on nature of the solute they only depend on its concentration.
- So properties that depend only on the number of solute particles and not on their identity named – ***colligative properties***:
  - Vapor pressure lowering
  - Boiling point elevation
  - Freezing Point depression



# Colligative Properties



On adding a solute to a solvent, the properties of the solvent are modified.

- Vapor pressure decreases
- Melting point decreases
- Boiling point increases
- Osmosis is possible (osmotic pressure)

These changes are called **COLLIGATIVE PROPERTIES**.

They depend only on the **NUMBER** of solute particles relative to solvent particles, not on the **KIND** of solute particles.



# Vapour pressure depression

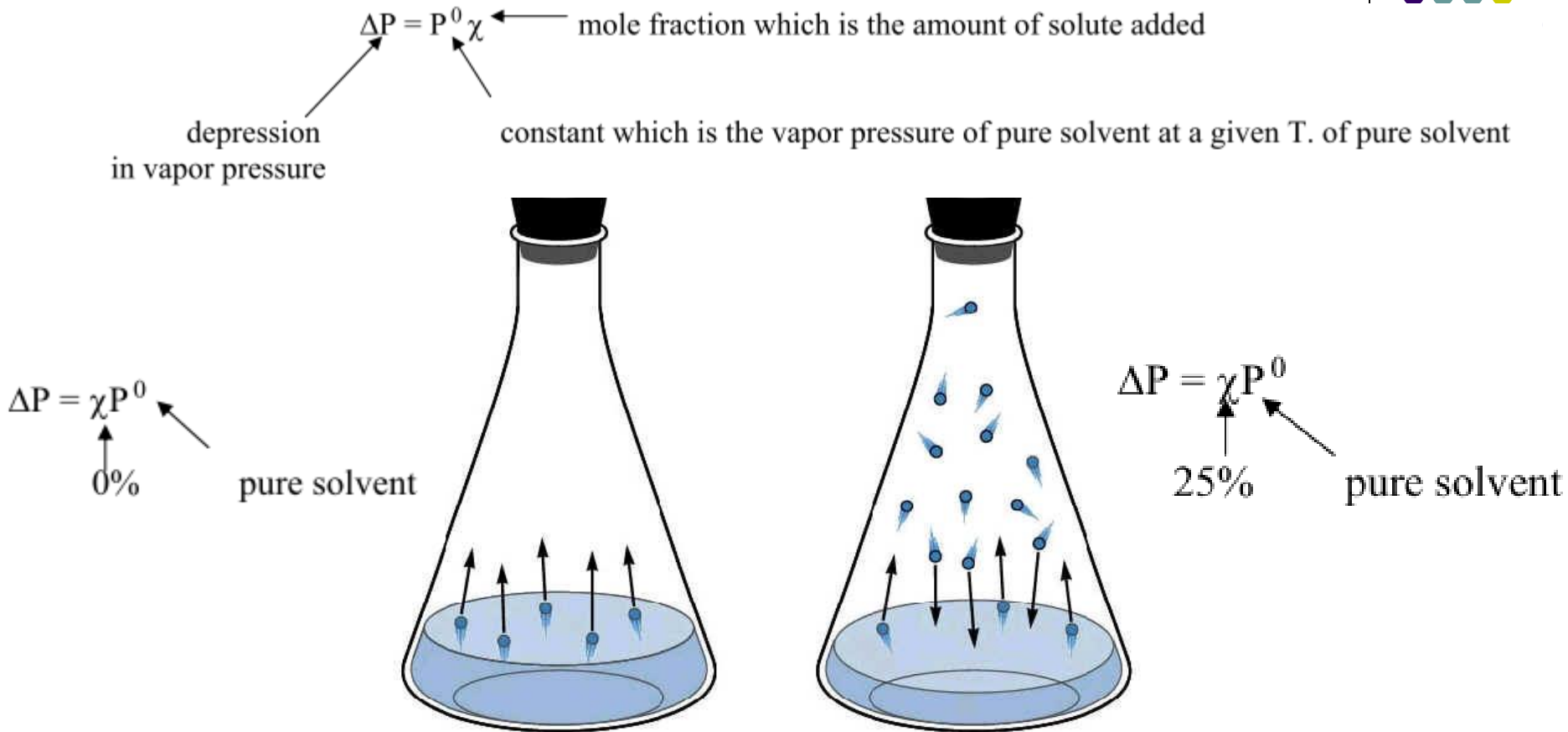
**The Raoult's law:** The relative depression of the saturated vapour pressure of a solvent over a solution equals the mole fraction of the solute.

The mathematical expression of the first Raoult's law is:

$$\frac{P_o - P_s}{P_o} = \frac{n_2}{n_1 + n_2},$$

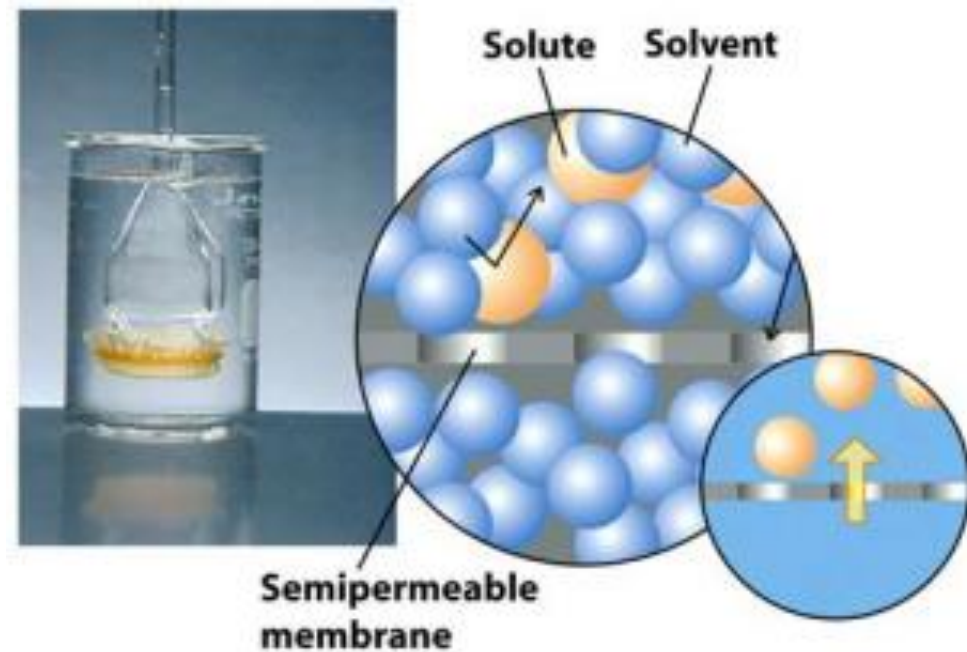
where  $p^o$  – vapour pressure of the solvent,  
 $P_s$  – vapour pressure of the solution,  
 $n_2$  – number of moles of the solute,  
 $n_1$  – number of moles of the solvent,

# Vapour pressure depression



- The particles of solute are surrounded by and attracted to particles of solvent.
- Now the solvent particles have less kinetic energy and tend less to escape into the space above the liquid.
- So the vapor pressure is less.

$$P_{\text{osmotic}} = C_M RT$$



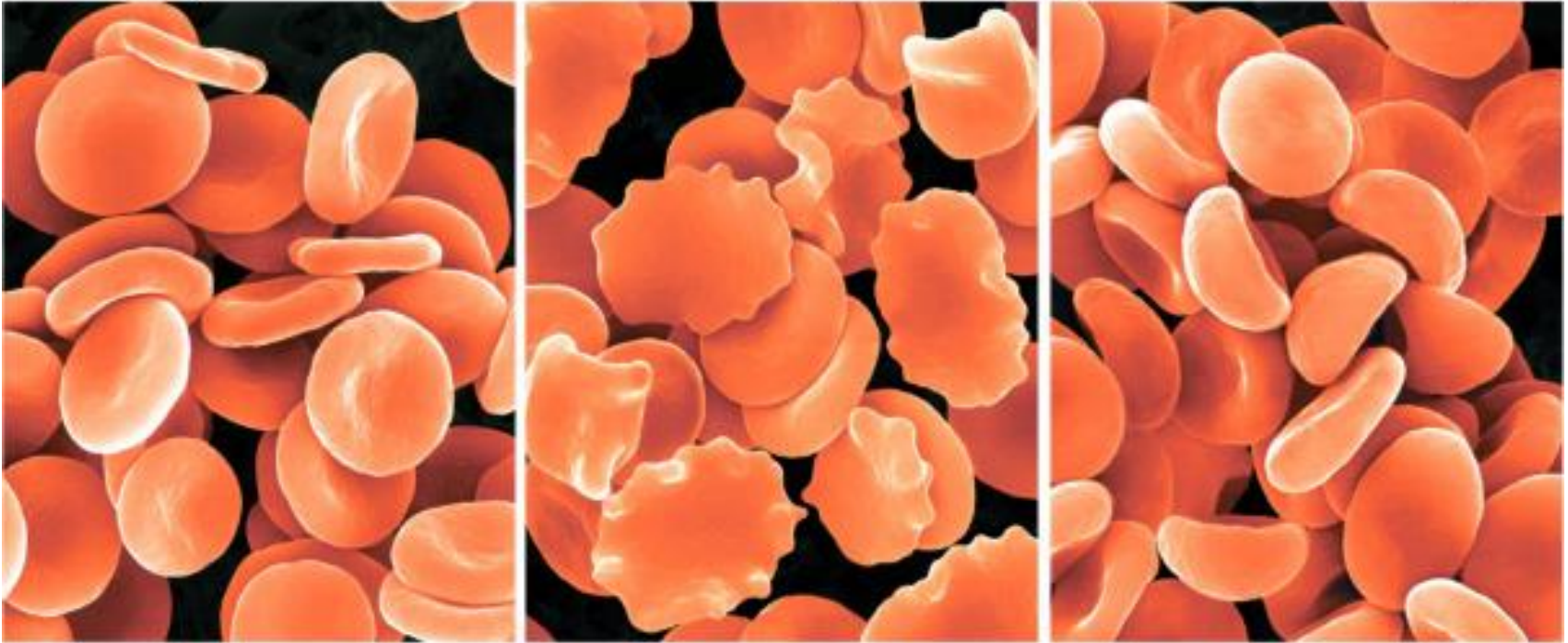
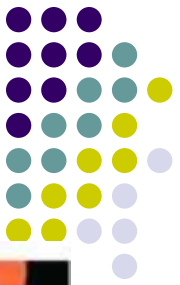
$\pi = MRT$   
Osmotic pressure  
increase because of solute

← molarity which is the amount of solute added  
constant,  $RT$ , which seems to pop up everywhere

The constants  $R$  and  $T$  are the ideal gas law constant and the system temperature

So the osmotic pressure change is  $\pi = MRT = (1.17 \text{ M})(0.082)(298 \text{ K}) = \mathbf{28.6 \text{ atm}}$

# Isotonic, hypotonic and hypertonic solutions



The solution around the cells has the same concentrations like inside of them (first picture), too high concentration in which case the water flows out of the cell and they shrivel up (second picture), or too low concentration in which case water flows into the cells and they lyse or rupture (third picture.)

- **Example** 1.2 L of solution contains 20.5 mg of sucrose ( $C_{12}H_{22}O_{11}$ ). Calculate the osmotic pressure of the solution at 22°C.
- **Solution**



$$M(C_{12}H_{22}O_{11}) = 342 \text{ g / mol}$$

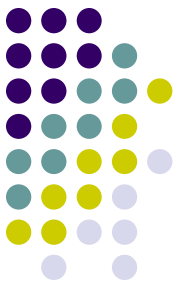
$$C_M = \frac{n(C_{12}H_{22}O_{11})}{V(\text{solution})} = \frac{m(C_{12}H_{22}O_{11})}{M(C_{12}H_{22}O_{11}) \cdot V(\text{solution})} = \frac{20.5 \cdot 10^{-2} \text{ g}}{342 \text{ g / mol} \cdot 1.2 \text{ L}} = 5.0 \cdot 10^{-5} \text{ mol / L}$$

$$P_{\text{osmotic}} = C_M RT = 5.0 \cdot 10^{-5} \text{ mol/L} \cdot 8.314 \text{ J/mol} \cdot \text{K} \cdot 295 \text{ K} = 0.123 \text{ kPa} = 123 \text{ Pa.}$$

Answer:  $P_{\text{osmotic}} = 123 \text{ Pa.}$

# Ionic vs Molecular Solutes

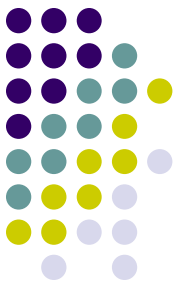
## Electrolytes vs Non electrolytes



- Ionic solutes produce two or more ion particles in solution.
- They affect the colligative properties proportionately more than molecular solutes (that do not ionize).
- The effect is proportional to the **number of particles of the solute in the solution.**



# Freezing Point Depression and Boiling Point Elevation



## Boiling Point Elevation

- $\Delta T_b = k_b C_m$  (for water  $k_b = 0.52 \text{ }^\circ\text{C/m}$ )
- Freezing Point Depression
- $\Delta T_f = k_f C_m$  (for water  $k_f = 1.86 \text{ }^\circ\text{C/m}$ )
- **Note:**  $C_m$  is the molality of the particles, so if the solute is ionic, multiply by the *i-Vant Hoff coefficient* of particles it dissociates to ( $v$ ).
- $i = 1 + \alpha(v - 1)$



# Boiling Point Elevation and Freezing Point Depression



$$\Delta T = K \cdot C_m \cdot i$$

$i$  = van't Hoff factor = number of particles produced per molecule/formula unit. For covalent compounds,  $i = 1$ . For ionic compounds,  $i =$  the number of ions present (both + and -)

<u>Compound</u>	<u>Theoretical Value of <math>i</math></u>
glycol	1
NaCl	2
CaCl <sub>2</sub>	3
Ca <sub>3</sub> (PO <sub>4</sub> ) <sub>2</sub>	5

# Boiling Point Elevation and Freezing Point Depression



$$\Delta T = K \cdot C_m \cdot i$$

**$C_m$  = molality**

**$K$  = molal freezing  
point/boiling point constant**

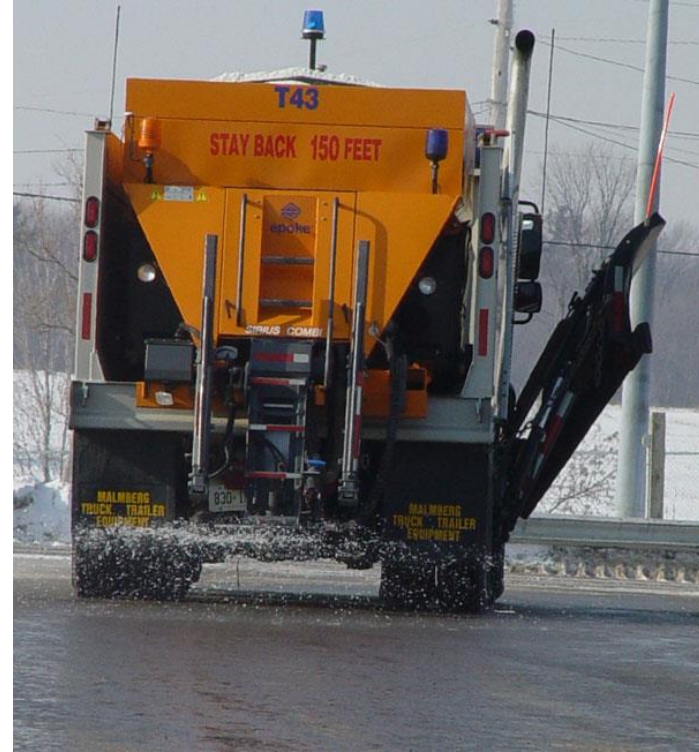
Substance	$K_f$
benzene	5.12
camphor	40.
carbon tetrachloride	30.
ethyl ether	1.79
water	1.86

Substance	$K_b$
benzene	2.53
camphor	5.95
carbon tetrachloride	5.03
ethyl ether	2.02
water	0.52



How many particles do each of the following give upon solvation?

- NaCl
- CaCl<sub>2</sub>
- Glucose



Which is more effective for lowering the freezing point of water?  
NaCl, CaCl<sub>2</sub> or Glucose

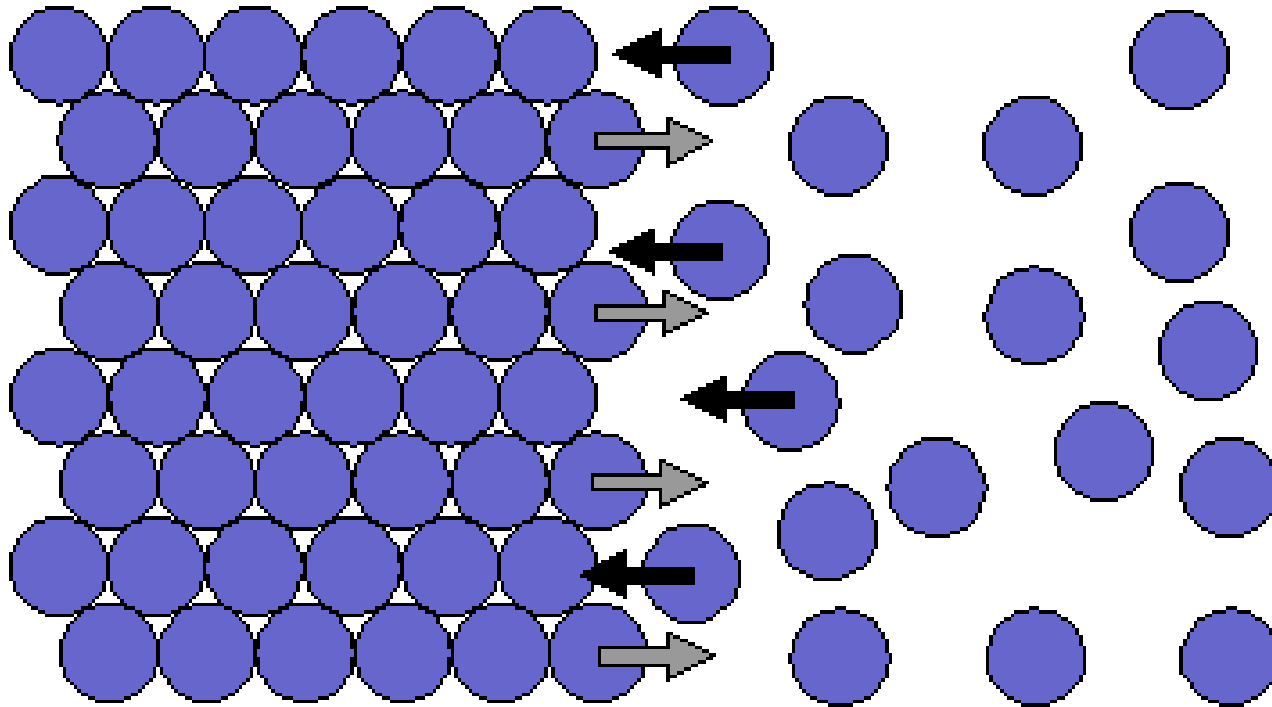
$$\Delta T_f = K_f K \cdot \text{kg/mol} \cdot \frac{m(\text{solute})\text{g} \cdot 1000}{M(\text{solute}) \text{g/mol} \cdot m(\text{H}_2\text{O})\text{g}}$$

$$\Delta T_b = K_b K \cdot \text{kg/mol} \cdot \frac{m(\text{solute})\text{g} \cdot 1000}{M(\text{solute}) \text{g/mol} \cdot m(\text{H}_2\text{O})\text{g}}$$

# Freezing Point Depression



$$\Delta T_f = K_f \cdot Cm \quad \Delta T_f = i \cdot K_f \cdot Cm$$

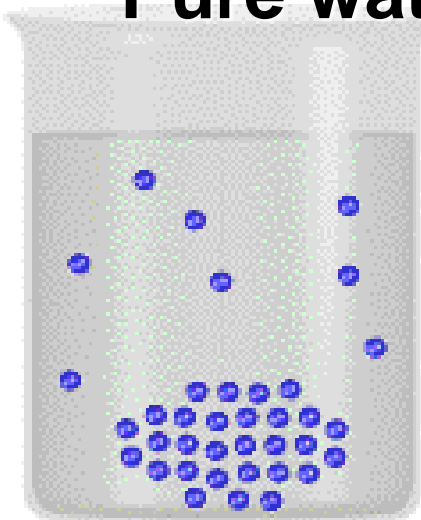


# Change in Freezing Point

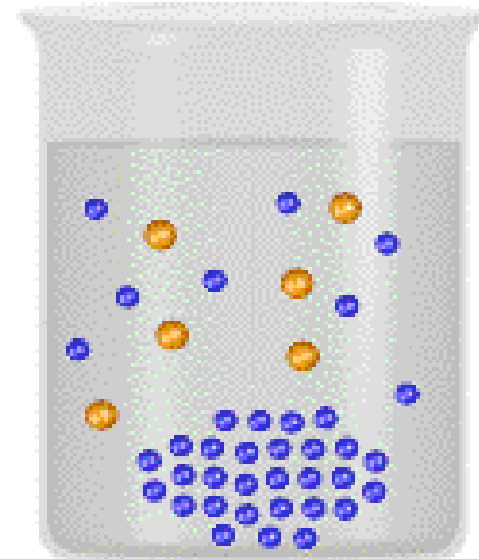


## Ethylene glycol/water solution

### Pure water



Pure water  
(without solute)



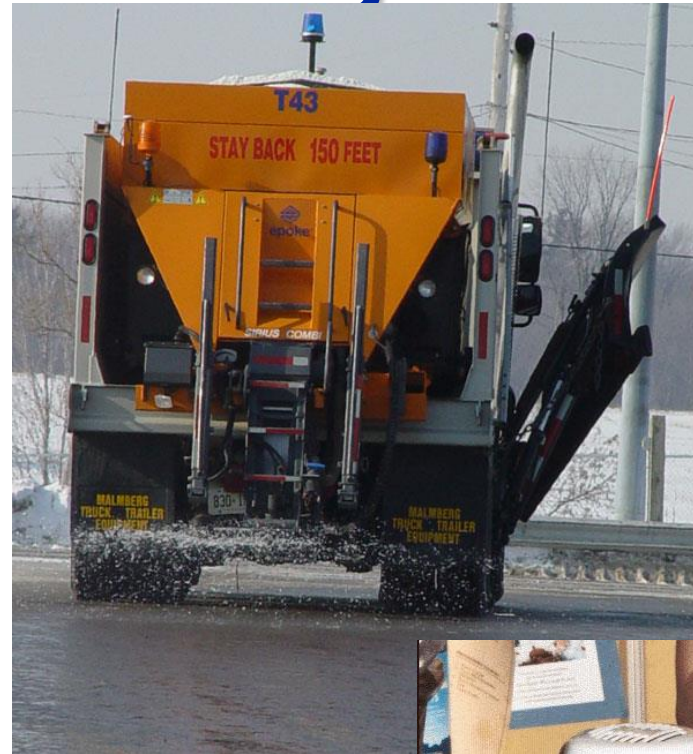
Ethylene glycol  
solution

The freezing point of a solution is **LOWER** than that of the pure solvent

# Change in Freezing Point



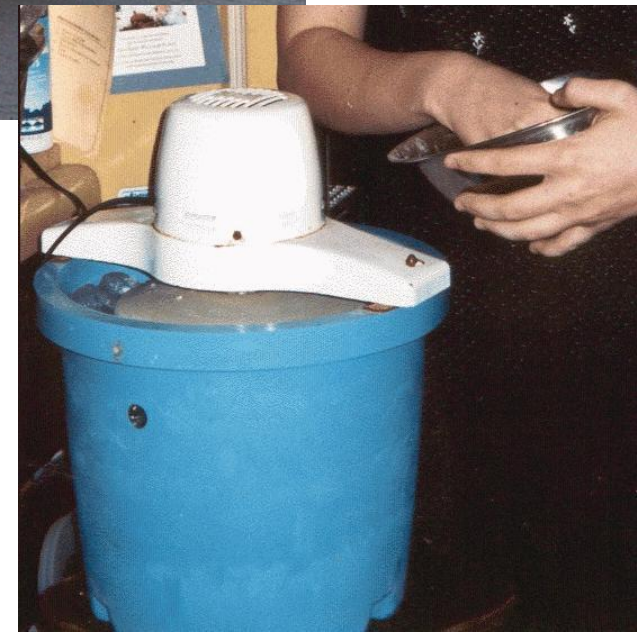
Common Applications  
of Freezing Point  
Depression

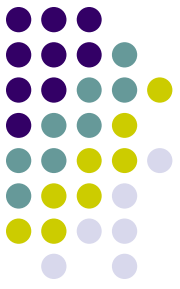


Propylene glycol

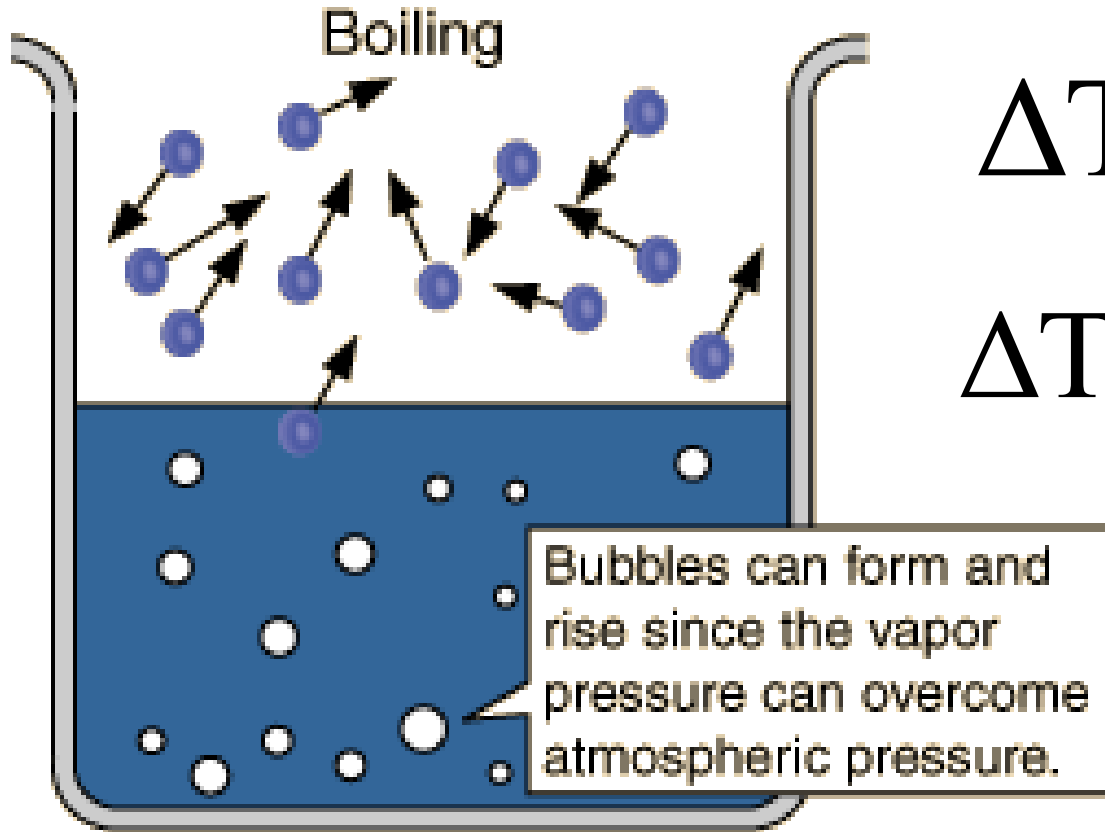


Ethylene glycol –  
deadly to  
small  
animals





# Boiling Point Elevation



$$\Delta T_f = K_f \cdot C_m$$

$$\Delta T_f = i \cdot K_f \cdot C_m$$

# Change in Boiling Point



## Common Applications of Boiling Point Elevation



**-84°** **+276°**  
All Weather Protection

PROTECTION	FREEZE UP PROTECTION	BOILOVER PROTECTION*	CORROSION PROTECTION
MINIMUM: 50% PRESTONE 50% WATER	-34°F	+265°F	EXCEEDS ALL ASTM AND SAE STANDARDS FOR CORROSION PROTECTION
MAXIMUM: 70% PRESTONE 30% WATER	-84°F	+276°F	

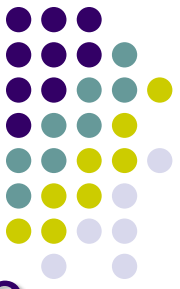
\* USING A 15 LB. PRESSURE CAP

**WARNING** Do not drink antifreeze or solution. If swallowed, give two glasses of water and...





# Change in Boiling Point



Dissolve 62.1 g of glycol (1.00 mol) in 250. g of water. What is the boiling point of the solution?

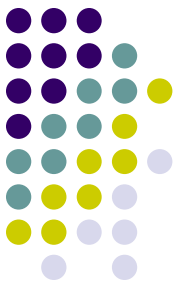
$K_b = 0.52 \text{ }^\circ\text{C/molal}$  for water (see  $K_b$  table).

**Solution**

$$\Delta T_{BP} = K_b \cdot Cm \cdot i$$

1. Calculate solution molality = 4.00 m
2.  $\Delta T_{BP} = K_b \cdot Cm \cdot i$   
 $\Delta T_{BP} = 0.52 \text{ }^\circ\text{C/molal} (4.00 \text{ molal}) (1)$   
 $\Delta T_{BP} = 2.08 \text{ }^\circ\text{C}$   
 $BP = 100 + 2.08 = 102.08 \text{ }^\circ\text{C}$   
(water normally boils at 100)

# Freezing Point Depression



At what temperature will a 5.4 molal solution of NaCl freeze?

## Solution

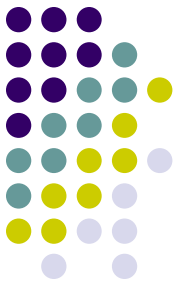
$$\Delta T_{\text{FP}} = K_f \cdot C_m \cdot i$$

$$\Delta T_{\text{FP}} = (1.86 \text{ }^\circ\text{C/molal}) \cdot 5.4 \text{ mol/kg} \cdot 2$$

$$\Delta T_{\text{FP}} = 20.1 \text{ }^\circ\text{C}$$

$$\text{FP} = 0 - 20.1 = -20.1 \text{ }^\circ\text{C}$$

Thank you for your attention!



Think like a  
**Proton**



and stay  
**Positive**