Osmotic Pressure

DR. MIOY T. HUYNH
YALE UNIVERSITY
CHEMISTRY 161
FALL 2019

www.mioy.org/chem161

WHAT (ELSE) HAPPENS WHEN I ADD SOLUTE TO A SOLVENT?

We've already seen the effect of adding solute to a solvent in terms of <u>vapor pressure</u> (decreases with increasing solute) via Raoult's Law.

All of these properties are dependent <u>only</u> on the <u>concentration</u> of solute, and *not* their identity, and are called <u>colligative</u> properties.

WHAT (ELSE) HAPPENS WHEN I ADD SOLUTE TO A SOLVENT?

We've already seen the effect of adding solute to a solvent in terms of <u>vapor pressure</u> (decreases with increasing solute) via Raoult's Law.

We've also seen the effect of adding solute to a solvent in terms of <u>phase changes</u>; e.g. freezing point depression and boiling point elevation.

FREEZING POINT DEPRESSION

$$\Delta T_f = i K_f m$$

BOILING POINT ELEVATION

$$\Delta T_{\rm b} = i K_{\rm b} m$$

All of these properties are dependent <u>only</u> on the <u>concentration</u> of solute, and *not* their identity, and are called <u>colligative</u> properties.

WHAT (ELSE) HAPPENS WHEN I ADD SOLUTE TO A SOLVENT?

We've already seen the effect of adding solute to a solvent in terms of <u>vapor pressure</u> (decreases with increasing solute) via Raoult's Law.

We've also seen the effect of adding solute to a solvent in terms of <u>phase changes</u>; e.g. freezing point depression and boiling point elevation.

FREEZING POINT DEPRESSION

BOILING POINT ELEVATION

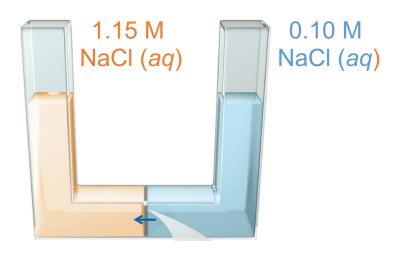
$$\Delta T_f = i K_f m$$

$$\Delta T_{\rm b} = i K_{\rm b} m$$

Lastly, we will consider the effect of adding solute to a solvent in terms of water flow through a semipermeable membrane (osmosis) through the concept of <u>osmotic pressure</u>.

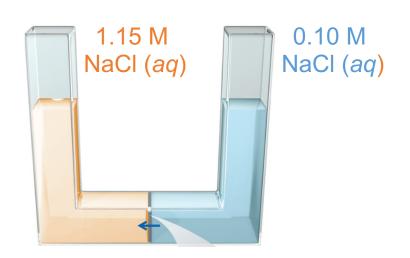
All of these properties are dependent <u>only</u> on the <u>concentration</u> of solute, and *not* their identity, and are called <u>colligative</u> properties.

Consider the following contraption where the left side is an aqueous solution of 1.15 M NaCl and the right side is an aqueous solution of 0.10 M NaCl.



Consider the following contraption where the left side is an aqueous solution of 1.15 M NaCl and the right side is an aqueous solution of 0.10 M NaCl.

Separating these two solutions is a <u>semi-permeable membrane</u>, which allows water to flow between the two sides but not the ions Na⁺ and Cl⁻.



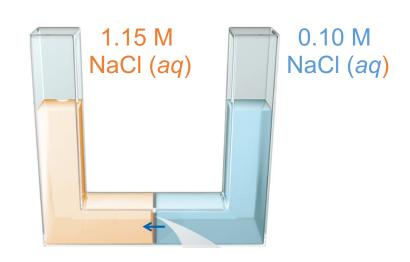
Consider the following contraption where the left side is an aqueous solution of 1.15 M NaCl and the right side is an aqueous solution of 0.10 M NaCl.

Separating these two solutions is a <u>semi-permeable membrane</u>, which allows water to flow between the two sides but not the ions Na⁺ and Cl⁻.

The two solutions each exert a <u>pressure</u> against this membrane, and this pressure depends on the molarity of the solution via:

$$\Pi = iMRT$$

 Π = pressure (atm) i = number of dissociated particles per mole of solute M = molarity (mol/L or M) R = 0.08206 L·atm/mol·K T = temperature (K)



Consider the following contraption where the left side is an aqueous solution of 1.15 M NaCl and the right side is an aqueous solution of 0.10 M NaCl.

Separating these two solutions is a <u>semi-permeable membrane</u>, which allows water to flow between the two sides but not the ions Na⁺ and Cl⁻.

The two solutions each exert a <u>pressure</u> against this membrane, and this pressure depends on the molarity of the solution via:

$$\Pi = iMRT$$

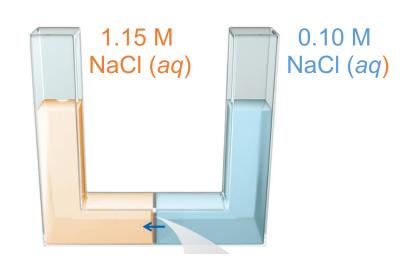
Π = pressure (atm)

i = number of dissociated particles per mole of solute

M = molarity (mol/L or M)

R = 0.08206 L·atm/mol·K

T = temperature (K)



Water always flows to <u>balance</u> the concentrations, so water flows *from* the solution of low concentration ($low \Pi$) to the solution of high concentration ($low \Pi$).

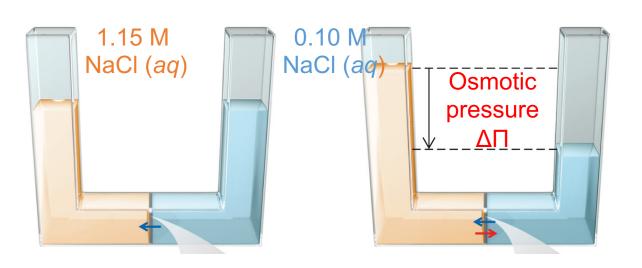
Consider the following contraption where the left side is an aqueous solution of 1.15 M NaCl and the right side is an aqueous solution of 0.10 M NaCl.

Separating these two solutions is a <u>semi-permeable membrane</u>, which allows water to flow between the two sides but not the ions Na⁺ and Cl⁻.

The two solutions each exert a <u>pressure</u> against this membrane, and this pressure depends on the molarity of the solution via:

$$\Pi = iMRT$$

 Π = pressure (atm) i = number of dissociated particles per mole of solute M = molarity (mol/L or M) R = 0.08206 L·atm/mol·K T = temperature (K)



Osmotic pressure ($\Delta\Pi$) is the pressure required to achieve equilibrium (stop water flow), which is equal to the difference in the two pressures.

Water always flows to <u>balance</u> the concentrations, so water flows *from* the solution of low concentration ($low \Pi$) to the solution of high concentration ($low \Pi$).

Solution A: 100. mL of 0.982 M CaCl₂ (strong electrolyte) Solution B: 16 g NaCl (strong electrolyte) in 100. mL water

In which direction will solvent flow across a membrane separating the two?

Your first actual step in these types of problem is to determine what kind of compound you have! Because:

- Molecular compounds that dissolve have an i = 1 since they do not dissociate.
- Insoluble ionic compounds do not dissociate in water, so no changes are observed!
- Soluble ionic compounds have theoretical i values equal to the number of ions per mole compound.

Your first actual step in these types of problem is to determine what kind of compound you have! Because:

- Molecular compounds that dissolve have an i = 1 since they do not dissociate.
- Insoluble ionic compounds do not dissociate in water, so no changes are observed!
- Soluble ionic compounds have theoretical i values equal to the number of ions per mole compound.

For both solutions, we can simply calculate the concentration of ions (*i*M):

Your first actual step in these types of problem is to determine what kind of compound you have! Because:

- Molecular compounds that dissolve have an i = 1 since they do not dissociate.
- Insoluble ionic compounds do not dissociate in water, so no changes are observed!
- Soluble ionic compounds have theoretical i values equal to the number of ions per mole compound.

For both solutions, we can simply calculate the concentration of ions (*i*M):

$$iM_A = (3) \times 0.982 M = 2.95 M$$

Your first actual step in these types of problem is to determine what kind of compound you have! Because:

- Molecular compounds that dissolve have an i = 1 since they do not dissociate.
- Insoluble ionic compounds do not dissociate in water, so no changes are observed!
- Soluble ionic compounds have theoretical i values equal to the number of ions per mole compound.

For both solutions, we can simply calculate the concentration of ions (*i*M):

$$iM_A = (3) \times 0.982 M = 2.95 M$$

$$iM_{\rm B} = (2) \times \frac{16 \text{ g NaCl} \times \frac{1 \text{ mol NaCl}}{58.44 \text{ g}}}{100. \text{ mL} \times \frac{1 \text{ L}}{1000 \text{ mL}}} = 5.5 \text{ M}$$

Your first actual step in these types of problem is to determine what kind of compound you have! Because:

- Molecular compounds that dissolve have an i = 1 since they do not dissociate.
- Insoluble ionic compounds do not dissociate in water, so no changes are observed!
- Soluble ionic compounds have theoretical i values equal to the number of ions per mole compound.

For both solutions, we can simply calculate the concentration of ions (*i*M):

$$iM_A = (3) \times 0.982 M = 2.95 M$$

$$iM_{\rm B} = (2) \times \frac{16 \text{ g NaCl} \times \frac{1 \text{ mol NaCl}}{58.44 \text{ g}}}{100. \text{ mL} \times \frac{1 \text{ L}}{1000 \text{ mL}}} = 5.5 \text{ M}$$

Solution B has a greater concentration of ions, so water will flow from side A to side B.

Your first actual step in these types of problem is to determine what kind of compound you have! Because:

- Molecular compounds that dissolve have an i = 1 since they do not dissociate.
- Insoluble ionic compounds do **not** dissociate in water, so **no changes** are observed!
- Soluble ionic compounds have theoretical i values equal to the number of ions per mole compound.

Working backwards, we can solve for the molarity (M) since we know i = 1:

$$\Pi = iMRT$$
9.94 Torr × $\frac{1 \text{ atm}}{760 \text{ Torr}} = (1)(M) \left(0.08206 \frac{L \cdot \text{atm}}{\text{mol} \cdot \text{K}}\right) (23.6 + 273.1_5 \text{ K})$

$$M = 5.37 \times 10^{-4} \frac{\text{mol}}{L}$$

Finally, we can back-calculate the molar mass from the molarity:

$$M = \frac{n_{\text{solute}}}{V_{\text{solution}}} = \frac{m_{\text{solute}} \times \frac{1}{MM_{\text{solute}}}}{V_{\text{solution}}}$$

Your first actual step in these types of problem is to determine what kind of compound you have! Because:

- Molecular compounds that dissolve have an i = 1 since they do not dissociate.
- Insoluble ionic compounds do **not** dissociate in water, so **no changes** are observed!
- Soluble ionic compounds have theoretical i values equal to the number of ions per mole compound.

Your first actual step in these types of problem is to determine what kind of compound you have! Because:

- Molecular compounds that dissolve have an i = 1 since they do not dissociate.
- Insoluble ionic compounds do **not** dissociate in water, so **no changes** are observed!
- Soluble ionic compounds have theoretical i values equal to the number of ions per mole compound.

Working backwards, we can solve for the molarity (M) since we know i = 1:

$$\Pi = iMRT$$
9.94 Torr × $\frac{1 \text{ atm}}{760 \text{ Torr}} = (1)(M) \left(0.08206 \frac{L \cdot \text{atm}}{\text{mol} \cdot \text{K}}\right) (23.6 + 273.1_5 \text{ K})$

$$M = 5.37 \times 10^{-4} \frac{\text{mol}}{L}$$

Your first actual step in these types of problem is to determine what kind of compound you have! Because:

- Molecular compounds that dissolve have an i = 1 since they do not dissociate.
- Insoluble ionic compounds do **not** dissociate in water, so **no changes** are observed!
- Soluble ionic compounds have theoretical i values equal to the number of ions per mole compound.

Working backwards, we can solve for the molarity (M) since we know i = 1:

$$\Pi = iMRT$$
9.94 Torr × $\frac{1 \text{ atm}}{760 \text{ Torr}} = (1)(M) \left(0.08206 \frac{L \cdot \text{atm}}{\text{mol} \cdot \text{K}}\right) (23.6 + 273.1_5 \text{ K})$

$$M = 5.37 \times 10^{-4} \frac{\text{mol}}{L}$$

Finally, we can back-calculate the molar mass from the molarity:

$$M = \frac{n_{solute}}{V_{solution}} = \frac{m_{solute} \times \frac{1}{MM_{solute}}}{V_{solution}}$$

$$MM_{solute} = \frac{m_{solute}}{M \times V_{solution}} = \frac{0.02740 \text{ g}}{\left(5.37 \times 10^{-4} \frac{\text{mol}}{\text{L}}\right) (0.1000 \text{ L})} = 510. \frac{\text{g}}{\text{mol}}$$