

Chapter 12

Solutions

Sections: 12.1 – 12.8

• Types of Solution

Solution: a homogenous mixture of two or more substances.

المحلول: خليط متجانس من مادتين أو أكثر.

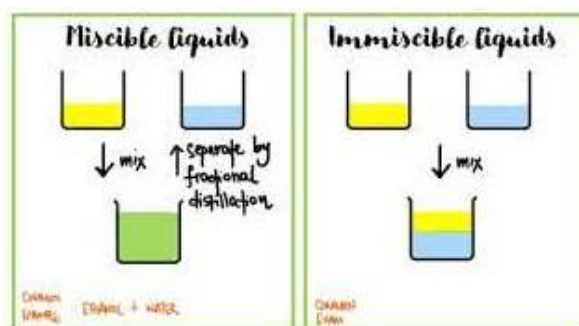
Solute: is(are) the substance(s) present in the smaller amount(s).

المذاب: هو المادة أو المواد الموجودة بكمية أو كميات أصغر.

Solvent: is the substance present in a larger amount.

المذيب: هو المادة الموجودة بكمية أكبر.

Fluids that mix with or dissolve in each other in all proportions are said to be **miscible**.



Fluids that do not dissolve in each other are said to be **immiscible**.

Example 1:

Identify the solute(s) and solvent(s) in the following solutions.

- 70 g of Cr and 12 g of Mo
- 5 g of MgCl_2 dissolved in 1500 g of H_2O
- 39% N_2 , 41% Air, and the rest O_2

Answer:

- The 12 g of Mo is the solute; the 70 g of Cr is the solvent.
- MgCl_2 is the solute; H_2O is the solvent.
- O_2 and N_2 are the solutes; Ar is the solvent.

• Solubility and the Solution Process

- **saturated solution:** contains the maximum amount of a solute that will dissolve in a given solvent at a specific temperature.
 - **المحلول المشبع:** يحتوي على أكبر كمية من المادة المذابة التي تذوب في مذيب معين عند درجة حرارة معينة.
- **unsaturated solution:** contains less solute than the solvent can dissolve at a specific temperature.
 - **المحلول غير المشبع:** يحتوي على كمية من المذاب أقل مما يمكن للمذيب أن يذوب عند درجة حرارة معينة.
- **supersaturated solution:** contains more solute than is present in a saturated solution at a specific temperature.
 - **المحلول فوق المشبع:** يحتوي على نسبة مذابة أكبر من الموجودة في المحلول المشبع عند درجة حرارة معينة.
- **Solubility:** the amount of solute that dissolves in a given quantity of solvent at a given temperature.
 - **الذائبية:** كمية المذاب التي تذوب في كمية معينة من المذيب عند درجة حرارة معينة.

For example, The **solubility** of sodium chloride in water is **36.0 g/100 mL at 20°C**.

Three types of interactions in the solution process:

- **solvent-solvent** interaction
- **solute-solute** interaction
- **solvent-solute** interaction

$$\Delta H_{\text{solution}} = \Delta H_1 + \Delta H_2 + \Delta H_3$$

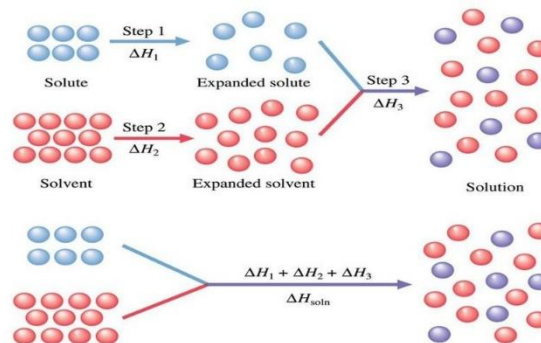
If $\Delta H_{\text{solution}} > 0$ لا يحدث التفاعل ولا يتكون محلول

If $\Delta H_{\text{solution}} < 0$ يحدث التفاعل ويتكون محلول

If $\Delta H_{\text{solution}} = 0$ محلول مثالي

If $\Delta H_{\text{solution}}$ small and positive يحدث التفاعل ويتكون محلول

If $\Delta H_{\text{solution}}$ big and positive لا يحدث التفاعل ولا يتكون محلول



like dissolves like:

Two substances with similar **intermolecular** forces are likely to be soluble in each other.

- **non-polar** molecules are soluble in **non-polar** solvents, ex: CCl_4 in C_6H_6
- **polar** molecules are soluble in **polar** solvents, ex: $\text{C}_2\text{H}_5\text{OH}$ in H_2O
- **ionic** compounds are more soluble in **polar** solvents, ex: NaCl in H_2O or NH_3

Miscible liquids; Similar polarity ex: $\text{CH}_3\text{CH}_2\text{OH}$ and water (H_2O)

Immiscible Liquids;

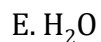
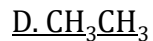
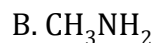
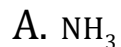
- Two insoluble liquids; Do not mix; Get two separate phases.
- Different polarity

ex: C_3H_8 and H_2O

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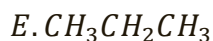
Example 2:

Which of the following molecules is most soluble in hexane, C_6H_{14} ?



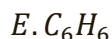
Example 3:

Which of the following compounds is likely to be **more soluble** in water?



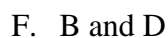
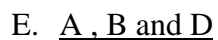
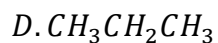
Example 4:

Which of the following compounds is likely to be more soluble in KCl ?



Example 5:

Which of the following compounds is likely to be **soluble** in Br_2 ?



• Ways of Expressing Concentration

Concentration: the amount of solute present in a given quantity of solvent or solution.

$$\text{number of moles (mol)} = \frac{\text{mass (Kg)}}{\text{molar mass } \left(\frac{\text{Kg}}{\text{mol}}\right)} = n = \frac{m}{M_w}$$

$$\text{Density} = \frac{\text{mass}}{\text{volume}} = \frac{m}{V}$$

$$\text{solution} = \text{solute} + \text{solvent}$$

$$\text{Mass fraction} = \frac{\text{mass of solute}}{\text{mass of solution}} = \frac{\text{mass of solute}}{\text{mass of solute} + \text{mass of solvent}}$$

$$\text{Percentage by mass} = \frac{\text{mass of solute}}{\text{mass of solution}} \times 100\%$$

$$\text{Mole fraction} = \frac{\text{mol of solute}}{\text{mol of solution}} = \frac{\text{mol of solute}}{\text{mol of solute} + \text{mol of solvent}}$$

$$\text{Mole percent} = \frac{\text{mol of solute}}{\text{mol of solution}} \times 100\%$$

• Molarity (concentration) (mol/L)

$$\text{Molarity (M)} = \frac{\text{moles of solute (mol)}}{\text{volume of solution (L)}}$$

• Molality (mol/Kg)

$$\text{Molality (m)} = \frac{\text{moles of solute}}{\text{mass of solvent}}$$

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| | |
|---------------------------|---|
| Molarity | $M = \frac{\text{moles of solute}}{\text{liters of solution}}$ |
| Percentage by mass | Percentage by mass = $\frac{\text{mass of solute}}{\text{mass of solution}} \times 100\%$ |
| Mass fraction | Mass fraction = $\frac{\text{mass of solute}}{\text{mass of solution}}$ |
| Molality | $m = \frac{\text{mol of solute}}{\text{kg of solvent}}$ |
| Mole fraction | $X_A = \frac{n_A}{n_A + n_B + n_C + n_D + \cdots + n_Z}$ |
| Mole percent | mol percent $A = \frac{n_A}{n_A + n_B + n_C + n_D + \cdots + n_Z} \times 100 \text{ mol}\%$ |
| Percentage by mass–volume | $\%(\text{w/v}) = \frac{\text{grams solute}}{\text{mL solution}} \times 100\%$ |

Example 6:

How many grams of hydrogen chloride (HCl) are required to prepare 4 liters of 5M HCl in water?

The molar mass of HCl is 36.5 g/mol.

Solution:

$$\text{Molarity (M)} = \frac{\text{moles of solute (mol)}}{\text{volume of solution (L)}}$$

$$n = M \times V = 5 \times 4 = 20 \text{ mol}$$

$$\text{mass} = n \times M_w = 20 \times 36.5 = 730 \text{ g}$$

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Example 7:

The density of a solution prepared by dissolving 120 g of urea ($M_w = 60 \text{ g/mol}$) in 1000 g of water is 1.15 g/mL. The molarity of urea is:

- a) 2.05 M
- b) 1.02 M
- c) 0.50 M
- d) 1.78 M

Solution:

$$\text{Total mass} = m_{\text{solute}} + m_{\text{solvent}} = 120 + 1000 = 1120 \text{ g}$$

$$V = \frac{m_{\text{Total}}}{\text{density}} = \frac{1120}{1.15} = 973.91 \text{ ml} = 0.97391 \text{ L}$$

$$n_{\text{urea}} = \frac{m}{M_w} = \frac{120}{60} = 2 \text{ mol}$$

$$\text{Molarity} = \frac{n}{V} = \frac{2}{0.97391} = \mathbf{2.05 \text{ M}}$$

Example 8:

Calculate the molality of a sulfuric acid solution containing 30.4 g of sulfuric acid in 193 g of water. The molar mass of sulfuric acid is 98.09 g/mol.

Solution:

$$\text{Moles of } H_2SO_4 = \frac{\text{mass of } H_2SO_4}{M_w \text{ of } H_2SO_4} = \frac{30.4}{98.09} = 0.31 \text{ mol}$$

$$\text{Molality (m)} = \frac{\text{moles of solute (} H_2SO_4 \text{)}}{\text{mass of solvent (water)}} = \frac{0.31 \text{ mol}}{0.193 \text{ Kg}}$$

$$= \mathbf{1.6 \text{ m}}$$

Example 9:

What is the molarity of the 42.6%(w/v) solution of sodium hydroxide (40g/mole) ?

- A. 18 M
 B. 11 M
 C. 9.7 M
 D. 1.065 M
 E. 101.65 M

Solution:

Let volume of solution = 100 ml

$$\text{mass of sodium hydroxide} = \frac{42.6 \text{ g}}{100 \text{ ml}} \times 100 \text{ ml} = 42.6 \text{ g}$$

$$n = \frac{m}{M_w} = \frac{42.6}{40} = 1.065 \text{ mol}$$

$$\text{Molarity} = \frac{n}{V} = \frac{1.065}{0.1} = \mathbf{10.65M}$$

Example 10:

The density of a 2.45 M aqueous solution of methanol (CH₃OH) is 0.976 g/mL.

What is the molality of the solution?

The molar mass of methanol is 32.04 g/mol.

Solution:

Let volume of solution = 1 L

$$\text{mass of solution} = \text{density} \times V_{\text{solution}} = 0.976 \frac{\text{g}}{\text{ml}} \times 1000 \frac{\text{ml}}{\text{L}} \times 1 \text{ L} = 976 \text{ g}$$

$$\text{moles of solute (CH}_3\text{OH)} = 2.45 \text{ M} \times 1 \text{ L} = 2.45 \text{ mol}$$

$$\text{mass of solvent (H}_2\text{O)} = \text{mass of solution} - \text{mass of solute}$$

$$= 976 - \left(2.45 \text{ mol} \times 32.04 \frac{\text{g}}{\text{mol}} \right) = 976 - 78.5 = 897.5 \text{ g} = 0.8975 \text{ Kg}$$

$$\text{Molality (m)} = \frac{\text{moles of solute (CH}_3\text{OH)}}{\text{mass of solvent (H}_2\text{O)}} = \frac{2.45 \text{ mol}}{0.8975 \text{ Kg}} = \mathbf{2.73 \text{ m}}$$

Example 11:

Calculate the molality of a 40.4 percent (by mass) aqueous solution of phosphoric acid (H_3PO_4).

The molar mass of phosphoric acid is 97.99 g/mole.

Solution:

$$\text{Molality (m)} = \frac{\text{moles of solute (H}_3\text{PO}_4\text{)}}{\text{mass of solvent (water)}}$$

Let mass of solution = 100 g

$$\text{moles of H}_3\text{PO}_4 = \frac{\text{mass}}{\text{molar mass}} = \frac{40.4 \text{ g}}{97.99 \text{ g/mol}} = 0.41 \text{ mol}$$

mass of solvent = 100 – 40.4 = 59.6 g = 0.0596 Kg

$$\text{Molality} = \frac{0.41}{0.0596} = 6.88 \text{ m}$$

Example 12:

What is the concentration of the 45%(w/w) solution of sodium hydroxide (40g/mole) if its density is 1.629 g/mL?

- A. 18 M C. 1.9 M
B. 1.25 M D. 0.76 M

Solution:

Let mass of solution = 100 g

$$n = \frac{m}{M_w} = \frac{45}{40} = 1.125 \text{ mol}$$

$$V = \frac{m}{\text{density}} = \frac{100}{1.629} = 61.39 \text{ ml} = 0.06139 \text{ L}$$

$$\text{Molrity} = \frac{n}{V} = \frac{1.125}{0.06139} = \mathbf{18.3 \text{ M}}$$

Example 13:

What **mass fraction** and mass **percentage by mass** of NH_4Cl (53.49g/mole) must be dissolved in 100 mL of H_2O to make a 0.250 M solution?

The density of the solution of NH_4Cl is 55000 g/m^3 .

Solution:

$$\text{Molarity} = \frac{\text{moles of } \text{NH}_4\text{Cl}}{\text{volume of solution}} \rightarrow n_{\text{NH}_4\text{Cl}} = V \times M = 0.1 \times 0.250 = 0.025 \text{ mol}$$

$$\text{mass of } \text{NH}_4\text{Cl} = n \times Mw = 0.025 \times 53.49 = 1.34 \text{ g}$$

$$\text{mass of solution} = \text{density} \times \text{volume} = 55000 \frac{\text{g}}{\text{m}^3} \times \frac{1 \text{ m}^3}{1000 \text{ L}} \times 0.1 \text{ L} = 5.5 \text{ g}$$

$$\text{mass fraction of } \text{NH}_4\text{Cl} = \frac{\text{mass of } \text{NH}_4\text{Cl}}{\text{mass of solution}} = \frac{1.34}{5.5} = \mathbf{0.24}$$

$$\text{percentage by mass of } \text{NH}_4\text{Cl} = \frac{\text{mass of } \text{NH}_4\text{Cl}}{\text{mass of solution}} \times 100\% = \mathbf{24\%}$$

Example 14:

What **mole fraction** and mass **percentage by mole** of NH_4Cl (53.49g/mole) must be dissolved in 100 mL of H_2O to make a 0.250 M solution?

The density of the solution of NH_4Cl is 55000 g/m^3 .

Solution:

$$\text{Molarity} = \frac{\text{moles of } \text{NH}_4\text{Cl}}{\text{volume of solution}} \rightarrow n_{\text{NH}_4\text{Cl}} = V \times M = 0.1 \times 0.250 = 0.025 \text{ mol}$$

$$\text{mass of solution} = \text{density} \times \text{volume} = 55000 \frac{\text{g}}{\text{m}^3} \times \frac{1 \text{ m}^3}{1000 \text{ L}} \times 0.1 \text{ L} = 5.5 \text{ g}$$

$$\text{mass of } \text{NH}_4\text{Cl} = n \times Mw = 0.025 \times 53.49 = 1.34 \text{ g}$$

$$\text{mass of } \text{H}_2\text{O} = 5.5 - 1.34 = 4.16 \text{ g}, \quad n_{\text{H}_2\text{O}} = \frac{4.16}{18} = 0.23 \text{ mol}$$

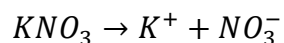
$$\text{mole fraction of } \text{NH}_4\text{Cl} = X_{\text{NH}_4\text{Cl}} = \frac{\text{moles of } \text{NH}_4\text{Cl}}{\text{moles of solution}} = \frac{0.025}{0.025 + 0.23} = \mathbf{0.098}$$

$$\text{mole percent of } \text{NH}_4\text{Cl} = \frac{\text{moles of } \text{NH}_4\text{Cl}}{\text{moles of solution}} \times 100\% = \mathbf{9.8\%}$$

- **the Stoichiometric Factors**

Example 15:

What are the concentrations of potassium and nitrate ions in a 0.025 M solution of KNO_3 ? The dissociation reaction for potassium nitrate (KNO_3) is:

**Solution:**

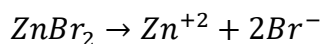
Concentration of K^+ : $1 \times 0.025 M = 0.025 M$

Concentration of NO_3^- : $1 \times 0.025 M = 0.025 M$

Example 16:

What are the concentrations of zinc and bromide ions in a 0.030 M solution of $ZnBr_2$?

The dissociation reaction for zinc bromide ($ZnBr_2$) is:

**Solution:**

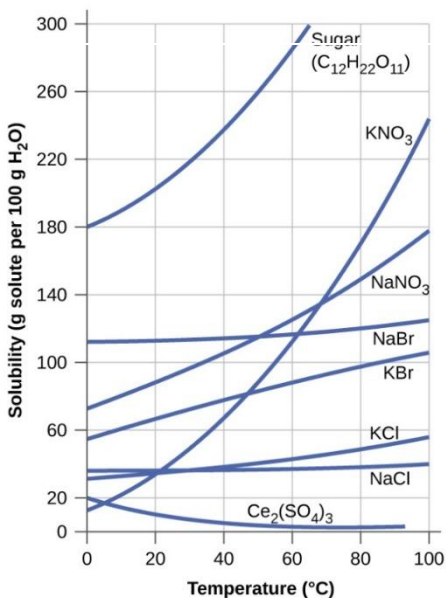
$$[Zn^{+2}] = 1 \times 0.030 M = 0.030 M$$

$$[Br^-] = 2 \times 0.030 M = 0.060 M$$

• **Effects of Temperature and Pressure on Solubility**

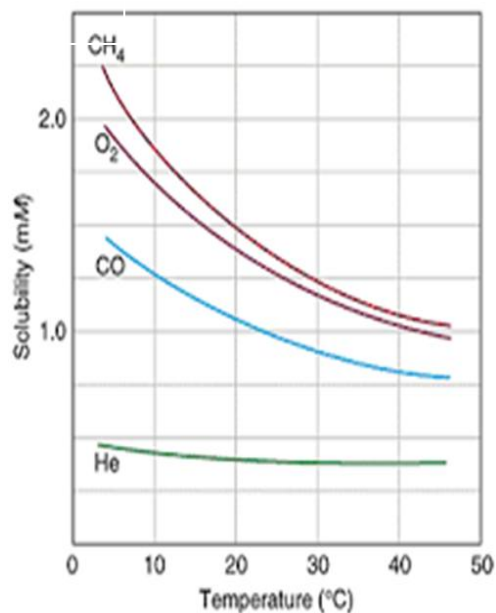
• **Effect of temperature on Solubility of Gases and solids**

• **Solid** solubility and temperature



The Solubility in most solids **increases** with temperature.

• **Gas** solubility and temperature



The Solubility in most gases **decreases** with temperature.

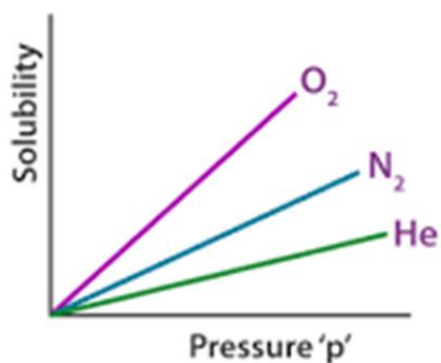
- **Effect of Pressure on Solubility of Gases**

Henry`s law: The solubility of a gas in a liquid is proportional to the pressure of the gas over the solution

$$C = KP$$

$$\text{, or } \frac{C_1}{C_2} = \frac{P_1}{P_2}$$

- $C (S)$: is the concentration (M) of the dissolved gas.
- P : is the pressure of the gas over the solution
- K : is a constant for each gas ($mol/L \cdot atm$) that depends only on temperature.



The Solubility in gases **increases** with pressure.

Example 17:

At 25°C 24 grams of CH_4 (16 g/mol) are dissolved in 37.5 L water at a partial pressure of 2.0 atm. Calculate the solubility of methane in water when the partial pressure is decreased to 0.5 atm at the same temperature.

Solution:

$$\text{Moles of } CH_4 = \frac{\text{mass}}{\text{molar mass}} = \frac{24}{16} = 1.5 \text{ mol}$$

$$[CH_4](\text{solubility}) = \frac{1.5 \text{ mol}}{37.5 \text{ L}} = 0.04 \text{ M (mol/L)}$$

$$S_1 = KP_1 \rightarrow K = \frac{S}{P_1} = \frac{0.04}{2} = 0.02 \frac{\text{mol}}{\text{L}} \cdot \text{atm}$$

$$S_2 = KP_2 = 0.02 \times 0.5 = \mathbf{0.01 \text{ mol/L}}$$

Example 18:

The solubility of nitrogen gas (N_2) in water at 25°C is found to be 3.2×10^{-4} mol/L when the partial pressure of nitrogen gas above the water is 1.5 atm. Calculate the solubility of nitrogen gas in water when the partial pressure is increased to 3.0 atm at the same temperature.

- A. 640 M
- B. 6.4×10^{-4} L/mol
- C. 6.4×10^{-4} M
- D. 6.9×10^{-4} M

Solution:

$$\frac{S_1}{S_2} = \frac{P_1}{P_2} \text{ or } S = kP \quad \rightarrow \quad S_2 = S_1 \frac{P_2}{P_1} = 3.2 \times 10^{-4} \times \frac{3}{1.5} = \mathbf{6.4 \times 10^{-4} \text{ mol/L}}$$

• Colligative Properties

Colligative properties: are properties that depend only on the number of solute particles in solution and not on the nature of the solute particles.

الخواص التي تعتمد فقط على عدد جزيئات المذاب في المحلول وليس على طبيعة جزيئات المذاب.

Amount of solute \uparrow \rightarrow *Vapor pressure* \downarrow

- Vapor-Pressure Lowering
- Boiling-Point Elevation
- Freezing-Point Depression
- Osmotic Pressure (π)

• Vapor-Pressure Lowering

1. When a **non-volatile**, insoluble solid dissolves in a given solution, then we use **Raoult`s law**.

عند اذابة مادة صلبة غير متطايرة وغير قابلة للتفكك لمحلول معين.

Raoult`s law:

$$P = X_1 P^\circ$$
$$\Delta P = P^\circ - P$$

Example 19:

Calculate the **vapor pressure** of a solution formed by dissolving 25.6 g of sucrose (molar mass = 342.3 g/mol) in 200 mL of ethanol (molar mass = 46 g/mol) at 40°C. What is the **vapor-pressure lowering**?

The vapor pressure of pure ethanol at 40°C is 100 mmHg. Assume the density of ethanol is 0.789 g/mL.

Solution:

$$n_2 (\text{sucrose}) = \frac{25.6}{342.3} = 0.075 \text{ mol}$$

$$n_1 (\text{ethanol}) = 0.789 \frac{\text{g}}{\text{ml}} \times 200 \text{ ml} \times \frac{1}{46 \frac{\text{g}}{\text{mol}}} = 3.43 \text{ mol}$$

$$X_1 = \frac{n_1}{n_1 + n_2} = \frac{3.34}{3.34 + 0.075} = 0.978$$

$$P = X_1 * P^\circ = 0.978 * 100 = \mathbf{97.8 \text{ mmHg}}$$

$$\Delta P = P^\circ - P = 100 - 97.8 = \mathbf{2.2 \text{ mmHg}}$$

- Vapor-Pressure Lowering

For nonvolatile solute: - $P < P^\circ$

2. If there are two liquids (A and B) **completely miscible** and both are **volatile**.

2. في حال وجود سائلين (A , B) ذائبين ببعضهما البعض وكلاهما متطاير.

$$P_A = X_A * P_A^\circ \quad , \quad P_B = X_B * P_B^\circ$$

$$P_T = P_A + P_B$$

$$P_T = X_A * P_A^\circ + X_B * P_B^\circ \quad X_A + X_B = 1$$

so we can use also: $P_T = (1 - X_B) * P_A^\circ + X_B * P_B^\circ$

Example 20:

At 40°C, the vapor pressure of pure acetone (molar mass = 58.08 g/mol) is 365.5 torr, while the vapor pressure of pure ethanol (molar mass = 46.07 g/mol) is 92.4 torr. If a solution is prepared by mixing 50.0 g of acetone and 30.0 g of ethanol at this temperature, what would be the total vapor pressure of the mixture?

Solution:

$$n_1 (\text{acetone}) = \frac{50 \text{ g}}{58.08 \text{ g/mol}} = 0.86 \text{ mol} \quad , \quad n_2 (\text{ethanol}) = \frac{30 \text{ g}}{46.07 \text{ g/mol}} = 0.65 \text{ mol}$$

$$X_1 = \frac{n_1}{n_1 + n_2} = \frac{0.86}{0.86 + 0.65} = 0.57$$

$$X_2 = \frac{n_2}{n_1 + n_2} = \frac{0.65}{0.86 + 0.65} = 0.43$$

$$P_1 = X_1 * P_1^\circ = 0.57 * 365.5 = 208.3 \text{ torr} \quad , \quad P_2 = X_2 * P_2^\circ = 0.43 * 92.5 = 39.8 \text{ torr}$$

$$P_T = P_1 + P_2 = 208.3 + 39.8 = \mathbf{248.1 \text{ torr}}$$

Or, $P_T = X_A * P_A^\circ + X_B * P_B^\circ = 0.57 * 365.5 + 0.43 * 92.5 = \mathbf{248.1 \text{ torr}}$

• Vapor Pressure of a Solution

- Vapor pressure for a **nonideal** solution.

❖ تعمل معادلة **(Raoult's law)** بشكل جيد مع المحلول المثالي **(ideal)**: المذيب والمذاب متساويان تمامًا نفس التفاعلات بين الجزيئات **(Intermolecular interaction)**

- Real solutions are not ideal, deviation from this equation is **expected**.

- Solution showing positive deviation from Raoult's law:

- $P_{\text{solution}} > P_{\text{ideal}}$
- $V_{\text{solution}} > V_{\text{solute}} + V_{\text{solvent}}$
-

Ex: London + H-bonding → London

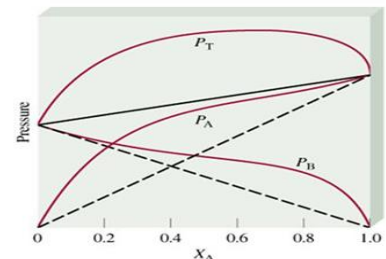
- Solution showing negative deviation from Raoult's law:

- $P_{\text{solution}} < P_{\text{ideal}}$
- $V_{\text{solution}} < V_{\text{solute}} + V_{\text{solvent}}$
-

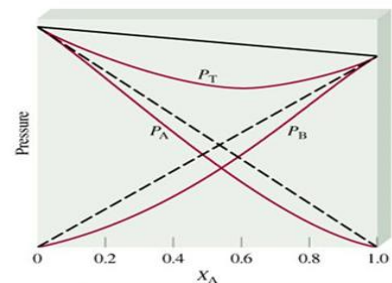
Ex: Dipole-Dipole + Dipole-Dipole → H-bonding

Non ideal solution

- ❖ For solution showing positive deviation from Raoult's law.
 $\Delta H > 0$ and $\Delta V > 0$
 $P_s > P_{\text{ideal}}$



- ❖ For solution showing negative deviation from Raoult's law.
 $\Delta H < 0$ and $\Delta V < 0$
 $P_s < P_{\text{ideal}}$



• Boiling-Point Elevation and Freezing-Point Depression

• Boiling point elevation (الارتفاع في درجة الغليان)

❖ Amount of solute ↑ → boiling point ↑

❖ كل ما زادت كمية المذاب زادت درجة الغليان

$$\Delta T_b = K_b * m$$

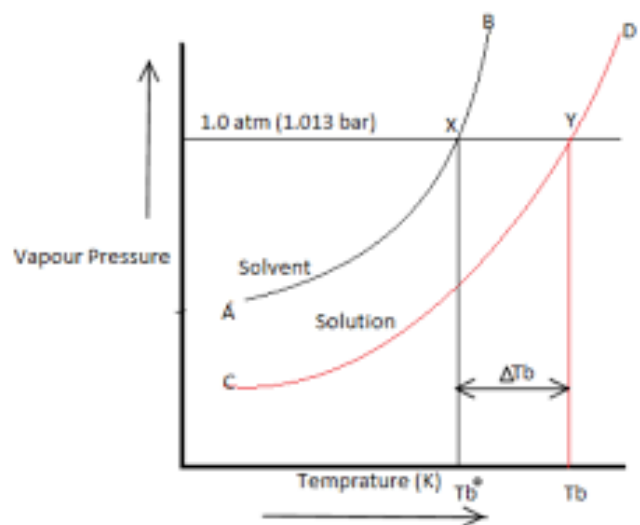
$$\Delta T_b = T_b - T_b^\circ \quad , \quad T_b > T_b^\circ$$

T_b : boiling point of solution

T_b° : boiling point of pure solvent

m : molality of the solution (mol/Kg)

K_b : molal boiling-point elevation constant for a given solvent ($^\circ\text{C}/m$)



• **Freezing point depression (الانخفاض في درجة التجمد)**

•

❖ Amount of solute ↑ → freezing point ↓

❖ كل ما زادت كمية المذاب تقل درجة التجمد

$$\Delta T_f = K_f * m$$

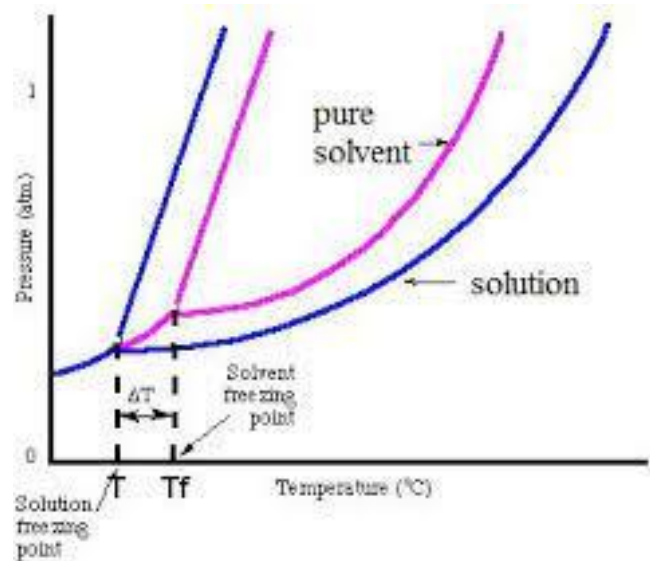
$$\Delta T_f = T_f^\circ - T_f \quad , \quad T_f < T_f^\circ$$

T_f : freezing point of solution

T_f° : freezing point of pure solvent

m : molality of the solution (mol/Kg)

K_f : molal freezing point depression constant for a given solvent ($^\circ\text{C}/m$)



Example 21:

Propylene glycol ($C_3H_8O_2$) ($M_w = 76.09 \text{ g/mol}$) is another common antifreeze used in automobile cooling systems. It is also water-soluble and has a boiling point of 188.2°C . Calculate the freezing point and boiling point of a solution containing 800 g of propylene glycol in 1500 g of water.

$$K_b = 3.55 \text{ }^\circ\text{C/m} \quad , \quad K_f = 1.86 \text{ }^\circ\text{C/m}$$

Solution:

$$n_{C_3H_8O_2} = \frac{800 \text{ g}}{\frac{76.09 \text{ g}}{\text{mol}}} = 10.5 \text{ mol}$$

$$\text{molality (m)} = \frac{\text{mole of solute}}{\text{mass of solvent (Kg)}} = \frac{10.5 \text{ mol}}{1.5 \text{ Kg}} = 7$$

$$\Delta T_b = K_b * m = 3.55 * 7 = 24.85 \text{ }^\circ\text{C}$$

$$\Delta T_b = T_b - T_b^\circ \rightarrow T_b = \Delta T_b + T_b^\circ = 24.85 + 100 = \mathbf{124.85 \text{ }^\circ\text{C}}$$

$$\Delta T_f = K_f * m = 1.86 * 7 = 13 \text{ }^\circ\text{C}$$

$$\Delta T_f = T_f^\circ - T_f \rightarrow T_f = T_f^\circ - \Delta T_f = 0 - 13 = \mathbf{-13 \text{ }^\circ\text{C}}$$

Example 22:

A 6.25 g sample of a compound with the empirical formula C_3H_3 is dissolved in 250 g of toluene. The freezing point of the solution is found to be 2.3°C below that of pure toluene. Calculate the **molar mass** of the compound.

$$K_f = 3.6 \text{ }^\circ\text{C/m}$$

Solution:

freezing point depression \rightarrow *molality* \rightarrow *number of moles* \rightarrow *molar mass*

$$\Delta T_f = K_f * m \rightarrow m = \frac{\Delta T_f}{K_f} = \frac{2.3}{3.6} = 0.64 \text{ m}$$

$$m = \frac{\text{mole of solute}}{\text{mass of solvent (Kg)}} \rightarrow n_{C_3H_3} = 0.64 \times \frac{\text{mol}}{\text{Kg}} 0.250 \text{ Kg} = 0.16 \text{ mol}$$

$$\text{molar mass} = \frac{\text{mass}}{n} = \frac{6.25}{0.16} = \mathbf{39.06 \text{ g/mol}}$$

$$\text{Or, } M_w (\text{solute}) = \frac{\text{mass of solute} \times K_f}{\text{mass of solvent} \times 10^{-3} \times \Delta T_f} = \frac{6.25 \text{ g} \times 3.6 \text{ }^\circ\text{C/m}}{250 \times 10^{-3} \times 2.3 \text{ }^\circ\text{C}} = \mathbf{39.06 \text{ g/mol}}$$

- **Osmosis**

- **Osmotic pressure (π) (الضغط الاسموزي)**

❖ الماء ينتقل من المحلول الذي يمتلك تركيز أقل إلى المحلول الذي يمتلك تركيز أعلى.

Osmotic pressure (π): the pressure required to stop osmosis.

Osmotic pressure:

الضغط اللازم لتوقف انتقال الماء بين المحلولين.

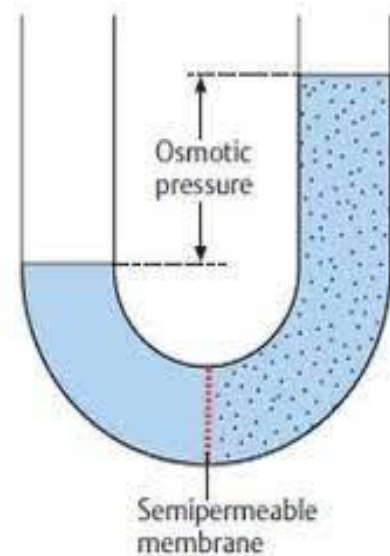
$$\pi = MRT$$

M : the molarity of the solution

T : the temperature (in K)

R : gas constant (0.0821 L . atm/K . mol)

π : osmotic pressure (atm , mmHg , ... ,)



Example 23:

A solution is prepared by dissolving 18.5 g of glucose in enough water to make up 500 mL in volume. If the osmotic pressure of the solution is found to be 8.2 mmHg at 25°C, calculate the **molar mass** of glucose.

Solution:

Osmotic pressure → *molarity* → *number of moles* → *molar mass*

| |
|------------------------------------|
| $1 \text{ atm} = 760 \text{ mmHg}$ |
|------------------------------------|

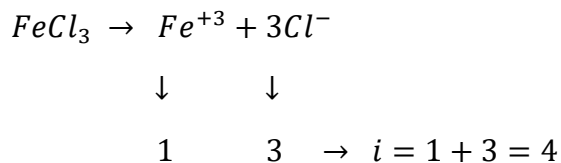
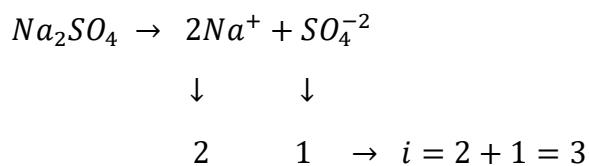
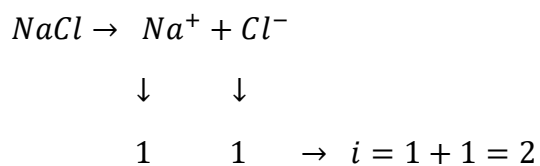
$$\pi = MRT \rightarrow M = \frac{\pi}{RT} = \frac{8.2 \text{ mmHg} \times \frac{1 \text{ atm}}{760 \text{ mmHg}}}{0.0821 \left(L \cdot \frac{\text{atm}}{K} \cdot \text{mol} \right) \times (25 + 273) K^\circ} = \frac{0.01}{0.0821 \times 298} = 0.00041 M$$

$$n = M * V = 0.00041 * 0.5 = 0.000205 \text{ mol}$$

$$M_w = \frac{\text{mass}}{n} = \frac{18.5}{0.000205} = 9.02 \times 10^4 \text{ g/mol}$$

- Osmosis**

van't Hoff factor (i) = $\frac{\text{actual number of particles in soln after dissociation}}{\text{number of formula units initially dissolved in solution}}$



Colligative Properties of Electrolyte Solutions

Boiling-Point Elevation $\Delta T_b = i K_b m$

Freezing-Point Depression $\Delta T_f = i K_f m$

Osmotic Pressure (π) $\pi = i MRT$

Example 24:

The osmotic pressure of a 0.020 M solution of sucrose at 39°C is found to be 1.23 atm. Calculate the **van't Hoff factor** for sucrose at this concentration.

- A. 1.4
- B. 2.5
- C. 2.4
- D. 3
- E. 2

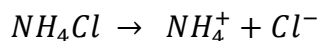
Solution:

$$\pi = iMRT \rightarrow i = \frac{\pi}{MRT} = \frac{1.23 \text{ atm}}{0.020 \text{ M} \times 0.0821 \text{ (L} \cdot \text{atm/K} \cdot \text{mol)} \times (39 + 273) \text{ K}^\circ} = \frac{1.23}{0.51} = 2.4$$

Example 25:

A 8.75 g sample of a compound with the empirical formula NH_4Cl (53.49 g/mol) is dissolved in 350 g of water. The boiling point elevation of the solution is found to be 1.2°C above that of pure water. Calculate the **molar mass** of the compound

$$K_b = 0.512 \text{ }^\circ\text{C/m}$$

Solution:

Boiling point elevation → *molality* → *number of moles* → *molar mass*

$$\Delta T_b = i * K_b * m \rightarrow m = \frac{\Delta T_b}{i * K_b} = \frac{1.2}{2 * 0.512} = 1.17 \text{ m}$$

$$m = \frac{\text{mole of solute}}{\text{mass of solvent (Kg)}} \rightarrow n = 1.17 \frac{\text{mol}}{\text{Kg}} \times 0.350 \text{ Kg} = 0.41 \text{ mol}$$

$$\text{molar mass} = \frac{\text{mass}}{n} = \frac{8.75}{0.41} = 21.3 \text{ g/mol}$$

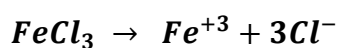
$$\text{Or, } M_w (\text{solute}) = \frac{\text{mass of solute} \times i \times K_b}{\text{mass of solvent} \times 10^{-3} \times \Delta T_b} = \frac{8.75 \text{ g} \times 2 \times 0.512^\circ\text{C/m}}{350 \text{ g} \times 10^{-3} \times 1.2^\circ\text{C}} = 21.3 \text{ g/mol}$$

Example 26:

A mole fraction of 0.097 of a compound with the empirical formula $FeCl_3$ (162.2 g/mol) is dissolved in 450 g of water.

Calculate **the freezing point** of the compound.

$$K_f = 1.86 \text{ }^\circ\text{C/m}$$

Solution:

$$n_{\text{water}} = \frac{450}{18} = 25 \text{ mol}$$

$$X_{FeCl_3} = \frac{\text{mole of } FeCl_3}{\text{mole of solution}} \rightarrow X = \frac{n_{FeCl_3}}{n_{FeCl_3} + n_{\text{water}}} \rightarrow 0.097 = \frac{n}{n + 25}$$

$$\rightarrow 0.097(n + 25) = n \rightarrow n = 2.69 \text{ mol}$$

$$m = \frac{\text{mole of solute}}{\text{mass of solvent (Kg)}} = \frac{2.69}{0.450} = 5.98 \text{ m}$$

$$\Delta T_f = i * K_f * m = 4 * 1.86 * 5.98 = 44.5 \text{ }^\circ\text{C}$$

$$\Delta T_f = T_f^\circ - T_f \rightarrow T_f = \Delta T_f + T_f^\circ = 44.5 + 0 = \mathbf{44.5 \text{ }^\circ\text{C}}$$

Problems:

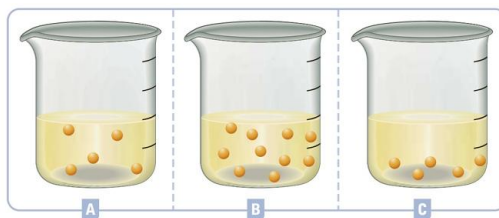
- 1) The mole fraction of copper (II) nitrate in an aqueous copper (II) nitrate solution is 0.135. What is the molality of the copper (II) nitrate solution?
 - a) 7.49 *m*
 - b) 8.01 *m*
 - c) 8.66 *m*
 - d) 9.34 *m*
 - e) 8.06 *m*

- 2) A 5.1 *g* sample of CaCl_2 is dissolved in a beaker of water. Which of the following statements is true of this solution?
 - a) The solution will freeze at a lower temperature than pure water.
 - b) The solution has a higher vapor pressure than pure water.
 - c) The solution will boil at a lower temperature than pure water
 - d) Water is the solute in this solution.
 - e) None of the other statements (a–d) are true.

- 3) All of the following are colligative properties except:
 - a) vapor pressure lowering
 - b) freezing point depression
 - c) osmotic pressure
 - d) density elevation
 - e) boiling point elevation

Ch 12

4) Consider the following three beakers that contain water and a non-volatile solute. The solute is represented by the orange spheres.



- Which solution would have the highest vapor pressure?
 - Which solution would have the lowest boiling point?
 - What could you do in the laboratory to make each solution have the same freezing point?
- 5) Arrange the following substances in order of increasing solubility in hexane, C_6H_{14} : CH_2OHCH_2OH , $C_{10}H_{22}$, H_2O .
- $CH_2OHCH_2OH < C_{10}H_{22} < H_2O$
 - $C_{10}H_{22} < CH_2OHCH_2OH < H_2O$
 - $H_2O < CH_2OHCH_2OH < C_{10}H_{22}$
 - $H_2O < C_{10}H_{22} < CH_2OHCH_2OH$
 - None of these
- 6) The solubility of carbon dioxide in water is $0.161\text{ g } CO_2$ in 100 mL of water at $20\text{ }^\circ\text{C}$ and 1.00 atm . A soft drink is carbonated with carbon dioxide gas at 5.50 atm pressure. What is the solubility of carbon dioxide in water at this pressure?

- $0.886\text{ g}/100\text{ml}$
- $0.55\text{ g}/100\text{ml}$
- $1.886\text{ g}/100\text{ml}$
- $0.75\text{ g}/100\text{ml}$
- $0.04\text{ g}/100\text{ml}$

Ch 12

7) What mass of solution containing 6.50% sodium sulfate, Na_2SO_4 , by mass contains 1.75 g Na_2SO_4 ?

- a) 34 g
- b) 27 g
- c) 33 g
- d) 30.1 g
- e) 0.75 g

8) An automobile antifreeze solution contains 2.50 kg of ethylene glycol, CH_2OHCH_2OH ($M_w = 62.07 \text{ g/mol}$), and 2.25 kg of water. Find the mole fraction of ethylene glycol in this solution. What is the mole fraction of water?

- a) 0.0078 , 0.9922
- b) 0.5 , 0.5
- c) 0.6 , 0.4
- d) 0.04 , 0.96
- e) 0.24 , 0.76

9) An antiseptic solution contains hydrogen peroxide, H_2O_2 , in water. The solution is 0.610 m H_2O_2 . What is the mole fraction of hydrogen peroxide?

- a) 0.9×10^{-2}
- b) 1.1×10^{-3}
- c) 2.1×10^{-2}
- d) 1.1×10^{-2}
- e) 1.1×10^{-9}

Ch 12

10) A solution was prepared by dissolving 0.800 g of sulfur S_8 ($M_w = 256.56 \text{ g/mol}$), in 100.0 g of acetic acid, $HC_2H_3O_2$. If the freezing point of pure acetic acid is 16.6°C and the boiling point of pure acetic acid is 118.1°C . Calculate the freezing point and boiling point of the solution.

$$K_f = 3.9^\circ\text{C}/m, K_b = 3.07^\circ\text{C}/m$$

- a) 17.9°C , 88.1°C
- b) 16.5°C , 118.1°C
- c) 17.9°C , 848.1°C
- d) 17.9°C , 88.9°C
- e) 55°C , 118.2°C

11) Which aqueous solution has highest osmotic pressure?

- a) 0.15 molar $Ba(NO_3)_2$ ($0.15 \times 3 \text{ particles} = 0.45 M$)
- b) 0.1 molar $Al(NO_3)_3$ ($0.1 \times 4 \text{ particles} = 0.4 M$)
- c) 0.1 molar $CaCl_2$ (three particle)
- d) 0.15 molar $NaCl$ (two particle)
- e) 0.2 molar NH_3 (one particle)

12) The osmotic pressure of blood at 37°C is 7.7 atm . A solution that is given intravenously must have the same osmotic pressure as the blood. What should be the molarity of a glucose solution to give an osmotic pressure of 7.7 atm at 37°C ?

- a) 0.6 M
- b) 0.9 M
- c) 0.3 M
- d) 0.1 M
- e) 0

Answers:

- 1) c (8.66 m)
- 2) a (The solution will freeze at a lower temperature than pure water)
- 3) d (density elevation)
- 4) a) C b) C c) Make the amount of **solute** in the three solutions equal.
- 5) c ($H_2O < CH_2OHCH_2OH < C_{10}H_{22}$)
- 6) A (0.886 g/100ml)
- 7) b (27 g)
- 8) e (0.24 , 0.76)
- 9) d (1.1×10^{-2})
- 10) b (16.5 °C , 118.1 °C)
- 11) a (0.15 molar $Ba(NO_3)_2$)
- 12) c (0.3 M)