

# Chap 11 Properties of Solutions 1

Table 11.1 Various Types of solutions

- 11.1 Solution composition
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- 11.3 Solubility
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- 11.5 Boiling-Point Elevation
- ⋮
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- 11.7 Electrolyte
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## § 11.1 Solution Composition

$$\text{mass percent} = \frac{\text{mass of solute}}{\text{mass of solution}} \times 100$$

$$\text{mole fraction of A} = \chi_A = \frac{n_A}{n_A + n_B} \quad \text{for a two-component sol'n}$$

$$\text{molality (m)} = \frac{\text{mole of solute}}{\text{kilograms of solvent}}$$

(M) molarity =  $\frac{\text{mole of solute}}{\text{L of solution}}$

normality: number of equivalents  $\rightarrow$   
per liter of solution

(N)

Ex. 11.1 & 11.2

1.00 g ethanol w/ 100.0 g water  
final volume = 101 mL  
calculate the molarity, mass percent, mole fraction, and molality.

$$\text{molality (m)} = \frac{\text{mole of solute}}{\text{kg solvent}}$$

$$= \frac{1.00 \text{ g}}{26.07 \text{ g/mol}}$$

molality (M)

m : molality

M : molarity

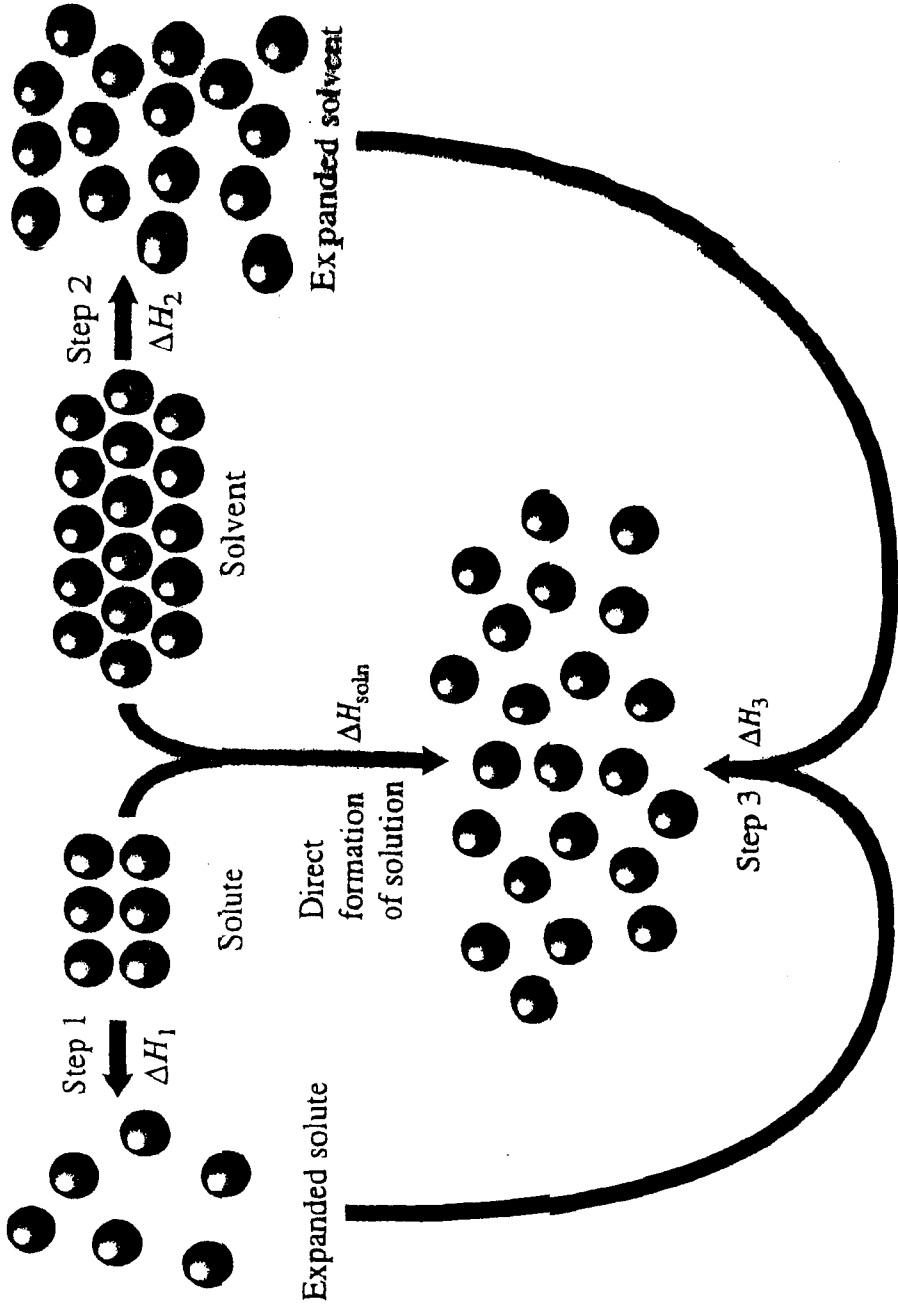


Figure 11.1  
The steps in the dissolving process

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§ 11.2 The Energies of Solution Formation

DTT: 在体内不易代谢. fat-soluble

BaSO<sub>4</sub>: X-ray 显影剂

影响 solubility 的因素  
like dissolves like

Formation of a liquid solution

1. Breaking up the solute into individual components (expanding the solute).  $\Delta H_1$
2. Overcoming intermolecular forces in the solvent to make room for the solute (expanding the solvent).  $\Delta H_2$
3. Allowing the solute and solvent to interact to form the solution.  $\Delta H_3$

Fig. 11.1 & Fig 11.2

enthalpy of solution =  $\Delta H_1 + \Delta H_2 + \Delta H_3$   
( $\Delta H_{sd}$ )

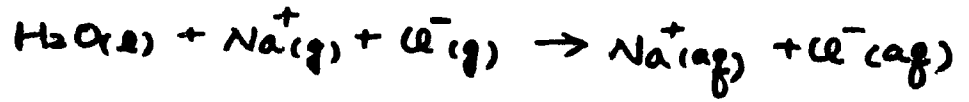
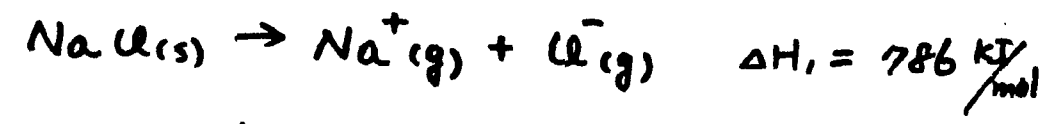
Fig. 11.1

- (a)  $\Delta H_{sol} < 0$  (放熱)
- (b)  $\Delta H_{sol} > 0$  (吸熱)

10-5

Enthalpy of hydration ( $\Delta H_{hyd}$ )  
 $= \Delta H_2 + \Delta H_3$

以 NaCl 溶於水為例



$$\Delta H_{hyd} = \Delta H_2 + \Delta H_3 = -783 \text{ kJ/mol}$$

$$\Delta H_{sol} = \Delta H_1 + \Delta H_{hyd} = 3 \text{ kJ/mol}$$

Ex 11.3.

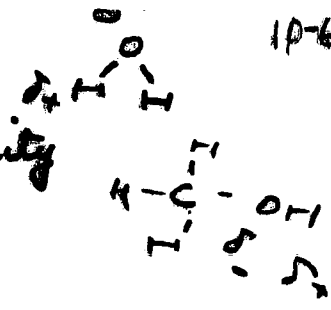
$C_6H_{12}$  (hexane) 己烷 } solvent  
 $CH_3OH$  (methanol) }

$C_{20}H_{42}$  } solute  
 KI }

Table 11.3.

10-6

§ 11.3 Factors Affecting Solubility  
structure effect



以維他命為例

vitamins: fat-soluble (A, D, E, K)  
 water-soluble (B, C)

See Fig. 11.4.

vitamin A: hydrophobic  
 vitamin C: hydrophilic

pressure effect

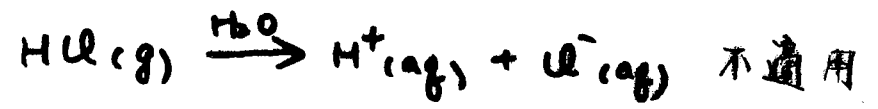
汽水

Henry's law:

$$P = k C \quad \rightarrow \text{concentration of dissolved gas}$$

↑  
gas pressure

(of dilute solutions of gases that do not dissociate  
 in or react w/ the solvent)



11.4 calculations Using Henry's Law (p.7)

The Henry's law const for  $\text{CO}_2 = 32 \text{ L} \cdot \text{atm} / \text{mol}$

$$P_{\text{CO}_2} = 4.0 \times 10^{-4} \text{ atm at } 1 \text{ atm } P$$

Sol: unopened:

$$P_{\text{CO}_2} = K C_{\text{CO}_2}$$

$$\frac{P_{\text{CO}_2}}{K} = C_{\text{CO}_2} = \frac{5.0 \text{ atm}}{32 \text{ L} \cdot \text{atm} / \text{mol}} = 0.16 \text{ mol/L}$$

opened:

$$\frac{P_{\text{CO}_2}}{K} = C_{\text{CO}_2} = \frac{4.0 \times 10^{-4} \text{ atm}}{32 \text{ L} \cdot \text{atm} / \text{mol}} = 1.2 \times 10^{-5} \text{ mol/L}$$

Temperature effect

溶解度与温度的关系

Fig 11.6 & 11.7

thermal pollution

影响溶解度

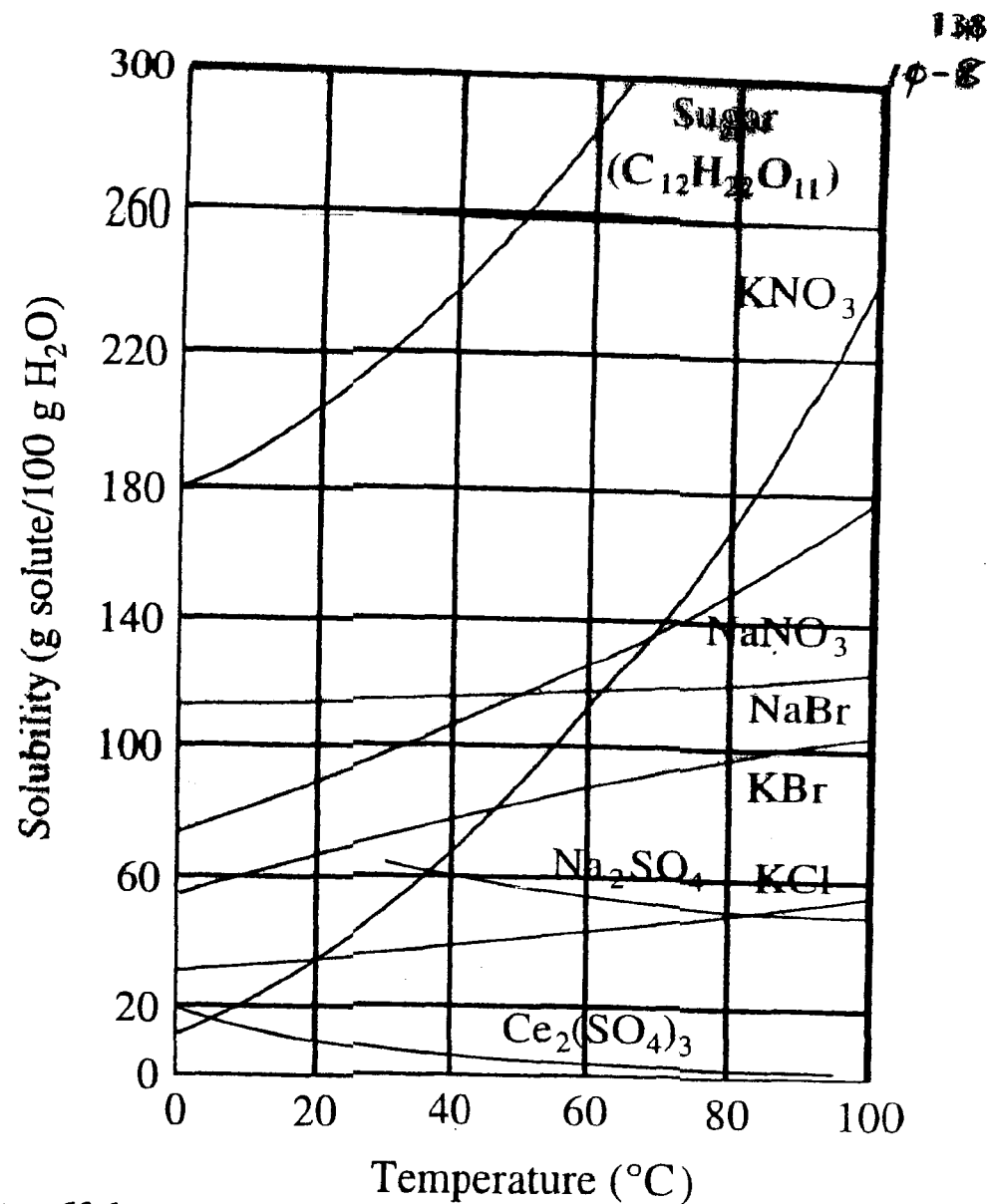


Figure 11.6

The temperature dependence of solubility for various solids

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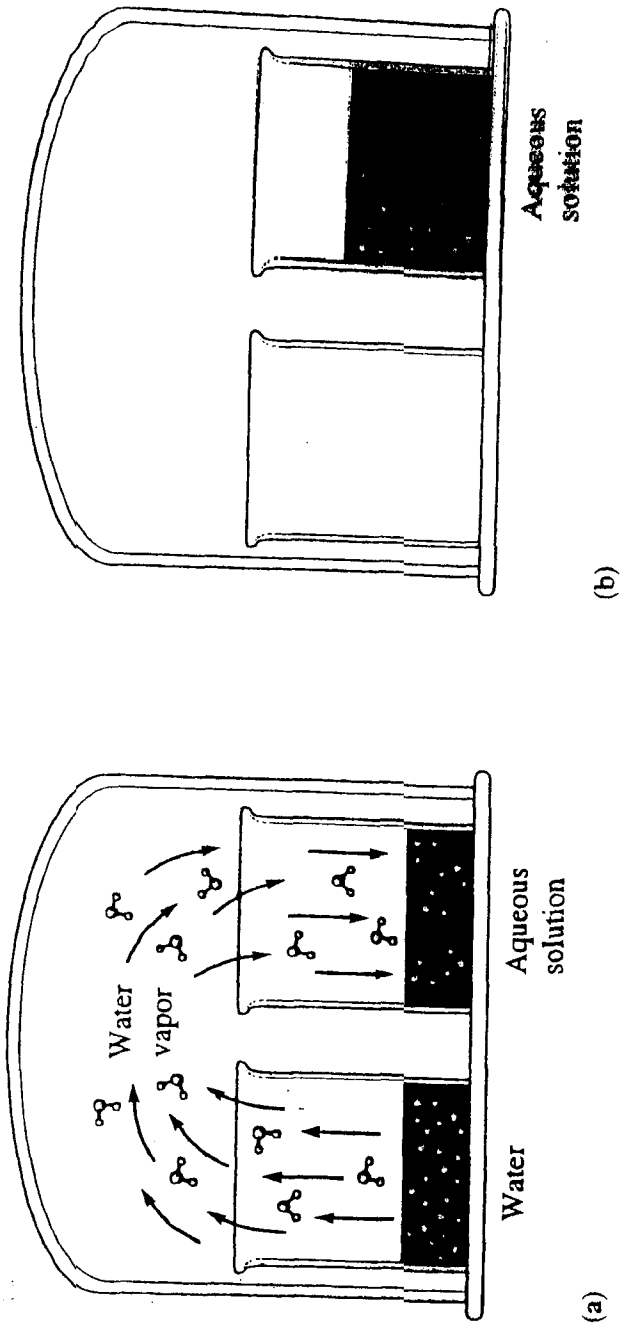


Figure 11.9

An aqueous solution and pure water in a closed environment

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139  
11.9

## § 11.4 The vapor pressures of solution

Fig 11.9

non-volatile solute lowers the vapor pressure of a solvent

Raoult's law:

$$P_{\text{sol}} = \underbrace{\chi_{\text{solvent}}}_{\substack{\uparrow \\ \text{mole fraction of solvent}}} P_{\text{solvent}}^{\circ} + \underbrace{\chi_{\text{solute}}}_{\downarrow} P_{\text{solute}}^{\circ} \approx 0$$

Ex. 11.5

158.0g sugar in 643.5 cm<sup>3</sup> water  
at 25°C, H<sub>2</sub>O density 0.9971 g/cm<sup>3</sup>  
vapor pressure = 23.76

Sol:

$$\chi_{\text{solvent}} = \frac{n_{\text{solvent}}}{n_{\text{solute}} + n_{\text{solvent}}} = \frac{643.5 \cdot 0.9971}{\frac{158.0}{342.3} + \frac{643.5 \cdot 0.9971}{18}} = 0.9873$$

$$P_{\text{sol}} = \chi_{\text{solvent}} \cdot P_{\text{solvent}}^{\circ} = 0.9873 \cdot 23.76 \text{ torr} = 23.46 \text{ torr}$$

ex 11.6.

35.0 g solid  $\text{Na}_2\text{SO}_4$  w/ 175 g water <sup>10-11</sup>

$P_{\text{H}_2\text{O}}^\circ = 23.76 \text{ torr at } 25^\circ\text{C}$

$n_{\text{H}_2\text{O}} = 175 \text{ g H}_2\text{O} \times \frac{1 \text{ mol H}_2\text{O}}{18.0 \text{ g H}_2\text{O}} = 9.72 \text{ mol H}_2\text{O}$

$n_{\text{Na}_2\text{SO}_4} = 35.0 \text{ g} \times \frac{1 \text{ mol Na}_2\text{SO}_4}{142 \text{ g Na}_2\text{SO}_4} = 0.246 \text{ mol Na}_2\text{SO}_4$

$n_{\text{solute}} = n_{\text{Na}^+} + n_{\text{SO}_4^{2-}} = 3 \times n_{\text{Na}_2\text{SO}_4}$

$= 3 \times 0.246 = 0.738 \text{ mol}$

$\chi_{\text{H}_2\text{O}} = \frac{9.72 \text{ mol}}{9.72 \text{ mol} + 0.738 \text{ mol}} = 0.929$

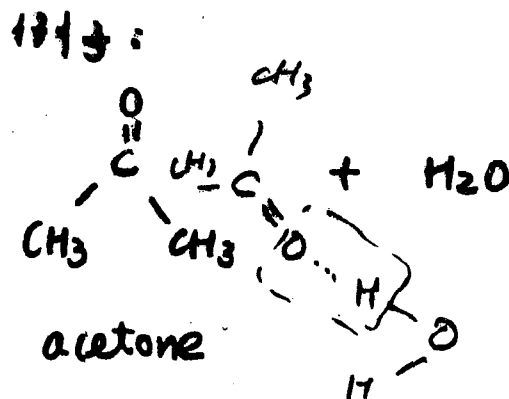
$P_{\text{soln}} = \chi_{\text{H}_2\text{O}} \cdot P_{\text{H}_2\text{O}}^\circ = (0.929)(23.76 \text{ torr}) = 22.1 \text{ torr}$

Nonideal Solutions

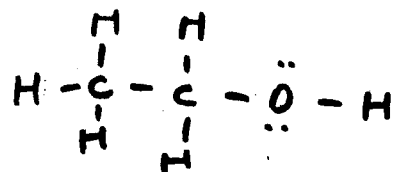
$P_{\text{Total}} \neq P_A + P_B = \chi_A P_A^\circ + \chi_B P_B^\circ$

Fig. 11.12 & 11.13.

- Non-ideal solutions <
- (a) ideal solution
  - (b) solvent-solute  $\Delta H < 0$
  - (c) solvent-solute  $\Delta H > 0$



11-12  
enthalpy of solution < 0  
negative deviation from Raoult's law



11-13  
enthalpy of solution > 0  
positive deviation from Raoult's law



very similar, enthalpy of solution ~ 0 ideal solution

Table 11.4:  
summary of the behavior of various types of solutions.

Ex. 11.7

5.81 g acetone (M.W. = 58.1 g/mol)

10-13

11.9 g chloroform (M.W. = 119.4 g/mol)

at 35°C, the solution has a total vapor pressure of 260 torr. Is the solution ideal?

$$P_{\text{acetone}}^{\circ} = 345 \text{ torr}$$

$$P_{\text{chloroform}}^{\circ} = 293 \text{ torr}$$

$$P_{\text{total}} = \chi_A P_A^{\circ} + \chi_B P_B^{\circ}$$

A: acetone

B: chloroform

$$\chi_A = \frac{\frac{5.81}{58.1} \text{ mol}}{\frac{5.81}{58.1} + \frac{11.9}{119.4}} = 0.5$$

$$\chi_B = \frac{\frac{11.9}{119.4}}{\frac{5.81}{58.1} + \frac{11.9}{119.4}} = 0.5$$

$$P_{\text{total}} = 0.5 \times 345 + 0.5 \times 293 = 319 \text{ torr}$$

$$260 \text{ (observed)} < 319 \text{ torr}$$

$$n_{\text{solute}} = 0.10 \text{ mol} = \frac{18.00 \text{ g}}{\text{molar mass}} \quad 10-1$$

molar mass of solute (glucose)

$$= \frac{18.00 \text{ g}}{0.10 \text{ mol}} = 180 \frac{\text{g}}{\text{mol}}$$

Freezing-Point Depression

$$\Delta T = K_f M_{\text{solute}}$$

$K_f$ : molal freezing-point depression constant

# § 11.5 Boiling-point Elevation and 10-15 Freezing-point Depression

$$\Delta T = K_b m_{\text{solute}}$$

$K_b$ : molal boiling-point elevation const  
 $m_{\text{solute}}$  = molarity of the solute in the solution

Fig. 11.14. & Table 11.5

## Ex. 11.8

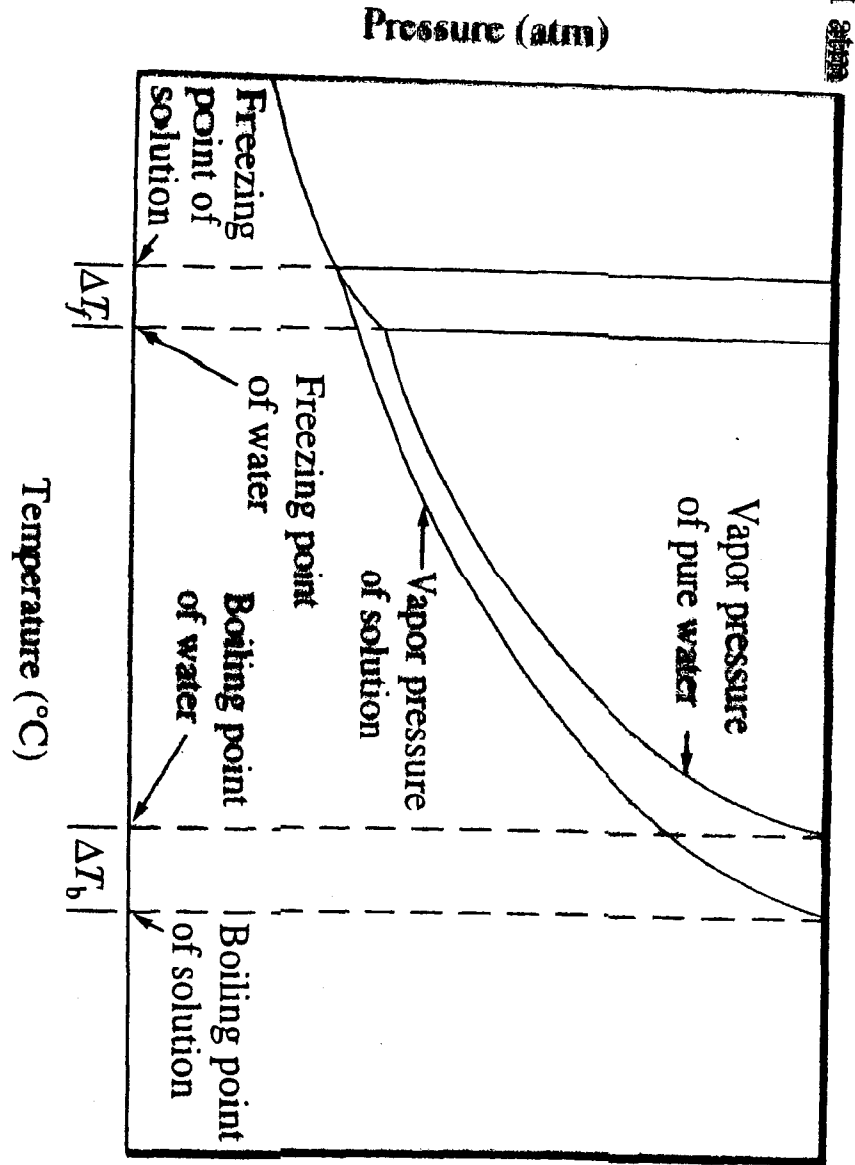
a solution was prepared by 18.00 g glucose in 150.0 g water. The resulting solution was found to have a boiling point = 100.34°C, calculate the molar mass of glucose.

Sol:  $\Delta T = K_b m_{\text{solute}}$

$$0.34^\circ\text{C} = K_b 0.51 \frac{\text{kg}}{\text{mol}} \cdot m_{\text{solute}}$$

$$M = \frac{0.67 \text{ mol}}{\text{kg}} = \frac{n_{\text{solute}}}{m_{\text{solute}}}$$

Figure 11.14  
 Vapor diagram for an aqueous solution  
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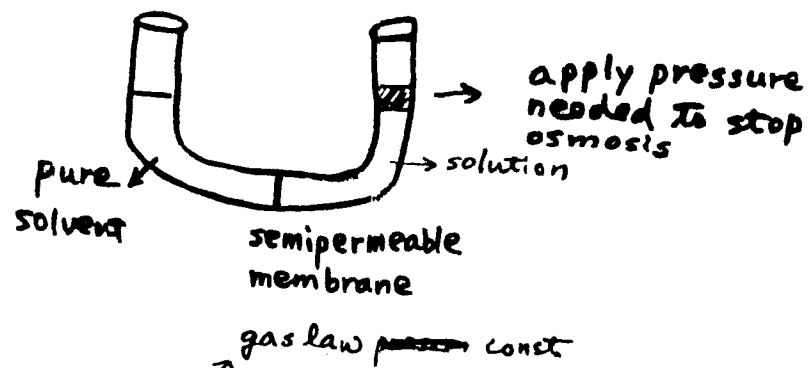
9 11.6 Osmotic Pressure (滲透壓), 10-17

semipermeable membrane (半透膜):

allow solvent but not solute molecules to pass through.

see Fig 11.16 & 11.17

As time passes, the volume of the solution increases and that of the solvent decreases.



$$\pi = M R T$$

↑ osmotic pressure (in atm)

↑ molarity (M)

↑ kelvin temp

See Fig 11.18

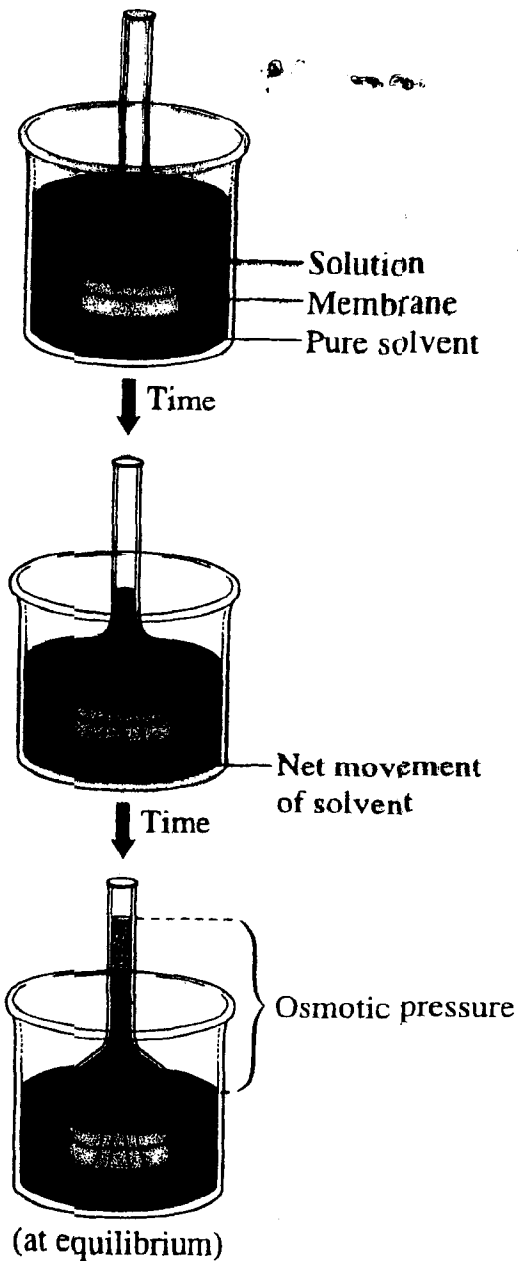


Figure 11.14  
The development of osmotic pressure

5) 1141  
10  
E

Ex. 11.11

10-19

To determine the molar mass of a certain protein,  $1.00 \times 10^{-3} \text{ g}$  in  $1.00 \text{ ml}$  of solution

osmotic pressure =  $1.12 \text{ torr}$  at  $25.0^\circ \text{C}$

→ calculate the molar mass of the protein

sol.

$$\pi = M R T$$

$$\frac{1.12 \text{ torr}}{760 \frac{\text{torr}}{\text{atm}}} = M \cdot 0.002 \frac{\text{L} \cdot \text{atm}}{\text{K} \cdot \text{mol}} \cdot (273 + 25.0)$$

$$M = 6.01 \times 10^{-5} \text{ mol/L}$$

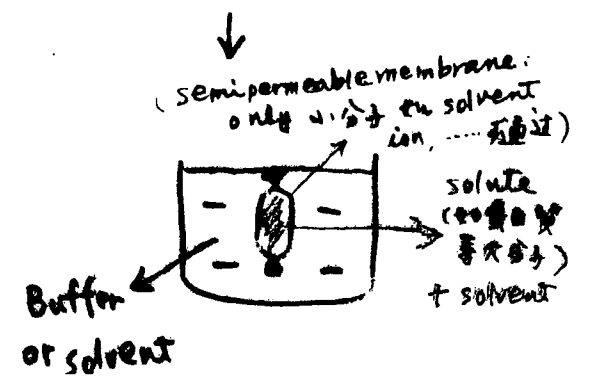
$$= \frac{1.00 \times 10^{-3} \text{ g}}{\frac{\text{M.W.}}{1.00 \times 10^{-3} \text{ L}}}$$

$$\therefore \text{MW} = 1.66 \times 10^4 \frac{\text{g}}{\text{mol}}$$

應用：洗腎機 & 透析 (dialysis)

see Fig. 11.19

(=清洗 or filter out waste molecule)  
小分子



Isotonic solutions: solutions that have identical osmotic pressure.  
(等滲壓液)

Ex. 1. 靜脈注射 (必須與血液是 isotonic)

If hypertonic (osmotic pressure > 紅血球的 osmotic pressure)

→ 血球內 solvent and small molecules 會流出 → 血球會萎縮 (crenation)

If hypotonic (osmotic pressure < 紅血球的 osmotic pressure)

→ solvent and small molecules 會流入 紅血球 → 紅血球會破裂 (hemolysis)

Ex. 2. 保存食物. 高鹽 or 高糖.

Ex. 11.12

如何 make isotonic solution using NaCl and let  $\pi = 7.10 \text{ atm}$  at  $25^\circ \text{C}$  (生理)

Sol.

$$\pi = M R T$$

$$7.10 \text{ atm} = M \cdot 0.082 \frac{\text{atm} \cdot \text{L}}{\text{mole} \cdot \text{K}} \cdot (273 + 25)$$

$$M = 0.315 \text{ M.}$$

∴ 0.315 M

## Reverse Osmosis (逆渗透):

10-21

如果溶液外加一于大于渗透压的压力, 此时溶剂会由溶液穿过 semi-permeable membrane 到溶剂.

应用: 淡化海水 → 食用水

See Fig 11.20 & 11.21 & 11.22

Apply  $800 \text{ lb/in}^2$  的压力

## § 11.7 Colligative Properties of Electrolyte Solutions

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

$$\Delta T_b = K_b \cdot m = K_b \cdot i \cdot m$$

$$\pi = M R T = i M R T$$

但此时  $m$  &  $M$  皆为溶液中的

"solute particles" 的浓度

∴ 再乘一于  $i$  (Van't Hoff factor)

$$i = \frac{\text{moles of particles in solution}}{\text{moles of solute dissolved}}$$

See Table 11.4

## § 11.8 Colloids 胶体

10-22

A suspension of tiny particles in some medium is called a colloidal dispersion or a colloid.

Ex. 泥土加水搅拌

→ 光会散射 (scatter)

又行 Tyndall effect

(The scattering of light by particles)

See Fig 11.24 The Tyndall effect

Table 11.7 Types of colloids

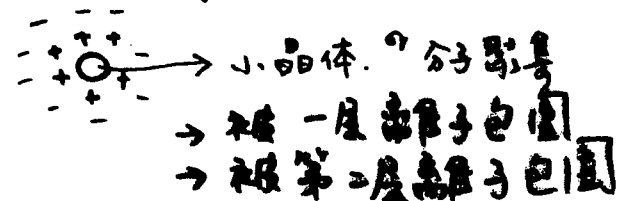
Colloid: the suspended particles

大约 1 to 1000 nm.

particles 形成原因与悬浮原因:

electrostatic repulsion

see Fig 11.25



Colloid 可凝集 heating  
• adding an electrolyte } ZB = 毒<sup>10-23</sup>

→ coagulation 凝集法

如: 1. 帶正電之可凝流之注入海 ∴ 碰到 electrolyte

→ coagulation 形成 = 角洲

2. 火煙灰 形成 (see Fig 11.26)

→ use charged plates to attract  
the colloidal particles

→ remove colloidal particles  
from smoke