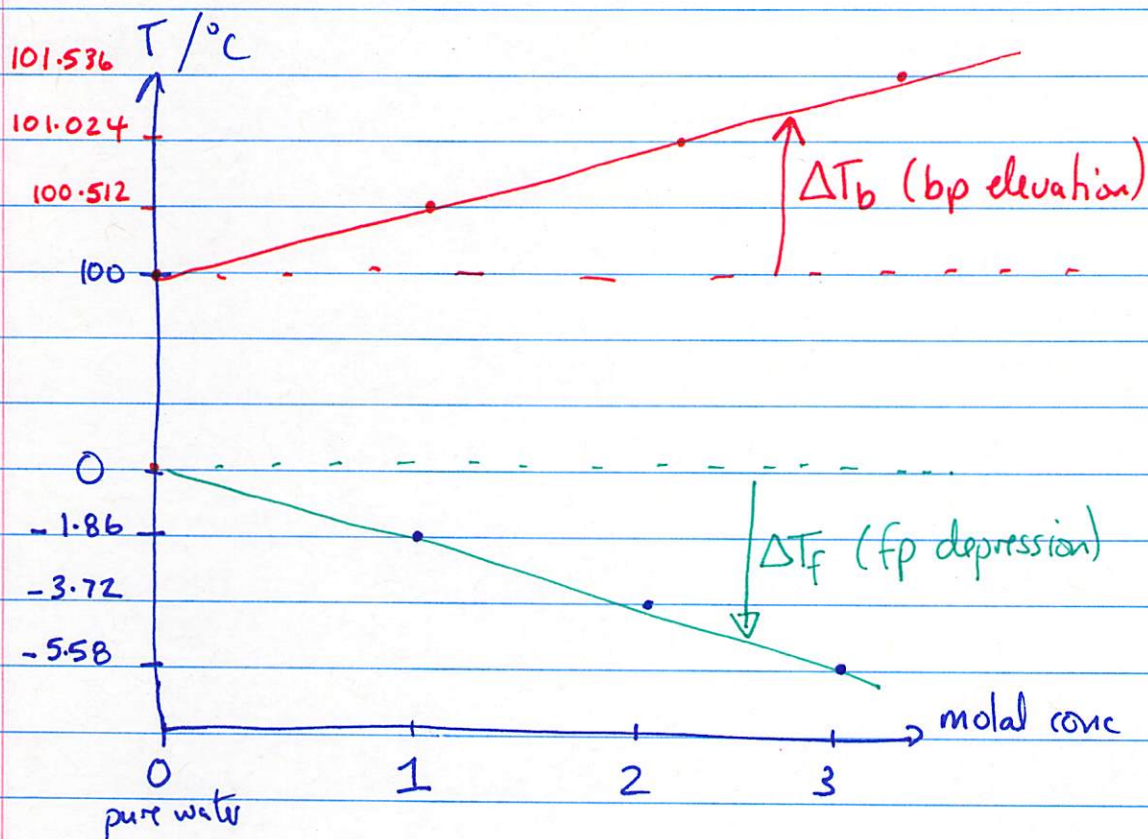


consider H<sub>2</sub>O



2/11/19

for H<sub>2</sub>O:  $\Delta T_b = m \times 0.512^\circ\text{C}/m$   
 $\Delta T_f = m \times 1.86^\circ\text{C}/m$

bp elevation constant

in general, for all solvents:  $\Delta T_b = m \times K_b$   
 $\Delta T_f = m \times K_f$

fp depression constant

ex: what's fp of 1.7m CH<sub>2</sub>O dissolved in benzene?

(normal fp)  $T_f^\circ = 5.5^\circ\text{C}$ ,  $K_f = 5.12^\circ\text{C}/m$ . Find:  $T_f$ ?

depressed by this amount

Do it!  $\Delta T_f = K_f \cdot m = 5.12^\circ\text{C}/m \cdot 1.7m = 8.704^\circ\text{C}$

$$T_f = T_f^\circ - \Delta T_f = 5.5^\circ\text{C} - 8.704^\circ\text{C} = -3.2^\circ\text{C}$$

**TABLE 13.8** Freezing Point Depression and Boiling Point Elevation Constants for Several Liquid Solvents

Solvent	Normal Freezing Point (°C)	$K_f$ (°C/m)	Normal Boiling Point (°C)	$K_b$ (°C/m)
Benzene (C <sub>6</sub> H <sub>6</sub> )	5.5	5.12	80.1	2.53
Carbon tetrachloride (CCl <sub>4</sub> )	-22.9	29.9	76.7	5.03
Chloroform (CHCl <sub>3</sub> )	-63.5	4.70	61.2	3.63
Ethanol (C <sub>2</sub> H <sub>5</sub> OH)	-114.1	1.99	78.3	1.22
Diethyl ether (C <sub>4</sub> H <sub>10</sub> O)	-116.3	1.79	34.6	2.02
Water (H <sub>2</sub> O)	0.00	1.86	100.0	0.512



What's molar mass of a solute if  $5.0\text{g}$  is added to  $125\text{g H}_2\text{O}$ , causing its bp to become  $100.43^\circ\text{C}$ ?

$$M_b = \frac{\#g \text{ solute}}{\#mol \text{ solute}}$$

$$\Delta T_b = 100.43^\circ - 100^\circ\text{C} = 0.43^\circ\text{C} \text{ (exact)}$$

✓

look-up:  $0.512^\circ\text{C}/m$

bp elevation:  $\Delta T_b = K_b \times m \Rightarrow m = \frac{\Delta T_b}{K_b} = \frac{0.43^\circ\text{C}}{0.512^\circ\text{C}/m}$

$$\Rightarrow m = 0.8398\text{m}$$

$\frac{\text{mol}}{\text{kg}}$

means:

#mol solute??

$$\frac{0.8398 \text{ mol solute}}{1 \text{ kg H}_2\text{O}}$$

$$125\text{g H}_2\text{O} \times \frac{\text{kg}}{10^3\text{g}} \times \frac{0.8398 \text{ mol solute}}{1 \text{ kg H}_2\text{O}} = 0.105 \text{ mol solute}$$

$$M_b = \frac{\#g}{\#mol} = \frac{5.0\text{g}}{0.105\text{mol}} = 48\text{g/mol}$$

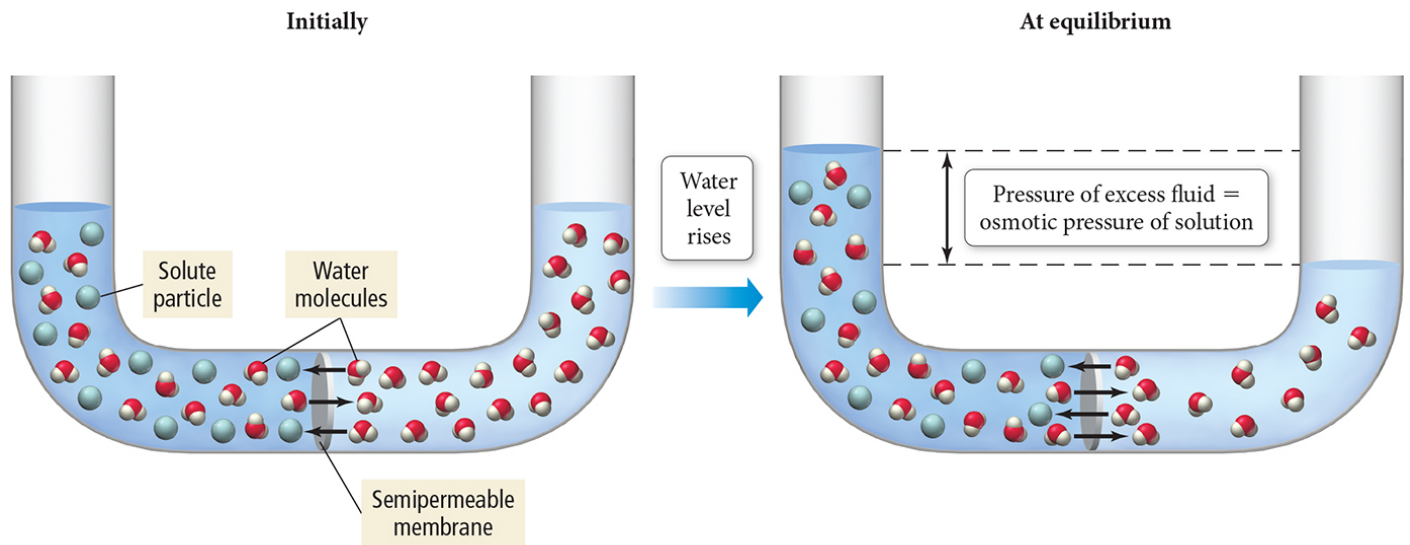
Osmotic pressure

net flow of molecules from high conc  $\rightarrow$  low conc

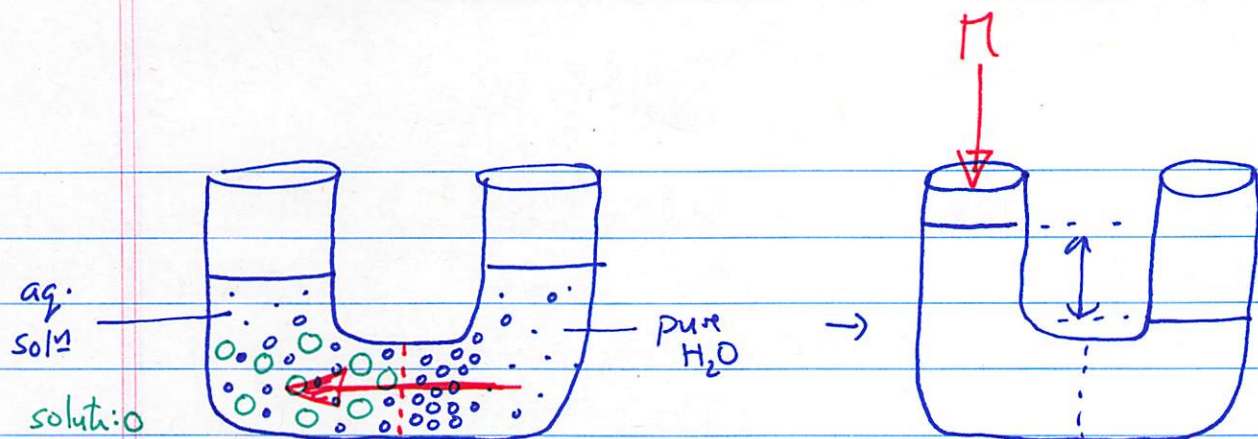
Osmosis = Diffusion of small, solvent molecules ( $\text{H}_2\text{O}$ ) through a semi-permeable membrane.

membrane w/ small holes

## Osmosis and Osmotic Pressure

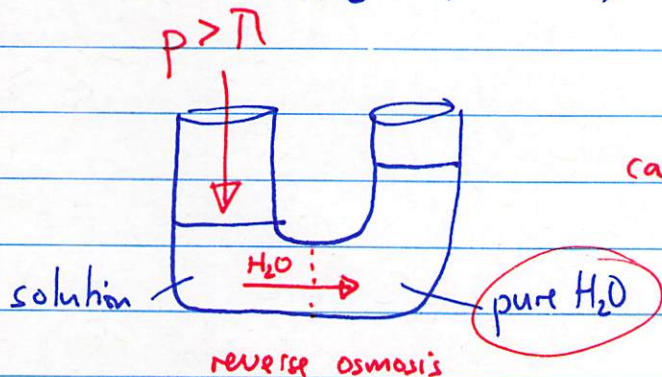






$\leftarrow$  semi-permeable membrane  
 $[Solute] \uparrow \xrightarrow{\text{diff}} [Solute] \downarrow \times$   
 $[H_2O] \downarrow \xleftarrow{\text{diff.}} [H_2O] \uparrow \checkmark$   
**OSMOSIS**

If we apply the osmotic pressure,  $\Pi$  ~ can stop osmosis  
 + if we apply  $p > \Pi$ , can reverse osmosis!



can make pure water  
 from (say) sea water!  
 - If we apply  $p > \Pi$ .

$\leftarrow$  gas constant

$$\Pi = M \cdot R \cdot T$$

$\leftarrow$  molar conc of solution      $\leftarrow$  temp/K

$$22 + 273.15 = 295.15$$

ex: Osmotic pressure of tree-sap,  $M = 0.75 M$  @  $22^\circ C$

$$\Pi = 0.75 \frac{\text{mol}}{L} \times 0.08206 \frac{\text{atm} \cdot K}{\text{mol} \cdot K} \times 295 K = 18 \text{ atm}$$

Ex 13-10 (using  $\Pi$  to find  $M$ ) p604.