

February 1, 2002

- **Quiz Today!**
- **Office Hours**
 - By appointment only, today
- **Supplemental Instruction**
 - Schedule on Web
- **Problem Set Solutions**
 - Chapter 14 now online

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Solvent Purification

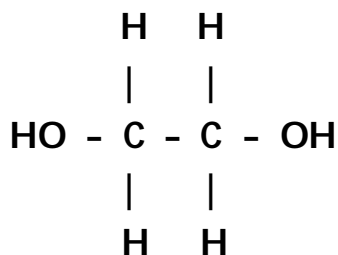
Freezing *process* often involves **solvent freezing first**

- Solid, initially, is **pure**
(zone refining, water purification)
- Solute concentration in remaining solution *increases* as solvent freezes
(FP is lowered even more)

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Automobile Antifreeze

- Add *Ethylene Glycol* ($C_2H_6O_2$) to water in radiator:



$$\text{MW} = 62.0 \text{ g/mol}$$

$$d = 1.12 \text{ g/mL}$$

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How Much EG to Add?

- Calculate the FP of a solution made by mixing 50.0 mL of EG with 50.0 g of H_2O .

1) Calculate *molality* of resulting solution.

$$50.0 \text{ mL EG} \times \frac{1.12 \text{ g EG}}{\text{mL EG}} \times \frac{1 \text{ mol EG}}{62.0 \text{ g EG}} = 0.90323 \text{ mol EG}$$

$$\text{So: } m = \frac{0.90323 \text{ mol EG}}{0.0500 \text{ kg } H_2O} = 18.06 \text{ mol EG/kg } H_2O$$

How does this compare with solution *molarity*?

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On to the solution!

2) Substitute and solve:

$$\Delta T_f = m K_f = (18.06 \text{ mol/kg})(1.86 \text{ K}\cdot\text{kg/mol})$$


$$\Delta T_f = 33.6 \text{ K} \approx 34 \text{ }^\circ\text{C}$$

So:

$$T_f = T_f^0 - \Delta T_f = 0 \text{ }^\circ\text{C} - 34 \text{ }^\circ\text{C} = \underline{\underline{-34 \text{ }^\circ\text{C}}}$$

Reality Check! Why is the *actual* FP between $-35 \text{ }^\circ\text{C}$ and $-37 \text{ }^\circ\text{C}$?

-29 °F



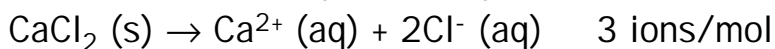
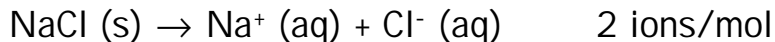
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Melting the Ice

➤ **"Salt" is added to icy roads to melt the ice**

- FP of H_2O lowered, so ice melts
- CaCl_2 more effective than NaCl **Why?**

➤ **How many ions released per mol of compound?**

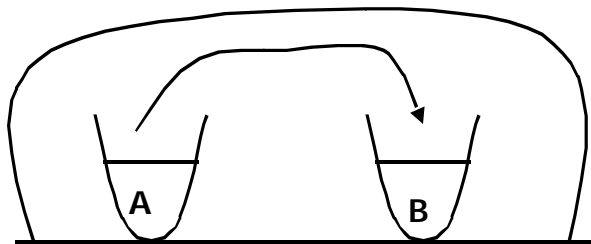


Need to consider *dissociation* of ionic compounds and the *total number* of species dissolved in solution.

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Osmosis

Consider the following experiment:



Pure
Solvent

Solution

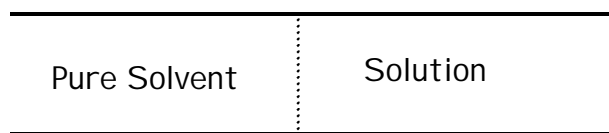
Recall that: $P_A > P_B$

So: net transfer of solvent from A to B

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Osmosis . . . *Really!*

Now, consider the following:



•Solvent "pressure" is greatest for pure solvent, so solvent flows into solution.

Semi-permeable membrane
(solvent only can pass through)

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Quantifying Osmosis

- “Pressure” of pure solvent = **Osmotic Pressure (P)**
- Magnitude will depend on *solution concentration*:

$$P = M R T$$

Osmotic Pressure (atm) Molarity (mol/L) Gas Constant (0.08206 L-atm/mol-K)

Absolute Temp (K)

Why can we use *molarity* here instead of **molality**?

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Osmosis: What is it good for?

- **Biological Systems**
 - Cell membranes
- **Water Desalinization**
 - *Reverse Osmosis*
- **MW Determination**
 - No solvent dependence of effect
 - Easier to measure:

e.g., 30 g of a 10,000 g/mol protein in 1.00 L H₂O

$$\Delta VP = 10^{-7} \text{ atm}$$

$$\Delta T_b = 10^{-4} \text{ }^\circ\text{C} \quad \Delta T_f = 10^{-3} \text{ }^\circ\text{C}$$

$$\Pi = 10^{-2} \text{ atm } (-10 \text{ torr})$$

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