

- The first rule of solubility is "likes dissolve likes"
- Polar or ionic substances are soluble in polar solvents
- Non-polar substances are soluble in nonpolar solvents



Solutes and Solution

- There must be a reason why a substance is soluble in a solvent:
 - either the solution process lowers the overall enthalpy of the system ($\Delta H_{rxn} < 0$)
 - Or the solution process increases the overall entropy of the system (ΔS_{rxn} > 0)
- Entropy is a measure of the amount of disorder in a system—entropy must increase for any spontaneous change

Solutes and Solution

- The forces that drive the dissolution of a solute usually involve both enthalpy and entropy terms
 - $\Delta H_{soln} < 0$ for most species
 - The creation of a solution takes a more ordered system (solid phase or pure liquid phase) and makes more disordered system (solute molecules are more randomly distributed throughout the solution)



- If we have enough solute available, a solution can become saturated—the point when no more solute may be accepted into the solvent
- Saturation indicates an equilibrium between the pure solute and solvent and the solution

K_C

solute + solvent ↔ solution

Saturation and Equilibrium

solute + solvent \Leftrightarrow solution K_c

- The magnitude of K_c indicates how soluble a solute is in that particular solvent
- If K_c is large, the solute is very soluble
- If K_C is small, the solute is only slightly soluble





Colligative Properties

- Colligative properties are a set properties that depend only on the amount of solute in a solution, and not on the chemical identity of the solute
- Colligative properties include:
 - Vapor pressure lowering
 - Freezing point depression
 - Boiling point elevation
 - Osmotic pressure

Colligative Properties

Vapor pressure lowering

- When solute is added to a pure solvent, solvent molecules are "tied up" in keeping the solute molecules in solution
- Because solvent molecules are more strongly attracted to the solute than to themselves, it requires more energy to remove them from the solution compared to the pure solvent

Colligative Properties

Vapor pressure lowering

- As a consequence, the vapor pressure of the solution is lowered
- Raoult's Law states:

$$\mathsf{P}_1 = \mathsf{X}_1 \, \mathsf{P}_1^{\mathsf{o}}$$

- P_1 = vapor pressure of the solution
- X_1 = mole fraction of solvent
- P_1^{o} = vapor pressure of the pure solvent

Colligative Properties

Vapor pressure lowering

 $\label{eq:example: what is the vapor pressure of a saturated NaCl solution at 25 °C? P^{o} = 23.76 Torr \\ \rho_{H2O} = 0.99707 g/mL \\ 35.7 g NaCl per 100 mL H_2O \\ \end{tabular}$ Step 1—Determine mole fraction of solution $35.7 g NaCl/58.443 g/mol = 0.611 mol NaCl (100 mL)(.9971 g/mL)/(18.0152 g.mol) \\ = 5.53 mol H_2O \\ \end{tabular}$



Vapor pressure lowering

<u>Example:</u> What is the vapor pressure of a saturated NaCl solution at 25 °C? Step 1—Determine mole fraction of solution

 $X = \frac{5.53 \text{ mol}}{2(.611 \text{ mol}) + 5.53 \text{ mol}} = .819$

Colligative Properties

Vapor pressure lowering

<u>Example:</u> What is the vapor pressure of a saturated NaCl solution at 25 °C? Step 2—Determine vapor pressure of solution

P = X P^o = (.819)(23.76 Torr)

= 19.5 Torr

The vapor pressure over a saturated NaCl solution is nearly 20% lower than that of pure water

Colligative Properties

Boiling Point Elevation

- Because the vapor pressure of solution is lower than the vapor pressure of the pure solvent, the solution's boiling point will be elevated
- Remember that the boiling point is that temperature where the vapor pressure of the solution is equal to the pressure over the solution

Boiling Point Elevation $\Delta T_{b} = T_{b}(soln) - T_{b}(solvent)$ $= K_{b} m$ $K_{b} = boiling point elevation constant$ m = molality of solute

Colligative Properties

Boiling Point Elevation

Example: What is boiling point of a NaCl solution that is saturated at 25 °C? $K_b(H_2O) = 0.512 \text{ K kg/mol}$ $m = (1.22 \text{ mol solute})/(.09771 \text{ kg } H_2O)$ = 12.5 m $\Delta T_b = (12.5 \text{ mol/kg})(0.512 \text{ K kg/mol}) = 6.40 \text{ K}$ $T_b = 106.4 \text{ °C}$

Colligative Properties

Freezing Point Depression

 The freezing point of a solution will be lower than that of the pure solvent because the solute molecules interrupt the crystal structure of the solid solvent

 $\Delta T_{f} = K_{f} m$

- K_f = freezing pt depression constant
- m = molality of solute

Colligative Properties

Freezing Point Depression

<u>Example</u>: Determine the freezing point of a solution that is 40% by volume ethylene glycol in water $K_{f}(H_{2}O) = 1.86 \text{ K kg/mol}$

 $\rho(C_2H_6O_2) = 1.109 \text{ g/mL}$

Colligative Properties

Freezing Point Depression

<u>Example:</u> Determine the freezing point of a solution that is 40% by volume ethylene glycol in water

Step 1—Assume we have 1.00 L of the solution; determine molality of ethylene glycol

(400 mL C₂H₆O₂)(1.109 g/mL)/(62.069 g/mol)

 $= 7.15 \text{ mol } C_2 H_6 O_2$

 $m = (7.15 \text{ mol})/(.600 \text{ kg H}_2\text{O}) = 11.9 \text{ m}$

