

Solutions and Their Behavior

What is a solution?

Quantitative descriptions of solutions:

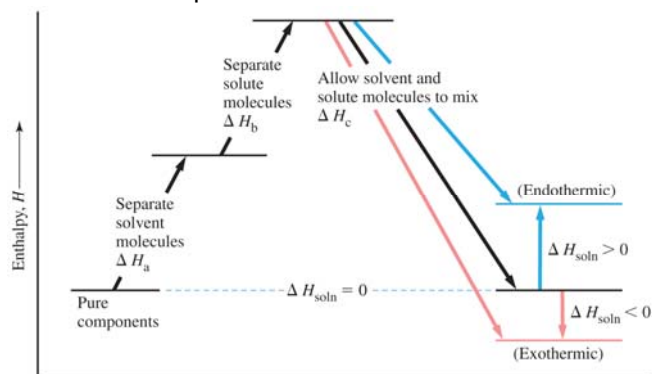
- Molarity (M):
- Molality (m):
- Mole Fraction (X):
- Weight Percent:
- Parts per (thousand/million/billion/trillion...):

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A Closer Look at Dissolution:

Largely a thermodynamic process

- Combination of Enthalpy (ΔH) and Entropy (ΔS)
- Energy "cost" for the process must not be too high
- Look at energy cost of IM forces disrupted compared to energy return of IM forces produced



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A Closer Look at Dissolution: "like dissolves like"...usually

Example: Ethylene Glycol and Hexane in Water

	Strongest IM force in pure compound	Strongest IM force in solution w/ water
$\begin{array}{c} \text{OH} \quad \text{OH} \\ \quad \\ \text{H}_2\text{C}-\text{CH}_2 \end{array}$		
$\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_3$		
H_2O		

Remember, the **tendency** for any process indicated by its Free Energy (Gibbs Energy, ΔG), which has enthalpy (ΔH) and entropy (ΔS) components! ($\Delta G = \Delta H - T\Delta S$)

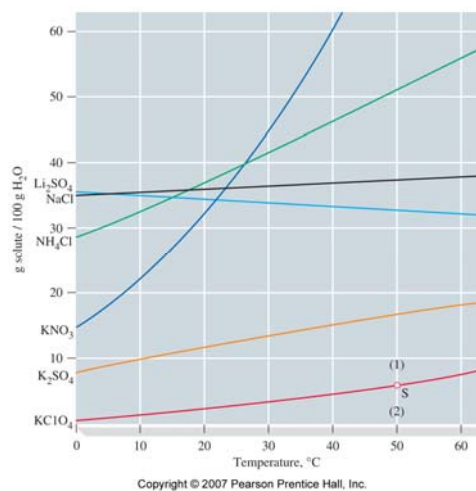
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Manipulating the Solution Process

Often **equilibrium** based.

Solubility and Temperature:

- Because of enthalpy and entropy components, we can use heat to drive dissolution
 - Impact depends on size of ΔH and ΔS parts.
- Consider heat as a reactant or product in the solubility process. What happens when we add (or remove) heat?
 - Le Chatelier's Principle:

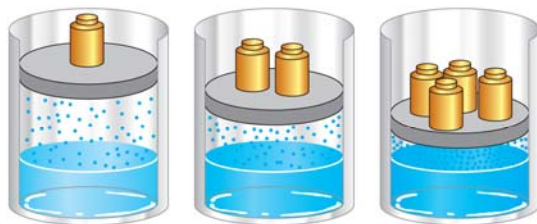


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Manipulating the Solution Process

Gas solubility and pressure: \uparrow Pressure \rightarrow \uparrow Solubility

– WHY?



- Henry's Law: $S_g = k_H P_g$

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Consequences of Forming Solutions

Colligative Properties: depend on # solute particles relative to # of solvent particles

1. Vapor Pressure Decrease

- Described by Raoult's Law: $P_{\text{solvent}} = X_{\text{solvent}} P^{\circ}_{\text{solvent}}$
- WHY?

2. Boiling Point Increase

$$\Delta t_{\text{bp}} = K_{\text{bp}} m_{\text{solute}}$$

- K_{bp} values are tabulated by solvent
- WHY?

3. Freezing Point Depression

$$\Delta t_{\text{fp}} = K_{\text{fp}} m_{\text{solute}}$$

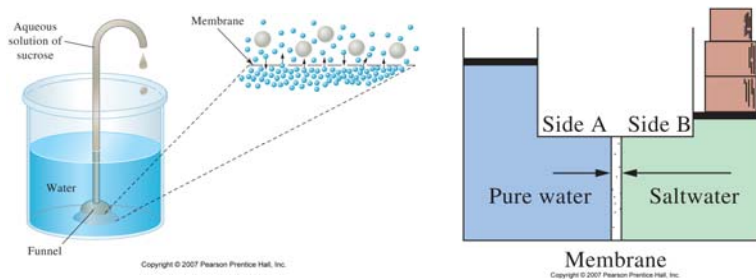
- K_{fp} values are tabulated by solvent
- WHY?

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Odds and Ends

1. Osmosis and Osmotic Pressure

- Attempt to reach a uniform concentration on both sides of semipermeable membrane.
- Results in change in pressure: $\Pi = M \times RT$



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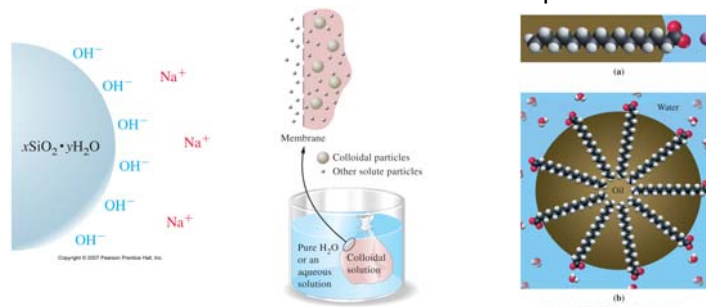
Odds and Ends

2. Electrolytes: need to account for dissociation – Van't Hoff Factor (i)

TABLE 13.3 Variation of the van't Hoff Factor, i , with Solution Molality

Solute	Molality, m					Inf. dil. ^a
	1.0	0.10	0.010	0.0010	...	
NaCl	1.81	1.87	1.94	1.97	...	2
MgSO ₄	1.09	1.21	1.53	1.82	...	2
Pb(NO ₃) ₂	1.31	2.13	2.63	2.89	...	3

3. Colloids and Surfactants: "Solutions" of small particles



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