

# Lecture 27: More Solutions

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Henry's Law, Solution Accounting, Phase Diagrams

## Solution Accounting Definitions

Quantity	Definition
Mole fraction	$\chi_n = \frac{\text{mol } n}{\text{mol total}}$
Molality	$m = \frac{\text{mol solute}}{\text{kg solvent}}$
Molarity	$M = \frac{\text{mol solute}}{\text{L solution}}$
Mass percent	$\% \text{ by mass} = \frac{\text{mass solute}}{\text{total mass}} \times 100\%$

## Example: Solution Accounting

An aqueous solution of  $\text{H}_2\text{O}_2$  is 30.0% by mass and has a density of  $1.11 \frac{\text{g}}{\text{mL}}$ . Calculate its molarity, molality, and mole fraction of  $\text{H}_2\text{O}_2$ .

Key idea: Ratios, so can assume 1000 mL solution

$$1.11 \frac{\text{g}}{\text{mL}} \times 1000 \text{ mL} = \text{1110 g solution}$$

$$\frac{x \text{ g}}{1110 \text{ g}} \times 100\% = 30\% \implies x = \text{333 g H}_2\text{O}_2$$

$$333 \text{ g H}_2\text{O}_2 \times \frac{1 \text{ mol}}{34 \text{ g}} = \text{9.79 mol H}_2\text{O}_2$$

$$1110 \text{ g solution} - 333 \text{ g H}_2\text{O}_2 = \text{777 g H}_2\text{O}$$

$$777 \text{ g H}_2\text{O} \times \frac{1 \text{ mol}}{18 \text{ g}} = \text{43.17 mol H}_2\text{O}$$

## Example: Solution Accounting

$$n_{\text{H}_2\text{O}_2} = 9.79 \text{ mol} \quad n_{\text{H}_2\text{O}} = 43.17 \text{ mol} \quad \text{kg solution} = 0.777 \text{ kg} \quad \text{L solution} = 1.000 \text{ L}$$

$$\chi_{\text{H}_2\text{O}_2} = \frac{n_{\text{H}_2\text{O}_2}}{n_{\text{total}}} = \frac{9.79 \text{ mol}}{9.79 \text{ mol} + 43.17 \text{ mol}} = \boxed{0.185}$$

$$m = \frac{n_{\text{H}_2\text{O}_2}}{\text{kg solvent}} = \frac{9.79 \text{ mol}}{0.777 \text{ kg}} = \boxed{12.6 \frac{\text{mol}}{\text{kg}} \text{ H}_2\text{O}_2}$$

$$M = \frac{n_{\text{H}_2\text{O}_2}}{\text{L solution}} = \frac{9.79 \text{ mol}}{1.000 \text{ L}} = \boxed{9.79 \frac{\text{mol}}{\text{L}} \text{ H}_2\text{O}_2}$$

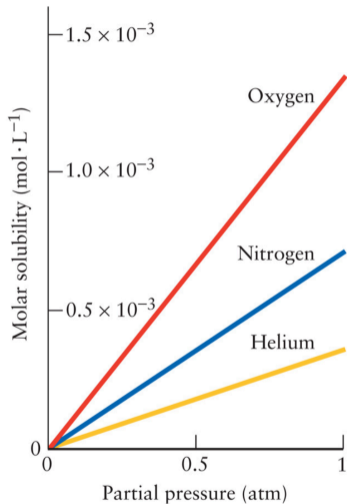
# Henry's Law

## Henry's Law

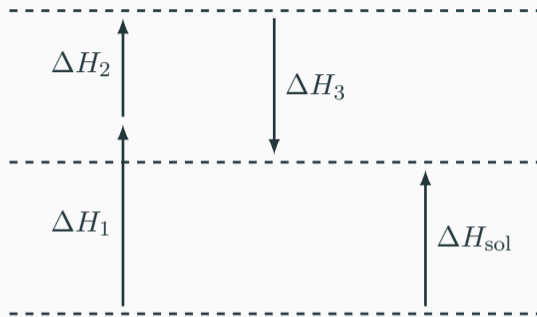
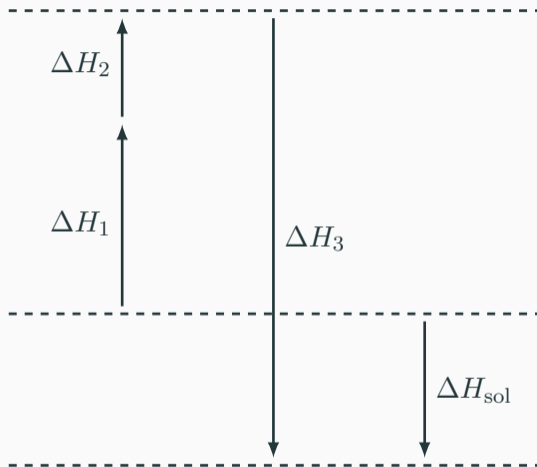
Solubility of a gas in solution is proportional to partial pressure

$$s = k_H \cdot P$$

$k_H$ : Henry's Law constant,  $T$  dependent



# Sign of $\Delta H_{\text{sol}}$



- $\Delta H_1$ : break solute IMF
- $\Delta H_2$ : break solvent IMF
- $\Delta H_3$ : form solute–solvent IMFs

$\Delta H_{\text{sol}}^{(\text{solid})}$  can be positive or negative, but  $\Delta H_{\text{sol}}^{(\text{gas})} < 0$

## Example: Henry's Law

Water is added to an empty 1.0 L container.  $\text{H}_2\text{S}(\text{g})$  is then added. When equilibrium is reached at 298 K, the pressure of  $\text{H}_2\text{S}(\text{g})$  reads 0.85 bar and there is 250 mL of solution. How many moles of  $\text{H}_2\text{S}$  are in each of the gas and aqueous phases?  $k_H(\text{H}_2\text{S}, 298 \text{ K}) = 0.10 \frac{\text{mol}}{\text{L} \cdot \text{bar}}$

Aqueous phase mols

$$s = k_H P$$

$$s = \left(0.10 \frac{\text{mol}}{\text{L} \cdot \text{bar}}\right) (0.85 \text{ bar})$$

$$= 0.085 \frac{\text{mol}}{\text{L}}$$

$$n_{\text{H}_2\text{S}(\text{aq})} = 0.085 \frac{\text{mol}}{\text{L}} \cdot 0.250 \text{ L}$$

$$= \boxed{0.02125 \text{ mol}}$$

Gas phase mols

$$n_{\text{H}_2\text{S}(\text{g})} = \frac{PV}{RT}$$

$$= \frac{(0.85 \text{ bar})(0.75 \text{ L})}{(0.08314 \frac{\text{L} \cdot \text{bar}}{\text{mol} \cdot \text{K}})(298 \text{ K})}$$

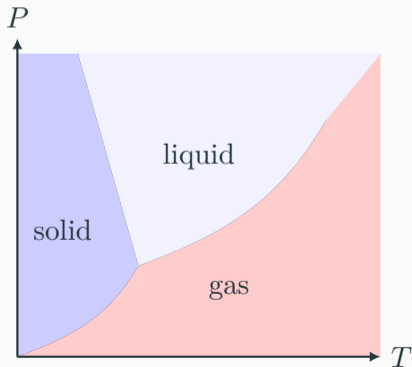
$$= \boxed{0.0257 \text{ mol}}$$

# Effect of Pressure on Boiling Point

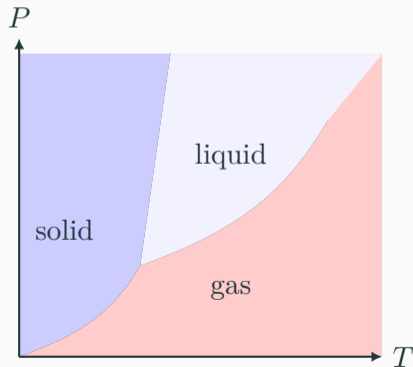


Higher external pressure favors liquid, more heat required, higher  $T_b$

Water phase diagram



Most things phase diagram



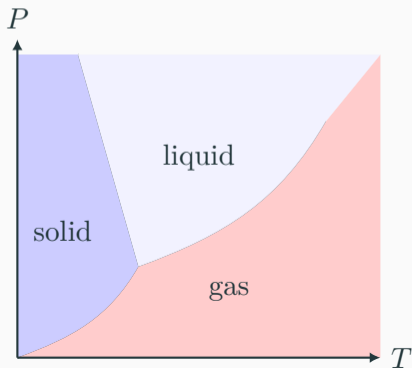
# Effect of Pressure on Melting Point



Higher external pressure favors denser phase

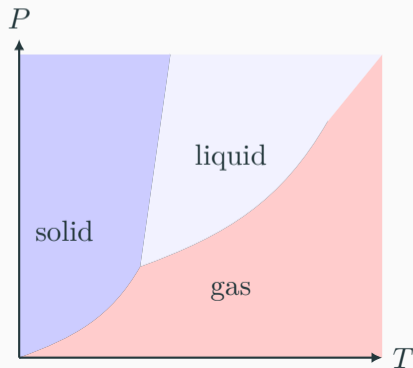
Density liquid > Density solid

$$\uparrow P \implies \downarrow T_m$$



Density solid > Density liquid

$$\uparrow P \implies \uparrow T_m$$



Exam 3: Lecture 19 – 27 (Up through solutions)

Lecture 28: Colligative

Activity 28: Applications of solutions, Colligative

Exam 4: Cumulative, emph on Lecture 29 – 37 (Colligative, Kinetics, Mechanisms)