

Colligative Properties

These are properties of solutions in the **dilute** limit, where there is a solvent "A" and a solute "B" where $n_A \gg n_B$.

These properties are a direct result of $\mu_A^{\text{mix}}(l, T, p) < \mu_A^{\text{pure}}(l, T, p)$

Use two measures of concentration:

- a. Mole Fraction: $x_B = n_B / (n_A + n_B) \sim n_B / n_A$
- b. Molality: $m_B = (\text{moles solute}) / (\text{kg solvent}) = n_B / (n_A M_A)$
Where M_A is the mass in kg of one mole of solvent.

There are **FOUR Colligative Properties**:

1. Vapor pressure lowering: $\Delta p_A = p_A - p_A^* = -x_B p_A^*$

2. Boiling point elevation: $\Delta T_b = T_b - T_b^* = K_b m_B$

$$K_b = \frac{M_A R (T_b^*)^2}{\Delta H_{\text{vap}}}$$

3. Freezing point depression: $\Delta T_f = T_f - T_f^* = -K_f m_B$

$$K_f = \frac{M_A R (T_f^*)^2}{\Delta H_f}$$

4. Osmotic pressure: $\pi = RT\tilde{c}$

where $\tilde{c} = \frac{n_B}{V}$ is concentration of solute

Let's go through these one at a time:

1. Vapor pressure lowering: This is just Raoult's Law.

$$p_A = x_A p_A^* = (1 - x_B) p_A^* \quad \text{So} \quad \Delta p_A = p_A - p_A^* = -x_B p_A^* (<0)$$

2. Boiling point elevation:

Let's derive this. Start with $\mu_A(l, T, p) = \mu_A(g, T, p)$

$$\text{So,} \quad \mu_A^*(l, T, p) + RT \ln x_A = \mu_A(g, T, p)$$

$$\text{And} \quad \ln x_A = \frac{1}{RT} [\mu_A(g, T, p) - \mu_A^*(l, T, p)] = \frac{\Delta \bar{G}_{\text{vap}}}{RT}$$

$$\text{But } \ln x_A = \ln(1 - x_B) \sim -x_B = -n_B / (n_A + n_B) \sim -n_B / n_A = -(M n_B) / (M n_A)$$

Where M is the total mass of A ,

So, $\ln x_A \sim m_B M_A$, where M_A is the molar mass of A .

$$\text{Putting it all together then,} \quad m_B = \frac{-\Delta \bar{G}_{\text{vap}}}{M_A RT}$$

But we need ΔT in there!

$$\dots \left(\frac{\partial m_B}{\partial T} \right)_p = \frac{-1}{M_A R} \left(\frac{\partial (\Delta \bar{G}_{\text{vap}} / T)}{\partial T} \right)_p = \frac{-\Delta \bar{H}_{\text{vap}}}{M_A R T^2}$$

$$\text{which gives us } \delta T = \frac{M_A R T^2}{\Delta \bar{H}_{\text{vap}}} \delta m_B$$

If $\delta m_B = m_B - 0$ (mixed - pure) and m_B is very small

Then

$$\delta T = T_b - T_b^* = \frac{M_A R (T_b^*)^2}{\Delta \bar{H}_{\text{vap}}} m_B = K_b m_B$$

3. Freezing point depression:

Same arguments as above, replacing

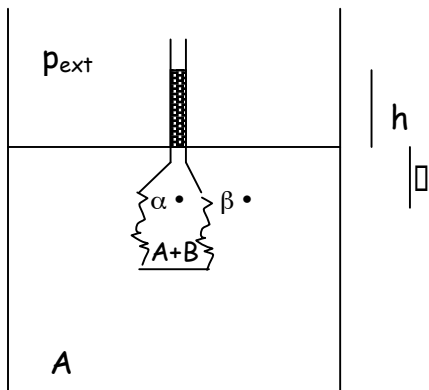
ΔG_{vap} with $-\Delta G_f$

ΔH_{vap} with $-\Delta H_f$

T_b with T_f

K_b with K_f

4. Osmotic Pressure:



The pressures at points:

$$\beta: p = p_{\text{ext}} + \rho g h$$

$$\alpha: p + \pi = p_{\text{ext}} + \rho g h$$

At equilibrium: $\mu_A(l, p + \pi, T) = \mu_A^*(l, p, T)$

Using Raoult's Law: $RT \ln x_A + \mu_A^*(l, p + \pi, T) - \mu_A^*(l, p, T) = 0$

At constant T: $dG = V dp$, or $d\mu_A^* = \bar{V}_A^* dp$

Integrating...
$$\mu_A^*(l, p + \pi, T) - \mu_A^*(l, p, T) = \int_p^{p+\pi} \bar{V}_A^* dp = \bar{V}_A^* \pi$$

(this assumed an incompressible liquid, where volume does not depend on p)

So then $RT \ln x_A + \bar{V}_A^* \pi = 0$

Again using $\ln x_A \sim -n_B/n_A$

Then $RT(-n_B/n_A) + (V_A/n_A)\pi = 0$

But $V_A \sim V_A + V_B = V$ ($V_B \ll V_A$)

So finally,
$$\pi V = RT n_B$$

This is the Van't Hoff Equation. It looks like the ideal gas law!

If we replace $\tilde{c} = n_B/V$ in the Van't Hoff Eq., then we get the osmotic pressure relation:

$$\pi = RT \tilde{c}$$