Chemistry 163B Colligative Properties Challenged Penpersonship Notes

colligative properties of solutions



colligative One entry found.

Main Entry: col·li·ga·tive Pronunciation: 'kä-lə-_lgā-tiv, kə-^lli-gə-tiv Function: adjective

: depending on the number of particles (as molecules) and not on the nature of the particles ressure is a colligative property>

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quantitative treatment of colligative properties

Handout #52

- A. Freezing point depression
- B. Boiling Point Elevation
- C. Osmotic Pressure

quantitative treatment of colligative properties

- The pure solvent (component B) is originally in equilibrium in the two
- II. Addition of solute (component A) lowers the chemical potential of the solvent in the solution phase
- III. Temperature (freezing point depression, boiling point elevation) or pressure (osmotic pressure) must be altered to reestablish equilibrium between the solution and the pure solvent phase.
- IV. Obtain relationships between $\boldsymbol{X}_{\!A}$ or $\boldsymbol{X}_{\!B}$ and change in T or P.

freezing point depression (solid ≠ solution)

I. pure solvent is originally in equilibrium in the two phases

pure solid
$$_{s}^{*} \rightleftharpoons pure \ liquid_{B}^{*} \quad at \ T_{_{f}}^{*} \quad the normal melting \ T_{fusion}$$

$$\mu_{B}^{**}(T_{_{f}}^{*}) = \mu_{B}^{(*)}(T_{_{f}}^{*})$$

$$\Delta \mu_B(T_f^{\bullet}) = \mu_B^{\ell \bullet}(T_f^{\bullet}) - \mu_B^{s \bullet}(T_f^{\bullet}) = 0$$

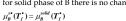
$$\Delta \overline{\mu}(T^{\bullet}) = \Delta \overline{\mu}$$

for solid \longrightarrow liquid $\Delta \overline{H}(T_{f}^{\bullet}) = \Delta \overline{H}_{B \text{ melting}} > 0$

freezing point depression (solid ≠ solution)

II. Still at $T_f^{\:\raisebox{3.5pt}{\text{\circle*{1.5}}}}$, add solute A to solvent with resulting mole fractions X_A and X_B

for solid phase of B there is no change:



for the solvent (B) in solution:

$$\mu_B^\ell(T_f^\bullet) \equiv \mu_B^{solvent} \equiv \mu_B^{\ell\,(in\,so\,\ln)}(T_f^\bullet) = \mu_B^{\ell\bullet}(T_f^\bullet) + RT_f^\bullet \ln\left(\gamma_B X_B\right)$$

so now
$$\Delta \mu_B(T_f^*) = \mu_B^\ell(T_f^*) - \mu_B^{\star\star}(T_f^*) = \Delta \mu_B^{\star}(T_f^*) + RT_f^* \ln(\gamma_B X_B)$$

where $\Delta \mu_A^{\star}(T_f^*) = \mu_B^{\star\star}(T_f^*) - \mu_B^{\star\star}(T_f^*)$

and $\Delta \mu_{\bullet}^{\bullet}(T_{\bullet}^{\bullet}) = 0$ since pure liquid and solid are in equilibrium at T_{\bullet}^{\bullet}

so the forward reacton (melting of the solid) would now occur spontaneously at T

freezing point depression (solid ≠ solution)

III. Alter temperature to restore equilibrium $T_f^{\:\raisebox{3.5pt}{\text{\circle*{1.5}}}} o T_f$

$$\left(\frac{\partial \frac{\Delta \mu}{T}}{\partial T}\right) = -\frac{\Delta \overline{H}}{T^2}$$

$$\begin{split} & \int\limits_{T_f}^{T_f} d\left(\frac{\Delta \mu_B}{T}\right)_p = -\int\limits_{T_f}^{T_f} \frac{\Delta \overline{H}_{B \, moling}}{T^2} dT \\ & \left(\frac{\Delta \mu_B(T_f)}{T_f}\right)_p - \left(\frac{\Delta \mu_B(T_f')}{T_f'}\right)_p = -\int\limits_{T_f}^{T_f} \frac{\Delta \overline{H}_{B \, moling}}{T^2} dT \end{split}$$

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freezing point depression (solid ≠ solution)

III. Alter temperature to restore equilibrium (continued)

$$\begin{split} \left(\frac{\Delta\mu_{B}(T_{f})}{T_{f}}\right)_{p} - \left(\frac{\Delta\mu_{B}(T_{f}^{*})}{T_{f}^{*}}\right)_{p} = -\int_{T_{f}^{*}}^{T_{f}} \frac{\Delta\overline{H}_{B \, moding}}{T^{2}} dT \\ \left(\frac{\Delta\mu_{B}(T_{f})}{T_{f}}\right)_{p} = 0 \text{ since at 'new' equilibrium } T_{f} \text{ , } \Delta\mu_{B}(T_{f}) = 0 \end{split}$$

$$\left(\begin{array}{c} T_f \\ \end{array} \right)_p = 0 \text{ since at new equilibrium } I_f \text{ , } \Delta \mu_B (I_f)$$
 and
$$\left(\begin{array}{c} \Delta \mu_B (T_f^{\star}) \\ T_f^{\star} \end{array} \right)_p = R \ln \left(\gamma_B X_B \right) \quad \text{from eqn in II.}$$

$$-R\ln(\gamma_{B}X_{B}) = -\int_{T}^{T_{f}} \frac{\Delta \overline{H}_{B \text{ melting}}}{T^{2}} dT$$

$$R\ln(\gamma_B X_B) + \left[-\int_{T}^{T_f} \frac{\Delta \overline{H}_{B \text{ melting}}}{T^2} dT \right] = 0$$

ange in Δμ_e due to adding solute change in Δμ

change in $\Delta\mu_{\rm B}$ due to temperature change

freezing point lowering

IV. Obtain relationships between X_B and change in T

$$R\ln\left(\gamma_{B}X_{B}\right) = \int_{T_{c}}^{T_{f}} \frac{\Delta \overline{H}_{B \text{ melting}}}{T^{2}} dT$$

 $\Delta \overline{H}_{B melting} \sim \text{independent of } T$

$$R\ln(\gamma_B X_B) = -\Delta \overline{H}_{B \text{ melting}} \left[\frac{1}{T_f} - \frac{1}{T_f^*} \right]$$

since $lhs < 0 \implies T_f < T_f^{\bullet}$ (freezing point **depression**)

$$\gamma_B X_B = \exp\left[-\frac{\Delta \overline{H}_{B melting}}{R} \left[\frac{1}{T_f} - \frac{1}{T_f^*}\right]\right]$$
 (integration of eqn 9.31 E&R)

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freezing point lowering

IV. Obtain relationships between X_{B} and change in T (cont)

$$\gamma_{_{B}}X_{_{B}} = \exp \left[-\frac{\Delta \overline{H}_{^{B} faction}}{R} \left[\frac{1}{T_{_{f}}} - \frac{1}{T_{_{f}}^{\star}} \right] \right]$$

$$\begin{split} &-\frac{R}{\Delta \overline{H}_{B \text{ fusion}}} \ln \left(\gamma_B X_B\right) + \frac{1}{T_f} = \frac{1}{T_f} \\ &T_f = \frac{T_f^* \Delta \overline{H}_{B \text{ fusion}}}{\Delta \overline{H}_{B \text{ fusion}} - R T_f^* \ln \left(\gamma_B X_B\right)} \quad (\sim \text{ eqn } 9.32 \text{ E\&R}) \end{split}$$

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Osmotic Pressure Equilibrium $P = P_o + \pi$ not permeable to solute aqueous soln. left right

 $osmotic\ pressure\ (pure\ solvent\ \rightleftarrows\ solution\ [solvent\ +\ solute]\)$

I. pure solvent at P_{left} is originally in equilibrium with pure solvent at P_{right} ; i.e. P_{left} = P_{right} = P_0

 $\begin{aligned} &pure\ \ell iquid_*(P_0,left) \rightleftharpoons pure\ \ell iquid_*(P_0,right) \quad at\ T \end{aligned}$ left' and 'right' refer to compartments separated by solute impermeable membrane $&\mu_{\rm B}(P_0,left) = \mu_{\rm B}^*(P_0,right) \end{aligned}$

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osmotic pressure (II add solute to left compartment)

II. in left hand compartment add solute A to solvent with resulting mole fractions X_A and X_B

 $add \ X_{\scriptscriptstyle A} \ solute \ to \ liquid \ in \ 'left' \ compatment \ resulting \ in \ X_{\scriptscriptstyle B} \ for \ solvent$ $\mu_{\scriptscriptstyle B}^{\prime}(P_0, left) = \mu_{\scriptscriptstyle B}^{\prime *}(P_0, left) + RT \ln \left(\gamma_{\scriptscriptstyle B} X_{\scriptscriptstyle B}\right) < \mu_{\scriptscriptstyle B}^{\prime *}(P_0, right)$

 $\mu_{b}^{\ell}(P_{0}, left) < \mu_{b}^{\ell} \cdot (P_{0}, right)$ so the solvent B moves spontaneously left \leftarrow right (i.e. diluting solution on left)

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osmotic pressure (III, alter pressure)

III. alter Pressure: $P_{left} \rightarrow P_0 + \pi$ to restore equilibrium

 $solution \ (X_{\scriptscriptstyle B}, P_{\scriptscriptstyle 0} + \pi, left) \rightleftarrows pure \ solvent(P_{\scriptscriptstyle 0}, right)$

$$\left(\frac{\partial \mu_B^{left}}{\partial P}\right)_T = \overline{V}_B$$

$$\int_{P_{0}}^{P_{0}+\pi} d\mu_{B}^{left}(X_{B}) = \int_{P_{0}}^{P_{0}+\pi} \overline{V}_{B} dP$$

assuming solvent is incompressible

 $(\overline{V}_{\!\scriptscriptstyle B}$ doesn't change with pressure at constant T)

$$\mu_{\scriptscriptstyle B}^{\scriptscriptstyle left}\left(X_{\scriptscriptstyle B}\,,P_{\scriptscriptstyle 0}+\pi\right)=\mu_{\scriptscriptstyle B}^{\scriptscriptstyle left}\left(X_{\scriptscriptstyle B}\,,P_{\scriptscriptstyle 0}\right)+\left[P_{\scriptscriptstyle 0}+\pi-P_{\scriptscriptstyle 0}\right]\overline{V}_{\scriptscriptstyle B}$$

$$\mu_B^{left}(X_B, P_0 + \pi) = \mu_B^{left}(X_B, P_0) + \pi \overline{V}_B$$

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osmotic pressure (III, alter pressure, continued)

$$\mu_B^{left}\left(X_B, P_0 + \pi\right) = \mu_B^{left}\left(X_B, P_0\right) + \pi \overline{V}_B$$

$$\mu_B^{left}(X_B, P_0 + \pi) = \overline{\mu_B^{\bullet}(P_0) + RT \ln(\gamma_B X_B)} + \pi \overline{V}_B$$

want π to restore equilibrium such that $\mu_B^{left}\left(\boldsymbol{X}_B,\boldsymbol{P}_0+\pi\right)=\mu_B^{\bullet right}\left(\boldsymbol{P}_0\right)$

$$\underline{\mu_{B}^{\bullet}(P_{0}) + RT \ln(\gamma_{B}X_{B}) + \pi \overline{V}_{B}} = \underline{\mu_{B}^{\bullet}(P_{0})}$$

$$\pi = -rac{RT\lnig(\gamma_{_B}X_{_E}}{ar{V}_{_B}}$$

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osmotic pressure (a little more manipulation)

$$\pi = -\frac{RT\ln\left(\gamma_{\scriptscriptstyle B} X_{\scriptscriptstyle B}\right)}{\bar{V}_{\scriptscriptstyle B}}$$

for
$$\gamma_B \approx 1$$
 and $X_B = 1 - X_A$

$$\pi = -\frac{RT \ln(1 - X_A)}{\sqrt{2}}$$

$$ln(1+x) \approx x$$
 for small x (i.e. dilute solution, X_A small)

$$\pi = \frac{X_A RT}{\sqrt{T}}$$

$$X_{\scriptscriptstyle A} = \frac{n_{\scriptscriptstyle A}}{n_{\scriptscriptstyle A} + n_{\scriptscriptstyle B}} \quad and \; n_{\scriptscriptstyle A} + n_{\scriptscriptstyle B} \approx n_{\scriptscriptstyle B} \quad for \; dilute \; solution$$

$$\pi \approx \frac{n_{\scriptscriptstyle A}RT}{n_{\scriptscriptstyle B}\overline{V_{\scriptscriptstyle B}}}$$

$$\pi V_{\scriptscriptstyle B} = n_{\scriptscriptstyle A} R T$$

$$\pi V_{solution} = n_{solute} RT$$

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quantitative treatment of colligative properties

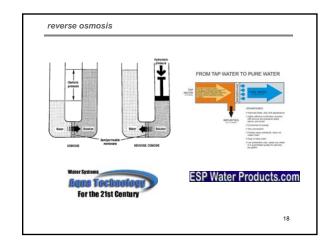
Handout #52 Colligative Properties

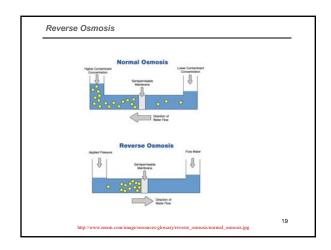
from relationships for Chem 163B final:

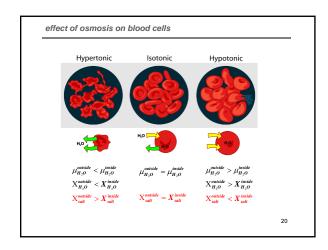
Colligative properties:

- freezing point lowering: $\gamma_g X_g = \exp \left[-\frac{\Delta \vec{H}_{factor}}{R} \left[\frac{1}{T_f} \frac{1}{T_f^*} \right] \right]$
- boiling point elevation: $\gamma_3 X_3 = \exp \left| \frac{\Delta H_{uppropulse}}{R} \right| \frac{1}{T} \frac{1}{T^*}$
- $\pi = \frac{-RT \ln(\gamma_g X_g)}{V_g}$ osmotic pressure:

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End of lecture