

Change of State Previous Year Problems

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By how much the freezing point of benzene 5.53°C , be reduced if 10g hexane added to 100 g benzene?

(ΔH_f for benzene = $9.836 \text{ kJ mol}^{-1}$).

formula $\Delta T_f = K' x_B$; $\left(K' = \frac{RT^{*2}}{\Delta H_f} \right)$

Given: $T^* = 5.53^{\circ}\text{C} + 273 \text{ K} = 278.53 \text{ K}$

$R = 8.314 \text{ J K}^{-1} \text{ mol}^{-1}$

$\Delta H_f = 9.836 \text{ kJ mol}^{-1} = 9.836 \times 10^3 \text{ J mol}^{-1}$

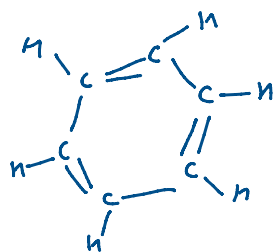
A = Solvent = Benzene
 ↑
 large quantity

B = Solute = Hexane.
 ↑
 less quantity.

no. of moles of A = $\frac{\text{given wt. of A}}{\text{Molecular wt. of A}}$

no. of moles of B = $\frac{\text{given wt. of B}}{\text{Molecular wt. of B}}$

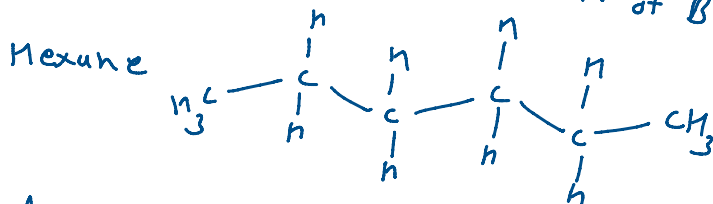
\therefore Molecular weight of benzene



$\text{C}_6\text{H}_6 = (12 \times 6) + (1 \times 6)$
 $\rightarrow = 78 \text{ gm mol}^{-1}$

$n_A = \frac{100 \text{ gm}}{78 \text{ gm mol}^{-1}}$

$n_A = 1.2820 \text{ mol}$



M. wt = $\text{C}_6\text{H}_{14} = (12 \times 6) + (1 \times 14) = 86 \text{ gm mol}^{-1}$

$\therefore n_B = \frac{10 \text{ gm}}{86 \text{ gm mol}^{-1}} = 0.1162 \text{ mol}$

$\therefore x_B = \frac{n_B}{n_A + n_B} = \frac{0.1162}{1.2820 + 0.1162}$

$x_B = 0.0831$

$\Delta T = RT^{*2}$

$$\therefore \Delta T = \frac{RT^{\circ 2}}{\Delta n_f} x_B$$

$$= \frac{8.314 \cancel{\text{ J K}^{-1} \text{ mol}^{-1}} \times (278.53 \text{ K})^2 \cancel{\text{ K}^2}}{9.836 \times 10^3 \cancel{\text{ J mol}^{-1}} \times 0.0831}$$

$$\Delta T = 5.4492 \text{ K}$$

When 5.25 g of a substance is dissolved in 565 g of benzene at 25°C, the boiling point is raised by 0.625°C. Evaluate the molecular weight of the substance. [$K_b = 2.53 \text{ K Kg mol}^{-1}$].

Formula:

$$\Delta T_b = K_b \times b$$

Given $\Delta T_b = 0.625^\circ\text{C} = 0.625 \text{ K}$ bcz, diff.

| weight of solvent = 565 gm = 0.565 kg
| weight of solute = 5.25 gm

$K_b = 2.53 \text{ K kg mol}^{-1}$

$$b = \text{molarity of solute} = \frac{\text{no. of moles of solute}}{\text{weight of solvent in kg}}$$

$$b = \frac{\text{wt. of solute} / \text{M. wt. of solute}}{0.565 \text{ kg}}$$

$$b = \frac{5.25 \text{ gm}}{\text{M. wt. of solute}} \div 0.565 \text{ kg}$$

$$b = \frac{5.25 \text{ gm}}{\text{M. wt. of solute} \times 0.565 \text{ kg}}$$

$$\Delta T_b = K_b \times b$$

$$b = \frac{\Delta T_b}{K_b}$$

$$\frac{5.25 \text{ gm}}{\text{M. wt. of solute} \times 0.565 \text{ kg}} = \frac{0.625 \text{ K}}{2.53 \text{ K kg mol}^{-1}}$$

$$\therefore \text{M. wt. of solute} = \frac{5.25 \text{ gm} \times 2.53 \text{ K kg mol}^{-1}}{0.565 \text{ kg} \times 0.625 \text{ K}}$$

$$= 37.61 \text{ gm mol}^{-1}$$

Estimate the molar solubility of oxygen in water at 25°C and partial pressure of 160 torr. (Henry's constant $K = 3.3 \times 10^7$ torr).

Given ::

$$P_B = 160 \text{ torr}$$

$$K_B = 3.3 \times 10^7 \text{ torr}$$

$$b_B = \text{molality of solute. (mol kg}^{-1}\text{)}$$

formula ::

$$P_B = K_B b_B$$

← Henry's law →

$$P_B = K_B x_B \Rightarrow x_B = \frac{P_B}{K_B}$$

$$b_B \propto x_B \Rightarrow b_B$$

ie

$$b_B = \frac{n_B}{\text{kg of solvent}}$$

$$x_B = \frac{n_B}{n_A + n_B}$$

$$b_B = \frac{P_B}{K_B} = \frac{160 \text{ torr}}{3.3 \times 10^7 \text{ torr kg mol}^{-1}} = 4.84 \times 10^{-6} \text{ mol kg}^{-1}$$

Molar solubility, expressed in mol lit⁻¹

$$\text{density of water} = 1 \text{ gm/ml} = 1 \text{ kg lit}^{-1}$$

$$\text{ie } 1 \text{ kg} = 1 \text{ lit.}$$

$$\therefore \text{molar solubility} = 4.84 \times 10^{-6} \text{ mol lit}^{-1}$$

$$\hookrightarrow = 0.00484 \times 10^{-3} \text{ mmol lit}^{-1}$$

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A water alcohol mixture is 40% in alcohol by mass, the density of water is 1 gram/cc and density of alcohol is 0.785 gram/cc. Find the total volume of 1 kg mixture [Given : Partial molar volume of water = 17.5 cc mole⁻¹, and partial molar volume of ethanol = 55.0 cc mole⁻¹].

A = solvent = water

B = solute = alcohol

Total volume of solution

$$V = n_A V_A + n_B V_B$$

Given

$$V_A = 17.5 \text{ cm}^3 \text{ mol}^{-1}$$

$$V_B = 55.0 \text{ cm}^3 \text{ mol}^{-1}$$

We have to find n_A & n_B

40% alcohol (B) & 60% water (A)

$$\text{no. of moles of A} = \frac{\text{given weight of A (gm)}}{\text{M. wt. of A (gm mol}^{-1}\text{)}} = \text{mol}$$

Total 1 kg, 60% water = 600 gm water

$$\rightarrow \frac{600 \text{ gm}}{18 \text{ gm mol}^{-1}} = 33.33 \text{ mol.}$$

$$\text{no. of moles of B} = \frac{\text{given wt of B (gm)}}{\text{M. wt of B in gm mol}^{-1}}$$

Total 1 kg, 40% alcohol = 400 gm

$$\text{M. wt of alcohol} = \text{CH}_3\text{CH}_2\text{OH} = (12 \times 2) + 16 + 1 = 46 \text{ gm mol}^{-1}$$

$$\begin{aligned} \text{M. wt of alcohol} &= \text{CH}_3\text{CH}_2\text{OH} = (12 \times 2) + 16 + 1 = 46 \text{ gmol}^{-1} \\ &\rightarrow = \frac{400 \text{ gm}}{46} = 8.69 \text{ mol} \end{aligned}$$

$$\begin{aligned} \therefore V &= n_A V_A + n_B V_B \\ &= 33.33 \times 17.5 + 8.69 \times 55.0 \\ &= 1061.225 \text{ ml} \end{aligned}$$

At 25°C the density of 50% by mass of ethanol-water mixture is 914 kg/m³. Find the Partial molar volume of ethanol.

[Partial Molar volume of water = 17.4 cm³ mole⁻¹]

Given, $V_A = 17.5 \text{ cm}^3 \text{ mol}^{-1} = \text{water}$

$V_B = ? = \text{ethanol}$

$$V = n_A V_A + n_B V_B$$

$$\therefore V_B = \frac{V - n_A V_A}{n_B}$$

Let's take 100 cm³ of solⁿ as convenient sample.

$$\text{Density} = \frac{\text{Mass}}{\text{Volume}}$$

$$\therefore \text{Mass} = \text{Density} \times \text{Volume}$$

$$= 914 \text{ kg m}^{-3} \times 100 \text{ cm}^3$$

$$\text{Mass} = \underline{0.914 \text{ gm cm}^{-3}} \times \underline{100 \text{ cm}^3}$$

$$\text{Mass} = 0.914 \text{ gm cm}^{-3} \times 100 \text{ cm}^3$$

$$\text{Total Mass} = 91.4 \text{ gm}$$

Given that 50%.

Now weight of water & weight of ethanol is

$$\text{ie } 0.5 \times 91.4 = 45.7 \text{ gm}$$

$$0.5 \times 91.4 = 45.7 \text{ gm}$$

$$\text{No. of moles of } H_2O = \frac{\text{given wt}}{\text{M. wt.}}$$

$$n_A = \frac{45.7 \text{ gm}}{18 \text{ gm mol}^{-1}}$$

$$n_A = 2.538 \text{ mol}$$

$$n_B = \frac{45.7 \text{ gm}}{\text{M. wt.}} = \frac{45.7 \text{ gm}}{46 \text{ gm mol}^{-1}}$$

$$n_B = 0.993 \text{ mol}$$

$$\therefore V_B = \frac{V - n_A V_A}{n_B} = \frac{100 - 2.538 \times 17.5}{0.993}$$

$$V_B = 55.97 \text{ cm}^3 \text{ mol}^{-1}$$