

## 2. SOLUTION

2.36 100g of liquid A (molar mass  $140 \text{ g mol}^{-1}$ ) was dissolved in 1000g of liquid B (molar mass  $180 \text{ g mol}^{-1}$ ). The vapour pressure of pure liquid B was found to be 500 torr. Calculate the vapour pressure of pure liquid A and its vapour pressure in the solution if the total vapour pressure of the solution is 475 torr.

Solution:

$$\text{No. of moles of solute, } n_2 = \frac{100}{140} = \frac{5}{7} \text{ mole}$$

$$\text{No. of moles of solvent, } n_1 = \frac{1000}{180} = \frac{50}{9} \text{ mole}$$

Mole fraction of solute,

$$x_2 = \frac{n_2}{n_1 + n_2} = \frac{5/7}{5/7 + 50/9} = 0.114$$

$$\text{Mole fraction of solvent, } x_1 = (1 - x_2) = (1 - 0.114) = 0.886$$

According to Raoult's law

$$P_A = x_A P_A^\circ = 0.114 \times P_A^\circ$$

$$P_B = x_B P_B^\circ = 0.886 \times 500 = 443 \text{ torr}$$

$$P_{\text{Total}} = P_A + P_B$$

$$475 = 0.114 P_A^\circ + 443$$

$$P_A^\circ = \frac{475 - 443}{0.114} = 280.7 \text{ torr}$$

$$\therefore P_A = 0.114 \times 280.7 = 32 \text{ torr.}$$

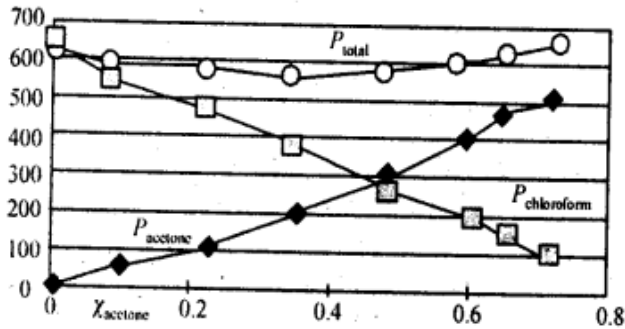
2.37 Vapour pressures of pure acetone and chloroform at 328 K are 741.8 mm Hg and 632.8 mm Hg respectively. Assuming that they form ideal solution over the entire range of composition, plot  $P_{\text{total}}$ ,  $P_{\text{chloroform}}$  and  $P_{\text{acetone}}$  as a function of  $\chi_{\text{acetone}}$ . The experimental data observed for different compositions of mixtures is:

$100 \times \chi_{\text{acetone}}$	0	11.8	23.4	36.0	50.8	58.2	64.5	72.1
$P_{\text{acetone}}/\text{mm Hg}$	0	54.9	110.1	202.4	322.7	405.9	454.1	521.1
$P_{\text{chloroform}}/\text{mm Hg}$	632.8	548.1	469.4	359.7	257.7	193.6	161.2	120.7

Plot this data also on the same graph paper. Indicate whether it has positive deviation or negative deviation from the ideal solution.

Solution:

$\chi_{\text{acetone}}$	0.0	0.118	0.234	0.360	0.508	0.582	0.645	0.721
$P_{\text{acetone}}/\text{mm Hg}$	0	54.9	110.1	202.4	322.7	405.9	454.1	521.1
$P_{\text{chloroform}}/\text{mm Hg}$	632.8	548.1	469.4	359.7	257.7	193.6	161.2	120.7
$P_{\text{total}}$	632.8	603.0	579.5	562.1	580.4	599.5	615.3	641.8



As the plot for  $P_{\text{total}}$  dips downwards, hence the solution shows negative deviation from the ideal behaviour.

Mole fraction of  $\text{C}_6\text{H}_5\text{CH}_3$ ,

$$x_T = \frac{1.087}{1.026 + 1.087} = 0.514$$

According to Raoult's Law,

$$P_B = x_B \times P_B^\circ = 0.486 \times 50.71 = 24.65 \text{ mm}$$

$$P_T = x_T \times P_T^\circ = 0.514 \times 32.06 = 16.48 \text{ mm}$$

Mole fraction of  $\text{C}_6\text{H}_6$  in vapour phase

$$= \frac{P_B}{P_B + P_T} = \frac{24.65}{24.65 + 16.48} = 0.599$$

2.38 Benzene and toluene form ideal solution over the entire range of composition. The vapour pressure of pure benzene and toluene at 300 K are 50.71 mm Hg and 32.06 mm Hg respectively. Calculate the mole fraction of benzene in vapour phase if 80g of benzene is mixed with 100g of toluene.

Solution:

$$\text{Molar mass of } \text{C}_6\text{H}_6 = 78 \text{ g mol}^{-1}$$

$$\text{Molar mass of } \text{C}_6\text{H}_5\text{CH}_3 = 92 \text{ gmol}^{-1}$$

$$\text{No. of moles of } \text{C}_6\text{H}_6 = \frac{80}{78} = 1.026 \text{ mole}$$

$$\text{No. of moles of } \text{C}_6\text{H}_5\text{CH}_3 = \frac{100}{92} = 1.087 \text{ mole}$$

Mole fraction of  $\text{C}_6\text{H}_6$ ,

$$x_B = \frac{1.026}{1.026 + 1.087} = 0.486$$

2.39 The air is a mixture of a number of gases. The major components are oxygen and nitrogen with approximate proportion of 20% is to 79% by volume at 298 K.

The water is in equilibrium with air at a pressure of 10 atm. At 298 K if the Henry's law constants for oxygen and nitrogen are  $3.30 \times 10^7$  mm and  $6.51 \times 10^7$  mm respectively, calculate the composition of these gases in water.

Solution:

Air containing 20% oxygen and 79% nitrogen by volume means

$$\begin{aligned} \text{Partial pressure of O}_2 (P_{\text{O}_2}) &= \frac{20}{100} \times 10 = 2 \text{ atm} \\ &= 2 \times 760 \text{ mm} = 1520 \text{ mm} \end{aligned}$$

$$\begin{aligned} \text{Partial pressure of N}_2 (P_{\text{N}_2}) &= \frac{79 \times 10}{100} = 7.9 \text{ atm} \\ &= 7.9 \times 760 \text{ mm} = 6004 \text{ mm} \end{aligned}$$

According to Henry's Law,

$$P_{\text{O}_2} = K_H x_{\text{O}_2}$$

$$x_{\text{O}_2} = \frac{P_{\text{O}_2}}{K_H} = \frac{1520}{3.30 \times 10^7} = 4.61 \times 10^{-5}$$

$$P_{\text{N}_2} = K_H x_{\text{N}_2}$$

$$x_{\text{N}_2} = \frac{P_{\text{N}_2}}{K_H} = \frac{6004}{6.51 \times 10^7} = 9.22 \times 10^{-5}$$

2.40 Determine the amount of  $\text{CaCl}_2$  ( $i = 2.47$ ) dissolved in 2.5 litre of water such that its osmotic pressure is 0.75 atm at  $27^\circ\text{C}$ .



Solution:

Using relation,  $\pi = iCRT = i \frac{n}{V} RT$

$$n = \frac{\pi V}{iRT} = \frac{0.75 \times 2.5}{2.47 \times 0.0821 \times 300} = 0.0308 \text{ mole}$$

Molar mass of  $\text{CaCl}_2 = 40 + 2 \times 35.5 = 111 \text{ g mol}^{-1}$

$\therefore$  Amount of  $\text{CaCl}_2$  dissolved

$$= 0.0308 \times 111 = 3.42 \text{ g.}$$

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