

LAKSHYA JEE

LAKSHYA KO HAR HAAL ME PAANA HAI



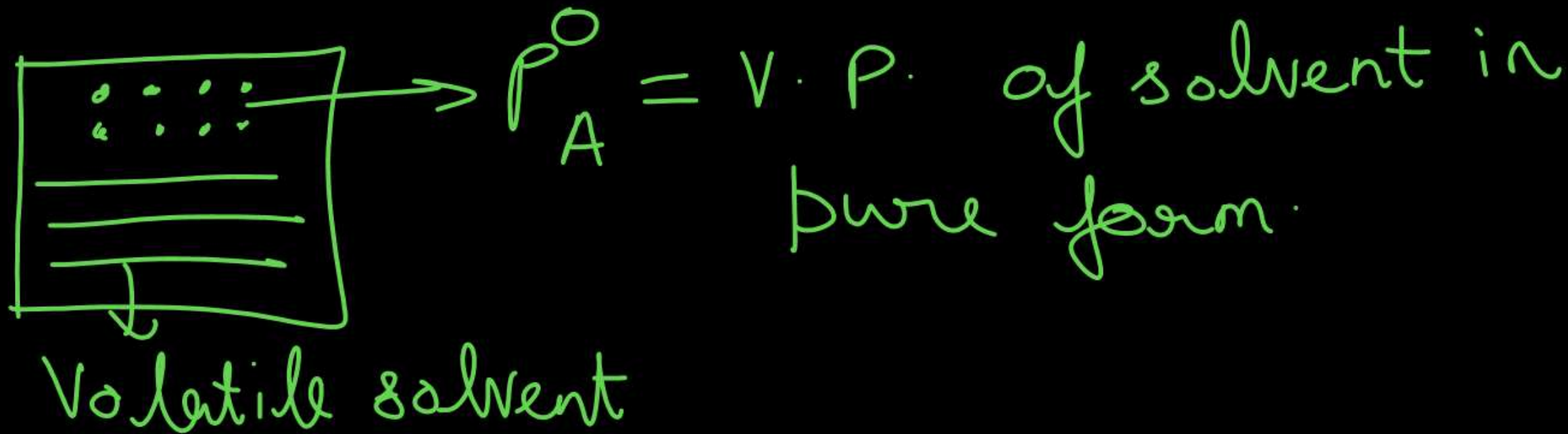
SOLUTION


By
Amit Mahajan Sir

- **RAOULT'S LAW** ✓
- **IDEAL SOLUTION** ✓



Raoult's law :

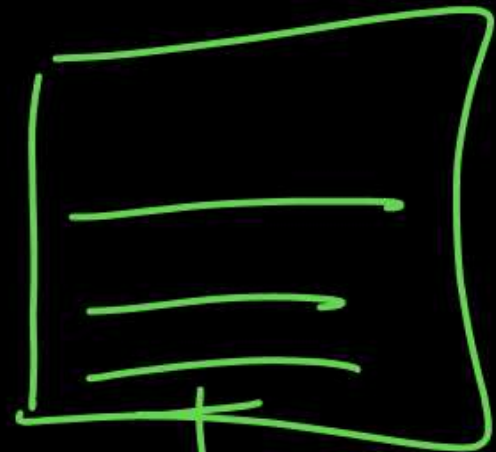




A rectangular container is shown with a horizontal line near the top. Above this line are four small dots. Below the line is a wavy line representing a liquid surface. An arrow points downwards from the wavy line to the text "Solvent".

Solvent
↓
Volatile
 $P_A^0 \neq 0$

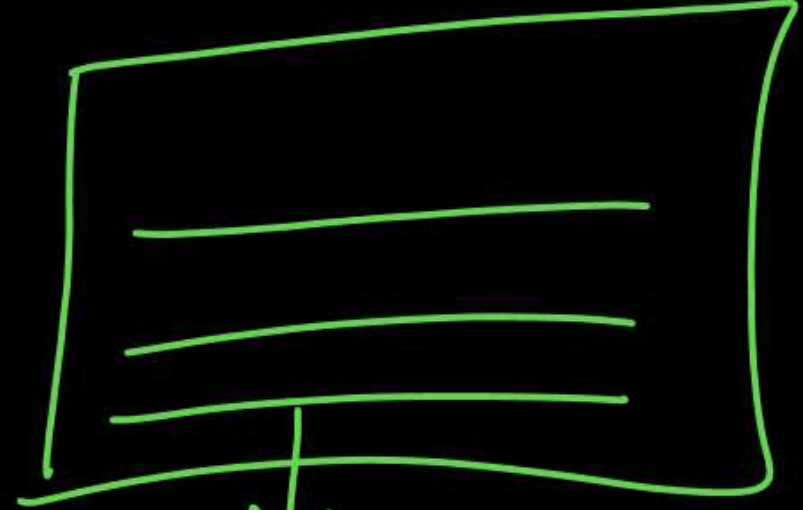
+



A rectangular container is shown with three horizontal lines. An arrow points downwards from the bottom line to the text "Solute".

Solute

→



A rectangular container is shown with three horizontal lines. An arrow points downwards from the bottom line to the text "Solution".

Solution.

$P_A = \text{V.P. of solvent}$
in solution

$$P_A = P_A^0 x_A$$

Raoult's law :-

V.P. of any Component in solution is equal to product V.P. of Component in pure form & it's mole-fraction in Solution

$$P_A = P_A^{\circ} X_A$$

P_A = V.P. of A in solution.

$$P_B = P_B^0 X_B$$



v.p. of solute in solution.

P_B^0 = v.p. of solute in pure form

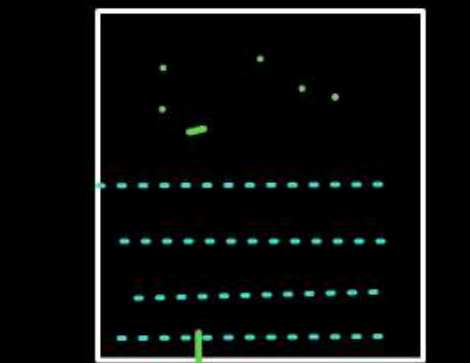
X_B = mole-fraction of B in solution.

$$\textcircled{x_A} + x_B = 1 \Rightarrow x_B = 1 - x_A$$

$$\frac{n_A}{n_A + n_B} + \frac{n_B}{n_A + n_B} = 1$$

$$\frac{\cancel{n_A} + n_B}{\cancel{n_A} + n_B} = 1$$

@ Raoult's law for Volatile Solute

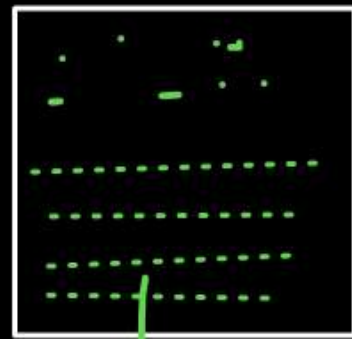


Solvent

↓
Volatile

↓
 $P_A \neq 0$

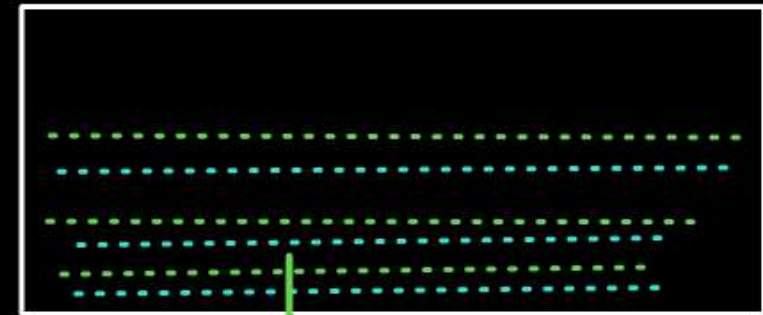
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Solute

↓
Volatile

↓
 $P_B \neq 0$



Solution

$P_S = V.P. \text{ of solution}$

$$P_S = P_A + P_B$$

$$P_S = P_A^{\circ} \chi_{A} + P_B^{\circ} \chi_{B}$$

$$P_S = P_A^{\circ} (1 - \chi_B) + P_B^{\circ} \chi_B$$

$$P_A = P_A^{\circ} \chi_A$$

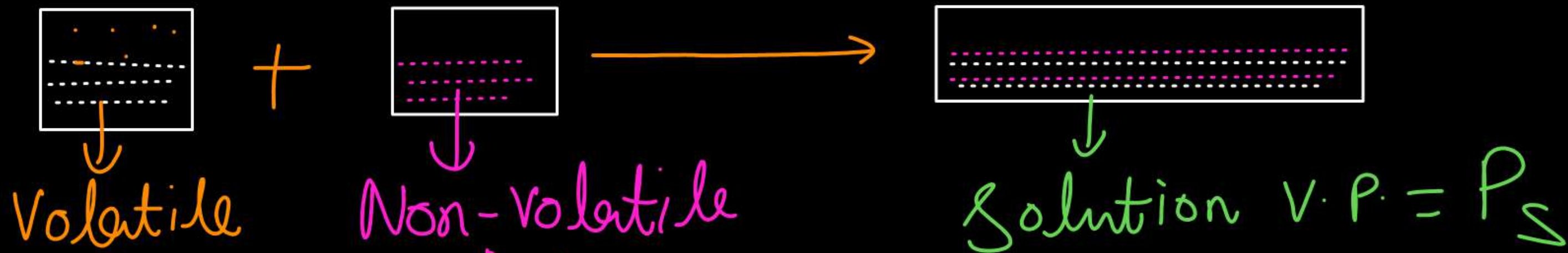
$$P_B = P_B^{\circ} \chi_B$$

$$P_S = P_A^{\circ} - P_A^{\circ} \chi_B + P_B^{\circ} \chi_B$$

$$P_S = (P_B^{\circ} - P_A^{\circ}) \chi_B + P_A^{\circ}$$

$$P_S = P_A^{\circ} + (P_B^{\circ} - P_A^{\circ}) \chi_B$$

⑥ Raoult's law for non-volatile solute.



$P_A^0 \neq 0$

$P_B^0 = 0$

$$P_S = P_A^0 X_A + P_B^0 X_B$$

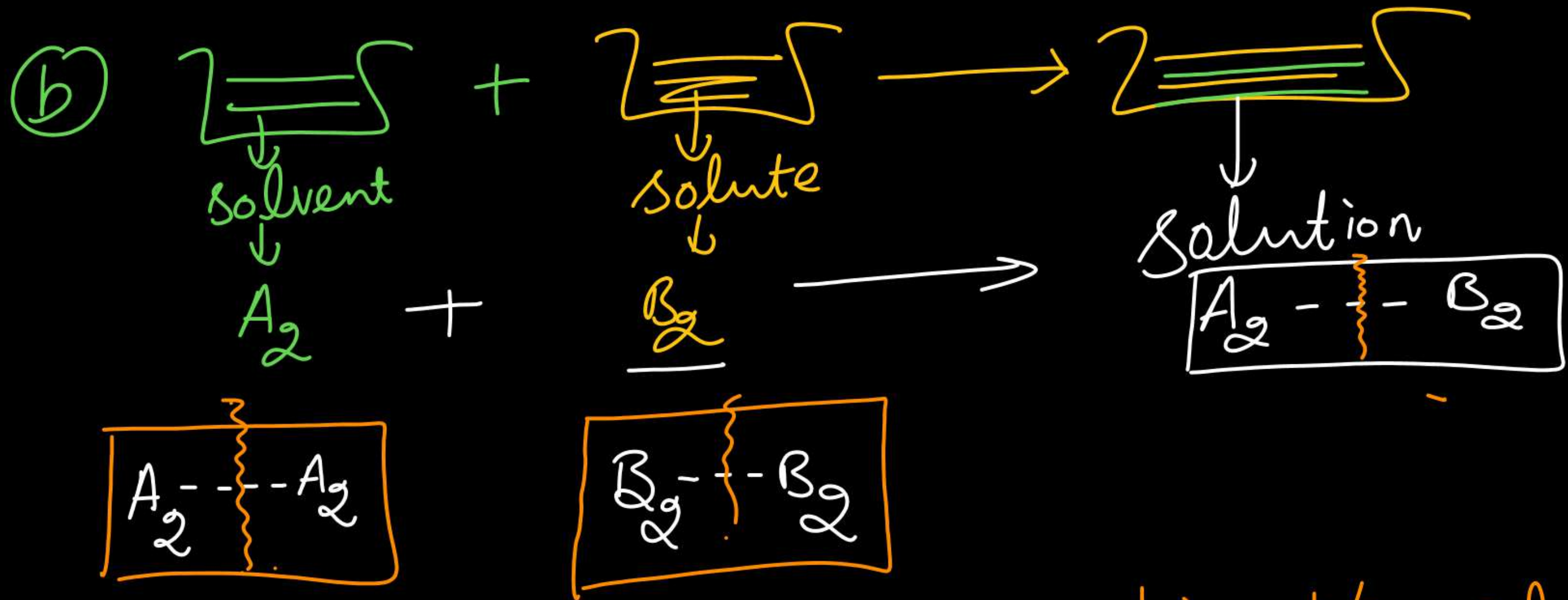
as $P_B^0 = 0$

$$P_S = P_A^0 \chi_A$$

Ideal solution :-

(a) Solutions which obey Raoult's law at all temperature & pressure.

$$P_S = P_A^{\circ} X_A + P_B^{\circ} X_B$$



When Forces of interaction b/w solvent solvent or solute-solute are similar as that of solution.

Properties of Ideal Solution

$$(a) P_S = P_A^{\circ} x_{oA} + P_B^{\circ} x_{oB}$$

$$(b) \Delta H_{\text{mixing}} = 0 \quad (c) \Delta V_{\text{mixing}} = 0$$

$$(d) \Delta G_{\text{mixing}} = (-)ve \quad (e) \Delta S_{\text{mixing}} = (+)ve$$

$$(f) \Delta U_{\text{mixing}} = 0$$

$$\Delta H_{\text{mix}} = \Delta U_{\text{mix}} + P \Delta \overset{\checkmark}{V}_{\text{mix}}$$

g

$$\Delta U_{\text{mix}} = 0$$

↓
Ideal solution

for ex - same homologous series.



↓
Ideal solution

✓
a) n-Octane + n-nonane

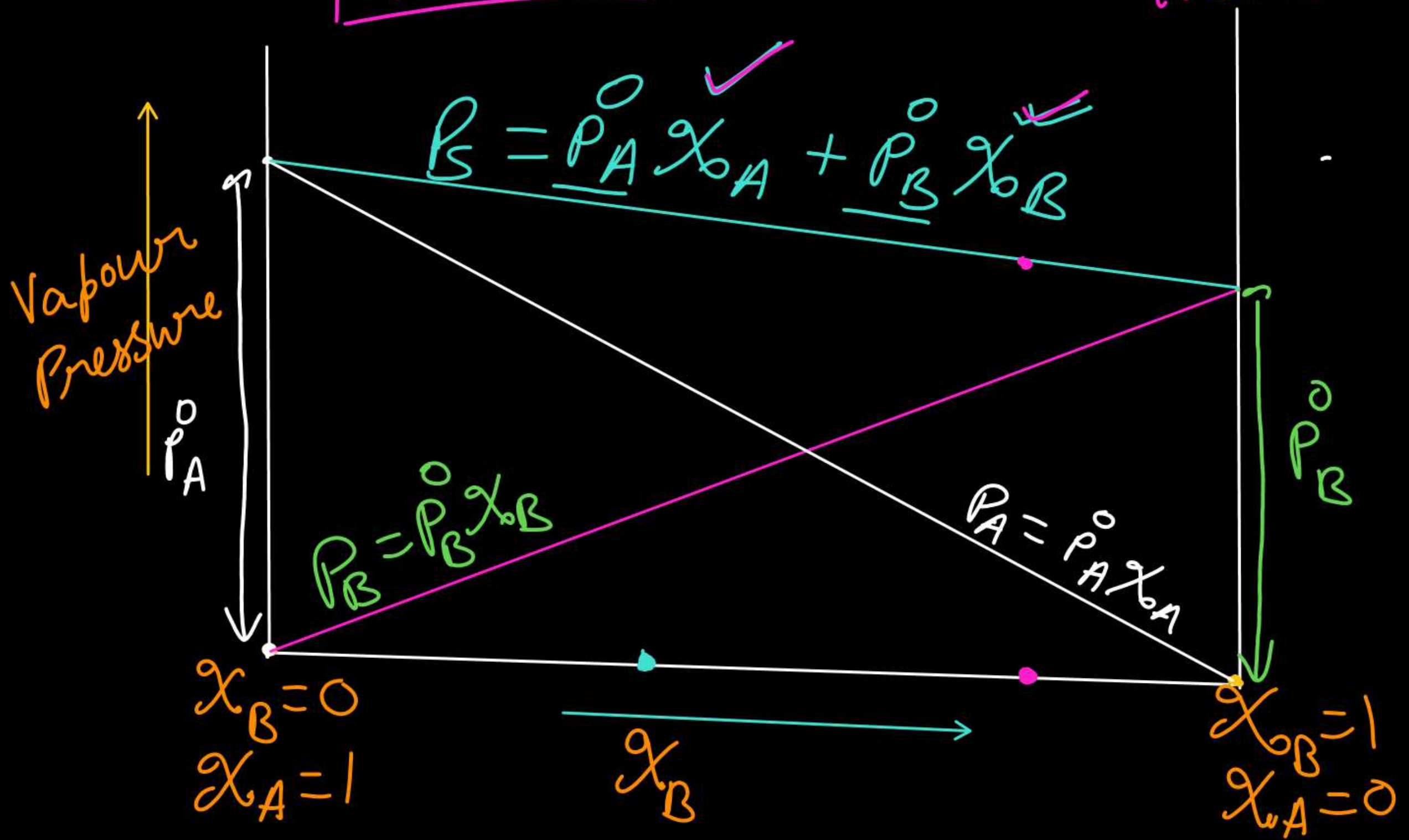
b) C_2H_5Br + C_2H_5I

c) C_2H_5Br + C_3H_7Br

d) n-Heptane + n-octane

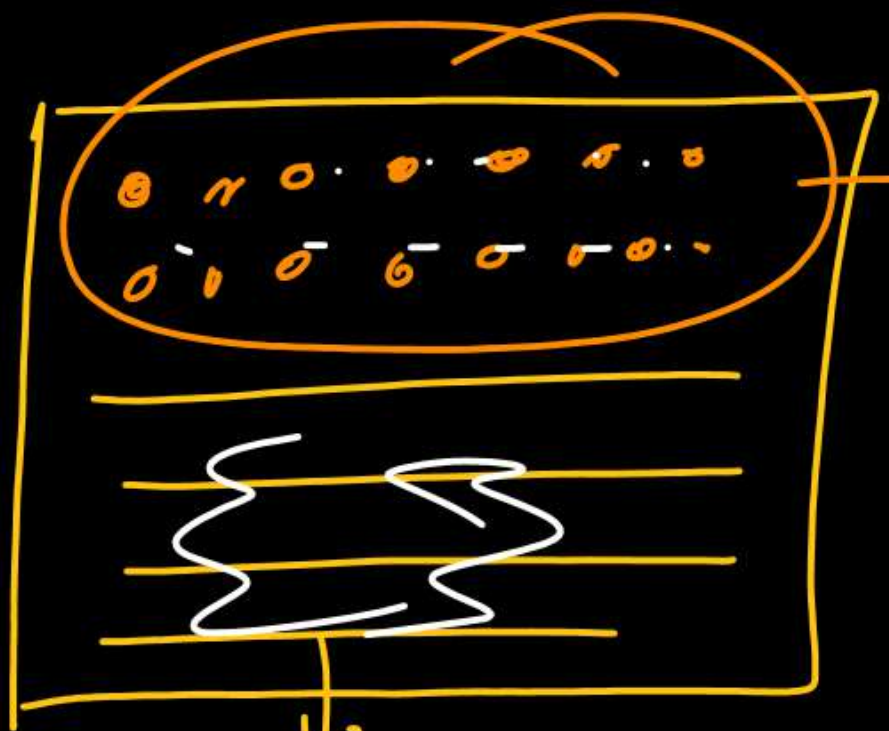
e)  + 
Benzene + Toluene

$P_A^0 > P_B^0 \rightarrow$ A is more volatile than B



$P_B = P_B^0 x_B$

$P_A = P_A^0 x_A$



Vapours of A & B

A + B
↓
Solution.

x_A & x_B

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

Mole fraction of A & B in Vapor phase is denoted by y_A & y_B

Dalton's law of Partial Pressure.

$$P_A = Y_A P_S$$

$$Y_A = \frac{P_A}{P_S} = \frac{P_A^0 X_A}{P_A^0 X_A + P_B^0 X_B}$$

$$Y_B = \frac{P_B}{P_S} = \frac{P_B^0 X_B}{P_A^0 X_A + P_B^0 X_B}$$

$$X_A + X_B = 1$$

||| clearly

$$Y_A + Y_B = 1$$

$$\frac{P_A^0 X_A}{P_S} + \frac{P_B^0 X_B}{P_S} = 1$$

$$\frac{P_A^0 X_A + P_B^0 X_B}{P_S} = 1$$

$$\frac{P_S}{P_S} = 1$$

$$Y_A + Y_B = 1$$

$$Y_B = 1 - Y_A$$


$\downarrow B$ $\downarrow A$
Benzene and toluene form a ideal solution and V.P. of pure benzene and toluene are 160 mm of Hg and 60 mm of Hg. Calculate partial pressure of benzene and toluene and total pressure also

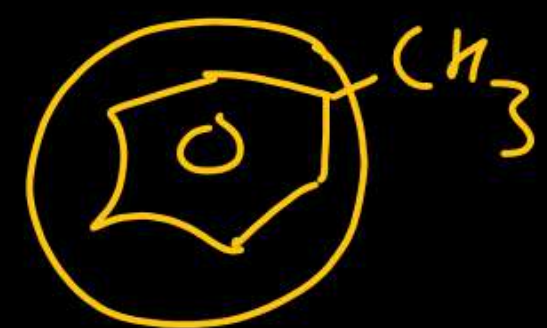
- (a) Containing equal mass of both benzene and toluene
- (b) Containing equal molecules of both benzene and toluene
- (c) Containing 1 mole of benzene and 4 moles of toluene
- (d) Also calculate mole fraction of Benzene and toluene in vapour phase if equal moles of benzene and toluene mixed

$$\begin{array}{l|l|l}
 p_B^0 = 160 \text{ mm of Hg} & P_A = ? & P_S = ? \\
 p_A^0 = 60 \text{ mm of Hg} & P_B = ? &
 \end{array}$$



(a) Containing equal mass of Benzene & toluene

B = Benzene  $C_6H_6 \Rightarrow$ Molar mass of B = 78g

A = Toluene  $C_7H_8 \Rightarrow$ Molar mass of A = 92g

$$W_A = W_B = 7176$$

$$M_A = 92, M_B = 78g$$

$$P_A = P_A^0 \chi_A$$

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

$$n_A = \frac{W_A}{M_A} = \frac{7176}{92} = 78$$

$$n_B = \frac{W_B}{M_B} = \frac{7176}{78} = 92$$

$$X_A = \frac{n_A}{n_A + n_B} = \frac{78}{92 + 78} = \frac{78}{170} = 0.46$$

$$X_B = 1 - X_A$$

$$X_B = 0.54$$

$$P_S = P_A^{\circ} \chi_{0A} + P_B^{\circ} \chi_{0B}$$

$$P_S = (60 \times 0.46 + 160 \times 0.54) \text{ mm of Hg}$$

(b) Same molecules \therefore same no. of moles

$$n_A = n_B = x$$

$$x_A = \frac{x}{x+x} = \frac{x}{2x} = \frac{1}{2} = 0.5$$

$$x_B = 1 - x_A = 0.5$$

$$P_A = P_A^{\circ} x_A = 60 \times 0.5 = 30 \text{ mm of Hg}$$

$$P_B = P_B^{\circ} x_B = 160 \times 0.5 = 80 \text{ mm of Hg}$$

$$P_{\Sigma} = P_A^{\circ} x_A + P_B^{\circ} x_B = 30 + 80 = 110 \text{ mm of Hg}$$

$$\textcircled{c} \quad n_A = 4, \quad n_B = 1$$

$$x_A = \frac{4}{5} = 0.8$$

$$x_B = 1 - x_A = 0.2$$

$$P_A = P_A^0 x_A = 60 \times 0.8 = 48 \text{ mm of Hg}$$

$$P_B = P_B^0 x_B = 160 \times 0.2 = 32 \text{ mm of Hg}$$

$$P_S = 48 + 32$$

$$P_S = 80 \text{ mm of Hg}$$

$$\textcircled{d} \quad n_A = n_B$$

$$x_A = \frac{1}{2}$$

$$x_B = \frac{1}{2}$$

$$P_S = 110 \text{ mm of Hg}$$

$$P_A = 30 \text{ mm of Hg}$$

$$P_B = 80 \text{ mm of Hg}$$

$$Y_A = \frac{P_A}{P_S} = \frac{30}{110} \text{ mm of Hg}$$

$$Y_B = 1 - Y_A = 1 - \frac{3}{11}$$

$$Y_B = \frac{8}{11} \text{ mm of Hg}$$

At a given temperature, the vapour pressure in mm of Hg of a solution of two volatile liquids A and B is given by equation $P = 120 - 80 \chi_B$

Calculate V.P. of pure A and B at same temperature

$$P_S = 120 - 80 \chi_B$$

$$P_S = P_A^0 + (P_B^0 - P_A^0) \chi_B$$

$$P_A^0 = 120 \text{ mm of Hg}$$

$$P_A^0 = ? \quad , \quad P_B^0 = ?$$

$$P_B^0 - P_A^0 = -80$$

$$P_B^0 - 120 = -80$$

$$P_B^0 = 40 \text{ mm of Hg}$$



Two liquids A and B form an Ideal solution at 300 K the V.P. of solution having 1 mole of A and 3 mole of B is 550 mm of Hg. At same temperature if 1 more mole of B is added to solution, V.P. of solution increases by 10 mm of Hg. Determine V.P. of A and B in pure state. [IIT]

Ans

$$n_A = 1, n_B = 3$$

$$P_S = 550 \text{ mm of Hg}$$

$$\%_A = \frac{1}{4} = 0.25$$

$$\%_B = 0.75$$

$$n_A = 1$$

$$n_B = 4$$

$$P'_S = 560 \text{ mm of Hg}$$

$$\%'_A = 0.2$$

$$\%'_B = 0.8$$



$$\text{Let } P_A^0 = a$$

$$P_B^0 = b$$

L.H.S.

$$P_S = P_A^0 \gamma_{A'} + P_B^0 \gamma_{B'}$$

$$550 = 0.25a + 0.75b$$

$$P_S' = P_A^0 \gamma_{A'} + P_B^0 \gamma_{B'}$$

$$560 = \underline{0.2a} + \underline{0.8b}$$

$$550 = \underline{0.25a} + \underline{0.75b}$$

For an ideal solution, the correct option is

(a) $\Delta_{\text{mix}} S = 0$ at constant T and P ✗

(b) $\Delta_{\text{mix}} V \neq 0$ at constant T and P

✓ (c) $\Delta_{\text{mix}} H = 0$ at constant T and P

(d) ✗ $\Delta_{\text{mix}} G = 0$ at constant T and P



Which one is not equal to zero for an ideal solution?

(a) $\Delta P = P_{\text{observed}} - P_{\text{Raoult}}$

(b) ΔH_{mix}

(c) ΔS_{mix}

(d) ΔV_{mix}

$$\Delta P = P_{\text{observed}} - P_{\text{Raoult's law}} = 0$$



Thank You Lakshyians