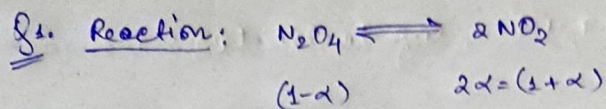


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$$\alpha = \frac{D_0 - D}{(2-1)D}$$

D_0 : Expected Vapour density from the formula weight

$$= \frac{D_0 - D}{D}$$

D : Actual Vapour density at the given T and P.

The calculated density of N_2O_4 : $D_0 = \frac{M}{2} = 46$

Observed Density : $D = 30.2$

Hence, $\alpha = \frac{46 - 30.2}{30.2} = \underline{\underline{0.523}}$

Hence, the fracⁿ of gram-moles of N_2O_4 decomposed = 0.523

Thus, percentage of NO_2 molecules by wt. = $\underline{\underline{52.3\%}}$

Now, the ratio of grammoles of NO_2 & N_2O_4 in the mixture

$\Rightarrow \frac{2\alpha}{1+\alpha} : \frac{1-\alpha}{1+\alpha}$ (This is same as the ratio of their volumes)

Therefore, % age of NO_2 by volume = $\frac{2 \times 0.523}{1 + 0.523} \times 100\%$
 $= \underline{\underline{68.7\%}}$

Q2. $400\text{g H}_2 + 1400\text{g N}_2$ ($P = 200\text{ atm}$)

$$P = P_{\text{H}_2} + P_{\text{N}_2}$$

$\text{H}_2 : 200\text{ mol}$ and $\text{N}_2 : 50\text{ mol}$.

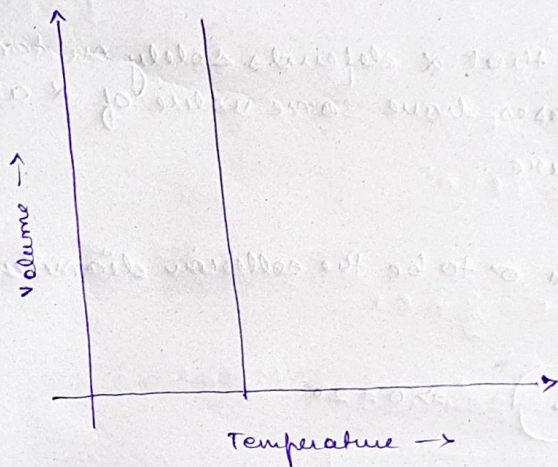
$$\therefore P_{\text{H}_2} = \frac{200}{250} \times 200\text{ atm.} \quad \left(P_a = \frac{n_a}{n} \times P \right)$$

$$= \underline{\underline{160\text{ atm}}}$$

$$P_{\text{N}_2} = \frac{50}{250} \times 200\text{ atm}$$

$$= \underline{\underline{40\text{ atm}}}$$

Q3. (i) Volume vs. Temperature Graph for Isothermal Expansion of a gas.



The graph stays vertical as the process is "isothermal" i.e., the temperature remains constant throughout. The pressure and volume are the only changing parameters in the graph / isothermal expansion.

(ii) α : Coefficient of thermal expansion.

According to definition,

$$\alpha = \frac{1}{V} \left(\frac{\partial V}{\partial T} \right)_P$$

For 1 mol ideal gas, $PV = RT$

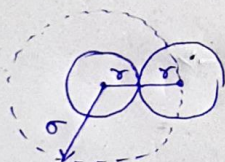
$$\text{Hence, } \left(\frac{\partial V}{\partial T} \right)_P = \frac{R}{P}$$

$$\text{So, } \alpha = \frac{1}{V} \times \frac{R}{P} = \frac{R}{RT} = \frac{1}{T}$$

$$\text{Or, } \boxed{\alpha = \frac{1}{T}}$$

This proves that α depends solely on temperature and all gases have same value of α at a given temperature.

Q4. Consider σ to be the collision diameter.



$$\Rightarrow \sigma = 2r$$

The sphere of radius σ occupies a space unavailable for a pair of molecules.

Thus, Excluded Volume = $\frac{4}{3} \pi \sigma^3$ for one pair of molecules.

$$\Rightarrow \text{Effective Volume of a single molecule} = \frac{1}{2} \times \frac{4}{3} \pi \sigma^3$$

$$= \frac{2}{3} \pi \sigma^3$$

and $b = \frac{2}{3} \pi N_A \sigma^3$ ~~Excluded volume correction~~
(b : Van der Waals' Volume correction Factor)

which is the effective volume of Avogadro no. of molecules present in 1 mol gas.

$$\text{Thus, } b = \frac{2}{3} \pi N_A \sigma^3 = 4 \times \frac{4}{3} \pi N_A \sigma^3$$

\therefore , Effective volume "b", is four times the actual volume of 1 mol gas molecules.

Q5. $T = 40^\circ\text{C} = 273 + 40 = 313\text{K}$

$$PV = nRT$$

$$n = \frac{100}{44} = 2.2727$$

$$V = 5\text{L} \Rightarrow P = \frac{nRT}{V} \approx \underline{11.68\text{ atm}} \quad (\text{Ideal}).$$

$$\left(P_i + \frac{an^2}{V^2}\right)(V_i - nb) = nRT \quad \begin{aligned} (a &= 3.59\text{ atm L}^2\text{ mol}^{-2}) \\ (b &= 0.43\text{ L mol}^{-1}) \end{aligned}$$

\Rightarrow Substituting all values, we get,

$$P_{\text{real}} = \underline{13.7765\text{ atm}} \quad (\text{Real})$$

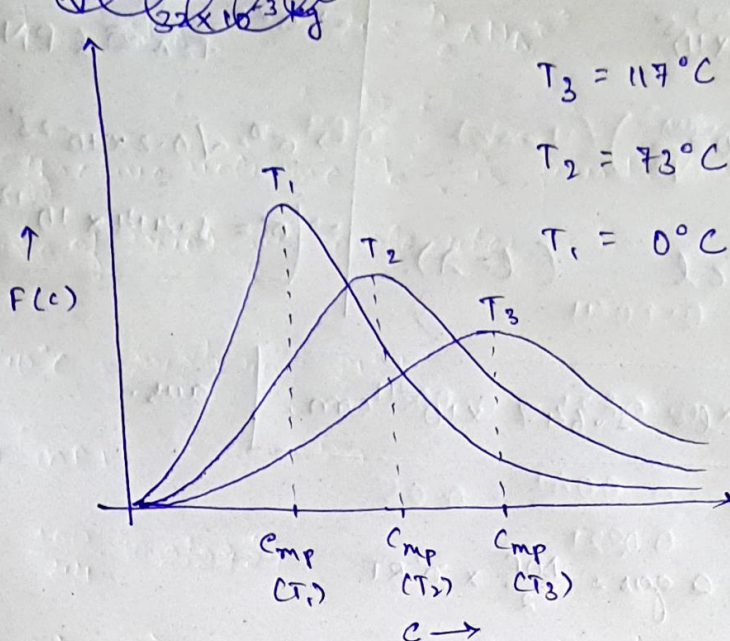
$$\underline{\underline{Q7.}} \quad c_{mp} = \sqrt{\frac{2RT}{M}} = \sqrt{\frac{2 \times 8.314 \times 300 \times 10^7}{32}}$$

$$\approx 39482 \text{ cm s}^{-1}$$

$$\approx \underline{\underline{394.83 \text{ ms}^{-1}}}$$

ans.

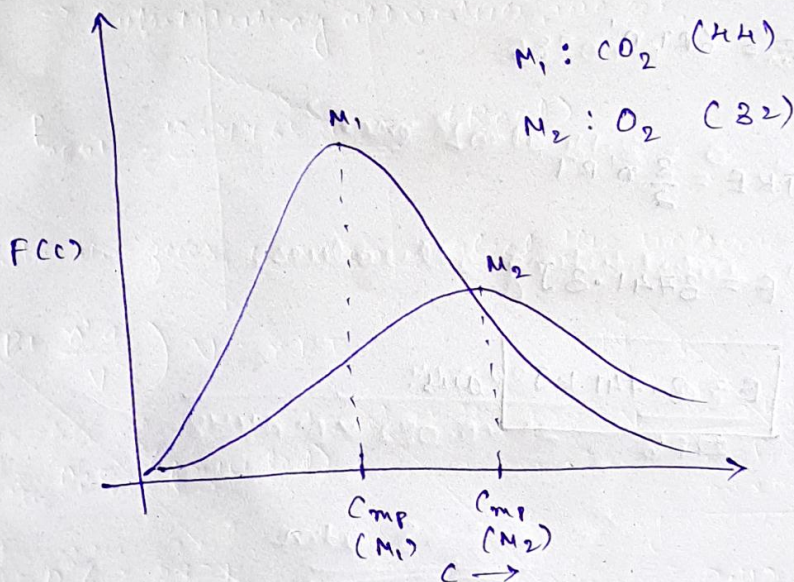
Q8. (a)



c : speed.

$F(c)$: Maxwell's speed distribution function

(b)



Q9. Given

$$\eta = 8.4 \times 10^{-5} \text{ poise}$$

$$\bar{c}_{\text{avg}} = 1.7 \times 10^5$$

$$\rho = 9 \times 10^{-5}$$

$$\eta = \frac{1}{3} \rho \bar{c} l \quad (l = \lambda)$$

$$\Rightarrow \boxed{\lambda = l = 1.647 \times 10^{-5} \text{ cm.}} \quad \text{ans.}$$

$$l = \frac{RT}{\sqrt{2} \pi \sigma^2 P N_A}$$

$$\Rightarrow \sigma^2 = \frac{RT}{\sqrt{2} \pi l P N_A}$$

$$\Rightarrow \sigma = \sqrt{0.2 \times 10^{-23}}$$

$$\Rightarrow \boxed{\sigma = 1.414 \times 10^{-12} \text{ cm.}}$$

Q10. KE for a gas = $\frac{\text{DOF}}{2} \times n' RT$

$$n' = \frac{8.5}{17}$$

Molar mass of $\text{NH}_3 = 17$.

$$\text{TKE} = \frac{3}{2} RT n'$$

$$\text{RKE} = \frac{3}{2} RT n'$$

$$\text{VKE} = 6 RT n'$$

$$\therefore \text{TKE} = \frac{3}{2} n' RT$$

$$\Rightarrow E = 3741.8 \text{ J}$$

$$\Rightarrow \boxed{E = 3.741 \text{ kJ}} \quad \text{ans.}$$

Q11. Rise in capillary Tube

$$h = \frac{2\tau}{\rho g}$$

$$g = 980 \text{ cm s}^{-2}$$

$$h = 6.30 \text{ cm}$$

T: Surface Tension

ρ : density

$$r: \text{radius} = \frac{0.21}{2} \text{ mm}$$

$$\approx 0.01 \text{ cm}$$

$$6.30 = \frac{2 \times T}{0.79 \times 0.01 \times 980}$$

$$\Rightarrow \boxed{T = 24.33 \text{ dyne cm}^{-1}} \quad \text{ans.}$$

$$\underline{\underline{Q12.}} \quad \left[P + \frac{0.00786}{V^2} \right] (V - 0.0024) = 0.0041 (273 + t)$$

We know,

$$\left(P + \frac{an^2}{V^2} \right) (V - nb) = nRT$$

$$nR = 0.0041$$

$$\Rightarrow n \times 0.0821 = 0.0041$$

$$\Rightarrow n = \frac{0.0041}{0.0821} = 0.05$$

$$nb = 0.00224$$

$$\Rightarrow b = \frac{0.00224}{0.05} = 0.0448 \text{ L mol}^{-1}$$

$$an^2 = 0.00786$$

$$\Rightarrow a = 3.14 \text{ atm L}^2 \text{ mol}^{-2}$$

Now, substituting all values,

$$T_c = \frac{8a}{27Pb} = \frac{8 \times 3.14}{27 \times 0.082 \times 0.0448}$$

$$= \underline{\underline{253.58 \text{ atm}}}$$

Q6. $\left(p + \frac{an^2}{V^2}\right) (V - nb) \overset{\text{neglect}}{=} nRT$

$$\Rightarrow Z = 0.5$$

$$T = 273K$$

$$Z = \frac{P}{P_i} \Rightarrow Z P_i = P$$

$$\Rightarrow Z P_i = \left(P_i + \frac{an^2}{V^2}\right)$$

$$\Rightarrow \boxed{a = 1.255} \text{ atm}.$$