

Gas Diffusion Coefficient: Measurement & Prediction

| | Gas | Liquids |
|------------|--------------------------|---|
| D_{AB} : | 0.1-1 cm ² /s | (0.1-1) × 10 ⁻⁵ cm ² /s |

Gases: (i) $D_{AB} \propto \frac{T^{1.5-1.75}}{P}$

(ii) Larger gas molecules diffuse slowly compared to the smaller molecules

Measurement of Gas Diffusion Coefficient-

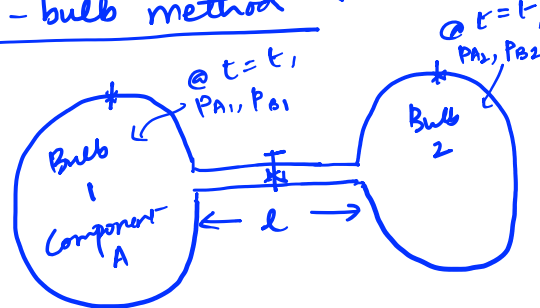
(i) Twin-bulb method :

Time $t=0$:

Comp

Bulb 1: $P_A = P$
 $P_B = 0$

Bulb 2: $P_A = 0$
 $P_B = P$



Initial condition @ $t=0$
(Before the valve connecting the bulb is opened)

Assumptions

(i) Volume of bulbs 1 & 2 is large compared to the tube connecting them

(ii) Bulbs 1 & 2 are "well mixed"

(iii) Temp. is constant

(iv) Total pressure remains same (=P).

(v) Steady state

• At $t > 0$, we open the valve connecting the two bulbs. As a result, components A & B start to diffuse from bulb 1 & 2, respectively.

• Since the total pressure is same in both the bulbs, equimolar counter-diffusion can be assumed in the tube connecting them.

• Flux for equimolar counter-diffusion (@ steady state) =

$$N_A = \frac{D_{AB} (P_{A1} - P_{A2})}{RTl}$$

• Cross section ^{area} of the tube = 'a'

⇒ Rate of mass transfer = $a N_A = -a N_B$

↙ cross section area of the tube.

Relating change in composition of bulbs 1 & 2 with the flux.

V_1 = Volume of bulb 1
 C_{A1} = Conc. of component A in bulb 1 @ $t = t$.

$$-V_1 \frac{dC_{A1}}{dt} = a N_A \Rightarrow -\frac{V_1}{RT} \frac{dP_{A1}}{dt} = a N_A \quad \text{---(i)}$$

Similarly: $V_2 \frac{dC_{A2}}{dt} = a N_A \Rightarrow \frac{V_2}{RT} \frac{dP_{A2}}{dt} = a N_A \quad \text{---(ii)}$

(i) & (ii)

$$\Rightarrow -\frac{d(P_{A1} - P_{A2})}{dt} = a RT \left(\frac{1}{V_1} + \frac{1}{V_2} \right) N_A$$

$$\Rightarrow -\frac{d(P_{A1} - P_{A2})}{dt} = \frac{a D_{AB} (P_{A1} - P_{A2})}{l} \left(\frac{1}{V_1} + \frac{1}{V_2} \right)$$

$$\Rightarrow \int \frac{-d(P_{A1} - P_{A2})}{(P_{A1} - P_{A2})} = \int_0^t \frac{a D_{AB}}{l} \left(\frac{1}{V_1} + \frac{1}{V_2} \right) dt$$

$$\Rightarrow \ln \left(\frac{P}{P_{A1}' - P_{A2}'} \right) = \frac{a D_{AB}}{l} \left(\frac{1}{V_1} + \frac{1}{V_2} \right) t$$

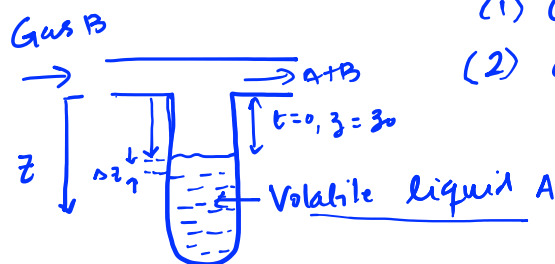
$$\underline{D_{AB}} = \frac{l}{a \left(\frac{1}{V_1} + \frac{1}{V_2} \right)} \ln \left(\frac{P}{P_{A1}' - P_{A2}'} \right)$$

$t=0$: $P_{A1} - P_{A2} = P - 0 = P$

@ $t = t$: $P_{A1} - P_{A2} = P_{A1}' - P_{A2}'$

- P_{A1}' = partial pressure of A in bulb 1 after time t
- P_{A2}' = partial pressure of A in bulb 2 after time t .

2) Stefan Tube:



- This method can be applied for:
- (1) Component A is volatile
 - (2) Component B is not soluble in Component A

Measure the change in liquid level vs time.

$N_B = 0$ { This case is similar to : "Diffusion of A in non-diffusing B"

$$N_A = \frac{D_{AB} P}{RTz} \ln \left(\frac{P - P_{A2}}{P - P_{A0}} \right)$$

Lets assume the cross section of the tube = 'a'

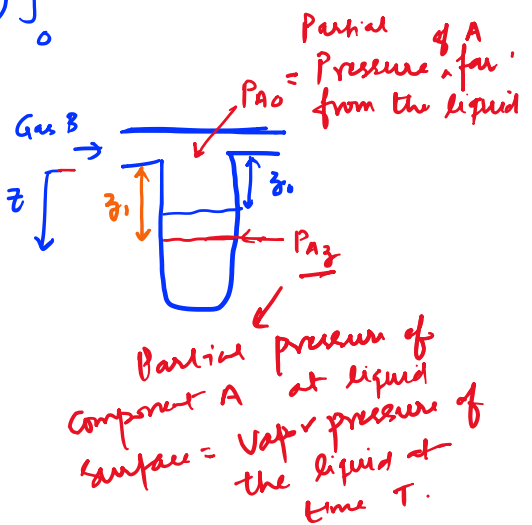
$$\frac{a dz \delta}{m_A} = a N_A dt$$

↑
mole of component
A evaporated in time
'dt'

$$\frac{a \delta dz}{m_A} = a \frac{D_{AB} P}{RTz} \ln \left(\frac{P - P_{A2}}{P - P_{A0}} \right) dt$$

$$\Rightarrow \frac{\delta_A}{m_A} \int_{z_0}^{z_1} z dz = \frac{D_{AB} P}{RT} \ln \left(\frac{P - P_{A2}}{P - P_{A0}} \right) \int_0^t dt$$

$$\Rightarrow D_{AB} = \frac{\delta_A RT (z_1^2 - z_0^2)}{2 P \ln \left(\frac{P - P_{A2}}{P - P_{A0}} \right) m_A t}$$



—————X—————X—————

Diffusion in liquids :

$$N_A = x_A (N_A + N_B) - \frac{D_{AB}}{m} \left(\frac{\delta}{m} \right)_{av} \frac{dx_A}{dz}$$

$\left(\frac{\delta}{m} \right)_{av}$ = Average molar concentration