

الجامعة التكنولوجية

قسم الهندسة الكيمائية

المرحلة الثالثة

انتقال كتلة

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(*) What is chemical Engineering ?

Chemical Engineering deals with industrial process in which raw materials are converted or separated into useful products.

(*) Who is the Chemical Engineer ?

An engineer, who apply the chemistry of a particular industrial process through the use of scientific and engineering principles.

He must **develop**, **design** and **engineer** both the industrial process and the equipment used.

To do the above purposes, the chemical engineer need to know well about :-

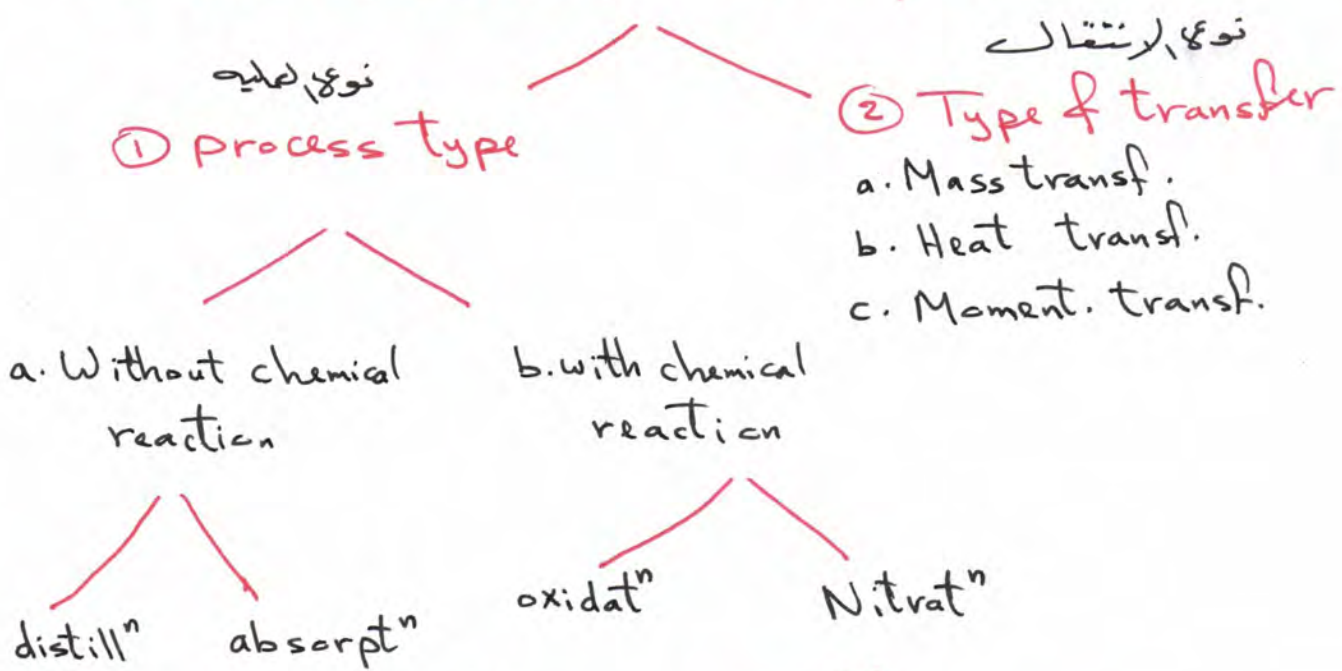
1. Stoichiometry
2. Thermodynamics
3. Kinetic
4. Unit operation
5. Economic

(*) What is Unit-Operation ?

The physical operation for manufacturing chemicals deals mainly with the transfer and change of materials and Energy principles by physical and physical-chemical means, such as :-

Mass, heat and momentum transfer.

- Classification of Unit-operation -



* Mass-Transfer and Diffusion.

When mass is being transferred from one distinct phase to another or through a single phase, the basic mechanisms are the same whether the phase is gas, liquid or solid.

$$\text{Rate of transfer process} = \frac{\text{driving force}}{\text{resistance}}$$

i.e. $\psi = -D \cdot \frac{dx}{dy}$

$$\therefore \psi = N_A = -D_{AB} \cdot \frac{dC_A}{dz}$$

$$\psi = q = -k \cdot \frac{dT}{dz}$$

$$\psi = \tau = -\mu \cdot \frac{du}{dz}$$

Mass Transfer

Fick's Law

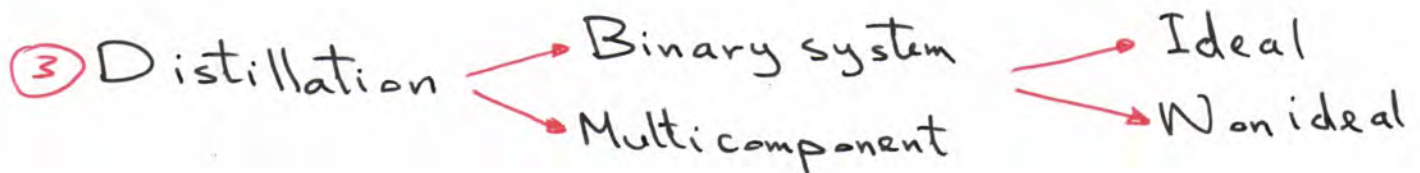
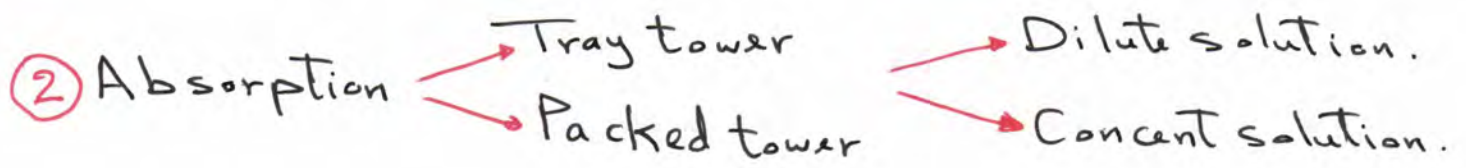
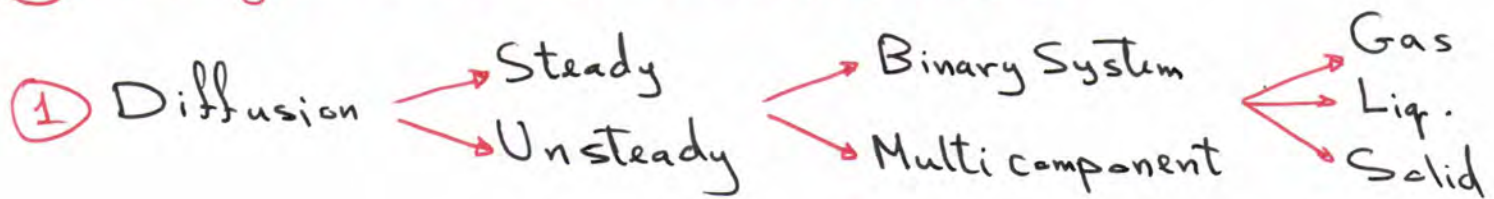
Heat Transfer

Fourier's Law

Moment. Transf.

Newton's Law

* Subjects to be Studied are :-



④ Mechanical separation or Size reduction.

⑤ Boundary Layer.

Diffusion

The transport of molecules from a higher concentration to lower concentration is often called "diffusion".

Diffusion through

- (a) Stagnant media called molecular diffusion
- (b) Turbulent media called eddy or turbulent diffusion.

Molecular diffusion is important when the medium is stagnant or laminar flow flowing. But does not have a significant role to play if the medium is turbulent except in region very close to the phase boundary.

Example - vaporization of a layer of water

على سطح بئر جليده ماء في الهواء ساكن، وفي الهواء متحرك

Molecular - Diffusion

Since most of mass transfer operations are based on diffusional phenomena, the study of molecular diffusion is essential for understanding these phenomena.

Molecular diffusion or molecular transport can be defined as the transfer or movement of individual molecule through a fluid by means of the ~~new~~ random individual movement of the molecules.

Since the molecules travel in random path, it is often called (Random-Walk-process).

The rate of molecular diffusion is connected with :-

1. molecular velocity
2. concentration difference (driving or potential force)

* Concentration, Velocity and Flux

① Concentration :- Generally expressed as :-

$$\rho_i = \text{mass-conc}^n \text{ (kg/m}^3\text{)}.$$

$$\rho = \text{total mass : conc}^n = \text{density (kg/m}^3\text{)} = \sum \rho_i$$

$$w_i = \rho_i / \rho = \text{mass fraction. } \sum w_i = 1$$

$$C_i = \text{molar. conc}^n \text{ (kmol/m}^3\text{)}.$$

$$C = \text{Total molar. conc}^n \text{ (kmol/m}^3\text{)} = \sum C_i$$

$$x_i = C_i / C = \text{mole fraction. } \sum x_i = 1$$

We usually denote to :-

$$\text{gas mole fraction by } (y_i) = \rho_i / \rho$$

$$\text{liquid mole fraction by } (x_i) = C_i / C$$

* Mass average - velocity (u)

$$u = \frac{1}{\rho} \sum \rho_i u_i \quad \text{for n-component mixture}$$

* Molar - average - velocity (U)

$$U = \frac{1}{C} \sum C_i u_i$$

$$\text{or } \begin{cases} v_i = \frac{N_i}{C} & \text{species} \\ v_m = \frac{N_T}{C_T} & \text{mixture} \end{cases}$$

Prove [$u = U$ if the M.wt of all species are equal].

* Species diffusion velocity relative to molar av. velocity

$$v_{iD} = v_i - v_m \Rightarrow v_i = v_{iD} + v_m$$

3 * Flux :-

The net rate at which a species in a solution passes through a unit area in unit time. It is expressed in $\text{kg/m}^2 \cdot \text{s}$ or $\text{kmol/m}^2 \cdot \text{s}$

Flux $\begin{cases} \text{mass flux} \\ \text{molar flux} \end{cases}$

a mass flux

1. relative to a stationary observer $\Rightarrow n_i = \rho_i u_i$
2. relative to an observer moving with mass avg. velocity $\Rightarrow i_i = \rho_i (u_i - u)$
3. relative to an observer moving with molar. avg. velocity $\Rightarrow j_i = \rho_i (u_i - u)$

b Molar flux.

1. $N_i = C_i u_i$
2. $I_i = C_i (u_i - u)$
3. $J_i = C_i (u_i - u)$

Example (1) :- A gas mixture ($N_2 = 5\%$, $H_2 = 15\%$, $NH_3 = 76\%$ and $Ar = 4\%$) flows through a pipe 25.4 mm in diameter at 4.05 bar total pressure.

If the velocities of the respective components are 0.03 m/s, 0.035 m/s, 0.03 m/s and 0.02 m/s.

Calculate :- mass. avg, molar avg and vol. avg. velocities of the mixture?

Take Mwt for :-

$$N_2 = 28, \quad NH_3 = 17, \quad Ar = 40$$

Solution :-

① to find mass. avg. velocity

$$u = \frac{1}{\rho} (\rho_1 u_1 + \rho_2 u_2 + \rho_3 u_3 + \rho_4 u_4)$$

$$\therefore \rho_i = P_i M_i / R.T, \quad \rho_T = P_T M_T / R.T$$

$$\therefore u = \frac{R.T}{P_T M_T} \left(\frac{P_1 M_1}{R.T} \cdot u_1 + \frac{P_2 M_2}{R.T} \cdot u_2 + \frac{P_3 M_3}{R.T} \cdot u_3 + \frac{P_4 M_4}{R.T} \cdot u_4 \right)$$

$$u = \frac{1}{M_T} \left(\frac{P_1}{P_T} M_1 u_1 + \frac{P_2}{P_T} M_2 u_2 + \frac{P_3}{P_T} M_3 u_3 + \frac{P_4}{P_T} M_4 u_4 \right)$$

$$\therefore P_i / P_T = y_i, \text{ therefore}$$

$$u = \frac{1}{M_T} (y_1 M_1 u_1 + y_2 M_2 u_2 + y_3 M_3 u_3 + y_4 M_4 u_4)$$

$$\Rightarrow M_T = y_1 M_1 + y_2 M_2 + y_3 M_3 + y_4 M_4$$

$$= 0.05 \times 28 + 0.15 \times 2 + 0.76 \times 17 + 0.04 \times 40$$

$$\therefore M_T = 16.22$$

$$u = \frac{1}{16.22} [0.05 * 28 * 0.03 + 0.15 * 2 * 0.03 + 0.76 * 17 * 0.03 + 0.04 * 40 * 0.02]$$

$$u = 0.029 \text{ m/sec.}$$

② to find molar avg. velocity (U)

$$U = \frac{1}{C_T} [C_1 U_1 + C_2 U_2 + C_3 U_3 + C_4 U_4]$$

$$y_i = x_i = \frac{C_i}{C_T} \text{ for vap. and gas}$$

$$U = 0.05 * 0.03 + 0.15 * 0.035 + 0.76 * 0.03 + 0.04 * 0.02$$

$$U = 0.0303 \text{ m/sec}$$

③ Vol. avg. velocity = molar avg. velocity
(for gases)

H.W :- Show that U_m can be expressed as:-

$$U_m = x_A \sqrt{V_A} + x_B \sqrt{V_B}$$

Fick's Law

The basic law of diffusion called "Fick's Law".

Flux of A in B $\rightarrow \bar{J}_A \propto \frac{dC_A}{dz} \Rightarrow \bar{J}_A = -D_{AB} \frac{dC_A}{dz}$

where D_{AB} = diffusion coeff. or diffusivity.

Diffusion is a M.T. process occurs when concⁿ difference between two phases exists. Fick's Law express the molar flux (\bar{J}) with respect to an observer moving with the molar avg. velocity. In practice, it is more useful to use expression for (N_A) instead of (\bar{J}_A).

$$\bar{J}_A = -D_{AB} \cdot \frac{dC_A}{dz} = C_A (u_A - u)$$

$$\begin{aligned} -D_{AB} \frac{dC_A}{dz} &= C_A u_A - C_A \cdot u \\ &= N_A - C_A \cdot \frac{1}{C} (C_A u_A + C_B u_B) \\ &= N_A - \frac{C_A}{C} (N_A + N_B) \end{aligned}$$

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$$N_A = \underbrace{\frac{C_A}{C} [N_A + N_B]}_{\text{bulk flow}} - \underbrace{D_{AB} \cdot \frac{dC_A}{dz}}_{\text{molecular diffusion}}$$

For gas mixture: $P = C \cdot R \cdot T \Rightarrow C = \frac{P}{RT} \Rightarrow C_A = \frac{P_A}{RT}$

$$dP = RT dC \Rightarrow dC = \frac{dP}{RT} \Rightarrow dC_A = \frac{dP_A}{RT}$$

$$N_A = \frac{P_A / RT}{P_T / RT} [N_A + N_B] - D_{AB} \frac{dP_A}{RT \cdot dz}$$

$\frac{\partial}{\partial y_A}$
pressi

To convert (N_A) in terms of mole fraction (y_A)

$$y_A = \frac{P_A}{P_T} \Rightarrow P_A = y_A P_T \Rightarrow dP_A = P_T dy_A$$

$$\therefore N_A = [N_A + N_B] \cdot y_A - \frac{D_{AB}}{R \cdot T} \cdot P_T \frac{dy_A}{dz}$$

$$\therefore N_A = [N_A + N_B] y_A - C_T D_{AB} \frac{dy_A}{dz}$$

$\frac{\partial}{\partial y_A}$
mole fraction

Note:- All the above equations for component (A) is derived in same way for species (B).

Then the total flux will be :-

$$N_T = N_A + N_B$$

Diffusion in Gases

The most important cases to be studied are :-

- ① Equimolar - Counter diffusion (EMD).
- ② Diffusion of species (A) through stagnant non-diffusing (B). } No bulk movement
- ③ Diffusion of gas (A) and (B) plus-convection.
- ④ Diffusion through a varying :-
 - Ⓐ Path-length (time dependent).
 - Ⓑ Cross-sectional area.

① Equimolar - Counter - diffusion (EMD)

For species (A, B), both diffuses at equal rates but in opposite direction.

A real life example for this case is :-

Ⓐ (partical of carbon burning in air)



The particle is surrounded by an air-film, through which the molecules of oxygen diffuse and reach the surface of the particle to sustain combustion.

If $[O_2]$ molecule diffuses to the surface, a molecule of CO_2 is formed which diffuses out through the air-film.

So (O_2 and CO_2) undergo (EMD).



Ⓑ vapour rising in the distillation column remain in contact with the down-flowing liquid



$$N_A = -N_B$$

$$\therefore N_A = \frac{P_A}{P_T} [N_A + N_B] - \frac{D_{AB}}{RT} \frac{dP_A}{dz}$$

for stationary bulk flow

$$\therefore N_A = -D_{AB} / RT \frac{dP_A}{dz}$$

$$N_A \cdot dz = \frac{-D_{AB}}{RT} dP_A$$

by integration =

$$N_A = \frac{D_{AB}}{RT} \cdot \frac{P_{A1} - P_{A2}}{z}$$

for EMD

* In this case, total pressure (P_T) must be constant, therefore the net moles of (A) diffusing to the right must equal moles of (B) diffusing to the left.

$$* N_A = -N_B$$

$$\therefore -D_{AB} \frac{dC_A}{dz} = +D_{BA} \frac{dC_B}{dz}$$

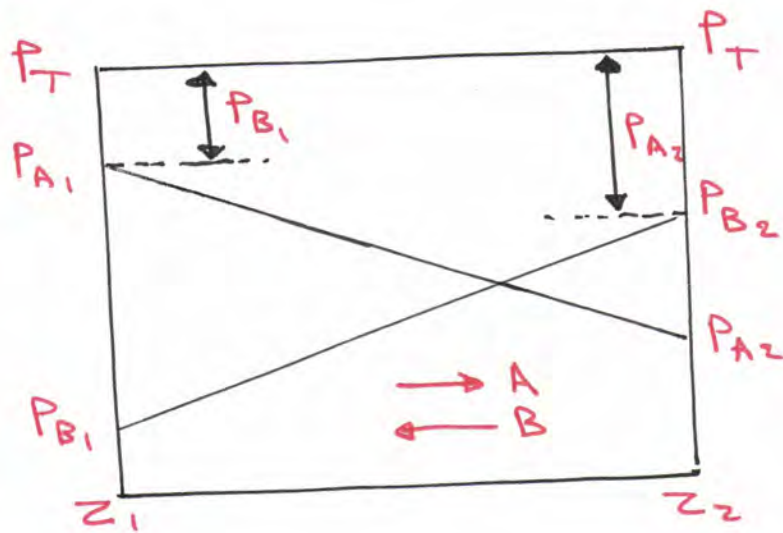
$$C_T = C_A + C_B \Rightarrow dC_A + dC_B = dC_T = 0$$

$$\therefore -dC_A = +dC_B$$

$$\therefore D_{AB} = D_{BA} = D$$

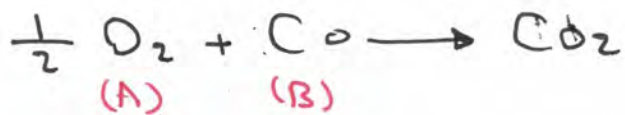
∴ for binary gas mixture the diffusivity of both (A) and (B) are the same.

The distribution of the partial pressure of (A) and (B) along the diffusion path in this case are linear.



(*) For a Non-equimolar-counter diffusion (NEMD)

Producing (CO₂) from (O₂) and hot-char-particle



hence $N_A = \frac{1}{2} N_B$

$$N_A = (N_A + N_B) \cdot \frac{P_A}{P_T} + (-D_{AB}) \frac{dP_A}{dz}$$

Now substitute $N_B = 2N_A$ and then integrate.

Example (1) 2. A uniform tube (0.1) m long and (0.01 m di), containing N_2 gas, Ammonia gas (A) diffusing through the pipe. At point (1), $P_{A1} = 1.013 \times 10^4$ pas, and at point (2) $P_{A2} = 0.507 \times 10^4$ pas. The total pressure is 1.013×10^5 pas and the diffusivity $D_{AB} = 0.23 \times 10^{-4} \text{ cm}^2/\text{sec}$. Calculate :-
 1- Fluxes of (A) and (B), at Temp = 298 K.
 2- The rate of M.T for species (A) and (B).

Solution 2-

1- Closed system with constant pressure and Temp., then the diffusion flow is EMD :-

$$\begin{aligned} \therefore N_A &= \frac{D_{AB}}{RT} \cdot \frac{P_{A1} - P_{A2}}{z_2 - z_1} \\ &= \frac{0.23 \times 10^{-4}}{8.314 \times 298} \cdot \frac{1.013 \times 10^4 - 0.507 \times 10^4}{0.1 - 0.0} \end{aligned}$$

$$N_A = 4.7 \times 10^{-7} \text{ kgmol/m}^2 \cdot \text{Sec}$$

For species (B), $P_{B2} = P_T - P_{A2}$, $P_{B1} = P_T - P_{A1}$

$$\text{or } N_B = -4.7 \times 10^{-7} \text{ kgmol/m}^2 \cdot \text{Sec.}$$

2- mass transfer rate (n_A)

$$\therefore N_A = \frac{n_A}{\text{Area}} \Rightarrow n_A = N_A \times A$$

$$A = \frac{\pi}{4} d_i^2 \Rightarrow \frac{\pi}{4} (0.01)^2 \Rightarrow 7.8 \times 10^{-5} \text{ m}^2 \quad \therefore n_A = 4.7 \times 10^{-7} \times 7.8 \times 10^{-5}$$

$$\therefore n_A = 36.66 \times 10^{-12} \frac{\text{kgmol}}{\text{Sec}}$$

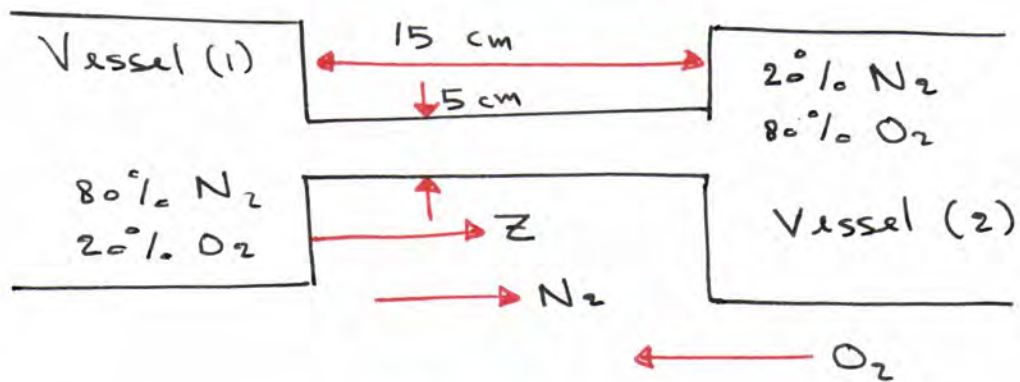
$$\therefore n_B = -36.66 \times 10^{-12} \frac{\text{kgmol}}{\text{Sec}}$$

3- Try to calculate (U_A, U_B)

$$U_A = \frac{N_A}{C_A} \quad \& \quad C_A = \frac{P_A}{RT}$$

Example (2) - Two large vessels are connected as shown in Figure. Vessel (1) contains 80% N_2 (A) and 20% O_2 (B), Vessel (2) contains 20% N_2 and 80% O_2 . The temp. ($20^\circ C$) and the total pressure (2 atm). Calculate :-

- ① The flux and rate of transport of N_2 from vessel 1 to 2, and same for O_2 .
 - ② The partial pressure of N_2 and its gradient in the tube (0.05 m) from vessel (1). (dP_A/dz).
 - ③ The net mass flux (n_T).
- Given that diffusivity of N_2-O_2 pair is $1.01 \times 10^{-5} m^2/s$ at $20^\circ C$ and 2 atm.



Solution:-

$$\textcircled{1} N_A = \frac{+D_{AB}}{RT} \frac{P_{A1} - P_{A2}}{z_2 - z_1}$$

$$D_{AB} = 1.01 \times 10^{-5} (m^2/s), R = 0.0821, T = 293 K$$

$$z = z_2 - z_1 = 0.15 m, \text{ To find } P_{A1}, P_{A2}$$

$$P_{A1} = y_{A1} \cdot P_T$$

$$= 0.8 * 2$$

$$= 1.6 \text{ atm}$$

$$P_{A1} = 1.62 \times 10^5 \frac{N}{m^2}$$

$$P_{A2} = y_{A2} \cdot P_T$$

$$= 0.2 * 2$$

$$= 0.4 \text{ atm}$$

$$P_{A2} = 0.406 \times 10^5 \frac{N}{m^2}$$

$$\therefore N_A = 3.3 \times 10^{-6} \text{ kmol/m}^2 \cdot \text{Sec.}$$

$$\text{rate of transport} = n_A = N_A \cdot \text{Area} \leftarrow \frac{\pi}{4} d_i^2 \leftarrow \frac{\pi}{4} (0.05)^2$$

$$= n_A = 6.6 \times 10^{-9} \text{ kmol/Sec.}$$

$$N_B = -N_A = -3.36 \times 10^{-6} \text{ kmol/m}^2 \cdot \text{s}$$

$$n_A = -n_B = -6.6 \times 10^{-9} \text{ kmol/Sec.}$$

② Partial press. changes linearly with diffusion Path.

$$\therefore \frac{dP}{dz} = \frac{P_{A2} - P_{A1}}{z} = \frac{(0.4 - 1.6)}{0.15} = -8 \text{ atm/m}$$

at point 0.05

$$P_A = P_T + \left[\frac{dP_A}{dz} \right] \cdot \Delta z \quad \text{قانون هينري}$$

$$= 1.6 + (-8) \cdot 0.05$$

$$P_A \frac{dP}{dz} = 1.2 \text{ atm.}$$

$$\textcircled{3} n_T = N_A \cdot M_{wtA} + N_B M_{wtB}$$

$$= (3.36 \times 10^{-6} \times 28) - (3.36 \times 10^{-6} \times 32)$$

$$n_T = -1.344 \times 10^{-5} \text{ kmol/m}^2 \cdot \text{s}$$

② Diffusion of (A) through non-diffusing (B)

In this case one boundary at the end of the diffusion path is impermeable to other species, so it can not pass through.

A real life example: Dry air in sulphuric acid plant.

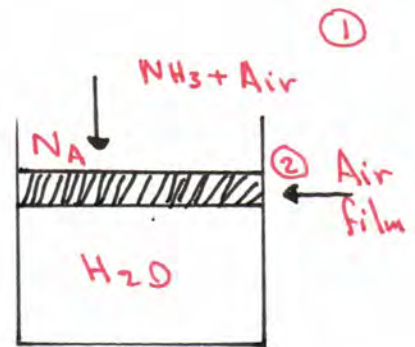
تحتاج الهواء الجاف لمرحلة هزجة الكبريت في مصنع إنتاج حامض H_2SO_4 لذلك يمر الهواء الرطب على حامض كبريتيك مركز في زجاج محسوس، الرطوبة تنفذ خلال طبقة من الهواء وتصل إلى سطح الحامض وبعد ما تمتصت لكن الهواء لا يزيد في الحامض وبالتالي لا يستطيع التنازل.

∴ moisture (H_2O) has $\begin{cases} \text{Source (bulk of air)} \\ \text{Sink (the acid)} \end{cases}$

but (dry air) has (source) but no (sink).

- Another example (NH_3 in Air) removal.

NH_3 diffuses easily across air film to water, but (Air) is insoluble.



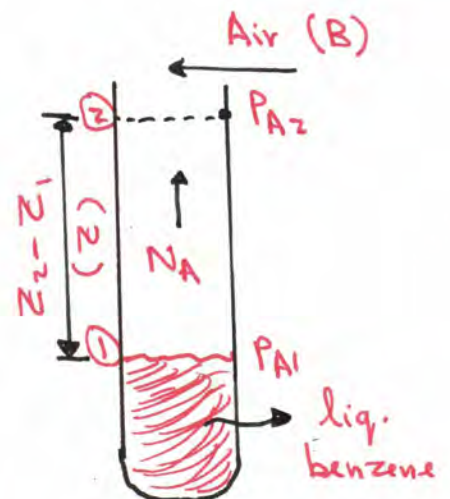
$$\therefore N_A = (N_A + N_B) \frac{P_A}{P_T} - \frac{D_{AB}}{RT} \frac{dP_A}{dz}$$

$$\therefore N_B = 0$$

$$\therefore N_A = N_A \frac{P_A}{P_T} - \frac{D_{AB}}{RT} \frac{dP_A}{dz}$$

$$\therefore N_A * dz = - \frac{D_{AB} P_T}{RT (P_T - P_A)} dP_A$$

integrating over $z = 0, P_A = P_{A1}$
and $z = L, P_A = P_{A2}$



$$N_A = \frac{D_{AB} P_T}{RTZ} \ln \frac{P_T - P_{A2}}{P_T - P_{A1}}$$

$\frac{P_{B2}}{\dots}$ above $P_T - P_{A2}$
 $\frac{P_{B1}}{\dots}$ below $P_T - P_{A1}$

It is more convenient to use [Log mean partial pressure]

$$N_A = \frac{D_{AB} P_T}{RTZ} \cdot \frac{P_{A1} - P_{A2}}{P_{BM}}$$

For $N_B = 0$

where $P_{BM} = \frac{P_{B2} - P_{B1}}{\ln \frac{P_{B2}}{P_{B1}}}$

This equation expresses the flux in terms of driving force of species (A), $(P_{A1} - P_{A2})$.

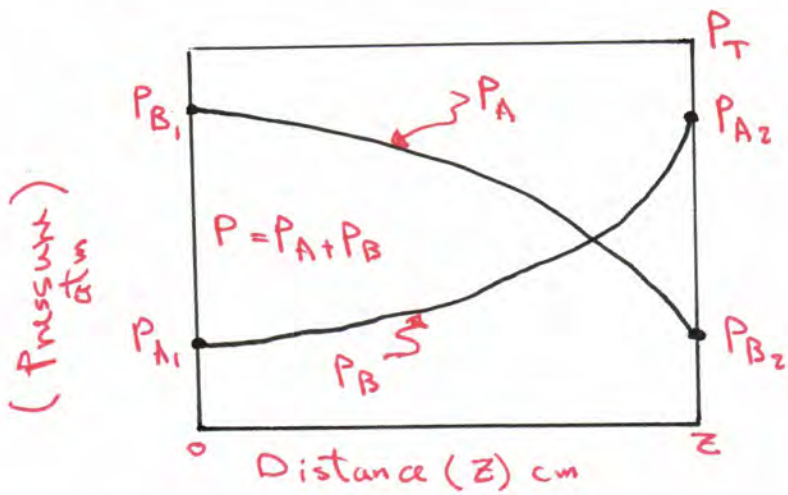
We also can calculate the partial pressure (P_A) of (A) at any intermediate point and the distribution of the partial pressure of (A) along the diffusion path, by using :-

$$N_A = \frac{D_{AB} P_T}{RTZ} \ln \frac{P_T - P_A}{P_T - P_{A1}}$$

H.W 3 Try To Find

$$N_A = \frac{C_T D_{AB}}{Z_2 - Z_1} \cdot \frac{X_{A1} - X_{A2}}{X_{BM}}$$

Typical distribution of the partial pressure of the components along the diffusion path



هنا عندنا (A) ينفذ من $z=0$ الى $z=L$
 كذلك (B) ينفذ من $z=L$ الى $z=0$
 اية له $P_{\text{mass. gradient}}$
 ولكن عندنا $N_B = 0$
 اية $\text{flux} = 0$ مثال =

سوف تسبب يسرع باتجاه معاكس لجريان الماء. فالذي نراه اننا لسوف
 لا تتحرك تقريباً في الماء، ولكن في الحقيقة اننا لمحركه، ولكن
 حركة الماء المعاكس تسمى سرعة الماء وبالتالي نحسب
 $\text{flux لها} = \text{معر}$.