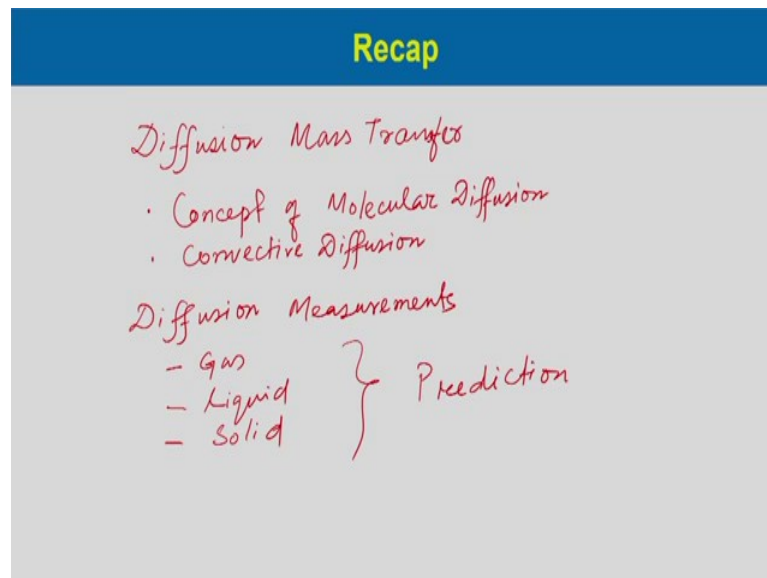


Mass Transfer Coefficients
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Lecture – 09
Mass transfer coefficient concept and classifications

Welcome to the second module of the course mass transfer operation. In this module we will discuss Mass Transfer Coefficients, before going to the today's first lecture let us have small recap on our previous lecture. In our previous know module particularly we have discussed diffusion mass transfer.

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Under which we have a discussed concept of molecular diffusion and convective diffusion. We have also considered diffusion measurements, gas then liquid and solid, we have also discussed its prediction.

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Module 2: Lecture 1

Concept of Mass Transfer Coefficients

- What is mass transfer coefficient?
- Types of mass transfer coefficient
- Diffusion of A through non diffusing B
- Equimolar counterdiffusion of A and B
- Relation between the gas-phase mass transfer coefficients, K_G and K_Y

So, in this lecture, we will consider concept of mass transfer coefficient. In the first lecture we will mostly concentrate on the what is mass transfer coefficient, type of mass transfer coefficient, diffusion of A through non diffusing B and in that case what should be the mass transfer coefficient. Then we will discuss equimolar counter diffusion of A and B and we find out the different mass transfer coefficient related to the system and finally, we will discuss the relation between the gas phase mass transfer coefficients K_G and K_Y .

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Concept of Mass Transfer Coefficients

- **The mass transfer coefficient is defined as follows:**
 - Rate of mass transfer \propto concentration driving force (i.e. the difference in concentration)
 - Rate of mass transfer \propto Area of contact between the phases
- If, W_A = Rate of mass transfer (kmol/s) of the solute A,
 ΔC_A = Concentration driving force between two points,
 a = area of mass transfer,
 $\rightarrow W_A \propto a \Delta C_A$
 $\Rightarrow W_A = k_c a \Delta C_A$ where k_c the proportionality constant, called the mass transfer coefficient.

The mass transfer coefficient is defined as the rate of mass transfer as we know is proportional to the concentration driving force that is the difference in concentration and also the rate of mass transfer is proportional to the area of contact between the phases.

So, if we consider W_A is the rate of mass transfer in terms of kilomole per second of particular solute A and if the concentration driving force between the two points is ΔC_A , that is C_{A1} and C_{A2} , $C_{A1} - C_{A2}$ ΔC_A is the concentration driving force and the area of mass transfer is a in that case we can write W_A is proportional to $a \Delta C_A$ concentration driving force. So, from this we can write W_A would be equal to $k_c a \Delta C_A$. This k_c over here is basically the proportionality constant and it is called the mass transfer coefficient.

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Concept of Mass Transfer Coefficients

- If N_A is the molar flux (expressed as $\text{kmol/m}^2\cdot\text{s}$, say), we may write

$$W_A = a N_A = k_c a \Delta C_A$$
- ⇒ **Mass transfer coefficient**, $k_c = \frac{N_A}{\Delta C_A} = \frac{\text{molar flux}}{\text{concentration driving force}}$.
- For the purpose of comparison, we may recall the definition of heat transfer coefficient:

$$\Rightarrow \text{Heat transfer co-efficient, } h = \frac{\text{heat flux}}{\text{temperature driving force, } \Delta T}$$

So, this is the mass transfer coefficient. Now, if we considered N_A is the molar flux which is expressed in kilomole per metre square second we may write W_A would be equal to $a N_A$ because W_A is the molar flow rate that is kilomole per second. So, if we multiplied by the area a then the unit would be molar flow rate that is k mole per second. So, then if you substitute N_A , it would be $k_c a \Delta C_A$ and from this we can write mass transfer coefficient. So, mass transfer coefficient k_c would be equal to from this relation we can write k_c would be equal to N_A by ΔC_A ; that means, the molar flux divided by the concentration driving force.

For the purpose of comparison, we may recall the definition of heat transfer coefficient. We know for heat transfer coefficient h , which is the heat flux divided by the temperature driving force. So, it is in the similar fashion we can write the mass transfer coefficient is the molar flux divided by the concentration driving force.

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Concept of Mass Transfer Coefficients

- The inverse of mass transfer coefficient is a measure of the **mass transfer resistance**.
- If the driving force is expressed as the difference in concentration (kmol/m^3 , say):
 - the unit of mass transfer coefficient is **m/s**

(or cm/s, ft/s, etc. which is same as the unit of velocity).

The inverse of mass transfer coefficient is a measure of the mass transfer resistance. So, 1 by mass transfer coefficient would be the resistance for mass transfer. If the driving force is expressed as the difference in concentration that is kilomole per metre cube say, then we can write the unit of mass transfer coefficient would be meter per second. So, this is a unit of velocity. So, we can also write in terms of centimetre per second or in feet per second with no different length in it which is the same of the unit of the velocity.

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Concept of Mass Transfer Coefficients

- If the mass transfer coefficient is expressed as the ratio of the local flux and the local driving force, it is called the **local mass transfer coefficient**.

$$\text{Local mass transfer coefficient} = \frac{\text{local flux}}{\text{local driving force}}$$

- When it is expressed as the ratio of the average flux (over a surface) and the average driving force, it is known as the **average mass transfer coefficient**.

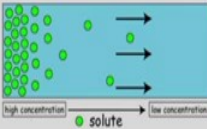
$$\text{Average mass transfer coefficient} = \frac{\text{average flux}}{\text{average driving force}}$$

Now, if the mass transfer coefficient is expressed as the ratio of the local flux and the local driving force, then it is called the local mass transfer coefficient. So, the local mass transfer coefficient can be written as local flux divided by the local driving force. Now, when it is expressed as the ratio of the average flux that is over a surface if the flux changes and if we take the average then the average flux divided by the average driving force then it should be named as average mass transfer coefficient. So, average mass transfer coefficient would be equal to average flux divided by the average driving force.

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Types of Mass Transfer Coefficients

- Convective mass transfer can occur in a gas or a liquid medium.
- A few choices of the driving force
 - Difference in concentration
 - Difference in partial pressure
 - Difference in molar fraction
- In the case of heat transfer, the temperature difference is the only driving force.



Convective mass transfer can occur in a gas or in a liquid medium, but it does not occur in the solid medium. So, in these two cases a few choices of the driving force can be written, one is the difference in concentration. So, in case of concentration driving force it diffuses from the high concentration to the low concentration. So, the solute moves from higher concentration to the lower concentration, we can also write the driving force in terms of the partial pressure difference of a particular component. So, partial pressure is another driving force. We can also write in terms of the molar fraction.

So, mole fraction can be written as the driving force. But in case of the heat transfer the temperature difference is the only driving force.

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Types of Mass Transfer Coefficients

- Different types of mass transfer coefficients have been defined depending upon:
 - (i) Whether mass transfer occurs in the gas phase or in the liquid phase
 - (ii) The choice of the driving force
 - (iii) Whether it is a case of diffusion of A through non-diffusing B or a case of counterdiffusion.
- If the transport of mass occurs through a stagnant film of thickness δ , we may write down

Flux = mass transfer coefficient \times driving force

So, different type of mass transfer coefficient have been defined depending upon the driving force and also it depends on whether the mass transfer occur in the gas phase or in the liquid phase. And second is choice of driving force, as we said before it is also depends on the driving force and the third one is whether it is a case of diffusing of component A through non diffusing B or whether it is a equimolar counter current diffusion or the counter diffusion. If the transport of mass occur through a stagnant film of thickness delta, then we can write down the flux as mass transfer coefficient into driving force.

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Diffusion of A through Non-diffusing B

- Mass transfer in the gas phase :**
$$N_A = k_G(p_{A1} - p_{A2}) = k_y(y_{A1} - y_{A2}) = k_c(C_{A1} - C_{A2})$$

where $k_G, k_y,$ and k_c = the gas phase mass transfer coefficients
The unit of mass transfer coefficient (k_y) is $\text{kmol/m}^2\text{s}(\Delta y)$
where Δy stands for driving force in mol fraction unit.
- Mass transfer in the liquid phase:**
$$N_A = k_x(x_{A1} - x_{A2}) = k_L(C_{A1} - C_{A2})$$

where k_x and k_L = liquid phase mass transfer coefficients, the subscript 1 and 2 refer to two positions in a medium

So, the mass transfer in the gas phase in that case we can write N_A would be equal to $k_G p_{A1} - p_{A2}$ which would be equal to $k_y (y_{A1} - y_{A2})$ is equal to $k_c (C_{A1} - C_{A2})$. So, this is termed as equation 1. In this case the k_G, k_y and k_c these are the gas phase mass transfer coefficient; k_G that is when we define the driving force in terms of the partial pressure difference, when we define the driving force as a mole fraction unit then it is k_y and when we define the concentration unit we define the mass transfer coefficient as k_c .

So, three different mass transfer coefficient is defined over here in case of gas phase depending on the driving force we use. The unit of mass transfer coefficient k_y we can easily calculate from these flux equation and which is kilomole per metre square second divided by the driving force. Here Δy stands for the driving force in mole fraction unit. In case of the liquid we can write N_A is the flux is equal to $k_x (x_{A1} - x_{A2})$ is equal to $k_L (C_{A1} - C_{A2})$. So, this is equation 2 for the mass transfer in the liquid phase.

So, the mass transfer coefficient k_x and k_L these two are the mass transfer coefficient in the liquid phase and the subscript for both the equations, subscript 1 and 2 refer to the two position in the medium it is considered.

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Diffusion of A through Non-diffusing B

- **If the gas phase is ideal:**

$$N_A = k_c(p_{A1} - p_{A2}) = k_y(y_{A1} - y_{A2}) = k_c(C_{A1} - C_{A2})$$

The concentration term in eq. (1) is given by

$$C_A = p_A/RT$$

Where p_A = The partial pressure of A.
- **Supposing that the distance between the two locations 1 and 2 is δ (the film thickness), the expression of mass transfer coefficient can be obtained by comparing eq.(1) with**

$$N_A = \frac{D_{AB} p_t}{RT(x_2 - x_1)p_{BLM}} (p_{A1} - p_{A2}) = \frac{D_{AB} p_t}{RT l p_{BLM}} (p_{A1} - p_{A2})$$

Where p_{BLM} = logarithmic mean partial pressure of species B

$$p_{BLM} = \frac{p_{B2} - p_{B1}}{\ln \left(\frac{p_{B2}}{p_{B1}} \right)}$$

Now, if we consider the gas phase is ideal in that case, the concentration term in equation one which is this the concentration term C_A we can write as p_A by $R T$, the partial pressure of component a divided by $R T$. So, in that case p_A is the partial pressure of component A. Now, suppose that the distance between the two locations A and 1 and 2 is delta that is the film thickness.

The expression of mass transfer coefficient can be obtained by comparing equation 1 with this equation, this we have already defined in case of diffusion N_A would be equal to $D_{AB} p_t$, p_t is the total pressure by $R T \times 2$ minus x_1 P_{BLM} into p_{A1} minus p_{A2} would be equal to $D_{AB} P t$ divided by $R T l p_{BLM}$, here l is the thickness over here x_2 minus x_1 . Later we will define as delta into p_{BLM} into partial pressure difference where, the p_{BLM} is the logarithmic mean partial pressure of species B. We can define it p_{B2} minus p_{B1} divided by $\ln p_{B2}$ by p_{B1} .

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Diffusion of A through Non-diffusing B

- The expression of mass transfer coefficient can be obtained by comparing eq.(2) with

$$N_A = \frac{D_{AB} (\rho/M)_{avg}}{\ell x_{BLM}} (x_{A1} - x_{A2})$$

$N_A = k_x(x_{A1} - x_{A2}) = k_L(C_{A1} - C_{A2})$
2

Where x_{BLM} = logarithmic mean *mole fraction* partial pressure of species B

$$x_{BLM} = \frac{x_{B2} - x_{B1}}{\ln \left(\frac{x_{B2}}{x_{B1}} \right)}$$

So, the expression of mass transfer coefficient can be obtained by comparing equation 2 with this equation; equation 2 in the liquid phase with the other equation in case of the diffusion that is N_A would be equal to $D_{AB} \rho$ by M average divided by $l \times x_{BLM}$ into $x_{A1} - x_{A2}$. In this case x_{BLM} it would be logarithmic mean of mole fraction of component B not partial pressure mole fraction of species B which is $x_{B2} - x_{B1}$ divided by $\ln x_{B2}$ by x_{B1} .

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Diffusion of A through Non-diffusing B

- Gas phase:** $k_G = \frac{D_{AB} P_t}{RT \delta p_{BLM}}$, $k_y = \frac{D_{AB} P_t^2}{RT \delta p_{BLM}}$, $k_c = \frac{D_{AB} P_t}{\delta p_{BLM}}$ 3
- The relation among the three types of gas phase mass transfer coefficients (i.e. k_G , k_y , and k_c) can be easily obtain from eq. 3.
- Liquid Phase:** $k_x = \frac{D_{AB} (\rho/M)_{avg}}{\delta x_{BLM}}$, $k_L = \frac{D_{AB}}{\delta x_{BLM}}$ 4

Here, $l = \delta$

- the relation between the two types of liquid – phase mass transfer coefficients, k_x and k_L can be obtained from eq. 4.

$$k_c = RT k_G; \quad k_y = P_t k_G \quad k_x = (\rho/M)_{avg} k_L$$

In case of gas phase we can define k_G is equal to $\frac{D_{AB} P}{\ell}$ by $\frac{R T \Delta p}{\Delta p}$ BLM. Similarly, for k_y in terms of the mole fraction unit of the component gas phase mole fraction unit that is equal to $\frac{D_{AB} P}{\ell}$ square divided by $\frac{R T \Delta p}{\Delta p}$ BLM and k_c would be equal to $\frac{D_{AB} P}{\ell}$ by Δp BLM. So, this is equation 3. The relation among the three types of gas phase mass transfer coefficient that is k_G , k_y and k_c among these 3 can easily be obtained from these 3 relations. In case of liquid phase, we can write k_x would be equal to $\frac{D_{AB} \rho}{M_{avg}}$ divided by Δx BLM and k_L would be $\frac{D_{AB}}{\ell}$ by Δx BLM.

So, this is equation 4 and here we have considered the ℓ the thickness between the two points where the diffusion is occurring as Δx the diffusion length. The relation between the two types of liquid phase mass transfer coefficient similarly to the gas phase we can obtained from k_x and k_L relations in equation 4 among these two. So, k_c would be equal to $\frac{R T}{P} k_G$, k_y would be equal to $\frac{P}{R T} k_G$ and k_x would be $\frac{\rho}{M_{avg}} k_L$. So, this is the final relations we can obtained among these mass transfer coefficient.

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Equimolar Counterdiffusion of A and B

- The set of notations for mass transfer coefficients are used here with a **prime (')**
- Gas phase: $N_A = k'_G (p_{A1} - p_{A2}) = k'_y (y_{A1} - y_{A2}) = k'_c (C_{A1} - C_{A2})$
6
- Liquid Phase: $N_A = k'_x (x_{A1} - x_{A2}) = k'_L (C_{A1} - C_{A2})$
7
- Comparing eq. (6) for gas-phase transport with

$$N_A = \frac{D_{AB} P}{R T \ell} (y_{A1} - y_{A2}) = \frac{D_{AB} P}{R T \ell} (p_{A1} - p_{A2})$$
- Comparing eq. (7) for liquid-phase transport with

$$N_A = \frac{D_{AB} (\rho / M)_{avg}}{\ell} (x_{A1} - x_{A2})$$

Now, the set of notations for mass transfer coefficient used in case of equimolar counter diffusion of a and b. So, if we considered equimolar counter diffusion here we have defined the mass transfer coefficient with a sign of prime. So, like in gas phase, we can write N_A would be equal to k_G dash into the partial pressure driving force p_{A1} minus p_{A2} which is equal to k_y dash into the mole fractions driving force that is y_{A1} minus

y_{A2} is equal to k_c dash into the concentration driving force C_{A1} minus C_{A2} this is equation 6. In case of liquid phase, similarly we can write N_A would be equal to k_x dash x_{A1} minus x_{A2} is equal to k_L dash C_{A1} minus C_{A2} .

Now if we consider equation 6, for the gas phase transport with this relations N_A equal to $D_{AB} P_t$ by RT $1 y_{A1}$ minus y_{A2} is equal to D_{AB} by RT $1 p_{A1}$ minus p_{A2} , then we can also compare equation 7 for the liquid phase transport with this relation N_A would be equal to $D_{AB} \rho$ by $M_{average}$ divided by $l x_{A1}$ minus x_{A2} .

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Equimolar Counterdiffusion of A and B

- We can have the following expressions for the mass transfer coefficients in this case:

Gas phase: $k'_G = \frac{D_{AB}}{RT\delta}$ $k'_y = \frac{D_{AB} P_t}{RT}$ $k'_c = \frac{D_{AB}}{\delta}$ 8

Liquid Phase: $k'_x = \frac{D_{AB} (\rho/M)_{av}}{\delta}$ $k'_L = \frac{D_{AB}}{\delta}$ 9

Conversion: $k'_c = \frac{RT}{P_t} k'_y = RT k'_G$;

$k'_x = (\rho/M)_{avg} k'_L = C_{avg} k'_L$

Then we can have the following expression for the mass transfer coefficient in this case. Gas phase we will get k_G dash would be equal to D_{AB} by RT into delta, k_y dash would be equal to $D_{AB} P_t$ by RT and k_c dash would be D_{AB} by delta.

Similarly, this is equation 8. Similarly, we can get for liquid phase k_x dash would be equal to $D_{AB} \rho$ by $M_{average}$ by delta and k_L dash would be D_{AB} by delta. So, this is equation 9. Now, if we convert among them then we will get k_c dash would be equal to RT by P_t k_y dash which would be equal to $RT k_G$ dash and k_x dash would be equal to ρ by $M_{average}$ k_L dash would be equal to $C_{average}$ k_L dash.

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Equimolar Counterdiffusion of A and B

- If the concentration of A is expressed in the mole ratio unit, the mass transfer coefficients k_Y and k_X are expressed as:

Conversion: $N_A = k_Y(Y_{A1} - Y_{A2})$ for the gas phase

and $N_A = k_X(X_{A1} - X_{A2})$ for the liquid phase
- where Y_A and X_A are the concentrations of A in the gas or in the liquid phase in mole ratio unit [note that similar expressions can be written using the mass ratio (mass A/ mass B) unit as well].

Note that $Y_A = \frac{y_A}{1 - y_A}$ ✓ and $X_A = \frac{x_A}{1 - x_A}$

Now, if the concentration of A is expressed in mole ratio unit, the mass transfer coefficient k_Y and k_X are expressed as N_A would be equal to $k_Y(Y_{A1} - Y_{A2})$ for the gas phase.

That means here k_Y and k_X they are not mole fraction unit they are mole ratio unit and similarly for the liquid phase N_A would be $k_X(X_{A1} - X_{A2})$ here, X_{A1} and X_{A2} are the mole ratio unit in the liquid phase. So, Y_A and X_A are the concentration of A in the gas or in the liquid phase in mole ratio unit, note that similar expression can be written using the mass ratio unit that is mass of A by mass of B unit as well. So, here in this case, as we said it is mole ratio unit; that means, it is a Y_A would be equal to small y_A divided by $1 - y_A$.

That is the solute free basis, this we need to consider when we consider different know systems or applications of mass transfer coefficient for different mass transfer operations. Similarly for liquid phase X_A would be equal to small x_A divided by $1 - x_A$.

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Different types of mass transfer coefficients				
Diffusion of A through non diffusing B		Equimolar counterdiffusion of A and B		Unit of the mass transfer coefficient
Flux, N_A	Mass transfer coefficient	Flux, N_A	Mass transfer coefficient	
Gas-phase mass transfer				
$k_G(p_{A1} - p_{A2})$	$k_G = \frac{D_{AB} P}{RT \delta p_{BM}}$	$k_G'(p_{A1} - p_{A2})$	$k_G' = \frac{D_{AB}}{RT \delta}$	$\frac{\text{mol}}{(\text{time})(\text{area})(\Delta p_A)}$
$k_y(y_{A1} - y_{A2})$	$k_y = \frac{D_{AB} P^2}{RT \delta p_{BM}}$	$k_y'(y_{A1} - y_{A2})$	$k_y' = \frac{D_{AB}}{RT \delta}$	$\frac{\text{mol}}{(\text{time})(\text{area})(\Delta y_A)}$
$k_c(C_{A1} - C_{A2})$	$k_c = \frac{D_{AB} P}{\delta p_{BM}}$	$k_c'(C_{A1} - C_{A2})$	$k_c' = \frac{D_{AB}}{\delta}$	$\frac{\text{mol}}{(\text{time})(\text{area})(\Delta C_A)}$

Now, different types of mass transfer coefficient as we have discussed for both diffusion of A through non diffusing B and equimolar counter diffusion of A and B, we will summarize them in a tabular format for gas phase mass transfer the flux we write k_G into the partial pressure unit say $p_{A1} - p_{A2}$ and its mass transfer coefficient k_G is $D_{AB} P$ or $P t$ divided by $R T \delta p_{BM}$.

And in case of equimolar counter diffusion of A and B, the flux is k_G' dash $p_{A1} - p_{A2}$ and the mass transfer coefficient is k_G' dash is equal to D_{AB} by $R T$ into delta. The unit of mass transfer coefficient over here is mole per time in per area per partial pressure gradient. Similarly, if we write in terms of the mole fraction unit k_y the flux in case of diffusion of A through non diffusing B, we can write the flux k_y is N_A would be equal to $k_y y_{A1} - y_{A2}$ and its know mass transfer coefficient in this case is $D_{AB} P t$ square divided by $R T \delta p_{BM}$.

And in case of equimolar counter diffusion, flux N_A would be k_y dash into $y_{A1} - y_{A2}$ and then, in this case the mass transfer coefficient k_y dash $D_{AB} P t$ by $R T \delta$. In this case, the unit of mass transfer coefficient would be mole per unit time per unit area per the mole fraction difference. Similarly, in terms of the concentration we can write $k_c C_{A1} - C_{A2}$, k_c would be equal to $D_{AB} P t$ by δp_{BM} and in case of equimolar counter diffusion, it is k_c dash $C_{A1} - C_{A2}$ and k_c dash is equal to D_{AB} by delta.

So, here instead of all other remain same it will be the concentration driving force in case of the unit of mass transfer coefficient remaining all other same.

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Different types of mass transfer coefficients				
Diffusion of A through non diffusing B		Equimolar counterdiffusion of A and B		Unit of the mass transfer coefficient
Flux, N_A	Mass transfer coefficient	Flux, N_A	Mass transfer coefficient	
Liquid –phase mass transfer				
$k_L(C_{A1} - C_{A2})$ ✓	$k_L = \frac{D_{AB}}{\delta x_{BM}}$ ✓	$k_L'(C_{A1} - C_{A2})$	$k_L' = \frac{D_{AB}}{\delta}$	$\frac{\text{mol}}{(\text{time})(\text{area})(\Delta C_A)}$
$k_x(x_{A1} - x_{A2})$ ✓	$k_x = \frac{D_{AB}C}{\delta x_{BM}}$ ✓	$k_x'(x_{A1} - x_{A2})$	$k_x' = \frac{CD_{AB}}{\delta}$	$\frac{\text{mol}}{(\text{time})(\text{area})(\Delta X_A)}$
Conversion				
$k_G RT = \frac{RT}{P_c} k_y = k_c; k_L = \frac{k_x}{C_{av}}$		$k_c' = RT k_G' = \frac{RT}{P_c} k_y'; k_L' = \frac{k_x'}{C_{av}}$ ✓		

Now, if we consider the liquid phase mass transfer, similarly for liquid phase also we have considered k_L and k_x unit which is given over here k_L into the concentration driving force then k_x into the mole fraction driving force and then their mass transfer coefficient is written in the respective terms. And here also it is the unit of mass transfer coefficient will depend on the driving force we are considering for the mass transfer.

Now, if we convert among these all these mass transfer coefficient we can write $k_G RT$ would be equal to $\frac{RT}{P_c} k_y$ would be equal to k_c and k_L would be equal to k_x by C_{av} . In case of the liquid phase, we can write k_c' would be equal to $RT k_G'$ k_y' would be equal to $\frac{RT}{P_c} k_y'$ and k_L' would be equal to k_x' by C_{av} . So, this is the conversion among the mass transfer coefficient both in the gas phase as well as in the liquid phase.

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Different types of mass transfer coefficients

- The former class of coefficients ($k_G, k_y, k_c, k_x,$ and k_L) are inherently associated with the log mean concentration of the other species (B) which is non-diffusing.
- Accordingly this type of mass transfer coefficient has a dependence on concentration because of the term p_{BLM} or x_{BLM} (this dependence can however be ignored at low concentrations of A).
- On the contrary, the coefficients ($k_G', k_y', k_c', k_x',$ and k_L') do not have dependence on concentration.
- The second type of coefficient, k_c' is called 'Colburn-Drew mass transfer coefficient'

The former mass transfer coefficient; that means, in case of the known diffusion of component A through non-diffusing B that is k_G, k_y, k_c, k_x and k_L these are inherently associated with the log mean concentration of the other species B which is non-diffusing. So, if we consider the species B is non-diffusing then, these mass transfer coefficients would be associated with the log mean concentration gradient. Accordingly these type of mass transfer coefficient has a dependence on concentration because of the term p_{BLM} or x_{BLM} .

This dependence can however, be ignored at low concentration. So, if we considered very low concentration this dependency of the logarithmic term or the p_{BLM} or x_{BLM} term in case of gas and the liquid phase we can ignore when the concentration of diffusing species that is component A is very low. On the other hand, in case of the equimolar counter diffusion of component A and B in that case the coefficient k_G', k_y', k_c', k_x' and k_L' do not have dependence on concentration. The second type of mass transfer coefficient or like k_c' is called Colburn Drew mass transfer coefficient.

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F-type coefficient

- Another type of mass transfer coefficient called the F-type coefficient has been proposed (Treybal 1980, Benitez 2002).
- This coefficient, similar to the Colburn-Drew mass transfer coefficient, is not concentration dependent even in the case of 'diffusion of A through non diffusing B'.
- If we integrate over a film of thickness δ , we get

$$N_A = \frac{N_A}{N_A + N_B} \frac{D_{AB} P_t}{RT} \ln \frac{\left[\frac{N_A}{(N_A + N_B)} \right] - y_{A2}}{\left[\frac{N_A}{(N_A + N_B)} \right] - y_{A1}}$$

$$= \frac{N_A}{N_A + N_B} F \ln \frac{\left[\frac{N_A}{(N_A + N_B)} \right] - y_{A2}}{\left[\frac{N_A}{(N_A + N_B)} \right] - y_{A1}}$$

Where
 $y_A = p_A/P_t$
 and
 $\frac{D_{AB} P_t}{RT} = F$

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Now there is another type of mass transfer coefficient which is called F-type coefficient and proposed by Treybal and Benitez in 1980 and then 2002, this is F-type mass transfer coefficient. This coefficient similar to the Colburn Drew mass transfer coefficient and is not concentration dependent even in case of the diffusion of A through non diffusing B ok. So, if it is on for both the cases that is equimolar counter diffusion of component A and B and also the case where diffusion of A through non diffusing B both cases it is independent.

This F-type mass transfer coefficient is independent on the concentration terms. Now if we integrate over a film of thickness delta from this F-type mass transfer coefficient we can obtain N_A would be equal to N_A by N_A plus N_B $D_{AB} P_t$ by $R T$ \ln N_A divided by N_A plus N_B minus y_{A2} divided by N_A by N_A plus N_B minus y_{A1} . So, we can write N_A by N_A plus N_B into F which is this part is written as F into \ln N_A by N_A plus N_B minus y_{A2} divided by N_A by N_A plus N_B minus y_{A1} . So, this is equation 10; y_A is the mole fractions it is p_A by P_t . So, partial pressure of component a divided by the total pressure and $D_{AB} P_t$ by $R T$ is F .

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F-type coefficient

- The F type mass transfer coefficient which is independent of the concentration of the diffusing species.
- If we compare eq. (10) with eq. (3) for the case of diffusion of A through non diffusing B, it is easy to find out that

$$F = k_G p_{BLM}$$

$$k_G = \frac{D_{AB} P_t}{RT \delta p_{BLM}}, k_y = \frac{D_{AB} P_t^2}{RT \delta p_{BLM}}, k_c = \frac{D_{AB} P_t}{\delta p_{BLM}}$$

$$N_A = \frac{N_A}{N_A + N_B} \frac{D_{AB} P_t}{RT} \ln \left[\frac{N_A / (N_A + N_B) - y_{A2}}{N_A / (N_A + N_B) - y_{A1}} \right]$$

$$= \frac{N_A}{N_A + N_B} F \ln \left[\frac{N_A / (N_A + N_B) - y_{A2}}{N_A / (N_A + N_B) - y_{A1}} \right]$$

The F type mass transfer coefficient which is independent of the concentration of the diffusing species and if we compare equation 10 with equation 3 for the case of diffusion of A through non diffusing B, it is very easy to find out F is equal to k G p BLM and this is the equation 3 and 10. So, we can from there if we compare we can write F would be equal to k G p BLM.

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F-type coefficient

- For equimolar counterdiffusion:

F and k' G are related as $F = k'_G P_t$

which is independent of partial pressure of the diffusing species.

Now, for equimolar counter diffusion, F and k G are k G dash are related as F is equal to k G dash P t, which is independent of partial pressure of diffusing species.

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Relation between k_G and k_Y

Solution:

The driving force in the mole ratio unit, Y between two points 1 and 2 can be written as

$$(Y_{A1} - Y_{A2}) = \frac{p_{A1}}{P_t - p_{A1}} - \frac{p_{A2}}{P_t - p_{A2}}$$

$$= \frac{P_t p_{A1} - p_{A1} p_{A2} - P_t p_{A1} + p_{A1} p_{A2}}{(P_t - p_{A1})(P_t - p_{A2})} = \frac{P_t(p_{A1} - p_{A2})}{p_{B1} p_{B2}}$$

$$\rightarrow p_{A1} - p_{A2} = (Y_{A1} - Y_{A2}) \frac{p_{B1} p_{B2}}{P_t}$$

Now, what is the relation between the k_G and k_Y . So, now, we will find out the driving force in case of mole ratio unit Y that is k_Y between the two points 1 and 2 we can write as $Y_{A1} - Y_{A2}$ would be equal to $\frac{p_{A1}}{P_t - p_{A1}} - \frac{p_{A2}}{P_t - p_{A2}}$. Now if we just re solve it then we will get $\frac{P_t p_{A1} - p_{A1} p_{A2} - P_t p_{A1} + p_{A1} p_{A2}}{(P_t - p_{A1})(P_t - p_{A2})}$ which would be equal to $\frac{P_t(p_{A1} - p_{A2})}{p_{B1} p_{B2}}$. So, this we will obtain.

Now, if we just rearrange this relations just to calculate the partial pressure difference we can write $p_{A1} - p_{A2}$ would be equal to $(Y_{A1} - Y_{A2}) \frac{p_{B1} p_{B2}}{P_t}$ and then multiplied by over here $\frac{p_{B1} p_{B2}}{P_t}$. So, after rearranging we will obtain this partial pressure difference.

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Relation between k_G and k_Y

Solution:

Now we have

$$N_A = k_G(p_{A1} - p_{A2}) = \frac{D_{AB} P_t}{RT \delta p_{BLM}} (p_{A1} - p_{A2})$$
$$= \frac{D_{AB} P_t}{RT \delta p_{BLM}} (Y_{A1} - Y_{A2}) \frac{p_{B1} p_{B2}}{P_t} = k_Y (Y_{A1} - Y_{A2})$$

Now, we have N_A is equal to $k_G p_{A1} - p_{A2}$. Now, if we substitute that $p_{A1} - p_{A2}$ which we have obtained we will get $D_{AB} P_t$ divided by $R T \delta p_{BLM} p_{A1} - p_{A2}$. So, we will have finally, k_Y into $Y_{A1} - Y_{A2}$.

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Relation between k_G and k_Y

- since $N_A = k_Y(Y_{A1} - Y_{A2})$, we may write

$$k_Y = \frac{D_{AB} p_{B1} p_{B2}}{RT p_{BLM}} = k_G \frac{p_{B1} p_{B2}}{P_t}$$

- A similar relation between k'_G and k'_Y (in case of equimolar counterdiffusion) applies

Since N_A is equal to k_Y capital $Y_{A1} - Y_{A2}$, we may write capital k_Y would be equal to $D_{AB} p_{B1} p_{B2}$ divided by $R T p_{BLM}$ would be equal to $k_G p_{B1} p_{B2}$ divided by P_t . A similar relations between k'_G and k'_Y in case of the equimolar counter diffusion applies.

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Relation between k'_G and k'_Y

Solution: We know that $p_{A1} - p_{A2} = (Y_{A1} - Y_{A2}) \frac{p_{B1}p_{B2}}{P_t}$ ✓

Now we have $N_A = k'_G(p_{A1} - p_{A2}) = \frac{D_{AB}}{RT\delta}(p_{A1} - p_{A2})$
 $= \frac{D_{AB}}{RT\delta}(Y_{A1} - Y_{A2}) \frac{p_{B1}p_{B2}}{P_t} = k'_Y(Y_{A1} - Y_{A2})$

since $N_A = k'_Y(Y_{A1} - Y_{A2})$, we may write

$$k'_Y = \frac{D_{AB}p_{B1}p_{B2}}{RT\delta} = k'_G \frac{p_{B1}p_{B2}}{P_t}$$
 ✓

Now, we know that $p_{A1} - p_{A2}$ would be equal to $(Y_{A1} - Y_{A2}) \frac{p_{B1}p_{B2}}{P_t}$. So, we have $N_A = k'_G(p_{A1} - p_{A2}) = \frac{D_{AB}}{RT\delta}(p_{A1} - p_{A2})$.

So, if we substitute $\frac{D_{AB}}{RT\delta}$ into $(Y_{A1} - Y_{A2}) \frac{p_{B1}p_{B2}}{P_t}$ into $p_{B1}p_{B2}$ divided by P_t . So, this part we can write k'_Y into $(Y_{A1} - Y_{A2})$, from this we can write k'_Y since $N_A = k'_Y(Y_{A1} - Y_{A2})$ we can write k'_Y would be equal to $\frac{D_{AB}p_{B1}p_{B2}}{RT\delta}$. So, we can get this would be equal to $k'_G \frac{p_{B1}p_{B2}}{P_t}$.

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Relation between k'_G and k'_Y

Solution:

- Conversion among the Gas Phase Mass Transfer Coefficients

$$F = k_G p_{BLM} = k_Y \frac{p_{BLM}}{P_t} = k_c \frac{p_{BLM}}{RT} = k'_G P_t = k'_Y = k'_c \frac{P_t}{RT}$$

- Conversion among the Liquid Phase Mass Transfer Coefficients

$$F = k_x x_{BLM} = k_L x_{BLM} C = k'_L C = k'_L \frac{\rho}{M} = k'_x$$

Conversion among the gas phase mass transfer coefficient, we can write F would be equal to $k_G p_{BLM}$ would be equal to $k_Y p_{BLM}$ divided by P_t which is equal to $k_c p_{BLM}$ by RT .

We can write $k_G P_t$ would be equal to k_Y which is equal to $k_c P_t$ by RT . Now conversion among the liquid phase mass transfer coefficient we can write F is equal to $k_x x_{BLM}$ would be equal to $k_L x_{BLM} C$ is equal to $k'_L C$ would be equal to $k'_L \rho$ by M is equal to k'_x . So, this is in case of the liquid phase and this is in case of the gas phase, for both the systems one is equimolar counter diffusion and another one is diffusion of A through non diffusing B.

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Typical Values of Mass Transfer Coefficients and Film Thickness

- Gas phase mass transfer coefficients, $k_c \sim 10^{-2}$ m/s; film thickness, $\delta \sim 1$ mm
- Liquid phase mass transfer coefficients, $k_c \sim 10^{-5}$ m/s; film thickness, $\delta \sim 0.1$ mm

Now, typical values for the mass transfer coefficient and the film thickness, for gas phase mass transfer coefficient it is k_c is around 10^{-2} metre per second and for a film thickness of δ of 1 millimetre. In case of liquid phase mass transfer coefficient k_c is equal to or k_L is equal to 10^{-5} metre per second and the film thickness is close to δ it would be about 0.1 millimetre.

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Example 1.1

Large volume of pure N_2 gas at atmospheric pressure is flowing over a pool of liquid of methanol, which is evaporating. N_2 is assumed to be insoluble in the liquid. The gas phase mass transfer coefficient of methanol is $k_G = 2 \times 10^{-5}$ kmol/m²s kPa. Assume vapour pressure of methanol at 298K is 10 kPa.

- Calculate k_y , k_c , k_Y and F
- If the diffusivity of methanol at 298K is 2×10^{-5} m²/s, calculate the thickness of the gas film

Now, let us take an example large volume of pure nitrogen gas at atmospheric pressure is flowing over a pool of liquid of methanol, which is evaporating. Nitrogen is assumed to

be insoluble in liquid. The gas phase mass transfer coefficient of methanol which is given k_G is 2×10^{-5} kilomole per metre square second kilopascal.

Assume vapour pressure of methanol at 298 Kelvin is 10 kilopascal. Now, we need to calculate k_y , k_c , k_Y and F . If the diffusivity of methanol at 298 Kelvin is 2×10^{-5} metre square per second calculate the thickness of the gas film. So, here the diffusivity values are also given which is equal to the mass transfer coefficient in the gas phase.

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Example 1.1: Solution

Large volume of pure N_2 gas at atmospheric pressure is flowing over a pool of liquid of methanol, which is evaporating. N_2 is assumed to be insoluble in the liquid. The gas phase mass transfer coefficient of methanol is $k_G = 2 \times 10^{-5}$ kmol/m²s kPa. Assume vapour pressure of methanol at 298K is 10 kPa.

(i) Calculate k_p , k_c , k_Y and F
(ii) If the diffusivity of methanol at 298K is 2×10^{-5} m²/s, calculate the thickness of the gas film

Solution: In this case diffusion of methanol occurs through non diffusing N_2

(i) Given, $P_t = 1 \text{ atm} = 101.3 \text{ kPa}$

$$R = 0.082 \text{ (m}^3 \text{ atm/kmol K)} = 0.082 \times 101.3 \text{ (m}^3 \text{ kPa/kmol K)}$$

$$= 8.3066 \text{ (m}^3 \text{ kPa/kmol K)}$$

$T = 298 \text{ K}$.

$$k_y = k_G P_t = 2 \times 10^{-5} \text{ (kmol/m}^2 \text{ s kPa)} \times 101.3 \text{ kPa} = 2.03 \times 10^{-3} \text{ (kmol/m}^2 \text{ s } \Delta y)$$

Now, in this case the diffusion of methanol occurs through non diffusing nitrogen. So, the first thing it is given that the pure nitrogen is diffusing at atmospheric pressure. So, the total pressure P_t is 1 atmosphere which is 101.3 kilopascal. Now, R is known to us is 0.082 metre cube atmosphere per kilomole Kelvin which is we can write in terms of the metre cube kilopascal kilomole Kelvin in this unit.

And which is 8.3066 metre cube kilopascal per kilomole Kelvin. So, and temperature is given is 298 Kelvin. So, k_y is equal to $k_G P_t$. So, we can just substitute k_G which is given and P_t is known to us. So, k_y would be equal to $2 \times 10^{-5} \times 101.3$ kilometre per metre square second Δy .

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Example 1.1: Solution

Large volume of pure N_2 gas at atmospheric pressure is flowing over a pool of liquid of methanol, which is evaporating. N_2 is assumed to be insoluble in the liquid. The gas phase mass transfer coefficient of methanol is $k_G = 2 \times 10^{-5}$ kmol/m²s kPa. Assume vapour pressure of methanol at 298K is 10 kPa.

(i) Calculate k_p , k_c , k_Y and F

(ii) If the diffusivity of methanol at 298K is 2×10^{-5} m²/s, calculate the thickness of the gas film

Solution: (Cont..)

$$k_c = k_G RT = 2 \times 10^{-5} \text{ (kmol/m}^2\text{s kPa)} \times 8.3066 \text{ (m}^3 \text{ kPa/kmol K)} \times 298\text{K}$$

$$= 0.0495 \text{ m/s}$$

$$p_{A1} = 10 \text{ kPa}$$

$$\text{i.e. } p_{B1} = P_t - p_{A1} = (101.3 - 10) \text{ kPa} = 91.3 \text{ kPa}$$

$$\text{Also, } p_{A2} = 0, \text{ i.e. } p_{B2} = P_t = 101.3 \text{ kPa}$$

k_c would be equal to k_G into $R T$ which is equal to 2 into 10 to the power minus 5 kilomole per metre square second kilopascal into R into T . If we substitute R and T then we can get is 0.0495 metre per second. So, the partial pressure of component A that is nitrogen is given as 10 kilopascal. So, we can calculate p_{B1} which P_t minus p_{A1} which is 91.3 kilopascal and p_{A2} is 0 that is p_{B2} would be equal to P_t which is equal to 101.3 kilopascal. So, both p_{A1} then p_{A2} p_{B1} and p_{B2} are known to us.

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Example 1: Solution

Large volume of pure N_2 gas at atmospheric pressure is flowing over a pool of liquid of methanol, which is evaporating. N_2 is assumed to be insoluble in the liquid. The gas phase mass transfer coefficient of methanol is $k_G = 2 \times 10^{-5}$ kmol/m²s kPa. Assume vapour pressure of methanol at 298K is 10 kPa.

(i) Calculate k_p , k_c , k_Y and F

(ii) If the diffusivity of methanol at 298K is 2×10^{-5} m²/s, calculate the thickness of the gas film

Solution: (Cont..)

$$k_Y = k_G \frac{p_{B1} p_{B2}}{P_t} = 2 \times 10^{-5} \frac{91.3 \times 101.3}{101.3}$$

$$= 1.83 \times 10^{-3} \frac{\text{kmol}}{(\text{s})\text{m}^2(\Delta Y)}$$

$$F = k_G p_{BLM} = 2 \times 10^{-5} \times \frac{p_{B2} - p_{B1}}{\ln\left(\frac{p_{B2}}{p_{B1}}\right)} = 2 \times 10^{-5} \times \frac{101.3 - 91.3}{\ln\left(\frac{101.3}{91.3}\right)}$$

$$= 1.92 \times 10^{-3} \text{ kmol/m}^2 \text{ s}$$

So, we can calculate capital k Y would be equal to k G p B 1 into p B 2 by P t. So, if we substitute this would be equal to 1.83 into 10 to the power minus 3 kilomole per metre square second into the driving force delta Y. F is equal to k G P BLM. So, if we substitute we can calculate P BLM and then if we substitute over here it would be equal to 2 into 10 to the power minus 5 and it is 1.92 into 10 to the power minus 3 kilomole per metre square second.

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Example 1.1: Solution

Large volume of pure N₂ gas at atmospheric pressure is flowing over a pool of liquid of methanol, which is evaporating. N₂ is assumed to be insoluble in the liquid. The gas phase mass transfer coefficient of methanol is $k_G = 2 \times 10^{-5}$ kmol/m²s kPa. Assume vapour pressure of methanol at 298K is 10 kPa.

(i) Calculate k_y , k_c , k_Y and F
(ii) If the diffusivity of methanol at 298K is 2×10^{-5} m²/s, calculate the thickness of the gas film

Solution: (ii) $k_G = \frac{D_{AB} P_t}{R T k_G p_{BLM}} \Rightarrow \delta = \frac{D_{AB} P_t}{R T k_G p_{BLM}}$

$D_{AB} = 2 \times 10^{-5}$ m²/s; $P_t = 101.3$ kPa

$R = 0.082$ (m³ atm/kmol K) = 8.3066 (m³ kPa/kmol K)

$T = 298$ K. $k_G = 2 \times 10^{-5}$ kmol/m²s kPa

So, F is calculated and we need to calculate the thickness of the gas film for k G as we know it is D AB P t by R T delta, delta is missing over here delta and from here we can write, if we rearrange delta would be equal to D AB P t by R T k G p BLM.

So, delta from here we can write delta would be equal to D AB P t divided by R T k G p BLM. So, D AB is given to us and is 2 into 10 to the power minus 5 metre square per second P t is 101.3 kilopascal and R is known to us, temperature is also known to us and k G is given. So, if we substitute D AB P t R T k G and p BLM then.

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Example 1.1: Solution

Large volume of pure N_2 gas at atmospheric pressure is flowing over a pool of liquid of methanol, which is evaporating. N_2 is assumed to be insoluble in the liquid. The gas phase mass transfer coefficient of methanol is $k_G = 2 \times 10^{-5}$ kmol/m²s kPa. Assume vapour pressure of methanol at 298K is 10 kPa.

(i) Calculate k_p , k_c , k_y and F

(ii) If the diffusivity of methanol at 298K is 2×10^{-5} m²/s, calculate the thickness of the gas film

Solution: (ii) (Cont.)

$$p_{BLM} = \frac{p_{B2} - p_{B1}}{\ln\left(\frac{p_{B2}}{p_{B1}}\right)} = \frac{101.3 - 91.3}{\ln\left(\frac{101.3}{91.3}\right)}$$

$$= 96.213$$

$$\delta = 0.425 \text{ mm}$$

P BLM as we have seen earlier p BLM we can calculate for in this case it is 96.213. So, we can calculate delta using the earlier equations which is equal to 0.425 millimetre.

Thank you very much for attending this lecture. In the next lecture we will continue for dimensionless analysis of the mass transfer coefficient.