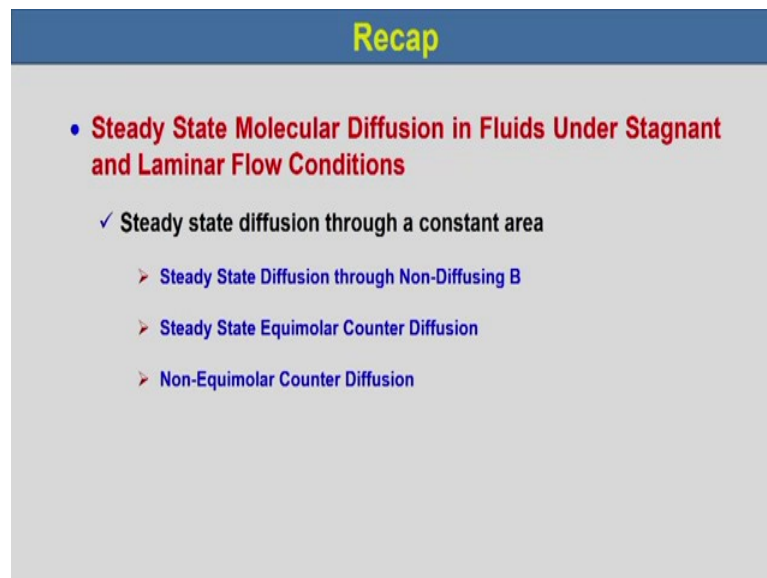


Mass Transfer Operations - I
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Diffusion Mass Transfer - II
Lecture - 05
Diffusion through variable cross-sectional area

Welcome to the 5th lecture on Mass Transfer Operation -I. We are discussing a diffusion mass transfer. Before going through this discussion, let us have recap on our previous lecture.

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The image shows a slide titled "Recap" with a blue header. The main content is a bulleted list of topics covered in the previous lecture. The first bullet point is "Steady State Molecular Diffusion in Fluids Under Stagnant and Laminar Flow Conditions". Under this, there are three sub-bullets: "Steady state diffusion through a constant area", "Steady State Diffusion through Non-Diffusing B", "Steady State Equimolar Counter Diffusion", and "Non-Equimolar Counter Diffusion".

- **Steady State Molecular Diffusion in Fluids Under Stagnant and Laminar Flow Conditions**
 - ✓ Steady state diffusion through a constant area
 - Steady State Diffusion through Non-Diffusing B
 - Steady State Equimolar Counter Diffusion
 - Non-Equimolar Counter Diffusion

We have considered Steady State Molecular Diffusion in fluids under stagnant and laminar flow conditions. Under this the first case we have considered Steady state molecular diffusion through a constant area. Under constant area we have considered the first case is Steady State Diffusion through Non-Diffusing B, Steady State Equimolar Counter Diffusion and the 3rd case we have considered Non-Equimolar Counter Diffusion.

In this lecture we will consider Steady State diffusion through a variable area.

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Steady State Molecular Diffusion Through Variable Area

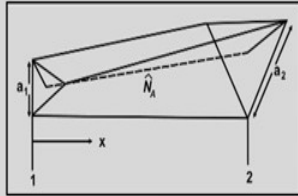


Figure 1: Schematic of diffusion of A through a uniformly tapered geometry

altitude : $\left(\frac{\sqrt{3}}{2} a\right)$ for equilateral triangle

$A = \frac{1}{2} \times a \times \frac{\sqrt{3}}{2} a = \frac{\sqrt{3}}{4} a^2$

- Consider a component A is diffusing at steady state through a equilateral triangle conduit which is tapered uniformly.
- A is diffusing through stagnant, non-diffusing B
- ✓ For an equilateral triangle, the formula for area (where a = length of one side) is:

$$A = \frac{1}{2} \times \text{side} \times \text{altitude}$$

The left hand side of this figure shows the schematic of diffusion of a through a uniformly tapered geometry. So, as you can see the area on the right hand side of the figure of a triangular triangle whose side is a 2 to which is tapered towards the left side of this figure which is that the triangular shape no face which is area of a 1. Obviously, a 1 is at point 1 and a 2 is at point 2 and a 1 is less than a 2. So, consider a component a is diffusing at steady state through a equimolar triangle conduit which is tapered uniformly.

Let us consider A is diffusing through stagnant non-diffusing B component. Now, for an equilateral triangle the formula for area where a is the length of one side can be written as A is equal to half into side into altitude. The altitude of triangle, equilateral triangle is root 3 by 2 into a. A is the side of the triangle. Now, the area A would be half into side is a into the altitude which is root 3 by 2 a which is equal to root 3 by 4 a square.

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Steady State Molecular Diffusion Through Variable Area

Returning to the Fick's Law formula, at position x the flux of A through a triangle of stagnant B can be written as:

$$N_A \left(1 - \frac{p_A}{P_t}\right) = \frac{-D_{AB}}{RT} \frac{dp_A}{dx}$$

Now, $N_A = \frac{\widehat{N}_A}{A} = \frac{4 \widehat{N}_A}{\sqrt{3} a^2}$ Where $\widehat{N}_A =$ rate of diffusion of A kmol/s

$$\Rightarrow -\frac{RT}{D_{AB}} \times \frac{4 \widehat{N}_A}{\sqrt{3} a^2} dx = P_t \frac{dp_A}{P_t - p_A}$$

So, now returning to the Fick's law formula at position x of A through a triangle of stagnant B can be written as $N_A \left(1 - \frac{p_A}{P_t}\right) = \frac{-D_{AB}}{RT} \frac{dp_A}{dx}$.

N_A is the flux, molar flux and p_A is the partial pressure of a component, P_t is the total pressure D_{AB} is the binary diffusion coefficient of component A into the non-diffusing B, R is the universal gas constant, T is the temperature and $\frac{dp_A}{dx}$ is the partial pressure gradient with respect to x . Now, if we define $N_A = \frac{\widehat{N}_A}{A}$ where \widehat{N}_A is the molar flow rate which would be equal to $\frac{4 \widehat{N}_A}{\sqrt{3} a^2}$.

If we substitute the area in this equations as we have calculated before, so N_A as we said $N_A = \frac{\widehat{N}_A}{A}$ is the rate of diffusion of A in kilomole per second. So, the above equations can be written as $-\frac{RT}{D_{AB}} \times \frac{4 \widehat{N}_A}{\sqrt{3} a^2} dx = P_t \frac{dp_A}{P_t - p_A}$.

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Steady State Molecular Diffusion Through Variable Area

Before limits are imposed, it must be remembered that a is a function of x , as the size of triangle uniformly tapers with distance along the duct:

$$a = a_1 + \frac{a_2 - a_1}{x_2 - x_1}(x - x_1)$$

$$= a_1 + \frac{a_2 - a_1}{x_2 - x_1}x - \frac{a_2 - a_1}{x_2 - x_1}x_1$$

$$\therefore \frac{da}{dx} = 0 + \frac{a_2 - a_1}{x_2 - x_1}(1) - \frac{a_2 - a_1}{x_2 - x_1}(0) = \frac{a_2 - a_1}{x_2 - x_1}$$

Or, $dx = \frac{x_2 - x_1}{a_2 - a_1} da$

Now, before limits are imposed it must be remembered that a is a function of x . That means, a is varying with respect to the distance x as the size of the triangle uniformly tapered with distance along the duct. So, we need to have a relation between a and x . Now, a would be equal to a_1 which is at point 1 in this figure plus $a_2 - a_1$ divided by $x_2 - x_1$ into $x - x_1$.

So, this is the relation between the a and the distance. if we just simplify, it would be equal to a_1 plus $a_2 - a_1$ by $x_2 - x_1$ into $x - x_1$. So, if we just differentiate with respect to x this equation, then it would be da/dx would be equal to 0 plus $a_2 - a_1$ by $x_2 - x_1$ into 1 minus $a_2 - a_1$ divided by $x_2 - x_1$ into 0 , so which will be simplified to $a_2 - a_1$ by $x_2 - x_1$. Therefore, dx would be equal to $x_2 - x_1$ divided by $a_2 - a_1$ into da . So, we have a relation between dx and da .

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Steady State Molecular Diffusion Through Variable Area

Substituting x for a , and integrating with limits of p_{A1} at triangle of side a_1 and p_{A2} when the triangle is of side a_2 :

$$\frac{-RT\widehat{N}_A(x_2 - x_1)}{D_{AB}(a_2 - a_1)} \int_{a_1}^{a_2} \frac{4}{\sqrt{3}a^2} da = P_t \int_{p_{A1}}^{p_{A2}} \frac{dp_A}{P_t - p_A}$$

$$\Rightarrow \frac{-4RT\widehat{N}_A(x_2 - x_1)}{\sqrt{3}D_{AB}(a_2 - a_1)} \left[\frac{-1}{a} \right]_{a_1}^{a_2} = -[\ln(P_t - p_A)]_{p_{A1}}^{p_{A2}}$$

$$\Rightarrow \frac{4RT\widehat{N}_A(x_2 - x_1)}{\sqrt{3}D_{AB}(a_2 - a_1)} \left(\frac{1}{a_1} - \frac{1}{a_2} \right) = \ln \left(\frac{P_t - p_{A2}}{P_t - p_{A1}} \right)$$

$$\Rightarrow \widehat{N}_A \left(\frac{a_1 - a_2}{a_1 a_2} \right) = \frac{\sqrt{3}D_{AB}P_t}{4RT} \left(\frac{a_2 - a_1}{x_2 - x_1} \right) \ln \left(\frac{P_t - p_{A2}}{P_t - p_{A1}} \right)$$

Now, if we substitute x for a and integrating with limits of partial pressure of component a at 1 at triangle of side a_1 and partial pressure of component a at point 2, that is p_{A2} when the triangle is at side a_2 , we can write minus $RT \widehat{N}_A$ cap divided by D_{AB} into $x_2 - x_1$ by $a_2 - a_1$ integral a_1 to a_2 4 divided by $\sqrt{3} a^2$ da which would be equal to P_t integral p_{A1} to p_{A2} dp_A divided by $P_t - p_A$.

So, this we can simplify we can integrate this and we will obtain minus $4 RT \widehat{N}_A$ cap divided by $\sqrt{3} D_{AB}$ into $x_2 - x_1$ divided by $a_2 - a_1$ into $\frac{-1}{a}$ with a limit a_1 to a_2 . So, this would be equal to $-\ln(P_t - p_A)$ with a limit p_{A1} to p_{A2} and hence, the left hand side can be written as $4 RT \widehat{N}_A$ cap divided by $\sqrt{3} D_{AB}$ into $x_2 - x_1$ divided by $a_2 - a_1$ into $\frac{1}{a_1} - \frac{1}{a_2}$ if we put the limit in the earlier equation which would be equal to $\ln(P_t - p_{A2})$ divided by $P_t - p_{A1}$.

If we simplify this it would be equal to \widehat{N}_A cap $\frac{a_1 - a_2}{a_1 a_2}$ which would be equal to $\frac{\sqrt{3} D_{AB} P_t}{4 RT} \frac{a_2 - a_1}{x_2 - x_1} \ln \left(\frac{P_t - p_{A2}}{P_t - p_{A1}} \right)$.

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Steady State Molecular Diffusion Through Variable Area

$$\Rightarrow \widehat{N}_A \left(\frac{a_1 - a_2}{a_1 a_2} \right) = \frac{\sqrt{3} D_{AB} P_t}{4RT} \left(\frac{a_2 - a_1}{x_2 - x_1} \right) \ln \left(\frac{P_t - p_{A_2}}{P_t - p_{A_1}} \right)$$
$$\therefore \widehat{N}_A = \frac{\sqrt{3} D_{AB} P_t}{4RT} \frac{a_1 a_2}{x_2 - x_1} \ln \left(\frac{P_t - p_{A_2}}{P_t - p_{A_1}} \right)$$

So, now this equation we can just rearrange and simplify from where we can write \widehat{N}_A would be equal to $\frac{\sqrt{3} D_{AB} P_t}{4RT} \frac{a_1 a_2}{x_2 - x_1} \ln \left(\frac{P_t - p_{A_2}}{P_t - p_{A_1}} \right)$. So, this is the final one equation for Steady State Molecular Diffusion Through Variable Area.

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Example 1

The CO_2 is diffusing through non-diffusing N_2 at steady state in a conduit of 2m long at 300K and a total pressure of 1 atmosphere. The partial pressure of CO_2 at the left end is 20kPa and 5kPa at the other end. The cross section of the conduit is in the shape of an equilateral triangle being 0.0025m at the left end and tapering uniformly to 0.05m at the right end. Calculate the rate of transport of CO_2 . The diffusivity is $D_{AB} = 2 \times 10^{-5} \text{ m}^2/\text{s}$.

Now, let us consider one example. The carbon dioxide is diffusing through non-diffusing nitrogen at steady state in a conduit of 2 metre long at 300 Kelvin and a total pressure of 1 atmosphere. The partial pressure of carbon dioxide at the left end is 20 kilopascal and 5

kilopascal at the other end. The cross section of the conduit is in the shape of an equilateral triangle being 0.025 metre at the left and tapering uniformly to 0.05 metre at the right end.

Now, we need to calculate the rate of transport of carbon dioxide. The diffusivity is given for D_{AB} equal to 2×10^{-5} metre square per second. So, the value which are given diffusion coefficient which is equal to 2×10^{-5} metre square per second.

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Example 1 : Solution

Given that

<p>✓ $D_{AB} = 2 \times 10^{-5} \frac{m^2}{s}$</p> <p>✓ $T = 300K$</p> <p>✓ $P_{A_1} = 20kPa = 20 \times 10^3 Pa$</p> <p>✓ $a_1 = 0.025 m$</p> <p>✓ $x_2 - x_1 = 2 m$</p>	<p>✓ $R = 8314 (m^3.Pa)/(kmol K)$</p> <p>✓ $P_t = 1 atm = 101.3 kPa = 1.013 \times 10^5 Pa$</p> <p>✓ $P_{A_2} = 5kPa = 5 \times 10^3 Pa$</p> <p>✓ $a_2 = 0.05 m$</p>
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R is known to us 8314 metre cube pascal per kilomole kelvin, T is the temperature which is given at 300 kelvin total pressure P_t is 1 atmosphere. So, it is 101.3 kilopascal which is equal to 101.3 into 10 to the power 5 pascal, P_{A_1} is equal to 20 kilopascal which is equal to 20 into 10 to the power 3 pascal, P_{A_2} is 5 kilopascal which is 5 into 10 to the power 3 pascal, a_1 is equal to 0.025 metre, a_2 is equal to 0.05 metre, $x_2 - x_1$ is 2 metre the distance between the two end.

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Example 1 : Solution

Now substituting values in the following equation:

$$\bar{N}_A = \frac{\sqrt{3} D_{AB} P_t}{4RT} \frac{a_1 a_2}{x_2 - x_1} \ln \left(\frac{P_t - p_{A_2}}{P_t - p_{A_1}} \right) \quad \bar{N}_A = \bar{N}_A$$

$$\bar{N}_A = \frac{\sqrt{3} \times 2 \times 10^{-5} \frac{\text{m}^2}{\text{s}} \times 101.3 \times 10^3 \text{ Pa} \times 0.025 \text{ m} \times 0.05 \text{ m}}{4 \times 8314 \frac{(\text{m}^3 \cdot \text{Pa})}{\text{kmol} \cdot \text{K}} \times 300 \text{ K} \times 2 \text{ m}} \ln \left(\frac{101.3 \times 10^3 \text{ Pa} - 5 \times 10^3 \text{ Pa}}{101.3 \times 10^3 \text{ Pa} - 20 \times 10^3 \text{ Pa}} \right)$$

$$\bar{N}_A = \frac{\sqrt{3} \times 2 \times 10^{-5} \times 101.3 \times 10^3 \times 0.025 \times 0.05}{4 \times 8314 \times 300 \times 2} \ln \left(\frac{101.3 \times 10^3 - 5 \times 10^3}{101.3 \times 10^3 - 20 \times 10^3} \right)$$

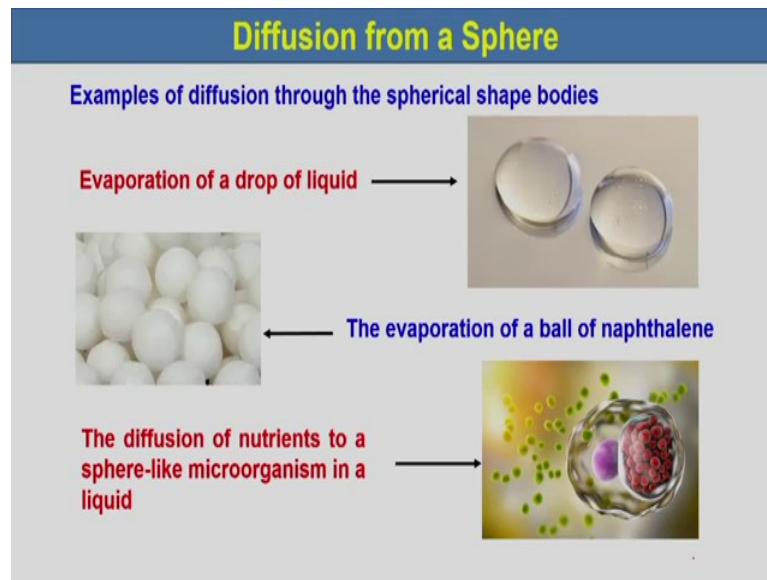
$$\bar{N}_A = \frac{0.22 \times 10^{-2}}{9976800} \ln \left(\frac{96.3 \times 10^3}{81.3 \times 10^3} \right) = 3.74 \times 10^{-11} \text{ kmol/s}$$

Now, substituting the values given in the following equations \bar{N}_A would be equal to $\frac{\sqrt{3} D_{AB} P_t}{4RT} \frac{a_1 a_2}{x_2 - x_1} \ln \left(\frac{P_t - p_{A_2}}{P_t - p_{A_1}} \right)$. So, this \bar{N}_A is equal to \bar{N}_A here. Now, if we substitute the values $\sqrt{3}$ into 2×10^{-5} metre square per second for diffusion coefficient, total pressure multiplied by the total pressure 101.3×10^3 pascal into 0.025 metre into 0.05 metre divided by 4×8314 metre cube pascal by k mole into Kelvin into temperature is 300 Kelvin, the distance $x_2 - x_1$ is 2 metre into \ln $\left(\frac{101.3 \times 10^3 \text{ Pa} - 5 \times 10^3 \text{ Pa}}{101.3 \times 10^3 \text{ Pa} - 20 \times 10^3 \text{ Pa}} \right)$.

The length of side at point 1 of the triangle which is 0.025 metre and then a_2 which is that side of the triangle at point 2, that is 0.05 metre divided by 4×8314 metre cube pascal by k mole into Kelvin into temperature is 300 Kelvin, the distance $x_2 - x_1$ is 2 metre into \ln $\left(\frac{101.3 \times 10^3 \text{ Pa} - 5 \times 10^3 \text{ Pa}}{101.3 \times 10^3 \text{ Pa} - 20 \times 10^3 \text{ Pa}} \right)$.

So, if we just calculate, it would be equal to 3.74×10^{-11} kilo mole per second. So, this is the molar flow rate which is given in the problem to calculate.

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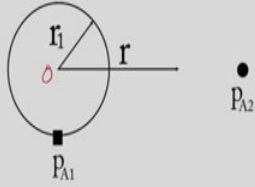
Now, let us consider diffusion from a sphere where the area will vary. Examples of diffusion through the spherical shape bodies one of them is the evaporation of a drop of liquid. So, if we take a drop of liquid and just put on a plane surface, the liquid droplet will slowly evaporate and its diameter will reduce slowly and hence, the area for the diffusions will also vary.

Another example is the evaporations of a ball of naphthalene if we just keep spherical naphthalene ball on to the air, then it will diffuse and slowly its diameter will decrease because the naphthalene will diffuse to the air. This is another example and then, the third examples is the diffusion of nutrient to a sphere like microorganism in a liquid.

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Diffusion from a Sphere

- Assume a constant number of moles N_A of A from a sphere (area = $4\pi r^2$) through stagnant B



From the Fick's law of diffusion, the rate of diffusion can be expressed as:

$$N_A \left(1 - \frac{P_A}{P_{total}} \right) = \frac{-D_{AB}}{RT} \frac{dP_A}{dr} \quad (1) \quad \text{where } N_A = \frac{\bar{N}_A}{4\pi r^2} \quad (2)$$

$$\frac{-RT \bar{N}_A}{4\pi D_{AB} r^2} dr = P_{total} \frac{dP_A}{P_{total} - P_A} \quad (3)$$

So, let us consider a sphere whose centre is O and the radius is r_1 and then, the area radius is varying with time and the partial pressure at the bulk is P_{A2} and at the surface of the sphere is P_{A1} .

Assume a constant number of moles N_A of A from a sphere whose area is equal to $4\pi r^2$ square through stagnant B. Now, if we consider Fick's law of diffusion, the rate of diffusion can be expressed as $N_A \left(1 - \frac{P_A}{P_{total}} \right) = \frac{-D_{AB}}{RT} \frac{dP_A}{dr}$ where N_A is equal to $\frac{\bar{N}_A}{4\pi r^2}$ square. So, we can just substitute this N_A is equal to $\frac{\bar{N}_A}{4\pi r^2}$ square. The equation 1 would be equal to $\frac{-RT \bar{N}_A}{4\pi D_{AB} r^2} dr = P_{total} \frac{dP_A}{P_{total} - P_A}$ by r^2 which would be equal to $P_{total} \frac{dP_A}{P_{total} - P_A}$.

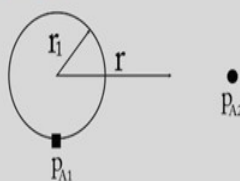
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Diffusion from a sphere

- Integrating with limits of P_{A2} at r_2 and P_{A1} at r_1 gives:

$$\frac{-RT\bar{N}_A}{4\pi D_{AB}P_{total}} \left(\frac{1}{r_1} - \frac{1}{r_2} \right) = \ln \left(\frac{P_{total} - P_{A2}}{P_{total} - P_{A1}} \right) \quad (4)$$
- As $r_1 \ll r_2$, then $1/r_2 \approx 0$:

$$\frac{\bar{N}_A}{4\pi r_1^2} = \frac{D_{AB}P_{total}(P_{A1} - P_{A2})}{RT P_{total} r_1} = N_{A1}, \text{ the flux at the surface} \quad (5)$$



This equation can be simplified if P_{A1} is small compared to P (a dilute gas phase), $P_{BLM} \approx P_{total}$. Also setting $2r_1 = D_1$ (diameter), $C_{A1} = P_{A1}/RT$

$$N_{A1} = \frac{2D_{AB}}{D_1} (C_{A1} - C_{A2})$$

Now, integrating with limits of P_{A2} at r_2 and P_{A1} at r_1 , it gives minus $RT\bar{N}_A$ divided by $4\pi D_{AB}P_{total}$ into $1/r_1 - 1/r_2$ is equal to $\ln(P_{total} - P_{A2} / P_{total} - P_{A1})$. Now, as r_1 is very very less than r_2 which is at far away, then $1/r_2$ would be approximately equal to 0. So, then the above equation 4 would be reduced to \bar{N}_A by $4\pi r_1^2$ into D_{AB} . The above equations would be reduced to \bar{N}_A divided by $4\pi r_1^2$ equal to $D_{AB}P_{total} / (RT P_{total} r_1) (P_{A1} - P_{A2})$ which would be equal to N_{A1} that is the flux at the surface.

P_{BLM} is the log mean partial pressure difference as we have discussed before. Now, these equations can be simplified if p_{A1} is small compared to p that is total pressure P_t since a dilute gas phase, then P_{BLM} would be approximately equal to P_{total} . Now, also if we set $2r_1$ is equal to D_1 that is diameter and then, C_{A1} would be equal to p_{A1} / RT the above equation, this equation. P_{BLM} and P_{total} will be cancelled out and P_{A1} / RT would be C_{A1} which is the concentration at point 1 of component A and P_{A2} by RT would be equal to C_{A2} that is at concentration of component A at point 2. So, the above equations will reduce to N_{A1} would be equal to twice D_{AB} by D_1 into $C_{A1} - C_{A2}$.

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Example 2

A sphere of naphthalene having a radius of 5 mm is suspended in a large volume of still air at 310 K and 1 atm. The partial pressure at the surface of naphthalene at 310K is 50 Pa. Assume dilute gas phase. The D_{AB} of naphthalene in air at 310 K is $6 \times 10^{-6} \text{ m}^2/\text{s}$. Calculate the rate of evaporation of naphthalene from the surface.

Now, let us consider one example. A sphere of naphthalene having a radius of 5 millimetre is suspended in a large volume of still air at 310 Kelvin and 1 atmospheric pressure. The partial pressure at the surface of naphthalene at 310 Kelvin is 50 Pascal. Assume dilute gas phase. The diffusion coefficient of component that is D_{AB} of naphthalene in air is at 310 Kelvin is given as 6 into 10 to the power minus 6 metre square per second.

Now, we need to calculate the rate of evaporation of naphthalene from the surface.

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Example 2 : Solution

A sphere of naphthalene having a radius of 5 mm is suspended in a large volume of still air at 310 K and 1 atm. The partial pressure at the surface of naphthalene at 310K is 50 Pa. Assume dilute gas phase. The D_{AB} of naphthalene in air at 310 K is $6 \times 10^{-6} \text{ m}^2/\text{s}$. Calculate the rate of evaporation of naphthalene from the surface.

Given:

$$D_{AB} = 6 \times 10^{-6} \text{ m}^2/\text{s} \quad P_{A1} = 50 \text{ Pa} \quad P_{A2} = 0$$

$$r_1 = (5/1000) \text{ m} \quad R = 8314 \text{ (m}^3 \cdot \text{Pa)} / (\text{kmol K})$$

$$P_{BLM} = P_t = 1 \text{ atm} = 101.3 \text{ kPa} = 1.013 \times 10^5 \text{ Pa}$$

So, what are the parameters given? Let us note them down. D_{AB} the diffusion coefficient of naphthalene into air. Air is component B which is equal to 6×10^{-6} to the power minus 6 metre square per second. Partial pressure of component A at point 1 is 50 Pascal and at P A2 is equal to 0 because it is dilute gas phase, r_1 is equal to 5 by 1000 metre since it is 5 millimeter, Capital R which is universal gas constant is known to us 8314 metre cube Pascal divided by kilo mole Kelvin. P_{BLM} since it is dilute gas P_{BLM} would be equal to the total pressure P_t which is equal to 1 atmosphere we can write this is 101.3 kilopascal, which is equal to 1.013×10^5 Pascal.

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Example 2 : Solution

Given: $D_{AB} = 6 \times 10^{-6} \text{ m}^2/\text{s}$ $P_{A1} = 50 \text{ Pa}$ $P_{A2} = 0$

$r_1 = (5/1000) \text{ m}$ $R = 8314 \text{ (m}^3 \cdot \text{Pa)} / (\text{kmol K})$

$P_{BLM} = P_t = 1 \text{ atm} = 101.3 \text{ kPa} = 1.013 \times 10^5 \text{ Pa}$

$$\frac{\bar{N}_A}{4\pi r_1^2} = \frac{D_{AB} P_{\text{total}} (P_{A1} - P_{A2})}{R T P_{BM} r_1} = N_{A1}, \text{ the flux at the surface}$$

$P_{BM} = P_{BLM}$

After substitution:

$$\frac{\bar{N}_A}{4\pi r_1^2} = \frac{6 \times 10^{-6} \times 1.013 \times 10^5 \times 50}{8314 \times 310 \times 1.013 \times 10^5 \times 0.005}$$

$$= \frac{303.9 \times 10^{-6}}{13054.22} = 0.023 \times 10^{-6} \text{ kmol} / \text{m}^2 \cdot \text{s}$$

Now, these are the parameters known to us and this is the relations of the flux at the surface. We know for the spherical surface N_A cap or N_A bar divided by $4 \pi r_1$ square is equal to $D_{AB} P_{\text{total}} (P_{A1} - P_{A2})$ divided by $R T P_{BM} r_1$. Here P_{BM} is P_{BLM} that is log mean partial pressure difference which is equal to N_{A1} , the flux at the surface. So, if you substitute the values which are given over there, it would be 6×10^{-6} into 1.013×10^5 into 50 divided by $8314 \times 310 \times 1.013 \times 10^5 \times 0.005$.

So, this would be equal to 303.9×10^{-6} divided by 13054.22 which is equal to 0.023×10^{-6} kilomole per metre square second. So, this is the flux for the ball at the naphthalene at the surface.

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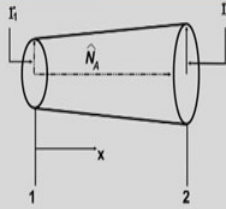
Steady State Equimolar Counter Diffusion

Component A is diffusing at steady state through a circular conduit. At point 1 the radius is r_1 and at point 2 it is r_2 .

We know that

$$N_A = -C D_{AB} \frac{dy_A}{dx} + y_A N$$

$$= -\frac{D_{AB}}{RT} P_t \frac{dy_A}{dx} + y_A N$$



At position x in the conduit, A is diffusing through B by equimolar counter diffusion.

Now for equimolar counter diffusion. $N_A = -N_B$, $N = N_A + N_B = 0$

Now, let us consider steady state equimolar counter diffusion. So, this is a cylindrical geometry uniformly tapered. So, component A is diffusing at steady state through a circular conduit. At point 1 the radius is r_1 and at point 2 it is r_2 .

So, we know that N_A is equal to minus $C D_{AB} \frac{dy_A}{dx}$ plus $y_A N$. Now, it would be minus $\frac{D_{AB}}{RT} P_t \frac{dy_A}{dx}$ plus $y_A N$. At position x in the conduit, A is diffusing through B by equimolar counter diffusion. Now, for equimolar counter diffusion we know that N_A would be equal to minus N_B . So, the total molar flux N would be equal to $N_A + N_B$ which would be equal to 0.

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Steady State Equimolar Counter Diffusion

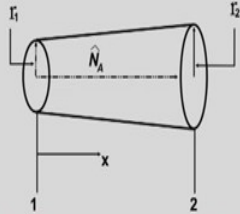
- At position x the flux can be written as

$$N_A = -\frac{D_{AB}}{RT} P_t \frac{d\left(\frac{p_A}{P_t}\right)}{dx} = \frac{-D_{AB}}{RT} \frac{dp_A}{dx} \quad \checkmark$$

$$\Rightarrow N_A = \frac{\widehat{N}_A}{\pi r^2} = \frac{-D_{AB}}{RT} \frac{dp_A}{dx}$$

- Using the geometry as shown, the variable radius r can be related to position x in the path as follows:

$$r = \left(\frac{r_2 - r_1}{x_2 - x_1}\right)x + r_1$$

$$\text{So, } \frac{\widehat{N}_A}{\pi} \int_{x_1}^{x_2} \frac{dx}{\left[\left(\frac{r_2 - r_1}{x_2 - x_1}\right)x + r_1\right]^2} = \frac{-D_{AB}}{RT} \int_{p_{A1}}^{p_{A2}} dp_A$$


At position x the flux can be written as N_A would be equal to minus D_{AB} by RT into P_t into dp_A by P_t divided by dx is equal to minus D_{AB} by RT into dp_A dx . Now, from this we can write N_A would be equal to \widehat{N}_A cap by πr^2 which is equal to minus D_{AB} by RT into dp_A dx . Now, using the geometry as shown the variable radius r can be related to position x in the path as follows r would be equal to r_2 minus r_1 divided by x_2 minus x_1 into x plus r_1 . Now, if we substitute this value of r into over here, the above equations will lead to \widehat{N}_A k f (cap) by π integral x_1 to x_2 dx divided by r_2 minus r_1 by x_2 minus x_1 into x plus r_1 whole square which would be equal to minus D_{AB} by RT integral p_{A1} to p_{A2} into dp_A .

(Refer Slide Time: 28:40)

Steady State Equimolar Counter Diffusion

Let, $C_1 = \left(\frac{r_2 - r_1}{x_2 - x_1} \right)$

$$\therefore \frac{\widehat{N}_A}{\pi} \int_{x_1}^{x_2} \frac{dx}{[C_1 x + r_1]^2} = \frac{-D_{AB}}{RT} \int_{p_{A1}}^{p_{A2}} dp_A$$

Let, $C_1 x + r_1 = z \quad \therefore dx = \frac{dz}{C_1}$

Now substituting:

$$\frac{\widehat{N}_A}{\pi} \int_{z_1}^{z_2} \frac{\frac{dz}{C_1}}{[z]^2} = \frac{-D_{AB}}{RT} \int_{p_{A1}}^{p_{A2}} dp_A$$

$$\Rightarrow \frac{-\widehat{N}_A}{\pi C_1} \left[\frac{1}{z} \right]_{z_1}^{z_2} = \frac{-D_{AB}}{RT} [p_A]_{p_{A1}}^{p_{A2}}$$

So, now let us consider C_1 is equal to $r_2 - r_1$ divided by $x_2 - x_1$. So, we can write the above equation as \widehat{N}_A cap by π integral x_1 to x_2 dx divided by $C_1 x + r_1$ whole square which would be equal to minus D_{AB} by RT integral p_{A1} to p_{A2} dp_A .

Let $C_1 x + r_1$ is equal to z . So, if we just differentiate, then it would be dx would be equal to dz by C_1 . Now, if we substitute the above equations, this equation would be \widehat{N}_A cap by π integral z_1 by z_2 dz by C_1 whole divided by z square would be equal to minus D_{AB} by RT integral p_{A1} to p_{A2} dp_A . So, if we integrate it would be minus \widehat{N}_A cap divided by πC_1 into $\left[\frac{1}{z} \right]_{z_1}^{z_2}$ which would be equal to minus D_{AB} by RT into p_A with a limit p_{A1} to p_{A2} .

(Refer Slide Time: 30:15)

Steady State Equimolar Counter Diffusion

$$\Rightarrow \frac{\widehat{N}_A}{\pi C_1} \left[\frac{1}{z_1} - \frac{1}{z_2} \right] = \frac{D_{AB}}{RT} [p_{A_1} - p_{A_2}]$$

$$\Rightarrow \frac{\widehat{N}_A}{\pi C_1} \left[\frac{1}{C_1 x_1 + r_1} - \frac{1}{C_1 x_2 + r_1} \right] = \frac{D_{AB}}{RT} [p_{A_1} - p_{A_2}]$$

$$\Rightarrow \frac{\widehat{N}_A}{\pi \left(\frac{r_2 - r_1}{x_2 - x_1} \right)} \left[\frac{1}{\left(\frac{r_2 - r_1}{x_2 - x_1} \right) x_1 + r_1} - \frac{1}{\left(\frac{r_2 - r_1}{x_2 - x_1} \right) x_2 + r_1} \right] = \frac{D_{AB}}{RT} [p_{A_1} - p_{A_2}]$$

$$\Rightarrow \widehat{N}_A = \frac{D_{AB}}{RT} \frac{\pi \left(\frac{r_2 - r_1}{x_2 - x_1} \right) [p_{A_1} - p_{A_2}]}{\left[\frac{1}{\left(\frac{r_2 - r_1}{x_2 - x_1} \right) x_2 + r_1} - \frac{1}{\left(\frac{r_2 - r_1}{x_2 - x_1} \right) x_1 + r_1} \right]}$$

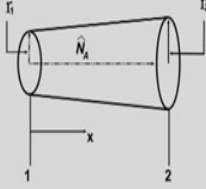
So, this would be written as \widehat{N}_A cap by π into C_1 whole into $\frac{1}{z_1} - \frac{1}{z_2}$ which is equal to $\frac{D_{AB}}{RT} [p_{A_1} - p_{A_2}]$.

So, from there if we substitute z_1 and z_2 , then it would be \widehat{N}_A cap by πC_1 into $\frac{1}{C_1 x_1 + r_1} - \frac{1}{C_1 x_2 + r_1}$ which would be equal to $\frac{D_{AB}}{RT} [p_{A_1} - p_{A_2}]$. So, now if we substitute C_1 , it would be \widehat{N}_A cap divided by $\pi \left(\frac{r_2 - r_1}{x_2 - x_1} \right)$ whole into $\frac{1}{\left(\frac{r_2 - r_1}{x_2 - x_1} \right) x_1 + r_1} - \frac{1}{\left(\frac{r_2 - r_1}{x_2 - x_1} \right) x_2 + r_1}$ which would be equal to $\frac{D_{AB}}{RT} [p_{A_1} - p_{A_2}]$.

So, finally if we rearrange the molar flow rate \widehat{N}_A cap would be equal to $\frac{D_{AB}}{RT}$ into $\pi \left(\frac{r_2 - r_1}{x_2 - x_1} \right) [p_{A_1} - p_{A_2}]$ whole divided by $\left[\frac{1}{\left(\frac{r_2 - r_1}{x_2 - x_1} \right) x_2 + r_1} - \frac{1}{\left(\frac{r_2 - r_1}{x_2 - x_1} \right) x_1 + r_1} \right]$. So, we would obtain this relation.

(Refer Slide Time: 32:12)

Steady State Equimolar Counter Diffusion

$$\Rightarrow \widehat{N}_A = \frac{D_{AB}}{RT} \frac{\pi \left(\frac{r_2 - r_1}{x_2 - x_1} \right) [p_{A1} - p_{A2}]}{\left[\frac{r_2 - r_1}{x_2 - x_1} x_2 + r_1 - \frac{r_2 - r_1}{x_2 - x_1} x_1 + r_1 \right]}$$


If $x_1 = 0$ and $x_2 = L$ i.e. $x_2 - x_1 = L$

$$\Rightarrow \widehat{N}_A = \frac{D_{AB}}{RT} \frac{\pi \left(\frac{r_2 - r_1}{L} \right) [p_{A1} - p_{A2}]}{\left[\frac{1}{r_2} - \frac{1}{r_1} \right]} = \frac{D_{AB}}{RT} \frac{\pi r_1 r_2}{L} (p_{A1} - p_{A2})$$

Now, after now if we put x_1 is equal to 0 and x_2 is equal to L and $x_2 - x_1$, that means x_2 minus x_1 is L . The total distance from point 1 to point 2 the above equations, we can write \widehat{N}_A would be equal to D_{AB} by RT into $\pi r_2 - r_1$ by L into $p_{A1} - p_{A2}$ divided by $1/r_2 - 1/r_1$ which would be equal to D_{AB} by RT into $\pi r_1 r_2$ into L into the partial pressure gradient $p_{A1} - p_{A2}$.

(Refer Slide Time: 32:55)

Example 3

The CO_2 is diffusing at steady state through N_2 by equimolar counter diffusion in a conduit 2 m long at 300K and a total pressure of 1 atmosphere. The partial pressure of CO_2 at the left end is 20 kPa and 5 kPa at the other end. The cross section of the conduit is in the shape of cylindrical of radius is 0.025m at the left end and tapering uniformly to a radius of 0.05m at the right end. Calculate the molar flow rate of CO_2 . The diffusivity is $D_{AB} = 2 \times 10^{-5} \text{ m}^2/\text{s}$.

Now, let us consider one example. The carbon dioxide is diffusing at steady state through nitrogen by equimolar counter diffusion in a conduit of 2 meter length at 300 Kelvin and

a total pressure of 1 atmosphere. The partial pressure of carbon dioxide at the left end is 20 kilo pascal and 5 kilo pascal at the other end. The cross section of the conduit is in the shape of cylindrical of radius 0.025 metre at the left end and tapering uniformly to a radius of 0.05 meter at the right end. We need to calculate the molar flow rate of carbon dioxide. The diffusivity of the component is D_{AB} is equal to 2×10^{-5} meter square per second.

(Refer Slide Time: 33:54)

Example 3 : Solution

The CO_2 is diffusing at steady state through N_2 by equimolar counter diffusion in a conduit 2 m long at 300K and a total pressure of 1 atmosphere. The partial pressure of CO_2 at the left end is 20 kPa and 5 kPa at the other end. The cross section of the conduit is in the shape of cylindrical of radius is 0.025m at the left end and tapering uniformly to a radius of 0.05m at the right end. Calculate the molar flow rate of CO_2 . The diffusivity is $D_{AB} = 2 \times 10^{-5} \text{ m}^2/\text{s}$.

Given that

$$D_{AB} = 2 \times 10^{-5} \frac{\text{m}^2}{\text{s}} \quad R = 8314 \text{ (m}^3 \cdot \text{Pa)} / (\text{kmol K})$$

$$T = 300\text{K} \quad P_t = 1 \text{ atm} = 101.3 \text{ kPa} = 1.013 \times 10^5 \text{ Pa}$$

$$P_{A_1} = 20 \text{ kPa} = 20 \times 10^3 \text{ Pa} \quad P_{A_2} = 5 \text{ kPa} = 5 \times 10^3 \text{ Pa}$$

$$a_1 = 0.025 \text{ m} \quad a_2 = 0.05 \text{ m} \quad L = x_2 - x_1 = 2 \text{ m}$$

Now, the data which are given can be noted D_{AB} is equal to 2×10^{-5} meter square per second, R is 8314 meter cube Pascal per kilo mol Kelvin, T is the temperature which is 300 Kelvin, P_t is the total pressure which is at atmospheric pressure which is equal to 101.3 kilo Pascal which equal to 1.013×10^5 Pascal, P_{A_1} is 20 kilo Pascal which is equal to 20×10^3 Pascal and P_{A_2} is 5 kilo Pascal which is 5×10^3 Pascal, a_1 is 0.025 meter and a_2 is 0.05 meter, the distance between the two end or two point x_1 and x_2 point which is L is equal to $x_2 - x_1$ which is 2 meter.

(Refer Slide Time: 35:03)

Example 3 : Solution

Given that

$$D_{AB} = 2 \times 10^{-5} \frac{\text{m}^2}{\text{s}} \quad R = 8314 \text{ (m}^3\text{.Pa)/(kmol K)}$$

$$T = 300\text{K} \quad P_t = 1 \text{ atm} = 101.3 \text{ kPa} = 1.013 \times 10^5 \text{ Pa}$$

$$P_{A_1} = 20\text{kPa} = 20 \times 10^3 \text{ Pa} \quad P_{A_2} = 5\text{kPa} = 5 \times 10^3 \text{ Pa}$$

$$a_1 = 0.025 \text{ m} \quad a_2 = 0.05 \text{ m} \quad L = x_2 - x_1 = 2 \text{ m}$$

Now substituting values in the following equation:

$$\bar{N}_A = \frac{D_{AB} \pi r_1 r_2}{RT L} (P_{A_1} - P_{A_2})$$

$$\bar{N}_A = \frac{2 \times 10^{-5} \frac{\text{m}^2}{\text{s}} \times 3.14 \times 0.025\text{m} \times 0.05\text{m}}{8314 \frac{\text{(m}^3\text{.Pa)}}{\text{kmol.K}} \times 300\text{K} \times 2\text{m}} (20 \times 10^3 - 5 \times 10^3)$$

Now, if we substitute the parameters which are given D_{AB} , R , T , P_t and partial pressure of component a at 1 and partial pressure of component two component a at 2 and then, the other values a_1 , a_2 and the length.

(Refer Slide Time: 35:30)

Example 3 : Solution

The CO_2 is diffusing at steady state through N_2 by equimolar counter diffusion in a conduit 2 m long at 300K and a total pressure of 1 atmosphere. The partial pressure of CO_2 at the left end is 20 kPa and 5 kPa at the other end. The cross section of the conduit is in the shape of cylindrical of radius is 0.025m at the left end and tapering uniformly to a radius of 0.05m at the right end. Calculate the molar flow rate of CO_2 . The diffusivity is $D_{AB} = 2 \times 10^{-5} \text{ m}^2/\text{s}$.

$$\bar{N}_A = \frac{2 \times 10^{-5} \frac{\text{m}^2}{\text{s}} \times 3.14 \times 0.025\text{m} \times 0.05\text{m}}{8314 \frac{\text{(m}^3\text{.Pa)}}{\text{kmol.K}} \times 300\text{K} \times 2\text{m}} (20 \times 10^3 - 5 \times 10^3)$$

$$= \frac{0.00785 \times 15}{4988400} \times 10^{-2} \text{ kmol/s}$$

$$\bar{N}_A = 2.36 \times 10^{-10} \text{ kmol/s}$$

So, if you substitute these values in the following equation, you would obtain \bar{N}_A bar would be equal to this which would be equal to 0.00785 into 15 divided by 4988400 into 10 to the power minus 2 kilo mole per second which is equal to \bar{N}_A bar is equal to 2.36 into 10 to the power minus 10 kilo mole per second.

Thank you very much for hearing this lecture.