

Fick's Law for Binary Systems

Molar Fluxes in Binary Systems

$$N_A = cx_A v_A$$

$$N = N_A + N_B$$

$$v_M = \frac{N}{c} = \frac{N_A + N_B}{c} = x_A v_A + x_B v_B$$

Here we have assumed 1-D.
We could use vectors for
fluxes & velocities...

- A species "A" in mixture of A & B.
- N_A molar flux of A.
- v_A velocity of A.
- x_A mole fraction of A.
- c mixture molar concentration.
- N total molar flux.
- v_M mixture molar-averaged velocity.

$$J_A = N_A - cx_A v_M$$

molar diffusive
flux of A
(relative to v_M)

molar convective
flux of A
(carried by v_M)

Fick's "Law" $J_A = -cD_{AB} \nabla x_A$

- Fick's law is a MODEL for J_A (has limitations!)
- $D_{AB} = D_{BA}$
- $J_A = -J_B$
- For $C > 2$ components, everything changes!
(graduate school, anyone?)

Often we know something about $x_A(z)$
and v_M . Fick's law lets us get N_A .

- J_A - molar diffusive flux of A relative to a molar averaged velocity. (motion of A relative to the mixture motion)
- Diffusive fluxes are only defined relative to a convective and total flux!
- Diffusive and convective fluxes are NOT independent (they must sum to N_A).

$$J_A = -cD_{AB} \frac{dx_A}{dz}$$

$$N_A = cx_A v_M - cD_{AB} \frac{dx_A}{dz}$$

$$= x_A N - cD_{AB} \frac{dx_A}{dz}$$

$$N_B = x_B N - cD_{AB} \frac{dx_B}{dz}$$

“Mixture Velocities”

Motorcycle: $u_m = 50$

Dump Truck: $u_t = 30$

If there are n_m motorcycles and n_t trucks on the road, what is the average velocity?

Number averaged: $v_{\#} = \frac{n_m v_m + n_t u_t}{n_m + n_t}$

$$u_t = v_{\#} + v_{t,diff}^{\#}$$

$$u_m = v_{\#} + v_{m,diff}^{\#}$$

what if
 $n_m = n_t$?

Mass averaged: $v = \frac{n_m m_m v_m + n_t m_t u_t}{n_m m_m + n_t m_t}$

$$u_t = v + v_{t,diff}$$

$$u_m = v + v_{m,diff}$$

what if
 $n_m = n_t$ and
 $m_t \gg m_m$?

Molar flux
(analogous to number flux)

$$N_A = c_A v_M + J_A$$

$$= x_A c v_M + J_A$$

$$= x_A N + J_A$$

$$v_A = \frac{N_A}{c_A} = \frac{N_A}{c x_A}$$

$$v_{A,diff} = J_A / c_A$$

$$v_M = \frac{N}{c} = \frac{N_A + N_B}{c}$$

Fick's Law gives us a relationship between J_A & x_A .

Steady-State Diffusion in Binary Systems

$$N_A = x_A \underbrace{(N_A + N_B)}_N + J_A$$

$$J_A = -cD_{AB} \frac{dx_A}{dz}$$

$$n_A = x_A \underbrace{(n_A + n_B)}_n + AJ_A$$

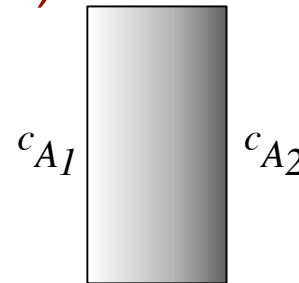
If there is no “bulk flow” ($N=0$) then

$$\begin{aligned} N_A &= -cD_{AB} \frac{dx_A}{dz} \\ &= -D_{AB} \frac{dc_A}{dz} \quad (\text{if } c \text{ is constant}) \end{aligned}$$

Be careful with areas!
(flux vs. flow rate)

Planar system with constant N_A (or n_A):

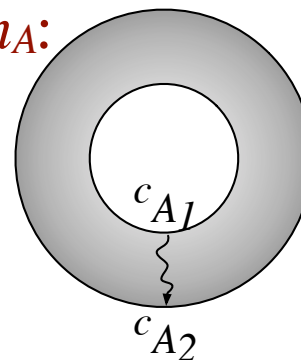
$$N_A = -D_{AB} \left(\frac{c_{A2} - c_{A1}}{z_2 - z_1} \right)$$



Note: in cylindrical & spherical coordinates, constant n_A does not imply constant N_A .

Concentric cylinders with constant n_A :

$$n_A = -2\pi L D_{AB} \left(\frac{c_{A2} - c_{A1}}{\ln(r_2/r_1)} \right)$$



Spherical shell with constant n_A :

$$n_A = -4\pi r_1 r_2 D_{AB} \left(\frac{c_{A2} - c_{A1}}{r_2 - r_1} \right)$$

The Molar Balance Equations (Again)

$$\mathbf{N}_i = c_i \mathbf{v}_M + \mathbf{J}_i = x_i \mathbf{N} + \mathbf{J}_i$$

Integral Forms

In terms of “total”
species fluxes:

$$\frac{d}{dt} \int_V c_i dV = - \int_S \mathbf{N}_i \cdot \mathbf{a} dS + \int_V S_i dV$$

In terms of “convective” and
“diffusive” species fluxes:

$$\frac{d}{dt} \int_V c_i dV = - \int_S x_i c \mathbf{v}_M \cdot \mathbf{a} dS - \int_S \mathbf{J}_i \cdot \mathbf{a} dS + \int_V S_i dV$$

Differential Forms

In terms of “total”
species fluxes:

$$\frac{\partial c_i}{\partial t} = -\nabla \cdot \mathbf{N}_i + S_i$$

In terms of “convective” and
“diffusive” species fluxes:

$$\frac{\partial c_i}{\partial t} = -\nabla \cdot (c_i \mathbf{v}_M) - \nabla \cdot \mathbf{J}_i + S_i$$

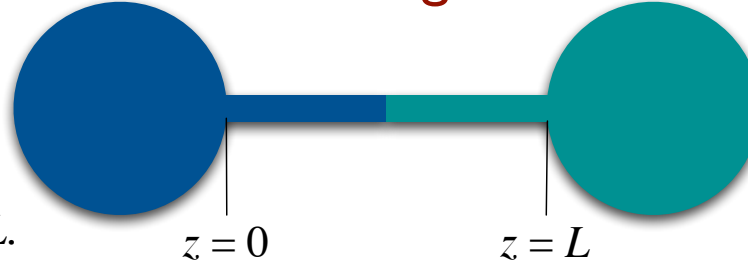
For a binary system of “A” and “B”: $\mathbf{J}_A = -cD_{AB} \nabla x_A$

Fick's Law Example: Equimolar Counter-diffusion

Assume:

- T, p are constant
- No reaction
- 1-D domain $z=[0, L]$
- Compositions are known at domain boundaries $z = 0, z = L$.

Find flux through the tube.



$$x_A = x_A^0 \quad z = 0$$

$$x_A = x_A^L \quad z = L$$

Equimolar counter-diffusion: for every mole of A that moves to the left, a mole of B moves to the right.

Total flux must be zero (closed system, constant T, p).

Therefore, $N = cv_M = 0$ so $v_M = 0$.

$$\frac{d}{dt} \int_V c_i dV = - \int_S x_i cv_M \cdot \mathbf{a} dS - \int_S \mathbf{J}_i \cdot \mathbf{a} dS + \int_V \mathcal{R}_i dV$$

pseudo-steady state v_M = 0 no reaction

Break the surface integral into pieces:

Tube sides: $\int_S \mathbf{J}_A \cdot \mathbf{a} dS = 0$

Tube face at $z=0$: $\mathbf{a} = -\hat{z} \int_S \mathbf{J}_A \cdot \mathbf{a} dS = -A_c J_A^0$

Tube face at $z=L$: $\mathbf{a} = \hat{z} \int_S \mathbf{J}_A \cdot \mathbf{a} dS = A_c J_A^L$

∴ Tube mole balance gives: $A_c J_A^L - A_c J_A^0 = 0$

$J_A^0 = J_A^L$

Note: we could have done this integral balance on any segment of the tube and arrived at the same conclusion regarding J_A . Therefore, it must be constant! Since $J_A = N_A$, N_A is also constant!

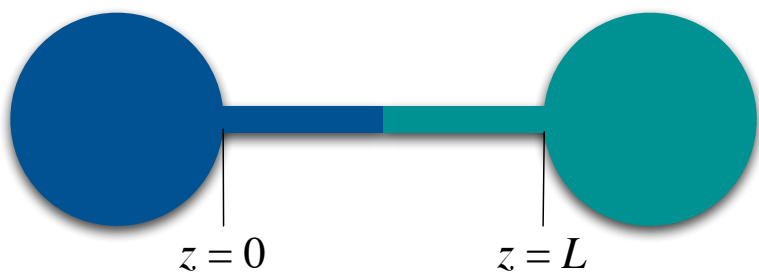
$$\begin{aligned} N_A &= x_A N + J_A \\ &= J_A = -cD_{AB} \frac{dx_A}{dz} \end{aligned}$$

$$N_A dz = -cD_{AB} dx_A$$

$$\int_0^L N_A dz = - \int_{x_A^0}^{x_A^L} cD_{AB} dx_A$$

$$N_A \int_0^L dz = -cD_{AB} \int_{x_A^0}^{x_A^L} dx_A$$

$$N_A = \frac{-cD_{AB}}{L} (x_A^L - x_A^0)$$



We previously showed:

- $N_A = J_A$ (no bulk flow/convection)
- N_A is constant (so J_A is constant)

What is the species mole fraction profile through the tube?

$$J_A = -cD_{AB} \frac{dx_A}{dz} = \alpha$$

$$dx_A = -\frac{\alpha}{cD_{AB}} dz$$

$$\int_{x_A^0}^{x_A} dx_A = -\frac{\alpha}{cD_{AB}} \int_0^z dz$$

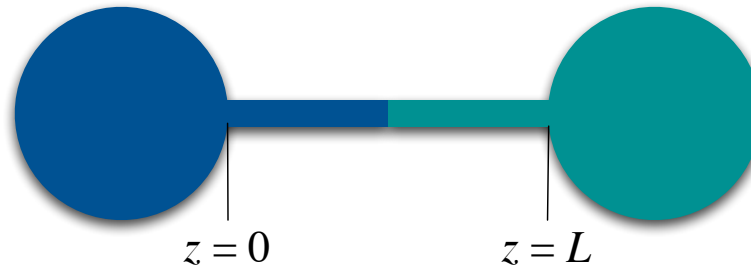
$$x_A - x_A^0 = -\frac{\alpha}{cD_{AB}} z$$

Note: to determine α , we can use $x_A(L) = x_A^L$

$$x_A = x_A^0 + (x_A^L - x_A^0) \frac{z}{L}$$

“Bulb” Balances

Find the composition in each bulb as a function of time.



$$N_A = \frac{-cD_{AB}}{L} (x_A^L - x_A^0)$$

Pseudo-steady state:

The tube is at steady state (adjusts to the slowly changing bulb compositions quickly).

$$\frac{d}{dt} \int_V c_i dV = - \int_S x_i c \cancel{v_M} \cdot \mathbf{a} dS - \int_S \mathbf{J}_i \cdot \mathbf{a} dS + \int_V \cancel{S_i} dV$$

(Note: v_M and S_i are crossed out with red arrows)

Mole balance on A relating to steady-state conditions:

$$\underbrace{cx_A^0 V_0 + cx_A^L V_L}_{\text{at time } t} = \underbrace{cx_A^\infty (V_0 + V_L)}_{\text{at } t=\infty}$$

$$x_A^L = x_A^\infty \left(1 + \frac{V_0}{V_L}\right) - x_A^0 \frac{V_0}{V_L}$$

For the “left” bulb: $cV_0 \frac{dx_A^0}{dt} = -N_A^0 A_c$ $V_0 \frac{dx_A^0}{dt} = \frac{A_c D_{AB}}{L} (x_A^L - x_A^0)$

We need to eliminate x_A^L

$$\frac{dx_A^0}{dt} = \frac{A_c D_{AB}}{V_0 L} [(x_A^\infty - x_A^0) (1 + V_0/V_L)]$$

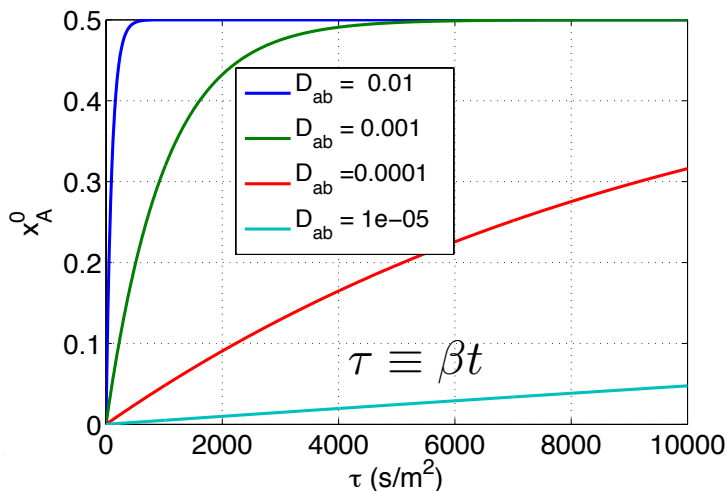
$$= \beta D_{AB} (x_A^\infty - x_A^0) \quad \beta \equiv \frac{A_c}{V_0 L} \left(1 + \frac{V_0}{V_L}\right)$$

“Geometry factor”

Separate & solve...

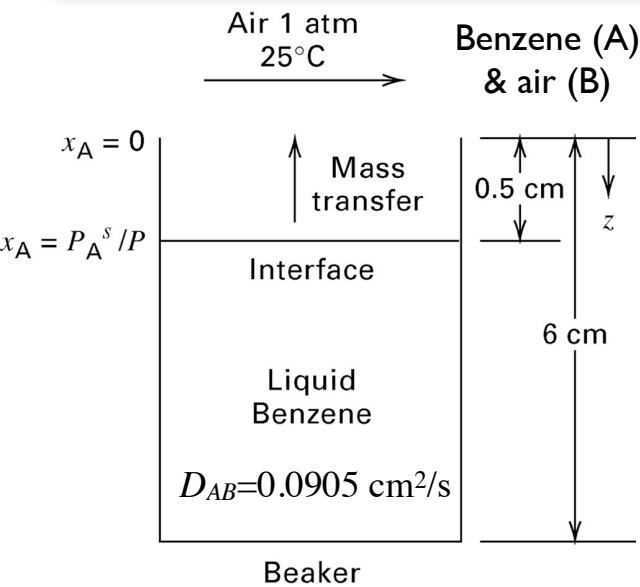
$$x_A^0 = x_A^\infty + (x_{A,0}^0 - x_A^\infty) \exp(-\beta D_{AB} t)$$

A nice way to determine D_{AB} experimentally!



Example: Evaporation from a Beaker

(Unimolecular Diffusion)



Determine $x_A(z)$.

$$N_A = x_A N + J_A$$

What do we know about N_A & N_B ?

$$\begin{aligned} N_A &= x_A N_A + J_A \\ &= \frac{J_A}{1 - x_A} \end{aligned}$$

(1- x_A) accounts for “bulk flow.”
In very dilute systems ($x_A \rightarrow 0$),
this effect is small relative to J_A .

$$N_A = \frac{-cD_{AB}}{1 - x_A} \frac{dx_A}{dz} \quad \longrightarrow \quad \frac{-N_A}{cD_{AB}} \int_{z_0}^z dz = \int_{x_A^0}^{x_A} \frac{dx_A}{1 - x_A}$$

$$N_A = \frac{cD_{AB}}{z - z_0} \ln \left(\frac{1 - x_A}{1 - x_A^0} \right)$$

or

$$x_A = 1 - (1 - x_A^0) \exp \left[\frac{N_A(z - z_0)}{cD_{AB}} \right]$$

Note: from $x_A(z)$ and N_A you can determine anything else about the system (e.g. $J_A(z)$).

1. Estimate gas-phase composition of benzene at the vapor-liquid interface (from equilibrium thermo).
2. Determine N_A from x_A at $z=0$.
3. Determine $x_A(z)$.

The Log-Mean

$$N_A = \frac{cD_{AB}}{z - z_0} \ln \left(\frac{1 - x_A}{1 - x_A^0} \right)$$

Log-mean of x_A at the two ends of the diffusion path.

$$(1 - x_A)_{LM} = (x_B)_{LM} = \frac{(1 - x_{A_2}) - (1 - x_{A_1})}{\ln \left[\frac{1 - x_{A_2}}{1 - x_{A_1}} \right]}$$
$$= \frac{x_{A_1} - x_{A_2}}{\ln \left[(1 - x_{A_2}) / (1 - x_{A_1}) \right]}$$

$$N_A = \frac{cD_{AB}}{(1 - x_A)_{LM}} \frac{-\Delta x_A}{\Delta z}$$

This makes things look a bit “cleaner” and allows us to express N_A in terms of Δx_A .

You will see this used more when we start dealing with Mass-Transfer Coefficients (soon)...

Comments on Fick's Law

- 📌 In this class, we typically assume that the total molar concentration, c , is constant.
 - This is usually reasonable for isothermal, isobaric systems or for liquid systems.
- 📌 We have only considered binary systems.
 - For multicomponent systems, things become considerably more complex.
- 📌 Other driving forces:
 - other species can cause strange diffusion (push a species against its gradient) for $C > 2$ components.
 - pressure gradients (centrifugation)
 - thermal gradients (Soret effect)
 - In general, the chemical potential is the correct driving force for diffusion.