Physical Principles in Biology Biology 3550 Spring 2024

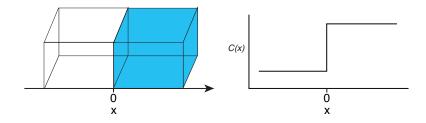
Lecture 16:

Diffusion Continued: Fick's First and Second Laws

Wednesday, 14 February 2024

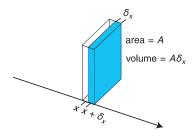
©David P. Goldenberg University of Utah goldenberg@biology.utah.edu

An Idealized Macroscopic Diffusion Experiment



■ How will plot of C(x) versus x change with time?

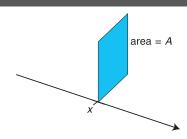
Diffusion Across a Thin Slice of Volume



- Slices are δ_x thick and have cross-sectional area across the *x*-axis of *A*
- Volume of each slice is $A\delta_{\times}$
- During time τ , all of the molecules will move (on average) the distance δ_x along the x-axis.
- In a given slice, 1/2 of the molecules will move to the right and half to the left.

Fick's First Law of Diffusion

$$J = -\frac{\delta_x^2}{2\tau} \frac{dC}{dx}$$



Symbols:

- J = flux of molecules per unit area per unit time:
- $\delta_x = \text{RMS}$ step length along the *x*-direction.
- τ = average duration of random steps.
- $\frac{dC}{dx}$ = derivative of concentration with x, "concentration gradient."
- If concentration increases with x, flux is in the negative x direction.

The Diffusion Coefficient, D

Consider the term from Fick's first law: $\frac{\delta_x^2}{2\tau}$

Both δ_{\times} and τ are parameters describing the random walk of molecules (or larger particles) undergoing diffusion, and are constant for a given type of particle under defined solution conditions.

■ Define a new parameter, the diffusion coefficient, *D*:

$$D=rac{\delta_x^2}{2 au}$$

The usual form of Fick's first law:

$$J = -D\frac{dC}{dx}$$

■ *D* can be experimentally determined, without knowing anything about the microscopic random walk steps.

What are the units of the diffusion coefficient: $D = \frac{\delta_x^2}{2\tau}$?

- A) mol/s
- B) m^2/s
- C) mol/m/s
- **D)** m/s

What are the units of the concentration gradient: $\frac{dC}{dx}$?

- A) mol/m
- B) mol/m²
- C) mol/m³
- D) mol/m⁴

Dimensional Analysis of Fick's First Law

- The diffusion coefficient: $D = \frac{\delta_x^2}{2\tau}$, units: m²/s
- The concentration gradient: $\frac{dC}{dx}$

C (concentration) units: moles/m³

x (distance) units: m

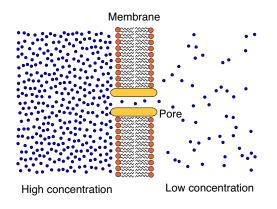
$$\frac{dC}{dx}$$
 has units of $\frac{\text{moles/m}^3}{\text{m}} = \frac{\text{moles}}{\text{m}^4}$

■ Flux, J

$$J = -D \frac{dC}{dx} = \frac{m^2 \text{ moles}}{s \cdot m^4} = \frac{\text{moles}}{s \cdot m^2}$$

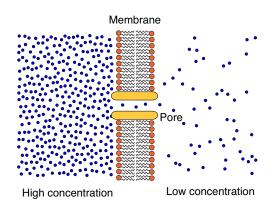
moles per unit time per unit area, as originally defined.

Diffusion of Molecules Across a Cellular Membrane



Some plausible values:

- Diffusion coefficient for a small molecule: $\approx 10^{-10} \, \text{m}^2/\text{s}$ (to be considered later)
- Thickness of a biological membrane: ≈ 3 nm
- Diameter of a molecular pore: ≈ 1 nm
- Concentrations: 50 mM and 5 mM



- Thickness of membrane: \approx 3 nm
- Diameter of pore: $\approx 1 \, \text{nm}$
- Concentrations: 50 mM and 5 mM

Estimate the concentration gradient:

$$\frac{dC}{dx} pprox \frac{\Delta C}{\Delta x}$$

- **A)** $1.5 \times 10^3 \, \text{moles/m}^4$
- **B)** $1.5 \times 10^6 \, \text{moles/m}^4$
- **C)** $1.5 \times 10^9 \, \text{moles/m}^4$
- D) $1.5 \times 10^{10} \, \text{moles/m}^4$
- **E)** $4.5 \times 10^{10} \, \text{moles/m}^4$

Applying Fick's First Law

Approximate the concentration gradient:

$$\frac{dC}{dx} \approx \frac{\text{concentration difference}}{\text{membrane thickness}} = \frac{50 \text{ mM} - 5 \text{ mM}}{3 \text{ nm}} = 15 \text{ mM/nm}$$

Convert this to units with moles and meters:

$$15 \, \text{mM/nm} = \frac{0.015 \, \text{moles/L}}{10^{-9} \, \text{m}} \times \frac{10^3 \, \text{L}}{\text{m}^3} = 1.5 \times 10^{10} \, \text{moles/m}^4$$

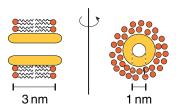
Flux from Fick's first law:

$$J = -D \frac{dC}{dx} = -10^{-10} \, \mathrm{m^2/s} \times 1.5 \times 10^{10} \, \mathrm{moles/m^4} = -1.5 \, \mathrm{moles/(s \cdot m^2)}$$

Negative sign indicates that diffusion is in the direction opposite to the concentration gradient.

Diffusion Through a Membrane Pore

■ The pore:



Diameter = 1 nm

Area =
$$\pi r^2 = \pi (0.5 \text{ nm})^2 = 7.8 \times 10^{-19} \text{ m}^2$$

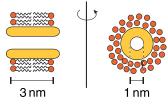
Flux:

$$J = -1.5 \, \text{moles/(s} \cdot \text{m}^2)$$

Moles per second through the pore:

$$1.5 \text{ moles/(s} \cdot \text{m}^2) \times 7.8 \times 10^{-19} \text{ m}^2 = 1.2 \times 10^{-18} \text{ moles/s}$$

Diffusion Through a Membrane Pore



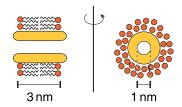
Moles per second through the pore:

$$1.5 \, \text{moles/(s} \cdot \text{m}^2) \times 7.8 \times 10^{-19} \, \text{m}^2 = 1.2 \times 10^{-18} \, \text{moles/s}$$

Molecules per second through the pore:

$$1.2 \times 10^{-18}$$
 moles/s $\times 6.02 \times 10^{23}$ molecules/mole = 7×10^{5} molecules/s

How Many Molecules are in the Pore?



Pore volume:

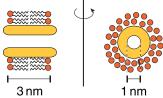
$$V = \pi r^2 \times I = \pi (0.5 \text{ nm})^2 \times 3 \text{ nm}$$

= 2.4 nm³ × $(10^{-9} \text{ m/nm})^3 = 2.4 \times 10^{-27} \text{ m}^3$
= 2.4 × $10^{-27} \text{ m}^3 \times 10^3 \text{ L/m}^3 = 2.4 \times 10^{-24} \text{ L}$

Assume that the concentration in the pore is the mean of the concentrations on the two sides:

$$C = (50 \text{ mM} - 5 \text{ mM})/2 = 22 \text{ mM} = 0.022 \text{ moles/L}$$

How Many Molecules are in the Pore?



- Concentration: 0.022 moles/L
- Moles in pore:

$$0.022 \, \text{moles} / \text{L} \times 2.4 \times 10^{-24} \, \text{L} = 5.3 \times 10^{-26} \, \text{moles}$$

Molecules in pore:

 $5.3 \times 10^{-26} \text{ moles} \times 6.02 \times 10^{23} \text{ molecules/mole} \approx 0.03 \text{ molecules}$

Most of the time, the pore is "empty"!

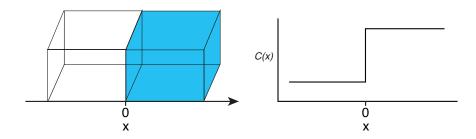
Warning!



Direction Change

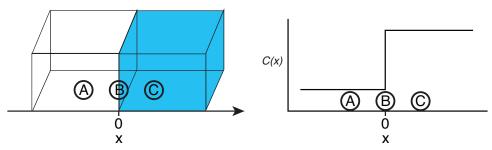
Change in Concentration with Time

An Idealized Macroscopic Diffusion Experiment



■ How will plot of C(x) versus x change with time?

Where will the flux, *J*, be greatest?

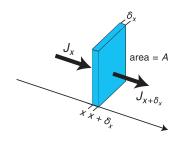


- At point B, where the concentration gradient is greatest.
- But, the molecules move at the same rate everywhere!

- How does the concentration at a given point change with time?
- Net number of molecules moving to the right at two sides of a slice, during interval dt:

$$N_x = J_x A dt$$

 $N_{x+\delta_x} = J_{x+\delta_x} A dt$



Change in number of molecules in the slice:

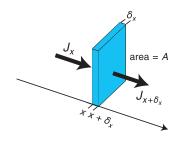
$$dN = N_{x} - N_{x+\delta_{x}}$$

$$= AJ_{x}dt - AJ_{x+\delta_{x}}dt$$

$$= Adt(J_{x} - J_{x+\delta_{x}})$$

Change in concentration in the slice:

$$dC = \frac{dN}{A\delta_{x}} = \frac{Adt(J_{x} - J_{x+\delta_{x}})}{A\delta_{x}}$$
$$= -dt \frac{J_{x+\delta_{x}} - J_{x}}{\delta_{x}}$$



■ In the limit of small dt and small δ_x :

$$\frac{dC}{dt} = -\frac{J_{x+\delta_x} - J_x}{\delta_x} = -\frac{dJ}{dx}$$

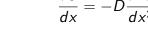
■ How does *J* change with *x*?

Fick's first law:

$$J = -D\frac{dC}{dx}$$

Derivative of J with respect to x:

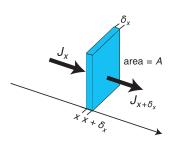
$$\frac{dJ}{dx} = -D\frac{d^2C}{dx^2}$$



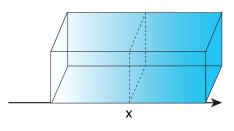
Fick's second law:

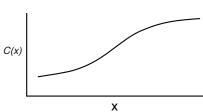
$$\frac{dC}{dt} = -\frac{dJ}{dx} = D\frac{d^2C}{dx^2}$$

Also called the diffusion equation. What is it good for? How do we use it?



Fick's First and Second Laws of Diffusion





First law:

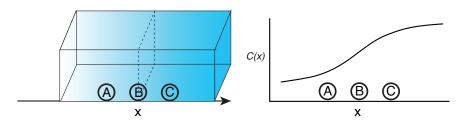
$$J = -D\frac{dC}{dx}$$

Flux, J, at position x is proportional to the concentration gradient at that position. Second law:

$$\frac{dC}{dt} = D\frac{d^2C}{dx^2}$$

Rate of change in concentration at position x is proportional to the derivative of the concentration gradient.

Where will the concentration increase most rapidly?



At point A, where the concentration gradient increases most rapidly with respect to x.

$$\frac{dC}{dt} = D\frac{d^2C}{dx^2}$$

- A "second-order differential equation".
- The solution to the equation is a function:

$$C = f(x, t)$$

that satisfies the equation:

$$\frac{df(x,t)}{dt} = D\frac{d^2f(x,t)}{dx^2}$$

- The trick is to find C = f(x, t).
- The solution depends on the shape of the volume and the initial concentrations, the boundary conditions.

