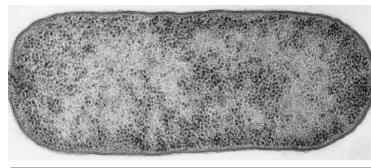


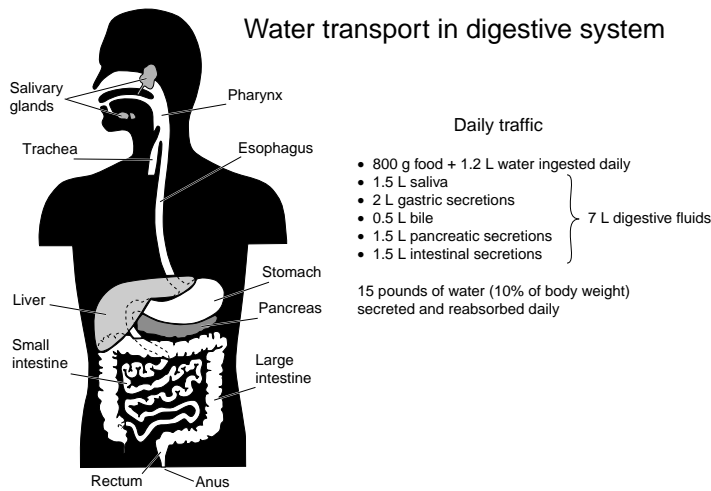
Figure 2.19



2.5 μm

Figure 1.1

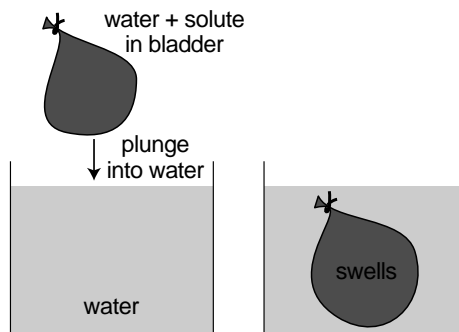
Water transport in digestive system



Osmosis Observations

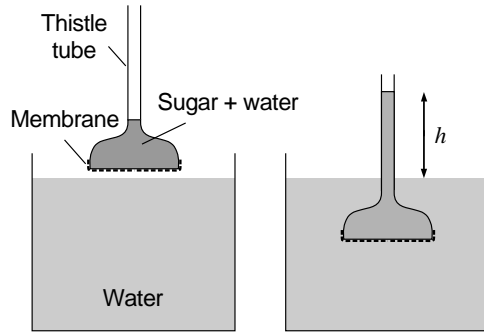
Henri Dutrochet (early 1800s)

- first described phenomenon and called it osmosis
- developed first osmometer: animal bladder filled with test solution, plunge into water, swells, turgid
- pressure greater for solutions with more solute



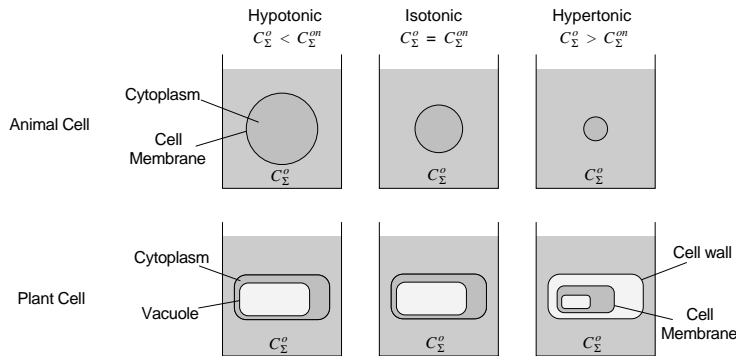
Wilhelm Pfeffer (mid 1800s)

- osmosis can be stopped with hydraulic pressure
- thistle tube + animal bladder (or artificial membrane by late 1800s)
 - water flows in direction to equalize sugar concentration
 - hydraulic pressure develops
 - flow stops when osmotic pressure = hydraulic pressure
- pressure proportional to concentration of solute
- pressure increases slightly with temperature



Hugo de Vries (late 1800s)

- studied osmosis in cells
- animal cell can shrink or swell depending on concentration
- isotonic (same "tension" as in cell's normal environment)
- plasmolysis – plant cell membrane separates from cell wall
- except for salts, plasmolysis occurs at same MOLAR concentration (does not depend on chemical properties of solute)
 - colligative property (freezing point depression, boiling point elevation)
- salts are different: ratios of small integers



Henricus van't Hoff (1886)

- formulated mathematical law
- count number of particles in volume V
- measure temperature T
- osmotic pressure = pressure produced by gas with same number of particles, same volume, and same pressure

van't Hoff's Law

$$\underbrace{\pi(x,t)}_{\text{osmotic pressure}} = \underbrace{R}_{\text{molar gas constant}} \underbrace{T}_{\text{absolute temperature}} \underbrace{\sum_n C_n(x,t)}_{\text{total solute concentration}}$$

8.314 J/(mol·K)

- salts are different

Svante Arrhenius (1884)

- PhD (age 25): dissolution of salts into ions
- $\text{NaCl} \rightarrow \text{Na}^+ + \text{Cl}^-$ (\therefore conducts electricity)
- count ions as separate particles
 \rightarrow van' t Hoff' s law works for salts as well

$$\underbrace{\pi(x,t)}_{\substack{\text{osmotic} \\ \text{pressure}}} = R T \sum_n C_n(x,t) = R T \underbrace{C_\Sigma(x,t)}_{\substack{\text{osmolarity} \\ [\text{osmol}/\text{m}^3]}}$$

[Pa = N/m²]

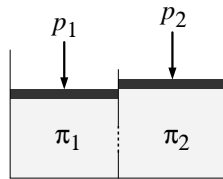
Summary

- Osmosis can be characterized by a pressure that is given by van' t Hoff' s Law:

$$\pi = R T \sum_n C_n = R T C_\Sigma$$

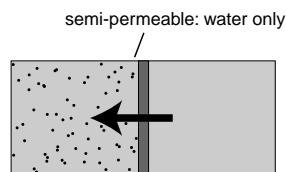
- Osmotic pressures cause water to flow
 ... similar to the flow caused by hydraulic pressure
 ... but in the opposite direction!
- Water flow stops when osmotic pressure difference is equal to the hydraulic pressure difference:

$$p_1 - p_2 = \pi_1 - \pi_2$$

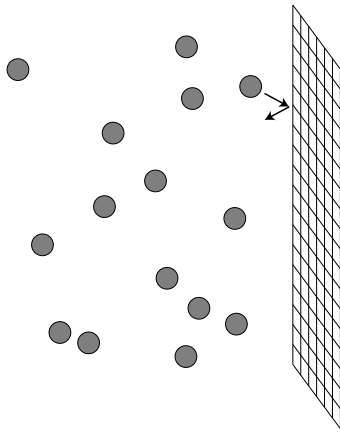


Controversy

- no question that van' t Hoff' s law is true
- but why?



- why should water go TOWARD the solute?
- large osmotic pressure ATTRACTS water!



solute collides with mesh

mesh exerts force on solute
→ changes solute momentum

solute collides with solvent
→ transfers momentum to solvent

change in solvent momentum
is equivalent to a hydraulic pressure

change in hydraulic pressure
= change in osmotic pressure

Macroscopic laws of solvent transport: hydraulic case

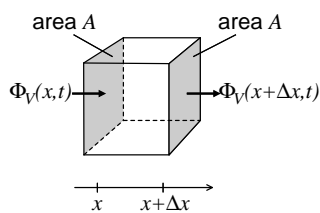
Darcy' s Law: flow through porous medium

$$\underbrace{\Phi_V(x,t)}_{\text{solvent flux}} = - \underbrace{\kappa}_{\text{hydraulic permeability}} \underbrace{\frac{\partial p(x,t)}{\partial x}}_{\text{hydraulic pressure gradient}}$$

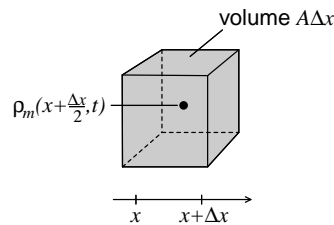
- analogous to Fick' s law for diffusion, Ohm' s law for electrical current, Fourier' s law for heat flow
- units for solvent flux different from units for solute flux

$$\phi_n \left[\frac{\text{mol}}{\text{m}^2 \text{s}} \right] \quad \Phi_V \left[\frac{\text{m}^3}{\text{m}^2 \text{s}} = \frac{\text{m}}{\text{s}} \right] \quad p \left[\text{Pa} = \frac{\text{N}}{\text{m}^2} \right]$$

Continuity Equation for Solvent Flow



Net mass of solvent entering through edges during $(t, t+\Delta t)$



Change in mass of solvent in volume from t to $t+\Delta t$

$$\text{left} \quad \text{right} \quad \text{time } t+\Delta t \quad \text{time } t$$

$$\rho_m(x, t+\frac{\Delta t}{2}) \Phi_V(x, t+\frac{\Delta t}{2}) A \Delta t - \rho_m(x+\Delta x, t+\frac{\Delta t}{2}) \Phi_V(x+\Delta x, t+\frac{\Delta t}{2}) A \Delta t = \rho_m(x+\frac{\Delta x}{2}, t+\Delta t) A \Delta x - \rho_m(x+\frac{\Delta x}{2}, t) A \Delta x$$

equal if solvent is neither created nor destroyed

$$\frac{\rho_m(x, t+\frac{\Delta t}{2}) \Phi_V(x, t+\frac{\Delta t}{2}) - \rho_m(x+\Delta x, t+\frac{\Delta t}{2}) \Phi_V(x+\Delta x, t+\frac{\Delta t}{2})}{\Delta x} = \frac{\rho_m(x+\frac{\Delta x}{2}, t+\Delta t) - \rho_m(x+\frac{\Delta x}{2}, t)}{\Delta t}$$

Take limit as $\Delta x \rightarrow 0$ and $\Delta t \rightarrow 0$

$$-\frac{\partial}{\partial x} [\rho_m(x,t) \Phi_V(x,t)] = \frac{\partial \rho_m(x,t)}{\partial t}$$

Macroscopic laws of water transport (hydraulic case):

• Darcy's law:

$$\Phi_V(x,t) = -\kappa \frac{\partial p(x,t)}{\partial x}$$

• Continuity

$$-\frac{\partial}{\partial x} [\rho_m(x,t)\Phi_V(x,t)] = \frac{\partial \rho_m(x,t)}{\partial t}$$

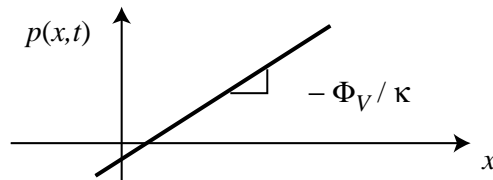
Assume water is incompressible

→ $\rho_m(x,t) = \rho_0 = \text{constant}$ in space and time

$$\rightarrow -\rho_0 \frac{\partial}{\partial x} [\Phi_V(x,t)] = 0$$

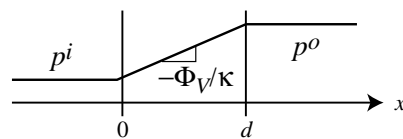
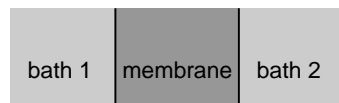
→ $\Phi_V(x,t) = \text{constant}$ in space

→ $p(x,t) = \text{linear function of space}$



very much like SS case for diffusion
but always true here since water is incompressible

Water flow through thin membrane
(hydraulic)



$$\frac{p^o(t) - p^i(t)}{d} = -\frac{\Phi_V(t)}{\kappa}$$

$$\Phi_V(t) = \mathcal{L}_V (p^i(t) - p^o(t)); \quad \mathcal{L}_V = \kappa / d$$

hydraulic conductivity

analogous to Fick's law for membranes

$$\phi_n(t) = P_n (c_n^i(t) - c_n^o(t)); \quad P_n = \frac{D_n k_n}{d}$$