

Last time: Tissue mechanics

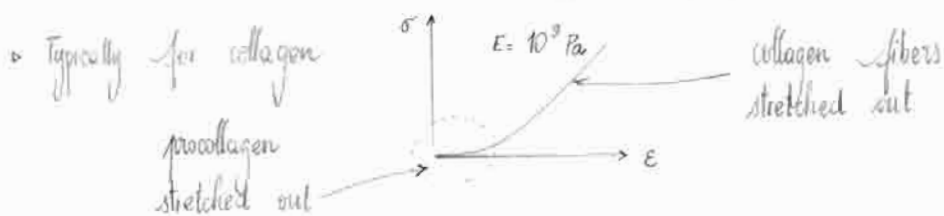
I. Molecular structure & composition of extracellular matrix (ECM)

- behavior of bone, muscle, tendon
- collagen superfamily (how many? fibrillar, fibril forming, sheet forming)
- processing of collagen
- nonlinear behavior (a wavy \gg collagen)

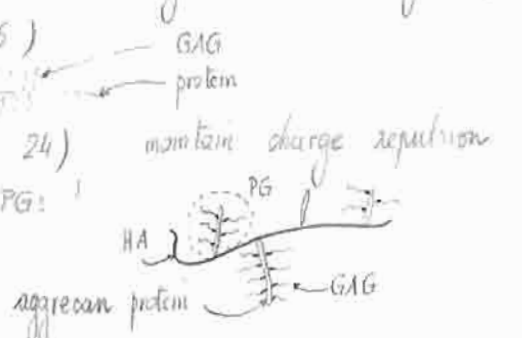
Today: Finish I: elastin, proteoglycans & glycosaminoglycans, adhesion proteins

Equilibrium equations

Energy methods (scaling analysis rather than precise solutions)

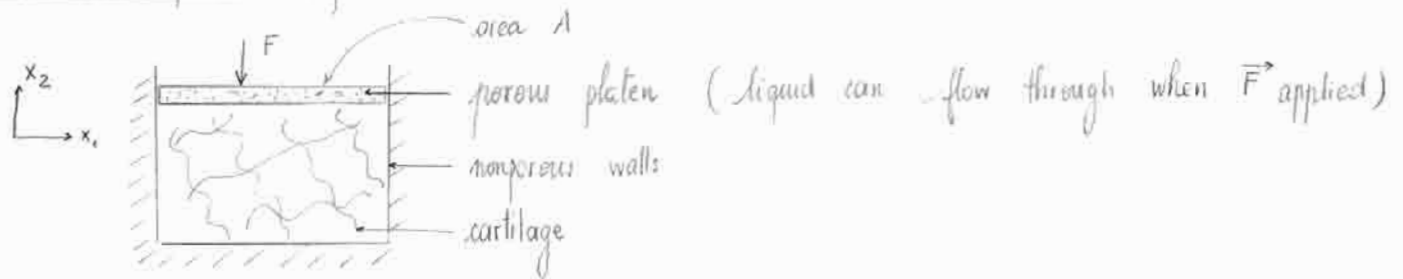


- connections between collagen fibers are proteoglycans or collagen (slide 15)
- elastin more random in character than collagen (straight on short distance, large persistence length)
- entropic elasticity (slide 16, 17)
- $E_{\text{elastin}} = 10^5 \text{ Pa}$ and strains up to 20% (huge but still linear pretty much)
- not much hysteresis (area difference = energy dissipated, not much!)
- elastic rather than viscoelastic response of elastin (slide 18)
- elastin: lung, blood vessels (collagen for resistance at high strains \Rightarrow don't break)
- radius as a function of pressure of arterial wall
- compliant artery first (elastin characteristic), stiffer than (collagen recruited \Rightarrow higher E)
- change stress of living material \Rightarrow remodeling (over time) of cells, they respond to environment
- mechanical properties evolve
- intima thickened by low shear stress (endothelial cells modified \Rightarrow atherosclerosis for instance)
- (slide 20)
- surface tension accounts for difference between air- and liquid-filled lung (slide 21)
- surfactants produced to reduce surface tension \hookrightarrow stiff \hookrightarrow larger volume for given pressure
- aggrecan can be as long as $1 \mu\text{m}$. huge! (slide 26)
- proteoglycan = core protein + GAGs attached to it
- retain proteoglycan structure (by freezing) for TEM (slide 24) maintain charge repulsion
- hyaluronan molecule can be the core of an array of PGs
- aggrecan is very space-filling
- gives rise to ECM compressibility



- highly charged GAGs: repel each other (slide 31)
equilibrium after all liquid is expelled
change modulus by changing solution ionic strength (charge shielding: of \ominus GAG charges by \oplus solution charges)
- 50% of stiffness from charge repulsion
50% " from collagen's own stiffness (slide 33)

o Confined compression experiment



$$\sigma_{22} A = -F \quad \text{or} \quad \sigma_{22} = -\frac{F}{A} = \lambda \epsilon_{kk} \delta_{ij} + 2G \epsilon_{22} = \underbrace{(\lambda + 2G)}_H \epsilon_{22}$$

$$\epsilon_{11} = \epsilon_{33} = 0$$

H confined compression modulus

$$H = \lambda + 2G$$

$$K = \frac{2}{3}G + \lambda = \frac{E}{3(1-2\nu)} \quad \text{bulk modulus}$$

$$\lambda = \frac{\nu E}{(1+\nu)(1-2\nu)} \quad \text{2nd Lamé coefficient}$$

$$G = \frac{E}{2(1+\nu)} \quad \text{shear modulus}$$

for our biological experiments $0.2 < \nu < 0.5$ typically, so "interchange" G and E

- $G \nearrow$ with collagen content
↳ accounts for 50% of stiffness
if you remove PGs, you lose ~ 50% of G
- charge repulsion (high distance d)
steric effects (~ 20 nm apart)
squeezing of water molecules: hydration
Van der Waals attraction



- cone-plate rheometer (rather than plate) = some shear rate (slide 37)
- storage modulus (very low for liquid) changes with ionic concentration (G' high after gelation \rightarrow solid)
as charges are neutralized (pH ok), gel formation is possible (slide 38, 39)
- glycocalyx (buffy coat) important for microcirculation (sticks ~ 1 μ m away from endothelial cells, and prevents red blood cells from approaching the vessel wall)

