

The two phase model for cornea

Solid phase	Extracellular fluid phase	Bath
Collagen Proteins	Negatively charged proteoglycans (PG's)	Water Na^+ Cl^-
	Water	
	Na^+ Cl^-	
		Diffusion both ways

Solid phase : $S = \{c\}$ collagen
 Extracellular fluid phase : $E = \{w, \text{PG}, \text{Na}, \text{Cl}\}$

Apart from mass and momentum balance, we have electroneutrality and incompressibility of all constituents.

The virtual work principle on the medium, in order to find the constitutive relations for the medium, we write

$$\delta \underline{W} = \underbrace{\underline{T} : \delta \underline{E}}_{\text{mechanical contribution}} + \underbrace{\sum_{K, k \in K} \mu_{Kk}^{ec} \delta m^{Kk}}_{\text{electrochemical contribution}}$$

$(\underline{T}, \underline{E})$: any energetically conjugate pair of stress and strain tensors, e.g

$\left(\begin{smallmatrix} \underline{\tau} \\ \underline{F} \end{smallmatrix} \right)$: 1st Piola-Kirchhoff stress tensor
 deformation gradient tensor

$\left(\begin{smallmatrix} \underline{\tau} \\ \underline{E}_G \end{smallmatrix} \right)$: 2nd Piola-Kirchhoff stress tensor
 Green strain tensor

Alternatively the virtual work principle can be written in terms of the mole based potentials g_{kk}^{ec} and moles N_{kk} .

$$\mathcal{J} \underline{W} = \sum_i \tilde{E}_i + \sum_{k, k \in K} g_{kk}^{ec} N_{kk}$$

The electrochemical potentials, μ_{kK}^{ec} , consist of the following

$$g_{kk}^{ec} = \underbrace{\hat{m}_k p_{kk}^\circ}_{\text{enthalpy of formation of new mass through e.g. chemical reaction}} + \underbrace{\hat{u}_k p_{kk}}_{\text{contribution significant for large concentrations of ions}} + \underbrace{RT \ln x_{kk}}_{\text{chemical contribution}} + \underbrace{\sum_k F \phi_k}_{\text{electrical contribution}}$$

At non-physiological ~~0~~ pH, we have for the phases (the extrafibrillar fluid phase)

$$E = \{w, pG, CQ, Na, H, H_2O\},$$

$$E_{\text{ions}} = \{ \text{Na}, \text{Cl}, \text{H}, \text{H}_2\text{O} \}$$

$$E_{mo} = E - \{PG\} \quad \text{no: mobile constituent,}$$

we have diffusion of masses

$$\vec{J}_K = \sum_{k \in K} n^{kk} (\vec{v}_{kk} - \vec{v}_s) \quad \vec{v}_s: \text{velocity of the solid phase}$$

We have transportation (motion) of electrical ~~charges~~ ^{carriers} (i.e. we have electrical current).

The electrical current density

$$\vec{I}_{ek} = F \sum_{k \in K} \tau_k \frac{N_{kk}}{V} \rightarrow v_{kk}$$

Mass balance equation for the fluid phase

$$\frac{1}{\det F} \frac{d_m k_E}{dt} + \operatorname{div} M_{k_E} = 0$$

for incompressibility in the whole medium we can write

$$\text{div } \vec{V}_s + \text{div } \vec{J}_E = 0$$

The electroneutrality condition

$$\text{div } \vec{I}_E = 0$$

$\vec{I}_E$: electric current density in the extrafibrillar phase

Momentum balance

$$\text{div } \underline{\underline{\sigma}} + \rho \vec{b} = \vec{0} \quad \vec{b}: \text{body forces}$$

We have matrix diffusion equation of the form

$$\vec{j} = -\kappa \vec{f}$$

κ : diffusion matrix

$\vec{j}$: diffusion vector

$\vec{f}$: gradient vector

$$\vec{j} = \begin{bmatrix} \vec{j}_{wE} \\ \vec{j}_{NaE} \\ \vec{j}_{ceE} \end{bmatrix}$$

$$\vec{f} = \begin{bmatrix} \vec{\nabla} g_{wE}^{ec} / \hat{v}_w \\ \vec{\nabla} g_{NaE}^{ec} \\ \vec{\nabla} g_{ceE}^{ec} \end{bmatrix}$$

Structure of the constitutive equations

$$\delta \underline{W} = \underline{T} : \delta \underline{E} + \sum_{e \in E} g_{eE} \delta N_{eE}^* + G_L \delta N_{ceE_L} + P_E \delta I_{inc} + \phi_E \delta I_{eE}$$

L : Ligands (extensions of collagen fibrils where ~~co~~ chlorine ions bind)

G_L : affinity potential (chemical affinity)

$\delta \underline{W}$ can be ~~written~~ written in a shifted form

as

$$\delta \underline{W} = \underline{\bar{T}} : \delta \underline{E} + \sum_{l \in E_{mo}} \bar{g}^{ec} \delta N_{eE}^* + G_L \delta N_{ceE_L}$$

where

$$\underline{\bar{T}} = \underline{T} + P_E \det \underline{F} \cdot \underline{E}^{-1} \cdot \underline{F}^{-T}$$

$$\bar{g}_{lE}^{ec} = g_{lE}^{ec} - \hat{v}_L P_E, \quad l \in E_{mo}$$

From the virtual work equation we can write the constitutive equation for the porous medium

$$\underline{\bar{T}} = \frac{\partial \underline{W}}{\partial \underline{E}}, \quad \bar{g}_{lE} = \frac{\partial \underline{W}}{\partial N_{eE}^*}, \quad G_L = \frac{\partial \underline{W}}{\partial N_{ceE_L}}$$

A particular ~~form~~ form of $\underline{W}$ is

$$\underline{W} = \underline{W}_{ch-mech}(\underline{E}, N_E^*) + \underline{W}_{ch}(N_E^*, N_{ceE_L}) + \underline{W}_{cf}(N_{ceE_L})$$

where

$\underline{W}_{ch-mech}$: chemomechanical term (contribution)

$\underline{W}_{ch}$: purely chemical term

$\underline{W}_{cf}$: term related to the binding of chloride ions to the collagen fiber

Another form of chemoelastic energy

$$\underline{W}_{ch-mech}(\underline{E}, N_E^*) = \underline{T}_0 : \underline{E} + \underline{W}_{ch,1}(\underline{E}, N_E^*) + \underline{W}_{ch,2}(N_E^*) (\underline{W}^{ss}(\underline{E}) + \underline{W}^c(\underline{E}))$$

where

$\underline{T}_0$: initial stress. The term $\underline{T}_0: \underline{E}$ refers to the history of the deformation process.

$\underline{W}^{gs}(\underline{E})$: mechanical energy of the ground substance

$\underline{W}^c(\underline{E})$: mechanical energy due to the activated collagen fibers

$W_{ch,1}(\underline{E}, N_E^*) = -p_{ch}(N_E^*)(\det \underline{F} - 1)$:
coupled chemomechanical contribution

$W_{ch,2}(N_E^*)$: chemical amplification term

A linear elastic relation can be derived from an energy potential of the form

$$\underline{W}^{gs} = \frac{1}{2} \lambda (\text{tr } \underline{E}_G)^2 + \mu \underline{E}_G : \underline{E}_G$$

The 2nd Piola-Kirchhoff stress tensor is given via the relation

$$\underline{\underline{\tau}}^{gs} = \frac{\partial \underline{W}^{gs}}{\partial \underline{E}_G} = \frac{1}{2} \lambda (\text{tr } \underline{\underline{C}} - 3) \underline{\underline{I}} + \mu (\underline{\underline{C}} - \underline{\underline{I}})$$

where $\underline{\underline{C}}$ is the right Cauchy-Green deformation tensor, for which the relation

$$\underline{E}_G = \frac{1}{2} (\underline{\underline{C}} - \underline{\underline{I}})$$

holds.

The energy potential may lead to a nonlinear stress-strain relation, via the multiplication with a nonlinear function $f(\underline{\underline{C}})$, i.e. we write

$$\underline{W}^{gs} = \underline{W}_0^{gs} f(\underline{\underline{C}})$$

where $\underline{W}^0$ is the above mentioned relation for $\underline{W}^0$, that led to the linear stress strain relation.

Transversely isotropic material with a single family of fibers (fibers ~~is~~ aligned along one direction)

For isotropic behavior the strain energy density is a function of the 3 invariants of the $\underline{\underline{C}}$ strain tensor, that is

$$\underline{W} = \underline{W}(I_1, I_2, I_3)$$

For transversely isotropic material

$$\underline{W} = \underline{W}(I_1, I_2, I_3, I_4, I_5)$$

is a function of the 3 invariants of $\underline{\underline{C}}$ plus the 2 pseudo invariants I_4, I_5 of $\underline{\underline{C}}$.