A CONTINUOUS MODEL FOR AN ARTERIAL TISSUE, INCORPORATING REMODELING AND VOLUMETRIC GROWTH

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A continuum-mechanics approach for the derivation of a model for the behavior, that is, the growth and remodeling, of an arterial tissue under a mechanical load is presented. This behavior exhibits an interplay between two phenomena: continuum mechanics and biology. The tissue is modeled as a continuous mixture of two components: elastin and collagen. Both components are incompressible, but the tissue as a whole can show volumetric growth due to the creation of collagen. Collagen is a fibrous structure, having a strain-induced preferred orientation. Remodeling of the tissue incorporates degradation of elastin and strain-induced creation and degradation of collagen fibers. Both elastin and collagen are considered to be nonlinear elastic media; elastin as a neo-Hookean material and collagen fibers behaving according to an exponential law. The modeling is based on the classical balance laws of mass and momentum.

1. Introduction

An aneurysm is a localized dilatation or ballooning of blood vessels. The size of an aneurysm was considered to be a critical indicator of the rupture potential and need for medical intervention. However, size is no longer considered to be an accurate parameter as there have been incidents of small aneurysms rupturing and large ones remaining intact. It is now believed that aneurysms rupture when the hemodynamically induced wall stress exceeds the wall strength. This necessitates a mechanical analysis of the biological tissue.

In modeling aneurysms and other cardiovascular pathologies, we will encounter the interplay of two phenomena: continuum mechanics and biology. Whereas traditional engineering materials passively respond to a change in their environment, biological tissues adapt to their environment by changing their configuration and material properties.

In this paper (originally presented at the TRECOP’07 conference on continuum physics) we will not aim at a physiological justification of our tissue model, but when needed we will use partial results from existing models in the literature. In contrast to many of these models, we will base our model strictly on the basic principles of continuum mechanics. Moreover, we refrain from giving an extensive literature review; for this we refer to the forthcoming [Machyshyn 2008]. Here, we only mention [Humphrey and Rajagopal 2002; Kroon and Holzapfel 2007; Baek et al. 2005; 2006].

We will present a continuous model for a tissue based on the basic laws of continuum mechanics coupled with considerations on the biological behavior of arterial vessels. A tissue of an arterial vessel is mainly built up of two components: elastin and collagen. Elastin behaves as an isotropic nonlinear elastic solid medium, and has as a special feature that it degrades (vanishes) during the formation of an aneurysm.

Keywords: arterial tissue, volumetric growth, strain-induced orientation, elastin, collagen.
Collagen has an anisotropic fibrous structure that can take up stresses in a nonlinear elastic way. Collagen remodels in two ways. On one hand collagen fibers can weaken, in that they elongate or disappear. On the other, new collagen can be laid down or passive fibers can become active, strengthening the collagen as a whole. We will propose a continuous mixture model for an arterial tissue in which remodeling is modeled by taking into account degradation of elastin, local changes of mass of collagen and elastin – resulting in volumetric growth of the tissue – and stress- or strain-induced preferred directions for the lay-down of collagen fibers. In order to more basically incorporate the idea of a distributed lay-down of fibers, Muschik et al. [2000] introduced the concept of mesoscopic continuum physics. This is a very elegant concept that also accounts for thermodynamical effects. However, these effects are not considered in this paper.

2. Basic model of a tissue

Our basic model for an arterial tissue is a three-dimensional continuous mixture of two components: elastin and collagen. These two components are both intrinsically incompressible, but the mixture as a whole can show volumetric growth caused by mass production. This is due to the degradation of elastin, modeled as vanishing of elastin particles, and the continuous creation (or lay-down) of new collagen fibers and degradation (removal) of old ones. By these processes, the total amount, or mass, of elastin and collagen continuously changes in time, which can result in growth or shrinkage of the tissue.

Two important state variables for the analysis to come are the volumetric fractions, \( n_e \) and \( n_c \), of elastin and collagen, respectively. They are defined as the relative amount of elastin or collagen in the mixture; they are dimensionless and they sum up to one, so

\[
\begin{align*}
n_e + n_c &= 1. 
\end{align*}
\]  

The elastin is modeled as an isotropic nonlinear elastic solid, and its elastic constitutive behavior is described by an incompressible neo-Hookean model. As elastin can only degrade, the amount of elastin is monotone decreasing once degradation has started.

On the other hand, collagen is an anisotropic fibrous medium. The elastic fibers can only take up stresses in their fiber direction. As constitutive equation for these elastic stresses we will adapt a nonlinear exponential law. Important fiber properties are their orientation (direction) and prestretch. At each moment in time, and thus at each configuration of the tissue, the fiber directions are described by a distribution function for the fiber orientations. This distribution function changes continuously in time, governed by the state of stretch of the tissue. A specific choice for this distribution function (see Section 6), which was first introduced by Baek et al. [2005; 2006], will be given further on.

Initially in an unloaded state, the collagen fibers are crimped, in which state they do not contribute to the strength of the tissue. When the tissue is loaded, the elastin will be stretched, and there will come a state in which the collagen fibers become uncrimped; the tissue stretch in this state in the direction of a collagen fiber is called the recruitment stretch. Here, we will consider the recruitment stretch as a state variable, which in some sense governs the adaptation of the newly created collagen fibers to the state of stretch of the tissue. This adaptation is modeled such that a newly laid-down collagen fiber is always in the same state of preferred stretch, called the attachment stretch. In contrast to the variable recruitment stretch, the attachment stretch is a constant material parameter in our model.
Before degradation of elastin starts, the tissue is in its healthy state. This is a loaded equilibrium state, in which the tissue is stretched and in which the collagen is always stretched to its attachment stretch \( \lambda_a \). The total amount of elastin (no degradation) and collagen do not change, and thus there is no mass production or volumetric growth in this state. However, also in this state there is a continuous turnover (creation and degradation) of collagen, but in such a way that the amount of degraded collagen is always equal to that of newly created collagen, thus keeping the total mass (and volume) of collagen constant.

Finally, for later use, we define the **metabolic equilibrium state** as the state in which the stretch in each collagen fiber is equal to the attachment stretch \( \lambda_a \).

### 3. Basics of volumetric growth for a one-component medium

In this section, we try to explain the peculiar behavior of a medium with internal mass production and volumetric growth by considering, as an example, a simple or single-component medium (not a mixture, as this case will be dealt with in the remainder of this paper). In so doing, we follow the approach of Kuhl et al. [2007].

To introduce volumetric growth due to mass production, we consider a one-component intrinsic incompressible medium having as reference configuration: \( G_r \) with reference position vector \( X \), and as deformed current configuration: \( G = \mathcal{G}(t) \) with current (at time \( t \)) position vector \( x = x(X, t) \). The density \( \rho \) of the medium is constant and uniform. The deformation gradient is \( \mathcal{F}(X, t) = \partial x / \partial X \), and the associated Jacobian is \( J = \det \mathcal{F} = J(X, t) \).

A material partial volume \( b \) with configuration \( g = g(t) \) at the current time \( t \), and reference configuration \( g_r \), is defined as a part of the whole body across the boundary of which no mass flux takes place. However, inside \( b \), mass sources can be active, causing changes of the total mass contained in \( b \).

The volume of \( b \) is

\[
V(t) = \int_{g(t)} \rho \, dv = \int_{g_r} J(X, t) \, dv_r ,
\]

and its mass is

\[
M(t) = \int_{g(t)} \rho \, dv = \rho \int_{g_r} J(X, t) \, dv_r .
\]

Since this mass \( M = M(t) \) is not constant, the balance of mass yields

\[
\frac{d}{dt} M(t) = \frac{d}{dt} \left[ \rho \int_{g_r} J(X, t) \, dv_r \right] = \rho \int_{g_r} J(X, t) \, dv_r = \rho \int_{g(t)} \frac{J}{J} \, dv = \int_{g(t)} \dot{m} \, dv ,
\]

where \( \dot{m} = \dot{m}(X, t) \) is the mass source, the rate of mass production per unit of current volume (in kg/m\(^3\)sec). This leads to the local mass equation

\[
\rho \frac{dJ}{dt} = J \dot{m} .
\]

The balance of momentum for a medium with volumetric growth reads

\[
\frac{d}{dt} \int_{g(t)} \rho v \, dv = \int_{\partial g(t)} \mathbf{t} \, dS + \int_{g(t)} \rho \mathbf{b} \, dv ,
\]
where now, however, the total body force must be split up into a purely mechanical part, $\rho b_m$, (the external body force) and a part due to the growth of mass, according to

$$\rho b = \rho b_m + \dot{m} v,$$

where $v$ is the velocity.

Using successively the equalities

$$\frac{d}{dt} \int_{g(t)} \rho v dv = \frac{d}{dt} \int_{g_r} \rho v J dV = \int_{g_r} \rho (v J + v J) dV = \int_{g(t)} (\rho \dot{v} + \dot{m} v) dv,$$

Cauchy’s stress law

$$t = \mathcal{T} n,$$

and (7), we obtain the local momentum balance

$$\rho \dot{v} + \dot{m} v = \text{div} \mathcal{T} + \rho b_m + \dot{m} v,$$

or

$$\rho \dot{v} = \text{div} \mathcal{T} + \rho b_m,$$

revealing that the effect of mass production in the local momentum balance has disappeared, and that the local momentum equation takes its classical form.

In (9), $t$ is the stress vector, or traction, $\mathcal{T}$ is the stress tensor, and $n$ is the unit outward normal vector on the boundary $\partial g$ of $g$.

### 4. Configurations and deformations

In this section, we consider the four different configurations depicted in Figure 1:

1. The unloaded state $G_r,0$: in this state both the elastin and the collagen are unloaded, but it is assumed that the collagen is not crimped, meaning that $\lambda_{\text{rec},0} = 1$.

2. The healthy state $G_r$: this state is an equilibrium state under a given external load; equilibrium implies here that the collagen stretch is equal to the attachment stretch $\lambda_a$; this healthy state is assumed to be known, and this state is in our further analysis considered as the reference state (note

![Figure 1. Configurations of tissue body $\mathcal{B}$.](image-url)
that this is not an undeformed or stress-free state). At the initial time $t = 0$, the tissue is in its healthy state and then the degradation followed by remodeling and volumetric growth starts.

(3) The intermediate state $\mathcal{G}(\tau)$: the intermediate time $\tau$ ranges from $t = 0$ to the current time $t$, and at each $\tau$ new collagen fibers are laid down.

(4) The current state $G = \mathcal{G}(t)$: this is the final deformed state we wish to determine.

To describe the deformation of the tissue $\mathcal{B}$, we consider an infinitesimal material volume element containing the material point $\mathcal{P}$. The position of $\mathcal{P}$ in $\mathcal{B}_{r,0}$ is given by its position vector $X_0$, and further by $X$ in $\mathcal{B}_r$, by $\xi$ in $\mathcal{G}(\tau)$, and by $x$ in $G$. Here, we consider $\xi$ and $x$ as functions of $X$ and $\tau$, and of $X$ and $t$, respectively,

$$\xi = \xi(X, t) \text{ , } x = x(X, t).$$  \hspace{1cm} (12)

Recalling that the total deformation gradient of the tissue, or the elastin, from $\mathcal{B}_{r,0}$ to $\mathcal{B}$ is $F_{\text{tot}}$, then

$$F_{\text{tot}} = \frac{\partial x}{\partial X_0} = \frac{\partial x}{\partial X} \frac{\partial X}{\partial X_0} = F = F_0,$$

where $\mathcal{F} = \mathcal{F}(X, t)$.

The partial deformation from $\xi$ to $x$ is described by

$$\hat{\mathcal{F}}(X, t, \tau) = \frac{\partial x}{\partial \xi} = \frac{\partial x}{\partial X} \frac{\partial X}{\partial \xi} = \mathcal{F}(X, t) F^{-1}(X, \tau),$$  \hspace{1cm} (14)

or

$$\mathcal{F}(X, t) = \hat{\mathcal{F}}(X, t, \tau) \mathcal{F}(X, \tau).$$  \hspace{1cm} (15)

Consider a collagen fiber created at time $\tau$ in direction $\gamma$ with initial stretch $\lambda_{\alpha}$, having the initial direction vector

$$e_{c}^{(0)}(X, \tau, \gamma) = \cos \gamma v_1(X, \tau) + \sin \gamma v_2(X, \tau),$$  \hspace{1cm} (16)

with $v_{1,2}$ in the principal directions of $\mathcal{F}(X, \tau)$. How the distribution of $e_{c}^{(0)}$ depends on the state of deformation at $\tau$ will be explained further on. The unit vectors $v_1$ and $v_2$ span a surface in which the main stretching takes place; in the third direction there is only shrinking. For instance, for an arterial tissue modeled as a tube under internal pressure and axial stretch, $v_1$ and $v_2$ are in the azimuthal and axial direction, while the radial direction is the third direction in which compression takes place.

Let $e_{c,0}$ be the direction vector representing $e_{c}^{(0)}$ in the unloaded state $\mathcal{B}_{r,0}$, then

$$e_{c,0}(X, \tau, \gamma) = \frac{F_{\text{tot}}^{-1}(X, \tau) e_{c}^{(0)}(X, \tau, \gamma)}{\| F_{\text{tot}}^{-1}(X, \tau) e_{c}^{(0)}(X, \tau, \gamma) \|}.$$  \hspace{1cm} (17)

The elastin stretch $\lambda$ at time $\tau$ in the direction $e_{c}^{(0)}$ is

$$\lambda(X, \tau, \gamma) = \| F_{\text{tot}}(X, \tau) e_{c,0}(X, \tau, \gamma) \| = \frac{1}{\| F_{\text{tot}}^{-1}(X, \tau) e_{c}^{(0)}(X, \tau, \gamma) \|}.$$  \hspace{1cm} (18)

where in the latter step we have used (17).
Since the collagen stretch $\lambda_c$ at the moment it is laid down is equal to $\lambda_a$, we obtain for the recruitment stretch

$$\lambda_{\text{rec}}(X, \tau, \gamma) = \frac{\lambda(X, \tau, \gamma)}{\lambda_a} = \lambda_a \frac{1}{\|\mathcal{F}^{-1}_{\text{tot}}(X, \tau) \ e^{(0)}_c(X, \tau, \gamma)\|}. \quad (19)$$

Hence, $\lambda_{\text{rec}}$ is not a function of the current time $t$, and we will use the above relation to eliminate $\lambda_{\text{rec}}$ from our further calculations.

The collagen stretch $\lambda_c(X, t, \tau, \gamma)$ at the current time $t$ of the fiber laid down at time $\tau$ in the direction $e^{(0)}_c$ is thus

$$\lambda_c(X, t, \tau, \gamma) = \frac{\lambda(X, t, \gamma)}{\lambda_{\text{rec}}(X, \tau, \gamma)} = \lambda_a \frac{\|\mathcal{F}^{-1}_{\text{tot}}(X, \tau) \ e^{(0)}_c(X, \tau, \gamma)\|}{\|\mathcal{F}^{-1}_{\text{tot}}(X, t) \ e^{(0)}_c(X, \tau, \gamma)\|}. \quad (20)$$

Let the initial direction at $\tau$, $e^{(0)}_c(X, \tau, \gamma)$, deform to the current direction vector $e_c$ at time $t$, then

$$e_c(X, t, \tau, \gamma) = \frac{\mathcal{F}(X, t, \tau) \ e^{(0)}_c(X, \tau, \gamma)}{\|\mathcal{F}(X, t, \tau) \ e^{(0)}_c(X, \tau, \gamma)\|}. \quad (21)$$

5. Mass balances

The tissue is here considered as a mixture of two components: elastin and collagen. Let $b$ be a material partial volume of $\mathcal{B}$ containing both elastin and collagen particles. In a material volume there is no mass flux across the boundaries of the volume, but due to the degradation and production of elastin and collagen, the mass of $b$ is not necessarily conserved. Moreover, although both elastin and collagen are modeled as intrinsically incompressible, the volume of $b$ is not conserved: there is a volumetric growth due to the mass production. This volumetric growth is represented by the value of the Jacobian $J = \det \mathcal{F}$, which is greater than one in case of positive volumetric growth, and less than one in case of tissue resorption. Because $J$ is related to the volumetric growth, we replace $J$ by $J_g = J_g(X, t)$. Since no volumetric growth takes place in the healthy phase, $J_0 = \det \mathcal{F}_0 = 1$.

Let $g = g(t)$ be the configuration of $b$ at time $t$, with $g_r$ its reference configuration in $G_r$. The volume of $b$ is given by

$$V(t) = \int_{g(t)} \ dv = \int_{g_r} J_g(X, t) \ dv_r, \quad (22)$$

which is comparable to Section 3.

The total mass of $b$ consists of the mass of elastin, $M_e$, and that of collagen, $M_c$. With $\rho_e$ and $\rho_c$ the constant intrinsic densities of elastin and collagen, respectively, and $n_e$ and $n_c$ their volume fractions, we have for the total mass of $b$

$$M(t) = M_e(t) + M_c(t), \quad (23)$$

where

$$M_e(t) = \int_{g(t)} \rho_e n_e(X, t) \ dv = \rho_e \int_{g_r} n_e(X, t) J_g(X, t) \ dv_r, \quad (24)$$

and

$$M_c(t) = \int_{g(t)} \rho_c n_c(X, t) \ dv = \rho_c \int_{g_r} n_c(X, t) J_g(X, t) \ dv_r. \quad (25)$$
We denote the rates of mass production per unit of current volume for elastin and collagen by \( \dot{m}_e \) and \( \dot{m}_c \) (kg/m\(^3\)sec), respectively. The change of elastin mass is only due to degradation, meaning that \( \dot{m}_e \) is negative, and is given by

\[
\dot{M}_e(t) = \int_{g(r)} \dot{m}_e(X, t) \, dv = \int_{g(r)} \dot{m}_e(X, t) J_g(X, t) \, dv_r .
\] (26)

For collagen, the situation is somewhat more complex: here, we have collagen production at every time \( \tau \) from \( t = 0 \) to the current time \( t \), but meanwhile the collagen produced at \( \tau \) is monotone decreasing during the period from \( \tau \) to \( t \), which is described by the monotone decreasing degradation function of collagen \( q_c = q_c(\tau) \). Here, we have assumed that the degradation function \( q_c \) is uniform over \( \mathcal{B} \).

Let us introduce a new rate of production of volume fraction of collagen \( \nu_c = \nu_c(X, t, \tau) \), such that \( \nu_c(X, t, \tau) \, d\tau \) is the volume fraction of collagen in \( V(t) \) at \( t \) produced from \( \tau \) to \( \tau + d\tau \) in \( V(\tau) \) and still surviving at \( t \). Thus, equating the rate of change of the collagen in \( V(t) \) due to the production in \( V(\tau) \) from \( \tau \) to \( \tau + d\tau \) multiplied by the collagen degradation function from \( \tau \) to \( t \), we obtain

\[
\rho_c \int_{g(r)} \nu_c(X, t, \tau) \, dv \, d\tau = \rho_c \int_{g(r)} \nu_c(X, t, \tau) J_g(X, t) \, dv_r \, d\tau
\]

\[
= \int_{g(r)} \dot{m}_c(X, \tau) \, dv \, \rho_c \, J_g(X, t) \, dq_c(t - \tau)
\]

\[
= \int_{g(r)} \dot{m}_c(X, \tau) J_g(X, t) \, dv_r \, q_c(t - \tau) \, d\tau ,
\] (27)

which, after localization, yields

\[
\nu_c(X, t, \tau) = \frac{\dot{m}_c(X, \tau) J_g(X, t)}{\rho_c J_g(X, t)} \, q_c(t - \tau) .
\] (28)

The latter relation implies that the total collagen mass of \( b \) at the current time \( t \) is given by

\[
M_c(t) = M_{c,r} + \int_0^t \rho_c \int_{g(r)} \nu_c(X, t, \tau) \, dv \, d\tau
\]

\[
= M_{c,r} + \int_0^t \int_{g(r)} \dot{m}_c(X, t) \, q_c(t - \tau) J_g(X, t) \, dv_r \, d\tau ,
\] (29)

where \( M_{c,r} \) is the collagen mass in \( b \) in the healthy state \( G_r \).

Differentiating this relation with respect to \( t \) and using that, by normalization, \( q(0) = 1 \), we find that the change of the collagen mass in \( b \) from \( t = 0 \) to the current time \( t \) is given by

\[
\dot{M}_c(t) = \int_{g(r)} \dot{m}_c(X, t) J_g(X, t) \, dv_r + \int_0^t \int_{g(r)} \dot{m}_c(X, t) \, q_c(t - \tau) J_g(X, t) \, dv_r \, d\tau .
\] (30)

Taking the time derivatives of \( M_c(t) \) and \( M_c(t) \) according to (26) and (30) in terms of the integrals over \( g_r \), equating the results to (25) and (26), respectively, and realizing that these results hold for arbitrary \( b \), we arrive at the local mass balance equations

\[
\dot{\nu}_e \frac{d}{dt} \mathcal{B}_n_e = J_g \dot{m}_e ,
\] (31)
for elastin, and

\[ \rho_c \frac{d}{dt} (J_g n_e) = J_g \dot{m}_e + \int_0^t \dot{m}_e (X, \tau) \dot{q}_c (t - \tau) J_g (X, \tau) \, d\tau , \tag{32} \]

for collagen. Note that \( \frac{d}{dt} \) stands for the material time derivative, so

\[ \frac{d}{dt} f = \frac{\partial f (X, t)}{\partial t} , \tag{33} \]

where \( f \) is arbitrary.

The initial conditions for (31) and (32) are \( n_e (X, 0) = n_{e,r} \) and \( n_c (X, 0) = n_{c,r} \). Dividing (31) by \( \rho_e \) and (32) by \( \rho_c \), adding them together and using the trivial relation

\[ n_e (X, t) + n_c (X, t) = 1 , \tag{34} \]

we obtain the integro-differential equation for \( J_g \):

\[ J_g = \frac{1}{\rho_e} J_g \dot{m}_e + \frac{1}{\rho_c} J_g \dot{m}_c + \frac{1}{\rho_c} \int_0^t \dot{m}_e (X, \tau) \dot{q}_c (t - \tau) J_g (X, \tau) \, d\tau , \tag{35} \]

with the initial condition \( J_g (X, 0) = 1 \).

At this point, we have derived four equations for the four unknowns \( J_g \), \( n_e \), \( n_c \), and \( \nu_c \), successively (35), (31), (32), and (28), under the momentary assumption that the production rates \( \dot{m}_e \) and \( \dot{m}_c \) are given. Inspection, especially with use of (30), shows us that

\[ n_c (X, t) = n_{c,r} + \int_0^t \nu_c (X, t, \tau) \, d\tau . \tag{36} \]

However, in contrast to what we said above, the production rates \( \dot{m}_e \) and \( \dot{m}_c \) are not explicitly given; we need constitutive equations for these quantities. We will not derive these constitutive equations herein, but rather use constitutive equations found in the literature; see specifically [Driessen 2006; Baek et al. 2005; 2006].

The elastin rate we choose here is constant and uniform, that is,

\[ \dot{m}_e (X, t) = - \mu_e , \tag{37} \]

for \( t \) running from \( t = 0 \) to the final time \( t = t_f \); before \( t = 0 \) and after \( t_f \) there is no degradation of elastin. Here, \( \mu_e \) is a given constant material parameter.

The collagen rate we split into two parts as

\[ \dot{m}_c (X, t) = \dot{m}_c^h (X, t) + \dot{m}_c^e (X, t) , \tag{38} \]

where the superindices ‘\( h \)’ and ‘\( e \)’ stand for ‘healthy’ and ‘extra’, respectively.

For the first term we take

\[ \dot{m}_c^h (X, t) = \frac{\rho_c}{Q_c} n_c (X, t) , \quad Q_c = \int_{-\infty}^t q_c (t - \tau) \, d\tau = \int_0^\infty q_c (\tau) \, d\tau . \tag{39} \]

This term is chosen in such a way that the net collagen mass production in the healthy state is zero.

The ‘extra’ production term can be related to either the state of stress or stretch of the collagen. Baek et al. [2005, Eq. (16)] use a stress-induced growth-rate relation, while the same authors discuss [2006]
three possible cases, the last of which is a stretch-induced growth-rate relation. In this study, we opt for the latter, and take for this the specific relation

\[ \dot{m}_e^c(X, t) = K_g(\bar{\lambda}_c(X, t) - \lambda_a), \]  

where \( \bar{\lambda}_c \) is a kind of averaged collagen stretch, averaged over the set of collagen fibers in the current state \( G = G(t) \) and defined as

\[ \bar{\lambda}_c(X, t) - \lambda_a = \frac{1}{\pi n_c(X, t)} \int_0^t v_c(X, t, \tau) \int_{-\pi/2}^{\pi/2} \lambda_c(X, t, \tau, \gamma) \, d\gamma \, d\tau. \]  

We see that the constitutive equation contains the unknown collagen stretch \( \lambda_c \), which is in turn related to the unknown deformation gradient \( \bar{\mathcal{F}} \) according to (20). For the determination of \( \bar{\mathcal{F}} \), we need the momentum equation as derived in Section 3. However, we will first discuss how to model the distribution of the collagen fibers over their different directions.

Finally, we make the following choice for the collagen degradation function \( q_c \):

\[ q_c(t) = e^{-t/T}, \]  

where \( T \) (in sec) is a time constant that is characteristic for the rate of degradation of the collagen. We note that the time scale \( t_f \) for the elastin degradation is much larger than that of remodeling, so \( T \ll t_f \).

For this choice, the coefficient \( Q_c \) becomes

\[ Q_c = \int_0^\infty e^{-\tau/T} \, d\tau = T. \]  

### 6. Distribution of collagen fibers

At each intermediate time \( \tau \) a new generation of collagen fibers is continuously created. These fibers are laid down in a plane spanned by the two unit vectors \( v_1 \) and \( v_2 \), the principal directions of the deformation gradient \( \bar{\mathcal{F}} = \bar{\mathcal{F}}(X, \tau) \) (see Equation (16)), such that for \( \lambda \in \mathbb{R} \),

\[ \bar{\mathcal{F}} v = \lambda v. \]  

The direction of each individual fiber is given by \( \gamma \), the angle between the fiber direction and the \( v_1 \)-axis. The distribution of these directions is assumed to be governed by a kind of double normal distribution \( D(X, \tau, \gamma) \) defined as

\[ D(X, \tau, \gamma) = e^{-1/\sigma} \left( \frac{\exp[2(\gamma - \mu)]}{\sigma} + \frac{\exp[2(\gamma + \mu)]}{\sigma} \right), \]  

where \( \pm \mu = \pm \mu(X, \tau) \) are the mean fiber directions, and \( \sigma = \sigma(X, \tau) \) is the width of the distribution. The value of \( \gamma \) runs from \(-\pi/2\) to \(\pi/2\); \(\gamma = 0\) corresponds with the \(v_1\)-direction, and \(\gamma = \pm\pi/2\) with the \(\pm v_2\)-direction. A plot of a specific distribution function of the type (45) is shown in Figure 2.

Let \( N_c(X, t, \tau, \gamma) \, d\tau \) denote the volume fraction of collagen at time \( t \) that is created at time \( \tau \), from \( \tau \) to \( \tau + d\tau \), having the direction \( \gamma \) and surviving at \( t \) (hence, \( N_c \) is not so much the volume fraction, but
rather the rate of change thereof). Then, clearly,

\[ \nu_c(X, t, \tau) = \int_{-\pi/2}^{\pi/2} N_c(X, t, \tau, \gamma) \, d\gamma. \]

(46)

Since the distribution of \( N_c \) is governed by \( D \), the latter relation implies that

\[ N_c(X, t, \tau, \gamma) = D(X, \tau, \gamma) \nu_c(X, t, \tau), \]

(47)

with

\[ \tilde{D}(X, \tau, \gamma) = \frac{D(X, \tau, \gamma)}{\int_{-\pi/2}^{\pi/2} D(X, \tau, \gamma) \, d\gamma}. \]

(48)

The mean \( \mu \) and width \( \sigma \) are related to the principal stretches of \( \mathcal{F}(X, \tau) \) in the state \( \Phi(\tau) \). The larger principle stretch \( \lambda_1 = \lambda_1(X, \tau) \) corresponds to the first principle direction \( v_1 = v_1(X, \tau) \), while smaller one, \( \lambda_2 \), corresponds to \( v_2 \). Moreover, since we only consider tissues that are principally stretched, we always have \( \lambda_1 > 1 \). Following [Driessen 2006, Sect. 8.2.2], we take

\[ \mu(X, \tau) = \arctan \left( \frac{g_2}{g_1} \right), \quad \sigma(X, \tau) = \frac{k}{g_1/g_2 - 1}, \]

(49)

where the so-called stimulus functions \( g_1 \) and \( g_2 \) are related to their corresponding principle stretches as

\[ g_1(X, \tau) = \lambda_1(X, \tau) - 1, \quad \text{and} \quad g_2 = \begin{cases} \lambda_2(X, \tau) - 1, & \lambda_2 > 1, \\ 0, & \lambda_2 < 1. \end{cases} \]

(50)

The volume fraction due to the distribution \( N_c \), as given in (47), will turn up in a constitutive equation for the stress given in the next section.

7. Momentum equation and stresses

Following Humphrey and Rajagopal [2002, p. 421] and many others in this field, we assume that we may consider the mixture of elastin and collagen as a constrained mixture that is homogenized with regard to the stresses. This implies that we only need to satisfy the overall momentum law for the mixture as a whole, and not the two partial momentum laws for the components separately. As we have shown in
Section 3, this relation keeps its classical form; see (11). In the following, we shall neglect inertia terms (the deformations we consider are extremely slow) and we exclude external body forces, by which the local momentum equation reduces to the simple equilibrium equation
\[ \text{div}\mathcal{T} = 0. \] (51)

We now need a constitutive equation for the stress tensor $\mathcal{T}$. For the constrained mixture we consider here, we may adapt a rule-of-mixture relation for the stress, stating that the partial stresses due to elastin or collagen are proportional to their volumetric fractions $n_e$ or $n_c$, respectively. Accordingly, we assume that the total stress in the tissue in the current configuration at time $t$ is built up of elastin stress and collagen stress according to
\[ \mathcal{T}(\mathbf{X}, t) = - p(\mathbf{X}, t) \mathcal{I} + \mathcal{T}_e(\mathbf{X}, t) + \mathcal{T}_c(\mathbf{X}, t) , \] (52)
where $p$ is a pressure term, needed to account for the incompressibility of elastin and collagen, and $\mathcal{I}$ is the unit tensor. The elastin is modeled as an incompressible isotropic neo-Hookean material with
\[ \mathcal{T}_e(\mathbf{X}, t) = c_e (\mathcal{B}(\mathbf{X}, t) - \mathcal{I}) , \] (53)
where $c_e$ is the elastin shear modulus and $\mathcal{B} = \mathcal{F}\mathcal{F}^T$ is the left Cauchy–Green tensor.

For the collagen stress, we follow van Oijen [2003, Sect. 3.2.4]. The collagen fibers can only take up tensile stress in their fiber direction, and the nonlinear stress-strain behavior of the collagen fibers is captured by an exponential form for the fiber stress. This leads to the formulation
\[ \mathcal{T}_c(\mathbf{X}, t) = \int_0^\tau \int_{-\pi/2}^{\pi/2} \left\{ N_c [\tau_f(\lambda_c) - \mathbf{e}_c \cdot \mathcal{T}_c \mathbf{e}_c] \mathbf{e}_c \otimes \mathbf{e}_c \right\} (\mathbf{X}, t, \tau, \gamma) \, d\gamma \, d\tau , \] (54)
where
\[ \tau_f(\lambda_c) = 2k_1 \lambda_c^2 (\lambda_c^2 - 1) \exp[k_2(\lambda_c^2 - 1)^2] , \] (55)
with $k_1$ and $k_2$ as material constants and $\lambda_c = \lambda_c(\mathbf{X}, t, \tau, \gamma)$ given by (20). For the $\mathcal{T}_e$ in the integral in (54) one must read $\mathcal{T}_e(\mathbf{X}, \tau)$.

In (52)–(54), the contribution of $n_e\mathcal{T}_e$ is split into $\mathcal{T}_e$ and $-n_c\mathcal{T}_c$, with the latter taken up in (54). However, in (54) this is only done in the fiber direction and not in the transverse direction. The reason for this is that the stress perpendicular to the fiber direction is uncoupled from the fiber fraction, which would mean that the transversal properties of the tissue are not affected by the fibers. This is not logical, as then the tissue would become unnaturally weak in the transverse direction leaving only the elastin to contribute to its stiffness. Oijen [2003] compensated for this by taking in the transverse direction the full $\mathcal{T}_e$, and not the partial $n_e\mathcal{T}_e$. Mechanically, this means that he gives the collagen fibers a transverse stiffness that is equal to that of the elastin. The stress perpendicular to the fiber direction is uncoupled from the fiber fraction, meaning that the transversal properties of the tissue are not affected by the fibers.

8. One-dimensional example

We consider as a first example a one-dimensional problem for a slender circular rod loaded by a fixed uniaxial tensile stress. Let $\mathbf{e}_3$ be the axial direction of the rod, then the axial normal stress $T_{33} = S$ is given, while all other stresses are zero, that is, $T_{ij} = 0, (i, j) \neq (3, 3)$. All collagen fibers are in the $\mathbf{e}_3$-direction, so the distribution of the fibers does not play a role here.
In the unloaded state $G_0$ the tissue is unstretched, whereas in the healthy state the rod is loaded and stretched such that the collagen stretch $\lambda_{c, r}$ is equal to its attachment stretch $\lambda_a$. By the (not very relevant) assumption that $\lambda_{\text{rec}} = 1$ in the healthy state, we see that the stretch of the tissue, and the elastin, in the $e_3$-direction is equal to $\lambda_a$. Since there is no volumetric growth in the healthy state, $\det F_0 = 1$ and thus the matrix of $F_0$ is of the form (also accounting for rotational symmetry of the problem)

$$F_0 = \begin{pmatrix} \lambda_a^{-1/2} & 0 & 0 \\ 0 & \lambda_a^{-1/2} & 0 \\ 0 & 0 & \lambda_a \end{pmatrix}. \quad (56)$$

We consider next the current state $G = \mathcal{G}(t)$. This state is here a homogeneous state, so there is no dependence on $X$ in this example. The total deformation gradient is $F_{\text{tot}}(t) = F(t)F_0$, implying that the tissue stretch in the $e_3$-direction is

$$\lambda(t) = \| F_{\text{tot}}(t)e_3 \| = \| F(t)F_0e_3 \| = \| F(t)\lambda_a e_3 \| = \lambda_a F_{33}. \quad (57)$$

Due to the rotational symmetry: $F_{11} = F_{22}$, and thus the matrix of $F(t)$ is

$$F(t) = \begin{pmatrix} F_{11}(t) & 0 & 0 \\ 0 & F_{11}(t) & 0 \\ 0 & 0 & \lambda(t)/\lambda_a \end{pmatrix}. \quad (58)$$

Since $\det F_0 = 1$, the Jacobian in $G$ is equal to

$$J_g(t) = \det F(t) = \frac{\lambda(t)}{\lambda_a} F_{11}^2(t), \quad (59)$$

yielding

$$F_{11}(t) = \sqrt{\frac{\lambda_a}{\lambda(t)}} J_g(t). \quad (60)$$

In the intermediate state $\mathcal{G}(\tau)$ we have

$$\mathcal{F}(\tau) = \mathcal{F}(t)\mathcal{F}^{-1}(\tau) = \begin{pmatrix} F_{11}(t)/F_{11}(\tau) & 0 & 0 \\ 0 & F_{11}(t)/F_{11}(\tau) & 0 \\ 0 & 0 & \lambda(\tau)/\lambda(t) \end{pmatrix}. \quad (61)$$

Moreover, the collagen stretch of a fiber created at time $\tau$ is

$$\lambda_c(\tau) = \lambda_a \frac{\lambda(t)}{\lambda(\tau)}. \quad (62)$$

From (35), we obtain with $\dot{m}_c(t) = -\mu_c$ and $\dot{m}_c(t) = \dot{m}_c^b(t) + \dot{m}_c^e(t)$, where $\dot{m}_c^b(t) = \rho_c n_c(t)/T$, and $\dot{m}_c^e(t) = K_g (\tilde{\lambda}_c(t) - \lambda_a)$. The ordinary differential equation for $J_g(t)$ is

$$\dot{J}_g(t) = -\frac{\mu_c}{\rho_c} J_g(t) + \frac{1}{T} J_g(t)n_c(t) + \frac{K_g}{\rho_c} (\tilde{\lambda}_c(t) - \lambda_a) J_g(t) - \frac{1}{T} J_g(t)n_c(t)$$

$$= \left[ -\frac{\mu_c}{\rho_c} + \frac{K_g}{\rho_c} (\tilde{\lambda}_c(t) - \lambda_a) \right] J_g(t). \quad (63)$$
For this one-dimensional problem, \( \tilde{\lambda}_c(t) \) follows from (41) as

\[
\tilde{\lambda}_c(t) - \lambda_a = \frac{1}{n_c(t)} \int_0^t v_c(t, \tau)\lambda_c(t, \tau) \, d\tau ,
\]  

(64)

where, according to (28)

\[
v_c(t, \tau) = \frac{J_e(\tau)}{\rho_c J_e(t)} \left( \rho_c \frac{n_c(\tau) + K_g(\tilde{\lambda}_c(\tau) - \lambda_a)}{T} \right) e^{-(t-\tau)/T} .
\]  

(65)

Substituting (65) and (62) into (64), we obtain the following integral equation for \( \tilde{\lambda}_c(t) \):

\[
\tilde{\lambda}_c(t) - \lambda_a = \frac{\lambda_a \lambda(t)}{\rho_c J_e(t) n_c(t)} \int_0^t \left[ \frac{\rho_c}{T} n_c(\tau) + K_g(\tilde{\lambda}_c(\tau) - \lambda_a) \right] \frac{J_e(\tau)}{\lambda(\tau)} e^{-(t-\tau)/T} \, d\tau .
\]  

(66)

Further, we find from (36) that

\[
n_c(t) = n_{c,e} + \frac{1}{\rho_c J_e(t)} \int_0^t \left[ \frac{\rho_c}{T} n_c(\tau) + K_g(\tilde{\lambda}_c(\tau) - \lambda_a) \right] J_e(\tau) e^{-(t-\tau)/T} \, d\tau .
\]  

(67)

At this point, we have with (63), (66) and (67) three equations for the four unknowns \( J_e(t) \), \( \tilde{\lambda}_c(t) \), \( n_c(t) \) and \( \lambda(t) \). The missing equation for \( \lambda(t) \) will follow from the equations for the stresses. Since we have here a homogeneous stress situation, the equilibrium equations are trivially satisfied. The pressure \( p \) will follow from the condition that \( T_{11} = T_{22} = 0 \), while the remaining equation \( T_{33} = S \) will yield the equation for \( \lambda(t) \) we are looking for.

The stresses \( T_{11} \) and \( T_{22} \) do not contain a collagen part, and they are given by

\[
T_{11} = T_{22} = -p(t) + c_e(F_{11}^2(t) - 1) = -p(t) + c_e \left( \frac{\lambda_a}{\lambda(t)} \right) J_e(t) - 1).
\]  

(68)

Hence, \( T_{11} = T_{22} \) yields

\[
p(t) = c_e \left( \frac{\lambda_a}{\lambda(t)} \right) J_e(t) - 1).
\]  

(69)

Next, \( T_{33} \) follows from (52)–(54) as

\[
T_{33}(t) = -p(t) + c_e(F_{33}^2(t) - 1) + \int_0^t v_c(t, \tau) \left[ \tau_f(\lambda_c) - T_{e,33} \right](t, \tau) \, d\tau
\]

\[
= c_e \left[ \frac{\lambda^2(t)}{\lambda_a^2} - \frac{\lambda_a}{\lambda(t)} \right] J_e(t) + \int_0^t v_c(t, \tau) \left[ \tau_f(\lambda_c(t, \tau)) - \tau_e(\tau) \right] \, d\tau ,
\]  

(70)

with \( v_c \) as given by (65), \( \tau_f(\lambda_c) \) by (55), and \( \lambda_c \) by (62), while \( \tau_e \) stands for

\[
\tau_e(\tau) = T_{e,33}(t) = c_e \left( \frac{\lambda^2(\tau)}{\lambda_a^2} - 1 \right).
\]  

(71)

The missing equation for \( \lambda(t) \) is the simple one

\[
T_{33}(t) = S.
\]  

(72)

Hence, we now have three equations for our three fundamental unknowns: \( J_e(t) \), \( n_c(t) \) and \( \lambda(t) \); the auxiliary variables \( v_c(t, \tau) \), \( \lambda_c(t, \tau) \) and \( \tilde{\lambda}_c(t) \) are determined by (65), (62), and (66), respectively.
However, even for this most simple example this set of equations is already very complex and cannot be solved analytically. Therefore, this set, which has only one independent variable, the time $t$, must be solved by numerical integration. At this stage, we refrain from doing this; it remains an option for further research.

9. Discussion and perspectives

In this paper, we have constructed a model for remodeling and volumetric growth in an arterial tissue, considered as a constrained mixture of elastin and collagen and based on a continuum-mechanics approach. This is in contrast to several other approaches, for example, [Machyshyn 2008] and [Driessen 2006], who built their models in a more discrete way, both in time and space, directly aiming at a finite element implementation. As far as we could compare the present continuous model with the discrete model developed in [Machyshyn 2008], we found complete correspondence.

We established a complete system for the four essential unknowns in the problem of a loaded tissue: the volumetric fraction of collagen, the Jacobian of the deformation (characteristic for the volumetric growth), the stretch of the collagen fibers, and the stretch of the tissue. The derivation is based on the classical balance laws of mass and momentum. Our model incorporates mass production, volumetric growth, degradation of elastin, strain-induced preferred fiber orientation and collagen creation, isotropic nonlinear (neo-Hookean) elastic behavior of elastin, and anisotropic (fibrous) nonlinear (exponential) elastic behavior of collagen.

In this paper, we gave the general derivation of the continuous model, but only applied it to a simple one-dimensional example. We did not perform explicit numerical calculations. This, together with a treatment of more complex examples, was beyond the scope of this article. In the near future, we hope to apply this model to more complex (tube-like) structures, and to do the necessary numerical calculations, with the ultimate goal of an adequate model for the growth of aneurysms in cerebral blood vessels.

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