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A relaxed growth modeling framework for controlling growth-induced residual stresses

M. Genet^{a,b,*}
 ^aLaboratoire de Mécanique des Solides, École Polytechnique / CNRS / Université
 Paris-Saclay, Palaiseau, France
 ^bM3DISIM team, INRIA / Université Paris-Saclay, Palaiseau, France

7 Abstract

8 (Abstract word count: 245.)

Background. Constitutive models of the mechanical response of soft tissues ٥ have been established and are widely accepted, but models of soft tissues re-10 modeling are more controversial. Specifically for growth, one important ques-11 tion arises pertaining to residual stresses: existing growth models inevitably 12 introduce residual stresses, but it is not entirely clear if this is physiological 13 or merely an artifact of the modeling framework. As a consequence, in sim-14 ulating growth, some authors have chosen to keep growth-induced residual 15 stresses, and others have chosen to remove them. 16

Methods. In this paper, we introduce a novel "relaxed growth" framework 17 allowing for a fine control of the amount of residual stresses generated dur-18 ing tissue growth. It is a direct extension of the classical framework of the 19 multiplicative decomposition of the transformation gradient, to which an ad-20 ditional sub-transformation is introduced in order to let the original unloaded 21 configuration evolve, hence relieving some residual stresses. We provide mul-22 tiple illustrations of the framework mechanical response, on time-driven con-23 strained growth as well as the strain-driven growth problem of the artery 24 under internal pressure, including the opening angle experiment. 25

Findings. The novel relaxed growth modeling framework introduced in this
 paper allows for a better control of growth-induced residual stresses compared
 to standard growth models based on the multiplicative decomposition of the
 transformation gradient.

^{*}martin.genet@polytechnique.edu

Interpretation. Growth-induced residual stresses should be better handled
 in soft tissues biomechanical models, especially in patient-specific models
 of diseased organs that are aimed at augmented diagnosis and treatment

4 optimization.

5 Keywords:

⁶ Finite growth, Residual stresses, Relaxation, Finite element method

 $_{7}$ (Total word count: *ca.* 3000.)

8 1. Introduction

Biomechanical models are widely considered as good candidates to im-9 prove patient diagnosis and treatment of various diseases, for instance in 10 vascular [Taylor and Figueroa, 2009], cardiac [Smith et al., 2011; Krishna-11 murthy et al., 2013, or respiratory [Roth et al., 2017] mechanics. To do so, 12 these models must first be formulated with patient-agnostic constitutive laws 13 and parameters that are developed alongside with experiments. And once 14 these generic models are set up, clinical data must be assimilated into to 15 personalize them and hence produce diagnosis [Xi et al., 2011; Genet et al., 16 2015a] and/or perform in silico treatment optimization [Sermesant et al., 17 2012]. 18

Today, most personalized modeling pipelines focus on the current state 19 of the tissue/organ, and few work focused on predicting the long term evo-20 lution of the system of interest, *i.e.*, prognosis [Clatz et al., 2005; Rausch 21 et al., 2017]. Similarly, constitutive models of the mechanical response of 22 soft tissues have been established and are widely accepted in the community, 23 but models of soft tissues remodeling are more controversial [Witzenburg 24 and Holmes, 2017]. This is explained by the increased complexity and as-25 sociated decreased understanding of the physical mechanisms at play during 26 remodeling. Specifically for growth, a remodeling mechanism associated with 27 the addition and/or removal of matter with unchanged properties [Taber. 28 1995; Kuhl, 2014, there are competing approaches for the very description 29 of growth (multiplicative decomposition of the transformation gradient [Ro-30 driguez et al., 1994; Kuhl, 2014] vs. constrained mixture theory [Humphrey 31 and Rajagopal, 2002; Valentín and Holzapfel, 2012; Cyron et al., 2016), the 32 growth driving force (chemistry vs. mechanics [Maillet et al., 2013; Kuhl, 33 2014], strain vs. stress [Rodriguez et al., 1994; Göktepe et al., 2010; Kerck-34 hoffs et al., 2012], static vs. oscillatory loading [Maillet et al., 2013; Lee et al., 35

¹ 2015], *etc.*). Nevertheless, the next generation of personalized modeling-² based clinical tools might include remodeling mechanisms such as growth for ³ prognosis and treatment optimization.

One important question associated with the modeling of growth is about 4 the induced residual stresses, or equivalently, prestrain. On the one hand 5 they are present in living tissues [Fung, 1993], on the other hand growth nat-6 urally induces residual stresses [Rodriguez et al., 1994; Skalak et al., 1996]; 7 however it is still largely unknown how much of the residual stresses in-8 duced by the physiological or pathological growth remain, and how much are q relaxed away by some relaxation mechanism [Fung, 1993; Taber, 1995]. Ac-10 tually, it was found experimentally, in studies involving hypertension-induced 11 pathological growth of cardiovascular tissues, that opening angle, and hence 12 residual stresses, were only very little correlated with growth [Liu and Fung, 13 1989; Omens et al., 1996]. As a consequence, in growth models of the lit-14 erature, some authors have chosen to keep growth-induced residual stresses 15 [Rodriguez et al., 1994; Lee et al., 2014; Genet et al., 2015b], and others have 16 chosen to remove them [Kroon et al., 2009; Lee et al., 2015, 2016]. Other 17 approaches have been proposed to better deal with growth-induced resid-18 ual stresses, notably a recent growth modeling framework that include some 19 "fluid-like" growth [Böl and Bolea Albero, 2014; Bolea Albero et al., 2014]. 20 In this paper, we introduce a novel "relaxed growth" model, in the gen-21 eral framework of the multiplicative decomposition of the transformation 22 gradient, which allows one to control the amount of growth-induced resid-23 ual stresses. The general idea is to add another sub-transformation in the 24 decomposition, which relaxes the tissue by transforming its unloaded config-25 uration. Illustrations are provided on time-driven constrained growth as well 26

²⁷ as the strain-driven growth problem of the artery under internal pressure.

28 2. Methods

29 2.1. Review of classical growth modeling

30 2.1.1. Kinematics

In this part, we review the standard formulation of finite growth modeling based on the multiplicative decomposition of the transformation gradient, originally proposed by [Rodriguez et al., 1994], and illustrated on Figure 1. Thus, let us consider a physical body \mathcal{B} that initially occupies a domain Ω_0 and that, after deformation and growth, occupies a domain Ω . The geometrical transformation between material neighborhoods of \mathcal{B} in Ω_0 and



Figure 1: Schematic of the multiplicative decomposition of the transformation gradient into growth, prestrain and loading parts, revealing the elastic part of the transformation [Rodriguez et al., 1994]. Because the growth part represents the transformation of material neighborhoods independently from each others and hence is incompatible (*i.e.*, does not derive from a continuous mapping), the grown configuration is incompatible (*i.e.*, discontinuous everywhere). Consequently, the prestrain and elastic parts of the transformation are also incompatible. Conversely, both the full transformation and the loading part of the transformation are compatible, though if the loading part does represent a continuous mapping between material points, the full transformation only represents a continuous mapping between material neighborhoods, in which mass has potentially been added or removed. This decomposition is generally represented without initial prestrain, *i.e.*, with $\underline{F_0} = \underline{1}$, in which case $\underline{F_{g0}} = \underline{1}$ and $\underline{F_{p0}} = \underline{F^p}$, such that $\underline{F^e} = \underline{F'}$.

¹ Ω is denoted $\underline{\Phi}$. Following the classical scheme of Figure 1, the transfor-² mation gradient $\underline{\underline{F}} := \underline{\underline{Grad}}(\underline{\Phi})$ is multiplicatively decomposed into growth, ³ prestrain, and load-induced parts:

$$\underline{\underline{F}} = \underline{\underline{F}^{l}} \cdot \underline{\underline{F}^{p}} \cdot \underline{\underline{F}^{g}},\tag{1}$$

⁴ where, in general, both $\underline{F^g}$ (the internal variable describing tissue growth) ⁵ and $\underline{F^p}$ (the prestrain) are incompatible second order tensor fields, and ⁶ $\underline{F^l} := \underline{Grad} (\underline{\Phi^l})$ is the gradient of the (compatible) transformation induced ⁷ by mechanical loading. The prestrain and load-induced parts can be multi-⁸ plicatively combined to form the elastic part of the transformation:

$$\underline{\underline{F}'} := \underline{\underline{F}^l} \cdot \underline{\underline{F}^p},\tag{2}$$

⁹ such that the full transformation gradient can be multiplicatively decomposed
 ¹⁰ into growth and elastic parts:

$$\underline{\underline{F}} = \underline{\underline{F'}} \cdot \underline{\underline{F^g}}.$$
(3)

¹¹ Equivalently, the elastic part of the transformation can be expressed from ¹² the full transformation gradient and the growth internal variable:

$$\underline{\underline{F}'} = \underline{\underline{F}} \cdot \underline{\underline{F}}^{g-1}.$$
(4)

¹³ The full right Cauchy-Green dilatation tensor is denoted $\underline{\underline{C}} := {}^{t}\underline{\underline{F}} \cdot \underline{\underline{F}}$. As ¹⁴ for the transformation gradient, the elastic part of the dilatation tensor can ¹⁵ be expressed from the full dilatation tensor and the growth internal variable: ¹⁶

$$\underline{\underline{C'}} := {}^{t}\underline{\underline{F'}} \cdot \underline{\underline{F'}} = {}^{t}\underline{\underline{F}}{}^{g-1} \cdot \underline{\underline{C}} \cdot \underline{\underline{F}}{}^{g-1}.$$
(5)

17 2.1.2. Free energy and Stresses

¹⁸ The elastic response is governed by the strain energy potential, W^e . Any ¹⁹ hyperelastic potential can be used. The main modeling assumption is that ²⁰ in fine the free energy $\rho_0 \Psi$ is not a function of the total transformation but ²¹ only its elastic part <u> F^e </u> [Rodriguez et al., 1994; Göktepe et al., 2010]:

$$\rho_0 \Psi\left(\underline{\underline{C}}, \underline{\underline{F}^g}\right) = W^e\left(\underline{\underline{C}^e} := {}^t\underline{\underline{F}^e} \cdot \underline{\underline{F}^e}\right).$$
(6)

²² Consequently, the second Piola-Kirchhoff stress tensor can be expressed as

$$\underline{\underline{\Sigma}} = \frac{\partial \rho_0 \Psi}{\partial \underline{\underline{E}}} = 2 \frac{\partial \rho_0 \Psi}{\partial \underline{\underline{C}}} = 2 \frac{\partial W^e}{\partial \underline{\underline{C}^e}} : \frac{\partial \underline{\underline{C}^e}}{\partial \underline{\underline{C}}}.$$
(7)

¹ In the canonical case described so far, we have

$$\underline{\underline{F}^e} = \underline{\underline{F}'} = \underline{\underline{F}} \cdot \underline{\underline{F}^{g-1}},\tag{8}$$

² such that

$$\underline{\underline{C}^{e}} = \underline{\underline{C}'} = {}^{t}\underline{\underline{F}^{g-1}} \cdot \underline{\underline{C}} \cdot \underline{\underline{F}^{g-1}}, \qquad (9)$$

3 and

$$\underline{\underline{\Sigma}} = \underline{\underline{F}}^{g-1} \cdot 2 \frac{\partial W^e}{\partial \underline{\underline{C}}^e} \cdot {}^t \underline{\underline{F}}^{g-1}.$$
 (10)

4 2.1.3. Growth evolution law

An evolution law must be formulated for the growth tensor to close the system. Several types have been proposed, including simple rates [Kuhl, 2014], strain- [Göktepe et al., 2010; Kerckhoffs et al., 2012; Genet et al., 2016] and stress- [Göktepe et al., 2010] driven laws. See for instance [Kuhl, 2014] for a recent review, and [Witzenburg and Holmes, 2017] for a detailed comparison of existing laws.

In case of tissues subjected to cyclic loading, it is necessary to uncouple 11 the temporal scales of loading and remodeling, by alternating between load-12 ing (*i.e.*, fast time scale) and growth (*i.e.*, slow time scale) steps [Kerckhoffs 13 et al., 2012; Lee et al., 2016]. Thus, for a given growth step, the driving 14 force for growth must be computed from the previous loading step. Aver-15 age [Lee et al., 2015] and maximum [Kerckhoffs et al., 2012] values of stress 16 and strain over a cycle have been proposed, in agreement with experimental 17 analysis [Holmes, 2004]. 18

¹⁹ 2.1.4. Case of initially prestrained material

For the sake of completeness, let us consider the case where the initial configuration Ω_0 has some potentially incompatible initial prestrain, denoted $\underline{F_0}$, or, equivalently, some autobalanced initial prestress $\underline{\sigma_0} = \frac{1}{J_0} \underline{\underline{F_0}} \cdot \underline{\underline{\Sigma_0}} \cdot \underline{\underline{F_0}}^{-1}$ with $J_0 := \det(\underline{F_0})$ and $\underline{\underline{\Sigma_0}} = 2\frac{\partial\rho_0 W^e}{\partial\underline{\underline{C^e}}}(\underline{\underline{C^e}} = \underline{\underline{C_0}})$ where $\underline{\underline{C_0}} := t\underline{\underline{F_0}} \cdot \underline{\underline{F_0}}$ (See Figure 1). Note that here, the prestrain must be seen as a local tensor state variable more than a gradient of some mapping. And in this case, the full elastic transformation, $\underline{\underline{F^e}}$, contains both $\underline{\underline{F'}}$ and the prestrain $\underline{\underline{F_0}}$:

$$\underline{\underline{F}^{e}} = \underline{\underline{F}} \cdot \underline{\underline{F}_{0}} \cdot \underline{\underline{F}^{g^{-1}}},\tag{11}$$

¹ such that the free energy is

$$\rho_0 \Psi\left(\underline{\underline{C}}, \underline{\underline{F}}^g, \underline{\underline{F}}_0\right) = W^e\left(\underline{\underline{C}}^e = {}^t\underline{\underline{F}}^{g-1} \cdot {}^t\underline{\underline{F}}_0 \cdot \underline{\underline{C}} \cdot \underline{\underline{F}}_0 \cdot \underline{\underline{F}}^{g-1}\right), \quad (12)$$

² and the second Piola-Kirchhoff stress tensor is

$$\underline{\underline{\Sigma}} = \underline{\underline{F}_0} \cdot \underline{\underline{F}}^{g-1} \cdot 2 \frac{\partial W^e}{\partial \underline{\underline{C}}^e} \cdot {}^t \underline{\underline{F}}^{g-1} \cdot {}^t \underline{\underline{F}}_0.$$
(13)

 $_3$ It is also interesting to express the prestrain in the grown configuration $_4$ Ω_g :

$$\underline{\underline{F}_{g0}} := \underline{\underline{F}^g} \cdot \underline{\underline{F}_0} \cdot \underline{\underline{F}^{g-1}}, \tag{14}$$

⁵ such that the full elastic transformation can also be expressed as

$$\underline{\underline{F}^e} = \underline{\underline{F}'} \cdot \underline{\underline{F}_{g0}}.$$
(15)

⁶ Similarly, the total prestrain in the new grown unloaded configuration Ω_p is

$$\underline{\underline{F}_{p0}} := \underline{\underline{F}^{p}} \cdot \underline{\underline{F}_{g0}} = \underline{\underline{F}^{p}} \cdot \underline{\underline{F}^{g}} \cdot \underline{\underline{F}^{g}} \cdot \underline{\underline{F}^{g-1}}, \qquad (16)$$

7 such that the full elastic transformation can also be expressed as

$$\underline{\underline{F}^{e}} = \underline{\underline{F}^{l}} \cdot \underline{\underline{F}_{p0}}.$$
(17)

- ⁸ 2.2. Relaxed growth modeling
- 9 2.2.1. Kinematics

We now describe a new formulation for finite growth, involving an additional internal sub-transformation for local relaxation. The associated schematic is represented on the Figure 2. The total transformation gradient is now decomposed into growth, relaxation, prestrain and loading parts:

$$\underline{\underline{F}} = \underline{\underline{F}}^{l} \cdot \underline{\underline{F}}^{p} \cdot \underline{\underline{F}}^{r} \cdot \underline{\underline{F}}^{g}, \tag{18}$$

where, again, both $\underline{F^g}$ (the internal variable describing tissue growth), $\underline{F^r}$ (the internal variable describing tissue relaxation) and $\underline{F^p}$ (the prestrain) are incompatible second order tensor fields, and $\underline{F^l} := \underline{Grad} (\underline{\Phi^l})$ is the gradient of the (compatible) transformation induced by mechanical loading. The elastic part of the transformation is still

$$\underline{\underline{F'}} := \underline{\underline{F}}^l \cdot \underline{\underline{F}}^p, \tag{19}$$



Figure 2: Schematic of the proposed multiplicative decomposition of the transformation gradient into growth, relaxation, prestrain and loading parts.

¹ such that

$$\underline{\underline{F}} = \underline{\underline{F}'} \cdot \underline{\underline{F}^r} \cdot \underline{\underline{F}^g}.$$
(20)

² Thus, here the elastic part of the transformation can be expressed from the
³ full transformation gradient, and both the growth and relaxation internal
⁴ variables:

$$\underline{\underline{F'}} := \underline{\underline{F}} \cdot \underline{\underline{F}}^{g-1} \cdot \underline{\underline{F}}^{r-1}.$$
(21)

⁵ Let us point out that the order of the decomposition is arbitrary.

⁶ The role of the relaxation is to update the unloaded configuration of ⁷ the grown body in order to control the amount of growth-induced residual ⁸ stresses. Thus, the updated unloaded configuration is now Ω_p , with prestrain ⁹

$$\underline{\underline{F}_{p0}} := \underline{\underline{F}^{p}} \cdot \underline{\underline{F}_{r0}}, \tag{22}$$

10 where

$$\underline{\underline{F_{r0}}} := \underline{\underline{F^r}} \cdot \underline{\underline{F_{g0}}} \cdot \underline{\underline{F^{r-1}}}, \qquad (23)$$

11 and

$$\underline{\underline{F}_{g0}} := \underline{\underline{F}^g} \cdot \underline{\underline{F}_0} \cdot \underline{\underline{F}^{g-1}}.$$
(24)

¹² 2.2.2. Free energy and Stresses

As for simple growth, we assume that the free energy $\rho_0 \Psi$ is only a function of the elastic part of the transformation $\underline{\underline{F}^e}$, see Equation (6). However, in the case of relaxed growth, we have

$$\underline{\underline{F}^{e}} = \underline{\underline{F}'} \cdot \underline{\underline{F}_{r0}} = \underline{\underline{F}} \cdot \underline{\underline{F}_{0}} \cdot \underline{\underline{F}^{g-1}} \cdot \underline{\underline{F}^{r-1}}, \qquad (25)$$

16 such that

$$\underline{\underline{C}}^{e} = {}^{t}\underline{\underline{F}}^{r-1} \cdot {}^{t}\underline{\underline{F}}^{g-1} \cdot {}^{t}\underline{\underline{F}}_{\underline{0}} \cdot \underline{\underline{C}} \cdot \underline{\underline{F}}_{\underline{0}} \cdot \underline{\underline{F}}^{g-1} \cdot \underline{\underline{F}}^{r-1}, \qquad (26)$$

17 and

$$\underline{\underline{\Sigma}} = \underline{\underline{F}}_{\underline{0}} \cdot \underline{\underline{F}}^{g-1} \cdot \underline{\underline{F}}^{r-1} \cdot 2 \frac{\partial W^e}{\partial \underline{\underline{C}}^e} \cdot {}^t \underline{\underline{F}}^{r-1} \cdot {}^t \underline{\underline{F}}^{g-1} \cdot {}^t \underline{\underline{F}}_{\underline{0}}.$$
(27)

¹⁸ Similarly, for the unloaded configuration the total elastic deformation is sim-¹⁹ ply

$$\underline{\underline{F}_{p0}} = \underline{\underline{F}^{u}} \cdot \underline{\underline{F}_{0}} \cdot \underline{\underline{F}^{g-1}} \cdot \underline{\underline{F}^{r-1}}, \qquad (28)$$

 $_{20}$ such that

$$\underline{\underline{C_{p0}}} := {}^{t}\underline{\underline{F_{p0}}} \cdot \underline{\underline{F_{p0}}} = {}^{t}\underline{\underline{F^{r-1}}} \cdot {}^{t}\underline{\underline{F^{g-1}}} \cdot {}^{t}\underline{\underline{F_{0}}} \cdot \underline{\underline{C^{u}}} \cdot \underline{\underline{F_{0}}} \cdot \underline{\underline{F^{g-1}}} \cdot \underline{\underline{F^{r-1}}}, \quad (29)$$

1 and

$$\underline{\underline{\Sigma}^{u}} = \underline{\underline{F}_{0}} \cdot \underline{\underline{F}^{g-1}} \cdot \underline{\underline{F}^{r-1}} \cdot 2 \frac{\partial W^{e}}{\partial \underline{\underline{C}_{p0}}} \cdot {}^{t} \underline{\underline{F}^{r-1}} \cdot {}^{t} \underline{\underline{F}^{g-1}} \cdot {}^{t} \underline{\underline{F}_{0}}.$$
(30)

² 2.2.3. Relaxation evolution law

As for the growth evolution, many choices can be made for the evolution of the relaxation internal variable. Since the role of the relaxation is to regulate the prestrain, in this article we will consider the simplest possible evolution law, directly based on the current prestrain level:

$$\underline{\underline{\dot{F}^r}} = \frac{1}{\tau^r} \cdot \underline{\underline{E_{p0}}},\tag{31}$$

⁷ where τ^r is a characteristic time for relaxation, and $\underline{\underline{E}_{p0}} := \frac{1}{2} \left(\underline{\underline{C}_{p0}} - \underline{\underline{1}} \right)$ is ⁸ the Green-Lagrange prestrain. The underlying hypothesis is that the tissue is ⁹ somehow able to sense the current level of prestrain. This is analogous to the ¹⁰ hypothesis made in constrained mixture theory-based growth models, where ¹¹ newly deposited matter has its own reference state that can be the one of ¹² the surrounding matter, the current state of deformation, or a combination ¹³ of both [Valentín et al., 2013; Cyron et al., 2016].

14 2.3. Numerical resolution

The relaxed growth problem can be solved like any nonlinear problem 15 with internal variables in mechanics. The only subtlety here is induced by 16 the choice of the relaxation evolution law, which requires the computation 17 of the new unloaded configuration together with the loaded configuration. 18 We propose a mixed formulation of the problem, the unknowns being U (the 19 total displacement), $\underline{U^u}$ (the displacement of the unloaded configuration), 20 <u> F^{g} </u> (the growth tensor) and <u> F^{r} </u> (the relaxation tensor). After quasi-static 21 assumption, and implicit, mid-point rule temporal discretization, the mixed 22

¹ variational formulation of the problem to solve at each time step is:

$$\int_{\Omega_0} \underline{\underline{\Sigma}} : d_{\underline{U};\underline{U^*}} \underline{\underline{E}} \ d\Omega_0 = \mathcal{W}_e \left(\underline{U};\underline{U^*}\right) \quad \forall \underline{U^*}$$
(32a)

$$\int_{\Omega_0} \underline{\underline{\Sigma}^u} : d_{\underline{U^u};\underline{U^{u*}}} \underline{\underline{E}^u} \ d\Omega_0 = 0 \quad \forall \underline{\underline{U^{u*}}}$$
(32b)

$$\int_{\Omega_0} \left(\underline{\underline{F}}^g - \left(\underline{\underline{F}}^{g,\text{old}} + \underline{\underline{\dot{F}}^{g,\text{mid}}} \Delta t \right) \right) : \underline{\underline{F}}^{g*} d\Omega_0 = 0 \quad \forall \underline{\underline{F}}^{g*} \qquad (32c)$$

$$\int_{\Omega_0} \left(\underline{\underline{F}^r} - \left(\underline{\underline{F}^{r,\text{old}}} + \underline{\underline{\dot{F}}^{r,\text{mid}}} \Delta t \right) \right) : \underline{\underline{F}^{r*}} \, d\Omega_0 = 0 \quad \forall \underline{\underline{F}^{r*}} \tag{32d}$$

where $d_{\underline{U};\underline{U^*}}\underline{\underline{E}}$ and $d_{\underline{U^u};\underline{U^{u*}}}\underline{\underline{E^u}}$ denote the first variations of the total and unloaded Green-Lagrange strain tensors with respect to the total and unloaded displacements, and \mathcal{W}_e denotes the virtual work (semi-linear form) associated to external loads. $\underline{F^{g,\text{mid}}}$ will be specified for each growth evolution law, and relaxation evolution law (31) leads to:

$$\underline{\dot{F}^{r,\text{mid}}}_{\underline{m}} = \frac{1}{\tau^r} \underline{E_{p0}^{\text{mid}}}$$
(33)

7 with $\underline{E_{p0}^{\text{mid}}} = \frac{1}{2} \left(\underline{E_{p0}^{\text{old}}} + \underline{E_{p0}} \right).$

In case of incompressible elastic deformation, two new unknowns are introduced: p (the Lagrange multiplier associated to the incompressibility constraint of the elastic strain, equal to the hydrostatic pressure within the tissue) and p^u (the Lagrange multiplier/hydrostatic pressure in the unloaded configuration), and the full mixed variational formulation becomes:

$$\left(\int_{\Omega_0} \left(\underline{\underline{\Sigma}} - pJ^e \underline{\underline{C}}^{e-1}\right) : d_{\underline{U};\underline{U}^*}\underline{\underline{E}} \ d\Omega_0 = \mathcal{W}_e\left(\underline{U};\underline{U}^*\right) \quad \forall \underline{U}^* \qquad (34a)$$

$$\int_{\Omega_0} \left(J^e - 1 \right) p^* \, d\Omega_0 = 0 \quad \forall p^* \tag{34b}$$

$$\int_{\Omega_0} \left(\underline{\underline{\Sigma}^u} - p^u J_{p0} \underline{\underline{C}_{p0}}^{-1} \right) : d_{\underline{U^u};\underline{U^{u*}}} \underline{\underline{E^u}} \ d\Omega_0 = 0 \quad \forall \underline{U^{u*}}$$
(34c)

$$\int_{\Omega_0} (J_{p0} - 1) p^{u*} d\Omega_0 = 0 \quad \forall p^{u*}$$
(34d)

$$\int_{\Omega_0} \left(\underline{\underline{F}^g} - \left(\underline{\underline{F}^{g,\text{old}}} + \underline{\underline{\dot{F}^{g,\text{mid}}}} \Delta t \right) \right) : \underline{\underline{F}^{g*}} \, d\Omega_0 = 0 \quad \forall \underline{\underline{F}^{g*}} \tag{34e}$$

$$\left(\int_{\Omega_0} \left(\underline{\underline{F}^r} - \left(\underline{\underline{F}^{r,\text{old}}} + \underline{\underline{\dot{F}}^{r,\text{mid}}}\Delta t\right)\right) : \underline{\underline{F}^{r*}} \, d\Omega_0 = 0 \quad \forall \underline{\underline{F}^{r*}} \tag{34f}$$

Problems (32) and (34) are spatially discretized using the standard finite element method. Second order elements are used for displacement unknowns (both $\underline{U} \& \underline{U^u}$), and first order elements for pressure unknowns (both p& p^u), preventing numerical locking in the incompressible limit [Hughes, 2000; Chapelle and Bathe, 2010]. First order elements are used for internal variables (both $\underline{F^g} \& \underline{F^r}$) as well.

The full scheme has been implemented in python, based on the FEniCS
library [Logg et al., 2012; Alnæs et al., 2015], and is freely available¹.

⁹ 3. Results

We will now present multiple illustrations of the relaxed growth responsein 2D.

¹² 3.1. Model response: constrained growth

Let us first illustrate the constitutive behavior described by the relaxed growth model. To do so, we consider a single material point for which the total deformation is blocked to zero, and subjected to time-driven isotropic growth:

$$\begin{cases} \underline{F^g} = (1 + \theta^g) \underline{1} \\ \overline{\theta^g} (t = 0) = 0 \\ \dot{\theta}^g = \frac{1}{\tau^g} \end{cases}$$
(35)

where τ^{g} is a characteristic time for growth. This growth evolution law leads, after temporal discretization, to the following expression:

$$\underline{\underline{\dot{F}}^{g,\text{mid}}}_{=} = \frac{1}{\tau^g} \underline{1}$$
(36)

For the sake of simplicity, and to focus on the relaxed growth framework introduced in this paper, we consider a simple compressible neo-hookean strain energy potential under the plane strain assumption [Ciarlet and Geymonat, 1982]:

$$W^{e}\left(\underline{\underline{C}^{e}}\right) = \frac{\lambda}{4} \left(J^{e^{2}} - 1 - 2\ln J^{e}\right) + \frac{\mu}{2} \left(\operatorname{tr}\left(\underline{\underline{C}^{e}}\right) - 2 - 2\ln J^{e}\right), \qquad (37)$$

¹https://gitlab.inria.fr/mgenet/dolfin_cm

¹ where $\lambda \& \mu$ are the bulk and shear modulus, taken as unity. Compressible ² mixed variational formulation (32) is used, with evolution laws (33) & (36).

This is the purest example of constrained growth. In case of simple 3 growth, this would lead to the development of compressive residual stresses, 4 which arise from constraining the grown tissue to its original volume. This 5 is also what happens in the relaxed growth framework; however, here the 6 amount of growth-induced stresses is controlled by the ratio of the characteristic time of relaxation to the characteristic time of growth. If relaxation 8 is much slower than growth, the response is similar to simple growth; conq versely, if relaxation is much faster than growth, then constrained growth 10 happens with almost no induced residual stress. This is well illustrated on 11 the Figure 3, which shows the normalized (with respect to shear modulus) 12 hydrostatic pressure within the material point as a function of normalized 13 (with respect to growth time constant) time, for various ratios of relaxation 14 over growth characteristic times. The relaxed growth framework allows to 15 control the amount of growth-induced residual stresses in constrained growth. 16

¹⁷ 3.2. A time-driven relaxed growth example: constrained vs. unconstrained ¹⁸ growth

Let us now illustrate the relaxed growth response in a first structural 19 case, inspired from [Kuhl, 2014]. Here we consider a simple square geometry, 20 with three fixed boundaries (left, bottom and right) and one free boundary 21 (top), subjected to the same time-driven growth (35). This case contains 22 both constrained (toward the bottom, where the total strain is restricted, 23 thus developing compressive residual stresses), and unconstrained (toward 24 the top, where the free edge limits the stress level, and large strains will 25 develop) growth [Kuhl, 2014]. In terms of elasticity, we now consider a simple 26 incompressible neo-hookean strain energy potential under the plane strain 27 assumption: 28

$$W^{e}\left(\underline{\underline{C}^{e}}\right) = \frac{\mu}{2}\left(\operatorname{tr}\left(\underline{\underline{C}^{e}}\right) - 2\right),\tag{38}$$

where μ is the shear modulus, taken as unity, and use the incompressible
mixed variational formulation (34) is used with evolution laws (33) & (36).
Figure 4 shows the response of the tissue modeled by relaxed growth, as a

³² function of normalized (with respect to growth time constant) time, and for
³³ various ratios of relaxation over growth characteristic times. Since growth is
³⁴ purely time-driven, the growth pattern is very similar in all cases. However,



Figure 3: Relaxed growth material response under constrained growth: normalized (with respect to shear modulus) hydrostatic pressure within a material point as a function of normalized (with respect to growth time constant τ^{g}) time, for various ratios of relaxation over growth characteristic times. For slow relaxation (*i.e.*, for large relaxation time constant τ^{r}), the model behaves as standard growth, with the development of residual stresses; conversely, for fast relaxation (*i.e.*, for small relaxation time constant τ^{r}), constrained growth happens with almost no induced residual stresses.

the level of growth-induced residual stresses varies drastically with the relaxation characteristic time: for very fast relaxation (compared to growth),
the muffin grows without developing large residual stresses, even in the constrained region.

⁵ 3.3. A strain-driven relaxed growth example: the artery

The final example is the canonical case of residual stresses in pressurized 6 arteries. We start from a simple disc representing an unloaded, stress-free 7 artery. The artery is loaded with some internal pressure, and allowed to grow 8 and relax for some time while the internal pressure is maintained. Then, the 9 loading is removed, and an opening angle experiment is simulated by making 10 a radial cut in the model, and letting the artery spring open by releasing 11 some residual stresses. Here we consider the simplest form of strain-driven 12 growth evolution law: 13

$$\begin{cases} \underline{F^g} = (1 + \theta^g) \underline{1} \\ \overline{\theta^g} (t = 0) = 0 \\ \dot{\theta}^g = \frac{\|\underline{E}^e\|}{\tau^g} \end{cases}$$
(39)

where τ^g still represents the growth time constant, and $\underline{\underline{E}^e} := \frac{1}{2} (\underline{\underline{C}^e} - \underline{\underline{1}})$ is the Green-Lagrange elastic strain. This growth evolution law leads, after temporal discretization, to the following expression:

$$\underline{\underline{\dot{F}}^{g,\mathrm{mid}}}_{g} = \frac{\|\underline{\underline{E}^{e,\mathrm{mid}}}\|}{\tau^g} \underline{\underline{1}}$$
(40)

We use the incompressible neo-hookean strain energy potential (38), and the
incompressible mixed variational formulation (34) with evolution laws (33)
& (40).

Figure 5 shows the response of the artery over time, for various levels of 20 relaxation characteristic times: it first inflates due to the applied pressure, 21 grows, deflates as the pressure is removed, and springs open. Note that since 22 residual stresses are here induced by heterogeneous, strain-driven growth de-23 scribed by growth evolution low (39), only a small amount of growth (with 24 respect to the pressure-induced deformation) is required to generate realistic 25 residual stresses, leading to physiological opening angles. This last example 26 illustrates the fact that the relaxed growth model allows to control the open-27 ing angle (*i.e.*, the residual stresses) induced by a given amount of growth. 28



Figure 4: Relaxed growth structural response under constrained and unconstrained growth: norm of Cauchy stress tensor superimposed onto the deformed domain as a function of normalized (with respect to growth time constant τ^r) time, for various ratios of relaxation over growth characteristic times. The faster the relaxation, the less residual stresses develop for the same amount of growth.



Figure 5: Relaxed growth response of a growing artery under pressure: circumferential strain superimposed onto the deformed artery at each step of the simulation (initial, after applying the internal pressure, after growth and relaxation, after removing the internal pressure, and after making a radial cut and letting the artery spring open), for various ratios of relaxation over growth characteristic times. The faster the relaxation, the less residual stresses, and hence a smaller opening angle develops for the same amount of growth.

¹ 4. Discussion

In this paper, we introduced a novel "relaxed growth" framework allowing 2 for a fine control of the amount of residual stresses generated during tissue 3 heterogeneous growth. It is a direct extension of the classical multiplicative 4 decomposition of the transformation gradient framework [Rodriguez et al., 5 1994, to which an additional sub-transformation is introduced in order to let 6 the original unloaded configuration to evolve, hence relaxing away some of 7 the residual stresses. To solve it numerically, we proposed here a monolithic mixed formulation, but a standard displacement formulation with internal 9 variables could be used equivalently. 10

This work could be extended in multiple directions. Most and foremost, 11 experimental characterization of the amount of residual stresses induced di-12 rectly by physiological and/or pathological growth is required to provide 13 elements of validation to relaxed growth models. Moreover, other relaxation 14 evolution laws could be formulated and tested, especially in case of cyclic 15 loading. 3D simulations could be run. More generally, since in the current 16 numerical procedure both the deformed and unloaded configurations are com-17 puted at once, it is straightforward to stop the simulation, for instance when 18 the total deformation becomes too important and the mesh too distorted. 19 transfer all the fields to the new unloaded configuration, and restart from 20 here. That way problems with extreme growth, for instance in organogene-21 sis, could be tackled, with a clear control in both growth and growth-induced 22 residual stresses. Finally, relaxed growth could be added personalized model-23 ing pipelines, at both modeling and estimation steps, in order to make more 24 objective and quantitative the handling of longitudinal clinical data, and 25 design the next generation of diagnosis and treatment optimization tools. 26

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30 References

Alnæs, M., Blechta, J., Hake, J., Johansson, A., Kehlet, B., Logg, A.,

Richardson, C., Ring, J., Rognes, M.E., Wells, G.N., 2015. The FEniCS

³³ Project Version 1.5. DOI:10.11588/ans.2015.100.20553.

Böl, M., Bolea Albero, A., 2014. On a new model for inhomogeneous volume
 growth of elastic bodies. Journal of the Mechanical Behavior of Biomedical
 Materials 29, 582–593. DOI:10.1016/j.jmbbm.2013.01.027.

⁴ Bolea Albero, A., Ehret, A.E., Böl, M., 2014. A new approach to the simulation of microbial biofilms by a theory of fluid-like pressure-restricted finite growth. Computer Methods in Applied Mechanics and Engineering 272, 271-289. DOI:10.1016/j.cma.2014.01.001.

⁸ Chapelle, D., Bathe, K.J., 2010. On the ellipticity condition for model⁹ parameter dependent mixed formulations. Computers & Structures 88,
¹⁰ 581–587. DOI:10.1016/j.compstruc.2010.01.009.

Ciarlet, P.G., Geymonat, G., 1982. Sur les lois de comportement en élasticité
non-linéaire compressible. Comptes Rendus de l'Académie des Sciences
Série II 295, 423—426.

Clatz, O., Sermesant, M., Bondiau, P.Y., Delingette, H., Warfield, S., Malandain, G., Ayache, N., 2005. Realistic simulation of the 3-D growth
of brain tumors in MR images coupling diffusion with biomechanical
deformation. IEEE Transactions on Medical Imaging 24, 1334–1346.
DOI:10.1109/TMI.2005.857217.

Cyron, C.J., Aydin, R.C., Humphrey, J.D., 2016. A homogenized constrained
 mixture (and mechanical analog) model for growth and remodeling of soft
 tissue. Biomechanics and Modeling in Mechanobiology 15, 1389–1403.
 DOI:10.1007/s10237-016-0770-9.

- Fung, Y.C., 1993. Biomechanics. Springer-Verlag, New York. DOI:10.1007/
 978-1-4757-2257-4.
- Genet, M., Lee, L.C., Baillargeon, B., Guccione, J.M., Kuhl, E., 2016. Modeling Pathologies of Diastolic and Systolic Heart Failure. Annals of Biomedical Engineering 44, 112–127. DOI:10.1007/s10439-015-1351-2.
- Genet, M., Lee, L.C., Ge, L., Acevedo-Bolton, G., Jeung, N., Martin, A.J.,
 Cambronero, N., Boyle, A.J., Yeghiazarians, Y., Kozerke, S., Guccione,
 J.M., 2015a. A Novel Method for Quantifying Smooth Regional Variations in Myocardial Contractility Within an Infarcted Human Left Ventricle Based on Delay-Enhanced Magnetic Resonance Imaging. Journal of
 Biomechanical Engineering 137. DOI:10.1115/1.4030667.

Genet, M., Rausch, M.K., Lee, L.C., Choy, S., Zhao, X., Kassab, G.S.,
 Kozerke, S., Guccione, J.M., Kuhl, E., 2015b. Heterogeneous growth induced prestrain in the heart. Journal of Biomechanics 48, 2080-2089.
 DOI:10.1016/j.jbiomech.2015.03.012.

Göktepe, S., Abilez, O.J., Kuhl, E., 2010. A generic approach towards finite
growth with examples of athlete's heart, cardiac dilation, and cardiac wall
thickening. Journal of the Mechanics and Physics of Solids 58, 1661–1680.

⁸ DOI:10.1016/j.jmps.2010.07.003.

Holmes, J.W., 2004. Candidate mechanical stimuli for hypertrophy during
volume overload. Journal of Applied Physiology 97, 1453-1460. DOI:10.
1152/japplphysiol.00834.2003.

Hughes, T.J.R., 2000. The Finite Element Method: Linear Static and Dy namic Finite Element Analysis. Dover Publications, Mineola, NY.

Humphrey, J.D., Rajagopal, K.R., 2002. A constrained mixture model for
growth and remodeling of soft tissues. Mathematical Models and Methods
in Applied Sciences 12, 407–430. DOI:10.1142/S0218202502001714.

Kerckhoffs, R.C.P., Omens, J.H., McCulloch, A.D., 2012. A single strain based growth law predicts concentric and eccentric cardiac growth during
 pressure and volume overload. Mechanics Research Communications 42,
 40-50. DOI:10.1016/j.mechrescom.2011.11.004.

Krishnamurthy, A., Villongco, C.T., Chuang, J., Frank, L.R., Nigam, V.,
Belezzuoli, E., Stark, P., Krummen, D.E., Narayan, S.M., Omens, J.H.,
McCulloch, A.D., Kerckhoffs, R.C.P., 2013. Patient-Specific Models
of Cardiac Biomechanics. Journal of computational physics 244, 4–21.
DOI:10.1016/j.jcp.2012.09.015.

Kroon, W., Delhaas, T., Arts, T., Bovendeerd, P.H.M., 2009. Computational modeling of volumetric soft tissue growth: Application to the cardiac
left ventricle. Biomechanics and Modeling in Mechanobiology 8, 301–309.
DOI:10.1007/s10237-008-0136-z.

Kuhl, E., 2014. Growing matter: A review of growth in living systems.
 Journal of the Mechanical Behavior of Biomedical Materials 29, 529-543.
 DOI:10.1016/j.jmbbm.2013.10.009.

Lee, L.C., Genet, M., Acevedo-Bolton, G., Ordovas, K., Guccione, J.M.,
Kuhl, E., 2014. A computational model that predicts reverse growth
in response to mechanical unloading. Biomechanics and Modeling in
Mechanobiology 14, 217–229. DOI:10.1007/s10237-014-0598-0.

Lee, L.C., Sundnes, J.S., Genet, M., Wall, S.T., 2016. Physics-based com puter simulation of the long-term effects of cardiac regenerative therapies.
 Technology 4, 23–29. DOI:10.1142/S2339547816400069.

⁸ Lee, L.C., Sundnes, J.S., Genet, M., Wenk, J.F., Wall, S.T., 2015. An
⁹ integrated electromechanical-growth heart model for simulating cardiac
¹⁰ therapies. Biomechanics and Modeling in Mechanobiology, 1–13DOI:10.
¹¹ 1007/s10237-015-0723-8.

Liu, S.Q., Fung, Y.C., 1989. Relationship between hypertension, hypertrophy, and opening angle of zero-stress state of arteries following aortic constriction. Journal of Biomechanical Engineering 111, 325–335.
DOI:10.1115/1.3168386.

Logg, A., Mardal, K.A., Wells, G. (Eds.), 2012. Automated Solution of
Differential Equations by the Finite Element Method: The FEniCS Book.
Number 84 in Lecture Notes in Computational Science and Engineering,
Springer, Heidelberg.

Maillet, M., van Berlo, J.H., Molkentin, J.D., 2013. Molecular basis of phys iological heart growth: Fundamental concepts and new players. Nature
 reviews. Molecular cell biology 14, 38–48. DOI:10.1038/nrm3495.

Omens, J.H., Rodriguez, E.K., McCulloch, A.D., 1996. Transmural Changes
 in Stress-free Myocyte Morphology During Pressure Overload Hypertrophy
 in the Rat. Journal of Molecular and Cellular Cardiology 28, 1975–1983.
 DOI:10.1006/jmcc.1996.0190.

Rausch, M.K., Zöllner, A.M., Genet, M., Baillargeon, B., Bothe, W., Kuhl,
E., 2017. A virtual sizing tool for mitral valve annuloplasty. International
Journal for Numerical Methods in Biomedical Engineering 33, e02788.
DOI:10.1002/cnm.2788.

Rodriguez, E.K., Hoger, A., McCulloch, A.D., 1994. Stress-dependent finite
 growth in soft elastic tissues. Journal of Biomechanics 27, 455–467.

 Roth, C.J., Yoshihara, L., Ismail, M., Wall, W.A., 2017. Computational modelling of the respiratory system: Discussion of coupled modelling approaches and two recent extensions. Computer Methods in Applied Mechanics and Engineering 314, 473–493. DOI:10.1016/j.cma.2016.08.
 010.

Sermesant, M., Chabiniok, R., Chinchapatnam, P., Mansi, T., Billet, F., Moireau, P., Peyrat, J.M., Wong, K.C.L., Relan, J., Rhode, K.S., Ginks, M.R., Lambiase, P.D., Delingette, H., Sorine, M., Rinaldi, C.A., Chapelle, D., Razavi, R., Ayache, N., 2012. Patient-specific electromechanical models of the heart for the prediction of pacing acute effects in CRT: A preliminary clinical validation. Medical Image Analysis 16, 201–15. DOI:10.1016/j. media.2011.07.003.

Skalak, R., Zargaryan, S., Jain, R.K., Netti, P.A., Hoger, A., 1996. Compatibility and the genesis of residual stress by volumetric growth. Journal of
Mathematical Biology, 26.

Smith, N.P., de Vecchi, A., McCormick, M., Nordsletten, D.A., Camara,
O., Frangi, A.F., Delingette, H., Sermesant, M., Relan, J., Ayache, N.,
Krueger, M.W., Schulze, W.H.W., Hose, R., Valverde, I., Beerbaum, P.,
Staicu, C., Siebes, M., Spaan, J., Hunter, P.J., Weese, J., Lehmann, H.,
Chapelle, D., Rezavi, R., 2011. euHeart: Personalized and integrated
cardiac care using patient-specific cardiovascular modelling. Interface focus
1, 349–64. DOI:10.1098/rsfs.2010.0048.

Taber, L.A., 1995. Biomechanics of Growth, Remodeling, and Morphogenesis. Applied Mechanics Reviews 48, 487. DOI:10.1115/1.3005109.

Taylor, C.A., Figueroa, C.A., 2009. Patient-specific modeling of cardiovascular mechanics. Annual Review of Biomedical Engineering 11, 109–134.
DOI:10.1146/annurev.bioeng.10.061807.160521.

Valentín, A., Holzapfel, G.A., 2012. Constrained Mixture Models as Tools
for Testing Competing Hypotheses in Arterial Biomechanics: A Brief Survey. Mechanics research communications 42, 126–133. DOI:10.1016/j.
mechrescom.2012.02.003.

Valentín, A., Humphrey, J.D., Holzapfel, G.A., 2013. A finite element-based
 constrained mixture implementation for arterial growth, remodeling, and

¹ adaptation: Theory and numerical verification. International journal for

- numerical methods in biomedical engineering 29, 822–49. DOI:10.1002/
 cnm.2555.
- $_{\rm 4}\,$ Witzenburg, C.M., Holmes, J.W., 2017. A Comparison of Phenomenologic
- ⁵ Growth Laws for Myocardial Hypertrophy. Journal of Elasticity DOI:10.
- 6 1007/s10659-017-9631-8.
- Xi, J., Lamata, P., Lee, J., Moireau, P., Chapelle, D., Smith, N.P., 2011.
 Myocardial transversely isotropic material parameter estimation from in silico measurements based on a reduced-order unscented Kalman filter.
- Journal of the mechanical behavior of biomedical materials 4, 1090–102.
- ¹¹ DOI:10.1016/j.jmbbm.2011.03.018.