# Towards the modelling of ageing and atherosclerosis effects in $ApoE^{-/-}$ mice aortic tissue

Tobias Waffenschmidt <sup>a,g</sup>, Myriam Cilla <sup>b</sup>, Pablo Sáez <sup>d</sup>, Marta M. Pérez <sup>e</sup>, Miguel A. Martínez <sup>c</sup>, Andreas Menzel <sup>a,f</sup>, Estefanía Peña <sup>c,\*</sup>

<sup>a</sup>Institute of Mechanics, Department of Mechanical Engineering, TU Dortmund, Germany

<sup>b</sup>Centro Universitario de la Defensa, Academia General Militar, Zaragoza, Spain

<sup>c</sup>Applied Mechanics and Bioengineering, Aragón Institute of Engineering Research (I3A), University
of Zaragoza, CIBER de Bioingeniería, Biomateriales y Nanomedicina (CIBER-BBN), Spain

<sup>d</sup>Laboratori de Calcul Numeric (LaCaN), Universitat Politecnica de Catalunya, Barcelona, Spain

<sup>e</sup>Department of Anatomy, Embryology and Genetics, Veterinary Faculty, University of Zaragoza

<sup>f</sup>Division of Solid Mechanics, Lund University, Sweden

<sup>g</sup>3M Deutschland GmbH, Carl-Schurz-Str. 1, D-41453 Neuss, Germany

#### Abstract

The goal of this work consists in a quantitative analysis and constitutive modelling of ageing processes associated to plaque formation in mice arteries. Reliable information on the characteristic evolution of pressure-stretch curves due to the ageing effects are extracted from previous inflation test experiments. Furthermore, characteristic age-dependent material parameters are identified on the basis of a continuum-mechanics-based parameter optimisation technique.

The results indicate that the aorta-stiffness of the healthy control mice remains basically constant irrespective of the diet-time and age. In contrast, significant differences exist within the material response and in consequence within the material parameters between the  ${\rm ApoE^{-/-}}$  and the control mice as well as for the different locations over the aorta which is underlined by our experimental observations. With regard to the temporal evolution of the material parameters, we observe that the material parameters for the  ${\rm ApoE^{-/-}}$  mice aortas exhibit a saturation-type increase with respect to age.

Key words: Ageing, anisotropic biological tissues, thick-walled tube, parameter identification.

<sup>\*</sup> Corresponding author. Estefanía Peña. Mechanical Engineering Department. c/ Maria de Luna s/n 50018. Zaragoza. Spain. Tel.: +34 876555233; Fax: +34 976762578

Email address: fany@unizar.es (Estefanía Peña ).

## 1 Introduction

- Recent experimental studies have shown that atherosclerosis significantly modifies the mechanical properties of the arterial tissue (Teng et al., 2014). In this regard, a combination of inflammatory, biological and mechanical processes tends to remodel the arterial wall structure and composition (Tracqui et al., 2011). Manifold animal species are commonly used to study the pathogenesis and potential treatment of the lesions of atherosclerosis (Buja et al., 1983; Faggiotto et al., 1984; Reitman et al., 1982; Schwartz et al., 1985; Wagner, 1978). Experimental tests on mice over a period of several months can be carried out and results may be extrapolated for the benefit of middle-aged or elderly humans.
- Studying the mechanical properties during atheroma plaque development is particularly im-10 portant in order to understand the mechanisms of vascular adaptation in response to changes 11 in physical stress. Several studies investigate the morphological, structural, and biochemical 12 changes of the aorta as well as the relation between these changes and the mechanics of the 13 aorta (Davis, 1995; Huang et al., 2006; Machii and Becker, 1997; Wells et al., 1999; Wong 14 and Langille, 1996; Tsamis et al., 2013). Most of these studies, however, mainly focus on the proximal agree, and none of them systematically characterise the geometrical and mechanical 16 properties along the entire length of the aorta. Even though Guo and Kassab (2003) study the 17 whole agrta, they exclusively focus on the standard laboratory mice (C57BL/6J mice). 18
- The modelling, simulation and experimental study of arteries and associated vascular diseases such as atherosclerosis is an area of current multidisciplinary research. In order to enable an individual patient-specific adaptation/modification of the medical treatment, it is necessary to understand the underlying mechanisms of atherosclerotic plaque evolution. In this context, modelling and simulation are becoming increasingly important research topics (Cilla et al., 2014). As *in-vivo* experiments are usually not feasible for the study of human arteries, both

modelling and simulation of ageing phenomena have become vital for the improvement of 25 the reliability of medical prognoses. For the description of the purely mechanical and passive 26 behaviour of arteries, several computational material models have already been developed in 27 the past decades (Holzapfel et al., 2000). In this context, we consider continuum mechanics 28 models and numerical approaches to validate and calibrate our constitutive models based on 29 the experimental investigations. However, the temporal evolution in these vessels, commonly 30 referred to as ageing, has hardly been discussed in the literature. Nevertheless, the incorporation 31 of such ageing effects—in combination with the detection, description, and especially interaction 32 with plaques—is of central importance for the understanding of atherosclerosis. 33

The scientific goal of this work is the mechanical analysis and constitutive modelling of age-34 ing processes usually associated with plaque formation in mouse arteries. Specifically speak-35 ing, reliable information on the evolution of pressure-stretch curves due to ageing effects is 36 extracted from previous experimental investigations (Cilla et al., 2015). A computational me-37 chanics framework is used to identify age-dependent material parameters on the basis of the 38 experimental data. We consider an inhomogeneous boundary value problem at large defor-39 mations for an incompressible thick-walled cylindrical tube, which is assumed to be a rough 40 approximation for a real artery. 41

## <sup>42</sup> 2 Materials and methods

#### $^{43}$ 2.1 Mechanical tests

Female apolipoprotein E-deficient transgenic mice (ApoE<sup>-/-</sup> mice) and common laboratory control mice (C57BL/6 mice) are used to obtain the pressure-diameter relation (Cilla et al., 2015). We analyse the progressive evolution of atherosclerotic lesions at three different locations of the aorta of (i) ApoE<sup>-/-</sup> mice put on a hyper-lipidic Western diet and (ii) C57BL/6 control

- mice put on a normal chow diet. Different sets of five  $ApoE^{-/-}$  and control mice are sacrificed after 10, 20, 30 and 40 weeks.
- Inflation tests are performed as described in Cilla et al. (2015). The aorta is preconditioned within three cycles from  $0 \ [mmHg] 200 \ [mmHg]$ . Thereafter, the perfusion pressure  $p_i$  is increased in steps of  $25 \ [mmHg]$  from  $p_i = 0 \ [mmHg] 250 \ [mmHg]$ , (Guo and Kassab, 2003), while the outer diameter  $d_o$  along the trunk of the aorta at each pressure step is recorded. The blood vessel is subdivided into a series of short regions of approximately  $3 \ [mm] 4 \ [mm]$  length per segment, and the external diameter and circumferential stretch are both measured pointwise. The outer circumferential stretch is evaluated as the ratio of the current and referential outer diameters  $d_o$  and  $D_o$ , i.e.  $\lambda_{\theta_o} = \frac{d_o}{D_o}$ , details on Cilla et al. (2015).
- An average pressure-circumferential-stress-relation for each of the segments is measured. Three
  different zones of the aorta are studied separately, i.e. the upper thoracic aorta, the lower
  thoracic aorta and the abdominal aorta. The upper thoracic zone is considered to reach from
  the aortic arch to the ninth thoracic vertebra, the lower thoracic up to the superior
  mesenteric branch and the abdominal part from there to the iliac bifurcation. They
  are not identical to ascending and descending thoracic aorta. The objective was to divide the
  aorta in regions of similar lengths. This description of the upper and lower thoracic aorta is
  included in the new version of the manuscript.

# $^{5}$ 2.2 Computational framework

We propose a computational mechanics framework to identify age-dependent material parameters of the data obtained by the experimental tets.

#### 2.2.1 Kinematics

78

Position vectors of particles in an undeformed reference configuration  $\mathcal{B}_0$  are denoted by X and position vectors in the deformed current configuration  $\mathcal{B}_t$  at time t by  $x = \varphi(X, t)$ . The kinematics of a thick-walled tube can conveniently be described by cylindrical polar coordinates. These coordinates are introduced as R,  $\Theta$  and Z with respect to a chosen reference configuration  $\mathcal{B}_0$  as well as r,  $\theta$  and z with respect to the current configuration  $\mathcal{B}_t$ . An orthonormal referential and spatial frame, respectively, can be defined in terms of these coordinates as

$$\boldsymbol{E}_{R}(\Theta) = \cos(\Theta) \, \boldsymbol{e}_{1} + \sin(\Theta) \, \boldsymbol{e}_{2}, \qquad \boldsymbol{e}_{r}(\theta) = \cos(\theta) \, \boldsymbol{e}_{1} + \sin(\theta) \, \boldsymbol{e}_{2}, \qquad (1)$$

$$\boldsymbol{E}_{\Theta}(\Theta) = -\sin(\Theta)\,\boldsymbol{e}_{1} + \cos(\Theta)\,\boldsymbol{e}_{2}\,, \qquad \boldsymbol{e}_{\theta}(\theta) = -\sin(\theta)\,\boldsymbol{e}_{1} + \cos(\theta)\,\boldsymbol{e}_{2}\,, \qquad (2)$$

$$\boldsymbol{E}_Z = \boldsymbol{e}_3, \qquad \boldsymbol{e}_z = \boldsymbol{e}_3. \tag{3}$$

wherein  $\{e_1, e_2, e_3\}$  is a Cartesian frame fixed in space. The geometry of the tube considered is visualised in Figure 1 and its material and spatial settings are specified by

[Fig. 1 about here.]

$$R_i \le R \le R_o \,, \qquad \qquad r_i \le r \le r_o \,, \tag{4}$$

$$0 \le \Theta \le 2\pi, \qquad 0 \le \theta \le 2\pi, \qquad (5)$$

$$0 \le Z \le L, \qquad 0 \le z \le l, \tag{6}$$

where  $R_i$ ,  $R_o$  and L represent the inner and outer radii and the length of the tube in a (undeformed) reference configuration  $\mathcal{B}_0$ , while  $r_i$ ,  $r_o$  and l represent the corresponding quantities in the (deformed) current configuration  $\mathcal{B}_t$ ; see Figure 1. The deformation modes of inflation and extension of an incompressible tube can be represented by the spatial position vector

$$\boldsymbol{x} = \boldsymbol{\varphi}(\boldsymbol{X}, t) = r \, \boldsymbol{e}_r(\theta) + z \, \boldsymbol{e}_z \tag{7}$$

 $_{83}$  specified by means of

$$r = \sqrt{\frac{R^2 - R_i^2}{\lambda_z} + r_i^2}, \qquad \theta = \Theta, \qquad z = \lambda_z Z,$$
(8)

where  $\lambda_z$  represents the axial stretch.

Representative deformation measures can be introduced with respect to the coordinates R,  $\Theta$ , Z and r,  $\theta$ , z. In particular, equation (7) is used together with  $\nabla_{\mathbf{X}} [\bullet] = \partial_R [\bullet] \otimes \mathbf{E}_R + \mathbf{E}_R [\bullet] \otimes \mathbf{E}_R$ 

$$\mathbf{F} = \lambda_r \, \mathbf{e}_r \otimes \mathbf{E}_R + \lambda_\theta \, \mathbf{e}_\theta \otimes \mathbf{E}_\Theta + \lambda_z \, \mathbf{e}_z \otimes \mathbf{E}_Z \,, \tag{9}$$

89 where the radial and circumferential stretch are each introduced as

$$\lambda_r = \frac{R}{r \, \lambda_z}$$
 and  $\lambda_\theta = [\lambda_r \lambda_z]^{-1} = \frac{r}{R}$ . (10)

 $\lambda_r$ ,  $\lambda_\theta$  and  $\lambda_z$  take the interpretation as principal stretches in radial, circumferential and axial direction.

#### 92 2.2.2 Constitutive model

We apply an orthotropic model with two families of fibres as introduced by Holzapfel et al. (2000). The strain energy function of this model is assumed to additively decompose into an isotropic part  $\Psi_{\rm iso}$ , representing the contribution of the non-collagenous ground material, and into an anisotropic part  $\Psi_{\rm ani}$ , representing the contributions of the different families of collagen fibres, i.e.

$$\Psi(\mathbf{F}, \mathbf{a}_{0i}) = \Psi_{\text{iso}}(\mathbf{F}) + \Psi_{\text{ani}}(\mathbf{F}, \mathbf{a}_{0i}). \tag{11}$$

Moreover,  $a_{0i}$  denotes a set of i = 1, ..., N referential unit-vectors characterising the mean orientations of the fibre families. The isotropic part of the strain energy is specified by a common neo-Hookean format

$$\Psi_{\rm iso}(\mathbf{F}) = \frac{c}{2} \left[ I_1 - 3 \right], \tag{12}$$

with  $I_1 = \mathbf{F} : \mathbf{F} = \lambda_{\theta}^2 + \lambda_z^2 + \lambda_{\theta}^{-2} \lambda_z^{-2}$  for the incompressible case, i.e.  $J = \det(\mathbf{F}) \doteq 1$ . The anisotropic part adopted takes the following exponential form

$$\Psi_{\text{ani}}(\boldsymbol{F}, \boldsymbol{a}_{0i}) = \frac{k_1}{2 k_2} \sum_{i=1}^{N} \left[ \exp\left(k_2 \langle E_i \rangle^2\right) - 1 \right], \tag{13}$$

wherein it is assumed that the fibres are mechanically equivalent. The notation  $\langle \bullet \rangle = [|\bullet| + \bullet]/2$ reflects the Macaulay brackets. These allow the activation of the fibre contributions in the tension regime only. A referential strain measure  $E_i$  is introduced as

$$E_i = \mathbf{a}_{0i} \cdot \mathbf{F}^{t} \cdot \mathbf{F} \cdot \mathbf{a}_{0i} - 1 = I_{4i} - 1.$$

$$(14)$$

The number of mechanically equivalent fibre families is restricted to N=2 and, moreover, their initial orientations are assumed as

$$\boldsymbol{a}_{01,2} = \sin(\beta) \, \boldsymbol{E}_Z \pm \cos(\beta) \, \boldsymbol{E}_{\Theta} \,, \tag{15}$$

so that the invariant introduced in equation (14) can be expressed as

$$I_{4i} = \boldsymbol{a}_{0i} \cdot \boldsymbol{F}^{t} \cdot \boldsymbol{F} \cdot \boldsymbol{a}_{0i} = \sin^{2}(\beta) \, \lambda_{z}^{2} + \cos^{2}(\beta) \, \lambda_{\theta}^{2}. \tag{16}$$

With these relations at hand, the Cauchy stress tensor  $\sigma = J^{-1} \partial_{\mathbf{F}} \Psi \cdot \mathbf{F}^{t}$  can be specified, namely

$$\boldsymbol{\sigma} = c \, \boldsymbol{F} \cdot \boldsymbol{F}^{t} + 4 \, k_{1} \, E \exp\left(k_{2} \left\langle E \right\rangle^{2}\right) \left[\, \boldsymbol{a}_{1} \otimes \boldsymbol{a}_{1} + \boldsymbol{a}_{2} \otimes \boldsymbol{a}_{2} \,\right], \tag{17}$$

with  $\boldsymbol{a}_{1,2} = \boldsymbol{F} \cdot \boldsymbol{a}_{0\,1,2}$ , and spectral form

$$\sigma_{rr} = c \,\lambda_{\theta}^{-2} \,\lambda_{z}^{-2} \,, \tag{18}$$

$$\sigma_{\theta\theta} = c \lambda_{\theta}^2 + 4\cos^2(\beta) k_1 \lambda_{\theta}^2 E \exp\left(k_2 \langle E \rangle^2\right), \qquad (19)$$

$$\sigma_{zz} = c \lambda_z^2 + 4 \sin^2(\beta) k_1 \lambda_z^2 E \exp\left(k_2 \langle E \rangle^2\right). \tag{20}$$

# 2.2.3 Equilibrium conditions

Neglecting body forces, the underlying local equilibrium conditions in terms of spatial arguments take the representation as

$$\mathbf{0} = \nabla_x \cdot \boldsymbol{\sigma} \qquad \text{in} \quad \mathcal{B}_t \quad , \tag{21}$$

$$t = \boldsymbol{\sigma} \cdot \boldsymbol{n}$$
 on  $\partial \mathcal{B}_t$ , (22)

$$\mathbf{t} = -p_i \mathbf{n}$$
 on  $\partial \mathcal{B}_{ti}$ . (23)

By analogy with the derivations reviewed above and with  $n=\pm e_r$ , the Euler-Lagrange equations can be summarised as

$$\mathbf{0} = \left[ \frac{\partial \sigma_{rr}}{\partial r} + \frac{\sigma_{rr} - \sigma_{\theta\theta}}{r} \right] \mathbf{e}_r + \frac{1}{r} \frac{\partial \sigma_{\theta\theta}}{\partial \theta} \mathbf{e}_{\theta} + \frac{\partial \sigma_{zz}}{\partial z} \mathbf{e}_z \quad \text{in} \quad \mathcal{B}_t , \quad (24)$$

$$\mathbf{t} = \sigma_{rr} \, \mathbf{e}_r$$
 on  $\partial \mathcal{B}_{to}$ , (25)

$$\mathbf{t} = p_i \, \mathbf{e}_r$$
 on  $\partial \mathcal{B}_{ti}$ . (26)

Due to geometrical and constitutive symmetry, the only non-trivial component of (24) is

$$\frac{\partial \sigma_{rr}}{\partial r} + \frac{\sigma_{rr} - \sigma_{\theta\theta}}{r} = 0, \qquad (27)$$

see, for example, Ogden (1997). From this equation and from boundary condition  $\sigma_{rr}|_{r=r_o} = 0$ on the outer surface of the tube, the radial Cauchy stress  $\sigma_{rr}$  may be calculated as

$$\sigma_{rr}(\xi) = \int_{\xi}^{r_o} [\sigma_{rr} - \sigma_{\theta\theta}] \frac{\mathrm{d}\xi}{\xi} \,. \tag{28}$$

The internal pressure  $p_i = -\sigma_{rr}|_{r=r_i}$  is then obtained in the form

$$p_i = \int_{r_i}^{r_o} [\sigma_{\theta\theta} - \sigma_{rr}] \frac{\mathrm{d}r}{r} \,. \tag{29}$$

## 2.2.4 Material parameter identification

The objective is to determine the characteristic evolution of material parameters in time on the basis of the experimental measurements. Several assumptions and approximations are included into the model. First, we idealise the aortic geometry by adopting a thick-walled tube and a single layer. Secondly, we replicate the experimental setting by imposing boundary conditions and kinematic constraints: fully incompressible, uniform internal pressure at the inner diameter, stress-free at the outer diameter of the tube (the outer circumferential stretch  $\lambda_{\theta_o}$ is one for zero pressure for all experimental pressure-stretch-curves), constant axial residual stretch, and circumferential residual stretch neglected. Thirdly, we adopt the following constitutive assumptions: hyperelastic (inelastic effects neglected), anisotropic (transversely isotropic or orthotropic), two mechanically equivalent families of fibres, fibres oriented in tangential plane of the tube, non-zero mechanical response of the fibres in tension-regime only, fibre dispersion neglected, and homogeneous material properties.

# [Table 1 about here.]

134

150

151

In order to investigate the age-dependent evolution of the material parameters in detail, we first fit the material parameters associated with the constitutive model to the pressure-outercircumferential-stretch curves recorded during the experiments  $(p_i - \lambda_{\theta_o}$ -curves).

For the computational fitting process, we apply a pressure-driven parameter identification pro-138 cedure where, by analogy to the experimental approach, the internal pressure  $p_i^{\text{exp}}$  is applied 139 to the tube, see Table 2 for algorithmic details. The equilibrium condition (24) is not fulfilled 140 a priori and therefore requires an additional iterative solution procedure. We apply a com-141 mon Newton-Raphson iteration scheme to iteratively satisfy the equilibrium condition of the 142 underlying boundary value problem as summarised in Table 2. This enables us to compute 143 an objective function f formulated in terms of the stretch-difference  $\lambda_{\theta_o} - \lambda_{\theta_o}^{\text{exp}}$ . The objective 144 function is then minimised with respect to the material parameters  $\nu$ , see Table 3. Generally, 145 such minimisation problems may conveniently be solved by typical optimisation techniques. In this study, the Matlab optimisation-algorithm fmincon is used which is based on a sequential 147 quadratic programming (SQP) method. We end up with a set of material parameters  $\nu^{\min}$ 148 providing optimal data fitting capabilities for the particular experimental curve of interest. 149

[Table 2 about here.]

[Table 3 about here.]

The material parameter identification procedure is based on a sequential optimisation of three 152 material parameters c,  $k_1$  and  $k_2$  as described in Sáez et al. (2014). In a first step, parameter 153 c is identified with  $k_1 = \text{const}$  and  $k_2 = \text{const}$  using the first two experimental data points, 154 in a second step  $k_1$  and  $k_2$  are identified with c = const using all experimental data points. 155 This non-standard sequential optimisation technique was applied, as it may occur that a single 156 optimisation including every experimental data point for all of the three material parameters 157 results in unphysical material parameters, e.g. such that c < 0, or in c = 1b for a constrained 158 optimisation including a lower bound 1b. 159

#### [Table 4 about here.]

Finally, we end up with a set of material parameters  $\nu^{min}$  providing optimal data fitting capabilities for the particular experimental curve of interest. The identified material parameters are then plotted over time/age in order to obtain an impression as to which parameter might be less/more affected due to the ageing effects.

#### 165 3 Results

160

172

#### 3.1 Mechanical properties

In order to illustrate the stiffening behaviour with respect to age, (Figure 2) shows the experimentally obtained circumferential stretch over different stages of ageing at three chosen pressure levels, i.e.  $p_i = \{50, 150, 250\}$  [mmHg].

We observed that the circumferential stretch  $\lambda_{\theta_o}$  decreases with age (stiffening) for the diseased ApoE<sup>-/-</sup> mice but stays relatively constant for the healthy control mice.

[Fig. 2 about here.]

As pointed out by Cilla et al. (2015), the ApoE<sup>-/-</sup> mice aorta stiffness increases dramatically with an increasing diet period. This holds for all the aortic zones studied. In contrast, the aorta stiffness of the control mice remains relatively constant for an increasing diet period. Comparing the ApoE<sup>-/-</sup> and control results, the ApoE<sup>-/-</sup> pressure-stretch curves after 10 [weeks] for the upper aorta and after 10 and 20 [weeks] for the lower and abdominal aorta are similar to the control curves for all ages.

# 3.2 Material parameter identification

179

184

185

The material parameters, i.e. c,  $k_1$  and  $k_2$ , which characterise the mechanical response of the artery are expected to evolve in time due to the ageing process. The fitting results of the material parameter identification procedure are depicted in Figure 3. The results in Figure 3 show excellent fitting capabilities.

[Fig. 3 about here.]

[Table 5 about here.]

The identified material parameters are summarised in Table 5 and their temporal evolution is 186 illustrated in Figure 4. With regard to the temporal evolution of the material parameters, we 187 expected the material parameters for the control mice to remain almost constant for a specific 188 location of the aorta. However, the parameters strongly deviate from each other with respect to 189 time, see parameter  $k_2$  in Figure 4. We also expected the material parameters for the ApoE<sup>-/-</sup> 190 mice aortas to exhibit a characteristic increase or decrease in time for one single location. And 191 we indeed observe that, e.g. parameter c first increases linearly within t = 10 - 30 [weeks]192 and saturates for t = 30 - 40 [weeks]. A similar behaviour holds for parameter  $k_1$ , whereas  $k_2$ , 193 remains almost constant within t = 10 - 30 [weeks] and then increases for t = 30 - 40 [weeks], see Figure 4. 195

#### 197 4 Discussion

A computational mechanics framework is used to identify age-dependent material parameters 198 on the basis of the experimental data. We consider an inhomogeneous boundary value problem 199 that allows for large deformations, i.e. an incompressible thick-walled cylindrical tube, which is 200 assumed to approximate a real artery. The basic deformation modes considered are represented 201 by combined inflation and axial extension. We make use of an orthotropic constitutive model 202 with two families of fibres and solve the underlying equilibrium conditions iteratively. We com-203 pute an objective function, represented by the difference between simulated and experimental 204 data, which is minimised with respect to the material parameters. As a main goal, we aim at 205 determining the characteristic age-dependent evolution of the material parameters over time. 206 Generally, the results show excellent fitting capabilities. Similar results have previously been 207 obtained by (Cheng et al., 2013). They used a fibre-based constitutive model to fit the me-208 chanical properties from the aorta during postnatal development of wild-type mice and elastin 209 haploinsufficient (Eln+/-) mice. Their results showed that a similar fibre-based constitutive 210 model was capable of distinguishing elastin content during development. 211

We observe significant deviations of the material parameters between the  ${\rm ApoE^{-/-}}$  and the 212 control mice as well as for the different locations over the aorta (upper, lower, abdominal) 213 which clearly underline our observations from the experiments. With regard to the temporal 214 evolution of the material parameters, we identify the material parameters for the control mice 215 aortas for one single location to remain almost constant. However, it becomes apparent that 216 the parameters quite strongly deviate from each other with respect to position mainly  $k_2$ . We 217 reporte the material parameters for the ApoE<sup>-/-</sup> mice aortas for one single location to exhibit 218 a characteristic increase/decrease in time. We indeed observe that parameter c first increase 219

linearly and saturate for t = 30 - 40 [weeks]. c parameter is the principle determinant of 220 stiffness in the small stretch region (pressure below physiological level), where the 221 crimped collagen molecules are primarily weaving. However, if there is a rupture of the elastin, there is no the transition region between the elastin-dominated and 223 the collagen-dominated parts of the stress-strain response, thus indicating earlier 224 collagen recruitment. For that reason, there is an increasing of the slope of the 225 stress-strain curve, so there is an increasing of the c parameter. This effect is due 226 to fragmentation of the elastin due to atherosclerosis and not due to ageing where 227 the content of elastin remains unaltered with age; however, its concentration is de-228 creased (Tsamis et al., 2013). A similar behaviour can be identified for parameter 229  $k_1$ , this parameter is the principle determinant of collagen stiffness in the small 230 stretch region of the stress-strain response, where the collagen molecules are pri-231 marily straightened and the tissue is resilient. Whereas  $k_2$  remained almost constant 232 within t = 10 - 30 [weeks] and then increased for t = 30 - 40 [weeks]. The main reason for 233 this fact could be that forty weeks may not suffice in order to study representative age-related 234 changes in elastic behavior for control groups, and that forty weeks is the start point of the 235 ageing process. However, additional studies are needed to understand and check this hypothesis. 236 Similar results were found on humans by Astrand et al. (2011). 237

To the best of the author's knowledge, not many studies containing all these characteristics appear to be available in the literature. Only Agianniotis and Stergiopulos (2012b) have so far compared the mechanical properties of young 10-12 week-old ApoE<sup>-/-</sup> mice without fat diet with C57BL/6J wild-type mice by extension-inflation mechanical tests. In their work, particular emphasis was placed on the computation of the Hudetz incremental elastic modulus. Previous studies suggest that 10% of collagen fibres are engaged at physiological pressure (Greenwald et al., 1997), whereas at higher pressures, the blood vessel becomes progressively less distensible as collagen fibres are recruited to support passive wall tension and to restrict aortic distension.

The pulse wave velocity increasement on ApoE<sup>-/-</sup> matches to the fragmentation of the elastic laminae in the arterial wall.

It should be pointed out that the study presented here contains some assumptions and limita-248 tions. First, the application of an inhomogeneous incompressible thick-walled cylindrical tube 249 which should represent a real artery is a significant idealisation and simplification. For future 250 studies, a non-cylindrical multi-layered problem setting should at least be considered. However, 251 the number of fitting parameters will increase, and to gain a realistic interpretation of their 252 physical sense some layer-specific information will be required. The assumption of circumfer-253 ential uniformity seems to be different between ApoE<sup>-/-</sup> mice, which have numerous plaques 254 concentrated around branches, and control mice, where the wall is much more uniform. The 255 plaques and geometry can influence the mechanical behaviour of the vessel and therefore distort 256 results due to the heterogeneous and inconsistent nature of their development. However, Trac-257 qui et al. (2011) measured the stiffness of the mice plaque and obtained much lower values with 258 respect to the vessel wall, so its effect on mechanical properties could be lower than expected. 259

In addition, material parameters can also be introduced as functions in time. To keep the fitting problem as simple as possible in order to avoid coupled mechanical effects, we did not
consider residual stretches. Although some investigations have previously been carried out to
study the aperture angle, these data cannot be uniquely related to the increase of residual
stresses. Moreover, circumferential residual stretches essentially require the acquisition of related experimental data which, in turn, is a non-trivial task and therefore not achievable at
this stage. In addition since these inflation tests are performed *in situ*, the measurement of
longitudinal forces, and thus the longitudinal stiffness cannot be assessed.

In spite of these limitations, the obtained results are promising and demonstrate substantial changes in the mechanical properties of  $ApoE^{-/-}$  mice aortas when compared to the results obtained from the control mice group. These findings are important for a better understanding

of the cardiovascular system of mice and could serve as a reference for future investigations of mechanical properties of blood vessels suffering atherosclerotic diseases.

# $_{ m 273}$ 5 Acknowledgements

Support from the Spanish Ministry of Economy and Competitiveness through the research projects DPI2010-20746-C03-01, PRI-AIBDE-2011-1216, and the CIBER initive and by the German Academic Exchange Service (Deutscher Akademischer Austauschdienst, DAAD) under PPP 54473666 are gratefully acknowledged.

#### 278 6 Conflict of interest

None of the authors of this work has conflict of interest with other people and organisations.

#### 280 References

- Agianniotis, A., Stergiopulos, N., 2012b. Wall properties of the apolipoprotein E-deficient mouse aorta. Atherosclerosis 223 (2), 314–320.
- <sup>283</sup> Astrand, H., Stalhand, J., Karlsson, J., Karlsson, M., Sonesson, B., , Länne, T., 2011. In vivo
- $_{284}$  estimation of the contribution of elastin and collagen to the mechanical properties in the
- human abdominal aorta: effect of age and sex. J Appl Physiol 110, 176–187.
- Buja, L., Kita, T., Goldstein, J., Watanabe, Y., Brown, M., 1983. Cellular pathology of progres-
- sive atherosclerosis in the WHHL rabbit. An animal model of familial hypercholesterolemia.
- Arteriosclerosis, Thrombosis, and Vascular Biology 3 (1), 87–101.
- <sup>289</sup> Cheng, J. K., Stoilov, I., Mecham, R. P., Wagenseil, J. E., 2013. A fiber-based constitutive

- model predicts changes in amount and organization of matrix proteins with development
- and disease in the mouse agrta. Biomech Model Mechanobiol 12, 497–510.
- <sup>292</sup> Cilla, M., Peña, E., Martínez, M. A., 2014. Mathematical modelling of atheroma plaque forma-
- tion and development in coronary arteries. J R Soc Interface 11, 20130866 1–16.
- <sup>294</sup> Cilla, M., Pérez, M. M., Peña, E., Martínez, M. A., 2015. Effect of diet and age on arterial
- stiffening due to atherosclerosis in ApoE<sup>-/-</sup> mice. Ann Biomed Eng In press.
- Davis, E. C., 1995. Elastic lamina growth in the developing mouse aorta. Journal of Histochem-
- 297 istry & Cytochemistry 43 (11), 1115–23.
- Faggiotto, A., Ross, R., Harker, L., 1984. Studies of hypercholesterolemia in the nonhuman
- primate. I. Changes that lead to fatty streak formation. Arteriosclerosis, Thrombosis, and
- 300 Vascular Biology 4 (4), 323–340.
- Greenwald, S. E., Moore, J. E. J., Rachev, A., Kane, T., Meister, J. J., 1997. Experimental
- investigation of the distribution of residual strains in the artery wall. ASME Journal of
- Biomechanical Engineering 119 (4), 438–444.
- Guo, X., Kassab, G. S., 2003. Variation of mechanical properties along the length of the aorta
- in C57BL/6 mice. American Journal of Physiology Heart and Circulatory Physiology 285,
- 306 Н2614-Н2622.
- Holzapfel, G., Gasser, T., Ogden, R., 2000. A new constitutive framework for arterial wall
- mechanics and a comparative study of material models. Journal of Elasticity 61, 1–48.
- Huang, Y., Guo, X., Kassab, G. S., 2006. Axial nonuniformity of geometric and mechanical
- properties of mouse agra is increased during postnatal growth. American Journal of Physi-
- ology Heart and Circulatory Physiology 290 (2), H657–H664.
- Machii, M., Becker, A. E., 1997. Morphologic features of the normal aortic arch in neonates,
- infants, and children pertinent to growth. The Annals of Thoracic Surgery 64 (2), 511–515.
- Ogden, R., 1997. Non-Linear Elastic Deformations. Dover.
- Ohayon, J., Mesnier, N., Broisat, A., Toczek, J., Riou, L., Tracqui, P., 2012. Elucidating
- $^{316}$  atherosclerotic vulnerable plaque rupture by modeling cross substitution of ApoE $^{-/-}$  mouse

- and human plaque components stiffnesses. Biomechanics and Modeling in Mechanobiology
- 11 (6), 801–813.
- Reitman, J., Mahley, R., Fry, D., 1982. Yucatan miniature swine as a model for diet-induced
- atherosclerosis. Atherosclerosis 43 (1), 119–132.
- Schwartz, C. J., Sprague, E. A., Kelley, J. L., Valente, A. J., Suenram, C. A., 1985. Aortic intimal
- monocyte recruitment in the normo and hypercholesterolemic baboon (*Papio Cynocephalus*).
- <sup>323</sup> Virchows Archiv 405, 175–191.
- Sáez, P., Peña, E., Martínez, M. A., 2014. A Structural Approach Including the Behavior of
- Collagen Cross-Links to Model Patient-Specific Human Carotid Arteries. Ann Biomed Eng
- <sup>326</sup> 42, 1158–1169.
- Teng, Z., Zhang, Y., Huang, Y., Feng, J., Yua, J., Lu, Q., Sutcliffe, M. P. F., Brown, A. J., Jing,
- 328 Z., Gillard, J. H., 2014. Material properties of components in human carotid atherosclerotic
- plaques: A uniaxial extension study. Acta Biomat 10, 5055–5063.
- Tracqui, P., Broisat, A., Toczek, J., Mesnier, N., Ohayon, J., Riou, L., 2011. Mapping elasticity
- moduli of atherosclerotic plaque in situ via atomic force microscopy. Journal of Structural
- Biology 174, 115–123.
- Tsamis, A., Krawiec, J. T., Vorp, D. A., 2013. Elastin and collagen fibre microstructure of the
- human aorta in ageing and disease: a review. J R Soc Interface 27, 10(83).
- Waffenschmidt, T., 2014. Modelling and simulation of adaptation and degradation in anisotropic
- biological tissues. Ph.D. thesis, Institut für Mechanik, Technische Universität Dortmund,
- http://hdl.handle.net/2003/31797.
- Waffenschmidt, T., Menzel, A., 2014. Extremal states of energy of a double-layered thick-
- walled tube Application to residually stressed arteries. Journal of the Mechanical Behavior
- of Biomedical Materials 29, 635–654.
- Wagner, W. D., 1978. Risk factors in pigeons genetically selected for increased atherosclerosis
- susceptibility. Atherosclerosis 31 (4), 453–463.
- Wells, S. M., Langille, B. L., Lee, J. M., Adamson, S. L., 1999. Determinants of mechanical

- properties in the developing ovine thoracic aorta. American Journal of Physiology Heart and Circulatory Physiology 277 (4), H1385–H1391.
- Wong, L. C. Y., Langille, B. L., 1996. Developmental Remodeling of the Internal Elastic Lamina of Rabbit Arteries: Effect of Blood Flow. Circulation Research 78 (5), 799–805.

Table 1 Setting of material, structural, geometrical and loading parameters used throughout all the following computations. The fibre angle  $\beta$  is based on data from Ohayon et al. (2012), the geometry is based on data from Cilla et al. (2015) and it has been obtained from histological analysis performed after the mechanical tests.

Type	Symbol	Description	Value	Unit
Material	c	elastic constant	to be identified	[kPa]
	$k_1$	elastic constant	to be identified	[kPa]
	$k_2$	elastic constant	to be identified	[-]
Structural	β	fibre angle	48.47	[deg]
Geometrical	$R_i$	inner referential radius	0.38197	[mm]
	H	referential wall thickness thoracic aorta 10 [weeks]	0.1	[mm]
	H	referential wall thickness abdominal aorta 10 [weeks]	0.08	[mm]
Loading	$\lambda_z$	axial stretch	1.0	[-]

Table 2 Algorithmic box for the material parameter identification procedure.

- (1) set up structural and geometrical parameters from Table 1 and collect these in pseudo-vector  $\boldsymbol{\kappa} = [\beta, R_i, H, \lambda_z]$
- (2) perform initial guess  $\nu = \nu^0 = \left[c^0, \, k_1^0, \, k_2^0\right]$  for the material parameters to be identified
- (3) set objective function to  $f(\mathbf{\nu}) = 0$
- (4) identify material parameters as argument of minimum of objective function f  $\boldsymbol{\nu}^{\min} = \arg\min_{\boldsymbol{\nu}} f(\boldsymbol{\nu}; \boldsymbol{\nu}^0, \boldsymbol{\kappa})$ ,

wherein  $f(\nu)$  is determined by the algorithm for the pressure-driven case and the minimisation can be performed by, e.g., the Matlab fmincon-optimisation-function

# Table 3 Algorithmic box for the pressure-driven deformation process were $h=1.0e^{-8}$ , $tol=1.0e^{-8}$ and w=1.0. All quantities are associated with $t_{n+1}$ .

- (1) given: internal pressure  $p_i^{\text{exp}}$  at time  $t_{n+1}$
- (2) perform initial guess  $\lambda_{\theta_o} = \lambda_{\theta_o}^0$  for the outer circumferential stretch
- (3) perform local Newton-Raphson iteration scheme
  - (a) compute residual

$$r(\lambda_{\theta_o}) = p_i^{\exp} - p_i(\lambda_{\theta_o})$$

wherein  $p_i(\lambda_{\theta_o})$  is determined by the algorithm described below

(b) compute linearisation of residual by means of forward difference scheme

$$\mathrm{d}r = \left[r(\lambda_{\theta_o} + h) - r(\lambda_{\theta_o})\right]/h$$
 with  $h \ll 1$ 

(c) Compute increment

$$\Delta \lambda_{\theta_o} = \mathrm{d}r/r$$

(d) compute update

$$\lambda_{\theta_o} \leftarrow \lambda_{\theta_o} - \Delta \lambda_{\theta_o}$$

(e) check tolerance with tol  $\ll 1$ 

if 
$$|r| < \text{tol go to } 4$$
.  
else go to 3. (a)

(4) compute objective function

$$f(\boldsymbol{\nu}) \leftarrow f(\boldsymbol{\nu}) + w \left[\lambda_{\theta_o} - \lambda_{\theta_o}^{\exp}\right]^2$$

#### Table 4

Algorithmic box for the determination of the internal pressure Waffenschmidt and Menzel (2014); Waffenschmidt (2014).

- (1) given: material parameters  $\boldsymbol{\nu} = [c, k_1, k_2]$ , structural and geometrical parameters  $\boldsymbol{\kappa} = [\beta, R_i, H, \lambda_z]$ , see Table 1, deformation in terms of the outer circumferential stretch  $\lambda_{\theta_o}$  at time  $t_{n+1}$
- (2) calculate referential outer radius  $R_o = R_i + H \label{eq:Ro}$
- (3) calculate current radii

$$r_o = \lambda_{\theta_o} R_o$$
 
$$r_i = \sqrt{r_o^2 - \left[R_o^2 - R_i^2\right]/\lambda_z}$$

- (4) apply m = 3-point Gaussian quadrature rule with quadrature points  $\xi_j = \left\{ -\sqrt{3/5} \ 0 \ \sqrt{3/5} \right\}$  and weights  $w_j = \left\{ 5/9 \ 8/9 \ 5/9 \right\}$ : loop over number of quadrature points  $j = 1, \ldots, m$ 
  - (a) calculate current radius

$$r_j = [[r_i + r_o] + \xi_j [r_o - r_i]]/2$$

(b) calculate referential radius

$$R_{j} = \sqrt{\lambda_{z} \left[ r_{j}^{2} - r_{i}^{2} \right] + R_{i}^{2}}$$

(c) calculate circumferential stretch

$$\lambda_{\theta j} = r_j / R_j$$

(d) calculate radial and circumferential stresses

$$\sigma_{rrj} = c \lambda_{\theta j}^{-2} \lambda_z^{-2}$$

$$\sigma_{\theta \theta j} = c \lambda_{\theta j}^2 + 4 \cos^2(\beta) k_1 \lambda_{\theta j}^2 E \exp\left(k_2 \langle E \rangle^2\right)$$
with  $E = \lambda_z^2 \sin(\beta)^2 + \lambda_{\theta j}^2 \cos(\beta)^2 - 1$  by means of equations (18, 19)

calculate internal pressure

$$p_i \approx \left[r_o - r_i\right]/2 \sum_{j=1}^{m} \left[\sigma_{\theta\theta j} - \sigma_{rrj}\right] w_j/r_j$$

Table 5 Identified material parameters by means of the pressure-driven material parameter identification procedure.

		diseased Apo $\mathrm{E}^{-/-}$ mice				healthy control mice			
	[weeks]	10	20	30	40	10	20	30	40
Upper aorta	c [kPa]	13.92	54.83	85.38	89.74	32.09	35.80	39.87	18.29
	$k_1 [kPa]$	116.70	155.08	167.36	160.41	128.96	125.49	142.81	113.99
	$k_2$ [-]	1.20	1.33	1.21	2.69	1.00	1.19	1.14	1.57
Lower aorta	c [kPa]	32.51	49.31	116.15	142.56	40.50	22.36	40.69	33.81
	$k_1 [kPa]$	123.90	142.81	171.43	158.89	136.79	118.49	150.56	131.24
	$k_2$ [-]	1.06	1.19	1.42	2.71	0.95	1.57	0.90	1.37
Abdominal aorta	c [kPa]	34.38	67.61	108.09	105.12	48.10	29.62	62.78	49.29
	$k_1 [kPa]$	122.75	144.24	173.31	180.15	132.20	112.06	150.12	147.65
	$k_2$ [-]	1.99	1.89	1.89	3.81	2.11	2.52	2.18	2.33

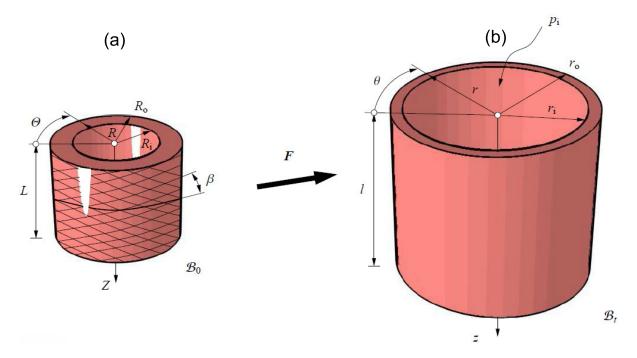


Fig. 1. Deformation modes of a thick-walled cylindrical tube: inflation (internal pressure  $p_i$ ) and axial extension (axial stretch  $\lambda_z = l/L$ ); arterial wall reinforced by two families of fibres with fibre angle  $\beta$  defined in (a) a stress-free reference configuration  $\mathcal{B}_0$  defined with respect to the circumferential direction; (b) residually stressed and loaded current configuration  $\mathcal{B}_t$ .

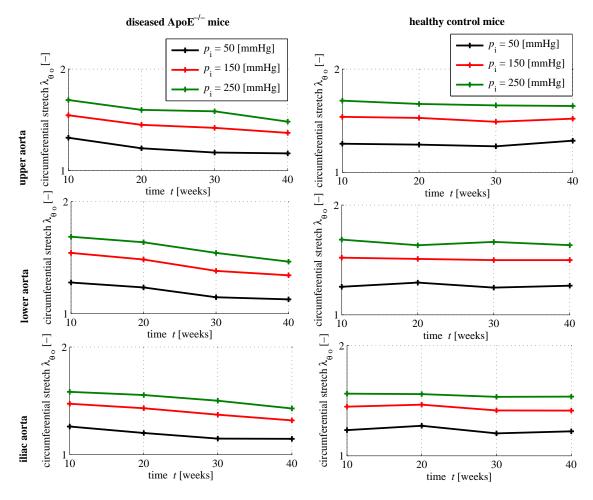


Fig. 2. Experimentally obtained circumferential stretch with respect to different stages of ageing at three chosen pressure levels, i.e.  $p_i = \{50, 150, 250\}$  [mmHg]. The left column corresponds to the diseased ApoE<sup>-/-</sup> mice, the right column corresponds to the healthy control mice. The three different rows top, middle, bottom are associated with different locations over the length of the considered arterial specimen (Cilla et al., 2015).

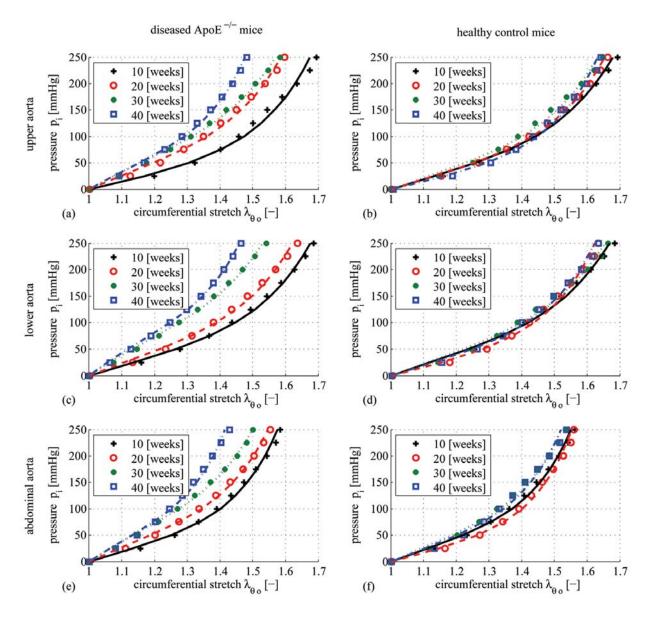


Fig. 3. Fitted experimental data based on the material parameter identification procedure. The discrete points represent the original experimental data from (Cilla et al., 2015), the lines the associated (simulated) fitting. The material parameter identification procedure is based on a sequential optimisation of three material parameters c,  $k_1$  and  $k_2$ : in a first step parameter c is identified with  $k_1 = \text{const}$  and  $k_2 = \text{const}$  using the first two experimental data points, in a second step  $k_1$  and  $k_2$  are identified with c = const using all experimental data points.

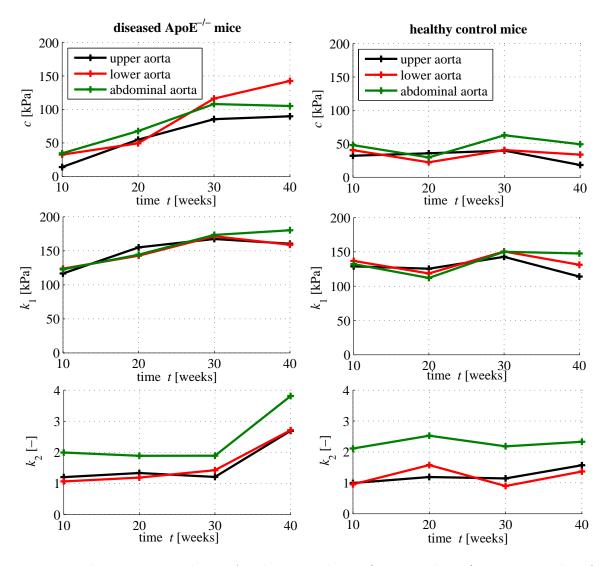


Fig. 4. Material parameter evolution for the stress-driven (pressure-driven) parameter identification procedure plotted over time, associated values given in Table 5. The left column corresponds to the diseased  ${\rm ApoE^{-/-}}$  mice, the right column corresponds to the healthy control mice.