

A CRACK PHASE-FIELD APPROACH TO MODEL AORTIC DISSECTIONS

Sandra P. Hager (1,2), Osman Gültekin (1,3), Gerhard A. Holzapfel (1,4)

1. Institute of Biomechanics, Graz University of Technology, Austria; 2. Faculty of Life Sciences and Medicine, King's College London, UK; 3. Department of Mechanical Engineering, Middle East Technical University, Ankara, Turkey; 4. Norwegian Institute of Science and Technology, Norway

Introduction

An aortic dissection describes a loss of structural integrity in the artery wall. The initial phase of this cardiovascular disease is a formation of a tear in the intimal sub-layer. During a cardiac cycle the heart jets additional blood in this region, which leads to a separation of sub-layers, mainly in the media. The cause for the formation of a crack and its propagation during a physiological cardiac cycle is still unclear. The aim of this abstract is to demonstrate a numerical method for the simulation of a crack formation between two medial sub-layers, as observed in patients.

Methods

The thermodynamic approach for the crack phase-field of fracture, as developed in [1] for an isotropic material, was used as a basis for the crack propagation model. The crack phase-field is described by the primary field variable d , which is interpolated between the intact state of the material ($d = 0$) and the ruptured state ($d = 1$). In general, a crack in a material describes a strong discontinuity in a domain (Fig. 1, solid bold curve). To smoothen this discontinuity over the domain, the length scale parameter l is used, defined by the mesh size.

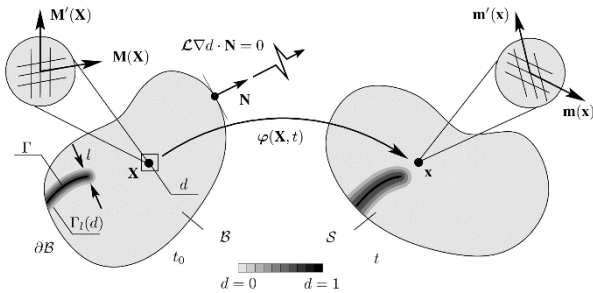


Figure 1: Overview of an anisotropic continuum body with a sharp crack surface Γ , and a diffusive crack surface $\Gamma_l(d)$.

The regularized anisotropic crack surface, which is based on the isotropic crack surface in [1], is described by

$$\Gamma_l(d) = \int_B \frac{1}{2l} (d^2 + \nabla d \cdot \mathcal{L} \nabla d) dV \quad (1)$$

where

$$\mathcal{L} = l^2 (\mathbf{I} + \omega_M \mathbf{M} \otimes \mathbf{M} + \omega_{M'} \mathbf{M}' \otimes \mathbf{M}') \quad (2)$$

is defined as a second-order anisotropic structure tensor. With the implementation of the tensor $\mathcal{L}$, the penalty term for the first fiber family ω_M and for the second fiber family $\omega_{M'}$ is included in the developed algorithm, see [2]. To solve the problem, the following considerations were used in their strong form, and

further with a standard Galerkin method in their weak form:

$$J \operatorname{div}(J^{-1} \tau) + \rho_0 \tilde{\gamma} = \mathbf{0} \quad (3)$$

$$\eta \dot{d} = (1 - d) \mathcal{H} - [d - \frac{1}{2} \operatorname{Div}(\mathcal{L} \nabla d)] \quad (4)$$

Equation (3) describes the balance of linear momentum in a continuum body for the kinematics of the mechanical problem, with the Kirchhoff stress tensor τ and the body force $\tilde{\gamma}$. Crack propagation is defined through Eq. (4), where the first term of the right hand side determines the crack driving force with the crack driving source term $\mathcal{H}$. This term can be defined by the combination of the isotropic (ground matrix) and the anisotropic (fibers) contribution of the material, and is further defined with the respective effective free-energy function and the regularized critical energy release rate. The second term in (4) defines the geometrical resistance of the material.

Results

To verify the model for an aortic dissection problem a multi-layer cylindrical domain with different sizes and shapes of an initial tear was used to simplify the aorta [2]. Two different cardiac cycles were applied to this domain, one physiological and one supra-physiological. Figure 2 illustrates the evolution of the crack phase-field.

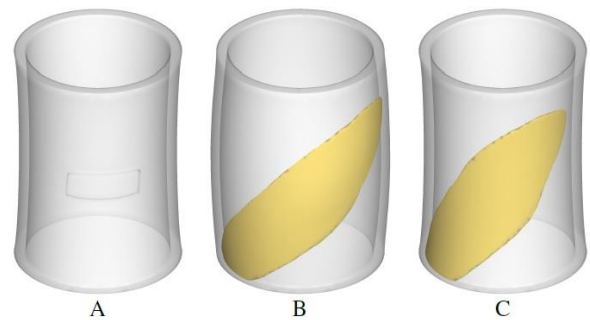


Figure 2: Evolution of the crack phase-field; A: start of simulation, B: peak supra-physiological cycle, C: end of simulation.

Discussion

With the discussed model, it is possible to reproduce the same pattern of crack propagation as observed clinically.

References

1. Miehe et al., Int. J. Numer. Meth. Eng., 83:1273-1311, 2010.
2. Gültekin et al., Biomech. Model. Mechanobiol., in press.

