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Analysis and Computation for a Fluid Mixture Model

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Abstract. A fluid mixture model of tissue deformations has been studied in this paper. The model is a mixed system of nonlinear hyperbolic and elliptic partial differential equations. Both theoretical linear stability and numerical analysis are presented. Comparisons between standard numerical methods that utilize Runge-Kutta methods coupled with the WENO scheme and the immersed interface methods are given. Numerical examples are also presented.

AMS subject classifications: 65N06, 65N12, 65M06, 76M20

Key words: Tissue deformations, immersed interface method, linear stability analysis, finite difference method.

1 Introduction

In this paper, we consider a mathematical model developed in [3,8,9] for modeling deformations of contractile mesenchymal tissues. The tissues are considered to be composed of two inter-penetrating material phases: an aqueous phase and a cell-fiber phase. The aqueous phase is composed of all the water and dissolved extracellular components of the tissues. The cell-fiber phase consists of the cells and the remaining, generally fibrous, extracellular components. It is assumed that: (1) the two phases occupy complementary portions of the space, (2) the aqueous phase behaves as a Stokes fluid, (3) the stresses in the cell-fiber phase are dissipated by permanent deformation on the relevant time scale and can also be treated as a Stokes flow. These assumptions lead to the following system

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of partial differential equations (in 1D):

$$\frac{\partial \theta}{\partial t} + \frac{\partial (\theta v)}{\partial x} = 0, \quad 0 < x < L, \tag{1.1}$$

$$\frac{\partial}{\partial x} \left(\frac{1-\theta}{\varphi \theta} \cdot \frac{\partial p}{\partial x} - v \right) = 0, \tag{1.2}$$

$$\frac{\partial}{\partial x} \left(2M \frac{\partial v}{\partial x} - p + \theta \psi + \sigma \ln(1 - \theta) \right) = 0, \tag{1.3}$$

where $0 < \theta < 1$ is the volume fraction of cells and fibers, v is the velocity of the cell-fiber phase, p is the pressure, φ is the drag coefficient, ψ is the contractility coefficient, σ is the swelling coefficient, and M is the viscosity coefficient of the cell-fiber fraction. Note that the parameters φ , M, ψ and σ are nonnegative and can depend on time, space, and θ .

A reasonable range of dimensional and non-dimensional parameters are presented in Table 1; see [3,8] for the references.

parameter	symbol	units	range
specific drag	φ	kg/m ³ -sec	$10^{12} \sim 10^{14}$
coefficient			
tissue viscosity	М	kg/m-sec	10 ⁵
specific contractility	ψ	kg/m-sec ²	$10^3 \sim 10^4$
coefficient			
swelling number	σ	kg/m-sec ²	$10 \sim 10^{3}$
volume fraction of	θ_0	-	$\epsilon \sim (1 - \epsilon)$
cell-fiber phase			

Table 1: Expected ranges of parameter values in which ϵ is a small positive number.

The boundary conditions (BC) are given as follows

$$v(0,t) = v(L,t) = 0, \quad \frac{\partial\theta}{\partial x}(0,t) = \frac{\partial\theta}{\partial x}(L,t) = 0, \quad \frac{\partial p}{\partial x}(0,t) = \frac{\partial p}{\partial x}(L,t) = 0. \tag{1.4}$$

One way to model two adjacent tissues is to simply include them in the same equations and account for their different densities with θ . Thus a simple interaction between two tissues can be modeled with piecewise constant initial condition (see Fig. 1),

$$\theta(x,0) = \begin{cases} \theta_l, & \text{if } 0 \le x < x_1 \text{ or } x_2 < x \le L, \\ \theta_u, & \text{if } x_1 \le x \le x_2, \end{cases}$$
(1.5)

where we use θ_l for the smaller constant (lower), and θ_u for the larger constant (upper).

In this paper, we will focus on simulating tissue deformations numerically for the one dimensional model. Note that the mathematical model is a non-linear, mixed (hyperbolic and elliptic) system of differential equations. Shock waves will be developed in