

MECHANICAL EFFECTS COUPLED WITH CALCIUM WAVES

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Abstract

In the paper we find explicit formulae for heteroclinic travelling wave solutions in the system of equations describing the dynamics of cytosolic calcium concentration and the accompanying mechanical phenomena.

Key Words: *Calcium waves, reaction-diffusion systems, mechanochemical coupling*

1 Introduction

This paper considers explicit travelling wave solutions to a system of equations describing the evolution of the calcium propagation and the associated mechanical phenomena in biological media (cells and tissues). This system of equations has the following form:

$$\frac{\partial c}{\partial t} = D\nabla^2 c + f(c) + \gamma\theta, \quad (1)$$

$$\nabla \cdot \boldsymbol{\sigma} = 0, \quad (2)$$

where $\theta = \nabla \cdot \mathbf{u}$ is the dilation and \mathbf{u} is the displacement. The stress tensor $\boldsymbol{\sigma} \equiv \sigma_{ij}$, where

$$\sigma_{ij} = \theta\lambda\delta_{ij} + 2G\epsilon_{ij} + \nu_1\theta_{,t}\delta_{ij} + \nu_2\epsilon_{ij,t} + \tau_{ij}. \quad (3)$$

In Eq.(1) c denotes the free cytosolic calcium concentration, D is its effective diffusion coefficient, $f(c)$ is the function describing calcium transport into and out of the cytosol. In Eq.(3) ϵ_{ij} are the components of the deformation tensor $\boldsymbol{\epsilon} = 1/2(\nabla\mathbf{u} + \nabla\mathbf{u}^T)$, λ and G are the Lamé coefficients, whereas ν_1 and ν_2 are the viscosity coefficients. τ_{ij} are the components of so called active traction tensor $\boldsymbol{\tau}$ (resulting from the actin-myosin interaction). Below, we assume that $\boldsymbol{\tau}$ is a diagonal tensor of the form

$$\boldsymbol{\tau} = \text{diag}(\tau_{11}, \tau_{22}, \tau_{33}).$$

In Eq.(2), the inertial terms have been neglected, due to the fact that the considered mechanical phenomena (connected with the wave of calcium concentration) induce relatively slow motion of the medium. Also, the divergence of the stress tensor is assumed to be equal to zero. This means that there are no external constraints for the expansion or contraction of different parts of the medium. This assumption will be changed in the last section of the paper. For the review of various aspects of calcium dynamics see, e.g. [2]. The linear form of the mechanical term $\gamma\theta$ in Eq.(1) is postulated in [8].

The dynamics of the local calcium concentration inside a cell or tissue coordinates many physiological processes. It plays a key role in transferring signals from the surrounding

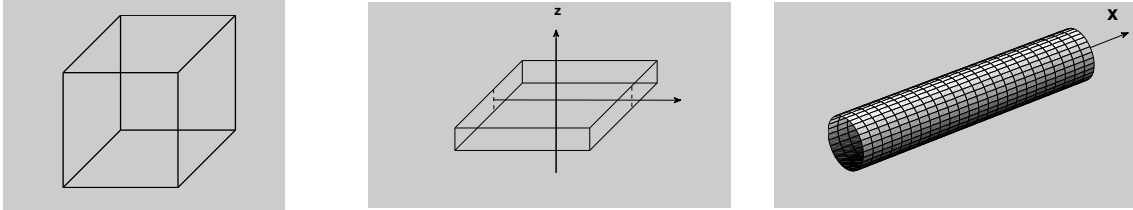


Figure 1: *The geometrical cases considered in the paper. From left to right: a. bulk medium (large in every direction) b. infinite plane layer of sufficiently small width $2d$ c. long cylinder of sufficiently small radius*

medium into the interior of the cell. It governs also the process of bridging between actin and myosin fiber proteins. This process is especially important in myocyte long cells and leads to their contraction. On the other hand, the local mechanical deformations of the medium influence the dynamics of the calcium through the term $\gamma\theta$ in Eq.(1). The explicit form for purely chemical travelling waves, i.e. for $\gamma = 0$ are well known (see e.g. [8]), however, up to our knowledge, the explicit travelling wave solutions to system (1)-(2) have not been studied.

In this paper we will find some explicit solutions of travelling wave type for system (1)-(2). This provides us some insight into the phenomenon of the mechano-chemical coupling described by this system of equations. We confine ourselves to three geometrical cases. These cases have been considered in [9] and [6] and are depicted on Fig. 1. Thus in the paper we consider the calcium waves in an unbounded bulk medium, thin infinite layers, which in their undeformed state are planes [1] and infinitely long cylinders of sufficiently small radius. The last situation can be physically realized e.g., in the case of long myocyte cells [7].

Remark Calcium dynamics is a complex phenomenon, consisting of numerous intracellular pathways and the exchange of calcium ions between extracellular matrix (ECM) and cell interior. However, in this paper we confine ourselves only to the processes related to the calcium activated, calcium release from endoplasmic reticulum vesicles into the cytoplasm. This simplifying assumption allows us to use a description based reaction-diffusion equation. \square

2 Analysis of the mechanical equation

For the reader's convenience we repeat here the analysis of the mechanical equation, which can be found in [9] or [6]. As we mentioned, we are interested in plane travelling waves

solutions to system (1)-(2) propagating along the x -axis. *As a result we may assume here that $c = c(x, t)$.*

Remark The last assumption is in fact a kind of homogenization assumption, by which we neglect the boundary effects or non-homogeneity of the internal endoplasmic calcium vessels distribution. \square

Bulk medium

In the case of bulk medium it is natural to assume that all the components of the deformation tensor ϵ depend only on x and t and that $\epsilon_{ij} \equiv 0$ except for ϵ_{11} . (Consequently, $u_2 \equiv 0$ and $u_3 \equiv 0$.) We thus have $\theta = \epsilon_{11}$ and the x -component of Eq.(2) gives

$$(\lambda\theta + 2G\theta + (\nu_1 + \nu_2)\theta_{,t} + \tau_{11})_{,x} = 0,$$

which after integrating and putting the integration constant to zero (assuming that there are no external forces) leads to the equation

$$\lambda\theta + 2G\theta + (\nu_1 + \nu_2)\theta_{,t} + \tau = 0,$$

with $\tau = \tau(c) = \tau_{11}(c)$.

Infinite layer

Now, let us consider the case of an infinite thin layer. We fix the system of coordinates in such a way that the x -axis is parallel to the layer and the z -axis is perpendicular to it (as in Fig. 1). As before, *we assume that all the components of the tensors σ and ϵ depend only on x and t , but do not depend on y and z .* According to the translational symmetry with respect to y , we can suppose that $u_2 \equiv 0$, hence $\epsilon_{22} \equiv 0$. We also demand the plane stress conditions on the boundary planes $\{(x, y, z) : z = \pm d\}$ (see [3]). That is to say, we suppose that for $z = \pm d$

$$\sigma_{i3} = 0, \quad i = 1, 2, 3. \tag{4}$$

Thus, in this case we obtain:

1. from the balance of mechanical forces:

$$(\lambda\theta + 2G\epsilon_{11} + \nu_1\theta_{,t} + \nu_2\epsilon_{11,t} + \tau_{11})_{,x} = 0 \tag{5}$$

2. from the boundary conditions on the boundary planes (by taking $i = 3$):

$$\lambda\theta + 2G\epsilon_{33} + \nu_1\theta_{,t} + \nu_2\epsilon_{33,t} + \tau_{33} = 0. \tag{6}$$

Though referring only to the boundary conditions at $z = \pm d$, Eq.(6) is valid in the whole of the thin layer, as we assume that the components of the tensor σ do not depend on the variable z . Integrating Eq.(5) and putting the integration constant zero we obtain

$$\sigma_{11} = \theta\lambda + 2G\epsilon_{11} + \nu_1\theta_{,t} + \nu_2\epsilon_{11,t} + \tau_{11} = 0, \tag{7}$$

hence by adding it to Eq.(6), we obtain the first order differential equation for the dilation θ

$$2(\lambda + G)\theta + (2\nu_1 + \nu_2)\theta_{,t} + \tau_{11} + \tau_{33} = 0. \tag{8}$$

Fibers

In the case of waves in fibers (e.g. long cells as myocytes) we assume cylindrical symmetry of the problem and that $\tau_{22} = \tau_{33}$. In this case we have $\epsilon_{22} = \epsilon_{33}$, so $\theta = \epsilon_{11} + 2\epsilon_{33}$. Formally the boundary conditions and the balance of forces have the same form as in the case of thin layers, then

$$(\lambda\theta + 2G\epsilon_{11} + \nu_1\theta_{,t} + \nu_2\epsilon_{11,t} + \tau_{11})_{,x} = 0$$

$$\lambda\theta + 2G\epsilon_{22} + \nu_1\theta_{,t} + \nu_2\epsilon_{22,t} + \tau_{22} = 0$$

$$\lambda\theta + 2G\epsilon_{33} + \nu_1\theta_{,t} + \nu_2\epsilon_{33,t} + \tau_{33} = 0$$

(We use the Cartesian system of coordinates keeping in mind however the radial symmetry.) Summing up these equations after the integration of the first one we obtain again a first order equation for the dilation θ

$$(3\lambda + 2G)\theta + (3\nu_1 + \nu_2)\theta_{,t} + \tau_{11} + 2\tau_{33} = 0.$$

Thus in all three cases we arrive at the same type of first order ODE. This equation has the form:

$$K\theta + \mu\theta_{,t} + \tau = 0. \tag{9}$$

The coefficients K , μ and τ depend on the case considered and τ is an appropriate function of the components of the tensor $\boldsymbol{\tau}$ and are *given explicitly* in Table 1.

Finally let us recall that the Young modulus E and Poisson ratio ν are related to the Lamé coefficients by the relations:

$$\lambda = \frac{E\nu}{(1+\nu)(1-2\nu)}, \quad G = \frac{E}{2(1+\nu)}$$

and

$$E = \frac{G(2G+3\lambda)}{\lambda+G}, \quad \nu = \frac{\lambda}{2(\lambda+G)}$$

3 Travelling wave solutions

Solutions of travelling wave type describe many important phenomena in biology [8], chemistry ([11]) and physics (e.g. different models of phase transitions [10]). Looking for travelling wave solutions we assume that

$$c(x, y, z, t) = c(x - vt), \quad \theta(x, y, z, t) = \theta(x - vt), \tag{10}$$

where v is the speed of the wave. Moreover, we assume that the displacement have also the form of a travelling wave in the x -direction, i.e.

$$\mathbf{u}(x, y, z, t) = \mathbf{u}(x - vt, y, z). \tag{11}$$

	$K(\lambda, G)$	$K(E, \nu)$	μ	τ
BULK MEDIUM	$\lambda + 2G$	$\frac{E(1-\nu)}{(1+\nu)(1-2\nu)}$	$\nu_1 + \nu_2$	τ_{11}
THIN LAYER	$2\lambda + 2G$	$\frac{E}{(1+\nu)(1-2\nu)}$	$2\nu_1 + \nu_2$	$\tau_{11} + \tau_{33}$
FIBER	$3\lambda + 2G$	$\frac{E}{1-2\nu}$	$3\nu_1 + \nu_2$	$\tau_{11} + 2\tau_{33}$

Table 1: The coefficients K , μ and τ in Eq.(9)

System (1)-(9) changes then to a system of ordinary differential equations of the form:

$$Dc'' + vc' + f(c) + \gamma\theta = 0. \quad (12)$$

$$-v\mu\theta' + K\theta + \tau = 0. \quad (13)$$

Here ' denotes the derivative with respect to $\xi = x - vt$. Thus, we are looking for heteroclinic solutions to system (12)-(13), that is to say $C^2(\mathbb{R}^1)$ functions c and θ such that $\lim_{\xi \rightarrow -\infty} c(\xi) = c_1$, $\lim_{\xi \rightarrow \infty} c(\xi) = c_3$ and $\lim_{|\xi| \rightarrow \infty} \theta(\xi) = 0$. Here c_1 and $c_3 > c_1$ denote the two stable equilibrium concentrations of the cytosolic calcium. When considering system (1)-(2), we also assume that

$$\tau(c_1) = \tau(c_3) = 0, \quad \text{and} \quad \tau(c) \geq 0. \quad (14)$$

The function $f(\cdot)$ is often modelled in the form $f(c) = A(c - c_1)(c - a)(c_3 - c)$ with appropriately chosen constants A , c_1 , c_3 and a . For considerations concerning the physical values of A and the effective diffusion coefficient D see e.g., [5] or [7]. Obviously, due the possibility of appropriate scaling, we can assume without losing generality that $c_1 = 0$ and $c_3 = 1$.

In the whole of the paper we will take the following assumptions.

Assumption 1. $f(c) = Ac(c - a)(1 - c)$ and $(1 - 2a) > 0$. □

Assumption 2. *The coefficients γ , λ , G , ν_1 , ν_2 are constants, whereas $\tau = \tau(c)$.* □

Remark The condition demanding that λ , G , ν_1 , ν_2 are constants can be relaxed to the condition that λ , G , ν_1 , ν_2 are appropriate functions of c . However, for clarity of exposition, we will not consider this generalization here. □

Remark concerning the form of $\tau(c)$ As we are not able to find explicit solutions in general, the form of $\tau = \tau(c)$ will be somehow adjusted to the form of the function θ . The precise form of $\tau = \tau(c)$ can depend on the kind of the tissue. It should be however positive for all c and must vanish for large c . In the paper, for sufficiently small relative values of the viscosity coefficient μ , $\tau = \tau(c)$ behaves approximately as $c(1 - c)$, so vanishes for $c = 0$ (as on the left panel of Fig.3). Only in the last section $\tau(0) > 0$ (as on the right panel of Fig.3). This agrees with the qualitative characterization of the traction terms, e.g. in [8]. □

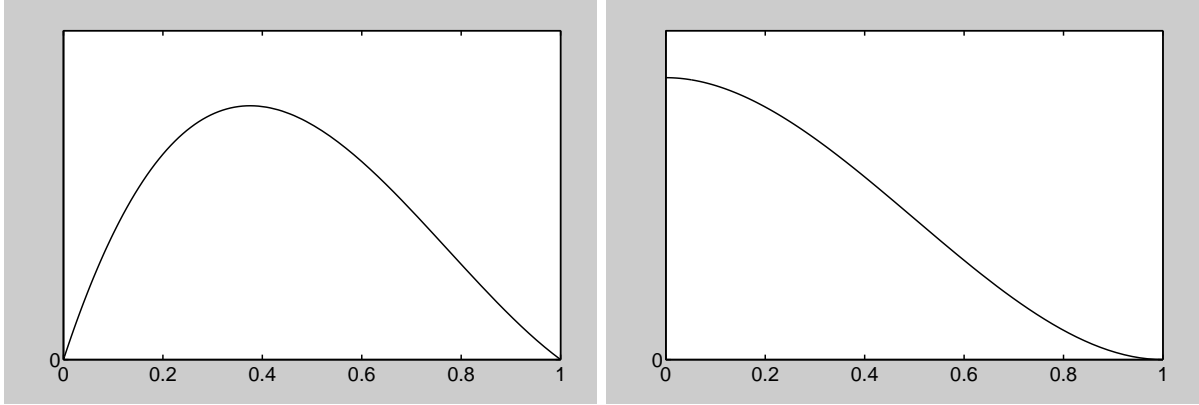


Figure 2: *Left: The qualitative shape of τ as a function of $c \in [0, 1]$ considered in section 3. Right: The shape of the function τ considered in section 4.*

It is known that, for $\theta = 0$, Eq.(12) has a heteroclinic solution connecting its constant steady states $c = 0$ and $c = 1$ for one and only one value of the parameter v equal to

$$v = -(AD/2)^{1/2}(1 - 2a).$$

The solution is of the form:

$$c(\xi) = \frac{1}{1 + \exp(-(\frac{A}{2D})^{1/2}\xi)}.$$

We prove that under some assumption about the form of τ a similar function satisfies the considered system. Let us make an ansatz:

$$c = \frac{1}{1 + \exp(-s\xi)}, \quad (15)$$

with $s \geq 0$. This function satisfies the identity:

$$c' = sc(1 - c) \quad (16)$$

and $c(0) = 1/2$. Next, let us suppose that

$$\theta(\xi) = -qc(\xi)(1 - c(\xi)) \quad (17)$$

for some positive $q \in \mathbb{R}^1$. Then

$$\theta' = -sqc(1-c)(1-2c) \quad (18)$$

Putting (17) into Eq.(12) determining the profile of the wave, we obtain

$$Dc'' + vc' + Ac(c-a)(1-c) - \gamma qc(1-c) = 0$$

or equivalently as

$$Dc'' + vc' + Ac(c-a + \gamma qA^{-1})(1-c) = 0.$$

This equation has a heteroclinic solution

$$c(\xi) = \frac{1}{1 + \exp(-(\frac{A}{2D})^{1/2}\xi)} \quad (19)$$

satisfying the condition $c(0) = 1/2$ **iff**

$$v = -(AD/2)^{1/2} (1 - 2(a + \gamma qA^{-1})). \quad (20)$$

This means in particular that $s = \sqrt{\frac{A}{2D}}$. According to (15), (19) and (20),

$$vs = -\frac{A}{2} (1 - 2(a + \gamma qA^{-1})). \quad (21)$$

Assumption 3. Let for τ defined in Table 1, $\tau = \tau(c)$ be such that the identity

$$qc(1-c) \left[K + \frac{A}{2} (1 - 2(a + \gamma qA^{-1})) \mu(1-2c) \right] = \tau(c) \quad (22)$$

is satisfied (with μ given in Table 1). \square

According to (21), the left hand side of (22) can be written as $qc(1-c) [K - vs\mu(1-2c)]$, so Eq.(13) is fulfilled.

Displacements

Bulk medium

In this case we have $\mathbf{u} \equiv (u_1, 0, 0)$ and $u_1(x, y, z, t) = \int_{-\infty}^{\xi} \theta(h) dh$, so demanding that $u_1 \rightarrow 0$ for $x - vt \rightarrow -\infty$, we have according to (18):

$$u_1 = -qc(x, y, z, t) \sqrt{\frac{2D}{A}}.$$

Infinite layer

This case is described by Eq.(8). For simplicity, we also assume that

$$\tau_{11} = \tau_{33}.$$

Then by (8)

$$(\lambda + G)\theta + (\nu_1 + \nu_2/2)\theta_{,t} + \tau_{11} = 0 \quad (23)$$

whereas from (7)

$$\sigma_{11} = \theta\lambda + 2G\epsilon_{11} + \nu_1\theta_{,t} + \nu_2\epsilon_{11,t} + \tau_{11} = 0. \quad (24)$$

From (24) we obtain:

$$\sigma_{11} = \theta(\lambda + G) + 2G\epsilon_{11} + (\nu_1 + \frac{\nu_2}{2})\theta_{,t} + [-G\theta - \frac{\nu_2}{2}\theta_{,t}] + \nu_2\epsilon_{11,t} + \tau_{11} = 0$$

and finally by subtracting (24) we get

$$2G\epsilon_{11} + [-G\theta - \frac{\nu_2}{2}\theta_{,t}] + \nu_2\epsilon_{11,t} = 0. \quad (25)$$

Similarly, using (8) and (6) one obtains

$$2G\epsilon_{33} + [-G\theta - \frac{\nu_2}{2}\theta_{,t}] + \nu_2\epsilon_{33,t} = 0. \quad (26)$$

Neglecting the viscosity coefficient by the time derivative of ϵ_{11} in Eq.(25) yields:

$$2G\epsilon_{11} = G\theta + \frac{\nu_2}{2}\theta_{,t}.$$

By assumption, $G\theta + \frac{\nu_2}{2}\theta_{,t} = -Gqc(1-c) - v\frac{\nu_2}{2}\theta' = \left(-G\frac{q}{s}c - v\frac{\nu_2}{2}\theta\right)'$. Using (11), we thus have

$$\begin{aligned} 2Gu_1(x, y, z, t) &= \int_{-\infty}^{\xi} \epsilon_{11}(h)dh = -G\frac{q}{s}c(\xi) - v\frac{\nu_2}{2}\theta(\xi) = \\ &= -G\frac{q}{s}c(\xi) - vq\frac{\nu_2}{2}c(\xi)(1-c(\xi)), \end{aligned}$$

so

$$u_1(x, y, z, t) = -qc(\xi)\sqrt{\frac{D}{2A}} \left[1 - A\frac{\nu_2}{2G}(1 - 2(a + \gamma qA^{-1}))(1 - c(\xi))\right].$$

The integration constant has been taken to equal to zero to assure that $u_1(-\infty) = 0$.

Now, let us analyze Eq.(26). By means of (18) we have

$$G\theta + \frac{\nu_2}{2}\theta_{,t} = -Gqc(1-c) - v\frac{\nu_2}{2}\theta' = -Gqc(1-c) + vs\frac{\nu_2}{2}qc(1-c)(1-2c),$$

so neglecting the viscosity coefficient by the time derivative of ϵ_{33} and using the fact that, by assumption, ϵ_{33} does not depend on z , we obtain by means of (21)

$$u_3(x, y, z, t) = -qc(\xi)(1-c(\xi)) \left[\frac{1}{2} + \frac{A}{2G}(1 - 2(a + \gamma q))\frac{\nu_2}{2}(1 - 2c(\xi))\right] z. \quad (27)$$

Similar calculations can be made for fibers and the bulk medium (with respect to the displacements in the x -direction). In the case of fibers (infinite cylinders) we use the fact of the radial symmetry, from which follows that the dependence of the radial displacement on the radius r is the same as the dependence of u_3 on z at the z -axis. That is why the displacement along the radius is equal to $\epsilon_{33}\mathbf{r}$. The qualitative behaviour of the displacements u_1 and u_3 (in the case of thin layers and fibers) are shown on Fig. 3 and Fig. 4 respectively.

Remarks

1. Eq.(25) can be solved explicitly also with $\nu_2 \neq 0$. However, the obtained solution would have much more complicated form. It is proved in [5] that for $\nu_2 \rightarrow 0$ this solution tends in $C^1(\mathbb{R})$ -norm to the solution of Eq.(25) with $\nu_2 = 0$. This reasoning is justified by the fact that in most of biological tissues the viscosity effects for displacements propagating with the speeds characteristic for cytosolic calcium waves ($10 - 100 \mu m/s$) are small with respect to elastic effects ([4],[8]).

2. As it follows from (27), $\epsilon_{13} \neq 0$ with $\epsilon_{13} = O(d)$. However, since the displacement vector is expressed up to linear terms in z , it is reasonable to take into account the strain tensor up to zero order terms in z as differentiation lowers the order of approximation by 1 (see [5]). In this sense the assumed condition that the components of σ and ϵ depend only on x is satisfied. Thus asymptotically as $d \rightarrow 0$, we have $u_3 \equiv 0$, whereas the expression (27) can be viewed as a first order perturbation. \square

We have thus shown the validity of the following theorem.

THEOREM 1. *Let Assumptions (1), (2) and (3) be satisfied. Then for all $q > 0$, system (12),(13) has a heteroclinic solution (v, c, θ) with v is given by (20), $c(\xi)$ by (19) and $\theta(\xi)$ by (17). The solution is unique up to a translation in ξ . \square*

The functions $c(x - vt)$ and $\theta(x - vt)$, together with the corresponding displacement functions \mathbf{u} (which were found above), satisfy the initial PDE system (1)-(3) exactly in the case of bulk unbounded medium. In the case of thin infinite layer and fibers, under the assumption of plane stress conditions on the boundary, system (1)-(3) is satisfied up to terms of order $O(d)$ as $d \rightarrow 0$.

4 Mechanochemical travelling waves with mechanical constraints

In this section we consider the travelling wave solutions of the system

$$\frac{\partial c}{\partial t} = D_c \nabla^2 c + f(c) + \gamma \theta, \quad (28)$$

$$\nabla \cdot \boldsymbol{\sigma} = k \mathbf{u}, \quad (29)$$

where $k = \text{const} > 0$. That is to say, we take into account the possibility of mechanical constraints which can counteract the displacements of the medium. The specific form of this constraint given by the right-hand side of Eq.(29) is called the Winkler model. We use the same methodology as in the previous sections. To be more precise, we exploit the explicit solution of the form (15) by appropriate choice of the function $\tau(c)$. For definiteness, we confine ourselves to the case of *bulk medium*. By the considerations of section 2, in the case a plane travelling wave solution propagating along the x -axis in the bulk medium, we can assume that $\theta = \epsilon_{11}$ and $\mathbf{u} = (u_1, 0, 0) =: (u, 0, 0)$. It follows that

$$\theta = u_{,x} = u',$$

where $'$ denotes the differentiation with respect to $\xi = x - vt$. Unlike the previous section, we demand that $u(\xi) \rightarrow 0$ as $\xi \rightarrow \pm\infty$ by assuming

$$u(\xi) = -\zeta c'(\xi).$$

It follows that $\theta(\xi) = u'(\xi) = -\zeta c''(\xi)$. If we suppose that $c(\xi)$ is given by (15), then

$$\theta = -s\zeta c(1-c)(1-2c) \quad (30)$$

and θ' is a fourth order polynomial in c vanishing for $c = 0$ and $c = 1$. Eq.(29) can thus be written as

$$v\mu\theta' + Ks\zeta c(1-c)(1-2c) - k\zeta c + \eta = \tau(c). \quad (31)$$

where η is an integration constant. Obviously, for $\eta > 0$ sufficiently large, $\tau(c) \geq 0$ for $c \in [0, 1]$ and $\tau(1) = 0$. Moreover, it follows from the implicit function theorem that for k sufficiently large with respect to K and μ sufficiently small, we can choose the constant $\eta > 0$ so large that $\tau(c) \geq 0$ for $c \in [0, 1]$ and $\tau(1) = 0$. An example of such a graph is depicted in the right panel of Fig. 2. Putting the form of θ into Eq.(28) we arrive, by using (30), at the equation

$$Dc'' + vc' + Ac(c-a)(1-c) - \gamma s\zeta c(1-c)(1-2c) = 0.$$

Hence, for $\tilde{\zeta} = \zeta/A$,

$$Dc'' + vc' + A(1+2\gamma\zeta s)c(1-c)(c - (a + \gamma\tilde{\zeta}s)(1+2\gamma\tilde{\zeta}s)^{-1}) = 0.$$

To calculate s we have to solve the equation

$$s = \left(\frac{A(1+2\gamma\tilde{\zeta}s)}{2D} \right)^{1/2}$$

which implies

$$s = \frac{A\gamma\tilde{\zeta} + \sqrt{A(2D + A\gamma^2\tilde{\zeta}^2)}}{2D} = \frac{\gamma\zeta}{2D} + \sqrt{\frac{A}{2D} + \frac{\gamma^2\zeta^2}{4D^2}}.$$

Having the parameter s , we can calculate the speed of the wave. Thus

$$v = -\sqrt{\frac{(A+2\gamma\zeta s)D}{2}} \left(1 - (a + \gamma\tilde{\zeta}s)(1+2\gamma\tilde{\zeta}s)^{-1} \right),$$

which can be written as

$$v = -Ds \left(1 - \frac{2a + 2\gamma\zeta s/A}{1 + 2\gamma\zeta s/A} \right).$$

This expression in its unfolded form is rather complicated. Asymptotically, for very large k , we have $\eta \cong \tau(0)$ and $\zeta \cong \tau(0)/k$, and the influence of mechanics on the speed of the wave is very small.

The graphs of displacement (u_1 - smooth line) and the dilation (θ - circled line) versus the coordinate x are given on Fig. 5.

5 Conclusions

In the paper we found explicit formulae for the travelling wave profiles as well as their speeds in a model describing the dynamics of cytosolic calcium and the accompanying mechanical effects under some simplifying assumptions concerning the form of the traction $\tau(c)$ and the cubic-like source term $f(c)$. The meaning of this result is twofold: firstly, the explicit solution can provide us some insight into the phenomena of mechano-chemical coupling, secondly, they can serve as a starting point of the analysis of more general problems.

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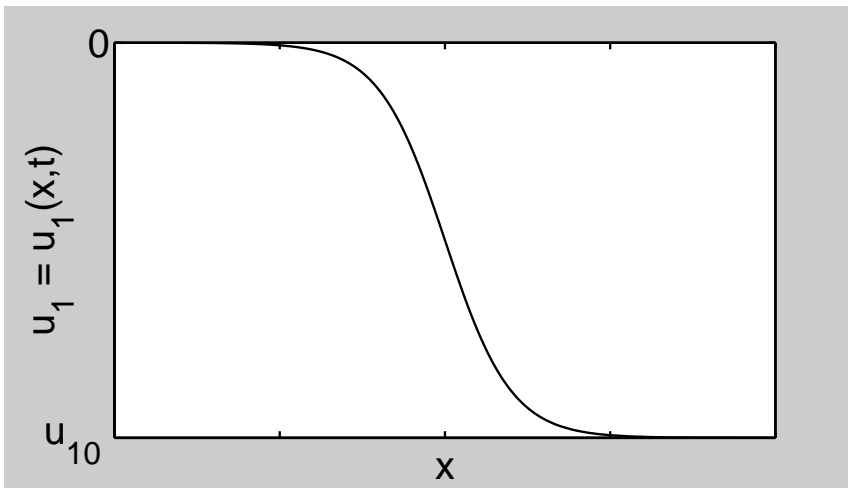


Figure 3: Displacements in the x -directions. The displacement vectors $u_1(x,t)$ are directed to the left. $u_1(x,t) \rightarrow 0$ as $x \rightarrow -\infty$ and $u_1(x,t) \rightarrow u_{10} < 0$ as $x \rightarrow \infty$.

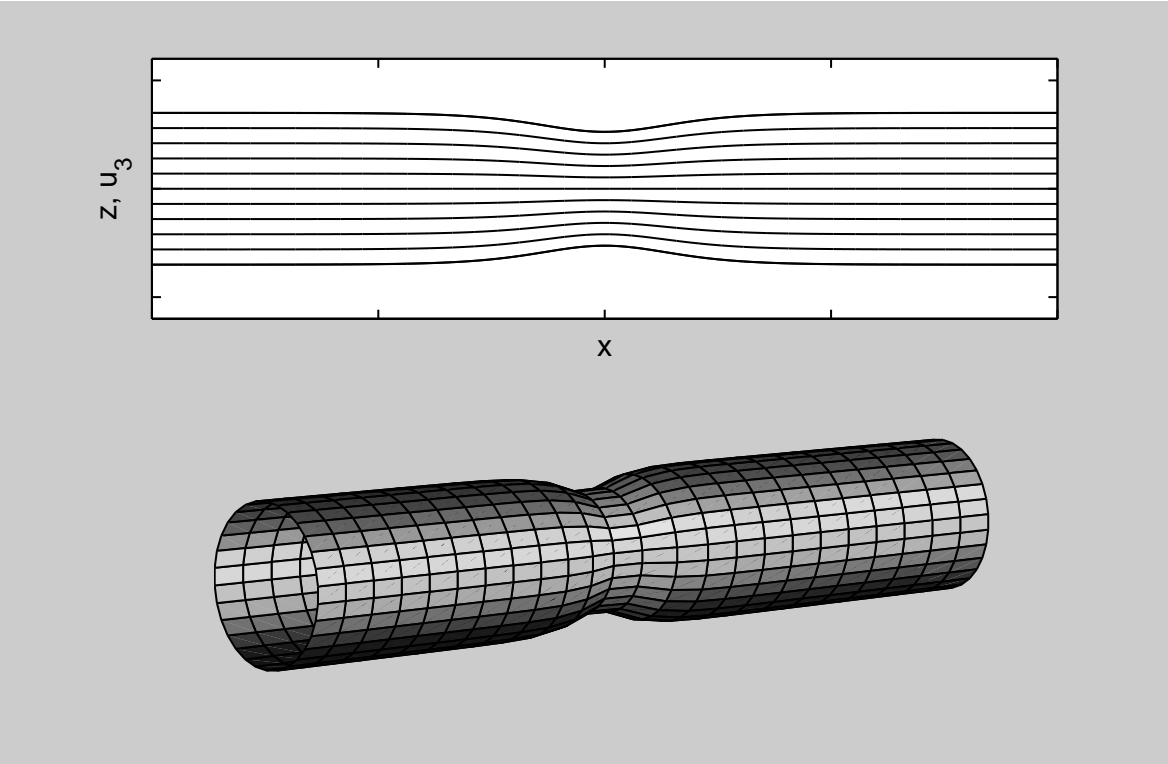


Figure 4: *Displacements in the z-direction. Upper panel: thin layer. Lower panel: long cylinder.*

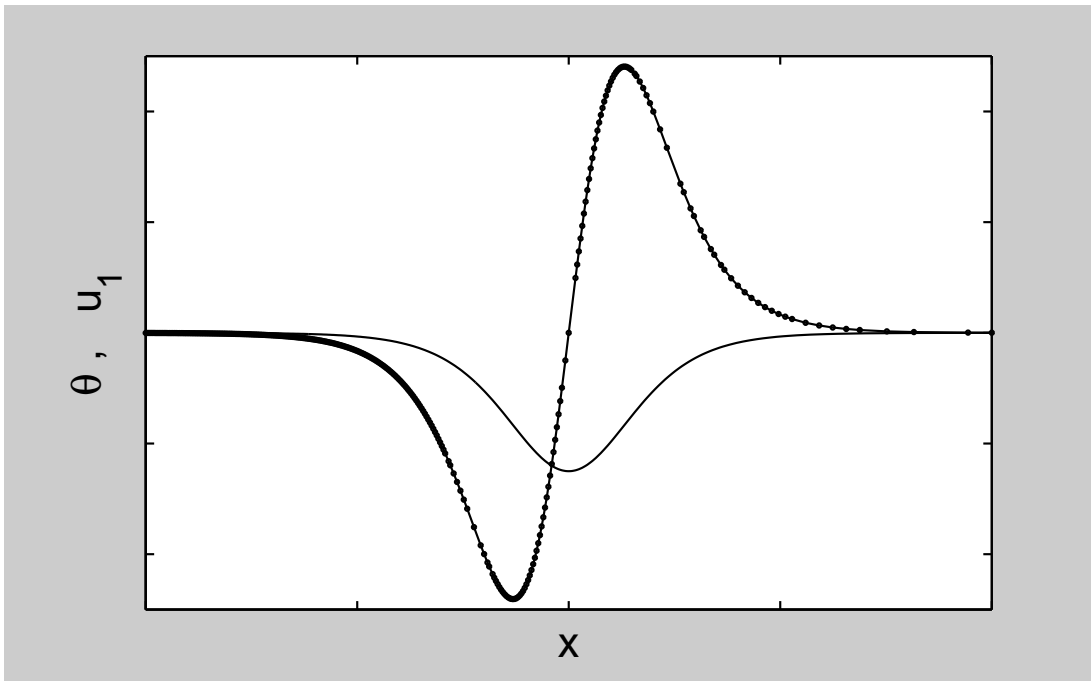


Figure 5: Displacement u_1 (smooth line) and the dilation θ (circled line) for system (28)-(29).