

Cell scale modeling of electropermeabilization by periodic pulses.

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Abstract: In this paper, we focus on the behaviour of periodic solutions to a cell- scale electroporabilization model previously proposed by Kavian et al. (JMB,2012). Since clinical permeabilization protocols mostly submit cancer cells to trains of periodic pulses, we investigate on parameters that modify significantly the resulting permeabilization. Theoretical results of existence and uniqueness of periodic solutions are presented, for two different models of membrane electric conductivity. Numerical simulations were performed to corroborate these results and illustrate the asymptotic convergence to periodic solutions, as well as the dependency on biological parameters such as the cell size and the extracellular conductivity.

Key-words: Cell modeling, Non-linear partial differential equations

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Modélisation de l'électroperméabilisation par pulses périodiques à l'échelle cellulaire.

Résumé : Dans ce papier, nous nous intéressons au comportement des solutions périodiques d'un modèle d'électroperméabilisation précédemment proposé par Kavian et al. (JMB,2012). La plupart des protocoles de perméabilisation soumettent les tumeurs à des trains de pulses périodiques, et nous souhaitons déterminer les paramètres qui modifient significativement la perméabilité résultante. Des résultats théoriques d'existence et d'unicité des solutions périodiques sont donnés, pour différents modèles de conductivité. Des simulations numériques ont été effectuées pour appuyer ces résultats et mettre en évidence la convergence asymptotique vers des solutions périodiques, ainsi que la dépendance des paramètres biologiques que sont la taille des cellules et la conductivité extracellulaire.

Mots-clés : Modélisation bio-cellulaire, Equations aux dérivées partielles non linéaires

1 Introduction

Electropermeabilization or *electroporation* is a phenomenon that occurs when a biological cell or a lipid vesicle is submitted to a high electric field. If a sufficiently large potential difference is applied to the membrane, its structure is altered and molecules that are usually not able to enter the cytoplasm can diffuse inside the cell. Reversible electropermeabilization is already used to improve delivery of drugs such as bleomycin in oncology [1, 4], and also to make the cell permeable to very large molecules for gene transfer [7].

This paper is an extent of a previous modeling work [5] in which we proposed a phenomenological approach of electropermeabilization, compared to the previous most achieved models of Neu, Debruin and Krassowska [9, 3].

Experiments show that a single pulse is often not sufficient to achieve cell electropermeabilization, with a wide variety of pulses, from nanopulses to millipulses [8, 12, 11]. The applied pulses are usually trains of several pulses, and can be considered as a periodic source. Here we focus theoretically and numerically on the behaviour of solutions to our equations in this particular case of sources. The objective of this paper is to show that the membrane conductivity reaches a stable periodic state when submitted to a periodic voltage.

When very intense electric pulses are clinically applied, healthy tissues in the vicinity of the electrodes are completely damaged due to the high electric field. Moreover, thermal effects induce cell destruction if pulses last for too long. It is therefore important to determine pulse characteristics (voltage, duration) that are sufficient to permeabilize the targeted tissue, minimizing at the same time the caused damage. Considering that, we will focus on the amount of time needed for the cell to reach a periodic conductive state, depending on which model is chosen to describe the membrane conductivity.

We will show, in particular, that using a static model of conductivity results in a high convergence speed of the solution to a periodic state. A first conclusion should be that a few pulses are needed to reach it. However, results given by a fully dynamical model infer that more pulses are required to actually achieve electropermeabilization.

After recalling in the first section the model on the electric potential in a biological cell, and especially across its membrane, we will justify the existence and uniqueness of periodic solutions in the second section. In the final section, we will illustrate how a cell at rest submitted to a periodic source reaches a periodic state. We will validate these results with numerical simulations.

2 Statement of the model

2.1 The electric potential in a biological cell

A biological cell is considered as an homogeneous medium \mathcal{O}_c , which is separated from an exterior bath \mathcal{O}_e by a phospholipidic membrane Γ (see fig. 1). Due to the high resistivity and the thickness of the membrane, it is considered as a single surface interface between the two domains. The capacity of this electric material is denoted by C_m , and its surface conductivity by S_m . Let σ be the conductivity of the medium, considered piecewise constant:

$$\sigma = \begin{cases} \sigma_e & \text{in the bath } \mathcal{O}_e, \\ \sigma_c & \text{in the cytoplasm } \mathcal{O}_c. \end{cases} \quad (1)$$

The electric potential U in both intra- and extracellular domains follows Poisson's law:

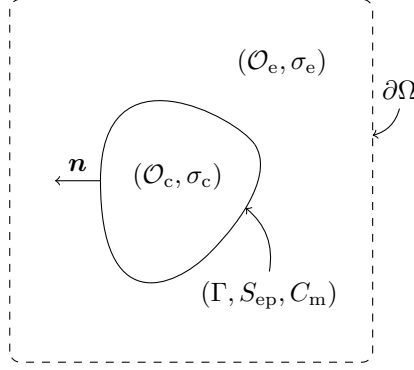


Figure 1: Geometry of the problem. The whole domain Ω is defined by $\Omega = \mathcal{O}_e \cup \overline{\mathcal{O}_c}$.

$$\Delta U = 0 \quad \text{in } \mathcal{O}_e \cup \mathcal{O}_c, \quad (2a)$$

$$U = g \quad \text{on } \partial\Omega, \quad \forall t > 0, \quad (2b)$$

with the transmission conditions on the flux and the transmembrane potential across the interface Γ :

$$[\sigma \partial_{\mathbf{n}} U]_{\Gamma} = 0, \quad (2c)$$

$$C_m \partial_t [U]_{\Gamma} + I_{ep} = \sigma_c \partial_{\mathbf{n}} U|_{\Gamma^-}, \quad (2d)$$

where g represents the potential that is applied by external electrodes, and I_{ep} is the total current due to electropermeabilization. In the mostly used models of electropermeabilization proposed by Neu, Krassowska and De Bruin [9, 3], this current was defined as the product of a pore density N_{ep} by the current flowing through a single pore i_{ep} . The pore current was defined by a highly non-linear function of the transmembrane potential.

Derived from a linearization of this total current I_{ep} , we proposed in our previous work [5] to replace this definition by the product of the surface conductivity of the membrane S_m by the transmembrane potential difference $[U]_{\Gamma}$. The transmission condition now holds :

$$C_m \partial_t [U]_{\Gamma} + S_m [U]_{\Gamma} = \sigma_c \partial_{\mathbf{n}} U|_{\Gamma^-}. \quad (3)$$

In order to take into account electropermeabilization, we write S_m as the sum of the lipid conductivity at rest S_0 and a non-linear part S_{ep} depending on $[U]_{\Gamma}$:

$$S_m([U]_{\Gamma}) := S_0 + S_{ep}([U]_{\Gamma}). \quad (4)$$

The next section introduces the different conductivity models we chose.

2.2 Models of membrane conductivity

Throughout this paper, we will focus on two models of membrane conductivity:

- a static conductivity model, which has a sigmoid profile:

$$\forall \lambda \in \mathbb{R}, S_{ep}(\lambda) = S_1 \beta(\lambda), \quad \text{with } \beta(\lambda) = (1 + \tanh(k_{ep}(|\lambda| - V_{rev}))) / 2 \quad (5)$$

where S_1 is another conductivity constant, much larger than S_0 . k_{ep} defines the speed of the switch between the rest state ($S_m = S_0$) and the fully permeabilized state ($S_m = S_0 + S_1$). V_{rev} designates the voltage threshold that needs to be crossed to permeabilize the membrane.

We will refer to this model as the β -model.

- a dynamical model which will be called X -model: for $t \geq 0$,

$$S_{ep}(t, \lambda) = S_1 X(t, \lambda), \quad \text{with} \quad \begin{cases} \partial_t X([U(t)], t) = \frac{\beta([U(t)]) - X(t)}{\tau_{ep}}, \\ X(t = 0) = X_0 \in [0, 1]. \end{cases} \quad (6)$$

where β is the same function as the β -model and τ_{ep} is the characteristic time constant of formation and resealing of pores. The X variable follows a *sliding-door* model around the electropermeabilization threshold V_{rev} . Using this model makes the dynamic of electropermeabilization intrinsic to the cell and not dependent of the applied pulse. We showed in [5] that X verifies:

$$\begin{cases} X \in C([0, +\infty)), \\ 0 \leq X(t, \lambda) \leq 1 \quad \forall t \geq 0. \end{cases} \quad (7)$$

In addition, we showed that there exists $K > 0$ such that:

$$\forall t \geq 0, \quad |\lambda_1 X(t, \lambda_1) - \lambda_2 X(t, \lambda_2)| \leq K |\lambda_1 - \lambda_2|. \quad (8)$$

All these models fall in the hypothesis of theorem 10 in [5], so existence and uniqueness of solutions to problem (2) holds.

The main topic is now to know if given a periodic source term g , there exists a unique solution that is also periodic. Also, starting from any initial condition, does the solution converge to the periodic state, and at which speed ?

3 Existence and uniqueness of periodic solutions

Notation For any functionnal space H and $T > 0$, we will designate by $C_T(H)$ the subspace of functions of $C([0, +\infty), H)$ that are T -periodic :

$$C_T(H) = \{f \in C([0, +\infty), H), \forall t \geq 0, f(t + T, \cdot) = f(t, \cdot)\}$$

And let

$$PH^s(\Omega) = \{f \in L^2(\Omega), f|_{\mathcal{O}_e} \in H^s(\mathcal{O}_e), f|_{\mathcal{O}_c} \in H^s(\mathcal{O}_c)\}.$$

This section will be dedicated to the proof of the following theorem :

Theorem 1.

Let $T > 0$ and g be a T -periodic function of $C_T(H^1(\partial\Omega)) \cap W^{1,1}((0, +\infty), L^2(\partial\Omega))$. There exists a unique T -periodic solution U to the following problem:

$$\begin{cases} U \in C_T(PH^1(\Omega)), \\ \Delta U = 0 & \text{in } \mathcal{O}_e \cup \mathcal{O}_c, \\ U = g & \text{in } \partial\Omega, \forall t \geq 0, \\ [\sigma \partial_{\mathbf{n}} U]_{\Gamma} = 0, \\ C_m \partial_t [U]_{\Gamma} + S_0 [U]_{\Gamma} + S_{\text{ep}}(t, [U]_{\Gamma}) [U]_{\Gamma} = \sigma_c \partial_{\mathbf{n}} U|_{\Gamma-} \end{cases} \quad (9)$$

Also, let $V_0 \in PH^1(\Omega)$ and V be the solution to

$$\begin{cases} V \in C([0, +\infty), PH^1(\Omega)), \\ \Delta V = 0 & \text{in } \mathcal{O}_e \cup \mathcal{O}_c, \\ V = g & \text{on } \partial\Omega, \forall t \geq 0, \\ V(t=0, \cdot) = V_0(\cdot) & \text{in } \Omega, \\ [\sigma \partial_{\mathbf{n}} V]_{\Gamma} = 0, \\ C_m \partial_t [V]_{\Gamma} + S_0 [U]_{\Gamma} + S_{\text{ep}}(t, [V]_{\Gamma}) [V]_{\Gamma} = \sigma_c \partial_{\mathbf{n}} V|_{\Gamma-} \end{cases} \quad (10)$$

then, for all $t \geq 0$,

$$\|[U(t)]_{\Gamma} - [V(t)]_{\Gamma}\|_{L^2(\Gamma)}^2 \leq e^{-\frac{C}{C_m} t} \|[U_0]_{\Gamma} - [V_0]_{\Gamma}\|_{L^2(\Gamma)}^2, \quad (11)$$

where C is a real constant.

Proof: Existence and uniqueness of the solution V is given by [5]. In order to prove existence and uniqueness of periodic solutions to problem (9), we introduce the Dirichlet-to-Neumann (or Steklov-Poincaré) operators Λ_c and Λ_e on the interface for the Laplacian. Denote by ν_c (resp. ν_e) the unitary normal to Γ directed from the inside to the outside of \mathcal{O}_c (resp. \mathcal{O}_e). Λ_c and Λ_e are defined by:

$$\Lambda_c : \begin{cases} H^1(\Gamma) \rightarrow L^2(\Gamma) \\ f \mapsto \nu_c \cdot \sigma_c \nabla U_c|_{\Gamma-} \end{cases} \quad \text{where } \begin{cases} \nabla \cdot (\sigma_c \nabla U_c) = 0 \text{ in } \mathcal{O}_c, \\ U_c|_{\Gamma} = f. \end{cases} \quad (12a)$$

$$\Lambda_e : \begin{cases} H^1(\Gamma) \rightarrow L^2(\Gamma) \\ f \mapsto \nu_e \cdot \sigma_e \nabla U_e|_{\Gamma+} \end{cases} \quad \text{where } \begin{cases} \nabla \cdot (\sigma_e \nabla U_e) = 0 \text{ in } \mathcal{O}_e, \\ U_e|_{\partial\Omega} = 0, \\ U_e|_{\Gamma} = f. \end{cases} \quad (12b)$$

Using these operators, the transmission condition (2d) of the transmembrane potential can be rewritten on the manifold Γ . For the sake of readability, we now denote by u the transmembrane potential difference $[U]_{\Gamma}$ across the membrane. The condition now reads:

$$C_m \partial_t u + \Lambda_c (\text{Id} + \Lambda_e^{-1} \Lambda_c)^{-1} u + S_0 u + S_{\text{ep}}(u) u = G, \quad (13)$$

where $G = \Lambda_c (\text{Id} + \Lambda_e^{-1} \Lambda_c)^{-1} \Lambda_e^{-1} \Lambda_0 g$ is the external source term which has been brought to

the interface *via* another Steklov-Poincaré operator:

$$\Lambda_0 : \begin{cases} H^1(\partial\Omega) & \rightarrow L^2(\Gamma) \\ g & \mapsto \nu_e \cdot \sigma_e \nabla U_e|_{\Gamma^+} \end{cases} \quad \text{where} \quad \begin{cases} \nabla \cdot (\sigma_e \nabla U_e) = 0 \text{ in } \mathcal{O}_e, \\ U_e|_{\partial\Omega} = g, \\ U_e|_{\Gamma} = 0. \end{cases} \quad (14)$$

Let $\mathcal{A} := \Lambda_c(\text{Id} + \Lambda_c^{-1}\Lambda_c)^{-1}$. In [5], we have already proven that the operator $(H^1(\Gamma), \mathcal{A})$ is m-accretive.

In order to prove the existence of periodic solutions, it is necessary to verify firstly the boundedness of the solution u to the following problem, equivalent of (2):

$$u \in C([0, +\infty), H^1(\Gamma)), \quad (15a)$$

$$C_m \partial_t u + \mathcal{A}u + S_0 u + S_{\text{ep}}(t, u)u = G, \text{ on } \Gamma, \forall t > 0 \quad (15b)$$

$$u(t = 0, \cdot) = u_0(\cdot) \text{ on } \Gamma. \quad (15c)$$

where $G \in C([0, +\infty), H^1(\Gamma))$ is not necessarily T -periodic, $u_0 \in H^1(\Gamma)$, \mathcal{A} is a m-accretive operator and S_{ep} the conductivities previously defined in eqs. (5) to (6). We have, for all $t > 0$,

$$\langle C_m \partial_t u, u \rangle + \langle \mathcal{A}u, u \rangle + S_0 \langle u, u \rangle + \langle S_{\text{ep}}(t, u)u, u \rangle = \langle G, u \rangle. \quad (16)$$

It is clear that for all conductivity models $\langle S_{\text{ep}}(t, u)u, u \rangle$ is positive. Hence thanks to Young's inequality:

$$\frac{1}{2} C_m \partial_t \|u\|_{L^2(\Gamma)}^2 + \langle \mathcal{A}u, u \rangle + S_0 \|u\|_{L^2(\Gamma)}^2 \leq \frac{\|u\|_{L^2(\Gamma)}^2}{2\alpha} + \frac{\alpha G_\infty^2}{2}$$

for any $\alpha > 0$, with $G_\infty := \max_{t \in [0, T]} \|G(t)\|_{L^2(\Gamma)}$. Since \mathcal{A} is m-accretive,

$$\frac{1}{2} C_m \partial_t \|u\|_{L^2(\Gamma)}^2 + S_0 \|u\|_{L^2(\Gamma)}^2 \leq \frac{\|u\|_{L^2(\Gamma)}^2}{2\alpha} + \frac{\alpha G_\infty^2}{2}.$$

We can now invoke Gronwall's lemma to prove boundedness of solutions in finite times: $\forall t > 0$,

$$\begin{aligned} \|u(t)\|_{L^2(\Gamma)}^2 &\leq \|u_0\|_{L^2(\Gamma)}^2 \exp\left(-\frac{2\alpha S_0 - 1}{\alpha C_m} t\right) + \frac{\alpha G_\infty^2}{C_m} \int_0^t \exp\left(-\frac{2\alpha S_0 - 1}{\alpha C_m} (t-s)\right) ds, \\ &\leq \|u_0\|_{L^2(\Gamma)}^2 \exp\left(-\frac{2\alpha S_0 - 1}{\alpha C_m} t\right) + \frac{\alpha^2 G_\infty^2}{2\alpha S_0 - 1} \left(1 - \exp\left(-\frac{2\alpha S_0 - 1}{\alpha C_m} t\right)\right). \end{aligned}$$

Setting $\alpha = \frac{1}{S_0}$ minimizes the coefficient $\frac{\alpha^2}{2\alpha S_0 - 1}$. Therefore,

$$\forall t > 0, \|u(t)\|_{L^2(\Gamma)}^2 \leq \|u_0\|^2 \exp\left(-\frac{S_0}{C_m} t\right) + \frac{G_\infty^2}{S_0^2} \left(1 - \exp\left(-\frac{S_0}{C_m} t\right)\right). \quad (17)$$

We will now focus on the existence and uniqueness of periodic solutions. Let G be a T -periodic source in $C_T(H^1(\Gamma))$, and \mathcal{A} a m-accretive operator in $\mathcal{L}(H^1(\Gamma))$. We will show that there exists a unique periodic solution u satisfying :

$$\begin{cases} u \in C_T(H^1(\Gamma)), \\ C_m \partial_t u + \mathcal{A}u + S_0 u + S_{\text{ep}}(t, u)u = G. \end{cases} \quad (18)$$

Let u_0 and $v_0 \in H^1(\Gamma)$ be two initial states and u, v the associated solutions to problem (18). Let $w = u - v$. w satisfies the homogeneous problem:

$$\begin{cases} C_m \partial_t w + \mathcal{A}w + S_0 w + S_{\text{ep}}(t, u)u - S_{\text{ep}}(t, v)v = 0, \\ w(t = 0, \cdot) = w(T, \cdot) = u_0(\cdot) - v_0(\cdot). \end{cases} \quad (19)$$

We have, for $t \geq 0$,

$$C_m \partial_t \|w\|_{L^2(\Gamma)}^2 + \langle \mathcal{A}w, w \rangle + S_0 \|w\|_{L^2(\Gamma)}^2 + \langle S_{\text{ep}}(t, u)u - S_{\text{ep}}(t, v)v, u - v \rangle = 0, \quad (20)$$

and since \mathcal{A} is m -accretive,

$$C_m \partial_t \|w\|_{L^2(\Gamma)}^2 + S_0 \|w\|_{L^2(\Gamma)}^2 + \langle S_{\text{ep}}(t, u)u - S_{\text{ep}}(t, v)v, u - v \rangle \leq 0. \quad (21)$$

We will show that for each conductivity model, the term $\langle S_{\text{ep}}(t, u)u - S_{\text{ep}}(t, v)v, u - v \rangle$ is positive.

- In the case of the β -model, it is easy to verify that for any λ_1, λ_2 with $\lambda_1 \neq 0$,

$$(\beta(\lambda_1)\lambda_1 - \beta(\lambda_2)\lambda_2)(\lambda_1 - \lambda_2) = \beta(\lambda_1)\lambda_1^2 \left(1 - \frac{\lambda_2}{\lambda_1}\right) \left(1 - \frac{\beta(\lambda_2)\lambda_2}{\beta(\lambda_1)\lambda_1}\right)$$

is positive, since β is an increasing function on $[0, +\infty)$.

- For the X -model, the ODE

$$\frac{d}{ds} X(s, \lambda(t)) = \frac{\beta(\lambda(t)) - X(s, \lambda(t))}{\tau_{\text{ep}}},$$

has the following solution

$$X(s, \lambda(t)) = e^{-\frac{s}{\tau_{\text{ep}}}} X(t, \lambda(t)) + \frac{\beta(\lambda(t))}{\tau_{\text{ep}}} \int_0^s e^{-\frac{s-s'}{\tau_{\text{ep}}}} ds'.$$

Therefore the mapping $\lambda \mapsto \lambda X(t, \lambda(t))$ is monotone. Using the same argument as for the β -model, we infer that for all $\lambda_1, \lambda_2, \lambda_1 \neq 0$

$$(X(\lambda_1)\lambda_1 - X(\lambda_2)\lambda_2)(\lambda_1 - \lambda_2) = X(\lambda_1)\lambda_1^2 \left(1 - \frac{\lambda_2}{\lambda_1}\right) \left(1 - \frac{X(\lambda_2)\lambda_2}{X(\lambda_1)\lambda_1}\right)$$

is positive.

For each conductivity model, we have, for $t \geq 0$,

$$C_m \partial_t \|w\|_{L^2(\Gamma)}^2 + S_0 \|w\|_{L^2(\Gamma)}^2 \leq 0,$$

and

$$\|w(t)\|_{L^2(\Gamma)}^2 \leq e^{-\frac{S_0}{C_m} t} \|w_0\|_{L^2(\Gamma)}^2. \quad (22)$$

Let $\Phi(u_0) := u(T)$. Writing the previous equation in $t = T$, we have

$$\|\Phi(u_0) - \Phi(v_0)\|_{L^2(\Gamma)} \leq e^{-\frac{S_0}{C_m} T} \|u_0 - v_0\|_{L^2(\Gamma)}, \quad (23)$$

with $e^{-\frac{S_0}{c_m}T} < 1$. Therefore, Φ is a contraction on $L^2(\Gamma)$ and there exists a unique initial state u_0 satisfying Eq. (18). We have also proven the asymptotic convergence of any solution to problem (10) to the periodic solution. \square

Remark 1. Note that the H^1 regularity is required on u and v so as their values are L^∞ , thanks to the injection $H^s \hookrightarrow L^\infty$ for $s > \frac{d}{2}$. If we had considered the three dimensional case, then more regularity would have been necessary: u_0 and v_0 should be in $H^{3/2}(\Gamma)$ and G should be in $C_T(H^{3/2}(\Gamma))$.

Remark 2. If the operator \mathcal{A} is coercive, with $\langle \mathcal{A}u, u \rangle > C_{\mathcal{A}} \|u\|_{L^2(\Gamma)}^2$ for all $u \in H^1(\Gamma)$, equation (20) becomes

$$C_m \partial_t \|w\|_{L^2(\Gamma)}^2 + S_0 \|w\|_{L^2(\Gamma)}^2 + \langle S_{\text{ep}}(t, u)u - S_{\text{ep}}(t, v)v, u - v \rangle \leq -C_{\mathcal{A}} \|w\|_{L^2(\Gamma)}^2$$

and

$$\|w(t)\|_{L^2(\Gamma)}^2 \leq e^{-\frac{S_0 + C_{\mathcal{A}}}{C_m}t} \|w_0\|_{L^2(\Gamma)}^2. \quad (24)$$

In this case, the convergence speed depends not only on the base conductivity of the membrane, but also on the shape of the cell and the extracellular conductivity, that are present in the definition of \mathcal{A} . We will detail in the next section a way to highlight the influence of these parameters for a circular cell.

4 Estimations of the convergence speed in the case of the circular cell.

4.1 Coercivity of the operator \mathcal{A}

Under specific conditions on the cell shape, it is possible to refine the previous estimation of the convergence rate to periodic solutions, by studying the operator \mathcal{A} .

Proposition 2.

Let \mathcal{O}_c be a circular disk of radius R_1 and \mathcal{O}_e a concentric ring around \mathcal{O}_c of outer radius $R_2 > R_1$. Let $H_p^{1/2}(\Gamma)$ be the space of functions of $H^{1/2}(\Gamma)$ with a zero mean value. Then the operator $(H_p^{1/2}(\Gamma), \mathcal{A}) : u \mapsto \mathcal{A}u = \Lambda_c(\text{Id} + \Lambda_e^{-1}\Lambda_c)^{-1}u$ with Λ_e and Λ_c defined by eqs. (12), is coercive.

Proof: We proceed by explicitly defining the Dirichlet-to-Neumann operators. Let us recall the definition of Λ_e :

$$\Lambda_e : u \mapsto -\sigma_e \partial_{\mathbf{n}} v \quad \text{with} \quad \begin{cases} \Delta v = 0 \text{ in } \mathcal{O}_e, \\ v = 0 \text{ on } \partial\Omega, \\ v = u \text{ on } \Gamma. \end{cases}$$

Expressing u and v with their Fourier transforms:

$$v = \sum_{k \in \mathbb{Z}^*} \alpha_k r^{|k|} e^{ik\theta} + \beta_k r^{-|k|} e^{ik\theta}, \quad u = \sum_{k \in \mathbb{Z}^*} u_k e^{ik\theta},$$

with constant terms $u_0 = v_0 = 0$, the boundary conditions on Γ and $\partial\Omega$ lead to

$$\begin{cases} \alpha_k R_1^{|k|} + \beta_k R_1^{-|k|} = u_k, \\ \alpha_k R_2^{|k|} + \beta_k R_2^{-|k|} = 0, \end{cases} \Rightarrow \partial_n v_k|_{R_1} = \frac{|k|}{R_1} \frac{R_2^{2|k|} + R_1^{2|k|}}{R_2^{2|k|} - R_1^{2|k|}} u_k.$$

Therefore

$$\Lambda_e^{-1} : v_k e^{ik\theta} \mapsto \frac{R_1}{\sigma_e |k|} \frac{R_2^{2|k|} - R_1^{2|k|}}{R_2^{2|k|} + R_1^{2|k|}} v_k e^{ik\theta}.$$

Using the same method for Λ_c leads to

$$\mathcal{A} = \Lambda_c (\text{Id} + \Lambda_e^{-1} \Lambda_c)^{-1} : u_k e^{ik\theta} \mapsto \frac{|k|}{R_1} \left(\frac{1}{\sigma_e} \frac{R_2^{2|k|} - R_1^{2|k|}}{R_2^{2|k|} + R_1^{2|k|}} + \frac{1}{\sigma_c} \right)^{-1} u_k e^{ik\theta}.$$

Then

$$\begin{aligned} \langle \mathcal{A}u, u \rangle &\geq \left(\frac{1}{\sigma_c} + \frac{1}{\sigma_e} \right)^{-1} \frac{1}{R_1} \sum_{k,l \in \mathbb{Z}} |k| u_k u_l \langle e^{ik\theta}, e^{il\theta} \rangle, \\ &= \left(\frac{1}{\sigma_c} + \frac{1}{\sigma_e} \right)^{-1} \frac{1}{R_1} \sum_{k,l \in \mathbb{Z}} |k| u_k u_l \int_0^{2\pi} e^{ik\theta} e^{-il\theta} d\theta, \\ &= \left(\frac{1}{\sigma_c} + \frac{1}{\sigma_e} \right)^{-1} \frac{1}{R_1} \sum_{k \in \mathbb{Z}} |k| u_k^2, \\ &\geq \left(\frac{1}{\sigma_c} + \frac{1}{\sigma_e} \right)^{-1} \frac{1}{R_1} \|u\|_{H^{1/2}(\Gamma)}^2, \\ \langle \mathcal{A}u, u \rangle &\geq C_{\mathcal{A}} \|u\|_{L^2(\Gamma)}^2 \end{aligned} \tag{25}$$

according to the definition of $\|\cdot\|_{H^{1/2}(\Gamma)}$ given by [10], p. 141, with

$$C_{\mathcal{A}} := \left(\frac{1}{\sigma_c} + \frac{1}{\sigma_e} \right)^{-1} \frac{1}{R_1}.$$

Therefore \mathcal{A} is coercive. □

Remark 3. Remark that we cannot extend this property to functions with a non-zero mean value, as a consequence of the non-invertibility of Λ_c . For example, constant functions $u = u_0 \neq 0$ verify $\langle \mathcal{A}u, u \rangle = 0 < \|u\|_{H^{1/2}(\Gamma)}^2$.

Remark 4. Also note that it is possible to consider a cell embedded in a uniform electric field by making R_2 grow to infinity and considering only the case $|k| = 1$. Under these conditions, that are the most frequently found in experiments on cells, the ratio $(R_2^{2|k|} - R_1^{2|k|}) / (R_2^{2|k|} + R_1^{2|k|})$ tends to 1 and $\|u\|_{H^{1/2}(\Gamma)} = \|u\|_{L^2(\Gamma)}$, so that

$$\langle \mathcal{A}u, u \rangle = \left(\frac{1}{\sigma_c} + \frac{1}{\sigma_e} \right)^{-1} \frac{1}{R_1} \|u\|_{L^2(\Gamma)}.$$

Remark 5. It is possible to extend the coercive property of \mathcal{A} to the three dimensional case, using spherical harmonic functions instead of usual Fourier series. Following the same computation,

the estimation on the coercivity constant differs from the 2D case by a factor 2 :

$$\langle \mathcal{A}u, u \rangle \geq \left(\frac{1}{\sigma_c} + \frac{1}{\sigma_e} \right)^{-1} \frac{1}{2R_1} \|u\|_{H^{1/2}(\Gamma)}^2.$$

Going back to problem (19), the coercivity of the operator \mathcal{A} leads to

$$\frac{1}{2} C_m \partial_t \|w\|_{L^2(\Gamma)}^2 \leq -(S_0 + C_A) \|w\|_{L^2(\Gamma)}^2.$$

and for all $t \geq 0$

$$\|w(t)\|_{L^2(\Gamma)} < e^{-\frac{(S_0 + C_A)}{C_m} t} \|w_0\|_{L^2(\Gamma)}, \quad (26)$$

In this case, the convergence speed also depends on parameters included in the constant C_A : the extracellular conductivity and the size of the cell. These dependencies will be illustrated in the next section.

4.2 Numerical validation

In order to solve numerically equation (2), the same scheme as in [5] was used. It is based on a method developed by Cisternino and Weynans [2], using second order finite differences for spatial discretization and adding unknowns at the intersection points of the interface Γ on the cartesian grid for a special treatment of fluxes. The details of the method are fully explicated and its convergence proven in a another previous publication dedicated to the numerical aspects of the problem [6].

Simulations were all performed in a two dimensional domain with the parameters presented in table 1. Since the numerical method is based on a cartesian grid, the simulated domain is a square $[x_{\min}, x_{\max}] \times [y_{\min}, y_{\max}]$ that is large enough compared to the radius of the cell, which is circular.

The chosen boundary conditions simulate a uniform electric field of amplitude E . It consists in a Dirichlet condition in the x direction, and a homogeneous Neumann condition in the y direction. In this case, effects due to the presence of the square-shaped boundary on the transmembrane potential difference (Δ TMP) can be neglected.

The problem to solve reads

$$\begin{cases} \Delta U = 0 & \text{in } \mathcal{O}_e \cup \mathcal{O}_c, \\ U(t=0, x, y) = 0 & \text{in } \Omega, \\ [\sigma \partial_{\mathbf{n}} U] = 0 & \text{on } \Gamma, \\ C_m \partial_t [U] + S_m(t, [U]) [U] = \sigma_c \partial_{\mathbf{n}} U|_{\Gamma^-} & \text{on } \Gamma \\ U(t, x, y) = Ex & \text{if } x = x_{\min} \text{ or } x = x_{\max}, \forall t > 0, \\ \partial_{\mathbf{n}} U(t, x, y) = 0 & \text{if } y = y_{\min} \text{ or } y = y_{\max}, \forall t > 0. \end{cases} \quad (27)$$

4.2.1 Results of the linear model

Before studying the influence of the electropermeabilization model on the convergence towards a periodic solution, we will focus on the variation of the coercivity constant C_A with the extracellular conductivity σ_e . In this section, we will consider a constant surface conductivity of the membrane.

The chosen pulse parameters to study the linear effects are not commonly used. Using a clinical pulse protocol with a linear model does not lead to relevant convergence results since

Table 1: Parameters of numerical simulations, that are mostly taken from [5]. EP stands for electropermeabilization, EPd for electropermeabilized.

Variable	Symbol	Value	Unit
Biological parameters:			
Intracellular conductivity	σ_c	0.455	S/m
Capacitance	C_m	1	F/m ²
Membrane surface conductivity	S_0	1.9	S/m ²
Cell radius	R_1	6	μm
Specific parameters of the models:			
EP threshold	V_{rev}	0.2	V
EP switch speed	k_{ep}	40	V ⁻¹
EP characteristic time	τ_{ep}	1×10^{-6}	s
EPd membrane surface conductivity	S_1	1×10^6	S/m ²
Numerical parameters:			
Simulation box size	L	30	μm
Grid points (each side)	N	100	
Time step	Δt	200	ns
Pulse parameters:			
Pulse duration	T_p	10	μs
Pulse period	T	20	μs
Intensity	E	40	kV/m
Duration of simulation	T_f	1000	μs

pulses are separated by a time that is long enough to let the cell retrieve its initial state. The solution is therefore already periodic for any pulse intensity. To prevent this, the pulses frequency was considerably augmented to avoid the complete discharge of the membrane between two pulses. Moreover, the membrane capacitance C_m was also changed to 1 (instead of ~ 0.01) to extend the duration of the charge and discharge of the membrane. This allows also direct comparison with the estimation of C_A in equation (25).

The periodic solution \bar{u} to problem (18) is obtained by performing long-time simulations of 50 periods, and verifying that the last obtained period does not vary from the previous one more than a ratio of 10^{-10} : if T_f designates the final time of the simulation,

$$\frac{\int_{T_f-T}^{T_f} \|u(t) - u(t-T)\|_{H^{1/2}(\Gamma)} dt}{\int_{T_f-T}^{T_f} \|u(t)\|_{H^{1/2}(\Gamma)} dt} < 10^{-10}. \quad (28)$$

Under these conditions, the last period is considered as the periodic solution. We then compute the L^2 error

$$e(t) := \|\bar{u}(t) - u(t)\|_{L^2(\Gamma)}.$$

The error is fitted with an exponential function

$$e(t) \sim A \exp^{-Bt}, \quad (29)$$

where A and B are constants, as illustrated in figure 2. The constant B can be considered as an estimation of the convergence speed C_A .

The process is repeated for different extracellular conductivities σ_e and radiuses R_1 . The Fourier coefficients of the solution are computed to check whether the influence of the square-shaped simulation box is important or not. All simulations showed a solution with greatest

coefficients $u_k, |k| \neq 1$ being $u_{|3|}$, 1000 times lower than the main harmonic $u_{|1|}$. Figure 3 shows that the dependency on σ_e and R_1 of the decay rate $B \leq S_0 + C_A$ is satisfied. The evaluated constant B has the same order of magnitude as the estimation of C_A in (25).

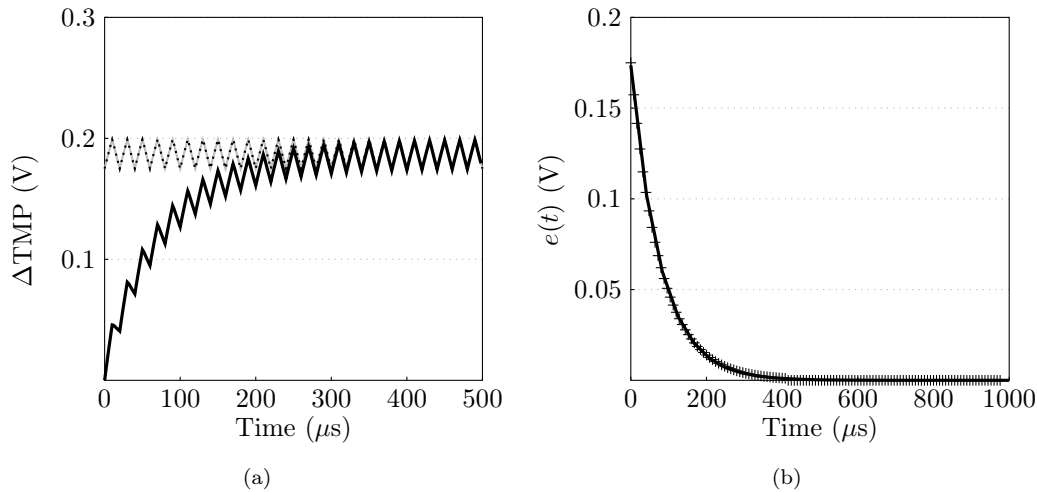


Figure 2: 2(a) : Transmembrane potential difference (solid) and periodic solution (dotted) of the linear problem, with $\sigma_e = 0.2 \text{ S.m}^{-1}$. 2(b) : Fit of the L^2 error $e(t)$ between the ΔTMP and the periodic solution, giving the constant C_A .

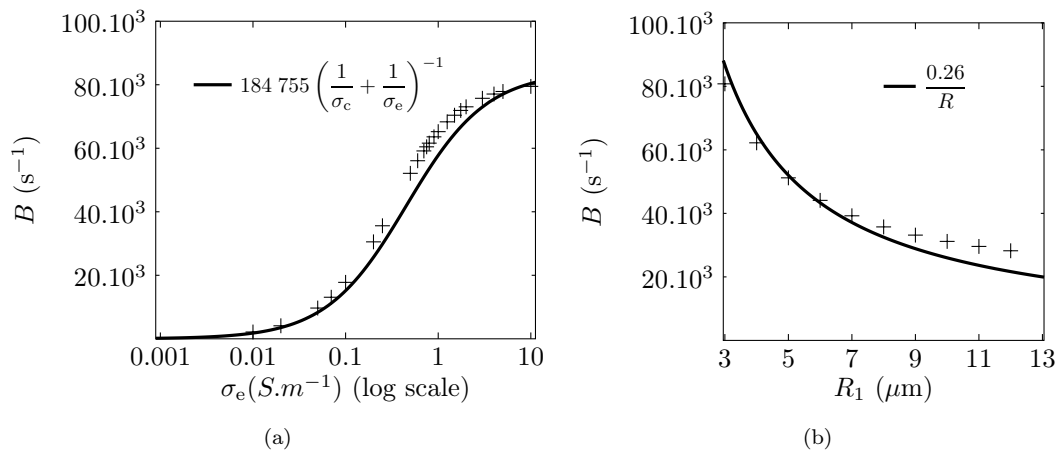


Figure 3: Linear conductivity model : evaluated decay rate B of the L^2 error depending on the extracellular conductivity σ_e with $R_1 = 6 \mu\text{m}$ (3(a)) and depending on the radius of the cell R_1 with $\sigma_e = 0.5 \text{ S/m}$ (3(b)). The first fitting constant has the same order of magnitude as $R_1^{-1} \sim 167\,000$. The other constant is of the order of $(\sigma_e^{-1} + \sigma_c^{-1})^{-1} \sim 0.24$.

4.2.2 Results of the static conductivity models

The same method was used to study the decay rate to periodic solutions when the membrane conductivity is a non-linear function of the transmembrane potential difference. Simulations showed that the convergence of the solution to the periodic state is still an exponential function of time:

$$e(t) \sim e^{-Ct}e(0).$$

even in the non-linear case. The constant C still includes the base conductivity of the membrane and the coercivity of the operator \mathcal{A} . We propose to note $C = S_0 + C_{\mathcal{A}} + C_{S_{ep}}$, where $C_{S_{ep}}$ accounts for the non-linearity of the conductivity.

Figure 4 shows the behaviour of the quantity C when the extracellular conductivity varies. The dependency on σ_e is similar to the linear case, the difference being only a constant factor. This can be explained by the increase in the maximum value of the Δ TMP that is obtained for each simulation. Lower conductivities induce a limited rise of the Δ TMP, contrary to high conductivities, for which the Δ TMP is close to the threshold voltage. Then the Lipschitz constant of $\lambda \mapsto \lambda S_{ep}$ vary accordingly to the maximum value of the Δ TMP λ_m :

$$C_{S_{ep}} \leq S_1 \max_{\lambda} |(\lambda\beta(\lambda))'| = S_1 (\beta(\lambda_m) + |\lambda_m| \beta'(\lambda_m)) \quad (30)$$

since $\lambda_m < V_{rev}$ in all our simulations. Figure 5 shows that the simulation results are in agreement with this estimation, although the proposed constant for the β model is one order of magnitude higher.

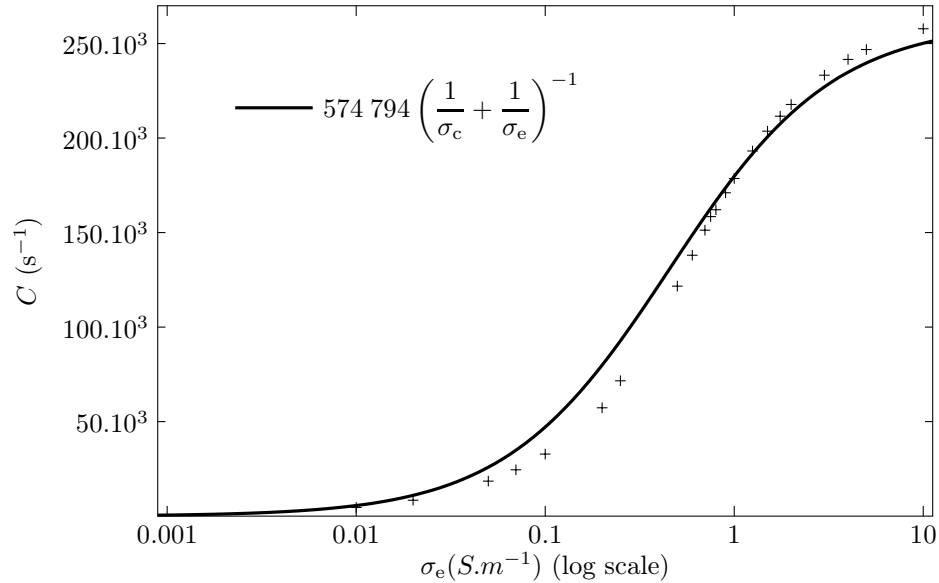


Figure 4: Evaluated convergence rate to the periodic solution for the case $S_m(\lambda) = S_0 + S_1\beta(\lambda)$.

4.2.3 Dynamical model

Figure 6 shows a different evolution of the non-linearity for the dynamical model. For lower conductivities, the behaviour is the same as the static cases, since the applied pulse train is not

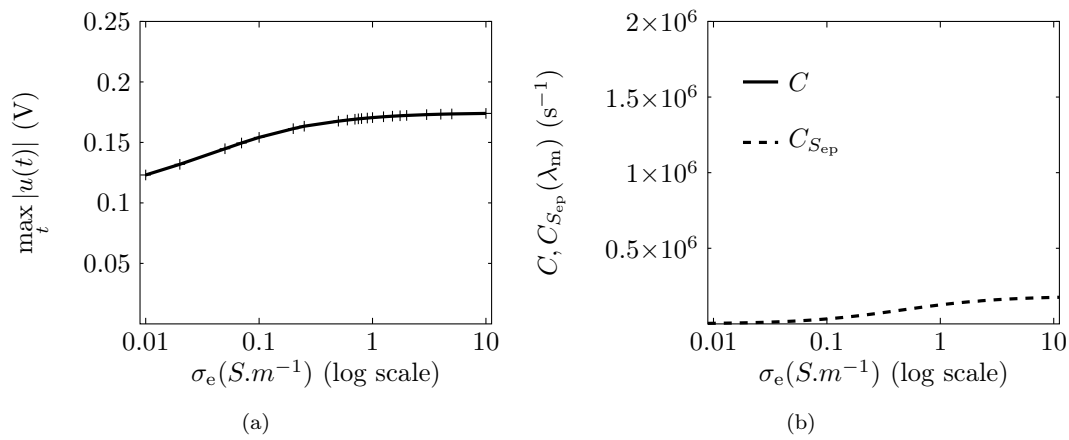


Figure 5: 5(a): Maximum Δ TMP during simulations for the β (solid) and λ^2 (dashed) static models. 5(b): Comparison of simulated and estimated Lipschitz constants $C_{S_{ep}}$ for the β model. Solid lines : difference between non-linear and linear decay rates. Dashed lines : estimations from eq (30) with λ_m from figure 5(a).

sufficient to reach the voltage threshold. However, as the extracellular conductivity rises, a drop of the convergence speed occurs, which depends on the dynamics of X . If τ_{ep} is low enough, the dynamical model has a behaviour which is the same as the β -model, whereas slower dynamics do not affect the convergence speed more than the linear constant.

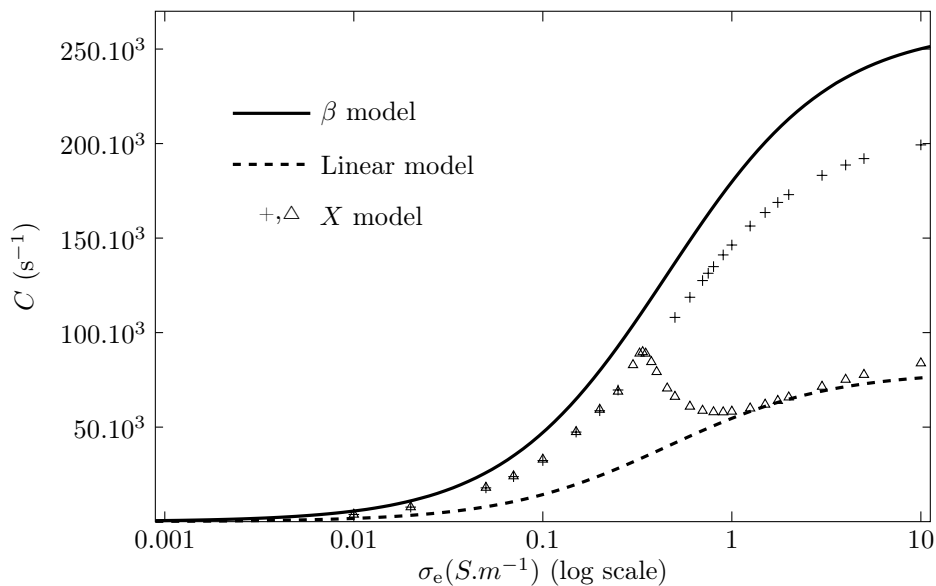


Figure 6: Decay rate to the periodic solution for the dynamical case $S_m(\lambda) = S_0 + S_1 X(t)$, compared to the evaluations already performed for the linear and the β models. Simulations of the dynamical model were made for several characteristic durations: $\tau_{ep} = 1 \mu s$ (+), $\tau_{ep} = 10 \mu s$ (Δ).

Since the choice of the model has a quantitative impact on the convergence speed to a periodic state, it may be important to translate the results of figure 6 in terms of pulse application. Figure 7 gives the number of pulses that are necessary to obtain a periodic response of the cell. This number is computed so as the solution is no more than 1% different from the periodic solution in $L^2(\Gamma)$ norm:

$$N(\sigma_e) := \left\lceil \frac{2 \ln 10}{C(\sigma_e)T} \right\rceil \quad (31)$$

It is shown that for the permeabilizing cases ($\sigma_e \geq 0.3$ S/m), three to four additional pulses are required for the dynamical model to achieve convergence, compared to the static model for which two pulses are sufficient. If it is intended to obtain a conductivity which corresponds to the periodic state, then considering only a static model of conductivity can lead to a lower level of permeabilization than expected.

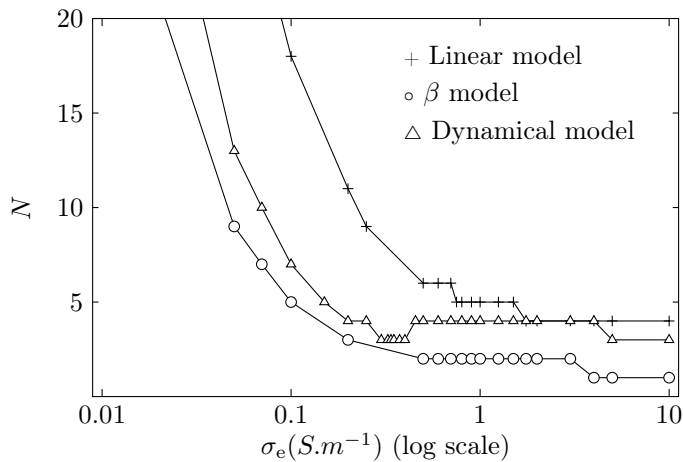


Figure 7: Number of pulses $N(\sigma_e)$ given by eq. (31) for the linear, β and dynamical models.

5 Conclusion

In this paper we study the evolution of solutions to several models of electropermeabilization derived from Kavian et al. [5] in the specific case of periodic sources. We show existence and uniqueness of periodic solutions, as well as convergence to these solutions given any initial condition. Using a simulation tool that was created specifically to solve electropermeabilization problems, we validate theoretical estimations of the convergence speed to periodic solutions.

We emphasize that static models of membrane conductivity give an overestimation of the convergence speed, compared to the dynamical model. Since the permeabilization is linked to the conductivity, those models predict a higher permeabilization level of the cell for short pulse treatments. The number of pulses needed to obtain a given conductivity is then lower than predictions of the dynamical model, and the cell could not be actually porated.

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