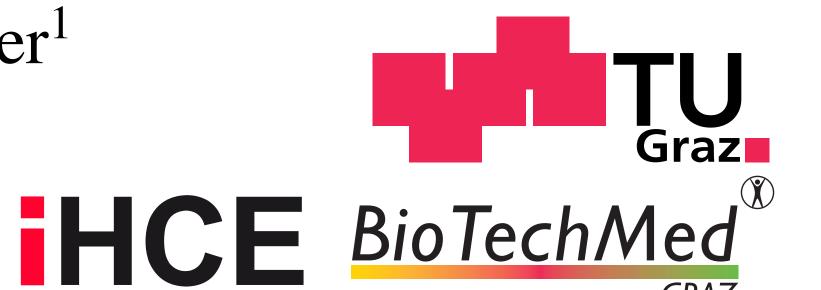
REAL-TIME INTERACTION WITH AN ELECTROPHYSIOLOGICAL CANCER CELL MODEL

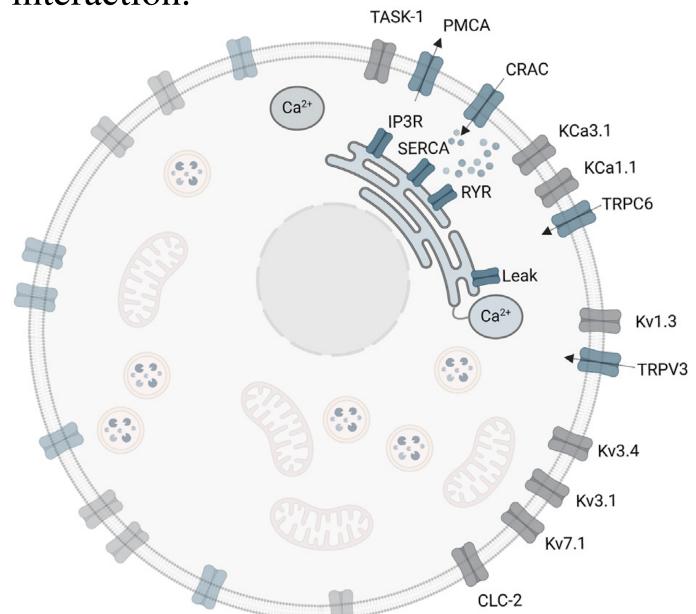
Peter Waldert^{1,2}, Theresa Rienmüller¹, Christian Baumgartner¹ and Sonja Langthaler¹

¹Institute of Health Care Engineering with European Testing Center of Medical Devices, TU Graz ²Institute of Computer Graphics and Knowledge Visualisation, TU Graz



Introduction

We improve on the A549 electrophysiological cancer cell model introduced in [1, 2], combining numerical methods with an efficient implementation to reduce simulation time to a level where it is feasible for live interaction.



[Image Source: [2]]. We accelerated the simulation with adaptive timestepping and a highly efficient implementation in the Rust programming language. The corresponding inverse problem is formulated into a quadratic program, solving it within milliseconds. We introduce a visualisation approach of the entire model in the form of a live simulation dashboard available online, running directly in the browser.

Live, In-Browser Cell Simulation Interface



This interface is available online via in-silico.hce.tugraz.at, try it out!

Model

The whole cell current $I: T \to \mathbb{R}$ over time $t \in T \subset \mathbb{R}^+$ is the sum of all individual channel contributions $I_k, k \in \{1, ..., M\}$ over $M \in \mathbb{N}$ channel types

$$I(t) := \sum_{k=1}^{M} N_k I_k(t) = \sum_{k=1}^{M} N_k g_k p_{o,k} \left(V(t) - E_k \right) ,$$

At each time step, the next state $s_{k,n+1} \in [0,1]^{N_{s,k}}$ of the k-th channel type is obtained by

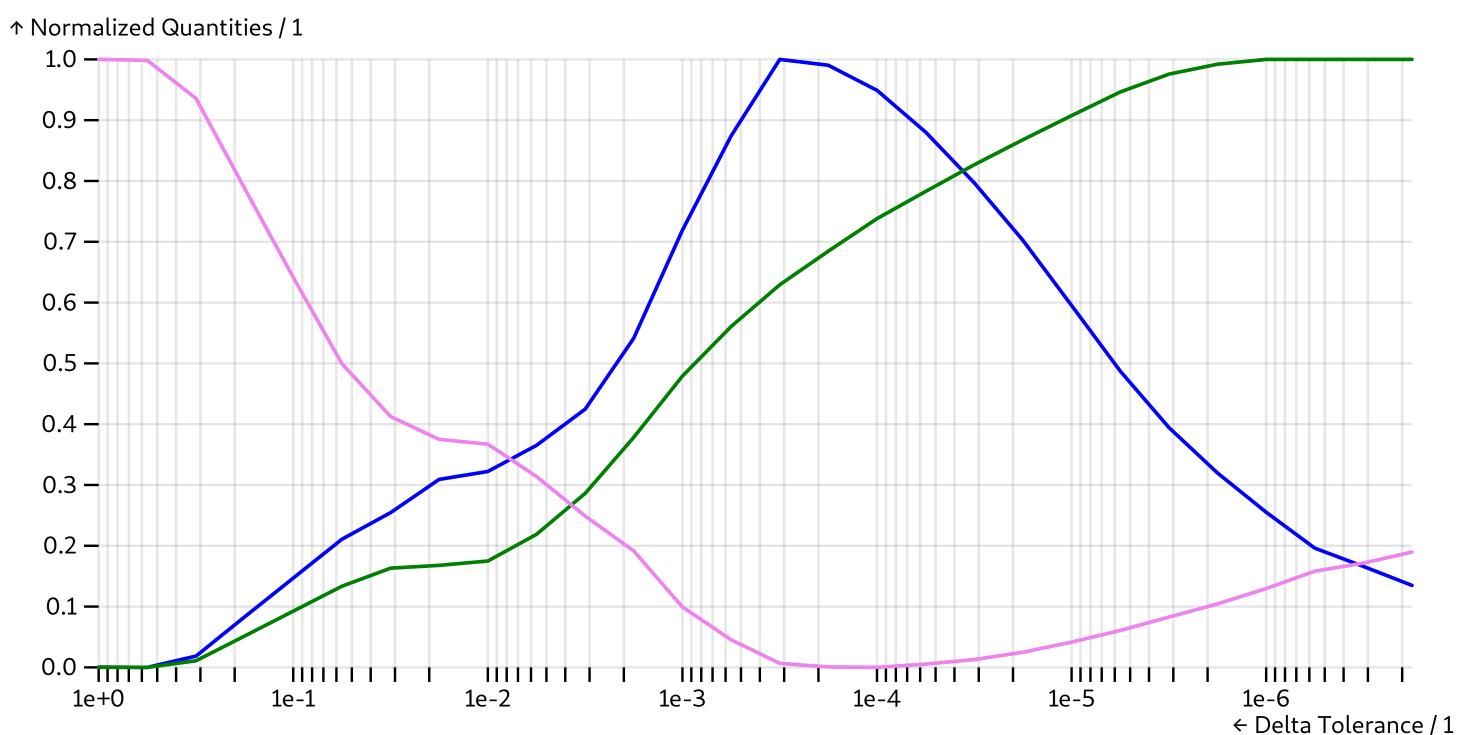
$$oldsymbol{s}_{k,n+1} = H_k\left(V(t_n), oldsymbol{C}(t_n), t_n\right) oldsymbol{s}_{k,n} \,, \quad ext{with} \quad t_n := \sum\limits_{i=0}^n (\Delta t)_i \,.$$

Adaptive Timestepping

In order to accelerate the simulation in areas where there is little change to the dynamics, we choose an adaptive step size $(\Delta t)_n$ based on

$$(\Delta t)_{n+1} = (\Delta t)_n \left(rac{\Delta^{ ext{tol}}}{\sum_{k=1}^M N_k \left\| oldsymbol{s}_{k,n+1} - oldsymbol{s}_{k,n}
ight\|_2}
ight)^{1/2} \,.$$

In order to find the optimal $\Delta^{tol} \in \mathbb{R}^+$ (the allowed state change in between steps):



Relative change of the average timestep Δt (in blue), simulation runtime (in violet) and step acceptance rate (in green) when varying the delta tolerance $\Delta^{\rm tol}$ on a log-scale. All three quantities were normalized from their individual extent to [0,1]. The most effective $\Delta^{\rm tol}$ is arguably on the order of 10^{-4} .

Formulation as a Quadratic Program

After simulation of the individual channel currents, we want to find

$$oldsymbol{N}_{ ext{opt}} = rg\min_{oldsymbol{N} \in \mathbb{N}_0^M} rac{1}{2} \left\| R oldsymbol{N} - oldsymbol{I}_{ ext{meas}}
ight\|_2^2 \ .$$

the number of channels per type, with $I_{\text{meas}} \in \mathbb{R}^{N_t}$ the experimentally measured current and $R \in \mathbb{R}^{N_t \times M}$ the matrix of all currents I_k per channel type. Letting $d := I_{\text{meas}}$ for brevity,

$$oldsymbol{N}_{ ext{opt}} pprox rg \min_{oldsymbol{x} \in \mathbb{R}_+^M} f(oldsymbol{x}) = rg \min_{oldsymbol{x} \in \mathbb{R}_+^M} {}^1 \! / \! {}_2 \left\| R oldsymbol{x} - oldsymbol{d}
ight\|_2^2 \; ,$$

with cost function $f: \mathbb{R}^M \to \mathbb{R}^+$, which we manipulate to

$$f(\boldsymbol{x}) = \frac{1}{2}(R\boldsymbol{x} - \boldsymbol{d})^{T}(R\boldsymbol{x} - \boldsymbol{d})$$

$$= \frac{1}{2}(\boldsymbol{x}^{T}R^{T}R\boldsymbol{x} - \boldsymbol{x}^{T}R^{T}\boldsymbol{d} - \boldsymbol{d}^{T}R\boldsymbol{x} + \boldsymbol{d}^{T}\boldsymbol{d})$$

$$= \frac{1}{2}(\boldsymbol{x}^{T}P\boldsymbol{x} + \boldsymbol{x}^{T}\boldsymbol{q} + \boldsymbol{q}^{T}\boldsymbol{x}) + \mathcal{O}(1)$$

$$= \frac{1}{2}\boldsymbol{x}^{T}P\boldsymbol{x} + \boldsymbol{q}^{T}\boldsymbol{x} + \mathcal{O}(1)$$

where we let $P := R^T R \in \mathbb{R}^{M \times M}$ and $\mathbf{q} := -R^T \mathbf{d} \in \mathbb{R}^M$ and leave out the constant $\mathbf{d}^T \mathbf{d}$ as $\mathcal{O}(1)$. We can express the nonnegativity constraint $\mathbf{x} \geq \mathbf{0}$ as an equality constraint using a slack variable $\mathbf{s} \in \mathbb{R}_+^M$,

$$-x + s = 0 \Leftrightarrow Ax + s = b \text{ with } A = -1 \in \mathbb{R}^{M \times M} \text{ and } b = 0 \in \mathbb{R}^{M}.$$

This leaves us with a constrained quadratic program,

$$\min_{oldsymbol{x} \in \mathbb{R}^M} \frac{1}{2} oldsymbol{x}^T P oldsymbol{x} + oldsymbol{q}^T oldsymbol{x}, \ s.t. \ A oldsymbol{x} + oldsymbol{s} = oldsymbol{b} \ , \ oldsymbol{s} \in \mathbb{R}_+^M \ .$$

The integer solution can then be obtained from rounding,

$$oldsymbol{N}_{ ext{opt}} = \lfloor oldsymbol{x}
ceil \in \mathbb{N}_0^M$$
 .

Comparison of Optimization Methods

The different approaches were evaluated on the G0 cell cycle phase with the activation voltage protocol. Runtime estimates were obtained on an IntelTMi7-5600U CPU.

Algorithm	Abbreviation	Runtime / ms	RMSE / pA
Particle Swarm Optimization	PSO	22571	27.69
Gradient Descent + More Thuente	GD	18924	32.34
Limited-Memory BFGS + Hager Zhang	LBFGS	4845	32.20
Non-Negative Least Squares [3]	NNLS	318	28.00
Quadratic Program	QP	18	28.13

Conclusion

Efforts to further enhance and complete the first electrophysiological cancer cell model simulation were fruitful, resulting in a new library implementation, an improved inverse problem solution technique and a live in-browser simulation dashboard.

References

^[1] Sonja Langthaler, Theresa Rienmüller, Susanne Scheruebel, Brigitte Pelzmann, Niroj Shrestha, Klaus Zorn-Pauly, Wolfgang Schreibmayer, Andrew Koff and Christian Baumgartner. 'A549 in-silico 1.0: A first computational model to simulate cell cycle dependent ion current modulation in the human lung adenocarcinoma'. In: *PLoS Comput. Biol.* 17.6 (June 2021), e1009091. DOI: 10.1371/journal.pcbi.1009091.

^[2] Sonja Langthaler, Christian Zumpf, Theresa Rienmüller, Niroj Shrestha, Julia Fuchs, Rui Zhou, Brigitte Pelzmann, Klaus Zorn-Pauly, Eleonore Fröhlich, Seth H. Weinberg and Christian Baumgartner. 'The bioelectric mechanisms of local calcium dynamics in cancer cell proliferation: an extension of the A549 in silico cell model'. In: Front. Mol. Biosci. 11 (May 2024), p. 1394398. DOI: 10.3389/fmolb.2024.1394398.

mechanisms of local calcium dynamics in cancer cell proliferation: an extension of the A549 in silico cell model. In: Front. Mol. Biosci. 11 (May 2024), p. 1394398. DOI: 10.3389/fmo1b.2024.1394398. [3] Rasmus Bro and Sijmen De Jong. 'A fast non-negativity-constrained least squares algorithm'. In: J. Chemom. 11.5 (Sept. 1997), pp. 393–401. ISSN: 0886-9383. DOI: 10.1002/(SICI) 1099–128X (199709/10) 11: 5<393::AID-CEM483>3.0.CO; 2-L.