

Simulation of Astrocytic Calcium Dynamics in Lattice Light Sheet Microscopy Images

Anaïs Badoual, Misa Arizono, Audrey Denizot, Mathieu Ducros, Hugues Berry, U Valentin Nägerl, Charles Kervrann

▶ To cite this version:

Anaïs Badoual, Misa Arizono, Audrey Denizot, Mathieu Ducros, Hugues Berry, et al.. Simulation of Astrocytic Calcium Dynamics in Lattice Light Sheet Microscopy Images. ISBI 2021 - IEEE International Symposium on Biomedical Imaging, Apr 2021, Nice, France. pp.1. hal-03516709

HAL Id: hal-03516709 https://inria.hal.science/hal-03516709v1

Submitted on 7 Jan 2022

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Simulation of Astrocytic Calcium Dynamics in Lattice Light Sheet Microscopy Images



A. BADOUAL¹, M. ARIZONO², A. DENIZOT³, M. DUCROS⁴, H. BERRY^{5,6}, U.V. NÄGERL² and C. KERVRANN¹

¹Inria, Centre de Recherche Bretagne Atlantique, Serpico Team, Rennes, France ³Computational Neuroscience Unit, Okinawa Institute of Science and Technology, Japan ⁵Inria, Centre de Recherche Rhone-Alpes, Beagle Team, F-69603, Villeurbanne, France

²Interdisciplinary Institute for Neuroscience, UMR5297 CNRS, Bordeaux, France ⁴Bordeaux Imaging Center, UMS3420 CNRS, US04 INSERM, Bordeaux, France ⁶Univ Lyon, LIRIS, UMR5205 CNRS, F-69621, Villeurbanne, France

ABSTRACT

Astrocytes regulate neuronal information processing through a variety of spatio-temporal calcium signals. Advances in calcium imaging started to reveal astrocytic activities, but the complexity of the recorded data strongly call for computational analysis tools. Their development is hindered by the lack of reliable annotations that are essential for their evaluation and for the design of learning-based methods. To overcome the labeling problem, we present a framework to simulate realistic astrocytic calcium signals in 3D+time lattice light sheet microscopy (LLSM) images by closely modeling calcium kinetics in real astrocytes.

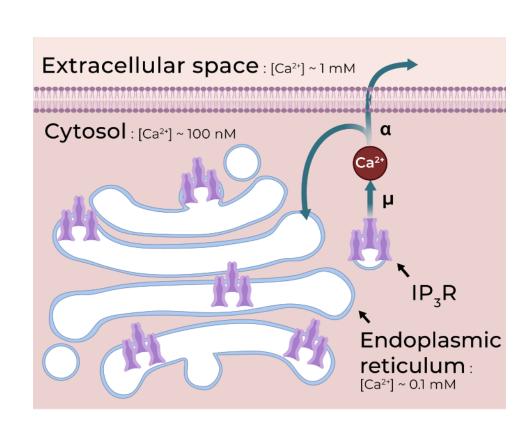
METHOD

Simulation of Calcium Puffs

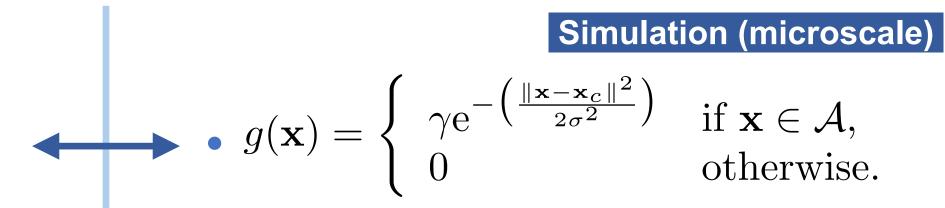
Kinetic model (nanoscale)

• Stimulus at time t_0 and location \mathbf{x}_c of an astrocyte triggering a γ variation of Ca^{2+3}

Cytosolic calcium fluxes:



Brownian motion



 ${\cal A}$: set of points belonging to astrocytes

 $\mathbb{1}_{\mathcal{O}_t}$: indicator function of the subset \mathcal{O}_t \mathcal{O}_t : set of open IP₃Rs at time t

 Δ_t : temporal resolution N_c : number of IP $_3$ Rs in a cluster

: sequence simulating one calcium event

div(⋅): divergence operator

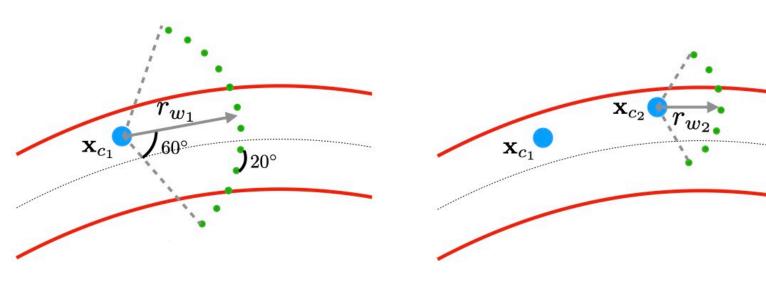
 ${\mathcal E}$: set of points belonging to astrocytes' edges

 β : maximum intensity $c(u) = \exp\left(-\left(\frac{u}{\kappa}\right)^{2}\right)$: conductivity function

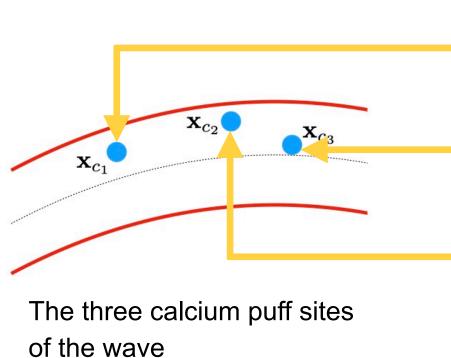
$$f_{\rm e}(\mathbf{x}, t + \Delta t_{\rm e}) = f_{\rm e}(\mathbf{x}, t) + \Delta t_{\rm e} \frac{\partial f_{\rm e}(\mathbf{x}, t)}{\partial t} + \Delta t_{\rm e} \left(\sum_{n=1}^{N_c} \mu \, \mathbb{1}_{\mathcal{O}_t}(n) - \alpha\right) \tag{1}$$

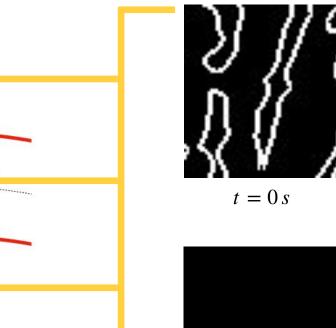
Simulation of Calcium Waves

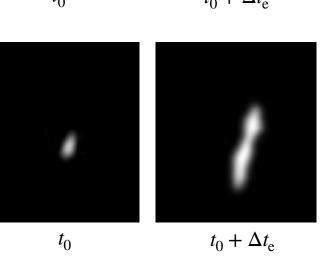
Propagation of calcium puffs from a IP₃R cluster to the neighboring ones

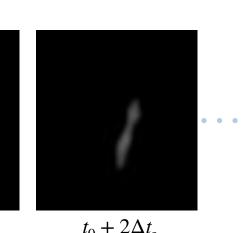


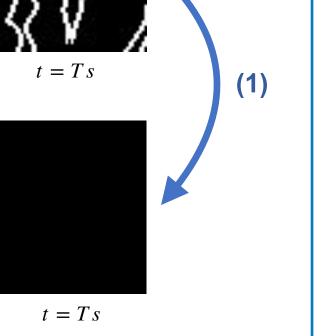
Determination of \mathbf{x}_{c} Determination of \mathbf{x}_{c}





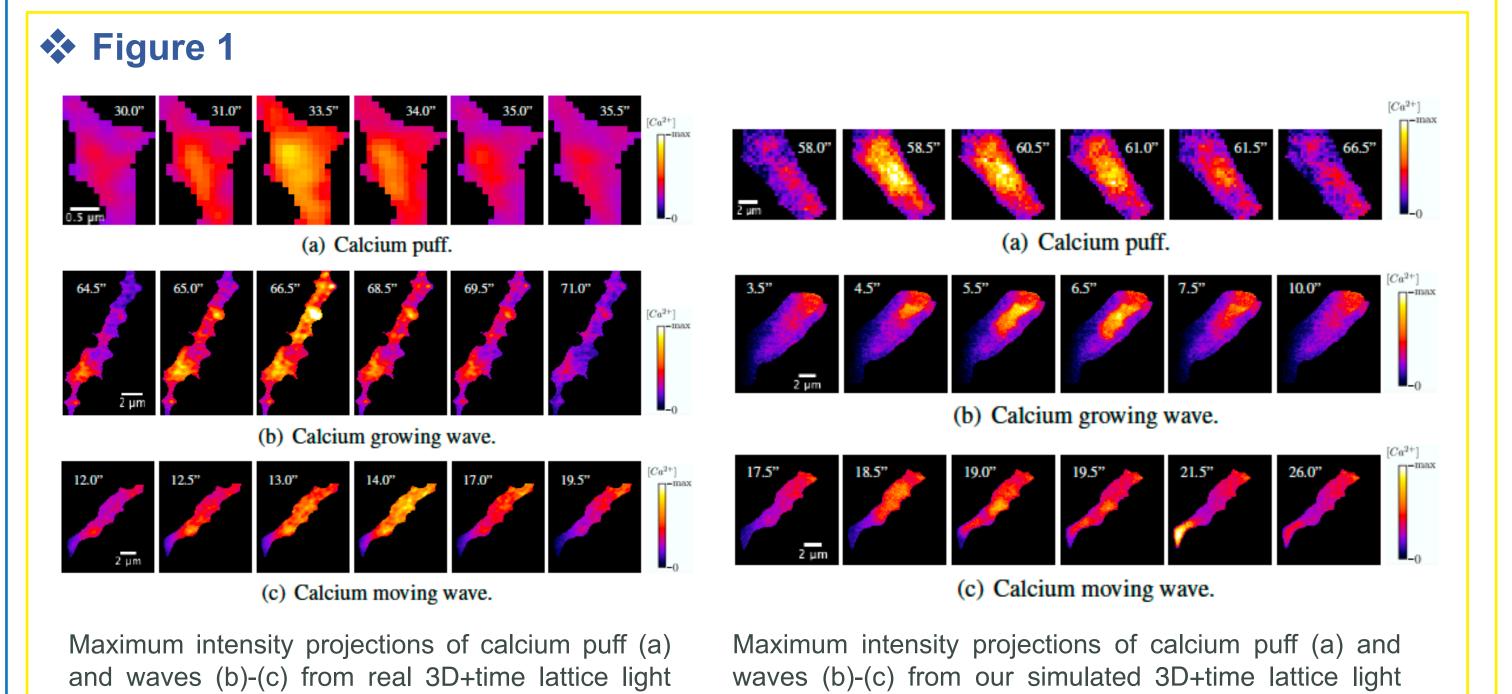






Edges of astrocytes

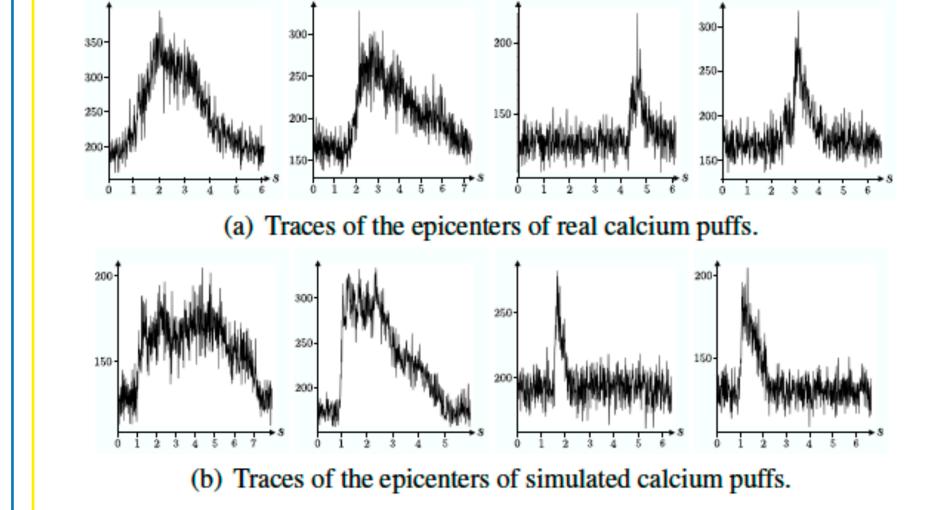
RESULTS



sheet microscopy images ($\Delta t = 0.5 s$)

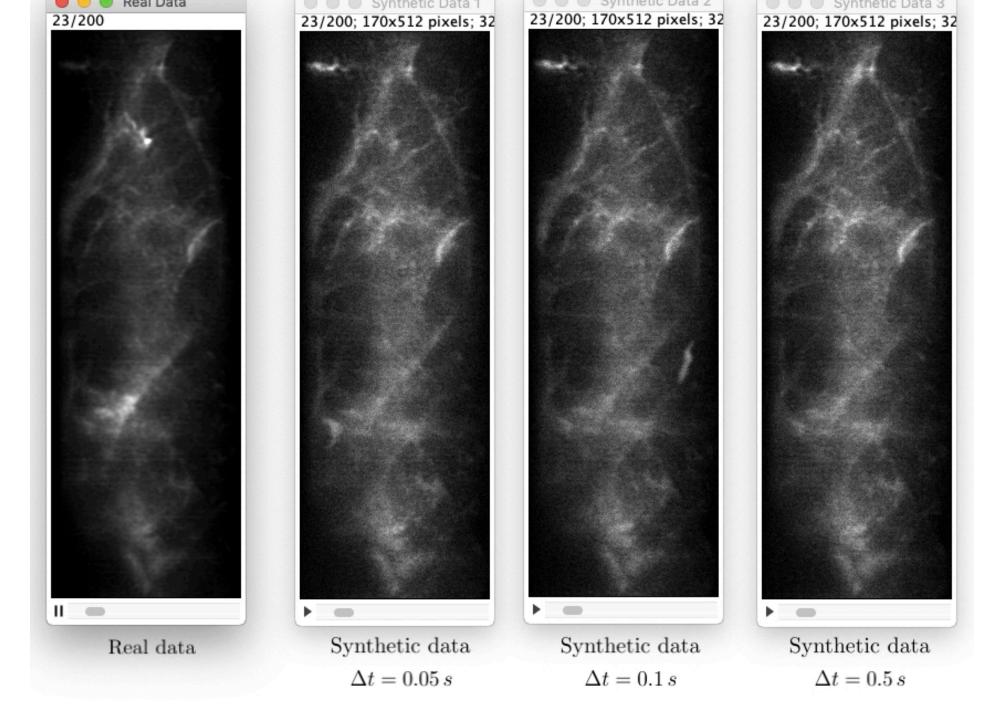


sheet microscopy images ($\Delta t = 0.5 s$).

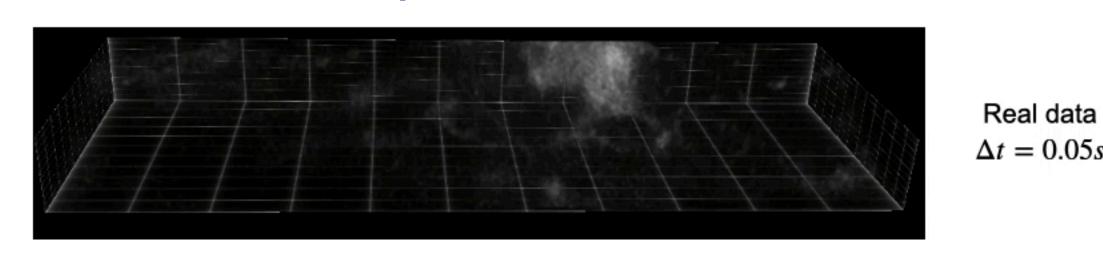


Traces of the epicenters of calcium puffs present in real (a) and synthetic (b) 2D+time lattice light sheet microscopy images ($\Delta t = 0.01 s$)

❖ Video 1: 2D+time sequences



❖ Video 2: 3D+time sequences



Synthetic data $\Delta t = 0.05s$

CONCLUSIONS

Contributions

- Generation of 3D+time LLSM images depicting astrocytic activity. First simulator of its kind
- Driven by a calcium kinetic model that we adapted from nanoscale to microscale
- Image background made of real astrocytic ramifications
- Simulation of experimental conditions of LLSM acquisitions (e.g., noise and blur)
- Implementation as an ImageJ plugin with parameters and user interface

Results

- Variety of synthetic datasets that realistically represents the complexity of experimental data
- Helpful to develop computerized learning-based methods to analyze astrocytic calcium signals in 3D+time images

REFERENCES

- A. Denizot et al., "Simulation of calcium signaling in fine astrocytic processes: Effect of spatial properties on spontaneous activity," PLoS Computational Biology, 2019
- M. Ducros et al., "Lattice light sheet microscopy and photostimulation in brain slices," Neural Imaging and Sensing, 2019
- M. Arizono et al., "Structural basis of astrocytic Ca²⁺ signals at tripartite synapses," Nature Communications, 2020
- E. Bindocci et al., "Three-dimensional Ca2+ imaging advances understanding of astrocyte biology," Science, 2017
- M. Santello et al., "Astrocyte function from information processing to cognition and cognitive impairment," Nature Neuroscience, 2019