

The nonlinear dynamics of the heart: chaos and synchronization in networks of cardiac cells

Ulrich Parlitz

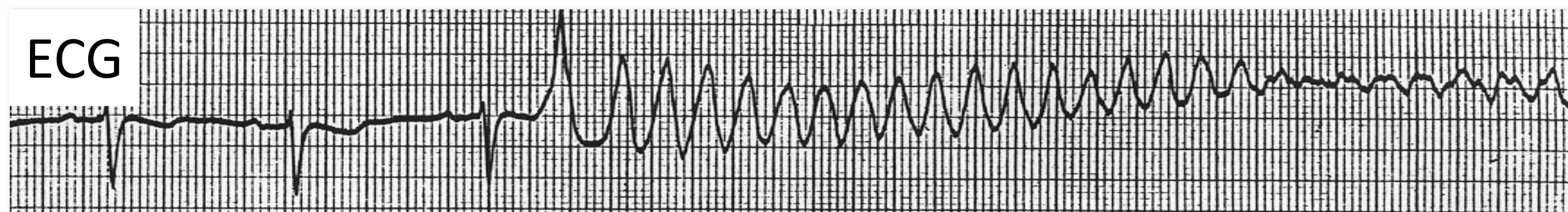
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Göttingen, Germany



Institute for the Dynamics of Complex Systems
University of Göttingen

Transitions to Cardiac Arrhythmias

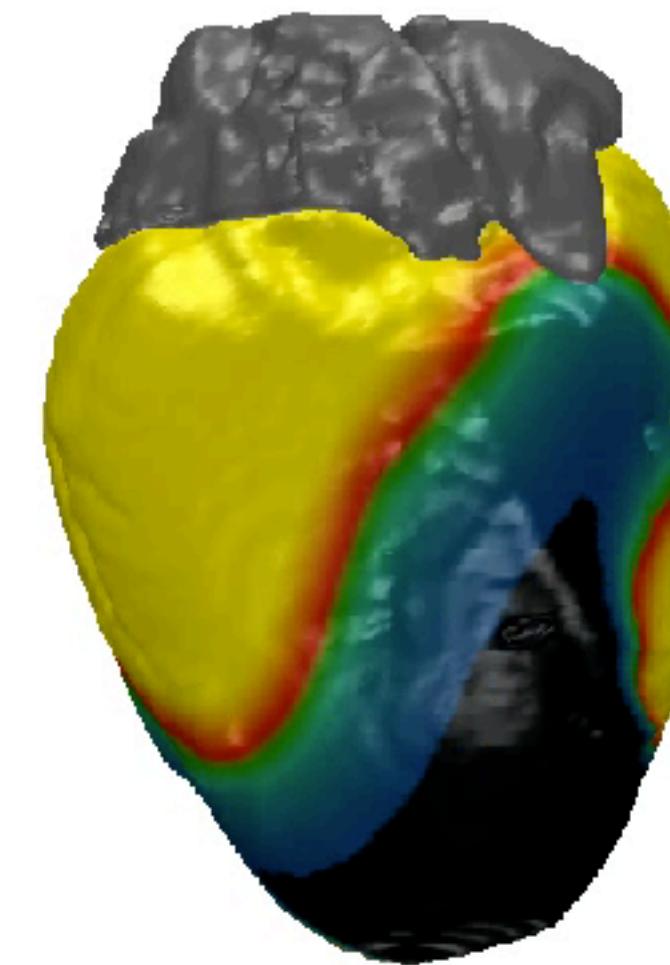
Normal Rhythm → Tachycardia → Fibrillation



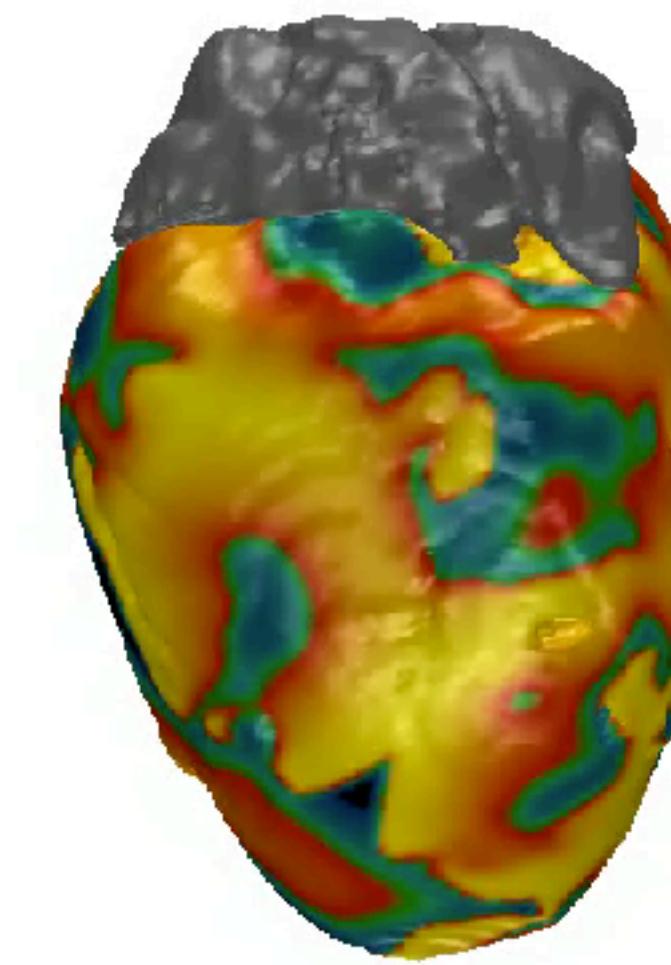
electrical excitation waves



plane waves



spiral waves



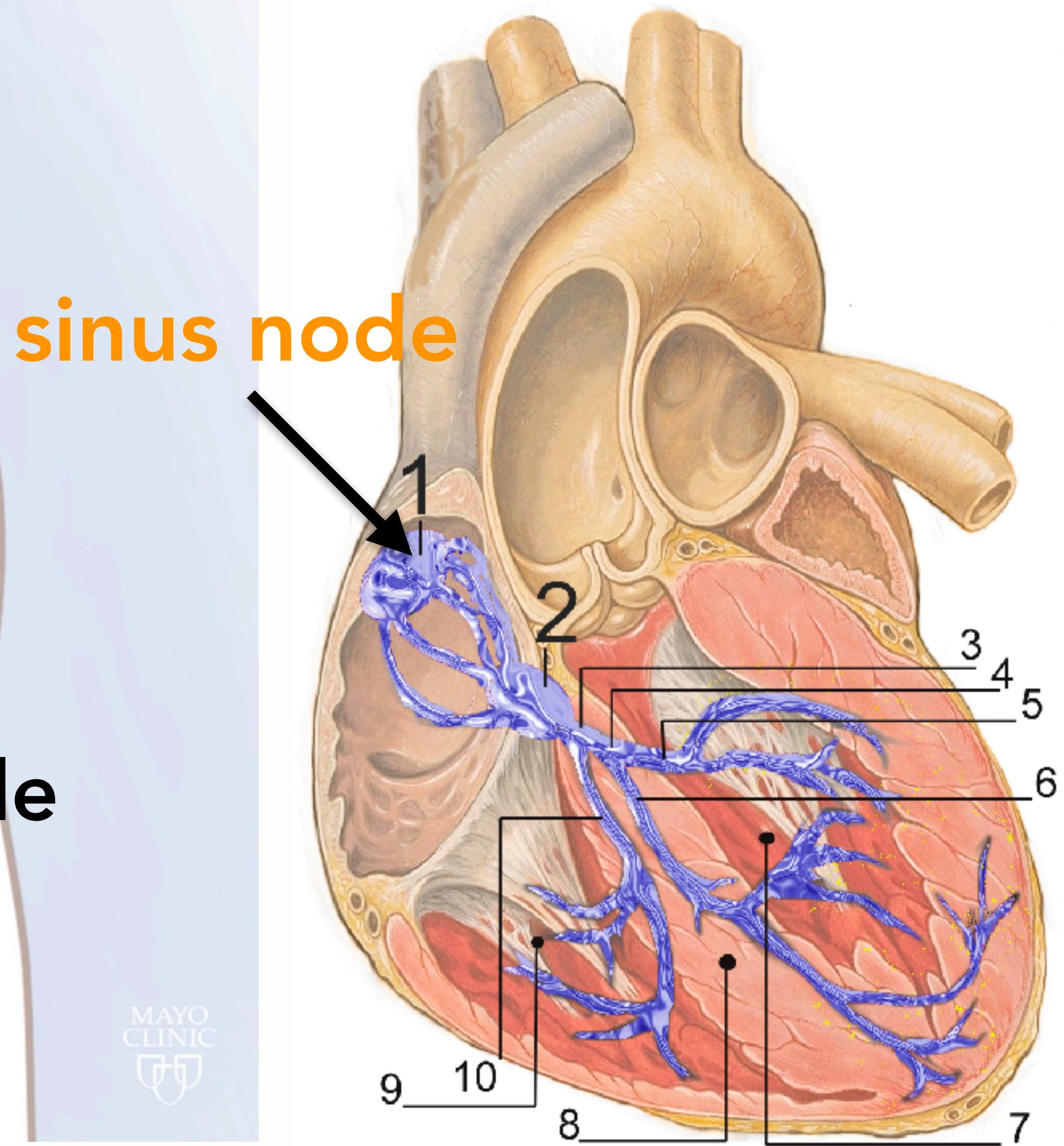
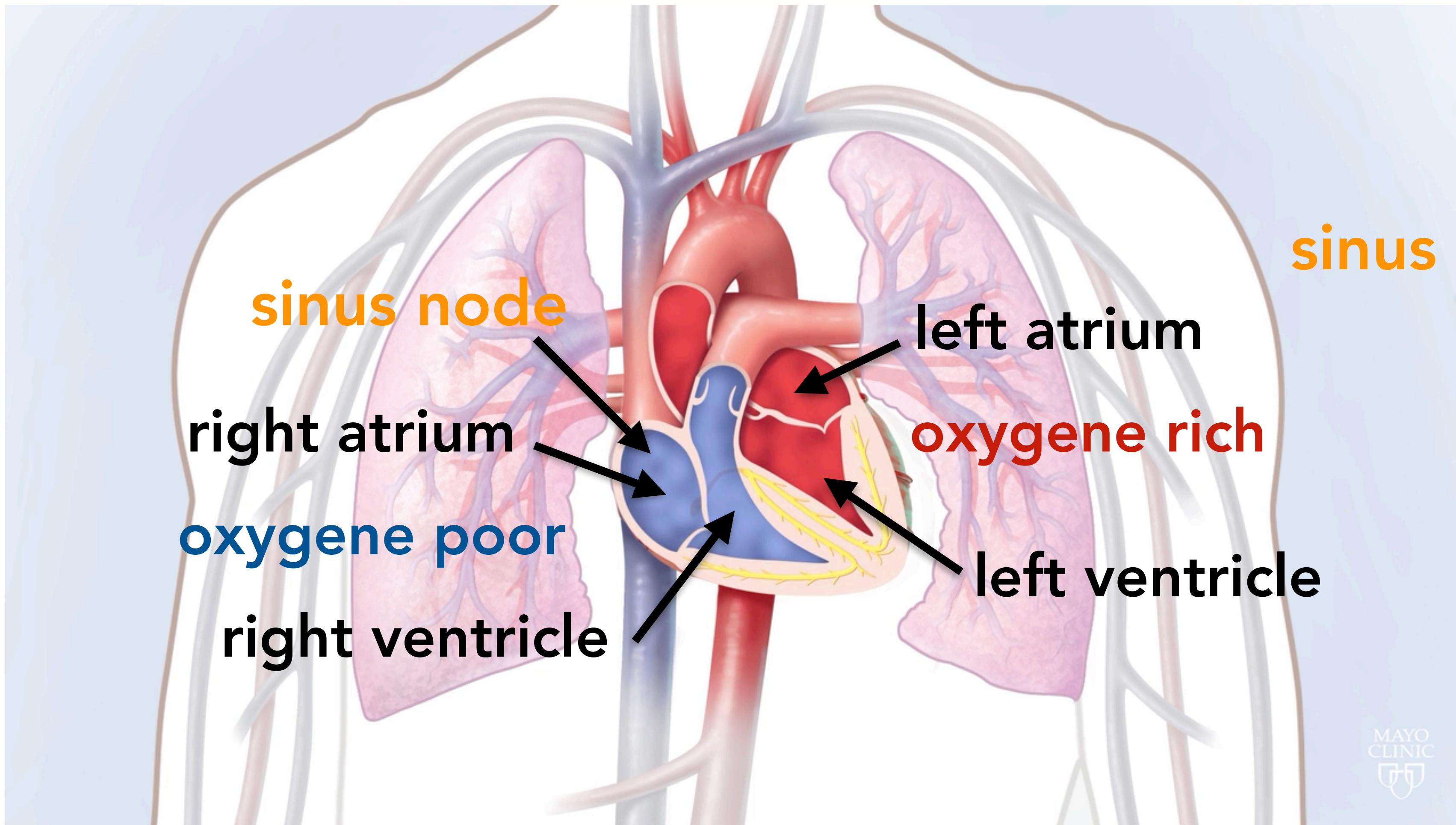
chaos

simulations: P. Bittihn

Outline

- the heart - a network of electrically and mechanically coupled contracting cardiac cells
- excitable media, (chaotic) spiral waves, and phase singularities
- virtual electrodes and low-energy defibrillation
- (transient) chaos and complexity in cardiac arrhythmias

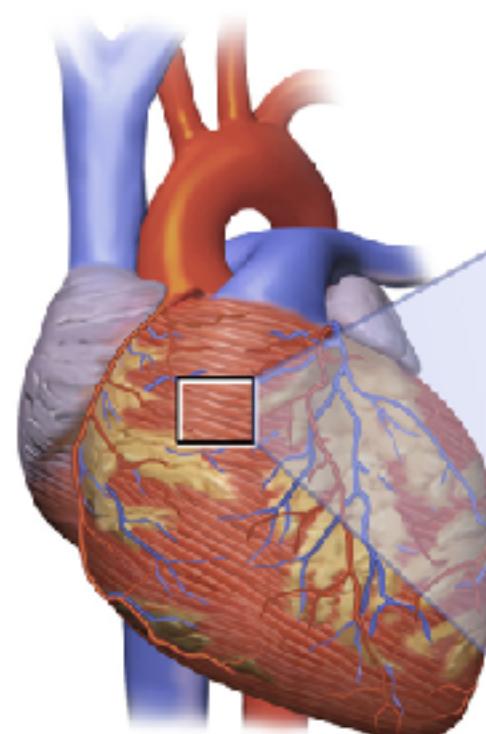
The Heart



<https://www.mayoclinic.org/diseases-conditions/heart-disease/multimedia/circulatory-system/vid-20084745>
J. Heuser, http://commons.wikimedia.org/wiki/File:RLS_12blauLeg.png
ISINP 2019

Network of Cardiomyocytes

cardiac muscle

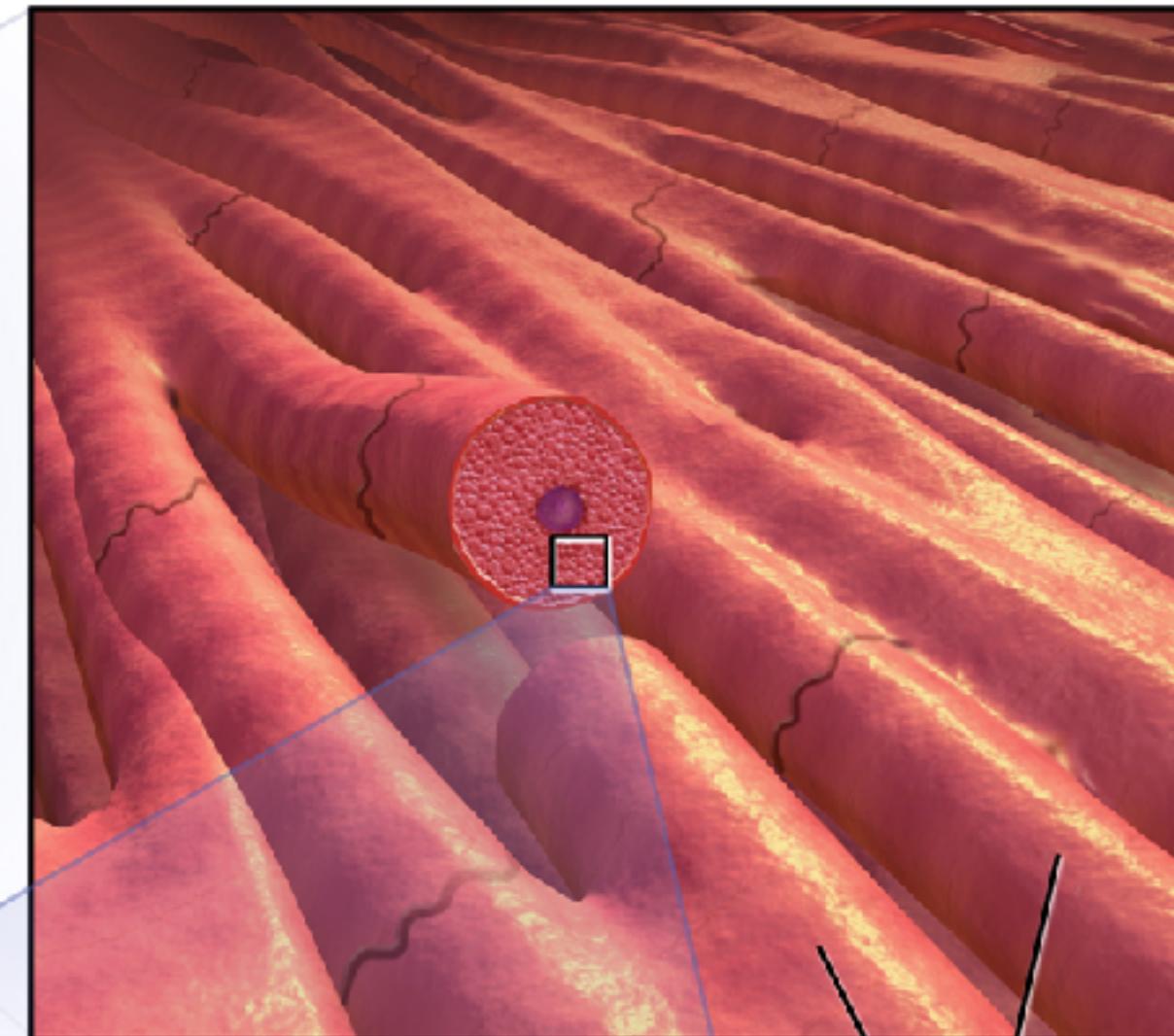
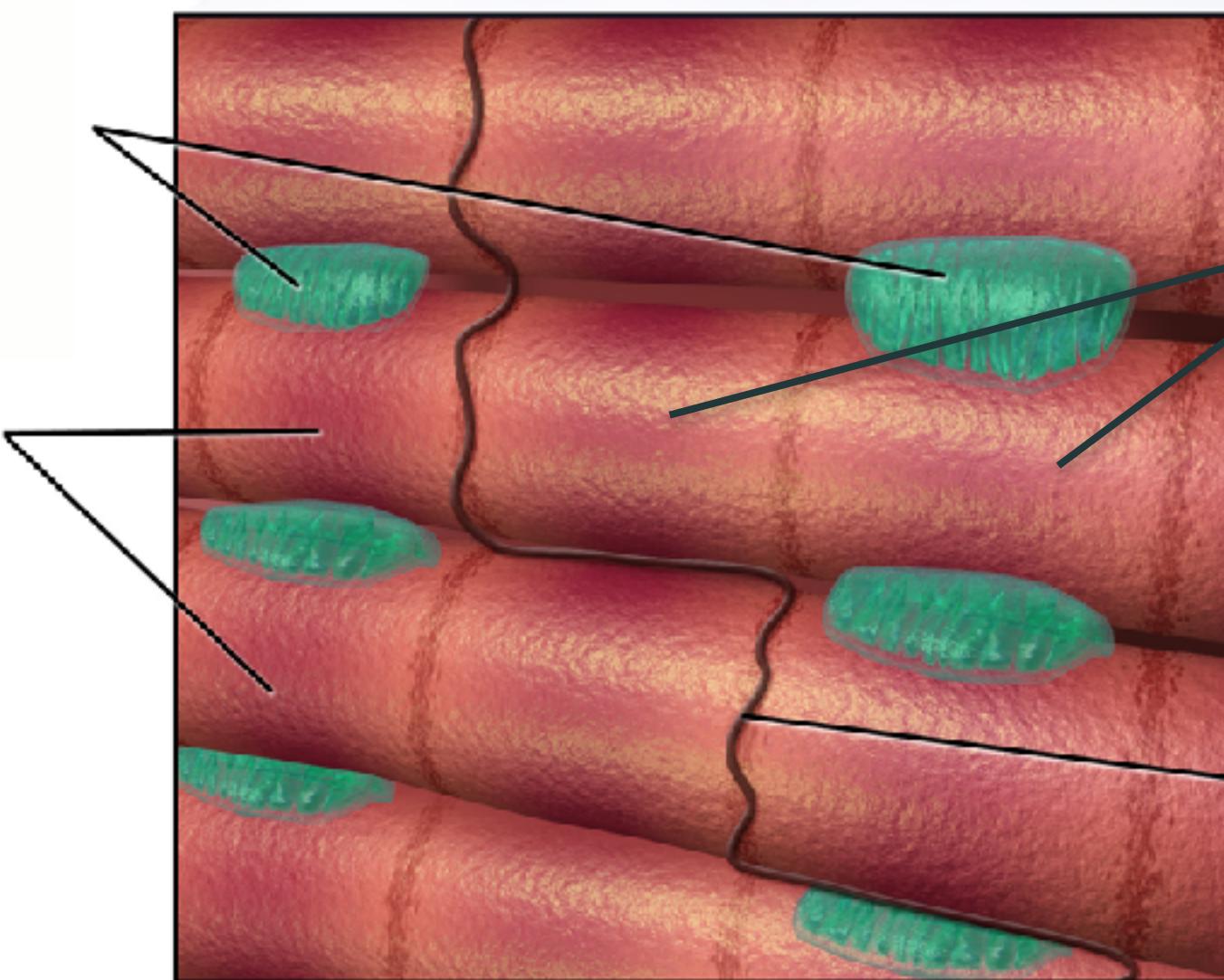


mitochondria

provide adenosine triphosphate (ATP) supply of the cell

myofibrils

provide mechanical contraction

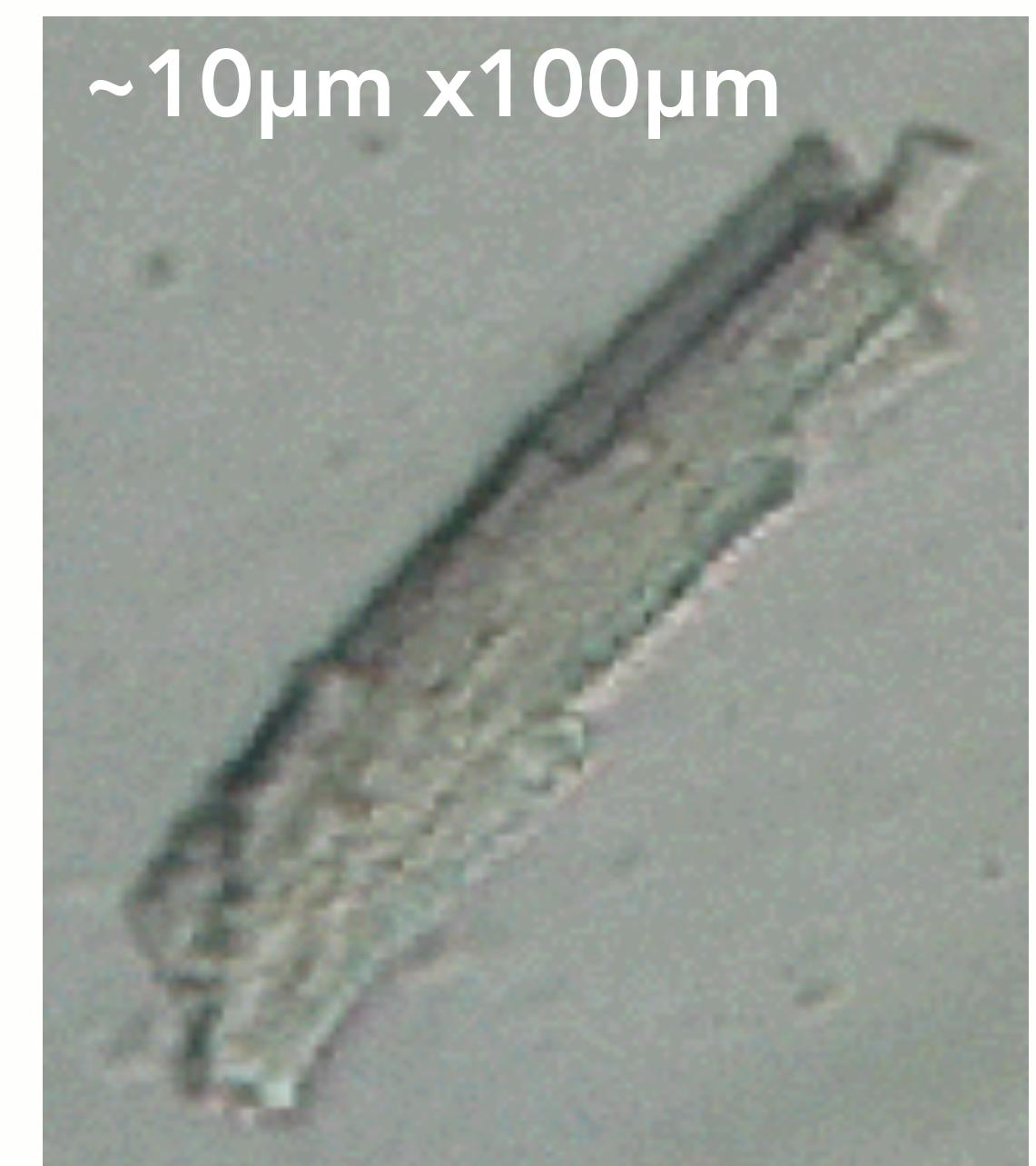


cardiac muscle fibers

BruceBlaus - Own work, CC BY-SA 4.0,
<https://commons.wikimedia.org/w/index.php?curid=44969447>

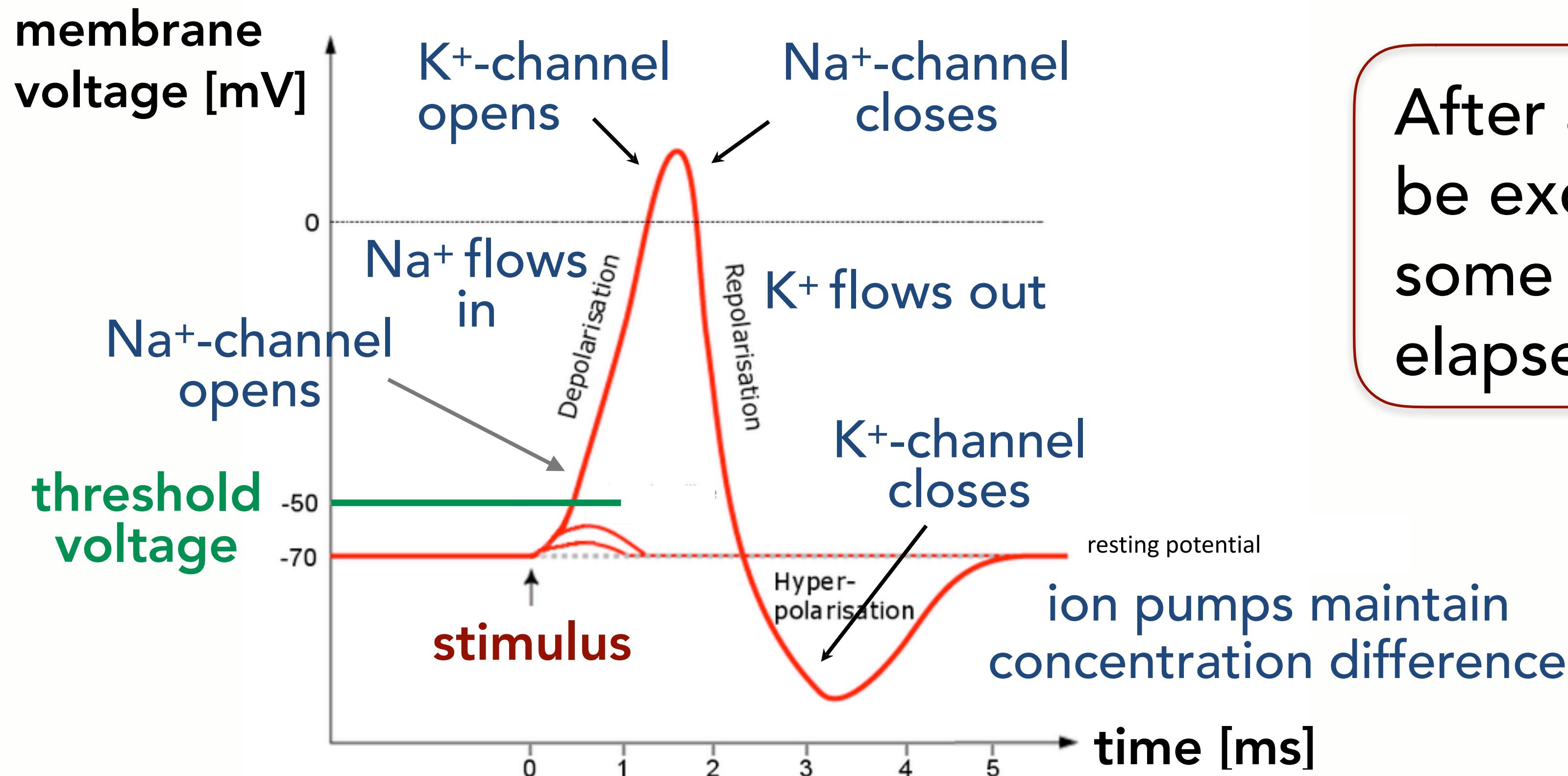
Ventricular Cell

~ $10\mu\text{m} \times 100\mu\text{m}$



© Kornreich & Fenton

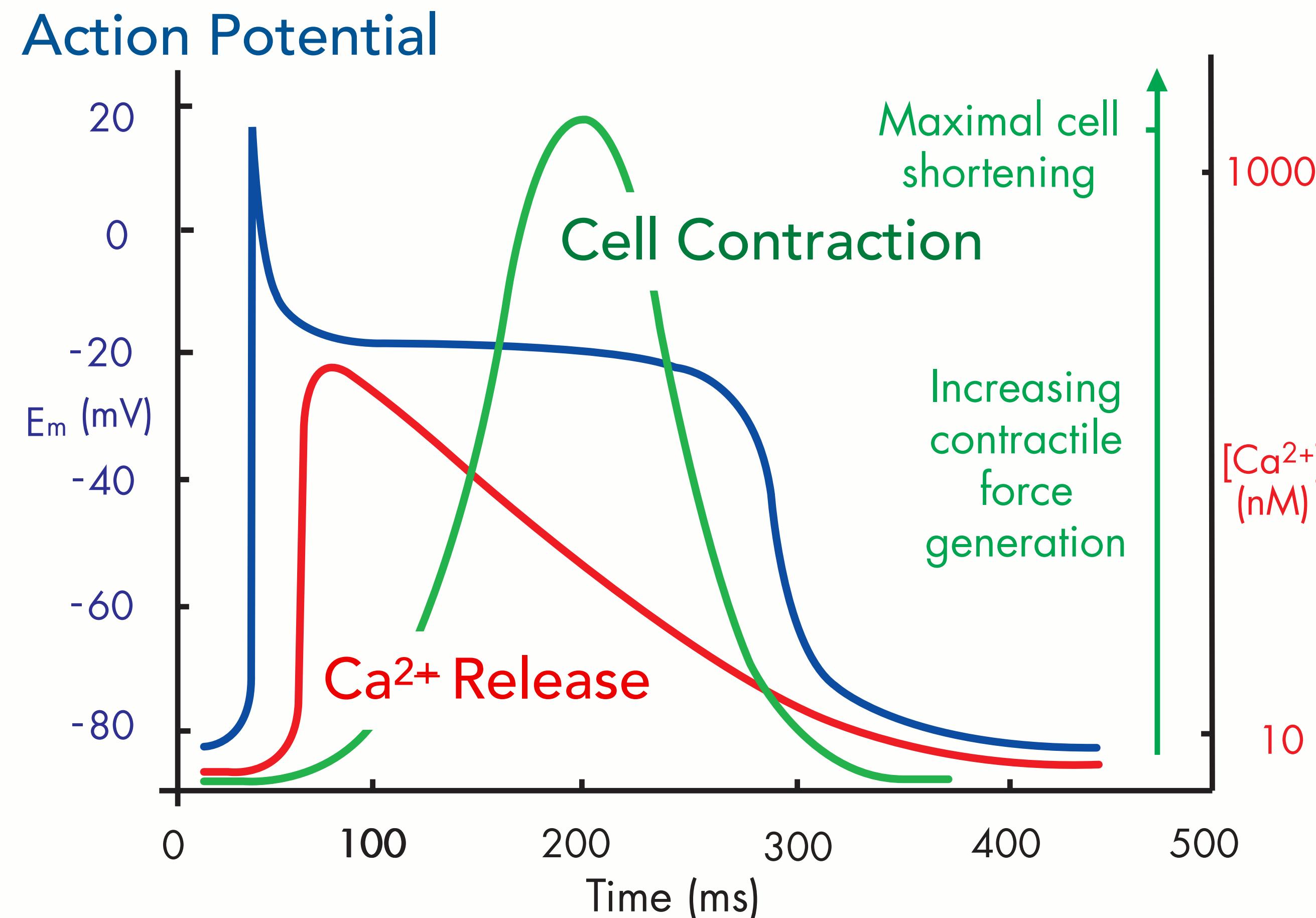
Generation of an Action Potential



After an excitation the cell can be excited again not before some **refractory phase** has elapsed.

adapted from Wikipedia

Excitation-Contraction Coupling



from: M. Scoote et al., Heart 89, 371–376 (2003)

electrical excitation

mechanical contraction

Mechanical perturbation
induces electrical stimulation via
stretch activated ion channels.

Mathematical Models of Cardiac Dynamics

continuum models averaging electrical behaviour of many cells

detailed ionic models: e.g., Luo-Rudy-II (15), Majahan (27), Bondarenko (44), ...

membrane
voltage

$$\frac{\partial V_m}{\partial t} = \nabla \cdot \underline{\mathbf{D}} \nabla V_m - I_{\text{ion}}(V_m, \mathbf{h})/C_m$$

ionic currents

$$\frac{\partial \mathbf{h}}{\partial t} = \mathbf{H}(V_m, \mathbf{h})$$

local cell dynamics (15-30 variables, 150 - 300 parameters!)

$$I_{\text{ion}}(V_m, \mathbf{h}) = \sum_x I_x(V_m, \mathbf{h}) + I_{\text{injection}}$$

generic qualitative models: e.g., Fenton-Karma (3), Beeler-Reuter (8), ...

simple qualitative models: e.g., Barkley (2), FitzHugh-Nagumo (2), Aliev-Panfilov (2), ...

see [Scholarpedia](#) article by F. Fenton and E. Cherry discussing 45 models of cardiac cells

Excitable Systems

Simple generic system: The Barkley model

$$\frac{\partial u}{\partial t} = \frac{1}{\epsilon} u(1-u)(u-u_{th})$$

$$\frac{\partial v}{\partial t} = u - v$$

controls
excitability
threshold

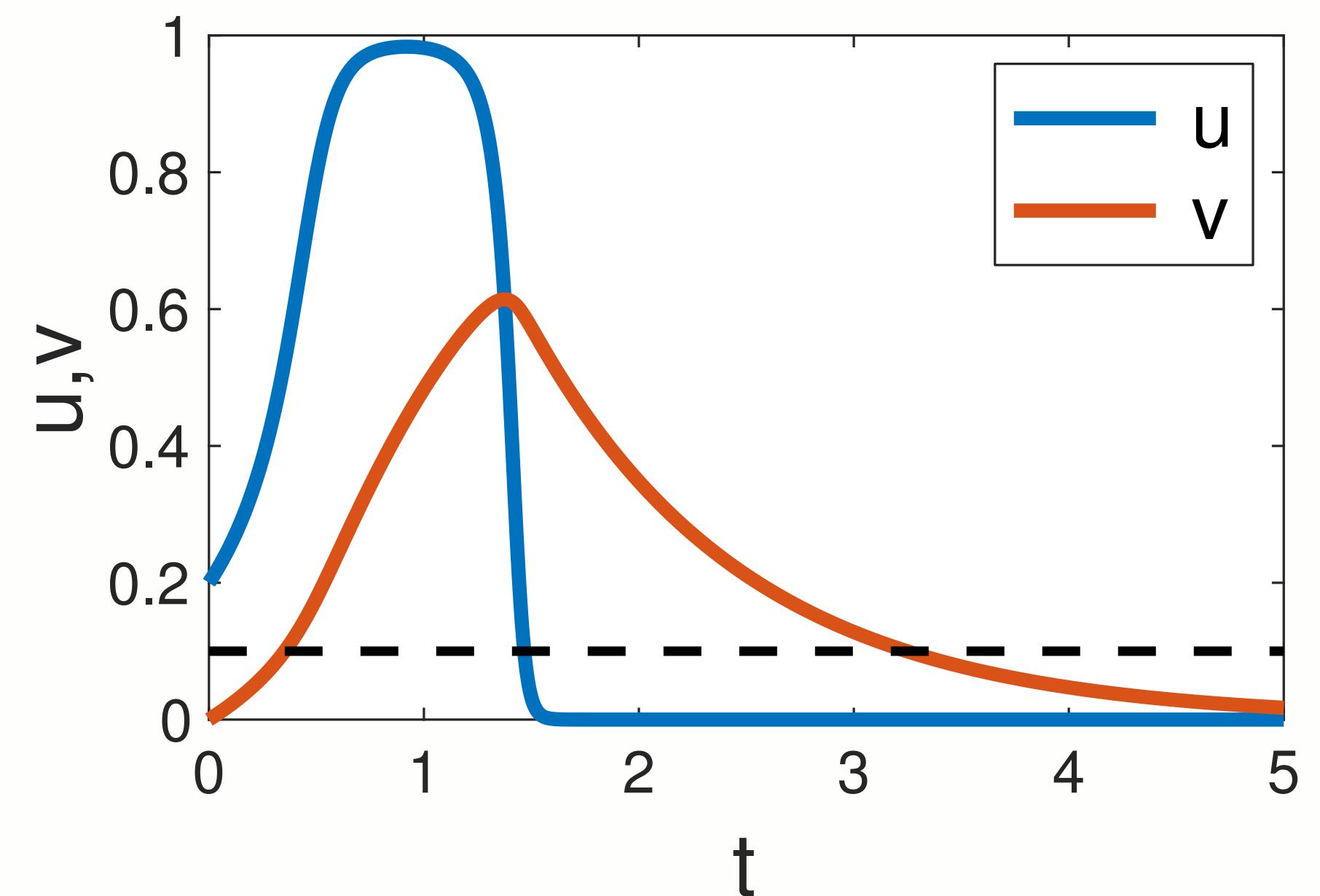
$1/\epsilon$ time scale of the fast variable u

a measure for action potential duration

b/a measure for excitation threshold

$$u_{th} = \frac{v + b}{a}$$

excitability
threshold



D. Barkley et al., Phys. Rev. A 4, 2489 (1990)

D. Barkley, Physica D 49, 6170 (1991)

http://www.scholarpedia.org/article/Barkley_model

The Barkley model

$$\begin{array}{lcl} \frac{\partial u}{\partial t} & = & \text{local dynamics} \\ & = & \frac{1}{\varepsilon} u(1-u)(u-u_{th}) + D \cdot \nabla^2 u \\ \frac{\partial v}{\partial t} & = & u - v \end{array} \quad \begin{array}{l} \text{diffusive coupling} \\ \text{with: } u_{th} = \frac{v+b}{a} \end{array}$$

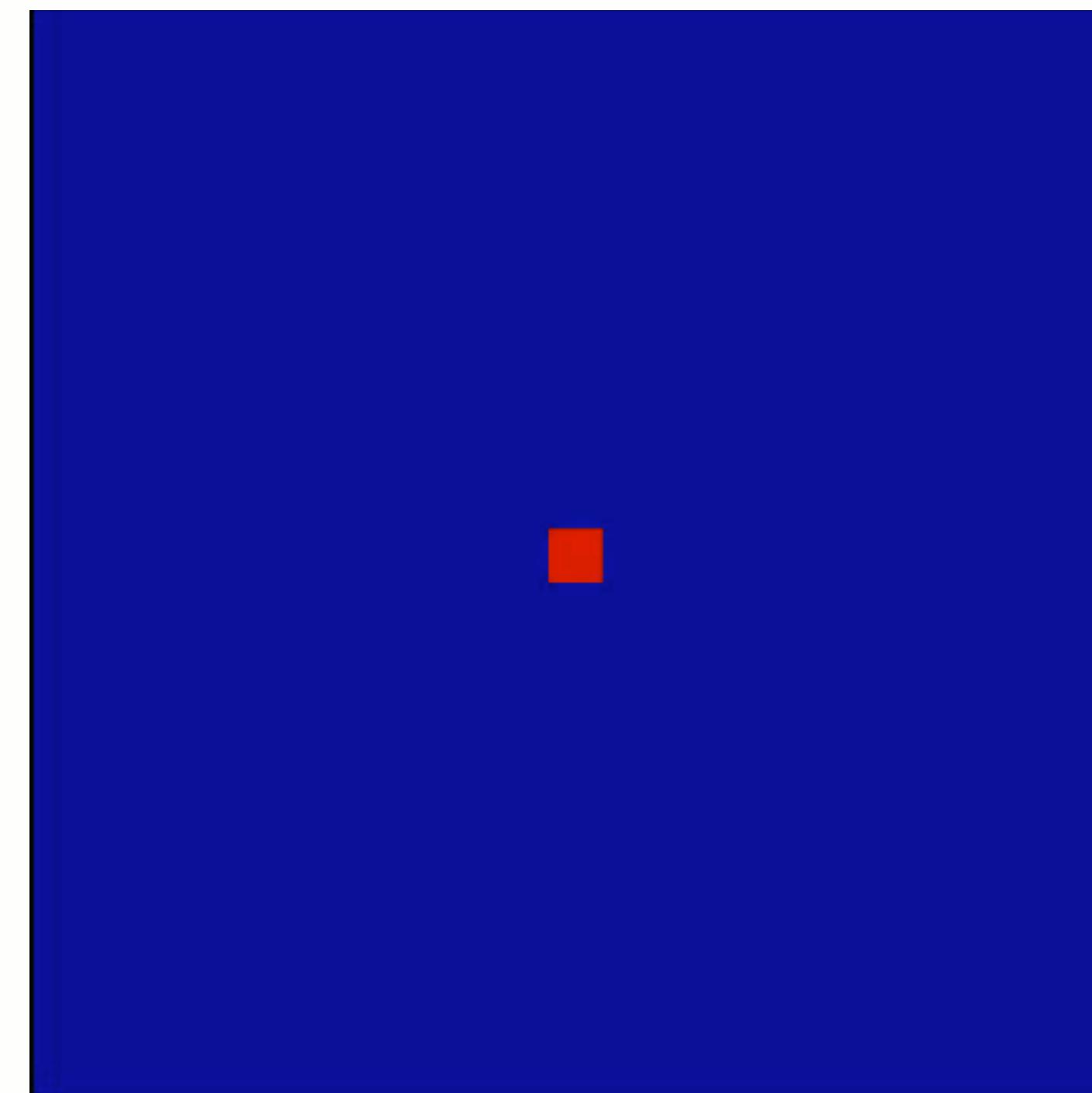
- $1/\varepsilon$ time scale of the fast variable u
- a measure for action potential duration
- b/a measure for excitation threshold

D. Barkley et al., Phys. Rev. A 4, 2489 (1990)
D. Barkley, Physica D 49, 6170 (1991)

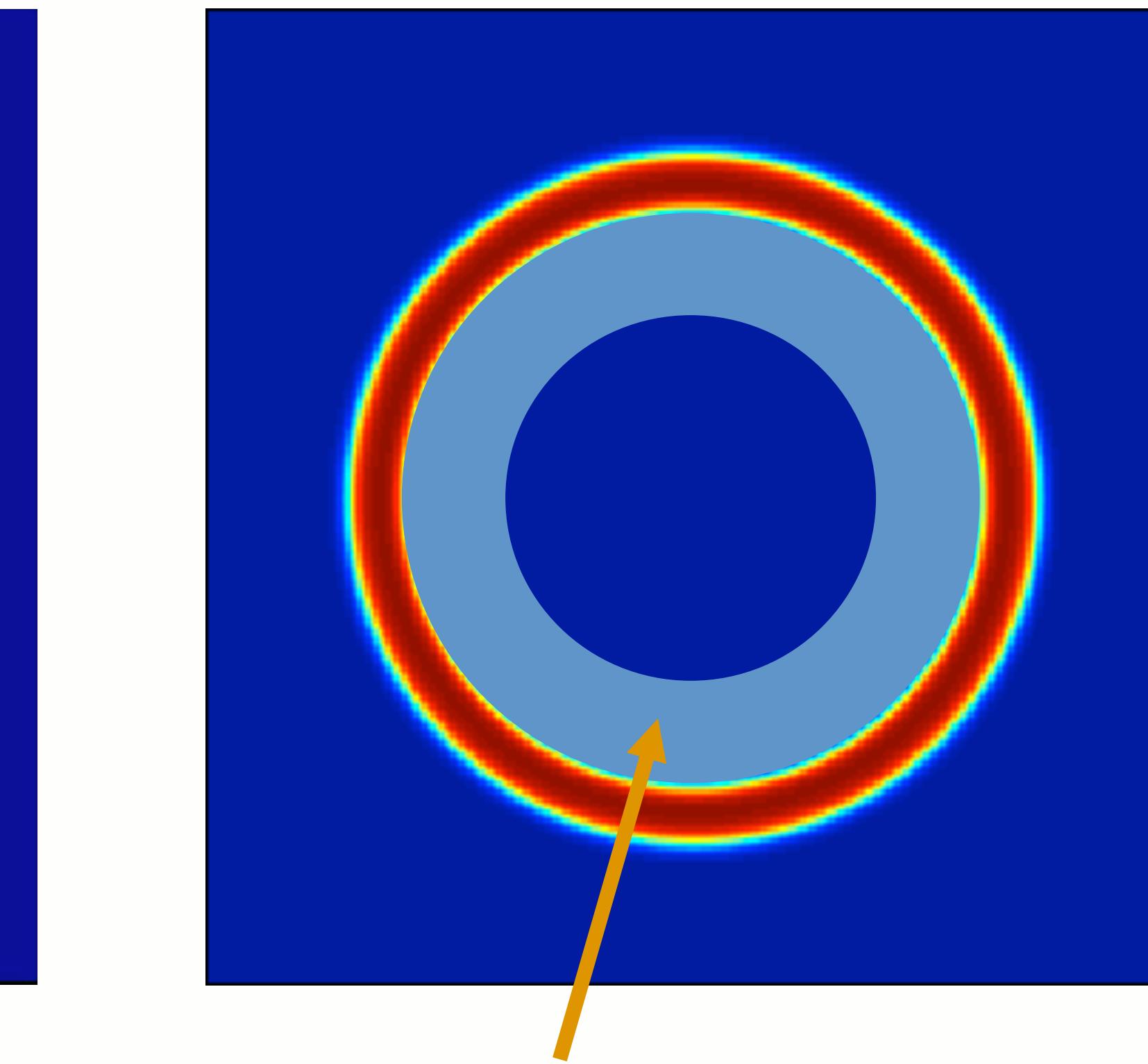
http://www.scholarpedia.org/article/Barkley_model

Excitation waves (Barkley model)

local stimulation
in the center



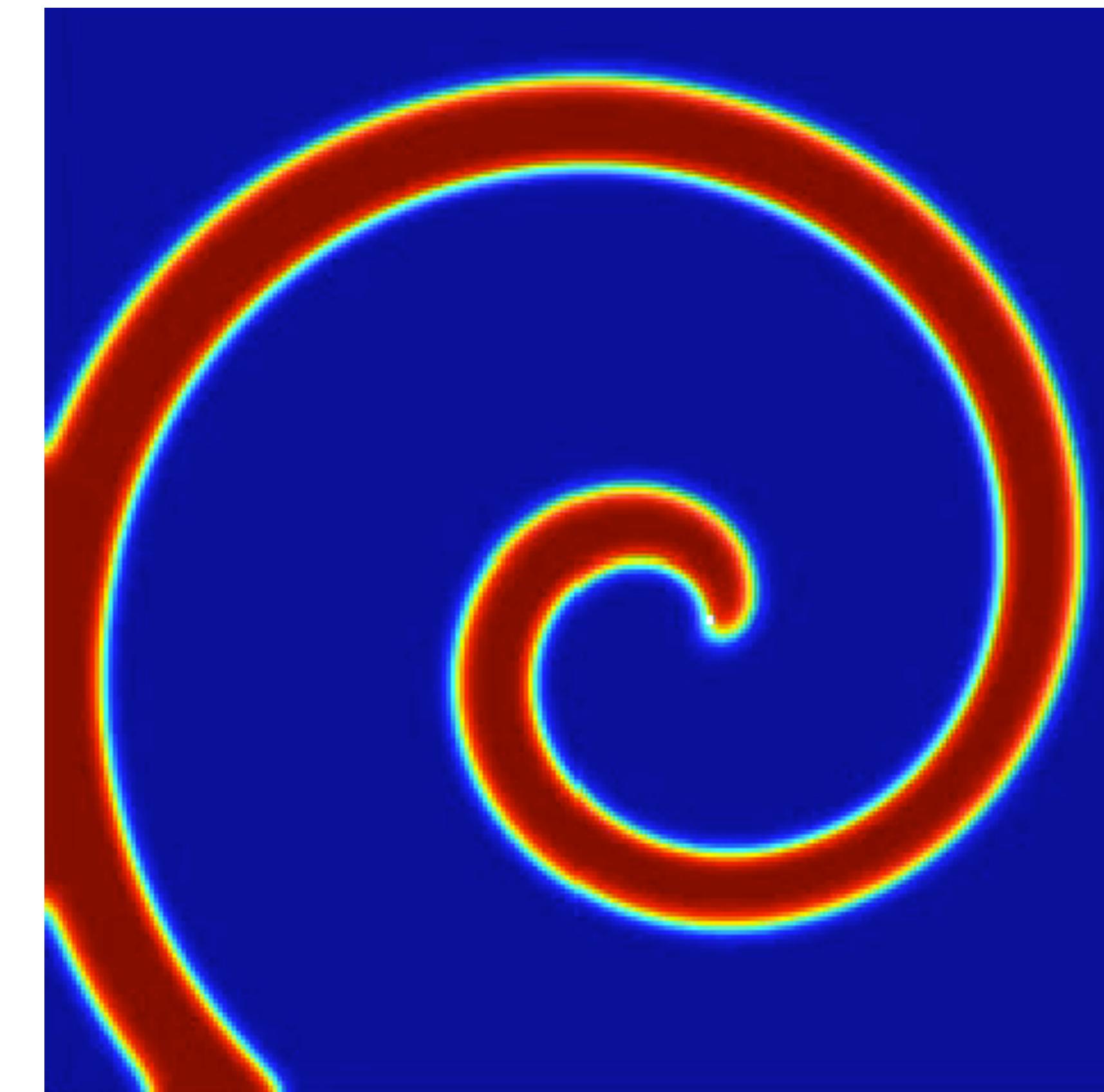
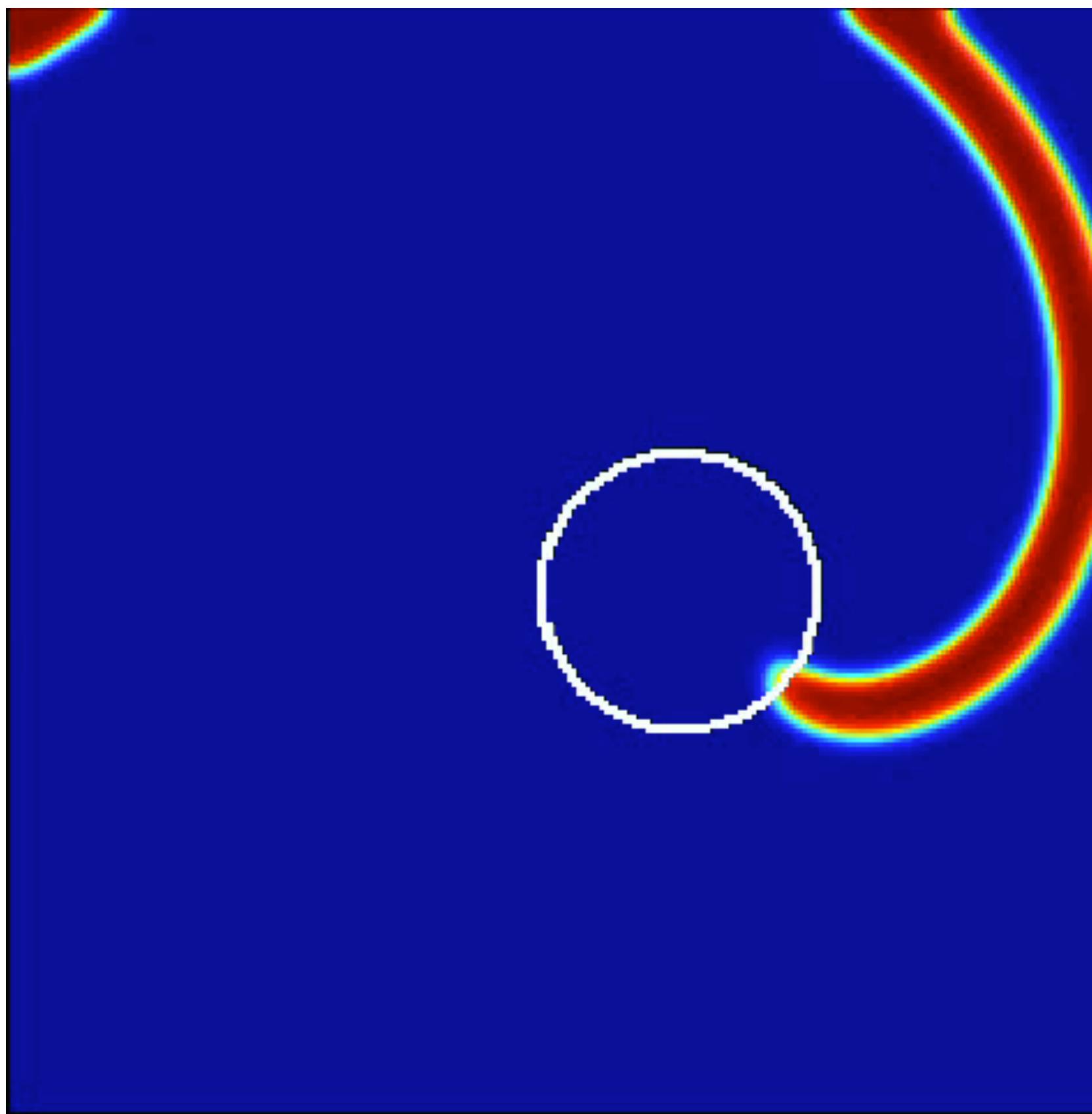
no flux boundary conditions



refractory region
(currently not excitable)

simulations: P. Bittihn

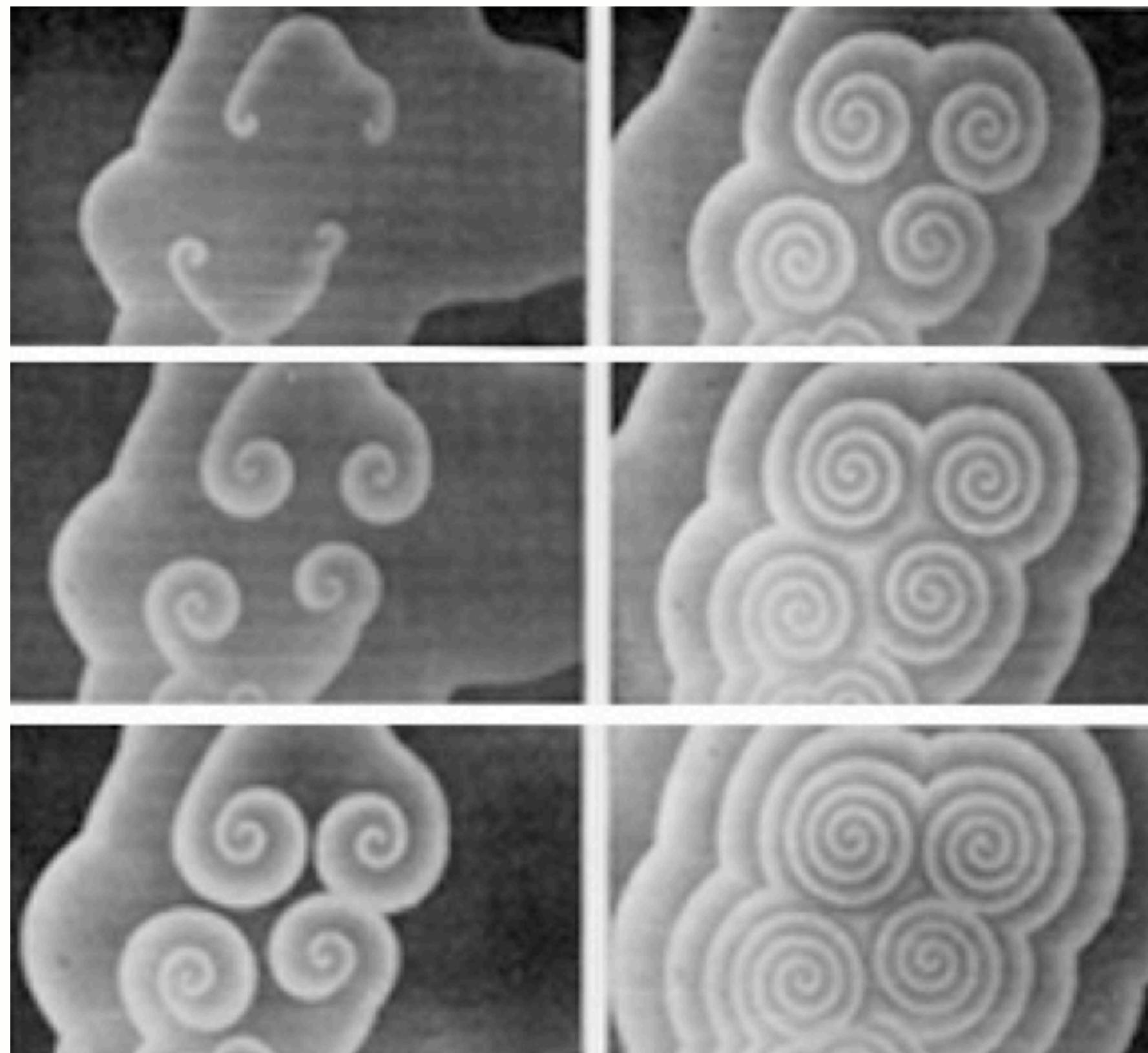
Spiral waves (Barkley model)



simulations: P. Bittihn

Excitable Media

The Belousov-Zhabotinsky (BZ) reaction



Development of spiral waves
after hydrodynamic breaking of
a concentric wave

www.scholarpedia.org

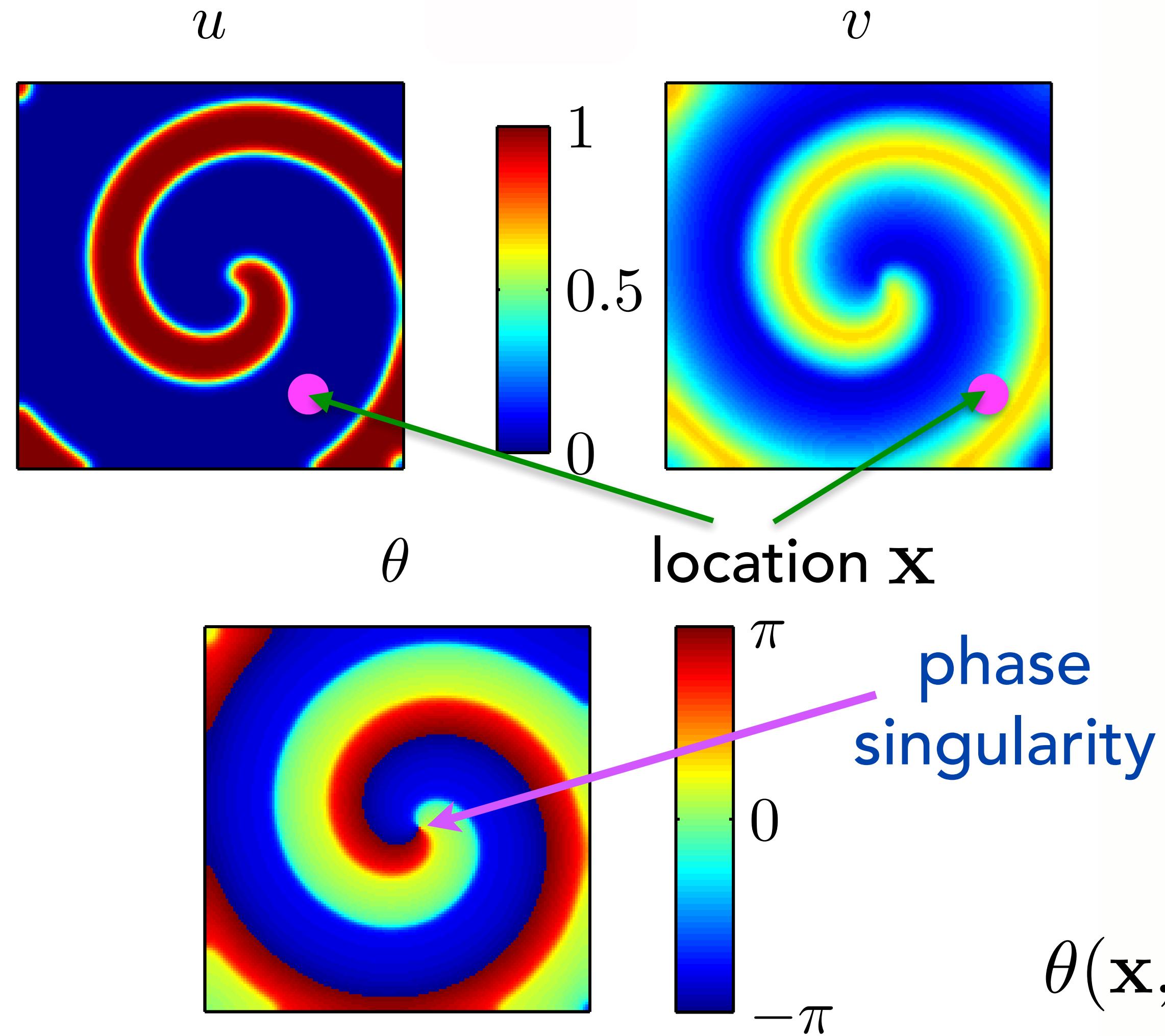
Geographic Tongue

inflammatory condition of the
mucous membrane of the tongue

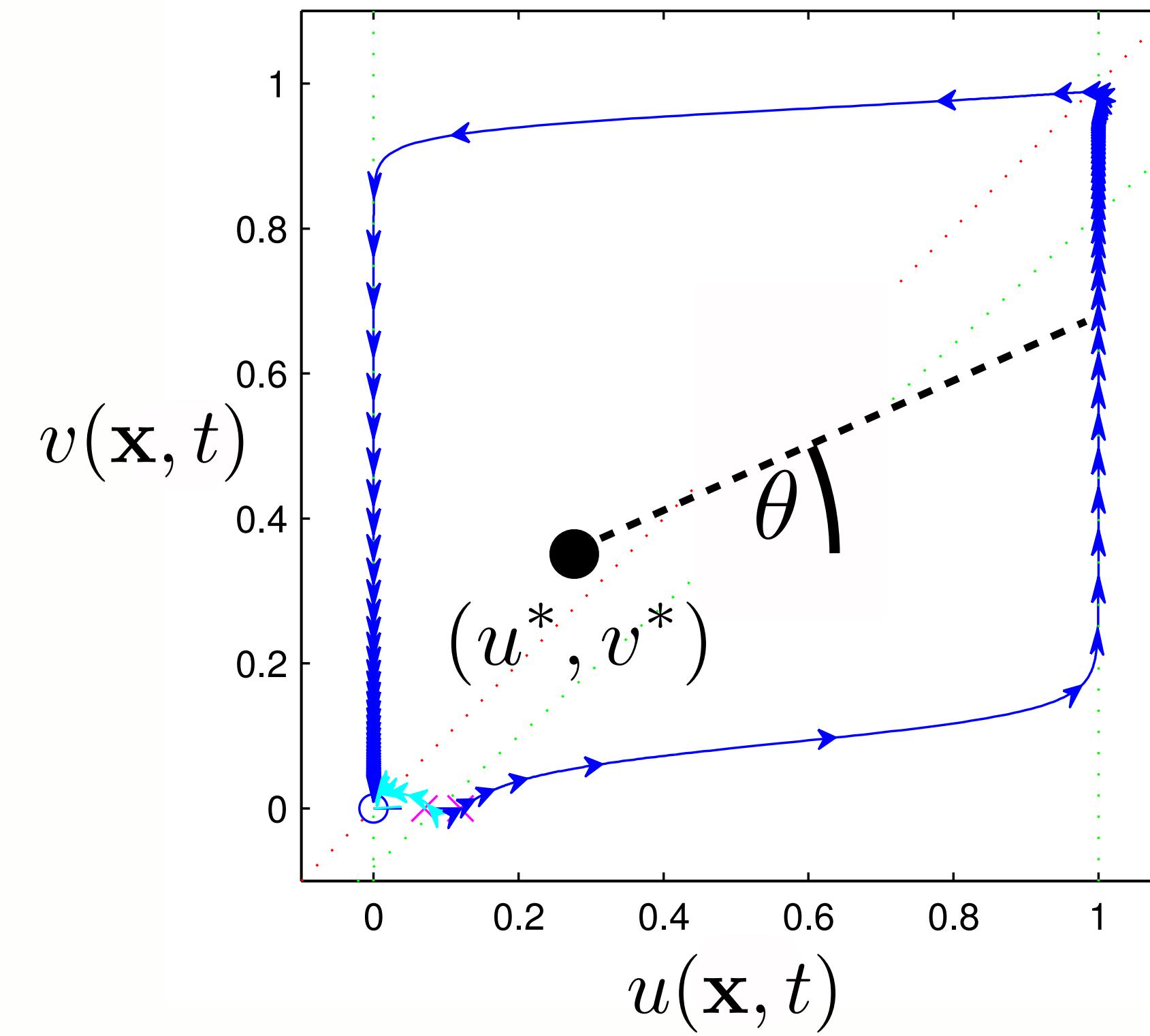


By Geographic_tongue.JPG: Martanopuederivative work: Jbarta -
This file was derived from: Geographic tongue.JPG;, CC BY-SA 3.0,
<https://commons.wikimedia.org/w/index.php?curid=24437119>

Spiral Tips and Phase Singularities

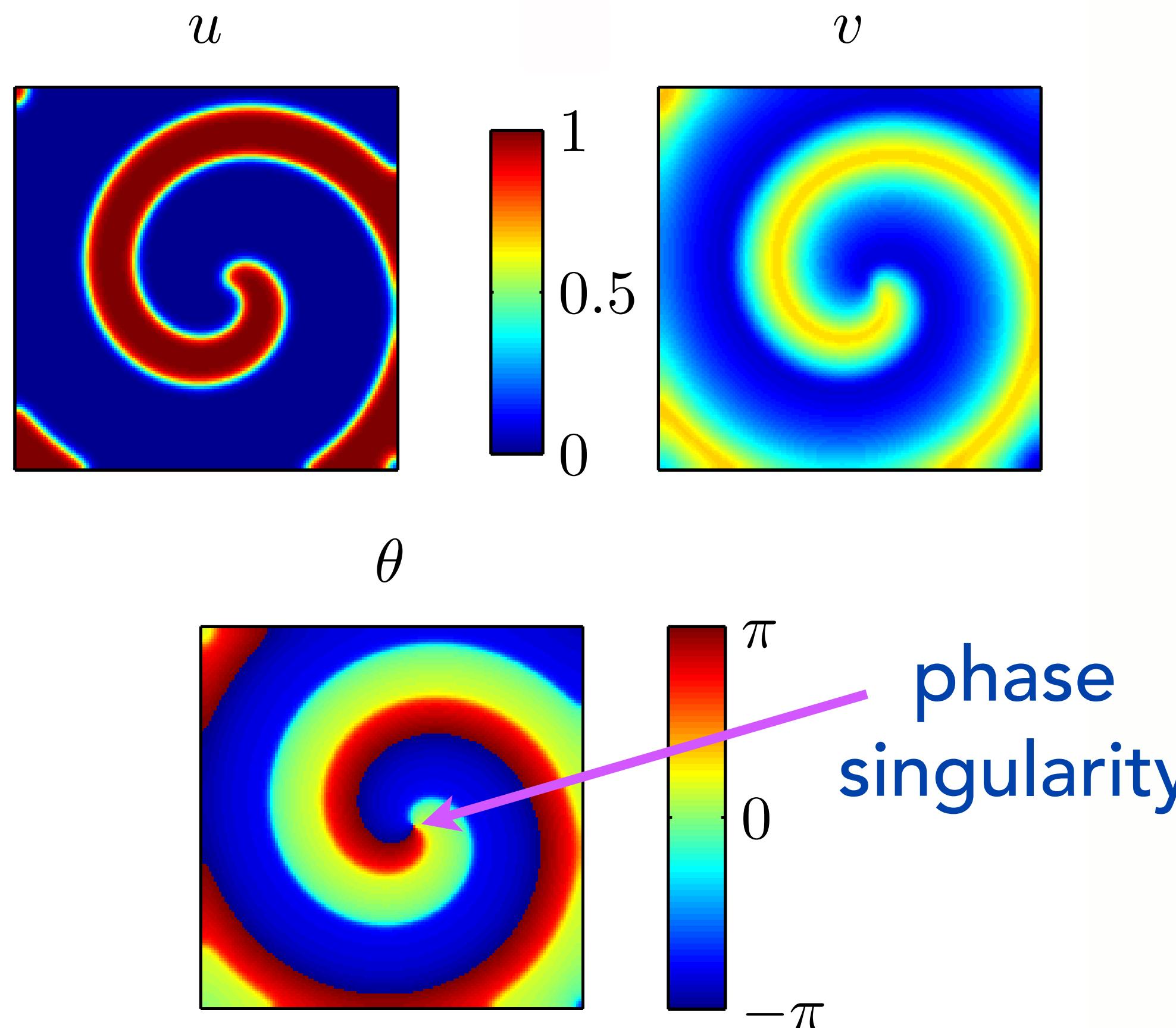


estimate phase at each location x

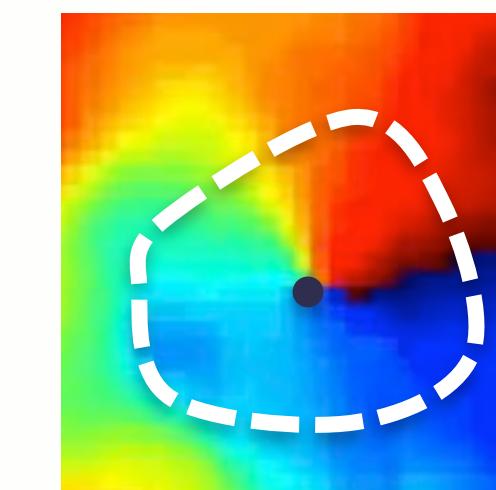


$$\theta(x, t) = \arctan 2(u(x, t) - u^*, v(x, t) - v^*)$$

Spiral Tips and Phase Singularities



sum of the topological charges in a domain \mathcal{D}



$$\oint_{\partial \mathcal{D}} \vec{\nabla} \theta \cdot d\vec{l} = 2\pi(n - m)$$

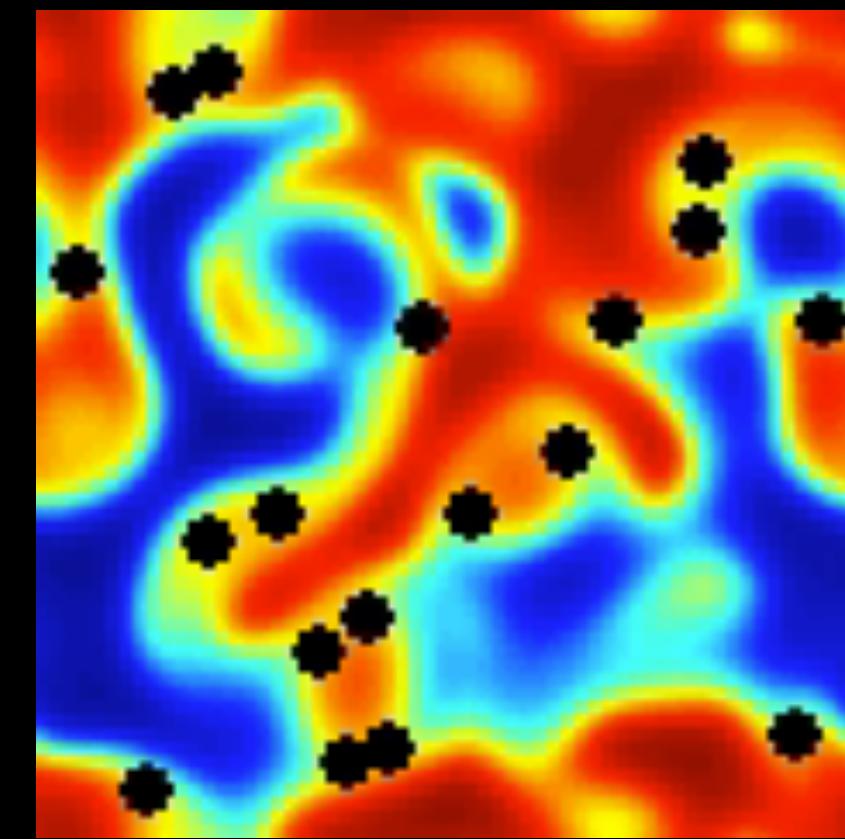
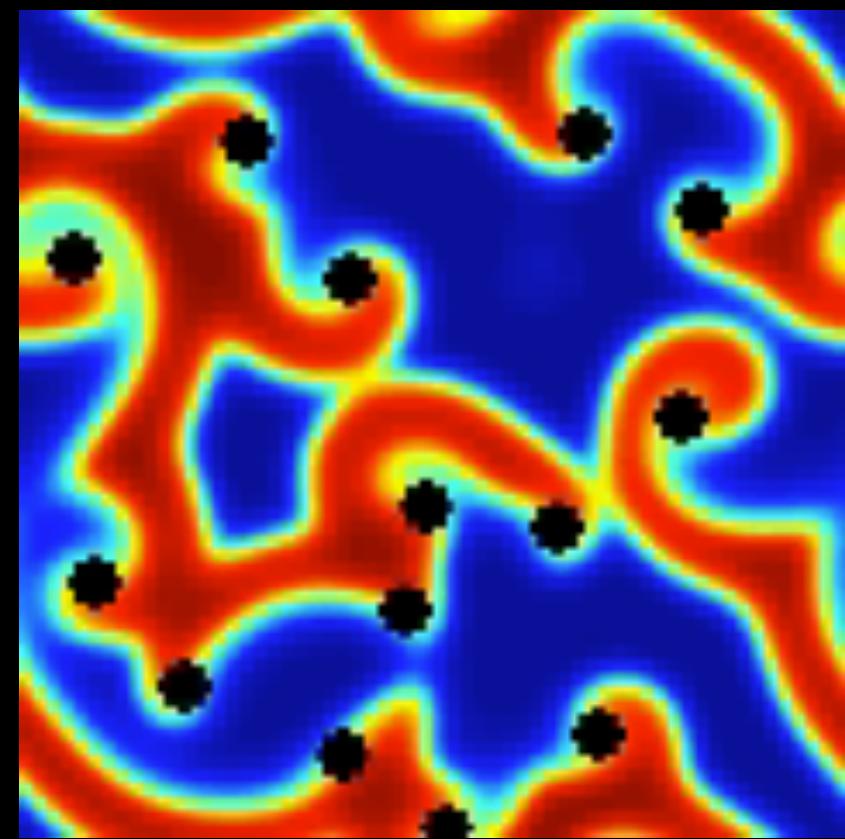
n # clockwise
 m # counter clockwise

rotating spirals

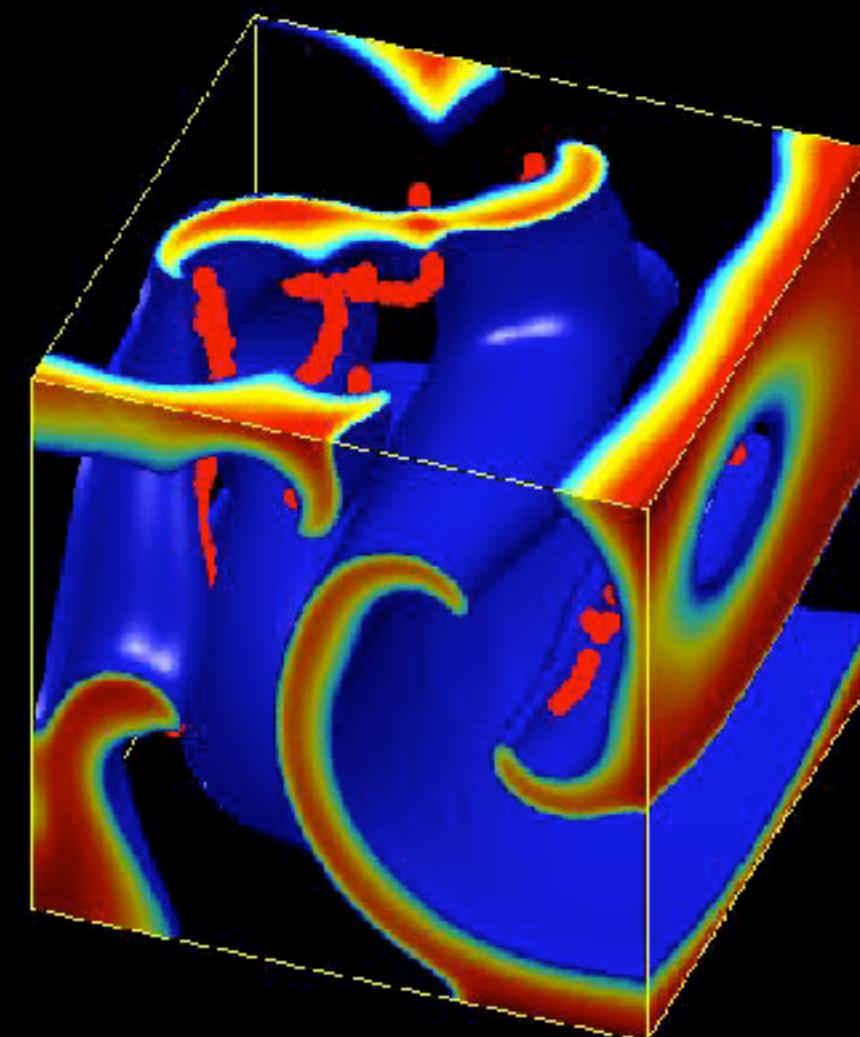
alternative approach: D.R. Gurevich and R.O. Grigoriev, Chaos 29, 053101 (2019)

Dynamics of Phase Singularities

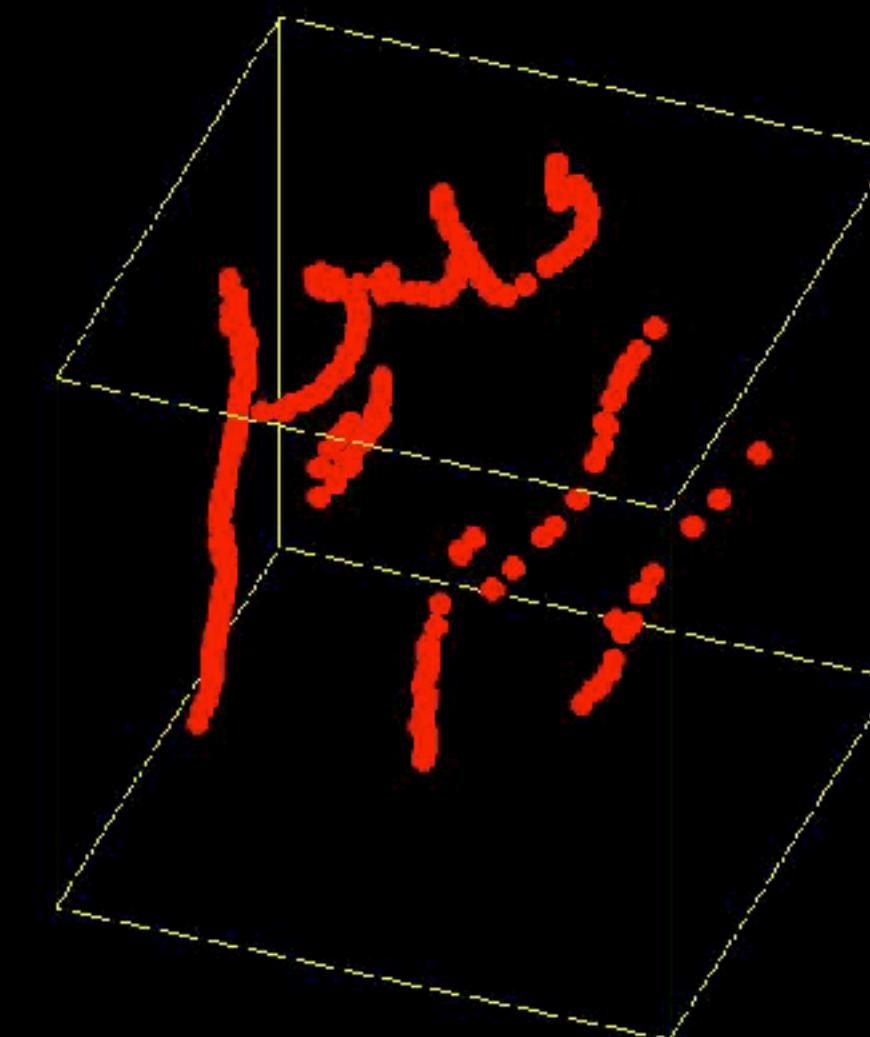
2D



3D



scroll wave

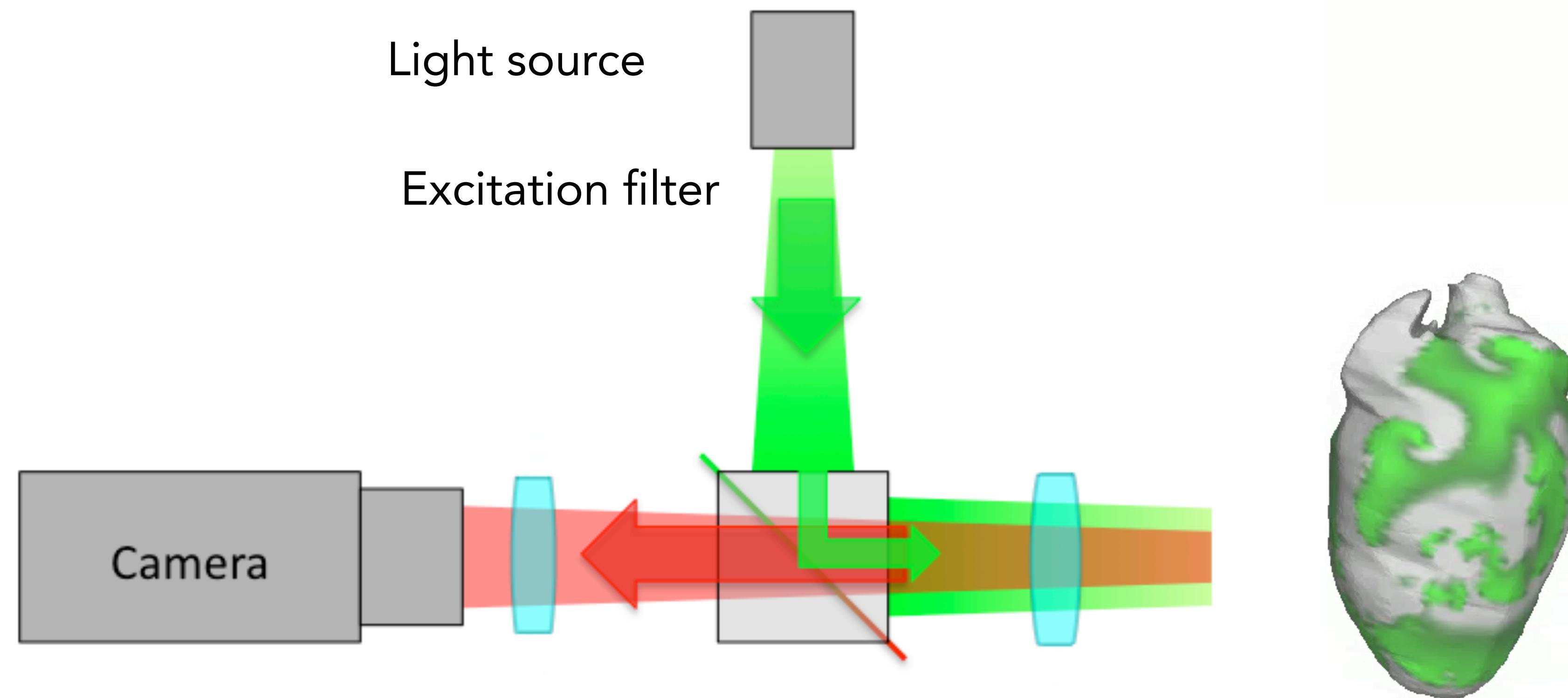


filaments

F. Fenton, E. Cherry
thevirtualheart.org
WebGL simulations

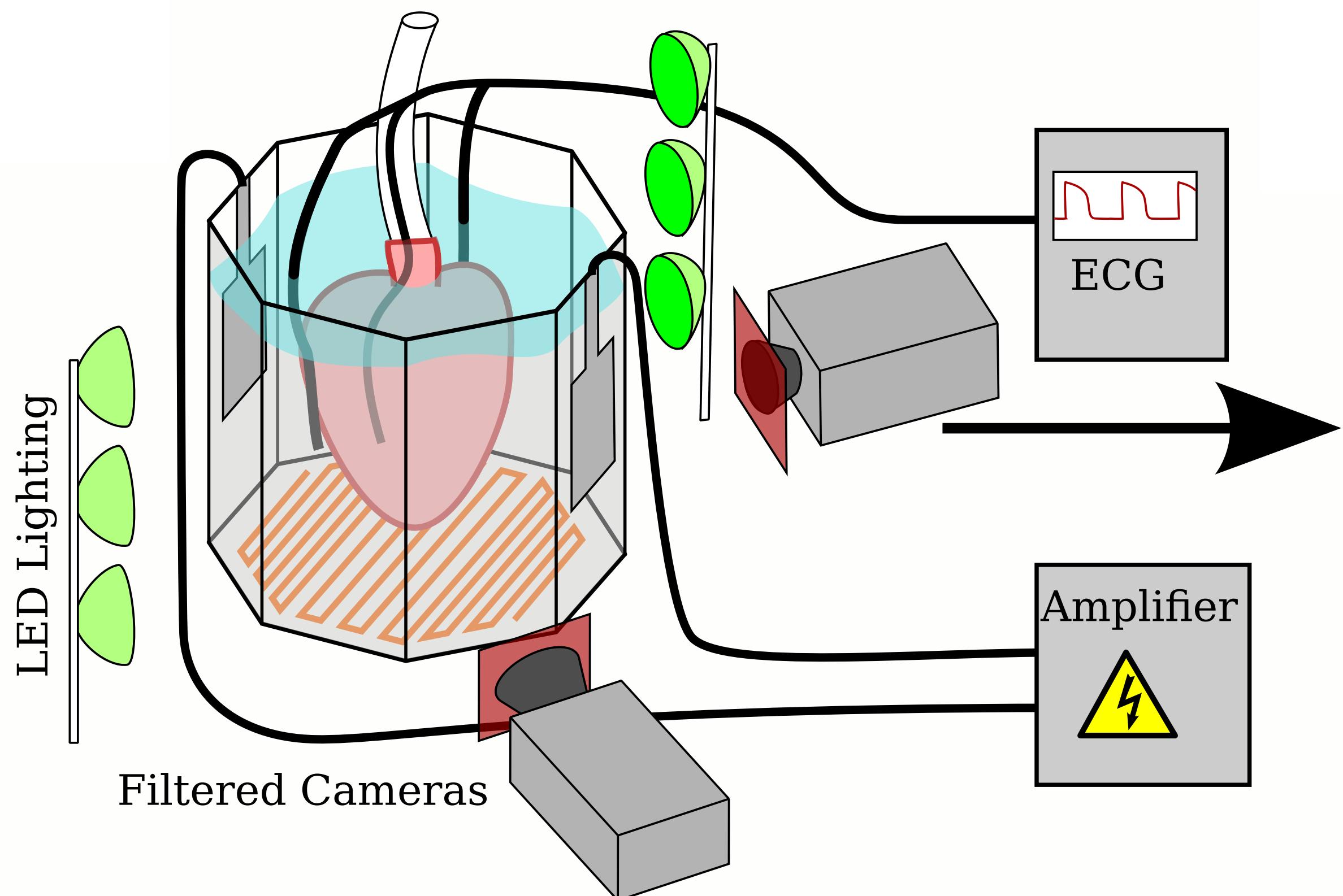
Optical Mapping

Visualisation of **membrane voltage** and **Ca⁺ concentration**
on the **surface of the heart** using **fluorescent dyes**

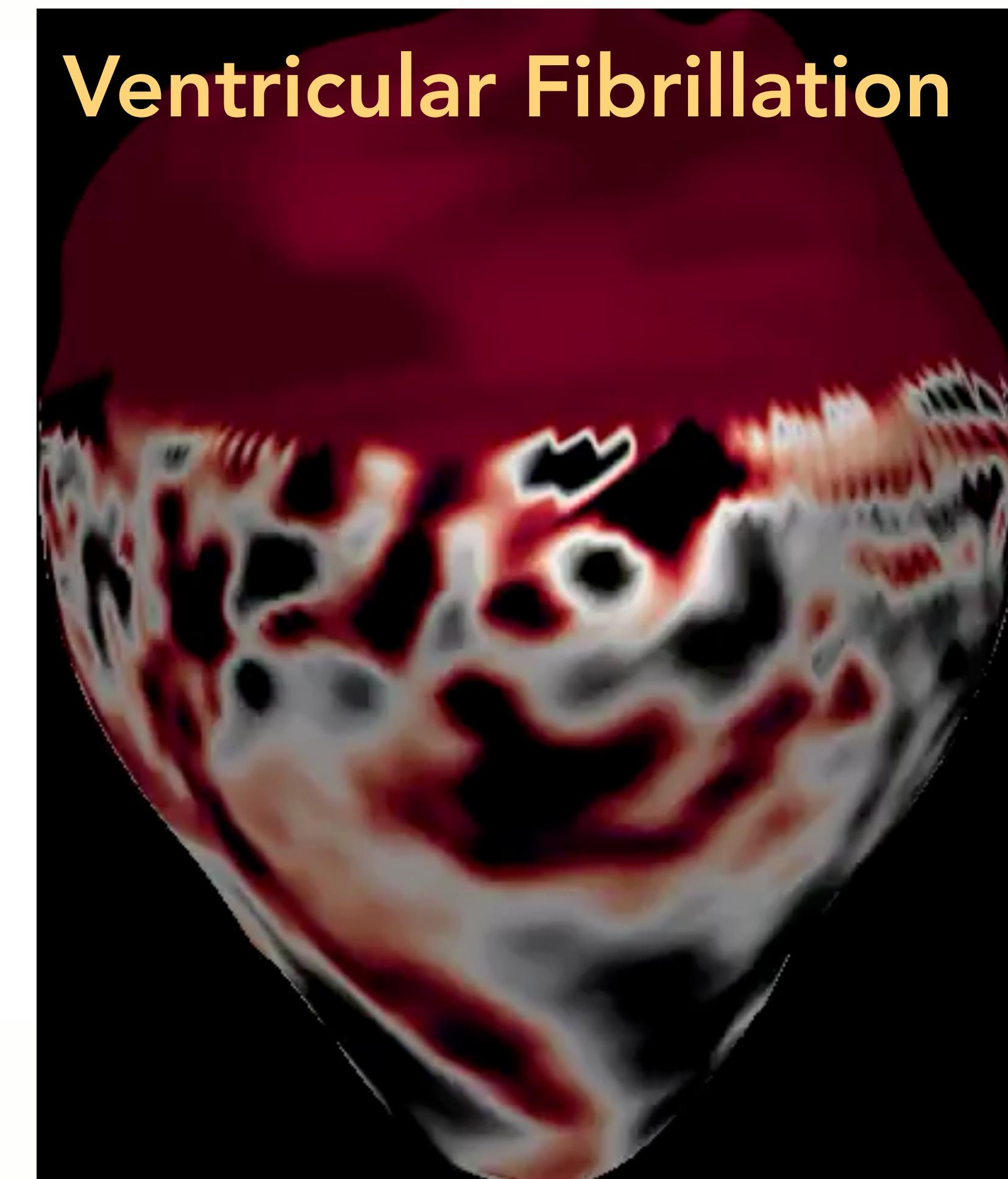


Optical mapping in Langendorff perfusion system

using voltage sensitive fluorescent dyes

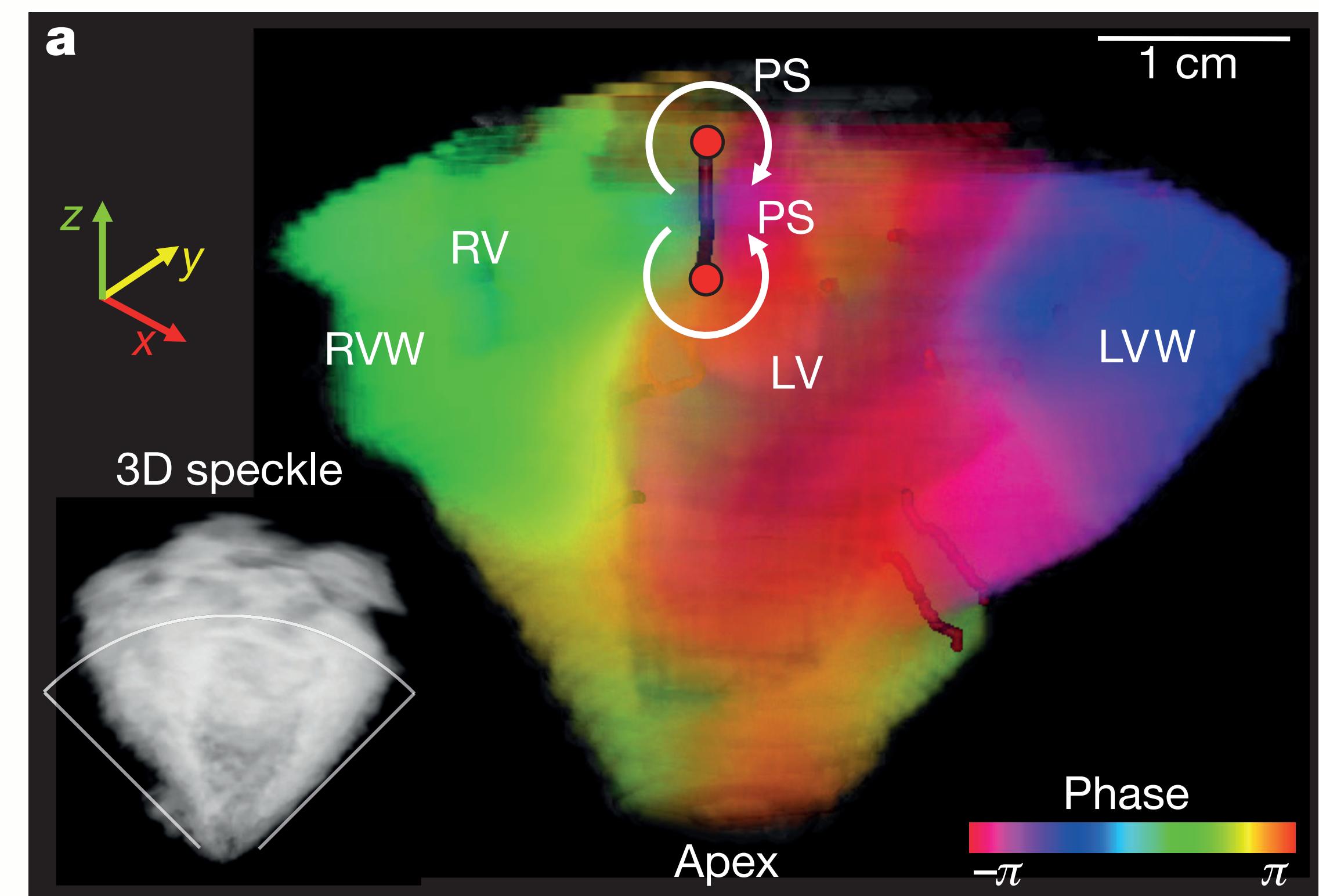
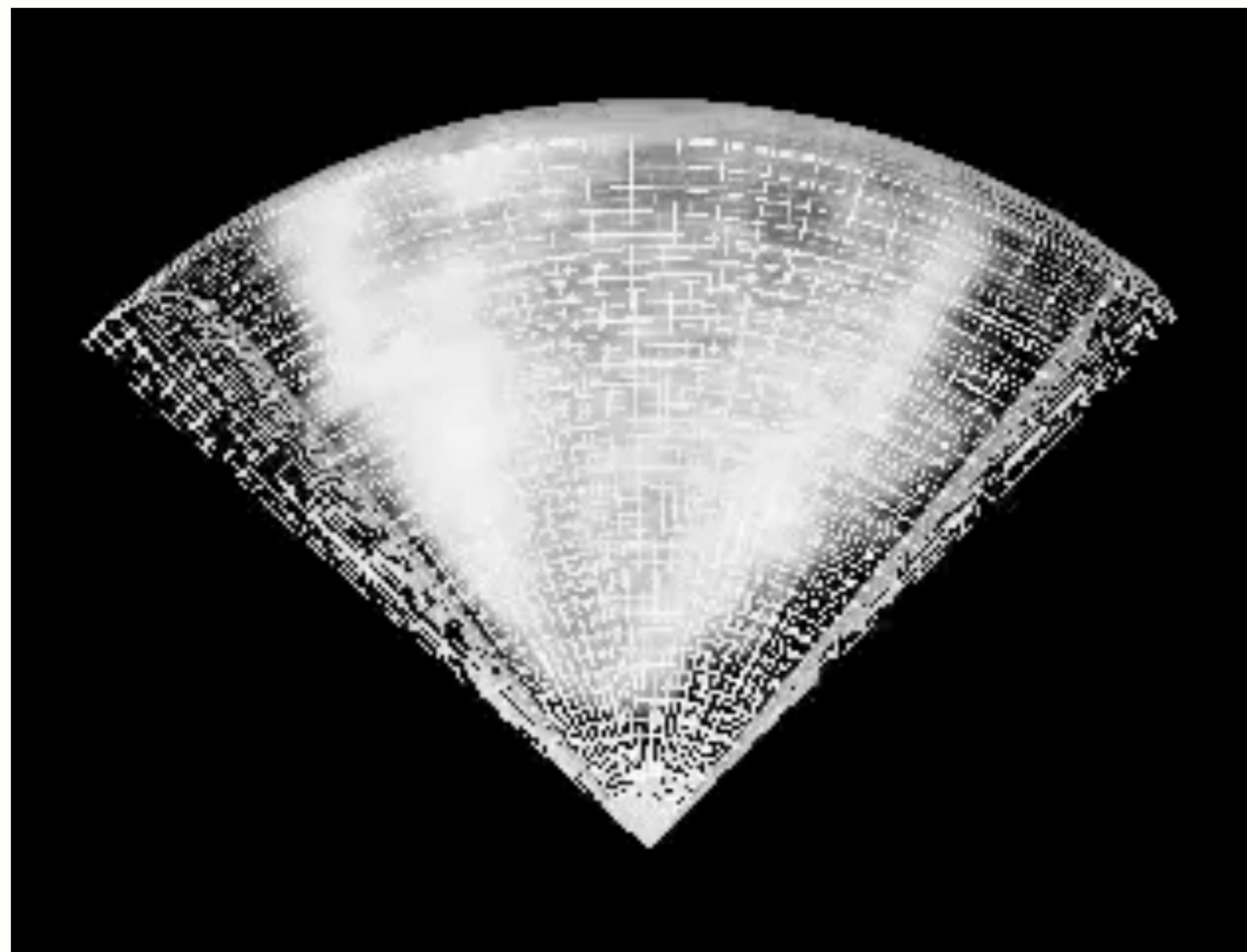


100.000 – 200.000 cases of **sudden cardiac deaths** in Germany per year



J. Schröder-Schötelig

Visualizing mechanical scroll waves within the heart muscle using highspeed ultrasound

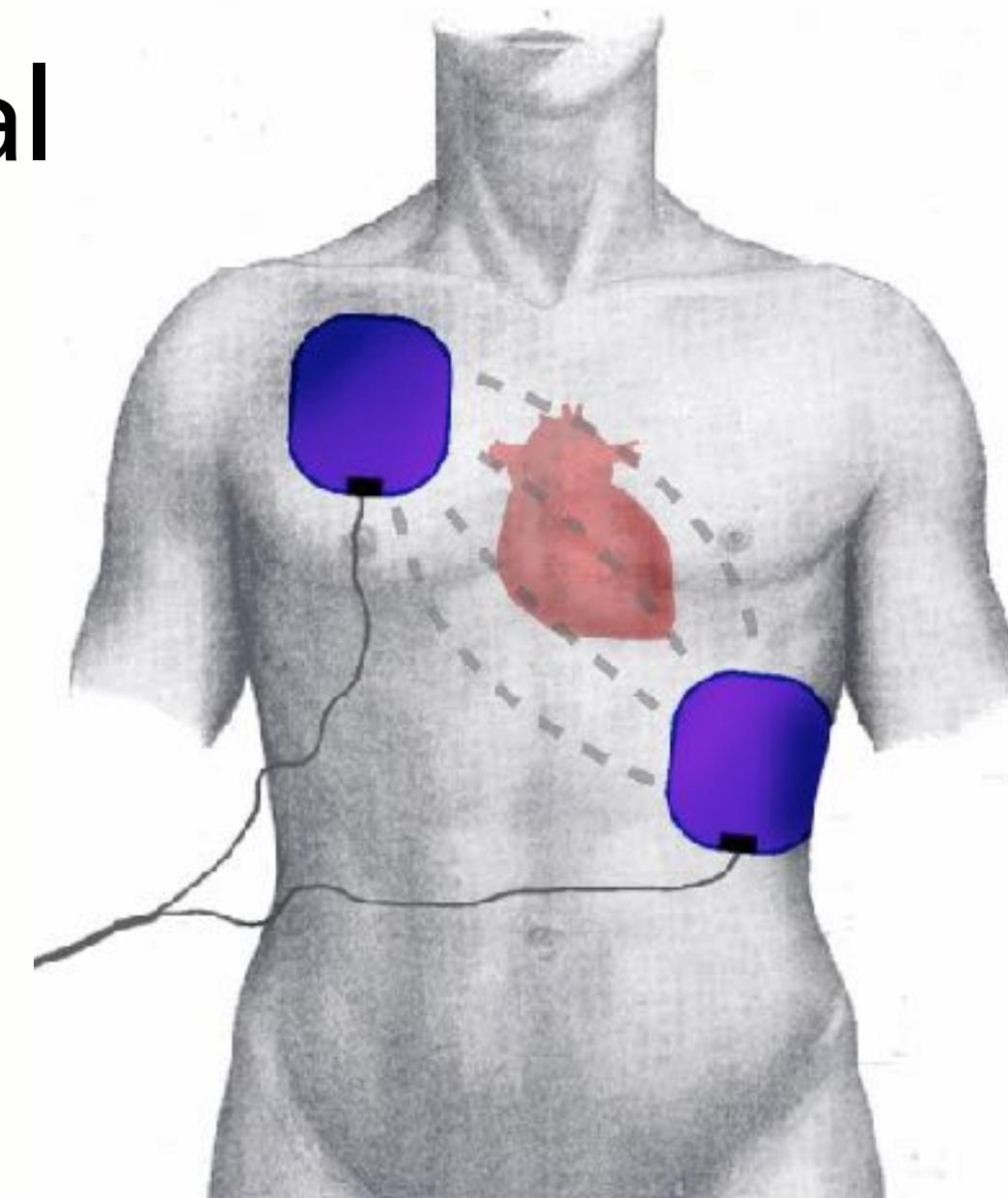


J. Christoph et al., Electromechanical vortex filaments during cardiac fibrillation, Nature 555, 667 (2018)

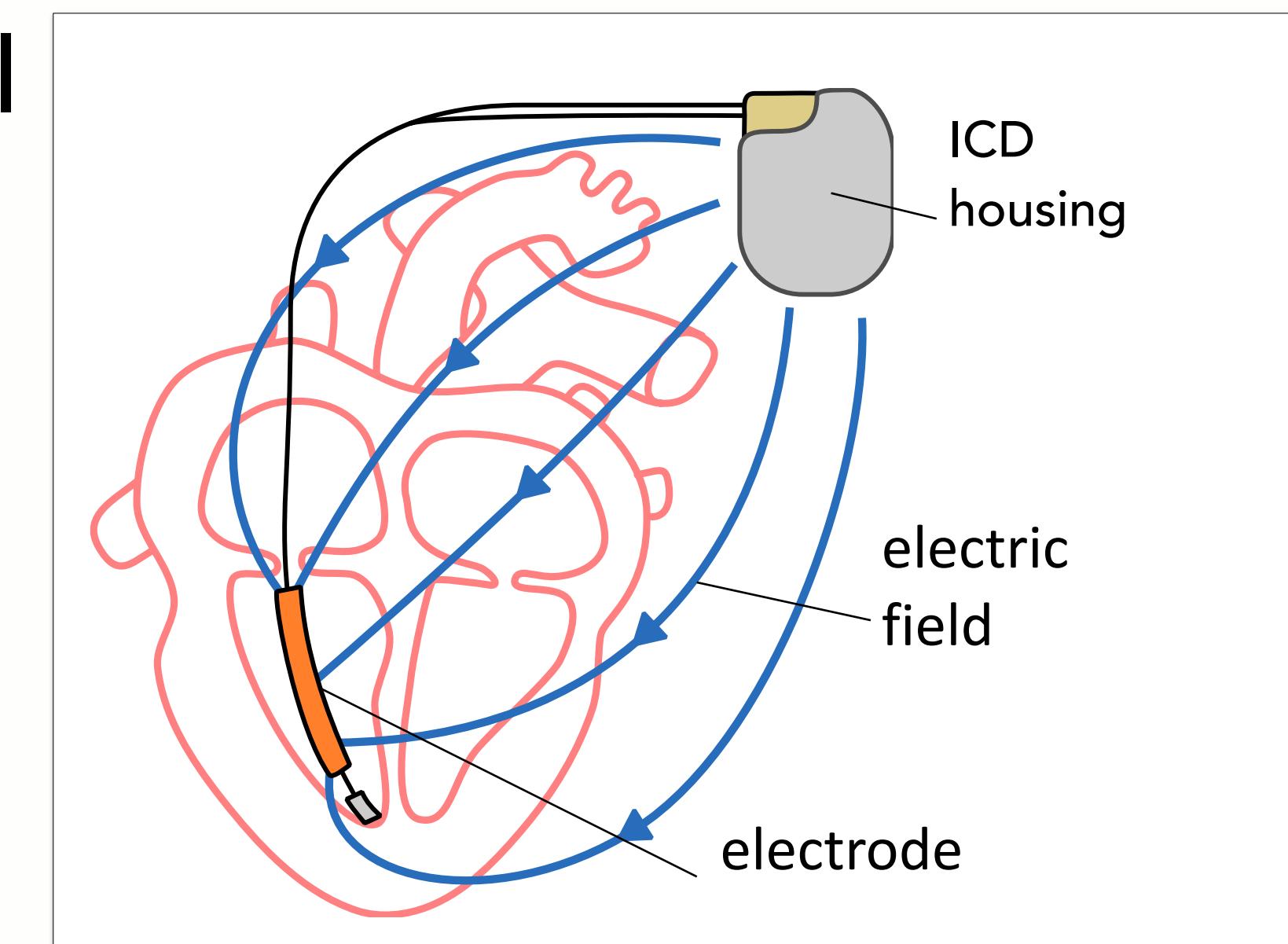
Defibrillation

Principle: Reset electrical activity of all cells by synchronous excitation

external



internal



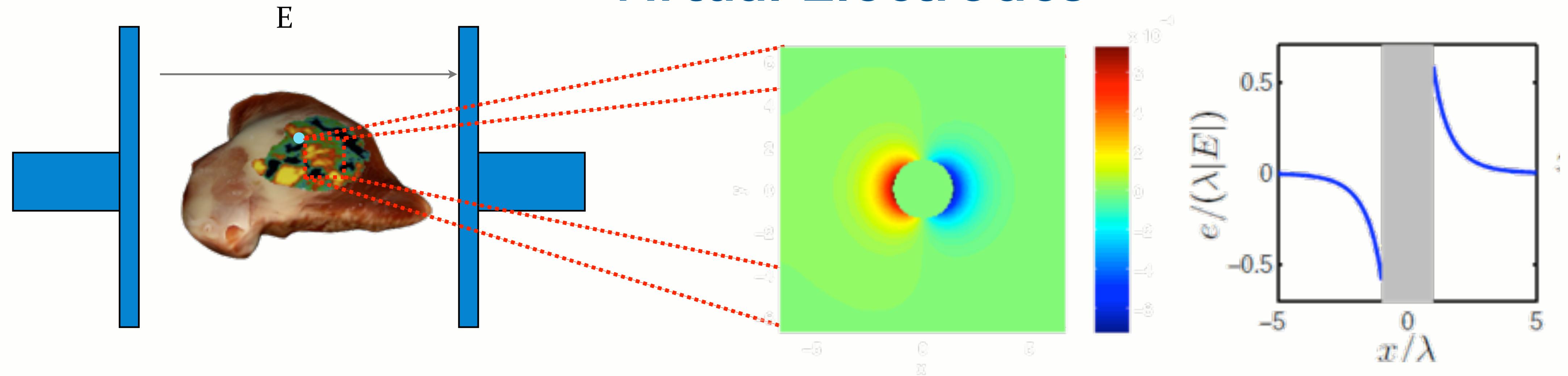
Electric shocks: energy 360J (external) 40 J (internal) 1000 V 30 A 12 ms

Severe side effects: tissue damage - traumatic pain

G.P. Walcott et al., Resuscitation 59, 59-70 (2003)

Terminating Cardiac Arrhythmias

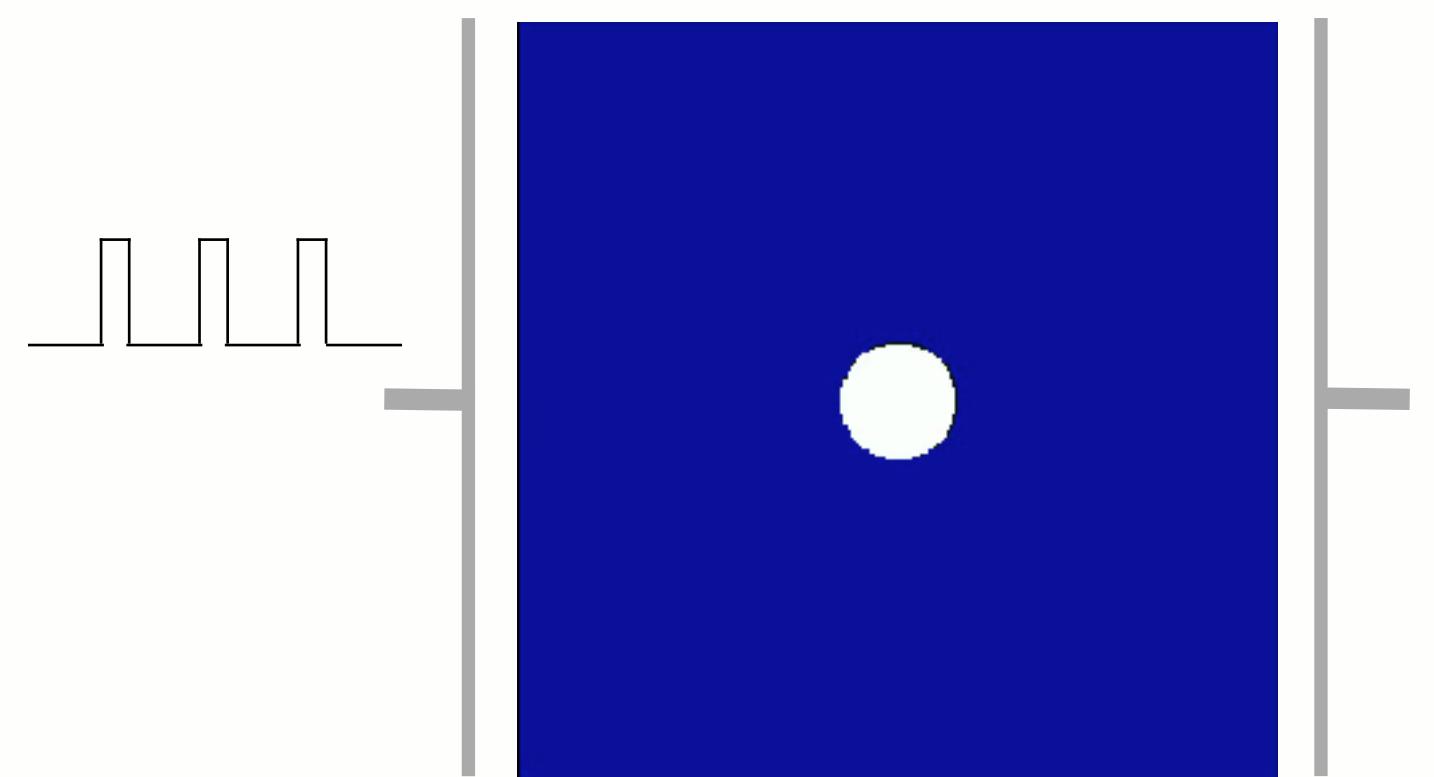
Virtual Electrodes



Blood vessels, scars, fatty tissue

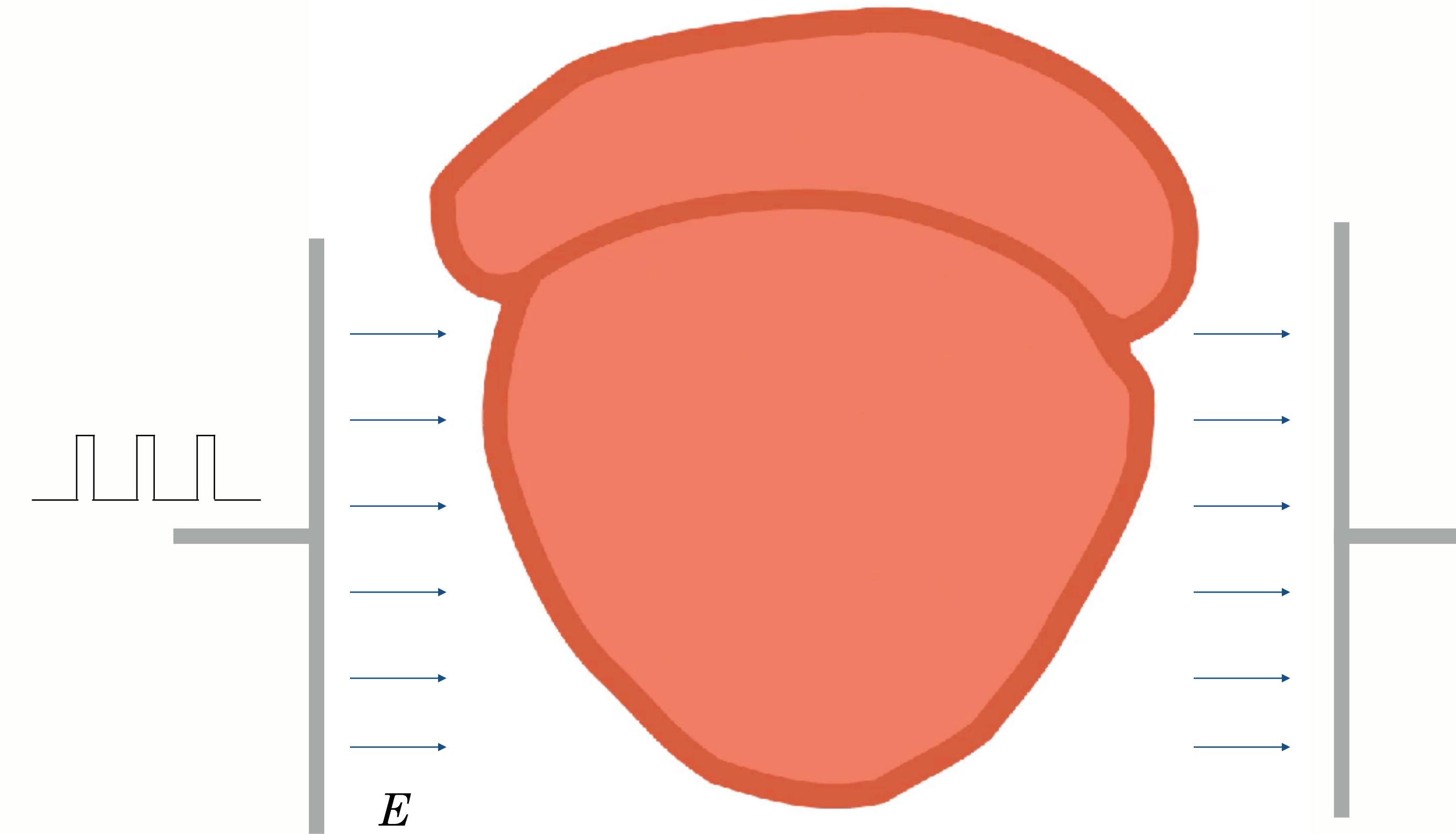
- are obstacles to electrical conduction
- may act as **virtual electrodes**

Super-threshold depolarization leads to **wave emission**
if a short **rectangular electric field pulse** is applied.



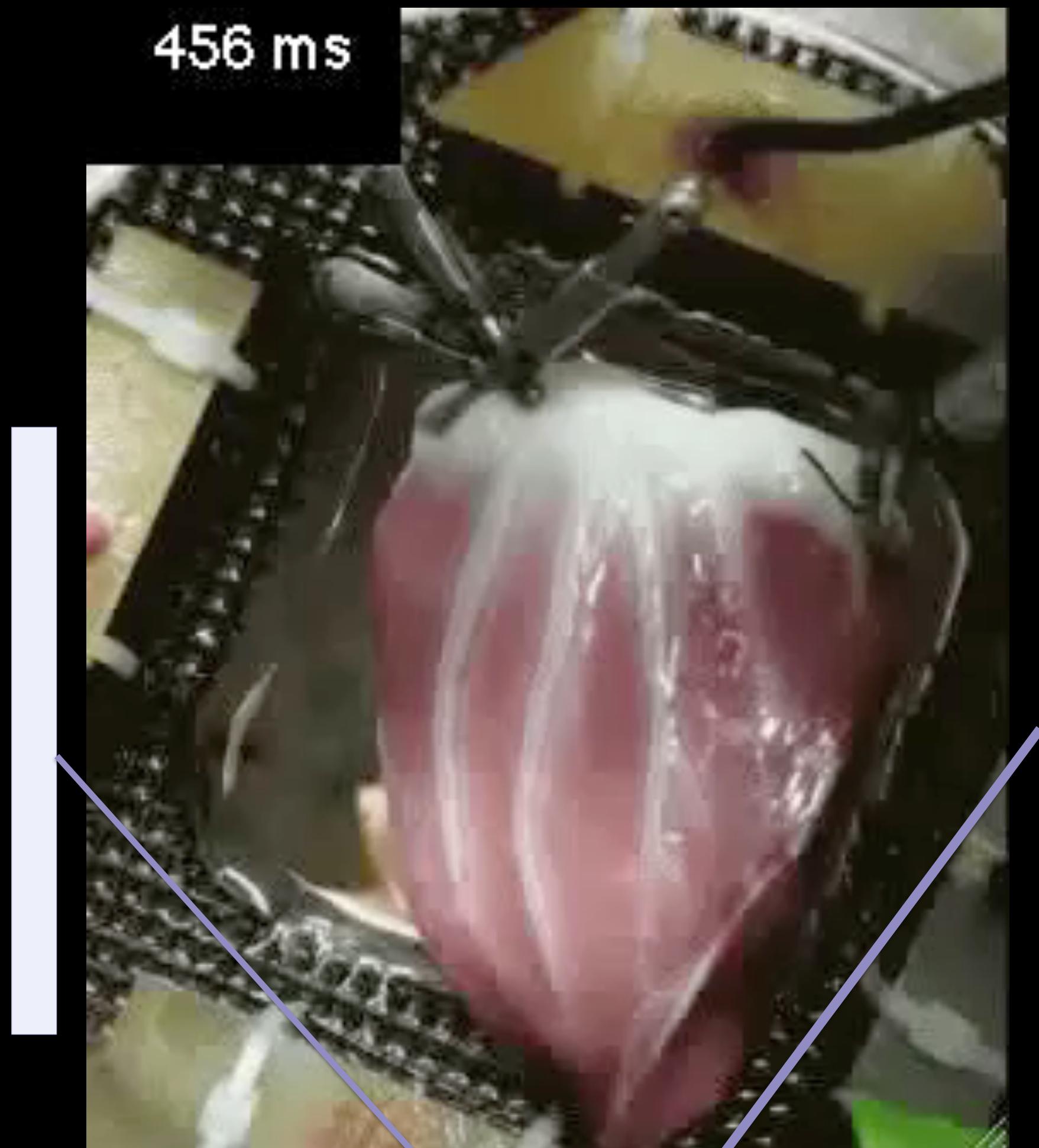
Terminating Cardiac Arrhythmias

Recruiting Networks of Virtual Electrodes for Terminating Cardiac Arrhythmias

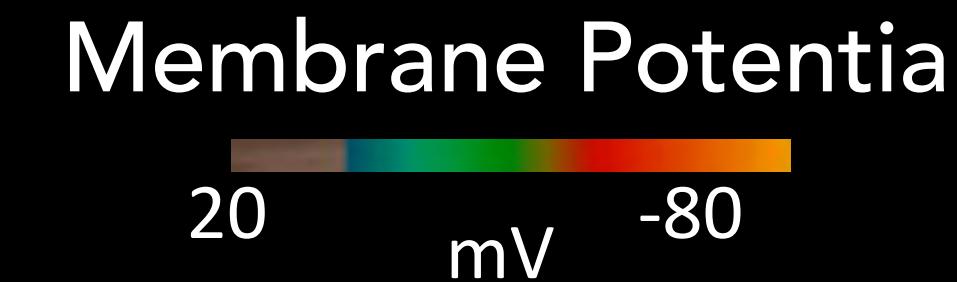


Animation: T. Lilienkamp

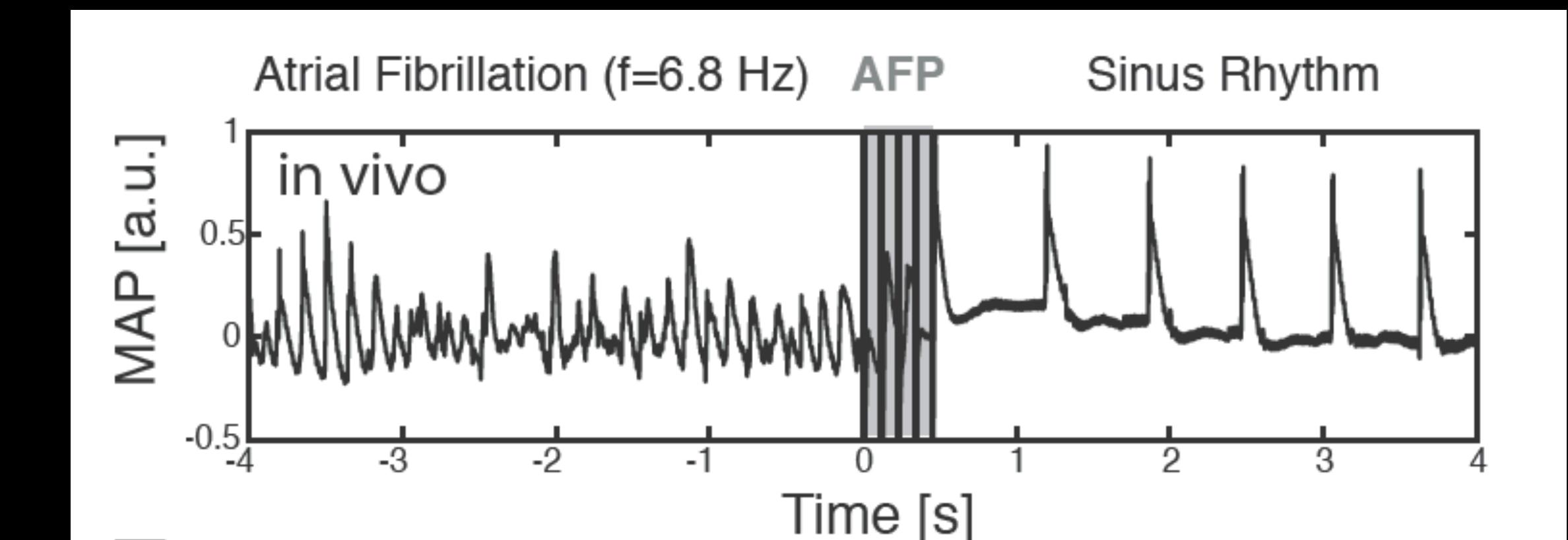
Low-Energy Anti-Fibrillation Pacing (LEAP)



Pulse Generator
Power Amplifier



N = 5 low energy pulses
 $E = 1.4 \text{ V/cm}$
 $dt = 90 \text{ ms}$

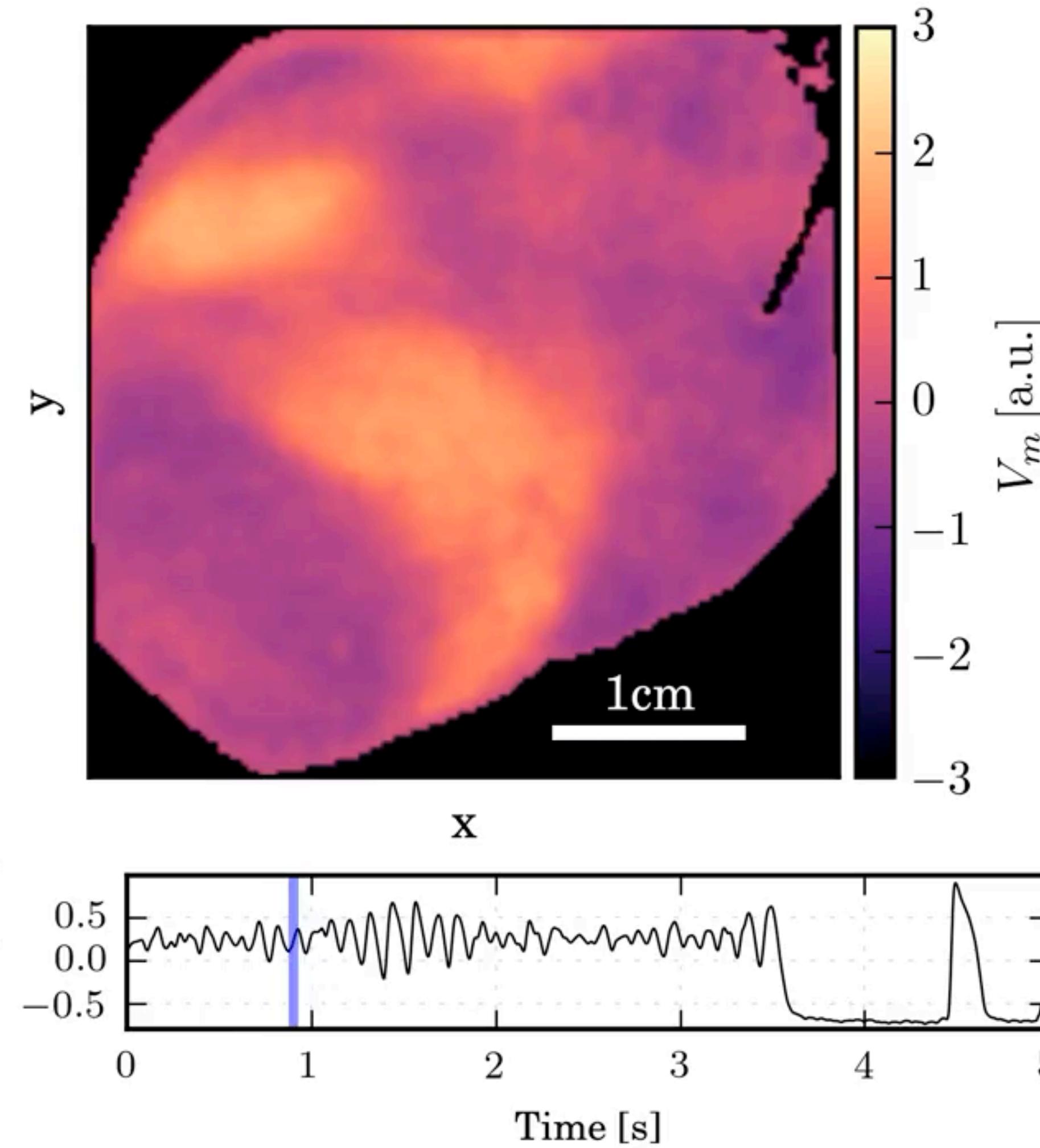


S. Luther et al., Nature 475, 235 (2011)

Transient Scroll Wave Dynamics during Ventricular Fibrillation

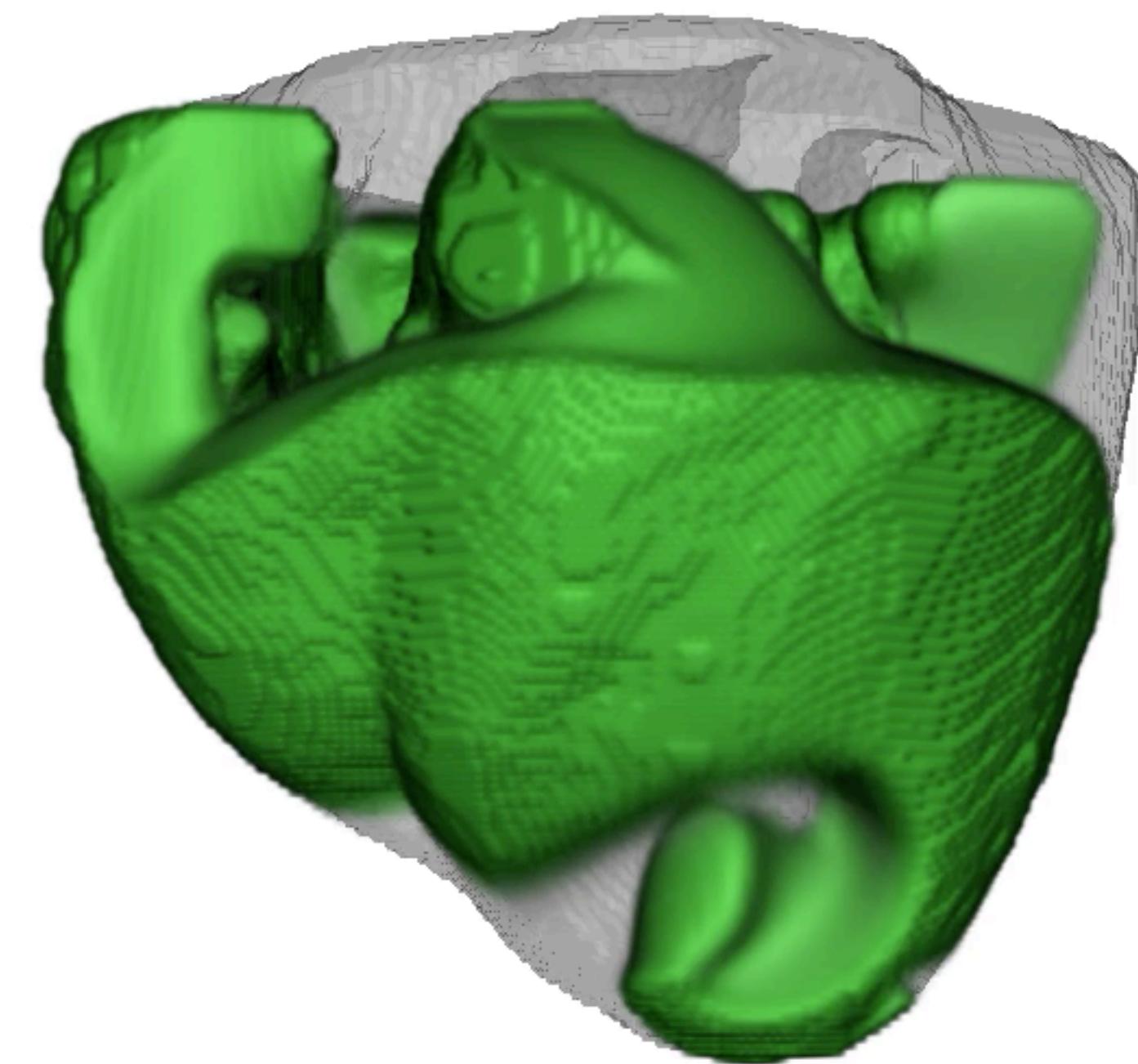
Experiment

Optical mapping
of a rabbit heart



Sebastian Berg
Daniel Hornung
Marion Kunze

Simulation
in a rabbit heart geometry



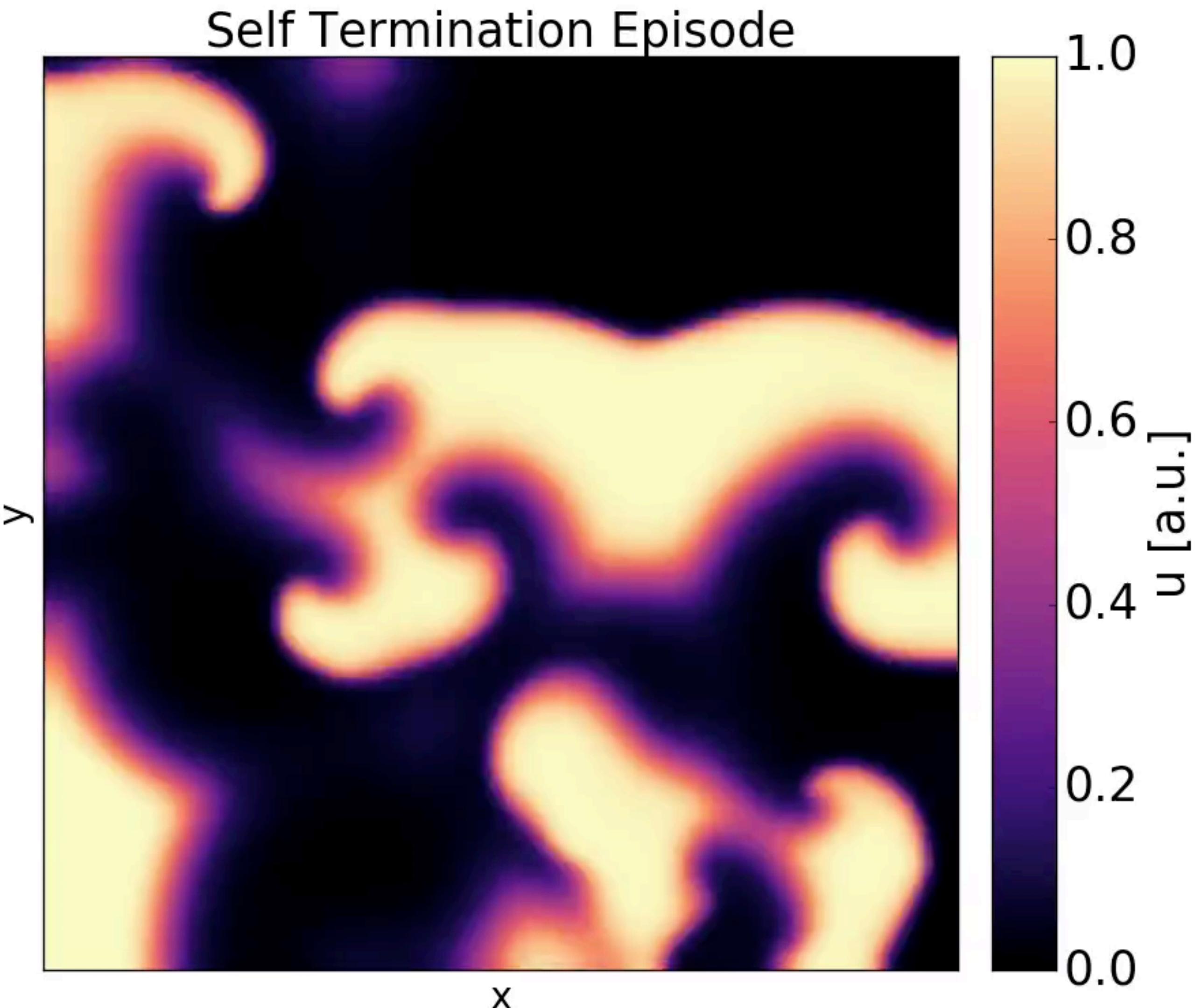
Thomas Lilienkamp

Transient Chaos

Simulation using the Fenton-Karma model

$$\begin{aligned}\frac{\partial u}{\partial t} &= \nabla \cdot \underline{\mathbf{D}} \nabla u - I_{Ion}(u, \mathbf{h})/C_m \\ \frac{\partial \mathbf{h}}{\partial t} &= \mathbf{g}(u, \mathbf{h})\end{aligned}$$

gating variables $\mathbf{h} = (v, w)$

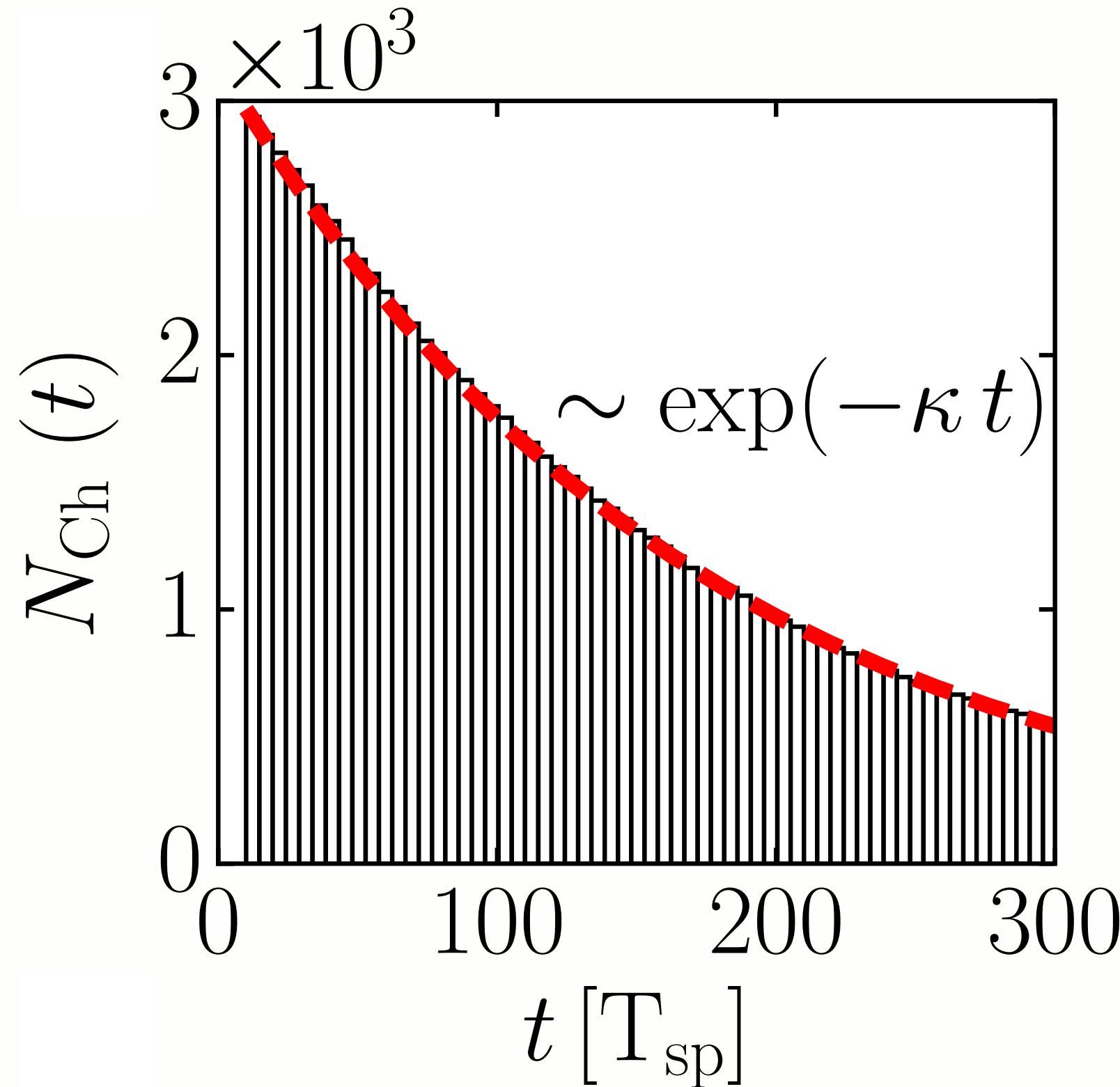


T. Lilienkamp and U. Parlitz, Phys. Rev. Lett. 120, 094101 (2018)

Chaotic transients and the average lifetime in 2D simulations

Fenton-Karma model 3000 initial conditions

fraction of trajectories still showing
chaotic dynamics at time t



$$N_{Ch}(t) \sim \exp(-\kappa t)$$

escape rate κ
quantifies **how fast trajectories**
from random initial conditions
escape the chaotic saddle and
reach the final (non-chaotic) state

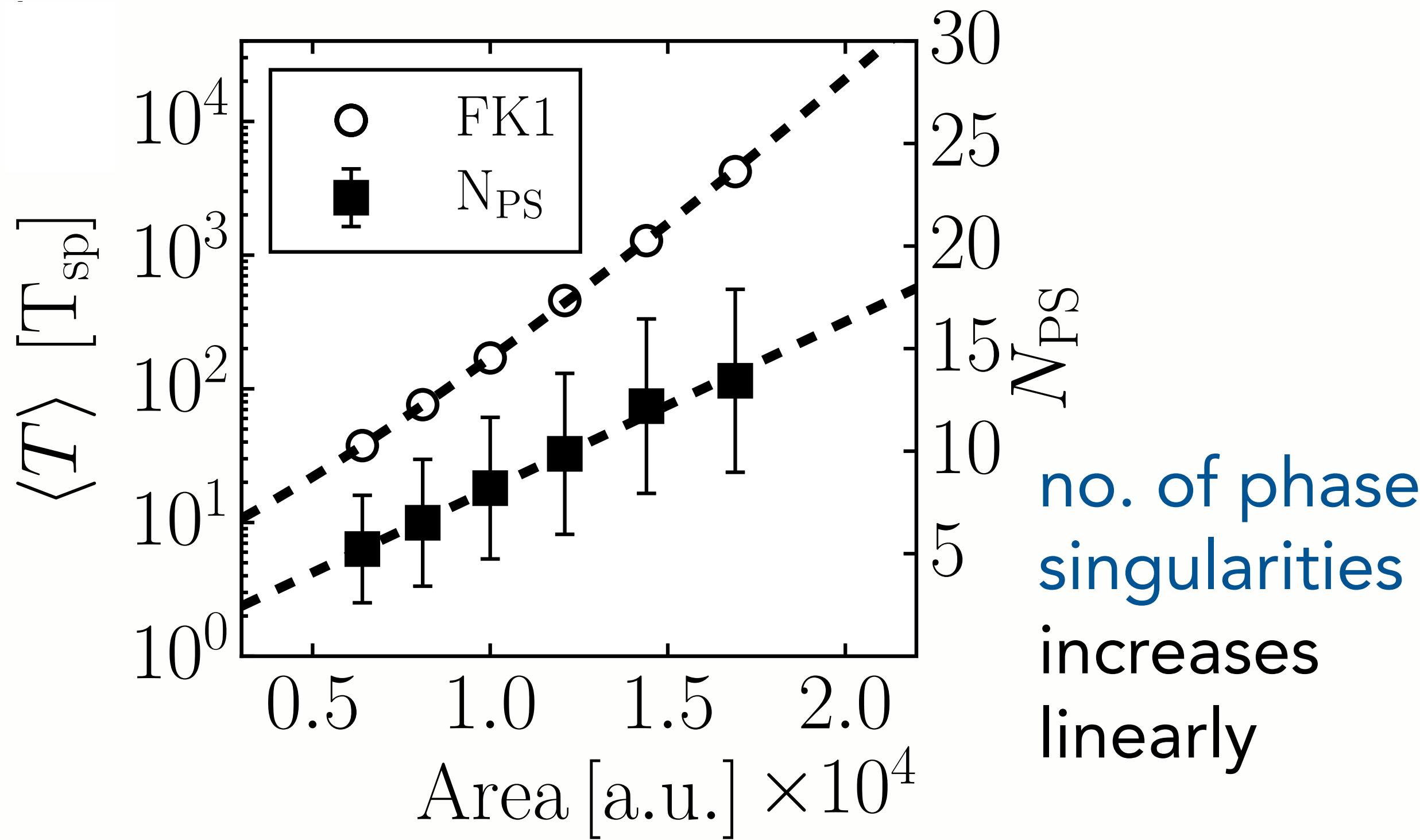
$$\langle T \rangle \approx \frac{1}{\kappa} \quad \text{average transient lifetime}$$

T. Lilienkamp et al., Phys. Rev. Lett. 119, 054101 (2017)

Chaotic transients and the average lifetime in 2D simulations

$\langle T \rangle \approx \frac{1}{\kappa}$ **average transient lifetime**

increases exponentially with system size



no. of phase singularities increases linearly

- Larger heart muscle volumes increase the risk of cardiac arrhythmias and related morbidity and mortality.
→ due to longer transients and more phase singularities (??)

- Impact of (finite) perturbations changes during some period of time prior to the end of the transient.

→ precursors for end of arrhythmia (??)

T. Lilienkamp and U. Parlitz,
Phys. Rev. Lett. 120, 094101 (2018),
Phys. Rev. E 98, 022215 (2018)

Summary

The heart

- consists of a **network of electrically and mechanically coupled excitable elements**
- forming an **excitable medium** that supports **plane waves, spiral waves, and**
- **(life-threatening) spatio-temporal chaos** (e.g., ventricular fibrillation)
- that can be transient and exhibits **complexity fluctuations** and
- provides an ambitious target for **(low-energy) control methods (defibrillation)**

Outlook: Interaction with other organs, in particular **heart & brain**

→ **Network Physiology**

Acknowledgement

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at the Max Planck Institute for Dynamics and Self-Organization, Göttingen



DZHK
DEUTSCHES ZENTRUM FÜR
HERZ-KREISLAUF-FORSCHUNG E.V.



Federal Ministry
of Education
and Research



Thank you!