



Towards Critical SNNs: Astrocytes Detect Chaos in Neuronal Dynamics

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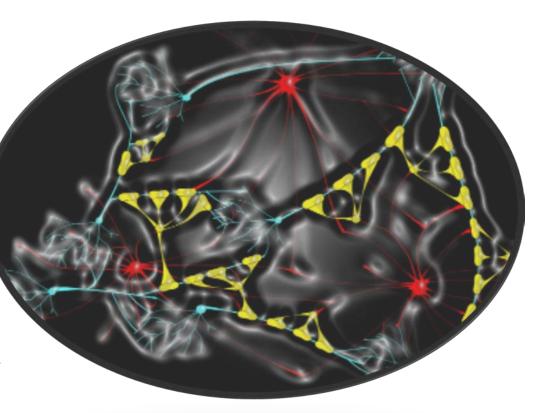
Main Message

• Brain cells have been computing long before we started recording neurons

 Neuronal processing is mediated by non-neuronal cells – and networks

• Non-neuronal computation: a critical untapped frontier to understand brain function

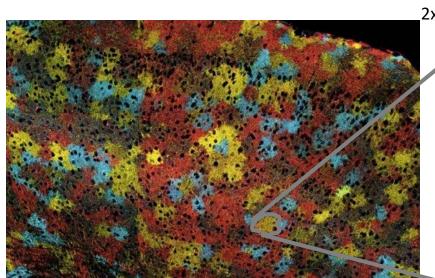
 Non-neuronal computing: rich opportunities for versatile and robust AI





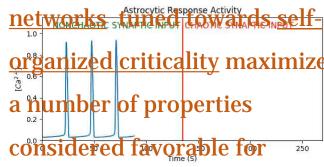
Astrocytes: Non-Neuronal Cells

Detecting Neuronal Chaos





Astrocytes oversee the neuronal activity with *non*-overlapping receptive fields



computation

organized criticality maximize

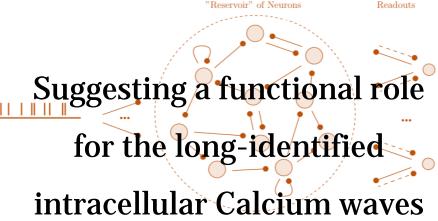
irontiars in SYSTEMS NEUROSCIENCE



Spike avalanches in vivo suggest a driven, slightly subcritical brain state

Viola Priesemann 1,2,3,4 *, Michael Wibral 5,6, Mario Valderrama 7, Robert Pröpper 8,9 Michel Le Van Quyen¹⁰, Theo Geisel^{1,2}, Jochen Triesch³, Danko Nikolić^{3,4,6,11} and Matthias H. J. Munk¹²

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- Department of Biomedical Engineering, University of Los Andes, Bogotá, Colombia
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- ⁹ Bernstein Center for Computational Neuroscience, Berlin, Germany
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Brodeur & Rouat ICANN 2012

Maass, Natschlager, Markram Neural Computation 2002



Real-time SNN-control of Robots



Real-world Environment

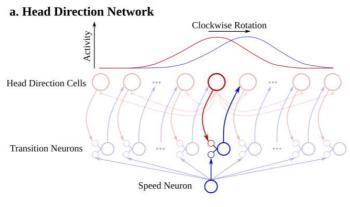


Experiment 1: os to 6os

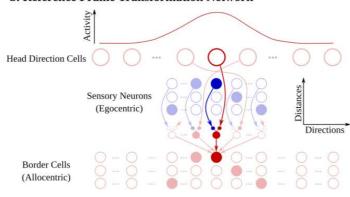
Speed x10

[NICE '17 & '18, CCN '17, ICONS '18]

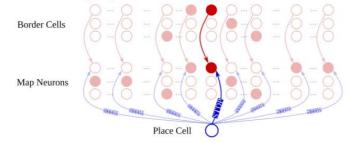
[Tang, Shah, Michmizos arXiv:1903.02504]



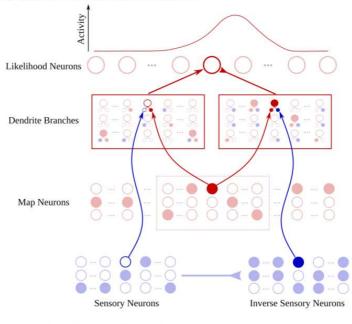
b. Reference Frame Transformation Network



c. Distance Mapping Network

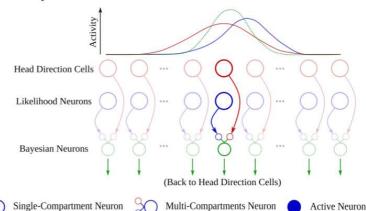


d. Observation Likelihood Network



e. Bayesian Inference Network

Excitatory Synapse



— Inhibitory Synapse

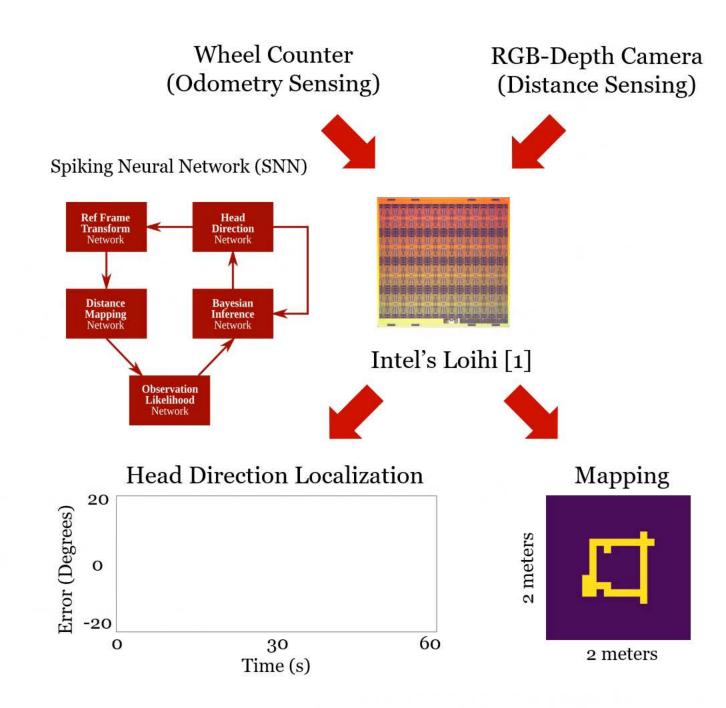


Real-world Environment



Experiment 1: os to 6os

Speed x10





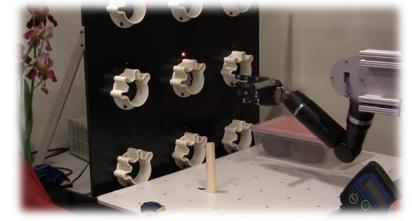
Record



1932. Edgar Adrian, Nobel Prize Single-Neuron Recordings



1971. David Cohen, MIT Magnetoencephalography



2013. Motor neurons control a robotic arm for paraplegic patients (BrainGate)

Information = f (electrical activity)

neurons



1997. Deep Brain Stimulation for alleviating Parkinson's disease

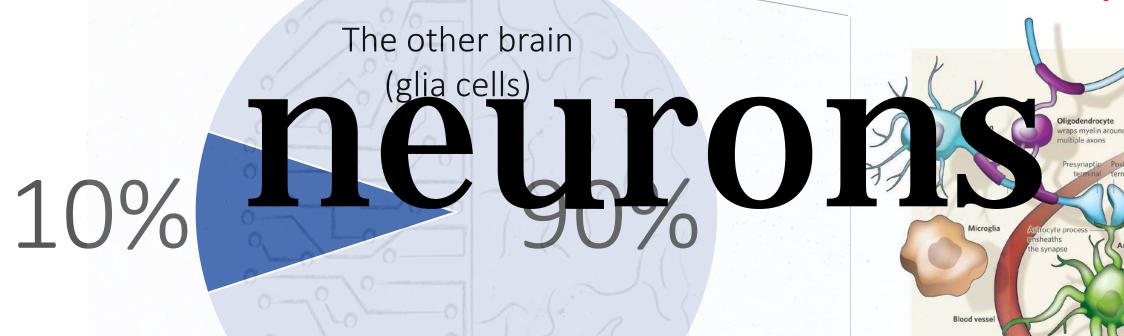


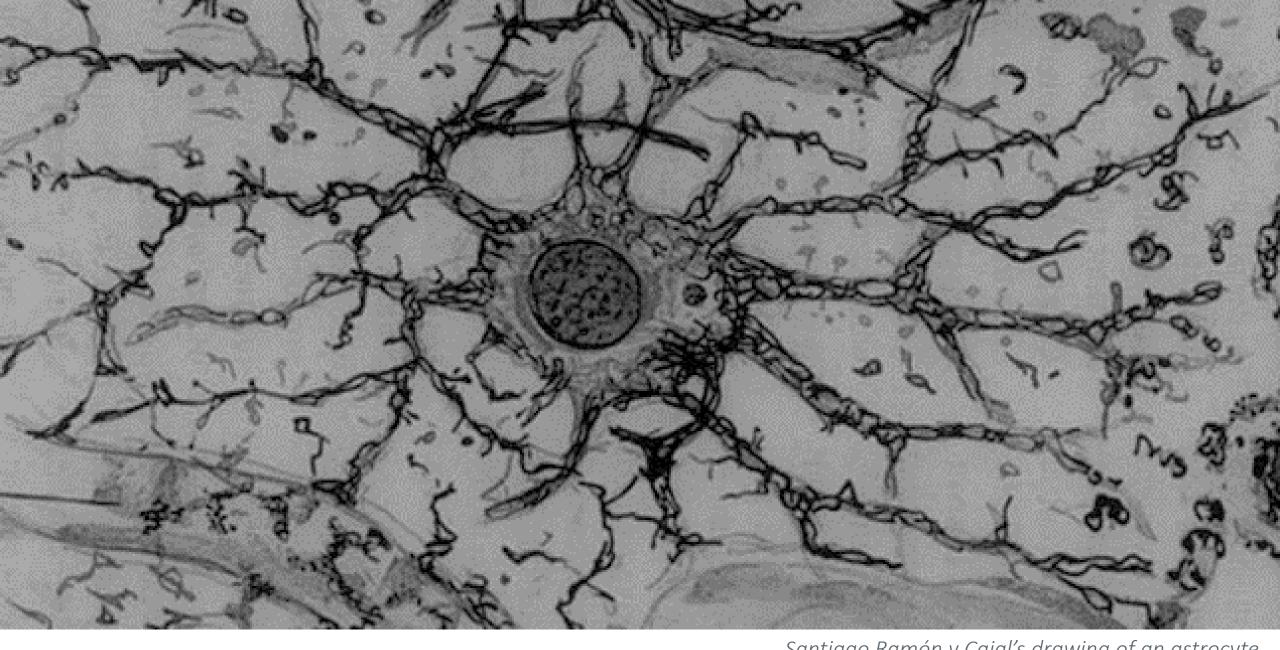
2013. TMS applied to the motor cortex induces hand movement

manipulate



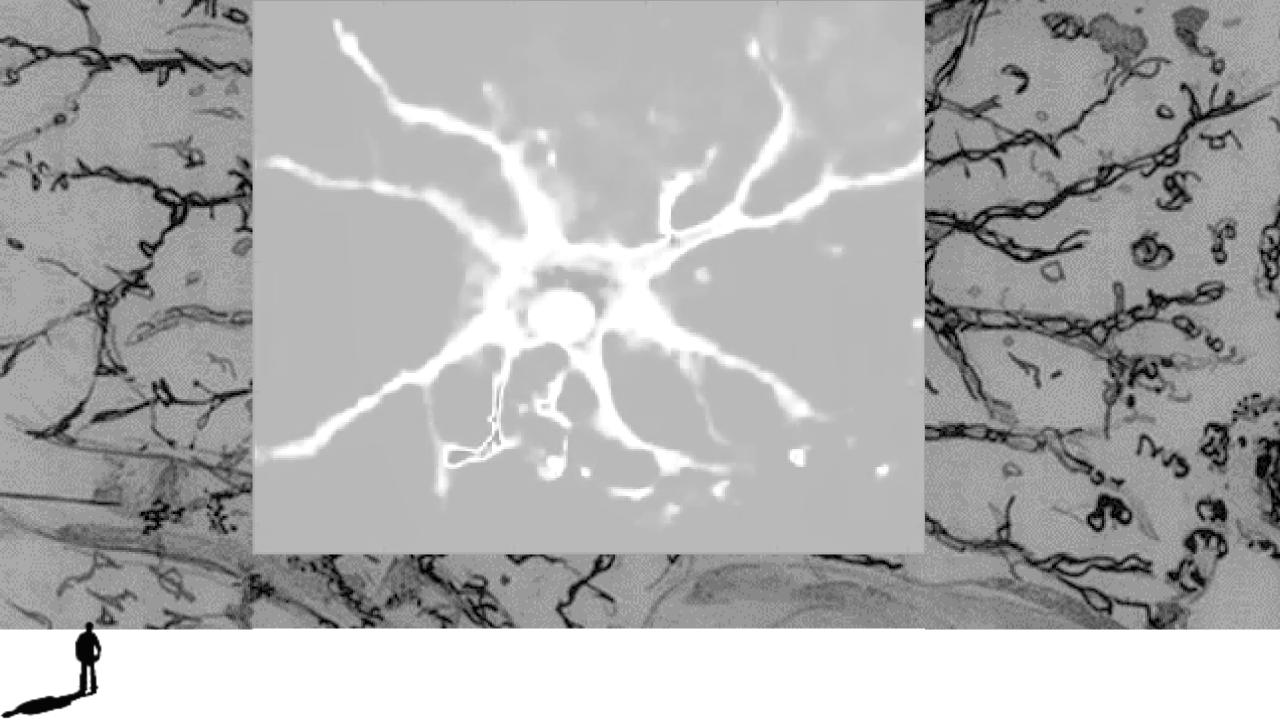
Non-neuronal brain cells are electrically silent

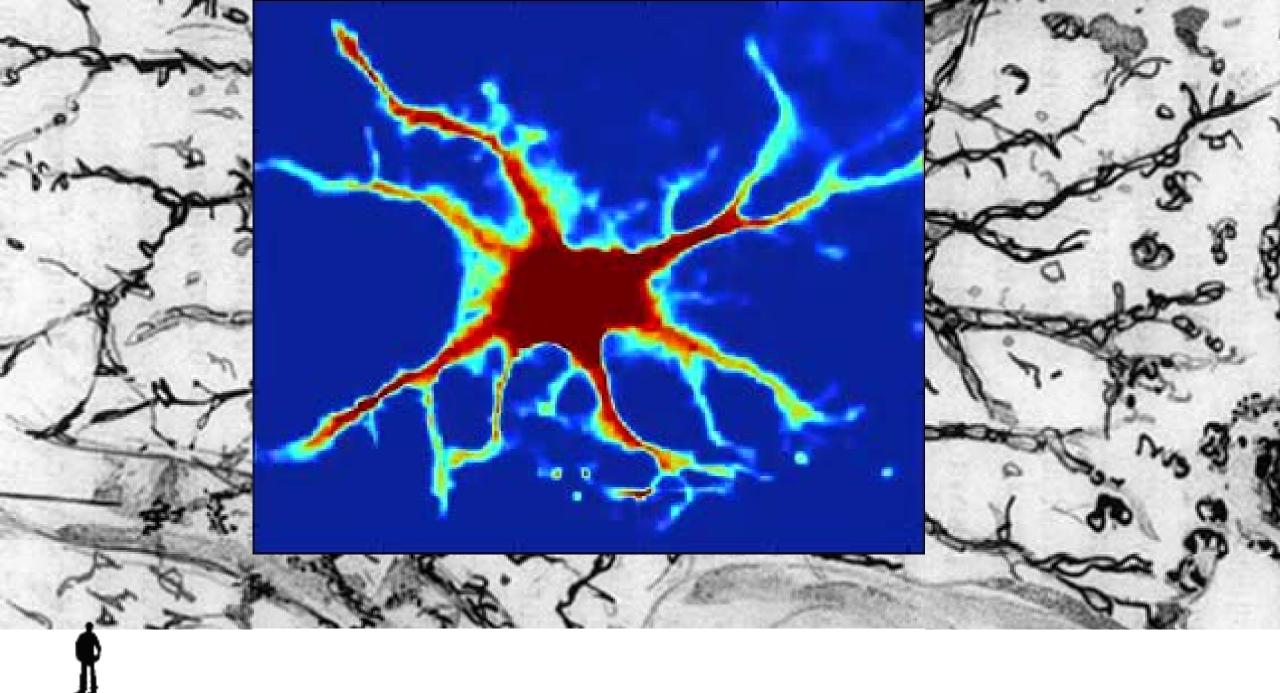




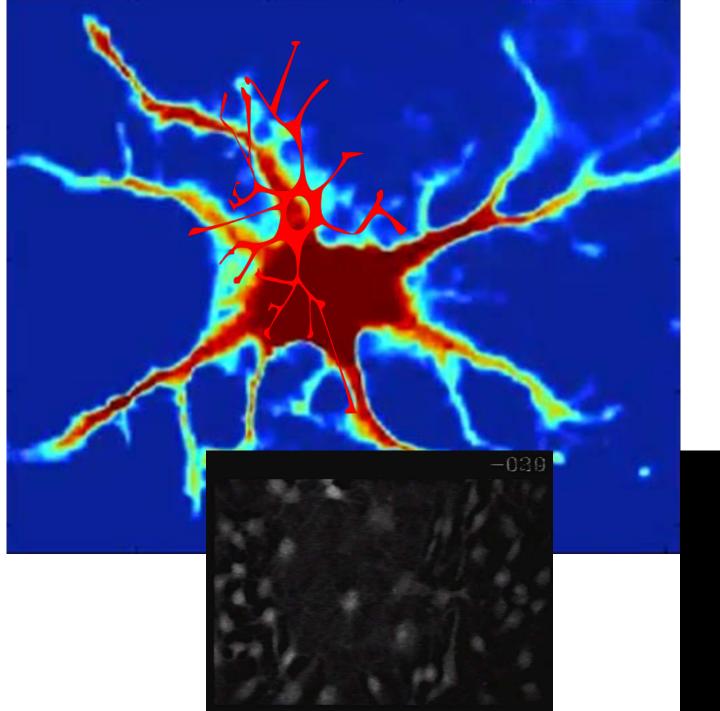
Santiago Ramón y Cajal's drawing of an astrocyte

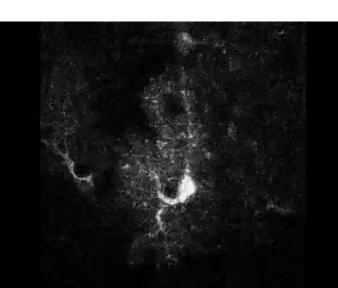
Ramón y Cajal S. Something about the physiological significance of neuroglia. Revista Trimestral Micrografía 1, 3–47, 1897





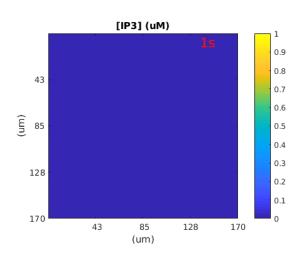


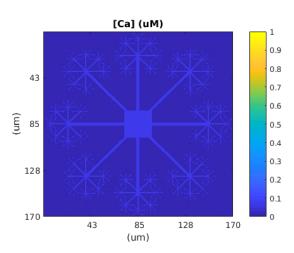


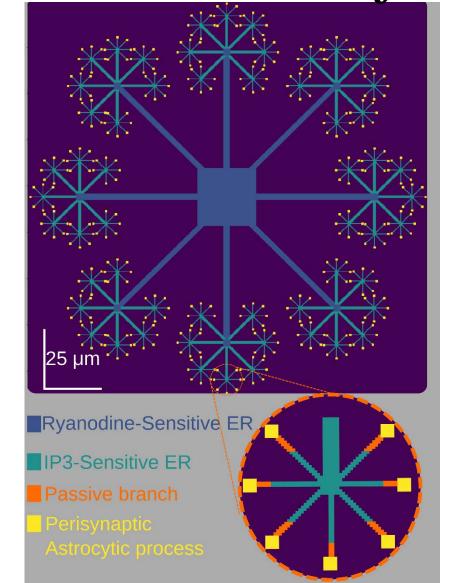


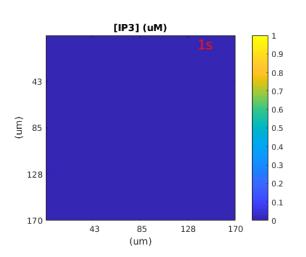


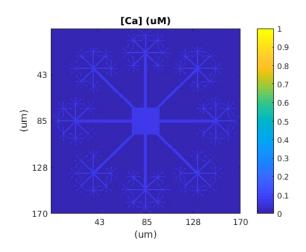
Computational Astrocyence





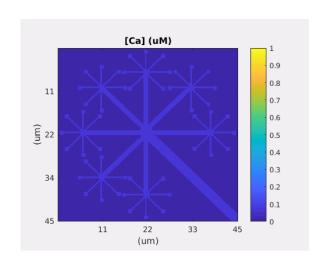


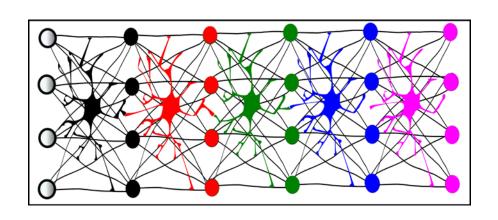


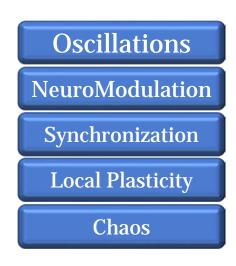




Computational Astrocyence





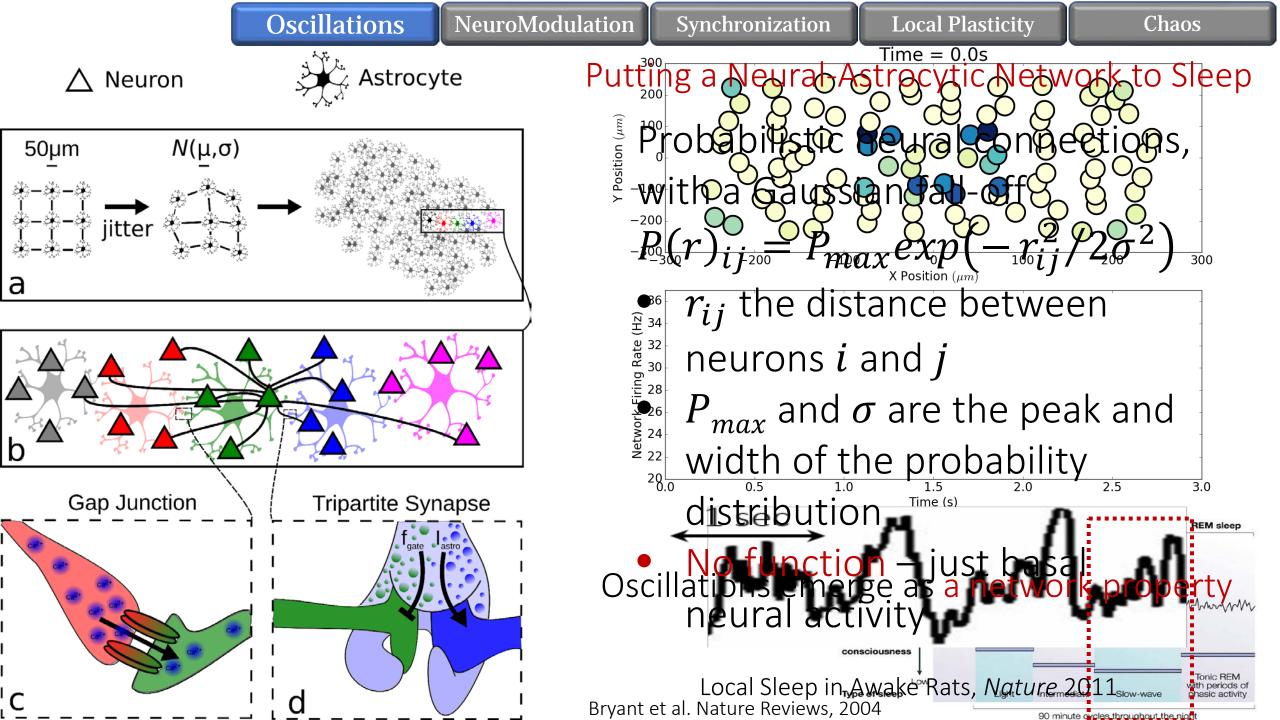


Astrocytic Cell

Neuronal-Astrocytic Network

Network Behavior





SCIENTIFIC REPORTS

OPEN

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Extensive astrocyte synchronization advances neuronal coupling in slow wave activity *in vivo*

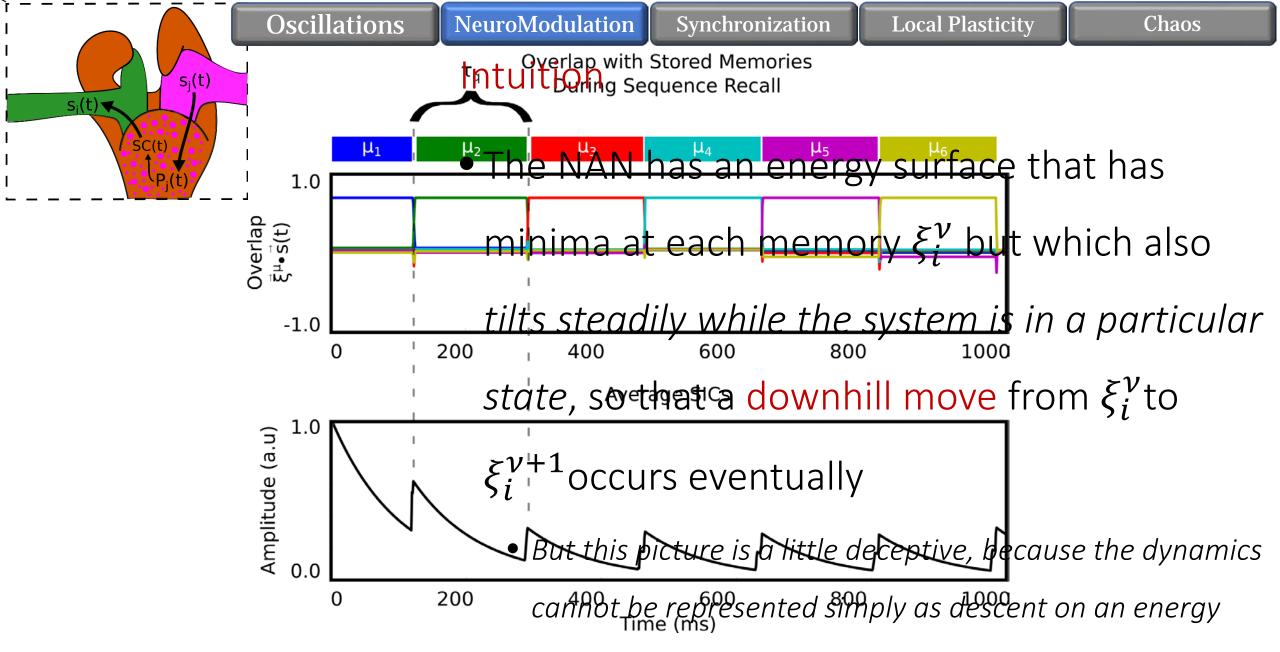
Zsolt Szabó 601, László Héja 601, Gergely Szalay2, Orsolya Kékesi1, András Füredi3,6, Kornélia Szebényi3,6, Árpád Dobolyi4, Tamás I. Orbán3, Orsolya Kolacsek3, Tamás Tompa2, Zsombor Miskolczy5, László Biczók 605, Balázs Rózsa2, Balázs Sarkadi3 & Julianna Kardos 601

Slow wave activity (SWA) is a characteristic brain oscillation in sleep and quiet wakefulness. Although the cell types contributing to SWA genesis are not yet identified, the principal role of neurons in the emergence of this essential cognitive mechanism has not been questioned. To address the possibility of astrocytic involvement in SWA, we used a transgenic rat line expressing a calcium sensitive fluorescent protein in both astrocytes and interneurons and simultaneously imaged astrocytic and neuronal activity in vivo. Here we demonstrate, for the first time, that the astrocyte network display synchronized recurrent activity in vivo coupled to UP states measured by field recording and neuronal calcium imaging. Furthermore, we present evidence that extensive synchronization of the astrocytic network precedes the spatial build-up of neuronal synchronization. The earlier extensive recruitment of astrocytes in the synchronized activity is reinforced by the observation that neurons surrounded by active astrocytes are more likely to join SWA, suggesting causality. Further supporting this notion, we demonstrate that blockade of astrocytic gap junctional communication or inhibition of astrocytic Ca²⁺ transients reduces the ratio of both astrocytes and neurons involved in SWA. These in vivo findings conclusively suggest a causal role of the astrocytic syncytium in SWA generation.

Increasing body of evidence substantiating the impact of astrocytes on neuronal activity prompted a paradigm shift from the neurocentric philosophy of nervous system function. Accordingly, astrocytes are increasingly recognized as major players in the modulation of neuronal function under both physiological and pathophysiological conditions $^{4-7}$. Beyond the local astroglial control over synaptic activity $^{8-12}$, however, little is known about the role of astrocytic networks in modulating large-scale neuronal ensembles. Exploration of the role of large-scale strocytic networks in information processing and cognition still lags behind its neuronal counterpart 15,16 . We conceived that fundamental properties of networking astrocytes may underlie physiological network-network interaction between astrocytes and neurons. Astrocytes are capable of 1) detecting neuronal activity, 2) responding to this activity by raising local Ca $^{2+}$ transients, 3) propagating the local changes over extended spatial scales by Ca $^{2+}$ waves traveling through the directly and densely interconnected astrocytic syncytium and 4) modulating neuronal activity at multiple locations by releasing gliotransmitters and other neuromodulatory substances or regulating ionic homeostasis 15 . Thus, astrocytes are ideally positioned to induce or contribute to synchronization of large-scale neuronal networks. Along this line, we have previously demonstrated that the astrocytic and

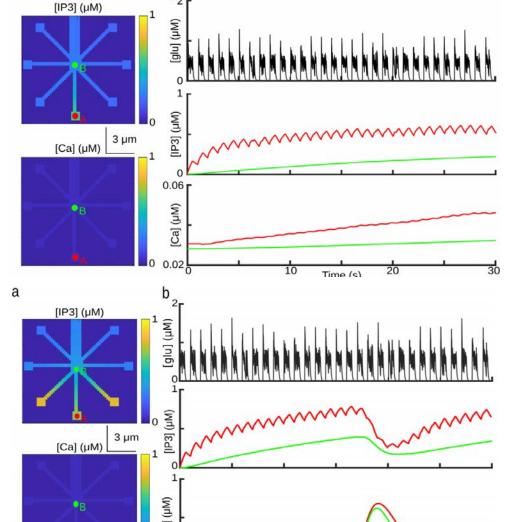
¹Institute of Organic Chemistry, Research Centre for Natural Sciences, Hungarian Academy of Sciences, Magyar tudósok körúţia 2, 1117, Budapest, Hungary. ²Institute of Experimental Medicine, Hungarian Academy of Sciences, Szigony 43, 1083, Budapest, Hungary. ³Institute of Enzymology, Research Centre for Natural Sciences, Hungarian Academy of Sciences, Magyar tudósok körúţia 2, 1117, Budapest, Hungary. *MTA-ELTE Laboratory of Molecular and Systems Neurobiology, Department of Physiology and Neurobiology, Eötvös Loránd University, Pázmány Péter sétány 1C, 1117, Budapest, Hungary. *Institute of Materials and Environmental Chemistry, Research Centre for Natural Sciences, Hungarian Academy of Sciences, Magyar tudósok körúţia 2, 1117, Budapest, Hungary. *Institute of Cancer Research, Medical University Wien, Borschkegasse 8a, 1090, Wien, Austria. Zsolt Szabó and László Héja contributed equally to this work. Correspondence and requests for materials should be addressed to L.H. (email: heja.laszlo@ttk.mta.hu)

hypothesis was reinforced modelling study showing that intercellular Ca2+ signaling potentially can introduce slow oscillation neurons [our Ref]. Our experimental this hypothesis supports by strongly demonstrating that increasing astrocytic influence neurons indeed drives them to join oscillatory activity (Fig. 5C). In this context it is also important to note, that the ratio of astrocytes involved in the SWA was found to start decreasing after virtually all neurons (Figs activity simultaneous observation further supports the astrocytic activity corresponds to the generation or maintenance, rather than termination of SWA."



The overlap of the neural network state was allowed by the state of the overlap of the neural network state of the overlap of the overlap of the neural network state of the overlap of the neural network state of the overlap of the overlap of the overlap of the neural network state of the overlap of

Astrocytes **sense** and **impose** synchronized neuronal activity



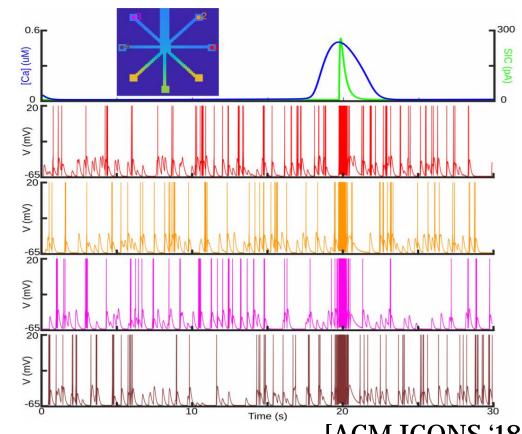
Time (s)

KEY POINT #1

Astrocytes encode synchronous synaptic activity into the extent of their Ca2+ waves

KEY POINT #2

They can impose synchronization to a group of postsynaptic neurons



[ACM ICONS '18] [ACM BI '18]

[IEEE BHI '19]

Oscillations

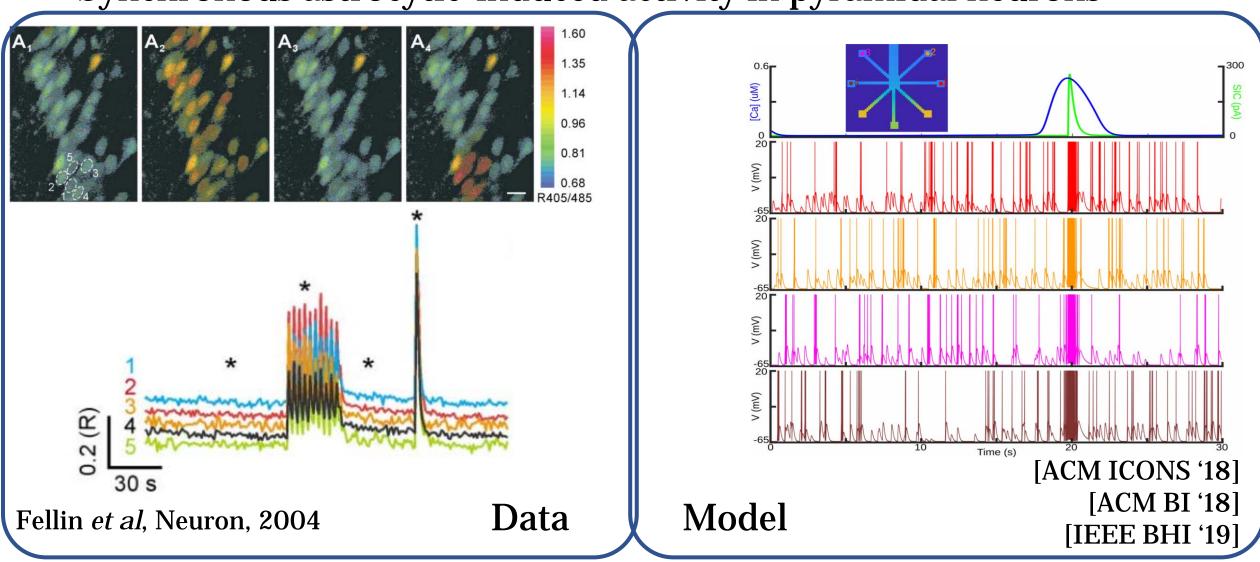
NeuroModulation

Synchronization

Local Plasticity

Experimentally-verified Computational Results

Synchronous astrocytic-induced activity in pyramidal neurons



Oscillations NeuroModulation

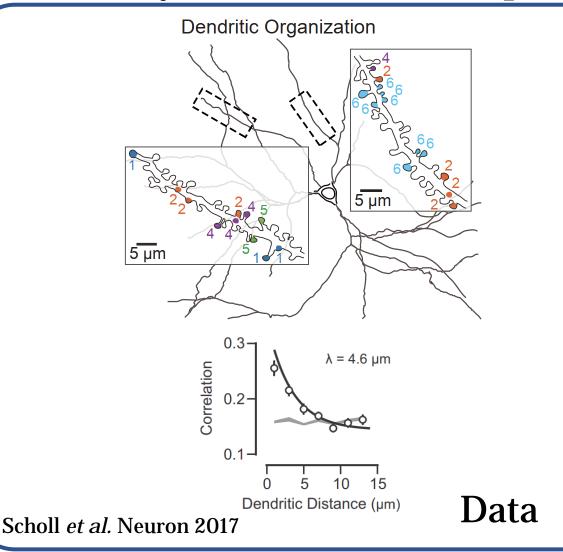
Synchronization

Local Plasticity

Chaos

Computationally-explained Experimental Results

Astrocytic-induced local plasticity

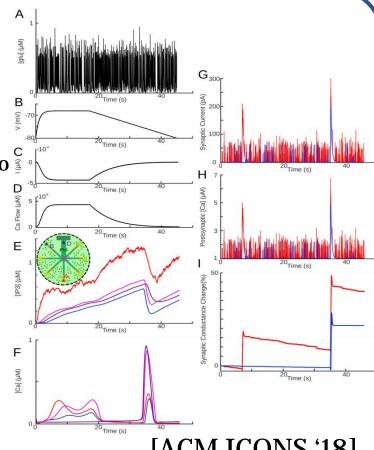


KEY POINT #3

- Two Ca²⁺ waves in the micro-domain: a)
 VGCCs and b)Glu-IP₃
- Process-specific VGCC

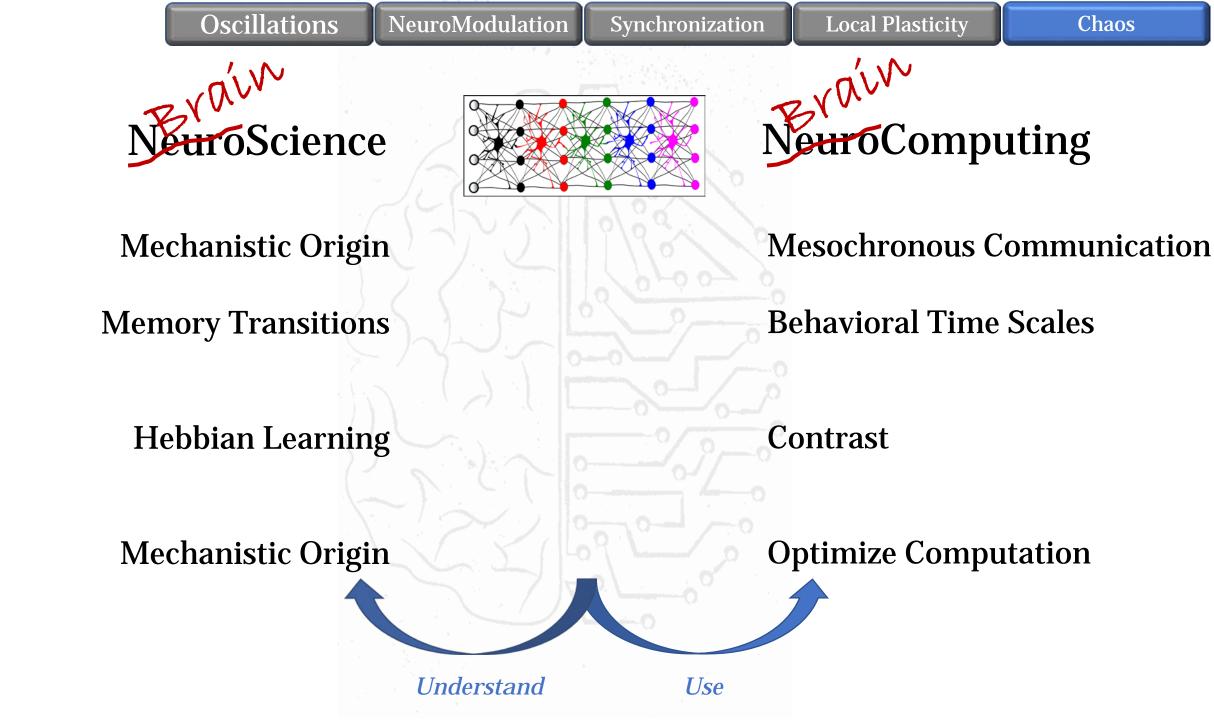
 Ca²⁺ waves restricted to processes with active presynaptic sites
- Microdomain-wide Ca²⁺ waves induced heteroshynaptic plasticity in inactive synapses
- Possible astrocytic mechanism for local clustering of dendritic spines within 15 µm

Model

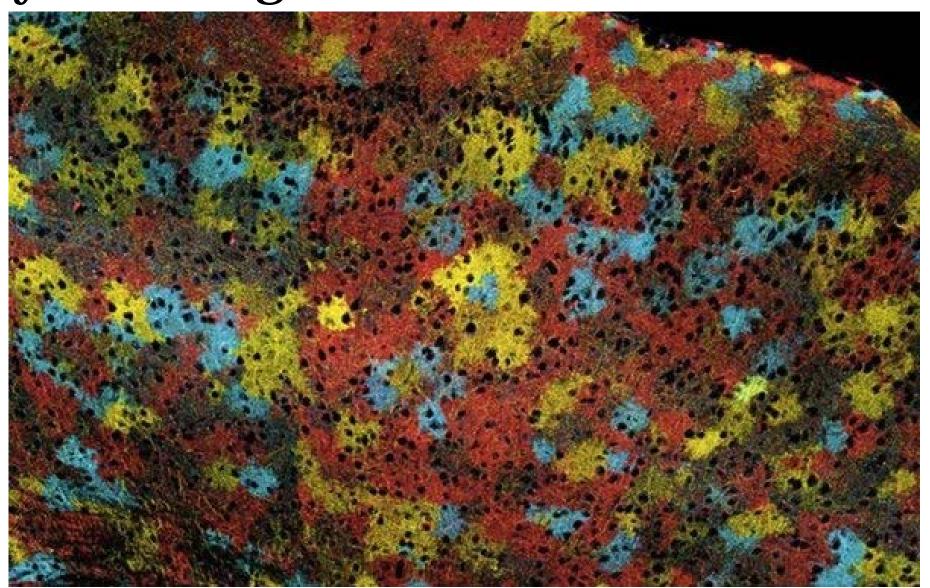


[ACM ICONS '18] [ACM BI '18]

[IEEE BHI '19]



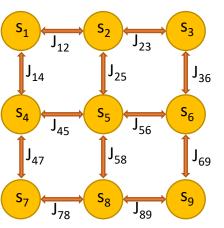
Astrocyte: A single cell Detects Neuronal Chaos





Synaptic Activity as an Ising Model

- Spins $s_i, s_j \in \{1, -1\}$
- Uniform Couplings $\forall i, j J_{ij} = 1$

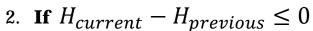


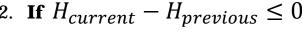
• **Why**?

- Fraiman et al. 2009
- Brain-like near-critical dynamics
 - Second-order phase transition between order and chaos
- Online control of dynamics using a <u>single</u> parameter *T*



- Minimize Energy function: $H = \sum_{i=1}^{n} -J_{ij}s_is_j$
- MC algorithm:
- 1. Make Random Spin flip



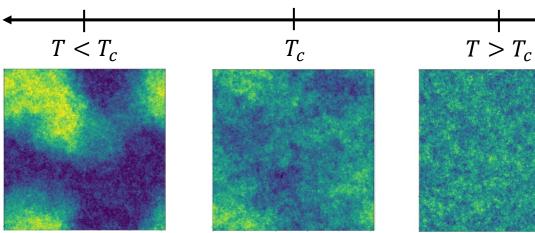


Accept flip

Else

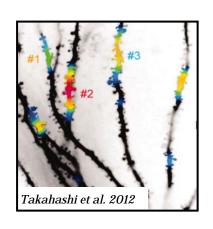
Accept flip with probability: $p = e^{-(H_{current} - H_{previous})/kT}$

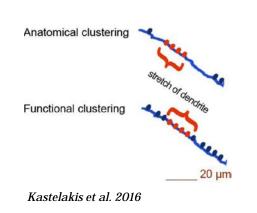
sotropic

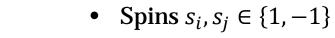




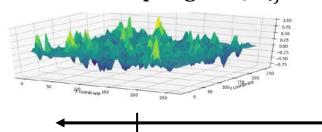
Synaptic Activity as an Ising Model

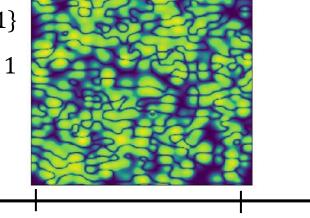








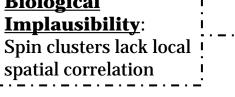


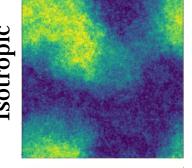


Experimental Data:

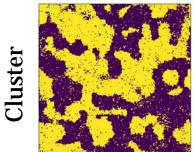
- Spatially correlated synaptic activity
- Functional and synchronous clusters i of neurons

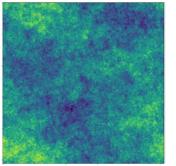


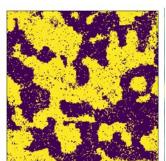


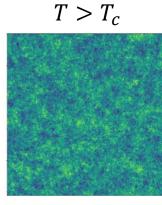


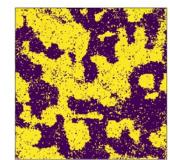
 $T < T_c$













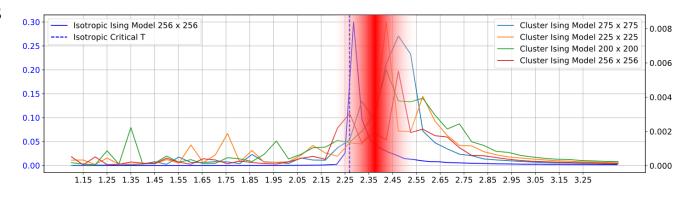
Finding 2^{nd} order Phase Transition in the Ising Model

- Confirmed phase transition with measures of variability in macroscopic Ising variables
- Magnetic susceptibility:

$$\chi = \frac{1}{T} < M^2 > - < M >^2$$

M Ising magnetization for N spins, $\frac{\sum_{i}^{N} s_{i}}{N}$ <> temporal average

Magnetic Susceptibility

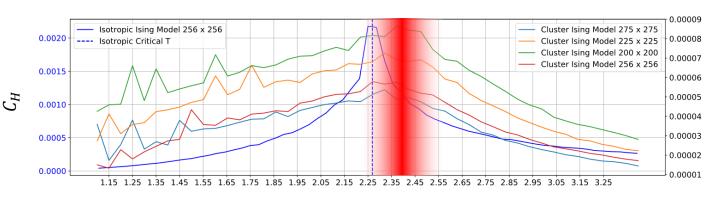


Heat Capacity

• Heat capacity:

$$C_H = \frac{1}{T^2} < H^2 > - < H >^2$$

H is system energy



 $< Degree > \approx 26$



Brain-Like Critical Transition in the Ising Model

- Correlation Network
 - Constructed pair-wise spin correlation matrix *C*

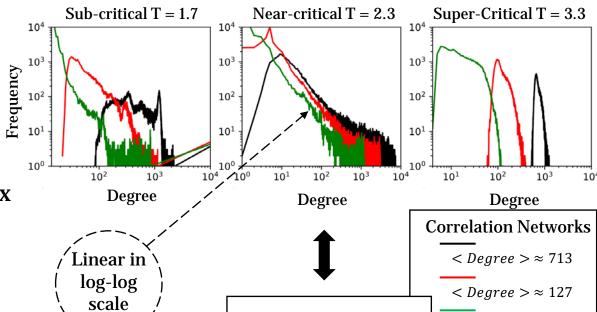
$$C_{ij} = \frac{\langle s_i(t)s_j(t) \rangle - \langle s_i(t) \rangle \langle s_j(t) \rangle}{\left(\langle s_i^2(t) \rangle - \langle s_i(t) \rangle^2\right)\left(\langle s_j^2(t) \rangle - \langle s_j(t) \rangle^2\right)}$$

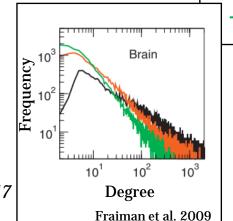
- Selected various correlation thresholds ρ
- Binarize correlation network connectivity matrix

$$W: W_{ij} = \begin{cases} 1, & C_{ij} > \rho \\ 0, & C_{ij} \le \rho \end{cases}$$

- Power law scaling at T_{nc}
 - Agrees with second order phase transition
 - Near-critical dynamics similar to fMRI data -

Correlation Network Degree Distributions





<u>Disclaimer</u>

Power-law statistics alone are not sufficient indications of criticality

Touboul & Destexhe 2017



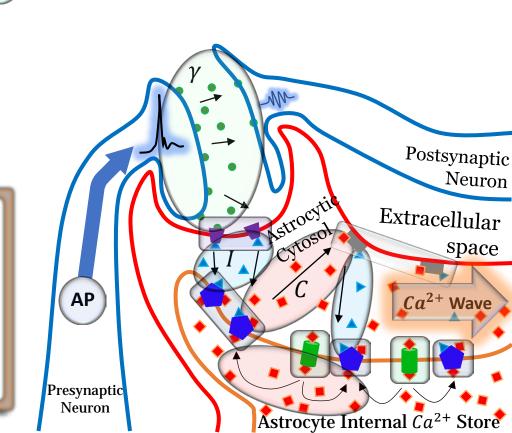
From Synaptic Activity to Astrocytic Ca²⁺ waves

• Comprehensive 3D model of astrocytic biochemical pathway:

IP3:
$$\frac{dI}{dt} C \text{ viosoffic IP3 (I)} \left(\frac{V^{0.7}}{I + K_R} \left(1 + \left(\frac{K_p}{K_R} \right) \left(\frac{C}{C + K_\pi} \right) \right) \right) + \left(\frac{V_{\delta}}{1 + \frac{I}{K_{\delta}}} \right) \left(\frac{C^2}{C^2 + K_{PLC\delta}} \right) - v_{3K} \left(\frac{C^4}{C^4 + K_D} \right) \left(\frac{I}{I + K_3} \right) - r_{5P} I$$

$$Ca^{2+}$$
: $\frac{dC}{dt}$ v_{ER} v_{ER} v_{ER} v_{ER} v_{ER}

- Quite i He quad suit is i suit subte
- Located on a serio strate delicitario
- Removed from cytosol by SERCA
- Rumps



De Pitta et al. 2009

From Synaptic Activity to Astrocytic Ca²⁺ waves

$$IP3: \quad \frac{dI}{dt} = \underbrace{\frac{1}{N}}_{V_{\beta_{i}}} \underbrace{\frac{\gamma^{0}7\gamma_{i}^{0.7}}{v_{\beta_{i}}} \underbrace{\frac{\gamma^{0}7\gamma_{i}^{0.7}}{K_{R}} \underbrace{\left(\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{R}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_{P}}\underbrace{\frac{K_{P}}{K_$$

$$Ca^{2+}: \quad \frac{dC}{dt} = (r_C m_\infty^3 n_\infty^3 h^3 + r_L)(C_0 - (1 + c_1)C) - v_{ER} \left(\frac{C^2}{C^2 + K_{ER}^2}\right)$$

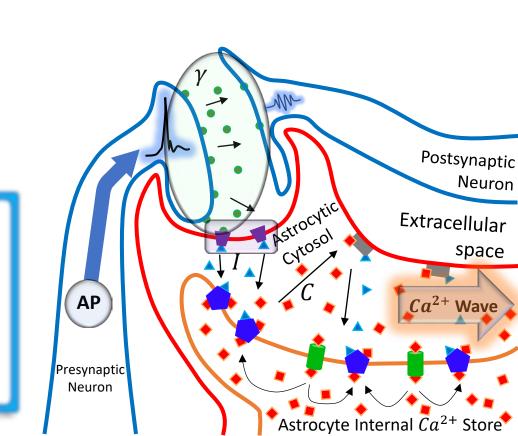
$$h: \quad \frac{dh}{dt} = \quad \frac{h_{\infty}(C, I) - h}{\tau_h(C, I)}$$

Deputations:

• **Expression are the Expression principle** (\mathbf{tis}_{β_i}) per synapse i



De Pitta et al. 2009



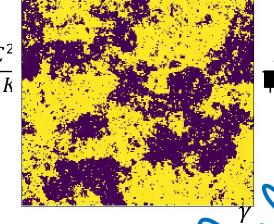


From Synaptic Activity to Astrocytics Ca2+ waves raining Rates r

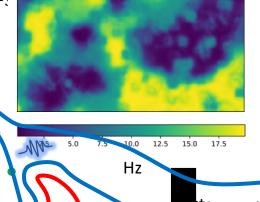
$$IP3: \quad \frac{dI}{dt} = \frac{1}{N} \sum_{i=1}^{N} \left[v_{\beta_i} \left(\frac{\gamma_i^{0.7}}{\gamma_i^{0.7} + K_R \left(1 + \left(\frac{K_p}{K_R} \right) \left(\frac{C}{C + K_{\pi}} \right) \right)} \right) \right] + \left(\frac{v_{\delta}}{1 + \frac{I}{\kappa_{\delta}}} \right) \left(\frac{C^2}{C^2 + K} \right)$$

$$\frac{Ca^{2+}}{dt} = (r_C m_\infty^3 n_\infty^3 h^3 + r_L)(C_0 - (1+c_1)C) - v_{ER} \left(\frac{C^2}{C^2 + K_{ER}^2}\right)$$

$$h: \quad \frac{dh}{dt} = \quad \frac{h_{\infty}(C, I) - h}{\tau_h(C, I)}$$



Experimental Data:

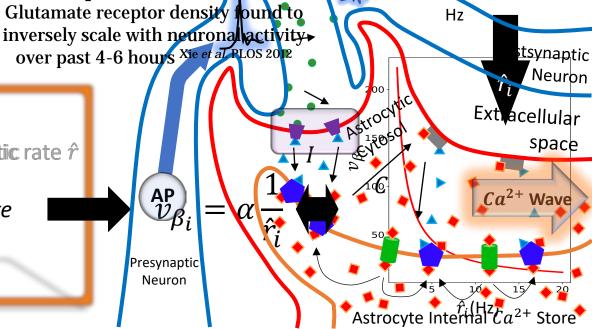


Oleight déforations:

- ${\it Dephinology}$ inputs
- Lieverrsien grædicheger eta Grtstringarte Receptor density (v_{eta_i})

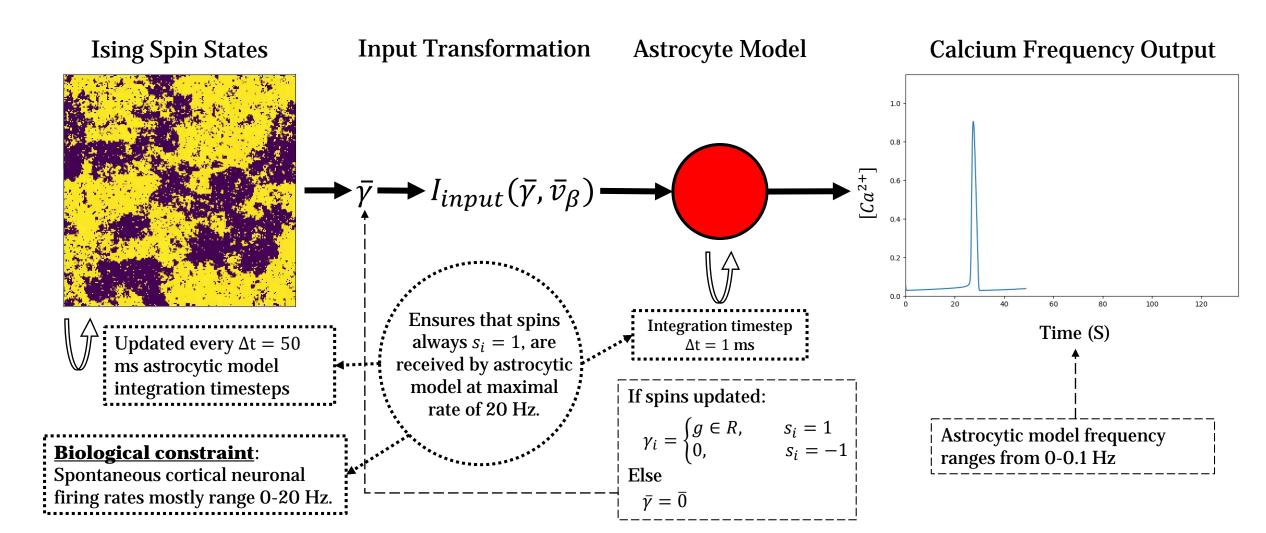


De Pitta et al. 2009



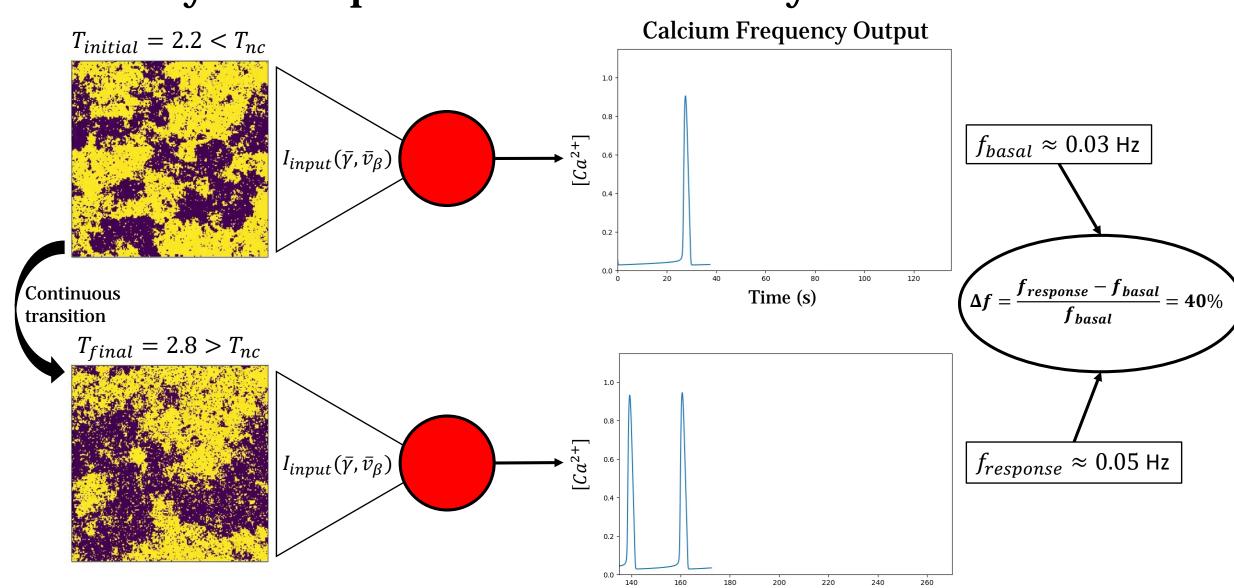
Simulation

• Ising model spin states were transformed into biological scales of time and quantity





Astrocytic Response to Neuronal Dynamics



Time (s)

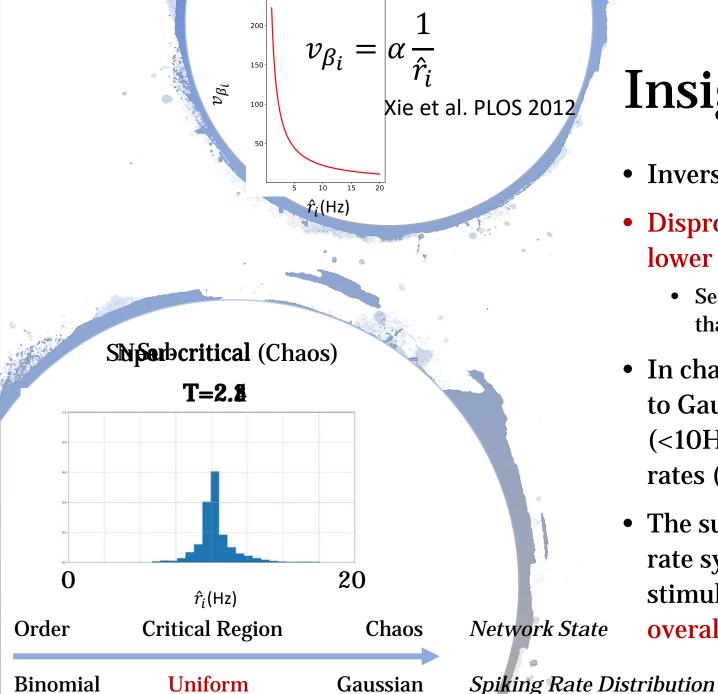
Both near-critical and chaotic T_{ini} did not alter astrocytic response



Astrocyte detects transitions from order to disorder

- Astrocytic response existed only for transition $T_{ini} < T_{near\ critical} \rightarrow T_{final} > T_{nc}$
- Response averaged over 5 cluster Ising models
 - Each transition $T_{ini} \rightarrow T_{final}$ was averaged over 10 simulations

Sub-critical $T_{ini} = 2.2$ Near-critical $T_{ini} = 2.3$ Super-critical $T_{ini} = 2.5$ Astrocytic frequency changes at the 0.07 transition across the near-critical range, $T_{nc} = [2.25, 2.45]$, where a 2nd Fitted Hyperbolic Tangent 0.06 0.06 0.06 $[Ca^{5+}]_{0.05}$ Basal Rate order phase transition exists Response Rate 0.05 0.05 between order and chaos. 0.04 -0.5 -0.50.0 0.5 1.0 -1.0Fitted Hyperbolic Tangent 20 20 Hp-GCh-I % Change in Rate 10 -1.0Changes in magnetization did $^*M^{\sim}$ not drive astrocytic response $\Delta M^* = < M_{T_{final}} > - < M_{T_{initial}} >$ 1.0 1.0 ΔT ΔT ΔT



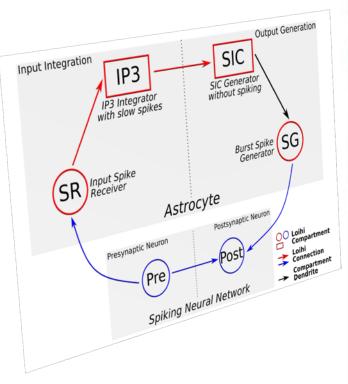
Is this magic? Insight

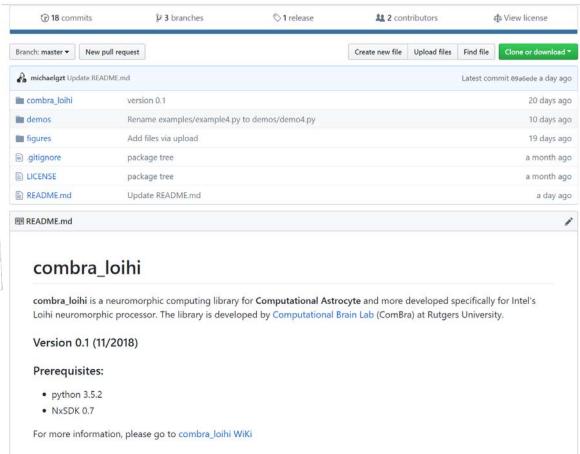
- ?
- Inverse scaling in learning the Glu receptor density
- Disproportional learning of Glu weights between lower synaptic rates and higher synaptic rates
 - Sensitivity to Glu in lower-rate synapses is much higher than the insensitivity of the higher rate synapses
- In chaos, the distribution of spiking rates changes to Gaussian μ =10Hz: Synapses that had low rates (<10Hz) increase to ~10Hz and synapses with high rates (>10Hz) decrease to ~10Hz
- The sudden increase in rates in the learned as lowrate synapses provides disproportionally larger Glu stimulation to the astrocyte, which results in an overall increase in Ca²⁺ frequency



Astrocytic module @ Loihi chip

released December 2018







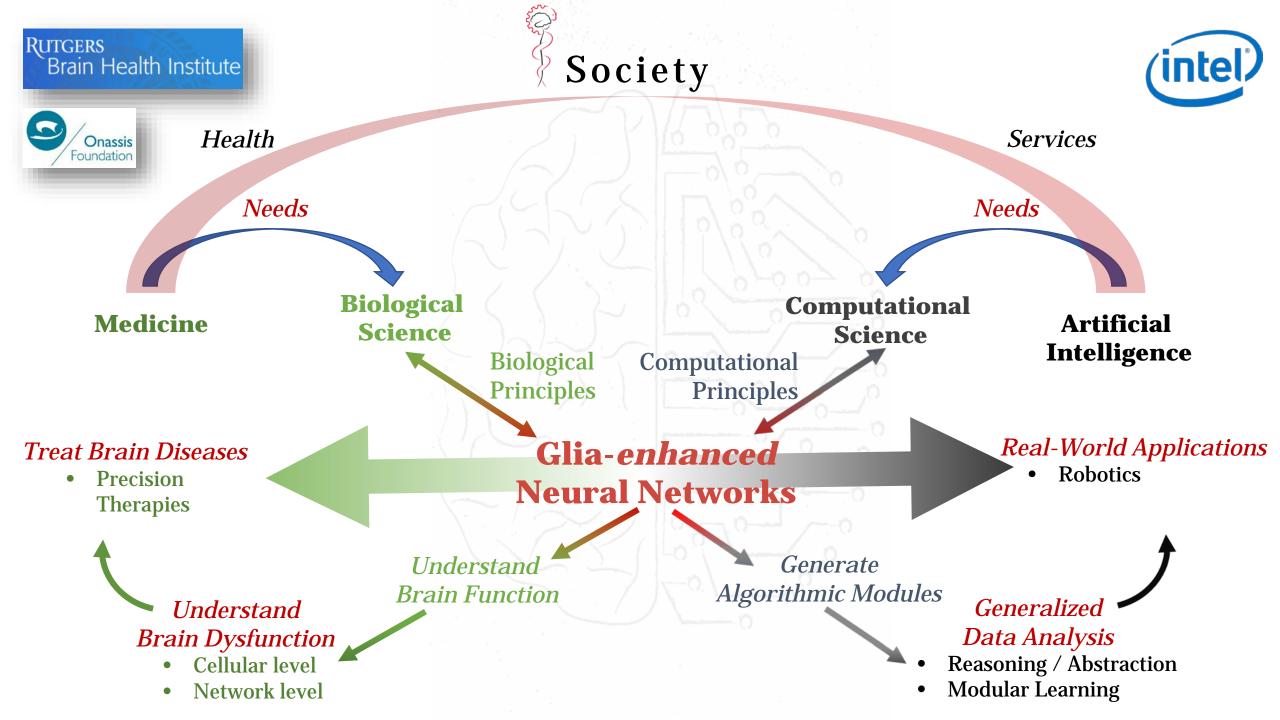


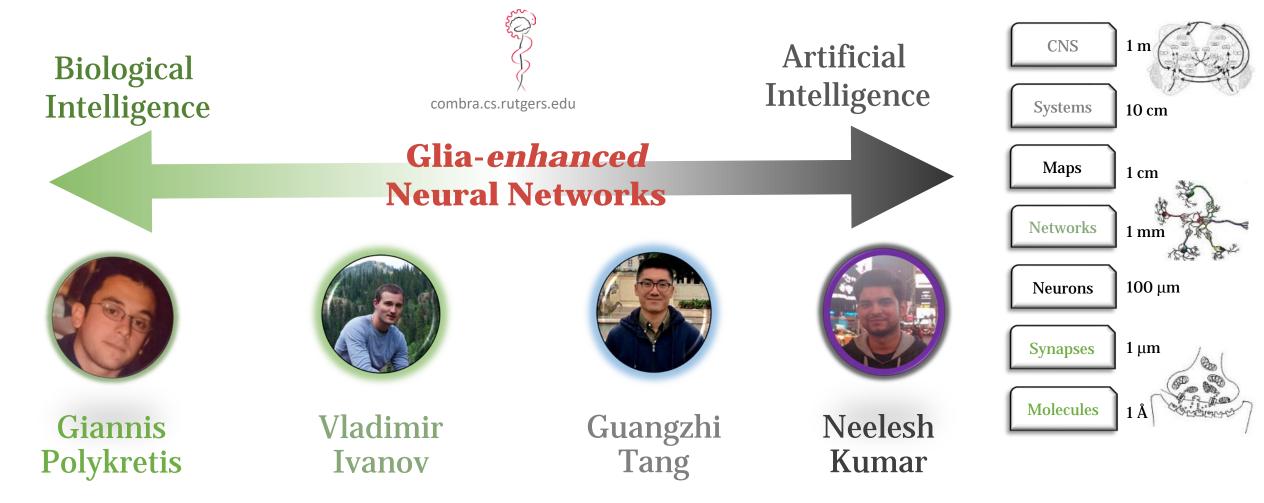


Computations at the edge of Chaos

- Carrying out computations at the edge of chaos is based on the hypothesis that complex systems can show an extended computational power when performing in between ordered and chaotic dynamics
- A Liquid State Machine whose reservoir shows critical dynamics could perform efficient computations in real-time
- Not much is yet known about the computational power of neural networks that are constructed consistently with neurobiology

• Our results pave the way for an SOC, self-organized criticality





Astrocytic Mechanisms for Astrocytic Computation for Neuromorphic Hardware for *Less* Machine Learning

- Neural Synchronization Network Oscillations
- **Astrocytic Processing**
- Reservoir Networks

Synaptic Changes

- Order/Chaotic network •
- **Astrocytic Learning**
- **Learning Methods**



Computational Astrocyence @ ComBra Lab

- G Tang, A Shah, **KP Michmizos**, "<u>Spiking Neural Network on Neuromorphic Hardware for Energy-Efficient Unidimensional SLAM</u>", arXiv preprint arXiv:1903.02504, **2019**
- I Polykretis, V Ivanov, **KP Michmizos**, "Computational Astrocyence: Astrocytes encode inhibitory activity into the frequency and spatial extent of their calcium elevations," IEEE Brain Health Informatics, Chicago, May 2019
- V Ivanov, I Polykretis, **KP Michmizos**, "Axonal Conduction Velocity Impacts Neuronal Network Oscillations," IEEE Brain Health Informatics, Chicago, May 2019
- I Polykretis, V Ivanov, **KP Michmizos**, "The Astrocytic Microdomain as a Generative Mechanism for Local Plasticity," International Conference on Brain Informatics, **2018** (pp.10)
- *I Polykretis*, *V Ivanov*, **K.P. Michmizos**, " <u>A Neural-Astrocytic Network Architecture: Astrocytic calcium waves modulate synchronous neuronal activity</u>," International Conference on Neuromorphic Systems (ICONS), Knoxville, Tennessee, July 23-26, 2018 (pp. 8)
- *G Tang*, **KP Michmizos**, " <u>Gridbot: An autonomous robot controlling by a spiking neural network mimicking the brain's navigational system</u>," International Conference on Neuromorphic Systems (ICONS), Knoxville, Tennessee, July 23-26, 2018 (pp. 8)
- *G. Tang*, **KP Michmizos**, " <u>Gridbot: A Spiking Neural Network Model of the Brain's Navigation System for Autonomous Robots</u>," 6th Neuro Inspired Computational Elements Workshop (NICE 2017), Intel Hillsboro, OR, February 27th- March 1st, 2018
- *L. Kozachkov*, **KP. Michmizos**, "<u>Brain-morphism: Astrocytes as Memory Units</u>," 6th Neuro Inspired Computational Elements Workshop (NICE 2017), Intel Hillsboro, OR, February 27th- March 1st, 2018
- L. Kozachkov, KP Michmizos, "A Computational Role for Astrocytes in Memory," arXiv:1707.05649, 2017
- L. Kozachkov, **KP Michmizos**, " The Causal Role of Astrocytes in Slow-Wave Rhythmogenesis: A Computational Modelling Study," arXiv:1702.03993, 2017





Towards Critical SNNs: Astrocytes Detect Chaos in Neuronal Dynamics

Back-UP Slides

Konstantinos Michmizos

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Computer Science | Rutgers University

NANS=Neural-Astrocytic Networks

Research on capsules is now at a similar stage to research on recurrent neural networks for speech recognition at the beginning of this century. There are fundamental representational reasons for believing that it is a better approach but it probably requires a lot more small insights before it can out-perform a highly developed technology. The fact that a simple capsules system already gives unparalleled performance at segmenting overlapping digits is an early indication that capsules are a direction worth exploring. New time and learning dimensions