

Larger Connection Radius Increases Hub Astrocyte Number in a 3-D Neuron–Astrocyte Network Model

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Abstract—Astrocytes - a prominent glial cell type in the brain - form networks that tightly interact with the brain’s neuronal circuits. Thus, it is essential to study the modes of such interaction if we aim to understand how neural circuits process information. Thereby, calcium elevations, the primary signal in astrocytes, propagate to the adjacent neighboring cells and directly regulate neuronal communication. It is mostly unknown how the astrocyte network topology influences neuronal activity. Here, we used a computational model to simulate planar and 3D neuron-astrocyte networks with varying topologies. We investigated the number of active nodes, the shortest path, and the mean degree. Furthermore, we applied a graph coloring analysis that highlights the network organization between different network structures. With the increase of the maximum distance between two connected astrocytes, the information flow is more centralized to the most connected cells. Our results suggest that activity-dependent plasticity and the topology of brain areas might alter the amount of astrocyte controlled synapses.

Index Terms—Simulation, astrocytes, gap junctions, neurons, network topology.

I. INTRODUCTION

THE PROPAGATION of information inside the brain is historically considered to be based on the communication between neurons. However, recent experimental studies show that astrocytes actively modulate neuronal activity. They are tightly linked to neurons via the so-called tripartite synapse [1] forming the neuron-astrocyte molecular communication system. The primary signaling mediator evoked by the synaptic communication in astrocytes is calcium [2]–[4]. Calcium propagates inside the complex astrocytic branch trees and from cell to cell through gap junction coupling (GJC) [3], [5]. Thereby, astrocytes form non-overlapping domains coupled only to the nearest neighbors [6]. Mainly

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unknown is how the astrocyte network topology influences the neuronal activity.

Different types of computational models are used to investigate the intra- and intercellular pathways of astrocytes and the communication with neurons (reviewed in [7], [8]). Lallouette *et al.* simulated five different 3D topologies of only-astrocyte networks [9], [10], of which some included long-distance connections and hubs. The propagation range of calcium in astrocytes was mostly given by the absence of long-distance GJC and independent of the presence of hubs. The correct functioning of this network could easily be perturbed under the effect of diseases, for example, with the increase of the overall network complexity. Barros *et al.* investigated the impact of calcium propagation in astrocyte network topologies in a model with altered intracellular dynamics to mimic Alzheimer’s disease [11]. They concluded that calcium transmission was differently affected in healthy or disease states - intracellularly and in the network propagation patterns. However, this theoretical study did not include experimental data, and the signaling was only implemented from the neurons towards the astrocytes but not vice versa. Neuronal networks have also been studied using computational models, such as [12], where they analyze the activity and noise of neuronal networks under small-world topologies.

Recently, we implemented a combined neuron-astrocyte network model called INEXA [13], [14]. Therewith, we investigated the astrocyte’s influence on the neuronal network stability. The simulations resembled *in vitro* experiments with planar multielectrode arrays (MEAs). In the INEXA model, the astrocytes and neurons communicated through the tripartite synapse by exchanging activating and depressing transmitters [13]. In the model, each astrocyte was connected to several hundreds of synapses. In Genocchi *et al.* [14], we varied the number of astrocytic GJCs and the noise levels applied to the presynaptic terminals to simulate low, high, and hyperactivity. The astrocytes increasingly downregulated high and hyperactivity with increased astrocytic GJCs.

Here, we extend our previous studies with the overall objective of investigating the importance of the astrocyte network topology and the role of the most active astrocytes in regulating the network activity. Therefore, we analyze the astrocyte network organization, where nodes are classified from most influencing nodes to the least. Since network nodes with higher degrees exist, an astrocyte connected to more than 75% of its neighbors is defined as a hub astrocyte, which presents the strongest influence in the overall network activity [9], [15]. First, we create planar and 3D neuron-astrocyte networks using

the previously-mentioned INEXA model [13] and compare their characteristics. Second, we vary the distance range within which two astrocytes are coupled to each other in the 3D network. According to Lallouette *et al.* [9], we choose the link radius, which seems to reflect the biological network topology the closest. Subsequently, we quantify the resulting neuronal network activity and topological measures of the astrocytic network. Third, to link network structure to network organization, we use a graph coloring algorithm to analyze the node organization over different distance values. We show that the organization and node structure are interlinked with the astrocyte activity levels regarding distance to neighboring astrocytes and the cell's position in the network.

II. METHODS

A. Neuron-Astrocyte Network Model

In the discrete-time model INEXA [13], the neuronal firing rate λ_i of a postsynaptic neuron i is calculated for each time slice t_k of 5 ms as follows:

$$\lambda_i(t_k) = \max \left(0, c_i + \sum_j y_{ij} \cdot s_j(t_{k-1}) - \sum_j y_{Astro} \cdot A_{ija}(t_{k-1}) \right), \quad (1)$$

where c_i denotes the noise of neuron i , which is sampled from a triangular distribution between 0 and an upper bound value, C_{max} . In the original model, c was implemented to study several noise levels of neurons. The term y_{ij} is the synaptic strength between the presynaptic neuron j to postsynaptic neuron i , which can vary over time. Thereby, the synapse can be either excitatory (y_{ij} between 0 and 1) or inhibitory (y_{ij} between -1 and 0). The parameter s_j is a binary parameter that indicates whether a spike has been emitted by neuron j in the previous time step ($s_j = 1$ if a spike has been emitted, else $s_j = 0$). The second part in the equation denotes the depressing effect exerted by the astrocytes. A_{ija} is a binary term that describes whether the synapse ij is wrapped by astrocyte "a" and if astrocyte "a" was in the active state at the previous time step. Otherwise A_{ija} is equal to zero, removing the astrocytic effect on the neuronal firing rate. If an active astrocyte wraps the synapse ij , the astrocyte applies a depressing effect, y_{Astro} , on the synapse.

The second governing equation is the one for the intracellular calcium dynamics $[Ca^{2+}]_{ija}$ in an astrocyte "a" that wraps synapse ij :

$$\begin{aligned} [Ca^{2+}]_{ija}(t_k) &= [Ca^{2+}]_{ija}(t_{k-1}) + \Omega_{acc} \\ &\times \left([IP_3]_{ija}(t_k) - [Ca^{2+}]_{ija}(t_{k-1}) \right). \end{aligned} \quad (2)$$

The calcium concentration is a sum of the calcium concentration remained from the last time slice ($Ca_{ija}^{2+}(t_{k-1})$), of the IP_3 -mediated Ca^{2+} -induced Ca^{2+} -release from the endoplasmic reticulum (ER) stores, and of the uptake of Ca^{2+} back

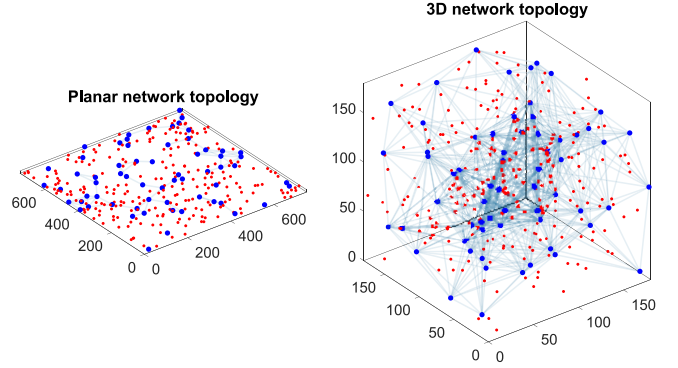


Fig. 1. Topologies for the planar (left) and 3D (right) neuron-astrocyte network. Note, both networks have the same total volume. Astrocytes and their connections are shown in blue and neurons in red. Axes are in μm .

to the ER by the SERCA pumps. The ER regulation of Ca^{2+} concentration is modeled in the second part of the equation. To reflect the slow dynamics of the calcium release (up to seconds) we multiplied the ER term by the time scale Ω_{acc} .

We extended the INEXA model [13] with the following modifications: To change from a planar to a 3D topology, we kept a similar volume as described in this letter (Figure 1). The network space from planar dimensions ($750 \times 750 \times 10 \mu m^3$, which resembles the culture area of a planar MEA) was altered to $180 \times 180 \times 180 \mu m^3$ in 3D. We used 250 neurons with 80% excitatory and 20% inhibitory, and 63 astrocytes for all simulations. The neuron-neuron and astrocyte-neuron connections, as well as the GJC, was modeled as described in Lenk *et al.* [13].

In the first step, the astrocytes were connected based on the spatially-constrained link radius topology model [9], where all astrocytes in the distance d equal to $100 \mu m$ were connected in the planar dimension and 3D. In the second step, we concentrated solely on 3D and ranged d between 70 to $120 \mu m$. For each distance d , the simulation was run ten times with the same topology, each with a simulated time of five minutes.

Initially, we fixed the upper boundary for the neuronal noise level C_{max} to 0.02 and the upper boundary for synaptic weights to 0.7 and -0.7 for excitatory and inhibitory neurons, respectively [13]. Keeping the rules for creating the topologies for the planar and 3D networks resulted in very high connectivity in 3D ($>80\%$). Thus, we reduced the probability that two neurons connect with each other, which resulted in similar connection numbers (Table I). We then obtained the neuronal activity as spikes (i.e., Dirac function of action potentials) per cell over time.

B. Graph Theory Analysis

We developed a graph of the 3D astrocyte network $G_{astro} = (V_{astro}, E_{astro})$, where V_{astro} was the set of vertices (cell bodies) and E_{astro} the set of nodes (connection between two cells). We collected V_{astro} and E_{astro} based on the topology of the INEXA model. For each simulation, we determined the total number of astrocytes activated at least once during a simulation, N_{act} . We calculated the mean degree k , which quantifies the number of connected cell pairs (number of GJCs). Furthermore, the number of hub cells, i.e., nodes

TABLE I

COMPARISON OF THE CHARACTERISTICS OF THE PLANAR AND THE 3D NETWORKS. SPIKE RATE, BURST RATE, AND BURST DURATION ARE FEATURES OF THE NEURONAL NETWORK ACTIVITY AND DISPLAYED WITH MEAN AND STANDARD DEVIATION OVER THE TEN SIMULATION RUNS. MEAN DEGREE k , SHORTEST PATH L AND NUMBER OF ACTIVE CELLS N_{act} ARE FEATURES OF THE ASTROCYTE NETWORK TOPOLOGY. k AND L ARE DISPLAYED WITH MEAN AND STANDARD DEVIATION OVER ALL NODES. N_{act} HAS THE MEAN AND STANDARD DEVIATION OVER ALL SIMULATION RUNS

Characteristics	Planar network	3D network
Dimensions [μm^3]	750x750x10	180x180x180
Cell numbers	250 neurons, 63 astrocytes	250 neurons, 63 astrocytes
Noise and synaptic strength	$c = 0.02, y = \pm 0.7$	$c = 0.02, y = \pm 0.7$
Spike rate [spikes/min]	69.18 ± 3.17	301.47 ± 37.79
Burst rate [bursts/min]	1.08 ± 0.17	26.80 ± 4.05
Burst duration [ms]	36.22 ± 2.76	193.89 ± 46.93
Max. amount of NN connections	62250	62250
Average neuron connections to other neurons	66.52	69.31
Network connectivity [%]	26.72	27.83
Average neurons connection length [μm]	209.67	101.55
Two directional connections between neurons	4924	2719
Average gap junction connections	2.70	19.61
Highest / lowest gap junction amount	7 / 0	35 / 7
Average distance between connected astrocytes [μm]	66.82	72.05
Average number of astrocyte-neuron connections	169.91	220.85
Excitatory synapses without an astrocyte	2500	0
% of not astrocyte controlled excitatory synapses	18.93	0.00
Mean degree k	2.70 ± 1.53	19.61 ± 7.30
Shortest path L	[(a) 4.57 ± 2.37 , (b) 3.89 ± 2.20 , (c) 1.33 ± 0.49]	1.82 ± 0.65
Number of activated cells N_{act}	8.00 ± 3.20	4.20 ± 0.91

which are coupled to more than 75% of the neighbors, was determined. The shortest path L denoted the minimal number of GJCs one must cross to connect the two astrocytes. Finally, we employed a graph coloring algorithm based on greedy coloring, i.e., on a coloring spectrum with the maximum amount of colors equal to the maximum number of degree. Our greedy coloring scheme is close to a Welsh-Powell algorithm [15], but with fewer colors available since we adjusted that number to match the number of degrees we got from the 3D astrocyte topologies.

C. Quantification of the Neuronal Activity

For all simulations in the planar dimension and 3D, we analyzed the spontaneous activity inside the neuronal network. From the activity-temporal series of each neuron, we calculated the features of spike rate, burst rate, and burst duration, according to [16]. We averaged each feature over all neuronal cells in each simulation and over the number of simulation runs per distance d .

III. RESULTS

A. Comparison of Planar and 3D Networks

As we kept the cell number the same and the percent of connectivity between the cells similar, it resulted in shorter neuronal connections in 3D than in the planar networks (Table I). However, the average distance between astrocytes increased from planar to 3D. Less bidirectional connections between a pair of neurons were counted. As there were more neighbors in a certain distance in 3D, the number of gap junctions between astrocytes and the average number of neuron-astrocyte connections was increased. In 3D, all excitatory synapses were occupied by an astrocyte branch, unlike in the planar network.

Measuring the neuronal activity showed a five-fold higher spike rate in 3D (Table I). Also, the burst rate and duration significantly increased when changing from a planar to a 3D neuron-astrocyte network. The graph theory analysis revealed that the mean degree k increased from planar (2.70 on average) to 3D (19.51 on average). In the planar networks, the network was divided into three subnetworks (Table I, (a)-(c) for parameter L). The number of active astrocytes, N_{act} , decreased with higher dimensionality.

B. Alteration of the Link Radius of Astrocytes in 3D Networks

The graph analysis for the astrocytes networks showed that the mean degree k increased when increasing the distance d between astrocytes (Figure 2). The shortest path L instead decreased since the increase in connections available for each node of the network allows cells to reach other cells with fewer node hops. The number of active astrocytes N_{act} varied significantly over the ten simulations ran for each distance (Figure 2), especially in the simulations with short distances ($d = 70 \mu\text{m}$ and $80 \mu\text{m}$). Also, N_{act} showed a decreasing tendency when increasing the distance. In Figure 3, the network organization for astrocytes with link distances of 70 and 120 μm is displayed. The color and size of the node varied with the number of connections per node ranging from 2 to 16 for $d = 70 \mu\text{m}$ and 10 to 51 for $d = 120 \mu\text{m}$. In the case of $d = 120 \mu\text{m}$, we counted four hub astrocytes (indicated by an arrow in Figure 3) and none for the other distances. This indicates that extending the distance d increases the network connectivity. Hence, the nodes with the highest degrees were found in the center of the topology.

The neuronal activity describing features spike rate, burst rate, and burst duration increased from $d = 70 \mu\text{m}$ to

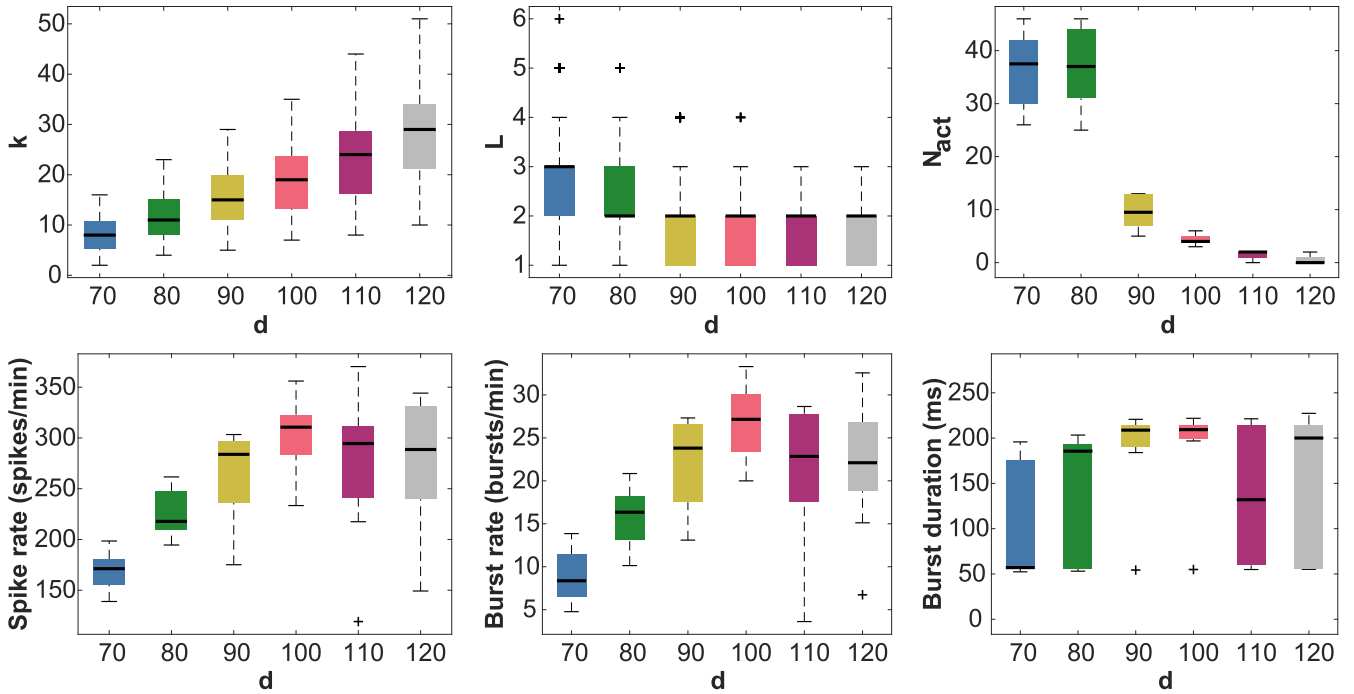


Fig. 2. Astrocytic network features, such as mean degree k , shortest path L and number of active astrocytes N_{act} , for each link radius d are shown in the upper row. Neuronal features, such as spike rate (spikes/minute), burst rate (bursts/minute), and burst duration (ms) are displayed in the lower row. The bold lines in the box plots represent the median and the crosses represent the outliers. The lower and upper whiskers represent the 25th and 75th percentiles of the data distribution, respectively.

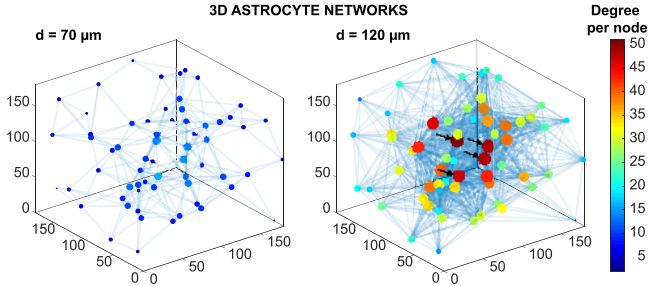


Fig. 3. Colored graphs for the astrocyte networks build with distance $d = 70 \mu\text{m}$ (left) and $d = 120 \mu\text{m}$ (right) indicating the degree per node using the greedy coloring scheme. Arrows indicate the hub astrocytes (connected to more than 75% of their neighbors). Axes are in μm .

$d = 100 \mu\text{m}$. For $d = 110 \mu\text{m}$ and $d = 120 \mu\text{m}$, they decreased again (Figure 2). Figure 4 displays the spike rate per neuron for each link radius d of the astrocytic network, respectively. Here, an example of how the spike rates were distributed in the neuronal network. The figure was reconstructed with data obtained from just one of the ten simulation runs per d . Since the neuronal network topology was fixed for all d , the spike rate variability across the six subfigures was a reflection of the astrocytic N_{act} .

IV. DISCUSSION AND CONCLUSION

In the last two decades, *in vitro* MEA experiments moved from planar to 3D in terms of the electrode shape [17], [18] and/ or scaffolds (e.g., by using hydrogels or microbeads) [19]–[21]. In 3D networks, Vogel [21] measured similar neuron projections ranging from 80 to 120 μm as we simulated in this study. In addition, when shifting from planar to 3D networks, she observed an increase of spontaneous

neuronal activity [21]. Noteworthy, our comparison between the planar and 3D network topologies shows that the astrocytes fully control the neuronal network in 3D. This increase in synaptic coverage is accompanied by a decreased astrocytic activation leading to a higher neuronal activity, which again highlights the astrocytic effect on controlling the neuronal activity as already shown in Lenk *et al.* [13] and Genocchi *et al.* [14]. This observation is relevant since the number of synapses ensheathed by astrocytes seem to vary between and within brain areas [22]. Our results suggest that activity-dependent plasticity, i.e., potentiation or depression of neurotransmitter release, and the topology of brain areas might alter the amount of astrocyte controlled synapses. The results from our graph analysis in the 3D network show that a shorter distance d between two astrocytes is followed by a larger number of active astrocytes N_{act} and shortest paths L , as well as a lower mean degree k , which is in accordance with Lallouette *et al.* [9], [10]. To summarize, longer distances between astrocytes seem to centralize the information propagation in astrocytes.

We will evaluate how structure and activity will be interlinked with network organization and information propagation in the next step. This quantification will help to understand the neuron-astrocyte dynamics and how the brain structure is plastic based on the cell molecular communications properties and patterns.

SOFTWARE AND DATA AVAILABILITY

The code and the resulting data used in this study can be found at https://github.com/kerstinlenk/INEXA_IIEEETransMolBiolMulti-ScaleCommun2020. The MATLAB code for the published INEXA model [13] is available in a

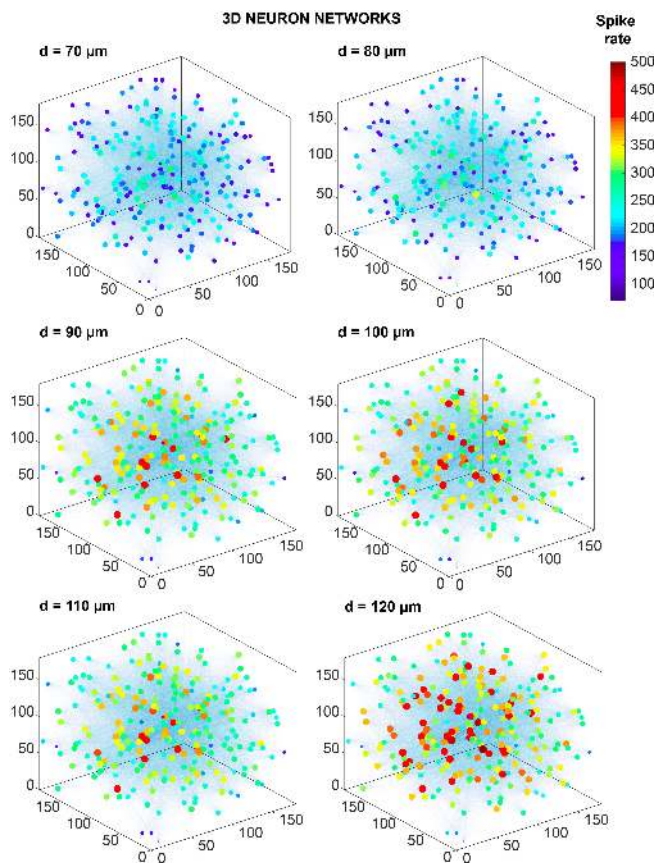


Fig. 4. Colored graphs for the neuronal networks, indicating the spike rate (spikes/min) per node using the greedy coloring scheme. Note that the neuronal network topology and the connections between neurons is the same for all the six subplots. The distances d , from $d = 70 \mu\text{m}$ to $d = 120 \mu\text{m}$, represent the link radii for the astrocyte network. Axes are in μm .

publicly accessible repository: https://github.com/kerstinlenk/INEXA_FrontCompNeurosci2020. The code for the burst analysis tool is stored at <https://doi.org/10.5281/zenodo.3883622>.

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