

POSTER PRESENTATION

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# Modeling astrocyte-neuron interactions in a tripartite synapse

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From The Twenty Third Annual Computational Neuroscience Meeting: CNS\*2014  
Québec City, Canada. 26-31 July 2014

Glial cells (microglia, oligodendrocytes, and especially astrocytes) play a critical role in the central nervous system by affecting in various ways the neuronal single cell level interactions as well as connectivity and communication at the network level, both in the developing and mature brain. Numerous studies (see, e.g., [1-3]) indicate an important modulatory role of astrocytes in brain homeostasis but most specifically in neuronal metabolism, plasticity, and survival. Astrocytes are also known to play an important role in many neurological disorders and neurodegenerative diseases. It is therefore important in the light of recent evidence to assess how the astrocytes interact with neurons, both in situ and in silico. The integration of biological knowledge into computational models is becoming increasingly important to help understand the role of astrocytes both in health and disease. We have previously addressed the role of transmitters and amyloid-beta peptide on calcium signals in rat cortical astrocytes [4]. In this work, we extend the work by using a modified version of the previously developed model [5] for astrocyte-neuron interactions in a tripartite synapse to explore the effects of various pre- and postsynaptic as well as extrasynaptic mechanisms on neuronal activity. We consider extending the model to include various additional mechanisms, such as the role of IP3 receptor function, recycling of neurotransmitters, K<sup>+</sup> buffering by the Na<sup>+</sup>/K<sup>+</sup> pump, and retrograde signaling by endocannabinoids. The improved tripartite synapse model for astrocyte-neuron interactions will provide an essential modeling tool for facilitating studies of local network dynamics in the brain. The model may also serve as an important step toward understanding

mechanisms behind induction and maintenance of plastic changes in the brain.

## Acknowledgements

The support from Tampere University of Technology Graduate School (R.H.) and Tampere University of Technology Foundation (M.-L.L.) is acknowledged.

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Published: 21 July 2014

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doi:10.1186/1471-2202-15-S1-P98

Cite this article as: Linne et al.: Modeling astrocyte-neuron interactions in a tripartite synapse. *BMC Neuroscience* 2014 **15**(Suppl 1):P98.

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