POSTER PRESENTATION



Open Access

Short desynchronization epochs in neural synchronization: detection, mechanisms, and functions

Leonid L Rubchinsky^{1,2*}, Sungwoo Ahn^{1,3}

From 24th Annual Computational Neuroscience Meeting: CNS*2015 Prague, Czech Republic. 18-23 July 2015

Synchrony of neural oscillations is believed to play important role in a variety of functions and dysfunctions of neural systems. However, neural synchrony can hardly be perfect for prolong intervals of time. If there is a significant synchrony strength observed over sufficiently long time interval, the imperfect nature of synchrony implies that for some intervals of time synchrony may be stronger, while for some intervals of time in may be weaker. If neural synchrony is to play some functional role, this role may depend on how synchrony is patterned in time. Roughly speaking, few long intervals of synchronous (or nonsynchronous) dynamics may be functionally different from many short intervals, although the synchrony strength may be the same on the average.

Recent developments in time-series analysis allowed exploring the patterning of synchrony in time on very short time-scales. If there is some synchrony level is present on the average, than one can look at each cycle of oscillations and detect of signals are synchronous or not resulting in a quantitative description of succession of episodes of synchronization and desynchronization [1]. We applied these methods to analyze neural signals of various natures: synchronization of single units and LFPs in subthalamic nucleus of Parkinsonian patients [2], synchronization of EEG signals in healthy humans [3], synchronization of LFPs within and between prefrontal and hippocampal circuits in normal rats and rats experiencing repetitive psychostimulant injections [4]). The temporal patterning of the neural synchrony was sensitive to behavioral states, drug status etc. However, all these studies yielded one important qualitative results: neural oscillations were observed to go out of synchrony primarily for a short time interval. They may go out of synch very frequently (leading to low synchrony levels), but all these different neural systems exhibit mostly very short desynchronization intervals (different from rare prolongs desynchronizations and a spectrum of possibilities in between, none of which were experimentally observed). On the contrary, non-neural systems can exhibit longer desynchronization durations [1,5].

These experimental results suggest that short desynchronization dynamics may be universal for synchronized activity of neural systems. Here we review the data analysis techniques and results of their application to experimental data. We then consider the potential mechanisms of the apparent universality of short desynchronization dynamics and its potential functional advantages. We consider simple neural circuits of Morris-Lecar-type neurons and identify conditions imposed on the parameter values to result in both short desynchronization dynamics and dynamics exhibiting longer and rarer desynchronized intervals. By maintaining the same levels of average synchrony (and the same frequency of oscillations) we show that fast conductances promote short desynchronization dynamics observed in experiments. By subjecting these model networks to oscillatory synaptic input, we found that systems with short desynchronization dynamics require weaker input signal in order to synchronize with this input. We finally conjecture that properties of ionic conductances responsible for short desynchronization dynamics may make neural systems more adaptable, as they are more efficient in generating transient synchronous patterns (synchronous assemblies) in response to synaptic or sensory inputs.



© 2015 Rubchinsky and Ahn This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited. The Creative Commons Public Domain Dedication waiver (http://creativecommons.org/ publicdomain/zero/1.0/) applies to the data made available in this article, unless otherwise stated.

^{*} Correspondence: leo@math.iupui.edu

¹Department of Mathematical Sciences, Indiana University Purdue University Indianapolis, Indianapolis, IN, USA

Full list of author information is available at the end of the article

Authors' details

¹Department of Mathematical Sciences, Indiana University Purdue University Indianapolis, Indianapolis, IN, USA. ²Stark Neurosciences Research Institute, Indiana University School of Medicine, Indianapolis, IN, USA. ³Present address: School of Mathematical and Statistical Sciences, Arizona State University, Tempe, AZ, USA.

Published: 18 December 2015

References

- Ahn S, Park C, Rubchinsky LL: Detecting the temporal structure of intermittent phase locking. Phys Rev E 2011, 84(1-2):016201.
- Park C, Worth RM, Rubchinsky LL: Fine temporal structure of beta oscillations synchronization in subthalamic nucleus in Parkinson's disease. J Neurophysiol 2010, 103(5):2707-2716.
- Ahn S, Rubchinsky LL: Short desynchronization episodes prevail in the synchronous dynamics of human brain rhythms. *Chaos* 2013, 23(1):013138.
- Ahn S, Rubchinsky LL, Lapish CC: Dynamical reorganization of synchronous activity patterns in prefrontal cortex - hippocampus networks during behavioral sensitization. *Cerebral Cortex* 2014, 24(10):2553-2561.
- Rubchinsky LL, Ahn S, Park C: Dynamics of synchronizationdesynchronization transitions in intermittent synchronization. Frontiers in Computational Physics 2014, 2:38.

doi:10.1186/1471-2202-16-S1-P3

Cite this article as: Rubchinsky and Ahn: **Short desynchronization epochs in neural synchronization: detection, mechanisms, and functions.** *BMC Neuroscience* 2015 **16**(Suppl 1):P3.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

BioMed Central

Submit your manuscript at www.biomedcentral.com/submit