

Oral presentation

Open Access

Fast and reliable methods for extracting functional connectivity in large populations

Yasser Roudi*¹, Joanna Tyrcha² and John Hertz^{1,3}

Address: ¹NORDITA, Roslagstullsbacken 23, 10691 Stockholm, Sweden, ²Dept of Mathematical Statistics, Stockholm University, 10691 Stockholm, Sweden and ³The Niels Bohr Institute, Copenhagen University, 2100 Copenhagen, Denmark

Email: Yasser Roudi* - yasser@nordita.org

* Corresponding author

from Eighteenth Annual Computational Neuroscience Meeting: CNS*2009 Berlin, Germany. 18–23 July 2009

Published: 13 July 2009

BMC Neuroscience 2009, **10**(Suppl 1):O9 doi:10.1186/1471-2202-10-S1-O9

This abstract is available from: <http://www.biomedcentral.com/1471-2202/10/S1/O9>

© 2009 Roudi et al; licensee BioMed Central Ltd.

The simplest model for describing multi-neuron spike statistics is the pairwise Ising model [1,2]. To start, one divides the spike trains into small time bins, and to each neuron i and each time bin t assigns a binary variables $s_i(t) = -1$ if neuron i has not emitted any spikes in that time bin and 1 if it has emitted one or more spikes. One then can construct an Ising model, $P(\mathbf{s}) = Z^{-1} \exp\{\mathbf{h}'\mathbf{s} + \mathbf{s}'\mathbf{J}\mathbf{s}\}$ for the spike patterns with the same means and pair correlations as the data, using Boltzmann learning, which is in principle exact. The elements J_{ij} of the matrix \mathbf{J} can be considered to be functional couplings. However, Boltzmann learning is prohibitively time-consuming for large networks. Here, we compare the results from five fast approximate methods for finding the couplings with those from Boltzmann learning.

We used data from a simulated network of spiking neurons operating in a balanced state of asynchronous firing with a mean rate of ~ 10 Hz for excitatory neurons. Employing a bin size of 10 ms, we performed Boltzmann learning to fit Ising models for populations of size N up to 200 excitatory neurons chosen randomly from the 800 in the simulated network. We studied the following methods: A) a naive mean-field approximation, for which \mathbf{J} is equal to the negative of the inverse covariance matrix, B) an independent-pair approximation, C) a low rate, small-population approximation (the low-rate limit of (B), which is valid generally in the limit of small Nrt , where r is the average rate (spikes/time bin) and t is the bin width [3], D) inversion of the TAP equations from spin-glass

theory [4] and E) a weak-correlation approximation proposed recently by Sessak and Monasson [5]. We quantified the quality of these approximations, as functions of N , by computing the RMS error and R^2 , treating the Boltzmann couplings as the true ones. We found, as shown in figure 1, that while all the approximations are good for small N , the TAP, Sessak-Monasson, and, in particular, their average outperform the others by a relatively large margin for N . Thus, these methods offer a useful tool for fast analysis of multineuron spike data.

References

1. Schneidman E, Berry MJ 2nd, Segev R, Bialek W: **Weak pairwise correlations imply strongly correlated network states in a neural population.** *Nature* 2006, **440**:1007-1012.
2. Shlens J, Field GD, Gauthier JL, Grivich MI, Petrusca D, Sher A, Litke AM, Chichilnisky EJ: **The structure of multi-neuron firing patterns in primate retina.** *J Neurosci* 2008, **28**:505-518.
3. Roudi Y, Nirenberg S, Latham P: **Pairwise maximum entropy models for large biological systems: when they can and when they can't work.** . arXiv:0811.0903v1 [q-bio.QM].
4. Tanaka T: **Mean-field theory of Boltzmann machine learning.** *Phys Rev E* 1998, **58**:2302-2310.
5. Sessak V, Monasson R: **Small-correlation expansions for the inverse Ising problem.** *J Phys A* 2009, **42**:055001.

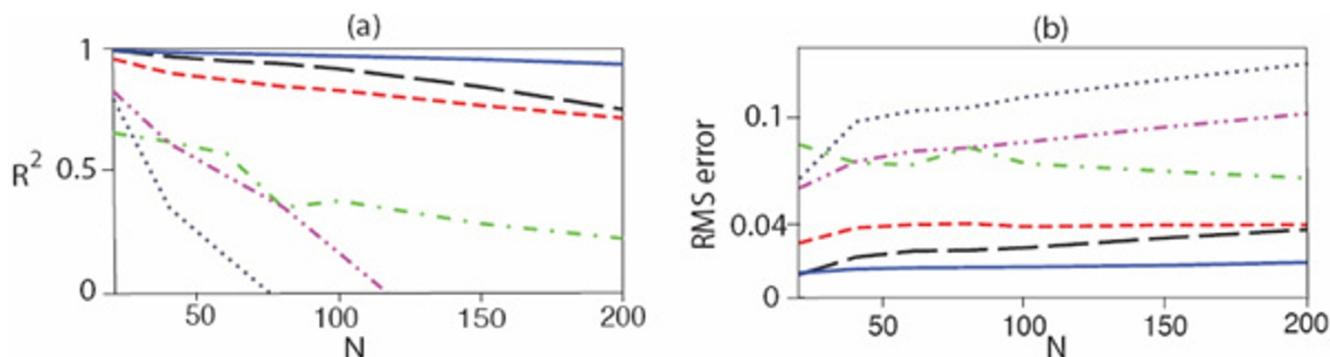


Figure 1
(a) R^2 and (b) RMS error for various approximate methods. Green (dashed dotted), naive mean-field; Purple (dashed double-dotted) low-rate, small N ; Gray (dotted) independent-pair; Red (dashed), TAP; Black (dashed), Sessak-Monasson; Blue, average of TAP and Sessak-Monasson.

Publish with **BioMed Central** and every scientist can read your work free of charge

"BioMed Central will be the most significant development for disseminating the results of biomedical research in our lifetime."

Sir Paul Nurse, Cancer Research UK

Your research papers will be:

- available free of charge to the entire biomedical community
- peer reviewed and published immediately upon acceptance
- cited in PubMed and archived on PubMed Central
- yours — you keep the copyright

Submit your manuscript here:
http://www.biomedcentral.com/info/publishing_adv.asp

