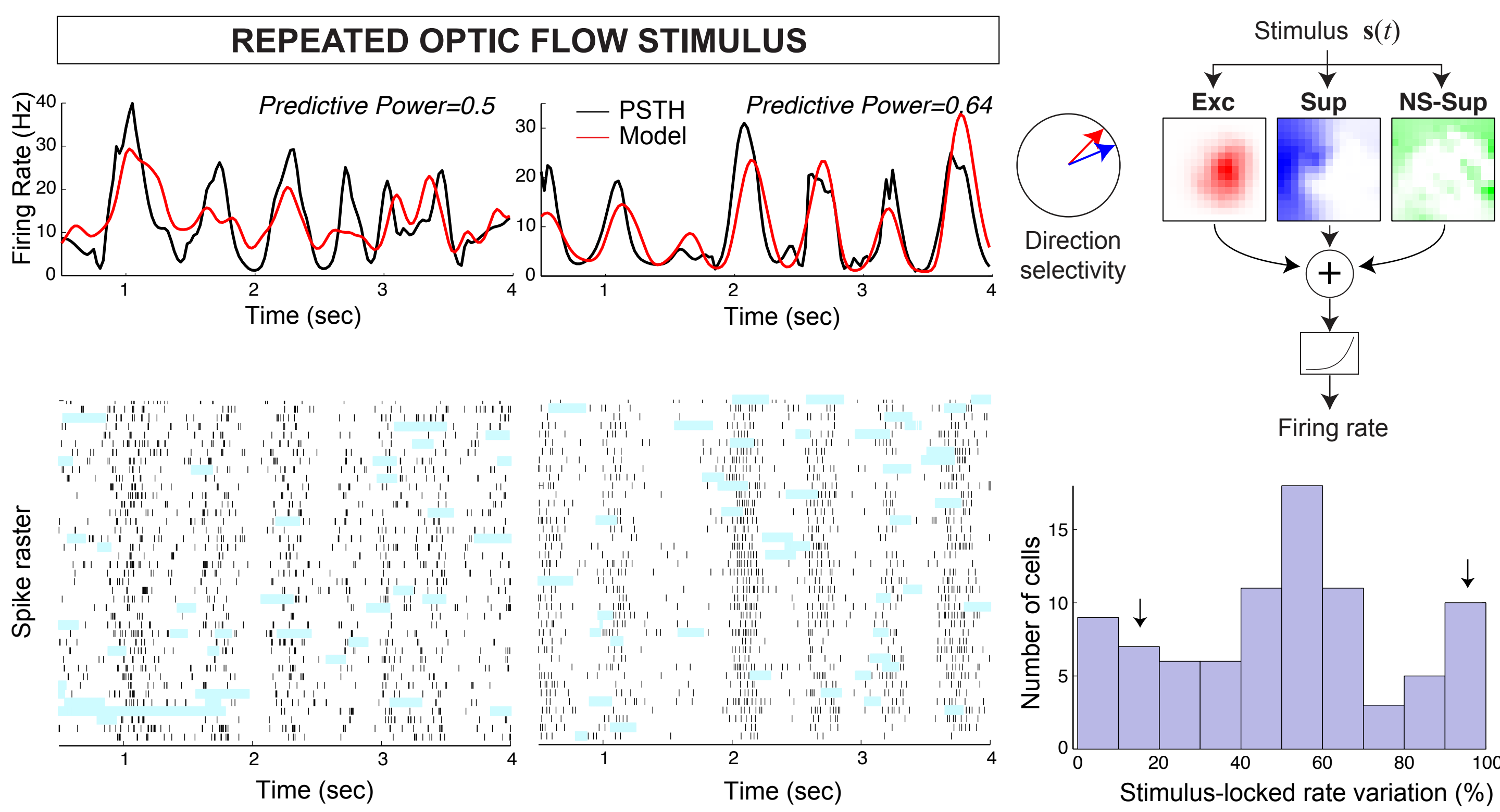


Motivation

While it is well understood that visual cortical neurons receive only a fraction of their inputs from upstream visual areas [1,2], a majority of studies of visual processing – both experimental and computational – necessarily focus on the role of feed-forward processing in vision. However, much recent work has described the possibility of inferring details of the network state from a common element of extracellular recordings, local field potentials [3-6]. In this work, we evaluate the impact of ongoing network activity on response variability of MT in both passive fixation conditions and during a motion discrimination task

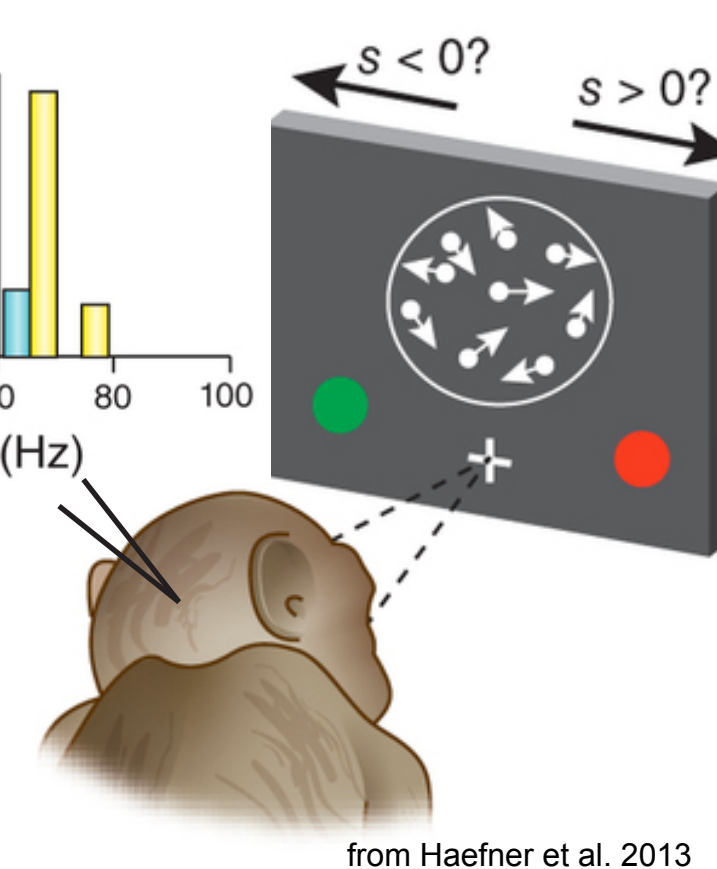
Response variability of MT neurons



Only ~50% of the firing rate variation is stimulus locked in a passive fixation task. Previous studies have shown that part of the response variability can be explained by ongoing network activity [4-5, 7].

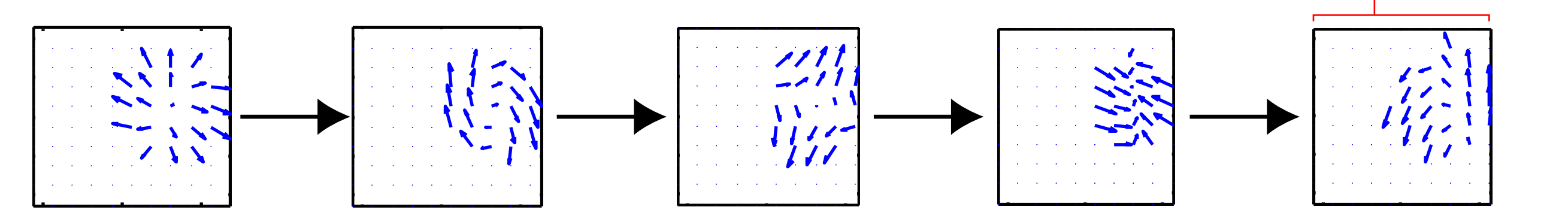
Response variability of MT is also related to perceptual decision in a motion discrimination task [8].

Can we explain spontaneous and decision-related cortical variability with ongoing network activity?



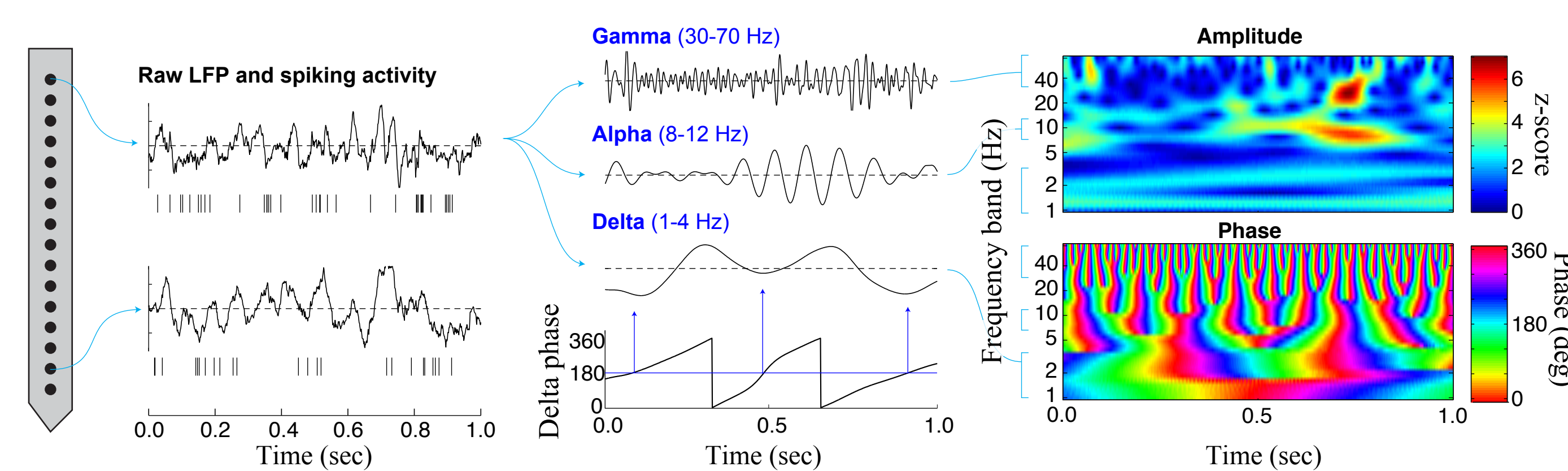
Experimental approach

Continuous Optic Flow Stimuli



Evolution of each optic flow component is independent low-passed Gaussian noise. A circular aperture is moving around slowly to explore the spatial profile of MT receptive fields.

Extracellular recordings were made in area MT in an awake macaque during a simple fixation task (e.g., see [10]). Eye position was monitored at 500Hz with an infrared eye tracker.



A Bayesian spike removal algorithm is applied to the raw LFP to remove artifactual correlations between spikes and LFPs ([9]).

Time frequency analysis to the LFPs was done using the continuous wavelet transform with a complex Morlet mother wavelet. Instantaneous phase and amplitude at each frequency were then given by the phase and modulus of the wavelet coefficients.

Modeling framework

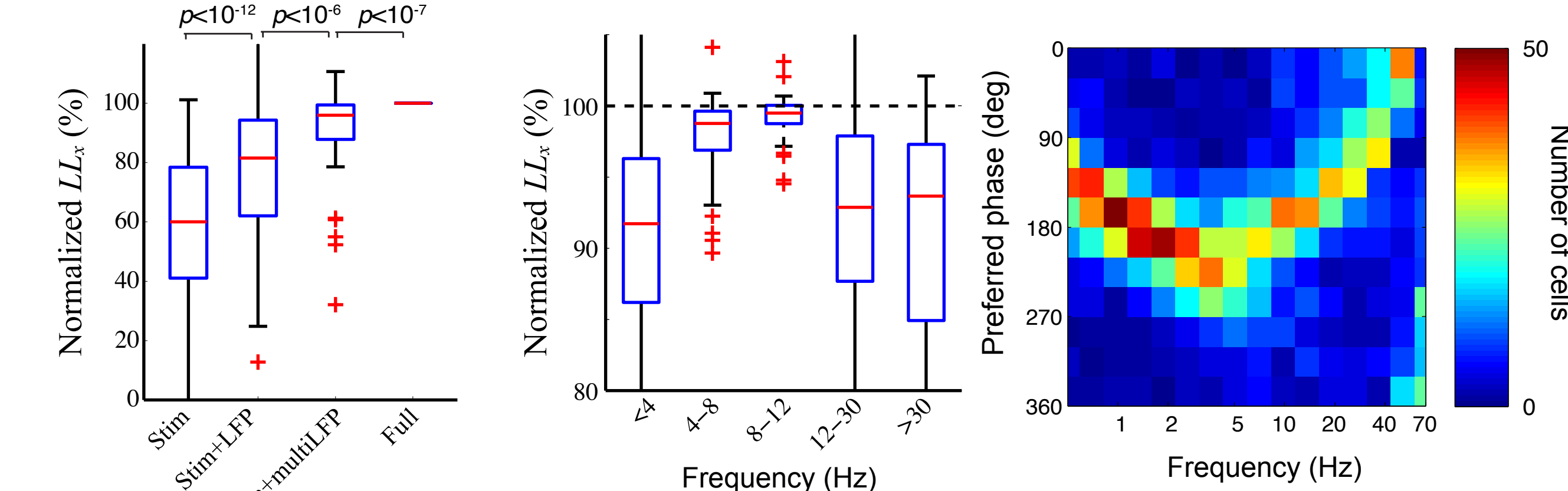
Neural response is modeled with a unified nonlinear framework using both stimulus (left, [10]) and network activity inferred from the LFP (middle) and MUA (right).

Contribution of LFP:

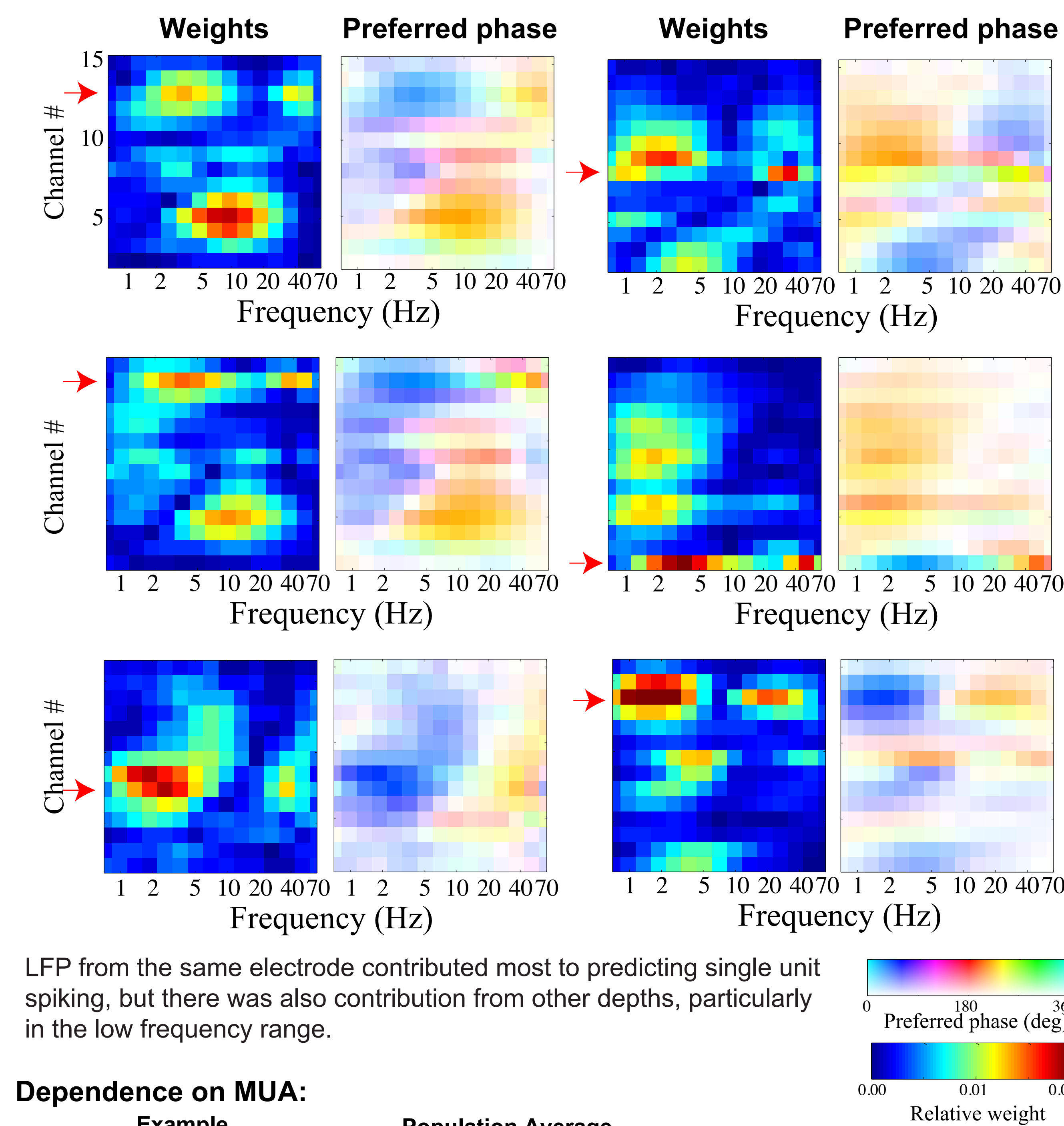
$$g_{LFP}(t) = \sum_j w_j \{A_j(f, t) \cos[\phi(f, t) - \phi_j^0]\} = \sum_j A_j(f, t) [\alpha_j \cos \phi(f, t) + \beta_j \sin \phi(f, t)] = k_{LFP} \cdot x_{LFP}(t)$$

Contribution of MUA: $g_{MUA}(t) = \sum_k k_{MUA}(t - \tau) = k_{MUA} \cdot x_{MUA}(t)$

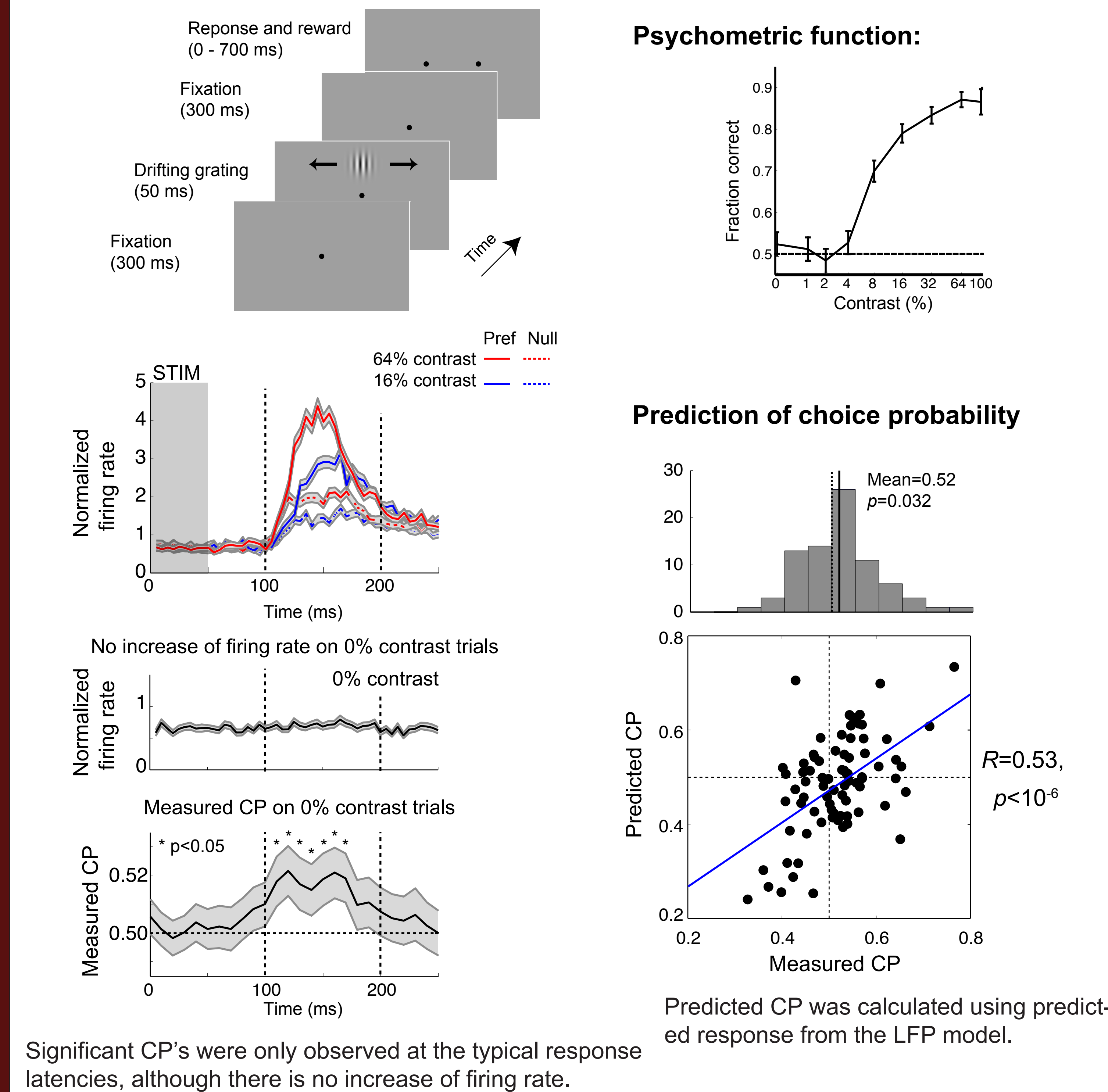
- Inclusion of LFP signals significantly improves model performance. Adding MUA provides little further improvement.
- Model performance depends on a wide range of LFP frequency bands (delta, beta, and gamma).
- Consistent phase preference across the population of cells.



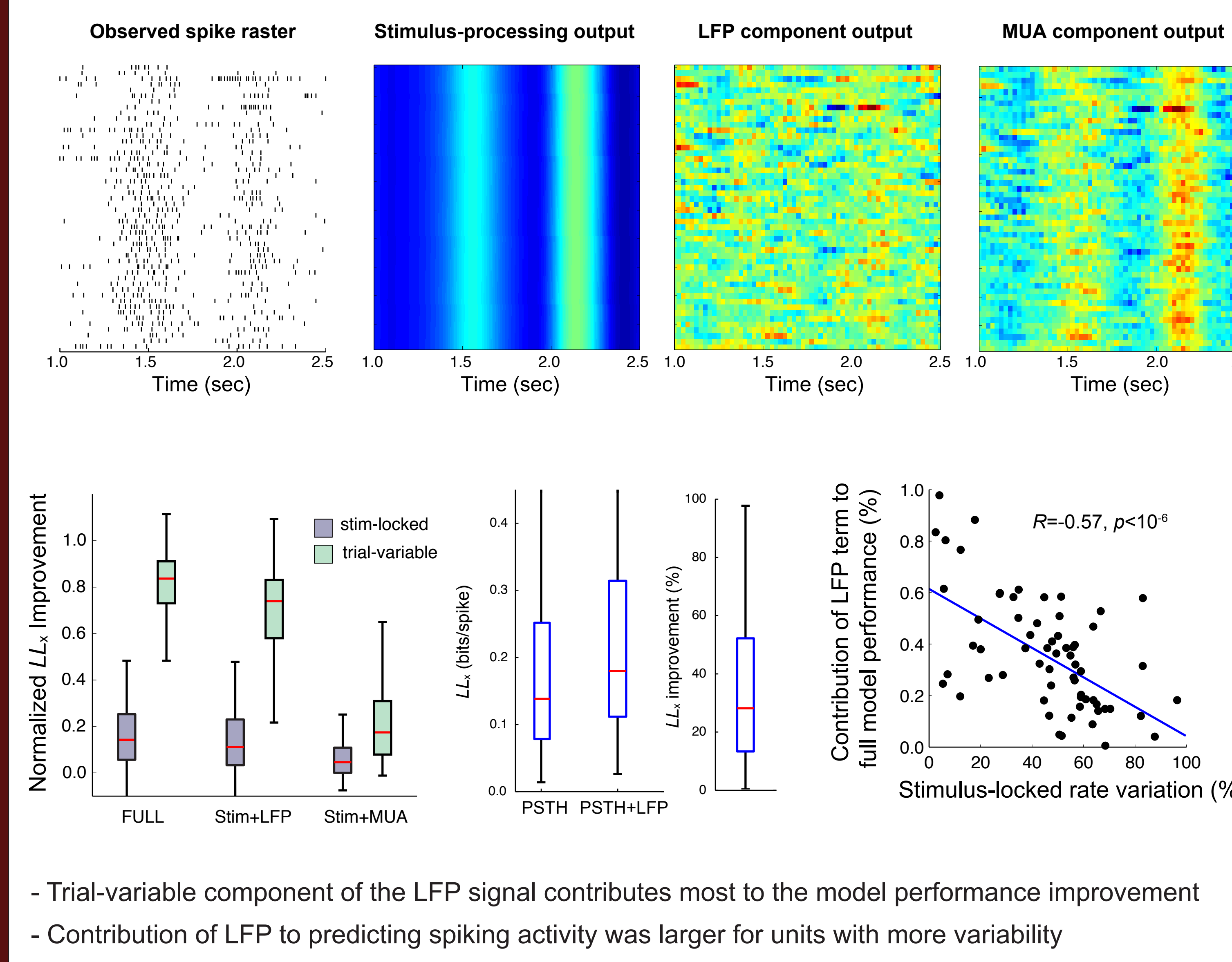
LFP modeling across cortical depth



Motion discrimination task

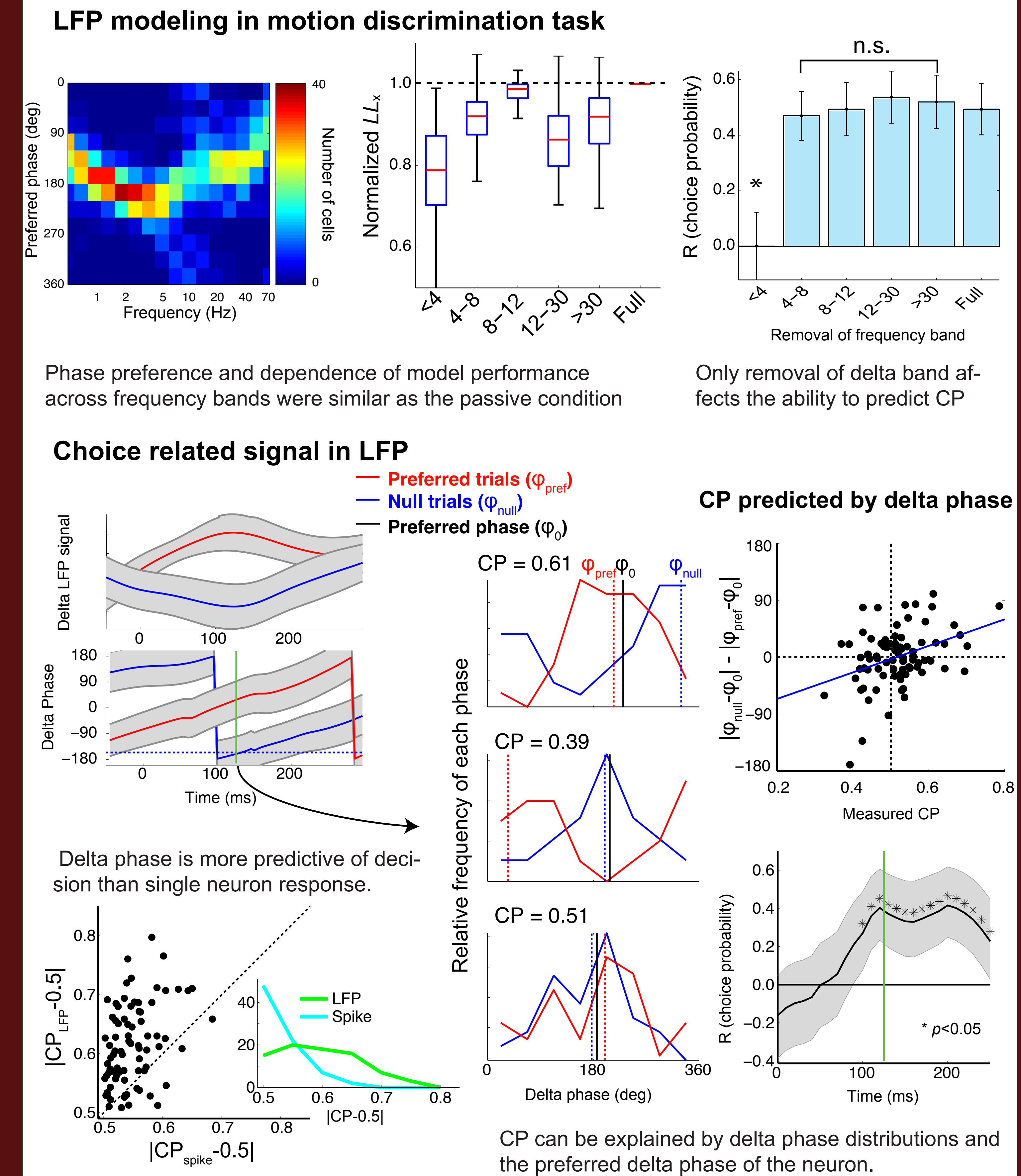


Network inputs predict variability



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Choice probability of MT neurons predicted by delta-band phase



Conclusions:

1. Only a fraction of cortical neuron response is stimulus driven. Part of the response variability can be explained by ongoing network activity.
2. Consistent dependence on LFP phase was observed across MT neurons and across task conditions.
3. Correlations between neuron activity and behavior (choice probability) can be predicted by low-frequency LFP components.
4. New modeling framework for incorporating network influences into stimulus processing.

References:

[1] Felleman DJ, Van Essen DC (1991) Cereb Cortex 1: 1-47.
[2] Salin PA, Bullier J (1995) Physiol Rev 75: 107-154.
[3] Buzsáki G (2004) Nat Neurosci 7: 446-451.
[4] Kelly RC, Smith MA, Kass RE, Lee TS (2010) Journal Comput Neurosci 29: 567-579.
[5] Haslinger R, Pipa G, Lima B, Singer W, Brown EN, et al. (2012) PLoS One 7(7): e39699
[6] Lakatos P, Shah AS, Knuth KH, Ulbert I, Karmos G, et al. (2005) J Neurophys 94: 1904-1911.
[7] Arieli, A., Sterkin, A., Grinvald, A. & Aertsen, A. Science 273, 1868-1871 (1996).
[8] Britten, K.H., Newsome, W.T., Shadlen, M.N., Celebri, S. & Movshon, J.A. Vis Neurosci 13, 87-100 (1996).
[9] Zanos, T.P., Mineault, P.J., and Pack, C.C. (2011) J Neurophys, 105, 474-486.
[10] Cui Y, Liu L, Khawaja FA, Pack CC, Butts DA (2013) J Neurosci 34(42): 16715-16728
[11] Bair W, Zohary E, Newsome, WT (2001) J Neurosci 21(5): 1676-97
[12] Haefner RM, Gerwinn S, Macke JH, Bethge M (2013) Nat Neurosci 16(2): 235-42