

Spatial extent of inputs to primate ganglion cells in natural viewing conditions

Nora Brackbill¹, Nishal Shah², Georges A. Goetz³, Alexandra Tikidji-Hamburyan³, Colleen Rhoades⁴, Alexander Sher⁵, Alan Litke⁵, E.J. Chichilnisky³

¹ Physics, Stanford University, Stanford, CA, United States.
² Electrical Engineering, Stanford University, Stanford, CA, United States.
³ Neurosurgery, Ophthalmology, and Hansen Experimental Physics Laboratory, Stanford University, Stanford, CA, United States.
⁴ Bioengineering, Stanford University, Stanford, CA, United States.
⁵ Santa Cruz Institute for Particle Physics, University of California, Santa Cruz, Santa Cruz, CA, United States.

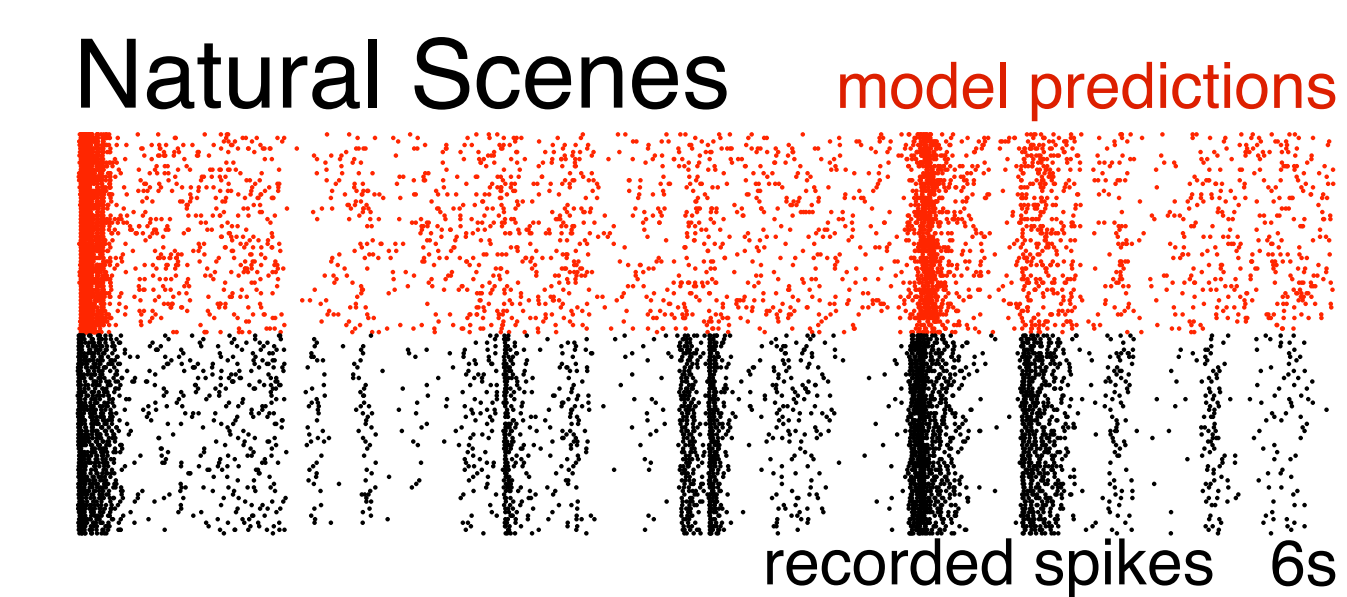
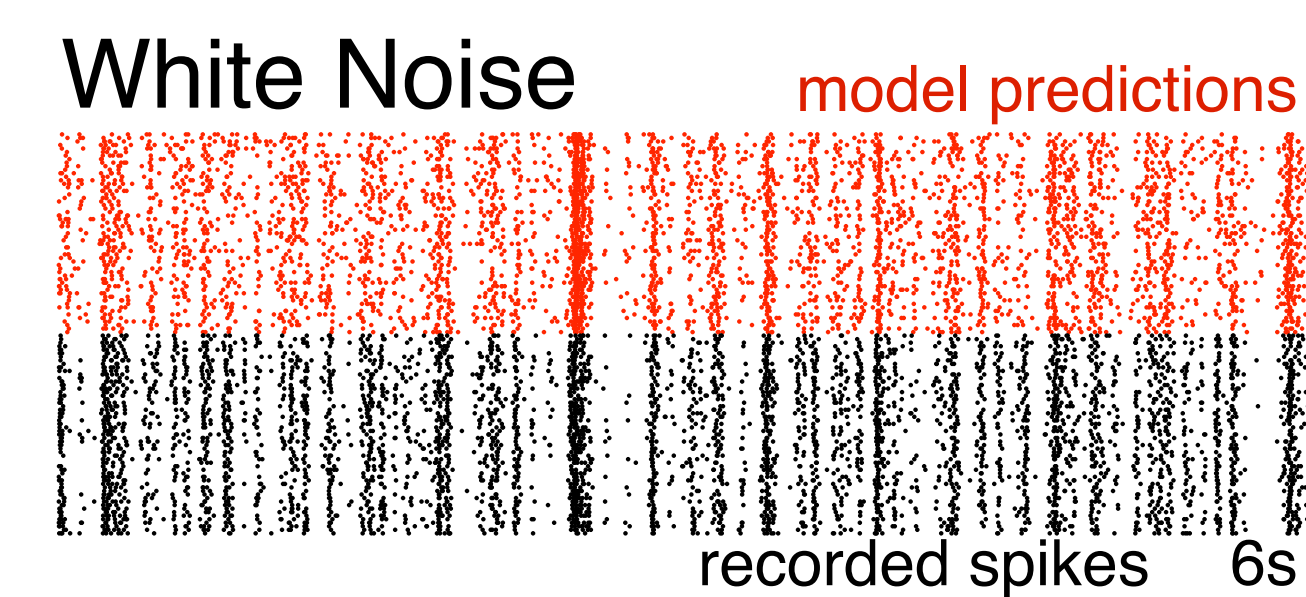
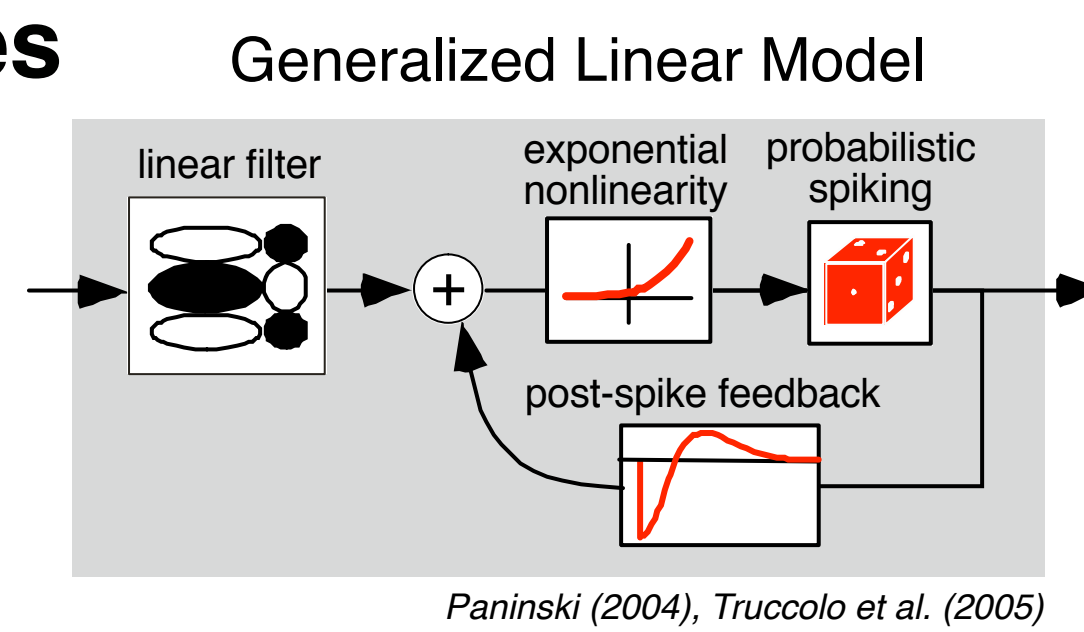
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Background

Modeling responses to natural scenes

Responses of primate retinal ganglion cells (RGCs) to natural scenes are poorly predicted by commonly used pseudo-linear models such as generalized linear models¹.



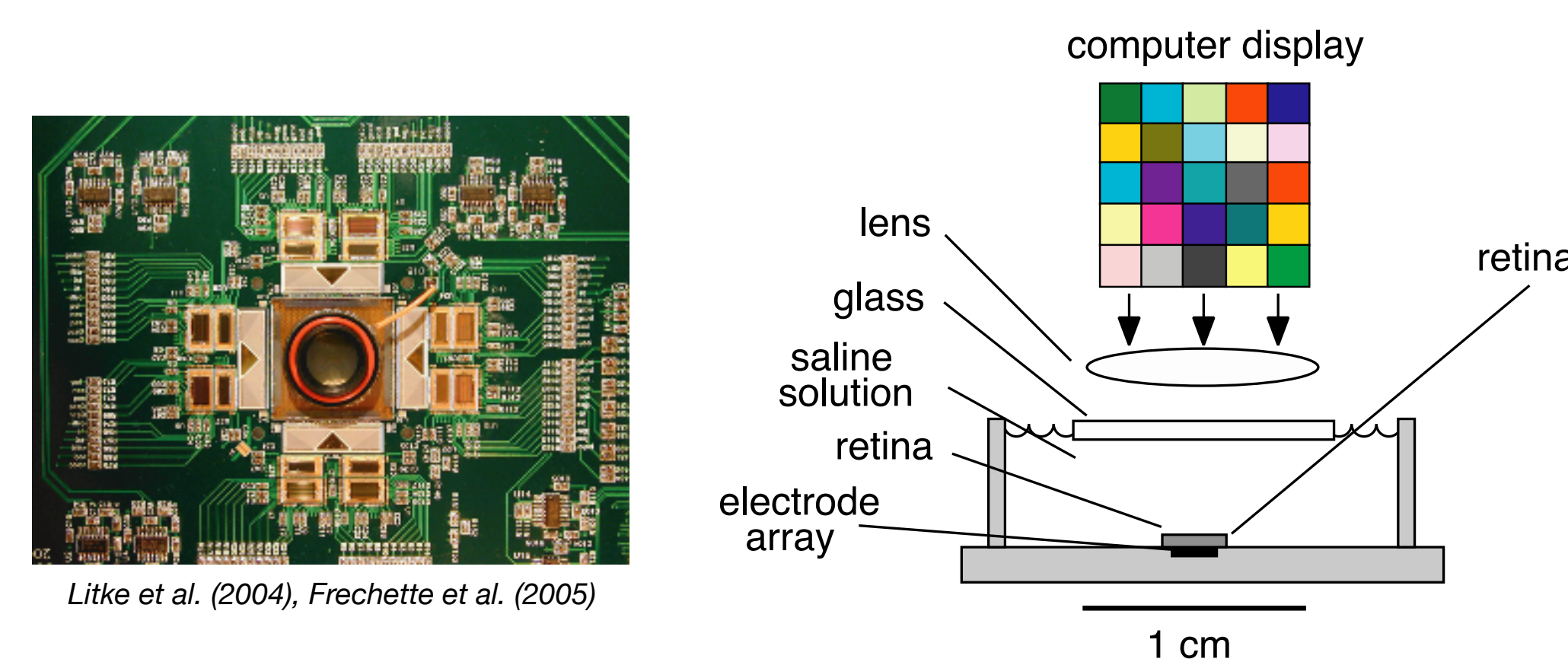
Extra-classical receptive field effects

Previous studies have shown that under certain conditions, RGCs in various species can receive peripheral input from outside their classical receptive fields²⁻⁶.

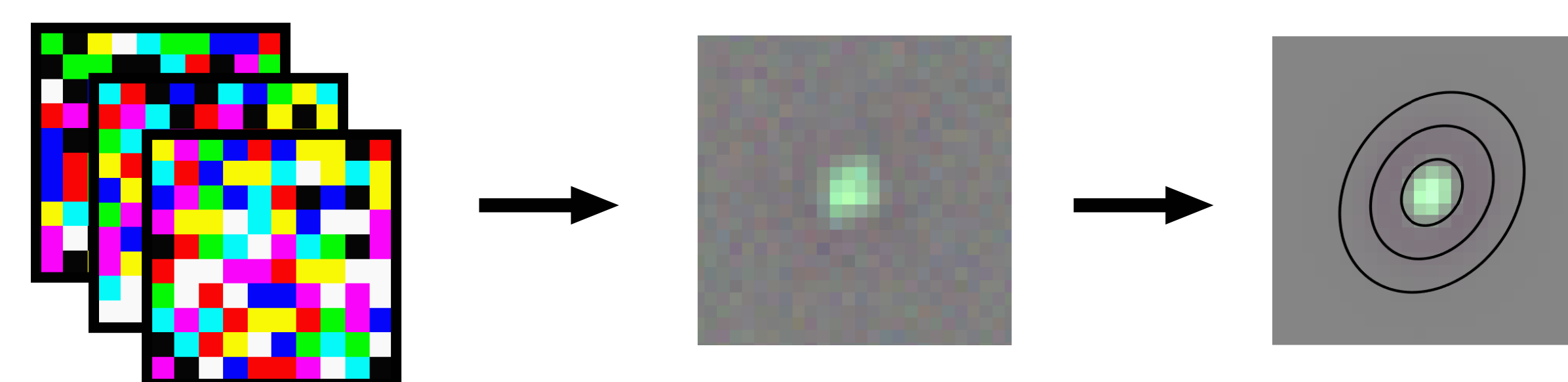
How do peripheral stimuli influence responses of primate RGCs in natural viewing conditions?

Methods

Large-scale multielectrode recordings were performed in peripheral macaque retina *ex vivo*.



ON and OFF parasol cells were identified, and their receptive fields were measured by reverse correlation with a white noise stimulus and fit with a Gaussian envelope with standard deviation σ , shown on right with contours of 2σ , 4σ , and 6σ .

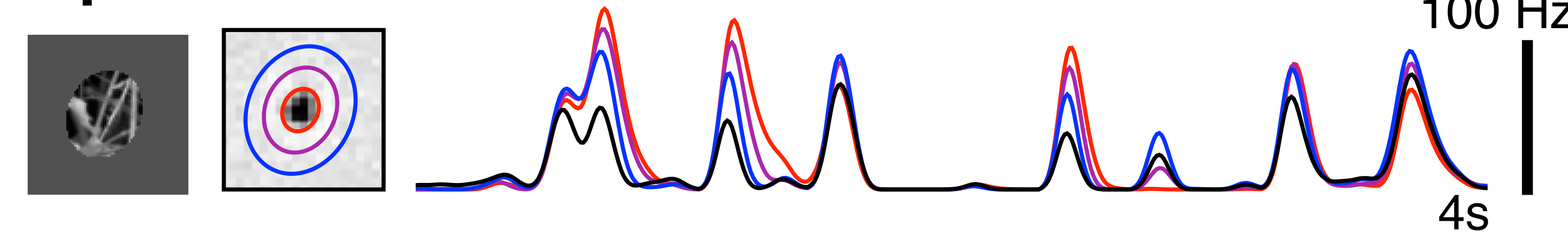


Natural scenes, consisting of images from the van Hateren database⁷ with fixational eye movements simulated by Brownian motion⁸, were presented in the three conditions.



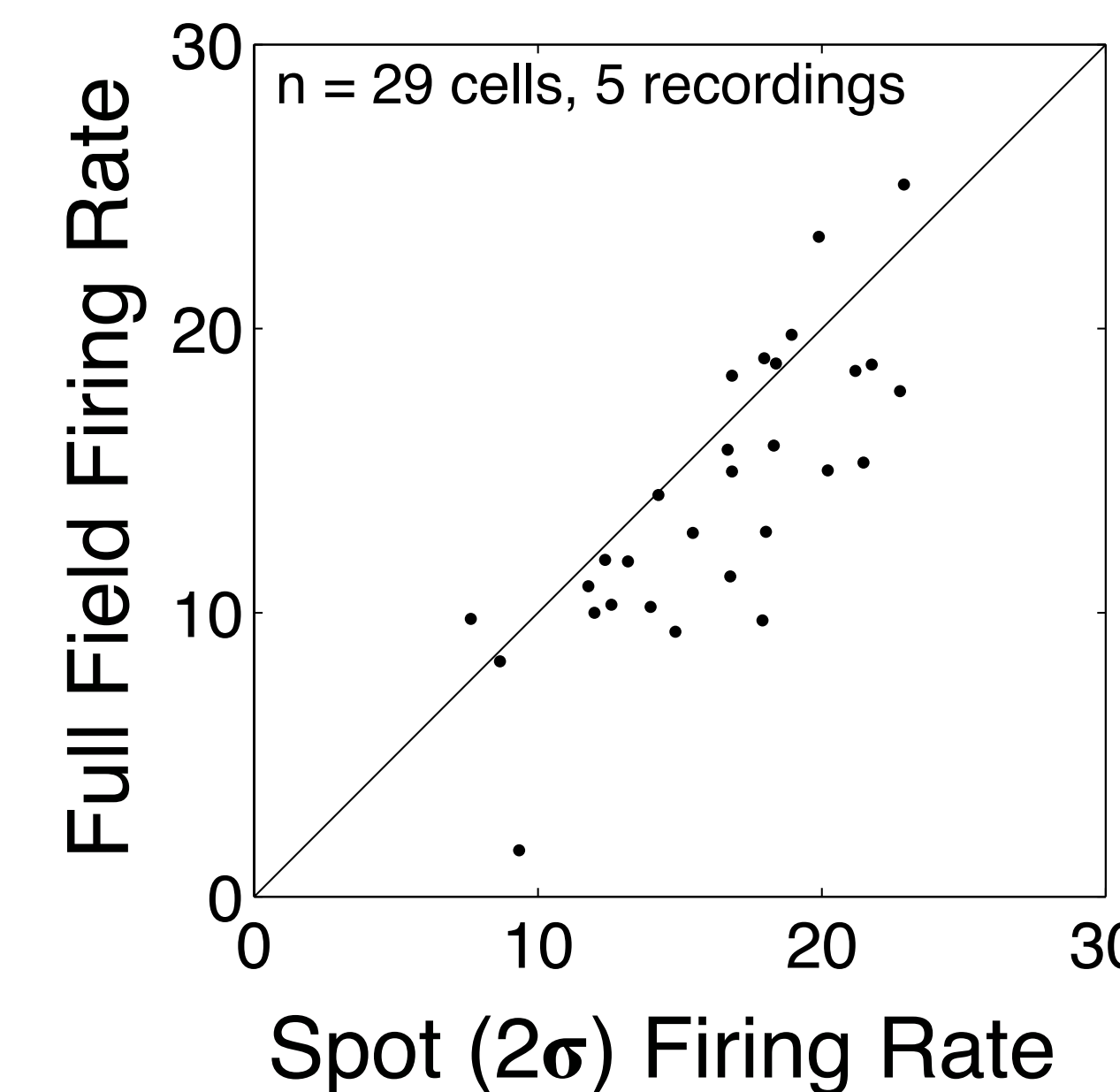
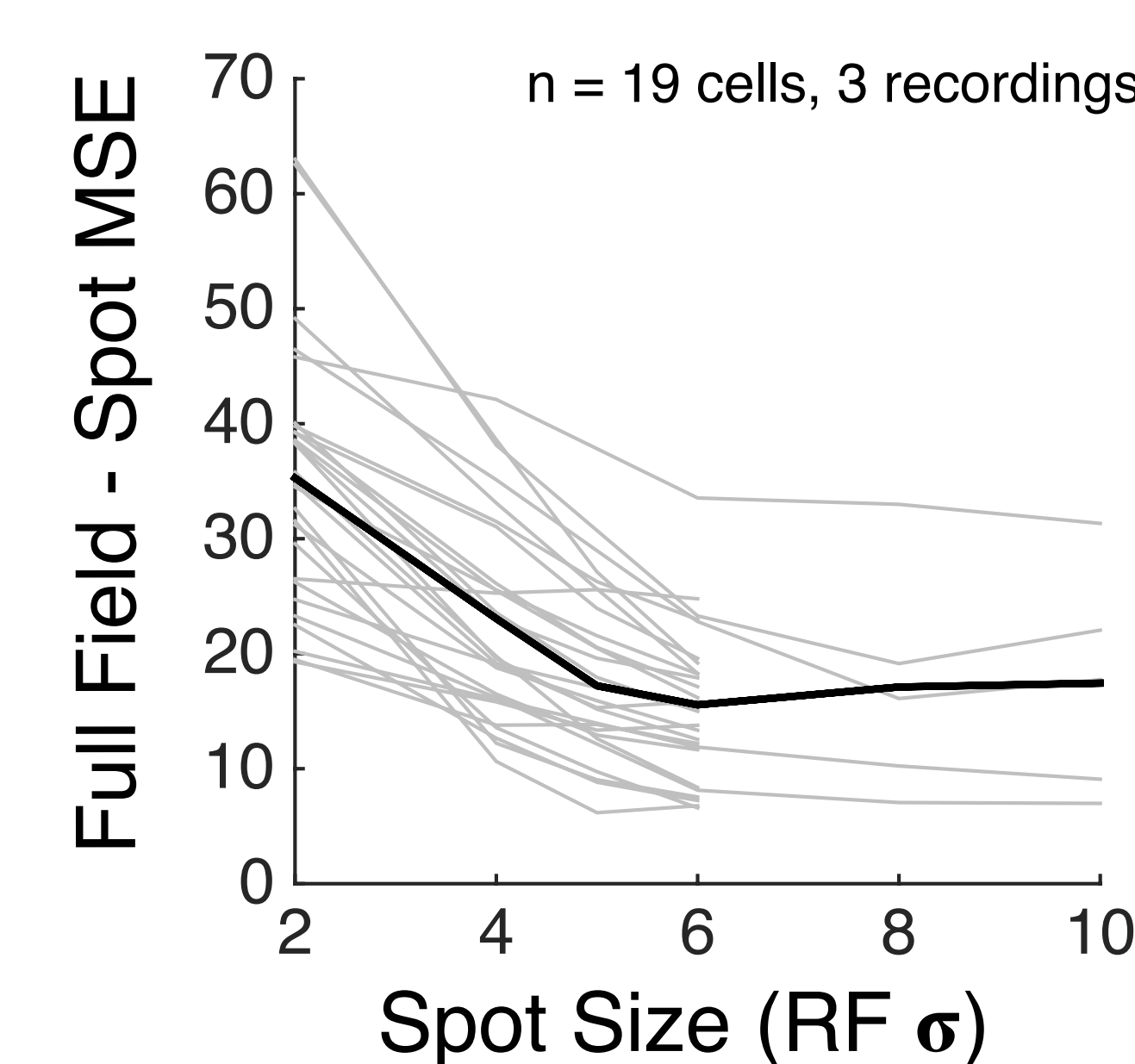
What effect does peripheral stimulation have on RGC responses under natural viewing conditions?

Spot

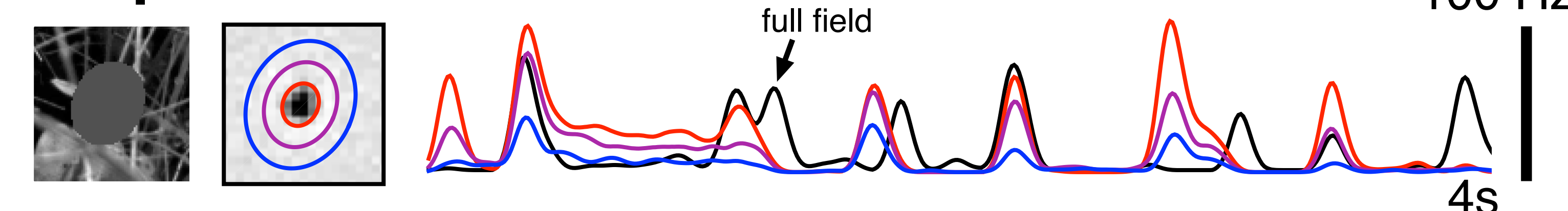


The response to the spot stimulus (above) had a time course **similar to the response to the full field** stimulus.

Consistent with previous findings, the periphery had a **mostly suppressive effect**.

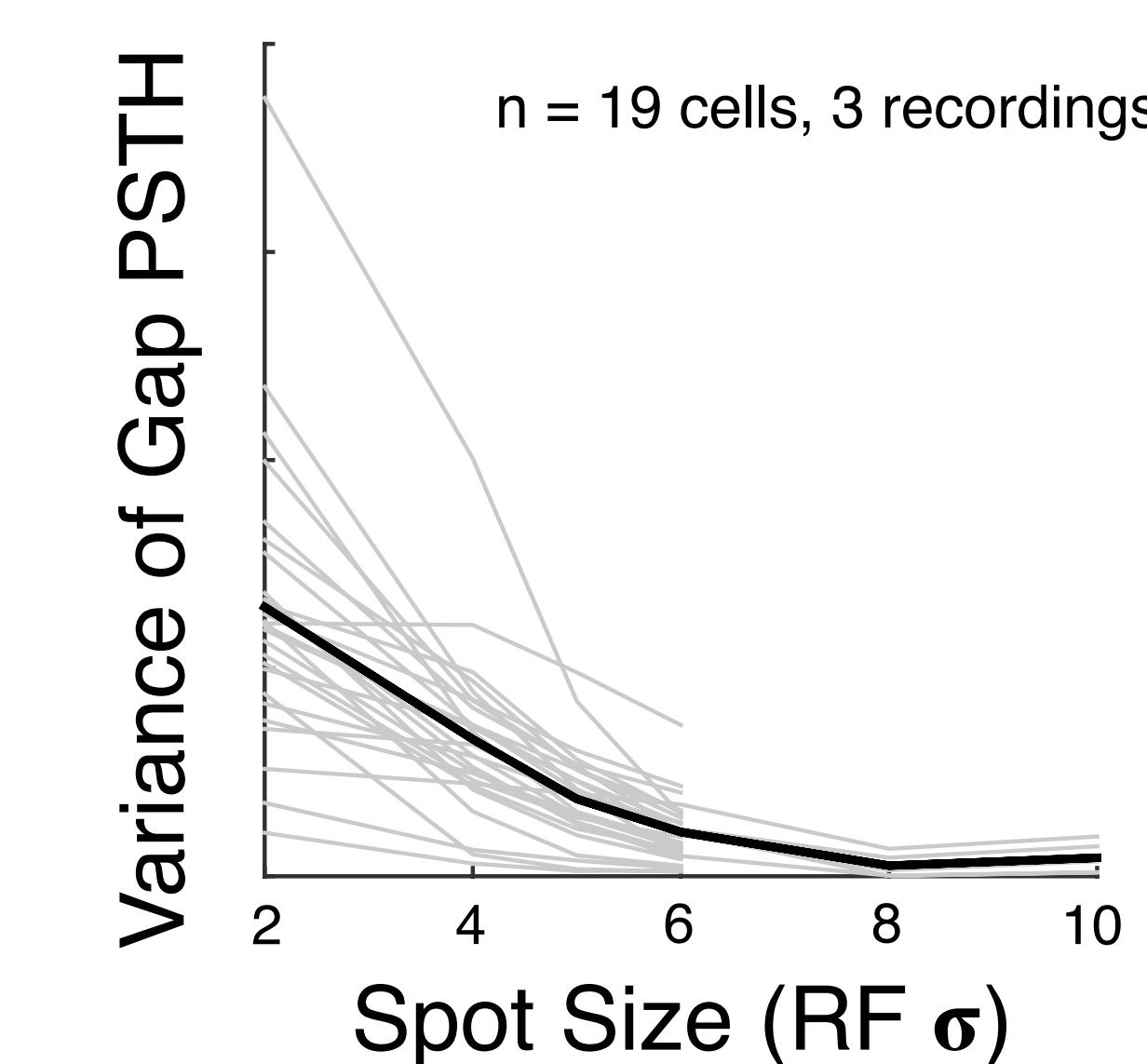


Gap



The response to the gap stimulus (above) had a time course **different than the response to the full field** stimulus.

In addition, the response to the gap stimulus did not disappear until greater than 6σ , **well outside the classical receptive field**.

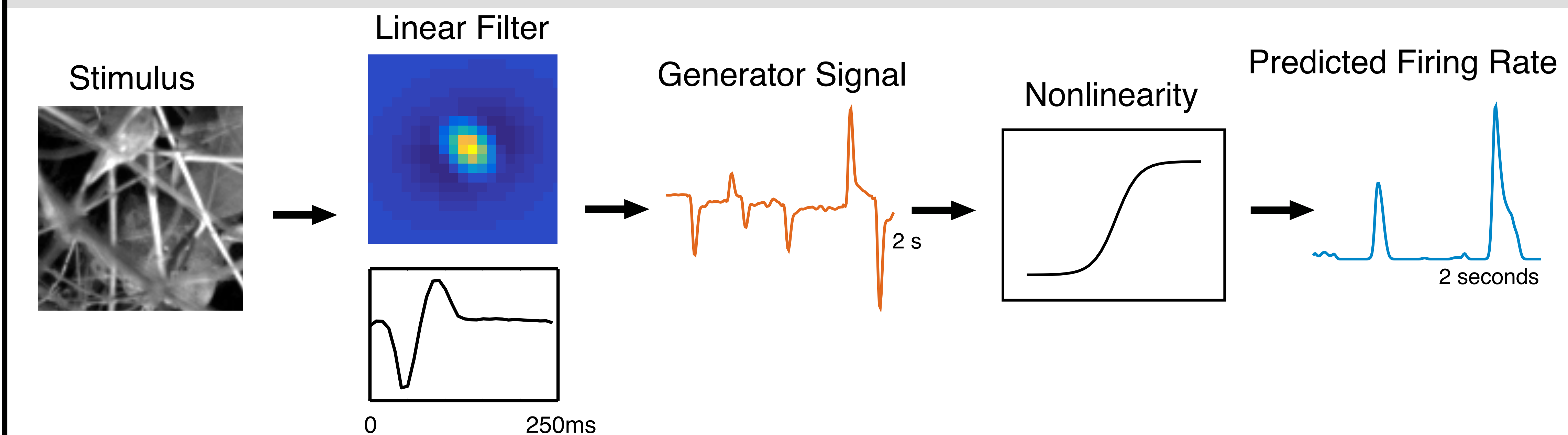


Peripheral stimulation far outside the classical RF had a significant effect during natural viewing conditions.

References

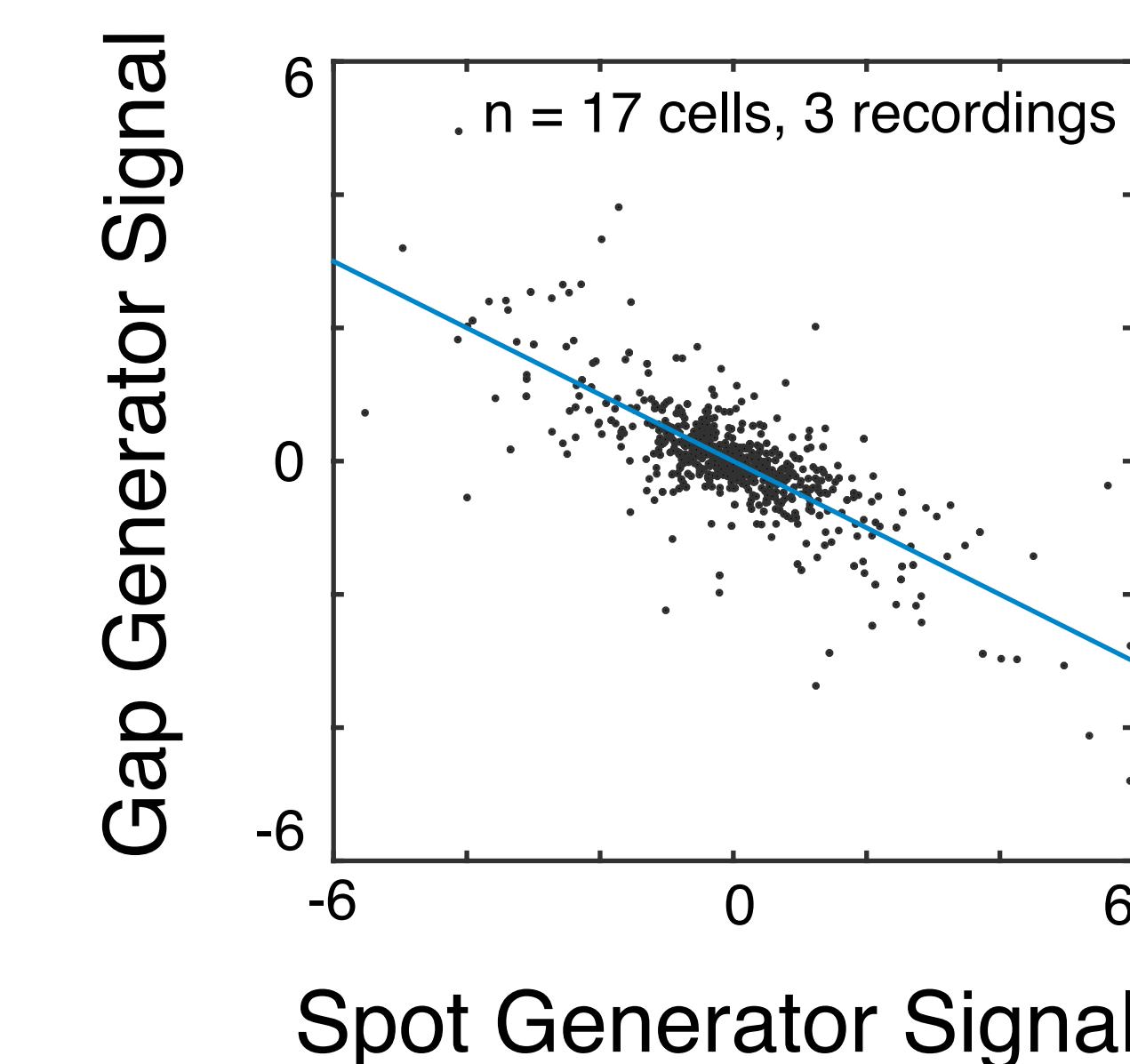
- [1] Alexander Heitman, Nora Brackbill, Martin Greschner, Alexander Sher, Alan M. Litke, and E.J. Chichilnisky. Testing pseudo-linear models of responses to natural scenes in primate retina. *bioRxiv*, 2016.
- [2] James T. McIlwain. Receptive fields of optic tract axons and lateral geniculate cells: peripheral extent and barbiturate sensitivity. *Journal of Neurophysiology*, 27(6):1154-1173, 1964.
- [3] H B Barlow, A M Derrington, L R Harris, and P Lennie. The effects of remote retinal stimulation on the responses of cat retinal ganglion cells. *The Journal of Physiology*, 289(1):177-194, 1977.
- [4] Christopher L. Passaglia, Daniel K. Freeman, and John B. Troy. Effects of remote stimulation on the modulated activity of cat retinal ganglion cells. *The Journal of Neuroscience*, 29(6):2407-2476, 02 2009.
- [5] Daisuke Takekoshi and Tim Gollisch. Nonlinear spatial integration in the receptive field surround of retinal ganglion cells. *The Journal of Neuroscience*, 34(22):7548-7561, 05 2014.
- [6] Kareem A. Zaghloul, Michael B. Manookin, Bart G. Borghuis, Kwabena Boahen, and Jonathan B. Demb. Functional circuitry for peripheral suppression in mammalian y-type retinal ganglion cells. *Journal of Neurophysiology*, 97(6):4327-4340, 06 2007.
- [7] J. H. van Hateren and A. van der Schaaf. Independent component filters of natural images compared with simple cells in primary visual cortex. *Proceedings: Biological Sciences*, 265(1394):359-366, Mar 1998.
- [8] Xutao Kuang, Martina Poletti, Jonathan D. Victor, and Michele Rucci. Temporal encoding of spatial information during active visual fixation. *Current Biology*, 22(6):510 - 514, 2012.
- [9] Liam Paninski. Maximum likelihood estimation of cascade point-process neural encoding models. *Network: Comput. Neural Syst.*, 15:243-262, 2004.
- [10] Wilson Truccolo, Uri T. Eden, Matthew R. Fellous, John P. Donoghue, Emery N. Brown, A Point Process Framework for Relating Neural Spiking Activity to Spiking History, Neural Ensemble, and Extrinsic Covariate Effects. *Journal of Neurophysiology*, 93(2):1074-1089, 2005.
- [11] Alan Litke et al. What Does the Eye Tell the Brain?: Development of a System for the Large-Scale Recording of Retinal Output Activity. *IEEE Transactions on Nuclear Science*, 51(4):1434-1440, 2004.
- [12] E. S. Fréchet, A. Sher, M. I. Grivich, D. Petrusca, A. M. Litke, E. J. Chichilnisky. Fidelity of the Ensemble Code for Visual Motion in Primate Retina. *Journal of Neurophysiology*, 94(1):119-135, 2005.

Can we understand these responses with linear-nonlinear models?

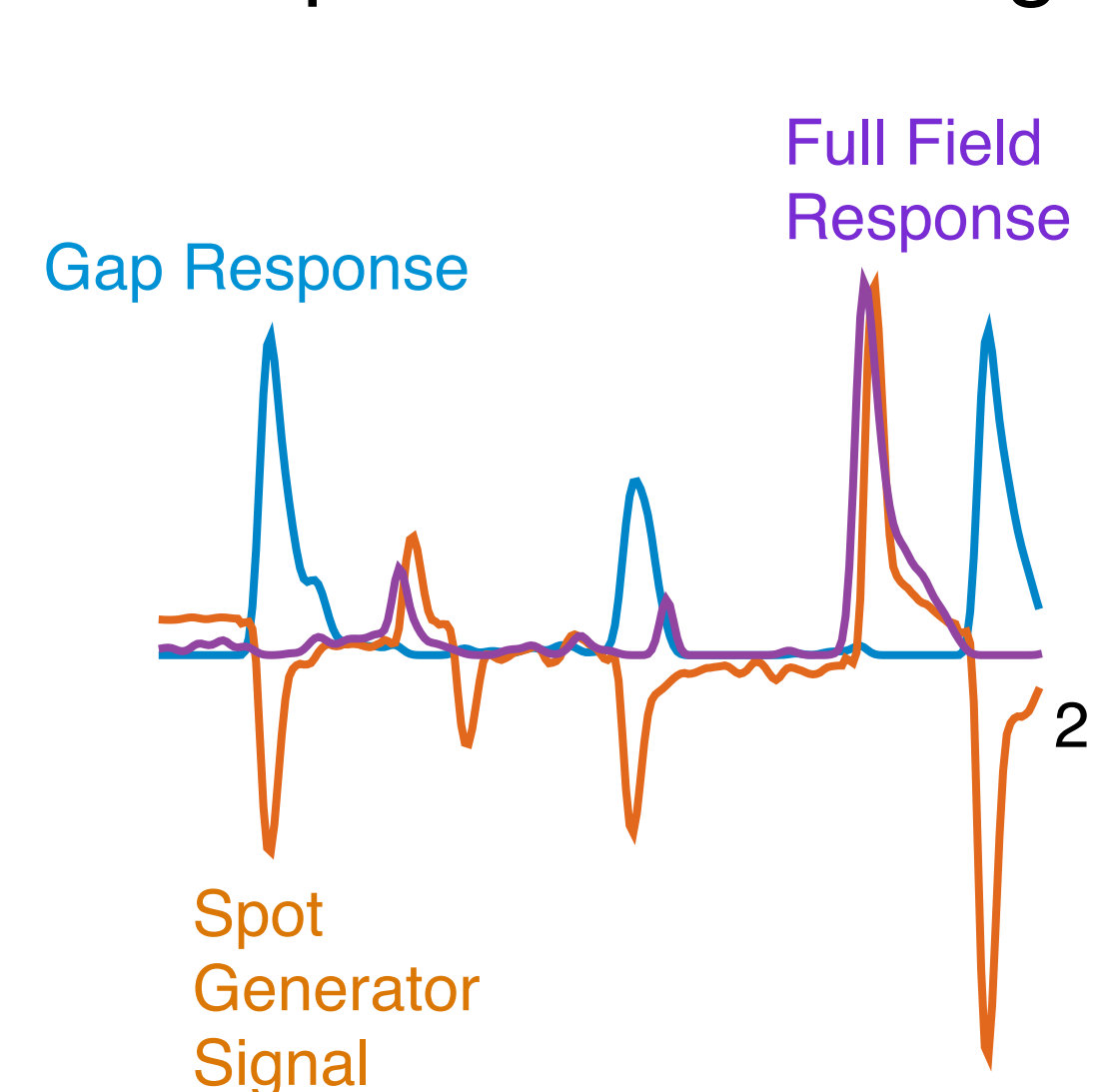


The observed effects can be explained by the fact that spatial correlations in natural scenes introduce anticorrelation in the gap and spot generator signals.

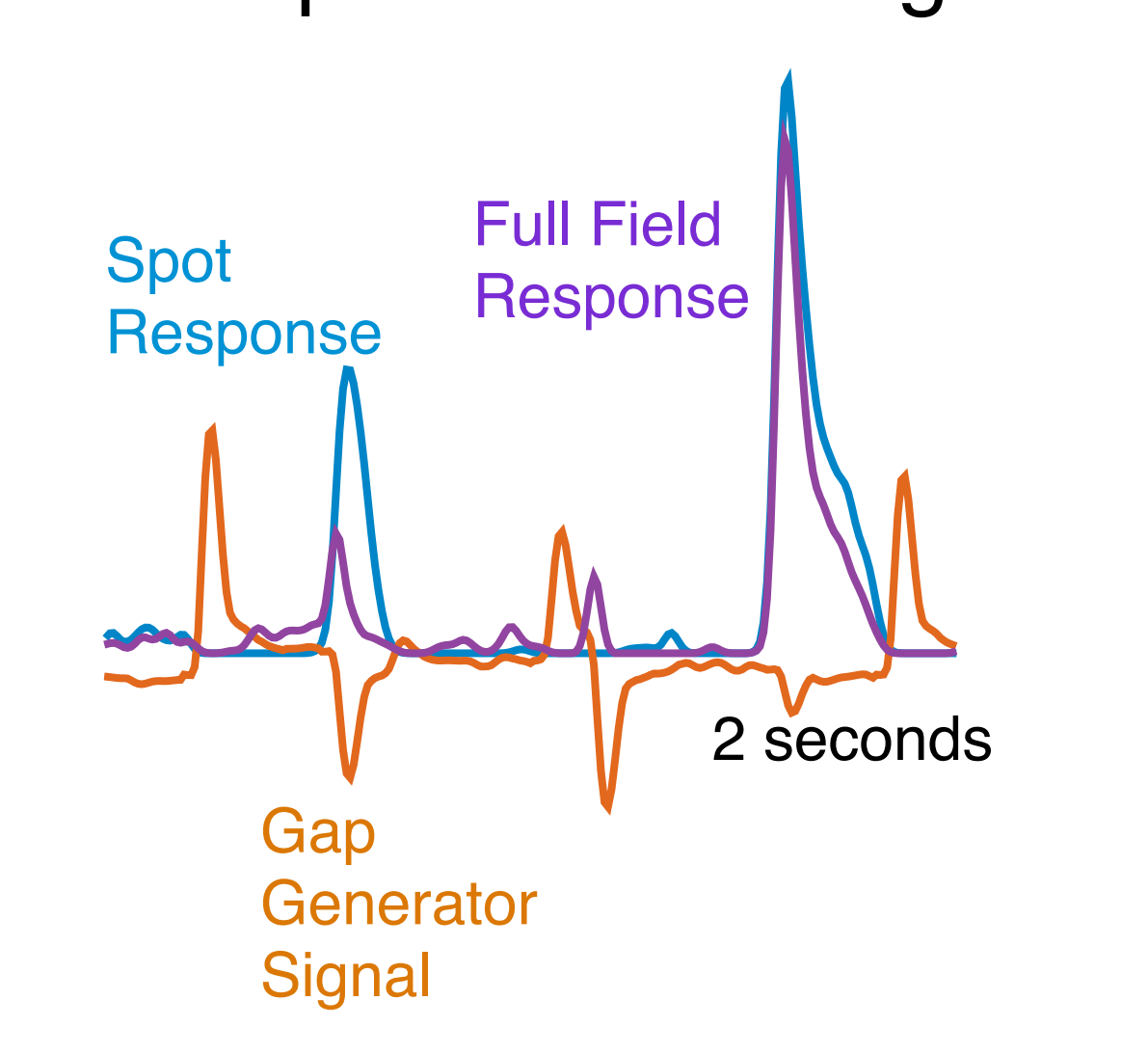
Since the spot generator signal tends to be stronger, gap response events tend to not be present during full field stimulation, while spot events tend to be reduced.



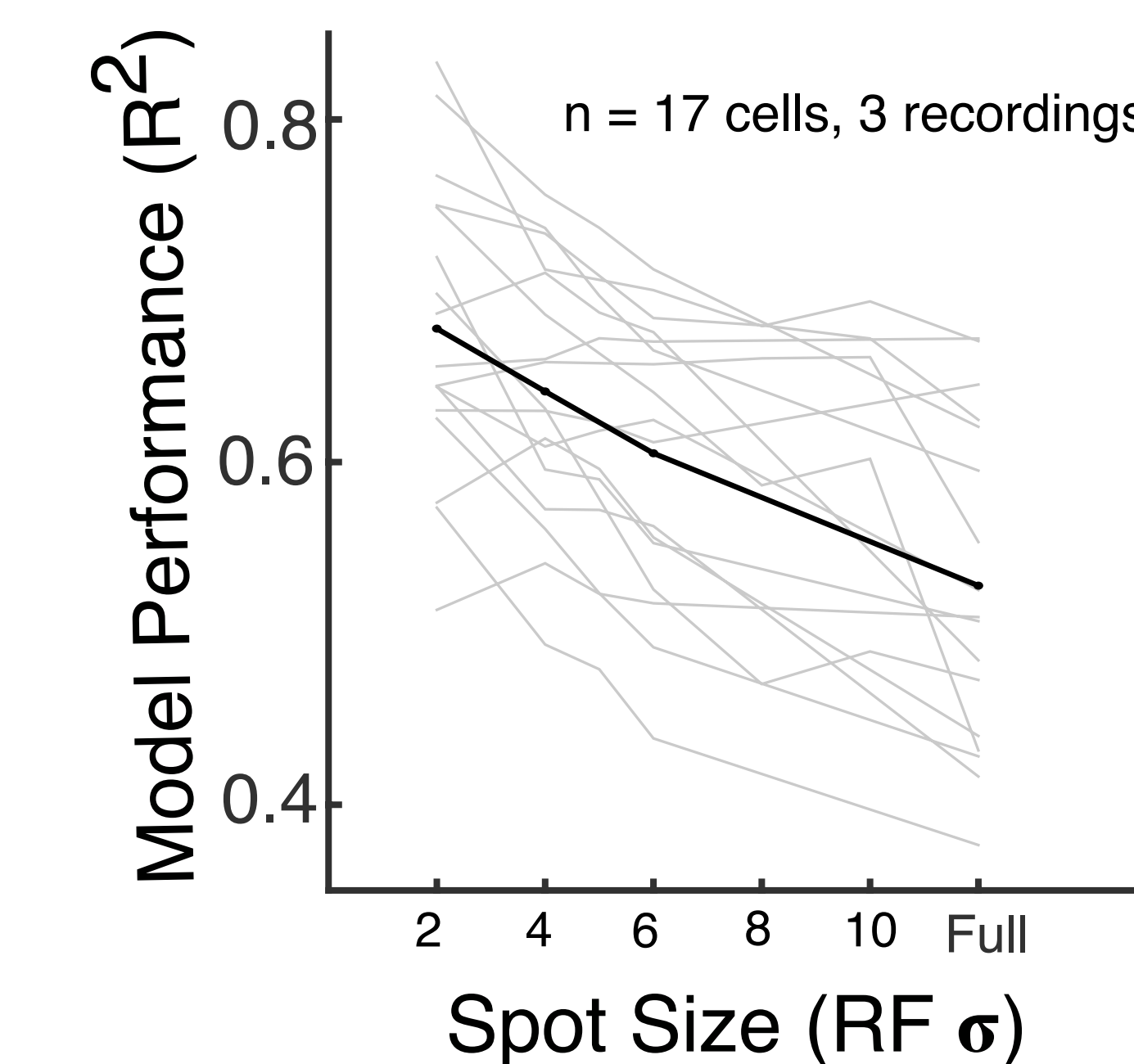
Gap Response vs Spot Generator Signal



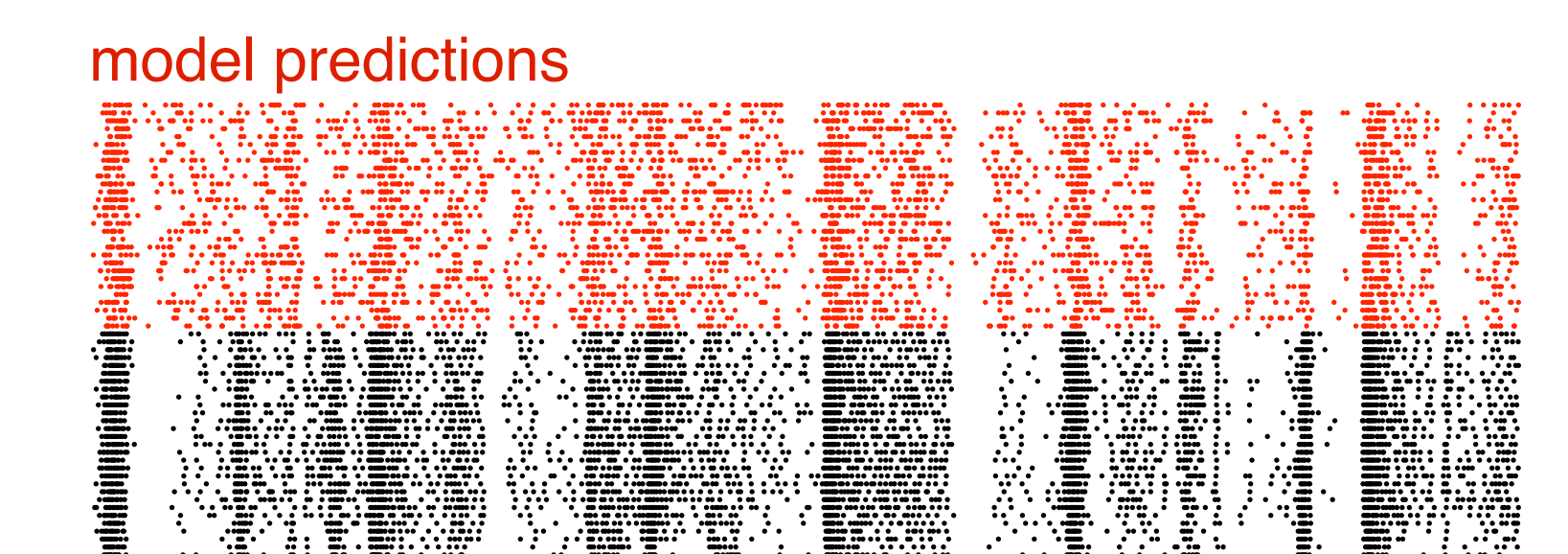
Spot Response vs Gap Generator Signal



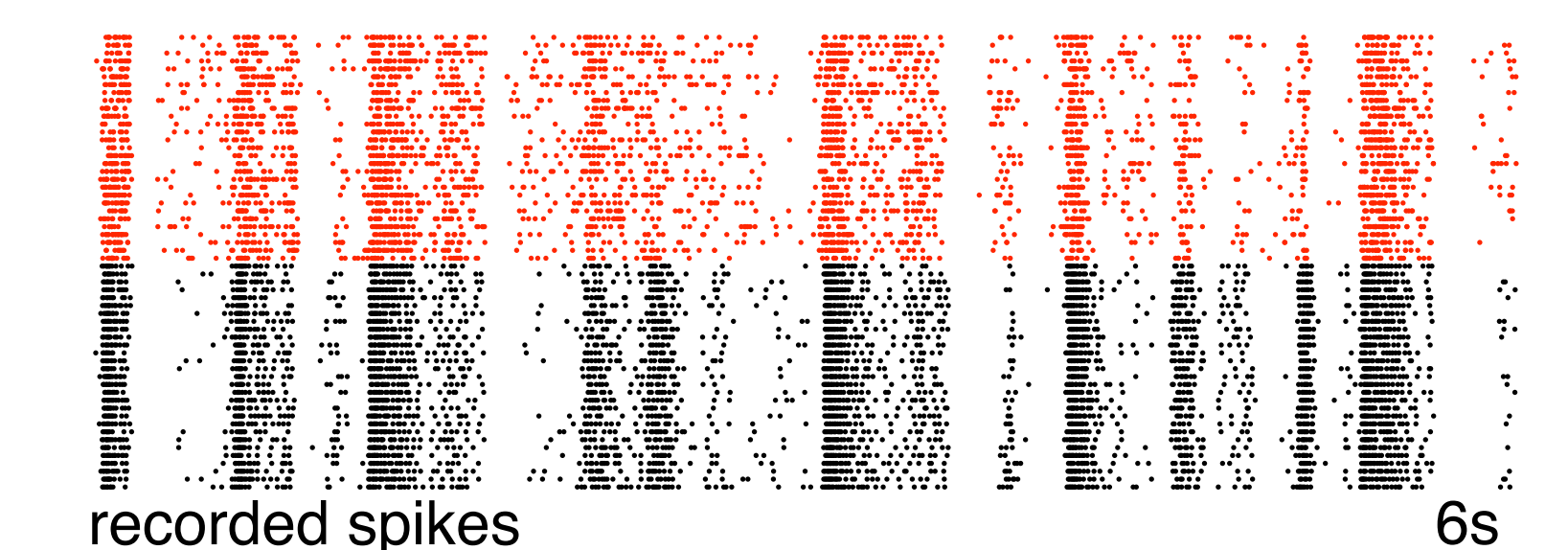
Surprisingly, this **model was more accurate for small spot sizes**. The model captured responses to stimuli restricted to the receptive field center more accurately than responses to stimuli in the surround and periphery.



Full Field



Spot (2 sigma)



While the observations can be qualitatively explained by linear summation of the spot and gap generator signals, a full linear-nonlinear model did not accurately capture responses outside the RF center.

Future work will focus on developing models that more effectively incorporate the surround and peripheral input.

Acknowledgements

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