# Stimuli for Fast Mapping of Feature Tuning in Mouse Visual Cortex

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# INTRODUCTION

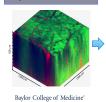
Parametric stimuli quickly characterize the basic visual response properties of sensory neurons. An effective stimulus evokes strong,

We present two new parametric visual stimuli, Monet and Trippy, designed to simultaneously and rapidly characterize linear receptive fields, motion tuning, and, for Trippy, the spatial and temporal frequency tunings. These stimuli are used in the MICrONS project for high-throughput functional circuit description.

# **MICrONS Study**

# Functional Description of All Neurons in 1 mm<sup>3</sup> 2-photon Ca<sup>2+</sup> Imaging 3-photon Ca<sup>2+</sup> Imaging Visual Stimulation: natural and









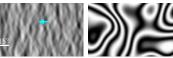
# Visual Stimuli

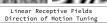
### "Monet"

Smoothened Gaussian noise with with coherent orientation motion of varying directions

#### "Natural" "Trippy"

Irregular and fluid ripples movies of natural scenes wriggling randomly across the synthetic movies of simulated visual field - synthesized as the cosine of a smooth Gaussian

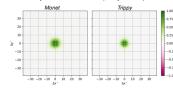




tuning spatial + temporal Localization of directional response

Modeling Ethological relevance

#### Spatial Autocorrelation (at any location)



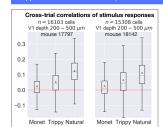
Despite the prominent global visual features present at any moment, the spatial and temporal autocorrelations of both Monet and Trippy are, by design, similar and are space-invariant, circularly symmetric (in visuotopic space), unbiased, and compact.

These properties enable computation of linear receptive fields by simple spiketriggered averaging

# **RESULTS**

# **Stimulus Drive**

strong reliable responses from neurons. We measure the stimulus drive as the correlations between the responses to two presentations of identical 15-s movie snippets shuffled across the experiment session.

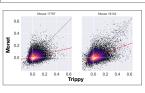


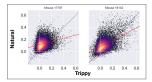
Natural movies produced strongest drive with mean cross-trial response correlations of 0.11 -0.13 in L2-4 of V1 when computed at 4 Hz.

Trippy with mean cross-correlations 0.05 - 0.07 significantly outperformed Monet (~0.03). p=0 in

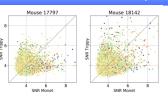
The increase from Monet to Trippy is mostly due

The increase from Trippy to Natural is mostly due to increased breadth of activation.



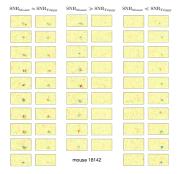


# Linear receptive fields



Trippy and Monet have similar autocorrelation functions: therefore, an ideal linear system would produce similar spike-triggered average receptive fields.

However, Trippy and Monet elicited responses from distinct subpopulations of cells that differed in their functional



Cells responsive to both types of stimuli produce similar receptive fields from both

Cells responsive to Monet but not to Trippy tend to be oriented and have multiple

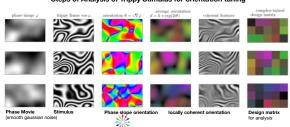
Cells selective to Trippy but not to Monet predominantly comprise a single negative punctate subfield (circular with a weak annular or crescent surround)

# **Directional Tuning**

The Monet stimulus comprises periods of coherent motion and orientation. Its analysis is not much different from that of conventional drifting grating stimuli.

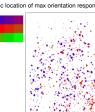
The Trippy stimulus is derived from an underlying phase movie, which provides a direct representation of local direction of motion and orientation, spatial and temporal frequency. This representation allows direct model fitting to these properties with locality information.

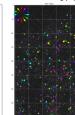
#### Steps of Analysis of Trippy Stimulus for orientation tuning

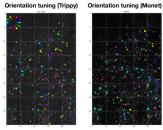


# **Pixelwise Orientation Map**

Visuotopic location of max orientation response







Monet and Trippy elicit responses from an overlapping population of cells. When cells respond to both, they

# **CONCLUSIONS AND FUTURE WORK**

Two synthetic stimuli with similar spectral properties but different global structures revealed the robustness of orientation tuning and linear receptive fields of subgroups of cortical neurons. But the comparison also differentiated subpopulations of cells with distinct functional properties in a consistent pattern. In ongoing work, we will relate structural and connectivity information reconstructed from electron microscopy to elucidate the anatomical regularities underlying these functional distinctions.

# **ACKNOWLEDGMENTS**

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