

Local and long-distance connectomics of the visual cortex

Clay Reid

Allen Institute for Brain Science, USA

Over the past decade, new tools have emerged for studying the structure and function of cortical circuits, an approach that has been given the unfortunate but now unavoidable name of functional connectomics. Our group has been using functional connectomics approaches that rely on two-photon calcium imaging, to assay the function of visual cortical neurons, followed by correlated anatomical studies. The first approach—serial section electron microscopy (Bock et al., 2011, Lee et al., 2016)—allows the tracing of synaptic connections between functionally imaged neurons. So far it has been possible to trace networks comprising ~ 100 neurons and ~ 1000 connections. Recent advances in imaging and automated segmentation should soon allow the study of networks that are several orders of magnitude larger.

Transsynaptic input mapping with G-deleted rabies (Wickersham et al., 2007) is another technique that can be combined with cellular imaging to study structure-function relationships. Recent technical advances in this technique allow the routine functional imaging of hundreds of inputs to a single postsynaptic neuron. While this many-to-one technique gives a more limited view of network structure than EM, it permits the study of long-distance feedforward and feedback connectivity between cortical areas. I will examine the near- and long-term prospects for functional connectomics and argue that long-standing questions of feedforward, feedback and recurrent connectivity in cortical circuits may be settled in the coming years.