

# Analysis of Neuronal Morphology in X-ray Synchrotron Images of *Drosophila Melanogaster* Brains

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Connectome, or the map of neural connections, provides crucial information for understanding operation principles of the brain. Although the fluorescent confocal microscopy and the electronic microscopy have been extensively used in large-scale projects on connectome mapping, we still lack a method that images neural circuits in a high-throughput fashion at the cellular resolution, and that generates samples from a large number of individuals efficiently. Without the availability of a large sample size, our understanding of the circuitry may become restricted. A novel imaging technique, Accelerated X-ray Observation of Neurons (AXON), was developed based on the synchrotron X-ray imaging and the Golgi stain. Using such a technique, we can acquire brain images rapidly with a fine resolution close to 0.5  $\mu\text{m}$ . In the present study, we develop imaging analysis tools for AXON and apply them on the *Drosophila* brain images. We show that similarity between images of neurons can be assessed based on the spatial innervation, the morphology, and the orientation of a neuron. Next, to perform image registration, we develop an algorithm to identify the fiber bundles. We treat the bundles as the skeletons of a brain and align different brain images based on these skeletons. Finally, we compare the AXON images with the fluorescent-image-based FlyCircuit database and identify new fiber bundles and features that are not available in the FlyCircuit database.

Keywords: A *Drosophila* connectome, AXON, the Golgi stain, neuronal classification, neuronal bundles

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