

# Research

---

## Research Overview

Nervous systems transform sensory signals and internal states into actions. Learning is a process by which this transformation is altered, producing different actions from the same stimulus. We study learning using the *Drosophila* model system. Flies can be trained to avoid an odor by pairing its presentation with electric shocks. This aversive memory is stored in a brain structure called the mushroom body that comprises roughly 2500 neurons per hemisphere. Olfactory signals arrive at the mushroom body neurons and cause a small fraction to fire action potentials. Pain signaling is thought to arrive via dopamine action on receptors linked to a calcium-stimulated adenylate cyclase. Elevated calcium resulting from action potentials coinciding with dopamine receptor activity are hypothesized to synergistically raise cyclic AMP levels. Cyclic AMP then mediates a range of synaptic plasticity responses that alter the fly's response to that odor.

We showed that aversive inputs in the mushroom body are mediated by dopaminergic cells in a single posterior cluster, projecting to restricted domains in the mushroom body. Our novel single-fly behavioral assay reveals differences between Pavlovian and operant conditioning. As there is no molecular framework for how action-contingency could influence learning, we are commencing a screen to find new genes and neurons that are involved in one process but not the other.

We can accept students through several programs, including the *Duke\_NUS doctoral program* and *a variety of programs offered by the A\*STAR Graduate Academy*.

<http://www.claridgechang.net/>