Optogenetic Modulation of Octopamine Neurons in Anesthetic Vulnerability in Drosophila melanogaster

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Results

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Introduction

Role of neurotransmitters in anesthetic usage remains largely unexplored, posing a significant risk of overlooking critical factors in human anesthetic administration

FIG 1: Control - Channel and Halo





- Acute activation of Octopamine was associated with less vulnerability to anesthetic and greater aversion
 - p value < 0.0001 → statistically significant
- Acute deactivation of Octopamine did not present a significant effect on anesthetic vulnerability
 ■ p value = 0.2201 → not statistically significant

 This study aimed to determine the effect of acute activation and deactivation of octopamine (OA) neurons on anesthetic vulnerability in fruit flies with the GAL4/UAS system (genetic tool to drive chosen gene expression)

Key terms:

Octopamine: neurotransmitter in fruit flies, equivalent of fight or flight in humans (norepinephrine)

- Findings highlight that Octopamine activation plays a significant role in anesthetic vulnerability
 - Highlights importance of considering additional factors in human anesthetic administration, such as norepinephrine levels and arousal states
 - Understanding the influence of octopamine on anesthetic susceptibility in Drosophila could inform better anesthetic management and personalized approaches in clinical settings, potentially improving patient outcomes

- Channelrhodopsin: facilitate neurotransmitter activation, are light-gated by wavelengths of approximately 700 nanometers (red light), Sodium ion channel
- Halorhodopsin: enable
 neurotransmitter deactivation,
 are light-gated by wavelengths
 of approximately 578
 nanometers (yellow light),
 Chloride ion channel







Image 1: Attempted Dissection



Image 2: Pupa under Leica Microscope









Offspring with OA neurons that inactivate by yellow light

Octopamine-GAL4

(female)



Image 3: Octopamine Diagram

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