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# How specific classes of retinal cells contribute to vision: A Computational Model

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## Introduction

Vision begins with the photoreceptors converting light from the visual scene into electrical signals, compressing our visual world into a code of action potentials sent to the brain by the **retinal ganglion cells (RGCs)**. A human retina contains almost **1 million RGCs** and each of these cells interprets **different features** of the visual scene (shape, motion, color, etc.). It is all these **parallel streams** of information received by the brain, that eventually lead to visual perception.

Currently, there exist over 30 RGCs subtypes based on:

- common anatomical features,
- functional properties,
- common gene expression.

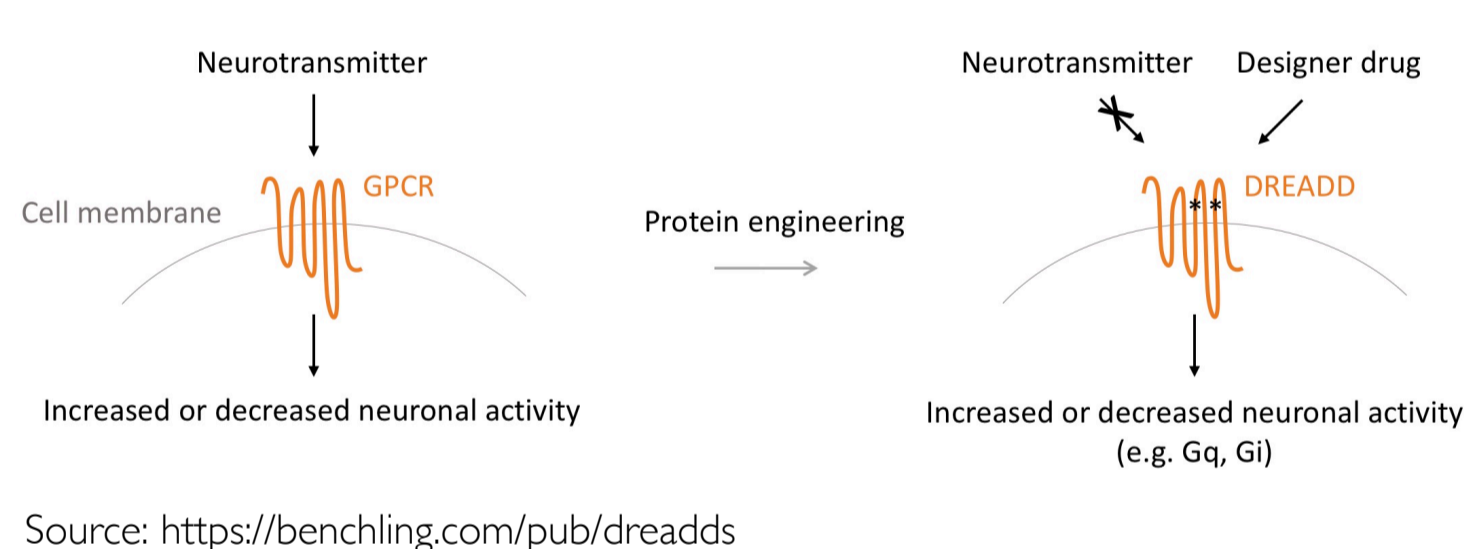
### Contemporary questions:

- ✓ What role does each RGC subtype play in vision?
- ✓ How is vision impaired if one of these subtypes is inactivated?

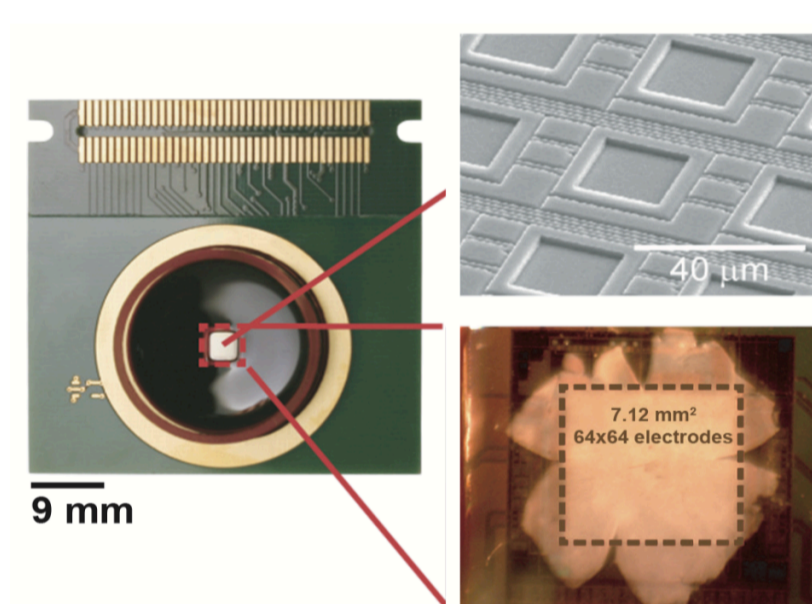
➤ We propose a novel approach combining for the first time **pharmacogenetics, electrophysiology, morphology, behavior and mathematical modelling** in order to **selectively inactivate specific RGCs types and decipher their role in vision**, both at the single cell and population level.

## Experiments

**Pharmacogenetics:** Modify neuronal activity **noninvasively and reversibly** by using the **Designer Receptors Exclusively Activated by Designer Drugs (DREADDs)** technique. DREADD-expressing RGCs' activity can be reduced or silenced when the synthetic designer drug clozapine N-oxide (**CNO**) is added to the solution.

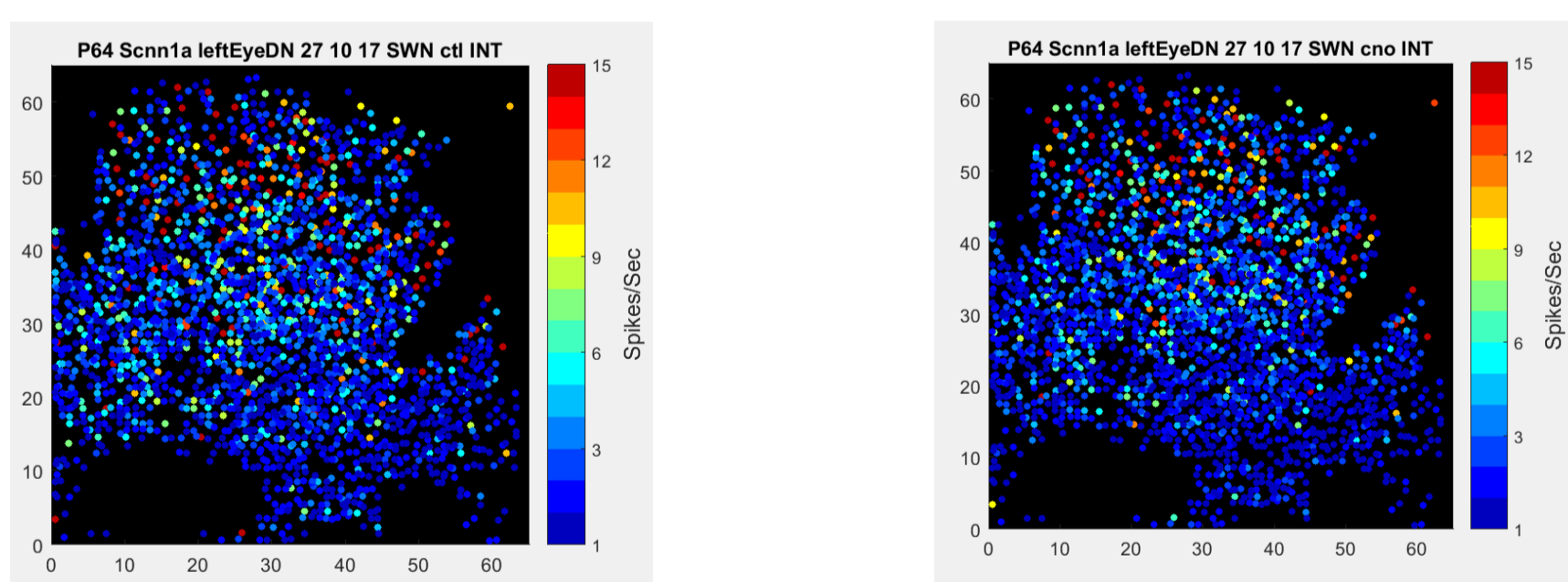


**Light stimulation and electrophysiological recordings:** Record light responses from the RGC layer at pan retinal level in vitro with the APS MEA in control and CNO conditions

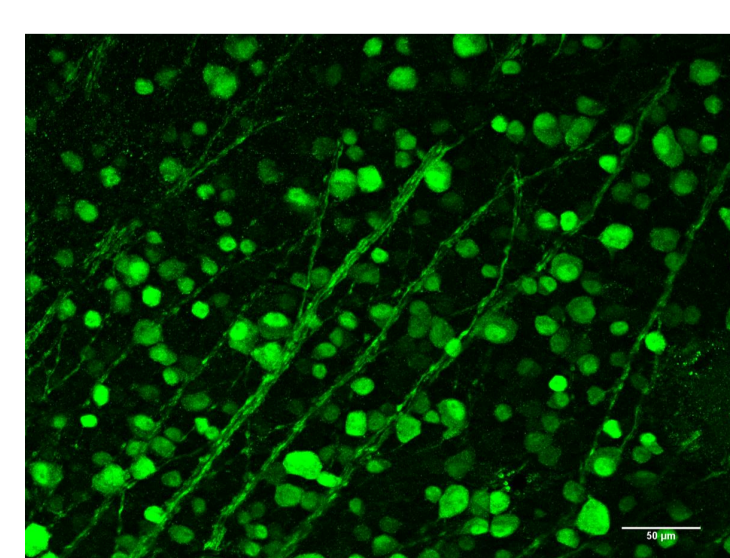


Active Pixel Sensor MEA Chip featuring 4,096 electrodes (42 μm spacing) arranged in a 64x64 configuration, covering an active area of 7.12 mm<sup>2</sup>. From Maccione et al., J Physiol 2014

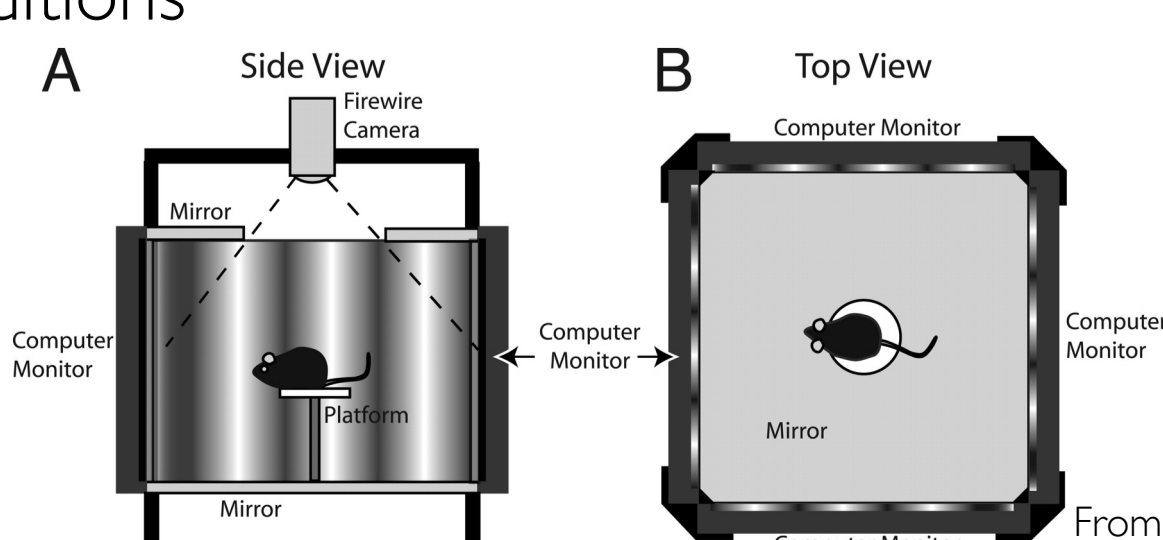
Pan-retinal recordings in control (left) and CNO (right) conditions



**Morphology:** Visualize cells and their dendritic trees by imaging under confocal microscopy post-recording.



**Behavioral testing:** Perform behavioral tests in control and CNO conditions



**Disclaimer:** All the experimental work is done at Pr. Sernagor's lab

## Computational analysis

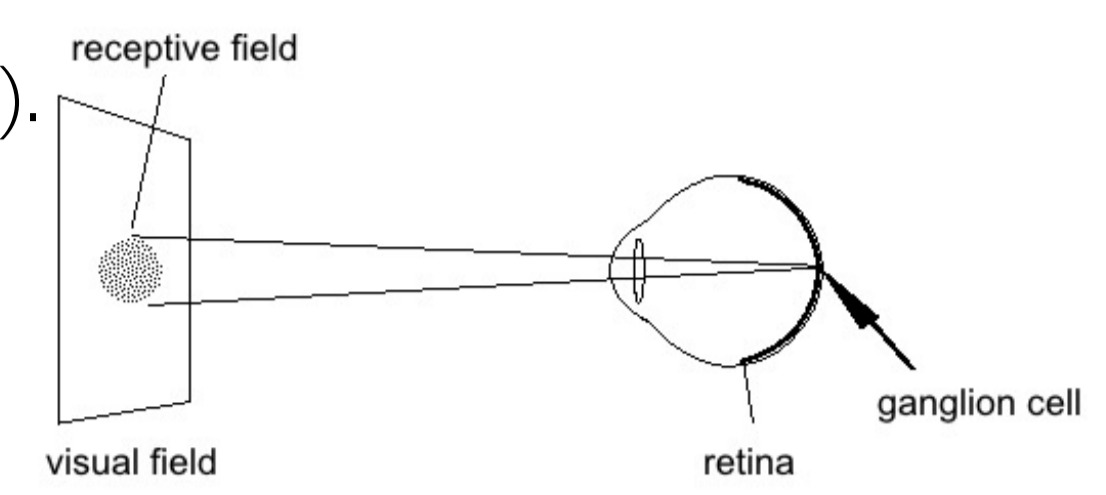
### How does CNO impact the response of RGCs to visual stimuli?

RGCs response to light stimuli is characterized by their **spatio-temporal receptive field (RF)**.

**Spatial profile:** the region of the visual field in which light stimuli evoke responses in a RGC.

The simplest RF is organized into a center-surround structure, responding oppositely to light.

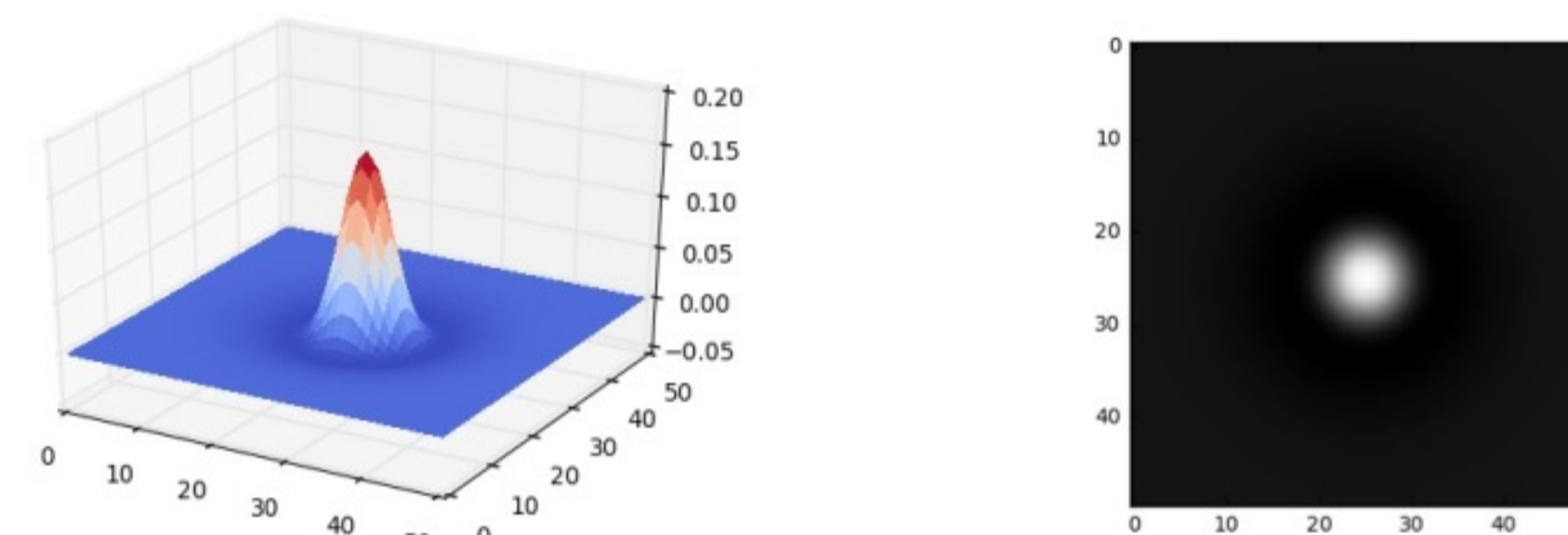
**Temporal profile:** the temporal course of this response.



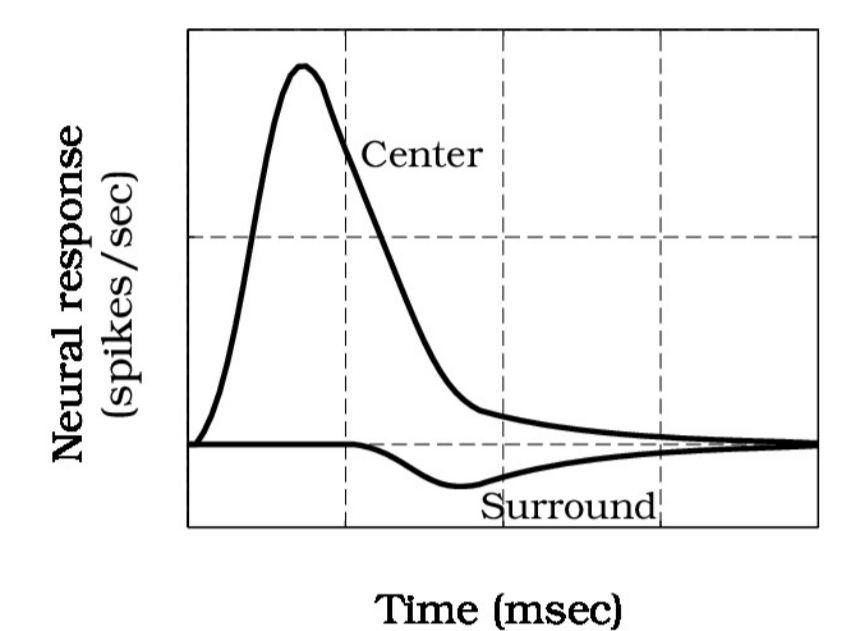
## Methods

### Use mathematical modelling

The difference of Gaussians function can model the **spatial RF**



The center-surround mechanism is **temporally biphasic**

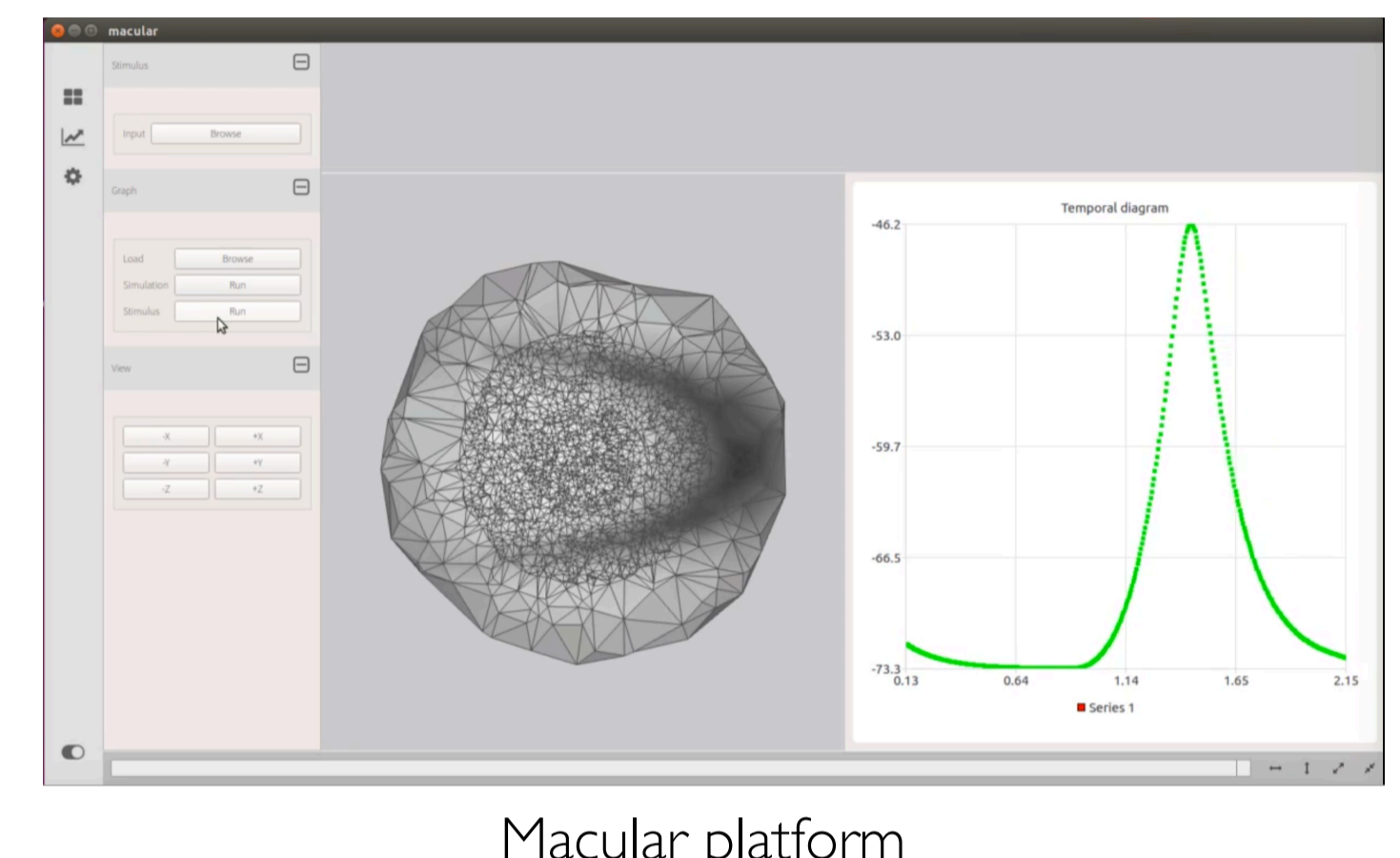


### Use numerical simulations

**PRANAS:** A platform for retinal analysis and simulation, developed by the Biovision team at Inria.

**MACULAR:** A platform for large scale simulations of the retina in pathological conditions (currently under co-development by the Biovision team and the CED engineering team at Inria).

**NEURON:** A simulation platform for modelling individual and networks of neurons, developed at Yale and Duke.



## Temporal profile

### Develop a generalized RGC model

Model single cell dynamics with **Hodgkin-Huxley** conductances using the NEURON simulation platform in order to reproduce what we observe in the experiments.

$$C \frac{dV}{dt} = (-g_L(V - V_L) - g_{Na}m^3h(V - V_{Na}) - g_Kn^4(V - V_K) - g_{CNO}(V - V_K))$$

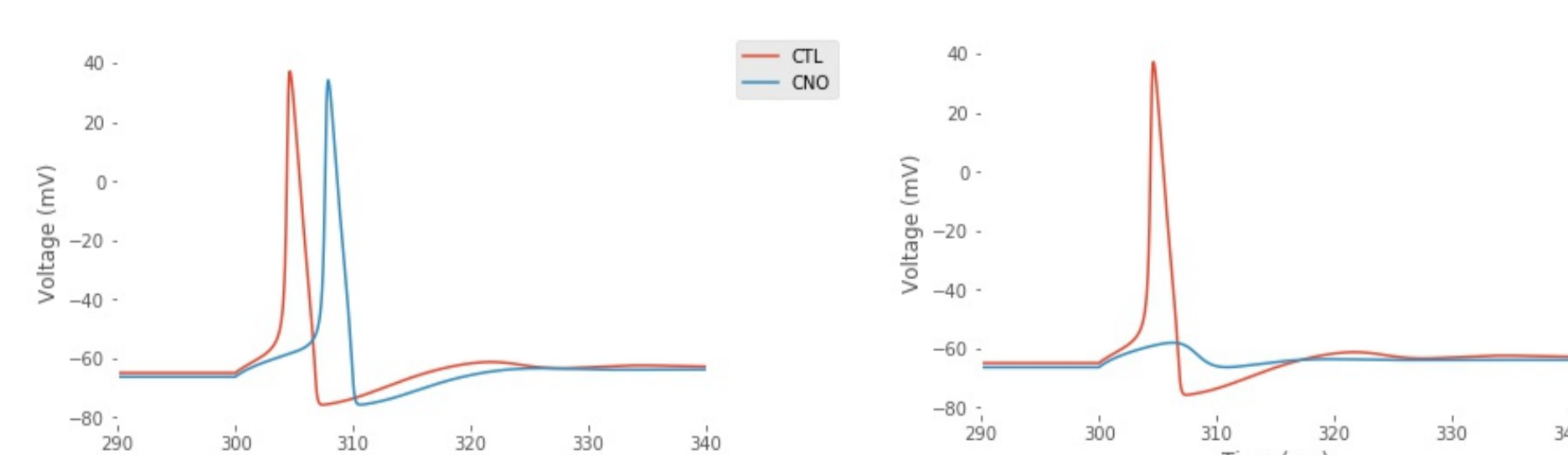
$$\frac{dm}{dt} = a_m(V)(1 - m) - m(V)m$$

$$\frac{dh}{dt} = a_h(V)(1 - h) - \beta_h(V)h$$

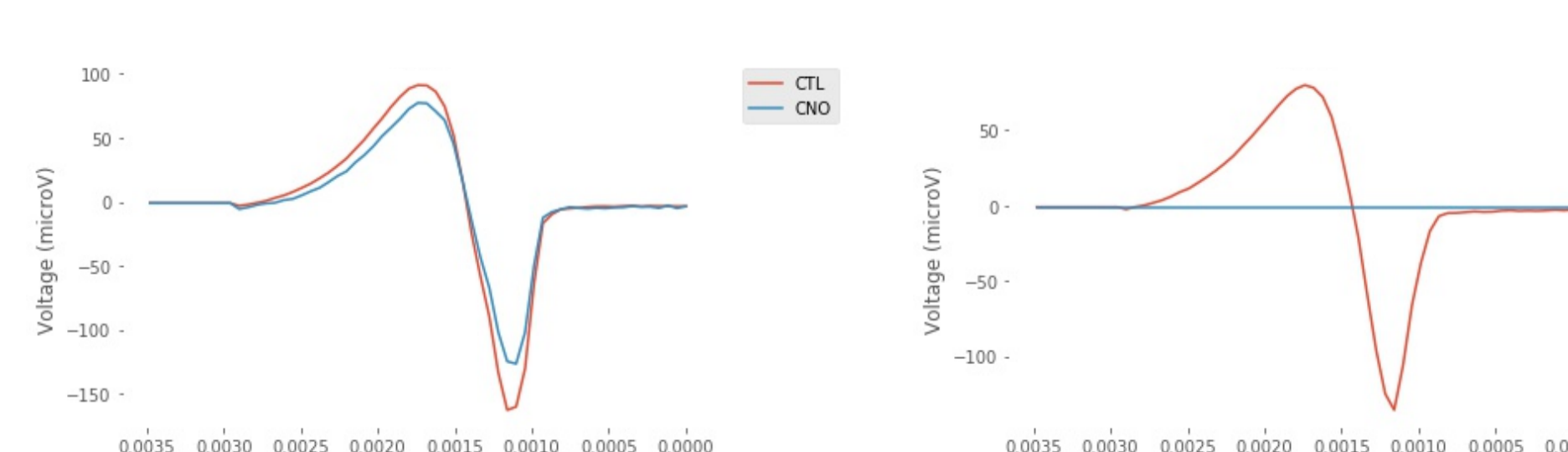
$$\frac{dn}{dt} = a_n(V)(1 - n) - \beta_n(V)n$$

To study the effect of CNO on the behavior of the cell, we added a leak potassium channel depended on [CNO], represented by the extra current  $g_{CNO}$ .

### Simulations



### VS Experiments



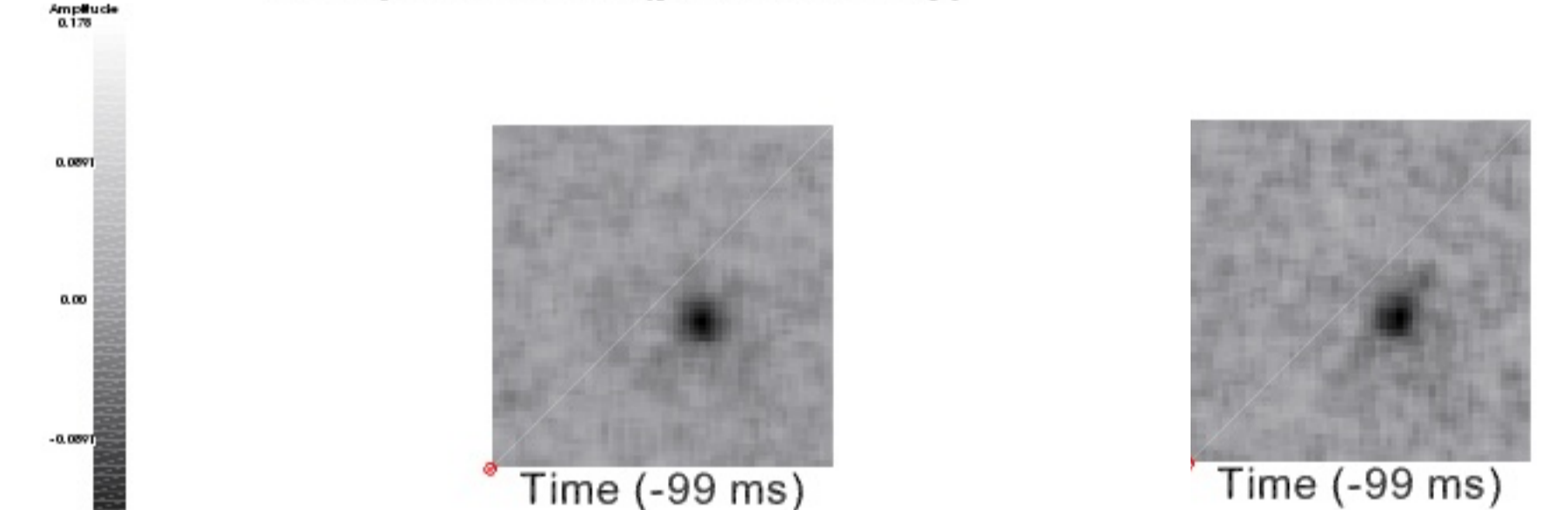
RGC's activity is reduced

RGC's activity is silenced

## Spatial profile

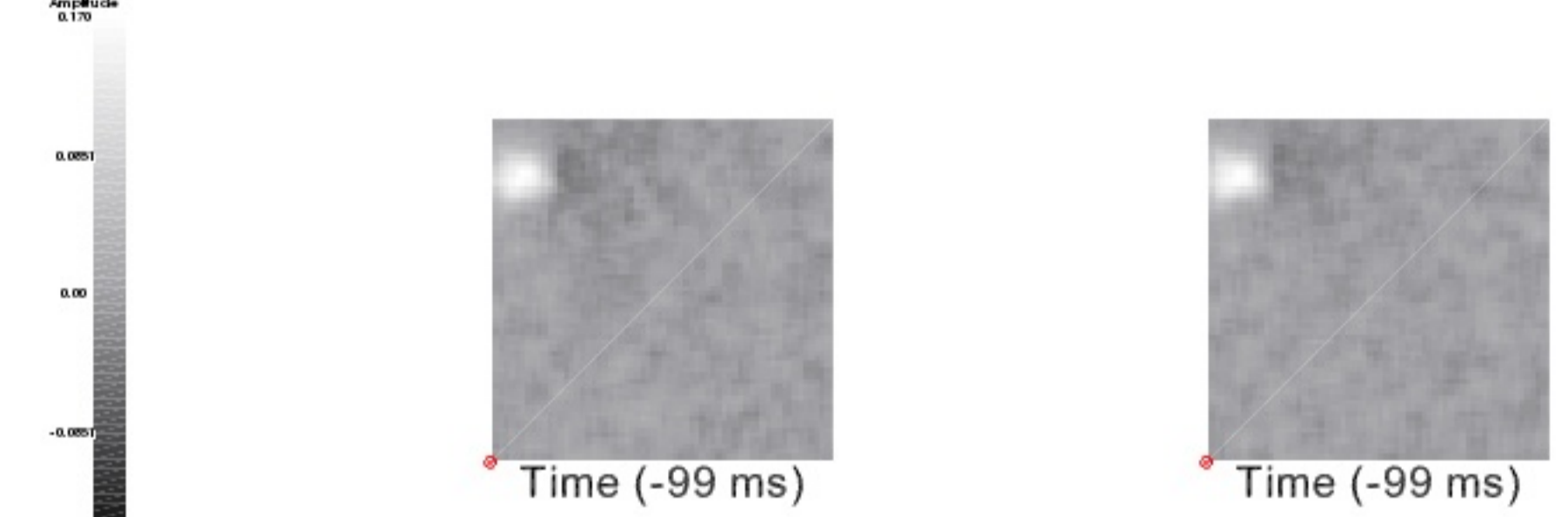
### RF estimation using PRANAS

#### Receptive fields (peak activity)



Estimated receptive field of an OFF RGC  
(A) In control condition (B) In CNO condition

#### Receptive fields (peak activity)



Estimated receptive field of an ON RGC  
(A) In control condition (B) In CNO condition

## Conclusions

- We reproduced the strong effect of CNO on the temporal part, as in the experiments.

### Next steps:

- Relate the CNO conductance to the CNO concentration
- Fit the experimental data, to tune the model's parameters
- There isn't a clear effect on the spatial part yet.
- There might be, if amacrine cells are also affected by CNO
- Mimic experimental setup with MACULAR.

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LEVERHULME TRUST

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