

Physiological Properties of Neurons in the Medial Amygdala

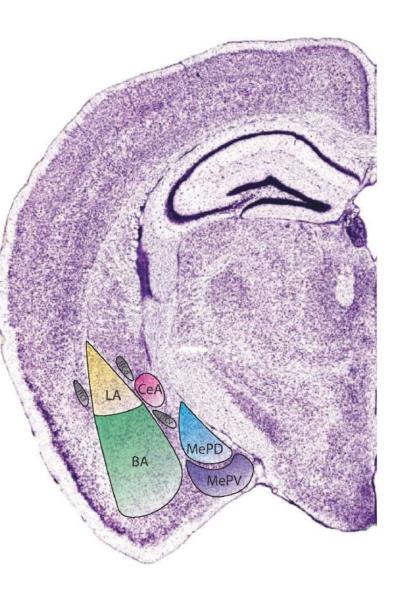


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Introduction

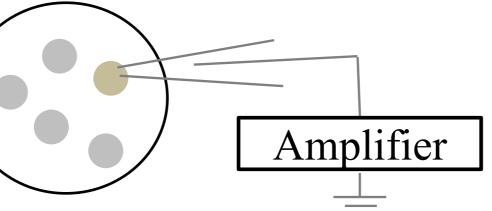


Medial Amygdala and Behavior

The amygdala is a conserved structure compose of multiple sub-nucleus (Janak et al., 2015). Recently, the roles of the medial amygdala in innate social and emotional behaviors have been reported (Hong et al., 2014)(Chen et al., 2019). By processing sensory information from olfactory system, the medial amygdala mediates social behaviors like grooming and parenting. Its function in sex-specific social behaviors has been highlighted.

Results

Slice Recording

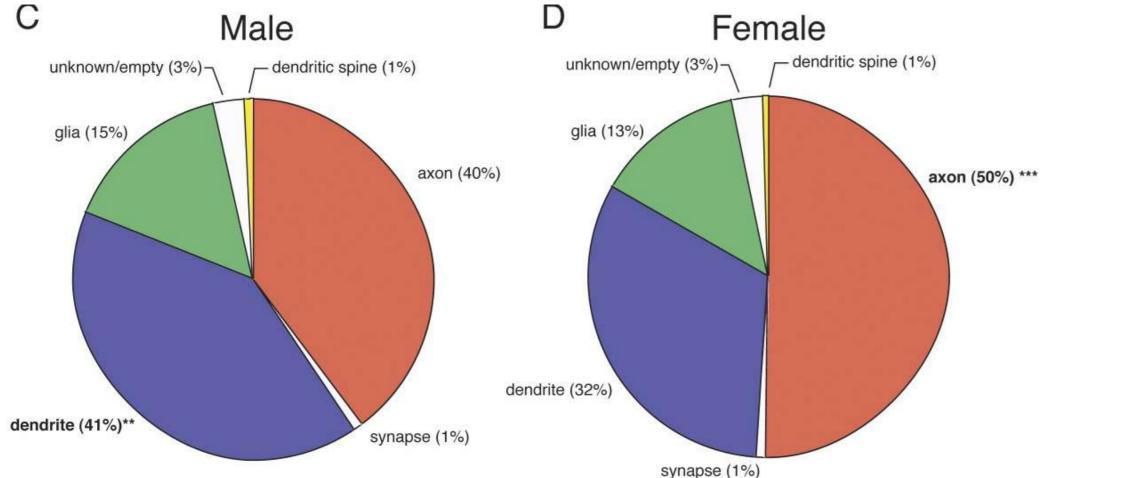


current-clamp: keep injecting current into the cell and see how the membrane potential change accordingly.

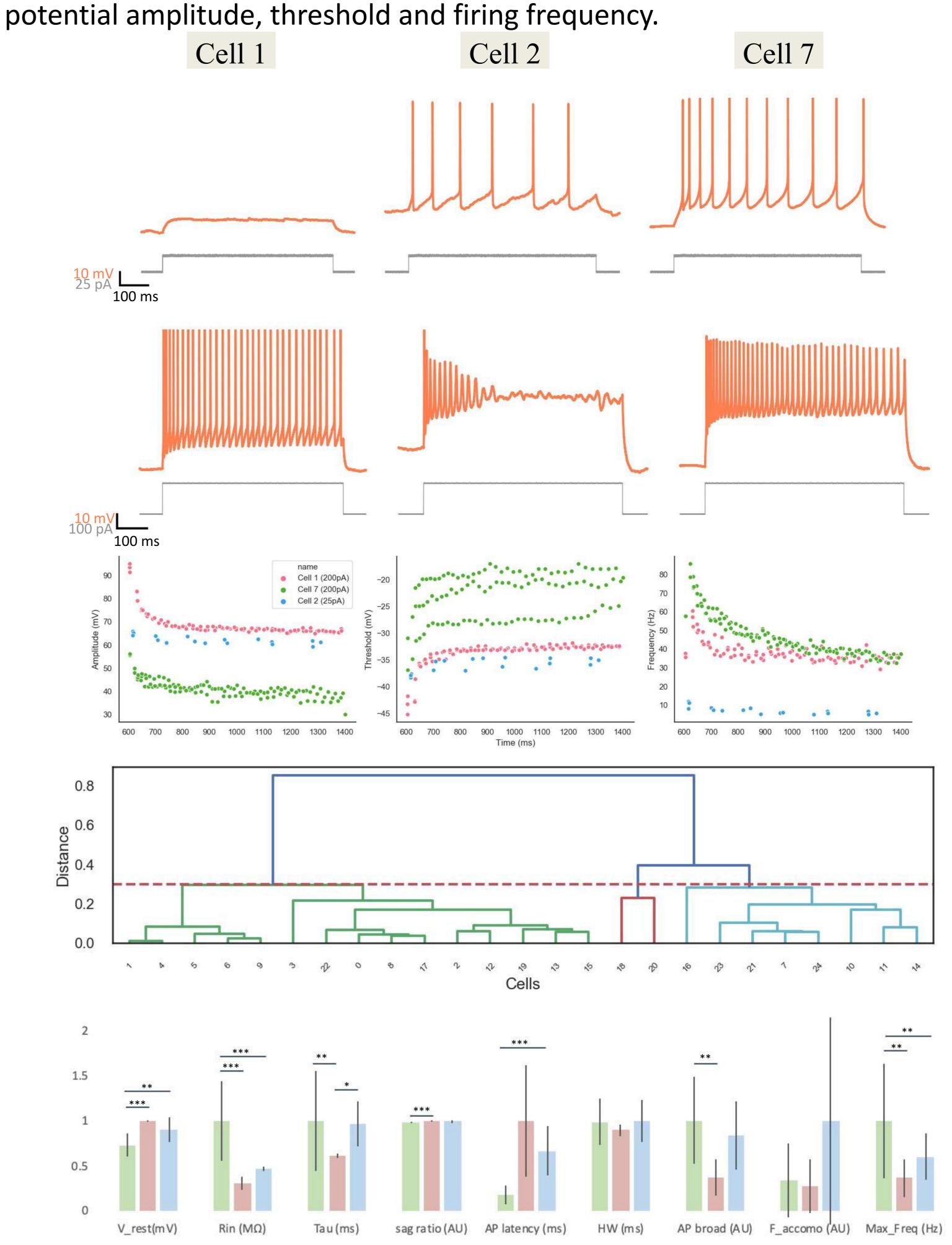
We injected 25pA and 200pA current into cells for 800ms and the traces are shown below. Neurons in the medial amygdala tend to have different action

Physiological Feature

The medial amygdala is one of the sexually dimorphic brain regions. Morphological differences between sexes have been reported (Cooke et al., 2007). High heterogeneity in molecular level implies distinctive populations among neurons in the medial amygdala (Chen et al., 2019).



Although the medial amygdala has been known to mediate several types of behaviors, little is known about its underlying neuronal mechanisms. Our project aims to analyze physiological properties of neurons in the



medial amygdala, including its electrophysiological parameters and projection sites.

Previous characterization of neurons in the medial amygdala using electrophysiological data (Keshavarzi et al., 2014) has mainly focused on male, here we introduce female mice into this characterization.

Summary

We used in-vitro electrophysiology to record neuron activity in the medial amygdala. Our preliminary results coordinate with the high heterogeneity previously reported. iDISCO imaging illustrates the projection pattern of neurons that serves as an important part of physiological properties, and also implies possible functional connectivity among brain subsets.



Automatic Tracking

We use subtraction, fast Fourier transform and low-pass filter to track the mass center of mouse on a frame basis. The heatmap and data analysis

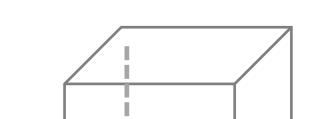
iDISCO

iDISCO enables whole-mount immunolabeling with volume imaging of large cleared samples. We tested this method on antero-traced MeA neurons.

PFA fixation

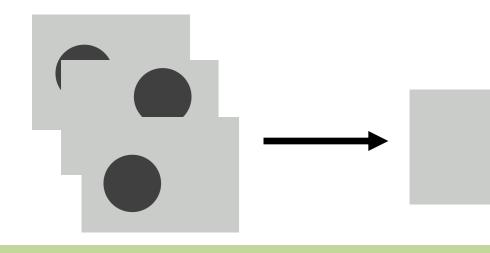
Imaging





were then conducted using location information.

Average 100 frames to get reference image





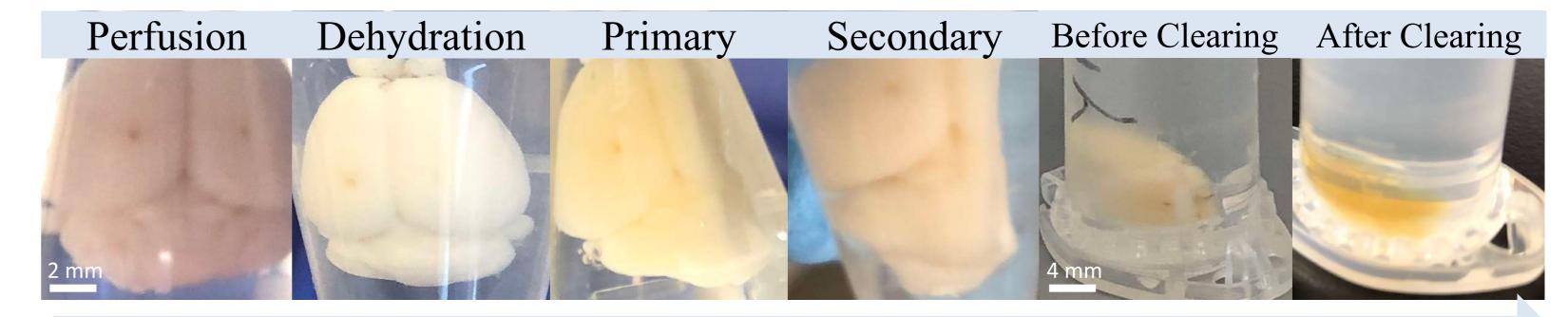
References

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Immunolabeling Clearing







Lateral (50um interval)

