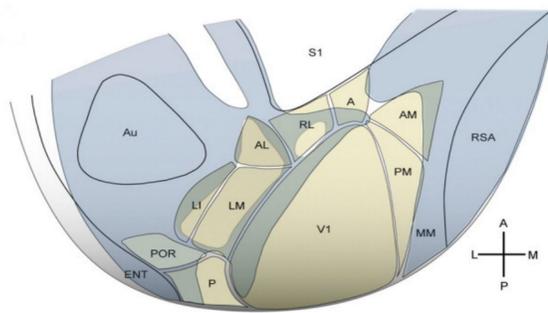


## Introduction

- **Goal: Explore effects of visual stimulus on firing rates of individual and ensembles of neurons in six distinct mouse brain regions.**
- Recordings were made from **six mouse brain regions** presumably involved in visual processing: the Primary Visual Cortex (**V1**), the Lateromedial Area (**LM**), the Anterolateral Area (**AL**), the Rostrolateral Area (**RL**), the Posteromedial Area (**PM**), and the Anteromedial Area (**AM**).



## Data and the experiment

- Data were collected from three individual mice, using **six NeuroPixels probes, which provided simultaneous recordings of many spiking neurons from all six brain regions** mentioned above.
- Each mouse was presented with drifting gratings under different combination of parameters, **15 times each for two seconds**, which provided **600 trials** of data.
- The two parameters accounted for **orientation (0°, 45°, 90°, 135°, 180°, 225°, 270°, 315°)** and **temporal frequency (1 Hz, 2 Hz, 4 Hz, 8 Hz, and 15 Hz)** of the gratings.

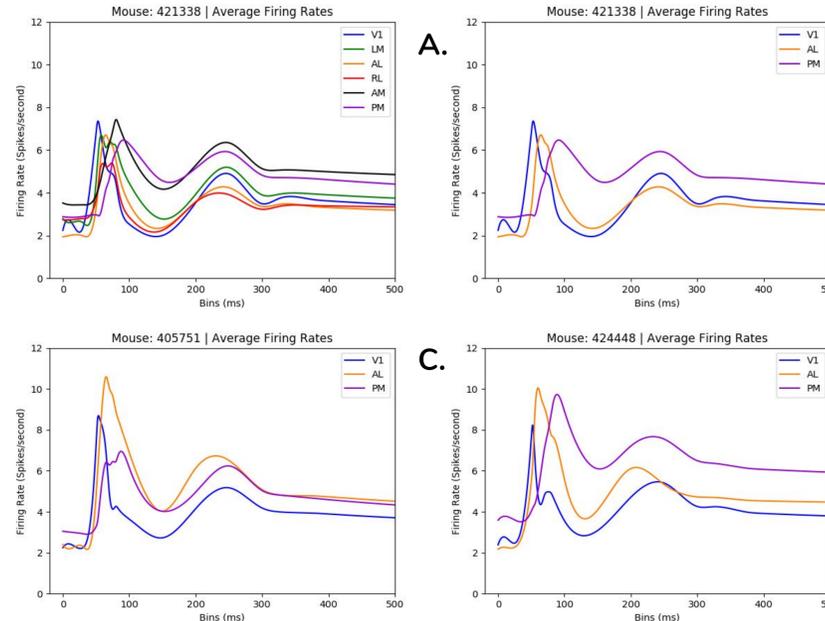


## Methods

- **For each brain area, univariate regression splines with 17 knots were used to smooth population peri-stimulus time histograms based on spikes pooled across all neurons within each region. We used the fitted curves to find the time of maximal population firing rate.**
- **Within-trial analyses** were based on two-way analysis of variance (**ANOVA**) and **paired differences**.

## Population spiking rates reveal order of firing

- In all cases, **V1 is the first region to reach its maximum firing rate and PM is the last.**
- All curves have roughly the same behavior, peaking for the first time between 50ms and 100ms and again between 200ms and 250ms. Let time of the **first peak** be  $t_{max1}$  and the **second** be  $t_{max2}$ .



- Average firing rate graphs for Mice **A, B, and C** demonstrates ordering based on  $t_{max1}$ :

$$V1 < AL < PM$$

- Table 1 solidifies that paired differences between **AL - V1** and **PM - AL**, based on above ordering, are highly significant since p values for the pairs in all mice are less than  $10^{-6}$ .

Table 1:  $t_{max1}$

Mouse	Paired Difference	Mean Time (ms)	95% Confidence Interval	p-value
A	AL - V1	6.2	(5.2, 7.3)	$p \ll 10^{-6}$
A	PM - AL	4.8	(3.6, 6.1)	$p \ll 10^{-6}$
B	AL - V1	6.5	(5.4, 7.6)	$p \ll 10^{-6}$
B	PM - AL	3.0	(1.8, 4.1)	$p = 6.0 \times 10^{-7}$
C	AL - V1	4.4	(3.2, 5.6)	$p \ll 10^{-6}$
C	PM - AL	5.5	(4.3, 6.8)	$p \ll 10^{-6}$

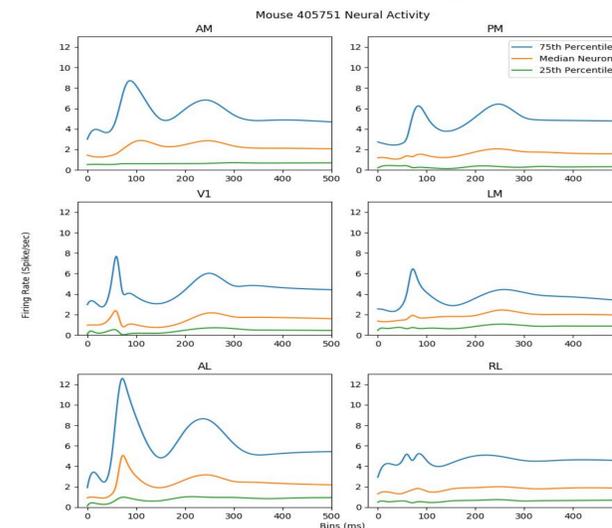
- Results from two-way ANOVA shows that **area means are highly significantly different**  $p \ll 10^{-6}$  ( $R^2 = 7.9\%$ ).
- **Trial-to-trial variation is highly significant**  $p \approx 10^{-6}$  ( $R^2 = 19.6\%$ ).
- **All paired differences between AL, RL, and LM, and also between AM and PM were insignificant**,  $p = 0.023$  with exceptions in mouse B (RL-AL,  $p \approx 10^{-6}$ ) and in mouse C (AL-RL,  $p \approx 10^{-6}$ , PM-AM,  $p \approx 10^{-6}$ ).

## Percentiles show limited variation in the lowest firing neurons

- For every time bin we compute the 25th, 50th, and 75th percentiles of firing rates among neurons, in each brain region. Curves are shown for mouse B.

**75th percentile** → follows the shape of the population average

**25th percentile** → shows almost no variation



## Discussion

- We took advantage of the simultaneous records from each area by using a within-trial analysis, which provided very strong evidence of the time ordering:

$$V1 < \begin{pmatrix} RL \\ AL \\ LM \end{pmatrix} < \begin{pmatrix} AM \\ PM \end{pmatrix}$$

- Many of our analyses revealed substantial noise in the measurements. This suggests that finding additional relationships may be challenging.

## Future plans

- Analyze separately neurons with high firing rates for each given direction.
- Determine correlation between change points of maximal firing rate curves, across all trials and across all regions.
- Refine the regression spline methodology in order to do addition within trial analysis.

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