# Interfacing with the brain: improving materials and devices for neural recording

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### 1. Why simultaneously monitor the activity of large number of neurons?

 $(x_0, y_0, z_0)$ 

100 µm

- Considerable progress is still necessary to reliably increase the neurons that are number of identified recorded and during simultaneously extracellular recordings.
- Extracellular recordings face a disagreement between the typical number of neurons observed and the number expected based on





#### anatomical considerations.

Figure 1. a) Scheme of a silicon probe comprising a dense electrode array. Geometric visualization of spontaneous neural activity from motor cortex in layer 5; b) photograph of 100-µm-thick section stained from M1 of the rat. The number of neurons in the detectable volume (within 50 µm) of this probe should be around 89 cells and; c) the signal transference from the tissue to the electrode depends on its impedance.

275 µm

## 2. Precision dual-probe setup for evaluating ultra-high density devices, new materials and spike sorting algorithms



Figure2. Automated surgery-physiology setup composed by a microscope to align both electrodes and high precision motors.

> Simultaneous extracellular and juxtacellular recordings from the same neurons can provide the 'ground-truth' data required for the probe and materials evaluation. Juxtacellular signature provides unambiguous detection of single neuron.



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time (s)

time (ms)

### 4. Material electrode-brain interface



Recordings from two extracellular probes within 100um were achieved by using a ThomasTetrode.

# **3. Electrode material**

> We are evaluating different interfaces for improving recording and isolation methods, as well as the effect of electrode features on the signal transfer from tissue to electrode.









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