

# Brain Signal Acquisition

# Partially Invasive Approach

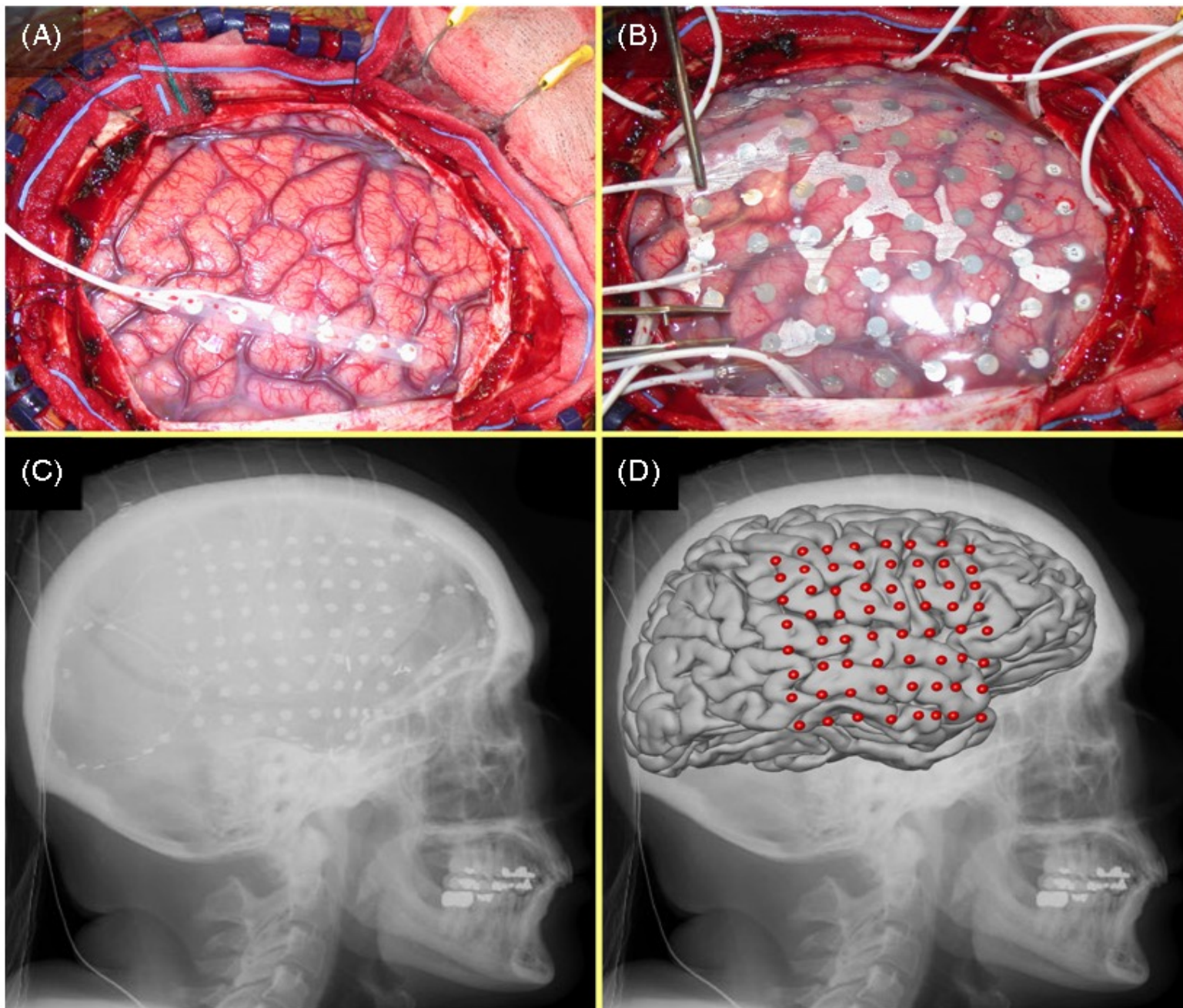
## **Electrocorticography (ECoG):**

- *Electrocorticography* (ECoG) is a technique for recording brain signals that involves placing electrodes on the surface (cortex) of the brain.
- The procedure requires making a surgical incision into the skull to implant the electrodes on the brain surface

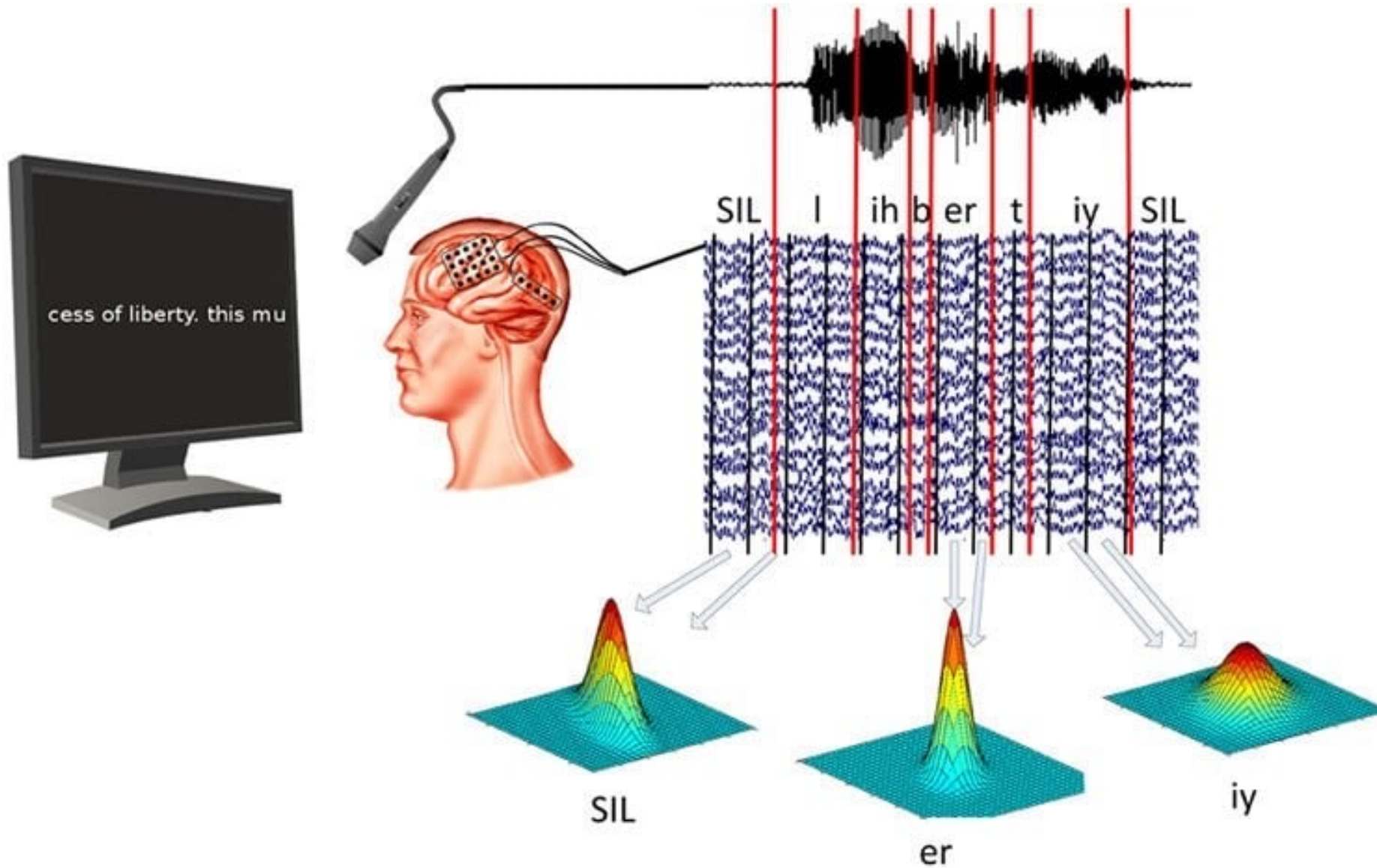
# Partially Invasive Approach

- ECoG electrodes can record the electrical fluctuations caused by the **coherent activity of large populations of neurons** (several tens of thousands).
- **Safer** than arrays implanted inside the brain.
- ECoG electrodes may also be **less likely to wear out** compared to brain penetrating electrodes
- ECoG offers greater **spatial resolution**





(from (Miller et al., 2007)).



Automatic Speech Recognition from Neural Signals: A Focused Review” by Christian Herff and Tanja Schultz in *Frontiers in Neuroscience*. Published online September 27 2016 [doi:10.3389/fnins.2016.00429](https://doi.org/10.3389/fnins.2016.00429)

ECoG and audio data are recorded at the same time. Speech decoding software is then used to determine timing of vowels and consonants in acoustic data. ECoG models are then trained for each phone individually by calculating the mean and covariance of all segments associated with that particular phone.

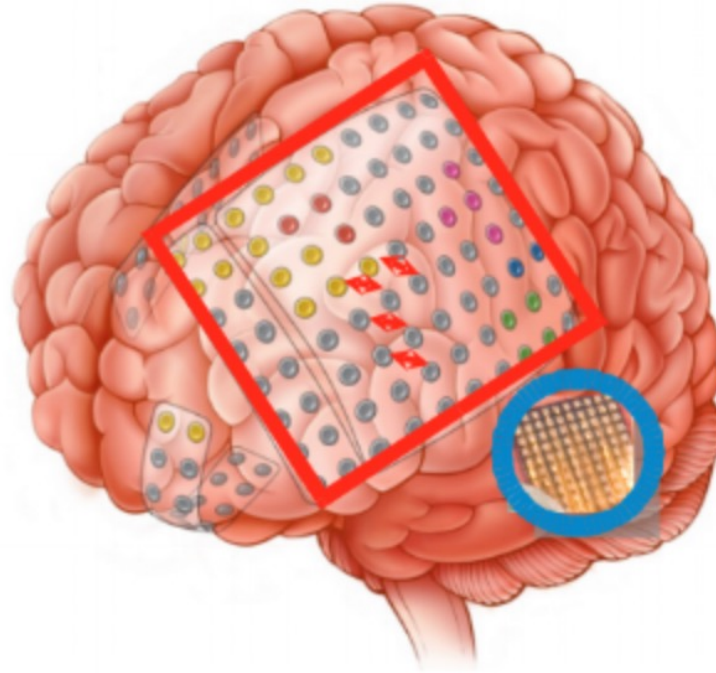
# Partially Invasive Approach

## **MicroECoG:**

- One disadvantage of ECoG, is the relatively large size of ECoG electrodes
- These microelectrodes are only a fraction of a millimeter in diameter and spaced only 2–3 mm apart in a grid
- Allows detection of neural activity at a much finer resolution than traditional ECoG.
- Decoding fine movements, such as the movements of individual fingers, or even speech, without actually penetrating the brain.



# Partially Invasive Approach



## Current ECoGs

- Large area
- Low resolution



## Current $\mu$ ECoGs

- Small area
- High resolution



# Partially Invasive Approach

## Optical Recording: Voltage-Sensitive Dyes and Two-Photon Calcium Imaging:

- Voltage-sensitive dyes
- Voltage-sensitive dye changes its absorbance and fluorescence intensity when membrane potential changes in a stained brain or heart tissue.
- By using voltage-sensitive dyes as chemical probes and capturing changes in light intensity with the use of a high-speed imaging device, it is possible to image in real time the activity of where, when, and how much excitation or inhibition occurred, in the brain and heart.



# Partially Invasive Approach

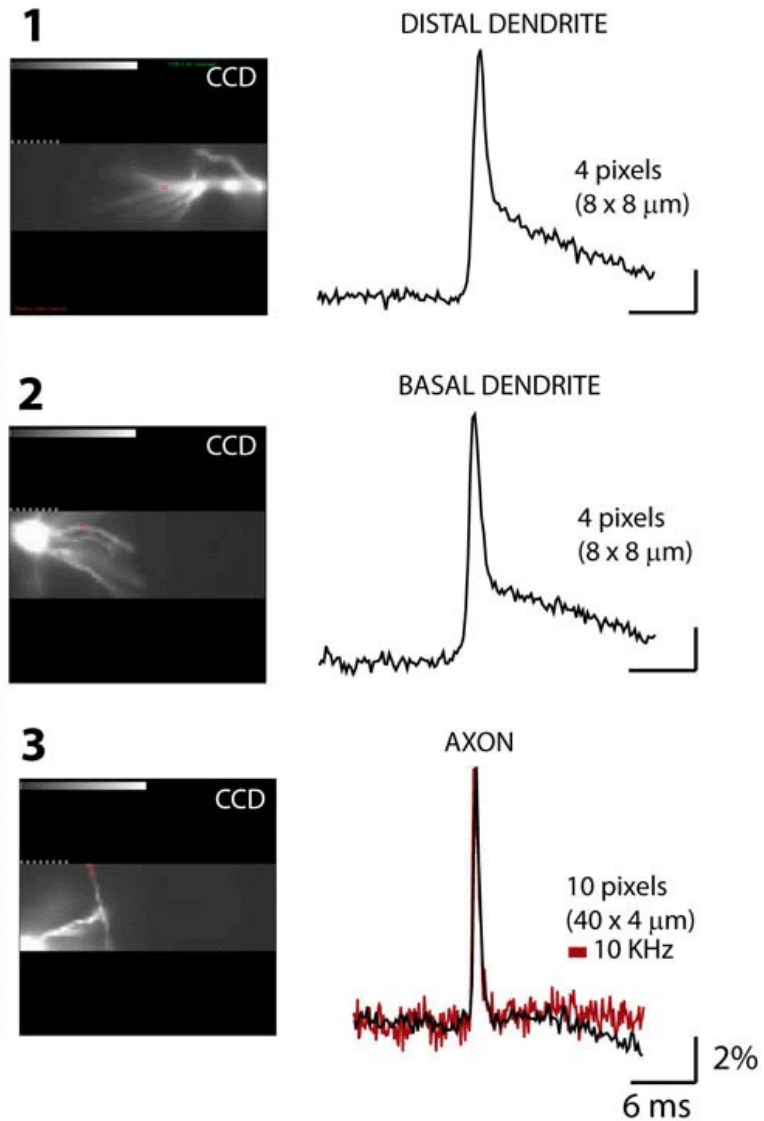
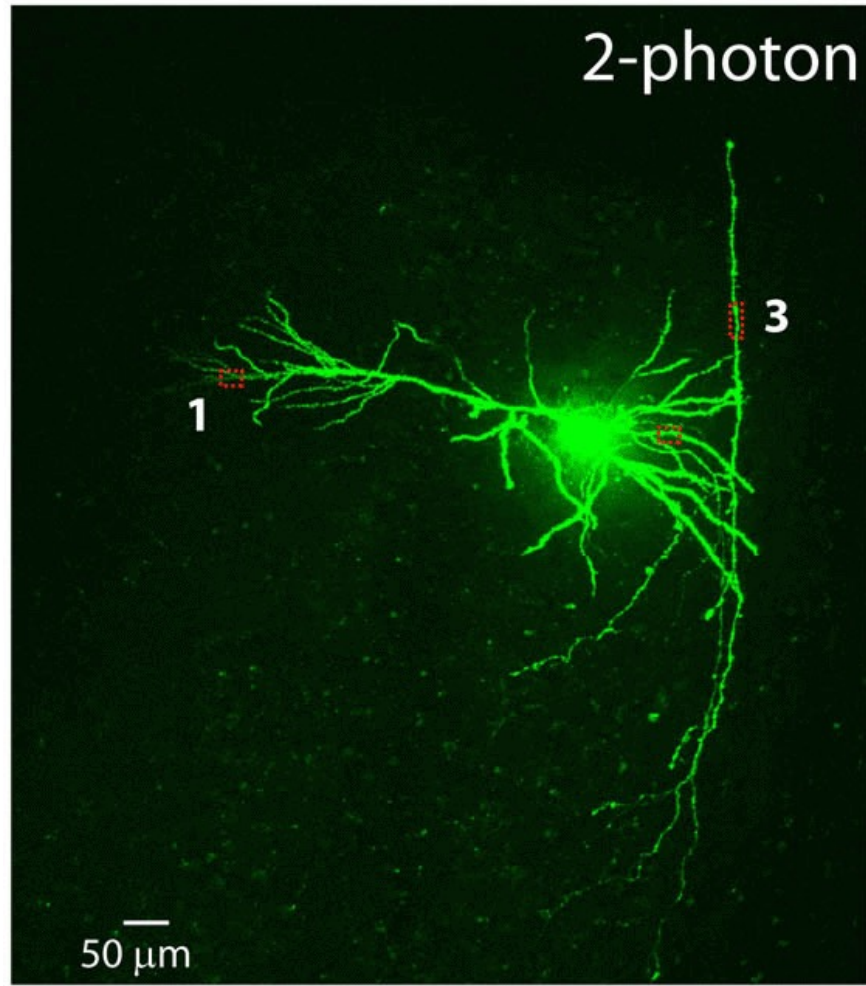
## Voltage-sensitive dyes

<https://www.scimedia.com/applications/neuro/vsd/data/>

- Neurons are **stained** with a voltage-sensitive dye
- **Dye responds** to changes in membrane potential by changing its **absorption** and/or **fluorescence**
- Recorded optical signals correspond to **summed responses** from several **simultaneously active neurons**.
- Useful for **imaging macroscopic features** of the brain.

# VOLTAGE-SENSITIVE DYE IMAGING

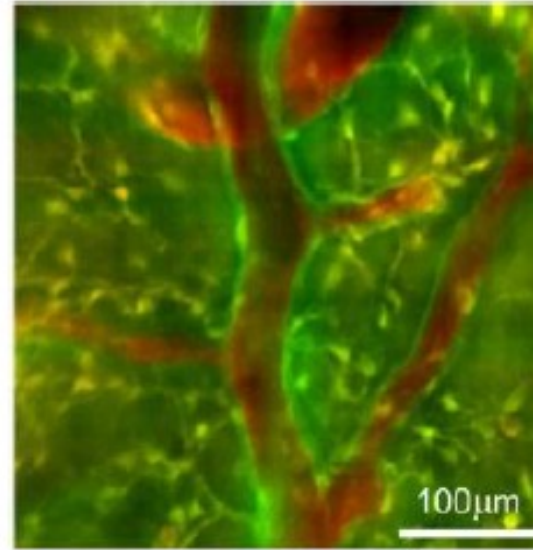
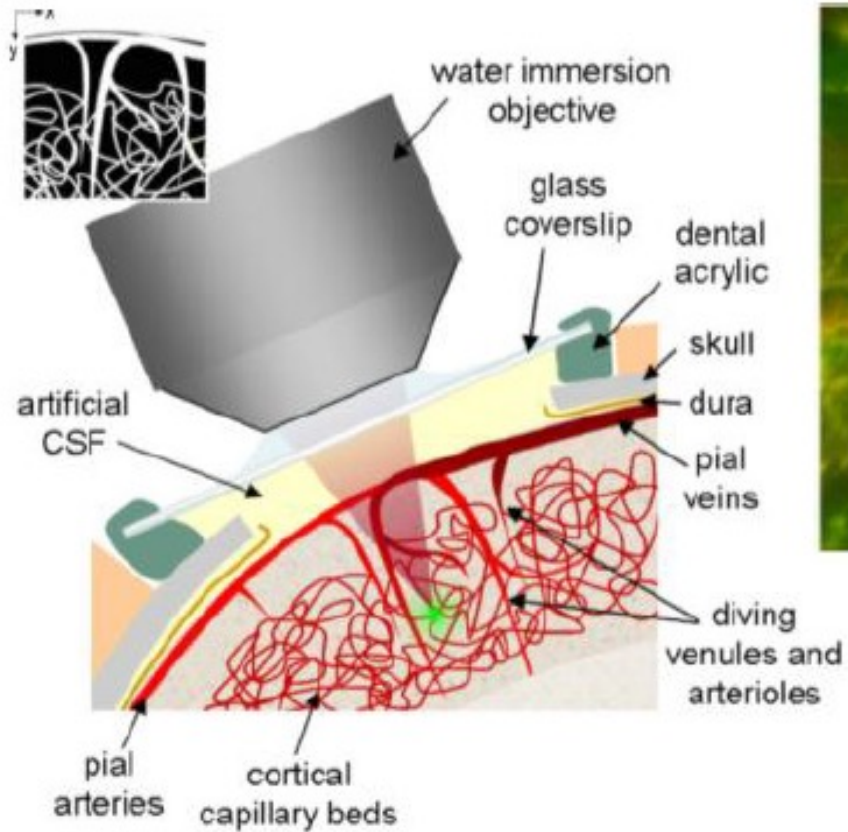
SINGLE TRIAL RECORDINGS AT 5 KHz



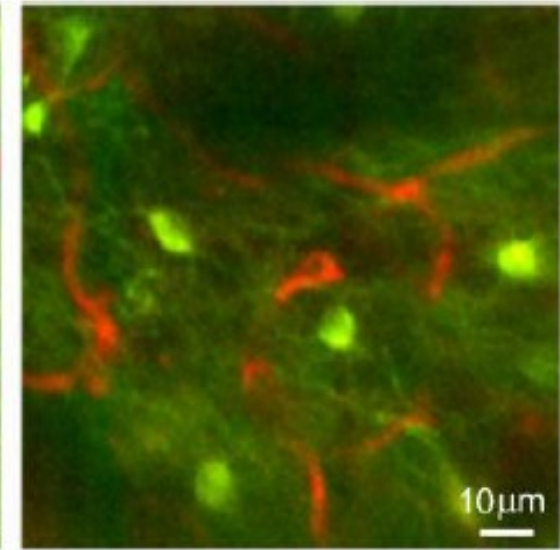
(image: Scholarpedia [http://www.scholarpedia.org/article/Voltage-sensitive\\_dye](http://www.scholarpedia.org/article/Voltage-sensitive_dye)).

# Partially Invasive Approach

- Two-photon calcium imaging
- <https://elifesciences.org/articles/26839#content>
- Based on the fact that electrical activity in neurons is typically associated with changes in calcium concentration.
- Two-photon calcium imaging allows us to observe the activity of multiple neurons up to  $\sim 500 \mu\text{m}$  below the cortical surface without cortical invasion.
- Photon calcium imaging involves:
  - (1) using pressure ejection to load neurons with fluorescent calcium-indicator dyes
  - (2) monitoring changes in calcium fluorescence during neural activity using two-photon microscopy.



Oregon Green  
calcium  
sensitive dye  
stained  
neurons



Transgenic mouse  
expressing green  
fluorescent protein  
(GFP) in a  
subpopulation of  
neurons

Image from Kherlopian et al., 2008).