

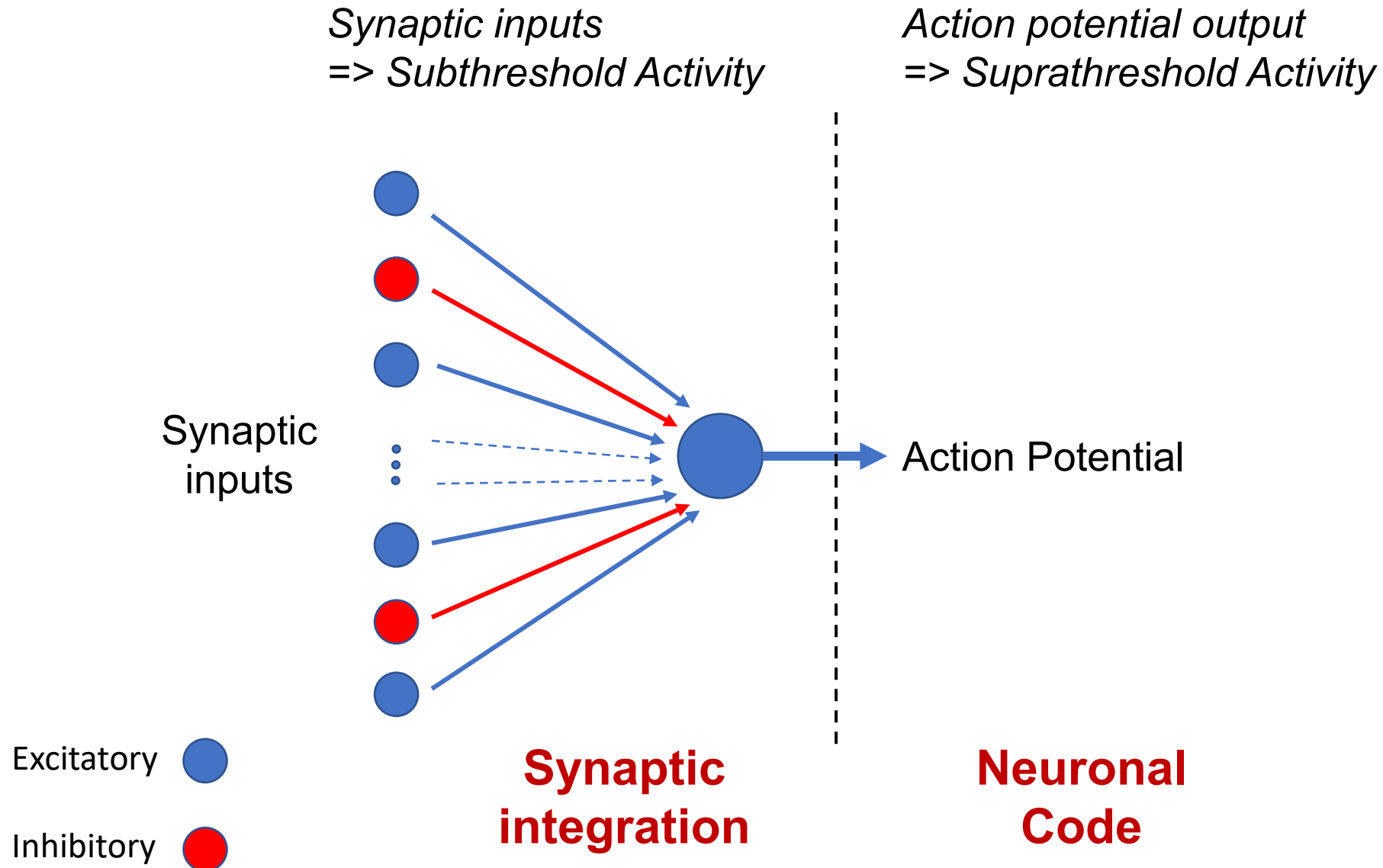
BIO-482 Neuroscience. Cellular and Circuit Mechanisms

**What can we learn from membrane potential
recordings in awake behaving mice?**

Sylvain Crochet & Carl Petersen

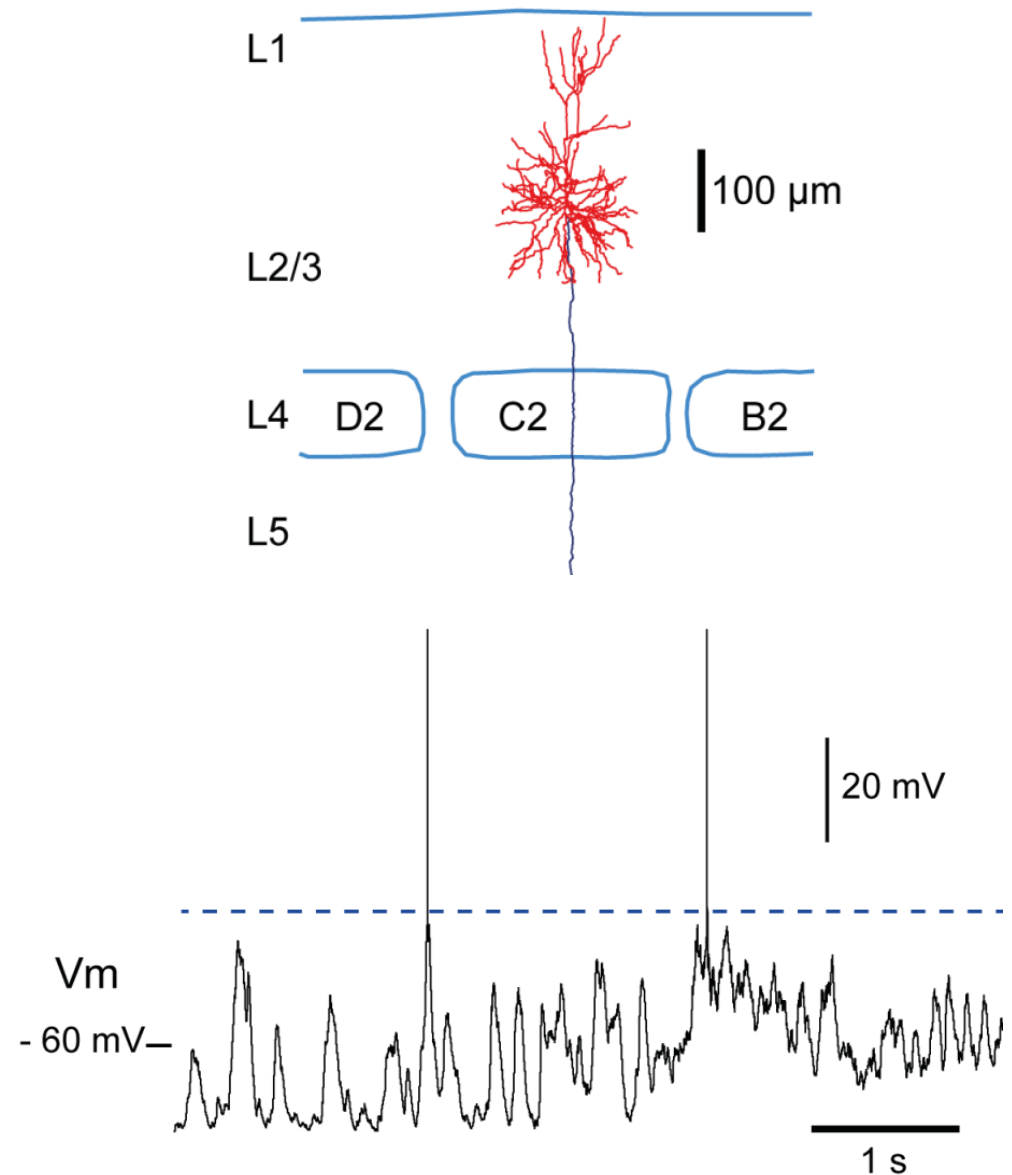
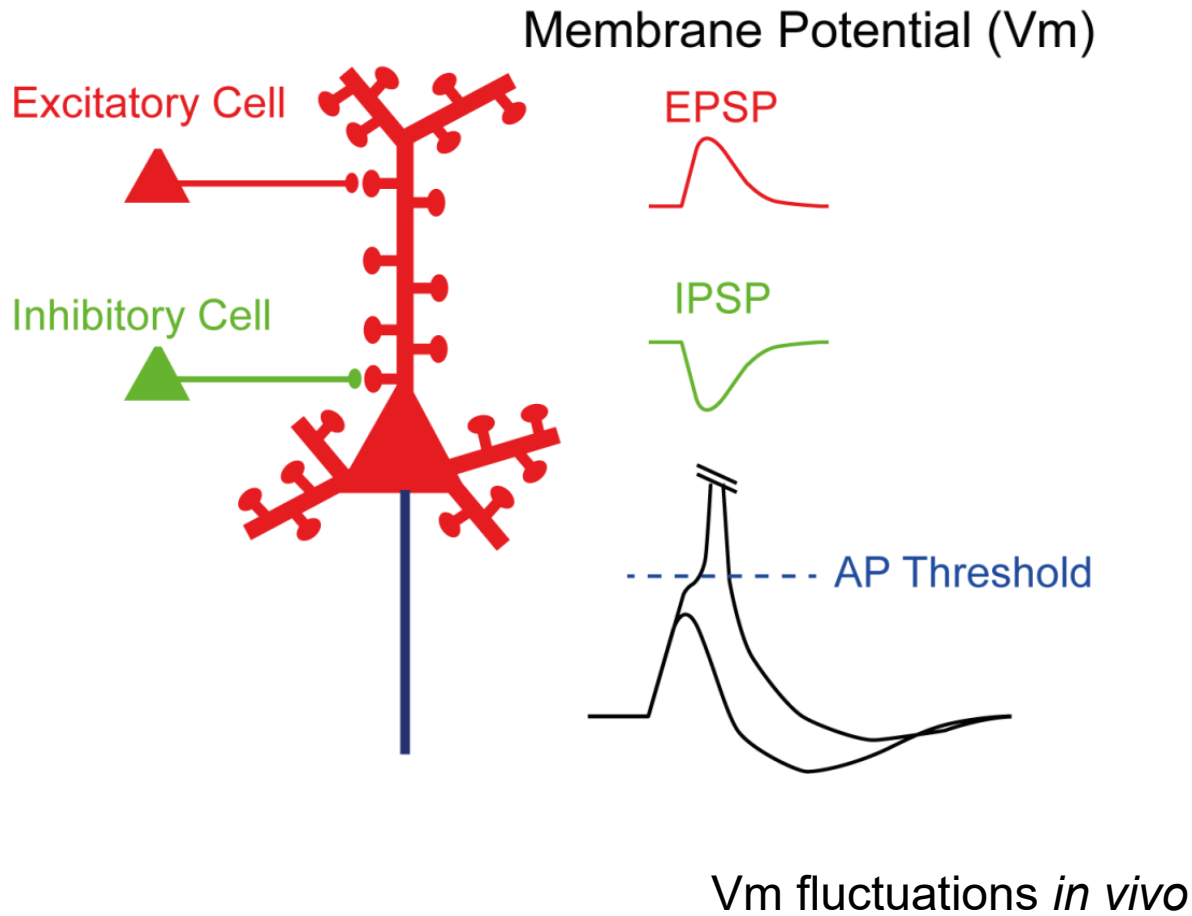
Laboratory of Sensory Processing - LSENS

■ Cortical neurons as synaptic integrators



Cortical neurons as synaptic integrators

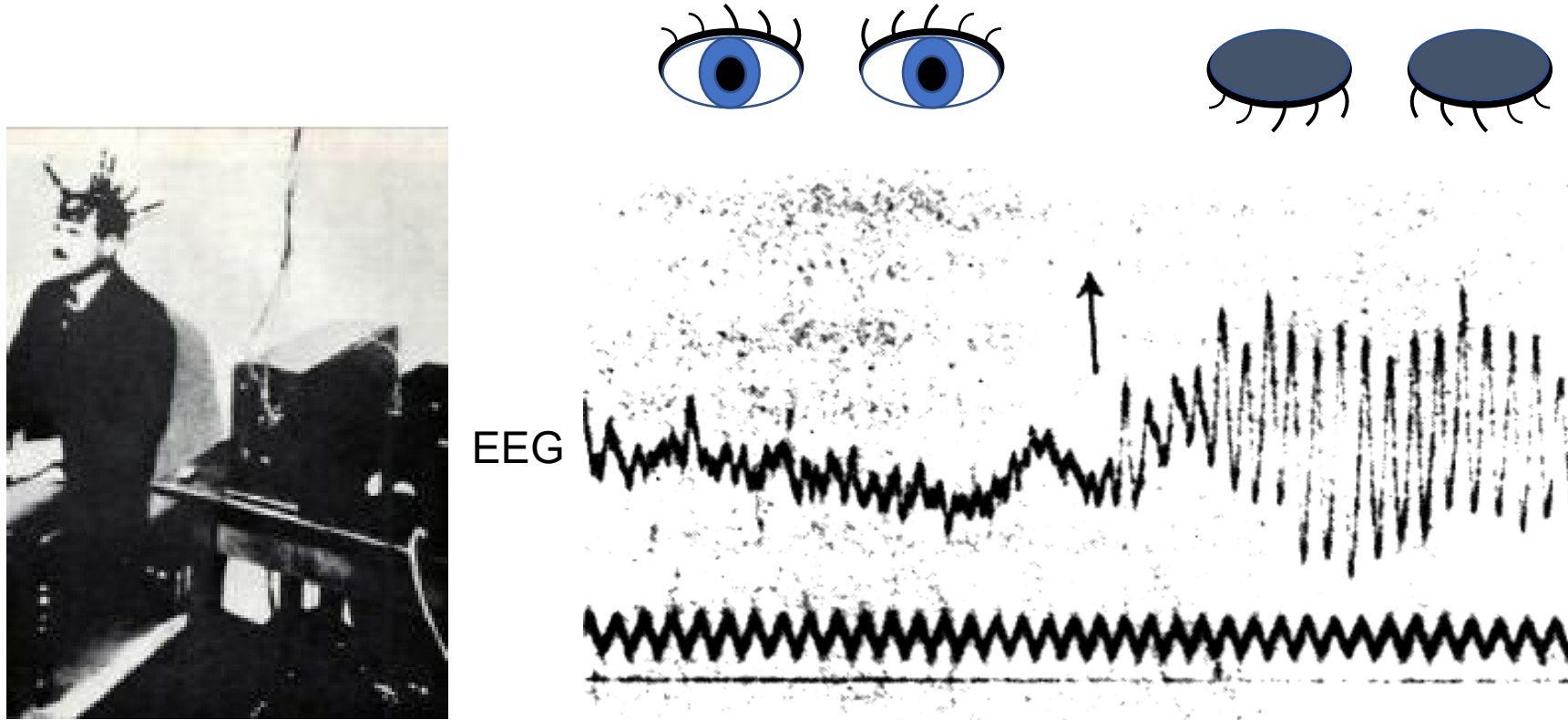
- . Cortical neurons receive ~1000 synaptic inputs
- . Mostly from other cortical neurons
- . ~80% excitatory / 20% inhibitory



1. Cellular mechanisms underlying cortical state changes in awake mice

- At the origin of *cortical states*

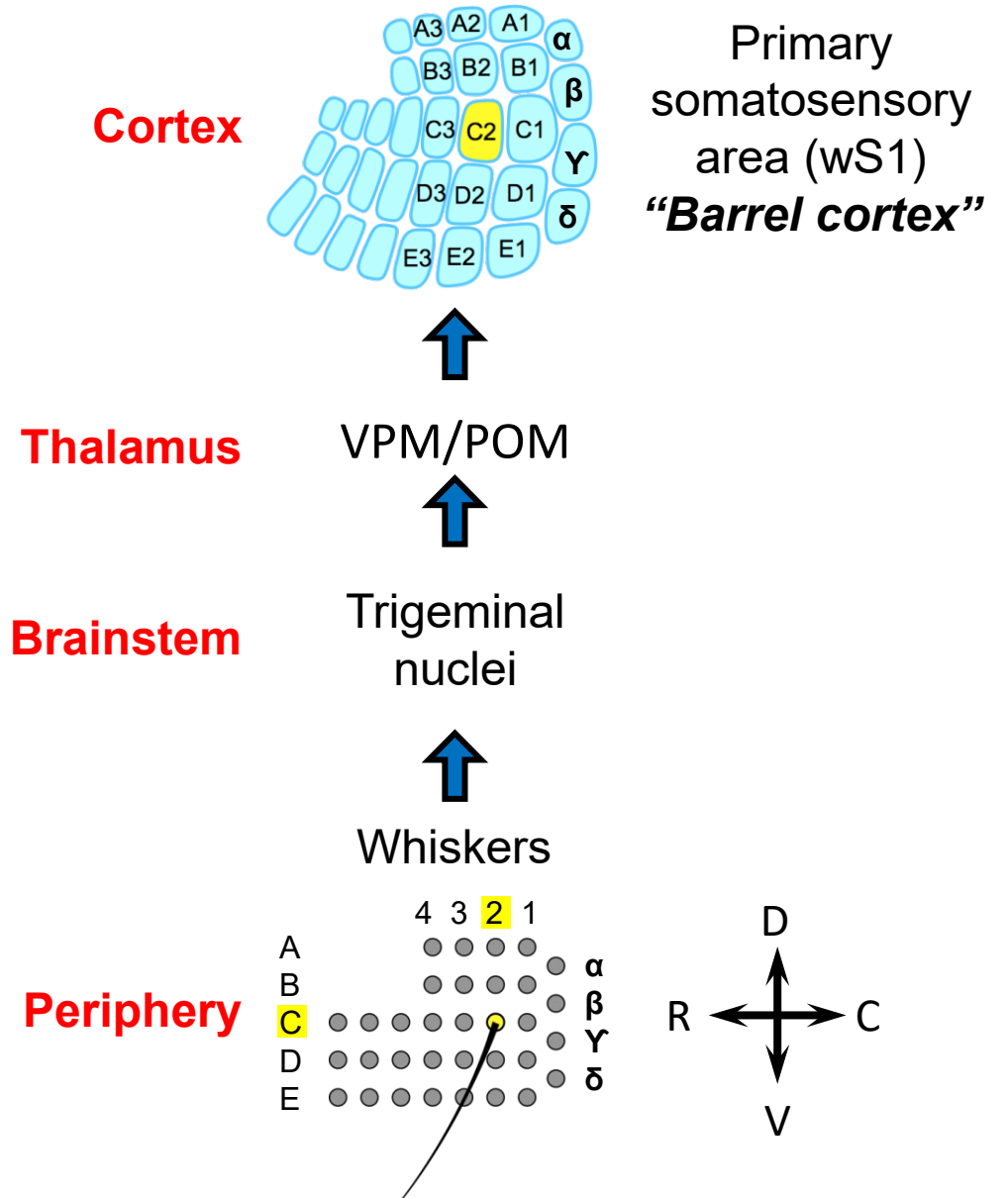
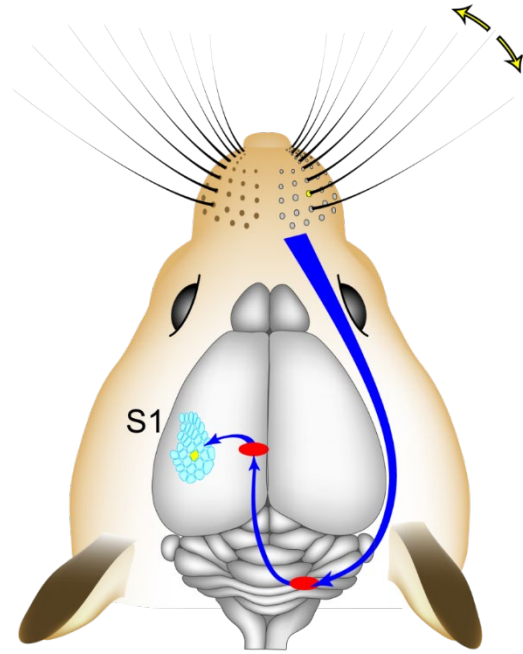
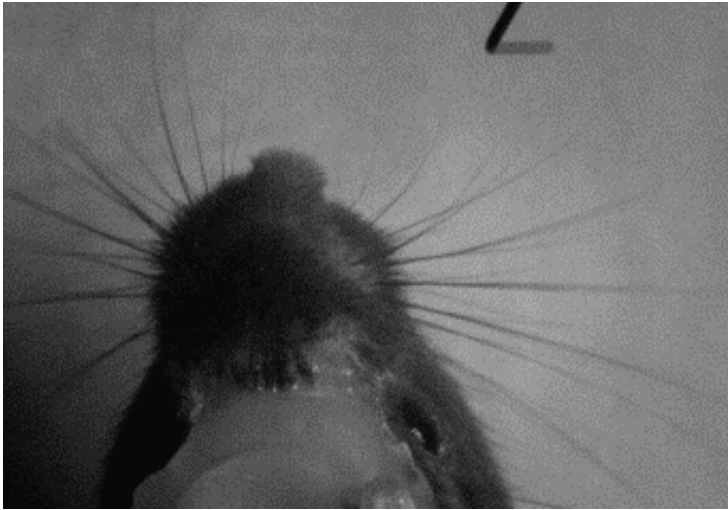
Cortical states are usually defined by the spontaneous activity of the cortical network



(Hans Berger, 1929)

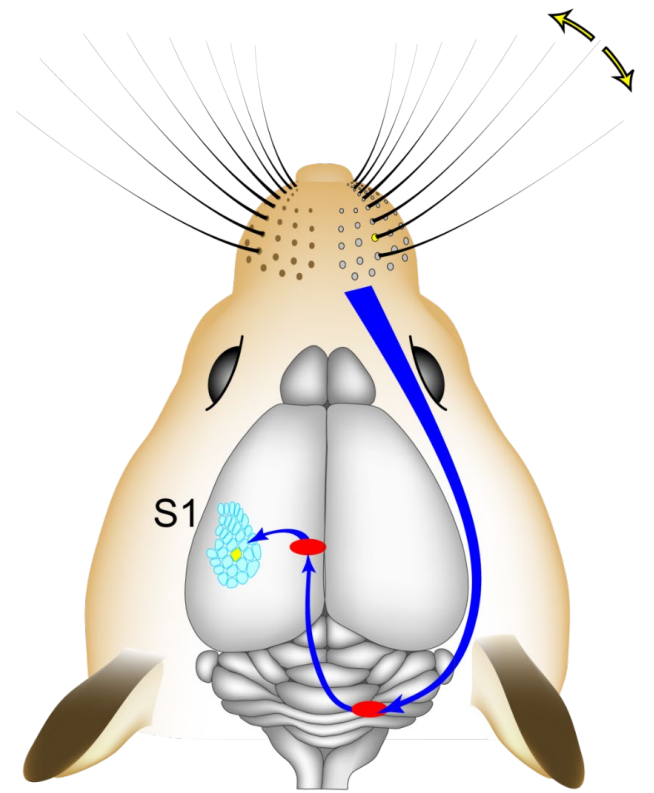
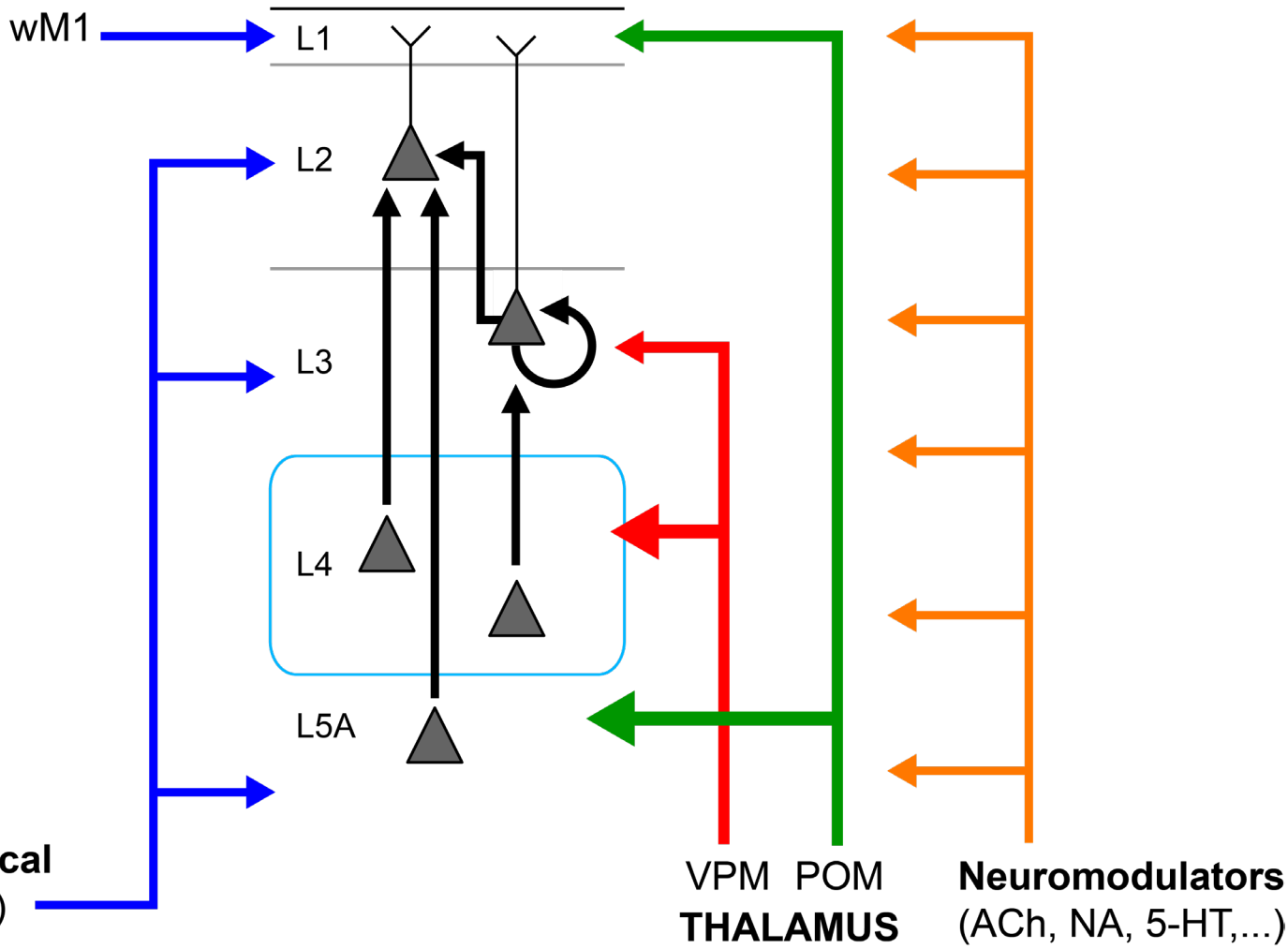
First demonstration of cortical state change correlated with behavior

Experimental model

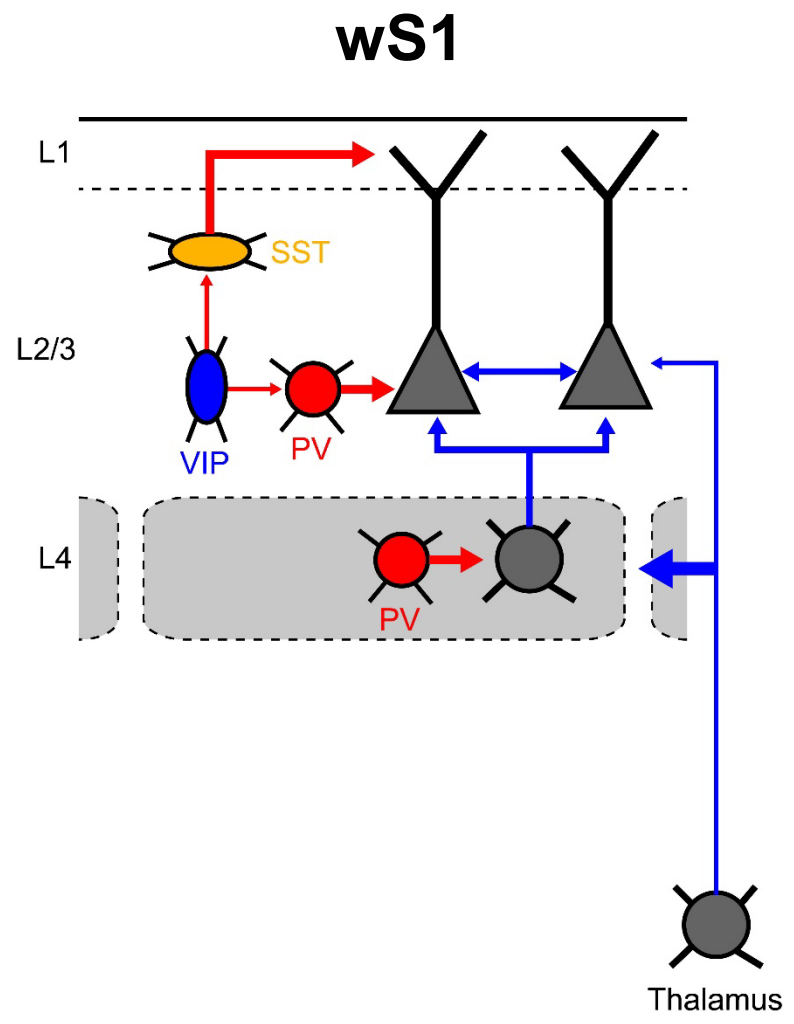
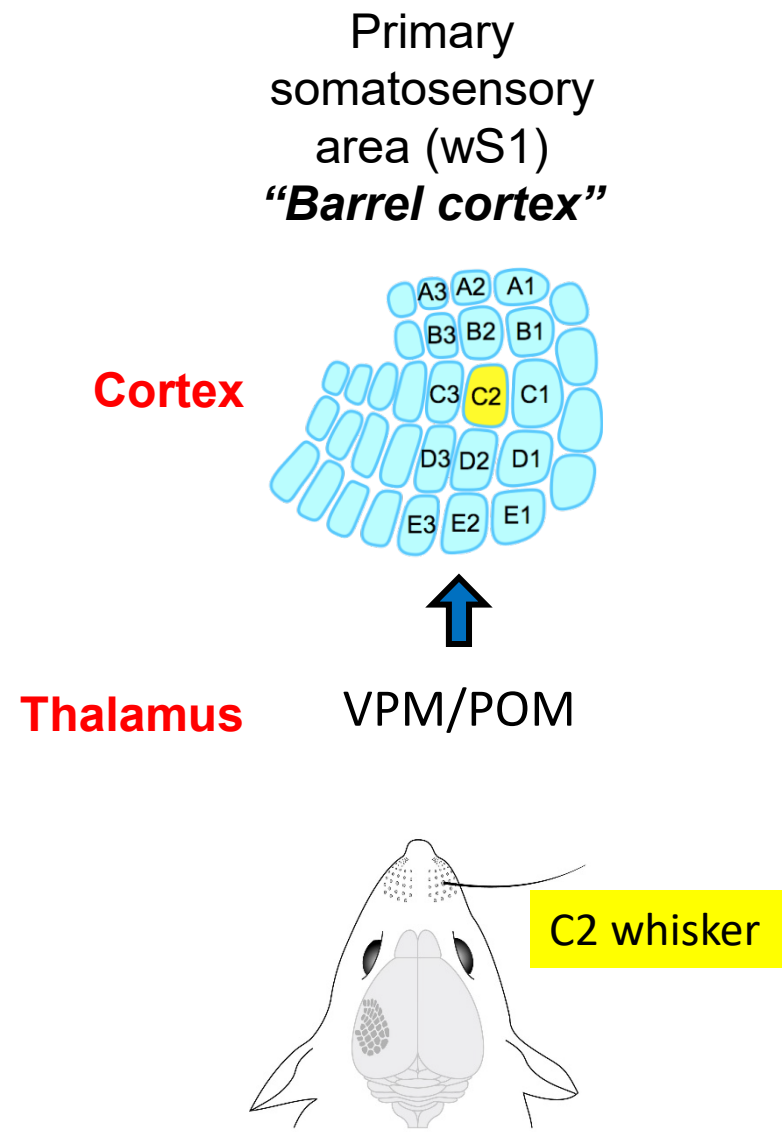


Synaptic inputs to L2/3 of the *barrel* cortex (wS1)

Whisker primary somatosensory cortex (wS1)

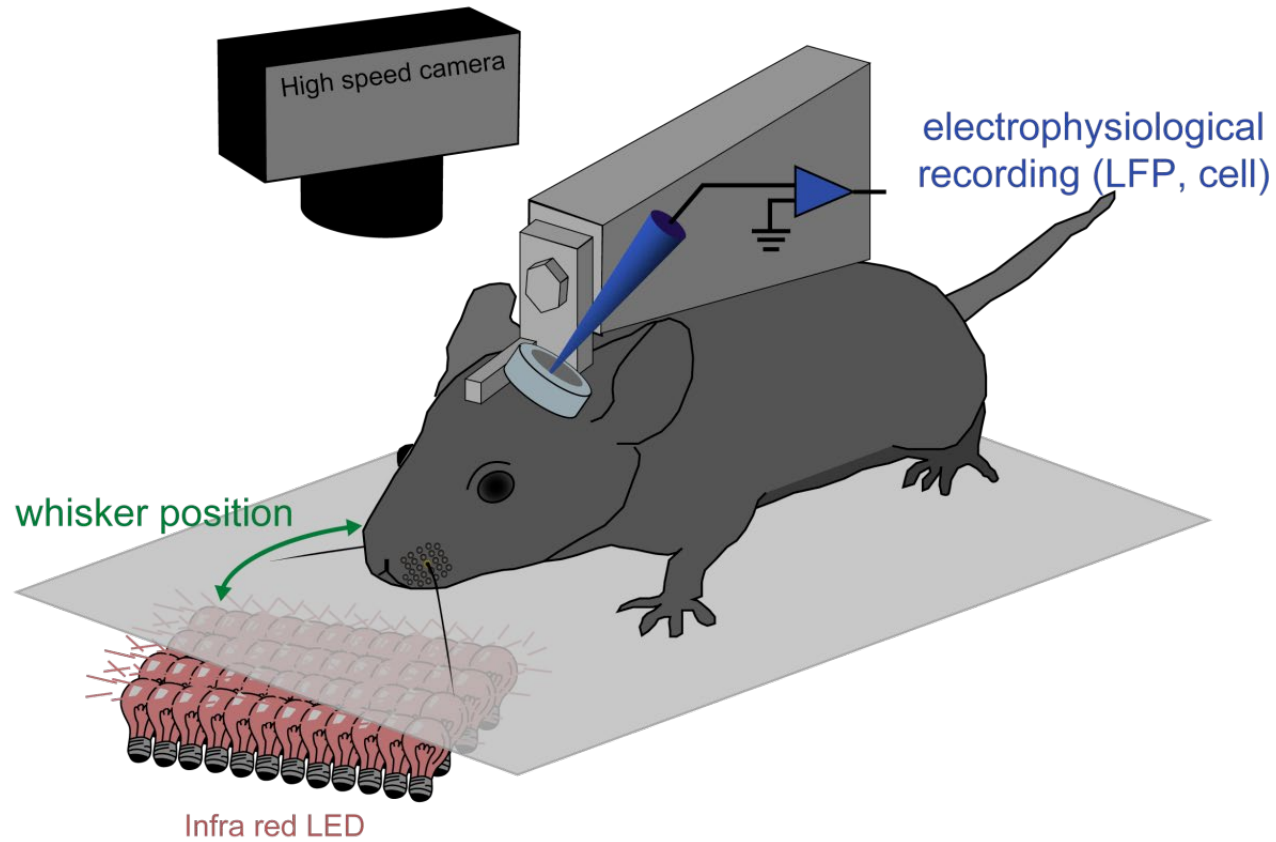


Local neuronal circuit of L2/3 in wS1

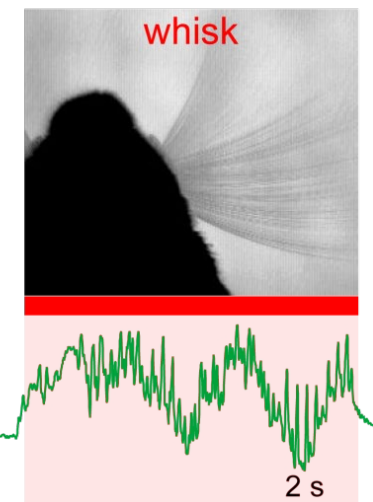
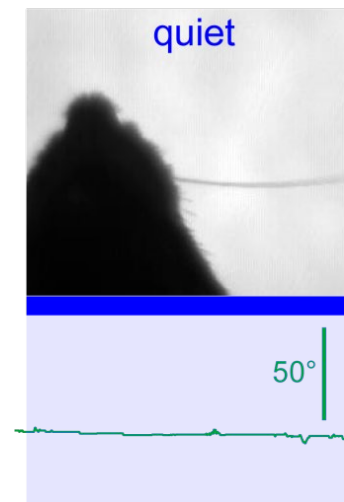
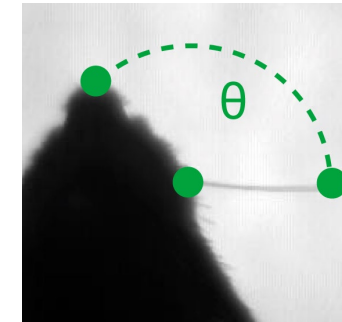


- ▲ ● Excitatory cells (Glutamate)
- Inhibitory interneurons (GABA):
 - Parvalbumine (PV)
 - Vasointestinal peptide (VIP)
 - Somatostatin (SST)

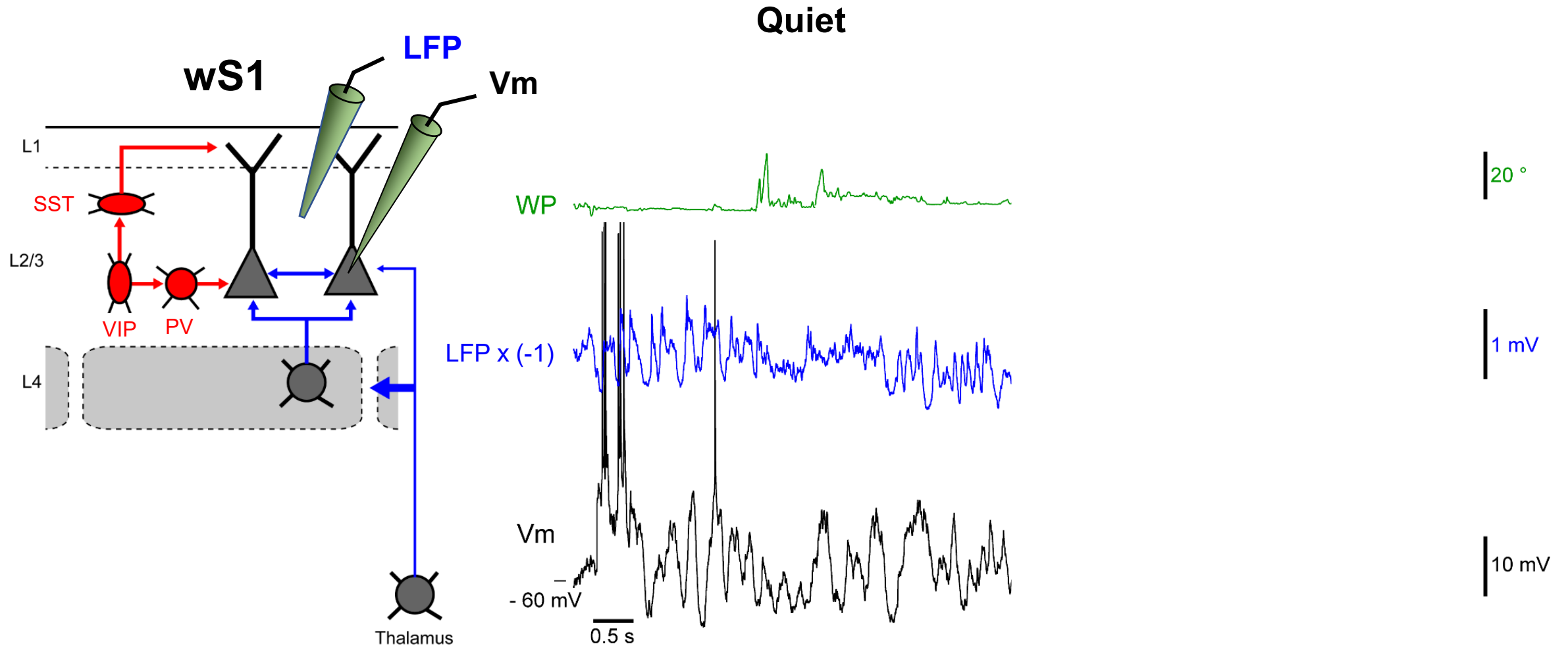
Membrane potential (V_m) recording in awake mice



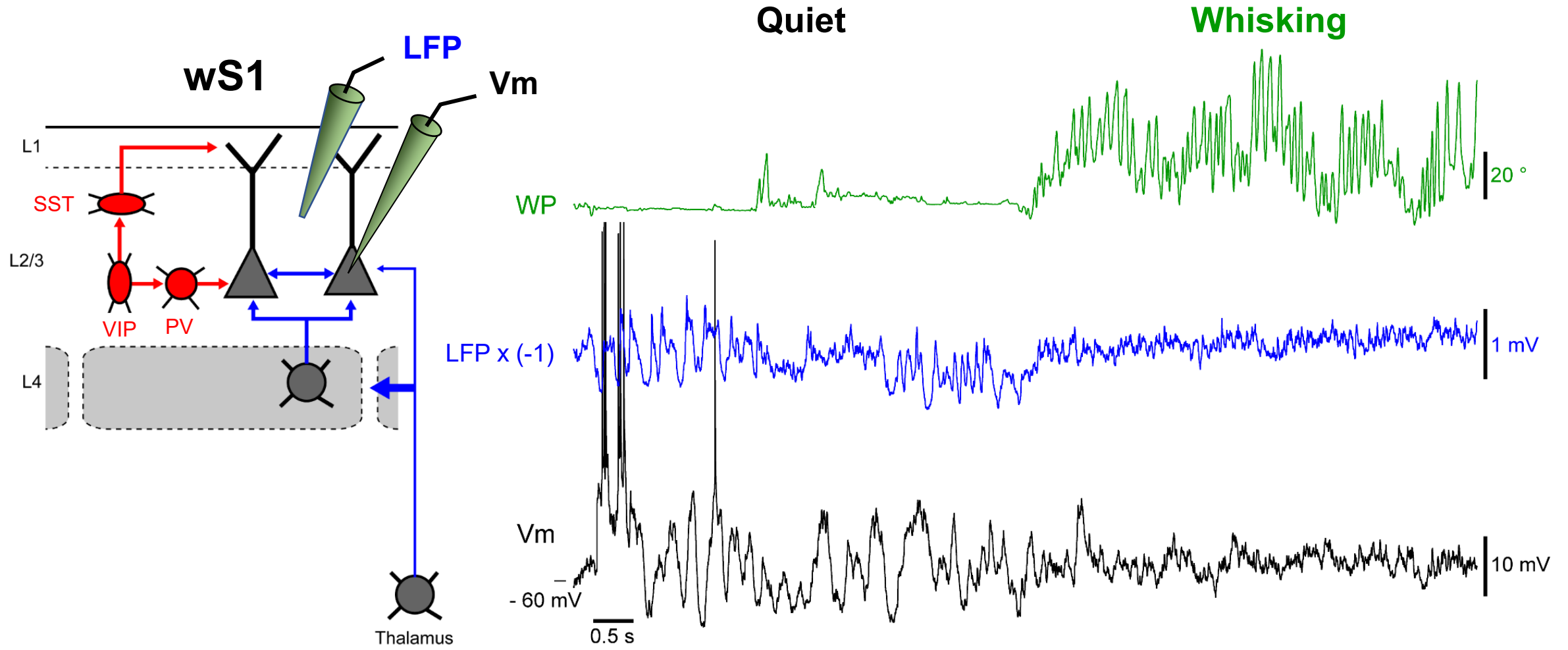
Whisker
Position (WP)



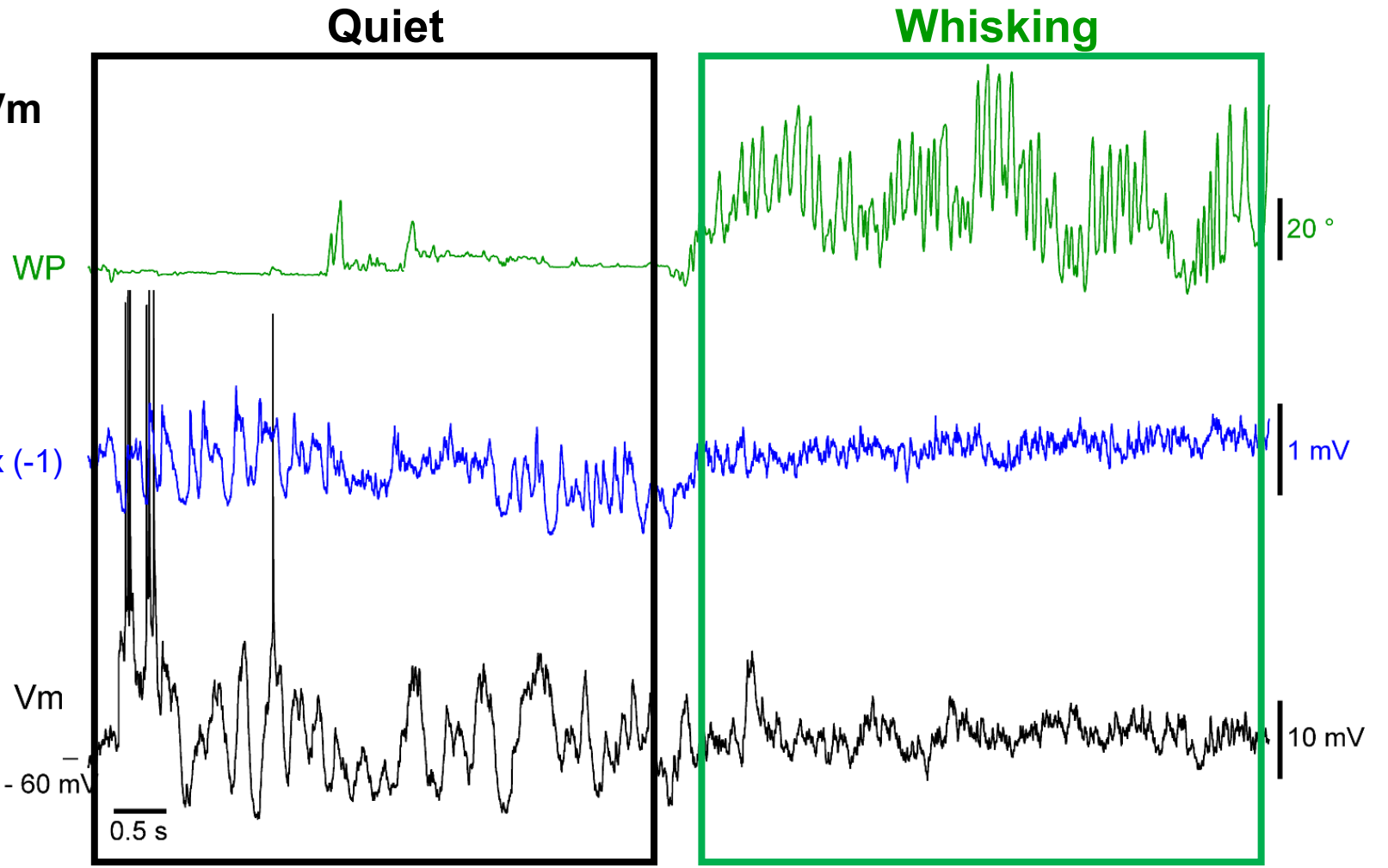
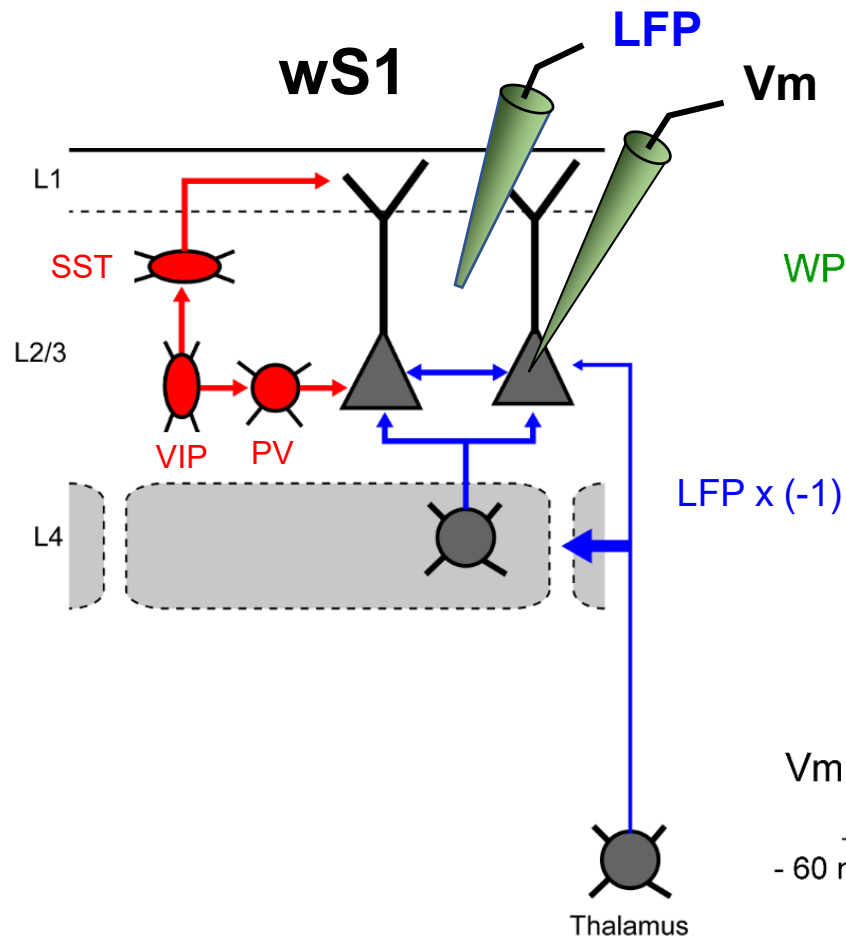
State change in wS1



State change in wS1

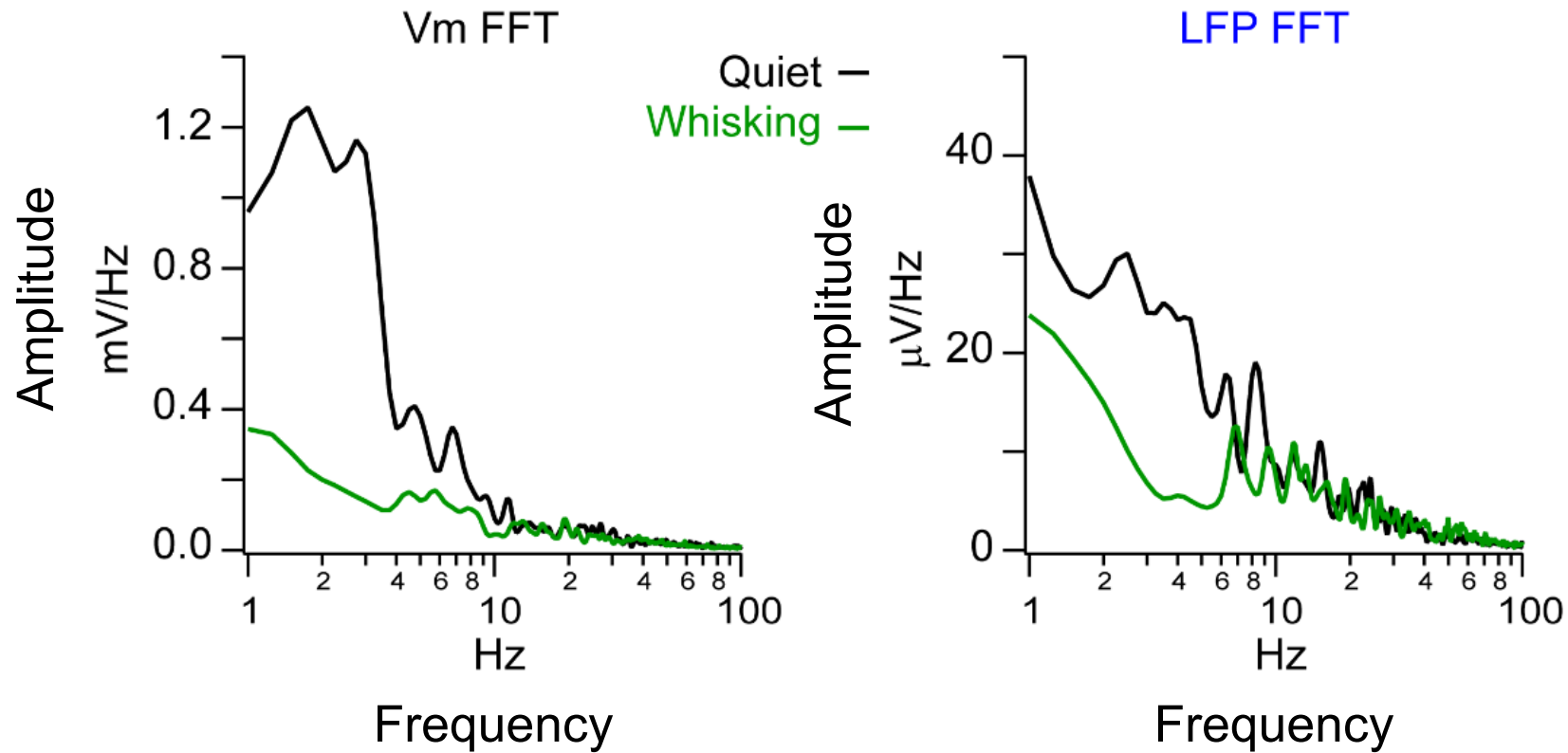


State change in wS1



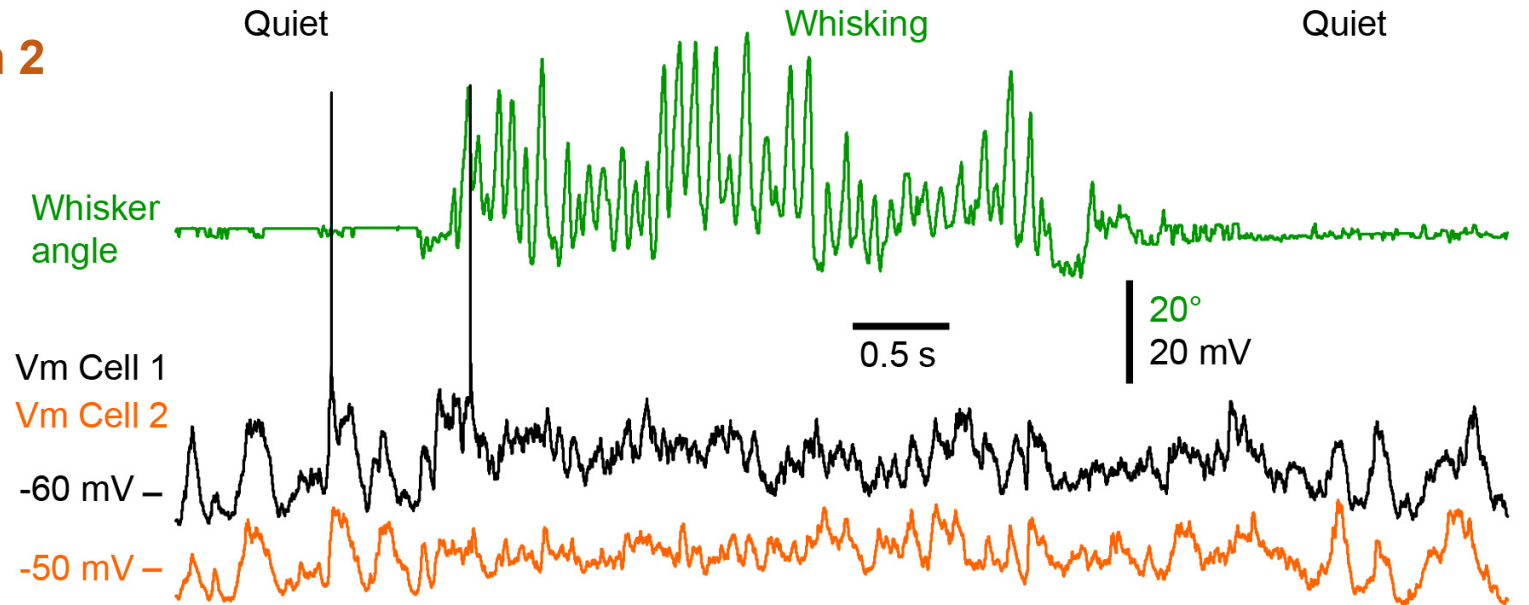
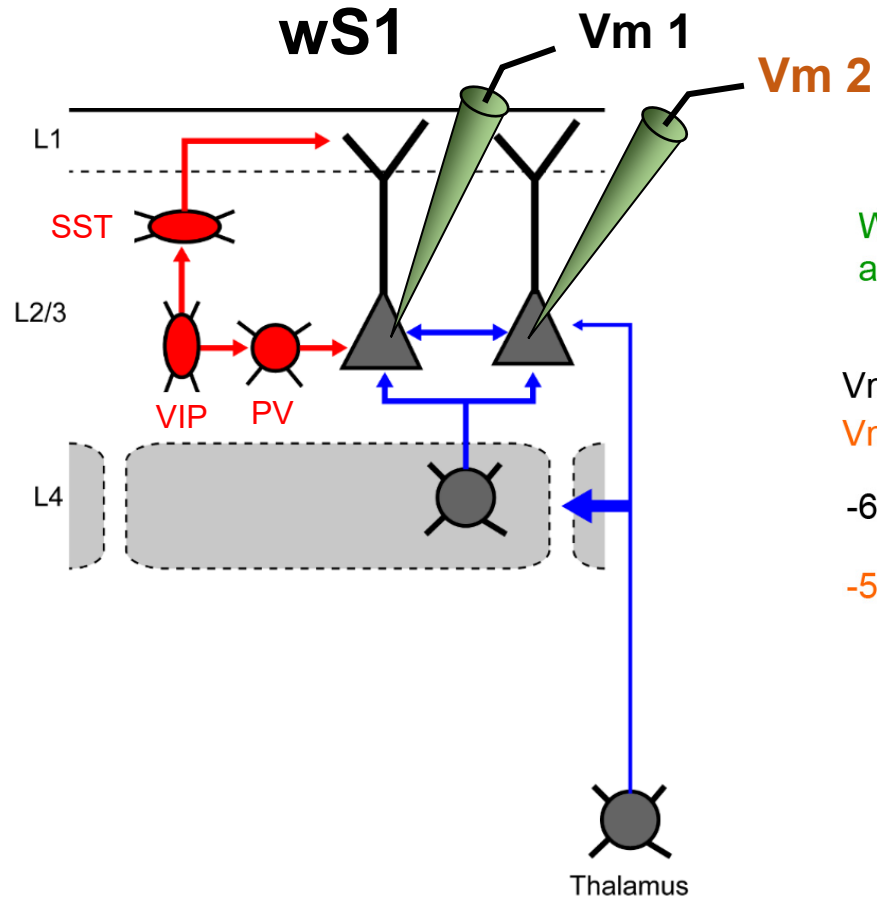
State change in wS1

Fast-Fourier Transform => frequency domain

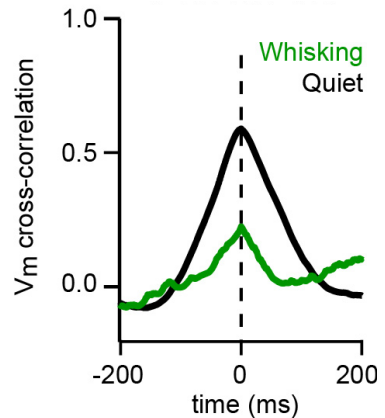


=> Decrease in low-frequency (1-10 Hz) activity during whisking

■ State change in wS1



Vm-Vm cross-correlograms



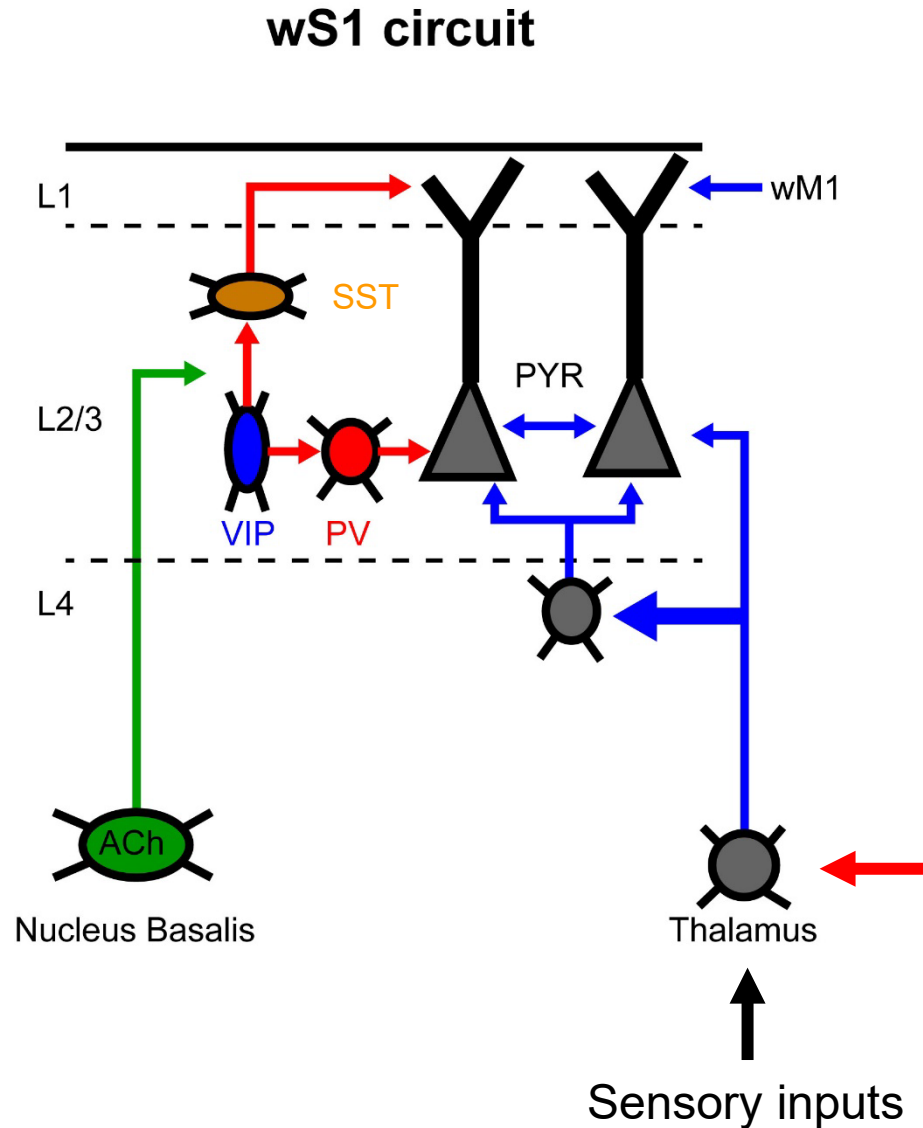
(Poulet and Petersen, Nature 2008)

- **State change in wS1**

Quiet => **Whisking** :

- **Strong decrease in low-frequency (1-10 Hz) Vm fluctuations**
- **Strong decrease in Vm variance**
- **Decrease in Vm correlation in nearby neurons (*desynchronization*)**
- **Slight but significant depolarization**
- **No significant change in firing rate at population level**

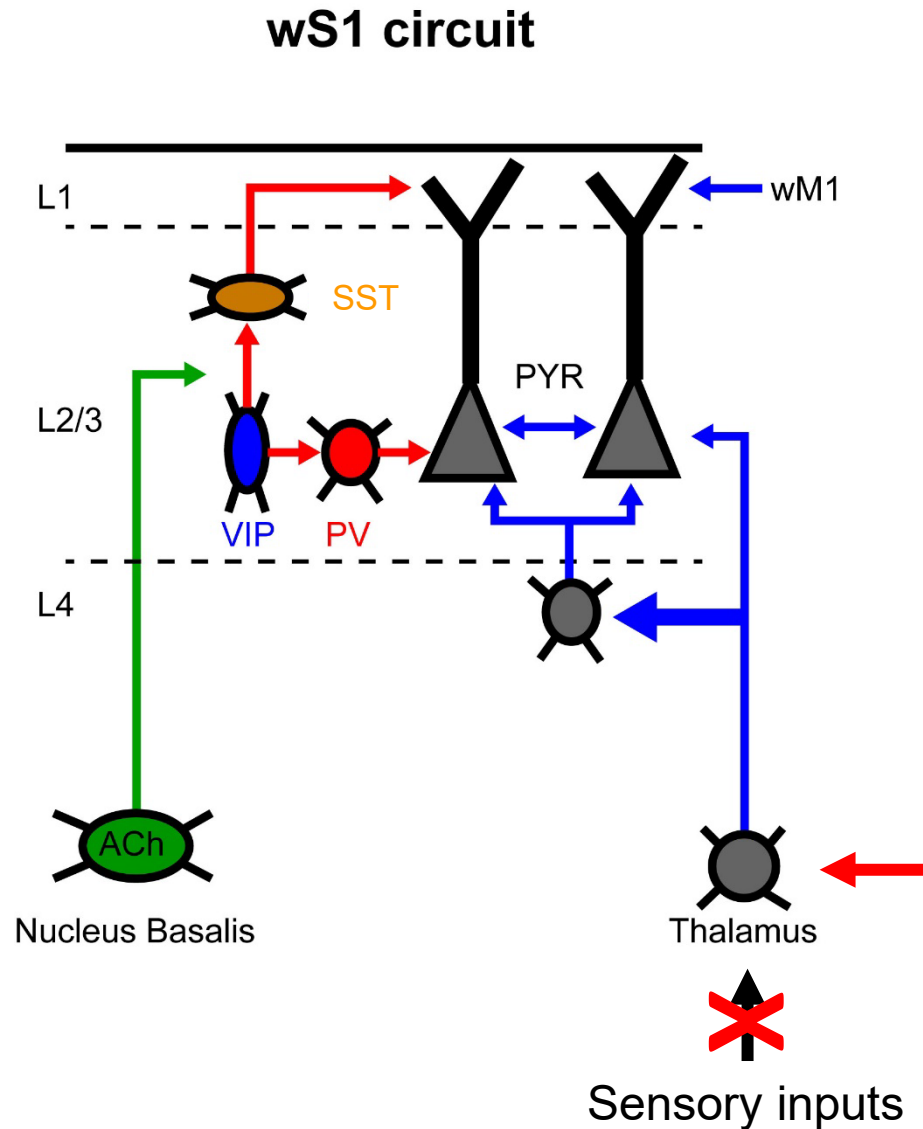
- The origins of state change in wS1



What are the possible origins of the state change?

Modulatory inputs
Brainstem (ACh, NA ...)

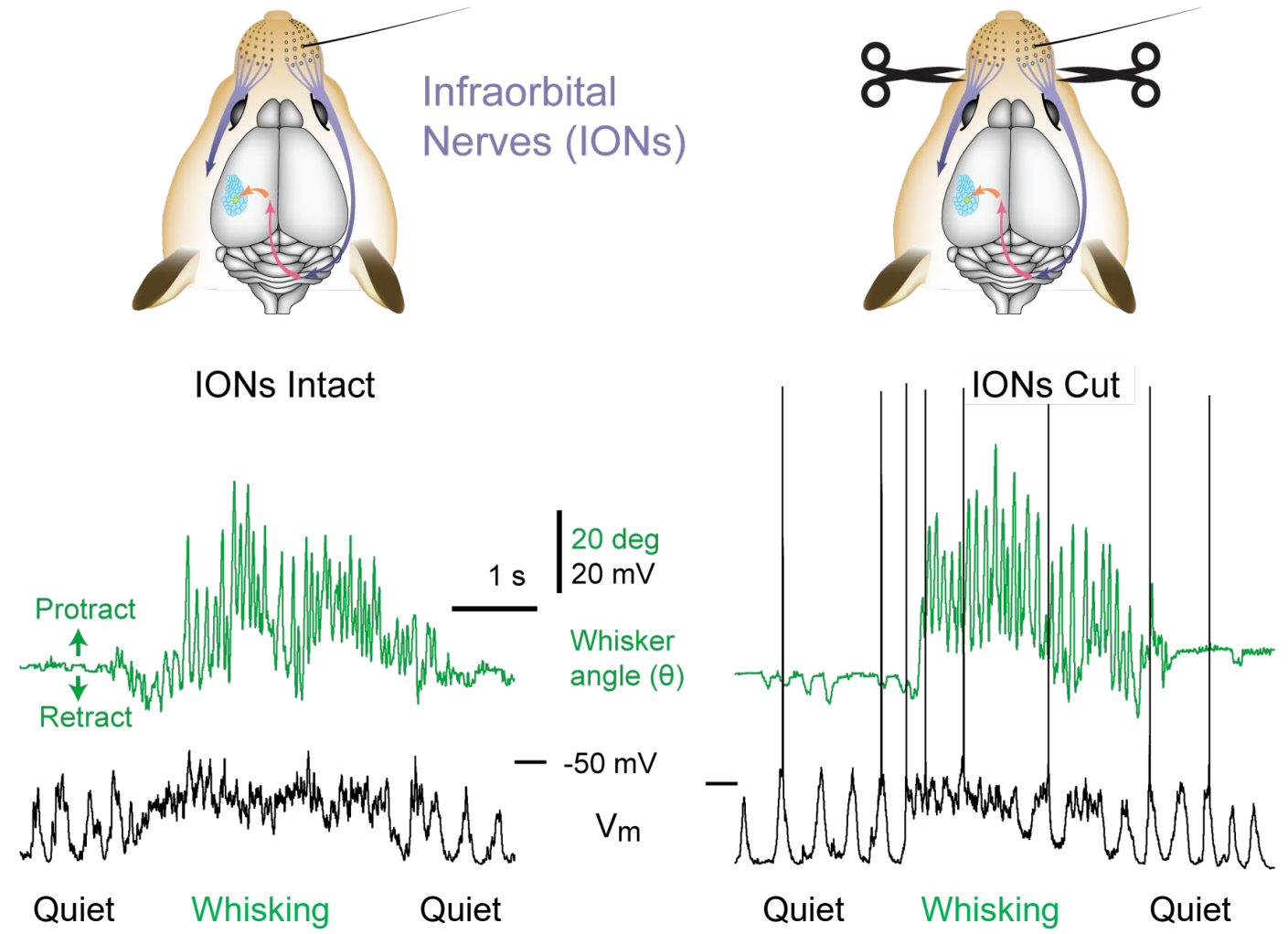
■ The origins of state change in wS1



What is the contribution of sensory inputs?

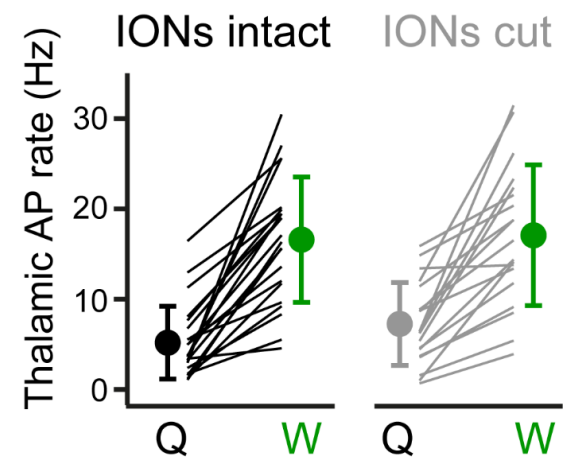
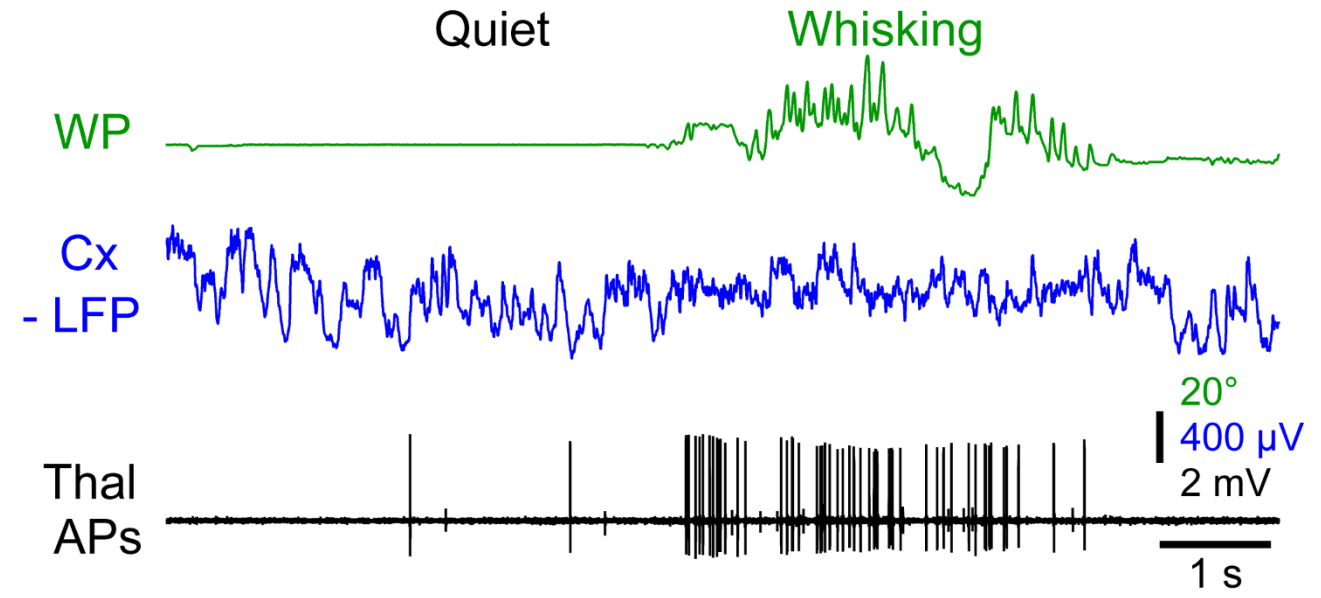
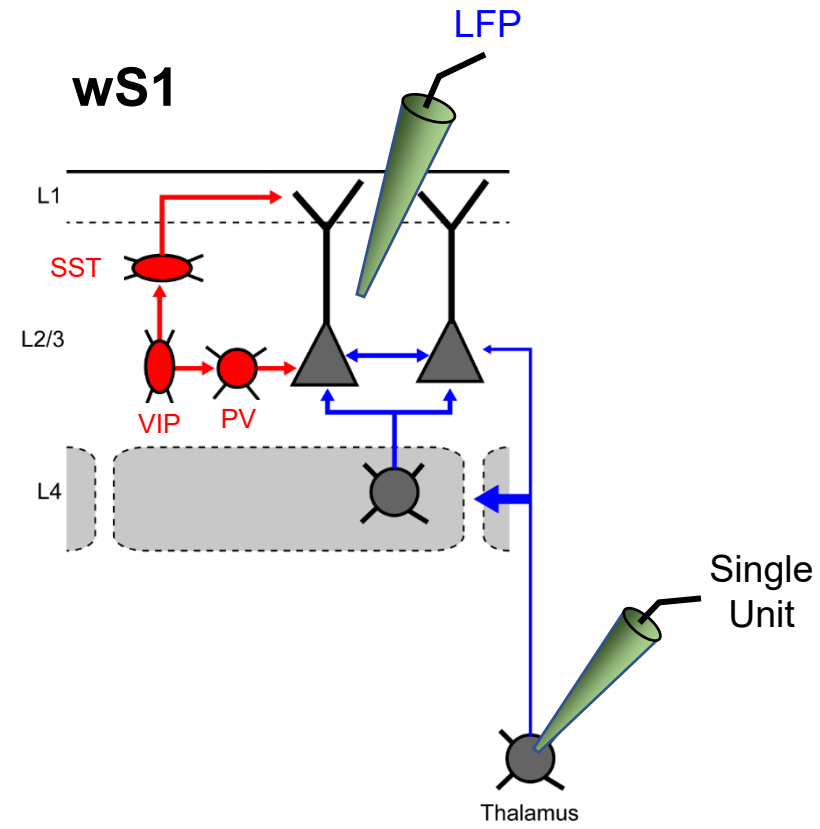
Modulatory inputs
Brainstem (ACh, NA ...)

State change in wS1 is centrally generated



(Poulet and Petersen, Nature 2008; Poulet, Fernandez, Crochet et al., Nat. Neurosci. 2012)

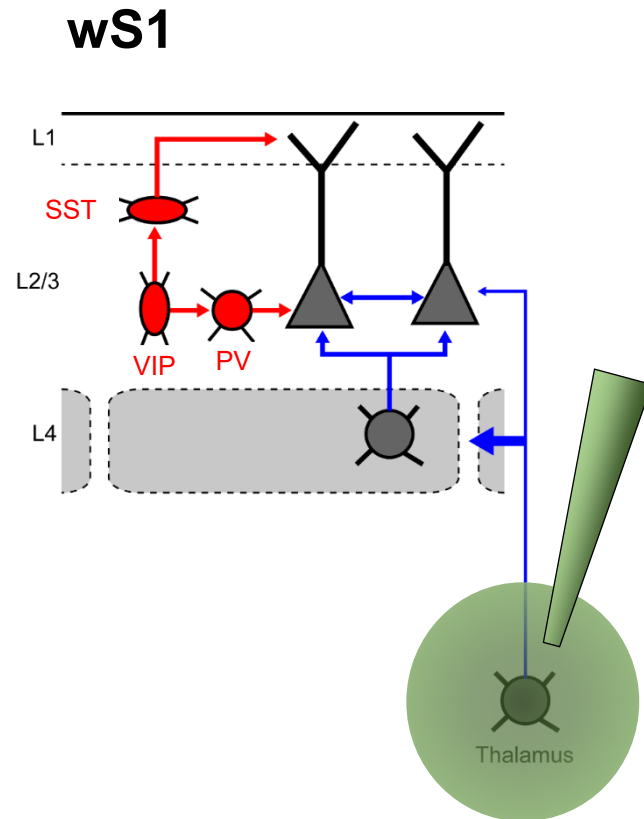
Thalamic contribution



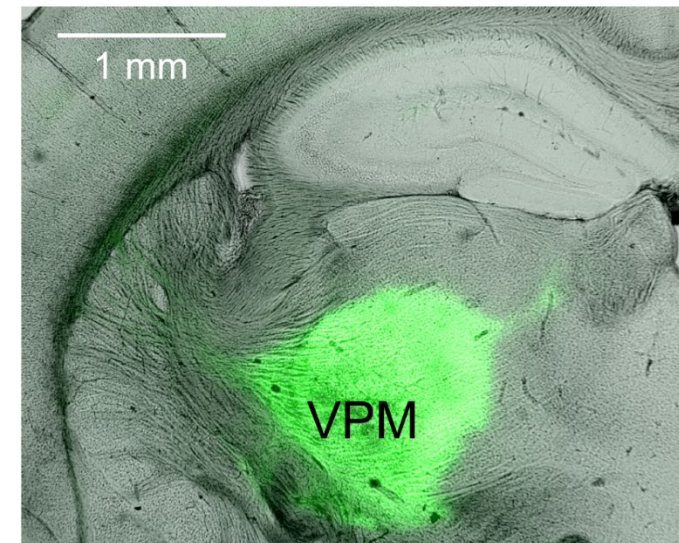
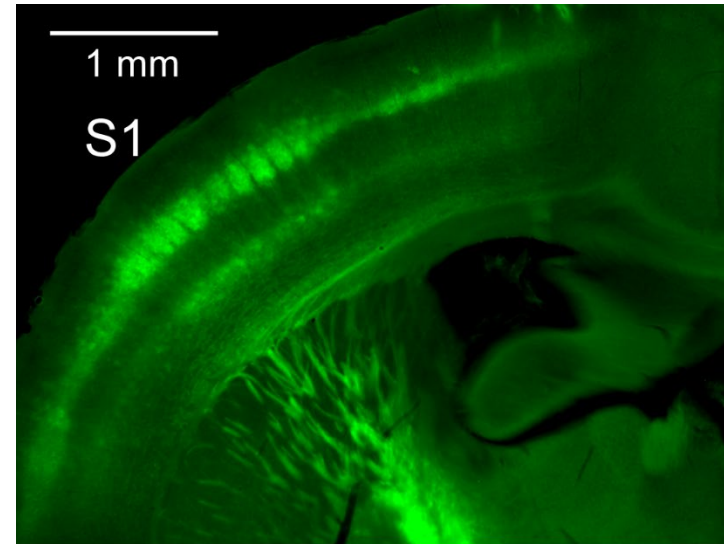
=> Activity of thalamic neurons increases with whisking and correlates with state change in wS1

■ Thalamic contribution

Optogenetic stimulation of thalamic neurons

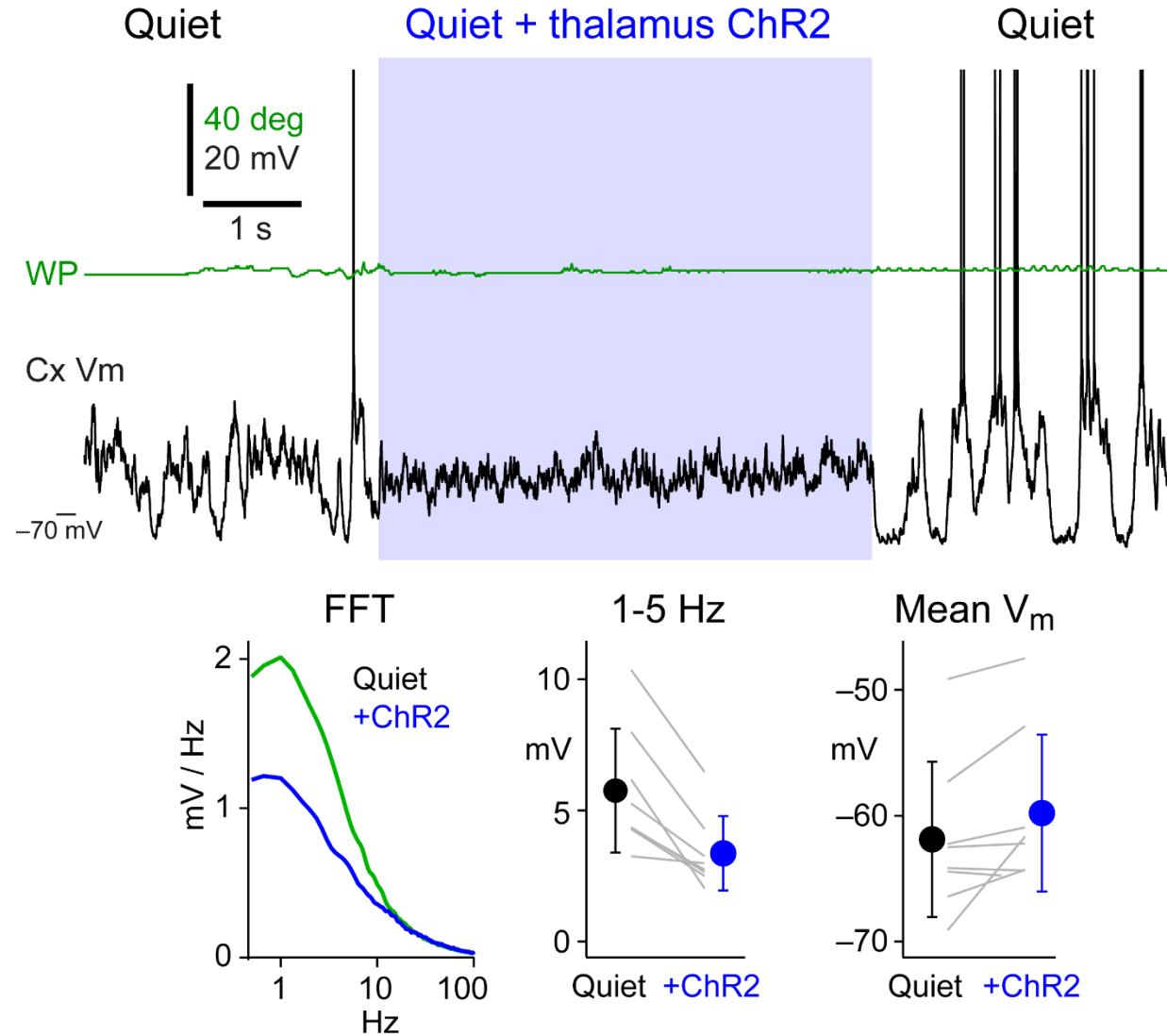
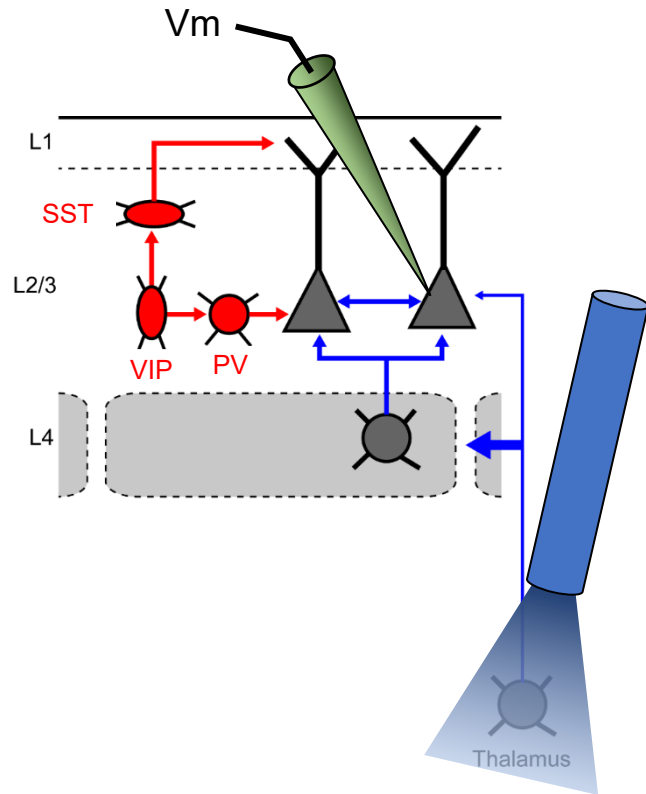


AAV2/1.CAG.ChR2-Venus.W.SV40



Thalamic contribution

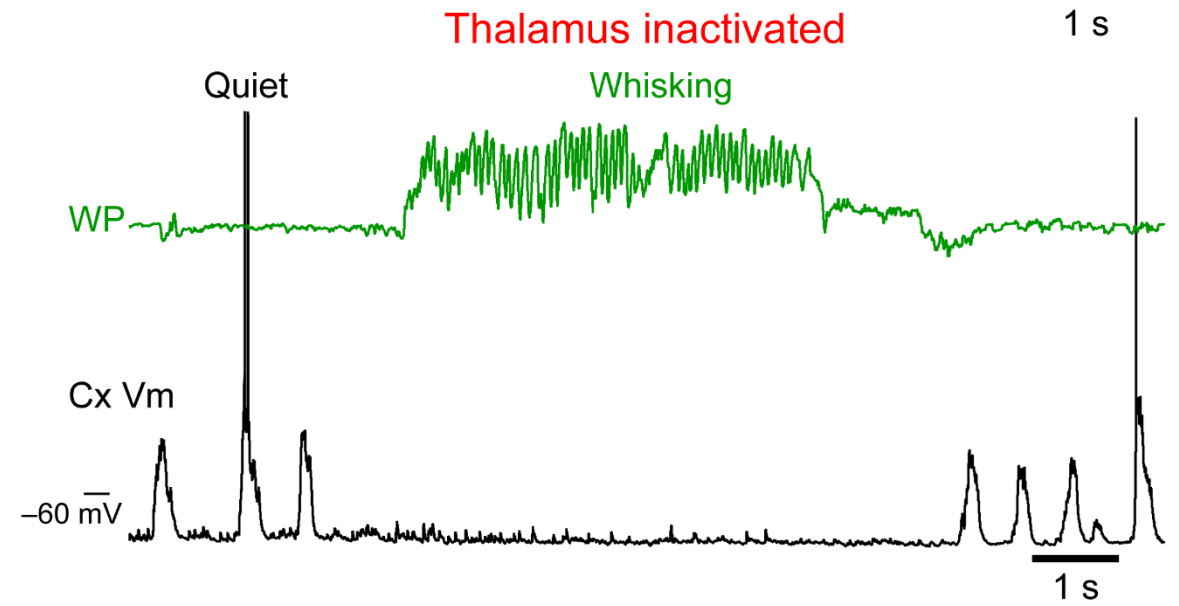
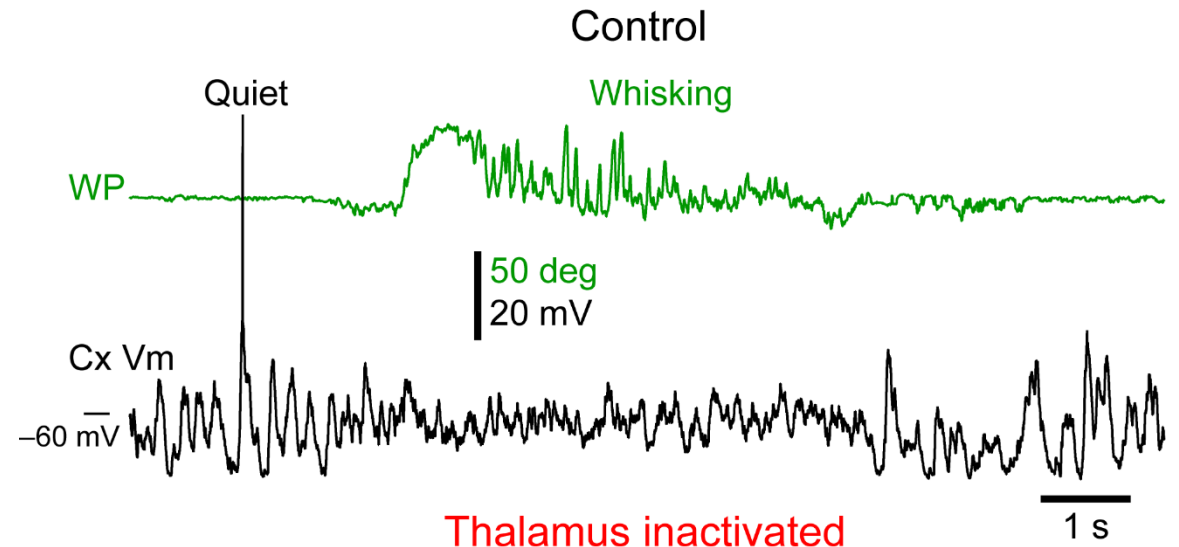
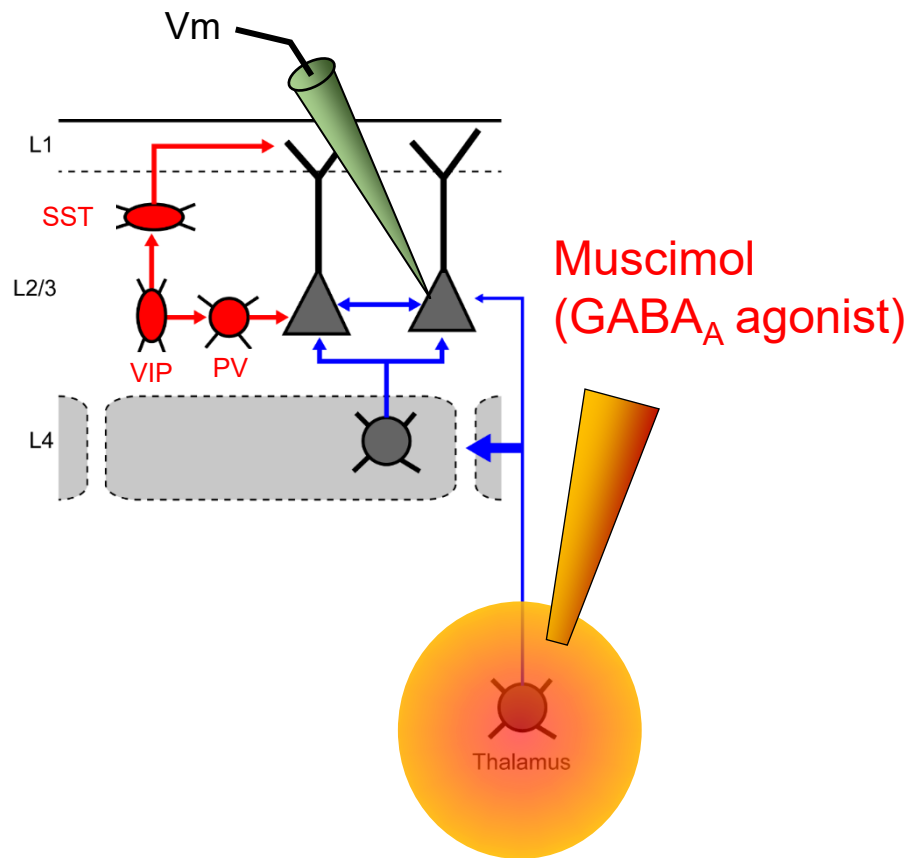
Optogenetic stimulation of thalamic neurons



=> Activation of thalamic neurons induces state change in wS1 during quiet wakefulness

Thalamic contribution

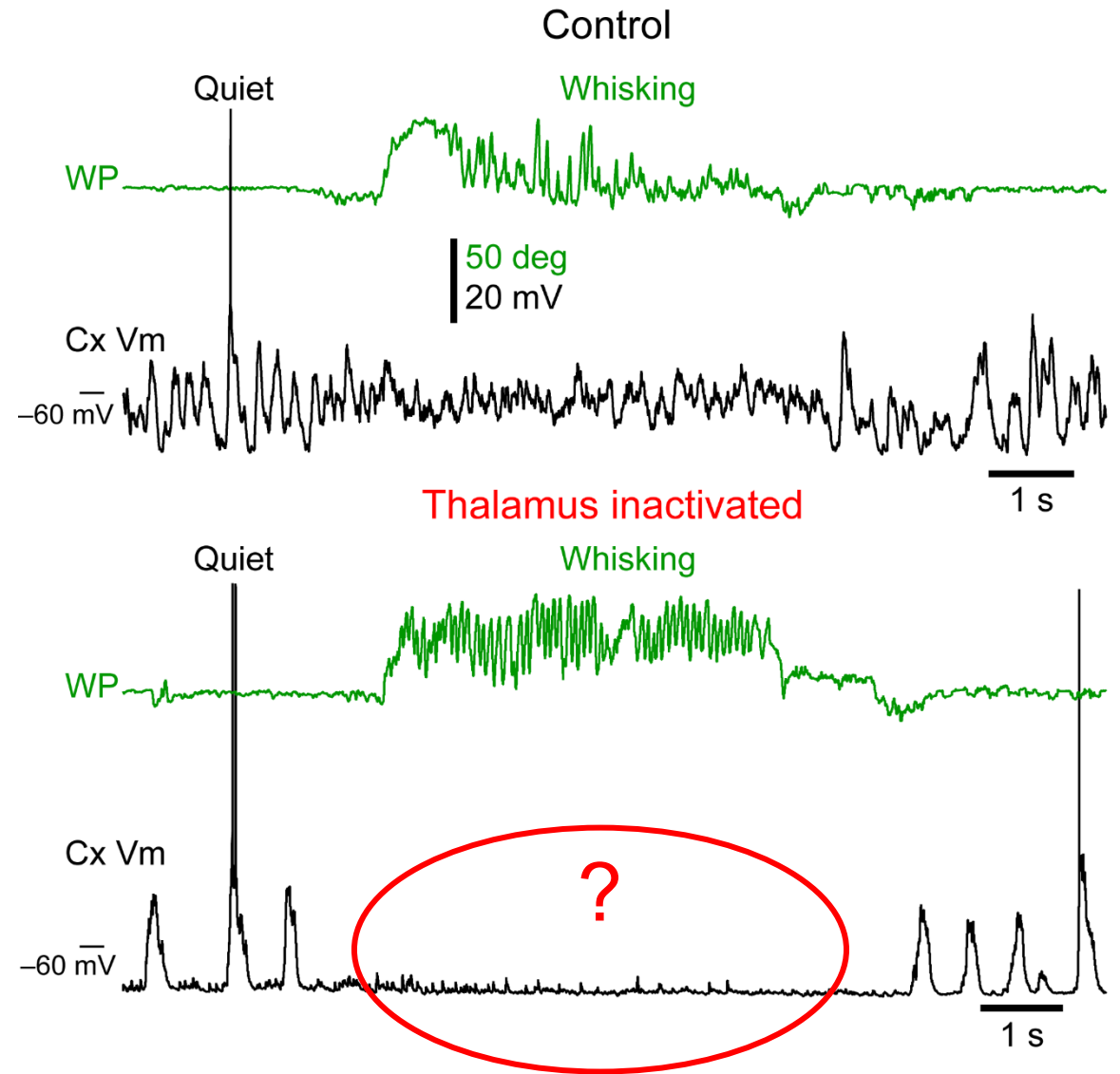
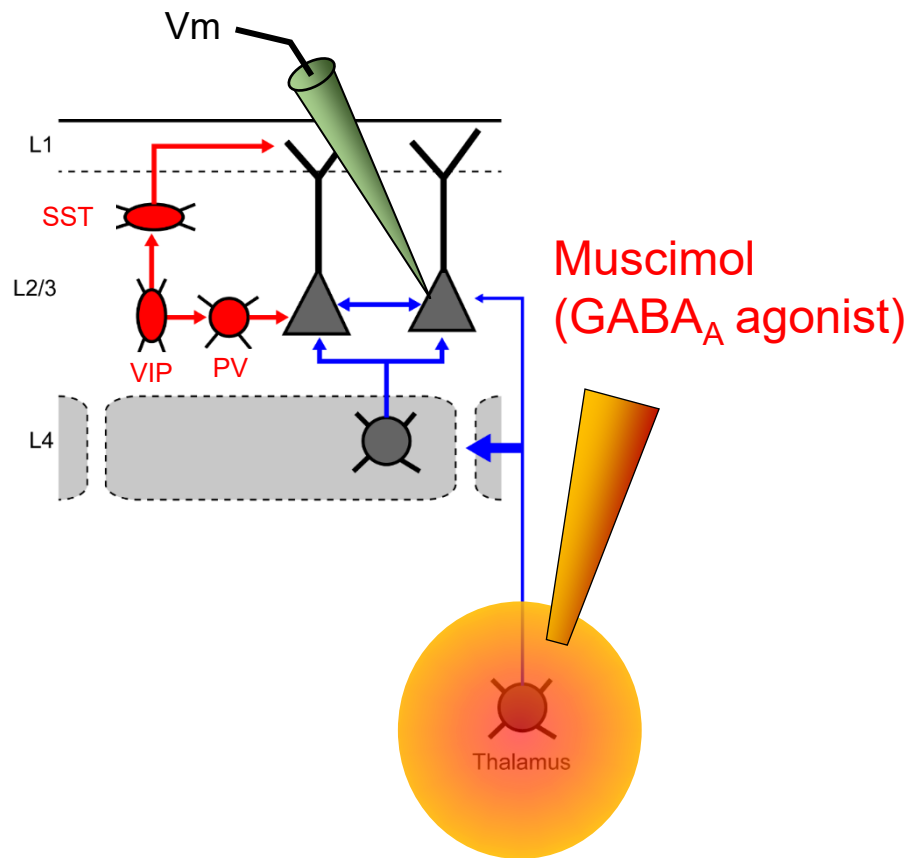
Pharmacological inhibition of thalamic neurons



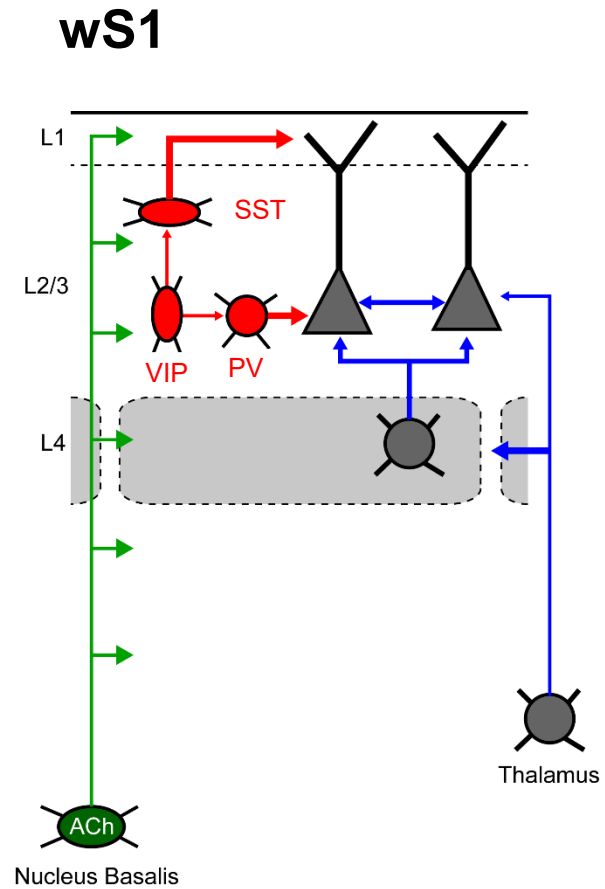
=> Inhibition of thalamic neurons suppresses the high-frequency activity in wS1 during whisking

Thalamic contribution

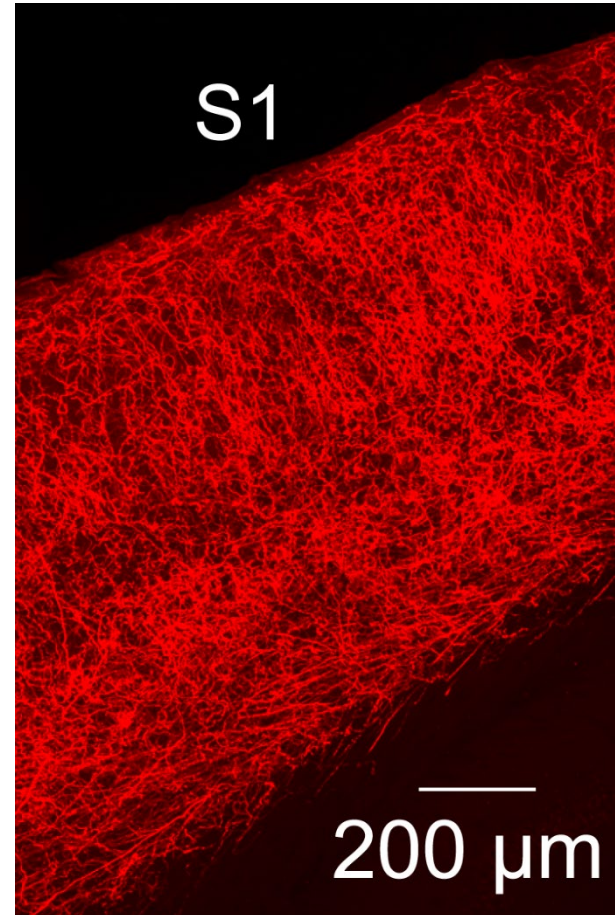
Pharmacological inhibition of thalamic neurons



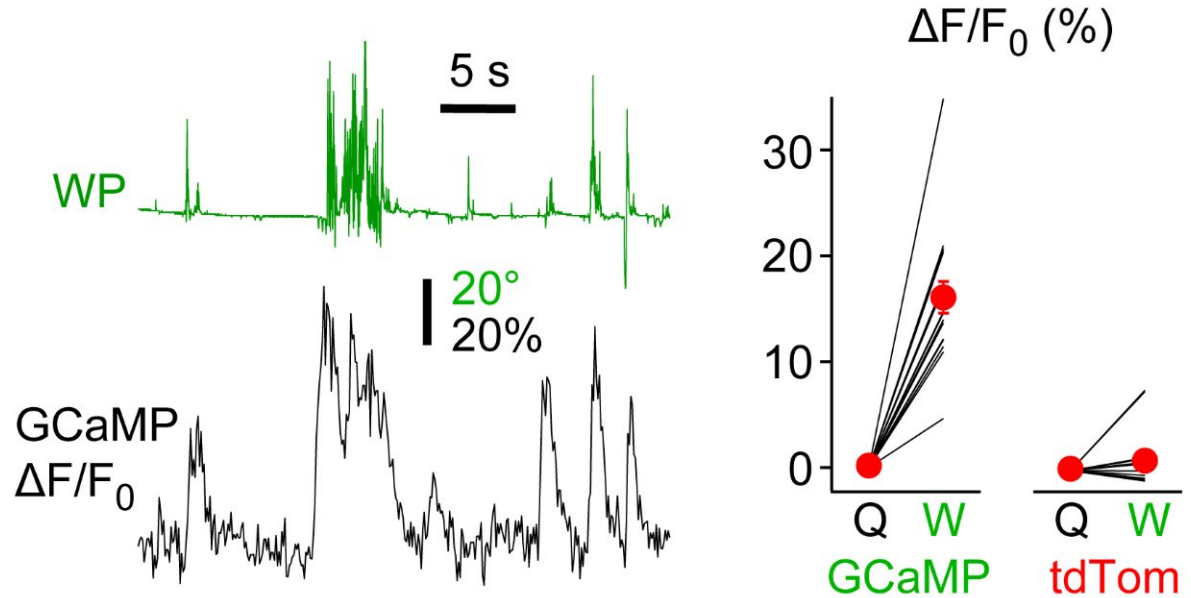
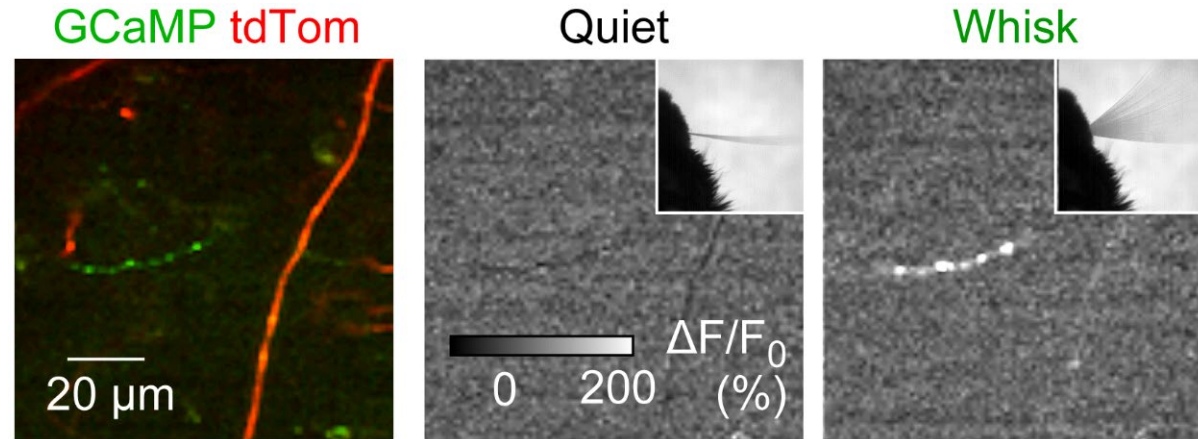
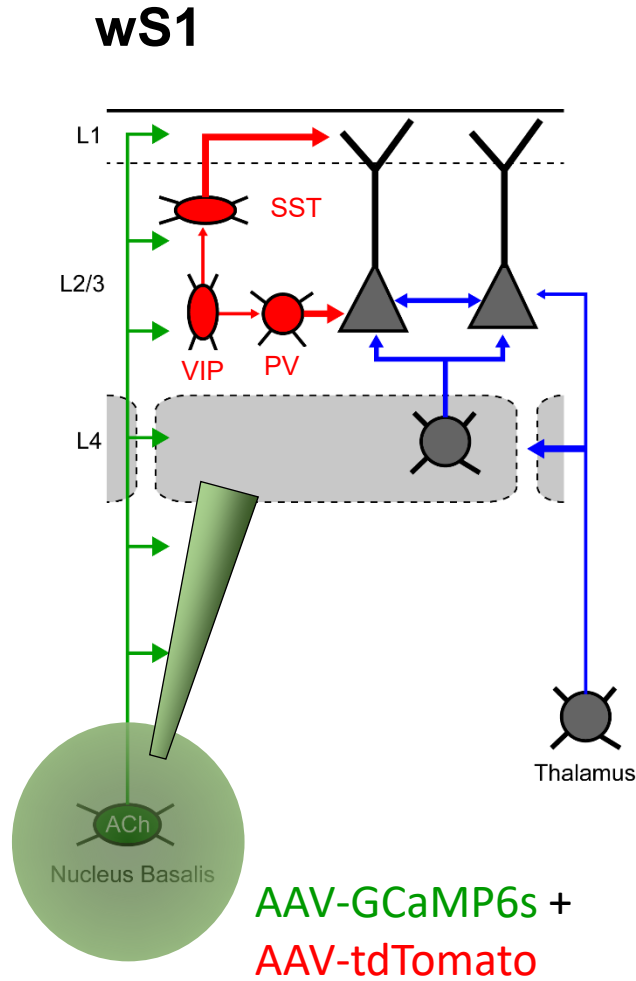
■ Cholinergic contribution



Cholinergic axons in S1 (tdTomato)



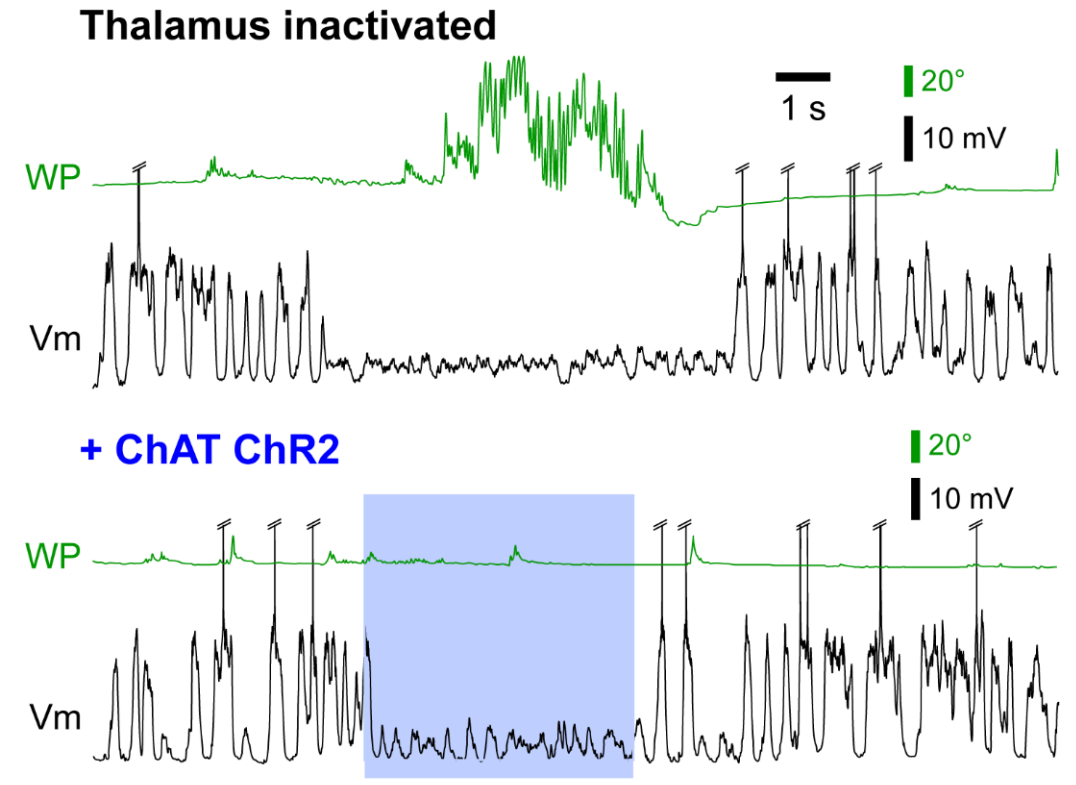
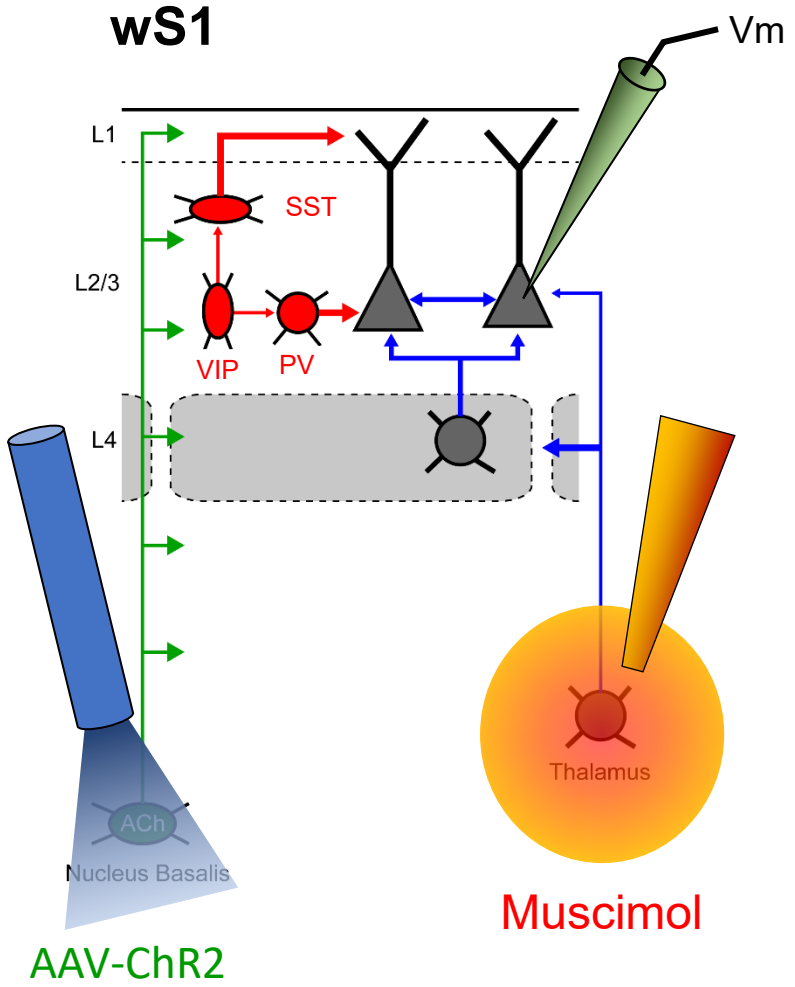
■ Cholinergic contribution



=> ACh axons in wS1 increase activity during whisking

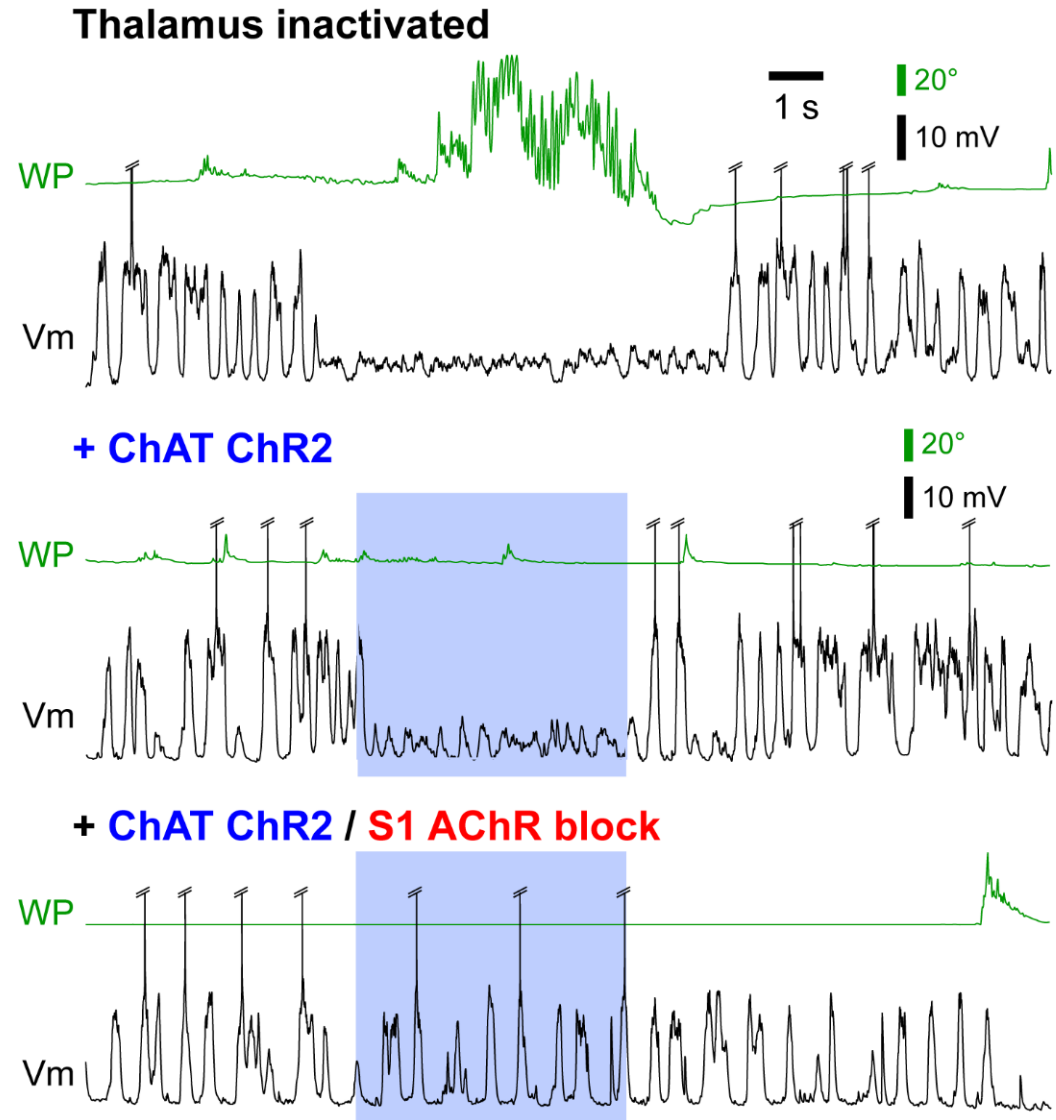
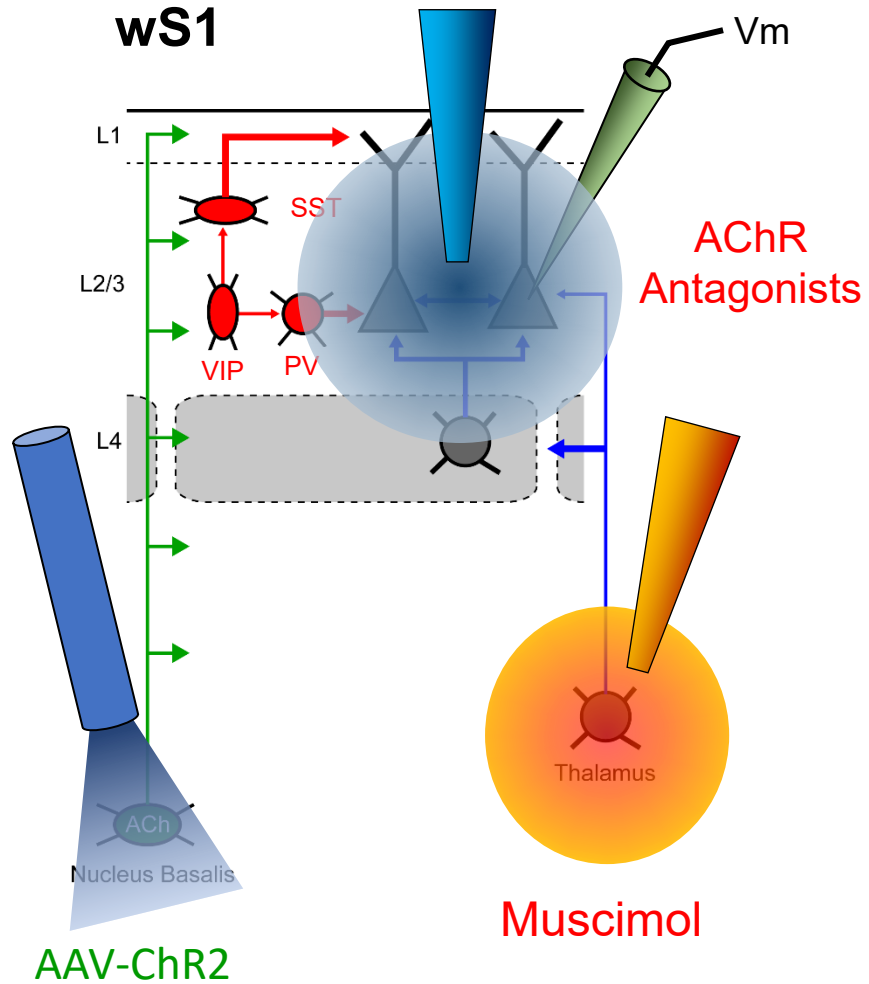
Cholinergic contribution

Optogenetic activation of cholinergic neurons



■ Cholinergic contribution

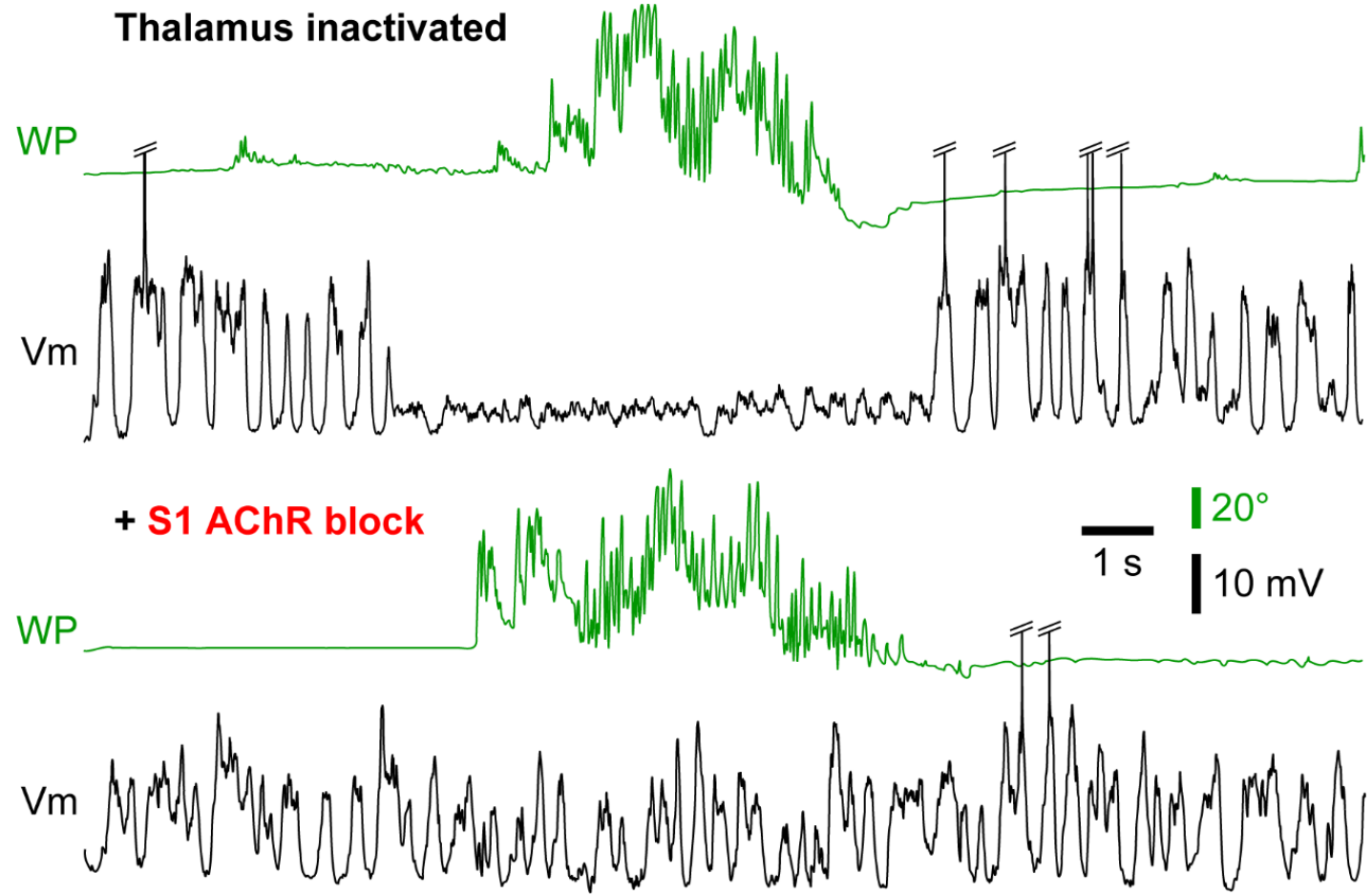
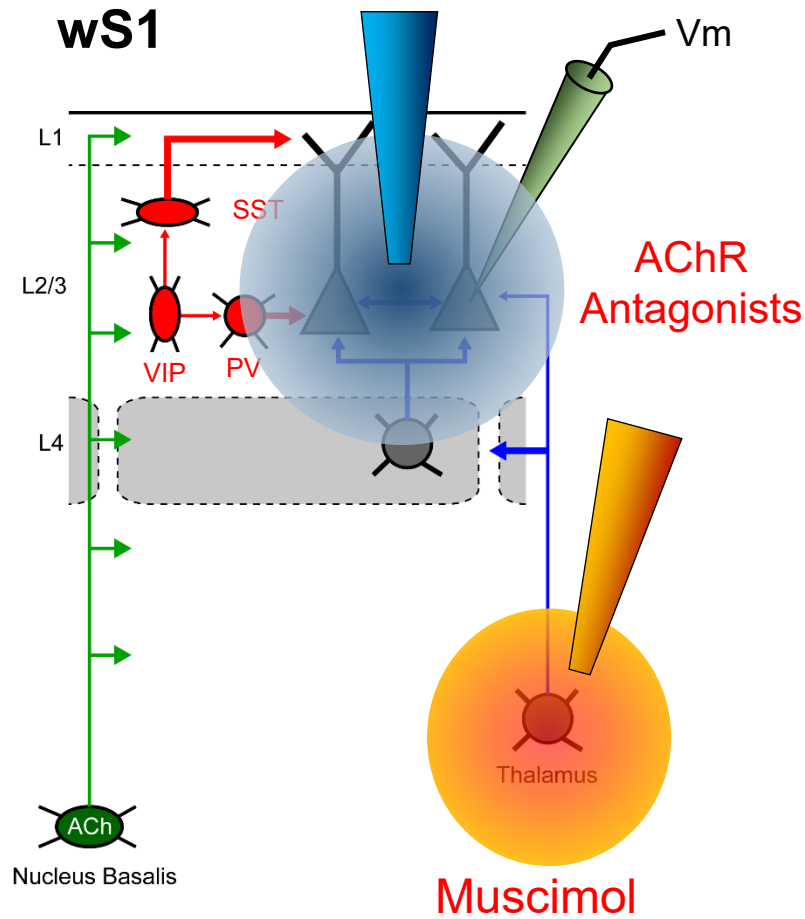
Pharmacological blockade of ACh receptors



=> Activation of ACh neurons induces state change in wS1

■ Cholinergic contribution

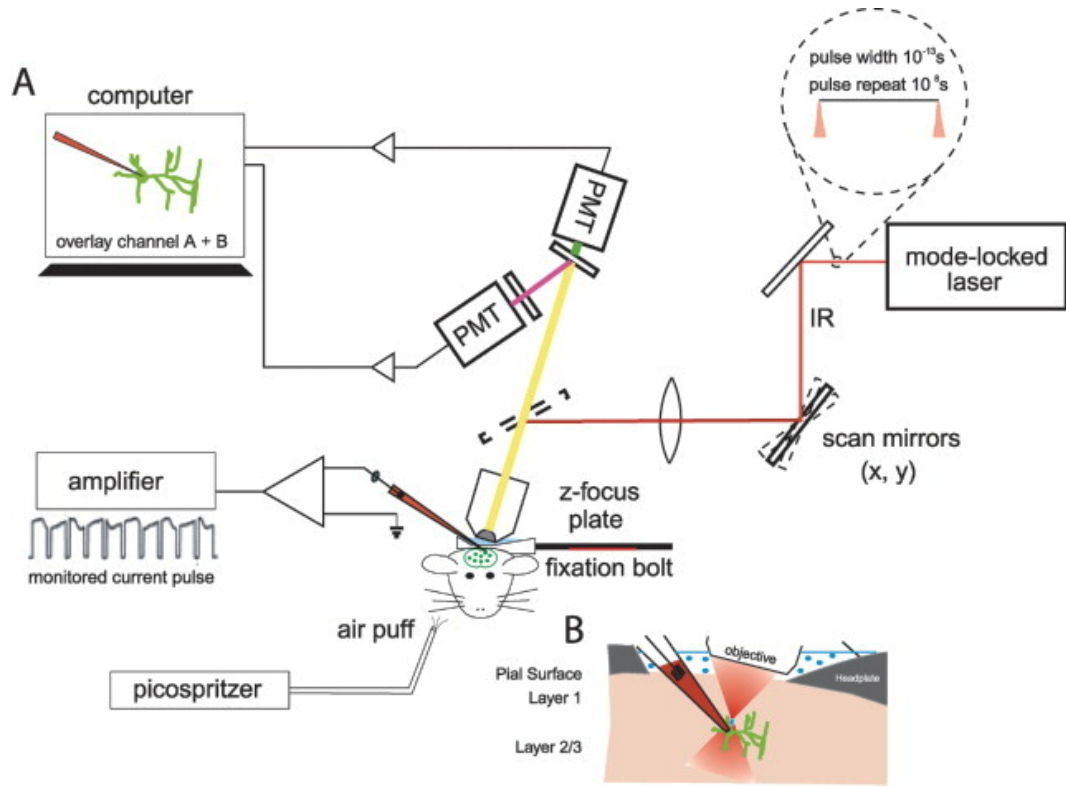
Pharmacological blockade of ACh receptors



=> Blockage of AChR in wS1 & inhibition of thalamus block the state change in wS1

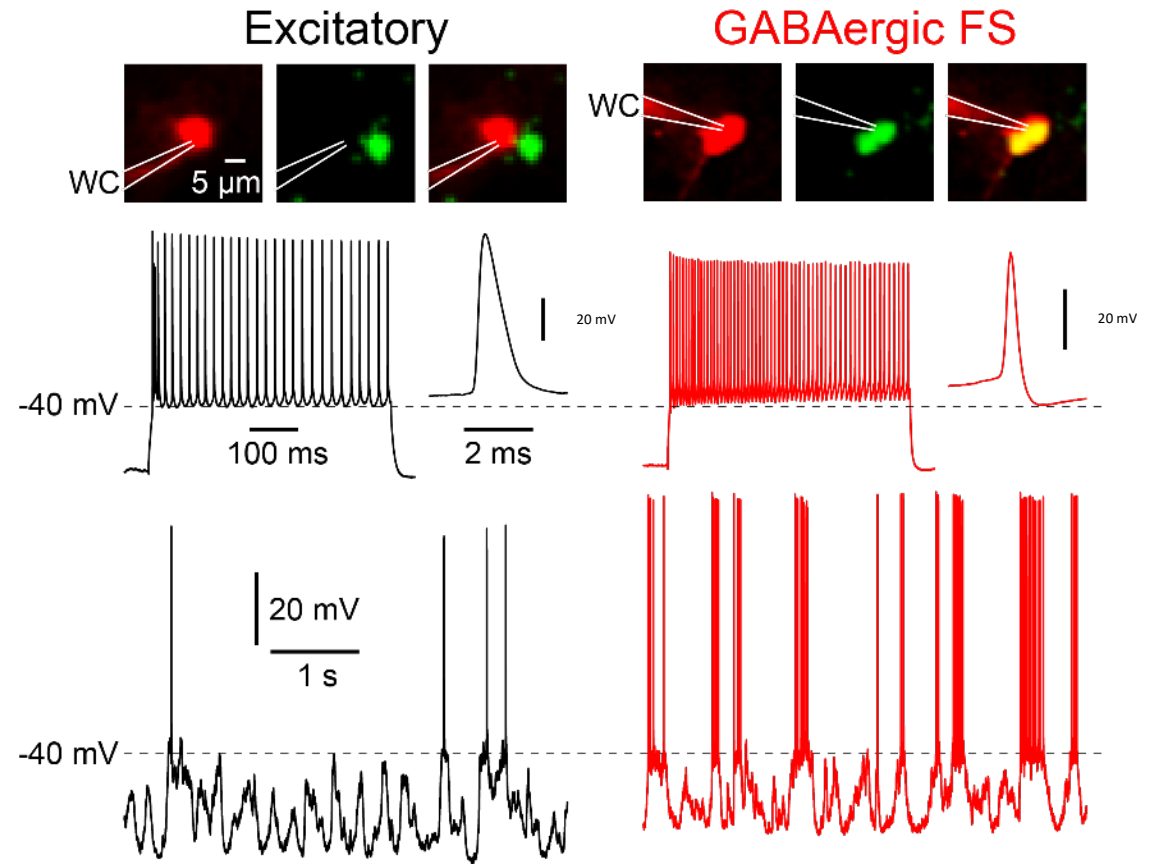
Cell-type specific changes with cortical states

Targeted Whole-Cell Recordings in the Mammalian Brain In Vivo



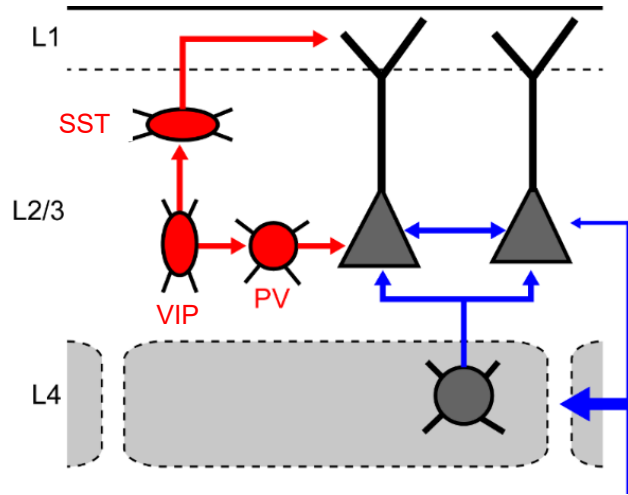
(Margrie et al., 2003; Komai et al., 2006)

Targeted Patch-Clamp recording in GAD67-GFP mouse



(Gentet et al., Neuron 2010 ; Gentet et al., Nat. Neurosci. 2012)

Cell-type specific changes with cortical states

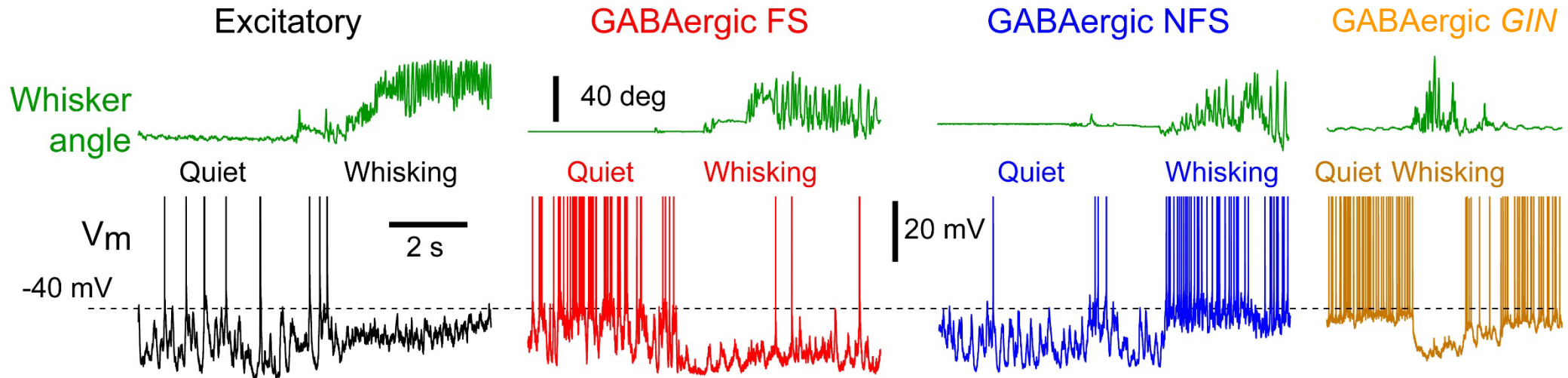


Excitatory Pyramidal

Fast Spiking (FS) ~ GABAergic PV

Non Fast-Spiking (NFS) ~ GABAergic VIP

GIN ~ GABAergic SST

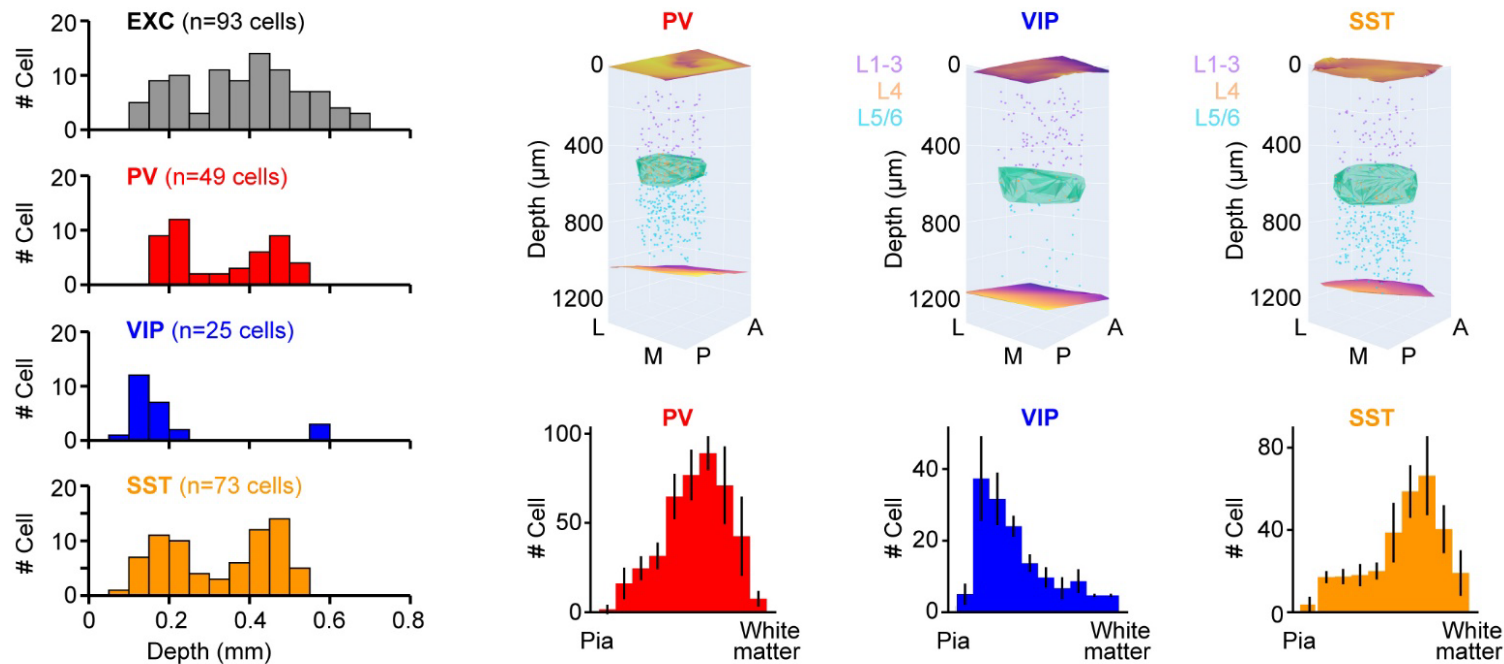


=> But cell-type identification approximative and limitation to superficial L2/3

Cell-type specific changes with cortical states

New study – Kiritani et al., PLOS One 2024

- Better cell-type identification => PV-Cre, VIP-Cre and SST-Cre mouse lines
- Recordings at lower cortical depth => improved 2-photon targeting

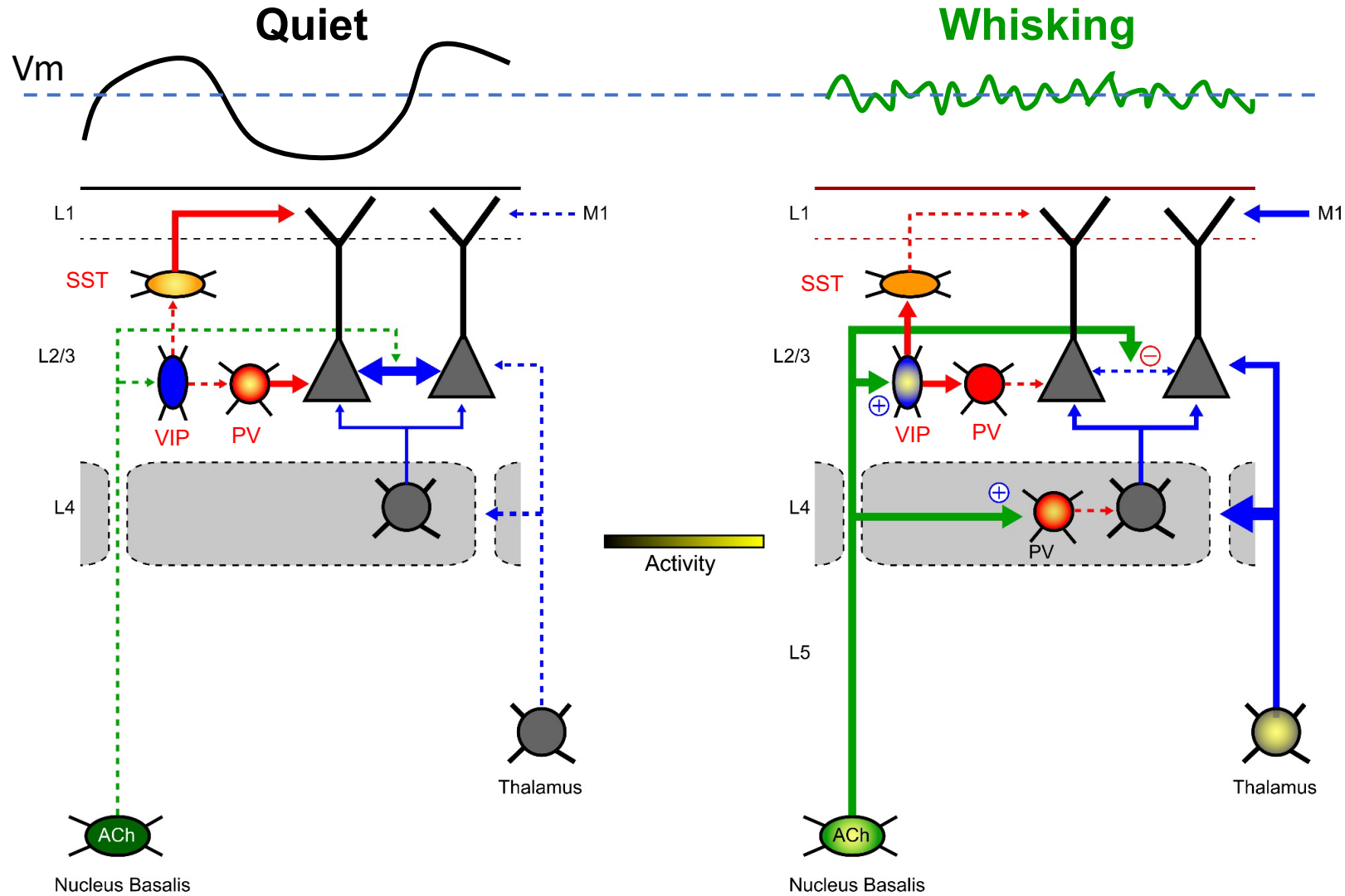


(Kiritani et al., PLOS One 2024)

=> BIO482 Miniproject:

Learn more about these different cell-types by yourself

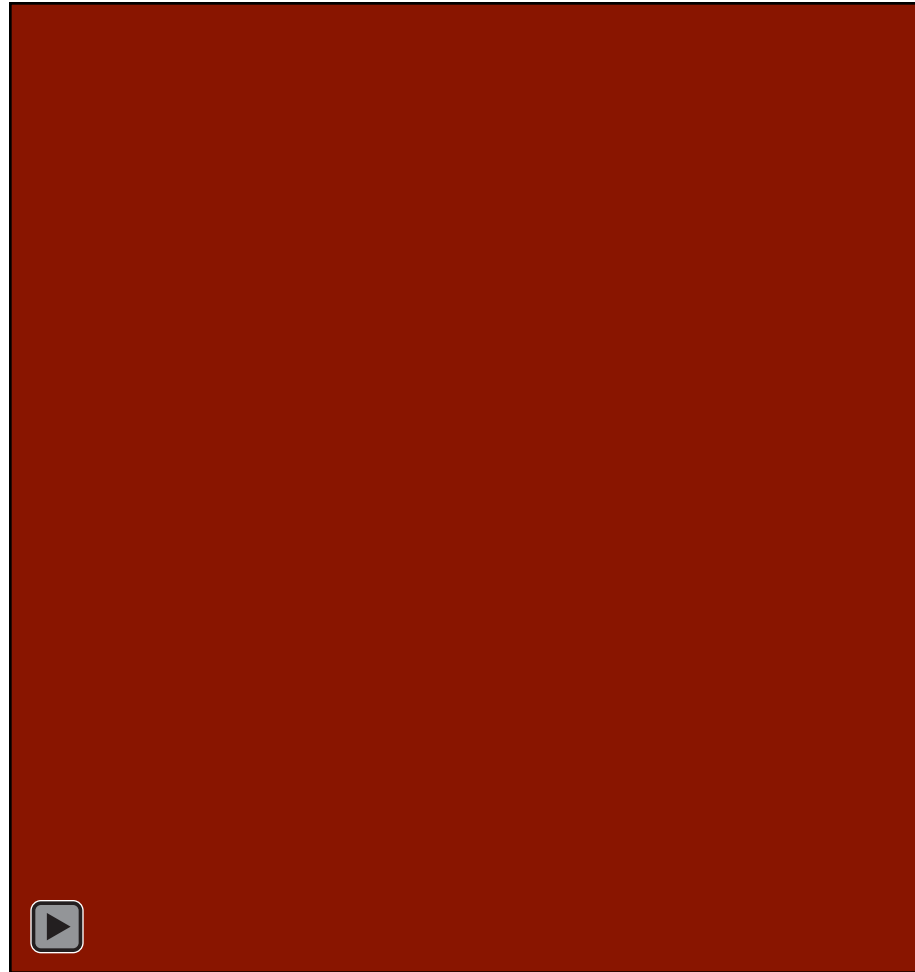
Cellular mechanisms of the state change in wS1



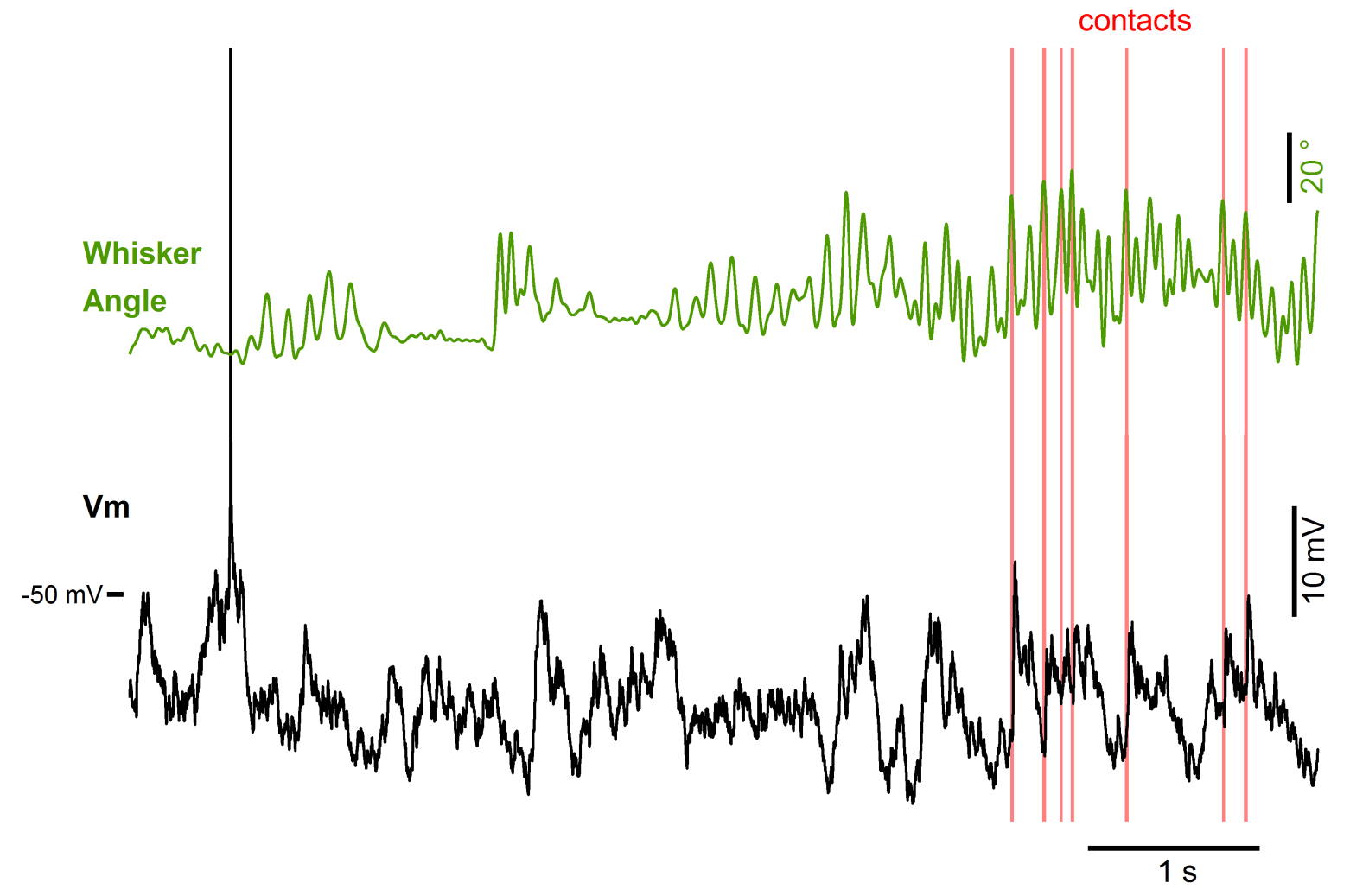
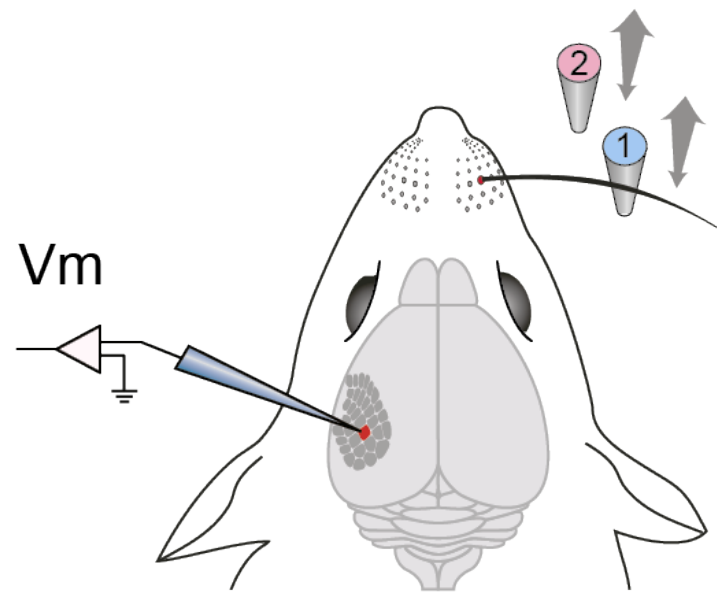
See <https://www.frontiersin.org/journals/systems-neuroscience/articles/10.3389/fnsys.2018.00064/full>

2. Sparse representation of sensory inputs in layer 2/3

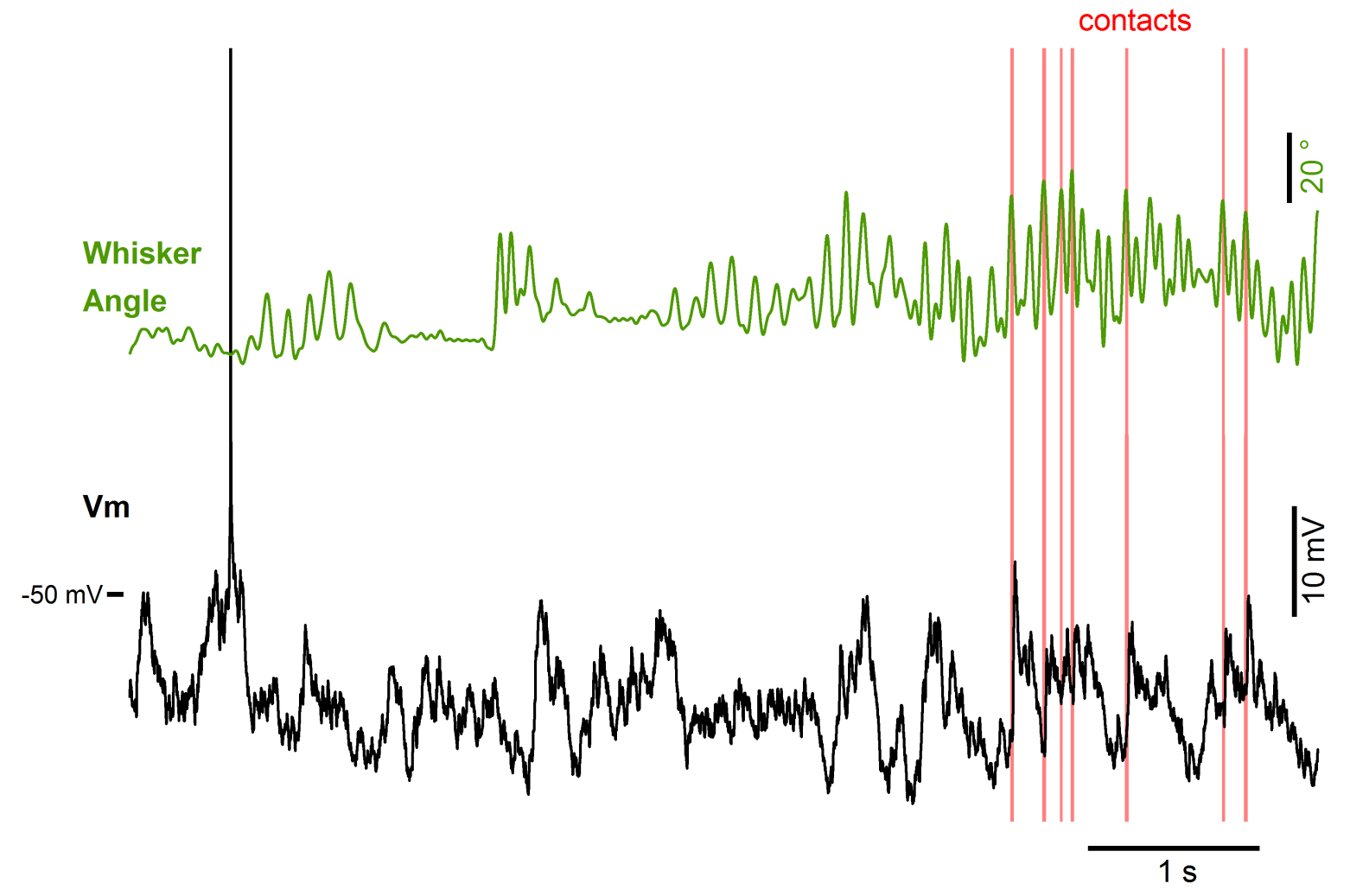
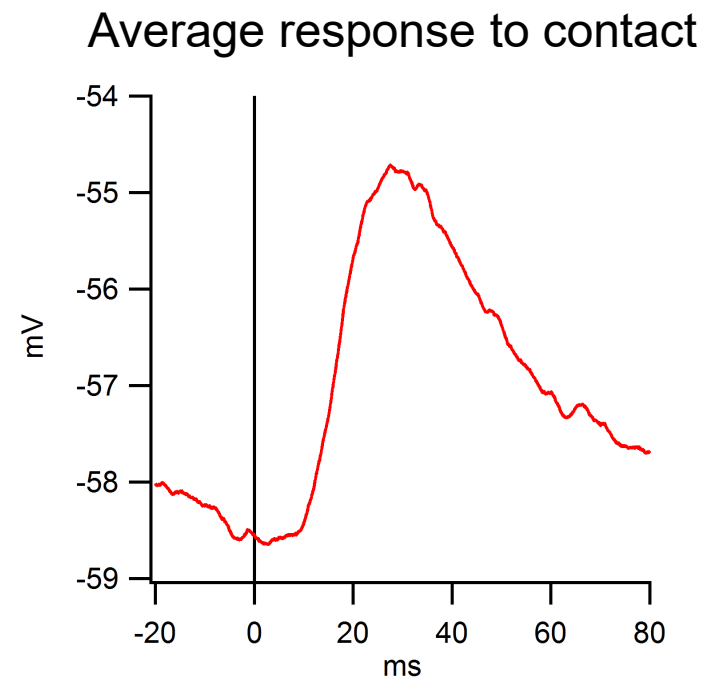
■ Membrane potential recording during active touch



Membrane potential recording during active touch



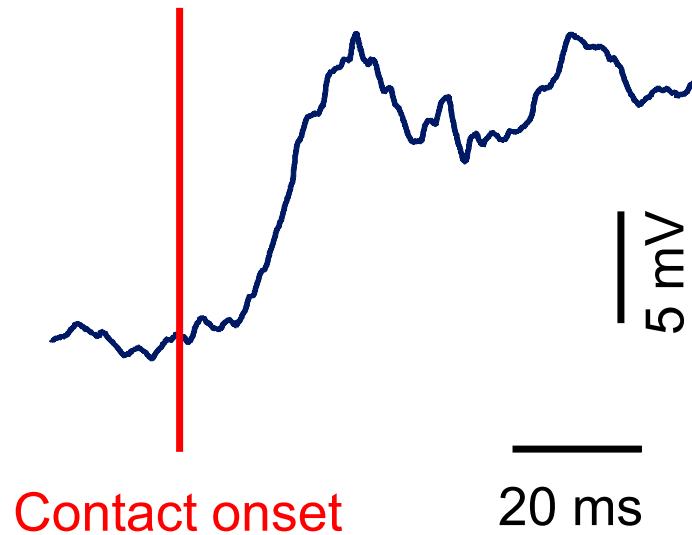
Membrane potential recording during active touch



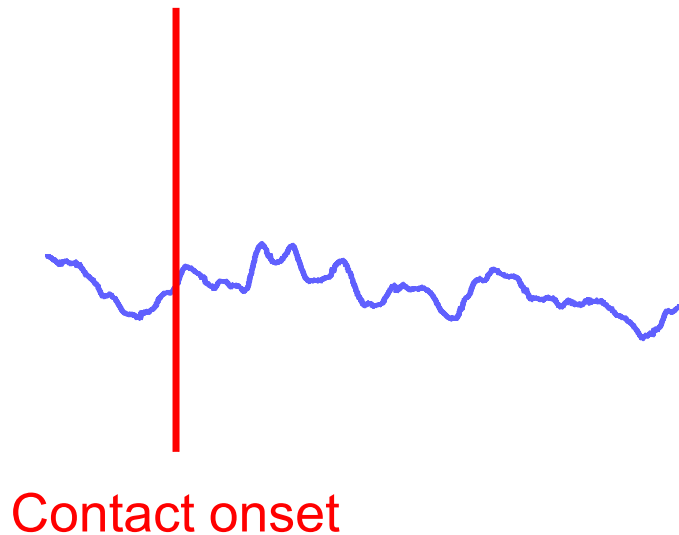
Variable evoked responses to active touch

Example of individual postsynaptic potentials (PSPs) evoked by contact

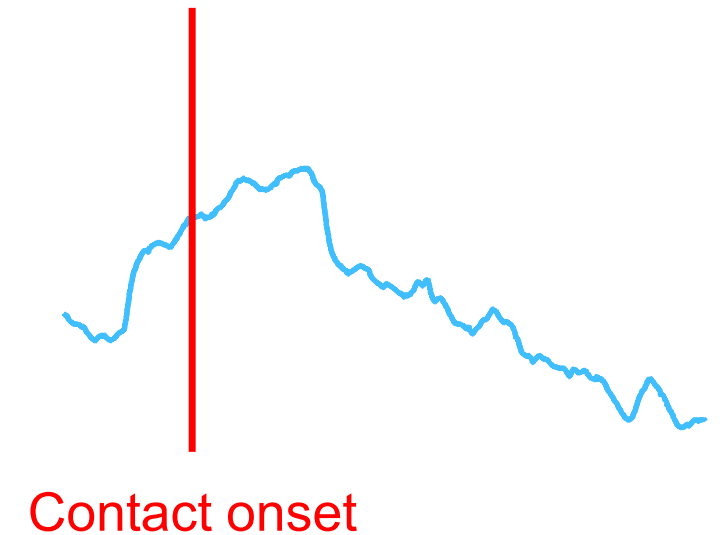
depolarizing response



small response

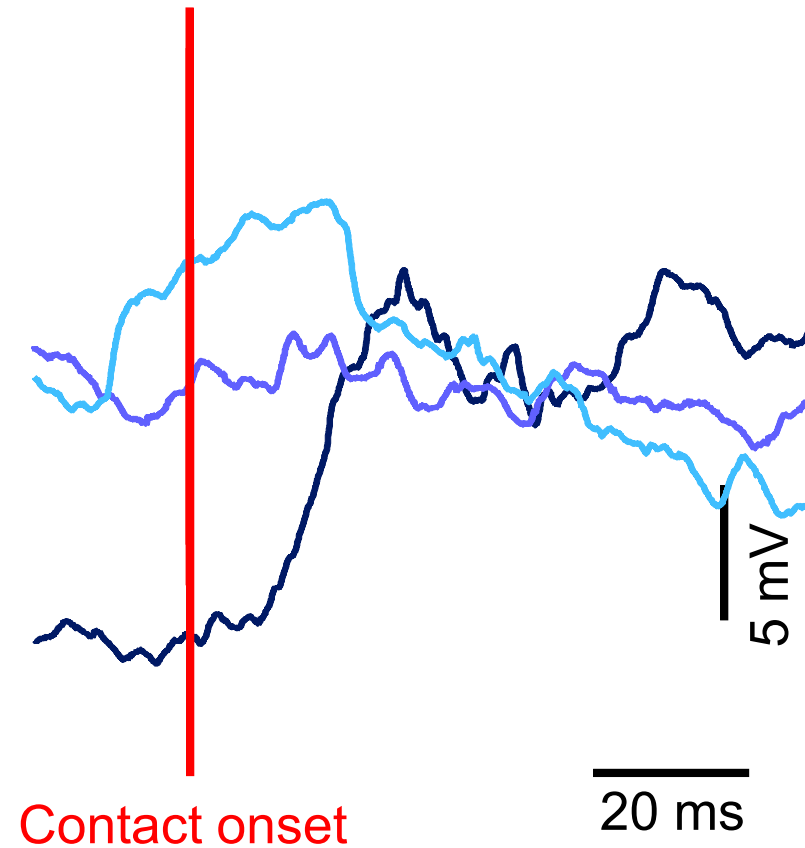


hyperpolarizing response



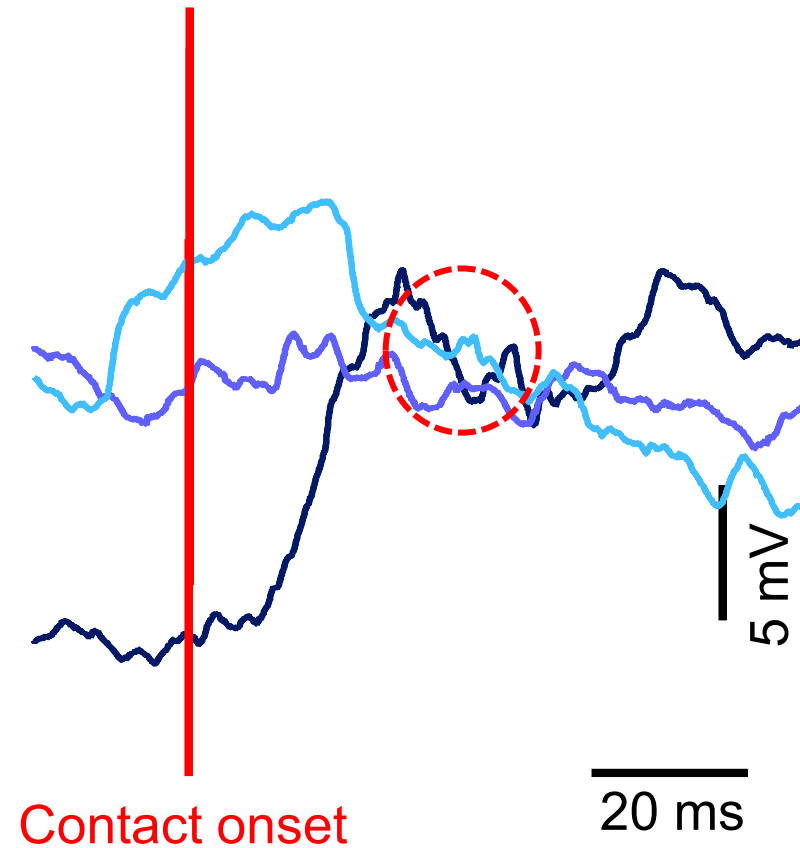
Variable evoked responses to active touch

Example of individual postsynaptic potentials (PSPs) evoked by contact



■ Variable evoked responses to active touch

Example of individual postsynaptic potentials (PSPs) evoked by contact

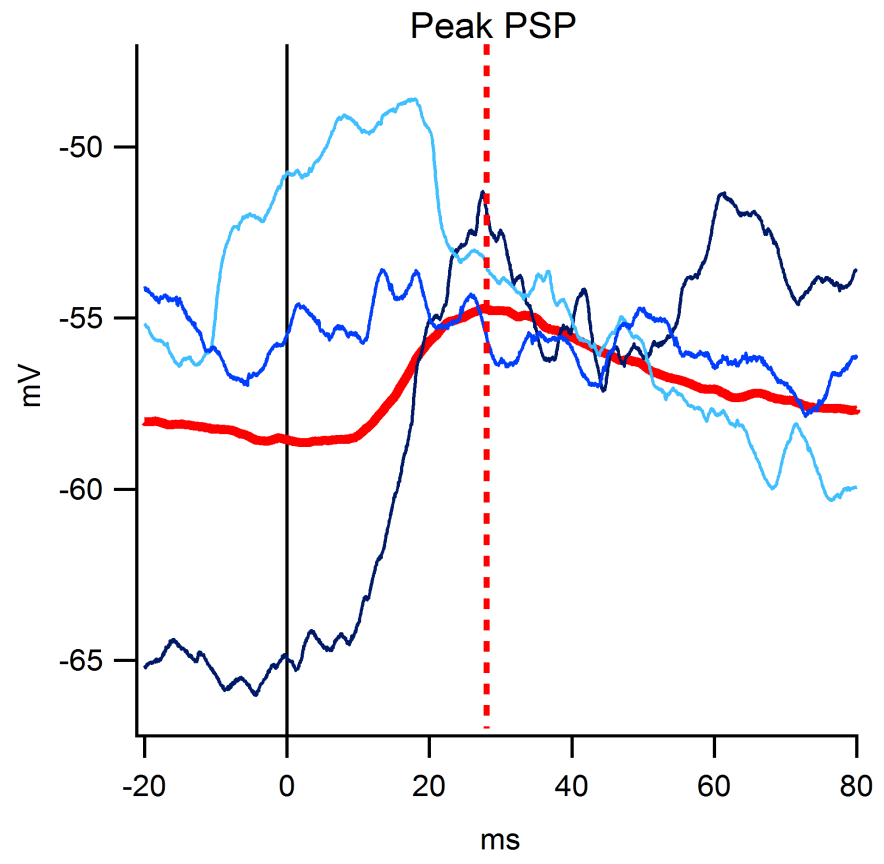


=> Individual PSPs converge to a fixed-point of the V_m

Variable evoked responses to active touch

Individual PSPs

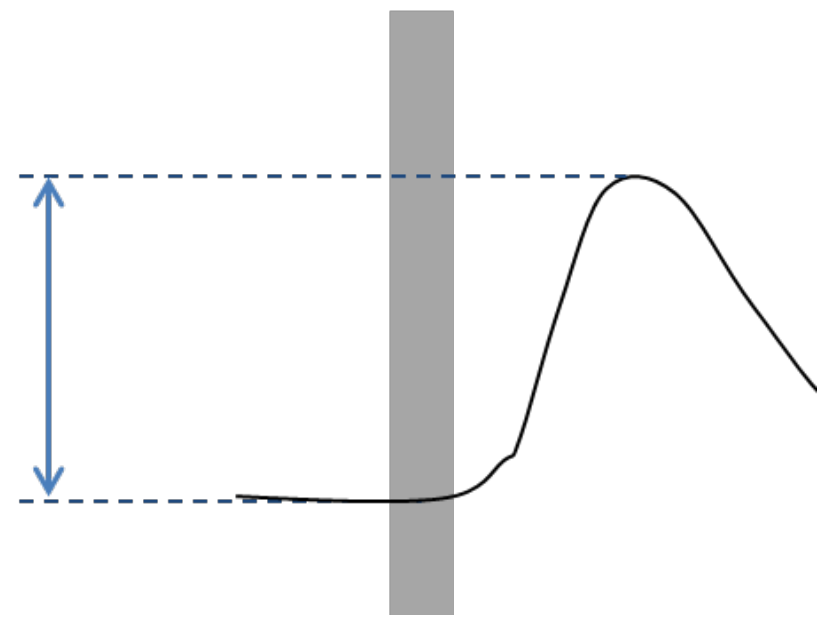
Average PSP



Contact

PSP amplitude

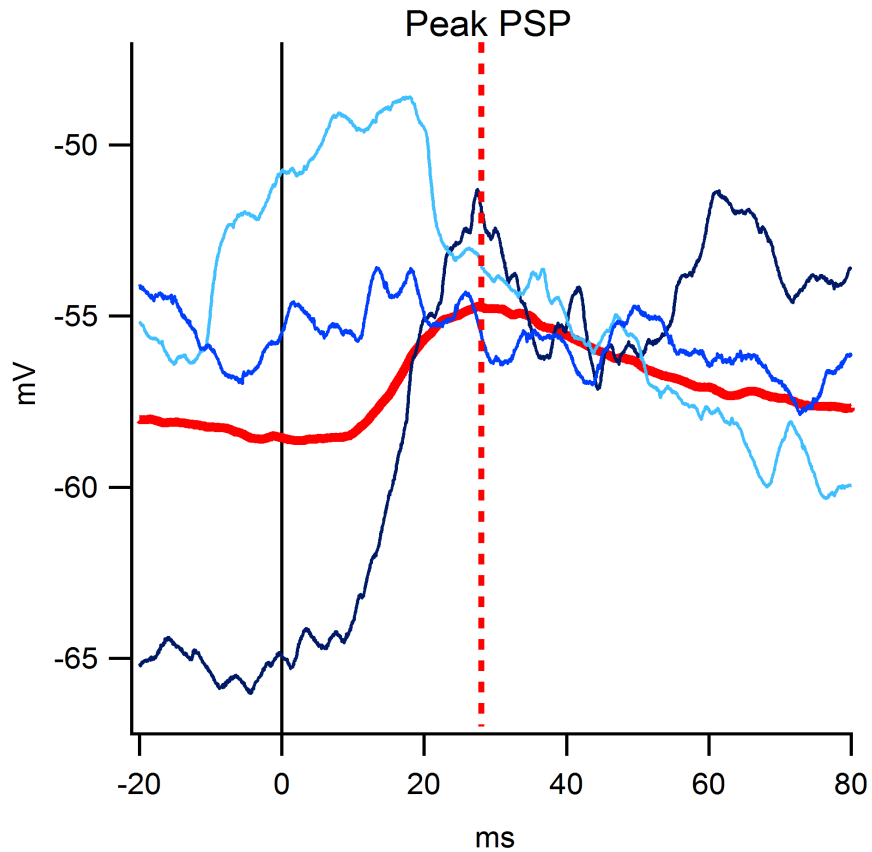
Baseline Vm



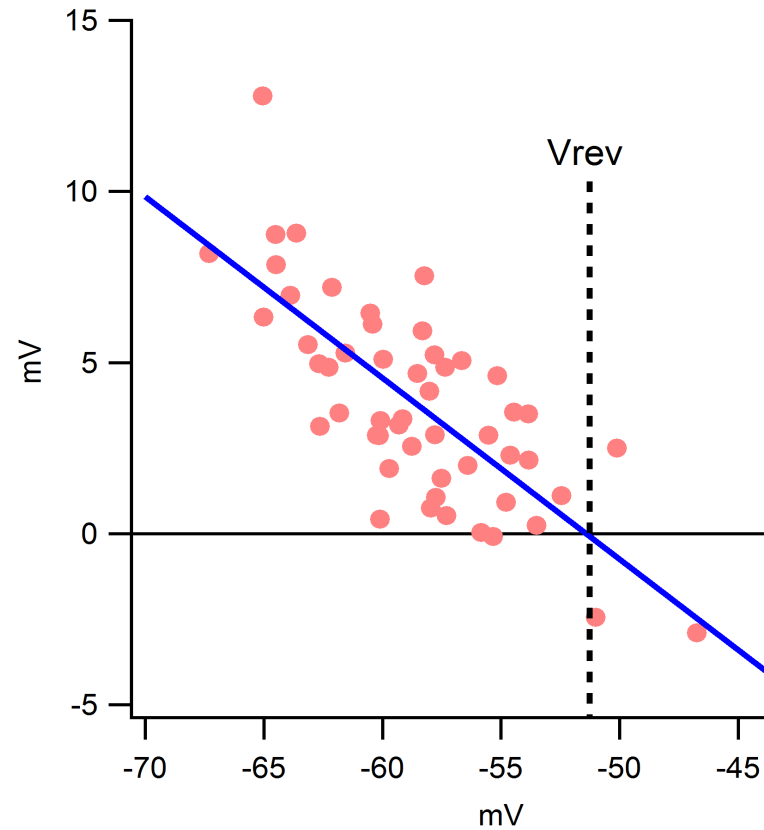
■ Reversal potential (V_{rev}) of the sensory evoked response

Individual PSPs

Average PSP



PSP amplitude vs Baseline V_m

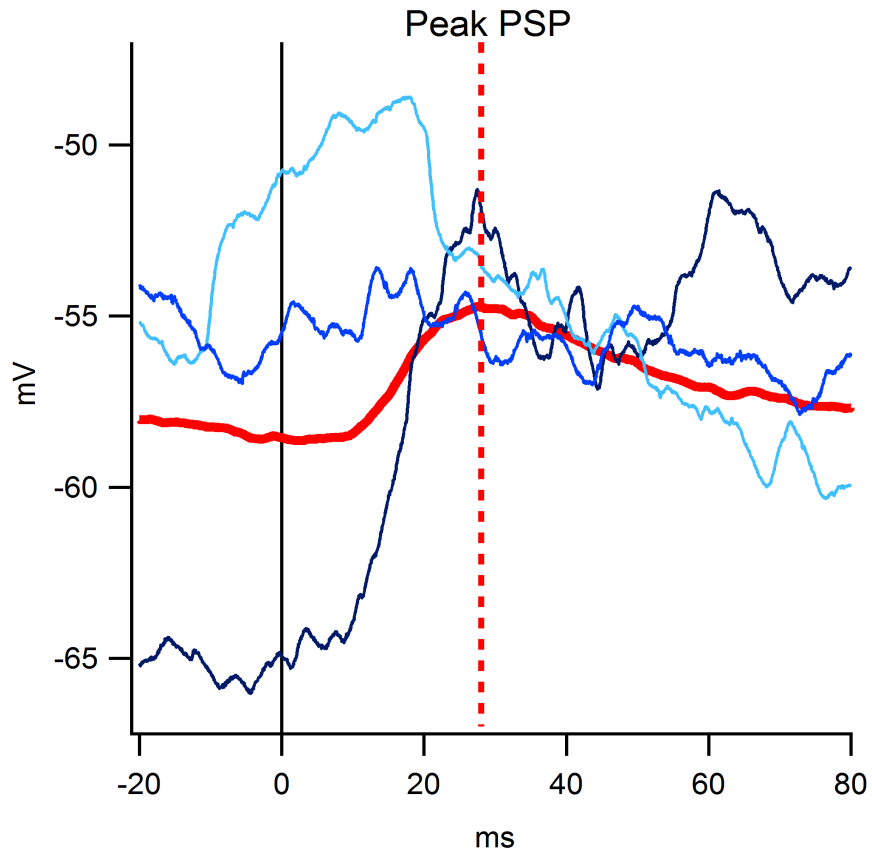


=> Negative V_{rev} of the PSP

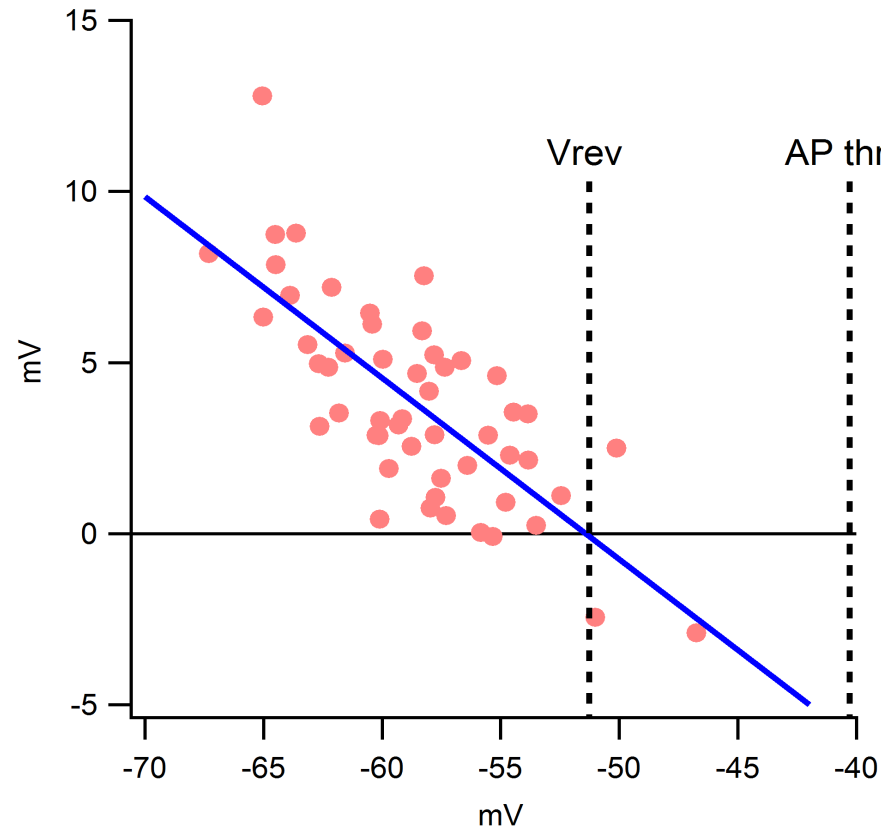
Reversal potential (V_{rev}) of the sensory evoked response

Individual PSPs

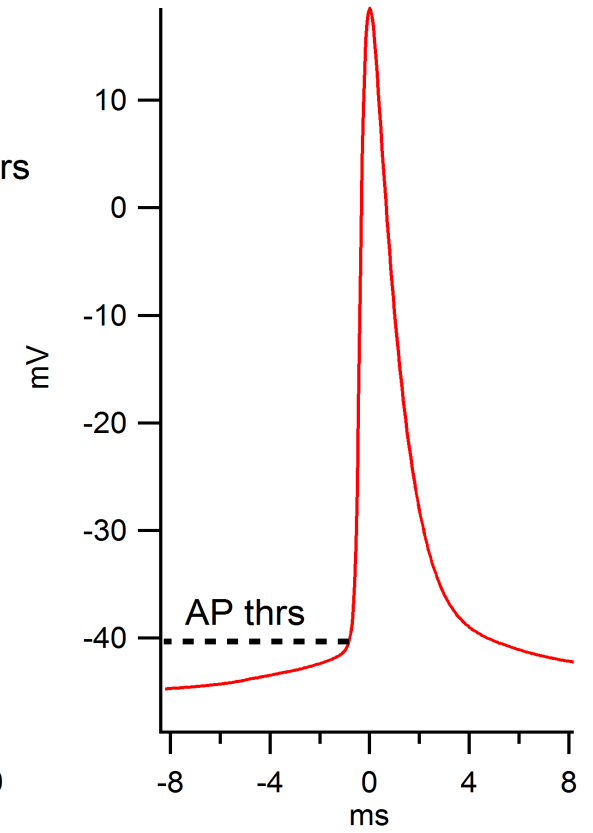
Average PSP



PSP amplitude vs Baseline V_m

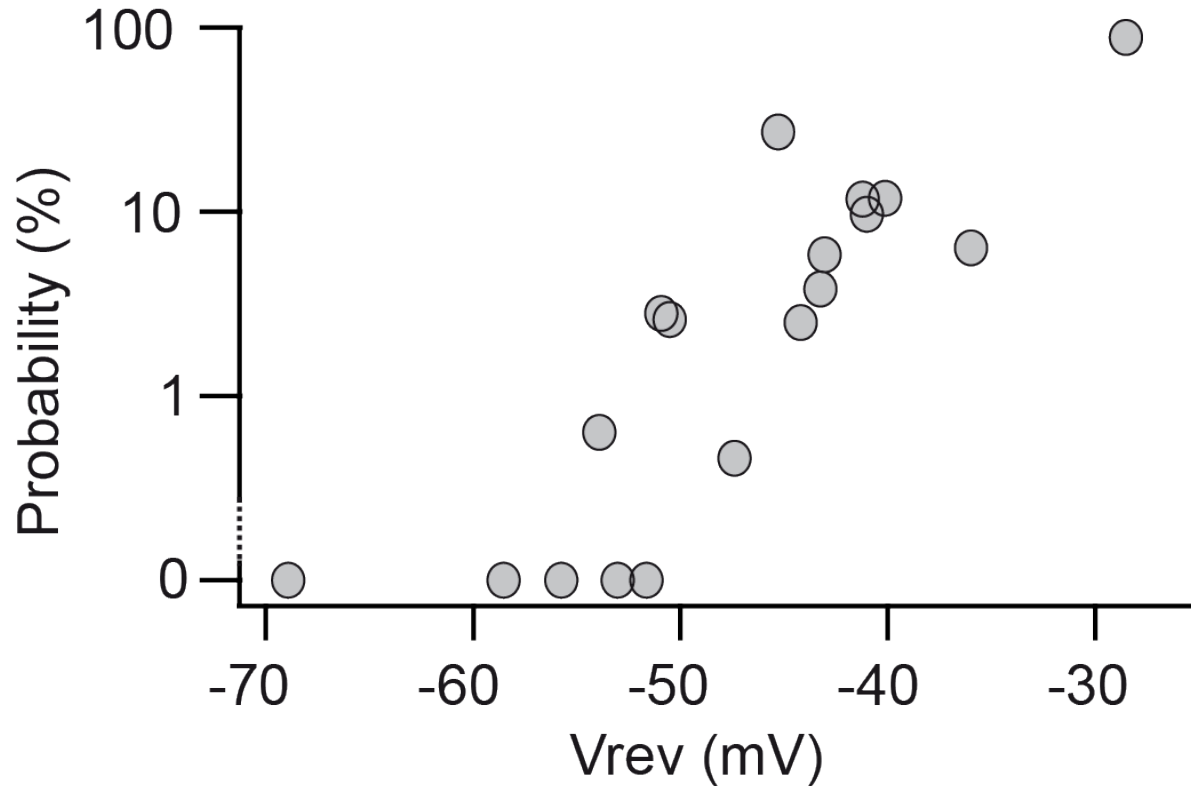
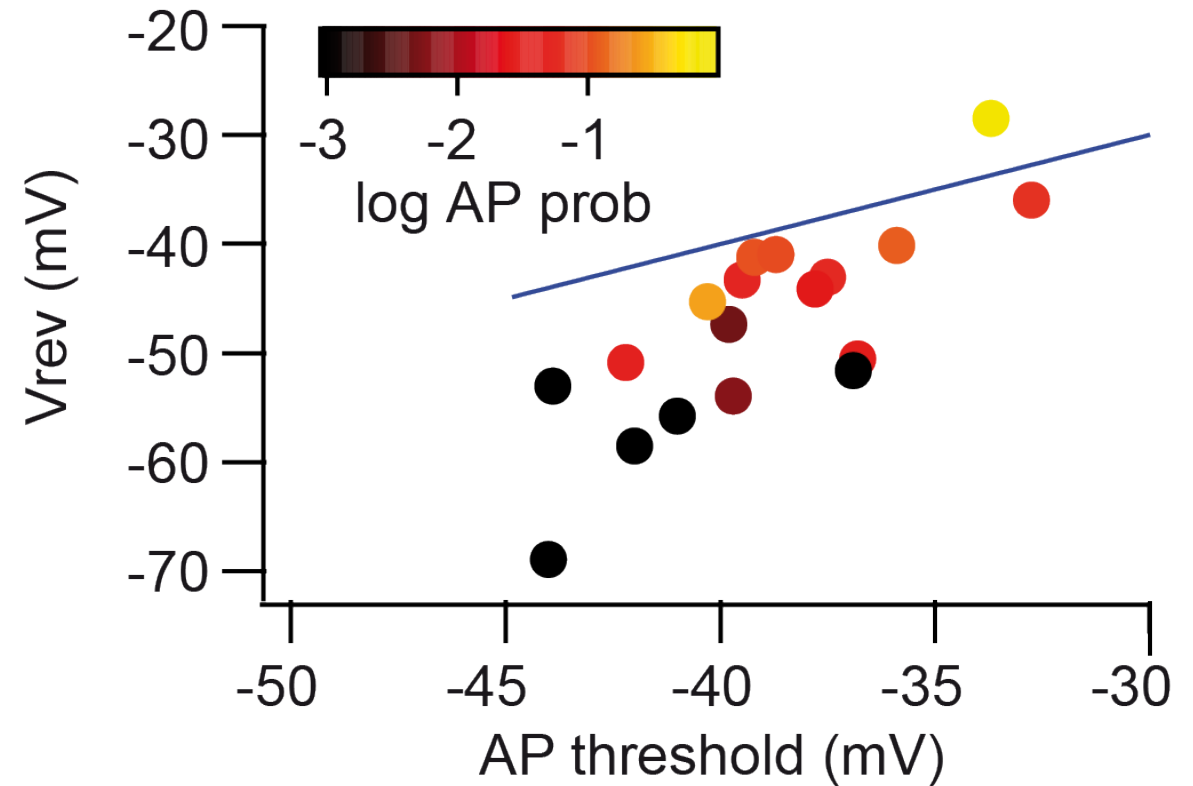


AP Threshold



=> Negative V_{rev} of the PSP

- V_{rev} of the evoked response accounts for AP probability

AP probability vs V_{rev}  V_{rev} vs AP threshold

=> V_{rev} of individual cell correlates with AP firing probability

■ Cortical inhibition mediates sparse response

Synaptic Inputs =>

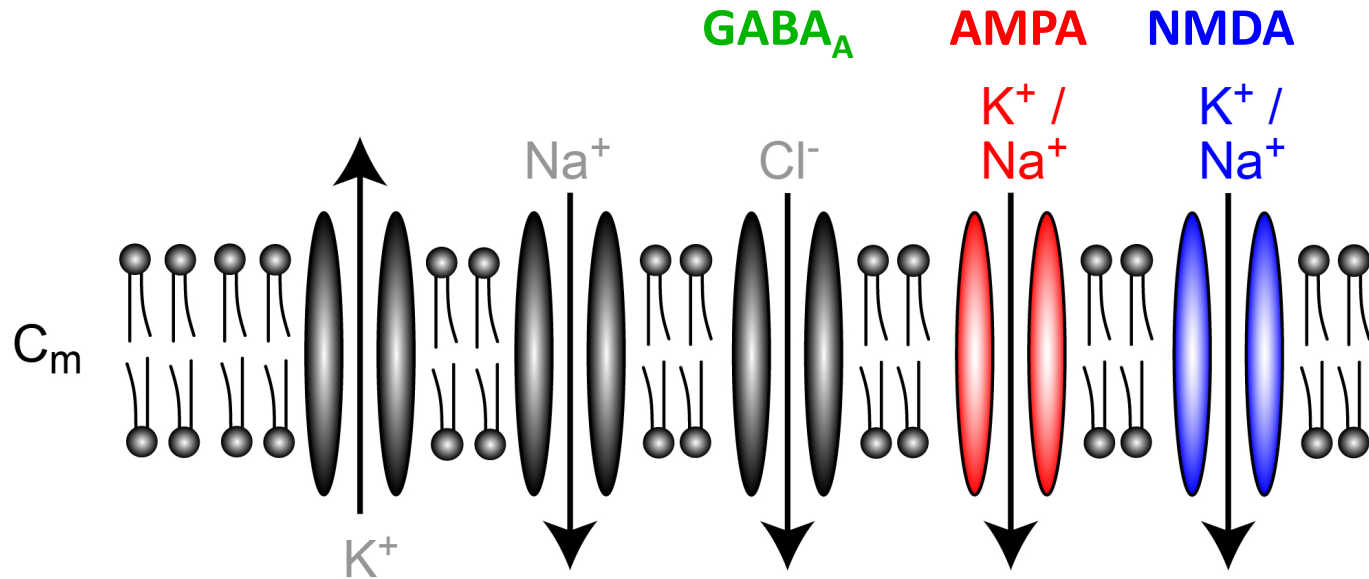
Inhibition (GABA)

Excitation (Glutamate)

$E_{rev} \text{ GABA}_A \sim -80 \text{ mV}$

$E_{rev} \text{ AMPA} \sim 0 \text{ mV}$

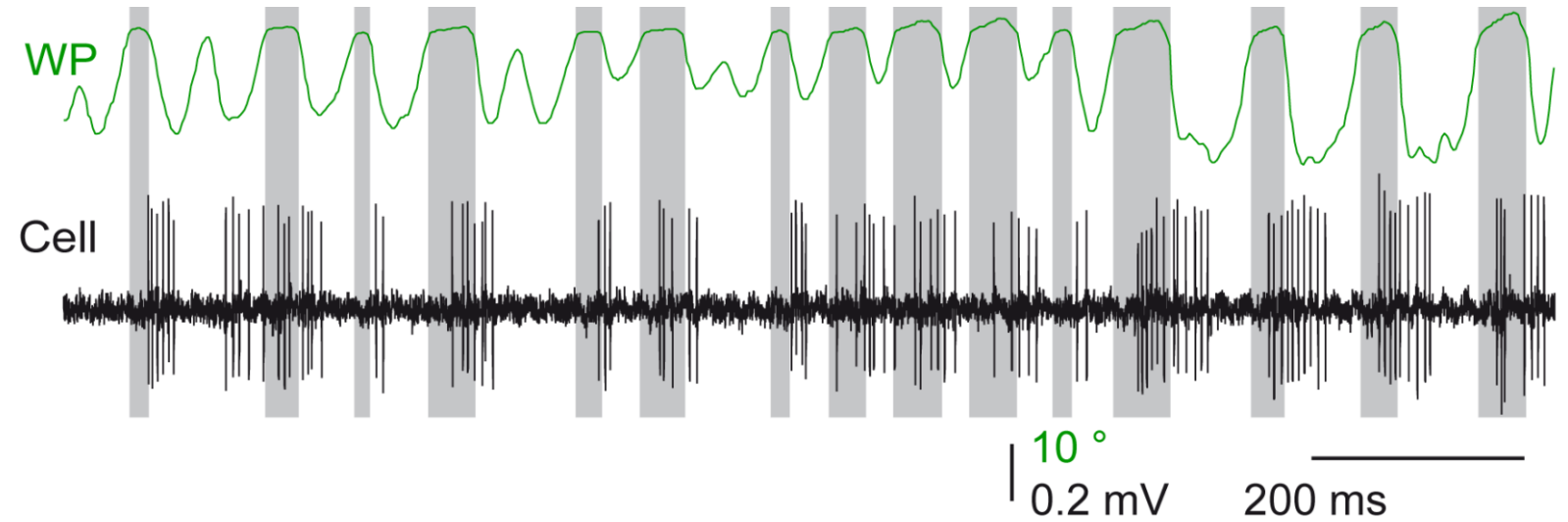
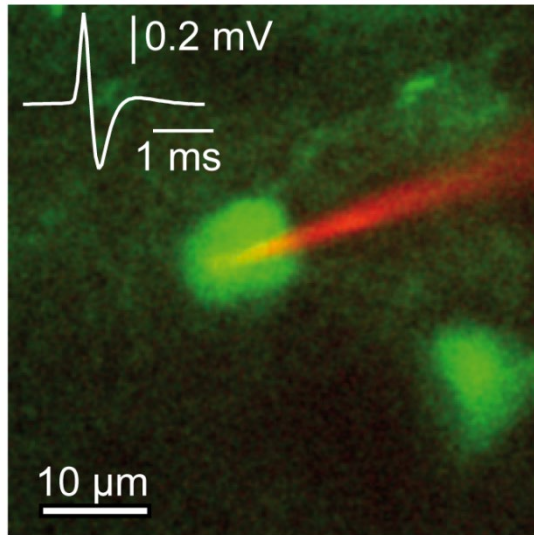
$E_{rev} \text{ NMDA} \sim 0 \text{ mV}$



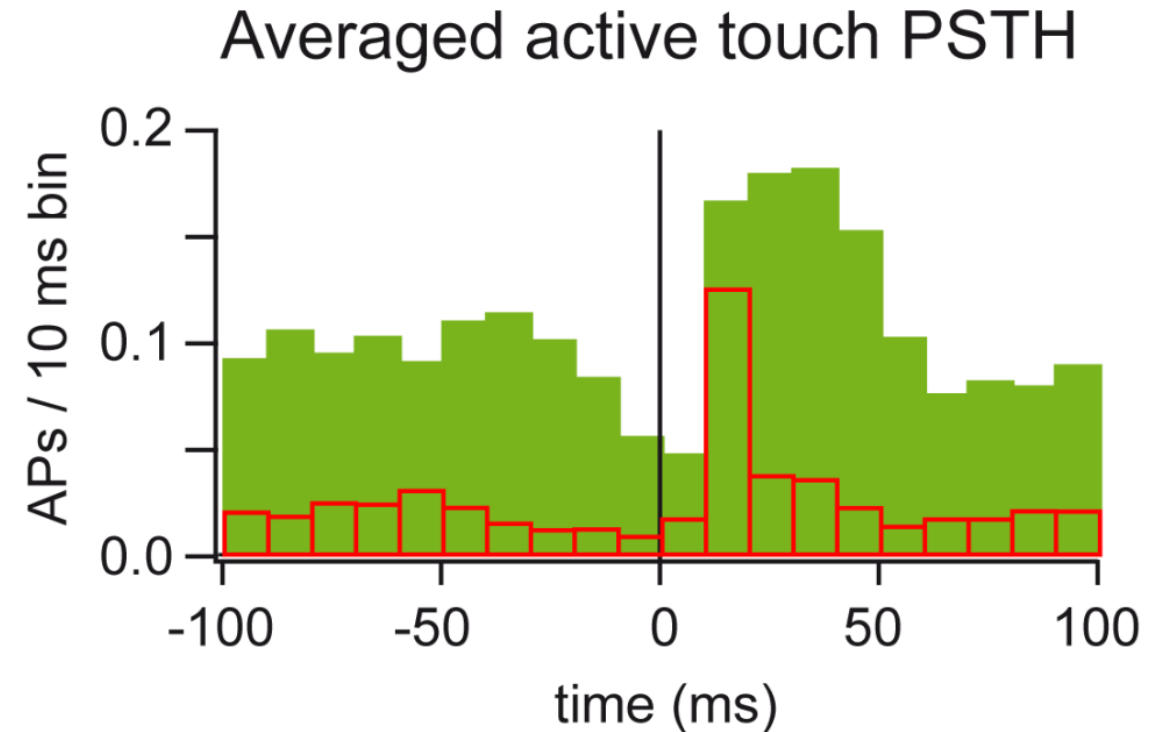
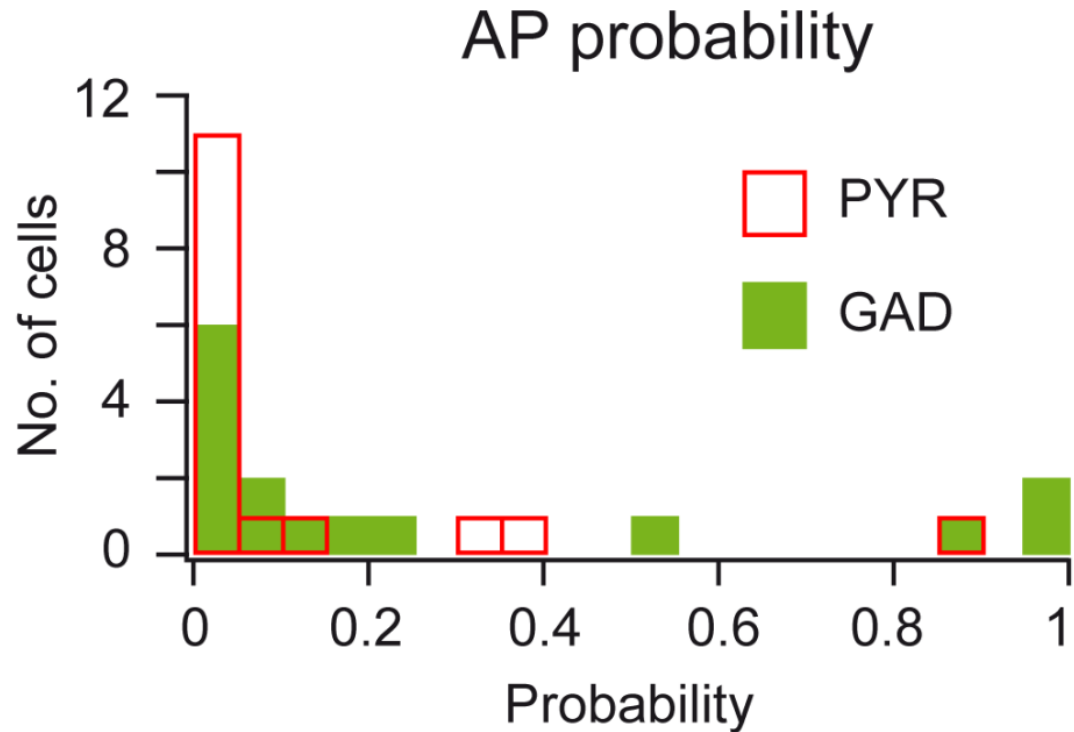
=> Negative V_{rev} of the PSP implies contribution of inhibition

■ Cortical inhibition mediates sparse response

Targeted juxtacellular recording in
GAD67-GFP knockin mouse



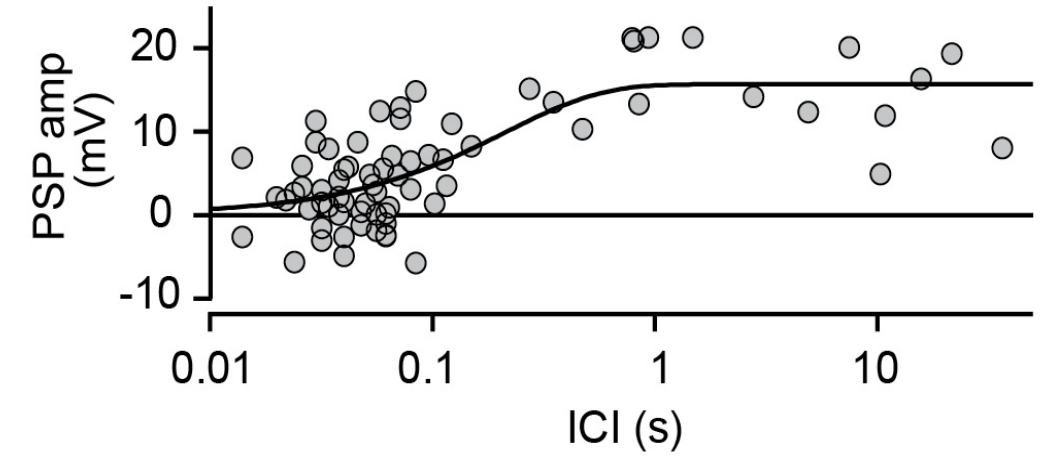
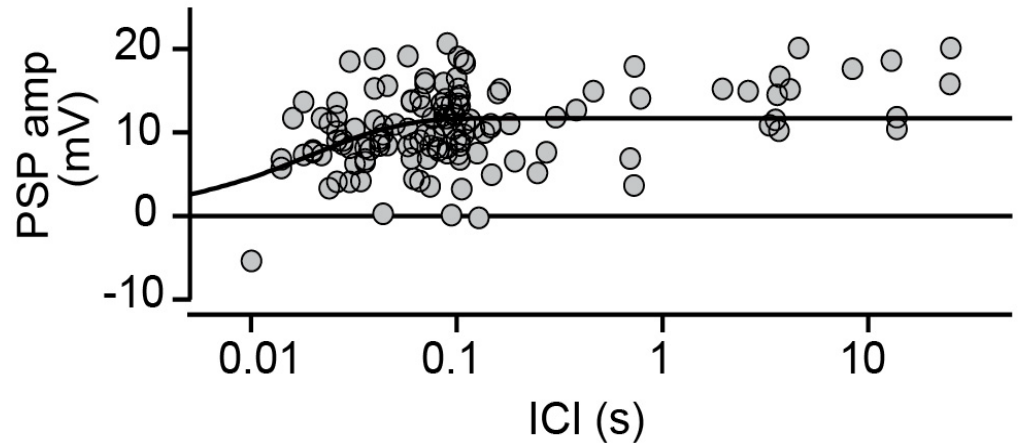
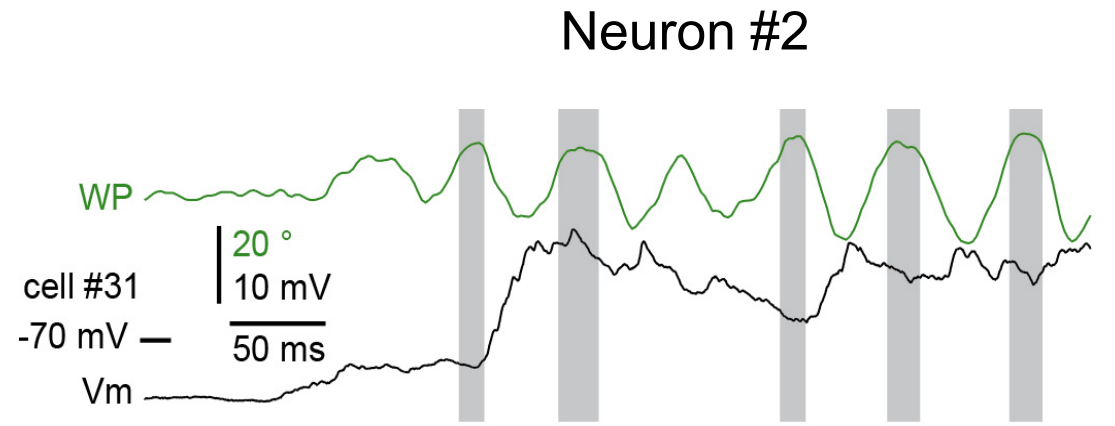
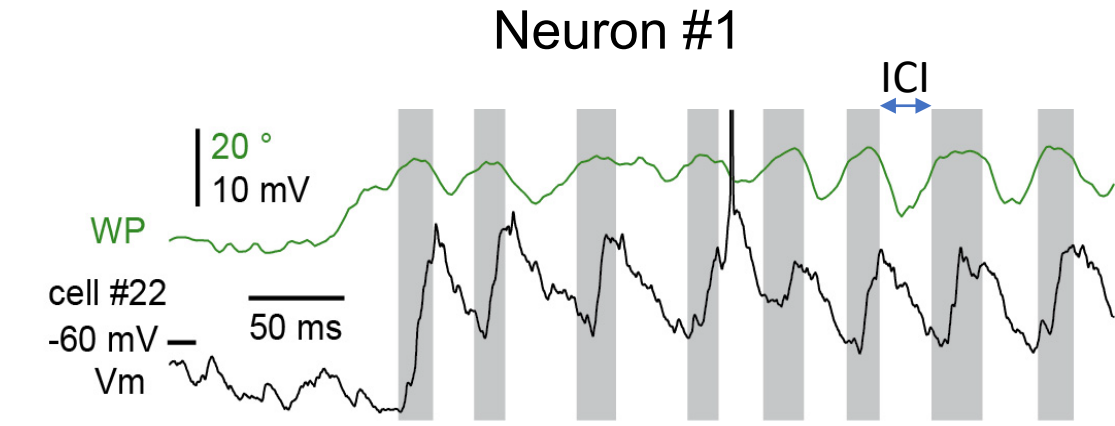
■ Cortical inhibition mediates sparse response



(Crochet et al., Neuron 2011)

=> GABAergic neurons fire more reliably than EXC neurons in response active touch

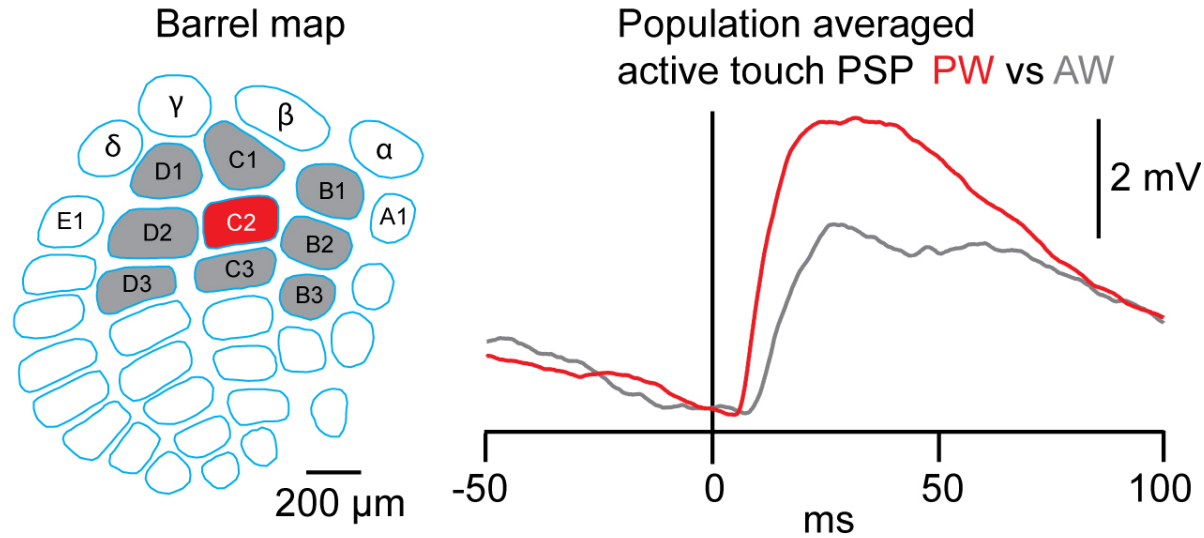
■ Short-term dynamics



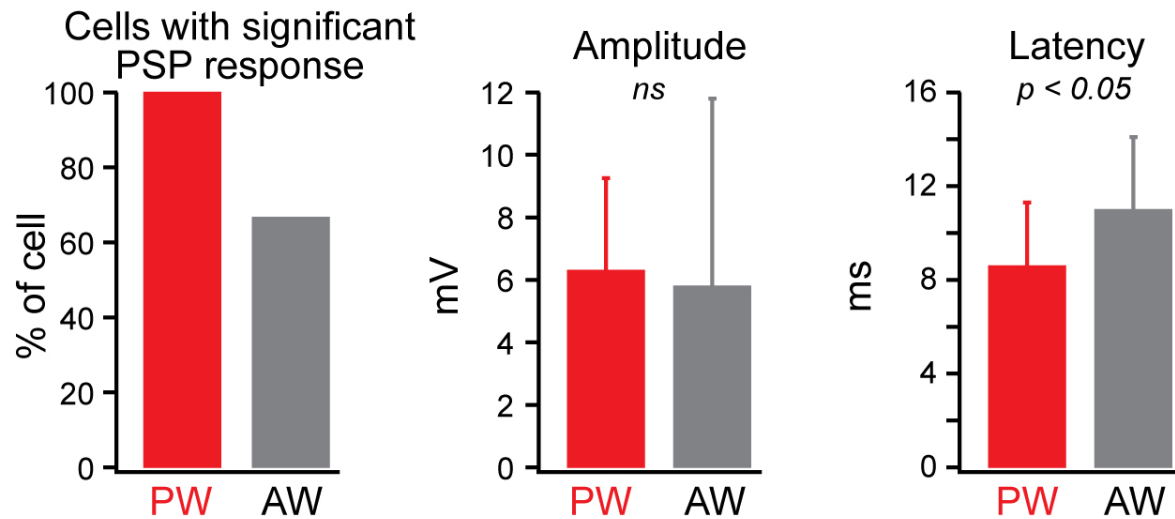
(Crochet et al., Neuron 2011)

=> EXC neurons have different short-term dynamics in response to successive active contacts

Principal vs surrounding cortical columns



=> Decreased response probability and longer latencies in surrounding vs principal cortical columns

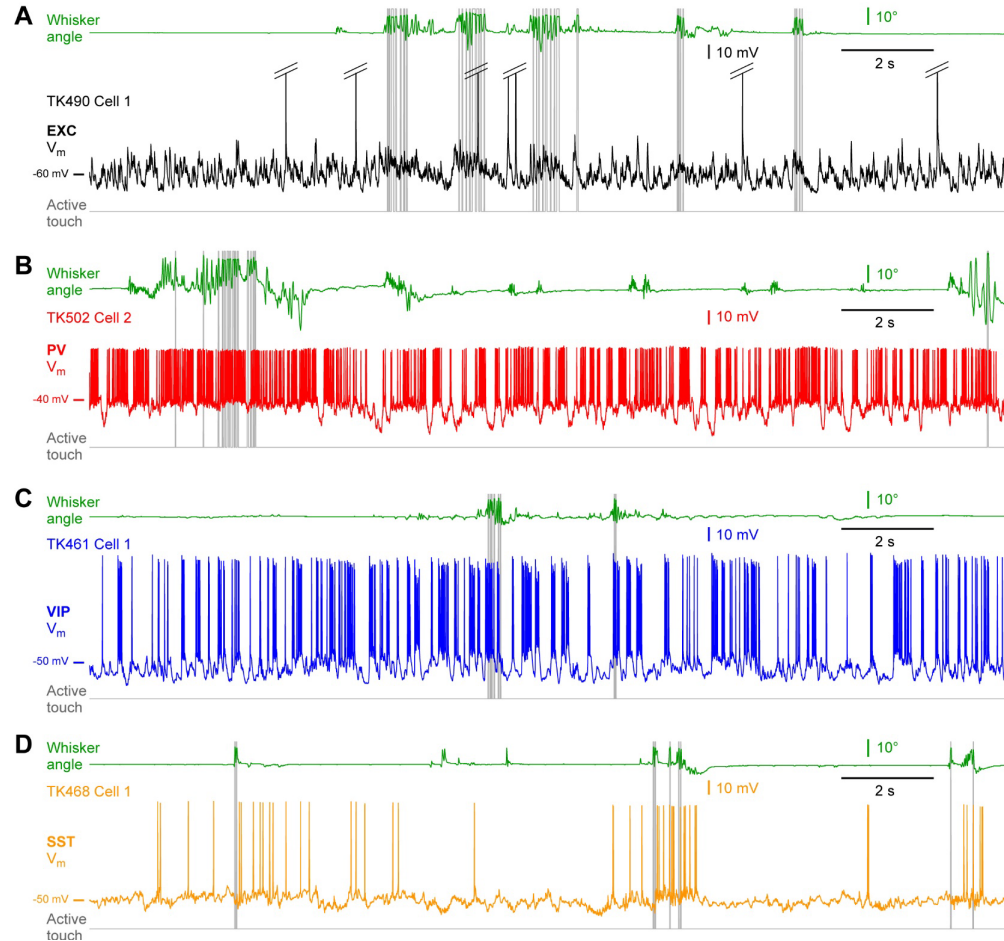


(Crochet et al., Neuron 2011)

Active touch responses in different cell-types

New study – Kiritani et al., PLOS One 2024

- Also active touch



=> BIO482 Miniproject:

Learn more about active touch responses in different cell-types by yourself

(Kiritani et al., PLOS one 2024)

QUESTIONS ?