

Inference of Global States Stability in Cortical Networks

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The Default Global State of the Cerebral Cortex [1]

Phenomenology

- ▶ Slow Oscillations ($\leq 1\text{Hz}$)
 - ▷ at the neuron level (membrane potential and firing rate)
 - ▷ at the **network level** (extracellular electrical activity: LFP and MUA)
- ▶ Observed in:
 - ▷ slow wave sleep
 - ▷ deep anaesthesia
 - ▷ deafferentation
 - ▷ **cortical slices**

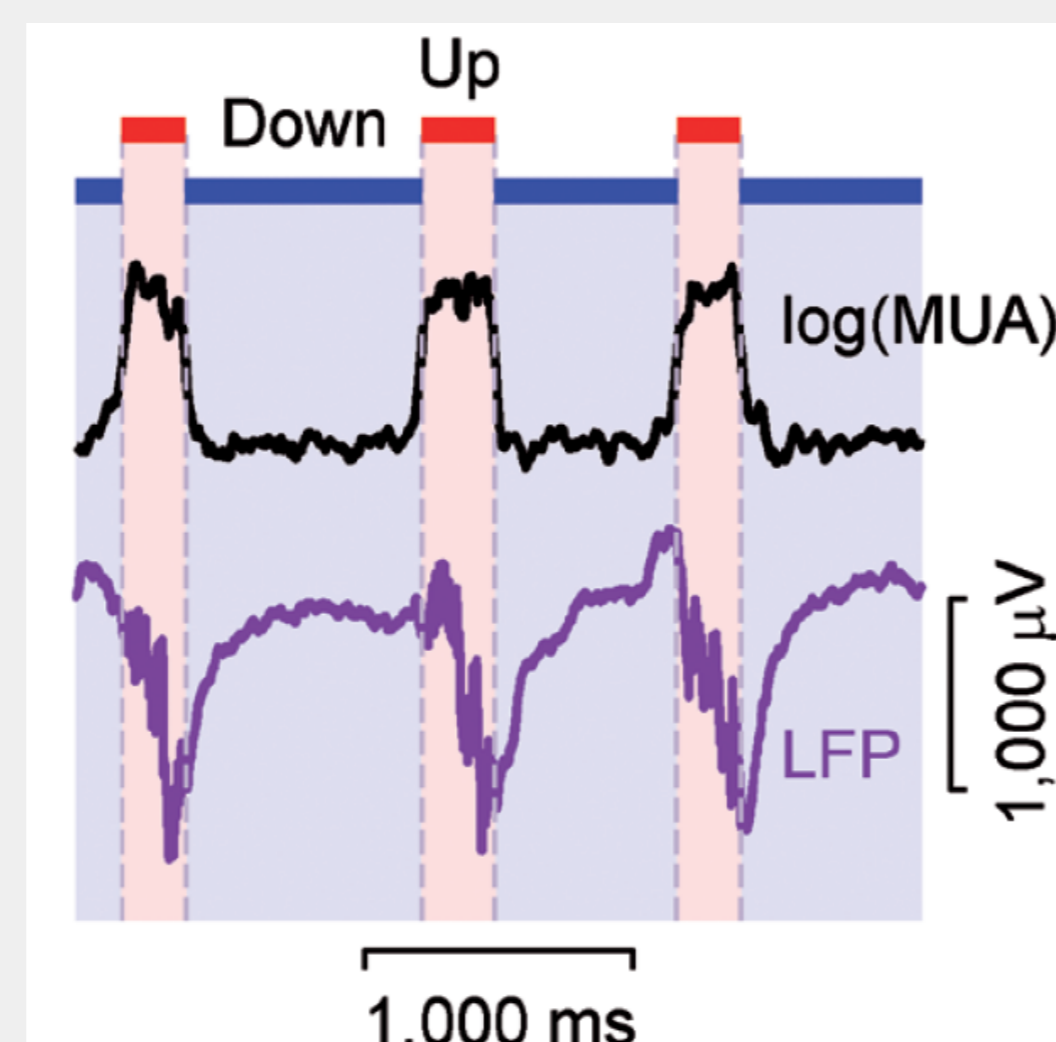


Figure 1: Slow oscillations recorded from the frontal cortex of an anaesthetized mouse [1]

Key Features

- ▶ Bistability
 - ▷ existence of two attractors: UP state and DOWN state
- ▶ Intrinsic fluctuations between these attractors
- ▶ Regularity of the UP/DOWN states alternance

→ behaves as a relaxation oscillator

Advantages

- ▶ Resilience to perturbances:
 - ▷ the relaxation-oscillator regime acts as an equilibrium of the network.
- ▶ Facilitation of the transition towards more connected, awake-like states.

Motivation

- ▶ Can we detect and characterize other global network states apart from the SO regime?

Cortical Networks Parameters: Literature Review

In fact, the dynamics of the network states are not fully understood [3]. The stability of such states seems to be strongly influenced by:

- ▶ the input stimulus [4]
- ▶ the connectivity properties of the network
 - ▷ either unshaped or structured in clusters [5]
- ▶ the excitatory-inhibitory balance [6]
- ▶ the network architecture
 - ▷ either predominantly feedforward or recurrent [7]
- ▶ the kind of noise
 - ▷ intrinsic or extrinsic [8]

Problem: Although multiple mechanistic hypotheses have been proposed in models, the current analysis tools do not enable us to discern empirically the dynamics of the network states.

References

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Cortical Slices Recordings

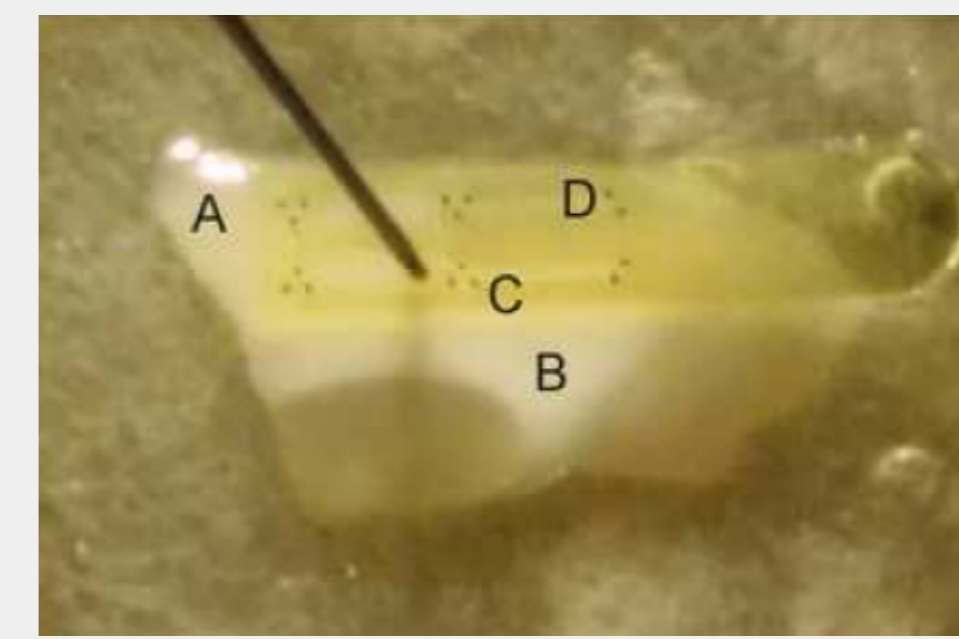


Figure 2: Cortical slice recording setup: (A) cortex, (B) white matter, (C) infragranular layers' electrodes and (D) supragranular layers' electrodes

Experimental Conditions [2]

- ▶ Pharmacological Modulations
 - ▷ adding Carbachol ($0.5 \mu\text{M}$) + Norepinephrine ($50 \mu\text{M}$)
 - ▷ reducing extracellular Calcium (to $0.8\text{-}0.9 \text{mM}$)
 - ▷ Reducing temperature (to $31\text{-}32 \text{ }^\circ\text{C}$)
- ▶ Electrical Stimuli
 - ▷ $150 \mu\text{A}$ pulses every 10 s at layer 5.

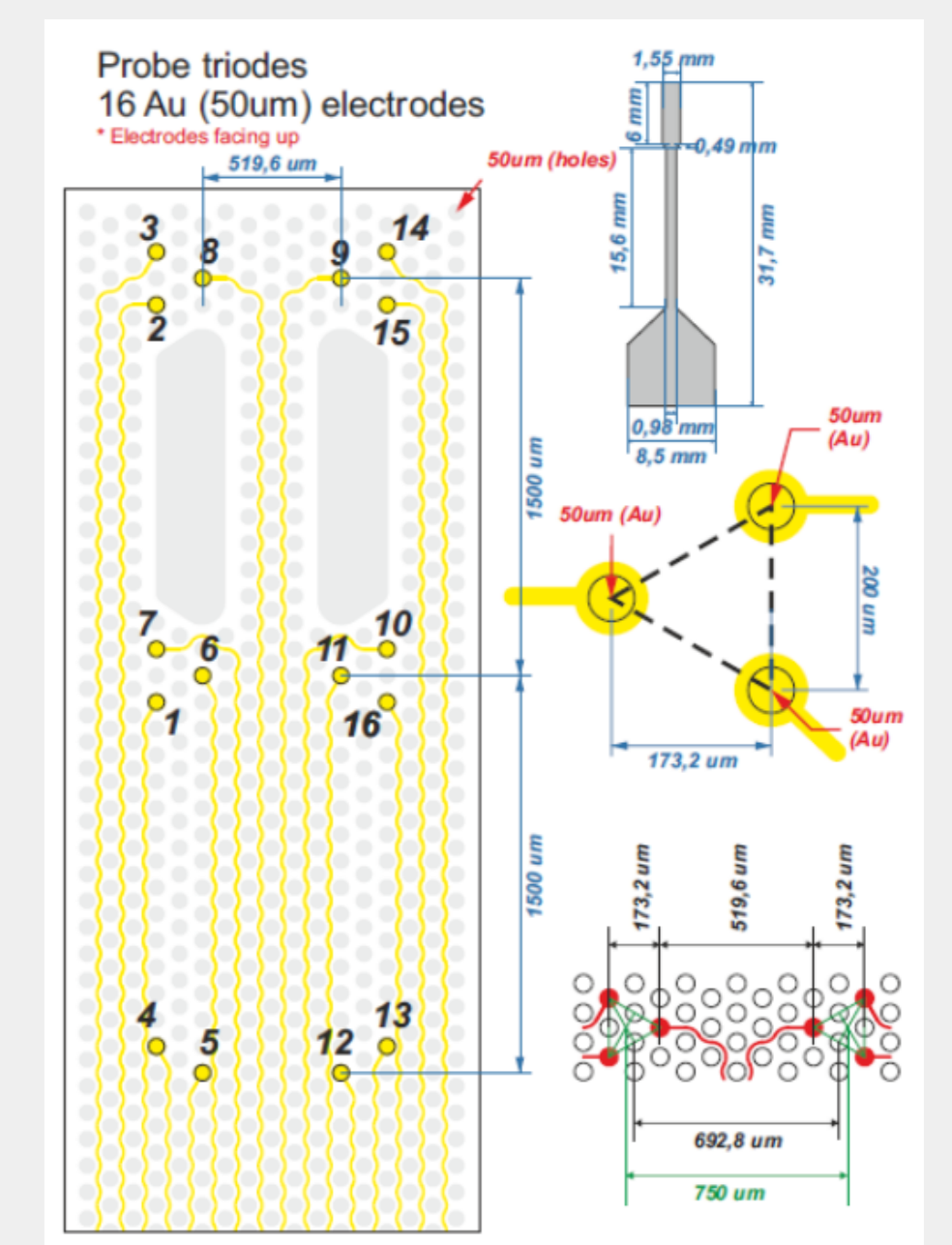


Figure 3: Schematic of the 16-channel SU-8-based flexible microprobe used for the recordings

→ Experimental model to explore the transitions from a state of slow oscillations towards a higher complexity state (awake-like asynchronous state).

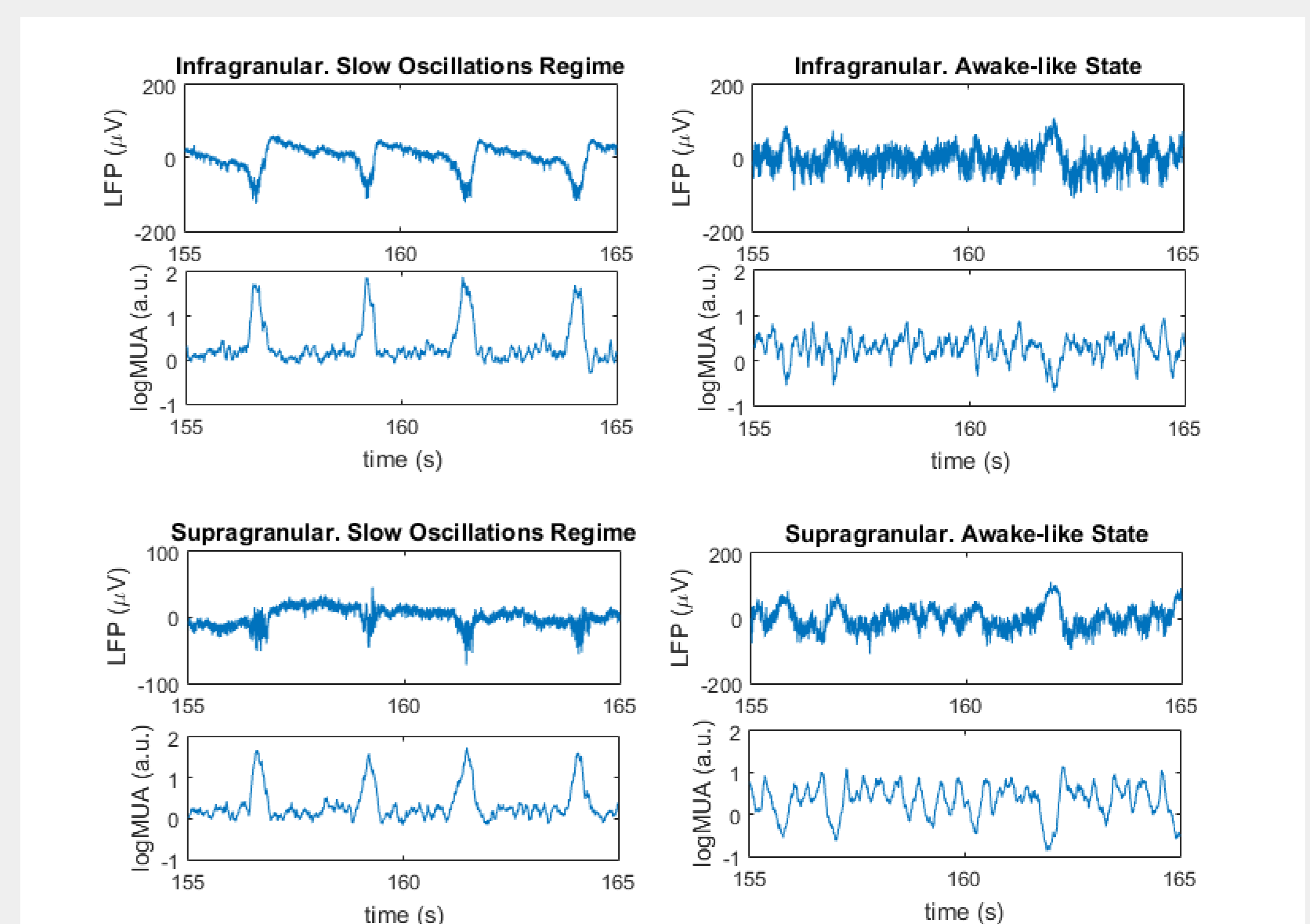


Figure 4: Example of extracellular activity (LFP and logMUA) issued from recordings at two different layers under two different experimental conditions

Our Approach

Aims

- ▶ To identify the **stability of the global network states** in isolated cortical networks under experimental manipulations that alter key network parameters (excitability, input, connectivity, etc.).
- ▶ To develop a novel theoretical tool which empirically captures the **metastable regions of the network**, i.e. transient states that temporarily behave as attractors.

Methodology

- ▶ With the aid of kernel mean embedding techniques for clustering [9], we will detect the convergence regions of the system.
- ▶ By studying how the phase portrait of the system evolves when the slow-oscillation regime is perturbed, we will map the bifurcations or transient states with the network parameters.