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Context

Brain circuits display **modular architecture** at different scales of organization [6]. Such neural assemblies are typically associated to functional specialization, but the mechanisms leading to their emergence and consolidation still remain elusive. In particular, it has been shown that **inhibitory** plasticity has a decisive impact on the dynamics and adaptation capacity of the brain, notably in the **long-term maintenance of memory** [4].

Although many studies using **coupled oscillator** models attempt to simulate this process, few consider the case of **distinct inhibitory and excitatory units** having an **adaptation mechanism** on their synaptic weights and thus impacting the network state.

Therefore, this study [1] investigates the impact of **inhibition** on the modulation of **synchronization** and on the creation and consolidation of **modular structures** in **coupled oscillator networks** subject to **adaptation**. These results are then extended to **spilling neural networks** with STDP.

Creating and maintaining modular structures

- Learning of **2 temporally alternating stimuli** applied to separate areas of the network.
- We observe the evolution of the network at rest over the short and long term in networks of: (1) excitatory and inhibitory neurons, (2) with only excitatory neurons, and (3) with unlabelled neurons (both excitatory and inhibitory).
- Without distinct inhibitory units, the 2 clusters created synchronize at rest and consequently are forgotten in the long term.
- The presence of inhibition allows the structure to consolidate, where each inhibitory neuron associates with a cluster.

Inhibitory T0	T1	T2	Т3

to adaptation. These results are then extended to spiking neural networks with STDP.

Model

Network of N θ -neurons all-to-all connected [2]

$$\frac{d\theta_i}{dt} = (1 - \cos(\theta_i)) + (1 + \cos(\theta_i)) \left[\eta_i + \frac{g}{N} \left(\sum_{j=1}^N \kappa_{ij} \sin(\theta_j - \theta_i) \right) + I_i(t) + \xi_i(t) \right]$$

With the phase $\theta_i \in [-\pi, \pi]$, the excitability parameter η_i , the global coupling g, κ_{ij} the coupling weight from pre-synaptic neuron j to post-synaptic neuron i, the external input $I_i(t)$ and the additive Gaussian noise $\xi_i(t)$.

Weights adaptation



$\frac{d\kappa_{ij}}{dt} = [\epsilon_1 + \epsilon_2 H(I_j(t))]\kappa_{ij}(1 - \kappa_{ij})\Lambda(\Delta\theta)$







 $\frac{d\kappa_{ij}}{dt} = \epsilon_1(\kappa_{ij} + 1)(-\kappa_{ij})\Lambda(\Delta\theta)$



With $\epsilon_1 \ll \epsilon_2 \ll 1$ the learning rates for slow and fast adaptations and H(x) the Heaviside function.

Symmetric phase difference-dependent plasticity rule [3, 5]

Δθ





Discussion

θ -neurons

- The creation of clusters is induced by the spatio-temporal correlations of the stimuli applied to the network and by the resulting adaptation mechanism, thus maintaining a certain synchrony within them.
- Networks made of purely excitatory neurons or of neurons not respecting Dale's principle are unable to maintain the modular architecture induced by the entrained stimuli.
- By preserving a specific dynamics to each cluster, **inhibitory** neurons allow to **maintain** and

Figure 2. Experiment with 2 clusters using θ-neurons. Graphs (a) represent the weight matrices at times T0, T1, T2 and T3: the color denotes if the connection is excitatory (red) or inhibitory (blue) or absent (white). Graphs (b) are raster plots at times T0, T1, T2 and T3, displaying the firing times of excitatory (red dots) and inhibitory (blue dots) neurons. Note that the inhibitory neurons are sorted by phases at time T3 in our model to better visualize clusters, otherwise the associations with each group are random. The neurons are also sorted by phases in each cluster for the last model.

Extension to spiking neurons

- Network of quadratic integrate and fire (QIF) neurons [2] subject to spike-timing-dependent plasticity.
- Similar protocol with learning of 2 temporally alternating stimuli applied to separate groups of excitatory and inhibitory neurons.
- At rest, we observe **asynchronous irregular dynamics**, with random epochs of **spontaneous recall** of the two learned items.



consolidate the learned structure over the **long-term** \rightarrow memory consolidation.

Spiking neurons

Feed-back and feed-forward inhibition favour the formation of the modular structures and their spontaneous recall during a resting state of asynchronous irregular dynamics.
Spontaneous recall of the learned items seems to promote their maintenance and consolidation over the long-term → memory consolidation.

Figure 3. Experiment with 2 clusters using QIF neurons. Graphs (a) represent the weight matrices at times T0, T1 and T2: the color denotes if the connection is excitatory (red) or inhibitory (blue) or absent (white). Graph (b) is the raster plot displaying the firing times of excitatory (red dots) and inhibitory (blue dots) neurons.

References

[1] Raphaël Bergoin, Alessandro Torcini, Gustavo Deco, Mathias Quoy, and Gorka Zamora-Lopez. Inhibitory neurons control the consolidation of neural assemblies via adaptation to selective stimuli. Scientific Reports, 13(1):6949, 2023.

[2] G Bard Ermentrout and Nancy Kopell. Parabolic bursting in an excitable system coupled with a slow oscillation. SIAM journal on applied mathematics, 46(2):233–253, 1986.

[3] Leonhard Lücken, Oleksandr V Popovych, Peter A Tass, and Serhiy Yanchuk. Noise-enhanced coupling between two oscillators with long-term plasticity. Physical Review E, 93(3):032210, 2016.

[4] Gianluigi Mongillo, Simon Rumpel, and Yonatan Loewenstein. Inhibitory connectivity defines the realm of excitatory plasticity. *Nature neuroscience*, 21(10):1463–1470, 2018.

[5] Jafar Shamsi, María José Avedillo, Bernabe Linares-Barranco, and Teresa Serrano-Gotarredona. Oscillatory neural networks. In 2020 XXXV Conference on Design of Circuits and Integrated Systems (DCIS), pages 1–6. IEEE, 2020.

[6] Gorka Zamora-López, Changsong Zhou, and Jürgen Kurths. Exploring brain function from anatomical connectivity. Frontiers in neuroscience, 5:83, 2011.