

# Modeling neural nets by interacting systems of chains with memory of variable length.

Joint work with A. Duarte, A. Galves, G. Ost

II. Neuromat Workshop, November 2016

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- Once the neuron has spiked, its potential is reset to a resting potential (here = 0). Then : Restart accumulating potentials coming from other neurons.
- This is what is called **Variable length memory** : the memory of a given neuron goes back in past up to its last spike.

- **Spike trains** : for each neuron  $i \in \mathcal{I}$ , we indicate if there is a spike or not at time  $t$ ,  $t \in \mathbb{Z}$ .

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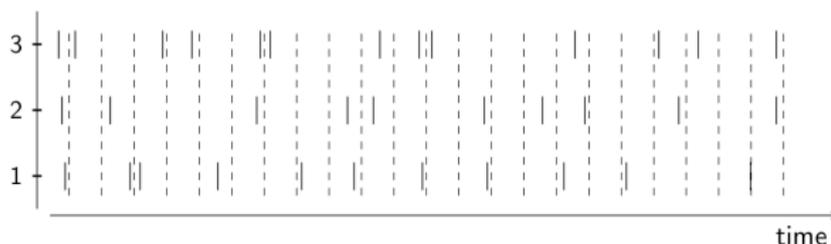


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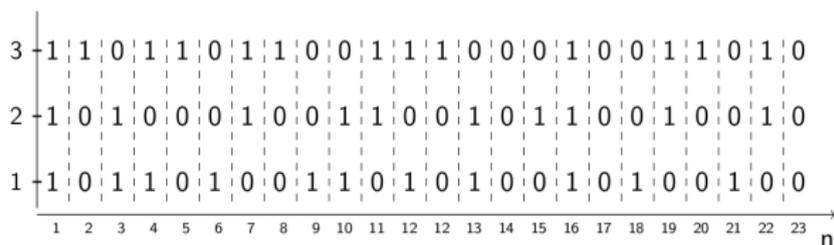
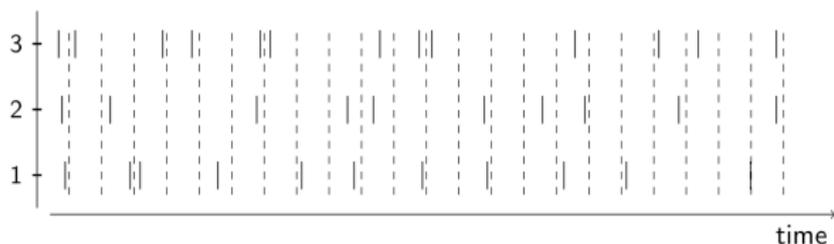


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- 2 these spikes are weighted by the synaptic weight  $W_{j \rightarrow i}$  of neuron  $j$  on neuron  $i$
- 3 they are also weighted by an aging factor which describes the loss of potential since the appearance of the spike of neuron  $j$  and the present time  $t$ .

The formula for the membrane potential of neuron  $i$  at time  $t$  :

$$U_t(i) = \sum_j W_{j \rightarrow i} \sum_{s=L_t^i+1}^{t-1} g_j(t-s)X_s(j),$$

where

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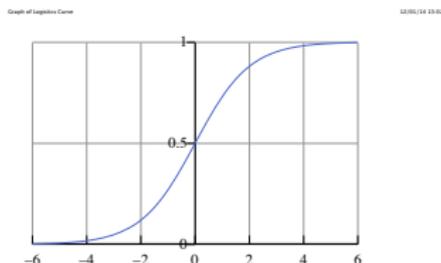
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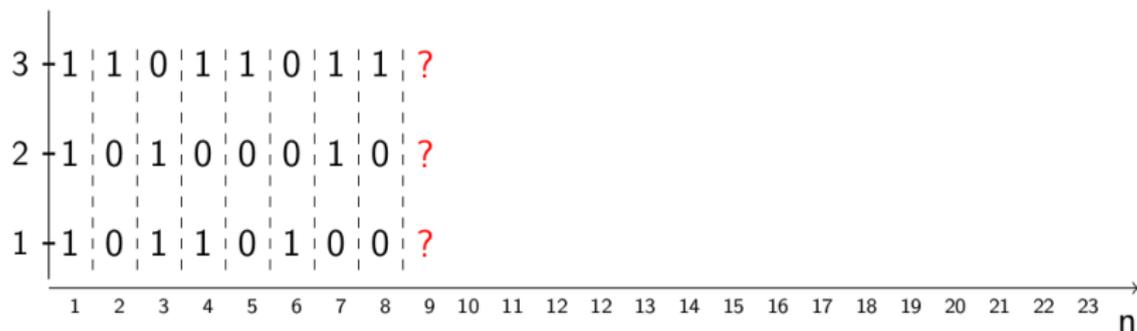
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Thus :  $X_{t-1}(i) = 1$  (spike at time  $t - 1$ )  $\Rightarrow U_t(i) = 0$ .
- $g_j : \mathbb{N} \rightarrow \mathbb{R}_+$  describes a leak effect.

$$P(i \text{ spikes at time } t) = \Phi_i(U_t(i) + S_t(i)),$$

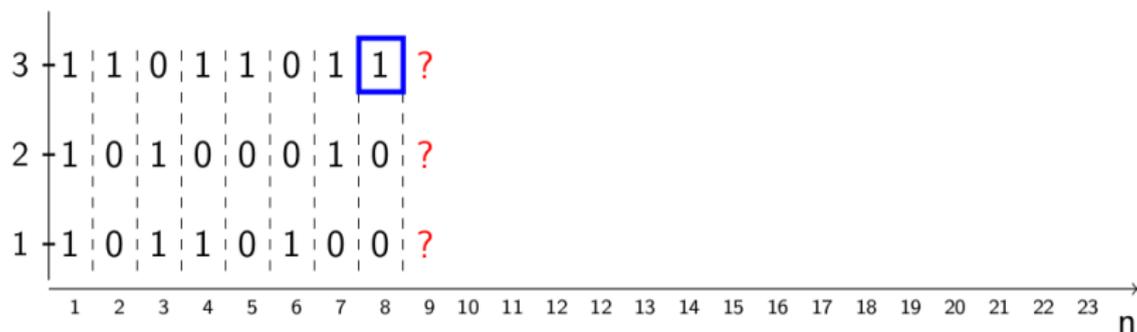
- $\Phi_i$  spiking rate function of neuron  $i$  : this is an **increasing** function.

It can have a logistic shape.



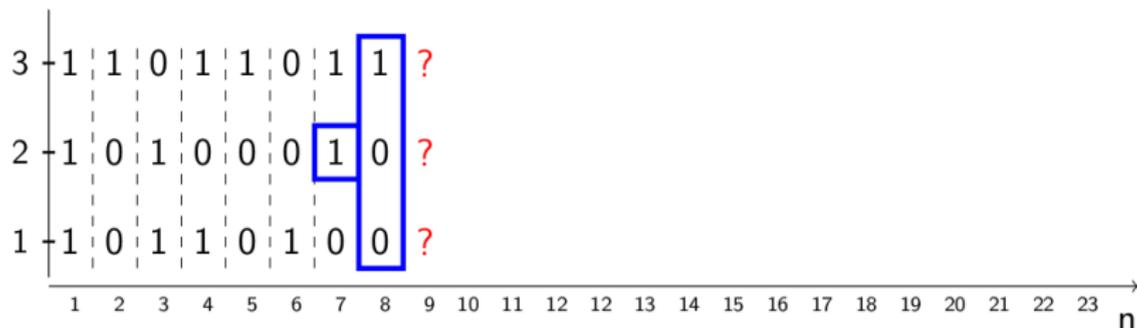


$$P(X_9(i) = 1 | \text{past}) = \begin{cases} \phi_i(0), & \text{if } L_9^i = 8 \\ \phi_i\left(\sum_{j \in I} W_{j \rightarrow i} \sum_{s=L_9^i+1}^8 g_j(t-s) X_s(j)\right), & \text{otherwise} \end{cases}$$



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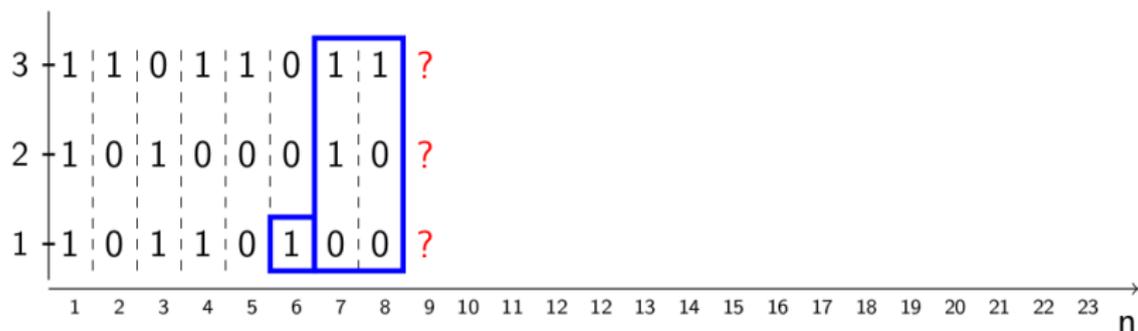
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$$L_9^1 = 6 \implies P(X_9(1) = 1 | \text{past}) = \phi_1(W_{3 \rightarrow 1}(g_3(1) + g_3(2)) + W_{2 \rightarrow 1} g_2(2))$$

# Comparison with other models

**Our model is a generalization of the classical LIF model, with random thresholds  $\Theta_t(i)$ ,  $t \in \mathbb{Z}$ ,  $i \in I$  which are i.i.d.**

$$\phi_i(U_t(i)) = P(U_t(i) > \Theta_t(i))$$

is the probability that the membrane potential  $U_t(i)$  exceeds the random threshold  $\Theta_t(i)$ .

# Comparison with other models

If we choose  $g_j(1) = 1, g_j(n) = 0$  for all  $n \geq 2$  and  $\Phi_i(x) = x$ , then our model is (almost) the

## Kinouchi-Copelli model with only one refractory period :

- each neuron has two states : passive (0) or active (spiking, 1)
- if  $j$  has just spiked, then  $i$  has a **transition from 0  $\rightarrow$  1** with probability  $W_{j \rightarrow i}$ .

## About the model

- ▶ Introduced in our joint paper with Antonio Galves in 2013.
- ▶ The spiking probability of neuron  $i$  depends on the activity of the system since the last spike time.
- ▶ The chain  $(X_t(i))_{t \in \mathbb{Z}}$  is a **chain with memory of variable length**.
- ▶ The model is a **system of interacting chains with memory of variable length**.

## More about the model

- Study of the model in **continuous time**, mean-field approximation and **Propagation of chaos** :  
De Masi, Galves, L., Presutti (2015), Fournier and L. (2016), Robert and Touboul (2014), Duarte, Ost and Rodriguez (2016) for a spatially structured model, Drougoul and Veltz (2016), Brochini, Costa, Abadi, Roque, Stolfi and Kinouchi (2016).
- Duarte and Ost (2016) : Finite systems of interacting neurons without external stimulus and with some leak effect stop spiking ...
- Estimation of the spiking rate function : Hodara, Krell, L. (2016)

# The interaction graph

Neurons who have a direct influence on  $i$  are those belonging to

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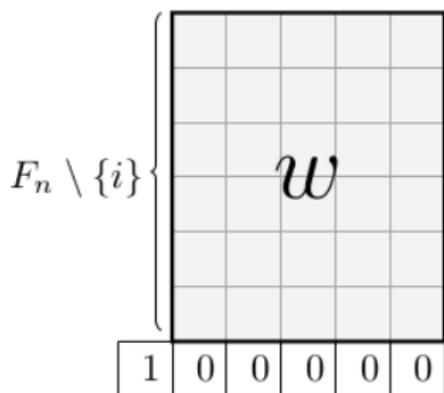
Goal of this talk : Estimate the **Interaction neighborhood**  $\mathcal{V}_i$  of a fixed neuron  $i$ — based on an observation of the process in discrete time  $[1, \dots, n]$ , within growing spatial windows.

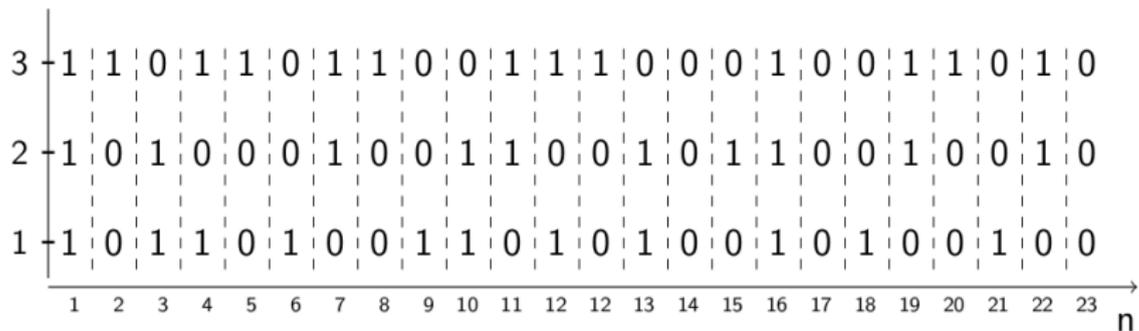
# Estimation procedure

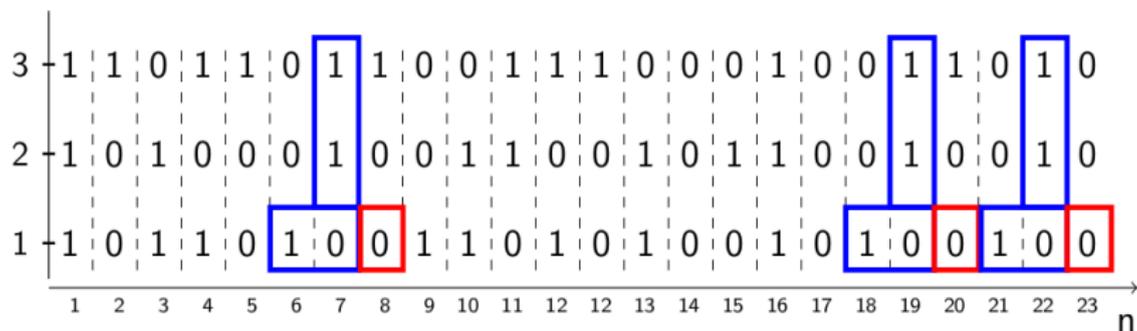
AIM : Estimate  $\mathcal{V}_i$ !

- Growing sequence of finite windows  $F_n$  - centered around site  $i$ .
- For a test-raster block  $w \in \{0, 1\}^{\{-\ell, \dots, -1\} \times F_n \setminus \{i\}}$  :

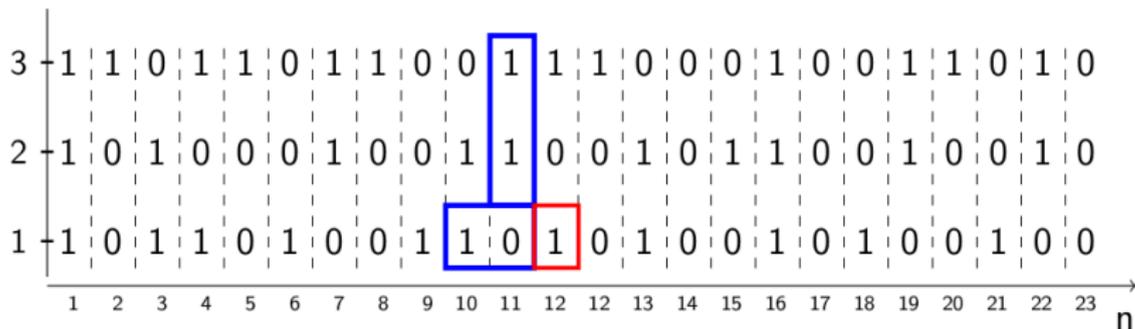
$N_{(i,n)}(w, 1)$  counts the number of occurrences of  $w$  followed by a spike of neuron  $i$  in the sample  $X_1(F_n), \dots, X_n(F_n)$ , when the last spike of neuron  $i$  has occurred  $\ell + 1$  time steps before in the past.



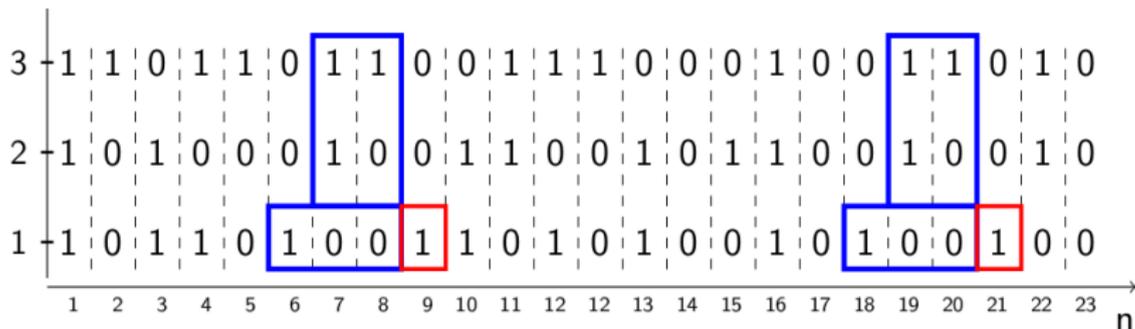




$$N_{(1,23)}(w, 0) = \sum_{m=3}^n 1\{X_{m-2}^{m-1}(1) = 10, X_{m-1}(F_n \setminus \{1\}) = w, X_m(1) = 0\}.$$



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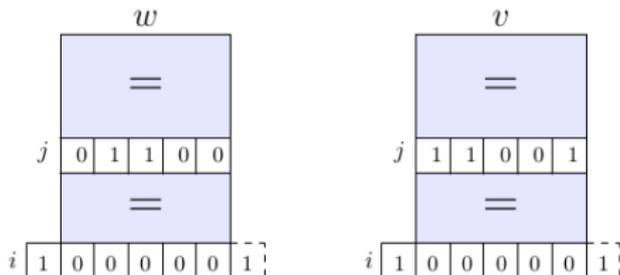


$$N_{(1,23)}(w, \mathbf{1}) = \sum_{m=4}^n \mathbb{1}\{X_{m-3}^{m-1}(1) = 10^2, X_{m-2}^{m-1}(F_n \setminus \{1\}) = w, X_m(1) = \mathbf{1}\}.$$

- Estimated spiking probability  $\hat{p}_{(i,n)}(1|w) = \frac{N_{(i,n)}(w,1)}{N_{(i,n)}(w)}$ .

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- Test statistics to test the influence of neuron  $j$  on neuron  $i$  :

$$\Delta_{(i,n)}(j) = \max_{w,v: v_{\{j\}^c} = w_{\{j\}^c}} |\hat{p}_{(i,n)}(1|w) - \hat{p}_{(i,n)}(1|v)|.$$



## Definition

For any positive threshold parameter  $\epsilon > 0$ , the *estimated interaction neighborhood of neuron  $i \in F_n$* , at accuracy  $\epsilon$ , given the sample  $X_1(F_n), \dots, X_n(F_n)$ , is defined as

$$\hat{V}_{(i,n)}^{(\epsilon)} = \{j \in F_n \setminus \{i\} : \Delta_{(i,n)}(j) > \epsilon\}.$$

## Conditions

1) Spiking rate functions  $\phi_i$  are *strictly increasing*, Lipschitz

$$|\phi_i(z) - \phi_i(z')| \leq \gamma |z - z'|$$

and bounded from above and below. 2) Uniform summability of the synaptic weights

$$r := \sup_i \sum_j |W_{j \rightarrow i}| < \infty.$$

3)  $\sup_i g_i(n) < \infty$  for all  $n$ .

## Theorem

Fix  $i \in \mathcal{I}$  and suppose :

- ▶  $V_i$  finite and
- ▶  $|F_n| = o(\log n)$ .

Let  $X_1(F_n), \dots, X_n(F_n)$  be a sample produced by the stochastic chain  $(X_t)_{t \in \mathbb{Z}}$  satisfying our assumptions. Then for  $\epsilon_n = O(n^{-\xi/2})$ , for some  $\xi > 0$ ,

$$\hat{V}_{(i,n)}^{(\epsilon_n)} = V_i \text{ eventually almost surely.}$$

Can be extended to the case when the interaction neighborhood of  $i$  is infinite !

## Remarks

- ▶ We are not estimating the weights of interaction, only the existence or not of interactions. (An oriented graph which is not weighted! )
- ▶ We are able to estimate the existence of interactions without assuming the knowledge of the spiking probability functions or the leak functions.
- ▶ We can deal with the case in which the system has several invariant states.

# Open questions

- Multi-unit recordings of spiking activity show only a picture of a very tiny part of the brain ...
- What does the observation of a very small part of the net tells us about the global behavior of the system ?

- In this talk : estimation of the **anatomical graph of interactions** between single neurons.
- What about the graph describing the **functional interactions** between components or regions of the brain ?
  - Usually derived from **correlations**
  - absence of correlations is not equivalent with independence!!!! (this would only be true for Gaussian systems)
  - need : **new mathematical definition of functional dependence.**

- Take into account the **characteristics of the graph of interactions (anatomic and/or functional) of the brain.**
- Types of graphs that have been considered : critical Erdős-Rényi random graphs, small world, rich-club networks, ...
- **Statistical model selection for systems of spiking neurons with interaction graphs belonging to one of these classes ?**

# Study of the relaxation period

- **Relaxation period** = interval during which the system transits from an initial condition to an asymptotic state (= stable state).
- Study how the limit **law of the last spiking time** (which is finite, see Duarte and Ost (2016)), rescaled by its mean value, depends on the **leak rate for a fixed spiking rate function**.
- Is there a notion of **criticality** as in Kinouchi and Copelli (Optimal Dynamical Range of Excitable Networks at Criticality, 2006) - and an associated **dynamical phase transition** ?

# Some literature

Paper is on arXiv :

- DUARTE, A., GALVES, A., L.E., OST, G., Estimating the interaction graph of stochastic neural dynamics. 2016, arXiv.

