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ICMNS 2018 - ANTIBES Synchronization of stochastic mean field networks of Hodgkin-Huxley neurons with noisy channels

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The model

V^{*t*} is the voltage of the neuron, m_t , h_t , $n_t \in [0, 1]$, are the proportions of open: activation Sodium channels, deactivation Sodium channels and activation Potassium channels respectively.

 $dV_t = F(V_t, m_t, n_t, h_t)dt$

According to the Hodgkin-Huxley model, the dynamics of a neuron's voltage is the result of the passage of ions through its membrane. This ion flux occurs through specific proteins which act as gated channels.

 $F(V, m, n, h) = I - g_{K}n^{4}(V - V_{K}) - g_{Na}m^{3}h(V - V_{Na})$ $-q_L(V-V_L)$, (1)

represents the effect on the voltage of the ionic channels and of an external current *I* (assumed constant for simplicity).

Single Neuron Model

We consider a population of *N* neurons $\{1, \ldots, i, \ldots, j, \ldots, N\}$, and two type of interactions:

With each neuron *j* we associate a new variable $y^{(j)} \in [0, 1]$ which represents its proportion of open synaptic channels at each time. Chemical synapses coming from neuron j induce on the voltage $V^{(i)}$ of the neuron *i* an instantaneous variation at time *t* of

Pre-synaptic neuron *j* contributes to the variation of the voltage of post-synaptic neuron *i* by the amount

*− J*E *N* (*V* (*i*) $s^{(i)}-V$ (*j*) ${s^{(J)}\choose s},$

where $J_{\rm E} \geq 0$ is the electrical conductance.

Interaction: Synapses

For the specific form of σ_x we refer to Pakdaman et al. [2] and the references therein, where the authors obtain a similar dynamic for the channels as the fluctuations of a Piecewise Deterministic Markov Process.

We consider the model for a network fully connected studied by Bossy et al. in [1]. The state of the neuron $i = 1, \ldots, N$ will represented by *X* (*i*) $t^{(\ell)}_t=(V)$ (*i*) $\tilde{t}^{(\iota)}, m$ (*i*) $\overset{\leftarrow}{t}^{\prime},n$ (*i*) $\boldsymbol{h}_t^{(i)}, \boldsymbol{h}_t^{(i)}, y$ (*i*) $t^{(l)}(t)$, and we denote by

Chemical Synapses: A neurotransmitter is released to the intercellular media from a pre-synaptic neuron to the post-synaptic one through synaptic channels.

 $\bar{V}_t^N =$ 1 *N* \sum *i*=1 *V* (*i*) $\bar{y}^{(i)}_t, \;\; \bar{y}^N_t =$ 1 *N* \sum *i*=1 *y* (*i*) *t ,*

where $(W^{x,i} : i \in \mathbb{N}, x = m, n, h, y)$ are one dimensional Brownian motions independent of each other and independent of the initial condition.

$$
-\frac{J_{\rm Ch}}{N}y_t^{(j)}(V_t^{(i)}-V_{\rm rev}),
$$

where $J_{\text{Ch}} \geq 0$ is the chemical conductance.

Electrical Synapses: Neurons directly connected with each other

through an intercellular channel called gap junction.

Numerical simulations show when increasing $J_{\rm E}$, the neurons synchronize. This phenomenon is independent of the number of neurons and the phase difference is only affected by the size of the noise.

Noise

Figure 2: Trajectories for the Voltage of 100 neurons under strong interaction and different level of noise: *Left:* $\sigma = 0.5$. *Right:* $\sigma = 1$.

To consider the intrinsic noise present in the ion and neurotransmitter channels we add a noise to the dynamic of the channels:

 $dx_t = \rho_x(V_t)(1 - x_t) - \zeta_x(V_t)x_t dt + \sigma_x(V_t, x_t)dW_t^x,$

where W^x , $x = m, n, h, y$ are independent Brownian motions.

As a measure of the synchronicity of the network we compute the empirical variance of Voltage of the system (2) : $N^{-1} \sum_{i=1}^{N} (V - 1)$ $\bar{V}_t^{(i)} - \bar{V}_t^N)^2,$ and to obtain statistically meaningful information, we perform Monte Carlo simulations to estimate the evolution of the expected value of this quantity for different values of N and σ .

 $M_{\rm A}$ and $A_{\rm A}$ and $A_{\rm 100}$ 0 500

Figure 3: Time evolution of the expected empirical variance of the Voltage for different level of noise: *Left:* $\sigma = 0.5$. *Right:* $\sigma = 1$.

Model for a Network

Partial Conclusion: The numerical evidence shows that for big enough $J_{\rm E}$ the neurons get synchronized, meaning that the empirical variance becomes small in mean. This phenomena seems independent of the size of the network.

For $t_1 \geq 0$, let $(\widehat{X}_t^{t_1} : t \geq t_1)$ denote the solution of the ordinary differential equation

> $dV_t =$ $\sqrt{2}$ $F(V_t, \hat{m})$ $\widehat{m}_t, \widehat{n}$ \widehat{n}_t, h_t) *− J*_{Ch} \widehat{y} y_1 $\tilde{t}(V_t - V_{\text{rev}})$] *dt,* $d\widehat{x}_t =$ $\overline{\Gamma}$ $\rho_x(V_t)(1 - \hat{x})$ $\boldsymbol{\psi}$ $f(t) - \zeta_x(V_t)\hat{x}_t$] *dt*

N N

We denote $(\hat{X}_{\underline{t}}^{t_1,E}: t \geq t_1)$ the solution of (3) with random initial condition $\widehat{X}_{t_1}^{\overline{t_1},E}$ $=\bar{X}^N_{t_1}$ *t*1 $:= (\bar{V}^N_{t_1})$ *t*1 $, \bar{m}^N_{t_1}$ *t*1 $, \bar{n}_{t_1}^N$ *t*1 $, \bar{h}^N_{t_1}$ *t*1 $, \bar{y}_{t_1}^N$ *t*1)*.*

the empirical means for the voltage and the proportion of open neurotransmitter channels respectively.

The dynamics will be:

V (*i*) $t^{(t)} = V$ (*i*) $0^{(t)} +$ \int_0^t $\overline{0}$ *F*(*V* (*i*) $s^{\left(\iota\right)},m$ (*i*) $\binom{l}{s},n$ $\binom{(i)}{s},h_{S}^{(i)})ds$ *−* \int_0^t $\overline{0}$ $J_{\rm E}$ ⁽ V $\bar{V}^{(i)}_s - \bar{V}^N_s$ $\big) - J_\mathrm{Ch} \bar y_s^N$ *s* (*V* $V_{\rm se}^{(i)}-V_{\rm rev}\big)ds,$ *x* (*i*) $t^{(t)}_t = x$ (*i*) $_{0}^{\left(\nu \right) }$ + \int_0^t $\overline{0}$ $\rho_x(V)$ (*i*) $J_s^{(\iota)}$ $(1 - x)$ (*i*) $\binom{v}{s}$ *−* $\zeta_x(V)$ (*i*) ${s^{(\iota)}\choose s}x$ (*i*) \int_{s}^{t}/ds $+$ \int_0^t $\overline{0}$ *σx*(*V* (*i*) $s^{\left(\imath\right)},x$ $\int_s^{(i)}$) $dW^{x,i}_s$, (2)

with $\chi : \mathbb{R} \to [0,1]$ a Lipschitz continuous function with support contained in [0, 1] and $\sigma \geq 0$. 2. The *leak conductance gL* is strictly positive. 3. One has (*m* (*i*) $\binom{0}{0},n$ (*i*) $_0^{(\iota)}, h$ (*i*) $_0^{(\iota)},y$ (*i*) $\binom{1}{0}$ $\in [0, 1]^4$ a.s. 4. The initial voltages are bounded uniformly in *N*:

> sup *i*=1*,...,N* $|V|$ (*i*) $|V_0^{(i)}| \leq V_0^{\max}$ $\begin{array}{cc} \eta \text{max} & a.s. \end{array}$

Simulating the Model

 $\lambda^0 > 0$ not depending on $N \geq 1$, $\sigma \geq 0$ or X_0 , and there exists a time $t_0 \geq 0$ not depending on $N \geq 1$ or $\sigma \geq 0$, such that for each $J_{\rm E} > J_{\rm E}^0$ E^U the solution *X* of (2) satisfies, for every $t \geq t_0$ and each *i ∈ {*1*, . . . , N}*:

Figure 1: Left: The evolution of the Voltage and Channels for one neuron. *Right:* The evolution of the Voltage of 100 neurons $J_{\rm E}$ small.

Emergence of Synchrony

a) There exists $(\mu_t : t \ge 0)$ in $C(\mathbb{R}^+; \mathcal{P}_2(\mathbb{R} \times [0, 1]^4))$ a global solution (in the sens of distribution) of the non linear McKean-Vlasov Fokker Planck equation

Time [ms]

 $W_2^2(\mu_t, \delta_{\langle \mu_t \rangle}) \le W_2^2(\mu_{t_0}, \delta_{\langle \mu_{t_0} \rangle})e^{-\lambda^0(t-t_0)} + \sigma^2$ C^0_{ζ} *ζ,ρ λ* 0 *.* In particular, $\limsup_{t\to\infty} \mathcal{W}_2^2(\mu_t, \delta_{\langle \mu_t \rangle}) \leq \sigma^2 \frac{C_{\zeta}^0}{\lambda^0}$ *ζ,ρ* $\frac{\partial \zeta, \rho}{\partial \lambda^0}.$ b) For every $t_1 \geq t_0$ and $\delta \geq 0$ we have: sup *t*1*≤t≤t*1+*δ* $\mathcal{W}_2^2(\mu_t, \delta_{\widehat{X}})$ *t*1 *,∞ t*) *≤ K*0 *∧* 2 $\sqrt{2}$ $(K'_0e^{-\lambda_0(t_1-t_0)} + \sigma^2)$ C^{0}_{ζ} *ζ,ρ λ* 0 $(1 + \delta K_{\delta})$] *.*

[2] K. Pakdaman, M. Thieullen, and G. Wainrib. Fluid limit theorems for stochastic hybrid systems with aplication to neuron models. *Adv. in Appl. Probab.*, 42(3):761–794, 09 2010.

0 20 40 60 80 100 Time [ms]

0

500

Reduce Synchronized Dynamics

(3)

Empirical Case

Mean Field Case

For $(\mu_t : t \ge 0) \in C([0, \infty), \mathcal{P}_1(\mathbb{R} \times [0, 1]^4))$ we denote by $(\widehat{X}^{t_1,\infty}_t : t \geq t_1)$ the solution of (3) with deterministic initial condition $\widehat{X}_{t_1}^{t_1,\infty} = \langle \mu_{t_1} \rangle$, where $\langle \mu_{t_1} \rangle$ is the vector of means of μ_{t_1} .

Hypothesis (H)

1. ρ_x and ζ_x are strictly positive, locally Lipschitz continuous functions on \mathbb{R} . The function $\sigma_x : \mathbb{R}^2 \to \mathbb{R}$ are given by

> $\sigma_x(v,z)=\sigma$ √ $|\rho_x(v)(1-z) + \zeta_x(v)z|\chi(z),$

Main Results

Theorem 1: Synchronization

Assume (H) and that the initial condition is an exchangeable random vector.

a) **Synchronization.** There exist constants $J_{\rm E}^0 > 0$, $C_{\zeta,\rho}^0 > 0$ and

$$
\mathbb{E}\left(|X_t^{(i)} - \bar{X}_t^N|^2\right) \le \mathbb{E}\left(|X_{t_0}^{(i)} - \bar{X}_{t_0}^N|^2\right) e^{-\lambda^0(t - t_0)} + \sigma^2 \frac{\tilde{\zeta}, \rho}{\lambda^0}
$$

In particular, $\limsup_{t \to \infty} \mathbb{E}\left(|X_t^{(i)} - \bar{X}_t^N|^2\right) \le \sigma^2 \frac{C_{\zeta, \rho}^0}{\lambda^0}.$

b) **Synchronized dynamics.** Assume $J_{\text{E}} > J_{\text{E}}^{0}$ E . Then, there are constant $K_0, K'_0 > 0$ depending only on the parameters of the voltage dynamics in (1) and, for each $\delta \geq 0$, constants $K_{\delta}, K'_{\delta} > 0$ depending on the coefficients in (2) and on δ (increasingly) but not on *N*, such that for every $t_1 \geq t_0$ and each $i \in \{1, \ldots, N\}$:

.

sup *t*1*≤t≤t*1+*δ* \mathbb{E} $\sqrt{2}$ *|X* (*i*) $\hat{X}_t^{(i)} - \hat{X}_t^{t_1,E} |^2$ *≤ K*0 *∧* 2 \lceil $K'_0 e^{-\lambda_0 (t_1 - t_0)} + \sigma^2$ C^0_ζ *ζ,ρ λ* 0 \setminus $(1 + \delta K_{\delta}) + \delta K_{\delta}'$ *σ* 2 *N* $C^0_\mathcal{C}$ *ζ,ρ*]

.

 $dx_t = [\rho_x(V_t)(1 - x_t) - \zeta_x(V_t)x_t] dt$

where, here and in the sequel, x generically represents the m, n, h components and $F : \mathbb{R} \times [0,1]^4 \to \mathbb{R}$, defined by

Theorem 2: Mean Field Limit

Assume (H) and that for all $N \geq 1$ the initial condition are i.i.d. random vectors with (compactly supported) common law $\mu_0 \in \mathcal{P}(\mathbb{R} \times$ $[0, 1]^4$ not depending on N.

Consider the notations:

$$
\Phi(w, z, v, u) := F(v, u_m, u_n, u_h) - J_E(v - w) - J_{Ch}z(v - V_{rev})
$$

\n
$$
b_x(v, u) := \rho_x(v)(1 - u_x) - \zeta_x(v)u_x.
$$

\n
$$
a_x(v, u) := (\rho_x(v)(1 - u_x) + \zeta_x(v)u_x)\chi(u_x)
$$

\n
$$
\langle \mu \rangle := (\langle \mu^V \rangle, (\langle \mu^x \rangle)_{x = m, n, h, y})
$$

$$
\partial_t \mu_t = \partial_v \left(\Phi(\langle \mu_t^V \rangle, \langle \mu_t^y \rangle, \cdot, \cdot) \mu_t \right) + \sum_{x=m,n,h,y} \frac{1}{2} \sigma^2 \partial_{u_x u_x}^2 (a_x \mu_t) - \partial_{u_x} (b_x \mu_t)
$$

with initial condition μ_0 .

for each $T > 0$, the process of empirical measures of the system (2) $(\mu_t^N = N^{-1} \sum_{i=1}^N \delta_i)$ $\mathbf{X}_t^{(i)}$: $t \in [0,T]$ converges in law *t* on $C([0,T]; \mathcal{P}_2(\mathbb{R} \times [0,1]^4))$, when *N* tends to infinity, to a deterministic and uniquely determined flow of probability measures $(\mu_t : t \in [0, T])$ defined on part a).

c) There is a constant $C(T) > 0$ depending on V_0^{max} U_0^{max} , $T>0$ and on the coefficients of system (2) , but not on *N*, such that

> sup *t∈*[0*,T*] \mathbb{E} $\left(\mathcal{W}_2^2(\mu_t^N)\right)$ $_{t}^{IV},\mu_{t})$ \setminus *≤ C*(*T*)*N −*2*/*5 *,*

where \mathcal{W}_2 is the Wasserstein distance of order 2.

Synchronization + Mean Field Limit

Under the assumptions of Theorem 2 and for the same constants as in Theorem 1, whenever $J_{\rm E} > J_{\rm E}^0$ E we have:

a) For every $t \geq t_0$,

References

[1] M. Bossy, O. Faugeras, and D. Talay. Clarification and complement to "mean-field description and propagation of chaos in networks of hodgkin–huxley and fitzhugh–nagumo neurons". *The Journal of Mathematical Neuroscience (JMN)*, 5(1):1–23, 2015.