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Attractor neural networks with activity-dependent synapses: The role of synaptic facilitation

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Abstract

We studied an autoassociative neural network with dynamic synapses which include a facilitating mechanism. We have developed a general mean-field framework to study the relevance of the different parameters defining the dynamics of the synapses and their influence on the collective properties of the network. Depending on these parameters, the network shows different types of behaviour including a retrieval phase, an oscillatory regime, and a non-retrieval phase. In the oscillatory phase, the network activity continuously jumps between the stored patterns. Compared with other activity-dependent mechanisms such as synaptic depression, synaptic facilitation enhances the network ability to switch among the stored patterns and, therefore, its adaptation to external stimuli. A detailed analysis of our system reflects an efficient—more rapid and with lesser errors—network access to the stored information with stronger facilitation. We also present a set of Monte Carlo simulations confirming our analytical results.

1. Introduction

Associative memory illustrates the simplest possible manner in which collective neural computation can work. The basic problem is to store a set of different patterns in a recurrent network, in such a way that when a new stimulus is presented, the network dynamics drives the system to the most similar—according to a previously defined distance—among all different ones. Thus, the recall of stored patterns during the dynamics appears, and the patterns are the attractors of the dynamics. The number of different attractors determines the number of patterns that the system can store and retrieve. Many different variants, including asymmetric synapses, have been studied in the last decades \cite{2,3}. In these models, after learning, neurons are considered as the only dynamical variables. That is, synapses are static, with fixed strengths. However, it has recently been emphasized that a neural dynamics with

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necessary to produce an appreciable synaptic plasticity [1] which is fundamental for the development and adaptation of the nervous system, and they are believed to be on the basis of higher functions as learning and memory.

2. The model

Our starting point is a network of $N$ binary neurons $s_i = 1, 0$ representing, respectively, firing or silent states. Each neuron follows the probabilistic dynamics:

\[ \text{Prob}(s_i(t+1) = 1) = \frac{1}{2}[1 + \tanh(2\beta h_i(t))]. \]  

(1)

$\beta = 1/T$ where $T$ is the temperature, and $h_i(t)$ represents the total presynaptic current arriving to the postsynaptic neuron $i$. In order to consider dynamic synapses with facilitation and/or depression, we assume $h_i(t) = \sum_{j=1}^{N} o_{ij} x_j(t) + \theta_i$. Here, $\theta_i$ is the threshold of neuron $i$ to fire, and $o_{ij}$ are the static synaptic weights where the system stored patterns of information during a previous slow learning process. The stored patterns are codified in states of neural activity, namely, $\xi^v = \{\xi_j^v\}, v = 1, \ldots, P$. For instance, one can choose the standard covariance rule, namely, $\omega_v = 1/(Nf(1-f))\sum_{j=1}^{P}(\xi_j^v - f)(\bar{\xi}_j^v - f)$ with $\bar{\xi}_j^v = f$. The complete dynamics for depression $D_j(t)$ and facilitation $F_j(t)$ was reported in [7]. Here, we use a simplified version of such model in which $D_j(t) \equiv x_j(t)$ and $F_j(t) \equiv U + (1 - U)x_j(t)$, being $x_j(t)$ the fraction of neurotransmitters which are in a recovered state. A fraction of these neurotransmitters, namely, $U x_j(t)$, is ready to be released after the arrival of a presynaptic action potential ($s_j = 1$). The remaining, $(1 - U)x_j(t)$, can also be released by facilitating mechanisms whose dynamics is driven by the variable $u_j(t)$. For simplicity we assume that the complete dynamics is described by the discrete system

\[ x_j(t+1) = x_j(t) + \frac{1 - x_j(t)}{\tau_{rec}} - U x_j(t) s_j(t) \]

\[ u_j(t+1) = u_j(t) - \frac{u_j(t)}{\tau_{fac}} + U[1 - u_j(t)] s_j(t), \]

(2)

where $\tau_{rec}$ and $\tau_{fac}$ are the time constant for depressing and facilitating mechanisms, respectively.

3. Results

We solved the system described by Eqs. (1)-(2) within the standard mean field assumption $s_j \approx \langle s_j \rangle$ and in the limit of only one stored pattern, $x \equiv P/N = 0$ and $f = \frac{1}{2}$, although most of our conclusions are also valid for many patterns. The result is a discrete map $y_{j+1} = F(y_j)$ where $y \equiv \{m_+, m_-, x_+, x_-, u_+, u_+\}$ is a vector whose components are order parameters which measure the overlap with the stored pattern ($m$), the mean depression level ($x$) and the mean facilitation level ($u$), in the neurons that are, respectively, active (+) or inactive (−) in the pattern. The local stability of the steady state solutions can be studied by analysing the behavior of the eigenvalues, namely $\lambda_j$ associated to the local dynamics of this map (see Ref. [6] for details). In particular, fixed points become unstable when the maximum absolute value of all eigenvalues, namely $|\lambda|_{\text{max}}$ is bigger than one. Fig. 1(left) shows $|\lambda|_{\text{max}}$ as a function of $\tau_{rec}$ for $U = T = 0.1$ and $\tau_{fac} = 20$ ms. Then, the analysis of the stability of fixed points reveals three different regimes in the behaviour of the system. First, a ferromagnetic-like phase associated to the stored pattern $P$ appears for $\tau_{rec} < \tau_{rec}^*$. Second a paramagnetic-like or non-memory phase occurs for $\tau_{rec} > \tau_{rec}^*$. Finally, an oscillatory phase in which the network activity is jumping.

![Fig. 1](image-url)
between different memories appears for $\tau_{\text{rec}}^{} < \tau_{\text{rec}} < \tau_{\text{rec}}^*$. Fig. 1(right) shows the emergence of these three phases when one continuously varies $\tau_{\text{rec}}$ between 3 and 18 during 3 s. Fig. 2 shows phase diagrams obtained by plotting the critical lines at which transitions between these three phases occur for different values of the parameters $\tau_{\text{rec}}$, $\tau_{\text{fac}}$ and $U$. By inspection of these diagrams, one observes that the width of the oscillatory phase enlarges for increasing values of $\tau_{\text{fac}}$. Moreover, a detailed analysis of the oscillatory phase shows that the access to the stored memories and the error in the retrieval of such memories strongly depends on facilitation and on its competition with depression. This is shown in Fig. 3 where the half period of the oscillations in the overlap with a pattern, $m \equiv m_+ - m_-$, and its maximum absolute value are represented as function of $\tau_{\text{fac}}$, for different values of $\tau_{\text{rec}}$ and $U = T = 0.1$.

4. Discussion

Using a realistic model of dynamical synapses which includes short-term facilitation and depression, we show that, depending on the competition between these two mechanisms, different type of behaviour occurs in ANN. For low depression, a ferromagnetic or memory phase occurs. For very large depression or facilitation, a paramagnetic or non-memory phase occurs. For intermediate facilitation and/or depression, an oscillatory phase with the network activity jumping between the attractors shows up. We also observed that a high facilitation enhances the network ability to switch among the stored patterns, as well as its adaptation to external stimuli, as it is reported with more detail in [6]. Other interesting new phenomena are, for instance, that the memory phase disappears earlier for a fixed degree of depression and temperature. Moreover, in the oscillatory phase we observe that its width, in the corresponding phase diagram, increases with facilitation, as shown in Fig. 2. In addition, the frequency of the oscillations also increases with facilitation. As a consequence, we conclude that facilitation allows to recover information with less error but during a shorter period of time. This supports the idea that synaptic facilitation has influence in processes of short-term

memory. The facility to switch could be interesting to code both spatial and temporal information, and could explain, for instance, the spatio-temporal dynamics in the early olfactory processes which is described in [4].

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References


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