How to enhance the dynamic range of excitatory-inhibitory excitable networks

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We investigate the collective dynamics of excitatory-inhibitory excitable networks in response to external stimuli. How to enhance dynamic range, which represents the ability of networks to encode external stimuli, is crucial to many applications. We regard the system as a two-layer network (E-Layer and I-Layer) and explore the criticality and dynamic range on diverse networks. Interestingly, we find that phase transition occurs when the dominant eigenvalue of E-layer's weighted adjacency matrix is exactly one, which is only determined by the topology of E-Layer. Meanwhile, it is shown that dynamic range is maximized at critical state. Based on theoretical analysis, we propose an inhibitory factor for each excitatory node. We suggest that if nodes with high inhibitory factors are cut out from I-Layer, dynamic range could be further enhanced. However, because of the sparseness of networks and passive function of inhibitory nodes, the improvement is relatively small compared to original dynamic range. Even so, this provides a strategy to enhance dynamic range.

I. INTRODUCTION

In last decades, the theory of complex networks has enjoyed tremendous development in many fields as diverse as neural science, epidemic control, social activities, economics, etc. Especially, in the research of neural networks, many enlightening theoretical results, which are verified by experimental systems, are obtained. For example, Beggs et al. shows that neural avalanches, which have a power law size distribution, are important for cortical information processing and storage; Soriano et al. relates neural cultures with percolation on a graph, obtaining a percolation transition in connectivity characterized by a power law.

In applications, many biological, social and engineering problems are accurately described as networks of coupled excitable systems. The studies of how such networks respond to external stimuli reveal that, although single nodes usually respond to stimuli with small ranges, the collective response of the entire system can encode stimuli spanning several orders of magnitude. This property of broad dynamic range (the range of stimulus intensities resulting in distinguishable network response) is of particular significance for information processing in sensory neural networks.

In order to explain this phenomenon, a model of an excitable network based on Erdős–Rényi random graphs is proposed by Kinouchi and Copelli. It is shown that such networks have their sensitivity and dynamic range maximized at the critical point of a non-equilibrium phase transition. Later on, models on other diverse networks including those with scale-free, degree-correlated, and assortative topologies are discussed. More recently, a general theoretical approach to study the effects of network topology on dynamic range is presented by Larremore, Shew and Streppe. Interestingly, results show that the dynamic range is governed by the largest eigenvalue of the weighted network adjacency matrix.

All the models discussed above have only considered excitatory nodes. However, in realistic neural systems, the excitatory and inhibitory neurons are coexisting. Such excitatory-inhibitory (E-I) networks arise in many regions throughout the central nervous system and display complex patterns of activity. The behavior of E-I networks is critical for understanding how neural circuits produce cognitive function. Therefore, huge effort on E-I networks has been made. It is shown that excitation/inhibition balance is crucial for transmission of rate code in long feedforward networks, signal propagation in spiking neural networks, and discharge of cortical neurons. Although some other profound studies on E-I networks are presented, how such networks respond to external stimuli is largely unknown. Moreover, the investigation for the criticality and dynamic range of E-I networks with arbitrary topology still remains open.

In this paper, we investigate the criticality and dynamic range of E-I network models, providing a strategy to further enhance dynamic range. In section II, we propose the excitatory-inhibitory network model and give some basic definitions about criticality and dynamic range. Then in section III and IV, we conduct a theoretical study on ER random networks and scale-free networks respectively. It is proved that the criticality only relates to the topology of E-layer. In section V, we investigate methods to further enhance dynamic range of given networks by analyzing the mutual effects of E-layer and I-layer. We analyze the upper bound on improvement and explain why it is small compared to original dynamic range. Lastly, in section VI, we conclude the results and give a discussion about the further research.
II. THE EXCITATORY-INHIBITORY NETWORK MODEL

In the present model, each excitable element \(i = 1, \ldots, N\) has \(n\) states: \(s_i = 0\) is the resting state, \(s_i = 1\) corresponds to excitation and the remaining \(s_i = 2, \ldots, n - 1\) are refractory states. Here two types of nodes are considered: excitatory and inhibitory nodes. The function of excitatory nodes is to transmit excitation signals, increasing the probability of excitation of their neighbors, while the inhibitory ones decrease this probability. To be precise, at discrete times \(t = 0, 1, \ldots\) the states of the nodes \(s_i^t\) are updated as follows: (i) If node \(i\) is in the resting state, \(s_i^t = 0\), it can be inhibited by another excited inhibitory neighbor \(j\), \(s_j^t = 1\) with probability \(A_{ij}\). In this case, the state of node \(i\) will remain 0 in the next time step. Otherwise, it can be excited by its excited excitatory neighbor \(j'\) with probability \(A_{ij'}\), or independently by an external stimulus with probability \(\eta\). (ii) The dynamics of the nodes that are excited or in a refractory state is deterministic: if \(s_i = 1\), then in the next time step its state changes to \(s_i = 2\), and so on until the state \(s_i = n - 1\) leads to the state \(s_i = 0\) resting state, see Fig.1(a).

The network topology and strength of interactions between the nodes are described by the weighted adjacency matrix \(A = \{A_{ij}\}\). Notice that this matrix contains the information of both the excitatory and inhibitory links. In order to simplify the analysis, we only consider undirected networks here, so \(A\) is symmetrical. Besides, in this model, \(\eta\) will be assumed to be proportional to the stimulus level. Each element receives external signals independently.

Considering the two distinct types of nodes in this model, we can regard the system as a layered network [33]. The upper layer is composed of excitatory nodes, while the lower layer only contains inhibitory ones. We denote them as E-layer and I-layer respectively. An illustration of this two-layer model is shown in Fig.1(b). Assume the E-layer has \(N_e\) nodes, and I-layer has \(N_i\) nodes, then \(N_e + N_i = N\). Denote \(f_e\) and \(f_i\) as the fraction of excitatory and inhibitory nodes. We have \(f_e = N_e/N, f_i = N_i/N\). For the convenience of analysis, we rearrange the indices of nodes so that elements with index \(1 \leq i \leq N_e\) represent excitatory nodes and the others are inhibitory ones. Therefore, we obtain the following weighted adjacency matrix

\[
A_{N \times N} = \begin{pmatrix}
A_{EE} & A_{EI} \\
A_{EI} & A_{II}
\end{pmatrix}. \quad (1)
\]

Here \(A_{EE} = \{A_{ij}^{EE}\}_{N_e \times N_e}\) describes the topology and interaction strength of E-layer. \(A_{IE} = \{A_{ij}^{IE}\}_{N_e \times N_i}\) represents the effect of I-layer on E-layer. The other two matrices have similar meanings.

To analyze the dynamics of this system, we denote the probability that a given node \(i\) is excited \((s = 1)\) at time \(t\) by \(p_i^t\). In this model, we define the network instantaneous activity \(p_i^t\) at time \(t\) as the fraction of excited excitatory nodes, i.e. \(p_i^t = \frac{1}{N_e} \sum_{j=1}^{N_e} p_j^t\). Notice that we only care about the excitatory nodes here. We also define the average activity \(F = \frac{1}{T} \sum_{t=1}^{T} \sum_{i} p_i^t\), where \(T\) is a large time window. As a function of the stimulus intensity \(\eta\), networks have a minimum response \(F_0\) and a maximum response \(F_{max}\). We define the dynamic range \(\Delta = 10 \log(\eta_{\text{high}}/\eta_{\text{low}})\) as the range of stimuli that is distinguishable based on the system’s response \(F\), discarding stimuli that are too weak to be distinguished from \(F_0\) or too close to saturation. The range \([\eta_{\text{low}}, \eta_{\text{high}}]\) is found from its corresponding response interval \([F_{\text{low}}, F_{\text{high}}]\), where \(F_x = F_0 + x(F_{\text{max}} - F_0)\). The choice of interval is arbitrary and does not affect our results.

In the following text, we will investigate this model on networks with various topologies. As typical examples of homogeneous and heterogeneous networks, random networks and scale-free networks will be analyzed separately.

III. RANDOM NETWORKS

The network with \(N\) nodes is an Erdös – Rényi (ER) undirected random graph, with \(NK/2\) links being assigned to randomly chosen pairs of nodes. This produces an average degree \(K\). We randomly choose \(N_e = f_eN\) nodes as excitatory elements, and the rest as inhibitory ones. For simplicity, we set the strength of interactions between the nodes as \(S\) uniformly. In a mean-field approximation, the average branching ratio \(\beta\) of excitatory nodes \(\sigma = KS\) corresponds to the average number of excitations created in the next time step by an excited excitatory element. Of all these excited nodes, \(f_e\) nodes are excitatory and the others are inhibitory.

In the case of random networks, the update equations
for both E-layer and I-layer are same. Assume that the events of neighbors of a node being excited at time $t$ are statistically independent. It is shown that this approximation yields good results even the network has a non-negligible amount of short loops [9]. Therefore, we obtain the following mean-field map for $p^t$ at sufficient long time $t$:

$$p^{t+1} = (1 - (n - 1)p^t)(1 - Sp^t)^{f,K} \times \{\eta + (1 - \eta)[1 - (1 - Sp^t)^{f,K}]\}. \quad (2)$$

In the stationary state, $p^{t+1} = p^t = p$. Thus $F = \frac{1}{T} \sum_{t=1}^{T} p^t \approx p$ for large $T$. To check the critical behavior without an external field, we set $\eta = 0$ and linearize the term $(1 - Sp^t)^{f,K}$ and $(1 - Sp^t)^{f,K}$ in Eq $(2)$ around $p^t = 0$, obtaining

$$p \approx (1 - (n - 1)p)(1 - f_i \sigma p) f_i \sigma p. \quad (3)$$

Up to first order, we have the nonzero solution

$$p \approx \frac{\sigma - 1/f_e}{\sigma(f_i \sigma + n - 1)}. \quad (4)$$

Therefore, the critical point is $\sigma_c = \frac{1}{f_e}$, corresponding to the condition that the average number of excitations in E-layer caused by an excited node at each time step is exactly one. In particular, $\lim_{\eta \to 0} F = 0$ if $\sigma < \sigma_c$ and $\lim_{\eta \to 0} F > 0$ if $\sigma > \sigma_c$.

Next we will analyze the effect of a vanishing field at the critical point. In the limit of $p \to 0$, Eq $(2)$ can be approximated by

$$p = (1 - (n - 1)p)e^{-f_i \sigma p} \{\eta + (1 - \eta)(1 - e^{-f_i \sigma p})\}. \quad (5)$$

At the critical point $\sigma_c = 1/f_e$, we expand Eq $(5)$ to second order in the case of $\eta \to 0$. Then we have

$$p \approx \sqrt{\frac{\eta}{\sigma + n - 3/2}}. \quad (6)$$

Making use of Eq $(5)$, we can also predict the dynamic range. We can solve stimulus $\eta$ for any given response $F$. For a system with refractory time $n$, the maximal response $F_{\text{max}} = 1/n$. Once we find $F_0$, $F_{\text{low}}$ can be obtained. Then we can get $\eta_{\text{low}}$ with Eq $(6)$. For $\eta_{\text{high}}$, Eq $(6)$ is invalid because it is only valid for $p \to 0$. In order to approximate $\eta_{\text{high}}$, we set $\eta_{\text{high}}$ to one. Therefore, we have

$$\Delta = -10 \log[1 - e^{f_i \sigma F_{\text{low}}} + \frac{F_{\text{low}}}{1 - (n - 1)F_{\text{low}}}e^{\sigma F_{\text{low}}}] \quad (7)$$

Now we check the critical point of phase transition via simulations. Fig $(2)$ shows the relationship between response $F_{\eta \to 0}$ and branching ratio $\sigma$ for different excitatory proportions $f_e = 0.6$, 0.8 and 1. The critical point of each situation is in accordance with the theoretical value, satisfying $\sigma_c = 1/f_e$. Meanwhile, the behavior near the critical point is well captured by Eq $(4)$.

Then we check the effect of a vanishing field on the system. The response curves from $\sigma = 0$ to 2 (in intervals of 0.25) for $f_e = 0.8$ are presented in Fig $(3)$. The theoretical prediction of Eq $(6)$ fits the simulation data well. The phase transition occurs at $\sigma = 1.25$. In critical regime, the power law exponent $m = 1/2$, compatible with Eq $(6)$. Meanwhile, in subcritical regime, $m = 1$. The inset of Fig $(3)$ shows the critical state for different branching ratios. The theoretical lines from Eq $(6)$ agree with the simulations.

Fig $(4)$ shows the dynamic range for $f_e = 0.5$ and 2/3. As can be seen, dynamic range versus branching ratio is...
IV. SCALE-FREE NETWORKS

For networks with heterogeneous topology, we have to analyze each node separately. Assume the connectivity matrix is \( A = \{A_{ij}\}_{N \times N} \). Then the update equation for \( p_i^t \) is

\[
p_i^{t+1} = (1 - (n - 1)p_i^t) \prod_{j=N_e+1}^N (1 - A_{ij}p_j^t) \times \{\eta + (1 - \eta)[1 - \sum_{j=1}^{N_e} (1 - A_{ij}p_j^t)]\},
\]

To examine the critical point, we set \( \eta = 0 \) and expand Eq. (8) to first order in the limit of \( p_i^t \to 0 \). We have

\[
p_i^{t+1} = \sum_{j=1}^{N_e} A_{ij}p_j^t.
\]

Based on matrix \( A \), we create new matrices \( A^E \) and \( A^I \) as follows

\[
A^E = \begin{pmatrix} A^{EE} & 0 \\ A^{EI} & 0 \end{pmatrix},
\]

\[
A^I = \begin{pmatrix} 0 & A^{IE} \\ 0 & A^{II} \end{pmatrix}.
\]

Thus, Eq. (9) can be transformed into

\[
p^{t+1} = A^E p^t,
\]

where \( p^t = (p_1^t, \ldots, p_N^t)^T \) is the probability vector at time \( t \). We can see the stability of the solution \( p = 0 \) is governed by the dominant eigenvalue \( \lambda \) of matrix \( A^E \). Therefore, the phase transition happens at \( \lambda_c = 1 \). Notice that \( A^E_c \geq 0 \), so \( \lambda \) is real and positive according to Perron-Frobenius theorem. Recall Eq. (10), \( \lambda \) is in fact the dominant eigenvalue of matrix \( A^{EE} \). This means that the critical point only determined by the topology of E-layer.

Relating Eq. (12) with the power method in numerical analysis, we find that for small \( p \) and \( \eta \), \( p \) should be almost proportional to the normalized right eigenvector \( u \) of \( A^E \) corresponding to \( \lambda \). Thus we assume \( p_i = C u_i + \epsilon_i \) where \( C \) is a constant and \( \epsilon_i \) is an error term. Based on this, we obtain

\[
F = \frac{1}{N_e} \sum_{i=1}^{N_e} p_i \approx \frac{1}{N_e} \sum_{i=0}^{N_e} C u_i = C \langle u \rangle_c.
\]

Here \( \langle u \rangle_c = \sum_{i=1}^{N_e} u_i/N_e \). Near the critical state, we can approximate the product terms of Eq. (8) with exponential ones. Then in the steady state, we have

\[
p_i = (1 - (n - 1)p_i) \exp(-\sum_{j=N_e+1}^N A_{ij}p_j^t) \times \{\eta + (1 - \eta)[1 - \exp(-\sum_{j=1}^{N_e} A_{ij}p_j^t)]\}.
\]

To solve the stationary state, set \( \eta = 0 \). Then using \( p_i = C u_i + \epsilon_i \) and \( A^E u = \lambda u \), we expand Eq. (13) to second order for \( p_i \to 0 \).

\[
C u_i + \epsilon_i = (A^E \epsilon)_i + C \lambda u_i - C^2 \lambda u_i (A^I u)_i - ((n - 1)\lambda + \frac{1}{2} \lambda^2) C^2 u_i^2.
\]

In order to eliminate the error term \( \epsilon_i \), we may use \( u \) and \( v \) and simplify the calculation. Assume the right and left eigenvector of \( A^{EE} \) corresponding to the dominant eigenvalue \( \lambda \) is \( u^E \) and \( v^E \) respectively. We can express \( u \) and \( v \) using \( u^E \) and \( v^E \):

\[
u = (u^E, \frac{1}{\lambda} A^{EI} u^E)^T, v = (v^E, 0).
\]
The maximal dynamic range is observed at criticality \( \lambda_c = 1 \), see Fig. 6. Just as the case of ER random networks, in the subcritical regime, dynamic range increases monotonically with the dominant eigenvalue \( \lambda \) due to enhancement of link strength. On the contrary, in supercritical region, dynamic range decreases because of the increase of self-sustained activity \( F \).

Relating to the result of ER random networks, assume the mean degree of E-layer is \( \langle k \rangle_e \). Because the E-layer and I-layer are randomly connected, and the fraction of excitatory nodes is \( f_e \), we have \( \langle k \rangle_e = K f_e \).

In the case of ER random networks, the largest eigenvalue of \( A^{EE} \) can be approximated by \( \lambda \approx \langle d \rangle_e \), where \( \langle d \rangle_e = \frac{1}{N_e} \sum_{i,j=1}^{N_e} A^{EE}_{ij} \approx S(k) \sigma_c f_e \). Then the critical point \( \sigma_c \) satisfies \( \sigma_c f_e = 1 \), which is identical to the result of Eq. (14).

In the next section, we will explore the way to further enhance dynamic range of a given network.
the inhibitory factor $\delta_i$ for each excitatory node $i$ as follows:

$$
\delta_i = v_i u_i \sum_k \sum_j A_{ij}^E A_{jk}^{EI} u_k.
$$

(19)

We now explain the meaning of the inhibitory factor $\delta_i$. In the case of weak stimulus, the influence of inhibitory nodes is relatively passive. According to the rule, only the inhibitory nodes are activated can they release inhibitory signals. Thus in this condition, excitatory nodes actually have two effects: activating their excitatory neighbors to transmit excitations and inhibitory neighbors to release inhibitory signals. The latter effect is implemented via the inhibitory paths, which start from excitatory nodes and end in E-layer through one step in I-layer, see Fig.1(b). For the inhibitory paths which start at node $i$ and ends at node $k$, the interaction strength is $\sum_j A_{ij}^E A_{jk}^{EI}$. Because in stationary state, excitatory nodes have different active probabilities, each inhibitory path should have its own weight. Here we set the weight as $u_i u_k$, reflecting the active probability of node $i$ and $k$. After summarizing the weighted interaction strength of inhibitory paths starting from node $i$, we have $u_i \sum_k \sum_j A_{ij}^E A_{jk}^{EI} u_k$. Lastly, we multiply it with $v_i$ to reflect the active probability of node $i$, obtaining the definition of inhibitory factor $\delta_i$. Therefore, $\delta_i$ can be interpreted as the ability of node $i$ to cause inhibition in stationary state. In the procedure of enhancing dynamic range, we only need to cut out the nodes with large $\delta_i$ from I-layer.

Now we check the relationship between the inhibitory factor and degree. In Fig.7 we present the simulation results of both ER random networks and BA scale-free networks. The solid dots represent the mean value and the vertical lines show the standard deviation. For ER random networks, nodes with small degree always have small $\delta_i$. However, for nodes with large degree, there are huge fluctuations in $\delta_i$. We quantify these fluctuations with standard deviations. As can be seen, the standard deviations are comparable with mean values, making the ranges of inhibitory factor for different degrees overlap with each other. More simulations with different network sizes show that these huge relative fluctuations also exist for other system sizes. As for scale-free networks, the values of $\delta_i$ are divided into two groups: for most nodes with small degree, their inhibitory factors are negligible; whereas, highly-connected nodes have larger $\delta_i$, which also shows great fluctuations. Also, these fluctuations are unrelated to system size. It can be seen there is no clear relations between inhibitory factor and degree. Consequently, it is unreasonable to just cut out the nodes with high degree from the I-layer.

We give an analysis of improvement of dynamic range. By Eq.(18) and definition of dynamic range, we have the upper bound on the improvement in dynamic range

$$
\Delta_u = 10 \log\left(1 + \frac{u(A_i u_{ji})_e}{(n - 1/2)u_{ji}^2 e}\right).
$$

(20)

For ER random networks, we can simplify the calculation as follows. For E-I networks with mean degree $K$, on average, each excitatory node has $f_i K$ edges pointing to I-layer, and each inhibitory node has $f_e K$ edges pointing back to E-layer. So we approximate $\sum_j A_{ij}^E A_{jk}^{EI} \approx f_i K S_f \cdot f_e K S_e / N_e$. Substituting this term in Eq.(19), we have

$$
\delta_i \approx v_i u_i f_e f_i K^2 S^2 (u)_e.
$$

(21)

Then we use Eq.(20) to obtain

$$
\Delta_u = 10 \log\left(1 + \frac{f_e f_i K^2 S^2 (u)_e}{N_e (n - 1/2)u_{ji}^2 e}\right).
$$

(22)

As for scale-free networks, we can check the upper bound through numerical calculations.

Fig.8 shows the simulations of upper bound $\Delta_u$. For both cases, $\Delta_u$ decreases with the increasing fraction of excitatory nodes $f_e$. The only difference is that scale-free networks present larger fluctuation, which cannot affect the decreasing trend. In the insets, we explore the effect of system size on upper bound. It can be seen for both cases, $\Delta_u$ is not influenced by system size. Also, the scale-free networks show larger fluctuation. Another conclusion we get from Fig.8 is that the improvement is small compared to the original dynamic range. This can be explained as follows. Firstly, $A_i^E$ and $A_i^{EI}$ are sparse, so there are not much inhibitory paths exist. This makes the term $\sum_j A_{ij}^E A_{jk}^{EI}$ very small. Therefore, by Eq.(19) and (20), $1 + \frac{u(A_i u_{ji})_e}{(n - 1/2)u_{ji}^2 e}$ is close to 1, which makes $\Delta_u$ small. Secondly, in the critical state, each excitatory node can only excite a small number of inhibitory nodes. Because the function of inhibitory nodes is passive (i.e. only they are excited can they exert inhibitory impact), this weakens the influence of inhibitory nodes. Consequently, cutting out the inhibitory nodes can only enhance dynamic range.
range a little. In fact, by this method, we cannot change the functional form $\eta \propto F^2$. We can only change the constant parameter.

To check the effect of our method, we perform a comparison of random remove and targeted remove. Denote $p_r$ as the fraction of nodes which are cut out from I-layer. In random remove, we choose the nodes randomly, while in targeted remove we select the nodes with top $p_r$ percent inhibitory factors. In Fig. 8, we see for both cases, dynamic range of targeted remove increases faster than that of random remove. However, the improvement is relatively small compared with the original dynamic range.

VI. CONCLUSIONS AND DISCUSSIONS

In this paper, we propose an excitatory-inhibitory excitable network model. To analysis the criticality and dynamic range of this model, we divide this network into two layers: E-layer which consist of excitatory nodes and I-Layer which only contains inhibitory ones. Based on it, we give a theoretical analysis on random networks and scale-free networks respectively. It is proved that, the critical state occurs at $\sigma_c = 1/f_e$ for random networks and the dynamic range is maximized at criticality. As for scale-free networks, the phase transition happens when the largest eigenvalue of the E-layer’s weighted adjacency matrix $A^{EE}$ is just one. Similarly, the dynamic range is also optimized at critical point. It is interesting that the critical point is not affected by the I-layer and the links between these two layers. Then, we discuss the method to enhance dynamic range. Based on the analysis of mutual effects, we propose an inhibitory factor $\delta_i$ for each excitatory node to quantify their inhibitory ability. By cutting out the excitatory nodes with high inhibitory ability from I-Layer, the dynamic range can be further improved. However, the improvement is relatively small. We give an analysis to the upper bound on improvement in dynamic range and explain why the enhancement is small.

For further study, it is meaningful to investigate networks with heterogenous connectivity strength [35]. Meanwhile, the study of real-world neural or sensor networks is also of great significance for the applications of the theory.

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