Functional Modularity of Background Activities in Normal and Epileptic Brain Networks

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We analyze the connectivity structure of weighted brain networks extracted from spontaneous magnetoencephalographic signals of healthy subjects and epileptic patients (suffering from absence seizures) recorded at rest. We find that, for the activities in the 5–14 Hz range, healthy brains exhibit a sparse connectivity, whereas the brain networks of patients display a rich connectivity with a clear modular structure. Our results suggest that modularity plays a key role in the functional organization of brain areas during normal and pathological neural activities at rest.

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From the brain to the Internet and to social groups, the characterization of the connectivity patterns of complex systems has revealed a wiring organization that can be captured neither by regular lattices, nor by random graphs [1]. In neurosciences, it is widely acknowledged that the emergence of several pathological states is accompanied by changes in brain connectivity patterns [2]. Recently, it has been found that functional connectivity patterns obtained from magnetoencephalography (MEG) and electroencephalography (EEG) signals during different pathological and cognitive brain states (including epilepsy) display small-world (SW) properties [3]. Empirical studies have also lead to the hypothesis that brain functions rely on the coordination of a scattered mosaic of functionally specialized brain regions (modules), forming a weblike structure of neural assemblies [2]. Modularity is a key concept in complex networks from RNA structures to social networks [4,5]. A module is usually defined as a subset of units within a network, such that connections between them are denser than connections with the rest of the network. In biological systems, it is generally acknowledged that modularity results from evolutionary constraints and plays a key role in robustness, flexibility, and stability [6].

Absence seizures are the most characteristic expression of nonconvulsive generalized epilepsy. Their main characteristic in brain signals is the occurrence of high amplitude, rhythmic spikewave discharges synchronized over wide cortical areas, which manifests suddenly from a normal background. Current studies of brain connectivity mainly focus on the onset and evolution of epileptic discharges. Nevertheless, little is known about the organization of brain networks during spontaneous background activities of patients, and the role of these connectivity patterns on the emergence of absence seizures. In this Letter we study the modular organization of brain networks extracted from spontaneous MEG signals of epileptic patients and healthy subjects. The results of our analysis reveal a nonrandom structural organization in both normal and pathological brain networks. In particular, the functional networks of control subjects are characterized by a sparse connectivity between the modules. In contrast, brain connectivity of epileptic patients (recorded out of seizures) displays a configuration where nodes in a functional module are connected to different functional modules. This modular configuration might play a key role in the integration of large scale brain activities, facilitating the emergence of epileptic discharges.

The data used in this study were acquired from 5 healthy subjects and 5 epileptic patients suffering from absence seizures. The study was performed with written consent of the subjects and with the approval of the local ethics committee. During the recordings, subjects and patients were instructed to rest quietly, but alert, and keep their eyes closed. The brain signals were acquired with a wholehead MEG system (151 sensors; VSM MedTech, Coquitlam, BC, Canada), digitized at 1.25 kHz with a bandpass of 0–200 Hz. All the analyses were performed on 338 nonoverlapping quasistationary segments (206 for all the patients and 132 for the healthy group) of 5 s without eyes or muscular artifacts, nor epileptic activities (as, e.g., seizures or epileptiform activity) and far (at least 10 s) from recent epileptic discharge. In agreement with previous findings, surrogate data tests revealed that less than 4% of interdependencies between the spontaneous brain activities were nonlinear [7]. Thus, weighted brain networks were constructed by means of a definition of functional links based on linear coherence. The squared modulus of the coherence between two time series \( x_i(t) \) and \( x_j(t) \) (normalized to zero mean and unit variance) was defined as \( |\Gamma_{ij}(f)|^2 = \frac{|S_{ij}(f)|^2}{S_{ii}(f)S_{jj}(f)} \), where \( S_{ii} \) and \( S_{ij} \) (the spectral and cross-spectral densities) were estimated using the Welch’s averaged periodogram method [8].

Recent results show that correlations between magnetic fields sensors located at a distance less than 4 cm cannot distinguish between spontaneous activities of epileptic
patients and control subjects [9]. To reduce the influence of these spurious correlations between MEG signals, we have excluded the nearest sensors (separated less than 5 cm) from the computation of coherence values. To perform the statistical analysis of coherence values, we used Fisher’s Z transform of $\Gamma_{ij}$: $Z_{ij} = 0.5 \ln \left( \frac{1 + \Gamma_{ij}}{1 - \Gamma_{ij}} \right)$. Under the hypothesis of independence, $Z_{ij}$ has a normal distribution with expected value 0 and variance $1/2N_{ij}$, where $N_{ij}$ is the number of nonoverlapping blocks used in the estimation of spectral densities [8]. To correct for multiple testing, the false discovery rate (FDR) method was applied to each matrix of $\Gamma_{ij}$ values [10]. With this approach, the threshold of significance $\Gamma_{th}$ was set such that the expected fraction of false positives is restricted to $q \approx 0.01$. Finally, in the network construction, a functional connection between two of the $N = 151$ nodes (brain sites) was assumed as an undirected weighted link; i.e., we set the weight of the link between $i$ and $j$ as $w_{ij} = w_{ji} = \Gamma_{ij}$ when $\Gamma_{ij} > \Gamma_{th}$, and $w_{ij} = 0$ otherwise.

To characterize the network structure of healthy subjects and epileptic patients, we evaluated a list of measures for weighted graphs [1]. Namely, the node strength $s_i = \sum_k w_{ik}$, the weighted clustering coefficient $c_i$ [11], the efficiency $E_i$ of a node $i$ [11], as well as the respective averages over the graph nodes $S = 1/N \sum_i s_i$, $C_w = 1/N \sum_i c_i$ and the global efficiency of the graph $E$. As null model, brain networks were compared to equivalent random graphs, obtained by randomly rewiring the links of the original networks while keeping the same degree distribution [12]. The statistical significance of a given metric $\theta$ is assessed by quantifying its statistical deviation from values obtained in the ensemble of randomized networks. Let $\mu$ and $\sigma$ be the mean and SD of the parameter $\theta$ computed from such an ensemble. The significance is given by the ratio $\Sigma_\theta = |\theta - \mu|/\sigma$ whose $p$ value is given by the Chebyshev’s inequality [13]. To assess significant differences between topological features of the two groups, we used standard nonparametric permutation methods, which account for spatial dependences in the data [14]. We used exhaustive permutations ($10^5$) to estimate the empirical distribution under the null hypothesis of no difference between the two groups.

Although we applied our approach to connectivity graphs obtained from brain oscillations at the conventional frequency bands ($f < 5$ Hz, $5 < f < 15$ Hz, $15 < f < 24$ Hz, $24 < f < 35$, and $f > 35$ Hz), statistically significant differences between normal and epileptic brain networks were observed only for the brain activities in the extended alpha range ($5–14$ Hz). Henceforth, all results presented here refer to functional networks obtained at this frequency band. The basic network properties (averaged over subjects) are summarized in Table I. The structure of functional brain networks was found to be significantly different from that of randomized counterparts. Namely, brain networks of both patients and healthy subjects yielded a clustering coefficient $C_w$ larger than that of randomized graphs ($p < 10^{-3}$), and values of efficiency as large as those of random graphs, indicating a small-world behavior. These results agree with previous findings suggesting that brain sites have an optimal interaction with most other brain regions [3]. Furthermore, average network properties ($S$, $C_w$, and $E$) of epileptic patients were found to be different ($p < 10^{-3}$) from those of control subjects, supporting the hypothesis that neural disturbances are correlated with changes in functional network architectural features [2].

More detailed information on the differences between patients and control subjects can be acquired by the analysis of the network at the level of node properties. In Fig. 1 we report the spatial distribution of node measures $s_i$, $c_i$, and $E_i$ (averaged over all control subjects and over all patients) for each sensor of the network. Results indicate that epileptic patients have a richer node connectivity than control subjects. Difference maps clearly identify the centroparietal regions as those brain areas with the highest contrast between patients and control group. Although the node strength of epileptic patients is twice as large as that of healthy subjects, if links of real networks are randomly rewired by keeping the same degree distribution, no sig-

| TABLE I. | Network properties in control subjects (CTL) and patients (PAT); mean node strength $S$, average weighted clustering coefficient $C_w$, global efficiency $E$, maximal modularity $Q$, and number of modules $N_m$. $\mu_\theta$ denotes the average of metric $\theta$ obtained from 20 random graphs. |
|-----------|-----------------|-----------------|-----------------|-----------------|
| $S$       | $C_w(\mu_\theta)$ | $E(\mu_\theta)$ | $Q(\mu_\theta)$ | $N_m$          |
| CTL       | 4.04            | 0.233 (0.096)   | 0.649 (0.892)   | 0.538 (0.001)   | 13.7           |
| PAT       | 7.34            | 0.300 (0.160)   | 0.893 (1.101)   | 0.503 (0.006)   | 8.30           |

FIG. 1 (color). Topographic distribution of node strength $s_i$ (a), weighted clustering coefficient $c_i$ (b), and efficiency $E_i$ (c). Control subjects (CTL), patients (PAT), and difference maps (p values) of their comparison are reported for real and equivalent random configurations.
significant differences are observed between the two groups. This rules out the possibility that the difference in the number of connections alone could account for differences in \( C_w \) and \( E \). No significant difference between healthy subjects and patients was observed by direct comparison of the spatial maps of the \( \alpha \)-activity power [15].

A potential modularity of brain networks is suggested by the fact that the networks display a clustering coefficient larger than that obtained in random graphs [16]. Previous studies over brain networks have used clustering methods to identify similar groups of brain activities. However, classical approaches such as those based on principal components analysis (PCA) and independent components analysis (ICA), make very strong statistical assumptions (orthogonality and statistical independence of the retrieved components, respectively) with no physiological justification [17]. To find the network modules we have instead used an algorithm based on a spectral embedding of graphs [18] (similar theoretical frameworks have been recently proposed [19]). The algorithm is based on the definition of a Markov chain with a transition probability matrix \( P \) with entries \( p_{ij} = \frac{w_{ij}}{\sum_j w_{ij}} \). If \( P^t \) is the \( t \)th iterate of matrix \( P \), the element \( p_{ij} (t) \) encodes the probability of moving from node \( i \) to node \( j \) through a random walk of length \( t \). For an undirected and connected graph, the Perron Frobenius theorem assures that \( \mu_1 = \frac{1}{\sum_i s_i} \) is the unique stationary distribution of the Markov chain, such that \( \lim_{t \rightarrow \infty} \sum_i p_{ij} (t) \mu_i (0) = \mu_j \). For directed networks, recent approaches are proposed to ensure this convergence [20].

The random walk gives rise to a geometric diffusion with an associated distance between nodes \( i \) and \( j \) defined as [18]:
\[
d_{ij}^2 (t) = \sum_{k=0}^{t} \frac{p_{i,k}(t) - p_{i,k}(t)}{\mu_i(t)},
\]
where the term \( \mu_i^* \) is supposed to compensate for discrepancies in local densities. By construction, distance between nodes is strongly ruled by the connectivity of the graph, and it takes small values if nodes are connected by many paths. Considering the spectral representations of matrix \( P \), one has a set of eigenvalues \( |\lambda_0| \geq |\lambda_1| \geq \ldots \geq |\lambda_{N-1}| \) and eigenvectors \( \phi_0 \) and \( \phi_k \) such that \( \phi_k^T P = \lambda_k \phi_k^T \) and \( P \phi_k = \lambda_k \phi_k \). The diffusion distance can be written as
\[
d_{ij}^2 (t) = \sum_{k=0}^{\beta} \lambda_k^2 [\phi_k (i) - \phi_k (j)]^2,
\]
where \( \phi_k (j) \) denotes the component \( j \) of eigenvector \( k \). The diffusion distance can be approximated (note that \( \phi_0 = \mu^* \) and \( \psi_0 \) is a constant vector) to a relative precision using the first \( \beta \) nontrivial eigenvectors and eigenvalues:
\[
d_{ij}^2 (t) = \sum_{k=1}^{\beta} \lambda_k^2 [\psi_k (i) - \psi_k (j)]^2.
\]
This is equivalent to embedding the graph in a low dimensional space \( R^\beta \), converting the diffusion distance between nodes of the graph into Euclidean distance in \( R^\beta \) [18]. This approach has the main advantage that it defines a meaningful representation of the graph and it leads to explicitly define a distance metric on the space \( R^\beta \) that reflects the connectivity of the network. Graph modules are then extracted by a \( k \)-means clustering algorithm in the embedding space. The algorithm starts with a random assignment of \( N_m \) cluster center. Then, the partition is updated by repeating the following steps: (i) each point is assigned to the nearest center; (ii) the new geometric center of each cluster are recomputed. The algorithm stops when the assignments of nodes are no longer changed. The algorithm reaches convergence in very few iterations after the initial generations, but it only guarantees a convergence towards a local optimum. In fact, the \( k \) means is sensitive to the initial choice of centroids. To overcome this drawback, we run the algorithm several times (500) for different number of modules \( N_m = 2, \ldots, 15 \) and return the partition yielding the largest modularity \( Q \). The modularity \( Q \) for a given partition \( S \) of a weighted network is defined as [4]
\[
Q(S) = \sum_{i=1}^{N} \left[ \frac{l_i}{2L} - \left( \frac{d_i}{2\hat{d}_i} \right)^2 \right],
\]
where \( L \) is the total weight of all connections in the network, \( l_i \) is the weight of links between vertices in module \( s \), and \( d_i \) is the total weight of links in module \( s \). To select the optimal number of diffusion coordinates, we have repeated the clustering algorithm for all possible values of \( \beta \), and looked for the largest value of \( Q \).

The method has been tested on synthetic networks [21], and used to find modules in brain networks. The presence of modules is actually confirmed by the high values of \( Q \) obtained for the optimal partition of the brain networks of both control subjects and patients. The main result is that brain networks of epileptic patients have a structure consisting in a smaller number of modules \( N_m = 8.30 \) with respect to that of control subjects \( N_m = 13.7 \). To assess the stability of the partition structure across healthy subjects and patients we used the adjusted Rand index \( J \) [22], which yields a normalized value between 0 (if the two partitions are randomly drawn) and 1 (for an identical partition structure). The values of \( J \) indicate a high stability of the partition structure across all the epileptic patients \( J = 0.733 \) and certain variability in the modular structure of control subjects \( J = 0.479 \).

Figure 2(a) illustrates the spatial distribution of the modules obtained for healthy subjects and patients. Despite the differences, some modules fit well some known brain regions including occipital, parietal, and frontoral areas. Although a one-to-one assignment of anatomical brain areas to the retrieved modules is difficult to define, modules assignment provides the basis for the analysis of nodes according to their patterns of intra- and intermodules connections [5]. The within-module degree \( z \) score measures how well connected the node \( i \) is to other nodes in the same module, and is defined as \( z_i = \frac{k_{ij} - \bar{k}_i}{\sigma_k} \), where \( k_{ij} \) is total weight of links of node \( i \) to other nodes in its module \( j \), while \( \bar{k}_i \) and \( \sigma_k \) are average node strength and standard deviation of nodes in \( s \). Thus node \( i \) will display a large value of \( z_i \) if it has a large number of intramodular connections relative to other nodes in the same module. The participation coefficient \( p_{ij} = 1 - \frac{\sum_{k=1}^{\beta} (\phi_k)^2}{\sum_{k=1}^{\beta} (\phi_k)^2} \) quantifies instead to which extent a node \( i \) is connected to different modules. This coefficient takes values of zero if a node has most of its links exclusively with other nodes of
A modular description of brain networks might provide, more in general, meaningful insights into the functional organization of brain activities recorded with others neuro-imaging techniques (EEG, MEG, or functional magnetic resonance imaging) during diverse cognitive or pathological states. Applied to other multivariate data (e.g., financial or epidemiological time series), our approach could provide new insights into the network structure of spatially extended systems.

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[21] The performance of the method has been tested on computer generated graphs with tunable modularity structure. Even when half of a node’s links points to outside modules, the accuracy of the method is high (∼80%) and comparable to that of the algorithm in Ref. [4].
[23] For all the topological features, a nominal type I error control by permutation testing revealed no significant differences between two randomly subsets of each group data.