



NETWORK NEURO SCIENCE

an open access  journal



Citation: Nunes, R. V., Reyes, M. B., Mejias, J. F., & de Camargo, R. Y. (2021). Directed functional and structural connectivity in a large-scale model for the mouse cortex. *Network Neuroscience*, 5(4), 874–889. https://doi.org/10.1162/netn_a_00206

DOI:
https://doi.org/10.1162/netn_a_00206

Supporting Information:
https://doi.org/10.1162/netn_a_00206

Received: 12 March 2021
Accepted: 9 August 2021

Competing Interests: The authors have declared that no competing interests exist.

Corresponding Author:
Ronaldo V. Nunes
ronaldovnunes@gmail.com

Handling Editor:
Olaf Sporns

Copyright: © 2021
Massachusetts Institute of Technology
Published under a Creative Commons
Attribution 4.0 International
(CC BY 4.0) license



RESEARCH

Directed functional and structural connectivity in a large-scale model for the mouse cortex

Ronaldo V. Nunes¹, Marcelo B. Reyes¹, Jorge F. Mejias², and Raphael Y. de Camargo¹

¹Center for Mathematics, Computing, and Cognition, Universidade Federal do ABC, São Bernardo do Campo, Brazil

²Swammerdam Institute for Life Sciences, University of Amsterdam, Amsterdam, The Netherlands

Keywords: Large-scale brain models, Directed functional connectivity, Causality, Generalized partial directed coherence

ABSTRACT

Inferring the structural connectivity from electrophysiological measurements is a fundamental challenge in systems neuroscience. Directed functional connectivity measures, such as the generalized partial directed coherence (GPDC), provide estimates of the causal influence between areas. However, the relation between causality estimates and structural connectivity is still not clear. We analyzed this problem by evaluating the effectiveness of GPDC to estimate the connectivity of a ground-truth, data-constrained computational model of a large-scale network model of the mouse cortex. The model contains 19 cortical areas composed of spiking neurons, with areas connected by long-range projections with weights obtained from a tract-tracing cortical connectome. We show that GPDC values provide a reasonable estimate of structural connectivity, with an average Pearson correlation over simulations of 0.74. Moreover, even in a typical electrophysiological recording scenario containing five areas, the mean correlation was above 0.6. These results suggest that it may be possible to empirically estimate structural connectivity from functional connectivity even when detailed whole-brain recordings are not achievable.

AUTHOR SUMMARY

We analyzed the relationship between structural and directed functional connectivity by evaluating the effectiveness of generalized partial directed coherence (GPDC) to estimate the connectivity of a ground-truth, data-constrained computational model of a large-scale network model of the mouse cortex. We show that GPDC values provide a reasonable estimate of structural connectivity even in a typical electrophysiological recording scenario containing few areas. These results suggest that it may be possible to empirically estimate structural connectivity from functional connectivity even when detailed whole-brain recordings are not achievable.

INTRODUCTION

The communication between brain regions is often analyzed using structural and functional connectivity (Avena-Koenigsberger, Misic, & Sporns, 2018). The former refers to anatomical connections between brain regions generally quantified using tracer injections or diffusion magnetic resonance imaging (Ambrosen et al., 2020). The map of these connections is called “connectome” (Sporns, Tononi, & Kötter, 2005). Network measures are usually used to analyze the connectome, whereas nodes represent brain regions and edges refer to axonal projections

(Bassett & Sporns, 2017; Bassett, Zurn, & Gold, 2018). Functional connectivity estimates brain communication from statistical relations between recorded brain signals (Avena-Koenigsberger et al., 2018; Reid et al., 2019). Particularly, directed functional connectivity methods use the concept of causality to infer both the intensity and the direction of the connections between brain regions (Bastos & Schoffelen, 2016). Even though there is some association between structural and functional connectivity, the relationship between them is not straightforward (Avena-Koenigsberger et al., 2018). While the former is practically static and composes the map of possible pathways for information flow between brain regions, the latter changes continuously and depends, for example, on the dynamical states of brain regions, noise, and strength of structural connections (Nunes, Reyes, & De Camargo, 2019).

During electrophysiological procedures, researchers typically record brain signals using electrodes positioned in different depths of brain regions. Even with the improvement in technologies for recording signals, it is usually possible to record signals only from a few areas compared with the number of sources of activity in the brain (Harris, Quiroga, Freeman, & Smith, 2016; Hong & Lieber, 2019; Schölvinck, Leopold, Brookes, & Khader, 2013). Thus, the functional connectivity analysis presents a problem because many unrecorded regions may indirectly influence other regions as common inputs (Bastos & Schoffelen, 2016; Reid et al., 2019; Sanchez-Romero & Cole, 2019). Therefore, the comparison between structural and functional connectivity becomes more complicated since spurious inferred causality relations can lead to misinterpretations of electrophysiological data.

Previous simulation studies evaluated the relation between directed functional connectivity and structural connections (Baccalá & Sameshima, 2001; Barnett & Seth, 2014; Mi, Cheng, & Zhang, 2014; Nunes et al., 2019). However, most of these studies used either autoregressive (Novelli, Wollstadt, Mediano, Wibral, & Lizier, 2019) or rate-based models (Mi et al., 2014) for the dynamics of each cortical area. These studies provided essential steps towards evaluating the reliability of causality measures. However, the time series obtained from autoregressive and rate models are distant from electrophysiological signals obtained in experimental laboratory conditions. Using **spiking models**, we can capture the dynamic of neuronal networks while generating simulated **local field potential (LFP)** signals from the synaptic currents. Also, most studies do not consider the impact of accessing only part of the activity in the brain.

In this work, we investigate the relationship between directed functional connectivity and structural connectivity in a large-scale network model of the cortex, derived from a cortical connectome of the mouse obtained using tracer injections (Gămănuț et al., 2018). We used generalized partial directed coherence (GPDC), a frequency-domain method based on **multivariate vector autoregressive (MVAR)** models, which provides estimates of directed functional connectivity (Baccalá, Sameshima, & Takahashi, 2007; Sameshima & Baccalá, 2014). The mean correlation between the fraction of labeled neurons (FLN) and GPDC remained high ($r > 0.6$) even when only a few cortical areas were considered in the GPDC calculation, indicating that this causality measure provides reliable results in typical experimental conditions in which only recordings from a subset of areas are available.

METHODS

Neuron Model

We modeled the neurons using a single-compartment Hodgkin–Huxley-type model, where the membrane potential of the i -th neuron described by,

$$C_m \frac{dV_i}{dt} = -g_{Na} m_i^3 h_i (V - E_{Na}) - g_K n_i^4 (V - E_K) - g_L (V - E_L) + I_{ext,i} + I_{syn,i}, \quad (1)$$

Spiking models:

A network model composed of spiking neurons (nodes) connected by synapses (edges). A spiking neuron is a simplified neuron model, which generates discrete spike events. It is also referred to as a spiking neuronal population model.

Local field potential (LFP):

A transient electrical activity in the extracellular medium resulting from ionic flows in multiple neurons.

Multivariate vector autoregressive (MVAR) model:

Autoregressive models permit predicting future values in time series from past values. Multivariate vector models extend them to work with multiple interdependent time series.

the membrane capacitance C_m is 0.50 nF (0.25 nF) for excitatory (inhibitory) neurons. The maximal conductances values were $g_{Na} = 12.5 \mu S$, $g_K = 4.74 \mu S$, and $g_L = 0.025 \mu S$. The reversal potentials $E_{Na} = 40$ mV, $E_K = -80$ mV, and $E_L = -65$ mV correspond to the sodium, potassium, and leakage channel, respectively (Gutfreund, Yarom, & Segev, 1995). The dynamics of the voltage-gated ion channels are described by activation and inactivation variables m , n , and h , where m and n account for the dynamics of Na channels and h for K channels. The probability that an ion channel is open evolves according to a set of ordinary differential equations (San Cristóbal, Vicente, & Garcia-Ojalvo, 2014),

$$\frac{dm}{dt} = \alpha_m(V)(1 - m) - \beta_m(V)m, \quad (2)$$

$$\frac{dh}{dt} = \alpha_h(V)(1 - h) - \beta_h(V)h, \quad (3)$$

$$\frac{dn}{dt} = \alpha_n(V)(1 - n) - \beta_n(V)n, \quad (4)$$

where

$$\alpha_m(V) = 0.1 \frac{(V + 16)}{1 - \exp(-(V + 16)/10)}, \quad (5)$$

$$\beta_m(V) = 4 \exp(-(V + 41)/18), \quad (6)$$

$$\alpha_h(V) = 0.07 \exp(-(V + 30)/20), \quad (7)$$

$$\beta_h(V) = [1 + \exp(-V/10)]^{-1}, \quad (8)$$

$$\alpha_n(V) = 0.01 \frac{(V + 20)}{1 - \exp(-(V + 20)/10)}, \quad (9)$$

$$\beta_h(V) = 0.125 \exp(-(V + 30)/80). \quad (10)$$

The parameters used in this neuron model were previously reported and applied in some studies that modeled cortical neuronal populations (Barardi, San Cristóbal, & Garcia-Ojalvo, 2014; De San Cristóbal, Vicente, Sancho, & Garcia-Ojalvo, 2013; San Cristóbal et al., 2014).

Spiking Neuronal Population Model

Each spiking neuronal population was composed of 2,000 neurons, 1,600 excitatory and 400 inhibitory. Connections between neurons within each spiking neuronal population are

Table 1. Synaptic weights for intra-areal connections. Mean synaptic weight w and standard deviation σ_w for all possible synapses. E, I, and input represent excitatory neurons, inhibitory neurons, and external input, respectively. The arrow indicates the direction of the connection.

Synapses	\bar{w} (nS)	σ_w (nS)
$E \rightarrow E$	2.5	1.0
$E \rightarrow I$	2.5	1.0
$I \rightarrow E$	240	10
$I \rightarrow I$	240	10
$Input \rightarrow E$	3.2	1.0
$Input \rightarrow I$	3.2	1.0

random with connection probability $p_{\text{intra}} = 10\%$. The synaptic current I_{syn} that arrives to post-synaptic neuron i is modeled by

$$I_{\text{syn},i}(t) = \sum_{j \in \text{presyn}} g_{\text{syn},i,j}(t) [E_{\text{syn}} - V_i(t)], \quad (11)$$

where the index j represents a presynaptic neuron connected to neuron i , and the sum over j accounts for all the synapses that impinge on neuron i . E_{syn} is the synaptic reversal potential, which is 0 mV for excitatory and -70 mV for inhibitory synapses. The dynamics of synaptic conductance $g_{\text{syn},i,j}$ is described by an exponential function as follows (Tomov, Pena, Zaks, & Roque, 2014):

$$g_{\text{syn},i,j}(t) = we^{-(t-t_j-d)/\tau}. \quad (12)$$

The characteristic decay time τ is 2 ms and 8 ms for excitatory and inhibitory synapses, respectively. When a presynaptic neuron j fires a spike at time t_j , $g_{\text{syn},i,j}$ is incremented by a synaptic weight w after the axonal delay d , which was set as 1 ms for all intra-areal connections (San Cristóbal et al., 2014). The value of w depends on the excitatory/inhibitory nature of the presynaptic and postsynaptic neurons. Furthermore, all neurons receive a background input given by a heterogeneous Poisson-process spiking activity with a rate of 7.3 kHz (San Cristóbal et al., 2014). The background input acts as an excitatory synaptic current. To add heterogeneity in our model, all synaptic weights w for recurrent connections and background input were taken from a Gaussian distribution (Table 1).

Mouse Large-Scale Cortical Network

The mouse cortex's large-scale network model is composed of 19 areas where a spiking neuronal population models each area with long-range and recurrent synapses. Parameters related to recurrent synapses were described in the previous session. Neurons from different areas are randomly connected with probability $p_{\text{inter}} = 5\%$. The synaptic weights between cortical areas are based on the previously published anatomical connectivity dataset for the mouse cortex (Gămănuț et al., 2018) obtained by retrograde tracer injections (Markov et al., 2014).

This technique consists of injecting a tracer that flows from the target synapses to the cell bodies, allowing us to identify neurons projecting to the target area. The fraction of labeled neurons (FLN) was measured as the ratio of the number of labeled neurons in a source area to the total quantity of labeled neurons in all source areas, where labeled neurons considered are

extrinsic to the injected area (Markov et al., 2014; Theodoni et al., 2020; Vezoli et al., 2020). We defined the edge measure FLN_{ij} as the number of neurons projecting from area j to area i , divided by the number of neurons projecting to area i from all the areas except i (de Lange, Ardesch, & van den Heuvel, 2019; Joglekar, Mejias, Yang, & Wang, 2018). The synaptic weights for directed long-range connections are the FLNs scaled by the global scaling parameters $\mu_E = 50$ and $\mu_I = 25$,

$$w_{lr,E}^i = \mu_E \sum_{j=1}^N FLN_{ij}, \quad (13)$$

$$w_{lr,I}^i = \mu_I \sum_{j=1}^N FLN_{ij}. \quad (14)$$

Long-range connections are excitatory, targeting either excitatory or inhibitory neurons with synaptic weight, $w_{lr,E}^i$, and $w_{lr,I}^i$, respectively. The index j represents the source area, i represents the target area, and N is the total number of simulated cortical areas. The axonal delay for long-range connections is given by the ratio between the inter-areal anatomical distance estimates between cortical areas and the conduction speed set as 3.5 m/s (Choi & Mihalas, 2019).

LFP Signal

We computed the local field potential (LFP) signal as a sum of the currents' absolute values acting upon excitatory neurons in a spiking neuronal population (Mazzoni et al., 2015; Mazzoni, Panzeri, Logothetis, & Brunel, 2008). Thus, for a cortical area in our model, the LFP signal will be given by

$$LFP = R \frac{\sum_i^{N_E} (|I_{E,i}| + |I_{I,i}| + |I_{bkg,i}|)}{N_E}. \quad (15)$$

$I_{E,i}$ accounts for both the local (within population) and global (inter-areal projections) excitatory synaptic currents, while $I_{I,i}$ corresponds to the local inhibitory current. $I_{bkg,i}$ is the synaptic current related to the background Poisson input. R represents the resistance of a typical electrode used for extracellular measurements, here chosen to be 1 M Ω (San Cristóbal et al., 2014). N_E is the number of excitatory neurons in each neuronal population.

The mean was subtracted from the simulated LFP signal. The resultant signal was filtered using a 1 kHz low-pass filter to avoid aliasing and downsampled to 1 kHz.

Generalized Partial Directed Coherence

Generalized partial directed coherence (GPDC) is a frequency-domain method of directed functional connectivity established on a multivariate vector autoregressive (MVAR) model (Baccalá et al., 2007). The MVAR model for a set $\mathbf{x}(t) = [x_1(t) \cdots x_N(t)]^T$ of simultaneously observed time series is defined as

$$\mathbf{x}(t) = \sum_{k=1}^p \mathbf{A}_k \mathbf{x}(t-k) + \boldsymbol{\varepsilon}(t), \quad (16)$$

where p is the MVAR model order. \mathbf{A}_k are coefficient matrices in which the element $A_{ij,k}$ define the effect of $x_j(t-k)$ on $x_i(t)$, where k is the time lag. The term $\boldsymbol{\varepsilon}(t)$ is a vector of N white noises

with covariance matrix Σ . The GPDC from the time series x_j to the time series x_i at frequency λ is defined as

$$\text{GPDC}_{ij}(\lambda) = \left| \frac{\frac{1}{\sigma_i} \bar{A}_{ij}(\lambda)}{\sqrt{\sum_{k=1}^N \frac{1}{\sigma_k^2} \bar{A}_{kj}(\lambda) \bar{A}_{kj}^*(\lambda)}}} \right|^2, \quad (17)$$

where

$$\bar{A}_{ij}(\lambda) = \begin{cases} 1 - \sum_{k=1}^p A_{ij,k} e^{-2\pi\sqrt{-1}\lambda k}, & \text{if } i = j \\ - \sum_{k=1}^p A_{ij,k} e^{-2\pi\sqrt{-1}\lambda k}, & \text{if } i \neq j, \end{cases} \quad (18)$$

and σ_i^2 refers to the variance of white noise $\epsilon_i(t)$ (Baccalá et al., 2007). λ is a normalized frequency where $|\lambda| \leq 0.5$ so that $\lambda = 0.5$ means one-half of the sampling rate f_s (Sameshima & Baccalá, 2014).

The MVAR model was estimated by the method of ordinary least squares (OLS; Hamilton, 1994). We used Akaike's information criterion (AIC) to select model order (Supplementary Equation S1), choosing the order $p \leq 50$ that had the minimum AIC (Supplementary Figure S6) value.

GPDC has values in the range from 0 to 1 and is invariant to scale, so the normalization of time series is unnecessary (Baccalá et al., 2007; Sameshima & Baccalá, 2014). Similar to other (directed) functional connectivity measures, unrecorded time series can lead to spurious estimates. Therefore, the reliability of estimates depends on the number of time series included in the estimates. For all analysis we used the peak GPDC value over all frequencies $[0, \frac{f_s}{2}]$.

Estimated Activity

The activity flow mapping measures the propagation of neural activity by estimating the activation of a target region. It is defined as the sum of the activity in each source region multiplied by the functional connectivity with the target region (Cole, Ito, Bassett, & Schultz, 2016). We adapted the idea of activity flow by defining two measures of estimated activity arriving in a cortical area i mediated by pathways of structural connectivity (FLNs) and directed functional connectivity (GPDC peak),

$$p_i^{\text{FLN}} = \sum_{j \neq i \in N} \text{FLN}_{ij} r_j, \quad (19)$$

$$p_i^{\text{GPDC}} = \sum_{j \neq i \in N} \text{GPDC}_{ij} r_j, \quad (20)$$

where FLN_{ij} is the FLN from area j to area i , r_j is the firing rate for area j , GPDC_{ij} is the peak of GPDC from area j to area i , and N is the total number of simulated cortical areas.

Centrality Measure

We computed the nodal in-strength for the mouse cortical connectome. The nodal in-strength for a node i is given by

$$s_i = \sum_{j \neq i \in N} \text{FLN}_{ij}, \quad (21)$$

where j is the source area, i is the target area, and N is the total number of simulated cortical areas (Fornito, Zalesky, & Bullmore, 2016).

Numerical Simulations

All simulations were performed using the simulator Brian2 (Stimberg, Brette, & Goodman, 2019) applying the exponential Euler method (Bower & Beeman, 2012) to integrate the differential equations with an integration step of 0.1 ms. Each simulation was 30 s long, generating sufficient data points to apply GPDC on the simulated LFP signals (Sommariva, Sorrentino, Piana, Pizzella, & Marzetti, 2019).

RESULTS

The large-scale network model of the mouse cortex contains 19 spiking neural populations with recurrent connections and excitatory long-range connections between populations, constrained by the directed and weighted structural connectome (Figure 1A and Figure 1B). The dynamical behavior of each simulated cortical area is predominantly asynchronous with transient spike synchronization (Palmigiano, Geisel, Wolf, & Battaglia, 2017; Uhlhaas et al., 2009) (Figure 1C), with the typical **power spectral density (PSD)** of LFP signals displaying a peak in the gamma band (Figures 1D and 1E) (Buzsáki & Wang, 2012). The firing rate of inhibitory neurons is 4.74 ± 0.11 , higher than the excitatory neurons' rate of 3.64 ± 0.42 (Figure 1F). Differences in population behavior are mostly due to inputs from other areas since we sample their parameters from the same distributions.

We first compared the FLN values with the average GPDC over 10 simulations of the model. Most medium to strong connections from the structural connectome were also captured by the directed functional connectivity (Figure 2A and Figure 2B). We used the GPDC largest value (peak), but other approaches such as the average of GPDC over frequencies and area under the GPDC curve (Supplementary Figure S1) produced similar results.

Although the graph density of the structural connectome is 97% (Gămănuț et al., 2018), most structural connections are weak, which leads to a prevalence of weak average GPDC values. Weak structural connections is a characteristic shared by connectomes from different mammals, with FLNs varying by several orders of magnitude, log-normally distributed (Buzsáki

Power spectral density (PSD): Description of the distribution of the power of a signal in terms of its frequencies.

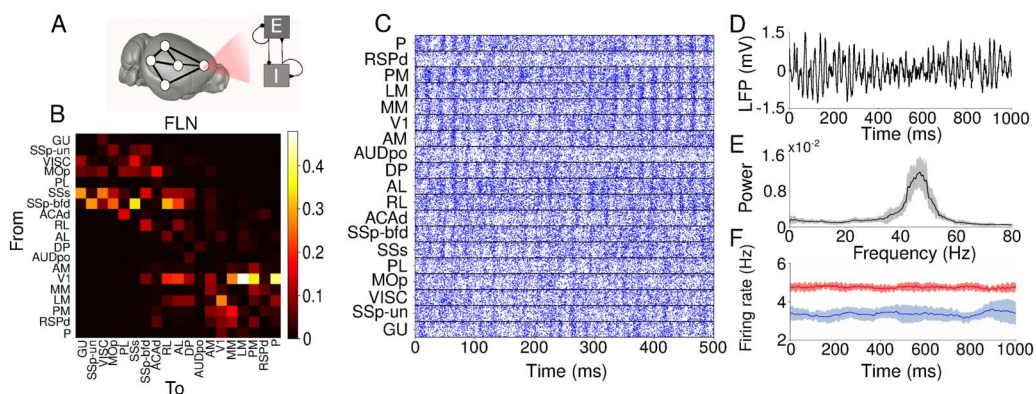


Figure 1. Large-scale cortical network. (A) Local neuronal population where E and I are populations of spiking neurons (Bakker, Tiesinga, & Kötter, 2015; Lein et al., 2007). (B) Map of structural connectivity given by the FLNs. These values define the strength of long-range projections in the large-scale network model. (C) Raster plot of 500 ms of activity for each cortical area. (D) Simulated LFP signal for an area in the large-scale network model. (E) Power spectral density for simulated LFP signal for one area. The continuous black line corresponds to the average over 10 simulations, and the gray shaded area delimits its standard deviation. (F) Firing rate for excitatory (blue) and inhibitory (red) populations computed using a sliding window of 100 ms. The continuous line corresponds to an average firing rate over 10 simulations, and the shaded area is the standard deviation. To exemplify, we used data from area MOp in (D), (E), and (F).

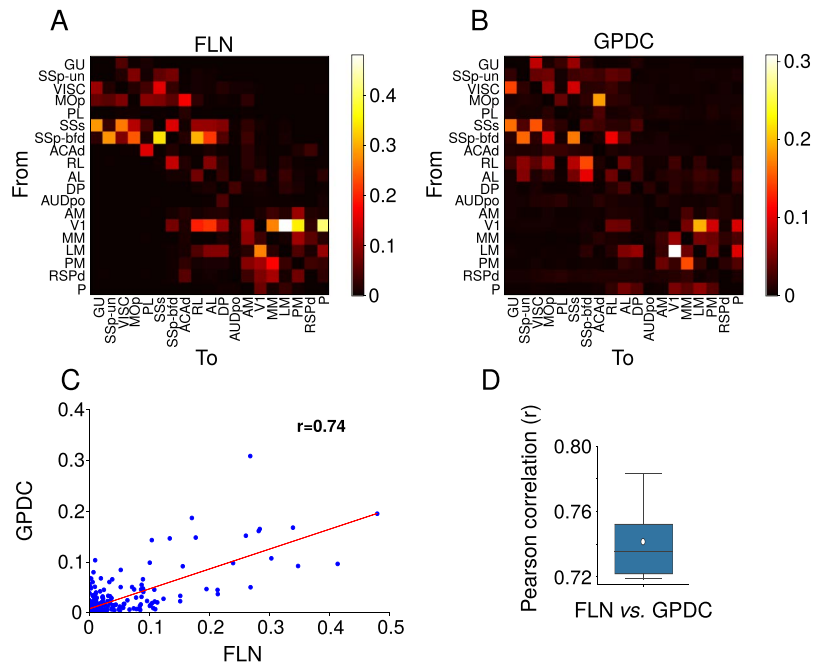


Figure 2. Relation between structural and directed functional connectivity. (A) Map of structural connectivity given by the FLNs. (B) Map of directed functional connectivity given by GPDC peaks for one simulation. GPDC from a cortical area to itself was set as 0. (C) Scatterplot of FLNs versus GPDC peak for one simulation. The red line corresponds to the linear fit. The Pearson correlation between FLNs and GPDC is 0.74. (D) Box plot showing the distribution of Pearson correlation between FLN and GPDC for 10 simulations. The white circle represents the average Pearson correlation over 10 simulations, which is $\bar{r} = 0.74$.

& Mizuseki, 2014; Gămănuț et al., 2018; Markov et al., 2014; Theodoni et al., 2020). To evaluate the relation between structural and directed functional connectivity, we plotted GPDC values from 10 simulations against FLNs and fitted a linear model, obtaining the Pearson correlation r (Figure 2C). The scatterplot presents most points close to the origin due to the predominance of small values for the GPDC and FLN. The average Pearson correlation between FLN and GPDC is 0.74 (Figure 2D). We also verified that the average correlation between GPDC and FLN over bootstrap samples of 80 randomly chosen edges is 0.74 (Supplementary Figure S2). This correlation level is close to those obtained by other works that analyzed different structural connectomes using functional connectivity applied to empirical data ($r \approx 0.79$; Hagmann et al., 2008) or firing rate models ($r \approx 0.73$; Chaudhuri, Knoblauch, Gariel, Kennedy, & Wang, 2015).

The centrality of the cortical area seems to influence the variability of GPDC estimates over simulations. The variability of directed functional connectivity was measured by the **coefficient of variation** of GPDC (Figure 3A). The centrality, measured by the nodal in-strength (i.e., the sum of inward FLNs to a cortical area; Figure 3B), is positively correlated ($r = 0.64$) to the sum of the coefficients of variation (CVs) of the connections emerging from that area (source; Figure 3C). When the cortical area is considered the target of directed functional connectivity, the correlation with nodal in-strength is negative ($r = -0.52$) (Figure 3D). We performed the same analysis correlating the sum of coefficient of variation with eigenvector centrality (Supplementary Figure S3), and we obtained the same relationship, but with smaller Pearson correlation coefficients ($r = 0.59$ and $r = -0.44$). We should note that in both cases (source and target), the actual variability (standard deviation) increases with larger nodal in-strength values (Supplementary Figure S4).

Coefficient of variation (CV):
The ratio between the standard deviation and the mean. It measures the relative dispersion around the mean.

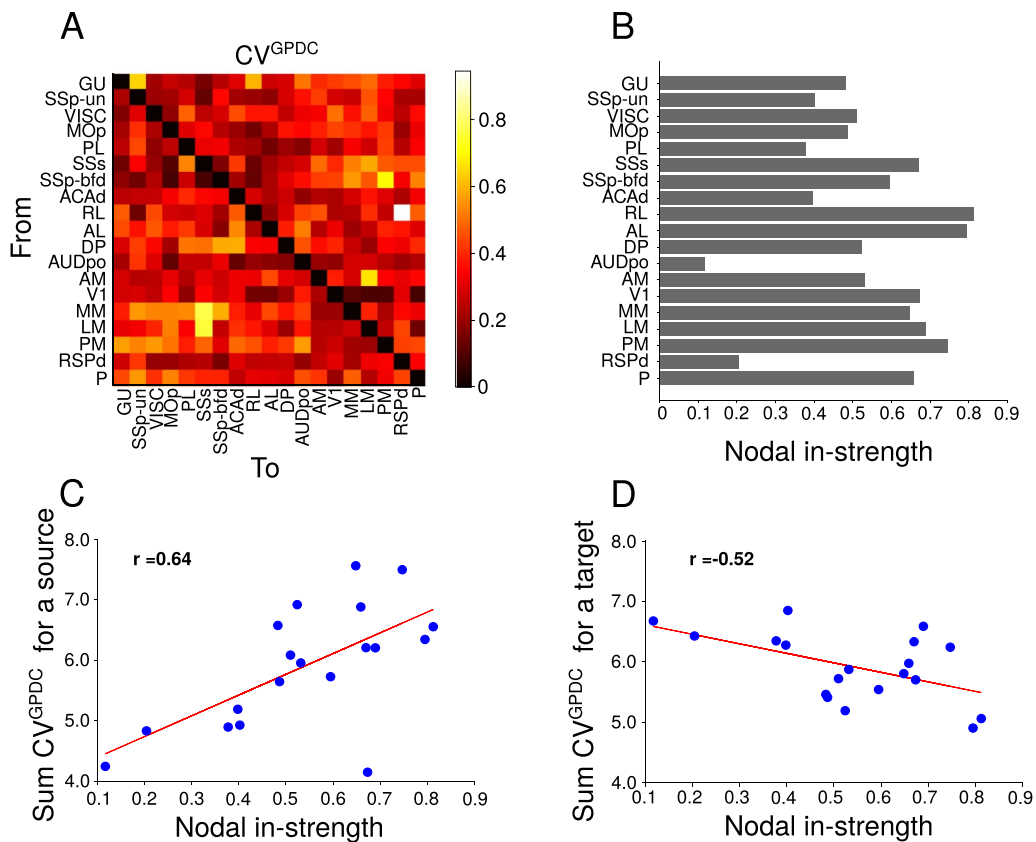


Figure 3. Relationship between nodal in-strength and variability of GPDC. (A) Coefficient of variation for GPDC (CV^{GPDC}). (B) Nodal in-strength for all cortical areas. (C) Sum of CV^{GPDC} for a source (sum of columns in A) versus nodal in-strength. (D) Sum of CV^{GPDC} for a target (sum of rows in A) versus nodal in-strength.

We also investigated the relationship between the firing rate in a cortical area and the estimated activity that is arriving at this cortical area mediated by structural or directed functional connectivity pathways. The propagation of activity in the cortex is constrained by direct anatomical connections between areas and indirect paths (Vélez-Rodríguez, Liu, Hagmann, & Misic, 2020), with the propagation of activity occurring mainly through the strongest long-range projections (Joglekar et al., 2018). The estimated activity mediated by FLNs is strongly correlated to the target areas' firing rate (Figure 4A), while the correlation of estimated activity mediated by GPDCs and firing rates was 0.54 (Figure 4B). This indicates that GPDC estimates can be used to infer the propagation pathways, although less reliably than when using FLN values directly.

We analyzed the behavior of GPDC estimates when considering a reduced number of areas, reproducing typical experimental setups. We considered a visual and a frontoparietal cluster, each containing seven cortical areas (Gămănuț et al., 2018) (Figure 5A). We evaluated the distribution of correlation between FLN and GPDC when GPDC estimates between all areas of each cluster are conditioned on the whole connectome, conditioned on the areas in each cluster, and using only pairwise (bivariate) estimates (Figures 5B and 5C). This analysis simulates the situations where an electrophysiologist has information only from a single cluster of cortical areas or a pair of areas. The highest correlations between the GPDC and FLN occurred when we conditioned GPDC to the whole connectome, followed by GPDC conditioned to the cluster area, and pairwise GPDC. Also, the correlation for the frontoparietal cluster was higher than for the visual cluster in all scenarios.

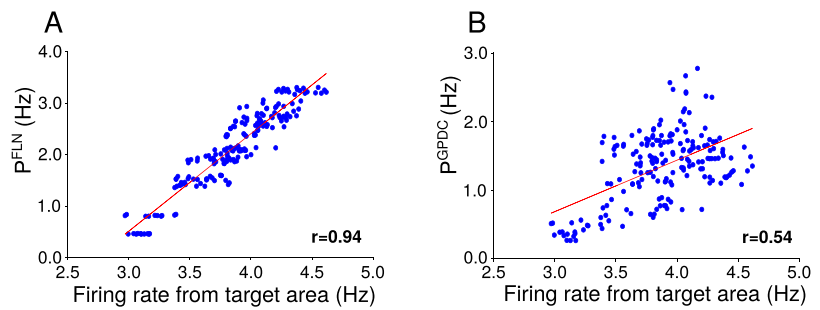


Figure 4. Estimated activity through structural and directed functional pathways. (A) Estimated activity mediated by structural connectivity versus firing rate for target areas ($r = 0.94$). (B) Estimated activity mediated by directed functional connectivity versus firing rate for target areas ($r = 0.54$). Red lines are linear fits.

We extended the analysis to evaluate the effect of cluster size on GPDC correlation to FLN. We used cluster sizes ranging from 3 to 15 areas. We created 150 random clusters sampled from all areas in the connectome for each cluster size and computed the Pearson correlation for the GPDC (a) conditioned on the whole connectome, (b) conditioned on the cluster areas, and (c) evaluated using pairwise data. For cases (a) and (b), the Pearson correlation increases, and the standard variation decreases as we increase the cluster size (Figure 6), showing that it is advantageous to include more areas in the GPDC calculation. Surprisingly, the correlation between structural and directed functional connectivity when using simulated signals from a few cortical areas (blue dots) is similar to using signals from the whole cortex (black dots), with most points showing statistically different results. The bivariate GPDC (Figure 6) had a statistically significant lower average Pearson correlation for all cluster sizes with four or more areas, indicating that these measures are affected by interference from ignored signals.

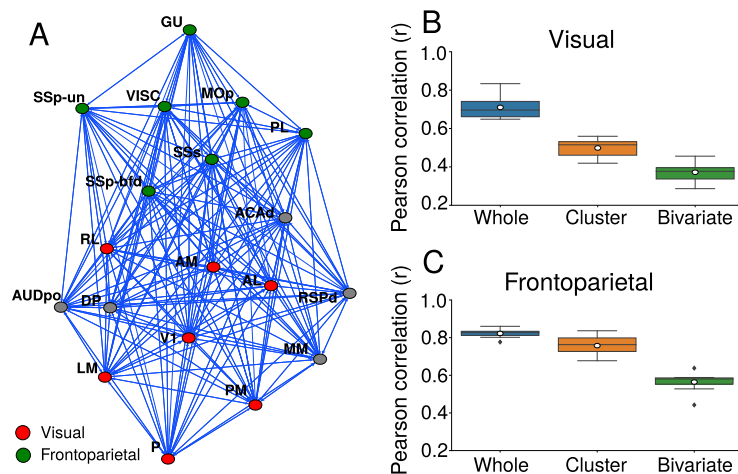


Figure 5. Correlation between FLN and GPDC for the visual and frontoparietal clusters. (A) Graph representing the mouse cortical connectome. Nodes represent cortical areas and edges, directed long-range projections between them. Green nodes are cortical areas belonging to the frontoparietal cluster. Red nodes are cortical areas belonging to the visual cluster. Each cluster contains seven cortical areas. Box plot of Pearson correlation between FLN and GPDC for the visual cluster (B) and frontoparietal cluster (C). GPDC was computed considering the whole connectome (blue box), only the cluster (orange box), and pairwise (green box).

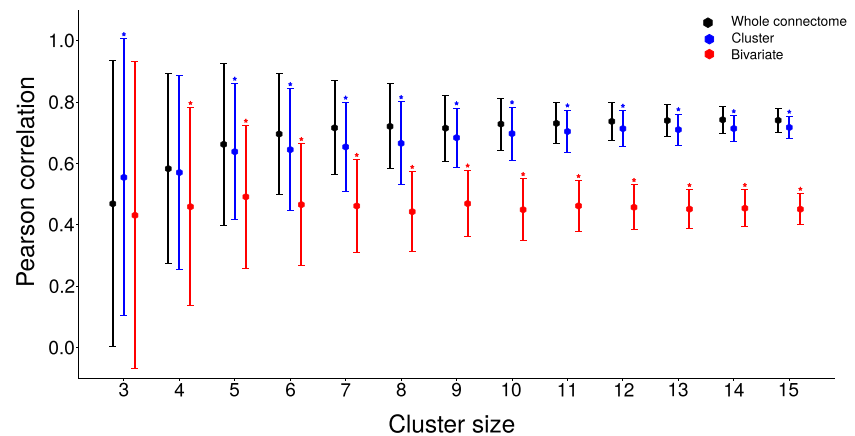


Figure 6. Pearson correlations between FLN and GPDC for the different cluster sizes with randomly chosen areas. The graph shows the average (dots) and error bars of the GPDC conditioned to the whole connectome (in black), to areas in the cluster (in blue), and to pairwise signals (in red). The error bar is the standard deviation of Pearson correlation considering values for all randomly chosen areas and all simulations. Stars represent statistically different averages when compared with the whole connectome (Holm-Bonferroni corrected Welch *t* test).

DISCUSSION

Our results shed light on the relationship between structural and directed functional connectivity in circumstances similar to those faced by electrophysiologists. They indicate that the reliability of directed functional connectivity estimates and their relationship with structural connectivity depends on the number of areas considered. Nevertheless, the GPDC conditioned on few cortical areas had similar results to the GPDC conditioned on all areas, providing evidence that it is possible to obtain a reasonable relationship between structural and directed functional connectivity in electrophysiological experiments even with signals from few areas.

Previous studies evaluated the relationship between structural and functional network connectivity strength on electrophysiological data (Straathof, Sinke, Dijkhuizen, & Otte, 2019), with some using undirected functional connectivity measures (Grandjean, Zerbi, Balsters, Wenderoth, & Rudin, 2017; Stafford et al., 2014). But in electrophysiology studies, researchers do not have access to signals from unrecorded areas and have only estimates of structural strengths from tracers. Using large-scale network models solves this problem, as the researcher has access to all variables in the system, allowing a better understanding of the obtained functional connectivity results.

The relationships between structural and functional connectivity have been largely unexplored through large-scale network models (Bansal, Nakuci, & Muldoon, 2018), and the existing models use neural mass descriptions (rate models) to describe each area's activity (Honey et al., 2009; Novelli & Lizier, 2021). However, information propagated between brain regions can be characterized not only by the rate code but also by the temporal code (Bieler et al., 2017; Hahn, Ponce-Alvarez, Deco, Aertsen, & Kumar, 2019; Kumar, Rotter, & Aertsen, 2010; Luczak, McNaughton, & Harris, 2015; Seth, 2015), and hypotheses are pointing to spike-timing and spike coherence as essential components of cortical communication (Hahn, Bujan, Frégnac, Aertsen, & Kumar, 2014; Palmigiano et al., 2017; Tiesinga & Sejnowski, 2010). Spiking neuronal populations have richer dynamical behaviors than rate models and better resemble cortical activity; through spiking neuronal networks it is possible

to investigate the consequences of spike synchronization (Palmigiano et al., 2017), model different approaches for the propagation of information (Hahn et al., 2019; Joglekar et al., 2018), and generate simulated LFP signals from the synaptic currents, which better resemble biological LFP signals (Mazzoni et al., 2015). Moreover, our obtained correlations are in the same range as the studies using more complex electrophysiological data (Straathof et al., 2019).

The centrality of a cortical area affects the variability of GPDC estimates in different ways when such area is examined as the target or source of functional connections. Strong functional connectivity generally occurs between areas with direct structural connections (Honey et al., 2009), and network measures applied to structural connections can help predict the resting-state functional connectivity (Goñi et al., 2014). However, as far as we know, no previous work has indicated that the variability of directed functional connections could be partially explained by centrality measures applied to structural connectivity. We also noticed that synchronization is strongly correlated to the centrality of the node (Supplementary Figure S5). So it is likely that stronger long-range connections targeting an area increase the synchronization of spikes in this area, and the increased synchronization changes the variability in directed functional connectivity. Indeed, it was observed in previous work that synchronization has an important role in directed functional connectivity (Palmigiano et al., 2017).

The firing rate of cortical areas is explained by the estimated activity flow, as proposed by Cole et al. (2016). When using GPDC as an estimate of structural connections, the correlation between actual and estimated activity in the target area decreases to 0.54. This indicates that directed functional connectivity can be used to estimate the activity flow. Although it is less reliable than when using the actual structural connection strengths, researchers may only have access to directed functional measures.

The relationship between structural and directed functional connectivity is the largest when GPDC is conditioned to all areas in the connectome and decreases as we reduce the number of areas. Gămănuț et al. (2018) identified six clusters in the mouse connectome (prefrontal, frontal, parietal, cingulate, temporal, and visual) based on the same approach used to investigate the macaque cortex (Ercsey-Ravasz et al., 2013). We evaluated the relationship between GPDC and FLNs in the visual cluster and in a combination of the prefrontal, frontal, and parietal clusters, which we called frontoparietal. We did not use the other clusters, which had a small number of regions. The average correlation was in the range of correlation obtained for random clusters, with $r = 0.76$ for the frontoparietal and $r = 0.50$ for the visual cortex. This indicates that within anatomical clusters the relationship between GPDC and FLNs does not change in relation to randomly selected areas. These results also show that GPDC estimates provide statistical information on structural connections even when considering only a few areas. However, when considering individual connections, there can be large differences between GPDC estimates and actual structural connection strengths.

Our large-scale network model has some limitations. First, modeled neuronal population parameters are drawn from the same distributions with activity in the gamma band range (Figure 1). The activity of cortical areas in mice occurs in multiple frequency ranges (Sherman et al., 2016; Tsurugizawa, Djemai, & Zalesky, 2019) and the relationship between structural and functional connectivity depends on the frequency (Vezoli et al., 2020). A second limitation is that we do not model changes in network states, which are known to influence functional connectivity (Stitt et al., 2017). Some studies in computational neuroscience have already explored multistability and temporal patterns of functional connectivity (Deco & Jirsa, 2012; Golos, Jirsa, & Daucé, 2015; Orio et al., 2018). Finally, we considered only cortical areas in our large-scale network model, excluding subcortical areas, which have a more

complex dynamic (Erö, Gewaltig, Keller, & Markram, 2018; Knox et al., 2018). Future studies can overcome these limitations by creating richer spiking network models, with different operating frequencies and evolving neuronal dynamics. These models are difficult to create but would allow one to compare functional connectivity values with structural connection strength in more dynamic settings.

SUPPORTING INFORMATION

Supporting information for this article is available at https://doi.org/10.1162/netn_a_00206.

AUTHOR CONTRIBUTIONS

Ronaldo V. Nunes: Conceptualization; Investigation; Methodology; Project administration; Software; Validation; Visualization; Writing – original draft. Marcelo B. Reyes: Conceptualization; Methodology; Supervision; Writing – review & editing. Jorge F. Mejias: Conceptualization; Methodology; Supervision; Writing – review & editing. Raphael Y. de Camargo: Conceptualization; Methodology; Supervision; Writing – review & editing.

FUNDING INFORMATION

Ronaldo V. Nunes, Coordenação de Aperfeiçoamento de Pessoal de Nível Superior (<https://dx.doi.org/10.13039/501100002322>), Award ID: Finance Code 001.

REFERENCES

- Ambrosen, K. S., Eskildsen, S. F., Hinne, M., Krug, K., Lundell, H., Schmidt, M. N., ... Dyrby, T. B. (2020). Validation of structural brain connectivity networks: The impact of scanning parameters. *NeuroImage*, 204, 116207. <https://doi.org/10.1016/j.neuroimage.2019.116207>, PubMed: 31539592
- Avena-Koenigsberger, A., Misić, B., & Sporns, O. (2018). Communication dynamics in complex brain networks. *Nature Reviews Neuroscience*, 19(1), 17. <https://doi.org/10.1038/nrn.2017.149>, PubMed: 29238085
- Baccalá, L. A., & Sameshima, K. (2001). Partial directed coherence: A new concept in neural structure determination. *Biological Cybernetics*, 84(6), 463–474. <https://doi.org/10.1007/PL00007990>, PubMed: 11417058
- Baccalá, L. A., Sameshima, K., & Takahashi, D. (2007). Generalized partial directed coherence. In *2007 15th international conference on digital signal processing* (pp. 163–166). <https://doi.org/10.1109/ICDSP.2007.4288544>
- Bakker, R., Tiesinga, P., & Kötter, R. (2015). The scalable brain atlas: Instant web-based access to public brain atlases and related content. *Neuroinformatics*, 13(3), 353–366. <https://doi.org/10.1007/s12021-014-9258-x>, PubMed: 25682754
- Bansal, K., Nakuci, J., & Muldoon, S. F. (2018). Personalized brain network models for assessing structure–function relationships. *Current Opinion in Neurobiology*, 52, 42–47. <https://doi.org/10.1016/j.conb.2018.04.014>, PubMed: 29704749
- Barardi, A., Sancristóbal, B., & Garcia-Ojalvo, J. (2014). Phase-coherence transitions and communication in the gamma range between delay-coupled neuronal populations. *PLoS Computational Biology*, 10(7), e1003723. <https://doi.org/10.1371/journal.pcbi.1003723>, PubMed: 25058021
- Barnett, L., & Seth, A. K. (2014). The MVGC multivariate Granger causality toolbox: A new approach to Granger-causal inference. *Journal of Neuroscience Methods*, 223, 50–68. <https://doi.org/10.1016/j.jneumeth.2013.10.018>, PubMed: 24200508
- Bassett, D. S., & Sporns, O. (2017). Network neuroscience. *Nature Neuroscience*, 20(3), 353–364. <https://doi.org/10.1038/nn.4502>, PubMed: 28230844
- Bassett, D. S., Zurn, P., & Gold, J. I. (2018). On the nature and use of models in network neuroscience. *Nature Reviews Neuroscience*, 19(9), 566–578. <https://doi.org/10.1038/s41583-018-0038-8>, PubMed: 30002509
- Bastos, A. M., & Schoffelen, J.-M. (2016). A tutorial review of functional connectivity analysis methods and their interpretational pitfalls. *Frontiers in Systems Neuroscience*, 9, 175. <https://doi.org/10.3389/fnsys.2015.00175>, PubMed: 26778976
- Bieler, M., Sieben, K., Cichon, N., Schildt, S., Röder, B., & Hanganu-Opatz, I. L. (2017). Rate and temporal coding convey multisensory information in primary sensory cortices. *eNeuro*, 4(2). <https://doi.org/10.1523/ENEURO.0037-17.2017>, PubMed: 28374008
- Bower, J. M., & Beeman, D. (2012). *The book of GENESIS: Exploring realistic neural models with the general neural simulation system*. Springer Science & Business Media.
- Buzsáki, G., & Mizuseki, K. (2014). The log-dynamic brain: How skewed distributions affect network operations. *Nature Reviews Neuroscience*, 15(4), 264–278. <https://doi.org/10.1038/nrn3687>, PubMed: 24569488

- Buzsáki, G., & Wang, X.-J. (2012). Mechanisms of gamma oscillations. *Annual Review of Neuroscience*, *35*, 203–225. <https://doi.org/10.1146/annurev-neuro-062111-150444>, PubMed: 22443509
- Chaudhuri, R., Knoblauch, K., Gariel, M.-A., Kennedy, H., & Wang, X.-J. (2015). A large-scale circuit mechanism for hierarchical dynamical processing in the primate cortex. *Neuron*, *88*(2), 419–431. <https://doi.org/10.1016/j.neuron.2015.09.008>, PubMed: 26439530
- Choi, H., & Mihalas, S. (2019). Synchronization dependent on spatial structures of a mesoscopic whole-brain network. *PLoS Computational Biology*, *15*(4), e1006978. <https://doi.org/10.1371/journal.pcbi.1006978>, PubMed: 31013267
- Cole, M. W., Ito, T., Bassett, D. S., & Schultz, D. H. (2016). Activity flow over resting-state networks shapes cognitive task activations. *Nature Neuroscience*, *19*(12), 1718–1726. <https://doi.org/10.1038/nn.4406>, PubMed: 27723746
- Deco, G., & Jirsa, V. K. (2012). Ongoing cortical activity at rest: Criticality, multistability, and ghost attractors. *Journal of Neuroscience*, *32*(10), 3366–3375. <https://doi.org/10.1523/JNEUROSCI.2523-11.2012>, PubMed: 22399758
- de Lange, S. C., Ardesch, D. J., & van den Heuvel, M. P. (2019). Connection strength of the macaque connectome augments topological and functional network attributes. *Network Neuroscience*, *3*(4), 1051–1069. https://doi.org/10.1162/netn_a_00101, PubMed: 31637338
- De Sanctis, B., Vicente, R., Sancho, J. M., & Garcia-Ojalvo, J. (2013). Emergent bimodal firing patterns implement different encoding strategies during gamma-band oscillations. *Frontiers in Computational Neuroscience*, *7*, 18. <https://doi.org/10.3389/fncom.2013.00018>, PubMed: 23532173
- Ercsey-Ravasz, M., Markov, N. T., Lamy, C., Van Essen, D. C., Knoblauch, K., Toroczkai, Z., & Kennedy, H. (2013). A predictive network model of cerebral cortical connectivity based on a distance rule. *Neuron*, *80*(1), 184–197. <https://doi.org/10.1016/j.neuron.2013.07.036>, PubMed: 24094111
- Erő, C., Gewaltig, M.-O., Keller, D., & Markram, H. (2018). A cell atlas for the mouse brain. *Frontiers in Neuroinformatics*, *12*, 84. <https://doi.org/10.3389/fninf.2018.00084>, PubMed: 30546301
- Fornito, A., Zalesky, A., & Bullmore, E. (2016). *Fundamentals of brain network analysis*. Academic Press.
- Gămănuț, R., Kennedy, H., Toroczkai, Z., Ercsey-Ravasz, M., Van Essen, D. C., Knoblauch, K., & Burkhalter, A. (2018). The mouse cortical connectome, characterized by an ultra-dense cortical graph, maintains specificity by distinct connectivity profiles. *Neuron*, *97*(3), 698–715. <https://doi.org/10.1016/j.neuron.2017.12.037>, PubMed: 29420935
- Golos, M., Jirsa, V., & Daucé, E. (2015). Multistability in large scale models of brain activity. *PLoS Computational Biology*, *11*(12), e1004644. <https://doi.org/10.1371/journal.pcbi.1004644>, PubMed: 26709852
- Goñi, J., van den Heuvel, M. P., Avena-Koenigsberger, A., De Mendizabal, N. V., Betzel, R. F., Griffa, A., ... Sporns, O. (2014). Resting-brain functional connectivity predicted by analytic measures of network communication. *Proceedings of the National Academy of Sciences*, *111*(2), 833–838. <https://doi.org/10.1073/pnas.1315529111>, PubMed: 24379387
- Grandjean, J., Zerbi, V., Balsters, J. H., Wenderoth, N., & Rudin, M. (2017). Structural basis of large-scale functional connectivity in the mouse. *Journal of Neuroscience*, *37*(34), 8092–8101. <https://doi.org/10.1523/JNEUROSCI.0438-17.2017>, PubMed: 28716961
- Gutfreund, Y., Yarom, Y., & Segev, I. (1995). Subthreshold oscillations and resonant frequency in guinea-pig cortical neurons: Physiology and modelling. *Journal of Physiology*, *483*(3), 621–640. <https://doi.org/10.1113/jphysiol.1995.sp020611>, PubMed: 7776248
- Hagmann, P., Cammoun, L., Gigandet, X., Meuli, R., Honey, C. J., Wedeen, V. J., & Sporns, O. (2008). Mapping the structural core of human cerebral cortex. *PLoS Biology*, *6*(7), e159. <https://doi.org/10.1371/journal.pbio.0060159>, PubMed: 18597554
- Hahn, G., Bujan, A. F., Frégnac, Y., Aertsen, A., & Kumar, A. (2014). Communication through resonance in spiking neuronal networks. *PLoS Computational Biology*, *10*(8), e1003811. <https://doi.org/10.1371/journal.pcbi.1003811>, PubMed: 25165853
- Hahn, G., Ponce-Alvarez, A., Deco, G., Aertsen, A., & Kumar, A. (2019). Portraits of communication in neuronal networks. *Nature Reviews Neuroscience*, *20*(2), 117–127. <https://doi.org/10.1038/s41583-018-0094-0>, PubMed: 30552403
- Hamilton, J. (1994). *Times series analysis*. Princeton University Press. <https://doi.org/10.1515/9780691218632>
- Harris, K. D., Quiroga, R. Q., Freeman, J., & Smith, S. L. (2016). Improving data quality in neuronal population recordings. *Nature Neuroscience*, *19*(9), 1165–1174. <https://doi.org/10.1038/nn.4365>, PubMed: 27571195
- Honey, C. J., Sporns, O., Cammoun, L., Gigandet, X., Thiran, J.-P., Meuli, R., & Hagmann, P. (2009). Predicting human resting-state functional connectivity from structural connectivity. *Proceedings of the National Academy of Sciences*, *106*(6), 2035–2040. <https://doi.org/10.1073/pnas.0811168106>, PubMed: 19188601
- Hong, G., & Lieber, C. M. (2019). Novel electrode technologies for neural recordings. *Nature Reviews Neuroscience*, *20*(6), 330–345. <https://doi.org/10.1038/s41583-019-0140-6>, PubMed: 30833706
- Joglekar, M. R., Mejias, J. F., Yang, G. R., & Wang, X.-J. (2018). Inter-areal balanced amplification enhances signal propagation in a large-scale circuit model of the primate cortex. *Neuron*, *98*(1), 222–234. <https://doi.org/10.1016/j.neuron.2018.02.031>, PubMed: 29576389
- Knox, J. E., Harris, K. D., Graddis, N., Whitesell, J. D., Zeng, H., Harris, J. A., ... Mihalas, S. (2018). High-resolution data-driven model of the mouse connectome. *Network Neuroscience*, *3*(1), 217–236. https://doi.org/10.1162/netn_a_00066, PubMed: 30793081
- Kumar, A., Rotter, S., & Aertsen, A. (2010). Spiking activity propagation in neuronal networks: Reconciling different perspectives on neural coding. *Nature Reviews Neuroscience*, *11*(9), 615–627. <https://doi.org/10.1038/nrn2886>, PubMed: 20725095
- Lein, E. S., Hawrylycz, M. J., Ao, N., Ayres, M., Bensinger, A., Bernard, A., ... Jones, A. R. (2007). Genome-wide atlas of gene expression in the adult mouse brain. *Nature*, *445*(7124), 168–176. <https://doi.org/10.1038/nature05453>, PubMed: 17151600

- Luczak, A., McNaughton, B. L., & Harris, K. D. (2015). Packet-based communication in the cortex. *Nature Reviews Neuroscience*, 16(12), 745–755. <https://doi.org/10.1038/nrn4026>, PubMed: 26507295
- Markov, N. T., Ercsey-Ravasz, M., Ribeiro Gomes, A., Lamy, C., Magrou, L., Vezoli, J., ... Kennedy, H. (2014). A weighted and directed interareal connectivity matrix for macaque cerebral cortex. *Cerebral Cortex*, 24(1), 17–36. <https://doi.org/10.1093/cercor/bhs270>, PubMed: 23010748
- Mazzoni, A., Linden, H., Cuntz, H., Lansner, A., Panzeri, S., & Einevoll, G. T. (2015). Computing the local field potential (LFP) from integrate-and-fire network models. *PLoS Computational Biology*, 11(12). <https://doi.org/10.1371/journal.pcbi.1004584>, PubMed: 26657024
- Mazzoni, A., Panzeri, S., Logothetis, N. K., & Brunel, N. (2008). Encoding of naturalistic stimuli by local field potential spectra in networks of excitatory and inhibitory neurons. *PLoS Computational Biology*, 4(12), e1000239. <https://doi.org/10.1371/journal.pcbi.1000239>, PubMed: 19079571
- Mi, X., Cheng, N., & Zhang, T. (2014). Performance comparison between GPDC and PCMI for measuring directionality of neural information flow. *Journal of Neuroscience Methods*, 227, 57–64. <https://doi.org/10.1016/j.jneumeth.2014.02.006>, PubMed: 24548795
- Novelli, L., & Lizier, J. T. (2021). Inferring network properties from time series using transfer entropy and mutual information: Validation of multivariate versus bivariate approaches. *Network Neuroscience*, 5(2), 373–404. https://doi.org/10.1162/netn_a_00178, PubMed: 34189370
- Novelli, L., Wollstadt, P., Mediano, P., Wibral, M., & Lizier, J. T. (2019). Large-scale directed network inference with multivariate transfer entropy and hierarchical statistical testing. *Network Neuroscience*, 3(3), 827–847. https://doi.org/10.1162/netn_a_00092, PubMed: 31410382
- Nunes, R. V., Reyes, M. B., & De Camargo, R. Y. (2019). Evaluation of connectivity estimates using spiking neuronal network models. *Biological Cybernetics*, 113(3), 309–320. <https://doi.org/10.1007/s00422-019-00796-8>, PubMed: 30783758
- Orio, P., Gatica, M., Herzog, R., Maidana, J. P., Castro, S., & Xu, K. (2018). Chaos versus noise as drivers of multistability in neural networks. *Chaos: An Interdisciplinary Journal of Nonlinear Science*, 28(10), 106321. <https://doi.org/10.1063/1.5043447>, PubMed: 30384618
- Palmigiano, A., Geisel, T., Wolf, F., & Battaglia, D. (2017). Flexible information routing by transient synchrony. *Nature Neuroscience*, 20(7), 1014. <https://doi.org/10.1038/nn.4569>, PubMed: 28530664
- Reid, A. T., Headley, D. B., Mill, R. D., Sanchez-Romero, R., Uddin, L. Q., Marinazzo, D., ... Cole, M. W. (2019). Advancing functional connectivity research from association to causation. *Nature Neuroscience*, 22(11), 1751–1760. <https://doi.org/10.1038/s41593-019-0510-4>, PubMed: 31611705
- Sameshima, K., & Baccalá, L. A. (2014). *Methods in brain connectivity inference through multivariate time series analysis*. CRC Press.
- Sanchez-Romero, R., & Cole, M. W. (2019). Combining multiple functional connectivity methods to improve causal inferences. *Journal of Cognitive Neuroscience*, 1–15. https://doi.org/10.1162/jocn_a_01580, PubMed: 32427070
- Sancristóbal, B., Vicente, R., & Garcia-Ojalvo, J. (2014). Role of frequency mismatch in neuronal communication through coherence. *Journal of Computational Neuroscience*, 37(2), 193–208. <https://doi.org/10.1007/s10827-014-0495-7>, PubMed: 24519733
- Schölvinck, M. L., Leopold, D. A., Brookes, M. J., & Khader, P. H. (2013). The contribution of electrophysiology to functional connectivity mapping. *NeuroImage*, 80, 297–306. <https://doi.org/10.1016/j.neuroimage.2013.04.010>, PubMed: 23587686
- Seth, A. K. (2015). Neural coding: Rate and time codes work together. *Current Biology*, 25(3), R110–R113. <https://doi.org/10.1016/j.cub.2014.12.043>, PubMed: 25649819
- Sherman, M. A., Lee, S., Law, R., Haegens, S., Thorn, C. A., Hämäläinen, M. S., ... Jones, S. R. (2016). Neural mechanisms of transient neocortical beta rhythms: Converging evidence from humans, computational modeling, monkeys, and mice. *Proceedings of the National Academy of Sciences*, 113(33), E4885–E4894. <https://doi.org/10.1073/pnas.1604135113>, PubMed: 27469163
- Sommariva, S., Sorrentino, A., Piana, M., Pizzella, V., & Marzetti, L. (2019). A comparative study of the robustness of frequency-domain connectivity measures to finite data length. *Brain Topography*, 32(4), 675–695. <https://doi.org/10.1007/s10548-017-0609-4>, PubMed: 29168017
- Sporns, O., Tononi, G., & Kötter, R. (2005). The human connectome: A structural description of the human brain. *PLoS Computational Biology*, 1(4), e42. <https://doi.org/10.1371/journal.pcbi.0010042>, PubMed: 16201007
- Stafford, J. M., Jarrett, B. R., Miranda-Dominguez, O., Mills, B. D., Cain, N., Mihalas, S., ... Fair, D. A. (2014). Large-scale topology and the default mode network in the mouse connectome. *Proceedings of the National Academy of Sciences*, 111(52), 18745–18750. <https://doi.org/10.1073/pnas.1404346111>, PubMed: 25512496
- Stimberg, M., Brette, R., & Goodman, D. F. (2019). Brian 2, an intuitive and efficient neural simulator. *eLife*, 8, e47314. <https://doi.org/10.7554/eLife.47314>, PubMed: 31429824
- Stitt, I., Hollensteiner, K. J., Galindo-Leon, E., Pieper, F., Fiedler, E., Stieglitz, T., ... Engel, A. K. (2017). Dynamic reconfiguration of cortical functional connectivity across brain states. *Scientific Reports*, 7(1), 1–14. <https://doi.org/10.1038/s41598-017-08050-6>, PubMed: 28821753
- Straathof, M., Sinke, M. R., Dijkhuizen, R. M., & Otte, W. M. (2019). A systematic review on the quantitative relationship between structural and functional network connectivity strength in mammalian brains. *Journal of Cerebral Blood Flow and Metabolism*, 39(2), 189–209. <https://doi.org/10.1177/0271678X18809547>, PubMed: 30375267
- Theodoni, P., Majka, P., Reser, D. H., Wójcik, D. K., Rosa, M. G., & Wang, X.-J. (2020). Structural attributes and principles of the neocortical connectome in the marmoset monkey. *bioRxiv*. <https://doi.org/10.1101/2020.02.28.969824>
- Tiesinga, P. H., & Sejnowski, T. J. (2010). Mechanisms for phase shifting in cortical networks and their role in communication through coherence. *Frontiers in Human Neuroscience*, 4, 196. <https://doi.org/10.3389/fnhum.2010.00196>, PubMed: 21103013

- Tomov, P., Pena, R. F., Zaks, M. A., & Roque, A. C. (2014). Sustained oscillations, irregular firing, and chaotic dynamics in hierarchical modular networks with mixtures of electrophysiological cell types. *Frontiers in Computational Neuroscience*, 8, 103. <https://doi.org/10.3389/fncom.2014.00103>, PubMed: 25228879
- Tsurugizawa, T., Djemai, B., & Zalesky, A. (2019). The impact of fasting on resting state brain networks in mice. *Scientific Reports*, 9(1), 1–12. <https://doi.org/10.1038/s41598-019-39851-6>, PubMed: 30814613
- Uhlhaas, P., Pipa, G., Lima, B., Melloni, L., Neuenschwander, S., Nikolić D., & Singer, W. (2009). Neural synchrony in cortical networks: History, concept and current status. *Frontiers in Integrative Neuroscience*, 3, 17. <https://doi.org/10.3389/neuro.07.017.2009>, PubMed: 19668703
- Vezoli, J., Vinck, M., Bosman, C. A., Bastos, A. M., Lewis, C. M., Kennedy, H., & Fries, P. (2020). The role of anatomical connection strength for interareal communication in macaque cortex. Available at SSRN 3751057. <https://doi.org/10.2139/ssrn.3751057>
- Vézquez-Rodríguez, B., Liu, Z.-Q., Hagmann, P., & Misic, B. (2020). Signal propagation via cortical hierarchies. *Network Neuroscience*, 4(4), 1072–1090. https://doi.org/10.1162/netn_a_00153, PubMed: 33195949