

Accepted Manuscript
Accepted Manuscript (Uncorrected Proof)

Title: Functional Connectivity Alterations of Within and Between Networks in Schizophrenia:
A Retrospective Study

Running Title: Networks' dysconnectivity in Schizophrenia

Authors: Farzaneh Keyvanfard^{1,*}, Anna-Katharina Schmid², Abbas Nasiraei Moghaddam³

1. *Post-Doctoral Research Fellow, School of Cognitive Sciences, Institute for Research in Fundamental Sciences (IPM), Tehran, Iran.*
2. *M.Sc. Student of Neurosciences, Department of Health Sciences and Technology (D-HEST), ETH Zürich, Zürich, Switzerland.*
3. *Associate Professor, Biomedical Engineering Department, Amirkabir University of Technology, Tehran, Iran.*

***Corresponding Author:** Farzaneh Keyvanfard, Post-Doctoral Research Fellow, School of Cognitive Sciences, Institute for Research in Fundamental Sciences (IPM), Tehran, Iran. Email: f.keyvanfard@ipm.ir, f.keyvanfard@gmail.com

To appear in: **Basic and Clinical Neuroscience**

Received date: 2022/01/10

Revised date: 2022/03/27

Accepted date: 2022/05/22

This is a “Just Accepted” manuscript, which has been examined by the peer-review process and has been accepted for publication. A “Just Accepted” manuscript is published online shortly after its acceptance, which is prior to technical editing and formatting and author proofing. *Basic and Clinical Neuroscience* provides “Just Accepted” as an optional and free service which allows authors to make their results available to the research community as soon as possible after acceptance. After a manuscript has been technically edited and formatted, it will be removed from the “Just Accepted” Web site and published as a published article. Please note that technical editing may introduce minor changes to the manuscript text and/or graphics which may affect the content, and all legal disclaimers that apply to the journal pertain.

Please cite this article as:

Keyvanfard, F., Schmid, A. K., Nasiraei Moghaddam, A. (In Press). Functional Connectivity Alterations of Within and Between Networks in Schizophrenia: A Retrospective Study. *Basic and Clinical Neuroscience*. Just Accepted publication Aug. 15, 2022. Doi: <http://dx.doi.org/10.32598/bcn.2022.3928.2>

DOI: <http://dx.doi.org/10.32598/bcn.2022.3928.2>

Accepted Manuscript (Uncorrected Proof)

Abstract

Schizophrenia (SZ) is a chronic brain disorder characterized by diverse cognitive dysfunctions due to abnormal brain connectivity. Evaluating these connectivity alterations between and within such networks (intra- and inter connectivity) may improve the understanding of disrupted information processing patterns in SZ patients. For this reason, resting-state fMRI analysis was performed on 24 SZ patients and 27 matched healthy controls. A functional connectivity matrix was constructed for each participant based on 129 gray matter regions. All regions were classified into eight distinct functional networks. Afterwards, all functional connections were segregated into inter- and intra-network connections considering the eight networks. The mean values of connectivity weights and nodal strength were examined for within-and between-network connections in SZ patients and healthy controls. This analysis revealed that the within-network connections in the somatomotor network were significantly reduced (p -value <0.001) in SZ patients. Additionally, intra-network connections within the visual and the ventral attention networks were proven to be significantly lower (p -value <0.01) in the SZ group. Moreover, disrupted intra-network connectivity was detected between the following network pairs: the visual-limbic, the somatomotor-limbic, the dorsal attention-limbic, and the ventral attention-dorsal attention system. Overall, an extensive reduction in functional connectivity strength for SZ patients was illustrated, with a particularly significant decrease in intra-network connections when compared to the inter-networks. These findings can impact the understanding of the important dysregulated connections that are implicated in the incidence of Schizophrenia.

Keywords: Schizophrenia, Cognitive dysfunction, fMRI, Resting state networks, Inter-network connectivity.

1. Introduction

Schizophrenia (SZ) is a mental disorder affecting approximately 1% of the world's population. It is primarily associated with dopamine dysfunction (Stahl, 2018; Yang & Tsai, 2017) and characterized by diverse clinical appearances such as the disintegration of emotional responsiveness and thought processes. The emergence of SZ is a result of alterations in interactions between two or more brain regions, not regionally isolated pathologies (Andreasen et al., 1998; Friston & Frith, 1995; Stephan et al., 2009; Van Den Heuvel & Fornito, 2014). The findings of previous brain functional (Correa et al., 2009; Lawrie et al., 2002, 2008; J. Liu et al., 2009; Manoach et al., 2000) and structural studies (Dietsche et al., 2017; Karlsgodt et al., 2010; Lawrie et al., 2008; Stephen et al., 2013) have established that a considerable of brain regions are involved in this disorder.

Resting-state functional magnetic resonance imaging (rs-fMRI) is a popular analyzing method for functional connectivity (FC) assessment in brain networks. The rs-fMRI method is based on the temporally correlated blood oxygen level-dependent (BOLD) signals between distinct brain regions; thereby, FC analysis in a resting state condition can detect the coherent spontaneous neuronal activity within brain networks [12]. Many studies have employed a seed-based approach in which a prior hypothesis about a varied region of interest (ROI) in SZ was used. However, it is also common practice to identify the alterations of a functionally connected brain at a “whole-brain” level in different disorders.

A number of SZ studies have examined the relationship strength between pairs of brain regions across the entire brain (Fornito et al., 2012; Karbasforoushan & Woodward, 2013; Lynall et al., 2010). Furthermore, network characteristics have been widely investigated by many researchers with the graph theory method (Cabral et al., 2012; Karbasforoushan & Woodward, 2013; Y. Liu et al., 2008; Lynall et al., 2010; Micheloyannis, 2012). Overall, findings of previous studies supported the existence a functional dysconnectivity in SZ patients

through the decrease of FC strength (Fornito et al., 2012; Skåtun, Kaufmann, Doan, Alnæs, Córdova-Palomera, Jönsson, Fatouros-Bergman, Flyckt, KaSP, et al., 2017; Skudlarski et al., 2010; Tu et al., 2013; Wei et al., 2018) and a reduction in other network measurements such as the clustering coefficient and efficiency (Algunaid et al., 2018; He et al., 2012; Karbasforoushan & Woodward, 2013; Y. Liu et al., 2008; Rubinov & Bullmore, 2013; Yu et al., 2011).

On the other hand, the complex brain network can be considered as a number of modulated (sub)networks which are functionally linked brain regions sharing specific roles and responsibilities (Keyvanfard et al., 2020; Sporns, 2013; Sporns & Betzel, 2016; van den Heuvel & Hulshoff Pol, 2010).

The within- and between connectivity of these networks change during brain development and in disorder progress (Chan et al., 2014; Sporns & Betzel, 2016). Therefore, the characterization of alterations of within- and between modules SZ is important and has attracted increasing attention in recent years. For instance, decreased FC between the frontoparietal network and the visual networks were reported for SZ patients Wu et al., 2017. Other studies have reported mixed findings. An increased FC was found between the default mode and the central executive networks (Manoliu et al., 2014), an increased FC between the sensory processing and the default mode network, (Tu et al., 2013) and a decrease in FC within the default mode network (Li et al., 2019) were detected in SZ patients. Moreover, prominent cortico-subcortical disconnections within the fronto-parietal network (Tu et al., 2013) and disrupted connectivity among their nodes (Shinn et al., 2015) were demonstrated for SZ patients.

However, the reason for this discrepancy within and between established resting-state networks (RSNs) remains unclear due to mixed and different results (Hummer et al., 2020; Manoliu et al., 2014). Several studies have examined within and between-network connectivity

separately (Skåtun, Kaufmann, Doan, Alnæs, Córdova-Palomera, Jönsson, Fatouros-Bergman, Flyckt, KaSP, et al., 2017; Skudlarski et al., 2010) or focused on considering particular networks and not all RSNs (Mamah et al., 2013; Manoliu et al., 2014; Tu et al., 2013). In other words, their analysis was not performed comprehensively covering whole the brain.

This study aimed to understand how interactions inside RSNs or between them changed in SZ patients in compare to healthy participants. Therefore, alterations in brain functional dysconnectivity between and within RSNs were examined in SZ patients and healthy controls (HC) along with a whole brain network analysis. Additionally, nodal strength variation was assessed to characterize an alteration pattern for inter-and intra-network comparison to whole brain. It was hypothesized that the between network connectivity is most affected by the SZ disorder.

2. Materials and Methods

2.1. Participants

Rs-fMRI analysis was performed in 27 SZ subjects (mean age, 41.9 ± 9.6) and 27 age-matched healthy individuals (mean age, 35 ± 6.8). The dataset is available on the Zenodo platform (Gutiérrez-Gómez et al., 2020; Vohryzek et al., 2020). The patients in the schizophrenic group were recruited from the Service of General Psychiatry at the Lausanne University Hospital. They met DSM-IV criteria for schizophrenic and schizoaffective disorders (American psychiatry association, 2000¹). Healthy control with major mood, psychotic, or substance-use disorders and having first-degree relative with a psychotic disorder were excluded in this database. Moreover, a history of neurological disease was an exclusion criterion for all subjects (Gutiérrez-Gómez et al., 2020). The data analysis was further carried out in 24 out of 27 patients of the SZ group because only 24 subjects were under an equivalent

¹ American Psychiatric Association (2000): Diagnostic and Statistical Manual of Mental Disorders, 4th ed. DSM-IV-TR. American Psychiatric Pub, Arlington, VA22209, USA.

medication dose of chlorpromazine (CPZ) (average medication 431 ± 288 mg) (Andreasen et al., 2010). The written consent was obtained for all subjects in accordance with the institutional guidelines of the Ethics Committee of Clinical Research of the Faculty of Biology and Medicine, University of Lausanne, Switzerland, #82/14, #382/11, #26.4.2005). Data of all subjects was fully anonymized.

2.2. Data acquisition and preparation

For each participant, a T1-weighted anatomical image and a rs-fMRI scan were acquired using a 3T Siemens Trio Scanner equipped with a 32-channel head coil. The magnetization-prepared rapid acquisition gradient echo (MPRAGE) sequence was applied for T1-weighted imaging with a resolution of $1 \times 1 \times 1.2$ mm³ and TI/TE/TR = 900/2.98/2300 ms. Gray matter was partitioned into 129 cortical regions of interest (ROI) including 114 cortical ROIs, 15 subcortical nuclei along with the brain stem, by employing the Desikan Killian atlas (Desikan et al., 2006). Each rs-fMRI scan covered a time span of 8 minutes with a 3.3mm voxel size (isotropic) and TE/TR=30/1920ms. Data preprocessing included exclusion of the first four timepoints of signals, regressing out of physiological signals (white-matter and cerebrospinal fluid), motion correction, spatial smoothing, bandpass filtering, and a linear registration to the T1-weighted image was performed in the whole dataset. Finally, the functional matrices were obtained by computing the absolute value of the Pearson correlation between individual brain regions' time courses to extract the strength of the connections. Consequently, one connectivity matrix was generated with the $K \times K$ dimension (with K=129 numbers of brain regions) for each participant. Further details of the data acquisition and the preparation steps can be found in Vohryzek et al., 2020. This workflow is shown in Fig. 1.

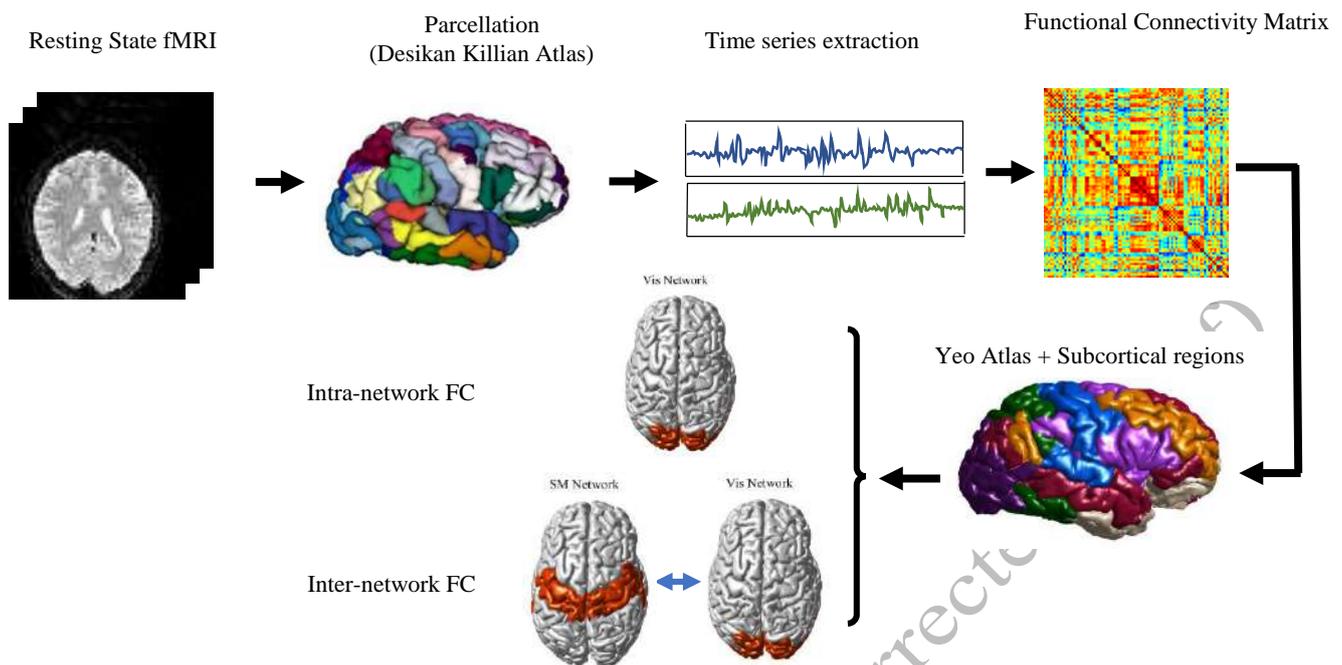


Fig. 1.

2.3. Data Analysis and Statistics

For the resting-state connectivity analysis of this study, a well-known RSNs map from Yeo, (Yeo et al., 2011) was adopted, which introduces seven functional subnetworks. In their parcellation, the cortex was divided into the visual (VIS), somatomotor (SM), fronto-parietal (FP), dorsal attention (DA), ventral attention (VA), default mode (DMN), and limbic (Limb) functional systems (Yeo et al., 2011). In addition to these seven cortical regions, there is one subnetwork that comprised all subcortical (non-cerebellar) parts of the brain. Subsequently, each brain region was allocated to one of these in total eight functional subnetworks.

The group difference analysis between the control and the SZ group was conducted in three steps. First, in order to compare the connectivity strengths between both groups, the mean weight for every connection was computed aside from the networks they belong to. Likewise, nodal strength was calculated with the Brain Connectivity Toolbox (Rubinov & Sporns, 2010)

and the values of the two groups were compared. In the second step, all connections were classified into two sets of inter- and intra-network connections. Inter-network connections are defined as the links between pairs of regions from two different networks, whereas intra-network connections are determined as the connections between pairs of regions inside each network. To understand the severity of changes in inter- and intra-network connections, the mean values of these two connection sets were assessed between SZ patients and HC. As the third step, the inter- and intra-network connectivity of all eight brain networks were statistically compared between the control and patient group. All statistical analyses involved a paired sample t-test. Bonferroni correction was used to correct for multiple comparisons. Finally, results with a p-value < 0.05 were considered statistically significant. The entire analysis was performed using the MATLAB version 2021a.

3. Results

3.1. Whole brain connectivity comparison

The statistical analysis of all connections in SZ patients compared to the ones in healthy individuals revealed a significant whole-brain difference for 1176 connections among 8256 edges (14.24%). For a clearer visual representation, the altered brain connectivity was mapped onto a cortical surface map. Fig. 2(a) displays the regions including nodes with altered connections; accordingly, the different colors represent the normalized values in $[0, 1]$. The normalized value is determined by the nodal strength divided by the largest value of strength. More specifically, one of the compromised regions with several changed edges was related to a part of the somatomotor network, can be seen in red in Fig. 2(a).

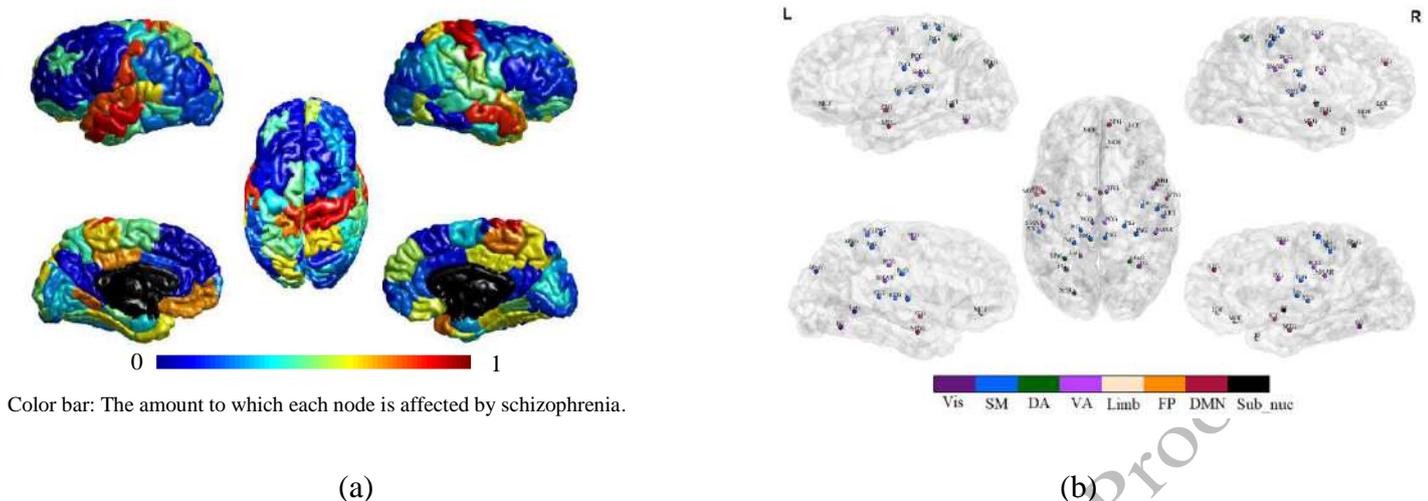


Fig. 2.

Regarding the inter-and intra-network variations perspective, a higher percentage of intra-network edges (19.47%) seemed to be changed when compared to the inter-network (13.41%). The group comparison of the nodal strength was accomplished. It is commonly defined as the summation of all connection weights linked to the node. This comparison exhibited a significant difference in 37 nodes. As displayed in Fig. 2(b), the differences of the cortex nodal strength were observed in the bilateral precentral gyrus (PrG), posterior cingulate gyrus (PCG), superior frontal gyrus (STG), postcentral gyrus (PoG), paracentral lobule (PaG), superior marginal gyrus (SMAR), superior parietal gyrus (SPaG), superior frontal gyrus (SFG), Medial orbito frontal gyrus (MOF), Fusiform gyrus (FG), middle temporal gyrus (MTG), insula (Ins) and right temporal pole (TP), left lingual gyrus (LgG) and transverse temporal gyrus (trTG). These regions partly overlap with the map of the altered edges in Fig. 2(a). The abbreviations of all region names are listed in Table. 1.

Table 1.

Abbreviation	Region Name	Abbreviation	Region Name
LOF	Lateral orbito frontal gyrus	SPaG	Superior parietal gyrus
PORB	Pars orbitalis	IPaG	Inferior parietal gyrus
FP	Frontal pole	PCUN	Precuneus
MOF	Medial orbito frontal gyrus	Cun	Cuneus
PTRI	Pars triangularis	PCAL	Pericalcarine cortex
POPE	Pars opercularis	LOGG	Lateral occipital gyrus
rosMFG	Middle frontal gyrus, rostral	LgG	Lingual gyrus
SFG	Superior frontal gyrus	FG	Fusiform gyrus
caMFG	Middle frontal gyrus, caudal	PHG	Para-hippocampal gyrus
PrG	Precentral gyrus	EC	Entorhinal cortex
PaG	Paracentral lobule	TP	Temporal pole
rosACG	Anterior cingulate gyrus, rostral	ITG	Inferior temporal gyrus
caACG	Anterior cingulate gyrus, caudal	MTG	Middle temporal gyrus
PCG	Posterior cingulate gyrus	bnkST	Bankssts
ICG	Isthmus cingulate gyrus	STG	Superior temporal sulcus
PoG	Postcentral gyrus	trTG	Transverse temporal gyrus
SMAR	Super marginal gyrus	Ins	Insula

3.2. Intra-and inter-network comparisons

After the classification of all connections into inter-and intra-networks, a statistical analysis was applied for both connectivity groups, the SZ patients and the control subjects. A t-test was carried out from two different viewpoints. In the first approach, the weights for inter-and intra-networks were averaged for each subject of the two groups and therefore, two vectors with 27 and 24 elements (representing the number of participants in the healthy and SZ group) were obtained. The control group showed a significantly higher intra-network FC than the SZ groups. On the other hand, no significant difference was detected between the inter-network edges as shown in Fig. 3. Furthermore, patients generally showed greater variance in both inter-

and intra-network connections (p-value <0.0001) which illustrates greater diversity in functional connectivity in SZ patients (Bassett et al., 2012).

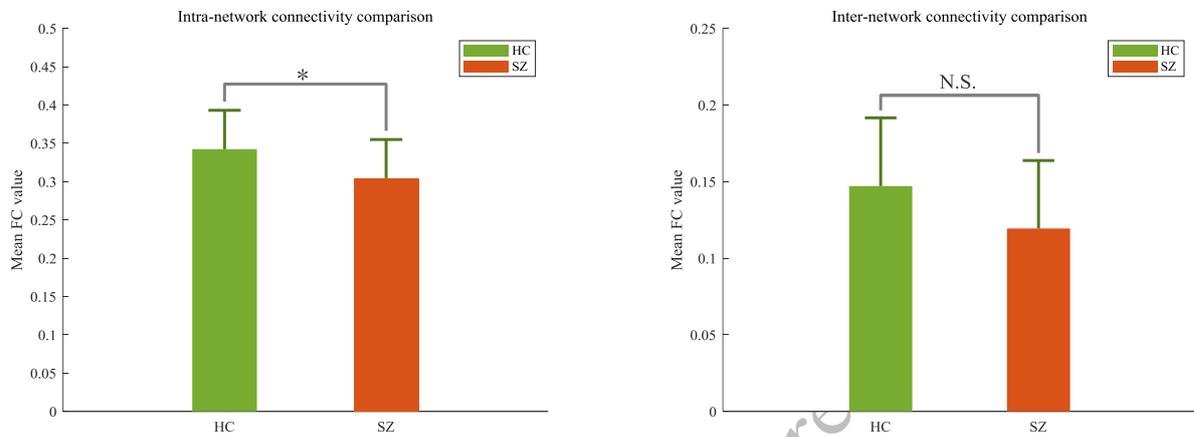


Fig. 3.

In the second comparison approach, a two-sampled t-test was applied on each inter-network connection as well as on the intra-network connection, which revealed a significant change in 955 out of 7121 (13.41 %) and 221 connections out of 1135 (19.47 %) for the inter-and intra-network respectively.

3.3. Intra-and inter-network comparisons of eight networks

Comparing the inter-and the intra-networks between healthy individuals and patients indicated that control subjects had higher connectivity weights within the RSNs (Fig. 4 (a)). However, this difference was significant for the visual, somatomotor, and ventral attention networks. The inter-network connectivity assessment revealed that connections between the limbic system and three networks including the visual, the somatomotor, and the dorsal attention networks were disrupted in SZ. In addition, the connections between the dorsal attention and the ventral attention networks were significantly weaker in the SZ group (Fig. 4 (a-c)). No other between-network differences appeared to be remarkably affected (Fig. 4 (b)).

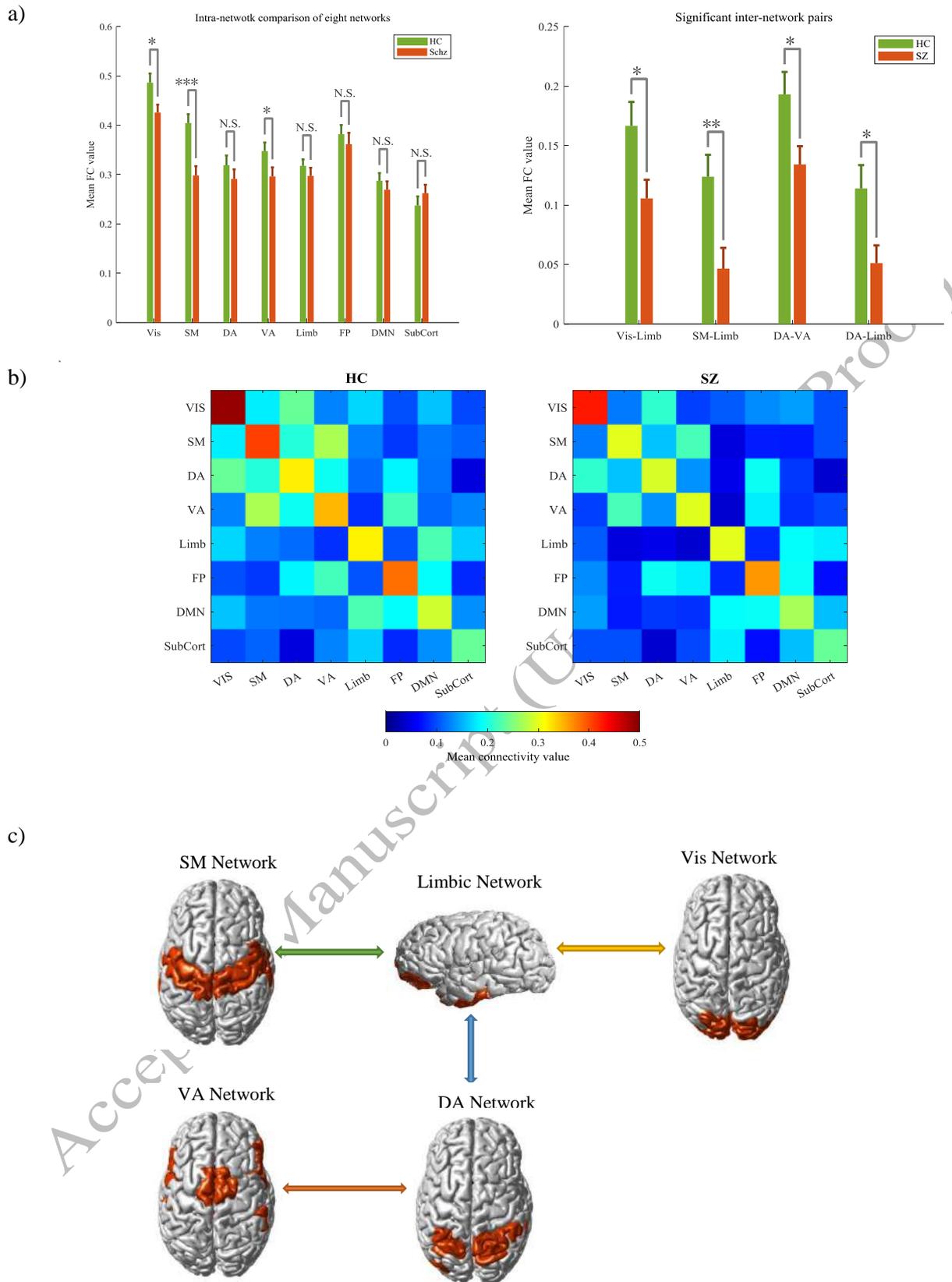


Fig. 4.

The affected nodal strength is illustrated in Fig. 5 indicating that the networks demonstrated a reduced inter- and intra-network connectivity. As illustrated in Fig. 5 (a), the affected nodal strength of intra-networks is concentrated in the somatomotor network. Nodes that are involved in varying inter-network connectivity between the four mentioned networks are also shown in Fig. 5 (b).

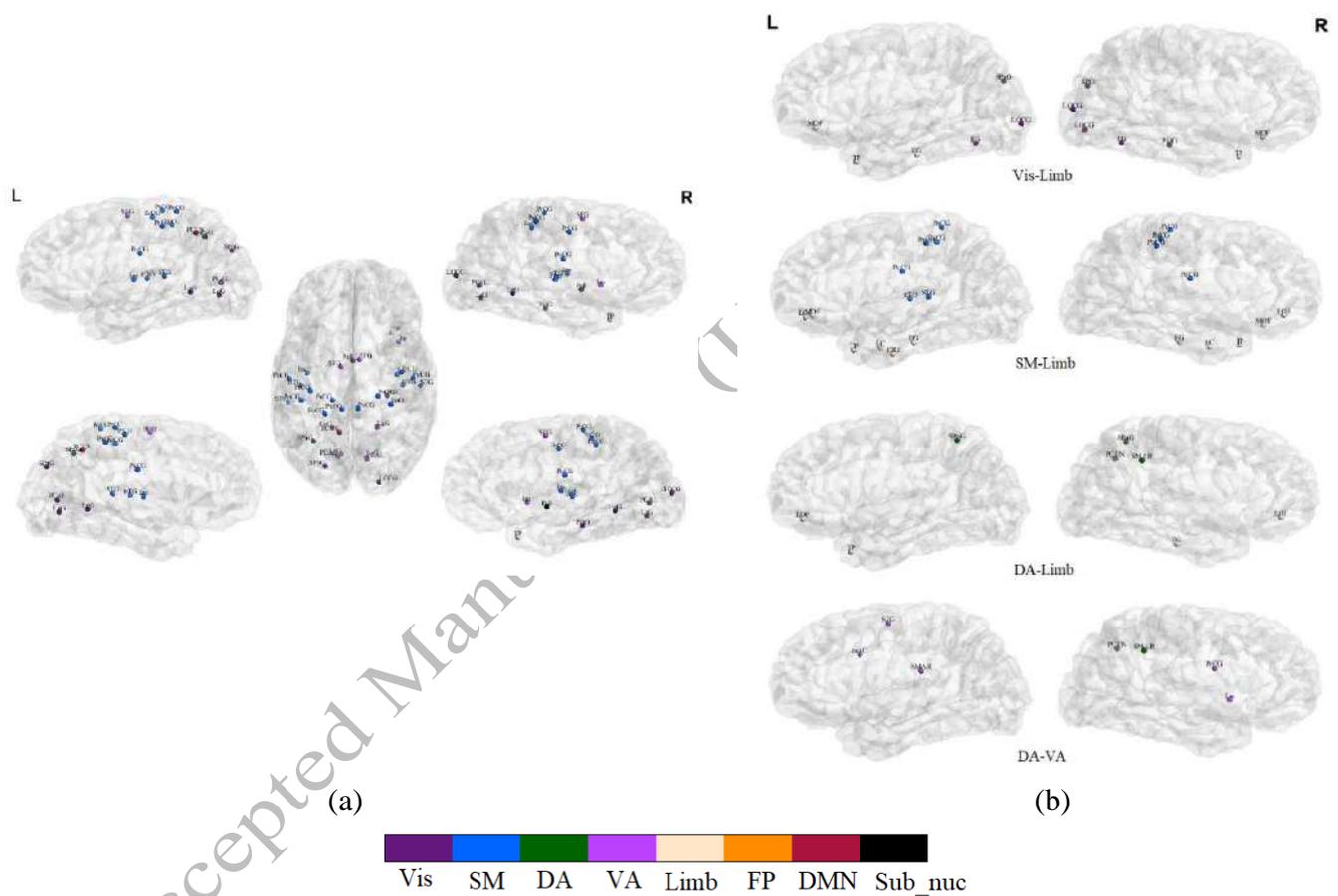


Fig. 5.

4. Discussion

In this study, we aimed to investigate to which extent schizophrenia disorder affects inter- and intra-network functional connections, as well as brain networks. Regarding the inter- and intra-network variations perspective, a higher percentage of intra-network edges (19.47 %) seemed to be changed in compare to inter-networks (13.41 %). One possible explanation for this observation would be that some basic symptoms of SZ such as motor disturbances, impaired bodily sensations, and fatigue (Larson et al., 2010; Miret et al., 2016) can be related to within-network connection impairments.

The significant intra-network alterations are allocated to the visual, somatomotor, and ventral attention networks. Prior research has confirmed that visual impairments are one of the most important manifesting features in SZ (Kogata & Iidaka, 2018; Silverstein et al., 2015; van de Ven et al., 2017). Therefore, it can be assumed that the drastic impairment of the within-network connectivity in the visual system can compromise the vision of these patients. Moreover, slow movement or useless and excessive movement is another important symptom of SZ, which can be attributed to somatomotor connectivity impairment (Adhikari et al., 2019; Kebets et al., 2019; Potvin et al., 2021; Shinn et al., 2015; Singh et al., 2014; Skåtun, Kaufmann, Doan, Alnaes, Córdova-Palomera, Jönsson, Fatouros-Bergman, Flyckt, Melle, et al., 2017; Walther & Strik, 2012). The ventral attention network is obtained as the third network experiencing intra-network alterations and it must be underlined that this network is closely related to the so-called “salience network.” The salience network is proven to be implicated in the pathophysiology of schizophrenia and its dysfunction results in the incorrect assignment of salience. This can in turn lead to the key symptoms of schizophrenia including delusions (Jimenez et al., 2016; L & PF, 2012; Smucny et al., 2016; Wynn et al., 2015).

Regarding the inter-network connectivity, the patient group demonstrated lower connectivity between the four networks. Most notably, functional connections between the

limbic system and three cortical networks were significantly affected in SZ patients. In line with this finding, several studies have reported impaired cortico-limbic functional connectivity in schizophrenia patients (Comte et al., 2018; Vai et al., 2015; Wang et al., 2021). Although the examination of between-network dysconnectivity in SZ is limited to a few studies, recent work has provided evidence for alterations in FC in between-networks for the visual- limbic (Cao et al., 2016), the somatomotor-limbic (Hummer et al., 2020) and the dorsal attention-limbic systems (Comte et al., 2018). Furthermore, a between-network dysconnectivity was found between the dorsal attention-ventral attention network, which was previously reported to be impaired in SZ patients (Hummer et al., 2020).

Abnormal connectivity between functional networks indicated impaired communication between these networks. This abnormality is potentially harming the ability to connect separate psychological and neurobiological constructs into a cohesive whole necessary for daily functioning. The interaction of the limbic system with cortical networks supports a variety of functions including emotion processing and excitation/inhibition in behavior and memory, whereas abnormal expression and regulation of emotions and impairment in the memory and cognition of patients with SZ are confirmed (J. Guo et al., 2019; Gur & Gur, 2013; Tu et al., 2013; Zhang et al., 2019). Remarkably, connections inside the limbic system were not proven to be significantly different and therefore its interaction with other networks is assumed to have a stronger effect on the symptoms of these patients.

Contrary to our findings, several researchers have reported a notable alteration (reduction or increment) of DMN and FP connections in addition to other networks in SZ patients (Galindo et al., 2017; W. Guo et al., 2017; Whitfield-Gabrieli et al., 2009). In this study, reduced connectivity within and between networks failed to survive the significant threshold. One possible reason for this discrepancy could be distinct data analyzing procedures. Data (pre-) processing steps including the selection of brain regions and network definition may cause

these differences. Here, a priori defined network based on a well-known atlas was utilized for the analysis, as opposed to data-driven approaches (such as independent component analysis (ICA)). Certainly, network definition has an undeniable impact on the number of connections and mean value of within- and between- networks. In this study, we defined nodes as distinct brain regions through automatic parcellation of each participant's brain via anatomical landmarks (FreeSurfer), as opposed to spatially uniform nodes (Baker et al., 2014) or performing a voxel-wise analysis (Gong et al., 2016).

In the assessment of the nodal strength, it is noteworthy that more significant changes occurred in the brain modular viewpoint. The strength of several nodes had not shown a remarkable difference in the whole brain evaluation (Error! Reference source not found. (b)). In other words, the total number of altered nodes in a modular viewpoint is 58 nodes, which is 21 more than the number of nodes with reduced nodal strength throughout the whole brain. This suggests that some connections may compensate for the effect of other edges' weight reduction. However, further research is required to examine this hypothesis.

4.1 Limitations

Some of the limitations of this study should be mentioned. In the current study, a dataset was used retrospectively, which included a rather low number of individuals. This may affect the precision of the obtained results. Therefore, a larger sample size is needed to confirm our findings in the future. Moreover, all subcortical regions were determined as one section while the cortex was parcellated into functionally defined regions. Consequently, a finer parcellation obtained with a higher image resolution might capture cortical-subcortical associations better.

5. Conclusion

In conclusion, it was found that SZ has a stronger effect on within-network connectivity than on between network connectivity. In a modular viewpoint of the intra-network assessment, the visual, somatomotor, and ventral attention regions demonstrated a significant reduction in

the patient group. Furthermore, the main alterations of inter-networks were attributed to interactions between the limbic system and three other cortical networks which highlighted the importance of the limbic system implicated in daily life activities. In addition, a higher number of regions with reduced nodal strength were discovered through the network viewpoint. This may illustrate the brain's attempt to compensate FC reductions and represents a topic of further research. Future studies could further investigate other graph-theoretic parameters such as e.g., the efficacy, and path length in each subnetwork.

Ethical Considerations

Compliance with ethical guidelines

All ethical principles were considered in this article.

Funding

No funding.

Conflict of interest statement

The authors declared no conflict of interest in this study.

Accepted Manuscript (Uncorrected Proof)

References

- Adhikari, B. M., Hong, L. E., Sampath, H., Chiappelli, J., Jahanshad, N., Thompson, P. M., Rowland, L. M., Calhoun, V. D., Du, X., Chen, S., & Kochunov, P. (2019). Functional network connectivity impairments and core cognitive deficits in schizophrenia. *Human Brain Mapping, 40*(16), 4593–4605. <https://doi.org/10.1002/HBM.24723>
- Algunaid, R. F., Algumaei, A. H., Rushdi, M. A., & Yassine, I. A. (2018). Schizophrenic patient identification using graph-theoretic features of resting-state fMRI data. *Biomedical Signal Processing and Control, 43*, 289–299. <https://doi.org/10.1016/j.bspc.2018.02.018>
- Andreasen, N. C., Paradiso, S., & O’Leary, D. S. (1998). “Cognitive Dysmetria” as an Integrative Theory of Schizophrenia: A Dysfunction in Cortical-Subcortical-Cerebellar Circuitry? *Schizophrenia Bulletin, 24*(2), 203–218. <https://doi.org/10.1093/OXFORDJOURNALS.SCHBUL.A033321>
- Andreasen, N. C., Pressler, M., Nopoulos, P., Miller, D., & Ho, B. C. (2010). Antipsychotic Dose Equivalents and Dose-Years: A Standardized Method for Comparing Exposure to Different Drugs. *Biological Psychiatry, 67*(3), 255–262. <https://doi.org/10.1016/J.BIOPSYCH.2009.08.040>
- Baker, J. T., Holmes, A. J., Masters, G. A., Yeo, B. T. T., Krienen, F., Buckner, R. L., & Öngür, D. (2014). Disruption of Cortical Association Networks in Schizophrenia and Psychotic Bipolar Disorder. *JAMA Psychiatry, 71*(2), 109–118. <https://doi.org/10.1001/JAMAPSYCHIATRY.2013.3469>
- Cabral, J., Kringelbach, M. L., & Deco, G. (2012). Functional graph alterations in schizophrenia: a result from a global anatomic decoupling? *Pharmacopsychiatry, 45 Suppl 1*(May). <https://doi.org/10.1055/s-0032-1309001>
- Cao, H., Bertolino, A., Walter, H., Schneider, M., Schäfer, A., Taurisano, P., Blasi, G., Haddad, L., Grimm, O., Otto, K., Dixon, L., Erk, S., Mohnke, S., Heinz, A., Romanczuk-Seiferth, N., Mühleisen, T. W., Mattheisen, M., Witt, S. H., Cichon, S., ... Meyer-Lindenberg, A. (2016). Altered Functional Subnetwork During Emotional Face Processing: A Potential Intermediate

- Phenotype for Schizophrenia. *JAMA Psychiatry*, 73(6), 598–605.
<https://doi.org/10.1001/JAMAPSYCHIATRY.2016.0161>
- Chan, M. Y., Park, D. C., Savalia, N. K., Petersen, S. E., & Wig, G. S. (2014). Decreased segregation of brain systems across the healthy adult lifespan. *Proceedings of the National Academy of Sciences*, 111(46), E4997–E5006. <https://doi.org/10.1073/PNAS.1415122111>
- Comte, M., Zendjidian, X. Y., Coull, J. T., Cancel, A., Boutet, C., Schneider, F. C., Sage, T., Lazerges, P.-E., Jaafari, N., Ibrahim, E. C., Azorin, J.-M., Blin, O., & Fakra, E. (2018). Impaired cortico- limbic functional connectivity in schizophrenia patients during emotion processing. *Social Cognitive and Affective Neuroscience*, 13(4), 381. <https://doi.org/10.1093/SCAN/NSX083>
- Correa, N. M. N., Li, Y.-O. Y., Adali, T., & Calhoun, V. D. (2009). Fusion of fMRI, sMRI, and EEG data using canonical correlation analysis. *Acoustics, Speech and Signal Processing, 2009. ICASSP 2009. IEEE International Conference On*, 385–388. <https://doi.org/10.1109/ICASSP.2009.4959601>
- Desikan, R. S., Ségonne, F., Fischl, B., Quinn, B. T., Dickerson, B. C., Blacker, D., Buckner, R. L., Dale, A. M., Maguire, R. P., Hyman, B. T., Albert, M. S., & Killiany, R. J. (2006). An automated labeling system for subdividing the human cerebral cortex on MRI scans into gyral based regions of interest. *NeuroImage*, 31(3), 968–980. <https://doi.org/10.1016/j.neuroimage.2006.01.021>
- Dietsche, B., Kircher, T., & Falkenberg, I. (2017). Structural brain changes in schizophrenia at different stages of the illness: A selective review of longitudinal magnetic resonance imaging studies. *Australian and New Zealand Journal of Psychiatry*, 51(5), 500–508. <https://doi.org/10.1177/0004867417699473>
- Fornito, A., Zalesky, A., Pantelis, C., & Bullmore, E. T. (2012). Schizophrenia, neuroimaging and connectomics. *NeuroImage*, 62(4), 2296–2314. <https://doi.org/10.1016/j.neuroimage.2011.12.090>
- Friston, K. J., & Frith, C. D. (1995). Schizophrenia: a disconnection syndrome. *Clin Neurosci*, 3(2), 89–97.

- Galindo, L., Bergé, D., Murray, G. K., Mané, A., Bulbena, A., Pérez, V., & Vilarroya, O. (2017). Default Mode Network Aberrant Connectivity Associated with Neurological Soft Signs in Schizophrenia Patients and Unaffected Relatives. *Frontiers in Psychiatry*, 8(JAN), 8. <https://doi.org/10.3389/FPSYT.2017.00298>
- Gong, Q., Hu, X., Pettersson-Yeo, W., Xu, X., Lui, S., Crossley, N., Wu, M., Zhu, H., & Mechelli, A. (2016). Network-Level Dysconnectivity in Drug-Naïve First-Episode Psychosis: Dissociating Transdiagnostic and Diagnosis-Specific Alterations. *Neuropsychopharmacology* 2017 42:4, 42(4), 933–940. <https://doi.org/10.1038/npp.2016.247>
- Guo, J., Ragland, J., & Carter, C. (2019). Memory and Cognition in Schizophrenia. *Molecular Psychiatry*, 24(5), 633. <https://doi.org/10.1038/S41380-018-0231-1>
- Guo, W., Liu, F., Chen, J., Wu, R., Li, L., Zhang, Z., Chen, H., & Zhao, J. (2017). Hyperactivity of the default-mode network in first-episode, drug-naïve schizophrenia at rest revealed by family-based case–control and traditional case–control designs. *Medicine*, 96(13). <https://doi.org/10.1097/MD.0000000000006223>
- Gur, R. C., & Gur, R. E. (2013). Memory in health and in schizophrenia. *Dialogues in Clinical Neuroscience*, 15(4), 399. <https://doi.org/10.31887/DCNS.2013.15.4/RGUR>
- Gutiérrez-Gómez, L., Vohryzek, J., Chiêm, B., Baumann, P. S., Conus, P., Cuenod, K. Do, Hagmann, P., & Delvenne, J. C. (2020). Stable biomarker identification for predicting schizophrenia in the human connectome. *NeuroImage: Clinical*, 27, 102316. <https://doi.org/10.1016/j.nicl.2020.102316>
- He, H., Sui, J., Yu, Q., Turner, J. A., Ho, B.-C., Sponheim, S. R., Manoach, D. S., Clark, V. P., & Calhoun, V. D. (2012). Altered small-world brain networks in schizophrenia patients during working memory performance. *PloS One*, 7(6), e38195. <https://doi.org/10.1371/journal.pone.0038195>
- Hummer, T. A., Yung, M. G., Goñi, J., Conroy, S. K., Francis, M. M., Mehdiyoun, N. F., & Breier, A. (2020). Functional network connectivity in early-stage schizophrenia. *Schizophrenia Research*,

218(xxxx), 107–115. <https://doi.org/10.1016/j.schres.2020.01.023>

- Jimenez, A. M., Lee, J., Wynn, J. K., Cohen, M. S., Engel, S. A., Glahn, D. C., Nuechterlein, K. H., Reavis, E. A., & Green, M. F. (2016). Abnormal ventral and dorsal attention network activity during single and dual target detection in schizophrenia. *Frontiers in Psychology*, 7(MAR), 1–11. <https://doi.org/10.3389/fpsyg.2016.00323>
- Karbasforoushan, H., & Woodward, N. D. (2013). Resting-State Networks in Schizophrenia. *Current Topics in Medicinal Chemistry*, 12(21), 2404–2414. <https://doi.org/10.2174/1568026611212210011>
- Karlsgodt, K. H., Sun, D., & Cannon, T. D. (2010). Structural and functional brain abnormalities in schizophrenia. In *Current Directions in Psychological Science* (Vol. 19, Issue 4, pp. 226–231). <https://doi.org/10.1177/0963721410377601>
- Kebets, V., Holmes, A. J., Orban, C., Tang, S., Li, J., Sun, N., Kong, R., Poldrack, R. A., & Yeo, B. T. T. (2019). Somatosensory-Motor Dysconnectivity Spans Multiple Transdiagnostic Dimensions of Psychopathology. *Biological Psychiatry*, 86(10), 779–791. <https://doi.org/10.1016/J.BIOPSYCH.2019.06.013>
- Keyvanfard, F., Nasiraei-Moghaddam, A., & Hagmann, P. (2020). Interindividual Covariations of Brain Functional and Structural Connectivities Are Decomposed Blindly to Subnetworks: A Fusion-Based Approach. *Journal of Magnetic Resonance Imaging*, 51(6), 1779–1788. <https://doi.org/10.1002/jmri.26988>
- Kogata, T., & Iidaka, T. (2018). A review of impaired visual processing and the daily visual world in patients with schizophrenia. *Nagoya Journal of Medical Science*, 80(3), 317. <https://doi.org/10.18999/NAGJMS.80.3.317>
- L, P., & PF, L. (2012). Does the salience network play a cardinal role in psychosis? An emerging hypothesis of insular dysfunction. *Journal of Psychiatry & Neuroscience: JPN*, 37(1), 17–27. <https://doi.org/10.1503/JPN.100176>
- Larson, M. K., Walker, E. F., & Compton, M. T. (2010). Early signs, diagnosis and therapeutics of the

- prodromal phase of schizophrenia and related psychotic disorders. *Expert Review of Neurotherapeutics*, 10(8), 1347–1359. <https://doi.org/10.1586/ern.10.93>
- Lawrie, S. M., Buechel, C., Whalley, H. C., Frith, C. D., Friston, K. J., & Johnstone, E. C. (2002). Reduced frontotemporal functional connectivity in schizophrenia associated with auditory hallucinations. *Biological Psychiatry*, 51(12), 1008–1011. [https://doi.org/10.1016/S0006-3223\(02\)01316-1](https://doi.org/10.1016/S0006-3223(02)01316-1)
- Lawrie, S. M., McIntosh, A. M., Hall, J., Owens, D. G. C., & Johnstone, E. C. (2008). Brain structure and function changes during the development of schizophrenia: The evidence from studies of subjects at increased genetic risk. *Schizophrenia Bulletin*, 34(2), 330–340. <https://doi.org/10.1093/schbul/sbm158>
- Li, S., Hu, N., Zhang, W., Tao, B., Dai, J., Gong, Y., Tan, Y., Cai, D., & Lui, S. (2019). Dysconnectivity of multiple brain networks in schizophrenia: A meta-analysis of resting-state functional connectivity. *Frontiers in Psychiatry*, 10(JULY), 1–11. <https://doi.org/10.3389/fpsy.2019.00482>
- Liu, J., Pearlson, G., Windemuth, A., Ruano, G., Perrone-Bizzozero, N. I., & Calhoun, V. (2009). Combining fMRI and SNP data to investigate connections between brain function and genetics using parallel ICA. *Human Brain Mapping*, 30(1), 241–255. <https://doi.org/10.1002/HBM.20508>
- Liu, Y., Liang, M., Zhou, Y., He, Y., Hao, Y., Song, M., Yu, C., Liu, H., Liu, Z., & Jiang, T. (2008). Disrupted small-world networks in schizophrenia. *Brain*, 131(4), 945–961. <https://doi.org/10.1093/brain/awn018>
- Lynall, M. E., Bassett, D. S., Kerwin, R., McKenna, P. J., Kitzbichler, M., Muller, U., & Bullmore, E. (2010). Functional connectivity and brain networks in schizophrenia. *Journal of Neuroscience*, 30(28), 9477–9487. <https://doi.org/10.1523/JNEUROSCI.0333-10.2010>
- Mamah, D., Barch, D. M., & Repovš, G. (2013). Resting state functional connectivity of five neural networks in bipolar disorder and schizophrenia. *Journal of Affective Disorders*, 150(2), 601–609. <https://doi.org/10.1016/j.jad.2013.01.051>
- Manoach, D. S., Gollub, R. L., Benson, E. S., Searl, M. M., Goff, D. C., Halpern, E., Saper, C. B., &

- Rauch, S. L. (2000). Schizophrenic subjects show aberrant fMRI activation of dorsolateral prefrontal cortex and basal ganglia during working memory performance. *Biological Psychiatry*, 48(2), 99–109. [https://doi.org/10.1016/S0006-3223\(00\)00227-4](https://doi.org/10.1016/S0006-3223(00)00227-4)
- Manoliu, A., Riedl, V., Zherdin, A., Mühlau, M., Schwerthöffer, D., Scherr, M., Peters, H., Zimmer, C., Förstl, H., Bäuml, J., Wohlschläger, A. M., & Sorg, C. (2014). Aberrant dependence of default mode/central executive network interactions on anterior insular salience network activity in schizophrenia. *Schizophrenia Bulletin*, 40(2), 428–437. <https://doi.org/10.1093/schbul/sbt037>
- Micheloyannis, S. (2012). Graph-based network analysis in schizophrenia. *World Journal of Psychiatry*, 2(1). <https://doi.org/10.5498/wjp.v2.i1.EDITORIAL>
- Miret, S., Fatjó-Vilas, M., Peralta, V., & Fañanás, L. (2016). Basic symptoms in schizophrenia, their clinical study and relevance in research. *Revista de Psiquiatria y Salud Mental*, 9(2), 111–122. <https://doi.org/10.1016/j.rpsm.2015.10.007>
- Potvin, S., Giguère, C. É., & Mendrek, A. (2021). Functional connectivity during visuospatial processing in schizophrenia: A classification study using lasso regression. *Neuropsychiatric Disease and Treatment*, 17, 1077–1087. <https://doi.org/10.2147/NDT.S304434>
- Rubinov, M., & Bullmore, E. (2013). Schizophrenia and abnormal brain network hubs. *Dialogues in Clinical Neuroscience*, 15(3), 339–349. <https://doi.org/10.1016/j.siny.2015.10.004>
- Rubinov, M., & Sporns, O. (2010). Complex network measures of brain connectivity: Uses and interpretations. *NeuroImage*, 52(3), 1059–1069. <https://doi.org/10.1016/j.neuroimage.2009.10.003>
- Shinn, A. K., Baker, J. T., Lewandowski, K. E., Öngür, D., & Cohen, B. M. (2015). Aberrant cerebellar connectivity in motor and association networks in schizophrenia. *Frontiers in Human Neuroscience*, 9(MAR), 134. <https://doi.org/10.3389/FNHUM.2015.00134>
- Silverstein, S., Keane, B. P., Blake, R., Giersch, A., Green, M., & Kéri, S. (2015). Vision in schizophrenia: why it matters. *Frontiers in Psychology*, 6(FEB). <https://doi.org/10.3389/FPSYG.2015.00041>

- Singh, S., Goyal, S., Modi, S., Kumar, P., Singh, N., Bhatia, T., Deshpande, S. N., & Khushu, S. (2014). Motor function deficits in schizophrenia: An fMRI and VBM study. *Neuroradiology*, *56*(5), 413–422. <https://doi.org/10.1007/s00234-014-1325-3>
- Skåtun, K. C., Kaufmann, T., Doan, N. T., Alnæs, D., Córdova-Palomera, A., Jönsson, E. G., Fatouros-Bergman, H., Flyckt, L., KaSP, Melle, I., Andreassen, O. A., Agartz, I., & Westlye, L. T. (2017). Consistent Functional Connectivity Alterations in Schizophrenia Spectrum Disorder: A Multisite Study. *Schizophrenia Bulletin*, *43*(4), 914–924. <https://doi.org/10.1093/SCHBUL/SBW145>
- Skåtun, K. C., Kaufmann, T., Doan, N. T., Alnæs, D., Córdova-Palomera, A., Jönsson, E. G., Fatouros-Bergman, H., Flyckt, L., Melle, I., Andreassen, O. A., Agartz, I., & Westlye, L. T. (2017). Consistent Functional Connectivity Alterations in Schizophrenia Spectrum Disorder: A Multisite Study. *Schizophrenia Bulletin*, *43*(4), 914–924. <https://doi.org/10.1093/schbul/sbw145>
- Skudlarski, P., Jagannathan, K., Anderson, K., Stevens, M. C., Calhoun, V. D., Skudlarska, B. A., & Pearlson, G. (2010). Brain connectivity is not only lower but different in schizophrenia: a combined anatomical and functional approach. *Biological Psychiatry*, *68*(1), 61–69.
- Smucny, J., Olincy, A., & Tregellas, J. R. (2016). Nicotine restores functional connectivity of the ventral attention network in schizophrenia. *Neuropharmacology*, *108*, 144–151. <https://doi.org/10.1016/J.NEUROPHARM.2016.04.015>
- Sporns, O. (2013). Structure and function of complex brain networks. *Dialogues in Clinical Neuroscience*, *15*(3), 247–262. <https://doi.org/10.1137/S003614450342480>
- Sporns, O., & Betzel, R. F. (2016). Modular brain networks. *Annual Review of Psychology*, *67*, 613–640. <https://doi.org/10.1146/annurev-psych-122414-033634>.Modular
- Stahl, S. M. (2018). Beyond the dopamine hypothesis of schizophrenia to three neural networks of psychosis: dopamine, serotonin, and glutamate. *CNS Spectrums*, *23*(3), 187–191. <https://doi.org/10.1017/S1092852918001013>
- Stephan, K. E., Friston, K. J., & Frith, C. D. (2009). Dysconnection in Schizophrenia: From Abnormal Synaptic Plasticity to Failures of Self-monitoring. *Schizophrenia Bulletin*, *35*(3), 509–527.

<https://doi.org/10.1093/SCHBUL/SBN176>

- Stephen, J. M., Coffman, B. a., Jung, R. E., Bustillo, J. R., Aine, C. J., & Calhoun, V. D. (2013). Using joint ICA to link function and structure using MEG and DTI in schizophrenia. *NeuroImage*, *83*, 418–430. <https://doi.org/10.1016/j.neuroimage.2013.06.038>
- Tu, P. C., Lee, Y. C., Chen, Y. S., Li, C. T., & Su, T. P. (2013). Schizophrenia and the brain's control network: Aberrant within- and between-network connectivity of the frontoparietal network in schizophrenia. *Schizophrenia Research*, *147*(2–3), 339–347. <https://doi.org/10.1016/j.schres.2013.04.011>
- Vai, B., Papa, G. S., Poletti, S., Radaelli, D., Donnici, E., Bollettini, I., Falini, A., Cavallaro, R., Smeraldi, E., & Benedetti, F. (2015). Abnormal cortico-limbic connectivity during emotional processing correlates with symptom severity in schizophrenia. *European Psychiatry*, *30*(5), 590–597. <https://doi.org/10.1016/J.EURPSY.2015.01.002>
- van de Ven, V., Rotarska Jagiela, A., Oertel-Knöchel, V., & Linden, D. E. J. (2017). Reduced intrinsic visual cortical connectivity is associated with impaired perceptual closure in schizophrenia. *NeuroImage: Clinical*, *15*, 45–52. <https://doi.org/10.1016/J.NICL.2017.04.012>
- Van Den Heuvel, M. P., & Fornito, A. (2014). Brain networks in schizophrenia. In *Neuropsychology Review* (Vol. 24, Issue 1, pp. 32–48). Springer. <https://doi.org/10.1007/s11065-014-9248-7>
- van den Heuvel, M. P., & Hulshoff Pol, H. E. (2010). Exploring the brain network: A review on resting-state fMRI functional connectivity. *European Neuropsychopharmacology*, *20*(8), 519–534. <https://doi.org/10.1016/j.euroneuro.2010.03.008>
- Vohryzek, J., Aleman-Gomez, Y., Griffa, A., Raoul, J., Cleusix, M., Baumann, P. S., Conus, P., Cuenod, K. Do, & Hagmann, P. (2020). *Structural and functional connectomes from 27 schizophrenic patients and 27 matched healthy adults*. <https://doi.org/10.5281/ZENODO.3758534>
- Walther, S., & Strik, W. (2012). Motor Symptoms and Schizophrenia. *Neuropsychobiology*, *66*(2), 77–92. <https://doi.org/10.1159/000339456>

- Wang, X., Yin, Z., Sun, Q., Jiang, X., Chao, L., Dai, X., & Tang, Y. (2021). Comparative Study on the Functional Connectivity of Amygdala and Hippocampal Neural Circuits in Patients With First-Episode Schizophrenia and Other High-Risk Populations. *Frontiers in Psychiatry, 0*, 1444. <https://doi.org/10.3389/FPSYT.2021.627198>
- Wei, Y., Chang, M., Womer, F. Y., Zhou, Q., Yin, Z., Wei, S., Zhou, Y., Jiang, X., Yao, X., Duan, J., Xu, K., Zuo, X. N., Tang, Y., & Wang, F. (2018). Local functional connectivity alterations in schizophrenia, bipolar disorder, and major depressive disorder. *Journal of Affective Disorders, 236*, 266–273. <https://doi.org/10.1016/j.jad.2018.04.069>
- Whitfield-Gabrieli, S., Thermenos, H. W., Milanovic, S., Tsuang, M. T., Faraone, S. V., McCarley, R. W., Shenton, M. E., Green, A. I., Nieto-Castanon, A., LaViolette, P., Wojcik, J., Gabrieli, J. D. E., & Seidman, L. J. (2009). Hyperactivity and hyperconnectivity of the default network in schizophrenia and in first-degree relatives of persons with schizophrenia. *Proceedings of the National Academy of Sciences of the United States of America, 106*(4), 1279–1284. <https://doi.org/10.1073/pnas.0809141106>
- Wu, X. jie, Zeng, L. L., Shen, H., Yuan, L., Qin, J., Zhang, P., & Hu, D. (2017). Functional network connectivity alterations in schizophrenia and depression. *Psychiatry Research - Neuroimaging, 263*, 113–120. <https://doi.org/10.1016/j.psychres.2017.03.012>
- Wynn, J. K., Jimenez, A. M., Roach, B. J., Korb, A., Lee, J., Horan, W. P., Ford, J. M., & Green, M. F. (2015). Impaired target detection in schizophrenia and the ventral attentional network: Findings from a joint event-related potential-functional MRI analysis: Target stimulus ERP/fMRI analysis in schizophrenia. *NeuroImage: Clinical, 9*, 95–102. <https://doi.org/10.1016/j.nicl.2015.07.004>
- Yang, A. C., & Tsai, S. J. (2017). New Targets for Schizophrenia Treatment beyond the Dopamine Hypothesis. *International Journal of Molecular Sciences, 18*(8). <https://doi.org/10.3390/IJMS18081689>
- Yeo, B. T. T., Krienen, F. M., Sepulcre, J., Sabuncu, M. R., Lashkari, D., Hollinshead, M., Roffman, J. L., Smoller, J. W., Zollei, L., Polimeni, J. R., Fischl, B., Liu, H., & Buckner, R. L. (2011). The

organization of the human cerebral cortex estimated by intrinsic functional connectivity. *Journal of Neurophysiology*, *106*, 1125–1165. <https://doi.org/10.1152/jn.00338.2011>.

Yu, Q., Sui, J., Rachakonda, S., He, H., Pearlson, G., & Calhoun, V. D. (2011). Altered small-world brain networks in temporal lobe in patients with schizophrenia performing an auditory oddball task. *Frontiers in Systems Neuroscience*, *5*(FEBRUARY 2011), 1–13. <https://doi.org/10.3389/fnsys.2011.00007>

Zhang, Y., Kuhn, S. K., Jobson, L., & Haque, S. (2019). A review of autobiographical memory studies on patients with schizophrenia spectrum disorders. *BMC Psychiatry*, *19*(1), 1–36. <https://doi.org/10.1186/s12888-019-2346-6>

Accepted Manuscript (Uncorrected Proof)