



Published in final edited form as:

Mol Psychiatry. 2020 October ; 25(10): 2431–2440. doi:10.1038/s41380-018-0288-x.

Functional Connectome Organization Predicts Conversion to Psychosis in Clinical High-Risk Youth from the SHARP Program

Guusje Collin, MD, PhD^{1,2,3}, Larry J. Seidman, PhD^{1,†}, Matcheri S. Keshavan, MD¹, William S. Stone, PhD¹, Zhenghan Qi, MD, PhD^{1,4}, Tianhong Zhang, MD, PhD⁵, Yingying Tang, PhD⁵, Huijun Li, PhD⁶, Sheeba Arnold Anteraper, PhD^{2,7}, Margaret A. Niznikiewicz, PhD⁸, Robert W. McCarley, MD^{8,†}, Martha E. Shenton, PhD^{3,9,10}, Jijun Wang, MD⁵, Susan Whitfield-Gabrieli, PhD^{2,11}

¹Department of Psychiatry, Beth Israel Deaconess Medical Center, Harvard Medical School, Boston, MA, USA

²McGovern Institute for Brain Research, Department of Brain and Cognitive Sciences, Massachusetts Institute of Technology, Cambridge, MA, USA

³Psychiatry Neuroimaging Laboratory, Department of Psychiatry, Brigham and Women's Hospital, Harvard Medical School, Boston, MA, USA

⁴Department of Linguistics and Cognitive Science, University of Delaware, Newark, DE, USA

⁵Shanghai Key Laboratory of Psychotic Disorders, Shanghai Mental Health Center, Shanghai Jiao Tong University School of Medicine, Shanghai, China

⁶Florida A&M University, Department of Psychology, Tallahassee, FL, USA

⁷Alan and Lorraine Bressler Clinical and Research Program for Autism Spectrum Disorder, Massachusetts General Hospital, Boston MA, USA

⁸Department of Psychiatry, VA Boston Healthcare System, Brockton Division, Brockton, MA, USA

⁹Research and Development, VA Boston Healthcare System, Brockton Division, Brockton, MA, USA

¹⁰Department of Radiology, Brigham and Women's Hospital, Harvard Medical School, Boston, MA, USA

¹¹Poitras Center for Affective Disorders, Department of Brain and Cognitive Sciences, Massachusetts Institute of Technology, Cambridge, MA, USA

Users may view, print, copy, and download text and data-mine the content in such documents, for the purposes of academic research, subject always to the full Conditions of use:http://www.nature.com/authors/editorial_policies/license.html#terms

Correspondence to: Guusje Collin, Beth Israel Deaconess Medical Center, Department of Psychiatry, Massachusetts Mental Health Center, 75 Fenwood Road, Boston, MA 02115, USA; Phone: +1 (617) 626 9300; gcollin@bidmc.harvard.edu. Jijun Wang, Shanghai Key Laboratory of Psychotic Disorders, Shanghai Mental Health Center, Shanghai Jiao Tong University School of Medicine, 600 Wanping Nan Road, Shanghai, 200030, China. Phone: +86 (21) 3477 3065. jijunwang27@163.com.

Conflict of interest

The authors declare no conflict of interest.

Supplementary Information accompanies the paper on the Molecular Psychiatry website (<http://www.nature.com/mp>)

† Dr. Larry Seidman passed away on September 7, 2017 and Dr. Robert McCarley passed away on May 27, 2017. Professors Seidman and McCarley were two of the initiators and principal investigators of the Shanghai At Risk for Psychosis (SHARP) study.

Abstract

The emergence of prodromal symptoms of schizophrenia and their evolution into overt psychosis may stem from an aberrant functional reorganization of the brain during adolescence. To examine whether abnormalities in connectome organization precede psychosis onset, we performed a functional connectome analysis in a large cohort of medication-naïve youth at risk for psychosis from the Shanghai At Risk for Psychosis (SHARP) study. The SHARP program is a longitudinal study of adolescents and young adults at Clinical High Risk (CHR) for psychosis, conducted at the Shanghai Mental Health Center in collaboration with neuroimaging laboratories at Harvard and MIT. Our study involved a total of 251 subjects, including 158 CHRs and 93 age-, sex-, and education-matched healthy controls. During one-year follow-up, 23 CHRs developed psychosis. CHRs who would go on to develop psychosis were found to show abnormal modular connectome organization at baseline, while CHR non-converters did not. In all CHRs, abnormal modular connectome organization at baseline was associated with a three-fold conversion rate. A region-specific analysis showed that brain regions implicated in early-course schizophrenia, including superior temporal gyrus and anterior cingulate cortex, were most abnormal in terms of modular assignment. Our results show that functional changes in brain network organization precede the onset of psychosis and may drive psychosis development in at-risk youth.

Introduction

Schizophrenia is a psychiatric disorder that manifests early in life and derails social, cognitive, and academic development. The development of the illness typically follows a sequential trajectory that includes a premorbid phase with subtle and nonspecific deviations from normative development,¹ a prodromal phase with sub-threshold symptoms and declining functioning,²⁻⁴ and a first psychotic episode that marks the formal onset of the illness.⁵ In recent years, the focus of schizophrenia research shifted from the first episode to earlier stages of illness development. Studies of the prodromal phase aim to elucidate the biological and environmental factors that guide the trajectory from elevated risk to established illness, in order to contribute to the development of early detection and intervention strategies for schizophrenia.⁶

The prodromal or clinical high risk (CHR) phase of schizophrenia is characterized by attenuated or transient psychotic symptoms such as unusual thought content, suspiciousness, or mild perceptual abnormalities that typically manifest in adolescence or early adulthood.^{2,4} The CHR syndrome has a large heterogeneity in clinical outcome ranging from complete remission to full-blown psychosis.^{7,8} It has been suggested that inter-individual differences in brain circuitry may underlie the differences in outcome for high-risk individuals.⁹ Indeed, recent studies suggest that abnormalities in functional brain connectivity and organization may differentiate at-risk individuals who will develop psychosis from those who do not transition.^{9,10} These studies may help to elucidate the neurobiological events that precipitate and possibly drive the manifestation of psychotic symptoms.

The typical timing of the CHR syndrome in middle to late adolescence coincides with a crucial phase of brain development during which psychosocial factors interact with genetically mediated brain changes to reshape the brain's functional organization. The brain is organized into a collection of functional networks that form identifiable modules in the brain's network.¹¹ This modular organization is thought to allow specialized circuits to focus on specific tasks by limiting the interference of regions processing different types of neural information.^{12,13} Neuroimaging studies indicate that a considerable reorganization of the brain's functional modules takes place between late childhood and early adulthood.^{11,14,15} We hypothesize that the modular reorganization of the brain during this developmental window may go awry in at-risk youth, resulting in aberrant connectivity patterns that may contribute to the development of psychotic symptoms.¹⁶

To examine modular brain organization in at-risk youth, we draw from the field of connectomics, an emerging branch of neuroscience that uses graph theory to examine the brain's connectivity network known as the connectome.¹⁷ By assessing the modular organization of the functional connectome in a large sample of adolescents and young adults at risk for psychosis, we aim to determine whether abnormalities in modular connectome organization exist before the onset of psychosis and predispose to psychotic convergence.

Materials and methods

Participants

This study involved a total of 251 participants, including 158 Clinical High Risk (CHR) subjects and 93 Healthy Controls (HC) matched to CHRs based on age, sex, and education in years. The large majority of CHRs was naïve to psychotropic medication at baseline clinical assessment (>95%) and neuroimaging (>80%). Participants were recruited at the Shanghai Mental Health Center (SMHC), as part of the Shanghai At Risk for Psychosis (SHARP) program.¹⁸ This NIMH-funded study is a collaboration between SMHC, Harvard Medical School at Beth Israel Deaconess Medical Center (BIDMC) and Brigham and Women's Hospital, and Massachusetts Institute of Technology (study details including power analysis and in/exclusion criteria in Supplementary Information 1.1). The study was approved by Institutional Review Boards of BIDMC and SMHC. All subjects or their legal guardians gave written informed consent. Table 1 provides demographic and clinical characteristics of all participants.

Clinical and cognitive assessment

Prodromal symptoms were assessed using a validated Chinese version¹⁹ of the Structured Interview for Prodromal Symptoms (SIPS).²⁰ Total IQ was estimated using the Wechsler Abbreviated Scale of Intelligence (WASI).²¹

Conversion criteria

During a mean (sd) follow-up of 392 (77) days, 23 CHRs developed psychosis (CHR+), while 135 did not (CHR-). Conversion to psychosis was determined using the SIPS operational definition of psychosis onset,²² with at least one psychotic level symptom (rated "6" on the SIPS positive scale) with either sufficient frequency or duration. For CHR+, the

date of conversion was recorded and time to psychosis computed as the number of days between study inclusion and psychosis onset.

Image acquisition

Magnetic Resonance Imaging (MRI) scans were acquired on a 3T Siemens MR B17 (Verio) system, 32-channel head coil at the SMHC and included an anatomical T1-weighted MRI scan (MP-RAGE; TR=2300 ms, TE=2.96 ms, FA=9 degree, FOV=256mm, voxel size: 1×1×1mm, 192 contiguous sagittal slices, duration 9'14'') and resting-state fMRI (rs-fMRI) scan (149 functional volumes; TR=2500 ms, TE=30 ms, FA=90 degree, FOV=224mm, voxel size: 3.5×3.5×3.5mm, 37 contiguous axial slices, duration 6'19'').

Image preprocessing

Figure 1 provides an overview of all analysis steps. As connectivity and connectome metrics are sensitive to methodological factors including image preprocessing and node selection,^{23,24} two processing streams were used to process our data to ensure that results were not specific to any one methodology (Figure 1A): a *surface-based* method, the results of which are presented as primary findings, and an *MNI-based* method used to verify primary findings.

Image processing is described in detail in Supplementary Information 1.2. Briefly, FreeSurfer (v6.0)²⁵ and CONN (v17d)²⁶ software were used to preprocess T1 and rs-fMRI data. For *surface-based* processing, a total of 162 subject-specific ROIs derived from FreeSurfer were used as nodes, including 148 regions comprising the Destrieux Atlas and 14 subcortical structures (bilateral thalamus, hippocampus, amygdala, nucleus accumbens, caudate nucleus, putamen and global pallidus). For *MNI-based* processing, T1 and rs-fMRI scans were normalized to MNI space and the Harvard-Oxford Atlas was used to divide the cerebrum into a total of 105 brain regions,²⁷ including 91 cortical brain regions and the same 14 subcortical structures. Rigorous motion correction and artifact removal were performed to deal with spurious correlations.²⁶ There were no significant group-differences in motion parameters or the number of rejected fMRI volumes (Supplementary Information 1.2).

Functional connectome reconstruction

A functional connectome was constructed for each participant, consisting of 162 subject-specific (or 105 atlas-based) nodes representing the aforementioned brain regions. The level of functional connectivity between each node pair was computed as the normalized z-score of the Pearson's correlation between the noise-corrected timeseries of each pair of brain regions and stored in a functional connectivity matrix (Figure 1B).

Modular organization of the functional connectome

Various methods exist to examine the modular organization of complex networks.¹³ Here, we use the Louvain community detection method (<https://sites.google.com/site/bctnet/>), which partitions a network so as to maximize metric Q, representing the strength of edges inside communities relative to edges between communities.²⁸ The method is suitable for functional connectome analysis as it can take both positive and negative edge weights (i.e.,

connectivity estimates) into account without requiring an arbitrary connectivity threshold.
13,28,29

Group-networks.

As a first step to assess the modular organization of the functional connectome, one group-averaged functional network was constructed for HC, CHR-, and CHR+ groups. Each group-network's modular organization was assessed using the Louvain method. As the algorithm searches for high modularity partitions in a heuristic fashion, resulting partitions differ slightly from run to run.^{13,29,30} Therefore, the algorithm was run 10,000 times for each group-network and the partition associated with the highest level of Q selected (Figure 1C). A consensus similarity method³¹ was used as an alternative method to select modular partitions for both group and individual networks (Supplementary Information 1.3).

Individual subjects.

Second, the Louvain algorithm was applied to individual connectome reconstructions. To assess how similar the modular organization of each individual network was to an average healthy network, network partitions of individual subjects were compared to the group-averaged HC network using the Rand similarity coefficient (S_R),³² providing an intuitive measure (between 0 and 1) of the similarity between two partitions (details in Supplementary Information 1.3). Network resolution parameter γ was set to 1.5, in order to identify more fine-grained modules while not overinflating the total number of modules, as this would hamper comparisons of network partitions.³² In addition, modular organization was examined across a range of γ (Supplementary Information 1.3).

Region-specific alterations in modular connectome organization

As an exploratory analysis to assess which brain regions are most abnormal in terms of module assignment in CHR+ versus CHR-, individual networks were again compared to the HC network, now determining for each node i in the network the fraction of neighboring nodes with equal module assignment (details in Supplementary Information 1.3).

Code availability

All image processing and graph analyses were performed using freely available software. Version and access details are provided in Supplementary Information 1.2 and 1.3.

Statistical analysis

Group analysis.—Analysis of Covariance (ANCOVA) was used to compare S_R among subject groups. Assumptions of normality and homogeneity of variance were met (Supplementary Information 1.4). Age, sex, and the number of rejected fMRI volumes were included in the model as covariates, and group-covariate interactions were assessed. Medication status was included as a covariate of non-interest as a minority of CHRs (<20%) were on psychotropic medication by the time of scanning. Region-specific metrics were analyzed using the same ANCOVA model, applying FDR-correction ($q = .05$) to account for multiple comparisons.

Psychosis-free survival analysis.—To determine whether abnormal modular connectome organization at baseline predicted conversion to psychosis, CHRs were divided by median split into two groups with above and below-average S_R , reflecting normal and abnormal modular organization respectively. Kaplan Meier analysis was used to assess psychosis-free survival for each group. Survival functions were compared using log-rank tests. Cox regression analysis was used to assess how baseline modular organization and clinical characteristics (i.e., age, sex, IQ, SIPS symptoms, and GAF functioning) predicted time to conversion.

Results

Modular connectome organization – group-networks

Figure 2A shows the modular organization of group-averaged functional networks. Community detection in the HC network yielded five modules, largely reflecting known functional networks. Modular organization of the CHR- network was similar to HCs. The CHR+ network showed a number of qualitative differences, including a separation of orbitofrontal regions from the (para)limbic module, and with bilateral superior temporal gyrus changing assignment from the sensorimotor to the (para)limbic module. Moreover, a sixth cingulo-opercular module was observed in the CHR+ network. This module was not present in HC and CHR- group-networks at this resolution but did show up at higher levels of network resolution (Supplementary Figure 2). Using consensus similarity to identify modular partitions produced highly similar partitions (Supplementary Information 1.3).

Modular connectome organization – individual subjects

The Rand similarity coefficient, reflecting how similar the modular organization of individual networks was to the averaged healthy network, showed a significant main effect of group ($F_{(2, 245)} = 4.08, p = .018$). Post-hoc bivariate comparison indicated that modular partitions of CHR+ subjects were significantly less similar to the average healthy network than both HCs ($F_{(1, 110)} = 4.32, p = .039$) and CHR- ($F_{(1, 152)} = 7.87, p = .006$) (Figure 2B). There was no significant difference between HCs and CHR- ($F_{(1, 222)} = 0.02, p = .898$). An additional analysis to ensure that HC results were not biased by the fact that individual HCs contributed to the group-averaged HC network confirmed our findings (Supplementary Information 1.3). Repeating the analyses using the MNI-based processing method largely corroborated our findings (details in Supplementary Information 1.3 and Supplementary Figure 3 and 4). Consensus partitions were very similar to the original partitions and reanalysis of our main finding using consensus partitions produced a trend-level effect but did not change the nature of our findings (Supplementary Information 1.3). Assessing group-effects across different levels of resolution parameter γ showed that the main effect holds for a range of γ (see Supplementary Information 1.3 and Supplementary Figure 1 and 2). Of note, there were no significant group-differences in the overall level of modularity Q ($F_{(2,250)} = 1.08, p = .340$) or overall connectivity strength ($F_{(2,250)} = 1.7, p = .185$).

Region-specific alterations in modular organization

Using *surface-based* data, no regional effects survived FDR-correction. Using *MNI-based* data, the right superior temporal gyrus (STG) was the only region surviving FDR-correction

($F_{(1, 152)} = 9.20, p < .001$). As exploratory results, regional effects at uncorrected $p < .05$ from the *surface-based* and *MNI-based* analysis are summarized in Figure 3A and 3B respectively. Regions with (marginal) effects in both atlases included bilateral STG and temporal plane, and right anterior cingulate gyrus, fusiform cortex, and amygdala.

Psychosis-free survival analysis

Kaplan-Meier analysis indicated significantly different ($z = 2.41, p = .016$) psychosis-free survival functions for CHRs with typical versus atypical modular connectome organization (Figure 4), with a hazard ratio of 3.1 indicating a three-fold relative event rate (i.e. conversion to psychosis) in CHRs with atypical modular organization at baseline. Combining baseline S_R with baseline clinical characteristics in one cox regression model indicated that abnormal baseline connectome organization ($z = -2.37, p = .018$), lower IQ ($z = -2.48, p = .013$) and male sex ($z = 1.92, p = .036$) predicted shorter time to conversion. These findings were confirmed using MNI-processed data (Supplementary Information 1.5 and Supplementary Figure 4).

Discussion

This study examined functional connectome organization in a large cohort of adolescents and young adults at clinical high-risk for psychosis. Our findings suggest that abnormalities in the modular organization of the functional connectome precede the first psychotic episode. We find that baseline modular connectome organization is abnormal in CHRs who go on to develop psychosis, but not in CHRs who do not convert. Moreover, conversion to psychosis was over three times more likely in CHRs with abnormal connectome organization at baseline as compared to CHRs with typical baseline connectome organization. Functional changes in brain network organization that precede the formal onset of psychosis may be involved in the manifestation of (prodromal) psychotic symptoms.

Our findings of abnormal modular organization of the functional connectome in youth at risk for schizophrenia are supported by three previous graph analytical studies of functional connectivity data in schizophrenia patients and at-risk youth. Two studies in schizophrenia patients demonstrated a reorganization of modular brain network topology in patients with established illness.^{33,34} In addition, a recent study in 88 at-risk individuals, including 12 who later developed psychosis, identified changes in the modular organization of the functional connectome in at-risk subjects who transitioned to psychosis.⁹ Our current study confirms and extends these previous results by showing that abnormal functional organization of the connectome precedes the first psychotic episode and develops in the absence of psychotropic medication.

Two competing hypotheses have been developed on modular brain network organization in schizophrenia. The first is that the connectome is *more* modular in schizophrenia. An early version of this theory was proposed by Hoffman and McGlashan, who modeled the effects of excessive pruning in neural network simulations. They concluded that the resulting fragmentation of the brain network gives rise to functionally autonomous modules that act as 'parasitic foci' that repeatedly introduce the same output into the brain's information flow, which may underlie auditory hallucinations or delusions of control.³⁵⁻³⁷ The second

hypothesis, first articulated in a critique of Hoffman and McGlashan's work, argues that the brain's network is *less* modular in schizophrenia.³⁸ A less modular network could result in reduced information encapsulation and "overflow" of neural information from e.g., language into perceptual systems, and thereby invoke symptoms such as thought insertion and auditory hallucinations. Empirical evidence appears to favor the first theory, with a functional connectivity study showing more and smaller modules in schizophrenia patients³⁹ and two structural connectome studies showing higher modularity in schizophrenia and CHR.^{40,41} In contrast, a study showing reduced modularity of functional brain networks in childhood-onset schizophrenia is more in line with the second theory.⁴² Our current findings and previous results^{33,34} add a third possibility. Namely, that the brain is not just more or less modular, but that there is a qualitative reorganization of the brain's modular organization in schizophrenia, resulting in abnormal functional interaction patterns between a range of brain regions, which may contribute to psychotic and cognitive symptom development.

Finding abnormalities in modular connectome organization before the onset of psychosis suggests that a maladaptive reorganization in the modular topology of the functional connectome may take place in the months to years preceding the first psychotic episode. In typical brain development, the maturation of different functional systems occurs at different times in the course of development, with e.g., sensorimotor networks maturing before those mediating higher cognitive functions.⁴³ Adolescent behaviors such as impulsivity and risk taking have been attributed to the asynchronous maturation of limbic and prefrontal systems, giving rise to heightened sensitivity to motivational cues in the context of immature cognitive control.⁴⁴ This critical window of limbic and cognitive system development and high sensitivity to socio-environmental inputs may represent a window of vulnerability for youth at risk for psychosis. Any deviation from the process of modular reorganization during this developmental window could give rise to complex patterns of hypo- and hyper-connectivity between brain regions, as have been observed in schizophrenia patients.^{45–47} These aberrant functional connectivity patterns could have a particularly profound and lasting impact on the brain's functional organization and may contribute to psychotic symptom development.

While the global modular organization of the brain was the main focus of our study, we note that the brain regions showing region-wise changes in modular assignment were regions that are commonly associated with schizophrenia. Examining two distinct brain atlases, overlapping regions in terms of (marginal) group-effects included STG and temporal plane, anterior cingulate cortex, fusiform gyrus, and amygdala. These regions are among the most consistently implicated brain regions in early-course schizophrenia.^{48–52} Examining modular network organization across levels of network resolution also indicated a change in modular assignment of STG from the sensorimotor to the limbic module. Intriguingly, a separation of STG from the larger somatosensory community was recently reported in a study of modular brain organization in schizophrenia.³³ Other consistent abnormalities across resolutions included changes in the modular assignment of orbitofrontal cortex, striatum, and insula, in line with recent findings of salience module abnormalities in at-risk individuals.⁹ Moreover, the latter study reported visual areas to extend into the limbic module.⁹ Both our current and previous investigations in at-risk youth thus find primary

sensory regions to become embedded in the limbic system in prodromal psychosis. These findings may fit in with theoretical models attributing psychotic symptoms to aberrant memory activations or the attribution of erroneous salience to the internal representation of a percept or memory.⁵³ Moreover, findings in schizophrenia patients indicate that the most prominent break-up of functional modules involves sensory, auditory, and visual areas.³³ Together, these previous and our current findings suggest that an initial separation of primary auditory and visual regions followed by a more generalized fragmentation of sensory processing may underlie disease progression in prodromal psychosis.

A number of issues should, however, be taken into account when interpreting our findings. First, physiological and head motion artifacts are known to influence fMRI-derived measures of functional connectivity.^{54,55} To deal with these issues, we used the anatomical CompCor (aCompCor) method⁵⁶ for physiological noise reduction and the Artifact Detection Tool (*art*) for efficient rejection of motion and artefactual time points.²⁶ Second, a much-debated issue in the context of functional connectivity is the biological validity of negative, or anti-correlations.⁵⁷ A recent study indicates that when physiological and other noise sources are effectively removed, anti-correlations are present in the absence of global signal regression, suggesting a biological origin.⁵⁸ We therefore chose to include both positively and negatively weighted connectome edges. A third and related issue is the application of thresholds in functional network analyses. In graph theoretical studies, thresholds are commonly applied to obtain a more sparsely connected representation of the functional connectome. Even when the network is examined over a range of different thresholds, the impact of imposing a threshold on the resulting graph metrics is non-trivial. Moreover, thresholding typically removes negative correlations, thereby discarding neurobiologically relevant information.¹³ To avoid these limitations, we used a community detection method equipped to deal with fully weighted networks including both positive and negative edge weights.²⁸ Lastly, the changes in functional connectome organization observed in our study may indicate abnormalities in synchronous neural activity and modular interactions. However, given the indirect and correlational nature of functional connectivity measurements derived from resting-state fMRI, we cannot rule out the possibility that non-neural factors including hemodynamic response function variability^{59,60} may have influenced our results.

This study finds that the modular organization of the functional connectome is abnormal in CHR youth that go on to develop a psychotic episode, but not in CHRs that do not develop psychosis. In addition, we show that abnormal modular connectome organization precedes overt psychosis and is predictive of psychotic development. Our results provide new insights into functional mechanisms on the connectome level that may underlie the development of psychotic symptoms and are of key interest to efforts to identify biomarkers for transition to psychosis.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgements

This study was supported by the Ministry of Science and Technology of China (2016 YFC 1306803) and US National Institute of Mental Health (R21 MH 093294, R01 MH 101052, and R01 MH 111448). Dr. Collin was supported by an EU Marie Curie Global Fellowship (Grant no. 749201); Dr. Keshavan was supported by an NIMH Grant (R01 MH 64023); Drs. Shenton and McCarley were supported by a VA Merit Award.

References

1. Liu CH, Keshavan MS, Tronick E, Seidman LJ. Perinatal risks and childhood premorbid indicators of later psychosis: Next steps for early psychosocial interventions. *Schizophr Bull* 2015; 41: 801–816. [PubMed: 25904724]
2. Keshavan MS, Delisi LE, Seidman LJ. Early and broadly defined psychosis risk mental states. *Schizophr Res* 2011; 126: 1–10. [PubMed: 21123033]
3. Fusar-poli P, Bonoldi I, Yung AR, Borgwardt S, Kempton MJ, Valmaggia L et al. Predicting psychosis: Meta-analysis of transition outcomes in individuals at high clinical risk. *JAMA Psychiatry* 2012; 69: 220–229.
4. Yung AR, McGorry PD. The Prodromal Phase of First-episode Psychosis: Past and Current Conceptualizations. *Schizophr Bull* 1996; 22: 353–370. [PubMed: 8782291]
5. Tandon R, Nasrallah HA, Keshavan MS. Schizophrenia, ‘just the facts’ 4. Clinical features and conceptualization. *Schizophr Res* 2009; 110: 1–23. [PubMed: 19328655]
6. Insel TR. Rethinking schizophrenia. *Nature* 2010; 468: 187–193. [PubMed: 21068826]
7. Simon AE, Borgwardt S, Riecher-Rössler A, Velthorst E, de Haan L, Fusar-Poli P. Moving beyond transition outcomes: Meta-analysis of remission rates in individuals at high clinical risk for psychosis. *Psychiatry Res* 2013; 209: 266–272. [PubMed: 23871169]
8. Schlosser DA, Jacobson S, Chen Q, Sugar CA, Niendam TA, Li G et al. Recovery from an at-risk state: Clinical and functional outcomes of putatively prodromal youth who do not develop psychosis. *Schizophr Bull* 2012; 38: 1225–1233. [PubMed: 21825282]
9. Wang C, Lee J, Ho NF, Lim JKW, Poh JS, Rekhi G et al. Large-Scale Network Topology Reveals Heterogeneity in Individuals With at Risk Mental State for Psychosis: Findings From the Longitudinal Youth-at-Risk Study. *Cereb Cortex* 2017; Epub ahead: 1–10. [PubMed: 28365777]
10. Anticevic A, Haut K, Murray JD, Repovs G, Yang GJ, Diehl C et al. Association of thalamic dysconnectivity and conversion to psychosis in youth and young adults at elevated clinical risk. *JAMA Psychiatry* 2015; 72: 882–891. [PubMed: 26267151]
11. Gu S, Satterthwaite TD, Medaglia JD, Yang M, Gur RE, Gur RC et al. Emergence of system roles in normative neurodevelopment. *Proc Natl Acad Sci USA* 2015; 112: 13681–13686. [PubMed: 26483477]
12. Meunier D, Lambiotte R, Bullmore ET. Modular and hierarchically modular organization of brain networks. *Front Neurosci* 2010; 4: 200. [PubMed: 21151783]
13. Sporns O, Betzel RF. Modular Brain Networks. *Annu Rev Psychol* 2016; 67: 613–640. [PubMed: 26393868]
14. Fair DA, Cohen AL, Power JD, Dosenbach NUF, Church JA, Miezin FM et al. Functional brain networks develop from a ‘local to distributed’ organization. *PLoS Comput Biol* 2009; 5: e1000381. [PubMed: 19412534]
15. Satterthwaite TD, Wolf DH, Ruparel K, Erus G, Elliott MA, Eickhoff SB et al. Heterogeneous impact of motion on fundamental patterns of developmental changes in functional connectivity during youth. *Neuroimage* 2013; 83: 45–57. [PubMed: 23792981]
16. Collin G, Keshavan MS. Connectome development and a novel extension to the neurodevelopmental model of schizophrenia. *Dialogues Clin Neurosci* 2018; 20: 101–110. [PubMed: 30250387]
17. Sporns O. Contributions and challenges for network models in cognitive neuroscience. *Nat Neurosci* 2014; 17: 652–60. [PubMed: 24686784]
18. Zhang T, Li H, Tang Y, Niznikiewicz MA, Shenton ME, Keshavan MS et al. Validating the Predictive Accuracy of the NAPLS-2 Psychosis Risk Calculator in a Clinical High-Risk Sample

- From the SHARP (Shanghai At Risk for Psychosis) Program. *Am J Psychiatry* 2018; 175: 906–908.
19. Zheng L, Wang J, Zhang T, Li H, Li C, Jiang K. The Chinese version of the SIPS/SOPS: a pilot study of reliability and validity. *Chinese Ment Heal J* 2012; 26: 571–576.
 20. Miller TJ, McGlashan TH, Rosen JL, Cadenhead K, Ventura J, McFarlane W et al. Prodromal assessment with the Structured Interview for Prodromal Syndromes and the Scale of Prodromal Symptoms: Predictive validity, interrater reliability, and training to reliability. *Schizophr Bull* 2003; 29: 703–715. [PubMed: 14989408]
 21. Wechsler D WASI Manual Psychological Corporation, Harcourt Brace, San Antonio, TX, 1999.
 22. McGlashan T, Walsh B, Woods S. *The Psychosis-Risk Syndrome: Handbook for Diagnosis and Follow-up*. New York: Oxford University Press, 2010.
 23. Fornito A, Zalesky A, Breakspear M. Graph analysis of the human connectome: Promise, progress, and pitfalls. *Neuroimage* 2013; 80: 426–444. [PubMed: 23643999]
 24. Sohn WS, Yoo K, Lee YB, Seo SW, Na DL, Jeong Y. Influence of ROI selection on resting functional connectivity: An individualized approach for resting fMRI analysis. *Front Neurosci* 2015; 9: 1–10. [PubMed: 25653585]
 25. Fischl B FreeSurfer. *Neuroimage* 2012; 62: 774–81. [PubMed: 22248573]
 26. Whitfield-Gabrieli S, Nieto-Castanon A. Conn: A functional connectivity toolbox for correlated and anticorrelated brain networks. *Brain* 2012; 2: 125–141.
 27. Makris N, Meyer JW, Bates JF, Yeterian EH, Kennedy DN, Caviness VS. MRI-Based Topographic Parcellation of Human Cerebral White Matter and Nuclei. *Neuroimage* 1999; 9: 18–45. [PubMed: 9918726]
 28. Blondel VD, Guillaume J-L, Lambiotte R, Lefebvre E. Fast unfolding of communities in large networks. *J Stat Mech* 2008; : P10008.
 29. Rubinov M, Sporns O. Weight-conserving characterization of complex functional brain networks. *Neuroimage* 2011; 56: 2068–2079. [PubMed: 21459148]
 30. Good BH, de Montjoye YA, Clauset A. Performance of modularity maximization in practical contexts. *Phys Rev E* 2010; 81: 46106.
 31. Doron KW, Bassett DS, Gazzaniga MS. Dynamic network structure of interhemispheric coordination. *PNAS* 2012; 109: 18661–18668. [PubMed: 23112199]
 32. Traud AL, Kelsic ED, Mucha PJ, Porter MA. Comparing Community Structure to Characteristics in Online Collegiate Social Networks. *SIAM Rev* 2008; 53: 526–543.
 33. Bordier C, Nicolini C, Forcellini G, Bifone A. Disrupted modular organization of primary sensory brain areas in schizophrenia. *NeuroImage Clin* 2018; 18: 682–693. [PubMed: 29876260]
 34. Lerman-Sinkoff DB, Barch DM. Network community structure alterations in adult schizophrenia: Identification and localization of alterations. *NeuroImage Clin* 2016; 10: 96–106. [PubMed: 26793435]
 35. Hoffman R, Dobschka S. Cortical Pruning and the Development of Schizophrenia: A Computer Model. *Schizophr Bull* 1989; 15: 477–490. [PubMed: 2814376]
 36. Hoffman R, McGlashan T. Parallel Distributed Processing and the Emergence of Schizophrenic Symptoms. *Schizophr Bull* 1993; 19: 119–140. [PubMed: 8451607]
 37. Hoffman RE, McGlashan TH. Synaptic elimination, neurodevelopment, and the mechanism of hallucinated ‘voices’ in schizophrenia. *Am J Psychiatry* 1997; 154: 1683–1689. [PubMed: 9396946]
 38. David AS. Dysmodularity: a neurocognitive model for schizophrenia. *Schizophr Bull* 1994; 20: 249–255. [PubMed: 8085128]
 39. Yu Q, Plis SM, Erhardt EB, Allen EA, Sui J, Kiehl KA et al. Modular Organization of Functional Network Connectivity in Healthy Controls and Patients with Schizophrenia during the Resting State. *Front Syst Neurosci* 2012; 5: 1–16.
 40. Van den Heuvel MP, Sporns O, Collin G, Scheewe T, Mandl RCW, Cahn W et al. Abnormal rich club organization and functional brain dynamics in schizophrenia. *JAMA psychiatry* 2013; 70: 783–92. [PubMed: 23739835]

41. Schmidt A, Crossley NA, Harrisberger F, Smieskova R, Lenz C, Riecher-Rössler A et al. Structural network disorganization in subjects at clinical high risk for psychosis. *Schizophr Bull* 2017; 43: 583–591. [PubMed: 27481826]
42. Alexander-Bloch AF, Gogtay N, Meunier D, Birn R, Clasen L, Lalonde F et al. Disrupted modularity and local connectivity of brain functional networks in childhood-onset schizophrenia. *Front Syst Neurosci* 2010; 4: 147. [PubMed: 21031030]
43. Supekar K, Musen M, Menon V. Development of large-scale functional brain networks in children. *PLoS Biol* 2009; 7: e1000157. [PubMed: 19621066]
44. Casey B, Jones R, Somerville L. Braking and Accelerating of the Adolescent Brain. *J Res Adolesc* 2011; 21: 21–33. [PubMed: 21475613]
45. Skudlarski P, Jagannathan K, Anderson K, Stevens MC, Calhoun VD, Skudlarska BA et al. Brain connectivity is not only lower but different in schizophrenia: a combined anatomical and functional approach. *Biol Psychiatry* 2010; 68: 61–9. [PubMed: 20497901]
46. Pettersson-Yeo W, Allen P, Benetti S, McGuire P, Mechelli A. Dysconnectivity in schizophrenia: where are we now? *Neurosci Biobehav Rev* 2011; 35: 1110–24. [PubMed: 21115039]
47. Zalesky A, Cocchi L, Fornito A, Murray MM, Bullmore E. Connectivity differences in brain networks. *Neuroimage* 2012; 60: 1055–1062. [PubMed: 22273567]
48. Honea R, Sc B, Crow TJ, Ph D, Passingham D, Ph D et al. Regional Deficits in Brain Volume in Schizophrenia: A Meta-Analysis of Voxel-Based Morphometry Studies. *Am J Psychiatry* 2005; 162: 2233–2245. [PubMed: 16330585]
49. Glahn DC, Laird AR, Ellison-Wright I, Thelen SM, Robinson JL, Lancaster JL et al. Meta-Analysis of Gray Matter Anomalies in Schizophrenia: Application of Anatomic Likelihood Estimation and Network Analysis. *Biol Psychiatry* 2008; 64: 774–781. [PubMed: 18486104]
50. Vita A, De Peri L, Deste G, Sacchetti E. Progressive loss of cortical gray matter in schizophrenia: A meta-analysis and meta-regression of longitudinal MRI studies. *Transl Psychiatry* 2012; 2: e190–13. [PubMed: 23168990]
51. Shenton ME, Dickey CC, Frumin M, Mccarley RW. A review of MRI findings in schizophrenia. *Schizophr Res* 2001; 49: 1–52.
52. Jung WH, Borgwardt S, Fusar-Poli P, Kwon JS. Gray matter volumetric abnormalities associated with the onset of psychosis. *Front Psychiatry* 2012; 3: 1–21. [PubMed: 22347194]
53. Tracy DK, Shergill SS. Mechanisms underlying auditory hallucinations - understanding perception without stimulus. *Brain Sci* 2013; 3: 642–669. [PubMed: 24961419]
54. Satterthwaite TD, Wolf DH, Loughhead J, Ruparel K, Elliott MA, Hakonarson H et al. Impact of in-scanner head motion on multiple measures of functional connectivity: Relevance for studies of neurodevelopment in youth. *Neuroimage* 2012; 60: 623–632. [PubMed: 22233733]
55. Power JD, Barnes KA, Snyder AZ, Schlaggar BL, Petersen SE. Spurious but systematic correlations in functional connectivity MRI networks arise from subject motion. *Neuroimage* 2012; 59: 2142–2154. [PubMed: 22019881]
56. Behzadi Y, Restom K, Liau J, Liu TT. A component based noise correction method (CompCor) for BOLD and perfusion based fMRI. *Neuroimage* 2007; 37: 90–101. [PubMed: 17560126]
57. Murphy K, Birn RM, Handwerker DA, Jones TB, Bandettini PA. The impact of global signal regression on resting state correlations: are anti-correlated networks introduced? *Neuroimage* 2009; 44: 893–905. [PubMed: 18976716]
58. Chai XJ, Castañán AN, Öngür D, Whitfield-Gabrieli S. Anticorrelations in resting state networks without global signal regression. *Neuroimage* 2012; 59: 1420–1428. [PubMed: 21889994]
59. Murphy K, Birn RM, Bandettini PA. Resting-state FMRI confounds and cleanup. *Neuroimage* 2013; 80: 349–359. [PubMed: 23571418]
60. Rangaprakash D, Wu GR, Marinazzo D, Hu X, Deshpande G. Hemodynamic response function (HRF) variability confounds resting-state fMRI functional connectivity. *Magn Reson Med* 2018; 80: 1697–1713. [PubMed: 29656446]

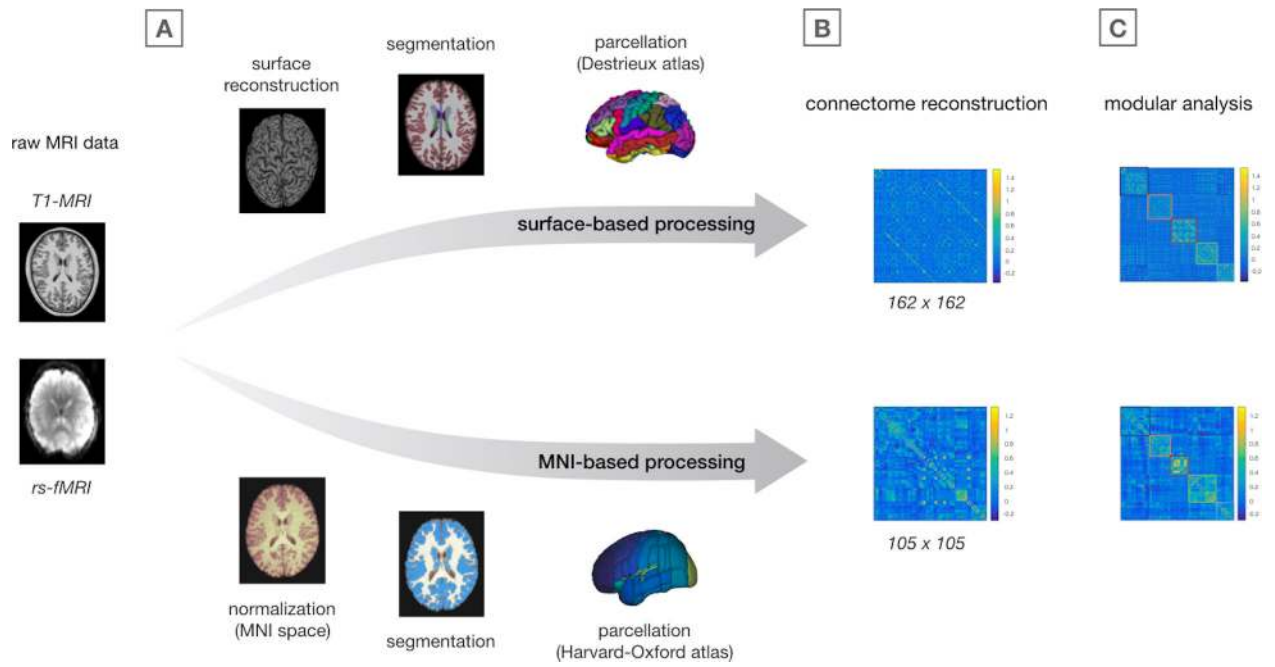


Figure 1.

Image preprocessing and connectome analysis steps

Overview of image preprocessing (A), connectome reconstruction (B), and modular community detection analysis (C). Of note, the colors of the modules shown in the connectivity matrices in panel C correspond to the modules as shown in Figure 2A.

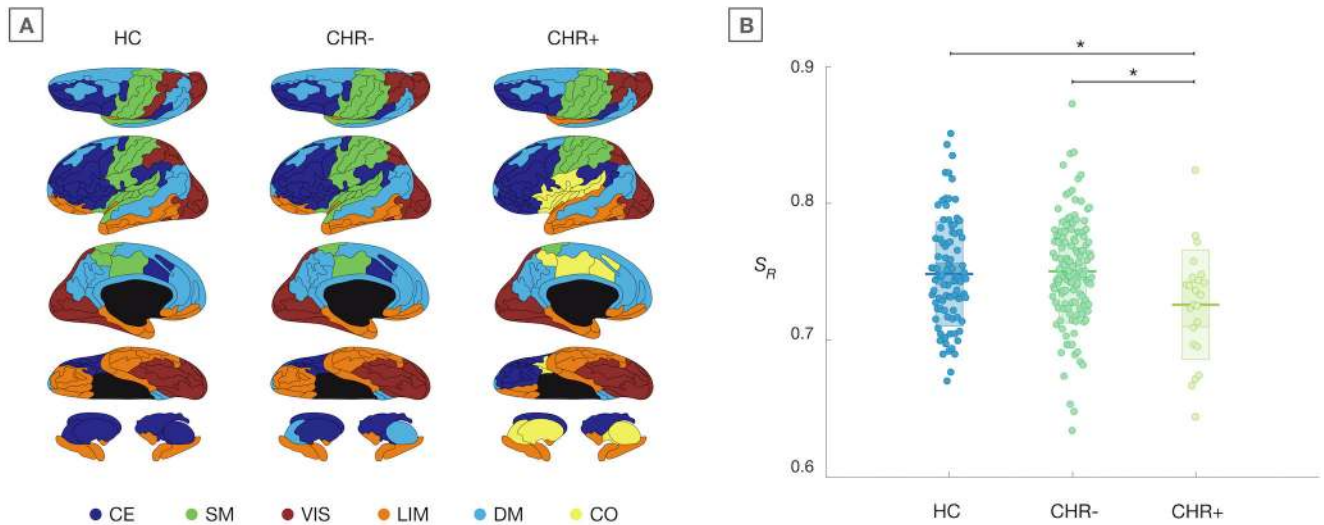


Figure 2.

Modular organization of the functional connectome

A) Modular partitions of group-networks, plotted on cortical surface from superior, lateral, medial, and inferior angle, and subcortical structures (top to bottom row respectively). Colors indicate separate modules, with the prefrontal central-executive (CE) module in dark blue, the central sensorimotor (SM) module in green, the posterior visual (VIS) module in red, the (para)limbic (LIM) module in orange, the medial default mode (DM) module in light blue, and the cingulo-opercular (CO) module in yellow. B) Degree of similarity to average healthy network (S_R) for individual subjects. Jittered data are plotted for each group, with mean (sd) values represented by the box behind the raw data. * indicates significant group-difference.

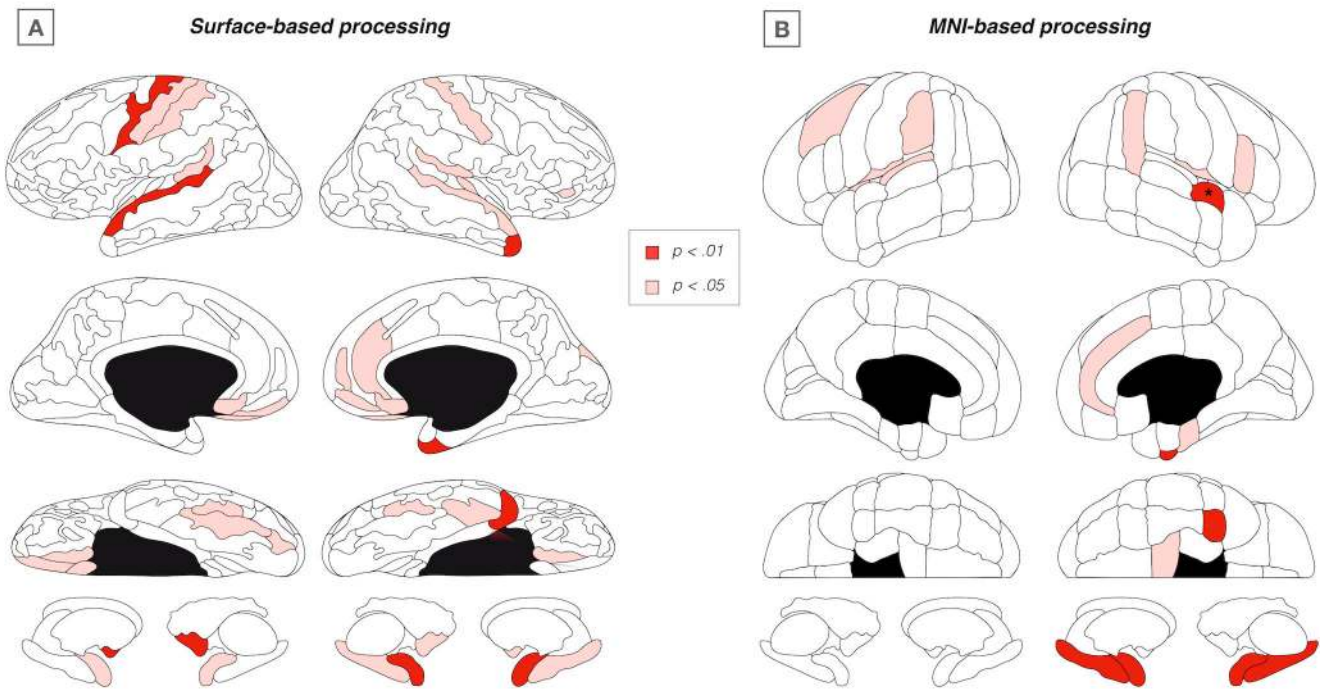


Figure 3.
 Regional findings of abnormal module assignment
 Surface plots showing exploratory regional findings (S_{Rnode}) at uncorrected $p < .05$ for surface-based (A) and MNI-based (B) processing methods respectively. * FDR-corrected significant effect for right superior temporal gyrus, anterior division.

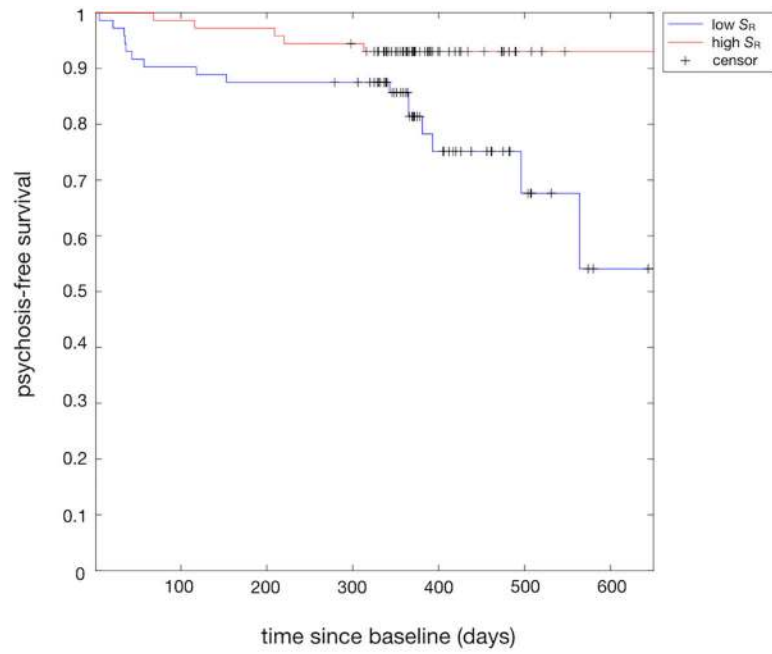


Figure 4. Psychosis-free survival for typical vs. atypical baseline connectome organization
Kaplan-Meier plot showing psychosis-free survival functions for CHRs with above-average (red) and below-average (blue) levels of S_R (reflecting typical and atypical connectome organization respectively) as a functional of time since baseline (days).

Table 1.

Demographic and clinical characteristics

Statistical comparison was performed using analysis of variance (ANOVA) tests for continuous, and chi-squared tests for categorical variables.

	CHR+N = 23	CHR-N = 135	ControlsN = 93	Statistics
Age, mean (sd) [range]	19.2 (5.2)[14 – 34]	18.7 (4.9)[13 – 32]	18.7 (4.6)[12 – 35]	$F = 0.10, p = .905$
Sex, male/female	16 / 7	64 / 71	49 / 44	$\chi^2 = 3.96, p = .138$
Education in years, mean (sd) [range]	10.3 (2.2)[7 – 16]	10.5 (2.9)[4 – 19]	10.8 (2.3)[6 – 17]	$F = 0.52, p = .0597$
IQ, mean (sd) [range]	92.1 (19.0)[52 – 112]	99.9 (11.2)[67 – 128]	104.2 (11.1)[75 – 133]	$F = 8.53, p < .001^I$
Baseline SIPS scores				
Positive, mean (sd) [range]	10.0 (3.3)[4 – 17]	10.1 (3.7)[0 – 21]		$F = 0.03, p = .871$
Negative, mean (sd) [range]	12.1 (6.4)[3 – 26]	11.6 (6.1)[1 – 27]		$F = 0.16, p = .687$
Disorganized, mean (sd) [range]	6.5 (3.0)[2 – 13]	6.6 (3.3)[1 – 19]		$F = 0.02, p = .891$
General, mean (sd) [range]	9.0 (2.9)[3 – 14]	9.1 (3.3)[1 – 17]		$F = 0.01, p = .902$
Total, mean (sd) [range]	37.6 (10.7)[16 – 65]	37.3 (10.9)[13 – 79]		$F = 0.01, p = .922$
Psychotropic medication				
At inclusion, N (%)	1 (4.3)	6 (4.4)		$\chi^2 < 0.01, p = .983$
At baseline MRI, N (%)	7 (30.4)	22 (16.3)		$\chi^2 = 2.62, p = .105$
Antipsychotics, N (%)	6 (26.1)	18 (13.3)		$\chi^2 = 2.48, p = .115$
Antidepressants, N (%)	2 (8.7)	5 (3.7)		$\chi^2 = 1.57, p = .282$
Other, N (%)	1 (4.3)	1 (0.7)		$\chi^2 = 2.05, p = .153$
GAF highest, mean (sd) [range]	77.6 (2.6)[73 – 83]	77.3 (4.9)[47 – 85]		$F = 0.08, p = .776$
GAF current, mean (sd) [range]	52.7 (7.7)[43 – 78]	54.1 (8.4)[21 – 76]		$F = 0.58, p = .449$

^I post-hoc analysis (Tukey-Kramer) indicates a significant IQ difference between each pair of subject groups (HC vs. CHR+, $p = .002$; HC vs. CHR-, $p = .022$; CHR- vs CHR+, $p = .039$).