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Frontostriatothalamic effective connectivity and dopaminergic function in the psychosis continuum

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Abstract

Dysfunction of fronto-striato-thalamic (FST) circuits is thought to contribute to dopaminergic dysfunction and symptom onset in psychosis, but it remains unclear whether this dysfunction is driven by aberrant bottom-up subcortical signaling or impaired top-down cortical regulation.

We used spectral dynamic causal modelling of resting-state functional magnetic resonance imaging (fMRI) to characterize the effective connectivity of dorsal and ventral FST circuits in a sample of 46 antipsychotic-naïve first-episode psychosis patients and 23 controls and an independent sample of 36 patients with established schizophrenia patients and 100 controls. We also investigated the association between FST effective connectivity and striatal [^{18}F]DOPA uptake in an independent healthy cohort of 33 individuals who underwent concurrent fMRI and positron emission tomography.

Using a posterior probability threshold of 0.95, we found that midbrain and thalamic connectivity were implicated as dysfunctional across both patient groups. Dysconnectivity in first-episode psychosis patients was mainly restricted to the subcortex, with positive symptom severity being associated with midbrain connectivity. Dysconnectivity between the cortex and subcortical systems was only apparent in established schizophrenia patients. In the healthy [^{18}F]DOPA cohort, we found that striatal dopamine synthesis capacity was associated with the effective connectivity of nigrostriatal and striatothalamic pathways, implicating similar circuits to those associated with psychotic symptom severity in patients.

Overall, our findings indicate that subcortical dysconnectivity is evident in the early stages of psychosis, that cortical dysfunction may emerge later in the illness, and that nigrostriatal and striatothalamic signaling are closely related to striatal dopamine synthesis capacity, which is a robust marker for psychosis.

Keywords: effective connectivity; dopamine; frontostriatal; psychosis; schizophrenia

Abbreviations: DCM: dynamic causal model; dlPFC: dorsolateral prefrontal cortex; FEP: first-episode psychosis; FST: fronto-striato-thalamic; PEB: parametric empirical bayes; Pp: posterior probability; SN: substantia nigra; vmPFC: ventromedial prefrontal cortex; VTA: ventral tegmental area

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Introduction

Dysfunction of fronto-striato-thalamic (FST) circuits linking the caudate and putamen with the prefrontal cortex is thought to be central to the emergence of psychotic symptoms.[1–6] Two such circuits are particularly relevant: 1) a ventral 'limbic' system involved in emotional and reward processing, which connects the orbital and ventromedial prefrontal cortices and subcortical limbic structures (e.g., hippocampus and amygdala) with the nucleus accumbens (NAcc); and 2) a dorsal 'associative' system, which links the dorsolateral prefrontal cortex (dlPFC) with the dorsal striatum, and subserves associative learning and executive functions.[1,7] Feedback loops passing through the pallidum and the thalamus connect both circuits back to the cortex.[1]

The striatum is a major target for dopamine projections from the ventral tegmental area (VTA) and substantia nigra (SN), with the ventral and the dorsal regions respectively forming part of the mesolimbic and nigrostriatal pathways.[8] Dysregulated dopamine signaling is proposed to contribute to psychosis onset by influencing the capacity of the striatum to filter and relay information to the thalamus, thus affecting broader FST function.[9] In-vivo positron emission tomography (PET) has revealed elevated presynaptic dopamine synthesis capacity, measured using 3,4-dihydroxy-6-[^{18}F]-fluoro-L-phenylalanine ([^{18}F]DOPA), in the dorsal striatum of at-risk groups, especially in individuals who transition to psychosis.[10,11] Increased dopamine binding and release in the dorsal striatum have also been shown in psychosis patients and at-risk individuals.[12,13] Moreover, [^{18}F]DOPA elevations in the ventral striatum of schizophrenia patients and of at-risk individuals have been reported.[14,15] In parallel, resting-state functional magnetic resonance imaging (fMRI) studies have identified reduced functional connectivity between the dorsal striatum, thalamus, and dlPFC in first episode psychosis (FEP) patients, their unaffected first-degree relatives, at-risk individuals, chronic unmedicated patients, and healthy

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people with psychosis-like experiences.[2–5,16,17] Increased functional connectivity between the ventral striatum, limbic regions, anterior cingulate cortex, ventromedial prefrontal and orbitofrontal cortices has also been found in these groups.[2,18–20] Correlations between striatal [^{18}F]DOPA activation of prefrontal and medial temporal areas have been reported in early and chronic stages of psychotic illness.[21–23]

Functional connectivity quantifies statistical dependencies between regional activity and does not distinguish causal interactions. It is therefore unclear whether FST dysfunction in psychosis arises from altered bottom-up signaling or disrupted top-down regulation of subcortical systems. Evidence for a primary bottom-up pathology comes from reports of aberrant molecular function, activity, and functional connectivity of the midbrain across at-risk and schizophrenia groups.[24–26] Others have proposed, largely based on ex-vivo and preclinical findings, that subcortical changes are secondary to deficient top-down control of midbrain neurons, which is caused by GABA/glutamate imbalance in the cortex or hippocampus.[9,27,28] However, in-vivo imaging findings have been mixed, with reports of increased, decreased, or no changes in prefrontal excitatory or inhibitory metabolites in FEP[29–33] and people with established schizophrenia.[30,31,34,35]

Top-down and bottom-up interactions between distinct elements of FST systems can be disentangled through models of effective connectivity. Effective connectivity refers to the causal influence that one neural system exerts over another[36] which can be estimated from fMRI data using dynamic causal modelling (DCM). DCM uses a Bayesian framework for optimizing a generative model of directed, or causal, influences between regions comprising a distributed neural system.[36] In DCM, inter-regional connections are referred to as extrinsic connections[37] and they can be used to infer top-down influences from cortical to subcortical areas or bottom-up subcortical-to-cortical interactions. Another important feature of DCM is that it allows the investigation of the intrinsic functioning of a brain region through the estimation of recurrent inhibition, modelled as a self-connection, which can uncover intra-regional pathology.[37] Here, we used spectral DCM DCM[38] of resting-state fMRI data to characterize disruptions of extrinsic and recurrent intrinsic effective connectivity within dorsal and ventral FST systems in antipsychotic-naïve FEP patients and people with established schizophrenia (SCZ), as well as associations with psychotic symptoms, which have historically been linked to FST dysfunction.[39] In an independent healthy cohort who underwent concurrent fMRI and [^{18}F]DOPA PET we then identified the specific FST connections that are associated with dorsal and ventral striatal [^{18}F]DOPA levels. This approach allowed us to cross-sectionally map effective dysconnectivity of FST circuits across different illness stages and to identify putative directed influences associated with striatal dopamine synthesis capacity, a robust marker of illness risk.[10,40,41]

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Materials and methods

Participants

We examined three independent cohorts: (1) a FEP cohort comprising antipsychotic-naïve patients within six months of psychosis onset and healthy controls;[42] (2) a SCZ cohort comprising schizophrenia patients and healthy controls obtained through the UCLA Consortium for Neuropsychiatric Phenomics open dataset;[43] and (3) an [^{18}F]DOPA cohort comprising healthy participants recruited from the community who underwent concurrent PET/fMRI scans. The FEP and SCZ cohorts allowed us to examine FST effective connectivity at distinct illness stages, while the [^{18}F]DOPA cohort allowed us to identify specific connections associated with striatal dopamine synthesis capacity. Separate analyses were also performed for a subgroup of the FEP cohort with a schizophrenia diagnosis (FEP-SCZ) to examine whether there were diagnosis-specific effects within the FEP group. Group numbers and demographic details are provided in Table 1. The recruitment of our FEP cohort and controls, and the [^{18}F]DOPA cohort was in accordance with the Melbourne Health Human Research Ethics Committee (reference numbers 2007.616 and 1443066.1). Below are the recruitment details for each of our cohort. Detailed description of the number of recruited participants excluded from our analyses are outlined in Supplementary Material.

First-episode psychosis (FEP)

Participants were aged 15–25 years, presenting with FEP to an early psychosis intervention centre at Orygen Youth Health, Melbourne, Australia.[42] FEP was defined as fulfilling Structured Clinical Interview for DSM-5 (SCID-5) criteria for a psychotic disorder. Patients with substance-induced psychosis were excluded in the present study. Healthy controls had no history of psychiatric or neurological illness, as determined by self-reports, SCID, and the Comprehensive Assessment of At-Risk Mental States. Recruitment period occurred from April 2008 until December 2016. All participants provided informed consent. For FEP patients, additional strict inclusion criteria include comprehension of English language, no contraindication to MRI scanning, less than six months of duration of untreated psychosis, living in stable accommodation, low risk to self or others (score of < 5 on the Brief Psychiatric Rating Scale version 4 (BPRS-4) Suicidality and Hostility subscales), and minimal previous exposure to antipsychotic medication (less than 7 days of use or lifetime 1750mg chlorpromazine equivalent exposure). Healthy controls were age matched to patients. Additional exclusion criteria for controls included history of psychiatric or neurological illness in first-degree relatives and current use of psychotropic medications. A total of 46 patients and 23 controls were included in our final analyses. We also analyzed a subsample of FEP patients with schizophrenia diagnoses (FEP-SCZ*; n* = 17) to identify the differences in connectivity based on differential diagnoses.

Established schizophrenia (SCZ)

Data were obtained through the UCLA Consortium for Neuropsychiatric Phenomics (CNP) open dataset.[43] The initial sample consisted of 121 right-handed healthy controls and 51 schizophrenia

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patients aged between 21–50 years old. Healthy controls were excluded if they had a lifetime history of mental illness, including schizophrenia or other psychotic disorders, and bipolar disorders Healthy controls were also excluded on current diagnosis of major depressive disorder, suicidality, anxiety disorder or attention deficit hyperactivity disorder (ADHD). Additional exclusion criteria for all participants included left-handedness, pregnancy, neurological disease, history of head injury with loss of consciousness or cognitive sequelae, use of psychoactive medications, substance abuse or dependence, and other MRI contraindications (e.g., claustrophobia). SCZ patients were also excluded if they had comorbidity with either bipolar disorder or ADHD. Diagnoses were based on the DSM-IV and SCID-1. All SCZ patients were on stable medication. Mean lifetime chlorpromazine-equivalent dose[44] for patients with available medication information is 662.51mgs (SD = 764.91mgs; *n* = 28). No significant correlations between antipsychotic exposure and effective connectivity were identified (i.e., all $p > .05$, corrected). A total of 36 patients and 100 controls were included in our analyses after quality checking.

Healthy [^{18}F]DOPA cohort

A total of 52 healthy participants aged 18–28 years old were recruited from the community through online advertisements. Recruitment period was between May 2016 until May 2019. Each participant provided informed consent. Exclusion criteria included current or history of psychiatric or neurological illnesses, significant medical history, intellectual disability, and first-degree relative with a mental illness. Participants underwent simultaneous resting-state fMRI and PET imaging with [^{18}F]DOPA tracer. A total of 33 participants were included in our final analyses.

Symptom measures

Positive symptoms were assessed in patients using the positive subscale of Brief Psychiatric Rating Scale (BPRS) version 4.[45] The subscales were derived from a five factor solution outlined in Dazzi et al.[45] Psychotic symptom scores were derived from the positive symptom subscale which consists of five items: unusual thought content, suspiciousness, hallucinations, grandiosity, and bizarre behavior. Secondary analyses also considered negative symptoms assessed with the same instrument. The negative symptom subscale consists of three items: blunted affect, emotional withdrawal, and motor retardation. Each item is ranked on a Likert-like scale between 1 to 7, with a score of 1 denoting the absence of a measured symptom.

fMRI acquisition

The scanning acquisition protocol for each participant group is described in the following.

FEP

Whole-brain T2*-weighed echo-planar images (EPIs) and anatomical T1-weighted (T1w) scans were acquired for each participant using a 3T Siemens Trio Tim scanner, equipped with a 32-channel head coil, located at the Royal Children's Hospital in Melbourne, Australia. Participants were instructed to lie still in the scanner while maintaining wakefulness with eyes closed. A total of 234 functional volumes

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with 37 slices each were acquired using the following parameters: repetition time = 2000ms; echo time = 32ms; flip angle = 90°; field of view = 210mm; slice thickness of 3.5 mm, and 3.3 x 3.3 x 3.55 mm voxels. A total of 176 slices were acquired for each participant's T1-weighted image using an interleaved acquisition using the following parameters: TR = 2.3s; TE = 2.98ms; flip angle of 9°; FOV of 256mm; voxel size of 1.1 x 1.1. x 1.2 mm.

SCZ

The UCLA CNP dataset[43] was acquired on one of two Siemens Trio 3T scanners located at the Ahmanson-Lovelace Brain Mapping Center and the Staglin Center for Cognitive Neuroscience at UCLA. Details of the resting-state EPI scan are TR = 2s, TE = 30ms , flip angle = 90°, 4mm slice thickness, 152 volumes with 34 slices each. Participants were instructed to keep their eyes open. Details of the T1 scan are TR = 1.9s, TE = 2.26ms, flip angle of 90°, 176 slices with $1mm^3$ voxels.

[^{18}F]DOPA

Data were acquired using a 3T Siemens Magnetom Biograph simultaneous MR-PET scanner equipped with a 20-channel head and neck coil at Monash Biomedical Imaging, Melbourne, Australia. Resting-state whole-brain T2*-weighed echo-planar image (EPIs) was also acquired for each subject using an interleaved acquisition with the following parameters: TR = 2.89s; TE = 30ms; 152 volumes with 44 slices per volume; flip angle = 90°; FOV = 190mm; slice thickness = 3mm; voxel size of $3mm^3$. A high resolution T1-weighted anatomical image was acquired for each subject using an ascending acquisition (176 slices; TR = 1640ms; TE = 2.34ms; flip angle = 8°; field of view (FOV) = 256mm; slice thickness = 1mm; voxel size = $1mm^3$).

MRI processing

fMRI and T1-weighted data were processed in the same way across all three cohorts using FMRIPREP software version 1.1.1.[46] Each T1w scan was corrected for non-uniformity in intensity and subsequently skull stripped. Brain surfaces were reconstructed using FreeSurfer version 6.0.1. Tissue masks were generated using FreeSurfer. Spatial normalization of the skull stripped T1w images to the ICBM 152 Nonlinear Asymmetrical template version 2009c was performed using a nonlinear registration in ANTs version 2.1.0. Similarly, for each participant, tissue masks were registered from the surface space to the MNI template.

EPIs were slice-timed corrected using AFNI version 16.2.07 and realigned to a mean reference image using FSL. EPIs were distortion corrected using fieldmaps (phasediff-based workflow; https://fmriprep.readthedocs.io/en/stable/api/index.html#sdc-phasediff). For participants with missing fieldmaps, a "fieldmap-less" distortion correction was performed by co-registering the functional image to the intensity-inverted T1w image constrained with an EPI distortion atlas.[47] Following distortion correction, EPIs were co-registered with their corresponding T1w using boundary-based registration with nine degrees of freedom using the bbregister routine in FreeSurfer. The motion-correcting transformations, field-distortion-correcting warp, EPI-to-T1w transformation, and T1w-to-MNI template

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warp were concatenated and applied in a single step using ANTs version 2.1.0. In-scanner head motion was defined as excessive according to previously-defined stringent exclusion criteria;[48] namely, if any of the following were met: 1) mean framewise displacement (FD) > 0.20mm; (2) sum of suprathreshold FD spikes > 20%; and (3) any FD > 5mm. FD was calculated using the root mean squared volume-to-volume displacement of all voxels, derived from the six head motion parameter (3 translations, 3 rotations).[48] ICA-based Automatic Removal of Motion Artifacts (AROMA) was used to generate noise regressors.[49] ICA-AROMA noise components were non-aggressively regressed out from the EPI data. Next, white matter and cerebrospinal fluid signals were estimated from each subject using restricted tissue masks and were removed out from the data via linear regression. A high-pass filter of 200 seconds was the applied to each EPI image prior to spatial smoothing using a Gaussian kernel of 4mm full-width half-maximum implemented using the 3dBlurToFWHM function in AFNI (version 16.2.16). We did not perform global signal regression as DCM incorporates noise parameter estimates that capture observation noise due to scanner and physiological noise[38,50] and also because it may remove real neuromodulatory fluctuations in neuronal activity, which are of key interest in the present study (for a discussion, see Aquino et al.[51] and Glasser et al.[52]).

PET acquisition, processing, and analysis

PET data in the [^{18}F]DOPA group were acquired on a MR-PET Siemens Biograph scanner. All participants received carbidopa (150 mg) and entacapone (400 mg) orally 60 minutes before imaging to reduce the formation of radiolabeled metabolites that can cross the blood-brain barrier and thus confound tracer availability in the striatum.[53,54] All subjects were administered approximately 150Mbq of [^{18}F]DOPA via interbolus injection at the start of the PET imaging. Participants were instructed to lie still in the scanner with eyes closed. Participants were instructed to lie still in the scanner with eyes closed.

The pseudo-CT attenuation correction method[55,56] was used to correct PET images during image reconstruction. Dynamic PET images were reconstructed using the Siemens e7tools software with image volume size 344×344×127(2.09×2.09×2.03mm^3). The Ordinary Poisson-Ordered Subset Expectation Maximization (OP-OSEM) algorithm (3 iterations, 21 subsets) was used with the point spread function (PSF) for partial volume correction. A 5-mm FWHM Gaussian filter was applied to each 3D image volume. For correction of subject motion, the list mode dataset was first binned into 95 frames consisting of one 30-second background frame and ninety-four 60-second frames. Dynamic motion was corrected based on an image registration approach,[57] where each frame was registered to the last frame using rigid-body transformation implemented in the FSL toolbox.[58] The final reconstructed dynamic PET images were registered to the corresponding T1 MPRAGE MRI image. Patlak graphical analysis[59] was performed using Qmodeling software[60] to quantify [^{18}F]DOPA influx rate constants (K_i^{cer} values) for dorsal and ventral striatal regions-of-interest (ROIs)[61] (Supplementary Fig. S1) relative to the cerebellum[62] in each participant's anatomical space. We only analyzed the left hemisphere consistent with our DCM (see below).

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Dynamic causal modeling

Model space selection

Regions of interest (ROIs) spanning the dorsal and ventral FST systems, and the midbrain were selected using stereotactic coordinates of past findings or of peak signals identified using functional connectivity. Effective connectivity explains the change of activity in one region as a function of activity in another region and does not depend on signal correlations between two areas. As such, selecting regions based on functional connectivity peaks does not result in biased ROI selection. We modelled 47 biologically plausible connections, identified through tract tracing or human functional connectivity, between eight ROIs (Fig. 1). The first eigenvariate of time-series of each ROI was extracted and entered into first-level DCM analysis. We focused on the left hemisphere given prior evidence of more consistent functional connectivity effects in left-lateralized FST circuits in patients.[2,3] Spherical ROIs were created with a radius of 6mm for cortical regions (i.e., dlPFC and vmPFC) and 3.5mm for each subcortical ROI. For vmPFC and thalamus ROIs with centroids that were close to the anatomical or functional boundary of the region, we used an anatomical mask from the FreeSurfer package when extracting the first eigenvariate for the ROIs to exclude signal from neighboring regions. Details of ROI selection for each region are as below.

Nucleus accumbens (NAcc) and dorsal caudate (DC)

For the ventral and dorsal striatum, we seeded the NAcc and DC consistent with our previous works[2,4,63] using a functional parcellation of the striatum[64] that was delineated based on a meta-analysis of striatal activation in fMRI and PET studies.[65] The pallidum, which is the primary output structure of the basal ganglia, was omitted to limit model dimensionality. Its effects will thus be captured through indirect (unmodelled) influences in the DCM.

Dorsolateral prefrontal cortex (dlPFC)

To capture variations associated with risk for psychosis as well as disruptions in clinical groups, the dlPFC ROI was selected based on a peak in which functional connectivity with the DC was reduced in the healthy relatives of FEP patients compared to healthy controls.[2] This peak overlapped with an area that also showed reduced connectivity in patients.

Ventromedial prefrontal cortex (vmPFC)

To choose a region that was most relevant to the ventral circuit, we chose a peak in the vmPFC that showed strong functional connectivity with the NAcc in an independent cohort of 353 healthy adults.[4]

Thalamus

The thalamus coordinate was selected from a peak in which functional connectivity with the dorsal caudate was reduced in at-risk mental state individuals compared to healthy controls.[63]

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Hippocampus and amygdala

ROIs for these regions were selected using a similar method. Peak coordinates within anatomical masks for the hippocampus and amygdala were selected in a separate group-level functional connectivity analysis of each ROIs connectivity with the whole brain in an independent cohort of 353 individuals.[4] For the hippocampus, we chose the anterior region, using of a hippocampal mask provided in the Freesurfer package.[66] The anterior hippocampus is a hippocampal region that is most frequently implicated as dysfunctional in schizophrenia,[67] and it was defined as the region having the MNI coordinate of less than y = -22.[68] Similar to the hippocampus, we used an amygdala mask provided in FreeSurfer.

Midbrain

The midbrain is a challenging area to identify due to the lack of tissue contrast afforded by anatomical T1 scans. We therefore employed a midbrain mask derived from a comprehensive study of functional connectivity between the ventral tegmental area (VTA), substantia nigra (SN) and the rest of the brain, which has been that was validated in two independent datasets.[69] Due to the coarse resolution of the functional scans, we combined VTA and SN into one midbrain ROI, which limits our ability to delineate mesolimbic and nigrostriatal pathways. We selected a functional connectivity signal peak in the same cohort that was used in the selection of our hippocampal, amygdala, and vmPFC ROIs.

Model estimation

We modelled effective connectivity in the spectral domain by fitting a complex cross spectral density using a generative model, parametrized by a power-law model of endogenous fluctuations, implemented in SPM12 (DCM 12; revision 7487).[38] Details are provided in Supplementary Methods. Briefly, subject-specific first-level analyses were used to estimate directed (causal) influences between regions (in Hz), and the (inhibitory) recurrent or self-connectivity (i.e., self-inhibition) within each region.

Subject-specific connectivity parameters were then passed to a group-level general linear model (GLM) implemented in the parametric empirical Bayes (PEB) framework to estimate between-subject variability.[70] PEB models were run separately for each cohort. FEP and SCZ patients were compared to their respective control groups. Symptom associations in patients were modelled separately for each group for each connection in the model space, with both positive and negative symptoms included as covariates. Associations with [^{18}F]DOPA were modelled in healthy individuals only, with dorsal and ventral striatal K_i^{cer} values as covariates. Age, sex, and mean framewise displacement[71] were used as nuisance covariates for all models to address unequal distributions of sex across groups, as well as age and head motion as confounding variables. Scanner site was also used as a covariate in the SCZ group. We only report effects with a posterior probability threshold above 0.95. This threshold is based upon the differences in model evidence, or marginal likelihood, observed when comparing models with and without a particular connection. Critically, PEB is a multivariate (Bayesian) GLM in which we fit all model parameters at once; hence, no correction for multiple comparisons is required. A typical effect size for effective connectivity between regions is 0.1 Hz.[38] Self-connections are inhibitory to reflect activity decay, with a negative sign indicating reduced self-inhibition (i.e., disinhibition, or slower activity decay)

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and a positive sign indicating increased self-inhibition (i.e., quicker activity decay). Note that a group difference of 0 Hz in effective connectivity does not imply the absence of that connection in either group, but instead signifies that the groups do not differ in their effective connectivity. A mean connectivity of 0 Hz within a group denotes an absence of effective connectivity, but it does not necessarily imply a lack of anatomical connectivity. Technical details are outlined in Supplementary Methods.

Checks on variance explained

Connectivity methods based on correlations such as functional connectivity depend on covariances between brain regions, which are influenced by observation noise and changes in signal-to-noise ratio.[37] In contrast to descriptive correlational methods, DCM accounts for noise by modelling observation noise in the data, which includes thermal and physiological noise[50] (see equations S2 and S3 in Supplementary Methods). The DCM routine also incorporates a diagnostic check to determine the accuracy of model inversion by calculating the percentage of variance explained by the model based on the predicted BOLD signal.[72] Based on this diagnostic check, we excluded subjects with <75% variance explained by DCM for subsequent analyses following first-level model inversion (details in Supplementary Methods).

Data availability

Data from the UCLA Consortium for Neuropsychiatric Phenomics are available on an open dataset platform on https://openneuro.org/datasets/ds000030/versions/00016. Derived data from our FEP and PET cohorts are available from the corresponding author on request.

Results

Group differences in effective connectivity

FEP vs. healthy controls

To examine FST disruptions at the earliest signs of illness, before exposure to antipsychotic medication, we first investigated effective connectivity disruptions in antipsychotic-naive FEP patients relative to healthy controls. Patients showed reduced self-inhibition of VTA/SN, increased self-inhibition of dlPFC, greater inhibitory influence of the thalamus on NAcc, and a greater excitatory influence of the amygdala on NAcc (Fig. 2A; Table 2). The subset of FEP-SCZ patients (Fig. 2B; Table 2) showed similar changes but did not show increased dlPFC self-inhibition. FEP-SCZ patients additionally demonstrated a relative disinhibition of the amygdala, increased excitatory influence of VTA/SN on NAcc, and increased inhibitory influence of VTA/SN and NAcc on hippocampus. Connections between cortical and subcortical ROIs were not implicated as dysfunctional in FEP patients.

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SCZ vs. healthy controls

We next examined, in an independent sample of patients and controls, whether people with established schizophrenia show similar disruptions of FST effective connectivity to those observed in FEP. As with FEP, we found that SCZ patients showed disinhibition of VTA/SN and a greater inhibitory influence of thalamus on NAcc relative to controls (Fig. 2C; Table 2). SCZ patients also showed increased inhibitory influence of vmPFC on dlPFC, of dlPFC on thalamus, and of thalamus and amygdala on hippocampus. FEP-SCZ patients and SCZ patients both showed altered bottom-up connectivity of the VTA/SN. In FEP-SCZ, these disruptions primarily affected the limbic circuit; in SCZ patients, they affected the dorsal system, with increased inhibitory influence on the dorsal caudate (DC) on VTA/SN and an increased excitatory influence running in the reverse direction, from DC to VTA/SN.

Associations with positive symptoms

Having identified both common and distinct disruptions of FST circuitry in FEP and SCZ patients, we next examined the association between specific FST connections (i.e., how much a region influences another) and psychotic symptom severity in both groups.

FEP

Greater positive symptom severity was predominantly associated with subcortical connectivity (Fig. 3A, Table 3). Specifically, more severe symptoms were associated with a relatively stronger influence of VTA/SN on NAcc and amygdala, of NAcc and hippocampus on VTA/SN. More severe symptoms were also associated with a weaker influence of VTA/SN on DC and hippocampus, and with reduced self-inhibition of the amygdala. In cortex, higher positive symptom ratings were associated with weaker influence of dlPFC on vmPFC and weaker self-inhibition of vmPFC.

The FEP-SCZ subgroup (Fig. 3B, Table 3) showed similar associations with vmPFC self-inhibition and subcortical connectivity, although associations with hippocampal to VTA/SN connectivity and amygdala self-inhibition were absent. Positive symptom severity was additionally associated with a weaker influence of NAcc on hippocampus, of hippocampus on amygdala, of DC on thalamus, and reduced dlPFC self-inhibition, coupled with a stronger influence of dlPFC on thalamus and stronger thalamic self-inhibition.

SCZ

Both cortical-subcortical and subcortical-cortical influences were associated with positive symptoms in SCZ patients (Fig. 3C, Table 3). Specifically, more severe positive symptoms were associated with a stronger top-down influence of vmPFC on VTA/SN and hippocampus; stronger bottom-up influence of VTA/SN on dlPFC; stronger influence of DC on thalamus and of thalamus on hippocampus; and greater dlPFC self-inhibition. Higher positive symptom ratings were also associated with weaker vmPFC and

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VTA/SN self-inhibition and a weaker influence of dlPFC on thalamus, thalamus on VTA/SN, VTA/SN on amygdala, hippocampus on vmPFC. Some common connections were implicated in both FEP-SCZ and SCZ patients (i.e., VTA/SN–amygdala, DC–thalamus, dlPFC–thalamus, and dlPFC self-inhibition), albeit with opposing polarity, suggesting that the link between positive symptoms and altered FST effective connectivity varies across illness stages.

Associations with negative symptoms

Associations with negative symptom for FEP, FEP-SCZ, and SCZ patients are presented in Supplementary Fig. S2 and Table S1. Associations between negative symptom severity and amygdala self-inhibition were consistent, albeit with reversed polarity, across the last two groups. Top-down cortical to subcortical connections were associated with negative symptom severity in FEP-SCZ patients, whereas SCZ patients showed associations with bottom-up subcortical to cortical connections. Similar to positive symptoms, negative symptom associations in FEP patients mainly implicated subcortical connections.

Associations with striatal dopamine synthesis capacity

The analysis of clinical patients identified a prominent role for subcortical dysconnectivity in both FEP and SCZ groups. Dopamine dysregulation plays a central role in the genesis of psychotic symptoms[73–75] and there is robust evidence for elevated pre-synaptic dopamine synthesis capacity, as measured with [^{18}F]DOPA, in the dorsal striatum of patients and at-risk individuals,[10,40,41]. Dopaminergic transmission is influenced by, and influences, FST function.[8,9,76,77] We therefore set out to identify FST circuit elements that are associated with striatal [^{18}F]DOPA in an independent sample of healthy individuals who underwent concurrent PET-fMRI.

Dorsal striatum

Significant associations with dorsal striatal [^{18}F]DOPA in healthy individuals largely implicated the thalamus (Fig. 4; Table 4). Specifically, higher [^{18}F]DOPA was associated with stronger thalamic self-inhibition; stronger thalamic influence over the VTA/SN and DC; stronger amygdala influence over thalamus; and a weaker influence of DC on thalamus. Stronger bottom-up influence of the amygdala on vmPFC was also associated with higher dorsal striatal [^{18}F]DOPA.

Ventral striatum

Ventral striatal [^{18}F]DOPA was associated with a distributed set of extrinsic effective connections centered on midbrain and thalamus (Fig. 4; Table 4). Specifically, higher [^{18}F]DOPA was associated with weaker self-inhibition of NAcc and amygdala, a weaker influence of these two regions on the thalamus, and a stronger influence of hippocampus on thalamus. Ventral striatal [^{18}F]DOPA was also associated with greater self-inhibition of the VTA/SN; weaker bottom-up influences of VTA/SN on the vmPFC and DC; weaker and stronger top-down influences of DC and hippocampus on VTA/SN, respectively; and greater DC self-inhibition.

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Discussion

FST dysfunction has been identified in psychotic illnesses,[2–5,16,18–20] but a characterization of how causal influences within these circuits are altered at distinct illness stages has been lacking. Using spectral DCM, we mapped the effective connectivity of dorsal and ventral FST circuits in FEP and established SCZ. Both clinical groups showed consistent disinhibition of VTA/SN and a stronger top-down inhibitory influence of the thalamus on NAcc. Altered top-down connectivity from the cortex to subcortex was only identified in established illness. Positive symptom severity was associated with a relative disinhibition of the VTA/SN in both illness stages. Additional associations were otherwise largely confined to subcortical connectivity in FEP and a distributed set of cortical and subcortical connections in SCZ patients. Concurrent PET-MRI in healthy individuals revealed distinct sets of FST connections associated with dorsal and ventral dopamine synthesis capacity, with DC and striatothalamic connectivity associated with both striatal dopamine synthesis and positive symptom severity. Our findings indicate that midbrain dysfunction and subcortical dysconnectivity is prominent in early illness stages, that cortical dysfunction becomes more salient in established illness, and that striatothalamic and nigrostriatal connectivity are related to both striatal dopamine synthesis capacity and positive symptom severity in patients.

Effective dysconnectivity of fronto-striato-thalamic circuitry

We observed VTA/SN disinhibition across both illness stages, suggesting that it represents a core feature of FST pathology in psychosis. Midbrain disinhibition is consistent with elevated striatal dopamine synthesis.[10] Given that we found no evidence of disrupted top-down influences on VTA/SN in FEP in our modelled regions, our findings thus suggest that prior reports of elevated striatal dopamine in [^{18}F]DOPA and other PET studies across different stages of psychosis may be linked to intrinsic dysregulation of the midbrain.[28,78]

Both FEP and SCZ groups also showed increased inhibitory influence of thalamus over NAcc. Excitatory thalamostriatal projections provide feedback for the striatum to maintain bottom-up signaling to thalamus and cortex in support of specific actions or behaviors.[79] A greater inhibitory influence of thalamus on NAcc in both FEP and SCZ thus implies a dysregulation of the thalamostriatal feedback pathway. Although this dysregulation was not directly tied to VTA/SN function in our findings, animal models suggest that NAcc dysregulation can disinhibit the midbrain in young adult rats.[28,80] Notably, we have recently found that antipsychotics rapidly and preferentially impact thalamo-striatal and thalamo-cortical functional connectivity in FEP patients,[81] suggesting that a remediation of thalamic interactions with the rest of the brain may be a key therapeutic mechanism.

SCZ patients showed increased inhibitory influence of VTA/SN over DC, which may disrupt the capacity of this area to filter information through the FST circuits.[9,82] SCZ patients also showed increased top-down excitatory influence of DC over VTA/SN, which may reflect a compensatory response to regulate disinhibited signaling from the midbrain. Given that VTA/SN disinhibition was also observed in FEP, our findings suggest that an early bottom-up pathology of the midbrain may evolve to affect striatal

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function and potentially dysregulate feedback loops within the dorsal FST with illness progression. However, we cannot rule out an effect of medication in the SCZ group and longitudinal data are required to test this hypothesis.

In FEP, cortical dysfunction was limited to increased dlPFC self-inhibition. Post-mortem work has identified robust changes in GABAergic neuron function in patients with established schizophrenia,[83] but in vivo studies of FEP patients have been inconsistent, with reports of lower, higher, or no differences in prefrontal GABA levels.[32,33,84,85] Our findings thus suggest that while DLPFC dysfunction is evident early in psychosis, it does not to play a prominent role in broader FST dysfunction.

Taken together, our DCM analysis indicates that the early phase of psychosis is associated with prominent dysfunction of subcortical systems, with alterations in cortico-subcortical connectivity emerging in later illness stages. Intrinsic dysfunction of the VTA/SN and dlPFC, and altered thalamic regulation over the NAcc, appear to be stable features of the illness.

Fronto-striato-thalamic connectivity and symptom severity

A primary role for VTA/SN dysregulation in psychosis is supported by our observations of consistent correlations between the connectivity of this region and positive symptom severity across illness stages.[26,86–88] However, the specific connections implicated varied across the patient groups. In the full FEP cohort, greater positive symptom severity was associated with connectivity between VTA/SN and limbic subcortical regions, which accords with a role for hippocampal–midbrain dysregulation identified in rodent models of psychosis.[28] Disinhibition of the vmPFC, another component of the limbic FST system, was negatively associated with positive symptom severity across the FEP, FEP-SCZ, and SCZ patients.[89,90]

Reduced influence of VTA/SN on DC was also associated with greater positive symptom severity in both FEP-SCZ and SCZ patients. When taken with evidence of robust [^{18}F]DOPA elevations in the DC in early psychosis,[10,11,41] our findings of intrinsic dysfunction of the midbrain across illness stages (see also references 23–25), and anatomical studies indicating that midbrain afferents to the dorsal striatum primarily originate in the SN,[8] our results identify a close link between nigrostriatal signaling and the expression of psychotic symptoms.

Several additional FST elements were consistently associated with positive symptom severity in both SCZ and FEP-SCZ patients, although the polarity of associations was often reversed between FEP and SCZ groups. This reversal may be due to several factors, including differences in the neural correlates of psychotic symptoms at different illness stages, an effect of medication in the SCZ group, or a combination of both. Dopamine is proposed to regulate neuronal signalling in complex ways, and it is as yet unclear how dopamine levels at different illness stages relate to measures of functional or effective connectivity.[91,92] Precisely understanding these effects, and the impact of antipsychotic medication, is an important topic for further investigation.

There was less consistency across the cohorts with respect to negative symptom associations, with different sets of connections being associated with symptom severity in the FEP and SCZ groups. In

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general, the SCZ cohort demonstrated associations that were more widespread across the cortical and subcortical regions compared to the FEP and FEP-SCZ groups. Across both FEP and SCZ cohorts, the dlPFC and subcortical limbic regions, including the amygdala and the hippocampus, were generally implicated, consistent with evidence of prefrontal and limbic involvement in this symptom dimension.[93–95]

Fronto-striato-thalamic connectivity and striatal dopamine synthesis capacity

Dorsal and ventral striatal [^{18}F]DOPA levels were associated with effective connectivity of distinct FST circuit elements. Most associations with dorsal striatum [^{18}F]DOPA were limited to the dorsal FST circuit and involved the thalamus. Two thalamic connections—an afferent input from DC and the thalamic self-connection—were also associated with positive symptom severity in FEP-SCZ patients. The modulation of cortical and thalamic glutamatergic signals by striatal dopamine controls striatothalamic filtering and information flow to the cortex, which is thought to play a central role in the pathogenesis of psychosis.[9] The close link between striatal [^{18}F]DOPA and thalamic connectivity that we identify here may be related to evidence that antipsychotics preferentially impact thalamo-striatal and thalamo-cortical functional connectivity in FEP, and can recover FST connectivity following treatment.[81,96]

Bidirectional connectivity between the VTA/SN and DC was associated with dorsal striatal [^{18}F]DOPA. This connection was identified as dysfunctional in SCZ patients and was correlated with positive symptom severity in both SCZ and FEP-SCZ. This consistency of findings suggests a close link between nigrostriatal signaling, striatal dopamine synthesis capacity, and positive symptom severity in patients. Notably, positive symptom severity in FEP was associated with a reduced influence of VTA/SN on DC, whereas higher dorsal striatal [^{18}F]DOPA was associated with an increased afferent influence of VTA/SN. Since patients are known to show elevated [^{18}F]DOPA in the dorsal striatum,[10,11] one might expect a negative association between striatal [^{18}F]DOPA and VTA/SN-DC connectivity. One potential explanation is that elevated dopamine levels lead to compensatory changes in circuit function that affect the optimal balance of signaling between different regions, thus resulting in different circuit mechanisms that regulate dopamine levels in the brains of psychotic and non-psychotic individuals. Concurrent PET-fMRI in both patients and controls would help to test this possibility.

Connections associated with ventral striatal [^{18}F]DOPA were distributed across the ventral and dorsal systems. NAcc projections are widely distributed in the midbrain and drive dorsal striatal dopamine levels.[8] Accordingly, we found that ventral striatal [^{18}F]DOPA was associated with VTA/SN–DC connectivity and inhibition within both regions. VTA/SN–DC connectivity was also disrupted in SCZ patients and associated with positive symptoms in FEP, suggesting a link between the NAcc's regulation of midbrain activity and dorsal circuit dysfunction in patients.[97]

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Limitations and future directions

We used independent samples to cross-sectionally characterize effective dysconnectivity across distinct illness stages. This approach offers a test of consistency across cohorts, but inferences about the progression of circuit dysfunction must be confirmed longitudinally, especially given the differences in sample size and symptom severity across the three cohorts.

Psychosis is a highly heterogenous syndrome and diagnostic differences in our cohort may potentially complicate interpretation of our results. FST and dopamine dysfunction have been implicated in the onset of psychotic symptoms, regardless of the underlying diagnosis. In our past work, we have identified common disruptions of FST functional connectivity across people with FEP,[2] first episode mania with psychosis,[98] at-risk mental state with psychosis,[3] and in association with high levels of subclinical schizotypy.[4,99] Given this evidence for transdiagnostic effects, and that first-episode diagnoses can sometimes be unstable,[100,101] it is critical to examine the FEP cohort as a group. The similarity in findings obtained with the full FEP and FEP-SCZ sub-groups with respect to subcortical and midbrain dysconnectivity and symptom associations indicate that diagnostic heterogeneity does not make a major contribution to our findings.

As the SCZ patients were medicated, the effects of antipsychotics on the brain should also be considered when interpreting the findings.[81] Although we did not find and significant associations between antipsychotic dose and effective connectivity parameters, only limited information on medication exposure was available for this open dataset;[43] therefore, further investigation of the specific effects of antipsychotic treatment on FST effective connectivity is warranted.

Our PEB analysis of the PET data identifies associations with, but not causal influences on, striatal [^{18}F]DOPA. Further investigation of patients would be required to identify precisely how effective dysconnectivity of FST systems leads to dopamine dysregulation in patients. One interesting future avenue of work could use effective connectivity measures to develop multivariate predictive models of symptoms and dopamine synthesis capacity across different cohorts.

We restricted our analysis to left hemisphere FST ROIs previously implicated in past work as showing disrupted functional connectivity or as being key elements of FST circuitry.[2,3] This focus facilitates efficient estimation of DCMs but may miss influences from other key regions extending beyond the circuits studied here, such as the anterior cingulate cortex and anterior insula or even other areas of lateral PFC or thalamus.. Recent improvements in the scalability of DCM[102] may be used to generate more anatomically comprehensive maps of effective dysconnectivity in psychosis.

Conclusions

Our analysis characterized FST effective dysconnectivity in FEP and established schizophrenia. Our findings indicate that subcortical dysfunction features prominently in early illness stages, with cortical abnormalities becoming more apparent later in the illness. In early psychosis, positive symptoms are associated with midbrain connectivity, suggesting that aberrant bottom-up signals emanating from the

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midbrain may present a key FST feature in the pathogenesis of psychotic symptoms. In light of other findings strongly implicating cortical pathology in early illness stages,[103,104] our results suggest that this pathology may emerge after subcortical dysfunction, or that it may arise in cortical areas that were not modelled in our analysis. Nigrostriatal and striatothalamic connectivity are closely linked to striatal dopamine levels, while also being tied to symptom expression in patients. Together, our findings suggest a prominent role for subcortical systems in driving FST dysfunction in psychosis.

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Competing interests

The authors declare no financial relationships with commercial interests.

Supplementary material

Supplementary material is available at *Brain* online.

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Figure legends

Figure 1. Parent model space of fronto-striato-thalamic systems encompassing dorsal and ventral circuits. Left shows anatomical locations of ROIs on axial slices. Right shows the parent model, including 47 biologically plausible connections including self-connections (circular arrows) for a network comprising eight regions. Centroids of ROIs in MNI coordinates (x, y z) are presented in the bottom panel. VTA/SN: ventral tegmental area/substantia nigra (i.e., the midbrain).

Figure 2. Group differences in fronto-striato-thalamic effective connectivity identified in the FEP (*n = 46*), FEP-SCZ (n **= 17), and SCZ (*n = 36*) patients relative to their respective control groups.** Differences in effective connectivity between FEP patients and healthy controls are shown in panel **(A)**, subgroup of FEP patients with schizophrenia-spectrum diagnosis are shown in panel **(B),** and patients with established schizophrenia and healthy controls are shown in panel **(C)**. Boxes show mean connectivity values in each group. For connections between regions, dashed arrows represent connections for which patients show an increased inhibitory influence compared to controls; solid arrows represent connections for which patients show increased excitatory influence compared to controls. For self-connections, dashed arrows represent reduced inhibition and solid arrows indicate increased inhibition in patients compared to controls. Gray arrows represent modeled connections that were not significantly different from the prior. Inter-regional connectivity parameters are in Hz. Self-connection values are in negative log scale. Connections were thresholded at Pp > 0.95 which represents strong evidence. DC: dorsal caudate; NAcc: nucleus accumbens: VTA/SN: ventral tegmental area/substantia nigra (midbrain).

* denotes connections for which the mean in each group has low posterior model probability, but in which differences between groups surpassed Pp > 0.95.

Figure 3. Associations between positive symptoms and fronto-striato-thalamic effective connectivity parameters in the FEP (*n = 46*), FEP-SCZ (*n* = 17), and SCZ (*n = 36*) patients. Panels from left to right depict the results of PEB models mapping associations between FST effective connectivity and positive symptoms in **(A)** FEP, **(B)** FEP-SCZ, and **(C)** SCZ patients. For between-region connections, solid arrows denote positive associations and dashed arrows depict negative associations between effective connectivity parameters and symptoms. For self-connections, solid arrows represent positive associations between symptom severity and self-inhibition, whereas dashed arrows denote negative associations such as more severe symptoms were associated with reduced inhibition. Gray arrows show modelled associations that were not (significantly) different from the prior. Connections were thresholded at Pp > 0.95. DC: dorsal caudate; NAcc: nucleus accumbens: VTA/SN: ventral tegmental area/substantia nigra (midbrain).

Figure 4. Associations between fronto-striato-thalamic connectivity and striatal dopamine synthesis capacity. Panel **(A)** depicts connections associated with dopamine synthesis in the dorsal circuit. Panel

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(B) illustrates associations with dopamine synthesis in the ventral circuit. Solid arrows: positive associations between effective connectivity parameters and dopamine synthesis; dashed arrows: negative associations between effective connectivity parameters and dopamine synthesis; gray arrows: modelled connections that were not (significantly) different from the prior. For self-connections, solid arrows represent positive associations between striatal dopamine synthesis and self-inhibition, whereas dashed arrows denote negative associations between striatal dopamine synthesis and self-inhibition. Connections were thresholded at Pp > 0.95. DC: dorsal caudate; NAcc: nucleus accumbens: VTA/SN: ventral tegmental area/substantia nigra (midbrain).

Table 1 Participants Descriptive Statistics

	FEP			SCZ		[^{18}F]DOPA
	HC (*n* = 23)	**FEP (*n* = 46)**	**FEP-SCZ (*n* = 17)**	**HC (*n* = 100)**	**SCZ (*n* = 36)**	**Healthy (*n* = 33)**
Males, N (%)	14 (60.87)	20 (43.48)	9 (41.18)	55 (55)	26 (72.22)	15 (45.45)
Age, mean (SD)	21.74 (1.92)	19.12 (2.97)	20.51 (2.87)	30.6 (8.87)	35.81(8.49)	22.30 (2.21)
Age range	18–26	15–25	15–25	21–50	22–49	18–28
BPRS Total						
Mean (SD)	–	56.91 (10.34)	61.94 (9.41)	–	49.39 (14.03)	–
Range	–	38–80	46–80	–	26–77	–
BPRS Positive						
Mean (SD)	–	15.65 (4.19)	17.94 (4.31)	–	14.11 (6.17)	–
Range	–	10–27	13–27	–	5–29	–
BPRS Negative						
Mean (SD)	–	5.76 (2.6)	7 (3.16)	–	5.14 (2.54)	–
Range	–	3–12	3–12	–	3–11	–

FEP: first-episode psychosis group; FEP-SCZ: first-episode psychosis subgroup with a diagnosis of schizophrenia spectrum disorder; SCZ: schizophrenia group; HC: healthy controls; [^{18}F]DOPA: healthy group who underwent a simultaneous PET-fMRI protocol

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Table 2 Summary of Group Differences in Fronto-striato-thalamic Effective Connectivity in FEP (*n* = *46*), FEP−SCZ (*n* = 17), and SCZ (*n* = *36*) Patients Respective to Healthy Controls

Connection	Increased (+) or decreased (−) in patients	Excitatory (E) or inhibitory (I) in patients	Effect size (Hz)	90% Posterior Confidence Interval (lower bound, upper bound)	Patients (mean)	Controls (mean)
FEP vs HCs						
dlPFC → dlPFC	+	E	*0.10*	*0.03, 0.20*	*0.64*	*0.44*
Thal → NAcc	−	I	0.10	−0.20, −0.01	−0.12	0.09
Amyg → NAcc	+	E	0.11	0.02, 0.21	0.00	−0.24
VTA/SN → VTA/SN	−	*I*	*0.14*	*−0.24, −0.04*	*−0.04 (−0.48)*	*0.00 (−0.05)*
FEP−SCZ vs HCs						
Thal → NAcc	−	I	0.13	−0.24, −0.03	−0.13	0.09
Amyg → NAcc	+	E	0.13	0.02, 0.23	0.00	−0.25
Amyg → Amyg	−	*I*	*0.11*	*−0.22, 0.00*	*−0.16 (−0.43)*	*0.00 (−0.05)*
NAcc → Hipp	−	I	0.15	−0.26, −0.04	−0.28	0.00
VTA/SN→ Hipp	−	I	0.09	−0.20, −0.02	−0.14	0.00
VTA/SN→ NAcc	+	E	0.09	−0.01, 0.20	0.00	−0.14
VTA/SN→ VTA/SN	−	*I*	*0.08*	*−0.19, 0.03*	*−0.06* (−0.47)*	*0.07* (−0.54)*
SCZ vs HCs						
dlPFC → Thal	−	I	0.11	−0.19, −0.03	−0.14	0.00
vmPFC → dlPFC	−	I	0.08	−0.16, 0.00	−0.06*	0.06*
Thal → Hipp	−	I	0.10	−0.18, −0.02	−0.20	0.00
Thal → NAcc	−	I	0.09	−0.17, −0.01	−0.11	0.00
Amyg → Hipp	−	E	0.09	−0.17, −0.01	0.00	0.16
VTA/SN → VTA/SN	−	*I*	*0.16*	*−0.25, −0.07*	*−0.26 (−0.39)*	*−0.09 (−0.46)*
VTA/SN → DC	−	I	0.14	−0.22, −0.06	−0.19	0.00
DC → VTA/SN	+	E	0.16	0.08, 0.25	0.12	−0.11

FEP: first−episode psychosis group; FEP−SCZ: first−episode psychosis subgroup with a diagnosis of schizophrenia spectrum disorder; SCZ: schizophrenia group; HC: healthy controls. Only connections surpassing a posterior probability threshold of 0.95 are presented in this table, and all reported connections have the posterior probability value of 1.00. All parameters for between region connections are in Hz. Self−connections are italicized, and values are log−transformed to ensure prior negativity (i.e., inhibitory) constraints on self−connections. A positive value for self−connection denotes increased inhibition, a negative value signifies reduced inhibition. Self−connection parameters in Hz are displayed in parentheses.

*Denotes connections for which the mean in each group has low posterior model probability, but in which differences between groups surpassed Pp > 0.95.

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Table 3 Summary of Connections Associated with Severity of Positive Symptoms in FEP (*n* = *46*), FEP-SCZ (*n* = 17), and SCZ (*n* = *36*) Patients

Connection	Positive (+) or negative (-) association	Effect size (Hz)	90% Posterior Confidence Interval (lower bound, upper bound)
Positive symptoms			
FEP			
dlPFC → vmPFC	−	0.10	−0.20, 0.01
vmPFC → vmPFC	−	*0.16*	*−0.27, −0.06*
Amyg → Amyg	−	*0.08*	*−0.19, 0.02*
Hipp → VTA/SN	+	0.10	0.00, 0.20
NAcc → VTA/SN	+	0.09	−0.01, 0.20
VTA/SN → NAcc	+	0.09	−0.02, 0.20
VTA/SN → Hipp	−	0.17	−0.27, −0.06
VTA/SN → Amyg	+	0.14	0.04, 0.25
VTA/SN → DC	−	0.08	−0.19, 0.02
FEP−SCZ			
vmPFC → vmPFC	−	*0.19*	*−0.33, −0.05*
dlPFC → dlPFC	−	*0.09*	*−0.23, 0.04*
dlPFC → Thal	+	0.11	−0.02, 0.24
Thal → Thal	+	*0.13*	*0.00, 0.26*
Hipp → Amyg	−	0.15	−0.28, −0.02
NAcc → Hipp	−	0.09	−0.22, 0.03
NAcc → VTA/SN	+	0.12	−0.01, 0.25
DC → Thal	−	0.12	−0.25, 0.02
VTA/SN → NAcc	+	0.13	0.01, 0.26
VTA/SN → Hipp	−	0.14	−0.27, −0.01
VTA/SN → Amyg	+	0.10	−0.03, 0.23
VTA/SN → DC	−	0.12	−0.25, 0.01
SCZ			
vmPFC → vmPFC	−	*0.09*	*−0.20, 0.03*
vmPFC → Hipp	+	0.16	0.05, 0.27
vmPFC → VTA/SN	+	0.08	−0.03, 0.20
dlPFC → dlPFC	+	*0.09*	*−0.03, 0.21*
dlPFC → Thal	−	0.18	−0.24, −0.06
Thal → Hipp	+	0.11	0.00, 0.22
Thal → VTA/SN	−	0.13	−0.24, −0.02
Hipp → vmPFC	−	0.09	−0.21, 0.02
DC → Thal	+	0.08	−0.03, 0.19
VTA/SN → VTA/SN	−	*0.15*	*−0.27, −0.04*
VTA/SN → Amyg	−	0.10	−0.21, 0.01
VTA/SN → dlPFC	+	0.11	0.00, 0.21

FEP: first-episode psychosis group; FEP-SCZ: first-episode psychosis subgroup with a diagnosis of schizophrenia spectrum disorder; SCZ: schizophrenia group

Only connections surpassing a posterior probability threshold of 0.95 are presented in this table, and all reported connections have the posterior probability value of 1.00.

All parameters for between region connections are in Hz. Self-connections are italicized, and values are log-transformed to ensure prior negativity (i.e., inhibitory) constraints on self-connections. A positive value for self-connection denotes increased inhibition, a negative value signifies reduced inhibition. Positive symptoms measured with BPRS Positive subscale.

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Table 4 Summary of Connections Associated with Striatal Dopamine Synthesis

Connection	Positive (+) or negative (−) association	Effect size (Hz)	90% Posterior Confidence Interval (lower bound, upper bound)
Dorsal Striatum			
Thal → Thal	+	*0.10*	*−0.04, 0.23*
Thal → DC	+	0.09	−0.04 0.22
Thal → VTA/SN	+	0.12	−0.01, 0.26
Amyg → vmPFC	+	0.08	−0.05, 0.21
Amyg → Thal	+	0.11	−0.02, 0.24
DC → Thal	−	0.13	−0.26, 0.00
Ventral Striatum			
Hipp → Thal	+	0.16	0.03. 0.30
Hipp → VTA/SN	+	0.16	0.03, 0.29
Amyg → Thal	−	0.12	−0.25, 0.02
NAcc → Thal	−	0.11	−0.24, 0.02
Amyg → Amyg	−	*0.14*	*−0.27, 0.01*
NAcc → NAcc	−	*0.14*	*−0.28, 0.00*
DC → VTA/SN	−	0.12	−0.25, 0.01
DC → DC	+	*0.10*	*−0.03, 0.22*
VTA/SN → VTA/SN	+	*0.10*	*−0.05, 0.25*
VTA/SN → vmPFC	+	0.15	0.02, 0.29
VTA/SN → DC	+	0.12	−0.01, 0.25

Only connections surpassing a posterior probability threshold of 0.95 are presented in this table, and all reported connections have the posterior probability value of 1.00.

All parameters for between region connections are in Hz. Self-connections are italicized, and values are log-transformed to ensure prior negativity (i.e., inhibitory) constraints on self-connections. A positive value for self-connection denotes increased inhibition, a negative value signifies reduced inhibition.

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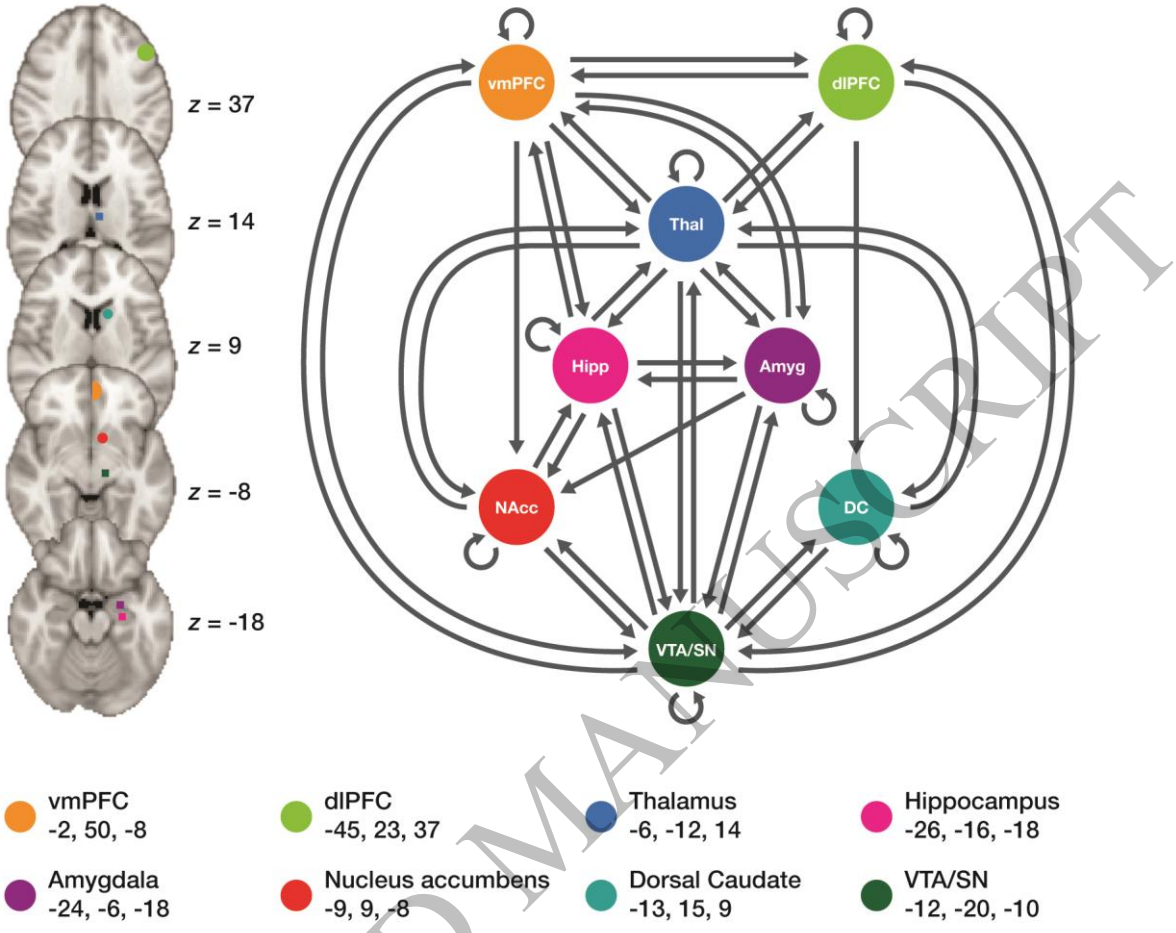


Figure 1
156x134 mm (71 x DPI)

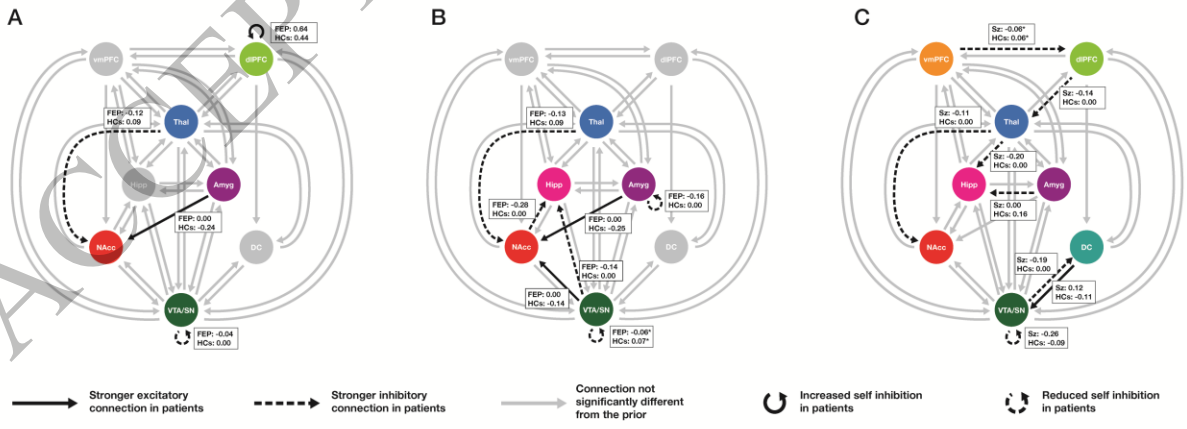


Figure 2
319x119 mm (71 x DPI)

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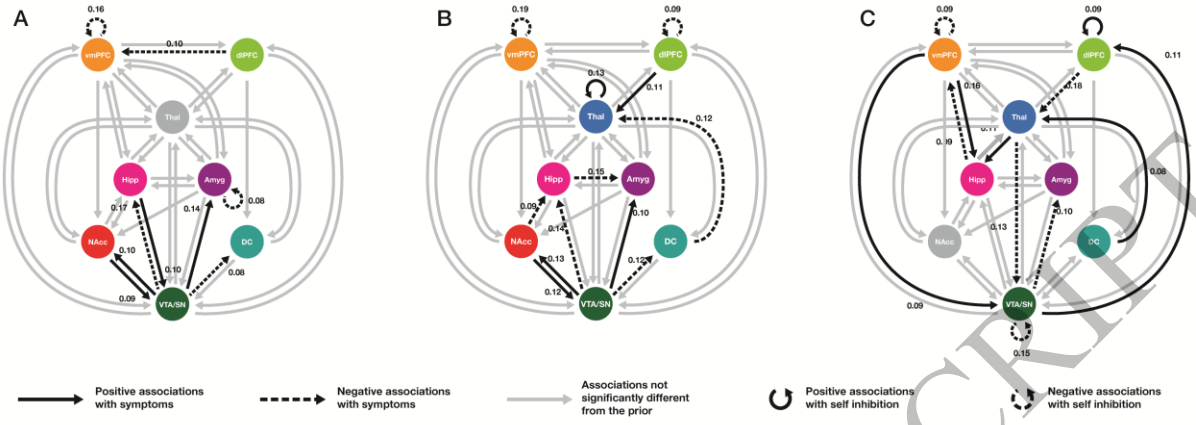


Figure 3
319x119 mm (71 x DPI)

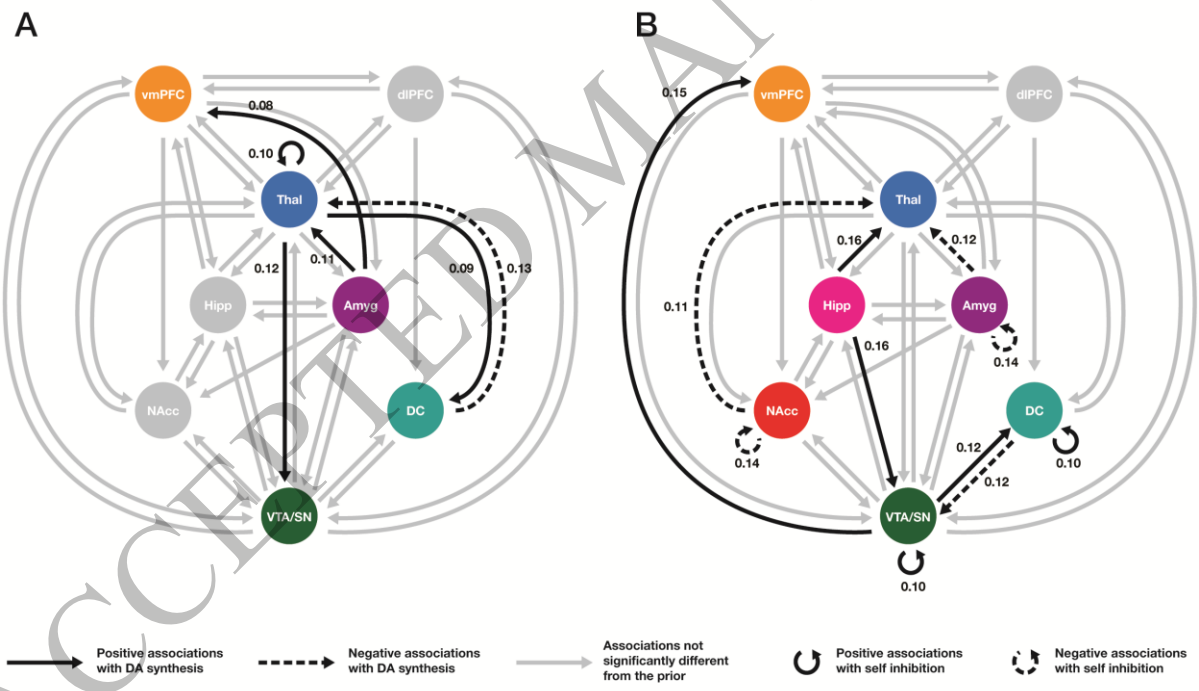


Figure 4
210x127 mm (71 x DPI)

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