Anterior limb of the internal capsule tractography: relationship with capsulotomy outcomes in obsessive-compulsive disorder

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ABSTRACT

Objectives  Surgical procedures targeting the anterior limb of the internal capsule (aLIC) can be effective in patients with selected treatment-refractory obsessive-compulsive disorder (OCD). The aLIC consists of white-matter tracts connecting cortical and subcortical structures and show a topographical organisation. Here we assess how aLIC streamlines are affected in OCD compared with healthy controls (HCs) and which streamlines are related with post-capsulotomy improvement.

Methods  Diffusion-weighted MRI was used to compare white-matter microstructure via the aLIC between patients with OCD (n=100, 40 women, mean of age 31.8 years) and HCs (n=88, 39 women, mean of age 29.6 years). For each individual, the fractional anisotropy (FA) and streamline counts were calculated for each white-matter fibre bundle connecting a functionally defined prefrontal and subcortical region. Correlations between tractography measures and pre-capsulotomy and post-capsulotomy clinical outcomes (in obsessive-compulsive, anxiety and depression scores 6 months after surgery) were assessed in 41 patients with OCD.

Results  Hierarchical clustering dendograms show an aLIC organisation clustering lateral and dissociating ventral and dorsal prefrontal–thalamic streamlines, findings highly relevant to surgical targeting. Compared with HCs, patients with OCD had lower aLIC FA across multiple prefrontal cortical–subcortical regions (p<0.0073, false discovery rate-adjusted). Greater streamline counts of the dorsolateral prefrontal–thalamic tracts in patients with OCD predicted greater post-capsulotomy obsessive-compulsive improvement (p=0.016). In contrast, greater counts of the dorsal cingulate–thalamic streamlines predicted surgical outcomes mediated by depressive and anxiety improvements.

Conclusions  These findings shed light on the critical role of the aLIC in OCD and may potentially contribute towards precision targeting to optimise outcomes in OCD.

INTRODUCTION

The pathophysiology of obsessive-compulsive disorder (OCD) implicates cortico–striato–thalamic networks1 2 whose tracts course through the anterior limb of the internal capsule (aLIC), a key white-matter structure in this network.14 Diffusion tensor imaging (DTI) studies in OCD have shown heterogeneous findings but suggest white-matter abnormalities in patients with OCD in the cingulate bundle, the corpus callosum and the aLIC.15-19

Capsulotomy and deep brain stimulation targeting the aLIC and subththalamic nucleus (STN) can be effective in alleviating the clinical symptoms of severe medical-refractory patients with OCD.10-13 Capsulotomy is a neurosurgical procedure involving ablation with radiotherapy, gamma knife or high-frequency ultrasound of the aLIC, and reduces OCD symptoms by an average of a little over 50% with a range of 37%–80%.14 15 A single-shot ablative lesion targeting the mid-aLIC is less effective than a further lesion expansion using ‘double-shot’ or ‘triple-shot’ techniques which include the ventral aLIC.12 16 Thus, clinical outcome studies tracking the effects of ventral–dorsal lobar lesions are crucial for assessing which tracts through the aLIC connecting cortical and subcortical structures may be most clinically relevant in predicting capsulotomy outcomes.

Converging evidence from primate tract tracing and human DTI studies shows that different regions of the aLIC carry fibres from different prefrontal regions with a ventral–dorsal, medial–lateral and anterior–posterior organisational topography.1 4 Imaging studies have highlighted specific tracts and connections for predicting postoperative clinical outcomes. A preoperative tractography study following ventral internal capsule deep brain stimulation (VIC DBS) showed that the most clinically effective contacts stimulated thalamo–cortical aLIC fibres to the dorsolateral prefrontal cortex (dIPFC), rostral dorsal anterior cingulate (dACC) and mesial PFC.17 In a second study comparing preoperative tractography in VIC and STN DBS for OCD, similar aLIC fibre streamlines were identified to the rostral cingulate and mesial PFC across both DBS targets as predictive of optimal clinical outcome.18 Furthermore, a predictive resting state functional MRI (fMRI) study of capsulotomy outcomes from our group highlighted functional connectivity between the dorsal cingulate and striatum as predictive of obsessive-compulsive symptom improvement 1 month after surgery.19 These studies investigating the predictive role of imaging are potentially critical in guiding more precise postsurgical targeting.20


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and suggest particular abnormality of white-matter tracts passing through the aLIC may exist in OCD and may also be related with clinical outcome by neurosurgery targeting at aLIC.

The goal of the present study was twofold: first, we identified which cortical–subcortical white-matter streamlines passing through the aLIC were abnormal in OCD. Here, we focused on specific functionally defined prefrontal regions implicated in OCD including dIPFC, orbitofrontal cortex (OFC), ventrolateral PFC (vIPFC) (or inferior frontal cortex), dACC and ventromedial (vmPFC), which were used in our previous studies.19 These cortical regions have been suggested to be impaired in OCD based on convergent cognitive, task-based fMRI studies related to decision-making and impulsivity.12 Here, using a tractographic analysis, we focused on the quantitative characteristics of the white matter streamlines via the aLIC of these prefrontal regions connecting to subcortical regions of striatum, thalamus and STN in OCD. Second, focusing on the regions shown to be predictive of capsulotomy and DBS outcomes, namely the dIPFC and dACC,17–19 we ask in a subset of these patients whether these streamlines are correlated with improvements in obsessive-compulsive, depressive and anxiety symptoms following capsulotomy for severe treatment-refractory OCD.

MATERIALS AND METHODS
Subjects and clinical assessment
Patients with OCD who were considered for neurosurgery at Ruijin Hospital from August 2013 to September 2018 were recruited in the imaging study (n=100). Age and gender-matched healthy controls (HCs) were recruited via community-based advertisements (n=88). OCD diagnosis was confirmed using the Diagnostic and Statistical Manual of Mental Disorders-IV-TR criteria by a psychiatrist specialising in OCD. Exclusion criteria for patients and for age-matched and gender-matched HCs for imaging included: age below 16 or above 65, a history of severe head trauma, substance use disorders, other major psychiatric or major neurological disorders, claustrophobia, and metal or electronic implants. Inclusion criteria and surgical details for capsulotomy have been previously reported.19 Inclusion for capsulotomy included a Yale-Brown Obsessive Compulsive Scale (Y-BOCS) score of more than 25 or one subscale of more than 15; chronicity of greater than 5 years; treatment refractory defined as more than three antidepressant trials of greater than 12 weeks at maximally tolerated doses and adequate failed psychotherapy. Clinical information was collected using the Y-BOCS, Hamilton Anxiety Rating Scale (HAM-A) and the 17-item version of Hamilton Depression Rating Scale (HAM-D).

Capsulotomy and follow-up
Among the patients with OCD (n=100) included in the current study, 78 underwent capsulotomy. Details of the surgical procedure can be found in our previous work19 and are briefly summarised here. A stereotactic frame was mounted on the patient’s head under local anaesthesia, and preoperative MRI was immediately scanned. The target was localised between the anterior and middle thirds of the aLIC. Stereotactic target coordinates and the trajectory angle for the target region were calculated according to stereotactic MRI. The radiofrequency lesions were generated by radiofrequency ablation with a 2 mm diameter electrode at 75°C for 60 s, and a lesion region 4–5 mm in diameter and 10–12 mm in length along the contoured target was produced.

After surgery (n=78), 41 completed assessments at least 6 months after surgery (data included in the analyses related to postsurgical improvement), 11 completed assessments between 1 and 5 months after surgery and 26 either could not be contacted after surgery or were unable to attend a follow-up visit.

Image acquisition
Diffusion-weighted imaging was conducted on a Siemens Tim Trio 3-Tesla scanner (Siemens, Erlangen, Germany) using a 12-channel phased-array head coil. A diffusion spectrum imaging (DSI) sequence was acquired with a total of 515 diffusion directions covering a hemisphere with the maximum b-value of 7000 s/mm² (field of view=192×192 mm², matrix size=80×80, in-plane resolution=2.4 mm, slice thickness=2.4 mm, repetition time=10 100 ms, echo time=152 ms, flip angle=90°, in-plane acceleration factor=2).21 Due to low signal-to-noise ratio (SNR) at high b-values, only 171 directions up to the b-value of 4800 s/mm² were used in further analyses. A T1-weighted image was also collected with a magnetisation prepared rapid gradient-echo sequence (field of view=256×256 mm², matrix size=256×256, in-plane resolution=1 mm, slice thickness=1 mm, repetition time=2300 ms, echo time=3 ms, inversion time=1000 ms, flip angle=9°).

Denoising, diffusion modelling and deterministic tractography
As the diffusion data were DSI rather than DTI with low SNR with very high b-values and in the ventromedial prefrontal regions, we adapted our processing and analytic algorithm specifically for these issues. We used local principal component analysis (local PCA) for denoising of the diffusion-weighted images (https://sites.google.com/site/pierrickcouce/softwares/denosing-for-medical-imaging/dwi-denosing/dwi-denosing-software).24 PCA was performed on a sliding local-block at each voxel over diffusion directions and the data were reconstructed without noisy components as determined by their associated eigenvalues (see online supplemental figure 1 for denoising). Eddy current correction could not be applied as it is not implemented for DSI data in any publicly available software (eg, see https://fsl.fmrib.ox.ac.uk/fsl/fslwiki/eddy/UsersGuide#A--data_is_shelled).

The denoised diffusion data were analysed using DSI Studio (http://dsi-studio.labsolver.org). The SNR, particularly in the frontal areas, was low, thus a high-order model such as generalised q-sampling imaging25 resulted in implausible and spurious results. A lower-order model is less sensitive to such noise because it does not model complex functions and only more general trends. Therefore, we sacrificed the high angular resolution to improve SNR by fitting with a traditional Gaussian tensor with only three orthogonal axes. Based on the tensor models, we used a deterministic fibre tracking algorithm26 to find tract bundles connecting pairs of cortical and subcortical regions of interest (ROIs) seeding from all voxels within the aLIC. The deterministic approach was chosen to find the best estimation with computational efficiency over the probabilistic approach.27 The aLIC was defined using the Harvard-Oxford atlas in FSL (FSL-HO (https://fsl.fmrib.ox.ac.uk/fsl/fslwiki/Atlases)).28 Six cortical ROIs (figure 1) were included: vIPFC, dIPFC, vmPFC, dACC, pre-supplementary motor area (pre-SMA) and OFC. The ROIs of the vIPFC, dIPFC, vmPFC, dACC and pre-SMA were manually defined and used in previous publications from our group (see online supplemental materials in Morris et al for more details on manual definitions).21 22 Five subcortical ROIs (figure 1) were included: thalamus, caudate, putamen, nucleus accumbens (NAcc) and STN. The ROIs of the OFC, thalamus, caudate nuclei, putamen and NAcc were identified using the...
streamlines were estimated from T1-weighted images using the Segment module and the Tissue Volume utility in SPM12 (http://www.fil.ion.ucl.ac.uk/spm).  
In order to assess the effect of group membership or clinical measures while regressing out the effects of covariates, a t-statistic is calculated from \( t = \frac{c \hat{\beta}}{\sqrt{\hat{\sigma}^2 (XX)^{-1}c}} \) where \( c \) is a contrast vector \([0, 0, 0, 1]^T\), \( \hat{\sigma}^2 \) is a sample variance and \( X \) is a design matrix.

Multicollinearity of regressors was assessed using a MATLAB implementation (https://github.com/brian-lau/colldiag). For model (1), CIs did not exceed 11 with only weak collinearity between Y-BOCS baseline scores and TIV (CI=11, variance decomposition proportions (VDPs)=82.5%, 90.1%). For model (2), CIs did not exceed 12 (age and TIV, CI=12, VDPs=96.2%, 96.2%). Overall, there was only weak collinearity between age and TIV below a threshold of 30.

P values were estimated using the Monte Carlo method (100,000 permutations) as implemented in the FieldTrip MATLAB Toolbox (http://www.fieldtriptoolbox.org/), with a false discovery rate (FDR) adjustment for multiple comparisons when necessary. A family-wise alpha level of 0.05 was used.

Tract-based spatial statistics
In order to assess whether our group findings of streamlines within the aLIC were specific to the aLIC or more generalised white matter differences between OCD and HC groups, we also conducted an analysis of local diffusion properties in a voxel-wise fashion using whole-brain tract-based spatial statistics (TBSS) in FSL. FA images were non-linearly registered to the FMRIB58_FA template, skeletonised and thresholded at 0.05. We focused here on FA as our aLIC measures focus on FA. Note that the range of the FA estimates was lower than usual DTI data. Two GLMs (equations 1–2) were non-parametrically tested using threshold-free cluster enhancement, as implemented in FSL, with 10,000 permutations each. Family-wise alpha level of 0.05 was used.

Unless otherwise noted, custom MATLAB (R2019a V9.6.0.1153713; MathWorks, Natick, Massachusetts, USA) functions were used. Streamline visualisation was conducted using DSI-Studio.

RESULTS
Demographic and clinical measures
The patients with OCD (n=100; 31.8 (SD 9.7) years old) and HCs (n=88; 29.6 (SD 8.0) years old) were matched in age (\( t=-1.64, p=0.102 \), gender (OCD female ratio 0.40; HC 0.44; \( \chi^2=0.21; p=0.649 \)) and total intracranial volume (OCD in litres 1.45 (SD 0.13); HC 1.42 (SD 0.14); \( t=-1.55; p=0.122 \)). Clinical measures and obsessive-compulsive symptoms at different time points are reported in table 1 and online supplemental table 1. The 41 patients who underwent capsulotomy and completed assessments at least 6 months after surgery were included in the analyses related to postsurgical improvement. The YBOCS, HAMD and HAMA scores decreased post-capsulotomy (one-sample t-test, \( p<0.0001 \)) (online supplemental figure 3).

TRACTOGRAPHY
General tractography and group comparison
The number of subjects that showed any streamlines varied across connections between cortical and subcortical ROIs (online supplemental table 2). Streamlines were found in more than 90% of subjects (>169) for 12 connections: (i) thalamus–vlPFC, (ii) thalamus–dlPFC, (iii) thalamus–dACC, (iv)
streamlines was similar between OCD and HC groups (χ² < 90% of subjects), the proportion of subjects who showed

tical regions. For each connection (including those found in

illustration of the streamlines between cortical and subcor-

spatially segregated within the aLIC, an important question

cant findings and hence are not reported here.

SMA.

Ham-D or NC (FA or NC) and baseline clinical severity (ie, baseline YBOCS/

also assessed the relationship between streamline measures

pre-

(vii) putamen–vlPFC, (viii) putamen–dlPFC, (ix) putamen–

pre-

SMA, (v) thalamus–OFC, (vi) caudate–dlPFC,

thalamus–pre-SMA, (v) thalamus–OFC, (vi) caudate–dlPFC,

(vii) putamen–vlPFC, (viii) putamen–dlPFC, (ix) putamen–

pre-

SMA. See figure 2 for averaged track density images and an illustration of the streamlines between cortical and subcor-

regions. For each connection (including those found in

<90% of subjects), the proportion of subjects who showed streamlines was similar between OCD and HC groups (χ²
tests; p>0.4395, uncorrected; p>0.9773, FDR-adjusted). We also assessed the relationship between streamline measures (FA or NC) and baseline clinical severity (ie, baseline YBOCS/ Ham-D/Ham-A scores) or obsessive-compulsive symptom types (online supplemental table 1); there were no significant

findings and hence are not reported here.

We then asked whether the identified aLIC streamlines were spatially segregated within the aLIC, an important question relevant to surgical targeting. Across all subjects, spatial distributions of the thalamocortical streamlines within the aLICs were consistent with previous studies.3 4 See figure 2 for thalamocortical streamlines from a representative subject. To quanti-

fy the spatial distribution, the centre of gravity in the aLICs in the normalised track-density images was compared between thalamocortical connections (online supplemental table 3). See figure 4 for individual centres of gravity in the aLICs. Subjects with all five thalamocortical connections were used (n=156). Streamlines connecting dorsal and ventral structures were separated. One-way multivariate analysis of variance revealed clear separation in coordinates between cortical targets in the 0-th dimension (ie, group means are not at the same point) and the 1st dimension (ie, group means are not in a straight line) (left, p<0.0001, p<0.0001, p=0.0645; right, p<0.0001, p<0.0001, p=0.3212 for each dimension). Using hierarchical clustering dendrograms based on shortest linkage, bilaterally, the lateral PFC regions are clustered first, followed by clustering with OFC, then dACC and finally pre-SMA (figure 4).

Using model (1), we found significant decreases in FA of streamlines in patients with OCD in all 12 connections tested (p<0.0073, FDR-adjusted) with no differences in NC (p>0.7762, FDR-adjusted) compared with HCs (figure 5).

Relationship between tractography and capsulotomy outcomes

Using model (2), we found significant correlations between clinical improvement after capsulotomy and NC in the thalamocortical connections of the dIPFC and dACC (r=0.34, p<0.0436). Specifically, Y-BOCS improvements (ie, decreases in Y-BOCS at follow-up compared with baseline) were positively correlated with NC of thalamus–dIPFC (r=0.36,

Figure 2  Maximal intensity projection of averaged track-density images. Colour coding is scaled proportionally to the maximal density. Cortical (orange) and subcortical (blue) ROIs and aILCs (grey) are marked in contours. Regions of interest. Caud, caudate nucleus; dACC, dorsal anterior cingulate cortex; dIPFC, dorsolateral prefrontal cortex; OFC, orbitofrontal cortex; pre-SMA, anterior portion of supplementary motor area; Put, putamen; NAcc, nucleus accumbens; STN, subthalamic nucleus; Thal, thalamus; vlPFC, ventrolateral prefrontal cortex; vmPFC, ventromedial prefrontal cortex.

Figure 3  Thalamocortical streamlines in the anterior limb of the internal capsule (aLIC). Streamlines of a representative subject are rendered with colour-coding for cortical targets (purple: pre-supplementary motor area (pre-SMA), yellow: dorsal cingulate (dACC), orange: dorsolateral prefrontal cortex (dIPFC), blue: ventrolateral prefrontal cortex (vlPFC), green: orbitofrontal cortex (OFC)). Isosurfaces of thalamus (pale brown) and the aILCs (white) are also shown. (A) Front view (radiological orientation), (B) oblique view (from the left), (C) lateral view (from the right).

Table 1  Clinical measures of patients with OCD

<table>
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<th></th>
<th>n</th>
<th>Mean</th>
<th>SD</th>
<th>Min</th>
<th>Max</th>
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<td>Duration of illness (year)</td>
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<td>10.3</td>
<td>5.8</td>
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<td>25</td>
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<td>Age at onset of illness (year)</td>
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<td>21.0</td>
<td>8.6</td>
<td>7</td>
<td>53</td>
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<td>Y-BOCS (baseline: all patients with OCD)</td>
<td>96</td>
<td>30.4</td>
<td>6.7</td>
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<td>40</td>
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<td>Y-BOCS (baseline: ≥6-month follow-up patients)</td>
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<td>30.1</td>
<td>7.1</td>
<td>15</td>
<td>40</td>
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<tr>
<td>Y-BOCS (follow-up: ≥6-month follow-up patients)</td>
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<td>12.7</td>
<td>12.0</td>
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<td>38</td>
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<tr>
<td>Y-BOCS follow-up time after surgery (month)</td>
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<td>25.8</td>
<td>23.0</td>
<td>6</td>
<td>91</td>
</tr>
<tr>
<td>Ham-A (baseline: all patients with OCD)</td>
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<td>17.8</td>
<td>10.0</td>
<td>0</td>
<td>46</td>
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<td>9.2</td>
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<td>8.5</td>
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<td>25.8</td>
<td>23.0</td>
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<td>Ham-D follow-up time after surgery (months)</td>
<td>38</td>
<td>23.3</td>
<td>19.4</td>
<td>6</td>
<td>69</td>
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</tbody>
</table>

The 41 patients who underwent capsulotomy and were clinically assessed at least 6 months after surgery were included in the follow-up. Ham-A, Hamilton Anxiety Rating Scale; Ham-D, Hamilton Depression Rating Scale; n, number of available data points out of 100 patients with OCD; Y-BOCS, Yale–Brown Obsessive Compulsive Scale.
p=0.0347) and thalamus–dACC (r=0.34, p=0.0402) but not with FA (figure 6). We then repeated model (2) with the covariates of time after surgery and changes in scores of depression and anxiety to assess the impact of these factors on our findings. We again show a positive correlation with YBOCS improvement with the NC of thalamus–dlPFC (p=0.016) but the thalamus–dACC results were no longer significant with inclusion of either depression or anxiety changes.

TBSS: whole brain analysis
We further investigated whether these aLIC findings were due to a decrease of FA specifically in the aLIC or whole brain white matter in general, or whether they were specific for thalamo–cortical/striato–cortical streamlines using whole brain TBSS analysis. From the two models (equations 1–2), we only found significant group differences in forceps minor, superior longitudinal fasciculi, and other cerebral and subcortical white matter (p<0.05, threshold free cluster enhancement (TFCE)-corrected). See figure 7 and online supplemental table 4 for significant clusters. We also found decreased FA in a posterior tip of the left aLIC, but very few streamlines pass through this region (0.24%±0.51% of maximal track-density), except thalamus–pre-SMA (25.52%), suggesting that the cluster is at the border as a part of the thalamus–pre-SMA connectivity and does not overlap with other streamlines. There was no correlation between TBSS and clinical outcome measures. Therefore, the TBSS analysis confirmed that the tractography was not confounded by the common seed (ie, aLIC) but specifically quantified structural connectivity of the thalamo–cortical/striato–cortical tracts.

Figure 4
Centres of gravity and comparison of thalamo–cortical streamlines through the anterior limb of the internal capsule (aLIC) tracts. (A) A rear view of the left aLIC, (B) rear view of the right aLIC, (C) a top view of the left aLIC, (D) a top view of the right aLIC. Each circle represents a centre of gravity of one subject (total n=156) for each thalamocortical connection (purple: pre-SMA, yellow: dACC, orange: dlPFC, blue: viPFC, green: OFC). Montreal Neurological Institute (MNI) coordinates are marked. (E) A dendrogram of a hierarchical clustering using the single linkage method based on the distances among clusters estimated by a multivariate analysis of variance (MANOVA) model in the left aLIC. (F) A dendrogram for the right aLIC. P values from one-way MANOVA are marked. dACC, dorsal anterior cingulate cortex; dlPFC, dorsolateral prefrontal cortex; OFC, orbitofrontal cortex; pre-SMA, anterior portion of supplementary motor area; viPFC, ventrolateral prefrontal cortex.

Figure 5
Group differences in fractional anisotropy in all connections tested (A–L). HC, healthy control, OCD, obsessive–compulsive disorder. P values are adjusted by false discovery rate (FDR). dACC, dorsal anterior cingulate cortex; dlPFC, dorsolateral prefrontal cortex; OFC, orbitofrontal cortex; pre-SMA, anterior portion of supplementary motor area; STN, subthalamic nucleus; viPFC, ventrolateral prefrontal cortex.

Figure 6
Correlation between connectivity measures and obsessive–compulsive symptom improvement. (A–B) Correlation between normalised streamline counts (NC) of the dorsolateral prefrontal cortex and thalamus (Thal–dlPFC) and dorsal cingulate and thalamus (Thal–dACC) and improvements of Yale Brown Obsessive Compulsive Symptom (Y-BOCS) scores following capsulotomy (postsurgery and presurgery). (C) Correlation between NC of the thal–dlPFC tracts with the inclusion of clinical variables including improvements in anxiety and depression scores following capsulotomy as covariates. dACC, dorsal anterior cingulate cortex; dlPFC, dorsolateral prefrontal cortex; Thal, thalamus.

Figure 7
Voxel-wise group differences in fractional anisotropy (FA). Using tract-based spatial statistics, FA values were compared and thresholded at p<0.05 (TFCE corrected). Signed p values are visualised in a log-scale for a contrast of patients with obsessive–compulsive disorder (OCD) minus healthy controls (HCs). Reddish colours mark increases while bluish colours mark decreases in OCDs compared with HCs. Green contours mark anterior limbs of internal capsules.
DISCUSSION

In this large-scale diffusion-weighted study, we focus on prefrontal–subcortical streamline pathways coursing through the aLIC highlighting differences between OCD subjects and healthy controls and showing a potential relationship with capsulotomy outcomes. We first show that compared with healthy controls, OCD was associated with lower FA in the aLIC across multiple prefrontal cortico-thalamic and cortico-striatal streamlines, namely bilateral dlPFC, vlPFC, dorsal cingulate, pre-SMA, OFC and vmPFC. Similarly, lower prefrontal–subthalamic streamline counts from dlPFC, vlPFC and pre-SMA were observed possibly reflecting hyper-direct projections. Critically, these findings may be specific to these prefrontal–subcortical streamline pathways coursing through the aLIC as whole-brain TBSS showed the aLIC in OCD to be no different from healthy controls, except in a posterior region with limited prefrontal–subcortical tracts. Instead, the TBSS analysis highlighted lower FA in various prefrontal regions including the connection between frontopolar regions (forceps major), streamlines within insular and middle frontal gyrus, and the long-range superior longitudinal fasciculus linking posterior and anterior cerebral regions of parieto-occipital and fronto-temporal cortices.

Critically, we highlight a relationship between presurgical aLIC tractography and capsulotomy outcomes with higher streamline count in the dlPFC–thalamic tracts predicting greater obsessive-compulsive symptom improvement post-capsulotomy. We show a potential dissociation between improvements in obsessive-compulsive symptoms as compared with anxious-depression symptoms with greater streamline count in the dorsal cingulate–thalamic tracts predicting surgical outcome but likely mediated by greater improvements in anxiety or depressive symptomatology, features commonly comorbid in OCD.

Anatomical organisation of prefrontal–thalamic tracts in the aLIC

Across all subjects, we show an anterior–posterior ventral–dorsal axis organisation demonstrating clustering of lateral prefrontal tracts and dissociating from ventral and dorsal prefrontal tracts. These findings have relevance to potential neurosurgical targeting influencing differing prefrontal regions. A DTI study in primates and human subjects that segmented human aLIC into five regions based on the positions of axons from different cortical regions revealed that fibres from vmPFC, dorsomedial PFC, dACC, vlPFC and dlPFC have some degree of topographical arrangement within the aLIC. This finding is homologous with rodent studies demonstrating that descending PFC fibres in the form of white-matter fascicles embedded within the striatum are arranged topographically.

Here, we confirm a statistically significant topographical arrangement based on functionally defined prefrontal regions and further show that within the aLIC a hierarchical clustering anatomical organisation: the lateral prefrontal cortices appear to cluster together, followed by OFC, then dACC and finally pre-SMA. This clustering arrangement has implications for DBS targeting and contact stimulation or more precise ablative lesioning within the aLIC. For instance, our findings suggest that a ventral anterior target (volume of tissue activation or lesion) of the aLIC is likely to simultaneously influence both dorsolateral and ventrolateral prefrontal cortices and may be difficult to dissociate in the context of precision targeting. Greater extension in the aLIC may then involve OFC with a more ventrally and anteriorly extension or contact then influencing dACC and pre-SMA.

Dorsolateral prefrontal cortex: capsulotomy outcomes

Our findings highlight a relationship between dlPFC and capsulotomy outcome: OCD subjects with higher dlPFC–thalamic streamline counts have better obsessive-compulsive symptom improvement outcomes after capsulotomy controlled for anxiety and depression scores. This finding highlighting the dlPFC tracts is consistent with observations from DBS studies. Obsessive-compulsive symptomatology improvement after VIM DBS in OCD was predicted by DBS contacts specifically targeting dlPFC tracts. The effect of VIM DBS has also been suggested to be partly explained by the enhancement of cognitive control involving prefrontal regions including the dlPFC. Furthermore, in a DBS study targeting the anterior limbic-associative STN in OCD, stimulation was associated with an improvement in reflection impulsivity correlating with dlPFC and STN resting state functional connectivity.

Here we show that FA of the dlPFC is lower in patients with OCD relative to healthy controls, but no differences were observed in streamline counts. These FA findings are consistent with observations of impairments of the dlPFC in OCD particularly with impaired functioning in goal-directed control, working memory, strategic planning and reflection impulsivity. However, the count of dlPFC–thalamic streamlines was not a marker for the differences between OCD relative to healthy controls nor a marker of baseline severity. As the location of capsulotomy is highly likely to target these dlPFC-tracts, those with a greater baseline number of streamlines may have a greater sensitivity and capacity for Postsurgical plasticity or reorganisation and hence symptom improvement. The final common mechanisms underlying stimulation and reorganisation post-capsulotomy might overlap, suggesting that the thalamic-dlPFC tracts may be important for optimal clinical outcome. Further longitudinal studies are required to address these specific questions.

Dorsal cingulate: capsulotomy outcomes

We further show that greater counts of thalamic-dorsal cingulate streamlines are also related with capsulotomy outcomes but suggest this may be mediated by improvements in anxiety and depressive symptoms. The dorsal cingulate is commonly implicated in OCD pathophysiology involving enhanced error monitoring with a greater error related negativity and greater aversive processing.

The dorsal cingulate, and more specifically the rostral cingulate and mesial prefrontal cortex at the genu of the corpus callosum was implicated in a tractography study predicting the clinical improvement of obsessive-compulsive symptoms for both DBS targeting the VIC and the STN. Cingulotomy targeting the cingulum bundle has also shown efficacy in alleviating treatment refractory OCD. We have previously shown that resting state functional connectivity of the dorsal cingulate and striatum was decreased in patients with OCD following capsulotomy and that connectivity predicted improvement of obsessive-compulsive scores 1 month post-capsulotomy. We have also shown normalisation and improvement of postoperative glucose metabolism in dorsal cingulate, OFC, caudate and thalamus following OCD capsulotomy in eight patients, regions which were hypometabolic relative to healthy controls. Our current finding differs from these previous predictive studies in highlighting the role of depressive and anxiety symptoms which may be related to our inclusion of anxiety assessments, differences between neurosurgical procedures (DBS and capsulotomy) or methodological and clinical outcomes (resting state fMRI 1 month after...
surgery as compared with tractography 6 months after surgery). These findings focusing on depressive improvements are consistent with a dual stimulation study in OCD demonstrating that tracts from the VIC DBS target were associated with the medial OFC and vmPFC linked with improving depressive symptoms whereas STN DBS tracts were associated with other prefrontal regions linked with improving behavioural flexibility. Further studies focusing on linking cognitive processes and anatomical targets on postsurgical outcomes are indicated.

Limitations and summary
This study is not without limitations. First, we note that specific conventional DTI processing steps such as Eddy current correction are not appropriate for DSI data. Second, as the frontal SNR was low, a high-order model such as generalised q-sampling imaging resulted in implausible results. Therefore, we limited ourselves to a traditional Gaussian tensor model, which is more robust to noise. Third, even though, poor SNR may still have limited the identification of these tracts to thalamic or subthalamic nuclei; in some cases of subcortical–cortical pairs, no streamline was found in some subjects. Thus, we only performed group-wise analyses for those ROI pairs, for which streamlines revealed in more than 90% of subjects. Alternative ROIs using Brodmann Areas were considered but did not result in more streamlines that allows comparisons between all possible subcortico–cortical pairs. This limitation excluded many of interesting subcortical regions such as NAcc and medial/lateral OFC from the current analyses. In exploration, we conducted a separate analysis of the vmPFC and NAcc (see online supplemental materials and figure S) without constraining within the aLIC ROI. The streamlines were found in 89% of subjects and consistent with the anatomy of vmPFC tracts but did not necessarily pass through the aLIC ROI. Data with higher SNR are necessary to enable more comprehensive analysis. Fourth, the current study was based on the assumption that the aLIC regions are similarly affected in patients who underwent capsulotomy. This assumption should be confirmed by post-capsulotomy images, which are still in the process. However, in a separate study, we reported the relationship between OCD improvement and the lesion size and location using structural images acquired 1–5 months after surgery (see online supplemental figure S for the overlap between the aLIC seed masks in the current study and the lesion probability map from a subset of patients from the separate study). A coming study will be able to stratify patients of treatment targets for both invasive and non-invasive neuro-modulation across other psychiatric disorders.

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Contributors
CZ, S-GK, BS, ZW and VV contributed to the conception and design of the study. CZ, S-GK, YZ, QL, KZ, HG, HW and WL contributed to acquisition, post-processing and analysis of the data. CZ, S-GK, JL, ZW and VV drafted the text and prepared the figures. All authors approved the final version of the manuscript.

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Competing interests
None declared.

Patient consent for publication
Not required.

Ethics approval
All procedures were approved by the ethics committee of Ruijin Hospital Shanghai Jiao Tong University School of Medicine (approval number: 2011R4). All subjects were fully informed regarding experimental procedures and provided written consent in accordance with the Declaration of Helsinki.

Provenance and peer review
Not commissioned; externally peer reviewed.

Data availability statement
Data are available upon reasonable request. The MRI datasets generated and analysed during the current study are not publicly available due to privacy regulations of patient data, but are available from the corresponding author upon reasonable request.

Supplemental material
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Subthalamic nucleus stimulation in severe obsessive-compulsive disorder.


Li N, Baldermann JC, Kibler A. Toward a unified connectomic target for deep brain stimulation in obsessive-compulsive disorder. bioRxiv 2019;608786.


Supplementary materials

Methods
The vmPFC was a specifically hypothesized ROI; however, none of the vmPFC to subcortical tracts passing through the aLIC had sufficient numbers of subjects with tracts (<90%). As we were uncertain if this was due to the low SNR in this region or that these tracts may be missed by limiting to tracts within the aLIC, we conducted a separate analysis including both vmPFC and Brodmann Area 25 to the NAcc, thalamus and STN without constraining to the aLIC.

Results
In the separate analysis for vmPFC (and BA 25) and NAcc without constraining in the aLIC, 89% of subjects showed tracts between this cortical-subcortical region (supplementary figure 3). The number of individuals with tracts between vmPFC (and BA 25) and STN (19%) and thalamus (65%) without constraining in the aLIC was still low with the latter including multiple implausible connections on visualization. We showed lower FA in OCD subjects compared to healthy controls ($t(165)=2.87, p=0.005$).
Supplementary Tables

Supplementary Table 1. Distribution of OCD symptoms.

<table>
<thead>
<tr>
<th>Symptom</th>
<th>OCD individuals reporting each symptom (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Symmetry</td>
<td>24</td>
</tr>
<tr>
<td>Taboo thoughts</td>
<td>31</td>
</tr>
<tr>
<td>Checking</td>
<td>30</td>
</tr>
<tr>
<td>Contamination/Cleaning</td>
<td>41</td>
</tr>
<tr>
<td>Hoarding</td>
<td>3</td>
</tr>
<tr>
<td>Other</td>
<td>21</td>
</tr>
</tbody>
</table>

Note: Totals exceed 100% given that each subject was allowed to report multiple obsessions and compulsions.
**Supplementary Table 2. Number of subjects with tractography results.**

<table>
<thead>
<tr>
<th></th>
<th>vlPFC</th>
<th>dlPFC</th>
<th>vmPFC</th>
<th>dACC</th>
<th>preSMA</th>
<th>OFC</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thalamus</td>
<td>180</td>
<td>187</td>
<td>33</td>
<td>182</td>
<td>187</td>
<td>184</td>
</tr>
<tr>
<td>Caudates</td>
<td>161</td>
<td>182</td>
<td>33</td>
<td>163</td>
<td>105</td>
<td>135</td>
</tr>
<tr>
<td>Putamen</td>
<td>179</td>
<td>183</td>
<td>29</td>
<td>157</td>
<td>170</td>
<td>104</td>
</tr>
<tr>
<td>NAcc</td>
<td>0</td>
<td>0</td>
<td>10</td>
<td>0</td>
<td>0</td>
<td>2</td>
</tr>
<tr>
<td>STN</td>
<td>179</td>
<td>171</td>
<td>2</td>
<td>152</td>
<td>172</td>
<td>169</td>
</tr>
</tbody>
</table>

vlPFC: ventrolateral prefrontal cortex, dlPFC: dorsolateral prefrontal cortex, vmPFC: ventromedial prefrontal cortex, dACC: dorsal anterior cingulate cortex, preSMA: anterior portion of supplementary motor area, OFC: orbitofrontal cortex, NAcc: nucleus accumbens, STN: subthalamic nucleus. The numbers in bold indicate tracts found in more than 90% of subjects (>169).
**Supplementary Table 3. Center of gravity of thalamocortical connections.**

<table>
<thead>
<tr>
<th>Cortical targets of thalamocortical connections</th>
<th>Min (mm)</th>
<th>Max (mm)</th>
<th>Mean (mm)</th>
<th>SD (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Left vlPFC</td>
<td>(-22, -1, 2)</td>
<td>(-15, 12, 17)</td>
<td>(-17, 8, 8)</td>
<td>(0.8, 1.7, 2.3)</td>
</tr>
<tr>
<td>Left dlPFC</td>
<td>(-20, 4, 1)</td>
<td>(-14, 11, 18)</td>
<td>(-17, 8, 9)</td>
<td>(0.8, 1.3, 2.7)</td>
</tr>
<tr>
<td>Left dACC</td>
<td>(-21, -4, 8)</td>
<td>(-16, 9, 17)</td>
<td>(-18, 5, 12)</td>
<td>(0.7, 1.7, 1.8)</td>
</tr>
<tr>
<td>Left preSMA</td>
<td>(-22, -4, 13)</td>
<td>(-17, 4, 17)</td>
<td>(-20, 1, 15)</td>
<td>(0.7, 1.5, 0.8)</td>
</tr>
<tr>
<td>Left OFC</td>
<td>(-19, 7, 2)</td>
<td>(-15, 11, 12)</td>
<td>(-17, 10, 6)</td>
<td>(0.6, 0.7, 1.6)</td>
</tr>
<tr>
<td>Right vlPFC</td>
<td>(12, -3, -2)</td>
<td>(22, 11, 17)</td>
<td>(18, 8, 8)</td>
<td>(1.1, 2.1, 2.7)</td>
</tr>
<tr>
<td>Right dlPFC</td>
<td>(16, 5, 2)</td>
<td>(20, 11, 15)</td>
<td>(18, 9, 8)</td>
<td>(0.7, 1.1, 2.1)</td>
</tr>
<tr>
<td>Right dACC</td>
<td>(17, 1, 6)</td>
<td>(22, 15, 18)</td>
<td>(19, 6, 12)</td>
<td>(0.8, 1.9, 2.3)</td>
</tr>
<tr>
<td>Right preSMA</td>
<td>(19, -5, 11)</td>
<td>(23, 4, 18)</td>
<td>(20, 1, 15)</td>
<td>(0.7, 1.9, 1.1)</td>
</tr>
<tr>
<td>Right OFC</td>
<td>(14, 6, -2)</td>
<td>(22, 21, 9)</td>
<td>(17, 10, 4)</td>
<td>(0.9, 1.2, 2.1)</td>
</tr>
</tbody>
</table>

Min: minimum, Max: maximum, SD: standard deviation. Note: Coordinates are in the MNI space.
## Supplementary Table 4. Clusters with voxel-wise group differences in FA.

<table>
<thead>
<tr>
<th>Cluster #</th>
<th>Anatomical region</th>
<th>Peak t-stat.</th>
<th>p-values (TFCE)</th>
<th>Size (mm$^3$)</th>
<th>X (mm)</th>
<th>Y (mm)</th>
<th>Z (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Anterior thalamic radiation</td>
<td>-4.6320</td>
<td>0.0159</td>
<td>31</td>
<td>-28</td>
<td>28</td>
<td>25</td>
</tr>
<tr>
<td>2</td>
<td>Anterior thalamic radiation</td>
<td>-4.3275</td>
<td>0.0229</td>
<td>23</td>
<td>-24</td>
<td>21</td>
<td>14</td>
</tr>
<tr>
<td>3</td>
<td>Forceps minor</td>
<td>-4.0792</td>
<td>0.0001</td>
<td>3147</td>
<td>-19</td>
<td>47</td>
<td>9</td>
</tr>
<tr>
<td>4</td>
<td>Forceps minor</td>
<td>-4.8755</td>
<td>0.0001</td>
<td>1161</td>
<td>-34</td>
<td>15</td>
<td>19</td>
</tr>
<tr>
<td>5</td>
<td>Cerebral WM in IFG</td>
<td>-5.0075</td>
<td>0.0001</td>
<td>760</td>
<td>-34</td>
<td>15</td>
<td>19</td>
</tr>
<tr>
<td>6</td>
<td>Inferior fronto-occipital fasciculus</td>
<td>-5.1242</td>
<td>0.0176</td>
<td>21</td>
<td>-29</td>
<td>47</td>
<td>-4</td>
</tr>
<tr>
<td>7</td>
<td>Cerebral WM in insular cortex</td>
<td>-5.9961</td>
<td>0.0001</td>
<td>372</td>
<td>-27</td>
<td>17</td>
<td>4</td>
</tr>
<tr>
<td>8</td>
<td>Cerebral WM in insular cortex</td>
<td>-4.0820</td>
<td>0.0403</td>
<td>16</td>
<td>27</td>
<td>15</td>
<td>11</td>
</tr>
<tr>
<td>9</td>
<td>Cerebral WM in insular cortex</td>
<td>-4.5232</td>
<td>0.0304</td>
<td>11</td>
<td>29</td>
<td>3</td>
<td>15</td>
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<td>10</td>
<td>Cerebral WM in MFG</td>
<td>-4.9230</td>
<td>0.0076</td>
<td>73</td>
<td>28</td>
<td>13</td>
<td>34</td>
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<tr>
<td>11</td>
<td>Cerebral WM in MFG</td>
<td>-5.0323</td>
<td>0.0055</td>
<td>73</td>
<td>-26</td>
<td>14</td>
<td>34</td>
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<tr>
<td>12</td>
<td>Cerebral WM</td>
<td>-4.1726</td>
<td>0.0069</td>
<td>250</td>
<td>-27</td>
<td>-7</td>
<td>21</td>
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<tr>
<td>13</td>
<td>Cerebral WM</td>
<td>-4.2759</td>
<td>0.0067</td>
<td>204</td>
<td>-26</td>
<td>-9</td>
<td>20</td>
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<tr>
<td>14</td>
<td>Subcortical WM</td>
<td>-3.0231</td>
<td>0.0458</td>
<td>3</td>
<td>25</td>
<td>1</td>
<td>31</td>
</tr>
<tr>
<td>15</td>
<td>Subcortical WM near putamen</td>
<td>-4.9229</td>
<td>0.0241</td>
<td>7</td>
<td>27</td>
<td>17</td>
<td>8</td>
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<tr>
<td>16</td>
<td>Superior longitudinal fasciculus</td>
<td>-3.9833</td>
<td>0.0210</td>
<td>64</td>
<td>38</td>
<td>-40</td>
<td>25</td>
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<tr>
<td>17</td>
<td>Superior longitudinal fasciculus</td>
<td>-4.2858</td>
<td>0.0235</td>
<td>24</td>
<td>39</td>
<td>41</td>
<td>-32</td>
</tr>
<tr>
<td>18</td>
<td>Superior longitudinal fasciculus</td>
<td>-4.3635</td>
<td>0.0288</td>
<td>18</td>
<td>53</td>
<td>-46</td>
<td>-15</td>
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<tr>
<td>19</td>
<td>Superior longitudinal fasciculus</td>
<td>-4.1833</td>
<td>0.0429</td>
<td>2</td>
<td>34</td>
<td>-22</td>
<td>33</td>
</tr>
<tr>
<td>20</td>
<td>Uncinate fasciculus</td>
<td>-4.5199</td>
<td>0.0251</td>
<td>22</td>
<td>-32</td>
<td>31</td>
<td>-7</td>
</tr>
</tbody>
</table>

WM: white matter, IFG, inferior frontal gyrus, MFG, middle frontal gyrus. Coordinates are in the MNI space. Note: T-statistics were computed for the contrast of healthy controls subtracted from patients with OCD.
Supplementary Figures

Supplementary Figure 1. Denoising using local PCA. Raw DSI data (upper row) and denoised DSI data (lower row) using local PCA. B-values are marked.
Supplementary Figure 2. Examples of registration of regions of interest. For randomly chosen four subjects (two health controls, HC; two patients with OCD, OCD), regions of interest (ROI) registered in the native diffusion-spectrum imaging (DSI) space are shown. Color-coding (the same scheme as Figure 1 in the main text) marks integer labels: 1 = ventrolateral prefrontal cortex, 2 = dorsolateral prefrontal cortex, 3 = ventromedial prefrontal cortex, 4 = dorsal anterior cingulate cortex, 5 = anterior portion of supplementary motor area, 6 = orbitofrontal cortex, 7 = thalamus, 8 = caudate nucleus, 9 = putamen, 10 = nucleus accumbens, 11 = subthalamic nucleus.
Supplementary Figure 3. Clinical measures at baseline and follow-up. A: Yale-Brown Obsessive-Compulsive Scale (Y-BOCS), B: Hamilton Anxiety Rating Scale (HAM-A), C: Hamilton Depression Rating Scale (HAM-D).
Supplementary Figure 4. Ventromedial prefrontal and nucleus accumbens connectivity.
A: tracts from a representative healthy control subject. B: group-averaged track-density image. C: Group difference in fractional anisotropy of the tracts.
Supplementary Figure 5. Overlap between aLIC masks and lesion probability map. The masks for anterior limbs of internal capsules (aLIC) were taken from the ICBM-DTI-81 white-matter atlas (as included in FSL) and used as seeding masks in the current study (blue contours). The lesion probability map is from a separate study from our group\(^1\); from a subset of OCD patients (\(n = 31\)), lesion masks were manually delineated by two human raters, averaged, and non-linearly registered to MNI-template space. Because the ICBM-DTI-81 labels were conservatively drawn without overlaps with adjacent structures (e.g., anterior corona radiata in green), the lesion is overlapped with the anterior part of the aLIC mask.