Mapping the human praxis network: an investigation of white matter disconnection in limb apraxia of gesture production

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Left hemispheric cerebral stroke can cause apraxia, a motor cognitive disorder characterized by deficits of higher-order motor skills such as the failure to accurately produce meaningful gestures. This disorder provides unique insights into the anatomical and cognitive architecture of the human praxis system. The present study aimed to map the structural brain network that is damaged in apraxia. We assessed the ability to perform meaningful gestures with the hand in 101 patients with chronic left hemisphere stroke. Structural white matter fibre damage was directly assessed by diffusion tensor imaging and fractional anisotropy mapping. We used multivariate topographical inference on tract-based fractional anisotropy topographies to identify white matter disconnection associated with apraxia. We found relevant pathological white matter alterations in a densely connected fronto-temporo-parietal network of short and long association fibres. Hence, the findings suggest that heterogeneous topographical results in previous lesion mapping studies might not only result from differences in study design, but also from the general methodological limitations of univariate topographical mapping in uncovering the structural praxis network. A striking role of middle and superior temporal lobe disconnection, including temporo-temporal short association fibres, was found, suggesting strong involvement of the temporal lobe in the praxis network. Further, the results stressed the importance of subcortical disconnections for the emergence of apractic symptoms. Our study provides a fine-grain view into the structural connectivity of the human praxis network and suggests a potential value of disconnection measures in the clinical prediction of behavioural post-stroke outcome.

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Abbreviations: ABA-2 = Apraxia Battery for Adults-2; DTI = diffusion tensor imaging; FA = fractional anisotropy; FDR = false discovery rate; FOV = field of view; IFG = inferior frontal gyrus; IPL = inferior parietal lobe; MNS = mirror neuron system; MTG = middle temporal gyrus; SMG = supramarginal gyrus; STG = superior temporal gyrus; SVR = support vector regression; SVR-LSM = support vector regression-based lesion-symptom mapping; TBSS = tract-based spatial statistics; TE = echo time; TI = inversion time; TR = repetition time; WAB-R = Western Aphasia Battery-Revised

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Introduction

Stroke to the left hemisphere of the brain often causes disorders of higher-order motor control. These disorders are summarized under the term ‘apraxia’ and include symptoms such as the inability to perform, imitate or recognize different kinds of gestures. In the present study, we investigated apraxia in the production of meaningful gestures, including the pantomime of tool use and communicative gestures like ‘waving goodbye’. For simplicity, we hereafter refer to these deficits, which belong to symptoms of the classical category of ideomotor apraxia, as ‘apraxia’. Many previous studies aimed to identify the neural correlates of apraxia. These studies suggest damage in parietal, temporal and frontal cortical regions mainly in the left hemisphere was associated with apraxia. 

Further studies, e.g. using electroencephalography, functional MRI and fibre tracking in healthy subjects, support this assumption.

The idea that apraxia results from disruptions of a complex brain network is as old as its first case descriptions by Liepmann. Although theories about the architecture of this network have undergone adaptations throughout the past century, the notion of a fronto-(temporo-)parietal praxis network still dominates the field. However, no single, widely accepted view of how the proposed network is structured exists. Further, most network theories of apraxia so far focused on cortical nodes of hypothesized networks, while white matter contributions were a minor topic in only a few previous lesion-behaviour mapping studies. White matter tracts associated with apraxia of pantomime were found below the precentral gyrus and in ventral fibres of the extreme capsule. A multivariate lesion-behaviour mapping study found focal lesion damage to several major fibre bundles, including fronto-parietal connections, to be associated with apraxia. Garcea et al. mapped the praxis network in reference to both structural lesion information.
and indirect connectome-based lesion mapping. Their study mainly implicated disconnection of parietal and ventral temporal regions, as well as disconnection of a few superior frontal regions.

However, the identification of white matter damage based on lesion-behaviour mapping bears limitations. First, univariate statistical approaches might lack statistical power to identify a full brain network, including relevant fibre tracks. Second, topographical structural lesion–brain inference only indirectly assesses white matter contributions to symptoms via reference to brain atlases and conclusions depend on what brain atlas is chosen to interpret results. Further, a holistic understanding of such a network is difficult to achieve. While the sum of all previous lesion-behaviour mapping studies suggests a set of frontal, parietal and temporal grey matter nodes, marked heterogeneity between topographical results exists. It is unclear to what extent the heterogeneity should be attributed to differences in sample characteristics, methodology or apraxia assessment, and to what extent to the above-mentioned limitations of univariate statistics.

The present study aimed to directly assess white matter damage contributions to stroke patients’ apractic deficits in the production of meaningful gestures, in contrast to previous studies that only indirectly assessed white matter disconnection. We did so by combining multivariate lesion-behaviour mapping with tract-based spatial statistics (TBSS) of fractional anisotropy (FA). A similar methodology has previously been applied in stroke patients to identify the contribution of white matter degeneration or abnormalities to spatial neglect. We utilized multivariate spatial statistics that potentially solve the limitations of univariate statistics in identifying brain networks and mapped tract-based FA abnormalities in chronic left hemisphere stroke patients. We aimed to identify (i) direct effects of lesion damage to white matter tracts and (ii) remote or long-term lesion effects on white matter integrity, as, for example, induced by Wallerian degeneration.

**Materials and methods**

**Patients and behavioural assessment**

We evaluated 101 individuals who had suffered a first-time major left hemisphere ischaemic stroke from an archival database at the University of South Carolina. This database contains test results from several research studies investigating aphasia and its clinical management. Patients were included in these research studies if they were between the ages of 21 and 80 and at least 5 months post-stroke. Exclusion criteria included: (i) stroke involving the right hemisphere, brainstem and/or cerebellum; (ii) other neurologic illness/injury affecting the brain and (iii) history of developmental speech and/or language disorder. Of note, we included participants in the present study regardless of aphasia diagnosis, and the database also includes patients with only mild or no aphasia at all. All participants signed an informed consent form that was approved by the Institutional Review Board at the University of South Carolina or the Medical University of South Carolina. All study procedures adhered to the principles set forth by the revised Declaration of Helsinki. Descriptive statistics concerning demographic, behavioural and lesion data are shown in Table 1.

Participants were evaluated during the chronic stage of stroke. Aphasia was tested with the Western Aphasia Battery-Revised (WAB-R). For patients with a diagnosis of aphasia (i.e. a WAB-R Aphasia Quotient < 93.8), a speech-language pathologist confirmed that their understanding was sufficient for task completion, and patients with severe comprehension difficulties were excluded. Participants were included regardless of the presence of apraxia of speech and unilateral primary motor deficits.

Apraxia was assessed using the limb apraxia subscale from the Apraxia Battery for Adults. The test contains 10 gestures that are to be executed upon oral instruction (‘make a fist’, ‘wave good-bye’, ‘snap your fingers’, ‘throw a ball’, ‘hide your eyes’, ‘make a hitchhiking sign’, ‘make a pointing sign’, ‘salute’, ‘play the piano’, ‘scratch’; for gestures involving objects, those objects were not present and participants had to pantomime their use). The gesture was executed using the preferred hand post-stroke and rated between zero (inability to complete the gesture even after demonstration of the gesture by the examiner) and five for best performance (i.e. the participant provides an accurate and prompt gestural response). Four points were given if the examinee produced an incorrect gesture, but self-corrected it; three points were given if the gesture was basically correct but crude and defective; two points were given if the examinee produced the correct gesture after an additional demonstration by the examiner; one point was given if the gesture after the additional demonstration was basically correct but crude and defective. Testing was conducted by an American Speech-Language-Hearing Association-certified speech-language pathologist, who administered and scored all assessments.

**Imaging and image processing**

Brain imaging was acquired using a 3 T Siemens Trio scanner with a 12-channel head coil. T1-weighted scans were obtained with an MP-RAGE (TFE) sequence with voxel size = 1 mm³, FOV = 256 mm × 256 mm, 192 sagittal slices, 9° flip angle, TR = 2250 ms, TI = 925 ms and TE = 4.15 ms, GRAPPA = 2, 80 reference lines. T2-weighted scans were obtained with 3D SPACE protocol with voxel size = 1 mm³, FOV = 256 mm × 256 mm, 160 sagittal slices, variable flip angle, TR = 3200 ms, TE = 352 ms, no slice acceleration and the same slice centre and angulation as in the T1 imaging. FA was computed using diffusion tensor imaging (DTI). We acquired diffusion imaging by a mono-polar sequence with 82 isotropic (2.3 mm) volumes (× 10 B = 0, × 72 B = 1000), TR = 4987 ms, TE = 79.2 ms, 90 × 90 matrix, with parallel imaging GRAPPA = 2 and 50 contiguous slices. The sequence was carried out in two series.
(41 volumes in each series) with opposite phase encoding polarity. We denoised images using MRtrix (https://www.mrtrix.org/) and undistorted them using FSL’s topup and Eddy tools. We converted 4D DTI images, bvecs and bvals to FA maps with FSL’s DTIFIT tool. We further processed FA data according to the standard procedures for TBSS\textsuperscript{27} in FSL, with the addition that during non-linear registration, each patient’s lesion was masked to avoid distortions. TBSS projects each subject’s FA data onto a mean FA tract skeleton (Fig. 1).

The stroke lesions were manually delineated on the T2-weighted images by trained personnel under the supervision of a neurologist (L.B.) and of a senior lab manager (Dr Roger Newman-Norlund, PhD), using MRICron (https://www.nitrc.org/projects/mricron/). Each lesion was drawn by one trainee and the quality was checked by L.B. or by R.N.N. When necessary, the lesion was adjusted or redrawn by L.B. or R.N.-N. to ensure anatomical correctness. All personnel (trainees, L.B. and R.N.-N.) were blinded to the behavioural scores. We co-registered T2-weighted scans and lesion masks to T1-weighted imaging. Lesion masks were normalized by warping T1-weighted images to age-specific normalization templates of elderly healthy controls using SPM 12 (http://www.fil.ion.ucl.ac.uk/spm/software/spm12/) and Clinical Toolbox\textsuperscript{34} with enantiomorphic-segmentation normalization (Fig. 1).

Table 1 Demographic and clinical data for patients with and without apractic deficits

|                      | Apractic patients | Non-apractic patients | Test statistics |
|----------------------|-------------------|-----------------------|-----------------|
| Age (years)          | 57.7 (11.9)       | 60.8 (10.6)           | $t_{99} = 1.297; P = 0.457$ |
| Sex (f/m)            | 15/16             | 22/48                 | $\chi^2 (1, n = 101) = 2.66; P = 0.103$ |
| Lesion size (cm$^3$) | 174.8 (95.7)      | 108.4 (84.4)          | $U = 1565; P < 0.001$ |
| ABA-2 apraxia score  | 35.4 (9.3)        | 48.3 (1.9)            | $U = 0; P < 0.001$ |
| WAB aphasia score    | 42 (16.8)         | 67.9 (21.5)           | $U = 387; P < 0.001$ |
| Aphasia (y/n)        | 31/0              | 66/4                  | $\chi^2 (x) = 1.84; P = 0.174$ |
| Time post-stroke at screening (months) | 53.5 (59.7) | 38.5 (39) | $U = 1145; P = 0.492$ |

Based on ABA-2 norms, patients with a score of 44 or less were put in the apraxia group. The group distinction was made solely for descriptive purposes and all further analyses were based on the full continuous data. Continuous values are presented as mean (standard deviation). Assessments of potential group differences are reported in the right column. The distribution of all ABA-2 apraxia scores is reported in the Supplementary material. ABA-2, Apraxia Battery for Adults-2.

Figure 1 Lesion topography and mean FA. (A) The lesion topography after enantiomorphic normalization in SPM indicating the number of overlapping lesions per voxel for all 101 patients. (B) The average skeletonized FA topography after normalization and skeletonization with TBSS projected on the average non-skeletonized, normalized FA maps. Numbers above slices indicate the Z-coordinate in MNI space.
Support vector regression-based FA mapping

We investigated the relation between voxel-wise FA and apraxia scores with a multivariate topographical analysis based on support vector regression (SVR). This multivariate topographical inference method was adopted from the field of lesion-behaviour mapping, where SVR-based lesion-symptom mapping (SVR-LSM) was previously used to identify the neural correlates of apraxia of pantomime. This method involved two major steps: first, we trained a SVR to predict behavioural scores based on FA data. Second, we statistically evaluated the contribution of individual voxels to the model with a permutation approach. In detail, we modelled apraxia scores with an SVR with a linear kernel using voxel-wise, skeletonized FA topographies. Hyperparameter C was optimized to maximize model fit and reproducibility of feature weights with five times 5-fold cross-validation procedure in the range of $C = [2^{-20}, 2^{-19}, \ldots, 2^{19}, 2^{20}]$. We evaluated model fit by SVR modelling of four-fifths of the data and assessed the correlation between the real and predicted scores in the last fifth of the data. To evaluate reproducibility, we computed the average correlation coefficient $\tau$ of feature weights between each pair of cross-validation subsets and then the average $\tau$ of all possible combinations of subsets. With the optimized $C$, we computed an SVR using data from all 101 patients and determined the feature weights, i.e. the contribution of each voxel-wise FA value to the model. A second step employed a permutation procedure to identify the features that provide a statistically significant contribution to this model. Following the SVR modelling steps as described above, voxel-wise feature weights were also computed 25 000 times for data sets consisting of the original FA data and a random permutation of the original behavioural data. Next, for each voxel, a statistical $P$-value was assigned to each feature weight in the original feature weight map by reference to the permutation-based feature weights. The required correction for multiple comparisons was carried out by false discovery rate (FDR) correction at $q \geq 0.05$. Further analyses were conducted with the FDR-corrected topography and with small clusters <20 voxels removed.

Statistical analysis: region-based evaluation of structural disconnectivity

To identify region pair-wise disconnection associated with significant FA abnormalities, we evaluated the disconnection underlying the TBSS-based SVR-LSM analysis. The exact procedure can be found in Supplementary materials. In short, we created a whole-brain tractogram with MRtrix (https://www.mrtrix.org/) using healthy controls’ data from the IIT Human Brain Atlas v.5.0 (https://www.nitrc.org/projects/iit/) representing the healthy connectome based on 84 regions of interest of the Desikan–Kilkenny atlas and the whole-brain tractogram. We then quantified the disconnection underlying our statistical group-level results by extracting all tracts crossing the three-dimensional statistical SVR-FA-map. We chose to interpret the absolute number of disconnected streamlines as the main outcome variable. Compared with the proportion of disconnected streamlines, this strategy prevented us from overestimating the role of originally faintly connected areas.

In a second analysis, the clusters of significant voxels in the SVR-FA mapping were referenced to a probabilistic white matter atlas based on DTI to identify a possible affection of major fibre tracks. Probabilistic maps of long association, projection and commissural fibres were binarized at $P \geq 0.4$. Binary maps were overlaid with the SVR-FA-map and the overlapping volume was identified. Since such anatomical interpretation largely depends on the choice of white matter atlas to interpret results, we additionally referenced results to a probabilistic cytoarchitectonic atlas in the Supplementary material and make topographies publicly available in our online materials.

Data availability

All analysis scripts and resulting topographies are publicly available at http://dx.doia.org/10.17632/dcpst33wc7.3. The clinical data sets analysed in the current study are not publicly available due to the data protection agreement approved by the local ethics committees and signed by the participants. They are available from J.F. (fridriks@mailbox.sc.edu) on reasonable request.

Results

The hyperparameter optimization found $C = 2^{-18}$ to provide both a high reproducibility of $\tau = 0.84$ and a relatively high model fit of $r^2 = 0.25$. The permutation-based and FDR-controlled SVR-FA-map identified larger clusters of significant FA-behaviour relations in several left hemisphere regions, where increased FA was associated with more severe apraxia (Fig. 2A). These included the largest clusters in parieto-temporal and temporal white matter and further clusters in inferior frontal, parietal and subcortical white matter.

Since the present study analysed DTI data from neurological patients and examined them for inter-regional discontinuities using DTI data from healthy subjects, we used a DTI-based atlas for referencing clusters of significant voxels to white matter anatomy (Table 2). It implicated the involvement of several major fibre tracts in apraxia, including all segments of the superior longitudinal fascicle, the uncinate fascicle, the inferior fronto-occipital fascicle and the inferior longitudinal fascicle. Further, some significant clusters were assigned to projection fibres, including the corticospinal tract. Additionally, we referenced clusters of significant voxels to a probabilistic cytoarchitectonic atlas, as documented in the online materials (http://dx.doia.org/10.17632/dcpst33wc7.3).
This analysis found a role of the inferior fronto-occipital fascicle, the corticospinal tract and callosal body.

In regions outside of the direct lesion areas, no significant FDR-corrected FA alterations could be detected. This means that the analysis did not identify any supra-threshold remote effects of FA related to apractic deficits. However, several sub-FDR-threshold clusters in the uncorrected SVR-FA-map (Fig. 2B) hinted at possible remote white matter alterations. Small clusters were scattered across frontal and posterior parts of the corpus callosum and in the right hemisphere.

The region-based evaluation of structural disconnectivity identified a large number of inter-regional disconnections of
Table 2 Overlap between fibre tracts as defined by Zhang et al. and the significant voxels found in our analysis

| Fibre tract category | Number of affected voxels/mm³ |
|----------------------|-------------------------------|
| Long association     |                               |
| Inferior fronto-occipital f. | 307                           |
| Inferior longitudinal f. | 103                           |
| Superior longitudinal f. (FP) | 98                            |
| Superior longitudinal f. (FT) | 268                           |
| Superior longitudinal f. (PT) | 474                           |
| Uncinate fasciculus | 279                           |
| Corticospinal tract | 326                           |
| Thalamus—precentral gyrus | 38                            |
| Thalamus—superior frontal gyrus | 127                        |
| Thalamus—superior occipital gyrus | 36                         |

Projection

| Corticospinal tract | 326 |
| Thalamus—preccentral gyrus | 38 |
| Thalamus—superior frontal gyrus | 127 |
| Thalamus—superior occipital gyrus | 36 |

* f, fasciculus; FP, fronto-parietal; FT, fronto-temporal; PT, parieto-temporal. There was no overlap with commissural fibre tracts. Fibre tracts with <20 mm³ overlap are not reported.

The left hemisphere grey matter areas that displayed the highest numbers of disconnected streamlines and that potentially execute a hub-like function in the praxis network, were found to be widely spread across the left cerebral hemisphere (see Table 3). The largest amount of disconnections was found in the basal grey matter, which is not possible with the SVR-FA analysis. We used the same SVR algorithms as in the SVR-FA analysis with a linear kernel and 25 000 permutations to map the relation between apraxia and binary lesion maps. The hyperparameter optimization suggested $C = 2^{−17}$, for which the reproducibility was $r = 0.72$ and the model fit $r^2 = 0.22$. No significant results remained after correction by FDR. The lower reproducibility and model fit hint at possible limitations of the linear SVR in modelling structural lesion data. Therefore, we report an additional SVR-lesion-behaviour mapping analysis with different specifications, including a non-linear kernel, in the Supplementary material.

**Discussion**

The present analysis used magnetic resonance diffusion imaging and tract-based, multivariate topographical inference to uncover white matter damage related to chronic apraxia in the production of meaningful gestures. We identified a fronto-parieto-temporal praxis network with a parieto-temporal focus. Considerable apraxia-related disconnection was found for short association fibres within the temporal lobe, as well as the parietal and the frontal lobe, and long association fibres connecting the frontal and temporal lobe with parietal areas. Notably, marked disconnection of middle and superior temporal regions was implicated, suggesting the temporal lobe as one of the main hubs in the praxis network. Further, the basal ganglia were prominently implicated with disconnection of the putamen and caudate, including connections with the frontal cortex and the insula. Significant remote effects of lesions on white matter integrity were not identified, but non-significant clusters in the right hemisphere and the callosal body hinted at potential effects that were not accessible with our methods.

**A structural praxis network**

The associations found between reduced FA in white matter and apractic deficits largely correspond with some existing network models. A recent study on indirect connectivity measures by Garcea et al. mapped disconnections associated with hand posture errors in tool use gestures. They found fronto-temporo-parietal disconnection to underlie deficits in this task. They also found a focus on parietal and temporal areas. A meta-analysis by Niessen et al. on pantomime—i.e. meaningful gestures with an imaginary tool—points in the same direction. They suggested that...
Apraxia of pantomime involves a network containing inferior parts of the parietal lobe, the frontal gyrus and temporal regions. We found disconnections involving the areas proposed by Niessen et al., as well as subcortical regions and succeeded in providing a more fine-grain description of connections underlying apraxia. Our results indicate the importance of temporo-parietal connections between lateral temporal and posterior parietal areas. Our parcellation also allowed more detailed conclusions concerning frontal region-wise disconnections. In the middle frontal gyrus,

**Figure 3 Region-based evaluation of structural disconnectivity (1).** Line colour indicates the number of disconnected fibre bundles between grey matter areas in the Desikan atlas in the fibre tracking analysis in MRtrix. The threshold was set to 130 fibres to depict the strongest inter-regional disconnections.

**Figure 4 Region-based evaluation of structural disconnectivity (2).** Heatmap of the 20 areas found to be most severely disconnected. Cell colour refers to the number of disconnected streamlines between the brain areas represented by the corresponding row and column. Full disconnectivity data are available in the online materials. A heatmap for the proportion of disconnected streamlines for the same regions is shown in the Supplementary material.
rostral and caudal portions showed high numbers of disconnected fibres. In the IFG, in accordance with Garcea et al., particularly the pars orbitalis and pars triangularis were disconnected. Notably, the current study succeeded in identifying all these areas in association with apraxia in one single sample and with one single apraxia measure. Hence, we could show that differences in topographical results in previous studies might not only result from differences in study design or apraxia assessment, but also from the general methodology that was unsuited to identify a praxis network.

The study’s multivariate approach might account for disconnections in most brain regions so far hypothesized to be relevant in apraxia, while univariate statistical approaches in previous studies tended to find only parts of this praxis network (see also Sperber et al.14). Another limitation of many previous studies might be the focus on structural lesion data. In an additional analysis, we modelled and mapped the relation between apraxia and structural lesion data with the same algorithms as in the SVR-FA mapping. Not only did this result in worse model performance, but also a null result after correction for multiple comparisons. This result could indicate that focal lesion data alone are less informative and predictive of apraxia. On the other hand, it is also possible that this finding simply hints at limitations of the modelling algorithm, as the prediction performance can differ across algorithms.39 Accordingly, when we performed an additional SVR-lesion-behaviour mapping with a non-linear kernel, we found significant results in the left basal ganglia, the left inferior and middle temporal cortex and the left lateral orbito-frontal cortex. Importantly, however, our white matter analysis uncovered the disconnection of areas that were not revealed by the grey matter analysis. For example, both left inferior parietal and superior frontal areas were found to be disconnected in our SVR-FA mapping while our grey matter analysis did not reveal lesions in these areas to be associated with apraxia.

### Table 3 Grey matter areas in the left hemisphere which displayed the highest disconnection counts (>1000) in the region-based evaluation of structural dysconnectivity

| Brain parcellation | Number of disconnections | Corresponding fibre damage (%) |
|--------------------|--------------------------|-------------------------------|
| Putamen            | 1904                     | 17.1                          |
| Superior temporal  | 1832                     | 23.5                          |
| Caudate            | 1633                     | 17.5                          |
| Supramarginal      | 1574                     | 17.4                          |
| Middletemporal     | 1506                     | 18.8                          |
| Precentral         | 1487                     | 12.0                          |
| Inferiorparietal   | 1357                     | 13.8                          |
| Insula             | 1205                     | 14.5                          |
| Superior frontal   | 1071                     | 8.0                           |
| Lateralorbitofrontal | 1070                  | 14.9                          |

The areas were defined by a volumetric grey matter atlas.33 The last column indicates the damage in relation to all existing fibers connecting the area.

### Table 4 Strongest direct disconnections between grey matter pairs which displayed the highest disconnection counts (>125) in the region-based evaluation of structural dysconnectivity

| Disconnected brain areas                  | Number of disconnections | Corresponding fibre damage (%) |
|-------------------------------------------|--------------------------|-------------------------------|
| Transversetemporal—superiortemporal       | 388                      | 39.2                          |
| Caudate—putamen                           | 374                      | 11.5                          |
| Superiortemporal—supramarginal            | 320                      | 26.8                          |
| Banksts—middletemporal                    | 301                      | 12.1                          |
| Partriangularis—lateralorbitofrontal      | 292                      | 37.5                          |
| Supramarginal—inferiorparietal            | 279                      | 10.4                          |
| Parsorbitalis—lateralorbitofrontal        | 240                      | 16.4                          |
| Middletemporal—inferiorparietal           | 239                      | 21.3                          |
| Superiortemporal—banksts                  | 224                      | 10.7                          |
| superiorparietal—inferiorparietal         | 203                      | 10.2                          |
| Rostralmiddlefrontal—caudate              | 202                      | 29.5                          |
| Precentral—putamen                        | 195                      | 37.5                          |
| Supramarginal—superiorparietal            | 181                      | 19.1                          |
| Putamen—lateralorbitofrontal              | 176                      | 48.8                          |
| Insula—putamen                            | 174                      | 14.1                          |
| Superiorfrontal—putamen                   | 173                      | 40.4                          |
| Superiorfrontal—caudate                   | 158                      | 22.5                          |
| Precentral—thalamus                       | 148                      | 38.8                          |
| Superiorfrontal—thalamus                  | 147                      | 48.4                          |
| Lateralorbitofrontal—caudate              | 135                      | 40.1                          |
| Precentral—inferiorparietal               | 132                      | 62.0                          |
| Precentral—caudate                        | 131                      | 48.5                          |
| Precentral—supramarginal                  | 130                      | 35.2                          |
| Putamen—rostralmiddlefrontal              | 127                      | 60.5                          |
| Superiorparietal—middletemporal           | 125                      | 77.6                          |

The areas were defined by a volumetric grey matter atlas.33 Column 3 indicates the damage in relation to all existing fibers connecting the areas. All affected structures were located in the left hemisphere.
analysis. However, the uncorrected statistical topography hints at possible effects. Sub-FDR-threshold clusters were apparent in the splenium and the genu of the corpus callosum. This is in line with previously reported and rare cases of callosal apraxia, which is caused by damage to the splenial part of corpus callosum. Non-callosal apraxia might therefore indirectly affect white matter integrity in remote areas that are directly damaged in cases of callosal apraxia. The finding is also in line with studies using resting-state functional MRI. Future studies are required to confirm findings on the praxis network structure—including the present study—into a comprehensive network model that also accounts for interhemispheric communication.

The interpretation of the statistical results with the probabilistic fibre tract atlas found a large overlap with the left corticospinal tract split across two clusters. One cluster was close to the left lateral ventricle and the other below the left thalamus and reached into the brainstem. This was a surprising finding, as we tested all subjects with the ipsilesional, non-paretic hand. One possible explanation could be the common co-morbidity of apraxia and right-sided hemiparesis. Both are typical symptoms of left-sided middle cerebral artery stroke that can co-occur and mapping one function might, to a small degree, also map the other function. Another explanation could be that these results reflect an actual impact of the ipsilesional primary motor system on the function of the ipsilesional upper limb. Recent studies found evidence for the role of both ipsi- and contralateral M1 on skilled motor function (for review, see Bradnam et al. and its recovery after stroke. However, the role of uncrossed fibres of the corticospinal tract in the adult human brain is still under discussion. As we did not assess contralesional primary motor deficits in the current study, future studies are required to elaborate on this issue and how possible effects might relate to apraxia.

Grey matter nodes in the praxis network

The region-wise interpretation of our topographical results validated the importance of frontal, temporal and parietal nodes in a single praxis network. Doing so, the present study integrates largely heterogeneous findings from previous lesion-behaviour mapping studies.

The IPL conventionally consists of both angular gyrus and SMG, but the atlas applied in the current study denotes an extra parcellation to SMG. The IPL was often seen as one of the most central areas in a praxis network. Our results support this notion. The SMG was among those areas with the highest number of disconnected streamlines, closely followed by the rest of the IPL, including the angular gyrus. Further, we found relevant short intra-parietal disconnections between neighbouring parietal areas, or even between subregions of the same structure. While the presence of short association fibres within a single cortical area might seem counterintuitive at first glance, it was shown that these intra-regional fibres can be identified and tracked with modern DTI. The parietal regions of the praxis network were previously assigned to process spatio-temporal aspects of acquired movements. The parietal lobe seems to store motor programmes involved in both tool use and the equivalent pantomimes and to code object positions in relation to oneself and other functions assigned to the parietal lobe include planning, predicting and choosing appropriate actions.

Disconnections of pars triangularis and pars orbitalis in the IFG are in accordance with accounts highlighting a major role of the IFG in apraxia of pantomime. Goldenberg et al. argued that activations in the IFG might be linked to the selection of pantomime-specific action features from semantic memory. While they focused on apraxia of pantomime exclusively, the present study investigated pantomimes of object use (such as ‘play the piano’) and communicative gestures (such as ‘wave good-bye’). An alternative explanation, accounting for pantomimes and emblems, is that frontal activations are concerned with the gestures’ inherent symbolism. The IFG is popular for its role in language, a skill similarly depending on symbolism. Its involvement is not restricted to spoken language but also applies to other forms of communication. While IFG activations were greater for deaf signers of American Sign Language when attending to signs than when watching pantomimes, hearing non-signers showed higher activations for familiar pantomimes than for the unfamiliar sign language. These results coincide with more recent work by Goldenberg, which argued tool use pantomimes to be communicative gestures, rather than replications of real tool use. IFG activations and their connections might further indicate involvement of the mirror neuron system (MNS), which engages the same neurons both when watching an action and executing it. The activations found when watching communicative gestures in aforementioned studies might therefore also be involved in the execution of such gestures and lesions in said areas might impair processing a gesture’s inherent symbolism. Another crucial MNS region, the IPL also displayed a high number of disconnections in this study. Further damage was found in fronto-parietal fibre tracks including the fronto-parietal superior longitudinal fascicle.

Disconnection of the precentral gyrus already held an important role in classical models of apraxia. They are believed to process motor programmes forwarded from parietal areas. Accordingly, EEG activity was found to begin in parietal areas and spread to left premotor areas in the course of movement preparation. This role of premotor areas was argued to be generally applicable for complex motor tasks. Finally, the insula, which was proposed to be critically involved in apraxia of pantomime, displayed reduced connectivity. Like IPL and IFG, the insula is associated with the MNS, apparently connecting the frontoparietal MNS to the limbic system.

While the basal ganglia are known to be an important component of motor control circuits, research on apraxia indicates that subcortical areas play a less prominent role in
gesture production. Although cases of ideomotor apraxia following subcortical insults have been reported, a re-evaluation by Pramstaller and Marsden of all published cases of ideomotor apraxia after a subcortical stroke before 1994 revealed that only a very small fraction of lesions (8 out of 82) were isolated subcortical insults without white matter involvement. A more recent study investigated sub-symptoms of ideomotor apraxia in greater detail and included both cortical and subcortical lesions. Also here, at least seven out of nine subcortical lesions stretched into surrounding white matter. Taken together, the existing literature on subcortical area involvement in meaningful gesture production only provides limited evidence on the role of the basal ganglia in apraxia. The number of cases with isolated subcortical lesions is vanishingly low. It might therefore be that disconnections to rather than lesions in said regions are relevant for apraxia since the basal ganglia indubitably play an important role in the control of motor skills. Different connections between the basal ganglia and several cortical structures, such as the IPL, have been identified as part of the so-called lateral grasping network in primate models and suggest the existence of several basal ganglia loops concerned with hand-related actions. Further praxis-related skills with involvement of the basal ganglia include movement timing and the execution of automatized movements.

Temporal contributions to a praxis network

So far, only a few lesion-behaviour mapping studies have implicated temporal regions. Since the present study found strong temporo-parietal and temporo-temporal disconnections associated with apraxia, not only cortical lesions to the MTG seem to be associated with apraxia but also decreased connectivity to that very region. We speculate that traditional mass-univariate lesion-behaviour mapping may have low statistical power for identifying large functional regions where small injuries are sufficient to yield impairment. In such cases, different individuals can show similar symptoms with mutually exclusive lesion locations. Since lesion-behaviour mapping identifies regions that are reliably damaged in patients with an impairment and reliably spared in those without symptoms, each of these individuals acts as a counter-example for the other. Our methods of identifying fibres as well as using multivariate analyses may mitigate these issues.

Network theories of praxis most commonly assigned action recognition to temporal areas. Recognition capabilities related to this area are not only limited to actions but also include objects. White matter lesions affecting the posterior part of the MTG were further associated with impairing conceptual aspects of action knowledge. Similarly, the relevance of MTG found in the present study might reflect conceptual issues impairing action knowledge and thus contribute to apraxia.

Results concerning the superior temporal sulcus are again prompting towards a possible role of action recognition, more precisely, the perception of biological motion. If this important foundation for recognizing actions plays a role in apraxia remains speculation, which requires future studies to be investigated. Further, the STG might be involved in evaluating the context-specific plausibility of action intentions. Since action understanding was not explicitly tested, it remains an enigma if and how understanding action and performing it relate to one another. Involvement of the temporal lobe has also been found to be relevant in the context of anosognosia for limb apraxia.

There is more to praxis and its network than a single test can measure

‘Apraxia’ is often used as an umbrella term and refers to a multitude of deficits of higher-order motor skills, including, among others, the execution of meaningful gestures. Deficits of higher-order motor skills can both behaviourally and anatomically dissociate into sub-deficits. The present study focused on the execution of meaningful gestures after verbal instruction. The gesture classes tested in this study, namely transitive and intransitive gestures, are known to dissociate in their cognitive load and neural correlates.

It is yet to be answered how different apraxia subtypes are related. Subtypes of apraxia occur together but are also known to dissociate occasionally. The classification based on action form (e.g. here meaningful gestures) can even be further subdivided according to error type. To conclude, given the complexity of praxis skills and apraxia, the delineation of the entire praxis network is not feasible within a single study. Whatever behavioural measure is chosen to access certain praxis skills, other aspects will not be assessed. Likewise, this study did not identify the entire praxis network, but the praxis sub-network related to the production of meaningful gestures—a skill commonly impaired in apraxia.

Limitations

There are some limitations to the present study. First, disconnection severity in the region-wise interpretation can either refer to the absolute number or the proportion of disconnected fibres. Both variables bear some limitations. We chose to primarily interpret the absolute numbers to avoid the overestimation of complete disconnections of originally faintly connected areas. Second, like in most studies on apraxia, many of the investigated patients also suffered from aphasia. This might create complications in disentangling lesion foci of these symptoms. However, the present patient sample was recruited independent of aphasia diagnosis, it included a wide range of clinical variance both for aphasia and for apraxia, and the multivariate statistical inference approach was utilized to map the clinical variance of the apraxia score. Therefore, it can be assumed that the
analysis indeed significantly reflects the anatomy of apraxia. Notably, this point reflects a general limitation of lesion-behaviour mapping, which is not necessarily overcome by covariate control. However, the complementary use of neuroscientific methods has the potential to alleviate this issue. Third, the present sample was tested in the chronic post-stroke stage at least several months after stroke. This allowed us to also account for chronic long-term changes in structural network integrity. On the other hand, post-stroke brain plasticity might have altered common brain anatomy. The current study provides insights into the hyper-critical parts of the praxis network, whose damage cannot be easily compensated by brain reorganization. Hence, to fully understand the human praxis network, acute neurological studies or studies on healthy subjects would synergize well with the current study.

**Conclusion**

The present study suggests the involvement of frontal, temporal and parietal cortical brain regions connected by short and long association fibres into a complex fronto-temporo-parietal network. A striking finding was the importance of temporal regions and tempo-temporal short association fibres, suggesting a possible hub-like function of the temporal lobe in praxis skills. Further, our results implicated several cortico-subcortical connections, albeit subcortical grey matter structures were rarely implicated in previous lesion mapping studies. It seems that disconnections rather than lesions to subcortical grey matter areas might be relevant for the emergence of apractic symptoms. A future challenge is to deepen our understanding of all the essential cognitive functions underlying apraxia and to identify the parts of the fronto-temporo-parietal network that constitute their neural correlates. As a clinical implication of the present study, our findings suggest that long-term stroke outcome prediction based on imaging data might profit from topographical variables representing white matter damage.

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**Competing interests**

No conflicts of interest are reported.

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**Supplementary material**

Supplementary material is available at *Brain Communications* online.

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