Corticospinal Tract Anatomy and Functional Connectivity of Primary Motor Cortex in Autism

Ruth A. Carper, PhD, Seraphina Solders, Jeffrey M. Treiber, BA, Inna Fishman, PhD, Ralph-Axel Müller, PhD

Objective: Growing evidence indicates that autism spectrum disorder (ASD) stems from abnormal structural and functional connectivity of neural networks. Although diagnostic symptoms are sociocommunicative, motor-related functions (beyond repetitive mannerisms) are also impaired. However, evidence on connectivity at the level of basic motor execution is limited, which we address here.

Method: We compared right-handed children and adolescents (aged 7–18 years) with ASD (n = 44) to matched typically developing participants (TD, n = 36) using magnetic resonance imaging (MRI). Diffusion-weighted imaging and probabilistic tractography measured microstructure of the corticospinal tract (CST). Intrinsic functional connectivity MRI examined whole-brain voxelwise correlations, both with identical precentral gyrus (PCG) seeds.

Results: In the group with ASD, radial and mean diffusivity were increased bilaterally in the CST, particularly in superior segments, and a leftward asymmetry of CST volume detected in the TD group was reversed. Functionally, overconnectivity was found for both left and right PCG with prefrontal, parietal, medial occipital, and cingulate cortices. The group with ASD also showed significantly reduced asymmetry of functional connectivity for both left and right PCG seeds. Finally, in the group with ASD, significant correlations were found for functional overconnectivity of the right PCG seed with anisotropy and mean diffusivity in the right CST.

Conclusion: The findings, implicating both functional and anatomical connectivity of the primary motor cortex, suggest that network anomalies in ASD go well beyond sociocommunicative domains, affecting basic motor execution. They also suggest that even in right-handed adolescents with ASD, typical left hemisphere dominance is reduced, both anatomically and functionally, with an unusual degree of right hemisphere motor participation.

Key Words: autism spectrum disorder, diffusion tensor imaging, probabilistic tractography, functional connectivity MRI, primary motor cortex


Although autism spectrum disorders (ASD) are diagnosed based on socio-communicative impairments and restricted or repetitive behaviors, there is increasing evidence that ASD is associated with abnormal motor-related functions beyond repetitive motor mannerisms, including hypotonia, abnormalities of gait, and dyspraxia or apraxia. Delays in motor development and impairments of gross and fine motor functions have also been reported, which may be present as early as infancy and may persist into adulthood.

In a typically developing (TD) brain, voluntary movements are controlled by primary motor cortex (M1) and its outputs through the corticospinal tract (CST), with indirect modulation through the basal ganglia and cerebellar circuits. Motor abnormalities in ASD are often interpreted as involving the basal ganglia and/or cerebellum, whereas evidence related to the motor control system itself remains limited. In 8- to 12-year-olds with ASD, increased white matter volume in the motor/premotor area (precentral gyrus) was found to be predictive of poorer motor function, a reversal of the correlation seen in typically developing children. The authors suggested that this effect primarily reflected local (rather than long-distance) connections. In the present study, we examined connectivity of the motor control system in ASD, including anatomical organization of its output pathway along the CST and functional connectivity of primary motor cortex, using diffusion tensor imaging (DTI) and intrinsic functional connectivity (iFC) magnetic resonance imaging (MRI).

Studies using DTI indicate atypical trajectories for white matter maturation. Toddlers with ASD show increased fractional anisotropy (FA), whereas school-aged children and adolescents show reduced FA, increased mean diffusivity (MD), and radial diffusivity (RD). This suggests an early acceleration of maturation followed by a plateau, further supported by longitudinal findings and similar to what is seen in volumetric studies of gray matter. Abnormalities have been reported in frontal, temporal, and parietal white matter, cerebellar peduncles, and large association tracts. Although none of these studies has systematically examined CST, a few authors have reported limited findings within the tract. Shukla et al. and Jou et al. reported reduced FA and increased...
MD in segments of the white matter “skeleton” corresponding to CST. One region-of-interest study found decreased FA in the posterior limb of the internal capsule bilaterally and in the right inferior CST (cerebellar peduncle), whereas another found increased MD but no effects on FA.

Complementing the investigation of anatomical connectivity through DTI, functional connectivity MRI (fcMRI) assesses functional coordination between spatially distributed brain regions \cite{20}. Intrinsic functional connectivity (iFC)—inferred from interregional cross-correlations of the blood oxygen level-dependent (BOLD) signal—can be detected at rest, in the absence of an overt task. Importantly, iFC patterns correspond to brain networks recruited during specific cognitive processes \cite{29,30} and likely reflect functional networks associated with functional specialization. The iFC patterns are largely consistent with anatomical connectivity \cite{33,34} and are highly reliable across individuals.\cite{27,33,35}

A growing number of fcMRI studies point to widespread abnormalities in interregional connections in ASD, including decreased connectivity within a motor system (M1, thalamus, and cerebellum),\cite{38} partially replicated by a study showing reduced iFC between (pre)motor cortex and thalamus.\cite{39} Another study suggested reduced differentiation between M1 and lower limb regions in ASD.\cite{40}

The present study provides a comprehensive and multimodal investigation of anatomical and functional connectivity of the motor control system in children and adolescents with ASD compared to TD children. We used DTI and probabilistic tractography to examine the major output pathway of M1, the CST, and iFC to examine the functional connectivity of M1.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>ASD(^a) (n = 44, 8 Female)</th>
<th>Mean ± SD</th>
<th>Range</th>
<th>TD(^b) (n = 36, 9 Female)</th>
<th>Mean ± SD</th>
<th>Range</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, y</td>
<td>13.2 ± 2.9</td>
<td>7–18</td>
<td></td>
<td>12.8 ± 2.4</td>
<td>8–17</td>
<td></td>
<td>.50</td>
</tr>
<tr>
<td>Nonverbal IQ</td>
<td>104.5 ± 17.9</td>
<td>69–140</td>
<td></td>
<td>105.4 ± 11.8</td>
<td>83–129</td>
<td></td>
<td>.79</td>
</tr>
<tr>
<td>Verbal IQ</td>
<td>100.7 ± 20.4</td>
<td>56–147</td>
<td></td>
<td>106.1 ± 12.4</td>
<td>73–133</td>
<td></td>
<td>.17</td>
</tr>
<tr>
<td>Full Scale IQ</td>
<td>102.4 ± 18.7</td>
<td>66–141</td>
<td></td>
<td>106.5 ± 11.9</td>
<td>79–132</td>
<td></td>
<td>.26</td>
</tr>
<tr>
<td>SRS Total</td>
<td>79.7 ± 10.4</td>
<td>57–100</td>
<td></td>
<td>41.9 ± 5.0</td>
<td>35–52</td>
<td></td>
<td>&lt;.001</td>
</tr>
<tr>
<td>ADOS Social</td>
<td>9.0 ± 3.5</td>
<td>4–21</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>ADOS Communication</td>
<td>4.3 ± 2.5</td>
<td>0–13</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>ADOS S+I-C</td>
<td>12.6 ± 4.9</td>
<td>1–22</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ADOS Repetitive Behavior</td>
<td>2.2 ± 1.6</td>
<td>0–6</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>ADI Social</td>
<td>19.1 ± 5.9</td>
<td>6–28</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>ADI Communication</td>
<td>15.2 ± 5.9</td>
<td>2–25</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>ADI Repetitive Behavior</td>
<td>6.5 ± 2.3</td>
<td>3–12</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DTI Total Motion Index [TMI]</td>
<td>2.61 ± 4.71</td>
<td>−1.50–18.70</td>
<td></td>
<td>0.87 ± 2.53</td>
<td>−1.97–10.88</td>
<td></td>
<td>.07</td>
</tr>
<tr>
<td>iFC-MRI RMSD</td>
<td>0.10 ± 0.07</td>
<td>0.02–0.30</td>
<td></td>
<td>0.09 ± 0.07</td>
<td>0.02–0.30</td>
<td></td>
<td>.46</td>
</tr>
</tbody>
</table>

Note: ADI = Autism Diagnostic Interview—Revised; ADOS = Autism Diagnostic Observation Schedule; DTI = diffusion tensor imaging; iFC-MRI = intrinsic functional connectivity magnetic resonance imaging; RMSD = root mean square of displacement; S+I-C = Social + Communication subscale; SRS = Social Responsiveness Scale; TD = typically developing.

\(^a\)IQ scores unavailable for 2 participants, SRS unavailable for 1 participant, ADOS scores unavailable for 4 participants, ADIR scores unavailable for 2 participants.

\(^b\)SRS scores unavailable for 2 participants.

METHOD

Participants

All participants with ASD met DSM-V criteria for ASD.\cite{1} Prospective participants with ASD were administered the Autism Diagnostic Observation Schedule (ADOS),\cite{41} and parents were administered the Autism Diagnostic Interview—Revised (ADI-R)\cite{42} with final diagnosis confirmed by an experienced clinical psychologist. Children with known neurological disorders other than ASD (e.g., Fragile X syndrome, epilepsy) were excluded. Prospective TD participants with personal or family history of autism, or personal history of other neurological or psychiatric conditions, were excluded. Additional assessments included the Wechsler Abbreviated Scale of Intelligence (WASI),\cite{43} the Developmental Test of Visual–Motor Integration, 6th Edition (VMI),\cite{44} and the Edinburgh Handedness Inventory.\cite{45} The study was approved by the University of California–San Diego and San Diego State University institutional review boards, with written informed consent and assent provided by all participants and caregivers.

We scanned 108 children and adolescents, excluding 28 because: of: non–right-hand preference (4 ASD, 6 TD), exclusionary finding on MRI or other measures (6 ASD, 1 TD), or incomplete or poor-quality data in multiple imaging modalities (8 ASD, 3 TD; additional exclusions for single modalities [DTI or iFC] described in Results; for quality criteria, see Analysis sections). The final sample included 44 participants with ASD (8 female) and 36 TD individuals (9 female) aged 7 to 18 years, all right-handed (Table 1) and matched for age and IQ.

MRI Data Acquisition

MRI data (GE Discovery MR750 3.0T, 8-channel head coil) included the following: T1-weighted anatomical scan (fast spoiled gradient echo; TR = 8.108; TE = 3.172 milliseconds; flip angle = 8°; resolution = 1 mm\(^3\)); diffusion weighted images (two-dimensional [2D] echo planar imaging [EPI]; 61 noncollinear diffusion directions at \(b = 1,000 \, s/mm^2\), 1 at \(b = 0 \, s/mm^2\); TR = 8500 milliseconds; TE = 84.9 milliseconds; flip angle = 90°; resolution = 1.875 × 1.875 × 2 mm\(^3\));
matching field map to correct inhomogeneities (2D gradient recalled acquisition in the steady state; TR = 1,097 milliseconds; TE = 9.5 milliseconds; flip angle = 45°); functional T2*-weighted EPI (6:10 minute resting-state scan; 185 time points; TR = 2,000 milliseconds; TE = 30 milliseconds; flip angle = 90°; resolution = 3.4 mm isotropic); matching field map (2D GRASS; TR = 614 milliseconds; TE = 6.5 milliseconds; flip angle = 45°). Participants viewed a white fixation cross displayed on a black background throughout the resting state scan.

**DTI Analysis**

Preprocessing used the FMRIB Software Library (FSL, v.5.0)\(^{46}\) and AFNI (v. 2011_12_21_1014),\(^{47}\) including correction of field inhomogeneities, resampling to \(1 \times 1 \times 1 \text{ mm}^3\), removal of non-brain tissue, and eddy current correction. Diffusion tensors, FA, MD, RD, and axial diffusivity (AD) were calculated, and the FA map non-linearly registered to the FMRIB58 FA template. Resulting transformation matrices were inverted and saved to facilitate fiber tracking in native space using standard-space atlases. We assessed all scans for motion through both visual inspection of eddy-corrected data (for signal dropout, image noise, shifts of head placement) and quantification of artifacts.\(^{48-50}\) Scans with visible evidence of moderate or severe motion were excluded. As our study included children as young as 7 years, we did not exclude scans with mild motion. Motion and related artifacts were then quantified according to Yendiki et al.,\(^{50}\) and this motion measure was used as a covariate in all analyses. Mean image translation and rotation applied during eddy correction were recorded, as were severity and frequency of signal drop-outs across slices. These quantities were combined into a total motion index (TMI), which has been demonstrated to reduce spurious findings related to motion when used as a nuisance regressor.\(^{50}\)

Fiber tracking was performed using probabilistic tractography (BEDPOSTX, ProbTrackX)\(^{51,52}\) in native space. A binary seed mask of precentral gyrus (PCG) was derived from the Harvard-Oxford atlas in standard Montreal Neurological Institute (MNI) space, and a cerebral peduncle target mask was generated manually on the standard brain at \(z = -18\) (see Figure S1, available online) with a stop mask 1 slice inferior. One thousand streamlines were initiated from each of the \(\sim 37,865\) seed voxels, giving a probability map of connectivity (0.5-mm step length, curvature threshold = 0.2). A minimum threshold for number of streamlines was applied at the 75th percentile (see Figure S2, available online), and the output binarized (see Figure S3, available online). These masks were applied to the DTI maps to obtain mean MD, AD, RD, and FA values for each CST, whereas mask volumes provided a measure of overall CST size in native space. The CST mask was also separated into 3 segments along its length based on the ICBM-DTI-81 white-matter labels atlas. These corresponded to the following: inferior CST \((-18 \leq z < -4)\) lying inferior to the thalamus; middle CST \((-4 \leq z < 19)\) lying within the internal capsule, adjacent to the thalamus; and superior CST \((z \geq 19)\) located within the corona radiata. Total brain volume (TBV), including all brain parenchyma but excluding CSF, was derived from the eddy-corrected \(b = 0\) image using FSL’s FAST.

**Functional Connectivity Analysis**

Functional connectivity processing used AFNI and FSL. The first 5 frames were discarded, resulting in 180 total whole-brain volumes. Data were slice-time and motion corrected by realigning to the middle time point, corrected for field inhomogeneity, co-registered to the anatomical image, resampled to \(3.0\)-mm isotropic voxels, standardized to the MNI template, and spatially smoothed with an isotropic Gaussian filter to an effective full-width at half-maximum of \(6\) mm, to allow for statistical parametric analysis. Band-pass filtering at \(0.08 < f < 0.08\) Hz isolated frequencies at which intrinsic network-specific blood oxygenation level–dependent (BOLD) correlations predominate.\(^{28}\) Time points with head displacement >1 mm (computed as the root sum of square of displacement

<table>
<thead>
<tr>
<th>TABLE 2</th>
<th>Diffusion Measures in Corticospinal Tract</th>
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<tbody>
<tr>
<td></td>
<td><strong>ASD (n = 31)</strong></td>
</tr>
<tr>
<td></td>
<td><strong>Mean</strong></td>
</tr>
<tr>
<td>TBV (cm(^3))</td>
<td>1,271.95</td>
</tr>
<tr>
<td>CST volume (cm(^3))</td>
<td>\n</td>
</tr>
<tr>
<td>Right</td>
<td>29.05</td>
</tr>
<tr>
<td>FA</td>
<td>\n</td>
</tr>
<tr>
<td>Right</td>
<td>0.4259</td>
</tr>
<tr>
<td>MD (10(^{-3}) mm(^2)/s)</td>
<td>\n</td>
</tr>
<tr>
<td>Right</td>
<td>0.8161</td>
</tr>
<tr>
<td>AD (10(^{-3}) mm(^2)/s)</td>
<td>\n</td>
</tr>
<tr>
<td>Right</td>
<td>1.2065</td>
</tr>
<tr>
<td>RD (10(^{-3}) mm(^2)/s)</td>
<td>\n</td>
</tr>
<tr>
<td>Right</td>
<td>0.6210</td>
</tr>
</tbody>
</table>

**Note:** AD = axial diffusivity; ANCOVA = analysis of covariance; ASD = autism spectrum disorders; CST = corticospinal tract; FA = fractional anisotropy; MD = mean diffusivity; RD = radial diffusivity; TBV = total brain volume; TD = typically developing.
between any 2 time points) and their immediately adjacent time points were censored, as were blocks of time points with <10 usable consecutive images. Mean percentage of data censored from all participants was <1.5%, with no difference between groups (mean_{ASD} = 1.6%; mean_{TD} = 1.2%; p = 0.81). Nuisance variables for linear trend, 6 rigid-body motion parameters (3 rotational, 3 translational), mean white matter (WM) and ventricular (CSF) signals, and their temporal derivatives (band-pass filtered at 0.08 < f < 0.08 Hz) were modeled and removed with regression. WM and CSF masks were derived by combining FSL’s automated tissue segmentation (fsl_anat/FAST) with structural templates of ventricular and subcortical zones and further eroding by 1 voxel. Finally, the root mean square of displacement (RMSD) across the time series was calculated for each participant and used for group matching.

The average BOLD time course was extracted from left and right PCG using identical seeds as for probabilistic tractography, but excluding nongray voxels, and Pearson’s correlations were performed with the time courses of all other brain voxels in each participant. Resulting correlation coefficients were Fisher transformed to z values and entered into 1- and 2-independent-sample(s) t tests to examine within- and between-group iFC effects. Statistical maps were corrected for multiple comparisons to a cluster-size-corrected threshold of p < .05 using Monte Carlo simulation.53

RESULTS
Diffusion Tensor Imaging
Eleven participants were excluded from diffusion analyses for excessive motion or other DTI artifacts (6 ASD, 5 TD); another 11 (7 ASD, 4 TD) received an fMRI but not a DTI scan, leaving 31 participants with ASD and 27 TD participants for the following analyses. Groups did not significantly differ in age, IQ, or TMI (see Table S1, available online).

Repeated-measures analyses of covariance (ANCOVAs) were conducted separately on FA, MD, AD, and RD. Cerebral hemisphere was the within-subjects variable, diagnosis was the between-subjects variable, and age and TMI were covariates. For analyses of CST volume, TBV was also a covariate to control for individual differences in brain size. CST volume showed a significant interaction between diagnosis and hemisphere but no main effects for either variable (Table 2). The interaction was due to a shift in asymmetry of CST volume, with TD participants having larger CST volumes in the left hemisphere, but ASD participants showing the reverse. CST volume was significantly related to TMI (p = .046) but not TBV or age. To further examine hemispheric differences, we calculated a volume asymmetry index (AI; [Left-Right/Left + Right] × 100), which also differed significantly between groups (F1,53 = 4.04, p = .05; Figure 1A). ANCOVA performed on FA values showed no main effects or interactions, although there was a significant relationship to the age covariate (F1,54 = 14.57, p < .001), with FA increasing with age when collapsing across diagnostic group and hemisphere. There was no significant relationship with TMI. Both RD and MD were significantly higher in ASD than TD. Age was a significant covariate for both tests, with RD (F1,54 = 9.83, p = .003) and MD (F1,54 = 6.54 p = .02) values decreasing with age when collapsing across diagnostic group and hemisphere. There were no significant findings for AD. Effects of age within each diagnostic group are further investigated below.

Given the possible confounding effects of motion and the significant relationship found between TMI and CST volume, the above analyses were repeated with tightly motion-matched (p = .9) subgroups of 23 ASD and 25 TD participants. Significant findings were similar (see Table S2, available online).

To better localize group differences along the CST, we separately analyzed superior, middle, and inferior segments (see Table S3, available online). Volume of superior CST (sCST) showed a significant diagnosis-by-hemisphere interaction similar to that for the whole CST (F1,53 = 6.62, p = .01) but no main effects. Significant main effects of group were found for MD (F1,54 = 5.48, p = .02), RD (F1,54 = 6.45, p = .01), and FA (F1,54 = 4.32, p = .04), with MD and RD higher in the group with ASD and FA lower. No significant group differences were found for middle or inferior segments, suggesting that group differences in the main analyses were primarily driven by sCST within the corona radiata.

Reduced FA localized to corona radiata could be a secondary outcome of an increased number of crossing association or thalamo-cortical fibers. To test this, we examined volume fractions of primary (f1) and secondary (f2) fiber components estimated by bedpostX, controlling for age and TMI. Within sCST, f1 was significantly reduced in ASD compared to TD (F1,54 = 5.40, p = .02), but groups did not significantly differ on f2 (F1,54 = 1.91, p = .17), and marginal f2 means were lower in ASD (0.106 ± 0.003) than TD (0.112 ± 0.004). Such findings do not support the likelihood of an
increased volume of crossing fibers. No main or interaction effects were found for hemisphere.

**Functional Connectivity**

Eleven participants (8 ASD, 3 TD) were excluded from iFC analyses for missing data or insufficient image quality, and 3 participants with ASD were excluded to improve motion matching. Therefore, iFC analyses included 33 ASD and 33 TD participants. Groups did not significantly differ in age, IQ, RMSD, or individual motion parameters (see Table S1, available online).

In both groups, both left and right PCG showed functional connectivity bilaterally with the postcentral, middle cingulate, and inferior and middle temporal gyri, supplementary motor area, inferior parietal cortex, and portions of dorsolateral prefrontal cortex, as well as putamen, pallidum, thalamus, and cerebellum (Figure 2; details in Table S4, available online). Direct group comparisons revealed several significant clusters of increased connectivity (ASD > TD) for both left and right PCG with the left inferior prefrontal cortex (middle orbital and inferior frontal gyri), bilateral superior and middle frontal gyri, and left inferior parietal lobule and supramarginal gyrus (see Table S5, available online). Right PCG had additional clusters of overconnectivity with bilateral lingual and pericalcarine gyri, cuneus, precuneus, and bilateral middle cingulate gyrus. A single underconnectivity effect (ASD < TD) was detected for left PCG in the left lateral occipital lobe.

Given the hemispheric differences for both iFC and CST volume findings, 2 asymmetry indices (AIs) were calculated,

**FIGURE 2** Within-group ($p < 1 \times 10^{-6}$, corrected) and between-group ($p < .05$, corrected) statistical maps of functional connectivity of the left and right precentral gyri (PCG; lighter colors represent greater connectivity). Note: ASD = autism spectrum disorder; TD = typically developing.
Functional overconnectivity of right precentral gyrus (PCG) in autism spectrum disorders (average z scores derived from all significant clusters showing overconnectivity effects) is (A) correlated with fractional anisotropy (FA; $r = 0.594, p = .004$) and (B) inversely correlated with mean diffusion (MD; $r = -0.621, p = .002$) of the corticospinal tract (CST; $n = 22$).

1 for each PCG seed. For each hemisphere, we summed the $z$ scores across all voxels within significant connectivity clusters (excluding the seed) from the within-group connectivity maps (threshold: $p < 1 \times 10^{-6}$). We applied the formula $(L-R/L+R) \times 100$ to those values, with L and R being $z$ scores from the left and right hemispheres, providing separate AIs for left and right PCG seeds. One-way ANCOVA covarying for age showed significantly reduced asymmetry in the group with ASD for both left and right PCG seeds. Relationships between structural (FA and MD in left and right CST) and functional (average $z$ scores) connectivity were examined with Pearson correlations separately for each group and for PCG seeds in each hemisphere. In the group with ASD, significant correlations were found between iFC overconnectivity of the right PCG seed and FA ($r = 0.594, p = .004$) and MD ($r = -0.621, p = .002$) of the right CST (Figure 3). No other correlations for either group survived correction for multiple comparisons. iFC asymmetry scores were calculated as described above, with no significant correlations between these and DTI asymmetry indices (CST volume, FA, MD) for either group.

Cross-Modality Correlations
A total of 46 participants (22 ASD, 24 TD) had high-quality data for both diffusion and iFC analyses (see Table S6, available online). Average $z$ scores for areas of significant iFC group differences were calculated, again separating overconnectivity and underconnectivity effects for each seed (left, right PCG). Relationships between structural (FA and MD in left and right CST) and functional (average $z$ scores) connectivity were examined with Pearson correlations separately for each group and for PCG seeds in each hemisphere. In the group with ASD, significant correlations were found between iFC overconnectivity of the right PCG seed and FA ($r = 0.594, p = .004$) and MD ($r = -0.621, p = .002$) of the right CST (Figure 3). No other correlations for either group survived correction for multiple comparisons. iFC asymmetry scores were calculated as described above, with no significant correlations between these and DTI asymmetry indices (CST volume, FA, MD) for either group.

DISCUSSION
Although socio-communicative deficits garner the most attention in autism research, motor abnormalities are also prominent, including repetitive behaviors, atypical gait, and dyspraxia. These deficits are often thought to arise from abnormalities in brain areas that modulate motor behavior, such as cerebellum and basal ganglia, whereas imaging evidence implicating the motor execution system itself remains limited. Here, we examined both whole-brain functional connectivity of the PCG and the
anatomical microstructure of the outputs of the motor execution system, namely the CST, using identical seeds.

Anatomical Compromise of CST

The group with ASD showed significantly greater MD and RD in the CST, both in our complete sample and in a tightly motion-matched subsample, suggesting structural abnormalities similar to those reported for large association tracts and other white matter sectors. Our findings are consistent with previous studies that included part of the CST, using other methods. In addition, we detected reduced FA in superior CST at the level of the corona radiata.

Although the cellular basis of group differences cannot be conclusively determined from tensor measures, possibilities include differences in myelination, axon caliber, intra-axonal structure, or presence of crossing fibers. However, the volume fraction of the secondary fiber compartment was not increased, arguing against an increase in crossing fibers, whereas differences in radial, but not axial, diffusivity suggest reduced myelination or reduced axon number or packing density. Reduced number or packing density (and reduced FA) would be most evident in superior CST, where tightly packed axons begin to fan to reach the expanse of M1, as observed here. Both reduced myelination and reduced axon number could result from reduced CST activity, given the known effects of neural activity on myelination and cell survival.

In the TD brain, FA increases as tracts mature, with connections to unimodal areas peaking earlier than those to multimodal areas. FA changes in CST are not always detected in adolescence, possibly due to an asymptote earlier in childhood. As seen in our partial-correlation analyses, age-related FA increase was nonsignificant in our TD group but robust and significant in our group with ASD, possibly reflecting delayed but ongoing motor maturation.

Another finding in ASD was atypical asymmetry. Our TD group showed larger CST volumes in the dominant left hemisphere than the right, similar to typical right-handed adults, but asymmetry was absent or reversed in ASD, although all participants were right-handed. While absolute measures of tract volume can be affected by image quality, relative volumes and asymmetry ratios, such as those used here, should be more stable and reliable. In addition, rightward shifts of structural and functional asymmetry are common findings in language areas in ASD, including rightward shifts for receptive language as early as 12 months of age. A recent iFC study showed rightward asymmetry shifts in numerous sensorimotor, visuospatial, frontal-executive, and frontoparietal networks, suggesting that atypical functional asymmetry may be pervasive in ASD.

Cortical Functional Overconnectivity of M1

Functional connectivity analyses identified overconnectivity clusters in ASD for both left and right PCG seeds, including in frontal and parietal association areas. Right PCG was also overconnected with bilateral calcarine cortex, cuneus, and bilateral middle cingulate gyrus. This atypical connectivity between primary motor and primary visual cortices may relate to a general increase in visual activity in ASD, as identified in a meta-analysis and to fMRI findings of increased occipito-frontal connectivity.

M1 was predominantly overconnected with brain regions outside the motor network proper, as seen in the TD group, suggesting reduced functional segregation of the motor execution system in children with ASD. By contrast, typical development is characterized by increasing network integration (strengthening connections within neurotypical networks) and segregation. This typical trajectory of network maturation appears disrupted in ASD. Increased M1 iFC observed here is thus in line with recent evidence of reduced or delayed network segregation (or differentiation) in ASD, possibly due to impaired synaptic pruning. Our findings add to growing evidence incompatible with the theory of generally reduced long-distance connectivity in ASD or the more specific hypothesis of frontoparietal underconnectivity.

Functional overconnectivity in the ASD group was more pronounced for right than for left PCG and was correlated with white matter microstructure of right CST. This suggests that right PCG overconnectivity is an aspect of pervasive shifts in structural and functional asymmetry, as discussed above. Functional asymmetry of unilateral PCG seeds also differed, being predominantly ipsilateral in the TD group, whereas in the group with ASD, functional connectivity with contralateral regions was atypically strengthened. Thus, both functional and structural connectivity were atypical in ASD with the most prominent differences related to asymmetry.

Our multimodal analyses benefited from complementary strengths of DTI and iFC to examine different components of the motor execution system. DTI analyses examined the output tract of PCG, but although cortical seeds were identical for iFC, no directly corresponding analysis was possible. Instead, iFC was well suited to examine cortical connections. Despite unavoidable differences, both analyses tapped into the motor execution system, as reflected in correlations detected between our iFC and DTI findings.

Although the core symptoms of ASD lie in the socio-communicative domain, our findings show that systems supporting motor execution are affected with respect to both functional and anatomical connectivity. Remarkably, children with ASD showed both white matter compromise, particularly in superior CST, and predominant functional overconnectivity of the primary motor cortex, suggesting reduced differentiation of motor execution networks. Finally, our findings add to growing evidence of atypical asymmetries in ASD. These appear to reflect general and potentially pervasive anomalies of cerebral lateralization.

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