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Altered brain morphometry in carpal tunnel syndrome is associated with median nerve pathology

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Abstract

Objective: Carpal tunnel syndrome (CTS) is a common median nerve entrapment neuropathy characterized by pain, paresthesias, diminished peripheral nerve conduction velocity (NCV) and maladaptive functional brain neuroplasticity. We evaluated structural reorganization in brain gray matter (GM) and white matter (WM) and whether such plasticity is linked to altered median nerve function in CTS.

Methods: We performed NCV testing, T1-weighted structural MRI, and diffusion tensor imaging (DTI) in 28 CTS and 28 age-matched healthy controls (HC). Voxel-based morphometry (VBM) contrasted regional GM volume for CTS versus HC. Significant clusters were correlated with clinical metrics and served as seeds to define associated WM tracts using DTI data and probabilistic tractography. Within these WM tracts, fractional anisotropy (FA), axial (AD) and radial (RD) diffusivity were evaluated for group difference and correlation with clinical metrics.

Results: For CTS subjects, GM volume was significantly reduced in contralesional S1 (hand-area), pulvinar and frontal pole. GM volume in contralesional S1 correlated with median NCV. NCV was also correlated with RD and was negatively correlated with FA within U-fiber cortico-cortical association tracts identified from the contralesional S1 VBM seed.

Conclusions: Our study identified clear morphometric changes in the CTS brain. This central morphometric change is likely secondary to peripheral nerve pathology and altered somatosensory afference. Enhanced axonal coherence and myelination within cortico-cortical tracts connecting primary somatosensory and motor areas may accompany peripheral nerve deafferentation. As structural plasticity was correlated with NCV and not symptomatology, the former may be a better determinant of appropriate clinical intervention for CTS, including surgery.

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1. Introduction

Carpal tunnel syndrome (CTS) is a common neuropathy associated with median nerve conduction block, as well as pain, numbness and paresthesia in the median nerve innervated territory of the affected hand. CTS is mainly driven by partial deafferentation secondary to the compression of the median nerve within the carpal tunnel (Kiernan et al., 1999). In addition to the peripheral sensorimotor manifestations of CTS, cortical digit representations in the brain display expansion, amplification, and/or shifted locus in the primary somatosensory cortex (S1), as evaluated by both fMRI (Napadow et al., 2006, 2007; Zanette et al., 2006) and MEG (Dhond et al., 2012; Tecchio et al., 2002). These studies suggest that the function of the entire hierarchy of the somatosensory system from the peripheral to the central sites may be altered in CTS. However, it is unclear whether this neuroplastic change extends to morphological reorganization in
2.2. Data acquisition

formed consent was obtained from all subjects.

CTS subjects in order to exclude CTS.

with the Boston Carpal Tunnel Questionnaire (BCTSQ) (Levine et al., 2000). Symptomatology was assessed with median and ulnar sensory nerve conduction velocities (Cadwell Sierra myelopathy, 7) generalized peripheral neuropathy, or 8.) severe the-

entrapment other than median nerve, 6) cervical radiculopathy or arthritis, 3) wrist fracture with direct trauma to median nerve,

were 1) contraindications to MRI, 2) history of diabetes mellitus or major cardiovascular, respiratory, neurological illnesses, rheumatoid myelopathy, 7) generalized peripheral neuropathy, or 8.) severe ther-

ary atrophy. All subjects were examined by a physiatrist at Spaulding Rehabilitation Hospital for eligibility, which included the testing of median and ulnar sensory nerve conduction velocities (Cadwell Sierra EMG/NCS Device, Kennewick, WA). Symptomatology was assessed with the Boston Carpal Tunnel Questionnaire (BCTSQ) (Levine et al., 1999).

HC were also enrolled and evaluated with the same procedures as CTS subjects in order to exclude CTS.

All study protocols were approved by the Massachusetts General Hospital (MGH) and Partners Human Research Committee and in-

formed consent was obtained from all subjects.

2.2. Data acquisition

Structural MRI was obtained on a 3.0 T Siemens Trio (Siemens Medical, Erlangen, Germany) equipped with 32-channel head coil at the MGH Athinoula A. Martinos Center for Biomedical Imaging (Charlestown).

A multi-echo MPRAGE T1-weighted pulse sequence was used (TR = 2530 ms, TE1/TE2 = 1.64/30.0 ms, TI = 1200 ms, flip angle = 7°, FOV = 256x256, slices = 176, sagittal acquisition, spatial resolution = 1 x 1 x 1 mm³). Diffusion-weighted MRI was also obtained (b-value = 700 s/mm², directions = 60, 32-channel coil, FOV = 256 mm, slice thickness = 2 mm, 64slices, voxel size = 2x2x2 mm, TR/TE = 8040/84 ms).

2.3. Voxel based morphometry (VBM) analysis

VBM was accomplished by first preprocessing the structural T1-weighted MRI data using SPM software (SPM8, Institute of Neurology, London, UK) running under Matlab (version 7.7, Mathworks). All individual images were processed in a generative model, which included tissue classification, image registration and bias correction. Nonlinear deformations for warping GM and WM images were deter-

mined by DARTEL (Diffeomorphic Anatomical Registration using Exponentiated Lie algebra: DARTEL toolbox8 (Ashburner, 2007)). The warped GM and WM images were smoothed with a Gaussian kernel (FWHM = 10 mm), modulated, and registered to MNI space (i.e. the final DARTEL template brain).

Voxel-wise general linear model (GLM) analysis was performed contrasting GM volumes from CTS versus HC. Significant differences between CTS and HC were determined with age and gender as regressors of no interest. Statistical maps were corrected for multiple comparisons by false discovery rate (FDR, p < 0.05) on the voxel level. We then ac-

cepted only clusters greater than 100 voxels in size in order to limit the potential for small cluster false positives. In order to avoid possible edge effects at the GM/WM border, we excluded all voxels with GM value lower than 0.2. In order to better understand the association between GM volume and CTS-related pathology, we correlated GM volumes from all subjects with a) NCV and b) symptom severity. For the latter, pain symptoms were summarized as the average of questions 1–5, while paresthesia symptoms (numbness/tingling) were summa-

rized as the average of questions 6–10 from the BCTSQ.

In order to account for variability in the laterality of the more affected hand, the imaging data were also analyzed by mirroring data across the mid-sagittal plane in CTS subjects whose more affected hand was the left hand. This analysis was important for brain regions known to be lateralized relative to peripheral lesions in the somatosen-

sory system such as S1, M1, and ventro-posterolateral nucleus of the thalamus. This mirrored analysis was used to interpret only these regions, while the conventional, non-mirrored analysis was used for all other brain areas.

2.4. Diffusion tensor imaging (DTI) analysis

DTI data processing was performed using the FMRIB Software Library (FDT v2.0, FSL v.4.1.9). Prior to fitting the diffusion tensor model, all datasets were skull stripped (Brain Extraction Tool) and eddy current corrected (eddy_correct). The DTI tensor model was then fit using voxel-wise linear least squares (diftit). Probabilistic tractography was performed in subject-space using Bayesian Estima-

tion of Diffusion Parameters Obtained using Sampling Techniques (BedpostX). In order to perform tracking, seeds were created using the results from VBM analyses contrasting CTS and HC (see above). These MNI-registered regions of interest (ROIs) were transformed to subject space using FMRIB Non-linear Image Registration Tool (FNIRT), which calculated an alignment warp between the subject-

space FA maps and the MNI-space FMRIBS58 FA map. The VBM-derived ROIs in subject space were then radially dilated by 1 voxel to assure intersection with WM and a WM mask was used to exclude GM voxels. Probabilistic tractography (FA threshold = 0.2, max steps 2000, step length = 0.5 mm, 5000 iterations per voxel) was then performed on each subject for each of the seed ROIs. Tracts were nor-

malized across all subjects by dividing each voxel in the tract by the
total number of waypoints counted in the entire tract (waytotal) (Johansen-Berg and Behrens, 2009). These subject space tracts were then warped to MNI space to create a group averaged map, restricted to a 5% threshold of the normalized waytotal, a previously validated methodology (Johansen-Berg and Behrens, 2009).

The tracts determined from all 3 ROIs were used to investigate seed-specific WM connections, and as masks to evaluate any significant group differences (CTS-HC) in DTI metrics including FA, AD ($\lambda_1$), and RD (average of $\lambda_2$ and $\lambda_3$). In WM, AD reflects diffusion parallel to axon bundles and is a marker of axonal integrity (Budde et al., 2009; Song et al., 2003). RD is the magnitude of diffusion orthogonal to the axonal axis and is modulated by axonal membrane properties and myelination along fiber tracts (Song et al., 2002, 2003). FA provides a summary measure of relative difference in AD and RD (degree of anisotropy) and is scaled from 0 (isotropic) to 1 (anisotropic). In order to evaluate if WM microstructure within tracts of interest was associated with CTS-related pathology, we correlated CTS subjects’ FA, AD and RD with median NCV, pain, and paresthesia, as defined above.

The existence of crossing fibers is an important potential confound to DTI estimation. To evaluate the probability of crossing fibers, we calculated the mode of the diffusion tensor, which refers to tensor region of the U-fiber cortico-cortical tract connecting contralesional S1 with M1, we found that FA was negatively correlated ($r = -0.65$), while RD was positively correlated ($r = 0.72$) with NCV (Fig. 4). Within both of these FA and RD clusters, the DTI mode was positive for CTS subjects (FA cluster: 0.23 ± 0.20; RD cluster: 0.02 ± 0.14) and was not correlated with NCV (FA cluster: $r = -0.06$; RD cluster: $r = 0.14$), suggesting that crossing fibers did not influence these results. No correlations to NCV were noted for any other DTI metrics. We also did not find any group differences nor correlations in any other tract space (thalamus and frontal pole seed spaces), nor significant correlation between CTS symptomatology and DTI metrics in any of the tracts noted above.

### 3.2. VBM analyses

VBM analysis revealed that GM volume was reduced in left S1 (Table 1, Fig. 1). This result was more prominent when VBM data were mirrored across the mid-sagittal plane for left hand affected CTS (i.e. reduced GM was in the contralesional S1 hand area). GM volume was also reduced in the right posterior thalamus (consistent with the pulvinar) and right frontal pole (Table 1, Fig. 1). A linear regression analysis in our study sample revealed that NCV was positively correlated to GM volume in contralesional S1 ($r = 0.45$, $p < 0.01$, Fig. 2). We found a weak (trending) correlation within the CTS group ($r = 0.24$, $p = 0.1$), while in the HC group, the correlation was not significant ($r = 0.18$, $p = 0.2$). Neither CTS symptomatology nor symptom duration had significant correlation with GM volume decrease in these ROIs.

### 3.3. Diffusion tensor imaging (DTI) analyses

DTI-based tractography identified a U-fiber cortico-cortical association tract connecting S1 (seed cluster defined from the result above) with pre-central gyrus (putative M1, Fig. 3). WM tracts connecting to this S1 seed also included superior longitudinal fasciculus (SLF), consistent with SLF II (Makris et al., 2005). The right posterior thalamus seed generated tracts consistent with the fornix, stria-terminalis, and inferior fronto-occipital fasciculus. The right frontal pole seed generated a tract consistent with the uncinate fasciculus. There was also slightly sub-threshold evidence for connecting fibers of the inferior fronto-occipital fasciculus and frontal aspects of the corpus callosum.

Within the S1-associated WM tract space, DTI metrics (FA, AD, RD, and mode) did not differ between CTS and HC. However, in the saddle region of the U-fiber cortico-cortical tract connecting contralesional S1 with M1, we found that FA was negatively correlated ($r =-0.65$), while RD was positively correlated ($r = 0.72$) with NCV (Fig. 4). Within both of these FA and RD clusters, the DTI mode was positive for CTS subjects (FA cluster: 0.23 ± 0.20; RD cluster: 0.02 ± 0.14) and was not correlated with NCV (FA cluster: $r = -0.06$; RD cluster: $r = 0.14$), suggesting that crossing fibers did not influence these results. No correlations to NCV were noted for any other DTI metrics. We also did not find any group differences nor correlations in any other tract space (thalamus and frontal pole seed spaces), nor significant correlation between CTS symptomatology and DTI metrics in any of the tracts noted above.

### 4. Discussion

This multimodal study investigated CTS-associated morphological reorganization in the brain’s GM and WM, and determined its association with altered peripheral nerve function. Our main finding was that CTS subjects demonstrated reduced GM volume in somatosensory (S1), visuomotor (thalamic pulvinar), and multisensory integration/cognitive (frontal pole) processing regions. GM reduction in S1 was specific to the contralesional cortical representation of the hand region. Moreover, GM volume in S1 was significantly correlated with NCV. Thus, the slower the median NCV across the wrist, the more pronounced the S1 GM volume reduction. We then used DTI to evaluate WM microstructure in the specific WM tracts connected to this S1 region. Interestingly, we found that NCV was negatively correlated with FA, and positively correlated with RD, within the U-fiber cortico-cortical tract connecting contralesional S1 with M1. Our results suggest that CTS, a chronic peripheral neuropathy, also results in structural remodeling and neuroplasticity in the brain, which is closely linked to reduced peripheral nerve conduction.

#### 4.1. Reduced GM volume in CTS

While ambiguity exists as to whether GM reductions represent degenerative atrophy or neuroplastic reorganization (maladaptive or compensatory), reduced GM volume in CTS subjects is consistent with findings observed in other chronic pain conditions in humans (Apkarian et al., 2004, 2011; Ass-Sanie et al., 2012) and animal models (Seminowicz et al., 2009), and may also relate to previous VBM findings of use-dependent plasticity (Draganski et al., 2006). In fact, S1 functional plasticity has been correlated with both diminished GM volume and changes in the diffusion tensor that suggest the growth
of new lateral connections in spinal cord injury patients (Henderson et al., 2011). These findings are interesting as we have previously shown that CTS patients also demonstrate functional neuroplasticity within S1 representations of median nerve innervated territories (Napadow et al., 2006). However, it should also be noted that not all GM morphometry studies demonstrate GM volume decrease in chronic pain patients. For instance, a recent study noted that patients suffering from temporomandibular disorder (a non-neuropathic pain syndrome) did not show any GM decrease, while trigeminal neuropathic pain patients did demonstrate GM decrease in the thalamus and other brain regions (Gustin et al., 2011). These results are consistent with our demonstration of GM volume decrease in CTS patients, as CTS can also be characterized as a neuropathic pain disorder.

4.2. Reduced GM volume in S1

In CTS, the median nerve is compressed at the carpal tunnel, resulting in ischemia (Seiler et al., 1989), inflammation, and elevated tunnel pressure. This pathophysiological sequence of events triggers altered afferent input from affected digits, and decreased nerve conduction along median nerve sensory fibers (Kiernan et al., 1999). Thus, S1 GM decreases in CTS may be the consequence of partial deafferentation or desynchronization in S1 inputs, and Taylor et al. recently found that peripheral nerve transection (a more severe form of deafferentation) was also associated with reduced GM in contralesional S1 (Taylor et al., 2009). However, this previous study did not associate contralesional S1 GM reductions with slowing of peripheral conduction velocities, an important finding in our study that strongly links peripheral pathophysiology with altered brain morphology. While histological changes supporting this structural plasticity are unknown, a number of animal studies have demonstrated that use-dependent structural plasticity is associated with GM increase mediated by neuronal (Lerch et al., 2011), and dendritic spine remodeling (Thomas and Baker, 2012; Xu et al., 2009). In turn, GM reduction has been hypothesized to be mediated by events ranging from neuronal or glial death (May, 2008) to loss of dendritic spine density (Metz et al., 2009).

As GM reduction in S1 was specifically associated with decreased impulse conduction along the median nerve, similar to what we found for functional brain reorganization in CTS (Napadow et al., 2006, 2007), we propose that cortical reorganization in CTS extends to structural change, triggered by chronically altered peripheral afference. Interestingly, GM decreases in contralesional S1 correlated with reduced median NCV and not symptomatology. Multiple CTS studies have found that median nerve conduction does not correlate well with symptomatology (Green et al., 2012; Mondelli et al., 2000). Symptoms such
as pain and paresthesias can fluctuate and are self reported. On the other hand, NCV is an objective electrophysiological marker evaluating the severity of an impaired median nerve, and was directly correlated with reduced GM in contralesional S1.

4.3. WM microstructure in S1/M1 cortico-cortical tract associated with NCV

We further found that greater FA and lower RD in a saddle region of the cortico-cortical U-fiber WM tract connecting S1 to M1 was correlated with NCV. This result suggests that enhanced axonal coherence and myelination which can produce lower RD (Song et al., 2003, 2002) within the WM connecting the pre- and post-central gyri, is also specifically associated with reduced peripheral nerve function. Myelination in the CNS can be induced or inhibited by increased or decreased neuronal activity, respectively (Demerens et al., 1996). Thus, WM microstructure in association with NCV may represent compensatory mechanisms for facilitating communication between S1 and M1, S1/M1 communication, as well as S1 communication with premotor and other prefrontal sensorimotor integration centers via the SLF (which was also found to be connected to our specific S1 seed), is critical for fine motor control (Shinoura et al., 2005), a functional deficiency in CTS (Radwin et al., 2004). Additionally, previous studies have suggested that GM volume reduction occurs secondary to functional brain plasticity (Seminowicz et al., 2011). Our previous studies have noted significant functional plasticity for somatosensory afference reaching S1 (Napadow et al., 2006, 2007). While confirmatory evidence awaits longitudinal studies, we suggest that the GM and WM changes are downstream from the previously noted functional neuroplasticity.

4.4. Reduced GM volume in thalamus

Reduced GM was also found in the right thalamus, consistent with the pulvinar nucleus. The pulvinar is a higher order relay important for cortico-cortical communication in the visual system (Sherman, 2007) and may also contribute in visuomotor integration (Grieve et al., 2000). Deprivation of afferent input leads to GM reduction in the posterolateral thalamus (Draganski et al., 2006), and other thalamic nuclei, such as the pulvinar, may also be affected due to thalamothalamic interactions (Elias et al., 2012). Reduced pulvinar GM may relate to disrupted visuomotor integration underlying the deficient psychomotor performance noted in CTS (Radwin et al., 2004). DTI analysis found that WM tracts leading to/away from this pulvinar region were consistent with the fornix, stria-terminalis, and inferior fronto-occipital fasciculus, tracts likely to relay visuospatial information to prefrontal cortical regions supporting executive control over fine motor commands (Grieve et al., 2000). Interestingly, pulvinar involvement in pain was also recently highlighted by Sprenger et al., who found that lesions in the ventral posterior/pulvinar border

Fig. 3. Significant clusters from the VBM analysis served as seeds to evaluate the WM tracts connected to regions showing reduced GM. [Left column] To create the seeds, significant VBM clusters (red) were dilated and masked by their WM intersection (green). [Right column] Probabilistic WM tractography identified that the contralesional S1 VBM cluster was mainly connected to contralesional M1 via a U-fiber cortico-cortical association tract. The right posterior thalamus (pulvinar) seed generated tracts consistent with the fornix, stria-terminalis, and inferior fronto-occipital fasciculus. The right frontal pole seed generated a tract consistent mainly with the uncinate fasciculus.
zone most readily predicted whether or not post-thalamic stroke patients developed pain (Sprenger et al., 2012).

4.5. Reduced GM volume in frontal pole

Reduced GM was also found in the frontal pole, which appears to be important for monitoring cognitive and motor outcomes as well as multi-sensory integration and executive motor control (Tsujimoto et al., 2010). Decreased GM in the frontal pole may contribute to impaired executive control of motor responses demonstrated in psychomotor testing (Radwin et al., 2004). However, our DTI analysis found that WM tracts leading to/away from this frontopolar region were consistent with the uncinate fasciculus, a limbic tract that has been associated with affective and cognitive dysfunction (Tartaglia et al., 2012), and which connects the frontal pole with the medial temporal lobe. Thus, future studies should further explore how affective dimensions of CTS symptomatology and psychomotor disruption relate to reduced GM in the pulvinar and frontal pole, as well as plasticity in WM tracts connected to these regions.

4.6. Limitations

A limitation of our study was that we did not include severe CTS subjects, and future research will need to clarify if severe CTS with motor dysfunction and atrophy will also demonstrate GM decrease in motor processing regions. Additionally, as structural plasticity following chronic pain may be reversible (Seminowicz et al., 2011), future longitudinal studies should evaluate if effective treatment that ameliorates symptomatology and psychomotor dysfunction improves median NCV can also normalize GM volume. Finally, a significant correlation between GM volume and median NCV was only found in the combined CTS and HC group, and may have been partially due to the significant group differences noted between these groups. However, a trending correlation was also noted within the CTS group, suggesting that this relationship was not solely due to the group difference. Ultimately, greater power may be needed for this relationship to reach significance in CTS subjects, who presented with a reduced dynamic range compared to the combined CTS and HC groups.

5. Conclusions

CTS demonstrated significant GM reductions in the contralesional hand area of S1, as well as right frontal pole and pulvinar. S1 GM reduction was correlated with NCV. In addition, WM microstructure within cortico-cortical U-fiber tracts connecting contralesional S1 with M1 was also correlated with NCV. Thus, structural plasticity in cortical GM and WM occurred as a consequence of peripheral neuropathy in CTS, which supports the view that CTS is not just a peripheral disorder but is accompanied by CNS remodeling. Finally, these results underscore the importance of nerve conduction studies in determining appropriate clinical interventions, including surgery.

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