FUNCTIONAL CONNECTIVITY, TISSUE MICROSTRUCTURE AND T2 AT 11.1 TESLA DISTINGUISHES NEUROADAPTIVE DIFFERENCES IN TWO CONTUSIVE BRAIN INJURY MODELS IN RATS

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Short title: Neuroimaging of contusion brain injury rat models

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Acknowledgements: This study was funded by a National Institute on Neurological Disorders and Stroke grant (UG3 NS106938, PI: Kevin Wang). The contents of this manuscript are solely the responsibility of the authors and do not necessarily represent the official views of the funding agencies. This work was performed in the McKnight Brain Institute at the National High Magnetic Field Laboratory's AMRIS Facility, which is supported by National Science Foundation Cooperative Agreement No. DMR-1644779 and the State of Florida.

ABSTRACT

The damage caused by contusive traumatic brain injuries (TBIs) is thought to involve breakdown in neuronal communication through focal and diffuse axonal injury along with alterations to the neuronal chemical environment, which adversely affects neuronal networks beyond the injury epicenter(s). In the present study, functional connectivity along with brain tissue microstructure coupled with T2 relaxometry were assessed in two experimental TBI models in rat, controlled cortical impact (CCI) and lateral fluid percussive injury (LFPI). Rats were scanned on an 11.1 Tesla scanner on days 2 and 30 following either CCI or LFPI. Naive controls were scanned once and used as a baseline comparison for both TBI groups. Scanning included functional magnetic resonance imaging (fMRI), diffusion weighted images (DWI), and multi-echo T2 images. fMRI scans were analyzed for functional connectivity across laterally and medially located region of interests (ROIs) across the cortical mantle, hippocampus, and dorsal striatum. DWI scans were processed to generate maps of fractional anisotropy, mean, axial, and radial diffusivities (FA, MD, AD, RD). The analyses focused on cortical and white matter (WM) regions at or near the TBI epicenter. Our results indicate that rats exposed to CCI and LFPI had significantly increased contralateral intra-cortical connectivity at 2 days post-injury. This was observed across similar areas of the cortex in both groups. The increased contralateral connectivity was still observed by day 30 in CCI, but not LFPI rats. Although both CCI and LFPI had changes in WM and cortical FA and diffusivities, WM changes were most predominant in CCI and cortical changes in LFPI. Our results provide support for the use of multimodal MR imaging for different types of contusive and skull-penetrating injury.

INTRODUCTION

Traumatic brain injury (TBI) afflicts millions worldwide, with mortality rates over 30%[1]. Although mild-to-moderate repetitive closed head injuries are the most prevalent among the different types of TBI, focal contusions and skull penetrating injuries caused by forceful blows, jolts, bumps to the head, neck, or spinal cord result in devastating outcomes. The long-term, lifelong sensorimotor, emotional, and cognitive impairments, and permanent disruptions to brain function and structure, with severe TBI exert a heavy emotional and financial toll on patients and caregivers. The heterogeneity of damage to the brain with severe TBI makes it difficult to clinically manage. The pathophysiology of contusive injuries involves mechanistically heterogeneous and complex cellular and molecular mechanisms extending beyond the primary site of injury. These so-called secondary injuries can involve aberrant inflammatory neuroimmune activity, diffuse axonal injury (DAI) and other forms of white matter (WM) damage, blood-brain barrier (BBB) disruption, microvasculature bleeds, intracranial hematomas, cerebral and subdural hemorrhage and edema, epileptiform activity, and other adaptive central nervous system (CNS) pathologies[2]. Clinical TBI treatment and management relies on the earliest possible interventions to curb the progression of these secondary injuries[3]. Despite the reported neuroprotective effects of experimental drugs in animal models of TBI, none have proven effective in clinical trials[4]. This failure can be attributed in part to a lack of data on *in vivo*, minimally invasive translational CNS markers to bridge preclinical mechanisms to clinical TBI[4].

Functional neuroimaging offers translational data that may help close in on clinically relevant mechanisms of disease progression and behavioral recovery in TBI, especially when analyzed together with other quantitative magnetic resonance imaging modalities. For instance, neuroimaging of a diverse cohort of mild-to-severe TBI identified stronger deactivations of the default mode network (DMN) regions during cognitive task performance and a negative relationship between posterior cingulate functional connectivity and mean diffusivity (MD) in splenium[5]. Similar brain areas were observed to have reduced connectivity following mild TBI[6],

although hyper-connectivity within DMN regions has also been reported[7] and is associated with metacognitive recovery following moderate to severe TBI[8]. Both increases and decreases in functional connectivity between limbic, sensorimotor, and association cortical areas have been observed over extended periods following moderate to severe TBI[9], and in mild concussive injuries[10]. Individuals sustaining sports-related mild concussions have decreased long-distance and increased local functional connectivity[11]. Thus, despite heterogeneity in the extent of direct or secondary damage caused by TBI, consistency can be said to emerge from results showing temporally evolving functional connectivity reductions and increases across cortical subregions. Whether or not such changes are specifically restricted to cognitive, motor, sensory, or association cortices over time remains unclear. Animal studies recapitulating mild concussive injuries caused by sports injuries or automobile accidents are necessary to model prevalent forms of TBI in humans[12,13]. Like clinical TBI, however, these produce diffuse and heterogenous pathology that, while invaluable to determining construct validity of preclinical TBI models, have so far shown limited ability to reveal fundamental mechanisms involved in cortical reorganization observed with contusions and penetrating cortical injuries. Preclinical studies using controlled cortical impact (CCI) and lateral fluid percussive impact (LFPI) models consistently demonstrate functional reorganization of the cortex near and distant from the TBI epicenter[14]. The large-scale functional network reorganization continues over 30 to 60 days[15,16] and may involve excitatory-inhibitory rebalancing following TBI[17]. These cortical changes are likely accompanied by changes in WM microstructure[18] and chemical changes measurable by tissue T2 relaxometry[19]. The objective of the present study was to assess functional connectivity together with brain tissue microstructure and T2 relaxometry in CCI and LFPI models to evaluate salient similarities and differences between these models.

MATERIALS AND METHODS

Subjects

Female and male Sprague-Dawley rats (220-300g) were obtained from Charles River Laboratories (Raleigh, NC, USA.). Rats were housed in sex-matched pairs in a temperature and humidity-controlled vivarium with a 12 h light cycle (lights on at 0700 h) and food and water provided ad libitum. Rats were randomly assigned to one of three experimental conditions: controls (15 male and 8 female rats), CCI (19 male and 12 female rats), and LFPI (7 male and 5 female rats). A subgroup of CCI rats was imaged 2 days post-injury (16 male and 7 female rats) and another group (3 male and 5 female rats) at 30 days. All LFPI rats were imaged 2 days post-injury, and 7 males were scanned once again at 30 days. All procedures received prior approval from the Institutional Animal Care and Use Committee of the University of Florida and followed all applicable NIH guidelines.

Controlled Cortical Impact

The CCI procedure was carried out using a Leica Impact One device (Leica Microsystems Inc). Anesthetic levels were induced with 4% isoflurane gas mixed with 100% oxygen and maintained under 2% isoflurane for the rest of the procedure. The body temperature was regulated to 37°C using a thermal pad while rats were prepared for surgery on a stereotaxic frame. A parasagittal craniectomy (center at anteroposterior [AP], -4.0 mm; L, +2.8 mm from lambda) 5 mm in diameter was performed to expose the brain and allow impactor tip access to the cortical surface. The impactor had a 4-mm flat-face tip. CCI at a depth of 1.5 mm at 4 m/s and a dwell time of 240ms was carried out. All injuries occurred in the right hemisphere. The surgical area was sutured, and recovery was monitored by tail pinch and righting reflexes. Control rats were naïve and received no sham surgical procedures.

Lateral Fluid Percussion Injury

To model diffuse and focal, open-head TBI, and as part of the translational outcomes project in neurotrauma (TOP-NT), we used the rat LFPI model. LFPI is a well-characterized model of TBI that captures clinically relevant deficits and pathologies[20-22]. LFPI also applies a localized, mechanical insult through a parasagittal craniectomy; however, LFPI is administered via a pressurized fluid pulse (2.0 ±0.2atm). This injury results in a moderated diffuse and focal injury on the right sensorimotor cortex and expansive and progressing white matter injury[2]. Even in cases where limited histopathology is observed, LFPI subjects exhibit manifold neurobehavioral deficits[21,23]. This is consistent with the literature where the majority of TBI cases do not present overt pathology on imaging assessments, yet patients experience lasting and debilitating symptoms.

Magnetic Resonance Imaging

Images were collected in an 11.1 Tesla MRI scanner (Magnex Scientific Ltd., Oxford, UK) with a Resonance Research Inc. gradient set (RRI BFG-240/120-S6, maximum gradient strength of 1000 mT/m at 325 Amps and a 200 µs risetime; RRI, Billerica, MA) and controlled by a Bruker Paravision 6.01 console (Bruker BioSpin, Billerica, MA). A custom-made 2.5 cm x 3.5 cm quadrature radiofrequency (RF) surface transmit/receive coil tuned to 470.7MHz (^{1}H resonance) was used for B1 excitation and signal detection (RF engineering lab, Advanced Magnetic Resonance Imaging and Spectroscopy Facility, Gainesville, FL). Rats were scanned under a continuous flow of 1.5 % isoflurane (delivered at 0.1L/min mixed with medical-grade air containing 70% nitrogen and 30% oxygen). Respiratory rates were monitored continuously, and body temperature was maintained at 36-37°C using a warm water recirculation system (SA Instruments, Inc., New York).

For each rat, a high-resolution T2 weighted anatomical scan followed by a functional magnetic resonance imaging (fMRI) scan, a diffusion weighted imaging (DWI) scan, and a multi-

echo time image series for T2 mapping were acquired. T2-weighted Turbo Rapid Acquisition with Refocused Echoes (TurboRARE) scans were acquired with the following parameters: effective echo time (TE) = 37 ms, repetition time (TR) = 5 seconds, RARE factor = 16, number of averages = 14, field of view (FOV) of 24 mm x 18 mm and 0.9 mm thick slice, and a data matrix of 256 x 192 and 25 interleaved ascending coronal (axial) slices covering the entire brain from the rostral-most extent of the anterior frontal cortical surface, caudally towards the upper brainstem and cerebellum. Functional images were collected using a single-shot spin-echo echo planar imaging (EPI) sequence with the following parameters: TE = 15 ms, TR = 2 seconds, 300 repetitions, FOV = 24 x 18 mm and 0.9 mm thick slice, and a data matrix of 64 x 48 with 25 interleaved ascending coronal slices in the same space as the T2 anatomical. Respiratory rates, isoflurane concentration, body temperature, lighting, and room conditions were kept constant across subjects. Diffusion images used a 4-shot spin echo EPI readout with TE=18ms, TR=4 seconds, 4 averages, FOV = 24 x 18 mm and 0.9 mm slices and a data matrix of 128 x 96 x 25, 30 directions with a b value = 900 s^2/mm and 4 b = 0 scan, pulse duration = 3 ms, and gradient separation = 8 ms. Multi-slice multi-echo scans were collected with TE=6-60 ms in 6 ms steps per image volume, TR=2 seconds, 2 averages, FOV = 24 x 18 mm with 0.9 mm slices and a data matrix of 128 x 128 x 25.

Functional MRI Processing

Image processing was conducted using software tools available in Analysis of Functional NeuroImages (AFNI)[24], FSL[25], and Advanced Normalization Tools (ANTs)[26] and manual mask drawing and editing functions in ITKSNAP[27]. Masks outlining the brains of rats on fMRI, DWI, and T2 anatomical scans were generated using *3dAutomask* in AFNI. These were used for brain extraction in various steps in the image processing workflows. The processing of fMRI scans included the following steps: (1) timeseries spike identification and removal using *3dDespike* in AFNI, (2) motion correction using *3dvolreg* in AFNI, (3) removal of slow signal

drifts (<0.009Hz) using *3dTproject* in AFNI, (4) identification and suppression of nuisance signals using *fsl_regfilt* in FSL, (5) low-pass filtering (>0.12Hz) and spatial smoothing (0.6mm FWHM), and (6) linear affine followed by non-linear registrations of cropped fMRI scans to cropped anatomical scans using *flirt* in FSL and the *antsRegistrationSyNQuick* script for ANTs. Anatomical co-registration matrices were applied to fMRI scans to align these to a multi-T2 scan template upon which ROIs were created. The template was generated by first using rigid-body and affine co-registration steps and then a final nonlinear symmetric normalization step in ANTs. A total of 52 bilateral ROIs were manually created in ITKSNAP. These spanned in an axial slice arrangement the medial and lateral regions of the rostral-to-caudal cortical mantle, hippocampus and dorsal striatum (**Figure 1A-C**). These ROIs were used for signal extraction for seed based functional connectivity.

This generated 52 individual ROI text files per subject that contained L2-normalized resting state signals as a vector of 300 data points. The timeseries files were used in cross-correlations and in calculations of Pearson r coefficients for every pairwise combinations of ROIs (using *corrcoef* function in MATLAB). The resulting number of pairwise correlations was 1,274 per subject (after removing 52 self-correlations). Correlation coefficients were Fisher's transform to ensure normality prior to statistical analyses. The 52 ROIs and template were visualized in 3D using BrainNet viewer [28]. Weighted undirected matrices derived from the Pearson correlations were analyzed using Brain Connectivity Toolbox [29] in MATLAB (Mathworks, Natick, MA). Global graph metrics were calculated for edge density thresholds ranging from 2-40%. Global network measures for this density range were converted to area under the curve (AUC) values prior to statistical analyses. Node-specific (ROI) network measures were calculated at a 12% threshold. The functional network analysis was focused on measuring node strength (sum of edge weights/node) in subregions of the cortex, hippocampus and striatum.

Diffusion Weighted Image Processing

ITKSNAP was used to segment contralateral and ipsilateral cortical and white matter ROIs. Diffusion MRI scans were processed using tools available on FSL [30]. DWI scans were first screened for volumes with low signal-to-noise or artefacts. Eddy correction was used for adjusting slight movements during image acquisition and gradient files rotated according to the motion correction vectors. After eddy correction, tensor element reconstruction and estimates of eigenvectors and eigenvalues were performed using weighted least squares regression on *dtifit* in FSL. This last step generated independent images of FA, mean, axial and radial diffusivities (MD, AD, and RD, respectively).

T2 Map processing

T2 parametric maps were created using the MRI processor plugin for ImageJ[31,32]. The simplex minimization method was used for the non-linear fit of multi-TR images to the equation $S_{TE} = S_0 * e^{-TE/T2}$). ITKSNAP was used to segment contralateral and ipsilateral cortical and white matter ROIs, similar to DWI scans. From T2 maps, the mean T2 is calculated per ROI and exported for statistical analysis.

Statistical Analysis

Statistical analyses and data plotting were carried out using GraphPad Prism 9. Functional connectivity data were analyzed using analysis of variance (ANOVA) with repeated measures. A two-stage step-up false discovery rate (FDR) method was used for multiple comparisons correction. DWI scalars and T2 data were analyzed using a two-way ANOVA with Tukey's honest significant difference post hoc test.

RESULTS

Differential Patterns of Recovery of Functional Connectivity in CCI and LFPI

Functional connectivity within and between contra and ipsilateral nodes was evaluated in the medial and lateral areas of the cortex, as well as the hippocampus and striatum, on days 2 and 30 post-injury (ROIs are shown in **Figure 1**). First, contra-to-ipsilateral homotopic connectivity was examined across bilateral ROI representations in the cortex, but no differences were found between controls and TBI rats (data not shown). **Figure 2** highlights the average node strength in 3D connectome maps of control and CCI. The 3D maps illustrate a higher node strength in the contralateral cortex of CCI animals relative to controls. The greater node strength is observed on both days 2 and 30. The 3D maps also show an increase in intra-hemispheric functional connectivity (an increase in the number of node-connecting edges) in the contralateral cortex. A closer evaluation of the CCI group in the heat maps in **Figure 2B** revealed that in the contralateral medial cortex, the rostral and most caudal areas are significantly greater in node strength on days 2 and 30 compared to controls (group x ROI: $F_{102,3468}=2.4$, $p<0.0001$; **Figure 2B** shows FDR corrected pairwise t-test comparisons). The lateral subregion of the contralateral cortex of CCI rats had significantly greater node strength on both days 2 and 30 post-injury compared to controls but in a limited single slice. In the ipsilateral cortex, the medial subregion had significantly lower node strength on both days 2 and 30 compared to controls (**Figure 2B** shows FDR-corrected pairwise t-test comparisons). This was limited to the more caudal areas of the cortex, within the range of the injury epicenter (**Figure 2B**). The CCI-induced reductions in node strength extended to the lateral areas of the ipsilateral cortex (**Figure 2B**).

Rats that received LFPI had increased node strength in the medial and lateral regions of the contralateral cortex (**Figure 3A**). The increased node strength in the contralateral cortex was observed 2 days, but not 30 days, post-injury (group x ROI: $F_{102,2937}=3.3$, $p<0.0001$; **Figure 3B** shows FDR corrected pairwise t-test comparisons). The ipsilateral cortex had significant

reductions in node strength, most prominently in the medial cortical areas, although lateral areas of the posterior region of the cortex also had low node strength in LFPI rats relative to controls (**Figure 3B** shows FDR corrected pairwise t-test comparisons). The observed reductions in ipsilateral cortical node strength in LFPI rats were observed on day 2 post-injury and recovered to control levels in both lateral and medial areas of the cortex by day 30 (**Figure 3B**).

Figure 4 summarizes the functional changes observed in cortical regions of CCI and LFPI rats. Both CCI and LFPI rats had significant contralateral increases and ipsilateral decreases in node strength, which we use here as a surrogate marker of functional connectivity. The observed functional changes were observed in primary sensory areas (e.g., somatosensory, auditory, visual) and primary motor cortex on day 2 post-injury in both models (**Figure 4A** and **4C**). By 30 days after injury, significant decreases in ipsilateral node strength are still observed near the injury epicenter in CCI rats. No differences were observed in LFPI rats.

Distinct White Matter and Cortical Diffusion Changes and 30-day Recovery in CCI and LFPI Models

Figure 5A shows a representative DWI scan collected in a control rat at 11.1 Tesla, illustrating the slice range for the analyzed scalar maps. We analyzed FA, MD, AD, and RD in ipsi and contralateral WM and cortex. The ROIs included the cortical TBI epicenter and its corresponding section of corpus callosum WM tissue. CCI rats had reduced FA in ipsilateral WM (group x day: $F_{1,94}$=29.6, p<0.0001; post hoc tests: day 2 controls vs day 2 CCI p < 0.0001, day 2 CCI vs day 30 CCI, p < 0.0001) and contralateral WM (group x day: $F_{1,94}$=7.3, p=0.008; post hoc tests: day 2 controls vs day 2 CCI p < 0.0001, day 2 CCI vs day 30 CCI, p = 0.005) and in the ipsilateral cortex on day 2 post-injury (group effect: $F_{1,94}$=7.5, p=0.008; post hoc test: day 2 controls vs day 2 CCI p = 0.008) (**Figure 5B**). FA returned to control levels in these brain

tissues by day 30 post-CCI. MD in ipsilateral WM was greater in CCI compared to controls on day 2, but not day 30 (group effect: $F_{1,94}$=11.2, p=0.001; post hoc test: day 2 control vs day 2 CCI, p = 0.01). No differences in MD between control and CCI rats were observed in contralateral WM, and ipsilateral and contralateral cortex (**Figure 5C**). AD in ipsilateral and contralateral WM was reduced in CCI rats on day 30 (group x day (ipsi): $F_{1,94}$=13.1, p=0.0005; post hoc tests: day 2 CCI vs day 30 CCI, p = 0.0001; day 30 control vs day 30 CCI, p = 0.04; group x day (contra): $F_{1,93}$=7.9, p=0.006; post hoc tests: day 2 CCI vs day 30 CCI, p = 0.004; day 30 controls vs day 30 CCI, p = 0.009). No differences in AD between controls and CCI rats were observed in the cortex (**Figure 5D**). RD in ipsilateral WM was reduced in CCI rats on day 30 (group x day: $F_{1,89}$=9.5, p=0.003; post hoc test: day 2 CCI vs day 30 CCI, p = 0.002). No differences in RD between controls and CCI rats were observed in contralateral WM and cortex (**Figure 5E**).

Compared to the control groups, LFPI rats had reduced FA in ipsilateral WM on day 2 post-TBI (group x day: F1,75=7.9, p = 0.006; post hoc test: day 2 controls vs day 2 LFPI, p < 0.0001; day 2 LFPI vs day 30 LFPI, p = 0.009) and in the ipsilateral cortex on day 30 (group effect: $F_{1,75}$=11.7, p=0.001; post hoc test: day 30 controls vs day 30 LFPI, p = 0.01)(**Figure 6A**). No differences in FA between controls and LFPI rats were observed in contralateral WM and contralateral cortex (**Figure 6A**). MD increased compared to controls in ipsilateral cortex on day 30 post-TBI (group effect: $F_{1,75}$=14.6, p=0.0003; post hoc test: day 30 controls vs day 30 LFPI, p = 0.01) and was reduced in contralateral cortex on day 2 post-injury in LFPI rats (group effect: $F_{1,75}$=12.3, p=0.0008; post hoc test: day 2 control vs day 2 LFPI, p = 0.03) (**Figure 6B**). No differences in MD between control and LFPI rats were observed in WM ROIs (**Figure 6B**). AD was lower in contralateral WM (group effect: $F_{1,75}$=9.6, p=0.0003; post hoc tests: day 2 LFPI vs day 30 LFPI, p = 0.03) and greater in the contralateral cortex on day 30 in LFPI rats compared to controls (group x day interaction: $F_{1,75}$=34, p<0.0001; post hoc tests: day 2 LFPI vs day 30 LFPI, p < 0.0001; day 30 controls vs day 30 LFPI, p < 0.0001) No differences in AD between

control and LFPI rats were observed in ipsilateral ROIs (**Figure 6C**). RD was lower in contralateral WM (group effect: $F_{1,72}$=16.3, p = 0.0001; post hoc tests: day 2 controls vs day 2 LFPI, p = 0.04; day 30 controls vs day 30 LFPI, p = 0.02) and contralateral cortex of LFPI compared to controls on days 2 and 30 (group effect: $F_{1,72}$=22.5, p < 0.0001; post hoc tests: day 2 controls vs day 2 LFPI, p = 0.008; day 30 controls vs day 30 LFPI, p = 0.006). A similar difference was observed in ipsilateral WM, although post hoc tests were not significant (group effect: $F_{1,72}$=9.4, p = 0.003) (**Figure 6D**). Compared to the control group, LFPI rats had higher RD on day 30 post-TBI (group effect: $F_{1,72}$=16.1, p = 0.0001; post hoc tests: day 2 LFPI vs day 30 LFPI, p < 0.0001; day 30 controls vs day 30 LFPI, p < 0.0001).

Distinct White Matter and Cortical T2 Relaxation Changes and 30-day Recovery in CCI and LFPI Models

Figure 7A shows a representative T2 map collected in a control rat at 11.1 Tesla, illustrating the slice range for the analysis. We analyzed tissue T2 values (in ms) in ipsilateral and contralateral WM and cortex. Compared to controls, T2 was increased in ipsilateral WM (group x day: $F_{1,53}$=12.4, p = 0.0009; post hoc tests: day 2 controls vs day 2 CCI, p < 0.0001; day 2 CCI vs day 30 CCI, p < 0.0001), contralateral WM (group effect: $F_{1,53}$=7.7, p = 0.008; post hoc test: day 2 controls vs day 2 CCI, p = 0.007) and in ipsilateral cortex (group x day: $F_{1,53}$=6.5, p = 0.01; post hoc tests: day 2 controls vs day 2 CCI, p = 0.004; day 2 CCI vs day 30 CCI, p = 0.04) of CCI rats on day 2 post-TBI. T2 values in these brain areas of CCI rats recovered to control levels on day 30 (**Figure 7B**). No differences in T2 between control and CCI rats were observed in the contralateral cortex. Compared to controls, T2 values in ipsilateral WM (group effect: $F_{1,47}$=12.4, p = 0.001; post hoc tests: day 2 controls vs day 2 LFPI, p = 0.03) and contralateral WM (group effect: $F_{1,47}$=5.8, p = 0.02; post hoc tests: day 2 controls vs day 2 CCI, p = 0.04) were greater in LFPI rats on day 2. T2 in these WM areas of LFPI rats recovered to

control values by day 30 post-injury (**Figure 7C**). No differences in T2 between control and LFPI rats were observed in ipsilateral and contralateral cortex (**Figure 7C**).

DISCUSSION

The results show important similarities and differences in cortical functional connectivity between CCI and LFPI groups. Both CCI and LFPI rats had significantly increased contralateral intra-cortical connectivity at 2 days post-injury. This was observed across a similar section of the cortex in the medial and lateral areas of the cortical mantle. The increased contralateral connectivity was still observed by day 30 in CCI, but not LFPI rats. While we did not include additional measures, based on the existing hypothesis suggesting excitatory-inhibitory rebalancing[14], it is possible that such changes in neuroadaptive processing continue in CCI animals, and it is established in LFPI. As expected, and reported previously in CCI rats by Yang et al[15], we observed functional connectivity reductions in the ipsilateral cortex in both groups. However, once again, we observed that reduced functional connectivity at or near the injury site is no longer observed by day 30 in LFPI rats, but it endures in both medial and lateral cortical areas in CCI rats. The cortical areas affected in CCI and LFPI included cross-modal sensory regions and motor cortex. These results suggest that comparisons between CCI and LFPI might provide insight into potential cortical neuronal adaptive mechanisms that may be important to functional recovery. To determine links between cortical functional changes and tissue microstructure, we assessed DTI-processed DWI scans. Although both CCI and LFPI had changes in WM and cortical FA and diffusivities, WM changes were most predominant in CCI and cortical changes in LFPI. This result could be related to the above proposition that LFPI rats had an appreciable recovery by day 30, possibly in association with a rebalancing of neuronal activity across the sensorimotor cortex. The hypothesized changes could involve axonal modifications to compensate for functional losses, as observed in recovery processes in

other injury models[33-35], and this, in turn, can impact axial and radial diffusivities. In the contralateral cortex of LFPI rats we observed an increased axial diffusivity and reduced radial diffusivity, which is suggestive of an increase in the directionality of fibers. Finally, T2 values in ipsilateral- and contralateral WM and cortex were similar in CCI and LFPI rats. The increased T2 values on day 2 recovered to control levels by day 30. The similarities between ipsilateral and contralateral regions in both models are interesting. While increased edematous tissue at 2 days could increase T2 at the epicenter, it may be plausible that this also infiltrates the contralateral region[19]. Histological assessments, particularly 3D whole-brain methods may be able to provide mechanistic insight.

Changes in cortical excitability have been reported in rats exposed to either CCI or LFPI, and this has been largely attributed to the development of post-trauma epilepsy (PTE). Both spontaneous and evoked discharges were reported in layer 5 pyramidal neurons, which additionally showed increased burst durations during the first 2 weeks post-CCI[36]. Reduced frequency of spontaneous inhibitory postsynaptic currents (sIPSC), possibly involving reduced gamma amino butyric acid (GABA) release locally, has also been reported[17]. Increases in functional connectivity in the contralateral cortex in both LFPI and CCI models may thus correspond to widespread changes in excitability involved in PTE. However, there is evidence suggesting the mechanisms may be more nuanced. For example, decreased functional connectivity in the cortex has been reported in rats also experiencing shorter latencies to seize and increased interictal epileptic discharges (IEDs)[37]. Besides the role altered neurovascular coupling is likely to play in both reductions and increases in functional connectivity following TBI, increased spontaneous firing across multiple neocortical areas should substantially elevate neuronal oxygen demand and increase blood oxygenation level-dependent (BOLD) signal fluctuations. Synchronous BOLD activity is expected to increase non-specifically, and in a widespread manner, in this scenario. Harris and co-workers reported evidence that this hyperexcitability can be reduced by local inhibitory effects of infused potassium in the

contralateral cortex [38]. They posit that this hyperexcitability may be due to the loss of ipsilateral cross-callosal excitatory fibers that modulate contralateral inhibitory neurons (contributing to overexcitability by releasing inhibitory modulation)[38]. The relationship between changes in neuronal excitability and synchronous BOLD activity is not a linear one. Inhibition of mitochondrial function reduces both spontaneous neuronal discharges and functional connectivity, while elevated mitochondrial activity increases neuronal discharges but still reduces cortical functional connectivity[39]. Our neuroimaging results, and behavioral results reported by others[40], suggest significant functional recovery over 30 days or more, further reinforcing the notion of compensatory neuroadaptations that may involve reorganization of neuronal activity in the cortex of CCI and LFPI rats. PTE-related changes may be accompanied by changes in functional network topology (perhaps contributing to behavioral recovery)[14,15].

The observed functional connectivity changes can also be attributed to altered neurovascular coupling[41]. Increased BBB permeability, astrocytosis, synaptic loss, and damage to microvessels are indicative of a failing neurovascular unit. A result of TBI-induced neurovascular changes is an alteration in cerebral blood flow (CBF). CBF changes have been reported early – within hours - post-injury and at 1 year[42,43]. The CBF changes with TBI involve changes in hypercapnia reactivity, suggesting changes in smooth muscle vasodilatory responsiveness in contralateral and ipsilateral cortices[44-46]. At 3 h, CBF is reduced globally by CCI by as much as 85% in ipsilateral and 49% in contralateral cortex[47]. Mild LFPI produced slight but significant differences in CBF and cerebral blood volume (CBV) between ipsilateral and contralateral cortex at 12 hours post-injury[42]. On the other hand, at 1-year post-injury, CCI reduced CBF at the TBI epicenter, including part of the hippocampus. However, no differences in CBF were observed in contralateral cortex[43]. This latter result suggests a certain degree of neurovascular recovery takes place in off-epicenter sites of the cortex, particularly in the contralateral cortex. Thus, functional connectivity changes in these off-TBI regions are likely to involve neuronal activity changes and are not entirely due to altered CBF mechanisms.

Our DTI analysis focused on corpus callosum segments circumventing the TBI epicenter and in the contralateral cortex. We also assessed the cortical epicenter area and corresponding contralateral cortex to evaluate tissue microstructural differences at 2 and 30 days. Based on the literature, the reported changes in DTI scalars, either using a similar ROI analysis or whole-brain statistics, appear highly heterogeneous across laboratories. Using the CCI model, Harris and colleagues reported ipsilateral WM FA reductions and corresponding diffusivity changes at 1 week, which persisted through 4 weeks[18]. The results were supported by histological data showing demyelinated fibers[18]. This contrasts with our present results using an a priori ROI approach and the difference is likely due to their more comprehensive WM tract-based statistics approach. Changes in WM can surpass localized changes near the epicenter, particularly at later stages post-injury[18]. Conversely, CCI-induced reductions in FA in callosal WM recovered to control levels by day 28 post-TBI, but MD persisted to this time point[48]. Others have reported reductions in FA that were observed 6 months after CCI exposure in rats[16]. This was accompanied by decreases in AD and increased RD[16]. The observed reductions in FA (and increased MD) are associated with significant increases in activated microglia marker translocator protein 18kDa (TSPO), *in vivo*[49]. Yet another study using CCI model reports reductions in FA and increases in MD at 2 days but not at 3 h or 7 days or later[50]. While these results call for improvements in the standardization of DWI methods to facilitate cross-laboratory comparisons, they also suggest the possibility that the neuroadaptations in WM carry large inter-subject (and perhaps, intra-subject) variability. The cellular and molecular changes guiding WM restructuring and axonal remodeling following TBI are likely to involve a certain degree of spatial and temporal ‘randomness’, even with focal contusion models, that could make it difficult to consistently measure the same results using DWI/DTI across and within research laboratories. Our results also indicate increases in T2 in ipsilateral and contralateral cortex and WM at day 2 that is no longer observed on day 30. CCI-induced increased T2 correlates with tissue edema and this in turn has been linked with increases in aquaporin-4 in cortex[19]. The

temporal evolution of changes in T2 varies across laboratories as well. Recovery of T2 to control levels in the perilesional or contralateral cortex has been reported at 7 days[50], 14 days[45], and 30 days (present study), or not recovery by a 42nd day of imaging post-CCI[19]. Manninen et al report increases in T2 at day 2 post-FPI and reductions in T2 compared to naïve controls by day 7, with the same effect observed again on day 21[51]. The latter could be a sign of permanent tissue damage at later time points.

In conclusion, our results provide additional support for the use of multimodal MR imaging to determine the *in vivo*, progressive changes in function and tissue microstructure following contusive and skull penetrating injury. Both fMRI and DWI are potentially capable of distinguishing pathological changes between LFPI and CCI models. However, comparisons with the literature strongly suggest the need for methodological standardization, data harmonization, and statistical approaches to facilitate cross-laboratory comparisons, feature extraction, and/or classifications with enough information to evaluate potentially important preclinical findings relevant to clinical TBI[52,53].

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FIGURES

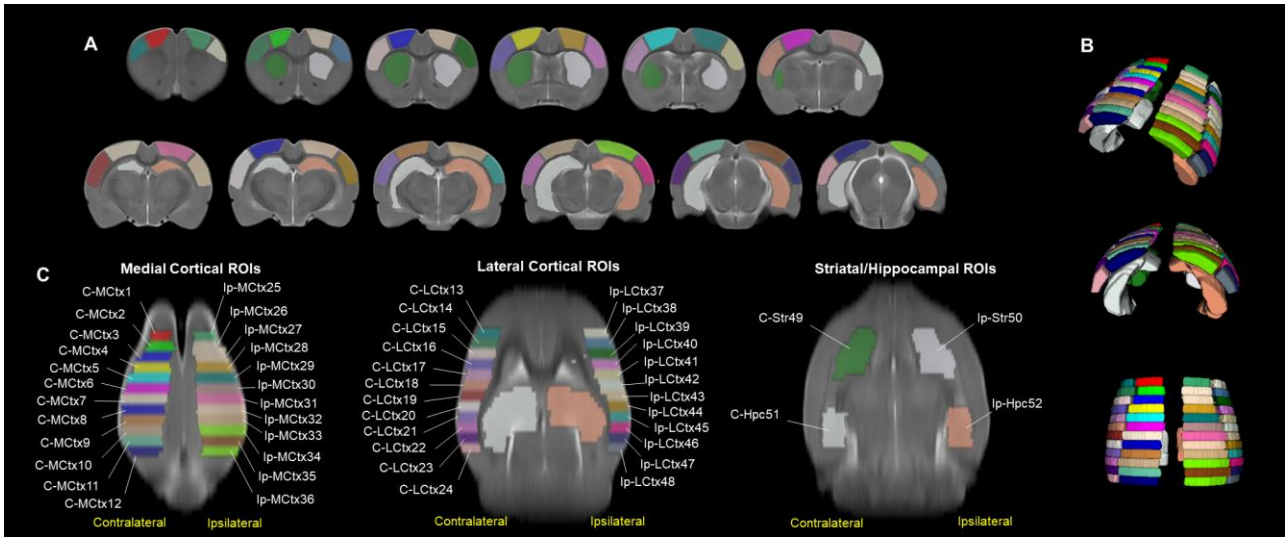

Figure 1. Regions of interest were manually created over a multi-subject template to extract fMRI signals for connectivity analysis. A) Coronal view of the distribution of bilateral cortical, hippocampal and striatal ROIs. B) 3D renditions of the ROIs to illustrate the sampling. C) Nomenclature used to identify ROIs individually across the cortical mantle, medially and laterally in each hemisphere, and in subcortical regions. Abbreviations: C-MC, contralateral-medial cortex; Ip-MC, ipsilateral-medial cortex; C-LC, contralateral-lateral cortex; Ip-LC, ipsilateral-lateral cortex; C-Str, contralateral striatum; Ip-Str, ipsilateral striatum; C-Hpc, contralateral hippocampus; Ip-Hpc, ipsilateral hippocampus.

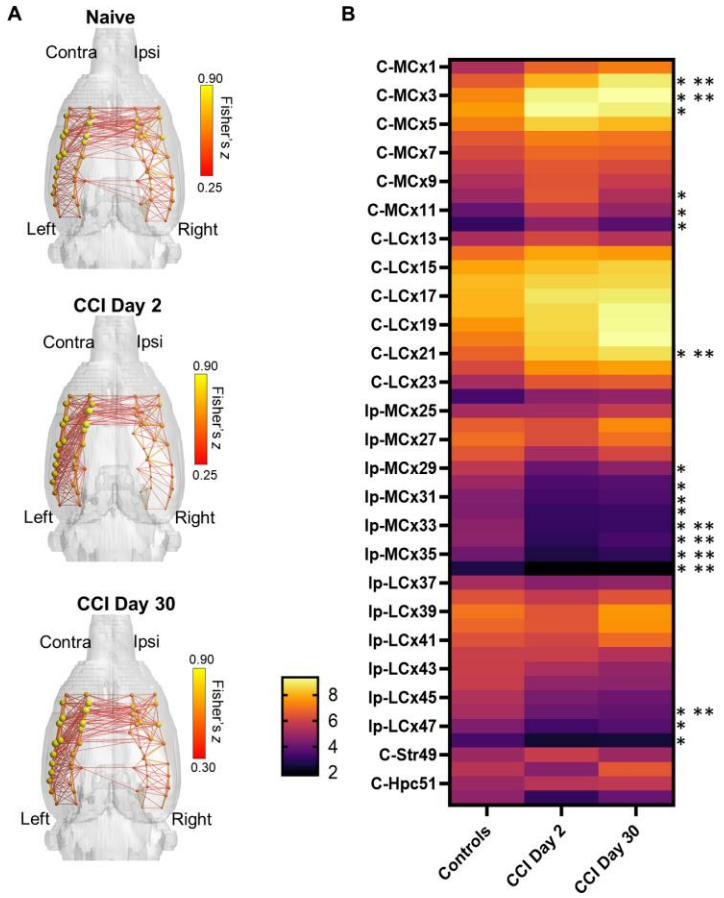

Figure 2. Controlled cortical impact (CCI) increased node strength in contralateral cortex and reduced it in ipsilateral cortex. A) group averaged 3D functional connectome maps illustrate contralateral increases in node strength on days 2 and 30 post-TBI and reductions in ipsilateral cortex. Node size is representative of the calculated ROI node strength and edges (connecting lines) are Fisher's z-transformed Pearson correlation coefficient (maps threshold is 12% edge density; scale bar besides each map indicates z range). B) Heat maps of node strength across the sampled ROIs. Left y axis shows ROI name and x axis group. Asterisks on the right indicate significant difference between CCI day 2 vs controls* and CCI day 30 vs controls (FDR corrected $p<0.05$). The scale bar to the left indicates averaged node strength value.

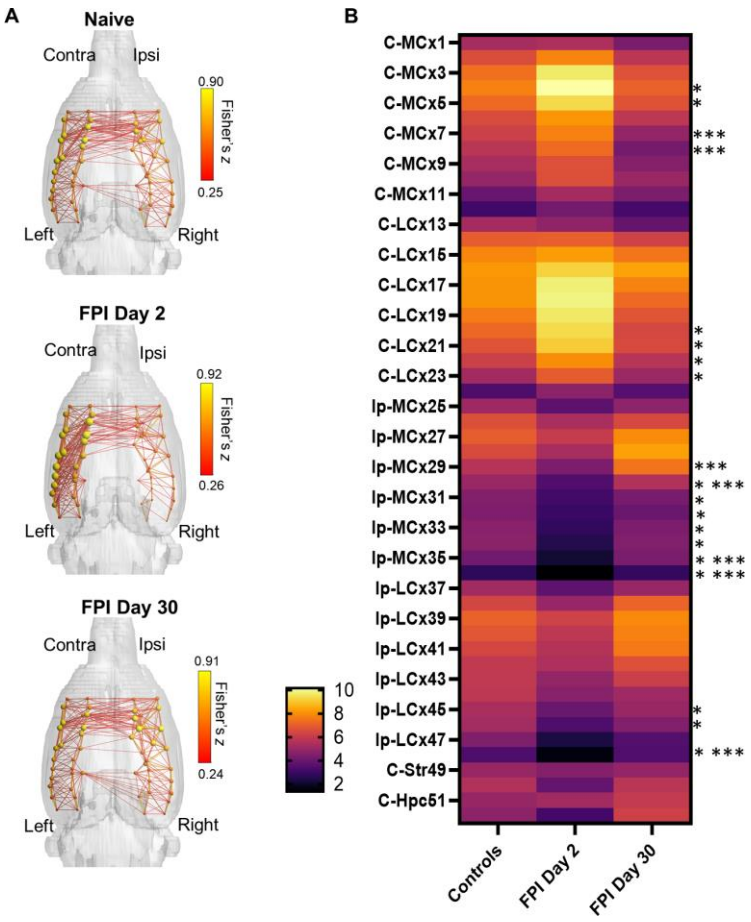

Figure 3. Lateral fluid percussion injury (LFPI) increased node strength in contralateral cortex and reduced it in ipsilateral cortex. A) group averaged 3D functional connectome maps illustrate contralateral increases in node strength on days 2 post-TBI and reductions in ipsilateral cortex. Node size is representative of the calculated ROI node strength and edges (connecting lines) are Fisher's z-transformed Pearson correlation coefficient (maps threshold is 12% edge density; scale bar besides each map indicates z range). B) Heat maps of node strength across the sampled ROIs. Left y axis shows ROI name and x axis group. Asterisks on the right indicate significant difference between CCI day 2 vs controls* and CCI day 30 vs CCI day 2 (FDR corrected $p<0.05$). The scale bar to the left indicates averaged node strength value.

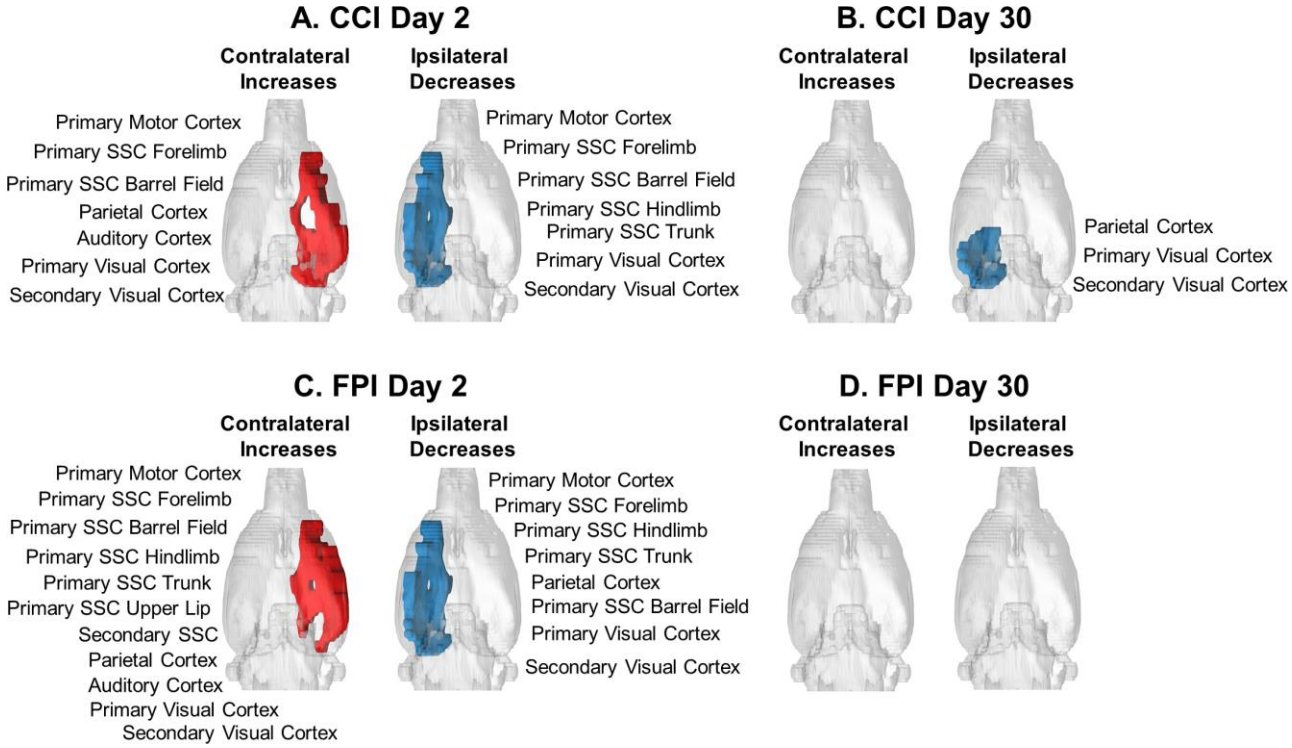

Figure 4. Cortical brain areas affected by CCI vs LFPI. The brain regions were identified by overlaying group averaged ROI's showing significant differences between either CCI or LFPI and controls onto a rat brain template and parcellation[13]. A) Regions with increases and decreases in node strength on day 2 following CCI include sensorimotor areas in both left and right hemispheres. B) By day 30 after CCI, parietal and visual areas shown decreases in ipsilateral cortex. C) Regions affected by LFPI include many of the same cortical areas as CCI on day 2. D) Functional connectivity differences are no longer measured on day 30 following LFPI.

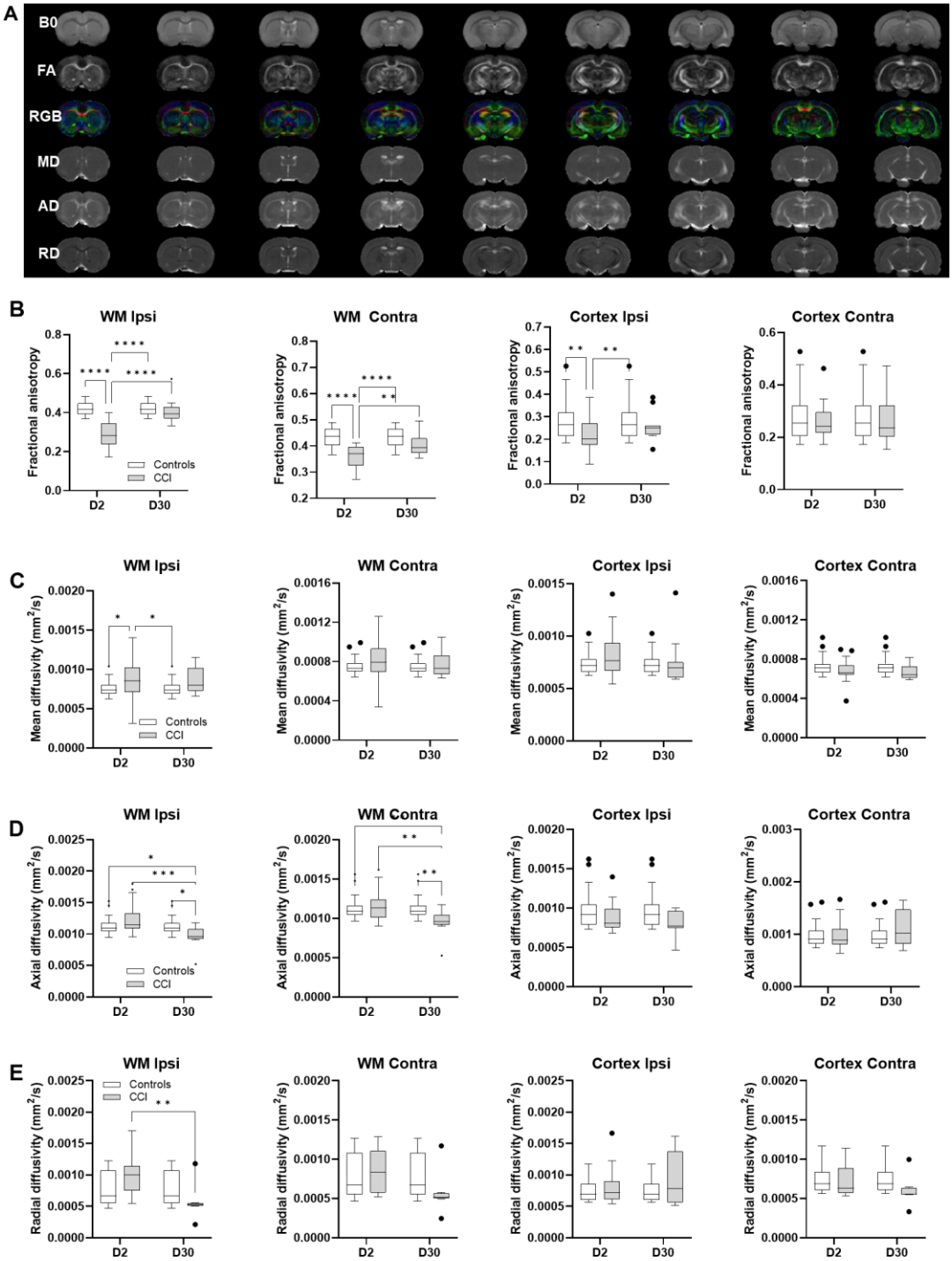

Figure 5. White matter (WM) and cortical tissue diffusion changes at 2 and 30 days following CCI. A) Representative diffusion tensor maps of a control animal. Shown are a B0 image, fractional anisotropy (FA) grey scale map and FA with red green blue (RGB) color coded overlay of 1st eigenvector map, and mean, axial and radial diffusivities (MD, AD, and RD, respectively). B-E) FA, MD, AD, and RD in ipsi and contralateral WM and cortex. Data are shown as box and whisker plots. Lines and asterisks indicate significant differences at $p<0.05$ (Tukey's multiple comparison post hoc test following two-way ANOVA).

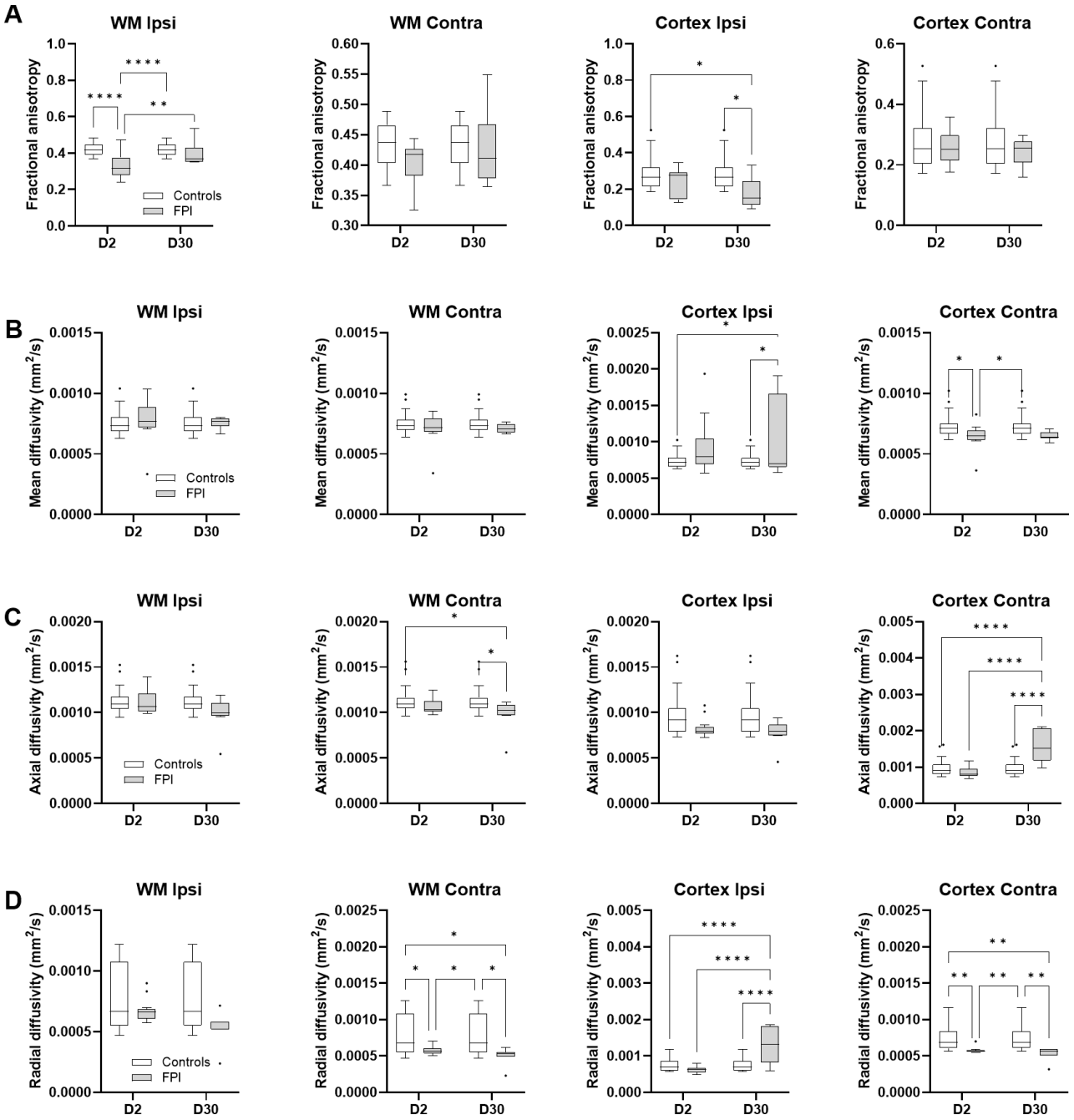

Figure 6. White matter (WM) and cortical tissue diffusion changes at 2 and 30 days following LFPI. A-D) FA, MD, AD, and RD in ipsi and contralateral WM and cortex. Data are shown as box and whisker plots. Lines and asterisks indicate significant differences at $p<0.05$ (Tukey's multiple comparison post hoc test following two-way ANOVA).

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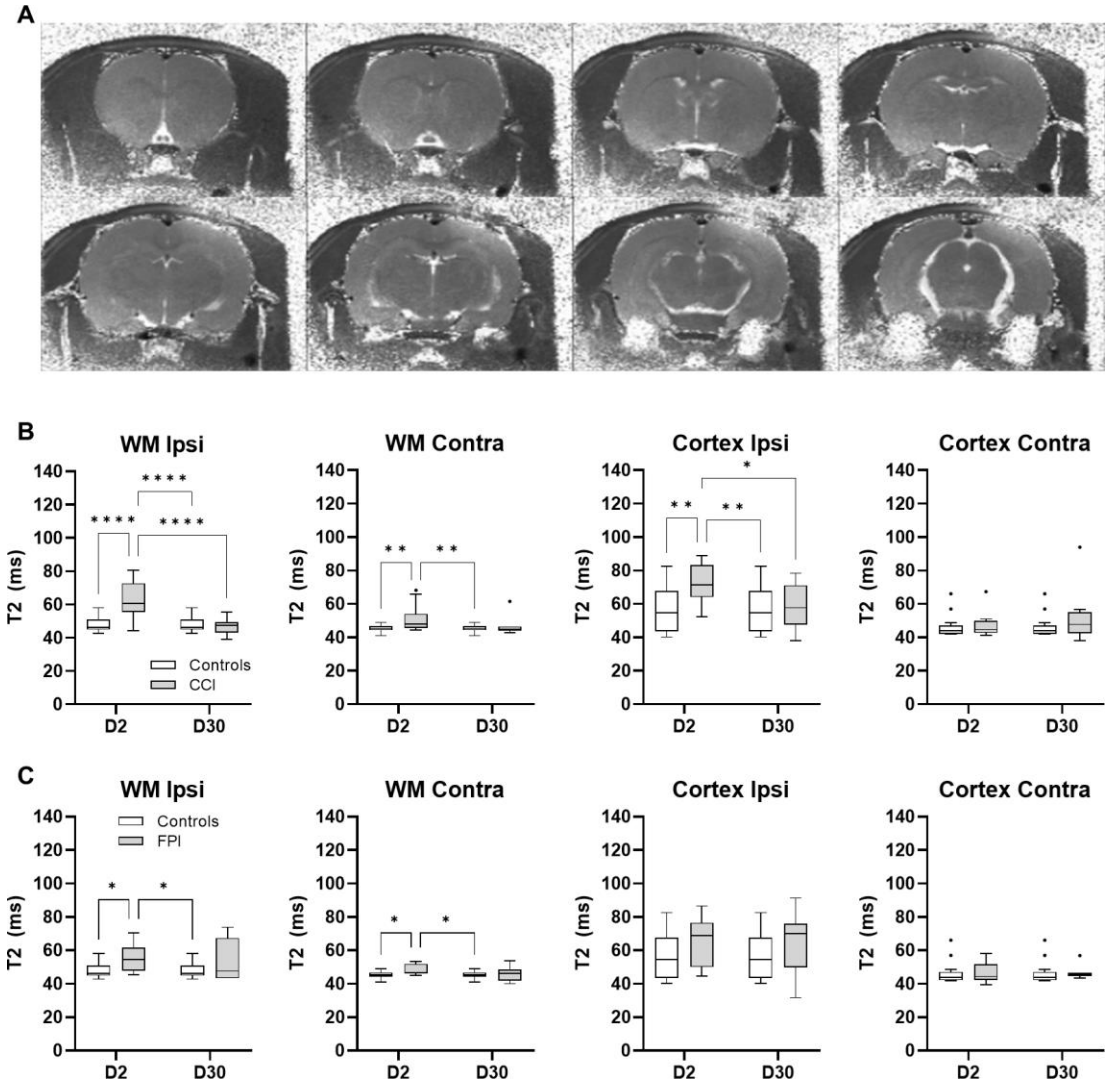

Figure 7. White matter (WM) and cortical tissue T2 relaxation at 2 and 30 days following CCI or LFPI. A) Representative T2 maps of a control animal. A-B) T2 (in ms) in ipsi and contralateral WM and cortex of control, CCI and LFPI exposed rats on days 2 and 30 post-TBI. Data are shown as box and whisker plots. Lines and asterisks indicate significant differences at $p<0.05$ (Tukey's multiple comparison post hoc test following two-way ANOVA).