Repetitive Model of Mild Traumatic Brain Injury Produces Cortical Abnormalities Detectable by Magnetic Resonance Diffusion Imaging, Histopathology, and Behavior

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Abstract
Noninvasive detection of mild traumatic brain injury (mTBI) is important for evaluating acute through chronic effects of head injuries, particularly after repetitive impacts. To better detect abnormalities from mTBI, we performed longitudinal studies (baseline, 3, 6, and 42 days) using magnetic resonance diffusion tensor imaging (DTI) and diffusion kurtosis imaging (DKI) in adult mice after repetitive mTBI (r-mTBI; daily ×5) or sham procedure. This r-mTBI produced righting reflex delay and was first characterized in the corpus callosum to demonstrate low levels of axon damage, astroglisis, and microglial activation, without microhemorrhages. High-resolution DTI-DKI was then combined with post-imaging pathological validation along with behavioral assessments targeted for the impact regions. In the corpus callosum, only DTI fractional anisotropy at 42 days showed significant change post-injury. Conversely, cortical regions under the impact site (M1–M2, anterior cingulate) had reduced axial diffusivity (AD) at all time points with a corresponding increase in axial kurtosis (Ka) at 6 days. Post-imaging neuropathology showed microglial activation in both the corpus callosum and cortex at 42 days after r-mTBI. Increased cortical microglial activation correlated with decreased cortical AD after r-mTBI \((r = -0.853; \ n = 5)\). Using Thy1-YFP-16 mice to fluorescently label neuronal cell bodies and processes revealed low levels of axon damage in the cortex after r-mTBI. Finally, r-mTBI produced social deficits consistent with the function of this anterior cingulate region of cortex. Overall, vulnerability of cortical regions is demonstrated after mild repetitive injury, with underlying differences of DTI and DKI, microglial activation, and behavioral deficits.

Keywords: concussion; diffusion kurtosis imaging; diffusion tensor imaging; microglia; Thy1-YFP

Introduction

Concussion, or mild traumatic brain injury (mTBI), is a blow or jolt to the head resulting in brief alteration in consciousness or post-traumatic amnesia lasting less than 24 h, loss of consciousness lasting less than 30 min, and a Glasgow Coma Score between 13 and 15.1 This type of head impact is common in contact sports, falls, motor vehicle accidents, and as a consequence of blast exposure. mTBI is the most common form of TBI, comprising over 82% of the 339,462 brain injuries sustained by military personnel between the years 2000 and 2015.2 Global incidence of mTBI may be upward of 40 million events per year.3 Although most individuals suffering from mTBI recover over a course of 8–12 weeks, approximately 15% of patients continue to experience symptoms, such as headache, dizziness, insomnia, and difficulties with balance, for much longer. Indeed, over 22% of mTBI patients were still functionally impaired at 12 months post-injury in a recent prospective study.4 Currently available diagnostic tools are not sufficient to accurately predict which patients may suffer persistent symptoms from mTBI. Further, it is critical to determine how repetitive concussive and even subconcussive impacts, that is, below the threshold for mTBI diagnosis, contribute to the long-term consequences of brain injury.

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Noninvasive modalities, such as neuroimaging techniques, have the potential to detect abnormalities in tissue microstructure and could be implemented for longitudinal studies of injury progression to better determine how early abnormalities predict long-term outcomes.

Conventional neuroimaging modalities, such as computed tomography (CT) and standard sequences for magnetic resonance imaging (MRI), do not detect mTBI. However, nuclear medicine imaging and advanced MRI techniques, such as hypercapnia/blood oxygen level dependent or diffusion tensor imaging (DTI), have shown promise in determining prognosis post-mTBI. To date, pre-clinical and clinical studies using DTI demonstrate both lower and higher fractional anisotropy (FA) in white matter, which indicates loss of structural integrity post-injury. Importantly, DTI and DKI changes are sensitive to tissue complexity, such as cortical regions. Therefore, the regions of interest (ROIs) in our longitudinal DTI and DKI studies were analyzed for the corpus callosum and the adjacent medial cerebral cortex. The immunohistochemical analysis of pathology in the corpus callosum was complemented by additional examination of the cortex in Thy1-YFP-16 mice, which have a high density of neurons expressing yellow fluorescent protein (YFP) throughout the cell body, axon, and dendritic arbor. Further, potential functional correlates of the DTI and DKI changes were evaluated using behavioral assessments targeted to the corpus callosum and the adjacent cortical regions within the imaging ROIs.

We show that the combination of DTI and DKI reveals important differences that add to the interpretation of the underlying pathology. Longitudinal studies of a given region across multiple time points aids in detection of abnormalities. Importantly, DTI and DKI changes in the cortical gray matter are associated with microglial activation and later-stage behavioral deficits resulting from repetitive traumatic brain injury.

Methods

Repetitive mild traumatic brain injury model

All animals were treated in accord with guidelines of the Uniformed Services University of the Health Sciences and the National Institutes of Health Guide for the Care and Use of Laboratory Animals. Seven-week-old male C57BL/6 mice or Thy1-YFP-16 mice, both from The Jackson Laboratory (Bar Harbor, ME), were housed in standard cages in a 12-h/12-h light-dark cycle with food and water ad libitum before sham or TBI procedures at 8 weeks of age. As a positive control for tau pathology, P301S mice expressing a mutant form of human tau were obtained from Tony Wyss-Coray (Stanford University, Palo Alto, CA) and used at 5 months of age when tau pathology develops in cortical regions.

The parameters of the mTBI model were based on pilot testing of multiple protocols (data not shown). An Impact One Stereotaxic Impactor (Leica, Wetzlar, Germany) was used to produce mTBI concussive impacts. A series of five impacts separated by 24-h intervals was chosen to target the period of decreased glucose uptake observed 24 h after a single mTBI in rats. Mice were anesthetized with 2.0% isoflurane in O2 and then the hair was shaved and depilated with Nair. Mice were positioned in a stereotaxic frame with rubber stoppers inserted between the external ear canal and ear bar. Impacts were made onto the skin approximately over bregma using a 3-mm-diameter tip (velocity set at 4.0 m/sec; depth of 1.0 mm; dwell time of 200 ms). Sham mice underwent identical procedures for the mTBI mice without receiving impacts. Body temperature was maintained with a warming pad. After each procedure, the duration of apnea and the righting reflex were recorded (Fig. 1). The repetitive injury study included 30 mTBI mice and 30 sham mice. A single impact (s-mTBI) model has been previously characterized and is included as supplementary data (Supplementary Fig. 1) (see online supplementary material at http://www.liebertpub.com) for direct comparison of assessment techniques with the novel s-mTBI model through 6 weeks post-injury.

Magnetic resonance image acquisition parameters

In vivo MRI was performed on a 7 Tesla small animal (20-cm bore) Bruker BioSpec scanner equipped with 12-cm diameter...
650 mT/m gradient coils (Bruker BioSpin GmbH, Reinstetten, Germany). Longitudinal scanning was used to evaluate changes in both the corpus callosum and the overlying cortical gray matter. All mice underwent pre-injury diffusion- and T2- weighted MR imaging and follow-up imaging at 3 days, 6 days, and 6 weeks post-injury. Mice were scanned as yoked pairs consisting of an mTBI mouse with a sham mouse. Two independent cohorts of 3 r-mTBI mice and 3 repetitive sham (r-sham) mice were run from surgery through analysis for a total of 6 r-mTBI mice and 6 r-sham mice scanned at all time points.

Animals were anaesthetized with 1% isoflurane and placed in an MR compatible mouse bed. Images were acquired with a four-channel phased array MR coil after volumetric radiofrequency excitation. A two-dimensional (2D) single-shot echo planar imaging sequence (repetition time/echo time [TR/TE]=4500/50 msec; four repetitions) was used to acquire four unweighted (b = 0 s/mm²) images and seven diffusion-weighted images (b = 400, 800, 1200, 1600, 2000, 2400, and 2800 s/mm²) using a Stejskal-Tanner diffusion preparation² with parameters of δ = 1 msec, δ = 5 msec, field of view (FOV) = 14 x 14 mm², matrix = 80 x 80, slice thickness = 0.75 mm, 18 slices, and 30 noncollinear diffusion gradient directions. A multiecho 2D rapid acquisition with relaxation enhancement (RARE) sequence³ was used to acquire T2-weighted images (TR = 4500 msec, TE = 20, 60, 100, and 140 msec, RARE factor 2, FOV =14 x 14 mm², matrix = 140 x 140 x 18 for s-mTBI and TE = 10, 30, 50, 70, 90, 110, and 130 msec and matrix = 100 x 100 x 18 for r-mTBI, with slice thickness = 0.75 mm). Body temperature was maintained at 36°C by circulating hot water. Respiration and heart rate were monitored throughout each 2-h imaging session.

Magnetic resonance image analysis

Each ROI for analysis was manually drawn on five consecutive coronal slices of the non-diffusion-weighted (b = 0 s/mm²) images (one rostral to bregma, one at the coronal level of bregma, and three caudal to bregma). A single-voxel width was used for the corpus callosum ROI. ROIs in the medial cerebral cortex were drawn in both hemispheres superior to the corpus callosum, while keeping one voxel distance of separation from the corpus callosum and cortical surface to avoid partial volume effects. The peak of the cingulum was used to delineate the border of the medial cortical ROIs. Multiple ROIs were drawn on test scans for standardization before analysis of the study data sets. The individual drawing the ROI for the quantitative analysis was blinded to the animal condition.

Diffusion data were analyzed using TORTOISE software.³⁴ An average non-diffusion-weighted image (b = 0 s/mm²) and all seven diffusion-weighted images were used in the analysis. Diffusion weighted imaging artifacts, including motion, eddy current, and concomitant field distortions, were calculated and combined to enable the image correction to be applied in a single step. This process was used to minimize interpolation errors in the corrected data. Fractional anisotropy (FA) and mean (MD), axial (AD), and radial (RD) diffusivity maps were calculated by the linear least squares method of tensor estimation. Diffusion direction-encoded color (DEC) maps output from the TOROTOISE software were used to demonstrate fiber orientation in the corpus callosum and cortical regions.³⁵ Kurtosis data were pre-processed with TOROTOISE, and each voxel was analyzed using in-house-developed software in MATLAB (The MathWorks, Inc., Natick, MA) according to a previously published method provided by University of Maryland. Essentially, this software routine fits the logarithm of the normalized signal intensity versus the b-values using linear least squares to a quadratic function.²⁶ The signal-to-noise ratio (SNR) of the diffusion-weighted images was >10 for b = 2800 and >50 for b = 0 s/mm² images, respectively, where SNR was calculated as: ±(0.655 x signal)/standard deviation of an ROI in the background of the image). The average ROI (corpus callosum and cortex) signal was a factor of 3 greater than the average background signal for b = 2800 s/mm² and 14 for b = 0 s/mm². Quadratic fits to the raw data were robust for the ROIs interrogated with adjusted R² > 0.99. The mean kurtosis (MK) was calculated by averaging the apparent kurtosis (Kapp) in 30 directions: MK = 1/30 ∑i=130 (Kapp)i.

Axial kurtosis (Kₐ) and radial kurtosis (Kₚ) maps, which characterize the kurtosis along the axial and radial diffusion directions, respectively, were calculated from the three eigenvalues derived from the diffusion kurtosis tensor as described by Jensen and colleagues.³⁶ The corpus callosum and medial cortical ROIs were overlaid onto DTI and DKI maps for further quantitative analysis.

Tissue analysis

Mice were perfused with 4% paraformaldehyde. Brains were further post-fixed by immersion in the same fixative overnight, cryopreserved, and then cut as coronal sections (14 μm). Prussian blue staining combined with pararosaniline nuclear stain (Sigma-Aldrich, St. Louis, MO) was used to detect hemosiderin-laden macrophages indicative of previous microbleeds.

Immunohistochemical analysis was used to assess axon damage based on β-amyloid precursor protein (β-APP)³⁷ (polycrotal rabbit anti-β-APP antibody; 1:100; Life Technologies, Grand Island, NY). Microglia/macrophages were detected based on CD11b (monocrotal rat anti-CD11b; 1:100; Abcam, Cambridge, MA). Astrocytes were evaluated by immunostaining for glial fibrillary acidic protein (GFAP; polycrotal rabbit anti-GFAP; 1:1000; DAKO, Carpinteria, CA). Myelination was estimated by detection of myelin oligodendrocyte glycoprotein (MOG; 1:20, monocrotal mouse anti-MOG immunoglobulin G [IgG] β). The oligodendrocyte progenitor response was evaluated by immunostaining for neural antigen 2 (NG2; polycrotal rabbit anti-NG2 ectodomain; 1:500; gift from William Stallrup, La Jolla, CA). Accumulation of phosphorylated tau was examined using antibodies against

FIG. 1. Characterization of the progression of pathology in the corpus callosum in a repetitive mild traumatic brain injury (r-mTBI) model. (A and B) Post-surgical data for r-mTBI. Apnea was not detected (nd) in r-sham mice (A), but was significantly increased after the first and the second impact in r-mTBI mice (A). The righting time is significantly delayed in r-mTBI versus r-sham mice (B). (C–E) Immunohistochemistry for β-APP is present mainly in neuronal cell bodies in r-sham mice (C). β-APP accumulations are readily detected in damaged axons (arrows) in r-mTBI mice acutely (D), but less marked by 7 days post-injury (E). (F–H) Immunohistochemistry for CD11b identifies microglia, which exhibit a resting phenotype in r-sham mice (F) with progressive activation evident at 24 h (G) and 7 days (H). (I–K) Immunohistochemistry for GFAP identifies astrocytes that have normal morphology in r-sham mice (I) and increasing cells with a reactive phenotype in r-mTBI mice between 24 h (J) and 7 days (K). (L–O) Quantification of immunohistochemistry in the corpus callosum. Axonal profiles with β-APP accumulations were increased very early after r-mTBI mice (L). Astroglosis increases over the first week after r-mTBI (M). Microglial activation also increases significantly over the first week based on area of CD11b immunoreactivity (N) and an earlier reaction is detected by counting activated cells, which have intense CD11b and shorter, thicker processes (O). Dashed lines outline the corpus callosum. Values are mean ± standard error of the mean; n = 30 for post-surgical data; n = 5 per condition for histology data; scale bar (E), (H), and (K) = 50 μm; *p<0.05; **p<0.01; ***p<0.001; ****p<0.0001. β-APP, beta amyloid precursor protein; GFAP, glial fibrillary acidic protein; r-sham, repetitive sham.
anti-phospho-Ser396Tau (mouse monoclonal IgG, 355300; 1:400; Thermo Fisher Scientific, Waltham, MA) and ATX antibody (mouse monoclonal anti-phospho-PHF-tau pSer202+Thr205; 1:400; Thermo Fisher Scientific).

Astrogliosis, microglia/macrophage activation, and myelination were estimated from the area of immunoreactivity for GFAP, CD11b, or MOG within the CC or medial cortex using Metamorph (Molecular Devices, Downingtown, PA) as previously detailed. In addition, microglia/macrophage cells were also quantified by manual counting of the morphological phenotype for resting, activated, or amoeboid stages. Among mice perfused for post-imaging analysis, 1 r-sham and 1 r-mTBI mouse were omitted because of technical concerns limiting quantification across the full set of regions in the double immunostain for CD11b and GFAP at the coronal levels matched to DTI-DKI slice at bregma. Damaged axons were quantified based on β-APP immunoreactivity, which was counted as a single damaged axon for accumulation as a large end bulb or as multiple swellings aligned along a longitudinal axonal profile.

Thy1-YFP-16 mice were used to visualize a high density of neurons in their entirety, including axons, cell bodies, and dendritic processes. Sections were prepared as above for immunohistochemistry. Nuclei were stained blue with 4',6-diamidino-2-phenylindole (DAPI; Sigma-Aldrich) whereas rough endoplasmic reticulum in the cytoplasm of the cell bodies was stained red with a fluorescent version of Nissl histological stain (NeuroTrace red; Thermo Fisher Scientific). Damaged axons were quantified based on YFP in axons with swellings using the same morphological characteristics as for β-APP above.

Bright-field images were captured using an Olympus Nanozoomer (Olympus, Center Valley, PA). Fluorescent wide-field images were acquired on an Olympus IX-70 microscope (Olympus). Optical image fluorescent image stacks were acquired using a Zeiss LSM 700 (Carl Zeiss Microscopy, Thornwood, NY).

**Social interaction assessment**

The social interaction assessment was carried out using a three-chamber sociability test apparatus (Stoelting Co., Wood Dale, IL) as previously described, with minor modification. The test mouse was first placed in the middle chamber and allowed 10 min to explore all chambers for habituation to the apparatus. A small wire enclosure was introduced to each of the side chambers and the mouse was allowed to explore all chambers for 10 min for habituation to the enclosures as novel objects. Social approach was then tested by placing an unfamiliar male C57BL/6J mouse (stranger) into one of the wire enclosures. The test mouse was again placed into the middle chamber and allowed to explore among the three chambers for 10 min. The movement of the test mouse was video recorded to determine the time related to social approach (defined as movement toward, circling, or sniffing of the stranger mouse in the wire enclosure) using ANY-Maze software (Stoelting Co.) as analyzed by an observer blinded to the treatment condition. Mice were run in two independent cohorts of 12 mice that each had 6 mice assigned to each condition. The total assay included 24 mice, with 12 injured and 12 matched sham mice.

**Complex wheel-running assay**

Motor skill learning and bilateral sensorimotor coordination were assessed using a complex wheel-running assay. Running wheels were placed in home cages equipped with an optical sensor to detect wheel revolutions and determine the daily maximal running velocity using Activity Monitoring Software (Lafayette Instruments, Lafayette, IN) over a 3-week period. A regular “training” wheel with evenly spaced rungs was in the cages for 1 week to reduce variability in running behavior. Mice then underwent r-mTBI or r-sham procedures. Training wheels were then replaced with “complex” wheels, which have irregularly spaced rungs. Mice were run in two independent cohorts that each had 6 mice assigned to each condition. The total assay included 24 mice, with 12 injured and 12 matched sham mice. Social interaction cohorts were separate mice from those used on the wheel assay.

**Statistical analysis**

All data analyses were performed with GraphPad Prism software (version 6; GraphPad Software, Inc., La Jolla, CA). Two-way repeated measures analysis of variance (ANOVA) with post-hoc Bonferroni comparison was used to compare TBI with yoked shams across time points for longitudinal analyses. One-way ANOVA was used to compare data among multiple groups for a single condition or time point of tissue analysis, which included three tissue sections/mouse. Post-hoc comparisons were made among the relevant pairs (each TBI model vs. matched sham; single vs. repetitive TBI models). The Student’s t-test was used to compare between two groups for a single condition or time point. Pearson’s correlation was used for analysis of the relationship between radiological and pathological findings. Statistical significance was defined as p<0.05. MRI quantitative analysis was performed independent of post-imaging tissue analysis. All quantification was performed with blinding for the condition.

**Results**

**Initial characterization of repetitive mild traumatic brain injury severity based on progression of pathology in the corpus callosum**

The post-surgical data demonstrate apnea (Fig. 1A) and a delayed righting reflex (Fig. 1B) after r-mTBI compared to the r-sham procedures, which indicates concussive effects across this series of impacts. Mice were prepared for tissue analysis at multiple time points after r-mTBI to characterize the injury severity based on the extent of axon damage, astrogliosis, and microglial activation in the corpus callosum at the coronal level of impact at bregma (Fig. 1C–O). Accumulations of β-APP demonstrated axon damage with impaired fast axonal transport at early times after r-mTBI (Fig. 1C–EL). By 2 h after the fifth impact in r-mTBI mice, β-APP-labeled axons were significantly increased in the corpus callosum (Fig. 1L). Also, astrogliosis (Fig. 1I–K) and microglial activation (Fig. 1E–H) increased in the corpus callosum of r-mTBI mice by 7 days post-injury, as demonstrated by increases in the GFAP-immunolabeled area (Fig. 1M). CD1b-immunolabeled area (Fig. 1N), and density of activated microglia (Fig. 1O). Astrogliosis and microglial activation were less marked at 6 weeks post-injury (Fig. 1MLN) whereas the density of activated microglia continued to be significantly increased from r-sham levels (Fig. 1LO). Prussian blue reaction, which detects hemosiderin-laden macrophages as evidence of previous bleeding, indicated a lack of microhemorrhages whereas the nuclear counterstain demonstrated normal adult tissue cytoarchitecture (Fig. 2).

**Diffusion imaging detects mainly cortical abnormalities after repetitive mild traumatic brain injury**

Longitudinal MRI analysis for both DTI and DKI techniques was tested as a method to identify microstructural changes in the corpus callosum and medial cerebral cortex induced by r-mTBI (Fig. 3). Mainly cortical changes were observed after r-mTBI. The only DTI or DKI parameter with a significant change in the corpus callosum on post-hoc analysis was a decrease in FA at week 6 (Fig. 3D). In contrast, cortical changes in both DTI and DKI were observed after r-mTBI (Fig. 3LV). FA was reduced in the cortex at
FIG. 2. Microhemorrhages are not observed after r-mTBI. Prussian blue was used to detect hemosiderin-laden macrophages that are indicative of previous hemorrhage or bleeding. A positive control for the Prussian blue reaction (arrowed and inset) is shown from a mouse with an injury that was more severe than those used in this study (A). No positive staining was present in either the corpus callosum or cortex in r-sham mice (B) or after r-mTBI at any time points examined (C–F). *n = 3; scale bar = 250 μm. d, days; r-mTBI, repetitive mild traumatic brain injury; r-sham, repetitive sham; w, weeks.

FIG. 3. Diffusion imaging abnormalities after r-mTBI. Corpus callosum (CC) analysis is shown in upper graphs (A–K), for CC region outlined in yellow on upper image) and medial cerebral cortex values are shown in lower graphs (L–V, for cortical region outlined in yellow on the lower image). Each region is shown at the coronal level containing the anterior commissure, which is under the impact site at bregma. r-sham and r-mTBI mice were scanned as yoked pairs with all mice being scanned at four time points—baseline (BL) and post-injury at 3 days, 6 days, and 6 weeks. Five coronal slices (one rostral to bregma, one at bregma = 0, and 3 caudal to bregma) were analyzed. (A–D) FA values in the CC showing all five coronal slices with locations noted as relative to bregma. A significant reduction in FA was noted in r-mTBI mice only at the bregma level at 6 weeks (D). (E–K) CC values in the individual coronal slice at bregma are shown across time points for the full set of parameters analyzed—diffusion tensor imaging (FA, MD, AD, and RD) and diffusion kurtosis imaging (MK, Kₓ, and Kᵧ). No parameters show a significant injury effect that reaches significance at any of the post-injury time points. (L–O) Analysis across coronal slices in the medial cortex shows a significant reduction in FA in r-mTBI mice versus sham near the impact site on days 3 and 6 (M and N). (P–S) Analysis of the individual slice under the impact site across time points for all diffusion parameters shows a significant injury effect in the axial dimension for both AD with diffusion tensor imaging (R) and Kₓ with diffusion kurtosis imaging (U). Values are mean ± standard deviation, n = 6 for each condition at each time point; *p < 0.05; **p < 0.01. AD, axial diffusivity; FA, fractional anisotropy; Kₓ, axial kurtosis; Kᵧ, radial kurtosis; MD, mean diffusivity; MK, mean kurtosis; RD, radial diffusivity; r-mTBI, repetitive mild traumatic brain injury; r-sham, repetitive sham.
days 3 and 6 in coronal slices near the impact site in r-mTBI mice (Fig. 3M,N). However, these FA changes are relatively small, and statistical significance could potentially be affected by variation in the yoked sham cohort during the longitudinal study. Analysis of the DTI and DKI subparameters further refined the cortical changes after r-mTBI. DTI showed significantly decreased AD values at all time points post-TBI (Fig. 3R), with no changes in RD (Supplementary Fig. 3; see online supplementary material at http://www.liebertpub.com). DKI also showed a significant complementary change in the axial dimension after r-mTBI based on increased $K_a$ values at day 6 (Fig. 3U). Taken together, these DTI and DKI findings confirm relatively little corpus callosum damage.
Conversely, and unexpectedly, our diffusion analysis revealed cortical abnormalities in this r-mTBI model.

To test the specificity of these findings in r-mTBI, we performed several additional analyses. Further r-mTBI analysis of diffusion DEC maps (Fig. 4A,B) demonstrated high anisotropy in the corpus callosum as well as marked directionality in the medial region of the cerebral cortex, which supports effective DTI and DKI analyses in these cortical areas. Structural analysis after r-mTBI using T2-weighted images and calculated T2 maps did not detect any overt structural changes or evidence of edema or microhemorrhages in the cortex under the impact site for any time point or condition (Fig. 4C,D and data not shown). The overlying scalp shows T2 hyperintensity attributed to the impacts onto the skin in the r-mTBI mice, which are not observed in r-sham mice (Fig. 4C,D).

Post-imaging analysis of repetitive mild traumatic brain injury pathology in corpus callosum and cortex

After completing the final MRI scan at the 6-week time point, mice were perfused for post-imaging analysis of the underlying pathology in the corpus callosum and cortical ROIs from the DTI and DKI analysis. We first examined neuroinflammation using CD11b and GFAP for detection of microglia and astrocytes, respectively. The r-mTBI tissues from our longitudinal MRI analysis showed mild, but persistent, neuroinflammation in the corpus callosum (Fig. 5A,B). Specifically, in r-mTBI mice compared to r-sham mice, the percent area immunolabeled was increased for CD11b (Fig. 5C). Morphological classification of individual CD11b-labeled cells also showed that the number of activated microglia increased after r-mTBI (Fig. 5D). Amoeboid CD11b-labeled cells were not found in any condition or time point (data not shown). After r-mTBI, astrogliosis in the corpus callosum was not significantly different from the r-sham condition (Fig. 5E). Overall, in the r-mTBI mice, neuroinflammation in the corpus callosum was much less robust relative to the s-mTBI model, which was used as a positive control comparison for corpus callosum pathology detectable by DTI (Supplementary Figs. 2 and 3) (see online supplementary material at http://www.liebertpub.com). Neuroinflammation in medial cortical regions, that is, directly under the impact site, in r-mTBI mice, showed a significant increase in microglial activation, but not astrogliosis (Fig. 5F–H). In contrast, the single-injury model did not exhibit significant neuroinflammation in the cortex with matched analysis of immunoreactivity for either CD11b (Supplementary Fig. 3F,G) (see online supplementary material at http://www.liebertpub.com) or GFAP (Supplementary Fig. 3H) (see online supplementary material at http://www.liebertpub.com) and did not have cortical abnormalities for DTI or DKI analysis (Supplementary Fig. 2) (see online supplementary material at http://www.liebertpub.com).

**Fig. 4.** In vivo diffusion direction encoded color (DEC) maps and T2-weighted imaging of r-mTBI. (A and B) DEC maps show the coronal level that includes the anterior commissure (ac), which aligns with bregma at the site of impact, 3 days after r-sham injury (A) or r-mTBI (B). The corpus callosum (cc) shows a high level of anisotropy. Note that the degree of anisotropy in the medial cortex is greater than in more lateral cortical regions. (C and D) Representative T2-weighted images at the coronal level of impact do not show signs of edema or overt structural damage in either the corpus callosum or cortex after r-sham injury (C) or r-mTBI (D). Images show 6 days post-TBI when significant FA changes are found in the medial cortex after r-mTBI. Scale bar = 1 mm. Colors indicate fiber direction: red: medial-lateral; blue: anterior-posterior; and green: superior-inferior. d, days; r-mTBI, repetitive mild traumatic brain injury; r-sham, repetitive sham.
We next examined changes related to myelination (Fig. 6). Immunohistochemistry for MOG myelin protein correlates well with electron microscopy and Luxol fast blue myelin stain to estimate demyelination within the corpus callosum. After r-mTBI, MOG immunolabeling was examined both in the corpus callosum of r-mTBI mice relative to sham (C and D). Atrophy in the corpus callosum was not significantly increased after r-mTBI (E). In the cortex, the r-mTBI mice had a significant increase in the area of CD11b immunolabeling and in the number of activated microglia (F and G). Astroglia in the medial cortex was not significantly increased after r-mTBI (H). Dashed lines outline the corpus callosum. Values are mean ± standard error of the mean; n = 5; scale bar = 50 μm; *p < 0.05. DAPI, 4′,6-diamidino-2-phenylindole; GFAP, glial fibrillary acidic protein; r-mTBI, repetitive mild traumatic brain injury; r-sham, repetitive sham.

NG2 cells give rise to new oligodendrocytes for remyelination and are also an important reactive cell population after cortical injury. Our previous studies in the s-mTBI model found that NG2 cell density increased in the corpus callosum and cortex.

FIG. 5. Post-imaging neuroinflammation in the corpus callosum and cortex after r-mTBI. Immunohistochemistry was used to detect microglia/macrophage cells with CD11b and astrocytes with GFAP in mice perfused after the final imaging scan, that is, 6 weeks post-injury or sham procedure. (A and B) Slight astroglia and microglia/macrophage activation were evident in r-mTBI mice (B) compared to r-sham mice (A). (C-H) Quantitative analysis shows a significant increase in microglial activation in the corpus callosum of r-mTBI mice relative to sham (C and D). Astroglia in the corpus callosum was not significantly increased after r-mTBI (E). In the cortex, the r-mTBI mice had a significant increase in the area of CD11b immunolabeling and in the number of activated microglia (F and G). Astroglia in the medial cortex was not significantly increased after r-mTBI (H). Dashed lines outline the corpus callosum. Values are mean ± standard error of the mean; n = 5; scale bar = 50 μm; *p < 0.05. DAPI, 4′,6-diamidino-2-phenylindole; GFAP, glial fibrillary acidic protein; r-mTBI, repetitive mild traumatic brain injury; r-sham, repetitive sham.
during the initial weeks post-injury with less of an elevation by 6 weeks post-injury. In r-mTBI mice, the NG2 cell population was not affected in the corpus callosum or cortex in the post-imaging 6-week cohort (Fig. 6C,D,F,H).

Further analysis of tau pathology was performed in the post-imaging 6-week cohort as a potential feature that may produce cortical changes after repetitive injury (Fig. 7). However, immunoreactivity for phosphorylated tau epitopes was not detected in r-mTBI mice using a pSer396 antibody (Fig. 7A) or AT8 antibody (data not shown). The sham condition also did not exhibit immunoreactivity for phosphorylated tau (data not shown). P301S human tau mice were used as a positive control using the same immunohistochemical protocols and showed clear tau pathology in cortical neurons for both pSer396 (Fig. 7B) and AT8 (data not shown).

Correlation studies were performed to more directly interpret the cortical abnormalities identified in the DTI-DKI analysis...
relative to the post-imaging pathological findings (Table 1; Fig. 8). At this 6-week time point, the significant changes between r-mTBI and r-sham were decreased AD values (Fig. 3R) and increased microglial activation (Fig. 5F). Values for these two parameters were strongly correlated (Table 1), and CD11b variation contributed to over 70% of the variation in AD based on the $r^2$ analysis (Fig. 8D). In contrast, cortical AD values did not correlate well with values for GFAP or MOG immunoreactivity (Table 1; Fig. 8E,F).

Next, we further examined the cortical changes after r-mTBI using Thy1-YFP-16 mice (Fig. 9), which have a high density of neurons with yellow fluorescent labeling. The labeling in cortical neurons enables visualization of the dendritic arbor and axonal processes, including callosal axons.30 Nissl staining in r-mTBI mice at 3 days post-injury and in matched r-sham mice showed a low density of neuronal cell bodies in layer I and a much higher density in layers II/III, indicating generally normal neuronal cytoarchitecture in the superficial cortex under the site of impact compared to the post-imaging pathological findings (Table 1; Fig. 8).

**Social interaction deficit resulting from repetitive mild traumatic brain injury**

Finally, behavioral assessments were targeted to the MRI ROIs to detect potential functional deficits in r-mTBI mice. The corpus callosum overlying the lateral ventricles is associated with motor skill learning that can be assessed using a running wheel with irregularly spaced rungs.45 Over a 3-week period, the r-mTBI mice appeared similar to r-sham mice in mastering the complex running-wheel task (Fig. 10A,B). The medial cortical region under the impact site used in this r-mTBI model includes the anterior cingulate cortex, which is considered a hub of social behaviors.51 A three-chamber social apparatus was used to compare the time each mouse spent interacting with an empty carrier or with a stranger mouse inside the carrier (Fig. 10C). The r-sham mice showed the normal preference to spend time interacting with the stranger mouse (Fig. 10C). The interaction time decreased from 190.3 ± 13.8 sec in r-sham mice to 127.9 ± 13.5 sec in r-mTBI mice (Fig. 10C), indicating impaired social approach behavior after r-mTBI. A cohort of mice subjected to the s-mTBI model that produces more-extensive damage in the corpus callosum than the cortex (Supplementary Figs. 2 and 3) (see online supplementary material at http://www.liebertpub.com) did not result in behavioral changes in either wheel running or social interaction (Supplementary Fig. 6) (see online supplementary material at http://www.liebertpub.com) and serves as a negative control comparison. Overall, the r-mTBI impairment on social behavior supports the interpretation that r-mTBI induces significant cortical changes.

**Discussion**

Our findings demonstrate the utility of longitudinal DTI combined with DKI for noninvasive analysis of white matter...
injury as well as more-subtle cortical abnormalities resulting from repetitive concussive impacts. These neuroimaging changes were supported by post-imaging pathological analysis of the corpus callosum and cerebral cortex at 6 weeks post-injury. The r-mTBI model had minimal DTI/DKI changes in the corpus callosum. Yet, the r-mTBI mice showed significant changes in the cortex for axial parameters with both DTI (AD) and DKI (Ka). Cortical changes in AD after r-mTBI correlated strongly with a significant microglial reaction on post-imaging histopathology. These cortical differences were supported by corresponding deficits in social behavior, which involves cortical regions under the impact site.

As a supplemental analysis, using a previously characterized single-impact model, we performed neuroimaging out to a sub-acute time point of 6 weeks with both DTI and DKI analysis followed by post-imaging tissue analysis. This single-impact model served as a positive control for corpus callosum pathology (Supplementary Fig. 3) (see online supplementary material at http://www.liebertpub.com) detectable by DTI (Supplementary Fig. 2) (see online supplementary material at http://www.liebertpub.com), to more accurately interpret the mainly negative findings for DTI and DKI in the corpus callosum of r-mTBI mice (Fig. 3).

Our radiological findings in the s-mTBI model are relatively consistent with recent studies of single-impact injuries, although...
differences in pathology must be taken into consideration. Closed head impact in rats resulted in reduced FA in the corpus callosum, which corresponded with axonal injury. CCI in rats produces a cortical contusion lesion along with pathology in a widespread distribution of white matter tracts and corresponds with bidirectional changes in FA. In this report, FA was reduced in the corpus callosum, but was increased in subcortical white matter as well as in some white matter tracts. Therefore, reduced FA was predicted to reflect fiber disorganization versus reorganization and/or higher fiber density in tracts with increased FA.

The novel r-mTBI model was initially developed in the current study based on analysis of the rostral corpus callosum (Fig. 1) and then expanded to overlying cortical regions to explore potential pathology after repetitive impacts. The 1-day interval choice was supported from our recent findings that glucose uptake is decreased at 24 h post-TBI and a second injury at 24 h worsens outcome. This r-mTBI model is anterior, but otherwise similar, to several other more posterior repetitive mTBI models, suggesting that paradigms of at least three to five mild impacts with 1- to 3-day intervals are effective and practical. Our expectation was that the impact used in the r-mTBI model would result in relatively mild initial damage in the corpus callosum whereas repetition may promote progression toward chronic white matter pathology, including neuroinflammation.

Indeed, the r-mTBI model showed no overt structural damage based on Prussian blue (Fig. 2) and Nissl staining (Fig. 9) or T2 MRI (Fig. 4). β-APP analysis at 24 h showed that the r-mTBI (Fig. 1L) resulted in less than half the amount of initial axon damage in the corpus callosum than was observed after s-mTBI in our previous study. A progression toward chronic white matter pathology was indicated by a significant reduction of FA that was only observed in the corpus callosum at 6 weeks post-injury in

FIG. 9. Thy1-YFP-16 mice show axonal pathology in the cortex and corpus callosum after r-mTBI. (A and B) Thy1-YFP (green) shows labeling of neurons throughout the cortex along with axons projecting through the corpus callosum in mice perfused 3 days after r-sham (A) or r-mTBI (B) procedure. Nissl labeling (red) of the cytoplasmic rough endoplasmic reticulum along with DAPI nuclei (blue) shows the overall cortical cytoarchitecture, which is not markedly disrupted after r-mTBI. Arrows (B) point to examples of swollen segments of damaged axons. (C and D) At higher magnification, Thy1-YFP (green) labeled axons in the corpus callosum exhibit a generally uniform longitudinal profile in r-sham mice (C) in contrast to the numerous swollen axonal segments observed in r-mTBI mice (D). (E and F) High-magnification Thy1-YFP (green) cortical regions analyzed in r-sham mice (E) and r-mTBI mice (F). Large swellings and thickened Thy1-YFP axonal profiles are evident in the cortex of r-mTBI mice (F, arrows) compared to the thin processes of Thy1-YFP axons in r-sham mice (E, arrow). Lack of DAPI nuclear labeling (blue) shows that enlarged Thy1-YFP regions (F) are not cell bodies. (G) Quantification of damaged axonal profiles (swollen, axonal bulbs, and varicosities) labeled with Thy1-YFP in the cingulate and M2 motor cortices that comprise the region of interest in the medial cortex for DTI/DKI analyses. A significant increase of damaged axons was observed in r-mTBI mice compared to r-sham. Values are mean ± standard error of the mean; n = 3; three to nine sections per animal. **p < 0.01; ****p < 0.0001. Scale bars = 200 μm (A and B) and 20 μm (C–F). Avg, average; d, days; DAPI, 4′,6-diamidino-2-phenylindole; DKI, diffusion kurtosis imaging; DTI, diffusion tensor imaging; r-mTBI, repetitive mild traumatic brain injury; r-sham, repetitive sham.
REPETITIVE MTBI PRODUCES CORTICAL ABNORMALITIES

The r-mTBI model produced significant radiological (Fig. 3) and pathological differences (Fig. 5) in cerebral cortex regions overlapping the corpus callosum at coronal levels near the impact site. Further, r-mTBI injury resulted in social behavior deficits (Fig. 10) associated with neural circuits of this medial cortical region, which includes anterior cingulate cortex. The r-mTBI mice had slight, but significant, differences in cortical FA values that were driven by reduced AD (Fig. 3). DKI detects changes in the non-Gaussian component of diffusion at higher b-values and is expected to reflect changes in diffusional heterogeneity, which may correspond to microstructural complexity.26,59 DKI showed a corresponding change in $K_a$, which is consistent with effects in the axial direction (Fig. 3). These cortical findings are somewhat unusual in that gray matter generally has low anisotropy so that DTI analysis is not expected to be informative. However, some cortical regions have more structural directionality, such as the medial cortical region analyzed in this study (Fig. 4).

Although the r-mTBI changes in AD and $K_a$ may be relatively small, they may be particularly informative in cortical regions with sufficient anisotropy. In white matter, FA appears most informative because of the combination of pathological factors that can decrease axial values, including axon damage, and increase radial values, including demyelination.60,61 After r-mTBI, in cortical regions, demyelination was not observed (Fig. 6) and axon damage was similar to s-mTBI (Figs. 9 and Supplementary Fig. 5) (see online supplementary material at http://www.liebertpub.com).

Yet, cortical regions showed significantly reduced AD and increased $K_a$, which typically corresponds to axon damage. Neuroinflammation has also been shown to influence DTI and DKI parameters in TBI models. For example, several reports have correlated astrogliosis with increased FA, driven by increased AD, or with kurtosis.18,26,52 In the current study, the decreased AD values in this medial cortical region correlated strongly with a significant increase in microglial reactivity on post-imaging tissue analysis (Table 1; Fig. 8A,D). It should be noted that diffusion studies have often evaluated FA values without reporting AD or RD changes,62 and this lack of specificity complicates the physiological interpretation of these diffusion changes. Even with clear changes in FA, AD, and RD throughout a time course of white matter disease, the specificity of each parameter is impaired when pathological features are present in combination.63 Previous animal and clinical studies have not necessarily agreed on FA, RD, or AD changes in gray matter post-mTBI.27,63,64 Further studies are needed to more completely examine the full extent of cortical pathology across the post-injury time course after repetitive injury.

The pathological and radiological abnormalities observed in the anterior cingulate and motor cortices after r-mTBI could be a precursor to cortical thinning that has been associated with concussion. In collegiate football players with a history of concussion, MRI revealed significant cortical thinning in frontotemporal regions, including the rostral anterior cingulate cortex and motor cortex, without changes in white matter FA.65 The anterior cingulate region was also identified as a region of cortical atrophy in a 1-year longitudinal study of mTBI patients with a single episode of concussion.66 Cortical abnormalities after repetitive concussive...
injury have also been associated with hyperphosphorylated tau accumulations in neurons and astrocytes, particularly at the depths of sulci and around small blood vessels.\(^6\)\(^7\)\(^8\) In our analysis of r-mTBI mice, hyperphosphorylated tau pathology was not evident using either AT8 staining or an antibody against phosphorylated serine 396 of tau (Fig. 7). Both antibodies readily detected tau pathology in transgenic P301S mice expressing mutant human tau, which served as a positive control. Therefore, tau pathology is unlikely to have contributed to the cortical abnormalities in AD and K\(_\alpha\) in our r-mTBI model. However, longer-term studies would be needed to fully appreciate the potential similarities of our r-mTBI findings to chronic-stage changes in TBI patients based on either cortical thinning or neurodegenerative disease.

To further understand the effects of the r-mTBI injury, functional assessments were targeted to the corpus callosum and cortical regions examined in our DTI/DKI analysis and post-imaging histopathology. The rostral area of corpus callosum is involved in motor skill learning on the complex running wheels.\(^4\)\(^5\) Axonal pathology in the corpus callosum or M2 motor cortex, which was evident at 3 days post-injury in Thy1-YFP-16 mice, did not impair motor skill learning (Fig. 10A,B). The r-mTBI mice had slower maximum velocities during the learning period of days 1–3 after introduction to the complex wheels, but the injury did not result in a statistical difference from sham. By day 4 on the complex wheels, the cohorts reached a plateau for the maximum velocity run each day, which was similar between injured and sham mice through 2 weeks post-injury. This plateau velocity has been shown to significantly decrease in conditions with extensive demyelination of the corpus callosum.\(^4\)\(^1\)\(^2\) The single-impact model with more-extensive corpus callosum pathology also did not alter the running on the complex wheel (Supplementary Fig. 6A) (see online supplementary material at http://www.liebertpub.com). The lack of impairment on both phases of this complex wheel assay indicates that either the current injuries do not impair function of a sufficient proportion of axons in each region or that broader pathology is needed to detect functional deficits with these assessment protocols.

The medial cortex region in our analysis includes the anterior cingulate cortex, which is involved in multiple cognitive domains and is considered a cortical hub for socially driven interactions in humans.\(^3\)\(^1\) In mice, deficits in social approach behavior are related to changes in intracellular signaling, cortical cytoarchitecture, and dendritic spine formation in the anterior cingulate cortex.\(^6\) The three-chambered task has been widely used to evaluate social interactions in mice.\(^3\)\(^9\)\(^7\)\(^1\) In TBI models, decreased social interaction has been reported in conditions of mild injury when combined with stress.\(^4\)\(^0\)\(^7\)\(^2\)\(^3\)\(^4\) We used this assessment of social approach behavior and found deficits after r-mTBI. Therefore, repetitive injury can impair social behavior without a stress paradigm. Social approach behavior was tested at 3 weeks post-injury or sham procedures as a time point that was midway between the 3-day and 6-week time points analyzed by MRI. This behavioral deficit at 3 weeks after r-mTBI indicates a potential predictive value of the 3- and 6-day abnormalities in DTI and DKI. These findings could be used to now develop an experimental design that would formally test predictive value in future studies.

Our findings highlight the need to evaluate changes in both white matter tracts and cortical regions post-mTBI. Using longitudinal MRI DTI combined with DKI, FA was most sensitive to varying severity of white matter injury (Fig. 3 and Supplementary Fig. 2) (see online supplementary material at http://www.liebertpub.com). Axial parameters, AD and K\(_\alpha\), were more informative for detecting subtle cortical abnormalities resulting from repetitive mild injury (Fig. 3). Cortical changes in AD after repetitive injury were strongly correlated with an indicator of a microglial reaction on post-imaging tissue analysis (Table 1; Fig. 8). The r-mTBI model used in the current study exhibits more pathology, based on \(\beta\)APP analysis (Fig. 1), than a report of DTI changes in the cortex only in MD after repetitive mTBI.\(^7\)\(^1\) Our studies are consistent with recent work in fluorescent reporter mice that characterized the microglia reaction as sensitive to cortical damage in mTBI.\(^7\)\(^3\) A recent study of acute repetitive closed head TBI in mice also reported changes in RNA expression indicating activation of microglia in the cortex along with changes in ex vivo DTI AD.\(^7\)\(^5\) Therefore, repetitive concussive or sub-concussive impacts may lead to a microglial reaction in the cortex that could be detected noninvasively with continued advances in the clinical application of diffusion imaging approaches. This cortical microglial reaction could have significant functional implications for behavioral deficits and cognitive impairment given the emerging roles of microglia in synaptic pruning and neurodegenerative disease.\(^7\)\(^6\)–\(^7\)\(^8\)

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