Microstructural Investigations of the Visual Pathways in Pediatric Epilepsy Neurosurgery: Insights From Multi-Shell Diffusion Magnetic Resonance Imaging

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Background: Surgery is a key approach for achieving seizure freedom in children with focal onset epilepsy. However, the resection can affect or be in the vicinity of the optic radiations. Multi-shell diffusion MRI and tractography can better characterize tissue structure and provide guidance to help minimize surgical related deficits. Whilst in adults tractography has been used to demonstrate that damage to the optic radiations leads to postoperative visual field deficits, this approach has yet to be properly explored in children.

Objective: To demonstrate the capabilities of multi-shell diffusion MRI and tractography in characterizing microstructural changes in children with epilepsy pre- and post-surgery affecting the occipital, parietal or temporal lobes.

Methods: Diffusion Tensor Imaging and the Spherical Mean Technique were used to investigate the microstructure of the optic radiations. Furthermore, tractography was used to evaluate whether pre-surgical reconstructions of the optic radiations overlap with the resection margin as measured using anatomical post-surgical T1-weighted MRI.

Results: Increased diffusivity in patients compared to controls at baseline was observed with evidence of decreased diffusivity, anisotropy, and neurite orientation distribution in contralateral hemisphere after surgery. Pre-surgical optic radiation tractography overlapped with post-surgical resection margins in 20/43 (46%) children, and where visual data was available before and after surgery, the presence of overlap indicated a visual field deficit.
**INTRODUCTION**

Resective surgery remains one of the most effective approaches to cure seizures in patients with drug resistant focal epilepsy (Ryvlin et al., 2014). In adults, the most common epilepsy surgery performed is anterior temporal lobe resection (ATLR), where the temporal pole, amygdala and anterior part of the hippocampus are removed (Kwan et al., 2011). Such surgery may damage the optic radiation (Winston et al., 2012). The optic radiations depart from the lateral geniculate nuclei (LGN) and subdivide into three main bundles: a posterior bundle and central bundle that run straight back to the occipital lobe and an anterior bundle, that runs anteriorly alongside the lateral ventricle, projecting deep within the temporal lobe before turning backward at the level of the temporal horn to join the first two bundles (Ebeling and Reulen, 1988; Rubino et al., 2005). The anterior bundle, most often denominated Meyer's Loop, is most at risk in the types of resections described above, and the correlation between the extent of its resection and resulting visual field deficits following surgery in adults is well established (Falconer and Wilson, 1958). Diffusion MRI allows extraction of measures that identify the main directions of water molecular displacement and from that a three-dimensional trajectory of fiber bundles can be reconstructed (tractography (Conturo et al., 1999) and has been used in numerous studies to reconstruct the optic radiations (Yogarajah et al., 2009; Clatworthy et al., 2010; Martinez-Heras et al., 2015; Chamberland et al., 2017)]. Despite the success of tractography in demonstrating involvement of the optic radiations in the resected tissue and subsequent correlation between the extent of optic radiation involvement and post-surgical visual fields in adults (Yam et al., 2010; Winston et al., 2011, 2012, 2014a), there is very little literature on the use of the technique in pediatric series. Addressing this issue is clearly an important step to determine the relevance of optic radiation tractography as a potential neurosurgical planning tool in children undergoing surgery for intractable epilepsy, where there is a greater variety of surgical approaches reflecting the broader and more diverse pathological and neuro-anatomical involvement. As such, surgical damage to the visual system may be more varied than in adults resulting in a wider variety of visual function disturbance than visual field deficits alone (Lee et al., 2002). Optic radiation tractography in healthy children has been well-documented in our previous work, including anatomical distances that are relevant to epilepsy surgery (Dayan et al., 2015b). Furthermore, based on studies in healthy adults, we have demonstrated optic radiation tractography to be robust and reproducible (Dayan et al., 2015a,b). Diffusion tensor imaging (DTI) (Basser et al., 1994) has been applied successfully in clinical practice but associated metrics such as fractional anisotropy are sensitive to numerous tissue properties including axonal density, diameter and orientation (Alexander et al., 2007; Jones et al., 2013). Building on the knowledge provided by DTI, more advanced diffusion signal modeling methods have been developed in recent years to explore the microscopic environment in clinically feasible scan times (Zhang et al., 2012; Farquharson et al., 2013; Lasić et al., 2014; Kaden et al., 2016a; Salama et al., 2018). This has been made possible due to developments in acquisition technology, such as multi-band echo planar imaging (Setsompop et al., 2012a,b) and multi-shell diffusion MRI in which a second shell or b-value of diffusion sensitizing directions is added to the initial single shell required for DTI. Here we focus on the spherical means technique (SMT), which is based on the observation that for any fixed b-value, the spherical mean of the diffusion signal over the gradient directions does not depend on the orientational structure (Kaden et al., 2016b). This insight permits disentangling microstructural tissue features from orientational effects such as fiber crossings and orientation dispersion that confound traditional DTI metrics.

**MAIN OBJECTIVES**

The aim of the present study was to utilize multi-shell diffusion MRI and tractography to delineate the posterior part of the visual pathway in children who have undergone epilepsy surgery involving the occipital, parietal and temporal lobes to:

1. Determine the frequency and location of optic radiation involvement relative to the resection and to relate these to information regarding measures of visual function where clinically available.
2. Investigate pre-existing differences in tissue microstructure between healthy subjects and the patient cohort before surgery and determine differences between the hemisphere where the operation would take place (“ipsilateral”), in comparison with the healthy hemisphere (“contralateral”).
3. Investigate the effects of surgery on the contralateral hemisphere by evaluating differences between the pre- and post-surgical optic radiation microstructure.

**MATERIALS AND METHODS**

**Ethics and Patient Information**

This manuscript results from a retrospective study registered as a Case Note Review, with access to previously collected, non-identifiable information/data under GAfREC 2011 and as a result is exempt from REC approval and from patient consent.
An approval was issued by the relevant Trust for the study to proceed. All patients were medically refractory to anti-epileptic drugs as per the ILAE definition (Téllez-Zenteno et al., 2014). Patients that had undergone resective epilepsy surgery affecting the temporal, parietal and occipital lobes and that may therefore have involved the optic radiations were selected for this study (age at operation 5–19 years). Patients undergoing hemispherectomy were excluded. In total 43 patients with median age of 10.70 years (Inter-quartile range 8.4; 22 male) was also selected to evaluate possible differences in optic radiation microstructure when compared to the patient group; further details are provided in the Table 1. Of the entire cohort there were 12 patients that had complete sets of data. Additionally, imaging data from a cohort of 50 healthy children and young adults with no visual or neurological conditions and median age of 11 years (Inter-quartile range 6.5; 30 male) was also selected to evaluate possible differences in optic projection microstructure when compared to the patient group; further details are provided in the main objectives section. Patient and control groups are both age (F = 1.31, df = 91, p-value = 0.2556) and gender (χ² = 0.73243, df = 1, p-value = 0.3921) matched.

**Imaging Data**

All imaging data were acquired on a Siemens 3.0 T Prisma scanner (Siemens, Erlangen, Germany), equipped with a 20-channel head receive coil. The protocol included a multi-shell diffusion sequence employing a diffusion-weighted single shot 2D EPI acquisition, with multi-band radio frequency pulses to accelerate volume coverage along the slice direction (Setsompop et al., 2012a,b; Auerbach et al., 2013). A multi-band factor of 2 was used to image 66 slices of 2 mm thickness with 0.2 mm slice gap. Diffusion gradients were applied over two ‘shells’: \( b = 1000 \text{ s/mm}^2 \) and \( b = 2200 \text{ s/mm}^2 \), with 60 non-collinear diffusion directions per shell in addition to 13 interleaved \( b = 0 \) (non-diffusion weighted) images. Other imaging parameters were: TR = 3050 ms, TE = 60 ms, field of view = 220 mm × 220 mm, matrix size = 110 × 110, in-plane voxel resolution = 2.0 mm × 2.0 mm, GRAPPA factor 2, phase-encoding (PE) partial Fourier = 6/8. An additional \( b = 0 \) acquisition was performed, with identical readout to the diffusion-weighted scan, but with the phase encode direction flipped by 180° (in the anterior-posterior direction), for correction of susceptibility-related artifacts. The total scan time for the multi-shell diffusion sequence (including the \( b = 0 \) acquisition with flipped PE) was 7 min 50 s. In addition, a T1-weighted MPRAGE structural image was acquired using 176 contiguous sagittal slices, FOV = 256 mm × 240 mm and 1 × 1 × 1 mm image resolution; TE = 4.9 ms and TR = 11 ms.

**Pre-processing and Reconstruction of Maps**

Multi-shell diffusion MRI data were denoised using MRtrix3’s implementation of the method developed by Veraart et al. (2016a,b). Furthermore, FSL (Jenkinson et al., 2012) commands TOPUP and EDDY were used to correct for susceptibility distortions and to perform motion and eddy current correction (Andersson and Sotiropoulos, 2016). After pre-processing, multi-shell diffusion MRI data was used in the MRtrix3 Constrained Spherical Deconvolution (CSD) pipeline, estimating both the multi-tissue response function and fiber orientation distribution (FOD) fields (Tournier et al., 2012). Freesurfer was used for bias field correction and skull stripping of pre-surgical structural T1-weighted scans and to provide parcellation of cortical and subcortical structures based on a standard atlas (Fischl, 2012). As Freesurfer was developed mainly for adult brains, in this study, regions from the derived cortical and subcortical parcellation, namely the thalamus and occipital cortices, were visually inspected and edited when required for each subject, before being used as regions of interest for performing optic radiation tractography. DTI maps were reconstructed with MRtrix3 – fractional anisotropy (FA), mean diffusivity (MD), axial diffusivity (AD), and radial diffusivity (RD). The SMT microscopic tensor was also fit and provided estimates of the microscopic diffusivities parallel (Long) and perpendicular to the neurites (Trans) from which the Microscopic fractional anisotropy (\( \mu \text{FA} \)) and Microscopic mean diffusivity (\( \mu \text{MD} \)) were calculated (Kaden et al., 2016b). Further to the measurement of \( \mu \text{FA} \) which reflects the degree of directionality arising from the local diffusion processes, the SMT technique also enables the reconstruction of the Orientation Dispersion Entropy (ODEntropy) reflecting the microdomain orientation distribution, taking higher values for regions constituted by highly coherent bundles and a lower value for more uniformly distributed substrates such as gray matter or the ventricular system.

**Tractography Reconstructions**

The Freesurfer parcelations were registered to diffusion space by registering the structural skull stripped data to the mean \( b = 0 \) image calculated for each subject, following a rigid body transformation and non-linear deformations applied with ANTS (Avants et al., 2014). Seeding and inclusion regions of interest, comprising the thalamus and a composite occipital region – consisting of lingual, pericalcarine, lateral occipital cortical regions – were generated for each subject. In order to determine the starting point of the optic radiation tractography – i.e., the LGN – the center of mass of the thalamus region was calculated and used to divide it in two halves; the posterior half of the thalamic region was then retained as the seed region. Additional regions of interest to exclude artifactual contributions were extracted from the Freesurfer parcelation, namely the contralateral gray matter mask. Finally sagittal, axial and coronal ROIs were drawn for each individual subject in
### TABLE 1 | Detailed clinical information for selected patient population.

| Patient | Age at surgery (years) | Scan after surgery (days) | Visual assessment after surgery (days) | Gender | Surgery | Seizure freedom > 1 year | Medication |
|---------|------------------------|---------------------------|--------------------------------------|--------|---------|--------------------------|------------|
| Patient01 | 16.0                  | 179                       | No data                              | F      | L temporal lesionectomy | Yes         | Keppra                 |
| Patient02 | 7.0                   | 137                       | No data                              | M      | L anterior temporal lobectomy | No          | Phenytoin, sodium valproate and zonisamide |
| Patient03 | 7.0                   | 236                       | No data                              | M      | L anterior temporal lobectomy and amygdalectomy | Yes         | Sodium valproate, lamotrigine |
| Patient04 | 13.0                  | 100 267                   | F                                    | L anterior temporal lobectomy and amygdalectomy | Yes         | Oxcarbazepine, zonisamide |
| Patient05 | 15.0                  | 167                       | No data                              | F      | L anterior temporal lobectomy and amygdalectomy | Yes         | Carbamazepine, sodium valproate, clobazam |
| Patient06 | 6.0                   | 559                       | No data                              | F      | L frontal, anterior temporal lobectomy and amygdalectomy | No          | Lacosamide, phenobarbitalne |
| Patient07 | 16.0                  | 200                       | No data                              | M      | L posterior temporal lesionectomy | Yes         | Oxcarbazepine, sodium valproate, levetiracetam |
| Patient08 | 12.0                  | 168                       | No data                              | M      | L posterior temporal lesionectomy | Yes         | Carbamazepine, clonazepam |
| Patient09 | 17.0                  | 82 140                    | F                                    | L temporal lesionectomy | Yes         | Tegretol                 |
| Patient10 | 6.0                   | 117                       | No data                              | M      | L temporal lesionectomy | Yes         | Levetiracetam, clobazam |
| Patient11 | 7.0                   | 286                       | No data                              | F      | L anterior temporal lesionectomy and amygdalectomy | No          | Topiramate, clobazam |
| Patient12 | 18.0                  | 102 320                   | M                                    | L anterior temporal lesionectomy (extended) | No          | Lamotrigine, lacosamide |
| Patient13 | 11.0                  | 4 50                      | M                                    | L anterior temporal lesionectomy and amygdalectomy (extended) | Yes         | Carbamazepine, sodium valproate, levetiracetam |
| Patient14 | 12.0                  | 158 161                   | M                                    | L temporo-occipito-parietal disconnection | Yes         | Perampanel, sodium valproate, levetiracetam |
| Patient15 | 12.0                  | 89 98                     | M                                    | L temporo-occipito-parietal disconnection | Yes         | Carbamazepine, topiramate |
| Patient16 | 11.0                  | 165                       | No data                              | M      | L temporo-occipito-parietal disconnection | Yes         | Carbamazepine, clobazam |
| Patient17 | 5.0                   | 137 54                    | M                                    | R fronto-temporo-parietal resection | Yes         | Levetiracetam, clobazam |
| Patient18 | 14.0                  | 81 64                     | F                                    | R occipital lesionectomy | Yes         | Sodium valproate, levetiracetam |
| Patient19 | 7.0                   | 186                       | No data                              | F      | R occipital lesionectomy | No          | Sodium valproate, levetiracetam |
| Patient20 | 10.0                  | 308                       | No data                              | F      | R anterior temporal lobectomy and amygdalectomy | Yes         | Sodium valproate, vigabatrin |
| Patient21 | 19.0                  | 237                       | No data                              | F      | R anterior temporal lobectomy and amygdalectomy | Yes         | Oxcarbazepine |
| Patient22 | 12.0                  | 411 264                   | M                                    | R anterior temporal lobectomy and amygdalectomy | No          | Oxcarbazepine, clobazam |
| Patient23 | 17.0                  | 176                       | No data                              | M      | R anterior temporal lobectomy and amygdalectomy | Yes         | Levetiracetam |
| Patient24 | 7.0                   | 160                       | No data                              | F      | R anterior temporal lobectomy and amygdalectomy | No          | Sodium valproate, lamotrigine, midazolam |
| Patient25 | 13.0                  | 108 286                   | M                                    | R temporal lesionectomy | Yes         | –                      |
| Patient26 | 10.0                  | 188                       | No data                              | F      | R temporal lesionectomy | Yes         | Levetiracetam |
| Patient27 | 6.0                   | 287                       | No data                              | F      | L temporal lesionectomy | Yes         | Sodium valproate, lamotrigine |
| Patient28 | 11.0                  | 242                       | No data                              | M      | L parietal tumor resection | Yes         | Zonisamide |
### TABLE 1 | Continued

| Patient | Age at surgery (years) | Scan after surgery (days) | Visual assessment after surgery (days) | Gender | Surgery | Seizure freedom > 1 year | Medication |
|---------|------------------------|--------------------------|---------------------------------------|--------|---------|--------------------------|------------|
| Patient29 | 6.0 | 259 | No data | F | R anterior temporal lesionectomy and amygdalo-hippocampectomy | Yes | Sodium valproate, topiramate |
| Patient30 | 6.0 | 154 | No data | M | L temporal lobectomy | No | Topiramate, keppra |
| Patient31 | 5.0 | 176 | 111 | F | R temporal lesionectomy | Yes | Keppra |
| Patient32 | 9.0 | No data | 333 | M | L temporal lobectomy lesionectomy | No | Lacosamide, levetiracetam, lamotrigine, risperidone, phenytoin |
| Patient33 | 10.0 | 85 | No data | F | L parietal lesionectomy | No | Sodium valproate, levetiracetam |
| Patient34 | 6.0 | No data | No data | F | R occipital lobectomy | Yes | Keppra, lamotrigine |
| Patient35 | 13.0 | 89 | No data | F | L temporal lobectomy and amygdalo-hippocampectomy | Yes | Carbamazepine |
| Patient36 | 8.0 | No data | No data | M | R temporal lobectomy | No | Oxcarbazepine, lacosamide |
| Patient37 | 8.0 | 81 | No data | M | L parietal lesionectomy | Available | Sodium valproate, oxcarbazepine |
| Patient38 | 17.0 | 73 | No data | F | R temporal lesionectomy and amygdalo-hippocampectomy | Available | Keppra, oxcarbazepine, clobazam |
| Patient39 | 16.0 | 155 | 134 | M | L temporo-occipito-parietal disconnection | Available | Carbamazepine, lacosamide, phenobarbitone |
| Patient40 | 18.0 | No data | No data | F | L temporal lesionectomy | Unavailable | Oxcarbazepine, midazolam |
| Patient41 | 18.0 | No data | No data | F | L parietal lesionectomy | Unavailable | Topiramate, lamotrigine, levothyroxine |
| Patient42 | 5.0 | 125 | 239 | M | R temporo-occipito-parietal disconnection | Unavailable | Carbamazepine, lacosamide, levetiracetam, lamotrigine |
| Patient43 | 15.0 | 102 | 299 | F | L temporo-occipito-parietal disconnection | Available | Lacosamide, levetiracetam, lamotrigine |

R, right; L, left; F, female; M, male.

In order to exclude lateral projections from the acoustic radiation, posterior projections of the corpus callosum, parts of the cingulum bundle and brainstem (Dayan et al., 2015a,b; Kammen et al., 2016; Nowell et al., 2016; Chamberland et al., 2017). These regions were used to perform tractography with a probabilistic algorithm in MRTRix3 with 2nd order integration over the fiber orientation distribution [iFOD2 (Tournier et al., 2010)] with a 90° angular threshold, 5000 selected streamlines and FA and FOD amplitude threshold of 0.15 and 0.1, respectively. The tractography were then multiplied by the individual DTI and SMT maps, and a weighted average of each metric calculated, resulting in single value for each metric within the resulting optic radiation region.

#### Resection Segmentation and Pre and Post-surgical MRI Registration

In order to assess overlap between the surgical resection and tractography prior to surgery, the pre-surgical reconstructions were first registered into post-surgical space. The first step consisted of semi-automatic segmentation of the resection, from the post-operative structural scan, using ITK-snap (Avants et al., 2014). The resulting resection mask was then visually inspected by a neurosurgeon (MT) and edited as required, to cover the entire extent of the resection. Following resection segmentation, the registration between pre and post-surgical scans was performed with the ANTS software (Tustison and Avants, 2013), selecting the derived segmentation as an exclusion mask in the registration procedure. The final step involved registering the pre-surgical tractography to post-surgical space. In order to achieve this, the original transformation files (from pre-structural to pre-diffusion space) were applied together with those derived in the previous step. Finally, the main axis along which the optic radiations maps (thresholded at 5%) are oriented was determined so that the optic radiations could be rotated to the axial plane (the same rotation was applied to the post-surgical structural volume and resected mask). The cross-sectional area was then calculated as the fraction of optic radiation voxels in common with the resected area over the whole optic radiation region.
radiation volume, at the coronal slice where overlap between both was maximum (Figure 2).

**Clinical Evaluation of Visual Pathways**
The patient's notes were retrospectively reviewed for pre- and post-operative ophthalmological examination undertaken, and specifically for documented visual field assessment. Visual fields were either tested by Goldmann perimetry or by confrontation methods using the two-examiner technique depending on the child's age and ability at the assessment (Heidary, 2016). As the patients in this study underwent surgery as part of a national service, pre and/or post-surgery ophthalmology has on occasion been performed by local services and as a result visual function data pre- and post-operatively is not available for the some of subjects in the selected cohort (further discussion in the limitations section). Pre-operatively 31 patients had visual field assessment and 2 were abandoned due to poor co-operation, 26 were normal and 3 were abnormal. Post-operatively 15 had visual field assessment out of which 2 were abandoned due to poor co-operation, 5 were normal and 8 were abnormal. Of the 12 patients who had pre- and post-operative testing, 5 were normal throughout, 1 had a deficit pre and post-operatively, the remaining 6 had acquired visual field deficits. Although the ophthalmological data in this report is limited, it was sufficient to establish the degree of visual field deficit (normal, quadrantanopia, or hemianopia) using appropriate methods.
Statistical Analyses
After extraction of all metrics a linear mixed-effects model was fit to the data in R (lme4 library in R\(^1\)) with age, gender, group (patient/control) and timepoint (baseline/after surgery) as fixed effects, and a random effect of subject-specific intercepts and hemisphere. Hemisphere was encoded as ipsilateral vs. contralateral in patients, and left vs. right differences in controls, as to account for the variability across group. The package lmerTest was then used to evaluate significant effects found with the fitted model the results of which can be found in Table 2.

RESULTS
Whilst age and gender displayed several significant associations, we were mostly interested in exploring the effect of surgery as well as group and so report on these: Pre-surgically, both DTI and SMT revealed significant increased diffusivity metrics in patients compared to controls, with further significant anisotropy decrease detected by SMT, while there were no differences in the neurite orientation dispersion as measured by ODEntropy. After surgery, we observed a decrease in contralateral hemisphere AD, FA, Long, \(\mu FA\), and ODEntropy while RD and Trans were increased in comparison to the pre-surgical optic radiations (a summary of these changes can be found in Table 3).

In 20/43 children there was overlap between the pre-surgical optic radiation tractography and post-surgical resection margin on T1-weighted MRI as measured by tractography. Where visual field data were available before and after surgery the presence of resection overlap with tractography corresponded to abnormal post-surgical visual function following normal pre-surgical visual function in every instance – patients 4, 13, 15, 18, 39, 42, and 43 (Figure 3). In patients with no pre-existing or post-surgical deficits no corresponding overlap of the optic radiations with the resection margin was found – patients 9, 17, 25, 31, and 32 (Figure 3). The overlap scores for the remaining patients can be found in the Supplementary Table 1.

| TABLE 2 | Mixed effect model results. |
| --- | --- | --- | --- | --- | --- | --- |
| **Effect** | **Estimate** | **Std. Error** | **t-value** | **Effect** | **Estimate** | **Std. Error** | **t-value** |
| MD | Intercept | 7.94E-04 | 8.04E-06 | 98.753*** | \(\mu MD\) | Intercept | 1.07E-03 | 8.53E-06 | 125.31*** |
| | Group | 3.01E-05 | 5.01E-06 | -6.011*** | Group | 3.98E-05 | 5.40E-06 | -7.372*** |
| | Timepoint | -5.71E-07 | 2.46E-06 | -0.232 | Timepoint | -6.29E-06 | 3.32E-06 | 1.986 |
| | Gender | 1.48E-05 | 4.92E-06 | 2.999** | Gender | 1.62E-05 | 5.20E-06 | 3.119** |
| | Age | -5.29E-06 | 6.12E-07 | -8.651*** | Age | -6.04E-06 | 6.47E-07 | -9.335*** |
| AD | Intercept | 1.23E-03 | 9.38E-06 | 130.786*** | Long | Intercept | 2.68E-03 | 1.72E-05 | 155.745*** |
| | Group | 4.13E-05 | 5.93E-06 | -6.965*** | Group | 7.96E-05 | 1.09E-05 | 7.307*** |
| | Timepoint | -3.95E-05 | 4.34E-06 | 9.109*** | Timepoint | -4.12E-05 | 7.70E-06 | 5.343*** |
| | Gender | 2.40E-05 | 5.72E-06 | 4.187*** | Gender | 3.38E-05 | 1.05E-05 | 3.213*** |
| | Age | -4.18E-06 | 7.12E-07 | -5.868*** | Age | -9.39E-06 | 1.31E-06 | 7.182*** |
| RD | Intercept | 5.81E-04 | 9.56E-06 | 60.733*** | Trans | Intercept | 2.61E-04 | 7.53E-06 | 34.687*** |
| | Group | 2.74E-05 | 5.97E-06 | -4.596*** | Group | 1.97E-05 | 4.68E-06 | 4.217*** |
| | Timepoint | 1.65E-05 | 3.15E-06 | 5.227*** | Timepoint | 1.03E-05 | 2.28E-06 | 4.53*** |
| | Gender | 1.21E-05 | 5.84E-06 | 2.067* | Gender | 9.82E-06 | 4.61E-06 | 2.132* |
| | Age | -5.92E-06 | 7.27E-07 | -8.146*** | Age | -4.27E-06 | 5.73E-07 | -7.446*** |
| FA | Intercept | 4.32E-01 | 7.68E-03 | 56.281*** | \(\mu FA\) | Intercept | 8.86E-01 | 3.36E-03 | 263.751*** |
| | Group | -8.37E-03 | 4.82E-03 | 1.737 | Group | -4.86E-03 | 2.10E-03 | 2.319* |
| | Timepoint | -2.80E-02 | 3.16E-03 | -8.885*** | Timepoint | -6.58E-03 | 1.20E-03 | -5.494*** |
| | Gender | -1.34E-03 | 4.69E-03 | -0.286 | Gender | -3.00E-03 | 2.05E-03 | -1.459 |
| | Age | 3.76E-03 | 5.84E-04 | 6.439*** | Age | 1.66E-03 | 2.56E-04 | 6.506*** |

Fixed effects and their significance are presented for the different metrics extracted within the optic radiations according to the following legend: *\(p < 0.05\), **\(p < 0.01\), and ***\(p < 0.001\).
FIGURE 3 | (A) Distribution of the resection margins for the 43 subjects in this study, per hemisphere, overlapped onto a template generated by registering all the individual structural scans. Hot color scale indicates the percentage of the total operations that were performed in a specific location. The maximum of the hot color scale indicates the highest incidence of surgical procedures for both hemispheres. (B) For a smaller subset of patients for which visual function data was available before and after surgery a small sketch of the resection locations and a table with overlap between pre-surgical tractography of the optic radiations and post-surgical structural scan are presented. R, right; L, left; HH, homonymous hemianopia; HSQ, homonymous superior quadrantanopia. *confrontational, +supported with electrophysiological data.

DISCUSSION

Advanced multi-shell diffusion imaging and tractography was used to detect changes in the structure of the visual pathways in a cohort of children undergoing epilepsy surgery involving the occipital, parietal, and temporal lobes. Additionally, tractography was used to determine optic radiation involvement in the resection margin and where follow-up ophthalmological data was available to determine the correspondence between these.

Investigation of Microstructural Properties of Optic Radiations

There were no significant differences in the microstructural properties of the optic radiations at baseline between the contralateral and ipsilateral sides in patients, nor between left and right hemisphere in controls (i.e., random effects were centered around zero – data not shown). Comparisons between patients and controls revealed statistically significant increases in MD, AD and RD which is consistent with previous findings in adults (Winston et al., 2014b; Bonilha et al., 2015). Further, anisotropy decreases in patients compared to controls were also found although only reaching significance as probed with $\mu$FA (from SMT) and not FA. SMT does not only provide metrics of tissue microstructure not confounded by orientational effects, but also estimates the orientation dispersion entropy, a measure of the neurite orientation distribution which was comparable between patients and controls. Remaining SMT parameters revealed significant increases in longitudinal, mean, and transverse microscopic diffusivities. These results are very likely explained by the increase in extracellular fluid that is observed in epilepsy, as well as altered myelination patterns and axonal loss (Concha et al., 2005). When comparing the contralateral hemisphere before and after surgery statistically significant decreases in AD, FA, Long, $\mu$FA, and OD Entropy as well as an increase in RD and transverse microscopic
and the nature of the developed visual field deficit have not
been well explored (Kivelev et al., 2012). Furthermore, whereas
in adults there have been many studies linking the impact of
surgery on brain function, in particular memory (Duncan, 2008),
this has yet to be addressed in pediatric cohorts (Berg, 2015),
where the incidence of neurological complications is higher than
in adults (Hader et al., 2013). One study in particular reported
a higher rate of major visual field deficits in children when
compared to adults (38% vs. 14%) and a lower rate of minor
visual field deficits in children when compared to adults (6.35
vs. 16%). The higher variability in estimates of the rate of visual
field deficits following epilepsy surgery in a pediatric cohort in
comparison to previous adult reports, likely arises due to the
more diverse surgeries performed as opposed to pure anterior
temporal lobectomies that are performed in adults (Hader et al.,
2013; Schmeiser et al., 2017) and due to the ability of the
brain to reorganize and revert to some extent the severity of
visual field deficit initially determined (Yam et al., 2010). In this
study, the optic radiations were found to be involved in the
resection in 20/43 (46%) patients. Where both pre- and post-
surgical ophthalmological data was available, 7/12 (58%) had a
demonstrable visual field deficit, corresponding to the presence
of a cross-sectional area (CSA) overlap of the resection with the
pre-surgical tractography (see Figure 3). Whilst typically the type
of analysis performed in adults when identifying relationships
between visual pathways and post-surgical deficits focus on the
Meyer’s loop – temporal pole distance (Yogarajah et al., 2009;
Mandelstam, 2012; Lilja et al., 2015), in children the variability in
surgery across patients which could affect any part of the optic
radiation demands a different approach, hence here the cross-
sectional overlap between the optic radiation and the resection
volume was determined.

Limitations and Future Directions
Whilst surgery still remains the most effective course of
action to drug resistant epilepsy (Schmidt and Stavem,
2009; Jobst and Cascino, 2015), it is well known that the
brain shifts position during surgery, and this has been
addressed using tractography during neurosurgical procedures
(Nimsky et al., 2005; Winston et al., 2012; Yang et al.,
2017). It has also been documented that there is degeneration
of the visual system over time following damage to post-
chiasmal visual pathways (Millington et al., 2014). In some
cases, the long-time interval between the pre-surgical scan
and surgery may have resulted in changes in the position
of the visual pathways by the time surgery was performed,
as well as in other brain regions reflecting brain plasticity
(Yogarajah et al., 2010). Furthermore, it is also possible, that
following surgery there may have been some morphological
changes over time, namely, the occupation of the resection
cavity by the remaining brain tissue. The timeframe of these
changes is not clear, but if this is the case, it is possible that
brain shift has contributed to an underestimation of
the segmented resection area. In the absence of proper a
validation technique other than post-mortem dissections, the
comparison between pre- and intra-operative tractography
reconstructions as well as the combination with other MRI
modalities and electrophysiological recordings should also

Clinical Correlations With Tractography
Despite the high anatomical variability and marked asymmetry
that is associated with the distance of Meyer’s Loop to the
temporal pole between hemispheres (Bürgel et al., 2006; Jeelani
et al., 2010; Dreessen de Gervai et al., 2014), in this study we
were able to successfully map the optic radiation in all subjects.
In particular, we were able to do so in a pediatric surgical cohort
and show, to the best of our knowledge, the first report that
relates the degree and location of resection extent with optic
radiation involvement in a pediatric surgical series. Although it
is also self-evident that disruption of optic pathways is related
to postoperative visual field deficit, the location of the resection
and the nature of the developed visual field deficit have not
diffusivity were found. Taken together these findings indicate
changes to the microstructural organization of the white matter
with greater dispersion of structures and microdomains being
less anisotropic. It is unclear exactly what processes relate to
these changes although it is possible that they represent a
process of re-organization. It has been documented previously
that as a result of surgery there is some reorganization that
takes place in the brain (Taylor et al., 2018), and that it is
generally associated to increased anisotropy in the contralateral
hemisphere (Yogarajah et al., 2010; Winston et al., 2014b).
However, a recent study (Li et al., 2019) shows that this trend
is more prominent after 6 months following surgery, following
an initial decrease in anisotropy documented at 3 months post-
surgery in the contralateral optic radiation. The pediatric cohort
studied here had varied intervals for follow-up imaging limiting
the conclusions that can be drawn regarding the timescale of
possible re-organization and its effect on diffusion metrics. Future
work should ideally obtain several follow-up scans at fixed time
points for all subjects in order to determine unambiguously the
evolution of diffusion parameters post-surgically.

Table 2

|                  | Patients vs. controls (at baseline) | Pre-surgical contralateral vs. Post-surgical contralateral hemisphere (patients) |
|------------------|------------------------------------|---------------------------------------------------------------------------------|
| DTI              |                                    |                                                                                 |
| MD               | ↑                                  | –                                                                               |
| AD               | ↑                                  | ↓                                                                               |
| RD               | ↑                                  | ↑                                                                               |
| FA               | –                                  | ↓                                                                               |
| SMT              |                                    |                                                                                 |
| µMD              | ↑                                  | –                                                                               |
| Long             | ↑                                  | ↓                                                                               |
| Trans            | ↑                                  | ↑                                                                               |
| µFA              | –                                  | ↓                                                                               |
| ODEntropy        | –                                  | ↓                                                                               |

Trend of non-significant changes (–) can be further investigated in Table 2. DTI, diffusion tensor imaging metrics; FA, fractional anisotropy; MD, mean diffusivity; AD, axial diffusivity; RD, radial diffusivity; SMT metrics, spherical mean technique; Long, parallel microscopic diffusivity; Trans, perpendicular microscopic diffusivity; µFA, microscopic fractional anisotropy; µMD, Microscopic mean diffusivity.
provide further assurance regarding the use of tractography for neurosurgical purposes (Schmitt et al., 2014; Alvarez et al., 2015; Nooij et al., 2015; Nimsky et al., 2016). A further limitation is the absence of a complete set of ophthalmology data in all the patients studied, although in the data available a clear correspondence between optic radiation involvement and visual field defect was demonstrated. Finally, SMT is able to provide an alternative to explore microstructural changes following surgery in epilepsy when compared to traditional DTI parameters. The technique is still relatively new but does provide unambiguous assessments of tissue dispersion and microscopic anisotropy both of which were found to have changed in the post-surgical optic radiations indicating a potential re-organization phenomenon. Further work is required however, to fully determine the time evolution and significance of these observations.

CONCLUSION

In this study tractography reconstructions of the optic radiations were successfully performed in a pediatric cohort undergoing epilepsy surgery and used to evaluate the overlap of pre-surgical reconstructions with the resected area after surgery. In a cohort of 43 children the optic radiations were involved in the resection margin in 46% of cases. Of those with follow-up ophthalmology data all of the 7 patients with a demonstrable overlap of the optic radiations with the resection volume had a visual field deficit whereas the remaining patients with no involvement of the optic radiations in the resection margin had no evidence of a visual field deficit. Microstructural differences of the optic radiations in patients with epilepsy when compared with controls were observed with higher diffusivity metrics and lower anisotropy, with a comparable (neurite) orientation distribution as measured by SMT. Evidence of changes in the contralateral hemisphere following surgery, namely decreases in anisotropy and (neurite) orientation distribution accompanied by increased diffusivity in the radial direction were also found and are possibly related to a process of microstructural re-organization. Finally, this is the first report in a pediatric series that highlights the relevance and importance of multi-shell diffusion MRI and optic radiation tractography for pre-surgical evaluation in children undergoing epilepsy surgery affecting the occipital, parietal or temporal lobes.

DATA AVAILABILITY STATEMENT

The datasets generated for this study are available on request to the corresponding author.

ETHICS STATEMENT

Ethical review and approval was not required for the study on human participants in accordance with the local legislation and institutional requirements. Written informed consent to participate in this study was provided by the participants’ legal guardian/next of kin.

AUTHOR CONTRIBUTIONS

LL, SH, JC, and CC contributed to conception and design of the study. JC performed the statistical analysis. LL wrote the first draft of the manuscript. All authors contributed to manuscript revision, read and approved the submitted version.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fnins.2020.00269/full#supplementary-material

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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

The handling Editor declared a past co-authorship with one of the authors EK.

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