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Bundle-Specific Tractography with incorporated anatomical and orientational priors

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Abstract

Anatomical white matter bundles vary in shape, size, length, and complexity, making diffusion MRI tractography reconstruction of some bundles more difficult than others. As a result, bundles reconstruction often suffers from a poor spatial extent recovery. To fill-up the white matter volume as much and as best as possible, millions of streamlines can be generated and filtering techniques applied to address this issue. However, well-known problems and biases are introduced such as the creation of a large number of false positives and over-representation of easy-to-track parts of bundles and under-representation of hard-to-track.

To address these challenges, we developed a Bundle-Specific Tractography (BST) algorithm. It incorporates anatomical and orientational prior knowledge during the process of streamline tracing to increase reproducibility, sensitivity, specificity and efficiency when reconstructing certain bundles of interest. BST outperforms classical deterministic, probabilistic, and global tractography methods. The increase in anatomically plausible streamlines, with larger spatial coverage, helps to accurately represent the full shape of bundles, which could greatly enhance and robustify tract-based and connectivity-based neuroimaging studies.

Keywords: Diffusion MRI, White Matter, Tractography, Tractometry, Connectivity

1. Introduction

Tractography algorithms are currently challenged by several important open issues and biases that need to be addressed to improve brain connectivity mapping [Jones, 2010; Jbabdi and Johansen-Berg, 2011; Maier-Hein et al., 2017]. In Maier-Hein et al. [2017], it was recently shown that whole-brain tractography can recover most existing true positive bundles but that these are poorly recovered in terms of spatial extent. In the

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7 simulations of 25 human-like synthetic white matter (WM) bundles, an average spatial
8 extent recovery of only 30% is reported [Maier-Hein et al., 2017, Fig. 3]. We propose
9 to term this tractography issue as the “*poor-spatial-extent*” challenge. Techniques that
10 have better spatial extent recovery always generate more invalid streamlines, and thus
11 fall into the usual *sensitivity-specificity curse*. In the current context, the notion of
12 invalid streamlines refers to anatomically improbable streamlines, meaning their path
13 is not reflective of known WM pathways. The quality of WM bundle reconstruction
14 using tractography is influenced by many variables: i) global geometry of the bundle like
15 curvature, thickness, length, ii) local crossing regions it traverses, position of the bundle
16 with respect to other bundles, and iii) extent and shape of cortical endpoints. As a result,
17 some WM bundles are harder-to-track than others, which leads to the poor-spatial-extent
18 challenge.

19 Why are some bundles more difficult to reconstruct than others? Complex local fiber
20 regions, even if well recovered locally in terms of fiber orientation distributions, can lead
21 to invalid tractography streamlines that respect the underlying local diffusion signal but
22 make clear invalid pathways at the global brain level [Maier-Hein et al., 2017]. Despite
23 the fact that almost all WM regions contain crossing fiber populations [Jeurissen et al.,
24 2013], some are well-know pitfalls for tractography methods and could be tackled with
25 better prior information. As an illustration, the *centrum semiovale*, as seen in Figure 1
26 and Figure 2, was and still is a major difficulty for tractography algorithms. Three major
27 WM pathways: i) the pyramidal tract, ii) the mid-body of the corpus callosum (anterior
28 and posterior), and iii) the arching streamlines connecting the frontal to the temporal
29 lobes, are crossing almost orthogonally in the most ventral part. At such an intersection,
30 even with multiple fiber techniques, tractography algorithms most often follow the easiest
31 path available. Bundles can seem to cover a large spatial extent and have an appropriate
32 amount of fanning when millions of streamlines are used. However, the density is far from
33 uniform within a bundle, even if it cannot be perceived visually, as seen in Figure 2. The
34 amount of crossing regions, overall length and curvature of the bundle will influence the
35 probability of a streamlines to reach its expected destination. Moreover, it is common
36 to observe streamlines abruptly changing direction to either switch to a different bundle
37 or follow an erroneous path, but end up in the right location. Crossing can become even
38 harder to distinguish as the main direction of the fiber populations kiss or come into a
39 bottleneck [Maier-Hein et al., 2017, Fig.7].

40 A partially reconstructed WM bundle could lead to erroneous interpretations and

Figure 1: Crossing region of the Corpus Callosum (red), the Arcuate Fasciculus (green) and the Pyramidal Tract (blue) in the centrum semiovale. A) Coronal cross-section, B) coronal 3D view and C) sagittal 3D view.

Figure 2: Reconstruction of pyramidal tract (PT) streamlines illustrating some of the challenges of tractography, seen on panel A, i) 2-way crossing ii) 3-way crossing, and iii) high curvature. In panel B, regions *a* and *b* encounter less difficulties in reconstruction due to only being subject to a 2-way crossing limitation, as illustrated in Panel A (i), leading to a higher proportion of streamlines reaching the cortex in these general regions. Conversely, region *c* in Panel B encounters 2 and 3 way crossings (Panel A (i) and (ii)), whereas, region *d* in Panel B encounters 2-way crossing and high-curvature limitations (Panel A (i) and (iii)), leading to a lower proportion of streamlines successfully reaching the cortex. Streamline coloring represents the local streamlines density (at the voxel-level), segments in red shows regions where a large proportion of streamlines pass through a bottleneck as opposed to segments in blue. Panel C illustrates that the majority of streamlines traverse through the core of the PT.

conclusions. At the tract-based analysis level [Dayan et al., 2016; Cousineau et al., 2017; Yeatman et al., 2012, 2018], the spatial extent of bundles (in cm^3) is a measurement often used to investigate morphological group differences, such as atrophy or asymmetry [Catani et al., 2007; Song et al., 2014; Chenot et al., 2018]. At the connectome-level, the number of streamlines in WM bundles connecting all possible pairs of cortical regions is usually used [Sotiropoulos and Zalesky, 2017]. However, because of a combination of local and global differences in brain geometry and WM configurations, certain subjects may have easier-to-track bundles than other subjects, which results in WM bundles that fill-up in spatial extent before others and thus have artificially larger number of streamlines. Two strategies exist to attempt overcoming these issues: 1) a whole brain tractography with millions of streamlines followed by manual, clustering or advanced filtering techniques, 2) a tailored ROI-based tractography.

Whole-brain tractography followed by filtering, clustering or manual dissection. In the first strategy, millions of streamlines are typically reconstructed using a whole-brain deterministic, probabilistic, or global fiber tracking algorithms to fill-up the whole WM volume and be able to capture a large spatial extent of all bundles [Wassermann et al., 2016; Jeurissen et al., 2017; Moyer et al., 2018; Calamante et al., 2015; Smith et al., 2015b]. With more streamlines, a larger extent of bundles is reconstructed (higher sensitivity) but also more false positive bundles created (lower specificity) [Côté et al., 2013;

Maier-Hein et al., 2017], thus leading to the *sensitivity-specificity curve* mentioned before. Generation of millions streamlines is often considered a brute force approach with a heavy computational burden. All *iterations* leading to false positive are considered wasted. Reducing such iterations increases the computational efficiency of a method, meaning that less effort is required while maintaining an acceptable sensitivity-specificity ratio.

Moreover, when millions of streamlines are desired in a whole brain tractogram, millions of seeds and hundreds of millions of 3D points are generated, which can be an important computational and memory limitation [Rheault et al., 2017]. In the case of a specific tract-based application, where only a single to a few WM bundles are desired, this whole brain strategy can be sub-optimal both in terms of bundle reconstruction quality and computational efforts. Increasing the number of streamlines to 10M or 100M would fill-up more of this hard-to-track fanning space, but would potentially never reach full extent coverage. Generating more streamlines will artificially increase the density of the central, easy-to-track part of the bundle and generate a large number of false positive streamlines. Advanced filtering techniques such as SIFT, LIFE, COMMIT [Smith et al., 2015a; Pestilli et al., 2014; Daducci et al., 2015] are able to filter some false positives and reduce the density bias by making sure streamlines explain the desired diffusion signal, but at the cost of removing some hard-to-track valid streamlines as well [Maier-Hein et al., 2017].

ROI-based tractography. In the second strategy, a ROI-based tractography strategy uses different masks, filtering ROI pass-ways and varying tractography parameters (step size, curvature, anisotropy thresholds) [Chamberland et al., 2014, 2017] to enhance the quality of results based on neuroanatomical prior knowledge. A ROI-based seeding strategy is an efficient way to quickly target a bundle of interest by removing unnecessary computations needed in whole brain tractography. Such strategy has already been used in multiple research projects to avoid generating unwanted streamlines [Basser et al., 2000; Catani et al., 2002; Mori and van Zijl, 2002; Behrens et al., 2007; Ghaziri et al., 2015; Renaud et al., 2016; Rozanski et al., 2017; Chamberland et al., 2017], mainly using manual delineations, or ROIs obtained from a segmentation tool such as FreeSurfer [Desikan et al., 2006], or the JHU template [Mori et al., 2005].

Here, ROI-based tractography can be more efficient but not necessarily lead to better spatial extent recovery. The poor-spatial-extent problem is temporarily “hidden” because one can now afford to launch millions of seeds just to reconstruct a single bundle. However, the same issues illustrated in Figure 2 remain, i.e. hard-to-track fanning regions are hard to reach and easy-to-track regions are amplified. Also, because of the ROI-based filtering process, all the false positive streamlines generated are thrown out and excluded from the result, but, computational efforts are still deployed to track them. Hence, overall, a ROI-based approach still suffers from poor efficiency, incomplete spatial-coverage and a similar underlying trade-off between sensitivity and specificity.

To address these issues, tractography needs the injection of more information and priors into the streamline tracing process itself. In Maier-Hein et al. [2017], it was clearly shown that following local orientations alone cannot resolve all the potential ambiguity and biases of fiber tracking. Bottlenecks are not tractable using directional information alone.

Several approaches have been proposed to inject more prior knowledge. An automated method called TRACULA [Yendiki et al., 2011] was proposed to include anatomical

priors from anatomical labeling. Pathway labels are used to initialize and constrain the probabilistic tractography algorithm. However, this approach uses a conservative reconstruction model (FACT, [Mori et al., 1999]) and strict anatomical definitions as training sets, which limits the spatial extent of bundles. Further, probability maps are computed instead of streamlines potentially limiting tract-based analyses. More recently, Wasserthal et al. [2018a] proposed a method to directly segment the volume occupied by bundle of interests. However this method does not rely on tractography and does not produce streamlines. Features of the raw diffusion signal are learned and regions of interests segmented where the characteristics of the signal match with bundles of interests.

On the other hand, global tractography algorithms [Kreher et al., 2008; Mangin et al., 2013; Christiaens et al., 2015; Neher et al., 2012] can sometimes overcome difficult local crossing regions, but resulting tractograms do not always represent the expected anatomical complexity of fasciculi. Global tracking algorithms are promising but are known to have their share of open challenges such as heavy computational needs, the generation of a large number of broken streamlines, and difficulty to impose anatomical priors [Maier-Hein et al., 2017; Jeurissen et al., 2017].

Other approaches exist to overcome fiber crossing difficulties. Chamberland et al. [2017] proposed a magnetic tracking (MAGNET) tool to manually influence directions in strategic regions, improving the reconstruction of the optic radiations bundle. MAGNET essentially pulls tractography towards the occipital lobe with an orientation prior, allowing fiber tracking to perform a “U-turn”-like reconstruction needed to fully reconstruct Meyer’s loop. Moreover, Dhollander et al. [2014] also suggested a method to sharpen orientations based on streamline distributions, called *Track-Orientation Distributions (TOD)*. TOD can help enhance tractography to cover longer distances along WM structures and address some of the biases of fiber tracking. However, since streamlines density is not uniform across a tractogram and does not represent the underlying anatomy, using such a method iteratively across WM will increase the density bias in easy-to-track regions. More recently, Wasserthal et al. [2018b] proposed an approach using bundle-specific orientation maps learned from multiple datasets to select a single fiber orientation distribution (FOD) peak in each voxel, thus improving reconstruction of bundles by simplifying regions with complex WM configuration. To the best of our knowledge, these techniques are the only existing methods trying to inject orientational prior knowledge into the tractography process. Other approaches have proposed to include an orientation-based regularization term in the local reconstruction of fiber orientations, but to our knowledge, not in the tractography procedure itself [Reisert and Kiselev, 2011; Portegies et al., 2015].

Tractography is therefore currently blind to the anatomy and does not have access to the anatomical knowledge accumulated over the past several hundred years by the neuroanatomy community. This neuroanatomical information, when available, could enhance tractography. For instance, priors could capture the “where” and “how” of bundles. That is where should the origin and target region of bundles be, and how difficult crossing, fanning, bending regions should be traversed. Our novel Bundle-Specific Tractography (BST) algorithm is designed to directly incorporate these anatomical and orientational priors in the tractography process itself. The main contribution is a novel methodology enhancing local fiber orientation distribution based on WM bundle priors coming from templates of streamlines. BST is fully automatic and yields better spatial

coverage, increases quality in the fanning extents and produces more plausible shape reconstructions of bundles in the centrum semiovale.

2. Methods

2.1. Bundle-Specific Tractography

Our novel bundle-specific tractography (BST) approach is composed of three steps. For each bundle of interest to be reconstructed, one needs to:

1. Build a **template of streamlines** that represents the shape and position of each bundle, covering as much geometric variability as possible (see first row of Figure 3 and Figure 4).
2. Build the **anatomical priors** by incorporating dilated versions of endpoints and spatial coverage maps defined from the template of streamlines in 1) (see second row of Figure 3).
3. Build the **orientation priors** by generating a field of enhanced fiber orientation distributions (E-FOD) from the track-orientation distribution of the template of streamlines in 1) (see Figure 4).

Our BST approach can be applied to any bundle of interest with the previous 3 steps. We now detail each of these steps before describing the datasets and how it was applied to bundles intersecting in the centrum semiovale seen in Figure 1.

Figure 3: The first row shows streamlines from the template on top of the FA template. In the second row, tracking masks (yellow) and seeding endpoints masks (red), automatically extracted from the template of streamlines of the first row.

Figure 4: a) Template in a standard space, b) orientation priors in native space, c) initial FOD from CSD, d) pre-computed enhanced FOD.

171 *Building the templates of streamlines*

172 This initial step is crucial since the template of streamlines affects all additional
 173 orientational and anatomical priors used by the BST algorithm. It involves four technical
 174 steps (see Figure B.9):

- 175 (i) Run a whole-brain or ROI-based tractography technique on the selected subjects.
- 176 (ii) Segment the bundles of interest.
- 177 (iii) Merge all streamlines of each bundle and each subject in a common space.
- 178 (iv) Adjust the density bias of streamlines distribution.

179 *i) Initial tractography.* Our approach works in combination with any existing whole-brain
 180 streamline-based or iterative tractography algorithm [Jeurissen et al., 2017]. BST will
 181 enhance the initial tractography algorithm and retain the properties of the chosen initial
 182 algorithm. Global tracking and graph-based/geodesic techniques could also be casted in
 183 a BST setting but this is outside the scope of the current paper and will be discussed
 184 later.

185 *ii) Segmentation of bundles.* Bundles of interest needed for a particular study are usually
 186 well-defined in terms of anatomical regions they traverse or connect to [Catani et al.,
 187 2002; Catani and De Schotten, 2008]. Moreover, the shape of the bundle is generally
 188 agreed upon by various experts [Catani and De Schotten, 2008; Hau et al., 2016] or have
 189 often been established from *ex vivo* Klingler blunt dissections [Fernández-Miranda et al.,
 190 2015; Benedictis et al., 2016; Hau et al., 2017]. Hence, this segmentation step of bundles
 191 is either done manually or with a semi-automatic filtering or clustering approach. Of

course, if the number of subjects is large, a manual approach can be tedious and prone to human error. Using ROIs defined by in WM and GM atlases to specify a sequence of rules describing a pathway can be a good alternative [Wassermann et al., 2016]. Another segmentation method is to select or discard streamlines based on shape of cluster using models or anatomical *a priori* [Voineskos et al., 2009; Guevara et al., 2011; O’donnell et al., 2013; Chekir et al., 2014; Garyfallidis et al., 2017; Zhang et al., 2018].

iii) Bring streamlines to a common space. A common space needs to be chosen. The usual MNI space is a possibility or a tailored template space defined for a specific group of subjects. Each subject is nonlinearly warped to that common space and the deformation field used to move each bundle of interest from each subject into the common space. The deformation involves transforming each point (vertex) of each streamline, using *tract-math* [Wassermann et al., 2016] or a in-house tool. Once aligned to a common space, bundles of interest are concatenated together into single track file, as seen in Figure 3. The intention of these templates is to represent the general shape and position of bundles.

iv) Correct for the density bias. An individual subject tractogram generally has low density in crossing regions, while extremely dense in regions of one fiber population (easy-to-track parts). The overall shape of a bundle influences this density, i.e. straight bundles will be denser than curved bundles. Since the concept of tract density (streamlines count) in tractography can be a major pitfall that does not accurately represent the underlying WM structure [Jones, 2010; Jbabdi and Johansen-Berg, 2011; Daducci et al., 2016], this non-uniform density bias needs to be removed from the template of streamlines as only the orientation is of interest for our priors. A smart streamline subsampling method based on hierarchical clustering algorithm and minimal direct flip (MDF) distance [Garyfallidis et al., 2012, 2016] is used to overcome this issue. A streamline within 2mm of its neighboring streamlines do not carry new information and is removed, without altering the general shape of the template.

As mentioned before, the template of streamlines essentially captures the *a priori* knowledge of “*where*” and “*how*”, i.e. where is the bundle originating from, where is going to, and how is it supposed to traverse complex crossing/kissing/branching/fanning regions.

Anatomical priors: Tractography map creation

The streamlines in the template can now be displaced using the same point-wise method from the template creation (using the inverse deformation field). Once the template of streamlines is in the subject space, a new tracking mask is automatically generated from the voxels intersected by the template streamlines. The streamlines end-points can also be used to generate a ROI seeding mask. To reduce gyral bias effects and to cover the potential variability across subjects, the seeding and tracking regions are all dilated by 5mm, as seen in Figure 3. These masks are used to generate probabilistic exclusion and inclusion maps as well as an interface GM-WM map needed for anatomically-constrained tractography [Smith et al., 2012; Girard et al., 2014]. To prevent streamlines from being generated outside of the white matter, all tracking masks were constrained by the subject’s white matter mask. This ensures that our method does not enforce its priors over a subject tissue segmentation, and only the general position within the white matter.

236 *Orientational priors: Enhancing fiber orientations*

237 During tractography, streamlines are generated by taking steps using the fiber orien-
238 tation distribution (FOD) to choose the next direction [Jeurissen et al., 2017]. The value
239 of the FOD is used to weigh the probability of picking a direction. The proposed BST
240 approach uses orientation priors estimated from the template of streamlines to modify
241 the weighting according to the general, or more global, *a priori* bundle direction. If a
242 streamline is following a specific fiber population, the most well-aligned FOD lobe needs
243 to be prioritized.

244 To achieve this “prioritization” of FOD lobes, the local orientation histogram from
245 nearby segments of streamlines is computed at every voxel. This method is based on a
246 TOD map estimated from the template of streamlines, as originally introduced [Dhol-
247 lander et al., 2014]. Improvements related to the FOD field aiming to constraint trac-
248 tography spatially were proposed in the past [Portegies et al., 2015; Reisert and Kiselev,
249 2011].

250 Then, at each voxel, a point-wise multiplication of the FOD and TOD orientation
251 priors is performed, followed by a normalization. As a result, when a lobe is well-
252 aligned with the directions of the priors, the values stay the same. This is observed
253 in Figure 5, where the top row shows TOD maps associated to the pyramidal tracts,
254 the corpus callosum and the arching streamlines respectively. The bottom row shows the
255 corresponding enhanced FOD (E-FOD) associated with each bundle intersecting through
256 the centrum semiovale. Therefore, when a streamline reaches this crossing region, the
257 probability of choosing an appropriate direction is increased. The orientational prior will
258 amplify the desired direction present in the TOD map, and thus improve the directionality
259 of streamlines at the crossing. This concept of using *a priori* information as well as the
260 observed signal to increase anatomical validity of the decision is borrowed from Bayesian
261 statistics, which is common in neuroimaging [Woolrich et al., 2009; Friston et al., 2002b,a].

262 If the template of streamlines have crossing or complex fanning, the TOD map will
263 correctly capture this and an appropriate weighting will be achieved. This can be seen in
264 the zoomed picture of Figure 4. On the contrary, lobes that are perpendicular decrease in
265 value as seen in Figure 6. It is important to note that other lobes are not removed, they
266 are simply decreased in amplitude as this information was still present in the diffusion
267 data. As the aim is not to enforced our orientation priors, but simply re-weighting the
268 FOD amplitude, these two scenarios show the impacts in a extreme case and a typical
269 situation.

270 To ensure that non-existent or erroneous information is not created, a minimal lobe-
271 value threshold is set on the sphere for the orientational priors. This is to ensure that if
272 there is no well-aligned lobe with an amplitude high enough compared to the others, the
273 operation will not increase the values in that direction. A similar threshold was used in
274 Raffelt et al. [2017] to exclude orientations that do not likely represent a fiber population,
275 and used to perform peak extraction on FOD, also to exclude unlikely fiber populations
276 [Tournier et al., 2012; Chamberland et al., 2014; Dell’Acqua et al., 2013]. In our case,
277 we set a relative amplitude threshold to 0.2 (or 20%).

278 *2.2. DWI Datasets*

279 For this work the templates and experiments were performed with the BIL&GIN
280 database [Mazoyer et al., 2016]. The datasets are composed of T1-weighted and diffusion-
281 weighted images from 39 healthy subjects. The dMRI acquisition consists of 21 gradient

Figure 5: Orientation distribution of the crossing section presented in Figure 1. The first row illustrates the track orientation distribution (TOD) maps generated from the different template (Pyramidal tract, corpus callosum, Arcuate Fasciculus). In the second row, the first FOD vignette presents the original fiber orientation distribution (FOD) in the crossing region followed by enhanced FOD (E-FOD), a combination of the associated template TOD (from the first row) and the original FOD.

Figure 6: The first row represents a voxel where the FOD (a) does not have any lobe aligned with the orientation prior (b), which results in an enhanced-FOD (c) almost identical to the original. The second row represents a voxel where a FOD (d) with a lobe of sufficient size (above the minimum relative amplitude threshold) globally aligned with the orientation prior (e), which results in an enhanced-FOD (f) with a increase amplitude for that lobe.

282 directions on half a sphere, the same direction were acquired on the other half of the
 283 sphere for averaging, on a single shell ($b = 1000 \text{ s/mm}^2$) with a 2 mm isotropic resolution.
 284 This scheme was acquired twice for averaging to increase SNR. At each voxel, the FOD
 285 was computed using constrained spherical deconvolution (CSD) with spherical harmonics

of order 6 [Tournier et al. \[2008\]](#); [Descoteaux et al. \[2009\]](#).

2.3. Template creation

For each subject, a whole brain tractogram was reconstructed using the anatomically-constrained probabilistic particle filtering tractography (PFT), with recommended default parameters [\[Girard et al., 2014\]](#). Each tractogram was then segmented using an automatic ROI-based method to extract five bundles of interest, these five bundles will simply be referred to as the left/right pyramidal tract (PT), the corpus callosum (CC) and the left/right arcuate fasciculus (AF). The complete anatomical definitions used by the neuroanatomy expert can be found in the [Anatomical Definition](#) Section. We built our own tailored anatomical template using the fractional anisotropy (FA) maps of the 39 subjects with ANTS Multivariate Template Construction script [\[Avants et al., 2008\]](#). This group average template was used as a common space for the template of streamlines. Subsequently, the 5 bundles of interest, from each subject, were moved into that common space using the nonlinear deformation field from ANTS. Once BOI are concatenated across subject, any redundant streamlines, up to a 2mm MDF distance threshold, are discarded. [Moreover, in this work, a leave-one-out approach was used to ensure that the template streamlines generalize to independent datasets. Any experiment using a template performed on a particular subject did not use the subject itself in that template.](#)

2.4. Experiments, Evaluation, and Statistical analysis

Probabilistic (Prob) and deterministic (Det), with or without PFT [\[Girard et al., 2014\]](#) were compared, as well as global tractography from *MITK Diffusion* [\[Neher et al., 2012\]](#). GM-WM interface seeding was performed, with 5 seeds per voxel, adapted to each bundle of interest (BOI) seeding region (BOI map). Three main sets of parameters were used to compare between classical tracking algorithm and the proposed BST method. To distinguish the impact of improvements from bundle-specific tracking anatomical prior masks and orientational prior enhanced-FOD, three tractography reconstructions were performed:

1. Original FOD with the original exclude/include maps
2. Original FOD with bundle-specific exclude/include maps
3. Enhanced FOD with bundle-specific exclude/include maps

Item 1 corresponds to classical HARDI tracking [\[Tournier et al., 2012; Jeurissen et al., 2017\]](#). The last item is the full BST method, while item 2 is a point of comparison to help separate the contributions of the anatomical and orientation priors. Finally, we also run global tractography [\[Kreher et al., 2008; Neher et al., 2012\]](#) to test against the proposed BST methods. Detailed descriptions of the experiments are available in the [Annexes](#) Section. These lead to five bundles, two tracking evolution approaches (Prob and Det), two algorithm (PFT/no-PFT), and the three compared method (classical, BOI map and BOI & E-FOD), for a total of 60 tractography files plus one global tractography result per subject. With these 2379 tractograms (61 tractograms times 39 subjects), the automatic bundle segmentation method mentioned earlier ([Anatomical Definition](#)) was applied to obtain each bundle.

Evaluation and statistical analysis were performed using the following measures: bundle volume, weighted-Dice coefficient, average streamlines length, percentage of valid streamlines and computational performance. The volume was obtained by computing a binary mask of every voxel intersected by streamlines, in a 1mm isotropic grid [Rheault et al., 2017]. The weighted-Dice was introduced by Cousineau et al. [2017] to obtain a value representing spatial agreement that takes account streamlines density. Using the Dice coefficient as a spatial agreement metric will be severely penalized for spurious streamlines. The overall shape of bundles of interests were expected to be similar across subjects. The strict anatomical definitions lead to comparable bundles across subjects once registered in a common space, especially for major WM pathways. In addition, an intra-subject approach would not have provided direct information regarding the quality of a bundle reconstruction. Lack of fanning reaching the cortex could be reproducible and the spatial agreement high without showing improvement compared to the expected shape. Since the bundles of interests in this project (AF, CC, PT) are major WM pathways, they are generally anatomically comparable across subjects, i.e. the spatial agreement is expected to be high, without necessarily reaching a perfect score.

The percentage of valid streamlines is the proportion of generated streamlines respecting the anatomical definition of a bundle from the expert (see Anatomical Definition Section) and a length threshold compared to the total amount of streamlines that were originally generated. All outputs from tractography have to respect the algorithm constraints (tracking mask, angular threshold, spherical function threshold), however the length threshold was applied only for bundles. segmentation. In the context of our experiments, the notion of valid and invalid streamlines vary with the bundle of interest investigated. A valid streamline for a particular BOI will be considered invalid for all other BOI. For example, a BOI interface of 100 voxels with 5 seeds per voxel would launch 500 streamlines, if 25 streamlines are segmented (considered valid), this would amount to 5% of valid streamlines. Computational performance (efficiency), represents the total number of tractography iterations, i.e. tracking steps, leading to valid streamline points. This is slightly different than the previous percentage of valid streamlines measure, especially with the BOI tracking mask, because most invalid streamlines end up having less than 50 steps while valid streamlines end up with more 500 steps. This metric is better than streamlines count to represent the improvement in terms of computational performance.

For statistical analysis, all subjects and bundles were analyzed separately. Then, to highlight general trends, results were averaged. The analysis was performed with all streamlines registered in the FA template space. This alignment, allowed to compute overlaps between subjects and to normalize the values obtained for the volume and average length. This procedure ensures that variations in volume or length does not come from disparities in brain size of the subject but rather from the tracking itself.

3. Results

To simplify reading of tables, graphs and figures, the focus will be on probabilistic tracking with particle filter tractography (PFT), but all evaluation data, graphs and figures are available in the Annexes Section. Observations and trends are similar and as expected. Moreover, MITK Global tractography was also benchmarked on the same measures to ensure an adequate comparison to state-of-the-art tractography algorithms.

Figure 7 shows segmentations of PT, CC, and AF bundles of interest with a probabilistic PFT algorithm, comparing the use of original FOD and PFT maps to the proposed BST approach using the enhanced FOD with the modified PFT maps. The reconstructed bundles of interest from the classical tracking show some degree of fanning, but the number of streamlines generated was not sufficient to fully recover the spatial extent of bundles. All bundles reconstructed using the masking priors and the enhanced FOD show an increase in both bundle coverage and quality of the fanning. For the PT, the lateral portions of the pre- and post-central gyri are better covered while they are almost non-existent with standard PFT. Similarly, the CC has the fanning fully covering the pre-central and post-central gyri. As for the AF, the spatial extent in the frontal and temporal lobes is increased, but streamlines are also reaching more laterally into the gyri. The quantitative impact of the improved fanning directly impacted the volumes of the reconstructed bundles, as seen in the first column of Table 1.

Figure 8 shows the trends for each measures of the analysis, where each graph represents the changes of all bundles across the three sets of tractography parameters. The impacts are not the same for each bundle, as the shape, size and the underlying anatomy are inherently different. Error bars in the histograms represent the average values and their standard deviation across 39 subjects. As the histograms can be hard to compare, values are also shown in Table 1.

In Figure 8 a), an increase in volume for all bundles is measured when the priors

Figure 7: Examples of resulting PFT and BST algorithms for a single subject. The first row shows results from standard tracking of the PT, CC and AF, while the second row shows results from the proposed bundle-specific tracking. Of note, at no point tracking is allowed outside of the initial white matter tracking mask. It involves that no streamline travel in gray matter or CSF and the impression of going over the sulci which can be observed in the second row is due to coronal view lacking perspective.

Left Pyramidal Tract

	Local model	Mask	Bundle vol. (cm^3)	Dice index	Average length (cm)	% of valid streamlines	Efficiency (%)
1.	FOD	WM	9.80 ± 3.42	0.63 ± 0.12	12.58 ± 0.38	0.29 ± 0.20	1.30 ± 0.92
2.	FOD	BOI	14.70 ± 4.00	0.71 ± 0.10	12.66 ± 0.32	0.67 ± 0.38	4.01 ± 2.45
3.	E-FOD	BOI	27.95 ± 6.24	0.86 ± 0.06	12.16 ± 0.30	3.92 ± 1.95	11.73 ± 6.49

Right Pyramidal Tract

	Local model	Mask	Bundle vol. (cm^3)	Dice index	Average length (cm)	% of valid streamlines	Efficiency (%)
1.	FOD	WM	6.56 ± 2.74	0.51 ± 0.15	12.03 ± 0.32	0.14 ± 0.10	0.58 ± 0.42
2.	FOD	BOI	11.08 ± 3.42	0.64 ± 0.13	12.07 ± 0.26	0.38 ± 0.22	2.21 ± 1.32
3.	E-FOD	BOI	25.27 ± 5.33	0.83 ± 0.06	11.59 ± 0.24	2.42 ± 1.19	7.26 ± 3.95

Corpus Callosum

	Local model	Mask	Bundle vol. (cm^3)	Dice index	Average length (cm)	% of valid streamlines	Efficiency (%)
1.	FOD	WM	25.78 ± 6.58	0.65 ± 0.11	11.25 ± 0.37	0.72 ± 0.33	2.89 ± 1.39
2.	FOD	BOI	49.51 ± 7.40	0.78 ± 0.07	11.49 ± 0.31	2.68 ± 0.85	14.09 ± 5.17
3.	E-FOD	BOI	87.23 ± 7.71	0.86 ± 0.06	11.53 ± 0.26	12.20 ± 3.46	30.32 ± 9.96

Left Arcuate Fasciculus

	Local model	Mask	Bundle vol. (cm^3)	Dice index	Average length (cm)	% of valid streamlines	Efficiency (%)
1.	FOD	WM	54.21 ± 9.49	0.70 ± 0.08	10.61 ± 1.30	1.23 ± 0.45	4.92 ± 1.94
2.	FOD	BOI	56.27 ± 10.06	0.70 ± 0.08	10.78 ± 1.26	1.48 ± 0.55	7.99 ± 3.28
3.	E-FOD	BOI	110.03 ± 10.23	0.75 ± 0.06	12.29 ± 0.72	9.74 ± 2.80	32.26 ± 11.14

Right Arcuate Fasciculus

	Local model	Mask	Bundle vol. (cm^3)	Dice index	Average length (cm)	% of valid streamlines	Efficiency (%)
1.	FOD	WM	35.89 ± 9.97	0.58 ± 0.13	9.35 ± 2.44	0.77 ± 0.36	2.71 ± 1.60
2.	FOD	BOI	37.99 ± 10.26	0.59 ± 0.13	9.55 ± 2.37	0.95 ± 0.44	4.62 ± 2.69
3.	E-FOD	BOI	81.84 ± 12.99	0.80 ± 0.06	11.60 ± 0.82	5.95 ± 2.46	18.89 ± 9.15

Table 1: Quantitative results are shown separately for each bundle of interest from three experiments, measures of the 39 validation subjects were averaged together. Values for average and standard deviation are formatted to simplify interpretation. All measures, except average length, are improved by our approach when compared to classical tractography (approach 1 and 2), the difference is statistically significant ($p < 0.01$). Average length remains stable across method, however it was significantly higher for both arcuate fasciculus.

are used. The AF are the only bundles with no significant increase between the tracking using no *a priori* masking and the tracking using only the *a priori* masking, however the volume significantly increases when enhanced FOD are used. The standard deviation, for methods 2 and 3, in proportion to the average, is decreasing. The volume of the reconstructed PT using method 3 is approximately 400% of the volume obtained from classical tracking, 300% for the CC and 200% for the AF.

The reproducibility across datasets was measured by the Weighted-Dice coefficient. Tractography without priors poorly performed with values between 0.5 (PT right) and 0.7 (AF left). In Figure 8 b), an increase in overlap for all BOI is observed when E-FOD are used, with Weighted-Dice values all more or less 0.8. Similarly to the volume, the AF are the only bundles with no significant increase between methods 1 and 2, while

the overlap significantly increases when enhanced FOD are used. For methods 2 and 3, the standard deviation of the Weighted-Dice coefficient is decreasing as the average coefficient is increasing. For all bundles, except the left AF, the Weighted-Dice increases by 20% when method 3 is used instead of the classical tracking.

The average length and standard deviation was computed to evaluate the shape

Figure 8: Quantitative measures for each BOI across the 3 tracking algorithms. Results from global tractography is added to provide more context, but comparison is partly "unfair" as many variables cannot be controlled in the same way. The Results are shown for probabilistic tracking with PFT. Graphs for deterministic and probabilistic versions of the algorithms with and without PFT are also available in the [Annexes](#) Section

coherence of each bundle. Globally, the average length of most BOI remained similar for each method, as seen in Figure 8 c). However, for the AF, going from methods 2 to 3, the average length increased and the standard deviation decreased. This can be observed more easily in Table 1. As mentioned earlier (Figure 7), this is due to streamlines reaching further inside the lateral gyri.

Finally Figure 8 d) and e) and the last two columns in Table 1 show measures of efficiency. The use of adapted tracking masks (method 2) is not remarkably influencing the percentage of valid streamlines. However, integrating enhanced FOD (method 3) significantly increases efficiency. A tractogram with one million streamlines, each streamline with more or less one hundred points, would need one hundred million iterations. However, not all streamlines have the same number of points depending on their length. When adapted tracking masks (method 2) are used, most invalid streamlines are quickly stopped and thus a large number of iterations avoided. Also, it is important to note that the percentage of valid streamlines and efficiency are improved but that they are still under 15% and 40% respectively, which leaves plenty of room for further improvements.

Similar trends can be observed in the Annexes Section for probabilistic tractography (without particle filtering). Bundle-specific tractography also improves reconstruction when using deterministic algorithms, the difference between classical deterministic and deterministic bundle-specific tractography was statistically significant ($p < 0.01$). This shows that most algorithms using FOD as an input need orientational priors to better reconstruct bundle of interest.

Results from MITK global tractography [Neher et al., 2012] were overall superior to all classic tractography algorithms used for the experiments, as seen in Annexes Figures B.10, B.11, B.12, B.13. However, bundle-specific probabilistic tractography always generate bundles with higher volume and higher reproducibility across subjects compared to global tractography. The orientational prior used with probabilistic tractography (with or without PFT) achieve better results while decreasing computational needs.

4. Discussion

Our bundle-specific tractography algorithm is designed to address the poor-spatial-extent challenge and the sensitivity-specificity trade-off of tractography. Contributions are: i) novel bundle-specific anatomical and orientational priors directly incorporated in the tractography process to boost streamline tracking in hard-to-track regions, ii) results that show increased reproducibility, sensitivity, and specificity of PT, CC, AF bundle reconstructions in the complex centrum semiovale region, and iii) a new BST algorithm with increased efficiency and reduced overall computation time. The main methodological innovation is the novel local fiber orientation distribution enhancement methodology based on white matter bundle priors. These additions were shown to yield better spatial coverage of the pyramidal tract, corpus callosum, and arcuate fasciculus, increasing the quality of the extent of fanning reaching the cortex, helping to accurately represent the shape of each fascicle. Increase in volume is not a direct measure of the quality of a bundle reconstruction. The use of a strict anatomical definition based on the literature is fundamental to correctly address this concern. The definition was designed to reduce as much as possible the number of false positives, meaning that significant

increase in volume cannot be accounted by an increase in outliers, spurious streamlines or contributions from other bundles. As mentioned earlier, tracking is never allowed outside of the initial white matter tracking mask. Hence, increase in volume cannot come from “new voxels” added by our algorithm. The increase in volume was observed in regions that initially lack fanning reaching the cortex, which is a common difficulty for classical tractography [Girard et al., 2014; St-Onge et al., 2017]. It is important to note that the FOD enhancement affects the mathematical properties of the FOD. The E-FOD should not be used to compute FOD metrics such as the AFD [Raffelt et al., 2017] or even NuFO [Tournier et al., 2008] since lobes amplitude changed and no longer agree with the DWI signal.

Template creation needed for Bundle-Specific Tractography

The creation of an appropriate template of streamlines is crucial for the proposed method, as mentioned in the Methods Section. The input bundle of interest of the template needs to be strictly segmented and cleaned of spurious streamlines, while still representing all the potential anatomical variability across subjects. We acknowledge that anatomical definition of WM bundles can vary among experts and that there exist several open controversies in the WM neuroanatomy literature [Forkel et al., 2014; Meola et al., 2015; Hau et al., 2017]. However, having a consensus on the definition of the bundle is not necessary to showcase the potential of our Bundle-Specific Tractography. Neuroanatomists can debate and converge on the precise definitions of anatomical bundles, which can then be adjusted into our BST method.

Moreover, the nonlinear deformation of the template to the subject’s native space is done under the assumption that ANTS performs an adequate registration in-between FA maps. In the case of a different study, with pathology or large brain alterations, such as a tumor, cyst, severe atrophy, or missing brain regions, the quality of the alignment would need to be guaranteed manually. Using a tailored template based on a specific cohort enables our proposed BST method to be fully automatic without relying on manually placed ROIs to enforce certain directions and ending regions. Pathological brains already raise numerous challenges for tissue segmentation, registration, classical tractography and analysis, which all require case-by-case decisions and tuning. For example, WM lesions for Alzheimer’s disease, multiple sclerosis, traumatic brain injuries do not have the same underlying causes and tractography could sometimes be allowed to go through them or not. When all conditions are met and potential pitfalls related to the pathology are accounted for, BST can yield interesting results and can have positive impacts for challenging datasets, as illustrated on the fornix reconstruction in an aging study [Rheault et al., 2018].

As atlases of streamlines become more widely available, such as the one of Yeh and Tseng [2011]; Wasserthal et al. [2018a], different research groups will be able to use these without having to build their own. As an attempt to facilitate its use and guarantee the quality of the anatomical priors for well-known pathways, a WM atlas dedicated to the BST has been designed. This is a step in the right direction to include anatomical priors to improve tractography and address some of the challenges raised in Maier-Hein et al. [2017]. To facilitate future investigations and help other researchers to use our method, a *Nextflow* pipeline [Di Tommaso et al., 2017] has been developed. By providing configuration files, a lightweight WM atlases of bundles and a singularity environment [Kurtzer et al., 2017], this pipeline will simplify the usage of our method. Also, researchers

with specific needs can easily customize the pipeline and the configuration file and obtain exactly what they need.

Bundle volume and spatial coverage

The priors on the regions of interests and local orientations noticeably improve the virtual dissection results by increasing the total volume and the fanning extent. FOD enhancement helps to overcome tracking difficulties in the crossing regions, allowing more streamlines to respect the anatomical definition of the bundle of interest (see [Anatomical Definition](#) Section). Difficulties encountered in the crossing region would have required a higher number of seeds per voxel to obtain an appropriate reconstruction for the classical method, however this approach does not ensure to fully capture the spatial extent. When FODs are enhanced to support the main directionality of a bundle of interest, crossing regions become less disorienting during the tractography, thus increasing the number of streamlines to be considered during segmentation. All this while being more efficient, in term of valid streamlines after segmentation.

As the fanning component of the bundles represent the majority of their volume, improved fanning has a quantitative impact on the volume. For example, the fanning of the PT and CC occurs in the 3-way crossing region (Figure 1), traditionally resulting in a small preponderance of streamlines reaching lateral portions of the primary motor and somatosensory cortex. The priors largely impact the volume of the bundles (first column of Table 1). The use of *a priori* seeding tracking masks increases volume as particle filtering tractography enforces streamline termination in GM regions. The usage of enhanced FOD increases the volume even more by reducing the potential confusion in crossings, as picking a bad direction at each step quickly accumulates in terms of error.

Reproducibility

Bundles of interest considered in this project are known to have anatomical variability at the micro-scale, but at the macro-scale (on the whole bundle level) the AF, CC and PT are expected to cover the same regions (once in a common space) after the segmentation described in the [Anatomical Definition](#) Section. We acknowledge that an identical coverage is not expected, meaning the weighted-dice should not be expected to be 1.0. However, poor spatial agreement (inter-subject) with the classical tractography method mainly comes from poor fanning, a typical solution to this problem is to seed more until the expected shape is obtained and fanning is adequate. The low weighted-dice value seen from classical method is influenced by the high variability of the fanning (CC, PT) or from the early-stop of streamlines in the frontal and temporal lobe (AF). The goal was not to obtain a value of 1.0. The goal was to reduce the variability in these regions as they were caused by tractography limitations and not explained by the underlying anatomy. Intra-subject reproducibility could have hidden this pitfall as a lack of fanning could be reproducible (intra-subject) and would not have reflected if the expected anatomical shape was achieved.

Computational Performance

Immediately stopping any streamline getting out of the bundle of interest region using a priori imposed dilated masks directly reduces the number of tractography iterations and saves computation time without altering the resulting dissection. Such stopping criteria also decreases the rate of false positives, as streamlines must follow a reasonable

path to be considered valid. The efficiency of BST is also increased since streamlines follow the main direction of the BOI instead of leaving prematurely the tracking mask without reaching ROI endpoints. By following the appropriate direction in the crossing, the streamlines have more chance to respect all the bundle definitions, resulting in a higher amount of valid streamlines.

Performance is not just about number of streamlines. The increase in performance combined with the increase in volume and overlap between subjects (Table 1) shows that using the proposed method helps to reach stable results more rapidly, while improving the quality of the bundle of interest.

Future Work

Since the output of the proposed method is in the form of FOD in a widespread format (spherical harmonics) and binary masks, any algorithm that accepts such input could be easily adapted. Other forms of tracking (Dipy [Garyfallidis et al., 2014], FSL [Behrens et al., 2007], MRtrix [Tournier et al., 2012], Real-time tractography [Chamberland et al., 2014], even Global tractography [Kreher et al., 2008; Christiaens et al., 2015; Reisert et al., 2011]) could be tried. Algorithms designed to add more *a priori* could also be tried alongside enhanced-FOD such as surface-enhanced tractography [St-Onge et al., 2017] and microstructure-informed tractography (AxTract) [Girard et al., 2015]).

5. Conclusion

We developed a new bundle-specific tracking (BST) algorithm incorporating novel anatomical orientational priors directly into the streamline tracing process to address the poor-spatial-extent challenge and sensitivity-specificity trade-off of WM bundle reconstruction using tractography. This fully automatic method exploits information extracted from a template of streamlines of bundles of interest to enhance the local modeling in the desired direction using the *a priori* fiber orientation distribution. It was shown that the proposed approach improves the spatial coverage and increases the quality of the fanning in crossing regions, while reducing computational need. Since a stable volume and sufficient anatomically valid streamlines are reconstructed faster than with classical techniques, this method helps the exploration of structural connectivity with more confidence and less biases in white matter bundles metrics. This could have a positive impact on the neurosciences community using dMRI tractography tract-based and connectivity-based analyses.

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References

- Avants, B.B., Epstein, C.L., Grossman, M., Gee, J.C., 2008. Symmetric diffeomorphic image registration with cross-correlation: evaluating automated labeling of elderly and neurodegenerative brain. *Medical image analysis* 12, 26–41.

583 Basser, P.J., Pajevic, S., Pierpaoli, C., Duda, J., Aldroubi, A., 2000. In vivo fiber tractography using
584 dt-mri data. *Magnetic resonance in medicine* 44, 625–632.

585 Behrens, T.E., Berg, H.J., Jbabdi, S., Rushworth, M.F., Woolrich, M.W., 2007. Probabilistic diffusion
586 tractography with multiple fibre orientations: What can we gain? *Neuroimage* 34, 144–155.

587 Benedictis, A., Petit, L., Descoteaux, M., Marras, C.E., Barbareschi, M., Corsini, F., Dallabona, M.,
588 Chioffi, F., Sarubbo, S., 2016. New insights in the homotopic and heterotopic connectivity of the
589 frontal portion of the human corpus callosum revealed by microdissection and diffusion tractography.
590 *Human brain mapping* 37, 4718–4735.

591 Calamante, F., Smith, R.E., Tournier, J.D., Raffelt, D., Connelly, A., 2015. Quantification of voxel-wise
592 total fibre density: investigating the problems associated with track-count mapping. *Neuroimage* 117,
593 284–293.

594 Catani, M., Allin, M.P., Husain, M., Pugliese, L., Mesulam, M.M., Murray, R.M., Jones, D.K., 2007.
595 Symmetries in human brain language pathways correlate with verbal recall. *Proceedings of the National*
596 *Academy of Sciences* 104, 17163–17168.

597 Catani, M., De Schotten, M.T., 2008. A diffusion tensor imaging tractography atlas for virtual in vivo
598 dissections. *cortex* 44, 1105–1132.

599 Catani, M., Howard, R.J., Pajevic, S., Jones, D.K., 2002. Virtual in vivo interactive dissection of white
600 matter fasciculi in the human brain. *Neuroimage* 17, 77–94.

601 Chamberland, M., Scherrer, B., Prabhu, S.P., Madsen, J., Fortin, D., Whittingstall, K., Descoteaux,
602 M., Warfield, S.K., 2017. Active delineation of meyer’s loop using oriented priors through magnetic
603 tractography (magnet). *Human brain mapping* 38, 509–527.

604 Chamberland, M., Whittingstall, K., Fortin, D., Mathieu, D., Descoteaux, M., 2014. Real-time multi-
605 peak tractography for instantaneous connectivity display. *Frontiers in neuroinformatics* 8, 59.

606 Chekir, A., Descoteaux, M., Garyfallidis, E., Côté, M.A., Boumghar, F.O., 2014. A hybrid approach
607 for optimal automatic segmentation of white matter tracts in hardi, in: *Biomedical Engineering and*
608 *Sciences (IECBES), 2014 IEEE Conference on, IEEE*. pp. 177–180.

609 Chenot, Q., Tzourio-Mazoyer, N., Rheault, F., Descoteaux, M., Crivello, F., Zago, L., Mellet, E., Jobard,
610 G., Joliot, M., Mazoyer, B., et al., 2018. A probabilistic atlas of the human pyramidal tract in 410
611 healthy participants. *bioRxiv* , 251108.

612 Christiaens, D., Reisert, M., Dhollander, T., Sunaert, S., Suetens, P., Maes, F., 2015. Global tractog-
613 raphy of multi-shell diffusion-weighted imaging data using a multi-tissue model. *Neuroimage* 123,
614 89–101.

615 Côté, M.A., Girard, G., Boré, A., Garyfallidis, E., Houde, J.C., Descoteaux, M., 2013. Tractometer:
616 towards validation of tractography pipelines. *Medical image analysis* 17, 844–857.

617 Cousineau, M., Jodoin, P.M., Garyfallidis, E., Côté, M.A., Morency, F.C., Rozanski, V., GrandMaison,
618 M., Bedell, B.J., Descoteaux, M., 2017. A test-retest study on parkinson’s pmi dataset yields
619 statistically significant white matter fascicles. *NeuroImage: Clinical* 16, 222–233.

620 Daducci, A., Dal Palú, A., Descoteaux, M., Thiran, J.P., 2016. Microstructure informed tractography:
621 pitfalls and open challenges. *Frontiers in neuroscience* 10, 247.

622 Daducci, A., Dal Palú, A., Lemkaddem, A., Thiran, J.P., 2015. Commit: convex optimization modeling
623 for microstructure informed tractography. *IEEE transactions on medical imaging* 34, 246–257.

624 Dayan, M., Monohan, E., Pandya, S., Kuceyeski, A., Nguyen, T.D., Raj, A., Gauthier, S.A., 2016.
625 Profilmetry: a new statistical framework for the characterization of white matter pathways, with
626 application to multiple sclerosis. *Human brain mapping* 37, 989–1004.

627 Dell’Acqua, F., Simmons, A., Williams, S.C., Catani, M., 2013. Can spherical deconvolution provide
628 more information than fiber orientations? hindrance modulated orientational anisotropy, a true-tract
629 specific index to characterize white matter diffusion. *Human brain mapping* 34, 2464–2483.

630 Descoteaux, M., Deriche, R., Knosche, T.R., Anwander, A., 2009. Deterministic and probabilistic
631 tractography based on complex fibre orientation distributions. *IEEE transactions on medical imaging*
632 28, 269–286.

633 Desikan, R.S., Ségonne, F., Fischl, B., Quinn, B.T., Dickerson, B.C., Blacker, D., Buckner, R.L., Dale,
634 A.M., Maguire, R.P., Hyman, B.T., et al., 2006. An automated labeling system for subdividing the
635 human cerebral cortex on mri scans into gyral based regions of interest. *Neuroimage* 31, 968–980.

636 Dhollander, T., Emsell, L., Van Hecke, W., Maes, F., Sunaert, S., Suetens, P., 2014. Track orientation
637 density imaging (todi) and track orientation distribution (tod) based tractography. *NeuroImage* 94,
638 312–336.

639 Di Tommaso, P., Chatzou, M., Floden, E.W., Barja, P.P., Palumbo, E., Notredame, C., 2017. Nextflow
640 enables reproducible computational workflows. *Nature biotechnology* 35, 316–319.

641 Fernández-Miranda, J.C., Wang, Y., Pathak, S., Stefaneau, L., Verstynen, T., Yeh, F.C., 2015. Asym-

metry, connectivity, and segmentation of the arcuate fascicle in the human brain. *Brain Structure and Function* 220, 1665–1680.

Forkel, S.J., de Schotten, M.T., Kawadler, J.M., Dell’Acqua, F., Danek, A., Catani, M., 2014. The anatomy of fronto-occipital connections from early blunt dissections to contemporary tractography. *Cortex* 56, 73–84.

Friston, K.J., Glaser, D.E., Henson, R.N., Kiebel, S., Phillips, C., Ashburner, J., 2002a. Classical and bayesian inference in neuroimaging: applications. *Neuroimage* 16, 484–512.

Friston, K.J., Penny, W., Phillips, C., Kiebel, S., Hinton, G., Ashburner, J., 2002b. Classical and bayesian inference in neuroimaging: theory. *NeuroImage* 16, 465–483.

Garyfallidis, E., Brett, M., Amirbekian, B., Rokem, A., Van Der Walt, S., Descoteaux, M., Nimmo-Smith, I., 2014. Dipy, a library for the analysis of diffusion mri data. *Frontiers in neuroinformatics* 8, 8.

Garyfallidis, E., Brett, M., Correia, M.M., Williams, G.B., Nimmo-Smith, I., 2012. Quickbundles, a method for tractography simplification. *Frontiers in neuroscience* 6, 175.

Garyfallidis, E., Côté, M.A., Rheault, F., Descoteaux, M., 2016. Quickbundlesx: sequential clustering of millions of streamlines in multiple levels of detail at record execution time. *ISMRM2016 (Singapore)*.

Garyfallidis, E., Côté, M.A., Rheault, F., Sidhu, J., Hau, J., Petit, L., Fortin, D., Cunanne, S., Descoteaux, M., 2017. Recognition of white matter bundles using local and global streamline-based registration and clustering. *NeuroImage*.

Ghaziri, J., Tucholka, A., Girard, G., Houde, J.C., Boucher, O., Gilbert, G., Descoteaux, M., Lippé, S., Rainville, P., Nguyen, D.K., 2015. The corticocortical structural connectivity of the human insula. *Cerebral cortex* 27, 1216–1228.

Girard, G., Fick, R., Descoteaux, M., Deriche, R., Wassermann, D., 2015. Atract: microstructure-driven tractography based on the ensemble average propagator, in: *International Conference on Information Processing in Medical Imaging*, Springer. pp. 675–686.

Girard, G., Whittingstall, K., Deriche, R., Descoteaux, M., 2014. Towards quantitative connectivity analysis: reducing tractography biases. *Neuroimage* 98, 266–278.

Guevara, P., Poupon, C., Rivière, D., Cointepas, Y., Descoteaux, M., Thirion, B., Mangin, J.F., 2011. Robust clustering of massive tractography datasets. *NeuroImage* 54, 1975–1993.

Hau, J., Sarubbo, S., Houde, J.C., Corsini, F., Girard, G., Deledalle, C., Crivello, F., Zago, L., Mellet, E., Jobard, G., et al., 2017. Revisiting the human uncinate fasciculus, its subcomponents and asymmetries with stem-based tractography and microdissection validation. *Brain Structure and Function* 222, 1645–1662.

Hau, J., Sarubbo, S., Perchey, G., Crivello, F., Zago, L., Mellet, E., Jobard, G., Joliot, M., Mazoyer, B.M., Tzourio-Mazoyer, N., et al., 2016. Cortical terminations of the inferior fronto-occipital and uncinate fasciculi: anatomical stem-based virtual dissection. *Frontiers in neuroanatomy* 10, 58.

Jbabdi, S., Johansen-Berg, H., 2011. Tractography: where do we go from here? *Brain connectivity* 1, 169–183.

Jeurissen, B., Descoteaux, M., Mori, S., Leemans, A., 2017. Diffusion mri fiber tractography of the brain. *NMR in Biomedicine*.

Jeurissen, B., Leemans, A., Tournier, J.D., Jones, D.K., Sijbers, J., 2013. Investigating the prevalence of complex fiber configurations in white matter tissue with diffusion magnetic resonance imaging. *Human brain mapping* 34, 2747–2766.

Jones, D.K., 2010. Challenges and limitations of quantifying brain connectivity in vivo with diffusion mri. *Imaging in Medicine* 2, 341.

Kreher, B., Mader, I., Kiselev, V., 2008. Gibbs tracking: a novel approach for the reconstruction of neuronal pathways. *Magnetic Resonance in Medicine* 60, 953–963.

Kurtzer, G.M., Sochat, V., Bauer, M.W., 2017. Singularity: Scientific containers for mobility of compute. *PloS one* 12, e0177459.

Maier-Hein, K.H., Neher, P.F., Houde, J.C., Côté, M.A., Garyfallidis, E., Zhong, J., Chamberland, M., Yeh, F.C., Lin, Y.C., Ji, Q., et al., 2017. The challenge of mapping the human connectome based on diffusion tractography. *Nature communications* 8, 1349.

Mangin, J.F., Fillard, P., Cointepas, Y., Le Bihan, D., Frouin, V., Poupon, C., 2013. Toward global tractography. *Neuroimage* 80, 290–296.

Mazoyer, B., Mellet, E., Perchey, G., Zago, L., Crivello, F., Jobard, G., Delcroix, N., Vigneau, M., Leroux, G., Petit, L., et al., 2016. Bil&gin: a neuroimaging, cognitive, behavioral, and genetic database for the study of human brain lateralization. *Neuroimage* 124, 1225–1231.

Meola, A., Comert, A., Yeh, F.C., Stefanescu, L., Fernandez-Miranda, J.C., 2015. The controversial

existence of the human superior fronto-occipital fasciculus: Connectome-based tractographic study with microdissection validation. *Human brain mapping* 36, 4964–4971.

Mori, S., Crain, B.J., Chacko, V.P., Van Zijl, P.C., 1999. Three-dimensional tracking of axonal projections in the brain by magnetic resonance imaging. *Annals of Neurology: Official Journal of the American Neurological Association and the Child Neurology Society* 45, 265–269.

Mori, S., Wakana, S., Van Zijl, P.C., Nagae-Poetscher, L., 2005. *MRI atlas of human white matter*. Elsevier.

Mori, S., van Zijl, P., 2002. Fiber tracking: principles and strategies—a technical review. *NMR in Biomedicine* 15, 468–480.

Moyer, D., Thompson, P.M., Steeg, G.V., 2018. Measures of tractography convergence. *arXiv preprint arXiv:1806.04634*.

Neher, P.F., Stieltjes, B., Reisert, M., Reicht, I., Meinzer, H.P., Fritzsche, K.H., 2012. Mitk global tractography, in: *Medical Imaging 2012: Image Processing, International Society for Optics and Photonics*. p. 83144D.

O’donnell, L.J., Golby, A.J., Westin, C.F., 2013. Fiber clustering versus the parcellation-based connectome. *NeuroImage* 80, 283–289.

Pestilli, F., Yeatman, J.D., Rokem, A., Kay, K.N., Wandell, B.A., 2014. Evaluation and statistical inference for human connectomes. *Nature methods* 11, 1058.

Portegies, J.M., Fick, R.H.J., Sanguinetti, G.R., Meesters, S.P., Girard, G., Duits, R., 2015. Improving fiber alignment in hardi by combining contextual pde flow with constrained spherical deconvolution. *PloS one* 10, e0138122.

Raffelt, D.A., Tournier, J.D., Smith, R.E., Vaughan, D.N., Jackson, G., Ridgway, G.R., Connelly, A., 2017. Investigating white matter fibre density and morphology using fixel-based analysis. *Neuroimage* 144, 58–73.

Reisert, M., Kiselev, V.G., 2011. Fiber continuity: an anisotropic prior for odf estimation. *IEEE transactions on medical imaging* 30, 1274–1283.

Reisert, M., Mader, I., Anastasopoulos, C., Weigel, M., Schnell, S., Kiselev, V., 2011. Global fiber reconstruction becomes practical. *Neuroimage* 54, 955–962.

Renauld, E., Descoteaux, M., Bernier, M., Garyfallidis, E., Whittingstall, K., 2016. Semi-automatic segmentation of optic radiations and lgn, and their relationship to eeg alpha waves. *PloS one* 11, e0156436.

Rheault, F., Houde, J.C., Descoteaux, M., 2017. Visualization, interaction and tractometry: Dealing with millions of streamlines from diffusion mri tractography. *Frontiers in neuroinformatics* 11, 42.

Rheault, F., Roy, M., Cunnane, S., Descoteaux, M., 2018. Bundle-specific fornix reconstruction for dual-tracer pet-tractometry. *bioRxiv*, 423459.

Rozanski, V.E., da Silva, N.M., Ahmadi, S.A., Mehrkens, J., da Silva Cunha, J., Houde, J.C., Vollmar, C., Bötzel, K., Descoteaux, M., 2017. The role of the pallidothalamic fibre tracts in deep brain stimulation for dystonia: a diffusion mri tractography study. *Human brain mapping* 38, 1224–1232.

Smith, R.E., Tournier, J.D., Calamante, F., Connelly, A., 2012. Anatomically-constrained tractography: improved diffusion mri streamlines tractography through effective use of anatomical information. *Neuroimage* 62, 1924–1938.

Smith, R.E., Tournier, J.D., Calamante, F., Connelly, A., 2015a. The effects of sift on the reproducibility and biological accuracy of the structural connectome. *Neuroimage* 104, 253–265.

Smith, R.E., Tournier, J.D., Calamante, F., Connelly, A., 2015b. Sift2: Enabling dense quantitative assessment of brain white matter connectivity using streamlines tractography. *Neuroimage* 119, 338–351.

Song, J.W., Mitchell, P.D., Kolasinski, J., Ellen Grant, P., Galaburda, A.M., Takahashi, E., 2014. Asymmetry of white matter pathways in developing human brains. *Cerebral cortex* 25, 2883–2893.

Sotiropoulos, S.N., Zalesky, A., 2017. Building connectomes using diffusion mri: Why, how and but. *NMR in Biomedicine*.

St-Onge, E., Daducci, A., Girard, G., Descoteaux, M., 2017. Surface-enhanced tractography (set). *NeuroImage*.

Tournier, J., Calamante, F., Connelly, A., et al., 2012. Mrtrix: diffusion tractography in crossing fiber regions. *International Journal of Imaging Systems and Technology* 22, 53–66.

Tournier, J.D., Yeh, C.H., Calamante, F., Cho, K.H., Connelly, A., Lin, C.P., 2008. Resolving crossing fibres using constrained spherical deconvolution: validation using diffusion-weighted imaging phantom data. *Neuroimage* 42, 617–625.

Voineskos, A.N., O’donnell, L.J., Lobaugh, N.J., Markant, D., Ameis, S.H., Niethammer, M., Mulsant, B.H., Pollock, B.G., Kennedy, J.L., Westin, C.F., et al., 2009. Quantitative examination of a novel

760 clustering method using magnetic resonance diffusion tensor tractography. *Neuroimage* 45, 370–376.

761 Wang, R., Benner, T., Sorensen, A.G., Wedeen, V.J., 2007. Diffusion toolkit: a software package for

762 diffusion imaging data processing and tractography, in: *Proc Intl Soc Mag Reson Med*, Berlin.

763 Wassermann, D., Makris, N., Rathi, Y., Shenton, M., Kikinis, R., Kubicki, M., Westin, C.F., 2016. The

764 white matter query language: a novel approach for describing human white matter anatomy. *Brain*

765 *Structure and Function* 221, 4705–4721.

766 Wasserthal, J., Neher, P., Maier-Hein, K.H., 2018a. Tractseg-fast and accurate white matter tract

767 segmentation. *arXiv preprint arXiv:1805.07103* .

768 Wasserthal, J., Neher, P.F., Maier-Hein, K.H., 2018b. Tract orientation mapping for bundle-specific

769 tractography. *arXiv preprint arXiv:1806.05580* .

770 Woolrich, M.W., Jbabdi, S., Patenaude, B., Chappell, M., Makni, S., Behrens, T., Beckmann, C.,

771 Jenkinson, M., Smith, S.M., 2009. Bayesian analysis of neuroimaging data in fsl. *Neuroimage* 45,

772 S173–S186.

773 Yeatman, J.D., Dougherty, R.F., Myall, N.J., Wandell, B.A., Feldman, H.M., 2012. Tract profiles of

774 white matter properties: automating fiber-tract quantification. *PloS one* 7, e49790.

775 Yeatman, J.D., Richie-Halford, A., Smith, J.K., Keshavan, A., Rokem, A., 2018. A browser-based tool

776 for visualization and analysis of diffusion mri data. *Nature communications* 9, 940.

777 Yeh, F.C., Tseng, W.Y.I., 2011. Ntu-90: a high angular resolution brain atlas constructed by q-space

778 diffeomorphic reconstruction. *Neuroimage* 58, 91–99.

779 Yendiki, A., Panneck, P., Srinivasan, P., Stevens, A., Zöllei, L., Augustinack, J., Wang, R., Salat, D.,

780 Ehrlich, S., Behrens, T., et al., 2011. Automated probabilistic reconstruction of white-matter pathways

781 in health and disease using an atlas of the underlying anatomy. *Frontiers in neuroinformatics* 5, 23.

782 Zhang, F., Wu, W., Ning, L., McNulty, G., Waber, D., Gagoski, B., Sarill, K., Hamoda, H.M., Song,

783 Y., Cai, W., et al., 2018. Suprathreshold fiber cluster statistics: Leveraging white matter geometry

784 to enhance tractography statistical analysis. *NeuroImage* .

785 Zhang, Y., Zhang, J., Oishi, K., Faria, A.V., Jiang, H., Li, X., Akhter, K., Rosa-Neto, P., Pike, G.B.,

786 Evans, A., et al., 2010. Atlas-guided tract reconstruction for automated and comprehensive examina-

787 tion of the white matter anatomy. *Neuroimage* 52, 1289–1301.

788 Appendix A. Anatomical Definition

789 The PT was extracted following the anatomical definition proposed by [Chenot et al.](#)
790 [\[2018\]](#) based on a precise manual positioning of individual subcortical regions of interest
791 (ROI) along the descending pathway of the PT. Three ROIs were therefore drawn in
792 each hemisphere within the internal capsule, the midbrain and the medulla oblongata
793 using both b0 and color FA maps. This was performed with TrackVis [\[Wang et al., 2007\]](#).
794 Additional anatomical ROIs were used from the JHU template [\[Zhang et al., 2010\]](#) once
795 warped to the native diffusion space of each subject using ANTS non-linear registration.
796 Streamlines passing through the 3 ROIs, terminating either in the frontal lobe or the
797 parietal lobe, and not terminating in the cerebellum were considered to compose the PT.

798 The AF was extracted in each hemisphere from a first set of streamlines with one
799 termination in the frontal lobe and the other either in the temporal lobe or in the oc-
800 cipital lobe. Then, streamlines that do not pass through the ventral part of the external
801 capsule (vEC) were considered as composing the AF. Such streamlines are anatomically
802 constraint to leave the frontal lobe from the superior part of the external capsule and
803 then present an arcuate shape defining the AF. The vEC ROI was composed of the region
804 of the JHU template located in the ventral part of the external capsule, namely the infe-
805 rior fronto-occipital and uncinate ROIs. Note that frontal, temporal and occipital lobes
806 were composed of the frontal, temporal and occipital cortical gray matter and superficial
807 white matter regions of the JHU template, respectively.

808 The CC was extracted from a first set of commissural streamlines with one termination
809 in each hemisphere. A precentral-postcentral ROI, composed of the gray matter and
810 superficial white matter part of the precentral and postcentral gyri of the JHU template
811 was created for each hemisphere. Then, the present CC bundle was composed of callosal
812 streamlines with one termination in each of the precentral-postcentral ROI and passing
813 through the corpus callosum.

814 It is important to mention that streamlines considered for segmentation come from
815 a tractogram generated with certain constraints and conditions. All streamlines are
816 within the subject white matter, respect the length threshold (20mm-200mm), the an-
817 gular threshold (45 degrees cone aperture for probabilistic and 20 degrees cone aperture
818 for deterministic) and the minimum relative spherical function threshold of 0.1)

819 Appendix B. Annexes

820 Appendix B.1. Pipeline overview

Figure B.9: Overview of the template creation pipeline. From whole brain tractogram (a) bundles are extracted (b) and then warped and concatenated in a common space (c), finally the density bias is reduced by a spatial subsampling (d)

821 Appendix B.2. Results from all tractography algorithms

822 In this section results from all tractography type are shown, comparaison to global
823 tractography from the Medical Imaging Toolkit (MITK) [Neher et al., 2012; Kreher et al.,
824 2008] was added. The tracking was performed with the original FOD and WM mask,
825 default parameters and 1×10^8 iterations. For a fair comparaison, each subject whole brain
826 tractogram was filtered to be similar to the classic tractography. Tractogram was forced
827 to end in the BOI adapted interface, subsampled to the same number of streamlines and
828 then segmented using the same automatic method.

Figure B.10: Results for each BOI across the 4 tracking variations with probabilistic particle filtering tractography, where the fourth method is the global tracking.

Figure B.11: Results for each BOI across the 4 tracking variations with probabilistic tractography, where the fourth method is the global tracking.

Figure B.12: Results for each BOI across the 4 tracking variations with deterministic particle filtering tractography, where the fourth method is the global tracking.

Figure B.13: Results for each BOI across the 3 tracking variations with deterministic tractography, where the fourth method is the global tracking.