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Registration and Analysis of White Matter Group Differences with a Multi-Fiber Model

Maxime Taquet^{1,2}, Benoît Scherrer¹, Olivier Commowick³, Jurriaan Peters^{1,4},
Mustafa Sahin⁴, Benoît Macq², and Simon K. Warfield¹

¹ Computational Radiology Laboratory, Children’s Hospital Boston, Harvard, USA

² ICTEAM Institute, Université catholique de Louvain, Louvain-La-Neuve, Belgium

³ INRIA, INSERM, VisAGeS U746 Unit/Project, F-35042 Rennes, France

⁴ Department of Neurology, Children’s Hospital Boston, Harvard, USA

Abstract. Diffusion magnetic resonance imaging has been used extensively to probe the white matter in vivo. Typically, the raw diffusion images are used to reconstruct a diffusion tensor image (DTI). The incapacity of DTI to represent crossing fibers led to the development of more sophisticated diffusion models. Among them, multi-fiber models represent each fiber bundle independently, allowing the direct extraction of diffusion features for population analysis. However, no method exists to properly register multi-fiber models, seriously limiting their use in group comparisons. This paper presents a registration and atlas construction method for multi-fiber models. The validity of the registration is demonstrated on a dataset of 45 subjects, including both healthy and unhealthy subjects. Morphometry analysis and tract-based statistics are then carried out, proving that multi-fiber models registration is better at detecting white matter local differences than single tensor registration.

Keywords: Diffusion Imaging, Multi-Fiber Models, Registration, White Matter

1 Introduction

Diffusion magnetic resonance imaging offers the ability to investigate in vivo the white matter microstructure. The representation of the signal by diffusion tensor images (DTI) has proven useful for population analysis in two ways [1]. First, scalar features extracted from DTI, such as the fractional anisotropy (FA), may indicate the presence of brain diseases. Second, the use of DTI in registration improves the detection of morphometric differences, compared to scalar images.

The single tensor diffusion model has, however, proven inaccurate for two main reasons. First, it cannot represent the signal arising from multiple fibers with heterogeneous orientations in one voxel. Second, it does not account for the non-monoexponential decay observed when imaging at high b-values. Novel models addressing one or both of these issues have been introduced [2] : Q-ball imaging, spherical deconvolution, 4th order tensors, DOT, and others. Most of them focus on describing the general shape of the diffusion at each voxel.

neighborhood), smoothing (the value at a grid voxel is replaced by a weighted combination of the values in a neighborhood) and atlas construction (the value at one voxel is the average of the values in the aligned subjects’).

Gaussian mixture simplification (GMS) was introduced to efficiently compute weighted combinations of diffusion mixture [7]. In this section, we underline the important aspects of this method. The idea behind GMS is that computing weighted combinations of mixtures would be trivial if the number of components of the result could be arbitrarily large. Indeed, the linear combination of K mixture models with N components is a mixture models with $M = KN$ components:

$$\mathbf{M}_C = \sum_{k=1}^K w_k \mathbf{M}_k = \sum_{k=1}^K w_k \sum_{j=1}^N f_j^k S_j^k(\mathbf{x}) = \sum_{i=1}^M g_i S_i(\mathbf{x}). \quad (2)$$

We refer to this mixture as the *complete* mixture. GMS optimizes the parameters of a *simplified mixture* $\mathbf{M}_S = \sum_{j=1}^N h_j R_j(\mathbf{x})$ with $N < M$ components that best approximates \mathbf{M}_C . The energy function to be minimized is the cumulative differential entropy (the reference to the location \mathbf{x} is omitted for clarity):

$$D(\mathbf{M}_C, \mathbf{M}_S) = \sum_{j=1}^N \sum_{i:\pi_i=j} g_i D(S_i || R_j) = \sum_{j=1}^N \sum_{i:\pi_i=j} g_i \int S_i(\mathbf{g}) \log \frac{S_i(\mathbf{g})}{R_j(\mathbf{g})} d\mathbf{g}, \quad (3)$$

where \mathbf{g} is the gradient vector and where latent variables π_i cluster the components of the complete mixture S_i in N clusters each represented by a single component of the simplified mixture, R_j ; $\pi_i = j$ means that S_i is best represented by R_j . Following the recent developments in probabilistic clustering, an EM scheme is used to minimize (3). Banerjee *et al* showed that both the E-step and the M-step can be solved in closed form for mixtures of exponential distributions [8]. For Gaussian mixtures, the E-step consists in optimizing the latent variables π_i by computing the Burg matrix divergence between the covariance matrices of each component of \mathbf{M}_C (Σ_i^S) and each component of \mathbf{M}_S (Σ_j^R):

$$\pi_i = \arg \min_j B(\Sigma_i^S, \Sigma_j^R) = \arg \min_j \text{Tr}(\Sigma_i^S \Sigma_j^{R-1}) - \log |\Sigma_i^S \Sigma_j^{R-1}|. \quad (4)$$

As for the M-step, it sums up to calculating:

$$\Sigma_j^R = \frac{\sum_{i:\pi_i=j} f_i \Sigma_i^S}{\sum_{i:\pi_i=j} f_i} \quad \text{and} \quad h_j = \sum_{i:\pi_i=j} f_i. \quad (5)$$

Alternating (4) and (5) until convergence provides the parameters (h_j and Σ_j^R) of the resulting mixture. A log-Euclidean version of this interpolation scheme is obtained by replacing all covariance matrices by their logarithm.

4 Generalized Correlation Coefficient for Mixtures

The correlation coefficient, invariant under linear transformations of the voxel intensities, is widely used in mono-modal image registration. The inter-subject

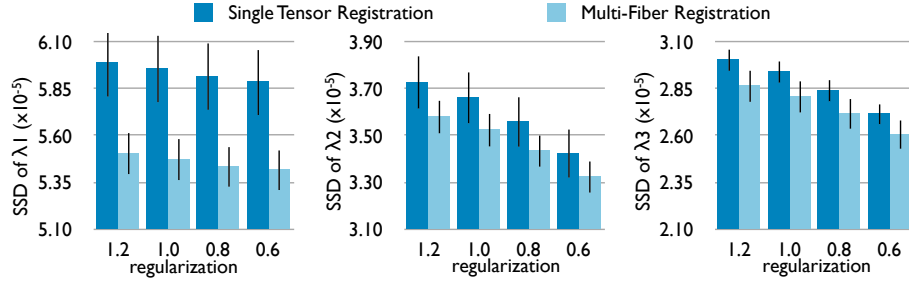


Fig. 1. Comparison of the single tensor and multi-fiber registration in terms of the SSD between eigenvalues after alignment, for different regularization parameter values [9]. Multi-fiber registration significantly improves the quality of the registration.

variability of diffusivity values motivates the introduction of a generalized correlation coefficient, invariant under these differences. In DTI, this variability has been reported and partially accounted for in some registration methods [1]. The correlation coefficient between blocks F and G is defined as the scalar product of the normalized blocks:

$$\rho(F, G) = \left\langle \frac{F - \mu_F}{\|F - \mu_F\|}, \frac{G - \mu_G}{\|G - \mu_G\|} \right\rangle,$$

where μ_F is the mean of the image values in the block. It is invariant if F (and/or G) is replaced by $aF + b$. It has been generalized to vector images by redefining the means μ_F and μ_G as the projection of the block onto a constant block T [10]:

$$F - \mu_F = F - F, T \frac{T}{\|T\|^2}.$$

The corresponding generalized correlation coefficient is invariant if F is replaced by $aF + bT$ where T is now any constant vector block. The definition of a scalar product between two blocks of mixture models seems impractical if not impossible. We therefore further generalize the correlation coefficient by substituting the inner product by a more general scalar mapping, $m(\mathbf{M}_f, \mathbf{M}_g)$:

$$\rho(\mathbf{M}_f, \mathbf{M}_g) = m\left(\frac{\mathbf{M}_f - m(\mathbf{M}_f, \mathbf{T})\mathbf{T}}{n_m(\mathbf{M}_f - m(\mathbf{M}_f, \mathbf{T})\mathbf{T})}, \frac{\mathbf{M}_g - m(\mathbf{M}_g, \mathbf{T})\mathbf{T}}{n_m(\mathbf{M}_g - m(\mathbf{M}_g, \mathbf{T})\mathbf{T})}\right),$$

where $n_m(\mathbf{M})^2 = m(\mathbf{M}, \mathbf{M})$ is a generalization of the norm. This definition does not guarantee the invariance property of the metric for any scalar mapping. One can show that the invariance is preserved as long as the scalar mapping is linear with respect to the constant block \mathbf{T} :

$$m(a\mathbf{M}_f + b\mathbf{T}, \mathbf{T}) = a m(\mathbf{M}_f, \mathbf{T}) + b m(\mathbf{T}, \mathbf{T}). \quad (6)$$

To preserve the interpretability of ρ as a similarity metric, it needs to be symmetric, equal to one in case of perfect match and lower than one in any other

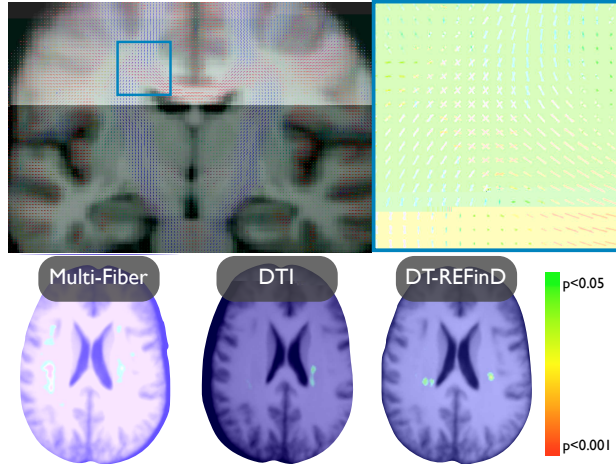


Fig. 2. (top) The two-tensor atlas built by means of the developed registration method reveals crossing pathways common to all anatomies. (bottom) White matter volume shrinkage in tuberous sclerosis represented by the p-value maps. Multi-fiber registration reveals more differences than single tensor registration and DT-REFinD [11]

case. These constraints on ρ translate into the following constraints on m :

$$m(\mathbf{M}_f, \mathbf{M}_g) = m(\mathbf{M}_g, \mathbf{M}_f) \quad (7)$$

$$n_m(a\mathbf{M}_f) = a n_m(\mathbf{M}_f) \quad (8)$$

$$|m(\mathbf{M}_f, \mathbf{M}_g)| \leq n_m(\mathbf{M}_f)n_m(\mathbf{M}_g). \quad (9)$$

The latter is a generalized form of the Cauchy-Schwartz inequality for inner products. Conditions (6-9), the choice of \mathbf{T} and the definitions of the addition $\mathbf{M} + \mathbf{T}$ and multiplication by a scalar $a\mathbf{M}$, stand together as a model to define a correlation coefficients in potentially any space. For DTI, if \mathbf{T} is an isotropic tensor block ($\mathbf{T}(\mathbf{x}) = D\mathbf{I}_{3 \times 3}$), m is the log-Euclidean scalar product, and the log-Euclidean algebra is used, then ρ is invariant under linear transformations of the eigenvalues in the log-domain [12]. For multi-tensor images, we fix $\mathbf{T}(\mathbf{x}) = \{(\frac{1}{N}, D\mathbf{I}_{3 \times 3}), \dots, (\frac{1}{N}, D\mathbf{I}_{3 \times 3})\}$, and we define the addition of \mathbf{T} , and the multiplication by a scalar component-wise in the log-domain. The scalar mapping $m(\mathbf{M}_f, \mathbf{M}_g)$ is defined by pairing the tensors in each voxel to maximize the linear combination of pairwise scalar products. Let $\mathbf{M}_f(\mathbf{x}) = \{(f_1, \mathbf{F}_1), \dots, (f_N, \mathbf{F}_N)\}$ and $\mathbf{M}_g(\mathbf{x}) = \{(g_1, \mathbf{G}_1), \dots, (g_N, \mathbf{G}_N)\}$ defined on a domain Ω , we have:

$$m(\mathbf{M}_f, \mathbf{M}_g) = \sum_{\mathbf{x} \in \Omega} \max_{\pi} \sum_{i=1}^N f_i g_{\pi(i)} \langle \mathbf{F}_i, \mathbf{G}_{\pi(i)} \rangle,$$

where π is a pairing function associating one tensor of \mathbf{M}_g to each tensor of \mathbf{M}_f . This scalar mapping satisfies conditions (6-9). Interestingly, the resulting generalized correlation coefficient is invariant under any global (within the block)

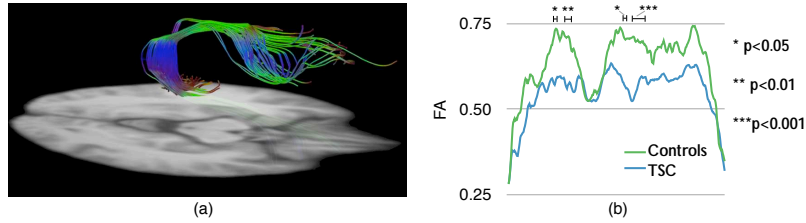


Fig. 3. (a) Arcuate fasciculus, a set of fibers involved in language, on which tract based statistics was performed, (b) The FA profile in TSC patients shows significantly disrupted white matter fascicules in different clusters, indicated by the stars.

linear transformation of all eigenvalues in the log-domain. This similarity metric is therefore robust to the inter-subject variability of diffusivities.

5 Implementation and Complexity

The developed methods were integrated in the efficient block matching registration algorithm described in []. The parameters used are the following: 4 pyramid level, 10 iterations per level, block size: $5 \times 5 \times 5$, outlier removal rate: 2%. The implementation was multi-threaded. On a 8 core workstation, with $224 \times 224 \times 176$ two-fiber images, the entire registration takes 1.5 hour. All weighted combinations were computed until complete convergence of the soft clustering. The average number of iterations required for that convergence is 4.

6 Results

The registration was applied to a clinical dataset of 45 subjects, 13 controls and 32 patients with tuberous sclerosis complex (TSC), a rare genetic disease associated with impaired white matter integrity. A DTI and a multi-tensor model with three components (one isotropic and two anisotropic) were reconstructed for each subject [13].

6.1 Validation

An alternative to the method presented in this paper would be to select one of the two tensors in each voxel (e.g. the one with the highest FA) and to perform single tensor registration on this image. Here, we validate that our method works better than this simple alternative. The quality of the alignment is assessed by the sum of square differences of each eigenvalue after alignment of control subjects. Indeed, while the diffusivities can significantly differ in diseased brain, they are approximately equal for healthy subjects. We performed 26 randomly chosen registrations with four levels of regularization, totalizing 104 registrations. In each voxel, the eigenvalues were averaged between the two anisotropic components (weighted by their fractions). Results show that multi-fiber registration performs significantly better than single tensor registration (Fig. 1).

6.2 Atlas Construction

An atlas was constructed using our registration and alternating three steps: aligning all subjects to the current atlas (initially a randomly chosen subject), averaging the aligned subjects (using the weighted combination of mixtures), applying the mean inverse field to the resulting average [14]. This atlas remarkably shows areas where multiple fibers are consistently present in all subjects (Fig. 2).

registration. We therefore believe that this registration method opens new doors to understanding brain disorders based on multi-fiber models.

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