

White matter characterization by massively multidimensional diffusion-correlation MRI



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Introduction

Massively multidimensional diffusion-correlation MRI uses free gradient waveforms to obtain nonparametric $D(\omega)$ - R_1 - R_2 -distributions^[1].

- Squared normalized anisotropy D_{Δ}^2
- Isotropic diffusivity **D**_{iso}
- Relaxation rates **R**₁ and **R**₂

To visualize the tissue fractions from the obtained $D(\omega)$ - R_1 - R_2 -distributions, a set of arbitrary "bins" limits are selected on the D_{Δ}^2 and D_{iso} plane.

To expand the within-tissue classification to not only in the 2D D_{Δ}^2 and D_{iso} plane, here we use the unsupervised k-means clustering method including the information of D_{Δ}^2 , D_{iso} , R_1 and R_2 to characterize the components in white matter.



Methods

- 5 fixed rat brains with 4% paraformaldehyde (PFA).
- Both optic nerves were removed from each brain

Data acquisition and processing

- Bruker Avance-III HD 11.7 T, MIC-5 probe 3 T/m
- Each scan consisted of 737 images using free gradient waveforms^[4] varying *b*-value, b_{Δ} , (θ , φ), $\omega_{cent}/2\pi$, τ_{R} , and τ_{E} . The total scan time was ~20 hours.
- Images were denoised^[5] and the Monte Carlo inversion algorithm was applied to obtain the nonparametric distributions^[6]
- The mean distributions $E[D_{\Delta}^{2}]$, $E[D_{iso}]$, $E[R_{1}]$ and $E[R_{2}]$ across the 10 optic nerves were calculated for k-means clustering







Acquisition scheme



nal



Results

In the 3D scatter plot can be observed that most part of the distribution is localized in high $E[D_{\Delta}^{2}]$, as expected for white matter.

The resulted clusters in white matter shows that group 1 has the biggest contribution from $E[D_{\Delta}^{2}]$, $E[R_{1}]$ and $E[R_{2}]$; group 2 shows a significant reduction of $E[D_{\Delta}^{2}]$, as well of $E[R_{1}]$ and $E[R_{2}]$; and cluster 3 has the smallest contribution of $E[D_{\Delta}^{2}]$ and almost no contribution of $E[R_{1}]$.

The cluster 1 is conformed by 88% percent of the data, cluster 2 by 9 % and cluster 3 by 3%. Each cluster might be related to the different elements in white matter.



3D scatter plot with the averaged distribution of $E[D_{\Delta}^2]$, $E[D_{iso}]$ and $E[R_1]$ (on the axes) and $E[R_2]$ (as the color-code).



Conclusion





Our results show the potential of massively multidimensional diffusion-relaxation correlation acquisitions to disentangle the per voxel microstructure information of white matter. Furthermore, the characterization of the clusters by diffusivities, anisotropy, and relaxation rates of white matter would allow a better understanding of the subtle changes during pathologies. We thank Maarit Pulkkinen for her assistance in animal and tissue handling.

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