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INTRODUCTION

The **free water elimination (FWE) diffusion model** accounts for partial volume effects that occur when voxels in diffusion tensor imaging (DTI) volumes contain both a tissue and a free water compartment^{1,2}. A downside of FWE is that the **model fitting problem is ill-conditioned**^{3,4}. Advanced parameter estimation techniques that incorporate regularization usually succeed in stabilizing the model fit but, as a trade-off, impose model assumptions that are likely to bias the results^{1,5-6}. In this work, **we exploit that the T_2 relaxation times of white matter and cerebrospinal fluid are very different**. By accounting for the associated echo time (T_E) dependency of the signal decay, the model parameters can be estimated more precisely, accurately, and robustly.

RESULTS

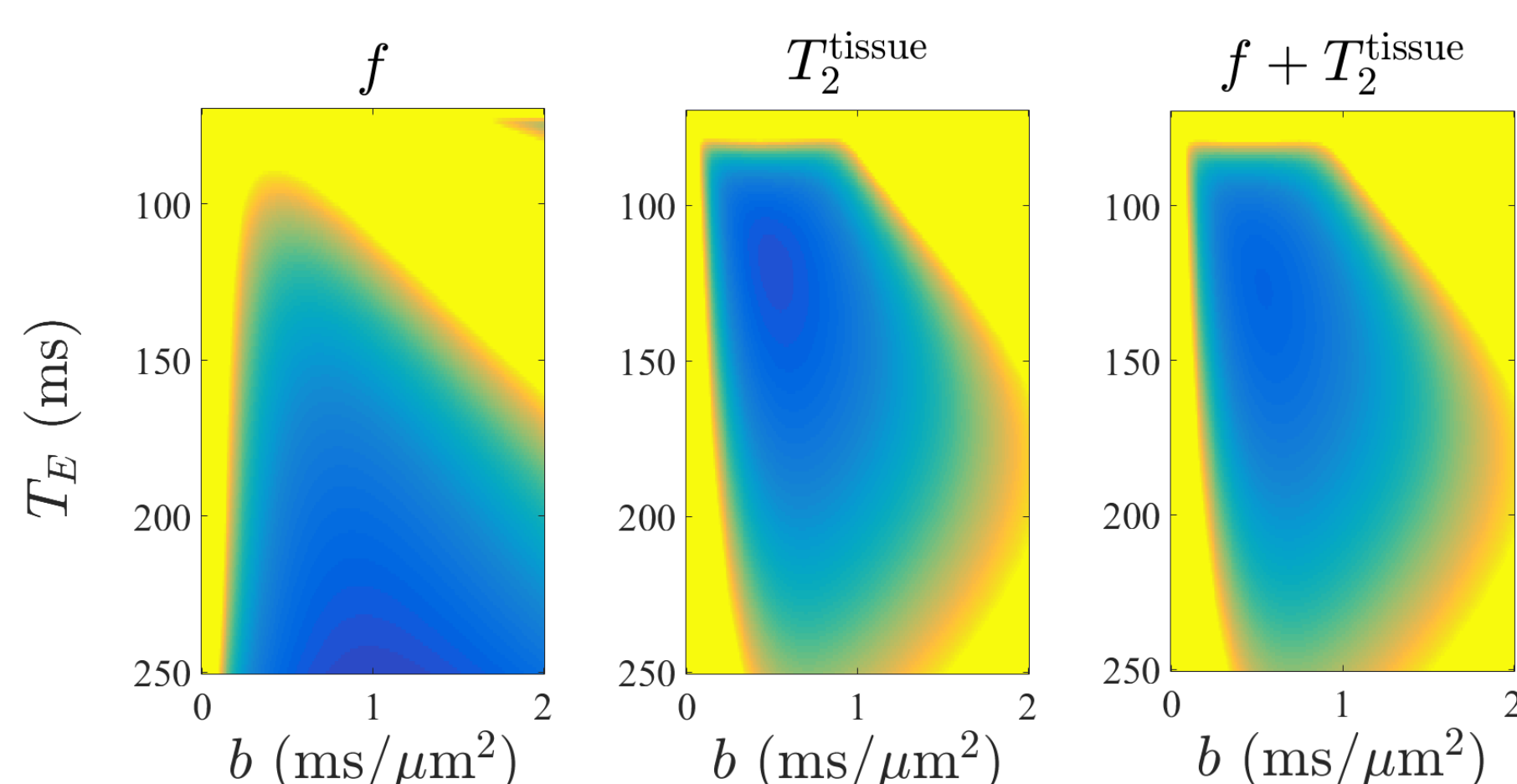


Figure 1: The **coefficient of variation** for the FWE- T_2 model with T_2^{fw} estimation, derived from the Cramér-Rao lower bound (assuming Gaussian noise), for f , T_2^{tissue} , and a combination of both. A typical WM voxel with free water was assumed. Protocol: $5 \times b = 0$, $30 \times b = 1$ with $T_E = 70\text{ms}$ and 1 extra shell with variable b and T_E (30 dirs).

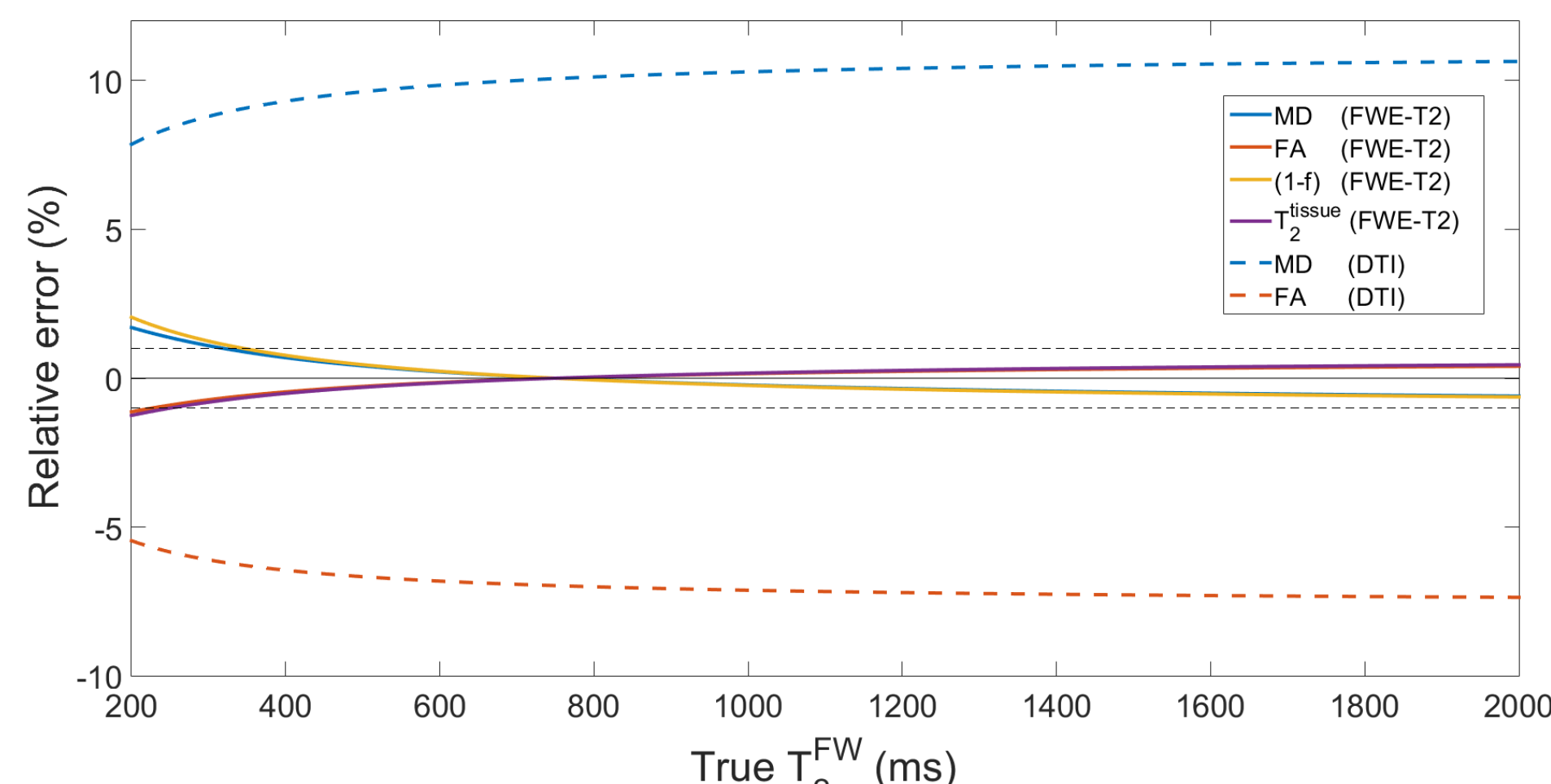


Figure 3: **Relative error (%)** of FWE- T_2 and DTI parameters in function of the true value of T_2^{fw} (FWE- T_2 model assumes a fixed $T_2^{fw} = 750\text{ms}$).

DISCUSSION AND CONCLUSION

In this work, we propose an extension to the FWE model that incorporates the T_E dependencies of both the tissue and free water compartments. We showed that in addition to a standard DTI acquisition protocol, ideally, **only a sparsely sampled additional shell should be acquired** with $b \approx 0.5 \text{ ms}/\mu\text{m}^2$ and $TE \approx 120\text{ms}$ (fig. 1). Monte Carlo simulations indicate that the FWE- T_2 model with a prior estimated T_2^{fw} , is substantially better compared to FWE or FWE- T_2 with estimation of T_2^{fw} , in terms of both the precision and accuracy of f and MD estimation. **Fixing the T_2 of CSF is justified** because the increase in precision (fig. 2: red versus yellow) outweighs the potential drop in accuracy for the FWE- T_2 model parameters, where the errors will typically be smaller than 1% (fig. 3). Finally, clinical data shows that **neglecting to account for free water partial volume effects, biases the estimation of diffusion properties** of brain tissue close to CSF regions.

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FWE- T_2 MODEL

$$S_i = S_0 \left(\underbrace{(1-f)e^{-T_E/T_2^{tissue}}}_{\text{Tissue}} \underbrace{e^{-b_i \mathbf{g}_i^T \mathbf{D} \mathbf{g}_i}}_{\text{Diffusion weighting}} + \underbrace{f e^{-T_E/T_2^{fw}}}_{\text{free water}} \underbrace{e^{-b_i d}}_{\text{Diffusion weighting}} \right)$$

S_0 = signal without diffusion weighting on $T_E = 0$; b_i , \mathbf{g}_i = diffusion weighting strength and direction; \mathbf{D} = diffusion tensor; d = diffusion of free water at body temperature ($= 3 \mu\text{m}^2/\text{ms}$)

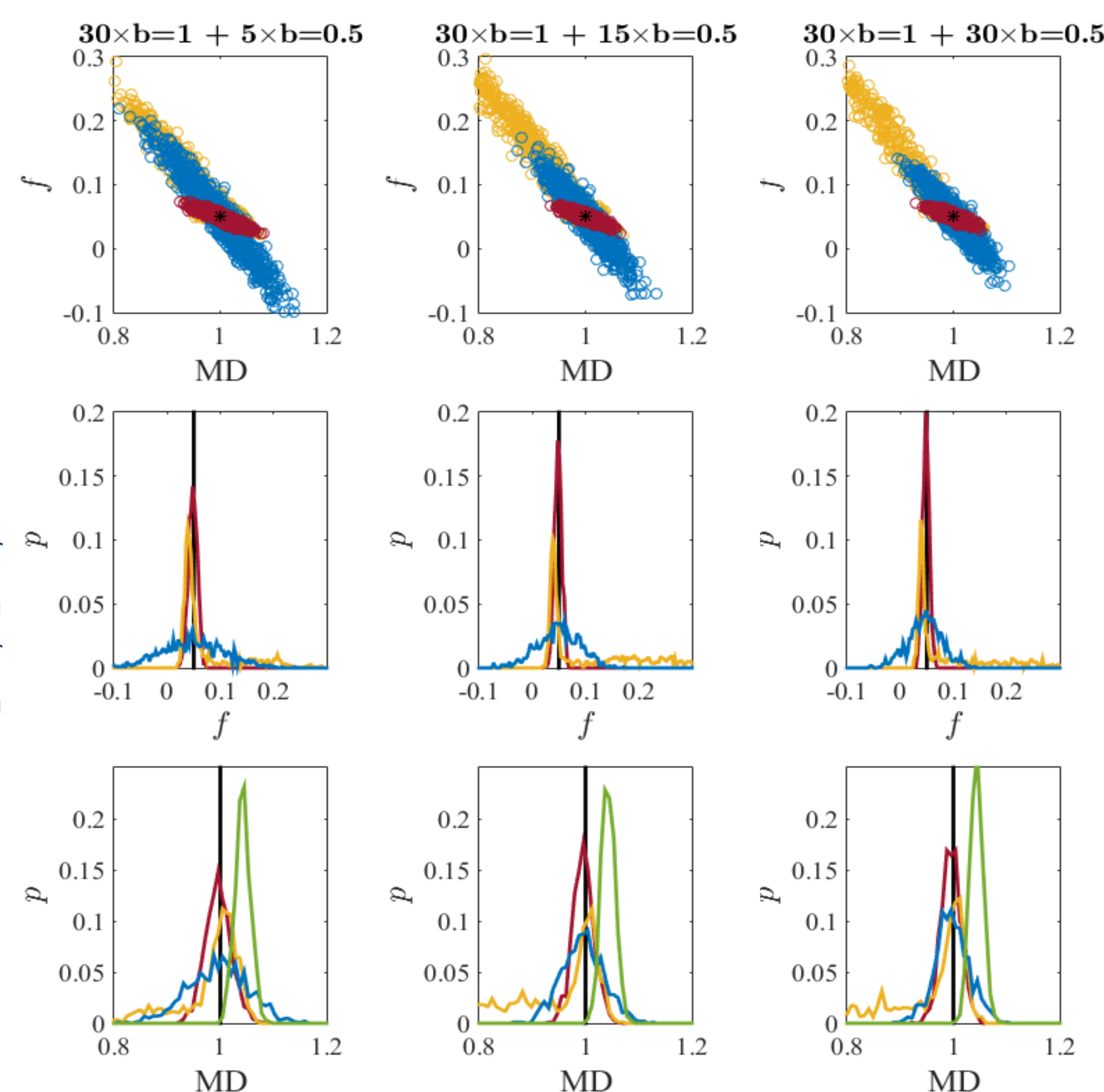


Figure 2: Single voxel **Monte Carlo simulations** assuming a typical WM voxel with a free water compartment and $\text{SNR}=50$ on $b = 0$ and $T_E = 100\text{ms}$ signal. Comparison between: FWE- T_2 w/o T_2^{fw} est., FWE- T_2 with T_2^{fw} est., FWE, DTI and true value. Protocol: $5 \times b = 0$ ($T_E = 70\text{ms}$), $30 \times b = 1$ ($T_E = 70\text{ms}$) and $5/15/30 \times b = 0.5$ ($T_E = 120\text{ms}$ or 70ms for DTI and FWE).

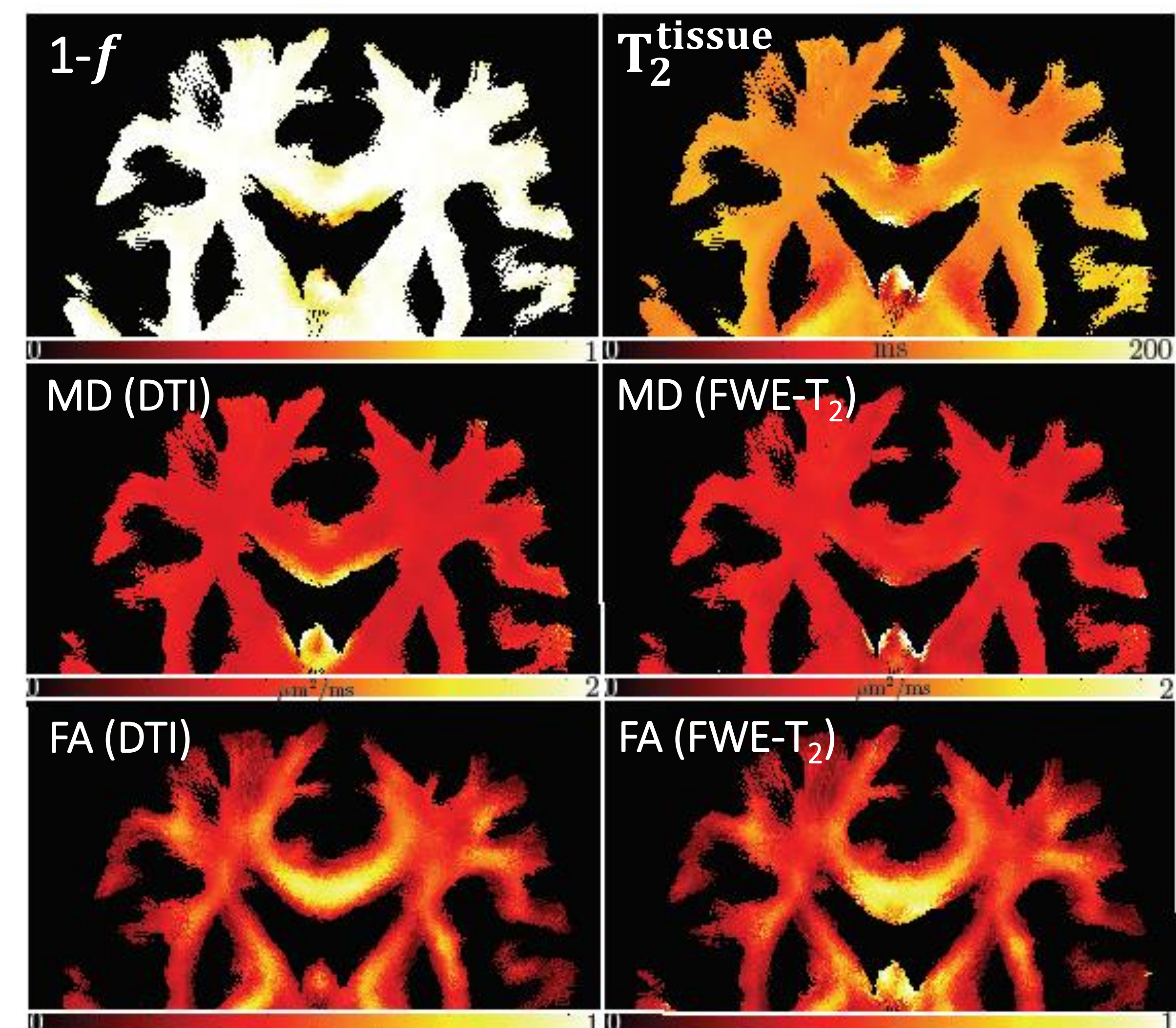


Figure 4: **Real data** results of FWE- T_2 (w/o estimation of free water T_2 value) and DTI. A healthy volunteer was scanned on a 3T scanner with $1 \times b = 0$, $30 \times b = 0.5$ ($T_E = 120\text{ms}$) and $30 \times b = 1$ ($T_E = 70\text{ms}$). The data was denoised⁷ and subsequently corrected for Gibbs ringing⁸, eddy current distortions and motion⁹.