

Motion Correction for 3D T₁ Mapping using GRICS: Phantom Validation

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INTRODUCTION

T₁ mapping is a valuable tool to analyze myocardial fibrosis or liver perfusion. For better characterization, 3D imaging is highly desirable. Unfortunately, these 3D acquisitions exceed tolerable breath hold time. Moreover, most T₁ mapping methods, such as the variable flip angles method, require several acquisitions with varying imaging parameters. But motion artifacts and misregistrations affect the quantification. In this work, we propose a T₁ mapping reconstruction method that includes motion correction. This method is a variant of T₂ GRICS proposed by Lohezic et al. [1], adapted to T₁ and 3D.

THEORY

T₁ weighting: The approximation of short flip angles (eq. 1) is used to describe the image weighting, where ρ_i stands for the T₁ weighted image, ρ₀ the image without T₁ weighting, and θ_i the current flip angle.

$$\rho_i = \rho_0 \frac{\sin \theta_i \left(1 - e^{-\frac{TR}{T_1}}\right)}{1 - e^{-\frac{TR}{T_1}} \cos \theta_i} \quad (1)$$

Motion Model: Each physiological motion is described by a non-rigid motion model α_k. Thus, global motion U can be described by a linear combination of the motion models weighted by the associated physiological sensors S_k (ECG, respiratory belt, etc.) (eq. 2).

$$U(x, t) = \sum_k \alpha_k(x) S_k(t) \quad (2)$$

GRICS algorithm: GRICS is an adaptive reconstruction method proposed by Odille et al. [2] that takes the physiological sensors into account in order to correct for motion that occurs during the acquisition. It is based on the inversion of coupled linear systems that describes the imaging system (fig. 1). If the motion model is known, weighted images ρ_i can be obtained (eq. 3). Then, the fitting of equation 1 gives us the T₁ map. If T₁ weighting is also simulated in the imaging system, we obtain the morphological image ρ₀ shared by all the acquisitions (eq. 4). Since the motion model is shared between all acquisitions, misregistration problems are also avoided. The motion model is updated by solving for a linearized expression of the residual error with respect to the error in motion model parameters δα_k (eq. 5). A fixed point multiresolution scheme is used to invert the system of equations formed by equations 3, 4 and 5.

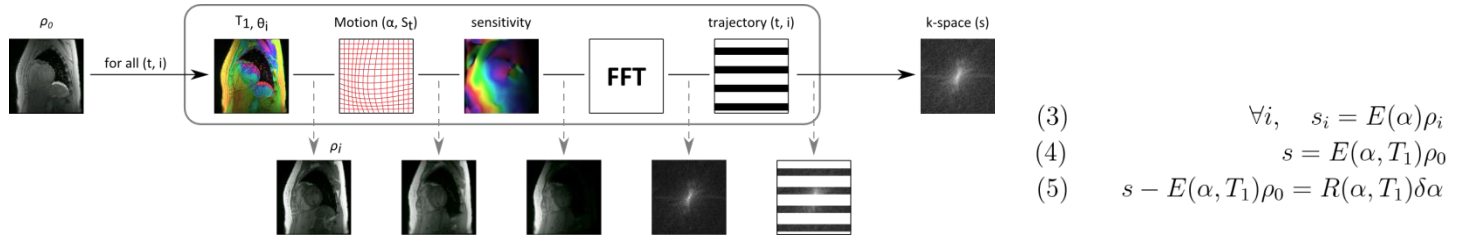


Figure 1 - The imaging system is made up of a weighting, a distortion, a sensitivity model, a Fourier transformation and a trajectory mask.

MATERIALS AND METHODS

Acquisition: At 3T (SIGNA HDxt, GE Healthcare, Milwaukee, WI), two data sets were acquired on a homemade phantom made up of eight tubes of gadolinium dissolution (0 to 0.875 mmol.L⁻¹) bathing in water. 3D FSPGR sequences (TE=3.3ms, TR=6.7ms, FOV=16x16x10cm) with variable flip angles (2, 5 and 11 , each repeated three times) were used. With a static phantom, the first data set was acquired and excitation field (B1) was reconstructed from two EPI acquisitions (same parameters with FA1/FA2=60/120  and 120/240 ). Next, the phantom was animated thanks to a breathing apparatus translating a platform in plane and through plane. Then, the second data set was acquired while signal from a respiratory belt was collected by a custom Maglife patient monitoring system (Schiller Medical, France) and recorded with a dedicated home-made hardware presented in [3].

Data analysis: Three T₁ maps were generated. Two maps were obtained with linear fitting on the two sets of data (with and without motion) and the third one with the new 3D T₁ GRICS reconstruction with the moving phantom. A B₁ correction was applied to each map following the method proposed by Cheng et al. [4]. Results were presented as the mean ± standard deviation of T₁ values in ROIs encompassing the tubes.

RESULTS

T₁ values obtained with GRICS correction show significant differences (p<0.001) compared to those with uncorrected motion (fig. 2) and are in good agreement with those without motion. Uncorrected T₁ values are always overestimated.

DISCUSSION & CONCLUSION

High T₁ values obtained with uncorrected motion show that T₁ values can be corrupted by surrounding T₁ values due to motion artifacts. In addition, since T₁ values obtained without motion can be considered as the reference, GRICS significantly improves the accuracy of T₁ for acquisitions taken while motion occurs. This method can be applied to correct breathing artifacts in myocardial T₁ mapping, or liver and breast perfusion.

REFERENCES

- [1] Lohezic et al., Proc. ISMRM 2010, 2958 ; [2] Odille et al., MRM 60:146–157 (2008); [3] Odille et al., IEEE TBME 54: 630-640 (2007); [4] Cheng et al., MRM 55:566–574 (2006)

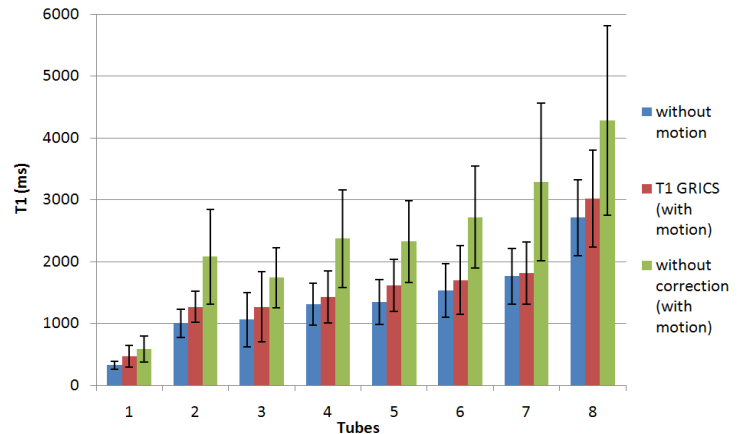


Figure 2 - Comparison of T₁ maps obtained without motion and with motion, with and without GRICS correction