

## Intrinsic motion correction for radial cardiac T<sub>2</sub> mapping through alternating T<sub>2</sub> preparation duration

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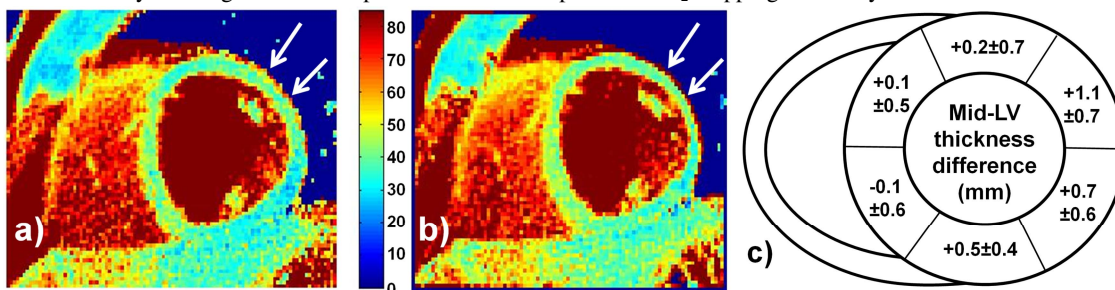
**Introduction:** T<sub>2</sub> mapping through variation of the T<sub>2</sub> preparation (T<sub>2</sub>Prep) duration has been increasingly used to detect and quantify cardiac edema in response to myocardial injury<sup>1</sup>. However, if images with incremental T<sub>2</sub>Prep duration are acquired in a sequential fashion (Fig.1a), irregular breathing patterns and heart rates may adversely affect the quality of the T<sub>2</sub> maps due to misalignment of the source images. A logical alternative is then to acquire all images in an alternating manner (Fig.1b), where the T<sub>2</sub>Prep duration changes cyclically from one heartbeat to the next. Combined with a radial signal readout, this may minimize the vulnerability to respiratory or RR variability. We therefore simulated, implemented and tested the use of an alternating magnetization preparation approach to T<sub>2</sub> mapping.

**Methods:** A navigator-gated ECG-triggered radial gradient-recalled-echo pulse sequence (20 lines per heartbeat, ECG trigger every 3 heartbeats) was implemented to obtain source images for the T<sub>2</sub> maps<sup>2</sup>, with the possibility to apply the T<sub>2</sub>Prep durations of 60/30/0ms in both an alternating and sequential manner. The sequential T<sub>2</sub>Prep source images were co registered before T<sub>2</sub> fitting<sup>3</sup>, while the alternating T<sub>2</sub>Prep images were not. Bloch equation simulations were performed in order to estimate the longitudinal magnetization residual due to T<sub>1</sub> relaxation<sup>2</sup>, as well as the accuracy over a range of heart rates. The sequences were validated in agar-NiCl<sub>2</sub> phantoms at 3T (12-channel surface coil array, on a Magnetom Trio, Siemens, Erlangen, Germany) by comparing the resulting T<sub>2</sub> maps to gold-standard spin-echo (SE) T<sub>2</sub> maps. A mid-ventricular short-axis T<sub>2</sub> map was then acquired with both (alternating and cyclical T<sub>2</sub>Prep) pulse sequences in 9 healthy adult volunteers. The total myocardial surface area and AHA-standard<sup>4</sup> segmental left ventricular (LV) wall thickness were measured in the T<sub>2</sub> maps, after which a paired Student's t-test was applied to detect differences.

**Results:** The Bloch equation simulations demonstrated that the T<sub>2</sub> value in the alternating method was most accurately fitted with a longitudinal magnetization residual of 0.13 and that it was as robust to heart rate variation as its sequential counterpart: ~3.4ms vs. ~2.4ms variation in fitted T<sub>2</sub> value between 40 and 90bpm for the alternating and sequential methods, respectively. Its accuracy was confirmed in the phantoms: T<sub>2</sub>=45.4±0.7ms for the alternating method vs. 45.3±0.7ms for the sequential method and 45.1±0.7ms for the spin-echo gold standard. The myocardial surface area was larger in the alternated T<sub>2</sub> maps of the volunteers (8.4±1.8cm<sup>2</sup> vs. 7.5±1.8cm<sup>2</sup>, p<0.001) (Fig.2), while the average midventricular T<sub>2</sub> value slightly differed between the alternated and sequential methods (T<sub>2</sub>=36.5±2.2ms alternated vs. 39.1±2.7ms sequential, p<0.001). The LV wall thickness measurements demonstrated that when the alternating method was used, the lateral segments had a higher thickness increase than the septal segments (Fig. 2c). The average thickness increase when comparing the alternating to the sequential method was 12±13% p<0.01.

**Discussion:** The alternating method demonstrated a larger LV surface area and LV wall thickness than the sequential method, while their T<sub>2</sub> fitting robustness was similar. The larger LV wall surface area and thickness in the alternating T<sub>2</sub> maps may be explained by the intrinsic source image alignment of this method: the source images obtained from the alternating method are less subject to transient changes of the end-expiratory position or RR changes during the scan. In contrast, and for a sequential acquisition, such transient changes inevitably lead to misalignment of the source images and ultimately also a smaller number of pixels that are available for analysis.

**Conclusions:** We successfully implemented and tested a T<sub>2</sub> mapping methodology in which the T<sub>2</sub> preparation is alternated. The *in vivo* T<sub>2</sub> maps demonstrate that this alternating method results in a better registration of the source images, which in turn results in a larger myocardial thickness and the availability of a larger number of pixels that can be exploited for T<sub>2</sub> mapping. This may allow for more accurate T<sub>2</sub> quantification.



**Figure 2 – a) and b)** T<sub>2</sub> map of volunteer acquired with the alternating (a) and sequential (b) method. Note that consistent with the quantitative findings, the antero-lateral myocardium is thicker when acquired with the alternating method (arrows). **c)** Myocardial LV thickness increase (in mm) when using the alternating method.

**References:** 1. Giri et al., J Cardio Magn Reson 2009;11:56, 2. van Heeswijk et al., JACC Cardio Imag 2012; 5(12):1231, 3. Giri et al., Magn Reson Med 2012; 68(5):1570, 4. Cerqueira et al.; Circulation 2002;105:539