

## T<sub>2</sub> and T<sub>2</sub>\* Relaxometry in the Meniscus using a Novel, Rapid Multi-Echo Steady State Sequence

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**Introduction:** Imaging the meniscus is challenging due to its short T<sub>2</sub> relaxation time and highly organized collagen architecture. The meniscus plays an important role in osteoarthritis and it is becoming increasingly clear that detecting early degeneration, before gross morphological changes occur, is essential for evaluating disease progression and treatments. T<sub>2</sub> and T<sub>2</sub>\* relaxation time can be used to detect early meniscal degeneration<sup>1,2</sup>; however, lengthy scans are required (10 to 20 minutes per measure). In this work we show that a novel multiecho steady-state sequence can be used to estimate T<sub>2</sub> and T<sub>2</sub>\* relaxation times in the meniscus *simultaneously in under 5 minutes*.

**Methods:** A quantitative DESS sequence<sup>3,4</sup> was modified to include multiple gradient echo readouts (“MEDESS”, Figure 1). In this sequence, the S<sup>+</sup> signal has the usual T<sub>2</sub>\* decay; however, the S<sup>-</sup> signal has T<sub>2</sub> decay with rephasing, similar to gradient-and-spin echo methods<sup>5,6</sup>. In TR<sub>1</sub>, we have included two S<sup>+</sup> and two S<sup>-</sup> signals. This sequence is then repeated with an offset of ΔTE in TR<sub>2</sub>. Interleaving TR<sub>1</sub> and TR<sub>2</sub>, we have four S<sup>+</sup> and four S<sup>-</sup> signals within one acquisition. T<sub>2</sub>\* relaxation time can be estimated by fitting a monoexponential decay curve to the four S<sup>+</sup> signals. T<sub>2</sub> relaxation time can be estimated using S<sup>+</sup> and S<sup>-</sup> signal pairs (ie:1 & 8, 2 & 7) and a signal model  $(T_2 = (TE_{S+,1} - (TR + TE_{S-,8})) / \log(S_{-,8} / S_{+,1}))^4$ . This signal model does not consider the mixed T<sub>1</sub>/T<sub>2</sub> contrast present in the S<sup>+</sup> signal which causes an underestimation of T<sub>2</sub>. Extended phase graph (EPG) simulations can be used to correct the estimate (assuming T<sub>1</sub> ≈ 800 ms, based on previous laboratory measurements).

We compared estimates of T<sub>2</sub>, corrected T<sub>2</sub> and T<sub>2</sub>\* relaxation times obtained using the MEDESS sequence to standard measures in four cadaver knee specimens. All scans were acquired in the sagittal plane (matrix: 256 x 256, field of view: 20 cm, slice thickness: 3 mm). 2D spin echo (SE) was used as the standard measure of T<sub>2</sub> (eight scans, TE: 10 to 24 ms, TR: 1 s, total time: 35 minutes). A 3D multi-echo GRE (MEGRE) was the standard measure of T<sub>2</sub>\* (eight echoes, TE: 2 to 20 ms, TR: 100 ms, FA: 30°, time: 19 minutes). Monoexponential curve fits were used for both standard measures. Differences between MEDESS and standard measures were described as the root mean square (RMS) absolute difference for a single slice.

**Results:** T<sub>2</sub> and T<sub>2</sub>\* relaxation times estimated using MEDESS were similar to standard measures (Figures 2 and 3); the RMS absolute difference was 3.5, 2.9 and 2.0 ms, for T<sub>2</sub>, corrected T<sub>2</sub> and T<sub>2</sub>\*, respectively. MEDESS consistently underestimated the T<sub>2</sub> and T<sub>2</sub>\* relaxation times (Figure 2); this was true for mean differences and pixel-wise differences (Figure 3).

**Discussion:** The EPG-based correction provided a modest improvement to the T<sub>2</sub> estimate. However, since the variation within the maps was consistent between the methods, the observed bias is not of great concern; it will still be possible to detect changes over time and between patient groups. This novel sequence, can be used for estimation of meniscal T<sub>2</sub> and T<sub>2</sub>\* relaxation times in under 5 minutes, about a 75% time reduction compared to acquiring both of these measures with other techniques.

**References:** 1. Rauscher et al, Radiology, 2008; 2) Williams et al, Osteoarthritis Cartilage, 2012; 3) Staroswiecki et al, MRM, 2012; 4) Welsch et al, MRM, 2009; 5) Ma et al. JMR, 1996; 6) Yablonskiy et al. MRM, 1997.

**Acknowledgements:** NIH R01-EB 002524, NSERC PDF, GE Healthcare.

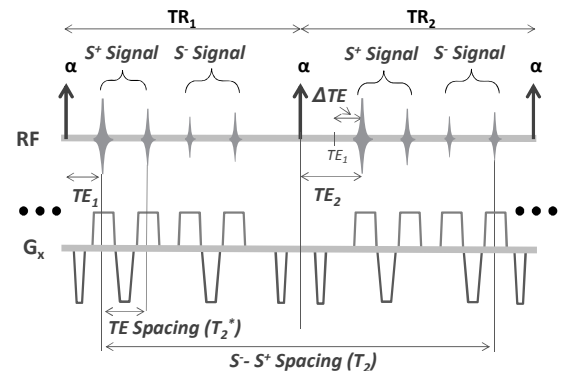


Figure 1: Multi-echo Steady State Sequence. Parameters include: TR = 12 ms, S<sup>+</sup> TE = 2.0, 3.2, 4.3, 5.4 ms, S<sup>-</sup> TE = 6.5, 7.7, 8.8 and 9.9 ms, FA = 30°, time = 4.5 minutes. (TE relative to α pulse, echo spacing = 2 \* ΔTE)

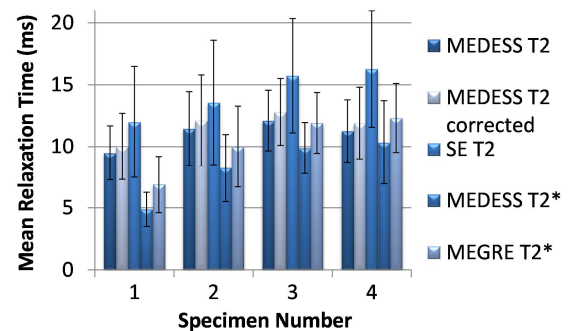


Figure 2: Multi-echo steady state (MEDESS) vs. standard measures of T<sub>2</sub> and T<sub>2</sub>\* relaxation times.

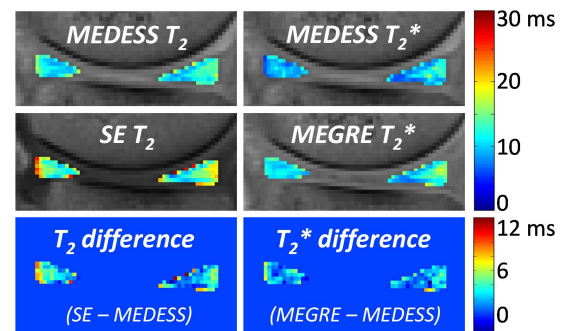


Figure 3: MEDESS consistently agreed with standard measures. Using specimen 3 as an example, the pixel-wise differences were 3.5±2.9, 2.7±2.7 and 2.1±2.0 ms, for T<sub>2</sub>, corrected T<sub>2</sub> and T<sub>2</sub>\* relaxation times, respectively, similar to the RMS absolute difference. \*note different scales