

Interleaved T1- and T2*-weighted Imaging Can Evaluate Dynamic Oxygen Challenge: A Feasibility Study

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INTRODUCTION

Dynamic oxygen challenge evaluated by nuclear magnetic resonance T₁ and T₂* relaxation times (DOCENT) is a robust non-invasive prognostic test that reveals tumor hypoxia (1,2). It accomplishes this by exploiting tissue oxygenation level dependent (TOLD) and blood oxygenation level dependent (BOLD) MRI contrast techniques that assess tumor oxygenation in vivo. Currently, no effective simultaneous T₁- and T₂*-weighted dynamic acquisition method has been developed for clinical applications at 3 Tesla field strength. It has been shown previously (3) that a two point 3D modified fast inversion recovery method can provide T₁- and T₂-weighted images simultaneously in a short scan time. However, the technique lacks acquisition of T₂* information, which is essential for the study of the BOLD effect. An interleaved T₁ and T₂* approach was presented by Winkelman, et al in brain studies at 1.5T, although this apparently did not work well for abdominal exams performed at higher field-strength. Therefore, a fast 2D respiratory-triggered interleaved T₁- and T₂*-weighted sequence, capable of monitoring BOLD and TOLD effects simultaneously, has been developed and tested on the human spleen and kidney.

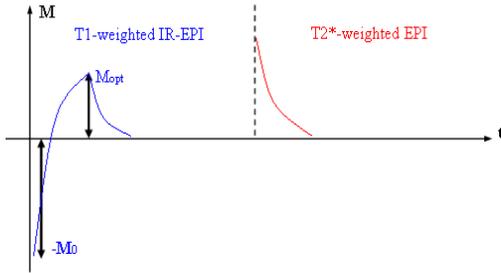


Fig. 1. Schematic plot - Interleaved T₁W and T₂*W sequence.

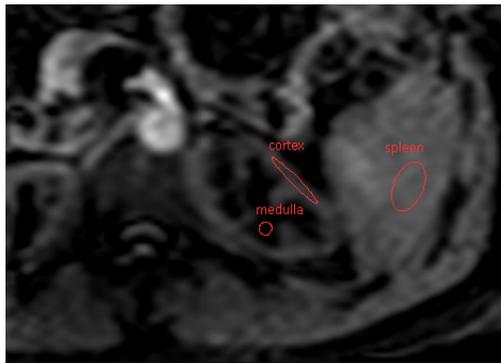


Fig. 2. T₁-weighted image with ROIs

DISCUSSION AND CONCLUSION

This study has shown that the present 2D respiratory-triggered interleaved T₁- and T₂*-weighted sequence provides a promising means to study TOLD and BOLD response simultaneously without the need for registration and with good temporal resolution (less than 30 seconds). Our observation of TOLD (but no BOLD) response in spleen, confirms that reported previously by O'Connor et al at 1.5T (6). This method also has the potential to calculate T₁ and T₂* maps with multi-echo acquisition applied (3).

REFERENCES

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Fig 4.(right) Changes in relative signal in response to 100% oxygen breathing in TOLD for spleen and kidney.

METHODS

Imaging has been performed on a clinical 3.0T scanner (Achieva, Philips Medical Systems) using a SENSE cardiac coil (6-element). A 2D T₁-weighted inversion recovery echo planar imaging (IR-EPI) experiment has been fused together (interleaved) with a T₂*-weighted EPI sequence (Fig. 1). For the T₁-weighted imaging, the optimum TI was set at 1400ms with TE = 5.6ms (4). The TE value of the T₂*-weighted portion was equal to 26ms. All other parameters for both sequences were identical (TR ~ 2 respiratory cycle, FOV=25cm, matrix size=128×128, slice thickness=5 mm, partial Fourier acquisition factor=0.625, SENSE factor=2, multi-shot, SPIR fat suppression). The FOV was selected to include spleen, left kidney, and paraspinal muscle (Fig. 2). Measurements were acquired during gentle breathing with respiratory triggering. Total scan time for each T₁-weighted and T₂*-weighted images was around 24s. Ten baseline measurements were collected while breathing room air, followed by 20 measurements during inhalation of 100% oxygen. The first *in vivo* experiments were conducted in 3 volunteers.

RESULTS

Three regions of interest (ROI) were drawn by Y.D. (Fig. 2), and dynamic BOLD (Fig.3) and TOLD (Fig. 4) effects with ~24 sec temporal resolution were generated from interleaved T₂*- and T₁-weighted dynamic images. Both spleen and renal cortex showed large increases in T₁-weighted signal intensity (mean maximum Rel. Signal Change (%) = 16.9 ± 4.8 and 10.8 ± 3.0, respectively), whereas renal medulla showed little TOLD response (mean maximum Rel. Signal Change (%) = 0.7 ± 6.6). Conversely, only renal medulla showed a large BOLD effect (mean maximum Rel. Signal Change (%) = 14.3 ± 3.9). A different pattern of BOLD response was observed in spleen and renal cortex, minimal in both (mean maximum Rel. Signal Change (%) = 0.3 ± 1.7 and 1.2 ± 3.1, respectively)

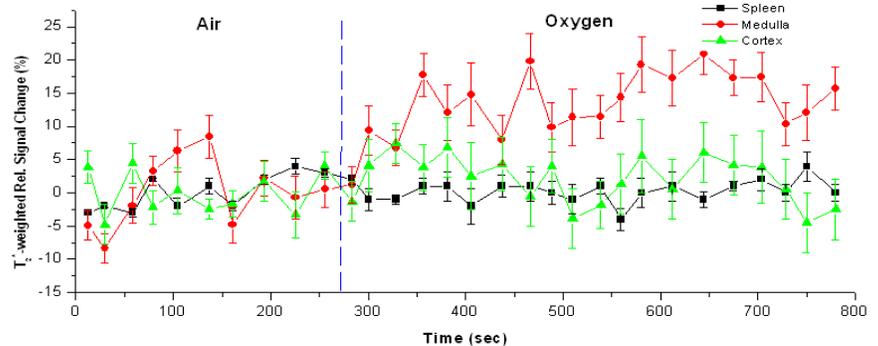


Fig 3. Changes in relative signal in response to 100% oxygen breathing in BOLD for spleen and kidney.

