

POSTER PRESENTATION

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T₂-dependent errors in MOLLI T₁ values: simulations, phantoms, and in-vivo studies

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Background

Diffuse myocardial fibrosis occurs in various cardiomyopathies and can be indirectly assessed with blood and myocardial T₁ mapping at baseline and after gadolinium administration. The widely used MODified Look-Locker Inversion-recovery (MOLLI) [1] sequence is known to underestimate myocardial T₁ at higher heart rates, but its dependence on T₂ has not been explored. We investigate MOLLI's T₁ accuracy in phantoms and confirm with simulations and in-vivo studies. T₁ values are further compared with a saturation-recovery T₁ mapping sequence [2].

Methods

Phantoms

14 NiCl₂ agarose phantoms with a broad range of T₁ and T₂ values were imaged with a gold-standard inversion-recovery spin-echo (IR-SE) sequence, MOLLI, and a new SATuration-recovery single-SHOT Acquisition (SASHA) technique (Siemens Avanto 1.5T). **IR-SE:** 16 TIs 100-5000ms, TE=11ms, TR>5s, 90° flip. **MOLLI:** 2 inversion sets of 3 and 5 images, 75% partial Fourier, T_{1min}=110ms with 80ms increment, 35° flip, TE/TR=1.3/2.9ms, simulated HR=60bpm. **SASHA:** single-shot SSFP images from 10 consecutive heartbeats with incremented TI spanning the RR interval in the last 9 images (no saturation in the first image), 70° flip, TE/TR=1.3/2.6ms, full k-space, simulated HR=60bpm. T₂: spin-echo (SE) with 7 TEs 11-200ms. Simulations: Bloch equation simulations of MOLLI and SASHA were performed in MATLAB using actual acquisition and physiology parameters and SE measured T₁ and T₂ values.

In-Vivo

For 10 healthy volunteers (5 male, 28.8±6.6yrs), blood and myocardial T₁s were measured using MOLLI and SASHA (parameters as above) in a mid-ventricular short-axis slice at baseline and 20 minutes following 0.1mmol/kg Magnevist.

Results

In blood-like phantoms with long T₂ (179-196ms), SASHA and MOLLI T₁s agree well with IR-SE (0.7±0.5% and 2.2±1.8% absolute difference respectively), while shorter T₂ (46-76ms) tissue-like phantoms have greater underestimation with MOLLI (8.4±3.5%) than SASHA (0.9±0.6%) (Fig. 1). MOLLI simulations predict underestimated T₁s, with 1.3±0.9% absolute difference from observed values (vertical lines, Fig. 1). SASHA simulations also agree well with observations (0.8±0.5%, not shown). In healthy volunteers (63.3±8.4bpm), MOLLI T₁s also show greater underestimation compared to SASHA in tissue than blood, although the difference is larger than observed in phantoms or predicted by simulations in all cases (Table 1).

Conclusions

MOLLI significantly underestimates T₁s in shorter T₂ tissue-like phantoms but less so in longer T₂ blood-like phantoms, as predicted by simulations. Similar trends were observed in-vivo with MOLLI, although with greater T₁ underestimation (compared to SASHA) than predicted. SASHA had good agreement with IR-SE T₁ phantom measurements and simulations and can be acquired in less time than MOLLI.

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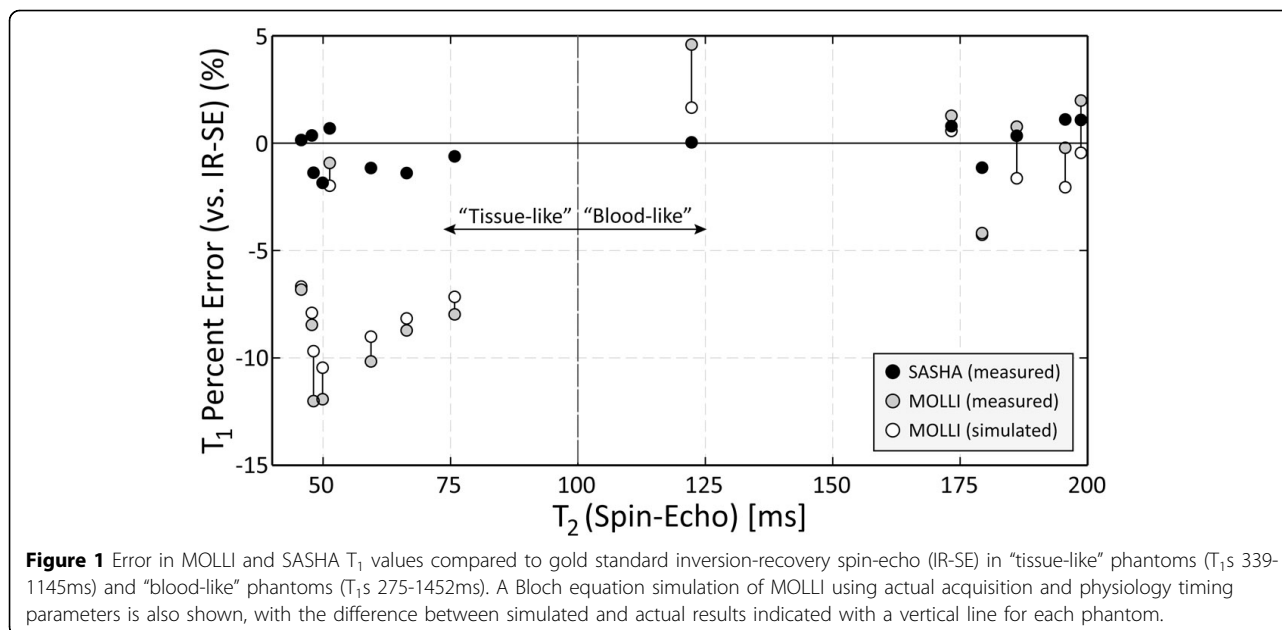


Table 1 Comparison of MOLLIs and SASHA T_1 values in 10 healthy volunteers prior to and 20 minutes following 0.1 mmol/kg Magnevist. All comparisons between MOLLIs and SASHA are significant ($p < 0.01$, two-tailed, paired Student’s t-test).

T_1 [ms]	Myocardium (mean±std)		Blood (mean±std)	
	Baseline	Post Gd (20 min)	Baseline	Post Gd (20 min)
MOLLIs	935.5±24.9	614.4±33.8	1514.1 ±107.5	524.9±55.2
SASHA	1175.2 ±27.6	752.9±48.2	1687.4±85.8	542.6±56.3

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