

POSTER PRESENTATION

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Characterization of myocardial T_1 and partition coefficient as a function of time after gadolinium delivery in healthy subjects

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Background

Diffuse myocardial fibrosis is associated with myocardial infarction [1], heart failure [2] and dilated cardiomyopathy [3]. Conventional T_1 -weighted late gadolinium enhancement (LGE) imaging highlights focal scarring in contrast to remote reference tissue, but it cannot detect global changes in T_1 associated with diffuse fibrosis. Quantitative T_1 imaging permits assessment of diffuse fibrosis by eliminating the use of reference tissue, but the dependence of the derived partition coefficient (lambda) on the time post-contrast injection (t_{post}) is not well established.

Purpose

Determine blood and myocardial T_1 values as a function of t_{post} and the resulting dependence of the blood-tissue partition coefficient.

Methods

Nine healthy subjects (22.0 ± 5.5 yrs, 6 male) were imaged using a Siemens Avanto 1.5T MRI. T_1 mapping was performed on a mid-ventricular short-axis slice using a custom saturation recovery single-shot TrueFISP sequence at baseline and one-minute intervals for 15 minutes following a bolus injection of gadopentetate dimeglumine (0.1 mmol/kg). At each time point, one "no-saturation" image and nine images with varying saturation recovery times spanning the cardiac cycle were acquired during a single breath-hold.

The myocardium was divided into 18 segments and mean values were fitted to a 3 parameter saturation

recovery curve to determine T_1 values for each segment at every time point. Blood T_1 values were computed using a region of interest within the left ventricular cavity. Lambda was computed using $\{\lambda = [R_1(\text{myocardium}_{\text{post}}) - R_1(\text{myocardium}_{\text{pre}})] / [R_1(\text{blood}_{\text{post}}) - R_1(\text{blood}_{\text{pre}})]\}$, where $R_1 = 1/T_1$.

Results

Figure 1 shows myocardial T_1 , blood T_1 , and lambda values averaged over all segments and subjects as a function of t_{post} . Average within-subject standard deviations of T_1 and lambda for t_{post} from 3-15 min were 34.1 ms and 0.046 respectively. Linear regression for lambda and t_{post} (3-15 min) shows an increase in lambda of 0.001 min^{-1} ($R^2 = 0.75$). Quantitative T_1 imaging is likely to be added to a clinical protocol following LGE imaging (t_{post} 10-15 min), where T_1 values increase by $5.9 \pm 1.6\%$ and lambda increase by $1.1 \pm 2.7\%$.

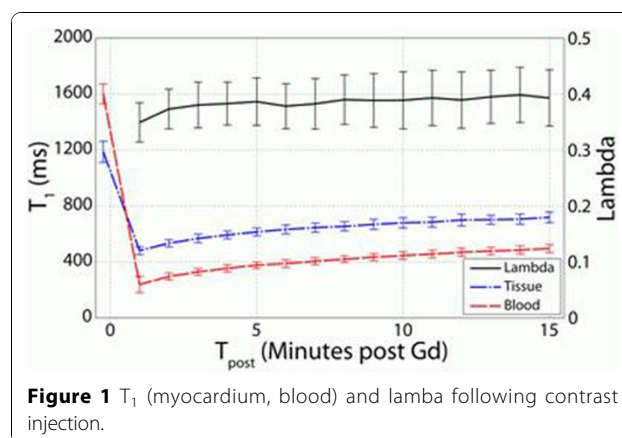


Figure 1 T_1 (myocardium, blood) and lambda following contrast injection.

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Conclusion

Saturation recovery SSFP T_1 mapping can be performed in a single breath-hold with derived blood-tissue partition coefficient (λ) values in good agreement with previous measurements³. In the post-LGE window of 10-15 min after contrast bolus, derived λ values show less time dependence than myocardial T_1 .

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