

POSTER PRESENTATION

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Native T₁ mapping identifies subclinical left ventricular myocardial t1 abnormality in patients with atrial fibrillation referred for pulmonary vein isolation

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Background

Native T₁ mapping has emerged as a noninvasive magnetic resonance imaging (MRI) method to assess the diffuse myocardial fibrosis without an exogenous contrast agents. To date, no data are available regarding native left ventricular (LV) T₁ relaxation time in AF patients referred for pulmonary vein isolation (PVI). The purpose of this study was to compare native myocardial T₁ value between AF patients undergoing PVI and control subjects.

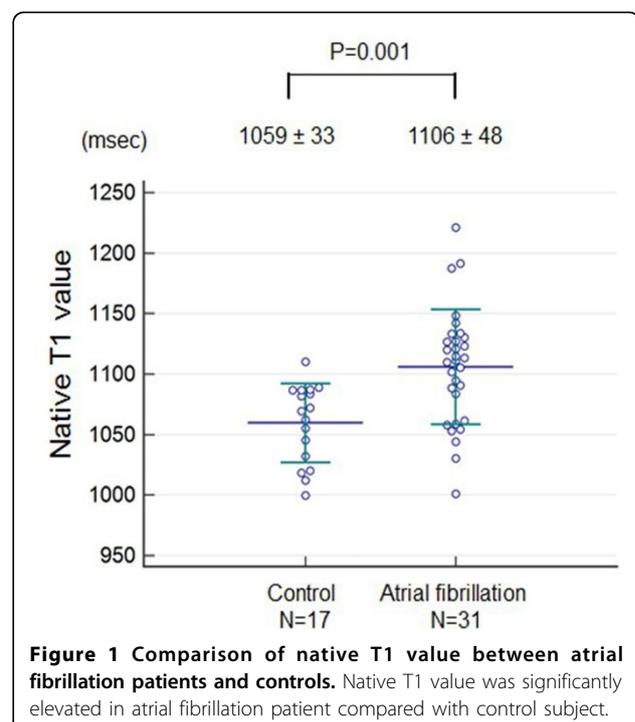
Methods

Thirty one AF patients referred for PVI (58±8 yrs; 23 m) and 17 control subjects (50±15 yrs; 13 m) were retrospectively identified. All patients were in sinus rhythm during the CMR scan. No AF patient had a history of myocardial infarction but 7 of 31 (23%) AF patients had history of hypertension. Non-contrast T₁ mapping images were acquired using a Modified Look-Locker imaging sequence (MOLLI) in 3 short-axis planes (basal, mid and apical slices) using an ECG-triggered single-shot acquisition with a balanced SSFP readout (TR 3.1; TE 1.5; FA 35°; FOV 360×337 mm²; acquisition matrix 188×135; voxel size 1.9 × 2.5 mm²; slice thickness 8 mm). Late gadolinium enhanced (LGE) MRI was acquired to evaluate for myocardial scar.

Results

LV ejection fraction was similar between groups (AF: 60 ±7%, controls: 58±7%, p=0.27). No LGE myocardial scar

was observed in any AF patient or control subject. Myocardial native T₁ was significantly elevated in AF patients (1106±48 msec vs 1059±33 msec, p=0.001). Multivariate linear regression analysis selected presence of AF as an independent and significant predictor of elevated native T₁ after adjustment for age and gender (OR: 25.8, 95%CI: 2.9 - 228.3, p=0.003).



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Conclusions

Our study suggests that there are subclinical differences (vs. normal controls) in native myocardial T_1 between AF patients referred for PVI. These data suggests that T_1 mapping may be useful for early detection of subclinical LV myocardial abnormalities in the AF population.

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