

# **POSTER PRESENTATION**

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# The native $T_1$ in remote myocardium of patients with prior chronic infarction is not normal

Steven Bellm\*, Shingo Kato, Ravi V Shah, Sophie Berg, Kraig V Kissinger, Beth Goddu, Long Ngo, Warren J Manning, Reza Nezafat

From 19th Annual SCMR Scientific Sessions Los Angeles, CA, USA. 27-30 January 2016

#### **Background**

Global left ventricular (LV) remodeling after myocardial infarction frequently occurs. Late Gadolinium Enhancement (LGE) CMR allows imaging of focal myocardial scar with areas remote from scar having no hyperenhancement. Myocardial  $T_1$  mapping allows quantification of interstitial fibrosis and may be a surrogate for LV remodeling. We sought to determine if there were  $T_1$  abnormalities in remote regions (no LGE positive areas) in patients with prior myocardial infarction.

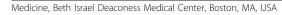
## Methods

In a prospective IRB-approved study, 12 patients with a history of coronary artery disease (CAD) and chronic

myocardial infarction (61  $\pm$  9 years, 9 males) and 10 healthy subjects (52  $\pm$  10 years, 8 males) were recruited to undergo CMR scans. All subjects were in sinus rhythm during CMR study. We assessed native  $T_1$  mapping using the slice interleaved  $T_1$  sequence in 5 short axis-slices (from apical to basal). The sequence was acquired in a free-breathing ECG-triggered slice-selective bSSFP.  $T_1$  mapping of each scan was estimated by voxel-wise curve fitting using a 2-paramter fit model. All images were corrected for in-plane motion between different  $T_1$  weighted scans. Native myocardial  $T_1$  in healthy subjects were measured over the three mid-ventricular slices by manually drawing epicardial and endocardial contours. The native  $T_1$  times of the remote myocardium of the CAD patients

Table 1 Subject characteristics of CAD patients and healthy cohorts with p-values of comparison.

	CAD Patients $(N = 12)$	Healthy Cohorts $(N = 10)$	Comparison (p-value)
Male, %	75% (N = 9)	80% (N = 8)	
Age, years	61 ± 9	52 ± 10	0.029
Height, cm	174 ± 8	174 ± 8	0.936
Weight, kg	82 ± 18	81 ± 17	0.893
Hypertension, %	75% (N = 9)		
Type 2 Diabetes, %	25% (N = 4)		
BSA, m2	1.99 ± 0.24	1.98 ± 0.25	0.918
SBP, mmHg	115 ± 16	112 ± 14	0.63
DBP, mmHg	64 ± 12	61 ± 11	0.633
HR, beats per minute	64 ± 11	58 ± 9	0.188
EDV-Cine, ml	231 ± 61	175 ± 27	0.011
ESV-Cine, ml	146 ± 52	73 ± 19	0.0001
SV-Cine, ml	84 ± 24	102 ± 9.5	0.033
EF-Cine, %	37 ± 9	59 ± 5	0.0001
LV-mass, grams	128 ± 32	$102 \pm 14$	0.022





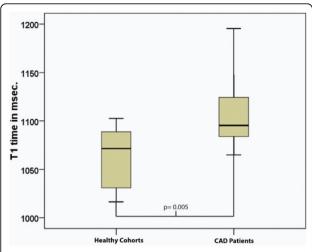


Figure 1 boxplot with mean (line in box), interquartile range (box) and complete range (whisker) of native T1 time in remote myocardium of CAD patients and healthy cohorts (p = 0.0005).

were measured by manually drawing a region of interest (ROI) on the three mid-ventricular slices and excluding the infarct area. An unpaired-samples T-test analysis was used to test for statistically significant differences between the two groups.

### Results

Patient characteristics are summarized in Table 1. LGE hyperenhancement was observed in all CAD patients. The mean native  $T_1$  time in the remote area myocardium of the CAD patients was significantly *higher* than the native  $T_1$  value of the myocardium in the healthy group (1107  $\pm$  36 ms. vs. 1061  $\pm$  32 ms.; p = 0.005) (Figure 1).

#### **Conclusions**

Our data suggest there are diffuse changes in remote/ normal myocardium resulting in abnormal/higher native  $T_1$  times in CAD patients with prior myocardial infarction. Further studies are needed to assess the prognostic value of an abnormal native  $T_1$  in the remote region among CAD patients.

Published: 27 January 2016

doi:10.1186/1532-429X-18-S1-P102

Cite this article as: Bellm et~al: The native  $T_1$  in remote myocardium of patients with prior chronic infarction is not normal. *Journal of Cardiovascular Magnetic Resonance* 2016 **18**(Suppl 1):P102.

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