

ORAL PRESENTATION

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Visual detection and characterization of chronic myocardial infarctions in patients using native T₁ maps at 3T

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Background

Late Gadolinium Enhancement (LGE) Cardiovascular Magnetic Resonance (CMR) is routinely used for characterizing chronic myocardial infarctions (cMIs), but it is contraindicated in patients with end-stage chronic kidney disease. We investigated whether native T₁ mapping at 3T can be used to detect and characterize cMIs in patients with prior STEMI and NSTEMI.

Methods

Breath-held 2D native T₁ maps (8 TIs with 2 Look-Locker cycles of 3+5 images; minimum TI = 120ms; TI increment = 80 ms; flip angle = 35°; bandwidth = 1085 Hz/pixel; voxel size = 1.5x1.5x8mm³) and LGE images (IR-prepared FLASH; optimal TI to null remote myocardium; TR/TE = 6.54/3.27ms; flip angle = 20°; bandwidth = 460 Hz/pixel; voxel size = 1.2x1.2x8mm³) were acquired in patients with prior STEMI (n=15) and NSTEMI (n=17) at 3T at a median of 13.6 years after acute MI. cMI location, size and transmuralities were determined using Mean+5SD criterion relative to remote myocardium. Visual detection of cMI territories on LGE images and T₁ maps were assessed by two independent reviewers.

Results

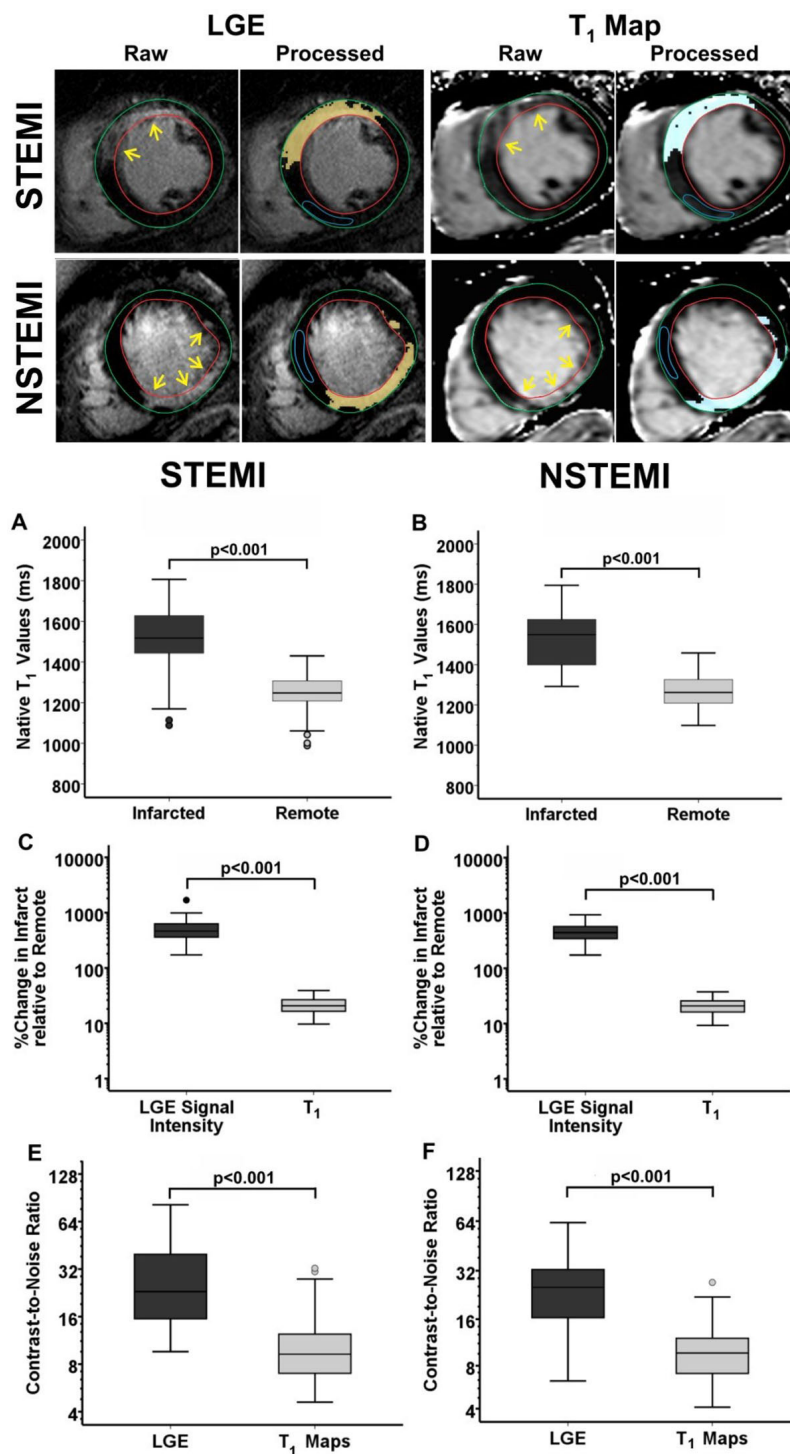
Representative native T₁ maps and LGE images from two patients, one with prior STEMI, and one with prior NSTEMI are shown in Fig. 1. Relative to remote myocardium, median T₁ of the cMI was 271ms higher in STEMI patients (Infarct: 1517ms; Remote: 1247ms;

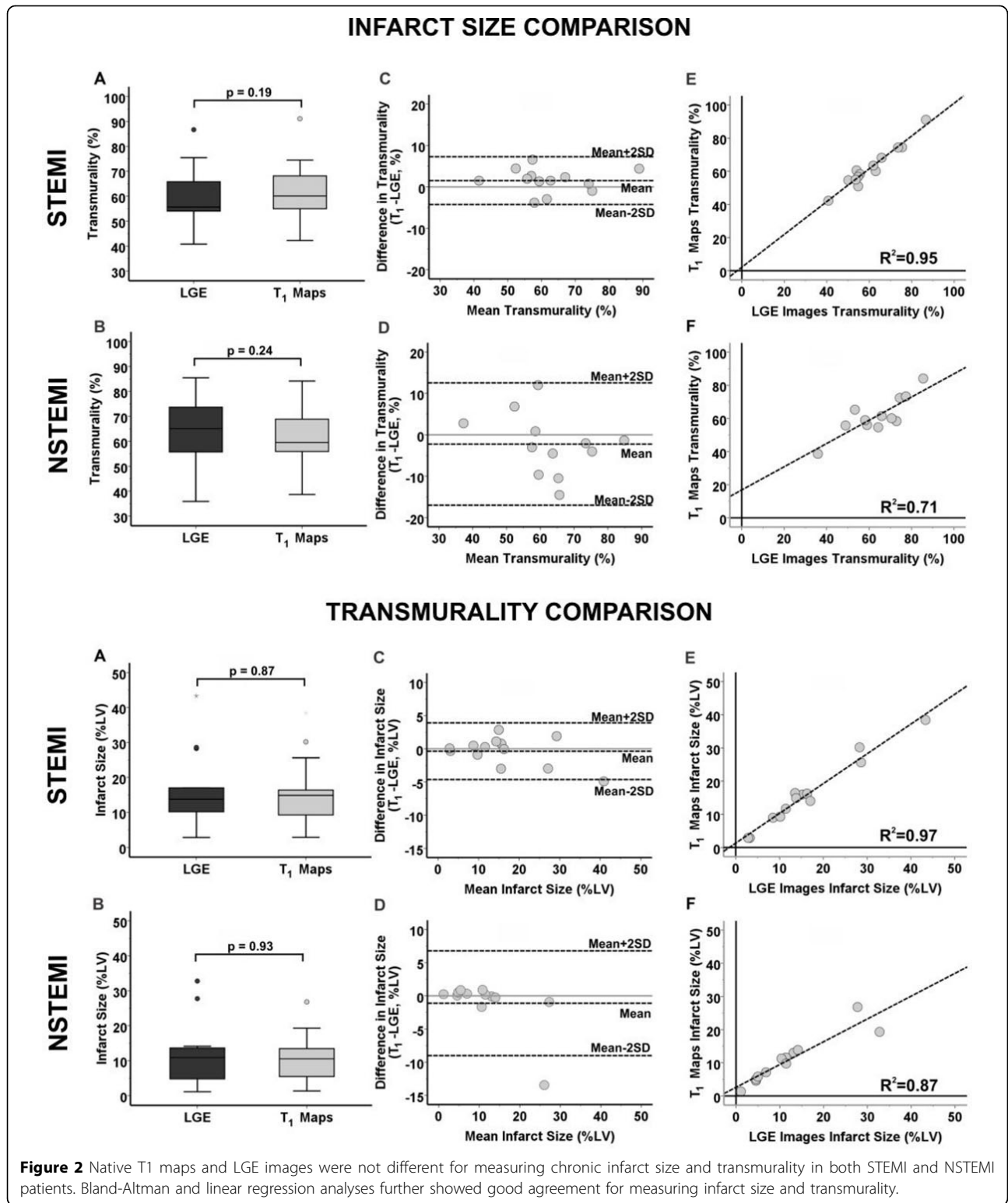
p<0.001; Fig. 1), and 229ms higher in NSTEMI patients (Infarct: 1549ms, Remote: 1262ms; p<0.001; Fig. 1). Median percentage change in LGE signal intensity (LGE-SI) of the cMI relative to remote myocardium was significantly higher than that of percentage change in T₁ in both STEMI (LGE: 465%, T₁: 21%; p<0.001) and NSTEMI (LGE: 441%, T₁: 20%; p<0.001) patients. Median CNR of LGE images was also 2.5-fold higher relative to that of T₁ maps in both STEMI (LGE: 23.1; T₁: 9.2; p<0.001) and NSTEMI (LGE: 25.3; T₁: 9.7; p<0.001) patients. LGE images and native T₁ maps were not different for measuring cMI size (STEMI - LGE: 13.8%; T₁: 14.9%; p=0.87; NSTEMI - LGE: 10.9%; T₁: 10.5%; p=0.93; Fig. 2) and transmuralities (STEMI - LGE: 55.6%; T₁: 60.1%; p=0.19; NSTEMI - LGE: 64.3%; T₁: 60.9%; p=0.24). Statistical analyses showed good agreement between LGE images and T₁ maps for measuring cMI size (STEMI: bias=-0.4±2.1%; R²=0.97; NSTEMI: bias=-1.1±3.9%; R²=0.87) and transmuralities (STEMI: bias=1.5±2.9%; R²=0.99; NSTEMI: bias=-2.2±7.4%; R²=0.71). Sensitivity and specificity of native T₁ maps for detecting cMIs based on threshold criterion were 93% and 97% respectively (STEMI); and 93% and 92% respectively (NSTEMI). Sensitivity and specificity of native T₁ maps for visual detection of cMI were: 61% and 85% (STEMI); and 67% and 90% (NSTEMI).

Conclusions

Native T₁ maps can reliably detect and characterize cMIs in STEMI and NSTEMI patients when the location of remote myocardium is known. Further increase in image contrast may be necessary to improve visual

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detection sensitivity of chronic MI territories to the levels observed with LGE.

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