



Acute Coronary Syndromes

PROGNOSTIC SIGNIFICANCE OF INFARCT CORE PATHOLOGY IN ST-ELEVATION MYOCARDIAL INFARCTION SURVIVORS REVEALED BY QUANTITATIVE T2-WEIGHTED CARDIAC MAGNETIC RESONANCE

Poster Contributions

Poster Hall B1

Sunday, March 15, 2015, 3:45 p.m.-4:30 p.m.

Session Title: From Cardiac Arrest, LV Failure to Myocardium Salvage

Abstract Category: 2. Acute Coronary Syndromes: Clinical

Presentation Number: 1209-053

Authors: *David Carrick, Caroline Haig, Sam Rauhalampi, Nadeem Ahmed, Ify Mordi, Margaret McEntegart, Mark Petrie, Hany Eteiba, Stuart Hood, Stuart Watkins, Mitchell Lindsay, Ahmed Mahrous, Aleksandra Radjenovic, Ian Ford, Niko Tzemos, Keith Oldroyd, Colin Berry, University of Glasgow, BHF Glasgow Cardiovascular Research Center, Glasgow, United Kingdom, Golden Jubilee National Hospital, Clydebank, United Kingdom*

Background: Myocardial transverse relaxation time (T2, ms) is a fundamental magnetic property of tissue that is related to water content and mobility. The pathophysiological and prognostic importance of native myocardial T2 in acute STEMI patients is unknown.

Methods: We performed a prospective single center cohort study in reperfused STEMI patients who underwent CMR 2 days and 6 months post-MI. T2-weighted CMR was measured in myocardial regions-of-interest. The infarct territory and microvascular obstruction were depicted with late gadolinium enhancement CMR. All-cause death or heart failure hospitalization was a pre-specified outcome that was assessed during follow-up.

Results: 324 STEMI patients (mean age 59 years, 237 males) gave informed consent (14 July 2011 - 22 November 2012). Of these, 295 STEMI patients had evaluable T2 maps. All 324 had follow-up assessments (median duration 860 days). Infarct size was $18 \pm 14\%$ of LV mass. 164 (51%) patients had late microvascular obstruction whereas 197 (61%) patients had an infarct core revealed by native T2. Native T2 within the infarct core (53.9 ± 4.8) was higher than in the remote zone (49.7 ± 2.1 ms; $p < 0.01$) but lower than in the area-at-risk (62.9 ± 5.1 ms) ($p < 0.01$). Native T2 in the infarct core was negatively multivariably associated with heart rate, Killip class, and peak neutrophil count at presentation (all $p < 0.05$). Baseline T2 core (ms) was univariably associated with baseline LVEF (0.31 (0.04, 0.58); $p = 0.023$), but not associated with LVEF or volumes at 6 months. Thirty (10.4%) patients died or experienced a heart failure event. These events included 5 cardiovascular deaths, 3 non-cardiovascular deaths and 22 episodes of heart failure (Killip Class 3 or 4 heart failure ($n = 20$) or defibrillator implantation ($n = 2$)). T2-core (ms) was associated with a reduced risk of all-cause death or heart failure hospitalization (hazard ratio 0.786, 95% CI 0.658, 0.939; $p = 0.008$) including after adjustment for LVEF at baseline ($p = 0.017$) or LV end-diastolic volume at baseline ($p = 0.009$) (Figure 3).

Conclusion: Infarct core revealed by native T2 was common and independently associated with all-cause death or heart failure hospitalization post-discharge.