# Journal of Cardiovascular Magnetic Resonance



Oral presentation Open Access

# Pre-operative ischaemia on CMR stress perfusion is a marker for prolonged post-operative stay after coronary artery bypass grafting

Joyce Wong\*<sup>1</sup>, Anthony Mathur<sup>2</sup>, Peter G Mills<sup>2</sup>, Redha Boubertakh<sup>1</sup>, Rakesh Uppal<sup>2</sup>, Alan Wood<sup>3</sup>, Mark Westwood<sup>1</sup> and LCeri Davies<sup>1</sup>

Address: ¹CMR Dept, London Chest Hospital, Barts and the London NHS Trust, London, UK, ²London Chest Hospital, Barts and the London NHS Trust, London, UK and ³St Bartholomew's Hospital, Barts and the London NHS Trust, London, UK

\* Corresponding author

from 13th Annual SCMR Scientific Sessions Phoenix, AZ, USA. 21-24 January 2010

Published: 21 January 2010

Journal of Cardiovascular Magnetic Resonance 2010, 12(Suppl 1):O39 doi:10.1186/1532-429X-12-S1-O39

This abstract is available from: http://jcmr-online.com/content/12/\$1/O39 © 2010 Wong et al; licensee BioMed Central Ltd.

### **Background**

Stress perfusion CMR accurately identifies inducible perfusion defects, but its role prior to coronary artery bypass surgery (CABG) is unclear. Similarly, late gadolinium enhancement (LGE) in CMR is well established in the identification of viability, but prognostic value post-CABG is uncertain. Out of the 2600 stress perfusion studies performed at our centre from 2008-2009, early post-operative outcomes were assessed in 56 consecutive patients who had CABG following a CMR scan. 28 patients underwent adenosine stress perfusion imaging, while all 56 underwent LGE imaging.

## Methods

56 patients (10 females, age 64 +/- 12 years) were imaged on a 1.5 Tesla MR Scanner (Philips Achieva, Best, Netherlands) within 2 months of cardiac catheterisation demonstrating significant multivessel disease, prior to urgent or elective CABG. Adenosine (140 mcg/kg/min) was administered for 3 minutes to achieve myocardial hyperaemia following a standard CMR stress perfusion protocol, with single-bolus injection of gadoterate meglumine contrast (0.1 mmol/kg, Dotarem, Guerbet, SA). Early post-operative outcomes were assessed over a mean follow up period of 6 months. Continuous variables were reported as mean ± standard error; groups (based on the number of viable or ischaemic segments) were compared using either Mann-Whitney, t-test or Fisher's exact test.

#### Results

Across the cohort, mean left ventricular (LV) end diastolic volumes (EVD) were increased at  $204 \pm 10$  mls, with a low mean ejection fraction (EF) of  $28 \pm 2.5\%$ . 24 patients out of 56 had LGE with 3 or more non-viable segments, and had a significantly longer post-operative stay ( $11.1 \pm 1.2$  vs  $7.4 \pm 0.6$  days, p < 0.01), and more severe LV impairment and dilatation (EF  $27 \pm 2.0$  vs  $50 \pm 3.5\%$ , p < 0.01, EDV  $242 \pm 13$  vs  $161 \pm 12$  mls, p < 0.0001). This group were also more likely to have had an unstable presentation with an acute coronary syndrome (ACS) (16/24 vs 12/40, p < 0.008). Patients with 6 or more ischaemic segments on stress CMR perfusion imaging also had a longer post-operative stay ( $11.6 \pm 2.0$  n = 12 vs  $6.8 \pm 0.6$  days p < 0.05, n = 17) but this group had no differences in EDV or EF.

At 6 months, the incidence of angina and heart failure symptoms were similar across all groups. All cause mortality and cardiac readmissions were also similar across all groups.

#### **Conclusion**

6 or more ischaemic segments on stress perfusion imaging are associated with a longer post-operative stay after CABG. Similarly, 3 or more non-viable segments detected by LGE are also associated with a longer post-operative stay. Although promising, further prospective studies are required to establish the prognostic role of stress perfusion in revascularisation.

Publish with **Bio Med Central** and every scientist can read your work free of charge

"BioMed Central will be the most significant development for disseminating the results of biomedical research in our lifetime."

Sir Paul Nurse, Cancer Research UK

Your research papers will be:

- available free of charge to the entire biomedical community
- $\bullet$  peer reviewed and published immediately upon acceptance
- cited in PubMed and archived on PubMed Central
- $\bullet$  yours you keep the copyright

Submit your manuscript here: http://www.biomedcentral.com/info/publishing\_adv.asp

