

POSTER PRESENTATION

Open Access

# A comparison of late gadolinium enhancement magnetic resonance imaging and left atrial endocardial voltage

James Harrison<sup>1,2\*</sup>, Nick Linton<sup>1,2</sup>, Steven Williams<sup>1,2</sup>, Rashed Karim<sup>1</sup>, Kawal Rhode<sup>1</sup>, Matthew Wright<sup>1,2</sup>, Tobias Schaeffter<sup>1</sup>, Reza Razavi<sup>1</sup>, Mark O'Neill<sup>1,2</sup>

From 16th Annual SCMR Scientific Sessions  
San Francisco, CA, USA. 31 January - 3 February 2013

## Background

Following catheter ablation for atrial fibrillation (AF), late gadolinium enhancement magnetic resonance imaging (LGE MRI) may be able to visualise areas of fibrosis and therefore reduced endocardial voltage. Unlike previous studies describing qualitative visual comparisons, we have developed a new quantitative technique for comparison of left atrial (LA) endocardial voltage with 3D LGE MRI signal intensity and have applied it in patients undergoing repeat left atrial ablation.

## Methods

Ten patients who had undergone previous catheter ablation for AF, and who represented with either paroxysmal AF (n=4) or atrial tachycardia (AT) (n=6) underwent pre-ablation LGE MRI. A 3D LA reconstruction was created by projecting the LGE data on to a segmented LA shell (signal intensities were displayed as the number of standard deviations (SD) from the mean signal intensity of the atrial blood pool).

During the ablation procedure, high density (mean number of points:  $368 \pm 127$ ) LA endocardial voltage maps were acquired in either sinus rhythm or AT using CARTO 3 (Biosense Webster).

The 3D LGE MRI reconstructions and endocardial voltage maps were manually segmented into LA regions (left and right WACA, roof, mitral line, anterior, posterior,

inferior, septum and lateral) using custom-written software to derive the mean signal intensity and endocardial voltage for each segment.

## Results

Mean LGE MRI signal intensity and endocardial voltage were calculated for a total of 131 atrial segments in the ten patients.

In the atrial segments in which the mean LGE MRI signal intensity was low (0 to 3 SD from the mean intensity of the atrial blood pool - 'healthy'), the mean  $\pm$  SD endocardial voltage was  $0.86 \pm 0.7$  mV, whereas when the mean LGE MRI signal intensity was high ( $>3$  SD - 'scar'), the mean endocardial voltage was significantly lower at  $0.51 \pm 0.4$  mV ( $p < 0.004$ ) (Figure).

## Conclusions

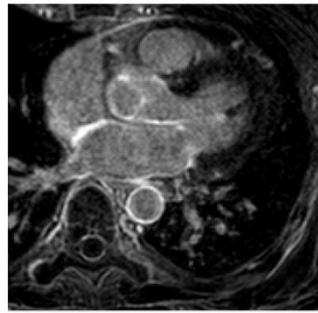
High LGE MRI signal intensity correlates with areas of low endocardial voltage, however, low voltage can still occur in areas of low signal intensity on LGE MRI. Further refinement of this technique is needed to permit the non-invasive assessment of atrial substrate prior to repeat catheter ablation.

## Funding

British Heart Foundation Clinical Research Training Fellowship.

<sup>1</sup>Division of Imaging Sciences & Biomedical Engineering, King's College London, London, UK

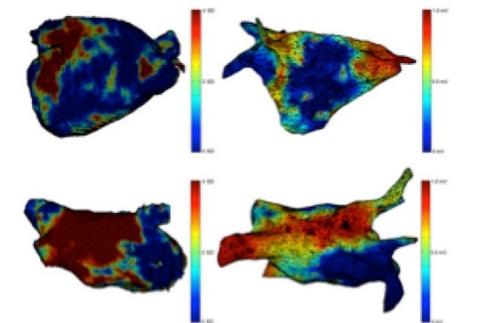
Full list of author information is available at the end of the article



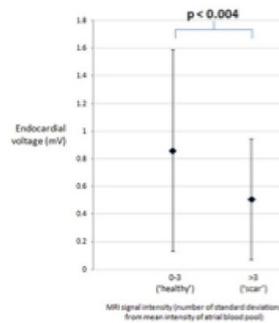
Transverse slice from a 3D LGE MRI of a patient with previous pulmonary vein isolation for paroxysmal AF



Manual segmentation of a 3D MRI reconstruction into LA regions



Corresponding posteroanterior (top row) and superior (bottom row) views of a 3D LGE MRI (left column) and a CARTO endocardial voltage map (right column) from two different patients



Graph showing endocardial voltage (mean  $\pm$  SD) for low ('healthy') and high ('scar') MRI signal intensities

**Figure 1**

#### Author details

<sup>1</sup>Division of Imaging Sciences & Biomedical Engineering, King's College London, London, UK. <sup>2</sup>Department of Cardiology, St Thomas' Hospital, London, UK.

Published: 30 January 2013

doi:10.1186/1532-429X-15-S1-P67

**Cite this article as:** Harrison *et al.*: A comparison of late gadolinium enhancement magnetic resonance imaging and left atrial endocardial voltage. *Journal of Cardiovascular Magnetic Resonance* 2013 **15**(Suppl 1):P67.

**Submit your next manuscript to BioMed Central and take full advantage of:**

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at  
[www.biomedcentral.com/submit](http://www.biomedcentral.com/submit)

 **BioMed Central**