

Poster presentation

Open Access

## Myocardial $T_1$ -Mapping in chronic myocardial infarction: preliminary results of unenhanced and contrast enhanced MR imaging using Gadobutrol

Kerstin U Bauner<sup>\*1</sup>, Andreas Biffar<sup>1</sup>, Daniel Theisen<sup>1</sup>, Torleif Sandner<sup>1</sup>, Andreas Greiser<sup>2</sup>, Maximilian F Reiser<sup>1</sup> and Bernd Wintersperger<sup>1</sup>

Address: <sup>1</sup>Ludwig-Maximilian-University Munich, Campus Grosshadern, Munich, Germany and <sup>2</sup>Siemens Medical Solutions, Erlangen, Germany

\* 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):P154 doi:10.1186/1532-429X-12-S1-P154

This abstract is available from: <http://jcmr-online.com/content/12/S1/P154>

© 2010 Bauner et al; licensee BioMed Central Ltd.

### Purpose

At a given field strength tissues present with specific  $T_1$ -values. Reference values for normal unenhanced myocardium have been established. We hypothesize, that infarcted myocardial tissue can be delineated from normal myocardium by means of  $T_1$ -maps in unenhanced and contrast-enhanced scans.

### Materials and methods

13 patients with chronic myocardial infarction were examined at 1.5 T (Magnetom Avanto, Siemens Healthcare). A modified Look-Locker inversion recovery (MOLLI) sequence (TR/TE 200.7/1.03 msec; TI 100-4000 msec; flip 35°) was performed pre- and 10 min post-contrast (0.15 mmol/kg gadobutrol, Bayer Schering Pharma) at an apical, midmyocardial and basal short axis position. For calculation of  $T_1$ -values signal intensities of myocardial and infarcted tissue were measured at 11 points of time [1] with two blocks of 3 and a third block 5 consecutive image acquisitions. Within each block TI increased by steps of 80 msec.

15 minutes after contrast medium application a single slice IR GRE was employed for imaging of delayed enhancement.

Data were post-processed with an in-house built software (PMI 0.4).  $T_1$  maps were created on the basis of unenhanced (fig. 1) and enhanced (fig. 2) data. Areas of normal and infarcted myocardial tissue were identified on

delayed enhancement images and the regions of interest were copied to the unenhanced and enhanced MOLLI images. The analyses of  $T_1$ -values were performed for normal myocardium (MYO), infarcted myocardium (CMI) and the left ventricular cavity (LVC). In addition  $T_1$ -ratios of MYO/LVC and CMI/LVC were calculated.

Student's t-test was used for statistical analysis of acquired and calculated data.

### Results

The comparison of  $T_1$ -values of MYO (fig. 3) and CMI (fig. 4) revealed significant differences in pre-contrast scans ( $1028 \pm 36$  vs.  $1210 \pm 63$  msec;  $p < 0.001$ ), as well as CMI  $T_1$ -values in comparison to LVC ( $1210 \pm 63$  vs.  $1509 \pm 70$  msec;  $p < 0.001$ ). The calculated ratios of MYO/LVC and CMI/LVC were also significantly different ( $0.68 \pm 0.04$  vs.  $0.79 \pm 0.06$ ;  $p < 0.001$ ) in pre-contrast scans. In post-contrast evaluations differences of  $T_1$ -values in MYO and CMI were equally high ( $360 \pm 46$  vs.  $224 \pm 55$  msec;  $p < 0.001$ ) resulting in significantly different  $T_1$ -ratios of MYO/LVC ( $1.5 \pm 0.21$ ) and CMI/LVC ( $0.9 \pm 0.10$ ;  $p < 0.001$ ).

### Conclusion

MR-measurements of  $T_1$ -values with the LVC as reference allow for differentiation of infarcted areas from normal myocardium. Further studies are warranted for a normalization of values in order to reduce the dependency on contrast timing, dose and agent.

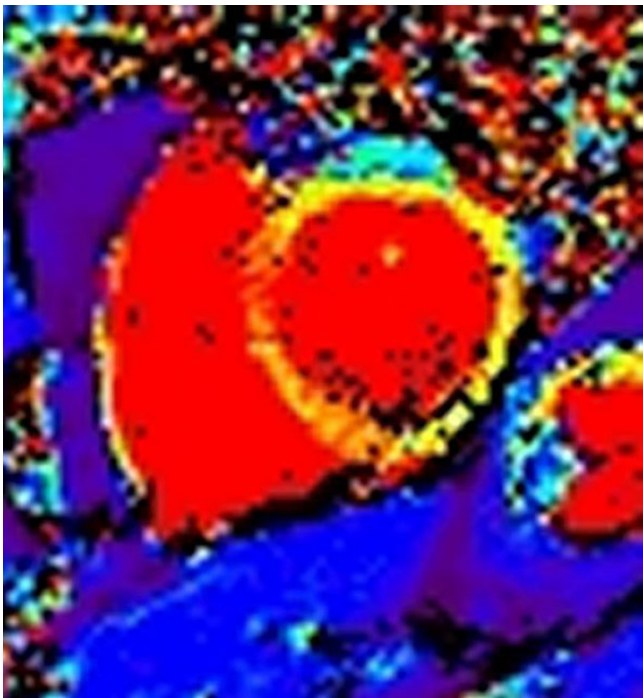


Figure 1

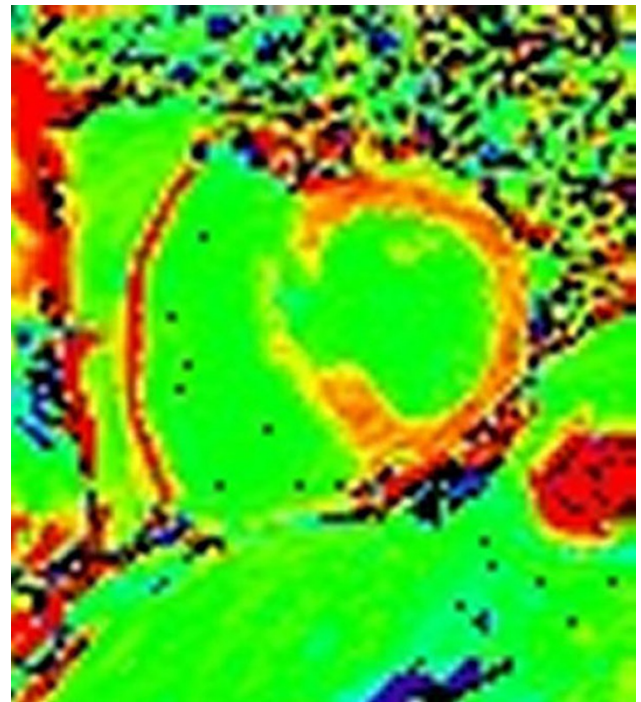


Figure 2

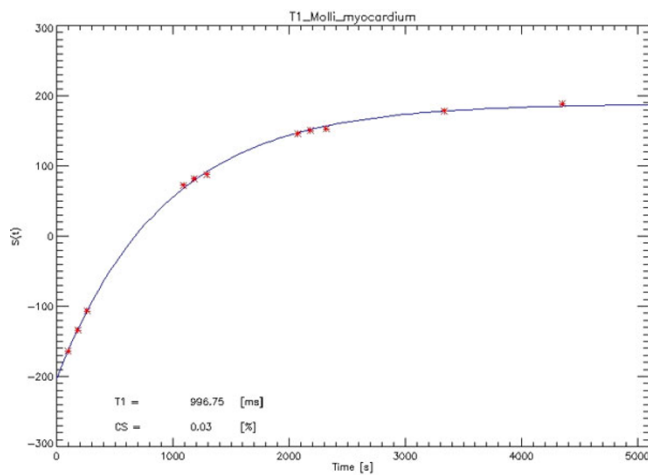


Figure 3

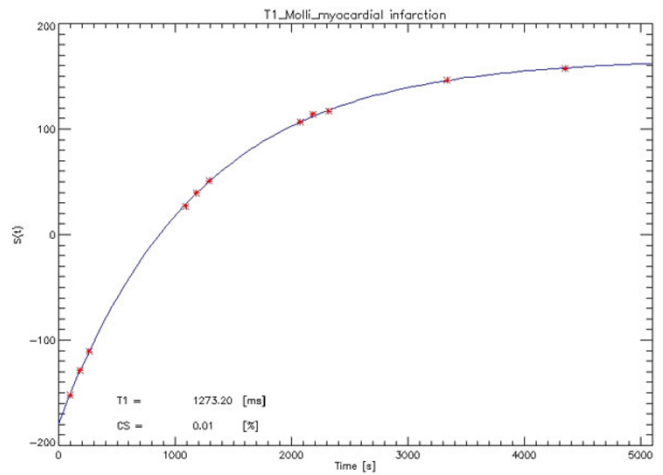


Figure 4

## References

1. Daniel Messroghli R, et al.: **Optimization and Validation of a Fully-Integrated Pulse Sequence for Modified Look-Locker Inversion-Recovery (MOLLI) T1 Mapping of the Heart.** *JMRI* 2007, **26**:1081-1086.

Publish with **BioMed 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
- peer reviewed and published immediately upon acceptance
- cited in PubMed and archived on PubMed Central
- yours — you keep the copyright

Submit your manuscript here:  
[http://www.biomedcentral.com/info/publishing\\_adv.asp](http://www.biomedcentral.com/info/publishing_adv.asp)

