

POSTER PRESENTATION

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# Quantitative whole-heart three-dimensional magnetic resonance myocardial perfusion imaging in systole and diastole at 3.0T

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## Background

Two-dimensional (2D) perfusion-CMR has been shown to have greater diagnostic accuracy than single-photon emission computed tomography but remains limited by a lack of complete myocardial coverage. Three-dimensional (3D) whole-heart myocardial perfusion CMR addresses this limitation and has recently been shown to be clinically feasible. However, the feasibility and potential clinical utility of *quantitative* 3D perfusion measurements, as already shown with 2D-perfusion-CMR and positron emission tomography, has yet to be evaluated. The purpose of this study was to establish the feasibility of quantitative 3D-perfusion-CMR to detect coronary artery disease (CAD). Additionally, as 3D-perfusion-CMR offers the opportunity to select the phase of acquisition, a secondary objective was to determine differences between systolic and diastolic estimates of myocardial blood flow (MBF).

## Methods

35 patients underwent 3D-perfusion-CMR (Philips 3T Achieva TX) with data acquired at both end-systole and mid-diastole (Fig 1). Systolic and diastolic perfusion images were analyzed in separate reporting sessions in random order. Image quality (0=non-diagnostic, 1=poor, 2=adequate and 3=excellent) and the occurrence of artifact related to respiratory-motion, k-t reconstruction or dark-rim artifact (0=none, 1=mild, 2=moderate and 3=severe) were scored. MBF and myocardial perfusion reserve (MPR) were estimated on a per patient and per

territory basis by Fermi function deconvolution. CAD was defined as luminal stenosis  $\geq 70\%$  on quantitative coronary angiography.

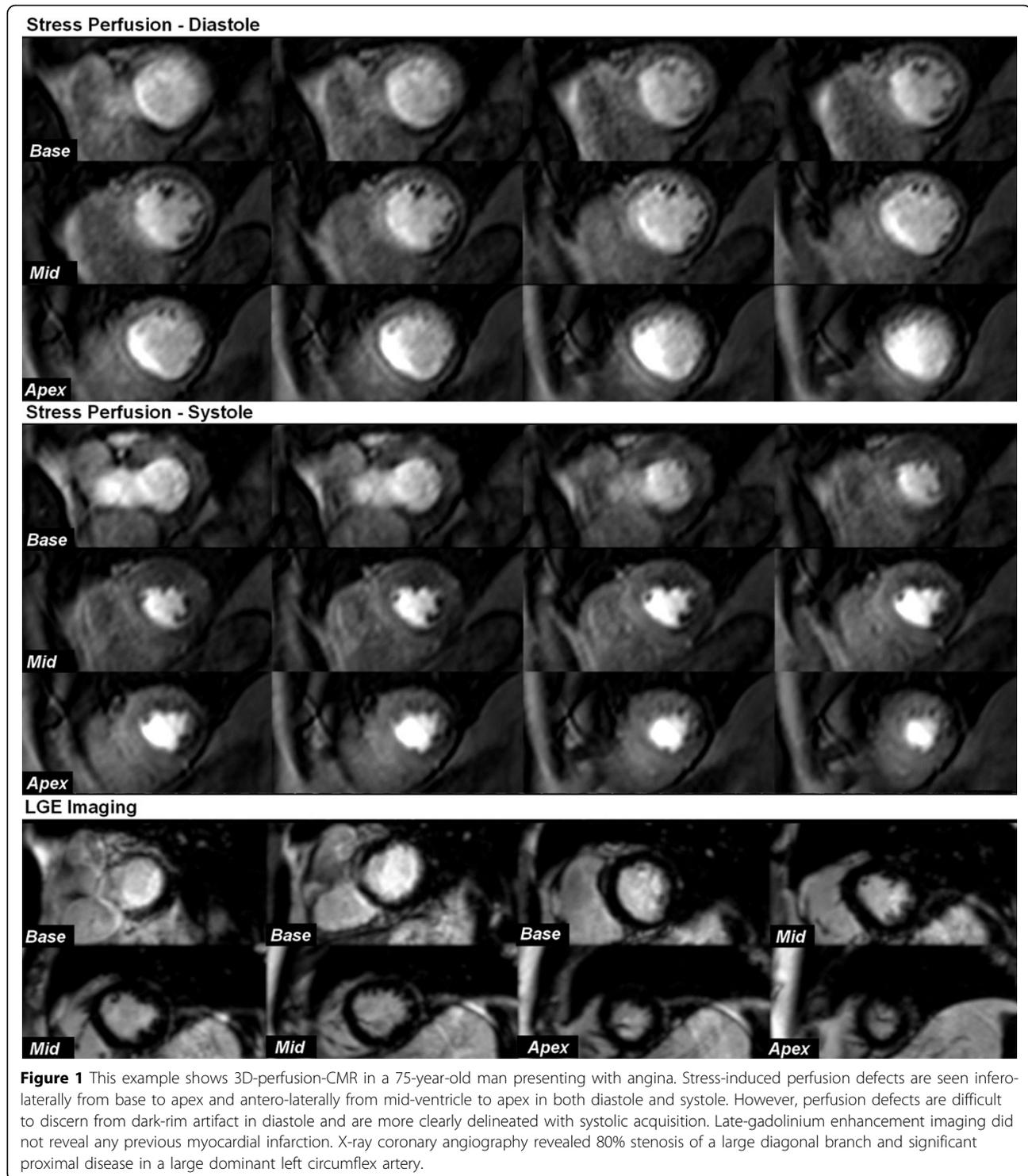
## Results

38 coronary territories had significant CAD. MPR had a high diagnostic accuracy for the detection of CAD, in both systole and diastole (area under curve: 0.92 vs. 0.94;  $p=0.41$ ) (Fig 2). At rest, systolic and diastolic MBF estimates were similar - in both normal and diseased territories (no CAD:  $1.24 \pm 0.15$  vs.  $1.25 \pm 0.15$  ml/g/min,  $p=0.27$ ; CAD:  $1.24 \pm 0.15$  vs.  $1.26 \pm 0.14$  ml/g/min,  $p=0.20$ ). At stress, diastolic MBF estimates were significantly greater than systolic estimates (no CAD:  $3.21 \pm 0.50$  vs.  $2.75 \pm 0.42$  ml/g/min,  $p<0.0001$ ; CAD:  $2.13 \pm 0.45$  vs.  $1.98 \pm 0.41$  ml/g/min,  $p<0.0001$ ). The diastolic/systolic stress MBF ratio was significantly reduced in territories with CAD (CAD:  $1.08 \pm 0.06$  vs. no CAD:  $1.17 \pm 0.11$ ;  $p<0.0001$ ). Systolic acquisition had a higher overall image quality score (median: 3 vs. 2,  $p=0.002$ ) and was less prone to artifact than diastolic acquisition (median artifact score: 0 vs. 1;  $p<0.0001$ ). In particular, there was a greater frequency of dark-rim artifact in diastole compared to systole (19 vs. 9 patients).

## Conclusions

We have shown that quantitative 3D-perfusion-CMR is feasible and can be used to detect CAD with high diagnostic accuracy. Image quality and less artifact, make systole the

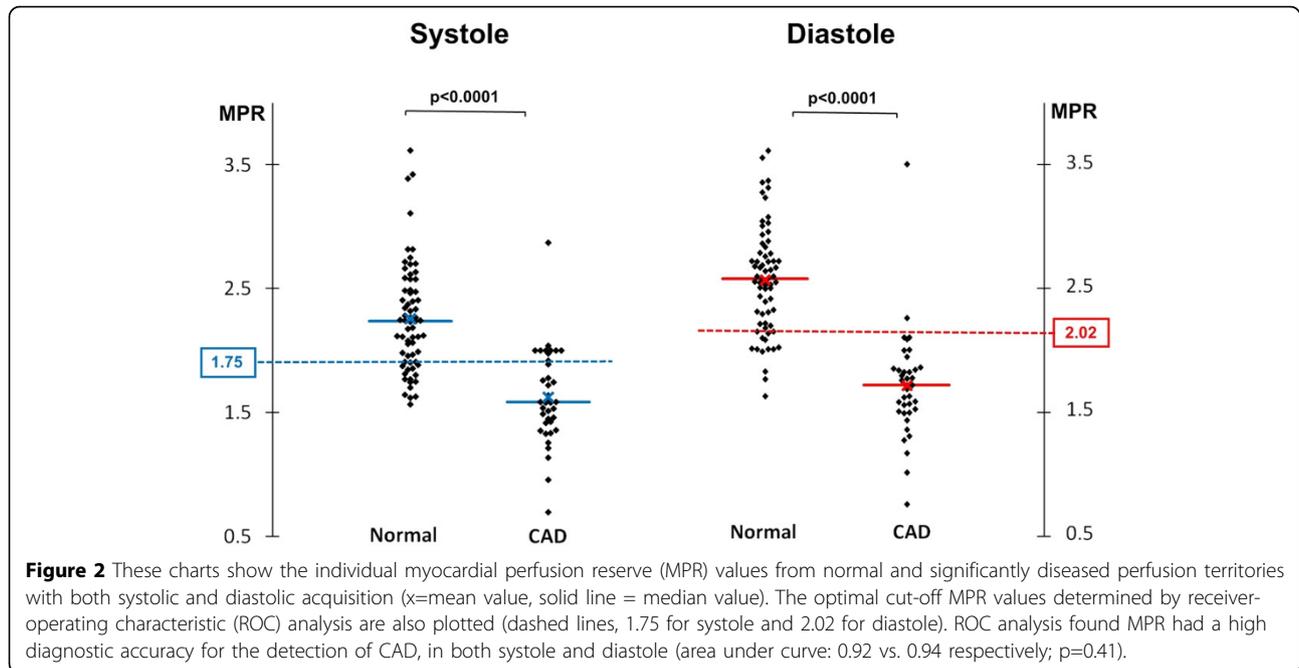
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preferred phase for acquisition in 3D-perfusion-CMR. Finally, there were significant differences in systolic and diastolic MBF estimates and therefore the phase of acquisition should always be stated in future quantitative studies.

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