Additional Diagnostic Value of Systolic Dysfunction Induced by Dipyridamole Stress Cardiac Magnetic Resonance Used in Detecting Coronary Artery Disease

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Introduction and objectives. Dipyridamole stress perfusion cardiac magnetic resonance (CMR) is used to detect coronary artery disease (CAD). However, few data are available on the diagnostic value of the systolic dysfunction induced by dipyridamole. This study investigated whether the induction of systolic dysfunction supplements the diagnostic information provided by perfusion imaging in the detection of CAD.

Methods. Overall, 166 patients underwent dipyridamole CMR and quantitative coronary angiography, with CAD being defined as a stenosis ≥70%. Systolic dysfunction at rest, systolic dysfunction with dipyridamole, induced systolic dysfunction, and stress first-pass perfusion deficit (PD) and delayed enhancement were quantified.

Results. In the multivariate analysis, PD (hazard ratio [HR] = 1.6; 95% confidence interval [CI], 1.33-1.91; \( P < .0001 \)) and induced systolic dysfunction (OR=1.8; 95% CI, 1.18-2.28; \( P < .007 \)) were independently associated with CAD and had a sensitivity and specificity of 92% and 62% and 43% and 96%, respectively. Patients were categorized as having no ischaemia (Group 1), PD but no induced systolic dysfunction (Group 2), or induced systolic dysfunction irrespective of PD (Group 3). In Group 3, the prevalence of CAD was higher than in Group 1 or 2 (86% vs 22% and 70%, respectively; \( P < .001 \)) and the risk of CAD was 2-fold higher than in Group 2 (OR=2.34; 95% CI, 1.07-5.13; \( P = .034 \)). Compared with Group 2, more hypoperfused segments were observed in Group 3 (6.2 [2.6] vs 7.4 [3.4]; \( P = .044 \)), and more diseased vessels (1.4 [1.0] vs 1.8 [0.9]; \( P = .036 \)). Adding induced systolic dysfunction to perfusion and clinical data improved the multivariate model’s C-statistic for predicting CAD (0.81 vs 0.87; \( P = .02 \)).

Conclusions. Combining induced systolic dysfunction with perfusion imaging increases the diagnostic accuracy of detecting CAD and enables patients with severe ischaemia and a high probability of CAD to be identified.
INTRODUCTION

Stress perfusion cardiac magnetic resonance imaging (CMR) is used for the evaluation of patients suspected of having coronary artery disease (CAD) in daily clinical practice.1 In a single setting, CMR allows the assessment of abnormal wall motion (AWM) and myocardial perfusion deficits (PD), 2 important parameters used for diagnosis of significant CAD.2-5 The sequential occurrence of these 2 phenomena during myocardial ischemia is part of what has been described as the “ischemic cascade.”6,7 Nevertheless, since the advent of myocardial perfusion imaging using vasodilators, the focus of attention has shifted away from stress inducible AWM to stress induced PD using CMR. Perfusion studies at stress have yielded promising results with a high accuracy for the detection of angiographic significant coronary lesions.4,8 CMR offers good temporal and excellent spatial resolution advocating this modality for the assessment of subtle changes in myocardial contractility during pharmacological stress, however there is little data on the diagnostic use of AWM at vasodilator stress for the detection of significant angiographic lesions.9

We have recently reported that the extent of AWM at peak dipyridamole stress is the CMR index most closely related to prognosis in patients with chest pain of possible coronary origin undergoing stress CMR.10 The objective of the present study was to analyze if inducible AWM affords additional diagnostic value for the detection of CAD.

METHODS

Study Group

Out of a total population of 600 patients who underwent stress CMR at our institution between January 2003 and May 2007, we retrospectively examined a study population of 166 patients (51 women (31%); age, 65 (11); range, 30-85; see Table 1) with chest pain of possible coronary origin who underwent dipyridamole stress CMR and coronary angiography within 3 months before or after CMR. The majority (88%) of patients underwent coronary angiography after CMR imaging. Patients underwent CMR studies because of inconclusive exercise testing in 18% cases, abnormal ECG at rest in 26%, inability to exercise in 28%, and as the first choice for ischemia evaluation in 28% of the cases. Twenty patients (12%) underwent CMR examination after coronary angiography due to evaluation of intermediate lesions. Patients, who had undergone a coronary revascularization, had a myocardial infarction within 3 months before the CMR study or any contraindication for dipyridamole were excluded from analysis. The study protocol was approved by the local ethics committee and all subjects gave informed consent.

TABLE 1. Clinical Characteristics (n=166)

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<table>
<thead>
<tr>
<th></th>
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<tbody>
<tr>
<td>Male sex (%)</td>
<td>115 (69)</td>
<td></td>
</tr>
<tr>
<td>Age, mean (SD), y</td>
<td>65 (11)</td>
<td></td>
</tr>
<tr>
<td>Diabetes, n (%)</td>
<td>64 (39)</td>
<td></td>
</tr>
<tr>
<td>Hypertension, n (%)</td>
<td>109 (66)</td>
<td></td>
</tr>
<tr>
<td>Hypercholesterolemia, n (%)</td>
<td>90 (54)</td>
<td></td>
</tr>
<tr>
<td>Smoker, n (%)</td>
<td>34 (21)</td>
<td></td>
</tr>
<tr>
<td>Previous IHD, n (%)</td>
<td>91 (55)</td>
<td></td>
</tr>
<tr>
<td>Previous MI, n (%)</td>
<td>45 (27)</td>
<td></td>
</tr>
<tr>
<td>Previous PTCA, n (%)</td>
<td>26 (16)</td>
<td></td>
</tr>
<tr>
<td>Previous bypass, n (%)</td>
<td>19 (11)</td>
<td></td>
</tr>
<tr>
<td>LVEF, %</td>
<td>60 (13)</td>
<td></td>
</tr>
<tr>
<td>LVEDD, mm</td>
<td>69 (23)</td>
<td></td>
</tr>
<tr>
<td>LVESD, mm</td>
<td>31 (25)</td>
<td></td>
</tr>
</tbody>
</table>

IHD indicates ischemic heart disease; LVEF, left ventricular ejection fraction; LVEDD, left ventricular end-diastolic diameter; LVESD, left ventricular end-systolic diameter; MI, myocardial infarction; PTCA, percutaneous transluminal coronary angioplasty. The data express mean (SD) or n (%).

Cine Images at Rest

Left ventricular function was assessed using cine images in 2-, 3-, 4 chamber, and short-axis views using a true fast imaging with steady-state precession sequence (SSFP) (repetition time/echo time 2.8 ms/1.2 ms, flip angle 58°, matrix 256 × 256, field of view 320 × 270 mm, slice thickness 6 mm). In
each patient, of the stack of short-axis views of cine images at rest, we selected 3 reference locations at base, mid-ventricle and apex which were repeated to assess perfusion, wall motion at stress and delayed enhancement.

First Pass Perfusion Imaging

Vasodilatation was induced with dipyridamole (0.56 mg/kg body weight over 4 min and if well tolerated by the patient up to 0.84 mg/kg delivered intravenously over 6 min). After the end of dipyridamole infusion, 0.1 mmol/kg gadopentate dimeglumine (Magnografl, Schering, Berlin, Germany) was injected intravenously at a speed of 5 mL/s. Then 5 sections equally separated in the short-axis view and 2 in the 2- and 4-chamber long-axis views were acquired for first-pass perfusion imaging every other heart beat (SSFP with a notched saturation pulse; inversion time, 125 ms; repetition time/echo time, 202 ms/1 ms; flip angle, 50°; matrix, 192 × 96; field of view, 350 × 220 mm; slice thickness, 8 mm).

Cine Images at Stress

Once peak myocardial enhancement was reached, to assess left ventricular function within the peak dipyridamole-induced vasodilatation (approximately 2-3 min after infusion) we used a fast-acquisition multislice SSFP cine image sequence (repetition time/echo time, 35 ms/1.1 ms; flip angle, 65°; matrix, 192 × 159; field of view, 350 × 304 mm; slice thickness, 8 mm) acquiring over a 14-s period in a single breath hold 4 slices in the short-axis view including the same 3 reference locations defined in the cine images at rest.

Delayed Enhancement Imaging

Delayed enhancement imaging was performed 10 min after contrast injection in the same locations evaluated for cine images at rest (segmented inversion recovery SSFP; repetition time/echo time, 700 ms/1.26 ms; flip angle, 45°; matrix, 256 × 184; field of view, 340 × 235 mm; slice thickness, 8 mm). The inversion time was adjusted to null normal myocardium.

CMR Data Analysis

CMR studies were analyzed by an experienced observer (M.P.L.L., 8 years of experience) blinded to all angiographic and patient data and unaware of the chronological order of the 2 examinations, using customized software (Syngo, Siemens, Erlangen, Germany). Left ventricular end-diastolic and end-systolic diameter (mm) and ejection fraction (Simpson method, %) were quantified by direct planimetry of left ventricular contours in the stack of short-axis cine images at rest. Segments location was defined in cine-images sequences applying the 17-segment model.11 We evaluated 5 dipyridamole stress CMR-derived indexes:

1. AWM at rest (AWM-rest). Number of segments showing hypokinesis, akinesis, or dyskinesis at baseline.12
2. AWM with dipyridamole (AWM-D). Number of segments showing hypokinesis, akinesis, or dyskinesis at stress.
3. Inducible AWM. Number of segments in which wall motion worsened with dipyridamole from normokinesis to hypokinesis or from hypokinesis to akinesis.
4. PD with dipyridamole. Number of segments showing a persistent delay (in at least 3 consecutive temporal images, compared to the other segments in the same slice) in the visual analysis of the enhancement pattern during the first pass of contrast through the myocardium.13
5. Delayed enhancement (DE). Number of segments showing enhancement in delayed enhancement imaging.

In a group of 20 patients evaluated for ischemic chest pain and not included in this study, interobserver agreement for the presence or absence of >1 segment with abnormal CMR indexes was as follows: AWM, 95%; AWM-D, 95%; abnormal perfusion, 90%; and delayed enhancement, 100%.10

According to the pathophysiologic considerations of the ischemic cascade,6,7 patients were categorized into 3 groups (Figure 2):
– Group 1: No evidence of ischemia. Normal perfusion study and no inducible AWM
– Group 2: Abnormal perfusion study but no inducible AWM
– Group 3: Inducible AWM present regardless of PD

**Coronary Angiography**

Angiographic data was evaluated by an experienced cardiologist (V.B., 8 years of experience), blinded to all CMR and patient data and unaware of the chronological order of the 2 examinations, using quantitative coronary angiography on a standard digital imaging system (HM3000, Philips, Best, The Netherlands). CAD was defined as ≥70% narrowing of the coronary lumen of 1 of the 3 major coronary arteries (or in a principal branch measuring ≥2 mm in diameter) or as ≥50% narrowing of the left main artery.

**Statistical Analysis**

Continuous data is expressed as the mean (standard deviation), and was compared by the unpaired t test. Categorical variables were compared with the χ² test. For calculation of sensitivity, specificity, positive predictive value, and negative predictive value variables were categorized according to the best cut-off value obtained in receiver-operating characteristic (ROC) curve analysis (0-1 segment vs >1 segment for AWM-rest, AWM-D, inducible AWM and DE; 0-2 segments vs >2 segments for PD). The higher cut-off value for PD was chosen due to susceptibility to artifacts and the inherent very high sensitivity and lower specificity of this index. The association of CMR with angiographic data was assessed with a forward logistic regression model adjusted for clinical parameters showing a P value <.2 in univariate analysis. For the 3 different groups another logistic regression was performed, also adjusted by clinical parameters with a P<.2 in univariate analysis. Odds ratios (OR) with 95% confidence interval (95% CI) were obtained. The additional value of inducible AWM was assessed by comparing the C-statistic of a model including baseline characteristics and PD with a model resulting by adding inducible AWM. A P value less than .05 was considered statistically significant. SPSS statistical package (Version 13.0, SPSS Inc, Chicago, Illinois, USA) and STATA (Version 9.0, StataCorp, College Station, Texas, USA) were used.
of the patients displayed PD, and 68 (41%) patients had abnormal delayed enhancement imaging.

**Angiographic Results**

Coronary angiography was abnormal in 119 patients (72%). Coronary angiography detected 209 significant coronary lesions (67 in the left anterior descending artery, 65 in the circumflex artery, and 77 in the right coronary artery). There were 58 patients (35%) with 1-vessel disease, 33 patients (20%) had 2-vessel disease, and 29 patients (17%) had 3-vessel disease.

**Detection of CAD**

A comparison of patients with and without CAD is displayed in Table 2. Patients with an abnormal angiography were mostly of male gender and tended to have a history of previous ischemic heart disease. Patients with angiographic CAD displayed a significant larger extent of all 5 CMR indexes. The diagnostic performance of the 5 CMR indexes for detecting CAD is displayed in Table 3. In the multivariate analysis, adjusted for clinical parameters, PD (1.6 [1.33-1.91]; \(P<.0001\), per segment), inducible AWM (1.8 [1.18-2.28]; \(P<.007\),...
DISCUSSION

Using a comprehensive approach of 5 CMR-derived parameters, our study shows that inducible AWM by dipyridamole offers additional diagnostic value to perfusion data for the detection of angiographically significant lesions in patients presenting with chest pain and with a high probability of CAD. Additional assessment of inducible AWM allows the identification of a group of patients with a higher probability of having angiographic CAD and a larger extent of ischemia in terms of hypoperfused segments and number of affected vessels.

Myocardial Perfusion Imaging

Since the advent of myocardial perfusion imaging, the focus of vasodilator stress CMR has shifted from the detection of inducible AWM to the detection of PD. Indeed, there are few CMR studies evaluating the diagnostic value of vasodilator inducible AWM for detecting CAD.9,14 PD is usually the consequence of significant epicardial coronary artery stenosis with a certain limitation due to microvascular disease15,16 and collateral circulation17 and therefore can be used to non-invasively diagnose and assess the severity of a luminal narrowing in CAD. In this study we used a widely accepted qualitative method by visual analysis. Using this approach, earlier studies have obtained encouraging results for diagnosing angiographic significant CAD.5,13,18,19 In our series of patients, we demonstrated a comparable diagnostic performance of PD for detection of CAD with a sensitivity of 0.92 and a specificity of 0.62. The presence of a PD is a very sensitive but in some cases

**TABLE 3. Diagnostic Performance of the CMR Parameters for Detection of Coronary Artery Disease (n=166)**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Sensitivity (95% CI)</th>
<th>Specificity (95% CI)</th>
<th>PPV (95% CI)</th>
<th>NPV (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>AWM-rest</td>
<td>0.49 (0.40-0.58)</td>
<td>0.77 (0.65-0.89)</td>
<td>0.84 (0.75-0.93)</td>
<td>0.37 (0.26-0.49)</td>
</tr>
<tr>
<td>n/No.</td>
<td>58/119</td>
<td>36/47</td>
<td>58/69</td>
<td>36/97</td>
</tr>
<tr>
<td>AWM-D</td>
<td>0.77 (0.60-0.85)</td>
<td>0.72 (0.69-0.84)</td>
<td>0.88 (0.81-0.94)</td>
<td>0.55 (0.45-0.64)</td>
</tr>
<tr>
<td>n/No.</td>
<td>91/119</td>
<td>34/47</td>
<td>91/104</td>
<td>34/62</td>
</tr>
<tr>
<td>Inducible AWM</td>
<td>0.43 (0.34-0.52)</td>
<td>0.96 (0.90-1.0)</td>
<td>0.96 (0.91-1.0)</td>
<td>0.40 (0.27-0.53)</td>
</tr>
<tr>
<td>n/No.</td>
<td>51/119</td>
<td>45/47</td>
<td>51/53</td>
<td>45/113</td>
</tr>
<tr>
<td>PD</td>
<td>0.92 (0.88-0.97)</td>
<td>0.62 (0.48-0.76)</td>
<td>0.86 (0.80-0.92)</td>
<td>0.76 (0.69-0.84)</td>
</tr>
<tr>
<td>n/No.</td>
<td>110/119</td>
<td>29/47</td>
<td>110/128</td>
<td>29/38</td>
</tr>
<tr>
<td>DE</td>
<td>0.49 (0.40-0.58)</td>
<td>0.79 (0.67-0.90)</td>
<td>0.85 (0.77-0.94)</td>
<td>0.38 (0.26-0.49)</td>
</tr>
<tr>
<td>n/No.</td>
<td>58/119</td>
<td>37/47</td>
<td>58/68</td>
<td>37/98</td>
</tr>
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</table>

AWM indicates abnormal wall motion; AWM-D, abnormal wall motion at dipyridamole stress; AWM-rest, abnormal wall motion at rest; CI, confidence interval; DE, delayed enhancement; PD, perfusion deficit; NPV, negative predictive value; PPV, positive predictive value.
a less specific parameter for the presence of a flow limiting stenosis at stress, due to overestimation of clinical significance of the ischemia detected.20

Pathophysiological Considerations. Role of the “Ischemic Cascade”

With increasing severity, myocardial ischemia leads from myocardial perfusion abnormalities to the occurrence of diastolic and systolic AWM and electrocardiographic changes and finally anginal symptoms, a sequence usually referred to as “the ischemic cascade.”6,7 With pharmacological stress CMR it is possible to assess 2 of these steps of the ischemic cascade; inducible AWM and the PD preceding it. Consequently in our study, 96% of the patients with inducible AWM presented an abnormal perfusion study. Inducible AWM was not a sensitive but a highly specific parameter with an excellent positive predictive value. In fact only 2 patients displaying inducible AWM did not have severe angiographic lesions. One case was a female patient who had 3 patent coronary bypass grafts on 3 occluded coronary arteries and the other case was a man who had a 60% lesion in the territory of inducible AWM. After adjusting for clinical characteristics and CMR parameters the occurrence of inducible AWM afforded independent additional diagnostic value. Indeed, patients with inducible AWM had a 2-fold higher probability of having a pathologic angiography than did patients with only an abnormal perfusion study.

Patient Classification

From a pathophysiological point of view, this study validates the ischemic cascade by allowing the classification of patients according to the severity of ischemia. By applying inducible AWM, it was possible to identify a group of patients displaying more hypoperfused segments and more diseased vessels. It has been shown that patients with inducible AWM are also at highest risk for adverse outcome and benefit most from revascularization.20 Moreover, our results are in line with data derived from a recent meta-analysis comparing myocardial perfusion imaging with AWM assessed by stress echocardiography. This meta-analysis concludes that perfusion imaging is a sensitive modality with a somewhat low specificity for the detection of CAD while the contrary is true for AWM assessed by stress echo.21 Our study demonstrates that using CMR, inducible AWM offers additional diagnostic value in combination with perfusion data, making stress CMR a truly comprehensive approach in the evaluation of ischemia. PD, since it occurs early in the ischemic cascade, is very sensitive for the detection of ischemia.
ischemia, nevertheless due to detection of clinically irrelevant perfusion deficits and susceptibility to artifacts can yield false positive results. Inducible AWM with dipyridamole mainly occurs in severe epicardial stenosis (>75%). The low sensitivity might be owed to worse detection of lesser grade stenosis. Thus, inducible AWM might balance out the lack of specificity of perfusion data.

Different Pharmacological Stressors

The diagnostic and prognostic usefulness of dobutamine stress CMR has already been validated. Dobutamine more frequently provokes systolic dysfunction than vasodilators. However, as a consequence of the resulting high heart rate, dobutamine is not the ideal stressor to evaluate perfusion due to the inherent difficulties to reconstruct perfusion images when cardiac cycles shorten. Inducible AWM using vasodilators is not a frequent phenomenon and when it occurs it relates to severe ischemia. Paetsch et al observed that adenosine-inducible AWM occurred in segments with at least 75% transmural perfusion deficit and related to high grade (>75%) coronary stenosis. In contrast, dobutamine-induced AWM was observed even in areas without perfusion deficit and related to less severe (>50%) coronary stenosis.

Therefore, the use of dipyridamole, albeit being less potent than dobutamine for inducing AWM, might be advantageous for identifying a group of patients with severe and clinically significant ischemia, and by that might improve of specificity in stress CMR. Moreover, the additional diagnostic value of inducible AWM that could be demonstrated in the present study can be obtained in a very short period of time (during a 14 second breath hold) not prolonging examination time, thereby making this approach easily applicable in clinical practice.

Study Limitations

Our study is limited by its retrospective nature. Out of all 600 patients who underwent a dipyridamole stress CMR study those with an abnormal CMR study were more likely to undergo coronary angiography than were patients with an entirely normal CMR examination. This probably limits the generalizability of our data and might account for the high prevalence of CAD and the rather low specificity of PD in our study.

CONCLUSIONS

Using dipyridamole stress CMR, adding inducible AWM to perfusion imaging offers additional diagnostic value for the detection of CAD in patients presenting with chest pain and known or suspected CAD. According to the ischemic cascade, the presence of inducible AWM allows the identification of a group of patients with more severe ischemia and with a higher probability of having angiographic CAD.
REFERENCES