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Diagnostic Accuracy of Dobutamine Stress Echocardiography for Detection of Cardiac Allograft Vasculopathy in Orthotopic Heart Transplant Patients

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Abstract

Objective: Cardiac allograft vasculopathy is one of the leading causes of late graft failure and subsequent death in orthotopic heart transplant. Although invasive coronary angiography is the gold standard modality for detection of cardiac allograft vasculopathy, dobutamine stress echocardiography has been recently frequently used as an alternative. Our aim was to evaluate the diagnostic performance of dobutamine stress echocardiography for detection of cardiac allograft vasculopathy in transplant patients.

Methods: A retrospective analysis was conducted using a total of 150 dobutamine stress echocardiographic exams that were performed on 99 patients in our institution, with paired coronary angiogram and no acute rejection, within a median of 538 [interquartile range 371–816] days. Sensitivity and specificity of dobutamine echocardiography to detect allograft vasculopathy was evaluated. Allograft vasculopathy was defined as Grade 1 or higher based on ISHLT criteria. A positive dobutamine stress echo result was defined by new or worsening wall motion abnormality.

Results: Median age of the population at transplant was 34 [interquartile range 22–46] years; 76 (77%) patients were male. Allograft vasculopathy was present in 31 (20.6%) out of 150 coronary angiograms. Only 7 (4.6%) of that number were positive on dobutamine stress echocardiography. Allograft vasculopathy was defined as Grade 1 or higher based on ISHLT criteria. A positive dobutamine stress echo result was defined by new or worsening wall motion abnormality.

Conclusions: Overall, positivity of dobutamine stress echocardiography in patients after heart transplant is low. It has high specificity, but very low sensitivity for detection of cardiac allograft vasculopathy. Dobutamine stress echocardiography should only be cautiously used as an alternative to coronary angiography.

Keywords: Dobutamine echocardiography, Stress echocardiography, Cardiac allograft vasculopathy, Heart transplantation, Myocardial ischemia, Diagnostic imaging tools

1. Introduction

Orthotopic heart transplant (OHT) is an established method for treating advanced heart failure that is unresponsive to conventional therapy. Since its establishment, the technique has continuously improved as is evident with the significant increase in post-OHT survival, where the
median survival after transplant is currently over 10 years [1].

However, cardiac allograft vasculopathy (CAV) remains one of the dominant causes of late graft failure and death [1]. CAV has an unpredictable course and can quickly transform into diffuse obstructive disease [2–6]. Various immunologic and non-immunologic factors play a role in the pathophysiology of CAV. Increased CRP (C-reactive protein) is a strong predictor of CAV development after heart transplantation, highlighting the role of inflammation [6]. Significant myocardial ischemia and infarctions may go unnoticed due to atypical presentations in denervated hearts. Early detection of coronary involvement with ischemia is therefore of paramount importance. Invasive coronary angiography is the gold standard method for the assessment of coronary artery disease [5], but in CAV its performance might be impaired by the diffuse nature of coronary involvement [7]. Furthermore, repeating coronary angiography for frequent assessment is not optimal due to its invasive nature, contrast use and radiation exposure. Intravascular ultrasound (IVUS) tends to be more sensitive [8], given the diffuse longitudinal concentric hypertrophy due to fibrous intimal hyperplasia that appears in the affected coronary arteries with CAV. However, it also needs invasive coronary access.

Dobutamine stress echocardiography (DSE), which has acceptable sensitivity and specificity, was established in the 1990s as a feasible alternative to invasive coronary angiography [9]. The International Society of Heart and Lung Transplantation (ISHLT) guidelines classified DSE as Class IIa for CAV screening in patients that cannot undergo invasive evaluation or possibly in children [5]. DSE is the best validated noninvasive method, with the sensitivity of 85% as compared to angiography and IVUS and an outstanding negative predictive value [9,10]. Recently, however, the performance of DSE in this setting has been questioned [3,4]. A systematic review and meta-analysis study found that DSE had a limited sensitivity to detect early CAV, but its specificity was much higher [11]. A few studies have suggested using speckle tracking echocardiography or contrast echocardiography to improve the diagnostic accuracy of DSE in detecting CAV [12,13].

We were unable to find published literature on this concept in the Middle Eastern population. The purpose of this study is to evaluate the performance of DSE as a non-invasive screening modality in determining the presence of coronary vasculopathy in our population of patients with OHT at our tertiary care center, the only institution that performs OHT in the entire Middle East: King Faisal Specialist Hospital & Research Centre (KFSH&RC).

2. Materials and methods

2.1. Population

A retrospective analysis on a consecutive set of 99 patients was performed. The main design of the study is shown in the Figure. These patients were evaluated in our clinic after orthotopic heart transplant. They underwent both coronary angiography (with or without endomyocardial biopsy) and DSE. All included heart transplants were performed at the King Faisal Specialist Hospital, Riyadh, KSA.

Post-transplant, endomyocardial biopsy, echocardiography, chest X-ray, and laboratory evaluation are performed on week number 3–4 depending on whether the patient is being discharged. From the week of discharge, the biopsies were done every month until 6 months and then every two months until 12 months. Echocardiogram is performed more frequently if the patient presents pericardial effusion, or whenever it was clinically indicated. A total of approximately 10 biopsies and 5–6 echocardiograms were performed during the first year. Annual HT admission at 12 months with tests is performed (including cardiac catheterization with biopsy and echocardiography when clinically indicated).

After 1 year, a transthoracic echocardiography is usually performed every 6 months. Biopsy is done annually for 5 years and then every 2–3 years as required.

Abbreviations

CAV Cardiac Allograft Vasculopathy
CHD Coronary Heart Disease
DCM Dilated Cardiomyopathy
DSE Dobutamine Stress Echocardiography
FCMP Familial Cardiomyopathy
HCM Hypertrophic Cardiomyopathy
ICA Invasive Coronary Angiography
ICM Ischemic Cardiomyopathy
ISHT International Society of Heart and Lung Transplantation
IQR Interquartile range
IVUS Intravascular ultrasound
LAD Left Anterior Descending artery
LM Left Main artery
OHT Orthotopic Heart Transplant
RHD Rheumatic Heart Disease
RCM Restrictive Cardiomyopathy
This medical chart retrospective study was approved by the institutional ethics committee (IRB number 2001055).

2.2. Echocardiography

Both resting echocardiography and DSE were performed using E7 or E9 (General Electric, Milwaukee, Wisconsin, USA) or iE33 or EPIQ 7 machines (Philips, Amsterdam, The Netherlands).

Routine resting echocardiography was performed in all patients according to current recommendations [14,15].

For the DSE, routine protocol was used as previously described [16]. Briefly, standard dosing of dobutamine was used, from 10 to 40 ug/kg/min with 10 ug/kg/min steps every 3 minutes to achieve ≥ 85% of the maximal target heart rate. Up to 1 mg of atropine was added if heart rate could not be achieved using dobutamine alone. Wall motion abnormalities, ECG changes, heart rate and blood pressure and occurrence of symptoms were evaluated at every stage.

Positive DSE was based on the appearance of or worsening of wall motion abnormality in at least two myocardial segments based on the 16 segments ASE model [15].

2.3. Invasive coronary angiography

Invasive coronary angiography and endomyocardial biopsy were performed in a routine manner at our institution whenever indicated as necessary by clinical findings and routine surveillance. Coronary angiography was evaluated for CAV based on the recommendations of the ISHLT [17]. Briefly, CAV Grade 0 was defined as no detectable angiographic lesions; Grade 1 (mild) was defined by angiographic left main (LM) < 50% stenosis or primary/branch vessel with a maximum lesion of <70% stenosis, including diffuse narrowing; Grade 2 (moderate) was angiographic LM < 50% stenosis, a single primary vessel ≥70% stenosis or isolated branch stenosis ≥70% in branches of two systems; Grade 3 (severe) was angiographic LM ≥ 50% stenosis, two or more primary vessels ≥70% stenosis, isolated branch stenosis ≥70% in all systems or graft dysfunction/evidence of restrictive physiology. Acute rejection was graded based on the revised ISHLT criteria [18]. Follow up/death of patients was obtained using patients’ charts and/or hospital information.

2.4. Statistical analysis

Due to common non-normal distribution, continuous values are presented as median [25th-75th percentile] or median with interquartile range (IQR) and binary data as proportions. Due to non-normal distributions, differences in parameters of DSE were evaluated using Mann–Whitney U or chi-square tests, as appropriate. Results were evaluated using contingency tables and reported as sensitivity and specificity ± standard error (SE). Because of
clustered nature of data, a correction for clustered observations using ratio estimator method was performed [19]. A p value < 0.05 was considered significant. Analysis was done using R software version 3.0.2 (R Foundation for Statistical Computing, Vienna, Austria).

3. Results (figure)

3.1. Baseline characteristics

For evaluation of CAV, catheterizations with acute rejection on endomyocardial biopsy were excluded. In total, there were 150 DSE performed on 99 patients that had paired DSE and coronary angiography with no acute rejection. Baseline characteristics of our cohort are shown in Table 1. During follow-up period, 21 patients died.

3.2. Prevalence of CAV

There were 150 coronary angiographies done on the 99 patients, out of which 31 (20.6%) were positive for CAV. Out of the 31, 24 had Grade 1 CAV, 4 had Grade 2, and 3 had Grade 3.

3.3. Characteristics of DSE

Baseline characteristics by DSE are shown in Table 2. No significant differences were seen when comparing positive and negative DSE, but there was a trend towards lower baseline LVEF in the positive patients (p = 0.086).

3.4. Dobutamine stress echocardiography for detection of CAV

Overall, DSE and ICA were done 538 (IQR 371–816) days apart. Cross tabulation is shown in Table 3 and reveals sensitivity of 3.2 ± 3.3% and specificity of 94 ± 2.9% for CAV on invasive angiography.

We have analyzed the 23 false negative dobutamine echocardiograms and 3 of them had Grade 2

Table 1. Baseline characteristics of the study population.

<table>
<thead>
<tr>
<th>Total number</th>
<th>99</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age at transplant (years)</td>
<td>34 [22, 47]</td>
</tr>
<tr>
<td>Male gender</td>
<td>76 (77%)</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>31 (31%)</td>
</tr>
<tr>
<td>Atrial fibrillation</td>
<td>10 (10%)</td>
</tr>
<tr>
<td>Creatinine (umol/l)</td>
<td>84 [66, 107]</td>
</tr>
<tr>
<td>Arterial Hypertension</td>
<td>29 (29%)</td>
</tr>
<tr>
<td>Heart failure etiology</td>
<td>29 (29%)</td>
</tr>
<tr>
<td>DCM</td>
<td>59 (60%)</td>
</tr>
<tr>
<td>ICM</td>
<td>25 (26%)</td>
</tr>
<tr>
<td>RHD</td>
<td>2 (2%)</td>
</tr>
<tr>
<td>HCM</td>
<td>2 (2%)</td>
</tr>
<tr>
<td>CHD</td>
<td>2 (2%)</td>
</tr>
<tr>
<td>Peripartum</td>
<td>2 (2%)</td>
</tr>
<tr>
<td>FCMP</td>
<td>1 (1%)</td>
</tr>
<tr>
<td>Chemotherapy induced</td>
<td>1 (1%)</td>
</tr>
<tr>
<td>RCM</td>
<td>2 (2%)</td>
</tr>
<tr>
<td>Other/unknown</td>
<td>3 (3%)</td>
</tr>
</tbody>
</table>

DCM - Dilated Cardiomyopathy, ICM - Ischemic Cardiomyopathy, RHD - Rheumatic Heart Disease, HCM - Hypertrophic Cardiomyopathy, CHD - Coronary Heart Disease, FCMP - Familial Cardiomyopathy, RCM - Restrictive Cardiomyopathy.

Table 2. Characteristics of dobutamine stress echocardiography.

<table>
<thead>
<tr>
<th>Total (n = 150)</th>
<th>Positive for RWMA (n = 8)</th>
<th>Negative for RWMA (n = 142)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Years from heart transplant</td>
<td>4.26 [2.87, 6.68]</td>
<td>6.00 [4.10, 6.72]</td>
<td>4.19 [2.87, 6.67]</td>
</tr>
<tr>
<td>LV EF (%)</td>
<td>63 [59, 70]</td>
<td>61 [57, 63]</td>
<td>63 [60, 71]</td>
</tr>
<tr>
<td>Mitral regurgitation Grade ≥ 2</td>
<td>3 (2%)</td>
<td>0 (0%)</td>
<td>3 (2.1%)</td>
</tr>
<tr>
<td>Maximal dobutamine dose (μg/kg/min)</td>
<td>40 [30,40]</td>
<td>40 [38, 40]</td>
<td>40 [30, 40]</td>
</tr>
<tr>
<td>Atropine used</td>
<td>14 (9.3%)</td>
<td>1 (12%)</td>
<td>13 (9.2%)</td>
</tr>
<tr>
<td>At Baseline</td>
<td>87 [80, 96]</td>
<td>89 [88, 92]</td>
<td>85 [79, 96]</td>
</tr>
<tr>
<td>Heartrate (s⁻¹)</td>
<td>124 [116, 135]</td>
<td>122 [114,132]</td>
<td>124 [116, 135]</td>
</tr>
<tr>
<td>SBP (mmHg)</td>
<td>78 [70, 86]</td>
<td>82 [78, 85]</td>
<td>78 [69, 86]</td>
</tr>
<tr>
<td>DBP (mmHg)</td>
<td>145 [135, 154]</td>
<td>148 [143, 158]</td>
<td>145 [134, 154]</td>
</tr>
<tr>
<td>At max dobutamine</td>
<td>154 [138, 170]</td>
<td>142 [114, 163]</td>
<td>154 [140, 170]</td>
</tr>
<tr>
<td>Heartrate (s⁻¹)</td>
<td>75 [62, 82]</td>
<td>64 [57, 90]</td>
<td>75 [63, 82]</td>
</tr>
</tbody>
</table>

DBP – diastolic blood pressure, LV EF – left ventricular ejection fraction, RWMA – regional wall motion abnormality, SBP – systolic blood pressure.
CAV while the remaining had Grade 1. There were 7 false positive DSE, two of them were from one patient who did have a myocardial bridge in the mid left anterior descending artery (LAD).

3.5. Dobutamine stress echocardiography in patients with acute cellular rejection

There were 2 patients with DSE that were excluded from the main analysis of our cohort of 99 patients because they had shown acute cellular rejection, both of them Grade 1R. There were 3 DSEs performed on these patients, 1883 days before, 617 days before and 798 days after their ICA. All of the DSE were negative.

4. Discussion

DSE remains used and recommended for non-invasive detection of CAV [5]; however, recent studies have questioned its validity and predictive value [3,4]. Our data seem to support this conclusion. We have found low sensitivity of DSE to detect CAV. This can be explained by a relatively low prevalence of significant CAV in our population and/or a mild degree of CAV. Indeed, our data are in agreement with recent publications where sensitivity was as low as 7% when patients with Grade 1 CAV were included and has risen to 28% after only patients with Grade 2 or higher were considered [3]. In another study, a sensitivity as low as 0% has been reported [4]. We have shown a relatively high specificity that is comparable to previous reports [2]. The specificity can be probably even higher, since one of our patients without significant CAV had myocardial bridge in mid LAD which might have reasonably caused the positivity of DSE [20].

Recently, it has also been shown that DSE might not be predictive of outcome in patients after OHT [3], while other studies have shown prognostic significance [9,21]. Unfortunately, our data is too small to draw any definitive conclusions.

Other improvements for echocardiographic detection of allograft vasculopathy have been proposed. Coronary blood flow assessment can be used alone or in conjunction with dobutamine stress echocardiography to improve detection [13,22]. Other studies have evaluated myocardial deformation imaging during stress echocardiography have found significant association with CAV [12]. However, these methods are not yet reflected in current guidelines and it remains to be seen whether they will be able to improve echocardiographic detection of CAV in clinical practice.

The incidence of CAV based on invasive coronary angiography was comparable to some previous studies [22] while others have reported somewhat higher prevalence [23]. However, local differences in our population might be accountable for that.

Interestingly, all our patients with acute cellular rejection had negative DSE. This can be probably explained by large time differences between DSE and ICA as well as the mild degree of cellular rejection (Grade 1R) in both patients.

4.1. Limitations

The main limitation remains the retrospective nature of our study where indication for DSE and coronary angiography was driven by clinical need, making a systematic assessment difficult. Furthermore, low prevalence of significant CAV in our young patient population, and large interval time between DSE and ICA reduces the statistical power of the study. We have not used intravascular ultrasound as that was not the routine practice at our institution, therefore the true prevalence of CAV on ICA might be underestimated [8,9].

5. Conclusion and clinical perspectives

This study indicates that DSE has a very low sensitivity and excellent specificity for the detection of CAV in the OHT patient population. Based on our findings, which are in agreement with previous series, the routine clinical use of DSE as a diagnostic tool for the assessment of CAV could not be the most suitable technique. Therefore, other imaging modalities should be considered for the evaluation of CAV post OHT.

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Author contribution

Conception and design of Study: Mohammed Mahmoodurrahman, Hamzah Ruxshan Juhardeen, Talal Al Otaibi, Bahaa M Fadel, Dania Mohty. Literature review: Mohammed Mahmoodurrahman, Josef Marek, Hamzah Ruxshan Juhardeen, Vera

None.

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