Mitral regurgitation determined by radionuclide cardiography: dependence on posture and exercise

Henning Kelbek, Jan Aldershvile, Knud Skagen, Per Hildebrandt, Steen Levin Nielsen

Abstract

Objectives—To evaluate dependence of posture and exercise on the degree of mitral regurgitation using combined first pass and equilibrium radionuclide cardiography.

Design—24 patients with clinically stable chronic mitral regurgitation and sinus rhythm were studied by first pass list mode and simultaneous multigated frame mode equilibrium radionuclide cardiography using red cells labelled with technetium-99m.

Results—When patients changed posture from supine to sitting upright, left ventricular volumes decreased considerably. Regurgitation tended to increase in patients with valve prolapse but decreased in patients with ischaemic heart disease and dilated cardiomyopathy. During submaximal bicycle exercise cardiac output increased without dilatation of the left ventricle. The increase in left ventricular forward stroke volume was more pronounced than that in the total stroke volume, leading to a considerable decrease in the regurgitant flow through the mitral valve. The repeatability and observer variability of radionuclide determination of regurgitation was acceptable, with limits of agreement within about 10%.

Conclusions—Change in posture induces a normal haemodynamic response in most patients with chronic mitral regurgitation; the effect of posture on regurgitation depends on the underlying disease. Mild to moderate exercise causes no deterioration in the severity of regurgitation.

Patients and methods

STUDY PROTOCOL

To evaluate the impact of posture on mitral regurgitation 18 patients were examined on the same day while sitting at 60° and while lying supine. Half of the patients were randomly allocated to have the examinations in the upright position performed first. Ten patients were examined upright at rest and during bicycle exercise to determine the influence of physical stress on the severity of the disease.

The reproducibility of the radionuclide technique was evaluated in two ways. Hour to hour variation was measured by examining 13 supine patients twice on the same day, the examinations being one hour apart. Day to day variation was measured by examining 14 patients twice, the examinations being two weeks apart. In addition, data from 20 patients were stored and processed by two independent observers.

Exercise was performed on an ergometer bicycle, and the workload was progressively adjusted with increments of 25 W every two minutes until the heart rate was increased by 35–40% compared with the value at rest. No attempt was made to provoke cardiac symptoms during exercise. Data were collected one minute after attaining the highest workload which was maintained for up to 10 minutes. All patients gave their informed consent, and the study protocol was approved by the local ethics committee.

PATIENTS

We studied 24 patients (14 men and 10 women, aged 27 to 82 (mean 59)) with moderate to severe chronic mitral valve regurgitation as judged from the results of contrast ventriculography or Doppler echocar-
Table 1  Mean (SD) values of haemodynamic variables, in supine and upright positions in 18 patients with mitral regurgitation.

<table>
<thead>
<tr>
<th>Supine</th>
<th>Upright</th>
<th>95% Confidence interval for difference upright – supine</th>
</tr>
</thead>
<tbody>
<tr>
<td>Heart rate (beats/min)</td>
<td>72 (14)</td>
<td>76 (14)</td>
</tr>
<tr>
<td>Blood pressure (mm Hg):</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Systolic</td>
<td>126 (19)</td>
<td>127 (21)</td>
</tr>
<tr>
<td>Diastolic</td>
<td>78 (7)</td>
<td>84 (8)</td>
</tr>
<tr>
<td>Left ventricular ejection fraction (%)</td>
<td>58 (18)</td>
<td>60 (18)</td>
</tr>
<tr>
<td>Total stroke volume (ml)</td>
<td>132 (60)</td>
<td>109 (42)</td>
</tr>
<tr>
<td>Regurgitant volume (ml)</td>
<td>70 (52)</td>
<td>57 (41)</td>
</tr>
<tr>
<td>Regurgitation fraction (%):</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Patients with mitral prolapse and chorda rupture</td>
<td>52 (19)</td>
<td>57 (15)</td>
</tr>
<tr>
<td>Patients with secondary mitral regurgitation</td>
<td>27 (12)</td>
<td>12 (11)</td>
</tr>
<tr>
<td>Whole group</td>
<td>43 (22)</td>
<td>43 (25)</td>
</tr>
</tbody>
</table>

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graphy. Ten patients had mitral valve prolapse, six ischaemic heart disease, three rheumatic valve disease, two ruptured chordae (primarily valve prolapse), two previous endocarditis, and two dilated cardiomyopathy. The patients’ conditions were clinically stable (New York Heart Association classification groups I and II). For technical reasons (see below) only patients with sinus rhythm were included. Reasons for exclusion were pregnancy, lactation, left sided cardiac valve disease other than mitral valve regurgitation, and treatment with vasodilatory drugs.

Investigations

Heart rate was measured in a three lead electrocardiograph, and blood pressure was recorded sphygmomanometrically.

Radionuclide cardiography was performed by a combined first pass and multigated equilibrium technique after in vitro labelling of autologous red blood cells with technetium-99m. To determine the cardiac output from a bolus transit curve, roughly 500 MBq (14 mCi) of the radiotracer was injected through a cannula in the right basilic vein for each investigation. First pass data were acquired in list mode in a 256 x 256 matrix with a mobile gamma-camera equipped with a low energy all purpose collimator (Siemens) in the left anterior oblique projection; data were processed in a 128 x 128 frame format with a dedicated computer. A 10 second static image was acquired in the same format with simultaneous drawing of a blood sample to determine the activity in blood after complete mixing of the tracer. Multigated equilibrium cardiography was performed in the same projection in a 64 x 64 matrix and a duration of 16 frames per cardiac cycle.

Forward stroke volume was calculated as the activity recorded in the left ventricular region of interest multiplied by the tracer distribution volume and divided by the product of the area under the time activity curve in the same region and the heart rate.

The left ventricular ejection fraction was determined as 1 - the end systolic activity divided by end diastolic activity; the left ventricular end diastolic volume was calculated as the end diastolic frame activity corrected for specific blood activity and attenuation. The total (absolute) stroke volume was determined as the product of the left ventricular end diastolic volume and ejection fraction as assessed by the equilibrium method.

The regurgitant volume was calculated as the difference between the total and the forward stroke volume, and the regurgitation fraction was calculated as the regurgitant volume divided by the total stroke volume.

Statistical analysis

Data from the investigations performed while the patient was upright and supine and from those performed while the patient was resting and exercising were processed by one observer and evaluated from the 95% confidence intervals of the differences.

Repeat investigations performed to assess the hour to hour and day to day variation were also processed by one person, but two independent observers evaluated the data to determine the observer variation of the radionuclide technique. Owing to limited capacity of data storing, measurements of observer variation were restricted to data obtained while patients were supine. Repeated observer variation were evaluated by comparison of the difference against mean method described previously. The limits of agreement were expressed as 1-96 standard deviations of the differences between the two determinations.

Results

Posture dependent haemodynamic changes

From the supine to the upright position heart rate increased by 6% and diastolic blood pressure by 8% (table 1). The mean left ventricular volume decreased at end diastole from 228 ml to 185 ml (95% confidence interval for the difference -60 to -25 ml), with a concomitant 17% reduction in the total stroke volume (table 1 and fig 1). The volume changes were independent of left ventricular size. No change was recorded in the left ventricular ejection fraction. While patients were upright the mean forward stroke volume was reduced from 63 ml to 53 ml (-18 to -4 ml) compared with that recorded while they were supine. The decrease in the regurgitant volume was similar to the change in total stroke volume, leaving the regurgitation fraction for the whole group unaltered. Patients with mitral valve prolapse and ruptured chordae alone, however, tended to show increased regurgitation when they were upright, whereas patients with secondary mitral regurgitation due to ischaemic heart disease or dilated cardiomyopathy showed significantly less regurgitation when they sat up (table 1 and fig 1).

Exercis induced haemodynamic changes

The exercise induced increases in heart rate and systolic blood pressure were 37% and 17%, respectively, compared with the values at rest (table 2). No significant change occurred in left ventricular end diastolic volume, whereas small increments were seen in both total stroke volume and left ventricular
Table 2  Mean (SD) values of haemodynamic variables at rest and during submaximal bicycle exercise in 10 patients with mitral regurgitation

<table>
<thead>
<tr>
<th></th>
<th>Rest</th>
<th>Exercise</th>
<th>95% Confidence interval for difference (exercise - rest)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Heart rate (beats/min)</td>
<td>82 (13)</td>
<td>112 (11)</td>
<td>23 to 38</td>
</tr>
<tr>
<td>Blood pressure (mm Hg):</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Systolic</td>
<td>128 (23)</td>
<td>150 (18)</td>
<td>13 to 30</td>
</tr>
<tr>
<td>Diastolic</td>
<td>84 (10)</td>
<td>83 (15)</td>
<td>-6 to 5</td>
</tr>
<tr>
<td>Left ventricular ejection fraction (%)</td>
<td>68 (12)</td>
<td>70 (12)</td>
<td>0 to 4</td>
</tr>
<tr>
<td>Total stroke volume (ml)</td>
<td>109 (41)</td>
<td>115 (38)</td>
<td>1 to 12</td>
</tr>
<tr>
<td>Regurgitant volume (ml)</td>
<td>64 (38)</td>
<td>48 (40)</td>
<td>-31 to -2</td>
</tr>
</tbody>
</table>

Ejection fraction (table 2 and fig 2). A more pronounced increase of 26% (13% to 40%) was recorded in the forward stroke volume (fig 2), resulting in a considerable reduction in the regurgitant volume (table 2) and a 24% decrease (-38% to -10%) in regurgitation fraction during exercise (fig 2).

Reproducibility of Radionuclide Technique

Haemodynamic variables at rest lying down determined by the combined radionuclide technique after an hour and after 14 days showed high levels of reproducibility with no significant differences between the two determinations (table 3). The mean difference between regurgitation fractions determined after an hour was 2% (limits of agreement -7% to 10%), and the corresponding value after 14 days was 1% (limits of agreement -11% to 9%) (fig 3).

Table 4 shows the values determined directly by the observers. A small but significant difference was found in the two observers' determinations of the left ventricular ejection fraction. None of the derived variables was significantly different, and the limits of agreement for the regurgitation fraction calculated by the two observers were -9% to 9% (fig 4).

Discussion

The severity of mitral regurgitation is dependent on the afterload that opposes left ventricular ejection and on the systolic pressure gradient from the left ventricle to the left atrium.13-15 This gradient is most likely to increase when the patient stands up so the regurgitation fraction might be expected to be more pronounced while standing than while lying supine. The severity of regurgitation, however, would be expected to depend not only on left ventricular volumes but also on the nature of the underlying disease. Our results show that when the patients changed posture from supine to sitting upright, the changes in left ventricular volume were similar to those in healthy subjects.16-17 The results also indicate that when the patients were examined as a group no significant alteration

Table 3  Hour to hour and day to day variation in mean (SD) values of haemodynamic variables at rest in 14 supine patients with mitral regurgitation

<table>
<thead>
<tr>
<th></th>
<th>Hour to hour (n = 13)</th>
<th>Day to day (n = 14)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>First determination</td>
<td>Second determination</td>
</tr>
<tr>
<td>Heart rate (beats/min)</td>
<td>77 (17)</td>
<td>76 (16)</td>
</tr>
<tr>
<td>Blood pressure (mm Hg):</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Systolic</td>
<td>127 (25)</td>
<td>128 (25)</td>
</tr>
<tr>
<td>Diastolic</td>
<td>77 (11)</td>
<td>79 (9)</td>
</tr>
<tr>
<td>Left ventricular end diastolic volume (ml)</td>
<td>222 (81)</td>
<td>217 (77)</td>
</tr>
<tr>
<td>Left ventricular ejection fraction (%)</td>
<td>55 (19)</td>
<td>55 (19)</td>
</tr>
<tr>
<td>Stroke volume (ml):</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>120 (62)</td>
<td>119 (59)</td>
</tr>
<tr>
<td>Forward</td>
<td>60 (23)</td>
<td>62 (22)</td>
</tr>
<tr>
<td>Regurgitant volume (ml)</td>
<td>60 (47)</td>
<td>57 (44)</td>
</tr>
</tbody>
</table>

*1-96 SD of difference between first and second determination.
occurred in the severity of regurgitation during a change in posture. When patients were divided according to their underlying disease, however, we found significant differences in regurgitation that were dependent on posture. When failure of cusp-apation was due to mitral valve prolapse and ruptured chordae moving from the supine to the upright position tended to increase regurgitation, probably because the defective leaflet prolapsed further into the left atrium (the two patients with ruptured chordae also had some degree of valve prolapse). In patients with secondary mitral valve disease, however, the corresponding posture induced change in left ventricular volume caused a decrease in regurgitation because of diminished dilatation of the ventricle in the upright position.

Both forward and total stroke volume and hence regurgitant flow through the mitral valve altered similarly with posture changes. These findings confirm the previously reported impact of reducing ventricular volume size on the severity of regurgitation. Volume changes, however, were of the same magnitude throughout the cardiac cycle, leaving relative measures, including the left ventricular ejection fraction and regurgitation fraction, unaffected. The small increments in heart rate and diastolic blood pressure when our patients changed posture from supine to sitting upright were also indistinguishable from those reported in normal subjects.

In healthy subjects left ventricular end diastolic volume increases during mild to moderate exercise while the end systolic volume decreases. The resultant increase in stroke volume is most pronounced at lower levels of work. The cardiovascular response to exercise in cardiac disease has been most thoroughly investigated in patients with coronary artery disease. In most of these patients the left ventricle dilatates in response to exercise in the upright position, but, in contrast with the changes in healthy subjects, a dilatation is recorded both at end diastole and end systole, resulting in a smaller exercise induced increase or even a decrease in the left ventricular ejection fraction. Dehmer et al found that exercise induced end systolic dilatation was seen only in patients with disease of more than one vessel. Although the left ventricular ejection fraction was only slightly increased from rest to exercise in the 10 patients in our study, the forward stroke volume increased more obviously during exercise, the increase being considerably larger than that in total stroke volume.

The relation between preload and afterload of the left ventricle is probably particularly important for the severity of mitral regurgitation. In normal subjects the increase in left ventricular filling pressure as indicated by the pulmonary artery wedge pressure during exercise is more pronounced than the increase in systemic blood pressure. Similar findings were recorded in patients with coronary artery disease with and without moderate heart failure. Stevenson et al examined 10 patients with advanced heart failure (New York Heart Association functional class III to IV) and chronic mitral regurgitation using a combination of the thermodilution and radionuclide technique. They did not find any significant changes in either forward or total stroke volume during exercise before treatment, indicating that the increase in cardiac output during exercise was entirely attributable to a rise in heart rate. After tailored vasodilatation the exercise induced increase in cardiac output was also a result of a rise in forward stroke volume. Thus, exercise induced haemodynamic changes in patients with mitral regurgitation without signs of overt heart failure are similar to those recorded in normal subjects, and the severity of regurgitation is reduced during exercise.

The reproducibility of the determination of the regurgitation fraction using the combined radionuclide first pass and equilibrium technique was found to be high. The repeatability in terms of day to day variation was of the same magnitude as the hour to hour variation, the standard deviations of the difference being about 5%. A small but significant difference was found in the two observers' measurements of the left ventricular ejection fraction, but the limits of agreement corresponded to

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**Table 4** Observer variation in mean (SD) values of haemodynamic variables at rest in 20 supine patients with mitral regurgitation

<table>
<thead>
<tr>
<th>Observer 1</th>
<th>Observer 2</th>
<th>Limits of agreement*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Left ventricular end diastolic volume (ml)</td>
<td>212 (79)</td>
<td>210 (77)</td>
</tr>
<tr>
<td>Left ventricular ejection fraction (%)</td>
<td>59 (15)</td>
<td>57 (14)</td>
</tr>
<tr>
<td>Forward stroke volume (ml)</td>
<td>57 (16)</td>
<td>55 (16)</td>
</tr>
</tbody>
</table>

*1.96 SD of difference between observer 1 and observer 2. 95% confidence interval of bias −3 to 0.
those previously reported from our laboratory, with a standard deviation of the difference of 2.9-6. Although radionuclide cardiography may not be the most widespread procedure for determining cardiac performance, it is highly reproducible, even in the upright position and during exercise.29 30 In contrast with the combined radionuclide technique, measurements of mitral regurgitation from methods based on completely different principles, such as the thermodilution and radionuclide techniques in combination, may be influenced by tricuspid regurgitation.31 In addition, we used a radionuclide method to determine the regurgitation fraction, which has the advantage of reducing one of the key problems inherited in nuclear imaging—the subtraction of background activity—in that identical procedures for subtracting background are used in determining the forward and the total stroke volume.3 We emphasise that multigated cardiography is extremely troublesome in patients with cardiac arrhythmias and that the results of our study therefore apply primarily to patients with sinus rhythm. Thus, the findings of our study cannot uncritically be extrapolated to patients with atrial fibrillation, which is present in roughly 30% of patients with pro- mised mitral regurgitation.13

In conclusion, left ventricular volumes are reduced in patients with clinically stable chronic mitral regurgitation when they change from lying supine to sitting upright. In patients with mitral valve prolapse and ruptured chordae regurgitation increases when sitting, whereas regurgitation is decreased in patients with ischaemic heart disease and dilated cardiomyopathy. Exercise in the upright position causes an increase in forward stroke volume and a decrease in the regurgit- ant volume, leading to a reduced regurgitation fraction. These results indicate that the severity of the disease is not worsened by exercise, provided that no symptoms are provoked.

17 Manyari DE, Kostuk WP, Purves PP. Left and right ven- tricular function at rest and during bicycle exercise in the supine and sitting positions in normal subjects and patients with coronary artery disease. Am J Cardiol 1983;51:36-42.
23 Manyari DE, Kostuk WP, Purves PP. Left and right ven- tricular function at rest and during bicycle exercise in the supine and sitting positions in normal subjects and patients with coronary artery disease. Am J Cardiol 1983; 51:36-42.
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