Right ventricular diastolic abnormalities in systemic sclerosis. Relation to left ventricular involvement and pulmonary hypertension

Anna Giunta, Enrico Tirri, Stefania Maione, Sara Cangianiello, Alessandro Mele, Amalia De Luca, Gabriele Valentini

Abstract

Objectives—To investigate right ventricular diastolic function in systemic sclerosis (SSc) and its relation to clinical features of the disease.

Methods—Seventy seven unselected SSc patients and 33 healthy subjects were submitted to echocardiography and echo Doppler study to assess left and right systolic as well diastolic function and to estimate maximal arterial systolic pulmonary pressure (PAP). In addition, the patients were investigated to define the SSc subset and the extent of skin and internal organ involvement.

Results—An abnormal right ventricular filling, as expressed by an inverted tricuspid (Tr) E/A ratio (Tr E/A ratio <1), was detected in 31 of the 77 SSc patients (40%) and in 0 of the 36 controls (p<0.001). All the 31 patients with an inverted Tr E/A ratio were found to have a PAP > 30 mm Hg. Twenty resulted to have an inverted mitral (Mit) E/A ratio (Mit E/A ratio <1), indicating an abnormal left ventricular filling. In multiple regression analysis, Tr E/A ratio resulted to be independently correlated to both PAP (r=−0.35;p<0.003) and Mit E/A ratio (r=0.39;p<0.001).

Conclusions—This study points out an impaired right ventricular filling in a significant percentage of SSc patients whatever the subset. This alteration is independently correlated to both PAP and left ventricular filling abnormalities.

Systemic sclerosis (SSc) is a multisystem disorder of connective tissue characterised by widespread vascular lesions and fibrosis of the skin and distinct internal organs.12

Some authors7–9 and we ourselves10 have pointed out an impaired left ventricular filling in a significant percentage of SSc patients in whom no other cause of altered diastolic function had been detected. Diastolic abnormalities in SSc are likely to depend on either myocardial fibrosis or myocardial ischaemia, or both.1–6 As myocardial fibrosis as well as small intramyocardial coronary vessel involvement are known to affect both left and right ventricles in SSc,7–8 an altered right ventricular filling is likely to occur in this disease. Nevertheless, such aspect has received little attention so far. In this study, we have investigated right ventricular diastolic function in SSc and its relation to clinical features of the disease.

Methods

PATIENTS

Seventy seven unselected SSc patients (74 women and three men, aged from 24 to 79 years, median 56, mean (SD) 53.4 (11.6)) admitted to the Institute of Clinical Medicine and Rheumatology of the 2nd University of Naples were studied. All of them were investigated by history, clinical examination and instrumental investigations to define the disease duration, the subset according to Giordano et al and the severity of various disease manifestations (that is, general conditions, peripheral vascular involvement, skin sclerosis, joint/tendons, muscle, gut, lung, heart and kidney involvement), which were scored from 0 (absent) to 4 (end stage) according to the scleroderma severity index developed by Medsger et al.10

CONTROLS

Thirty six subjects (33 women, three men), (aged from 28 to 75 years, median age 50, mean (SD) 49.6 (10.3)) without any past or present evidence of heart and/or lung disease acted as controls.

ECHOCARDIOGRAPHY AND ECHO DOPPLER STUDY

All examinations were obtained with a phased array system (HP), with a 2.5 or 3.5 MHz transducer. Complete 2D echocardiography as well as pulsed and continuous Doppler examination were performed in a standard manner.11 12

The left ventricular ejection fraction (LVEF) was used as an index of left ventricular systolic pump function and was calculated by a modification of the method of Quinones et al.13 The percentage of shortening in right ventricle (RV) area during the systole (per cent fractional area shortening) (FAS%) was used as an index of RV systolic function and was calculated, in
Right ventricular diastolic abnormalities in systemic sclerosis

RV inflow velocities were assessed by the parasternal short axis view at the level of the tricuspid valve with the sampling window placed at the tricuspid anulus. We chose this validated\textsuperscript{11} approach because most of our SSc patients are thin women in whom keeping the transducer in good contact with the skin at the apical window is hampered by the narrow intercostal spaces. According to Nishimura et al.,\textsuperscript{11} the following diastolic parameters were measured: tricuspid (Tr) peak early inflow velocity (Tr peak E), peak late velocity (Tr Peak A), their ratio (Tr E/A) and deceleration time (Tr DT). As the RV diastolic filling is affected by respiration, measurement of beats was timed with respiration. In particular, at least three beats from the end inspiration and three beats from the end expiration were recorded, and their values were averaged.

LV inflow velocities were measured from the apical four chamber views, with the sample volume placed at the level of the leaflets tips of the mitral valve.\textsuperscript{11} The following LV filling parameters were measured: mitral (Mit) peak E, Mit peak A velocities, Mit E/A ratio and Mit DT. Moreover, isovolumic relaxation time of the left ventricle (LV-IVRT) was also assessed as the time interval elapsing from the aortic valve closure to the mitral valve opening. Each value was obtained as the mean of the measures detected during at least three cardiac cycles.

Systolic pulmonary arterial pressure (PAP) was estimated when a tricuspidal regurgitation was detected by continuous wave Doppler echocardiogram. It was calculated by measuring the peak systolic pressure gradient across the tricuspid valve (peak regurgitant velocity) and adding the estimated right atrial pressure (10 mm Hg). In absence of a tricuspidal regurgitation, PAP was considered normal.

**STATISTICAL ANALYSIS**

Unpaired Student’s $t$ test, $\chi^2$ with Yates’s correction, Fisher’s exact test, Spearman’s correlation and multiple regression analysis were used when appropriate. All data were expressed as mean (SD); a $p$ value $<0.05$ was considered statistically significant.

**Results**

Of the 77 patients, 23 were affected with limited cutaneous SSc (lc SSc), 38 with intermediate cutaneous SSc (ic SSc), 16 with diffuse cutaneous SSc (dc SSc). The disease duration ranged from 1 to 48 years, median 18, mean (SD) 17.3 (9.0).

The above mentioned Scleroderma Severity Index considers nine items: any alteration of each of them (score 1–4) was detected in the following percentages: peripheral vascular system (77 of 77, 100%); skin sclerosis (77 of 77, 100%); joint/tendons (18 of 77, 23%); muscles (6 of 77, 7%); gastrointestinal tract (65 of 77, 84%); lung (51 of 77, 66%); heart (17 of 77, 22%); kidney (2 of 77, 2%); general conditions (12 of 77, 15%). It is worth noting, that heart involvement scoring in this index is only evaluated on the basis of clinically evident cardiac manifestations, including arrhythmias and congestive heart failure, ECG, conduction defects, and LVEF.

Table 1 shows right and left echocardiographic and echo Doppler indices of systolic and diastolic function in SSc patients and controls. No significant difference was detected between patients and controls in any index of systolic function. However, two patients were found to have a LVEF <50% and three patients were found to have a FAS <30%—that is, below the lower normal values.

Table 1. Echographic and echo Doppler indices (mean (SD)) of right and left systolic and diastolic function in 77 SSc patients and 36 controls

<table>
<thead>
<tr>
<th></th>
<th>Patients (n=77)</th>
<th>Controls (n=36)</th>
<th>$p$ Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>FAS (%)</td>
<td>40.3 (3.8)</td>
<td>41.2 (2.9)</td>
<td>NS</td>
</tr>
<tr>
<td>Tr peakE (cm/s)</td>
<td>41.4 (14.4)</td>
<td>45.1 (8.5)</td>
<td>NS</td>
</tr>
<tr>
<td>Tr peakA (cm/s)</td>
<td>37.9 (15.2)</td>
<td>36.5 (9.7)</td>
<td>NS</td>
</tr>
<tr>
<td>Tr E/A</td>
<td>1.2 (0.4)</td>
<td>1.2 (0.2)</td>
<td>NS</td>
</tr>
<tr>
<td>Tr DT (ms)</td>
<td>190.5 (37.0)</td>
<td>185.3 (14.1)</td>
<td>NS</td>
</tr>
<tr>
<td>LVEF (%)</td>
<td>59.9 (4.0)</td>
<td>60.0 (3.6)</td>
<td>NS</td>
</tr>
<tr>
<td>Mit peakE (cm/s)</td>
<td>65.7 (17.0)</td>
<td>70.0 (8.5)</td>
<td>NS</td>
</tr>
<tr>
<td>Mit peakA (cm/s)</td>
<td>57.7 (17.4)</td>
<td>46.6 (8.8)</td>
<td>NS</td>
</tr>
<tr>
<td>Mit E/A</td>
<td>1.2 (0.5)</td>
<td>1.5 (0.1)</td>
<td>NS</td>
</tr>
<tr>
<td>Mit DT (ms)</td>
<td>172.4 (28.7)</td>
<td>159.4 (10.0)</td>
<td>NS</td>
</tr>
<tr>
<td>LV-IVRT (ms)</td>
<td>78.5 (1.3)</td>
<td>59.3 (0.9)</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

FAS = per cent fractional area shortening of the right ventricle; Tr peak E = tricuspidal peak early inflow velocity; Tr peak A = tricuspidal late velocity; Tr E/A = tricuspidal E/A ratio; Tr DT = tricuspidal deceleration time; LVEF = left ventricular ejection fraction; Mit peak E = mitral peak early inflow velocity; Mit peak A = mitral flow late velocity; Mit E/A = mitral E/A ratio; Mit DT = mitral deceleration time; LV-IVRT = left ventricle isovolumic relaxation time.
Table 2 shows echo Doppler parameters of left and right ventricular filling (mean (SD)) in controls and in SSc patients with (group I) and without (group II) an inverted tricuspidal E/A ratio.

Table 2  Echo Doppler parameters of left and right ventricular filling (mean (SD)) in controls and in SSc patients with (group I) and without (group II) an inverted tricuspidal E/A ratio

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Group I (n=31)</th>
<th>Group II (n=46)</th>
<th>Controls (n=36)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tr peak E (cm/s)</td>
<td>37.0 (12.4)§*</td>
<td>43.9 (14.6)</td>
<td>45.1 (8.5)</td>
</tr>
<tr>
<td>Tr peak A (cm/s)</td>
<td>48.1 (17.0)**</td>
<td>31.1 (8.7)</td>
<td>36.5 (9.7)</td>
</tr>
<tr>
<td>Tr E/A</td>
<td>0.8 (0.1)†***</td>
<td>1.4 (0.3)</td>
<td>1.2 (0.2)</td>
</tr>
<tr>
<td>Tr DT (ms)</td>
<td>188.2 (34.0)</td>
<td>188.8 (36.5)</td>
<td>185.3 (14.1)</td>
</tr>
<tr>
<td>Mit peak E (cm/s)</td>
<td>59.9 (17.2)§*</td>
<td>69.7 (15.9)</td>
<td>70.0 (8.3)</td>
</tr>
<tr>
<td>Mit peak A (cm/s)</td>
<td>63.9 (18.2)***</td>
<td>52.3 (14.1)**</td>
<td>46.6 (8.8)</td>
</tr>
<tr>
<td>Mit E/A</td>
<td>1.0 (0.3)***</td>
<td>1.4 (0.5)</td>
<td>1.5 (0.1)</td>
</tr>
<tr>
<td>Mit DT (ms)</td>
<td>173.5 (30.5)†</td>
<td>170.5 (27.1)§</td>
<td>159.4 (10.0)</td>
</tr>
<tr>
<td>IVRT (ms)</td>
<td>82.3 (13.5)††</td>
<td>76.3 (10.1)§§</td>
<td>90.3 (5.9)</td>
</tr>
<tr>
<td>PAP (mm Hg)</td>
<td>38.3 (9.8)†***</td>
<td>32.9 (5.3)</td>
<td>26.0 (5.3)</td>
</tr>
</tbody>
</table>

‡Number of patients in whom pulmonary artery pressure could be estimated. *p<0.05 versus group II. †p<0.01 versus group II. ‡Number of patients in whom pulmonary artery pressure could be estimated. *p<0.05 versus controls. **p<0.01 versus group II patients. ***p<0.001 versus controls. IVRT = isovolumic relaxation time; PAP = systolic pulmonary arterial pressure. Other abbreviations as Table 1.

Pulmonary hypertension (that is, an estimated maximal PAP >30 mm Hg) was detected in 31 of 31 (100%) of group I patients and in 18 of 46 (39%) group II patients (p<0.0001) and in 0 of 36 controls (p<0.0001). The prevalence of pulmonary hypertension was also statistically greater in group II patients than in controls (p<0.0001). Moreover, a PAP >40 mm Hg was found in 17 of 31 group I and five of 46 group II patients (p<0.0001) and a PAP >50 mmHg was detected in 14 of 31 group I and one of 46 group II patients (p<0.0001).

Tr E/A ratio (dependent variable) was found to be inversely correlated to PAP (r=−0.41; p<0.0001) and positively correlated to Mit E/A ratio (r=−0.35; p<0.0001). These correlations were then investigated in multiple regression analysis in the 70 SSc patients in whom PAP could have been valued. TrE/A ratio resulted to be independently correlated to both Mit E/A ratio (r=0.39; p<0.001) and PAP (r=−0.35; p<0.003). No correlation was found between Mit E/A ratio and PAP.

Of the 31 patients with an inverted Tr E/A ratio, all presented a PAP >30 mm Hg and 17 also impaired left ventricular filling (that is, a Mit E/A ratio <1).

No correlation was found between altered right ventricular filling and any epidemiological (sex, age, disease duration), or clinical parameters of the disease—that is, either the presence or the severity score of the nine organ system considered. In particular, the Tr E/A ratio did not correlate with either forced vital capacity or diffusing lung capacity for CO (r=0.151, p=0.278; r=0.058, p=0.736, respectively). Moreover, no difference was detected in the prevalence of an abnormal right ventricular filling in patients belonging to different SSc subsets. As a matter of fact, a Tr E/A ratio <1 was detected in eight of 16 patients with dc SSc (50%); in 13 of the 38 patients with ic SSc (33%) and in 10 of the 23 patients with lc SSc (43%). Moreover, no difference was detected in the mean Tr E/A ratio among the three subsets: 1.1 (0.4) in dc SSc; 1.2 (0.4) in ic SSc; 1.1 (0.4) in lc SSc respectively (p>0.05).

Our series is characterised by a very low number of men (n=3). As the exclusion of the three male patients did not affect the results, we chose to include them in the analysis.

Discussion

We have investigated right ventricular diastolic function in SSc and its relation to clinical and epidemiological features of the disease. We discovered an abnormal right ventricular filling in 40% of SSc patients. Such alteration was detected in many patients without clinically evident cardiac disease and resulted to be correlated with both left ventricular diastolic abnormalities and pulmonary hypertension.

Scleroderma heart disease is historically subclassified into primary and secondary.7 Primary SSc cardiac disease depends on the involvement of myocardium and/or pericardium and/or small intramyocardial vessels by SSc heart disease itself; secondary cardiac involvement develops either in patients with systemic arterial hypertension mainly induced by renal scleroderma (left ventricular disease) or in those with vascular and/or interstitial lung disease (right ventricular disease).

The pathological hallmark of SSc heart disease is myocardial fibrosis.13-14 Myocardial fibrosis and myocardial ischaemia have long been known to affect ventricular filling.15-17 SSc myocardial fibrosis is different from that occurring in patients with coronary atherosclerosis.
Actually, SSC myocardial fibrosis is equally distributed throughout the right and left ventricles, does not involve the immediate subendocardial layers, is not related to the distribution of the epicardial coronary vessels and is not associated with haemosiderin deposits.7 10–21 The detection of left ventricular filling abnormalities in SSC patients6 8 led us to study right ventricular filling in this disease. Such an aspect has received little attention so far. In point of fact, no contribution has been made in that regard except for that by Candell-Riera et al.,22 who detected a significantly low Tr E/A ratio in 63 patients with lc SSC.

In our study, we have found right diastolic abnormalities in 31 of 77 patients (40%), with no difference in either the prevalence of an inverted Tr E/A ratio or in the Tr E/A ratio values among patients belonging to different subsets.

In our SSC patients with an abnormal right ventricular filling, we detected a significant increase in Tr peak A, suggestive of abnormal relaxation.23 This pattern is classically caused by ventricular hypertrophy (that is, secondary to hypertension or aortic stenosis, etc) in which prominent atrial contraction is important for ventricular filling. A similar pattern, however, has also been observed in patients with cardiac amyloidosis with no or little evidence of increased right ventricular free wall thickness. It was ascribed to early amyloid infiltration possibly interfering with the relaxation process.24 It was ascribed to early amyloid infiltration possibly interfering with the relaxation process.24

Nevertheless, in our whole SSc series. Moreover, excluding three male patients from the analysis did not change the results. In conclusion, we have found an impaired right ventricular filling in patients with SSC. Such alteration is detectable in patients without any clear cut evidence of cardiac disease. Nevertheless, its prognostic significance awaits to be defined.


Right ventricular diastolic abnormalities in systemic sclerosis. Relation to left ventricular involvement and pulmonary hypertension

Anna Giunta, Enrico Tirri, Stefania Maione, Sara Cangianiello, Alessandro Mele, Amalia De Luca and Gabriele Valentini

doi: 10.1136/ard.59.2.94