Cardiac function in borderline hypothyroidism: a study by pulsed wave tissue Doppler imaging

Sandra Zoncu, Francesca Pigliaru, Claudia Putzu, Lorella Pisano, Sara Vargiu, Martino Deidda, Stefano Mariotti and Giuseppe Mercuro

Departments of Cardiovascular Sciences and Endocrinology, Department of Medical Sciences, University of Cagliari, Sardinia, Italy

(Correspondence should be addressed to G Mercuro; Department of Cardiovascular Sciences, University of Cagliari, S.S. 554, bivio di Sestu, 09042 Monserrato, Cagliari, Italy; Email: mercuro@pacs.unica.it)

Abstract

Objective: In subclinical hypothyroidism (SH), impaired diastolic function has been documented at rest and on effort, while systolic dysfunction has only been assessed on effort.

Design: The aim of the present study was: (a) to further assess systolic function at rest in SH; and (b) to ascertain whether cardiac dysfunction could precede TSH increase in euthyroid patients with a high risk of developing SH.

Methods: We studied 32 patients with classical Hashimoto’s thyroiditis (22 with increased serum TSH (>3 mU/ml – group A), and 10 with normal serum TSH (<3 mU/ml – group B)); a third group (C), which included 13 healthy controls. All subjects underwent pulsed wave tissue Doppler imaging (PWTDI) to accurately quantify the global and regional left ventricular function.

Results: When compared with group C, PWTDI indices showed that in both groups A and B there was a significant impairment of systolic ejection (P<0.001 and P<0.05, respectively), a delay in diastolic relaxation (P<0.001 and P<0.05, respectively) and a decrease in the compliance to the ventricular filling (P<0.05). Several significant correlations were found between PWTDI parameters and serum-free T3 and T4 and TSH concentrations.

Conclusion: PWTDI is a sensitive technique that allows detection of both diastolic and systolic abnormalities, not only in patients with SH, but also in euthyroid subjects with a high risk of developing thyroid failure. Furthermore, the significant correlations of several PWTDI indices with serum FT3 and TSH concentrations strongly support the concept of a continuum spectrum of a slight thyroid failure in autoimmune thyroiditis extending to subjects with serum TSH still within the normal range.

European Journal of Endocrinology 152 527–533

Introduction

Subclinical hypothyroidism (SH) is a frequent condition defined by elevated thyroid stimulating hormone (TSH) secretion in the presence of normal concentrations of circulating thyroid hormones (1–3). Most cases of SH are due to the slow progression of thyroid failure caused by autoimmune thyroiditis and it is generally believed that the majority of patients with SH will eventually become frankly hypothyroid (2). Clinical symptoms and signs are often non-specific, and diagnosis and monitoring of therapy depend crucially on measurements of thyroid hormones and TSH in the blood (3–5). On the other hand, patients with unequivocal evidence of autoimmune thyroiditis may report symptoms (mainly chronic fatigue) even in the presence of normal serum TSH concentrations: the term ‘subclinical hypothyroidism’ has been suggested for this condition (6), but this remains open to speculation (7).

The clinical relevance of SH is still unclear, in spite of several reports of an increased risk of low density lipoprotein (LDL)-cholesterol and negative influence on the hemostatic profile, and of increased occurrence of coronary and peripheral arterial disease and depression (1–4). Several mild cardiac abnormalities, such as impairment of left ventricular diastolic function at rest and of systolic function on effort have been described in SH (8–12). These anomalies are probably responsible for a wide spectrum of symptoms suggestive of thyroid failure observed in patients with SH (13).

In the present study, taking advantage of the peculiar sensitivity of the heart to the action of the thyroid hormone, we assessed diastolic and systolic function in patients with SH and in subjects with euthyroid
autoimmune thyroiditis with normal serum TSH concentration, but at risk of developing thyroid failure. For this purpose, we used pulsed wave tissue Doppler imaging (PWTDI), a technique based on the Doppler principle, which is able to precisely assess the ventricular wall motion (14).

Subjects and methods

Study population

We studied 32 patients (31 females, one male) with documented classical Hashimoto’s thyroiditis (see below for the diagnostic criteria), whose main features are reported in Table 1. Twenty-two of them had SH with increased serum TSH (>3 mU/ml: 5.40±2.08 mU/ml, range 3.31–10.7 mU/ml; group A), and 10 had normal serum TSH (<3 mU/ml: 2.27±0.52 mU/ml, range 1.20–2.89 mU/ml; group B). Serum-free T3 (FT3) concentration was within the normal range in both groups. A third group included 13 healthy euthyroid controls (12 females, one male; group C). These were subjects who came to our laboratory for a thyroid function test and were found within normal values. All groups were matched for age, sex and body mass index (BMI) (Table 1). No significant differences were observed among the groups in heart rate, blood pressure and serum FT3 concentration, but at risk of developing thyroid failure.

Table 1 Biophysical characteristics, cardiovascular measurements and hormonal data of patients and controls.

<table>
<thead>
<tr>
<th>Patients</th>
<th>Group A</th>
<th>Group B</th>
<th>Group C</th>
</tr>
</thead>
<tbody>
<tr>
<td>TSH (mU/l)</td>
<td>3 mU/l</td>
<td>&lt;3 mU/l</td>
<td>&lt;3 mU/l</td>
</tr>
<tr>
<td>(n = 22)</td>
<td>(n = 10)</td>
<td>(n = 13)</td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td>42.5±9.9</td>
<td>40.9±1.9</td>
<td>39.0±8.2</td>
</tr>
<tr>
<td>Male/female</td>
<td>1/21</td>
<td>0/10</td>
<td>1/12</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>64.1±10.6</td>
<td>64.6±10.9</td>
<td>62.1±11.4</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>158.2±5.4</td>
<td>160.1±4.4</td>
<td>162.6±10.5</td>
</tr>
<tr>
<td>BSA (m²)</td>
<td>1.6±0.1</td>
<td>1.7±0.1</td>
<td>1.6±0.2</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>25.6±4.2</td>
<td>25.1±4.6</td>
<td>23.5±3.7</td>
</tr>
<tr>
<td>HR (bpm)</td>
<td>69.9±9.5</td>
<td>72.0±13.3</td>
<td>76.2±11.6</td>
</tr>
<tr>
<td>SBP (mmHg)</td>
<td>126.4±12.7</td>
<td>124.8±19.2</td>
<td>122.2±7.1</td>
</tr>
<tr>
<td>DBP (mmHg)</td>
<td>80.5±7.7</td>
<td>76.1±6.5</td>
<td>77.8±7.1</td>
</tr>
<tr>
<td>FT3 (pg/ml)</td>
<td>3.2±0.5</td>
<td>3.2±0.4</td>
<td>3.5±0.2</td>
</tr>
<tr>
<td>FT4 (pg/ml)</td>
<td>7.8±1.7</td>
<td>10.2±1.4</td>
<td>10.2±1.7</td>
</tr>
<tr>
<td>TSH (mU/l)</td>
<td>5.4±2.1</td>
<td>2.3±0.5</td>
<td>1.2±0.5</td>
</tr>
</tbody>
</table>

*P < 0.01 vs Group C; **P < 0.01 vs Group B and Group C.
BMI, body mass index; BSA, body surface area; DBP, diastolic blood pressure; HR, heart rate; SBP, systolic blood pressure. Values are given as means±S.D.

Study protocol

The Ethical Committee of our University approved the present study, and informed written consent was obtained from all subjects. Participants were familiarized with instrumentation and medical environment of echocardiographic laboratory before testing. All subjects underwent physical examination, 12-lead electrocardiogram and M-mode, 2D and Doppler-echocardiography. Finally, subjects underwent PWTDI study, using an ultrasound system equipped with tissue Doppler imaging capabilities (SSA-380A; Toshiba Corp., Toshigi, Japan). A single experienced echocardiographer, who was unaware of the subjects’ thyroid conditions, carried out all examinations with subjects in the left lateral decubitus position. A simultaneous electrocardiographic tracing was also obtained.

Doppler echocardiography

A complete M-Mode, 2D, spectral- and color-Doppler recordings were performed using a 2.5-MHz transducer. Left ventricular mass (LVM) was measured according to the Devereux method (15): LVM index (LVMI) was obtained by dividing LVM by body surface area. Ejection fraction (EF) was calculated using the Simpson method (16). Pulsed Doppler transmitral flow velocities were recorded from the 4-chamber apical view, with the sample volume placed at the level of the mitral valve leaflet tips. Early (Em) and late (Am) diastolic velocities of transmitral flow were measured and the Em/Am ratio was derived. Isovolumic relaxation time (IVRT) was measured as the time interval between the end of systolic output flow and transmitral Em wave onset, by placing the sample volume between outflow tract and the mitral valve.

PWTDI

PWTDI uses the Doppler principle to assess the ventricular wall motion velocity by positioning the sample volume within the myocardium (14). In PWTDI, low-amplitude blood-flow signals are eliminated by gain adjustment to allow only high-amplitude Doppler signals from wall motion to enter the velocity calculation circuit. By means of a 4-chamber apical view, a mm sample volume was placed both at the level of basal lateral and infero-septal mitral annulus, with the ultrasonic Doppler beam in a position as parallel as possible to the motion of the myocardial wall. Three distinct waves were obtained through PWTDI in each of the two mitral annular sites during the cardiac cycle: a systolic wave (S); an early diastolic wave (E) and a late diastolic wave (A). The following measurements were made from the PWTDI recordings: peak systolic velocity (Sₐ, cm/sec), acceleration time of S (ATSₐ, sec), deceleration time of S (DTSₐ, sec), peak early diastolic velocity (Eₐ, cm/sec), peak late diastolic velocity
In comparison with controls, E_m and E_m/Am ratio was then calculated. Mean acceleration and deceleration rates of S (ARS_s, cm/sec^2; DRS_s, cm/sec^2) and of E (ARE_m, cm/sec^2; DRE_m, cm/sec^2) were obtained as S_a and E_a divided by their respective time intervals. These measurements were repeated on recordings of three consecutive cardiac cycles and their mean value was obtained. Then, values of basal lateral and interseptal mitral annulus were averaged. Finally, we considered the peak early diastolic velocity of lateral mitral annulus (E al, cm/sec) as a relaxation index. We then calculated the ratio between E_m and E al (Em/E al), a parameter that takes into account the influence of cardiac relaxation on mitral flow (17). Reproducibility of PWTDI parameters in our laboratory had been previously documented (18).

Assessment of thyroid function

Serum TSH, FT_3, and FT_4, anti-thyroglobulin and anti-thyroid peroxidase antibodies (TgAb, TPOAb) were measured in all subjects. FT_4 and FT_3 were assayed by a direct method with chromatographic separation in Lisophase columns (Technogenetics, Milano, Italy). The normal ranges for FT_4 and FT_3 were 6.6–16 pg/ml and 2.8–5.6 pg/ml, respectively. Serum TSH was measured by immunochemiluminescent assay (Ortho-Clinical Diagnostic, Amersham, UK, normal range 0.3–3.0 mU/l). TPOAb were determined by RIA (ICN; normal range <10 UI/ml; TgAb by passive haemagglutination (Fujirebio Inc. Pharmaceuticals, Tokyo, Japan: normal range <1/100). TPOAb ranged 64–3000 UI/ml (median 369) in group A, 134–3000 UI/ml (median 379) in group B, and were negative in controls. Thyroid ultrasound was performed in all cases using a 7.5 mHz linear electronic transducer, and thyroid echogenicity was subjectively evaluated by a conventional gray scale. The diagnosis of Hashimoto’s thyroiditis was made on the basis of hypoechoic goiter (19) associated to high levels of TPOAb and/or TgAb (20, 21).

Statistical analysis

Data from all groups are reported as mean ± S.D. Differences between control subjects and patients were assessed by the Student’s 2-tailed t-test for unpaired observations. Correlations between echocardiographic parameters and PWTDI indices, and hormones (FT_3, FT_4 and TSH) were evaluated using Pearson’s correlation coefficient. P values < 0.05 were considered to be significant.

Results

Doppler echocardiography

In comparison with controls, E_m and E_m/Am ratio showed a significant decrease (P < 0.05 and P < 0.001, respectively), and IVRT a significant prolongation (P < 0.001) in group A (Table 2). Conversely, these parameters remained unmodified in group B. On the other hand, none of the echocardiographic parameters showed relevant differences between patients of group A and B. No single parameter of conventional echocardiography was correlated to serum TSH or FT_3 concentration, with the exception of E_m/Am ratio, which showed a significant inverse correlation (r = –0.36, P < 0.01) with serum TSH.

PWTDI systolic function

PWTDI systolic indices analysis revealed a significant impairment of the systolic ejection in both groups A and B, in comparison with group C, as shown by a decreased S_a and DRS_a (P < 0.001 in group A, P < 0.05 in group B). Moreover, patients of group A showed an increase of TS_a and DTS_a (P < 0.05), and a decrease of ARS_a and DRS_a in respect to group C (P < 0.001). None of the systolic PWTDI indices showed significant differences between groups A and B. Table 3 shows PWTDI systolic indices in all the groups.

Table 4 details correlations between PWTDI systolic indices and serum FT_3, FT_4 and TSH when patients of groups A, B and C were analyzed together. Taken collectively, the results showed significant correlations between most systolic indices and serum FT_3 or TSH, whereas, with the exception of DRS_a, no correlation was found with serum FT_4. Some of these correlations are depicted in Fig. 1.

PWTDI diastolic function

PWTDI diastolic recordings analysis revealed a significant impairment of both diastolic relaxation (decreased E_a, ARE_a, DRE_a and E al, P < 0.001) and compliance to the ventricular filling (decreased E_m/A_m ratio, P < 0.001, and increased E_m/E_a ratio, P < 0.05) in...
groups A and B as compared with group C. None of the diastolic PWTDI indices showed significant differences between Groups A and B. Table 5 displays PWTDI diastolic indices in all the three groups.

Table 6 shows correlations between PWTDI diastolic indices and serum FT3, FT4 and TSH when patients of groups A, B, and C were analyzed together. Again, in keeping with the important role of this hormone in determining cardiac function, most diastolic parameters were significantly associated to FT3. Surprisingly, three diastolic indices (Ea/Aa, Eal, Em/Eal) were strongly correlated to serum FT4 and TSH but not to serum FT3, thus suggesting a different sensitivity to circulating T3 and T4 concentrations. The correlations between serum TSH and selected PWTDI diastolic parameters are depicted in Fig. 2.

**Discussion**

In the present study we carried out an extensive investigation of cardiac function in patients with borderline or mild thyroid failure using PWTDI, a method of wall motion-velocity assessment for quantifying the dynamics of the cardiac cycle. The results obtained confirmed the presence of a diastolic dysfunction in patients with SH and provided further evidence that an impairment of systolic function is present in mild thyroid failure, and also at rest. Moreover, our results provided the first evidence that a subtle impairment in both

---

Table 3 Systolic mitral annulus indices assessed by PWTDI.

<table>
<thead>
<tr>
<th>Patients</th>
<th>Group A</th>
<th>Group B</th>
<th>Group C</th>
</tr>
</thead>
<tbody>
<tr>
<td>TSH &lt; 3 mIU/l</td>
<td>TSH &gt; 3 mIU/l</td>
<td>n = 22</td>
<td>n = 10</td>
</tr>
<tr>
<td>Sa (cm/sec)</td>
<td>8.68±1.08</td>
<td>9.03±1.54</td>
<td>10.65±1.37</td>
</tr>
<tr>
<td>TSa (sec)</td>
<td>0.29±0.02</td>
<td>0.28±0.01</td>
<td>0.27±0.02</td>
</tr>
<tr>
<td>ATSa (sec)</td>
<td>0.06±0.01</td>
<td>0.05±0.01</td>
<td>0.05±0.01</td>
</tr>
<tr>
<td>DTSa (sec)</td>
<td>0.23±0.03</td>
<td>0.22±0.01</td>
<td>0.21±0.04</td>
</tr>
<tr>
<td>ARSa (cm/sec²)</td>
<td>155.74±41.35</td>
<td>176.55±56.37</td>
<td>227.13±113.58</td>
</tr>
<tr>
<td>DRSa (cm/sec²)</td>
<td>37.85±5.67</td>
<td>40.58±7.57</td>
<td>53.62±19.26</td>
</tr>
</tbody>
</table>

*P < 0.05, **P < 0.001 vs Group C.

ARSa: mean acceleration of Sa; ATSa: acceleration time of Sa; DRSa: mean deceleration of Sa; DTSa: deceleration time of Sa; Sa: peak systolic velocity of mitral annulus; TSa: ATSa + DTSa. Values are given as means±S.D.

Table 4 Correlations between PWTDI systolic indices and thyroid hormones in all subjects of Groups A, B and C.

<table>
<thead>
<tr>
<th>FT3</th>
<th>FT4</th>
<th>TSH</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sa</td>
<td>0.37</td>
<td>0.28</td>
</tr>
<tr>
<td>TSa</td>
<td>−0.49</td>
<td>0.15</td>
</tr>
<tr>
<td>ATSa</td>
<td>0.06</td>
<td>−0.06</td>
</tr>
<tr>
<td>DTSa</td>
<td>−0.47</td>
<td>−0.20</td>
</tr>
<tr>
<td>ARSa</td>
<td>0.12</td>
<td>0.20</td>
</tr>
<tr>
<td>DRSa</td>
<td>0.53</td>
<td>0.30</td>
</tr>
</tbody>
</table>

For abbreviations see Table 3.

Figure 1 Selected correlations between plasma TSH concentrations and PWTDI systolic indices. □ TSH > 3 mIU/l patients (group A); ○ TSH < 3 mIU/l patients (group B); ● control subjects (group C).
Eal (cm/sec) 12.65
DREa (cm/sec²) 146.50
Ea (cm/sec) 10.99
DTEa (sec) 0.08
Ea/Aa 1.04

ATEa: acceleration time of Ea; DREa: mean deceleration of Ea; DTEa:
transmitral flow, e.g. Em/Am, which are influenced by
phy indices of left ventricular filling assessed by Doppler
tial confounding effects of conventional echocardiogra-
wedge pressure (17). This approach avoided the poten-
Eal: early diastolic peak velocity of lateral mitral annulus; E m: mitral early
peak velocity. Values are given as means ± S.D.

AREa 0.41
DREa 0.46
Ea/Aa 0.29
Em/Eal 0.30

*P < 0.05, **P < 0.001 vs Group C.

As far as the diastolic function is concerned, our
results with PWTDI are in keeping with several pre-
vious studies (for extensive reviews see (10, 11)) carried
out by conventional echocardiography, documenting
the presence of a significant diastolic impairment in
patients with SH. With respect to conventional echocar-
diography, PWTDI allowed the use of Eal, a relaxa-
tion index independent from the preload, and of
Em/Eal, a filling index closely correlated with the
wedge pressure (17). This approach avoided the poten-
tial confounding effects of conventional echocardiogra-
phy indices of left ventricular filling assessed by Doppler
transmitral flow, e.g. Em/Am, which are influenced by
several confounding factors such as age, heart rate,
preload and afterload (22).

In contrast with the consistent findings on diastolic
function, the literature provided conflicting data on sys-
tolic function in patients with SH. Biondi et al. (8)
reported a significant reduction of the mean aortic ac-
celeration, whereas peak aortic flow velocity, cardiac
output, fractional shortening, and mean velocity of cir-
cumferential fiber shortening remained unaltered (8).

An impairment of systolic function was observed by
means of the systolic time interval, which shows the
limits of an indirect method, inferred by the integration
of electrocardiographic and Doppler data (9, 12). Systolic
abnormalities were detected with certainty in SH only in
the course of physical exercise, when the evaluation of
the cardiac performance is favored by the discrepancy
between the energy request and the functional cardiac
reserve (8, 9, 12). In this context, SH has been found to
be associated with a deranged cardio-respiratory func-
tion, both at the level of anaerobic threshold and at
peak exercise (23). Evidence suggesting systolic abnor-
malities in SH has recently been reported by other tech-
niques such as ultrasonic myocardial textural analysis
(24), impedance cardiography (25), and radionuclide
ventriculography (26).

By means of recordings performed at several sites of
the mitral annulus, the PWTDI technique produces
an average figure of the structure displacement velocity,
which represents a reliable index of global systolic func-
tion (27). This methodological approach allowed us to
demonstrate that an impairment of the systolic phase
is also present at rest in patients with SH.

On account of the assumption that PWTDI is the
most sensitive technique to assess the effects of subtle
thyroid failure on heart contractility, we decided to
use it to test the possibility that patients with auto-
immune thyroiditis and normal TSH levels (i.e. those
with a high risk of developing SH or overt thyroid fail-
ure) may actually be slightly hypothyroid; this con-
dition has been recently envisaged (6, 28) but, to our
knowledge, no investigation (and particularly no
studies on heart contractility) has been carried out to
directly address the question.

Unlike conventional echocardiography, which did not
show any significant alteration in both systolic and
diastolic function of patients with ‘euthyroid’ auto-
immune thyroiditis, PWTDI allowed us to detect a
subtle but significant impairment in the whole cardiac
function of these individuals. In detail, our findings
were consistent in showing a significant impairment of
left ventricular ejection (decreased S a and DRS a),
diastolic relaxation (decreased E a, AREa, DREa and Eal)
and ventricular filling (decreased Eal/Aa ratio and in-
creased Em/Eal).

The degree of derangement of the above parameters
found in patients with ‘euthyroid’ autoimmune thy-
roiditis was always intermediate between patients
with SH and normal euthyroid controls.

These results, along with the significant correlations
of a number of PWTDI indices with thyroid hormones

---

**Table 5** Diastolic mitral annulus indexes assessed by PWTDI.

<table>
<thead>
<tr>
<th>Patients</th>
<th>Group A</th>
<th>Group B</th>
<th>Group C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>TSH &gt; 3 mU/l</td>
<td>TSH &lt; 3 mU/l</td>
<td>TSH &lt; 3 mU/l</td>
</tr>
<tr>
<td></td>
<td>n = 22</td>
<td>n = 10</td>
<td>n = 13</td>
</tr>
<tr>
<td>Ea (cm/sec)</td>
<td>10.99±2.35**</td>
<td>12.10±2.58**</td>
<td>15.21±2.5</td>
</tr>
<tr>
<td>Aa (cm/sec)</td>
<td>10.84±1.94</td>
<td>11.34±2.15</td>
<td>10.38±1.72</td>
</tr>
<tr>
<td>E2/Ea</td>
<td>1.04±0.28**</td>
<td>1.13±0.37**</td>
<td>1.53±0.47</td>
</tr>
<tr>
<td>ATEa (sec)</td>
<td>0.05±0.00</td>
<td>0.05±0.01</td>
<td>0.18±0.01</td>
</tr>
<tr>
<td>DTEa (sec)</td>
<td>0.08±0.01*</td>
<td>0.08±0.02</td>
<td>0.08±0.01</td>
</tr>
<tr>
<td>AREa (cm/sec²)</td>
<td>226.19±57.83**</td>
<td>244.29±59.42**</td>
<td>336.25±71.15</td>
</tr>
<tr>
<td>DREa (cm/sec²)</td>
<td>146.50±37.01**</td>
<td>166.45±44.45**</td>
<td>212.30±40.63</td>
</tr>
<tr>
<td>Em (cm/sec)</td>
<td>12.65±3.30**</td>
<td>13.42±3.20**</td>
<td>17.94±3.36</td>
</tr>
<tr>
<td>Em/Eal</td>
<td>5.00±1.28*</td>
<td>4.85±1.12*</td>
<td>4.63±0.95*</td>
</tr>
</tbody>
</table>

*P < 0.05, **P < 0.001 vs Group C.

---

**Table 6** Correlations between PWTDI systolic indices and thyroid hormones in all subjects of Group A, B and C.

<table>
<thead>
<tr>
<th>FT3</th>
<th>FT4</th>
<th>TSH</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

For abbreviations see Table 5.
and TSH concentrations, strongly support the concept of a continuum spectrum of a slight thyroid failure in autoimmune thyroiditis extending to subjects with serum TSH still within the normal range. The majority of PWTDI indices were correlated to serum FT3 rather than FT4 concentrations. This proves the pivotal role of T3 in cardiac function. Moreover, some diastolic indices such as Ea/Aa, Eal and Em/Eal were significantly correlated to FT4 but not to FT3. The reason for such findings is unclear and cannot be derived from the present data. We are inclined to think that some indices may be more affected by local generation of T3 (from circulating T4) than by circulating T3. Indeed, autoimmune thyroid disease is so common in the population (up to 40% of women with lymphocytic infiltration of the thyroid and 10–15% with thyroid autoantibodies) (1–4), that laboratory reference ranges derived from apparently healthy subjects could easily be affected by diseased individuals. In keeping with this concept, increased prevalence of TPOAb has been found in subjects with serum TSH concentrations outside the narrow range 0.2–1.9 mU/l, providing evidence that TSH in the upper reference range is often associated with abnormal pathology in the thyroid (7).

In conclusion, by showing an impaired systolic function in the basal condition of patients with SH, the tissue Doppler imaging has extended our knowledge of the cardiac involvement in this disease. However, the clinical relevance of this finding and the effects of L-thyroxine substitution therapy remain matters for further investigation.

Acknowledgements

The authors thank Enrico Lampis, MA, for his technical assistance in preparing the manuscript.

References

3 Surks MI, Ortiz E, Daniels GH, Sawin CT, Col NF , Cobin RH, Franklyn JA, Hershman JM, Burman KD, Denke MA, Gorman C, Cooper RS & Weissman NJ. Subclinical thyroid disease: scientific review and guidelines for diagnosis and management. JAMA 2004 291 228–238.
7 Dayan C. Whose normal thyroid function is better – yours or mine? Lancet 2002 360 353–354.
9 Monzani F, Di Bello V, Caraccio N, Bertini A, Giorgi D, Giusti C & Ferrannini E. Effect of levothyroxine on cardiac function and structure in subclinical hypothyroidism: a double blind,
23 Kahaly GJ. Cardiovascular and atherogenic aspects of subclinical hypothyroidism. Thyroid 2000 10 665–679.