Clinical, ECG and echocardiographic clues to the diagnosis of TTR-related cardiomyopathy

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ABSTRACT
Background: Signs of cardiac transthyretin (TTR) amyloidosis (ATTR) in patients with echocardiographic increase in interventricular septal thickness (IVST) are lacking.

Objectives: To identify clinical and ECG/echocardiographic signs associated with increased IVST in ATTR.

Methods: Analysis of patients with baseline echocardiography in the Transthyretin Amyloidosis Outcomes Survey (THAOS) registry (N=1682). Patients were categorised into IVST classes according to the American Society of Echocardiography classification adapted to gender (ie, normal, mild, moderate, severe); then into two combined IVST classes (normal-mild and moderate-severe).

Results: 425 patients were included: 336 with a TTR mutation (m-TTR) and 89 with wild-type TTR (WT-TTR). 72% were men. Median (25th, 75th centile) age was 62 (45, 72) years. Non-Val30Met and WT-TTR were frequent in moderate (41% and 35%) and severe (50% and 33%) IVST classes. Median IVST was 15 mm (14, 16) (moderate) and 20 mm (18, 22) (severe). In the combined moderate-severe class, 85% of patients were ≥55 years of age; 81% were men; 86% had blood pressure <140 mm Hg; and 77% had increased right ventricle thickness (≥7 mm). Up to 66% of patients had cardiac sparkling. Systolic dysfunction (left ventricular ejection fraction <50%), restrictive pattern and low voltage were less frequent, and observed in 49%, 18% and 33% of patients, respectively.

Conclusions: Increased IVST, especially in men ≥55 years with normal systolic blood pressure, increase in right ventricle free wall and valve thicknesses, and sparkling, should alert practitioners to the possibility of ATTR. Absence of restrictive pattern and low voltage should not rule out the suspicion.

Trial registration number: NCT00628745 (clinicaltrials.gov).

KEY QUESTIONS

What is already known about this subject?
- Amyloidosis is a result of continuous accumulation of insoluble fibril proteins in the extracellular matrix in various organs including the heart. The two major proteins involved in this process are immunoglobulin light chains (AL) and transthyretin (TTR). Cardiac amyloid infiltration is the classic form of infiltrative hypertrophic cardiomyopathy (HCM) and is associated with abnormal increased interventricular septal thickening.

What does this study add?
- We investigated and identified the clinical and echocardiographic signs that should alert practitioners to the possibility of TTR cardiac amyloidosis in patients with abnormal increased IVST. In future, additional research using new sensitive imaging techniques and including the other major type of cardiac amyloidosis, that is, light chain amyloidosis, and other types of HCM, should be undertaken to identify and validate specific diagnostic markers of the different types of cardiac amyloidosis.

How might this impact on clinical practice?
- Cardiac amyloidosis is of poor prognosis and should be considered in the differential diagnosis of patients with abnormal increased IVST. When suspected, patients should be referred for genetic testing and/or other imaging evaluation (MRI and bone scintigraphy), and/or biopsy analysis should be identified.

INTRODUCTION

Recently, the European Society of Cardiology (ESC) published a position statement concerning hypertrophic cardiomyopathy, to raise clinicians’ awareness of the possible spectrum of abnormalities in patients, and eventually their families, with heart muscle diseases. Cardiac hypertrophy is generally identified by echocardiography and defined by an increase in interventricular septal thickness (IVST) and/or the left ventricular (LV) posterior wall thickness (PWT), with different cut-off values according to gender. This definition has been endorsed in the 2014 ESC guidelines. However, IVST

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increase alone cannot provide accurate information concerning the aetiology of heart disease. Hence, there is a need to develop appropriate diagnostic strategies based on clues from medical and family histories, physical examination, and non-invasive investigations such as ECG and echocardiography. Cardiac hypertrophy is now defined as a morphological increase in thickness of the LV wall that results from different conditions: cardiomyocyte hypertrophy or infiltration of either cardiomyocytes or the extracellular matrix.\(^1\) Cardiac transthyretin amyloidosis (ATTR) is a classic form of an infiltrative cardiomyopathy. It is a progressive lethal disease caused by continuous accumulation of insoluble TTR fibrils in the extracellular matrix.\(^3\)

The hereditary form of ATTR expresses different phenotypes involving neuropathy and/or cardiomyopathy.\(^5\) More than 100 mutations have been reported.\(^7\) The Val30Met mutation is by far the most frequent mutation reported to cause ATTR, with two distinguishable forms depending on the age of onset: early (<50 years) and late (≥50 years).\(^6\) Early-onset Val30Met exhibits neuropathy as the major presenting feature and is endemic in Portugal, Japan and Brazil.\(^9\) Late-onset Val30Met patients exhibit neurological as well as cardiac abnormalities.\(^6\) Patients with non-Val30Met mutations were also reported to have cardiomyopathy and/or neuropathy.\(^4\)

Wild-type (WT) ATTR, also known as senile systemic amyloidosis, usually affects the heart and primarily affects older male patients, with a prevalence of 25–36% observed in an autopsy series of elderly patients.\(^9\)

Regardless of the genotype, identifying signs that may alert clinicians to the presence of ATTR cardiomyopathy are vital, as typical therapies for heart failure are not well tolerated; emerging treatments are designed to prevent progression of amyloid infiltration and prognosis is directly related to cardiac dysfunction, and is poor for later-stage disease.\(^10\)

Accordingly, the aim of this study was to identify clinical, echocardiographic and ECG parameters associated with IVST increase in cardiac ATTR using the largest available international cohort (the Transthyretin Amyloidosis Outcomes Survey (THAOS)) on this disease to raise clinician’s awareness of the diagnosis of cardiac ATTR.

**MATERIALS AND METHODS**

**Patient population and data collection**

The THAOS registry began enrolling patients in 2007. Details of this registry have been previously published.\(^11\)\(^12\)

Data were obtained from the information given by the reporting centres, using a standardised dedicated website. Data entered into the database included, but were not limited to, demographics, clinical status, medical history, type of neuropathy when present, ECG and echocardiographic data, and TTR genotype.\(^11\)\(^12\)

All the participating clinical sites had received authorisation from their local ethical committees for use of the registry. Patients entered in the registry were required to be ≥18 years of age. Privacy and confidentiality of participating individuals were assured and each individual gave written consent.

**Data extraction**

By 30 June 2013, THAOS included data on 1682 patients, registered from 17 countries. This study reports the data of the 425 patients included in THAOS with a baseline echocardiography and IVST measurement in the range (3–40 mm), prespecified in the registry. Three patients with baseline echocardiography data were excluded due to out-of-range IVST values.

**Definition of the interventricular septal thickness and genotype classifications**

Patients were divided according to the IVST classification set by the American Society of Echocardiography.\(^2\)\(^3\) As stated, gender was also taken into consideration to limit misclassification.\(^2\)\(^3\) Classification was as follows: normal IVST group if thickness ≤9 mm in female subjects or ≤10 mm in male subjects; mildly abnormal if ≥10 and ≤12 mm in females, or ≥11 and ≤13 mm in males; moderately abnormal if ≥13 and ≤15 mm in females, or ≥14 and ≤16 mm in males;\(^2\)\(^3\) and severely abnormal if ≥16 mm in females or ≥17 mm in males.\(^2\)\(^3\) In a subsequent analysis, the normal/mild groups and moderate/severe groups were pooled to calculate clinical, ECG and echocardiographic prevalence signs associated with marked IVST increase.

Patients were also divided into four categories depending on ATTR genotype and onset: Val30Met early onset (<50 years), Val30Met late onset (≥50 years), non-Val30Met and WT-ATTR.

**Definition of ECG and echocardiographic variables recorded**

ECG analysis and measurements included heart rate, PR interval, QRS duration, QT interval, atrial fibrillation, pathological Q waves and low voltage. The latter was defined as QRS voltage amplitude <0.5 mV in all limb leads or <1 mV in all precordial leads.

Echocardiographic measurements were obtained in accordance with the American Society of Echocardiography recommendations. Wall and valve thickness (IVST; PWT; RV free-wall thickness; mitral, aortic and tricuspid valvular thickening) as well as LV and LA size (LV end-diastolic diameter (LVEDD) and left atrial diameter (LAD)) were measured on transthoracic echocardiograms. LV ejection fraction (LVEF) was measured using Simpson’s biplane method. Peak E and A wave velocities were measured on Doppler mitral flow. Systolic pulmonary artery pressure (syst PAP) was estimated from the modified Bernoulli equation.\(^13\) Peak Ea velocity was measured on tissue Doppler images recorded at the annular mitral valve. Mitral regurgitation (MR) was quantified by measuring the MR area of the
colour Doppler signal divided by the left atrial area measured in the four-chamber view.\textsuperscript{14} MR was considered ‘mild or less’ if MR area/LA area was <10\%, ‘severe’ if MR area/LA area was >40\% and ‘moderate’ if between the two.

Definition of selected clinical, ECG and echocardiographic classes

Patients were divided according to baseline symptoms. Symptoms were defined as ‘any symptom classified as possibly or definitely related to TTR amyloidosis as they were recorded in medical history or general examination in the THAOS registry’. To study the prevalence of potential signs associated with cardiac ATTR, clinical, ECG and echocardiographic characteristics were defined as follows: advanced age if \( \geq 55\) years, high systolic blood pressure (SBP) if \( \geq 140\) mm Hg,\textsuperscript{15} abnormal LVEF if \(<50\%\),\textsuperscript{16} hypertrophied posterior wall if PWT >13 mm for women and \( \geq 14\) mm for men,\textsuperscript{4} asymmetric hypertrophy if IVST/PWT ratio >1.3,\textsuperscript{17} restrictive filling pattern if E/A transmitral flow velocities >2\textsuperscript{18} and increased RV free-wall if RV thickness \( \geq 7\) mm.\textsuperscript{19}

Statistical analysis

Continuous variables are reported as median, and 25th and 75th centiles. For nominal qualitative data, n and percentage were reported. Statistical differences between patient groups were calculated using \( \chi^2 \) test for categorical variables. For continuous variables, the Mann-Whitney test was used when two groups were compared and the Kruskal-Wallis test when more than two groups were compared. Statistical analysis was performed using SAS software; \( p<0.05 \) was considered significant.

RESULTS

Baseline characteristics of the echocardiographic THAOS subpopulation

The median (25th and 75th centile) age of the study population of 425 patients was 62 (45, 72) years and 73\% were men. Figure 1 presents the flow chart of the study, and prevalence of men and women, according to the IVST classification. A total of 336 patients had a diagnosis of m-ATTR and 89 of WT-ATTR (table 1). Val30Met mutation alone accounted for 50\% of the m-ATTR subpopulation, of whom 104 were in early-onset and 63 were in late-onset groups (table 1).

Clinical, biological and ECG characteristics of the THAOS echo-subgroup according to the IVST classes

Baseline demographic, genetic, echocardiographic, ECG and biological parameters with respect to the four IVST classes, are shown in table 1. Briefly, moderate or severe IVST was observed more frequently in non-Val30Met and in WT-TTR amyloidosis. Of the early Val30Met patient group, 90\% had normal or mild increased IVST.

Older age, male gender, cardiac symptoms, history of heart failure, higher brain natriuretic peptide (BNP) values, conduction abnormalities, low voltage and pathological Q waves, were more frequent in the moderate and severe IVST classes. No differences were observed in body mass index (BMI), modified BMI (mBMI), heart rate, troponin I and T values, and there was equal prevalence of pacemaker implant between the different IVST classes.

Echocardiographic characteristics of the THAOS echo-subgroup according to the IVST classes

Baseline echocardiographic data according to the IVST classification are summarised in table 2. Increased IVST...
Severity was associated with higher sparkling, decreased LVEF, prevalence of moderate to severe MR, valvular thickening, increased LAD and increased syst PAP. E/A was significantly different in the four IVST classes, and was more elevated in patients with normal and severe IVST. In all groups, the median of E/A was <1.5.

The prevalence of clinical, echocardiographic and ECG abnormal signs in patients with respect to IVST classes is shown in Figure 2. The most frequent signs (>60%) are dyspnoea (New York Heart Association (NYHA) II–IV), age >55 years, male gender, LAD enlargement, increased RV free wall thickness, LV dysfunction measured by ejection fraction, and myocardial sparkling. IVST/PWT ratio >1.3, a sign of asymmetric septal hypertrophy encountered most commonly in sarcomeric cardiomyopathy, was reported in 21%; whereas low voltage, believed to be a strong sign of ATTR, was observed in only a third of these patients.

Detailed comparisons of clinical, echocardiographic and ECG abnormal signs between the normal-mild IVST

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**Table 1** Comparison of baseline clinical, biological and ECG variables among patients with normal, mild, moderate and severe IVST

| Variables                  | N  | Normal | Mild | Moderate | Severe | p Value  |
|----------------------------|----|--------|------|----------|--------|----------|
| N                          | 425| 80     | 93   | 81       | 171    |          |
| Genetic                    |    |        |      |          |        |          |
| ATTR FAP early Val30Met, n (%) | 104| 50 (63) | 44 (47) | 8 (10) | 2 (1)  | <0.0001  |
| ATTR FAP late Val30Met, n (%) | 63 | 7 (9)  | 17 (18) | 12 (15) | 27 (16) |          |
| ATTR FAP non-Val30Met, n (%) | 169| 22 (28) | 29 (31) | 33 (41) | 85 (50) |          |
| Wild type, n (%)            | 89 | 1 (1)  | 3 (3) | 28 (35) | 57 (33) |          |

Demographic and clinical

| Variable                   | N  | Normal | Mild | Moderate | Severe |
|----------------------------|----|--------|------|----------|--------|
| Age, years                 | 425| 35 (31.50) | 49 (39.64) | 67 (58.76) | 69 (61.75) | <0.0001 |
| Male, n (%)                | 425| 52 (65) | 55 (59) | 62 (77) | 143 (84) | <0.0001 |
| Pacemaker, n (%)           | 221| 2 (12)  | 6 (21) | 12 (24) | 35 (28) | 0.4524  |
| Myocardial infarction, n (%) | 421| 0 (0)  | 0 (0) | 2 (3)  | 3 (2)  | 0.3007  |
| Heart failure, n (%)       | 421| 8 (10)  | 16 (18) | 45 (56) | 124 (73) | <0.0001 |
| Palpitations, n (%)        | 421| 6 (8)   | 5 (6)  | 11 (14) | 32 (19) | 0.0081  |
| Syncope, n (%)             | 421| 5 (6)   | 5 (6)  | 12 (15) | 32 (19) | 0.0044  |
| BMI, kg/m²                 | 376| 24 (21.26) | 24 (22.27) | 25 (22.27) | 25 (23.27) | 0.4255 |
| Karnofsky index, %         | 360| 90 (80.90) | 90 (80.90) | 80 (70.90) | 75 (60.80) | <0.0001 |
| NYHA II–IV vs I, n (%)     | 190| 6 (75)†  | 12 (75) | 40 (89) | 112 (93) | 0.8666 |
| SBP, mm Hg                 | 374| 121 (110,130) | 122 (114,132) | 120 (105,130) | 112 (104,130) | 0.0025 |
| DBP, mm Hg                 | 374| 77 (70.84) | 78 (71.83) | 70 (66.80) | 70 (65.80) | 0.0002  |
| Age at onset, years        | 423| 31 (27.44) | 43 (33.57) | 60 (49.70) | 62 (53.70) | <0.0001 |
| Liver transplant, n (%)    | 425| 16 (20) | 11 (12) | 11 (14) | 12 (7) | 0.2689  |
| Age at liver transplant, years | 50 | 40 (35.52) | 44 (32.54) | 55 (48.61) | 50 (39.60) | 0.0675 |
| Motor neuropathy, n (%)    | 425| 31 (39) | 33 (36) | 33 (41) | 69 (40) | 0.8688  |
| Sensory neuropathy, n (%)  | 425| 69 (86) | 79 (85) | 54 (67) | 92 (54) | <0.0001 |
| Autonomic neuropathy, n (%) | 425| 59 (74) | 59 (63) | 45 (56) | 98 (57) | 0.0529  |

Biological

| Variable                   | N  | Normal | Mild | Moderate | Severe |
|----------------------------|----|--------|------|----------|--------|
| BNP, pg/mL                 | 141| 32 (14.83) | 72 (37.179) | 591 (206,1027) | 552 (216,939) | 0.0031 |
| NT-proBNP, pg/mL           | 152| 55 (27.108) | 100 (47.345) | 939 (330,3445) | 3507 (1658,8982) | 0.4320 |
| Troponin I, ng/mL          | 37 | 0.06 (0.06,0.06) | 0.06 (0.02,0.1) | 0.06 (0.02,0.09) | 0.12 (0.07,0.15) | 0.1329 |
| Troponin T, ng/mL          | 122| 0.01 (0.01,0.015) | 0.01 (0.01,0.04) | 0.04 (0.02,0.06) | 0.05 (0.03,0.08) | 0.1828 |

ECG

| Variable                   | N  | Normal | Mild | Moderate | Severe |
|----------------------------|----|--------|------|----------|--------|
| Heart rate, bpm            | 349| 74 (68.85) | 74 (65.78) | 72 (64.82) | 74 (65.85) | 0.6633 |
| PR interval, ms            | 236| 166 (148,188) | 161 (149,200) | 178 (152,208) | 193 (176,220) | 0.0011 |
| QRS interval, ms           | 294| 94 (86,100) | 98 (88,106) | 110 (88,128) | 116 (100,148) | <0.0001 |
| QT interval, ms            | 280| 378 (360,397) | 394 (361,413) | 424 (390,461) | 438 (398,470) | <0.0001 |
| Low voltage, n (%)         | 265| 2 (4)  | 5 (9)  | 20 (39) | 34 (30) | <0.0001 |
| Pathologic Q waves, n (%)  | 229| 3 (17) | 2 (7)  | 17 (31) | 51 (41) | 0.0013 |

*Only TTR FAP was included in the analysis (wild type excluded).
†Data for this parameter were not reported in 90% of the patients of this group. Continuous variables are presented as median (25th, 75th centile). Percentage indicates proportion of patients with the variable within each IVST category.

ATTR, transthyretin amyloidosis; BMI, body mass index; BNP, brain natriuretic peptide; DBP, diastolic blood pressure; FAP, familial amyloid polyneuropathy; IVST, interventricular septal thickness; NT-proBNP, N-terminal proBNP; NYHA, New York Heart Association; SBP, systolic blood pressure.
and moderate-severe IVST groups are shown in Table 3. Briefly, patients with moderate to severe IVST were older, more symptomatic, male gender, and presented low voltage and Q waves, LV systolic dysfunction, increased filling pressure (Ea >15), and left atrial enlargement, and had more myocardial sparkling and valvular thickening, and increased RV wall and PWTs, than normal to mild IVST patients.

Table 2 Comparison of baseline echocardiographic characteristics among patients according to IVST classification

| IVST               | N  | Normal          | Mild            | Moderate        | Severe          | p Value   |
|--------------------|----|-----------------|-----------------|-----------------|-----------------|-----------|
| IVST, mm           | 425| 9 (8, 10)       | 11 (10, 12)     | 15 (14, 16)     | 20 (18, 22)     | –         |
| PWT, mm            | 403| 9 (8, 9)        | 10 (9, 11)      | 14 (12, 15)     | 18 (16, 20)     | <0.0001   |
| LVEDD, mm          | 379| 45 (42, 48)     | 46 (42, 49)     | 45 (41, 48)     | 43 (39, 47)     | 0.0129    |
| Sparkling, n (%)   | 299| 17 (33)         | 23 (38)         | 30 (53)         | 93 (72)         | <0.0001   |
| LVEF, %            | 327| 62 (56, 70)     | 60 (55, 67)     | 55 (45, 60)     | 43 (35, 60)     | <0.0001   |
| RV wall thickness, mm | 87 | 6 (5, 7)        | 6 (5, 7)        | 8 (6, 9)        | 9 (7, 10)       | 0.0086    |
| Mitral thickening, n (%) | 228 | 5 (14)        | 4 (9)           | 14 (33)         | 51 (48)         | <0.0001   |
| Aortic thickening, n (%) | 225 | 6 (16)        | 9 (21)          | 12 (29)         | 42 (40)         | 0.0184    |
| Tricuspid thickening, n (%) | 221 | 0 (0)         | 2 (5)           | 2 (5)           | 18 (18)         | 0.0031    |
| LAD, mm            | 359| 33 (30, 35)     | 36 (33, 40)     | 42 (39, 46)     | 45 (42, 49)     | <0.0001   |
| Syst PAP, mm Hg    | 140| 24 (19, 25)     | 25 (20,28)      | 30 (26, 40)     | 40 (30,45)      | <0.0001   |
| E, cm/s            | 139| 74 (62, 85)     | 70 (63, 81)     | 88 (65, 102)    | 81 (67, 101)    | 0.0107    |
| A, cm/s            | 114| 54 (48, 65)     | 64 (56, 75)     | 70 (56, 104)    | 54 (40, 81)     | 0.0048    |
| E/A                | 109| 1.3 (1, 1.7)    | 1.1 (0.9, 1.2)  | 0.9 (0.7, 1.6)  | 1.3 (0.9, 2.4)  | 0.0390    |
| E/Ea, mean±SD      | 60 | 7.3±4.0         | 7.8±4.3         | 14.4±6.7        | 16.8±6.7        | 0.0001    |

Continuous variables are presented as median (25th, 75th percentile). Percentage indicates proportion of the patients with the variable within each IVST category.

IVST, interventricular septal thickness; LAD, left atrial diameter; LVEDD, left ventricular end-diastolic diameter; LVEF, left ventricular ejection fraction; PWT, posterior wall thickness; RV, right ventricle; syst PAP, systolic pulmonary arterial pressure.

Figure 2 Prevalence of the clinical, ECG and echocardiographic signs in the group with moderate and severe IVST increase in the THAOS population with echocardiography data available at baseline (N=252). *Women >13 mm, men >14 mm (IVST, interventricular septal thickness; n, number of patients with individual sign; N, number of patients with evaluations available for individual sign; NYHA, New York Heart Association; PWT, posterior wall thickness, RV, right ventricle; SBP, systolic blood pressure; THAOS, Transthyretin Amyloidosis Outcomes Survey).
Prevalence of clinical, echocardiographic and ECG abnormal signs in the moderate and severe group, according to genotype and disease onset

Of the overall early-onset and late-onset Val30Met, non-Val30Met and WT-ATTR, 10%, 62%, 70% and 96%, had moderate or severe IVST, respectively. Percentages of signs depending on genotype and/or disease onset for Val30Met in the moderate to severe IVST group are presented in table 4. Briefly, the prevalence of age ≥55 years and male gender were higher in the WT versus other ATTR groups (table 4). Neuropathy was more frequent in late Val30Met compared with non-Val30Met and WT patients. Hypertension or asymmetric ‘hypertrophy’ was more frequently observed in late Val30Met than in patients with non-Val30Met and WT-ATTR. Prevalence of dyspnoea, Q waves, left atrial dilation and sparkling was similar in the late Val30Met, non-Val30Met and WT-ATTR groups (table 4). However, it is important to note that no specific clinical, ECG or echocardiographic signs were exclusively associated with a given ATTR genotype.

**DISCUSSION**

This study identified clinical, ECG and echocardiographic features that should alert cardiologists to suspect amyloidosis in patients with increased IVST, using the THAOS registry.

First, we demonstrated that advanced age, male gender, dyspnoea, increased RV free-wall thickness and echocardiographic myocardial sparkling, were frequently associated with moderate to severe IVST in patients with mostly normal SBP <140 mm Hg. Second, we showed that asymmetric LV ‘hypertrophy’ and absence of trans-mitral restrictive pattern or low voltage should not rule out a diagnosis of ATTR in these patients; and third, we pointed out that these methods could not discriminate the different TTR genotypes.

**Interest of a large international registry**

The THAOS registry represents the largest available database to document transversal ATTR characteristics and offers the unique opportunity to analyse data according to different genotypes and phenotypes. The international nature of the study allows for the generalisation of the findings, as it includes data from various countries and centres. A further advantage of such a large database is the use of standardised forms and the uniform manner in which data from a large number of patients with different genotypes are collected.

**IVST measured by echocardiography as a useful tool to screen symptomatic patients for ATTR**

Increased wall thickness of the heart is frequently diagnosed by echocardiography in patients; however, it could result from a number of different conditions. Progress

### Table 3: Prevalence of indicators in symptomatic subjects by IVST classification in patients classified in normal-mild and moderate-severe IVST classes

| IVST                  | N   | Normal-mild (n=173) | Moderate-severe (n=252) | p Value |
|-----------------------|-----|---------------------|-------------------------|---------|
| **Clinical**          |     |                     |                         |         |
| Age ≥55 years, n (%)  | 425 | 56 (32)             | 214 (85)                | <0.0001 |
| NYHA II–IV, n (%)     | 190 | 18 (75)             | 152 (92)                | 0.0134  |
| Male, n (%)           | 425 | 107 (62)            | 205 (81)                | <0.0001 |
| SBP ≥140 mm Hg, n (%) | 374 | 19 (12)             | 30 (14)                 | 0.7141  |
| Sensory neuropathy, n (%) | 425 | 148 (86)         | 146 (58)                | <0.0001 |
| Motor neuropathy, n (%) | 425 | 64 (37)            | 102 (41)                | 0.4698  |
| **ECG**               |     |                     |                         |         |
| Low voltage, n (%)    | 265 | 7 (7)               | 54 (33)                 | <0.0001 |
| Q waves, n (%)        | 229 | 5 (10)              | 68 (38)                 | 0.0002  |
| **Echo**              |     |                     |                         |         |
| LVEF <50%, n (%)      | 327 | 18 (14)             | 98 (49)                 | <0.0001 |
| Posterior wall moderate or severe hypertension (women ≥13 mm, men ≥14 mm), n (%) | 403 | 11 (7) | 49 (21) | 0.0001 |
| IVST/PWT ratio >1.3, n (%) | 403 | 11 (7) | 49 (21) | 0.0001 |
| E/A >2, n (%)         | 109 | 5 (7)               | 7 (18)                  | 0.0705  |
| E/Ea mean >15, n (%)  | 60  | 4 (8)               | 4 (40)                  | 0.0066  |
| RV free wall thickness ≥7 mm, n (%) | 87  | 20 (42) | 30 (77) | 0.0009 |
| LAD ≥40 mm, n (%)     | 359 | 26 (18)             | 171 (79)                | <0.0001 |
| Sparkling, n (%)      | 299 | 40 (36)             | 123 (66)                | <0.0001 |
| Aortic thickening, n (%) | 225 | 15 (19) | 54 (37) | 0.0052 |
| Mitral thickening, n (%) | 228 | 9 (11)  | 65 (44) | <0.0001 |
| Tricuspid thickening, n (%) | 221 | 2 (3)  | 20 (14) | 0.0053 |

IVST, interventricular septal thickness; LAD, left atrial diameter; LVEF, left ventricular ejection fraction; NYHA, New York Heart Association; PWT, posterior wall thickness; RV, right ventricle; SBP, systolic blood pressure.
In molecular biology, it has elucidated the underlying genetic abnormalities in many cases of cardiac hypertrophy; however, in daily practice, cardiologists have access first to the phenotype of patients rather than its genetic defects. As was pointed out in the recent ESC statement concerning hypertrophic cardiomyopathy, there is still a need to develop appropriate diagnostic strategies based on clues from medical and family histories, physical examination and non-invasive investigations such as echocardiography.1 Hence, identification of clinical signs associated with specific genetic disease, such as ATTR, is an important step before performing appropriate genetic testing.

Clinical and echocardiographic signs associated with increased IVST in ATTR

In the present study, we identified signs that may provide guidance in a possible diagnosis of ATTR. We demonstrated that age >55 years and male gender are frequently associated with increased IVST ≥13 mm in female, and ≥14 mm in male patients with ATTR. WT-ATTR, non-Val30Met and late-onset Val30Met patients showed higher IVST than early Val30Met patients. This may be explained by the relationship between increased IVST and age, and male gender, as shown in a previous study.16 Accordingly, in our study, WT-ATTR, non-Val30Met and late-onset Val30Met patients were older, and more frequently male, than early Val30Met patients. However, mechanisms other than age and gender might be involved in IVST increase, such as TTR fibril composition. Unfortunately, this information is lacking in the THAOS registry. Late-onset Val30Met and WT-ATTR showed fragmented TTR protein; conversely, whole TTR protein is observed in early-onset Val30Met.20 21

The striking difference in penetrance according to gender has been reported in several previous studies.22 23 Of note, 94% of patients with WT-ATTR were men; which is in accordance with reports in the literature.24–26 In Val30Met, as described previously,22 male gender dominated among late-onset patients compared to those with early onset. To the best of our knowledge, no pathophysiological explanation for this gender discrepancy has emerged.

In our study, most patients with increased IVST had normal blood pressure and dyspnoea. Hypertension is a ubiquitous cause of LV hypertrophy, with advanced age. Therefore, the combination of increased IVST and normal blood pressure should increase cardiologists’ awareness of the possibility of cardiac amyloidosis. Conversely, in daily practice, dyspnoea might not be useful to discriminate ATTR amyloidosis from other hypertrophic cardiomyopathies. Of course, these signs are also frequent in

### Table 4

| IVST Moderate and severe | Val30Met Early onset | Val30Met Late onset | non-Val30Met | WT |
|--------------------------|----------------------|---------------------|--------------|-----|
| N                        | 10                   | 39                  | 118          | 85  |
| Clinical                 |                       |                     |              |     |
| Age ≥55 years, n (%)     | 5 (50)               | 39 (100)            | 85 (72)      | 85 (100) |
| NYHA II–IV, n (%)        | 2 (100)              | 13 (87)             | 65 (93)      | 72 (91)  |
| Gender, male, n (%)      | 6 (60)               | 31 (80)             | 88 (75)      | 80 (94)  |
| SBP ≥140 mm Hg, n (%)    | 3 (33)               | 13 (37)             | 8 (8)        | 6 (8)   |
| Sensory neuropathy, n (%)| 9 (90)               | 37 (95)             | 75 (64)      | 25 (29)  |
| Motor neuropathy, n (%)  | 8 (80)               | 30 (77)             | 57 (48)      | 7 (8)   |
| ECG                      |                       |                     |              |     |
| Low voltage, n (%)       | 1 (17)               | 2 (11)              | 27 (38)      | 24 (35)  |
| Q waves, n (%)           | 1 (17)               | 8 (44)              | 31 (38)      | 28 (38)  |
| Echocardiography         |                       |                     |              |     |
| LVEF <50%, n (%)         | 0 (0)                | 2 (15)              | 53 (51)      | 43 (55)  |
| Posterior wall hypertrophy*, n (%) | 1 (10) | 17 (53) | 80 (71) | 67 (79)  |
| IVST/PWT ratio ≥ 1.3, n (%) | 3 (30) | 13 (41) | 23 (21) | 10 (12)  |
| E/A >2, n (%)            | 0 (0)                | 1 (20)              | 3 (14)       | 3 (30)  |
| E/Ea mean >15, n (%)     | 0 (0)                | NA                  | 3 (75)       | 1 (25)  |
| RV free wall thickness ≥7 mm, n (%) | 1 (33) | 7 (88) | 14 (74) | 8 (89) |
| LAD >40 mm, n (%)        | 5 (56)               | 28 (80)             | 69 (73)      | 69 (89)  |
| Sparkling, n (%)         | 4 (67)               | 25 (74)             | 56 (65)      | 38 (62)  |
| Aortic thickening, n (%) | 1 (33)               | 2 (18)              | 33 (46)      | 18 (30)  |
| Mitral thickening, n (%) | 0 (0)                | 2 (18)              | 37 (51)      | 26 (43)  |
| Tricuspid thickening, n (%) | NA | 1 (9) | 12 (17) | 7 (12) |

*Moderate or severe hypertrophy—women >13 mm and men >14 mm; NA, no data available for tricuspid valve thickening in early-onset Val30Met and for E/Ea in late onset for Val30Met.

IVST, interventricular septal thickness; LAD, left atrial diameter; LVEF, left ventricular ejection fraction; NYHA, New York Heart Association; PWT, posterior wall thickness; RV, right ventricle; SBP, systolic blood pressure.
amyloid light-chain (AL) amyloidosis, and this diagnosis should be ruled out by identification of amyloidogenic monoclonal proteins (serum and urine immunofixation combined with free light-chain quantification) and demonstration of TTR amyloid deposition in a tissue specimen.

Increased RV wall thickness, valvular thickening and granular sparkling of the myocardium were frequently observed in patients with moderate to severe IVST, in accordance with previous findings.\textsuperscript{25,27,28} TTR infiltrates all the structures of the heart (left and right ventricle (RV) and valves) in contrast to hypertension, which only increases LV wall thickness.\textsuperscript{29} Increases in aortic valve and mitral valve thickening were frequently observed. In previous studies, atrioventricular thickening has been described as a marker for cardiac amyloidosis.\textsuperscript{28} Rapezzi \textit{et al.}\textsuperscript{60} showed recently that half of patients with ATTR exhibited abnormal valvular thickness whereas only 3% exhibited sarcomeric hypertrophic cardiomyopathies. Myocardial sparkling, a qualitative—but subjective and highly dependent on the echocardiographic technique (harmonic)—sign of amyloid infiltration, was observed in 65% of the patients with moderate to severe IVST in our study, which is in agreement with previous reports.\textsuperscript{31-33} Combination of increased RV wall thickness, valvular thickening and sparkling, are of particular relevance to the diagnosis of ATTR in patients with increased IVST. However, increased RV wall thickness can also be observed in cases of symmetric LV hypertrophy in Fabry disease or in mitochondrial cytopathy, and in cases of asymmetric LV hypertrophy in sarcomeric hypertrophic cardiomyopathy. Thus it is more the combination of signs rather than the sign alone that may suggest a diagnosis of TTR amyloidosis.

**Absence of signs should not rule out the diagnosis**
Cardiac amyloidosis is often described as a restrictive cardiomyopathy with symmetric hypertrophic pattern, preserved ejection fraction and ECG low voltage.\textsuperscript{34}

In our study, the restrictive transmitral pattern was observed in only 18% of patients with ATTR moderate to severe IVST. Asymmetric LV hypertrophy pattern was not rare, as it was exhibited in 21% of patients with moderate and severe IVST. This asymmetric pattern was more frequently observed in late-onset Val30Met and in non-Val30Met than in WT-ATTR. Forty-nine per cent of patients with moderate to severe IVST had an ejection fraction <50%. Low voltage was observed in less than one-third of the patients. All of this suggests that absence of a restrictive pattern or low voltage, or presence of asymmetric hypertrophy pattern or LV systolic dysfunction, should not rule out the diagnosis of ATTR amyloidosis.

**Study limitations**
There are several limitations in investigations based on data collected in observational surveys such as THAOS. As data were collected during routine clinical practice from many centres/countries and at the discretion of the patient’s physician, there are inevitably variations in the type of clinical investigations conducted. In particular, at the time when these data were reviewed (data cut-off date 30 June 2013), echocardiography was not considered as a routine clinical examination in many participating neurology centres. Thus, patients with primarily neurological manifestations of their disease are generally attended by neurologists and do not undergo heart examinations to a similar extent as those presenting with primarily heart problems and followed by cardiologists. Thus, there was a selection bias in investigations performed, with heart examinations rarely performed in patients with early-onset ATTR Val30Met disease. The high prevalence of heart failure (NYHA II–IV) noted in the group of patients with normal-mild increased IVST probably represents a selection bias, where patients with symptoms of dyspnoea are registered as having heart failure and undergo echocardiographic examination. Similarly, heart examinations are generally not carried out on asymptomatic carriers of an ATTR gene mutation, a group that otherwise would be interesting to analyse. Furthermore, the conclusion of this study may be applicable only for patients with the same characteristics as those included in this study.

Non-Val30Met includes many different mutations with phenotype heterogeneity that could not be addressed in this study due to the limited number of patients with echocardiography.

Echocardiographic measurements such as MR measurement by vena-contraction or proximal isovelocity surface area, or new echocardiographic techniques such as strain or strain rate, were not performed worldwide and are, therefore, not reported in the THAOS registry. Thus, although they are of interest in this disease,\textsuperscript{45} evaluation of MR severity and LV systolic function was limited. Not all echocardiography machines used were able to measure strain by Doppler tissue imaging or two-dimensional speckle-tracking, measurements may differ between brands of device. This makes it difficult to calculate a mean value.

Lastly, this study focused only on TTR amyloidosis, whereas most of the signs described are also present in AL cardiac amyloidosis. A diagnosis of AL cardiac amyloidosis should be ruled out by other means, such as demonstration of TTR amyloid deposition in a tissue specimen. Future studies are needed comparing patients with hypertensive heart disease and hypertrophic cardiomyopathy with patients having ATTR amyloidosis to determine sensitivity and specificity, and cut-off value of each sign, to predict ATTR amyloidosis.

**CONCLUSION**
TTR cardiac amyloidosis should be suspected in the presence of increased echocardiographic IVST, particularly in patients >55 years of age, of male gender, with normal SBP, increased thickness of RV free wall and
valves, LAD enlargement and granular sparkling of the myocardium. Female gender or absence of restrictive pattern, or absence of low voltage or LV dysfunction, should not rule out the possibility of cardiac ATTR. ATTR genetic testing should be performed in patients with suspected TTR cardiomyopathy, as the criteria above are not specific and were present across the different ATTR genotypes and in WT-ATTR. Further studies, including control groups, are needed to determine the specificity, sensitivity and cut-off values of the criteria described in this study.

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Ethics approval
Local institutional review boards approved the study.

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No additional data are available.

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