Progression of myocardial sympathetic denervation assessed by $^{123}$I-MIBG imaging in familial amyloid polyneuropathy and the effect of liver transplantation

Maria da Conceição Azevedo Coutinho$^{a,*}$, Nuno Cortez-Dias$^a$, Guilhermina Cantinho$^b$, Isabel Conceição$^c$, Tatiana Guimarães$^a$, Gustavo Lima da Silva$^a$, Miguel Nobre Menezes$^a$, Ana Rita Francisco$^a$, Rui Plácido$^a$, Fausto J. Pinto$^a$

$^a$ Serviço de Cardiologia, Hospital Universitário de Santa Maria, Centro Cardiovascular da Universidade de Lisboa, Centro Académico de Medicina de Lisboa, Lisboa, Portugal

$^b$ Instituto de Medicina Nuclear, Faculdade de Medicina, Universidade de Lisboa, Portugal

$^c$ Departamento de Neurociências, Hospital Universitário de Santa Maria, Lisboa, Portugal

Received 18 June 2016; accepted 11 August 2016
Available online 24 May 2017

KEYWORDS
Amyloid; Scintigraphy; Liver transplantation

Abstract

Introduction: Familial amyloid polyneuropathy (FAP) is a rare disease caused by systemic deposition of amyloidogenic variants of the transthyretin (TTR) protein. The TTR-V30M mutation is caused by the substitution of valine by methionine at position 30 and mainly affects the peripheral and autonomic nervous systems. Cardiovascular manifestations are common and are due to autonomic denervation and to amyloid deposition in the heart. Cardiac sympathetic denervation detected by iodine-123 labeled metaiodobenzylguanidine (MIBG) is an important prognostic marker in TTR-V30 M FAP. Liver transplantation, widely used to halt neurological involvement, appears to have a varying effect on the progression of amyloid cardiomyopathy. Its effect on the progression of cardiac denervation remains unknown.

Methods: In this observational study, patients with the TTR-V30 M mutation underwent annual cardiac assessment and serial MIBG imaging with quantification of the late heart-to-mediastinum (H/M) ratio.

Results: We studied 232 patients (median age 40 years, 54.7% female, 37.9% asymptomatic at the time of inclusion) who were followed for a median of 4.5 years and underwent a total of


* Corresponding author.

E-mail address: cacoutinho@sapo.pt (M.d.C. Azevedo Coutinho).

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Introduction

Familial amyloid polyneuropathy (FAP) due to a mutation in the transthyretin (TTR) protein, in which most frequently valine is replaced by methionine in position 30 (TTR-V30M), is a multisystem hereditary disease characterized by progressive involvement of sensory, motor and autonomic nerve fibers. Neurological manifestations are the main clinical feature, but cardiovascular involvement due to autonomic neuropathy and cardiac amyloid deposition is also common.

Myocardial scintigraphy with iodine-123 labeled metaiodobenzylguanidine (123I-MIBG) is a non-invasive method for quantifying cardiac sympathetic innervation.
Previous studies have demonstrated that myocardial sympathetic innervation as detected by \(^{123}\)I-MIBG is decreased in the initial stages of FAP\(^{8,9}\) and that reduction in the late heart-to-mediastinum (H/M) ratio is an important predictor of prognosis that is strongly associated with long-term risk of mortality.\(^{4}\) However, the progression of cardiac denervation during the clinical course of TTR-V30 M FAP has never been assessed by serial \(^{123}\)I-MIBG scans, and the added value of repeated measurement of late H/M for prognostic stratification is unknown.

Liver transplantation is widely used to slow the progression of FAP and until recently was the only treatment option for these patients.\(^{7}\) Transplantation halts neurological involvement and improves survival, especially if performed early in the symptomatic stages of the disease.\(^{6}\) However, amyloid deposition in the heart may continue after transplantation, leading to progressive myocardial thickening, arrhythmias and conduction disturbances.\(^{8,9,12,13}\) Little is known of the course of cardiac denervation following liver transplantation.

The aim of this study was to assess the effect of liver transplantation on the progression of cardiac denervation in patients with TTR-V30 M FAP.

**Methods**

This was a prospective observational study of consecutive patients with the TTR-V30 M mutation. All subjects underwent annual cardiac assessment and periodic measurement of the late H/M ratio. Progression of cardiac denervation was determined by comparing the H/M ratio in successive exams.

\(^{123}\)I-MIBG scintigraphy was performed following premedication with Lugol’s solution (iodine-potassium iodide) to block absorption of free \(^{123}\)I by the thyroid gland. Chronic medication was not suspended before the exam. Planar anterior thoracic images were acquired 15 min (early) and 3 h (late) after intravenous injection of a fixed dose of 185 MBq of \(^{123}\)I-MIBG. A dual-head gamma camera (Millennium, General Electric Healthcare) equipped with a low-energy, high-resolution parallel-hole collimator was used for image acquisition and a 20% energy window was centered over the 159 keV photopeak of \(^{123}\)I.

\(^{123}\)I-MIBG uptake was quantified through the H/M ratio, which was calculated by dividing the mean counts per pixel in the region of interest around the heart by the mean counts per pixel in the mediastinum, without correction for background activity. The myocardial washout rate was calculated as the percentage reduction of myocardial counts between early and late images:

\[
\text{Washout rate} = \frac{[\text{early H/M} - \text{late H/M}]}{\text{early H/M}} \times 100
\]

**Statistical analysis**

Continuous variables with normal distribution were expressed as means ± standard deviation and were compared using the Student’s t test or analysis of variance, and those with non-normal distribution were expressed as medians and interquartile range (IQR) and compared with the Mann-Whitney or Kruskal-Wallis tests. Scintigraphic parameters were correlated with age, duration of symptoms and time before and after liver transplantation, using Pearson’s and Spearman’s correlation coefficients.

The prognostic impact of scintigraphic parameters on all-cause mortality during follow-up was determined by Cox univariate and multivariate regression analysis adjusted for age. To avoid bias, stepwise methods were used for the multivariate analysis, with non-significant variables being removed from the model at each step (p-value 0.05 for inclusion and 0.10 for removal). The accuracy of predictors of mortality was assessed by the area under the curve (AUC) on receiver operating characteristic analysis for different durations of follow-up. Cumulative event rates by quartiles of H/M ratio were assessed by the Kaplan-Meier method and the log-rank test.

The statistical analysis was performed with SPSS version 21.0 (IBM SPSS, Chicago, IL). A p-value <0.05 was considered statistically significant.

**Results**

**Population characteristics**

Between September 1998 and July 2015, 305 carriers of the TTR-V30 M mutation underwent annual cardiac assessment. A total of 73 patients were excluded from the analysis: 45 for not having undergone MIBG imaging and 28 because their first MIBG scan was performed after liver transplantation. The final study population thus consisted of 232 patients, median age 40 years (IQR 32-55), 54.7% female (n=127).

At the initial cardiac assessment, 144 patients (62.1%) had documented neurological involvement, the other 88 (37.9%) being asymptomatic. Of the symptomatic patients, 99 had developed clinical manifestations by the age of 50, while the other 45 had late onset (after the age of 50). The mean age of symptom onset was 38 years (IQR 31-54) and the mean duration of symptoms at initial assessment was two years (IQR 0-3). Nineteen carriers of the pathogenic mutation aged over 50 were still symptom-free.

**\(^{123}\)I-MIBG scintigraphy**

At the initial assessment, the mean early and late H/M ratios were 1.85±0.23 and 1.83±0.03, respectively, and median washout rate was 2.5 (IQR 2.3 to 8.5). Early and late H/M ratios were significantly lower and the washout rate was significantly higher in symptomatic patients (Table 1).

During a median follow-up of 4.5 years (IQR 2.1-7.7; maximum 15.6), 121 patients underwent serial MIBG imaging, of whom 36 underwent at least five exams (maximum nine) (Supplementary Table 1). Including both initial and subsequent assessments, a total of 558 MIBG scans were performed. In these exams, the late H/M ratio decreased with age (by 0.082/year, p<0.001) and with duration of symptoms (by 0.066/year, p<0.001) (Figure 1 and Supplementary Figure 1). Figure 2 presents the progression of late H/M in serial MIBG scans of two typical patients, illustrating the slow reduction in the presymptomatic stage of the disease,
Table 1  Comparison of scintigraphic parameters according to neurological involvement.

<table>
<thead>
<tr>
<th></th>
<th>Symptomatic (n=144)</th>
<th>Asymptomatic (n=88)</th>
<th>p</th>
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<tbody>
<tr>
<td>Early H/M</td>
<td>1.75±0.33</td>
<td>2.03±0.28</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Late H/M</td>
<td>1.70±0.37</td>
<td>2.09±0.31</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Washout rate</td>
<td>3.98 (IQR 0.0 to 10.0)</td>
<td>0 (IQR -6.26 to 4.66)</td>
<td>&lt;.001</td>
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H/M: heart-to-mediastinum ratio; IQR: interquartile range.

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Figure 1  Changes in late H/M ratio by age (A) and duration of symptoms (B). H/M: heart-to-mediastinum.

Figure 2  Changes in late H/M ratio over time in two patients with TTR-V30 M familial amyloid polyneuropathy, showing reduction in the ratio with disease progression and apparent stabilization after liver transplantation. H/M: heart-to-mediastinum.

the more marked decline after symptom onset, and the apparent stabilization after liver transplantation.

Prognostic stratification

During follow-up, 47 patients (20.3%) died. The most frequent cause of death (40%) was terminal FAP (Supplementary Table 3). MIBG scintigraphy at inclusion was a strong predictor of prognosis, with the risk of death increasing by 27.8% (95% confidence interval [CI] 17.5-39.1) for each one-tenth reduction in the late H/M ratio (hazard ratio [HR] 0.78; 95% CI 0.72-0.85; p<0.001). Of the MIBG parameters, only late H/M was an independent prognostic predictor (Table 2). Mortality was nine times higher in patients with lower H/M (first quartile, <1.55) at inclusion than in those with H/M above the median for the study population (≥1.83; HR 9.36; 95% CI 4.27-20.56; p<0.001). Mortality was also four times higher in patients with moderate reductions (1.55-1.83) in late H/M (HR 4.27; 95% CI 1.68-9.05; p=0.002) (Figure 3A).

To determine the potential added value of serial late H/M assessment for prognostic stratification, we assessed the accuracy of all the MIBG scans (558 scans, 2246 patient-years of follow-up) for predicting post-exam survival, which was found to be significantly reduced with lower late H/M
Table 2  Mortality during follow-up by age and MIBG parameters at inclusion.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Survival (n=185, 79.7%)</th>
<th>All-cause death (n=47, 20.3%)</th>
<th>Univariate Cox regression</th>
<th>Multivariate Cox regression</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>HR 95% CI p</td>
<td>HR 95% CI p</td>
</tr>
<tr>
<td>Median age, years (IQR)</td>
<td>37 (31-49)</td>
<td>59 (42-69)</td>
<td>1.067 1.046-1.089 &lt;0.001</td>
<td>1.052 1.046-1.089 &lt;0.001</td>
</tr>
<tr>
<td>Early H/M ratio</td>
<td>1.91±0.33</td>
<td>1.62±0.33</td>
<td>0.085 0.035-0.206 &lt;0.001</td>
<td>- - NS</td>
</tr>
<tr>
<td>Late H/M ratio</td>
<td>1.90±0.37</td>
<td>1.58±0.40</td>
<td>0.086 0.037-0.199 &lt;0.001</td>
<td>0.183 0.075-0.450 &lt;0.001</td>
</tr>
<tr>
<td>Washout rate</td>
<td>2.50</td>
<td>3.85</td>
<td>0.137 1.008-1.066 0.012</td>
<td>- - NS</td>
</tr>
<tr>
<td></td>
<td>(-2.30-7.14)</td>
<td>(0.0-12.50)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

CI: confidence interval; H/M: heart-to-mediastinum ratio; HR: hazard ratio; IQR: interquartile range; NS: non-significant.

![Figure 3](image_url) Survival by quartiles of H/M ratio assessed by first scintigraphic scan (A) and all scans performed during follow-up (B). CI: confidence interval; H/M: heart-to-mediastinum; HR: hazard ratio.

The risk of death increased by 27.8% for every one-tenth reduction in the late H/M ratio (HR 0.774; 95% CI 0.73-0.82; p=0.001), and was nine times higher in patients in the first H/M quartile (HR 9.15; 95% CI 5.44-15.38; p<0.001) (Figure 3B).

The accuracy of prediction based on serial exams was assessed by the area under the curve (AUC) on receiver operating characteristic analysis for different time intervals and compared with prognostic stratification based only on the initial MIBG scan (Table 3). Prognostic stratification based on annual repetition of MIBG imaging was slightly superior, but the differences were trivial for time intervals less than the median follow-up (4.5 years). The statistical power of prognostic stratification based on a single exam and on serial exams was similar, and very high (99%).

**Effect of liver transplantation on progression of myocardial sympathetic denervation assessed by 123I-MIBG scintigraphy**

Seventy patients (30.2%) underwent liver transplantation. In 39 of these patients surgery was performed less than one year after initial assessment. Perioperative mortality was 7.1% (5/70). The effect of transplantation on progression of sympathetic denervation was determined by comparing reductions in the late H/M ratio in images before (116 exams) and after (100 exams) transplantation. Of the former, 53 were performed less than a year before surgery.

In symptomatic patients requiring transplantation, late H/M decreased by 0.19/year (p<0.001) until transplantation; early H/M also decreased progressively and the
The cardiovascular manifestations of TTR-V30 M FAP are caused by autonomic dysfunction (disrupted control of blood pressure and heart rate) and by cardiac amyloid deposition (infiltrative cardiomyopathy, arrhythmias and conduction disturbances). However, phenotypic differences are seen in TTR-V30 M carriers according to geographic origin: cohort studies of Swedish patients report a high prevalence of restrictive cardiomyopathy,\(^3\),\(^4\) which is rare in Portugal, where cardiac autonomic neuropathy and conduction disturbances tend to dominate the clinical picture.\(^5\)

Previous studies have shown that MIBG scintigraphy is an indirect way to visualize the impairment of cardiac sympathetic innervation by amyloid deposition. Nakata et al.\(^1\) were the first to report a case of a patient with TTR-V30 M FAP with no myocardial activity on MIBG imaging, indicating involvement of the sympathetic nerves. Subsequent small studies demonstrated not only that myocardial denervation was strongly correlated with severity of polyneuropathy\(^6\) but also that MIBG uptake may be reduced before the onset of clinically apparent cardiac disease.\(^7\) In a previous study of 143 individuals, we showed that cardiovascular abnormalities as assessed by electrocardiography, Holter monitoring, ambulatory blood pressure monitoring, echocardiography and MIBG scintigraphy are common in TTR-V30 M FAP, particularly in patients with neurological involvement.\(^6\) Unlike other forms of systemic amyloidosis, the cardiovascular involvement in our study was mostly subclinical and obvious cardiac manifestations were rare. However, we did find that subclinical cardiovascular manifestations often preceded neurological involvement, which offers the possibility of earlier recognition of the onset of disease in asymptomatic carriers. We also demonstrated the prognostic value of MIBG imaging, which was in fact the strongest prognostic predictor of all the cardiac and neurological parameters assessed: patients with late H/M of less than 1.60 had a seven-fold higher risk of death during follow-up.

The present study confirms previous findings in a larger population and broadens the results to include serial H/M assessments. As expected, late H/M decreased significantly over time, but progression of cardiac denervation was so slow that annual repetition of MIBG imaging did not increase its accuracy in predicting all-cause death. Our results therefore suggest that repeated assessment of the H/M ratio is not justified for the purposes of prognostic stratification, at

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**Table 3** Accuracy of initial and serial late H/M ratios in MIBG scans for predicting mortality assessed by receiver operating characteristic analysis.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Late H/M assessment at inclusion (232 scans in 232 patients)</th>
<th>Serial late H/M assessments (558 scans in 232 patients)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Patient-years</td>
<td>No. of events</td>
</tr>
<tr>
<td>At any time</td>
<td>1171.5</td>
<td>38</td>
</tr>
<tr>
<td>12 months</td>
<td>109.1</td>
<td>3</td>
</tr>
<tr>
<td>24 months</td>
<td>352.7</td>
<td>9</td>
</tr>
<tr>
<td>36 months</td>
<td>467.3</td>
<td>15</td>
</tr>
<tr>
<td>48 months</td>
<td>534.4</td>
<td>17</td>
</tr>
<tr>
<td>60 months</td>
<td>553.8</td>
<td>22</td>
</tr>
</tbody>
</table>

AUC: area under the curve on receiver operating characteristic analysis. All other abbreviations as in Table 2.

**Figure 4** Changes in late H/M over time before and after liver transplantation. H/M: heart-to-mediastinum.

MIBG washout rate increased. By contrast, all scintigraphic parameters stabilized after transplantation and no statistically significant differences were detected over time (Table 4 and Figure 4).

**Discussion**

In this large prospective study of carriers of the TTR-V30 M mutation followed for a median of 4.5 years, we analyzed the progression of cardiac sympathetic denervation as assessed by the H/M MIBG uptake ratio, the prognostic value of this index for long-term survival, and the effect of liver transplantation. The main results are as follows: (1) cardiac denervation progresses slowly in the pre-symptomatic stages of the disease but tends to accelerate after symptom onset; (2) MIBG scintigraphy at inclusion is a strong predictor of prognosis, with the risk of death increasing by 27.8% for each one-tenth reduction in the late H/M ratio; and (3) liver transplantation stabilizes cardiac denervation in long-term follow-up.
least not more often than at intervals of less than five years (the median duration of follow-up in the present study).

Liver transplantation has been widely used for the last twenty years to remove the main source of mutant transthyretin, in order to slow the progression of clinical manifestations, including peripheral neuropathy, gastrointestinal symptoms and other visceral complications. Transplantation also improves long-term survival, especially when performed in the early stages of symptomatic disease. However, the effect of liver transplantation on the progression of cardiomyopathy in TTR-V30 M FAP is the subject of debate, with some studies suggesting that transplanted patients do not develop overt cardiomyopathy, while others – in Swedish, French and Japanese patients – report progression of arrhythmias and myocardial thickening due to amyloid deposition. This adverse outcome appears to be more frequent in Swedish patients with late-onset TTR-V30 M FAP.

It has been suggested that the main component of cardiac amyloid deposits following liver transplantation is normal (wild-type) transthyretin. Nevertheless, none of the studies that monitored the progression of amyloid infiltration by echocardiography following liver transplantation assessed its effect on cardiac autonomic denervation. In a small group of 31 French patients, Delahaye et al. compared the H/M ratio before and after liver transplantation, complemented with echocardiography; as in our study, no change was seen in sympathetic innervation following transplantation in a short follow-up (24±15 months), even though cardiac amyloid infiltration, as measured by myocardial thickness, had progressed. In the present study, of a larger group (70) of transplanted patients, the rate of decrease in H/M was assessed by comparing MIBG scintigraphy before (116 exams) and after (100 exams) transplantation. Of note, all scintigraphic parameters stabilized after surgery. There thus appears to be a distinction between amyloid deposition as assessed by myocardial thickness on echocardiography, which may progress after transplantation, and autonomic nerve involvement, which tends to stabilize.

### Conclusions

Cardiac denervation is common during the progression of TTR-V30 M FAP and quantification of the late H/M ratio on MIBG scintigraphy is valuable for prognostic stratification of these patients. Liver transplantation stabilizes cardiac denervation. Following transplantation, cardiac 123I-MIBG uptake remains stable over time, with no further recovery or deterioration. It is thus essential to perform transplantation sufficiently early in the natural history of the disease.

### Ethical disclosures

**Protection of human and animal subjects.** The authors declare that the procedures followed were in accordance with the regulations of the relevant clinical research ethics committee and with those of the Code of Ethics of the World Medical Association (Declaration of Helsinki).

**Confidentiality of data.** The authors declare that they have followed the protocols of their work center on the publication of patient data.

**Right to privacy and informed consent.** The authors declare that no patient data appear in this article.

### Conflicts of interest

The authors have no conflicts of interest to declare.

### Appendix A. Supplementary Material

Supplementary material associated with this article can be found in the online version available at [http://dx.doi.org/10.1016/j.repce.2016.08.009](http://dx.doi.org/10.1016/j.repce.2016.08.009).

### References