**Troponin-T Monitoring in Outpatients With Nonischemic Heart Failure**

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**Introduction and objective.** The usefulness of prolonged troponin-T (TnT) monitoring in outpatients with nonischemic heart failure (HF) is not clear. The aim of this study was to investigate the incidence, prognostic value, and determinants of a raised TnT level.

**Methods.** The study involved 80 outpatients (age 56[14] years, 69% male) with chronic stable HF (mean left ventricular ejection fraction, 24[9]; 51 in New York Heart Association class II and 29 in class III) of non-ischemic origin, as confirmed with coronary angiography. The TnT level was measured at study entry and at every outpatient visit (median interval, 3.1 months; interquartile range [IQR], 1.8-5.0 months) in a follow-up period of 22.2(10.6) months. Patients were TnT+ if the level was measurable (ie, >0.01 ng/mL).

**Results.** At study entry, 7 (9%) patients were TnT+. By 5 years, the cumulative incidence had reached 53%, and the median TnT level was 0.059 ng/mL (IQR, 0.023-0.100 ng/mL; range, 0.013-0.500 ng/mL). Beta-blocker therapy was associated with a reduction in incidence (hazard ratio [HR]=0.220; 95% confidence interval [CI], 0.089-0.540; P=.001) while the incidence increased with the N-terminal probrain natriuretic peptide (NT-proBNP) level (HR per 500 pg/mL=1.057; 95% CI, 1.023-1.092; P=.001).

**Conclusions.** A measurable TnT level was frequently observed during clinical monitoring of outpatients with non-ischemic HF and indicated a poor prognosis, even when the level was low.

**Key words:** Heart failure. Troponin T. Prognosis. NT-proBNP.

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**Valor de la determinación seriada de troponina T en pacientes ambulatorios con insuficiencia cardiaca no isquémica**

Introducción y objetivos. El valor del seguimiento prolongado de las determinaciones de troponina T (TnT) en pacientes ambulatorios con insuficiencia cardiaca no isquémica no está aclarado. Estudiamos la incidencia de las concentraciones aumentadas de TnT, sus determinantes y su valor pronóstico.

Métodos. Se incluyó a 80 pacientes ambulatorios (56 ± 14 años; el 69% varones) con insuficiencia cardiaca crónica (fracción de eyeción del ventrículo izquierdo, 24% ± 9%), situación estable (NYHA II, 51; NYHA III, 29) y etiología no isquémica confirmada por coronariografía. Los valores de TnT se midieron a la inclusión y en cada visita ambulatoria (median, 3 [intervalo intercuartílico, 1,8-5] meses), durante un seguimiento de 22,2 ± 10,6 meses. Se definió TnT+ como aquella detectable (> 0,01 ng/mL).

Resultados. En la visita inicial 7 (9%) pacientes presentaban TnT+, y la incidencia acumulada a 5 años alcanzó el 53%, con una mediana de 0,059 (intervalo intercuartílico, 0,023-0,1; mínimo, 0,013; máximo, 0,5) ng/mL. La incidencia de TnT+ estuvo disminuida en caso de tratamiento con bloqueadores beta (hazard ratio [HR] = 0,220; intervalo de confianza [IC] del 95%, 0,089-0,540; p = 0,001) y aumentada en relación con el nivel de NT-proBNP (HR = 1,005; IC del 95%, 1,001-1,010; p = 0,021). Presentaron eventos durante el seguimiento 14 (17,5%) pacientes (9 muertes cardiacas y 5 trasplantes urgentes), que ocurrieron en 12 (50%) de 24 con TnT+ frente a 2 (3,6%) de 56 con TnT– (p < 0,001). Tras el ajuste multivariable de Cox, la aparición de TnT+ fue predictora de eventos adversos (0,01 ng/mL: HR = 1,359; IC del 95%, 1,037-1,782; p = 0,026) independientemente del NT-proBNP level (HR per 500 pg/mL=1.057; 95% CI, 1.023-1.092; P=.001).

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INTRODUCTION

In spite of the numerous therapeutic advances, the prognosis for heart failure (HF) continues to be poor. It is important to develop awareness of the prognostic factors in this clinical setting, as these can easily be reproduced, and will help us to identify patients with the most unfavourable clinical evolution. Cardiac troponin (Tn) is a specific, sensitive marker of damage to the myocardium the introduction of which into clinical practice has changed the definition and management of acute coronary syndromes.

A variable percentage of patients hospitalized as a result of acute HF (23%-80%) have detectable levels of Tn, which appears to be related to the severity of the clinical condition and a higher incidence of death and re-hospitalization as a result of HF during follow up.4-9 It has been suggested that the presence of myocyte necrosis phenomena during acute decompensation may be contributing to the progression of the disease. The aetiology of HF in these studies was predominantly ischemic.4-7 With regard to patients with stable and outpatient HF, there are fewer data on the value of Tn, and these have also been gathered predominantly from populations with predominantly ischemic aetiology.10-12 On the other hand, in these studies Tn was measured transversally meaning the usefulness of prolonged Tn monitoring in patients with dilated nonischemic cardiomyopathy and its incidence was not clarified.12

The hypothesis of our study was the long-term monitoring of troponin values (TnT) in an outpatient clinic and in a population whose HF was not caused by ischemia, which may be helpful for prognostic stratification and clinical management. We therefore studied the incidence of its positivisation (TnT+), its clinical determinants and its prognostic value in long-term follow-up.

METHODS

Study Population

The study population consisted of 80 consecutive patients, attending an outpatient service specializing in HF during 2002-2004 and meeting the following criteria: a) an established diagnosis of chronic HF;13 b) stable clinical situation, without hospitalization during the previous 3 months and functional class of the New York Heart Association (NYHA) II-III (II/III 51/29); c) diagnosis of systolic function defined as a left ventricular ejection fraction (LVEF) <40% in the ultrasound study performed during the month prior to their inclusion; and d) nonischemic aetiology defined in all patients as the concurrence of absence of a history of ischemic cardiopathy and absence of coronary lesions. All patients had at least 1 coronary angiographic study and the study excluded all those with coronary lesions of any severity or with any history of acute coronary syndrome, as well as those on chronic haemodialysis. The aethiology was established as idiopathic in 62 (78%), of patient, valvular in 8 (10%), familiar in 6 (7%) and alcoholic in 4 (5%).

Their parameters and clinical, ECG, and analytical characteristics were included with the data from the initial visit. Patients were subsequently monitored at the outpatient clinic with an average frequency of 3.1 months (1.8-5.0 months). At each visit, the clinical evaluation included evaluation of the clinical status and blood sampling for TnT determination. All the patients had a conventional ECG during the month before they were included in the study (Sonos 550, Phillips, Andover, Massachusetts). The left ventricular ejection fraction was calculated using the modified Simpson method, using image with second harmonic second, and the projections and standard measurements were performed for the study of anatomy and cardiac function.14

The written, informed consent was obtained from all patients before they were included in the study.

Laboratory Parameters

Blood samples were obtained by venous puncture when the patients arrived to surgery, between 8 am and 10 am, fasting and after resting for 20 minutes. The cardiac TnT was measured using Electrochemiluminescence Immunoassay with a Roche Elecsys 2010 (Roche Diagnostics, Manheim, Germany) modular analyser. The detection limit provided by the supplier was <0.01 ng/mL and the variability coefficient was 5.6% to 0.04 ng/mL. Raised TnT was defined in accordance with the supplier, as a percentage value in excess of 99 percentile (>0.01

NT-proBNP (500 pg/ml: HR = 1,057; IC del 95%, 1,023-1,092; p = 0,001).

Conclusiones. En el seguimiento ambulatorio de pacientes con insuficiencia cardiaca no isquémica, la aparición de TnT+ es frecuente e identifica un peor pronóstico, incluso con bajas concentraciones.

Palabras clave: Insuficiencia cardiaca. Troponina T. Prognóstico. NT-proBNP.
ng/mL) obtained from a population of 1951 healthy volunteers. The patients with a value of >0.01 ng/mL during the follow-up period were identified as TnT+, while patients with values <0.01 ng/mL during the entire follow-up period were classed as TnT-. Additionally, the index of TnT was calculated as the relationship between TnT maximum/0.01 ng/mL. The plasma levels of NT-proBNP were diagnosed using the test “proBNP assay” using the Elesys 2010 (Roche Diagnostics, Manheim, Germany) analyser. The precision of the assay was from 1.8% at 800 pmol/L to 2.7% at 20.7 pmol/L and the detection limits were 0.6 pmol/L and 4130 pmol/L. The conversion ratio of pmol/L to pg/ml was ×8457. The haemoglobin and hematocrit concentrations were determined with the automatic haematological analyserXE-2100 (Symex, Kobe, Japan) and the biochemical measurements were obtained with the Roche/Hitachi Modular analyser (Roche diagnosis, Manheim, Germany). The glomerular filtration rate (GFR) was calculated using the simplified MDRD formula (mL/min/1.73 m², 186.3 × [plasma creatinine]–1.154 × [age]–0.203) (the correction factor for women was ×0.742).

Follow-up and Clinical Events

All patients were followed up, obtaining the vital status of all of them. The main events in the study were death for cardiac causes, defined as death due to heart failure and/or sudden death and the need for emergency heart transplant, the decision for which were not influenced by any of the biochemical findings of the study. Patients dying from non-cardiac causes or elective transplant were followed up until death, but this was not considered an event.

Analysis of Results and Statistics

Test the normality of the distribution of the quantitative variables was checked using Kolmogorov-Smirnov test; the variables with abnormal distribution were expressed as medium [interquartile range, IQR], the rest as mean (standard deviation [SD]) for quantitative variables and number (%) for qualitatives. At baseline level, the comparison of quantitative variables among patients with TnT+ and TnT- was performed using the Wilcoxon Student t test for independent samples, as appropriate, and the Fisher’s test for qualitative variables. The variables with P<.1 in the univariable analysis were introduced in the Cox multiple regression for the identification of TnT+ predictors. The Kaplan-Meier analysis and the statistical log rank were used for the description of the accumulated incidence of TnT+ and the differences between groups. The association between levels of TnT and the appearance of events is studied using the Wilcoxon test for independent samples and the receiver operating characteristic curve (COR). The Cox regression analysis was used for the univariable study of event predictors.

The variables with P<.1 in the univariable were introduced in a multivariable Cox model, that was also adjusted by known risk factors (LVEF, class NYHA) and by the baseline characteristics associated in a significant manner with TnT+. The TnT+ and the TnT index were introduced separately into the model. The adherence to the proposed proportionality of risks was checked using logarithmic representations of the accumulated risks, the analysis of the Schoenfeld residuals and the Ln representation (-Ln S[t]) for each variable after its categorization. The event-free survival after the appearance of an initial TnT+ value was studied using Kaplan-Meier curves and the log rank statistic.

A P<.05 was considered significant. The statistical analysis was performed using the statistical software package for social sciences (SPSS v. 14.0 for Windows, SPSS Inc., Chicago, Illinois, USA).

RESULTS

Clinical Characteristics and TnT+

The study population consisted of 80 patients, whose clinical characteristics are set out in Table 1. At the initial visit, 7 patients (8.7%) presented TnT+. The Kaplan-Meier analysis showed an accumulated incidence of TnT+ of 52.8% at 5 years during the follow up (Figure 1). In TnT+ patients, the levels measured presented a median of 0.059 ng/mL (IQR, 0.023-0.100; minimum, 0.013; and maximum, 0.500). At baseline, the variables associated with the appearance of a TnT+ were: raised NT-proBNP maximum, 0.500). At baseline, the variables associated in a significant manner with TnT+. The TnT+ and the TnT index were introduced separately into the model. The adherence to the proposed proportionality of risks was checked using logarithmic representations of the accumulated risks, the analysis of the Schoenfeld residuals and the Ln representation (-Ln S[t]) for each variable after its categorization. The event-free survival after the appearance of an initial TnT+ value was studied using Kaplan-Meier curves and the log rank statistic.

A P<.05 was considered significant. The statistical analysis was performed using the statistical software package for social sciences (SPSS v. 14.0 for Windows, SPSS Inc., Chicago, Illinois, USA).

PROGNOSIS AND TnT

During the clinical follow-up (median, 28.5 months [IQR, 12.9-39.4]), a total of 14 (17.5%) patients presented adverse clinical events: 9 cardiac deaths (2 sudden deaths
TABLE 1. Clinical Characteristics

<table>
<thead>
<tr>
<th></th>
<th>Total population (n=80)</th>
<th>TnT: (n=214)</th>
<th>TnT- (n=5)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, y</td>
<td>56 (14)</td>
<td>59 (16)</td>
<td>55 (13)</td>
<td>.18</td>
</tr>
<tr>
<td>Sex, % men</td>
<td>55 (69)</td>
<td>16 (67)</td>
<td>39 (70)</td>
<td>.79</td>
</tr>
<tr>
<td>IMC, kg/m²</td>
<td>28 (25-31)</td>
<td>27 (26-29)</td>
<td>28 (25-32)</td>
<td>.54</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>22 (27)</td>
<td>9 (37)</td>
<td>13 (23)</td>
<td>.19</td>
</tr>
<tr>
<td>Hypertension</td>
<td>32 (40)</td>
<td>10 (42)</td>
<td>22 (39)</td>
<td>.84</td>
</tr>
<tr>
<td>Active smoker</td>
<td>14 (17)</td>
<td>2 (8)</td>
<td>12 (21)</td>
<td>.20</td>
</tr>
<tr>
<td>Hypercholesterolaemia</td>
<td>26 (32)</td>
<td>5 (21)</td>
<td>21 (37)</td>
<td>.14</td>
</tr>
<tr>
<td>NYHA II/III, %</td>
<td>64/36</td>
<td>66/34</td>
<td>59/42</td>
<td>.56</td>
</tr>
<tr>
<td>Previous hospitalisations for HF</td>
<td>26 (33)</td>
<td>12 (50)</td>
<td>16 (29)</td>
<td>.06</td>
</tr>
<tr>
<td>Duration HF, y</td>
<td>7.0 (5.2)</td>
<td>6.4 (4.9)</td>
<td>7.3 (5.4)</td>
<td>.49</td>
</tr>
<tr>
<td>LVEF, %</td>
<td>23.6 (9.4)</td>
<td>23.5 (9.0)</td>
<td>23.7 (9.6)</td>
<td>.96</td>
</tr>
<tr>
<td>Auricular fibrillation</td>
<td>15 (19)</td>
<td>3 (13)</td>
<td>12 (21)</td>
<td>.53</td>
</tr>
<tr>
<td>Chronic pneumopathy</td>
<td>4 (5)</td>
<td>1 (4)</td>
<td>3 (5)</td>
<td>.82</td>
</tr>
<tr>
<td>Creatinine, mg/dL</td>
<td>1.0 (0.8-1.2)</td>
<td>1.2 (0.9-1.5)</td>
<td>1.0 (0.8-1.2)</td>
<td>.01</td>
</tr>
<tr>
<td>GFR, mL/min/1.73 m²</td>
<td>74 (69-89)</td>
<td>64 (45-77)</td>
<td>77 (63-96)</td>
<td>.01</td>
</tr>
<tr>
<td>Troponin T, ng/mL</td>
<td>0.01 (0.0-0.02)</td>
<td>0.06 (0.02-0.10)</td>
<td>–</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>NT-proBNP, pg/mL</td>
<td>839 (321-2218)</td>
<td>2312 (1619-3537)</td>
<td>542 (245-1243)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>C-reactive protein, mg/dL</td>
<td>0.5 (0.2-0.9)</td>
<td>0.6 (0.3-1.2)</td>
<td>0.4 (0.2-0.8)</td>
<td>.10</td>
</tr>
<tr>
<td>Haemoglobin, g/dL</td>
<td>14 (1.5)</td>
<td>13.7 (1.7)</td>
<td>14.2 (1.4)</td>
<td>.15</td>
</tr>
<tr>
<td>Sodium, mEq/L</td>
<td>139 (3.5)</td>
<td>139 (4.7)</td>
<td>139 (2.9)</td>
<td>.59</td>
</tr>
<tr>
<td>Uric acid, mg/dL</td>
<td>6.9 (2.3)</td>
<td>7.7 (2.9)</td>
<td>6.6 (1.9)</td>
<td>.13</td>
</tr>
<tr>
<td>Beta-blockers</td>
<td>66 (82)</td>
<td>16 (67)</td>
<td>50 (89)</td>
<td>.02</td>
</tr>
<tr>
<td>ACE inhibitors/ARA II</td>
<td>80 (100)</td>
<td>24 (100)</td>
<td>56 (100)</td>
<td>1</td>
</tr>
<tr>
<td>ASA diuretics</td>
<td>70 (87)</td>
<td>22 (92)</td>
<td>48 (86)</td>
<td>.71</td>
</tr>
<tr>
<td>Statins</td>
<td>18 (22)</td>
<td>6 (25)</td>
<td>12 (21)</td>
<td>.72</td>
</tr>
<tr>
<td>Spironolactone</td>
<td>51 (64)</td>
<td>14 (58)</td>
<td>37 (66)</td>
<td>.50</td>
</tr>
<tr>
<td>Anticoagulation</td>
<td>40 (50)</td>
<td>13 (54)</td>
<td>27 (48)</td>
<td>.81</td>
</tr>
<tr>
<td>Antiaggregation</td>
<td>17 (21)</td>
<td>6 (25)</td>
<td>11 (20)</td>
<td>.59</td>
</tr>
</tbody>
</table>

The values are expressed in mean (standard deviation), median (interquartile range), and number (%). BMI indicates body mass index; NYHA, New York Heart Association; HF, heart failure; LVEF, left ventricular ejection fraction; GFR, glomerular filtration rate; NT-proBNP, N-terminal probrain natriuretic peptide type B; ACEI, inhibitors of angiotensin-converting enzyme; ARAII, angiotensin II receptor antagonists.

Figure 1. Accumulated incidence of troponin T>0.01 ng/mL (TnT+).
and 7 due to refractory heart failure) and 5 emergency code heart transplants; and no patients presented acute coronary syndrome. The levels of TnT were the highest in patients developing events ($P < .001$) (Figures 3A and 3B). The ROC analysis of the levels of TnT showed an area under the curve of 0.83 (95% CI, 0.71-0.95) ($P < .001$) for the detection of events during the follow up (Figure 4). The optimum cut off point was a detectable value of TnT (>0.01 ng/mL or TnT+), with a sensitivity of 86%, specificity 82%, positive predictive value 50% and negative 96%.

During the follow-up, the adverse events occurred in 12 (50%) of 24 patients with TnT+ compared with 2 (3.6%) of 56 with TnT- ($P < .001$). Of the patients with TnT+, 7 (29%) died of cardiac causes and 5 (21%) required an emergency heart transplant. The Kaplan-Meier survival analysis (Figure 5) showed that the first adverse event occurred in the fourth month of follow up after positivization of TnT, the survival being significantly decreased among patients with TnT+ (log rank < 0.001).

The results of the Cox regression analysis for the appearance of events is set out in Table 2; after adjusting the multivariable model, a value of TnT+ (>0.01 ng/mL) was the most powerful predictor of adverse events (model 1), and when considering the TnT index (model 2), this showed that each increase of 0.010 ng/mL in the TnT was associated with a 36% increase in the risk of adverse events (Table 2).

**DISCUSSION**

This study evaluated the usefulness of monitoring TnT in an outpatient population with nonischemic HF and severe systolic dysfunction. In this population we found: *a*) the frequency of positivization of TnT occurs in a high population of patients during a broad time range; *b*) the risk of positivization was greater in the presence of greater severity of the clinical situation and greater myocardial stress (NT-proBNP) while the use of beta blocker therapy was associated with lower levels of risk; *c*) the appearance of a TnT+, even at low levels, led to a poor prognosis, which is independent of other clinical, analytical and ECG variables; and *d*) after the positivization of the TnT, the occurrence of the adverse events occurs after several months (>4 months), which could be interpreted as an opportunity to treat these patients more intensely.

**Prevalence, Frequency, and Determinants**

The majority of the information is drawn from transversal prevalence studies, that show variable percentages (15%-80%) depending on the population evaluated. Most studies were performed with hospitalized populations with acute HF, where the greatest prevalence is reported (45%-80%). There are fewer data gathered from outpatient populations with stable HF: Missov (n=33; 55% ischemic) found a prevalence...
of 15%, Perna et al\textsuperscript{10} (n=115; 50% ischemic) of 32% and Hudson et al\textsuperscript{11} (n=136; 62% ischemic) of 24%; which suggests a lower prevalence in the out-patient setting. Our study showed a prevalence at inclusion of only 9%, which could be due partly to the absence of coronary disease, as some authors have suggested the frequency is higher in the presence of ischemic aetiology.\textsuperscript{7,21}

On the other hand, different works have shown an association between TnT+ and lower LVEF, worse NYHA class, greater BNP, and deterioration of renal function.\textsuperscript{11,16} These associations suggest that the variability of prevalence is also a consequence of the differences in the severity of the HF syndrome. Thus, recently Miller et al describe a frequency of TnT+ of 54% in 190 out-patient patients, most of them with ischemic aetiology (55%).\textsuperscript{12} However, this work included patients who had recently been hospitalised with acute HF, all of whom were in NYHA III-IV and with associated deterioration of renal function (TFG, 45 mL/min/1.73 m\textsuperscript{2}). Compared to this, our population did not present recent hospitalizations, the majority (64%) were in NYHA II

![Figure 3](https://example.com/fig3.png)  
**Figure 3.** Block diagram showing the values of troponin T in accordance with the presence of absence of death (A) and of death or transplant (B) during the follow-up.

![Figure 4](https://example.com/fig4.png)  
**Figure 4.** Analysis of the characteristic operator/receptor curve for the levels of troponin T and the appearance of events during the follow-up.

### Table 2. Variables Associated With a Higher Risk of Adverse Events in the Cox Regression Analysis

<table>
<thead>
<tr>
<th>Variable</th>
<th>Univariable</th>
<th>Multivariable (Model 1)\textsuperscript{a}</th>
<th>Multivariable (Model 2)\textsuperscript{a}</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>P</td>
<td>HR (95% CI)</td>
<td>HR (95% CI)</td>
</tr>
<tr>
<td>TnT+, &gt;0.01 ng/mL</td>
<td>&lt;.001</td>
<td>14.3 (2.87-71.4)</td>
<td>.001</td>
</tr>
<tr>
<td>TnT index, &gt;0.01 ng/mL</td>
<td>.004</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>NT-proBNP, 500 pg/mL</td>
<td>&lt;.001</td>
<td>1.036 (1.000-1.074)</td>
<td>.060</td>
</tr>
<tr>
<td>Age, y</td>
<td>.037</td>
<td>1.042 (1.003-1.064)</td>
<td>.030</td>
</tr>
<tr>
<td>Haemoglobin, g/L</td>
<td>.040</td>
<td>0.659 (0.442-1.029)</td>
<td>.067</td>
</tr>
</tbody>
</table>

NT-proBNP indicates N-terminal probrain natriuretic peptide type B; TnT, troponin T; TnT indexed, TnT peak/0.01 ng/mL. \textsuperscript{a}Adjusted by class NYHA, LVEF, GFR, previous hospitalisations due to heart failure and beta blocker treatment.

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and the GFR was clearly greater (74 mL/min/1.73 m²). In the light of all these findings, it seems to be that the characteristics of the population and the severity of the HF syndrome determine the variability in the prevalence described for TnT+.

The greatest contribution of our work is the description of the frequency or positivization of TnT during the long-term follow-up. The accrued frequency is high (53%) but during a wide range of time (5 years). Previously, only the work carried out by Miller et al had contributed data regarding monitoring TnT. However, as has been indicated, in this study, the initial prevalence was already high, being 54%, and reaching 71% after 2 years of follow-up. The increased severity of the HF syndrome appears to be responsible for the high initial prevalence, however, the annual rate of incidence was similar; being 8.2% in the work by Miller and 8.9% according to our findings.

An important finding of our study is the decreased risk of the appearance of a TnT+ in patients receiving treatment with beta-blockers, which, together with the increased risk associated with NT-proBNP, was independent after the multivariable adjustment. The association between the levels of NT-proBNP and TnT has been described in a standard manner by several authors showing a greater degree of ventricular stress. However, none of the studies have shown the protective effects of beta-blockers on positivization of TnT, and consequently of myocyte necrosis. It is a known fact that circulating catecholamine are predictors of the severity and prognosis in HF and that in other clinical situations, a high adrenergic influx can lead to cardiomyopathy with myocyte necrosis even in the absence of coronary disease. In experimental models the stimulation of the β1 receptors leads to necrosis and apoptosis, leading to progressive of dysfunction and adverse ventricular remodelling. In animal models of HF, blocking the β1 receptors protected against necrosis and apoptosis of the cardiomyocytes, suggesting this blockade as one of the mechanisms explaining the benefits of beta-blockers. Our Study illustrates this protection in a clinical setting, regardless of the NT-proBNP level, and supports that in all the aetiopathogenic mechanisms suggested, excess β1 stimulation might be the main cause contributing to nonischemic myocardial damage in HF.

**Prognosis**

Our work confirmed the significant association between the presence of TnT+ levels and a worse prognosis in patients with nonischemic HF. During recent years, the presence of TnT+ in hospitalized patients with acute HF has been related to a greater frequency of death, transplant and rehospitalization due to HF. In the outpatient setting, raised levels have also been associated with adverse
events.\textsuperscript{5-10,12} In our work, the levels of TnT detected were minimum (range, 0.013-0.500) and independently of the amount, the presence of measurable levels (>0.01 ng/mL) is associated with increased risk, which was independent of other clinical, analytical, and ECG variables. The prognostic value of TnT, even minimum detections, has also been shown by others, and in the case of the TnT the majority of authors describe an increase risk after it is detected.\textsuperscript{5,7,10,11,17}

One interesting observation is the timeline of appearance of the events after TnT+ has been detected. In earlier studies, the appearance of events is earlier and described during the first month, however, the results of this are not comparable as these are prevalence studies on hospitalized populations with predominantly ischemic aetiology.\textsuperscript{6,8,10,11} In our study, the events occurred as from the fourth month after TnT+ was detected, although we cannot rule out the possibility that close monitoring meant this was detected sooner before the occurrence of the first event. In any case our work shows that in the outpatient population with HF, the detection of TnT+ can open a therapeutic window during which the treatment given to patients can be intensified, thus helping to improve their prognosis.

The results obtained from this study draw attention to the importance and usefulness of monitoring TnT when following up outpatients with chronic stable HF. In our opinion, from all the findings described, it is evident that TnT is a useful tool in this area. In the future, the use of more sensitive methods for detecting TnT and their integration together with the measurement of NT-proBNP could contribute to new applications. One advantage of TnT is the absence of a variable cut-off point or population dependent cut-off point, meaning that a single detection is associated with increased risk, while the cut-off point for BNP is dependent on the population being tested.\textsuperscript{30}

In our analysis, both were quantitative predictors of risk, which, together with the numerous works identifying both as independent prognostic markers, suggests there is a need to use both jointly.

Limitations

The main limitation of the study was the selective nature of the population, which could have led to a bias that interfered with the extrapolation of the results to other populations. On the other hand, the analytical monitoring was not standard and, although programmed every 3 months, it must be acknowledged that there was a variable inherent to the healthcare practices themselves, neither could we exclude that although all the samples were outpatient and scheduled, some were taken during phases of decompensation. Lastly, the observational nature of the association between beta-blockers and a decreased elevation of TnT, does not enable us to infer the protective role of the same, but establishes the need to investigate this aspect more deeply.

CONCLUSIONS

In outpatients with stable chronic HF of a nonischemic nature, the detection of TnT, even at low levels, means an increased risk of adverse cardiac events, which is independent of other clinical, analytical and ECG variables. Treatment with beta-blockers had a protective effect, which was independent of the level of NT-proBNP.

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