Non-ST-Elevation Acute Myocardial Infarction With Normal Coronary Arteries: Predictors and Prognosis

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Introduction and objectives. Occasionally, coronary arteries without significant stenosis are observed during invasive treatment of acute non-ST-elevation myocardial infarction (NSTEMI). The aim was to investigate predictive factors and prognosis in these patients.

Methods. The study involved 504 patients admitted for NSTEMI who underwent cardiac catheterization. The primary end-point was the observation of coronary arteries without significant stenosis, and the secondary end-point was death or myocardial infarction within a median of 3 years. In evaluating the secondary end-point, a control group of 160 patients with a normal troponin level and no significant coronary artery stenosis who were admitted for chest pain during the same period was included.

Results. Overall, 64 patients (13%) had coronary arteries without significant lesions. The predictors were: female sex (odds ratio [OR] = 6.6; P = .0001), age <55 years (OR = 3.0; P = .001), and the absence of diabetes (OR = 2.4; P = .02), previous antiplatelet treatment (OR = 3.9; P = .007) or ST-segment depression (OR = 2.4; P = .008). The composite variable of female sex plus at least 2 additional predictive factors had a specificity of 85% and a sensitivity of 53% for coronary angiography showing no significant stenosis. The absence of coronary artery stenosis decreased the probability of death or myocardial infarction during follow-up (hazard ratio = 0.3, 95% confidence interval, 0.2-0.9; P = .03). Among all patients without significant stenosis (n = 224), there was no difference in the event rate between those with elevated and normal troponin levels.

Conclusions. In NSTEMI, female sex, age <55 years and the absence of diabetes, previous antiplatelet treatment, or ST-segment depression were all associated with coronary angiography showing no significant stenosis. The long-term prognosis in these patients was good.

Key words: Myocardial infarction. No significant stenosis on coronary angiography. Predictors. Prognosis.

Infarto de miocardio sin elevación del ST con coronarias normales: predictores y pronóstico

Introducción y objetivos. El manejo invasivo del infarto agudo de miocardio sin elevación del ST (IAMSEST) detecta en ocasiones arterias coronarias sin estenosis significativas. Nuestro objetivo fue evaluar los factores asociados y el pronóstico de esta población.

Métodos. Estudiamos a 504 pacientes ingresados por IAMSEST y sometidos a cateterismo cardíaco. El objetivo primario fue el hallazgo de coronarias sin estenosis significativas y el secundario, la mortalidad o el infarto a una mediana de 3 años. Para evaluar el objetivo secundario, se utilizó un grupo control de 160 pacientes ingresados por dolor torácico durante el mismo periodo con troponina normal y coronarias sin estenosis significativas.

Resultados. Encontramos coronarias sin lesiones significativas en 64 (13%) pacientes. Los predictores fueron: ser mujer (odds ratio [OR] = 6.6; p = 0.0001), edad < 55 años (OR = 3.0; p = 0.001) y ausencia de diabetes (OR = 2.4; p = 0.02), tratamiento antiagregante previo (OR = 3.9; p = 0.007) o descenso del ST (OR = 2.4; p = 0.008). La variable ser mujer con al menos dos variables adicionales identificó una coronariografía sin estenosis significativas con especificidad del 85% y sensibilidad del 53%. La ausencia de estenosis coronarias significativas disminuyó la probabilidad de muerte o infarto durante el seguimiento (hazard ratio = 0.3; intervalo de confianza del 95%, 0.2-0.9; p = 0.03). En el grupo de pacientes sin estenosis coronarias significativas (n = 224), no hubo diferencias en la tasa de sucesos entre los pacientes con troponina elevada y normal.

Conclusiones. El sexo femenino, la edad < 55 años y la ausencia de diabetes, tratamiento antiagregante previo o descenso del ST se asocian a una coronariografía sin estenosis significativas en el IAMSEST. El pronóstico a largo plazo de esta población fue bueno.

January 20, 2001 and April 1, 2008. The following criteria had to be met for inclusion in the study: 

- **a)** no persistent ST elevation in the initial cardiogram; 
- **b)** elevated troponin I in serial measurements on arrival at hospital and 8-12 hours after the onset of pain; 
- **c)** no prior history of significant coronary artery stenosis determined by coronary angiography; and 
- **d)** cardiac catheterization and coronary angiography during admission. 

The study population accounted for 62% of all patients admitted for NSTEMI without prior significant coronary artery stenosis recorded during the study period; the remaining 38% of the population did not undergo catheterization.

In our hospital, 2 different troponin I tests were used during the study: Immulite (Diagnostic Products Corporation, Los Angeles, California, US; upper limit of normal, 0.5 ng/mL) until October 2003 and Dimension (Dade Behring, Newark, Delaware, USA; upper limit of normal, 0.2 ng/mL) thereafter.

The management of the patients and the indication for catheterization were according to the judgment of the attending cardiologist.

### Variables Recorded

The following clinical variables were recorded for each patient: including coronary risk factors (age, smoking habit, hypertension, hypercholesterolemia, diabetes mellitus, family history), history of myocardial infarction, peripheral artery disease, stroke, and prior antiplatelet therapy. An ST-segment depression was recorded if more than 1 mm decrease occurred, compared to the initial electrocardiogram (ECG). In all patients, creatinine was measured on admission and renal failure defined as levels greater than 1.3 mg/dL. Coronary stenosis was considered significant if it exceeded 50% in one of the 3 main epicardial vessels or the left coronary artery.

### Objectives

The primary objective was to analyze the factors predictive of the absence of significant coronary stenosis (>50%). The secondary objective was to determine total mortality or acute myocardial infarction during follow-up. The median follow-up period was 3 years (interquartile range, 64-184 weeks). Follow-up exceeded 6 months in 97% of the patients and 1 year in 91%.

### Control Group

Prognosis in patients with NSTEMI and normal coronary arteries was compared with that of a control group of 160 consecutive patients who were
admitted during the same period for chest pain of possible coronary origin according to the judgment of the cardiologist on duty. These patients had normal troponin I levels, and the coronary angiography performed during hospitalization showed lack of significant coronary stenosis. In total, 37 patients showed ST-segment depression (>1 mm) in the initial ECG; 13 had a negative T wave (>1 mm), and 10 had left bundle branch block. Coronary angiography was done directly in 94 patients. For the remaining patients, prior to catheterization, exercise testing was done in 59 patients (28 with a positive result and 31 with an inconclusive result), and magnetic resonance imaging with dipyridamole in 15. The diagnosis on discharge was, in all cases, chest pain of unknown origin.

**Statistical Analysis**

The variables associated with the primary objective were analyzed using the χ² test. In order to transform age into a qualitative variable, its association with the primary outcome measure was analyzed using quartiles and the cutpoint corresponding to the upper limit of the first quartile (55 years) was chosen for its greater discriminative power. A multivariate logistic regression analysis was subsequently performed with the variables that were associated with the primary endpoint in the univariate analysis with P≤.1. The entry criterion in the model was P<.05. Odds ratios (OR) and 95% confidence intervals (CI) were calculated.

For the secondary objective, a univariate analysis using a χ² test was performed, along with a multiple Cox regression (backward conditional method) with the variables of prognostic value in the univariate analysis. The hazard ratio (HR) and corresponding 95% confidence intervals (CI) were calculated. Statistical significance was set at a value of P less than .05. The SPSS statistical analysis program, version 9.0 (SPSS Inc., Chicago, Illinois, USA) was used.

**RESULTS**

**Characteristics of the Study Population**

Table 1 shows the characteristics of the study population. Normal coronary arteries were observed in 64 patients (13%; 95% CI, 15.6-9.7). Sixty-five patients (13%) died during follow-up, 80 (16%) had an acute myocardial infarction, and 122 (24%) had an acute myocardial infarction or death.

**Factors Related to Normal Coronary Angiogram**

Table 1 shows the differences in the baseline characteristics between patients with and without significant coronary stenosis. Table 2 shows the results of the multivariate analysis. Female sex was the variable most strongly associated with absence of significant coronary stenosis (OR=6.6; 95% CI, 3.5-12.5; P=.0001). Other variables were age less than 55 years (OR=3.0; 95% CI, 1.5-5.8; P=.001), absence of diabetes (OR=2.4; 95% CI, 1.2-4.8; P=.02), prior antiplatelet therapy (OR=3.9; 95% CI, 1.9-7.9; P<.001), and family history of ischemic heart disease (OR=3.8; 95% CI, 1.4-10.0; P=.007).

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**Table 1. Baseline Characteristics of the Patients With Non-ST-Elevation Myocardial Infarction and Prognosis According to Presence or Absence of Significant Coronary Artery Stenosis**

<table>
<thead>
<tr>
<th>Variable</th>
<th>Without Coronary Stenosis (n=64)</th>
<th>With Coronary Stenosis (n=440)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>60 (12.5)</td>
<td>66 (11.3)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Heart failure</td>
<td>3 (4.7)</td>
<td>49 (11.1)</td>
<td>.128</td>
</tr>
<tr>
<td>Men</td>
<td>27 (42.2)</td>
<td>325 (73.9)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Smoking</td>
<td>17 (26.6)</td>
<td>142 (32.3)</td>
<td>.391</td>
</tr>
<tr>
<td>Hypertension</td>
<td>33 (51.6)</td>
<td>264 (60)</td>
<td>.222</td>
</tr>
<tr>
<td>Hypercholesterolemia</td>
<td>23 (35.9)</td>
<td>205 (46.6)</td>
<td>.139</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>13 (20.3)</td>
<td>167 (38)</td>
<td>.005</td>
</tr>
<tr>
<td>Peripheral artery disease</td>
<td>0</td>
<td>35 (8)</td>
<td>.015</td>
</tr>
<tr>
<td>Creatinine on admission</td>
<td>0.98 (0.34)</td>
<td>1.13 (0.40)</td>
<td>.007</td>
</tr>
<tr>
<td>ST-segment depression</td>
<td>17 (26.6)</td>
<td>199 (45.2)</td>
<td>.005</td>
</tr>
<tr>
<td>Prior antiplatelet use</td>
<td>5 (7.8)</td>
<td>148 (33.6)</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Family history of ischemic heart disease</td>
<td>4 (6.3)</td>
<td>39 (8.9)</td>
<td>.634</td>
</tr>
<tr>
<td>History of myocardial infarction</td>
<td>2 (3.1)</td>
<td>65 (14.8)</td>
<td>.009</td>
</tr>
<tr>
<td>History of heart failure</td>
<td>2 (3.1)</td>
<td>9 (2)</td>
<td>.638</td>
</tr>
<tr>
<td>History of stroke</td>
<td>2 (3.1)</td>
<td>30 (6.8)</td>
<td>.408</td>
</tr>
<tr>
<td>Acute myocardial infarction during follow-up</td>
<td>2 (3.1)</td>
<td>78 (17.7)</td>
<td>.001</td>
</tr>
<tr>
<td>Death during follow-up</td>
<td>2 (3.1)</td>
<td>63 (14.3)</td>
<td>.009</td>
</tr>
</tbody>
</table>

Data are shown as means (SD).
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rate of death or infarction during follow-up (6% vs 27%; \( P = .0001 \)) after adjusting for other variables of prognostic value (HR=0.3; 95% CI, 0.2-0.9; \( P = .03 \)) (Figure). The other associated variables were heart failure on admission (HR=2.6; 95% CI, 1.7-4.0; \( P = .0001 \)), age (per year, HR=1.02; 95% CI, 1.0-1.04; \( P = .05 \)), diabetes mellitus (HR=1.5; 95% CI, 1.0-2.2; \( P = .04 \)), prior myocardial infarction (HR=1.6; 95% CI, 1.1-2.4; \( P = .03 \)), and renal failure (HR=1.7; 95% CI, 1.1-2.6; \( P = .02 \)).

The outcomes for the patients with NSTEMI and normal coronary arteries were compared with those of the control group. Table 3 shows the baseline differences between the 2 groups. Patients in the control group were older and more frequently had hypercholesterolemia and prior treatment with antiplatelet agents. Likewise, there was a tendency toward a greater proportion of men and systemic hypertension. During follow-up, among the entire population of patients with normal coronary arteries (n=224), 23 patients (10%) died or had an infarction. The death rates and rates of infarction in the subgroup with elevated troponin and normal troponin were 4 (6%) and 19 (12%) (HR=0.7; 95% CI, 0.2-2.2; \( P = .6 \)).

**DISCUSSION**

In our series, we found that 13% of the patients with NSTEMI lacked significant coronary stenosis according to the coronary angiogram procedure performed during the initial stay in hospital. Female sex, younger age, absence of diabetes, and prior antiplatelet treatment,

<table>
<thead>
<tr>
<th>Variable</th>
<th>OR (95% CI)</th>
<th>( P )</th>
</tr>
</thead>
<tbody>
<tr>
<td>Female sex</td>
<td>6.6 (3.5-12.5)</td>
<td>.0001</td>
</tr>
<tr>
<td>Age &lt;55 years</td>
<td>3.0 (1.5-5.8)</td>
<td>.001</td>
</tr>
<tr>
<td>Without diabetes mellitus</td>
<td>2.4 (1.2-4.8)</td>
<td>.02</td>
</tr>
<tr>
<td>No prior antiplatelet treatment</td>
<td>3.9 (1.5-10.2)</td>
<td>.007</td>
</tr>
<tr>
<td>No ST-segment depression in ECG</td>
<td>2.4 (1.2-4.4)</td>
<td>.008</td>
</tr>
</tbody>
</table>

CI indicates confidence interval; ECG, electrocardiogram; OR, odds ratio.

CI, 1.5-10.2; \( P = .007 \), and presentation without ST-segment depression in the ECG (OR=2.4; 95% CI, 1.2-4.4; \( P = .008 \)). The C statistic for the model with the 5 variables was 0.80, indicating an optimum predictive capacity.

In order to assess what proportion of patients were appropriately classified with the model, a variable indicative of high probability of coronary arteries without significant stenosis was created. This consisted of being a woman (given the greater OR) and at least another 2 of the 4 remaining variables. This variable identified a coronary angiogram without significant stenosis with a specificity of 85% and a sensitivity of 53%.

**Prognosis of Acute Non-ST-Elevation Myocardial Infarction With Normal Coronary Arteries**

In the NSTEMI population, the lack of significant coronary artery stenosis was associated with a lower rate of death or infarction during follow-up (6% vs 27%; \( P = .0001 \)) after adjusting for other variables of prognostic value (HR=0.3; 95% CI, 0.2-0.9; \( P = .03 \)) (Figure). The other associated variables were heart failure on admission (HR=2.6; 95% CI, 1.7-4.0; \( P = .0001 \)), age (per year, HR=1.02; 95% CI, 1.0-1.04; \( P = .05 \)), diabetes mellitus (HR=1.5; 95% CI, 1.0-2.2; \( P = .04 \)), prior myocardial infarction (HR=1.6; 95% CI, 1.1-2.4; \( P = .03 \)), and renal failure (HR=1.7; 95% CI, 1.1-2.6; \( P = .02 \)).

The outcomes for the patients with NSTEMI and normal coronary arteries were compared with those of the control group. Table 3 shows the baseline differences between the 2 groups. Patients in the control group were older and more frequently had hypercholesterolemia and prior treatment with antiplatelet agents. Likewise, there was a tendency toward a greater proportion of men and systemic hypertension. During follow-up, among the entire population of patients with normal coronary arteries (n=224), 23 patients (10%) died or had an infarction. The death rates and rates of infarction in the subgroup with elevated troponin and normal troponin were 4 (6%) and 19 (12%) (HR=0.7; 95% CI, 0.2-2.2; \( P = .6 \)).

**DISCUSSION**

In our series, we found that 13% of the patients with NSTEMI lacked significant coronary stenosis according to the coronary angiogram procedure performed during the initial stay in hospital. Female sex, younger age, absence of diabetes, and prior antiplatelet treatment,
and presentation without ST-segment depression in the ECG were the variables associated with coronary angiography showing no significant lesions. The long-term prognosis of this population was good.

**Prevalence of NSTEMI and Normal Coronary Arteries**

The frequency of myocardial infarction and normal coronary arteries reported in the literature ranges from 8% to 12%. The differences in prevalence between the studies could depend on the different characteristics of the populations studied. Thus, in some series troponin is not used as a marker of necrosis, while others include patients if they have an ECG consistent with diagnosis of acute coronary syndrome in absence of elevated necrosis biomarkers. In addition, all studies represent the subgroup of patients with NSTEMI selected for catheterization, and the criterion used for this selection, whether broader or more restrictive, can also influence the angiographic results obtained. In the present study, diagnosis of infarction was done with troponin and catheterization was indicated in 62% of the patients with NSTEMI; this rate of catheterization is congruent with normal clinical practice in Spain for NSTEMI.

**Factors Predictive of Acute Non-ST-Elevation Myocardial Infarction With Normal Coronary Arteries**

Female sex was the variable most strongly associated with absence of significant coronary artery stenosis. Other associated factors were age less than 55 years, absence of diabetes, lack of prior antiplatelet treatment, and presentation without ST-segment depression in the ECG. The predictive model with these 5 variables showed an optimal discriminative power (C statistic = 0.80). In addition, we constructed a simple model consisting of an obligatory variable (female sex) and at least 2 of the 4 remaining variables. This simple model allowed us to identify a coronary angiogram without stenosis with good specificity (85%), although the sensitivity was low (53%).

Age and female sex are the predictive variables detected in other studies. In the PURSUIT study, a model was developed for predicting normal coronary angiography. This model included the variables of female sex, younger age, and absence of elevated necrosis markers, prior angina, diabetes, or ST-segment depression. In that study, unlike ours, patients were included without elevated necrosis markers.

**Prognosis**

The lack of significant coronary artery stenosis was associated with better prognosis within the NSTEMI population. While there is agreement about the lower risk in patients with NSTEMI who have normal coronary arteries, their prognosis in the long term is subject of some debate. Thus, some series question the good prognosis of these patients. In our series, the long-term prognosis was better than that of the remaining patients with NSTEMI.
and similar to those of patients with chest pain, without elevated troponin levels and coronary angiography showing no significant stenosis. These data reinforce the good prognosis of this entity.

Mechanisms

Several mechanisms might explain why some patients with NSTEMI had no significant stenosis in the coronary angiogram, such as a thrombotic process that has already resolved when catheterization is done, microvascular dysfunction, or coronary embolism.19,21-24 This could also underline substantial arterial disease that leads to an eccentric expansion of the atherosclerotic plaque that cannot be detected by coronary angiography.5-7 However, only a minority of patients have etiologic factors related to myocardial infarction and normal coronary arteries. Thus, a vasospasm test with ergonovine was positive in 15% of the patients and a coagulation disorder was detected in 12%.13

CONCLUSIONS

Clinical practice guidelines recommend an early invasive strategy in NSTEMI.25 In our study, we found coronary arteries without significant stenosis in 13% of the patients with NSTEMI and associated factors have been identified. Likewise, a predictive model has been constructed. Although this information could be relevant for assessing the strategy of systemic cardiac catheterization in certain patients with a high probability of normal coronary angiography, the predictive model has not been compared with any external cohort, and so it is hard to gauge its true usefulness.

Limitations

As the primary objective was to analyze the variables predictive of coronary arteries without significant stenosis, patients were excluded with a prior documented history of coronary artery stenosis. In addition, catheterization was indicated according to the judgment of the attending cardiologist. All these factors might have influenced the proportion of patients whose coronary angiogram showed no significant stenosis. Finally, the small number of patients in the group without significant coronary artery stenosis (n=224), where comparison is made between the subgroup with NSTEMI and the control group with normal troponin, means that any conclusions concerning the relationship with the prognostic value of troponin should be drawn with caution.

REFERENCES