Lack of a relationship between serum ferritin levels and coronary atherosclerosis evaluated by coronary arteriography

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Abstract

Many clinical and epidemiological studies have demonstrated the relationship between serum ferritin and ischemic heart disease. In the present study we evaluated the relationship between coronary heart disease (CHD) and serum ferritin levels in patients submitted to coronary arteriography. We evaluated 307 patients (210 (68.7%) males; median age: 60 years) who were submitted to coronary angiography, measurement of serum ferritin and identification of clinical events of ischemic heart disease. Serum ferritin is reported as quartiles. Ninety-six patients (31.27%) had normal coronary angiography (group 1) and 211 (68.73%) had coronary heart disease (group 2). Of the patients with CHD, 61 (28.9%) had serum ferritin levels higher than 194 ng/ml (4th quartile), as opposed to only 14 (14.58%) of those without CHD (P = 0.0067). In the 2nd quartile, 39 patients (18.48%) had normal coronary angiography (group 1) and 211 (68.73%) had coronary heart disease (group 2). Of the patients with CHD, 61 (28.9%) had serum ferritin levels higher than 194 ng/ml (4th quartile), as opposed to only 14 (14.58%) of those without CHD (P = 0.0067). In the 2nd quartile, 39 patients (18.48%) had normal coronary arteries (P = 0.00064). Multivariate analysis of the data showed that the difference between groups was not statistically significant (P = 0.33). We conclude that there is no independent relationship between coronary heart disease and increased levels of serum ferritin.

Introduction

Most deaths in the United States of America are due to ischemic heart disease (IHD) and several risk factors are known to be related to this condition (1-3). Among these factors, it has been demonstrated that total cholesterol and its low density lipoprotein (LDL) fraction are main factors in the genesis and development of coronary atherosclerosis and are related to the prognosis of the disease (4-6).

Atherosclerosis is a pathological process of the arterial wall characterized by deposition of lipids, especially LDL cholesterol, on the subendothelial layer inside mononuclear cells (foam cells) (7). It has been demonstrated that free oxygen radicals directly act on the endothelial cell membrane, determining alterations in its defense mechanisms. At
the same time, free radicals have an important action on lipid peroxidation, causing a modification of LDL and facilitating LDL deposition, with the consequent formation of atherosclerotic plaque (8-11). It has been further demonstrated that in vivo, at physiological pH, free radicals are produced at low rates through the reaction of Haber-Weiss. Nevertheless, the reaction may be accelerated when it is catalyzed by iron, with a greater production of free radicals (12-18).

This is one of the explanations for the greater incidence of ischemic events in males compared to females in the premenopausal state, since during this period the concentration of ferritin is three times higher in men than in women (19-29).

The putative role of iron in the development of atherosclerosis is based on clinical and epidemiological studies. However, no investigations correlating serum concentration of ferritin and anatomic diagnosis of coronary atherosclerosis by coronary arteriography are available in the literature. Thus, the objective of the present study was to evaluate the relationship between concentration of serum ferritin and presence of coronary atherosclerosis.

**Population and Methods**

In a transversal study we allocated 307 patients of both sexes with a previous diagnosis of IHD and with a formal indication of coronary arteriography from a physician. Before being submitted to hemodynamic studies, the patients signed an informed consent by which they authorized the collection of 10 ml of blood for laboratory tests. Patients were interviewed according to a standard questionnaire about the presence and clinical aspects of the different forms of angina pectoris. Coronary risk factors such as smoking, systemic arterial hypertension (SAH), diabetes mellitus (DM) and obesity were also investigated. Smoking was defined as the use of more than five cigarettes per day, at least for one year during the last five years; SAH and DM were defined as a previous medical diagnosis or the use of diet or tablets for their treatment; obesity was defined as a body mass index (BMI) higher than 27.3 for females and higher than 27.8 for males, as proposed by the National Institutes of Health Consensus Development (30). Normal values of total cholesterol and LDL cholesterol were considered to be less than 200 mg/dl and 130 mg/dl, respectively.

Coronary arteriography studies were performed according to the technique of Sones (31) or Judkins (32) after a 12-h fast. For statistical analysis of the sample, any coronary stenosis was considered as coronary heart disease (CHD), the major endpoint. Two experienced observers blindly interpreted the angiographic film.

Ferritin, glucose, total cholesterol and LDL cholesterol were measured in the blood sample. Serum ferritin was measured by enzyme immunoassay using a Bayer Stratus apparatus (Miami, FL, USA).

Levels of serum ferritin were analyzed as quartiles and the means were compared. The cut-off points used were as follows: 1st quartile - serum ferritin under 68 ng/ml; 2nd quartile - serum ferritin from 68 to 103 ng/ml; 3rd quartile - serum ferritin from 103 to 194 ng/ml; 4th quartile - serum ferritin higher than 194 ng/ml.

Statistical analysis was performed using the chi-square test, ANOVA for paired data, Kruskal-Wallis for non-parametric data and logistic regression for multivariate analysis. Statistical significance was considered as P<0.05.

**Results**

Of the 307 patients studied, 96 (31.27%) had normal coronary arteries (group 1) and 211 (68.73%) had CHD (group 2).

Sample characteristics are listed in Table 1. Risk factors for coronary heart disease were compared between the group of pa-
patients with CHD and those without CHD. A statistically significant difference was found between groups when considering age, sex, smoking, obesity, serum total cholesterol and serum LDL cholesterol by univariate analysis. When these variables were included in a logistic regression model, obesity and total serum cholesterol were not statistically significant.

Comparison of the serum ferritin quartiles between groups showed that of the patients without CHD 14 (14.58%) were in the 4th quartile, as opposed to 61 (28.9%) patients with CHD ($P = 0.0067$) (Figure 1). In the 2nd quartile, 39 patients (36.46%) had CHD, while 35 patients (18.48%) had normal coronary arteries ($P = 0.00064$). However, when this relationship was controlled for possible confounding factors such as age, sex, race, smoking, obesity, DM, SAH, total serum cholesterol and serum LDL cholesterol by multivariate analysis, no difference was observed ($P = 0.33$) (Figure 1).

**Discussion**

Based on results of experiments with animals, the hypothesis that high concentrations of iron in the organism are associated with accelerated free radical reaction and with increased lipid peroxidation seems to be plausible. More specifically, Balla and colleagues (33) demonstrated that the combination of physiological concentrations of hydrogen peroxide and the heme component of hemoglobin induces a rapid peroxidation of LDL *in vitro*, liberating free iron from this reaction. It has also been demonstrated that liberation of iron by ferritin stimulates the formation of hydroxyl radicals through superoxide radicals and hydrogen peroxide (34,35).

Sullivan (16,19,36) proposed a possible relationship between iron concentration and greater incidence of CHD in men and postmenopausal women. This hypothesis was based on the high incidence of CHD in patients with hereditary diseases of iron storage such as hemochromatosis, as well as the simultaneous increase of cardiovascular risk and serum iron in women after menopause and in old men. Nevertheless, Aronow (37) found no relationship between serum ferritin and CHD in men and women over 62 years.

The objective of the present study was to identify a possible association between serum ferritin concentration and coronary heart disease as evaluated by coronary arteriography. The choice of serum ferritin levels to evaluate iron concentration in the organism was based on previous studies which demonstrated that this is the best method to evaluate the actual condition of iron in the blood (38-40).

**Table 1 - Sample characteristics.**

<table>
<thead>
<tr>
<th></th>
<th>Coronary heart disease</th>
<th>Control patients</th>
<th>$P$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (mean ± SD, years)</td>
<td>61.6 ± 10.6</td>
<td>56.7 ± 11.4</td>
<td>0.0003</td>
</tr>
<tr>
<td>Race (white, %)</td>
<td>90</td>
<td>85.4</td>
<td>0.3</td>
</tr>
<tr>
<td>Sex (male, %)</td>
<td>68.7</td>
<td>43.7</td>
<td>0.00005</td>
</tr>
<tr>
<td>Smokers (%)</td>
<td>33.7</td>
<td>15.6</td>
<td>0.0018</td>
</tr>
<tr>
<td>SAH (%)</td>
<td>52.1</td>
<td>54.2</td>
<td>0.83</td>
</tr>
<tr>
<td>DM (%)</td>
<td>20.4</td>
<td>14.6</td>
<td>0.29</td>
</tr>
<tr>
<td>Serum cholesterol (mean ± SD, mg/dl)</td>
<td>190.9 ± 43</td>
<td>176.9 ± 39.1</td>
<td>0.007</td>
</tr>
<tr>
<td>Serum LDL cholesterol (mean ± SD, mg/dl)</td>
<td>122.6 ± 37.9</td>
<td>119.3 ± 82.4</td>
<td>0.018</td>
</tr>
<tr>
<td>Obesity (%)</td>
<td>13.7</td>
<td>26</td>
<td>0.013</td>
</tr>
<tr>
<td>Stable angina (%)</td>
<td>31.2</td>
<td>30.6</td>
<td>0.92</td>
</tr>
<tr>
<td>Unstable angina (%)</td>
<td>59.4</td>
<td>60</td>
<td>0.93</td>
</tr>
<tr>
<td>Atypical angina (%)</td>
<td>9.4</td>
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</table>

Figure 1 - Patient distribution according to ferritin quartiles. CHD = Coronary heart disease.
In this sample, main risk factors for CHD indeed proved to be higher in the group of patients with this condition. By analyzing our data about the main endpoint, we conclude that there is no independent relationship between serum ferritin and CHD. In spite of the evidence in literature for a direct role of iron in the pathogenesis of atherosclerosis, serum ferritin levels do not reflect this process. Probably there are local factors that determine the iron activity, independently of the systemic concentration of iron. Studies focusing on this subject should be performed to elucidate this important question.

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References


