Abstract

Objective: Fibromyalgia (FM) is a chronic pain syndrome with unknown etiology, which affects predominantly women. Mitochondrial alteration could have a role in the pathophysiological mechanisms of inflammatory conditions as FM and periodontitis. The aim of the present study was to assess the relationship between both diseases and mitochondrial dysfunction.

Patient and Methods: We studied the presence of periodontitis in twelve patients diagnosed with FM and mitochondrial dysfunction described. The diagnosis of FM was established according to ACR criteria and clinical symptoms were evaluated using the Fibromyalgia Impact Questionnaire (FIQ) and Beck Depression Inventory (BDI).

Results: Only one of the twelve patients included and agreed to participate in the study was diagnosed with periodontitis.

Conclusions: Pending studies with larger numbers of patients, we can conclude that mitochondrial dysfunction in FM is itself a condition not related with periodontitis. Periodontitis could be considered an exclusion criterion in all studies about mitochondrial dysfunction in patients.

Key Words: Periodontitis, fibromyalgia, mitochondrial dysfunction, oxidative stress.
Introduction
Fibromyalgia (FM) is a common chronic pain syndrome accompanied by other symptoms such as depression, anxiety, fatigue or sleep disturbances (1). The prevalence of FM in Spain is 2.4% in the general population, being significantly more frequent in females (4.2%) than in males (0.2%) (2). However, the pathogenic mechanism of the disease is still unknown.

In recent years, new researches have shown evidence showing that mitochondrial dysfunction and oxidative stress may play an important role in the pathophysiology of FM (3,4). Oxidative stress, as an event characteristic of mitochondrial dysfunction is involved in other diseases such as: atherosclerosis, hypertension, insulin resistance, heart failure, and the aging process (5). Recently, it has also seen the presence of mitochondrial dysfunction and oxidative stress in patients with periodontitis (6).

Periodontitis is a chronic infectious disease characterized by destructive inflammatory process affecting the tooth-supporting tissues resulting in loss of dentition. The cause is due to an ecological imbalance between the microbial biofilm on teeth and an impaired host inflammatory response. Oxidative stress is one of the main factors studied that could explain the pathophysiological mechanisms of inflammatory conditions that occur in the FM and periodontitis.

In order to demonstrate that mitochondrial dysfunction present in patients with FM is produced in the disease per se and not as a result of the presence of periodontitis, we propose in this paper the study of signs of periodontitis and the relationship with symptoms in a simple of patients with fibromyalgia.

Patient and Methods
We studied a sample of 20 patients of the Sevillian Fibromyalgia Association (AFIBROSE) who met the criteria of the American Collage of Rheumatology (ACR). Two were males and 18 females, with mean age of (SD) 50.8±8.6 years, with mean duration of disease of 13.65±9.19 years and a mean of 14.9±3.1 tender points. The parameters of mitochondrial dysfunction in these patients were previously described in detail (4). Exclusion criteria were: rheumatic concomitant diseases, significant systemic diseases (cardiopulmonary, neurological, renal or febrile), patients with severe psychopathology or psychoactive substance dependence and chronic pain patients of different origin to the FM. The study was performed with the informed consent of all participants and the approval of the local ethical committee. Patients did not take any drugs or nutritional or vitamin supplements for a period of 15 days prior to sampling and reported a sedentary lifestyle.

In patients, we evaluated the impact of the disease by testing the Spanish version of Fibromyalgia Impact Questionnaire (FIQ), with an interval of 0-80, pain by visual analog scale (VAS), with a range of 0-10, and depression using the standard version of the Beck Depression Inventory (BDI), with an interval of 0-63 (0-9 no depression; 10-18 mild depression; 19-29 moderate depression; greater than 30 severe depression).

A baseline periodontal examination was performed, and a single examiner collected full medical and dental histories. A single trained dental examiner recorded periodontal data. The periodontal probing depth (PD) and the recession of the gingival margin (GM) relative to the cementoenamel junction at six sites per tooth were recorded. Clinical attachment level (CAL) was calculated by adding recession to PD. PD and CAL were recorded to the nearest highest millimeter by means of the North Carolina periodontal probe (Hu-Friedy, Chicago, IL, USA), 15 mm in length and 0.35 mm in diameter. According to the criteria established by Machtei et al. (7), the clinical entity of periodontitis is based on the presence of CAL ≥ 6mm in two or more teeth and one or more sites with PD ≥ 5mm. Patients were divided into two groups: one with periodontitis (n = 1) and the other without periodontitis (n = 19). We also collected plaque and bleeding indices. The bleeding index of Van der Velden (8) was measured at six sites per tooth, while the plaque index of Silness and Löe (9) was valued at four sites per tooth.

Results
Patients showed high levels of pain determined by VAS, depression (BDI) and a high disease impact determined by the FIQ test (mean 7.2±2.2, 27±2.5 and 59.6±7.7, respectively) characteristic of the disease (Table 1). With respect to periodontal disease, only 1 of 20 patients who met the inclusion criteria and agreed to participate in the study were diagnosed with periodontitis. Table 2 summarizes the results of periodontal examination detailing of the parameters studied (GM, PD, CAL, dental plaque and bleeding on probing). As expected, the sample of patients with FM do not have periodontitis, with just one exception.

| Age (years) | 50.8 (8.6) | _ |
| Trigger points | 14.9 (3.1) | 0-18 |
| Disease duration (years) | 13.65 (9.19) | _ |
| Sex (female/male) | 18/2 | _ |
| BMI (Kg/m²) | 24.8 (2.1) | _ |
| FIQ total score, range 0 80 | 59.6 (7.7) | 0-80 |
| VAS total score, range 0 10 | 7,2 (2,2) | 0-10 |
| BDI total score, range 0 63 | 27 (2,5) | 0-63 |

BDI: Beck depression inventory; BMI: body mass index; FIQ: fibromyalgia impact questionnaire; VAS: visual analog scale. All values are mean (SD).

Table 1. Clinical characteristics of patients
can say that mitochondrial dysfunction found in a sample of patients with fibromyalgia is characteristic of the disease and does not occur as a result of the presence of periodontal disease.

On the other hand, another factor to consider in future research is that the presence of periodontitis should be considered as an exclusion criteria for any study whose objective the presence of mitochondrial dysfunction.

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