Macrophage migration inhibitory factor gene polymorphisms as exacerbating factors of apical periodontitis

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Abstract

Background. Two polymorphisms in the macrophage migration inhibitory factor (MIF) gene have been associated with inflammatory diseases (-794 CATT5–8 and -173G>C); however, so far there are no reports of studies related to oral health.

Objectives. To genotype the -794 CATT5–8 and -173G>C MIF polymorphisms in Mexican patients with apical periodontitis as a genetic risk of exacerbation.

Material and methods. The study involved 120 patients with apical periodontitis: 60 with a diagnosis of acute apical periodontitis (Group A) and 60 without previous episodes of exacerbation (Group B). Allelic discrimination was performed from peripheral blood DNA; the repeat polymorphism -794 CATT5–8 was genotyped with sequencing, while the -173G>C polymorphism was determined using real-time polymerase chain reaction (RT-PCR) using TaqMan probes. The associations between MIF polymorphisms, haplotypes and the risk of exacerbated apical periodontitis were assessed.

Results. The allele CATT7 was associated with the risk of a stage of acute inflammation (OR = 4.13; 95% CI = 1.82–9.63; p = < 0.001). Regarding the -173G>C polymorphism, a process of inflammation exacerbation was only associated with the CC genotype (OR  = 4.1; 95% CI = 1.02–20.84; p = 0.045). The analysis of the haplotype showed that the combination CATT7/C increases the risk of exacerbation of apical periodontitis (OR = 3.57; 95% CI = 1.038–13.300; p = 0.021).

Conclusions. The polymorphisms -794 CATT5–8 and -173G>C MIF seem to significantly influence the development of a state of exacerbated inflammation in patients with apical periodontitis.

Key words: apical periodontitis, genetic polymorphisms, macrophage migration inhibitory factor
Introduction

Periodontitis is an inflammatory condition which affects areas such as the gingiva, periodontal ligaments and alveolar bones because of viral or bacterial infections. In acute episodes, there is pain because of an increase in hypersensitivity to touch.\(^1\) According to several studies, the prevalence of periodontitis increases with age and could be a problem in more than 60% of the world population.\(^2\)\textsuperscript{-4} When acute inflammation becomes apparent in the tissue, cells of the immune system such as macrophages, lymphocytes and plasma cells are wrapped in collagenous connective tissue as part of the antimicrobial response. As a result, the cytokines block not only the osteoclastic activity but also bone reabsorption, leading to asymptomatic latent or inactive granuloma without visible changes in a radiographic image.\(^1\)

When the balance of the periapical area is broken, bacterial proliferation can be triggered towards the radicular conduct and periapical tissues, with subsequent exacerbation of chronic periodontitis, which manifests itself in the formation of a secondary abscess.\(^5\) In this process, interleukins such as tumor necrosis factor α (TNF-α), interleukin 1b (IL-1b), IL-6, and IL-8 intervene as mediators of inflammation.\(^6\) The macrophage migration inhibitory factor (MIF) is another regulatory cytokine of the innate immune response expressed by macrophages, monocytes, B cells and dendritic cells as well as granulocytes.\(^7\)\textsuperscript{,8}

The MIF is considered an important component in the defense against bacterial infections, since this cytokine promotes the release of pro-inflammatory molecules.\(^9\)\textsuperscript{,10}; however, its overexpression could lead to an acute exacerbated response. In fact, MIF has been used as a biomarker of diseases with a relevant inflammatory component.\(^11\)\textsuperscript{,12}

Two polymorphisms located in the promoter region of the MIF gene are associated with inflammatory diseases. First, the short tandem repeat CATT\(_{5-8}\) is a tetrancleotide that is repeated between 5 and 8 times in position -794; increases in the numbers of repetitions produce a corresponding increase in MIF promoter activity.\(^13\) On the other hand, C-173G, a single nucleotide polymorphism (SNP), acts to enhance the union of the transcription factor AP4 and the subsequent overexpression of MIF.\(^14\) Both polymorphisms have been studied as factors related to the pathogenesis of sepsis, severe acute respiratory syndrome, asthma, arthritis, glomerulonephritis, and inflammatory bowel disease.\(^12\)\textsuperscript{,15}

However, we have found no scientific reports that associate MIF polymorphisms with acute inflammatory processes in the oral cavity. Therefore, the goal of this study was to evaluate the risk of exacerbation in Mexican patients with a diagnosis of apical periodontitis by analyzing the distribution of alleles, genotypes and haplotypes in 2 groups: patients without previous episodes of exacerbation, and patients with a medical history of exacerbation.

Material and methods

Characteristics of the study groups

A cross-sectional study of 120 patients including 46 males (38.3%) and 74 females (61.7%) between 18 and 72 years of age diagnosed with apical periodontitis at the endodontic clinic at the University of San Luis Potosí, Mexico, was carried out. Anthropometric data and clinical histories were collected from all patients. The individuals were classified into 2 groups according to clinical assessment.

Group A included 60 patients with pulp necrosis associated with acute apical abscess or phoenix abscess. Symptoms could vary from moderate to severe, including intraoral/extraoral edema, exudate, tumefaction, and pain on palpation or percussion. In severe cases, fever and/or lymphadenopathy could also be observed. When the diagnosis was acute apical abscess, no periapical lesion could be detected radiographically, while for a phoenix abscess, a periapical lesion could be greater than 2 mm.

Group B also included 60 patients with chronic apical periodontitis, but without a previous history of exacerbation. A negative response to a cold thermic test was diagnosed as pulp necrosis. The patients could have mild or no pain on palpation or percussion. Radiographically, a periapical lesion with a diameter wider than 2 mm needed to be detected.

This project was previously approved by the bioethics committee at the Autonomous University of San Luis Potosí (approval No. CEJ-FE-009-014), and informed consent was obtained from all the participants prior to initiating our study. All the procedures performed were in accordance with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards.

DNA purification

DNA was isolated from blood obtained by venous puncture and preserved at –80°C in accordance with a previously described protocol.\(^16\) Briefly, 1 mL of blood was treated with a pH 7.5 lysis solution (0.3 M sucrose, 10 mM Tris–HCl pH 7.5, 5 mM MgCl\(_2\), and 1% Triton X-100). Leukocytes were obtained with centrifugation and washed with the lysis solution. Subsequently, the pellet was suspended in 10 mM Tris–HCl pH 8 and lysed with 20 mg/mL lysozyme detergent. Proteins were precipitated with 5 M NaCl and the supernatant was treated with cold 96% ethanol. The DNA was washed with 70% ethanol and suspended in nuclease-free water. The samples were analyzed with the help of a UV-spectrophotometer at 260 nm and 280 nm. The ratio of the absorbance at 260 nm and 280 nm (A260/280) was used as indicator of purity of DNA; a value between 1.4 and 2 was considered acceptable. The DNA samples were standardized to a final concentration of 30 ng/µL and frozen until use. All the reagents were purchased from Sigma-Aldrich (St. Louis, USA).
Genotyping

CATT\textsubscript{5–8} repetitions in the position -794 of the MIF gene were identified with sequencing from a polymerase chain reaction (PCR) product obtained using the following oligonucleotides: (F) 5′-TGTCTCTTCTCTGCTATGTC/(R) 5′-CCTAATGTTAAACTCGGGG -3′.\textsuperscript{17} The final volume of each PCR was 25 µL with the following composition: 5 µL 5 x iProof\textsuperscript{TM} HF Buffer (BioLabs, Cambridge, USA), 120 ng of DNA and 0.125 µL of Taq polymerase (1.25 units). For each primer, the final concentrations were 200 nM and 200 µM for deoxynucleotide triphosphates (dNTPs) (BioLabs). The amplification conditions were 1 cycle at 95°C for 1 min; 35 cycles at 95°C for 30 s, 60°C for 30 s and 72°C for 30 s; and finally, 1 cycle at 72°C for 2 min. All PCR assays were performed in duplicate in a T100™ Thermal Cycler (Bio-Rad Laboratories Inc., Hercules, USA). The amplification products were previously verified in 3% agarose gel electrophoresis, and the sequencing service was provided by LANBAMA (IPICYT, San Luis Potosí, Mexico).

The MIF -173G>C polymorphism was detected with real-time PCR (RT-PCR) using allele-specific oligonucleotide probes for both wild-type and mutant alleles, labeled with different fluorescent tags. The primers used were as follows: (F) 5′-CCAGCAACCGCCTAAG-3′/(R) 5′-TGCCGACCTAAACATCGGTGA-3′, and the probe for the -173G allele was [Q705]- ACCGCTCCAACCTGTT-[BHQ2], while the probe for the -173C allele was [Cy5]- CCGCTCAAAGCTCATT-[BHQ2]. The primers and probes were designed by RealTimeDesign software and purchased from BioSearch Technologies Inc. (Petaluma, USA). The qPCR was carried out with 30 ng of genomic DNA. Each sample (10 µL) was analyzed in duplicate and contained 5 µL of iQTM Multiplex Powermix (Bio-Rad) and 300 nmol of each primer and probe. The PCR cycles were programmed on a CFX96 Touch Real-time PCR detection system (Bio-Rad) and consisted of hot-start incubation (95°C, 3 min) and amplification for 45 cycles (95°C, 15 s; 59°C, 1 min). Genotypes were distinguished by post-read PCR fluorescence of normalized reported values for wild-type and mutant alleles. The genotype of each sample was determined using a multicomponent algorithm, generating 3 allelic clusters: GG, GC and CC genotypes.

Statistical analysis

Geometric means and standard deviations (SD) were calculated for continuous variables, percentages by gender and frequencies for alleles, genotypes and haplotypes. Comparisons of means between the 2 study groups were analyzed using the Mann–Whitney U test, while differences in the distribution of alleles and genotypes were analyzed using the χ\textsuperscript{2} test. The Hardy–Weinberg equilibrium for the alleles was measured using the χ\textsuperscript{2} test, and the linkage disequilibrium (LD) between the loci was calculated using a 2-locus LD calculator.

The association magnitude was quantified with odds ratios (OR) with 95% confidence intervals (95% CI) to calculate the risk of exacerbation of apical periodontitis. A binomial logistic regression analysis was used after categorization by alleles, genotypes and haplotypes, considering the clinical diagnosis as a dependent variable with a binary outcome (acute vs chronic apical periodontitis). Results were considered statistically significant when p < 0.05. All the analyses were performed with an IBM SPSS v. 19.0 statistic software package (IBM Corp., Armonk, USA).

Results

A total of 120 patients with diagnoses of apical periodontitis were included in this study, 50% with a history of at least 1 acute episode (Group A) and the other half with chronic apical periodontitis without exacerbations (Group B). The percentages of women and men were 62.7 and 37.3, respectively; the average age was 36.4 years. With respect to the body mass index (BMI), 43.2% corresponded to normal weight, 35.2% to overweight and 21.6% to obesity. Age, the proportion of women and men and BMI did not differ between the 2 groups when the data was statistically analyzed. These results are summarized in Table 1.

The allelic and genotyping frequencies, as well as the distribution of them between the 2 study groups, are shown in Table 2. In the case of the MIF -794 CATT\textsubscript{5–8} polymorphisms, frequencies of 0.23, 0.57 and 0.2 were found for CATT\textsubscript{5}, CATT\textsubscript{6} and CATT\textsubscript{7}, respectively; the CATT\textsubscript{8}
allele was not found in any of the participants. Regarding the -173G/C polymorphism, the frequencies for G and C alleles were 0.625 and 0.375, respectively, following the Hardy–Weinberg equilibrium ($\chi^2 = 2.28$, $p = 0.13$). Both MIF polymorphisms were found in linkage disequilibrium ($D' = 0.635$, $r^2 = 0.179$, $\chi^2 = 17.72$). When the distribution of alleles and genotypes of -794 CATT and -173G/C variants were analyzed between the 2 study groups, we found that only the CATT polymorphism had a statistically significant distribution between the groups ($p = 0.001$). Odds ratios were measured for MIF polymorphisms according to the classification of apical periodontitis (acute vs chronic). As shown in Table 3, an association was established in individuals carrying the alleles -794 CATT<sub>5</sub> (OR = 2.8; 95% CI = 1.44–5.62; $p = 0.002$) and CATT<sub>7</sub> (OR = 4.13; 95% CI = 1.82–9.63; $p < 0.001$), while no association with the -173C allele was found. The analysis by genotype showed that the CATT risk alleles are associated with acute episodes of apical periodontitis only in a homozygous model. In fact, CATT<sub>7/7</sub> had a greater strength of association and increased odds with respect to CATT6/6 (OR = 4.39, 95% CI = 1.09–22.88 and $p = 0.03$ vs OR = 6.65, 95% CI = 1.29–42.49 and $p = 0.02$). In the case of the -173C/C genotype, an association was observed in a homozygous model (OR = 4.1; 95% CI = 1.02–20.84; $p = 0.045$) as well as a recessive model (OR = 3.9; 95% CI = 1.06–18.50; $p = 0.019$).

An analysis of haplotypes was conducted considering the -794CATT<sub>5</sub>-173G/C combination as a reference (Table 4).

### Table 2. Allelic and genotypic frequencies of -794 CATT<sub>5–8</sub> and G-173C MIF polymorphisms in the study group

| MIF polymorphism | Total n (%) | Group A (AP*) n | Group B (CP*) n | p-value<sup>ab</sup> |
|------------------|-------------|----------------|----------------|-----------------|
| -794 CATT<sub>5–8</sub> |             |                |                |                |
| Allele | | | | |
| CAT T<sub>5</sub> | 55 (23) | 16 | 39 | 0.001 |
| CAT T<sub>6</sub> | 136 (57) | 73 | 63 | |
| CAT T<sub>7</sub> | 49 (20) | 31 | 18 | |
| CAT T<sub>8</sub> | 0 | 0 | 0 | |
| Genotype | | | | |
| CAT T<sub>5</sub>/5 | 12 (10) | 3 | 9 | 0.060 |
| CAT T<sub>5</sub>/6 | 26 (22) | 8 | 18 | |
| CAT T<sub>5</sub>/7 | 5 (4) | 2 | 3 | |
| CAT T<sub>6</sub>/6 | 50 (42) | 30 | 20 | |
| CAT T<sub>6</sub>/7 | 10 (8) | 5 | 5 | |
| CAT T<sub>7</sub>/7 | 17 (14) | 5 | 12 | |
| G-173C | | | | |
| Allele | | | | |
| G | 150 (62.5) | 68 | 82 | 0.0125 |
| C | 90 (37.5) | 50 | 40 | |
| Genotype | | | | |
| GG | 43 (36) | 19 | 24 | 0.0122 |
| GC | 64 (53) | 30 | 34 | |
| CC | 13 (11) | 10 | 3 | |

<sup>ab</sup>χ² test; *AP – acute periapical periodontitis; CP – chronic periapical periodontitis.

### Table 3. Analysis of the association of MIF alleles and genotypes with the risk of exacerbation of periapical periodontitis

| MIF polymorphisms | OR (95% CI) | p-value |
|-------------------|-------------|---------|
| MIF alleles | | |
| -784 CATT<sub>5–8</sub> | | |
| CATT<sub>5</sub> | 1 | – |
| CATT<sub>6</sub> | 2.8 (1.44–5.62) | 0.002 |
| CATT<sub>7</sub> | 4.13 (1.82–9.63) | <0.001 |
| -173G/C | | |
| G | 1 | – |
| C | 0.81 (0.52–1.28) | 0.18 |
| MIF genotypes | | |
| -784 CATT<sub>5–8</sub> | | |
| 5/5 | 1 | – |
| 5/6 | 1.32 (28.00–7.48) | 0.74 |
| 5/7 | 1.91 (0.16–20.30) | 0.58 |
| 6/6 | 1 | – |
| 6/7 | 4.39 (1.09–22.28) | 0.03 |
| 7/7 | 4.1 (1.02–20.84) | 0.045 |
| -173G/C | | |
| GG co-dominant | 1 | – |
| GC | 1.18 (0.54–2.60) | 0.67 |
| CC | 4.1 (1.02–20.84) | 0.045 |
| GG dominant | 1 | – |
| GC+CC | 1.36 (0.64–2.91) | 0.42 |
| GG+GC recessive | 1 | – |
| CC | 3.9 (1.06–18.50) | 0.019 |

OR – odds ratio.

### Table 4. Haplotype frequencies of the -784 CATT<sub>5–8</sub> and -173G/C MIF polymorphisms and the risk of exacerbation of periapical periodontitis

| Haplotype | Acute cases | Chronic cases | OR (95% CI) | p-value |
|-----------|-------------|---------------|-------------|---------|
| CATT<sub>5</sub>/G | 10 | 27 | 1.6 (0.67–4.16) | 0.13 |
| CATT<sub>5</sub>/GC | 42 | 43 | 2.15 (0.85–5.58) | 0.046 |
| CATT<sub>7</sub>/G | 18 | 12 | 0.99 (0.28–3.42) | 0.49 |
| CATT<sub>7</sub>/GC | 30 | 20 | 2.51 (0.96–6.83) | 0.03 |
| CATT<sub>7</sub>/CC | 13 | 6 | 3.57 (1.03–13.3) | 0.021 |

Total 120 | 120 | – | – |
In our results, the CATT<sub>7</sub>/G and CATT<sub>6</sub>/C haplotypes had statistically significant associations with exacerbations of periodontitis (OR = 2.15, 95% CI = 0.85–7.58 and p = 0.046 vs OR = 2.51, 95% CI = 0.96–6.83 and p = 0.03, respectively). However, the OR for the CATT<sub>7</sub>/C haplotype was greater than the other 2 combinations, taking into consideration that both MIF alleles were associated with a higher risk of inflammation (OR = 3.57; 95% CI = 1.038–13.300; p = 0.021).

**Discussion**

Both MIF -794 CATT<sub>5</sub>–<sub>8</sub> and -173G>C polymorphisms have been related to the risk of increasing the severity of the inflammatory response. Consequentially, we studied both variants as genetic risk factors for exacerbation of inflammation in patients with apical periodontitis. When patients with acute apical periodontitis and chronic apical periodontitis were compared, no correlations were observed in terms of age, sex or BMI in either study group; therefore, possible effects of age, sex or obesity on the exacerbation of periodontitis were ruled out (Table 1). The most frequent allele was CATT<sub>5</sub> (57%), while the CATT<sub>6</sub> allele was not detected, which is consistent with previous reports carried out in the Mexican population. Regarding MIF -173G>C polymorphism, a frequency of 37.5% for risk allele C was found. This was slightly higher than previously reported for the Mexican population, in which a value around 30% has been observed; however, those studies were carried out in other geographical regions of Mexico. The -794 CATT alleles showed a significant difference in distribution between the 2 groups, in contrast with the -173G>C allele, which did not show a different proportion between the acute and chronic groups (Table 2).

The genetic risk of exacerbated apical periodontitis is analyzed in Table 3, noting that CATT<sub>5</sub> and CATT<sub>7</sub> carriers have almost 3 or 4 times higher potential risk of developing an episode of acute apical periodontitis than CATT<sub>6</sub> carriers do; however, either a homozygous condition or CATT<sub>6</sub>/7 combination seems to be necessary for an acute episode. Indeed, the homozygous carriers of CATT<sub>7</sub> showed the highest risk of exacerbation, with an OR value of 6.6 (95% CI = 1.29–42.49; p = 0.02). In fact, the CATT<sub>6</sub>/7 genotype has been considered a vulnerability factor for other health risks in the Mexican population, such as acute coronary syndrome and rheumatoid arthritis.

On the other hand, risk allele C for the MIF -173 polymorphism was only associated with acute apical periodontitis in a homozygous condition (OR = 4.1; 95% CI = 1.02–20.84; p = 0.045), but has also been observed in a recessive model where GG+GC genotypes were used as references (OR = 3.9; 95% CI = 1.064–18.500; p = 0.019). Very recently, several studies have reported that the C allele in the MIF -173 position represents a genetic risk factor associated with diverse disorders or diseases, such as osteoporosis; pulmonary arterial hypertension in patients with systemic sclerosis; tuberculosis; fibrosis in biliary atresia patients; breast cancer, especially among older patients; childhood asthma; and autoimmune hepatitis with acute symptomatic presentation. However, differences in allele frequency among racial groups is a factor to be considered when interpreting the results. In the present study, none of these diseases was reported by carriers of the -173C allele.

When a haplotype analysis was performed considering all the allele combinations between -794 CATT and -193 G>C MIF polymorphisms (Table 4), the association of CATT<sub>7</sub/>CATT<sub>6</sub> showed the greatest risk of exacerbating periodontitis, which concurs with other studies where the CATT<sub>7</sub>/CATT<sub>6</sub> haplotype has been shown to be related to an increased inflammatory process. Particularly, the CATT<sub>7</sub>/CATT<sub>6</sub> combination has been considered a risk factor for death in carriers with severe sepsis, as well as a risk factor for inflammatory polyarthritis.

Since MIF -794 CATT<sub>7</sub> and -173C polymorphisms can modify the gene expression and therefore the production of MIF, carriers of this allele combination would have higher levels of circulating MIF, with a consequent predisposition to an increased response to inflammatory processes; however, plasma MIF levels were not determined in this study. More investigations into the role of MIF in the development and severity of apical periodontitis as well as the genetic predisposition to exacerbation are needed.

**Conclusions**

Although several studies have shown that MIF plays a role in the pathogenesis of apical periodontitis as a promoter of other pro-inflammatory molecules, MIF polymorphisms have only been studied in systemic types of inflammatory disorders, such as sclerosis, tuberculosis, fibrosis, and asthma. Very few studies have looked at MIF polymorphisms in diseases of the oral cavity. There are few reports in the literature about polymorphisms in pro-inflammatory cytokine genes related to apical periodontitis. This study is the first report where a positive association has been found between acute stages in patients with apical periodontitis and -794 CATT<sub>7</sub> and MIF -173C polymorphisms.

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