Case Report

CTCFL (BORIS) mRNA Expression in a Peripheral Giant Cell Granuloma of the Oral Cavity

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Peripheral giant cell granuloma (PGCG) is a relatively common benign reactive lesion of the oral cavity which can occur at any age; it consists of a soft tissue injury mainly originating from the periosteum or periodontal membrane following local irritation or chronic trauma. The main clinical feature of PGCG is a red-purple nodule located in the region of the gums or the alveolar edentulous, mainly in the lower jaw [1, 2].

PGCG is of osteoclastic nature and is not considered a true neoplasm. It has been termed as PGCG “abnormal reparative”; however, this function has not been fully established, and its osteoclastic activity seems doubtful [3]. Although the presence of calcitonin membrane receptors, as well as osteoclastic activity, has been demonstrated by immunohistochemistry [4, 5], some authors report that the lesion is formed by cells of the fagocito mononuclear system [6]. In this respect, some authors propose that PGCG is a process in which the fibroblasts overexpress cytokines and growth factors, which induce or activate macrophages to become giant cells [7].

CTCF (CCCTC-binding factor) is an essential protein encoded by the gene of the same name, which is ubiquitously expressed and plays an important role in the regulation of gene expression, genomic imprinting, and nuclear chromatin insulators regulation. BORIS expression promotes cell immortalization and growth while CTCF has tumor suppressor activity; the expression pattern may reflect the reverse transcription silencing of BORIS. The aim of this work was to describe a histopathological and molecular approach of an 8-year-old pediatric male patient with PGCG diagnosis. It was observed that the PGCG under study expressed CTCF as well as BORIS mRNAs alongside with the housekeeping gene GAPDH, which may be related to possible genetic and epigenetic changes in normal cells of oral cavity.

1. Introduction

Peripheral giant cell granuloma (PGCG) is a relatively common benign reactive lesion of the oral cavity which can occur at any age; it consists of a soft tissue injury mainly originating from the periosteum or periodontal membrane following local irritation or chronic trauma. The main clinical feature of PGCG is a red-purple nodule located in the region of the gums or the alveolar edentulous, mainly in the lower jaw [1, 2].

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CTCF (CCCTC-binding factor) is an essential protein encoded by the gene of the same name, which is ubiquitously expressed and plays an important role in the regulation of gene expression; the multiple activities of CTCF in mammals include transcriptional activation and repression, gene silencing, constitutive chromatin insulation, and functional reading of imprinted states, reasons for which it has been named the master weaver of the genome [8]. CTCF has been proposed as a novel tumor suppressor gene because the CTCF expression suppresses tumor cell proliferation [9].
Some mutations in the *CTCF* gene have been localized and characterized in various types of cancer, including breast [10]. Recently was identified a parologue of the *CTCF* gene that was called *CTCFL* (*CTCF* like) and also BORIS (Brother of the Regulator of Imprinted Sites, name that will be used in this paper), which encodes a protein of the same name [11]. *CTCF* and BORIS are zinc-finger proteins sharing the same II zinc fingers and interact with similar DNA sequences, although they have different amino and carboxyl ends. Normally *CTCF* and BORIS are expressed in a mutually exclusive pattern that correlates with the reestablishment of methylation marks during male germ cell differentiation [11, 12].

Apart of male germ cells, scarce or null expression of BORIS has been reported in normal human tissues and cells. Thus, it has been suggested that BORIS expression is controlled at epigenetic level by promoter DNA methylation, and their activation requires demethylation [13]. In cancer patients, high expression of BORIS is correlated with the size and grade of the tumor, especially in breast, endometrium, prostate, and colon [14]. An inhibitor of DNA methylation, 5-az-2′ deoxy-cytidine (5-azadC), as well as histone deacetylase inhibitors, induces or enhances the BORIS gene expression in various carcinoma cell lines [15].

**2. Case Report**

An 8-year-old pediatric male patient, referred from private practice to the Department of Clinical Pathology of the Faculty of Dentistry of the Juarez University of the State of Durango, Mexico, presented a lobular nodular lesion at the level of dental organs 31 and 32, similar in color to the mucosa, with a pediculated base in the gingival region, with an evolution time of approximately 6 months. The patient underwent excisional biopsy; the specimen was grayish-white, irregularly shaped, and had a firm consistency. The measures of the specimen were 1.1 × 0.6 × 0.7 cm. The sample was formalin-fixed and paraffin-embedded (FFPE) for histopathological and molecular analysis.

**Microscopic Description.** H&E staining was performed and analyzed with a Leica DMD108 Optic Microscopic. By H&E staining a stratified squamous parakeratinized epithelium, with the presence of elongated epithelial anastomosing nails and pseudoepitheliomatous hyperplasia, was observed. Epithelial atrophic areas presenting a continuum with the stroma, dense fibrous hyalinized connective tissue, heavy hemorrhagic areas, blood vessels, and the presence of multinucleated giant cells, some of which with vesicular nuclei and abundant mitotic figures, were also observed. Furthermore, a lymphoplasmocitary inflammatory response dominated by positive surgical margins was present (Figure 1).

**Molecular Description.** A descriptive study of a FFPE sample with histopathologic diagnosis of PGCG was conducted. Total RNA was obtained with the QuickExtract FFPE RNA Extraction Kit (Illumina, San Diego, CA) and cDNA was synthesized with the cDNA Synthesis Kit iScript (BIO-RAD). BORIS, CTCF, and GAPDH (used as housekeeping gene) mRNA expression was carried out by real-time PCR using the QuantiTect SYBR Green PCR kit system and Quantitect Primers (Qiagen Germantown, MD. Catalogue numbers: BORIS, QT00023191, amplicon size 125 bp; CTCF, QT00045437, amplicon size 119 bp; GAPDH, QT01192646, amplicon size 119 bp).

The amplification conditions consisted of incubation at 50°C for 30 min, followed by polymerase activation at 95°C for 15 min and 45 cycles of denaturation at 94°C for 15 s, and annealing and extension at 60°C for 1 min. Amplification, acquisition, and data analysis were performed with an Eco Illumina Real-Time PCR System and Eco Version 0.1753.9 Software.

The specificity of the reaction was verified by melting analysis of the amplified products; the size of the amplicons was analyzed by electrophoresis in agarose gels stained
4. Conclusion

The coexpression of BORIS and CTCF genes in PGCG may be associated with the alteration of genetic and epigenetic mechanisms in the normal function of the cells in the oral cavity.

Conflict of Interests

The authors have no conflict of interest with respect to the publication of this paper.

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