An Unusual Presentation of Peripheral Ossifying Fibroma in an Elderly Male - A Case Report

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INTRODUCTION

Peripheral ossifying fibroma (POF) is a reactive process of the gingiva that develops due to irritation or minor trauma. Females are more affected than males suggesting a hormonal influence. This case report describes a case of peripheral ossifying fibroma in a 48-year-old male patient in 33, 34 regions in contrast to its common occurrence in young females. Histopathological examination is necessary to confirm the diagnosis. Surgical excision is the treatment of choice to prevent recurrence. One year follow up of the case showed no signs of recurrence.

Gingival overgrowth is a common feature of the various types of gingival disease in the oral cavity. Gingival diseases and conditions can be due to genetic disorder, specific infections, inflammatory and immune conditions and lesions, reactive processes, neoplasms, endocrine, nutritional, and metabolic diseases, traumatic lesions and gingival pigmentation. Reactive processes are non-neoplastic nodular swellings that develop in response to local irritation or minor trauma. The term epulis is exophytic processes confined to the gingiva. Kfir et al. classified epulides into fibrous epulis, peripheral ossifying fibroma, pyogenic granuloma (vascular epulis), peripheral giant cell granuloma (or central).

Ossifying fibromas in the oral cavity can be classified into central and peripheral type. The central type expands from the medullary cavity of the bone, arising from the endosteum or the periodontal ligament (PDL) adjacent to the root apex. Peripheral type arises from the soft tissues overlying the alveolar process which is contiguous with the periodontal ligament.⁴ This article presents a case of POF in a male patient.

PRESENTATION OF CASE

A 48-year- old male patient reported to the department of periodontics with the chief complaint of growth on the gums in relation to the lower left canine-premolar region since 4 months. Initially, it started with a small nodule and gradually increased to the present size. Growth was associated with bleeding on brushing and discomfort during mastication. (Figure 1) There was no relevant medical and dental history and patient did not give any history of trauma.

On intraoral examination, there was a solitary, reddish-pink, exophytic mass measuring about $0.9~\rm cm~x~0.6~\rm cm~x~0.5~cm$ with a yellowish patch on the surface. The growth was emerging from the buccal interdental papilla. It was extending from distal of 33 to mesial of 34, up to the level of the occlusal surface (Figure 2). The growth was firm with tenderness on palpation. There was no mobility of associated teeth. Therefore, a provisional diagnosis was given as irritation fibroma.

An intraoral periapical radiograph (IOPA) revealed no findings pertaining to the lesion. (Figure 3)

DIFFERENTIAL DIAGNOSIS

The differential diagnosis included irritation fibroma, pyogenic granuloma and peripheral giant cell granuloma.

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DISCUSSION OF MANAGEMENT

One week after phase-I therapy, there was no change in the size of the gingival overgrowth (Figure 4). After routine blood investigations, an excisional biopsy was done using the scalpel down to the periosteum. (Figure 5) The tissue was then sent for histopathological examination.



Figure 1.
Pre- Operative
- First Visit



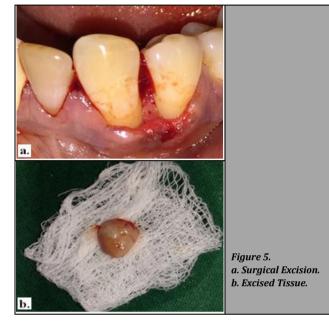
Figure 2. Solitary, Reddish-Pink, Exophytic Mass Measuring about 0.9 cm x 0.6 cm x 0.5 cm with a Yellowish Patch on the Surface

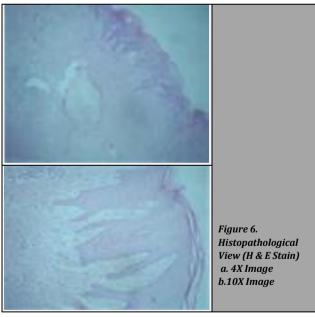


Figure 3. IOPA IRT 33, 34



Figure 4. One Week after Phase-I Therapy





Histological examination of the haematoxylin & eosin (H & E) stained sections revealed epithelium and underlying connective tissue. The epithelium was stratified squamous parakeratinized epithelium of varying thickness. Areas of ulceration were seen with fibrous exudates and numerous capillaries with dense acute and chronic inflammatory cells chiefly neutrophils and lymphocytes. The underlying connective tissue was fibrous with dense bundles of collagen fibers and blood vessels. Focal calcification of varying size and number was evident and was surrounded by plump

fibroblasts. (Figure 6) Thus, a final diagnosis of peripheral ossifying fibroma with respect to 33 and 34 regions was established.

After 15 days, the surgical site showed satisfactory healing. The case was followed up for a period of 1 year, and had shown normal healing with no signs of recurrence. (Figure 7)



Figure 7. (a) 15 Days Post-Operative View (b) 1 Year Post-Operative View

DISCUSSION

POF is a solitary, slow-growing, nodular gingival overgrowth originating from cells of the PDL and periosteum.⁵ It occurs mostly on the maxillary and mandibular gingiva, anterior to the molars emerging from the interdental papilla.⁶ In this case report, the origin of overgrowth was considered as cells of PDL because of the proximity of gingiva (interdental papilla) to PDL and the presence of dense collagen fibers.

In 1972, Eversole and Rovin described POF as a reactive lesion of gingiva. In 1982, Gardner coined the term POF.⁷ A lesion may develop as a result of trauma or local irritants such as subgingival plaque and calculus, dental appliances, faulty dental restorations, food lodgements and iatrogenic factors.⁸ In this case, it could be due to the presence of subgingival plaque and calculus which was evident on the first visit. The lesion is usually smaller than 1.5 cm in diameter with an ulcerated surface. The base of the lesion is either pedunculated or sessile.⁹ This lesion generally affects females in the second or third decade of life.¹⁰ However, in this case, POF occurred in an aged male patient which contradicts the concept of hormonal influence in the aetiology of the disease.

In this case, the clinical features led to the differential diagnosis of irritation fibroma, pyogenic granuloma or peripheral giant cell granuloma. However, confirmatory diagnosis of POF was made based on the histopathological examination which is characterized by ulcerated stratified squamous surface epithelium, a high degree of cellularity, acute and chronic inflammatory cells, and dystrophic calcification. Pyogenic granuloma is more vascular than POF. Peripheral giant cell granuloma is less cellular and contains giant cells when compared to POF. Various other terminologies of POF are calcifying fibroblastic granuloma,

peripheral fibroma with calcification, ossifying fibroid epulis, and peripheral cementifying fibroma.¹³

The management of POF is by the elimination of local irritants and surgical excision of the lesion including the underlying periosteum. 14 The recurrence rate of POF is 8 - 20%, which can be due to the incomplete removal of the growth. 15

Abbreviations

- 1. POF Peripheral ossifying fibroma
- 2. PDL Periodontal ligament.
- 3. IOPA- Intraoral periapical radiograph
- 4. H & E Haematoxylin & eosin

CONCLUSIONS

A slowly progressing growth, anterior to the molar region in an elderly male patient with high degree of cellularity, inflammatory cells and dystrophic calcification confirmed the diagnosis of POF. Therefore, it is important for clinicians to think out of the box as regards the aetiology of POF.

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