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# Clinicopathological Study of Peripheral Ossifying Fibroma in a Kashmiri Population

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### **Abstract**

Peripheral cement ossifying fibromas are reactive rather than of neoplastic nature, the pathogenesis being uncertain. The lesion is seen mostly in adolescents and young adults, with the highest prevalence in 10-19 year age group and affecting females more than males(two to four times more), maxilla affected more than mandible. The present clinicopathological study investigated 150 cases of peripheral ossifying fibroma in a Kashmiri population.

**Keywords:** Peripheral cementifying fibroma, peripheral ossifying fibroma, pyogenic granuloma

#### Introduction

The peripheral ossifying fibroma (POF) makes about 3.1% of all oral tumors[1] and for 9.6% of all gingival lesions.[2] This lesion was first described In 1872 by Menzel.[3] The literature reports a great incidence in infants and in young adults, mainly in the age group of 10 to 19 years,[4] mainly affecting females (two to four times more),[5] suggesting some hormonal influences as well.[1] The pathogenesis of this tumor is uncertain; however, the pluripotent cells of the periodontal ligament have the apparent ability to transform or metaplastically change into osteoblasts, cementoblasts or fibroblasts, in

irritants such as calculus, bacterial response plaque, orthodontic appliances, ill-adapted crowns, and irregular restorations, and are therefore, capable of producing a unique inflammatory hyperplasia, the peripheral ossifying fibroma.[1,6-10] The cells of the periosteum or periodontal ligament also synthesize the mineralized product of this entity. The POF, representing a reactive benign lesion of the connective tissue, is not the soft tissue counterpart of the central ossifying fibroma, which represents an osteogenic neoplasm.[11] The central type of ossifying fibroma arises from the endosteum or the periodontal ligament (PDL) adjacent to the root apex and expands from the medullary cavity of the bone. On the other hand, the peripheral type shows a contiguous relationship with the PDL, occurring solely on the soft tissues overlying the alveolar process. Clinically POF presents as a solitary, slow-growing, and well-demarcated nodular mass that exhibits a smooth surface, usually with normal-colored mucosa. It has a sessile or pedunculated base and is generally of a hard consistency.[12] Such lesions are generally smaller than 1.5 cm in diameter,[13] although there have been reports of some large 4 cm lesions as well.[14] About 60% of such lesions occur in the maxilla and more than 50% of all cases affect the

region of the incisors and canines; more precisely n the interdental papilla.[4,13-15] In the vast majority of cases, there is no apparent underlying bone involvement visible on the roentgenogram. However, on rare occasions, there does appear to be superficial erosion of the bone. A possibility of tooth migration due to the presence of a cemento-ossifying peripheral fibroma has been reported.[16] The lesions must be surgically excised and microscopically examined for confirmation of diagnosis. The extraction of the adjacent teeth is seldom necessary. However, the lesions do occur with some frequency, and in fact, repeated recurrences are not uncommon. Clinical differential diagnosis for such gingival growths includes fibroma, peripheral giant cell granuloma, pyogenic peripheral odontogenic granuloma, fibroma, and peripheral ossifying fibroma.[17]

#### **Materials and Methods**

The study was conducted jointly in the departments of periodontics and conservative dentistry and endodontics, government Dental college srinagar. Detailed clinical and histological information of 150 surgically removed POFs formed the basis of the study. The criteria of patients for inclusion in the study were:

- a)Age group 10-19 years
- b)Presence of atleast one clinical lesion in the mouth
- c) kashmiri origin patients
- 150 lesions of POF diagnosed clinically and histologically were evaluated for their biological considerations.

# Clinical manifestation of the lesions

Majority of the POF lesions included in the study were slow-growing, painless, soft tissue masses. The lesions usually started as small papules. According to the patients, the lesions had been present for approximately two months before diagnosis. As reported by the patients, the lumps were interfering with bite making them uncomfortable. Occasionally, bleeding occurred on tooth

brushing. The lesions initially occurred on one side of the mouth and in a subsequent course of time involved both sides. Both the maxillary and mandibular gingivae were affected and the growths appeared as nodular masses, with erythematous and ulcerated surfaces. The growths were sessile and firm in consistency measuring anywhere between 5 mm and 3 cm in diameter. The lesions appeared reddish-pink in color and were slightly pedunculated, with what appeared to be a broad-based attachment. The lesions were not fluctuant, nor did they blanch with pressure, but had a rubbery consistency. The lesions were tender to firm pressure, but not to light palpation.

## **Radiographic examination**

Panoramic radiographs were obtained [Figure 3]. The radiographs did not reveal any abnormality and there was no finding pertaining to the multiple exophytic lesions.

## **Biochemical investigations**

The laboratory tests performed included, a complete blood hemogram and lipid and thyroid profiles. All the test results were within normal limits except for the total leukocyte count,.

which was slightly more than normal in some cases (12800/mm3 of blood).

# Histopathological examination

On histopathological examination, upon low power magnification (4x), the lesional tissues exhibited a keratinized stratified squamous epithelium (gingiva), overlying a fibrous connective tissue stroma exhibiting dense interlacing bundles of collagen and numerous ossifications [Figure 4]. High power magnification (40x) showed pink homogenous calcified tissue (ossification), with a presence of osteocytes entrapped in the lacunae [Figure 5]. The histology for all the lesions was the same. The picture is consistent with the clinical diagnosis of POF.

### **Results and Statistical Analysis**

The POF was found more frequently in females(70%), (Table 1).76% of the POF lesions were found in the Maxillary gingiva and 24% of the lesions were found in the mandibular gingiva, (Table 2). Majority of the lesions were seen in the canine/premolar region(70%).

Table 1

Total	POF	Females		Males	
lesions		No	%	No	%
150		105	70	45	30

Table 2

Total	POF	Maxillary gingival\		Mandibular gingival	
lesions		No	%	No	%
150		114	76	36	24

#### **Discussion**

Intraoral ossifying fibromas have been described in literature since the late 1940s. Many names have been given to similar lesions, such as, epulis, peripheral fibromas with calcification, peripheral ossifying fibromas, calcifying fibroblastic granuloma, peripheral cementifying fibroma, peripheral fibroma with cementogenesis, and peripheral cemento-ossifying fibroma.[3,6,18-21] The varied nomenclature used for fibroblastic gingival lesions indicates that there is much controversy surrounding the classification of these lesions.[6,22] The main etiological factors of POF are trauma and chronic irritation, particularly from the subgingival plaque calculus.[3,5,8,23] Eversole and Rovin[3] stated that the constant irritation present during exfoliation of the deciduous teeth and eruption of the permanent teeth may also result in an increased incidence of reactive lesions. which originate from the periodontal ligament. This hypothesis is based on the fact that POFs arise exclusively on the gingiva, the subsequent proximity of the gingiva to the periodontal ligament, and the inverse correlation between age distribution of patients presenting with POF

and the number of missing teeth with associated periodontal ligament.[1,23] Despite reports of POF in children with deciduous or mixed dentition, little data is available regarding the specific occurrence of this lesion, nvolvement of the deciduous dentition. The with literature also reflects the association of natal teeth with ossifying peripheral fibroma.[24] Defective odontogenesis and/or retarded eruption of teeth have also been found to be associated with similar histological features in the dental follicles.[25 Ossifying fibromas may be clinically and radiographically impossible to separate from cementifying fibromas.[26] Much of the research study has been carried out to distinguish nossifying fibromas from cementifying fibromas, by using immunohistochemical analysis for keratin sulfate and chondroitin-4 sulfate, in which the cementifying fibromas show significant immunoreactivity for keratin sulfate and the ossifying fibromas show intensive immunostaining for chondroitin-4 sulfate.[27] Such lesions may be present for a number of months to years before excision, depending on the degree of ulceration, discomfort, and interference with function.[4,18]The present study evaluated the occurance of POF in a Kashmiri population, further studies are needed to evaluate the condition in the current population using contemporary gene analysis techniques.



Figure 1: Multiple oral lesions of peripheral ossifying fibroma in both the maxilla as well as the mandible



Figure 2: Occlusal clinical view of the lesions showing their expansile nature



Figure 3: Orthopantogram of a patient showing mixed stage of dentition, without any abnormality pertaining to the multiple exophytic lesions



Figure 4: Histopathological examination of the lesional tissue at ×4 magnification.

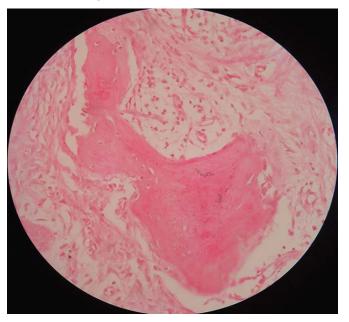


Figure 5: Histopathological examination of the lesional tissue at  $\times 40$  magnification



Figure 6: Post treatment photograph

## Conclusion

A peripheral cement-ossifying fibroma is a slowly progressing lesion, the growth of which is generally limited. Many cases will progress for long periods before patients seek treatment, because of the lack of symptoms associated with the lesion. Treatment consists of surgical excision, including the periosteum and scaling of the adjacent teeth. Close postoperative follow-up is required because of the growth potential of incompletely removed lesions and an 8 - 20% recurrence rate.

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