

A Retrospective Descriptive Study of Clinical Profile and Histopathological Diagnoses of Solitary Gingival Enlargements

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Abstract: Solitary gingival enlargements are similar in appearance and often do not show specific symptoms. Moreover in peripheral lesions we do not have radiographic findings to aid in the diagnosis. Histopathological diagnosis may vary from the clinical diagnosis to a great extent. This study was conducted to describe the clinical profile and histopathological diagnosis of solitary gingival enlargements that underwent excision biopsy over a period of one year. The study showed varied histopathological diagnosis among the solitary gingival enlargements studied, though the majority of cases were pyogenic granuloma.

Keywords: Solitary gingival enlargements, Pyogenic granuloma, Peripheral ossifying fibroma, Peripheral cementifying fibroma, Peripheral Ameloblastoma.

I. Introduction

Gingival enlargements are a common manifestation of several local and systemic diseases. They are the most often encountered lesions in the oral cavity¹. Due to the similarity in the clinical presentation and variety of overlapping terminologies, gingival enlargements remain a challenge for the clinician in proper diagnoses, classification and nomenclature². Most of the lesion occurs due to trauma or irritation¹. They differ in their etiopathogenesis, location, size and propensity for local destruction. Based on distribution they can be localized or generalized. The localized gingival enlargements are mostly inflammatory /reactive in nature example: irritational fibroma, pyogenic granuloma, peripheral ossifying fibroma and peripheral giant cell lesion/granuloma. Rarely some of these lesions are neoplastic that may be benign or malignant.³ Hyperplasia and neoplasia need to be clearly differentiated; long standing hyperplastic lesions in the presence of chronic irritation can become neoplastic⁴. An accurate diagnosis is critically important for the management and prevention of recurrence of these lesions^{3,5}. Histopathological diagnoses are found to vary from clinical diagnoses to a great extent in many cases⁵. The present study was conducted to describe the clinical profile of solitary gingival enlargements and note the histopathological diagnoses obtained for each.

II. Methodology

2.1 Study Design: Retrospective study

2.2 Study Population: All excisional biopsies submitted for histopathological examination from March 2015 to February 2016.

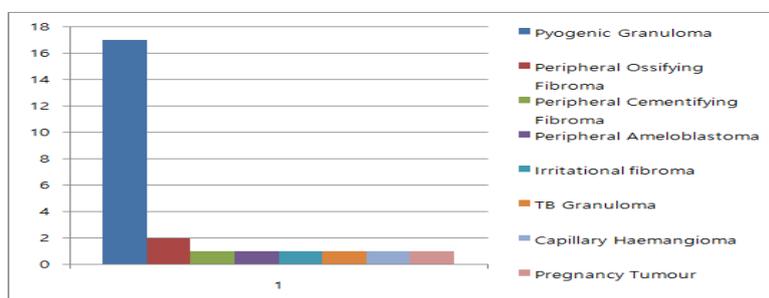
2.3 Inclusion Criteria: Solitary gingival enlargements that underwent excisional biopsy.

III. Results

Table 1: Distribution of Gingival Enlargements

S.No	AGE	GENDER	LOCATION	HISTOPATHOLOGICAL FEATURE
1	55	FEMALE	46	Capillary Haemangioma
2	53	FEMALE	11,21	Pyogenic granuloma
3	44	FEMALE	24,25	Peripheral Ossifying Fibroma
4	32	MALE	31,41	Pyogenic Granuloma
5	51	FEMALE	46,47	Pyogenic Granuloma
6	34	FEMALE	33,34	Pyogenic Granuloma
7	65	FEMALE	11,12,13	Peripheral Ameloblastoma
8	35	FEMALE	11,21	Peripheral Cementifying Fibroma
9	22	MALE	22,23	Pyogenic Granuloma
10	46	FEMALE	17,18,27,28	Irritational Fibroma
11	44	MALE	24,25,26	Pyogenic Granuloma
12	36	MALE	36	Pyogenic Granuloma
13	44	FEMALE	12,13	Pyogenic granuloma
14	25	FEMALE	35,36	Pregnancy Tumour
15	53	FEMALE	21,22	Pyogenic granuloma
16.	42	FEMALE	45,46	Pyogenic granuloma

17.	53	MALE	31,32,41,42	Tuberculous Granuloma
18.	56	FEMALE	27,28	Pyogenic granuloma
19.	33	MALE	33	Pyogenic granuloma
20.	60	FEMALE	13	Pyogenic granuloma
21.	42	FEMALE	11,12	Peripheral Ossifying Fibroma
22.	24	FEMALE	22	Pyogenic granuloma
23.	47	FEMALE	45,46	Pyogenic granuloma
24.	46	FEMALE	36	Pyogenic granuloma
25.	35	FEMALE	17	Pyogenic granuloma



Graph1 : Distribution of Gingival Enlargements

IV. Discussion

Soft tissue enlargements of the oral cavity often present a diagnostic challenge because of diverse etiopathogenesis. Among the soft tissue enlargements of oral cavity gingival enlargements are a common finding in clinical practice and pose a diagnostic dilemma to the clinician due to their similarity in clinical appearance. The most common form of enlargement is due to plaque induced inflammation of the adjacent gingival tissues and may be localized or generalized. Gingival enlargements are influenced by hormonal effects, as found in puberty and pregnancy, and also by certain systemic medications. The present study aims to describe the clinical profile of solitary gingival enlargements and to note the histopathological diagnoses.

In our study Pyogenic granuloma (Fig 1)(Table 1) was the most frequently encountered pathology comprising 68% of total cases of solitary lesions. Females were more affected than males with a wide age range having a peak incidence in the fourth and fifth decade of life. Similar observations were reported by Kfir et al⁶ and Angelopoulos⁷. Poor oral hygiene may be a precipitating factor in many PG patients⁸. In our study maxilla was more affected than mandible, anterior region more affected than posterior and facial regions were more affected than lingual aspects. Similar findings were recorded by Vilman et al⁹.

Pyogenic granuloma of the gingiva develops in up to 5% of pregnant females¹⁰. In the present study pregnancy tumour (Fig2) (Table 1) constitutes about 4% of total cases. The development of this particular kind of gingival lesion, typical in pregnancy, which is clinically similar to pyogenic granuloma in non pregnant women, suggests the existence of a relationship between the gingival lesions and the hormonal condition observed in pregnancy. Progesterone and estrogen hormones render the gingival tissue more susceptible to chronic irritation caused by plaque and calculus^{6,7}.

In this study, irritational fibroma (Fig3) (Table 1) constituted 4% of the cases. The inflammatory or reactive hyperplasia of gingiva may be the pyogenic granuloma at different stages of histological maturation^{6,11}. Irritational fibroma could represent a fibrous maturation of PG especially in those lesions with long duration. Peripheral ossifying fibroma (Fig4) (Table 1) which is also known as Ossifying fibrous epulis or peripheral fibroma with calcification occurred more frequently in females than males by a ratio of 2.25:1^{6,12}. In the present study Peripheral ossifying fibroma comprised of 8% of the total cases with female predilection. In the present study lesions were observed in the fourth decade of life, in contrast to the findings of Eversole and Rovin¹³. They suggested that the loss of periodontium that accompany tooth loss in old age may explain the greater occurrence of Peripheral ossifying fibroma in the younger age group. Superficial periodontal ligament which contains cells capable of producing bone are considered to be the etiopathogenesis of the lesion.

Peripheral Cementifying Fibroma (Fig5) (Table 1) comprises of 1%-3% of the gingival lesions^{14,15,16,17}, with a peak incidence in 2nd and 3rd decade of life having a female predilection¹⁸. PCF is encountered more often in maxillary anterior region¹⁹. Our study showed similar findings.

Peripheral Ameloblastoma (Fig6) (Table 1) is a rare odontogenic tumor that accounts for 1% of all ameloblastomas²⁰. It is typically a slow, benign, single, sessile, asymptomatic lesion. Histologically, this lesion is identical to the classic intraosseous ameloblastoma but appears exclusively in the oral mucosa over the alveolar processes of both jaws. Radiographic studies usually are negative for any bony destruction or invasion except for sclerization of the underlying periosteum²¹. The etiology of PA is unclear. The tumor may be derived from the extraosseous epithelial remnants of the dental lamina or from the basal cell layer of the oral mucosa, which is believed

eved to have odontogenic potential²². In our study we observed a case with similar clinical and radiographic features in the maxillary anterior region of a female patient which was histopathologically diagnosed as peripheral ameloblastoma.

Tuberculosis is a chronic granulomatous infectious disease caused by *Mycobacterium tuberculosis*; it can affect any part of the body including the oral cavity. Extrapulmonary tuberculosis is rare, occurring in 10% to 15% of all cases²³. Diagnosis of oral tuberculosis is difficult as the clinical presentation may take various forms and the typical constitutional features are absent in most cases. The usual manifestation is as an ulcer or localized granular mass^{24,25}. In the present study we encountered a case of granular mass involving the mandibular anterior gingiva in male patient with tuberculosis (Fig7) (Table 1).

Capillary hemangiomas are considered one of the common soft tissue tumors of the head and neck; it is relatively rare in the oral cavity. Capillary hemangiomas are composed of many small capillaries lined with a single layer of endothelial cells supported in connective tissue stroma of varying density^{26,27}. In the present study we encountered a lesion with similar histopathological features involving the mandibular posterior region (Fig8) (Table 1).

V. Conclusion

Our study though limited for a short period of one year and consisting of only excised lesions showed a plethora of histopathologic diagnoses in clinically similar lesions. This highlights the importance of biopsying gingival enlargements especially in patients with long standing lesions and those with systemic diseases. Pyogenic granuloma is the most common lesion in our study, as found by other investigators. Hence it would be prudent to remove the most probable etiologic factor of this lesion, namely local irritants, as the first line of treatment of solitary gingival enlargements. Lesions which do not regress however need to be biopsied without fail.

Reference

- [1]. Sumona Pal, Shruthi Hegde, Vidya Ajila. The varying Clinical Presentations of Peripheral ossifying Fibroma: A report of three cases Rev OdontoCience 2012;27(3):251-255
- [2]. Kiran Kumar Ganji, Arun Kumar Bhimashankar Chakki, Sharanbasappa Chandrashelar et al. peripheral Cemento-ossifying Fibroma: Case series Literature review. Case Reports in Dentistry Vol 2013 Article ID 930870, 5 pages.
- [3]. Amit Arvind Agarwal. Gingival Enlargements: Differential Diagnosis and review of literature. World J Clin Cases 2015 September 16;3(9):779-788
- [4]. Mani AM, Marawar PP, Pendyala G, Mhaisekar R, Bhadange S. series of Gingival Enlargements: Case Reports. Parvata Med Rev 2014;6(1):23-28
- [5]. Devanorkar A, Guttiganur N, Dawarakanath C D, Savitha A N. Biopsy the gold standard in the final diagnosis of solitary gingival enlargements: Case reports and an overview. Int J Oral Health Sci 2013;3:65-9
- [6]. Kfir Y, Buchner A, Hansen LS : Reactive lesions of the gingiva. A clinico-pathological study of 741 cases. J Periodontol. 1980; 51: 655-61.
- [7]. Angelopoulos AP: Pyogenic granuloma of the oral cavity: Statistical analysis of its clinical features. J Oral Surg. 1971; 29: 840-47.
- [8]. Regezi JA, Sciubba JJ, Jordan RCK: Oral Pathology: Clinical Pathologic consideration. 4th ed; WB Saunders, Philadelphia ; 2003. p . 115-116.
- [9]. Vilmann A, Vilmann P, Vilmann H: Pyogenic granuloma: evaluation of oral conditions. Br J Oral Maxillofac Surg. 1986; 24:376-82.
- [10]. Sills ES, Zegarelli DJ, Hoschander MM, Strider WE: Clinical diagnosis and management of hormonally responsive oral pregnancy tumor (pyogenic granuloma). J Reprod Med . 1996; 41:467-70.
- [11]. Daley TD, Wysocki GP, Wysocki PD, Wysocki DM: The major epulides: Clinico pathological correlations. J Can Dent Assoc. 1990; 56: 627-30.
- [12]. Buchner A, Calderon S, Ramon Y: Localized hyperplastic lesions of the gingiva: a clinicopathological study of 302 lesions. J Periodontol. 1977; 93:305-9.
- [13]. Eversole LR, Rovin S: Reactive lesions of the gingiva. J Oral Pathol 1972;1:30-38.
- [14]. Cuisia ZE, Brannon RB. (2001). Peripheral ossifying fibroma- a clinical evaluation of 134 pediatric cases. Pediatr Dent, 23, 245-248.
- [15]. 15. Bhaskar SN, Jacoway JR. (1966). Peripheral fibroma and peripheral fibroma with calcification: Report of 376 cases. J Am Dent Assoc, 73, 1312-20.
- [16]. 16. Kenney JN, Kaugars GE, Abbey LM. (1989). Comparison between the peripheral ossifying fibroma and peripheral odontogenic fibroma. J Oral Maxillofac Surg, 47(4), 378-82
- [17]. Das S, Das AK. (1993). A review of pediatric oral biopsies from a surgical pathology service in a dental school. Pediatr Dent, 15(3), 208-11.
- [18]. Iqbal Ali, Gazala Parveen, Sameer Singh, Manish Dubey, Akheel M.D. (2014). Peripheral Ossifying Fibroma of Maxilla: A case report & Review. Journal of Head & Neck physicians and surgeons, 2(2), 128-35.
- [19]. Terry Farquhar, Jennifer Mac Lellan, Heather Dymont, Ross D. Anderson. (2008). Peripheral Ossifying Fibroma: A Case Report. Journal of Canadian dental association, 74, 809-12.
- [20]. Neville BW, Damm DD, Allen CM, Bouquot JE. Oral & maxillofacial pathology. Pennsylvania: W.B. Saunders Company; 1995, p 519.
- [21]. Nauta JM, Paunders AK, Schoots CJF, Varmey A, Roodenburg JLN. Peripheral ameloblastoma. A case report and review of literature. Int J Oral Maxillofac Surg 1992;21: 40-4.
- [22]. Buchner A, Sciubba JJ. Peripheral epithelial odontogenic tumors: a review. Oral Surg 1987;63:688.
- [23]. Memon GA, Khushk IA. Primary tuberculosis of tongue. J Coll Physicians Surg Pak 2003; 13(10):604-5.
- [24]. Aneksuk V. [Primary tuberculous ulcer of the gingiva: report of 1 case]. J Dent Assoc Thai 1988; 38(3):111-5. Thai.

- [25]. Kobayashi T, Sato M, Onoi Y, Ohshiro S, Nishii K, Ishihara T, and others. [A case of pulmonary tuberculosis complicated with gingival lesions.] *Kekkaku* 1997; 72(6):411-4. Japanese.
- [26]. Bayrak S, Dalci K, Tansel H. Capillary hemangioma of the palatal mucosa: Report of an unusual case. *SÜ Dişhek Fak Derg* 2010;19:87- 9.
- [27]. Chen YK, Lin LM, Huang HC, Lin CC, Yan YH. A retrospective study of oral and maxillofacial biopsy lesions in a pediatric population from southern Taiwan. *Pediatr Dent* 1998;20:404- 10.

FIGURES



Fig 1 Pyogenic Granuloma



Fig 2 Pregnancy Tumour



Fig 3 Irritational Fibroma



Fig 4 Peripheral Ossifying Fibroma



Fig 5 Peripheral Cementifying fibroma



Fig 6 Peripheral Ameloblastoma



Fig 7 Tuberculous Granuloma



Fig 8 Capillary Haemangioma