

Orofacial Pathologic Lesions in Children and Adolescents: A Clinicopathological Study in Southern Iran

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Received: Mar 25, 2013; Accepted: Dec 23, 2013; First Online Available: Apr 18, 2014

Abstract

Objective: Oral and maxillofacial lesions vary regarding their clinical presentation in different populations. Until now, oral and maxillofacial lesions in Iranian children and adolescents have not been studied. The aim of this study was to determine the type and distribution of biopsied oral lesions among children and adolescents in Southern Iran.

Methods: All the patients referred to the pathology department of Shiraz Faculty of Dentistry from 1991-2009 were enrolled in this retrospective, case-series study. The information regarding the patients' age, gender as well as the histopathologic type and anatomic location of the biopsied oral lesions in patients under 18 years was collected from patients' medical documents and were analyzed by SPSS version 11.

Findings: Out of 2984 patients, 576 (19.3 %) cases were children and adolescents under 18 years. The most prevalent category was soft tissue lesions (45.5 %). The most common lesion was peripheral giant cell granuloma (15.6%) followed by dentigerous cyst (14.2%) and pyogenic granuloma (11.3%). Gingiva was the most common affected site. Male to female ratio was 1.2.

Conclusion: Our results revealed that near 20% of orofacial lesions occur in children and adolescents with rather equal male to female ratio. The majority of lesions were soft tissue diseases with a reactive nature. Unlike other studies we had higher rates of soft tissue lesions. These data can help dentists and surgeons for more accurate management of their patients.

Iranian Journal of Pediatrics, Volume 24 (Number 3), Jun 2014, Pages: 307-312

Key Words: Pathologic Lesion; Oral Pathology; Maxillofacial Malignancy

Introduction

Oral and maxillofacial lesions are heterogeneous group of diseases with wide spectrum of clinicopathologic characteristics. Initial diagnosis of these disorders is essentially based on their clinical presentation. Relative incidence of any pathologic lesion plays a critical role in differential and definitive diagnosis. There are many studies published in recent years about overall prevalence

of oral and maxillofacial lesions in adults. Also, many epidemiologic researches that have been done in children and adolescents, were focused on specific lesions such as maxillofacial malignancies^[1], salivary gland tumors^[2,3] and odontogenic lesions^[4]. However, there are not enough studies concerning the pathologic oral lesions in children and adolescents. Since hereditary and geographical factors influence the incidence of oral lesions; further studies are

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important to evaluate the clinical and epidemiologic characteristics in different populations^[5]. Regarding this fact, to the best of our knowledge, a few studies performed on Asian children are available^[3,6,7].

The aim of our study was to define the frequency and type of maxillofacial lesions in a group of Iranian children and adolescents.

Subjects and Methods

In this retrospective study all pathologic reports at oral and maxillofacial pathology department of Shiraz Dental School in southern Iran from 1989 to 2009 were recruited.

The demographic data were obtained from patients' medical records and documents. Anatomic location of lesions and their histopathologic type were recorded as well as the patients' age and gender. Patients were classified into two age groups: under 12 years old (children) and 13-18 years old (adolescents). Lesions were categorized in 8 groups based on the tissue of origin where the lesions aroused (Neville et al, with a few modifications)^[8]: soft tissue lesions, odontogenic cysts, odontogenic tumors, salivary gland lesions, bone lesions, epithelial lesions, hematologic diseases and pulp and periapical diseases. Data were analyzed using SPSS software (Version 11, Chicago).

Findings

Out of 2984 patients, 576 (19.3%) cases were under 18 years of age. In our study, male to female ratio was 1:2 and the most common involved site was gingiva. Tables 1-4 show the distribution of lesions according to the location of the lesion and patients' gender. The most common histopathologic type was soft tissue lesions (45.5%), followed by odontogenic cysts (20%), and bone lesions (10.2%) (Table1). Overall, peripheral giant cell granuloma (PGCG) with 920 cases, dentigerous cyst with 820 cases, and pyogenic granuloma with 650 cases were the most common lesions.

In the pediatric age group, the most common lesion was PGCG. In the adolescents, the most common lesion was pyogenic granuloma and peripheral ossifying fibroma.

Discussion

This study evaluated the oral pathologic lesions in a group of children and adolescents and found that these patients comprise 19.2 % of total cases.

This rate was higher than the rate reported by the previous researchers in USA (7%)^[9], U.K (8.2%)^[10], China (5.5%)^[3] and Thailand (15%)^[7] but Lawoyin et al^[12] have reported a prevalence of 25% for the pathologic lesions in Nigerian children and adolescents under 17 years. It should

Table 1: Soft tissue lesions

	Number		% of total	Male/Female ratio	Most common location
	adolescents	children			
PGCG	76	16	35/1	2	Gingiva
Pyogenic Granuloma	33	34	24/8	1	Gingiva
Peripheral Ossifying Fibroma	9	32	16/4	1/1	Gingiva
Irritation Fibroma	7	15	8/4	0/6	Gingiva
Fibroepithelial Polyp	8	9	6/5	0/5	Gingiva
Hemangioma	4	6	3/8	4	Buccal mucosa
Fibrous Histiocytoma	2	3	2	1/5	Gingiva
Giant cell Fibroma	2	1	1/2	0:3	Gingiva
Neurofibroma	3	-	1/2	0:3	Buccal mucosa
Desmoplastic Fibroma	2	-	0/8	1	Gingiva
Total	146	116	-	-	

Table 2: Odontogenic cysts and tumors

Lesions	Number		% of total	Male/Female ratio	Most common location
	adolescents	children			
Dentigerus Cyst	53	29	71/4	2/03	Mand
KOT	6	20	22/6	1/16	Max = Mand
Eruption Cyst	6	0	5/2	6:0	Mand
Calcifying Odontogenic Cyst	0	1	0/8	0:1	Max
Odontoma	4	5	27/3	2	Max
Ameloblastoma	1	8	27/3	1/25	Mand
AOT	-	5	15/2	0/25	Mand
Odontogenic Fibroma	1	3	12/2	1	Max
Cementoblastoma	-	3	9	0/5	Mand
Ameloblastic Fibroma	2	-	6	0:2	Max = Mand
Myxoma	1	-	3	1:0	Mand
Total	74	74			

KOT: Keratocystic odontogenic tumor; AOT: Adenomatoid odontogenic cyst

be noted that the age range of the patients have been different in these studies (under 15 to under 18 years old)^[6,7,9,10]. Evaluation of clinical characteristics of pathologic lesions can help clinicians for a better clinical management of the any pathologic lesion, demographic data of the patient should be considered. These data are influenced by geographic and ethnic factors, so epidemiologic studies in any population seem to be important.

The frequency of oral lesions was almost equal in male and female groups. This finding was in agreement with other studies that have shown M:F ratio to be 1-1.2^[6,7,10,11]. In the current study, the oral lesions in children and adolescents showed the same incidence. The studies that have evaluated the oral lesions in a long-term have stated that the prevalence of lesions increased with age^[6,9,10,11]. In a study that has been done in

Thailand, the patients were classified into three groups regarding their dental age: permanent, mixed and deciduous dentition. The authors stated that the prevalence of oral lesions was higher in mixed dentition group^[7]. Also, the incidence of oral lesions in children and adolescents was respectively 52% and 48% in Jones et al study^[10] and 48.2% and 51.8% in Chen et al study^[6]. It seems that the overall prevalence of oral pathologic disorders was almost equal between children and adolescent groups. Moreover, it should be considered that the prevalence of any specific lesion may be different in the mentioned groups.

Regarding the histopathologic type of the subjects, the present study, in line with the previous studies, showed that reactive soft tissue lesions as well as cystic lesions occurred more frequently than other lesions^[6,7,9,10], but one study

Table 3: Salivary gland disease and epithelial pathology and hematologic disorders

Lesions	Number		% of total	M/F ratio	Most common location
	adolescents	children			
Muscocele	20	25	93/7	1/5	Lower lip & Floor of mouth
Mucoepidermoid Carcinoma	-	2	4	1	Parotid
Adenoid Cystic Carcinoma	1	-	2	1:0	Palate
Necrotizing Sialometaplasia	1	-	2	1:0	Palate
Pleomorphic Adenoma	1	-	2	0:1	Parotid
Epithelial Hyperplasia	3	4	53/5	1/3	Gingiva
Squamous Papilloma	2	3	39	0:5	Lip and tongue
Nevus	-	1	7/5	0:1	Buccal mucosa
Lymphoma	4	1	71/5	5:0	Vestibule
Eosinophilic granuloma	-	2	28/5	1	Mand = Max
Total	32	38	-	-	-

Table 4: Pulp and periapical diseases and bone pathology

Lesions	Number		% of total	M/F ratio	Most common location
	adolescents	children			
Radicular cyst	15	18	91/7	1/3	Max
Residual cyst	1	2	8/3	2	Mand
CGCG	16	11	45	0/7	Mand
Fibrous Dysplasia	2	7	15	1	Max
Central Ossifying Fibroma	3	5	13/3	1	Max
Aneurysmal Bone Cyst	3	4	11/7	0:7	Mand
Traumatic Bone Cyst	0	6	10	1	Mand
Osteosarcoma	1	1	3/4	1	Max
Chondrosarcoma	0	1	1/6	0:1	Max
Total	41	55			

CGCG: central giant cell granuloma

done on African children and adolescents has shown that neoplasms were more common than inflammatory lesions^[12]. This finding may have been the result of considering reactive lesions (such as ossifying fibroma) as a neoplasm in that study and also, due to the endemic occurrence of Burkitt's lymphoma in African children.

PGCG was the most common lesion in our study; however, other authors have reported mucocele, dentigerous cyst, radicular cyst, and pyogenic granuloma as the most common oral lesion, respectively in southern Taiwan and Chile^[6,13,14], Greece^[15,16], Thailand^[7] and Brazil^[11,17]. Trauma and poor oral hygiene are the major etiologic factors for the mentioned lesions, except dentigerous cyst. In contrast to our results in children, pyogenic granuloma in adults shows a definitive female predilection, probably because feminine hormones affect the vascular events^[8,18].

Odontogenic cysts were the second most common group which constituted 20% of all lesions. This incidence was lower than that of the studies performed in Africa^[12], Thailand^[7] and Taiwan^[6] (22-35%), but higher than in Turkey (12%)^[19]. In the most studies, dentigerous cyst was the most common cyst^[6,7,9,12,20]. In our study, dentigerous cyst and then odontogenic keratocyst (OKC) were reported more frequently. This finding has been confirmed by other researchers^[4,7,13]. Odontogenic cysts arise from odontogenic epithelium of tooth germ, and formation of these lesions during development of teeth in the first two decades of life occur frequently^[8].

Bone lesions were the third most common group of lesions which constituted 10.3% of all. Other studies have been reported this incidence

from 3.2 to 4.8%^[10,20]. The present study and all the other studies showed that central giant cell granuloma and fibrous dysplasia were the most common lesions in this group. Respectively, mandible and maxilla were the most frequent sites of involvement, as reported by Lawoyin et al^[12]. However, Maya et al reported that both lesions were more frequent in mandible^[17].

In salivary gland pathology, mucocele was the most frequently reported lesion detected in lower lip. The lesion was found predominantly in females. Other authors have stated that mucocele is the most common oral lesion in children^[6,10,11,13,15,17,20], but its incidence was lower in our study. Also, Shulma^[21] in a large group of patients under 17 years found only 5 cases of mucocele. This lesion almost equally affected the males and females, but Nico et al have found a female predilection^[22].

In the group of pulp and periapical lesions, similar to Maya et al, radicular cyst was the most common lesion which involved maxillary bone more frequently than mandible^[17]. Many authors have included this cyst in odontogenic cysts group and have reported this lesion as the second common cyst after dentigerous cyst^[23].

Within odontogenic tumors, odontoma and ameloblastoma were the most frequently found hamartoma and tumor in maxilla and mandible respectively. It is in agreement with other studies that have reported these lesions as first and second common odontogenic tumor^[7,10,11,17,20]. Maya et al have found odontoma in mandible more frequently^[17]. The male to female ratio for odontoma and ameloblastoma was 2:1 and 1.25:1 respectively, which is comparable with other studies^[6,12,24]. However, there are some reports

that have demonstrated female or male predilection or an equal gender distribution for both lesions.

Although in the new classification of World Health Organization (WHO), OKC have been defined as an odontogenic tumor^[8] and authors who have considered this classification, have reported keratocystic odontogenic tumor (KOT) as the most common odontogenic tumor^[4,6,10,11,13,17,20]. Regarding the anatomic location, we found predominance of maxillary bone. Authors in Africa^[12], Thailand^[7] and Nigeria^[24] have reported results which are in line with our findings, but Latrou et al^[16], studied odontomas in Greece population and have noted that mandible and maxilla have been involved equally.

We found only 13 cases of epithelial lesions. Gingiva was the most frequently affected site. Epithelial hyperplasia and squamous papilloma were the most common lesions in this category. Unfortunately, there are not many available researches about these lesions. Wang et al^[13] have demonstrated only a few cases of epithelial hyperplasia. These lesions showed female predilection in our study which is in agreement with a study in UK^[10] but Wang et al have found male predominance in Taiwan^[13]. However, due to the limited number of recorded cases, this difference is acceptable.

The incidence of hematopoietic disorders was 1.2%, this included 5 cases of Burkitt's lymphoma and 2 cases of eosinophilic granuloma. In other studies this incidence was lower than 5%^[6,10,17,20]. But in Lowoyin et al's study the prevalence was 11%^[12]. This high incidence is due to the endemic occurrence of Burkitt's lymphoma in African children. Out of five cases of Burkitt's lymphoma, 4 cases occurred in males. This male predilection is similar to other studies^[6,12,20].

In this study we found only 0.86% malignant neoplasms including 5 cases of Burkitt's lymphoma, 2 cases of mucoepidermoid carcinoma and one case of adenoid cystic carcinoma, with a M:F ratio of 0.7:1. The incidence of malignant lesions was similar to the previous studies in Taiwan^[6], UK^[10] and Thailand^[7]. However, the studies on African populations have demonstrated a high rate of malignancy; most of them were lymphomas^[12,13]. These results might be explained by some deficiencies and genetic susceptibilities in African children^[12,13].

Conclusion

Our results revealed that nearly 19.2% of lesions occurred in patients less than 18 years with an equal gender distribution. The majority of lesions were soft tissue diseases with a reactive nature. However, there were some differences in comparison with other studies; for instance, our rate was higher in soft tissue lesions and a lower prevalence of mucocele. These data could help the dentists and surgeons for more accurate management of their patients.

Similar studies in various groups of patients should be designed to obtain an actual prevalence and accurate demographic data for any oral disease.

Acknowledgment

This article is based on the thesis by Dr. Zahra Ahmadi-Sheshdeh. The authors thank the vice-chancellor of Shiraz University of Medical Sciences, for supporting the research. Also the authors would like to thank Dr Sh. Hamedani (DDS, MSc) for critical review of the manuscript and helping with English and editorial assistance.

Authors' Contribution

Z. Jaafari: conceived and designed the study and prepared the manuscript.
Z. Ahmadi: performed the literature search, collected data, initially analyzed and interpreted the data.
F. Kamali: contributed in writing of initial manuscript.
All authors read and approved the final version of the manuscript.

Conflict of Interest: None

References

1. Kamulegeya A, Lakor F. Oral maxillofacial tumors and tumor-like conditions: a Ugandan survey. *Pediatr surg int* 2011; 27(9):925-30.
2. Sultan I, Rodriguez-Galindo C, Al-Sharabati S, et al. Salivary gland carcinomas in children and adolescents: a population-based study, with comparison to adult cases. *Head Neck* 2011; 33(10): 1476-81.
3. Liu B, Liu JY, Zhang WF, et al. Pediatric parotid tumors: clinical review of 24 cases in a Chinese population. *Int J Pediatr Otorhinolaryngol* 2012; 76(7):1007-11.

4. Servato JP, Prieto-Oliveira P, de Faria PR, et al. Odontogenic tumours: 240 cases diagnosed over 31 years at a Brazilian university and a review of international literature. *Int J Oral Maxillofac Surg* 2013; 42(2):288-93.
5. Jaafari-Ashkavandi Z, Ashraf MJ. A clinico-pathologic study of 142 orofacial tumors in children and adolescents in southern Iran. *Iran J Pediatr* 2011; 21(3):367-72.
6. Chen YK, Lin LM, Huang HC, et al. A retrospective study of oral and maxillofacial biopsy lesions in a pediatric population from southern Taiwan. *Pediatr Dent* 1998; 20(7):404-10.
7. Dhanuthai K, Banrai M, Limpanaputtajak S. A retrospective study of paediatric oral lesions from Thailand. *Int J Paediatr Dent* 2007; 17(4):248-53.
8. Neville BW, Damm DD, Allen CM, et al. Oral and maxillofacial pathology. 3rd ed. Philadelphia: Saunders. 2009; Pp:362-741.
9. Shah SK, Le MC, Carpenter WM. Retrospective review of pediatric oral lesions from a dental school biopsy service. *Pediatr Dent* 2009; 31(1):14-9.
10. Jones AV, Franklin CD. An analysis of oral and maxillofacial pathology found in children over a 30-year period. *Int J Paediatr Dent* 2006; 16(1):19-30. .
11. Sousa FB, Etges A, Correa L, et al. Pediatric oral lesions: a 15-year review from Sao Paulo, Brazil. *J Clin Pediatr Dent* 2002 Summer;26(4):413-8.
12. Lawoyin JO. Paediatric oral surgical pathology service in an African population group: a 10 year review. *Odonto-Stomatol Trop* 2000;23(89):27-30.
13. Wang YL, Chang HH, Chang JY, et al. Retrospective survey of biopsied oral lesions in pediatric patients. *J Formos Med Assoc* 2009; 108(11):862-71.
14. Zuniga MD, Mendez CR, Kauterich RR, et al. Paediatric oral pathology in a Chilean population: a 15-year review. *Int j paediatr dent* 2013; 23(5):346-51.
15. Sklavounou-Andrikopoulou A, Piperi E, Papanikolaou V, et al. Oral soft tissue lesions in Greek children and adolescents: a retrospective analysis over a 32-year period. *J Clin Pediatr Dent* 2005; 29(2):175-8.
16. Iatrou I, Vardas E, Theologie-Lygidakis N, et al. A retrospective analysis of the characteristics, treatment and follow-up of 26 odontomas in Greek children. *J Oral Sci* 2010; 52(3):439-47.
17. Maia DM, Merly F, Castro WH, et al. A survey of oral biopsies in Brazilian pediatric patients. *ASDC J Dentist Child* 2000; 67(2):128-31, 83.
18. Regezi JA, Sciubba J, Jordan R. Red-blue lesions. In: Oral pathology clinical-pathology correlations. 5th ed. Philadelphia: Saunders; 2008: Pp:111-112.
19. Gultelkin SE, Tokman B, Turkseven MR. A review of paediatric oral biopsies in Turkey. *Int Dent J* 2003; 53(1):26-32.
20. Lima Gda S, Fontes ST, de Araujo LM, et al. A survey of oral and maxillofacial biopsies in children: a single-center retrospective study of 20 years in Pelotas-Brazil. *J Appl Oral Sci* 2008; 16(6):397-402.
21. Shulman JD. Prevalence of oral mucosal lesions in children and youths in the USA. *Int J Paediatr Dent* 2005; 15(2):89-97.
22. Nico MM, Park JH, Lourenco SV. Mucocele in pediatric patients: analysis of 36 children. *Pediatr Dermatol* 2008; 25(3):308-11.
23. Bodner L. Cystic lesions of the jaws in children. *Int J Pediatr Otorhinolaryngol* 2002; 62(1):25-9.
24. Adebayo ET, Ajike SO, Adekeye EO. Odontogenic tumours in children and adolescents: a study of 78 Nigerian cases. *J Craniomaxillofac Surg* 2002; 30(5): 267-72.