

Retrospective study of the clinical and histopathological parameters of the dental lesions

Shilpi Daveshwar¹, Hiral Samir Shah^{2*}, Meena Rajiv Daveshwar³

¹PG student, Manubhai Patel Dental College and Dental Hospital, Baroda.

²Tutor, ³Associate Professor, S.S.G. Hospital and Medical College, Baroda.

Email: hiralsamir07@yahoo.in

Abstract

Background: Oral cavity is the site of myriads of lesions that may be categorized as inflammatory/reactive, cystic and neoplastic. The possibility of identifying the frequency of each lesion is fundamental in order to understand disease patterns within populations and support differential diagnosis. The aim of our study was to investigate the frequency and spectrum of oral lesions biopsied in a hospital population in order to provide insight into the prevalence of them.

Methods: A retrospective study of 94 specimens received in Pathology Department from Dental OPD from January 2015 to December 2019 was conducted. The histopathology slides and reports were retrieved from Pathology Department. The histopathological diagnosis was divided into categories considering several variables (type of lesion, gender and age of the affected patient). A statistical analysis was conducted in order to evaluate the prevalence for each cluster of pathology and its relationship with the other identified variables. **Results:** In group 1, there were 29 samples which showed mainly nonspecific inflammation. Group 2, which consisted of total 23 samples of cystic lesions among which radicular cyst was the most frequent diagnosis. Maximum number of lesions were in group 3 which consisted of 42 neoplastic diseases (44.6%). In third group, malignant tumors represented 25.5% (24/94 cases) of the samples followed by benign tumours and premalignant lesions. Males represented 58.5% of the sample, while females were 41.5%. The overall male to female ratio was 1.4:1. The patients' age range was from 2 to 79 years and mean age was 34.9 year. **Conclusions:** At present time, the most reliable diagnostic tool is represented by histopathological analysis, which is essential in terms of morphological characterization of the lesion and still remains the gold standard for obtaining a definitive diagnosis. Many of our findings are consistent with those reported in literature, suggesting that the study of demographic characteristics and their association with occurrence of lesions should be considered in performing differential diagnosis.

Key Words: Biopsy, Dental Lesions, Histopathological Examination.

*Address for Correspondence:

Dr. Hiral Samir Shah, Address: Above Samir Hospital, Lakdipul, Main Road Dandiabazar, Baroda.390001. Gujarat. INDIA.

Email: hiralsamir07@yahoo.in

Received Date: 17/10/2019 Revised Date: 09/11/2019 Accepted Date: 23/12/2019

DOI: <https://doi.org/10.26611/10191232>

Access this article online

Quick Response Code:



Website:

www.medpulse.in

Accessed Date:
26 October 2019

INTRODUCTION

The word biopsy originates from the Greek terms bios (life) and oopsis (vision): vision of life. Biopsy is defined

as "The removal of tissue from a living person for microscopic examination to confirm or to establish the diagnosis."¹ The term was coined by Ernst Henry in 1879. The possibility of identifying the frequency of each lesion is fundamental in order to understand disease patterns within populations and support differential diagnosis. According to the American Academy of oral and Maxillofacial Pathology, any abnormal tissue removed from the oral region should be submitted to pathologist for final histopathological diagnosis. The exceptions are in cases such as tori, exostosis, carious teeth lacking attached soft tissue, extirpated dental pulp and clinically normal tissues.² It is important for the clinician to decide whether a lesion needs to be biopsied or not before treating it. With regard to oral soft tissues, any lesion in

How to cite this article: Shilpi Daveshwar, Hiral Samir Shah, Meena Rajiv Daveshwar. Retrospective study of the clinical and histopathological parameters of the dental lesions. *MedPulse – International Journal of Dentistry*. December 2019; 12(3): 19-24. <http://www.medpulse.in/Dentistry>

question, if persisting for more than 2 weeks even after the removal of the irritating factor (if any), biopsy should be performed. Biopsy is also advisable in bony lesions that cannot be diagnosed radiographically and which are usually accompanied by pain, sensation alterations or other symptoms³. Biopsy has been used since more than 150 years to establish the diagnosis of an unknown medical condition.⁴ It is widely used in the medical field, but the practice is not quite widespread in dental practice. Biopsy is one of the oldest, gold standard and most reliable currently available method that can establish the definitive diagnosis of clinical abnormalities. The diagnosis of dental lesions is established from the different clinical and radiological features though the final diagnosis is based on histopathological examination of the lesion.^{5,6} Thus Oral biopsy is considered essential for i) To establish a definitive diagnosis as early as possible so that correct treatment may be initiated without delay, ii) to establish a prognosis in malignant or premalignant lesions iii) To determine whether an abnormality has been completely removed, iv) act as a document with medico legal value. Failure to diagnose oral disease may have profound implications for both the patient and the dentist. Although absolute contraindications are not present, there are some conditions where decision to proceed with biopsy should be done with caution. These are bleeding diathesis secondary to anticoagulation, lesion located near vital structures that could be injured by biopsy and in medical conditions that do not allow for the use of local anesthetics. Biopsy is not advised in the case of multiple neurofibromas due to the risk of neurosarcomatous transformation, or in tumors of the major salivary glands. Such biopsies must be performed by specialized surgeons in order to avoid damaging the nearby anatomical structures and causing the spread of tumor cells, as this would adversely affect the prognosis.³ Aim of present study is to classify and determine the frequency of distribution of various inflammatory, cystic, preneoplastic and neoplastic dental lesions received for histopathological examination from dental department at tertiary care center. Final histopathological diagnosis was given in each case in correlation with the clinical findings.

METHODS

This retrospective study was performed at histopathology section of Pathology department, SSG Hospital and Medical College, Baroda. The biopsies of oral lesions were received from dental department of SSG Hospital. The patient records of histopathology sections from January 2015 to December 2019 were reviewed

considering gender, age and histopathological diagnosis. Lesions were divided into three major categories based on their histopathologic diagnosis: Group 1: Inflammatory/reactive oral lesions, Group 2: Cystic lesions and Group 3: Benign, premalignant and malignant lesions.^{7,8} The results obtained were tabulated and analyzed.

Statistical analysis

All results were analyzed using aspects of descriptive statistics. The data was analyzed in different age groups and gender by using frequency, ratio and percentage methods. For evaluation of association between mean age group and number of cases, χ^2 test was applied. Significance was considered for values of $p < 0.05$.

RESULTS

Total 94 cases were studied and classified in three groups. Distribution of cases among these three groups were analyzed. In Group 1 there were 29 cases (30.9%), group 2 had 23 cases (24.5%) and group 3 consisted of 42 cases (44.6%). In view of gender distribution, 55 cases (61.2%) were males and 39 (38.8%) cases were females. The overall male to female ratio was 1.4:1. The patients' age range was from 2 to 79 years and mean age was 34.9 year (Table 1). Group 1 included inflammatory or reactive conditions, had mean age of 35 year with M:F ratio 1.4:1. Histopathological examination showed 17 cases (58.6%) of chronic inflammatory cell infiltration along with other specific inflammatory lesions like mucormycosis, osteomyelitis, pericoronitis, pyogenic granuloma, inflamed epulis, fibroepithelial polyp and submucosal fibrosis. Group 2 included cystic lesions, with mean age of 24.5 year and M:F ratio was 1.5:1. In this group, the most prevalent was radicular cyst (n=8, 34.7%), followed by nasolabial cyst (n=4, 13.04%). Group 3 consisted of majority cases; including benign, premalignant and malignant lesions. Here mean age was 42.4 year and M:F ratio was 1.3:1. Among benign lesions, Fibroma and its subtypes and ameloblastoma were the most common tumors along with a each case of inflammatory myofibroblastic tumor, adenomatoid odontogenic tumor, lipoma, neurofibroma and psuedoepitheliomatous hyperplasia. Each case of verrucous hyperplasia and carcinoma in situ were included as premalignant conditions. Both were male with mean age of 50 year. Among malignant lesions, squamous cell carcinoma (n=20, 47.6%) was the most common tumor along with two cases of intraosseous mucoepidermoid carcinoma (n=2, 4.8%) and each case of plasmacytoma (n=1, 2.3%) and oral malignant melanoma (n=1, 2.3%).

Table 1: Shows Frequency of Various Lesions according to Number of Cases, Gender, Mean Age and Histopathological Diagnosis

Group	Total Number of Cases	M:F	Mean Age	Diagnosis	Number of Cases				
Group 1 (inflammatory/ reactive conditions)	29	1.4:1	35 year	Chronic Mixed inflammation	17				
				Pyogenic granuloma	4				
				Fibroepithelial polyp	2				
				Osteomyelitis	2				
				Submucosal fibrosis	1				
				Mucormycosis	1				
				Pericoronitis	1				
				Inflamed epulis	1				
				Radicular cyst	8				
				Nasolabial cyst	4				
Group 2 (cystic lesions)	23	1.5:1	24.5 year	Dentigerous cyst	3				
				Periapical cyst	3				
				Mucus retention cyst	2				
				Orthokeratinised cyst	1				
				Epidermoid cyst	1				
				Odontogenic keratocyst	1				
				Group 3 Benign	42(Total) 16	1.3:1 1:1	42.4 year 29.4 year	Fibroma and its subtype	8
								Ameloblastoma	3
				Preneoplastic	2	--	50 year	Inflammatory myofibroblastic tumor	1
								Adenomatoid odontogenic tumor	1
Pseudoeplitheliomatous hyperplasia	1								
Neurofibroma	1								
Lipoma	1								
Verrucous hyperplasia	1								
Carcinoma in situ	1								
Malignant	24	1.4:1	48 year	Squamous cell carcinoma	20				
				Mucoepidermoid carcinoma	2				
				Plasmacytoma	1				
				Malignant melanoma	1				
Total Cases	94			Mean age: 34.9 year					

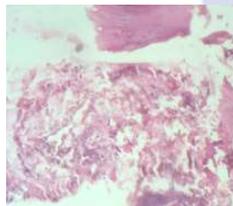


Figure 1

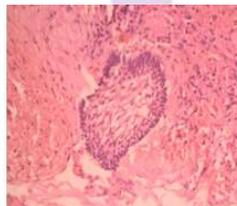


Figure 2



Figure 3

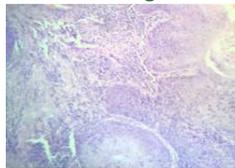


Figure 4

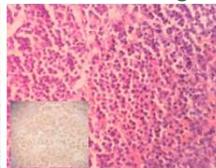


Figure 5

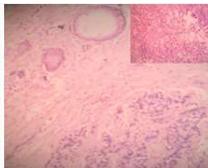


Figure 6

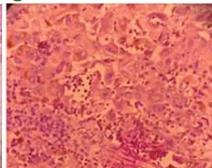


Figure 7

Figure 1: Mucormycosis Showing Large, Nonseptate, Branching Fungal Hyphae. Tissue Necrosis is seen as it is Angioinvasive.(HandE Stain 400X); **Figure 2:** Ameloblastoma Showing Odontogenic Epithelial Islands with Peripheral Palisading of Columnar Cells and Stellate Reticulum like Cells.(HandE Stain 400X); **Figure 3:** Hypercellular Fibrous Connective Tissue within which Irregularly Shaped Calcifications suggestive of cemento-ossifying fibroma (HandE Stain 100X); **Figure 4:** Carcinoma in Situ showing Severely Dysplastic Squamous Epithelium with Intact Basement Membrane .(HandE Stain 400X); **Figure 5:** Plasmacytoma showing Monomorphous Population of Plasma Cells having Eccentric Placed Nucleus and Abundant Cytoplasm. (HandE Stain 400X)(Inset Shows CD 138 Positivity on Immunohistochemistry); **Figure 6:** Mucoepidermoid Carcinoma showing Cords and Sheets of Mucous Cells, some clear cells and Nests of Squamous Cells. Inset Shows Extracytoplasmic Mucin. (HandE Stain 400X); **Figure 7:** The cells are highly pleomorphic in nests with dense intra and extra- cytoplasmic melanin pigment. Inset shows positivity for Melan A confirms the diagnosis of malignant melanoma.

DISCUSSION

It is an accepted fact that microscopic analysis is the gold standard for the diagnosis of most lesions. Present study demonstrate general profile of oral lesions and attempts best to report and classify them. It is based on histopathological examination of oral lesions in correlation with clinical findings to predict significant level of patients' prognostic outcome. It was difficult to compare the results with other studies as they were performed in a specific group of lesions^{9,10} and age.¹¹ Teye *et al* studied 638 jaw swellings and broadly categorized as benign tumors, cysts, infective lesions and malignant neoplasms. M: F ratio was 1.1:1. Mean age was 32.36 ± 17.14 years with a range of 3 to 84 years. Ameloblastoma was the commonest odontogenic tumor followed by fibromyxoma.⁷ Elif Pecker *et al* studied jaw lesions for correlation of clinical and histopathological diagnosis. In this study the lesions were divided into three groups that is inflammatory/developmental lesions, cystic lesion and tumor and tumor like lesions. Here distribution of lesions in each group was 26.9%, 53.6% and 19.5% respectively. Mean age was 40 ± 1.9 years.⁸ We followed the same simple and most acceptable way of classification for easy implementation of treatment and prognosis part for clinician. Typically inflammatory swellings are expected to be the type of oral lesions that are caused by mechanical and chemical trauma, radiation injury, infections and immunological mechanisms. In our study they formed 30.9% of all oral lesions. Occurrence of this group of lesions is little more frequently seen in males(1.4:1) as seen in this study and was also supported by Mosby *et al*.¹² Alveolar bone mucormycosis is rare that frequently occurs after tooth extraction in immunocompromised patients and requires aggressive clinical management(Figure 1).¹³ Osteomyelitis of the jaws also occur in immuno-compromised patients both locally and generally following dental treatment. It is a rare condition in jaw characterized by exposed bone in mouth which fails to heal after appropriate intervention.¹⁴ The embryonic "resting" epithelium (Also termed cell rests) is usually dormant or, undergoes atrophy but when stimulated may be due to inflammation, may form a cyst.(15) Within group 2 lesions, radicular cysts were the most prevalent cyst(n=8, 34.7 %) followed by nasaolabial cysts(n=4,17.4 %). A greater incidence was in males than in females(1.5:1) which was confirmed by conclusion of other study.(16)Radicular cysts were the most biopsied lesions followed by dentigerous cysts and residual cysts in cystic lesions and these data support the data presented by Nuñez Urrutia *et al*.¹⁷ The cyst lining Epithelium may be derived in some case from 1) Respiratory epithelium from maxillary sinus when the periapical communicates with the maxillary sinus 2) Oral epithelium from a

fistulous tract 3) Oral epithelium proliferating apically from a periodontal pocket. Their incidence is highest in third and fourth decade of lifes with male dominance.¹⁸ Third major group includes benign, preneoplastic and malignant lesions(n=42, 44.6%). The most frequent lesion included in benign category was fibroma, a benign neoplasm of fibroblastic origin, is reactive in nature and represents a reactive hyperplasia of fibrous connective tissue in response to local irritation or trauma rather than true neoplasm.¹⁹ Ameloblastoma is a benign tumor that is "usually unicentric, nonfunctional, intermittent in growth, anatomically benign and clinically persistent." About 80% of ameloblastomas occurs in the mandible.(20)The World Health Organization (WHO) (1991) defined ameloblastoma as a benign but locally aggressive tumor with a high tendency to recur, consisting of proliferating odontogenic epithelium lying in a fibrous stroma (Figure 2).²¹ The main modality of treatment is surgery, with wide resection recommended due to the high recurrence rate. Cement forming periodontogenic tumor, cementossifying fibroma showed histologically hypercellular fibrous connective tissue within which is irregularly shaped calcifications.(Figure 3) The recommended treatment is enucleation of smaller ossifying fibromas, curettage of lesions where no clear radiolucency is present around the lesion and mono-bloc resection with bone reconstruction for larger tumors in close proximity to the inferior border of the mandible.²² Despite the easy access of oral cavity for direct visual examination and advances in treatment for Oral Squamous Cell Carcinoma, the 5-year survival rate remains at <60%.²³ Identification of premalignant lesions or early carcinoma with intensified follow up assessments is an ideal strategy to improve survival rates. The premalignant and malignant lesions are closely related lesions which cannot be diagnosed clinically. Therefore, the diagnosis must be established histopathologically. Accurate histopathologic diagnosis depends on an adequate depth of the biopsy specimen and the adjacent normal epithelium.²⁴ Present study also includes two rare preneoplastic lesions that are verrucous hyperplasia and carcinoma in situ (Figure 4). Plasmacytoma is a disorder characterized by uncontrolled monoclonal proliferation of plasma cells that causes overproduction of immunoglobulins in blood (Figure 5).²⁵ Plasmacytoma is clinical entity where it is very difficult to confirm the diagnosis without radiological, histopathological, immunohistochemical and other supportive investigative modalities. In our case radiological findings were correlated with histopathological findings and for establishing the final diagnosis confirmed by Immunohistochemistry (IHC) positivity for CD138. Mucoepidermoid carcinomas infrequently arise

centrally within the mandible as in our two cases (Figure 6). They usually arise in the salivary glands or sinonasal cavities. On radiograph, mucoepidermoid carcinoma in the jaw typically appear more infiltrative.²⁶ Intraosseous mucoepidermoid carcinoma is more common in middle-aged adults and have a slight female predilection. It is three times more common in the mandible than in the maxilla and are most often found in the area of the molars and mandibular ramus. The most frequent symptom is cortical bulging, although some lesions may be discovered as an accidental finding on radiographs. The main modality of treatment for patients with this neoplasm is radical surgical resection, offering a greater chance of cure than the more conservative procedures, such as enucleation or curettage, that have great possibility of recurrence and tumor metastasis. Oral mucosal malignant melanoma is rare representing about 0.5% of oral malignancies and less than 0.01% of all oral biopsies (Figure 7). It arise in adults with an average age of about 55 years. Though surgical excision is the gold standard treatment, it may be combined with chemotherapy, radiotherapy and immunotherapy²⁷. The malignant lesions are The global burden because of the aging and growth of the world population alongside an increasing adoption of cancer-causing behaviors like smoking, betel quid chewing and alcohol consumption.²⁸ Oral Cancer is the 6th most frequent malignant tumour.²⁹ There are nearly 300 Dental Colleges in all over the India. Despite number of Dental surgeon passing out from so many Dental institutions, Still India stands first among high incidence of oral cancer cases. In support of present study, Monika *et al* also proves the maximum number of cases of malignant lesions.¹⁶

CONCLUSIONS

The diagnosis of dental lesions should be based on clinical, radiographic and histopathologic features. Oral and Maxillofacial Surgeons must establish the histological diagnosis of their cases by routine biopsy and provide an adequate treatment, which might involve further procedures. This will prevent unnecessary treatments, delayed surgical operations and disease related mortality.

REFERENCES

1. Joblonski S. Illustrated Dictionary of Dentistry. WB Saunders Company. 1982: 104.
2. Tissue submission policy Available from: <http://www.aaomp.org/general/tissue.htm>. [Last accessed on May 2011].
3. Mota-Ramírez A, Silvestre FJ, Simó JM. Oral biopsy in dental practice.
4. Raymond JM *et al*. The use of biopsy in dental practice: The position of the American Academy of Oral and maxillofacial pathology. *General Dentistry* 2007; 457–461.
5. Nary Filho H, Matsumoto MA, Fraga SC, Gonçalves ES, Sérvulo F. Periapical radiolucency mimicking an odontogenic cyst. *Int Endod J* 2004;37:337-44.
6. Koivisto T, Bowles WR, Rohrer M. Frequency and distribution of radiolucent jaw lesions: A retrospective analysis of 9,723 cases. *J Endod.* 2012;38:729–32. [PubMed: 22595103].
7. Taye JemilatLasisi *et al*. Appraisal of jaw swellings in a Nigerian tertiary healthcare facility. *J Clin Exp Dent.* 2013;5(1):e42-7.
8. ElifPeker, *et al*. A 5 year retrospective study of biopsied jaw lesions with the assessment of concordance between clinical and histopathological diagnoses. *J Oral MaxillofacPathol.* 2016 Jan-Apr; 20(1): 78–85.
9. Regezi JA. Odontogenic cysts, odontogenic tumors, fibrousseous, and giant cell lesions of the jaws. *Mod Pathol* 2002;15:331-41.
10. Mosqueda-Taylor A, Irigoyen-Camacho ME, Diaz-Franco MA, Torres-Tejero MA. Odontogenic cysts. Analysis of 856 cases. *Med Oral* 2002;7:89-96.
11. Wang YL, Chang HH, Chang JY, Huang GF, Guo MK. Retrospective survey of biopsied oral lesions in pediatric patients. *J Formos Med Assoc* 2009;108:862-71.
12. Mosby S, Magar S, Magar S, Ranpise SG, Agarwal PK, Agarwal S. Assessment of clinic-pathologic discrepancy in diagnosis of Jaw lesions: A retrospective analysis. *IAIM*, 2016; 3(8): 140-145.
13. Orval E. Brown, Richard Finn, DDS. Mucormycosis of the mandible. 1986;44(2)132-136.
14. Reid IR. Osteonecrosis of the jaw: who gets it, and why. *Bone.* 2009;44:4–10.
15. Prockt AP, Schebela CR, Maito FD, Sant'Ana-Filho M, Rados PV. Odontogenic cysts: analysis of 680 cases in Brazil. *Head Neck Pathol* 2008;2:150-6.
16. Monika Aroquiadasse, Mariappan J Daniel, Subramanian V Srinivasan, Vannathan K Jimsha, Jignesh Modha, DurgadeviPancharethinam. A retrospective study of cysts and tumors of the oral cavity. *Journal of Indian academy of oral medicine and radiology* 2017. 29(1)2-6
17. Nuñez Urrutia S, Figueiredo R, Gay Escoda C. Retrospective clinicopathological study of 418 odontogenic cysts. *Med Oral Patol Oral Cir Bucal.*, 2010; 15: e767 73.
18. P Venkatalakshmi Aparna *et al*. Bilateral radicular cyst of the mandible: A rare case report. 2018;9(1)37-39.
19. Barker DS, Lucas RB. Localised fibrous overgrowths of the Oral mucosa. *Br J Oral Surg* 1967;5:86-92
20. Becelli R, Carboni A, Cerulli G, Perugini M, Iannetti G. Mandibular ameloblastoma: analysis of surgical treatment carried out in 60 patients between 1977 and 1998. *J Craniofac Surg.* 2002 May; 13(3):395-400;
21. Kramer IR, Pindborg JJ, Shear M. 2nd ed. Berlin: Springer-Verlag; 1992. *Histological Typing of Odontogenic Tumours.* WHO International Histological Classification of Tumours; pp. 11–14.

22. Titinchi F, Morkel J Ossifying Fibroma: Analysis of Treatment Recurrence Patterns. *J Oral Maxillofac Surg.* 2016 Methods and Dec; 74(12):2409-2419.
23. Liu SY, Lu CL, Chiou CT, *et al.* Surgical outcomes and prognostic factors of oral cancer associated with betel quid chewing and tobacco smoking in Taiwan. *Oral Oncol.* 2010;46(4):276-282.
24. Preeti Sharma, Vijay Wadhwan, Pooja Aggarwall, Anamika Sharma Oral verrucous hyperplasia versus oral verrucous carcinoma: A clinicopathologic dilemma revisited using p53 as immunohistochemical marker. 2016; 20(3):362-368
25. Rodriguez-Caballero B, Sanchez-Santolino S, Gracia-Montesinos-Perea B, Gracia-Reija MF, Gomez-Roman J, Saiz-Bustillo R. Mandibular solitary plasmacytoma of the jaw: a case report. *J Med Oral Pathol Cir Bucal.* 2011;16(5):e647-50.
26. Weber AL. Mandible. In: Som PM, eds. *Head and neck imaging.* 2nd ed. St Louis, Mo: Mosby-Year Book, 1991; 399.
27. Raman Preet Kaur Bhullar, Amandeep Bhullar, Mamata S. Kamat *et al.* Primary melanoma of oral mucosa: A case report and review of literature. *Dent Res J (Isfahan).* 2012 May-Jun; 9(3): 353-356
28. Parkin DM *et al.* Global Cancer Statistics 2002. *CA J Clin.* 2005; 55:74-108.
29. Sciubba JJ. Oral cancer. The importance of early diagnosis and treatment. *Am J Clin Dermatol.* 2001; 2: 239-51.

Source of Support: None Declared
Conflict of Interest: None Declared

