

UNIVERSIDADE ESTADUAL DE CAMPINAS FACULDADE DE ODONTOLOGIA DE PIRACICABA

ANDERSON MAURICIO PAIVA E COSTA

FREQUÊNCIA DAS LESÕES EM ASSOALHO DE BOCA: UM ESTUDO MULTICÊNTRICO DE 4.287 CASOS.

FREQUENCY OF FLOOR OF THE MOUTH LESIONS: A MULTICENTER STUDY OF 4,287 CASES.

Piracicaba 2020

ANDERSON MAURICIO PAIVA E COSTA

FREQUÊNCIA DAS LESÕES EM ASSOALHO DE BOCA: UM ESTUDO MULTICÊNTRICO DE 4.287 CASOS.

FREQUENCY OF FLOOR OF THE MOUTH LESIONS: A MULTICENTER STUDY OF 4,287 CASES.

Dissertação apresentada à Faculdade de Odontologia de Piracicaba da Universidade Estadual de Campinas como parte dos requisitos exigidos para a obtenção do título de Mestre em Estomatopatologia, na Área de Estomatologia.

Dissertation presented to the Piracicaba Dental School of the University of Campinas in partial fulfillment of the requirements for the degree of Master in Oral Medicine and Oral Pathology, in Stomatology area.

ORIENTADOR: PROF. DR. HELDER ANTONIO REBÊLO PONTES COORIENTADOR: PROF. DR. FELIPE PAIVA FONSECA

ESTE **EXEMPLAR** CORRESPONDE A VERSÃO FINAL DA DISSERTAÇÃO DEFENDIDA PELO ALUNO **ANDERSON** MAURICIO PAIVA E COSTA E ORIENTADA PELO PROF. DR. REBÊLO ANTONIO HELDER PONTES.

> Piracicaba 2020

Ficha catalográfica Universidade Estadual de Campinas Biblioteca da Faculdade de Odontologia de Piracicaba Marilene Girello - CRB 8/6159

C823f	Costa, Anderson Mauricio Paiva e, 1989- Frequência das lesões em assoalho de boca : um estudo multicêntrico de 4.287 casos / Anderson Mauricio Paiva e Costa. – Piracicaba, SP : [s.n.], 2020.
	Orientador: Helder Antonio Rebêlo Pontes. Coorientador: Felipe Paiva Fonseca. Dissertação (mestrado) – Universidade Estadual de Campinas, Faculdade de Odontologia de Piracicaba.
	 Epidemiologia. 2. Soalho bucal. I. Pontes, Helder Antonio Rebêlo. II. Fonseca, Felipe Paiva, 1986 III. Universidade Estadual de Campinas. Faculdade de Odontologia de Piracicaba. IV. Título.

Informações para Biblioteca Digital

Γ

Título em outro idioma: Frequency of floor of the mouth lesions: multicenter study of 4.287 cases **Palavras-chave em inglês:** Epidemiology Mouth floor **Årea de concentração:** Estomatologia **Titulação:** Mestre em Estomatopatologia **Banca examinadora:** Helder Antonio Rebêlo Pontes [Orientador] Sérgio Elias Vieira Cury Alan Roger dos Santo Silva **Data de defesa:** 14-02-2020 **Programa de Pós-Graduação:** Estomatopatologia

Identificação e informações acadêmicas do(a) aluno(a) - ORCID do autor: https://orcid.org/0000-0001-8953-4065 - Currículo Lattes do autor: http://lattes.cnpq.br/3955490148793360



UNIVERSIDADE ESTADUAL DE CAMPINAS Faculdade de Odontologia de Piracicaba

A Comissão Julgadora dos trabalhos de Defesa de Dissertação de Mestrado, em sessão pública realizada em 14 de fevereiro de 2020, considerou o candidato ANDERSON MAURICIO PAIVA E COSTA aprovado.

PROF. DR. HELDER ANTONIO REBÊLO PONTES

PROF. DR. SÉRGIO ELIAS VIEIRA CURY

PROF. DR. ALAN ROGER DOS SANTOS SILVA

A Ata da defesa, assinada pelos membros da Comissão Examinadora, consta no SIGA/Sistema de Fluxo de Dissertação/Tese e na Secretaria do Programa da Unidade.

DEDICATÓRIA

Dedico este trabalho na sua integralidade ao Prof. Dr. Helder Antonio Rebêlo Pontes, pelo exemplo de profissional, por não medir esforços para ajudar a quem dele precisa, além de ser um pai para todos que o cercam e aprendem diariamente com seus ensinamentos profissionais e pessoais.

AGRADECIMENTOS

Ao Prof. Dr. Hélder Antônio Rebelo Pontes, pela orientação incondicional de mais essa etapa da minha formação profissional e pessoal.

À Universidade Estadual de Campinas (UNICAMP), na pessoa dos professores da Faculdade de Odontologia de Piracicaba (FOP) Dr. Oslei Paes de Almeida, Dr. Márcio Ajudarte Lopes, Dr. Alan Roger, Dr. Pablo Vargas.

Aos meus familiares por estarem sempre presentes, sendo o pilar da minha trajetória.

A minha namorada Joyce Rêgo, inseparável e incansável em todos os momentos, sempre presente e uma das minhas maiores motivadoras nessa jornada.

A todos os colegas de pós-graduação, em especial à Lígia Akiko, e ao meu amigo agora Dr. Ricardo Carvalho, parceiros inseparáveis.

Ao professor Dr. Felipe Fonseca pelas orientações e preciosas informações para a elaboração desse trabalho.

À professora Dra. Flávia Pontes, pelos ensinamentos diários.

Aos colegas de residência, pela parceria e crescimento profissional.

Aos Amigos Lucas Lacerda, pelo esforço imensurável para a realização desse trabalho, e Raíra Brito pelo acolhimento durante minha estadia em Piracicaba e amizade incontestável.

RESUMO

O objetivo deste estudo foi investigar a frequência em importantes centros de patologia oral na América latina de lesões em assoalho bucal. Um estudo transversal foi conduzido com biópsias obtidas de janeiro de 1978 a dezembro de 2018 em nove centros latino-americanos de patologia oral e maxilofacial. Gênero, idade e diagnóstico histopatológico foram avaliados. Os arquivos foram analisados utilizando-se métodos descritivos. De 114.893 amostras, 4.287 lesões (3,73%) ocorreram no soalho bucal. Brasil demonstrou a maior quantidade de casos (4.030 casos; 94%), seguido pelo México (198 casos; 4,6%) e Argentina (59 casos; 1,4%). Lesões benignas representaram 67,2% (2.883 casos), seguido por 32,8% (1.404 casos) de desordens malignas. Lesões de origem epitelial foram mais frequentes (1.964 cases; 45,8%). Seguido por glândulas salivares (1.245 casos; 29%) e lesões de tecido mole (473 casos; 11%). Os subtipos histológicos mais comuns foram carcinoma de células escamosas oral (1.347 casos; 31,4%), rânula (724 casos; 16,8%), leucoplasia oral (476 casos; 11,1%) e hiperplasia fibrosa inflamatória (239 casos; 5,6%). Homens (2.118 cases) foram mais afetados que mulheres (1.900 casos). No presente estudo, lesões em assoalho bucal representaram 3,73% das biópsias submetidas aos serviços de patologia oral e carcinoma de células escamosas, rânula e leucoplasia foram as lesões mais comuns.

Palavras-chave: Epidemiologia, Assoalho bucal.

ABSTRACT

The aim of this study was to investigate the frequency of oral lesions in the floor of the mouth from representative oral pathology centers in Latin America. A cross-sectional study was conducted on biopsies obtained from January of 1978 to December of 2018 at nine Latin America oral and maxillofacial pathology centers. Gender, age and histopathological diagnosis were evaluated. Data were analyzed using descriptive methods. From 114,893 samples, 4,287 lesions (3.73%) occurred in the floor of the mouth. Brazil showed the highest amount of cases (4,030 cases; 94%), followed by Mexico (198 cases; 4.6%) and Argentina (59 cases; 1.4%). Benign lesions represented 67.2% (2,883 cases), followed by 32.8% (1,404 cases) of malignant disorders. Lesions of epithelial origin were more frequent (1,964 cases; 45.8%), followed by salivary glands (1,245 cases; 29%) and soft tissue lesions (473 cases; 11%). The most common histological subtypes were oral squamous cell carcinoma (1,347 cases; 31.4%), ranula (724 cases; 16.8%), oral leukoplakia (476 cases; 11.1%) and inflammatory fibrous hyperplasia (239 cases; 5.6%). Males (2,118 cases) were more affected than females (1,900 cases). In the current study, lesions in the floor of the mouth represented 3.73% of biopsies submitted to oral pathology services and oral squamous cell carcinoma, ranula and leukoplakia were the most common lesions.

Keywords: Epidemiology, Floor of mouth.

SUMÁRIO

1	INTRODUÇÃO10
2	ARTIGO: FREQUENCY OF FLOOR OF THE MOUTH LESIONS: A MULTICENTER STUDY OF 4.287 CASES
3	CONCLUSÃO
	REFERÊNCIAS
	ANEXOS
	Anexo 1- Verificação de originalidade e prevenção de plágio

1 INTRODUÇÃO

O assoalho bucal é uma localização anatômica de interesse dos profissionais da saúde, em especial os cirurgiões dentistas. Neste local, inúmeras lesões benignas e malignas podem se fazerem presentes. Esta estrutura anatômica é constituída totalmente por tecido mole, em forma de "U", situado abaixo da língua, tendo sua porção principal localizado logo à frente desta, enquanto outras duas porções estendem-se na região lateral. Patologias ou alterações de desenvolvimento localizadas nessa região podem afetar diretamente, a fonação e a deglutição. (La porte, 2011).

A região de assoalho de boca é constituída por estruturas de tecidos de várias origens (adiposo, glandular, muscular, epitelial), o que pode explicar os diversos tipos de patologias que podem se originar nesta região anatômica (Boko E, 2014; Carlon ER, 2016). Para o melhor de nosso entendimento inexistem estudos que avaliem a frequência de lesões em assoalho de boca. Assim como estudos que associem tais lesões com parâmetros clínicos. Portanto, o objetivo deste trabalho foi realizar um estudo multicêntrico sobre a frequência das patologias situadas no assoalho bucal, assim como correlacionar as lesões com um perfil clínico.

2 ARTIGO

WHAT IS THE FREQUENCY OF FLOOR OF THE MOUTH LESIONS? A DESCRITIVE STUDY OF 4,287 CASES.

Anderson Mauricio Paiva e Costa^{1,2}, Flávia Sirotheau Corrêa Pontes¹, Lucas Lacerda de Souza^{1,2}, Ana Carolina Carneiro Cardoso¹, Michelle Carvalho de Abreu^{1,2}, Priscilla Flores Silva Gonçalves^{1,2}, Márcio Ajudarte Lopes², Alan Roger Santos-Silva², Pablo Agustin Vargas², Bruno Augusto Benevenuto de Andrade³, Kelly Tambasco Bezerra³, Mário José Romañach³, Ricardo Santiago Gomez⁴, Rafael Ferreira e Costa⁴, Maria Sissa Sant'Ana⁴, Décio dos Santos Pinto Júnior⁵, Danyel Elias da Cruz Perez⁶, Jurema Freire Lisboa de Castro⁶, Tatiana Nayara Libório-Kimura⁷, Jeconias Câmara⁷, Paulo Victor Mendes Penafort⁷, Felipe Paiva Fonseca^{4,11}, Oslei Paes de Almeida², Adalberto Mosqueda-Taylor⁸, Mireya Olmedo-Campos⁸, Ramiro Alejandro Tomasi⁹, Ruth Salomé Ferreyra⁹, Waqas Khan¹⁰, Hélder Antônio Rebelo Pontes^{1,2}.

¹Service of Oral Pathology, João de Barros Barreto University Hospital, Federal University of Pará, Belém/Brazil.

²Oral Diagnosis Department (Pathology and Semiology Areas), Piracicaba Dental School, University of Campinas, Piracicaba/Brazil.

³Department of Oral Diagnosis and Pathology, School of Dentistry, Federal University of Rio de Janeiro, Rio de Janeiro/Brazil.

⁴Department of Oral Surgery and Pathology, School of Dentistry, Federal University of Minas Gerais, Belo Horizonte/Brazil.

⁵Oral Pathology Department, Dental School, University of São Paulo, São Paulo/Brazil.

⁶Oral Pathology Department, Dental School, Federal University of Perrnambuco, Recife/Brazil.

⁷Department of Pathology and Legal Medicine, Federal University of Amazonas, Manaus/Brazil.

⁸Health Care Department, Universidad Autónoma Metropolitana Xochimilco, Mexico City/Mexico.

⁹Department of Oral Pathology, Dental School, National University of Córdoba, Córdoba/Argentina.

¹⁰The School of Clinical Dentistry, Sheffield/UK.

¹¹Department of Oral Pathology and Oral Biology, School of Dentistry, University of Pretoria, Pretoria, South Africa.

Corresponding author: Prof. Dr Helder Antônio Rebelo Pontes; João de Barros Barreto University Hospital, Department of Surgery and Oral Pathology, Mundurucus Street, nº 4487, Zip Code 66073-000, Belém, Pará, Brazil. Telephone +55 91 32016786. E-mail address: https://www.harp@ufpa.br

ABSTRACT

The aim of this study was to investigate the frequency of oral lesions in the floor of the mouth from representative oral pathology centres in Latin America. This study was conducted on biopsies obtained from January of 1978 to December of 2018 at nine Latin America oral and maxillofacial pathology centres. Gender, age and histopathological diagnosis were evaluated. Data were analysed using descriptive methods. From 114,893 samples, 4,287 lesions (3.73%) occurred in the floor of the mouth. Brazil showed 4,030 cases (94%), Mexico 198 cases (4.6%) and Argentina 59 cases (1.4%). Benign lesions represented 67.2% (2,883 cases), followed by 32.8% (1,404 cases) of malignant disorders. Lesions of epithelial origin were more frequent (1,964 cases; 45.8%), followed by salivary glands (1,245 cases; 29%) and soft tissue lesions (473 cases; 31.4%), ranula (724 cases; 16.8%), oral leukoplakia (476 cases; 11.1%) and inflammatory fibrous hyperplasia (239 cases; 5.6%). The lesion affected males in 2,118 cases and females in 1,900 cases. In the current study, lesions in the floor of the mouth represented 3.73% of biopsies submitted to oral pathology services and oral squamous cell carcinoma, ranula and leukoplakia were the most common lesions.

Keywords: Epidemiology; floor of the mouth; benign; malignant.

INTRODUCTION

The floor of the mouth (FOM) is a horizontally aligned U-shaped space situated in the part of the oral cavity that lies beneath the tongue¹. It is commonly associated with a wide variety of local and systemic pathologies². In recent years, epidemiological studies have been performed worldwide and have shown that FOM lesions represented a notable number, varying in different countries³⁻⁶. Thus, the aim of this study is to present a multicenter study of the lesions affecting the FOTM from nine representative Latin America oral and maxillofacial pathology centres.

MATERIALS AND METHODS

Study design and ethical approval

This study was performed in the files from nine independent oral and maxillofacial diagnostic centres from Latin America. The samples of the centres were retrieved up to a period of 40-years, from January of 1978 to December of 2018. A total of 114,893 samples were analysed. The diagnosis centres were from Brazil (Service of Oral Pathology, João de Barros Barreto University Hospital, Federal University of Pará, Belém/Brazil; Oral Diagnosis Department (Pathology and Semiology), Piracicaba Dental School, University of Campinas, Piracicaba/Brazil; Oral Pathology, Dental School, Federal University of Rio de Janeiro, Rio de Janeiro/Brazil; Department of Oral Surgery and Pathology, School of Dentistry, Federal University of Minas Gerais, Belo Horizonte/Brazil; Oral Pathology Department, Dental School, University of São Paulo, São Paulo/Brazil; Oral Pathology Department, Dental School, Federal University of Pernambuco, Recife/Brazil; Department of Pathology and Legal Medicine, Federal University of Amazonas, Manaus/Brazil), Mexico (Oral Pathology Private Clinics, Coyoacán/Mexico) and Argentina (Department of Stomatology, Dental School, National University of Córdoba, Córdoba/Argentina). The expert oral pathologists of

each centre evaluated the samples. The ethical committee of the João de Barros Barreto University Hospital approved this work under approval number 3.381.233. The patient's identity remained anonymous according to the Declaration of Helsinki.

Samples

Lesions in the FOM were retrieved, and each of the analysed centres recovered data regarding age, sex and final histopathological diagnosis. All lesions were reclassified according to the World Health Organisation Classification of Head and Neck Tumours Update published in 2017⁷ and other categories were classified according to Textbook of Oral and Maxillofacial Pathology, 4th Edition⁸. The exclusion criteria were samples that did not present lesion in the FOM, as well as those whose final diagnosis did not follow the criteria established in the current study. The lesions in which diagnosis was not possible to be performed were excluded of the current study.

Data analysis

The collected data were tabulated in Microsoft Excel® for epidemiologic analysis. Descriptive and quantitative data analysis was performed using the Statistical Package for Social Sciences (SPSS) software, version 22.0 (SPSS Inc., Chicago, IL, USA).

RESULTS

From 114,893 samples, 4,287 (3.73%) were in the FOM. Figure 1 shows the distribution of the lesions in in the centres analysed, evidencing that Brazil showed 4,030 cases (94%), Mexico 198 cases (4.6%) and Argentina 59 cases (1.4%). Table 1 and 2 show benign and malignant lesions found in the current study, respectively. Benign disorders represented 67.2% (2,883 cases), followed by 32.8% (1,404 cases) of malignancies. Lesions of epithelial

origin were more frequent (1,964 cases; 45.8%), followed by salivary glands (1,245 cases; 29%) and soft tissue (473 cases; 11%). The most common histological subtypes were oral squamous cell carcinoma (1,347 cases; 31.4%), ranula (724 cases; 16.8%), oral leukoplakia (476 cases; 11.1%) and inflammatory fibrous hyperplasia (239 cases; 5.6%). Males (2,118 cases) were more affected than females (1,900 cases). The patients demonstrated a mean age of 39.5 years old (range of 8-89-year-old).

Table 3 shows the frequency of the lesions according to the age and gender. Salivary gland lesions (SGL) represented salivary glands (1,245 cases; 29%) of the samples. Ranula (724 cases; 16.8%), non-specific chronic inflammatory process (204 cases; 17.5%) and sialolitiasis (183 cases; 15.7%) were the most common lesions. Patients affected by this group of lesions were mostly in age lower than 20 years old (397 cases) and female (710 cases) were more affected than male (452 cases) with a M:F ratio of 1:1.57. Epithelial lesions (EL) were presented in (1,964 cases; 45.8%). Oral squamous cell carcinoma (1,347 cases; 31.4%), oral leukoplakia (476 cases; 11.1%) and papilloma (71 cases; 3.8%) were the most frequently diagnosed lesions. Individuals older than 60 years old (826 cases) and males (1,322 cases) were more affected than females (546 cases) in a M:F ratio of 2.4:1. Soft tissue tumors (STT) represented 447 cases (10.4%). Inflammatory fibrous hyperplasia (226 cases; 50.6%), fibrous hyperplasia (153 cases; 34.2%) and lipoma (37 cases; 8.2%) were the most frequent lesions of this group. Individuals older than 60 years were the most frequently affected by this group of lesions (212 cases) and females (338 cases) were more commonly detected than males (109 cases) in a M:F ratio of 1:3.1.

Infectious diseases (ID) were seen in 30 cases (0.7%). The two lesions observed in the current group were paracoccidioidomycosis (28 cases; 93.3%) and candidiasis (2 cases; 6.7%), and mostly affected patients with ages from 50 to 60 years old. Males (27 cases) were affected more frequently than females (3 cases) with a M: F ratio of 1:9. Dermatological

diseases (DEDI) were observed in 27 cases (0.6%). Lichen planus (16 cases; 53.3%), pemphigus vulgaris (9 cases; 30%) and pemphigoid (2 cases; 6.6%) were the DEDI observed in the FOM. Patients affected by this group of lesions mainly were older than 60 years old (9 cases) and females (21 cases) were more affected than males (6 cases). Physical and chemical injuries (PCI) represented 158 cases of the sample (3.6%). The lesions detected on this group were amalgam tattoo (108 cases; 68.3%), ulcers (42 cases; 26.6%) and exogenous pigmentation (8 cases; 5%). Individuals with age ranging from 50 to 60 years old were mostly identified and females (108 cases) were more affected than male, with a M: F ratio of 1:2.1.

Developmental disorders (DD) were seen in 93 cases (2.1%). Lymphoepithelial cyst (33 cases; 35.5%), dermoid and teratoid cyst (20 cases; 21.5%) and epidermoid cyst (20 cases; 21.5%). People with age lower than 20 years old and ranging from 50 to 60 years old were mostly seen, representing 26 cases each group. DD demonstrated a slight frequency in females (48 cases) than males (45 cases), with a M: F ratio of 1:1.06. Hematological diseases (HD) were six cases (0.13%) and small cell B-cell lymphomas (2 cases; 33.3%) and large B-cell lymphoma (2 cases; 33.3%) were the most common histological types. Patients with age higher than 60 years old represented three cases. This group of lesions did not demonstrate any gender predominance. Oral manifestation of systemic diseases represented two cases and all were patients affected by amyloidosis on the FOM with age older than 60 years old. This group were seen in one man and one woman, with M: F ratio of 1:1.

DISCUSSION

The findings of the current study showed that 3.73% of the analysed cases were presented in the FOM and can be compared with the existing literature of other countries. In a systematic review of oral cancer in Arab countries, Al-Jaber and collaborators observed that FOM was affected in 50% of all sample³. In addition, a study developed in Zimbabwe observed that 18.5% of cases were lesions located in the FOM⁹. Nemes *et al.* investigated the

prevalence of oral lesions in Northeastern Hungary and showed that 27.7% affected the FOM ¹⁰. These differences seem to be related to the number of FOM lesions included in each study, and the number of individuals examined. Besides differences in age, sex and habitual differences between populations, methodology and criteria of diagnosis could influence the results.

In the current study, SGL represented 27.1% of all lesions in the FOM. In a Chinese epidemiological study, it was observed that SGL in the FOM represented 1.5% of all samples¹¹. In the current study, ranula (685 cases) and non-specific chronic inflammatory process (204 cases) and sialolitiasis (183 cases) were most common benign lesions. Previous studies demonstrated that pleomorphic adenoma was the most common lesion in the FOM^{12,13}. According De Oliveira et al. SGL lesions in the FOM tend to be presented as malignancies¹³, which is not similar with the current study. Clinically, SGL tended to affect female patients in age lower than 20 years old. Li *et al.* observed that 1.5% of lesions affected the FOM and male patients were more frequently seen¹¹.

EL were observed in 43.5% and oral squamous cell carcinoma (1,281 cases), oral leukoplakia (454 cases) and papilloma (71 cases) were most commonly seen. A Brazilian study observed that EL in the FOTH comprised just 9.5% of all sample¹⁴. Under clinical evaluation, males older than 60 years old were more affected by this group of lesions^{15,16}. In addition, a previous study of our group showed that oral squamous cell carcinoma of the FOM affected males older than 60 years old¹⁷. It is important emphasize that leukoplakia had higher mean ages than the average age of the patients affected by oral squamous cell carcinoma, suggesting that the majority of oral squamous cell carcinoma was not originated from pre-existing leukoplakias.

STT were observed in 10.4% of cases and inflammatory fibrous hyperplasia (226 cases), fibrous hyperplasia (153 cases) and lipoma (37 cases) were the most common lesions

seen on this study. In Venezuela, a study exploring oral lesions in elderly people showed that STT represented 18% of all analysed sample¹⁸. Clinical aspects evidenced that females older than 60 years old were more affected, consistent with previous epidemiological studies in Chile and Mexico^{19,20}. In a multicentre study above oral sarcomas in Brazil, it was observed that osteosarcoma, Kaposi's sarcoma and chondrosarcoma were the most common malignant STT. This information is not consistent with the current study, since leiomyosarcoma, Kaposi's sarcoma and carcinosarcoma were the most commonly lesion seen²¹.

ID were seen in 0.7% of the analysed samples and paracoccidioidomycosis (28 cases) was the most common entity found this group of lesions. Two of the centres included this group are located in an endemic region for paracoccidioidomycosis. Dutra and collaborators observed that FOM was not a common location for ID, representing just 2% of all cases of paracoccidioidomycosis of their sample²². Under clinical evaluation, ID affected males with age ranging from 50 to 60 years old, consistent with previous literature^{23,24}.

DEDI were shown in 0.6% of the present samples and presented lichen planus (16 cases), pemphigus (9 cases) and pemphigoid (2 cases) as the most common lesion in the FOM. Sultan et al. demonstrated that pemphigus is the most common dermatological lesions with oral manifestation in USA²⁵. Clinically, lesions tended to be presented in females older than 60 years old, as well as previous studies²⁶⁻²⁸. In addition, this group of lesions is not commonly observed in the FOM^{27,28}. PCI corresponded to 3.6% of our cases and the mostly observed lesions were amalgam tattoo (108 cases), ulcers (42 cases) and exogenous pigmentation (8 cases). A Brazilian study observed that amalgam tattoo was also the most common lesion and that FOM was not a frequent location to observe PCI²⁹. In the current study, females were more affected at an age ranging from 50 to 60 years old ²⁹⁻³¹.

DD were observed in 2.1% of all cases seen in the current study and lymphoepithelial cyst (33 cases), dermoid and teratoid cyst (20 cases) and epidermoid cyst (20 cases) were the

most common lesions. Dovigi and colleagues recognized that in USA, DD corresponded to 6.5% of 51,781 cases analysed on their study³². In the present research, the patients of this group of lesions showed a slight female frequency, and people with age younger than 20 years old and ranging from 50 to 60 years old were mostly seen. Nonaka et al. observed that females in the fourth decade of life were more affected by this group of lesions³³.

Regarding HD, they were uncommon lesions in the FOM, representing just 0.13% of all lesions. Deng et al. showed that lymphomas are rare on the FOM³⁴. Clinically, our study demonstrated that these lesions affects patients on age lower than 20 years old and higher than 60 years old, consistent with previous literature^{35,36}. Oral manifestation of systemic diseases were also reported in our study and we had two cases of amyloidosis affecting the floor of the mouth. A Latin America study showed that FOM is affected in 14.3% of all cases³⁷. Under clinical evaluation, this group of lesions commonly affects patients older than 60 years old and demonstrated no gender predominance. Previous literature showed that males with a peak age of 50 years old are more described^{37,38}.

CONCLUSION

This manuscript shows the trends of the oral pathologies of the FOM, and oral squamous cell carcinoma, ranula and leukoplakia were the most common histological types

ACKNOWLEDGMENTS

No acknowledgments.

CONFLICT OF INTEREST STATEMENT

None.

TABLES

Table 1. Benign lesions in the floor of the mouth.

Table 2. Malignant lesions in the floor of the mouth.

Table 3. Frequency of floor of the mouth lesions in Latin America according to age and gender.

REFERENCES

1 - LA'PORTE SJ, JUTTLA JK, LINGAM RK. Imaging the floor of the mouth and the sublingual space. Radiographic. 2011 Sep-Oct;31(5):1215-30.

2 - PURICELLI E, BARREIRO BOB, QUEVEDO AS, PONZONI D. Occurrence of dermoid cyst in the floor of the mouth: the importance of differential diagnosis in pediatric patients. J Appl Oral Sci. 2017; 25(3):341–345.

3 - AL-JABER A, AL-NASSER L, EL-METWALLY A. Epidemiology of oral cancer in Arab countries. Saudi Med J. 2016; 37(3):249–255.

4 - JÉHANNIN-LIGIER K, DEJARDIN O, LAPÔTRE-LEDOUX B, BARA S, COUREAU G, GROSCLAUDE P, et al. Oral cancer characteristics in France: Descriptive epidemiology for early detection. J Stomatol Oral Maxillofac Surg. 2017 Apr; 118(2):84-89.

5 - SAGGI S, BADRAN KW, HAN AY, KUAN EC, ST JOHN MA. Clinicopathologic Characteristics and Survival Outcomes in Floor of Mouth Squamous Cell Carcinoma: A Population-Based Study. Otolaryngol Head Neck Surg. 2018 Jul;159(1):51-58.

6 - SCHWANKE TW, OOMEN KP, APRIL MM, WARD RF, MODI VK. Floor of mouth masses in children: proposal of a new algorithm. Int J Pediatr Otorhinolaryngol. 2013 Sep; 77(9):1489-94.

7 - EL-NAGGAR AK, CHAN JK, GRANDIS JR, TAKATA T, SLOOTWEG PJ. World Health Organization classification of head and neck tumours. Lyon, France: IARC Press; 2017.

8 - NEVILLE BW, DAMM DD, ALLEN CM, et al. Oral & Maxillofacial Pathology, ed 4, Philadelphia, WB Saunders, 2016.

9 - CHIDZONGA MM. Oral malignant neoplasia: a survey of 428 cases in two Zimbabwean hospitals. Oral Oncol. 2006 Feb;42(2):177-83.

10 - NEMES JA, REDL P, BODA R, KISS C, MÁRTON IJ. Oral cancer report from Northeastern Hungary. Pathol Oncol Res. 2008 Mar;14(1):85-92.

11 - LI LJ, LI Y, WEN YM, LIU H, ZHAO HW. Clinical analysis of salivary gland tumor cases in West China in past 50 years. Oral Oncol. 2008 Feb;44(2):187-92.

12 - SUBHASHRAJ K. Salivary gland tumors: a single institution experience in India. Br J Oral Maxillofac Surg. 2008 Dec;46(8):635-8.

13 - DE OLIVEIRA FA, DUARTE EC, TAVEIRA CT, et al. Salivary gland tumor: a review of 599 cases in a Brazilian population. Head Neck Pathol. 2009;3(4):271–275.

14 - PEREIRA JDOS S, CARVALHO MDE V, HENRIQUES AC, DE QUEIROZ CAMARA TH, MIGUEL MC, FREITAS RDE A. Epidemiology and correlation of the clinicopathological features in oral epithelial dysplasia: analysis of 173 cases. Ann Diagn Pathol. 2011 Apr;15(2):98-102.

15 - NEVILLE BW, DAY TA. Oral cancer and precancerous lesions. CA Cancer J Clin. 2002 Jul-Aug;52(4):195-215.

16 - CARVALHO MDE V, IGLESIAS DP, DO NASCIMENTO GJ, SOBRAL AP. Epidemiological study of 534 biopsies of oral mucosal lesions in elderly Brazilian patients. Gerodontology. 2011 Jun;28(2):111-5.

17 - PONTES FS, CARNEIRO JT JR, FONSECA FP, DA SILVA TS, PONTES HA, PINTO DDOS S JR. Squamous cell carcinoma of the tongue and floor of the mouth: analysis of survival rate and independent prognostic factors in the Amazon region. J Craniofac Surg. 2011 May;22(3):925-30.

18 - MUJICA V, RIVERA H, CARRERO M. Prevalence of oral soft tissue lesions in an elderly venezuelan population. Med Oral Patol Oral Cir Bucal. 2008 May 1;13(5):E270-4.

19 - CASTELLANOS JL, DÍAZ-GUZMÁN L. Lesions of the oral mucosa: an epidemiological study of 23785 Mexican patients. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 2008 Jan;105(1):79-85.

20 - ESPINOZA I, ROJAS R, ARANDA W, GAMONAL J. Prevalence of oral mucosal lesions in elderly people in Santiago, Chile. J Oral Pathol Med. 2003 Nov;32(10):571-5.

21 - DE CARVALHO WRS, DE SOUZA LL, PONTES FSC, UCHÔA DCC, CORRÊA DL, DE CÁCERES CVBL, et al. A multicenter study of oral sarcomas in Brazil. Oral Dis. 2020 Jan; 26(1):43-52.

22 - DUTRA LM, SILVA THM, FALQUETO A, PEÇANHA PM, SOUZA LRM, GONÇALVES SS, et al. Oral paracoccidioidomycosis in a single-center retrospective analysis from a Brazilian southeastern population. J Infect Public Health. 2018 Jul - Aug; 11(4):530-533.

23 - GIRARDI FM, SCROFERNEKER ML, GAVA V, PRUINELLI R. Head and neck manifestations of paracoccidioidomycosis: an epidemiological study of 36 cases in Brazil. Mycopathologia. 2012 Mar;173(2-3):139-44.

24 - TRINDADE AH, MEIRA HC, PEREIRA IF, DE LACERDA JCT, DE MESQUITA RA, SANTOS VR. Oral paracoccidioidomycosis: Retrospective analysis of 55 Brazilian patients. Mycoses. 2017 Aug; 60(8):521-525.

25 - SULTAN AS, VILLA A, SAAVEDRA AP, TREISTER NS, WOO SB. Oral mucous membrane pemphigoid and pemphigus vulgaris-a retrospective two-center cohort study. Oral Dis. 2017 May; 23(4):498-504.

26 - SULIMAN NM, ÅSTRØM AN, ALI RW, SALMAN H, JOHANNESSEN AC. Clinical and histological characterization of oral pemphigus lesions in patients with skin diseases: a cross sectional study from Sudan. BMC Oral Health. 2013 Nov 21; 13:66.

27 - CASPARIS S, BORM JM, TEKTAS S, KAMARACHEV J, LOCHER MC, DAMERAU G, et al. Oral lichen planus (OLP), oral lichenoid lesions (OLL), oral dysplasia, and oral cancer: retrospective analysis of clinicopathological data from 2002-2011. Oral Maxillofac Surg. 2015 Jun; 19(2):149-56.

28 - BAGAN J, JIMÉNEZ Y, MURILLO J, BAGAN L. Oral mucous membrane pemphigoid:A clinical study of 100 low-risk cases. Oral Dis. 2018 Mar;24(1-2):132-134.

29 - TAVARES TS, MEIRELLES DP, DE AGUIAR MCF, CALDEIRA PC. Pigmented lesions of the oral mucosa: A cross-sectional study of 458 histopathological specimens. Oral Dis. 2018 Nov;24(8):1484-1491.

30 - GONDAK RO, DA SILVA-JORGE R, JORGE J, LOPES MA, VARGAS PA. Oral pigmented lesions: Clinicopathologic features and review of the literature. Med Oral Patol Oral Cir Bucal. 2012 Nov 1; 17(6):e919-24.

31 - KAUZMAN A, PAVONE M, BLANAS N, BRADLEY G. Pigmented lesions of the oral cavity: review, differential diagnosis, and case presentations. J Can Dent Assoc. 2004 Nov; 70(10):682-3.

32 - DOVIGI EA, KWOK EY, EVERSOLE LR, DOVIGI AJ. A retrospective study of 51,781 adult oral and maxillofacial biopsies. J Am Dent Assoc. 2016 Mar;147(3):170-6.

33 - NONAKA CF, HENRIQUES AC, DE MATOS FR, DE SOUZA LB, PINTO LP. Nonodontogenic cysts of the oral and maxillofacial region: demographic profile in a Brazilian population over a 40-year period. Eur Arch Otorhinolaryngol. 2011 Jun; 268(6):917-22. 34 - DENG D, WANG Y, LIU W, QIAN Y. Oral and maxillofacial non-Hodgkin lymphomas: Case report with review of literature. Medicine (Baltimore). 2017 Sep; 96(35):e7890.

35 - ABDELWAHED HUSSEIN MR. Non-Hodgkin's lymphoma of the oral cavity and maxillofacial region: a pathologist viewpoint. Expert Rev Hematol. 2018 Sep; 11(9):737-748.

36 - TRIANTAFILLIDOU K, DIMITRAKOPOULOS J, IORDANIDIS F, GKAGKALIS A. Extranodal non-hodgkin lymphomas of the oral cavity and maxillofacial region: a clinical study of 58 cases and review of the literature. J Oral Maxillofac Surg. 2012 Dec;70(12):2776-85.

37 - GOUVÊA AF, RIBEIRO AC, LEÓN JE, CARLOS R, DE ALMEIDA OP, LOPES MA. Head and neck amyloidosis: clinicopathological features and immunohistochemical analysis of 14 cases. J Oral Pathol Med. 2012 Feb; 41(2):178-85.

38 - VIGGOR SF, FREZZINI C, FARTHING PM, FREEMAN CO, YEOMAN CM, THORNHILL MH. Amyloidosis: an unusual case of persistent oral ulceration. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 2009 Nov; 108(5):e46-50.

FIGURE LEGENDS

Figure 1. Distribution of sample in the oral pathology diagnosis centres.

Table 1. Benign lesions in the floor of the mouth.

	Ber	nign Lesions		
Benign floor of the mouth pathologies	M (%)	F (%)	Mean age (years) at diagnosis ± Standard deviation	Total (%)
Salivary gland				
Ranula	274 (37.29%)	450 (62.70%)	24.16 ± 16.46	724 (27.7%)
Non-specific chronic inflammatory process	85 (37.84%)	137 (63.06%)	52.28 ± 18.78	222 (8.49%)
Sialolitiasis	88 (44.62%)	108 (55.38%)	46.14 ± 19.89	196 (7.46%)
Retention cyst	22 (37.04%)	33 (62.96%)	32.39 ± 19.39	55 (2.07)
Adenomatous hyperplasia	5 (71.43%)	2 (28.57%)	43 ± 13.38	7 (0.27%)
Pleomorphic adenoma	2 (50%)	2 (50%)	43 ± 6.36	4 (0.15%)
Epithelial				
Oral leukoplakia	227 (47.90%)	249 (51.68%)	60.27 ± 14.84	476 (18.21%)
Hyperkeratosis	19 (43.59%)	20 (56.41%)	55.75 ± 16.27	39 (1.49%)
Papilloma	45 (58.67%)	30 (41.33%)	50.9 ± 18.42	75 (2.87%)
Verruca Vulgar	3 (42.86%)	4 (57.14%)	57.43 ± 20.90	7 (0.27%)
Condyloma	4 (57.14%)	3 (42.86%)	50.14 ± 16.81	7 (0.27%)
Verruciform xanthoma	1 (100%)	0 (0%)	40 ± 12	2 (0.08%)
Soft tissue tumors				
Inflammatory fibrous hyperplasia	45 (17.57%)	194 (82.42%)	60.7 ± 12.77	239 (9.14%)
Fibrous hyperplasia	36 (22.50%)	124 (77.50%)	58.2 ± 14.61	160 (6.12%)
Lipoma	16 (40%)	24 (60%)	57.85 ± 14.67	40 (1.53%)
Hemangioma	2 (33.33%)	7 (66.67%)	63.5 ± 20.48	9 (0.34%)
Neurofibroma	6 (75%)	2 (25%)	29.75 ± 21.87	8 (0.31%)
Pyogenic granuloma	2 (25%)	6 (75%)	50 ± 19.49	8 (0.31%)
Rhabdomyoma	2 (100%)	0 (0%)	49.5 ± 3.5	2 (0.08%)
Lymphangioma	1 (50%)	1 (50%)	53.5 ± 16.5	2 (0.08%)
Myofibroma	1 (100%)	0 (0%)	54 ± 0	1 (0.04%)

Hamartoma	0 (0%)	1 (100%)	3 ± 0	1 (0.04%)
Infectious diseases				
Paracoccidioidomycosis	28 (96.55%)	1 (3.45%)	51.64 ± 10.11	29 (1.11%)
Candidiasis	0 (0%)	2 (100%)	70 ± 4	2 (0.08%)
Dermatological diseases				
Lichen planus	4 (25%)	12 (75%)	54.14 ± 11.56	16 (0.61%)
Pemphigus	2 (20%)	8 (80%)	43.98 ± 16.11	10 (0.38%)
Pemphigoid	0 (0%)	2 (100%)	58 ± 9	2 (0.08%)
Physical and chemical injuries				
Amalgam Tattoo	29 (25.22%)	86 (74.78%)	48.6 ± 15.23	115 (4.4%)
Ulcers	21 (38.30%)	26 (61.70%)	60.14 ± 14.30	47 (1.8%)
Exogenous pigmentation	5 (55.56%)	4 (44.44%)	50.13 ± 10.36	9 (0.34%)
Oral manifestations of systemic diseas	es			
Amiloidosis	1 (50%)	1 (50%)	72 ± 1	2 (0.08%)
Developmental disorders				
Lymphoepithelial cyst	11 (30.56%)	25 (69.44%)	43.28 ± 17	36 (1.38)
Dermoid and teratoid cyst	15 (72.73%)	7 (27.27%)	22.44 ± 15.47	22 (0.84)
Epidermoid cyst	12 (60%)	7 (40%)	29.79 ± 19.44	19 (0.77)
Thrombus	4 (33.33%)	8 (66.67%)	45.25 ± 15.17	12 (0.46)
Varicose veins	3 (37.5%)	5 (62.5%)	60.14 ± 10.71	8 (0.31)
Thyroglossal duct cyst	1 (50%)	1 (50%)	26 ± 18	2 (0.08)

Table 2. Malignant lesions in the floor of the mouth.

	Maligna	nt Lesions		
Malignant floor of the mouth pathologies	M (%)	F (%)	Mean age (years) at diagnosis ± Standard deviation	Total (%)
Salivary gland				
Adenoid cystic carcinoma	7 (36.84%)	12 (63.16%)	$56 \pm 20,89$	19 (1.35)

Mucoepidermoid carcinoma	7 (50%)	7 (50%)	49.82 ± 20.94	14 (1)
Adenocarcinoma	0 (0%)	3 (100%)	49.67 ± 7.54	3 (0.21)
Clear cell hyalinizing carcinoma	0 (0%)	1 (100%)	36 ± 0	1 (0.07)
Epithelial				
Oral squamous cell carcinoma	1,081 (80.62%)	266 (19.38%)	58.93 ± 16.71	1347 (95.94)
Verrucous carcinoma	5 (62.5%)	3 (37.5%)	67 ± 17.36	8 (0.57)
Squamous basaloid carcinoma	2 (100%)	0 (0%)	72 ± 10	2 (0.14)
Carcinoma Cuniculatum	0 (0%)	1 (100%)	86 ± 0	1 (0.07)
Soft tissue tumors				
Leiomyosarcoma	0 (0%)	1 (100%)	53 ± 0	1 (0.07)
Kaposi's sarcoma	0 (0%)	1 (100%)	29 ± 0	1 (0.07)
Carcinosarcoma	1 (100%)	0 (0%)	71 ± 0	1 (0.07)
Hematological diseases				
ALK-positive anaplastic large cell lymphoma	1 (100%)	0 (0%)	18 ± 0	1 (0.07)
MALT Lymphoma	0 (0%)	1 (100%)	64 ± 0	1 (0.07)
Small cell B-cell lymphomas	1 (50%)	1 (50%)	69 ± 43.1	2 (0.14)
Large B-cell lymphoma	1 (50%)	1 (50%)	43 ± 29.34	2 (0.14)

Table 3. Frequency of floor of the mouth lesions in Latin America according to age and gender.

					Age	(years)				
Floor of the mouth pat	thologies	Sex	<20 n (%)	20-30 n (%)	30-40 n (%)	40-50 n (%)	50-60 n (%)	>60 n (%)	NR (%)	Total n (%)
Salivary gland										
Ranula		F	231(51%)	88 (20%)	45(10%)	26 (6%)	21 (5%)	15 (3%)	24 (5%)	450(100%)
		Μ	120(43%)	36 (16%)	41(15%)	22 (8%)	26 (9%)	14 (4%)	15 (5%)	274(100%)
Non-specific inflammatory process	chronic	F	12 (8%)	5 (4%)	9 (7%)	20 (14%)	41 (30%)	41 (30%)	9 (7%)	137(100%)
		Μ	3 (4%)	7 (8%)	6 (7%)	15 (18%)	20 (24%)	25 (30%)	9 (8%)	85 (100%)

Sialolitiasis	F	9 (8%)	14 (13%)	14(13%)	18 (17%)	21 (19%)	27 (25%)	5 (5%)	108(100%)
	Μ	9 (10%)	13 (15%)	8 (9%)	13 (15%)	16 (18%)	21 (24%)	8 (9%)	88 (100%)
Retention cyst	F	6 (18%)	5 (15%)	3 (10%)	8 (24%)	5 (15%)	2 (6%)	4 (12%)	33 (100%)
-	Μ	7 (31,5%)	7 (31,5%)	2 (9%)	2 (9%)	2 (9%)	1 (5%)	1 (5%)	22 (100%)
Adenomatous hyperplasia	F	0 (0%)	0 (0%)	1 (50%)	0 (0%)	0 (0%)	0 (0%)	1 (50%)	2 (100%)
	Μ	0 (0%)	0 (0%)	2 (40%)	0 (0%)	0 (0%)	1 (20%)	2 (40%)	5 (100%)
Pleomorphic adenoma	F	0 (0%)	0 (0%)	0 (0%)	2 (100%)	0 (0%)	0 (0%)	0 (0%)	2 (100%)
	Μ	0 (0%)	0 (0%)	1 (50%)	1 (50%)	0 (0%)	0 (0%)	0 (0%)	2 (100%)
Adenoid cystic carcinoma	F	0 (0%)	0 (0%)	1 (8,5%)	4 (33%)	4 (33%)	2 (17%)	1 (8,5%)	12 (100%)
	Μ	0 (0%)	0 (0%)	1 (14%)	0 (0%)	1 (14%)	4 (58%)	1 (14%)	7 (100%)
Mucoepidermoid carcinoma	F	0 (0%)	0 (0%)	2 (29%)	0 (0%)	2 (29%)	2 (29%)	1 (13%)	7 (100%)
	Μ	0 (0%)	1 (14%)	1 (14%)	1 (14%)	2 (29%)	0 (0%)	2 (29%)	7 (100%)
Adenocarcinoma	F	0 (0%)	0 (0%)	1 (25%)	0 (0%)	2 (75%)	0 (0%)	0 (0%)	3 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Clear cell hyalinizing carcinoma	F	0 (0%)	0 (0%)	1(100%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Epithelial									
Oral leukoplakia	F	2 (1%)	3 (1%)	19 (8%)	23 (9%)	55 (22%)	139 (56%)	8 (3%)	249(100%)
	Μ	1 (1%)	2 (1%)	12 (5%)	39 (17%)	60 (26%)	99 (44%)	14 (6%)	227(100%)
Hyperkeratosis	F	0 (0%)	0 (0%)	0 (0%)	6 (32%)	5 (25%)	6 (32%)	2 (11%)	19 (100%)
	Μ	0 (0%)	2 (10%)	0 (0%)	2 (10%)	8 (40%)	7 (35%)	1 (5%)	20 (100%)
Papilloma	F	1 (3%)	3 (10%)	1 (3%)	5 (17%)	5 (17%)	13 (43%)	2 (7%)	30 (100%)
	Μ	2 (4%)	3 (7%)	8 (18%)	8 (18%)	10 (22%)	12 (27%)	2 (4%)	45 (100%)
Verruca Vulgaris	F	0 (0%)	0 (0%)	0 (0%)	2 (50%)	0 (0%)	2 (50%)	0 (0%)	4 (100%)
	Μ	1 (25%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	2 (75%)	0 (0%)	3 (100%)
Condyloma	F	0 (0%)	1(33,33%)	0 (0%)	0 (0%)	1(33,33%)	1(33,33%)	0 (0%)	3 (100%)
	Μ	0 (0%)	0 (0%)	1 (25%)	2 (50%)	1 (25%)	0 (0%)	0 (0%)	4 (100%)
Verruciform xanthoma	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
	Μ	0 (0%)	1 (50%)	0 (0%)	0 (0%)	1 (50%)	0 (0%)	0 (0%)	2 (100%)

Oral squamous cell carcinoma	F	0 (0%)	2 (1%)	8 (3%)	25 (9%)	74 (28%)	140 (53%)	17 (6%)	266(100%)
	Μ	1 (0%)	1 (0%)	17 (1%)	190(18%)	426(39%)	397(37%)	49 (5%)	1.081(100)
Verrucous carcinoma	F	0 (0%)	0 (0%)	0 (0%)	1 (25%)	0 (0%)	2 (75%)	0 (0%)	3 (100%)
	Μ	0 (0%)	0 (0%)	1 (20%)	0 (0%)	0 (0%)	3 (40%)	1 (20%)	5 (100%)
Squamous basaloid carcinoma	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
	Μ	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	2 (100%)	0 (0%)	2 (100%)
Carcinoma cuniculatum	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	1 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Soft tissue tumors									
Inflammatory fibrous hyperplasia	F	2 (1%)	0 (0%)	3 (2%)	24 (12%)	59 (30%)	94 (48%)	12 (7%)	194(100%)
	Μ	0 (0%)	2 (4%)	4 (9%)	6 (14%)	8 (18%)	24 (53%)	1 (2%)	45 (100%)
Fibrous hyperplasia	F	0 (0%)	1 (1%)	5 (4%)	18 (15%)	40 (32%)	55 (44%)	5 (4%)	124(100%)
	Μ	3 (8%)	1 (3%)	2 (6%)	5 (14%)	8 (22%)	15 (42%)	2 (5%)	36 (100%)
Lipoma	F	0 (0%)	0 (0%)	2 (8%)	6 (25%)	6 (25%)	7 (29%)	3 (13%)	24 (100%)
	Μ	0 (0%)	1 (6%)	0 (0%)	5 (31%)	4 (25%)	6 (38%)	0 (0%)	16 (100%)
Hemangioma	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (14%)	5 (72%)	1 (14%)	7 (100%)
	Μ	1 (50%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (50%)	0 (0%)	2 (100%)
Neurofibroma	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (50%)	1 (50%)	0 (0%)	2 (100%)
	Μ	4 (66%)	0 (0%)	1 (17%)	1 (17%)	0 (0%)	0 (0%)	0 (0%)	6 (100%)
Pyogenic granuloma	F	0 (0%)	1 (17%)	0 (0%)	1 (17%)	1 (17%)	1 (17%)	2 (32%)	6 (100%)
	Μ	0 (0%)	0 (0%)	1 (50%)	0 (0%)	0 (0%)	1 (50%)	0 (0%)	2 (100%)
Rhabdomyoma	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
	Μ	0 (0%)	0 (0%)	0 (0%)	1 (50%)	1 (50%)	0 (0%)	0 (0%)	2 (100%)
Lymphangioma	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	1 (100%)
	Μ	0 (0%)	0 (0%)	1(100%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)
Myofibroma	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
	Μ	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	0 (0%)	1 (100%)
Hamartoma	F	1 (100%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (%)

Leiomyosarcoma	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	0 (0%)	1 (100%)
-	М	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Kaposi's sarcoma	F	0 (0%)	1 (100%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)
-	М	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Carcinosarcoma	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
	М	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	1 (100%)
Infectious diseases									
Paracoccidioidomycosis	F	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)
-	М	0 (0%)	1 (3,5%)	3 (11%)	6 (21%)	12 (43%)	5 (18%)	1 (3,5%)	28 (100%)
Candidiasis	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	2 (100%)	0 (0%)	2 (100%)
	М	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Dermatological diseases									
Lichen planus	F	0 (0%)	1 (8%)	1 (8%)	2 (17%)	2 (17%)	6 (50%)	0 (0%)	12 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	1 (25%)	2 (50%)	1 (25%)	0 (0%)	4 (100%)
Pemphigus	F	2 (25%)	0 (0%)	0 (0%)	3 (38%)	2 (25%)	0 (0%)	1 (12%)	8 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (50%)	1 (50%)	0 (0%)	2 (100%)
Pemphigoid	F	0 (0%)	0 (0%)	0 (0%)	1 (50%)	0 (0%)	1 (50%)	0 (0%)	2 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Physical and chemical injuries									
Amalgam tattoo	F	3 (3%)	5 (6%)	12(14%)	14 (16%)	33 (39%)	15 (17%)	4 (5%)	86 (100%)
	Μ	0 (0%)	0~(0%)	5 (17%)	6 (22%)	10 (34%)	5 (17%)	3 (10%)	29 (100%)
Ulcers	F	1 (4%)	0 (0%)	2 (9%)	1 (4%)	4 (16%)	15 (59%)	3 (12%)	26 (100%)
	Μ	0 (0%)	2 (10%)	0 (0%)	2 (10%)	6 (28%)	9 (42%)	2 (10%)	21 (100%)
Exogenous pigmentation	F	0 (0%)	0 (0%)	0 (0%)	2 (50%)	0 (0%)	1 (25%)	1 (25%)	4 (100%)
	Μ	0 (0%)	0~(0%)	1 (20%)	1 (20%)	3 (60%)	0 (0%)	0 (0%)	5 (100%)
Developmental disorders									
Lymphoepithelial cyst	F	2 (8%)	2 (8%)	5 (20%)	3 (12%)	9 (36%)	2 (8%)	2 (8%)	25 (100%)
	Μ	0 (0%)	2 (18%)	2 (18%)	2 (18%)	3 (27%)	1 (9,5%)	1 (9,5%)	11 (100%)
Dermoid and teratoid cyst	F	3 (43%)	1 (14%)	1 (14%)	0 (0%)	0 (0%)	0 (0%)	2 (29%)	7 (100%)

	Μ	9 (61%)	2 (13%)	0 (0%)	2 (13%)	2 (13%)	0 (0%)	0 (0%)	15 (100%)
Epidermoid cyst	F	2 (25%)	3 (38%)	0 (0%)	1 (12%)	2 (25%)	0 (0%)	0 (0%)	8 (100%)
	Μ	8 (68%)	0 (0%)	1 (8%)	1 (8%)	1 (8%)	1 (8%)	0 (0%)	12 (100%)
Thrombus	F	1 (13%)	0 (0%)	1 (13%)	0 (0%)	5 (61%)	0 (0%)	1 (13%)	8 (100%)
	Μ	0 (0%)	1 (25%)	0 (0%)	2 (50%)	1 (25%)	0 (0%)	0 (0%)	4 (100%)
Varicose veins	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	2 (40%)	2 (40%)	1 (20%)	5 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	1(33,33)	1(33,33)	1(33,33)	0 (0%)	3 (100%)
Thyroglossal duct dyst	F	1 (100%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)
Hematological diseases									
ALK-positive anaplastic large cell lymphoma	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
• •	Μ	1 (100%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)
MALT Lymphoma	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	1 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)
Small cell B-cell lymphomas	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	1 (100%)
	Μ	1 (100%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)
Large B-cell lymphoma	F	0 (0%)	1 (100%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0(0%)	1 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	1 (100%)
Oral manifestations of systemic diseases									
Amiloidosis	F	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	1 (100%)
	Μ	0 (0%)	0 (0%)	0 (0%)	0 (0%)	0 (0%)	1 (100%)	0 (0%)	1 (100%)

Figure 1. Distribution of sample in the oral pathology diagnosis centres.



3 CONCLUSÃO

Em centros representativos de diagnóstico de patologia orais da América Latina, lesões em assoalho bucal representaram 3.73% de todas as amostras acessadas e carcinoma de células escamosas, rânula e leucoplasia foram os tipos histológicos mais comuns.

REFERÊNCIAS

Abdel Wahed Hussein MR. Non-Hodgkin's lymphoma of the oral cavity and maxillofacial region: a pathologist viewpoint. Expert Rev Hematol. 2018 Sep; 11(9):737-748.

Al-jaber A, Al-Nasser L, El-Metwally A. Epidemiology of oral cancer in Arab countries. Saudi Med J. 2016; 37(3):249–255.

Bagan J, Jiménez y, Murillo J, Bagan L. Oral mucous membrane pemphigoid: A clinical study of 100 low-risk cases. Oral Dis. 2018 Mar; 24(1-2):132-134.

Carvalho MDE V, Iglesias DP, Do Nascimento GJ, Sobral AP. Epidemiological study of 534 biopsies of oral mucosal lesions in elderly Brazilian patients. Gerodontology. 2011 Jun;28(2):111-5.

Carvalho WRS, De Souza LL, Pontes FSC, Uchôa DCC, Corrêa DL, De Cáceres CVBL, et al. A multicenter study of oral sarcoma in Brazil. Oral Dis. 2020 Jan; 26(1):43-52.

Casparis S, Borm JM, Tektas S, Kamara Chev J, Locher MC, Damerau G, et al. Oral lichen planus (OLP), oral lichenoid lesions (OLL), oral dysplasia, and oral cancer: retrospective analysis of clinicopathological data from 2002-2011. Oral Maxillofac Surg. 2015 Jun; 19(2):149-56.

Castellanos JL, Díaz-Guzmán L. Lesions of the oral mucosa: an epidemiological study of 23785 Mexican patients. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 2008 Jan; 105(1):79-85.

Chidzonga MM. Oral malignant neoplasia: a survey of 428 cases in two Zimbabwean hospitals. Oral Oncol. 2006 Feb; 42(2):177-83.

Deng D, Wang Y, LIU W, Qian Y. Oral and maxillofacial non-Hodgkin lymphomas: Case report with review of literature. Medicine (Baltimore). 2017 Sep; 96(35):e7890.

Dutra LM, Silva THM, Falqueto A, Peçanha PM, Souza LRM, Gonçalves SS, et al. Oral paracoccidioidomycosis in a single-center retrospective analysis from a Brazilian southeastern population. J Infect Public Health. 2018 Jul - Aug; 11(4):530-533.

Espinoza I, Rojas R, Aranda W, Gamonal J. Prevalence of oral mucosal lesions in elderly people in Santiago, Chile. J Oral Pathol Med. 2003 Nov; 32(10):571-5.

El-naggar AK, Chan JK, Grandis JR, Takata T, Slootweg PJ. World Health Organization classification of head and neck tumours. Lyon, France: IARC Press; 2017.

Girardi FM, Scroferneker ML, Gava V, Pruinelli r. Head and neck manifestations of paracoccidioidomycosis: an epidemiological study of 36 cases in Brazil. Mycopathologia. 2012 Mar; 173(2-3):139-44.

Gondak RO, Da Silva-Jorge R, Jorge J, Lopes MA, Vargas PA. Oral pigmented lesions: Clinicopathologic features and review of the literature. Med Oral Patol Oral Cir Bucal. 2012 Nov 1;17(6): e919-24.

Gouvêa AF, Ribeiro AC, León JE, Carlos R, De Almeida OP, Lopes MA. Head and neck amyloidosis: clinicopathological features and immunohistochemical analysis of 14 cases. J Oral Pathol Med. 2012 Feb; 41(2):178-85.

Dovigi EA, Kwok EY, Eversole LR, Dovigi AJ. A retrospective study of 51,781 adult oral and maxillofacial biopsies. J Am Dent Assoc. 2016 Mar; 147(3):170-6.

Jéhannin-Ligier K, Dejardin O, Lapôtre-Ledoux B, Bara S, Coureau G, Grosclaude P, et al. Oral cancer characteristics in France: Descriptive epidemiology for early detection. J Stomatol Oral Maxillofac Surg. 2017 Apr; 118(2):84-89.

Kauzman A, Pavone M, Blanas N, Bradley G. Pigmented lesions of the oral cavity: review, differential diagnosis, and case presentations. J Can Dent Assoc. 2004 Nov; 70(10):682-3.

La'porte SJ, Juttla JK, Lingam RK. Imaging the floor of the mouth and the sublingual space. Radiographics. 2011 Sep-Oct; 31(5):1215-30.

Li LJ, Li Y, Wen YM, Liu H, Zhao HW. Clinical analysis of salivary gland tumor cases in West China in past 50 years. Oral Oncol. 2008 Feb; 44(2):187-92.

Mujica V, Rivera H, Carrero M. Prevalence of oral soft tissue lesions in an elderly Venezuelan population. Med Oral Patol Oral Cir Bucal. 2008 May 1; 13(5): E270-4.

Neville BW, Damm DD, Allen CM, et al. Oral & Maxillofacial Pathology, ed 4, Philadelphia, WB Saunders, 2016.

Neville BW, Day TA. Oral cancer and precancerous lesions. CA Cancer J Clin. 2002 Jul-Aug;52(4):195-215.

Nemes JA, Redl P, Boda R, Kiss C, Márton IJ. Oral cancer report from Northeastern Hungary. Pathol Oncol Res. 2008 Mar;14(1):85-92.

Nonaka CF, Henriques AC, De Matos FR, De Souza LB, Pinto LP. Nonodontogenic cysts of the oral and maxillofacial region: demographic profile in a Brazilian population over a 40-year period. Eur Arch Otorhinolaryngol. 2011 Jun; 268(6):917-22.

Oliveira FA, Duarte EC, Taveira CT, et al. Salivary gland tumor: a review of 599 cases in a Brazilian population. Head Neck Pathol. 2009; 3(4):271–275.

Pereira JDOS S, Carvalho MDE V, Henriques AC, DE Queiroz Camara TH, Miguel MC, Freitas RDE A. Epidemiology and correlation of the clinicopathological features in oral epithelial dysplasia: analysis of 173 cases. Ann Diagn Pathol. 2011 Apr;15(2):98-102.

Pontes FS, Carneiro JT JR, Fonseca FP, Da Silva TS, Pontes HA, Pinto DDOS S JR. Squamous cell carcinoma of the tongue and floor of the mouth: analysis of survival rate and independent prognostic factors in the Amazon region. J Craniofac Surg. 2011 May; 22(3):925-30.

Puricelli E, Barreiro BOB, Quevedo AS, Ponzoni D. Occurrence of dermoid cyst in the floor of the mouth: the importance of differential diagnosis in pediatric patients. J Appl Oral Sci. 2017; 25(3):341–345.

Saggi S, Badran KW, Han AY, Kuan EC, ST John MA. Clinicopathologic Characteristics and Survival Outcomes in Floor of Mouth Squamous Cell Carcinoma: A Population-Based Study. Otolaryngol Head Neck Surg. 2018 Jul; 159(1):51-58.

Schwanke TW, Oomen KP, April MM, WARD RF, MODI VK. Floor of mouth masses in children: proposal of a new algorithm. Int J Pediatr Otorhinolaryngol. 2013 Sep; 77(9):1489-94.

Subhashraj K. Salivary gland tumors: a single institution experience in India. Br J Oral Maxillofac Surg. 2008 Dec; 46(8):635-8.

Sultan AS, Villa A, Saavedra AP, Treister NS, WOO SB. Oral mucous membrane pemphigoid and pemphigus vulgaris-a retrospective two-center cohort study. Oral Dis. 2017 May;23(4):498-504.

Suliman NM, Åstrøm AN, Ali RW, Salman H, Johannessen AC. Clinical and histological characterization of oral pemphigus lesions in patients with skin diseases: a cross sectional study from Sudan. BMC Oral Health. 2013 Nov 21; 13:66.

Tavares TS, Meirelles DP, DE Aguiar MCF, Caldeira PC. Pigmented lesions of the oral mucosa: A cross-sectional study of 458 histopathological specimens. Oral Dis. 2018 Nov;24(8):1484-1491.

Triantafillidou K, Dimitrakopoulos J, Iordanidis F, Gkagkalis A. Extranodal nonhodgkin lymphomas of the oral cavity and maxillofacial region: a clinical study of 58 cases and review of the literature. J Oral Maxillofac Surg. 2012 Dec; 70(12):2776-85.

Trindade AH, Meira HC, Pereira IF, DE Lacerda JCT, De Mesquita RA, Santos VR. Oral paracoccidioidomycosis: Retrospective analysis of 55 Brazilian patients. Mycoses. 2017 Aug;60(8):521-525.

Viggor SF, Frezzini C, Farthing PM, Freeman co, Yeoman CM, Thornhill MH. Amyloidosis: an unusual case of persistent oral ulceration. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 2009 Nov; 108(5): e46-50.

ANEXOS

Anexo 1 - Verificação de originalidade e prevenção de plágio



Anexo 2 - Certificado do Comitê De Ética em Pesquisa



PARECER CONSUBSTANCIADO DO CEP

DADOS DO PROJETO DE PESQUISA

Título da Pesquisa: Epidemiologia das lesões em assoalho de boca: Um estudo retrospectivo. Pesquisador: Anderson Mauricio Paiva e Costa Área Temática: Versão: 2 CAAE: 09429519.0.0000.0017 Instituição Proponente: Hospital Universitário João de Barros Barreto - UFPA Patrocinador Principal: Financiamento Próprio

DADOS DO PARECER

Número do Parecer: 3.381.233

Apresentação do Projeto:

As Lesões de assoalho bucal podem se apresentar como desafio diagnóstico para cirurgiões dentistas, devido há um vasto espectro de patologias envolvidas nesta região. Lesões em assoalho bucal, podem ser classificadas em tumores benignos e malignos, condições inflamatórias, malformações vasculares, anomalias de desenvolvimento. O objetivo deste estudo é realizar a análise das patologias bucais diagnosticadas em assoalho bucal, através da análise dos prontuários dos pacientes com lesão em assoalho de boca diagnosticado por exame histopatológico, do ano 2000 até 2019, do serviço de patologia bucal do Hospital Universitário João de Barros Barreto (HUJBB). Serão avaliados os seguintes dados clínicos dos pacientes: Sexo, idade, cor da pele, localização da lesão, hipótese clínica diagnóstica, diagnóstico histopatológico final. Os dados serão tabelados e feito análises estatísticas descritivas. Trata-se este de um estudo descritivo retrospectivo com abordagem quantitativa, que de acordo com Polit, Beck e Hungler (2004) é definido como a descrição objetiva sistemática do conteúdo analisado; se realiza com toda a informação numérica obtida através da investigação e que posteriormente será apresentada com o auxílio de quadros, tabelas e medidas que irão traduzir as informações que serão classificadas e analisadas por meio de técnicas estatísticas.

Situação do Parecer:

 Endereço:
 RUA DOS MUNDURUCUS 4487

 Bairro:
 GUAMA
 CEP:
 66.073-000

 UF:
 PA
 Município:
 BELEM

 Telefone:
 (91)3201-6754
 Fax:
 (91)3201-6663
 E-mail:
 cephujbb@yahoo.com.br

Pagina 04 de 05

PlataPorma Brasil

UFPA - HOSPITAL UNIVERSITÁRIO JOÃO DE BARROS BARRETO DA

Continuação do Parecer: 3.381.233

Aprovado Necessita Apreciação da CONEP:

Não

BELEM, 10 de Junho de 2019

Assinado por: João Soares Felicio (Coordenador(a))

Anexo3 - Documento de submissão do artigo

Brazilian Oral Research



WHAT IS THE FREQUENCY OF FLOOR OF THE MOUTH LESIONS? A DESCRITIVE STUDY OF 4,287 CASES.

Journal:	Brazilian Oral Research
Manuscript ID	BOR-2020-0007
Manuscript Type:	Original Research Report
Specialties:	Oral Pathology
CategorySelect your categories from the MeSH or DeCS lists.:</a </a 	Epidemiology, floor of the mouth, benign, malignant

SCHOLARONE[™] Manuscripts