Fibrolipoma of the hard palate: an uncommon location. Case report¹

Fibrolipoma del paladar duro: una ubicación poco frecuente. Reporte de caso¹

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Abstract

Fibrolipoma, a benign soft tissue adipose tumor, is a histological variant of lipoma. Clinically, it presents as a painless slow-growing mass, indistinguishable from other benign soft tissue tumors. In the oral cavity, it is mainly encountered in the buccal mucosa. Involvement of the palate is very rare; it accounts for around 7-14% of all oral fibrolipomas. In this article, we describe a case of fibrolipoma in the hard palate of a 70-year-old female, who presented with an asymptomatic pedunculated mass, characterized by a normal-colored and smooth surface, which have been present for 20 years. The lesion was excised, and histopathological study revealed a fibrolipoma. To the best of our knowledge, only 17 cases occurring in the palate have been reported in the fibrolipoma, mouth, literature. Since fibrolipoma presents clinical similarities with other benign soft tissue neoplasms, a thorough clinical examination and histopathological analysis are essential for obtaining diagnosis.

Resumen

El fibrolipoma es una neoplasia benigna de tejido adiposo, variante histológica del lipoma. Clínicamente se presenta como una tumoración de crecimiento lento, asintomática, indistinguible de otras neoplasias benignas de tejidos blandos. En cavidad oral se presenta principalmente en la mucosa yugal. El paladar es un sitio infrecuente, comprende del 7 al 14% de todos los fibrolipomas. En este reporte, describimos un caso de fibrolipoma de paladar duro en una mujer de 70 años de edad, que presentó una tumoración pediculada, asintomática, de superficie lisa, del mismo color de la mucosa adyacente, con un tiempo de evolución de 20 años. La lesión fue extirpada, y el estudio histopatológico reveló un fibrolipoma. En nuestro conocimiento, se han reportado en la literatura únicamente 17 casos de fibrolipoma de paladar. Siendo que el fibrolipoma presenta similitud clínica con otras neoplasias benignas de tejidos blandos, un examen clínico detallado, así como el estudio histopatológico son esenciales en la obtención del diagnóstico.

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INTRODUCTION

Lipomas are benign neoplasms of adipose tissue. They are commonly encountered as subcutaneous tumors of the head and neck, shoulders, and back. Clinically, lipomas typically present as painless slow-growing, well-circumscribed yellowish masses, usually soft and mobile.¹ Intraoral lipomas are infrequent, they represent less than 5% of all benign mesenchymal neoplasms of the oral cavity. They are predominantly found in the buccal mucosa, labial

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mucosa, and tongue; and their localization in the hard palate is infrequent. Histologically, the most common subtype is the conventional lipoma, followed by fibrolipoma, other variants such as spindle cell lipoma, angiolipoma, and chondrolipoma that represent less than 3%.²⁴

The prevalence of fibrolipomas reported in recent case series of intraoral lipomas varies from 19.6 to 33%.²⁻⁴ Approximately, 251 cases of intraoral fibrolipomas have been reported, according to a PubMed search performed on case series and case reports, in English language, from 2000 to 2021 and using the keywords "oral fibrolipoma" (Table 1). They occur mainly in females in their seventh decade of life and present as a painless solitary nodule, normal-colored and with a slight prevalence for a sessile base.^{3,4} Buccal mucosa is the most affected site, followed by labial mucosa, tongue, alveolar sulcus, floor of the mouth, retromolar pad, ptherygomandibular raphe, and gingiva as the less common site. Palatal fibrolipomas are uncommon, they represent 7-14% of intraoral fibrolipomas.^{4,5} Microscopically, they are characterized by sheets of mature adipocytes admixed with dense bands of collagenous fibrous tissue.⁴ Their treatment is the surgical excision, and histopathological study is mandatory since it presents clinical similarity with other benign soft tissue reactive and neoplastic lesions. In general, clinical(differential?) diagnoses for intraoral fibrolipomas include mostly fibrous hyperplasia, lipoma/fibrolipoma, and other reactive conditions.⁴

This case report aims to present a pedunculated fibrolipoma of the hard palate, which is considered a very infrequent anatomical site and clinical feature of the oral cavity?

Article type	Author	Number of cases	Gender/ Age	Duration (months)	Clinical presentation	Clinical differential diagnosis
Case series	Said al Naief, 2001 ¹¹	4	NS	NS	NS	NS
Case report	Fonseca, 2009 ¹²	1	M/35	NS	Firm, rubbery, normal-colored sessile nodule	Pleomorphic adenoma, lymphoma, malignant neoplasm of salivary glands
Case series	Manjunatha, 2010 ¹³	1	M/70	6	Painless, well-circumscribed, solitary, sessile, and normal- colored sub-mucosal nodule	Fibroma
Case series	Allon, 2011 ¹⁰	4	NS	NS	NS	NS
Case series	Naruse, 2015⁵	1	M/70	12	Painless, normal-colored	NS
Case series	Linares, 2019 ²	1	NS	NS	Painless nodule	NS
Case report	Hayashi, 2020 ¹⁴	1	F/85	84	Painless, smooth-surfaced, sessile, slightly yellowish mass	Lipoma
Case series	Pires, 2021 ⁴	4	NS	NS	NS	Fibrous hyperplasia, Lipoma/ Fibrolipoma, other reactive conditions
Case report	Present case	1	F/70	240	Painless, well-circumscribed, solitary, pedunculated, and normal-colored mass	Fibrous hyperplasia

Table 1. Cases of fibrolipoma affecting the palate reported in the literature

NS: not specified

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Source: by the authors

CASE REPORT

A 70-year-old female presented to the oral and maxillofacial surgeon complaining of mild pain on mastication. She mentioned that for 20 years she has had a swelling in her palate, which never caused pain. Intraoral examination revealed a well-circumscribed, pedunculated, elongated, and lobulated mass, arising from the anterior and lateral part of the palatal mucosa, particularly behind the upper left lateral incisor. The lesion was characterized by a pink and smooth non-ulcerated surface, measuring approximately 20 x 15 mm (Figure 1). During palpation, a fibroelastic consistency was detected. Her oral status was poor and multiple remaining roots were identified.

Based on the clinical features, fibrous hyperplasia was the main diagnostic hypothesis. However, the fact of being a long-standing lesion made it difficult to associate any source of trauma. Therefore, an excisional biopsy under local anesthesia was performed after obtaining the signed informed consent. The specimen was sent for histopathological analysis. Microscopically, it consisted of mature adipose tissue lobules, intersected by dense collagenous fibrous tissue bands covered by a squamous stratified parakeratinized epithelium. A fibrous capsule was seen surrounding the fibroadipose proliferation (Figure 2). The final diagnosis was fibrolipoma. The postoperative course was uneventful, and no recurrence was identified after 7 months of follow-up.



Figure 1. Clinical features: a normal-colored, elongated, and pedunculated swelling, with a smooth pink surface originating from the anterior and lateral hard palate mucosa

Source: by the authors

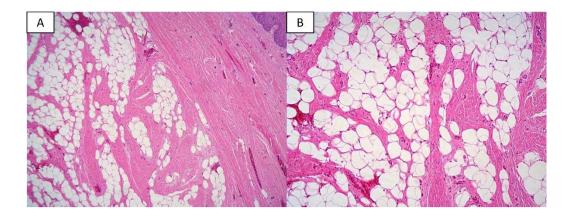


Figure 2. Histopathological features. A. Low-power microphotograph showing the fibrous capsule surrounding the proliferation of mature adipocytes and intersecting collagen bands (H&E, 5x). B. Detail of the adipocytes in a background of dense fibrous tissue (H&E, 10x)

Source: by the authors

DISCUSSION

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The hard palate is an infrequent site for adipose neoplasms as there is scarce adipose tissue in this area. In contrast, it is a common site for minor salivary gland neoplasms, harboring up to two-thirds of these cases. Most benign salivary gland tumors present with similar clinical features to other mesenchymal lesions, i.e. asymptomatic nodules with a smooth and normal-colored surface. However, and different from the present case, they are commonly sessile and found on the posterolateral aspect of the palate.⁶

Fibrous hyperplasia is the most common among the mesenchymal reactive lesions. It usually presents as a sessile nodule, and rarely exceeds 2 cm.⁷ Unlike this case, very often a source of chronic trauma is identified or related by the patients. Despite its high frequency, reactive hyperplastic lesions are uncommon in the palate, except for those associated with old or ill-fitting dentures.⁸

Regarding mesenchymal neoplasms, the palate is a frequent site for neural tumors. The most common in the oral cavity are neurofibroma and granular cell tumor, occurring in the palate and tongue, respectively. The hard palate is the most common intraoral site for palisaded encapsulated neuroma; however, it is usually a small submucosal nodule that rarely exceeds 1.0 cm.⁹

Classically, adipose lesions are associated with a yellowish discoloration; however, this color is only observed in 16.3% of intraoral lipomas. Most of them are covered by normal-colored mucosa as reported in the present case, and in these cases usually, lipoma is not considered a clinical differential diagnosis.³

Different from lipoma which has a male preponderance, fibrolipoma is more common in females, and regarding age, the latter is diagnosed in patients one decade older than simple lipoma, most of them in the seventh decade, as is our patient.¹⁰

Independent of histological variant, most of the intraoral lipomas are sessile based,⁴ particularly in fibrolipomas of the palate, when data were available, also most had a sessile base (Table 1). Interestingly, the actual case had a pedunculated base.

Due to their low frequency, there is few information about the clinical features and implications of the histological variants of intraoral lipoma. For this reason, it becomes important to properly diagnose and clinically characterize them as well as to differentiate them from other entities with similar characteristics.

CONCLUSION

Fibrolipoma is a benign lesion that mainly affects the buccal mucosa, and it is characterized as a painless, well-circumscribed sessile nodule with a normal-colored smooth surface. These clinical features are non-specific and could correspond to other entities, both reactive and neoplastic. The present case occurred in an unusual location for fibrolipoma, but common for salivary gland tumors, so histopathological examination is essential in order to achieve an accurate diagnosis.

CONFLICTS OF INTEREST

The authors declare that they have no conflict of interest.

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