INTRODUCTION

The peak incidence of lipomas is in the fifth or sixth decades of life while it is uncommon in children. Mostly lipomas develop in the subcutaneous tissues but deeper tissues may be involved as well; the oral cavity is not commonly affected.1 The overall incidence in the oral cavity is thought to be between 1% and 4.4% of all benign intraoral lesions.2,3 Oral lipomas can occur in various anatomic sites including the major salivary glands, buccal mucosa, lip, tongue, palate, vestibule, and floor of mouth.4,5 Although benign in nature, their continuous growth may cause interference with speech and mastication due to tumors’ dimension. Some studies showed a female preponderance while others did not found gender preference.4,6

CASE REPORT

A 60 years old patient came to OPD with a cystic swelling in the floor of mouth approximately 5 cm in size. (Fig.1) A longitudinal incision was given over the tumor. Blunt dissection was used through out and the lesion literally popped out from the surrounding.

DISCUSSION

Mostly lipomas develop in the subcutaneous tissues but deeper tissues may be involved as well; the oral cavity is not commonly affected. The overall incidence in the oral cavity is thought to be between 1% and 4.4% of all benign intraoral lesions. The present report shows a 60 years old male who presented with large intraoral mass on the floor of the tongue. Excision biopsy was performed and the histological report proved to be a lipoma.

CASE REPORT

LIPOMA AT THE FLOOR OF MOUTH

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ABSTRACT

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evaluation of superficial structures, like oral and maxillofacial region when the mass is difficult to identify on ultrasonogram, CT or MRI is necessary. Lipoma has a characteristic radiographic appearance. On CT scan it shows a high density from 83 to 143 Haunsfield units with well or poorly defined margins depending on the capsule. Despite the close histological similarity to normal adipose tissue, lipomas usually have chromosomal aberrations such as translocations involving 12q13-15, locus interstitial deletions of 13q, and rearrangements involving 8q11-13 locus. The clinical differential diagnosis includes ranula, dermoid cyst, thyroglossal duct cyst, ectopic thyroid tissue, pleomorphic adenoma and mucoepidermoid carcinoma, angiolipoma, fibrolipoma and malignant lymphoma. A histopathologic differential diagnosis appropriate to the oral cavity would include fibrosarcoma if spindle cells are not localized and numerous. Other lesions should be also considered: they include schwannoma, myxoid-neurofibroma, leiomyoma, nodular fasciitis, myxolipoma, fibrolipoma, malignant fibrous histiocytoma, myxoidliposarcoma, and myxoid solitary fibrous tumor.

The prognosis of this tumor is always good. In adults, the recurrence is rare after complete resection; nonetheless, Cao reported recurrence in patients under 18 years age and development of liposarcoma after several recurrences. Long-term follow-up is necessary in patients under 18 years old. Complete resection should be emphasized during the first surgical operation, which is the key factor in order to avoid recurrence. Well encapsulated lipomas, as the present case, easily shell out with no possibility of recurrence or damage to the surrounding structures.

REFERENCES


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