Lipomatous Pleomorphic Adenoma: A Rare Case Report

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Abstract
Pleomorphic adenoma (PA) is a benign salivary gland tumour characterized by cellular and architectural pleomorphism presented as histological heterogeneity with chondroid, myxoid or osseous components. Lipomatous differentiation is a rarely observed in PA. Here we are presenting a case report of PA with striking lipomatous differentiation, located in upper lip. Recognition of such rare subtype of PA is important for clinical management. As it is located in upper lip should be considered for the differential diagnosis of Adnexal tumour.

Keywords: PA, LPA, Adnexal tumour, Salivary gland tumour.

Introduction
Salivary gland tumour constitute an influential area in the field of the pathology. It is important to recognize that neoplasms may arise not only from major salivary glands but also from any of the numerous, diffuse ,minor salivary glands. Pleomorphic adenoma is most common salivary gland neoplasm and represents 45 -75% of all salivary gland tumours. Annual incidence is approximately 2-3.5 cases per 10,000 population. About 80% of pleomorphic adenomas arise in the parotid gland, 10% in the submandibular gland, and 10% in the minor salivary gland. Among minor salivary glands accounts, hard palate is the most common site approximately 50-60%, followed by upper lip (15-20%) and buccal mucosa (8-10%) 1

Pleomorphic adenoma (PA) is the most common salivary gland neoplasm and yet it has been deemed to be the most difficult SGT in terms of setting the descriptive criteria. Classically, pleomorphic adenoma is characterized by predominance of ductal ,myoepithelial cells along with foci of chondro-myxoid and hyaline stromal development2. It’s pleomorphism is chalked up to the proliferation of basal and myoepithelial cells.
in a capricious stroma matrix of mucoid, myxoid, cartilaginous or hyaline origin. Along with these routine stromal components PA has been known to show some unusual tissue differentiation in the form of osseous, lipomatous and adnexal areas. Rarely, lipomatous change is seen in pleomorphic adenoma of minor salivary gland. A PA is designated as Lipomatous pleomorphic adenoma (LPA) in presence of more than 90% lipomatous stromal component. We report one such case of PA of upper lip which exhibited extensive lipomatous metaplasia which is extremely rare variant of pleomorphic adenoma of minor salivary gland.

Case Report
A 30 year old male patient reported to the Department of oral pathology and microbiology with complain of painless swelling in the upper lip since last 8 months. Past medical history was not conclusive. Clinical examination revealed, small localized swelling in upper lip near the philtrum more towards the skin and intra orally presents as mucosal swelling, round to ovoid in shape measuring approximately 2x1cm in size. (fig 1 & 2). On palpation, the swelling was soft-firm, nodular, well circumscribed, freely movable and non-tender. A provisional clinical diagnosis of minor salivary gland neoplasm was made with a differential diagnosis of benign adnexal tumor. Routine hematological investigations were within normal limits. Surgical excision was done with all aseptic precautions.

On macroscopic examination, a lobulated, ovoid mass measuring about 1.5 × 1 cm in diameter was received. The cut surface of the resected lesion was smooth and yellowish in color. (fig.4) Histopathological finding showed a fibrous connective tissue capsule partially enclosing the lesional tissue and the lesional tissue was composed of basal and myoepithelial cells arranged in sheets, strands and nests. Stromal tissue exhibited abundant adipose tissue. Adipocytes were typically univacuolar and did not show cellular atypia. Focal areas of myxochondroid and hyaline deposition were also noted (fig.5&6). Histopathology was consistent with pleomorphic adenoma. Taking into consideration, the extensive presence of adipose tissue, in pleomorphic adenoma labeled as lipomatous variant of pleomorphic adenoma was made. Excision of the lesion had already been done and the patients was followed up for 18 months without any evidence of recurrence.
Discussion

Pleomorphic adenoma is the most common neoplasm of minor salivary glands, and classically consists of proliferation ductal and myoepithelial cells in chondromyxoid stroma. Other stromal components hyaline or fibrous stroma are found to be rare. But extremely rare variant shows osseous or lipomatous changes in the stroma. Pleomorphic adenomas with an extensive lipomatous stromal component are unusual and very rare histological variant. There are very few reports in the literature, shows pleomorphic adenoma with lipomatous variants. Jamal Musayer et al reported first case of lipomatous pleomorphic adenoma.5

The term lipomatous pleomorphic adenoma was coined by Seifert et al. in 1999. Was described by characterizing the tumour as atypical with more than 90% of its area composed of adipose components.6 Various forms of lipomatous tissue overtones within salivary glands have been defined such as lipoma, interstitial lipomatosis, lipoadenoma, oncocytic lipoadenoma, sialolipoma, and lipomatous atrophy. Granting that foci of adipose tissue are occasionally found within the stroma of pleomorphic adenoma, such an extensive replacement by adipose tissue like the current case is quite an unusual finding.6

Seifert et al showed that the fat-containing salivary gland tumors can be classified into two main groups based on their histopathological composition: monophasic true adipocytic neoplasms (lipoma and its variants) and hybrid lipoepithelial lesions composed of epithelial variants mixed with a variable fatty component. Alternative acknowledged classification divides fat-containing tumors and tumor-like lesions of salivary glands into 4 categories

1. Fat containing epithelial/myoepithelial tumors such as pleomorphic adenoma;
2. Mixed lipoepithelial tumors such as sialolipoma;
3. True adipose tumors like lipoma;
4. Fat-containing tumor-like salivary gland lesions in which diffuse lipomatosis falls.7

Sialolipoma is a benign tumour consisting of adipose tissue admixed with variable amount of adenomatous gland. The tumour present as usually slowly growing mass involving major salivary gland. Sialolipoma composed adipose cells and proliferated glandular tissue. Two possible mechanisms for histogenesis of LPA are suggested-

First doctrine states that the myoepithelial cells shows transformation to adipocytes. Pleomorphic adenoma is also termed as mixed tumor which comprises of both epithelial and mesenchymal tissue components. These mesenchymal tissues may be produced by modified Myoepithelial cells which also have great potential to undergo metaplasia. The second revolves around possible entrapment of fat tissue within the growing tumor.4

The demographic data pertaining to LPA is scanty owing to its rare occurrence. Jamal et al in their case report have recounted mean age for reported cases, and also including our case to be 37.83 years. The gender distribution is 2:1, in favour of female population. Commonly involved site is major salivary gland and in only 10% cases show involvement of minor salivary gland. Hard palate is the most frequently reported site for LPA of minor salivary gland. Our case presents upper lip involvement which is an unusual location for LPA. For clinical differential diagnosis adnexal tumour is considered. Lipoma was also thought as one of the tumour for differential diagnosis. Another salivary gland tumour that is associated

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with adipose tissue is the lipoadenoma or sialolipoma. It is a well circumscribed tumour that may be encapsulated and consist of adipose tissue and glandular elements including ductal, acinar, basal, and Myoepithelial cells, closely resembling normal salivary gland tissue. Although these tumour have both adipose component and epithelial elements, they differ in lipoadenomas lack the abnormal Myoepithelial cells, chondromyxoid stroma, and mesenchymal metaplasia.  

Spindle cell lipoma also considered in the differential diagnosis of a spindle cell lesion showing prominent lipocytic component. Spindle cell lipomas may be distinguished from Myoepithelial lesions by their characteristic ropy collagen and an inflammatory infiltrate including large number of mast cells. Another condition such as lipomatousatrophy, a condition associated with old age considered as one of the differentiating lesion. In this condition no true mass or capsule is present, but normal salivary structure are intermixed with adipose tissue. In present case, upper lip is the site involvement, therefore differentiated from the other lesions such as adenal tumour (chondrosyringoma). Microscopically, presence of metachromatically stained chondromyxoid stroma and epithelial/myoepithelial component indicative of PA is mandatory. Alternative stains such as Romanowsky type and alcohol-fixed dyes, could be applied to aid in differential diagnosis. Painless swelling in upper lip, should be clinically differentiated from the other lesions commonly occurring in that area such as lipoma, fibroma, mucocele, adenal tumours such as chondrosyringoma. Chondrosyringoma of skin is essentially identical to some of mixed tumour. Chondrosyringoma are composed of both epithelial and mesenchymal components with sweat gland elements set in a cartiligenous stroma. Literature review suggests that five histological criteria for diagnosis of chondromyxoid syringoma 1) nests of cuboidal cells 2)intercommunicating tubuloalveolar structures lined with two or more rows of cuboidal cells 3) ductal structures composed of one or two rows of cuboidal cells 4) occasional keratinous cysts 5) a matrix of varying composition.chondroidsyringomas may have all five characteristics or manifest only some. Chondroidsyringoma share similarities with pleomorphic adenomas, but thought to be arise from sweat glands. 

Conclusion

Pathologist should be aware of this rare variant of pleomorphic adenoma. As the site of the lesion in the present case is upper lip near philtrum, may mimick for the adenal tumours. Treatment modality is being same for the all variants of pleomorphic adenoma. Literature reveals that high recurrence 8-45% in pleomorphic adenoma. But the lipomatous variant does not show any recurrence. This rare subtype of pleomorphic adenoma must be known to pathologist in order to establish differential diagnosis, proper treatment modality related with this rare variant of pleomorphic adenoma.

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