

CASE REPORT

INTRADUCTAL PAPILOMA OF SALIVARY GLAND

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ABSTRACT: A case of intraductal papilloma of a parotid gland is presented because of its rarity. The lesion was clinically diagnosed as retention cyst of salivary gland. The excised specimen was histopathologically diagnosed as intraductal papilloma of parotid gland. According to the microscopic finding papillary growth of ductal epithelium with fibrovascular core projecting in to ductal space was seen.

KEYWORDS: Intraductal papilloma, Salivary gland.

INTRODUCTION: Intraductal papilloma of parotid gland is a rare salivary gland tumor^{1,2} and only a few cases have been reported in literature.

CASE REPORT: The patient developed a peanut size swelling just below his left ear lobe which progressively increased in size and caused cosmetic discomfort to the patient which brought him to the surgeon.

On examination a 3x3cms firm to cystic swelling was seen just below the earlobe. Margins were well defined. Swelling was freely mobile and non-tender.

RADIOLOGIC FINDINGS: Ultrasound and CT scan was done-both pointed towards a benign cystic lesion. Left superficial parotidectomy was done. Specimen was sent for histopathological examination which was diagnosed as intraductal papilloma of parotid gland.

FNAC FINDINGS: Smears studied revealed cytological finding highly suggestive of benign cystic papillary lesion of the parotid gland.

MACROSCOPIC FINDINGS: A grey black, grey brown specimen measuring 2.5x1.5cms was sent. On cut section a cystic space measuring 1.5x1cms filled with serous fluid was seen.

MICROSCOPIC FINDINGS: Sections studied revealed normal salivary gland tissue along with a cyst lined by benign low cuboidal to flattened cells. There were papillary structures composed of tubules of varying size lined by two layers of benign cuboidal to low columnar cells; some of the tubular structures revealed proteinaceous material. They were separated by thin fibrous tissue. Areas of hyalinization were also present. The cyst wall contained scattered inflammatory cells comprising of lymphocytes.

Features were consistent with intraductal papilloma of salivary gland

DISCUSSION: Intraductal papilloma of parotid gland is a rare salivary gland tumor most commonly seen involving excretory ducts of intraoral minor salivary gland, buccal mucosa and palate^{3,4}. Only few

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cases have been reported to affect the major salivary gland. The tumors form palpable nodules without any special features and the diagnosis is only made by Histopathological examinations.

Cytological findings are orderly papillary clusters of benign epithelial cells with oncocyctic features some showing sebaceous differentiation.^{5,6}

Clinically patients present with a painless oral or submucosal mass. Majority of patients are in their sixth or seventh decades. Men and women were equally affected. Grossly uniloculated cystic lesion can be seen. Microscopy will reveal cystically dilated duct with papillary epithelial proliferation composed of branching or anastomosing proliferation of single or double layered tall columnar cells with fibrovascular core.^{7,8,9} The cyst is in continuity with the salivary gland excretory duct. Tumor does not proliferate beyond the cyst wall.¹⁰ The lesion is surrounded by a thick fibrous capsule infiltrated by chronic inflammatory cells.

DIFFERENTIAL DIAGNOSES:

1. PAPILLARY CYSTADENOMA: They are multicystic structures.
2. INVERTED DUCTAL PAPILLOMA: Arises at interface of the salivary gland duct with the mucosal surface epithelium whereas intraductal papilloma arises more proximal to a duct.

CONCLUSION: Intraductal papilloma arising in salivary gland may show acinar differentiation, ductal cyst by obliteration and progress to adenocarcinoma. To prevent this complication surgery is the only mode of treatment.

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MICROSCOPIC PICTURES

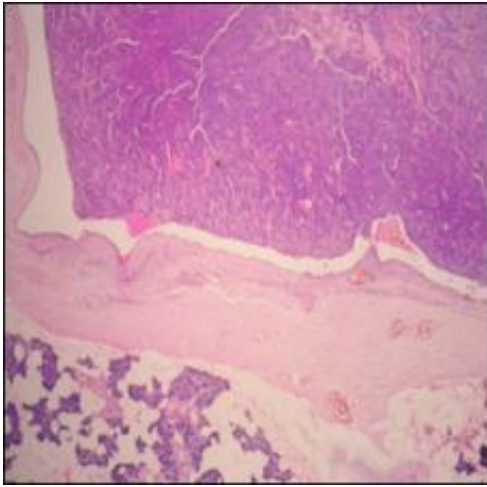


Fig. 1: 10x

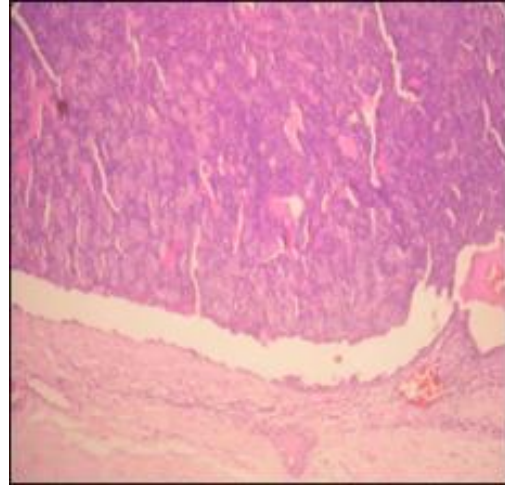


Fig. 2: 20x

Cyst lined by benign low cuboidal flattened cells is seen with papillary structures composed of tubular structures lined by double layer of cuboidal to low columnar cells which are separated by thin fibrovascular core.

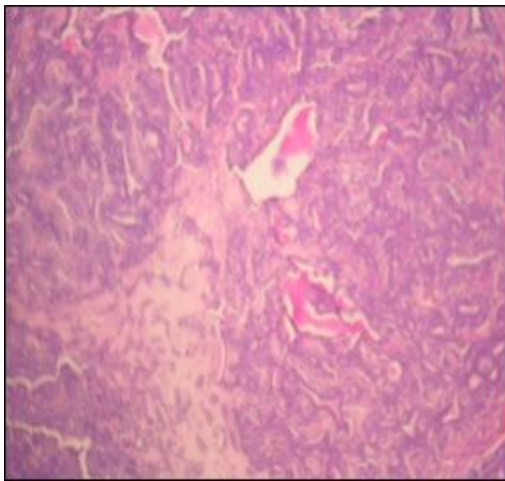


Fig. 3: 10x

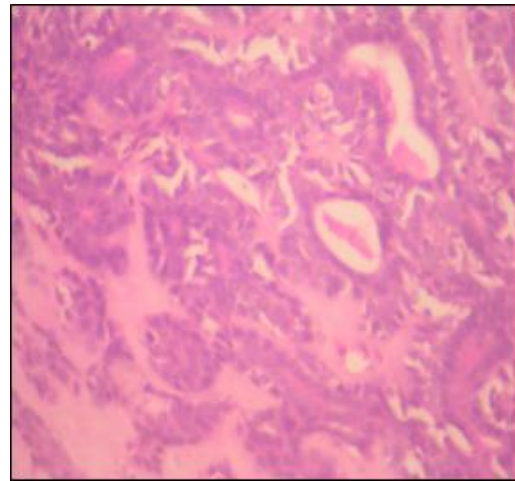


Fig. 4: 20x

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Areas of hyalinization

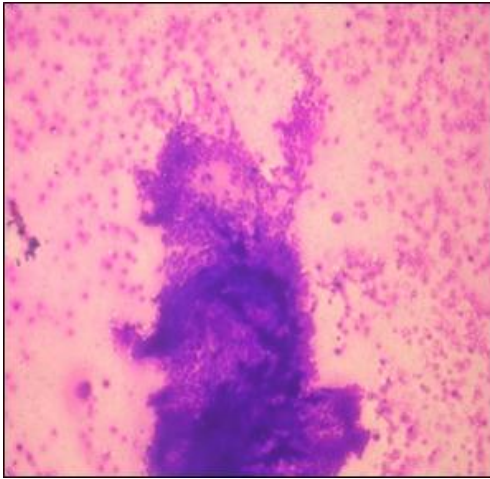


Fig. 5: 10x

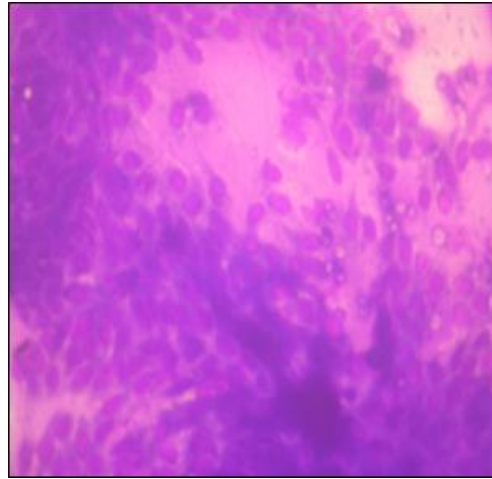


Fig. 6: 40x

Clusters of cells of ductal epithelial origin with a vague papillary pattern with a fibrovascular core are well differentiated with fibrous stroma.

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