www.jmscr.igmpublication.org Index Copernicus Value: 79.54 ISSN (e)-2347-176x ISSN (p) 2455-0450 crossref DOI: https://dx.doi.org/10.18535/jmscr/v7i6.150



Journal Of Medical Science And Clinical Research An Official Publication Of IGM Publication

Case Report

Nodular Hidradenoma- A Case Report

Authors

Parveen Rana Kundu¹, Anjali Sindhu², ^{*}Amrita Kulhria³, Swaran Kaur⁴, Kulwant Singh⁵

^{1,5}Associate Professor, Department of Pathology, B.P.S. Government Medical College For Women, Khanpur Kalan, Sonipat Haryana

²Junior Resident, Department of Pathology, B.P.S. Government Medical College For Women, Khanpur Kalan, Sonipat Haryana.

³Senior Resident, Department of Pathology, B.P.S. Government Medical College For Women, Khanpur Kalan, Sonipat Haryana

⁴Professor & Head, Department of Pathology, B.P.S. Government Medical College For Women, Khanpur Kalan, Sonipat Haryana

^{*}Corresponding Author

Amrita Kulhria

Addresses: Department of Pathology, B.P.S. Government Medical College For Women, Khanpur Kalan, Sonipat Haryana. 131305, India

Abstract

Introduction: Nodular Hidradenoma is rare, benign adnexal tumor arising from the eccrine sweat glands. It presents usually as a slow growing, nodular, solid or cystic cutaneous mass with serous discharge or ulceration.

Case Report: We report case of a 78-year-old lady who presented with a slowly progressive nodular mass over left cheek. Histopathological examination of excision biopsy show histomorphological features characteristic of Nodular Hidradenoma.

Conclusion: Nodular hidradenoma is a rare benign tumor. The clinician and the pathologist should have a strong index of suspicion to detect the lesion as malignant transformation can be there in these lesions. **Keywords**: Adnexal, Nodular, Eccrine.

Introduction

Hidradenoma is benign cutaneous tumor arising from sweat gland. Other names of this tumour are nodular hidradenoma, eccrine acrospiroma and solid cystic hidradenoma.^[1] The age of presentation is in the elderly population, with the peak in the fifth to sixth decade of life. It presents clinically as a firm, solitary intradermal nodule.^[2,3] The lesions are commonly seen on head, neck and limbs and are common in women than in men.^[4,5] The most common histological picture reveals both solid and cystic components with characteristic clear cells and cells with eosinophilic cytoplasm.

Case Report

A 78-year-old female presented in the skin OPD with a 3 years history of an asymptomatic

JMSCR Vol||07||Issue||06||Page 893-895||June

swelling over the left cheek. The swelling was insidious in onset and gradually progressive in nodular size. The swelling measured approximately $2cm \times 2cm$ in size. On local examination, swelling was nontender and mobile. The overlying skin was ulcerated at one point with evidence of serous discharge. No associated cervical lymphadenopathy was noted. The general condition of the patient was normal. Clinical diagnosis of pyogenic granuloma was made. Local excision of the lesion was performed. The excised was sent for histopathological specimen examination. Grossly, specimen comprised of a single grey brown soft tissue piece measuring 1x0.8x0.5cm. Histopathology revealed a well circumscribed neoplasm in the dermis, extending up to subcutaneous tissue. It showed both solid and cystic component (Figure 1). The cystic eosinophilic spaces contained homogenous material. Solid component consisted of nests of two cell types, one with polyhedral shape, a nucleus and slightly eosinophilic rounded cytoplasm and another cell type with round cells containing clear cytoplasm (Figure 2). Focal ulceration of overlying epithelium was noted. No atypical mitosis or necrosis seen. The findings were consistent with the diagnosis of nodular hidradenoma.



Figure 2

Discussion

Benign Nodular Hidradenoma is known by many names such as solid cystic hidradenoma, clear cell myoepithelioma, eccrine sweat gland adenoma, large cell hidradenoma and eccrine acrospiroma. It more often effects women and is commonly seen on scalp, face, anterior trunk and extremities. This tumour can occur at any age, but is commonly seen in the elderly. The first case of nodular hiradenoma was described in the literature by Delacretaz and Leresche.^[6] The origin of the lesion is reported as eccrine based on the number of mitochondria, glycogen granules, and enzyme histochemistry.^[7] Some mention it as of the apocrine origin. Histopathology shows both solid and cystic components. The solid portion contains two types of cells: Polyhedral cells with eosinophilic cytoplasm and glycogen containing clear cells with clear cytoplasm and Cystic areas probably result from degeneration of tumor cells. Clear cell hidradenoma shows positive staining with PAS, cytokeratin, epithelial membrane antigen, S-100 protein, smooth muscle actin, vimentinand p63.^[8,9]

The exact frequency of atypical clear cell hidradenoma and their risk of transformation into malignant tumors are not known. Malignant hidradenomas usually arise de novo.^[10] Malignant hidradenomas present with infiltrative growth pattern, deep extension, necrosis, nuclear pleomorphism and ≥ 4 mitoses per 10 HPFs. The management of benign, atypical and malignant hidradenomas includes wide local excision with adequate margins to reduce the risk of recurrence.

Conclusion

Nodular hidradenoma is a rare benign tumor of sweat glands. The clinician and the pathologist should have a strong index of suspicion to detect the lesion. Complete surgical excision is of the lesion is done with a close follow-up.

Financial Support and Sponsorship- Nil.

Conflicts of Interest- There are no conflicts of interest.

JMSCR Vol||07||Issue||06||Page 893-895||June

References

- 1. Winkelmann RK, Wolff K. Solid-cystic hidradenoma of the skin. Clinical and histopathologic study. Arch Dermatol 1968; 97: 651-61.
- Elder D, Elentisas R, Ragsdale BD (1997) .Tumors of the epidermal appendages, in David E (ed): Lever's Histopathology of the Skin, Philadelphia: Lippincott Raven Publisheres,47804.
- 3. Wilhelmi BJ, Appelt EA, Phillips LG. A rare case of atypical eccrine acrospiroma of the scalp and a literature review. Ann Plast Surg 1999; 42: 568-69.
- 4. Orsaria M, Mariuzzi L. Recurrent eccrine hidradenoma of the breast in a male patient: Problems in differential diagnosis. Our Dermatol Online 2013; 4: 215-17.
- Faulhaber D, Wörle B, Trautner B, Sander CA. Clear cell hidradenoma in a young girl. J Am Acad Dermatol 2000; 42: 693-95.
- Delacretaz J, Leresche A. Hidradenoma of the clear cells. Rev Med Suisse Romande1958; 78:130-35.
- Yildiz B, Ozdemir F, Cobanoglu U, Kavgaci H, Fidan E et al. Clear cell hidradenoma of the gluteal region: A case report. Acta Dermatovenerol Croat 2009; 17: 144-46.
- Gianotti R, Alessi E. Clear cell hidradenoma associated with the folliculosebaceous-apocrine unit. Histologic study of five cases. Am J Dermatopathol1997; 19: 351-57.
- Volmar KE, Cummings TJ, Wang WH, Creager AJ, Tyler DS et al. Clear cell hidradenoma: A mimic of metastatic clear cell tumors. Arch Pathol Lab Med 2005;129: 113-16
- Thami GP, Kaur S, Mohan H. Atypical clear cell hidradenoma. Indian J Dermatol Venereol Leprol 2003; 69: 43-45.