Giant Epidermoid Cyst: An Uncommon presentation of a Common Entity

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Abstract
Epidermoid cysts are the developmental cysts affecting people of any age and gender. Usually, the cysts are asymptomatic and require no treatment. But secondarily infected cysts cause intense pain and discomfort and warrant complete excision. Giant cysts can rarely lead to malignancies. Histopathology can help to confirm the diagnosis and rule out the malignant transformation.

Keywords: Epidermoid cyst, Giant cyst, Sebaceous Cyst, Developmental cyst, keratin cyst.

Introduction
Epidermoid cyst is a developmental cyst mainly originating in the head and neck region[1]. Also known as epidermal cyst, epidermal inclusion cyst, sebaceous cyst, epithelial cyst, keratin cyst[2]. On histology, the cysts are lined by stratified epithelium and filled with a keratinous material. An epidermal punctum is a hallmark for making the clinical diagnosis. People of any age and gender can be involved, though young males are affected most commonly[3]. Generally, the cysts are asymptomatic and require no treatment, but secondarily infected cysts need complete excision of the cyst wall and its contents to prevent recurrence and scarring. Giant epidermoid cysts more than 5 cm in diameter can cause problems, particularly in the head-and-neck region[4,5]. Malignant transformation of an epidermoid cyst can be a rare possible event. Squamous cell carcinoma, basal cell carcinoma, and Merkel cell carcinoma have been reported to arise in the epidermoid cysts[6,7,8]. We report a case of giant epidermoid cyst treated successfully with surgery.

Case Report
45 years old male presented to the Surgery OPD with a painful swelling over the left shoulder. Swelling was present for the last 20 years, was asymptomatic and slowly progressive. Developed throbbing pain and redness 10 days back. General and systemic examination revealed no abnormality. On local examination, there was single erythematous globular swelling of size around 14 * 10 cm over left shoulder region (Figure 1).
It was soft in consistency, fluctuant with smooth surface, well defined margins, fixed to the skin but not to the underlying structures, tender with discharging sinus over the skin with whitish cheesy material coming through sinus (Figure 2).

There was no regional lymphadenopathy or bruit or pulsations. A preliminary diagnosis of giant epidermoid cyst was proposed. Since the swelling was more than 5 cm, possibility of malignancy was also considered. Ultrasonography of swelling showed heterogenously hypoechoic lesion with hyperechoic areas in it and increased vascularity measuring 11.4 * 9.8 * 10 cm. No mobile echos were seen. Possibility of mesenchymal tumour was kept. Fine needle aspiration cytology showed features of keratinous cyst. Excision of the cyst under local anaesthesia was done and sent for histopathology (Figure 3). Grossly, it was cystic mass of size 15*10 cm in size, globular, greyish white in colour (Figure 4). Histopathology was suggestive of epidermoid cyst with no malignant changes. Post operative course was favourable.

Discussion
Epidermoid cysts are benign, slow growing, congenital or acquired, subcutaneous lesions presenting as nodules or tumours\(^9\). Epidermoid cysts are formed either by the sequestration and implantation of epidermal rest during the embryonal
period or by the occlusion of the pilosebaceous unit or the iatrogenic implantation of epithelium or due to the proliferation of epidermal cells within the dermis. Inflammation is in part mediated by the keratinous material contained in epidermoid cysts. Various causes like infection with Human Papilloma Virus\textsuperscript{10}, crush injuries\textsuperscript{11}, cases after skin transplantation\textsuperscript{12}, cryotherapy for wart\textsuperscript{13}, repeated physical loading\textsuperscript{14} on the area have been implicated as the additional etiological factors. Although seen commonly on the hair bearing areas with head and neck accounting for around 7% cases\textsuperscript{15}, they have been seen on the mucosae and glabrous skin. Various differential diagnosis include lipoma, dermoid cyst, trichelemmal cyst, pilomatricoma. Large size of the cyst in our case led to the possibility of malignant transformation, though rare, with the differential diagnosis of squamous cell carcinoma, basal cell carcinoma, Bowens disease. Histopathological examination, excluded malignancy in our patient. In present case, surgical removal followed by histopathology confirmed the diagnosis, ruled out malignancy and its success was confirmed by no recurrence of the lesion. Thus, the giant lesions warrant the use of histopathology to rule out malignancy.

Acknowledgement: Nil

References

