Hidradenoma Papilliferum of the Caruncle

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Abstract

Hidradenoma papilliferum (HAP) is known since the nineteenth century. The vast majority of cases occur on genital and perianal regions of women. Tumors found in other areas of the body are considered to be “ectopic”.

The lacrimal caruncle is a vestigial skin structure with a diverse histology including hair. To our knowledge, the presence of HAP on the caruncle was referred only once in a series of cases published in a gynecological journal. We report for the first time in the ophthalmic literature a case of a caruncular hidradenoma papilliferum (HAP), an unusual benign tumor of apocrinic differentiation.

Keywords: Caruncle; Apocrine; Hidradenoma

Introduction

The human skin has 3 to 4 million sweat glands distributed over most of the body surface. Most of these glands are eccrine which control body temperature by releasing their hypotonic secretion directly on the skin surface without any cell loss [1]. However, apocrine sweat glands, are larger than eccrine glands and secrete, by focal apical decapitation, a proteinaceous viscous sweat into a hair canal. They are concentrated on the axilla, the areola of nipple and genital and perianal areas [1]. The lacrimal caruncle is a vestigial skin structure with a diverse histology including hair [2]. We report for the first time in the ophthalmic literature a case of Caruncular Hidradenoma Papilliferum (HAP), an unusual benign tumor of apocrinic differentiation.

Case Report

A 69-year-old female was referred to the oculoplastic service for an atypical enlargement of the right caruncle. The patient denied any trauma or pain and stated that she first noticed the abnormal size of the caruncle 2 months prior to presentation. On examination there was a round 8.0 × 4.0 mm, reddish and firm mass on the right caruncle (Figure 1). The remaining ophthalmic evaluation was unremarkable. Complete surgical excision was performed under local anesthesia without complications and the specimen was sent for histopathological analysis.

Microscopic examination revealed a soft tissue lesion without connection to the overlying epithelium. The nodule showed tubules and elongated fronds with an arborizing pattern lined by double-layered epithelium. The inner layer was composed of cuboidal cells and the outer layer was tall and consisted of columnar apocrine cells. Periodic acid–Schiff (PAS) -positive diastase-resistant granules were present in the apices of these cells. The mass showed pseudo capsule and cystic areas with colloidal iron positive material. In focal areas infiltration by lymphocytes and plasma cells was observed. The epithelial lesion was positive with cytokeratin CK7 and Epithelial Membrane Antigen (EMA) stains. The inner myoepithelial layer was positive with smooth muscle actin (SMA) and p63 confirming apocrine origin; however the cytoplasmic gross-cystic disease fluid protein-15 stain (GCDP-15) was negative. The mitotic index using Ki 67 was low (Figures 2a-2f).

Discussion

According to Meeker et al. [3], who reviewed the literature on HAP in 1962, this peculiar adenoma is known since the nineteenth century. The vast majority of cases occur on genital and perianal regions of women. Tumors found in other areas of the body are considered to be “ectopic” [4]. The head and neck region is the most common site of non-genital HAP [4] As the gland of Molls are modified apocrine glands, it is not surprising that some cases of HAP related to the eye were reported on the eyelids [5-7] or eyebrow [8,9]. As the orbit has no sweat glands we believe that the only one true ectopic HAP was the orbital lesion recently reported [10]. Although the presence of sweat gland cysts have been found on the caruncle [11,12], HAP did not appear among the 1136 caruncular lesions described in the major series on pathology of the caruncle published in the English literature [11-16]. To our knowledge, the presence of HAP on the caruncle was referred only once on a series of cases published in a gynecological journal [17]. HAP is a benign tumor and does not recur if completely excised. Clinically it presents as an inconspicuous solitary nodule of varying size. The diagnosis is...
entirely dependent on the microscopic features of the tumor since it is a variant of apocrine adenoma with peculiar morphology. Usually the tumor is partly cystic with papillary and glandular components as described in our typical case. Immunohistochemical staining is helpful in confirming the diagnosis especially in highlighting the myoepithelial cell layer using SMA, p63 and GCDP-15 stains. The presence of this myoepithelial cell layer and the secretory granules are important distinguishing features between apocrine and eccrine lesions [6]. The mitotic index in HAP is variable; however it does not necessarily reflect the aggressive behavior, which is seen in hidradenocarcinoma.

Although the rate of malignant transformation of HAP is considered to be low [6], wide local excision is the treatment modality of choice, especially for lesions located intimately on the medial canthus, a notoriously difficult region to perform surgery.

References


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