Özet

Anahtar Kelimeler
Leiomyoadenomatoid Tümör; Epididim

Leiomyoadenomatoid Tümör / Leiomyoadenomatoid Tumor

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Abstract
Ebstein used leiomyoadenomatoid tumor description of a variant of adenomatoid tumor with a prominent smooth muscle component. Adenomatoid tumor is benign, rare neoplasm that may be found mostly in genital tract of women and men at 3rd - 4th decade. Microscopic examination reveals epitheloid appearance of vacuolated, signet-ring like cells, showing infiltrative distribution in fibrous or smooth muscle stroma in tubular, canalicular pleksiform patterns or lining cystically dilated spaces. Most of the ultrastructural and immunohistochemical studies revealed mesotelial origin of this lesion. The presence of smooth muscle in stroma has been generally reported in adenomatoid tumors originated from uterus. Herein we report a case of an adenomatoid tumor located in epididymis which is poor of smooth muscle that indicates the smooth muscle tissue in the stroma is a neoplastic component of adenomatoid tumor.

Keywords
Leiomyoadenomatoid Tumor; Epididymis
Introduction
Leiomyoadenomatoid tumor was first described by Ebstein, in 1992. The term “leiomyoadenomatoid” is indeed a descriptive name for the lesion, and reflects its histopathological appearance that is composed of prominent stromal smooth muscle proliferation accompanied by glandular structures [1]. Most of the ultrastructural and immunohistochemical studies indicate a mesothelial origin for the lesion. Adenomatoid tumors are benign, solitary lesions that occur mostly in genital tract of women and men at 3rd-4th decade [2]. Clinically, these lesions are usually found incidentally, and differential diagnosis of this rather rare lesion from malignant tumors which are more common in testicular region is crucial.

Case Report
A 76-year old man was admitted to urology outpatient clinic with swelling and pain in his right testis. In radiological evaluation of ultrasound showed 3.7x3.2 cm lobular mass, including echogenic areas on right testis cauda epididymis. The patient had a history of coronary by-pass surgery and was taking antihypertensive medication for 20 years. Laboratory analysis revealed ALK:126 IU/L, LDH: 179 IU/L, HCG: 1.2, PSA: 1.56 ng/ml. The patient underwent to right orchiectomy. Orchiectomy specimen was 94 g, and measured as 10.5x7.7 cm. The well circumscribed solid, firm, grayish white mass located between tunica albuginea and epididymis was 3.5x3x3 cm. Microscopically prominent fascicles of smooth muscle were infiltrated by cuboidal to flattened and vacuolated cells as well as tubular-glandular cystically dilated spaces (Figure 1,2). There was no mitoses, necrosis or pleomorphism. In immunohistochemical analysis the epithelial like cells were strongly positive for cytokeratin and calretinin (Figure 3a, 3b). Fascicles of prominent spindle cells were positive for smooth muscle actin (Figure 4).

The lesions were diagnosed as leiomyoadenomatoid tumor.

Discussion
Adenomatoid tumors are benign lesions located most frequently in paratesticular region, epididymis, spermatic cord and tunica albuginea in males and uterus, fallopian tubes and ovary in females. Besides genital region, pleura, lymph nodes and intestinal mesothelial localizations have been reported in both sexes [2,3,4]. Our case was localized in epididymis. Most of the adenomatoid tumors are well circumscribed, grayish white colored solid lesions, smaller than 3 cm in size, and include no hemorrhage or necrosis [5]. In our case, the lesion was also well circumscribed and 3.5x3x3 cm in dimension.

Histopathologically, leiomyoadenomatoid tumors consist of mostly cuboidal cells with epitheloid appearance, sometimes accompanied by vacuolated, signet-ring like cells, showing infiltrative distribution in fibrous or smooth muscle stroma in tubular, canalicular plexiform patterns or lining cystically dilated spaces. Epithelial cells have small round-oval shaped nuclei, without prominent nucleoli and bright eosinophilic cytoplasm. Since the lesion may present with variable histopathologic patterns, differential diagnosis may be challenging. The presence of glandular pattern may mimic infiltrative epithelial and mesothelial lesions, while small-transparent cells with vacuolated
cytoplasm may resemble liposarcoma, and signet-ring like cells may resemble signet-ring adenocarcinoma. In ovary, Leydig cell adenocarcinoma should also be included in the differential diagnosis because of the presence of cells with large, eosinophilic cytoplasm [1-4].

The cases are generally clinically asymptomatic and usually found in radiological scans incidentally. Radiologic evaluations cannot distinguish other benign lesions like leiomyoma [6]. Most other lesions located in the testis have malignant nature; it is important to distinguish adenomatoid tumor of the testis from malignant lesions with frozen section during surgery while the therapy of adenomatoid tumor is enucleation of the mass only. In cases which proper differential diagnosis could not be made, overtreatment with orchiectomy may be performed. Although Akhtar et al. has reported that inter glandular stroma in adenomatoid tumor may only be consist of smooth muscle tissue, stromas of most of the reported adenomatoid tumors are not rich of smooth muscle tissue [7]. Ebstein et al. has used the term 'leiomyoadenomatoid tumor' for adenomatoid tumors that has stroma with prominent smooth muscle tissue. Cases with leiomyoadenomatoid tumor reported in the English literature are rather few. Kausch et al. has reported a leiomyoadenomatoid tumor localized in epididymis [8]. It is not clear whether the presence of smooth muscle tissue is a component of the lesion or a marker of myometrial adjacency in adenomatoid tumors localized in uterus. On the other hand, prominent smooth muscle proliferations in the stroma of adenomatoid tumors localized in ovarian stroma, testis, and adrenal glands which are poor of smooth muscle tissue encourage the leiomyoadenomatoid tumor definition.

Although almost all of the studies indicate a mesothelial origin, histogenesis of adenomatoid tumors is still not clear. Immunohistochemically, the epithelial cells are positive for vimentin, cytokeratin, calretinin, HMME-1 and negative for EMA and CEA [1-4]. Mesothelial cell origin is comprehensible for the adenomatoid tumors that are located in serosa-related organs. In locations which are not directly related with mesothelial lining, such as subserosal myometrium, adenomatoid tumors may be originated from mesothelial inclusions. Another theory in adenomatoid tumor histogenesis is that the lesion may be developed from mesenchymal tissues. This theory can explain the presence of smooth muscle, fibrous tissues and vascular structures as a component of tumor along with the glandular structures [8]. In conclusion, most of the studies endorse the mesothelial cell origin of adenomatoid tumors. In the literature, the presence of smooth muscle in stroma has been generally reported in adenomatoid tumors originated from uterus. As is seen in our case, prominent smooth muscle tissue in the stroma of adenomatoid tumors located in organs like testis and ovary, that are normally poor of smooth muscle tissue, indicates that the smooth muscle tissue is a neoplastic component of adenomatoid tumor. We want to emphasize that defining this benign lesion, which has no clinical, radiological or prognostic difference from adenomatoid tumors, as "leiomyoadenomatoid tumor", explains morphological features much more clearly.

Competing interests

The authors declare that they have no competing interests.

References


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