Primary Mature Cystic Teratoma Mimicking as an Adrenal Mass in an Adult Male Patient

Yetişkin Erkek Hastada Adrenal Kitleyi Taklit Eden Primer Matür Kistik Teratom

Adrenal Kitleyi Taklit Eden Matür Teratom / Mature Teratoma Mimicking as Adrenal Mass

Özet

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Kistik Teratom; Adrenal Kitle; Erkek Hasta

Abstract
Teratomas are bizarre neoplasms derived from embryonic tissues that are typically found only in the gonadal and sacrococcygeal regions of adults. Primary retroperitoneal teratomas are rare and present challenging management options. We report a case of unilateral primary retroperitoneal mature cystic teratoma mimicking as an adrenal mass in 54-year-old male patient. Adrenal mass complete resection was performed by flank approach using the 11th rib resection. Because of the risk of malignancy, follow-up radiographic studies were performed to ensure the oncologic efficacy of resection. The patient has been free of recurrence for longer than 12 months.

Keywords
Cystic Teratoma; Adrenal Mass; Male Patient
Introduction
Primary Retroperitoneal teratomas are very rare in adults, typically occurring in this location only in infancy and childhood. Teratomas are tumors that are derived from embryonal tissue and composed of somatic cell types from two or more germ layers (ectoderm, mesoderm or endoderm) [1]. A teratoma is considered to be a non-seminomatous germ cell tumor and is typically located in either the sacrococcygeal region or in the gonads. Most teratomas in this region (retroperitoneal) are secondary to germ cell tumors of the testes or ovaries. Specifically, in male patients, retroperitoneal germ cell tumors are more likely to have metastasized from the testes than to present as primary tumors [2]. Computed tomography (CT) scan is very useful in differential diagnosis of this rare tumor. We report a case of unilateral primary mature cystic teratoma of the retroperitoneum mimicking as an adrenal mass in an adult male patient.

Case Report
A 54-year-old previously healthy male patient developed acute, left upper quadrant abdominal and left flank pain. This pain lasted several hours and then resolved spontaneously. He was admitted to the hospital where, on ultrasound and an abdominal and pelvic CT scan, he was found to have an 8x7x6 cm solid and cystic mass in the left (surrenal area) retroperitoneum mass containing bone and multiple different soft tissue densities (Figure 1). This tumor had calcification and was not well enhanced. There was no evidence of distant metastasis. This tumor was also examined by magnetic resonance imaging (MRI). We also diagnosed that the tumor originated in the left adrenal gland, because the normal adrenal gland could not be recognized by CT and MRI. The patient had hypertension for which he was on amlodipine 5 mg daily. However, levels of plasma catecholamines, rennin, aldosterone, adrenocorticotropic hormone and cortisol were within the normal range. Furthermore, some tumor markers, such as serum alpha-fetoprotein, lactate dehydrogenase, CA-125, neuron-specific enolase and CA19-9 were examined because it was possible that the tumor did not originate from the adrenal gland. He had no prior surgery. The patient underwent resection of this lesion through a flank incision by 11th rib resection. The retroperitoneal dissection was tedious and difficult, but the mass was excised in its entirety (Figure 2). The left adrenal gland was normal. The resected retroperitoneal cystic mass measured 8x7x6 cm and weighed 153 g (Figure 2). The final pathological evaluation of the tumor was mature teratoma without malignant components. Macroscopically the specimen was 6x5x4 cm fluctuating mass. On incision cystic material came off and a thin walled cyst wall was left. Cyst wall thickness was approximately 1 mm with only some minor excrescens on it. Histologically the cystic cavity was lined by ciliated columnar cells (Figure 3) and the excrescents were hyalin cartilage islands under this epithelium with some lobules of seromucinous glands around (Figure 3). No immature elements were seen and the diagnosis was a mature cystic teratoma pathologically. These tumors usually represent metastasis from other primary sites, additional imaging with CT of the chest and scrotal ultrasonography were performed. No other primary tumor was identified. Therefore, we diagnosed the mass as a primary retroperitoneal teratoma. The patient, after 12 months of followup, was free of recurrence.

Discussion
Teratomas are rare congenital neoplasms that develop from more than one and usually all three of the primordial germ cells which differentiate to form ectodermal, mesodermal and endodermal tissue elements. During the fourth week of embryologic development, germ cells originating from the yolk sac migrate in the midline of the fetus along the dorsal mesentery from the urogenital ridge to the developing gonads. Some of the cells do not complete the migration and survive in midline locations such as the pineal gland, anterior mediastinum, retroperitoneum and sacrococcygeal area where they differentiate into extragonadal teratomas [2,4]. Teratomas are classified as one of four variants: 1) mature, when they contain adult or differentiated tissue; 2) immature, when they are comprised of predominantly embryonic or undifferentiated tissue; 3) teratoma with malignant transformation; and 4) monodermal, when there is a predominance of tissue arising from one germ cell layer. Mature teratomas occur most often in the ovaries and testes. Extragonadal sites account for 15% of all teratomas, and the retroperitoneum is the least common location [5]. The diagnosis of retroperitoneal teratoma often can be made on the basis of radiologic imaging. Retroperitoneal teratomas can be predominantly cystic or completely solid in appearance.
A CT scan or MRI can identify various components of these tumors, including bone, soft-tissue density structures, adipose tissue, and sebaceous and serous-type fluids. These imaging studies also can display the precise location, morphology, and adjacent structures of the tumor, which provide better preoperative planning and increased likelihood of complete removal of the tumor with less iatrogenic damage [8].

A primary retroperitoneal mature cystic teratoma in a 54-year-old man is a rare phenomenon. Most retroperitoneal teratomas in adults represent metastases from a primary gonadal tumor [3]. Our patient had a retroperitoneal mature cystic teratoma that was not derived from any specific organ. Radiographic evaluation did not confirm the origin or the nature of the retroperitoneal mass.

Germ cell tumors in the retroperitoneum usually occur in pediatric populations [6]. Retroperitoneal mature cystic teratomas are characterized by a bimodal peak in incidence, occurring in the first six months of life and in early adulthood [2]. Primary retroperitoneal teratomas in adults are uncommon, with only 32 cases reported between 1937 and 1987 [2]. In adults, documented cases of retroperitoneal teratomas are often secondary sites of tumor genesis. Primary tumors metastasizing to this region have been identified in the breasts, lungs, and gonads. Specifically, in male patients, retroperitoneal germ cell tumors are more likely metastasized from the testes than to present as primary tumors [2]. Primary retroperitoneal teratomas in adults are usually found in the upper portion of the left kidney [2]. In the case of a retroperitoneal tumor, germ cell tumors should be considered and tumor markers examined before surgery [7].

The malignancy rate of 25.8% in adults is significantly higher than the 6.8% rate documented in children [2]. Regardless of the benign histological nature of mature teratomas, close follow up is recommended because the incidence of malignant transformation is approximately 3 to 6%. The patient, after 12 months of follow up, was free of recurrence.

In conclusion, primary retroperitoneal teratoma is a rare entity in adults. Although usually asymptomatic, large neoplasms can cause abdominal and flank pain. Preoperatively, the diagnosis can be established by its characteristic appearance on computed tomography. Although retroperitoneal teratoma can be radiologically recognized, it is important to note that the masses placed at suprarenal region are likely to be confused with suprarenal masses as in our case. The definitive primary treatment of retroperitoneal teratomas is surgical resection. Preoperative biopsy is not necessary if the neoplasm is thought to be completely removable.

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“This manuscript has been revised for language mistakes by a native speaker and professional medical editor of English (Safak Ugur, PhD English and American Literature- Writing Academic English-Middle East Technical University-Modern Languages dept.) for 17 years’

Competing interests

The authors declare that they have no competing interests.