Extrarenal Retroperitoneal Angiomyolipoma: A Rare Case

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Abstract

Angiomyolipomas are benign tumors consisting of smooth muscles, vessels, and fat tissue. They are generally located on the kidneys; extrarenal angiomyolipoma is rare. Apart from the kidneys, angiomyolipomas are seldom located in the retroperitoneal area. Moreover, it is quite difficult to diagnose angiomyolipomas located in the retroperitoneal area. Due to the low amount of fat, negative density may not be identified by computed tomography (CT). The purpose of this report was to discuss the case of a retroperitoneal angiomyolipoma that clinically resembles a surrenal tumor and cannot be clearly discriminated from surrenal tissue on CT.

Keywords: Angiomyolipoma, density, kidney, retroperitoneum, tomography

Introduction

An angiomyolipoma (AML) consists of mature fat tissue, thick blood vessels, and soft muscle cells (1). AMLs are frequently located on the kidneys. Although most cases are sporadic, 20% are accompanied by tuberous sclerosis (2). Retroperitoneal AML may cause abdominal pain and increased abdominal discomfort. The diagnosis of renal AML is established using computed tomography (CT), showing negative fat density [between -10 and -100 Hounsfield unit (HU)] with an accuracy rate of 86% (3,4). In extrarenal AMLs, negative density may not be revealed due to the low amount of fat tissue, which complicates diagnosis (5). In our study, we discussed the rarely encountered case of a retroperitoneal AML.

Case Presentation

A 45-year-old male patient presented to our clinic with the complaint of pain on the right side of his body. A mass lesion (5x6 cm), which could not be clearly discriminated from the surrenal tissue in the right retroperitoneal area, was identified using ultrasonography. Abdominal CT showed a mass lesion measuring 5x6.5 cm with a density of 40 HU on the right side that could not be clearly discriminated from the surrenal tissue (Figure 1). The patient, who occasionally suffered from blood pressure attacks, was pre-diagnosed with pheochromocytoma or adrenocortical carcinoma. He was prepared for surgery by the endocrine clinic. Informed consent was obtained from the patient.

The mass was reached employing the open retroperitoneal approach. The mass was found to be located between the...
kidney and the surrenal tissue, independent of both organs. The whole mass was resected. Pathological examination of the lesion revealed vascular structures with thick walls containing no elastic lamina, mature adipose tissue foci, and fascicles made of intense spindle cells around these structures. In the immunohistochemical examination, spindle cells were reactively identified through S100 SMA and Desmine HBM45. In the immunohistochemical study, S-100 reactivity was determined in the adipocyte areas, whereas CD34 and CD31 were positively identified in the vascular areas. The Ki-67 proliferative index was found to be low (5-10%) in all three components of the tumoral formation. Based on all the data, the case was interpreted in favor of “retroperitoneal AML” (Figure 2).

Discussion
AML was defined in 1951 by Morgan et al. (6). It is characterized by various amounts of vessels, muscles, and fat tissues. Although it is substantially observed to be sporadic, it is accompanied by tuberous sclerosis in 20% of cases. AML accompanied by tuberous sclerosis is frequent in the third or fourth decade of life and especially among women. These tumors are mostly observed to be asymptomatic, small, multifocal, and bilateral. Sporadic cases are observed as bigger, single, and unilateral masses (7) in patients aged between 40 and 70 years.

Tumors located in the retroperitoneal area are not common. The diagnosis before surgery is established via radiological screening. In the retroperitoneal area, lymph node metastasis may be observed after lymphoma, liposarcoma, leiomyosarcoma, schwannoma, paraganglioma, and testicular tumors (8). Retroperitoneal AML is a rare tumor. In the English literature, only 11 cases of retroperitoneal AML have been reported. Typically, these tumors grow towards the vascular pedicle and may cause abdominal pain and hemorrhagic shock (9). Although the radiological diagnosis of renal AML is straightforward, the diagnosis of extrarenal AML is difficult. Since fat density is low in these tumors, a negative density may not be identified, which complicates the diagnosis (10).

In our case, the density was measured as 40 HU due to the low amount of fat. As it could not be clearly discriminated from the surrenal tissue on CT, the mass lesion was thought to be primary surrenal tumor. However, in surgical exploration, the lesion was seen to be independent of the surrenal tissue. If AML could been diagnosed through radiology before surgery and the size of the tumor had been small, the patient could be managed via active surveillance. However, the patient has been operated on.

Consequently, the radiological diagnosis of renal AML is easy while the diagnosis of extrarenal retroperitoneal AML is more difficult. Since there is a lower fat density in these tumors, they may not be detected due to negative density.

Ethics
Informed Consent: Informed consent was obtained from the patient.
Peer-review: Externally peer-reviewed.

Authorship Contributions

Conflict of Interest: No conflict of interest was declared by the authors.

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References


