Intraglomerular Lesions of Renal Angiomyolipoma in Tuberous Sclerosis Complex: A Case Report

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Abstract
The association between tuberous sclerosis complex (TSC) and renal angiomyolipomas (RAML) is widely recognized, but the evidence of intraglomerular microlesions of angiomyolipoma in TSC is extremely rare. We report a 36-year-old woman with TSC, presenting as gross hematuria and hypovolemic shock. The patient was diagnosed with bilateral RAML and spontaneous rupture of right RAML. She underwent right nephrectomy. Microscopic examination revealed multiple bilateral RAML composed of dysmorphic blood vessels, smooth muscle, epithelioid cells and adipose tissue, as well as intraglomerular microlesions of these components in continuity with the glomerular basement membrane but without adhesion to the Bowman’s capsule. In this report we describe the histopathologic features and immunohistochemical markers of TSC-associated RAML with multiple parenchymatous lesions and isolated micro foci in the glomerular tuft. We also review recent discoveries related to possible pathogenesis of renal and intraglomerular lesions.

Keywords: angiomyolipoma, kidney, intraglomerular, Wunderlich’s syndrome

INTRODUCTION

Tuberous sclerosis complex (TSC) is an autosomal dominant disease with a mean incidence estimated in 1/6000 live births1. Approximately 70–85% of patients with TSC develop renal angiomyolipomas (RAML). The most common complication of RAML is Wunderlich’s syndrome, an urological emergency due to spontaneous, non-traumatic retroperitoneal hemorrhage, which occurs in 10% of the cases and can be a life-threatening condition2. Pathologically, RAML are characterized by variable admixture of adipocytes, smooth muscle cells and abnormal vasculature3. The presence of intraglomerular microlesions is extremely rare and there are a few number of reports in the literature4.

CASE REPORT

A 36-year-old woman with medical history of TSC and intellectual disability was admitted to our hospital with sudden-onset right-sided abdominal pain, gross hematuria and hypotension. Abdominal computed tomography (CT) revealed multiple and bilateral renal masses with heterogeneous fat density, which were consistent with RAML. The right kidney showed a large heterogeneous tumor measuring 12.7x 10.5x 9.5 cm, associated with haematoma, suggestive of ruptured RAML.

A diagnosis of bilateral, multifocal RAML with Wunderlich’s syndrome was made. After the patient was hemodynamically stable, right nephrectomy was performed. No complications were observed and the patient was discharged on the fourth postoperative day in a good condition.

Macroscopic examination revealed an enlarged kidney, measuring 14x 8x 6 cm, with multiple nodules range from 0.5-2 cm in diameter and a solid cystic tumor measuring 9.8x 7.5x 6 cm associated with a large perirenal hematoma.

Histologically, the tumors were composed of thick-walled, hyalinised vessels, many of them without
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internal elastic lamina, bundles of smooth muscle, mature adipose tissue and epithelioid cells of varying proportions. Necrosis and hemorrhage were also observed. Numerous microscopic tumor foci were found throughout the parenchyma, as well as isolated intraglomerular segmental nodules consisting of epithelioid, smooth muscle and adipose cells without attachment to the Bowman’s capsule (Fig. 1).

Neoplastic cells expressed melanocytic markers (HMB-45 and Melan-A), desmin, smooth muscle actin, vimentin and were negative for pan-cytokeratin AE1-AE3.

Figure 1. Intraglomerular microlesions of angiomyolipoma consisting of adipose and epithelial cells. A-C: hematoxylin-eosin; original magnification, 200x (A) and 1000x (B-C). D: periodic acid-Schiff; original magnification, 1000x

Discussion

TSC is an autosomal dominant disease characterized by hamartomatous tumors of the skin, brain, kidney, heart and lung. TSC is caused by mutations in two genes, TSC1 at position 9q34 and TSC2 at 16p13.3, which encode proteins that act as tumor suppressors. These proteins are called hamartin and tuberin, respectively. Kidneys are frequently involved in TSC and renal complications are the most common causes of morbidity and mortality in these patients. Renal disease includes angiomyolipomas, epithelial cysts and renal cell carcinoma. Multiple and bilateral RAML develop in up to 80% of patients with TSC. Although these tumors are usually asymptomatic, lesions larger than 3 cm in diameter are prone to bleeding. The risk of bleeding is proportional to the size of the lesion and up to 10% of TSC patients may experiment spontaneous massive and potentially fatal retroperitoneal hemorrhage or Wunderlich’s syndrome.

This report describes a case of bilateral RAML with multiple nodular lesions either inside or outside the glomeruli. All glomerular lesions were in continuity with the capillary wall. The lack of attachment to the Bowman’s capsule suggests that the origin of these microlesions occurs inside the glomerulus.
Intraglomerular Lesions of Renal Angiomyolipoma in Tuberous Sclerosis Complex: A Case Report

The intraglomerular microlesions in RAML were first reported in 1930 by Feriz et al. They described “epithelial and fatty inclusions” in the glomerular tufts. In 1973, Shinohara et al reported similar lesions, but the first detailed description was given by Nagashima et al in 1988.

The RAML are histologically classified as typical (triphasic) or atypical (monophasic or epithelioid). Most RAML contain variable admixtures of tumor cells that are histologically and molecularly similar to vascular, smooth muscle and fat lineages. Renal AML cells express molecular markers of melanocyte lineage such as HMB-45 and melan-A and muscular markers such as smooth muscle actin and less frequently desmin. It has been proposed that they might originate from undifferentiate cells of the neural crest or of an unidentified AML neoplastic stem cell.

RAML have been considered to be an hamartoma for a long time, but there is strong evidence supporting that they are true neoplasms with a clonal origin. Martignoni et al suggested that there is a group of lesions that originate from a progenitor cell called perivascular epithelioid cell (PEC) which is characterized by an epithelioid appearance, perivascular distribution and intracytoplasmic HMB-45 positivity. They described intraglomerular lesions with features of angiomyolipoma in patients with and without tuberous sclerosis and in the TSC2/PKD1 contiguous gene syndrome, a disease with a deletion disrupting both TSC2 and PKD1 (autosomal dominant polycystic disease gene).

Recently, Gonçalves et al described the first mouse model of RAML and provided evidence that these mesenchymal tumors originate from renal proximal tubule epithelial cells.

CONCLUSION

The intraglomerular microlesions of angiomyolipoma are extremely infrequent. Human RAML express melanocyte and smooth-muscle markers, as we found in our case. However, the accurate cell origin of AML or AML stem cell remain unknown.

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Intraglomerular Lesions of Renal Angiomyolipoma in Tuberous Sclerosis Complex: A Case Report


