

## ADENOMATOID TUMOR OF SUPRA-RENAL GLAND

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### ABSTRACT

Adenomatoid tumors of adrenal gland are rare, asymptomatic neoplasias, with benign behavior, and usually are diagnosed incidentally.

We report a case of a voluminous adenomatoid tumor of left adrenal gland in a 42-year old man who sought evaluation because of renal colic due to left nephrolithiasis. During the investigation, a tumor localized in left adrenal gland was identified by ultrasonography (14.3 x 10.5 x 19.0).

The patient underwent adrenalectomy and pyelolithotomy with histopathological and immunohistochemical diagnosis of adenomatoid tumor of adrenal gland, being the largest one described in the literature to the moment. The patient does not present any signs of recurring lesion after a 3-year follow-up.

**Key words:** adrenal glands; neoplasms; adenomatoid tumors; nephrolithiasis  
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### INTRODUCTION

Adenomatoid tumors of adrenal gland are rare, asymptomatic neoplasias, with benign behavior, and usually are diagnosed incidentally (1). Adenomatoid tumors appear more frequently in the genital system both in males and in females (2). The literature reports only 16 cases of adrenal adenomatoid tumors (3). In this work, we report one case of voluminous adenomatoid tumor affecting the left adrenal gland.

### CASE REPORT

Caucasian, 42-year old man was referred due to presenting, during investigation of renal colic, a mass measuring 14.3 x 10.5 x 19.0 cm localized on left adrenal gland in ultrasonographic imaging. The patient had a previous history of systemic arterial hypertension with moderate tension levels since he was

28 years old, and elimination of urinary stones at the ages 32 and 39. On the physical examination, he presented blood pressure of 150 x 100 mmHg (using captopril) and a mass with firm consistency and indefinite limits on left hypochondrium. Abdominal computerized tomography demonstrate a tumor measuring 10.5 x 9.9 cm localized on left adrenal gland, caudally displacing the ipsilateral kidney, showing peripheral impregnation by the contrast agent, presenting heterogeneous texture, with hypodense areas in its interior, possibly corresponding to necrotic areas, and a stone measuring 3.2 x 2.7 cm in left renal pelvis corresponding to renal lithiasis. Dosing for urinary vanillylmandelic acid, cortisoluria, 17-hydroxyprogesterone, hydro-epiandrosterone sulfate, androstenedione, aldosterone, follicle-stimulating hormone, luteinizing hormone, prolactin, total and free testosterone, thyrotrophin, thyroxine and cortisol (following 1 mg of dexamethasone) were within the normal range.

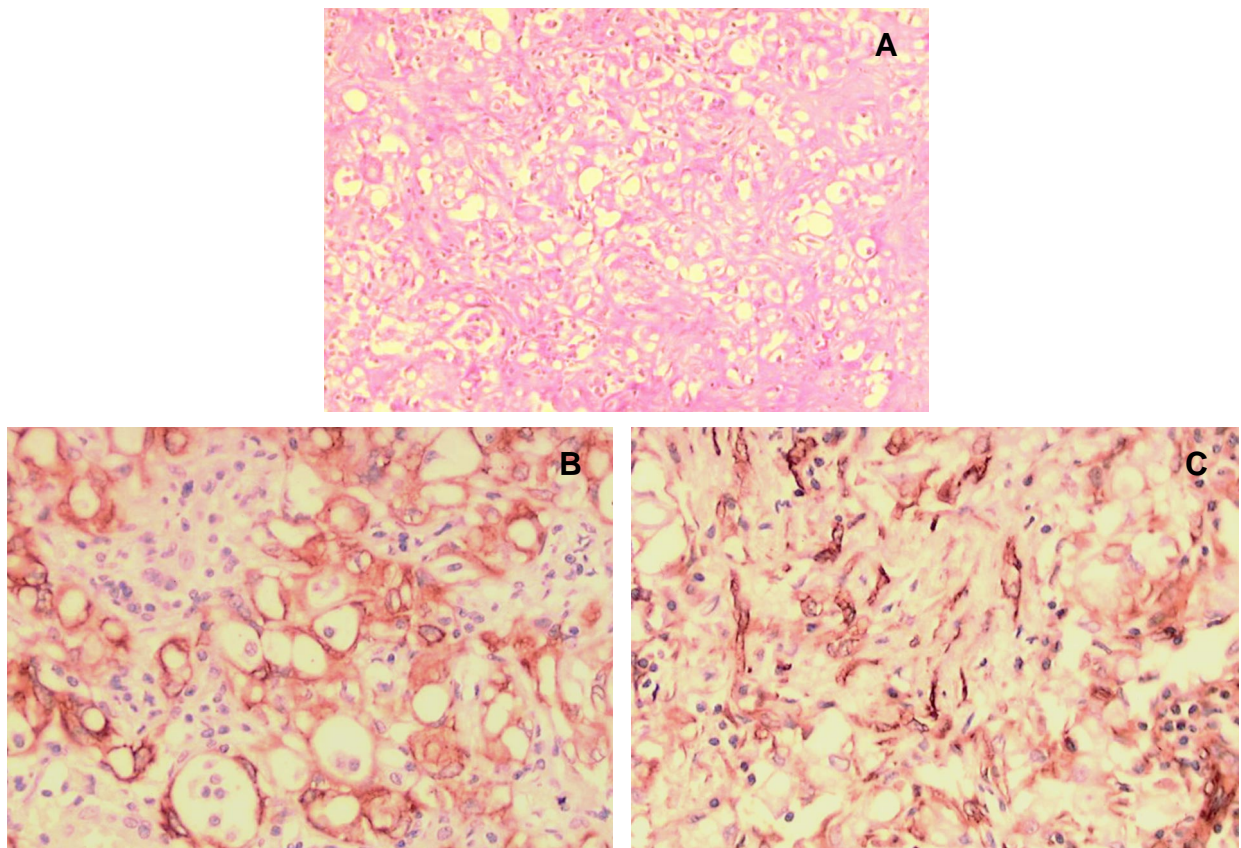
Adrenalectomy and pyelolithotomy were performed by open approach, with no intercurrences trans- or post-operatively. The adrenal gland weighted 535 g and upon sectioning it showed predominance of multicystic, yellowish and opaque tissue, with the tumor being predominantly solid. Histological examination with hematoxylin-eosin demonstrated mesothelial cells similar to epithelial lineage cells arranged as small tubules, cysts or string-shaped, with histological pattern compatible with adenomatoid tumor of adrenal gland (Figure-1). Immunohistochemical study by the immunoperoxidase technique using the markers AE1, AE3, vimentin, CEA and CD31, demonstrate that neoplastic cells were positive to AE1,

AE3 and vimentin (Figure-1), and negative to the other markers, confirming the diagnosis.

The patient has not shown signs of recurrent lesion after a 3-year follow-up.

## COMMENTS

Adenomatoid tumors are benign neoplasias with mesothelial origin, with cases rarely reported in extra-genital sites (2). There are 8 cases describing adenomatoid tumor of adrenal gland, occurring mostly in males, involving more frequently the left adrenal gland (1). With the increasing use of imaging examinations for diagnostic of other pathologies, adrenal



**Figure 1** – Histological and immunohistochemical studies of the adenomatoid tumor of adrenal gland. A) Proliferation of mesothelial cells similar to epithelial lineage cells, arranged as small tubules, cysts, or string-shaped (HE, X80). B) Positivity for cytokeratins in epithelial-like cells. CEA negative. Positive to AE1 and AE3 (Imunolabeling, X200). C) Positivity for vimentin expressed in mesothelial cells (Imunolabeling, X200).

tumors have been more diagnosed, usually in an incidental way. Apparently, there are no specific characteristics that enable us to radiologically distinguish adenomatoid tumors from other adrenal lesions (4). The differential diagnosis of an adrenal mass includes adrenal adenoma or carcinoma, myelolipoma, pheochromocytoma, cysts and metastatic neoplasias (1). There are no reports in the literature of lesion recurrence following complete surgical removal, and this treatment is definitive.

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