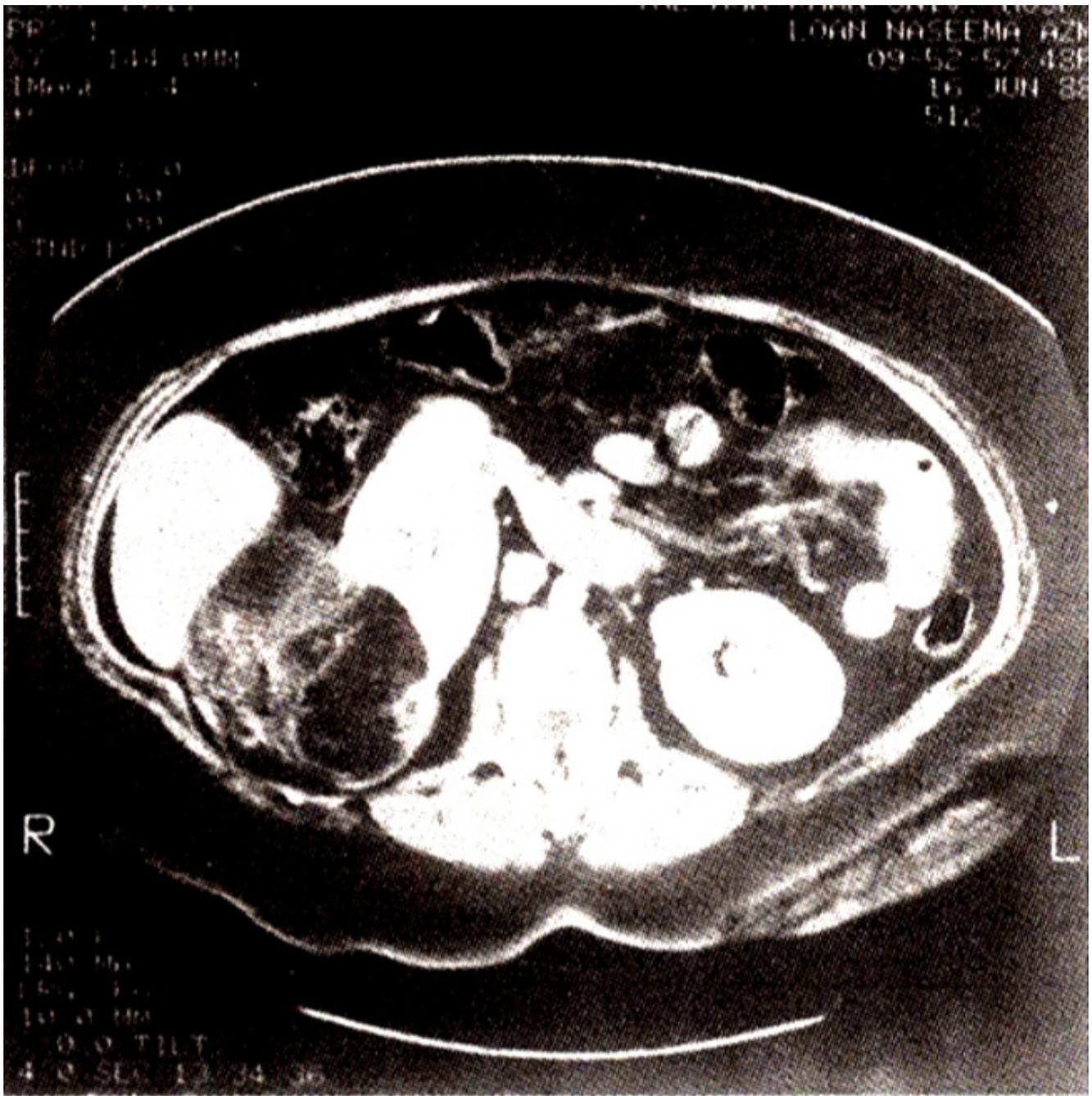


# RENAL ANGIOMYOLIPOMA: AN UNCOMMON TUMOUR

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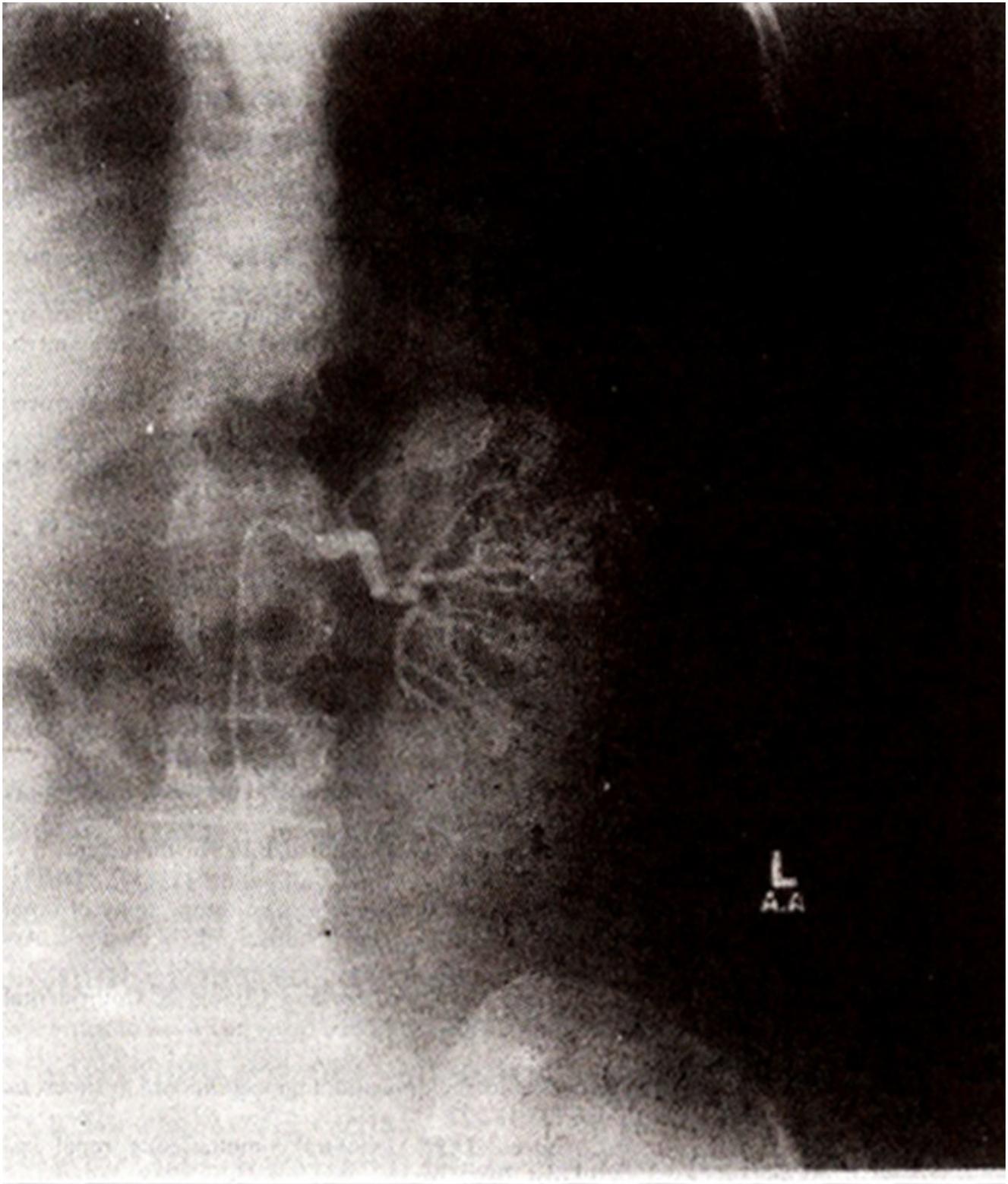
The term angiomyolipoma of the kidney was first introduced by Morgan et al<sup>1</sup> to denote an uncommon type of renal tumour with the histological components of thick-walled vessels, smooth muscle cells extending outwards from the vessels in a peritheliomatous fashion and fat cells of the adult and foetal types. Although the tumour is referred to as a renal hamartoma<sup>2</sup>, a mass containing tissues normally present in the organ but abnormal in their arrangement, in fact, renal angiomyolipoma (HAML) are choriostomas, as the fat is not the normal component of the human kidney<sup>3</sup>. Isolated angiomyolipomas are relatively rare renal tumours. Till 1988 only 250 cases were reported in English literature. The present study adds 3 additional cases and review the outstanding clinical features and advances in the diagnosis and management. Case 1. L.M.N. a 47 year old diabetic, hypertensive female presented in the outpatient clinic for evaluation for right sided flank pain of 5 months duration. There was no history of bowel or urinary symptoms. Examination revealed a tender, firm and smooth mass measuring approximately 20x15 cm occupying the right side of the abdomen. CBC, urine analysis, 24 hour urinary VMA and biochemical profile were normal. The abdominal ultrasonography and intravenous urography revealed a space occupying lesion in right kidney and fullness of the pelvicalyceal system. The CAT scan revealed a 9 cm nonhomogeneous mass with areas of low attenuation values (-65 Hounsfield units) in the right kidney (Figure 1).



**Figure 1. CAT scan showing nonhomogeneous mass of low attenuation value consistent with renal angiomyolipoma.**

In addition a 2.5 cm solid mass arising from the right adrenal gland was noted. There was no evidence of retroperitoneal lymphadenopathy. Renal cell carcinoma could not be excluded on CT scan appearances. At exploratory laparotomy after opening the gerota's fascia a tumour mass was seen which was mostly extrarenal but was growing into the right kidney, leaving only a thin rim of cortex. It was soft and spongy to feel and was not vascular. A separate adrenal mass hard in consistency could also be felt. The tumour mass, right kidney and adrenal gland were excised in toto. Histopathology confirmed angiomyolipoma of kidney and adrenal gland. The postoperative course was uneventful and the patient has remained well and symptom free throughout a two years follow-up. Case 2. S.F.N. a 45 year, hypertensive female presented with left lumbar pain of one year duration. Her symptoms had worsened since the past month and had become associated with generalized weakness, anorexia, nausea

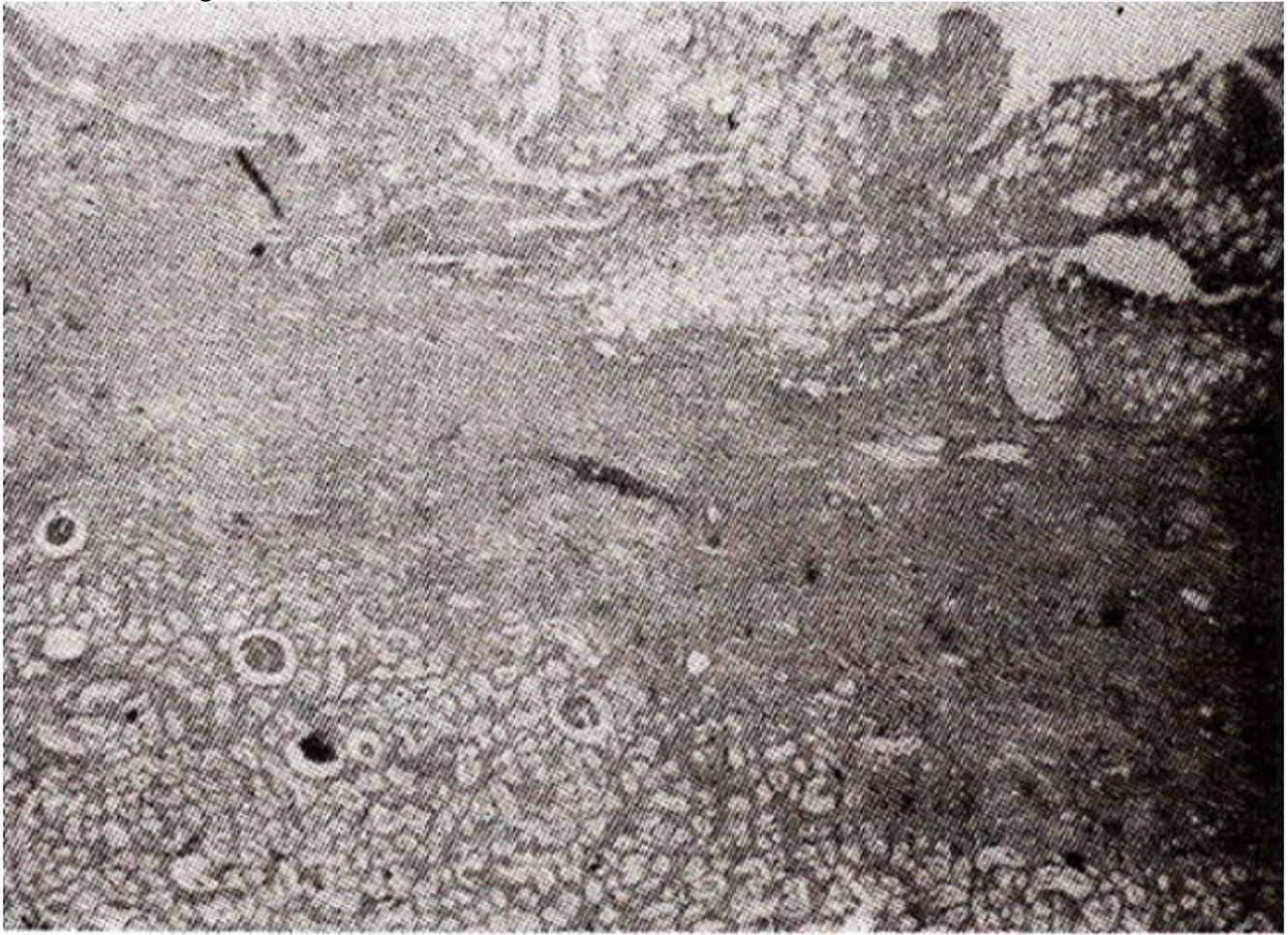
and vomiting. There was no history of fever and urinary symptoms. Examination revealed pallor and a bimanually palpable mass measuring approximately 30x20 cm in left lumbar region. Hemoglobin was 9.2 grn/dl with hematocrit 27.7%. A urinalysis and routine biochemical profile was normal. An intravenous pyelogram revealed an enlarged left kidney with adequate excretion of dye. An ultrasound (US) of the abdomen showed a hyperechoic mass arising from lateral aspect of left kidney. CT scan of abdomen showed a perinephric hematoma and an intrarenal mass measuring 6x6cm, showing decreased attenuation values (-60 hounsfield units) supportive of fat in the kidney. A renal angiography demonstrated a nonvascular mass, multisaculated aneurysmal dilatation of peripheral artery, sunburst appearance of veins and lack of arteriovenous shunting (Figure 2).



**Figure 2. Angiographic features of angiomyolipoma.**

A preoperative diagnosis of RAML was made. Exploratory laparotomy revealed a large perirenal hematoma contained by the gerota's fascia. Evacuation of hematoma revealed a 6x6 cm, well demarcated, yellowish firm mass arising from the lateral aspect of upper pole of left kidney. Hemostasis was secured by multiple suture ligation and the cavity was obliterated with an omental

pedicle. The diagnosis of RAML was confirmed on frozen section and again on final histopathological examination (Figure 3).



**Figure 3. Microscopic features of renal angiomyolipoma showing muscles, abnormal blood vessels and fat cells.**

Postoperative recovery was uneventful. The patient has remained symptom free throughout the 18 months follow-up. Case 3. K.A. 34 year old female who was previously in good health presented to the emergency room with acute flank pain associated with nausea, vomiting and sweating. There were no urinary or bowel symptoms. Examination revealed a tender, firm mass measuring approximately 15x10 cm occupying the right side of abdomen. Hemoglobin was 5.1 gm/dl with hematocrit 16.8%. Urinalysis revealed microscopic hematuria. Biochemical profile was normal. Intravenous pyelogram and ultrasound of abdomen revealed a space occupying lesion in the right kidney. CT scan showed perirenal haematoma, a 4 cm smooth mass in the right kidney and a huge retroperitoneal mass with low attenuation value (-65 negative hounsfield units) consistent with diagnosis of RAM!.. The patient normally a resident of London, returned there where she underwent a right nephrectomy. Histopathology revealed RAML.

## **DISCUSSION**

BAML are uncommon benign lesions that may occur as an isolated phenomenon or as seen in 20-50% of patients as part of the syndrome associated with tuberous sclerosis (Bournvill's disease<sup>4</sup>). About 50-80% of patients with tuberous sclerosis will have angiomyolipoma<sup>5</sup>. Therefore these patients require

periodic screening as these can be occasionally symptomatic<sup>6</sup>. RAML as an isolated phenomenon usually present as a large symptomatic unilateral mass. They occur predominantly in women in fifth and seventh decades<sup>7</sup>. Grossly RAML are greyish-yellow in colour and have propensity for profuse hemorrhage and local aggressive growth<sup>7</sup>. They may occasionally parasitize to external blood vessels and lymph nodes. The actual incidence of nodal involvement is not known. A review of literature reveals only 8 cases<sup>9</sup>. The involvement of regional lymph nodes and blood vessels has been attributed to the multicentric origin of the lesion rather than metastatic behaviour<sup>10-13</sup>. To date there has been no death due to metastatic spread. Microscopically angiomyolipoma consists of sheaths of smooth muscles, abnormal thick wall blood vessels and clusters of adipocytes in varying proportions and arrangements (Figure 3). Renal cell carcinoma may coexist with angiomyolipoma in the same kidney. Twelve such cases until 1989 have been reported in literature<sup>9,14,15</sup>. One case of simultaneous RAML and oncocytoma has also been reported<sup>16</sup>. The clinical presentation varies. Symptoms may be absent if the lesion is small (<4cm). When present, symptoms are usually a result of intrarenal or perirenal hemorrhage. More than 50% of patients with angiomyolipomas larger than 4cm have associated hemorrhage<sup>12</sup>. The most frequent signs and symptoms are acute flank/abdominal pain, palpable mass, hematuria, shock and anaemia. Hypertension and fever has been reported<sup>8,12</sup>. The diagnosis of angiomyolipoma should be borne in mind since 25% of patients with RAML described in the literature presented with severe abdominal pain with or without shock, which necessitated emergency laparotomy<sup>8</sup>. In the past, correct preoperative diagnosis has been extremely difficult with the result that most patients undergo nephrectomy for a mistaken diagnosis of renal cell carcinoma<sup>1,3,12</sup>. In recent years CT scan and ultrasonography have resolved the diagnostic dilemma, making definitive diagnosis of RAML possible in almost all cases<sup>8,13</sup>. Angiomyolipomas are the most echogenic form of renal tumours, due to their high fat content<sup>4</sup>. However this pattern is not always pathognomonic since other echogenic lesions, including renal cell carcinoma incorporating perirenal fat, liposarcoma, lipoma, teratoma, oncocytoma, cavernous hemangioma and renal sinus lipomatosis may also exhibit a similar picture. A CT diagnosis of angiomyolipoma requires identification of fat component in the tumour which usually stands out in the lesion as areas with low attenuation value (negative Hounsfield unit). Lesions smaller than 2.0 cm may be difficult to diagnose owing to partial volume effect. Moreover a hemorrhage within or outside the tumour may increase the attenuation value to above that of fat<sup>17</sup>. A 50-78% diagnostic accuracy of CT scan has been reported<sup>1,3,8,12,13,17</sup>. We were able to make tentative preoperative diagnosis in all of our cases with CT scan but were reluctant to confidently exclude carcinoma except in the case who had angiography (case No.2). Since CT scan cannot consistently diagnose all EAML, a fine needle aspiration biopsy (FNA) has been advocated by Scant and associates<sup>18</sup> to solve the diagnostic dilemma<sup>18</sup>. Selective renal angiography has been suggested both for diagnostic and therapeutic purposes<sup>12</sup>. The angiographic characteristic of an angiomyolipoma are hypovascular mass with circumferential tortuous dilated peripheral vessels having multiple aneurysms, sunburst or whorled appearance of the veins and lack of the arteriovenous shunting which is seen in renal cell carcinomas<sup>19-21</sup>. Selective arterial embolization has been shown to be effective in treating acute hemorrhage alone or in conjunction with surgical intervention<sup>4</sup>. With accurate preoperative diagnosis of the angiomyolipoma, most urologists currently recommend a conservative approach to the management. The nephrectomy rate has declined from 93% in 1976 to 50% in 1984<sup>12</sup>. For small (< 4cm) asymptomatic tumours, surveillance is all that is required. With large (>4cm) and symptomatic angiomyolipomas, renal conservation with tumour excision or partial nephrectomy are preferred if possible<sup>1,3,4,7,8,12</sup>. However it is important that conservation should never compromise completeness. Frozen section may be helpful for the diagnosis of coexistent renal cell carcinoma<sup>12</sup>. In conclusion,

renal angiomyolipoma, although a rare lesion, should be included in the differential diagnosis, when dealing with a patient presenting with severe flank pain and anaemia. Considering the benign nature of disease and improvement in preoperative diagnosis, a kidney conserving operation is desirable.

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