

Images in . . .

Ultrasound and CT imaging assessment of renal angiomyolipoma

Alexandros Kotis,¹ Filippos Lisgos,² Stylianos Karatapanis²¹ CT and MRI Department, General Hospital of Rhodes, Rhodes, Greece² Department of 1st Internal Medicine, General Hospital of Rhodes, Rhodes, Greece

Correspondence to Alexandros Kotis, kotisnet@otenet.gr

Summary

The authors report a case of a 41-year-old woman who was admitted to the emergency department of our hospital because of acute right flank pain. Laboratory investigations and cultures were negative. A transabdominal ultrasonography revealed a large mass of the upper pole of the right kidney as an incidental finding.

CASE PRESENTATION

A 41-year-old woman was admitted to the emergency department of our hospital because of acute right flank pain. Laboratory investigations and cultures were negative. A transabdominal colour doppler ultrasonography scan using head transducer 1.75–4 MHz (Siemens/Acuson; Aspen, Mountain view, California, USA) revealed a large (5×4 cm), highly hyperechoic well-demarcated exophytic mass with shadowing, of the upper pole of the right kidney as an incidental finding (figure 1). Subsequently abdominal CT

spiral scan (10/10 mm sections GE SX Prospeed before and after contrast injection) showed a mass with mixed hyperdense and fatty tissue elements (figures 2,3). Renal excretion and filling of the urinary bladder were both satisfactory. There was no evidence of lymph node enlargement or other space occupying lesion in the abdomen. All the diagnostic imaging findings were consistent with the diagnosis of angiomyolipoma. A follow-up was recommended.

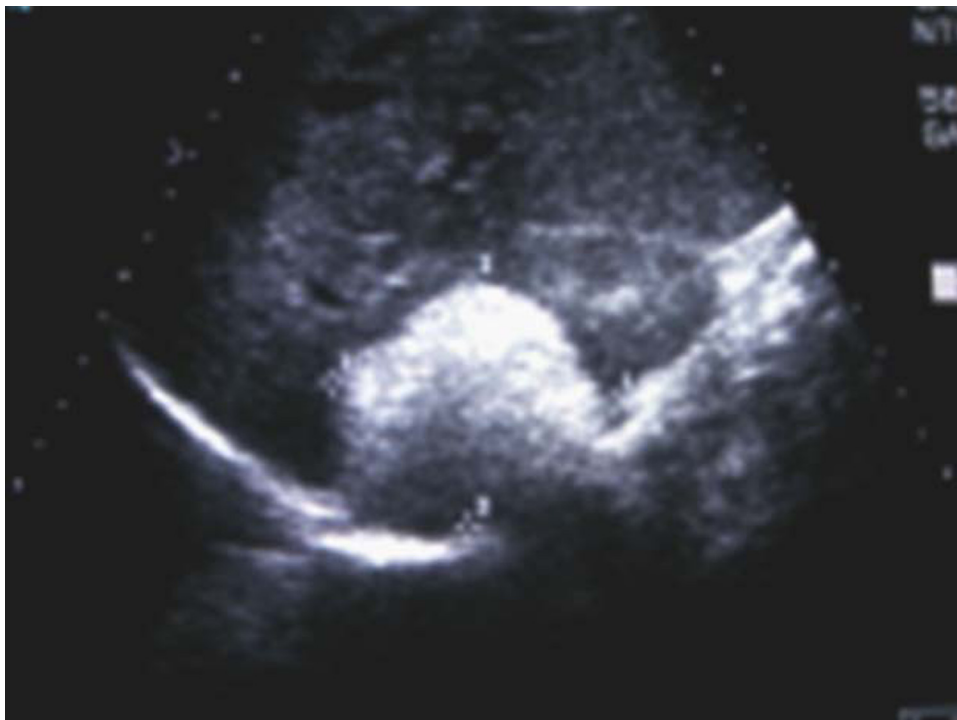


Figure 1 Large uniformly hyperechoic mass of the upper pole of the right kidney with sharp borders where echogenicity of the mass is at least equal to that of renal sinus fat. A picture of the classic ultrasonography appearance of acute myeloid leukaemia.

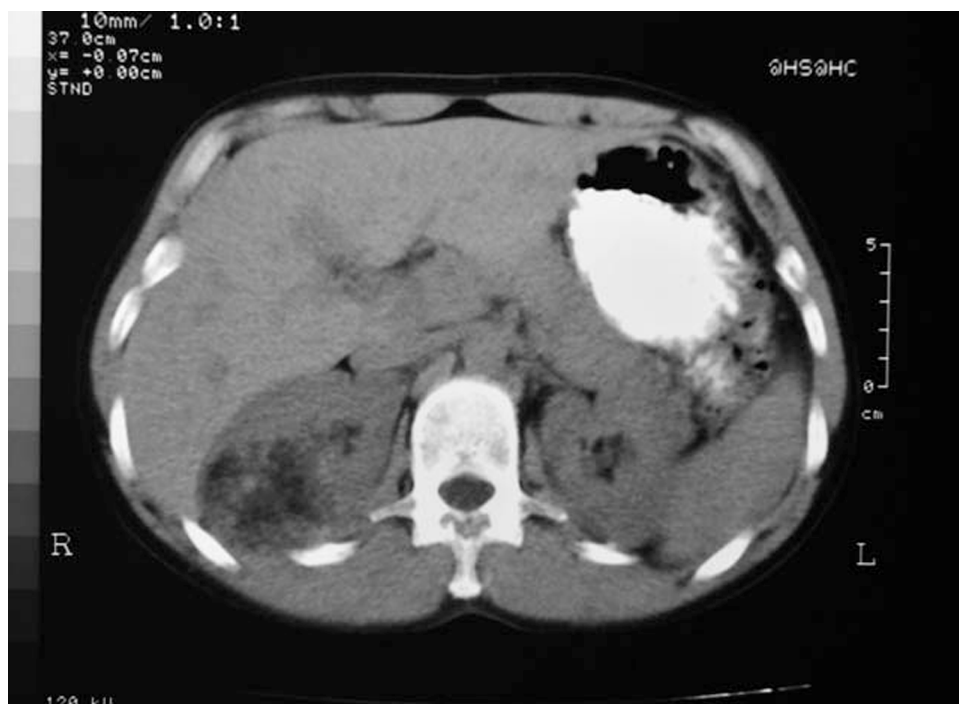


Figure 2 CT shows well-demarcated mass of the upper pole of the right kidney with mixed soft tissue and fatty tissue elements.



Figure 3 CT picture of angiomyolipoma. Enhancement of smooth muscle and vascular portions of the tumour after administration of contrast.

DISCUSSION

Angiomyolipoma is a benign renal neoplasm composed of fat, vascular and smooth muscle elements. It has a prevalence of about 0.3–3%. Two types are described: isolated angiomyolipoma and angiomyolipoma associated with tuberous sclerosis. Isolated angiomyolipoma occurs sporadically, is often solitary and accounts for 80% of all angiomyolipomas.¹ Shadowing on sonography is seen in 33% of acute myeloid leukaemias.^{2–3} In the general population in both sexes angiomyolipoma are most common in the age group 40–45 years. The mean age at presentation in patients with isolated angiomyolipoma is 43 years. This neoplasm is about four times more common in women than in men and interestingly 80% of cases involve the right kidney. Angiomyolipomas have thicker arteries than normal but abnormally weak vessel walls which predispose to aneurysm formation.⁴ Angiomyolipoma that is associated with tuberous sclerosis accounts for 20% of angiomyolipomas. The lesions are typically larger than isolated angiomyolipomas, and they are often bilateral and multiple. Angiomyolipomas occur in 80% of patients with tuberous sclerosis.¹ The male-to-female distribution of angiomyolipoma in patients with tuberous sclerosis is almost equal. Angiomyolipomas occur in young women with lymphangiomyomatosis without other stigmata of tuberous sclerosis. Although angiomyolipomas are considered benign, rare cases that are possibly related to multicentric disease have been reported regarding extension into the renal vein, the inferior vena cava, or both; deposits in the regional lymph nodes have also been reported. The risk is probably very low in tumours that are <3 cm in diameter. Apart from this group with tuberous sclerosis, follow-up may be reasonably restricted to patients with sporadic tumours >4 cm in

diameter, in whom the prevalence rate of haemorrhagic complications is higher. Thin section multidetector CT is the preferred method to demonstrate fat in problem cases (table 1).⁵

Table 1 Examination methods of angiomyolipoma

Plain abdominal radiography	Ultrasonography
CT scanning	Intravenous urography
Renal arterial embolisation	Isotope renography and dimercaptosuccinic acid
Angiography	
Percutaneous renal biopsy	DMSA scanning
MRI	Renal arterial embolisation

DMSA, dimercaptosuccinic acid.

Competing interests None.

Patient consent Obtained.

REFERENCES

1. **Rakowski** SK, Winterkorn EB, Paul E, *et al*. Renal manifestations of tuberous sclerosis complex: incidence, prognosis, and predictive factors. *Kidney Int* 2006;**70**:1777–82.
2. **Rumack** CM, Wilson SR, Charboneau JW. The urinary tract. In: Wilson SR, ed. *Diagnostic Ultrasound*. Third edition. St. Louis: Elsevier Mosby 2005:321–93.
3. **Siegel** CL, Middleton WD, Teefey SA, *et al*. Angiomyolipoma and renal cell carcinoma: US differentiation. *Radiology* 1996;**198**:789–93.
4. **Webb** WR, Brant WE, Major NM. Kidneys and ureters. In: **Webb** WR, Brant WE, Major NM, eds. *Fundamentals of Body CT*. Third edition. Philadelphia, PA: Saunders 2005:273–302.
5. **Brant** WE. Adrenal glands and kidneys. In: Brant WE, Helms CA, eds. *Fundamentals of Diagnostic Radiology*. Third edition. Baltimore: Lippincott Williams & Wilkins 2007:867–86.

This pdf has been created automatically from the final edited text and images.

Copyright 2010 BMJ Publishing Group. All rights reserved. For permission to reuse any of this content visit <http://group.bmj.com/group/rights-licensing/permissions>.
 BMJ Case Report Fellows may re-use this article for personal use and teaching without any further permission.

Please cite this article as follows (you will need to access the article online to obtain the date of publication).

Kotis A, Lisgos F, Karatapanis S Ultrasound and CT imaging assessment of renal angiomyolipoma. *BMJ Case Reports* 2010;10.1136/bcr.01.2010.2624, date of publication

Become a Fellow of BMJ Case Reports today and you can:

- ▶ Submit as many cases as you like
- ▶ Enjoy fast sympathetic peer review and rapid publication of accepted articles
- ▶ Access all the published articles
- ▶ Re-use any of the published material for personal use and teaching without further permission

For information on Institutional Fellowships contact consortiasales@bmjgroup.com

Visit casereports.bmj.com for more articles like this and to become a Fellow