## VASCULAR IMAGES

## Sarcoidosis with double saccular abdominal aortic aneurysms

Satoshi Numata, MD, PhD,<sup>a</sup> Keiichi Kanda, MD, PhD,<sup>a</sup> Tsuguru Hatta, MD, PhD,<sup>b</sup> Syuuji Tanda, MD, PhD,<sup>b</sup> Tomoya Inoue, MD,<sup>a</sup> Kiyoshi Doi, MD, PhD,<sup>a</sup> Susumu Sasaki, MD, PhD,<sup>b</sup> and Hitoshi Yaku, MD, PhD,<sup>a</sup> Kyoto, Japan

A 60-year-old man was noted to have proteinuria and a raised serum creatinine level (1.7 mg/dL) at an annual medical check-up. Laboratory investigations with abnormal findings were serum calcium, 9.9 mg/dL; serum complement C3, 151 mg/dL; C4, 53.9 mg/dL; CH50, 69 mg/dL; and angiotensin converting enzyme, 34.4 mg/dL. A tuberculin test result was negative. Rheumatoid factor, antinuclear antibody, and immunoglobulin levels were normal. A percutaneous renal biopsy revealed noncaseating sarcoid granulomas with giant cells and a lymphocytic infiltrate. A diagnosis of sarcoidosis was made.

Contrast-enhanced multislice computed tomography (CT) angiography with three-dimensional reconstruction revealed two infrarenal saccular aortic aneurysms arising from the right side of the aorta (Cover, anterior view; A, posterior view). Sclerotic changes were present along the entire length of the abdominal aorta.

We proceeded to surgical repair of the abdominal aortic aneurysm through a median laparotomy. At operation, there was significant para-aortic lymphadenopathy. The aortic wall was highly sclerotic, partially calcified, and friable. Histologic findings of the para-aortic lymph nodes revealed typical sarcoid lesions containing giant cells and noncaseating sarcoid granulomas (B). The aneurysmal wall contained atheromatous plaque with a nonspecific lymphocytic infiltration (C). These findings are consistent with a diagnosis of sarcoid aortitis.

## COMMENT

Sarcoidosis involving the aorta is extremely rare. Lesions can be found at various sites within the thoracic or abdominal aorta.<sup>1</sup> A few cases of surgical reconstruction for abdominal aortic aneurysms with sarcoidosis have been reported.<sup>2-3</sup>

Occasionally, sarcoid granulomas are not found in the aneurysmal wall.<sup>1-3</sup> In these cases, histologic findings are mainly atherosclerosis and myxoid degeneration. As in our patient, surgical reconstruction can be difficult because of the friability of aortic tissue.<sup>2</sup> This is secondary to the inflammatory process within the aortic wall. Preoperative systemic corticosteroid therapy may ameliorate this process and reduce tissue friability.<sup>3</sup>

We present a case of successful repair of a sarcoid abdominal aortic aneurysm. As there are very few cases of this condition in the literature, evidence for successful management is anecdotal. Furthermore, long-term outcome for surgical intervention for sarcoid aortic aneurysms is unknown. As such, we advocate close monitoring using conventional imaging techniques such as CT.



## REFERENCES

- 1. Maeda S, Murao S, Sugiyama T, Utaka I, Okamoto R. Generalized sarcoidosis with "sarcoid aortitis". Acta Pathol JPN 1983;33:183-8.
- 2. Naraynsingh V, Raju GC. Sarcoid aortic aneurysm: surgical difficulties. J R Coll Surg Edinb 1987;32(3):167-8.
- 3. Gedalia A, Shetty AK, Ward K, Correa H, Venters CL, Loe WA. Abdominal aortic aneurysm associated with childhood sarcoidosis. J Rheumatol 1996;23:757-9.

From the Department of Surgery, Division of Cardiovascular Surgery,<sup>a</sup> and Department of Medicine, Divi-

sion of Hypertension and Nephrology,<sup>b</sup> Kyoto Prefectural University of Medicine.

J Vasc Surg 2005;41:1065

0741-5214/\$30.00

Copyright © 2005 by The Society for Vascular Surgery. doi:10.1016/j.jvs.2004.10.043