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# Antineutrophil cytoplasmic antibody-associated vasculitis presenting as kidney rupture and gonadal artery hemorrhage

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A 70-year-old man was transferred to our department of nephrology for the evaluation of persistent fever and azotemia. He has been treated with anti-tuberculous (Tb) medication for Tb pleurisy in recent weeks. The laboratory findings at transfer were as follows: white blood cells, 25,030/mm<sup>3</sup>; hemoglobin, 11.8 g/dL; platelets, 545,000/ mm<sup>3</sup>; C-reactive protein, 13 mg/dL; protein/albumin, 6.1/2.6 g/dL; blood urea nitrogen, 28 mg/dL; and creatinine, 2.06 mg/dL. Urinalysis showed both proteinuria (4+) and hematuria (4+). Serologic markers including immunoglobulin and complements were within normal range, except that myeloperoxidase anti-neutrophil cytoplasmic antibody (ANCA) was positive. On suspicion of ANCA vasculitis, 500-mg methylprednisolone was administered. While planning renal biopsy, the patient complained of severe right flank pain, and his hemoglobin level decreased from 11 to 7 g/dL. Abdominopelvic computed tomography showed kidney rupture with a subcapsular hematoma, and renal angiography revealed innumerous punctate contrast staining and a minute leak in the vascular bed of right renal artery (Fig. 1); total embolization of the right renal artery was thus performed. After 2 days, a decrease in hemoglobin to 2.5 g/dL occurred. A focal contrast extravasation arising from the right gonadal artery was seen on further abdominal aortography; this culprit vascular branch was embolized again (Fig. 2). Thereafter, right nephrectomy with hematoma evacuation was done. Macroscopically, the excised kidney revealed diffuse hemorrhagic infarct, and microscopically showed total ischemic necrosis with suspected diffuse vasculitis (Fig. 3). Under a diagnosis of microscopic polyangiitis based on serology and histology, steroid, intravenous immunoglobulin, plasmapheresis, and continuous renal replacement therapy were applied. Taking cyclophosphamide after stabilization, he recovered enough to survive without dialysis. Creatinine level was around 2 mg/dL prior to discharge.

ANCA-associated vasculitis mainly involves small blood vessels, but can also rarely involve the middle arteries, which may be fatal if diagnosis and treatment are delayed. In the case of fever of unknown origin and renal impairment, this type of vasculitis should be considered in differential diagnosis.

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**Figure 1. Abdominopelvic computed tomography and right renal angiography.** (A) Nonenhanced computed tomography shows kidney rupture with a large subcapsular hematoma (arrows) in the right kidney. (B) Right renal angiography reveals contrast extravasation (arrow) from a branch of the upper polar area and innumerous punctate staining throughout the renal vascular bed.



**Figure 2.** Abdominal aortography. (A) Abdominal aortogram shows focal contrast extravasation (arrow). (B) The culprit branch arose from the right gonadal artery (arrows) and was embolized with an *n*-butyl-2-cyanoacrylate and lipiodol mixture.



**Figure 3. Macroscopic and microscopic features of the excised kidney.** (A) Macroscopically, sectioned surfaces of the kidney exhibited diffuse hemorrhagic infarct. (B) Microphotograph of the kidney shows total ischemic necrosis (H&E stain, ×100).

## **Conflicts of interest**

All authors have no conflicts of interest to declare.

#### **Data sharing statement**

The data presented in this study are available on request from the corresponding author.

### **Authors' contributions**

Writing-original draft: HSL Writing-review & editing: MJC, IHS, SMK All authors read and approved the final manuscript.

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