Ureteroarterial fistula: imaging diagnosis and endovascular management

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DESCRIPTION

A 65-year-old woman presented with haematuria and hypovolaemic shock. Her medical history includes International Federation of Gynaecology and Obstetrics (FIGO) stage IIIC2 serous endometrial carcinoma diagnosed 5 years previously, which was treated with transabdominal hysterectomy and bilateral salpingo-oophorectomy, adjuvant chemotherapy and radiotherapy. Two years after initial treatment, she was diagnosed with disease recurrence in the right ureter and new nodes in the coeliac, left para-aortic and right retrocrural regions. At that time, she underwent further chemotherapy and placement of a right ureteric stent for obstructive uropathy.

Her estimated glomerular filtration rate (GFR) was 36 mL/min/1.73 m². Cystoscopy and ureteroscopy performed showed normal bladder, left ureteric and distal right ureteric mucosa. It was not possible to advance the ureteroscope beyond the distal right ureter. CT scan showed clots within the right renal calyces. The right kidney also showed significant renal cortical atrophy on CT scan, suggestive of a non-functioning right kidney. This was confirmed by a subsequent dimercaptosuccinic acid (DMSA) scan, which showed 5% split function in the right kidney. The right kidney was thought to be the source of haematuria, and she proceeded to have a right nephrectomy and removal of the ureteric stent. The lower right ureter was not removed due to severe fixation.

Twelve days later, she had a further episode of massive haematuria. A repeat CT scan revealed an enhancing abnormality within the ureteric remnant, corresponding to the point of contact between the previous ureteric stent and the right common iliac artery (CIA) on the previous CT scan (figure 1). She proceeded to angiography that showed a CIA pseudoaneurysm. A 9×37 mm stent graft (BeGraft, Bentley, Hechingen, Germany) was deployed within the CIA across the defect (figure 2). This resulted in resolution of her haematuria by postprocedure day 1. She remained asymptomatic 9 months following the procedure.

Ureteroarterial fistula (UAF) is a rare condition with only approximately 150 cases reported in the literature.1 Risk factors include chronic indwelling ureteral stents, pelvic surgery and radiotherapy and systemic chemotherapy. The diagnosis of UAF is often delayed because it is an uncommon cause of haematuria. CT scan may show an iliac artery injury near the crossing of the ureter, with ancillary findings of blood clots in the collecting system. However, the diagnostic rates with CT scan are only 42%–50%.2 Angiographic findings of UAF vary from extravasation to a subtle intimal defect. Contrast extravasation into the ureter might not occur when a ureteral stent or clots are at the site of the leak. Open surgical repair in these patients...
Images in...

is often difficult due to prior intervention and radiation. Endovascular management, with placement of a stent graft to exclude the fistula, reported a success rate of 85%–100%. Although stent-graft infection is an inherent risk, such complication after treatment of UAF has not been described to date.

In summary, it is important to consider the possibility of a UAF in patients who present with haematuria, an appropriate history and in whom other common causes of haematuria have been excluded. Endovascular treatment is a feasible and safe approach in managing a patient with UAF.

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