Xanthogranulomatous pyelonephritis

Tahmina Begum, Muhammad Ekramul Huq, Mashrafi Ahmed

Texas Tech University, Amarillo,
Texas, USA

Correspondence to
Dr Mashrafi Ahmed,
mashrafi.ahmed@gmail.com

Accepted 2 June 2016

DESCRIPTION

A 48-year-old African-American woman was admitted to the hospital, because of weight loss and poor appetite for the past 3 months. The patient mentioned that she had nearly 30 lb of unintentional weight loss during this period. She had neither nausea, vomiting nor chest pain, but had felt mildly feverish with left-sided back and flank pain for the past 2 weeks. The pain was 5/10 in intensity. There was no change in urine frequency, and neither colour change in the urine nor burning on micturition. There was no blood in the urine. The patient mentioned occasional night sweats for the past 2 weeks as well. On two occasions, respectively, she measured her temperature to be 102°F and 101.6°F. At the time of admission, her temperature was 99.6°F and heart rate 120 bpm. Physical examination revealed left-sided costovertebral angle tenderness with an ill-defined mass extending up to the costal margin. Initial white cell count was $22.6 \times 10^3/\mu\text{L}$ with 85% neutrophils, and urinalysis was indicative of urine infection. A CT scan of the abdomen and pelvis with intravenous contrast revealed a markedly enlarged left kidney with diminished perfusion. There were multiple low-density areas throughout the kidney, suggestive of necrosis or abscess (figures 1 and 2; yellow arrow). There was a little stag horn calculus at the renal pelvis (figure 2; green arrow). There was also perinephric fat stranding with retroperitoneal lymphadenopathy. The diagnosis of xanthogranulomatous pyelonephritis was suggested. A renal biopsy was performed to rule out malignancy, which further confirmed chronic granulomatous inflammatory changes in the renal parenchyma. The patient was started on an intravenous antibiotic with ceftriaxone for initial management. Eventually, she underwent left-sided total nephrectomy.

Figure 2 (A) CT scan of the abdomen and pelvis showing xanthogranulomatous pyelonephritis (yellow arrow) and stag horn calculus at the renal pelvis (green arrow).

Figure 1 (A) CT scan of the abdomen and pelvis showing a large left kidney with multiple areas of low density suggestive of necrosis or abscess (yellow arrow).

Learning points

- ▶ Xanthogranulomatous pyelonephritis is a granulomatous inflammation of the renal parenchyma due to chronic urinary tract infection often associated with renal stones.
- ▶ On the ultrasound and CT scan image, it is detected as an enlargement of the kidney, with multiple hypoechoic areas, calyceal dilation, necrosis, abscess and, often, inflammation extending to the perinephric fat and retroperitoneum.
- ▶ Fine-needle aspiration biopsy is recommended to confirm the diagnosis.
- ▶ Since it is an irreversible condition, total nephrectomy is recommended.

Xanthogranulomatous pyelonephritis is the chronic inflammation of the renal parenchyma, and is rarely encountered in clinical medicine. It

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To cite: Begum T, Huq ME, Ahmed M. *BMJ Case Rep* Published online: [please include Day Month Year] doi:10.1136/bcr-2016-216025

is characterised by tubulointerstitial damage with chronic interstitial inflammation and lymphoid follicle formation. It also consists of solid sheets of lipid filled foamy macrophages, inflammatory cells and extensive fibrosis.¹ Multiple discrete abscesses are common. Xanthogranulomatous pyelonephritis is mostly diffuse (92%) and occupationally focal (8%).¹ It usually develops during prolonged kidney infection by obstructive uropathy due to renal stones. The commonly isolated organisms are *Proteus* spp. and *Escherichia coli*.² Ultrasound and CT scan are both sensitive diagnostic tools. Review of image findings of 21 cases by Kim³ reported presence of renal calculi in 76% and hydronephrosis in 81% of cases. Since the image findings may also mimic Wilms tumour, neuroblastoma, clear cell carcinoma, malacoplakia, pyonephrosis or minimal tuberculosis, tissue

biopsy to confirm the diagnosis is recommended. Although partial nephrectomy can be performed for focal disease, most cases require total nephrectomy for resolution of symptoms.

Competing interests None declared.

Patient consent Obtained.

Provenance and peer review Not commissioned; externally peer reviewed.

REFERENCES

- 1 Samuel M, Duffy P, Capps S, *et al*. Xanthogranulomatous pyelonephritis in childhood. *J Pediatr Surg* 2001;36:598–601.
- 2 Yiğiter M, Ilgici D, Celik M, *et al*. Renal parenchymal malacoplakia: a different stage of xanthogranulomatous pyelonephritis? *J Pediatr Surg* 2007;42:E35–8.
- 3 Kim JC. US and CT findings of xanthogranulomatous pyelonephritis. *Clin Imaging* 2001;25:118–21.

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