Emphysematous and xanthogranulomatous pyelonephritis: rare diagnosis

ABSTRACT

Pyelonephritis is a pyogenic infection of renal parenchyma that involves the renal pelvis. It is generally of easy diagnosis. The present case report aims to describe two different manifestations of this infection: xanthogranulomatous pyelonephritis and emphysematous pyelonephritis, which have poor prognosis and require a more effective treatment. The two cases were women in the fiftieth and sixtieth decade of life, with diabetes mellitus and history of weight loss. The diagnosis of the renal infection was established through computed tomography and the treatment was based in surgical procedure, with favorable outcome.

Keywords: xanthogranulomatous pyelonephritis, emphysematous pyelonephritis, urinary tract infection, nephrostomy, nephrectomy, surgery.

INTRODUCTION

Pyelonephritis is a pyogenic infection of renal parenchyma that involves the renal pelvis. It generally begins through an ascending infection that affects renal tubules, interstitium, glomeruli and vessels. The acute form manifests with fever and lumbar pain, which can be associated with inflammatory symptoms of the bladder. The chronic forms are the result of cicatricial effects and repeated infections, can present as an insidious disease and can lead to end-stage renal disease. The diagnosis of pyelonephritis is easy in the majority of cases, but there are unusual presentations that make diagnosis more difficult. The aim of this study was to present two hard to diagnose unusual cases of pyelonephritis (xanthogranulomatous pyelonephritis and emphysematous pyelonephritis).

CASE REPORT

Case 1

A 60-year-old woman was admitted to the emergency room with complaints of epigastric pain, vomiting, asthenia and weight loss for three weeks, complicated by digestive tract hemorrhage (hematemesis and melena), myalgia, obtundation and loss of consciousness. She had history of diabetes mellitus, with irregular use of glibenclamide, and smoking. At physical examination, she was confuse, with jaundice (2+/4+), blood pressure 100/40 mmHg, pulse 188 bpm, respiratory rate 28 rpm, temperature 36.5 °C, and distended abdomen, painful at palpation. Laboratory tests showed Ht 25.9%, Hb 9.2g/dL, WBC 43,800/mm³, with 78% neutrophils, creatinine 2.9 mg/dL, urea 84.7 mg/dL, glicemia > 500 mg/dL, LDH 2,141 mg/dL, and no electrolyte abnormality. Abdomen x-ray showed the presence of air in the renal topography (Figure 1). An abdominal ultrasound and CT scan showed bilateral emphysematous pyelonephritis, with no evidence of urinary stone obstruction (Figure 2). After antibiotic therapy the patient underwent left nephrectomy and had a favorable outcome.

Case 2

A 48-year-old woman was admitted to the emergency room with a history of lumbar pain for two weeks, associated with asthenia, macroscopic hematuria and severe weight loss (15 kg in three months). She also had diaze-
Figure 1: Abdomen plain x-ray shows the presence of air in both renal topography, more evident in the left side.

Figure 2: CT scan showing bilateral emphysematous pyelonephritis.

Figure 3: CT scan of xanthogranulomatous pyelonephritis. There is diffuse enlargement of the left kidney and the renal tissue is replaced by multiple low-attenuated masses. The contralateral kidney is normal.

DISCUSSION

Xanthogranulomatous pyelonephritis and emphysematous pyelonephritis are two rare, atypical and severe forms of renal parenchyma infection. Emphysematous pyelonephritis was first described by Kelly and MacCullum, in 1898, and xanthogranulomatous pyelonephritis, by Schlagenhaufer, in 1916, termed staphylocomycosis due to its similarity with actinomyces and the presence of Staphylococcus. In 1935, it was named xanthogranulomatous pyelonephritis by Oberling.

Xanthogranulomatous pyelonephritis is characterized by a chronic inflammatory process, with destruction of renal parenchyma, subsequently replaced by a granulomatous tissue, containing mononuclear macrophages and lipids (Xanthomam Cells). This clinical entity represents 1% of all renal infections. The disease is four times more frequent among women between the fiftieth and sixtieth decades of life, but can occur at any age. In the majority of cases the disease is unilateral, and the right kidney is more often involved. Bilateral cases are thought to be fatal. Patients with xanthogranulomatous pyelonephritis commonly have diabetes or immunodepression.

Emphysematous pyelonephritis is a necrotizing infection of the renal parenchyma characterized by the production of gas in the intra- and peri-renal tissues.
believed that high levels of glucose, in association with inadequate perfusion, lead to a favorable environment for the growth of anaerobic organisms. This disease affects individuals of all ages, but women are six times more likely to be affected.\textsuperscript{5,12,13}

Many conditions are responsible for the pathogenesis of both diseases, including urinary tract obstruction due to kidney stones, ineffective treatment of urosepsis, renal ischemia, lymphatic obstructions, abnormalities in the lipids metabolism and abnormal immune response.\textsuperscript{5,12}

Xanthogranulomatous pyelonephritis is associated with renal stones in 75-86% of cases. The most common associated infectious agents are \textit{Proteus} and \textit{E. coli}, which are responsible for 30-40% of cases.\textsuperscript{8} Approximately 10% of patients have negative cultures.\textsuperscript{7} In emphysematous pyelonephritis, the most common agents are \textit{P. mirabilis}, \textit{E. coli}, and \textit{K. pneumoniae}.\textsuperscript{5,12,13}

Xanthogranulomatous pyelonephritis is most frequently misdiagnosed as renal carcinoma due to its clinical presentation and radiographic appearance.\textsuperscript{14} Evidence of chronic urinary tract infection and CT scan findings usually make it easier to differentiate these disorders. Although rarely, the two disorders can occur together. Xanthogranulomatous pyelonephritis must also be distinguished histologically from two other inflammatory conditions, namely renal parenchymal malakoplakia and megalocytic interstitial nephritis.\textsuperscript{15}

CT scan is now the best tool to diagnose these infections, and it is also important to establish the presence and extension of extra-renal involvement.\textsuperscript{4,6,8,12,16} The most frequent findings in the CT scan are calculi, hydronephrosis, kidney enlargement and hypodense areas, with parenchyma destruction.\textsuperscript{5,12,13} Other diagnostic methods that can be used in these conditions are voiding cysotretherography, ultrasound and renal scintigraphy.\textsuperscript{4,10,12,16}

The gold-standard therapy for both infections is nephrectomy, which is total in the majority of cases. Circumjacent inflammatory tissues should be removed. In rare cases, partial nephrectomy can be successfully performed.\textsuperscript{1,2,4,6,8,12,13,16} Nephrostomy before nephrectomy can be considered a method that facilitates surgery, because it allows a reduction in renal mass and favors the access to the kidney at the time nephrectomy is done.\textsuperscript{5,12}

Antibiotics alone are not effective for these infections, but should be initiated before surgical procedure in order to control the infectious process and avoid systemic involvement (sepsis).\textsuperscript{2,4,12}

In conclusion, xanthogranulomatous pyelonephritis is an unusual variant of chronic pyelonephritis. Most cases occur in the setting of obstruction due to infected renal stones. Affected patients usually have massive destruction of the kidney due to granulomatous tissue; the appearance may be confused with renal malignancy.

\textbf{REFERENCES}