Case Report

Pyonephrosis Presenting as Lumbar Abscess—An Uncommon Clinicoradiological Entity

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Abstract

Pyonephrosis secondary to an obstructing calculus in renal pelvis, pelviureteric junction or ureter usually presents clinically with fever, loin pain and other signs of urinary tract infection. Rarely, severe thinning of renal parenchyma in pyonephrosis may allow direct fornical rupture into the retroperitoneum producing retroperitoneal abscess that might be the only presenting complaint without any complaints related to renal system. Prompt and aggressive management including optimal imaging and correct diagnosis is needed in such cases to prevent further complication. Hence, this article focuses on the rare spontaneous rupture of calculus pyonephrosis in to ipsilateral psoas & quadratus lumborum muscle to clinically present as a lumbar pain & abscess, presentation not described before in medical literature.

Introduction

Complete obstruction of renal collecting system causing moderate to severe hydroureteronephrosis can uncommonly undergo spontaneous decompression following a caliceal rupture [1-2]. This spontaneous rupture of collecting system is usually secondary to an obstructing calculus but less commonly may be due to posterior urethral valves, benign prostatic hyperplasia, pelvic tumors and rarely abdominal aortic aneurysm & retroperitoneal fibrosis [3-4]. Occasionally, the urine in collecting system gets infected forming pus (pyonephrosis) with parenchymal destruction rarely rupturing in to retroperitoneum in to perinephric space, psoas muscle and quadratus lumborum muscle [5,6]. In this article, we present a rare case of spontaneous rupture of pyonephrotic kidney secondary to pelviureteric junction calculus presenting as a lumbar abscess.

Case report

A 27-year old female presenting to outpatient department complained of fever with chills and pain in left lumbar region. Patient was visiting local doctors in rural area for fever for few weeks prior to present visit and wad receiving antibiotics without any laboratory or radiological investigations. Clinical examination revealed a soft & tender bulge in left lumbar region with signs of inflammation. Laboratory test revealed raised total leucocyte count (16000 cells/microliter of blood) with predominant neutrophilia (>90%). Blood urea nitrogen and serum creatinine values were within normal limits being 18mg/dL and 1.1mg/dL respectively. Urine microscopy revealed 10–11 pus cells/high-power field. Chest radiograph in posteroanterior view did not reveal any significant abnormality.

Ultrasoundography of abdomen revealed grossly-enlarged, hydroureteric left kidney measuring up to 170*98 mm [Superoinferior*Anteroposterior] with severe thinning of renal parenchyma, low–level internal echoes in collecting system and a large, calculus measuring up to 23–24 mm in maximum dimension located at the pelviureteric junction. In addition, a septeate collection measuring up to 110*110*70 mm (corresponding approximately up to 400–450 ml in volume) was noted in the ipsilateral psoas and quadratus lumborum muscles; latter corresponding to the clinically palpable lumbar swelling. The above-described collection was extending in to the psoas up to the left iliacus muscle causing its thickening.

A contrast-enhanced, computed tomography (CECT) of
abdomen revealed a large, calculus of 26*11*9 mm in maximum dimensions with 900–1000 HU attenuation value causing a gross hydronephrosis in left kidney which reveal very thin but enhancing, rim-parenchyma and high-attenuating fluid (25–30 HU) along with few subcentimeter secondary caliceal calculi in anterior midpolar calices without aerocales (Figure 1). In addition, there was a fairly-defined, thick-walled, septate collection without obvious internal calcification/air in left psoas & quadratus lumborum muscles extending up to left iliacus muscle measuring 500–550 ml in volume (Figure 2). The above-described myofascial collection was apparently communicating with posterior inferior-polar calices of left kidney associated with soft-tissue stranding in the visualised pararenal / retroperitoneal fat. No obvious abnormality was noted in the bony spine. No evidence of any free fluid was noted in the peritoneal cavity.

Based on the radiological findings, enlarged left kidney with Grade-III hydrenephrosis causing by PUJ calculus showing secondary caliceal calculi, signs of pyonephrosis and left iliopsoas & quadratus lumborum collection secondary to spontaneous caliceal rupture was made. Fine-needle aspiration was performed from both the collecting system of left kidney and quadratus lumborum muscle not only to confirm the presence of pus & detect etiological agent but also to establish the cause and effect relationship. Culture of pus revealed Escherichia coli from both sites.

The patient was treated nephrostomy with pyelolithotomy, placement of ureteral stent and drainage of myofascial collection. Significant clinicoradiological improvement in form of near-complete resolution of retroperitoneal & pararenal pus without any other complication was noted on one-month follow-up of patient. Due to financial constraints, CECT abdomen with urography phase was postponed to a later date.

Discussion

Retroperitoneal abscess especially psoas abscess secondary to caliceal rupture in calculus-induced pyonephrosis is a very rare phenomenon [5]. The psoas abscess thus formed may be detected more sensitively with computed tomography rather than with ultrasonography [7]. Rarely, it may further extend in to the muscles of the posterior abdominal wall especially quadratus lumborum when it may present as a lumbar abscess as in our index case. Management is usually with broad-spectrum antibiotics in addition to image-guided percutaneous or surgical drainage of pus collection by nephrostomy & ureteral stent [5]. Surgical removal of affected kidney is the treatment of choice with non-functioning, affected and normal-functioning contralateral kidney [8–9].

Thorough search in to the existing English medical literature revealed less than five reported cases of spontaneous rupture of pyonephrosis secondary to urolithiasis with psoas abscess formation [5]. Posterior abdominal wall abscess in lumbar region was an additional finding in our case not reported in literature. However, one case showing lumbar panniculitis with subcutaneous abscess secondary to pyonephrosis has been reported [10]. Rarely, peritoneal rupture of pyonephrosis causing peritonitis & splenic abscess has been also described in the literature [11-12].

Conclusion

Spontaneous retroperitoneal rupture in pyonephrosis forming a psoas & quadratus lumborum abscess is a rare occurrence. Furthermore rarer is the presentation of pyonephrosis as a lumbar abscess. Contrast-Enhanced Computed Tomography (CECT) abdomen is superior to ultrasonography abdomen in demonstrating the retroperitoneal rupture of pyonephrosis and its extent. Hence, pre-procedural CECT abdomen should form part of usual protocol for evaluation of all lumbar abscesses not only to rule out this rare entity by detecting renal cause of lumbar abscess but also to minimize morbidity by early diagnosis and management.
References


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