Giant Ureteral Stone in a Pediatric Patient: Case Report

Yalcin Comert HS¹, Salci G¹, Sarihan H¹, Bozoglu ST¹, Ozdogan EB² And Eyuboglu I³

¹Karadeniz Technical University, Faculty of Medicine, Department of Pediatric Surgery, Trabzon, Turkey
²Karadeniz Technical University, Faculty of Medicine, Department of Pediatric Nephrology, Trabzon, Turkey
³Karadeniz Technical University, Faculty of Medicine, Department of Radiology, Trabzon, Turkey

*Corresponding author: Yalcin Comert HS, MD, Assist. Prof, Karadeniz Technical University, Faculty of Medicine, Department of Pediatric Surgery, TR61080, Trabzon, Turkey, Fax: +90 462 3250518, Tel: +90 5327419246, E-mail: sonayyalcin@hotmail.com


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Abstract

A 9 year old boy referred to the hospital with abdominal pain insisting for 2 weeks. In his history there was any clinic symptom before. A giant opacification was detected in the distal part of the right ureter on plain abdominal film. Hematuria, hyperuricosuria, hyperoxaluria and hypocitraturia in his urinary analysis. Only a small right kidney (right kidney: 62x25mm; left kidney: 100x35mm) has found in the abdominal ultrasonography and an approximately a stone 42x17mm in seizes has detected in the magnetic resonance urography. In DMSA evaluation of the right kidney was 15.96% and the left kidney was 84.04%. Ureterolitotomy was performed and the giant distal ureteral stone has removed.

Keywords: Giant Stone; Ureter; Pediatric
Introduction

Primary ureteral stones are very rare in children. Ureteral stones which are bigger than 5cm defined “giant ureteral stones” in adults in literature [1]. However in children there is any definition of the sizes of ureteral stone. Here we present a child with a distal ureteral stone which was bigger than 4cm without any conspicuous clinic symptom history.

Case

A 9-year old boy presented with blunt abdominal pain insisting for two weeks. In his history there was any clinic symptom before. His systemic and urogenital examination was normal. Urinalysis showed hematuria (33), hyperuricosuria (18.5mg/kg), hyperoxaluria (180mg/1.73m²) and hypocitraturia (313mg/1.73m²). A giant opasification was detected in the distal part of the right ureter on plain abdominal film (Figure 1). Abdominal ultrasound revealed only a small right kidney (right kidney: 62x25mm; left kidney: 100x35mm). In magnetic resonance urography; the right kidney was 103x62mm in size and it was observed that the right calyceal structures and pelvis were dilated and blunting of the calyces. There was a right giant ureteral stone with dimensions of approximately 42x17mm with dilated proximal ureter and a dilatation up to 7mm in the proximal of the stone in the right ureter (Figure 2). We did voiding cystourethrogram for exclude vesicoureteral reflux. Right kidney function was 15.96% and left kidney function was 84.04% in 99m Tc-dimercaptosuccinic acid (DMSA). We did cystoscopy first and rule out any ureterocele or ureteric abnormalities. A right ureterolitotomy operation was performed with a Mc Burney incision. During the surgery a giant distal ureteral stone with dimensions of 5x2cm was removed through a longitudinal incision (Figure 3). Therefore a double J stent was inserted into the ureter for both any underlying ureterovesical junction stenosis and for the ureters'healing. Lastly the incision was closed.

The patient was discharged from the hospital on postoperative day 3 without any complication. Analysis of the stone showed it to be composed of “struvite (\(\text{NH}_4\text{MgPO}_4\cdot6\text{H}_2\text{O}\))”. The double J stent was removed under anesthesia 15 days postoperatively.

Figure 1: Plain abdominal film demonstrates a dense opacification in the line of right distal ureter
Figure 2: Coronal T2-HASTE MR image (a) reveals hypointense calculus in the right distal ureter and coronal multislice heavily T2-weighted (3D-T2-SPIR-TSE) MR image (b), shows filling defect in distal ureter indicating calculus. The pelvicalyceal system and ureter are dilated.

Figure 3: The giant ureteral stone after removal of the ureter
Discussion

The prevalence of urolithiasis is low and being more common in developing than the industrialized countries [2]. Primary ureteral stones are very rare in children and usually secondary stones originated from kidney [1,3]. Ureteral stones are uncommon due to the smooth mucosa of ureter and regularly flowing of urine [3]. They may cause many symptoms or may be symptomless [4]. In primary cases there are some underlying pathologies like ureterocele, ureteral duplication, ectopic ureter, tuberculosis, megareter, polyp or neoplasms [1-3,5-10]. Furthermore stones may be occur because of the stasis due to ureterovesical junction stenosis or vesicoureteral reflux. We were not able to detect any evidence or underlying anatomic predisposition that could favor stone formation in our case. Our patient had metabolic hyperoxaluria.

Ureteral stones are growing longitudinally and for a long time; thence they do not cause acute obstruction such as our patient. His stone was originated probably from the kidney and stayed in the ureter for a long time; than caused stasis. Stasis prompted pyelonephritis and small kidney opposite to the left one.

The American Urological Association (AUA) guidelines, indicated that up to 98% of ureteral stones 4 mm or smaller will pass spontaneously which are based on a meta-analysis of the literature for adults [9,11]. However for childhood there is any classification of sizes of the ureteral stones.

The etiology of urolithiasis in children is widely idiopathic; but some giant ureteral stones may occur with the effect of urinary pathologies or metabolic disorders [2,4]. Some predisposition metabolic disorders accompany to the urteral stones such as hypercalciuria, hypocitraturia, hyperoxaluria, hyperuricosuria and hypocitraturia. Rates of metabolic disorders are as high as 90% with the most common abnormalities being hypercalciumia and hypocitraturia [2]. Elmaci et al evaluated that primer ureteral stones were accompanied with hyperoxaluria and hypocitraturia in Turkey in 2014 [12]. Our patient had hyperuricosuria, hyperoxaluria and hypocitraturia entirely.

Calcium-oxalateephosphate stones accounting for 80% of all stones are the most stones in children; however struvite, uric acid and cystine stones are seen less frequently [7]. Our patients’ stone analysis was composed of struvite. Struvite stones are often seen in younger age and male gender [13]. Clinical presentation of these stones are usually with urinary infection, renal colic and fever, gross hematuria [13]. Our patient had hematuria and renal colic. In the treatment of struvite stones in children, the underlying pathology must first be corrected. Since the stone of our patient was struvite stone, nutrition and hydration were regulated because it was mostly associated with infection. In addition, it was followed up with protective dual antibiotic therapy.

Actually the therapy of urteral stones depends on the size of the stone, morphology and the function of the affected kidney [8,14]. Extra-corporeal shockwave lithotripsy (ESWL) and ureteroscopy are the most used noninvasive modalities of treatment [11]. However, these noninvasive techniques are not effective or usable for large size ureteral stones. Generally open surgery is using for large size ureteral stones [1,4]. We performed urterolitotomy and remove the stone in a short time without any complication. We have used Mc Burney incision but Wieslaw et al used Phannenstiel or an oblique Gibson incision leads to a wide dissection in the retroperitoneal space [15].

Conclusion

In conclusion the rarity and atipic clinic presentation of giant ureteral stones leads to silent progression and causes problems in differential diagnosis in childhood. Also we assumed that open surgery is more favourable treatment choice for giant pediatric ureteral stones.

Conflict of interest

All authors have no conflict of interest.
References


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