Case Presentation of Preureteral Vena Cava and Review of the Literature

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ABSTRACT

Retrocaval ureter, terms are anatomically descriptive but misleading in regard to development and results from altered vascular development. This anomaly is relatively uncommon, although it has clinical relevance. The ureter typically deviates medially behind the inferior vena cava, winding about and crossing in front of it from a medial to a lateral direction, to resume a normal course, distally, to the bladder. The renal pelvis and upper ureter typically appear elongated and dilated in a “J” or fishhook shape before passing behind the vena cava. Diagnoses were confirmed with intravenous urography and patient had an open surgical repair of the anomaly. The anomaly predominantly involves the right ureter, as was observed in these reported cases. Treatment is surgical allowing for correction of the anomaly with resolution of symptoms.

Keywords: Preureteral Vena Cava; Pediatric Urology

1. Introduction

This anomaly is commonly known as circumcaval or retrocaval ureter. This variety of vascular lesion can cause ureteral obstruction. The term preurteral vena cava emphasizes that the circumcaval ureter results from altered vascular, rather than ureteral, development [1].

It was initially considered as aberration in ureteric development; however several studies in embryology have led to it being considered as an aberration in the development of the inferior vena cava [2-4].

This disorder involves the right ureter, which typically deviates medially behind (dorsal to) the inferior vena cava, winding about and crossing in front of it from a medial to lateral direction, to resume a normal course, distally, to the bladder. The renal pelvis and upper ureter are typically elongated and dilated in a “J” or fishhook shape before passing behind the vena cava [5].

2. Case Report

A 5 year-old boy since four months, he admitted having had occasional sharp transient dull and intermittent pain in the right flank.

Physical examination was normal. Complete laboratory evaluation including urinalysis, complete blood picture, urea, creatinine and electrolytes were within normal limits. KUB ultrasound showed a moderate hydronephrosis. Left kidney, left ureter and urinary bladder were normal. An intravenous pyelogram showed prompt bilateral excretion from both kidneys and a normal left upper urinary tract. On the right side a moderate hydronephrosis associated with caliectasis was observed and it was noted that the upper ureter was S-shaped and was kinked medially towards the midline at the level of the transverse process of the third lumbar vertebra. The ureter could not be visualized beyond that point (Figure 1). Retrograde ureteropyelography demonstrates and S curve to the point of obstruction, with the retrocaval segment lying at the level of L3 or L4 suggesting the presence of a retrocaval ureter (Figure 2). The right ureter was explored through a right-flank incision. On exploration, proximal ureter was curved medially then posterior to IVC. Finally curved anteromedially to IVC and took a downward course (Figure 3). Surgical correction involves ureteral division, with relocation and ureteroureteral reanastomosis (Figure 4). A simple ureteral stent was inserted in an antegrade manner during operation. An intravenous pyelography, and renal ultrasonography were performed 3 months postoperatively, showed regression...
Figure 1. Intravenous pyelogram showed that the upper ureter was S-shaped and was kinked medially towards the midline at the level of the transverse process of the third lumbar vertebra.

Figure 2. Retrograde ureteropyelography demonstrates an S curve to the point of obstruction, with the retrocaval segment lying at the level of L3 or L4 suggesting the presence of a retrocaval ureter.

Figure 3. On exploration, proximal ureter was curved medially then posterior to IVC and finally curved anteromedially to IVC and took a downward course.

Figure 4. Following the confirmation of obstruction, surgery was indicated in the form of pyelic sectioning and ureteral transpositioning of the retrocaval segment.

Figure 5. The ureteric segment behind the vena cava was repositioned medially and anteromedially to anastomose with the pelvic ureter and returned to a normal ureteral position.

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3. Discussion

The first observed case of retrocaval ureters was described by Hochstetter in 1893 [6]. Though initially thought of as an anomaly of ureteric development, studies in embryology have revealed an anomaly related to the development of the inferior vena cava [7-9].

This anomaly is commonly known as circum-caval or retrocaval ureter [5]. The term of circumcaval ureter is preferred, because rarely a ureter may lie behind (dorsal to) the vena cava for some portion of its lumbar course, forming a “siphon” capable of causing urinary obstruction. The anomaly predominantly involves the right ureter, as was observed in these our reported cases. If it involves the left ureter then it is usually associated with

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either partial or complete situs inversus or duplication of the inferior vena cava (IVC) [10,11]. Duplication of the IVC (D-IVC): This is a relatively uncommon congenital anomaly with a reported incidence of 0.2%-3%. A majority of the cases are clinically silent and they are diagnosed incidentally during imaging studies which are done for other reasons [12]. Retrocaval ureter results from altered vascular, rather than ureteral, development. Bateson and Atkinson distinguished the two types of retrocaval ureters according to the radiological appearance and the site of the ureteral narrowing. These are:

Type I: The ureter crosses behind the IVC, at the level of the L3 vertebra and it exhibits an “S-shaped” deformity.

Type II: The renal pelvis and the upper ureter lie horizontally. The retrocaval segment of the ureter is at the same level as that of the renal pelvis and it exhibits a “sickle shaped” deformity [13].

The retrocaval ureter which was observed in our case classified into the Type I of the given classification. The incidence of preuretoral vena cava at autopsy is about one in 1500 cadavers, although the lesion is congenital, most patients do not present until the third or fourth decade of life [5]. Clinically, may present with symptoms of flank or abdominal pain or infection or the disorder may be discovered incidentally during other radiological tests. This disorder can cause varying degrees of ureteral obstruction. In order to reduce irradiation, the scintigraphy scan is likely to replace IV urography, CT urography and diuretic renography. Excretory urography often fails to visualize the portion of the ureter beyond the J hook, but retrograde ureteropyelography demonstrates an S curve to the point of obstruction with the retrocaval ureter lying at the level of L3 or L4 [14].

In our cases Intravenous pyelogram showed that the upper ureter was S-shaped and was kinked medially towards the midline at the level of the transverse process of the third lumbar vertebra. Also we perform the retrograde ureteropyelography and demonstrate the S curve of the retrocaval ureter. MRI can demonstrate the course of a preuretreal vena cava, and may be a more detailed and less invasive imaging procedure, compared with CT and retrograde Pyelography [15]. Surgical repair is indicated only when symptoms are present or significant obstruction exist that have repercussion in renal function. Surgical correction involves ureteral division, with relocation and ureteroureteral or ureteropelvic reanastomosis [16, 17]. Laparoscopic and robotic minimally invasive repair of the ureter has been described by a trans or retroperitoneal approach and should be considered before open surgery [18,19]. In our cases following the confirmation of obstruction, surgery was indicated in the form of proximal ureteric sectioning and ureteral transpositioning of the retrocaval segment.

4. Conclusion

Retrocaval ureter should be suspected in any case of pyelectasis and proximal uretereoclastasis respectively of the upper third ureter on the right side.

REFERENCES


**Abbreviations**

IVC: Inferior Vena Cava  
D-IVC: Duplication of the IVC  
KUB: Kidney Urinary Bladder