Case Report

We report a case of an 11-year-old male child who presented with left sided lump in abdomen since one year. Patient had a history of intermittent constipation but no history of any urinary tract complaints. He had no history of renal stone disease or lithuria in past. He had no significant past medical history or any urological intervention. Renal function test and hemogram was within normal limits. Urine culture for Tuberculosis (TB) and urine for TB polymerase chain reaction (PCR) were negative. On physical examination, left sided lump was palpable in lumbar region extending up to iliac region and crossing midline. Lump was cystic on palpation and non tender. The ultrasonography (USG) of abdomen revealed bilateral hydronephrosis with proximal hydroureters, massively enlarged left kidney with gross hydronephrosis & thinned out parenchyma and right kidney was moderately hydronephrotic with parenchymal thickness of 12 mm and proximal hydroureter. Bladder on USG was normal. Of note no ureteric dilatation was seen at the level of the bladder. On Intravenous pyelography the left kidney was non-functioning with no uptake of contrast, while the right kidney showed moderate hydronephrosis with dilatation of proximal ureter with delayed uptake and retention of contrast, which were suggestive of stricture in right sided mid ureter. Tc99m Diethylene Triamine Pentaoacetic Acid (DTPA) scan showed non-functioning left kidney and decreased function and retention on right side. We did a left nephroureterectomy and uretero-ureterostomy without tapering over a Double-J stent (DJ), with excision of stenosed segment on right side. Pathological review of the ureteric segments of both sides revealed proximal dilatation with non-specific thickening of the muscular wall of the stenosed segments of ureter. The differential diagnosis must additionally include ureteral valves and fibro epithelial polyps. Important role of intravenous pyelography in such cases must be emphasized to localize the pathology site. In addition, retrograde ureteropyelography is essential in management of this anomaly to delineate the stenosed segment of the ureter preoperatively, so that the surgeon can be well prepared with final plan of management.

Keywords: Hydroureteronephrosis, Retrograde ureteropyelography, Uretero-ureterostomy

Surgery Section

Congenital Bilateral Mid Ureteral Stenosis: A Rare Finding

Bilateral congenital mid ureteric stenosis is a very rare entity causing hydroureteronephrosis. The pathophysiology and aetiology of this condition is uncertain. Congenital functional obstruction of the mid ureter has only 17 reported cases, that too of unilateral pathology. We report a case of an 11-year-old male child who was presented with left sided lump in abdomen since one year. On bilateral retrograde ureteropyelography (RGP), bilateral mid ureteric stenosis was documented at the level of pelvic brim. Tc99m Diethylene Triamine Pentaoacetic Acid (DTPA) scan showed non functioning left kidney and decreased function and retention on right side. We did a left nephroureterectomy and uretero-ureterostomy without tapering over a Double-J stent (DJ), with excision of stenosed segment on right side. Pathological review of the ureteric segments of both sides revealed proximal dilatation with non-specific thickening of the muscular wall of the stenosed segments of ureter. The differential diagnosis must additionally include ureteral valves and fibro epithelial polyps. Important role of intravenous pyelography in such cases must be emphasized to localize the pathology site. In addition, retrograde ureteropyelography is essential in management of this anomaly to delineate the stenosed segment of the ureter preoperatively, so that the surgeon can be well prepared with final plan of management.
was excised revealing a pin-hole lumen. No valves or polyps were noted intraluminally. Ureter-ureterostomy was performed without tapering over a DJ stent. The post operative course was uneventful with good recovery. Cystoscopic DJ removal was done at 12 wk postoperatively. Follow up ultrasonography at three months showed improvement in hydronephrosis on right side. Follow up DTPA scan after Double-J stent removal showed improvement in drainage pattern of right kidney, with only 40% retention at 20 min

Pathological review of the ureteric segments of both sides revealed proximal dilatation with non-specific thickening of the muscular wall of the stenosed segments of the ureters. The left kidney histopathology report was consistent with obstructive changes. There were no structural changes in the muscular wall of the ureters on both sides.

DISSCUSSION

Bilateral Congenital mid ureteric stricture is a very rare entity causing hydroureteronephrosis [1], and the diagnosis can be confused with ureteropelvic or ureterovesical junction obstruction. Although ureteral stricture or strictures most commonly occur in distal ureter and are associated with megaureters, these segments also occur in mid ureters but are rare findings. The pathophysiology and aetiology of this condition is uncertain. Developmentally, narrowing probably results from a disturbance in embryogenesis around the 11th–12th weeks with disturbed development of mesenchyme contributing to the ureteral musculature. In children with congenital mid ureteral stenosis an association with contra lateral renal agenesis or athrophy has been reported, suggesting possible bilateral aberrant ureteral development [2]. The differential diagnosis must additionally include ureteral valves and fibro epithelial polyps. Congenital mid ureteric obstruction runs a more aggressive natural history than other causes of congenital hydronephrosis [3]. Congenital functional obstruction of the mid ureter has only 17 reported cases, that too of unilateral pathology [4]. After extensive literature search we found only a few case report of bilateral mid ureteral stenosis by Jayanthi VR et al., H Rode et al., and Domenichelli V et al., [1,4,5]. This feature is also suggested by our case where longstanding obstruction resulted in renal impairment. The pathological appearance of mid ureteric obstruction is given by Allen [6], in which the segments were stenosed but muscle fibres were normal which suggested functional obstruction. Term such as “Midureteral adynamic segment” has been in use for the last few years only [5].

Different theories are postulated for congenital mid ureteral stenosis such as

1) Compression by iliac vessels.
2) Failure of recanalization of ureter.
3) Acquired foetal inflammation of the ureter similar to biliary atresia.

Although ultrasonography may give us anatomic details but it fails to exact localize such unusual pathological lesions. Important role of intravenous pyelography in such cases must be emphasized to localize the pathology site. Magnetic resonance urography (MRU) provides excellent anatomic and functional detail of the collecting system, leading to accurate diagnosis and management of ureteral stricture in children [7]. In addition, retrograde ureteropyelography is essential in management of this anomaly to delineate the stenosed segment of the ureter preoperatively, so that the surgeon can be well prepared with final plan of management. Definitive management includes open or laparoscopic resection of the stenosed segment with uretero-ureterostomy. There are case reports of laparoscopic resection and ureteroureterostomy for congenital midureteral stricture [8].

CONCLUSION

There should be a systematic sequence of imaging studies for the evaluation of hydronephrosis such as ultrasonography, micturating cystourethrogram, renal nuclear scan, intravenous pyelography, magnetic resonance urography and if there is diagnostic dilemma then one should proceed with retrograde ureteropyelography (RGP) as it can help us to locate and exactly measure the stenosed segment length. With the help of these diagnostic modalities and their accuracy, we will be able to plan a specific surgical intervention, which can definitely improve our surgical outcome and benefit the patients.

REFERENCES


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