Congenital Posterior Urethral Diverticulum in a Male Child
Case Report

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Abstract
Bladder neck and posterior urethra are common sites for obstructive uropathy in children. Diverticula of posterior urethra are rare cause of obstruction in children. A six year old boy presented with features of bladder outlet obstruction since birth. Ultrasound findings were suggestive of posterior urethral valve. Micturating cystourethrogram and endoscopic examinations revealed posterior urethral diverticulum which was placed dorsally. Diverticulectomy and reconstruction of urethra was performed by midline perineal incision. This report emphasizes that a posterior urethral diverticulum may be considered in those cases where features are suggestive of posterior urethral valve bladder outlet obstruction.

Key words: Posterior urethral valve, Diverticulum, Diverticulectomy

Introduction
Posterior urethral valve causing bladder outlet obstruction with back pressure changes is known in children but similar picture may be caused by congenital posterior urethral diverticula. We report a case of six year old boy who had features of bladder outlet obstruction. Micturating cystourethrogram revealed posterior urethral diverticulum which was suspected to have posterior urethral valve on ultrasonography, subsequently reconfirmed on cystourethroscopy.

The Case
A six year old male presented with complaints of straining while micturition, dribbling and sensation of incomplete emptying of urine with nocturnal enuresis for three years. There was no history of operative intervention. Physical examination was unremarkable. Laboratory evaluation revealed normal hemogram, blood urea 50 mg/dl, serum creatinine 1.5 mg/dl. Microscopic examination of urine revealed pus cells and growth in urine culture. Ultrasonography revealed thickened, trabeculated bladder and dilated posterior urethra. The patient was stabilized with urethral catheter and antibiotic cover. Micturating cystourethrogram revealed cystic dilatation in the region of posterior urethra which was placed dorsally resulting in bladder outlet obstruction with large postvoid residual urine volume (Figure 1). Cystourethroscopy revealed a diverticulum in the region of posterior urethra distal to verumontanum. Intra venous urography showed both kidneys normal without any dilatation of upper tract. Diverticulum was approached through midline perineal incision. Cystoscope was advanced to locate extent of lesion thereby facilitating dissection. Dorsally placed diverticulum was rotated ventrally by hooking it with infant feeding tube. Diverticulectomy and urethroplasty was done over the infant feeding tube with the help of vicryl 5-0 suture along with reinforcement of corpous spongiosum (Figure 2). Post operatively patient was voiding with good urinary stream with insignificant post void residual urine. In one year follow up in outpatient department the patient was asymptomatic.

Fig 1: Photograph, MCU showing posterior urethral diverticulum.
Discussion

Urethral diverticulum is defined as saccular enbaginations of urethral mucosa. In contrary to anterior urethral diverticulum, posterior urethral diverticulum is generally acquired in origin. In a review of 95 diverticulae of posterior urethra, only six were of congenital origin. Trauma of different types in an elderly multiparous female give rise to acquired posterior urethral diverticulum.

Etiology for congenital posterior urethral diverticulum is proposed that it is due to faulty or incomplete fusion of a segment of urethral plate. Congenital etiology is supported on the basis of back pressure changes in the urinary bladder without any predisposing factor. Proposed etiopathogenesis of posterior urethral diverticulum is that it can swell up significantly during micturition compressing bladder neck and urethra from posterior aspect resulting in outlet obstruction and its antecedent complications.

We have noticed that posterior urethral diverticulum was deficient of corpous spongiosum. Common differential diagnosis of posterior urethral diverticulum are mullerian duct cyst and utricular cyst. The differentiating point of posterior urethral diverticulum with mullerian duct cyst and utricular cyst are opening which was paramedian and occupies distal to verumontanum in contrast to midline position in the later cysts. Congenital posterior urethral diverticulum may present at birth and in grown up children. Saurabh Agrawal et.al has reported posterior utethral diverticulum even in adult patient. Similar case was reported by Shahram Mousavi et.al in a nineteen year old male patient. Presenting features of posterior urethral diverticulum vary from asymptomatic patient to different features of lower urinary tract symptoms. Some neglected cases may present with calculi, persistent infection and even adenocarcinoma. In most cases of congenital posterior urethral diverticulum, diverticulum is placed ventrally but in the present case it was placed dorsally which is unique.

Conclusion

A high index of suspicion is required in order to diagnose posterior urethral diverticulum especially in bladder outlet obstruction cases. It is always should be kept in differential diagnosis where the diagnosis of posterior urethral valve is made.

References


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