Bilateral Panlobar Nephroblasyomatosis and Bladder Neck Polyp in A Newborn
Bilateral Panlobar Nephroblastomatoz ve Yenidoğanda Mesane Boynu Polipi

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Abstract
Nephroblastomatosis has been described as the existence of numerous or diffuse nephrogenic rests or their derivatives. It is mainly seen in neonates and characterized by multiple bilateral subcapsular masses. We report an unusual case of a term neonate with panlobar nephroblastomatosis. As far as we know, there is no coincidence of panlobar type nephroblastomatosis and bladder neck polyp in the previous literature.

Keywords: Nephroblastomatosis; newborn; bladder polyp

Introduction
Nephroblastomatosis (NB) is described as the presence of multifocal or diffuse nephrogenic rests¹ It occurs mostly in neonates with multiple bilateral subcapsular masses. Nephrogenic rests are foci of metanephric blastema that persists at birth. Nephroblastomatosis predisposes to the development of Wilms tumor.² In a previous study, the prevalence of renal rests (0.87 %) and the incidence of Wilms’ tumor was reported approximately 1 in 10000 infants. Panlobar nephroblastomatosis is an very rare entity.¹ Here, we report a case with bilateral nephromegaly detected prenatally, who had bladder neck polyp and diffuse NB.

Case
A one hour old male infant was admitted to our neonatal
intensive care unit (NICU) with respiratory insufficiency which has started since his birth. A month earlier, an antenatal ultrasound showed enlarged, echogenic, and cystic fetal kidneys. A presumptive diagnosis of autosomal recessive polycystic kidney disease (ARPK) was made. Second pregnancy of his mother had been terminated due to a presumptive diagnosis of ARPK also. He was born to a 24-year-old mother at 38 week’s gestation at a country hospital by cesarean section because of buttock presentation. The mother was G3P2A1. The parents were not consanguineous. The family history was negative for any renal disease. The neonate’s birth weight was 3070 gr. APGAR scores were 1, 3, and 6 at 1, 5, and 10 minutes, respectively. He was supported by using mechanical ventilation in NICU. On physical examination, abdominal distension with bilateral flank masses, bilateral undescended testis and club foot was noted. The faces featured a prominent forehead, a flat nose with a wide bridge, low-set ears, a long upper lip with inverted V-shape. Hematologic and serum biochemical markers were not abnormal. He died of respiratory insufficiency 10 hours after delivery despite NICU care. An autopsy was performed.

Pathological Findings

The lungs were normally situated. Both lungs were composed of two lobes. The right lung weighted 31g and the left lung weighted 29g. Microscopically, lungs were immature with hyaline membrane formation. Severe interstitial hemorrhage was also noted.

The kidneys were normally situated. The right and the left kidney weighted 61g and 84g, respectively. The right kidney was connected by a normal ureter to the bladder. The proximal portion of the left ureter showed a marked dilatation (Fig 1). The lower part of ureter inserted into dome of the bladder. Both kidneys were hyperlobulated. The cut surface of the kidneys had cerebriform appearance and loss of corticomedullary demarcation.

Microscopically, there were numerous foci of nephrogenic rests in the subcapsular areas, in columns of Bertini (Fig 2). Maturing nephrogenic rests consisted of glomeruloid structures and tubular structures lined by columnar epithelium. Mitosis was not determined. Blastemal cells were rare in the nephrogenic rests. Dilated tubules lined by cuboidal or columnar epithelium were shown in some areas (Fig 3). Urinary bladder was normally situated and empty. On cut section, a polyp was shown in the urinary bladder (Fig 4). It was joined with a stalk to the left lateral wall of the urinary bladder.

Discussion

Nephroblastomatosis is a rare congenital pre-neoplastic renal disease with abnormally proliferated renal blastoma. This disease is defined as the presence of diffuse or
multifocal nephrogenic rests and classified into four categories: perilobar, intralobar, combined, and universal. Universal (panlobar) type indicates total replacement of the renal lobe by nephrogenic tissue. Massive renal involvement and pulmonary hypoplasia due to compression of enlarged kidneys make this extremely uncommon disease usually incompatible with life.

Although nephroblastomatosis may be an isolated disease, it has been frequently accompanied with certain teratological disorders. The findings of visseromegaly, bilateral cryptorchidism and low-set ears in our case are similar to the characteristics of Beckwith-Widemann syndrome, but makroglossia, microcephaly and muscular hypertrophy were not observed. The findings in our case may resemble to those of Perlman syndrome, however polyhydramnios, macrosomia, macrocephaly, or hyperplasia of the thymus were not observed in this case. But this case may be a variation of Perlman syndrome of panlobar type. Bladder neck polyp was an unusual finding in our case, which has not been reported before. Recently Gen et al. have suggested that nephroblastomatosis is a syndromic constituent of ischiospinal disostosis, but spinal malformations were not noticed in our case. In view of the fact that, panlobar nephroblastomatosis is a very rare disease, we believe that, our case, together with the finding of a bladder neck polyp, may represent another extremely rare variety of nephroblastomatosis or Perlman syndrome.

References