Complete renal fusion in a child with recurrent urinary tract infection

Fusão renal completa em criança com infecção recorrente do trato urinário

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Abstract
Cake kidney, a rare anomaly of the urinary tract, may be diagnosed at any age range. During the investigation of recurrent urinary tract infection in a 12-year-old child, contrast-enhanced computed tomography demonstrated the presence of a right-sided ectopic kidney, with renal fusion, drained by two ureters. Prophylactic treatment with nitrofurantoin was instituted, and the patient currently remains asymptomatic.

Keywords: Kidney; Malformation; Cake kidney.

Resumo
O rim em bolo é uma rara anormalidade do trato urinário que pode ser diagnosticada em qualquer faixa etária. Durante investigação de infecção urinária recorrente em criança de 12 anos, foi revelada em tomografia computadorizada contrastada a presença de rim direito ectópico, com fusão renal, drenado por dois ureteres. Foi iniciado tratamento profilático com nitrofurantoína e o paciente se encontra assintomático.

Unitermos: Rim; Malformação; Rim em bolo.

INTRODUCTION

Cake kidney is a rare congenital abnormality of the genitourinary tract, with barely more than twenty cases described in the literature[1]. The term “cake kidney” or fused pelvic kidney was defined by Glenn in 1958 as “an abnormality in which all the renal system tissue are fused into a single mass lying at the bottom of the pelvis and which two ureters drain separately into the vesical trigone”[2]. The early diagnosis and the recognition of potential complications associated with such abnormality constitute relevant factors to prevent permanent renal injury[3].

In the present paper, the authors report a case of cake kidney drained by two ureters diagnosed at a tertiary health care unit specialized in urology.

CASE REPORT

A 12-year-old male patient with a history of recurrent urinary tract infections since childhood was referred to the urology unit of a tertiary school hospital for investigation of a mass on right kidney topography discovered on a ultrasonography examination. In his city of origin, the patient had already been hospitalized three times, because of acute pyelonephritis. A previous pelvic ultrasound report brought by the patient demonstrated a mass in the right kidney and absence of the left kidney.

Contrast-enhanced computed tomography of the pelvis was requested and demonstrated right-sided renal ectopia in the lower pelvis and presence of a fused cake kidney (Figure 1) drained by two distinct ureters (Figure 2), without any further alteration.

Considering the possibility of coexisting abnormalities, supplementary imaging
studies were requested but no other alteration was found. Additionally, creatinine testing (0.6 mg/dl) and DMSA renal scan were performed and did not demonstrate renal function compromise.

The patient is currently undergoing prophylactic treatment with nitrofurantoin and has not presented further episode of urinary tract infection and is asymptomatic since.

**DISCUSSION**

Cake kidney is a congenital abnormality defined as a complete fusion of both kidneys, representing only 2% of all renal fusion cases. Such abnormality may be diagnosed at any age range and likewise other renal fusion abnormalities, is most frequently found in men at a 2–3:1 ratio.

Such abnormality occurs at the early phases of the embryonic development. Under normal conditions, two masses of metanephrogenic tissue existing in the lower pelvis develop until taking their definitive positioning in the lumbar region after complex movements involving lateral and ascending migration, axial deflection and internal rotation. It is believed that during the formation of a cake kidney, nephrogenic blastemas are compressed by the umbilical arteries at the beginning of the cranial migration of ureteral buds, which could be the cause of the fusion. The fused kidneys do not ascend as they should in a normal development, remaining in an ectopic pelvic position.

Partial renal fusion is more frequently found, being principally represented by horseshoe kidney and crossed fused renal ectopia. Horseshoe kidneys correspond to 90% of all the renal abnormalities, with an incidence of up to 0.25%.

Anatomically, the cake kidney presents a lobulated anterior aspect while the posterior facet is smooth and homogeneous. The renal pelvis is located anteriorly to the kidney and except for some few cases, there are two ureters which drain into the bladder in the normal anatomical regions of the vesical trigone. Such congenital abnormalities may present some histological alterations, namely: immature glomeruli; cystic changes; widening and dilatation of tubules; or even evidences of chronic renal disease. In other cases, there may be signs of infarct or ischemia secondary to a blood supply abnormality.

This renal fusion abnormality can remain asymptomatic, or even be detected only in an autopsy exam.

The presence of a cake kidney does not indicate a poor prognosis with kidney malfunction or possible progressive deterioration of its function. The follow up is important for the patient for early diagnosis of complications such as: obstruction, calculi, infection, hematuria and uremia. These conditions also can be present in other urinary tract fusions. Besides, ruling out other concomitant congenital abnormalities and perform constant evaluation of the renal function is important to reduce the associated morbidity.

**REFERENCES**