Ectopic Intrathoracic Kidney Along with Right Anterior Diaphragmatic Hernia

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Dear Editor,

Intra-thoracic kidney (ITK) is a very rare congenital disease which is considered as an uncommon type of renal ectopia (1). ITK accounts for 0.1% of the total renal ectopia cases and it is twice as common in men as women. Additionally, ITK in the left hemi thorax is twice as common as on the right (2). Although ITK can be asymptomatic in the most of the cases, it is important to distinguish it from other significant conditions, such as pulmonary masses etc. (3). In this article we report a 5 years old female patient with right sided ITK which presented with chronic respiratory symptoms.

A 5-year-old female was brought to the general surgery clinic with chronic respiratory symptoms including occasional non-productive coughs since birth. The patient was referred to the radiology department and chest X-ray was carried out. There was no history of fever, trauma, surgery or underlying disease (such as renal dysfunction, respiratory distress or gastrointestinal diseases). On the chest X-ray, there was an ill-defined opacity in the lower zone of the right hemi thorax (Figure 1). In the abdominal sonography, there was no right kidney in the anatomical position and pelvis. Also, there was a suspicious view of the diaphragmatic defect in the anterior part of the diaphragmatic muscle in the right hemi-thorax.

Therefore, the patient underwent magnetic resonance urography (MRU), as the best radiation-free method, to confirm the diagnosis of intrathoracic renal ectopia. On MRU, there was a right ITK and intrathoracic intestinal herniation (Figure 2).

Renal development during the embryonic period divides into three stages: pronephros, mesonephros and metanephros (4). During the pronephros, which takes up to four weeks of the embryonic period, primary non-functional kidneys appear. Thereafter, the mesonephros period begins with functional kidneys developing in the embryo which takes up to nine weeks of embryonic period. These kidneys appear around the vertebral column in the form of two ductal structures. After this stage, metanephrin begins with the formation of metanephe-rogenic blastema (which develops into permanent kidneys) and ureteric bud (which develops into collecting systems) in the pelvis and begin to rise in their pathway in retroperitoneum to renal fossa (4). Any defect along this pathway (includes hypo acceleration, over acceleration and metanephrogenic blastema fusion) can lead to ectopic kidneys including pelvic, horseshoe, crossed or thoracic kidney.

Previous research has suggested some predisposing factors such as maternal vitamin A or folic acid deficiency, radiologic exposure, infectious agents in fetal period, teratogens, 18 trisomy and 21 trisomy for developing renal ectopia (5).

The prevalence of this disease has been reported less than one in 1000 births (5). Most of the ectopic kidneys are located in the pelvis or lower lumbar region. ITK is the rarest type of renal ectopia and has been reported one in 10000 cases of renal ectopia (6).

It can be diagnosed in the prenatal and postnatal period, sonography, X-ray, computed tomography (CT) scan, CT urography, magnetic resonance imaging (MRI) and scintigraphy are diagnostic methods for renal ectopia evaluation; however, ultrasound is still at the most cost-effective, non-invasive and reliable method for screening (7).

ITK was first described in 1930 by Campbell (8). The incidence of ITK has been reported 0.5 to 5% of renal ectopia and occurs in less than five in one million live births (9, 10). It is also more common in males and left hemi thorax than females and right hemi thorax. This condition has a strong association with the Buchdalek or Morgani hernia (11). This type of ectopic kidney usually has normal function and its
blood supply can be maintained from the abdominal aorta or the thoracic aorta (12).

When there is a combination of ITK and bowel herniation, the surgical nephropexy and diaphragmatic repair are considered appropriate, but in the case of pure ITK, observation has been recommended (12).

In terms of radiological studies, abnormal shape, intrathoracic origin of vessels, longer ureter, abnormal rotation and medial deviation of lower pole of the kidney have been observed (13).

ITK is classified into four groups based on its features: (1) Pure ITK with normal function, (2) ITK with diaphragmatic eventration, (3) ITK with diaphragmatic hernia which could be congenital (Type A) or acquired (Type B), and (4) Post traumatic diaphragmatic rupture and intrathoracic renal herniation (14).

In this case, there was an uncommon presentation of the ITK. As we discussed above, ITK is more common in males and in left hemi-thorax while in our case it was in a female and the right hemi-thorax. One of the reasons that can be considered for the higher incidence of ITK in the left hemi-thorax can be the presence of the liver on the right which acts as a mechanical barrier against the hyper acceleration of the kidney in embryonic period. An alternative explanation, can be earlier closing of the right hemi thorax diaphragm than the left (15).

Moreover, in this case, there was an anterior diaphragmatic hernia of the intestinal loops. There was a few reports of this coincidence of hernia and ITK but none of the studies reported the incidence of this type of ITK (16).
In conclusion, identifying the ITK subtype, based on radiological studies, is essential for treatment and management planning.

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