A 52-year-old man with a history of sarcoidosis presented with continuous right flank and back pain for four months. Contrast-enhanced CT abdomen and pelvis revealed a lobulated soft tissue lesion extending from the aortic bifurcation to the anterior margin of L5 that measured up to 8 cm in craniocaudal dimension, and contained peripheral ring like foci of calcification. The right kidney was not visualized. Review of a CT performed four years prior demonstrated stability of this lesion. The presumed diagnosis of ectopic kidney was confirmed with a cystoscopy with retrograde pyelography. The patient was counseled that the presence of the ectopic kidney was unlikely to be the cause of his pain. However, due to significant pain and patient anxiety, which was due in part to the delay in diagnosis, the ectopic kidney was surgically removed at the patient’s request. Postoperatively, the patient reported the right flank and back pain was improved for a “short period” but more recently he reported that it was coming back. He is being managed with pain medications, including Tylenol.

The initial contrast-enhanced CT abdomen and pelvis performed for workup of the patient’s abdominal pain demonstrated a lobulated soft tissue lesion at the level of the aortic bifurcation extending inferiorly to the anterior margin of L5 (Figure 1). It was slightly lower in attenuation compared to muscle, measured 2 x 3 x 8 cm (anteroposterior x transverse x craniocaudal), and contained peripheral ring like foci of calcifications. The right kidney was absent. The lesion was also partially visualized on two sequences (T2 and T1 45 minutes post-gadolinium delayed) of an MRI that was performed for further characterization of a gallbladder abnormality. The pelvic lesion demonstrated round T2 hyperintense cystic foci (Figure 2) and diffuse enhancement (Figure 3). The initial differential included neoplastic lesions, such as lymphoma and carcinoid. However, when an old CT abdomen and pelvis performed four years prior became available for review that demonstrated stability of the lesion, a benign etiology for the lesion, such as ectopic kidney, became the presumed diagnosis. In retrospect, a linear configuration of the right adrenal gland was observed on CT (Figure 4).

Cystoscopy with retrograde pyelogram was performed for confirmation of the diagnosis. On cystoscopy, the right ureteral orifice was noted along the right trigonal ridge; however, it was located significantly more distal and medial than would be expected. Retrograde pyelogram demonstrated a small but patent ureter that extended to the level of the lesion (Figure 5).

Dysplastic ectopic pelvic kidney. The diagnosis was confirmed pathologically with intraoperative (Figure 6).
Ectopic pelvic kidneys occur as a result of an abnormal halt in migration of the kidney from the pelvis to its normal location during embryogenesis. The kidney is more often dysplastic in appearance and may be smaller in size than normal with abnormalities in its shape and rotation. Additionally, because the kidney is not present in the renal fossa, rather than taking on its more common Y or V shaped appearance, the adrenal gland takes on a more globular or linear appearance, commonly referred to as the lying down or pancake adrenal gland. Blood supply is derived from neighboring vessels, and the renal artery and vein may not be seen at the level of the empty renal fossa. The ureter may also insert in an ectopic location.

Renal ectopia is asymptomatic in most patients. The ectopic kidney may function normally or have limited to no function at all. Ectopic kidneys are more disease prone than normal kidneys. They are more vulnerable to obstruction, vesicoureteral reflux, infection, calculi, renal vascular hypertension, and trauma. When a patient requires imaging of the abdomen and pelvis, the ectopic kidney may be mistaken for a retroperitoneal mass. The diagnosis of ectopic kidney is made with the use of contrast-enhanced CT imaging, MRI, or a retrograde pyelogram.
experiences symptoms from pathology within the ectopic kidney, it may present a diagnostic dilemma, as the presentation is atypical. Chronic obstruction may lead to cystic dysplasia. Multicystic dysplastic kidney may also be seen in an ectopic kidney.\(^5\)

Asymptomatic ectopic kidneys are managed conservatively; however, surgical management may be needed in the setting of refractory stones or infection.\(^5\)

There is an increased incidence of genital tract abnormalities in the setting of renal ectopia, such as hypospadias and cryptorchidism in males and agenesis of the uterus and vagina, or unicornuate uterus in females. Additionally, renal ectopia may also be associated with spinal abnormalities.\(^6\)

**CONCLUSION**

In a patient with an empty renal fossa, it is more common for the patient to have an ectopic kidney rather than renal agenesis. Ectopic kidneys may be dysplastic in appearance with variable size, shape, and rotation. It is important to be aware of the possibility of having an ectopic kidney and that it may have an atypical appearance to avoid misdiagnosis and potentially unnecessary surgery or biopsies.

**REFERENCES**


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