

## Bilateral Ectopic Pelvic Kidneys: The Novel Association with Liver Hemangioma

Aamir Jalal Al-Mosawi<sup>1</sup>

<sup>1</sup>Baghdad Medical City and the National Training and Development Center Iraqi ministry of Health Baghdad, Iraq

### \*Corresponding Author

Aamir Jalal Al-Mosawi, Baghdad Medical City and the National Training and Development Center Iraqi ministry of Health Baghdad, Iraq.

Submitted: 14 Dec 2022; Accepted: 30 Dec 2022; Published: 22 Jan 2023

**Citation:** Aamir Jalal Al-Mosawi, (2023). Bilateral Ectopic Pelvic Kidneys: The Novel Association with Liver Hemangioma, *World J Clin Med Img*, 2(1), 11-13.

### Abstract

#### Background

Renal ectopia is a congenital abnormality in which one or both kidneys are located in an unusual position because of failure of normal ascend from its origin in the true pelvis. Several types of renal ectopia with or without fusion or other associated renal and abdominal visceral abnormalities have been reported. We have previously reported the case forty-one of crossed unfused renal ectopia, and the aim of this paper is to report the association of bilateral ectopic pelvic kidneys with liver hemangioma.

#### Patients and methods

The case of a woman in her mid-forties with renal and hepatic abnormalities on pelvic and abdominal ultrasound was studied.

#### Results

A woman in her mid-forties was having recurrent lower abdominal pain or discomfort and recurrent urinary tract infection. Ultrasound showed that both kidneys were ectopic and were located in the pelvic cavity. Both kidneys were normal in size, texture, parenchymal thickness, and had normal cortico-modularly differentiation. There was mild splitting of the pelvi-calyceal system suggesting infection. Both ureters were normal. The liver was normal in size with homogenous texture, but there was a right lobe well-defined echogenic soft tissue mass, hemangioma. Renal function tests showed normal findings.

#### Conclusion

Bilateral ectopic pelvic kidneys is a rare congenital condition that has not been reported to occur in association with hepatic hemangioma. This paper reports the novel association of bilateral ectopic pelvic kidneys with hepatic hemangioma.

**Keywords:** Bilateral ectopic pelvic kidneys, hepatic hemangioma.

### Introduction

Renal ectopia is congenital abnormality in which one or both kidneys are located in an unusual position because of failure of normal ascend from its origin in the true pelvis. Several types of renal ectopia with or without with or without fusion or other associated renal and abdominal visceral abnormalities have been reported since the early report of Polk in 1882 [1, 2, 3]. We have previously reported the case forty-one of crossed unfused renal ectopia [1], and the aim of this paper is to report the association of bilateral ectopic pelvic kidneys with liver hemangioma.

### Patients and methods

The case of a woman in her mid-forties with renal and hepatic abnormalities on pelvic and abdominal ultrasound was studied.

### Results

A woman in her mid-forties was found to have renal and hepatic abnormalities on pelvic and abdominal ultrasound during the evaluation after treatment for urinary tract infection. She was having recurrent lower abdominal pain or discomfort and recurrent urinary tract infection. Ultrasound (Figure-1) showed that both kidneys were ectopic, and were located in the pelvic cavity. Both kidneys were normal in size, texture, parenchymal thickness, and had normal cortico-modularly differentiation. There was mild splitting of the pelvi-calyceal system suggesting infection. Both ureters were normal. The liver was normal in size with homogenous texture with a right lobe well-defined echogenic soft tissue mass, hemangioma. She had normal blood pressure and renal function tests showed normal findings.



**Figure 1:** Ultrasound showed that the both kidneys were ectopic and were located in the pelvic cavity

### Discussion

As early as 1931 Howard L Tolson emphasized that ectopic kidneys can be structurally and functionally normal and remain asymptomatic or associated with mild dull pain and urinary tract infections [4].

Boujnah et al (1989) from Tunisia emphasized that bilateral pelvic ectopic kidneys is a rare congenital condition, and they reported 50 cases of pelvic ectopic kidneys observed during 12 years. 47 patients had unilateral pelvic ectopic kidney, two patients had bilateral pelvic ectopic kidneys, and one patient had pelvic ectopic kidney solitary pelvic ectopic kidney. Eighteen patients had healthy ectopic kidneys. Thirty-two patients had diseased ectopic kidney including twenty-one patients with renal stones and seven patients with uretero-pelvic junction disease.

Boujnah et al from Tunisia emphasized the diagnostic value of renal ultrasound which can help in avoiding other useless and expensive investigations [5].

Hirano and colleagues (1992) reported the incidental detection of bilateral ectopic pelvic kidneys by radioisotope angiography [6].

Alonso Domínguez (1996) also emphasized the rarity of bilateral pelvic ectopic kidneys and reported a case with an unusual presentation [7].

Gokalp and colleagues (2010) reported a case with bilateral ectopic kidneys associated with vascular anomaly and hypertension and renal dysfunction [8].

Hemangiomas including liver hemangiomas are benign vas-

cular tumors that are generally observed during infancy. Liver hemangioma is the most common benign tumor of the liver [8, 9]. Brodsky et al (1987) reported that during the performance of high-resolution real-time abdominal sonography, hemangioma small echogenic hepatic masses are frequently discovered [10].

Lipman and Tumei (1990) from Boston, Massachusetts emphasized that the diagnosis of small liver hemangiomas, less than 3 cm can be made with abdominal ultrasound study [11].

Al-Durazi et al (2003) from Bahrain reported the incidental finding of a case of hepatic hemangioma while performing a routine renal and pelvic ultrasound in patients with urinary retention associated with benign prostatic hyperplasia [12].

Mungovan and colleagues (1994) emphasized that the size of the majority of hepatic hemangiomas don't increase in size for months and years, and an increase in size demands further evaluation [13].

### Conclusion

Bilateral ectopic pelvic kidneys is a rare congenital condition that has not been reported to occur in association with hepatic hemangioma. This paper reports the novel association of bilateral ectopic pelvic kidneys with hepatic hemangioma.

### References

1. Al-Mosawi, A. J. (2020). The Case Forty-One of Crossed Unfused Renal Ectopia. *International Journal of Recent Innovations in Medicine and Clinical Research* (ISSN: 2582-1075), 2(2), 14-18.
2. Cullen, T. S. (1910). Total absence of the vagina and uterus; right pelvic kidney; absence of the left kidney; the tubes and

- ovaries on both sides in the inguinal canal. *Am. J. Obstet. and Gynecol.*, NY, 62, 296.
3. Looney, W. W., & Dodd, D. L. (1926). An ectopic (pelvic) completely fused (cake) kidney associated with various anomalies of the abdominal viscera. *Annals of surgery*, 84(4), 522.
  4. Tolson HL. Ectopic (Pelvic) Kidney. *Ann Surg* 1931 Apr; 93(4):880-5. Doi: 10.1097/0 0000658-193104000-00011.
  5. Boujnah, H., Abid, I., Moalla, N., & Zmerli, S. (1989, January). Pelvic kidney. Apropos of 50 cases. In *Annales D'urologie* (Vol. 23, No. 1, pp. 11-16).
  6. Hirano T, Igarashi H, Mogi Y. Bilateral ectopic pelvic kidneys incidentally demonstrated by radioisotope angiography. *Clin Nucl Med*. 1992 Oct; 17(10): 831.
  7. DOMINGUEZ, F. (1996). Riñón ectópico pelviano bilateral. *Archivos españoles de urología*, 49(9), 977-978.
  8. Gokalp, G., Hakyemez, B., & Erdogan, C. (2010). Vascular anomaly in bilateral ectopic kidney: a case report. *Cases Journal*, 3(1), 1-4.
  9. Al-Mosawi, A. J. (2022). Esquirol-Séguin-Down Syndrome Associated with Hepatic Hemangioma: An Association not Previously Reported in the Literature.
  10. Al-Mosawi, A. J. (2022). Uncomplicated Cutaneous Infantile Strawberry Hemangioma: Educational Images and Evidence-Based Recommendation.
  11. Brodsky, R. I., Friedman, A. C., Maurer, A. H., Radecki, P. D., & Caroline, D. F. (1987). Hepatic cavernous hemangioma: diagnosis with <sup>99m</sup>Tc-labeled red cells and single-photon emission CT. *American Journal of Roentgenology*, 148(1), 125-129.
  12. Lipman, J. C., & Tumeh, S. S. (1990). The radiology of cavernous hemangioma of the liver. *Critical Reviews in Diagnostic Imaging*, 30(1), 1-18.
  13. Al-Durazi, M. H., Al-Helo, H. A., Al-Reefi, S. M., Al-Sanaa, S. M., & Abdulwahab, W. A. (2003). Routine ultrasound in acute retention of urine. *Saudi medical journal*, 24(4), 373-375.
  14. Mungovan, J. A., Cronan, J. J., & Vacarro, J. (1994). Hepatic cavernous hemangiomas: lack of enlargement over time. *Radiology*, 191(1), 111-113

**Copyright:** ©2023: Aamir Jalal Al-Mosawi. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.