Hematuria after endoscopic ultrasound-guided fine needle aspiration of a renal tumor in von Hippel–Lindau disease

A 35-year-old man presented at our hospital with weight loss and diarrhea. Computed tomography (CT) enteroclysis was carried out for evaluation of the small bowel to investigate the diarrhea. The scan revealed bilateral renal cysts with complex renal masses, as well as multiple pancreatic cysts and a mass in the genu (Fig. 1). The patient was suspected as having von Hippel–Lindau (VHL) disease, which was confirmed after a CT brain revealed a cerebellar hemangioblastoma.

Endoscopic ultrasound (EUS) evaluation of the pancreas showed multiple pancreatic cysts between the head and the tail (Fig. 2) with an 18 × 12 mm, well-defined hyper-echoic mass in the genu. Both kidneys were studded with cysts. In addition, a hyper-echoic mass (12 × 15 mm) was seen in the right kidney. We carried out EUS-guided fine needle aspiration (FNA) of the pancreatic mass, followed by the right renal mass (Video 1).

However, shortly afterward, the patient passed blood-stained urine, and hematuria was confirmed with a dipstick. As it was mild, the patient was managed conservatively, and the hematuria resolved several hours later. The renal aspirate revealed a renal cell carcinoma (RCC), while the pancreatic aspirate revealed a neuroendocrine tumor. The patient was offered surgery, but he declined.

VHL disease is an autosomal dominant condition characterized by tumors involving multiple organs [1]. We report the first case of EUS-guided FNA of a renal mass in VHL disease resulting in hematuria. This complication has never been documented before, and there are only four reports of uncomplicated EUS-guided FNA of renal masses [2–5]. The risks associated with percutaneous biopsy of renal lesions include hematoma, hematuria, pneumothorax, and needle tract seeding. Endoscopists should be aware of these complications because a renal mass may be accessible only by traversing through the parenchyma. Our case also illustrates the usefulness of EUS in the evaluation of both pancreas and kidneys in VHL disease.

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**References**


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