Vesicourachal Diverticulum: An Uncommon Incidental Finding on Staging $^{18}$F-FDG PET/CT in a Patient with Suspected Malignant Transformation of Neurofibromatosis

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In a 32-y-old man with neurofibromatosis type 1, $^{18}$F-FDG PET/CT incidentally revealed a vesicourachal diverticulum, a rare anatomic variant. The PET/CT, performed for staging a malignant peripheral nerve sheath tumor, highlighted a distinctive $^{18}$F-FDG–avid pattern crucial for accurate diagnosis. Recognizing such features enhances disease assessment and clarifies distinctions between benign urogenital anomalies and malignancies in $^{18}$F-FDG PET/CT staging.

Key Words: $^{18}$F-FDG PET/CT; vesicourachal diverticulum; CT

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The urachus connects the fetal bladder to the umbilicus, aiding waste removal. Second-trimester descent obliterates it, forming the median umbilical ligament. Occasionally, incomplete closure leads to urachal anomalies, affecting 1 in 5,000 adults (1). A vesicourachal diverticulum, a rare variant, manifests as an outpouching from the bladder to the umbilicus and is often asymptomatic. Identified uniquely on $^{18}$F-FDG PET/CT because of tracer excretion, it appears as an $^{18}$F-FDG–avid, fluid-filled lumen. Notably, this finding, usually associated with malignant transformation, stands out on imaging (2). In contrast, neurofibromatosis type 1, a genetic disorder causing neurologic tumors, rarely involves the bladder, with only 3 reported cases (3). In a single case, a neurofibroma originating from a urachal mass was noted, highlighting the exceptional nature of these associations (4). For the current case study, the need for informed consent was waived by the institutional review board (waiver form 118).

CASE REPORT

A 32-y-old man with neurofibromatosis type 1, who had a prior neurofibroma resection in his right lower extremity, presented with a rapidly enlarging mass in the same area. CT revealed a heterogeneous lesion in the right femur, and MRI displayed a T1-isointense, T2-weighted short-tau inversion recovery hyperintense lesion. Additionally, numerous T2-hyperintense neurofibromas were seen along the course of the sciatic nerve in this known case of neurofibromatosis (Fig. 1). Biopsy confirmed a malignant peripheral nerve sheath tumor. $^{18}$F-FDG PET/CT showed intense uptake in the tumor and a linear structure in the anterior lower abdomen, indicating physiologic tracer excretion into a vesicourachal diverticulum (Figs. 2 and 3A–C). CT confirmed a fluid-filled structure in the anterior lower abdomen, extending to the umbilicus (Fig. 3D). These findings emphasize the diagnostic significance of multimodal imaging in neurofibromatosis-related malignancies and the unique identification of a vesicourachal diverticulum on PET/CT.

DISCUSSION

A vesicourachal diverticulum, an outpouching from the bladder to the umbilicus, is often asymptomatic but may present complications such as malignancies or infections. On $^{18}$F-FDG PET/CT, it appears as an avid, fluid-filled tract. The association of neurofibromatosis type 1 with the bladder, particularly originating from a urachal mass, is rare (3,4).

Physiologic $^{18}$F-FDG excretion in the urinary system results in visualized activity in the kidneys, renal pelvis, and bladder. A patent urachus or vesicourachal diverticulum, continuous with the bladder lumen, can appear $^{18}$F-FDG–avid. Distinguishing features from malignancy include the location of the tracer uptake, a concurrent hypoplastic bladder, and the absence of a focal mass. This case highlights the unique appearance of vesicourachal diverticulum on $^{18}$F-FDG PET/CT.

FIGURE 1. Coronal anterior (A) and posterior (B) MR images of right femur. Heterogeneous, hyperintense lesion (arrow) in right proximal thigh is seen with mild adjacent edema. Additionally, numerous T2-hyperintense neurofibromas are seen along course of sciatic nerve (arrowheads).

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CONCLUSION

In $^{18}$F-FDG PET/CT disease staging, recognizing metabolically active tissue is crucial for assessing severity. This case emphasizes discerning urogenital anomalies during interpretation, enhancing the distinction between benign anatomic variants and malignancies.

DISCLOSURE

No potential conflict of interest relevant to this article was reported.

REFERENCES