Myopericytoma in urinary bladder: a case report

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Abstract

Background: Myopericytoma is reported to occur mainly in the skin and superficial soft tissue of the extremities. In contrast, occurrence in the urinary bladder is extremely rare.

Case presentation: We describe a 75-year-old Japanese man who developed a submucosal tumor at the right trigone of his bladder that led to interference with the discharge of right ureteral calculus. No invasive growth was observed by magnetic resonance imaging. Transurethral resection was successfully performed; histopathological analysis revealed perivascular proliferation of spindle-shaped to oval-shaped, cytologically bland tumor cells with eosinophilic cytoplasm. On immunohistochemical examination, the tumor cells were positive for alpha-smooth muscle actin, desmin, CD34 and h-caldesmon.

Conclusion: Cystoscopic and pathological findings were compatible with a diagnosis of myopericytoma of the urinary bladder.

Keywords: Myopericytoma, Bladder tumor, Submucosal tumor

Background

Mesenchymal tumor in the bladder accounts for less than 1 % of all bladder neoplasms [1]. Although benign mesenchymal tumors are also rare, leiomyoma is the most common, followed by hemangioma, solitary fibrous tumor, and neurofibroma [1–3].

Myopericytoma is a benign mesenchymal neoplasm composed of cytologically uniform presentation ranging from oval to spindle-shaped cells with a myoid appearance showing multilayered, perivascular concentric growth [2, 3]. Myopericytoma is reported to occur typically in skin and superficial soft tissue, commonly in the distal and proximal extremities, and occasionally in the trunk, head, and neck. In all locations the mass is solitary, painless, and slow growing [2–10]. In rare cases, the tumor is found in the oral cavity, nasal cavity, and thorax [2–10]. By contrast, occurrence in visceral organs such as the lungs, heart, gastrointestinal tract, and brain is extremely rare [2–10]. Furthermore, only 11 cases (ten cases in the kidney and one case in urinary bladder) in the urinary tract have been described [4–10].

In the present case report, we review our experience with a case of myopericytoma occurring in urinary bladder, and report the clinicopathological features.

Case presentation

A 75-year-old Japanese man was referred for sudden asymptomatic gross hematuria. Ultrasonography and abdominal X-ray suggested bladder calculus, and computed tomography (CT) was performed. CT showed intramural right ureteral calculus (1.8 cm in diameter) and a well-circumscribed hypervascular mass measuring approximately 2.1 cm at the right trigone of his urinary bladder (Fig. 1b). Hydronephrosis was absent in bilateral kidneys. We then performed a cystoscopy to confirm the bladder mass. Examination revealed a non-papillary nodular mass located at the right trigone (Fig. 1a). The mass appeared to be completely covered by normal urothelium, suggesting a submucosal tumor. The mucosa was elevated behind the mass; however, we were unable to confirm the right ureteral orifice and calculus preoperatively. Since
CT showed the mass to be adjacent to the calculus, the tumor may have interfered with the discharge of the calculus. No invasive growth was observed by magnetic resonance imaging (Fig. 1c). Urine cytology was negative for malignant cells. We then performed transurethral resection (TUR) of the mass. Following resection, we confirmed right ureteral orifice and intra-ureteral calculus. As a result, we performed transurethral lithotripsy. Our patient recovered without perioperative complications and has remained free of recurrence for 5 months.

Pathological features
On gross examination, the resected surface of the mass specimens was solid, firm, and tan-white to yellow without apparent necrosis and hemorrhage.

On histologic examination, the urothelium was largely denuded; however, residual urothelium showed no atypia. In the submucosal lesion, we observed spindle-shaped to oval-shaped cells that had mildly hyperchromatic nuclei and eosinophilic elongated or tapering cytoplasm arranged in loose fascicles or perivascular whorls around often-hyalinized blood vessels or in a vague storiform-growth fashion, embedded in variably fibromyxoid or edematous stroma associated with dilated thin-walled blood vessels (Fig. 2a, b). Mitotic figures are rare. On immunohistochemical examination, the tumor cells were positive for alpha-smooth muscle actin (SMA), desmin, CD34, and h-caldesmon, and CD99 was weakly and focally expressed (Fig. 2c-e). The cells were negative for S-100, CAM5.2, AE1/AE3, epithelial membrane antigen (EMA), STAT6, MUC4, and claudin-1.

Discussion
Myopericytoma is defined as a member of the perivascular tumor family, which includes myofibroma, angioleiomyoma, glomus tumor, and myopericytoma, by the World Health Organization (WHO) classification (of tumor of soft tissue and bone) [2]. Myopericytoma occurs in various age groups, reported as ranging from 10 to 87 years; however, most are seen in adults with a median age of 49 years [3]. Regarding gender, a slight male predominance was reported in a male to female ratio of 1.25:1 [3]. As mentioned above, the tumor generally arises in dermal and subcutaneous tissue, and occurrence in visceral or intracranial sites is extremely rare [2–10]. To the best of our knowledge, the current case is the second report in the English literature [5]. Compared with a previous report by Zhao et al. [5], the majority of clinical features are similar, except for gross hematuria. In our case, the position of the tumor relative to the ureteral calculus was very unusual. Because the tumor was located immediately adjacent to the right ureteral orifice, it may have interfered with the discharge of the calculus. We speculated that the gross hematuria might have been the result of mucosal injury caused by the calculus. However, the results of cystoscopic and enhanced CT were nearly identical in both cases.
On histopathologic examination, morphological features were consistent with myopericytoma. On immunohistochemical examination, the majority of the findings were also consistent; however, unusual intense immunoreactivity was observed in CD34. Although, myopericytoma is reported to be negative for CD34 in most cases, one case of strong and three cases of partial reactivity have been reported in the kidney [6, 7, 9].

Myopericytoma shares several morphologic and immunohistochemical features with myofibroma, angioleiomyoma, and glomus tumor [2, 3]. However, each tumor has distinct characteristics. Myofibroma is characterized by a multinodular and biphasic growth pattern of peripherally located mature spindle-shaped myoid cells arranged in bundles, and centrally located immature-appearing oval-shaped tumor cells associated with branching thin-walled vessels, which are the most significant appearance for differential diagnosis [2, 7]. Angioleiomyoma shares the appearance of concentric perivascular growth pattern with myopericytoma; however, angioleiomyoma tends to lack the distinctive thick-walled vessels and shows negative or focal reactivity for desmin immunohistochemically [2, 7].

Myopericytoma is generally considered a benign tumor with favorable clinical course, although local recurrence has been reported in a small number of cases [3–10]. However, a few cases of malignant myopericytoma have been reported [3–10]. The proposed pathological criteria for malignancy are reported as deep-seated location, infiltrative growth pattern, nuclear pleomorphism, high mitotic activity, and presence of tumor necrosis [7, 9]. In these reports, six of eight patients with malignant myopericytoma developed metastasis, and three patients died of the disease within 1 year [9]. The majority of these malignant tumors occurred in extremities, and one case each of neck, superior mediastinum, and left atrium were involved [9]. No recurrence or metastasis has been reported in cases occurring in the urinary tract [9]. In our case, cystoscopy and CT showed that the patient has remained free from recurrence for 5 months.

**Conclusions**

While myopericytoma commonly occurs in skin and superficial soft tissue, it is extremely rare in the urinary bladder. We treated a 75-year-old Japanese man with submucosal tumor adjacent to the right ureteral orifice. The tumor may have interfered with the discharge of ureteral calculus. Cystoscopic and pathological findings prompted the conclusion that the tumor was the urinary bladder counterpart of myopericytoma.

Abbreviations
CT: Computed tomography; EMA: Epithelial membrane antigen; SMA: Smooth muscle actin; TUR: Transurethral resection; WHO: World Health Organization

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Authors’ contributions
TN drafted the report, performed the surgical procedure, and approved the final version of the manuscript. TK, KI, MF, and HT performed examinations, cared for the patient, and approved the final version of the manuscript. YA and HK reviewed the pathological specimens, and approved the final version of the manuscript. SM and TK drafted the report and contributed to the final version of the manuscript. All authors read and approved the final manuscript.

Competing interests
The authors declare that they have no competing interests.

Consent for publication
Written informed consent was obtained for publication of this case report and accompanying images from the patient. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

Ethics approval and consent to participate
Not applicable.

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