CASE REPORT

Blastoid variant of mantle cell lymphoma of the female urethra mimicking a caruncle: A rare but highly aggressive subtype case with literature review

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Summary
Primary urethral lymphoma is a rare entity without a standardized treatment protocol. We report a case of an elderly woman presenting with a caruncle associated with vaginal spotting and intermittent dysuria. She underwent surgical excision of the lesion. Histological analysis revealed a blastoid variant of mantle cell lymphoma, a previously unreported subtype. The patient received chlorambucil assisting a rapid local disease progression. She died of disseminated disease 6 months after diagnosis. A review of the lymphomas of the urethra is included.

KEY WORDS: Caruncle; Lymphoma; Urethra; Genitourinary; Mantle cell lymphoma; Blastic variant.

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INTRODUCTION
Non-Hodgkin’s genitourinary lymphoma as a primary extra nodal invasion is a rare condition, with few reported cases affecting the urethra (1). The most common subtypes described are mucosa-associated lymphoid tissue-type (MALT) lymphoma and diffuse large B-cell lymphoma. Herein, we report the first documented case of a blastoid variant of mantle cell lymphoma resembling a caruncle. Furthermore, to shed light on clinical features of primary urethral lymphoma, we reviewed previously reported cases.

CASE REPORT
A 84-year-old woman presented with a recent onset, fast-growing genital node associated with vaginal spotting and intermittent dysuria. Her medical history was non-contributory, lacking fever, weight loss or any other systemic manifestations. There were no enlarged lymph nodes or organomegalies upon physical examination. Pelvic examination revealed a urethral caruncle arising from the posterior wall of the urethral meatus (Figure 1a, b). It was characterized by an erythematous appearance, firm at palpation, measuring approximately 4 cm at its maximum diameter. The patient underwent cystourethroscopy with a 21F cystoscope revealing no bladder involvement, and surgical excision of the urethral caruncle. Histologic examination revealed a urethral mucosa characterized by squamous metaplasia, subjigated by a widespread growth pattern of lymphoid proliferation, consisting of medium-sized, blastic-shaped, B-cell phenotype elements. Immunohistochemically, these cells were positive for CD20, CD5, Cyclin D1 and Bcl-2, negative for CD23, CD10, BCL6, MUM1, CD3, CD30, TdT and cytokeratin. The Ki-67 index was approximately 90%. A diagnosis of a primary blastoid variant of mantle cell urethral lymphoma was made. A chest, abdomen and pelvis contrast enhanced computed tomography (CT) was performed for the staging workup and revealed non- mediastinal lymphadenopathy, no focal lung lesions, a 2.5-mm angioma located at the right lobe of the liver, normal spleen, pancreas and kidneys, and no abdominal or pelvic lymphadenopathy. 18F-fluorodeoxyglucose positron emission tomography (FDG-PET) showed abnormal FDG uptake present in the external genitalia and vagina.

The patient was then referred to the Oncohematology Department and underwent two cycles of chlorambucil, despite this a rapid local disease progression was seen (Figure 1c). The patient died of disseminated disease six months after diagnosis.

DISCUSSION
Caruncles commonly affect the postmenopausal female urethra, appearing as erythematous nodules at the posterior lip of the urethral meatus. This lesion is considered to be neither neoplastic nor preneoplastic, being composed of chronic inflammatory cells, dilated vessels and hyperplastic epithelium. In light of this, an incorrect diagnosis of lymphoma or sarcoma may occasionally be made based on the presence of bizarre stromal cells or lymphoid infiltrate (2). On the other hand, 2.3% of the caruncles were found to be malignant tumors upon histopathological analysis, usually a carcinoma or a Bowen’s disease (2). In this context, a caruncle-like primary lymphoma is exceptionally rare. Since 1949, only 29 patients with urethral lymphoma have been reported (1-2), including the present case, with an age distribution ranging from 31 to 90 years old. Of these, a female
predominance has been shown, with 75.9% of cases affecting women. When the female urethra is involved, a caruncle-like aspect is the most common presentation, having been reported in 9 cases. Other associated symptoms include spotting, hematuria, dysuria and vulvar pruritis. Cancer-specific mortality is usually relegated to a disseminate disease at diagnosis, whereas the overall 4-year survival rate of caruncle-like tumors is 55.5%.

Each of the previously reported 29 cases involved non-Hodgkin’s lymphoma, with the most common subtypes being mucosa-associated lymphoid tissue-type lymphoma and diffuse large B-cell lymphoma. Treatments include excision, radiotherapy and chemotherapy. To our knowledge, this is the first blastoid variant of mantle cell caruncle-like lymphoma to be reported. According to Bernard et al. (3), the blastoid variant form of mantle cell lymphoma is considered to be a highly aggressive subtype of non-Hodgkin’s lymphoma with 46% of patients who do not respond to treatment dying quickly due to disseminated disease.

The overall mortality rate is 66% with a 14.5-month median survival time. Reported frontline therapy includes CHOP-like regimen or chlorambucil.

The blastic variant of mantle cell lymphoma is usually diagnosed upon initial presentation and more rarely in the course of common forms. Patients are predominantly male, over 60 years of age, and one third have B symptoms.

The majority of patients have an Ann Arbor stage IV disease (85%), lymphadenopathy (82%) and extranodal involvement (66%), especially in the lung and pleural cavities, more rarely with a gastro-intestinal infiltration. Neither Ann Arbor staging nor bone marrow or blood involvement have been found to influence clinical outcome and there is currently no standard therapy for this disease.

CONCLUSIONS

In conclusion, the correct recognition of a primary lymphoma of the urethra is of clinical importance and should be considered in the differential diagnosis of a caruncle. Being the extranodal lymphoid infiltrate a diagnostic dilemma, it is of utmost importance that the pathologist has expertise concerning these tumors. An universally accepted treatment scheme is needed, and for subtypes of lymphoma with a less favorable prognosis an aggressive treatment strategy is strongly recommended.

REFERENCES


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