

A Rare Cause for Hematuria: Lymphoma of the Urinary Bladder

Abstract

Primary lymphomas of the urinary bladder are rare and majority present with hematuria. Patients are usually elderly females with a history of recurrent cystitis. The diagnosis is clinched after imaging and biopsy of suspicious lesions in the bladder. The prognosis is good irrespective of the modality of treatment. With this background, we report a case of primary marginal zone lymphoma of the urinary bladder (Stage I_E) in a young lady with no history of cystitis, who presented with isolated hematuria. After treatment with radiotherapy alone, patient is in complete remission for the past 18 months. The timely diagnosis in a patient with hematuria holds the key to successful treatment and its importance cannot be overemphasized.

Keywords: Hematuria, marginal zone lymphoma, mucosa-associated lymphoid tissue, primary lymphoma, radiotherapy, urinary bladder

**Sherin P Mathew,
Sonia Parikh,
Asha Anand,
Vaibhava Srivastava**

*Department of Medical
Oncology, Gujarat Cancer and
Research Institute, Ahmedabad,
Gujarat, India*

Introduction

Primary lymphomas of the urinary bladder are rare, representing 0.2% of all known bladder cancers.^[1] The most common symptoms of bladder lymphomas are macroscopic hematuria, dysuria, and increased frequency of micturition.^[2] In the literature reported so far, patients are usually elderly and have a history of chronic cystitis. Herein, we report a case of a young female with no such history, who is in complete remission after treatment.

Case Report

A 35-year-old lady with no comorbidity presented to the urologist with 1-month history of painless hematuria. On physical examination, no pallor, lymphadenopathy, hepatosplenomegaly, or any mass in the pelvis was detected. The urine routine and microscopic examination was confirmative of hematuria. Thereafter, a contrast-enhanced computed tomography (CECT) of the abdomen and pelvis showed a large, lobulated mass, 96 mm × 46 mm × 63 mm, in the posterolateral wall of the urinary bladder with few subcentimeters internal iliac and left obturator lymph nodes. The mass was encasing the left vesicoureteral junction causing mild proximal hydronephrosis [Figures 1 and 2].

Subsequently, cystoscopy showed a well-defined, lobulated submucosal mass arising from the posterior wall from which biopsy was taken. Histopathological examination of the biopsy specimen revealed infiltration of the bladder mucosa by sheets of monomorphic population of small, atypical lymphocytes, suggestive of non-Hodgkin's lymphoma [Figure 3].

The immunohistochemistry was positive for leucocyte common antigen and also for CD20, [Figure 4] Bcl2, CD5, CD43, CD68, and MPO. About 40% of cells stained positive for MIB-1 [Figure 5]. CD10 and cyclin D1 were negative, consistent with marginal zone B-cell lymphoma (MZL) with a high proliferative index.

The patient had a hemoglobin of 11.5 g/dl and total leukocyte count of 5500/mm³ with 78% neutrophils and platelet count of 3.5 lakhs/mm³. Her lactate dehydrogenase was 139 U/L. Bone marrow aspiration and trephine bone biopsy did not show infiltration by lymphoma and CECT of the neck and thorax showed no disease elsewhere. Hence, the final diagnosis of primary MZL of the urinary bladder stage I_E was confirmed.

In view of localized disease, the patient was treated with curative intent, by means of radiation therapy of 50 Gy in 25 fractions. She tolerated the therapy well with no major side effects following which the hematuria subsided. A repeat CECT of

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Address for correspondence:

*Dr. Sherin P Mathew,
Department of Medical
Oncology, Gujarat Cancer
and Research Institute,
Ahmedabad, Gujarat, India.
E-mail: sherin.p.mat@gmail.
com*

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Figure 1: Contrast enhanced computed tomography of the abdomen and pelvis showing the tumor arising from the posterolateral bladder wall – Coronal section

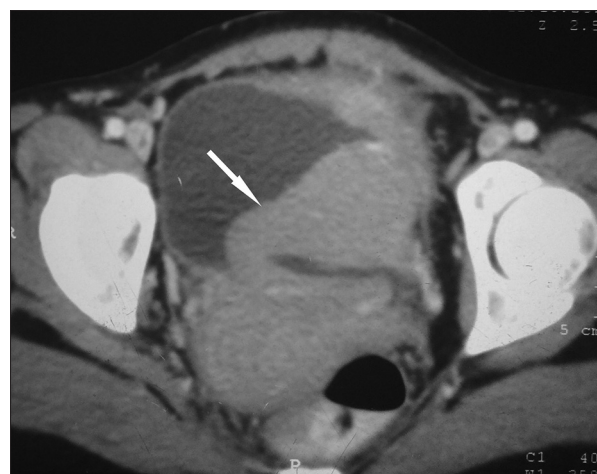


Figure 2: Axial section of the same contrast enhanced computed tomography showing the tumor

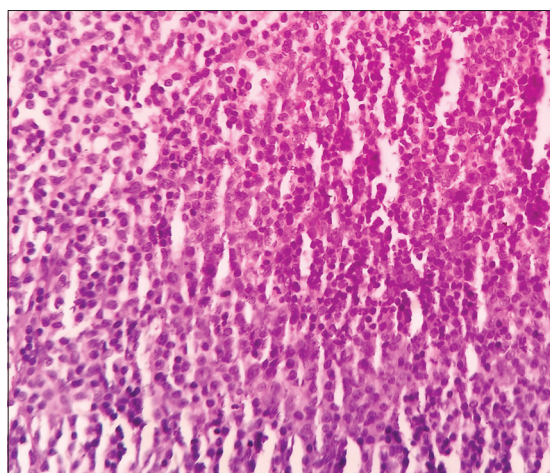


Figure 3: Atypical lymphocytes infiltrating the submucosa of the urinary bladder (H and E, $\times 40$)

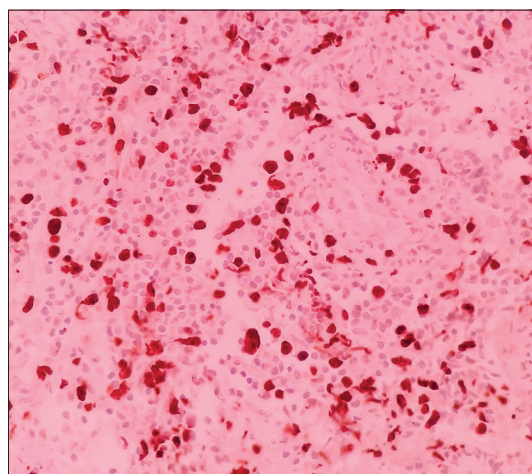


Figure 5: MIB-1 positivity among 40% of tumor cells ($\times 40$)

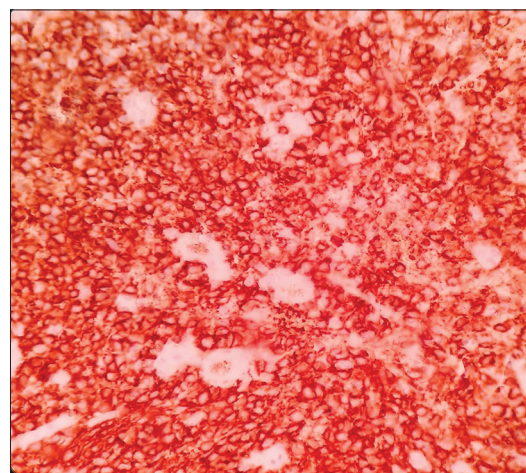


Figure 4: The tumor cells staining for CD20 on immunohistochemistry ($\times 40$)

After 2 months, it had reduced to a size of 17 mm \times 35 mm, and cystoscopic guided biopsy was negative for malignancy. She is under observation with CECT pelvis and urine cytology now. Her recurrence-free survival is 18 months.

Discussion

Since the first report of primary lymphosarcoma of the urinary bladder by Eve in 1885,^[2] much has come to light about this rare lymphoma. It was classified along with sarcomas and the treatment has evolved from surgery alone to adjuvant radiotherapy and then to multimodality therapy.

Extranodal marginal zone lymphomas or mucosa-associated lymphoid tissue (MALT) lymphomas were first described by Isaacson in 1983 where he postulated the MALT concept to explain the development of lymphomas in organs with no lymphoid tissues.^[1] This concept of acquisition of lymphoid tissue due to chronic antigenic stimulation by a bacterial infection is supported by the history of chronic cystitis, seen in around 20%–40% of the cases.^[3] On the other hand, in those cases with no such previous history, like ours, the

the pelvis showed soft-tissue density lesion with a size of 21 mm \times 37 mm \times 32 mm at the base of the urinary bladder – presumably residual disease or postradiation changes.

Table 1: All cases of marginal zone B-cell lymphoma urinary bladder reported since 2014

Author	Age (years)/gender	Presenting symptoms	Treatment	Outcome
D'Souza <i>et al.</i> ^[1]	56/male	Hematuria, clot retention	TURBT	CR at 3 years
Combaz and Kuhn ^[2]	67/female	Recurrent cystitis, asthenia	Rituximab (375 mg/m ² weekly, 4 weeks)	CR at 3 years
Isono <i>et al.</i> , 2018 ^[3]	77/female	Frequency, hematuria, cystitis	Radiotherapy (30.6 Gy in 17 fractions)	CR at 5 years
Ozawa <i>et al.</i> ^[5]	72/female	Thrombocytosis	TURBT	CR at 13 months
Matsuda <i>et al.</i> ^[7]	78/female	Refractory cystitis and renal dysfunction	Rituximab × 4 cycles	PR
Haddad-Lacle <i>et al.</i> , 2014 ^[8]	54/male	Low back pain	Radiotherapy (30.6 Gy in 17 fractions)	No recurrence at 3 years
Chen <i>et al.</i> ^[9]	63/female	Gross hematuria and urethral caruncle	Radiotherapy to bladder (36 Gy) and pelvic and inguinal lymph nodes (45 Gy)	Uneventful at 11 months
Hsu <i>et al.</i> , 2015 ^[10]	76/female	Recurrent cystitis	Radiotherapy	CR at 3 months

CR – Complete remission; PR – Partial response; TURBT – Transurethral resection of bladder tumor

alternative hypothesis is the development of tumors from the lymphoid tissue derived from the embryonic cloaca.^[2]

Extranodal marginal zone lymphomas are most common among primary urinary bladder lymphomas. More than 50 cases of primary bladder marginal zone lymphomas have been reported so far, after the first by Kuhara *et al.* in 1990.^[4] Patients are mostly elderly females, more than 60 years of age, with a history of chronic cystitis in the past, contrary to the patient reported here. Hematuria was the predominant symptom in 60%, with others having dysuria and urinary frequency.^[2]

Since there is no consensus on the optimal therapy, either of radiation, chemotherapy, targeted therapy, or a combination have been attempted and are successful in attaining remission. A few patients were treated with local therapy only, such as transurethral resection of bladder tumor^[5] and diathermy. There was one patient who attained CR with *Helicobacter pylori* eradication therapy alone,^[6] while another patient was treated exclusively with rituximab.^[2] In this context, all cases reported after 2014 are shown in Table 1. The prognosis remains good despite the modality of treatment used, but rarely, it can disseminate or transform into higher grade diffuse large B-cell lymphoma.^[2,7]

Conclusion

Although marginal zone lymphoma of the bladder is not a common diagnosis of a patient with hematuria, hematuria is its most common presentation. Hence, urine cytology and cystoscopy should be done in persistent cases of hematuria to rule out the same, as it has a very good prognosis with any form of either local or systemic therapy.

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent forms. In the form the patient(s) has/have given his/her/their consent for his/her/their images and other clinical information to be reported in the journal. The patients understand that their names and initials will not be published and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

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Nil.

Conflicts of interest

There are no conflicts of interest.

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