



Case report

Primary diffuse large B-Cell lymphoma of the uterine cervix with severe lower urinary tract Symptoms: A rare case report and review of the literature

Jiaosheng Li^a, Xiufen Zhang^b, You Liu^c, Qinhu Li^a, Yifan Guo^{b,*}, Haotian Yu^{a,d,*}

^a The Second School of Clinical Medicine, Southern Medical University, Guangzhou 510518, China

^b Department of Obstetric and Gynecology, Hainan Hospital of PLA General Hospital, Sanya 572013, China

^c Department of Pathology, Hainan Hospital of PLA General Hospital, Sanya, China

^d Department of Obstetrics and Gynecology, The Eighth Medical Center of PLA General Hospital, Beijing 100094, China

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ABSTRACT

Background: Diffuse large B-cell lymphoma (DLBCL) is a rare disease with a crude annual incidence rate of 3.8 cases per 100,000 people. Besides, primary cervical lymphoma is very rare, accounting for only 0.008% of cervical malignancies. (Nant et al., 2016) Although DLBCL patients often present with abnormal vaginal bleeding, it was not involved in this case. In this article, we present a rare case of primary cervical diffuse large B-cell lymphoma with urinary tract symptoms.

Case: A 71-year-old woman who had been suffering from dysuria for two months came to our hospital. A pelvic examination revealed a 10 cm cervical mass, while HPV and squamous cell carcinoma (SCC) antigen tests were negative. The bulky cervical mass invaded the posterior wall of the uterus, vagina, superior rectum, bladder, and bilateral lower ureters, resulting in dysuria and dilation of the upper ureter. Histopathological and immunohistochemical examination confirmed DLBCL and PET-CT suggested that it was stage IV. After two cycles of R-CHOP (rituximab, cyclophosphamide, doxorubicin, vincristine, and prednisone), the large lesions were eliminated. Unfortunately, the patient suffered an untimely death unrelated to her disease before the fourth cycle of R-CHOP could begin.

Conclusions: DLBCL of the cervix is a rare, but potentially curable disease if the diagnosis is made accurately, and doing so requires a high index of suspicion for cervical masses with an atypical presentation in which traditional diagnostic methods are equivocal. Obtaining adequate multilayered lesion biopsies containing both cervical epithelium and mesenchyme helps to avoid misdiagnoses. Histopathological biopsy and immunohistochemistry are the gold standards for diagnosis, and R-CHOP chemotherapy is an effective treatment.

1. Introduction

Diffuse large B-cell lymphoma (DLBCL) is a rare disease with a crude annual incidence rate of 3.8 cases per 100,000 people. Primary diffuse large B-cell lymphoma of the cervix occurs primarily in older patients, with a peak incidence between 54 and 60 years old. (Naniodu et al., 2017).

Perimenopausal or postmenopausal abnormal vaginal bleeding is the most common presentation of DLBCL. Other symptoms include pelvic pain, vaginal discharge, and post-coital bleeding. Occasionally, DLBCL of the cervix is an incidental finding, and pelvic examinations often

reveal a cervical mass or mass adjacent to the cervix.

Fox et al. (Fox et al., 1988) proposed the following criteria for diagnosing primary large B-cell lymphoma of the cervix: a) the lymphoma is confined to the cervix or adjacent organs; b) full examinations fail to reveal evidence of lymphoma elsewhere; c) no abnormal cells are found in the peripheral blood or bone marrow; d) several months pass between the appearance of lesions in the genital tract and extragenital lesions; e) there is no prior history of lymphoma.

Histopathology and immunohistochemistry are recognized as the benchmarks for the diagnosis of DLBCL. Additionally, DLBCL is a chemotherapy-sensitive tumor, and R-CHOP chemotherapy (rituximab,

* Corresponding authors at: Department of Obstetrics and Gynecology, Hainan Hospital of PLA General Hospital, Sanya 572013, China (G. Yifan) Department of Obstetrics and Gynecology, The Eighth Medical Center of PLA General Hospital, Beijing 100094, China (Y. Haotian).

E-mail addresses: guoyifan2016@126.com (Y. Guo), yhc200725@163.com (H. Yu).

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