

# Secondary B-cell lymphoma diagnosed by fine-needle aspiration cytology and flow cytometry following penile carcinoma: A case report

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**Abstract.** The number of studies reporting lymphoma as a secondary tumor has gradually increased. However, few studies have reported that occurrence of lymphoma as a secondary tumor following treatment for penile carcinoma, particularly cases in which the lymphoma was diagnosed by fine-needle aspiration cytology and flow cytometry. The present study reports the case of a 62-year-old male patient who was troubled with frequent urination and repeated chest tightness for 5 years. The diagnosis upon admission was penile carcinoma. Two months subsequent to the tumor removal surgery, enlarged lymph nodes were extracted from the patient using fine-needle biopsy, to be analyzed using light microscopy and flow cytometry. Smear results indicated a large number of abnormal cells scattered in the right axillary lymph node. Flow cytometry immunophenotyping of fine-needle aspiration samples indicated the increased expression of cluster of differentiation (CD)79a, CD19, CD20, CD38,  $\kappa$  chain and human leukocyte antigen-DR, which supported a diagnosis of B-cell lymphoma. Thus, the patient was diagnosed with B-cell lymphoma based on the results of the fine-needle aspiration biopsy and flow cytometry. The method of diagnosis and causes of therapy-related leukemia are discussed in the present report.

## Introduction

Non-Hodgkin's lymphomas (NHLs) are a heterogeneous group of lymphoproliferative disorders originating in B, T or natural killer (NK) lymphocytes. In the United States, B-cell lymphomas represent 80-85% of all NHL cases; 15-20% of cases are T- and NK-lymphomas (1). The National Comprehensive Cancer Network guidelines considers the following common subtypes of B-cell lymphoma: Diffuse large B-cell lymphoma, 31% of cases; follicular lymphoma, 22%; chronic lymphocytic leukemia/small lymphocytic lymphoma, 6%; mantle-cell lymphoma, 6%; mucosa-associated lymphoid tissue lymphoma, 5% (2). Overall NHL mortality rates are ~10.9 per 100,000 individuals in one year. However, there is little information concerning the mortality rates for the specific subtypes (3).

Recent studies have demonstrated that the diagnostic accuracy of fine-needle aspiration cytology (FNAC) may improve significantly when FNAC is used in combination with flow cytometry (FCM) or immunohistochemistry (4). FNAC offers several advantages: The procedure is quick, inexpensive and the aspiration procedure exhibits very few complications (5). At present, treatment options for B-cell lymphoma differ between patients. CHOP (cyclophosphamide, doxorubicin, vincristine and prednisone) chemotherapy has been the standard treatment for patients, and subsequently rituximab was added to CHOP to improve the outcomes for patients (2). High dose therapy with autologous stem cell rescue is another alternative for relapsed or refractory patients (3).

Non-Hodgkin's lymphoma as a secondary tumor has recently gained attention following decades of neglect during the diagnosis and treatment of primary tumors (6). Krikorian *et al* (7) were the first to report the increase risk of secondary NHL (sNHL) in patients successfully treated for a primary tumor. Krishnan and Morgan (8) reported that the lowest occurrence rates of sNHL was 0.07%, which was obtained from multinational population-based registries data in 1987, and the highest occurrence rate was 3%, which was obtained from the Norwegian Cancer Registry Database in 2002. Usually, sNHL develops after the first 5 years of initial therapy of primary cancer. The majority of sNHL cases

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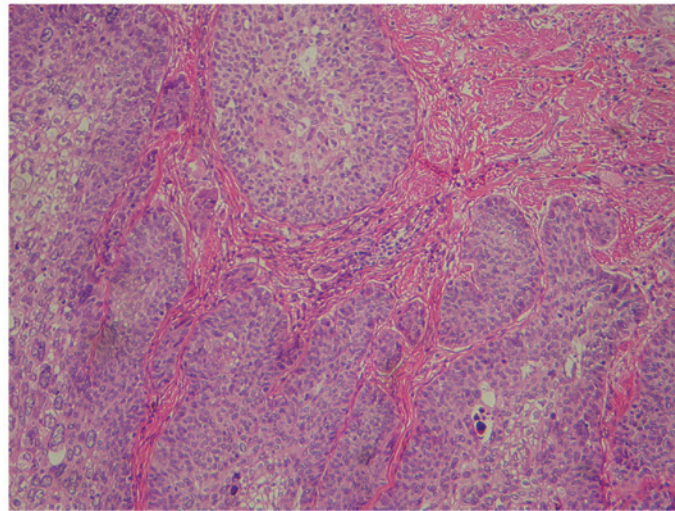


Figure 1. Histopathological analysis of the penile tumor that was removed during surgery. The penis tissue was composed of nests of heterogenous tumor cells with interstitial infiltration, and the tumor was determined to be moderately differentiated squamous cell carcinoma. The invasion depth was ~7 mm (hematoxylin and eosin staining; original magnification, x100).

develop after ~5 years of initial therapy for primary cancers. sNHLs lead to increased morbidity and mortality in these patients 8). Primary penile carcinoma has rarely been reported to lead to the development of B-cell lymphoma (9). To the best of our knowledge, the present study is the first to report a case of secondary B-cell lymphoma, diagnosed by FNAC and flow cytometry (FCM), following the treatment of penile carcinoma. Informed consent for the publication of this data was obtained from the patient.

### Case report

The patient, a 62-year-old man, had suffered from frequent urination, repeated chest tightness, and weakness for 5 years. The situation worsened 1 month prior to his admission to the People's Hospital of Zhejiang Province (Hangzhou, China) in May 2014. The patient was diagnosed with hyperplasia of the prostate gland and cardiomyopathy.

A physical examination of the patient revealed a long foreskin and a thick and swollen cauliflower-like mass (diameter, 3 cm) on the head of the penis. The right inguinal lymph nodes were swollen but painless. A preoperative biopsy of the lymph nodes indicated squamous cell carcinoma. The patient underwent a penis and scrotal skin excision, and the postoperative pathological diagnosis revealed a moderately differentiated squamous cell carcinoma (Fig. 1).

In July 2014, the patient returned to the People's Hospital of Zhejiang Province with a swollen and painful cheek. Upon physical examination, the superficial lymph nodes on the left neck, and right axillary and inguinal regions all demonstrated severe swelling, high activity and slight tenderness. Additional auxiliary examinations were performed. B-mode ultrasonography revealed that the lymph nodes of the bilateral axilla, groin and neck were enlarged. A chest computed tomography (CT) scan revealed multiple masses at the mediastinum, hilum of the right lung and the right axilla. In addition, bilateral lung lesions, a tubercle at the anterior side of the right upper lobe, which does not rule out lymph node infiltration, a calcified lesion at the apex

of the right lung, small levels of bilateral pleural effusion, and pericardial effusion were identified. The laboratory examination data were as follows: White blood cell count,  $4.64 \times 10^9$  cells/l (normal range,  $4-10 \times 10^9$  cells/l), including 77.1% neutrophils (normal range, 50-75%), 14.8% lymphocytes (normal range, 20-40%), 6.1% monocytes (normal range, 2-12%), 1.8% eosinophils (normal range, 0.5-5%) and 0.2% basophils (normal range, 0-2%); hemoglobin, 106 g/l (normal range, 120-160 g/l); platelet count,  $233 \times 10^9$ /l (normal range,  $100-300 \times 10^9$ /l);  $\text{Na}^+$ , 3.81 mmol/l (normal range, 135-145 mmol/l);  $\text{K}^+$ , 137 mmol/l (normal range, 3.5-5.5 mmol/l);  $\text{Cl}^-$ , 96.7 mmol/l (normal range, 96-108 mmol/l); total protein, 49.5 g/l (normal range 65-85 g/l); albumin, 27.1 g/l (normal range, 40-55 g/l); globulin, 22.4 g/l (normal range, 20-40 g/l); aspartate transaminase, 20 units/l (normal range, 0-50 units/l); alanine transaminase, 28 units/l (normal range, 10-52 units/l); alkaline phosphatase, 93 units/l (45-125 units/l); blood urea nitrogen, 7.75 mmol/l (normal range, 2.85-7.14 mmol/l); creatinine,  $93.3 \mu\text{mol/l}$  (normal range,  $44-133 \mu\text{mol/l}$ ); and uric acid,  $774 \mu\text{mol/l}$  (normal range, 210-440  $\mu\text{mol/l}$ ). A routine urine test revealed no abnormality; however, levels of tumor markers in the serum increased, including the levels of carbohydrate antigen 125 at 52.4 units/ml (normal range, 0-35 units/ml) and cytokeratin 19 at 5.9 ng/ml (normal range, 0-3.8 ng/ml). The other laboratory findings were also normal. A bone marrow smear analysis revealed the presence of hyperplastic myelocytes, with a large number of myelocyte and metamyelocyte cells and a low number of segmented granulocytes with toxic particles (data not shown). FNAC was performed on the lymph node of the present patient. The standard surgical procedure is as follows: Firstly, the skin above the mass to undergo biopsy is swabbed with an antiseptic solution; the skin, underlying fat and muscle may be numbed with a local anesthetic, although this is often not necessary with superficial masses. Secondly, a needle of extremely fine diameter is passed into the mass and withdrawn several times for sampling of cells. Finally, these cells are used in a smear and examined under a microscope or rendered into a suspension used for flow cytometry (10). A large

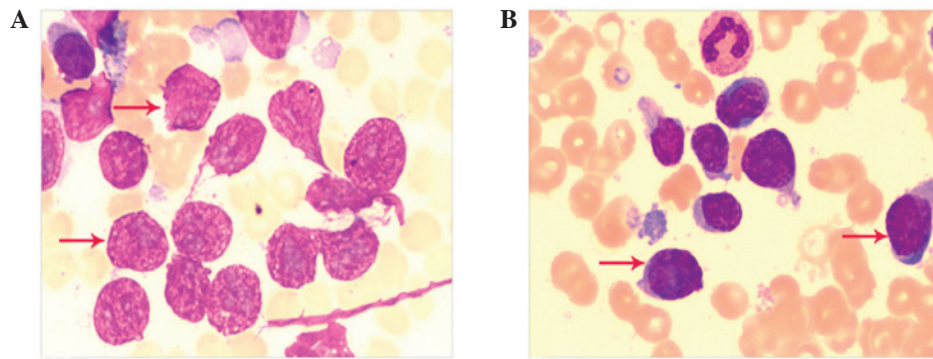


Figure 2. Smears of the lymph node biopsy samples obtained by fine-needle aspiration. (A) Large numbers of various sized and bare nuclei cells were present in the smear, and chromatin of the cells was dense (arrows). (B) Few round and large malformed cells were observed in the smear, the cytoplasm was smaller with no particles. Nuclear size was irregular and the nucleoplasm ratio was high. Chromatin was loose and colored blue and nucleolus were faintly visible (arrows). (Wright staining; original magnification,  $\times 1,000$ ).

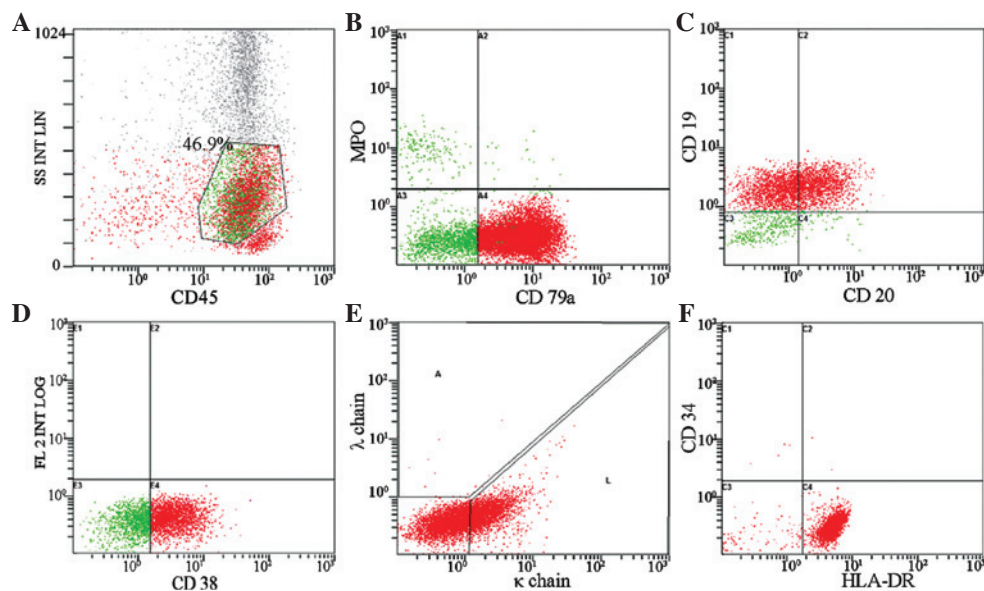


Figure 3. Flow cytometry immunophenotyping of the fine-needle aspiration sample of the lymph node. (A) Abnormal lymphocyte cell population demonstrated by forward and side scattered light. Increased expression of (B) CD79a, (C) CD20, (D) CD38, (E)  $\kappa$  chain and (F) HLA-DR. CD, cluster of differentiation; HLA-DR, human leukocyte antigen-DR; SS INT LIN, side scatter integral linear; MPO, myeloperoxidase; FL2 INT LOG, fluorescence channel 2 integral logarithmic.

number of abnormal and bare nuclei cells were distributed in the current smears obtained using FNAC, which were indicative of lymphoma cells (Fig. 2). In the present study, immunophenotyping of the aspirate obtained from FNAC revealed the following data, which supported the concept of a B-cell origin: Cluster of differentiation (CD)3<sup>+</sup>, 6.8%; CD3<sup>+</sup>CD4<sup>+</sup>, 1.9%; CD3<sup>+</sup>CD8<sup>+</sup>, 3.2%; natural killer, 2.7%; CD19<sup>+</sup>, 88.1%; CD10<sup>+</sup>, 0.2%; CD20<sup>+</sup>, 42.7%; CD79a, 84.8%; CD38, 64.9%; CD23<sup>+</sup>, 6.5%;  $\kappa$  chain, 31.0%; and  $\lambda$  chain, 1.4% (Fig. 3).

## Discussion

The number of studies reporting lymphoma as a secondary cancer has gradually increased, and the condition has been described following a number of types of primary tumor, including neuroblastoma (11), breast cancer (12), follicular thyroid carcinoma (13) and parotid gland cancer (14). There are a number of potential mechanisms acting in the pathogenesis of NHL, which may be associated with radiation,

cytotoxic drugs and immunosuppression (8). In the present report, a patient with penile cancer developed enlarged lymph nodes two months subsequent to penile cancer surgery. The recurrence of enlarged lymph nodes at the left neck, and right axillary and inguinal regions was the major reason for the readmission of the patient to the hospital. The tissues specimens that had been previously diagnosed with penile cancer were immunohistochemically examined and markers for B-cell lymphoma (CD19, CD20, CD22 and CD79a) were negative (data not shown); therefore, the possibility of penile cancer and B-cell lymphoma appearing at the same time was eliminated, and the B-cell lymphoma was concluded to be a secondary tumor that developed subsequent to penile cancer.

Immunophenotypic analysis may be performed using flow cytometry or immunohistochemistry, the choice depends on the antigens as well as the expertise and resources available to the hematopathologist. Immunohistochemistry has certain limitations on the diagnosis of certain tumors, including in cases where the lymph node is not easily accessible, such as



Hodgkin's lymphoma (15). FNAC is a simple, safe, minimally invasive and fast technique that is well tolerated by patients. In addition, the single cell suspension that is suitable for FCM analysis is easy to prepare (16). FCM is a rapid and sensitive multi-parameter analysis technique, which is important in the diagnosis of patients with lymph node enlargement (17). In the present study, a fine-needle aspiration biopsy of the lymph node was performed and FCM analyses were conducted. Immunophenotyping by FCM revealed increased expression of CD79a, CD19, CD20 and  $\kappa$  chain, and a diagnosis of B-cell lymphoma was confirmed.

The present study demonstrates that the combination of FNAC and FCM with lymph node aspiration analysis may improve the accuracy and sensitivity of lymphoma diagnoses. Therefore, the combined technique used in the present study may be used as the routine method in lymphoma diagnosis (18). The application of this technology in clinical practice will contribute to the earlier diagnosis and treatment for patients with secondary lymphoma.

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