

# Primary Testicular and Cutaneous Philadelphia Chromosome Positive B-Cell Lymphoblastic Lymphoma: A Rare Case and Review

Qiuxia Yu\*, Gaoxiang Wang\*, Jue Wang, Wei Zhang, Li Meng, Yang Cao

Department of Hematology, Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, Hubei, 430030, People's Republic of China

\*These authors contributed equally to this work

Correspondence: Yang Cao, Department of Hematology, Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology, No. 1095 Jiefang Avenue, Wuhan, Hubei, 430030, People's Republic of China, Tel +86 2783662680, Fax +86 2783662680, Email caoyangemma@163.com



**Abstract:** Philadelphia chromosome positive B cell lymphoblastic lymphoma (Ph+ B-LBL) is an extremely rare disease. We report a 27-year-old patient diagnosed with primary testicular and cutaneous Ph+ B-LBL without bone marrow involvement. The CCCG-LBL-2016 regimen (<https://clinicaltrials.gov/ct2/show/NCT02845882>) was initially administered due to the fast pathological diagnosis as B-LBL that was first obtained. To identify potential therapeutic targets, RNA sequencing (RNAseq) was also performed on lymph node specimens as a part of the routine diagnostic workup in our center. Unexpectedly, IKZF1 deletions and BCR-ABL1 fusion transcripts were detected. Based on these results, we retrospectively performed fluorescence in situ hybridization (FISH) for BCR/ABL1 rearrangements in the same lymph node specimen, and a 70% positive signal was detected. The patient subsequently received the CCCG-LBL-2016 protocol combined with the BCR-ABL tyrosine kinase inhibitor (TKI) dasatinib, along with prophylactic intrathecal infusion. Then, the patient underwent TBI-based haploidentical (haplo) allogeneic hematopoietic stem cell transplantation (haplo-allo-HSCT) as consolidation following the achievement of remission and continued taking dasatinib as maintenance therapy. The patient was still in complete remission 1 year after diagnosis. This case indicates that the detection of potential molecular targets, especially those targets that can be pharmacologically treated, such as BCR/ABL1 fusion transcripts, is of important value to both LBL diagnosis and therapeutic strategy choices. FISH, reverse transcriptase polymerase chain reaction (RT-PCR) and/or RNAseq should be routinely carried out in lymphoma specimens to depict its genetic landscape for the further execution of a precise therapy strategy.

**Keywords:** B-cell lymphoblastic lymphoma, Philadelphia chromosome, tyrosine kinase inhibitor, hematopoietic stem cell transplantation

## Background

Lymphoblastic lymphoma (LBL) is a rare neoplasm of immature lymphoid progenitors that arises from either B cell or T cell lineages. Due to their similar biological features, LBL and acute lymphoblastic leukemia (ALL) are considered a single entity (lymphoblastic leukemia/lymphoma) in the recent World Health Organization (WHO) classification of precursor lymphoid neoplasms.<sup>1</sup> Nevertheless, LBL is distinguished from ALL in clinical presentation. ALL mainly manifests as extensive bone marrow involvement. However, LBL is characterized by a predominantly nodal distribution of disease, and the percentage of bone marrow lymphoblast involvement is less than 25%. B-lineage LBL comprises approximately 10% of all LBLs, with a higher incidence in the younger age population.<sup>2</sup> Extranodal bone lesions, including the skin, soft tissues and osteolytic bone lesions, are frequently involved, but primary involvement sites of the bone marrow, visceral sites, testicles or central nervous system are rare.<sup>3</sup> The general characteristics of B-ALL/LBL and the different types of B-ALL/LBL with recurrent genetic abnormalities are detailed in the [Table S1](#).<sup>4-7</sup>

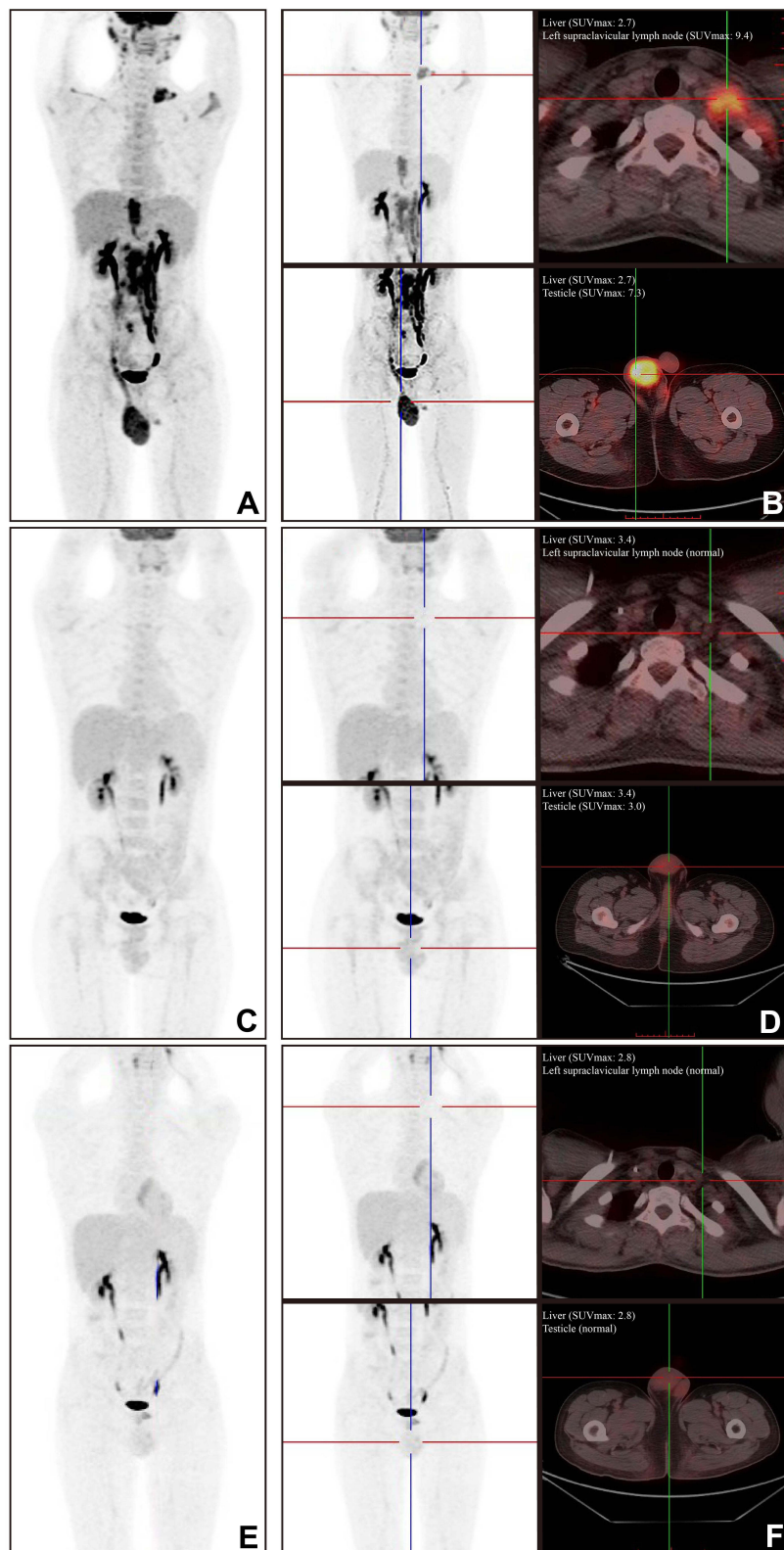
The Philadelphia (Ph) chromosome is the most common cytogenetic abnormality associated with chronic myeloid leukemia (CML) and ALL in adults, especially in elderly patients.<sup>8</sup> Therefore, Ph+ B-LBL is an extremely rare disease. To date, only 8 cases of Ph+ B-LBL have been reported in the published literature.<sup>9–16</sup> Herein, we report a case of a male with Ph+ B-LBL with unilateral swollen testis and scattered skin erythema as the first clinical presentations who successfully achieved complete remission through the high-risk chemotherapy protocol of CCCG-LBL-2016 combined with dasatinib. He maintained remission after TBI-based haplo-allo-HSCT and was subsequently maintained on dasatinib. This case study was approved by the Ethics Committee of Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology. The patient provided written informed consent according to the Declaration of Helsinki.

## Case Presentation

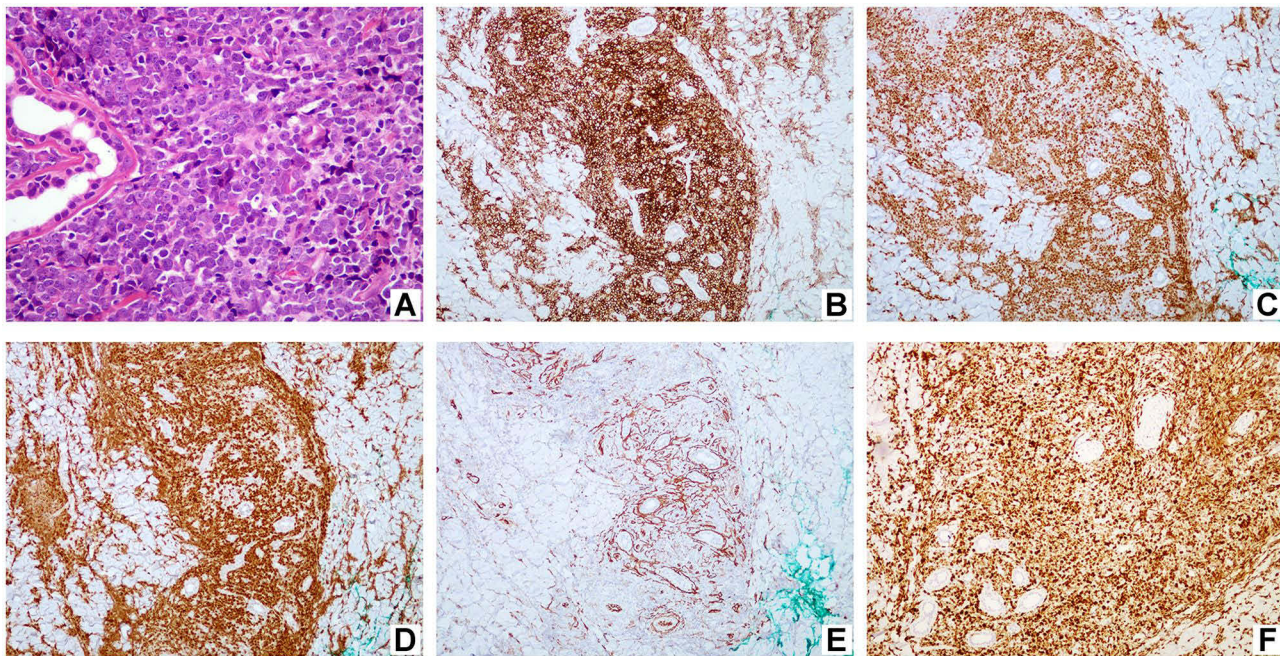
A previously healthy 27-year-old male first presented to the outpatient department in Tongji Hospital due to right testis swelling and scattered skin erythema onset for approximately 1 month. There were no other concomitant symptoms, such as fever, night sweat, or loss of weight. On physical examination, painless enlargement of submandibular and cervical lymph nodes was palpable. Then, he was admitted to our department for further diagnosis and treatment.

Routine laboratory evaluation demonstrated normal white blood cell (WBC) counts and hemoglobin and mildly increased platelet counts [ $376.0 \times 10^9/L$  (normal:  $150\text{--}350 \times 10^9/L$ )], lactate dehydrogenase [ $234 U/L$  (normal:  $135\text{--}225 U/L$ )] and  $\beta_2$  microglobulin [ $2.76 \text{ mg/L}$  (normal:  $0.8\text{--}2.2 \text{ mg/L}$ )]. Serology tests were negative for HIV, HBV, CMV and EBV infection. An abdominal ultrasound did not indicate hepatomegaly or splenomegaly. Bone marrow smear and histology revealed normal morphological, immunological flow cytometric (FCM), cytogenetic and molecular (MICM) results. No central nervous system infiltration was detected by cerebrospinal fluid flow cytometry. However, positron emission tomography/computed tomography (PET/CT) scans demonstrated abnormally increased metabolic activity in multiple lymph nodes (SUV max: 9.4), bones (right clavicle, left scapula, left 2nd rib, right accessory of 11th thoracic vertebra, 1st-2nd lumbar vertebra, bilateral femurs and bilateral superior tibial segment; SUV max: 6.2), the posterior walls of the nasopharynx (SUV max: 6.5), the bilateral ethmoid sinuses (SUV max: 5.0), and the bilateral testicular and right spermatic cords (SUV max: 7.3). There was also mildly increased metabolic activity in the subcutaneous nodules on the bilateral cheeks (SUV max: 2.8) (Figure 1A and B). The patient underwent ultrasound-guided fine-needle aspiration in the left enlarged supraclavicular lymph node. The pathological biopsy specimen was extensively replaced with tumor cells characterized by distinct nucleoli and scant cytoplasm. Immunohistochemical staining showed that representative B lymphoblastic tumor cells were positive for CD20 and terminal deoxynucleotidyl transferase (TdT) and negative for CD3, and approximately 90% of the neoplastic cells displayed nuclear Ki67 staining (Figure S1). Meanwhile, the cutaneous biopsy, also revealed consistency of the phenotype of the B lymphoblasts, were positive for CD20, CD19, PAX5, TDT, CD10, CD38, C-MYC, CD99, BCL-2 and CD43, but negative for CD3, CD34, CD7, BCL-6, MPO and MUM1, and Ki67 staining about 90% (Figure 2). Therefore, the diagnosis of B-LBL (Ann Arbor stage IIIA, IPI 3 scores) was confirmed.

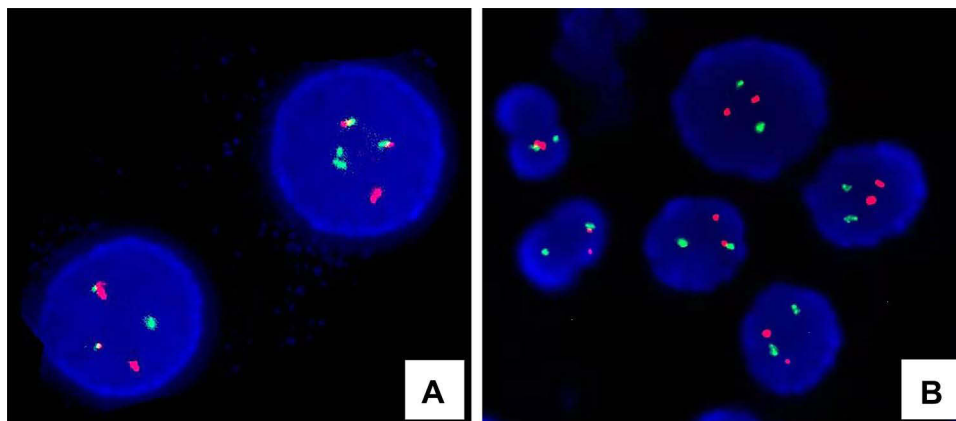
To control lymphoma in a timely manner, the patient received the high-risk chemotherapy protocol of CCCG-LBL-2016 (Induction I with prednisone, vincristine, daunorubicin, PEG-LASP, cyclophosphamide, cytarabine, 6-mercaptopurine plus prophylactic intrathecal infusion from Day 1 to Day 63). RNAseq, with reference to the GRCh37/hg19 genome and mainly to detect genetics disruption at the level of the human transcriptome, was performed on the left supraclavicular lymph node specimens as a part of routine diagnostic workup in our institution. Unexpectedly, IKZF1 deletions and BCR-ABL1 fusion transcripts were detected approximately 3 weeks after chemotherapy. Based on these results, we retrospectively validated BCR/ABL1 rearrangement in previous lymph node biopsy specimens by fluorescence in situ hybridization (FISH), which detected a 70% positive signal (Figure 3). Given that the modified diagnosis turned to Ph+ B-LBL, a combination of dasatinib (140mg, qd) was added to therapeutic regimen. PET/CT evaluation was carried out before Protocol M of CCCG-LBL-2016 began, and it showed complete remission (CR) (Figure 1C and D), Protocol M ( $5 \text{ g/m}^2$  methotrexate and 6-mercaptopurine) combined with dasatinib proceeded as planned for 4 cycles. Then, the patient finally underwent allogeneic transplantation with granulocyte colony-stimulating factor-mobilized bone marrow cells plus peripheral blood hematopoietic stem cells from his father, who was mismatched for five loci in March



**Figure 1** Representative of patient's PET imaging. (A and B) PET scan images revealed that FDG uptake was high in multiple lymph nodes (SUV max: 9.4) and bones (SUV max: 6.2), posterior wall of nasopharynx (SUV max: 6.5), bilateral ethmoid sinuses (SUV max: 5.0), bilateral testicle and right spermatic cord (SUV max: 7.3) and subcutaneous nodules on bilateral cheeks (SUV max: 2.8) before therapy. (C and D) PET/CT evaluation was carried out before Protocol M of CCCG-LBL-2016 beginning, and it showed complete remission. (E and F) Repeat PET/CT scan more than five months after transplantation showed normal metabolic activity. FDG, fluorodeoxyglucose; PET, positron emission tomography.



**Figure 2** Morphologic features of the skin lesion. (A) The core needle biopsy specimen is extensively replaced with tumor cells characterized by nucleoli and scant cytoplasm (H&E, 400x). (B–D) Immunohistochemical stains show that tumor cells were positive for CD20 ((B), 100x), PAX5 ((C), 100x), and TdT ((D), 100x). (E) Immunohistochemical stains show that tumor cells were negative for CD34 (100x). (F) About 90% of the neoplastic cells display nuclear Ki67 staining (100x). TdT, terminal deoxynucleotidyl transferase.



**Figure 3** FISH analysis was approached on suspension samples in interphase cells of fresh swelling lymph node specimen using commercial probe (LSI BCR/ABL DS probe, Vysis). (A) Cells with a signal pattern of 1 red, 1 green, and 2 yellow (fusion signal) are positive for BCR/ABL1 rearrangement. (B) Cells with 2 red and 2 green are normal. FISH, Fluorescence in situ hybridization.

2021. He received a TBI/VP/CY/ATG conditioning regimen consisting of total body irradiation (Day –9), etoposide (15 mg/kg/day on Days –8 and –7), cyclophosphamide (1.8 g/m<sup>2</sup>/day on Days –6 and –5), and ATG (thymoglobulin, 2.0 mg/kg/day on Days –4 to –1). The procedure was well tolerated, and he achieved successful engraftment on Day 15 post-stem cell infusion (SCI). He did not undergo any grade of acute or chronic graft-versus-host disease (GVHD) but only had temporary and asymptomatic Epstein–Barr virus (EBV)-reactivated viremia. All immunosuppressants were tapered and stopped within 2 months after SCI, and he took dasatinib 50 mg daily for maintenance. Monthly follow-ups were regularly performed, including the monitoring of bone marrow chimerism and BCR/ABL1 copies by polymerase chain reaction amplification of short tandem repeats (PCR-STR) and RT-PCR, respectively. Sustained complete chimerism and

negative BCR/ABL1 detection, along with recent PET/CT results (Figure 1E and F), indicated a satisfactory CR for more than 1 year.

## Discussion and Conclusion

Philadelphia chromosome-positive (Ph+) B cell lymphoblastic lymphoma is an extremely uncommon entity. To date, only 8 cases of Ph+ B-LBL have been published (see Table 1). Although B-LBL predominantly occurs in the younger-aged population, the 8 cases of Ph+ B-LBL that are currently reported involve all ages. B-LBL typically involves lymph nodes and extranodal sites, such as the skin, bone and soft tissues. Testicular infiltration occurs more frequently in disease progression and relapse and rarely appears as an initial manifestation. To our knowledge, this is the first case report of primary involvement of the testis and skin without evidence of bone marrow infiltration in B-LBL.

The lack of positive findings in the bone marrow would cause physicians to discount further detection of lymphoma tissues. Our case revealed that it may be clinically warranted to assess the molecular and cytogenetic characteristics in

**Table 1** Summary of Previous Case Reports

Case	Time	Sex	Age (Years)	Involvement Sites	First Detection of BCR-ABL1	Detection of BCR-ABL1 in BM	Treatment	SCT	Outcome	Reference
1	2013	F	43	Left parietal skull	FISH of the biopsy	Not tested	Hyper-CVAD/MA +dasatinib	Yes	CR (4 months)	Sadrzadeh et al <sup>9</sup>
2	2015	M	27	First: testis, Relapse: BM, PB	RT-PCR of BM (at relapse)	RT-PCR (+) (at relapse)	Hyper-CVAD/MA Relapse: imatinib +vincristine +prednisolone	No	Death (8 months)	Zhu et al <sup>10</sup>
3	2017	M	77	Testis, retroperitoneal lymph node, left pubic symphysis, L5 vertebral body Relapse: right kidney	RT-PCR of BM	FISH (-) RT-PCR (+)	Rituximab+hyper-CVAD/MA+ dasatinib→ imatinib→nilotinib	No	Death (4 years)	Boddu et al <sup>11</sup>
4	2018	M	26	Distal femur, pancreas, left kidney, multiple lymph nodes	FISH of the biopsies	Not tested	Vincristine +dexamethasone+ dasatinib	Yes	CR (4 months)	Alshomar et al <sup>12</sup>
5	2019	M	31	Paravertebral mass	RT-PCR of BM	FISH (-) RT-PCR (+)	Steroids+ imatinib→dasatinib; radiation	Yes	CR (30 months)	Dragani et al <sup>13</sup>
6	2019	M	65	Right humerus	FISH of the biopsy	FISH (-) RT-PCR (-)	CHOP, hyper-CVAD/MA +dasatinib, radiation	No	CR (5 years)	Takahashi et al <sup>14</sup>
7	2021	F	18	Right 5th rib, duodenum, pancreas tail	RT-PCR of BM	RT-PCR (+)	Prednisolone +dasatinib; Hyper-CVAD/MA +dasatinib	Yes	CR (4 months)	Yamada et al <sup>15</sup>
8	2021	M	19	Multiple bone	FISH of the biopsy	Not tested	Vinorelbine +daunorubicin +prednisone +imatinib, clarithromycin	No	CR (30 months)	Li et al <sup>16</sup>

lymphoma tissues at initial diagnosis, regardless of whether the bone marrow is involved. With the RNAseq results of BCR/ABL1 (p190) rearrangement and IKZF1 deletion, a precise diagnosis was ultimately made in our case. We noticed that a small number of B-LBL patients with KMT2A (MLL) rearrangements have also been reported previously.<sup>17,18</sup> Chromosomal translocations in B-ALL, such as the BCR/ABL, t(9;22)(q34;q11.2) alteration and MLL rearrangements involving chromosome 11 at 11q23, have also been associated with adverse outcomes. These B-LBL cases with ALL-associated BCR/ABL and KMT2A gene rearrangements support the interpretation of B-LBL and B-ALL as a spectrum of the same disease by the WHO classification. Routine assessment of molecular and cytogenetic characteristics in B-LBL specimens as in B-ALL bone marrow at initial diagnosis may be useful for risk stratification.

The advent of tyrosine kinase inhibitors (TKIs) has changed the prognosis of Ph+ ALL.<sup>19,20</sup> Although Ph+ B-LBL is a rare entity without standard treatment strategies, the current therapy program could utilize the Ph+ ALL protocol that predominantly contains tyrosine kinase inhibitors (TKIs), which target BCR/ABL1. Notably, Case 2, listed in Table 1, failed to perform a molecular abnormality screening at his initial diagnosis, leading to a loss of opportunity for targeted therapy with rapid relapse and very poor outcomes.<sup>10</sup> Therefore, this study clearly shows the importance and necessity of potential target analysis in B-LBL at first diagnosis.

The choice to use TKIs is another key point of concern. Because the testis is an immune-privileged site, primary testicular malignancies provide a sanctuary for cancer cells. They have a uniformly poor prognosis with a high risk of relapse in the CNS and the contralateral testis.<sup>21</sup> Studies have indicated that some patients with Ph+ ALL on prolonged imatinib therapy have progressed to isolated CNS leukemia due to the poor permeability of imatinib.<sup>22</sup> In contrast, dasatinib, a dual-specific SRC/BCR-ABL1 kinase inhibitor, can cross the blood–brain barrier and be present at detectable levels in the CSF to clear blasts.<sup>23</sup> Remarkably, in the previous reports of patients with testicular infiltration (shown in Table) and in our case, the patients were administered dasatinib, and they achieved CR without relapses in any immune-privileged sites.<sup>11</sup> Thus, Ph+ B-LBL with testis involvement could also benefit from dasatinib, which might effectively penetrate the blood–testis barrier as well as the blood–brain barrier, clear blasts and prevent relapse in these privileged sites.

Given its rapid proliferative features and the expansion of a BCR/ABL1 mutated dasatinib and imatinib-resistant clone, the treatment strategy for Ph+ ALL should be consolidated with allo-HSCT following achievement of remission. However, a recent study of TKI-based chemotherapy-free induction and consolidation as a first-line treatment in GIMEMA LAL2217 poses a challenge to the allo-HSCT approach at CR1 for the consolidation of Ph+ ALL.<sup>24</sup> The role of allo-HSCT in Ph+ B-LBL is also not clear. We observed that 4 of the 8 cases in Table were treated with a TKI followed by allo-HSCT at CR1, with disease-free survival (DFS) ranging from 4 to 30 months.<sup>9,12,13,15</sup> The outcomes of the other 4 cases without allo-HSCT varied widely. One patient without a TKI upfront relapsed 4 months after chemotherapy.<sup>10</sup> Another patient with a TKI upfront but without allo-HSCT relapsed 4 years after the initial diagnosis.<sup>11</sup> Interestingly, the 2 patients without allo-HSCT were probably due to age as contraindications and were finally sustained in a CR status for 4 and 5 years, respectively, using TKIs only for maintenance.<sup>11,14</sup> In light of the younger age and longer survival expectations of the patient in our study, we speculated that he would have benefit from allo-HSCT in CR1, although this should be confirmed in large-scale studies in the future.

In conclusion, we describe a rare case of primary testicular and cutaneous Ph+ B-LBL without bone marrow involvement. Therefore, it was very easy to neglect the screening of potential molecular targets, which would lead to a missed diagnosis. However, we successfully made an accurate diagnosis and implemented an appropriate therapeutic strategy at the initial hospitalization through the proper detection of lymphoma specimens. We suggest that FISH, reverse transcriptase polymerase chain reaction (RT-PCR) and/or RNAseq should be routinely carried out in lymphoma specimens. It is not only valuable for hematologists to tailor treatment but also for patients to improve their outcomes. Nevertheless, optimized therapeutic strategies for Ph+ B-LBL need further study.

## Data Sharing Statement

The materials and methods supporting the conclusion of this study have been included within the manuscript and supplementary materials.

## Ethics Approval and Consent to Participate

This case study was approved by the Ethics Committee of Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology. Institutional approval was not required to publish the case details. The patient provided written informed consent according to the Declaration of Helsinki.

## Consent for Publication

The patient provided written informed consent for the case details to be published.

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## Author Contributions

All authors contributed to data analysis, drafting or revising the article, have agreed on the journal to which the article will be submitted, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

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## Disclosure

The authors report no conflicts of interest in this work.

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