Hepatic Hodgkin Lymphoma Presenting as Solitary Hepatic Mass Following Other Iatrogenic Immunodeficiency-Associated Lymphoproliferative Disorder in a Patient With Rheumatoid Arthritis

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Abstract

Lymphoproliferative disorders (LPDs) occur frequently in patients with rheumatoid arthritis (RA) under methotrexate treatment. Some LPDs spontaneously regressed after methotrexate treatment, but classic Hodgkin lymphoma (CHL)-type LPDs frequently relapse, and chemotherapy is usually required for the treatment. CHL usually spreads in contiguous lymph nodes and then infiltrates in organs at an advanced stage. Thus, hepatic Hodgkin lymphoma (HHL) without lymphadenopathy is extremely rare at diagnosis. We present a case of methotrexate-associated LPDs associated with systemic lymphadenopathy and hepatosplenic mass in a 71-year-old woman with RA under methotrexate treatment over 10 years. Although spontaneous remission occurred after methotrexate discontinuation, she developed HHL presenting as a solitary hepatic mass without lymphadenopathy 3 years after spontaneous regression. She received brentuximab vedotin (BV) combination chemotherapy without bleomycin to avoid pulmonary toxicity. Complete metabolic response was achieved after four courses of BV combination chemotherapy, and the activity of RA was kept to be in remission. Our case suggested that the recurrence lesions of LPDs may present at unexpected site, which is not coincide with the primary site, and BV combination chemotherapy is a promising regimen for limited-stage CHL-type LPDs in patients with RA owing to its anti-lymphoma effect on CHL-type LPDs and a possible targeted therapy for RA.

Keywords: Classic Hodgkin lymphoma; Lymphoproliferative disorders; Hepatic mass; Methotrexate; Brentuximab vedotin; Combination chemotherapy

Introduction

Classic Hodgkin lymphoma (CHL) is a lymphoid hematological malignancy characterized by clinically lymph node enlargement and pathologically characteristic appearance of Hodgkin cells and Reed-Sternberg cells. It usually spreads in contiguous lymph nodes along the lymphatic system and then infiltrates in organs such as the liver, bone, and lung [1]. Thus, hepatic involvement usually occurs late during the disease or at an advanced stage, and hepatic presentation without lymphadenopathy is extremely rare [2]. The standard treatment regimen consists of doxorubicin, bleomycin, vinblastine, and dacarbazine (ABVD), but pulmonary sequelae due to bleomycin toxicity usually occur after ABVD chemotherapy. Brentuximab vedotin (BV) is monoclonal anti-CD30 antibody conjugated with monomethyl auristatin E. Hodgkin cells and Reed-Sternberg cells usually express CD30, and CD30 is the target antigen for BV. BV in combination with ABVD chemotherapy without bleomycin has recently been reported to show favorable efficacy for CHL [3]. We report here a case of hepatic Hodgkin lymphoma presenting as a solitary hepatic mass following other iatrogenic immunodeficiency-associated lymphoproliferative disorders (OII-LPDs) in a patient with rheumatoid arthritis (RA). Our patient received BV combination chemotherapy and achieved complete response. This is the first case of hepatic Hodgkin lymphoma in which complete metabolic response was achieved with BV combination chemotherapy in a patient with RA.

Case Report

A 71-year-old woman was admitted to our hospital for the evaluation of multiple hepatic masses in December 2015. She was diagnosed as having RA in 1992 at another hospital and had been taking methotrexate for RA for over 10 years. She had a fever of more than 38 °C for over 2 months, and her laboratory examination showed an elevated level of C-reactive protein (CRP). Computed tomography (CT) showed multiple lymphadenopathies in bilateral supraclavicular and mediastinal regions, and multiple masses in the liver and spleen. Methotrexate was discontinued before her admission to our hospit-
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Discussion

Here, we report a rare case of hepatic Hodgkin lymphoma following OII-LPD in a patient with RA. The standardized incidence ratio of LPDs is increased in RA patients treated with methotrexate [4, 5]. MTX-LPDs are complications of methotrexate treatment in patients with RA. The most frequent histological subtype of MTX-LPDs is diffuse large B-cell lymphoma followed by CHL. Momose et al [6] reported that CHL-type OII-LPDs (CHL-LPDs) account for about 13% of OII-LPDs, among which many showed involvement of lymph nodes or a combination of lymph nodes and other organs, such as the liver and lung. However, to the best of our knowledge, only one case of OII-LPD that involved solely the liver has been reported. Tsukazaki et al [7] reported the case of an 88-year-old female with RA who developed hepatosplenic Hodgkin lymphoma. She had been taking methotrexate and infliximab for 6 years, and she developed lymph node enlargement and hepatosplenomegaly. Hodgkin lymphoma was diagnosed on the basis of the finding of cervical lymph node biopsy. Spontaneous regression occurred after the withdrawal of methotrexate and infliximab, but after 1 year, her Hodgkin lymphoma relapsed presenting as hepatosplenomegaly without lymphadenopathy. She was treated with four cycles of ABVD chemotherapy and achieved a complete response. In our patient, multiple hepatosplenic mass and lymph nodes enlargement were found on CT scans when she was under methotrexate treatment; and about 3 years later, a solitary hepatic mass without lymphadenopathy was found on PET-CT scans when she was under the treatment of biologics such as etanercept and abatacept. She was treated with BV combination chemotherapy and achieved CMR, as shown by PET-CT. This is the second case of a patient with RA with a hepatic presentation of classic Hodgkin lymphoma recurring following OII-LPD.

Patients with primary hepatic lymphoma (PHL) presented with hepatic failure or encephalopathy. Imaging methods such as CT showed hepatomegaly without any mass [8]. This indicated a diffuse infiltration pattern, as in the case reported by Tsukazaki et al. Almost all cases with the diffuse infiltration pattern of PHL showed secondary hepatic involvement in an advanced stage of CHL. Otherwise, patients with PHL presented with no obvious constitutional symptoms such as fever, weight loss, and general malaise. Imaging methods showed a solitary mass or multiple masses in the liver without lymphadenopathy [2]. This indicated the nodular infiltration pattern, as in our case. In the case of primary non-Hodgkin lymphoma of the liver, Emile et al [9] showed that the diffuse infiltration pattern indicated a worse prognosis than the nodular infiltration pattern. However, in the case of CHL, there has been no report about whether the diffuse infiltration pattern has a worse prognosis than the nodular infiltration pattern. More cases should be collected to clarify this point.
Patients with RA frequently have pulmonary complications including interstitial pneumonia and are treated with immunosuppressive drugs such as methotrexate or tacrolimus, prednisolone, and biologics. These drugs lead to immunosuppression and complications, especially infectious pneumonia [10]. On the other hand, bleomycin leads to pulmonary fibrosis and fatal pulmonary complications in patients with CHL [11, 12]. Thus, extra care should be taken for patients with RA and CHL for pulmonary complications under treatment with ABVD chemotherapy. Although ABVD chemotherapy is the standard treatment for CHL, Abramson et al [13] reported that BV combination chemotherapy without consolidation radiotherapy is effective for nonbulky limited-stage CHL in a multicenter phase 2 study. Thirty-four patients were enrolled, and those who received four cycles of BV combination chemotherapy achieved a high complete response rate. Ichikawa et al

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**Figure 1.** Images of hepatic mass. (a) CT images showed a solitary low-density mass in the right lobe of the liver. MRI showed a low-intensity solitary hepatic mass on T1-weighted image (b), a high-intensity mass on T2-weighted image (c), and a high-intensity mass on diffusion-weighted image (d). (e) PET-CT images showed abnormal accumulation of fluorodeoxyglucose in liver (arrow); (upper panel) whole-body PET-CT image. (lower panel) transverse section of PET-CT image. (f) PET-CT images after four courses of BV combination chemotherapy showed no abnormal accumulation of fluorodeoxyglucose in whole body, which indicated that she remained in complete metabolic remission; (upper panel) whole-body PET-CT image, (lower panel) transverse section of PET-CT image. CT: computed tomography; MRI: magnetic resonance imaging; PET-CT: positron emission tomography computed tomography.
[14] reported that three patients with methotrexate-associated advanced-stage CHL received six courses of BV combination chemotherapy and achieved CMR without pulmonary toxicity. Our patient received four courses of BV combination chemotherapy and achieved CMR as shown by PET-CT. Thus, this is the first case in which BV combination chemotherapy was effective for limited-stage hepatic CHL-LPD in a patient with RA. Recently, Vachhani et al [15] and Nakazato et al [16] have reported that BV may represent a novel targeted therapy for RA. This point may support the choice of BV combination chemotherapy instead of ABVD chemotherapy for CHL-LPDs in patients with RA.

In conclusion, we report a case of hepatic Hodgkin lymphoma presenting as a solitary hepatic mass following OII-LPD in a patient with RA. Our patient achieved CMR after four courses of BV combination chemotherapy. Our case suggested that the recurrence lesions of LPDs may present at unexpected site, which is not coincide with the primary site. BV combination chemotherapy is a promising regimen for CHL-LPDs in patients with RA owing to its anti-lymphoma effect on CHL-LPDs and a possible targeted therapy for RA.

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Conflict of Interest

All authors declare no conflict of interest.

Informed Consent

We verbally obtained the informed consent from the patient for the publication of this case report and accompanying images.

Author Contributions

YT designed the study, collected the data and wrote the paper; SA provided all the pathological pictures; AH and IS treated this patient; IS supervised this study. All authors contributed to the editing of this manuscript.

Data Availability

The authors declare that data supporting the findings of this study are available within the article.

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