Gradually-Deteriorating Liver Function due to Iron Overload Over Four Years after Allogeneic Stem Cell Transplantation

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In allogeneic hematopoietic stem cell transplantation (allo-SCT) recipients with liver dysfunction, it is often difficult to determine the cause. Several cases of liver dysfunction may be interpreted as chronic graft versus host disease without a definitive diagnosis, resulting in continued immunosuppressive therapy for longer periods. Allo-SCT recipients commonly require frequent red blood cell transfusions during the course of treatment and transplantation, leading to significant iron overload, which could be one of causes of liver dysfunction. Here we report an allo-SCT recipient with chronic deteriorating liver dysfunction due to iron overload, despite maintaining transfusion independence for more than four years. Using magnetic resonance-based liver iron concentration (MR-LIC), iron overload-related liver dysfunction was diagnosed. It drastically improved with monthly phlebotomy and has not recurred following its termination. The observations from our case suggested that iron overload should be recognized as a cause of chronic liver dysfunction even in patients who remain transfusion-independent for several years and that MR-LIC analysis is a useful and reliable method for detecting iron overload and monitoring the effect of iron-reduction therapy.

Keywords: iron overload, liver dysfunction, transfusion independence, magnetic resonance imaging, phlebotomy

INTRODUCTION

In allogeneic hematopoietic stem cell transplantation (allo-SCT) recipients with liver dysfunction, it is often difficult to determine whether the cause of liver dysfunction is hepatic graft-versus-host disease (GVHD), viral liver dysfunction, drug toxicity, non-alcoholic steatohepatitis, or iron overload.1 Several cases of liver dysfunction may be often interpreted as chronic hepatic GVHD, resulting in continued immunosuppressive therapy for longer periods, despite liver dysfunction being the only symptom that could suggest chronic GVHD.

Allo-SCT recipients commonly require frequent red blood cell (RBC) transfusions during the course of treatment and transplantation, leading to significant post-transplantation iron overload. Tissue accumulation of iron may eventually lead to progressive liver, heart, and endocrine gland dysfunction. Pre-transplant iron overload has been significantly associated with an increased incidence of transplant-related mortality and decreased overall survival rate.2,3 However, to date, delayed post-transplant iron overload and its complications have not been well characterized, except in patients with thalassemia.4,5

Here we report the successful treatment by monthly phlebotomy of chronic deteriorating liver dysfunction in an allo-SCT recipient. Despite maintaining transfusion independence for more than four years, the patient was diagnosed with iron overload by serum ferritin (SF) levels and magnetic resonance-based liver iron concentration (MR-LIC) analyses.

CASE REPORT

A 58-year-old Japanese man developed chronic-phase chronic myeloid leukemia (CML) in September 2005. Treatment with imatinib was initiated; however, it resulted in severe pancytopenia. The patient received blood transfusions over six months; as a result of which, imatinib was eventually discontinued in the absence of a cytogenetic response. Bone marrow transplantation from an HLA-matched sibling donor...
was performed in September 2007, following conditioning with busulfan and fludarabine. A complete cytogenetic response was initially achieved; however, the disease relapsed with increased amounts of recipient cells, and the patient developed severe pancytopenia. In May 2008, the patient underwent allogeneic peripheral blood stem cell transplantation (PBSCT) from the same sibling donor, following conditioning with fludarabine and melphalan. For both transplantation procedures, Cyclosporin A (CsA) and short-term methotrexate were administered for GVHD prophylaxis. Engraftment was achieved on the 11th post-PBSCT day. However, 28 days post-PBSCT, the patient developed acute GVHD with skin involvement and diarrhea; hence, he was treated with 2 mg/kg of prednisolone. A complete molecular response was achieved 67 days post-PBSCT, and the patient remained independent of blood transfusions following engraftment. SF level on the 63rd post-PBSCT day was 2,520 ng/mL (reference range, 25-280 ng/mL). On the 105th post-PBSCT day, when the CsA dose was reduced to 20 mg, γ-glutamyl transpeptidase levels acutely increased to 560 IU/mL (normal, 10-47 IU/mL). Concurrently, serum alanine aminotransferase (ALT) and SF levels were also found to be elevated [177 IU/mL (reference range, 8-42 IU/mL) and 4,840 ng/mL, respectively]. Although there was no histological evidence, liver dysfunction was initially attributed to chronic hepatic GVHD due to the development of characteristic signs and symptoms of chronic GVHD, including sicca, gingival erosion, and bronchiolitis obliterans. The CsA dose was increased to 100 mg, leading to immediate improvement in liver dysfunction. On the 171st post-PBSCT day, previously elevated SF levels decreased to 2,710 ng/mL.

Over the four years following transplantation, liver function fluctuated and gradually deteriorated, despite of improvement in the other symptoms of chronic GVHD (Fig. 1). Tapering of CsA dose was attempted, however, it could not be decreased to less than 50 mg. Laboratory evaluation in January 2013 revealed the following biochemical results: total bilirubin, 0.5 mg/dL; aspartate aminotransferase, 160 IU/mL (reference range, 13-33 IU/mL); ALT, 177 IU/mL; lactate dehydrogenase, 221 IU/mL (reference range, 119-229 IU/mL); alkaline phosphatase, 288 IU/mL (reference range, 115-359 IU/mL); γ-glutamyl transpeptidase, 35 IU/mL; white blood cell count, 7.7 × 10^9/L; RBC count, 4,690 × 10^9/L; hemoglobin, 15.1 g/dL; platelets, 244 × 10^9/L; serum iron (Fe), 219 µg/dL; unsaturated iron-binding capacity, 167 µg/dL; total iron-binding capacity, 386 µg/dL; and SF, 4,529 ng/mL. Serological tests for hepatitis B and C virus were negative. Abdominal ultrasonography revealed no abnormal lesions in the liver or biliary system, except for elevated hepatorenal contrast. The patient received 56 RBC transfusion units in total since the onset of CML and remained RBC transfusion-independent for more than four years thereafter.

MR-LIC was calculated according to a well-validated algorithm described by Gandon and colleagues. Magnetic resonance imaging of the liver with various gradient-recalled echo sequences was performed using a 1.5-T system (EXCELART Vantage XGV, Toshiba Medical Systems Corporation, Japan). The correlation between the liver-to-muscle signal intensity ratio and liver iron concentration was calculated. An algorithm to calculate MR-LIC was developed using data from the aforementioned study and is publicly available online (http://www.radio.univ-rennes1.fr/index.html.en.html).

Iron overload-related liver injury was diagnosed with an MR-LIC of 200 µmol/g. Phlebotomy, drawing 400 mL of whole blood, was performed every four weeks after the initial
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Kamble et al. reported that these patients can be correctly diagnosed by liver biopsy and successfully treated with phlebotomy. In our case, the first liver dysfunction on the 105th post-PBSCT day was confirmed to be chronic hepatic GVHD because it was effectively treated with CsA, and the other characteristic symptoms of chronic GVHD were complicated. Therefore, the liver injury was assumed to be due to prolonged chronic hepatic GVHD for over four years. However, in fact, the primary cause of his liver dysfunction might be iron overload for four years following allo-SCT, because all the other symptoms suggestive of chronic GVHD had already improved, and the liver function drastically improved by only phlebotomy. His liver dysfunction was not severe, and the patient remained asymptomatic; however, it is essential to note that CsA administration, which was needlessly continued on a long-term basis, could have been terminated. CsA is expensive and has been known to have adverse effects, such as immune depression, glucose intolerance, and dyslipidemia.

In summary, our case highlights the susceptibility of RBC transfusion-independent patients to liver dysfunction due to hepatic iron overload even several years after allo-SCT. We demonstrate the efficacy of monthly phlebotomy for treating iron overload in transfusion-independent allo-SCT recipients. Finally, we propose MR-LIC analysis using Gandon’s algorithm as a useful and reliable method for detecting iron overload and monitoring the effect of iron-reduction therapy.

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REFERENCES


