Development of a secondary autoimmune disorder after hematopoietic stem cell transplantation for autoimmune diseases: role of conditioning regimen used

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Patients undergoing autologous hematopoietic stem cell transplantation (auto-HSCT) for autoimmune disease may have an added propensity to develop a second autoimmune disorder, given the genetic predisposition to autoimmunity. Therefore, we undertook a retrospective analysis of all patients who have undergone auto-HSCT for an autoimmune disease in our institution to determine the occurrence of a secondary autoimmune disorder and possible risk factors. In all, 155 patients underwent auto-HSCT for various autoimmune diseases; of those patients, 6 manifested a distinct secondary autoimmune disease at a median of 8.5 months (range, 2-30 months) after auto-HSCT.

Introduction

Autoimmune disorders have been reported to occur following both autologous and allogeneic hematopoietic stem cell transplantation (HSCT) for malignant and nonmalignant conditions, with autoimmune cytophenias being reported most frequently.1-4 The underlying mechanism for the development of autoimmunity is postulated to involve impaired function of regulatory T cells allowing self-reactive T cells to act unchecked.5 Delayed T-cell reconstitution following transplantation may thus predispose patients to the development of a secondary autoimmune disorder.

In addition to the immunologic milieu, a genetic predisposition to autoimmunity is well described, with epidemiologic studies demonstrating a higher rate of concordance among monozygotic compared with dizygotic twins, and association with certain HLA types.5,6 Patients with autoimmune disorders who have undergone autologous HSCT (auto-HSCT), with the combination of genetic predisposition and the lympho-deficient state after undergoing transplantation, may thus be at risk of developing another autoimmune disorder.

We therefore undertook a retrospective analysis of all patients with autoimmune diseases who had undergone auto-HSCT in our institution to determine the occurrence of a secondary autoimmune disorder. We studied the characteristics of affected patients, and the treatment and outcome of the secondary autoimmune disease to assess the factors that may predispose to their development. We also discuss herein the possible contributory role played by the agents used in the conditioning regimen, with particular reference to alemtuzumab and antithymocyte globulin (ATG).

Patients and methods

We carried out a retrospective analysis of all patients who had undergone an auto-HSCT for an autoimmune disease in our institution. All patients underwent transplantation on protocols approved by the Institutional Review Board of Northwestern University, Chicago, IL. Patients met eligibility criteria for the respective disease-specific protocols, which have been previously described.8-12 After informed consent in accordance with the Declaration of Helsinki, the patients received cyclophosphamide (2 g/m²) and subcutaneous filgrastim for mobilization, and had peripheral blood stem cells collected (PBSCs) that were either unmanipulated or manipulated with CD34 selection by either Isolex, Nexell, Irvine, CA; or Cellpro, Bothell, Washington, or unmanipulated. The conditioning regimens consisted of various combinations of cyclophosphamide with either ATG (rabbit or equine) or alemtuzumab or cyclophosphamide with either intravenous busulfan or total body irradiation (TBI), again depending on disease-specific protocols. PBSC infusion was on day 0, 36 hours after completion of cyclophosphamide. Subcutaneous filgrastim was administered from either day 0 or day +6 and continued until neutrophil recovery. Neutrophil engraftment was defined as the first day after transplantation when the absolute neutrophil count exceeded 0.5 × 10⁹/L. Platelet engraftment was defined as the first of 3 days when platelet counts exceeded 20 × 10⁹/L without transfusion. All blood products were irradiated and stored in an aseptic environment. Correlation of the various conditioning regimens with the development of a second autoimmune disorder was performed using survival analysis by the Kaplan-Meier method. Differences between the proportions of patients developing a second autoimmune disorder were assessed using the log-rank test. A P value less than .05 was considered to be significant.

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filtered and were cytomegalovirus safe. Packed red blood cells were transfused when hemoglobin declined below 80 g/L or where clinically indicated. Platelet transfusions were administered when platelet counts fell below 20 × 10^9/L or where clinically indicated.

Antimicrobial prophylaxis was with oral ciprofloxacin, valacyclovir or acyclovir, fluconazole or voriconazole, and trimethoprim-sulfamethoxazole or nebulized pentamidine. During the period of neutropenia, patients were given intravenous ceftazime or ceftriaxone prophylactically. Herpes virus prophylaxis was continued for 12 months after transplantation, whereas antifungal and Pneumocystis jiroveci prophylaxis was continued for at least 6 months, or longer if the patient remained on immunosuppressive therapy. The patients were followed up at 6 and 12 months after undergoing transplantation, then yearly thereafter.

For purpose of analysis, patients were grouped by conditioning regimen into 3 categories, namely ATG-containing, alemtuzumab-containing, and without lympho-depleting antibodies (ie, cyclophosphamide-TBI or busulfan-cyclophosphamide). Fisher exact test was used to compute the statistical significance of observed differences in the frequency of secondary autoimmune disorders occurring among the 3 groups. Comparisons were also made pair-wise. In addition, the impact of sex, type of ATG used, and CD34 selection of the PBSC product were evaluated, and Fisher exact test was used to determine statistical significance. All P values were 2-sided and the statistical software employed was R version 2.2.0 (R Foundation for Statistical Computing; http://www.r-project.org). The same software was used to compute exact confidence intervals and bounds for odds ratios in 2-by-2 tables. An approximation of the odds ratio was obtained in cases where n was 0 by the addition of 0.25 to each cell in the 2-by-2 table, where the odds ratio would otherwise be expressed as “infinity.”

Results

During the period from August 1996 to March 2006, 155 patients with various autoimmune disorders underwent an auto-HSCT. The autoimmune diseases represented were systemic lupus erythematosus (SLE; n = 60, 38.7%), systemic sclerosis (SSc; n = 13, 8.4%), multiple sclerosis (MS; n = 43, 27.7%), rheumatoid arthritis (n = 6, 3.9%), Crohn disease (n = 19, 12.3%), and others (n = 14, 9.0%). There were 6 patients (3 with SLE, 2 with MS, and 1 with SSc) who developed an autoimmune disorder distinct from their underlying autoimmune disease at a median of 8.5 months (range, 2-30 months) after auto-HSCT. The occurrence of a secondary autoimmune disorder and the possible contributory factors are summarized in Table 1. The secondary autoimmune conditions manifested were acquired factor VIII inhibitors (n = 2), autoimmune thrombocytopenia (n = 2), autoimmune hemolytic anemia (AIHA; n = 1), and autoimmune neutropenia/AIHA (n = 1). Their presentation, management, and outcomes are summarized in Table 2.

Table 1. Occurrence of secondary autoimmune disorders and possible risk factors

<table>
<thead>
<tr>
<th>Possible risk factors</th>
<th>Affected by secondary autoimmune disorder, no.</th>
<th>Unaffected by secondary autoimmune disorder, no.</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Conditioning regimen</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cy/ATG</td>
<td>2</td>
<td>100</td>
<td></td>
</tr>
<tr>
<td>Cy/alemtuzumab</td>
<td>4</td>
<td>21</td>
<td></td>
</tr>
<tr>
<td>Cy/TBI or Bu/Cy</td>
<td>0</td>
<td>28</td>
<td>.011</td>
</tr>
<tr>
<td><strong>Sex</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>6</td>
<td>106</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>0</td>
<td>43</td>
<td>.19</td>
</tr>
<tr>
<td><strong>Primary disease</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Systemic lupus erythematosus</td>
<td>3</td>
<td>57</td>
<td></td>
</tr>
<tr>
<td>Multiple sclerosis</td>
<td>2</td>
<td>41</td>
<td></td>
</tr>
<tr>
<td>Systemic sclerosis</td>
<td>1</td>
<td>12</td>
<td></td>
</tr>
<tr>
<td>Crohn disease</td>
<td>0</td>
<td>19</td>
<td></td>
</tr>
<tr>
<td>Rheumatoid arthritis</td>
<td>0</td>
<td>6</td>
<td></td>
</tr>
<tr>
<td>Others</td>
<td>0</td>
<td>14</td>
<td>ND</td>
</tr>
<tr>
<td><strong>Type of ATG</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rabbit</td>
<td>0</td>
<td>51</td>
<td></td>
</tr>
<tr>
<td>Equine</td>
<td>2</td>
<td>49</td>
<td>.50</td>
</tr>
<tr>
<td><strong>PBSC product</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>CD34 selected</td>
<td>2</td>
<td>85</td>
<td></td>
</tr>
<tr>
<td>Unmanipulated</td>
<td>4</td>
<td>64</td>
<td>.41</td>
</tr>
</tbody>
</table>

Two patients developed a secondary acquired factor VIII inhibitor. The first was a 36-year-old female with refractory SLE with multisystem involvement including antiphospholipid syndrome (APS) with recurrent venous thromboembolism. She underwent auto-HSCT after conditioning with cyclophosphamide and equine ATG uneventfully, achieving both clinical and serologic remission from both SLE and APS, permitting discontinuation of anticoagulation and immunosuppression by 7 months. At 30 months after auto-HSCT, she developed severe postoperative bleeding following umbilical hernia repair and insertion of a peritoneal dialysis catheter, developing large intra-abdominal and retroperitoneal hematomas. Investigations revealed a partial thromboplastin time (PTT) exceeding 150 seconds, a factor VIII activity of less than 1%, and a factor VIII inhibitor at a level of 16.4 Bethesda units. Her lupus anticogulant and anticardiolipin antibodies remained negative. The patient was treated with activated factor VII, prednisone, and rituximab and achieved remission from acquired hemophilia.

The second patient was a 28-year-old female with severe SLE, who was refractory to numerous immunosuppressants. She underwent auto-HSCT after conditioning with cyclophosphamide and
Secondary autoimmune cytopenias

Four patients developed secondary autoimmune cytopenias. The first was a 46-year-old female with relapsing-remitting MS failing interferon therapy. She underwent auto-HSCT after cyclophosphamide and alemtuzumab (20 mg). She achieved platelet engraftment by day +8 and remained well until 8 months after transplantation when she presented with diffuse petechiae. Her platelet counts fell to 2 × 10^9/L with mild anemia (hemoglobin 100 g/L) and leukopenia. A bone marrow aspiration and biopsy revealed normal cellularity and maturation in all cell lineages. The direct antiglobulin test (DAT) was positive with biochemical evidence of hemolysis. She was treated with intravenous immunoglobulin (IVIG) and methylprednisolone, mycophenolate mofetil (MMF), and prednisone. The factor VIII inhibitor cleared within several months but low-grade lupus activity has persisted.

Table 2. Summary of patients who developed a secondary autoimmune disorder

<table>
<thead>
<tr>
<th>Primary disease</th>
<th>Regimen</th>
<th>CD34-selected PBSCs</th>
<th>Current status of primary disease</th>
<th>Secondary autoimmune disorder (time of onset)</th>
<th>Treatment of secondary disorder</th>
<th>Outcome of secondary disorder</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 SLE</td>
<td>Cy/ATG</td>
<td>Yes</td>
<td>Remission</td>
<td>FVIII inhibitor (30 mo)</td>
<td>FVIIa, RTX, pred</td>
<td>Resolved</td>
</tr>
<tr>
<td>2 SLE</td>
<td>Cy/ATG</td>
<td>Yes</td>
<td>Persistent</td>
<td>FVIII inhibitor (9 mo)</td>
<td>FVIIa, MP, RTX, Cy, MMF</td>
<td>Resolved</td>
</tr>
<tr>
<td>3 MS</td>
<td>Cy/alemtuzumab</td>
<td>No</td>
<td>Remission</td>
<td>ITP (8 mo)</td>
<td>IVIG, MP, RTX, pred</td>
<td>Resolved</td>
</tr>
<tr>
<td>4 SSc</td>
<td>Cy/alemtuzumab</td>
<td>No</td>
<td>Remission</td>
<td>AIHA (5 mo)</td>
<td>IVIG, MP, MMF, pred, RTX, Cy</td>
<td>Controlled</td>
</tr>
<tr>
<td>5 SLE</td>
<td>Cy/alemtuzumab</td>
<td>No</td>
<td>Remission</td>
<td>Autoimmune neutropenia/AIHA (2 mo)</td>
<td>IVIG, MP, RTX, Cy, MMF, pred,</td>
<td>Controlled</td>
</tr>
<tr>
<td>6 MS</td>
<td>Cy/alemtuzumab</td>
<td>No</td>
<td>Remission</td>
<td>ITP (14 mo)</td>
<td>IVIG, MP, RTX, MMF</td>
<td>Resolved</td>
</tr>
</tbody>
</table>

PBSCs indicates peripheral blood stem cells; HSCT, hematopoietic stem cell transplantation; SLE, systemic lupus erythematosus; MS, multiple sclerosis; SSc, systemic sclerosis; Cy, cyclophosphamide; ATG, antithymocyte globulin; FVIII, factor VIII; ITP, immune thrombocytopenia purpura; AIHA, autoimmune hemolytic anemia; FVIIa, recombinant activated factor VII; RTX, rituximab; pred, prednisone; MP, methylprednisolone; MMF, mycophenolate mofetil; and IVIG, intravenous immunoglobulin.

equine ATG; SLE disease activity remained persistent after the transplantation, albeit reduced in severity. At 9 months after auto-HSCT, she suffered a spontaneous abortion complicated by severe bleeding after evacuation of the uterus. She also developed multiple spontaneous bruises and hematomas on her upper and lower extremities. Her PTT exceeded 150 seconds, factor VIII activity was below 1%, and the factor VIII inhibitor was 1600 Bethesda units. She received activated factor VII which arrested the bleeding, and was also treated with rituximab, intravenous cyclophosphamide, mycophenolate mofetil (MMF), and prednisone. The factor VIII inhibitor cleared within several months but low-grade lupus activity has persisted.

**Discussion**

This study describes for the first time the development of a secondary autoimmune complication among patients undergoing auto-HSCT for an autoimmune condition, and possible contributory factors involved. The limitation of this study is its retrospective and nonrandomized nature, relatively small numbers, and its involvement of patients with different autoimmune diseases recruited under several different disease-specific protocols. It does, however, represent the largest single-center experience of auto-HSCT for autoimmune diseases, with the benefit of standardization of procedures, evaluation, and management. The timing of follow-up varies, as the regimens have evolved with time. Our first patient to receive an alemtuzumab-based regimen was in 2003, whereas our experience with ATG- or TBI-based regimens dates from 1996. It is noteworthy therefore that, despite a shorter duration of follow-up, we have seen a higher frequency of secondary autoimmunity among the alemtuzumab group, an observation that prompted this retrospective analysis. Among the factors analyzed, only alemtuzumab exposure had a statistically significant association with this complication.
The patients described in our series developed second autoimmune disorders distinct from their primary disease, and which are not usually associated with the underlying disorder. The onset of the complication was at a median of 8.5 months after undergoing transplantation, occurring as late as 30 months in the case of the first patient with SLE who developed acquired hemophilia. While autoimmune cytopenias are common manifestations of SLE, the fifth patient described never had significant cytopenia prior to auto-HSCT. In addition, with the exception of the second case, the patients developed the disorders despite achieving remission from the primary autoimmune disease. The occurrence of more than one autoimmune disorder in susceptible individuals has been well described.\textsuperscript{14} This results from a genetic predisposition to autoimmune disease, which is supported by epidemiologic and genetic evidence.\textsuperscript{6,7} We believe, however, that the secondary autoimmune disorders in our patients occurred because of a combination of factors: an underlying propensity to autoimmunity, the immune dysregulation/reset resulting from the auto-HSCT, and the conditioning regimen drugs used.

Autoimmune diseases developing after both autologous and allogeneic HSCT for hematologic malignancies have been reported to occur at a frequency of between 2\% and 5\%.\textsuperscript{15-17} The spectrum of autoimmune disorders includes autoimmune cytopenias most frequently, but has also included autoimmune thyroiditis,\textsuperscript{18} myasthenia gravis,\textsuperscript{19} rheumatoid arthritis,\textsuperscript{20} and a lupuslike illness.\textsuperscript{2} Acquired hemophilia due to factor VII antibodies after HSCT has been described in 8 patients,\textsuperscript{21} but acquired hemophilia due to factor VIII inhibitors has been reported only once to our knowledge, having occurred in a patient with MS after auto-HSCT during which he received conditioning with ATG.\textsuperscript{22} The authors of the latter report concluded the coexistence of 3 factors (underlying autoimmunity, auto-HSCT, and use of interferon-beta) predisposed the patient to developing this autoimmune complication.

Acquired hemophilia due to factor VIII inhibitors is a rare condition with a reported incidence of only one person per million per year.\textsuperscript{23} It is usually idiopathic, although the association with autoimmune disorders such as SLE and rheumatoid arthritis has been described.\textsuperscript{24} In one of the largest series of acquired hemophilia published to date, with 215 patients, only 4\% of the cohort had coexistent SLE.\textsuperscript{25} The occurrence of this rare disorder with an infrequent association with SLE among 2 of our patients is thus notable. While acquired hemophilia is associated with pregnancy, it is usually mild in this subgroup of patients, with a median inhibitor titer of 20 Bethesda units, and spontaneous remissions often occur.\textsuperscript{26,27} In contrast, our patient described in the second case had an inhibitor titer of 1600 Bethesda units and required multiple immunosuppressive agents to achieve control. This presentation, although perhaps exacerbated by pregnancy, is not typical of a case of postpartum acquired hemophilia.

The development of autoimmunity following autologous and allogeneic HSCT is postulated to be due to the immunologic dysregulation resulting from the transplant.\textsuperscript{28} While high-dose immunosuppression and auto-HSCT have been employed in a variety of autoimmune disorders with varied success, the same process of immune ablation may alter the regulatory mechanisms within the immunologic system keeping further autoimmunity in check. The role of regulatory T cells in maintaining self-tolerance has been demonstrated by the development of autoimmunity in thymectomized animals,\textsuperscript{29} with restoration of self-tolerance through reconstitution with CD4\(^+\)CD25\(^+\) splenocytes.\textsuperscript{30} The role of these CD25\(^+\) regulatory T cells in maintaining tolerance is now well established.\textsuperscript{31} These T-cell homeostatic mechanisms are also altered during lymphopenia, with T-cell clones proliferating in response to self-antigens in the lymphopenic host.\textsuperscript{32} The rate of immune reconstitution following transplantation of the various lymphocyte subsets differs, with B-cell recovery preceding T cell, and CD8 subsets occurring earlier than CD4.\textsuperscript{33} These altered balances affect the immune rheostat and may set the stage for the emergence of autoimmune cytopenias and immune-mediated factor VIII deficiency. An analysis of T-cell subsets among our patients would serve to clarify the role regulatory T cells may play in the pathogenesis of secondary autoimmune disorders after transplantation.

In auto-HSCT for autoimmune diseases, the regimens chosen are selected specifically for lympho-depletion, with the aim of eliminating auto-reactive T-cell clones in the patient. Drugs like ATG and alemtuzumab are thus used for their lymphotoxicity. ATG induces a rapid and profound lymphopenia attributed to complement- and antibody-dependent cytolyis and macrophage phagocytosis\textsuperscript{34} and apoptosis of T cells.\textsuperscript{35} The effect of ATG on immune function has also been shown to persist for up to 12 months.\textsuperscript{36,37} The net result is a lasting quantitative and qualitative defect in T-cell function. There have been several reports of various autoimmune disorders precipitated by antithymocyte or antilymphocyte globulin including thyroid disease,\textsuperscript{38} Guillain-Barre syndrome,\textsuperscript{39} immune hemolysis,\textsuperscript{40} and alveolitis.\textsuperscript{41} There was a case of a patient who developed asymptomatic factor XII inhibitors after allogeneic HSCT with an ATG-containing regimen for acute myeloid leukemia.\textsuperscript{42} To our knowledge, the only other report of acquired hemophilia secondary to factor VIII inhibitors after ATG involved a patient with MS who underwent auto-HSCT after busulphan and ATG conditioning.\textsuperscript{22}

Alemtuzumab is a monoclonal antibody directed against CD52 which is expressed on virtually all T and B lymphocytes (although preferentially on T lymphocytes)\textsuperscript{43} and monocytes.\textsuperscript{44} It has the ability to induce a rapid and profound lymphopenia following a single administered dose.\textsuperscript{45} This lymphopenia persists beyond 6 months and the effect on the CD4\(^+\) and CD8\(^+\) subsets, compared with ATG, appears to be even more protracted, with some studies demonstrating decreased T-cell subsets beyond 18 months.\textsuperscript{46}

The lymphopenia effected by alemtuzumab, with decreased CD4\(^+\)/CD25\(^+\) regulatory T cells, may predispose to the development of autoimmunity. In addition, CD52 antigens are also expressed on monocyte-derived dendritic cells and thus antigen presentation in the host is affected following alemtuzumab exposure.\textsuperscript{47,48} Immature peripheral dendritic cells present antigen to T cells in a tolerogenic manner by silencing activated T cells, activating and causing expansion of regulatory T cells, and causing the differentiation of naive CD4\(^+\) cells to regulatory T cells.\textsuperscript{49,50} The effect of alemtuzumab in depleting these dendritic cells may therefore predispose to the development of autoimmunity. Furthermore, alemtuzumab has been shown to deplete natural killer cells; studies suggest these cells may play a role in both exacerbating and protecting against autoimmune mechanisms.\textsuperscript{51} It can be seen, therefore, that alemtuzumab induces a wide range of perturbations in the various components of the adaptive and innate immune system, from which recovery and reconstitution occur within different time frames and to different extents. This immune dysregulation may set the stage for the development of autoimmune disorders.

There have been several reports of autoimmune disorders occurring after therapy with alemtuzumab. In a report of patients with MS given the drug, 32\% developed Grave disease. The authors concluded that while the drug suppressed MS disease
activity, it permitted the development of autoimmunity, perhaps due to delayed CD4+ T-cell recovery. There have been 2 reports of fatal refractory ITP after treatment of chronic lymphocytic leukemia with alemtuzumab. In 3 cases, ITP (one case fatal) occurred during a clinical trial of alemtuzumab for MS. As with the published reports, we have found the autoimmune cytopenias developing after auto-HSCT and auto-ITP to be generally severe and refractory to steroid therapy alone. All our patients required multiple agents to treat the cytopenias, and demonstrated a response to rituximab, which appears to be an effective agent in refractory cases.

Interestingly, alemtuzumab has also been employed in the treatment of refractory autoimmune cytopenias with responses seen in two-thirds of patients, although a third subsequently relapsed. The drug also has demonstrated activity in treating MS, but the occurrence of autoimmune thyroid disease and severe delayed ITP has dampened enthusiasm toward its use in MS. As with 2 patients with MS who developed ITP at 8 and 14 months after transplantation, respectively, the development of ITP among the patients who received alemtuzumab for MS was relatively delayed and manifested up to one year after treatment. It is therefore essential to continue monitoring the blood cell counts in these patients beyond a year. Alemtuzumab thus appears to be a somewhat double-edged sword in autoimmunity: its effective long-term lympho-depletion removes auto-reactive T-cell clones, whereas the loss of T-cell homeostasis may favor subsequent development of another autoimmune condition.

Alemtuzumab has been used increasingly in solid organ and hematopoietic stem cell transplantation, and in treating lymphoid malignancies; nevertheless, the reports of autoimmune complications have been relatively few. In fact, in a long-term study of immune reconstitution in patients with chronic lymphocytic leukemia who had been treated with alemtuzumab, there was no reported increase in autoimmunity. In contrast, the occurrence of autoimmune complications among patients with underlying autoimmune disorders receiving alemtuzumab appears to be increased.

Among the 155 patients with autoimmune diseases who underwent transplantation at our institution thus far, the frequency of secondary autoimmune complications was 16.0% with alemtuzumab (4/25), 1.9% for ATG (2/102), and 0% for conditioning regimens without lympho-depleting antibodies (0/28), a difference that was found to be significantly higher with alemtuzumab exposure. In the interest of safety, therefore, we have ceased to use alemtuzumab in our conditioning regimens for auto-HSCT.

Growing evidence supports the potential of auto-HSCT in controlling severe autoimmune diseases, but a second autoimmune disorder may complicate the procedure. The underlying susceptibility of an “autoimmune-prone” patient, coupled with the immune dysregulation resulting from auto-HSCT and lympho-depleting agents used in the conditioning regimen, may set the stage for immune-mediated cytopenias and factor VIII deficiency, especially with long-term lympho-depleting agents such as alemtuzumab. With the increasing employment of auto-HSCT in the treatment of autoimmune diseases, physicians need to be alert to the occurrence of this complication. The increased likelihood of a secondary autoimmune disorder with alemtuzumab recommends against using it in autologous HSCT regimens for autoimmune diseases.

Authorship

Contribution: Y.L. collected and analyzed data, reviewed literature, and wrote and revised the paper; R.B. and Y.O. designed and formulated research, and revised the paper; B.J. performed statistical analysis; Y.O., L.S., K.Q., K.Y., E.G., R.C., D.S., B.C., W.B., Y.L., and R.B. performed the study and collected data.

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