Abstract: Mycosis Fungoides is typically an indolent disease in early stages. However, approximately 30% of patients have advanced staged disease at presentation and 20% will develop it at some time. These patients have a poorer prognosis with a median survival of 2-4 years. The only curative option for mycosis fungoides may be hematopoietic allogeneic stem cell transplantation. We report the case of a patient with mycosis fungoides in an advanced stage (IIB), refractory to treatment options. She underwent allogeneic hematopoietic stem-cell transplantation (allo-HSCT). The patient remains in complete remission nineteen months after allo-HSCT. Allogeneic transplantation can alter the natural history of mycosis fungoides and should be considered in patients who have refractory disease or short-lived responses with standard therapies.

Keywords: Hematopoietic stem cell transplantation; Lymphoma, T-Cell, cutaneous; mycosis fungoides

INTRODUCTION

Mycosis fungoides (MF) is the most common subtype of primary cutaneous T-cell lymphomas, with an annual incidence of 0.36 cases per 100,000 person-years.1,2 MF affects men twice as often as women, is more common in black people, and manifests with higher incidence over the age of 50.3 The prognosis for MF depends on the age and clinical stage of the patient.4

It is characterized initially by scaly, non-infiltrative, erythematous or hypopigmented plaques, persistent, of varying sizes, and featuring intense itching. In the advanced stages (> or = IIB) it becomes deeply infiltrated, ulcerated, violet staining, with nodules and tumors appearing that can become infected. Belatedly it can spread to lymph nodes and visceral organs.5,6 Approximately 30% of patients have advanced staged disease with generalized erythroderma or tumor at the time of presentation, and 20% will develop advanced disease.7,8 The correct diagnosis of MF can take years because of its similarity to other benign skin conditions. In the USA, it can take about six years from the time symptoms begin until the
diagnosis of MF, and some patients require multiple biopsies over the years.1

The chronic nature of the disease results in many patients being treated with multiple therapies, including: PUVA, radiotherapy, corticosteroids, alpha-interferon, chemotherapy, and immunotherapy. Patients with advanced disease have a median survival of only two to four years; refractory patients are considered incurable with chemotherapy alone and most of these therapies are palliative.3,6,7

There is hope that Hematopoietic Stem Cell Transplants (HSCT) have the potential to provide prolonged remissions or a possible cure for advanced MF; however, their role is not well defined.6-8 We report the case of a female patient with MF, with approximately 13 years of evolution and subjected to several treatments but obtaining low response. The advanced stage and the lack of therapeutic options had led to an indication of allogeneic HSCT (allo-HSCT). The effect of HSCT in the patient’s illness in a follow-up of 19 months is present here.

**CASE REPORT**

A 44-year-old white woman observed an erythema in the sternal region that progressed to involvement of the whole body in the form of urticarial plaques. The picture remained stable with topical and systemic steroids for eight years. Progression was observed with elevated and pruritic patches and plaques with purulent discharge from secondary infection on an extensive portion of the body surface.

When the patient came to our service, we performed a skin biopsy with immunohistochemistry that revealed cutaneous lymphoproliferative disease compatible with anaplastic large T-cell lymphoma focally CD30+ or Histiocytic sarcoma. Bone marrow biopsy was normal. CHOEP (cyclophosphamide, doxorubicin, etoposide, vincristine and prednisone) chemotherapy was initiated, totaling eight cycles, with minimal initial response in plaques and pruritus, but reappearing at the end of the eighth cycle. A new skin biopsy showed cutaneous T-cell lymphoma with epidermotropism, CD45+ and CD3+ (suggesting MF). Staged as IIB (T3-N0-M0), with 80% affected area, and presenting the most advanced lesions in the trunk, breasts and legs.

Subcutaneous interferon 3,000,000U was initiated, three times a week, gradually increasing to 12,000,000U and radiotherapy in tougher skin areas in the right groin. Major improvement of the lesions was obtained with interferon, but this had to be suspended because of non-hematological toxicity and disease progression. Gemcitabine, 1000mg/m² per week, was initiated. There was a significant regression of lesions, but recurrence was noticed at the end of the sixth cycle.

Her condition progressively worsened despite partial and temporary improvements with the various treatment modalities. The patient’s skin was diffusely infiltrated (Figures 1 and 2) with patches, plaques, and tumors. The patient was referred to Euryclides de Jesus Zerbini Transplants Hospital, in São Paulo, and a reduced-intensity conditioning (RIC) allogeneic HLA matched sibling transplant was performed. After conditioning with fludarabine and melfalan she received peripheral blood stem-cells from her brother. Graft-versus-host-disease (GVHD) prophylaxis was done with mycophenolate mofetil and cyclosporine.

Engraftment took place on D+11 and the patient was discharged from the hospital on D+14, with complete remission. Skin biopsy of residual hyperpigmented patches (Figure 3) showed GVHD without evidence of lymphocyte atypia. The patient has been in complete remission (Figures 4 and 5) for more than one year without treatment or any signs of GVHD, and peripheral blood mononuclear cells are 100% donor.
DISCUSSION

Advanced MF is associated with a poor outcome. Allo-HSCT has proved to be an effective therapy in MF, demonstrating a decrease in the relapse rate and an overall increase in disease-free survival compared with conventional therapy. Allogeneic transplantation from a healthy donor not only eliminates the potential of graft contamination by tumor cells but also provides an immunologic antitumor effect, known as graft-versus-lymphoma (GVL).6,9 Although only a few patients have been treated with allo-HSCT until now, the results are consistent and promising.6,7,9,10

In the Wu et al. meta-analysis, 70% of patients undergoing allo-HSCT exhibited GVHD, the majority being mild to moderate, particularly in skin tissue.10 In the Molina et al. series of transplants, complete remission of skin lesions, tumors and ulcerated areas, itching and pain was observed in 100% of patients after allo-HSCT.8 The results of Duarte et al. showed that one year after allo-HSCT, 42% of their patients remained in remission.9 Our patient has achieved early remission and is maintaining it for over a year after transplantation. More than two thirds of patients with advanced stage CTCL, after receiving an allo-HSCT, were alive and without evidence of disease for more than three years, which appears to be substantially higher than the median for disease-free survival, or even the overall rate for such patients.9

Most patients undergoing allo-HSCT previously received an average of seven different treatments before transplantation, similar to our patient who received five. However, despite these values, the optimal time for transplantation remains unknown; it is expected that, in the future, morbidity and the risk of recurrence may be even lower. Nevertheless, it is unquestionable that the allo-HSCT has a role of great importance in the management of CTCL, especially MF. The results observed in our patients are encouraging, and further studies are needed to answer the questions that still remain. 

FIGURE 3: Graft-versus-host-disease - hyperchromic macules on the back

FIGURE 4: Remission of mycosis fungoides lesions after allogeneic hematopoietic stem cell transplantation

FIGURE 5: Remission of mycosis fungoides lesions after allogeneic hematopoietic stem cell transplantation
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