Sweet Syndrome Associated With Myelodysplastic Syndrome: Report of a Case. Review of the Literature

Delia Reina,1,∗ Dacia Cerdà, a Daniel Roig, a Ramon Fíguls, a M. Luz Villegas, b Hèctor Corominas a

a Unidad de Reumatología, Hospital Sant Joan Despí Moisés Broggi, Sant Joan Despí, Barcelona, Spain
b Servicio de Medicina Interna, Hospital General de l’Hospitalet, Barcelona, Spain

Case report

Sweet Syndrome Associated With Myelodysplastic Syndrome: Report of a Case. Review of the Literature

Sweet’s syndrome or acute neutrophilic febrile dermatosis is a systemic disease of unknown etiology characterized by the appearance of skin lesions produced by a neutrophilic dermal infiltrate, fever, and peripheral leukocytosis. It may be associated with hematologic diseases, including leukemia, with immune diseases as rheumatoid arthritis, or can occur in isolation. The myelodysplasias are hematological disorders characterized by one or more cytopenias secondary to bone marrow dysfunction. We present the case of a patient with Sweet’s syndrome associated with myelodysplastic syndrome and treated with glucocorticoids who did not present a good clinical outcome. We discuss the different treatment of these diseases because in most cases glucocorticoids, which are the treatment of choice in Sweet’s syndrome, may be insufficient.

Introduction

Sweet’s syndrome (SS) is a skin disease of unknown etiology1,2 which is characterized by fever, leukocytosis, and painful erythematous skin plaques. Histology shows dense neutrophilic infiltrates. The presence of anemia and thrombocytopenia may be associated with an underlying neoplastic.3

We present the case of a patient with SS and anemia diagnosed as a myelodysplastic syndrome (MDS), worsening the prognosis. MDS are hematologic diseases characterized by cytopenias that result in dysmorphic states of the cells of the bone marrow. One type of MDS is refractory anemia with blast excess (RABE), with the number of blasts being greater than 5% and less than 10% in type 1, and between 10% and 20% in type 2. These diseases are refractory to chemotherapy and stem cell transplant offers a cure. In RABE, supportive treatment is performed with red blood cell transfusions.

Case Report

The patient is a 70 years old male with no history of interest, who had recurrent episodes of 2 weeks with high fever, leukocytosis, and...
The patient described had a case that progressed despite treatment. Pending the transplantation we were able to use some of the treatments described in the literature, although possibly the transplant would had been the only effective treatment.

Conclusion

SS is a rare dermatological entity, which when accompanied by an MDS may worsen the patient’s prognosis. The treatment of choice for SS is the use of glucocorticoids, although in most cases this may be insufficient and require alternative therapies.

Ethical disclosures

Protection of human and animal subjects. The authors declare that no experiments were performed on humans or animals for this investigation.

Confidentiality of Data. The authors declare that they have followed the protocols of their work centre on the publication of patient data and that all the patients included in the study have received sufficient information and have given their informed consent in writing to participate in that study.

Right to privacy and informed consent. The authors have obtained the informed consent of the patients and / or subjects mentioned in the article. The author for correspondence must be in possession of this document.

Disclosure

The authors have no disclosures to make.

References