

MEETING ABSTRACT

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Severe combined immunodeficiency: case report of allogenic, haploidentical bone marrow transplantation

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From 3rd WAO International Scientific Conference (WISC) 2014
Rio de Janeiro, Brazil. 6-9 December 2014

Background

Severe combined immunodeficiency (SCID) is characterized by important impairment in differentiation of T and/or B lymphocytes and occasionally Natural Killer cells, representing a pediatric emergency. A case of immunodeficiency is described emphasizing symptoms, diagnosis and answer after bone marrow transplantation.

Methods

Case report of a 2 years old male patient with severe combined immunodeficiency (SCID), diagnosed at 9 months after hospitalization due to failure to thrive, chronic diarrhea and pneumonia. Evolved with recurrent respiratory and gastrointestinal infections although using prophylaxis and immunoglobulin infusion. Allogenic, haploidentical transplantation was carried out with positive selection of CD34+, at 18 months of age due no compatible donor been found.

Results

Satisfactory answer after transplantation keeping infusion of IV immunoglobulin with clinical and laboratorial favorable evolution.

Conclusions

Bone marrow transplantation when succeeds is supposed to restore lymphocyte system diminishing risks of severe and fatal infections.

Consent

Written informed consent was obtained from the patient for publication of this abstract and any accompanying images. A copy of the written consent is available for review by the Editor of this journal.

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Published: 8 April 2015

doi:10.1186/1939-4551-8-S1-A122

Cite this article as: Silveira et al.: Severe combined immunodeficiency: case report of allogenic, haploidentical bone marrow transplantation. *World Allergy Organization Journal* 2015 8(Suppl 1):A122.

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