



PEER-REVIEW REPORT

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Manuscript NO: 57284

Title: Nephrotic syndrome in syngeneic hematopoietic stem cell transplantation recipients: A case report

Reviewer's code: 03850089

Position: Editorial Board

Academic degree: MD, PhD

Professional title: Emeritus Professor

Reviewer's Country/Territory: Brazil

Author's Country/Territory: China

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Reviewer chosen by: Pan Huang

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Scientific quality	<input type="checkbox"/> Grade A: Excellent <input checked="" type="checkbox"/> Grade B: Very good <input type="checkbox"/> Grade C: Good <input type="checkbox"/> Grade D: Fair <input type="checkbox"/> Grade E: Do not publish
Language quality	<input type="checkbox"/> Grade A: Priority publishing <input checked="" type="checkbox"/> Grade B: Minor language polishing <input type="checkbox"/> Grade C: A great deal of language polishing <input type="checkbox"/> Grade D: Rejection
Conclusion	<input type="checkbox"/> Accept (High priority) <input checked="" type="checkbox"/> Accept (General priority) <input type="checkbox"/> Minor revision <input type="checkbox"/> Major revision <input type="checkbox"/> Rejection
Re-review	<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No
Peer-reviewer statements	Peer-Review: <input checked="" type="checkbox"/> Anonymous <input type="checkbox"/> Onymous Conflicts-of-Interest: <input type="checkbox"/> Yes <input checked="" type="checkbox"/> No



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SPECIFIC COMMENTS TO AUTHORS

Manuscript 57284 - WJCC In this report, the authors present a case of a patient with T lymphoblastic lymphoma who developed a nephrotic syndrome two years after a syngeneic, allogeneic transplant. The description follows the CARE guidelines, is well documented, and presents correct diagnostic evidence that justifies the diagnosis of Autologous GVHD. The treatment effectively resolved the renal complication, and the discussion advances several aspects related to renal complications secondary to the allogeneic transplant matched sibling donors, or not related. It highlights B cells' role in the genesis of GVHD and autoimmune diseases and punctuates the situation of Autologous GVHD. It concludes by pointing out the causes and possible mechanisms of the nephrotic syndrome's appearance in that case. The references in number 24 are current and support the full description. Supplementary files follow the magazine's rules. Due to the rarity and absence of similar information in the literature, the suggestion is that the article has a quality for publication