



ESPS PEER-REVIEW REPORT

Name of journal: World Journal of Hematology

ESPS manuscript NO: 27283

Title: Successful lower leg microsurgical reconstruction in homozygous sickle cell disease: Case report

Reviewer's code: 03638253

Reviewer's country: United States

Science editor: Xue-Mei Gong

Date sent for review: 2016-05-23 15:08

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CLASSIFICATION	LANGUAGE EVALUATION	SCIENTIFIC MISCONDUCT	CONCLUSION
<input type="checkbox"/> Grade A: Excellent	<input type="checkbox"/> Grade A: Priority publishing	Google Search:	<input type="checkbox"/> Accept
<input checked="" type="checkbox"/> Grade B: Very good	<input checked="" type="checkbox"/> Grade B: Minor language polishing	<input type="checkbox"/> The same title	<input type="checkbox"/> High priority for publication
<input type="checkbox"/> Grade C: Good	<input type="checkbox"/> Grade C: A great deal of language polishing	<input type="checkbox"/> Duplicate publication	<input type="checkbox"/> Rejection
<input type="checkbox"/> Grade D: Fair	<input type="checkbox"/> Grade D: Rejected	<input checked="" type="checkbox"/> No	<input checked="" type="checkbox"/> Minor revision
<input type="checkbox"/> Grade E: Poor		BPG Search:	<input type="checkbox"/> Major revision
		<input type="checkbox"/> The same title	
		<input type="checkbox"/> Duplicate publication	
		<input type="checkbox"/> Plagiarism	
		<input checked="" type="checkbox"/> No	

COMMENTS TO AUTHORS

This is a case report of utilizing a novel surgical technique for management of ulcers related to sickle cell disease. Historically the outcome of such surgeries in patients with sickle cell disease has not been favorable. The author suggests more aggressive anticoagulation and lowering the hemoglobin S levels in blood circulation around the surgery time and was able to achieve better results with such measures. The manuscript is well written and I suggest a few minor changes as follows: 1) In Introduction: Please provide reference for the statement: "Patients with hematologic diseases and hypercoagulability are usually associated with high rates of complications, including anastomotic thrombosis and flap loss." 2) Discussion: Change the statement: "Sickle cell trait [SCT] results from the inheritance of one normal hemoglobin gene [HbA] and one mutated b1-globin gene , a sickle hemoglobin gene [HbS]." to "Sickle cell.....mutated Beta Globin Gene...HbS." 3) Provide reference for the statement: "Increased erythrocyte sickling occurs under hypoxia, acidosis, dehydration, and hypothermia. SCT has been associated with an increased rate of exercise-related deaths, fetal loss, pre-eclampsia, and venous thromboembolism" 4)Provide reference for the statement: "Patients



BAISHIDENG PUBLISHING GROUP INC

8226 Regency Drive, Pleasanton, CA 94588, USA

Telephone: +1-925-223-8242

Fax: +1-925-223-8243

E-mail: bpgoffice@wjgnet.com

<http://www.wjgnet.com>

with sickle cell trait or disease are usually considered a higher risk group for microsurgical reconstructions and there is some pessimism among microsurgeons when these procedures are required" 5) In the manuscript it is suggested to bring the Hemoglobin S level down to 30% but in the Table 1 it says 20%. I suggest you change it to 30% as well and provide reference for that.



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		<input type="checkbox"/> Duplicate publication	
		<input type="checkbox"/> Plagiarism	
		<input checked="" type="checkbox"/> No	

COMMENTS TO AUTHORS

This is an interesting and well written clinical case report. I would recommend publication, with suggested minor revision: check through the paper for typos and grammatical errors, and secondly revise Table 1 - it would be presented better as a flow chart and with text of larger font to make it easier to read.