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ESPS PEER-REVIEW REPORT

Name of journal: World Journal of Nephrology
ESPS manuscript NO: 25834
Title: New SLC12A3 disease causative mutation of Gitelman's syndrome
Reviewer's code: 00503255
Reviewer's country: Japan
Science editor: Xue-Mei Gong
Date sent for review: 2016-04-15 10:12
Date reviewed: 2016-04-15 11:58

CLASSIFICATION	LANGUAGE EVALUATION	SCIENTIFIC MISCONDUCT	CONCLUSION
[ ] Grade A: Excellent	[Y] Grade A: Priority publishing	Google Search:	[Y] Accept
[Y] Grade B: Very good	[ ] Grade B: Minor language polishing	[ ] The same title	[ ] High priority for publication
[ ] Grade C: Good	[ ] Grade C: A great deal of language polishing	[ ] Duplicate publication	[ ] Rejection
[ ] Grade D: Fair	[ ] Grade D: Rejected	[ ] Plagiarism	[ ] Minor revision
[ ] Grade E: Poor		[Y] No	[ ] Major revision
		BPG Search:	
		[ ] The same title	
		[ ] Duplicate publication	
		[ ] Plagiarism	
		[Y] No	

COMMENTS TO AUTHORS

The authors described brothers with Gitelman syndrome who had a new mutation of SLC12A3. The paper is well-written and interesting findings.

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ESPS PEER-REVIEW REPORT

Name of journal: World Journal of Nephrology
ESPS manuscript NO: 25834
Title: New SLC12A3 disease causative mutation of Gitelman's syndrome
Reviewer's code: 00503254
Reviewer's country: Japan
Science editor: Xue-Mei Gong
Date sent for review: 2016-04-15 10:12
Date reviewed: 2016-04-22 09:29

CLASSIFICATION	LANGUAGE EVALUATION	SCIENTIFIC MISCONDUCT	CONCLUSION
[Y] Grade A: Excellent	[ ] Grade A: Priority publishing	Google Search:	[Y] Accept
[ ] Grade B: Very good	[Y] Grade B: Minor language polishing	[ ] The same title	[ ] High priority for publication
[ ] Grade C: Good	[ ] Grade C: A great deal of language polishing	[ ] Duplicate publication	[ ] Rejection
[ ] Grade D: Fair	[ ] Grade D: Rejected	[ ] Plagiarism	[ ] Minor revision
[ ] Grade E: Poor		[Y] No	[ ] Major revision
		BPG Search:	
		[ ] The same title	
		[ ] Duplicate publication	
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		[Y] No	

COMMENTS TO AUTHORS

None

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ESPS PEER-REVIEW REPORT

Name of journal: World Journal of Nephrology
ESPS manuscript NO: 25834
Title: New SLC12A3 disease causative mutation of Gitelman's syndrome
Reviewer's code: 00631992
Reviewer's country: Italy
Science editor: Xue-Mei Gong
Date sent for review: 2016-04-15 10:12
Date reviewed: 2016-05-05 06:03

CLASSIFICATION	LANGUAGE EVALUATION	SCIENTIFIC MISCONDUCT	CONCLUSION
[ ] Grade A: Excellent	[Y] Grade A: Priority publishing	Google Search:	[ ] Accept
[Y] Grade B: Very good	[ ] Grade B: Minor language polishing	[ ] The same title	[ ] High priority for publication
[ ] Grade C: Good	[ ] Grade C: A great deal of language polishing	[ ] Duplicate publication	[ ] Rejection
[ ] Grade D: Fair	[ ] Grade D: Rejected	[ ] Plagiarism	[Y] Minor revision
[ ] Grade E: Poor		[Y] No	[ ] Major revision
		BPG Search:	
		[ ] The same title	
		[ ] Duplicate publication	
		[ ] Plagiarism	
		[Y] No	

COMMENTS TO AUTHORS

The authors present two brothers suffering from a rare genetic disorder with autosomal recessive inheritance, of nephrologic interest. The article is interesting, well written and certainly deserves to be published. Just a very few remarks: It would be appropriate to mention the possibility that point mutations that damage splice sites cause the phenomenon of "exon skipping" (see Tacheuchi et al., 2015). At page. 4 (Introduction): line 6, "experienced" should be "experience"; lines 26 and 27, review the phrase "this mutation is a disease causative of GS." At page. 8 (Discussion): line 7, "both patients, inherited" should be "both patients inherited"; line 11"The second mutation, inherited from the mother, carried by our patients, is a ..." would be better "The second mutation carried by our patients, inherited from the mother, is a ..."