

PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 59042

Title: A novel de novo mutation in the ASX gene in a Chinese boy with microcephaly and speech impairment: The Bainbridge-Ropers syndrome

Reviewer's code: 03478635

Position: Peer Reviewer

Academic degree: PhD

Professional title: Senior Research Fellow

Reviewer's Country/Territory: Japan

Author's Country/Territory: China

Manuscript submission date: 2020-08-28

Reviewer chosen by: AI Technique

Reviewer accepted review: 2020-08-28 03:28

Reviewer performed review: 2020-08-31 05:45

Review time: 3 Days and 2 Hours

Scientific quality	<input type="checkbox"/> Grade A: Excellent <input type="checkbox"/> Grade B: Very good <input checked="" type="checkbox"/> Grade C: Good <input type="checkbox"/> Grade D: Fair <input type="checkbox"/> Grade E: Do not publish
Language quality	<input checked="" type="checkbox"/> Grade A: Priority publishing <input 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 type="checkbox"/> Accept (General priority) <input checked="" 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

SPECIFIC COMMENTS TO AUTHORS

This is an important case report. Figure 2 may be revised to indicate more detailed information about each lines.

PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 59042

Title: A novel de novo mutation in the ASX gene in a Chinese boy with microcephaly and speech impairment: The Bainbridge-Ropers syndrome

Reviewer's code: 00646519

Position: Editorial Board

Academic degree: PhD

Professional title: Doctor

Reviewer's Country/Territory: Mexico

Author's Country/Territory: China

Manuscript submission date: 2020-08-28

Reviewer chosen by: AI Technique

Reviewer accepted review: 2020-08-30 13:52

Reviewer performed review: 2020-09-08 01:09

Review time: 8 Days and 11 Hours

Scientific quality	<input type="checkbox"/> Grade A: Excellent <input type="checkbox"/> Grade B: Very good <input checked="" 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 type="checkbox"/> Accept (General priority) <input checked="" type="checkbox"/> Minor revision <input type="checkbox"/> Major revision <input type="checkbox"/> Rejection
Re-review	<input checked="" type="checkbox"/> Yes <input 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

In the manuscript the authors present a case report of a novel mutation in the The Bainbridge-Ropers syndrome, these are my comments: Title should be modified “ A novel de novo”?, please correct it. Summary, please correct nomenclature (p.E599X). Specify if parents were young. Correct the text and the nomenclature “...the deletion of one base pair (G>T) and resulted in the replacement of the amino acid (histidine) (c.1795 G>T; p. E599X, 1650).....,” it was not a replacement of histidine, it was the substitution for a stop codon. Please describe properly the pedigree and electropherograms (codon and patient and parents). Figure 2 is unnecessary. Please include a couple of phrases about characteristics of the gene.

RE-REVIEW REPORT OF REVISED MANUSCRIPT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 59042

Title: Novel mutation in the ASX gene in a Chinese boy with microcephaly and speech impairment: A case report

Reviewer's code: 00646519

Position: Editorial Board

Academic degree: PhD

Professional title: Doctor

Reviewer's Country/Territory: Mexico

Author's Country/Territory: China

Manuscript submission date: 2020-08-28

Reviewer chosen by: Le Zhang

Reviewer accepted review: 2020-10-11 15:05

Reviewer performed review: 2020-10-11 15:11

Review time: 1 Hour

Scientific quality	<input type="checkbox"/> Grade A: Excellent <input 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 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 type="checkbox"/> Accept (General priority) <input type="checkbox"/> Minor revision <input type="checkbox"/> Major revision <input type="checkbox"/> Rejection



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Peer-reviewer statements	Peer-Review: <input type="checkbox"/> Y] Anonymous <input type="checkbox"/>] Onymous Conflicts-of-Interest: <input type="checkbox"/>] Yes <input type="checkbox"/> Y] No
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SPECIFIC COMMENTS TO AUTHORS

No comments