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PEER-REVIEW REPORT

Name of journal: *World Journal of Clinical Cases*

Manuscript NO: 71817

Title: A PIGN Mutation Case Report of Multiple Congenital Anomalies-Hypotonia-Seizures Syndrome 1

Provenance and peer review: Unsolicited manuscript; externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 02446734
Position: Peer Reviewer
Academic degree: DNB, FACC, FESC, MBBS, MD
Professional title: Academic Research, Doctor, Research Assistant Professor
Reviewer's Country/Territory: India
Author's Country/Territory: China

Manuscript submission date: 2021-09-23

Reviewer chosen by: Xin Liu (Online Science Editor)

Reviewer accepted review: 2021-12-21 05:15

Reviewer performed review: 2021-12-21 05:24

Review time: 1 Hour

Scientific quality	[ ] Grade A: Excellent [ ] Grade B: Very good [ ] Grade C: Good [Y] Grade D: Fair [ ] Grade E: Do not publish
Language quality	[ ] Grade A: Priority publishing [ ] Grade B: Minor language polishing [Y] Grade C: A great deal of language polishing [ ] Grade D: Rejection
Conclusion	[ ] Accept (High priority) [ ] Accept (General priority) [ ] Minor revision [Y] Major revision [ ] Rejection
Re-review	[Y] Yes [ ] No

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Peer-reviewer statements	Peer-Review: [ ] Anonymous [Y] Onymous Conflicts-of-Interest: [ ] Yes [Y] No

SPECIFIC COMMENTS TO AUTHORS

This a very common reported mutation in literature Its a well diagnosed case If can be rewritten as a review article, may help more Needs lot of English grammar corrections

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Provenance and peer review: Unsolicited manuscript; externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 00631867
Position: Peer Reviewer
Academic degree: MD

Professional title: Professor

Reviewer's Country/Territory: Australia

Author's Country/Territory: China

Manuscript submission date: 2021-09-23

Reviewer chosen by: Xin Liu (Online Science Editor)

Reviewer accepted review: 2021-12-14 08:12

Reviewer performed review: 2021-12-29 09:03

Review time: 15 Days

Scientific quality	[ ] Grade A: Excellent [Y] Grade B: Very good [ ] Grade C: Good [ ] Grade D: Fair [ ] Grade E: Do not publish
Language quality	[ ] Grade A: Priority publishing [Y] Grade B: Minor language polishing [ ] Grade C: A great deal of language polishing [ ] Grade D: Rejection
Conclusion	[ ] Accept (High priority) [ ] Accept (General priority) [Y] Minor revision [ ] Major revision [ ] Rejection
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Peer-reviewer statements	Peer-Review: [Y] Anonymous [ ] Onymous Conflicts-of-Interest: [ ] Yes [Y] No

SPECIFIC COMMENTS TO AUTHORS

The case represents a genetic basis for a form of epilepsy with developmental delay and cerebellar atrophy. The English could be improved as evidenced by the following quotation, " Here, we report…" . 'Here' is totally unnecessary and my preference is for third person rather than first person, the latter suggestive of personal bias. The same applies to 'here' in the following quotation "It should also be noted here that…" The authors persist with " anti-convulsive therapy" when common usage favours 'anti-seizure medication' The authors should avoid emotive language, such as "unfortunately", which suggests personal involvement rather than simple scientific reporting. The paper reports a positive result for genetic testing when the diagnosis was obscure and it reinforces the application of such testing even when the phenotypic presentation was somewhat unusual for the genetic findings but it does add a further dimension to the growing knowledge of genetic causes of epilepsy and for that reason alone merits publication.