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PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 68893

Title: A missense mutation in DYN H1 gene caused psychomotor developmental delay and muscle weakness: A case report

Reviewer's code: 05476795
Position: Peer Reviewer
Academic degree: MD
Professional title: Doctor
Reviewer's Country/Territory: Indonesia
Author's Country/Territory: China

Manuscript submission date: 2021-06-07
Reviewer chosen by: AI Technique
Reviewer accepted review: 2021-06-09 02:49
Reviewer performed review: 2021-06-09 06:36
Review time: 3 Hours

Scientific quality	[ ] Grade A: Excellent [Y] Grade B: Very good [ ] Grade C: Good [ ] Grade D: Fair [ ] Grade E: Do not publish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

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SPECIFIC COMMENTS TO AUTHORS

An interesting, eloquent report on the mutate DYNC1H1 gene in effecting psychomotor developmental delay and muscle weakness. Sentence fluency that is smooth and expressive, with no grammatical errors. A few things that the author(s) could consider adding to this manuscript: 1. What instrument was used to measure muscle strength and developmental delay on this patient? It would be great if the the author(s) comparison with normal measurements for children of the same age as a comparison to this patient. 2. What is the prognosis of this patient? 3. What is the follow-up plan? how will the author(s) plan to observe the progression of the disease?

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PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 68893

Title: A missense mutation in DYN H1 gene caused psychomotor developmental delay and muscle weakness: A case report

Reviewer's code: 05429607
Position: Peer Reviewer
Academic degree: MD

Professional title: Academic Fellow, Doctor

Reviewer's Country/Territory: Lithuania

Author's Country/Territory: China

Manuscript submission date: 2021-06-07

Reviewer chosen by: AI Technique

Reviewer accepted review: 2021-06-08 09:55

Reviewer performed review: 2021-06-17 10:36

Review time: 9 Days

Scientific quality	[ ] Grade A: Excellent [Y] Grade B: Very good [ ] Grade C: Good [ ] Grade D: Fair [ ] Grade E: Do not publish
Language quality	[Y] Grade A: Priority publishing [ ] 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
Re-review	[Y] Yes [ ] No
Peer-reviewer statements	Peer-Review: [Y] Anonymous [ ] Onymous Conflicts-of-Interest: [ ] Yes [Y] No

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SPECIFIC COMMENTS TO AUTHORS

Dear Authors, First of all, thank you for submitting your manuscript to the World Journal of Clinical Cases. This is an interesting, fluently written article about a rare genetical syndrome. However, it could be improved with some corrections and supplements. 1. Are there any prenatal signs of this rare syndrome? 2. In the discussion part, I suggest providing a table summarising previously reported cases. It would be 3. As you are planning to follow up this patient, it would add more scientific value to your article if you provide a follow-up plan and possible treatment and rehabilitation measures that could be used in such cases. In addition, an algorithm would supplement your case report greatly.