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PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 60505

Title: ACT mutation is responsible for multi-systemic smooth muscle dysfunction syndrome with seizures in a Chinese child: a case report and literature review

Reviewer's code: 01436649
Position: Editorial Board
Academic degree: MD, PhD
Professional title: Full Professor
Reviewer's Country/Territory: Croatia
Author's Country/Territory: China
Manuscript submission date: 2021-02-26
Reviewer chosen by: AI Technique
Reviewer accepted review: 2021-06-24 14:08
Reviewer performed review: 2021-06-24 14:46
Review time: 1 Hour

Scientific quality	[Y] Grade A: Excellent [ ] Grade B: Very good [ ] Grade C: Good [ ] Grade D: Fair [ ] Grade E: Do not publish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	[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

In this manuscript authors described a 7-year-old and eight-month-old girl who experienced seizures which are related to pathology of ACTA2 gene mutation. Paper was well prepared and written. However, there are few thing to mention: 1. Laboratory examinations have to be better described. 2. Please describe why patient got treatment which was described and cite treatment in literature for patients with this mutation