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PEER-REVIEW REPORT

Name of journal: *World Journal of Clinical Cases*

Manuscript NO: 86630

Title: Compound heterozygous mutations in TP Y cause rare autosomal recessive spinocerebellar ataxia type 7: a case report

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 05236189
Position: Editorial Board
Academic degree: MD
Professional title: Academic Research, Adjunct Associate Professor, Research Associate
Reviewer's Country/Territory: Brazil
Author's Country/Territory: China

Manuscript submission date: 2023-06-30

Reviewer chosen by: Geng-Long Liu

Reviewer accepted review: 2023-07-20 00:39

Reviewer performed review: 2023-07-20 01:01

Review time: 1 Hour

Scientific quality	[ ] Grade A: Excellent [Y] Grade B: Very good [ ] Grade C: Good [ ] Grade D: Fair [ ] Grade E: Do not publish
Novelty of this manuscript	[ ] Grade A: Excellent [Y] Grade B: Good [ ] Grade C: Fair [ ] Grade D: No novelty
Creativity or innovation of this manuscript	[ ] Grade A: Excellent [Y] Grade B: Good [ ] Grade C: Fair [ ] Grade D: No creativity or innovation

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Scientific significance of the conclusion in this manuscript	[ ] Grade A: Excellent [Y] Grade B: Good [ ] Grade C: Fair [ ] Grade D: No scientific significance
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) [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

SPECIFIC COMMENTS TO AUTHORS

1. Abstract. Please, remove "Eventually, the female was diagnosed with SCAR7" or rephrase it. 2. Case report. a. description of birth features should be provided—E.g., GA, head circumference, weight, baby's body length, and reflexes. b. Could the authors provide a history line of the clinical manifestations of the individual throughout the time? c. Why were used MMSE and MoCA instead of other scales specific for individuals with SCA, such as the CCAS scale? d. Could the authors describe the patient's vision?