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PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 65790

Title: SATB2-associated syndrome caused by a novel SATB2 mutation in a Chinese boy: A case report and literature review

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: 2021-03-16

Reviewer chosen by: AI Technique

Reviewer accepted review: 2021-03-16 09:54

Reviewer performed review: 2021-03-19 07:49

Review time: 2 Days and 21 Hours

Scientific quality	[ ] Grade A: Excellent [ ] Grade B: Very good [Y] Grade C: Good [ ] Grade D: Fair [ ] Grade E: Do not publish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
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 Author, First of all, thank you for submitting your manuscript to the World Journal of Clinical Cases. This is an interesting article about a rare genetical syndrome and providing some valuable insights about the importance of early interventions needed. However, it could be improved with some corrections and supplements. 1 What was the reason for genetic testing in this case? Was a patient referred by another specialist? Or did the parents reach out on their own? 2 It looks like the summary is too long (according to the manuscript guidelines, it should not exceed 250 words). You should also avoid abbreviations in summary as much as possible. 3 Line 19: Case description: treatment part – can you specify what exact special education and symptomatic treatment were performed? 4 Discussion - I suggest you provide a table with basic signs of the disease and possible treatment interventions. This might help to make a stronger impression about the importance of early diagnostics. 5 You can also make some reccomendations based on which the children with ID must be sent for genetic testing. 6 Figure 2: please indicate pictures A and B. 7 Some language polishing needed: several articles and commas are missing, some grammar mistakes. 8 Please correct the manuscript according to the provided guidelines for manuscript writing (format references indexes throughout the manuscript - superscript in square brackets– please check https://www.wjgnet.com/bpg/GerInfo/187.