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PEER-REVIEW REPORT

Name of journal: *World Journal of Clinical Cases*

Manuscript NO: 81790

Title: Acromicric dysplasia caused by a mutation of fibrillin 1 in a family: A case report

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 05238069
Position: Peer Reviewer
Academic degree: MD
Professional title: Doctor
Reviewer's Country/Territory: China
Author's Country/Territory: China

Manuscript submission date: 2022-11-29

Reviewer chosen by: AI Technique

Reviewer accepted review: 2022-12-02 00:11

Reviewer performed review: 2022-12-06 00:19

Review time: 4 Days

Scientific quality	[ ] Grade A: Excellent [ ] Grade B: Very good [Y] Grade C: Good [ ] Grade D: Fair [ ] Grade E: Do not publish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) [ ] Minor revision [Y] Major revision [ ] Rejection
Re-review	[ ] Yes [Y] No
Peer-reviewer	Peer-Review: [Y] Anonymous [ ] Onymous

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statements	Conflicts-of-Interest: [ ] Yes [Y] No

SPECIFIC COMMENTS TO AUTHORS

The authors described the clinical phenotype and genetic characteristics of a family with AD associated with a mutation in the FBN1 gene to enhance clinicians' understanding of the disease and to share the experience of a young patient receiving long-term rhGH therapy. However, some errors need to be corrected.

Comments to manuscript

Thank you for your kind invitation.

The authors described the clinical phenotype and genetic characteristics of a family with AD associated with a mutation in the FBN1 gene to enhance clinicians' understanding of the disease and to share the experience of a young patient receiving long-term rhGH therapy. However, some errors need to be corrected.

Query 1: There were some language errors in the manuscript that need to be revised as a native speaker.

Query 2: How to distinguish whether the boy's height increase is due to physiology or rhGH treatment? Maybe it can be analyzed according to the performances of his father and grandfather in the family.

Query 3: The pathogenicity criteria of the mutation had better be offered to the manuscript according to American College of Medical Genetics and Genomics (ACMGG).

Query 4: In the title "Acromicric dysplasia caused by a mutation of Fibrillin1 in a family: A case report", "Fibrillin1" should be written separately.

Query 5: In the part of background, did this sentence "which is easy to be confused with

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idiopathic short stature." describe appropriately?

Query 6: It's recommended that a general description of previously reported cases should be included in the manuscript to adequately describe the background, present status and significance of the study.

Query 7: There should be a proband mark in the pedigree map of the family.

Yours sincerely,

Miss Zhai.