



PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 69482

Title: A Rare Mutation in MKRN3 in Two Twin Sisters with Central Precocious Puberty:
A Case Report

Reviewer's code: 05117991

Position: Associate Editor

Academic degree: MD, MSc

Professional title: Associate Professor, Director, Surgeon

Reviewer's Country/Territory: Turkey

Author's Country/Territory: China

Manuscript submission date: 2021-07-01

Reviewer chosen by: AI Technique

Reviewer accepted review: 2021-07-02 05:57

Reviewer performed review: 2021-07-02 10:50

Review time: 4 Hours

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 Grade E: Do not publish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



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SPECIFIC COMMENTS TO AUTHORS

Dear Authors, I would like to thank you for submitting the case report entitled " A Rare Mutation in MKRN3 in Two Twin Sisters with Central Precocious Puberty: a Case Report". It describes twin sisters with central presenting with precocious puberty. This presentation was related to a specific genetic mutation that disrupts the hypothalamic-pituitary-gonadal axis. The mutated gene is located on chromosome 15q11.2. which codes the Makorin protein family. Case presentation, diagnostic work-up, genetic mutation, and treatment were described concisely and to the point. As a reviewer, I enjoyed reading the manuscript and find it acceptable for publishing. Many thanks again. Kind Regards, Sanem Guler Cimen



PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 69482

Title: A Rare Mutation in MKRN3 in Two Twin Sisters with Central Precocious Puberty:
A Case Report

Reviewer's code: 05916273

Position: Peer Reviewer

Academic degree: MBBS

Professional title: Doctor

Reviewer's Country/Territory: Nigeria

Author's Country/Territory: China

Manuscript submission date: 2021-07-01

Reviewer chosen by: AI Technique

Reviewer accepted review: 2021-07-01 16:54

Reviewer performed review: 2021-07-06 22:51

Review time: 5 Days and 5 Hours

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



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SPECIFIC COMMENTS TO AUTHORS

The finding is worth reporting. The case is well written except for the minor corrections highlighted in the review especially in the presentation of the history. overall, it is a good report.