PEER-REVIEW REPORT

Name of journal: World Journal of Clinical Cases

Manuscript NO: 68886

Title: Crouzon syndrome resulting from a novo mutation in a twin boy: a case report and review of the literature

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer’s code: 06124644

Position: Peer Reviewer

Academic degree: MD

Professional title: Doctor

Reviewer’s Country/Territory: Iran

Author’s Country/Territory: China

Manuscript submission date: 2021-12-13

Reviewer chosen by: AI Technique

Reviewer accepted review: 2021-12-14 19:36

Reviewer performed review: 2021-12-17 22:08

Review time: 3 Days and 2 Hours

Scientific quality

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[ ] Grade D: Fair  [ ] Grade E: Do not publish

Language quality

[ ] Grade A: Priority publishing  [ ] Grade B: Minor language polishing
[ Y] 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

[ ] Yes  [ Y] No
SPECIFIC COMMENTS TO AUTHORS

Dear author, It is well written rare case, quality is fair and as basis try to gain more cases
Name of journal: World Journal of Clinical Cases

Manuscript NO: 68886

Title: Crouzon syndrome resulting from a novo mutation in a twin boy: a case report and review of the literature

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer’s code: 03593092

Position: Editorial Board

Academic degree: MD, PhD

Professional title: Associate Professor, Chief Physician, Neurosurgeon

Reviewer’s Country/Territory: Taiwan

Author’s Country/Territory: China

Manuscript submission date: 2021-12-13

Reviewer chosen by: AI Technique

Reviewer accepted review: 2021-12-14 17:32

Reviewer performed review: 2021-12-26 17:24

Review time: 11 Days and 23 Hours

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<th>Scientific quality</th>
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SPECIFIC COMMENTS TO AUTHORS
This report presented a 6-year-old boy diagnosed with Crouzon syndrome caused by a de novo mutation in the fibroblast growth factor receptor 2 (FGFR2) gene - c.1026C>G (p.Cys342Trp). Since Crouzon syndrome is rare genetic disorder, this case is worthy of reporting. However, there are a few minor issues in the current manuscript. 1. To clarify boy/girl twins, “fraternal twins” may be used. 2. In Core tip, you mentioned “There are no case reports showing the Crouzon syndrome occurs in the twins.” But, this statement is NOT correct according to the following reference, which should be rephrased. Lloyd MS, Trost JG, Khechoyan DY, Hollier LH Jr, Buchanan EP. Identical Twins with Crouzon Syndrome: Eight-Year Follow-up, Genetic Considerations, and Operative Management. Craniomaxillofac Trauma Reconstr. 2017 Dec;10(4):286-291. doi: 10.1055/s-0036-1592091. Epub 2016 Sep 2. PMID: 29109840; PMCID: PMC5669987. 3. Please correct grammatical and spelling errors. For example: In the Title, “....resulting from a novo mutation...” “The most common feartures include...” “Since the twin sister and the parents did not present gene mutations and a similar presentation,” 4. Please remove “case report” from Key Words. 5. Please rephrase the Conclusion parts to emphasize the patient presented, rather than providing general statements regarding Crouzon syndrome.