



## PEER-REVIEW REPORT

**Name of journal:** World Journal of Clinical Cases

**Manuscript NO:** 65822

**Title:** Diagnosis and treatment discussion of congenital factor VII deficiency in pregnancy: A case report

**Reviewer's code:** 03818170

**Position:** Peer Reviewer

**Academic degree:** MD, PhD

**Professional title:** Doctor

**Reviewer's Country/Territory:** Turkey

**Author's Country/Territory:** China

**Manuscript submission date:** 2021-03-22

**Reviewer chosen by:** AI Technique

**Reviewer accepted review:** 2021-03-25 17:47

**Reviewer performed review:** 2021-03-28 09:43

**Review time:** 2 Days and 15 Hours

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 Grade E: Do not publish            |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



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#### **SPECIFIC COMMENTS TO AUTHORS**

The present study named "Diagnosis and treatment discussion on congenital Factor VII deficiency in pregnancy and a case report" was read and evaluated carefully. This is case report but it was written like a review. So there is a mixed situation in presentation. The present study is well written and rare condition and unusual case in literature. Because there is less than 20 studies in literature (PMID: 16834743, 6983546, 25525535, 30831446, 22197042, 11336760, 12759632, 11012708). The author should add the result of genetic disorder which include homozygote F7D (with mutation in exon). That will prevent the bias or false information. The study should include other case report in literature, if possible. Best regards..