Name of journal: World Journal of Clinical Cases

Manuscript NO: 87543

Title: A case report and literature review of congenital leukemia

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer’s code: 01551432

Position: Editorial Board

Academic degree: MD, PhD

Professional title: Doctor

Reviewer’s Country/Territory: Japan

Author’s Country/Territory: China

Manuscript submission date: 2023-08-20

Reviewer chosen by: AI Technique

Reviewer accepted review: 2023-08-27 04:45

Reviewer performed review: 2023-08-27 08:15

Review time: 3 Hours

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<th>Scientific quality</th>
<th>[ ] Grade A: Excellent</th>
<th>[Y] Grade B: Very good</th>
<th>[ ] Grade C: Good</th>
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<td>[ ] Grade D: Fair</td>
<td>[ ] Grade E: Do not publish</td>
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<th>Novelty of this manuscript</th>
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<td>[ ] Grade D: No novelty</td>
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<th>Creativity or innovation of this manuscript</th>
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<td>[ ] Grade D: No creativity or innovation</td>
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SPECIFIC COMMENTS TO AUTHORS

Dear Authors,

Thank you for your submitting the manuscript entitled, "A case report and literature review of congenital leukemia". The manuscript is well written and compactly summarized. This is an interesting, thought-provoking, and unusual case of DS-related AML diagnosed and identified by gene sequencing as a mutation in the GATA1 gene, well-documented and highly unusual. 

1. Please list several similar cases and add a new table summarizing age, sex, symptoms, characteristic physical and laboratory findings, treatment, course, and prognosis. Further, please add new considerations from that table. 

2. Please add such as macroscopic and gross images of the baby and invaded organs, and imaging findings from echocardiography and computed tomography (CT), if possible. 

3. Were there any abnormal prenatal echocardiographic findings, was prenatal diagnosis possible by amniotic fluid testing, cell chromosome analysis, etc? Is prenatal diagnosis possible in the first place?

Best regards,