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PEER-REVIEW REPORT

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

Scientific quality	[ ] Grade A: Excellent [Y] Grade B: Very good [ ] Grade C: Good [ ] Grade D: Fair [ ] Grade E: Do not publish
Novelty of this manuscript	[ ] Grade A: Excellent [Y] Grade B: Good [ ] Grade C: Fair [ ] Grade D: No novelty
Creativity or innovation of this manuscript	[ ] Grade A: Excellent [Y] Grade B: Good [ ] Grade C: Fair [ ] Grade D: No creativity or innovation

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Scientific significance of the conclusion in this manuscript	[ ] Grade A: Excellent [Y] Grade B: Good [ ] Grade C: Fair [ ] Grade D: No scientific significance
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) [Y] Minor revision [ ] Major revision [ ] Rejection
Re-review	[Y] Yes [ ] No
Peer-reviewer statements	Peer-Review: [Y] Anonymous [ ] Onymous
	Conflicts-of-Interest: [ ] Yes [Y] No

SPECIFIC COMMENTS TO AUTHORS

Dear Authers, Thank you for your submitting the manuscript entitled, "A case report and literature review of congenital leukemia". The manuscript is well written and compactky 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. minor) 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,