

Review Form 3

Journal Name:	Asian Journal of Case Reports in Medicine and Health
Manuscript Number:	Ms_AJCRMH_129502
Title of the Manuscript:	Genetic and Clinical Complexity in Jacobsen Syndrome: A Case of 11q23.3-q25 Deletion with Trigenocephaly and MYBPC3 Mutation
Type of the Article	Case report

General guidelines for the Peer Review process:

This journal's peer review policy states that **NO** manuscript should be rejected only on the basis of '**lack of Novelty**', provided the manuscript is scientifically robust and technically sound. To know the complete guidelines for the Peer Review process, reviewers are requested to visit this link:

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PART 1: Comments

	Reviewer's comment	Author's Feedback <i>(Please correct the manuscript and highlight that part in the manuscript. It is mandatory that authors should write his/her feedback here)</i>
Please write a few sentences regarding the importance of this manuscript for the scientific community. A minimum of 3-4 sentences may be required for this part.	This manuscript is essential for the scientific community as it explores the genetic and clinical intricacies of Jacobsen syndrome, a rare chromosomal disorder. By documenting a case with unique findings, including a pathogenic MYBPC3 mutation alongside the typical 11q23.3-q25 deletion, it broadens the understanding of phenotypic diversity within this condition.	
Is the title of the article suitable? (If not please suggest an alternative title)	a simpler or more concise title is preferred, it could be rephrased as: "Genetic and Clinical Insights into Jacobsen Syndrome: A Case Report."	
Is the abstract of the article comprehensive? Do you suggest the addition (or deletion) of some points in this section? Please write your suggestions here.	The abstract mentions the MYBPC3 mutation but does not fully explain its contribution to the case's uniqueness. A brief note on why this mutation is significant in the context of Jacobsen syndrome could add value. While the abstract mentions the importance of comprehensive management, it could briefly touch on the specific disciplines involved (e.g., neurology, cardiology, genetics) to emphasize the complexity of care. Include a statement on how the findings contribute to improving diagnostic accuracy or management strategies for Jacobsen syndrome. This would strengthen the link between the case and its importance to the scientific community. Certain details, such as "T2 prolongation of unmyelinated white matter," could be explained in simpler terms or omitted if they are not central to the abstract's purpose.	
Is the manuscript scientifically, correct? Please write here.	The manuscript is scientifically correct and well-structured.	
Are the references sufficient and recent? If you have suggestions of additional references, please mention them in the review form.	"Recent Advances in the Clinical and Molecular Aspects of Chromosomal Deletion Syndromes" (European Journal of Human Genetics, 2021). "The Role of Secondary Genetic Findings in Rare Diseases: Implications for Diagnosis and Management" (Genetics in Medicine, 2020). The current references are sufficient for the manuscript's core content, but adding a few recent, high-impact studies as outlined above would strengthen the manuscript and demonstrate engagement with the latest advancements in the field.	
Is the language/English quality of the article suitable for scholarly communications?	It is suitable.	
Optional/General comments		

PART 2:

	Reviewer's comment	Author's comment <i>(if agreed with reviewer, correct the manuscript and highlight that part in the manuscript. It is mandatory that authors should write his/her feedback here)</i>
Are there ethical issues in this manuscript?	<i>(If yes, Kindly please write down the ethical issues here in details)</i>	

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