Baishideng Publishing Group

7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA
Telephone: +1-925-399-1568
E-mail: office@baishideng.com
https://www.wjgnet.com

PEER-REVIEW REPORT

Name of journal: *World Journal of Clinical Cases*

Manuscript NO: 90798

Title: Detection of 4p16.3 deletion and 11p15.5p15.4 gain in a boy by comparative genomic hybridization array

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 05714329
Position: Peer Reviewer
Academic degree: MD

Professional title: Doctor

Reviewer's Country/Territory: Spain

Author's Country/Territory: Turkey

Manuscript submission date: 2023-12-13

Reviewer chosen by: AI Technique

Reviewer accepted review: 2023-12-14 09:23

Reviewer performed review: 2023-12-18 08:31

Review time: 3 Days and 23 Hours

Scientific quality	[ ] Grade A: Excellent [ ] Grade B: Very good [Y] Grade C: Good [ ] Grade D: Fair [ ] Grade E: Do not publish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

Baishideng Publishing Group

7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA
Telephone: +1-925-399-1568
E-mail: office@baishideng.com
https://www.wjgnet.com

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	[Y] Grade A: Priority publishing [ ] 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

The genetic characteristics of the clinical case presented are of interest and a correct methodology has been followed for its diagnosis. Although the genetic information is correct, the information regarding the facial characteristics present in a more explicit way. On the other hand, since we dot know the age of the case, we do not know if it would be pertinent to provide data referring to possible dental findings to complete the oral characteristics. In relation to the references provided on the WSH, I would suggest taking into account the incorporation of some pertinent to the clinical case, as follows: Related to mental retardation: Zollino M, Doronzio PN. Dissecting the Wolf-Hirschhorn syndrome phenotype: WHSC1 is a neurodevelopmental gene contributing to growth delay, intellectual disability, and to the facial dysmorphism. J Human Genet 2018;63:859-61. Related to oral findings: LimeresJ et al. Oral manifestations of Wolf-Hirschhorn syndrome: genotype-phenotype correlation analysis. J Clin Med 2020;4;9(11):3556