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## PEER-REVIEW REPORT

Name of journal: World Journal of Otorhinolaryngology

Manuscript NO: 88281

**Title:** Usher Syndrome: genetic diagnosis and current therapeutic approaches

Provenance and peer review: Unsolicited Manuscript; Externally peer reviewed

Peer-review model: Single blind

Reviewer's code: 01221812 Position: Peer Reviewer

Academic degree: MPhil, PhD

Professional title: Academic Research, Professor, Research Scientist, Teacher

Reviewer's Country/Territory: Pakistan

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

Manuscript submission date: 2023-09-18

Reviewer chosen by: Yu-Lu Chen

Reviewer accepted review: 2023-12-05 02:57

Reviewer performed review: 2023-12-09 09:32

**Review time:** 4 Days and 6 Hours

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	[ ] Grade D: Fair [ ] Grade E: Do not publish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

Genetic and molecular diagnosis of Usher Syndrome: can early diagnosis improve quality of life? Name of Journal: World Journal of Otorhinolaryngology This is an interesting review study, however, the manuscript would benefit from the following amendments: 1. Please give the OMIM number for each malformation and gene mentioned in the text. 2. Introduction: The text overlapping Table 1 should be reduced. 3. Introduction: The information on genes should be summarized into separate Table. 4. It would be worthwhile to present the list of representative mutations (and respective ethnic/geographic populations) found in genes linked with Usher syndrome. 5. Please give the differential diagnosis of Usher syndrome. 6. The text does not justify the title, "Genetic and molecular diagnosis of Usher Syndrome: can early diagnosis improve quality of life?" Please emphasize the early diagnosis in the manuscript. 7. It would be worthwhile to present a schematics showing the pathway of diagnosis pathway. 8. The text under Gene Therapy, Drug Therapy, Cochlear Implantation should be divided into sections.