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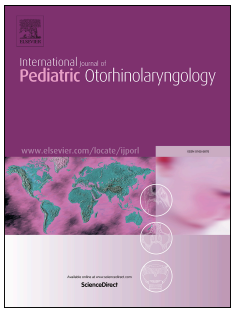

Utility of whole exome sequencing in the diagnosis of Usher syndrome: Report of novel compound heterozygous *MYO7A* mutations

Khushnooda Ramzan, Mohammed Al-Owain, Rozeena Huma, Selwa A.F. Al-Hazzaa, Sarah Al-Ageel, Faiqa Imtiaz, Moeenaldeen Al-Sayed

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Utility of whole exome sequencing in the diagnosis of Usher syndrome: report of novel compound heterozygous *MYO7A* mutations

Khushnooda Ramzan [1*], Mohammed Al-Owain [2,3], Rozeena Huma [2], Selwa A.F. Al-Hazzaa [3,4], Sarah Al-Ageel [5], Faiqa Imtiaz [1], Moeenaldeen Al-Sayed [2,3]

[1] Department of Genetics, King Faisal Specialist Hospital and Research Centre, Riyadh, Saudi Arabia

[2] Department of Medical Genetics, King Faisal Specialist Hospital and Research Centre, Riyadh, Saudi Arabia

[3] College of Medicine, Alfaisal University, Riyadh, Saudi Arabia

[4] Department of Ophthalmology, King Faisal Specialist Hospital and Research Centre, Riyadh, Saudi Arabia

[5] Department of Otolaryngology Head And Neck Surgery, King Faisal Specialist Hospital and Research Centre, Riyadh, Saudi Arabia

* Corresponding author: Khushnooda Ramzan, PhD, Department of Genetics, Research Centre, King Faisal Specialist Hospital & Research Centre, P.O.Box 3354, Riyadh 11211, Saudi Arabia.
Tel.: +966 11 4647272 Ext: 36484; fax: + 966 11 4424585
e-mail: kramzan@kfshrc.edu.sa; khushnooda@gmail.com

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