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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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