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Genome-Wide Association Study to Find Modifiers for Tetralogy of Fallot in the 22q11.2 Deletion Syndrome Identifies Variants in the GPR88 Locus on 5q14.3

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α-Galactosidase A Genotype N215S Induces a Specific Cardiac Variant of Fabry Disease

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A Comparison of Whole Genome Sequencing to Multigene Panel Testing in Hypertrophic Cardiomyopathy Patients

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Genomic Triangulation and Coverage Analysis in Whole-Exome Sequencing-Based Molecular Autopsies

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Care in Specialized Centers and Data Sharing Increase Agreement in Hypertrophic Cardiomyopathy Genetic Test Interpretation

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Clinical Genomic Cases

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