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Multiple Gene Variants in Hypertrophic Cardiomyopathy in the Era of Next-Generation Sequencing

Novel Genetic Triggers and Genotype–Phenotype Correlations in Patients With Left Ventricular Noncompaction

Editorial: Navigating Genetic and Phenotypic Uncertainty in Left Ventricular Noncompaction

Clinical Characteristics and Long-Term Outcome of Hypertrophic Cardiomyopathy in Individuals With a MYBPC3 (Myosin-Binding Protein C) Founder Mutation

Editorial: Founder Mutations in Myosin-Binding Protein C: Maybe Not So Benign After All

Fabry Disease in Families With Hypertrophic Cardiomyopathy: Clinical Manifestations in the Classic and Later-Onset Phenotypes

Editorial: Fabry Disease: A Rare Condition Emerging From the Darkness

CKM Glu83Gly Is Associated With Blunted Creatine Kinase Variation, but Not With Myalgia

Editorial: CKing Precision in the Interpretation of Diagnostic Biomarkers

Carotid Plaque Rupture Is Accompanied by an Increase in the Ratio of Serum circR-284 to miR-221 Levels


Investigating the Genetic Causes of Sudden Unexpected Death in Children Through Targeted Next-Generation Sequencing Analysis

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Editorial: Informational Quest

Advances in Genetics

Cardiovascular Disease and Long Noncoding RNAs: Tools for Unraveling the Mystery Lnc-ing RNA and Phenotype

Clinical Genomic Cases

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