Original Articles

**Lamin A/C-Related Cardiac Disease: Late Onset With a Variable and Mild Phenotype in a Large Cohort of Patients With the Lamin A/C p.(Arg331Gln) Founder Mutation**

**Editorial: LMNA Mutations Associated With Mild and Late-Onset Phenotype: The Case of the Dutch Founder Mutation p.(Arg331Gln)**

**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**

**Editorial: Towards Point-of-Care Measurements Using Noncoding RNAs: A Novel Tool to Monitor Aggravation of Advanced Atherosclerotic Lesions**

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

**Loss of Y Chromosome in Blood Is Associated With Major Cardiovascular Events During Follow-Up in Men After Carotid Endarterectomy**

**Editorial: Loss of Chromosome Y in Leukocytes and Major Cardiovascular Events**

**Fifteen Genetic Loci Associated With the Electrocardiographic P Wave**

**Editorial: Dissecting the Genetic Basis of the ECG as a Means of Understanding Mechanisms of Arrhythmia**

**PCSK9 Loss-of-Function Variants, Low-Density Lipoprotein Cholesterol, and Risk of Coronary Heart Disease and Stroke: Data From 9 Studies of Blacks and Whites**

**Effect of Disclosing Genetic Risk for Coronary Heart Disease on Information Seeking and Sharing: The MI-GENES Study (Myocardial Infarction Genes)**

**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**

Previously Unreported in Women *Galactosidase Alpha* Pro409Ser Variant Is Associated With Fabry Disease