Volume 10, Number 4, August 2017
ISSN 1942-3268
http://circgenetics.ahajournals.org

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


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