Editorials

Are Double Mutations Double Trouble?
Local Ancestry Association, Admixture Mapping, and Ongoing Challenges

Original Articles

Identification of Cadherin 2 (CDH2) Mutations in Arrhythmogenic Right Ventricular Cardiomyopathy

Screening of the Filamin C Gene in a Large Cohort of Hypertrophic Cardiomyopathy Patients

Nonfamilial Hypertrophic Cardiomyopathy: Prevalence, Natural History, and Clinical Implications

Prevalence and Clinical Implication of Double Mutations in Hypertrophic Cardiomyopathy: Revisiting the Gene-Dose Effect

Additional Candidate Genes for Human Atherosclerotic Disease Identified Through Annotation Based on Chromatin Organization

Admixture Mapping of Subclinical Atherosclerosis and Subsequent Clinical Events Among African Americans in 2 Large Cohort Studies

Socioeconomic Status Interacts with the Genetic Effect of a Chromosome 9p21.3 Common Variant to Influence Coronary Artery Calcification and Incident Coronary Events in the Heinz Nixdorf Recall Study (Risk Factors, Evaluation of Coronary Calcium, and Lifestyle)

Genome-Wide Association Study Meta-Analysis of Long-Term Average Blood Pressure in East Asians

Investigating the Genetic Architecture of the PR Interval Using Clinical Phenotypes

Heritability of the Severity of the Metabolic Syndrome in Whites and Blacks in 3 Large Cohorts

Genetic Analysis of Venous Thromboembolism in UK Biobank Identifies the ZFPM2 Locus and Implicates Obesity as a Causal Risk Factor

Advances in Genetics

Delivering Clinical Grade Sequencing and Genetic Test Interpretation for Cardiovascular Medicine

AHA Scientific Statement

Potential Impact and Study Considerations of Metabolomics in Cardiovascular Health and Disease: A Scientific Statement From the American Heart Association
Circ Cardiovasc Genet. 2017;10:

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