

# Circulation: Genomic and Precision Medicine

## ORIGINAL ARTICLES

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***SCN5A* (Na<sub>v</sub>1.5) Variant Functional Perturbation and Clinical Presentation: Variants of a Certain Significance**

Editorial

Predicting Penetrance of *SCN5A* Rare Variants: Peering Beyond the Black and White and Into the Shades of Grey

Common Coding Variants in *SCN10A* Are Associated With the Nav1.8 Late Current and Cardiac Conduction

Editorial

*SCN10A*-Dependent Late I<sub>Na</sub> Current: Never Too Late for Cardiac Conduction?

Common and Rare Coding Genetic Variation Underlying the Electrocardiographic PR Interval

Editorial

Genetic Variation and the Electrocardiograph: From Genome-Wide Association Studies to the Patient

Genetic Regulation of PCSK9 (Proprotein Convertase Subtilisin/Kexin Type 9) Plasma Levels and Its Impact on Atherosclerotic Vascular Disease Phenotypes

Editorial

Genetic Regulation of PCSK9 (Proprotein Convertase Subtilisin/Kexin Type 9) Plasma Levels: Another Piece of the Puzzle

CETP (Cholesteryl Ester Transfer Protein) Concentration: A Genome-Wide Association Study Followed by Mendelian Randomization on Coronary Artery Disease

Editorial

Application of Genetic Epidemiology to CETP (Cholesteryl Ester Transfer Protein) Concentration and Risk of Cardiovascular Disease

## PERSPECTIVE

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Frameshifts in Code and in Care: The Importance of Timely Genetic Evaluation

## RESEARCH LETTER

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Increased Prevalence of Congenital Heart Disease in Children With Diamond Blackfan Anemia Suggests Unrecognized Diamond Blackfan Anemia as a Cause of Congenital Heart Disease in the General Population: A Report of the Diamond Blackfan Anemia Registry

## CLINICAL VIGNETTES

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*TRPM4* Mutation in Patients With Ventricular Noncompaction and Cardiac Conduction Disease

## CORRESPONDENCE

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Letter by Ruiz-Guerrero et al Regarding Article, "Yield of the RYR2 Genetic Test in Suspected Catecholaminergic Polymorphic Ventricular Tachycardia and Implications for Test Interpretation"

Response by Kapplinger et al to Letter Regarding Article, "Yield of the RYR2 Genetic Test in Suspected Catecholaminergic Polymorphic Ventricular Tachycardia and Implications for Test Interpretation"

Letter by Vermeer et al Regarding Article, "Phenotypic Spectrum of HCN4 Mutations: A Clinical Case"

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