

Original Articles

Diminished *PRRX1* Expression Is Associated With Increased Risk of Atrial Fibrillation and Shortening of the Cardiac Action Potential

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Editorial: Connecting the Dots in Atrial Fibrillation

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Genome-Wide Association Study to Find Modifiers for Tetralogy of Fallot in the 22q11.2 Deletion Syndrome Identifies Variants in the *GPR98* Locus on 5q14.3

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Editorial: Modifying Mendel Redux: Unbiased Approaches Can Find Modifiers

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New Blood Pressure-Associated Loci Identified in Meta-Analyses of 475000 Individuals

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Editorial: Blood Pressure Genome-Wide Association Studies, Missing Heritability, and Omnigenics

Brian J. Morris, DSc, PhD e001943

Transcriptome and Functional Profile of Cardiac Myocytes Is Influenced by Biological Sex

Christa L. Trexler, PhD; Aaron T. Odell, BS; Mark Y. Jeong, MD; Robin D. Dowell, DSc; Leslie A. Leinwand, PhD e001770

Editorial: Sex Determines Cardiac Myocyte Stretch and Relaxation

Michael J. Coronado, PhD; DeLisa Fairweather, PhD; Katelyn A. Bruno, PhD e001950

MicroRNA Signature of Cigarette Smoking and Evidence for a Putative Causal Role of MicroRNAs in Smoking-Related Inflammation and Target Organ Damage

Christine M. Willinger, BS; Jian Rong, PhD; Kahraman Tanriverdi, PhD; Paul L. Courchesne, MBA; Tianxiao Huan, PhD; Gregory A. Wasserman, MS; Honghuang Lin, PhD; Josée Dupuis, PhD; Roby Joehanes, PhD; Matthew R. Jones, PhD; George Chen, BS; Emelia J. Benjamin, MD, ScM; George T. O'Connor, MD; Joseph P. Mizgerd, ScD; Jane E. Freedman, MD; Martin G. Larson, ScD; Daniel Levy, MD e001678

Editorial: A Smoking-Associated miRNA-mRNA Coexpression Network

Mete Civelek, PhD e001914

Reduced Cardiac Calcineurin Expression Mimics Long-Term Hypoxia-Induced Heart Defects in *Drosophila*

Rachel Zarndt, PhD; Stanley M. Walls, PhD; Karen Ocorr, PhD; Rolf Bodmer, PhD e001706

Editorial: High Heart: A Role for Calcineurin Signaling in Hypoxia-Influenced Cardiac Growth

TyAnna L. Lovato, PhD; Richard M. Cripps, DPhil e001919

Body Height and Incident Risk of Venous Thromboembolism: A Cosibling Design

Bengt Zöller, MD, PhD; Jianguang Ji, MD, PhD; Jan Sundquist, MD, PhD; Kristina Sundquist, MD, PhD e001651

Editorial: Insights From the Positive Association of Height With Incident Venous Thromboembolism

C. Mary Schooling, PhD e001911

Heritability of Mitral Regurgitation: Observations From the Framingham Heart Study and Swedish Population

Francesca N. Delling, MD, MPH; Xinjun Li, MD, MPH, PhD; Shuo Li, MS; Qiong Yang, PhD; Vanessa Xanthakis, PhD; Andreas Martinsson, MD; Pontus Andell, MD, PhD; Birgitta T. Lehman, RDCS; Ewa W. Osypiuk, MD, RDCS; Plamen Stantchev, MD; Bengt Zöller, MD, PhD; Emelia J. Benjamin, MD, ScM; Kristina Sundquist, MD, PhD; Ramachandran S. Vasan, MD; J. Gustav Smith, MD, PhD e001736

Editorial: Inheritance Impacts Mitral Valve Insufficiency

Daniel P. Judge, MD; Russell A. Norris, PhD e001920

Validation of Polygenic Scores for QT Interval in Clinical Populations

Michael A. Rosenberg, MD; Steven A. Lubitz, MD, MPH; Honghuang Lin, PhD; Gulum Kosova, PhD; Victor M. Castro, MS; Paul Huang, MD, PhD; Patrick T. Ellinor, MD, PhD; Roy H. Perlis, MD, MSc; Christopher Newton-Cheh, MD, MPH e001724

Editorial: QT Interval Determinant: Mutations, Rare Variants, or Single-Nucleotide Polymorphisms?

Takeshi Aiba, MD, PhD; Atsushi Takahashi, PhD e001945

Prediction for Intravenous Immunoglobulin Resistance by Using Weighted Genetic Risk Score Identified From Genome-Wide Association Study in Kawasaki Disease

Ho-Chang Kuo, MD, PhD; Henry Sung-Ching Wong, MSc; Wei-Pin Chang, PhD; Ben-Kuen Chen, PhD; Mei-Shin Wu, MSc; Kuender D. Yang, MD, PhD; Kai-Sheng Hsieh, MD; Yu-Wen Hsu, MSc; Shih-Feng Liu, MD; Xiao Liu, MSc; Wei-Chiao Chang, PhD **OPEN** e001625

Editorial: One Size Does Not Fit All: Genetic Prediction of Kawasaki Disease Treatment Response in Diverse Populations

Michael A. Portman, MD; Sadeep Shrestha, PhD, MHS, MS e001917

Impact of Selection Bias on Estimation of Subsequent Event Risk

Yi-Juan Hu, PhD; Amand F. Schmidt, PhD; Frank Dudbridge, PhD; Michael V. Holmes, PhD; James M. Brophy, MD, PhD; Vinicius Tragante, PhD; Ziyi Li, MS; Peizhou Liao, PhD; Arshed A. Quyyumi, MD; Raymond O. McCubrey, MS; Benjamin D. Horne, PhD; Aroon D. Hingorani, PhD; Folkert W. Asselbergs, MD, PhD; Riyaz S. Patel, MD; Qi Long, PhD; on behalf of the GENIUS-CHD Consortium e001616

Editorial: Biases in Genetic Association of Coronary Heart Disease Events May Be Less Likely Than Suspected: Here Is When to Check for Them

Jennifer R. Dungan, PhD, RN e001912

Predicting the Functional Impact of KCNQ1 Variants of Unknown Significance

Bian Li, MSc; Jeffrey L. Mendenhall, MSc; Brett M. Kroncke, PhD; Keenan C. Taylor, PhD; Hui Huang, PhD; Derek K. Smith, DDS, PhD; Carlos G. Vanoye, PhD; Jeffrey D. Blume, PhD; Alfred L. George, Jr, MD; Charles R. Sanders, PhD; Jens Meiler, PhD e001754

Editorial: Machine Learning and Rare Variant Adjudication in Type 1 Long QT Syndrome

John R. Giudicessi, MD, PhD e001944

Characterization of a Human Induced Pluripotent Stem Cell-Derived Cardiomyocyte Model for the Study of Variant Pathogenicity: Validation of a KCNJ2 Mutation

Roselle Gélinas, PhD; Nabil El Khoury, PhD; Marie-A. Chaix, MD, MSc; Claudine Beauchamp, MSc; Azadeh Alikashani, MSc; Nathalie Ethier, MSc; Gabrielle Boucher, MSc; Louis Villeneuve, MSc; Laura Robb, MSc; Frédéric Latour, MSc; Blandine Mondesert, MD; Lena Rivard, MD; Philippe Goyette, PhD; Mario Talajic, MD; Céline Fiset, PhD; John David Rioux, PhD e001755

Editorial: Commercially Available Human-Induced Pluripotent Stem Cell-Derived Cardiomyocytes: Another Piece in Our Tool Box, but Not a Swiss Army Knife Yet

Leif-Hendrik Boldt, MD; Abdul S. Parwani, MD; Frank R. Heinzel, MD, PhD e001913

α -Galactosidase A Genotype N215S Induces a Specific Cardiac Variant of Fabry Disease

Daniel Oder, MD; Dan Liu, MD; Kai Hu, MD; Nurcan Üçeyler, MD; Tim Salinger, MD; Jonas Müntze, MD; Kristina Lorenz, PhD; Reinhard Kandolf, MD; Hermann-Josef Gröne, MD; Claudia Sommer, MD; Georg Ertl, MD; Christoph Wanner, MD; Peter Nordbeck, MD e001691

Editorial: Clinical Characteristics of the GLA N215S Variant and Implications for the Diagnosis and Management of Nonclassic Fabry Disease

Chloe Reuter, MS; Julia Platt, MS e001918

A Comparison of Whole Genome Sequencing to Multigene Panel Testing in Hypertrophic Cardiomyopathy Patients

Allison L. Cirino, MS; Neal K. Lakdawala, MD; Barbara McDonough, RN; Lauren Conner, MS; Dale Adler, MD; Mark Weinfeld, MD; Patrick O'Gara, MD; Heidi L. Rehm, PhD; Kalotina Machini, PhD; Matthew Lebo, PhD; Carrie Blout, MS; Robert C. Green, MD, MPH; Calum A. MacRae, MD, PhD; Christine E. Seidman, MD; Carolyn Y. Ho, MD; for the MedSeq Project e001768

Editorial: Hypertrophic Cardiomyopathy Gene Testing: Go Big?

Megan J. Puckelwartz, PhD; Elizabeth M. McNally, MD, PhD e001951

Genomic Triangulation and Coverage Analysis in Whole-Exome Sequencing–Based Molecular Autopsies

Garrett W. Shanks, BA; David J. Tester, BS; Sneha Nishtala, MS; Jared M. Evans, MS; Michael J. Ackerman, MD, PhD e001828

Editorial: Genomic Triangulation in Sudden Unexplained Death in the Young: The Way to Go?

Emil Daniel Bartels, MD, PhD; Jacob Tfelt-Hansen, MD, PhD, DMSc; Bo Gregers Winkel, MD, PhD ... e001915

Care in Specialized Centers and Data Sharing Increase Agreement in Hypertrophic Cardiomyopathy Genetic Test Interpretation

Aisha Furqan, MS; Patricia Arscott, MS; Francesca Girolami, BS; Allison L. Cirino, MS; Michelle Michels, MD, PhD; Sharlene M. Day, MD; Iacopo Olivotto, MD; Carolyn Y. Ho, MD; Euan Ashley, MRCP, DPhil; Eric M. Green, MD, PhD; Colleen Caleshu, MS; on behalf of the SHARe Consortium e001700

Editorial: Pathogenicity of Hypertrophic Cardiomyopathy Variants: A Path Forward Together

Jodie Ingles, MPH, PhD; Charlotte Burns, MGC, MPH; Birgit Funke, PhD e001916

Perspective

What Do We Really Think About Human Germline Genome Editing, and What Does It Mean for Medicine?

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Clinical Genomic Cases

Heterozygous Null LDLR Mutation in a Familial Hypercholesterolemia Patient With an Atypical Presentation Because of Alcohol Abuse

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Letter by Brewster Regarding Article, “CKM Glu83Gly Is Associated With Blunted Creatine Kinase Variation, but Not With Myalgia”

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10 (5)

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