

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

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Eloisa Arbustini, MD; Valentina Favalli, PhD; Nupoor Narula, MD

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June-Wha Rhee, MD; Megan E. Grove, MS, LCGC; Euan A. Ashley, BSc, DPhil, FRCP

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Perry Elliott, MBBS, MD, FRCP, FESC, FACC

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AHA Scientific Statement

The Expressed Genome in Cardiovascular Diseases and Stroke: Refinement, Diagnosis, and Prediction: A Scientific Statement From the American Heart Association <i>Kiran Musunuru, MD, PhD, MPH, FAHA, Chair; Erik Ingelsson, MD, PhD, FAHA, Co-Chair; Myriam Fornage, PhD, FAHA; Peter Liu, MD, FAHA; Anne M. Murphy, MD, FAHA; L. Kristin Newby, MD, MHS, FAHA; Christopher Newton-Cheh, MD, MPH, FAHA; Marco V. Perez, MD; Deepak Voora, MD, FAHA; Daniel Woo, MD, MS, FAHA; on behalf of the American Heart Association Committee on Molecular Determinants of Cardiovascular Health of the Council on Functional Genomics and Translational Biology and Council on Epidemiology and Prevention; Council on Cardiovascular Disease in the Young; Council on Cardiovascular and Stroke Nursing; Council on Cardiovascular Surgery and Anesthesia; Council on Clinical Cardiology; and Stroke Council</i>	e000037
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Correction to: Prevalence and Clinical Implication of Double Mutations in Hypertrophic Cardiomyopathy Revisiting the Gene-Dose Effect	e000038
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