

High-risk APOL1 not tied to CVD, stroke in older black women

9 July 2018



mean of 11 years; no differences were seen for other outcomes. A significantly increased hazard of hospitalized HFpEF was seen among carriers of high-risk versus low-risk *APOL1* variants in adjusted models (hazard ratio, 1.58; 95 percent confidence interval, 1.03 to 2.41). After adjustment for baseline eGFR, the correlation with HFpEF was attenuated and no longer significant (hazard ratio, 1.5; 95 percent confidence interval, 0.98 to 2.3).

"These findings do not support an association of high-risk *APOL1* genotypes with [coronary heart disease](#), stroke, or mortality in postmenopausal African-American women," the authors write.

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(HealthDay)—For postmenopausal African-American women, high-risk *APOL1* genotype seems not to be associated with coronary heart disease, stroke, or mortality, according to a study published online July 3 in *JAMA Cardiology*.

Nora Franceschini, M.D., M.P.H., from the University of North Carolina in Chapel Hill, and colleagues used data from the Women's Health Initiative to examine whether high-risk *APOL1* genotypes are associated with cardiovascular disease and stroke in postmenopausal African-American women. *APOL1* variants were genotyped or imputed from whole-exome sequencing.

The researchers found that high-risk *APOL1* [variant](#) carriers had increased prevalence of hypertension, use of cholesterol-lowering medications, and reduced estimated glomerular filtration rate (eGFR). Carriers of high-risk *APOL1* variants had increased incidence of hospitalized heart failure with preserved ejection fraction (HFpEF) compared with low-risk carriers after a

APA citation: High-risk APOL1 not tied to CVD, stroke in older black women (2018, July 9) retrieved 11 June 2022 from <https://medicalxpress.com/news/2018-07-high-risk-apol1-tied-cvd-older.html>

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