

Protocol Title

A Study of the Prevalence of Apolipoprotein L1 (APOL1) Alleles Among Individuals With Proteinuric Kidney Disease Who Are of Recent African Ancestry or Geographic Origin

Study Contact

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Study Status

Open for Enrollment

Brief Summary of the Protocol

If you have symptoms of kidney disease, or a family history of kidney disease, a clinical research study may be an option for you. Participation includes investigational genetic testing to see if you were born with the variation in your APOL1 gene that causes genetic APOL1-mediated nondiabetic chronic kidney disease (CKD) with an increased level of protein in your urine.

Sponsor Information

Vertex Pharmaceuticals Inc.

Protocol Number

VX19-NEN-801