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
Nelson Chen, Clinical Research Coordinator
646-317-0785
nec9039@nyp.org

Principal Investigator
Jay Lakkis, MD
212-746-7647

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