

THIS IS NOT A TEST REQUEST FORM.
 Please fill out this form and submit it with the test request form or electronic packing list.

PATIENT HISTORY FOR CADASIL (*NOTCH3* GENE) TESTING

Patient Name _____ Date of Birth _____ Sex F M
 Physician _____ Physician Phone _____
 Practice Specialty _____ Physician Fax _____
 Genetic Counselor _____ Counselor Phone _____

Patient's Ethnicity (check all that apply)

African-American Asian Hispanic Native American
 Ashkenazi Jewish Caucasian Middle Eastern Other: _____

Does the patient have symptoms? No Yes (check all that apply and describe)

Ischemic Events Mood Disorders Cognitive defects Migraine with aura
 Stroke Depression Memory loss Seizures
 Transient ischemic attacks (TIA) Personality changes Attention deficits
 Other symptom(s): _____

Has the patient undergone an MRI for brain imaging studies? No Yes Unknown

If yes, describe results: _____

Has the patient undergone previous DNA testing for CADASIL? No Yes Unknown

If yes, describe the results: _____ Normal Abnormal Unknown

Is there any relevant family history? No Yes Unknown

If yes, attach a pedigree or specify the relative's relationship to the patient. List their symptoms and age of onset:

Has DNA testing been performed for the family member(s)? No Yes Unknown

If yes, attach a copy of the relative's DNA laboratory result. (REQUIRED for familial mutation testing)

Check the test you intend to order.

- 3000531 CADASIL: Cerebral Autosomal Dominant Arteriopathy with Subcortical Infarcts and Leukoencephalopathy (NOTCH3), Sequencing;**
 Sequence analysis of NOTCH3 coding regions; clinical sensitivity 95%
- 2001961 Familial Mutation, Targeted Sequencing.** Tests for a mutation previously identified in a family member; a copy of relative's lab result is REQUIRED.

Master Label

For questions, contact an ARUP genetic counselor at (800) 242-2787, ext. 2141