

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:** Female Male

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/White 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 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