

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

PATIENT HISTORY FOR TUBEROUS SCLEROSIS COMPLEX PANEL, SEQUENCING AND DELETION/DUPLICATION

Patient Name: _____ Date of Birth: _____ Sex: Female Male
 Ordering Provider: _____ Provider's Phone: _____
 Practice Specialty: _____ Provider's Fax: _____
 Genetic Counselor: _____ Counselor Phone: _____

Patient's Ethnicity/Ancestry (check all that apply)

African American/Black Asian Hispanic White Other: _____

List country of origin (if known): _____

Does the patient have a diagnosis of Tuberos Sclerosis Complex (TSC)? Confirmed Suspected Unknown

Does the patient have clinical symptoms? No Yes (check all that apply) Unknown

- Skin**
 - Hypomelanotic macules (# _____)
 - Confetti skin lesions
 - Facial angiofibromas (# _____)
 - Shagreen patch
 - Ungula fibromas (# _____)
- Central Nervous System**
 - Subependymal nodules
 - Cortical dysplasias
 - Subependymal giant cell astrocytoma
 - Seizures
- Other symptom(s):** _____
- Cognitive impairment (specify: _____)
- Renal angiomyolipoma
- Renal cysts (# _____)
- Cardiac rhabdomyoma
- Pulmonary lymphangiioleiomyomatosis (LAM)
- Retinal hamartomas
- Retinal achromic patches

Does the patient have a FAMILY HISTORY of TSC or individuals with findings of TSC? No Yes Unknown

If yes, attach a PEDIGREE or specify the relatives' RELATIONSHIP to the patient. List their symptoms & age of onset:

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

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

If yes, please describe test(s) and results: _____

Check the test you intend to order OR write the test name and number below:

Testing for Tuberos Sclerosis Complex

- 3002100 Tuberos Sclerosis Complex Panel, Sequencing and Deletion/Duplication: NGS and CGH of *TSC1* and *TSC2*.
- 2001961 Familial Mutation, Targeted Sequencing: Targeted testing for a known familial sequence variant; a copy of the affected relative's lab report is REQUIRED.

Prenatal Testing for Tuberos Sclerosis Complex:

- 3002096 Tuberos Sclerosis Complex Panel, Sequencing and Deletion/Duplication, Fetal: NGS and CGH of *TSC1* and *TSC2* on cultured fetal samples (cultured villi or amniocytes).
- 2001980 Familial Mutation, Targeted Sequencing, Fetal: Targeted testing for a known familial sequence variant on a fetal sample; a copy of the affected relative's lab report is REQUIRED.

Master Label

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