

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

PATIENT HISTORY FOR *PTEN* HAMARTOMA TUMOR SYNDROME (PHTS) TESTING

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): _____

Clinical Diagnosis/Reason for Referral:

Bannayan-Riley-Ruvalcaba syndrome Intellectual disability/autism Proteus-like syndrome
 Cowden syndrome Proteus syndrome Other: _____

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

Breast cancer.....(age: _____) Macrocephaly
 Endometrial cancer.....(age: _____) Nevi (connective tissue epidermal)
 Follicular thyroid cancer(age: _____) Papillary thyroid cancer(age: _____)
 Intellectual disability / developmental delay Papillomatous papules
 Keratoses (location: _____) Pigmented macules of glans penis
 Lhermitte-Duclos disease (cerebellar tumor) Tissue overgrowth
 Lipomas Trichilemmomas
 GI hamartoma (describe) location: _____ number: _____
 Skeletal abnormalities (describe): _____
 Vascular malformation (describe): _____
 Other symptom(s): _____

Has the patient had an allogeneic bone marrow or umbilical cord blood transplant?..... No Yes Unknown

Has the patient undergone previous DNA testing No Yes Unknown
 If yes, describe the test(s) and results: _____

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.

- 2002470 *PTEN*-Related Disorders, Sequencing and Deletion/Duplication:** Sequencing and deletion/duplication analysis of *PTEN* coding regions and intron/exon boundaries with a clinical sensitivity of 85% for Cowden, 65% for Bannayan-Riley-Ruvalcaba, and 20% for Proteus syndrome.
- 2001961 Familial Mutation, Targeted Sequencing:** Tests for a mutation previously identified in a family member; a copy of relative's lab result is **REQUIRED**.
- 3003144 Deletion/Duplication Analysis by MLPA:** Tests for large deletion/duplication previously identified in a family member; a copy of a relative's lab report is **REQUIRED**.



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