

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

NEUROFIBROMATOSIS TYPE 1 (NF1) AND LEGIUS SYNDROME PATIENT HISTORY FORM

Patient Name: _____ **Date of Birth:** _____

Sex Assigned at Birth: Female Male Intersex **Gender Identity (optional):** Female Male _____

Ordering Provider: _____ **Provider's Phone:** _____

Practice Specialty: _____ **Provider's Fax:** _____

Genetic Counselor: _____ **Counselor's 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 symptoms? No Yes (check all that apply and describe)

- | | |
|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------------------------|
| <input type="checkbox"/> Axillary or inguinal freckling | <input type="checkbox"/> Malignant peripheral nerve sheath tumor (MPNST) |
| <input type="checkbox"/> Café au lait macules (indicate number) <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input type="checkbox"/> 5 <input type="checkbox"/> 6 <input type="checkbox"/> 7+ | <input type="checkbox"/> Optic glioma (age at dx): _____ |
| <input type="checkbox"/> Dermal fibromas | <input type="checkbox"/> Osseous lesion (describe): _____ |
| <input type="checkbox"/> Learning disabilities | <input type="checkbox"/> Overgrowth (describe): _____ |
| <input type="checkbox"/> Lisch nodules (iris hamartomas) | <input type="checkbox"/> Scoliosis |
| | <input type="checkbox"/> Other symptoms: _____ |

Has the patient undergone previous germline DNA testing? No Yes Unknown

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

Does this patient have NF1 or SPRED1 genetic variant(s) previously identified in tumor/bone marrow?

..... No Yes Unknown

If yes, attach results and describe: _____

Is there any relevant family history? No Yes; NF1 Yes; Legius 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 variant testing).

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

Master Label