

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

FAMILIAL TRANSTHYRETIN (TTR) AMYLOIDOSIS PATIENT HISTORY

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 of any of the following TTR phenotypes? No Yes (check all that apply)

TTR amyloid polyneuropathy TTR leptomeningeal amyloidosis
 TTR cardiac amyloidosis TTR oculoleptomeningeal amyloidosis
 Other: _____

Does the patient have symptoms of TTR amyloidosis? No Yes (check all that apply)

<input type="checkbox"/> Peripheral sensory/motor neuropathy <ul style="list-style-type: none"> <input type="checkbox"/> Loss of sensation in feet or hands <input type="checkbox"/> Prickling/burning/numbness in extremities <input type="checkbox"/> Muscle weakness <input type="checkbox"/> Difficulty walking or using arms/hands <input type="checkbox"/> Autonomic neuropathy <ul style="list-style-type: none"> <input type="checkbox"/> Sweating too much or too little <input type="checkbox"/> Vomiting/delayed gastric emptying <input type="checkbox"/> Constipation/diarrhea <input type="checkbox"/> Sexual impotence <input type="checkbox"/> Orthostatic fainting <input type="checkbox"/> Urinary incontinence or retention <input type="checkbox"/> Nephropathy <input type="checkbox"/> Vitreous opacities	<input type="checkbox"/> CNS Findings <ul style="list-style-type: none"> <input type="checkbox"/> Visual impairment <input type="checkbox"/> Dementia <input type="checkbox"/> Ataxia/spasticity <input type="checkbox"/> Seizures <input type="checkbox"/> Psychosis <input type="checkbox"/> Hydrocephalus <input type="checkbox"/> Cardiac Disease <ul style="list-style-type: none"> <input type="checkbox"/> Cardiomegaly <input type="checkbox"/> Conduction block <input type="checkbox"/> Angina <input type="checkbox"/> Congestive heart failure <input type="checkbox"/> Aortic dissection/dilation <input type="checkbox"/> Other symptom(s): _____
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Other laboratory testing for TTR amyloidosis: Not performed Performed (describe below) Unknown

Congo red staining/immunochemistry Normal Abnormal: _____ Uncertain: _____
 Serum TTR testing by mass spec Normal Abnormal: _____ Uncertain: _____

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 variant testing).

Master Label

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