

This requisition form can be used to submit a specimen for Alnylam Act™, a sponsored genetic testing program brought to you by Alnylam Pharmaceuticals and Invitae Corporation. Please confirm that the patient meets the eligibility requirements for the program. To submit orders for genetic testing outside of this program, please order through Invitae's online portal or use a standard requisition form, accessible at www.invitae.com/order-forms.

PATIENT INFORMATION		
First name	MI	Last name
Date of birth (MM/DD/YYYY)	Sex <input type="radio"/> M <input type="radio"/> F	MRN (medical record number)
Ancestry <input type="radio"/> Asian <input type="radio"/> Black/African American <input type="radio"/> White/Caucasian <input type="radio"/> Ashkenazi Jewish <input type="radio"/> Hispanic <input type="radio"/> Native American <input type="radio"/> Pacific Islander <input type="radio"/> French Canadian <input type="radio"/> Sephardic Jewish <input type="radio"/> Mediterranean <input type="radio"/> Other:		
Phone	Email address	
Address		City
State	ZIP code	Country

ORGANIZATION INFORMATION	
Organization name and address	
Organization name	
Phone	Fax
Address	
State	ZIP code
Country	
Primary clinical contact	
Name	Role/title
Phone	NPI
Email address (for report access)	
Ordering physician	
<input type="radio"/> Same as primary clinical contact	
Name	NPI
Email address (for report access)	
Additional clinical or laboratory contact (optional)	
Name	Email address (for report access)

SPECIMEN INFORMATION	
Label each tube with the patient's full name, date of birth, and specimen collection date. A requisition form MUST accompany each specimen. www.invitae.com/specimen-requirements	
Specimen type : <input type="radio"/> Blood <input type="radio"/> Saliva <input type="radio"/> Assisted saliva <input type="radio"/> DNA - source: <i>DNA must be extracted in a CLIA or other suitably certified laboratory</i> <i>We are unable to accept blood/saliva from patients with:</i> <ul style="list-style-type: none"> Allogeneic bone marrow transplants Blood transfusion <2 weeks prior to specimen collection 	
Collection date (MM/DD/YYYY)	<i>If not provided, date will be 1 day prior to our receipt of specimen. For DNA, provide date retrieved from archive.</i>
Special cases : <input type="radio"/> History of/current hematologic malignancy <input type="radio"/> Resubmission	

REASON FOR TESTING	
Previous results (if applicable and not included in clinical criteria below)	

INVITAE PARTNER CODE	TTR
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ALNYLAM ACT (hATTR AMYLOIDOSIS) ELIGIBILITY CRITERIA

Patient must be age 18 or older

AND

Have a family history or suspected diagnosis of hATTR amyloidosis with one or more of the following signs and symptoms. Please check all that apply:

<input type="checkbox"/> Family history of hATTR amyloidosis	<input type="checkbox"/> Spinal stenosis or spinal radiculopathy
<input type="checkbox"/> Sensory and/or motor neuropathy (e.g., neuropathic pain, alternation sensation [sensitivity to pain and temperature], numbness and tingling, muscle weakness, impaired balance, difficulty walking)	<input type="checkbox"/> Renal abnormalities (e.g., renal insufficiency and/or proteinuria)
<input type="checkbox"/> Autonomic dysfunction (e.g., nausea and vomiting, changes in GI motility [diarrhea, constipation, gastroparesis, early satiety], orthostatic hypotension [fainting and dizziness upon standing], sexual dysfunction, bladder dysfunction)	<input type="checkbox"/> Ocular changes (e.g., blurred vision, blindness, dry eyes, glaucoma, visual field abnormalities, retinal detachment)
<input type="checkbox"/> Heart disease (e.g., shortness of breath, edema, palpitations, arrhythmias, conduction abnormalities, heart failure, abnormal cardiac imaging [echo, MRI, or technetium])	<input type="checkbox"/> Biopsy positive for amyloid
<input type="checkbox"/> Bilateral carpal tunnel syndrome	<input type="checkbox"/> Other (must be completed if checked): _____

FAMILY VARIANT TESTING			
Invitae's family variant testing programs involves full analysis of the gene in which the original family member's variant was identified. For more information, visit www.invitae.com/family-testing .			
Please attach the proband's clinical report or provide Invitae RQ#			
INVITAE PROBAND RQ#	RELATIONSHIP TO PROBAND	GENE(S)	VARIANT(S)

RE-REQUISITION

The Alnylam Act program offers one re-requisition at no additional charge within 90 days to the Invitae Comprehensive Neuropathies Panel and/or the Invitae Cardiomyopathy Comprehensive Panel. For more information and to request online, please visit www.invitae.com/re-requisition.

ASSAY

Invitae is a CAP-accredited and CLIA-certified clinical diagnostic laboratory performing full-gene sequencing and deletion/duplication analysis using next-generation sequencing technology (NGS). Search for details on the analysis of any gene in our test catalog at www.invitae.com/physician/search.

To request a complimentary specimen collection kit visit www.invitae.com/request-a-kit

SHIPPING INSTRUCTIONS

Please ship specimen overnight in insulated containers:

Attn: Invitae Client Services
1400 16th Street, San Francisco, CA 94103, USA

Invitae continually updates its panels based on the most recent evidence. Please note that if an order is placed using an older version of this form, Invitae reserves the right to upgrade any ordered panel(s) to the current version(s).

TESTS INCLUDED IN THE PROGRAM

Test code	Test name	# of genes	Gene list
<input type="radio"/> 02251	Invitae Cardiomyopathy Comprehensive Panel	50	ABCC9, ACTC1, ACTN2, AGL, BAG3, CACNA1C, CAV3, CRYAB, CSRP3, DES, DMD, DOLK, DSC2, DSG2, DSP, EMD, EYA4, FHL1, FKRP, FKTN, FLNC, GAA, GLA, HCN4, JUP, LAMP2, LMNA, MYBPC3, MYH7, MYL2, MYL3, PKP2, PLN, PRKAG2, RAF1, RBM20, RYR2, SCN5A, SGCD, SLC22A5, TAZ, TCAP, TMEM43, TNNC1, TNNI3, TNNT2, TPM1, TTN, TTR, VCL
<input type="radio"/> 03200	Invitae Comprehensive Neuropathies Panel	72	AARS, AIFM1, ATL1, ATL3, ATP7A, BICD2, BSCL2, CHCHD10, DCTN1, DNAJB2, DNMT2, DNMT1, DST, DYNC1H1, EGR2, FAM134B, FBXO38, FGD4, FIG4, GAN, GARS, GDAP1, GJB1, GNB4, HARS, HINT1, HSPB1, HSPB8, IGHMBP2, IKBKAP, INF2, KIF1A, LITAF, LMNA, LRSAM1, MED25, MFN2, MORC2, MPZ, MTMR2, NDRG1, NEFL, NGF, NTRK1, PDK3, PLEKHG5, PMP22, PRPS1, PRX, RAB7A, REEP1, SBF2, SCN11A, SCN9A, SH3TC2, SIGMAR1, SLC25A46, SLC52A2, SLC52A3, SLC5A7, SPG11, SPTLC1, SPTLC2, TFG, TRIM2, TRPV4, TTR, UBA1, VAPB, VRK1, WNK1, YARS
<input type="radio"/> 02265	Invitae Transthyretin Amyloidosis Test	1	TTR

By signing this form, the medical professional acknowledges that the individual/family member authorized to make decisions for the individual (collectively, the "Patient") has been supplied information regarding and consented to undergo genetic testing, substantially as set forth in Invitae's Informed Consent for Genetic Testing (www.invitae.com/patient-consent). And in connection with the Alnylam Act™ program, the Patient has been informed that Invitae may notify them of clinical updates related to genetic test results (in consultation with the ordering medical professional as indicated) and has been informed that de-identified Patient data may be used and shared with third parties, including Alnylam Pharmaceuticals, Inc. ("Alnylam"), for research and commercial purposes and, in the U.S., to contact their medical professional. For orders originating in Canada, the Patient has been informed that their personal information and specimen will be transferred to and processed in the U.S. and that de-identified Patient data may be used and shared for research and commercial purposes in the U.S. The medical professional warrants that he/she will not seek reimbursement for this sponsored test from any third party, including but not limited to U.S. federal healthcare programs. The medical professional also hereby acknowledges that organization and clinician contact information provided in the order may be shared with third parties, including Alnylam, that may contact the medical professional directly in connection with the Alnylam Act program, or Alnylam products. The medical professional understands that the use of this sponsored test is not intended to be, nor should it be construed as, either express or implied, an obligation or inducement for the medical professional to recommend, purchase, order, prescribe, promote, administer or otherwise support any Alnylam product or any other Invitae product or service. In addition to the above, I attest that I am the ordering physician, or I am authorized by the ordering physician to order this test, or I am authorized under applicable law to order this test.

Medical professional signature

Date