

Test Requisition Form

PATIENT INFORMATION (Patient must be 18 years or older)						
Name (Last, First, MI)		Date of Birth (MM/DD/YY)		Phone Number	Email	
Address		City	State	Zip	Biological Sex <input type="checkbox"/> F <input type="checkbox"/> M	Ethnicity: <input type="checkbox"/> African American <input type="checkbox"/> Asian <input type="checkbox"/> Caucasian <input type="checkbox"/> Hispanic <input type="checkbox"/> Jewish <input type="checkbox"/> Other:
SPECIMEN INFORMATION*						
Type(s) <input type="checkbox"/> Blood <input type="checkbox"/> Saliva <input type="checkbox"/> Send saliva kit to patient** <input type="checkbox"/> Phlebotomy request*				<input type="checkbox"/> Personal history of allogenic bone marrow or peripheral stem cell transplant		
Collection Date		Specimen ID			Medical Record #	
<small>*Blood or saliva from patients with active/recent hematological disease will undergo additional review and may not be accepted in some cases. For these, cultured fibroblasts or fresh/fresh frozen normal tissue are preferred. See ambrygen.com/specimen-requirements for details. **By checking this box and submitting the completed form, a saliva kit will be sent to the patient's address above. Your patient will be able to submit a saliva sample directly to Ambry for testing. * Available for US patients only. As the patient's clinician, I am unaware of any potential for complication or difficulty in drawing blood for the listed patient(s). I understand that the phlebotomist has full authority to refuse to draw any patient if the safety of the phlebotomist and/or patient(s) are in question.</small>						
ORDERING PHYSICIAN/SENDING FACILITY (Each listed person will receive a copy of the report)						
Facility Name (Facility Code)		Address		City	State /Country	Zip Phone
Ordering Licensed Provider Name (Last, First)(Code)		NPI#	Phone		Fax	Email
Additional Results Recipients						
Genetic Counselor or Other Medical Provider Name (Last, First) (Code)				Phone/Fax/Email		
PATIENT ELIGIBILITY: PLEASE READ CAREFULLY						
Patients must be 18 years and older and have a family history of hATTR amyloidosis OR have at least 2 of the following clinical symptoms/manifestations related to hATTR amyloidosis. Does the patient have a family history of hATTR amyloidosis? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Patient does not know SYMPTOM CHECKLIST (Please check if the patient has previously had or currently has any of the following clinical manifestations and indicate symptom onset age.)						
<input type="checkbox"/> Sensory dysfunction (e.g. numbness and tingling in feet and/or hands, sensitivity to pain and temperature, pain in extremities); Symptom onset age: _____			<input type="checkbox"/> Bilateral Carpal tunnel syndrome; Symptom onset age: _____			
<input type="checkbox"/> Motor dysfunction (e.g. muscle weakness, impaired balance, difficulty walking); Symptom onset age: _____			<input type="checkbox"/> Lumbar spinal stenosis; Symptom onset age: _____			
<input type="checkbox"/> Autonomic dysfunction (e.g., orthostatic hypotension, early satiety, nausea and vomiting, changes in GI motility, erectile dysfunction, bladder dysfunction); Symptom onset age: _____			<input type="checkbox"/> Unintentional weight loss; Symptom onset age: _____			
<input type="checkbox"/> Gastrointestinal (diarrhea or constipation not responding to typical therapy, or alternating bouts of diarrhea/constipation); Symptom onset age: _____			<input type="checkbox"/> Myocardial radiotracer (^{99m} Tc-PYP/DPD/HMDP) uptake on bone scintigraphy and the absence of a monoclonal protein in serum or urine; Symptom onset age: _____			
<input type="checkbox"/> Heart disease (e.g., shortness of breath, edema, fatigue, palpitations, and arrhythmias); Symptom onset age: _____			<input type="checkbox"/> Histological confirmation and typing of TTR amyloid; Symptom onset age: _____			
<input type="checkbox"/> Renal issues (proteinuria or renal insufficiency/failure); Symptom onset age: _____			<input type="checkbox"/> Other _____ Approximately, how many doctors has the patient seen about this condition? _____			
CHECK TO ORDER	TEST NAME	TEST CODE	# OF GENES	GENE LIST		
Please select only one test.						
<input type="checkbox"/>	Transthyretin amyloidosis	1560	1	TTR		
<input type="checkbox"/>	CardioNext (includes TTR)	8911	92	TTR, ABCC9, ACTC1, ACTN2, ALMS1, ALPK3, AKAP9, ANK2, ANKRD1, BAG3, CACNA1C, CACNA2D1, CACNB2, CALM1, CALM2, CALM3, CASQ2, CAV3, CRYAB, CSRP3, DES, DMD, DOLK, DSC2, DSG2, DSP, EMD, EYA4, FHL1, FKRP, FLNC, FKTN, FXN, GATA4, GATAD1, GLA, GPD1L, HCN4, JAG1, JPH2, JUP, KCND3, KCNE1, KCNE2, KCNE3, KCNH2, KCNJ2, KCNJ5, KCNJ8, KCNQ1, LAMA4, LAMP2, LDB3, LMNA, MYBPC3, MYH6, MYH7, MYL2, MYL3, MYOZ2, MYPN, NEXN, NKX2-5, PKP2, PLN, PRKAG2, PTPN11, RAF1, RBM20, RIT1, RYR2, SCN1B, SCN2B, SCN3B, SCN4B, SCN5A, SCN10A, SNTA1, SOS1, TAZ, TBX1, TBX20, TBX5, TCAP, TECRL, TGFβ3, TMEM43, TMPO, TNNC1, TNNT3, TNNT2, TPM1, TRDN, TRPM4, TTN, TXNRD2, VCL		
<input type="checkbox"/>	NeuropathySelect (includes TTR)	9570	80	TTR, AARS, AIFM1, APOA1, ATL1, ATL3, ATP7A, BICD2, BSCL2, CHCHD10, DCTN1, DNAJB2, DNM2, DNMT1, DST, DYNC1H, EGR2, FAM134B, FBXO38, FGD4, FIG4, FUS, GAN, GARS, GDAP1, GJB1, GNB4, GSN, HARS, HINT1, HSPB1, HSPB8, IGHMBP2, IKBKAP, INF2, KIF1A, LITAF, LMNA, LRSAM1, MARS, MFN2, MORC2, MPZ, MTMR2, NDRG1, NEFH, NEFL, NGF, NTRK1, OPTN, PDK3, PLEKHG5, PMP22, PRDM12, PRPS1, PRX, RAB7A, REEP1, SBF2, SCN11A, SCN9A, SETX, SH3TC2, SIGMAR1, SLC25A46, SLC52A2, SLC52A3, SLC5A7, SPG11, SPTLC1, SPTLC2, TARDBP, TFG, TRPV4, UBA1, VAPB, VRK1, WNK1, YARS, VCP		
Genetic Counseling: Ambry and Akcea have partnered with PWNHealth to offer no cost, pre- and/or post-test genetic counseling for your patients. Genetic counseling is not required for testing. By checking the boxes below, I agree to allow Ambry to facilitate the provision of pre-test and/or post-test genetic counseling services by PWNHealth. <input type="checkbox"/> Yes. I request a pre-test genetic counseling session for my patient. <input type="checkbox"/> Yes. I request a post-test genetic counseling session for my patient.						
CONFIRMATION OF INFORMED CONSENT AND MEDICAL NECESSITY FOR GENETIC TESTING The undersigned person (or representative thereof) ensures he/she is a licensed medical professional authorized to order genetic testing and confirms that the patient has given appropriate consent. I confirm testing is medically necessary and test results may impact medical management for the patient. Furthermore, all information on this ordering form is true to the best of my knowledge. I understand that organization and clinician contact information provided may be shared with third parties including Akcea Therapeutics and Backpack Health and may contact you in connection with the Genetic Pathway Program. My patient would like to receive more information from Backpack Health and consents to share patient's name and email with this third party (unless this box is checked <input type="checkbox"/>).						
Signature Required for Processing Medical Professional Signature:					Date:	
To request a complimentary specimen collection kit visit: ambrygen.com/clinician/order-sample-kit						