

COMPLETE ENTIRE FORM TO AVOID DELAYS

PATIENT INFORMATION						
Name (Last, First, MI)		Date of Birth (MM/DD/YY)	Date of Death (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 (Ashkenazi) <input type="checkbox"/> Portuguese <input type="checkbox"/> Other:	
SPECIMEN INFORMATION* (For phlebotomy service, select all services you are requesting)						
Type(s) <input type="checkbox"/> Blood (EDTA preferred) <input type="checkbox"/> Saliva <input type="checkbox"/> DNA <input type="checkbox"/> Other:			<input type="checkbox"/> Personal history of allogenic bone marrow or peripheral stem cell transplant			
Collection Date	Specimen ID		Medical Record #			
*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.						
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			
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 that testing is medically necessary and that test results may impact medical management for the patient. Furthermore, all information on this TRF is true to the best of my knowledge. My signature applies to the attached letter of medical necessity (unless this box is checked <input type="checkbox"/>).						
Signature Required for Processing Medical Professional Signature:				Date:		
INSURANCE BILLING (Include copy of both sides of insurance card)			INSTITUTIONAL BILLING			
Patient Relation to Policy Holder? <input type="checkbox"/> Self <input type="checkbox"/> Spouse <input type="checkbox"/> Child		Name and DOB of Policy Holder (if not self)		Facility Name <input type="checkbox"/> Send invoice to facility address above		
Insurance Company	Policy #	HMO Auth #		Address		
Ambry Genetics preverifies insurance coverage and will contact the patient after the patient's sample is received if the out-of-pocket amount for testing is estimated to exceed (Nothing checked defaults to >\$100): <input type="checkbox"/> \$100 <input type="checkbox"/> Any amount <input type="checkbox"/> Other \$			Contact Name		E-mail/Fax	
<input type="checkbox"/> Hold order pending patient contact and approval of payment terms regarding out-of-pocket. Patient preferred method of contact regarding out-of-pocket amount: <input type="checkbox"/> Email <input type="checkbox"/> Phone			Phone Number			
			<input type="checkbox"/> PATIENT PAYMENT		<input type="checkbox"/> Check (Payable to Ambry Genetics) <input type="checkbox"/> Credit Card (Call 949-900-5795)	
Representative Acknowledgement: I acknowledge that the information provided by me is true and correct. For payment by credit card: I hereby authorize Ambry Genetics Corporation to bill my credit card as indicated above. In order to expedite consideration for eligibility for Ambry's E.P.I.C. Program , please provide the total annual gross household income: \$ [] and the number of family members in the household supported by the listed income: []. I authorize Ambry Genetics Corporation to verify the above information for the sole purpose of assessing financial need, including the right to seek supporting documentation.						
FOR NY RESIDENTS:						
<input type="checkbox"/> I am a New York resident and I give Ambry Genetics permission to store my sample for longer than 60 days. NOTE: If left blank, consent is interpreted as "NO".						
Signature Required For Insurance/Self-Pay Patients and NY Sample Storage Consent:				Date:		

REQUIRED ORDERING CHECKLIST

- Clinic notes (with pedigree if available)
- ICD-10 code(s)
- Clinician & patient signatures
- Insurer-specific forms (i.e. ABN), if applicable
- Front/back copy of insurance card(s)

Sudden Death Test Requisition Form - Page 2 of 2

INDICATION(S) FOR TESTING

Diagnostic Family history Positive or normal control Other: _____
 ICD-10 code(s): _____

PATIENT HISTORY No personal history of cardiovascular disease No personal history of neurological disease

PLEASE SUPPLY CLINIC NOTES AND PEDIGREE
CARDIOVASCULAR MEDICAL HISTORY

Sudden cardiac arrest Y N (if yes): # Episodes: _____ Age first incident: _____
 # Episodes: _____ Age first incident: _____
 Syncope Y N (if yes): # Episodes: _____ Age first incident: _____
 History of cardiomyopathy Y N Age at dx: _____
 HCM DCM ARVD LVNC RCM
 Other cardiomyopathy Types: _____
 History of Arrhythmia Y N Age at dx: _____
 Long QT Short QT Brugada CPVT ARVD
 Other arrhythmia Types: _____
Other features/syndromes
 Clinical diagnosis of Marfan Syndrome
 Aortic Aneurysm/Dilation Age at dx: _____
 Other Aneurysm Location: _____ Age at dx: _____
 Aortic/Vascular Dissection Location: _____ Age at dx: _____
 Arterial tortuosity BAV MVP
 Congenital Heart Defect Type: _____
 Ectopia lentis Myopia Marfanoid habitus
 Pectus deformity Type: _____
 Scoliosis Joint Hypermobility Joint contractures
 Pneumothorax Craniosynostosis
 Facial clefting, Type : _____

Xanthoma(s) Epistaxis (nosebleeds) Telangiectasia
 AVM Location: _____
 Amyloidosis Age at dx: _____
 Neuromuscular disease Specify: _____
 Hearing Loss Describe: _____
 Genetic syndrome Specify: _____
 Other Specify: _____

NEUROLOGICAL MEDICAL HISTORY

Seizure History N/A Age at first unprovoked seizure: _____
 Was this patient diagnosed with an epilepsy syndrome? yes no unknown
 If yes, please specify: _____

Prior Testing and Procedures:
FAMILY HISTORY

Mother: Unaffected Affected, list symptoms/dx: _____
 _____ Dx age: _____
 Father: Unaffected Affected, list symptoms/dx: _____
 _____ Dx age: _____
 Relationship to patient: _____ Maternal Paternal
 Diagnosis: _____ Dx age: _____

Please check the box next to the test(s) being ordered below. All tests include gene sequence and deletion/duplication analyses, unless otherwise indicated. If this TRF is sent to Ambry without or ahead of the sample, it will be treated as a preverification. If test ordered is different than the test preverified, we will honor what is on the TRF order form with the sample.

For multiple test orders, testing will be run concurrently (multiple tests initiated at the same time) unless otherwise specified. To order reflexive testing (second test starts pending first test outcome), please clearly indicate the order of reflexive tests in the notes section or next to the test check box. For reflex test orders, any positive findings (pathogenic/likely pathogenic) in the first test will be reported out to the clinician, and the requested second test will be canceled; all other findings will automatically reflex (including VUS).

Check to order	Test Name	Test Code	Description	Check to order	Test Name	Test Code	Description
Arrhythmia				Other Cardiac Findings			
<input type="checkbox"/>	RhythmNext	8900	36 genes for long QT syndrome, Brugada syndrome, and other inherited arrhythmias	<input type="checkbox"/>	FHNext	8680	4 genes (APOB, LDLR, LDLRAP1, PCSK9)
<input type="checkbox"/>	CPVTNext	8902	6 genes for catecholaminergic polymorphic ventricular tachycardia	<input type="checkbox"/> Check this box if you would like to have the <i>SLC01B1</i> c.521T>C polymorphism reported, which has been associated in medical literature with statin-induced myopathies			
Cardiomyopathy				<input type="checkbox"/>	TAADNext	8789	22 genes for thoracic aortic aneurysms/dissections, Marfan syndrome, and related disorders
<input type="checkbox"/>	HCMNext	8936	27 genes for hypertrophic cardiomyopathy	Seizures/Epilepsy			
<input type="checkbox"/>	DCMNext	8884	36 genes for dilated cardiomyopathy	<input type="checkbox"/>	EpilepsyNext	7019	100 genes for epilepsy
<input type="checkbox"/>	ARVDNext	8904	9 genes for arrhythmogenic right ventricular dysplasia	SINGLE SITE ANALYSIS (Please include a copy of relative's report)			
<input type="checkbox"/>	LVNCNext	8906	8 genes for left ventricular non-compaction	Gene(s): _____ Mutation(s): _____			
<input type="checkbox"/>	CMNext with <i>TTN</i>	8886	55 genes for hereditary cardiomyopathy	Relative Name: _____			
Cardiomyopathy and Arrhythmia				Relationship to Relative: _____ Accession # (If tested at Ambry): _____			
<input type="checkbox"/>	CardioNext with <i>TTN</i>	8911	85 genes for hereditary cardiomyopathies and arrhythmias	Positive control sample: <input type="checkbox"/> Will be provided <input type="checkbox"/> Already at Ambry <input type="checkbox"/> Not available			
ADDITIONAL TESTS							
Test: _____		Test code: _____		Test: _____		Test code: _____	
Test: _____		Test code: _____		Test: _____		Test code: _____	