

Hereditary Cardiomyopathies and Arrhythmias: Patient Information

Instructions: Accurate interpretation and reporting of genetic results is contingent upon the reason for testing, clinical information, ethnic background/ancestry, and family history. To help provide the best possible service, supply the information requested below and **send paperwork with the specimen or return by fax to Mayo Clinic Laboratories, Attn: Molecular Technologies Laboratory Genetic Counselors at 507-284-1759. Phone: 800-533-1710 / International clients: +1-507-266-5700 or email MLIINT@mayo.edu**

Patient Information

Patient Name (Last, First Middle)		Birth Date (mm-dd-yyyy)
Sex Assigned at Birth <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Unknown <input type="checkbox"/> Choose not to disclose	Legal/Administrative Sex <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Nonbinary	

Referring Provider Information

Referring Provider Name (Last, First)	Phone	Fax*
Other Contact Name (Last, First)	Phone	Fax*

*Fax number given must be from a fax machine that complies with applicable HIPAA regulations.

Is this a postmortem specimen? Yes No If "Yes," attach autopsy report if available.

Reason for Testing

 Check all that apply.

Diagnosis Family history** Sudden death

**Genetic testing should be performed on an affected family member first, when possible. FMTT / Familial Variant, Targeted Testing should be ordered when there is a previous positive genetic test result in the family.

Clinical History

 Attach medical records/diagnostic tests.

Diagnosis

Is this patient affected by one or more of the following? Yes No If "Yes," check all that apply:

HCM DCM ARVC LVNC Other cardiomyopathy: _____

CPVT Brugada Long QT Other arrhythmia: _____

Other: _____

Age at diagnosis: _____

Has patient had:

Sudden cardiac arrest Yes No Describe: _____

Sudden cardiac death Yes No Describe: _____

Syncope Yes No Describe: _____

ARVC: RV fatty infiltration Yes No

Arrhythmia: Maximum QTc interval _____ msec

Conduction system disease Yes No Describe: _____

Cardiomyopathy:

LV hypertrophy Yes No Maximum LV wall thickness _____ mm

LV Dilation Yes No LV internal diameter, diastole _____ mm

Ejection fraction _____%

Other Relevant Information

Hereditary Cardiomyopathies and Arrhythmias: Patient Information (continued)

Patient Information (required)

Patient Name (Last, First Middle)	Patient ID (Medical Record Number)
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Family History

Are there similarly affected relatives? <input type="checkbox"/> Yes <input type="checkbox"/> No If "Yes," indicate relationship and symptoms: _____
Have any family member had genetic testing? <input type="checkbox"/> Yes*** <input type="checkbox"/> No <input type="checkbox"/> Unknown ***FMTT / Familial Variant, Targeted Testing should be ordered when there is a previous positive genetic test result in the family. Contact the lab for ordering assistance.
History of consanguinity: <input type="checkbox"/> No <input type="checkbox"/> Yes; relationship details: _____

Ancestry

<input type="checkbox"/> African American	<input type="checkbox"/> East Asian	<input type="checkbox"/> Latinx/Latine	<input type="checkbox"/> South Asian	<input type="checkbox"/> Choose not to disclose
<input type="checkbox"/> Ashkenazi Jewish	<input type="checkbox"/> European	<input type="checkbox"/> Middle Eastern	<input type="checkbox"/> None of the above	<input type="checkbox"/> Unknown

New York State Patients: Informed Consent for Genetic Testing is required. See Informed Consent for Genetic Testing (T576), Informed Consent for Genetic Testing – Spanish (T826), or Informed Consent for Genetic Testing for Deceased Individuals (T782).