



# CARDIOGENETICS PROGRAM REFERRAL FORM

**Patient Name:** \_\_\_\_\_

**Phone:** \_\_\_\_\_ **Date of Birth:** \_\_\_\_\_

**Date:** \_\_\_\_\_

**SERVICES:**

- Genetic counseling ONLY (with a certified genetic counselor)
- Evaluation with a cardiology specialist AND genetic counseling

**REASON FOR REFERRAL** (check all that apply):

|   |   |
|---|---|
| <b>CARDIOMYOPATHY</b>   | <b>AORTOPATHY</b>   |
| <input type="checkbox"/> Hypertrophic cardiomyopathy (HCM), obstructive<br><input type="checkbox"/> Hypertrophic cardiomyopathy (HCM), non-obstructive<br><input type="checkbox"/> Dilated cardiomyopathy<br><input type="checkbox"/> Peripartum cardiomyopathy<br><input type="checkbox"/> Left ventricular noncompaction (LVNC)<br><input type="checkbox"/> Arrhythmogenic right ventricular cardiomyopathy (ARVC/D)<br><input type="checkbox"/> Congestive heart failure (CHF) | <input type="checkbox"/> Aortic dissection<br><input type="checkbox"/> Aortic aneurysm, unspecified site<br><input type="checkbox"/> Thoracic aortic dissection<br><input type="checkbox"/> Thoracic aortic aneurysm<br><input type="checkbox"/> Family history of aortic problem<br><input type="checkbox"/> Marfan syndrome (pt. has clinical diagnosis)<br><input type="checkbox"/> Ehler’s-Danlos syndrome (pt. has clinical diagnosis) |
| <b>ARRHYTHMIA</b>   | <b>SIGNS/SYMPTOMS</b>   |
| <input type="checkbox"/> Long QT syndrome<br><input type="checkbox"/> Brugada syndrome<br><input type="checkbox"/> Catecholaminergic polymorphic ventricular tachycardia (CPVT)<br><input type="checkbox"/> Abnormal ECG (excludes LQTS)<br><input type="checkbox"/> Personal history of cardiac arrest<br><input type="checkbox"/> s/p ICD in situ<br><input type="checkbox"/> s/p Pacemaker in situ   | <input type="checkbox"/> Syncope and collapse<br><input type="checkbox"/> Dizziness/vertigo<br><input type="checkbox"/> Palpitations<br><input type="checkbox"/> Shortness of breath  |
| <b>CORONARY ARTERY DISEASE</b>  | <b>GENETICS/FAMILY HISTORY</b>  |
| <input type="checkbox"/> Familial hypercholesterolemia  | <input type="checkbox"/> Healthy patient who has a gene mutation<br><input type="checkbox"/> Family history of a gene mutation<br><input type="checkbox"/> Family history of sudden cardiac death<br><input type="checkbox"/> Family history of sudden cardiovascular disease<br><br>If known, describe gene mutation or family history:<br>_____<br>_____<br>_____<br>_____  |
| <b>OTHER</b>  |   |
| <input type="checkbox"/> _____<br><input type="checkbox"/> _____  |   |

**Referring Physician Name:** \_\_\_\_\_

**Phone Number:** \_\_\_\_\_ **Fax Number:** \_\_\_\_\_

**Address:** \_\_\_\_\_

**Signature:** \_\_\_\_\_

Please fax this form and, if available, pertinent medical records (most recent cardiology visit note, echocardiogram, cardiac MRI, lab results and electrocardiogram). Please also fax information regarding previously completed genetic testing to 310-423-6795.