HELICE
GENOMIC SCIENCES, INC.

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Received on (MMDDYYYY): _____

☐ Call ☐ Email Your Accession # _____

WELLNESS SCREENING REQUISITION FORM

PATIENT INFORMATION

Last Name	First Name	MI

Date of Birth	Gender	Email Address
	☐ M ☐ F	

Address	Home Number Mobile Number

Today's Date	Collection Date

Ethnicity ☐ African American ☐ Ashkenazi ☐ Asian ☐ Caucasian ☐ Hispanic
☐ Other (please specify) _____

REFERRING PHYSICIAN INFORMATION

Physician's Name:

License Number:

Specialty:

Affiliated Hospital:

Address:

Contact Number:

Email Address:

SEND REPORT TO

SERVICE MENU SELECT A TEST. WHEN APPLICABLE, ALSO SELECT NUMBER OF GENES.

- 01 **ArtheroGxOne™** (84 genes)
- 02 **CardioGxOne™** (213 genes)
- 03 **PGxCardio™** (213 genes)

CardioGxOne™ Subpanels:

- 04 Cardiomyopathies (149 genes)
- 05 Arrhythmias (141 genes)

- 06 Reflex to PGxCardio™
- 07 Reflex to PGxCardio™
- 08 Ventricular arrhythmia & sudden death without structural heart disease (53 genes)
- 09 Hypertrophic cardiomyopathy: ☐ 16 genes ☐ 90 genes
- 10 Dilated cardiomyopathy (81 genes)
- 11 Restrictive cardiomyopathy (20 genes)
- 12 Left ventricular non-compaction (36 genes)
- 13 Arrhythmogenic cardiomyopathy (17 genes)
- 14 Noonan, Leopard, Costello (12 genes)
- 15 Long QT Syndrome (24 genes)
- 16 Short QT Syndrome (7genes)
- 17 Brugada Syndrome (23 genes)
- 18 J wave Syndrome (23 genes)
- 19 Atrial fibrillation (38 genes)
- 20 Aortic vascular diseases (30 genes)
- 21 Congenital heart diseases (39 genes)
- 22 Skeletal myopathies (46 genes)
- 23 Other _____

CLINICAL INFORMATION (Attach clinical notes and current medication list)

CIRCLE ALL DIAGNOSIS CODES THAT APPLY (USE ADDITIONAL BLANKS IF NEEDED)

Cardio Related
- ❑ **Z82.41** Family History of Cardiac Death
- ❑ **Z84.81** Family History of Carrier of Genetic Disease
- ❑ **Z72.0** Tobacco Use
- ❑ **Z31.5** Encounter for Genetic Counseling
- ❑ **R01.1** Cardiac Murmur Unspecified
- ❑ **E78.9** Hyperlipidemia
- ❑ **I10** Essential Hypertension
- ❑ **R55** Syncope and Collapse
- ❑ **Z01.810** Encounter for Pre-procedural Cardiac Examination
- ❑ **Z31.5** Encounter for Genetic Counseling

Dyslipidemias
- ❑ **E78.5** Hyperlipidemia Unsp.
- ❑ **E78.0** Familial Hypercholesterolemia
- ❑ **E88.81** Metabolic Syndrome
- ❑ **E78.1** Pure Hyperglyceridemia
- ❑ **E78.2** Mixed Hyperlipidemia
- ❑ **124.8** Coronary insufficiency

Cardiopathies
- ❑ **I42.2** Hypertrophic Cardiomyopathy
- ❑ **I42.0** Dilated Cardiomyopathy
- ❑ **I42.5** Restrictive Cardiomyopathy
- ❑ **I34.1** Mitral Valve Prolapse
- ❑ **I71.01** Dissection of Thoracic Aorta
- ❑ **I71.2** Thoracic Aortic Aneurysm, no rupture
- ❑ **Q24.9** Congenital Malformation of Heart, Unspecified
- ❑ **Q87.1** Noonan's Syndrome
- ❑ **Q87.40** Marfan's Syndrome

Arrhythmias
- ❑ **R94.31** Abnormal Electrocardiogram
- ❑ **I45.81** Long QT Syndrome
- ❑ **I49.9** Short QT Syndrome
- ❑ **Q24.8** Brugada Syndrome

Other _____ Other _____ Other _____ Other _____ Other _____

SPECIMEN INFORMATION

❑ Blood (purple top vacutainer) ❑ Saliva

❑ gDNA Concentration _____

SIGNATURE

X _____

Signature over Printed Name

❑ **I confirm documented medical necessity for this test in the patient's file.**

COMMENTS

NOTES

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√	CHECKLIST OF ITEMS TO INCLUDE WITH PATIENT SAMPLE
	Physician's Signature
	Patient's Signature
	Patient Demographics/Insurance Information
	Medications List
	Patient History/Physical
	Patient Office Notes/Progress Notes
	Medical Necessity Has Been Documented in Patient Notes

ABOUT HELICE

Helice Genomic Sciences, Inc. a platform that connects physicians and individuals to global genomics laboratories and provides them a portfolio of the latest genomics-based tests and assays to arm them with powerful and actionable options for navigating the right medical care for the patient's individual needs.

For more information, go to **www.helicegenomics.com** or send us an email at **info@helicegenomics.com**.