

Test Definition: LQTSG

Long QT Syndrome Gene Panel, Varies

Reporting Title: Long QT Syndrome Gene Panel

Performing Location: Mayo Clinic Laboratories - Rochester Main Campus

Ordering Guidance:

This test is intended for genetic screening for and diagnosis of long QT syndrome.

For Brugada syndrome genetic testing, order SCN5A / Brugada Syndrome Multi-Gene Panel, Blood.

For comprehensive inherited cardiac arrhythmia genetic testing, order CARGG / Comprehensive Arrhythmia Gene Panel, Varies.

Customization of this panel and single gene analysis for any gene present on this panel are available. For more information see CGPH / Custom Gene Panel, Hereditary, Next-Generation Sequencing, Varies.

Targeted testing for familial variants (also called site-specific or known mutations testing) is available for the genes on this panel. See FMTT / Familial Variant, Targeted Testing, Varies. To obtain more information about this testing option, call 800-533-1710.

Shipping Instructions:

Specimen preferred to arrive within 96 hours of collection.

Necessary Information:

<u>Prior Authorization</u> is available, **but not required**, for this test. If proceeding with the prior authorization process, submit the required form with the specimen.

Specimen Requirements:

Patient Preparation: A previous bone marrow transplant from an allogenic donor will interfere with testing. Call 800-533-1710 for instructions for testing patients who have received a bone marrow transplant.

Specimen Type: Whole blood

Container/Tube:

Preferred: Lavender top (EDTA) or yellow top (ACD)

Acceptable: Any anticoagulant Specimen Volume: 3 mL Collection Instructions:

1. Invert several times to mix blood

2. Send whole blood specimen in original tube. **Do not aliquot. Specimen Stability Information:** Ambient (preferred)/Refrigerated

Forms:

- . **New York Clients-Informed consent is required.** Document on the request form or electronic order that a copy is on file. The following documents are available:
- -Informed Consent for Genetic Testing (T576)
- -Informed Consent for Genetic Testing (Spanish) (T826)
- 2. Hereditary Cardiomyopathies and Arrhythmias Patient Information
- 3. Long QT Syndrome Gene Panel (LQTSG) Prior Authorization Ordering Instructions



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4. If not ordering electronically, complete, print, and send a <u>Cardiovascular Test Request Form</u> (T724) with the specimen.

| Specimen Type | Temperature | Time | Special Container | |
|---------------|-------------|------|-------------------|--|
| Varies | Varies | | | |

Result Codes:

| Result ID | Reporting Name | Туре | Unit | LOINC® |
|-----------|------------------------|--------------|------|---------|
| 617352 | Test Description | Alphanumeric | | 62364-5 |
| 617353 | Specimen | Alphanumeric | | 31208-2 |
| 617354 | Source | Alphanumeric | | 31208-2 |
| 617355 | Result Summary | Alphanumeric | | 50397-9 |
| 617356 | Result | Alphanumeric | | 82939-0 |
| 617357 | Interpretation | Alphanumeric | | 69047-9 |
| 617358 | Additional Results | Alphanumeric | | 82939-0 |
| 617359 | Resources | Alphanumeric | | 99622-3 |
| 617360 | Additional Information | Alphanumeric | | 48767-8 |
| 617361 | Method | Alphanumeric | | 85069-3 |
| 617362 | Genes Analyzed | Alphanumeric | | 48018-6 |
| 617363 | Disclaimer | Alphanumeric | | 62364-5 |
| 617364 | Released By | Alphanumeric | | 18771-6 |

LOINC® and CPT codes are provided by the performing laboratory.

Supplemental Report:

Supplemental

CPT Code Information:

81403

81406 x2

81407

81479

81479 (if appropriate for government payers)

Reference Values:

An interpretive report will be provided.