Overview

Useful For
Confirming a clinical diagnosis of familial hypercholesterolemia or sitosterolemia

Cascade screening of at-risk family members and early diagnosis, treatment, and dietary modifications

Ascertaining carrier status of family members of individuals diagnosed with familial hypercholesterolemia for genetic counseling purposes

Genetics Test Information
This test includes next-generation sequencing (NGS) and supplemental Sanger sequencing to evaluate for variants in the ABCG5, ABCG8, APOB, LDLR, LDLRAP1, and PCSK9 genes. Additionally, NGS is used to test for the presence of large deletions and duplications in a subset of genes.

Highlights
Identification of a pathogenic variant may assist with prognosis, clinical management, familial screening, and genetic counseling.

Special Instructions
- Informed Consent for Genetic Testing
- Familial/Autosomal Dominant Hypercholesterolemia Patient Information
- Informed Consent for Genetic Testing (Spanish)

Method Name
Custom Sequence Capture and Targeted Next-Generation Sequencing Followed by Polymerase Chain Reaction (PCR) and Supplemental Sanger Sequencing or qPCR

NY State Available
Yes

Specimen

Specimen Type
Varies

Advisory Information
Targeted testing for familial variants (also called site-specific or known mutation testing) is available for the genes on this panel. See:

- KVAR1 / Known Variant Analysis-1 Variant, Varies
- KVAR2 / Known Variant Analysis-2 Variants, Varies
- KVAR3 / Known Variant Analysis-3+ Variants, Varies

Call 800-533-1710 to confirm the appropriate test for targeted testing.

Shipping Instructions
Specimen preferred to arrive within 96 hours of collection.

**Necessary Information**

1. Familial/Autosomal Dominant Hypercholesterolemia Patient Information is required, see Special Instructions. Testing may proceed without the patient information however it aids in providing a more thorough interpretation. Ordering providers are strongly encouraged to complete the form and send it with the specimen.

2. Include physician name and phone number with specimen.

**Specimen Required**

Submit only 1 of the following specimens:

**Specimen Type:** Whole blood

**Container/Tube:** Lavender top (EDTA)

**Specimen Volume:** 3 mL

**Collection Instructions:**

1. Invert several times to mix blood.

2. Send specimen in original tube.

**Specimen Stability Information:** Ambient (preferred)/Refrigerated

**Specimen Type:** DNA

**Container/Tube:** 2 mL screw top tube

**Specimen Volume:** 100 mcL (microliters)

**Collection Instructions:**

1. The preferred volume is 100 mcL at a concentration of 250 ng/mcL.

2. Include concentration and volume on tube.

**Specimen Stability Information:** Frozen (preferred)/Ambient/Refrigerated

**Forms**

1. 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 in Special Instructions:

   - Informed Consent for Genetic Testing (T576)
   - Informed Consent for Genetic Testing-Spanish (T826)

2. If not ordering electronically, complete, print, and send a Cardiovascular Test Request (T724) with the specimen.

**Specimen Minimum Volume**
Familial hypercholesterolemia (FH) is a genetic disorder characterized by elevated levels of low-density lipoprotein cholesterol (LDL-C). FH is associated with increased risk of cardiovascular disease, including premature coronary artery disease and myocardial infarction. Early diagnosis and treatment are crucial in mitigating these risks.

The most common form of FH is autosomal dominant heterozygous familial hypercholesterolemia (heFH), caused by loss-of-function variants in the **LDLR** gene. Loss of LDL-receptors reduces the clearance of LDL from circulation, leading to elevated LDL-C levels. Hundreds of variants in the **LDLR** gene have been identified, most of which are detectable by sequencing. Additional 10% of variants are large intragenic rearrangements, such as deletions and duplications.

A more severe form of FH is homozygous familial hypercholesterolemia (hoFH), caused by homozygous or compound heterozygous (biallelic) variants in the **LDLR** gene. Individuals with hoFH typically have severe hypercholesterolemia, with LDL-C levels exceeding 650 mg/dL, and often develop childhood coronary artery disease, with mortality occurring before the age of 20 due to myocardial infarction.

Another form of FH is familial defective apolipoprotein B-100 (FDB), caused by variants in the **APOB** gene, which reduce the binding affinity between the encoded apolipoprotein B (apoB) and the low-density lipoprotein receptor (LDLR). Individuals with FDB have elevated LDL-C levels, increased rates of coronary artery calcifications, and premature myocardial infarction. The R3500Q variant, prevalent in Northern European Caucasians, and the R3500W variant, prevalent in East Asians, are the most common FDB variants.

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sometimes misdiagnosed with heFH.

Autosomal dominant hypercholesterolemia can also be caused by gain-of-function variants in the PCSK9 gene. Variants in this gene are rare, but when present, they result in increased PCSK9 protein levels, leading to increased degradation of low-density lipoprotein receptors. Recently, drugs targeting PCSK9 (called PCSK9 inhibitors) have been developed. These drugs inhibit the binding of PCSK9 to LDL-receptors, thus reducing degradation of LDL-receptors and increasing the amount of LDL-C cleared in certain individuals.

Loss-of-function variants in the LDLRAP1 gene cause a rare form of familial hypercholesterolemia called autosomal recessive familial hypercholesterolemia. Once LDL-C binds to the LDL-receptor the LDLRAP1 protein binds to the complex and internalization of the complex, which results in degradation of either the LDL particle or the entire complex occurs. Unlike autosomal dominant hypercholesterolemia caused by heterozygous variants in LDLR, APOB, and PCSK9, biallelic variants in LDLRAP1 are required for elevated LDL-C levels. Individuals with homozygous or compound heterozygous LDLRAP1 variants typically have LDL-C levels above 400 mg/dL, cutaneous and tendon xanthomas, and coronary artery disease. Heterozygosity for LDLRAP1 variants does not result in elevated cholesterol levels, so the parents of children with biallelic LDLRAP1 variants are typically normocholesterolemic.

Sitosterolemia, a rare autosomal recessive inherited lipid metabolism disease, is caused by biallelic variants in the ABCG5 or ABCG8 genes and has similar clinical manifestations to familial hypercholesterolemia. Sitosterolemia is characterized by increased intestinal absorption of plant sterols (15% to 60% compared to <5% in unaffected individuals). These individuals also typically have elevated total cholesterol and LDL cholesterol levels, although individuals with normal LDL-C levels have also been reported. Untreated individuals with sitosterolemia exhibit tendon and tuberous xanthomas in childhood, premature atherosclerosis, myocardial infarction, and coronary heart disease. At least one report of an individual with sitosterolemia being misdiagnosed with homozygous FH has been reported. The authors noted that the Dutch Lipid Clinic Network diagnostic (DLCN) criteria could not distinguish between homozygous FH and sitosterolemia in this individual.

Identification of the genetic cause of an individual's clinical features helps to determine the appropriate treatment for their clinical features. Treatment is aimed at lowering plasma LDL levels and plasma sterol levels. Common treatments included statins, LDL apheresis, dietary modifications, and more recently PCSK9 inhibitors. Screening of at-risk family members allows for effective primary prevention by instituting appropriate therapy and dietary modifications at an early stage.

Table 1. Genes included in the Familial Hypercholesterolemia and Related Disorders Multigene Panel

<table>
<thead>
<tr>
<th>Gene Symbol (alias)</th>
<th>Protein</th>
<th>OMIM</th>
<th>Inheritance</th>
<th>Phenotype Disorder</th>
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<tr>
<td>ABCG5</td>
<td>ATP-binding cassette, subfamily G, member 5</td>
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<td>APOB</td>
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**Test Definition: FHRGP**

Hypercholesterolemia Gene Panel

<table>
<thead>
<tr>
<th>Gene</th>
<th>Description</th>
<th>Reference Value</th>
<th>Inheritance</th>
<th>Condition</th>
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<tr>
<td>LDLR</td>
<td>Low density lipoprotein receptor</td>
<td>606945</td>
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<td>LDLRAP1</td>
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AD: autosomal dominant

AR: autosomal recessive

**Reference Values**

An interpretive report will be provided.

**Interpretation**

Evaluation and categorization of variants is performed using the most recent published American College of Medical Genetics and Genomics (ACMG) recommendations as a guideline. Variants are classified based on known, predicted, or possible pathogenicity and reported with interpretive comments detailing their potential or known significance.

Multiple in silico evaluation tools may be used to assist in the interpretation of these results. The accuracy of predictions made by in silico evaluation tools is highly dependent upon the data available for a given gene, and predictions made by these tools may change over time. Results from in silico evaluation tools should be interpreted with caution and professional clinical judgment.

**Cautions**

**Clinical Correlations:**

Some individuals who have involvement of 1 or more of the genes on the panel may have a variation that is not identified by the methods performed (eg, promoter variants, deep intronic variants). The absence of a variant, therefore, does not eliminate the possibility of familial hypercholesterolemia (FH) or a related disorder.

Test results should be interpreted in context of clinical findings, family history, and other laboratory data. Misinterpretation of results may occur if the information provided is inaccurate or incomplete.

If testing was performed because of a family history of FH or a related disorder, it is often useful to first test an affected family member. Identification of a pathogenic variant in an affected individual would allow for more informative testing of at risk individuals.

**Technical Limitations:**

Next-generation sequencing may not detect all types of genetic variants. Additionally, rare polymorphisms may be present that could lead to false-negative or false-positive results. If the patient has had an allogeneic blood or marrow transplant or a recent (ie, <6 weeks from time of sample collection) heterologous blood transfusion, these results may be inaccurate due to the presence of donor DNA.
Reclassification of Variants Policy:

At this time, it is not standard practice for the laboratory to systematically review likely pathogenic variants or variants of uncertain significance that are detected and reported. The laboratory encourages health care providers to contact the laboratory at any time to learn how the status of a particular variant may have changed over time.

Contact the laboratory if additional information is required regarding the transcript or human genome assembly used for the analysis of this patient’s results.

Clinical Reference


Performance

Method Description

Next-generation sequencing (NGS) is performed using an Illumina instrument with paired-end reads. The DNA is prepared for NGS using a custom Agilent SureSelect Target Enrichment System. Data is analyzed with a bioinformatics software pipeline for sequence variants and the presence of large intragenic deletions and duplications. Supplemental Sanger sequencing or qPCR may be performed occasionally in regions where NGS is insufficient for data capture or not specific enough to correctly identify a variant. Sanger sequencing or qPCR may also be used for confirmatory testing. (Unpublished Mayo method)
The following genes are evaluated in this multi-gene panel, \( ABCG5 \), \( ABCG8 \), \( APOB \), \( LDLR \), \( LDLRAP1 \), and \( PCSK9 \).
PDF Report
No

Day(s) and Time(s) Test Performed
Monday; Varies

Analytic Time
2 weeks

Maximum Laboratory Time
4 weeks

Specimen Retention Time
2 months

Performing Laboratory Location
Rochester

Fees and Codes

Fees
- Authorized users can sign in to Test Prices for detailed fee information.
- Clients without access to Test Prices can contact Customer Service 24 hours a day, seven days a week.
- Prospective clients should contact their Regional Manager. For assistance, contact Customer Service.

Test Classification
This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.

CPT Code Information
81479
81406 x 2
81407

LOINC® Information

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