
Reporting Title: Hereditary Hemolytic Anemia, NGS**Performing Location:** Rochester**Ordering Guidance:**

Multiple hematology gene panels are available. For more information see [NHHA and Subpanel Comparison Gene List](#).

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 variants 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.

Additional Testing Requirements:

This test is best interpreted in the context of protein studies and peripheral blood findings. Prior to sending this test, Coombs testing should be negative, laboratory testing should indicate a hemolytic process, and consider evaluating a peripheral blood smear. In addition, protein analysis for hereditary causes of hemolytic anemia can be provided by ordering HAEV1 / Hemolytic Anemia Evaluation Profile, Blood. Fill out the information sheet and indicate that a next-generation sequencing test was also ordered. Additionally, providing complete blood cell count data and clinical notes will allow a more precise interpretation of results.

Shipping Instructions:

Specimen preferred to arrive within 96 hours of collection.

Necessary Information:

[1. Metabolic Hematology Next-Generation Sequencing \(NGS\) Patient Information](#) is required. Testing may proceed without the patient information; however, the information aids in providing a more thorough interpretation. Ordering providers are strongly encouraged to fill out the form and send with the specimen.

2. If form not provided, include the following information with the test request: clinical diagnosis, pertinent clinical history (ie, complete blood cell count results and relevant clinical notes), and differentials based on previous enzyme testing, clinical or morphologic presentation.

Specimen Requirements:

Specimen Type: Whole blood

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.

Container/Tube:

Preferred: Lavender top (EDTA)

Acceptable: Yellow top (ACD)

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) 4 days/Refrigerated

Forms:

1. [Metabolic Hematology Next-Generation Sequencing \(NGS\) Patient Information \(T816\)](#) is required.
2. **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 \(Spanish\) \(T826\)](#)
3. If not ordering electronically, complete, print, and send a [Benign Hematology Test Request \(T755\)](#)with the specimen.

Specimen Type	Temperature	Time	Special Container
Varies	Varies		

Result Codes:

Result ID	Reporting Name	Type	Unit	LOINC®
619034	Test Description	Alphanumeric		62364-5
619035	Specimen	Alphanumeric		31208-2
619036	Source	Alphanumeric		31208-2
619037	Result Summary	Alphanumeric		50397-9
619038	Result	Alphanumeric		82939-0
619039	Interpretation	Alphanumeric		59465-5
619040	Additional Results	Alphanumeric		82939-0
619041	Resources	Alphanumeric		99622-3
619042	Additional Information	Alphanumeric		48767-8
619043	Method	Alphanumeric		85069-3
619044	Genes Analyzed	Alphanumeric		82939-0
619045	Disclaimer	Alphanumeric		62364-5
619046	Released By	Alphanumeric		18771-6

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

Supplemental Report:

Supplemental

CPT Code Information:

81443

Reference Values:

An interpretive report will be provided.