

# CONCERT GENETIC TESTING: HEMATOLOGIC CONDITIONS (NON-CANCEROUS)

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[Coding implications](#)  
[Revision Log](#)

**See Important Reminder at the end of this policy for important regulatory and legal information.**

## OVERVIEW

Genetic testing for hematologic (non-cancerous) conditions may be used to confirm a diagnosis in a patient who has signs and/or symptoms of a specific hematologic condition. Confirming the diagnosis may alter aspects of management and may eliminate the need for further diagnostic workup. This document addresses genetic testing for common hematologic (non-cancerous) conditions.

## POLICY REFERENCE TABLE

### Coding Implications

This clinical policy references Current Procedural Terminology (CPT®). CPT® is a registered trademark of the American Medical Association. All CPT codes and descriptions are copyrighted 2022, American Medical Association. All rights reserved. CPT codes and CPT descriptions are from the current manuals and those included herein are not intended to be all-inclusive and are included for informational purposes only. Codes referenced in this clinical policy are for informational purposes only and may not support medical necessity. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

NOTE: Coverage is subject to each requested code's inclusion on the corresponding LDH fee schedule. Non-covered codes are denoted (\*) and are reviewed for Medical Necessity for members under 21 years of age on a per case basis.

. Please see the [Concert Genetics Platform](#) for a comprehensive list of registered tests.

<a href="#">Criteria Sections</a>	<b>Example Tests (Labs)</b>	<b>Common CPT Codes</b>	<b>Common ICD Codes</b>	<a href="#">Ref</a>
<b><a href="#">Known Familial Variant Analysis for Hematologic Conditions (non-cancerous)</a></b>				
<a href="#">Known Familial Variant Analysis for Hematologic Conditions (non-cancerous)</a>	Targeted Mutation Analysis for a Known Familial Variant	81403*, 81258*, 81362*		11
<b><a href="#">Inherited Thrombophilia</a></b>				
<a href="#">Factor V Leiden (F5) and Prothrombin (F2) Variant Analysis for Inherited Thrombophilia</a>	Factor V (Leiden) Mutation Analysis (Quest Diagnostics)	81241	D68.51, D68.2, D68.59, R79.1, Z86.2, I82.90	1, 5
	Prothrombin (Factor II) 20210G>A Mutation Analysis (Quest Diagnostics)	81240*	D68.52, D68.2, D68.59, R79.1, Z86.2, I82.90	
<b><a href="#">Hemoglobinopathies</a></b>				
<a href="#">HBA1/HBA2 and/or HBB Variant Analysis</a>	HBA1 Deletion/Duplication Analysis (GeneDx) HBA2 Deletion/Duplication Analysis (GeneDx) HBA1 Single Gene (Sequencing Only) (Fulgent Genetics) HBA2 Gene Sequencing (Fairview Diagnostic Laboratories)	81257*, 81259*, 81269*, S3845*, S3850*	D56.0, D56.9, D53.9, R70.1, D56.3, D56.8, Z86.2	2, 3, 4, 6
	HBB Sequencing Analysis (ARUP Laboratories) HBB Deletion/Duplication Analysis (GeneDx)	81361*, 81363*, 81364*, S3846*	D57.00- D57.819, D56.1, D64.9	
<b><a href="#">Hemophilia</a></b>				
<a href="#">F8 and/or F9 Variant Analysis</a>	F8 Deletion/Duplication Analysis (GeneDx) F8 Single Gene (Sequencing Only) (Fulgent Genetics)	81403*, 81406*, 81407*	D66, D67, I62.9, M25, N92.2, R04.0, R31	8, 9
	F9 Full Gene Sequencing and Deletion/Duplication (Invitae)	81238*, 81479		
<b><a href="#">Glucose-6-Phosphate Dehydrogenase (G6PD) Deficiency</a></b>				
<a href="#">G6PD Variant Analysis</a>	G6PD Targeted Variant - Single Test (GeneDx)	81247*, 81248*, 81249*	D55.0	7

	Carrier - G6PD Full Gene Sequencing and Deletion/Duplication (Invitae)			
<b><u>von Willebrand Disease</u></b>				
<a href="#"><u>GP1BA and/or VWF Variant Analysis</u></a>	GP1BA Gene Sequencing & Deletion/Duplication (Fairview Diagnostic Laboratories) VWF Targeted Variant - Single Test (GeneDx) Von Willebrand Disease (VWF) Sequencing (ARUP Laboratories)	81401*, 81403*, 81404*, 81405*, 81406*, 81408*, 81479	D68.0	10
<b><u>Other Covered Hematologic Conditions (non-cancerous)</u></b>				
<a href="#"><u>Other Covered Hematologic Conditions (non-cancerous)</u></a>	See list below	81400*-81408*		12, 13, 14

## OTHER RELATED POLICIES

This policy document provides criteria for Genetic Testing for Hematologic Conditions (Non-Cancerous). Please refer to:

- ***Oncology: Molecular Analysis of Solid Tumors and Hematologic Malignancies*** for criteria related to exome and genome sequencing of solid tumors and hematologic malignancies.
- ***Genetic Testing: Prenatal and Preconception Carrier Screening*** for criteria related to carrier screening in the prenatal, preimplantation, and preconception setting.
- ***Genetic Testing: Prenatal Diagnosis (via amniocentesis, CVS, or PUBS) and Pregnancy Loss*** for criteria related to prenatal and pregnancy loss diagnostic genetic testing for tests intended to diagnose genetic conditions following amniocentesis, chorionic villus sampling or pregnancy loss.
- ***Genetic Testing: Multisystem Inherited Disorders, Intellectual Disability, and Developmental Delay*** for criteria related to diagnostic genetic testing for conditions affecting multiple organ systems.
- ***Genetic Testing: Metabolic, Endocrine, and Mitochondrial Disorders*** for criteria related to genetic testing for *MTHFR*.
- ***Genetic Testing: General Approach to Genetic and Molecular Testing*** for criteria related to genetic testing for non-cancerous hematologic disorders that are not specifically discussed in this or another non-general policy.

## CRITERIA

It is the policy of Louisiana Healthcare Connections that the specific genetic testing noted below is **medically necessary** when meeting the related criteria:

### KNOWN FAMILIAL VARIANT ANALYSIS FOR HEMATOLOGIC CONDITIONS (NON-CANCEROUS)

- I. Targeted mutation analysis for a known familial variant (81403\*, 81258\*, 81362\*) for a non-cancerous hematologic condition is considered **medically necessary** when:
  - A. The member/enrollee has a [close relative](#) with a known pathogenic or likely pathogenic variant causing the condition.
- II. Targeted mutation analysis for a known familial variant (81403\*, 81258\*, 81362\*) for a non-cancerous hematologic condition is considered **investigational** for all other indications.

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## INHERITED THROMBOPHILIA

### Factor V Leiden (*F5*) and Prothrombin (*F2*) Variant Analysis for Inherited Thrombophilia

- I. *F5* (81241) and *F2* (81240\*) variant analysis to confirm or establish a diagnosis of an inherited thrombophilia may be considered **medically necessary** when:
  - A. The member/enrollee meets at least one of the following:
    1. A first unprovoked venous thromboembolism (VTE) younger than 50 years old, **OR**
    2. VTE at unusual sites (such as hepatic portal, mesenteric, and cerebral veins), **OR**
    3. Recurrent VTE, **OR**
    4. Personal history of VTE with at least one of the following:
      - a) Two or more family members with a history of VTE, **OR**
      - b) One [first-degree relative](#) with VTE at a young age, **OR**

5. Low activated protein C (APC) resistance activity, **OR**
  6. The member/enrollee is a female under the age of 50 who smokes tobacco and has a history of acute myocardial infarction, **OR**
  7. The member/enrollee has a [first-degree relative](#) known to be homozygous for factor V Leiden or factor II c.\*97G>A, **OR**
  8. The member/enrollee is an asymptomatic pregnant female or female contemplating pregnancy, with a [first-degree relative](#) with unprovoked VTE or VTE provoked by pregnancy or contraceptive use, **OR**
  9. The member/enrollee is a pregnant female or female contemplating pregnancy or estrogen use who has a [first-degree relative](#) with both of the following:
    - a) A history of VTE, **AND**
    - b) The member/enrollee is a known carrier for factor V Leiden and/or factor II c.97\*G>A variant, **OR**
  10. The member/enrollee is a pregnant female or female contemplating pregnancy with a previous non-estrogen-related VTE or VTE provoked by a minor risk factor.
- II. *F5* (81241) and *F2* (81240\*) variant analysis to confirm or establish a diagnosis of an inherited thrombophilia is considered **investigational** for all other indications, including:
- A. Fetal loss or adverse pregnancy outcomes (examples: placental abruption, fetal growth restriction, or preeclampsia).

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## HEMOGLOBINOPATHIES

### *HBA1/HBA2* and/or *HBB* Variant Analysis

- I. *HBA1/HBA2* variant analysis (81257\*, 81259\*, 81269\*, S3845\*, S3850\*), and/or *HBB* variant analysis (81361\*, 81363\*, 81364\*, S3846\*) to confirm or establish a diagnosis of a hemoglobinopathy (alpha-thalassemia, beta-thalassemia, or sickle cell disease) is considered **medically necessary** when:
  - A. The member/enrollee's hematologic screening results (examples: MCV, MCH, CBC, hemoglobin electrophoresis, or dichlorophenol indophenol (DCIP)) are positive for a hemoglobinopathy, **OR**

- B. The member/enrollee's hematologic screening results (examples: MCV, MCH, CBC, hemoglobin electrophoresis, or dichlorophenol indophenol (DCIP)) do not conclusively diagnose or rule out a hemoglobinopathy.
- II. *HBA1/HBA2* variant analysis (81257\*, 81259\*, 81269\*, S3845\*, S3850\*), and/or *HBB* variant analysis (81361\*, 81363\*, 81364\*, S3846\*) to confirm or establish a diagnosis of a hemoglobinopathy (alpha-thalassemia, beta-thalassemia, or sickle cell disease) is considered **investigational** for all other indications.

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## HEMOPHILIA

### *F8* and/or *F9* Variant Analysis

- I. *F8* variant analysis (81403\*, 81406\*, 81407\*) and/or *F9* variant analysis (81238\*, 81479) to confirm or establish a diagnosis of hemophilia A or B is considered **medically necessary** when:
- A. The member/enrollee has any of the following clinical features of hemophilia:
1. Hemarthrosis (especially with mild or no antecedent trauma), **OR**
  2. Deep-muscle hematomas, **OR**
  3. Intracranial bleeding in the absence of major trauma, **OR**
  4. Neonatal cephalohematoma or intracranial bleeding, **OR**
  5. Prolonged oozing or renewed bleeding after initial bleeding stops following tooth extractions, mouth injury, or circumcision, **OR**
  6. Prolonged, delayed bleeding, or poor wound healing following surgery or trauma, **OR**
  7. Unexplained GI bleeding or hematuria, **OR**
  8. Heavy or prolonged menstrual bleeding (especially with onset at menarche), **OR**
  9. Prolonged nosebleeds, especially recurrent and bilateral, **OR**
  10. Excessive bruising (especially with firm, subcutaneous hematomas), **OR**
- B. The following laboratory features:
1. Normal platelet count, **AND**

2. Prolonged activated partial thromboplastin time (aPTT), **AND**
  3. Normal prothrombin time (PT).
- C. *F8* variant analysis (81403\*, 81406\*, 81407\*) and/or *F9* variant analysis (81238\*, 81479) to confirm or establish a diagnosis of hemophilia A or B is considered **investigational** for all other indications.

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## GLUCOSE-6-PHOSPHATE DEHYDROGENASE (G6PD) DEFICIENCY

### *G6PD* Variant Analysis

1. *G6PD* variant analysis (81247\*, 81248\*, 81249\*) to confirm or establish a diagnosis\* of glucose-6-phosphate dehydrogenase deficiency is considered **investigational**.

\* Diagnosis of *G6PD* can be achieved by quantitative spectrophotometric analysis or, more commonly, by a rapid fluorescent spot test detecting the generation of NADPH from NADP.

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## VON-WILLEBRAND DISEASE

### *GP1BA* and/or *VWF* Variant Analysis

1. *GP1BA* and/or *VWF* variant analysis (81401\*, 81403\*, 81404\*, 81405\*, 81406\*, 81408\*, 81479) to confirm or establish a diagnosis\* of von-Willebrand disease is considered **investigational**.

\* Diagnosis of von-Willebrand disease can be achieved by standard laboratory and biochemical testing.

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## OTHER COVERED HEMATOLOGIC CONDITIONS (NON-CANCEROUS)

The following is a list of conditions that have a known genetic association. Due to their relative rareness, it may be appropriate to cover these genetic tests to establish or confirm a diagnosis.

1. Genetic testing to establish or confirm one of the following hematologic conditions (non-cancerous) to guide management is considered **medically necessary** when the

member/enrollee demonstrates clinical features\*\* consistent with the disorder (the list is not meant to be comprehensive, see II below):

- A. [Atypical Hemolytic-Uremic Syndrome \(aHUS\)](#)
  - B. [Complete Plasminogen Activator Inhibitor 1 Deficiency \(PAI-1\)](#)
  - C. [Diamond-Blackfan Anemia \(DBA\)](#)
  - D. [Hereditary Spherocytosis](#)
  - E. Factor VII Deficiency
  - F. Factor X Deficiency
  - G. Factor XI Deficiency (Hemophilia C)
  - H. Factor XII Deficiency
  - I. Factor XIII Deficiency
- II. Genetic testing to establish or confirm the diagnosis of all other non-cancerous hematologic conditions not specifically discussed within this or another medical policy will be evaluated by the criteria outlined in *General Approach to Genetic and Molecular Testing* (see policy for criteria).

\*\*Clinical features for a specific disorder may be outlined in resources such as [GeneReviews](#), [OMIM](#), [National Library of Medicine](#), [Genetics Home Reference](#), or other scholarly source.

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## NOTES AND DEFINITIONS

1. **Close relatives** include first, second, and third degree blood relatives on the same side of the family:
  - a. **First-degree relatives** are parents, siblings, and children
  - b. **Second-degree relatives** are grandparents, aunts, uncles, nieces, nephews, grandchildren, and half siblings
  - c. **Third-degree relatives** are great grandparents, great aunts, great uncles, great grandchildren, and first cousins

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## BACKGROUND AND RATIONALE

### Known Familial Variant Analysis for Hematologic Conditions (non-cancerous)

*Genetic Support Foundation*



The Genetic Support Foundation's Genetics 101 information on genetic testing says the following about testing for familial pathogenic variants:

Genetic testing for someone who may be at risk for an inherited disease is always easier if we know the specific genetic cause. Oftentimes, the best way to find the genetic cause is to start by testing someone in the family who is known or strongly suspected to have the disease. If their testing is positive, then we can say that we have found the familial pathogenic (harmful) variant. We can use this as a marker to test other members of the family to see who is also at risk.

### **Factor V Leiden (F5) and Prothrombin (F2) Variant Analysis for Inherited Thrombophilia**

*American College of Medical Genetics and Genomics (ACMG)*

The American College of Medical Genetics and Genomics (Zhang, 2018) published updated technical standards for genetic testing for variants associated with VTE, with a focus on factor V Leiden and factor II. Testing is recommended for factor V Leiden and factor II c.\*97G>A for the following indications:

- 1.) A first unprovoked VTE, especially <50 years old
- 2.) VTE at unusual sites (such as hepatic portal, mesenteric, and cerebral veins)
- 3.) Recurrent VTE
- 4.) Personal history of VTE with (a) two or more family members with a history of VTE or (b) one first-degree relative with VTE at a young age
- 5.) Patients with low activated protein C (APC) resistance activity (p. 1492)

In addition, this testing “may be considered” for the following indications:

- 1.) Females under the age of 50 who smoke tobacco and have a history of acute myocardial infarction
- 2.) Siblings of individuals known to be homozygous for factor V Leiden or factor II c.\*97G>A, because they have a 1 in 4 chance of being a homozygote
- 3.) Asymptomatic pregnant female or female contemplating pregnancy, with a first-degree relative with unprovoked VTE or VTE provoked by pregnancy or contraceptive use
- 4.) Pregnant female or female contemplating pregnancy or estrogen use who has a first-degree relative with a history of VTE and is a known carrier for factor V Leiden and/or factor II c.\*97G>A variant
- 5.) Pregnant female or female contemplating pregnancy with a previous non-estrogen-related VTE or VTE provoked by a minor risk factor, because knowledge of the factor V Leiden or factor II c.\*97G>A status may alter pregnancy-related thrombophylaxis (p. 1492-1493)

*American College of Obstetricians and Gynecologists (ACOG)*

ACOG also published Practice Bulletin 197 (2018) on Inherited Thrombophilias in Pregnancy which states that “...screening for inherited thrombophilias is not recommended for women with a

history of fetal loss or adverse pregnancy outcomes including abruption, preeclampsia, or fetal growth restriction because there is insufficient clinical evidence that antepartum prophylaxis with unfractionated heparin or low-molecular-weight-heparin prevents recurrence in these patients, and a causal association has not been established.” (p. e23).

### **Hemoglobinopathies - *HBA1/HBA2* and/or *HBB* Variant Analysis**

GeneReviews is an expert-authored review of current literature on a genetic disease, and goes through a rigorous editing and peer review process before being published online. The recommended hemoglobinopathy evaluation testing for Alpha-Thalassemia, Beta-Thalassemia, and Sickle Cell Disease is as follows:

#### *GeneReviews: Alpha-Thalassemia*

Hemoglobin Bart hydrops fetalis (Hb Bart) syndrome, which is caused by deletion or inactivation of all four alpha globin genes, exhibits the following hematologic findings: severe macrocytic hypochromic anemia (in the absence of ABO or Rh blood group incompatibility), reticulocytosis (may be >60%), and peripheral blood smear with large, hypochromic red cells, severe anisopoikilocytosis, and numerous nucleated red cells. In addition, hemoglobin analysis will typically display decreased amounts or complete absence of hemoglobin A and increased amounts of Hb Bart.

Hemoglobin H disease (HbH disease), which is caused by deletion or inactivation of three alpha globin genes, exhibits the following hematologic findings: mild-to-moderate (rarely severe) microcytic hypochromic hemolytic anemia, moderate reticulocytosis (3%-6%), Peripheral blood smear with anisopoikilocytosis, and very rarely nucleated red blood cells, Red blood cell supravital stain showing HbH inclusions ( $\beta_4$  tetramers) in 5%-80% of erythrocytes following incubation of fresh blood smears with 1% brilliant cresyl blue for one to three hours. In addition, hemoglobin analysis will typically display the presence of 0.8%-40% HbH and 60%-90% hemoglobin A.

#### *GeneReviews: Beta-Thalassemia*

Beta-Thalassemia typically displays the following hematologic findings of microcytic hypochromic anemia, anisopoikilocytosis with nucleated red blood cells on peripheral blood smear, and hemoglobin analysis that reveals decreased amounts or complete absence of hemoglobin A and increased amounts of hemoglobin F.

#### *GeneReviews: Sickle Cell Disease*

Laboratory features of sickle cell disease include: normocytic anemia; sickle cells, nucleated red blood cells, target cells, and other abnormal red blood cells on peripheral blood smear; Howell-Jolly bodies indicate hyposplenism; presence of hemoglobin S (HbS) on a hemoglobin assay (e.g., high-performance liquid chromatography [HPLC], isoelectric focusing, cellulose acetate electrophoresis, citrate agar electrophoresis) with an absence or diminished amount of HbA.

*Viprakasit V, Ekwattanakit S. Clinical classification, screening and diagnosis for thalassemia*

Viprakasit and Ekwattanakit (2018) published a clinical classification, screening and diagnosis for thalassemia article that states:

“In general, these mutation analyses would be critical for the confirmation of thalassemia diagnoses in only a few selected cases for whom the basic hematology and Hb analysis described could not provide a conclusive diagnosis. However, these molecular analyses would be indispensable in a program for the prevention and control of thalassemia syndromes because the mutation data would be required for genetic counseling, genetic risk calculation in the offspring, and prenatal and preimplantation genetic diagnosis. In addition, DNA analysis could help in predicting the clinical severity and guiding clinical management; milder b-globin mutations (b1-thal) usually are associated with milder phenotypes, as has been shown in HbE/b-thalassemia.” (p. 207)

### **Hemophilia - F8 and/or F9 Variant Analysis**

GeneReviews is an expert-authored review of current literature on a genetic disease, and goes through a rigorous editing and peer review process before being published online. The recommended hemoglobinopathy evaluation testing for Hemophilia A and Hemophilia B is as follows:

*GeneReviews: Hemophilia A and Hemophilia B*

Individuals with Hemophilia A (factor VIII deficiency) or Hemophilia B (factor IX deficiency) can exhibit the following clinical symptoms:

- Hemarthrosis, especially with mild or no antecedent trauma
- Deep-muscle hematomas
- Intracranial bleeding in the absence of major trauma
- Neonatal cephalohematoma or intracranial bleeding
- Prolonged oozing or renewed bleeding after initial bleeding stops following tooth extractions, mouth injury, or circumcision
- Prolonged or delayed bleeding or poor wound healing following surgery or trauma
- Unexplained GI bleeding or hematuria
- Menorrhagia, especially with onset at menarche
- Prolonged nosebleeds, especially recurrent and bilateral
- Excessive bruising, especially with firm, subcutaneous hematomas

The following are laboratory findings in individuals with Hemophilia A or Hemophilia B:

- Normal platelet count
- Prolonged activated partial thromboplastin time (aPTT)
- Normal prothrombin time (PT)

## Glucose-6-Phosphate Dehydrogenase Deficiency - G6PD Variant Analysis

*American Academy of Family Physicians*

Frank (2005) published guidelines in American Family Physician for evaluating individuals for G6PD deficiency, including specific laboratory tests which notably do not include genetic testing: “The diagnosis of G6PD deficiency is made by a quantitative spectrophotometric analysis or, more commonly, by a rapid fluorescent spot test detecting the generation of NADPH from NADP. The test is positive if the blood spot fails to fluoresce under ultraviolet light.” (p. 1278)”

## von Willebrand Disease - GPIBA and/or VWF Variant Analysis

*Centers for Disease Prevention and Control (CDC), via the National Heart Lung and Blood Institute, National Institutes of Health (NHLBI-NIH)*

Guidelines for diagnosis and management of von Willebrand disease (VWD) were developed for practicing primary care and specialist clinicians—including family physicians, internists, obstetrician-gynecologists, pediatricians, and nurse-practitioners—as well as hematologists and laboratory medicine specialists, which included recommendations for laboratory tests to aid in the diagnosis of VWD, which notably do not include genetic testing.

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Reviews, Revisions, and Approvals	Revision Date	Approval Date
Converted corporate to local policy.	09/23	11/27/23
Semi-annual review. Overview, coding, reference-table, background and references updated. Throughout policy: replaced “coverage criteria” with “criteria. For Policy Reference Table; under Hemoglobinopathies: added “(GeneDx)”; and added “HBA1 Single Gene...”; added “(ARUP Laboratories)”; under Hemophilia: removed “F8 Sequencing Analysis” and added “(GeneDx)...”; removed “Deletion/Duplication Analysis...”; added “Full Gene Sequencing...”; under Glucose-6-Phosphate Dehydrogenase...” removed “Mutation Analysis...”; and added “Variant-Single Test...”; under von Willebrand Disease: removed “Sequencing Analysis...” and added “Gene Sequencing...”. For Other Related Policies: added “and Molecular”. For Criteria; under Hemoglobinopathies: added “variant analysis”; under Von-Willebrand Disease: added “/or”; under Glucose-6-Phosphate Dehydrogenase (G6PD) Deficiency added “and Molecular”. For Background and Rationale; under Known Familial Variants: added “and Molecular”. For Background and Rationale; under Known Familial Variants: changed “inheritance patterns” to “genetic testing”.	12/23	2/27/24

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### **Important Reminder**

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. LHCC makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable LHCC administrative policies and procedures.

This clinical policy is effective as of the date determined by LHCC. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. LHCC retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members/enrollees. This clinical policy is not intended to recommend treatment for members/enrollees. Members/enrollees should consult with their treating physician in connection with diagnosis and treatment decisions.

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