

**THIS IS NOT A TEST REQUEST FORM.**

**The information below is required to perform hemoglobinopathy/thalassemia testing.  
Please fill out this form and submit it with the test request form or electronic packing list.**

**PATIENT HISTORY FOR HEMOGLOBINOPATHY/THALASSEMIA TESTING**

**Patient Name** \_\_\_\_\_ **Date of Birth** \_\_\_\_/\_\_\_\_/\_\_\_\_ **Gender**  F  M

**Physician** \_\_\_\_\_ **Physician Phone** (\_\_\_\_\_) \_\_\_\_\_ **Practice Specialty** \_\_\_\_\_

**Genetic Counselor** \_\_\_\_\_ **Counselor Phone** (\_\_\_\_\_) \_\_\_\_\_

**PATIENT'S ETHNICITY** (check all that apply)

- |   |   |   |
|---|---|---|
| <input type="checkbox"/> African American               | <input type="checkbox"/> Caucasian (N Europe) | <input type="checkbox"/> Middle Eastern |
| <input type="checkbox"/> African (specify region _____) | <input type="checkbox"/> Caucasian (S Europe) | <input type="checkbox"/> Puerto Rican   |
| <input type="checkbox"/> Asian Indian                   | <input type="checkbox"/> Chinese              | <input type="checkbox"/> Vietnamese     |
| <input type="checkbox"/> Asian Southeast                | <input type="checkbox"/> Hispanic             | <input type="checkbox"/> Other _____    |

**Does the patient have SYMPTOMS?**  No  Yes (check all that apply)

- |  |
|--|
| <input type="checkbox"/> Anemia (has iron deficiency been excluded? <input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> Unknown) |
| <input type="checkbox"/> Splenomegaly <input type="checkbox"/> Other _____   |

**Has the patient had a recent TRANSFUSION?**  No  Yes (date of transfusion: \_\_\_\_\_)  Unknown

**LABORATORY FINDINGS**

Hemoglobin evaluation by electrophoresis or HPLC (date performed : \_\_\_\_\_)

Hb A% \_\_\_\_\_ Hb A<sub>2</sub>% \_\_\_\_\_ Hb F% \_\_\_\_\_ Hb S% \_\_\_\_\_ Hb C% \_\_\_\_\_ Hb E% \_\_\_\_\_ Other \_\_\_\_\_  
 HGB: \_\_\_\_\_ HCT: \_\_\_\_\_ MCV: \_\_\_\_\_ Reticulocyte count: \_\_\_\_\_(\_\_\_\_\_%)

**FAMILY HISTORY OF HEMOGLOBINOPATHY/ THALASSEMIA?**  No  Yes  Unknown

If yes, what is the **RELATIONSHIP** of family member(s) to the patient? \_\_\_\_\_

Is the relative?  a healthy carrier  affected

List the **GENE and MUTATION(S)** identified or include a copy of the laboratory result: \_\_\_\_\_

**HAS DNA TESTING BEEN PERFORMED PREVIOUSLY FOR THIS PATIENT?**  No  Yes  Unknown

If yes, please check the completed test(s) and provide result or attach report.

- |   |               |
|---|---------------|
| <input type="checkbox"/> Alpha globin deletion analysis | Result: _____ |
| <input type="checkbox"/> Beta globin sequencing         | Result: _____ |
| <input type="checkbox"/> Other                          | Result: _____ |

**Circle the test you intend to order.**

<b>Initial screening tests for hemoglobinopathies/thalassemia:</b>	
<b>0050610</b>	<b>Hemoglobin Evaluation with Reflex to Electrophoresis and/or RBC Solubility – HPLC with reflex to electrophoresis and/or RBC solubility</b>
<b>2005792</b>	<b>Hemoglobin Evaluation Reflexive Cascade – HPLC with reflex to electrophoresis, solubility testing, or molecular analyses to identify Hb variants</b>
<b>Molecular tests for beta thalassemia/ hemoglobinopathies:</b>	
<b>2010117</b>	<b>Beta Globin (HBB) Sequencing and Deletion/Duplication – Clinical sensitivity for beta thalassemia ~99%.</b>
<b>0050578</b>	<b>Beta Globin (HBB) Sequencing- Clinical sensitivity for beta thalassemia ~97%.</b>
<b>2010113</b>	<b>Beta Globin (HBB) Deletion/Duplication- Clinical sensitivity varies by ethnicity.</b>
<b>2004686</b>	<b>Hemoglobin Lepore (HBD/HBB Fusion) 3 Mutations</b>
<b>Molecular tests for alpha thalassemia:</b>	
<b>2011708</b>	<b>Alpha Globin (HBA1 and HBA2) Sequencing and Deletion/Duplication- Clinical sensitivity is 99%</b>
<b>2011622</b>	<b>Alpha Globin (HBA1 and HBA2) Deletion/Duplication- Clinical sensitivity up to 95% Assesses for common, rare and novel deletions and duplications.</b>
<b>0051495</b>	<b>Alpha Thalassemia (HBA1 &amp; HBA2) 7 Deletions – Clinical sensitivity up to 90%. Assesses for 7 common large deletions.</b>
<b>2001582</b>	<b>Alpha Thalassemia (HBA1 &amp; HBA2) Sequencing – Clinical sensitivity is ~10%.</b>

**For questions, contact a genetic counselor at (800) 242-2787, ext. 2141.**

Master Label