



Instructions: The accurate interpretation and reporting of genetic results is contingent upon the reason for testing, ancestry, family history, and clinical history. To help provide the best possible service, supply the information requested below and **send this paperwork with the specimen, or return by fax to Mayo Clinic Laboratories, Attn: Molecular Hematopathology Genetics Lab Counselors at 507-284-1759. Phone: 800-533-1710 / International clients: 855-379-3115 or +1-507-284-9273, or email mliintl@mayo.edu**

Patient Information (required)

Patient Name (Last, First Middle)		Birth Date (mm-dd-yyyy)
Sex Assigned at Birth <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Unknown <input type="checkbox"/> Choose not to disclose	Legal/Administrative Sex <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Nonbinary	

Referring Healthcare Professional Information

Requesting Healthcare Professional Name (Last, First)	Phone	Fax*
Healthcare Professional Email		
Genetic Counselor Name (Last, First)	Phone	Fax*

**Fax number given must be from a fax machine that complies with applicable HIPAA regulations.*

Reason for Testing

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Ancestry

<input type="checkbox"/> African/African American	<input type="checkbox"/> East Asian	<input type="checkbox"/> Latinx/Latine	<input type="checkbox"/> South Asian	<input type="checkbox"/> Unknown
<input type="checkbox"/> Ashkenazi Jewish	<input type="checkbox"/> European	<input type="checkbox"/> Middle Eastern	<input type="checkbox"/> None of the above	<input type="checkbox"/> Choose not to disclose
<input type="checkbox"/> Other, specify: _____		<input type="checkbox"/> Indicate countries of origin: _____		

Family History

Are other relatives known to be affected? <input type="checkbox"/> Yes <input type="checkbox"/> No If "Yes," describe symptoms and relationship to patient:
<hr/> <hr/>
If a relative was tested at Mayo Clinic, family member name (Last, First Middle) _____

Clinical History

Reason for Testing		
<input type="checkbox"/> PK enzyme level: _____	<input type="checkbox"/> Neonatal anemia	<input type="checkbox"/> Known previous diagnosis: _____
<input type="checkbox"/> Chronic anemia	<input type="checkbox"/> Hyperbilirubinemia/Jaundice	
<input type="checkbox"/> Carrier testing	<input type="checkbox"/> Iron overload: _____	Previously tested at Mayo Clinic: <input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> Pigmented gallstones	<input type="checkbox"/> Other: _____	
RBC _____ HGB _____ HCT _____ MCV _____ RDW _____ MCH _____ Ferritin _____		
<input type="checkbox"/> Reticulocyte count _____ p50 _____	Coombs test: <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Not done Therapy _____	

Calcium Sensing Receptor (CASR) Gene Testing Patient Information (continued)

Clinical History (continued)

Patient recently transfused: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown
Last transfusion(s) date(s) (mm-dd-yyyy): _____
<input type="checkbox"/> Splenomegaly Splenectomy: <input type="checkbox"/> Yes <input type="checkbox"/> No Date (mm-dd-yyyy): _____
<input type="checkbox"/> Phototherapy <input type="checkbox"/> No therapy
Other Relevant Clinical Information

Peripheral blood smear shows:

