

**THIS IS NOT A TEST REQUEST FORM.**  
Please fill out this form and submit it with the test request form or electronic packing list.

**PATIENT HISTORY FOR G6PD DEFICIENCY TESTING**

Patient Name \_\_\_\_\_ Date of Birth \_\_\_\_\_ Sex  F  M  
 Physician \_\_\_\_\_ Physician Phone \_\_\_\_\_  
 Practice Specialty \_\_\_\_\_ Physician Fax \_\_\_\_\_  
 Genetic Counselor \_\_\_\_\_ Counselor Phone \_\_\_\_\_

**Patient's Ethnicity** (check all that apply)

<input type="checkbox"/> African-American	<input type="checkbox"/> Asian Indian	<input type="checkbox"/> Filipino	<input type="checkbox"/> Taiwanese
<input type="checkbox"/> African (specify region): _____	<input type="checkbox"/> Caucasian (Mediterranean)	<input type="checkbox"/> Hispanic	<input type="checkbox"/> Thai
	<input type="checkbox"/> Caucasian (N Europe)	<input type="checkbox"/> Middle Eastern	<input type="checkbox"/> Other: _____
	<input type="checkbox"/> Chinese	<input type="checkbox"/> Papua New Guinea	

Did the patient have an **abnormal newborn screen** for G6PD deficiency?  No  Yes  
 If yes, describe result: \_\_\_\_\_

Does the patient have **symptoms**?  No  Yes (check all that apply and describe)

<input type="checkbox"/> Acute hemolytic anemia after exposure to oxidative stress (infection/certain medications/fava beans)	<input type="checkbox"/> Splenomegaly
<input type="checkbox"/> Chronic non-spherocytic hemolytic anemia	<input type="checkbox"/> Cholelithiasis
<input type="checkbox"/> Jaundice or hyperbilirubinemia	<input type="checkbox"/> Other Symptom: _____

Has the patient undergone previous DNA testing for G6PD deficiency?  No  Yes  Unknown  
 If yes, describe the test(s) and results: \_\_\_\_\_

**Laboratory Findings**

G6PD quantitative enzyme level:  Normal  Abnormal  Unknown  Not Performed  
 Rapid fluorescent spot test:  Normal  Abnormal  Unknown  Not Performed  
 Other: \_\_\_\_\_  Normal  Abnormal

Is there any relevant **family history**?  No  Yes  Unknown  
 If yes, attach a pedigree or specify the relative's relationship to the patient. List their symptoms and age at diagnosis:  
 \_\_\_\_\_  
 \_\_\_\_\_

Has DNA testing been performed for the family member(s)?  No  Yes  Unknown  
 If yes, attach a copy of the relative's DNA laboratory result (REQUIRED for familial mutation testing).

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

- 0080135 Glucose-6-Phosphate Dehydrogenase:** Quantitative enzymatic test. Preferred initial test to confirm a diagnosis of G6PD deficiency.
- 2007163 Glucose-6-Phosphate Dehydrogenase (G6PD) Sequencing:** Clinical sensitivity for G6PD deficiency is expected to be greater than 98% for all ethnicities.
- 0051684 Glucose-6-Phosphate Dehydrogenase (G6PD) 2 Mutations:** Clinical sensitivity in African-Americans is 99%. Sensitivity in other ethnicities is unknown.
- 2001961 Familial Mutation, Targeted Sequencing:** Tests for a mutation previously identified in a family member; a copy of relative's lab result is REQUIRED.

Master Label

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