

THIS IS NOT A TEST REQUEST FORM. Please complete and submit with the test request form or electronic packing list.

G6PD DEFICIENCY TESTING PATIENT HISTORY FORM

Patient Name: _____ **Date of Birth:** _____ **Sex:** Female Male
Ordering Provider: _____ **Provider's Phone:** _____
Practice Specialty: _____ **Provider's Fax:** _____
Genetic Counselor: _____ **Counselor's Phone:** _____

Patient's Ethnicity/Ancestry (check all that apply)

African American/Black Asian Hispanic White Other: _____

List country of origin (if known): _____

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 individuals of African descent is 99%. Sensitivity in other ethnicities is unknown.
- 2001961 Familial Mutation, Targeted Sequencing:** Tests for a sequence variant 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.