

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 HEREDITARY HEMOLYTIC ANEMIA TESTING

Patient Name: _____ **Date of Birth:** _____ **Sex:** Female Male
Physician: _____ **Physician Phone:** _____
Practice Specialty: _____ **Physician Fax:** _____
Genetic Counselor: _____ **Counselor Phone:** _____

Patient Ethnicity/Ancestry (check all that apply)

African American/Black Asian Hispanic White Other _____

List countries of origin if known: _____

Suspected clinical diagnosis: _____

Does the patient have symptoms of a hereditary hemolytic anemia disorder? No Yes (check all that apply)

Anemia Fatigue Gallstones Hemolytic crisis Jaundice Splenomegaly

Other symptom(s): _____

Laboratory Findings (please attach recent CBC or provide values below)

CBC date: _____ RBC: _____ HGB: _____ HCT: _____ MCV: _____ MCHC: _____ RDW: _____ Retic: _____

Blood smear abnormalities: _____

Bilirubin..... Normal Not performed Abnormal: _____

Hemoglobin evaluation..... Normal Not performed Abnormal: _____

Osmotic Fragility..... Normal Not performed Abnormal: _____

EMA/RBC Band 3 Protein Reduction . Normal Not performed Abnormal: _____

G6PD..... Normal Not performed Abnormal: _____

Pyruvate kinase..... Normal Not performed Abnormal: _____

Other: _____ Normal Abnormal: _____

Other: _____ Normal Abnormal: _____

Has the patient undergone previous DNA testing for this condition?..... No Yes Unknown

If yes, describe the gene(s), methodology, and results: _____

Is there any relevant family history of hemolytic anemia disorder?..... No Yes Unknown

If yes, attach a pedigree or specify the relative's relationship to the patient. List their symptoms and age of onset: _____

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.

- 3000894** Hereditary Hemolytic Anemia Cascade
- 2012052** Hereditary Hemolytic Anemia Panel, Sequencing
- 3002059** Pyruvate Kinase Deficiency (PKLR) Sequencing:
Clinical sensitivity 98% for pyruvate kinase deficiency.
- 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.
- Other:** _____

Master Label

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