

Thalassemia/Hemoglobinopathy Patient Information

Instructions: The information requested below is important for interpretation of test results. To help us provide the best possible service, answer the questions completely and **send the paperwork with the specimen.** All answers will be kept confidential.

Patient Info	rmation						
Patient Name (La	st, First, Middle)					Birth Date (mm-dd-yyyy)	
Cov Assigned at	Dirth			Logol/Adm	iniotrotivo		
Sex Assigned at	b disclose						
□ Male □ Female □ Unknown □ Choose not to disclose □ Male □ Female □ Nonbinary							
Referring Provider Information							
Referring Provide	er Name <i>(Last, F</i>	irst)	Phone		Email		
Ethnic Background: Ethnic background is necessary to provide appropriate interpretation of test results.							
🗆 African 🗆 Ashkenazi Jewish 🗆 Asian 🗆 Arab 🗆 European 🗆 Other:							
Clinical History							
Reasons for Test	-						
□ Microcytosis □ Erythrocytosis □ Previously known hemoglobinopathy							
Prenatal/Carrier testing Cyanosis Diagnosis:							
□ Hemolytic anemia □ Abnormal newborn screen Previously tested at Mayo Clinic? □ Yes □ No							
□ Sickle monitor/treatment monitor □ Other:							
Family History If y Are other relatives known to be affected? Yes				If yes, explain disorder and relationship to patient:			
If relative was tested at Mayo Clinic, include the name of the family member.							
RBC	HGB	HCT	Recent transfusion history: Yes No Unknown				
		MCH	If yes, date(s) of last transfusion(s):				
		Retic count	Splenomegaly: Yes No Hydroxyurea treatment: Yes No				
Relevant Clinical							
As part of this e	evaluation:		·				
□ Do not perform molecular testing							
Perform molecular testing:							
Metabolic Laboratory Selected* molecular testing to explain: or exclude:							
Perform regardless of protein results: (check all that apply) □ Alpha Globin Gene MLPA† (α deletions)**							
\Box Beta Globin Gene Sequencing (β variants / β thal)							
\Box Beta Globin Cluster MLPA† (β cluster deletions: β thal, HPFH, $\delta\beta$ thal, $\epsilon\gamma\delta\beta$ thal)							
		uencing (α variants / nondeletional		, - ,			
* If checked, Ma	iyo experts sele	ect testing.					

- If a specific test is not performed, it can be added subsequently at client request (sample volume permitting).
- ** Assay available as a reflex test in the THEV1 / Thalassemia and Hemoglobinopathy Evaluation, Blood and Serum or separate order ATHAL test.
- † MPLA tests require a minimum of 3 mL of blood.