

## Metabolic Hematology 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 Informat	tion					
Name (Last, First, Middle)			Birth Date (mm-dd-yyyy)		Sex	
					☐ Male ☐ Female	
Referring Provider Nam	ne (Last, First)		Phone		Email	
Ethnic Origin/Ra	ace Check all that a	pply.				
		☐ Hispanic ☐ Mediterranear	n □ Southeast A	Asian		
Clinical History						
CBC Data	Relevant Clinical Information					
HGB:	□ Asymptomatic     □ Symptomatic:       □ Acquired     □ Lifelong/familial					
RBC:	Recent transfusion:	ansfusion date (mm-	fusion date (mm-dd-yyyy):			
MCH:	Hydroxyurea:	oxyurea:				
MCHC:	Family history:	☐ Yes ☐ No Disorde	er/relation to patien	nt:		
RDW:	Blood smear shows:					
Retics:						
	esting See Metabol	ic Hematology Profile Compariso	on Chart.		/ '   PE//E4	
Hemoglobin Disorder (consider THEV1 or HBEL1)  Genetic counseling or prenatal Abnormal newborn screen Anemia Microcytosis Other:		Hemolytic Anemia (consider HAEV1, RBCME, or EEEV1) Suspect		Erythrocytosis (consider REVE1)  JAK2 V617F: □ Pos □ Neg □ Not done  JAK2 Exon 12: □ Pos □ Neg □ Not done		
						☐ HS ☐ HE ☐ HP
		☐ Enzyme disorder: Phlebotomy: ☐ Yes		Phlebotomy:		
						☐ Monitoring of Hb fractions (order HGBCE) ☐ Cyanosis/Hypoxia (order MEV1 + P50B)
Test Reflex Opti	ons					
As part of HBEL1, THEV	1, HAEV1, REVE1 and	MEV1 evaluations, the following	g 4 options are avai	lable:		
1. Do <b>NOT</b> perforr	n molecular testing.					
2. Add only alpha	globin deletion/duplic	cation testing for common alpha	thalassemias.			
3. Mayo expert se	election of relevant mo	olecular testing (if needed) to ex	plain/exclude:			
4. Perform the fol	lowing tests regardles	ss of protein results:				
Additional Clinic						