

Laboratory Service Report

1-800-533-1710

Patient Name SAMPLEREPORT,HHEMO N	Patient ID SA00058757	Age 46	Gender F	Order # SA00058757
Ordering Phys CLIENT,CLIENT			•	DOB 06/10/1966
Client Order # SA00058757	Account Information			Report Notes
Collected 06/04/2013 00:00	C7028846-DLMP Rochester 3050 Superior Drive			
Printed 06/11/2013 15:19	Rochester, MN 55901			

Test	Flag Results	Unit	Reference Value	Perform Site*
Hemochromatosis HFE Gene Analysi	s,B			
RECEIVED: 06/04/2013 17:40 REPO	RTED: 06/10/2013 15:00			
Specimen	Blood			MCR
Specimen ID	1062039			MCR
Order Date	05 Jun 2013 (08:42		MCR
Method				MCR
A PCR-based assay was ut	ilized to test for the follo	wing		
three mutations in the H	FE gene: C282Y, H63D, and S6	5C.		
Because of the minimal e	ffect on iron metabolism ass	ociated		
with the S65C mutation,	it is only reported when it	is found		
with the C282Y mutation	(i.e. if the patient has the			
C282Y/S65C genotype).				
Results				MCR
C282Y: Not detected.				
H63D: Not detected.				
Interpretation				MCR

The diagnosis of HH cannot be excluded because approximately 5 to 8% of patients with HH in a North American Caucasian population do not have either the C282Y or H63D mutation. For other ethnic and racial groups, the percentage of patients with an unidentified mutation on both chromosomes may differ. This result does not rule out the presence of disease causing mutations in other regions of the HFE gene or in other genes associated with hemochromatosis. This

This result suggests a low risk for either a diagnosis of or

predisposition for hereditary hemochromatosis (HH).

The above interpretation assumes that testing is being performed for a possible diagnosis of HH. For carrier testing, the interpretation of this result depends on the family history and genotype of affected individuals.

result should be interpreted in the context of clinical presentation and results of other laboratory tests (e.g., serum transferrin-iron saturation and serum ferritin).

A genetic consultation may be of benefit. CAUTIONS:

Test results should be interpreted in context of clinical findings, family history, and other laboratory data. Misinterpretation of results may occur if the information provided is inaccurate or incomplete.

Rare polymorphisms exist that could lead to false negative

Performing Site Legend on Last Page of Report

Patient Name	Collection Date and Time	Report Status
SAMPLEREPORT,HHEMO N	06/04/2013 00:00	Final
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or positive results. If results obtained do not match the clinical findings, additional testing should be considered.

Bone marrow transplants from allogenic donors will interfere with testing. Call Mayo Medical Laboratories for instructions for testing patients who have received a bone marrow transplant.

Laboratory developed test. Reviewed By:

Emily Christine Lauer

Release Date 10 Jun 2013 14:57

* Performing Site:

MCR	Mayo Clinic Laboratories - Rochester Main Campus 200 First St SW Rochester, MN 55905	Lab Director: Franklin R. Cockerill, III, M.D.

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SAMPLEREPORT,HHEMO N	06/04/2013 00:00	Final
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