

### **Laboratory Service Report**

## 1-800-533-1710

Patient Name SAMPLEREPORT,CANW	Patient ID SA00007360	<b>Age</b> 32	Gender F	<b>Order #</b> SA00007360
Ordering Phys				<b>DOB</b> 04/04/1980
Client Order # SA00007360	Account Information			Report Notes
<b>Collected</b> 07/31/2012 13:00	C7028846-DLMP ROCHESTER 3050 SUPERIOR DRIVE			
<b>Printed</b> 08/15/2012 14:05	ROCHESTER,MN 559	901		

Test	Flag Results	Unit	Reference Value	Perform Site*
	-			
Canavan Disease, Mutation Analysis	5	REPORTED 0	8/14/2012 16:52	
Specimen	Blood			MCR
Specimen ID	976458			MCR
Order Date	01 Aug 2012 13:05			MCR
Reason For Referral				MCR
Carrier screen for Canava	n disease. Test for the presence of	E		
a mutation in the ASPA gen	ne.			
Method				MCR
A PCR-based assay was used	d to test for the following			
mutations in the ASPA gene	e: E285A, Y231X, 433(-2)A>G, and			
A305E.				
Results				MCR
None of the listed mutation	ons were detected.			
Interpretation				MCR
Having excluded the presen	nce of the listed mutations, the			
risk that this individual	is a carrier of another Canavan			

disease mutation is approximately 1/2001.

This risk calculation assumes that there is no family

history of Canavan disease. Additionally, this calculation is based on a mutation detection rate of 98% (for Ashkenazi Jewish ancestry) and a population carrier frequency of 1/41.

There is considerable variability both in the carrier frequency and detection rate among different ethnic groups. The risk estimate provided, therefore, is our best approximation given the information available to us for this patient and for this population.

#### 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 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.

#### \*\*\*Performing Site Legend on Last Page of Report\*\*\*

	<b>Collection Date and Time</b> 07/31/2012 13:00	Report Status Final
Page 1 of 2		>> Continued on Next Page >>



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Test Flag Results Unit Reference Value Site\*

Laboratory developed test.

Reviewed By:
Release Date

D Brian Dawson PhD
Release Date

MCR
MCR

#### \* Performing Site:

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

Patient Name	Collection Date and Time	Report Status
SAMPLEREPORT.CANW	07/31/2012 13:00	Final
Page 2 of 2	07/01/2012 10:00	** End of Report **

<sup>\*</sup> Report times for Mayo performed tests are CST/CDT