

<b>Patient Name</b> TESTINGRNV,REPORTS	<b>Patient ID</b> SA00057597	<b>Age</b> 20D	<b>Gender</b> F	<b>Order #</b> SA00057597
<b>Ordering Phys</b> CLIENT,CLIENT				<b>DOB</b> 05/01/2013
<b>Client Order #</b> SA00057597	<b>Account Information</b>			<b>Report Notes</b>
<b>Collected</b> 05/21/2013 00:00	C7028846-DLMP Rochester 3050 Superior Drive Rochester, MN 55901			
<b>Printed</b> 06/25/2013 13:38				

Test	Flag	Results	Unit	Reference Value	Perform Site*
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**Hunter Syndrome, Known Mutation**
**RECEIVED:** 05/24/2013 08:50 **REPORTED:** 05/24/2013 08:50

## Reason for Referral

MCR

Family history of mucopolysaccharidosis type II (Hunter syndrome). Test for the presence of the familial alteration in the IDS gene.

## Result

MCR

The p.Q531X alteration was NOT detected.

## Interpretation

MCR

Absence of the mutation previously identified in an affected family member indicates that this individual is at no greater risk than someone in the general population to be a carrier of or affected with mucopolysaccharidosis type II, also known as Hunter syndrome.

This assay does not rule out the presence of other disease causing mutations in this gene or other genes associated with metabolic disease. Errors in the diagnosis or pedigree provided to us may lead to an erroneous interpretation of test results.

A genetic consultation may be of benefit.

Unless reported or predicted to cause disease, alterations found deep in the intron or alterations that do not result in an amino acid substitution are not reported. These and common polymorphisms identified for this patient are available upon request.

**CAUTIONS:**

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.

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

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>Patient Name</b> TESTINGRNV,REPORTS	<b>Collection Date and Time</b> 05/21/2013 00:00	<b>Report Status</b> Final
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\* Report times for Mayo performed tests are CST/CDT

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Test	Flag	Results	Unit	Reference Value	Perform Site*
Laboratory developed test.					
Method		DNA sequence analysis was used to test for the presence of the p.Q531X (c.1591C>T) alteration in exon 9 of the IDS gene. Testing was performed for this specific alteration because it was previously identified in a family member of this individual. Mutation nomenclature is based on GenBank accession number NM_000202.5.			MCR
Specimen		Blood			MCR
Reviewed By		Devin Oglesbee PhD			MCR
Release Date		24 May 2013 08:48			MCR

\* Performing Site:

MCR	Mayo Clinic Laboratories - Rochester Main Campus 200 First St SW Rochester, MN 55905	Lab Director:
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