Patient Name	Patient ID	Age	Gender	Order #
SAMPLEREPORT,VHLKP ABNORMAL	SA00067258	47	F	SA00067258
Ordering Phys CLIENT,CLIENT				**DOB** 06/10/1966
Client Order # SA00067258	**Account Information**			**Report Notes**
Collected 04/29/2014 11:00	C7028846-DLMP Rochester SDSC 2 - Client Support Rochester, MN 55901			
Printed 05/02/2014 07:34				

Test	Flag	Results	Unit	Reference Value	Perform Site*
VHL Known Mutation					
VHL Known Mutation					
Reason For Referral		Familial mutation			MCR
A VHL gene mutation was previously identified in an affected family member. Test for the presence of the following mutation in the VHL gene: Exon 3 c.481C>T p.R161X					
Method		Familial mutation			MCR
Fluorescent DNA sequence analysis was used to test for the presence of a specific mutation in the VHL gene, which was previously identified in an affected family member of this individual(GenBank accession number: NM_000551.2).					
Result		Positive			MCR
This individual was shown to be heterozygous for the familial mutation: Exon 3 c.481C>T p.R161X					
Interpretation		Positive			MCR
The above mutation was previously identified in a family member affected with Von Hippel-Lindau (VHL) syndrome. The presence of this mutation in the individual tested here is consistent with a diagnosis of VHL syndrome for this individual. Therefore, this individual is at risk for developing symptoms associated with VHL. Appropriate screening procedures should be considered. Since a mutation has been identified in the VHL gene in this individual, genetic testing for this specific mutation in other family members is recommended. Please contact the Endocrine Laboratory at 1-800-533-1710 or the online test catalog at mayomedicallaboratories.com for information about how to order VHL Known Mutation (89084/VHLKP) for sequencing mutations.					
Comment		GC consult			MCR
A genetic consultation may be of benefit.					
Reviewed By		LYNETTE HANSEN			MCR
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. A list of common polymorphisms identified for this patient is available from the Endocrine Laboratory upon request. Rarely, individuals may have a mutation or deletion in the					

Performing Site Legend on Last Page of Report

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gene(s) tested that is not identified by the described testing methodology. In addition, the phenotype observed in the individual tested here may be due to a variant in a gene not analyzed by this test. Laboratory developed test.					
VHL Known Mutation Sequencing					
Sequencing		Performed			MCR

RECEIVED: 05/01/2014 15:03 **REPORTED:** 05/01/2014 15:34

* Performing Site:

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