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# **Laboratory Service Report**

# 1-800-533-1710

Patient Name	Patient ID	Age	Gender	Order #
TESTINGRNV,APO1K	SA00048293	67	M	SA00048293
Ordering Phys		•	•	DOB
				06/15/1945
Client Order #	Account Information			Report Notes
SA00048293				
Collected	C7028846-DLMP RO	CHESTER		
08/08/2012	3050 SUPERIOR DRIVE			
Printed	ROCHESTER,MN 559	01		
08/31/2012 14:47				

Test	Flag	Results	Unit	Reference Value	Perform Site*
1650	riag	Kesaics	OHIC	varue	DICE
POA1 Gene, Known Mutation			REPORTED (	08/08/2012 15:01	
Specimen		Blood			MCR
Specimen ID		1038564			MCR
Order Date		08 Aug 2012 14:56			MCR
Reason For Referral					MCR
Family history of apolipopro	tein A-I (AF	POA1) associated			
familial amyloidosis. Test	for the pres	sence of a mutation			
in the APOA1 gene.					
Method					MCR
DNA sequence analysis was us	sed to test f	for the presence of			
the c.148G>C (p.G50R) mutati	on in exon 2	of the APOA1 gene.			
Testing was performed for th	is specific	mutation because it			
was previously identified in	an affected	d family member of			
this individual. Mutation n	omenclature	is based on GenBank			
accession number; NM_000039.	1.				
Result					MCR
The following sequence chang	ge was detect	ed:			
Exon: 2					
DNA change: c.148G>C					
Amino acid change: p.G50R (G	ly50Arg)				
Legacy nomenclature: p.G26R	(Gly26Arg)				
Classification: DELETERIOUS					
Interpretation					MCR
This alteration is a known d	leleterious m	nutation.			
This result is consistent wi	th a diagnos	sis of APOA1			
associated familial amyloido	_				

Since a mutation has been identified, testing of at risk family members is possible.

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.

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

Patient Name	Collection Date and Time	Report Status	
TESTINGRNV,APO1K	08/08/2012	Final	
Page 1 of 2		>> Continued on Next Page >>	

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



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Test Flag Results Unit Value Site\*

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.

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 Melody Elizabeth Kimball Release Date

MCR

08 Aug 2012 14:59 MCR

### \* Performing Site:

MCR Mayo Clinic Laboratories - Rochester Main Campus Lab Director: Franklin R. Cockerill, III, M.D.

Patient Name	Collection Date and Time	Report Status
TESTINGRNV,APO1K	08/08/2012	Final
Page 2 of 2		** End of Report **

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