

### **Laboratory Service Report**

## 1-800-533-1710

Patient Name	Patient ID	Age	Gender	Order #
SAMPLEREPORT, CHEKK	SA00064002	27	M	SA00064002
Ordering Phys CLIENT,CLIENT				<b>DOB</b> 06/25/1986
Client Order # SA00064002	Account Information			Report Notes
<b>Collected</b> 10/30/2013 00:00	C7028846-DLMP Rochester SDSC 2 - Client Support			
<b>Printed</b> 12/24/2013 09:26	Rochester, MN 55901			

Test Flag Results Unit Value Site\*

CHEK2 Gene, Known Mutation

**RECEIVED:** 10/30/2013 13:21 **REPORTED:** 12/02/2013 12:58

Reason For Referral

MCR

Family history of breast cancer. Test for the presence of a mutation in the CHEK2 gene.

Result

ılt MCR

The c.1100delC mutation was NOT detected.  $\bar{\tau}$ 

Interpretation

MCR

Absence of the CHEK2 mutation previously identified in an affected family member may decrease this individual's risk of breast cancer given that the CHEK2 mutation may be contributing to an increased risk for breast cancer in this family (J Clin Oncol 2011 29:3747-3752). However, this assay does not rule out the presence of mutations within other genes associated with breast cancer susceptibility which may also be contributing to cancer risk in the patient and/or family.

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:

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.

Laboratory developed test.

Method

MCR

 ${\tt DNA}$  sequence analysis was used to test for the presence of

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

Patient Name	Collection Date and Time	Report Status
SAMPLEREPORT, CHEKK	10/30/2013 00:00	Final
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<sup>\*</sup> Report times for Mayo performed tests are CST/CDT



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Test	Flag Results	Unit	Reference Value	Perform Site*
the c.1100delC alteration in accession number, NM_007194;				
Specimen	Blood			MCR
Reviewed By				MCR
Matthew John Ferber PhD				
Release Date	02 Dec 2013 12:5	57		MCR

## \* Performing Site:

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

Patient Name	Collection Date and Time	Report Status
SAMPLEREPORT, CHEKK	10/30/2013 00:00	Final
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