

Laboratory Service Report

1-800-533-1710

Patient Name	Patient ID	Age	Gender	Order #	
SAMPLEREPORT,BA190	SA00067483	47	F	SA00067483	
Ordering Phys CLIENT,CLIENT		·		DOB 06/10/1966	
Client Order # SA00067483	Account Information			Report Notes	
Collected	C7028846-DLMP Roc	C7028846-DLMP Rochester			
05/14/2014 00:00		SDSC 2 - Client Support			
Printed	Rochester, MN 55901				
05/15/2014 14:45					

Reference Perform
Test Flag Results Unit Value Site*

BCR/ABL, p190, Quant, Monitor

Specimen Type Bone marrow MCR
Final Diagnosis: MCR

Bone marrow, BCR/ABL mRNA level analysis (p190 fusion form):

Negative. No BCR/ABL p190 mRNA transcripts were detected (%bcr/abl(p190):abl=0).

Signing Pathologist: Melissa Tricker-Klar Method summary-BCR/ABL, p190 fusion: BCR/ABL p190 mRNA transcript level was evaluated using quantitative, reverse transcription PCR. The detection limit for this assay is 0.01%. See Mayo Medical Laboratories Interpretive Handbook for method details.

The assay detects the most common fusion form (e1/a2), but does not detect other fusions, including the p210, which is the most common form found in chronic myelogenous leukemia. This assay should only be ordered for monitoring patients with a previously identified p190 fusion form. Test #89006(BCR/ABL, mRNA detection, RT-PCR, Quantitative, Diagnostic) should be ordered if the test is being performed in a diagnostic setting and test #89007(BCR/ABL, p210, mRNA detection, RT-PCR, Quantitative, Monitoring CML) should be ordered if this patient is being monitored for a known p210 fusion form. Please contact the Mayo Molecular Hematopathology Laboratory at 507-266-0489 with questions or if additional testing is required.

The reproducibility of this assay is such that results within 0.5 log should be considered equivalent. Trends in the level of BCR/ABL mRNA should be followed and clinically significant changes verified with a subsequent specimen. Please contact the Mayo Molecular Hematopathology Laboratory at 507-266-0489 with questions or if additional testing is required. Laboratory developed test.

RECEIVED: 05/15/2014 10:12 **REPORTED:** 05/15/2014 10:58

Performing Site Legend on Last Page of Report

Patient Name	Collection Date and Time	Report Status
SAMPLEREPORT,BA190	05/14/2014 00:00	Final
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* 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,BA190	05/14/2014 00:00	Final
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^{*} Report times for Mayo performed tests are CST/CDT