

Laboratory Service Report

1-800-533-1710

MCR

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Patient Name SAMPLEREPORT,BCRAB	Patient ID C7028846-001896	Age 47	Gender F	Order # X100067674
Ordering Phys DOC,DOC		·		DOB 01/10/1966
Client Order # X100067674	Account Information	Account Information		Report Notes
Collected 06/09/2013 00:00	3050 Superior Drive	·		
Printed 06/10/2013 14:46	Rochester, MN 55901			

Test Flag Results Unit Value Site*

BCR/ABL, p210, Quant, Monitor

RECEIVED: 06/10/2013 12:42 **REPORTED:** 06/10/2013 14:21

Specimen Type Peripheral blood

Final Diagnosis:

Peripheral blood, BCR/ABL mRNA level analysis (p210 fusion

form):

Negative. No BCR/ABL p210 mRNA transcripts were detected (% bcr/abl(p210):abl=0). The detection limit for this sample was 0.0002%.

Signing Pathologist: Melissa Tricker-Klar p210 mRNA Method summary - BCR/ABL, p210 fusion: transcript level was evaluated using quantitative, reverse transcription PCR. The assay detects the two most common fusion forms in chronic myelogenous leukemia: e13/a2 and e14/a2, which code for p210 proteins. It is intended for monitoring patients with neoplasms known to carry the p210 fusion form. The assay does not detect other fusions, including those for the p190 protein commonly present in acute lymphoblastic leukemia. This assay should not be used in the diagnostic setting, as it does not detect all bcr/abl fusion forms. If this has been ordered in a diagnostic setting and the result is negative, test #89006(BCR/ABL mRNA Detection, RT-PCR, Qualitative, Diagnostic) should be ordered to evaluate for all possible fusion forms. Please contact the lab at 1-507-266-0489 with questions or if additional testing is required. See the Mayo Medical Laboratories Interpretive Handbook for method details.

Typical clinical samples have detection limits ranging from 0.01% to 0.0001% bcr/abl:abl. Most CML patients at diagnosis have a bcr/abl:abl result in the range of 20% to 100%.

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. Laboratory developed test.

* Performing Site:

MCR	Mayo Clinic Laboratories - Rochester Main Campus 200 First St SW Rochester. MN 55905	Lab Director: Franklin R. Cockerill, III, M.D.
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Patient Name	Collection Date and Time	Report Status
SAMPLEREPORT,BCRAB	06/09/2013 00:00	Final
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^{*} Report times for Mayo performed tests are CST/CDT