

Patient Name SAMPLEREP, HV1CM A	Patient ID SA00046039	Age 40	Gender F	Order # SA00046039
Ordering Phys				DOB 09/28/1971
Client Order # SA00046039	Account Information			Report Notes
Collected 09/25/2012 13:41	C7028846-DLMP ROCHESTER 3050 SUPERIOR DRIVE ROCHESTER, MN 55901			
Printed 11/05/2012 15:08				

Test	Flag	Results	Unit	Reference Value	Perform Site*
HIV-2 Ab Eval, S		Negative		REPORTED 09/25/2012 13:44 Negative	SDL
<p>A cadaveric or hemolyzed specimen was received for testing. Hemoglobin could interfere with test results. Interpret result with caution. Testing is performed using the Bio-Rad GS HIV-2 antibody enzyme immunoassay.</p>					
HIV 2 Antibody, IBL		NEGATIVE		REPORTED 09/27/2012 15:09	Y03 8
<p>HIV 2 Ab, Immunoblot</p> <p>A Negative HIV-2 antibody Immunoblot result does not exclude HIV-2 infection since the time frame for seroconversion is variable. If acute HIV-2 infection is suspected, suggest testing a new plasma sample by an amplified nucleic acid test (HIV-2 DNA/RNA, unit code 49000). Testing for HIV-1 should also be considered.</p> <p>Reference range: NEGATIVE</p> <p>This test was performed using a kit that has not been cleared or approved by the FDA. The analytical performance characteristics of this test have been determined by Focus Diagnostics. This test should not be used for diagnosis without confirmation by other medically established means.</p>					
HIV-1 Ab Confirm by IFA, S		Negative		REPORTED 09/25/2012 13:44 Negative	SDL
<p>Repeat testing is recommended in 1 to 2 months for those at risk for HIV infection. A cadaveric or hemolyzed specimen was received for testing. Hemoglobin could interfere with test results. Interpret result with caution.</p>					
HIV-1/-2 Ab Screen Hemolyzed, S	AB	Reactive		REPORTED 09/25/2012 13:43 Negative	SDL
<p>Western blot confirmatory test is ordered. Confirmatory test is the definitive test for HIV-1/-2 infection.</p>					
HIV-1/-2 Ab Confirm Eval, S		Positive		REPORTED 09/25/2012 13:43 Negative	SDL
<p>HIV-1 Ab Confirm Western Blot, S</p> <p>REPORTABLE DISEASE. A positive result is based on the presence of at least two of the following three bands: p24,</p>					

Performing Site Legend on Last Page of Report

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* Report times for Mayo performed tests are CST/CDT

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gp41, and gp120/gp160. The U.S. Association of Public Health Laboratories recommends verification of first-time positive test results for the diagnosis of HIV infection. A second specimen should be submitted for testing to verify all such positive results. A cadaveric or hemolyzed specimen was received for testing. Hemoglobin could interfere with test results. Interpret result with caution.

* Performing Site:

SDL	Mayo Clinic Laboratories - Rochester Superior Drive 3050 Superior Dr. NW Rochester, MN 55901	Lab Director:
Y038	Focus Diagnostics, Inc. 5785 Corporate Avenue Cypress, CA 90630-4750	Lab Director:

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