MAYO CLINIC
Mayo Medical Laboratories

Laboratory Service Report 1-800-533-1710

Patient Name	Patient ID	Age	Gender	Order #
SAMPLEREPORT,HV1CD A	SA00046037	40	F	SA00046037
Ordering Phys				**DOB** 09/28/1971
Client Order # SA00046037	**Account Information**			**Report Notes**
Collected 09/25/2012 10:32	C7028846-DLMP ROCHESTER 3050 SUPERIOR DRIVE ROCHESTER,MN 55901			
Printed 11/05/2012 15:07				

Test	Flag	Results	Unit	Reference Value	Perform Site*
			REPORTED 09/25/2012 13:37		
HIV-2 Ab Eval, S		Negative		Negative	SDL
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.					
HIV 2 Antibody, IBL			REPORTED 09/27/2012 15:08		
HIV 2 Ab, Immunoblot		NEGATIVE			Y038
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. Reference range: NEGATIVE 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.					
			REPORTED 09/25/2012 13:37		
HIV-1 Ab Confirm by IFA, S		Negative		Negative	SDL
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.					
			REPORTED 09/25/2012 13:36		
HIV-1/-2 Cadaver/Hemolyzed, S	AB	Reactive		Negative	SDL
Western blot confirmatory test is ordered. Confirmatory test is the definitive test for HIV-1/-2 infection.					
HIV-1/-2 Ab Confirm Eval, S			REPORTED 09/25/2012 13:37		
HIV-1 Ab Confirm Western Blot, S	AB	Positive		Negative	SDL
REPORTABLE DISEASE. A positive result is based on the presence of at least two of the following three bands: p24,					

Performing Site Legend on Last Page of Report

* 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: