MAYO CLINIC
Mayo Medical Laboratories

Laboratory Service Report 1-800-533-1710

Patient Name	Patient ID	Age	Gender	Order #
SAMPLEREPORT,HIVE A	SA00049269	46	F	SA00049269
Ordering Phys				**DOB** 06/10/1966
Client Order # SA00049269	**Account Information**			**Report Notes**
Collected 09/25/2012 13:14	C7028846-DLMP ROCHESTER 3050 SUPERIOR DRIVE ROCHESTER,MN 55901			
Printed 10/11/2012 10:32				

Test	Flag	Results	Unit	Reference Value	Perform Site*
			REPORTED 09/25/2012 13:22		
HIV-2 Ab Eval, S		Negative		Negative	SDL
Testing is performed using the Bio-Rad GS HIV-2 antibody enzyme immunoassay.					
HIV 2 Antibody, IBL			REPORTED 09/27/2012 15:07		
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 13:17		
HIV-1/-2 Ab Evaluation, S	AB	Reactive		Negative	SDL
Not diagnostic. Western blot confirmatory testing ordered. Testing is performed using the Ortho Vitros Anti-HIV 1+2 chemiluminescence immunoassay.					
			REPORTED 09/25/2012 13:21		
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.					
HIV-1/-2 Ab Confirm Eval, S			REPORTED 09/25/2012 13:21		
HIV-1 Ab Confirm Western Blot, S		Negative		Negative	SDL
No bands detected. If acute HIV infection is suspected in a high-risk individual, HIV-1 and/or HIV-2 nucleic acid test is recommended, and a plasma specimen needs to be collected and submitted for testing.					

Performing Site Legend on Last Page of Report

* Report times for Mayo performed tests are CST/CDT

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>> Accession SA00049269 - Continued From Previous Page <<
>> Do Not Discard <<

* Performing Site:

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

Patient Name	Collection Date and Time	Report Status
SAMPLEREPORT,HIVE A	09/25/2012 13:14	Final
Page 2 of 2		** End of Report **

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