



BML496925

Laboratory Investigation Report

Name	: Ms. TIFFANY IMANEH	Ref No.	: 45102
DOB	: 01/12/1996	Sample No.	: 2412508740
Age / Gender	: 28 Y / Female	Collected	: 03/12/2024 19:00
Referred by	: CITICARE MEDICAL CENTER	Registered	: 04/12/2024 14:48
Centre	: CITICARE MEDICAL CENTER	Reported	: 04/12/2024 17:08

IMMUNOLOGY

Test	Result	Flag	Unit	Reference Range	Methodology
HEPATITIS B SURFACE ANTIGEN (HBSAG)	0.33		COI	Non-Reactive: <0.9 Borderline: =/>0.9 - <1.0 Reactive: =/>1.0	ECLIA

Note changes in method and reference range.
Source: Roche IFU.

INTERPRETATION NOTES :

A positive HBsAg test result means that the patient is infected with acute or chronic hepatitis B virus or chronic HBV carrier state. A negative result implies the patient is not infected with hepatitis B.

HEPATITIS C ANTIBODIES	0.04		COI	Non-Reactive: <0.9 Borderline: =/>0.9 - <1.0 Reactive: =/>1.0	ECLIA
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Source: Roche IFU.

INTERPRETATION NOTES :

A non-reactive screening test result does not exclude the possibility of exposure to or infection with HCV. Non-reactive screening results in individuals with prior exposure to HCV may be due to low antibody levels that are below the limit of detection of this assay or lack of reactivity to the HCV antigens used in this assay. Patients with acute or recent HCV infections (< 3 months from time of exposure) may have false-negative HCV antibody results due to the time needed for seroconversion (average of 8 - 9 weeks). Testing for HCV RNA and or RIBA is recommended.

A repeatedly reactive screening result is consistent with current HCV infection, or past HCV infection that has resolved, or biologic false positivity for HCV antibody. Testing for HCV RNA and or RIBA is recommended

HIV I & II ANTIBODY AND P24 ANTIGEN	0.21		S/CO	Non-Reactive: <1.0 Reactive: =/>1.0 Source: Roche IFU.	ECLIA
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INTERPRETATION NOTES :

1. A negative test result does not completely rule out the possibility of an infection with HIV. Serum or plasma samples from the very early (preseroconversion) phase or the late phase of HIV infection can occasionally yield negative findings. Yet unknown HIV variants can also lead to a negative HIV finding. The presence of antibodies to HIV is not a diagnosis of AIDS.

2. For diagnostic purposes, the results should always be assessed in conjunction with the patient's medical history, clinical examination and other

Dr. Adley Mark Fernandes
M.D (Pathology)
Pathologist

Gyoma V. Shah
Dr. Vyoma V Shah
M.D (Pathology)
Clinical Pathologist

Harshad Manikandan
HARSHAD MANIKANDAN
Laboratory Technician

This is an electronically authenticated report

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Test result pertains only to the sample tested and to be interpreted in the light of clinical history. These tests are accredited under ISO 15189:2012 unless specified by (^). Test marked with # is performed in an accredited referral laboratory.





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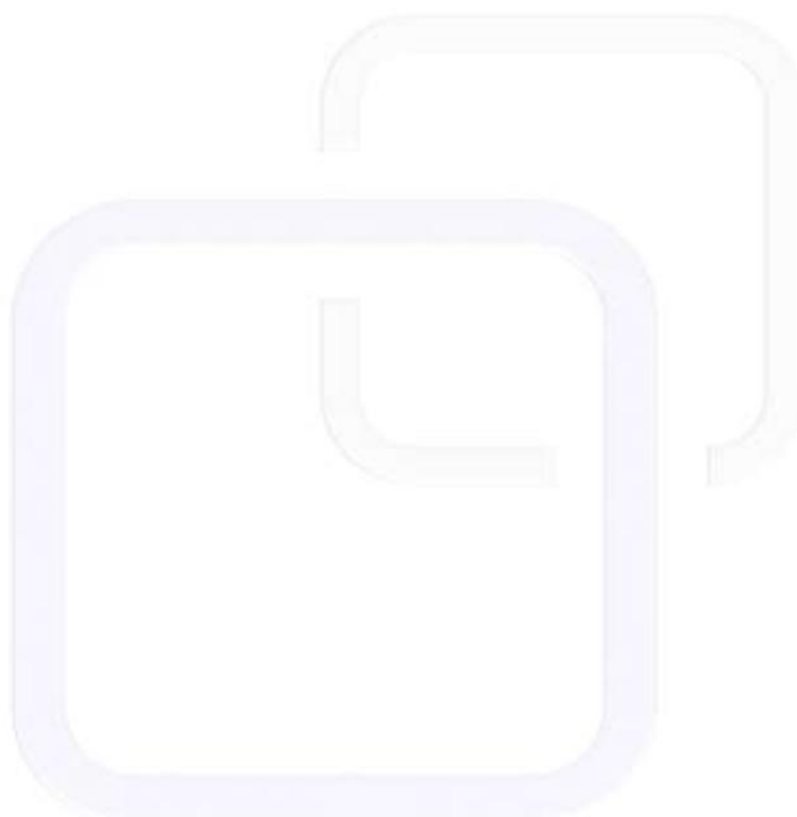
IMMUNOLOGY

Test	Result	Flag	Unit	Reference Range	Methodology
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findings.

3. This is a screening test.

Source: Roche Cobas IFU.



Dr. Adley Mark Fernandes
M.D (Pathology)
Pathologist

Vyoma V. Shah
Dr. Vyoma V Shah
M.D (Pathology)
Clinical Pathologist

Harshad Manikandan
HARSHAD MANIKANDAN
Laboratory Technician

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SEROLOGY

Test	Result	Flag	Unit	Reference Range	Methodology
RPR (RAPID PLASMA REAGEN)	Non-Reactive			Non-reactive	Carbon flocculation

INTERPRETATION NOTES :

Syphilis is a disease caused by infection with the spirochete *Treponema pallidum*. The infection is systemic and the disease is characterized by periods of latency.

Patients with primary or secondary syphilis should be reexamined clinically and serologically 6 months and 12 months following treatment.

Typically, rapid plasma reagin (RPR) titers decrease following successful treatment, but this may occur over a period of months to years.

Biological false-positive reactions with cardiolipin-type antigens have been reported in disease such as infectious mononucleosis, leprosy, malaria, lupus erythematosus, vaccinia, and viral pneumonia. Pregnancy, autoimmune diseases, and narcotic addictions may give false-positives. Pinta, yaws, bejel, and other treponemal diseases may also produce false-positive results with this test.

False negatives tend to be more common in the initial and end stages of infection. Among people who are in the secondary (middle) stage of infection, the RPR test result is nearly always positive. (Interpretation added on 28 Dec 2019).

Sample Type : Serum

End of Report

Dr. Adley Mark Fernandes
M.D (Pathology)
Pathologist

Vyoma V. Shah
Dr. Vyoma V Shah
M.D (Pathology)
Clinical Pathologist

Pradeep Dhamotharan
Pradeep Dhamotharan
Laboratory Technologist

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