

Patient Name : Mr MR.DUMMY

DOB/Age/Gender : 23 Y/Male

Patient ID / UHID : 8052567/RCL7249136

Referred By : Dr. X

Sample Type : Serum

Sample Collected : Apr 26, 2024, 01:00 PM

Report Date : May 25, 2024, 02:52 PM.

Barcode No : SI485384

Report Status : Final Report

Test Description	Value(s)	Unit(s)	Reference Range
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Syphilis Antibodies

Syphilis Antibodies (Treponema) EIA, SERUM	0.21	Ratio	Negative: < 1.00 Positive: >= 1.00
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Interpretation:**NOTE**

1. A negative result for an individual subject indicates absence of detectable anti - TP antibodies. However , a negative test result does not preclude the possibility of exposure to or infection with TP.
2. Some specimens containing unusually higher titre of hetrophile antibodies or rheumatoid factor may affect expected results.

INTERPRETATION

Syphilis antibody test is a solid phase enzyme linked immunosorbant assay for the qualitative detection of antibodies (IgG, IgM, IgA) against Treponema pallidum in human serum or plasma. Treponema pallidum, a spirochete bacterium, is the causative agent of the venereal diseases syphilis . The IgM antibody may be detectable towards the second week of infection , while IgG antibody appears later, at about 4 weeks . These antibodies could last for several years or even decades in the serum of a patient with untreated latent syphilis . Antibody to RPR antigen does not develop until 1-4 weeks after the appearance of the chancre, thus this antigen lacks of sensitivity to primary syphilis.

*** End Of Report ***

Disclaimer: Method given in report are only indicative and can be changed depending upon type of machine and kit available at time of testing.
Not all tests at all locations are under NABL scope. Availability of tests under NABL scope varies from lab to lab.



Booking Centre :- DEMO PARTNER CHENNAI, DEMO PARTNER CHENNAI
Processing Lab :-

 928-909-0609 ccsupport@redcliffelabs.com www.redcliffelabs.com

All Lab results are subject to clinical interpretation by qualified medical professional and this report is not subject to use for any medico-legal purpose.


Dr. Dummy

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1. The presented findings in the Reports are intended solely for informational and interpretational purposes by the referring physician or other qualified medical professionals possessing a comprehensive understanding of reporting units, reference ranges, and technological limitations. The laboratory shall not be held liable for any interpretation or misinterpretation of the results, nor for any consequential or incidental damages arising from such interpretation.
2. It is to be presumed that the tests performed pertain to the specimen/sample attributed to the Customer's name or identification. It is presumed that the verification particulars have been cleared out by the customer or his/her representation at the point of generation of said specimen / sample. It is hereby clarified that the reports furnished are restricted solely to the given specimen only.
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5. The Customers assume full responsibility for apprising the Company of any factors that may impact the test finding. These factors, among others, includes dietary intake, alcohol, or medication / drug(s) consumption, or fasting. This list of factors is only representative and not exhaustive.

DISCLAIMER

This is a sample report provided for demonstration purposes only and does not represent an actual patient report. Test results, reference ranges, methodologies, instrumentation, and report formats may vary depending on the laboratory performing the test. The format and representation shown are indicative of reports generated by the National Reference Laboratory of Redcliffe Labs, Noida. This sample report should not be used for medical interpretation, diagnosis, or treatment decisions.