LABORATORY REPORT

Redcliffe labs

Patient Name	: **Mr MR.DUMMY**		
DOB/Age/Gender	: 23 Y/Male	Sample Collected	: Apr 26, 2024, 01:00 PM
Patient ID / UHID	: 8052806/RCL7249292	Report Date	: May 24, 2024, 06:43 PM.
Referred By	: Dr. X	Barcode No	: SI484394
Sample Type	: Serum	Report Status	: Final Report

Test Description	Value(s)	Unit(s)	Reference Range
VDRL- RPR Titre (Semi Quant)			
RPR, Serum *Slide Flocculation*	Non Reactive		Non Reactive

Interpretation:

RESULTS	REMARKS
Reactive	Indicates presence of IgM & IgG antibodies against Treponemal Pallidum antigens
Non Reactive	Indicates absence of IgM & IgG antibodies against Treponemal Pallidum antigens

Note

1. Titres of ≥1: 8 and rising titres are significant.
2. Titres are reported only in reactive cases
3. Positive reaction indicates ongoing or recent infection and the diagnosis should be confirmed by specific Treponemal tests such as TPHA & FTA- AbS.
4. The reactivity will vary with Primary (60-86%), Secondary (99%) and Tertiary (98%) stage of Syphilis.
5. False positive reaction may be observed in patients of Malaria, Hepatitis, Mumps, Leprosy, Infectious Mononucleosis, Rheumatoid Arthritis and Collagen disease.
6. False negative reaction may be due to processing of sample collected early in the course of disease, immunosuppression and due to prozone effect.
7. The test is conducted on serum.
8. This is a Semi-quantitative test.

Uses

1. To screen for presence of Syphilis infection.
2. To monitor the progression of disease.
3. To asses the response to therapy (decreasing titres) in patients being treated for 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.

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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