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	: 8052367/RCL7248820	Report Date	: May 25, 2024, 01:59 PM.
Referred By	: Dr. Dr. X	Barcode No	: SI485048
Sample Type	: Serum	Report Status	: Final Report

Herpes Simplex Virus (HSV) 1 + 2 IgG & IgM Antibody Panel

Test Description	Value(s)	Unit(s)	Reference Range
Herpes simplex virus 1+2, IgG *EIA, SERUM*	1.28	RATIO	NEGATIVE: < 0.8
Herpes simplex virus 1+2, IgM *EIA, SERUM*	0.32	RATIO	NEGATIVE: < 0.8

Interpretation:

HSV 1 + 2 IgG

HSV 1 + 2 IgG RESULT IN RATIO	REMARKS
RATIO <0.80	NEGATIVE
RATIO >-0.80-<1.10	BORDERLINE
RATIO >-1.10	POSITIVE

Note 1. This assay is used for qualitative detection of specific IgG antibodies to Herpes Simplex virus (1+2) in serum samples only.
2. Positive result indicates past infection with Herpes Simplex virus or administration of HSV immunoglobulins. Pregnant females with positive HSV specific IgG antibodies are considered to be immune and hence risk of transmission of infection to fetus is minimal.
3. Borderline results should be re-tested in 10-14 days.
4. Negative result indicates person has not been exposed to Herpes Simplex virus in the past. Patients with negative results in suspected disease should be re-tested after 10-14 days. False negative results can be due to immunosuppression or due to low/undetectable level of IgG antibodies.
5. HSV serology cannot distinguish genital from nongenital infections.
6. The result should be interpreted in conjunction with clinical finding and other diagnostic tests.
7. The magnitude of the measured result is not indicative of the amount of antibody present.

HSV 1 + 2 IgM

HSV 1 + 2 IgM RESULT IN RATIO	REMARKS
RATIO <0.80	NEGATIVE
RATIO >-0.80-<1.10	BORDERLINE
RATIO >-1.10	POSITIVE

Note 1. This assay is used for qualitative detection of specific IgM antibodies to Herpes Simplex virus (1&2) in serum samples only. 2. Positive result for Herpes Simplex virus IgM may indicate acute infection, reinfection or reactivation of latent virus. Persistence of low level HSV IgM antibodies following post infection over a long period is not uncommon. False positive reaction may occur due to high levels of rheumatoid factor or during the course of other viral illnesses due to cross reactivity.
3. An borderline result requires repeat testing in 10-14 days.
4. Negative result for Herpes Simplex virus IgM indicates no Herpes Simplex virus infection. False negative reaction may be due to processing of sample collected early in the course of disease or absence of immune response.
5. HSV serology cannot distinguish genital from nongenital infections.
6. The result should be interpreted in conjunction with clinical finding and other diagnostic tests.
7. The magnitude of the measured result is not indicative of the amount of antibody present.

Dr. Dummy

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

*** 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

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

Terms and Conditions of Reporting

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.

3. It is to be noted that variations in results may occur between different laboratories and over time, even for the same parameter for the same Customer. The assays are performed and conducted in accordance with standard procedures, and the reported outcomes are contingent on the specific individual assay methods and equipment(s) used, as well as the quality of the received specimen.

4. This report shall not be deemed valid or admissible for any medico-legal purposes.

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.