

Patient Name : Mr MR.DUMMY

DOB/Age/Gender : 23 Y/Male

Patient ID / UHID : 8051913/RCL7248505

Referred By : Dr. Dr. X

Sample Type : Serum

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

Report Date : May 25, 2024, 12:01 PM.

Barcode No : SI484614

Report Status : Final Report

Test Description	Value(s)	Unit(s)	Reference Range
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Chikungunya IgM, Rapid Card

Chikungunya IgM, Rapid Card Immunochromatographic Assay	Negative		Negative
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Interpretation:**Note:**

- 1.It is a screening test
- 2.The Onsite Chikungunya IgM Rapid test is limited to the qualitative detection of IgM in human sera .
- 3.A negative result indicates absence of detectable IgM antibodies. However a negative result does not preclude the possibility of exposure or infection with CHIK.
- 4.Positive results in serum indicates ongoing or recent infection by Chikungunya virus causing rash, fever and severe joint pain (arthralgia). The test cannot be used to differentiate between primary and secondary infection.
- 5.Any reactive or non reactive report is recommended to be confirmed with additional test like PCR.
6. False positive results may be seen due to cross reactivity in infection with Epstein -BARR virus, Influenza A & B, Brucella and other mosquito borne diseases like Dengue and Zika Virus infections, in patients with high levels of heterophile antibodies and Rheumatoid factor.
5. False negative results may be due to processing of sample collected early in the course of disease and immunosuppression.
6. Test conducted on serum.

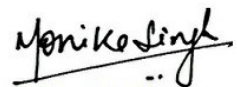
*** End Of Report ***

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.



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MD (Microbiology), MAMS



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Consultant Microbiologist



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.

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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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