

Patient Name : Mr MR.DUMMY
DOB/Age/Gender : 23 Y/Male
Patient ID / UHID : 8053740/RCL7249765
Referred By : Dr. Dr. X
Sample Type : Serum

Sample Collected : Apr 26, 2024, 01:00 PM
Report Date : May 25, 2024, 12:39 PM.
Barcode No : SI484754
Report Status : Final Report

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

Drugs Allergy Panel Test 12 allergens**Paracetamol Allergy Test**

IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Paracetamol EIA	0.66	IU/mL	<0.35

Diclofenac Allergy Test

IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Diclofenac EIA	0.66	IU/mL	<0.35

Ibuprofen Allergy Test

IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Ibuprofen EIA	0.66	IU/mL	<0.35

Penicilloyl G Allergy Test

IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Penicilloyl G EIA	0.66	IU/mL	<0.35

Penicilloyl V Allergy Test

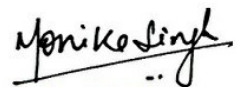
IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Penicilloyl V EIA	0.66	IU/mL	<0.35

Ciprofloxacin Allergy Test

IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Ciprofloxacin Has EIA	0.66	IU/mL	<0.35



Prof. Ashok Rattan
MD (Microbiology), MAMS



Dr. Monika Singh
M.D. (Microbiology)
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.

Patient Name : Mr MR.DUMMY
DOB/Age/Gender : 23 Y/Male
Patient ID / UHID : 8053740/RCL7249765
Referred By : Dr. Dr. X
Sample Type : Serum

Sample Collected : Apr 26, 2024, 01:00 PM
Report Date : May 25, 2024, 12:39 PM.
Barcode No : SI484754
Report Status : Final Report

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

Ampicillin Allergy Test

IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Ampicillin EIA	0.66	IU/mL	<0.35

Amoxicillin Allergy Test

IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Amoxicillin EIA	0.66	IU/mL	<0.35

Gentamycin Allergy Test

IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Gentamycin EIA	0.66	IU/mL	<0.35

Erythromycin Allergy Test

IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Erythromycin EIA	0.66	IU/mL	<0.35

Streptomycin Allergy Test

IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Streptomycin EIA	0.66	IU/mL	<0.35

Rifampicin Allergy Test

IMMUNOGLOBULIN IgE TOTAL SERUM ECLIA	232.9	IU/mL	0-100
Allergy Rifampicin EIA	0.66	IU/mL	<0.35

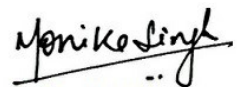
*** End Of Report ***

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.



Prof. Ashok Rattan
MD (Microbiology), MAMS



Dr. Monika Singh
M.D. (Microbiology)
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.

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.