LABORATORY REPORT

Patient Name	:			
DOB/Age/Gender	:	Bill Date	:	
Patient ID / UHID	:	Sample Collected	:	
Referred By	:	Sample Received	:	
Sample Type	:	Report Date	:	
Barcode No	:	Report Status	:	

Test Description	Value(s)	Unit(s)	Reference Range
SEROLOGY AND IMMUNOLOGY REPORT			
Typhidot IgG & IgM, Rapid Card			
TYPHI DOT/ SALMONELLA TYPHI IgG Method : Qualilative immunoassay,rapid card	Negative	-	Negative
TYPHI DOT/ SALMONELLA TYPHI IgM Method : Qualilative immunoassay,rapid card	Negative	-	Negative

Interpretation:

RESULTS	REMARKS
Positive	Indicates presence of IgM & IgG antibodies against Salmonella typhi.
Negative	Indicates absence of IgM & IgG antibodies against Salmonella spp.

Note:

1.Its positivity in serum indicates ongoing or recent infection by Salmonella typhi and the diagnosis should be confirmed by gold standard test such as Blood culture prior to start of antibiotics.

2.IgM antibodies are typically detectable 5-7 days post symptom onset, peaking in 2nd week and frequently remain elevated for 2-4 months following infection.

3.False positive results may be due to cross reactivity with other Salmonella spp., Dengue virus infection & in patients with high levels of Rheumatoid factor.

4. False negative reaction may be due to processing of sample collected early in the course of disease, antibiotic treatment during 1st week and immunosuppression.

5. Test conducted on serum.

Use

To diagnose infection due to Salmonella typhi (Enteric fever)

Dr. Neha Niharika
MD, DNB (Pathology)
Consultant Pathologist

Booking Centre :- WALK-IN REDCLIFFE LABS- TILAK NAGAR, WALK-IN REDCLIFFE LABS- TILAK NAGAR
Processing Lab :- Redcliffe Lifetech Pvt. Ltd., 4B2 Basement & Ground Floor, Tilak Nagar, New Delhi-110018.

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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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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.