

Patient Name	: Mr MR.DUMMY		
DOB/Age/Gender	: 23 Y/Male	Sample Collected	: Apr 26, 2024, 01:00 PM
Patient ID / UHID	: 8052180/RCL7248749	Report Date	: May 24, 2024, 06:22 PM.
Referred By	: Dr. X	Barcode No	: SI484338
Sample Type	: Serum	Report Status	: Final Report
Test Description	Value(s)	Unit(s)	Reference Range

Anti Soluble Liver Antigen (Anti SLA)

ANTI SOLUBLE LIVER ANTIGEN (SLA) (EIA)	11.0	U/mL	<12.00
---	------	------	--------

Interpretation:

RESULT IN U/mL	REMARKS
<12	Negative
12-18	Equivocal
>18	Positive

Comments

Antibodies to Soluble liver antigen (SLA) are specific for Autoimmune hepatitis (AIH) and present in 20% of these patients, many of whom are negative for other autoantibodies. AIH is a chronic progressive liver disease of unknown origin that responds well to immunosuppressive therapy and has a poor prognosis if untreated . Early and accurate diagnosis is hence of great importance. 70% of these patients also show presence of Anti nuclear antibodies, Anti smooth muscle antibodies and Liver Kidney microsomal antibodies which are of diagnostic value for AIH but not specific for the disease since they also occur in 10-15% patients with Viral hepatitis and other immune mediated diseases.

*** 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.



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.


Dr. Dummy

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.

DISCLAIMER

This is a sample report provided for demonstration purposes only and does not represent an actual patient report. Test results, reference ranges, methodologies, instrumentation, and report formats may vary depending on the laboratory performing the test. The format and representation shown are indicative of reports generated by the National Reference Laboratory of Redcliffe Labs, Noida. This sample report should not be used for medical interpretation, diagnosis, or treatment decisions.