

Patient Name : Mr MR.DUMMY

DOB/Age/Gender : 23 Y/Male

Patient ID / UHID : 8052154/RCL7248632

Referred By : Dr. Dr. X

Sample Type : Serum

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

Report Date : May 25, 2024, 02:32 PM.

Barcode No : SI485221

Report Status : Final Report

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

ANCA by IFA

p-ANCA	Negative		Negative
c-ANCA	Negative		Negative

Interpretation:**TEST DESCRIPTION :**

Anti neutrophil cytoplasmic antibody (ANCA) tests are used to diagnose and monitor inflammatory activity in the primary systemic small vessel vasculitides. ANCA in these diseases is best demonstrated by using a combination of Indirect Immunofluorescence of normal peripheral blood neutrophils and ELISAs that detect ANCA specific for proteinase 3 (PR3) or Myeloperoxidase(MPO).

1. c-ANCA is associated with Wegener's granulomatosis and less frequently with microscopic polyangiitis. Approximately 85% of patients with a C-ANCA pattern by IFA have antibodies specific for PR3.

2. Atypical C-ANCA has no clinical significance.

3. p-ANCA is associated with microscopic polyangiitis and focal necrotizing and crescentic glomerulonephritis. Churg-Strauss syndrome is associated with p-ANCA directed against MPO. Approximately 90% of patients with a P-ANCA pattern by IFA have antibodies specific for MPO.

4. Atypical P-ANCA is described with a mixed group of inflammatory diseases e.g sclerosing cholangitis and connective tissue disorders. Antigen targets include lactoferrin, elastase, cathepsin G and BPI.

Patients with a number of other diseases, such as ulcerative colitis and ankylosing spondylitis, will commonly have ANCA as well. However in these cases there is no associated vasculitis and the ANCA are thought to be incidental or epiphenomena rather than part of the disease itself.

TECHNIQUE :

• BIOCHIP slide with Hep 2 cells, primate liver, ethanol fixed granulocytes and formaldehyde fixed granulocytes is used. The technique avoids false positive reporting of ANCA in ANA positive patients.

• Indirect Immunofluorescence, Sample Screening dilution used is 1:10.

REMARK:

Decisions about treatments should not be solely based on results of ANCA tests.

*** 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 :-

 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.

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.