Thyrocare

Tests you can trust

Name : xxxxxxxxxxxxxxxx xxxxxxxxxxxxxxxx

Date : xx xx xxxx

Test Asked : Mmr Immunity Profile Igg Antibody

Report Status: Complete Report

Dummy Report



First National Diagnostic Chain to have **100% of its Labs with NABL Accreditation#**

98% Reports released within **06 Hours** of sample reaching the lab+

9 out of 10 Doctors Trust that Thyrocare reports are **Accurate & Reliable***

1200+ Tests & Profiles

Temperature-Controlled Sample Logistics

Unique Barcode Tracking

Fully Automated Machines Inspected Daily

Abnormal Values Re-Checked Twice

Reports Verified By Expert MD Pathologists Stationed at Every Lab

Your reports are digitally verifiable

Scan the QR code inside the report to check authenticity of reported values

QR code will remain active for 30 days from report release date

Accredited by

NABL From 2005

ISO 9001: 2015 – From 2015

CAP From 2007

PROCESSED AT :
Thyrocare

Thyrocare
Tests you can trust

Thyrocare Technologies Limited, D-37/3, TTC MIDC, Turbhe, Navi Mumbai - 400 703 9870666333 wellness@thyrocare.com

First National Diagnostic Chain to have 100% of its Labs with NABL Accreditation#

NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : MMR IMMUNITY PROFILE IGG ANTIBODY

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

Report Availability Summary

Note: Please refer to the table below for status of your tests.

3 Ready | **0** Ready with Cancellation | **0** Processing | **0** Cancelled in Lab

TEST DETAILS	REPORT STATUS
MMR IMMUNITY PROFILE IGG ANTIBODY	Ready
MEASLES ANTIBODY - IGG	Ready
MUMPS ANTIBODY - IGG	Ready
RUBELLA - IgG	Ready

Dummy Report

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NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : MMR IMMUNITY PROFILE IGG ANTIBODY

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS
MEASLES ANTIBODY - IGG	C.L.I.A	< 6.7	AU/mL

Bio. Ref. Interval. :
Negative : < 13.5
Equivocal : 13.5 - 16.5
Positive : > 16.5

Clinical Significance:
Measles is acute, highly contagious disease. Measles symptoms appear in two stages. In first stage, the individual may have runny nose, cough and a slight fever. The second stage consist of high fever, red blotchy rash.

Specifications:
Intra assay Precision: 8% (%CV), Inter Assay Precision: 15.8% (%CV), Sensitivity: 94.70%, Specificity: 97.40%

Kit Validation References:
TH.TULCHINSKY et.al measles control in developing and developed countries : the case for a two-dose policy. Bull World health organ ,71(1): 93-103(1993)

Method : SANDWICH CHEMI LUMINESCENT IMMUNO ASSAY

Please correlate with clinical conditions.

Dummy Report

Sample Collected on (SCT) : Sample collection time
Sample Received on (SRT) : Sample receiving time at Lab
Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

Doctor 1 Sign Doctor 2 Sign

Scan QR code to verify authenticity of reported results; active for 30 days from release time.

NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : MMR IMMUNITY PROFILE IGG ANTIBODY

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS
MUMPS ANTIBODY - IGG	C.L.I.A	< 4.5	AU/mL

Bio. Ref. Interval. :
Negative : < 9
Equivocal : 9 - 11
Positive : > 11

Clinical Significance:
Mumps is a viral infection caused by paramyxovirus that primarily affects saliva-producing (salivary) glands that are located near ears. Mumps can cause swelling in one or both of those glands. Signs and symptoms of mumps may include:
1. Pain in the swollen salivary glands on one or both sides of your face
2. Pain while chewing or swallowing
It spreads easily from person to person through infected saliva. If individual is not immune, they can contract mumps by breathing in saliva droplets from an infected person who has just sneezed or coughed. Assay aids in detection of IGG antibodies against MUMPS Virus in Patient serum. Presence of these IGG antibody indicates that the person had previous infection or was vaccinated against MUMPS.

Specifications:
Intra- assay : 8.34 % , Inter assay - 9.4 % ; Specificity - 98.2 %, Sensitivity - 98.5 %

Kit validation reference:
C.E. CAPLAN Mumps in the era of vaccines. C.M.A.J., 160 (6) : 865-866 (1999).
Method :

Please correlate with clinical conditions.

Dummy Report

Sample Collected on (SCT) : Sample collection time
Sample Received on (SRT) : Sample receiving time at Lab
Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

Doctor 1 Sign Doctor 2 Sign

PROCESSED AT :
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NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : MMR IMMUNITY PROFILE IGG ANTIBODY

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS
RUBELLA - IgG	C.M.I.A	< 2.5	IU/mL

Bio. Ref. Interval. :
Negative : < 5 || Equivocal : 5 - 10 || Positive : > 10

Method : FULLY AUTOMATED CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY

Please correlate with clinical conditions.

~~ End of report ~~

Dummy Report

Sample Collected on (SCT) : Sample collection time
Sample Received on (SRT) : Sample receiving time at Lab
Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

Doctor 1 Sign Doctor 2 Sign

Scan QR code to verify authenticity of reported results; active for 30days from release time.

CONDITIONS OF REPORTING

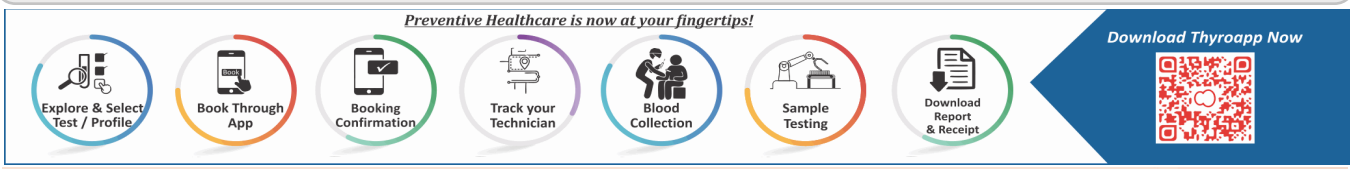
- The reported results are for information and interpretation of the referring doctor only.
- It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- Results of tests may vary from laboratory to laboratory and also in some parameters from time to time for the same patient.
- Should the results indicate an unexpected abnormality, the same should be reconfirmed.
- Only such medical professionals who understand reporting units, reference ranges and limitations of technologies should interpret results.
- This report is not valid for medico-legal purpose.
- Neither Thyrocare, nor its employees/representatives assume: (a) any liability, responsibility for any loss or damage that may be incurred by any person as a result of presuming the meaning or contents of the report, (b) any claims of any nature whatsoever arising from or relating to the performance of the requested tests as well as any claim for indirect, incidental or consequential damages. The total liability, in any case, of Thyrocare shall not exceed the total amount of invoice for the services provided and paid for.
- Thyrocare Discovery video link :- https://youtu.be/nbdYeRgYyQc

EXPLANATIONS

- Majority of the specimen processed in the laboratory are collected by Pathologists and Hospitals we call them as "Clients".
- **Name** - The name is as declared by the client and recored by the personnel who collected the specimen.
- **Ref.Dr** - The name of the doctor who has recommended testing as declared by the client.
- **Labcode** - This is the accession number in our laboratory and it helps us in archiving and retrieving the data.
- **Barcode** - This is the specimen identity number and it states that the results are for the specimen bearing the barcode (irrespective of the name).
- **SCP** - Specimen Collection Point - This is the location where the blood or specimen was collected as declared by the client.
- **SCT** - Specimen Collection Time - The time when specimen was collected as declared by the client.
- **SRT** - Specimen Receiving Time - This time when the specimen reached our laboratory.
- **RRT** - Report Releasing Time - The time when our pathologist has released the values for Reporting.
- **Reference Range** - Means the range of values in which 95% of the normal population would fall.

SUGGESTIONS

- Values out of reference range requires reconfirmation before starting any medical treatment.
- Retesting is needed if you suspect any quality shortcomings.
- Testing or retesting should be done in accredited laboratories.
- For suggestions, complaints, clinical support or feedback, write to us at **customersupport@thyrocare.com** or call us on **022-3090 0000**





*+T&C Apply, #As on 5th December 2024, *As per a survey on doctors' perception of laboratory diagnostics (IJARIIT,2023)*