Thyrocare

Tests you can trust

Name : XXXXXXXXXXXXXXXX XXXXXXXXXXXXXXXX

Date : XX XX XXXX

Test Asked : Torch All 8

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

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NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : TORCH ALL 8

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

Report Availability Summary

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

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

TEST DETAILS	REPORT STATUS
TORCH ALL 8	Ready
HERPES SIMPLEX VIRUS (HSV)-IgG	Ready
HERPES SIMPLEX VIRUS (HSV)-IgM	Ready
RUBELLA - IGG	Ready
RUBELLA - IGM	Ready
CYTOMEGALO VIRUS (CMV) - IgG	Ready
CYTOMEGALO VIRUS (CMV) - IgM	Ready
TOXO GONDII - IgG	Ready
TOXO GONDII - IgM	Ready

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NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : TORCH ALL 8

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS
HERPES SIMPLEX VIRUS (HSV)-IgG	E.L.I.S.A	< 0.4	OD Ratio
Bio. Ref. Interval. : Negative : < 0.80 Equivocal : 0.80 - 1.20 Positive : > 1.20 **Method :** Solid Phase Enzyme Immunoassay			
HERPES SIMPLEX VIRUS (HSV)-IgM	E.L.I.S.A	< 0.4	OD Ratio
Bio. Ref. Interval. : Negative : < 0.80 Equivocal : 0.80 - 1.20 Positive : > 1.20 **Method :** Solid Phase Capture Enzyme Immunoassay			

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 : TORCH ALL 8

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

RUBELLA - IGM	C.M.I.A	< 2.5	AU/mL

Bio. Ref. Interval. :

Negative : < 5
Equivocal : 5-8
Positive : > 8

Method : FULLY AUTOMATED CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY

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

NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : TORCH ALL 8

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS
CYTOMEGALO VIRUS (CMV) - IgG	C.M.I.A	< 5	AU/mL

Bio. Ref. Interval. :
Negative : < 10
Equivocal : 10 - 14
Positive : > 14
Method : FULLY AUTOMATED CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY

CYTOMEGALO VIRUS (CMV) - IgM	C.M.I.A	< 4	AU/mL

Bio. Ref. Interval. :
Negative : < 8
Equivocal : 8 - 12
Positive : > 12
Method : FULLY AUTOMATED CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY

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

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NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : TORCH ALL 8

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS
TOXO GONDII - IgG	C.M.I.A	< 0.41	IU/mL

Bio. Ref. Interval. :
Negative : < 0.81
Equivocal : 0.81 - 1.20
Positive : > 1.20
Method : FULLY AUTOMATED CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY

TOXO GONDII - IgM	C.M.I.A	< 3	AU/mL

Bio. Ref. Interval. :
Negative : < 6
Equivocal : 6 - 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 30 days from release time.

CONDITIONS OF REPORTING

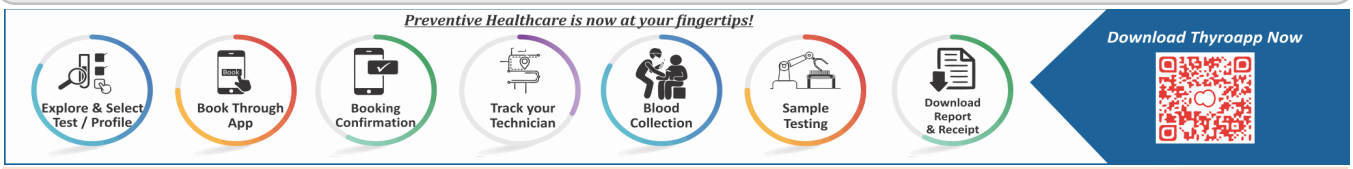
- The reported results are for information and interpretation of the referring doctor only.
- It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- Results of tests may vary from laboratory to laboratory and also in some parameters from time to time for 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)*