Thyrocare

Tests you can trust

Name : XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

Date : xx xx xxxx

Test Asked : Dengue Rapid Test (Igg/Igm)

Report Status: Complete Report



9 out of 10 Doctors trust that Thyrocare **reports are accurate & reliable***

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

Samples Processed in **NABL Accredited** Labs#

700+ Tests & Profiles

Temperature-Controlled Sample Logistics

Unique Barcode Tracking & Reports with QR Code Verification

Fully Automated Machines Inspected Daily

Abnormal Values Re-Checked Twice

Reports Verified By Expert MD Pathologists Stationed at Every Lab

Accredited by

NABL From 2005#

ISO 9001: 2015 - From 2015

CAP From 2007

Dummy Report

PROCESSED AT :

Thyrocare
Tests you can trust

Thyrocare Technologies Limited, D-37/3, TTC MIDC, Turbhe, Navi Mumbai - 400703 98706 66333 wellness@thyrocare.com

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

NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : DENGUE RAPID TEST (IGG/IGM)

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

Report Availability Summary

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

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

TEST DETAILS	REPORT STATUS
DENGUE RAPID TEST (IGG/IGM)	Ready

Dummy Report

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NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : DENGUE RAPID TEST (IGG/IGM)

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE
DENGUE - IGG RAPID TEST	IMMUNOASSAY	NEGATIVE

Bio. Ref. Interval. :-

Positive : Presence of Dengue IgG antibody
Negative : Absence of Dengue IgG antibody

Clinical significance :
1. This is a screening and Qualitative test and a positive report does not confirm diagnosis. All positive cases should be verified by confirmatory methods like ELISA.
2. Presence of IGG dengue antibodies indicates patient is recently exposed to or currently infected with Dengue virus
3. Cross reactivity seen in flavivirus group between dengue virus, yellow fever etc
4. Recommended test is Dengue NS1 antigen in first 5 days of fever and Dengue IgG and IgM after 7-10 days of fever.

Sensetivity : 97.2% ; Specificity : 96.2%

References :
Dengue Guidelines for Diagnosis and treatment, prevention and control. WHO . New edition 2009"

Please correlate with clinical conditions.
Method:- LATERAL FLOW IMMUNOASSAY

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

NAME : XXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : DENGUE RAPID TEST (IGG/IGM)

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE
DENGUE - IGM RAPID TEST	IMMUNOASSAY	NEGATIVE

Bio. Ref. Interval. :-

Positive : Presence of Dengue IgM antibody
Negative : Absence of Dengue IgM antibody

Clinical significance :
1. This is a screening and Qualitative test and a positive report does not confirm diagnosis. All positive cases should be verified by confirmatory methods like ELISA.
2. Presence of IGM dengue antibodies indicates patient is recently exposed to or currently infected with Dengue virus
3. Cross reactivity seen in flavivirus group between dengue virus, yellow fever etc
4. Recommended test is Dengue NS1 antigen in first 5 days of fever and Dengue IgG and IgM after 7-10 days of fever.

Sensetivity : 97.5% ; Specificity : 96.6%

References : Dengue Guidelines for Diagnosis and treatment, prevention and control. WHO . New edition 2009"

Please correlate with clinical conditions.
Method:- LATERAL FLOW IMMUNOASSAY

~~ 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

CONDITIONS OF REPORTING

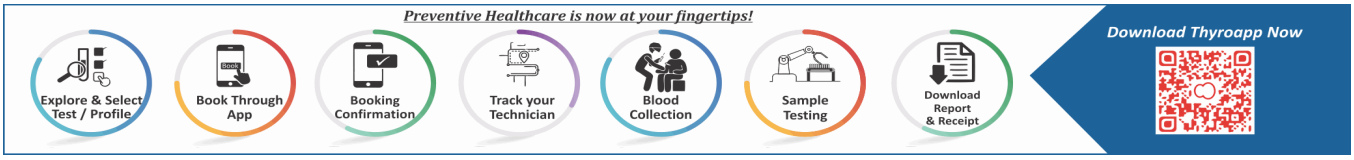
- The reported results are for information and interpretation of the referring doctor only.
- It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- Results of tests may vary from laboratory to laboratory and also in some parameters from time to time for 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 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.
- Thyrocare Discovery video link :- https://youtu.be/nbdYeRgYyQc
- For clinical support please contact @8450950852,8450950853,8450950854 between 10:00 to 18:00

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 or feedback, write to us at **info@thyrocare.com** or call us on **022-3090 0000 / 61 30 00**
- SMS:<Labcode No.>to **9666333**





*+T&C Apply, # Upto 95% Samples in NABL Accredited Labs, * As per a survey on doctors' perception of laboratory diagnostics (IJARIIT,2023)*