

Name : XX

Date : XX XX XXXX

Test Asked : Hcvab

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*

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NAME : XXXXXXXXXXXXXXXXXX

REF. BY : XXXXXXXXXXXXXXXXXX

TEST ASKED : HCVAB

SAMPLE COLLECTED AT : XX

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
HEPATITIS C ANTIBODY(HCVAB) RAPID TEST	Ready ✓

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NAME : XXXXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXXXX
TEST ASKED : HEPATITIS C ANTIBODY(HCVAB) RAPID TEST
SAMPLE COLLECTED AT :
XX

TEST NAME	TECHNOLOGY	VALUE
HEPATITIS C ANTIBODY(HCVAB) RAPID TEST	IMMUNOASSAY	REACTIVE

Reactive : Presence of antibodies to Hepatitis C virus
Non Reactive : Absence of antibodies to Hepatitis C virus

Clinical Significance

1. This is a screening and Qualitative test and a positive report does not confirm diagnosis. All Reactive tests should be confirmed with HCV RNA PCR, as per National guidelines.
2. The test result should be evaluated with other data by the physician.
3. False Reactive results seen in patients receiving mouse monoclonal antibodies and those with heterophile antibodies in serum
4. False negative / non reactive results seen if sample collected during early course of illness, in immunosuppression etc.

Sensitivity : 100%; Specificity : 99.74%

References : National Laboratory guidelines for viral hepatitis Lee SR, Yearwood GD, et al. Evaluation of a rapid, Point of care test device for the diagnosis of Hepatitis C infection. J Clin Virol.2010; 48:15-17.

Please correlate with clinical conditions.

Method:- RAPID IMMUNOCHROMATOGRAPHIC ASSAY

~~ End of 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 / 62 300**
- ✓ SMS: <Labcode No.>to **90666333**

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