Thyrocare

Tests you can trust

Name : XXXXXXXXXX

Date : XXXXXXXXX

Test Asked : Hive

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~

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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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Patient Name : XXXXXXXXXXXXX
Referred By : XXXXXXXXXXXXX
Sample Collected At : XXXXXXXXXXXXX

Tests Done : HIVE

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
HIV I and II	Ready

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Patient Name	: XXXXXXXXXXXXXX	Sample Collected on (SCT)	: XXXXXXXXXXXXX
Referred By	: XXXXXXXXXXXXXX	Sample Received on (SRT)	: XXXXXXXXXXXXX
Sample Collected At	: XXXXXXXXXXXXXX	Report Released on (RRT)	: XXXXXXXXXXXXX
		Sample Type | Barcode	: XXXXXXXXXXXXX

Method 1 : FULLY AUTOMATED CHEMILUMINESCENT MICROPARTICLE IMMUNOASSAY

TEST NAME	TECHNOLOGY	VALUE	UNITS	INTERPRETATION
HIV I and II	C.M.I.A	1	OD ratio	REACTIVE

Interpretation for Method 1 :

NONREACTIVE (Titre value <1):
- Indicates Absence of HIV-1 p24 antigens and antibodies specific to HIV-1 and HIV -2.

REACTIVE (Titre value >= 1)
- Indicates Presence of HIV-1 p24 antigens and antibodies specific to HIV-1 and HIV -2.

Method 2 : RAPID CARD IMMUNOCHROMATOGRAPHY ASSAY

TEST NAME	TECHNOLOGY	INTERPRETATION
HIV 1 RAPID	RAPID CARD	REACTIVE
HIV 2 RAPID	RAPID CARD	REACTIVE

Method 3 : RAPID CARD IMMUNOCONCENTRATION / IMMUNOFILTRATION TECHNIQUE

TEST NAME	TECHNOLOGY	INTERPRETATION
HIV 1 TRIDOT	RAPID CARD	REACTIVE
HIV 2 TRIDOT	RAPID CARD	REACTIVE

Interpretation for Method 2 & Method 3:

NONREACTIVE
- Indicates Absence of Antibodies specific to HIV-1 and/or HIV -2

REACTIVE
- Indicates Presence of Antibodies specific to HIV-1 and/or HIV -2

FINAL RESULT :

TEST NAME	INTERPRETATION
HIV 1	REACTIVE
HIV 2	REACTIVE

Dummy Report

Tests Done : HIV - 1 & II

Report Remarks : Labcode:1612080299/IT001

Doctor 1 Sign

Doctor 2 Sign

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Patient Name	: XXXXXXXXXXXXXX	**Sample Collected on (SCT)**	: XXXXXXXXXXXXX
Referred By	: XXXXXXXXXXXXXX	**Sample Received on (SRT)**	: XXXXXXXXXXXXX
Sample Collected At	: XXXXXXXXXXXXX	**Report Released on (RRT)**	: XXXXXXXXXXXXX
		Sample Type | Barcode	: XXXXXXXXXXXXX

Note :

The result is Reactive/ positive. This has been confirmed by three different assay methods, in accordance with the National HIV testing guidelines recommended by NACO.

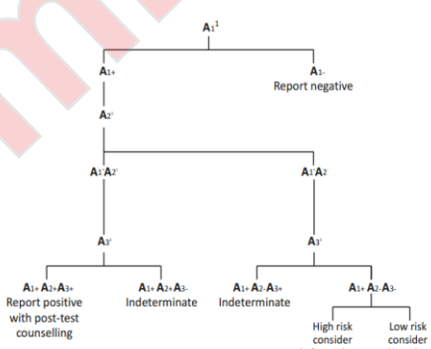
Clinical Significance :

- This is a screening test. Any positive or non-correlating result needs to be confirmed with a confirmatory test, such as Western Blot or HIV RNA PCR testing etc.
- All results should be interpreted in conjunction with the patient's clinical history and other findings.
- HIV reporting follows Strategy 3 of the National Guidelines of HIV Testing by NACO (National AIDS Control Organization), utilizing highly sensitive and specific assays.
- All indeterminate results require retesting with a fresh sample after 14-28 days or confirmation by a confirmatory test.
- Indeterminate cases are reported if one or two of the three assays report reactive results.

Limitations :

- Several conditions can lead to false positive or false negative results.
- Common causes of a false positive result include autoimmune diseases, alcoholic hepatitis, primary biliary cirrhosis, leprosy, multiple pregnancies, and cross-reactive antibodies.
- Common causes of a false negative result include testing during the window period or the end stage of the disease.

FINAL RESULT INTERPRETATION : As per the below Strategy 3 flow diagram of HIV Testing by NACO guidelines.



~~ End of report ~~

Tests Done : HIV - 1 & II

Report Remarks : Labcode:1612080299/IT001

Doctor 1 Sign **Doctor 2 Sign**

Dummy Report

CONDITIONS OF REPORTING

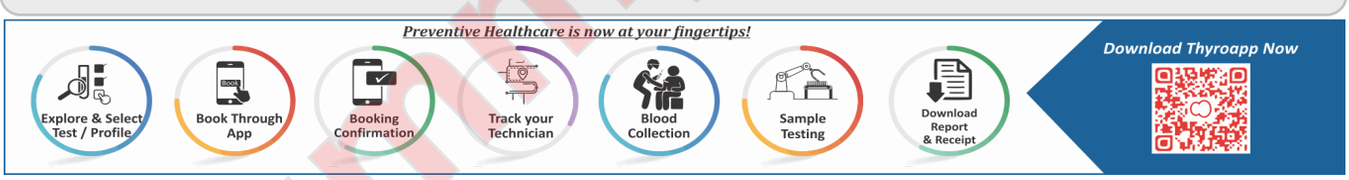
- The reported results are for information and interpretation of the referring doctor only.
- It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- Results of tests may vary from laboratory to laboratory and also in some parameters from time to time for 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 (Applicable for all company owned labs except Bhagalpur & Vijayawada),

⋆ As per survey on doctors' perception of laboratory diagnostics (IJARIIT, 2023), -Mumbai Reference Lab is CAP Accredited

Dummy Report