Thyrocare

Tests you can trust

Name : XXXXXXXXXX

Date : XXXXXXXX

Test Asked : Ttgaa

Report Status: XXXXXXXXXX



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~

Dummy Report

Processed At :

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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Patient Name : **XXXXXXXXXXXXXX**
Referred By : **XXXXXXXXXXXXXX**
Sample Collected At : **XXXXXXXXXXXXXX**

Tests Done : **TTGAA**

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
TISSUE TRANSGLUTAMINASE (TTG) ANTIBODY, IGA (CLIA)	Ready

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Patient Name : **XXXXXXXXXXXXXX**
Referred By : **XXXXXXXXXXXXXX**
Sample Collected At : **XXXXXXXXXXXXXX**

Sample Collected on (SCT) : **XXXXXXXXXXXXXX**
Sample Received on (SRT) : **XXXXXXXXXXXXXX**
Report Released on (RRT) : **XXXXXXXXXXXXXX**
Sample Type | Barcode : **XXXXXXXXXXXXXX**

TEST NAME	TECHNOLOGY	VALUE	UNITS
TISSUE TRANSGLUTAMINASE (TTG) ANTIBODY, IGA (CLIA)	CLIA	10	AU/mL

Bio. Ref. Interval. :-

<8AU/mL: Negative for autoantibodies of IgA class directed against tissue transglutaminase (tTG)
> or=8AU/mL: Positive for autoantibodies of IgA class directed against tissue transglutaminase (tTG)

NOTE
-Negative results do not fully exclude disease, especially in patients with IgA deficiency or in children < 2 years.
-A positive result requires confirmation with additional tests. False positives may occur in Type 1 Diabetes, chronic liver disease, rheumatoid arthritis, psoriatic arthritis.
-Results in range of 7.2 to 8.8 AU/mL are retested and reported only after thorough correlation to rule out assay variability and should be interpreted in conjunction with clinical history
CLINICAL SIGNIFICANCE
-Gluten-sensitive enteropathy or celiac disease is a chronic condition affecting genetically-susceptible children and adults.The disease is HLA-associated, and may strike at any age with peak onset in early childhood.
-Tissue transglutaminase has been identified as the major autoantigen associated with celiac disease.
-Anti-tTG IgA is a highly specific and sensitive marker for Celiac disease.
-Results should always be interpreted together with clinical presentation, dietary history, HLA typing (DQ2/DQ8), Anti EMA, Anti Gliadin and Anti Reticulin Antibodies and histological examination of the duodenal mucosa.
-Diagnosis of a disease should not be established on the basis of a single test result, but should be determined in conjunction with clinical findings and other diagnostic procedures as well as in association with medical judgement. Any therapeutic decision must also be taken on a case-by-case basis.

REFERENCES
1. E. BODIL ROTH et al. Biochemical and immuno-pathological aspects of tissue transglutaminase in coeliac disease. Autoimmunity, 36 : 221-226 (2003).
2. W. DIETERICH et al. Identification of tissue transglutaminase as the autoantigen of celiac disease. Nature Med., 3 : 797-801 (1997)

Please correlate with clinical conditions.
Method:- INDIRECT CHEMILUMINESCENT IMMUNOASSAY (CLIA)

~~ End of report ~~

Dummy Report

Tests Done : **TTGAA**

Doctor 1 Sign **Doctor 2 Sign**

CONDITIONS OF REPORTING

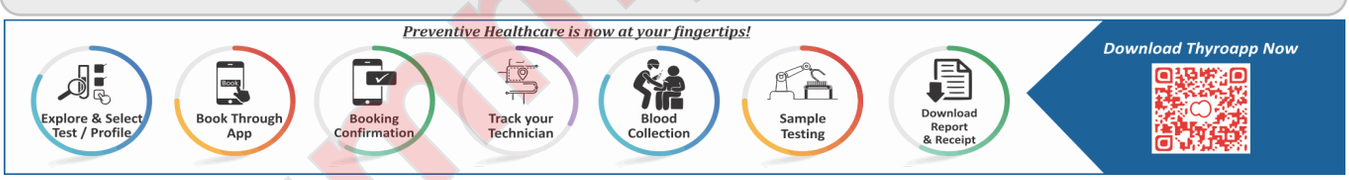
- The reported results are for information and interpretation of the referring doctor only.
- It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- Results of tests may vary from laboratory to laboratory and also in some parameters from time to time for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