Thyrocare

Tests you can trust

Name : XXXXXXXXXXXXXXXXXXXXXXXXX

Date : XXXXXXXX

Test Asked : Allergy Non-vegetarian Panel 2 (by Phadia)

Report Status : Complete Report

Dummy Report



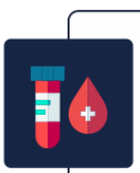
First National Diagnostic Chain to have **100% of its Labs with NABL Accreditation#**



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~

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 : **ALLERGY NON-VEGETARIAN PANEL 2 (BY PHADIA)**

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
ALLERGY NON-VEGETARIAN PANEL 2 (BY PHADIA)	Ready

Dummy Report

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

TEST NAME	TECHNOLOGY	VALUE	UNITS	Bio. Ref. Interval.
ALLERGY SHRIMP (BY PHADIA)	FEIA	0.25	kUA/L	< 0.35
ALLERGY FISH COD (BY PHADIA)	FEIA	0.19	kUA/L	< 0.35
ALLERGY LOBSTER (BY PHADIA)	FEIA	0.21	kUA/L	< 0.35
ALLERGY BLUE MUSSEL (BY PHADIA)	FEIA	0.15	kUA/L	< 0.35
ALLERGY SALMON (BY PHADIA)	FEIA	0.27	kUA/L	< 0.35
ALLERGY TUNA (BY PHADIA)	FEIA	0.22	kUA/L	< 0.35

Please correlate with clinical conditions.

Method :

APSHR - IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD
APFIS - IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD
APLOB - IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD
APBLU - IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD
APSA2 - IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD
APTUN - IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD

~~ End of report ~~

Dummy Report

Tests Done : ALLERGY NON-VEGETARIAN PANEL 2 (BY PHADIA)

Report Remarks : PatientId:A228618041 Labcode:0801001082/IT001

Doctor 1 Sign **Doctor 2 Sign**

CUSTOMER DETAILS

As declared in our data base

Name: XXXXXXXXXXXXXX **Age:** XX **Sex:** XX

Barcodes/Sample_Type	:	XXXXXXXXXXXXXX
Labcode	:	XXXXXXXXXXXXXX
Ref By	:	XXXXXXXXXXXXXX
Sample_Type/Tests	:	SERUM SPL:ALLERGY NON-VEGETARIAN PANEL 2 (BY PHADIA)
Sample Collected At	:	XXXXXXXXXXXXXX
Sample Collected on (SCT)	:	XXXXXXXXXXXXXX
Report Released on (RRT)	:	XXXXXXXXXXXXXX
Amount Collected	:	**XXXXXXXXXXXXX**

Thyrocare,D-37/1,MIDC,Turbhe,Navi Mumbai - 400703. | Phone:022 - 6712 3400 |www.thyrocare.com | info@thyrocare.com

Dummy Report

CONDITIONS OF REPORTING

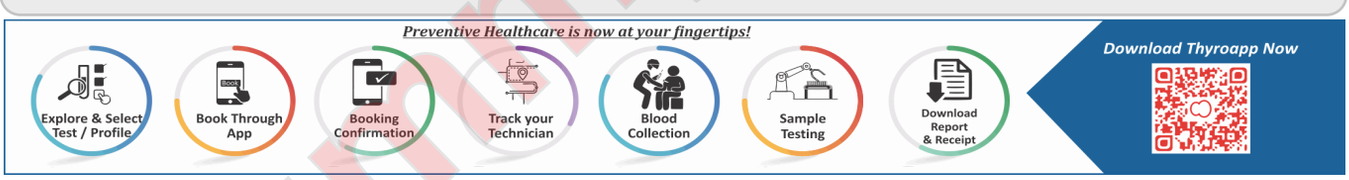
- The reported results are for information and interpretation of the referring doctor only.
- It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- Results of tests may vary from laboratory to laboratory and also in some parameters from time to time for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