Thyrocare

Tests you can trust

Name : XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

Date : XXXXXXXX

Test Asked : Allergy Inhalant Advanced Package (by Phadia)

Report Status : Complete Report

Dummy Report







Thyrocare
Tests you can trust

Thyrocare Technologies Limited, D-37/3, TTC MIDC, Turbhe, Navi Mumbai - 400 703. 9870666333 wellness@thyrocare.com

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

Patient Name : XXXXXXXXXXXX
Referred By : XXXXXXXXXXXX
Address : XXXXXXXXXXXX

Tests Done : **ALLERGY INHALANT ADVANCED PACKAGE (BY PHADIA)**

Report Availability Summary

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

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

TEST DETAILS	REPORT STATUS
ALLERGY INHALANT ADVANCED PACKAGE (BY PHADIA)	Ready
ALLERGY HOUSE DUST (BY PHADIA)	Ready
Allergy House Dust Mite - D.Pteronyssinus (By Phadia)	Ready
Allergy House Dust Mite- D.Farinae (By Phadia)	Ready
ALLERGY COCKROACH GERMAN (BY PHADIA)	Ready
ALLERGY BERMUDA GRASS (BY PHADIA)	Ready
ALLERGY CAT DANDER (BY PHADIA)	Ready
ALLERGY DOG DANDER (BY PHADIA)	Ready

Dummy Report

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Patient Name	: XXXXXXXXXXXX	Tests Done	: ALLERGY INHALANT ADVANCED PACKAGE (BY PHADIA)
Referred By	: XXXXXXXXXXXX		
Address	: XXXXXXXXXXXX		

Tests Outside Reference Range

Note: Please refer to the table below for tests outside reference range.

Test Name	Observed Value	Units	Bio. Ref. Interval.
ALLERGY			
ALLERGY CAT DANDER (BY PHADIA)	**0.44**	kUA/L	< 0.35
ALLERGY DOG DANDER (BY PHADIA)	**2**	kUA/L	< 0.35
ALLERGY HOUSE DUST (BY PHADIA)	**0.44**	kUA/L	< 0.35
Allergy House Dust Mite- D.Farinae (By Phadia)	**1**	kUA/L	< 0.35

Dummy Report

Disclaimer: The above listed is the summary of the parameters with values outside the BRI. For detailed report values, parameter correlation and clinical interpretation, kindly refer to the same in subsequent pages.

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

TEST NAME	TECHNOLOGY	VALUE	UNITS
ALLERGY BERMUDA GRASS (BY PHADIA)	FEIA	0.21	kUA/L

Bio. Ref. Interval. :-

Technology: FEIA

Methodology: ImmunoCAP, Fluorescent Enzyme Immunoassay (FEIA) method

Clinical Significance:
- Allergy test by Phadia (ImmunoCAP Specific IgE) is a test for the quantitative measurement of allergen specific IgE antibodies in human serum and plasma

- This assay gives an indication of the risk of clinical reactions to an allergen and assist in identifying the offending allergens for avoidance measures

- In food allergy, circulating IgE antibodies may remain undetectable despite a convincing clinical history, because the antibodies here may be directed towards the allergens that are revealed or altered during industrial processing, cooking or digestion, and therefore do not exist in the original food against which the patient is tested

- A negative result does not exclude allergy sensitization. Allergy panels include only common causative allergens and do not include all possible allergens. Normal IgE levels do not necessarily exclude the possibility of allergy because certain allergens can be Non IgE mediated.

- Very low levels of allergen specific IgE antibodies must be evaluated with caution when Total IgE (TIGE) values are >1000 kU/L

- The result interpretation should be on an individual case basis, in conjunction with clinical history and other findings

RAST (Radioallergosorbent Test) classification of Allergy is a scale from 0-6 that quantifies the level of specific IgE antibodies to an allergen in the blood. The grading is as follows:
- The results are expressed as Positive or negative, 0.35 (kUA/L) as Analytical cut off
- A positive result indicates specific IgE antibodies to one or more of the allergens in the Immuno CAP allergen mix
- Negative result indicates absence of the IgE antibodies to the components of Allergen mix

Please correlate with clinical conditions.

Method:- IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD

Dummy Report

Tests Done : ALLERGY INHALANT ADVANCED PACKAGE (BY PHADIA)

Report Remarks : Labcode:0904001682/IT001

Doctor 1 Sign

Doctor 2 Sign

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

TEST NAME	TECHNOLOGY	VALUE	UNITS
ALLERGY CAT DANDER (BY PHADIA)	**FEIA**	**0.44**	**kUA/L**

Bio. Ref. Interval. :-

Technology: FEIA

Methodology: ImmunoCAP, Fluorescent Enzyme Immunoassay (FEIA) method

Clinical Significance:
- Allergy test by Phadia (ImmunoCAP Specific IgE) is a test for the quantitative measurement of allergen specific IgE antibodies in human serum and plasma

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- Negative result indicates absence of the IgE antibodies to the components of Allergen mix

Please correlate with clinical conditions.

Method:- IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD

Dummy Report

Tests Done : ALLERGY INHALANT ADVANCED PACKAGE (BY PHADIA)

Report Remarks : Labcode:0904001682/IT001

Doctor 1 Sign

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

TEST NAME	TECHNOLOGY	VALUE	UNITS
ALLERGY COCKROACH GERMAN (BY PHADIA)	FEIA	0.21	kUA/L

Bio. Ref. Interval. :-

Technology: FEIA

Methodology: ImmunoCAP, Fluorescent Enzyme Immunoassay (FEIA) method

Clinical Significance:
- Allergy test by Phadia (ImmunoCAP Specific IgE) is a test for the quantitative measurement of allergen specific IgE antibodies in human serum and plasma

- This assay gives an indication of the risk of clinical reactions to an allergen and assist in identifying the offending allergens for avoidance measures

- In food allergy, circulating IgE antibodies may remain undetectable despite a convincing clinical history, because the antibodies here may be directed towards the allergens that are revealed or altered during industrial processing, cooking or digestion, and therefore do not exist in the original food against which the patient is tested

- A negative result does not exclude allergy sensitization. Allergy panels include only common causative allergens and do not include all possible allergens. Normal IgE levels do not necessarily exclude the possibility of allergy because certain allergens can be Non IgE mediated.

- Very low levels of allergen specific IgE antibodies must be evaluated with caution when Total IgE (TIGE) values are >1000 kU/L

- The result interpretation should be on an individual case basis, in conjunction with clinical history and other findings

RAST (Radioallergosorbent Test) classification of Allergy is a scale from 0-6 that quantifies the level of specific IgE antibodies to an allergen in the blood. The grading is as follows:
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- Negative result indicates absence of the IgE antibodies to the components of Allergen mix

Please correlate with clinical conditions.

Method:- IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD

Dummy Report

Tests Done : ALLERGY INHALANT ADVANCED PACKAGE (BY PHADIA)

Report Remarks : Labcode:0904001682/IT001

Doctor 1 Sign

Doctor 2 Sign

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

TEST NAME	TECHNOLOGY	VALUE	UNITS
ALLERGY DOG DANDER (BY PHADIA)	**FEIA**	**2**	**kUA/L**

Bio. Ref. Interval. :-

Technology: FEIA

Methodology: ImmunoCAP, Fluorescent Enzyme Immunoassay (FEIA) method

Clinical Significance:
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- Negative result indicates absence of the IgE antibodies to the components of Allergen mix

Please correlate with clinical conditions.

Method:- IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD

Dummy Report

Tests Done : ALLERGY INHALANT ADVANCED PACKAGE (BY PHADIA)

Report Remarks : Labcode:0904001682/IT001

Doctor 1 Sign

Doctor 2 Sign

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

TEST NAME	TECHNOLOGY	VALUE	UNITS
ALLERGY HOUSE DUST (BY PHADIA)	**FEIA**	**0.44**	**kUA/L**

Bio. Ref. Interval. :-

Technology: FEIA

Methodology: ImmunoCAP, Fluorescent Enzyme Immunoassay (FEIA) method

Clinical Significance:
- Allergy test by Phadia (ImmunoCAP Specific IgE) is a test for the quantitative measurement of allergen specific IgE antibodies in human serum and plasma

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Method:- IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD

Dummy Report

Tests Done : ALLERGY INHALANT ADVANCED PACKAGE (BY PHADIA)

Report Remarks : Labcode:0904001682/IT001

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Doctor 2 Sign

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Referred By	: XXXXXXXXXXXX	Sample Received on (SRT)	: XXXXXXXXXXXX
Address	: XXXXXXXXXXXX	Report Released on (RRT)	: XXXXXXXXXXXX
		Sample Type | Barcode	: XXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS
Allergy House Dust Mite- D.Farinae (By Phadia)	**FEIA**	**1**	**kUA/L**

Bio. Ref. Interval. :-

Technology: FEIA

Methodology: ImmunoCAP, Fluorescent Enzyme Immunoassay (FEIA) method

Clinical Significance:
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Method:- IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD

Dummy Report

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Report Remarks : Labcode:0904001682/IT001

Doctor 1 Sign

Doctor 2 Sign

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Referred By	: XXXXXXXXXXXX	Sample Received on (SRT)	: XXXXXXXXXXXX
Address	: XXXXXXXXXXXX	Report Released on (RRT)	: XXXXXXXXXXXX
		Sample Type | Barcode	: XXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS
Allergy House Dust Mite - D.Pteronyssinus (By Phadia)	FEIA	0.21	kUA/L

Bio. Ref. Interval. :-

Technology: FEIA

Methodology: ImmunoCAP, Fluorescent Enzyme Immunoassay (FEIA) method

Clinical Significance:
- Allergy test by Phadia (ImmunoCAP Specific IgE) is a test for the quantitative measurement of allergen specific IgE antibodies in human serum and plasma

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Please correlate with clinical conditions.

Method:- IMMUNOCAP, FLUORESCENT ENZYME IMMUNOASSAY (FEIA) METHOD

~~ End of report ~~

Dummy Report

Tests Done : ALLERGY INHALANT ADVANCED PACKAGE (BY PHADIA)

Report Remarks : Labcode:0904001682/IT001

Doctor 1 Sign **Doctor 2 Sign**

CUSTOMER DETAILS

As declared in our data base

Name: XXXXXXXXXXXXXXXX **Age:** XX **Sex:** XX

Barcodes/Sample_Type	:	XXXXXXXXXXXXXXXXXXX
Labcode	:	XXXXXXXXXXXXXXXXXXX
Ref By	:	XXXXXXXXXXXXXXXXXXX
Sample_Type/Tests	:	SERUM SPL:ALLERGY INHALANT ADVANCED PACKAGE (BY PHADIA)
Address	:	XXXXXXXXXXXXXXXXXXX
Sample Collected on (SCT)	:	XXXXXXXXXXXXXXXXXXX
Report Released on (RRT)	:	XXXXXXXXXXXXXXXXXXX
Amount Collected	:	**XXXXXXXXXXXXXXXXXX**

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