# **Thyrocare**

Plot No.265/A,Survey No.4625, T.P No.19,Moje MEM Nagar, Sabarmati,Ahmedabad-380009





Corporate office: Thyrocare Technologies Limited, ♥ D-37/3, TTC MIDC, Turbhe, Navi Mumbai - 400 703

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REPORT

NAME : YATIN KUMAR(33Y/M)

REF. BY : SELF

TEST ASKED : T3,T4,MEDIWHEEL 60+

**HOME COLLECTION:** 

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI ANAND NEAR IRMA BH GANESH CHOKDI

**PATIENTID** : YK20763898

 TEST NAME
 TECHNOLOGY
 VALUE
 UNITS

 CA-125
 C.L.I.A
 9.5
 U/ml

Reference Range :-

Less than 30.2 U/ml

Clinical Significance:

CA-125 is used to monitor therapy during treatment for Ovarian Cancer. CA125 is also to detect or monitor whether there is a recurrence of cancer or malignancy after surgical removal of tumor or radiation therapy or chemotherapy (antineoplastic drugs). This test is sometimes used to follow High-Risk women who have a family history of Ovarian Cancer. CA-125 may normally be increased in early pregnancy and during menstruation. It can also be increased in diseases such as Pelvic Inflammatory Disease or Endometriosis and sometimes in Hepatitis and Cirrhosis of the liver.

Specifications:

Precision: Intra Assay (%CV): 4.3 %, Inter Assay (%CV): 2.5%; Sensitivity: 2.0 U/ml

Kit Validation References:

Mackey SE, Creasman WT. Ovarian Cancer Screening. J. Clin Oncol 1995; 13(3); 783 - 93.

Please correlate with clinical conditions.

Method:- TWO SITE SANDWICH IMMUNOASSAY

Sample Collected on (SCT)
Sample Received on (SRT)

Report Released on (RRT)

Sample Type

Labcode

Barcode

: 26 Dec 2022 08:03

: 26 Dec 2022 15:58

: 26 Dec 2022 17:42

: SERUM

: 2612079273/DS853 Dr Yukti Shah, MD (Path)

: AH107761

(A)

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#### Thyrocare

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REPORT

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**REF. BY**: SELF

TEST ASKED : T3,T4,MEDIWHEEL 60+

PATIENTID : YK20763898

**HOME COLLECTION:** 

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI ANAND NEAR IRMA BH GANESH CHOKDI

TEST NAME TECHNOLOGY VALUE UNITS

RHEUMATOID FACTOR (RF) IMMUNOTURBIDIMETRY < 10 IU/mL

Reference Range: ADULT: <= 18

Clinical Significance:

Rheumatoid factor is an anti IgE autoimmune antibody. There are high concentration of rheumatoid factor in the serum of some disease, especially rheumatoid arthritis patients. It helps to diagnose rheumatism ,systematic lupus erythematosus, chronic hepatitis etc.

Specifications:

Precision %CV:- Intra assay %CV- 1.38%, Inter assay %CV-2.88%, Sensitivity:- 40 IU/mL.

Kit Validation Reference:

Anderson, S.G., Bentzon, M.W., Houba, V. and Kraq, P. Bull. Wld. Hlth. Org. 42: 311-318 (1970).

Method: LATEX ENHANCED IMMUNOTURBIDIMETRY

Please correlate with clinical conditions.

Sample Collected on (SCT) :26 Dec 2022 08:03

Sample Received on (SRT) : 26 Dec 2022 15:58
Report Released on (RRT) : 26 Dec 2022 17:42

Sample Type : SERUM

**Labcode** : 2612079273/DS853

Barcode : AH107761

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**PATIENTID** : YK20763898

TEST NAME TECHNOLOGY VALUE UNITS
25-OH VITAMIN D (TOTAL) C.L.I.A 20.65 ng/ml

Reference Range :-

DEFICIENCY: <20 ng/ml || INSUFFICIENCY: 20-<30 ng/ml SUFFICIENCY: 30-100 ng/ml || TOXICITY: >100 ng/ml

Clinical Significance:

Vitamin D is a fat soluble vitamin that has been known to help the body absorb and retain calcium and phosphorous; both are critical for building bone health. Decrease in vitamin D total levels indicate inadequate exposure of sunlight, dietary deficiency, nephrotic syndrome. Increase in vitamin D total levels indicate Vitamin D intoxication.

Specifications: Precision: Intra assay (%CV):5.3%, Inter assay (%CV):11.9%; Sensitivity:3.2 ng/ml.

Kit Validation Reference: Holick MF. Vitamin D Deficiency. N Engl J Med. 2007;357:266-81.

# Please correlate with clinical conditions.

Method:- Fully Automated Chemi Luminescent Immuno Assay

 Sample Collected on (SCT)
 : 26 Dec 2022 08:03

 Sample Received on (SRT)
 : 26 Dec 2022 15:58

 Report Released on (RRT)
 : 26 Dec 2022 17:42

Sample Type . SERUM

Labcode 2612079273/DS853 Dr Yukti Shah, MD (Path)

**Barcode** : AH107761 Page : 3 of 18

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NAME : YATIN KUMAR(33Y/M)

: SELF REF. BY

: T3,T4,MEDIWHEEL 60+ **TEST ASKED** 

**HOME COLLECTION:** 

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI ANAND NEAR IRMA BH GANESH CHOKDI

**PATIENTID** : YK20763898

**TEST NAME TECHNOLOGY** VALUE UNITS **HIGH SENSITIVITY C-REACTIVE PROTEIN (HS-CRP) IMMUNOTURBIDIMETRY** 4.49 mg/L

Reference Range :-

< 1.00 - Low Risk 1.00 - 3.00 - Average Risk >3.00 - 10.00 - High Risk

> 10.00 - Possibly due to Non-Cardiac Inflammation

Disclaimer: Persistent unexplained elevation of HSCRP >10 should be evaluated for non-cardiovascular etiologies such as infection , active arthritis or concurrent illness.

# Clinical significance:

High sensitivity C- reactive Protein ( HSCRP) can be used as an independent risk marker for the identification of Individuals at risk for future cardiovascular Disease. A coronary artery disease risk assessment should be based on the average of two hs-CRP tests, ideally taken two weeks apart.

# Kit Validation Reference:

- 1. Clinical management of laboratory date in medical practice 2003-3004, 207(2003).
- 2.Tietz: Textbook of Clinical Chemistry and Molecular diagnostics: Second edition: Chapter 47:Page no.1507-1508.

Please correlate with clinical conditions.

Method:- FULLY AUTOMATED LATEX AGGLUTINATION - BECKMAN COULTER

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Sample Type

. SERUM : 2612079273/DS853 Dr Yukti Shah, MD (Path) Labcode

Barcode : AH107761 Page: 4 of 18

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HOME COLLECTION:

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI ANAND NEAR IRMA BH GANESH CHOKDI

**PATIENTID** : YK20763898

TEST NAME TECHNOLOGY VALUE UNITS
VITAMIN B-12 C.L.I.A 186 pg/ml

Reference Range :-

Normal: 211 - 911 pg/ml

# Clinical significance:

Vitamin B12 or cyanocobalamin, is a complex corrinoid compound found exclusively from animal dietary sources, such as meat, eggs and milk. It is critical in normal DNA synthesis, which in turn affects erythrocyte maturation and in the formation of myelin sheath. Vitamin-B12 is used to find out neurological abnormalities and impaired DNA synthesis associated with macrocytic anemias. For diagnostic purpose, results should always be assessed in conjunction with the patients medical history, clinical examination and other findings.

Specifications: Intra assay (%CV):5.0%, Inter assay (%CV):9.2 %;Sensitivity:45 pg/ml

## Kit Validation reference:

Chen IW, Sperling MI, Heminger LA. Vitamin B12. In: Pesce AJ, Kaplan LA, eds. Methods in Clinical Chemistry. St. Louis: CV Mosby; 1987:569–73.

# Please correlate with clinical conditions.

Method:- COMPETITIVE CHEMI LUMINESCENT IMMUNO ASSAY

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REPORT

NAME : YATIN KUMAR(33Y/M)

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TEST ASKED : T3,T4,MEDIWHEEL 60+

**HOME COLLECTION:** 

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI ANAND NEAR IRMA BH GANESH CHOKDI

**PATIENTID** : YK20763898

TEST NAMETECHNOLOGYVALUEUNITSPROSTATE SPECIFIC ANTIGEN (PSA)C.L.I.A0.81ng/ml

Reference Range :-

Normal: < 4.00 ng/ml

Border line: 4.01 to 10.00 ng/ml

Clinical Significance:

Elevated levels of PSA are associated with prostate cancer, but may also be seen with prostatitis (Inflammation of the prostate) and benign prostatic hyperplasia (BPH). PSA test done along with free PSA provides additional information. Studies have suggested that the percentage of free PSA in total PSA is lower in patients with prostate cancer than those with benign prostate hyperplasia.

Specification:

Precision: Intra assay (%CV): 4.38%, Inter assay (%CV): 4.67%; Sensitivity: 0.01 ng/ml

Kit validation references:

Wang MC, Valenzuala LA, Murphy GP, and Chu TM. Purification of a human prostate-specific antigen. Invest. Urol. 1979; 17: 159

Please correlate with clinical conditions.

Method:- TWO SITE SANDWICH IMMUNOASSAY

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TEST ASKED : T3,T4,MEDIWHEEL 60+

R(33Y/M)

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI

ANAND NEAR IRMA BH GANESH CHOKDI

**HOME COLLECTION:** 

PATIENTID: YK20763898

TEST NAME	TECHNOLOGY	VALUE	UNITS	NORMAL RANGE
TOTAL CHOLESTEROL	PHOTOMETRY	216	mg/dl	< 200
HDL CHOLESTEROL - DIRECT	PHOTOMETRY	37	mg/dl	40-60
LDL CHOLESTEROL - DIRECT	PHOTOMETRY	160	mg/dl	< 100
TRIGLYCERIDES	PHOTOMETRY	150	mg/dl	< 150
TC/ HDL CHOLESTEROL RATIO	CALCULATED	5.9	Ratio	3 - 5
TRIG / HDL RATIO	CALCULATED	4.07	Ratio	< 3.12
LDL / HDL RATIO	CALCULATED	4.3	Ratio	1.5-3.5
HDL / LDL RATIO	CALCULATED	0.23	Ratio	> 0.40
NON-HDL CHOLESTEROL	CALCULATED	179.41	mg/dl	< 160
VLDL CHOLESTEROL	CALCULATED	30.06	mg/dl	5 - 40

## Please correlate with clinical conditions.

## Method:

CHOL - Cholesterol Oxidase, Esterase, Peroxidase

HCHO - Direct Enzymatic Colorimetric

LDL - Direct Measure

TRIG - Enzymatic, End Point

TC/H - Derived from serum Cholesterol and Hdl values

TRI/H - Derived from TRIG and HDL Values

LDL/ - Derived from serum HDL and LDL Values

HD/LD - Derived from HDL and LDL values.

NHDL - Derived from serum Cholesterol and HDL values

VLDL - Derived from serum Triglyceride values

## \*REFERENCE RANGES AS PER NCEP ATP III GUIDELINES:

TOTAL CHOLESTEROL	(mg/dl)	HDL	(mg/dl)	LDL	(mg/dl)	TRIGLYCERIDES	(mg/dl)
DESIRABLE	<200	LOW	<40	OPTIMAL	<100	NORMAL	<150
BORDERLINE HIGH	200-239	HIGH	>60	NEAR OPTIMAL	100-129	BORDERLINE HIGH	150-199
HIGH	>240			BORDERLINE HIGH	130-159	HIGH	200-499
				HIGH	160-189	VERY HIGH	>500
				VERY HIGH	>190		

Alert !!! 10-12 hours fasting is mandatory for lipid parameters. If not, values might fluctuate.

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Barcode : AH107761

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REPORT

NAME : YATIN KUMAR(33Y/M)

REF. BY : SELF

**TEST ASKED** : T3,T4,MEDIWHEEL 60+ **HOME COLLECTION:** 

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI

ANAND NEAR IRMA BH GANESH CHOKDI

**PATIENTID** : YK20763898

TEST NAME	TECHNOLOGY	VALUE	UNITS	NORMAL RANGE
ALKALINE PHOSPHATASE	PHOTOMETRY	72.94	U/L	45 - 129
BILIRUBIN - TOTAL	PHOTOMETRY	0.39	mg/dl	0.3-1.2
BILIRUBIN -DIRECT	PHOTOMETRY	0.05	mg/dl	< 0.3
BILIRUBIN (INDIRECT)	CALCULATED	0.34	mg/dl	0-0.9
GAMMA GLUTAMYL TRANSFERASE (GGT)	PHOTOMETRY	45.57	U/I	< 55
ASPARTATE AMINOTRANSFERASE (SGOT )	PHOTOMETRY	38.45	U/I	< 35
ALANINE TRANSAMINASE (SGPT)	PHOTOMETRY	67.65	U/I	< 45
SGOT / SGPT RATIO	CALCULATED	0.57	Ratio	< 2
PROTEIN - TOTAL	PHOTOMETRY	6.77	gm/dl	5.7-8.2
ALBUMIN - SERUM	PHOTOMETRY	3.78	gm/dl	3.2-4.8
SERUM GLOBULIN	CALCULATED	2.99	gm/dL	2.5-3.4
SERUM ALB/GLOBULIN RATIO	CALCULATED	1.26	Ratio	0.9 - 2

# Please correlate with clinical conditions.

# Method:

ALKP - Modified IFCC method

BILT - Vanadate Oxidation

**BILD** - Vanadate Oxidation

BILI - Derived from serum Total and Direct Bilirubin values

GGT - Modified IFCC method

SGOT - IFCC\* Without Pyridoxal Phosphate Activation

SGPT - IFCC\* Without Pyridoxal Phosphate Activation

OT/PT - Derived from SGOT and SGPT values.

PROT - Biuret Method

SALB - Albumin Bcg1method (Colorimetric Assay Endpoint)

SEGB - DERIVED FROM SERUM ALBUMIN AND PROTEIN VALUES

A/GR - Derived from serum Albumin and Protein values

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Dr Yukti Shah, MD (Path) Labcode : 2612079273/DS853

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**REF. BY**: SELF

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PATIENTID : YK20763898

**HOME COLLECTION:** 

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI ANAND NEAR IRMA BH GANESH CHOKDI

TEST NAME	TECHNOLOGY	VALUE	UNITS
PHOSPHOROUS	PHOTOMETRY	3.72	mg/dL
Reference Range :			

Adults: 2.4 - 5.1 mg/dL

## Clinical Significance:

In plasma and serum the majority of phosphate exists in the inorganic form (Pi), approximately 15% bound to protein and the remainder in complexes and free forms. Serum phosphate concentrations are dependent on diet and variation in the secretion of hormones such as Parathyroid Hormone (PTH).

Specifications:

Precision %CV :- Intra assay %CV- 1.55% , Inter assay %CV-2.99% , Sensitivity:-0.10 mmol/L

Kit Validation Reference:

Young DS. Effects of drugs on clinical laboratory tests, 5th ed. AACC Press, 2000.

Method: UNREDUCED PHOSPHOMOLYBDATE METHOD

Please correlate with clinical conditions.

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REPORT

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**REF. BY**: SELF

TEST ASKED : T3,T4,MEDIWHEEL 60+

PATIENTID : YK20763898

**HOME COLLECTION:** 

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI ANAND NEAR IRMA BH GANESH CHOKDI

Reference Range: ADULTS: 3.5-5.1 MMOL/L

# Clinical Significance:

An abnormal increase in potassium (hyperkalemia)can profoundly affect the nervous system and increase the chance of irregular heartbeats (arrhythmias), which ,when extreme ,can be fatal. The assay could be affected mildly and may result in anomalous values if serum samples have heterophilic antibodies, hemolyzed , icteric or lipemic. The concentration of Potassium in a given specimen may vary due to differences in assay methods, calibration and reagent specificity.

Method: ION SELECTIVE ELECTRODE

CHLORIDE I.S.E 103.6 mmol/l

Reference Range: ADULTS: 98-107 MMOL/L

# Clinical Significance:

An increased level of blood chloride (called hyperchloremia) usually indicates dehydration, but can also occur with other problems that cause high blood sodium, such as Cushing syndrome or kidney disease. Hyperchloremia also occurs when too much base is lost from the body (producing metabolic acidosis) or when a person hyperventilates (causing respiratory alkalosis). A decreased level of blood chloride (called hypochloremia) occurs with any disorder that causes low blood sodium. Hypochloremia also occurs with congestive heart failure, prolonged vomiting or gastric suction, Addison disease, emphysema or other chronic lung diseases (causing respiratory acidosis), and with loss of acid from the body (called metabolic alkalosis).

Method: ION SELECTIVE ELECTRODE

Please correlate with clinical conditions.

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**PATIENTID** 

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# REPORT

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REF. BY : SELF

TEST ASKED : T3,T4,MEDIWHEEL 60+

: YK20763898

....

**HOME COLLECTION:** 

4.1

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI

mg/dl

4.2 - 7.3

ANAND NEAR IRMA BH GANESH CHOKDI

**TEST NAME TECHNOLOGY VALUE** NORMAL RANGE UNITS BLOOD UREA NITROGEN (BUN) **PHOTOMETRY** 11.05 7 - 25 mg/dl CREATININE - SERUM **PHOTOMETRY** 0.66 0.6-1.1 mg/dl BUN / SR.CREATININE RATIO CALCULATED 16.74 9:1-23:1 Ratio Adult: 17-43 UREA (CALCULATED) CALCULATED 23.65 mg/dL UREA / SR.CREATININE RATIO **CALCULATED** 35.83 Ratio < 52 **CALCIUM PHOTOMETRY** 9.38 mg/dl 8.8-10.6 SODIUM I.S.E 139.1 mmol/I 136 - 145

**PHOTOMETRY** 

## Please correlate with clinical conditions.

# Method:

**URIC ACID** 

BUN - Kinetic UV Assay.

SCRE - Creatinine Enzymatic method

B/CR - Derived from serum Bun and Creatinine values

UREAC - Derived from BUN Value.

UR/CR - Derived from UREA and Sr.Creatinine values.

CALC - Arsenazo III Method, End Point.

SOD - ION SELECTIVE ELECTRODE

URIC - Uricase / Peroxidase Method

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REPORT

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HOME COLLECTION:
2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI

ANAND NEAR IRMA BH GANESH CHOKDI

**PATIENTID** : YK20763898

**TEST NAME TECHNOLOGY VALUE** UNITS REFERENCE RANGE C.L.I.A 130 60-200 TOTAL TRIIODOTHYRONINE (T3) ng/dl C.L.I.A **TOTAL THYROXINE (T4)** 11.9 µg/dl 4.5-12 0.3-5.5 C.L.I.A 1.45 μIU/ml THYROID STIMULATING HORMONE (TSH)

Comments: SUGGESTING THYRONORMALCY

Please correlate with clinical conditions.

#### Method:

T3 - Competitive Chemi Luminescent Immuno Assay

T4 - Competitive Chemi Luminescent Immuno Assay

TSH - SANDWICH CHEMI LUMINESCENT IMMUNO ASSAY

# Disclaimer:

Results should always be interpreted using the reference range provided by the laboratory that performed the test. Different laboratories do tests using different technologies, methods and using different reagents which may cause difference In reference ranges and hence it is recommended to interpret result with assay specific reference ranges provided in the reports. To diagnose and monitor therapy doses, it is recommended to get tested every time at the same Laboratory.

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REPORT

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REF. BY : SELF

TEST ASKED : T3,T4,MEDIWHEEL 60+

**HOME COLLECTION:** 

2 SMRURI APARTMENT NEAR IRMA BH GANESH

CHOKDI ANAND NEAR IRMA BH GANESH CHOKDI

**PATIENTID** : YK20763898

TEST NAMETECHNOLOGYVALUEUNITSEST. GLOMERULAR FILTRATION RATE (eGFR)CALCULATED127mL/min/1.73 m2

Reference Range :-

> = 90 : Normal 60 - 89 : Mild Decrease

45 - 59 : Mild to Moderate Decrease 30 - 44 : Moderate to Severe Decrease

15 - 29 : Severe Decrease

# Clinical Significance

The normal serum creatinine reference interval does not necessarily reflect a normal GFR for a patient. Because mild and moderate kidney injury is poorly inferred from serum creatinine alone. Thus, it is recommended for clinical laboratories to routinely estimate glomerular filtration rate (eGFR), a "gold standard" measurement for assessment of renal function, and report the value when serum creatinine is measured for patients 18 and older, when appropriate and feasible. It cannot be measured easily in clinical practice, instead, GFR is estimated from equations using serum creatinine, age, race and sex. This provides easy to interpret information for the doctor and patient on the degree of renal impairment since it approximately equates to the percentage of kidney function remaining. Application of CKD-EPI equation together with the other diagnostic tools in renal medicine will further improve the detection and management of patients with CKD.

## Reference

Levey AS, Stevens LA, Schmid CH, Zhang YL, Castro AF, 3rd, Feldman HI, et al. A new equation to estimate glomerular filtration rate. Ann Intern Med. 2009;150(9):604-12.

Please correlate with clinical conditions.

Method:- CKD-EPI Creatinine Equation

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Sample Type . SERUM

Report Released on (RRT)

Labcode 2612079273/DS853 Dr Yukti Shah, MD (Path)

: 26 Dec 2022 17:42

**Barcode** : AH107761 Page : 13 of 18

# **Thyrocare**

Plot No.265/A,Survey No.4625, T.P No.19,Moje MEM Nagar, Sabarmati,Ahmedabad-380009





Corporate office: Thyrocare Technologies Limited, ♥ D-37/3, TTC MIDC, Turbhe, Navi Mumbai - 400 703

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wellness@thyrocare.com 
www.thyrocare.com

REPORT

NAME : YATIN KUMAR(33Y/M)

REF. BY : SELF

TEST ASKED : T3,T4,MEDIWHEEL 60+

HOME COLLECTION :

2 SMRURI APARTMENT NEAR IRMA BH GANESH

CHOKDI ANAND NEAR IRMA BH GANESH CHOKDI

PATIENTID : YK20763898

TEST NAMETECHNOLOGYVALUEUNITSFASTING BLOOD SUGAR(GLUCOSE)PHOTOMETRY91.38mg/dL

# Reference Range:-

As per ADA Guide	line: Fasting Plasma Glucose (FPG)
Normal	70 to 100 mg/dl
Prediabetes	100 mg/dl to 125 mg/dl
Diabetes	126 mg/dl or higher

# Note:

The assay could be affected mildly and may result in anomalous values if serum samples have heterophilic antibodies, hemolyzed, icteric or lipemic. The concentration of Glucose in a given specimen may vary due to differences in assay methods, calibration and reagent specificity. For diagnostic purposes results should always be assessed in conjunction with patients medical history, clinical findings and other findings.

Please correlate with clinical conditions.

Method:- GOD-PAP METHOD

Sample Collected on (SCT) : 26 Dec 2022 08:03
Sample Received on (SRT) : 26 Dec 2022 15:58

Report Released on (RRT) : 26 Dec 2022 16:50
Sample Type : FLUORIDE

Labcode : 2612079267/DS853 Dr Yukti Shah, MD (Path)

**Barcode** : AO208264 Page : 14 of 18

#### Thyrocare

Plot No.265/A,Survey No.4625, T.P No.19,Moje MEM Nagar, Sabarmati,Ahmedabad-380009





REPORT

NAME : YATIN KUMAR(33Y/M)

**REF. BY**: SELF

**TEST ASKED**: T3,T4,MEDIWHEEL 60+

PATIENTID : YK20763898

**HOME COLLECTION:** 

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI ANAND NEAR IRMA BH GANESH CHOKDI

TEST NAME TECHNOLOGY VALUE UNITS
HbA1c - (HPLC)

(111 20)

H.P.L.C

5.7

%

Reference Range:

Reference Range: As per ADA Guidelines

Below 5.7% : Normal 5.7% - 6.4% : Prediabetic >=6.5% : Diabetic Guidance For Known Diabetics

Below 6.5% : Good Control 6.5% - 7% : Fair Control

7.0% - 8% : Unsatisfactory Control

>8% : Poor Control

Method: Fully Automated H.P.L.C. using Biorad Variant II Turbo

AVERAGE BLOOD GLUCOSE (ABG)

**CALCULATED** 

117

mg/dl

Reference Range :

90 - 120 mg/dl : Good Control 121 - 150 mg/dl : Fair Control

151 - 180 mg/dl: Unsatisfactory Control

> 180 mg/dl : Poor Control

Method: Derived from HBA1c values

Please correlate with clinical conditions.

Sample Collected on (SCT) :26 Dec 2022 08:03

Sample Received on (SRT) : 26 Dec 2022 15:57
Report Released on (RRT) : 26 Dec 2022 16:37

Sample Type : EDTA

**Labcode** : 2612079220/DS853

Barcode : AO208269

g

Dr Yukti Shah, MD (Path)

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# PROCESSED AT: **Thyrocare**

Plot No.265/A, Survey No.4625, T.P No.19, Moje MEM Nagar, Sabarmati, Ahmedabad-380009





: YATIN KUMAR(33Y/M) NAME

**REF. BY** : SELF

: T3,T4,MEDIWHEEL 60+ **TEST ASKED** 

**PATIENTID** : YK20763898

# **HOME COLLECTION:**

2 SMRURI APARTMENT NEAR IRMA BH GANESH

CHOKDI ANAND NEAR IRMA BH GANESH

CHOKDI

TEST NAME	VALUE	UNITS	REFERENCE RANGE
TOTAL LEUCOCYTES COUNT (WBC)	6.88	X 10 <sup>3</sup> / μL	4.0-10.0
NEUTROPHILS	58.4	%	40-80
LYMPHOCYTE PERCENTAGE	33.1	%	20-40
MONOCYTES	4.1	%	0-10
EOSINOPHILS	3.8	%	0.0-6.0
BASOPHILS	0.3	%	<2
IMMATURE GRANULOCYTE PERCENTAGE(IG%)	0.3	%	0-0.5
NEUTROPHILS - ABSOLUTE COUNT	4.02	$X~10^3$ / $\mu L$	2.0-7.0
LYMPHOCYTES - ABSOLUTE COUNT	2.28	$X~10^3$ / $\mu L$	1.0-3.0
MONOCYTES - ABSOLUTE COUNT	0.28	$X~10^3$ / $\mu L$	0.2-1
BASOPHILS - ABSOLUTE COUNT	0.02	$X~10^3$ / $\mu L$	0-0.1
EOSINOPHILS - ABSOLUTE COUNT	0.26	$X~10^3$ / $\mu L$	0-0.5
IMMATURE GRANULOCYTES(IG)	0.02	$X~10^3$ / $\mu L$	0-0.3
TOTAL RBC	5.99	X 10^6/μL	4.5-5.5
NUCLEATED RED BLOOD CELLS	Nil	$X~10^3$ / $\mu L$	<0.01
NUCLEATED RED BLOOD CELLS %	Nil	%	<0.01
HEMOGLOBIN	15.4	g/dL	13-17
HEMATOCRIT(PCV)	49	%	40-50
MEAN CORPUSCULAR VOLUME(MCV)	81.8	fL	83-101
MEAN CORPUSCULAR HEMOGLOBIN(MCH)	25.7	pq	27-32
MEAN CORP.HEMO.CONC(MCHC)	31.4	g/dL	31.5-34.5
RED CELL DISTRIBUTION WIDTH - SD(RDW-SD)	38.5	fL	39-46
RED CELL DISTRIBUTION WIDTH (RDW-CV)	13	%	11.6-14
PLATELET DISTRIBUTION WIDTH(PDW)	10	fL	9.6-15.2
MEAN PLATELET VOLUME(MPV)	9.4	fL	6.5-12
PLATELET COUNT	383	$X~10^3$ / $\mu L$	150-400
PLATELET TO LARGE CELL RATIO(PLCR)	20.1	%	19.7-42.4
PLATELETCRIT(PCT)	0.36	%	0.19-0.39

Please Correlate with clinical conditions.

Method: Fully automated bidirectional analyser (6 Part Differential SYSMEX XN-1000)

(This device performs hematology analyses according to the Hydrodynamic Focussing (DC method), Flow Cytometry Method (using a semiconductor laser), and SLS- hemoglobin method)

Sample Collected on (SCT) .26 Dec 2022 08:03 26 Dec 2022 15:57 Sample Received on (SRT)

26 Dec 2022 16:37 Report Released on (RRT)

**Sample Type** . EDTA

: 2612079220/DS853 Dr Yukti Shah, MD (Path) Labcode

**Barcode** : AO208269 Page: 16 of 18

# PROCESSED AT: **Thyrocare**

Plot No.265/A, Survey No.4625, T.P No.19, Moje MEM Nagar, Sabarmati, Ahmedabad-380009





**PATIENTID** 

**CRYSTALS BACTERIA** 

**PARASITE** 

**YEAST** 

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NAME : YATIN KUMAR(33Y/M)

**REF. BY** : SELF

**TEST ASKED** : T3,T4,MEDIWHEEL 60+

: YK20763898

**HOME COLLECTION:** 

2 SMRURI APARTMENT NEAR IRMA BH GANESH CHOKDI ANAND NEAR IRMA BH GANESH CHOKDI

Absent

Absent Absent

Absent

**OBSERVATION REFERENCE RANGE TEST NAME** UNITS **COMPLETE URINOGRAM VOLUME** >=5 mL PALE YELLOW Pale Yellow COLOUR HAZY Clear **APPEARANCE** SPECIFIC GRAVITY > 1.030 1.003-1.030 6 5 - 8 URINARY PROTEIN **ABSENT** mg/dl Absent **URINARY GLUCOSE ABSENT** Absent mg/dl **ABSENT** Absent URINE KETONE mg/dl **ABSENT** URINARY BILIRUBIN mg/dl Absent **UROBILINOGEN** < 0.2 mg/dl <=0.2 **BILE SALT ABSENT** Absent ABSENT Absent BILE PIGMENT **URINE BLOOD ABSENT** Cells/ul\* Absent **NITRITE ABSENT** Absent < 20 **MICROALBUMIN** 10 mg/l **MUCUS ABSENT** Absent **RED BLOOD CELLS ABSENT** Cells/ul\* Absent URINARY LEUCOCYTES (PUS CELLS) **ABSENT** Cells/ul\* Absent 0-4 **EPITHELIAL CELLS** 1-2 Absent **CASTS** ABSENT

Please correlate with clinical conditions.

Method: Fully Automated FUS2000-3 Urinalysis Dipstick Method, Microscopy

~~ End of report ~~

**ABSENT** 

**ABSENT** 

**ABSENT ABSENT** 

Sample Collected on (SCT)

Sample Received on (SRT)

Report Released on (RRT)

Sample Type Labcode

**Barcode** 

: 26 Dec 2022 08:03

: 26 Dec 2022 16:05

: 26 Dec 2022 17:08

: URINE

: 2612079664/DS853

: Z2143964

Dr Yukti Shah, MD (Path)

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<sup>\*</sup> To Obtain Counts in Cells / HPF Divide the Cells / ul by 5

## 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 :- <a href="https://youtu.be/nbdYeRgYyQc">https://youtu.be/nbdYeRgYyQc</a>
- For clinical support please contact @8450950852,8450950853,8450950854 between 10:00 to 18:00

## **EXPLANATIONS**

- v Majority of the specimen processed in the laboratory are collected by Pathologists and Hospitals we call them as "Clients".
- v **Name** The name is as declared by the client and recored by the personnel who collected the specimen.
- v 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).
- v SCP Specimen Collection Point This is the location where the blood or specimen was collected as declared by the client.
- v SCT Specimen Collection Time The time when specimen was collected as declared by the client.
- v 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 / 6712 3400
- SMS:<Labcode No.> to **9870666333**





**BOOK A TEST** 

Give a Call - 022 4128 2828

WhatsApp **TGS** to 8104112632

Email to tgs@focustb.com

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