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REPORT

NAME : XXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXX
TEST ASKED : ELEMENTS 22 (TOXIC AND NUTRIENTS),HbA1c,HEMOGRAM
PATIENTID :

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS	NORMAL RANGE
ARSENIC	ICP-MS	2.54	µg/L	< 5
CADMIUM	ICP-MS	1.06	µg/L	< 1.5
MERCURY	ICP-MS	2.6	µg/L	< 5
LEAD	ICP-MS	130.38	µg/L	< 150
CHROMIUM	ICP-MS	15.53	µg/L	< 30
BARIUM	ICP-MS	5.03	µg/L	< 30
COBALT	ICP-MS	0.3	µg/L	0.10 - 1.50
CAESIUM	ICP-MS	1.46	µg/L	< 5
THALLIUM	ICP-MS	0.04	µg/L	< 1
URANIUM	ICP-MS	0.04	µg/L	< 1
STRONTIUM	ICP-MS	22.19	µg/L	8 - 38
ANTIMONY	ICP-MS	7.65	µg/L	0.10 - 18
TIN	ICP-MS	1.46	µg/L	< 2
MOLYBDENUM	ICP-MS	2.7	µg/L	0.70 - 4.0
SILVER	ICP-MS	3.53	µg/L	< 4
VANADIUM	ICP-MS	0.71	µg/L	< 0.8
BERYLLIUM	**ICP-MS**	**0.03**	**µg/L**	**0.10 - 0.80**
BISMUTH	ICP-MS	0.13	µg/L	0.10 - 0.80
SELENIUM	ICP-MS	206.7	µg/L	60 - 340
ALUMINIUM	ICP-MS	10.15	µg/L	< 30
NICKEL	ICP-MS	8.07	µg/L	< 15
MANGANESE	ICP-MS	19.22	µg/L	7.10 - 20

Please correlate with clinical conditions.

Method :

ICP - MASS SPECTROMETRY
Note:Reference range has been obtained after considering 95% population as cutoff.

Dummy Report

Sample Collected on (SCT) : Sample collection time
Sample Received on (SRT) : Sample receiving time at Lab
Report Released on (RRT) : Report release time
Sample Type : EDTA
Labcode :
Barcode :

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REPORT

NAME : XXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : ELEMENTS 22 (TOXIC AND NUTRIENTS),HbA1c,HEMOGRAM
PATIENTID :

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS
HbA1c - (HPLC)			
	H.P.L.C	**6.7**	**%**

Reference Range :

Reference Range: As per ADA Guidelines	Guidance For Known Diabetics
Below 5.7% : Normal 5.7% - 6.4% : Prediabetic >=6.5% : Diabetic	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 method

AVERAGE BLOOD GLUCOSE (ABG)	**CALCULATED**	**146**	**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.

Dummy Report

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Report Released on (RRT) : Report release time
Sample Type : EDTA
Labcode :
Barcode :

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REPORT

NAME : XXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXX
TEST ASKED : ELEMENTS 22 (TOXIC AND NUTRIENTS),HbA1c,HEMOGRAM
PATIENTID :

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	VALUE	UNITS	REFERENCE RANGE
TOTAL LEUCOCYTES COUNT (WBC)	6.71	X 10³ / µL	4.0 - 10.0
NEUTROPHILS	60.6	%	40-80
LYMPHOCYTE PERCENTAGE	33.2	%	20-40
MONOCYTES	**1.9**	**%**	**2-10**
EOSINOPHILS	3.6	%	1-6
BASOPHILS	0.4	%	0-2
IMMATURE GRANULOCYTE PERCENTAGE(IG%)	0.3	%	0.0-0.4
NEUTROPHILS - ABSOLUTE COUNT	4.07	X 10³ / µL	2.0-7.0
LYMPHOCYTES - ABSOLUTE COUNT	2.23	X 10³ / µL	1.0-3.0
MONOCYTES - ABSOLUTE COUNT	**0.13**	**X 10³ / µL**	**0.2 - 1.0**
BASOPHILS - ABSOLUTE COUNT	0.03	X 10³ / µL	0.02 - 0.1
EOSINOPHILS - ABSOLUTE COUNT	0.24	X 10³ / µL	0.02 - 0.5
IMMATURE GRANULOCYTES(IG)	0.02	X 10³ / µL	0.0-0.3
TOTAL RBC	4.28	X 10^6/µL	3.8-4.8
NUCLEATED RED BLOOD CELLS	0.01	X 10³ / µL	<0.01
NUCLEATED RED BLOOD CELLS %	0.01	%	<0.01
HEMOGLOBIN	12.1	g/dL	12.0-15.0
HEMATOCRIT(PCV)	38.7	%	36.0-46.0
MEAN CORPUSCULAR VOLUME(MCV)	90.4	fL	83.0-101.0
MEAN CORPUSCULAR HEMOGLOBIN(MCH)	28.3	pq	27.0-32.0
MEAN CORP.HEMO.CONC(MCHC)	**31.3**	**g/dL**	**31.5-34.5**
RED CELL DISTRIBUTION WIDTH - SD(RDW-SD)	44.8	fL	39.0-46.0
RED CELL DISTRIBUTION WIDTH (RDW-CV)	13.6	%	11.6-14.0
PLATELET DISTRIBUTION WIDTH(PDW)	12.1	fL	9.6-15.2
MEAN PLATELET VOLUME(MPV)	10.4	fL	6.5-12
PLATELET COUNT	233	X 10³ / µL	150-410
PLATELET TO LARGE CELL RATIO(PLCR)	28.7	%	19.7-42.4
PLATELETCRIT(PCT)	0.24	%	0.19-0.39

Remarks : Alert!!! Predominantly normocytic normochromic with ovalocytes. Platelets:Appear adequate in smear.

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)

Dummy Report

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Labcode :
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REPORT

NAME : XXXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
ALPHA-1-ANTITRYPSIN (AAT)	IMMUNOTURBIDIMETRY	140.15	mg/dL

Reference Range :-

90 - 200 mg/dL

Clinical Significance ;
Alpha-1-antitrypsin (AAT) is a serine protease inhibitor and the third most abundant protein in circulation. AAT levels can increase 3 to 5 folds in states of systemic inflammation or infection. Increased levels are also seen in Covid-19 patients. Usually, the increase in AAT is directly proportional to the increase in IL6, supporting an anti inflammatory function. It is also considered that patients with Alpha-1-antitrypsin deficiency (AATD) might be at increased risk of SARS-CoV-2 infection and development of severe COVID-19.

Alpha-1 antitrypsin (AAT) deficiency can cause various medical complications of the lungs, liver, and skin, including COPD, emphysema, chronic bronchitis, cirrhosis of the liver and other side effects of the excess enzyme.

Specifications;
Precision (%CV) : Intra-assay %CV: 1.4 %, Inter-assay %CV: 1.36%

Kit Validation reference ;
1.Janciauskiene SM, Bals R, Koczulla R, et al. The discovery of Alpha1-antitrypsin and its role in health and disease. Respir Med, 2011, 105(8) : 1129-1139
2.Sun Z, Yang P. Role of imbalance between neutrophil elastase and alpha1-antitrypsin in cancer development and progression. Lancet Oncol, 2004, 5(3) : 182-190

Please correlate with clinical conditions.
Method:- Polyethylene glycol(PEG)-enhanced immunoturbidimetric

Dummy Report

Sample Collected on (SCT) : Sample collection time
Sample Received on (SRT) : Sample receiving time at Lab
Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

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REPORT

NAME : XXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
HOMOCYSTEINE	PHOTOMETRY	16.02	µmol/L

Reference Range :-

Normal: < 30 µmol/L

Clinical Significance:
Homocysteine is linked to increased risk of premature coronary artery disease, stroke and thromboembolism. Moreover, alzheimers disease, osteoporosis, venous thrombosis, schizophrenia, cognitive deficiency and pregnancy complications also elevates Homocysteine levels.

High Values:
Elevated homocysteine levels might be due to increasing age, genetic traits, drugs, renal dysfunction and dietary deficiency of vitamins or smoking. To lower your homocysteine, eat more green vegetables, stop smoking, alcohol. Folic acid helps lowering elevated levels.

Specifications:
Precision %CV :- Intra assay %CV- 4.5 % , Inter assay %CV-5.87 % Sensitivity : 0.4 umol/L

Kit Validation Reference:
Eikelboom JW, et al Ann Intern Med 131 : 363-75 (1999)

Please correlate with clinical conditions.
Method:- ENZYMATIC ASSAY

Dummy Report

Sample Collected on (SCT) : Sample collection time
Sample Received on (SRT) : Sample receiving time at Lab
Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

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REPORT

NAME : XXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
CYSTATIN C	IMMUNOTURBIDIMETRY	0.98	mg/L

Reference Range :-

<= 60 years: <= 1.03 mg/L
> 60 years : < 1.50 mg/L

Clinical significance

Cystatin c, is a small 13-kda protein and is a member of the cysteine proteinase inhibitor family, it is produced at a constant rate by all nucleated cells. Due to its small size it is freely filtered by the glomerulus and is not secreted but is fully reabsorbed and broken down by the renal tubules. This means that the primary determinate of blood Cystatin c levels is the rate at which it is filtered at the glomerulus making it an excellent gfr marker. Cystatin c is also a marker of inflammation and like many other markers of inflammation; its serum concentration may be higher in patients with decreased renal clearance. There is mounting evidence, however, that Cystatin c may be a predictor of adverse outcomes independent of renal function with its higher sensitivity to detect a reduced GFR than Creatinine determination, also in the so-called "Creatinine-blind" range. Thus, Cystatin c is suggested to be a better marker for GFR than the ubiquitous serum Creatinine.

Reference

1. Barrett aj, Davies me, Grubb a. the place of human gamma-trace (Cystatin c) among the cysteine proteinase inhibitors. Biochem biophys res common 1984; 120: 631-6.

2. Grubb a. diagnostic value of analysis of Cystatin c and protein HC in biological fluids. Clin Nephrol 1992; 38: S20-7.

Please correlate with clinical conditions.
Method:- LATEX ENHANCED IMMUNOTURBIDIMETRY

Dummy Report

Sample Collected on (SCT) : Sample collection time
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Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

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REPORT

NAME : XXXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
25-OH VITAMIN D (TOTAL)	E.C.L.I.A	30.7	ng/mL

Reference Range :
Deficiency : <=20 ng/ml || Insufficiency : 21-29 ng/ml
Sufficiency : >= 30 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):9.20%, Inter assay (%CV):8.50%
Kit Validation Reference : Holick M. Vtamin D the underappreciated D-Lightful hormone that is important for Skeletal and cellular health Curr Opin Endocrinol Diabetes 2002:9(1)87-98.
Method : Fully Automated Electrochemiluminescence Compititive Immunoassay

VITAMIN B-12	E.C.L.I.A	377	pg/mL

Reference Range :
Normal: 197-771 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):2.6%, Inter assay (%CV):2.3 %

Kit Validation Reference : Thomas L.Clinical laborator Diagnostics : Use and Assessment of Clinical laboratory Results 1st Edition,TH Books-Verl-Ges,1998:424-431
Method : Fully Automated Electrochemiluminescence Compititive Immunoassay

Please correlate with clinical conditions.

Dummy Report

Sample Collected on (SCT) : Sample collection time
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Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

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REPORT

NAME : XXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
APOLIPOPROTEIN - A1 (APO-A1)	IMMUNOTURBIDIMETRY	110	mg/dL

Reference Range :
Male : 86 - 152
Female : 94 - 162
Method : FULLY AUTOMATED RATE IMMUNOTURBIDIMETRY – BECKMAN COULTER

APOLIPOPROTEIN - B (APO-B)	IMMUNOTURBIDIMETRY	121	mg/dL

Reference Range :
Male : 56 - 145
Female : 53 - 138
Method : FULLY AUTOMATED RATE IMMUNOTURBIDIMETRY – BECKMAN COULTER

APO B / APO A1 RATIO (APO B/A1)	CALCULATED	1.1	Ratio

Reference Range :
Male : 0.40 - 1.26
Female : 0.38 - 1.14
Method : DERIVED FROM SERUM APO A1 AND APO B VALUES

Please correlate with clinical conditions.

Dummy Report

Sample Collected on (SCT) : Sample collection time
Sample Received on (SRT) : Sample receiving time at Lab
Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

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REPORT

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REF. BY : XXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
HIGH SENSITIVITY C-REACTIVE PROTEIN (HS-CRP)	**IMMUNOTURBIDIMETRY**	**7.8**	**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

Dummy Report

Sample Collected on (SCT) : Sample collection time
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Report Released on (RRT) : Report release time
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Labcode :
Barcode :

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REPORT

NAME : XXXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
FRUCTOSAMINE	PHOTOMETRY	214.8	µmol/L

Reference Range :

Normal < 286 µmol/L

Clinical Significance:

Fructosamine assay is useful in monitoring the degree of glycemia over short-to-intermediate time frames (1-3 weeks) concentration greater than the established normal range is an indication of prolonged hyperglycemia of 1-3 weeks or longer. The higher fructosamine value, poorer is the degree of glycemia control.

Specifications:

Precision %CV : Intra assay %CV- 3.2% , Inter assay %CV-4.0%, Sensitivity:- 290 umol/L

Kit Validation Reference:

Howey JEA, Browning MCK, Fraser CG. Assay of serum fructosamine that minimizes standardization and matrix problems: Use to assess components of biological va-riation. Clin Chem 1987; 33: 269- 272.

Method : NITROBLUE TETRAZOLIUM ASSAY (NBT)

BLOOD KETONE (D3HB)	PHOTOMETRY	0.47	mg/dL

Reference Range :
0.21-2.81 mg/dL

Clinical Significance:
Three types of ketones can be produced in body D-3- Hydroxybutyrate, Acetoacetate and Acetone. D-3- Hydroxybutyrate accounts for approximately 75% of the ketone bodies. During periods of ketosis, D-3- Hydroxybutyrate increases more than the other two. It has been shown to be a better index of ketoacidosis. In diabetics, D-3- Hydroxybutyrate is needed for the assessment of the severity of diabetic coma and to calculate insulin requirements.

Speficcation:
Precision: Intra assay (%CV): 4.53, Inter assay (%CV): 2.9, Sensitivity: 10.41 mg/dL.

Kit validation references:
Mcmurray, C.H., Blanchflower, W.J., Rice, D.A., ClinChem., 1984;30:No.3.

Method : ENZYMATIC (KINETIC)

Please correlate with clinical conditions.

Sample Collected on (SCT) : Sample collection time
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Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

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TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
Lipoprotein (a) [Lp(a)]	IMMUNOTURBIDIMETRY	29.8	mg/dL

Reference Range :-

Adults : < 30.0 mg/dl

Clinical Significance:
Determination of LPA may be useful to guide management of individuals with a family history of CHD or with existing disease. The levels of LPA in the blood depends on genetic factors; The range of variation in a population is relatively large and hence for diagnostic purpose, results should always be assessed in conjunction with the patient's medical history, clinical examination and other findings.

Specifications:
Precision %CV :- Intra assay %CV- 4.55% , Inter assay %CV-0.86 %

Kit Validation Reference:
Tietz NW,Clinical Guide to Laboratory Tests Philadelphia WB. Saunders 1995 : 442-444

Please correlate with clinical conditions.
Method:- LATEX ENHANCED IMMUNOTURBIDIMETRY

Dummy Report

Sample Collected on (SCT) : Sample collection time
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Labcode :
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REPORT

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TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
SERUM COPPER	**PHOTOMETRY**	**155.7**	**µg/dL**

Reference Range :-

Male : 63.5 - 150
Female : 80 - 155

Clinical significance:
Copper is an important trace element and a component of numerous enzymes and proteins involved in energy production, connective tissue formation, melanin synthesis, iron metabolism, development of central nervous system, angiogenesis as well as an antioxidant. Deficiency can cause- Malnourishment, cardiovascular disease, anemia & neuropathy, toxicity may be manifested as acute renal failure, gastroenteritis & chronic liver disease.

Specifications:
Precision: Intra assay (%CV): 1.17, Inter assay (%CV): 2.32.

Kit validation references:
Thomas L. Clinical Laboratory Diagnostics. 1st ed. Frankfurt: TH-Books Verlagsgesellschaft; 1998. p. 337-8

Please correlate with clinical conditions.
Method:- 3,5-DIBR-PAESA

Dummy Report

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Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

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REPORT

NAME : XXXXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
SERUM ZINC	**PHOTOMETRY**	**47.8**	**µg/dL**

Reference Range :-

52 - 286

Clinical Significance:
Zinc is one of the essential trace elements in the body. Its metalloenzymes play a key rple in protein and nucleic acid synthesis, gene expression, wound healing, as an antioxidant, etc. Deficiency can cause- Poor wound healing, gastroenteritis, impaired spermatogenesis, Alzheimer's disease, etc. Toxicity may be manifested as pancreatitis, gastric ulcer, anemia, pulmonary fibrosis.

Specifications:
Precision: Intra assay (%CV): 2.02, Inter assay (%CV): 2.22.

Kit Validation References:
Thomas L. Clinical Laboratory Diagnostics. 1st ed. Frankfurt: TH-Books Verlagsgesellschaft; 1998. p. 347-9

Please correlate with clinical conditions.
Method:- NITRO - PAPS

Dummy Report

Sample Collected on (SCT) : Sample collection time
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Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

Doctor 1 Sign Doctor 2 Sign

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022 - 3090 0000 / 6712 3400 9870666333 wellness@thyrocare.com www.thyrocare.com

REPORT

NAME : XXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
TESTOSTERONE	E.C.L.I.A	23.3	ng/dL

Reference Range :-

Males
20-49 yrs: 249-836 ng/dL
>=50 yrs : 193-740 ng/dL
Females
20-49 yrs : 8.4 - 48.1 ng/dL
>=50 yrs : 2.9 - 40.8 ng/dL

Clinical Significance: Clinical evaluation of serum testosterone, along with serum LH, assists in evaluation of Hypogonadal males. Major causes of lowered testosterone in males include Hypogonadotropic hypogonadism, testicular failure Hyperprolactinema, Hypopituitarism some types of liver and kidney diseases and critical illness.

Specifications: Precision: Intra assay (%CV): 11.50 %, Inter assay (%CV): 5.70%; Sensitivity: 7 ng/dL.

Kit Validation Reference: Wilson JD Foster DW (Eds) Williams Textbook of Endocrinology 8th Edition WB Saunders Piladelphia Pennsylvania .

Please correlate with clinical conditions.
Method:- Fully Automated Electrochemiluminescence Compitive Immunoassay

Dummy Report

Sample Collected on (SCT) : Sample collection time
Sample Received on (SRT) : Sample receiving time at Lab
Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

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REPORT

NAME : XXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
AMYLASE	PHOTOMETRY	64.9	U/L

Reference Range :-

Adults : 28-100 U/L

Interpretation:
Lipemic Sera (Hypertriglyceridemia) may contain inhibitors, Which falsely depress results. About 20% of patients with Acute Pancreatitis have abnormal lipids. Normal serum amylase may occur in Pancreatitis, Especially relapsing and chronic pancreatitis. Moderate increases may be reported in normal pregnancy.

Clinical Significance:
Causes of high Serum Amylase include Acute Pancreatitis, Pancreatic Pseudocyst, Pancreatic Ascites, Pancreatic Abscess, Neoplasm in or adjacent to Pancreas, Trauma to Pancreas, and common Duct Stones. Nonpancreatic Causes include inflammatory salivary lesions (Eg, Mumps), Perforated Peptic Ulcer, Intestinal Obstruction, Biliary Tract Disease, Peritonitis, Acute Appendicitis, Diabetic Ketoacidosis, and Extrapancreatic Carcinomas. Amylase levels more than 25-fold the upper limit of normal are often found when metastatic tumors produce Ectopic Amylase.

Specifications:
Precision: Intra assay (%CV): 2.82, Inter assay (%CV): 2.49, Sensitivity: 10.9 U/L.

Kit Validation References:
Rauscher, E., et coll., Fresenius Z. Analyt. Chem. 324 (1986) 304-305.

Please correlate with clinical conditions.
Method:- ENZYMATIC COLORIMETRIC TEST

Dummy Report

Sample Collected on (SCT) : Sample collection time
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Labcode :
Barcode :

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REPORT

NAME : XXXXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
IRON	PHOTOMETRY	65.4	µg/dL
Reference Range : Male : 65 - 175 Female : 50 - 170 **Method :** FERROZINE METHOD WITHOUT DEPROTEINIZATION			
TOTAL IRON BINDING CAPACITY (TIBC)	PHOTOMETRY	396	µg/dL
Reference Range : Male: 225 - 535 µg/dl Female: 215 - 535 µg/dl **Method :** SPECTROPHOTOMETRIC ASSAY			
% TRANSFERRIN SATURATION	CALCULATED	16.52	%
Reference Range : 13 - 45 **Method :** DERIVED FROM IRON AND TIBC VALUES			
UNSAT.IRON-BINDING CAPACITY(UIBC)	PHOTOMETRY	330.6	µg/dL
Reference Range : 162 - 368 **Method :** SPECTROPHOTOMETRIC ASSAY			

Please correlate with clinical conditions.

Dummy Report

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Sample Type : SERUM
Labcode :
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REPORT

NAME : XXXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
LIPASE	PHOTOMETRY	34.9	U/L

Reference Range :-

Adults : 5.6 - 51.3 U/L

Interpretation:

For diagnostic purposes, the results should always be assessed in conjunction with the patient's medical history, clinical examination and other findings like serum amylase. Serum Lipase is usually normal in patients with elevated serum amylase, having peptic ulcer, salivary adenitis, inflammatory bowel disease, intestinal obstruction, and macroamylasemia. Lipemic sera may interfere with results.

Clinical Significance:
High serum Lipase is a specific marker for pancreatitis; after acute pancreatitis the Lipase activity increases within 4-8 hours, reaches a peak after 24 hours and decreases after 8 to 14 days. However, there is no correlation between the Lipase activity determined in serum and the extent of damage to the pancreas.

Specifications:
Precision: Intra assay (%CV): 3.35, Inter assay (%CV): 2.46, Sensitivity: 3.5 U/L.

Kit Validation References:
Tietz Nw Et Al. Lipase In Serum - The Elusive Enzyme: An Overview. Clin Chem 1993; 39:746-756.

Please correlate with clinical conditions.
Method:- ENZYMATIC COLORIMETRIC ASSAY

Dummy Report

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REPORT

NAME : XXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

PATIENTID :

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS	NORMAL RANGE
TOTAL CHOLESTEROL	**PHOTOMETRY**	**210**	**mg/dL**	**< 200**
HDL CHOLESTEROL - DIRECT	**PHOTOMETRY**	**39**	**mg/dL**	**40-60**
HDL / LDL RATIO	**CALCULATED**	**0.25**	**Ratio**	**> 0.40**
LDL CHOLESTEROL - DIRECT	**PHOTOMETRY**	**160**	**mg/dL**	**< 100**
TRIG / HDL RATIO	CALCULATED	2.87	Ratio	< 3.12
TRIGLYCERIDES	PHOTOMETRY	113	mg/dL	< 150
TC/ HDL CHOLESTEROL RATIO	**CALCULATED**	**5.3**	**Ratio**	**3 - 5**
LDL / HDL RATIO	**CALCULATED**	**4.1**	**Ratio**	**1.5-3.5**
VLDL CHOLESTEROL	CALCULATED	22.56	mg/dL	5 - 40
NON-HDL CHOLESTEROL	**CALCULATED**	**170.5**	**mg/dL**	**< 160**

Please correlate with clinical conditions.

Method :

CHOL - Cholesterol Oxidase, Esterase, Peroxidase
HCHO - Direct Enzymatic Colorimetric
HD/LD - Derived from HDL and LDL values.
LDL - Direct Measure
TRI/H - Derived from TRIG and HDL Values
TRIG - Enzymatic, End Point
TC/H - DERIVED FROM SERUM CHOLESTEROL AND HDL VALUES
LDL/ - Derived from serum HDL and LDL Values
VLDL - DERIVED FROM SERUM TRIGLYCERIDE VALUES
NHDL - Derived from serum Cholesterol and HDL 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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Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

Doctor 1 Sign Doctor 2 Sign

Dummy Report

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REPORT

NAME : XXXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

PATIENTID :

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS	NORMAL RANGE
ALKALINE PHOSPHATASE	PHOTOMETRY	70.7	U/L	45-129
BILIRUBIN - TOTAL	PHOTOMETRY	0.76	mg/dL	0.3-1.2
BILIRUBIN -DIRECT	PHOTOMETRY	0.14	mg/dL	< 0.3
BILIRUBIN (INDIRECT)	CALCULATED	0.52	mg/dL	0-0.9
GAMMA GLUTAMYL TRANSFERASE (GGT)	PHOTOMETRY	20.5	U/L	< 38
SGOT / SGPT RATIO	CALCULATED	0.88	Ratio	< 2
ASPARTATE AMINOTRANSFERASE (SGOT)	PHOTOMETRY	25.5	U/L	< 31
ALANINE TRANSAMINASE (SGPT)	PHOTOMETRY	28.9	U/L	< 34
PROTEIN - TOTAL	PHOTOMETRY	6.79	gm/dL	5.7-8.2
ALBUMIN - SERUM	PHOTOMETRY	3.9	gm/dL	3.2-4.8
SERUM GLOBULIN	CALCULATED	2.89	gm/dL	2.5-3.4
SERUM ALB/GLOBULIN RATIO	CALCULATED	1.35	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
OT/PT - Derived from SGOT and SGPT values.
SGOT - IFCC* WITHOUT PYRIDOXAL PHOSPHATE ACTIVATION
SGPT - IFCC* WITHOUT PYRIDOXAL PHOSPHATE ACTIVATION
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

Dummy Report

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REPORT

NAME : XXXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

PATIENTID :

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

TEST NAME	TECHNOLOGY	VALUE	UNITS	NORMAL RANGE
CALCIUM	PHOTOMETRY	9.3	mg/dL	8.8-10.6
URIC ACID	PHOTOMETRY	6.07	mg/dL	3.2 - 6.1
UREA (CALCULATED)	CALCULATED	33.32	mg/dL	Adult : 17-43
BLOOD UREA NITROGEN (BUN)	PHOTOMETRY	15.57	mg/dL	7.94 - 20.07
UREA / SR.CREATININE RATIO	CALCULATED	37.44	Ratio	< 52
CREATININE - SERUM	PHOTOMETRY	0.89	mg/dL	0.55-1.02
BUN / SR.CREATININE RATIO	CALCULATED	17.49	Ratio	9:1-23:1
SODIUM	I.S.E	138	mmol/L	136 - 145
CHLORIDE	I.S.E	102	mmol/L	98 - 107

Please correlate with clinical conditions.

Method :

CALC - ARSENAZO III METHOD, END POINT.
URIC - Uricase / Peroxidase Method
UREAC - Derived from BUN Value.
BUN - KINETIC UV ASSAY.
UR/CR - Derived from UREA and Sr.Creatinine values.
SCRE - CREATININE ENZYMATIC METHOD
B/CR - DERIVED FROM SERUM BUN AND CREATININE VALUES
SOD - ION SELECTIVE ELECTRODE
CHL - ION SELECTIVE ELECTRODE

Dummy Report

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REPORT

NAME : XXXXXXXXXXXXXXXXXXXX

REF. BY : XXXXXXXXXXXXXXXXXXXX

TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS	REFERENCE RANGE
TOTAL TRIIODOTHYRONINE (T3)	E.C.L.I.A	100	ng/dL	80-200
TOTAL THYROXINE (T4)	E.C.L.I.A	8.7	µg/dL	4.8-12.7
TSH - ULTRASENSITIVE	E.C.L.I.A	1.3	µIU/mL	0.54-5.30

Comments : ***

The Biological Reference Ranges is specific to the age group. Kindly correlate clinically.

Method :

T3 - Fully Automated Electrochemiluminescence Compititive Immunoassay
T4 - Fully Automated Electrochemiluminescence Compititive Immunoassay
USTSH - Fully Automated Electrochemiluminescence Sandwich Immunoassay

Pregnancy reference ranges for TSH/USTSH :

Trimester || T3 (ng/dl) || T4 (µg/dl) || TSH/USTSH (µIU/ml)
1st || 83.9-196.6 || 4.4-11.5 || 0.1-2.5
2nd || 86.1-217.4 || 4.9-12.2 || 0.2-3.0
3rd || 79.9-186 || 5.1-13.2 || 0.3-3.5

References :

1. Carol Devilia, C I Parhon. First Trimester Pregnancy ranges for Serum TSH and Thyroid Tumor reclassified as Benign. Acta Endocrinol. 2016; 12(2) : 242 - 243
2. Kulhari K, Negi R, Kalra DK et al. Establishing Trimester specific Reference ranges for thyroid hormones in Indian women with normal pregnancy : New light through old window. Indian Journal of Contemporary medical research. 2019; 6(4)

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.

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REPORT

NAME : XXXXXXXXXXXXXXXXXXXX
REF. BY : XXXXXXXXXXXXXXXXXXXX
TEST ASKED : AAROGYAM 1.4 WITH UTSH

SAMPLE COLLECTED AT :
XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX

PATIENTID :

TEST NAME	TECHNOLOGY	VALUE	UNITS
EST. GLOMERULAR FILTRATION RATE (eGFR)	**CALCULATED**	**81**	**mL/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

~~ End of report ~~

Dummy Report

Sample Collected on (SCT) : Sample collection time
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Report Released on (RRT) : Report release time
Sample Type : SERUM
Labcode :
Barcode :

Doctor 1 Sign Doctor 2 Sign

CUSTOMER DETAILS

As declared in our data base

Name: XXXXXXXXXXXXX **Age:** XXX **Sex:** XX **Customer Mobile No:** XXXXXXXXXX

Barcodes/Sample_Type	:	
Labcode	:	
Ref By	:	XXXXXXXXXX
Sample_Type/Tests	:	EDTA:HBA , HEMOGRAM - 6 PART (DIFF) , ELEMENTS 22 (TOXIC AND NUTRIENTS) SERUM:AAROGYAM 1.4 WITH UTSH
Sample Collected At	:	
Sample Collected on (SCT)	:	Sample collection time
Report Released on (RRT)	:	Report release time

Thyrocare

Dummy Report

CONDITIONS OF REPORTING

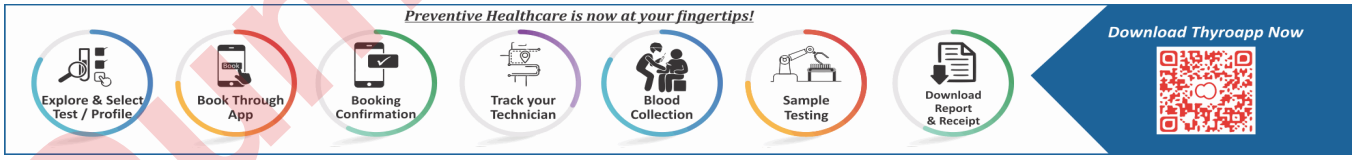
- The reported results are for information and interpretation of the referring doctor only.
- It is presumed that the tests performed on the specimen belong to the patient; named or identified.
- Results of tests may vary from laboratory to laboratory and also in some parameters from time to time for 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 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 :- https://youtu.be/nbdYeRgYyQc
- For clinical support please contact @8450950852,8450950853,8450950854 between 10:00 to 18:00

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 or feedback, write to us at **info@thyrocare.com** or call us on **022-3090 0000 / 6712 3400**
- SMS:<Labcode No.> to **9870666333**





Dummy Report