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Seshans Complex, No.140,  
Anna Salai, Saidapet,  
Chennai - 600 015



Tests you can trust

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

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**REPORT****NAME** : MRS NIVETHA M (31Y/F)**REF. BY** : SELF**TEST ASKED** : HbA1c, HEMOGRAM**HOME COLLECTION :**

1, 1ST FLOOR, THIRUMULLAI ILLAM,  
JANAKIRAMAN STREET, PERUNGUDI, CHENNAI -  
600096

**PATIENTID** : NM21881251

TEST NAME	TECHNOLOGY	VALUE	UNITS
HbA1c - (HPLC)	H.P.L.C	5.1	%

**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 method**AVERAGE BLOOD GLUCOSE (ABG)**

CALCULATED

100

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)** : 25 Jun 2023 08:45**Sample Received on (SRT)** : 25 Jun 2023 13:32**Report Released on (RRT)** : 25 Jun 2023 14:59**Sample Type** : EDTA**Labcode** : 2506075010/CHE33**Barcode** : AS338462

Dr Anusha B, MD (Path)

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**REF. BY** : SELF  
**TEST ASKED** : HbA1c, HEMOGRAM  
**PATIENTID** : NM21881251

**HOME COLLECTION :**

1, 1ST FLOOR, THIRUMULLAI ILLAM ,  
JANAKIRAMAN STREET , PERUNGUDI, CHENNAI -  
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TEST NAME	VALUE	UNITS	REFERENCE RANGE
TOTAL LEUCOCYTES COUNT (WBC)	8.47	X 10 <sup>3</sup> / μL	4.0 - 10.0
NEUTROPHILS	42.3	%	40-80
<b>LYMPHOCYTE</b>	<b>48.4</b>	<b>%</b>	<b>20-40</b>
MONOCYTES	2.1	%	2-10
<b>EOSINOPHILS</b>	<b>6.8</b>	<b>%</b>	<b>1-6</b>
BASOPHILS	0.2	%	0-2
IMMATURE GRANULOCYTE PERCENTAGE(IG%)	0.2	%	0.0-0.4
NEUTROPHILS - ABSOLUTE COUNT	3.58	X 10 <sup>3</sup> / μL	2.0-7.0
<b>LYMPHOCYTES - ABSOLUTE COUNT</b>	<b>4.1</b>	<b>X 10<sup>3</sup> / μL</b>	<b>1.0-3.0</b>
<b>MONOCYTES - ABSOLUTE COUNT</b>	<b>0.18</b>	<b>X 10<sup>3</sup> / μL</b>	<b>0.2 - 1.0</b>
<b>BASOPHILS - ABSOLUTE COUNT</b>	<b>0.02</b>	<b>X 10<sup>3</sup> / μL</b>	<b>0.02 - 0.1</b>
<b>EOSINOPHILS - ABSOLUTE COUNT</b>	<b>0.58</b>	<b>X 10<sup>3</sup> / μL</b>	<b>0.02 - 0.5</b>
IMMATURE GRANULOCYTES(IG)	0.02	X 10 <sup>3</sup> / μL	0.0-0.3
TOTAL RBC	4.12	X 10 <sup>6</sup> /μL	3.8-4.8
NUCLEATED RED BLOOD CELLS	0.01	X 10 <sup>3</sup> / μL	0.0-0.5
NUCLEATED RED BLOOD CELLS %	0.01	%	0.0-5.0
HEMOGLOBIN	12.1	g/dL	12.0-15.0
HEMATOCRIT(PCV)	39.9	%	36.0-46.0
MEAN CORPUSCULAR VOLUME(MCV)	96.8	fL	83.0-101.0
MEAN CORPUSCULAR HEMOGLOBIN(MCH)	29.4	pq	27.0-32.0
<b>MEAN CORP.HEMO.CONC(MCHC)</b>	<b>30.3</b>	<b>g/dL</b>	<b>31.5-34.5</b>
<b>RED CELL DISTRIBUTION WIDTH - SD(RDW-SD)</b>	<b>50.9</b>	<b>fL</b>	<b>39.0-46.0</b>
<b>RED CELL DISTRIBUTION WIDTH (RDW-CV)</b>	<b>14.2</b>	<b>%</b>	<b>11.6-14.0</b>
PLATELET DISTRIBUTION WIDTH(PDW)	10.7	fL	9.6-15.2
MEAN PLATELET VOLUME(MPV)	10	fL	6.5-12
PLATELET COUNT	380	X 10 <sup>3</sup> / μL	150-410
PLATELET TO LARGE CELL RATIO(PLCR)	23.5	%	19.7-42.4
PLATELETCRIT(PCT)	0.38	%	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)**

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**Labcode** : 2506075010/CHE33 Dr Anusha B, MD (Path)  
**Barcode** : AS338462

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**REPORT****NAME** : MRS NIVETHA M (31Y/F)**REF. BY** : SELF**TEST ASKED** : AAROGYAM C PRO WITH  
UTSH,FSH,FT4,LH,POTASSIUM,PROLACTIN (PRL)**PATIENTID** : NM21881251**HOME COLLECTION :**

1, 1ST FLOOR, THIRUMULLAI ILLAM,  
JANAKIRAMAN STREET, PERUNGUDI, CHENNAI -  
600096

TEST NAME	TECHNOLOGY	VALUE	UNITS
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**25-OH VITAMIN D (TOTAL)****C.L.I.A****15.98****ng/mL****Reference Range :**

DEFICIENCY : &lt;20 ng/ml || INSUFFICIENCY : 20-&lt;30 ng/ml

SUFFICIENCY : 30-100 ng/ml || TOXICITY : &gt;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.

**Method :** FULLY AUTOMATED CHEMI LUMINESCENT IMMUNO ASSAY**VITAMIN B-12****C.L.I.A****260****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.

**Method :** COMPETITIVE CHEMI LUMINESCENT IMMUNO ASSAY**Please correlate with clinical conditions.****Sample Collected on (SCT)** : 25 Jun 2023 08:45**Sample Received on (SRT)** : 25 Jun 2023 13:32**Report Released on (RRT)** : 25 Jun 2023 17:29**Sample Type** : SERUM**Labcode** : 2506075042/CHE33 Dr Anusha B, MD (Path)**Barcode** : AS338463

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**REPORT**

**NAME** : MRS NIVETHA M (31Y/F)

**REF. BY** : SELF

**TEST ASKED** : AAROGYAM C PRO WITH  
UTSH,FSH,FT4,LH,POTASSIUM,PROLACTIN (PRL)

**HOME COLLECTION :**

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JANAKIRAMAN STREET , PERUNGUDI, CHENNAI -  
600096

**PATIENTID** : NM21881251

TEST NAME	TECHNOLOGY	VALUE	UNITS
<b>HIGH SENSITIVITY C-REACTIVE PROTEIN (HS-CRP)</b>	<b>IMMUNOTURBIDIMETRY</b>	<b>4.6</b>	<b>mg/L</b>
<b>Reference Range :-</b>			

< 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 data 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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**REPORT**

**NAME** : MRS NIVETHA M (31Y/F)  
**REF. BY** : SELF  
**TEST ASKED** : AAROGYAM C PRO WITH  
UTSH,FSH,FT4,LH,POTASSIUM,PROLACTIN (PRL)

**HOME COLLECTION :**  
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JANAKIRAMAN STREET , PERUNGUDI, CHENNAI -  
600096

**PATIENTID** : NM21881251

TEST NAME	TECHNOLOGY	VALUE	UNITS
TESTOSTERONE	C.L.I.A	18.84	ng/dL

**Reference Range :-**

Adult Male

21 - 49 Yrs : 164.94 - 753.38 || 50 - 89 Yrs : 86.49 - 788.22

Adult Female

Pre-Menopause : 12.09 - 59.46 || Post-Menopause: < 7.00 - 48.93

Boys

2-10 Years : < 7.00 - 25.91

11 Years : < 7.00 - 341.53

12 Years : < 7.00 - 562.59

13 Years : 9.34 - 562.93

14 Years : 23.28 - 742.46

15 Years : 144.15 - 841.44

16-21 Years : 118.22 - 948.56

Girls

2-10 Years : < 7.00 - 108.30

11-15 Years : < 7.00 - 48.40

16-21 Years : 17.55 - 50.41

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 Hyperprolactinemia, Hypopituitarism some types of liver and kidney diseases and critical illness.

Specifications: Precision: Intra assay (%CV): 8.5 %, Inter assay (%CV): 12.6%; Sensitivity: 7 ng/dL.

Kit Validation Reference: Kicklighter EJ, Norman RJ. The gonads. In: Kaplan LA, Pesce AJ, eds. Clinical Chemistry: Theory, Analysis, Correlation. 2nd ed. St. Louis: CV Mosby; 1989:657-662.

**Please correlate with clinical conditions.**

**Method:-** COMPETITIVE CHEMI LUMINESCENT IMMUNO ASSAY

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TEST NAME	TECHNOLOGY	VALUE	UNITS
<b>IRON</b> <b>Reference Range :</b> Male : 65 - 175 Female : 50 - 170 <b>Method :</b> FERROZINE METHOD WITHOUT DEPROTEINIZATION	PHOTOMETRY	61.68	µg/dL
<b>TOTAL IRON BINDING CAPACITY (TIBC)</b> <b>Reference Range :</b> Male: 225 - 535 µg/dl Female: 215 - 535 µg/dl <b>Method :</b> SPECTROPHOTOMETRIC ASSAY	PHOTOMETRY	404.58	µg/dL
<b>% TRANSFERRIN SATURATION</b> <b>Reference Range :</b> 13 - 45 <b>Method :</b> DERIVED FROM IRON AND TIBC VALUES	CALCULATED	15.25	%
<b>UNSAT.IRON-BINDING CAPACITY(UIBC)</b> <b>Reference Range :</b> 162 - 368 <b>Method :</b> SPECTROPHOTOMETRIC ASSAY	PHOTOMETRY	342.9	µg/dL

Please correlate with clinical conditions.

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**HOME COLLECTION :**1, 1ST FLOOR, THIRUMULLAI ILLAM , JANAKIRAMAN  
STREET , PERUNGUDI, CHENNAI - 600096

TEST NAME	TECHNOLOGY	VALUE	UNITS	NORMAL RANGE
<b>TOTAL CHOLESTEROL</b>	<b>PHOTOMETRY</b>	<b>211</b>	<b>mg/dL</b>	<b>&lt; 200</b>
<b>HDL CHOLESTEROL - DIRECT</b>	<b>PHOTOMETRY</b>	<b>38</b>	<b>mg/dL</b>	<b>40-60</b>
<b>HDL / LDL RATIO</b>	<b>CALCULATED</b>	<b>0.24</b>	<b>Ratio</b>	<b>&gt; 0.40</b>
<b>LDL CHOLESTEROL - DIRECT</b>	<b>PHOTOMETRY</b>	<b>158</b>	<b>mg/dL</b>	<b>&lt; 100</b>
TRIG / HDL RATIO	CALCULATED	1.96	Ratio	< 3.12
TRIGLYCERIDES	PHOTOMETRY	74	mg/dL	< 150
<b>TC/ HDL CHOLESTEROL RATIO</b>	<b>CALCULATED</b>	<b>5.6</b>	<b>Ratio</b>	<b>3 - 5</b>
<b>LDL / HDL RATIO</b>	<b>CALCULATED</b>	<b>4.2</b>	<b>Ratio</b>	<b>1.5-3.5</b>
<b>NON-HDL CHOLESTEROL</b>	<b>CALCULATED</b>	<b>173.2</b>	<b>mg/dL</b>	<b>&lt; 160</b>
VLDL CHOLESTEROL	CALCULATED	14.78	mg/dL	5 - 40

**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

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.****Sample Collected on (SCT)** : 25 Jun 2023 08:45**Sample Received on (SRT)** : 25 Jun 2023 13:32**Report Released on (RRT)** : 25 Jun 2023 17:29**Sample Type** : SERUM**Labcode** : 2506075042/CHE33 Dr Anusha B, MD (Path)**Barcode** : AS338463

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UTSH,FSH,FT4,LH,POTASSIUM,PROLACTIN (PRL)**PATIENTID** : NM21881251**HOME COLLECTION :**

1, 1ST FLOOR, THIRUMULLAI ILLAM ,  
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600096

TEST NAME	TECHNOLOGY	VALUE	UNITS
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**FOLLICLE STIMULATING HORMONE (FSH)**

C.L.I.A

8.13

mIU/mL

**Reference Range :**

Females :

Normally Menstruating :

Follicular Phase : 2.5-10.2 | Midcycle Peak: 3.4 - 33.4 | Luteal Phase :1.5-9.1

Pregnant : &lt; 0.3 | Postmenopausal : 23.0 - 116.3

Males (13 - 70 Years) : 1.4-18.1

**Method :** Fully Automated Bidirectionally Interfaced Chemi Luminescent Immuno Assay**LUTEINISING HORMONE (LH)**

C.L.I.A

5.22

mIU/mL

**Reference Range :**

Females:

Normally Menstruating:

Follicular Phase : 1.9 - 12.5 | Midcycle Peak : 8.7 - 76.3

Luteal Phase :0.5 - 16.9 | Pregnant : 0.1 - 1.5

Postmenopausal : 15.9 - 54.0

Children : 0.1 - 6.0

Males (20 - 70 Years) : 1.5 - 9.3

&gt;70 Years : 3.1 - 34.6

**Method :** Fully Automated Bidirectionally Interfaced Chemi Luminescent Immuno Assay**PROLACTIN (PRL)**

C.L.I.A

0.37

ng/mL

**Reference Range :**

Females :

Normally Menstruating : 2.8 - 29.2

Pregnant : 9.7 - 208.5

Postmenopausal : 1.8 - 20.3

Male : 2.1 - 17.7

**Method :** Fully Automated Bidirectionally Interfaced Chemi Luminescent Immuno Assay**Please correlate with clinical conditions.****Sample Collected on (SCT)** :25 Jun 2023 08:45**Sample Received on (SRT)** : 25 Jun 2023 13:32**Report Released on (RRT)** : 25 Jun 2023 17:29**Sample Type** : SERUM**Labcode** : 2506075042/CHE33 Dr Anusha B, MD (Path)**Barcode** : AS338463



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**HOME COLLECTION :**1, 1ST FLOOR, THIRUMULLAI ILLAM , JANAKIRAMAN  
STREET , PERUNGUDI, CHENNAI - 600096

TEST NAME	TECHNOLOGY	VALUE	UNITS	NORMAL RANGE
ALKALINE PHOSPHATASE	PHOTOMETRY	86.78	U/L	45-129
BILIRUBIN - TOTAL	PHOTOMETRY	0.31	mg/dL	0.3-1.2
BILIRUBIN -DIRECT	PHOTOMETRY	0.1	mg/dL	< 0.3
BILIRUBIN (INDIRECT)	CALCULATED	0.21	mg/dL	0-0.9
GAMMA GLUTAMYL TRANSFERASE (GGT)	PHOTOMETRY	19	U/L	< 38
SGOT / SGPT RATIO	CALCULATED	1.1	Ratio	< 2
ASPARTATE AMINOTRANSFERASE (SGOT )	PHOTOMETRY	20.7	U/L	< 31
ALANINE TRANSAMINASE (SGPT)	PHOTOMETRY	18.8	U/L	< 34
PROTEIN - TOTAL	PHOTOMETRY	7.97	gm/dL	5.7-8.2
ALBUMIN - SERUM	PHOTOMETRY	4.35	gm/dL	3.2-4.8
<b>SERUM GLOBULIN</b>	<b>CALCULATED</b>	<b>3.62</b>	<b>gm/dL</b>	<b>2.5-3.4</b>
SERUM ALB/GLOBULIN RATIO	CALCULATED	1.2	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 BCG<sup>1</sup>METHOD (COLORIMETRIC ASSAY ENDPOINT)  
 SEGB - DERIVED FROM SERUM ALBUMIN AND PROTEIN VALUES  
 A/GR - DERIVED FROM SERUM ALBUMIN AND PROTEIN VALUES

**Sample Collected on (SCT)** : 25 Jun 2023 08:45**Sample Received on (SRT)** : 25 Jun 2023 13:32**Report Released on (RRT)** : 25 Jun 2023 17:29**Sample Type** : SERUM**Labcode** : 2506075042/CHE33 Dr Anusha B, MD (Path)**Barcode** : AS338463

**PROCESSED AT :****Thyrocare**

Seshans Complex, No.140,  
Anna Salai, Saidapet,  
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**REPORT**

**NAME** : MRS NIVETHA M (31Y/F)  
**REF. BY** : SELF  
**TEST ASKED** : AAROGYAM C PRO WITH  
UTSH,FSH,FT4,LH,POTASSIUM,PROLACTIN (PRL)  
**PATIENTID** : NM21881251

**HOME COLLECTION :**

1, 1ST FLOOR, THIRUMULLAI ILLAM , JANAKIRAMAN  
STREET , PERUNGUDI, CHENNAI - 600096

TEST NAME	TECHNOLOGY	VALUE	UNITS	REFERENCE RANGE
TOTAL TRIIODOTHYRONINE (T3)	C.L.I.A	99	ng/dL	60-200
TOTAL THYROXINE (T4)	C.L.I.A	7.3	µg/dL	4.5-12
TSH - ULTRASENSITIVE	C.L.I.A	1.378	µIU/mL	0.55-4.78
FREE THYROXINE (FT4)	C.L.I.A	1.03	ng/dL	0.7-1.8

**Comments :** \*\*\*

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

**Method :**

T3 - COMPETITIVE CHEMI LUMINESCENT IMMUNO ASSAY

T4 - COMPETITIVE CHEMI LUMINESCENT IMMUNO ASSAY

USTSH - Third Generation Ultrasensitive Chemi Luminescent Immuno Assay

FT4 - Competitive Chemi Luminescent Immuno Assay

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)

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Tests you can trust

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1, 1ST FLOOR, THIRUMULLAI ILLAM ,  
JANAKIRAMAN STREET , PERUNGUDI, CHENNAI -  
600096

TEST NAME	TECHNOLOGY	VALUE	UNITS
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**POTASSIUM**

I.S.E

4.14

mmol/L

**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**Please correlate with clinical conditions.****Sample Collected on (SCT)** : 25 Jun 2023 08:45**Sample Received on (SRT)** : 25 Jun 2023 13:32**Report Released on (RRT)** : 25 Jun 2023 17:29**Sample Type** : SERUM**Labcode** : 2506075042/CHE33 Dr Anusha B, MD (Path)**Barcode** : AS338463

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**PATIENTID** : NM21881251

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STREET , PERUNGUDI, CHENNAI - 600096

TEST NAME	TECHNOLOGY	VALUE	UNITS	NORMAL RANGE
<b>UREA (CALCULATED)</b>	<b>CALCULATED</b>	<b>16.22</b>	<b>mg/dL</b>	<b>Adult : 17-43</b>
<b>BLOOD UREA NITROGEN (BUN)</b>	<b>PHOTOMETRY</b>	<b>7.58</b>	<b>mg/dL</b>	<b>7.94 - 20.07</b>
UREA / SR.CREATININE RATIO	CALCULATED	22.85	Ratio	< 52
CREATININE - SERUM	PHOTOMETRY	0.71	mg/dL	0.55-1.02
BUN / SR.CREATININE RATIO	CALCULATED	10.68	Ratio	9:1-23:1
CALCIUM	PHOTOMETRY	9.67	mg/dL	8.8-10.6
URIC ACID	PHOTOMETRY	5.12	mg/dL	3.2 - 6.1
SODIUM	I.S.E	137.7	mmol/L	136 - 145
CHLORIDE	I.S.E	103.4	mmol/L	98 - 107

**Please correlate with clinical conditions.**

**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

CALC - ARSENAZO III METHOD, END POINT.

URIC - URICASE / PEROXIDASE METHOD

SOD - ION SELECTIVE ELECTRODE

CHL - ION SELECTIVE ELECTRODE

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600096

**PATIENTID** : NM21881251

TEST NAME	TECHNOLOGY	VALUE	UNITS
EST. GLOMERULAR FILTRATION RATE (eGFR)	CALCULATED	113	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 ~~

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