





# **Diagnostics & Speciality Centre**

MR NO. NAME Mr. RANA KUMAR MAUSUM : 21100455

AGE/SEX 34 Yrs / Male VISIT NO. 141946

DATE OF COLLECTION: 09-10-2021 at 09:18 AM REFERRED BY ·

> DATE OF REPORT : 09-10-2021 at 02:17 PM

REF CENTER : MEDIWHEEL 

**RESULT** REFERENCE RANGE **SPECIMEN TEST PARAMETER** 

### **HAEMATOLOGY**

### COMPLETE BLOOD COUNT (CBC) WITH ESR

**HAEMOGLOBIN** 16.2 gm/dL 13 - 18 gm/dL Colorimetric Method

**HEMATOCRIT (PCV)** 49.4 % 40 - 54 %

Calculated

4.5 - 5.9 million/cu.mm RED BLOOD CELL (RBC) COUNT 5.1 million/cu.mm

Electrical Impedance

PLATELET COUNT 1.7 Lakhs/cumm 1.5 - 4.5 Lakhs/cumm

Electrical Impedance

MEAN CELL VOLUME (MCV) 96.8 fl 80 - 100 fl

MEAN CORPUSCULAR HEMOGLOBIN (MCH) 31.7 pg 26 - 34 pg Calculated

MEAN CORPUSCULAR HEMOGLOBIN 32.8 % 31 - 35 %

**CONCENTRATION (MCHC)** 

Calculated

TOTAL WBC COUNT (TC) 5650 cells/cumm 4000 - 11000 cells/cumm

Electrical Impedance

**DIFFERENTIAL COUNT** 

**NEUTROPHILS** 58 % 40 - 75 % VCS Technology/Microscopic 25 - 40 % LYMPHOCYTES 35 % VCS Technology/Microscopic

**EOSINOPHILS** 02 % 0 - 7 % VCS Technology/Microscopic

**MONOCYTES** 05 % 1 - 8 %

VCS Technology/Microscopic **BASOPHILS** 00 %

Electrical Impedance

15 mm/hr 0 - 15 mm/hr **ESR** Westergren Method

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Dr. KRISHNA MURTHY Dr. VAMSEEDHAR.A Lab Seal

MD **BIOCHEMIST** 

D.C.P, M.D CONSULTANT PATHOLOGIST

The laboratory values And Normal values need to be interpreted based on patients clinical characteristics. The values in reference range is for an average normal individual which may vary depending upon age, sex and other characteristics.

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TEST PARAMETER RESULT REFERENCE RANGE SPECIMEN

"O" Positive

BLOOD GROUP & Rh TYPING

Tube Agglutination (Forward and Reverse)

GLYCATED HAEMOGLOBIN (HbA1C)

7.2 %

American Diabetic Association (ADA) recommendations:

Non diabetic adults: <5.7 %

At risk (Pre diabetic): 5.7 -

6.4%

Diabetic: >/= 6.5%

Therapeutic goal for glycemic control:

Goal for therapy: < 7.0%

Action suggested: > 8.0%

ESTIMATED AVERAGE GLUCOSE (eAG) 159.94 mg/dL

Calculation
Comments:

This assay is useful for diagnosing Diabetes and evaluating long term control of blood glucose concentrations in diabetic patients. It reflects the mean glucose concentration over the previous period of 8 to 12 weeks and is a better indicator of long term glycemic control as compared with blood and urine glucose measurements. This provides a additional criterion for assessing glucose control because glycated hemoglobin values are free of day-to-day glucose fluctuation and are unaffected by exercise or food ingestion.

After a sudden alteration in blood glucose concentration, the rate of change of HbA1c is rapid during initial 2 months, followed by more gradual change approaching steady state 3 months later.

Collegn. u.



A. Hurudhay

Dr. KRISHNA MURTHY

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**SPECIMEN** 



(A Unit of Vijayalakshmi Diagnostics Pvt. Ltd.)

REFERENCE RANGE

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CLINICAL BIOCHEMISTRY

FASTING BLOOD SUGAR 138.5 mg/dl 70 - 110 mg/dl

Hexokinase

**TEST PARAMETER** 

CREATININE 1.02 mg/dL 0.8 - 1.4 mg/dL

Jaffe Method

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MD

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TEST PARAMETER RESULT REFERENCE RANGE SPECIMEN

**LIPID PROFILE TEST** 

TOTAL CHOLESTEROL 179.1 mg/dL up to 200 mg/dL

Cholesterol Oxidase-Peroxidase (CHOD-POD)

Border Line: 200 – 240 mg/dL

High: > 240 mg/dL

TRIGLYCERIDES 161.5 mg/dL up to 150 mg/dL

Glycerol Peroxidase-Peroxidase (GPO-POD)

Desirable: <150 mg/dL

Border Line: 150 – 200 mg/dL High: >200 – 500 mg/dL Very High: > 500 mg/dL

HDL CHOLESTEROL - DIRECT 57.1 mg/dl 40 - 60 mg/dl

PEG-Cholesterol Esterase >/= 60mg/dL - Excellent (protects

against heart disease)
40-59 mg/dL - Higher the better
<40 mg/dL - Lower than desired (major risk for heart disease)

LDL CHOLESTEROL - DIRECT 89.7 mg/dL up to 100 mg/dL

Cholesterol Esterase-Cholesterol Oxidase 100-129 mg/dL- Near optimal/above

optimal

130-159 mg/dL- Borderline High 160-189 mg/dL- High 190->190 mg/dL - Very High

VLDL CHOLESTEROL 32.3 mg/dL 2 - 30 mg/dL

Calculation

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TEST PARAMETER	RESULT	REFERENCE RANGE	SPECIMEN
TOTAL CHOLESTROL/HDL RATIO Calculation	3.1	up to 3.5 3.5-5.0 - Moderate >5.0 - High	
LDL/HDL RATIO Calculation	1.6	up to 2.5 2.5-3.3 - Moderate >3.3 - High	
LIVER FUNCTION TEST (LFT)			
TOTAL BILIRUBIN Colorimetric Diazo Method	0.96 mg/dL	0.2 - 1.2 mg/dL	
DIRECT BILIRUBIN Colorimetric Diazo Method	0.24 mg/dL	0 - 0.4 mg/dL	
INDIRECT BILIRUBIN Calculation	0.72 mg/dl		
S G O T (AST)  IFCC Without Pyridoxal Phosphates	<b>46.7</b> U/L	up to 35 U/L	
S G P T (ALT)  IFCC Without Pyridoxal Phosphates	<b>67.2</b> U/L	up to 50 U/L	
ALKALINE PHOSPHATASE p-Nitrophenyl Phosphate	112.4 U/L	36 - 113 U/L	
SERUM GAMMA GLUTAMYLTRANSFERASE (GGT	7) 70.1 U/L	15 - 85 U/L	
TOTAL PROTEIN Biuret Colorimetric	6.52 g/dl	6.2 - 8 g/dl	
S.ALBUMIN Bromocresol Green (BCG)	4.23 g/dl	3.5 - 5.2 g/dl	
S.GLOBULIN Calculation	<b>2.3</b> g/dl	2.5 - 3.8 g/dl	
A/G RATIO	1.8	1 - 1.5	

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Calculation



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POST PRANDIAL BLOOD SUGAR Hexokinase	<b>196.3</b> mg/dl	80 - 150 mg/dl	
BLOOD UREA	16.6 mg/dL	15 - 50 mg/dL	

**RESULT** 

UREASE-GLUTAMATE DEHYDROGENASE (GLDH) 1.02 mg/dL 0.4 - 1.4 mg/dL **CREATININE** Jaffe Kinetic

**URIC ACID** 5.1 mg/dL 3 - 7.2 mg/dL Uricase-Peroxidase

SERUM ELECTROLYTES

SODIUM 139 mmol/L 136 - 145 mmol/L Ion Selective Electrode (ISE) 4.6 mmol/L 3.5 - 5.2 mmol/L **POTASSIUM** Ion Selective Electrode (ISE) **CHLORIDE** 103 mmol/L 97 - 111 mmol/L

Ion Selective Electrode (ISE)

**TEST PARAMETER** 

### **CLINICAL PATHOLOGY**

### **URINE ROUTINE & MICROSCOPIC** PHYSICAL EXAMINATION

Appearance Visual Method  Appearance Visual Method  Specific Gravity Strips Method pH  6.0  Fale Yellow Pale yellow- Yellow Pale Yellow Pa	CHEMICAL EXAMINATION (DIPSTICK)					
Visual Method  Appearance Visual Method  Specific Gravity  Clear  Clear  Clear/Transparent  1.015  1.005-1.035	pH	6.0	4.6-8.5			
Visual Method  Appearance Clear Clear/Transparent		1.015	1.005-1.035			
The second secon	• •	Clear	Clear/Transparent			
Colour Polo Vollow Polo vollow vollow	Colour Visual Method	Pale Yellow	Pale yellow- yellow			

Protein Nil Nil -Trace Strips Method

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**TEST PARAMETER RESULT** REFERENCE RANGE **SPECIMEN** 

Glucose Nil Nil Strips Method

Negative Negative Blood

Strips Method

Ketone Bodies Absent Negative Strips Method

Urobilinogen Normal Normal Strips Method

Bile Salt Negative Negative

Strips Method

Bilirubin Negative Negative

Bile Pigments Negative NIL

**MICROSCOPY** 

Cast

Pus Cells (WBC) 3 - 4 /hpf 0-5/hpf

1 - 2 /hpf 0-4/hpf **Epithelial Cells** 

Light Microscopic Not Seen /hpf **RBC** 0-2/hpf

Light Microscopic

NIL

Light Microscopic

Nil NIL Crystal Light Microscopic

**FASTING URINE SUGAR (FUS)** NIL NIL

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NIL

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**STOOL ROUTINE EXAMINATION** 

MACROSCOPIC EXAMINATION

**COLOUR Brownish** Light to Dark brown Manual

Semi Solid Well formed-semi solid CONSISTENCY

**MUCUS** Absent Absent

Manual

**BLOOD** Absent Absent

CHEMICAL EXAMINATION

Manual

1 - 2 Absent **PUS CELLS** Light Microscopy

**EPITHELIAL CELLS** 1-2 Few

**BACTERIA** Present (++) Light Microscopy

MICROSCOPIC EXAMINATION

**CYST** Not Seen Absent Light Microscopy

Not Seen **OVA** Absent

Light Microscopy

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### **IMMUNOASSAY**

### THYROID PROFILE

**TOTAL TRIIODOTHYRONINE (T3)** 1.06 ng/mL 0.87 - 1.78 ng/mL

**TOTAL THYROXINE (T4)** 8.47 µg/dL 6.09 - 12.23 µg/dL

THYROID STIMULATING HORMONE (TSH) 2.058 µIU/mL 0.38 - 5.33 µIU/mL

1st Trimester: 0.05 - 3.70 2nd Trimester: 0.31 - 4.35 3rd Trimester: 0.41 - 5.18

#### Note:

- TSH levels are subject to circadian variation, reaching peak levels between 2 4.a.m. and at a minimum between 6-10 pm. The variation is of the order of 50%, hence time of the day has influence on the measured serum TSH concentrations.
- Recommended test for T3 and T4 is unbound fraction or free levels as it is metabolically active.
- Physiological rise in Total T3 / T4 levels is seen in pregnancy and in patients on steroid therapy.

### Clinical Use:

- Primary Hypothyroidism
- Hyperthyroidism
- Hypothalamic Pituitary hypothyroidism
- Inappropriate TSH secretion
- Nonthyroidal illness
- Autoimmune thyroid disease
- Pregnancy associated thyroid disorders
- Thyroid dysfunction in infancy and early childhood

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**PROSTATIC SPECIFIC ANTIGEN (PSA)** 

: MEDIWHEEL

Up to 4ng/mL: Normal PROSTATIC SPECIFIC ANTIGEN (PSA) 0.45 ng/mL

4-10 ng/mL Hypertrophy & benign genito urinary

conditions.

>10 ng/mL Suspicious of

malignancy.

PSA is used for monitoring patients with a history of prostate cancer and as an early indicator of recurrence and response to treatment. The test is commonly used for Prostate cancer screening.

Dispatched by: Sumalatha

REF CENTER

\*\*\*\* End of Report \*\*\*\*

Printed by: Sumalatha on 09-10-2021 at 02:17 PM

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