

DEPARTMENT OF LABORATORY MEDICINE

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| Patient Name : Mr. SANTHOSH S | Order No : 1000072780 |
| UHID : UHJA23018273 | Registered On : 13/02/2024 08:28:57 AM |
| Age/Sex : 40/Years Male | Collected On : 13/02/2024 08:37:48 AM |
| Ward / Bed No : | Reported On : 13/02/2024 01:04:17 PM |
| Reference : Dr. Preventive Health Check Up | Bill No : OPBJA230022614 |
| Station : At Hospital | Mobile No : 9886105101 |
| Payer Name : Mediwheel | Report Status : Final Report |

| Test Name | Result | Unit | Bio. Ref. Interval |
|--|------------|--------|--|
| BIOCHEMISTRY | | | |
| FASTING GLUCOSE (Method: Hexokinase) | 119 | mg/dL | ADA Guidelines < 100 mg/dl - Normal 100 to 125 mg/dl - Prediabetes ≥ 126 mg/dl - Diabetes |
| POST PRANDIAL GLUCOSE (Method: Hexokinase) | 146 | mg/dL | 70-140 |
| GLYCOSYLATED HAEMOGLOBIN (HBA1C) | | | Sample: Whole blood (EDTA) |
| HBA1C (Method: HPLC) | 5.8 | % | ADA Guidelines < 5.7% - Normal 5.7 to 6.4% - Prediabetes ≥ 6.5% - Diabetes |
| Estimated Average Glucose (eAG) (Method: Calculated) | 119.75 | mg/dL | |
| THYROID PROFILE (TOTAL T3, TOTAL T4 & TSH) | | | Sample: Serum |
| TOTAL T3 (Method:CLIA) | 1.01 | ng/mL | 0.87-1.78 |
| TOTAL T4 (Method:CLIA) | 8.17 | µg/dL | 5.1-14.1 |
| THYROID STIMULATING HORMONE (TSH) (Method:CLIA: Ultra-sensitive) | 2.87 | µIU/mL | 0.34-5.60 |
| LIPID PROFILE | | | Sample: Serum |
| TOTAL CHOLESTEROL (Method:CHOD-POD) | 279 | mg/dL | ATP III Guidelines < 200 - Desirable 200-239 - Borderline high ≥ 240 - High |
| TRIGLYCERIDES (Method:Enzymatic GPO-POD) | 219 | mg/dL | < 150 - Normal 150-199 - Borderline High 200-499 - High ≥ 500 - Very High |
| HDL CHOLESTEROL (Method:ENZYMATIC METHOD) | 48.0 | mg/dL | < 40 - Low ≥ 60 - High |

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| LDL CHOLESTEROL (Method:ENZYMATIC METHOD) | 187.2 | mg/dL | <100 - Optimal 100-129 - Near or above optimal 130-159 - Borderline high 160-189 - High ≥190 - Very high |
| VLDL CHOLESTEROL (Method: Calculated) | 43.79 | mg/dL | < 30 |
| TOTAL CHOLESTEROL : HDL RATIO (Method: Calculated) | 5.8 | | Low Risk: 3.3 - 4.4 Average Risk: 4.5 - 7.1 Moderate Risk: 7.2 - 11.0 |
| LDL/HDL CHOLESTEROL RATIO (Method: Calculated) | 3.9 | | < 2.5 Optimal |
| NON HDL CHOLESTEROL (Method: Calculated) | 231 | mg/dL | < 130 |
| URIC ACID (Method:Uricase - POD(Enzymatic)) | 7.2 | mg/dL | 3.5-7.2 |
| BUN/CREATININE RATIO | | | Sample: Serum |
| BLOOD UREA NITROGEN(BUN) (Method:Urease GLDH - Kinetic) | 16 | mg/dL | 7.93-20.07 |
| CREATININE (Method:Modified Jaffe, Kinetic) | 0.89 | mg/dL | 0.9-1.3 |
| BUN/CRE-RATIO (Method: Calculated) | 17.97 | | 12-20 : 1 |
| LIVER FUNCTION TEST | | | Sample: Serum |
| TOTAL BILIRUBIN (Method:Dichlorophenyl Diazotization) | 0.66 | mg/dL | 0.3-1.2 |
| DIRECT BILIRUBIN (Method:Dichlorophenyl Diazotization) | 0.12 | mg/dL | 0.0-0.2 |
| INDIRECT BILIRUBIN (Method: Calculated) | 0.55 | mg/dL | 0.2-1.0 |
| TOTAL PROTEIN (Method:BIURET) | 7.2 | g/dL | 6.6-8.3 |

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| ALBUMIN (Method:BCG) | 4.17 | g/dL | 3.5-5.2 |
| GLOBULIN (Method: Calculated) | 3.03 | g/dL | 2.3-3.5 |
| AG RATIO (Method: Calculated) | 1.37 | | 2:1 |
| SERUM SGOT (Method:IFCC without P5P) | 24 | U/L | < 50 |
| SERUM SGPT (Method:IFCC without P5P) | 23 | U/L | < 50 |
| ALKALINE PHOSPHATASE, SERUM (Method:PNPP AMP Buffer) | 50 | U/L | 50-116 |
| GGT (Method:IFCC) | 71 | U/L | < 55 |
| PROSTATE SPECIFIC ANTIGEN (PSA) (Method:CLIA) | 0.50 | ng/mL | < 4.0 |

Interpretation Notes

Serum PSA concentrations should not be interpreted as absolute evidence for the presence or absence of malignant disease nor should serum PSA be used alone as a screening test for malignant disease. For diagnostic purposes, the results obtained by immunometric assay should always be used in combination with the clinical examinations, patient medical history and other findings. The concentration of PSA in a given specimen, determined with assays from different manufacturers, may not be comparable due to differences in assay methods, calibration, and reagent specificity.

| | | | |
|---|------|-------|-------|
| UREA (Method:Urease GLDH - Kinetic) | 35.3 | mg/dL | 17-43 |
|---|------|-------|-------|



Dr. Shanthakumar Muruda
Sr CONSULTANT BIOCHEMIST
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HAEMATOLOGY

COMPLETE BLOOD COUNT(CBC)

Sample: Whole blood (EDTA)

| | | | |
|---|-------|-------------|------------|
| HAEMOGLOBIN (Method:Photometric Measurement: Oxyhemoglobin method) | 14.67 | g/dL | 13.5-17.5 |
| PACKED CELL VOLUME/HEMATOCRIT (PCV/HCT) (Method: Calculated) | 44.9 | % | 42-52 |
| TOTAL WBC COUNT (TLC) (Method:Coulter Principle) | 4930 | Cells/Cum | 4000-11000 |
| DIFFERENTIAL COUNT | | | |
| NEUTROPHILS (Method:Optical/Impedance) | 64.45 | % | 40-75 |
| LYMPHOCYTES (Method:Optical/Impedance) | 23.35 | % | 20-45 |
| EOSINOPHILS (Method:Optical/Impedance) | 2.85 | % | 0-6 |
| MONOCYTES (Method:Optical/Impedance) | 8.87 | % | 2-10 |
| BASOPHILS (Method:Optical/Impedance) | 0.48 | % | 0-2 |
| RED BLOOD CORPUSCLES(RBC) (Method:Coulter Principle) | 5.65 | million/cum | 4.5-5.9 |
| MCV (Method:Derived from RBC Histogram) | 79.5 | fL | 78-100 |
| MCH (Method: Calculated) | 26.0 | pg | 27-31 |
| MCHC (Method: Calculated) | 32.7 | g/dL | 31-37 |
| RDW - CV (Method: Calculated) | 14.9 | % | 11.5-14.5 |
| PLATELET COUNT (Method:Electrical Impedance) | 2.46 | Lakhs/Cum | 1.5-4.5 |

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| MEAN PLATELET VOLUME(MPV) (Method:Derived from PLT Histogram) | 8.19 | fl | 9-13 |
| PLATELET DISTRIBUTION WIDTH (PDW) (Method: Calculated) | 20.2 | fl | 9-19 |
| ERYTHROCYTE SEDIMENTATION RATE(ESR) (Method:Modified Westergren Method) | 10 | mm/hour | 1-15 |

BLOOD GROUPING & RH TYPING

Sample: Whole blood (EDTA)

| | |
|---|----------|
| ABO Group (Method:Agglutination Gel Method) | A |
| Rh Factor (Method:Agglutination Gel Method) | Positive |

Interpretation Notes

Note: Both forward and reverse grouping performed

Naveen N

Dr. Naveen Kumar
CONSULTANT PATHOLOGIST
KMC NO : 71418

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CLINICAL PATHOLOGY
URINE EXAMINATION, ROUTINE

Sample: Urine

PHYSICAL EXAMINATION

| | | | |
|------------------|-------------|----|-------------|
| VOLUME | 25 | mL | |
| COLOUR | Pale Yellow | | |
| APPEARANCE | Clear | | |
| PH | 6.5 | | 5.0-8.0 |
| SPECIFIC GRAVITY | 1.025 | | 1.005-1.030 |

CHEMICAL EXAMINATION

| | | | |
|--|----------|--|----------|
| PROTEIN (Method:Protein Error of pH Indicator) | Absent | | Absent |
| GLUCOSE (Method:GOD-POD) | Absent | | Absent |
| KETONE BODIES (Method:Nitroprusside method/ Rothera's test) | Absent | | Absent |
| BILIRUBIN (Method:DIAZO/FOUCHET'S TEST) | Negative | | Negative |
| BILE SALT (Method:Hay's sulfur test) | Absent | | Absent |
| NITRITE (Method:Griess method) | Negative | | Negative |
| UROBILINOGEN (Method:Azo coupling method) | Normal | | |
| LEUKOCYTE ESTERASE (Method:Leukocyte Esterase activity) | Negative | | Negative |
| BLOOD (Method:Peroxidase Reaction) | Negative | | Negative |

MICROSCOPIC EXAMINATION

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| EPITHELIAL CELLS | 2-4 | /HPF | 0-5 |
| PUS CELLS | 2-4 | /HPF | 0-5 |
| RBCs | Nil | /HPF | 0-2 |
| CASTS | Nil | /LPF | |
| CRYSTALS | Nil | | |
| OTHERS | Nil | | |

Verified By
Parameshwar B

---End of Report---

Naveen M

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