

| Name | Mr.SIDDALINGAIAH | ID | MED121173651 |
|-----------------|------------------|------------|--------------|
| Age & Gender | 56/MALE | Visit Date | 04/07/2022 |
| Ref Doctor Name | MediWheel | | |

ABDOMINO-PELVIC ULTRASONOGRAPHY

LIVER is normal in shape, size (12.7 cm) and has uniform echopattern. No evidence of focal lesion or intrahepatic biliary ductal dilatation. Hepatic and portal vein radicals are normal.

GALL BLADDER is partially distended. CBD is not dilated.

PANCREAS has normal shape, size and uniform echopattern. No evidence of ductal dilatation or calcification.

SPLEEN shows normal shape, size (11.6 cm) and echopattern.

No demonstrable Para-aortic lymphadenopathy.

KIDNEYS

Right kidney: Normal in shape, size and echopattern. Cortico-medullary differentiation is well madeout. No evidence of calculus or hydronephrosis.

Left kidney: Normal in shape, size and echopattern. Cortico-medullary differentiation is well madeout. No evidence of calculus or hydronephrosis.

The kidney measures as follows:

| | Bipolar length (cm) | Parenchymal thickness (cm) |
|--------------|---------------------|----------------------------|
| Right Kidney | 10.8 | 1.4 |
| Left Kidney | 10.3 | 1.5 |

URINARY BLADDER shows normal shape and wall thickness. It has clear contents. No evidence of diverticula.

PROSTATE shows normal shape, size and echopattern. It measures 3.2 x 2.9 x 4.0 cm volume: 19.4 cc.

No evidence of ascites.

IMPRESSION:

• No significant abnormality detected.

REPORT DISCLAIMER

- 1. This is only a radiologincal imperssion. Like other investigations, radiological investication also have limitation. Therefore radiologincal reports should be interpreted in correlation with clinical and pathological findings.
- 2. The results reported here in are subject to interpretation by qualified medical professionals only.
- 3.Customer identities are accepted provided by the customer or their representative. 4.information about the customer's condition at the time of sample collection such as fasting, food
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- 5.If any specimen/sample is received from any others laboratory/hospital, its is presumed that the sample belongs to the patient identified or named.
- 6.Test results should be interpreted in context of clinical and other findings if any.In case of any clarification /doubt , the refrering doctor/patient can contact the respective section head of the laboratory.
- 7.Results of the test are influenced by the various factors such as sensitivity, specificity of the procedures of the tests, quality of the samples and drug interactions etc.,
- 8.If the test nexults are found not to be correlating clinically can contact the lab in charge for clarification or retesting where practicable within 24 hours from the time of issue of results.
- 9.Liability is limited to the extend of amount billed.
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- 11.Disputes, if any, with regard to the report findings are subject to the exclusive jurisdiction of the competent courts chennai only.



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DR. HEMANANDINI V.N

CONSULTANT RADIOLOGIST Hn/mj

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| Age / Sex | : 56 Year(s) / Male | Report On : 04/07/2022 2:01 PM | MEDALL |
| Туре | : OP | Printed On : 15/07/2022 11:51 AM | |
| Ref. Dr | : MediWheel | | |

| Investigation | <u>Observed</u> <u>Value</u> | <u>Unit</u> | <u>Biological</u> <u>Reference Interval</u> |
|---|---------------------------------|-------------|--|
| HAEMATOLOGY | | | |
| Complete Blood Count With - ESR | | | |
| Haemoglobin (EDTA Blood'Spectrophotometry) | 15.2 | g/dL | 13.5 - 18.0 |
| Packed Cell Volume(PCV)/Haematocrit (EDTA Blood) | 44.6 | % | 42 - 52 |
| RBC Count (EDTA Blood) | 5.08 | mill/cu.mm | 4.7 - 6.0 |
| Mean Corpuscular Volume(MCV) (EDTA Blood) | 87.8 | fL | 78 - 100 |
| Mean Corpuscular Haemoglobin(MCH) (EDTA Blood) | 30.0 | pg | 27 - 32 |
| Mean Corpuscular Haemoglobin concentration(MCHC) (EDTA Blood) | 34.2 | g/dL | 32 - 36 |
| RDW-CV | 13.1 | % | 11.5 - 16.0 |
| RDW-SD | 40.26 | fL | 39 - 46 |
| Total Leukocyte Count (TC) (EDTA Blood) | 8500 | cells/cu.mm | 4000 - 11000 |
| Neutrophils (Blood) | 51.5 | % | 40 - 75 |
| Lymphocytes (Blood) | 33.0 | % | 20 - 45 |
| Eosinophils (Blood) | 7.7 | % | 01 - 06 |



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|--|---------------------------------|------------------------|--|
| Monocytes (Blood) | 6.7 | % | 01 - 10 |
| Basophils (Blood) | 1.1 | % | 00 - 02 |
| INTERPRETATION: Tests done on Automated Five P | art cell counter. All a | abnormal results are r | eviewed and confirmed microscopically. |
| Absolute Neutrophil count (EDTA Blood) | 4.38 | 10^3 / µl | 1.5 - 6.6 |
| Absolute Lymphocyte Count (EDTA Blood) | 2.81 | 10^3 / µl | 1.5 - 3.5 |
| Absolute Eosinophil Count (AEC) (EDTA Blood) | 0.65 | 10^3 / µl | 0.04 - 0.44 |
| Absolute Monocyte Count (EDTA Blood) | 0.57 | 10^3 / µl | < 1.0 |
| Absolute Basophil count (EDTA Blood) | 0.09 | 10^3 / µl | < 0.2 |
| Platelet Count (EDTA Blood) | 253 | 10^3 / µl | 150 - 450 |
| MPV (Blood) | 6.5 | fL | 7.9 - 13.7 |
| PCT (Automated Blood cell Counter) | 0.16 | % | 0.18 - 0.28 |
| ESR (Erythrocyte Sedimentation Rate) (Citrated Blood) | 6 | mm/hr | < 20 |



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| BIOCHEMISTRY | | | |
| Liver Function Test | | | |
| Bilirubin(Total) (Serum/DCA with ATCS) | 0.32 | mg/dL | 0.1 - 1.2 |
| Bilirubin(Direct) (Serum/Diazotized Sulfanilic Acid) | 0.16 | mg/dL | 0.0 - 0.3 |
| Bilirubin(Indirect) (Serum/Derived) | 0.16 | mg/dL | 0.1 - 1.0 |
| SGOT/AST (Aspartate Aminotransferase) (Serum/ <i>Modified IFCC</i>) | 16.68 | U/L | 5 - 40 |
| SGPT/ALT (Alanine Aminotransferase) (Serum/ <i>Modified IFCC</i>) | 17.06 | U/L | 5 - 41 |
| GGT(Gamma Glutamyl Transpeptidase) (Serum/IFCC / Kinetic) | 41.25 | U/L | < 55 |
| Alkaline Phosphatase (SAP) (Serum/Modified IFCC) | 99.9 | U/L | 56 - 119 |
| Total Protein (Serum/Biuret) | 7.24 | gm/dl | 6.0 - 8.0 |
| Albumin (Serum/Bromocresol green) | 4.37 | gm/dl | 3.5 - 5.2 |
| Globulin (Serum/Derived) | 2.87 | gm/dL | 2.3 - 3.6 |
| A : G RATIO | 1.52 | | 1.1 - 2.2 |

(Serum/Derived)



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|---|---------------------------------|-------------|--|
| Lipid Profile | | | |
| Cholesterol Total (Serum/CHOD-PAP with ATCS) | 125.97 | mg/dL | Optimal: < 200 Borderline: 200 - 239 High Risk: >= 240 |
| Triglycerides (Serum/GPO-PAP with ATCS) | 167.23 | mg/dL | Optimal: < 150 Borderline: 150 - 199 High: 200 - 499 Very High: >=500 |

INTERPRETATION: The reference ranges are based on fasting condition. Triglyceride levels change drastically in response to food, increasing as much as 5 to 10 times the fasting levels, just a few hours after eating. Fasting triglyceride levels show considerable diurnal variation too. There is evidence recommending triglycerides estimation in non-fasting condition for evaluating the risk of heart disease and screening for metabolic syndrome, as non-fasting sample is more representative of the `usual_circulating level of triglycerides during most part of the day.

| HDL Cholesterol (Serum/Immunoinhibition) | 35.69 | mg/dL | Optimal(Negative Risk Factor): >= 60 Borderline: 40 - 59 High Risk: < 40 |
|--|-------|-------|---|
| LDL Cholesterol (Serum/ <i>Calculated</i>) | 56.9 | mg/dL | Optimal: < 100 Above Optimal: 100 - 129 Borderline: 130 - 159 High: 160 - 189 Very High: >=190 |
| VLDL Cholesterol (Serum/Calculated) | 33.4 | mg/dL | < 30 |
| Non HDL Cholesterol (Serum/Calculated) | 90.3 | mg/dL | Optimal: < 130 Above Optimal: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very High: >=220 |



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|---|---------------------------------|---|
| INTERPRETATION: 1.Non-HDL Cholesterol is now 2.It is the sum of all potentially atherogenic proteins inc co-primary target for cholesterol lowering therapy. | * | cardiovascular risk marker than LDL Cholesterol. LDL and chylomicrons and it is the "new bad cholesterol" and is a |
| Total Cholesterol/HDL Cholesterol Ratio (Serum/Calculated) | 3.5 | Optimal: < 3.3 Low Risk: 3.4 - 4.4 Average Risk: 4.5 - 7.1 Moderate Risk: 7.2 - 11.0 High Risk: > 11.0 |
| Triglyceride/HDL Cholesterol Ratio (TG/HDL) (Serum/ <i>Calculated</i>) | 4.7 | Optimal: < 2.5 Mild to moderate risk: 2.5 - 5.0 High Risk: > 5.0 |
| LDL/HDL Cholesterol Ratio (Serum/Calculated) | 1.6 | Optimal: 0.5 - 3.0 Borderline: 3.1 - 6.0 High Risk: > 6.0 |



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| Investigation Glycosylated Haemoglobin (HbA1c) | <u>Observed</u> <u>Value</u> | <u>Unit</u> | <u>Biological</u> <u>Reference Interval</u> | | |
|--|---------------------------------|-------------|---|--|--|
| HbA1C (Whole Blood/ <i>HPLC</i>) | 9.5 | % | Normal: 4.5 - 5.6 Prediabetes: 5.7 - 6.4 Diabetic: >= 6.5 | | |
| INTERPRETATION: If Diabetes - Good control : 6.1 - 7.0 %, Fair control : 7.1 - 8.0 %, Poor control >= 8.1 % | | | | | |

| Estimated Average Glucose | 225.95 | mg/dL |
|---------------------------|--------|-------|
| e | | • |

(Whole Blood)

INTERPRETATION: Comments

HbA1c provides an index of Average Blood Glucose levels over the past 8 - 12 weeks and is a much better indicator of long term glycemic control as compared to blood and urinary glucose determinations.

Conditions that prolong RBC life span like Iron deficiency anemia, Vitamin B12 & Folate deficiency,

hypertriglyceridemia, hyperbilirubinemia, Drugs, Alcohol, Lead Poisoning, Asplenia can give falsely elevated HbA1C values.

Conditions that shorten RBC survival like acute or chronic blood loss, hemolytic anemia, Hemoglobinopathies, Splenomegaly, Vitamin E ingestion, Pregnancy, End stage Renal disease can cause falsely low HbA1c.



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| Investigation | <u>Observed</u> Value | <u>Unit</u> | Biological Reference Interval | | |
|---|--------------------------|-----------------------|--------------------------------------|--|--|
| IMMUNOASSAY | | | | | |
| THYROID PROFILE / TFT | | | | | |
| T3 (Triiodothyronine) - Total (Serum/ECLIA) | 1.39 | ng/ml | 0.4 - 1.81 | | |
| INTERPRETATION: Comment : Total T3 variation can be seen in other condition like preg Metabolically active. | gnancy, drugs, neph | rosis etc. In such ca | ses, Free T3 is recommended as it is | | |
| T4 (Tyroxine) - Total (Serum/ <i>ECLIA</i>) | 8.70 | µg/dl | 4.2 - 12.0 | | |
| INTERPRETATION: Comment : Total T4 variation can be seen in other condition like pregnancy, drugs, nephrosis etc. In such cases, Free T4 is recommended as it is Metabolically active. | | | | | |
| TSH (Thyroid Stimulating Hormone) (Serum/ECLIA) | 1.91 | µIU/mL | 0.35 - 5.50 | | |
| INTERPRETATION: Reference range for cord blood - upto 20 1 st trimester: 0.1-2.5 2 nd trimester 0.2-3.0 3 rd trimester : 0.3-3.0 (Indian Thyroid Society Guidelines) Comment : 1.TSH reference range during pregnancy depends on Iodi 2.TSH Levels are subject to circadian variation, reaching | | | | | |
| 2.TSH Levels are subject to circadian variation, reaching peak levels between 2-4am and at a minimum between 6-10PM. The variation can be of the order of 50%, hence time of the day has influence on the measured serum TSH concentrations. | | | | | |

3.Values&lt;0.03 µIU/mL need to be clinically correlated due to presence of rare TSH variant in some individuals.



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|--|---|--|
| CLINICAL PATHOLOGY | | |
| <u>PHYSICAL EXAMINATION (URINE</u> <u>COMPLETE)</u> | | |
| Colour (Urine) | Pale Yellow | Yellow to Amber |
| Appearance (Urine) | Clear | Clear |
| Volume(CLU) (Urine) | 20 | |
| <u>CHEMICAL EXAMINATION (URINE</u> <u>COMPLETE)</u> | | |
| pH (Urine) | 5.5 | 4.5 - 8.0 |
| Specific Gravity (Urine) | 1.025 | 1.002 - 1.035 |
| Ketone (Urine) | Negative | Negative |
| Urobilinogen (Urine) | Normal | Normal |
| Blood (Urine) | Negative | Negative |
| Nitrite (Urine) | Negative | Negative |
| Bilirubin (Urine) | Negative | Negative |
| Protein (Urine) | Negative | Negative |



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The results pertain to sample tested.

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|--|---------------------------------|-------------|---|
| Glucose (Urine/GOD - POD) | Negative | | Negative |
| Leukocytes(CP) (Urine) | Negative | | |
| <u>MICROSCOPIC EXAMINATION</u> (URINE COMPLETE) | | | |
| Pus Cells (Urine) | 0-1 | /hpf | NIL |
| Epithelial Cells (Urine) | 0-1 | /hpf | NIL |
| RBCs (Urine) | NIL | /HPF | NIL |
| Others (Urine) | NIL | | |

INTERPRETATION: Note: Done with Automated Urine Analyser & Automated urine sedimentation analyser. All abnormal reports are reviewed and confirmed microscopically.



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Investigation

<u>Observed</u> <u>Value</u> <u>Unit</u>

Biological Reference Interval

IMMUNOHAEMATOLOGY

BLOOD GROUPING AND Rh TYPING (EDTA Blood/Agglutination)

'O' 'Positive'

INTERPRETATION: Note: Slide method is screening method. Kindly confirm with Tube method for transfusion.



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|---|---------------------------------|-------------|--|
| BIOCHEMISTRY | | | |
| BUN / Creatinine Ratio | 11 | | 6.0 - 22.0 |
| Glucose Fasting (FBS) (Plasma - F/GOD-PAP) | 145.46 | mg/dL | Normal: < 100 Pre Diabetic: 100 - 125 Diabetic: >= 126 |

INTERPRETATION: Factors such as type, quantity and time of food intake, Physical activity, Psychological stress, and drugs can influence blood glucose level.

| Glucose, Fasting (Urine) | Negative | | Negative |
|-----------------------------|----------|-------|----------|
| (Urine - F/GOD - POD) | | | |
| Glucose Postprandial (PPBS) | 190.16 | mg/dL | 70 - 140 |
| (Plasma - PP/GOD-PAP) | | | |

INTERPRETATION:

Factors such as type, quantity and time of food intake, Physical activity, Psychological stress, and drugs can influence blood glucose level. Fasting blood glucose level may be higher than Postprandial glucose, because of physiological surge in Postprandial Insulin secretion, Insulin resistance, Exercise or Stress, Dawn Phenomenon, Somogyi Phenomenon, Anti- diabetic medication during treatment for Diabetes.

| Urine Glucose(PP-2 hours) (Urine - PP) | Trace | Negative |
|--|------------|-----------|
| Blood Urea Nitrogen (BUN) (Serum/Urease UV / derived) | 12.9 mg/dL | 7.0 - 21 |
| Creatinine | 0.90 mg/dL | 0.9 - 1.3 |

(Serum/Modified Jaffe)

INTERPRETATION: Elevated Creatinine values are encountered in increased muscle mass, severe dehydration, Pre-eclampsia, increased ingestion of cooked meat, consuming Protein/ Creatine supplements, Diabetic Ketoacidosis, prolonged fasting, renal dysfunction and drugs such as cefoxitin ,cefazolin, ACE inhibitors ,angiotensin II receptor antagonists,N-acetylcyteine , chemotherapeutic agent such as flucytosine etc.

| Uric Acid | 3.57 | mg/dL | 3.5 - 7.2 |
|-------------------|------|-------|-----------|
| (Serum/Enzymatic) | | | |



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|--|---------------------------------|-------------|--|
| IMMUNOASSAY | | | |
| Prostate specific antigen - Total(PSA) (Serum/ <i>Manometric method</i>) | 0.467 | ng/ml | Normal: 0.0 - 4.0 Inflammatory & Non Malignant conditions of Prostate & genitourinary system: 4.01 - 10.0 Suspicious of Malignant disease of Prostate: > 10.0 |

-- End of Report --



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DR SHAMIM JAVED MD PATHOLOGY KMC 88902

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