Patient Name : MR. V. BABU JAGADESH Age / Gender : 35 years / Male

Patient ID: 8362

Source : MEDI WHEEL

Referral : SELF

Sample ID :

Collection Time : Aug 19, 2022, 09:45 a.m.

Reporting Time : Aug 19, 2022, 11:16 a.m.



| Test Description | Value(s) | Reference Range | Unit |
|---|----------|-----------------|------------|
| CBC; Complete Blood Count | | | |
| Hemoglobin (Hb)* Method : Cynmeth Photometric Measurement | 14.2 | 13.5 - 18.0 | gm/dL |
| Erythrocyte (RBC) Count* Method : Electrical Impedence | 5.0 | 4.7 - 6.0 | mil/cu.mm |
| Packed Cell Volume (PCV)* Method : Calculated | 44 | 42 - 52 | % |
| Mean Cell Volume (MCV)* Method : Electrical Impedence | 88 | 78 - 100 | fL |
| Mean Cell Haemoglobin (MCH)* Method : Calculated | 28 | 27 - 31 | pg |
| Mean Corpuscular Hb Concn. (MCHC)* Method : Calculated | 32 | 32 - 36 | gm/dL |
| Red Cell Distribution Width (RDW)* Method : Electrical Impedence | 14.8 | 11.5 - 14.0 | % |
| Total Leucocytes (WBC) Count* Method : Electrical Impedence | 6600 | 4000-10000 | cell/cu.mm |
| Neutrophils* Method : VCSn Technology | 53 | 40 - 80 | % |
| Lymphocytes* Method : VCSn Technology | 40 | 20 - 40 | % |
| Monocytes* Method : VCSn Technology | 5 | 2 - 10 | % |
| Eosinophils* Method : VCSn Technology | 2 | 1 - 6 | % |
| Basophils | 0 | 0 - 1 | |
| Platelet Count* Method : Electrical Impedence | 3.12 | 1.5 - 4.5 | 10^3/ul |
| Mean Platelet Volume (MPV)* Method : Electrical Impedence | 7.3 | 7.2 - 11.7 | f∟ |
| | | | |

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Reporting Time : Aug 19, 2022, 11:16 a.m.

Sample ID :

| | | | 669327321 | |
|---------------------|----------|-----------------|-----------|--|
| Test Description | Value(s) | Reference Range | Unit | |
| PCT* | 0.23 | 0.2 - 0.5 | % | |
| Method : Calculated | | | | |
| PDW* | 15.4 | 9.0 - 17.0 | % | |
| Method : Calculated | | | | |

Tests done on Automated Three Part Cell Counter. (WBC, RBC, Platelet count by impedance method, colorimetric method for Hemoglobin, WBC differential by flow cytometry using laser technology other parameters are calculated). All Abnormal Haemograms are reviewed confirmed microscopically.

| Esr, Erythrocyte Sedimentation Rate | | | |
|---|----|------|-------|
| Esr, Erythrocyte Sedimentation Rate (Westergren) | 21 | 0-10 | mm/hr |
| Interpretation: | | | |

- It indicates presence and intensity of an inflammatory process. It does not diagnose a specific disease. Changes in the ESR are . more significant than the abnormal results of a single test.
- It is a prognostic test and used to monitor the course or response to treatment of diseases like tuberculosis, bacterial ٠ endocarditis, acute rheumatic fever, rheumatoid arthritis, SLE, Hodgkins disease, temporal arteritis and polymyalgia rheumatica.
- It is also increased in pregnancy, multiple myeloma, menstruation, and hypothyroidism. ٠

| Urine Routine | | |
|--------------------------------------|--------------------|---------------|
| Colour* | Pale Yellow | Pale Yellow |
| Transparency (Appearance)* | Clear | Clear |
| Deposit* | Absent | Absent |
| Reaction (pH)* | 5.0 | 4.5 - 8 |
| Specific Gravity* | 1.025 | 1.010 - 1.030 |
| Chemical Examination (Automated Dips | tick Method) Urine | |
| Urine Glucose (sugar)* | Absent | Absent |
| Urine Protein (Albumin)* | Absent | Absent |
| Urine Ketones (Acetone)* | Absent | Absent |
| | | |

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Source : MEDI WHEEL

Sample ID :

| | | 669327321 | | |
|-------------------------------|----------|-----------------|------|--|
| Test Description | Value(s) | Reference Range | Unit | |
| | | | | |
| Blood* | Absent | Absent | | |
| Bile pigments* | Absent | Absent | | |
| Nitrite* | Absent | Absent | | |
| Urobilinogen* | Normal | Normal | | |
| Microscopic Examination Urine | | | | |
| Pus Cells (WBCs)* | 4-5 | 0 - 5 | /hpf | |
| Epithelial Cells* | 1-2 | 0 - 4 | /hpf | |
| Red blood Cells* | Absent | Absent | /hpf | |
| Crystals* | Absent | Absent | | |
| Cast* | Absent | Absent | | |
| Trichomonas Vaginalis* | Absent | Absent | | |
| Yeast Cells* | Absent | Absent | | |
| Amorphous deposits* | Absent | Absent | | |
| Bacteria* | Absent | Absent | | |

Blood Group & Rh Type

Blood Grouping & Rh Typing Method : Forward and Reverse By Tube Method

"O" + (POSITIVE)

Methodology

This is done by forward and reverse grouping by tube Agglutination method.

Interpretation

Newborn baby does not produce ABO antibodies until 3 to 6 months of age. So the blood group of the Newborn baby is done by ABO antigen grouping (forward grouping) only, antibody grouping (reverse grouping) is not required.Confirmation of the New-born's blood group is indicatedwhen the A and B antigen expression and the isoagglutinins are fully developed (2–4 years).

Fasting - Glucose

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| | | 60 | 9327321 |
|----------------------------------|----------|---|---------|
| Test Description | Value(s) | Reference Range | Unit |
| Glucose Fasting* | 70 | Normal: 70-100 | mg/dL |
| Method : Plasma, Hexokinase | | Impaired Fasting Glucose (IFG): | - |
| | | 100-125 | |
| | | Diabetes Mellitus: >= 126 | |
| | | (On more than one occasion) (American Diabetes Association | |
| | | guidelines 2017) | |
| | | | |
| Post Prandial Blood Sugar | | | |
| Blood Glucose-Post Prandial* | 120 | 80-140 | mg/dL |
| Method : Plasma - P, Hexokinase | | | |
| Fasting Urine Sugar | | | |
| Fasting Urine Sugar | NEGATIVE | NEGATIVE - | |
| Post Prandial Urine Sugar | | | |
| Post Prandial Urine Sugar | NEGATIVE | | |
| HBA1C (Glycosylated Haemoglobin) | | | |
| Glyco Hb (HbA1C) | 5.3 | Non-Diabetic: <=5.9 | % |
| Method : EDTA Whole blood, HPLC | | Pre Diabetic:6.0-6.4 | |
| | | Diabetic: >=6.5 | |
| Estimated Average Glucose : | 105 | | mg/dL |
| Interpretations | | | |

1. HbA1C has been endorsed by clinical groups and American Diabetes Association guidelines 2017 for diagnosing diabetes using a cut off point of 6.5%

2. Low glycated haemoglobin in a non diabetic individual are often associated with systemic inflammatory diseases, chronic anaemia (especially severe iron deficiency and haemolytic), chronic renal failure and liver diseases. Clinical correlation suggested.

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| Test Description | Value(s) | Reference Range | Unit |
| 3. In known diabetic patients, following values c | an be considered a | as a tool for monitoring the glycemic con | trol. |
| Excellent control-6-7 % | | | |
| Fair to Good control – 7-8 % | | | |
| Unsatisfactory control – 8 to 10 % | | | |
| Poor Control – More than 10 % | | | |
| Thyroid Function Test (TFT) | | | |
| THYROID STIMULATING HORMONE (TSH) | 1.6 | 0.46 – 8.10 : 1 Yrs – 5 Yrs | ulU/mL |
| Method : CLIA | | 0.36 – 5.80 : 6 Yrs – 18 Yrs | |
| | | 0.35 – 5.50 : 18 Yrs – 55 Yrs | |
| | | 0.50 – 8.90 : >55 Yrs | |
| | | Pregnancy Ranges:::: | |
| | | lst Tri :0.1 - 2.5 | |
| | | lind Tri :0.2 - 3.0 | |
| | | llird Tri:0.3 - 3.0 | |
| TOTAL TRIIODOTHYRONINE (T3) | 158 | 126 – 258 : 1 Yr – 5 Yr | ng/dl |
| Method : CLIA | | 96 – 227 : 6 Yr – 15 Yr | |
| | | 91 – 164 : 16 Yr – 18 Yr | |
| | | 60 – 181 : > 18 Years | |
| | | Pregnancy : | |
| | | 1st Trimester : 81 - 190 | |
| | | 2nd & 3rd Trimester:100 - 260 | <i>.</i> |
| TOTAL THYROXINE (T4) | 9.0 | | µg/dL |
| Method : CLIA | | 4.6 - 10.9 Decementaria | |
| | | Pregnancy: | |
| | | 4.6 – 16.5 : 1st Trimester | |
| | | 4.6 – 18.5 : 2nd & 3rd Tri | |

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|---|--|--|--|
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| Patient ID: 8362 | Reporting Time : Aug 19, 2 | 2022, 11:16 a.m. | |
| Source : MEDI WHEEL | | Sample ID : | 9327321 |
| Test Description | Value(s) | Reference Range | Unit |
| Comments: | IF NOT ON DRUGS SUGGESTED FT3 & FT4 ESTIMATION Please correlate with clinical conditions. Note : Serum T3, T4 and TSH form the three components of thyroid scre panel, useful in diagnosing various disorders of the thyroid gland. Primary Hypothyroidism is accompanied by depressed serum T3 and T4 values a elevated serum TSH levels. Although elevated TSH levels are nearly alwa indicative of Primary Hypothyroidism, rarely they can from TSH secreting pituitary tumors (Secondary hyperthyroidism)To confirm diagnosis - evalu FT3 and FT4. | | nts of thyroid screening oid gland. Primary 3 and T4 values and els are nearly always om TSH secreting |
| Cholesterol-Total Method : Serum, Cholesterol oxidase esterase, peroxidase | 138 | Desirable: <= 200 Borderline High: 201-239 High: > 239 Ref: The National Cholesterol Education Program (NCEP) Adult Treatment Panel III Report. | mg/dL |
| Triglycerides Method : Serum, Enzymatic, endpoint | 206 | Normal: < 150 Borderline High: 150-199 High: 200-499 Very High: >= 500 | mg/dL |
| Cholesterol-HDL Direct Method : Serum, Direct measure-PEG | 43 | Normal: > 40 Major Heart Risk: < 40 | mg/dL |
| LDL Cholesterol Method : Serum | 53.8 | Optimal: < 100 Near optimal/above optimal: 100-12 Borderline high: 130-159 High: 160-189 Very High: >= 190 | mg/dL 29 |

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| Test Description | Value(s) | Reference Range | Unit |
|------------------------------|----------|---------------------------------|-------|
| Non - HDL Cholesterol, Serum | 95 | Desirable: < 130 mg/dL | mg/dL |
| Method : calculated | | Borderline High: 130-159mg/dL | 5 |
| | | High: 160-189 mg/dL | |
| | | Very High: > or = 190 mg/dL | |
| VLDL Cholesterol | 41.2 | 6 - 38 | mg/dL |
| Method : calculated | | | |
| CHOL/HDL RATIO | 3.2 | 3.5 - 5.0 | ratio |
| Method : calculated | | | |
| LDL/HDL RATIO | 1.25 | Desirable / low risk - 0.5 -3.0 | ratio |
| Method : calculated | | Low/ Moderate risk - 3.0- 6.0 | |
| | | Elevated / High risk - > 6.0 | |
| HDL/LDL RATIO | 0.8 | Desirable / low risk - 0.5 -3.0 | ratio |
| Method : calculated | | Low/ Moderate risk - 3.0- 6.0 | |
| | | Elevated / High risk - > 6.0 | |

| KIDNEY FUNCTION TEST | | | |
|-----------------------------------|-----|-----------|-------|
| Urea * | 19 | 15- 50 | mg/dL |
| Method : Serum | | | |
| Blood Urea Nitrogen-BUN* | 8.8 | 7 - 24 | mg/dL |
| Method : Serum, Urease | | | |
| Uric Acid* | 4.9 | 3.5 - 7.2 | mg/dL |
| Method : Serum, Uricase/POD | | | |
| Creatinine* | 0.7 | 0.6 - 1.1 | mg/dL |
| Method : Serum, Jaffe IDMS | | | |
| Liver Funtion Test (LFT) with GGT | | | |
| Bilirubin - Total | 1.0 | 0.3 - 1.2 | mg/dL |

Method : Serum, Jendrassik Grof

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| Test Description | Value(s) | Reference Range | Unit |
|--|----------|----------------------------|-------------|
| | | | <i>(</i>), |
| Bilirubin - Direct | 0.4 | Adults and Children: < 0.2 | mg/dL |
| Method : Serum, Diazotization | 0.0 | 0.1.1.0 | and as (all |
| Bilirubin - Indirect | 0.6 | 0.1 - 1.0 | mg/dL |
| Method : Serum, Calculated | 40 | 50 | 11/1 |
| SGOT | 40 | < 50 | U/L |
| Method : Serum, UV with P5P, IFCC 37 degree | 25 | . 50 | 11/1 |
| SGPT | 35 | < 50 | U/L |
| Method : Serum, UV with P5P, IFCC 37 degree | 1.14 | 0.7 - 1.4 | ratio |
| SGOT/SGPT Method : calculated | 1.14 | 0.7 - 1.4 | ralio |
| GGT-Gamma Glutamyl Transpeptidae | 26 | < 55 | U/L |
| Method : Serum, G-glutamyl-carboxy-nitoanilide | 20 | < 55 | 0/L |
| Alkaline Phosphatase-ALPI | 105 | 30-120 | U/L |
| Method : Serum, PNPP, AMP Buffer, IFCC 37 degree | 100 | 30-120 | 0/2 |
| Total Protein | 7.0 | 6.6 - 8.3 | g/dL |
| Method : Serum, Biuret, reagent blank end point | 1.0 | 0.0 0.0 | 9,42 |
| Albumin | 4.0 | Adults: 3.5 - 5.2 | g/dL |
| Method : Serum, Bromcresol purple | | | 9, 42 |
| Globulin | 3.0 | 1.8 - 3.6 | g/dL |
| Method : Calculated | | | J |
| A/G Ratio | 1.33 | 1.2 - 2.2 | ratio |
| Method : Calculated | | | |

END OF REPORT

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