

Patient Name : MRS. SUMATHI R Age / Gender : 49 years / Female Patient ID : 74746

Referral : MediWheel

Collection Time : Dec 26, 2021, 08:21 a.m.

Reporting Time : Dec 26, 2021, 03:36 p.m.

Sample ID :



| | | | 001036021 |
|-----------------------------------|----------|-----------------|------------|
| Test Description | Value(s) | Reference Range | |
| COMPLETE BLOOD COUNT (CBC) | | | |
| ESR | 20 | 12.0 - 15.0 | mm/hr |
| Hemoglobin (Hb) | 14.8 | 12.0 - 15.0 | gm/dL |
| Erythrocyte (RBC) Count | 5.42 | 3.8 - 4.8 | mil/cu.mm |
| Packed Cell Volume (PCV) | 42.5 | 36 - 46 | % |
| Mean Cell Volume (MCV) | 78.41 | 83 - 101 | fL |
| Mean Cell Haemoglobin (MCH) | 27.31 | 27 - 32 | pg |
| Mean Corpuscular Hb Concn. (MCHC) | 34.82 | 31.5 - 34.5 | g/dL |
| Red Cell Distribution Width (RDW) | 12.7 | 11.6 - 14.0 | % |
| Total Leucocytes (WBC) Count | 4800 | 4000-10000 | cell/cu.mm |
| Neutrophils | 66 | 40 - 80 | % |
| Lymphocytes | 28 | 20 - 40 | % |
| Monocytes | 5 | 2 - 10 | % |
| Eosinophils | 1 | 1 - 6 | % |
| Basophils | 0 | 1-2 | % |
| Absolute Neutrophil Count | 3168 | 2000 - 7000 | /c.mm |
| Absolute Lymphocyte Count | 1344 | 1000 - 3000 | /c.mm |
| Absolute Monocyte Count | 240 | 200 - 1000 | /c.mm |
| Absolute Eosinophil Count | 48 | 20 - 500 | /c.mm |
| Absolute Basophils Count | 0 | 20 - 100 | /c.mm |
| Platelet Count | 190 | 150 - 410 | 10^3/ul |
| Mean Platelet Volume (MPV) | 10.2 | 7.2 - 11.7 | fL |
| PCT | 0.19 | 0.2 - 0.5 | % |
| PDW | 10.8 | 9.0 - 17.0 | % |

END OF REPORT

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URINE COMPLETE ANALYSIS,

| | Physical Exa | mination | |
|-------------------|------------------|--------------------|----|
| Quantity | 26 | | ml |
| Colour | Pale Yellow | Pale yellow/Yellow | |
| Appearance | Clear | Clear | |
| Specific Gravity | 1.020 | 1.005-1.025 | |
| рН | 6.5 | 5.0 - 8.0 | |
| Deposit | Present | Absent | |
| | Chemical Exa | mination | |
| Protein | Absent | Absent | |
| Sugar | Present (++) | Absent | |
| Ketones | Absent | Absent | |
| Bile Salt | Absent | Absent | |
| Bile Pigment | Absent | Absent | |
| Urobilinogen | Normal | Normal | |
| | Microscopic Exam | ination (/hpf) | |
| Pus Cell | 2-4 | Upto 5 | |
| Epithelial Cells | 1-2 | Upto 5 | |
| Red Blood Cells | Absent | Absent | |
| Casts | Absent | Absent | |
| Crystals | Absent | Absent | |
| Amorphous Deposit | Absent | Absent | |
| Yeast Cells | Absent | Absent | |
| Bacteria | Absent | Absent | |
| Other findings | Not seen | Not seen | |

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END OF REPORT

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| Test Description | Value(s) | Reference Range | |
| STOOL ROUTINE ANALYSIS | | | |
| Color | Brownish | Brown | |
| Consistency | Semisolid | Solid - Semi solid | |
| Reaction (pH) | Acidic | Acidic - Alkaline | |
| Method : Methyl Red & Bromothymol Blue | | | |
| Mucous | Absent | Absent | |
| Blood | Absent | Absent | |
| Pus cells | 2-3/hpf | Few | /hpf |
| Epithelial cells | 1-2/hpf | | /hpf |
| RBC | Absent | Absent | /hpf |
| Ova | Not found | Absent | /hpf |
| Cyst | Not found | Absent | /hpf |
| Starch granules | Absent | None to small amount | /hpf |
| Vegetable cells | Absent | | /hpf |
| Fat globules | Absent | Absent | /hpf |
| Others Method : Microscopy (Concentration technique) | Nil | | /hpf |

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| | | |
| Test Description | Value(s) | Reference Range |
| BLOOD GROUP & RH TYPING | | |
| Blood Group (ABO typing) Method : Manual-Hemagglutination | "B" | |
| RhD Factor (Rh Typing) Method : Manual hemagglutination | Positive | |

END OF REPORT

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| | | Sample ID : | 001036021 |
| Test Description | Value(s) | Reference Range | |
| Glycosylated HbA1c | | | |
| HbA1c (GLYCOSYLATED HEMOGLOBIN), BLOOD | 11.0 | | % |
| Method : (HPLC, NGSP certified) | | | |
| Estimated Average Glucose : | 269 | - | mg/dL |
| Interpretation | | | |
| As per American Diabetes Association (ADA) | | | |
| Reference Group | HbA1c in % | | |
| Non diabetic adults >=18 years | <5.7 | | |
| At risk (Prediabetes) | 5.7 - 6.4 | | |
| Diagnosing Diabetes | >= 6.5 | | |
| Therapeutic goals for glycemic control | Age > 19 years Goal of therapy: Action suggeste | | |
| Note: | Age < 19 years Goal of therapy: | <7.5 | |
| | | | |

- 1. Since HbA1c reflects long term fluctuations in the blood glucose concentration, a diabetic patient who is recently under good control may still have a high concentration of HbA1c. Converse is true for a diabetic previously under good control but now poorly controlled .
- 2. Target goals of < 7.0 % may be beneficial in patients with short duration of diabetes, long life expectancy and no significant cardiovascular disease. In patients with significant complications of diabetes, limited life expectancy or extensive co-morbid conditions, targeting a goal of < 7.0 % may not be appropriate.

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.

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Test Description

Value(s)

Reference Range

ADA criteria for correlation between HbA1c & Mean plasma glucose levels.

| HbA1c(%) | Mean Plasma Glucose (mg/dL) |
|----------|-----------------------------|
| 6 | 126 |
| 7 | 154 |
| 8 | 183 |
| 9 | 212 |
| 10 | 240 |
| 11 | 269 |
| 12 | 298 |

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Sample ID :

Third trimester : 0.3-3.0

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| | | 001036021 | |
|----------------------------|-----------|---------------------------|--------|
| Test Description | Value(s) | Reference Range | |
| THYROID PROFILE TEST - TOT | <u>4L</u> | | |
| T3-Total | 99.26 | 60 - 200 | ng/dL |
| T4-Total | 8.40 | 4.52 - 12 | ug/dL |
| TSH-Ultrasensitive | 3.68 | 0.32 - 5.5 | ulU/mL |
| Method : CLIA | | First Trimester : 0.1-2.5 | |
| | | Second Trimester : 0.2-3 | 3.0 |

Interpretation

| TSH | Т3 | Т4 | Suggested Interpretation for the Thyroid Function Tests Pattern |
|--------------------------|------------------------|------------------------|--|
| Raised | Within range | Within range | Raised Within Range Within Range .Isolated High TSHespecially in the range of 4.7 to 15 m1U/m1 is commonly associated with Physiological & Biological TSH Variability. Subclinical Autoimmune Hypothyroidism.Intermittent 14 therapy for hypothyroidism .Recovery phase after Non-Thyroidal illness" |
| Raised | Decreased | Decreased | Chronic Autoimmune Thyroiditis Post thyroidectomy, Post radioiodine Hypothyroid phase of transient thyroiditis" |
| Raised or within range | Raised | Raised or within range | Interfering antibodies to thyroid hormones (anti-TPO antibodies)Intermittent 14 therapy or T4 overdose •Drug interference- Amiodarone, Heparin,Beta blockers,steroids, anti-epileptics. |
| Decreased | Raised or within range | Raised or within range | Isolated Low TSH -especially in the range of 0.1 to 0.4 often seen in elderly & Range Range associated with Non-Thyroidal illness .Subclinical Hyperthyroidism .Thyroxine ingestion' |
| Decreased | Decreased | Decreased | Central Hypothyroidism .Non-Thyroidal illness .Recent treatment for Hyperthyroidism (TSH remains suppressed)" |
| Decreased | Raised | Raised | Primary Hyperthyroidism (Graves' disease).Multinodular goitre, Toxic nodule •Transient thyroiditis:Postpartum, Silent (lymphocytic), Postviral (granulomatous,subacute, DeQuervain's),Gestational thyrotoxicosis with hyperemesis gravidarum" |
| Decreased Within Rang | Raised | Within range | T3 toxicosis •Non-Thyroidal illness |
| Within range | Decreased | Within range | Isolated Low T3-often seen in elderly & associated Non-Thyroidal illness In elderly the drop in 13 level can be upto 25%. |

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Scan to Validate





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| Test Description | Value(s) | Reference Range | |
|--|----------|---|-------|
| LIPID PROFILE | | | |
| Cholesterol-Total Method : Spectrophotometry | 261 | Desirable level < 200 Borderline High 200-239 High >or = 240 | mg/dL |
| Triglycerides Method : Serum, Enzymatic, endpoint | 300 | Normal: < 150 Borderline High: 150-199 High: 200-499 Very High: >= 500 | mg/dL |
| HDL Cholesterol Method : Serum, Direct measure-PEG | 44 | Normal: > 40 Major Risk for Heart: < 40 | mg/dL |
| LDL Cholesterol Method : Enzymatic selective protection | 157 | Optimal < 100 Near / Above Optimal 100-129 Borderline High 130-159 High 160-189 Very High >or = 190 | mg/dL |
| VLDL Cholesterol Method : Serum, Enzymatic | 60 | 6 - 38 | mg/dL |
| CHOL/HDL Ratio Method : Serum, Enzymatic | 5.93 | 3.5 - 5.0 | |
| LDL/HDL Ratio Method : Serum, Enzymatic Note: | 3.57 | 2.5 - 3.5 | |
| 8-10 hours fasting sample is required. | | | |

END OF REPORT

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| Test Description | Value(s) | Reference Range | |
| RENAL PROFILE | | | |
| Urea Method : Uricase | 21 | 15-36 | mg/dL |
| Blood Urea Nitrogen-BUN Method : Serum, Urease | 9.80 | 7 - 17 | mg/dL |
| Creatinine Method : Serum, Jaffe | 0.7 | 0.52-1.25 | mg/dL |
| Uric Acid Method : Serum, Uricase | 5.9 | 2.5 - 6.2 | mg/dL |
| Remark: | | | |

In blood, Urea is usually reported as BUN and expressed in mg/dl. BUN mass units can be converted to urea mass units by multiplying by 2.14.

END OF REPORT

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| Test Description | Value(s) | Reference Range | | |
|---|----------|-----------------|----------|--|
| | | | | |
| LIVER FUNCTION TEST | | | <i>.</i> | |
| Total Protein | 8.1 | 6.3 - 8.2 | g/dL | |
| Method : Serum, Biuret, reagent blank end point | | | <i></i> | |
| Albumin | 4.9 | 3.5 - 5.0 | g/dL | |
| Method : Serum, Bromocresol green | | | | |
| Globulin | 3.20 | 1.8 - 3.6 | g/dL | |
| Method : Serum, EIA | | | | |
| A/G Ratio | 1.53 | 1.2 - 2.2 | | |
| Method : Serum, EIA | | | | |
| Bilirubin - Total | 0.3 | 0.2 - 1.3 | mg/dL | |
| Method : Serum, Jendrassik Grof | | | | |
| Bilirubin - Direct | 0.2 | < 0.3 | mg/dL | |
| Method : Serum, Diazotization | | | | |
| Bilirubin - Indirect | 0.1 | 0.0 - 1.1 | mg/dL | |
| Method : Serum, Calculated | | | | |
| SGOT | 17 | 14-36 | U/L | |
| Method : Serum, UV with P5P, IFCC 37 degree | | | | |
| SGPT | 14 | < 52 | U/L | |
| Method : Serum, UV with P5P, IFCC 37 degree | | | | |
| Alkaline Phosphatase | 104 | 38 - 126 | U/L | |
| Method : PNPP-AMP Buffer/Kinetic | | | | |
| GGT-Gamma Glutamyl Transpeptidae | 18 | < 43 | U/L | |
| Method : Serum, G-glutamyl-carboxy-nitoanilide | | | | |
| | | | | |

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| Test Description | Value(s) | Reference Range | | |
| <u>GLUCOSE (F)</u> | | | | |
| Glucose fasting Method : GOD-POD | 250 | Normal: 70 - 120 | mg/dL | |
| | | | | |

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| Test Description | Value(s) | Reference Range | |
| GLUCOSE (PP) | | | |
| Blood Glucose-Post Prandial Method : GOD-POD | 402 | 80- 140 mg/dL | |
| | | | |

END OF REPORT

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