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CID : 2132924997
Name : MR.AKASH DUBE
Age / Gender : 30 Years / Male
Consulting Dr. : -
Reg. Location : J B Nagar, Andheri East (Main Centre)

Collected : 26-Nov-2021 / 12:57
Reported : 26-Nov-2021 / 16:53

ARCOFEMI HEALTHCARE- BLOOD TEST

CBC (Complete Blood Count), Blood

| PARAMETER | RESULTS | BIOLOGICAL REF RANGE | METHOD |
|---|---------|----------------------|--------------------|
| <u>RBC PARAMETERS</u> | | | |
| Haemoglobin | 15.5 | 13.0-17.0 g/dL | Spectrophotometric |
| RBC | 4.38 | 4.5-5.5 mil/cmm | Elect. Impedance |
| PCV | 46.1 | 40-50 % | Measured |
| MCV | 105.2 | 80-100 fl | Calculated |
| MCH | 35.3 | 27-32 pg | Calculated |
| MCHC | 33.6 | 31.5-34.5 g/dL | Calculated |
| RDW | 18.4 | 11.6-14.0 % | Calculated |
| <u>WBC PARAMETERS</u> | | | |
| WBC Total Count | 10410 | 4000-10000 /cmm | Elect. Impedance |
| <u>WBC DIFFERENTIAL AND ABSOLUTE COUNTS</u> | | | |
| Lymphocytes | 36.1 | 20-40 % | |
| Absolute Lymphocytes | 3758.0 | 1000-3000 /cmm | Calculated |
| Monocytes | 7.2 | 2-10 % | |
| Absolute Monocytes | 749.5 | 200-1000 /cmm | Calculated |
| Neutrophils | 51.0 | 40-80 % | |
| Absolute Neutrophils | 5309.1 | 2000-7000 /cmm | Calculated |
| Eosinophils | 5.1 | 1-6 % | |
| Absolute Eosinophils | 530.9 | 20-500 /cmm | Calculated |
| Basophils | 0.6 | 0.1-2 % | |
| Absolute Basophils | 62.5 | 20-100 /cmm | Calculated |
| Immature Leukocytes | - | | |
| WBC Differential Count by Absorbance & Impedance method/Microscopy. | | | |
| <u>PLATELET PARAMETERS</u> | | | |
| Platelet Count | 219000 | 150000-400000 /cmm | Elect. Impedance |
| MPV | 9.5 | 6-11 fl | Calculated |
| PDW | 14.1 | 11-18 % | Calculated |
| <u>RBC MORPHOLOGY</u> | | | |
| Hypochromia | - | | |
| Microcytosis | - | | |
| Macrocytosis | + | | |



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| | |
|----------------------|---|
| Anisocytosis | + |
| Poikilocytosis | Mild |
| Polychromasia | - |
| Target Cells | - |
| Basophilic Stippling | - |
| Normoblasts | - |
| Others | - |
| WBC MORPHOLOGY | - |
| PLATELET MORPHOLOGY | Platelet count may not be representative due to presence of platelet aggregates |
| COMMENT | Leucocytosis |

Result rechecked.
Kindly correlate clinically.
Advice : Repeat estimation with a fresh EDTA/Citrate whole blood sample for accurate platelet count.

Specimen: EDTA Whole Blood

ESR, EDTA WB-ESR 26 2-15 mm at 1 hr. Westergren

*Sample processed at SUBURBAN DIAGNOSTICS (INDIA) PVT. LTD CPL, Andheri West
*** End Of Report ***



Amar Dasgupta
Dr. AMAR DASGUPTA, MD, PhD
Consultant Hematopathologist
Director - Medical Services

M. Millu Jain
Dr. MILLU JAIN
M.D.(PATH)
Pathologist

ADDRESS: 2nd Floor, Aston, Sundervan Complex, Above Mercedes Showroom, Andheri West - 400053

HEALTHLINE - MUMBAI: 022-6170-0000 | OTHER CITIES: 1800-266-4343

For Feedback - customerservice@suburbandiagnosics.com | www.suburbandiagnosics.com



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ARCOFEMI HEALTHCARE- BLOOD TEST

| <u>PARAMETER</u> | <u>RESULTS</u> | <u>BIOLOGICAL REF RANGE</u> | <u>METHOD</u> |
|--|----------------|---|---------------|
| GLUCOSE (SUGAR) PP, Fluoride Plasma PP/R | 87.2 | Non-Diabetic: < 140 mg/dl Impaired Glucose Tolerance: 140-199 mg/dl Diabetic: >/= 200 mg/dl | Hexokinase |
| TOTAL PROTEINS, Serum | 8.4 | 6.4-8.3 g/dL | Biuret |
| ALBUMIN, Serum | 4.8 | 3.5-5.2 g/dL | BCG |
| GLOBULIN, Serum | 3.6 | 2.3-3.5 g/dL | Calculated |
| A/G RATIO, Serum | 1.3 | 1 - 2 | Calculated |
| BLOOD UREA, Serum | 22.4 | 12.8-42.8 mg/dl | Kinetic |
| BUN, Serum | 10.5 | 6-20 mg/dl | Calculated |
| CREATININE, Serum | 0.89 | 0.67-1.17 mg/dl | Enzymatic |
| eGFR, Serum | 107 | >60 ml/min/1.73sqm | Calculated |
| URIC ACID, Serum | 4.2 | 3.5-7.2 mg/dl | Enzymatic |
| Urine Sugar (Fasting) | Absent | Absent | |
| Urine Ketones (Fasting) | Absent | Absent | |

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*** End Of Report ***



Anupa

Dr. ANUPA DIXIT
M.D.(PATH)
Consultant Pathologist & Lab Director

ADDRESS: 2nd Floor, Aston, Sundervan Complex, Above Mercedes Showroom, Andheri West - 400053

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ARCOFEMI HEALTHCARE- BLOOD TEST
GLYCOSYLATED HEMOGLOBIN (HbA1c)

| PARAMETER | RESULTS | BIOLOGICAL REF RANGE | METHOD |
|---|---------|---|------------|
| Glycosylated Hemoglobin (HbA1c), EDTA WB - CC | 5.2 | Non-Diabetic Level: < 5.7 % Prediabetic Level: 5.7-6.4 % Diabetic Level: >= 6.5 % | HPLC |
| Estimated Average Glucose (eAG), EDTA WB - CC | 102.5 | mg/dl | Calculated |

Intended use:

- In patients who are meeting treatment goals, HbA1c test should be performed at least 2 times a year
- In patients whose therapy has changed or who are not meeting glycemic goals, it should be performed quarterly
- For microvascular disease prevention, the HbA1C goal for non pregnant adults in general is Less than 7%.

Clinical Significance:

- HbA1c, Glycosylated hemoglobin or glycated hemoglobin, is hemoglobin with glucose molecule attached to it.
- The HbA1c test evaluates the average amount of glucose in the blood over the last 2 to 3 months by measuring the percentage of glycosylated hemoglobin in the blood.

Test Interpretation:

- The HbA1c test evaluates the average amount of glucose in the blood over the last 2 to 3 months by measuring the percentage of Glycosylated hemoglobin in the blood.
- HbA1c test may be used to screen for and diagnose diabetes or risk of developing diabetes.
- To monitor compliance and long term blood glucose level control in patients with diabetes.
- Index of diabetic control, predicting development and progression of diabetic micro vascular complications.

Factors affecting HbA1c results:

Increased in: High fetal hemoglobin, Chronic renal failure, Iron deficiency anemia, Splenectomy, Increased serum triglycerides, Alcohol ingestion, Lead/opiate poisoning and Salicylate treatment.

Decreased in: Shortened RBC lifespan (Hemolytic anemia, blood loss), following transfusions, pregnancy, ingestion of large amount of Vitamin E or Vitamin C and Hemoglobinopathies

Reflex tests: Blood glucose levels, CGM (Continuous Glucose monitoring)

References: ADA recommendations, AACC, Wallach's interpretation of diagnostic tests 10th edition.

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*** End Of Report ***



MC-2111

Dr. Leena Salunkhe

DR.LEENA SALUNKHE
M.B.B.S, DPB (PATH)
Pathologist

ADDRESS: 2nd Floor, Aston, Sundervan Complex, Above Mercedes Showroom, Andheri West - 400053

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Reported : 26-Nov-2021 / 19:02

**ARCOFEMI HEALTHCARE- BLOOD TEST
URINE EXAMINATION REPORT**

| <u>PARAMETER</u> | <u>RESULTS</u> | <u>BIOLOGICAL REF RANGE</u> | <u>METHOD</u> |
|---------------------------------------|----------------|-----------------------------|--------------------|
| <u>PHYSICAL EXAMINATION</u> | | | |
| Color | Yellow | Pale Yellow | - |
| Reaction (pH) | 5.0 | 4.5 - 8.0 | Chemical Indicator |
| Specific Gravity | 1.020 | 1.001-1.030 | Chemical Indicator |
| Transparency | Slight hazy | Clear | - |
| Volume (ml) | 30 | - | - |
| <u>CHEMICAL EXAMINATION</u> | | | |
| Proteins | Absent | Absent | pH Indicator |
| Glucose | Absent | Absent | GOD-POD |
| Ketones | Absent | Absent | Legals Test |
| Blood | Trace | Absent | Peroxidase |
| Bilirubin | Absent | Absent | Diazonium Salt |
| Urobilinogen | Normal | Normal | Diazonium Salt |
| Nitrite | Absent | Absent | Griess Test |
| <u>MICROSCOPIC EXAMINATION</u> | | | |
| Leukocytes(Pus cells)/hpf | 1-2 | 0-5/hpf | |
| Red Blood Cells / hpf | Occasional | 0-2/hpf | |
| Epithelial Cells / hpf | 0-1 | | |
| Casts | Absent | Absent | |
| Crystals | Absent | Absent | |
| Amorphous debris | Absent | Absent | |
| Bacteria / hpf | +(>20/hpf) | Less than 20/hpf | |
| Others | - | | |

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M Jain
Dr.MILLU JAIN
M.D.(PATH)
Pathologist

ADDRESS: 2nd Floor, Aston, Sundervan Complex, Above Mercedes Showroom, Andheri West - 400053

HEALTHLINE - MUMBAI: 022-6170-0000 | OTHER CITIES: 1800-266-4343

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ARCOFEMI HEALTHCARE- BLOOD TEST
BLOOD GROUPING & Rh TYPING

| PARAMETER | RESULTS |
|-----------|----------|
| ABO GROUP | B |
| Rh TYPING | POSITIVE |

NOTE: Test performed by automated column agglutination technology (CAT) which is more sensitive than conventional methods.

Specimen: EDTA Whole Blood and/or serum

Clinical significance:

ABO system is most important of all blood group in transfusion medicine

Limitations:

- ABO blood group of new born is performed only by cell (forward) grouping because allo antibodies in cord blood are of maternal origin.
- Since A & B antigens are not fully developed at birth, both Anti-A & Anti-B antibodies appear after the first 4 to 6 months of life. As a result, weaker reactions may occur with red cells of newborns than of adults.
- Confirmation of newborn's blood group is indicated when A & B antigen expression and the isoagglutinins are fully developed at 2 to 4 years of age & remains constant throughout life.
- Cord blood is contaminated with Wharton's jelly that causes red cell aggregation leading to false positive result
- The Hh blood group also known as Oh or Bombay blood group is rare blood group type. The term Bombay is used to refer the phenotype that lacks normal expression of ABH antigens because of inheritance of hh genotype.

References:

1. Denise M Harmening, Modern Blood Banking and Transfusion Practices- 6th Edition 2012. F.A. Davis company. Philadelphia
2. AABB technical manual

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*** End Of Report ***



MC-2111

Dr. Trupti Shetty
Dr. TRUPTI SHETTY
M. D. (PATH)
Pathologist

ADDRESS: 2nd Floor, Aston, Sundervan Complex, Above Mercedes Showroom, Andheri West - 400053

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ARCOFEMI HEALTHCARE- BLOOD TEST
LIPID PROFILE

| PARAMETER | RESULTS | BIOLOGICAL REF RANGE | METHOD |
|----------------------------------|---------|---|------------|
| CHOLESTEROL, Serum | 178.5 | Desirable: <200 mg/dl Borderline High: 200-239mg/dl High: >/=240 mg/dl | Enzymatic |
| TRIGLYCERIDES, Serum | 189.2 | Normal: <150 mg/dl Borderline-high: 150 - 199 mg/dl High: 200 - 499 mg/dl Very high:>/=500 mg/dl | Enzymatic |
| HDL CHOLESTEROL, Serum | 31.5 | Desirable: >60 mg/dl Borderline: 40 - 60 mg/dl Low (High risk): <40 mg/dl | Enzymatic |
| NON HDL CHOLESTEROL, Serum | 147 | Desirable: <130 mg/dl Borderline-high:130 - 159 mg/dl High:160 - 189 mg/dl Very high: >/=190 mg/dl | Calculated |
| LDL CHOLESTEROL, Serum | 109.0 | Optimal: <100 mg/dl Near Optimal: 100 - 129 mg/dl Borderline High: 130 - 159 mg/dl High: 160 - 189 mg/dl Very High: >/= 190 mg/dl | Calculated |
| VLDL CHOLESTEROL, Serum | 38.0 | < /= 30 mg/dl | Calculated |
| CHOL / HDL CHOL RATIO, Serum | 5.7 | 0-4.5 Ratio | Calculated |
| LDL CHOL / HDL CHOL RATIO, Serum | 3.5 | 0-3.5 Ratio | Calculated |

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ARCOFEMI HEALTHCARE- BLOOD TEST
THYROID FUNCTION TESTS

| PARAMETER | RESULTS | BIOLOGICAL REF RANGE | METHOD |
|---------------------|---------|----------------------|--------|
| Free T3, Serum | 5.3 | 3.5-6.5 pmol/L | ECLIA |
| Free T4, Serum | 17.7 | 11.5-22.7 pmol/L | ECLIA |
| sensitiveTSH, Serum | 1.23 | 0.35-5.5 microIU/ml | ECLIA |

Interpretation:

A thyroid panel is used to evaluate thyroid function and/or help diagnose various thyroid disorders.

Clinical Significance:

- 1)TSH Values between 5.5 to 15 microIU/ml should be correlated clinically or repeat the test with new sample as physiological factors can give falsely high TSH.
- 2)TSH values may be transiently altered because of non thyroidal illness like severe infections,liver disease, renal and heart severe burns, trauma and surgery etc.

| TSH | FT4 / T4 | FT3 / T3 | Interpretation |
|------|----------|----------|---|
| High | Normal | Normal | Subclinical hypothyroidism, poor compliance with thyroxine, drugs like amiodarone, Recovery phase of non-thyroidal illness, TSH Resistance. |
| High | Low | Low | Hypothyroidism, Autoimmune thyroiditis, post radio iodine Rx, post thyroidectomy, Anti thyroid drugs, tyrosine kinase inhibitors & amiodarone, amyloid deposits in thyroid, thyroid tumors & congenital hypothyroidism. |
| Low | High | High | Hyperthyroidism, Graves disease, toxic multinodular goiter, toxic adenoma, excess iodine or thyroxine intake, pregnancy related (hyperemesis gravidarum, hydatiform mole) |
| Low | Normal | Normal | Subclinical Hyperthyroidism, recent Rx for Hyperthyroidism, drugs like steroids & dopamine), Non thyroidal illness. |
| Low | Low | Low | Central Hypothyroidism, Non Thyroidal Illness, Recent Rx for Hyperthyroidism. |
| High | High | High | Interfering anti TPO antibodies, Drug interference: Amiodarone, Heparin, Beta Blockers, steroids & anti epileptics. |

Diurnal Variation:TSH follows a diurnal rhythm and is at maximum between 2 am and 4 am , and is at a minimum between 6 pm and 10 pm. The variation is on the order of 50 to 206%. Biological variation:19.7%(with in subject variation)

Reflex Tests:Anti thyroid Antibodies,USG Thyroid ,TSH receptor Antibody. Thyroglobulin, Calcitonin

Limitations:Samples should not be taken from patients receiving therapy with high biotin doses (i.e. >5 mg/day) until atleast 8 hours following the last biotin administration.

Reference:

- 1.O.koulouri et al. / Best Practice and Research clinical Endocrinology and Metabolism 27(2013)
- 2.Interpretation of the thyroid function tests, Dayan et al. THE LANCET . Vol 357
- 3.Tietz ,Text Book of Clinical Chemistry and Molecular Biology -5th Edition
- 4.Biological Variation:From principles to Practice-Callum G Fraser (AACC Press)

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M.D.(PATH)
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ARCOFEMI HEALTHCARE- BLOOD TEST
LIVER FUNCTION TESTS

| <u>PARAMETER</u> | <u>RESULTS</u> | <u>BIOLOGICAL REF RANGE</u> | <u>METHOD</u> |
|-----------------------------|----------------|-----------------------------|------------------|
| BILIRUBIN (TOTAL), Serum | 0.46 | 0.1-1.2 mg/dl | Colorimetric |
| BILIRUBIN (DIRECT), Serum | 0.17 | 0-0.3 mg/dl | Diazo |
| BILIRUBIN (INDIRECT), Serum | 0.29 | 0.1-1.0 mg/dl | Calculated |
| TOTAL PROTEINS, Serum | 8.4 | 6.4-8.3 g/dL | Biuret |
| ALBUMIN, Serum | 4.8 | 3.5-5.2 g/dL | BCG |
| GLOBULIN, Serum | 3.6 | 2.3-3.5 g/dL | Calculated |
| A/G RATIO, Serum | 1.3 | 1 - 2 | Calculated |
| SGOT (AST), Serum | 45.9 | 5-40 U/L | NADH (w/o P-5-P) |
| SGPT (ALT), Serum | 63.8 | 5-45 U/L | NADH (w/o P-5-P) |
| GAMMA GT, Serum | 22.2 | 3-60 U/L | Enzymatic |
| ALKALINE PHOSPHATASE, Serum | 126.5 | 40-130 U/L | Colorimetric |

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