

CID : 2317521933 Name : MRS.NETRA BENGALI Age / Gender : 34 Years / Female Consulting Dr. : -Reg. Location : Malad West (Main Centre) Authenticity Check

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Collected Reported :24-Jun-2023 / 09:55 :24-Jun-2023 / 13:49

AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE

	<u>CBC (Complet</u>	<u>e Blood Count), Blood</u>	
<u>PARAMETER</u>	<u>RESULTS</u>	BIOLOGICAL REF RANGE	<u>METHOD</u>
RBC PARAMETERS			
Haemoglobin	12.2	12.0-15.0 g/dL	Spectrophotometric
RBC	4.38	3.8-4.8 mil/cmm	Elect. Impedance
PCV	37.9	36-46 %	Calculated
MCV	86.6	80-100 fl	Measured
MCH	27.9	27-32 pg	Calculated
MCHC	32.2	31.5-34.5 g/dL	Calculated
RDW	16.4	11.6-14.0 %	Calculated
WBC PARAMETERS			
WBC Total Count	6870	4000-10000 /cmm	Elect. Impedance
WBC DIFFERENTIAL AND	ABSOLUTE COUNTS		
Lymphocytes	15.1	20-40 %	
Absolute Lymphocytes	1037.4	1000-3000 /cmm	Calculated
Monocytes	6.3	2-10 %	
Absolute Monocytes	432.8	200-1000 /cmm	Calculated
Neutrophils	72.7	40-80 %	
Absolute Neutrophils	4994.5	2000-7000 /cmm	Calculated
Eosinophils	5.3	1-6 %	
Absolute Eosinophils	364.1	20-500 /cmm	Calculated
Basophils	0.6	0.1-2 %	
Absolute Basophils	41.2	20-100 /cmm	Calculated
Immature Leukocytes	-		

WBC Differential Count by Absorbance & Impedance method/Microscopy.

PLATELET PARAMETERS Platelet Count 317000 150000-400000 /cmm Elect. Impedance MPV 6-11 fl Measured 8.6 PDW 15.1 11-18 % Calculated **RBC MORPHOLOGY** Hypochromia Microcytosis

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HEALTHLINE: 022-6170-0000 | E-MAIL: customerservice@suburbandiagnostics.com | WEBSITE: www.suburbandiagnostics.com

Corporate Identity Number (CIN): U85110MH2002PTC136144



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CID Name	: 2317521933 : MRS.NETRA				O R
Age / Gender Consulting Dr. Reg. Location	: 34 Years / : - : Malad West	Female : (Main Centre)	Collected Reported	Use a QR Code Scanner Application To Scan the Code : 24-Jun-2023 / 09:55 : 24-Jun-2023 / 12:40	т
Macrocytosis		-			
Anisocytosis		Mild			
Poikilocytosis		Mild			
Polychromasia		-			
Target Cells		-			
Basophilic Stipp	oling	-			
Normoblasts		-			
Others		Elliptocytes-occasional			
WBC MORPHO	DLOGY	-			
PLATELET MC	RPHOLOGY	-			
COMMENT		-			
Specimen: EDTA V	Vhole Blood				
ESR, EDTA WE	3-ESR	15	2-20 mm at 1 hr.	Sedimentation	

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



C. Solunda \mathcal{Y}

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

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:24-Jun-2023 / 09:55 :24-Jun-2023 / 12:45

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:2317521933

AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE				
PARAMETER	<u>RESULTS</u>	BIOLOGICAL REF RANGE	<u>METHOD</u>	
GLUCOSE (SUGAR) FASTING, Fluoride Plasma	81.5	Non-Diabetic: < 100 mg/dl Impaired Fasting Glucose: 100-125 mg/dl Diabetic: >/= 126 mg/dl	Hexokinase	
GLUCOSE (SUGAR) PP, Fluoride Plasma PP/R	104.0	Non-Diabetic: < 140 mg/dl Impaired Glucose Tolerance: 140-199 mg/dl Diabetic: >/= 200 mg/dl	Hexokinase	
BILIRUBIN (TOTAL), Serum	0.47	0.1-1.2 mg/dl	Colorimetric	
BILIRUBIN (DIRECT), Serum	0.21	0-0.3 mg/dl	Diazo	
BILIRUBIN (INDIRECT), Serum	0.26	0.1-1.0 mg/dl	Calculated	
TOTAL PROTEINS, Serum	7.2	6.4-8.3 g/dL	Biuret	
ALBUMIN, Serum	4.7	3.5-5.2 g/dL	BCG	
GLOBULIN, Serum	2.5	2.3-3.5 g/dL	Calculated	
A/G RATIO, Serum	1.9	1 - 2	Calculated	
SGOT (AST), Serum	10.0	5-32 U/L	NADH (w/o P-5-P)	
SGPT (ALT), Serum	13.8	5-33 U/L	NADH (w/o P-5-P)	
GAMMA GT, Serum	16.1	3-40 U/L	Enzymatic	
ALKALINE PHOSPHATASE, Serum	136.5	35-105 U/L	Colorimetric	
BLOOD UREA, Serum	11.2	12.8-42.8 mg/dl	Kinetic	
BUN, Serum	5.2	6-20 mg/dl	Calculated	
CREATININE, Serum eGFR, Serum	0.70 102	0.51-0.95 mg/dl >60 ml/min/1.73sqm	Enzymatic Calculated	

Note: eGFR estimation is calculated using MDRD (Modification of diet in renal disease study group) equation

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URIC ACID, Se	rum 4.8	2.4-5.7 mg/dl	Enzymatic	
Consulting Dr. Reg. Location	: - : Malad West (Main Centre)	Collected Reported	: 24-Jun-2023 / 09:55 :24-Jun-2023 / 19:14	
Age / Gender	: 34 Years / Female		Use a QR Code Scanner Application To Scan the Code	т
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PRECISE TESTING - HEAL	THIER LIVING			Р
DIAGNOSTI	c s			E

Urine Sugar (Fasting) Absent Absent Urine Ketones (Fasting) Absent Absent

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Anto

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

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Non-Diabetic Level: < 5.7 %

Prediabetic Level: 5.7-6.4 % Diabetic Level: >/= 6.5 %

: 24-Jun-2023 / 09:55 :24-Jun-2023 / 13:23

HPLC

Calculated

AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE **GLYCOSYLATED HEMOGLOBIN (HbA1c) BIOLOGICAL REF RANGE** RESULTS METHOD

mg/dl

PARAMETER

Glycosylated Hemoglobin

(HbA1c), EDTA WB - CC

5.6

Estimated Average Glucose 114.0 (eAG), EDTA WB - CC

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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Former

Dr.NAMRATA RAUL M.D (Biochem) **Biochemist**

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AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE URINE EXAMINATION REPORT

<u>PARAMETER</u>	<u>RESULTS</u>	BIOLOGICAL REF RANGE	<u>METHOD</u>
PHYSICAL EXAMINATION			
Color	Pale yellow	Pale Yellow	-
Reaction (pH)	8.0	4.5 - 8.0	Chemical Indicator
Specific Gravity	1.015	1.001-1.030	Chemical Indicator
Transparency	Slight hazy	Clear	-
Volume (ml)	50	-	-
CHEMICAL EXAMINATION			
Proteins	Absent	Absent	pH Indicator
Glucose	Absent	Absent	GOD-POD
Ketones	Absent	Absent	Legals Test
Blood	Absent	Absent	Peroxidase
Bilirubin	Absent	Absent	Diazonium Salt
Urobilinogen	Normal	Normal	Diazonium Salt
Nitrite	Absent	Absent	Griess Test
MICROSCOPIC EXAMINATION	<u>N</u>		
Leukocytes(Pus cells)/hpf	3-4	0-5/hpf	
Red Blood Cells / hpf	Absent	0-2/hpf	
Epithelial Cells / hpf	8-10		
Casts	Absent	Absent	
Crystals	Absent	Absent	
Amorphous debris	Absent	Absent	
Bacteria / hpf	+(>20/hpf)	Less than 20/hpf	
Others	-		

Kindly rule out contamination.

Interpretation: The concentration values of Chemical analytes corresponding to the grading given in the report are as follows:

- Protein:(1+ ~25 mg/dl, 2+ ~75 mg/dl, 3+ ~ 150 mg/dl, 4+ ~ 500 mg/dl)
- Glucose:(1+ ~ 50 mg/dl, 2+ ~100 mg/dl, 3+ ~300 mg/dl,4+ ~1000 mg/dl)

Ketone: (1+ ~5 mg/dl, 2+ ~15 mg/dl, 3+ ~ 50 mg/dl, 4+ ~ 150 mg/dl)

Reference: Pack insert

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M. Jain

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

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AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE BLOOD GROUPING & Rh TYPING

PARAMETER

RESULTS

ABO GROUP Rh TYPING

POSITIVE

0

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

Note: This sample is not tested for Bombay blood group.

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.

Refernces:

- 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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AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE LIPID PROFILE

PARAMETER	<u>RESULTS</u>	BIOLOGICAL REF RANGE	<u>METHOD</u>
CHOLESTEROL, Serum	194.7	Desirable: <200 mg/dl Borderline High: 200-239mg/dl High: >/=240 mg/dl	CHOD-POD
TRIGLYCERIDES, Serum	85.4	Normal: <150 mg/dl Borderline-high: 150 - 199 mg/dl High: 200 - 499 mg/dl Very high:>/=500 mg/dl	GPO-POD
HDL CHOLESTEROL, Serum	55.8	Desirable: >60 mg/dl Borderline: 40 - 60 mg/dl Low (High risk): <40 mg/dl	Homogeneous enzymatic colorimetric assay
NON HDL CHOLESTEROL, Serum	138.9	Desirable: <130 mg/dl Borderline-high:130 - 159 mg/dl High:160 - 189 mg/dl Very high: >/=190 mg/dl	Calculated
LDL CHOLESTEROL, Serum	122.0	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	16.9	< /= 30 mg/dl	Calculated
CHOL / HDL CHOL RATIO, Serum	3.5	0-4.5 Ratio	Calculated
LDL CHOL / HDL CHOL RATIO, Serum	2.2	0-3.5 Ratio	Calculated
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AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE **THYROID FUNCTION TESTS** RESULTS **BIOLOGICAL REF RANGE** PARAMETER METHOD Free T3, Serum 3.1 3.5-6.5 pmol/L **ECLIA** Free T4, Serum 13.2 11.5-22.7 pmol/L **ECLIA** First Trimester:9.0-24.7 Second Trimester: 6.4-20.59 Third Trimester: 6.4-20.59 sensitiveTSH, Serum 0.589 0.35-5.5 microIU/ml **ECLIA** First Trimester:0.1-2.5 Second Trimester: 0.2-3.0 Third Trimester: 0.3-3.0 mIU/ml

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Interpretation:

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

Clinical Significance:

1)TSH Values between high abnormal upto15 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 trasiently altered becuase 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:

1. 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.

2. Patient samples may contain heterophilic antibodies that could react in immunoassays to give falsely elevated or depressed results.

this assay is designed to minimize interference from heterophilic antibodies.

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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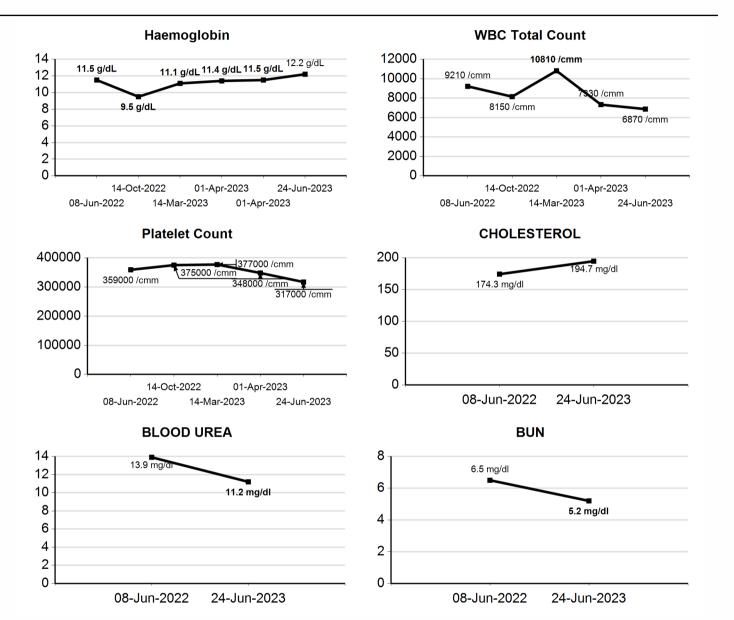
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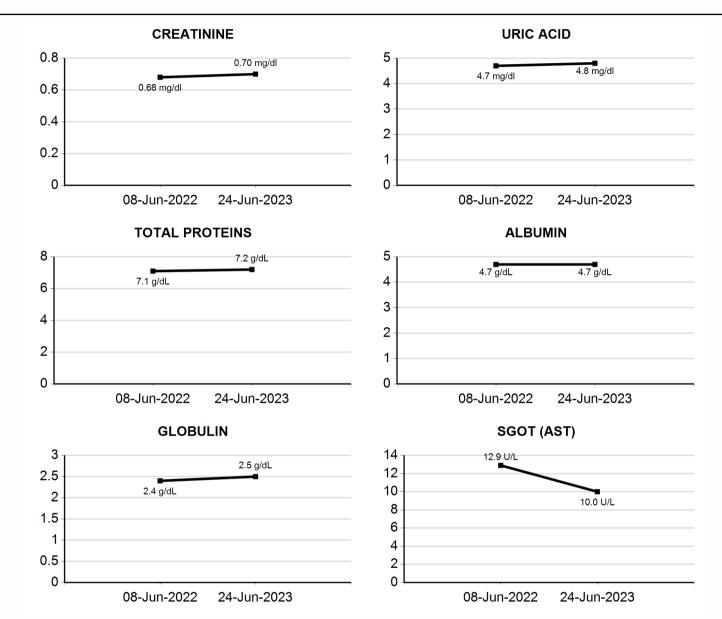
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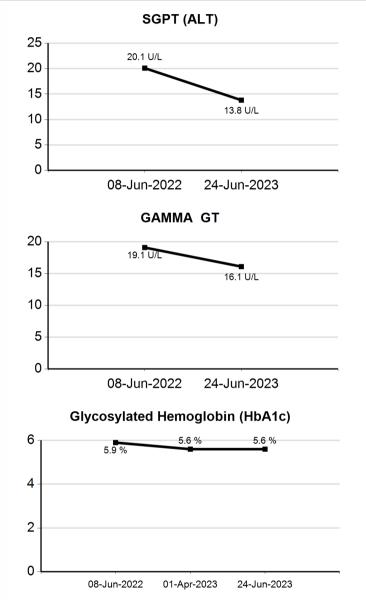
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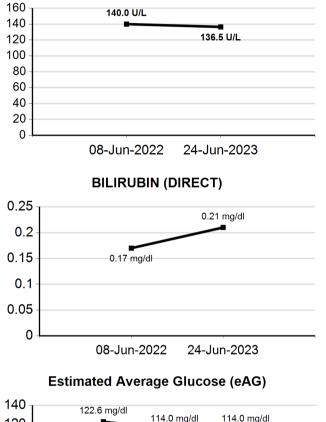
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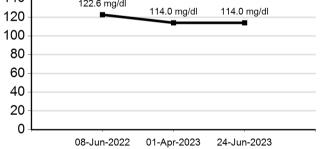
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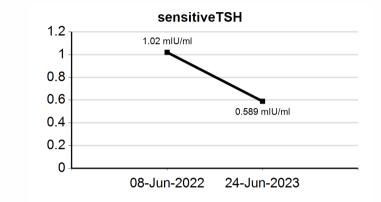


ALKALINE PHOSPHATASE





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