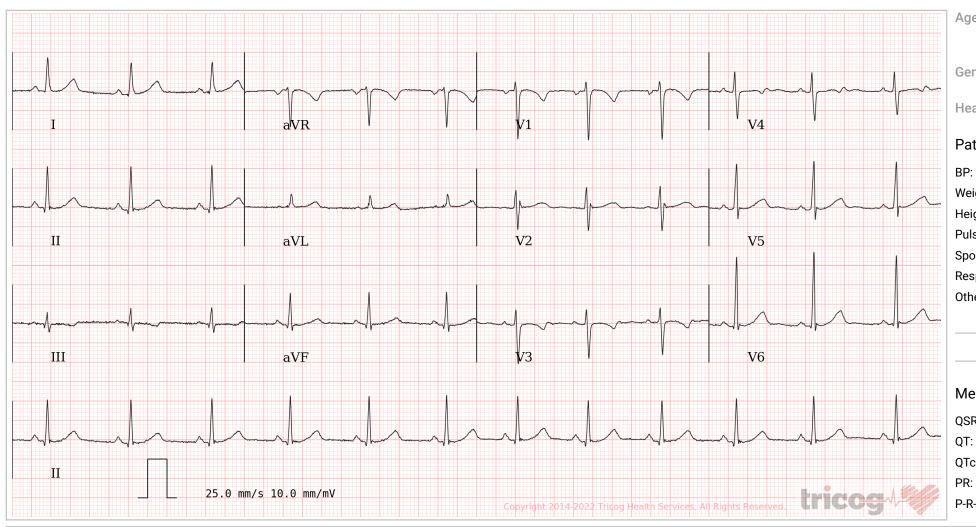
# SUBURBAN DIAGNOSTICS - ANDHERI WEST



Patient Name: SHWETA SINHA(ARCOFEMI)

Date and Time: 7th Feb 22 11:28 AM

Patient ID: 2203802121



Age years months days

Gender Female

Heart Rate 75 bpm

# **Patient Vitals**

110/70 mmHg BP:

NA

69 kg Weight:

Height: 167 cm

Pulse:

Spo2: NA

NA Resp:

Others:

Measurements

QSRD: 84 ms

406 ms

453 ms QTc:

PR: 128 ms

P-R-T: 45° 40° 32°

Sinus Rhythm, Normal Axis, rsr' in V2, Tinversion in leads V3-V4. Please correlate clinically.

REPORTED BY

DR RAVI CHAVAN MD, D.CARD, D. DIABETES Cardiologist & Diabetologist 2004/06/2468

Disclaimer: 1) Analysis in this report is based on ECG alone and should be used as an adjunct to clinical history, symptoms, and results of other invasive and non-invasive tests and must be interpreted by a qualified physician. 2) Patient vitals are as entered by the clinician and not derived from the ECG.



Name : MRS.SHWETA SINHA

Age / Gender : 35 Years / Female

Consulting Dr. : -

**Reg. Location**: Andheri West (Main Centre)



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# **AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE**

CBC (Complete Blood Count), Blood			
<u>PARAMETER</u>	<u>RESULTS</u>	<b>BIOLOGICAL REF RANGE</b>	<u>METHOD</u>
RBC PARAMETERS			
Haemoglobin	14.2	12.0-15.0 g/dL	Spectrophotometric
RBC	4.63	3.8-4.8 mil/cmm	Elect. Impedance
PCV	42.2	36-46 %	Measured
MCV	91.1	80-100 fl	Calculated
MCH	30.6	27-32 pg	Calculated
MCHC	33.6	31.5-34.5 g/dL	Calculated
RDW	15.1	11.6-14.0 %	Calculated
WBC PARAMETERS			
WBC Total Count	8640	4000-10000 /cmm	Elect. Impedance
WBC DIFFERENTIAL AND ABSO	LUTE COUNTS		
Lymphocytes	18.9	20-40 %	
Absolute Lymphocytes	1633.0	1000-3000 /cmm	Calculated
Monocytes	6.4	2-10 %	
Absolute Monocytes	553.0	200-1000 /cmm	Calculated
Neutrophils	73.0	40-80 %	
Absolute Neutrophils	6307.2	2000-7000 /cmm	Calculated
Eosinophils	1.5	1-6 %	
Absolute Eosinophils	129.6	20-500 /cmm	Calculated
Basophils	0.2	0.1-2 %	
Absolute Basophils	17.3	20-100 /cmm	Calculated

WBC Differential Count by Absorbance & Impedance method/Microscopy.

# **PLATELET PARAMETERS**

Platelet Count	102000	150000-400000 /cmm	Elect. Impedance
MPV	9.4	6-11 fl	Calculated
PDW	17.5	11-18 %	Calculated

**RBC MORPHOLOGY** 

Immature Leukocytes

Hypochromia Microcytosis -

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Macrocytosis -

Anisocytosis -

Poikilocytosis -

Polychromasia -

Target Cells -

Basophilic Stippling -

Normoblasts -

Others Normocytic, Normochromic

WBC MORPHOLOGY -

PLATELET MORPHOLOGY -

COMMENT -

Note: Collected sample received.

Result rechecked & Kindly correlate clinically

Advice: In view of low platelet count, repeat estimation with a fresh sample for confirmation, if clinically indicated before

taking clinical and therapeutic decision.

Specimen: EDTA Whole Blood

ESR, EDTA WB 38 2-20 mm at 1 hr. Westergren

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







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



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# AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE

<u>PARAMETER</u>	<u>RESULTS</u>	BIOLOGICAL REF RANGE	METHOD Hexokinase	
GLUCOSE (SUGAR) FASTING, Fluoride Plasma	77.9	Non-Diabetic: < 100 mg/dl Impaired Fasting Glucose: 100-125 mg/dl Diabetic: >/= 126 mg/dl		
GLUCOSE (SUGAR) PP, Fluoride Plasma PP/R	120.2	Non-Diabetic: < 140 mg/dl Impaired Glucose Tolerance: 140-199 mg/dl Diabetic: >/= 200 mg/dl	Hexokinase	
BILIRUBIN (TOTAL), Serum	0.33	0.1-1.2 mg/dl	Colorimetric	
BILIRUBIN (DIRECT), Serum	0.14	0-0.3 mg/dl	Diazo	
BILIRUBIN (INDIRECT), Serum	0.19	0.1-1.0 mg/dl	Calculated	
TOTAL PROTEINS, Serum	7.1	6.4-8.3 g/dL	Biuret	
ALBUMIN, Serum	3.9	3.5-5.2 g/dL	BCG	
GLOBULIN, Serum	3.2	2.3-3.5 g/dL	Calculated	
A/G RATIO, Serum	1.2	1 - 2	Calculated	
SGOT (AST), Serum	37.2	5-32 U/L	NADH (w/o P-5-P)	
SGPT (ALT), Serum	40.2	5-33 U/L	NADH (w/o P-5-P)	
GAMMA GT, Serum	15.9	3-40 U/L	Enzymatic	
ALKALINE PHOSPHATASE, Serum	167.0	35-105 U/L	Colorimetric	
BLOOD UREA, Serum	9.1	12.8-42.8 mg/dl	Kinetic	
BUN, Serum	4.2	6-20 mg/dl	Calculated	
CREATININE, Serum	0.48	0.51-0.95 mg/dl	Enzymatic	
eGFR, Serum	156	>60 ml/min/1.73sqm	Calculated	
URIC ACID, Serum	3.9	2.4-5.7 mg/dl	Enzymatic	

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:07-Feb-2022 / 18:44

Urine Sugar (Fasting)AbsentAbsentUrine Ketones (Fasting)AbsentAbsent

Urine Sugar (PP)AbsentAbsentUrine Ketones (PP)AbsentAbsent

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\*\*\* End Of Report \*\*\*







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

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:07-Feb-2022 / 15:46

# **AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE GLYCOSYLATED HEMOGLOBIN (HbA1c)**

### **BIOLOGICAL REF RANGE PARAMETER RESULTS** METHOD

Glycosylated Hemoglobin **HPLC** 4.8 Non-Diabetic Level: < 5.7 % (HbA1c), EDTA WB - CC

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

91.1 Estimated Average Glucose mg/dl Calculated

(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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Dr.MEGHA SHARMA M.D. (PATH), DNB (PATH) **Pathologist** 

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# **AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE** LIRINF FXAMINATION REPORT

ORINE EXAMINATION REPORT			
<u>PARAMETER</u>	<u>RESULTS</u>	<b>BIOLOGICAL REF RANGE</b>	<u>METHOD</u>
PHYSICAL EXAMINATION			
Color	Pale yellow	Pale Yellow	-
Reaction (pH)	7.0	4.5 - 8.0	Chemical Indicator
Specific Gravity	1.005	1.001-1.030	Chemical Indicator
Transparency	Clear	Clear	-
Volume (ml)	30	-	-
<b>CHEMICAL EXAMINATION</b>			
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	1-2	0-5/hpf	

Red Blood Cells / hpf Absent 0-2/hpf

Epithelial Cells / hpf 2-3

Casts Absent Absent Crystals **Absent Absent** Amorphous debris Absent Absent

Bacteria / hpf 3-4 Less than 20/hpf

Others







**Dr.SHASHIKANT DIGHADE** M.D. (PATH) **Pathologist** 

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

<u>PARAMETER</u> <u>RESULTS</u>

ABO GROUP 0

Rh TYPING POSITIVE

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

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# **AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE** LIPID PROFILE

<u>PARAMETER</u>	<u>RESULTS</u>	BIOLOGICAL REF RANGE	<u>METHOD</u>
CHOLESTEROL, Serum	184.9	Desirable: <200 mg/dl Borderline High: 200-239mg/dl High: >/=240 mg/dl	Enzymatic
TRIGLYCERIDES, Serum	182.4	Normal: <150 mg/dl Borderline-high: 150 - 199 mg/dl High: 200 - 499 mg/dl Very high:>/=500 mg/dl	Enzymatic
HDL CHOLESTEROL, Serum	72.2	Desirable: >60 mg/dl Borderline: 40 - 60 mg/dl Low (High risk): <40 mg/dl	Enzymatic
NON HDL CHOLESTEROL, Serum	112.7	Desirable: <130 mg/dl Borderline-high:130 - 159 mg/d High:160 - 189 mg/dl Very high: >/=190 mg/dl	Calculated l
LDL CHOLESTEROL, Serum	77.0	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	35.7	< /= 30 mg/dl	Calculated
CHOL / HDL CHOL RATIO, Serum	2.6	0-4.5 Ratio	Calculated
LDL CHOL / HDL CHOL RATIO, Serum	1.1	0-3.5 Ratio	Calculated

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# AERFOCAMI HEALTHCARE BELOW 40 MALE/FEMALE THYROID FUNCTION TESTS

<u>PARAMETER</u>	<u>RESULTS</u>	<b>BIOLOGICAL REF RANGE</b>	<u>METHOD</u>
Free T3, Serum	4.6	3.5-6.5 pmol/L	ECLIA
Free T4, Serum	9.3	11.5-22.7 pmol/L First Trimester:9.0-24.7 Second Trimester:6.4-20.59 Third Trimester:6.4-20.59	ECLIA
sensitiveTSH, Serum	2.42	0.35-5.5 microIU/ml First Trimester:0.1-2.5 Second Trimester:0.2-3.0 Third Trimester:0.3-3.0	ECLIA



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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 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: 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)







**Dr.VRUSHALI SHROFF** M.D.(PATH) **Pathologist** 

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