

PATIENT NAME : SONAL BHATT

REF. DOCTOR : SELF

CODE/NAME & ADDRESS : C000138362
 ACROFEMI HEALTHCARE LTD (MEDIWHEEL)
 F-703, LADO SARAI, MEHRAULISOUTH WEST
 DELHI
 NEW DELHI 110030
 8800465156

ACCESSION NO : 0030WB005257
 PATIENT ID : SONAF03106830
 CLIENT PATIENT ID:
 ABHA NO :

AGE/SEX : 54 Years Female
 DRAWN :
 RECEIVED : 25/02/2023 08:23:17
 REPORTED : 27/02/2023 15:46:59

Test Report Status	Preliminary	Results	Biological Reference Interval	Units
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MEDI WHEEL FULL BODY HEALTH CHECKUP ABOVE 40FEMALE**XRAY-CHEST****IMPRESSION**

NO ABNORMALITY DETECTED

TMT OR ECHO**TMT OR ECHO**

2D-ECHO DONE

2D ECHOCARDIOGRAPHY & COLOR DOPPLER STUDY

INDICATION - CARDIAC EVALUATION

HR - 80/min, sinus

CARDIAC CHAMBER DIMENSION AND FUNCTION

LA: Normal

LV: Normal, No wall motion abnormality

LV systolic function - Normal, LVEF - 60%

LV diastolic function - Grade I diastolic dysfunction

RA: Normal, RV: Normal

CARDIAC VALVES -

Mitral valve - Normal, No mitral regurgitation.

Aortic valve - Three leaflets, No aortic regurgitation

Tricuspid valve - Trivial tricuspid regurgitation, No PAH

Pulmonary valve - Normal

Septae (IAS/IVS) - Intact on trans-thoracic echo

Clot/Vegetation/Pericardial effusion - No

Great Arteries (Aorta/pulmonary artery) - Normal

IVC - Normal calibre and collapsibility

MEASUREMENTS -

AO	LA	IVS	PW	LVIDd	LVIDs
21	24	9	9	40	22

CONCLUSION:-

NORMAL CHAMBER DIMENSIONS

NO RWMA, NORMAL LV SYSTOLIC FUNCTION, LVEF - 60%

GRADE I LV DIASTOLIC DYSFUNCTION

NORMAL PA PRESSURE

DR JIGNESH PARIKH

DNB (MED), DNB (CARD)

CARDIOLOGIST

ECG**ECG**

WITHIN NORMAL LIMITS

MAMOGRAPHY (BOTH BREASTS)


Dr. Swati Pravin Mulani
 Lab Head

Page 1 Of 20



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Patient Ref. No. 775000002438805

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MAMOGRAPHY BOTH BREASTS

RIGHT BREAST - Shows mixed fatty and glandular parenchyma.
 No focal lesion is seen.

LEFT BREAST - Shows mixed fatty and glandular parenchyma.
 No focal lesion is seen.

No obvious axillary adenopathy noted on either side.

Clinical correlation.

SoS X-Mammography if clinically indicated.

MEDICAL HISTORY

RELEVANT PRESENT HISTORY
 RELEVANT PAST HISTORY
 RELEVANT PERSONAL HISTORY
 MENSTRUAL HISTORY (FOR FEMALES)
 LMP (FOR FEMALES)
 RELEVANT FAMILY HISTORY
 OCCUPATIONAL HISTORY
 HISTORY OF MEDICATIONS

K/C/O HYPERTENSION, UNDER TREATMENT
 ANGIOPLASTY IN 2019.
 NOT SIGNIFICANT
 REGULAR.
 LMP - 19-01-2023
 HIGH BLOOD PRESSURE, HEART DISEASE
 NOT SIGNIFICANT
 CAP. ROSUMAC GOLD, TAB. TELMA-CT

ANTHROPOMETRIC DATA & BMI

HEIGHT IN METERS
 WEIGHT IN KGS.
 BMI

1.57
 84
 34

mts
 Kgs

BMI & Weight Status as follows:
 Below 18.5: Underweight
 18.5 - 24.9: Normal
 25.0 - 29.9: Overweight
 30.0 and Above: Obese

GENERAL EXAMINATION

MENTAL / EMOTIONAL STATE
 PHYSICAL ATTITUDE
 GENERAL APPEARANCE / NUTRITIONAL
 STATUS
 BUILT / SKELETAL FRAMEWORK
 FACIAL APPEARANCE
 SKIN

NORMAL
 NORMAL
 OVERWEIGHT
 AVERAGE
 NORMAL
 NORMAL



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Page 2 Of 20



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UPPER LIMB	NORMAL			
LOWER LIMB	NORMAL			
NECK	NORMAL			
NECK LYMPHATICS / SALIVARY GLANDS	NOT ENLARGED OR TENDER			
THYROID GLAND	NOT ENLARGED			
CAROTID PULSATION	NORMAL			
TEMPERATURE	NORMAL			
PULSE	82/MIN REGULAR, ALL PERIPHERAL PULSES WELL FELT, NO CAROTID BRUIT			
RESPIRATORY RATE	NORMAL			
CARDIOVASCULAR SYSTEM				
BP	112/70 MM HG (SITTING)		mm/Hg	
PERICARDIUM	NORMAL			
APEX BEAT	NORMAL			
HEART SOUNDS	NORMAL			
MURMURS	ABSENT			
RESPIRATORY SYSTEM				
SIZE AND SHAPE OF CHEST	NORMAL			
MOVEMENTS OF CHEST	SYMMETRICAL			
BREATH SOUNDS INTENSITY	NORMAL			
BREATH SOUNDS QUALITY	VESICULAR (NORMAL)			
ADDED SOUNDS	ABSENT			
PER ABDOMEN				
APPEARANCE	NORMAL			
VENOUS PROMINENCE	ABSENT			
LIVER	NOT PALPABLE			
SPLEEN	NOT PALPABLE			
HERNIA	ABSENT			
CENTRAL NERVOUS SYSTEM				
HIGHER FUNCTIONS	NORMAL			
CRANIAL NERVES	NORMAL			
CEREBELLAR FUNCTIONS	NORMAL			



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Page 3 Of 20



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SENSORY SYSTEM	NORMAL
MOTOR SYSTEM	NORMAL
REFLEXES	NORMAL

MUSCULOSKELETAL SYSTEM

SPINE	NORMAL
JOINTS	NORMAL

BASIC EYE EXAMINATION

CONJUNCTIVA	NORMAL
EYELIDS	NORMAL
EYE MOVEMENTS	NORMAL
CORNEA	NORMAL
DISTANT VISION RIGHT EYE WITHOUT GLASSES	DISTANT VISION - 6/9
DISTANT VISION LEFT EYE WITHOUT GLASSES	DISTANT VISION - 6/9
NEAR VISION RIGHT EYE WITHOUT GLASSES	NEAR VISION - N 10
NEAR VISION LEFT EYE WITHOUT GLASSES	NEAR VISION - N 10
COLOUR VISION	NORMAL

BASIC ENT EXAMINATION

EXTERNAL EAR CANAL	NORMAL
TYMPANIC MEMBRANE	NORMAL
NOSE	NO ABNORMALITY DETECTED
SINUSES	NORMAL
THROAT	NO ABNORMALITY DETECTED
TONSILS	NOT ENLARGED

SUMMARY

RELEVANT HISTORY	K/C/O HYPERTENSION, UNDER TREATMENT
RELEVANT GP EXAMINATION FINDINGS	NOT SIGNIFICANT
RELEVANT LAB INVESTIGATIONS	HBA1C RAISED (6.1%) POST PRANDIAL BLOOD SUGAR LEVEL RAISED - 171 MG/DL TRIGLYCERIDE RAISED (182 mg/dL) HDL CHOLESTEROL LOW (32 mg/dL) DIRECT BILLIRUBIN RAISED - 0.35 MG/DL BLOOD DETECTED (+) IN URINE RBC'S 2-3 / HPF IN URINE



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Page 4 Of 20



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RELEVANT NON PATHOLOGY DIAGNOSTICS NO ABNORMALITIES DETECTED
 REMARKS / RECOMMENDATIONS ADV. REDUCE PROCESSED FOOD IN DIET
 INCREASE UNSATURATED FATS IN DIET
 REDUCE FRIED & OILY FOOD IN DIET
 REPEAT BILIRUBIN AFTER 15 DAYS
 DIABETIC DIET, REGULLAR EXRCISE.
 REDUCE INTAKE OF SWEETS, SUGAR & STARCH IN DIET.
 DO FASTING & POST PRANDIAL BLOOD SUGAR LEVEL AFTER 1 MONTH
 FOLLOW UP WITH DIABETOLOGIST.
 FOLLOW UP WITH EYE SPECIALIST
 FOLLOW UP WITH GASTROENTEROLOGIST.

FITNESS STATUS
 FITNESS STATUS FIT (WITH MEDICAL ADVICE) (AS PER REQUESTED PANEL OF TESTS)

Comments

 OUR DOCTORS ON PANEL FOR NON-PATHOLOGICAL REPORTS:
 1. DR. JIGNESH PARIKH: DNB (CARDIOLOGY), N.B.E
 (CONSULTANT CARDIOLOGIST)
 2. DR.SANJAY JOSHI, D M R D, DNB - RADIOLOGIST
 3. DR. SUCHARITA PARANJPE, MBBS, FCPS (OPHTHALMOLOGY)
 4. DR. (MRS.) MANJUSHA PRABHUNE - GYNAECOLOGIST.
 5. DR. (MRS.) NIMKAR - GYNAECOLOGIST.

This report bears the signature of the in-charge of the facility.
 Panel doctors are responsible for the results/reports of their individual specialty.



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MEDI WHEEL FULL BODY HEALTH CHECKUP ABOVE 40FEMALE**ULTRASOUND ABDOMEN****ULTRASOUND ABDOMEN****ULTRASONOGRAPHY OF ABDOMEN & PELVIS**

LIVER: Liver is normal in size. **Grade I / II changes of fatty liver are noted.** No focal intra-hepatic lesion is detected. Intrahepatic biliary radicals are not dilated. Portal vein is normal

GALL BLADDER: Gall bladder shows normal thickness of its walls. No calculi are seen. Common bile duct is normal .

PANCREAS: Pancreas is normal in size and echo pattern.

SPLEEN: Spleen is normal in size . It is normal in shape and position. Echoes are normal. Splenic vein is not dilated .

RIGHT KIDNEY: Normal in position, size and outline. Corticomedullary differentiation is maintained. Central sinus echoes are compact.No evidence of calculus is seen.No hydronephrosis.

LEFT KIDNEY: Normal in position, size and outline. Corticomedullary differentiation is maintained. Central sinus echoes are compact.No evidence of calculus is seen.No hydronephrosis.

URINARY BLADDER:Urinary bladder is normal in wall thickness with clear contents. Its walls show a smooth outline. There is no evidence of any intraluminal or perivesicle abnormality.

UTERUS: Uterus is normal in size & shape. Endometrium is central and normal in thickness. Myometrial echogenecity appears uniform. Cervix is normal.

OVARIES :Both ovaries are normal in size, shape and echo pattern.No abnormal adnexa mass lesion is seen.No free fluid is detected in pouch of Douglas and Morissons pouch.

No e/o any retroperitoneal lymphadenopathy.

No e/o any free fluid noted in abdomen.



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Page 6 Of 20



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Clinical correlation.
Sub optimal window due to obesity.

Interpretation(s)

MEDICAL HISTORY-*****
 THIS REPORT CARRIES THE SIGNATURE OF OUR LABORATORY DIRECTOR. THIS IS AN INVIOABLE FEATURE OF OUR LAB MANAGEMENT SOFTWARE. HOWEVER, ALL EXAMINATIONS AND INVESTIGATIONS HAVE BEEN CONDUCTED BY OUR PANEL OF DOCTORS.

 FITNESS STATUS-Conclusion on an individual's Fitness, which is commented upon mainly for Pre employment cases, is based on multi factorial findings and does not depend on any one single parameter. The final Fitness assigned to a candidate will depend on the Physician's findings and overall judgement on a case to case basis, details of the candidate's past and personal history; as well as the comprehensiveness of the diagnostic panel which has been requested for. These are then further correlated with details of the job under consideration to eventually fit the right man to the right job.

Basis the above, SRL classifies a candidate's Fitness Status into one of the following categories:

- Fit (As per requested panel of tests) - SRL Limited gives the individual a clean chit to join the organization, on the basis of the General Physical Examination and the specific test panel requestec for.
- Fit (with medical advice) (As per requested panel of tests) - This indicates that although the candidate can be declared as FIT to join the job, minimal problems have been detected during the Pre- employment examination. Examples of conditions which could fall in this category could be cases of mild reversible medical abnormalities such as height weight disproportions, borderline raised Blood Pressure readings, mildly raised Blood sugar anc Blood Lipid levels, Hematuria, etc. Most of these relate to sedentary lifestyles and come under the broad category of life style disorders. The idea is to caution an individual to bring about certain lifestyle changes as well as seek a Physician's consultation and counseling in order to bring back to normal the mildly deranged parameters. For all purposes the individual is FIT to join the job.
- Fitness on Hold (Temporary Unfit) (As per requested panel of tests) - Candidate's reports are kept on hold when either the diagnostic tests or the physical findings reveal the presence of a medical condition which warrants further tests, counseling and/or specialist opinion, on the basis of which a candidate can either be placed into Fit, Fit (With Medical Advice), or Unfit category. Conditions which may fall into this category could be high blood pressure, abnormal ECG, heart murmurs, abnormal vision, grossly elevated blood sugars, etc.
- Unfit (As per requested panel of tests) - An unfit report by SRL Limited clearly indicates that the individual is not suitable for the respective job profile e.g. total color blindness in color related jobs.



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HAEMATOLOGY - CBC

MEDI WHEEL FULL BODY HEALTH CHECKUP ABOVE 40FEMALE

BLOOD COUNTS, EDTA WHOLE BLOOD

HEMOGLOBIN (HB)	13.3	12.0 - 15.0	g/dL
RED BLOOD CELL (RBC) COUNT	4.86 High	3.8 - 4.8	mil/ μ L
METHOD : ELECTRICAL IMPEDANCE			
WHITE BLOOD CELL (WBC) COUNT	8.00	4.0 - 10.0	thou/ μ L
METHOD : ELECTRICAL IMPEDANCE			
PLATELET COUNT	366	150 - 410	thou/ μ L
METHOD : ELECTRICAL IMPEDANCE			

RBC AND PLATELET INDICES

HEMATOCRIT (PCV)	40.9	36 - 46	%
METHOD : CALCULATED			
MEAN CORPUSCULAR VOLUME (MCV)	84.0	83 - 101	fL
METHOD : CALCULATED			
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	27.5	27.0 - 32.0	pg
METHOD : CALCULATED			
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC)	32.6	31.5 - 34.5	g/dL
METHOD : CALCULATED			
RED CELL DISTRIBUTION WIDTH (RDW)	12.4	11.6 - 14.0	%
METHOD : CALCULATED			
MENTZER INDEX	17.3		
MEAN PLATELET VOLUME (MPV)	9.1	6.8 - 10.9	fL
METHOD : CELL COUNTER (CALCULATED)			

WBC DIFFERENTIAL COUNT

NEUTROPHILS	54	40 - 80	%
METHOD : ELECTRICAL IMPEDANCE/MICROSCOPY			
LYMPHOCYTES	34	20 - 40	%
METHOD : ELECTRICAL IMPEDANCE/MICROSCOPY			
MONOCYTES	8	2 - 10	%
EOSINOPHILS	4	1 - 6	%
METHOD : ELECTRICAL IMPEDANCE/MICROSCOPY			
BASOPHILS	0	0 - 2	%
METHOD : ELECTRICAL IMPEDANCE/MICROSCOPY			



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Page 8 Of 20



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ABSOLUTE NEUTROPHIL COUNT		4.32	2.0 - 7.0	thou/ μ L
METHOD : CALCULATED				
ABSOLUTE LYMPHOCYTE COUNT		2.72	1.0 - 3.0	thou/ μ L
METHOD : CALCULATED				
ABSOLUTE MONOCYTE COUNT		0.64	0.2 - 1.0	thou/ μ L
METHOD : CALCULATED				
ABSOLUTE EOSINOPHIL COUNT		0.32	0.02 - 0.50	thou/ μ L
METHOD : CALCULATED				
ABSOLUTE BASOPHIL COUNT		0.00 Low	0.02 - 0.10	thou/ μ L
METHOD : CALCULATED				
NEUTROPHIL LYMPHOCYTE RATIO (NLR)		1.6		
METHOD : CALCULATED				

MORPHOLOGY**REMARKS**

RBCS: PREDOMINANTLY NORMOCYTIC NORMOCHROMIC.
WBCS: WBCS ARE NORMAL IN NUMBER & MORPHOLOGY.
PLATELETS: ADEQUATE ON PERIPHERAL SMEAR.

Interpretation(s)

BLOOD COUNTS, EDTA WHOLE BLOOD- The cell morphology is well preserved for 24hrs. However after 24-48 hrs a progressive increase in MCV and HCT is observed leading to a decrease in MCHC. A direct smear is recommended for an accurate differential count and for examination of RBC morphology.
RBC AND PLATELET INDICES- Mentzer index (MCV/RBC) is an automated cell-counter based calculated screen tool to differentiate cases of Iron deficiency anaemia (>13) from Beta thalassaemia trait (<13) in patients with microcytic anaemia. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HbA2 remains the gold standard for diagnosing a case of beta thalassaemia trait.
WBC DIFFERENTIAL COUNT- The optimal threshold of 3.3 for NLR showed a prognostic possibility of clinical symptoms to change from mild to severe in COVID positive patients. When age = 49.5 years old and NLR = 3.3, 46.1% COVID-19 patients with mild disease might become severe. By contrast, when age < 49.5 years old and NLR < 3.3, COVID-19 patients tend to show mild disease.
(Reference to - The diagnostic and predictive role of NLR, d-NLR and PLR in COVID-19 patients ; A.-P. Yang, et al.; International Immunopharmacology 84 (2020) 106504
This ratio element is a calculated parameter and out of NABL scope.



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Page 9 Of 20



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HAEMATOLOGY

MEDI WHEEL FULL BODY HEALTH CHECKUP ABOVE 40FEMALE

ERYTHROCYTE SEDIMENTATION RATE (ESR), WHOLE BLOOD

E.S.R	15	0 - 20	mm at 1 hr
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METHOD : WESTERGREN METHOD

Interpretation(s)

ERYTHROCYTE SEDIMENTATION RATE (ESR), WHOLE BLOOD-TEST DESCRIPTION :-

Erythrocyte sedimentation rate (ESR) is a test that indirectly measures the degree of inflammation present in the body. The test actually measures the rate of fall (sedimentation) of erythrocytes in a sample of blood that has been placed into a tall, thin, vertical tube. Results are reported as the millimetres of clear fluid (plasma) that are present at the top portion of the tube after one hour. Nowadays fully automated instruments are available to measure ESR.

ESR is not diagnostic; it is a non-specific test that may be elevated in a number of different conditions. It provides general information about the presence of an inflammatory condition. CRP is superior to ESR because it is more sensitive and reflects a more rapid change.

TEST INTERPRETATION

Increase in: Infections, Vasculitides, Inflammatory arthritis, Renal disease, Anemia, Malignancies and plasma cell dyscrasias, Acute allergy Issue injury, Pregnancy, Estrogen medication, Aging.

Finding a very accelerated ESR (> 100 mm/hour) in patients with ill-defined symptoms directs the physician to search for a systemic disease (Paraproteinemias, Disseminated malignancies, connective tissue disease, severe infections such as bacterial endocarditis).

In pregnancy BRI in first trimester is 0-48 mm/hr (62 if anemic) and in second trimester (0-70 mm /hr (95 if anemic). ESR returns to normal 4th week post partum.

Decreased in: Polycythemia vera, Sickle cell anemia

LIMITATIONS

False elevated ESR : Increased fibrinogen, Drugs (Vitamin A, Dextran etc), Hypercholesterolemia

False Decreased : Poikilocytosis, (Sickle Cells, spherocytes), Microcytosis, Low fibrinogen, Very high WBC counts, Drugs (Quinine, salicylates)

REFERENCE :

1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition; 2. Paediatric reference intervals. AACC Press, 7th edition. Edited by S. Soldin; 3. The reference for the adult reference range is "Practical Haematology by Dacie and Lewis, 10th edition.

Page 10 Of 20



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Patient Ref. No. 775000002438805

PATIENT NAME : SONAL BHATT**REF. DOCTOR : SELF**

CODE/NAME & ADDRESS : C000138362

ACROFEMI HEALTHCARE LTD (MEDIWHEEL)

F-703, LADO SARAI, MEHRAULISOUTH WEST
DELHI

NEW DELHI 110030

8800465156

ACCESSION NO : **0030WB005257**

PATIENT ID : SONAF03106830

CLIENT PATIENT ID:

ABHA NO :

AGE/SEX : 54 Years Female

DRAWN :

RECEIVED : 25/02/2023 08:23:17

REPORTED : 27/02/2023 15:46:59

Test Report Status	<u>Preliminary</u>	Results	Biological Reference Interval	Units
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IMMUNOHAEMATOLOGY**MEDI WHEEL FULL BODY HEALTH CHECKUP ABOVE 40FEMALE****ABO GROUP & RH TYPE, EDTA WHOLE BLOOD**

ABO GROUP

TYPE B

METHOD : TUBE AGGLUTINATION

RH TYPE

POSITIVE

METHOD : TUBE AGGLUTINATION

Interpretation(s)

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD-

Blood group is identified by antigens and antibodies present in the blood. Antigens are protein molecules found on the surface of red blood cells. Antibodies are found in plasma. To determine blood group, red cells are mixed with different antibody solutions to give A,B,O or AB.

Disclaimer: "Please note, as the results of previous ABO and Rh group (Blood Group) for pregnant women are not available, please check with the patient records for availability of the same."

The test is performed by both forward as well as reverse grouping methods.


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		PATIENT ID : SONAF03106830	DRAWN :
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BIOCHEMISTRY

MEDI WHEEL FULL BODY HEALTH CHECKUP ABOVE 40FEMALE

GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD

HBA1C	6.1 High	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 Therapeutic goals: < 7.0 Action suggested : > 8.0 (ADA Guideline 2021)	%
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METHOD : HPLC

ESTIMATED AVERAGE GLUCOSE(EAG)	128.4 High	< 116.0	mg/dL
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GLUCOSE FASTING,FLUORIDE PLASMA FBS (FASTING BLOOD SUGAR)

FBS (FASTING BLOOD SUGAR)	98	74 - 99	mg/dL
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METHOD : HEXOKINASE

GLUCOSE, POST-PRANDIAL, PLASMA

PPBS(POST PRANDIAL BLOOD SUGAR)	171 High	Normal: < 140, Impaired Glucose Tolerance:140-199 Diabetic > or = 200	mg/dL
---------------------------------	-----------------	--	-------

METHOD : HEXOKINASE

LIPID PROFILE, SERUM

CHOLESTEROL, TOTAL	100	Desirable: <200 BorderlineHigh : 200-239 High : > or = 240	mg/dL
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TRIGLYCERIDES	182 High	Desirable: < 150 Borderline High: 150 - 199 High: 200 - 499 Very High : > or = 500	mg/dL
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METHOD : ENZYMATIC WITH GLYCEROL BLANK

HDL CHOLESTEROL	32 Low	< 40 Low > or = 60 High	mg/dL
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METHOD : DIRECT MEASURE - PEG



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CHOLESTEROL LDL		32	Adult levels: Optimal < 100 Near optimal/above optimal: 100-129 Borderline high : 130-159 High : 160-189 Very high : = 190	mg/dL
NON HDL CHOLESTEROL		68	Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL
VERY LOW DENSITY LIPOPROTEIN CHOL/HDL RATIO		36.4 3.1		mg/dL
LDL/HDL RATIO		1.0	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	

Interpretation(s)

LIVER FUNCTION PROFILE, SERUM

BILIRUBIN, TOTAL METHOD : DIAZONIUM ION, BLANKED (ROCHE)	0.91	0.0 - 1.2	mg/dL
BILIRUBIN, DIRECT METHOD : DIAZOTIZATION	0.35 High	0.0 - 0.2	mg/dL
BILIRUBIN, INDIRECT METHOD : CALCULATED PARAMETER	0.56	0.00 - 1.00	mg/dL
TOTAL PROTEIN METHOD : BIURET, REAGENT BLANK, END POINT	6.9	6.4 - 8.3	g/dL
ALBUMIN METHOD : BROMOCRESOL GREEN (BCG)	3.9	3.50 - 5.20	g/dL
GLOBULIN METHOD : CALCULATED PARAMETER	3.0	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO METHOD : CALCULATED PARAMETER	1.3	1.0 - 2.0	RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	17	UPTO 32	U/L

Page 13 Of 20


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ALANINE AMINOTRANSFERASE (ALT/SGPT)

16

UPTO 34

U/L

ALKALINE PHOSPHATASE

57

35 - 104

U/L

METHOD : PNPP - AMP BUFFER

GAMMA GLUTAMYL TRANSFERASE (GGT)

21

5 - 36

U/L

METHOD : GAMMA GLUTAMYL-3-CARBOXY-4-NITROANALIDE (IFCC)

LACTATE DEHYDROGENASE

160

135 - 214

U/L

METHOD : LACTATE -PYRUVATE

BLOOD UREA NITROGEN (BUN), SERUM

BLOOD UREA NITROGEN

6

6 - 20

mg/dL

METHOD : UREASE COLORIMETRIC

CREATININE, SERUM

CREATININE

0.62

0.50 - 0.90

mg/dL

METHOD : JAFFE'S ALKALINE PICRATE -IFCC IDMS STANDARDIZED

BUN/CREAT RATIO

BUN/CREAT RATIO

9.68

5.0 - 15.0

URIC ACID, SERUM

URIC ACID

4.8

2.6 - 6.0

mg/dL

METHOD : URICASE, COLORIMETRIC

TOTAL PROTEIN, SERUM

TOTAL PROTEIN

6.9

6.4 - 8.3

g/dL

METHOD : BIURET, REAGENT BLANK, END POINT

ALBUMIN, SERUM

ALBUMIN

3.9

3.5 - 5.2

g/dL

METHOD : BROMOCRESOL GREEN (BCG)

GLOBULIN

GLOBULIN

3.0

2.0 - 4.1

g/dL

METHOD : CALCULATED PARAMETER

ELECTROLYTES (NA/K/CL), SERUM

SODIUM, SERUM

136 Low

137 - 145

mmol/L

METHOD : ISE INDIRECT

POTASSIUM, SERUM

3.40 Low

3.6 - 5.0

mmol/L

METHOD : ISE INDIRECT

CHLORIDE, SERUM

102

98 - 107

mmol/L

METHOD : ISE INDIRECT

Interpretation(s)

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Interpretation(s)

GLYCOSYLATED HEMOGLOBIN(HBA1C), ED IA WHOLE BLOOD-Used For:

1. Evaluating the long-term control of blood glucose concentrations in diabetic patients.
 2. Diagnosing diabetes.
 3. Identifying patients at increased risk for diabetes (prediabetes).
- The ADA recommends measurement of HbA1c (typically 3-4 times per year for type 1 and poorly controlled type 2 diabetic patients, and 2 times per year for well-controlled type 2 diabetic patients) to determine whether a patient's metabolic control has remained continuously within the target range.
1. eAG (Estimated average glucose) converts percentage HbA1c to mg/dl, to compare blood glucose levels.
 2. eAG gives an evaluation of blood glucose levels for the last couple of months.
 3. eAG is calculated as eAG (mg/dl) = 28.7 * HbA1c - 46.7

HbA1c Estimation can get affected due to :

- I. Shortened Erythrocyte survival : Any condition that shortens erythrocyte survival or decreases mean erythrocyte age (e.g. recovery from acute blood loss, hemolytic anemia) will falsely lower HbA1c test results. Fructosamine is recommended in these patients which indicates diabetes control over 15 days.
- II. Vitamin C & E are reported to falsely lower test results. (possibly by inhibiting glycation of hemoglobin).
- III. Iron deficiency anemia is reported to increase test results. Hypertriglyceridemia, uremia, hyperbilirubinemia, chronic alcoholism, chronic ingestion of salicylates & opiates addition are reported to interfere with some assay methods, falsely increasing results.
- IV. Interference of hemoglobinopathies in HbA1c estimation is seen in
 - a. Homozygous hemoglobinopathy. Fructosamine is recommended for testing of HbA1c.
 - b. Heterozygous state detected (D10 is corrected for HbS & HbC trait.)
 - c. HbF > 25% on alternate platform (Boronate affinity chromatography) is recommended for testing of HbA1c. Abnormal Hemoglobin electrophoresis (HPLC method) is recommended for detecting a hemoglobinopathy

GLUCOSE FASTING, FLUORIDE PLASMA-TEST DESCRIPTION

Normally, the glucose concentration in extracellular fluid is closely regulated so that a source of energy is readily available to tissues and so that no glucose is excreted in the urine.

Increased in

Diabetes mellitus, Cushing's syndrome (10 - 15%), chronic pancreatitis (30%). Drugs: corticosteroids, phenytoin, estrogen, thiazides.

Decreased in

Pancreatic islet cell disease with increased insulin, insulinoma, adrenocortical insufficiency, hypopituitarism, diffuse liver disease, malignancy (adrenocortical, stomach, fibrosarcoma), infant of a diabetic mother, enzyme deficiency diseases (e.g., galactosemia), Drugs- insulin, ethanol, propranolol; sulfonylureas, tolbutamide, and other oral hypoglycemic agents.

NOTE: While random serum glucose levels correlate with home glucose monitoring results (weekly mean capillary glucose values), there is wide fluctuation within individuals. Thus, glycosylated hemoglobin (HbA1c) levels are favored to monitor glycemic control.

High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glycosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc.

GLUCOSE, POST-PRANDIAL, PLASMA-High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glycosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc. Additional test HbA1c

LIVER FUNCTION PROFILE, SERUM-LIVER FUNCTION PROFILE

Bilirubin is a yellowish pigment found in bile and is a breakdown product of normal heme catabolism. Bilirubin is excreted in bile and urine, and elevated levels may give yellow discoloration in jaundice. Elevated levels result from increased bilirubin production (eg, hemolysis and ineffective erythropoiesis), decreased bilirubin excretion (eg, obstruction and hepatitis), and abnormal bilirubin metabolism (eg, hereditary and neonatal jaundice). Conjugated (direct) bilirubin is elevated more than unconjugated (indirect) bilirubin in viral hepatitis, Drug reactions, Alcoholic liver disease Conjugated (direct) bilirubin is also elevated more than unconjugated (indirect) bilirubin when there is some kind of blockage of the bile ducts like in Gallstones getting into the bile ducts, tumors & Scarring of the bile ducts. Increased unconjugated (indirect) bilirubin may be a result of Hemolytic or pernicious anemia, Transfusion reaction & a common metabolic condition termed Gilbert syndrome, due to low levels of the enzyme that attaches sugar molecules to bilirubin.

AST is an enzyme found in various parts of the body. AST is found in the liver, heart, skeletal muscle, kidneys, brain, and red blood cells, and it is commonly measured clinically as a marker for liver health. AST levels increase during chronic viral hepatitis, blockage of the bile duct, cirrhosis of the liver, liver cancer, kidney failure, hemolytic anemia, pancreatitis, hemochromatosis. AST levels may also increase after a heart attack or strenuous activity. ALT test measures the amount of this enzyme in the blood. ALT is found mainly in the liver, but also in smaller amounts in the kidneys, heart, muscles, and pancreas. It is commonly measured as a part of a diagnostic evaluation of hepatocellular injury, to determine liver health. AST levels increase during acute hepatitis, sometimes due to a viral infection, ischemia to the liver, chronic hepatitis, obstruction of bile ducts, cirrhosis.

ALP is a protein found in almost all body tissues. Issues with higher amounts of ALP include the liver, bile ducts and bone. Elevated ALP levels are seen in Biliary obstruction, Osteoblastic bone tumors, osteomalacia, hepatitis, Hyperparathyroidism, Leukemia, Lymphoma, Paget's disease, Rickets, Sarcoidosis etc. Lower-than-normal ALP levels

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seen in Hypophosphatasia, Malnutrition, Protein deficiency, Wilson's disease. GGT is an enzyme found in cell membranes of many tissues mainly in the liver, kidney and pancreas. It is also found in other tissues including intestine, spleen, heart, brain and seminal vesicles. The highest concentration is in the kidney, but the liver is considered the source of normal enzyme activity. Serum GGT has been widely used as an index of liver dysfunction. Elevated serum GGT activity can be found in diseases of the liver, biliary system and pancreas. Conditions that increase serum GGT are obstructive liver disease, high alcohol consumption and use of enzyme-inducing drugs etc. Serum total protein, also known as total protein, is a biochemical test for measuring the total amount of protein in serum. Protein in the plasma is made up of albumin and globulin. Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenstrom's disease. Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage), Burns, Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic syndrome, Protein-losing enteropathy etc. Human serum albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by: Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance, malnutrition and wasting etc

BLOOD UREA NITROGEN (BUN), SERUM- Causes of Increased levels include Pre renal (High protein diet, Increased protein catabolism, GI haemorrhage, Cortisol, Dehydration, CHF Renal), Renal Failure, Post Renal (Malignancy, Nephrolithiasis, Prostatism)
 Causes of decreased level include Liver disease, SIADH.

CREATININE, SERUM- Higher than normal level may be due to:

- Blockage in the urinary tract
- Kidney problems, such as kidney damage or failure, infection, or reduced blood flow
- Loss of body fluid (dehydration)
- Muscle problems, such as breakdown of muscle fibers
- Problems during pregnancy, such as seizures (eclampsia), or high blood pressure caused by pregnancy (preeclampsia)

Lower than normal level may be due to:

- Myasthenia Gravis
- Muscular dystrophy


URIC ACID, SERUM- Causes of Increased levels:- Dietary (High Protein Intake, Prolonged Fasting, Rapid weight loss), Gout, Lesch nyhan syndrome, Type 2 DM, Metabolic syndrome

Causes of decreased levels- Low Zinc intake, OCP, Multiple Sclerosis

TOTAL PROTEIN, SERUM- Serum total protein, also known as total protein, is a biochemical test for measuring the total amount of protein in serum. Protein in the plasma is made up of albumin and globulin

Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C, Multiple myeloma, Waldenstrom's disease
 Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage), Burns, Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic syndrome, Protein-losing enteropathy etc.

ALBUMIN, SERUM- Human serum albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by: Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance, malnutrition and wasting etc.



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CLINICAL PATH - URINALYSIS

MEDI WHEEL FULL BODY HEALTH CHECKUP ABOVE 40FEMALE

PHYSICAL EXAMINATION, URINE

COLOR PALE YELLOW
APPEARANCE CLEAR

METHOD : DIPSTICK, MICROSCOPY

CHEMICAL EXAMINATION, URINE

PH	6.5	4.7 - 7.5	
METHOD : DIPSTICK			
SPECIFIC GRAVITY	<= 1.005	1.003 - 1.035	
METHOD : DIPSTICK			
PROTEIN	NOT DETECTED	NOT DETECTED	
METHOD : DIPSTICK			
GLUCOSE	NOT DETECTED	NOT DETECTED	
METHOD : DIPSTICK			
KETONES	NOT DETECTED	NOT DETECTED	
METHOD : DIPSTICK			
BLOOD	DETECTED (+)	NOT DETECTED	
METHOD : DIPSTICK			
BILIRUBIN	NOT DETECTED	NOT DETECTED	
METHOD : DIPSTICK (DIAZOTISED DICHLOROANILINE)			
UROBILINOGEN	NORMAL	NORMAL	
METHOD : DIPSTICK			
NITRITE	NOT DETECTED	NOT DETECTED	
METHOD : DIPSTICK			

MICROSCOPIC EXAMINATION, URINE

RED BLOOD CELLS	2 - 3	NOT DETECTED	/HPF
METHOD : MICROSCOPIC EXAMINATION			
PUS CELL (WBC'S)	2-3	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION			
EPITHELIAL CELLS	3-5	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION			
CASTS	NOT DETECTED		
METHOD : MICROSCOPIC EXAMINATION			
CRYSTALS	NOT DETECTED		



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Page 17 Of 20



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METHOD : MICROSCOPIC EXAMINATION

BACTERIA

NOT DETECTED

NOT DETECTED

METHOD : MICROSCOPIC EXAMINATION

REMARKS

URINE ANALYSIS : MICROSCOPIC EXAMINATION IS CARRIED OUT ON CENTRIFUGED URINARY SEDIMENT.

Interpretation(s)



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CYTOLOGY

MEDI WHEEL FULL BODY HEALTH CHECKUP ABOVE 40 FEMALE	RESULT PENDING
PAPANICOLAOU SMEAR	RESULT PENDING
LETTER	RESULT PENDING



View Details



View Report

PERFORMED AT :
 SRL Ltd
 Ground floor 365/6, Aaj Ka Aanand building, Shivaji Nagar
 PUNE, 411005
 MAHARASHTRA, INDIA
 Tel : 9111591115, Fax : 020 30251212
 CIN - U74899PB1995PLC045956
 Email : customercare.pune@srl.in



Patient Ref. No. 775000002438805

PATIENT NAME : SONAL BHATT

REF. DOCTOR : SELF

CODE/NAME & ADDRESS : C000138362

ACROFEMI HEALTHCARE LTD (MEDIWHEEL)

F-703, LADO SARAI, MEHRAULISOUTH WEST
DELHI

NEW DELHI 110030

8800465156

ACCESSION NO : 0030WB005257

PATIENT ID : SONAF03106830

CLIENT PATIENT ID:

ABHA NO :

AGE/SEX : 54 Years Female

DRAWN :

RECEIVED : 25/02/2023 08:23:17

REPORTED : 27/02/2023 15:46:59

Test Report Status	Preliminary	Results	Biological Reference Interval	Units
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SPECIALISED CHEMISTRY - HORMONE

MEDI WHEEL FULL BODY HEALTH CHECKUP ABOVE 40FEMALE

THYROID PANEL, SERUM

T3	128.29	58 - 159	ng/dL
T4	8.05	4.87 - 11.71	µg/dL
TSH (ULTRASENSITIVE)	1.478	0.350 - 4.940	µIU/mL

End Of Report

Please visit www.srlworld.com for related Test Information for this accession


CONDITIONS OF LABORATORY TESTING & REPORTING

1. It is presumed that the test sample belongs to the patient named or identified in the test requisition form.
2. All tests are performed and reported as per the turnaround time stated in the SRL Directory of Services.
3. Result delays could occur due to unforeseen circumstances such as non-availability of kits / equipment breakdown / natural calamities / technical downtime or any other unforeseen event.
4. A requested test might not be performed if:
 - i. Specimen received is insufficient or inappropriate
 - ii. Specimen quality is unsatisfactory
 - iii. Incorrect specimen type
 - iv. Discrepancy between identification on specimen container label and test requisition form
5. SRL confirms that all tests have been performed or assayed with highest quality standards, clinical safety & technical integrity.
6. Laboratory results should not be interpreted in isolation; it must be correlated with clinical information and be interpreted by registered medical practitioners only to determine final diagnosis.
7. Test results may vary based on time of collection, physiological condition of the patient, current medication or nutritional and dietary changes. Please consult your doctor or call us for any clarification.
8. Test results cannot be used for Medico legal purposes.
9. In case of queries please call customer care (91115 91115) within 48 hours of the report.

SRL Limited

Fortis Hospital, Sector 62, Phase VIII,
Mohali 160062

Page 20 Of 20



Dr. Swati Pravin Mulani
Lab Head



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