

CLIENT CODE : C000138394
CLIENT'S NAME AND ADDRESS :
 ACROFEMI HEALTHCARE LTD (MEDIWHEEL)
 F-703, F-703, LADO SARAI, MEHRAULI
 SOUTH WEST DELHI
 NEW DELHI 110030
 DELHI INDIA
 8800465156

SRL Ltd
 S.K. Tower, Hari Niwas, LBS Marg
 THANE, 400602
 MAHARASHTRA, INDIA
 Tel : 9111591115, Fax : CIN - U74899PB1995PLC045956
 Email : customercare.thane@srl.in

PATIENT NAME : MADHURI SHARMA **PATIENT ID : MADHF100879181**

ACCESSION NO : **0181WC000704** AGE : 43 Years SEX : Female

DRAWN : RECEIVED : 14/03/2023 08:23 REPORTED : 16/03/2023 14:05

REFERRING DOCTOR : SELF CLIENT PATIENT ID :

Test Report Status	Results	Biological Reference Interval	Units
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MEDI WHEEL FULL BODY HEALTH CHECKUP ABOVE 40FEMALE

BLOOD COUNTS, EDTA WHOLE BLOOD

HEMOGLOBIN (HB)	12.0	12.0 - 15.0	g/dL
<small>METHOD : SLS- HEMOGLOBIN DETECTION METHOD</small>			
RED BLOOD CELL (RBC) COUNT	4.09	3.8 - 4.8	mil/ μ L
<small>METHOD : HYDRODYNAMIC FOCUSING BY DC DETECTION</small>			
WHITE BLOOD CELL (WBC) COUNT	5.69	4.0 - 10.0	thou/ μ L
<small>METHOD : FLUORESCENCE FLOW CYTOMETRY</small>			
PLATELET COUNT	245	150 - 410	thou/ μ L
<small>METHOD : HYDRODYNAMIC FOCUSING BY DC DETECTION</small>			

RBC AND PLATELET INDICES

HEMATOCRIT (PCV)	37.5	36.0 - 46.0	%
<small>METHOD : CUMULATIVE PULSE HEIGHT DETECTION METHOD</small>			
MEAN CORPUSCULAR VOLUME (MCV)	91.7	83.0 - 101.0	fL
<small>METHOD : CALCULATED FROM RBC & HCT</small>			
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	29.3	27.0 - 32.0	pg
<small>METHOD : CALCULATED FROM THE RBC & HGB</small>			
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC)	32.0	31.5 - 34.5	g/dL
<small>METHOD : CALCULATED FROM THE HGB & HCT</small>			
RED CELL DISTRIBUTION WIDTH (RDW)	13.5	11.6 - 14.0	%
<small>METHOD : CALCULATED FROM RBC SIZE DISTRIBUTION CURVE</small>			
MENTZER INDEX	22.4		
MEAN PLATELET VOLUME (MPV)	10.8	6.8 - 10.9	fL
<small>METHOD : CALCULATED FROM PLATELET COUNT & PLATELET HEMATOCRIT</small>			

WBC DIFFERENTIAL COUNT

NEUTROPHILS	60	40 - 80	%
<small>METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING</small>			
LYMPHOCYTES	30	20 - 40	%
<small>METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING</small>			
MONOCYTES	6	2 - 10	%
<small>METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING</small>			
EOSINOPHILS	4	1 - 6	%
<small>METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING</small>			
ABSOLUTE NEUTROPHIL COUNT	3.38	2.0 - 7.0	thou/ μ L
<small>METHOD : FLOW CYTOMETRY WITH LIGHT SCATTERING</small>			



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METHOD : ENZYMATIC COLORIMETRIC ASSAY				
TRIGLYCERIDES	101		Normal: < 150 Borderline high: 150 - 199 High: 200 - 499 Very High: >= 500	mg/dL
METHOD : ENZYMATIC COLORIMETRIC ASSAY				
HDL CHOLESTEROL	62	High	Low HDL Cholesterol <40 High HDL Cholesterol >= 60	mg/dL
METHOD : ENZYMATIC, COLORIMETRIC				
CHOLESTEROL LDL	115	High	Adult levels: Optimal < 100 Near optimal/above optimal: 100-129 Borderline high : 130-159 High : 160-189 Very high : = 190	mg/dL
METHOD : ENZYMATIC COLORIMETRIC ASSAY				
NON HDL CHOLESTEROL	135	High	Desirable : < 130 Above Desirable : 130 -159 Borderline High : 160 - 189 High : 190 - 219 Very high : > / = 220	mg/dL
VERY LOW DENSITY LIPOPROTEIN CHOL/HDL RATIO	20.2		< OR = 30.0	mg/dL
LDL/HDL RATIO	3.2	Low	Low Risk : 3.3 - 4.4 Average Risk : 4.5 - 7.0 Moderate Risk : 7.1 - 11.0 High Risk : > 11.0	
	1.9		0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	

LIVER FUNCTION PROFILE, SERUM

BILIRUBIN, TOTAL	0.41		Upto 1.2	mg/dL
METHOD : COLORIMETRIC DIAZO				
BILIRUBIN, DIRECT	0.22		< 0.30	mg/dL
BILIRUBIN, INDIRECT	0.19		0.1 - 1.0	mg/dL
TOTAL PROTEIN	7.3		6.0 - 8.0	g/dL
METHOD : COLORIMETRIC				
ALBUMIN	4.5		3.97 - 4.94	g/dL
METHOD : COLORIMETRIC				
GLOBULIN	2.8		2.0 - 3.5	g/dL
ALBUMIN/GLOBULIN RATIO	1.6		1.0 - 2.1	RATIO





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ASPARTATE AMINOTRANSFERASE (AST/SGOT)		22	< OR = 35	U/L
METHOD : UV ABSORBANCE				
ALANINE AMINOTRANSFERASE (ALT/SGPT)		25	< OR = 35	U/L
METHOD : UV ABSORBANCE				
ALKALINE PHOSPHATASE		58	35 - 104	U/L
METHOD : COLORIMETRIC				
GAMMA GLUTAMYL TRANSFERASE (GGT)		21	0 - 40	U/L
METHOD : ENZYMATIC, COLORIMETRIC				
LACTATE DEHYDROGENASE		176	125 - 220	U/L
METHOD : UV ABSORBANCE				
BLOOD UREA NITROGEN (BUN), SERUM				
BLOOD UREA NITROGEN		7	6 - 20	mg/dL
METHOD : ENZYMATIC ASSAY				
CREATININE, SERUM				
CREATININE		0.59	0.5 - 0.9	mg/dL
METHOD : COLORIMETRIC				
BUN/CREAT RATIO				
BUN/CREAT RATIO		11.86	8.0 - 15.0	
URIC ACID, SERUM				
URIC ACID		4.7	2.4 - 5.7	mg/dL
METHOD : ENZYMATIC COLORIMETRIC ASSAY				
TOTAL PROTEIN, SERUM				
TOTAL PROTEIN		7.3	6.0 - 8.0	g/dL
METHOD : COLORIMETRIC				
ALBUMIN, SERUM				
ALBUMIN		4.5	3.97 - 4.94	g/dL
METHOD : COLORIMETRIC				
GLOBULIN				
GLOBULIN		2.8	2.0 - 3.5	g/dL
ELECTROLYTES (NA/K/CL), SERUM				
SODIUM, SERUM		133	Low 136 - 145	mmol/L
POTASSIUM, SERUM		4.14	3.5 - 5.1	mmol/L
CHLORIDE, SERUM		96	Low 98 - 107	mmol/L
PHYSICAL EXAMINATION, URINE				
COLOR		PALE YELLOW		
APPEARANCE		CLEAR		



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CHEMICAL EXAMINATION, URINE

PH	6.0	5.00 - 7.50	
SPECIFIC GRAVITY	1.010	1.010 - 1.030	
PROTEIN	NOT DETECTED	NOT DETECTED	
GLUCOSE	NOT DETECTED	NOT DETECTED	
KETONES	NOT DETECTED	NOT DETECTED	
BLOOD	DETECTED (TRACE)	NOT DETECTED	
UROBILINOGEN	NORMAL	NORMAL	
NITRITE	NOT DETECTED	NOT DETECTED	
LEUKOCYTE ESTERASE	NOT DETECTED	NOT DETECTED	

MICROSCOPIC EXAMINATION, URINE

RED BLOOD CELLS	1 - 2	NOT DETECTED	/HPF
PUS CELL (WBC'S)	2-3	0-5	/HPF
EPITHELIAL CELLS	1-2	0-5	/HPF
CASTS	NOT DETECTED		
CRYSTALS	NOT DETECTED		
BACTERIA	NOT DETECTED	NOT DETECTED	
YEAST	NOT DETECTED	NOT DETECTED	

THYROID PANEL, SERUM

T3	87.6	Non-Pregnant Women 80.0 - 200.0 Pregnant Women 1st Trimester: 105.0 - 230.0 2nd Trimester: 129.0 - 262.0 3rd Trimester: 135.0 - 262.0	ng/dL
METHOD : ELECTROCHEMILUMINESCENCE			
T4	9.02	Non-Pregnant Women 5.10 - 14.10 Pregnant Women 1st Trimester: 7.33 - 14.80 2nd Trimester: 7.93 - 16.10 3rd Trimester: 6.95 - 15.70	µg/dL
METHOD : ELECTROCHEMILUMINESCENCE			
TSH (ULTRASENSITIVE)	2.530	Non Pregnant Women 0.27 - 4.20 Pregnant Women 1st Trimester: 0.33 - 4.59 2nd Trimester: 0.35 - 4.10 3rd Trimester: 0.21 - 3.15	µIU/mL
METHOD : ELECTROCHEMILUMINESCENCE			



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PAPANICOLAOU SMEAR

TEST METHOD SNR
 METHOD : MICROSCOPIC EXAMINATION

MICROSCOPIC EXAMINATION, STOOL

REMARK SAMPLE NOT RECEIVED

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP TYPE A
 METHOD : GEL COLUMN AGGLUTINATION METHOD.

RH TYPE POSITIVE
 METHOD : GEL COLUMN AGGLUTINATION METHOD.

XRAY-CHEST

IMPRESSION NO ABNORMALITY DETECTED

TMT OR ECHO

TMT OR ECHO NEGATIVE

ECG

ECG WITHIN NORMAL LIMITS

MAMOGRAPHY (BOTH BREASTS)

MAMOGRAPHY BOTH BREASTS FEW FIBROCYSTIC CHANGES ARE NOTED BETWEEN 9-10 O'CLOCK POSITIONS IN RIGHT BREAST AND 11-12 O'CLOCK LARGEST CYST MEASURING 4.6mm ON RIGHT AND 5.4mm ON LEFT.

MEDICAL HISTORY

RELEVANT PRESENT HISTORY C/O JOINT PAINS ON & OFF.
C/O HEADACHE ON & OFF.
 RELEVANT PAST HISTORY PULMONARY KOCHS IN 2000.
 RELEVANT PERSONAL HISTORY MARRIED / 3 CHILD / VEG DIET / NO ALLERGIES / NO SMOKING / NO ALCOHOL.
 MENSTRUAL HISTORY (FOR FEMALES) REGULAR 28-32/1DAY C/O HYPOMENORRHOEA.
 LMP (FOR FEMALES) 13/3/2023.
 OBSTETRIC HISTORY (FOR FEMALES) 3FTNDA2L3
 LCB (FOR FEMALES) 14 YEARS BACK.
 RELEVANT FAMILY HISTORY NOT SIGNIFICANT
 HISTORY OF MEDICATIONS NOT SIGNIFICANT

ANTHROPOMETRIC DATA & BMI

HEIGHT IN METERS 1.63 mts
 WEIGHT IN KGS. 60 Kgs



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BMI	23	BMI & Weight Status as follows: kg/sqmts Below 18.5: Underweight 18.5 - 24.9: Normal 25.0 - 29.9: Overweight 30.0 and Above: Obese
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GENERAL EXAMINATION

MENTAL / EMOTIONAL STATE	NORMAL
PHYSICAL ATTITUDE	NORMAL
GENERAL APPEARANCE / NUTRITIONAL STATUS	HEALTHY
BUILT / SKELETAL FRAMEWORK	AVERAGE
FACIAL APPEARANCE	NORMAL
SKIN	NORMAL
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	REGULAR, ALL PERIPHERAL PULSES WELL FELT, NO CAROTID BRUIT
RESPIRATORY RATE	NORMAL

CARDIOVASCULAR SYSTEM

BP	120/80 MM HG (SUPINE)	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
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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, sulfonyleureas, 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-
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** results 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. Tissues 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, Pagets disease, Rickets, Sarcoidosis etc. Lower-than-normal ALP levels seen in Hypophosphatasia, Malnutrition, Protein deficiency, Wilsons 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.
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, Waldenstroms 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 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, Muscuophy
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- 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, Waldenstroms 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.
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.

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.



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Final		

MEDI WHEEL FULL BODY HEALTH CHECKUP ABOVE 40FEMALE

ULTRASOUND ABDOMEN
ULTRASOUND ABDOMEN
 GRADE I FATTY LIVER.

****End Of Report****
 Please visit www.srlworld.com for related Test Information for this accession

CONDITIONS OF LABORATORY TESTING & REPORTING

- | | |
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| <ol style="list-style-type: none"> 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: <ol style="list-style-type: none"> 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 | <ol style="list-style-type: none"> 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. |
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 Fortis Hospital, Sector 62, Phase VIII,
 Mohali 160062



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