

Units

PATIENT NAME: BABULAL HALEYAJI BARNDA **REF. DOCTOR: SELF** CODE/NAME & ADDRESS: C000138364 AGE/SEX

ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, LADO SARAI, MEHRAULISOUTH WEST

DELHI

**NEW DELHI 110030** 8800465156

**Test Report Status** 

ACCESSION NO: 0321XC000126

PATIENT ID : BABUM010578321

CLIENT PATIENT ID: ABHA NO

Results

DRAWN

**Biological Reference Interval** 

RECEIVED: 02/03/2024 14:34:47 REPORTED :02/03/2024 18:21:33

:45 Years

**Preliminary** 

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVR BOUMAREENDING **XRAY-CHEST RESULT PENDING ECG RESULT PENDING MEDICAL HISTORY RESULT PENDING** ANTHROPOMETRIC DATA & BMI RESULT PENDING **GENERAL EXAMINATION** RESULT PENDING CARDIOVASCULAR SYSTEM RESULT PENDING RESPIRATORY SYSTEM RESULT PENDING PER ABDOMEN RESULT PENDING **CENTRAL NERVOUS SYSTEM RESULT PENDING** MUSCULOSKELETAL SYSTEM **RESULT PENDING BASIC EYE EXAMINATION RESULT PENDING BASIC ENT EXAMINATION RESULT PENDING BASIC DENTAL EXAMINATION** RESULT PENDING **SUMMARY RESULT PENDING FITNESS STATUS** RESULT PENDING

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ARCOFEMI HEALTHCARE LTD (MEDIWHEEL
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Test Report Status Preliminary Results Units

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVR END MARTEN DINGULTRASOUND ABDOMENRESULT PENDINGTMT OR ECHORESULT PENDING

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**PATIENT NAME: BABULAL HALEYAJI BARNDA REF. DOCTOR: SELF** CODE/NAME & ADDRESS: C000138364 ACCESSION NO: 0321XC000126 AGE/SEX :45 Years ARCOFEMI HEALTHCARE LTD (MEDIWHEEL PATIENT ID : BABUM010578321 DRAWN F-703, LADO SARAI, MEHRAULISOUTH WEST CLIENT PATIENT ID: RECEIVED: 02/03/2024 14:34:47 DELHI REPORTED :02/03/2024 18:21:33 ABHA NO **NEW DELHI 110030** 8800465156

Test Report Status <u>Preliminary</u> Results Biological Reference Interval Units

HAEMATOLOGY - CBC				
MEDI WHEEL FULL BODY HEALTH CHECK UP AE	BOVE 40 MALE			
BLOOD COUNTS,EDTA WHOLE BLOOD				
HEMOGLOBIN (HB)	13.6	13.0 - 17.0	g/dL	
RED BLOOD CELL (RBC) COUNT	5.19	4.5 - 5.5	mil/μL	
WHITE BLOOD CELL (WBC) COUNT	5.56	4.0 - 10.0	thou/µL	
PLATELET COUNT	213.4	150 - 410	thou/µL	
RBC AND PLATELET INDICES				
HEMATOCRIT (PCV)	43.3	40.0 - 50.0	%	
MEAN CORPUSCULAR VOLUME (MCV)	83.5	83.0 - 101.0	fL	
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	26.2 Low	27.0 - 32.0	pg	
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC)	31.5	31.5 - 34.5	g/dL	
RED CELL DISTRIBUTION WIDTH (RDW)	16.0 High	11.6 - 14.0	%	
MENTZER INDEX	16.1			
MEAN PLATELET VOLUME (MPV)	9.4	6.8 - 10.9	fL	
WBC DIFFERENTIAL COUNT				
NEUTROPHILS	55	40 - 80	%	
LYMPHOCYTES	33	20 - 40	%	
MONOCYTES	10	2.0 - 10.0	%	
EOSINOPHILS	2	1.0 - 6.0	%	
BASOPHILS	0	0 - 1	%	
ABSOLUTE NEUTROPHIL COUNT	3.06	2.0 - 7.0	thou/µL	
ABSOLUTE LYMPHOCYTE COUNT	1.83	1.0 - 3.0	thou/µL	
ABSOLUTE MONOCYTE COUNT	0.56	0.2 - 1.0	thou/µL	
ABSOLUTE EOSINOPHIL COUNT	0.11	0.02 - 0.50	thou/µL	
ABSOLUTE BASOPHIL COUNT	0 Low	0.02 - 0.10	thou/µL	

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NEUTROPHIL LYMPHOCYTE RATIO (NLR) 1.7

MORPHOLOGY

NORMOCYTIC NORMOCHROMIC **RBC** 

**WBC** NORMAL MORPHOLOGY

**ADEQUATE PLATELETS** 

NO PREMATURE CELLS ARE SEEN. MALARIAL PARASITE NOT DETECTED. REMARKS

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)

This ratio element is a calculated parameter and out of NABL scope.

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Units

mg/dL

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**HAEMATOLOGY** 

PATIENT ID

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MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

**ERYTHROCYTE SEDIMENTATION RATE (ESR), EDTA BLOOD** 

15 High E.S.R 0 - 14

mm at 1 hr

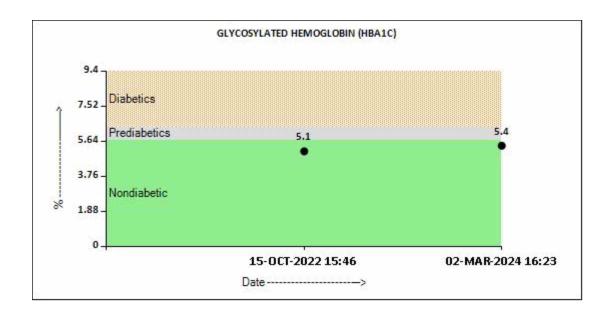
GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE **BLOOD** 

% HBA1C 5.4 Non-diabetic: < 5.7

Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5Therapeutic goals: < 7.0 Action suggested: > 8.0

< 116.0

(ADA Guideline 2021) ESTIMATED AVERAGE GLUCOSE(EAG) 108.3



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<b>Interpretation(s)</b>

ERYTHROCYTE SEDIMENTATION RATE (ESR), EDTA BLOOD-<b>TEST DESCRIPTION</b>:

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. <br/>
<br/>
<br/>
<br/>
<br/>
difficulty of the control of the con

<br/>
<br/> Pregnancy, Estrogen medication, Aging.

Finding a very accelerated ESR<b>(>100 mm/hour)</b> 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. <b > Decreased </b > in: Polycythermia vera, Sickle cell anemia

#### <b>LIMITATIONS</b>

<b>False elevated</b> ESR : Increased fibrinogen, Drugs(Vitamin A, Dextran etc), Hypercholesterolemia

<br/> <br/> False Decreased</b> : Poikilocytosis, (SickleCells, 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. GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD-<br/>b>Used For</b>:
- 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 wellcontrolled type 2 diabetic patients) to determine whether a patients metabolic control has remained continuously within the target range. 1. eAG (Estimated average glucose) converts percentage HbA1c to md/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

- <br/>
  <br/> anemia) will falsely lower HbA1c test résults.Fructosamine is recommended in these patients which indicates diabetes control over 15 days.
- 2. Vitamin C & E are reported to falsely lower test results. (possibly by inhibiting glycation of hemoglobin.
  3. Iron deficiency anemia is reported to increase test results. Hypertriglyceridemia, uremia, hyperbilirubinemia, chronic alcoholism, chronic ingestion of salicylates & opiates addiction are reported to interfere with some assay methods, falsely increasing results.

  4. 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 paltform (Boronate affinity chromatography) is recommended for testing of HbA1c.Abnormal Hemoglobin electrophoresis (HPLC method) is recommended for detecting a hemoglobinopathy

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# **IMMUNOHAEMATOLOGY**

## MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

**ABO GROUP & RH TYPE, EDTA WHOLE BLOOD** 

TYPE O **ABO GROUP POSITIVE** RH TYPE

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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CLIENT PATIENT ID:

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ABHA NO

# **BIOCHEMISTRY**

# MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

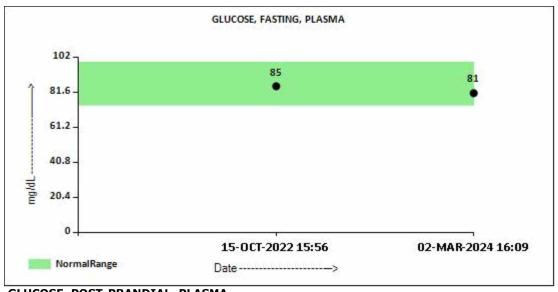
**GLUCOSE FASTING, FLUORIDE PLASMA** 

FBS (FASTING BLOOD SUGAR)

81

74 - 99

mg/dL



GLUCOSE, POST-PRANDIAL, PLASMA

PPBS(POST PRANDIAL BLOOD SUGAR)

66 Low

70 - 140

mg/dL

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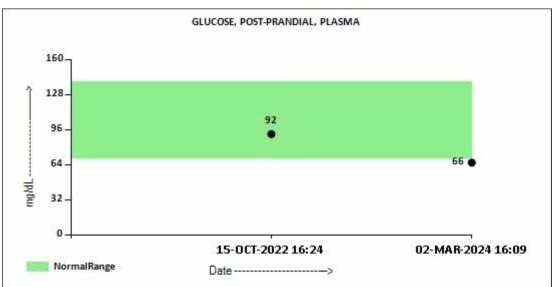
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LIPID PROFILE WITH CALCULATED LDL

CHOLESTEROL, TOTAL	186	Desirable: < 200 mg/dL BorderlineHigh: 200 - 239 High: > or = 240	
TRIGLYCERIDES	46	Desirable: $< 150$ mg/dL BorderlineHigh: $150 - 199$ High: $200 - 499$ Very High: $> or = 500$	
HDL CHOLESTEROL	77 High	< 40 Low mg/dL > or = 60 High	
CHOLESTEROL LDL	100	Adult levels: mg/dL Optimal < 100 Near optimal/above optimal: 100-129 Borderline high: 130-159 High: 160-189 Very high: = 190	
NON HDL CHOLESTEROL	109	Desirable: Less than 130 mg/dL Above Desirable: 130 - 159 Borderline High: 160 - 189	

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High: 190 - 219



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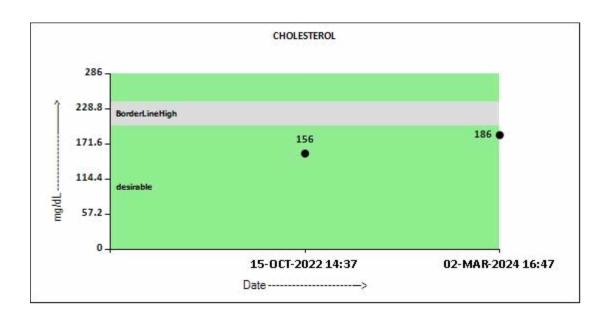
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Test Report Status <u>Preliminary</u>	Results	Biological Reference Interval Units	
		Very high: $>$ or $= 220$	
VERY LOW DENSITY LIPOPROTEIN	9.2	< or = 30 mg/dL	
CHOL/HDL RATIO	2.4 Low	3.3 - 4.4	
LDL/HDL RATIO	1.3	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate	
·			
		Risk	
		>6.0 High Risk	



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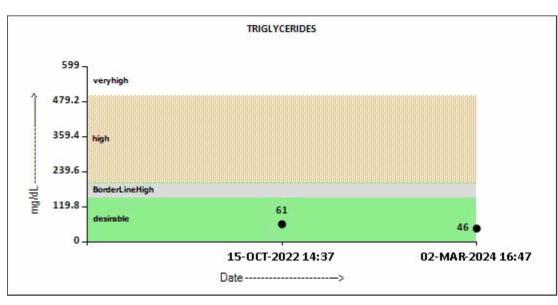
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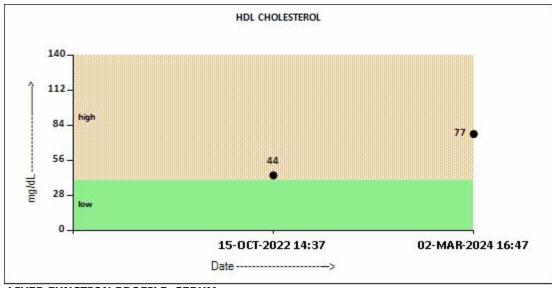
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Test Report Status Preliminary

Results

**Biological Reference Interval Units** 





LIVER FUNCTION PROFILE, SERUM

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BILIRUBIN, TOTAL	0.41	Upto 1.2	mg/dL
BILIRUBIN, DIRECT	0.21 High	Upto 0.2	mg/dL
BILIRUBIN, INDIRECT	0.20	0.00 - 1.00	mg/dL
TOTAL PROTEIN	7.5	6.4 - 8.3	g/dL
ALBUMIN	5.0	3.5 - 5.2	g/dL
GLOBULIN	2.5	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO	2.0	1.0 - 2.0	RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	19	0 - 40	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT)	8	0 - 41	U/L
ALKALINE PHOSPHATASE	99	40 - 129	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT)	13	8 - 61	U/L
LACTATE DEHYDROGENASE	181	135 - 225	U/L
BLOOD UREA NITROGEN (BUN), SERUM			
BLOOD UREA NITROGEN	9	6 - 20	mg/dL

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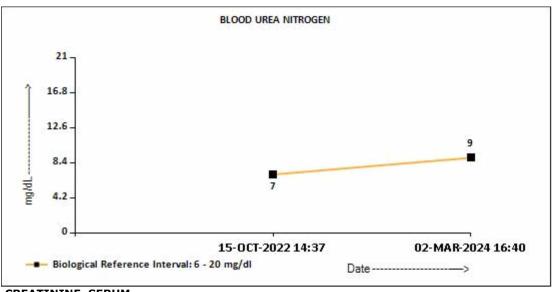
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**CREATININE, SERUM** 

CREATININE 0.80 0.70 - 1.30mg/dL

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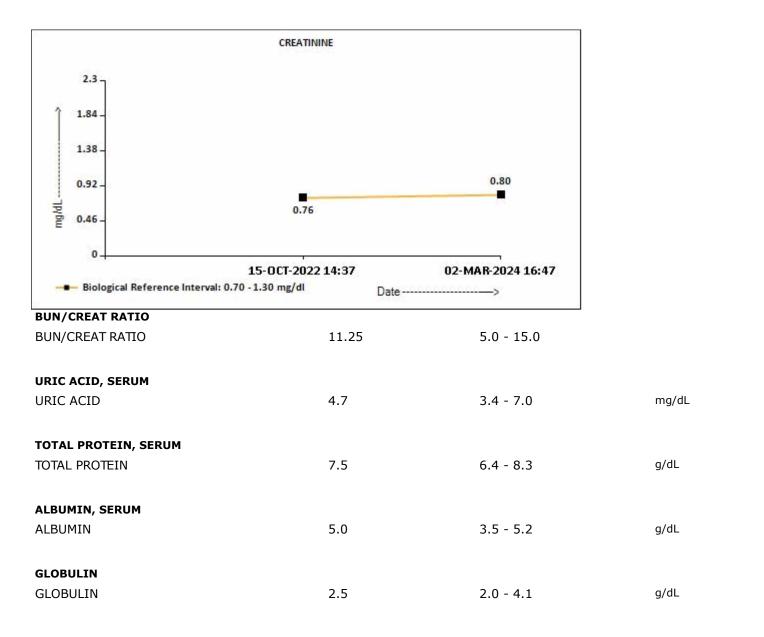
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# **ELECTROLYTES (NA/K/CL), SERUM**

SODIUM, SERUM	139.7	136 - 145	mmol/L
POTASSIUM, SERUM	4.45	3.3 - 5.1	mmol/L
CHLORIDE, SERUM	103.7	98 - 106	mmol/L

<b>Interpretation(s)</b>

GLUCOSE FASTING, FLUORIDE PLASMA-<b>TEST DESCRIPTION</b>

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

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sulfonylureas,tolbutamide,and other oral hypoglycemic agents.

<br/>
<br 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 Glyosuria, 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 Glyosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc.Additional test HbA1c LIVER FUNCTION PROFILE, SERUM-

<br/>
<br/> 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.

<br/>
<br/> 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.
<br/>
<br/> 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.

<a href="mailto:specification-normal-nor

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.

<br/>
<br/> disease.Lower-than-normal levels may be due to: Agammaglobulinemia,Bleeding (hemorrhage),Burns,Glomerulonephritis,Liver disease, Malabsorption,Malnutrition,Nephrotic syndrome,Protein-losing enteropathy etc.

<br/>

BLOOD UREA NITROGEN (BUN), SERUM-<br/>b>Causes of Increased<br/>/b> levels include Pre renal (High protein diet, Increased protein catabolism, GI haemorrhage, Cortisol, Dehydration, CHF Renal), Renal Failure, Post Renal (Malignancy, Nephrolithiasis, Prostatism)

<br/>

• 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,

**Dr.Miral Gaiera Consultant Pathologist** 





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CODE/NAME & ADDRESS: C000138364 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, LADO SARAI, MEHRAULISOUTH WEST

DELHI

**NEW DELHI 110030** 8800465156

ACCESSION NO: 0321XC000126

PATIENT ID : BABUM010578321

CLIENT PATIENT ID: ABHA NO

AGE/SEX DRAWN

RECEIVED: 02/03/2024 14:34:47

:45 Years

REPORTED :02/03/2024 18:21:33

**Test Report Status** Results **Biological Reference Interval** Units **Preliminary** 

such as breakdown of muscle fibers, Problems during pregnancy, such as seizures (eclampsia)), or high blood pressure caused by pregnancy (preeclampsia) <b>Lower than normal level may be due to:</b>• Myasthenia Gravis, Muscuophy URIC ACID, SERUM-<b>Causes of Increased levels:</b>-Dietary(High Protein Intake,Prolonged Fasting,Rapid weight loss),Gout,Lesch nyhan syndrome,Type 2

DM, Metabolic syndrome <br/>
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.<br/>
<br/>
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.<br/>
<br/>
<br cb>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. <b>Low blood albumin levels (hypoalbuminemia) can be caused by: </b> 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.

Dr.Miral Gajera **Consultant Pathologist** 



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CODE/NAME & ADDRESS: C000138364 ACCESSION NO: 0321XC000126 AGE/SEX :45 Years Male

ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, LADO SARAI, MEHRAULISOUTH WEST

DELHI

**NEW DELHI 110030** 

8800465156

PATIENT ID : BABUM010578321

CLIENT PATIENT ID: ABHA NO

DRAWN

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**Test Report Status** Results Biological Reference Interval Units **Preliminary** 

## **CLINICAL PATH - URINALYSIS**

## MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

PHYSICAL EXAMINATION, URINE

**COLOR** Yellow **APPEARANCE** Clear

## CHEMICAL EXAMINATION, URINE

PH	5.5	4.7 - 7.5
SPECIFIC GRAVITY	<=1.005	1.003 - 1.035
PROTEIN	NOT DETECTED	NOT DETECTED
GLUCOSE	NOT DETECTED	NEGATIVE
KETONES	NOT DETECTED	NOT DETECTED
BLOOD	NOT DETECTED	NOT DETECTED
BILIRUBIN	NOT DETECTED	NOT DETECTED
UROBILINOGEN	NORMAL	NORMAL
NITRITE	NOT DETECTED	NOT DETECTED
LEUKOCYTE ESTERASE	NOT DETECTED	NOT DETECTED

# MICROSCOPIC EXAMINATION, URINE

RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
PUS CELL (WBC'S)	0-1	0-5	/HPF
EPITHELIAL CELLS	0-1	0-5	/HPF

NOT DETECTED **CASTS** NOT DETECTED **CRYSTALS** 

**BACTERIA** NOT DETECTED NOT DETECTED YEAST **NOT DETECTED NOT DETECTED** 

MICROSCOPIC EXAMINATION OF URINE IS CARRIED OUT ON **REMARKS** 

CENTRIFUGED URINARY SEDIMENT.

**Dr.Miral Gaiera Consultant Pathologist** 



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Gujrat, India





PATIENT NAME: BABULAL HALEYAJI BARNDA REF. DOCTOR: SELF

CODE/NAME & ADDRESS: C000138364

ARCOFEMI HEALTHCARE LTD (MEDIWHEEL
F-703, LADO SARAI, MEHRAULISOUTH WEST

DELHI

NEW DELHI 110030 8800465156 ACCESSION NO : **0321XC000126** 

PATIENT ID : BABUM010578321

CLIENT PATIENT ID: ABHA NO : AGE/SEX :45 Years
DRAWN :

N :

RECEIVED : 02/03/2024 14:34:47 REPORTED : 02/03/2024 18:21:33

Test Report Status Preliminary Results Biological Reference Interval Units

Dr.Miral Gajera Consultant Pathologist



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CODE/NAME & ADDRESS: C000138364 ACCESSION NO: 0321XC000126 AGE/SEX ARCOFEMI HEALTHCARE LTD (MEDIWHEEL

F-703, LADO SARAI, MEHRAULISOUTH WEST

DELHI

**NEW DELHI 110030** 8800465156

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:45 Years

**Test Report Status** Results Biological Reference Interval Units **Preliminary** 

# **CLINICAL PATH - STOOL ANALYSIS**

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVR BOUMARENDING PHYSICAL EXAMINATION, STOOL RESULT PENDING **CHEMICAL EXAMINATION, STOOL** RESULT PENDING MICROSCOPIC EXAMINATION, STOOL RESULT PENDING

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**REF. DOCTOR: SELF** PATIENT NAME: BABULAL HALEYAJI BARNDA

CODE/NAME & ADDRESS: C000138364 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, LADO SARAI, MEHRAULISOUTH WEST

DELHI

**NEW DELHI 110030** 

8800465156

ACCESSION NO: 0321XC000126 AGE/SEX

PATIENT ID : BABUM010578321

CLIENT PATIENT ID: ABHA NO

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:45 Years

Test Report Status Results Biological Reference Interval Units **Preliminary** 

## **SPECIALISED CHEMISTRY - HORMONE**

## MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

#### THYROID PANEL, SERUM

ТЗ	93.73	80.0 - 200.0	ng/dL
T4	7.98	5.10 - 14.10	μg/dL
TSH (ULTRASENSITIVE)	1.540	0.270 - 4.200	μIU/mL

\*\*End Of Report\*\* Please visit www.agilusdiagnostics.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 AGILUS 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. AGILUS Diagnostics 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.
- Test results cannot be used for Medico legal purposes.
- 9. In case of gueries please call customer care (91115 91115) within 48 hours of the report.

# **Agilus Diagnostics Ltd**

Fortis Hospital, Sector 62, Phase VIII, Mohali 160062

**Dr.Miral Gaiera Consultant Pathologist** 





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