

**Biological Reference Interval** Units

PATIENT NAME : B D PRASAD SAMMANGI	REF. DOCTOR :	DR. ARCOFEMI HEALTHCARE LTD (MEDIWHEEL
CODE/NAME & ADDRESS : C000138398 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST DELHI NEW DELHI 110030 8800465156	ACCESSION NO : <b>0194XB002750</b> РАПЕНТ ID : BDPRM080284194 СЫТЕЛТВАПЕНТ ID:	AGE/SEX :40 Years Male DRAWN : RECEIVED :26/02/2024 09:36:33 REPORTED :26/02/2024 12:57:52

Results

MEDI WHEEL FULL	<b>BODY HEALTH CHECK UP</b>	ABOVERENDING

**Preliminary** 

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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Test Report Status <u>Preliminary</u>	Results	Units

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVR BOUMARENDING **ULTRASOUND ABDOMEN RESULT PENDING** 

TMT OR ECHO

**RESULT PENDING** 

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Results

н	AEMATOLOGY - CBC		
EDI WHEEL FULL BODY HEALTH CHECK UP A	BOVE 40 MALE		
LOOD COUNTS,EDTA WHOLE BLOOD			
EMOGLOBIN (HB)	14.9	13.0 - 17.0	g/dL
ED BLOOD CELL (RBC) COUNT	4.79	4.5 - 5.5	mil/µL
HITE BLOOD CELL (WBC) COUNT	6.70	4.0 - 10.0	thou/µL
LATELET COUNT	254	150 - 410	thou/µL
BC AND PLATELET INDICES			
EMATOCRIT (PCV)	45.6	40 - 50	%
IEAN CORPUSCULAR VOLUME (MCV)	95.0	83 - 101	fL
IEAN CORPUSCULAR HEMOGLOBIN (MCH)	31.2	27.0 - 32.0	pg
IEAN CORPUSCULAR HEMOGLOBIN ONCENTRATION (MCHC)	32.8	31.5 - 34.5	g/dL
ED CELL DISTRIBUTION WIDTH (RDW)	12.9	11.6 - 14.0	%
IENTZER INDEX	19.8		
IEAN PLATELET VOLUME (MPV)	7.9	6.8 - 10.9	fL
/BC DIFFERENTIAL COUNT			
EUTROPHILS	53	40 - 80	%
YMPHOCYTES	35	20 - 40	%
IONOCYTES	9	2 - 10	%
OSINOPHILS	3	1 - 6	%
ASOPHILS	0	0 - 2	%
BSOLUTE NEUTROPHIL COUNT	3.55	2.0 - 7.0	thou/µL
BSOLUTE LYMPHOCYTE COUNT	2.35	1.0 - 3.0	thou/µL
BSOLUTE MONOCYTE COUNT	0.60	0.2 - 1.0	thou/µL
BSOLUTE EOSINOPHIL COUNT	0.20	0.02 - 0.50	thou/µL
BSOLUTE BASOPHIL COUNT	0.00 Low	0.02 - 0.10	thou/µL



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NEUTROPHIL LYMPHOCYTE RATIO (NLR)	1.5	

RBC NORMOCYTIC NORMOCHROMIC RBC. NORMAL COUNT & DISTIBUTION , NO ABNORMAL CELLS / IMMATURE WBC CELLS. ADEQUATE & DISCRETELY PRESENT. NO HAEMOPARASITES SEEN. PLATELETS **IMPRESSION** NORMOCYTIC NORMOCHROMIC BLOOD PICTURE.

<b>Interpretation(s)</b>

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

(<15) In patients with mile ocycle anaema. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HDA2 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)

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



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Results

	HAEMATOLOGY		
MEDI WHEEL FULL BODY HEALTH CHECK UP	PABOVE 40 MALE		
ERYTHROCYTE SEDIMENTATION RATE (ESR BLOOD	),EDTA		
E.S.R	05	0 - 14	mm at 1 hr
GLYCOSYLATED HEMOGLOBIN(HBA1C), ED BLOOD	TA WHOLE		
HBA1C	5.7	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 ADA Target: 7.0 Action suggested: > 8.0	%
ESTIMATED AVERAGE GLUCOSE(EAG)	116.9 High	< 116.0	mg/dL

<b>Interpretation(s)</b>

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

**Preliminary** 

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. <b>TEST INTERPRETATION</b>

<b>Increase</b> in: Infections, Vasculities, Inflammatory arthritis, Renal disease, Anemia, Malignancies and plasma cell dyscrasias, Acute allergy Tissue injury, 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 <b>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.



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GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD-<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 well-

controlled 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.

eAG gives an evaluation of blood glucose levels for the last couple of months.
 eAG is calculated as eAG (mg/dl) = 28.7 \* HbA1c - 46.7

<b>HbA1c Estimation can get affected due to :</b>

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.

 Z.Vitamin C & E are reported to falsely lower test results. (possibly by inhibiting glycaridemia, 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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L	IMMUNOHAEMATOLOGY	
MEDI WHEEL FULL BODY HEAL	TH CHECK UP ABOVE 40 MALE	
ABO GROUP & RH TYPE, EDTA	WHOLE BLOOD	
ABO GROUP	TYPE A	
RH TYPE	POSITIVE	

<b>Interpretation(s)</b> 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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F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST DELHI NEW DELHI 110030 8800465156	EPILA 1000101121	RECEIVED : 26/02/2024 09:36:33 REPORTED :26/02/2024 12:57:52

Results

BIOCHEMISTRY					
MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE					
GLUCOSE FASTING, FLUORIDE PLASMA					
FBS (FASTING BLOOD SUGAR)	115 High	Normal : < 100 Pre-diabetes: 100-125 Diabetes: >/=126	mg/dL		
GLUCOSE, POST-PRANDIAL, PLASMA					
PPBS(POST PRANDIAL BLOOD SUGAR)	137	70 - 140	mg/dL		
LIPID PROFILE WITH CALCULATED LDL					
CHOLESTEROL, TOTAL	189	< 200 Desirable 200 - 239 Borderline High >/= 240 High	mg/dL		
TRIGLYCERIDES	119	< 150 Normal 150 - 199 Borderline High 200 - 499 High >/=500 Very High	mg/dL		
HDL CHOLESTEROL	45	< 40 Low >/=60 High	mg/dL		
CHOLESTEROL LDL	120 High	< 100 Optimal 100 - 129 Near optimal/ above optima 130 - 159 Borderline High 160 - 189 High >/= 190 Very High	mg/dL I		
NON HDL CHOLESTEROL	144 High	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	23.8	= 30.0</td <td>mg/dL</td>	mg/dL		



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F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST DELHI	ABHAN BATIENT ID:		RECEIVED : 26/02/2024 09:36:33	
NEW DELHI 110030			REPORTED :26/02/2024 12:57:52	
8800465156				
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CHOL/HDL RATIO	4.2	3.3 - 4.4		
		Low Risk		
		4.5 - 7.0		
		Average F 7.1 - 11.0		
		Moderate		
		> 11.0		
		High Risk		
LDL/HDL RATIO	2.7		Desirable/Low Risk Borderline/Moderate	
		Risk	Dordenine/Hoderate	
		>6.0 High	n Risk	
LIVER FUNCTION PROFILE, SERUM	1.20 High	0.2 1.0	ma/dl	
BILIRUBIN, TOTAL	-	0.2 - 1.0	mg/dL mg/dL	
BILIRUBIN, DIRECT	0.10	0.0 - 0.2	-	
BILIRUBIN, INDIRECT	1.10 High	0.1 - 1.0	mg/dL	
	7.9	6.4 - 8.2	g/dL	
ALBUMIN	4.4	3.4 - 5.0	g/dL	
	3.5	2.0 - 4.1	g/dL	
ALBUMIN/GLOBULIN RATIO	1.3	1.0 - 2.1	RATIO	
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	24	15 - 37	U/L	
ALANINE AMINOTRANSFERASE (ALT/SGPT)	29	< 45.0	U/L	
ALKALINE PHOSPHATASE	65	30 - 120	U/L	
GAMMA GLUTAMYL TRANSFERASE (GGT)	21	15 - 85	U/L	
LACTATE DEHYDROGENASE	176	85 - 227	U/L	
BLOOD UREA NITROGEN (BUN), SERUM				
BLOOD UREA NITROGEN	14	6 - 20	mg/dL	



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DATIENT NAME - D D DAGAD CAMMANCI			
PATIENT NAME : B D PRASAD SAMMANGI	KE	F. DOCTOR : DR. ARCOFEM (MEDIWHEEL	I HEALIHCARE LID
CODE/NAME & ADDRESS : C000138398	ACCESSION NO : 0194XB	002750 AGE/SEX :	40 Years Male
ARCOFEMI HEALTHCARE LTD (MEDIWHEEL	PATIENT ID : BDPRMO	30284194 DRAWN :	
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CREATININE, SERUM			
CREATININE	0.82 Low	0.90 - 1.30	mg/dL
BUN/CREAT RATIO			
BUN/CREAT RATIO	17.07 High	5.00 - 15.00	
URIC ACID, SERUM			
URIC ACID	6.8	3.5 - 7.2	mg/dL
	0.0	5.5 7.2	
	7.0	6.4.0.2	a /dl
TOTAL PROTEIN	7.9	6.4 - 8.2	g/dL
ALBUMIN, SERUM			
ALBUMIN	4.4	3.4 - 5.0	g/dL
GLOBULIN			
GLOBULIN	3.5	2.0 - 4.1	g/dL
ELECTROLYTES (NA/K/CL), SERUM			
SODIUM, SERUM	135.8 Low	137 - 145	mmol/L
POTASSIUM, SERUM	4.22	3.6 - 5.0	mmol/L



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CHLORIDE, SERUM

101.9

98 - 107

mmol/L

## Comments

NOTE: KINDLY CORRELATE THE RESULT WITH CLINICAL & THERAPEUTIC HISTORY.

<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 the urine

<b>Increased in</b>:Diabetes mellitus, Cushing's syndrome (10 – 15%), chronic pancreatitis (30%). Drugs:corticosteroids,phenytoin, estrogen, thiazides.<b>Decreased in </b>: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.

6b>NOTE:
Vb/lile 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 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-

cb>Bilirubin</b> 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.<b>Elevated levels</b> 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. <b>AST</b> 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 bild 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.

Ab>ALP
/b> 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

Ab reversised in trypophosphatasia, number donot declarate, wissing accesses.
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Ab reversised in th 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.

<b>Total Protein</b> 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.

<b>Albumin</b> 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-<b>Causes of Increased</b> levels include Pre renal (High protein diet, Increased protein catabolism, GI haemorrhage,

Cortisol, Dehydration, CHF Renal), Renal Failure, Post Renal (Malignancy, Nephrolithiasis, Prostatism)

<b>Causes of decreased</b> level include Liver disease, SIADH.

CREATININE, SERUM-<b>Higher than normal level may be due to:</b>

• 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,



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PATIENT NAME : B D PRASAD SAMMANGI		DR. ARCOFEMI HEALTHCARE LTD MEDIWHEEL
CODE/NAME & ADDRESS : C000138398 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST DELHI NEW DELHI 110030 8800465156	ACCESSION NO : <b>0194XB002750</b> PATIENT ID : BDPRM080284194 SEIFN BATIENT ID:	AGE/SEX :40 Years Male DRAWN : RECEIVED :26/02/2024 09:36:33 REPORTED :26/02/2024 12:57:52
Test Report Status <u>Preliminary</u>	Results Biological	Reference Interval Units

such as breakdown of muscle fibers, Problems during pregnancy, such as seizures (eclampsia)), or high blood pressure caused by pregnancy (preeclampsia)

Colore than normal level may be due to:
Myasthenia Gravis, Muscuophy
URIC ACID, SERUM-<br/>cb>Causes of Increased levels:</br>

URIC ACID, SERUM-<br/>cb>Causes of Increased levels: 

<b>Lower-than-normal levels may be due to:</b> 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.



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Dr.Kamidi Varun **Consultant Pathologist** 

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PATIENT NAME : B D PRASAD SAMMANGI		DR. ARCOFEMI HEALTHCARE LTD MEDIWHEEL
CODE/NAME & ADDRESS : C000138398 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST DELHI NEW DELHI 110030 8800465156	ACCESSION NO : <b>0194XB002750</b> РАТІЕНТ ID : BDPRM080284194 АНЕЛТРАПЕНТ ID:	AGE/SEX :40 Years Male DRAWN : RECEIVED :26/02/2024 09:36:33 REPORTED :26/02/2024 12:57:52
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PATIENT NAME : B D PRASAD SAMMANGI	REF. DOCTOR	: DR. ARCOFEMI HEALTHCARE LTD (MEDIWHEEL
CODE/NAME & ADDRESS : C000138398 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST DELHI NEW DELHI 110030 8800465156	ACCESSION NO : <b>0194XB002750</b> РАПЕНТ ID : BDPRM080284194 Сынат Матіент ID:	AGE/SEX :40 Years Male DRAWN : RECEIVED :26/02/2024 09:36:33 REPORTED :26/02/2024 12:57:52
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CLINI	CAL PATH - STOOL ANALYSIS
MEDI WHEEL FULL BODY HEALTH CHECK UP	ABOVREDUMARENDING
PHYSICAL EXAMINATION, STOOL	RESULT PENDING
CHEMICAL EXAMINATION, STOOL	RESULT PENDING
MICROSCOPIC EXAMINATION, STOOL	RESULT PENDING

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Biological Reference Interval Units

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Results

	SPECIALISED CHEMISTRY - I	IORMONE	
MEDI WHEEL FULL BODY HEALTH CH	IECK UP ABOVE 40 MALE		
THYROID PANEL, SERUM			
ТЗ	114.60	80.0 - 200.0	ng/dL
T4	6.91	5.10 - 14.10	µg/dL
TSH (ULTRASENSITIVE)	2.440	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	5. AGILUS Diagnostics confirms that all tests have been
named or identified in the test requisition form.	performed or assayed with highest quality standards,
2. All tests are performed and reported as per the	clinical safety & technical integrity.
turnaround time stated in the AGILUS Directory of Services.	6. Laboratory results should not be interpreted in
3. Result delays could occur due to unforeseen	isolation; it must be correlated with clinical information and
circumstances such as non-availability of kits / equipment	be interpreted by registered medical practitioners only to
breakdown / natural calamities / technical downtime or any	determine final diagnosis.
other unforeseen event.	<ol><li>Test results may vary based on time of collection,</li></ol>
<ol><li>A requested test might not be performed if:</li></ol>	physiological condition of the patient, current medication or
i. Specimen received is insufficient or inappropriate	nutritional and dietary changes. Please consult your doctor
ii. Specimen quality is unsatisfactory	or call us for any clarification.
iii. Incorrect specimen type	8. Test results cannot be used for Medico legal purposes.
iv. Discrepancy between identification on specimen	<ol><li>In case of queries please call customer care</li></ol>
container label and test requisition form	(91115 91115) within 48 hours of the report.
	·
	Agilus Diagnostics Ltd
	Fortis Hospital, Sector 62, Phase VIII,
	Mohali 160062



Dr.Kamidi Varun **Consultant Pathologist** 







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