



PATIENT NAME : RAJSHEKAR PABBA		REF. DOCTOR : SELF				
		ACCESSION N	0 : 0002WL024266	AGE/SEX	:46 Years M	lale
RAJSHEKAR PABBA		PATIENT ID	: RAJSM3110772	DRAWN	:16/12/2023 09	:01:42
		CLIENT PATIEN	IT ID:	RECEIVED	: 16/12/2023 09	:03:01
		ABHA NO	:	REPORTED	:16/12/2023 15	:55:32
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Test Report Status	<u>Preliminary</u>	Results	Biolog	ical Reference	e Interval Unit	ts

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVR BOUMARENDING

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

Page 1 Of 17



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PATIENT NAME : RAJSHEKAR PABBA	REF. DOCTOR :	SELF
	ACCESSION NO : 0002WL024266	AGE/SEX : 46 Years Male
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Test Report Status <u>Preliminary</u>	Results	Units

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE ULTRASOUND ABDOMEN **ULTRASOUND ABDOMEN** GRADE I FATTY LIVER. TMT OR ECHO

RESULT PENDING







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Test Report Status

Preliminary



Biological Reference Interval Units



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Results

HAEMATOLOGY - CBC				
MEDI WHEEL FULL BODY HEALTH CHECK UP AB	OVE 40 MALE			
BLOOD COUNTS, EDTA WHOLE BLOOD				
HEMOGLOBIN (HB)	16.0	13.0 - 17.0	g/dL	
RED BLOOD CELL (RBC) COUNT	5.51 High	4.5 - 5.5	mil/µL	
WHITE BLOOD CELL (WBC) COUNT	7.75	4.0 - 10.0	thou/µL	
PLATELET COUNT	247	150 - 410	thou/µL	
RBC AND PLATELET INDICES				
HEMATOCRIT (PCV)	48.8	40 - 50	%	
MEAN CORPUSCULAR VOLUME (MCV)	88.6	83.0 - 101.0	fL	
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	29.1	27.0 - 32.0	pg	
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC)	32.9	31.5 - 34.5	g/dL	
RED CELL DISTRIBUTION WIDTH (RDW)	13.4	11.6 - 14.0	%	
MENTZER INDEX	16.1			
MEAN PLATELET VOLUME (MPV)	10.4	6.8 - 10.9	fL	
WBC DIFFERENTIAL COUNT				
NEUTROPHILS	54	40 - 80	%	
LYMPHOCYTES	37	20 - 40	%	
MONOCYTES	6	2 - 10	%	
EOSINOPHILS	2	1 - 6	%	
BASOPHILS	1	0 - 1	%	
ABSOLUTE NEUTROPHIL COUNT	4.19	2.0 - 7.0	thou/µL	
ABSOLUTE LYMPHOCYTE COUNT	2.87	1.0 - 3.0	thou/µL	
ABSOLUTE MONOCYTE COUNT	0.47	0.2 - 1.0	thou/µL	
ABSOLUTE EOSINOPHIL COUNT	0.16	0.02 - 0.50	thou/µL	
ABSOLUTE BASOPHIL COUNT	0.08	0.02 - 0.10	thou/µL	
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	1.5			

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

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Page 3 Of 17

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Test Report Status Preliminary	Results Biological	Reference Interval Units		

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 4 Of 17



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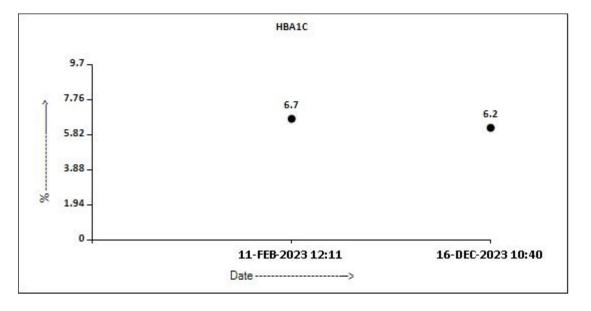






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Test Report Status <u>Prelimir</u>	nary Results	Biological Reference Interval Units			

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HAEMATOLOGY						
MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE						
ERYTHROCYTE SEDIMENTATION RATE (ESR) BLOOD	,EDTA					
E.S.R	2	= or < 10	mm at 1 hr			
GLYCOSYLATED HEMOGLOBIN(HBA1C), EDT/ BLOOD	A WHOLE					
HBA1C	6.2 High	Non-diabetic Adult < 5.7 Pre-diabetes 5.7 - 6.4	%			
		Diabetes diagnosis: > or = Therapeutic goals: < 7.0 Action suggested : > 8.0 (ADA Guideline 2021)	6.5			
ESTIMATED AVERAGE GLUCOSE(EAG)	131.2 High	< 116	mg/dL			



Comments

Advised : Kindly correlate clinically.

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Test Report Status	Preliminary	Results	Biolo	ogical F	Reference	Interval U	Jnits

Interpretation(s)

ERYTHROCYTE SEDIMENTATION RATE (ESR), EDTA 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, Vasculities, Inflammatory arthritis, Renal disease, Anemia, Malignancies and plasma cell dyscrasias, Acute allergy Tissue 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: Polycythermia vera, Sickle cell anemia

LIMITATIONS

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

False Decreased : 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-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 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

HbA1c Estimation can get affected due to :

1. 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.

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.

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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Page 6 Of 17

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IMMUNOHAEMATOLOGY				
MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE				
ABO GROUP & RH TYPE, EDTA WHOLE BLOOD				
ABO GROUP	В			
RH TYPE	POSITIVE			

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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Page 7 Of 17

View Report

Details



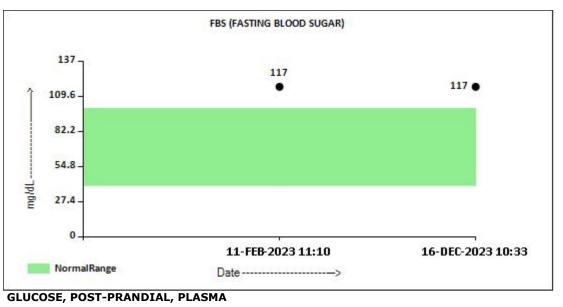
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Results Biological	Reference Interval Units
	CLIENT PATIENT ID: ABHA NO :

	BIOCHEMISTRY		
MEDI WHEEL FULL BODY HEALTH CHEC	K UP ABOVE 40 MALE		
GLUCOSE FASTING, FLUORIDE PLASMA			
FBS (FASTING BLOOD SUGAR)	117 High	Normal <100	mg/dL
		Impaired fasting glue	cose:100 to
		125	
		Diabetes mellitus: >	= 126 (on
		more than 1 occassio	on)



113

PPBS(POST PRANDIAL BLOOD SUGAR)

mg/dL Normal <140 Impaired glucose tolerance:140 to 199 Diabetes mellitus : > = 200 (on more than 1 occassion) ADA guideline 2021

(ADA guidelines 2021)

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Page 8 Of 17

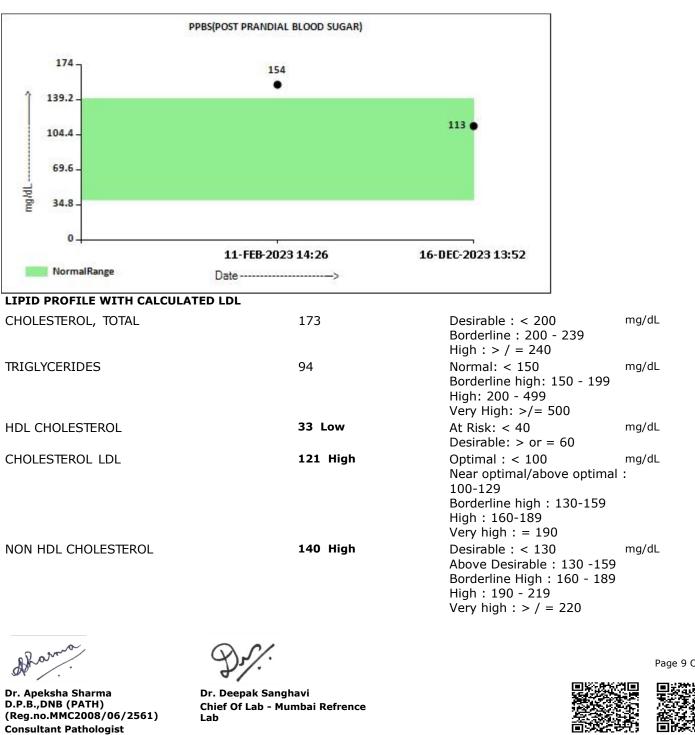








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Page 9 Of 17



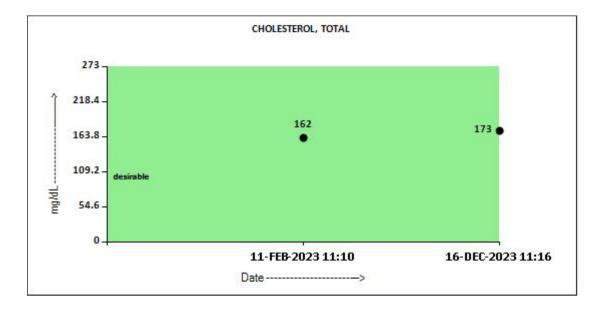






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	ACCESSION NO : 0002WL0	24266	AGE/SEX	:46 Years Male
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Test Report Status <u>Preliminary</u>	Results	Biological	Reference	Interval Units
VERY LOW DENSITY LIPOPROTEIN	19.0	< or = 30	.0	mg/dL
CHOL/HDL RATIO	5.2 High	Low Risk	: 3.3 - 4.4	
		5	Risk : 4.5 -	
		Moderate High Risk	Risk : 7.1	- 11.0
LDL/HDL RATIO	3.7 High	-	Low Risk :	05-30
				Risk : 3.1
		- 6.0	.,	

High Risk : > 6.0



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Page 10 Of 17





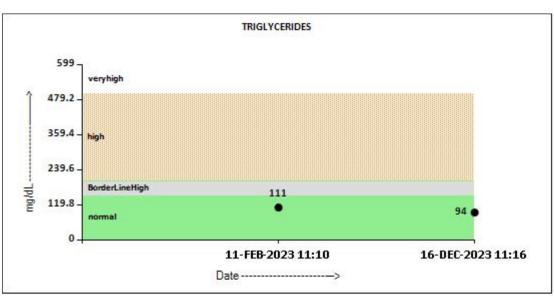
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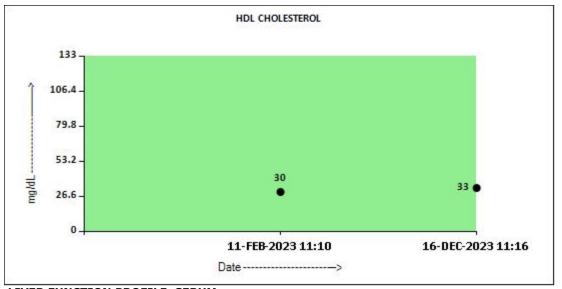






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Page 11 Of 17



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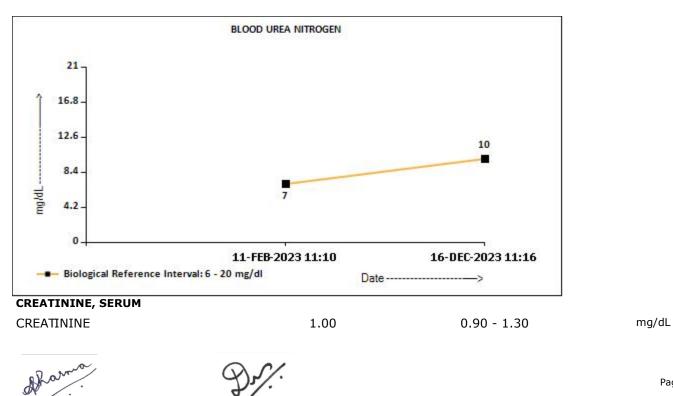






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Test Report Status <u>Preliminary</u>	Results	Biological Reference Int	erval Units
BILIRUBIN, TOTAL	0.66	Upto 1.2	mg/dL
BILIRUBIN, DIRECT	0.23	< or = 0.3	mg/dL
BILIRUBIN, INDIRECT	0.43	0.0 - 0.9	mg/dL
TOTAL PROTEIN	7.8	6.0 - 8.0	g/dL
ALBUMIN	4.7	3.97 - 4.94	g/dL
GLOBULIN	3.1	2.0 - 3.5	g/dL
ALBUMIN/GLOBULIN RATIO	1.5	1.0 - 2.1	RATIO
ASPARTATE AMINOTRANSFERASE(AST/SGOT)	20	Upto 40	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT)	28	Upto 41	U/L
ALKALINE PHOSPHATASE	69	40 - 129	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT)	19	< 60	U/L
LACTATE DEHYDROGENASE	172	< 232	U/L
BLOOD UREA NITROGEN (BUN), SERUM			
BLOOD UREA NITROGEN	10	6 - 20	mg/dL



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Page 12 Of 17



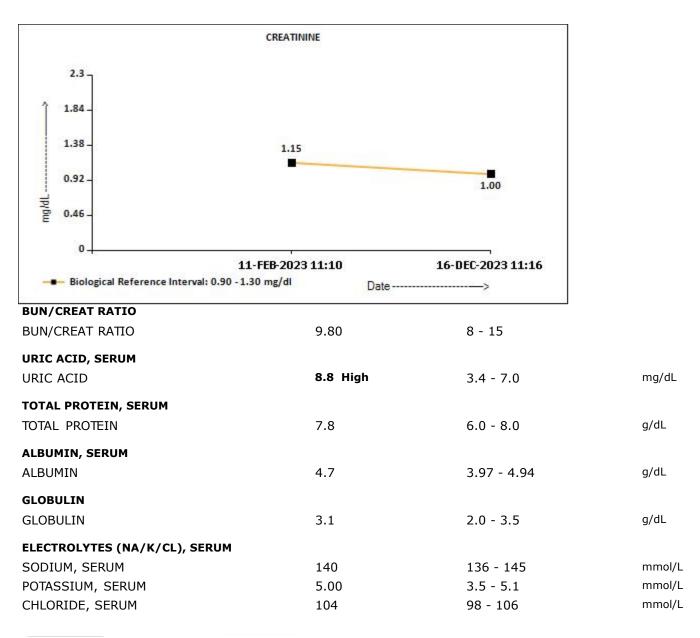
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Page 13 Of 17









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Test Report Status	<u>Preliminary</u>	Results	Biol	ogical I	Reference	e Interval 🛛 🛛	Jnits

Interpretation(s)

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 sothat 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 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-

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, (indirect) bilirubin in Viral hepatitis), and abnormal bilirubin metabolism (eg, hereditary and neonatal jaundice). Conjugated (direct) bilirubin is elevated more than unconjugated (indirect) bilirubin is also elevated more than unconjugated (indirect) bilirubin is also elevated more than unconjugated (indirect) bilirubin 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

BLODD 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.



Dr. Apeksha Sharma D.P.B., DNB (PATH) (Reg.no.MMC2008/06/2561) **Consultant Pathologist**

Page 14 Of 17





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Lab

Dr. Deepak Sanghavi

Chief Of Lab - Mumbai Refrence





PATIENT NAME : RA	JSHEKAR PABBA	REF. DOCTOR : SELF				
		ACCESSION NO	: 0002WL024266	AGE/SEX	:46 Years	Male
RAJSHEKAR PABBA		PATIENT ID	: RAJSM3110772	DRAWN	:16/12/2023	09:01:42
		CLIENT PATIENT	ID:	RECEIVED	: 16/12/2023	09:03:01
		ABHA NO	:	REPORTED	:16/12/2023	15:55:32
Test Report Status	Preliminary	Results	Biological	Reference	e Interval L	Jnits

CLINICA	AL PATH - URINALYSIS					
MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE						
PHYSICAL EXAMINATION, URINE						
COLOR	PALE YELLOW					
APPEARANCE	SLIGHTLY HAZY					
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	NOT DETECTED	NOT DETECTED				
BILIRUBIN	NOT DETECTED	NOT DETECTED				
UROBILINOGEN	NOT DETECTED					
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)	2-3	0-5	/HPF			
EPITHELIAL CELLS	2-3	0-5	/HPF			
CASTS	NOT DETECTED					
CRYSTALS	NOT DETECTED					
BACTERIA	NOT DETECTED	NOT DETECTED				
YEAST	NOT DETECTED	NOT DETECTED				

Dr. Deepak Sanghavi Chief Of Lab - Mumbai Refrence Lab



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Page 15 Of 17



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PATIENT NAME : RAJSHEKAR PABBA	REF. DOCTOR : SELF			
	ACCESSION NO : 0002WL024266	AGE/SEX :46 Years Male		
RAJSHEKAR PABBA	PATIENT ID : RAJSM3110772	DRAWN :16/12/2023 09:01:42		
	CLIENT PATIENT ID:	RECEIVED : 16/12/2023 09:03:01		
	ABHA NO :	REPORTED :16/12/2023 15:55:32		
Test Report Status <u>Preliminary</u>	Results Biological	Reference Interval Units		

,			
CLINIC	AL PATH - STOOL ANALYS	SIS	
MEDI WHEEL FULL BODY HEALTH CHECK UP	ABOVE 40 MALE		
PHYSICAL EXAMINATION, STOOL			
COLOUR	BROWN		
CONSISTENCY	SEMI FORMED		
MUCUS	NOT DETECTED	NOT DETECTED	
VISIBLE BLOOD	ABSENT	ABSENT	
ADULT PARASITE	NOT DETECTED		
CHEMICAL EXAMINATION, STOOL			
STOOL PH	6.0		
OCCULT BLOOD	NOT DETECTED	NOT DETECTED	
MICROSCOPIC EXAMINATION, STOOL			
PUS CELLS	NOT DETECTED		/hpf
RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
CYSTS	NOT DETECTED	NOT DETECTED	
OVA	NOT DETECTED		
LARVAE	NOT DETECTED	NOT DETECTED	
TROPHOZOITES	NOT DETECTED	NOT DETECTED	
FAT	ABSENT		
CHARCOT LEYDEN CRYSTALS	ABSENT		



Dr. Ekta Patil,MD Microbiologist

Page 16 Of 17



Vie<u>w Details</u>







PATIENT NAME : RAJSHEKAR PABBA	REF. DOCTOR : SELF		
DAISHEKAD DABBA	ACCESSION NO : 0002WL024266	AGE/SEX : 46 Years Male	
	PATIENT ID : RAJSM3110772	DRAWN :16/12/2023 09:01:42	
	CLIENT PATIENT ID:	RECEIVED : 16/12/2023 09:03:01	
	ABHA NO :	REPORTED :16/12/2023 15:55:32	
Test Report Status Preliminary	Results Biological	Reference Interval Units	

SPECIALISED CHEMISTRY - HORMONE				
MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE				
THYROID PANEL, SERUM				
ТЗ	134.0	80.0 - 200.0	ng/dL	
T4	8.44	5.10 - 14.10	µg/dL	
TSH (ULTRASENSITIVE)	2.050	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.

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.

Agilus Diagnostics Ltd

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Page 17 Of 17

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