

<b>PATIENT NAME : SANCHIT RAINA</b>		<b>REF. DOCTOR : SELF</b>	
<b>CODE/NAME &amp; ADDRESS</b> : C000138402 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST DELHI NEW DELHI 110030 8800465156	<b>ACCESSION NO</b> : <b>0202XA006875</b>	<b>AGE/SEX</b> : 33 Years	<b>Male</b>
	<b>PATIENT ID</b> : SANCM150191121	<b>DRAWN</b> :	
	<b>CLIENT PATIENT ID</b> :	<b>RECEIVED</b> : 27/01/2024 15:37:22	
	<b>ABHA NO</b> :	<b>REPORTED</b> : 02/02/2024 09:12:41	

Test Report Status	Preliminary	Results	Biological Reference Interval	Units
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**MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE**

**XRAY-CHEST**

>>>	BOTH THE LUNG FIELDS ARE CLEAR
>>>	BOTH THE COSTOPHRENIC AND CARIOPHRENIC ANGELS ARE CLEAR
>>>	BOTH THE HILA ARE NORMAL
>>>	CARDIAC AND AORTIC SHADOWS APPEAR NORMAL
>>>	BOTH THE DOMES OF THE DIAPHRAM ARE NORMAL
>>>	VISUALIZED BONY THORAX IS NORMAL
<b>IMPRESSION</b>	<b>NO ABNORMALITY DETECTED</b>

**ECG**

ECG WITHIN NORMAL LIMITS

**MEDICAL HISTORY**

RELEVANT PRESENT HISTORY	NOT SIGNIFICANT
RELEVANT PAST HISTORY	NO PAST H/O HT,TB,BA,DM,IHD,EPILEPSY NO SURGERY IN THE PAST
RELEVANT PERSONAL HISTORY	MARRIED,1 ISSUE,VEGETARIAN,NO SMOKING,NO ALCOHOL
RELEVANT FAMILY HISTORY	H/O MOTHER HTN FATHER DM
OCCUPATIONAL HISTORY	PVT.JOB

**ANTHROPOMETRIC DATA & BMI**

HEIGHT IN METERS	1.71	mts
WEIGHT IN KGS.	86	Kgs
BMI	29	kg/sqmts

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

**Dr.Hitesh Marwaha**  
Locum Pathologist



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 Tel : 9111591115, Fax : CIN - U74899PB1995PLC045956



**Patient Ref. No. 77500006214280**

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**GENERAL EXAMINATION**

GENERAL APPEARANCE / NUTRITIONAL STATUS	OVERWEIGHT
SKIN	NORMAL
UPPER LIMB	NORMAL
LOWER LIMB	NORMAL
TEMPERATURE	NORMAL
PULSE	74/MIN
RESPIRATORY RATE	16/MIN

**CARDIOVASCULAR SYSTEM**

BP	120/80	mm/Hg
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**BASIC EYE EXAMINATION**

DISTANT VISION RIGHT EYE WITH GLASSES	WITH GLASSES NORMAL
DISTANT VISION LEFT EYE WITH GLASSES	WITH GLASSES NORMAL
NEAR VISION RIGHT EYE WITHOUT GLASSES	N/6
NEAR VISION LEFT EYE WITHOUT GLASSES	N/6
COLOUR VISION	NORMAL (OUT OF 17 NUMBERED PLATES)

**BASIC DENTAL EXAMINATION**

TEETH	NORMAL
GUMS	HEALTHY



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**SUMMARY**

RELEVANT HISTORY	NOT SIGNIFICANT
RELEVANT GP EXAMINATION FINDINGS	NOT SIGNIFICANT
RELEVANT LAB INVESTIGATIONS	hard copy attached
RELEVANT NON PATHOLOGY DIAGNOSTICS	ECG:-WNL,CHEST X-RAY:-WNL,GP DONE
REMARKS / RECOMMENDATIONS	NONE

**FITNESS STATUS**

FITNESS STATUS	FIT (AS PER REQUESTED PANEL OF TESTS)
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**Comments**

OUR PANEL OF DOCTORS:

CONSULTANT CARDIOLOGIST AND PHYSICIAN - DR.PARMINDER SINGH RANA  
M.B.B.S., MD (Regd.No.22878)

CONSULTANT RADIOLOGIST - DR. AMRITA RANA - (M.D. RADIODIAGNOSIS)  
(Regd .No.24590)

OUR WELLNESS INVESTIGATIONS HAVE BEEN PERFORMED BY OUR PANEL OF DOCTORS;  
HOWEVER THIS REPORT CARRIES THE SIGNATURES OF OUR LAB MEDICINE DOCTORS  
WHICH IS AN INVIOABLE FEATURE OF OUR LABORATORY MANAGEMENT SOFTWARE



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**MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 YEARS** RESULT PENDING

**ULTRASOUND ABDOMEN** RESULT PENDING

**TMT OR ECHO**

**CLINICAL PROFILE**

Test pending

<b>Interpretation(s)</b>  
MEDICAL HISTORY-

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

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

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

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## HAEMATOLOGY - CBC

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

## BLOOD COUNTS, EDTA WHOLE BLOOD

HEMOGLOBIN (HB)	13.2	13.0 - 17.0	g/dL
RED BLOOD CELL (RBC) COUNT	<b>3.29 Low</b>	4.5 - 5.5	mil/ $\mu$ L
WHITE BLOOD CELL (WBC) COUNT	4.90	4.0 - 10.0	thou/ $\mu$ L
PLATELET COUNT	151	150 - 410	thou/ $\mu$ L

## RBC AND PLATELET INDICES

HEMATOCRIT (PCV)	<b>39.6 Low</b>	40 - 50	%
MEAN CORPUSCULAR VOLUME (MCV)	<b>120.0 High</b>	83 - 101	fL
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	<b>40.1 High</b>	27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC)	33.3	31.5 - 34.5	g/dL
RED CELL DISTRIBUTION WIDTH (RDW)	12.9	11.6 - 14.0	%
MENTZER INDEX	36.5		
MEAN PLATELET VOLUME (MPV)	<b>11.7 High</b>	6.8 - 10.9	fL

## WBC DIFFERENTIAL COUNT

NEUTROPHILS	48	40 - 80	%
LYMPHOCYTES	39	20 - 40	%
MONOCYTES	6	2 - 10	%
EOSINOPHILS	<b>7 High</b>	1 - 6	%
BASOPHILS	0	0 - 2	%
ABSOLUTE NEUTROPHIL COUNT	2.35	2.0 - 7.0	thou/ $\mu$ L
ABSOLUTE LYMPHOCYTE COUNT	1.91	1.0 - 3.0	thou/ $\mu$ L
ABSOLUTE MONOCYTE COUNT	0.29	0.2 - 1.0	thou/ $\mu$ L
ABSOLUTE EOSINOPHIL COUNT	0.34	0.02 - 0.50	thou/ $\mu$ L
ABSOLUTE BASOPHIL COUNT	<b>0.00 Low</b>	0.02 - 0.10	thou/ $\mu$ L



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**AGE/SEX : 33 Years Male**

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NEUTROPHIL LYMPHOCYTE RATIO (NLR) 1.2

<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 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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## HAEMATOLOGY

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

## ERYTHROCYTE SEDIMENTATION RATE (ESR), EDTA BLOOD

E.S.R	02	0 - 14	mm at 1 hr
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## GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD

HBA1C	5.0	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 Therapeutic goals: < 7.0 Action suggested : > 8.0 (ADA Guideline 2021)	%
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ESTIMATED AVERAGE GLUCOSE(EAG)	96.8	< 116.0	mg/dL
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## &lt;b&gt;Interpretation(s)&lt;/b&gt;

ERYTHROCYTE SEDIMENTATION RATE (ESR), EDTA BLOOD- &lt;b&gt;TEST DESCRIPTION&lt;/b&gt; :-

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.

## &lt;b&gt;TEST INTERPRETATION&lt;/b&gt;

<b>Increase</b> in: Infections, Vasculitides, 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 (Paraproteinemia, 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: Polycythemia vera, Sickle cell anemia

## &lt;b&gt;LIMITATIONS&lt;/b&gt;

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

<b>False Decreased</b> : Poikilocytosis, (Sickle Cells, spherocytes), Microcytosis, Low fibrinogen, Very high WBC counts, Drugs (Quinine,

salicylates)

## REFERENCE :

1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition 2. Paediatric reference intervals. AACC Press, 7th edition. Edited by S. Soldin 3. The reference



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for the adult reference range is "Practical Haematology by Dacie and Lewis,10th edition.  
 GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD-<b>Used For</b>:"

- Evaluating the long-term control of blood glucose concentrations in diabetic patients.
  - Diagnosing diabetes.
  - 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.
- 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.
  - Vitamin C & E are reported to falsely lower test results.(possibly by inhibiting glycation of hemoglobin.
  - 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
    - Homozygous hemoglobinopathy. Fructosamine is recommended for testing of HbA1c.
    - Heterozygous state detected (D10 is corrected for HbS & HbC trait.)
    - 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 BELOW 40 MALE

## ABO GROUP &amp; RH TYPE, EDTA WHOLE BLOOD

ABO GROUP	TYPE O
RH TYPE	POSITIVE

## Comments

FALSE NEGATIVE RH TYPING COULD BE DUE INHERITED CHARACTERISTIC IN SOME. TWO GRADES, HIGH GRADE DU AND LOW GRADE DU. FORMER AGGLUTINATED BY CERTAIN ANTISERA AND LOW GRADE DU ARE MOSTLY DETECTED BY AHG TEST. FALSE POSITIVE RH TYPING CAN OCCUR DUE TO SEVERAL REASONS INCLUDING SPONTANEOUS AGGLUTINATION OF RED CELLS WITH POSITIVE DAT, ROULEAUX FORMATION DUE TO PRESENCE OF COLD AUTOAGGLUTININS/ABNORMAL PROTEINS IN PATIENTS SERA, REAGENT CONTAMINATION, ANTIBODY TO LOW INCIDENCE ANTIGENS AND HUMAN ERROR. IN CASE OF ANY DISCREPANCY, REQUESTED TO REPORT TO THE LAB FOR FURTHER ACTION.

## &lt;b&gt;Interpretation(s)&lt;/b&gt;

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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**BIOCHEMISTRY**

**MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE**

**GLUCOSE FASTING, FLUORIDE PLASMA**

FBS (FASTING BLOOD SUGAR)	80	Normal : < 100 Pre-diabetes: 100-125 Diabetes: >/=126	mg/dL
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**LIPID PROFILE WITH CALCULATED LDL**

CHOLESTEROL, TOTAL	173	< 200 Desirable 200 - 239 Borderline High >/= 240 High	mg/dL
TRIGLYCERIDES	71	< 150 Normal 150 - 199 Borderline High 200 - 499 High >/=500 Very High	mg/dL
HDL CHOLESTEROL	<b>37 Low</b>	< 40 Low >/=60 High	mg/dL
CHOLESTEROL LDL	<b>122 High</b>	< 100 Optimal 100 - 129 Near optimal/ above optimal 130 - 159 Borderline High 160 - 189 High >/= 190 Very High	mg/dL
NON HDL CHOLESTEROL	<b>136 High</b>	Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL
VERY LOW DENSITY LIPOPROTEIN CHOL/HDL RATIO	14.2 <b>4.7 High</b>	</= 30.0 3.3 - 4.4 Low Risk 4.5 - 7.0 Average Risk 7.1 - 11.0 Moderate Risk > 11.0	mg/dL



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 Locum Pathologist



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 Amritsar, 143001  
 Punjab, India  
 Tel : 9111591115, Fax : CIN - U74899PB1995PLC045956



**Patient Ref. No. 77500006214280**

**PATIENT NAME : SANCHIT RAINA**

**REF. DOCTOR : SELF**

**CODE/NAME & ADDRESS :** C000138402  
 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL  
 F-703, F-703, LADO SARAI, MEHRAULISOUTH  
 WEST DELHI  
 NEW DELHI 110030  
 8800465156

**ACCESSION NO :** **0202XA006875**  
**PATIENT ID :** SANCM150191121  
**CLIENT PATIENT ID:**  
**ABHA NO :**

**AGE/SEX :** 33 Years Male  
**DRAWN :**  
**RECEIVED :** 27/01/2024 15:37:22  
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LDL/HDL RATIO	<b>3.3 High</b>	High Risk 0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	
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**LIVER FUNCTION PROFILE, SERUM**

BILIRUBIN, TOTAL	0.90	0.2 - 1.0	mg/dL
BILIRUBIN, DIRECT	0.15	0.0 - 0.2	mg/dL
BILIRUBIN, INDIRECT	0.75	0.1 - 1.0	mg/dL
TOTAL PROTEIN	7.6	6.4 - 8.2	g/dL
ALBUMIN	4.1	3.4 - 5.0	g/dL
GLOBULIN	3.5	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO	1.2	1.0 - 2.1	RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	22	15 - 37	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT)	<b>53 High</b>	< 45.0	U/L
ALKALINE PHOSPHATASE	89	30 - 120	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT)	31	15 - 85	U/L
LACTATE DEHYDROGENASE	161	85 - 227	U/L

**BLOOD UREA NITROGEN (BUN), SERUM**

BLOOD UREA NITROGEN	6	6 - 20	mg/dL
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**CREATININE, SERUM**

CREATININE	<b>0.66 Low</b>	0.90 - 1.30	mg/dL
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**BUN/CREAT RATIO**

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BUN/CREAT RATIO		9.09	5.00 - 15.00	
<b>URIC ACID, SERUM</b>				
URIC ACID		4.0	3.5 - 7.2	mg/dL
<b>TOTAL PROTEIN, SERUM</b>				
TOTAL PROTEIN		7.6	6.4 - 8.2	g/dL
<b>ALBUMIN, SERUM</b>				
ALBUMIN		4.1	3.4 - 5.0	g/dL
<b>GLOBULIN</b>				
GLOBULIN		3.5	2.0 - 4.1	g/dL
<b>ELECTROLYTES (NA/K/CL), SERUM</b>				
SODIUM, SERUM		137	136 - 145	mmol/L
POTASSIUM, SERUM		4.91	3.50 - 5.10	mmol/L
CHLORIDE, SERUM		104	98 - 107	mmol/L

**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

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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.  
**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.

**Dr.Hitesh Marwaha**  
Locum Pathologist



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**CLINICAL PATH - URINALYSIS**

**MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE**

**PHYSICAL EXAMINATION, URINE**

COLOR	PALE YELLOW
APPEARANCE	CLEAR

**CHEMICAL EXAMINATION, URINE**

PH	6.5	4.5 - 7.5
SPECIFIC GRAVITY	1.010	1.005 - 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	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)	2-3	0-5	/HPF
EPITHELIAL CELLS	NOT DETECTED	0-5	/HPF
CASTS	NOT DETECTED		
CRYSTALS	CALCIUM OXALATE DETECTED.		
BACTERIA	NOT DETECTED	NOT DETECTED	
YEAST	NOT DETECTED	NOT DETECTED	

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	<b>PATIENT ID :</b> SANCM150191121	<b>DRAWN :</b>	
	<b>CLIENT PATIENT ID:</b>	<b>RECEIVED :</b> 27/01/2024 15:37:22	
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**Comments**

URINE MICROSCOPIC EXAMINATION PERFORMED ON DEPOSIT AFTER CENTRIFUGATION.



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**CLINICAL PATH - STOOL ANALYSIS**

**MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE**

**MICROSCOPIC EXAMINATION,STOOL**

REMARK

TEST CANCELLED AS SPECIMEN NOT RECEIVED

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## SPECIALISED CHEMISTRY - HORMONE

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

## THYROID PANEL, SERUM

T3	108.04	60.0 - 181.0	ng/dL
T4	5.90	4.5 - 10.9	µg/dL
TSH (ULTRASENSITIVE)	<b>25.286 High</b>	0.550 - 4.780	µIU/mL

\*\*End Of Report\*\*

Please visit [www.agilusdiagnostics.com](http://www.agilusdiagnostics.com) for related Test Information for this accession

## CONDITIONS OF LABORATORY TESTING &amp; 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

Fortis Hospital, Sector 62, Phase VIII,  
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