



Patient Ref. No. 251000000166199



CLIENT CODE : C000049066

Cert. No. MC-5333

CLIENT'S NAME AND ADDRESS :

SRL JAIPUR WELLNESS CORPORATE WALK IN (CASH)
AAKRITI LABS PVT LTD. A-430, AGRASEN MARG

SRL Ltd

C/o Aakriti Labs Pvt Ltd, 3, Mahatma Gandhi Marg, Gandhi Nagar Mod,
Tonk Road
JAIPUR, 302015
Rajasthan, INDIAJAIPUR 302017
RAJASTHAN INDIA
9314660100

PATIENT NAME : TARUN MODI

PATIENT ID : TARUM261191251

ACCESSION NO : 0251VK002455 AGE : 31 Years SEX : Male

ABHA NO :

DRAWN : 26/11/2022 08:55:00

RECEIVED : 26/11/2022 11:36:25

REPORTED : 26/11/2022 16:02:14

REFERRING DOCTOR : SELF

CLIENT PATIENT ID : 012211260014

Test Report Status	Final	Results	Biological Reference Interval	Units
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MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE**BLOOD COUNTS, EDTA WHOLE BLOOD**

HEMOGLOBIN (HB)	14.2	13.0 - 17.0	g/dL
METHOD : CYANIDE FREE DETERMINATION			
RED BLOOD CELL (RBC) COUNT	5.18	4.5 - 5.5	mil/ μ L
METHOD : ELECTRICAL IMPEDANCE			
WHITE BLOOD CELL (WBC) COUNT	5.10	4.0 - 10.0	thou/ μ L
METHOD : ELECTRICAL IMPEDANCE			
PLATELET COUNT	205	150 - 410	thou/ μ L
METHOD : ELECTRONIC IMPEDANCE			

RBC AND PLATELET INDICES

HEMATOCRIT (PCV)	43.8	40 - 50	%
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR VOLUME (MCV)	85.0	83 - 101	fL
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	27.4	27.0 - 32.0	pg
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC)	32.4	31.5 - 34.5	g/dL
METHOD : CALCULATED PARAMETER			
RED CELL DISTRIBUTION WIDTH (RDW)	14.3	High 11.6 - 14.0	%
METHOD : CALCULATED PARAMETER			
MENTZER INDEX	16.4		
MEAN PLATELET VOLUME (MPV)	12.4	High 6.8 - 10.9	fL
METHOD : CALCULATED PARAMETER			

WBC DIFFERENTIAL COUNT

NEUTROPHILS	54	40 - 80	%
METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY			
LYMPHOCYTES	36	20 - 40	%
METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY			
MONOCYTES	06	2 - 10	%
METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY			
EOSINOPHILS	04	1 - 6	%
METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY			
BASOPHILS	00	0 - 2	%
METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY			



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ABSOLUTE NEUTROPHIL COUNT	2.75	2.0 - 7.0	thou/ μ L
METHOD : CALCULATED PARAMETER			
ABSOLUTE LYMPHOCYTE COUNT	1.84	1.0 - 3.0	thou/ μ L
METHOD : CALCULATED PARAMETER			
ABSOLUTE MONOCYTE COUNT	0.31	0.2 - 1.0	thou/ μ L
METHOD : CALCULATED PARAMETER			
ABSOLUTE EOSINOPHIL COUNT	0.20	0.02 - 0.50	thou/ μ L
METHOD : CALCULATED PARAMETER			
ABSOLUTE BASOPHIL COUNT	0	Low 0.02 - 0.10	thou/ μ L
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	1.5		
* ERYTHROCYTE SEDIMENTATION RATE (ESR), WHOLE BLOOD			
E.S.R	05	0 - 14	mm at 1 hr
METHOD : AUTOMATED (PHOTOMETRICAL CAPILLARY STOPPED FLOW KINETIC ANALYSIS)"			
GLUCOSE FASTING, FLUORIDE PLASMA			
FBS (FASTING BLOOD SUGAR)	86	74 - 99	mg/dL
METHOD : GLUCOSE OXIDASE			
GLYCOSYLATED HEMOGLOBIN (HBA1C), EDTA WHOLE BLOOD			
HBA1C	5.3	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 ADA Target: 7.0 Action suggested: > 8.0	%
METHOD : HIGH PERFORMANCE LIQUID CHROMATOGRAPHY (HPLC)			
ESTIMATED AVERAGE GLUCOSE (EAG)	105.4	< 116.0	mg/dL
METHOD : CALCULATED PARAMETER			
GLUCOSE, POST-PRANDIAL, PLASMA			
PPBS (POST PRANDIAL BLOOD SUGAR)	108	70 - 140	mg/dL
METHOD : GLUCOSE OXIDASE			
LIPID PROFILE, SERUM			
CHOLESTEROL, TOTAL	200	< 200 Desirable 200 - 239 Borderline High >= 240 High	mg/dL
METHOD : CHOLESTEROL OXIDASE			
TRIGLYCERIDES	164	High < 150 Normal 150 - 199 Borderline High 200 - 499 High >= 500 Very High	mg/dL



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METHOD : LIPASE/GPO-PAP NO CORRECTION

HDL CHOLESTEROL	40	< 40 Low >/=60 High	mg/dL
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METHOD : DIRECT CLEARANCE METHOD

CHOLESTEROL LDL	127	High < 100 Optimal 100 - 129 Near optimal/ above optimal 130 - 159 Borderline High 160 - 189 High >/= 190 Very High	mg/dL
NON HDL CHOLESTEROL	160	High Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL

METHOD : CALCULATED PARAMETER

CHOL/HDL RATIO	5.0	High 3,3 - 4,4 Low Risk 4,5 - 7,0 Average Risk 7,1 - 11,0 Moderate Risk > 11,0 High Risk	
LDL/HDL RATIO	3.2	High 0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	

VERY LOW DENSITY LIPOPROTEIN	32.8	High </= 30.0	mg/dL
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LIVER FUNCTION PROFILE, SERUM

BILIRUBIN, TOTAL	0.81	0 - 1	mg/dL
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METHOD : DIAZO WITH SULPHANILIC ACID

BILIRUBIN, DIRECT	0.21	0.00 - 0.25	mg/dL
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METHOD : DIAZO WITH SULPHANILIC ACID

BILIRUBIN, INDIRECT	0.60	0.1 - 1.0	mg/dL
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METHOD : CALCULATED PARAMETER

TOTAL PROTEIN	8.2	6.4 - 8.2	g/dL
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METHOD : BIURET REACTION, END POINT

ALBUMIN	4.9	High 3.8 - 4.4	g/dL
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METHOD : BROMOCRESOL GREEN

GLOBULIN	3.3	2.0 - 4.1	g/dL
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METHOD : CALCULATED PARAMETER



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ALBUMIN/GLOBULIN RATIO	1.5	1.0 - 2.1	RATIO
METHOD : CALCULATED PARAMETER			
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	40	High 0 - 37	U/L
METHOD : TRIS BUFFER NO P5P IFCC / SFBC 37° C			
ALANINE AMINOTRANSFERASE (ALT/SGPT)	51	High 0 - 40	U/L
METHOD : TRIS BUFFER NO P5P IFCC / SFBC 37° C			
ALKALINE PHOSPHATASE	104	39 - 117	U/L
METHOD : AMP OPTIMISED TO IFCC 37° C			
GAMMA GLUTAMYL TRANSFERASE (GGT)	32	11 - 50	U/L
METHOD : GAMMA GLUTAMYL-3 CARBOXY-4 NITROANILIDE (IFCC) 37° C			
LACTATE DEHYDROGENASE	392	230 - 460	U/L
METHOD : GERMAN METHODS 37° C			
BLOOD UREA NITROGEN (BUN), SERUM			
BLOOD UREA NITROGEN	13	5.0 - 18.0	mg/dL
METHOD : UREASE KINETIC			
CREATININE, SERUM			
CREATININE	0.91	0.8 - 1.3	mg/dL
METHOD : ALKALINE PICRATE NO DEPROTEINIZATION			
BUN/CREAT RATIO			
BUN/CREAT RATIO	14.29		
METHOD : CALCULATED PARAMETER			
URIC ACID, SERUM			
URIC ACID	6.9	3.4 - 7.0	mg/dL
METHOD : URICASE PEROXIDASE WITH ASCORBATE OXIDASE			
TOTAL PROTEIN, SERUM			
TOTAL PROTEIN	8.2	6.4 - 8.3	g/dL
METHOD : BIURET REACTION, END POINT			
ALBUMIN, SERUM			
ALBUMIN	4.9	High 3.8 - 4.4	g/dL
METHOD : BROMOCRESOL GREEN			
GLOBULIN			
GLOBULIN	3.3	2.0 - 4.1	g/dL
METHOD : CALCULATED PARAMETER			
ELECTROLYTES (NA/K/CL), SERUM			
SODIUM, SERUM	139.3	137 - 145	mmol/L



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METHOD : ION-SELECTIVE ELECTRODE

POTASSIUM, SERUM

4.13

3.6 - 5.0

mmol/L

METHOD : ION-SELECTIVE ELECTRODE

CHLORIDE, SERUM

101.5

98 - 107

mmol/L

METHOD : ION-SELECTIVE ELECTRODE

Interpretation(s)**PHYSICAL EXAMINATION, URINE**

COLOR

PALE YELLOW

METHOD : GROSS EXAMINATION

APPEARANCE

CLEAR

METHOD : GROSS EXAMINATION

CHEMICAL EXAMINATION, URINE

PH

6.5

4.7 - 7.5

METHOD : DOUBLE INDICATOR PRINCIPLE

SPECIFIC GRAVITY

1.005

1.003 - 1.035

METHOD : IONIC CONCENTRATION METHOD

PROTEIN

NOT DETECTED

NOT DETECTED

METHOD : PROTEIN ERROR OF INDICATORS WITH REFLECTANCE

GLUCOSE

NOT DETECTED

NOT DETECTED

METHOD : GLUCOSE OXIDASE PEROXIDASE / BENEDICTS

KETONES

NOT DETECTED

NOT DETECTED

METHOD : SODIUM NITROPRUSSIDE REACTION

BLOOD

NOT DETECTED

NOT DETECTED

METHOD : PEROXIDASE ANTI PEROXIDASE

BILIRUBIN

NOT DETECTED

NOT DETECTED

METHOD : DIPSTICK

UROBILINOGEN

NORMAL

NORMAL

METHOD : EHRlich REACTION REFLECTANCE

NITRITE

NOT DETECTED

NOT DETECTED

METHOD : NITRATE TO NITRITE CONVERSION METHOD

LEUKOCYTE ESTERASE

NOT DETECTED

NOT DETECTED

MICROSCOPIC EXAMINATION, URINE

RED BLOOD CELLS

NOT DETECTED

NOT DETECTED

/HPF

METHOD : MICROSCOPIC EXAMINATION

PUS CELL (WBC'S)

1-2

0-5

/HPF



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METHOD : DIPSTICK, MICROSCOPY

EPITHELIAL CELLS

0-1

0-5

/HPF

METHOD : MICROSCOPIC EXAMINATION

CASTS

NOT DETECTED

METHOD : MICROSCOPIC EXAMINATION

CRYSTALS

NOT DETECTED

METHOD : MICROSCOPIC EXAMINATION

BACTERIA

NOT DETECTED

NOT DETECTED

METHOD : MICROSCOPIC EXAMINATION

YEAST

NOT DETECTED

NOT DETECTED

Interpretation(s)**THYROID PANEL, SERUM**

T3

145.8

60.0 - 181.0

ng/dL

METHOD : CHEMILUMINESCENCE

T4

9.10

4.5 - 10.9

µg/dL

METHOD : CHEMILUMINESCENCE

TSH (ULTRASENSITIVE)

3.381

0,550 - 4,780

µIU/mL

METHOD : CHEMILUMINESCENCE

Interpretation(s)**STOOL: OVA & PARASITE**

COLOUR

SAMPLE NOT RECEIVED

METHOD : GROSS EXAMINATION

Interpretation(s)*** ABO GROUP & RH TYPE, EDTA WHOLE BLOOD**

ABO GROUP

TYPE A

METHOD : TUBE AGGLUTINATION

RH TYPE

POSITIVE

METHOD : TUBE AGGLUTINATION

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.



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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.

ERYTHROCYTE SEDIMENTATION RATE (ESR),WHOLE 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, 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 (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: Polycythemia 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.

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 so that 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; sulfonyleureas,tolbutamide, and other oral hypoglycemic agents.

NOTE:

Hypoglycemia is defined as a glucose of < 50 mg/dL in men and < 40 mg/dL in women.

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 glycaemic 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.

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 patient's metabolic control has remained continuously within the target range.

1. eAG (Estimated average glucose) converts percentage HbA1c to mg/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 $eAG (mg/dl) = 28.7 * HbA1c - 46.7$

HbA1c Estimation can get affected due to :

I. 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.

II. Vitamin C & E are reported to falsely lower test results. (possibly by inhibiting glycation of hemoglobin).

III. 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.



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

GLUCOSE, POST-PRANDIAL, PLASMA-High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glycosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc. Additional test HbA1c

LIVER FUNCTION PROFILE, SERUM-

LIVER FUNCTION PROFILE

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 result 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, Paget's disease, Rickets, Sarcoidosis etc. Lower-than-normal ALP levels seen in Hypophosphatasia, Malnutrition, Protein deficiency, Wilson's 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. Serum 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, Waldenstrom's disease. Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage), Burns, Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic syndrome, Protein-losing enteropathy etc. 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

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
- Muscular dystrophy

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-

Serum 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, Waldenstrom's disease

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

ALBUMIN, SERUM-

Human serum albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by: Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance, malnutrition and wasting etc.

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD-

Blood group is identified by antigens and antibodies present in the blood. Antigens are protein molecules found on the surface of red blood cells. Antibodies are found in plasma. To determine blood group, red cells are mixed with different antibody solutions to give A, B, O or AB.



Scan to View Details



Scan to View Report



Patient Ref. No. 251000000166199



CLIENT CODE : C000049066

Cert. No. MC-5333

CLIENT'S NAME AND ADDRESS :

SRL JAIPUR WELLNESS CORPORATE WALK IN (CASH)
AAKRITI LABS PVT LTD. A-430, AGRASEN MARGSRL Ltd
C/o Aakriti Labs Pvt Ltd, 3, Mahatma Gandhi Marg, Gandhi Nagar Mod,
Tonk Road
JAIPUR, 302015
Rajasthan, INDIAJAIPUR 302017
RAJASTHAN INDIA
9314660100

PATIENT NAME : TARUN MODI

PATIENT ID : TARUM261191251

ACCESSION NO : 0251VK002455 AGE : 31 Years SEX : Male

ABHA NO :

DRAWN : 26/11/2022 08:55:00

RECEIVED : 26/11/2022 11:36:25

REPORTED : 26/11/2022 16:02:14

REFERRING DOCTOR : SELF

CLIENT PATIENT ID : 012211260014

Test Report Status	Final	Results	Biological Reference Interval	Units
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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.

****End Of Report****

Please visit www.srlworld.com for related Test Information for this accession
TEST MARKED WITH '*' ARE OUTSIDE THE NABL ACCREDITED SCOPE OF THE LABORATORY.

Dr. Akansha Jain
Consultant Pathologist

Dr. Abhishek Sharma
Consultant Microbiologist



Scan to View Details



Scan to View Report



Aakriti Labs

3 Mahatma Gandhi Marg, Gandhi Nagar Mod
Tonk Road, Jaipur (Raj.) Ph.: 0141-2710661
www.aakritilabs.com
CIN NO.: U85195RJ2004PTC019563



Name : Mr. TARUN MODI
Age/Gender: 31 Y 6 M 4 D/Male
Patient ID : 012211260014
BarcodeNo : 10068593
Referred By : Self

Registration No: 47084
Registered : 26/Nov/2022 08:55AM
Analysed : 26/Nov/2022 12:22PM
Reported : 26/Nov/2022 12:22PM
Panel : Medi Wheel (ArcoFemi
Healthcare Ltd)

DIGITAL X-RAY CHEST PA VIEW

Soft tissue shadow and bony cages are normal.

Trachea is central.

Bilateral lung field and both CP angle are clear.

Domes of diaphragm are normally placed.


Transverse diameter of heart appears with normal limits.

IMPRESSION:- NO OBVIOUS ABNORMALITY DETECTED.

*** End Of Report ***

Page 1 of 1




Dr. Neera Mehta
M.B.B.S., D.M.R.D.
RMCNO.005807/14853



Aakriti Labs

3 Mahatma Gandhi Marg, Gandhi Nagar Mod
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Name : **Mr. TARUN MODI**

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Analysed : 26/Nov/2022 11:21AM

Reported : 26/Nov/2022 11:21AM

Panel : Medi Wheel (ArcoFemi
Healthcare Ltd)

USG: WHOLE ABDOMEN (Male)

LIVER : Is normal in size and shape with **mild bright** echogenecity.
The IHBR and hepatic radicals are not dilated.
No evidence of focal echopoor/echorich lesion seen.
Portal vein diameter and common bile duct appear normal.

GALL BLADDER : Is normal in size, shape and echotexture. Walls are smooth and regular with normal thickness. There is no evidence of cholelithiasis.

PANCREAS : Is normal in size, shape and echotexture. Pancreatic duct is not dilated.

SPLEEN : Is normal in size, shape and echogenecity. Splenic hilum is not dilated.

KIDNEYS : Right Kidney:-Size:103 x 43 mm, Left Kidney:-Size:101 x 47 mm.
Bilateral Kidneys are normal in size, shape and echotexture,
corticomedullary differentiation is fair and ratio appears normal.
Pelvi calyceal system is normal. No evidence of hydronephrosis/ nephrolithiasis.

URINARY BLADDER : Bladder walls are smooth, regular and normal thickness.

BLADDER : No evidence of mass or stone in bladder lumen.

PROSTATE: Is normal in size, shape and echotexture,
measures: 34 x 27 x 26 mm, wt:12 gms.
Its capsule is intact and no evidence of focal lesion.

SPECIFIC : No evidence of retroperitoneal mass or free fluid seen in peritoneal cavity.
No evidence of lymphadenopathy or mass lesion in retroperitoneum.
Visualized bowel loop appear normal. Great vessels appear normal.

IMPRESSION :- Mild fatty liver

*** End Of Report ***

Dr. NEERA MEHTA
MBBS, DMRC
RMC No. 00580714000

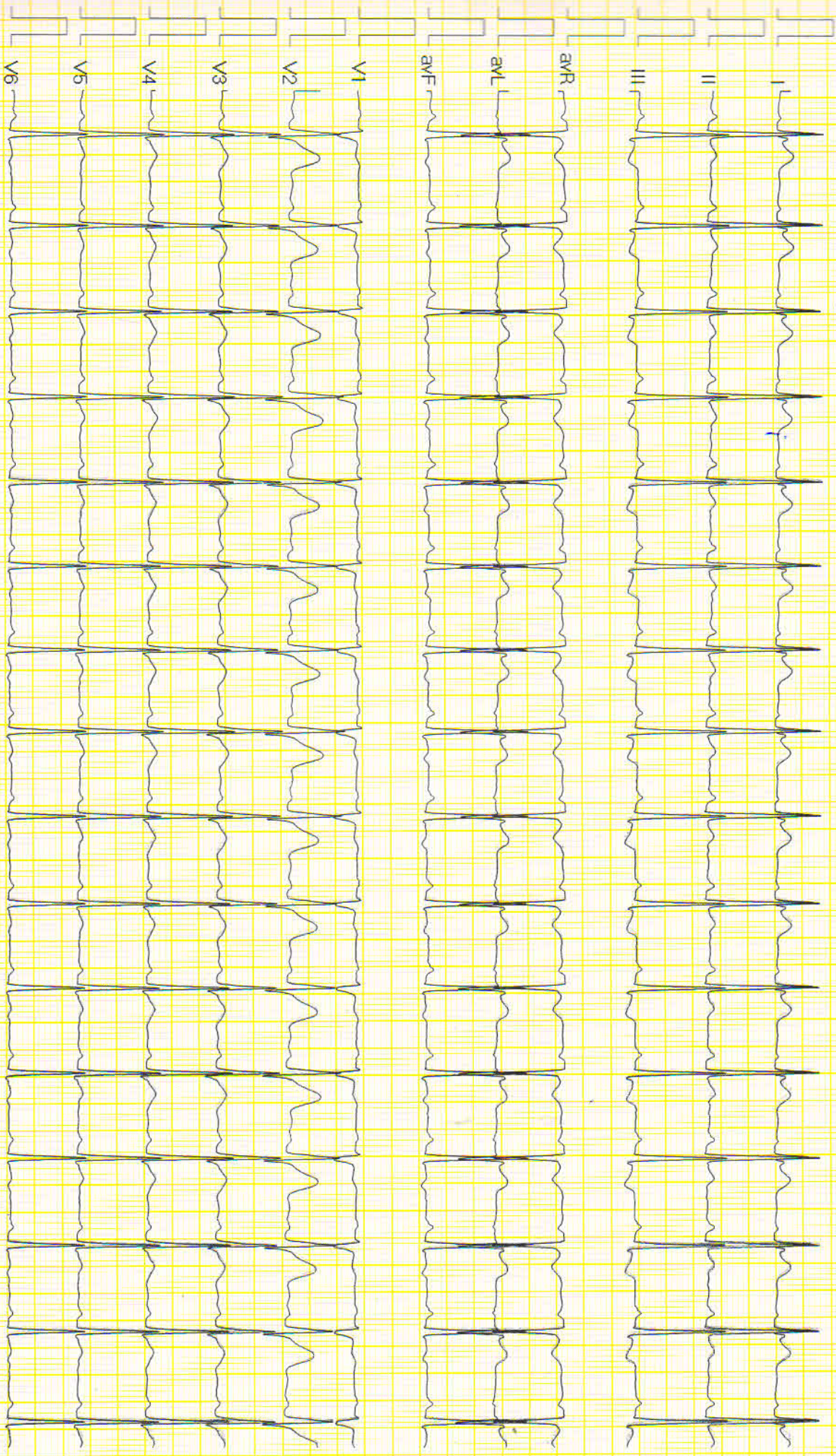
Page 1 of 1





MR. TARUN MODI / 31 Yrs / M / 10 cms / 10 Kg / HR 90

Date: 26 / 11 / 2022 BP: 125/85 mm/Hg BLC On Notch On HF 0.05 Hz LF 100 Hz



ST-T changes

KD

Dr. NITIZ GOYAL
M.B.B.S., M.D.
RMC - 023319



TARUN MODI / 31 Yrs / M / 0 Cms / 0 Kg
 Date: 26 / 11 / 2022

Stage	Time	Duration	Speed(mph)	Elevation	METS	Rate	%THR	BP	RPP	PVC	Comments
Supine	00:03	0:03	00.0	00.0	01.0	096	51%	125/85	120	00	
Standing	00:06	0:03	00.0	00.0	01.0	096	51%	125/85	120	00	
HV	00:19	0:13	00.0	00.0	01.0	096	51%	125/85	120	00	
Warm Up	00:33	0:14	00.0	00.0	01.0	101	53%	125/85	126	00	
ExStart	00:37	0:04	00.0	00.0	01.0	101	53%	125/85	126	00	
BRUCE Stage 1	03:37	3:00	01.7	10.0	04.7	126	67%	125/85	157	00	
BRUCE Stage 2	06:37	3:00	02.5	12.0	07.1	162	86%	125/85	202	00	
PeakEx	07:36	0:59	03.4	14.0	08.1	172	91%	125/85	215	00	
Recovery	08:36	1:00	00.0	00.0	01.2	143	76%	125/85	178	00	
Recovery	09:36	2:00	00.0	00.0	01.0	132	70%	155/80	204	00	
Recovery	10:40	3:04	00.0	00.0	01.0	119	63%	140/70	166	00	

REPORT :

Positive

FINAL IMPRESSION - TEST IS NEGATIVE FOR INDUCIBLE ISCHAEMIA

DR. NITIZ GOYAL
 M.B.B.S., M.D.
 DR. NITIZ GOYAL
 RMC - 023319