wellness partner

Aakriti Labs

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

Name	: MR. KHUSI RAM	Age/Sex	: 32Yrs/ MALE
Ref.By	: AAKRITI LABS	Date	: 28 January 2023

RADIOGRAPH OF CHEST : PA VIEW

Soft tissue and bony cage are normal.

Both lungs are normal.

Both domes of diaphragm are normal in position and contour.

Hilar shadows are normal.

Mediastinum is central.

Both costo-phrenic angles are clear.

Cardiac size and shape are within normal limits.

IMPRESSION:

- NO OBVIOUS ABNORMALITY.

DR. SHUBHAM SINGHAL
CONSULTANT RADIOLOGIST

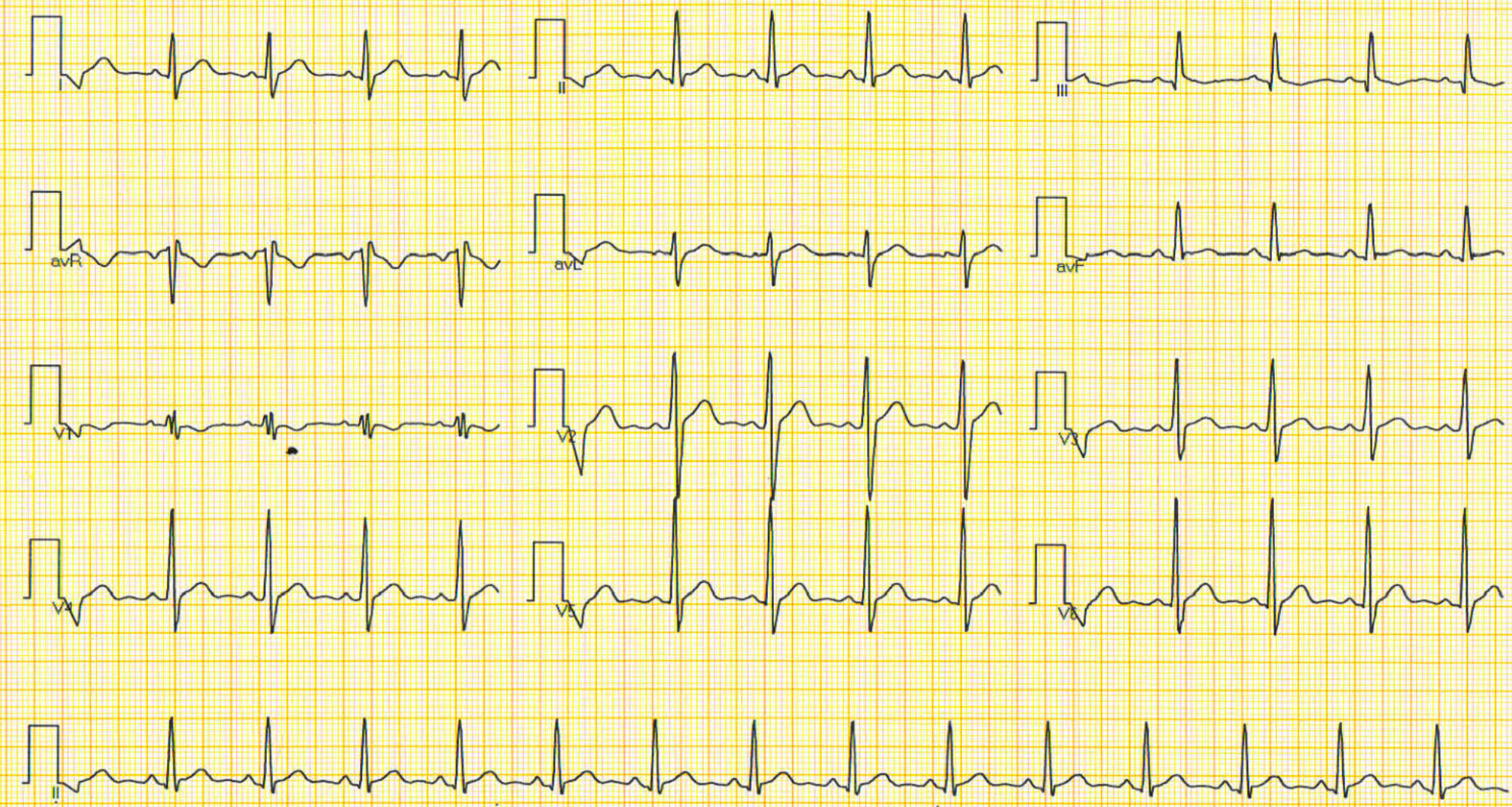
AAKRITI LABS PVT.LTD JAIPUR

ECG

48692 / MR. KUSHI RAM / 32 Yrs / M/ Smoker

Heart Rate : 88 bpm / Tested On : 28-Jan-23 12:03:05 / HF 0.05 Hz - LF 100 Hz / Notch 50 Hz / Sn 1.00 Cm/mV / Sw 25 mm/s

/ Refd By: MEDI WHEEL



Vent Rate : 88 bpm
PR Interval : 138 ms
QRS Duration: 100 ms
QT/QTc Int : 348/397 ms
P-QRS-T axis: 57.00• 68.00• 23.00•

Allengers ECG (Pisces)(PIS215190517)

Reported By:

DR. ARIF HUSSAIN KHAN
MBBS PGDCC
RMC NUMBER 023361

wellness partner

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CIN NO.: U85195RJ2004PTC019563

NAME	MR. KHUSHI RAM MEENA	AGE	32 Y	SEX	MALE
REF BY	MEDIWHEEL	DATE	28/01/2023	REG NO	

ECHOCARDIOGRAM REPORT

WINDOW- POOR/ADEQUATE/GOODVALVE

MITRAL	NORMAL	TRICUSPID	NORMAL
AORTIC	NORMAL	PULMONARY	NORMAL

2D/M-MOD

IVSD mm	9.8	IVSS mm	9.5	AORTA mm	22.7
LVID mm	38.2	LVIS mm	24.7	LA mm	31.8
LVPWD mm	8.1	LVPWS mm	9.8	EF%	60%

CHAMBERS

LA	NORMAL	RA	NORMAL
LV	NORMAL	RV	NORMAL
PERICARDIUM	NORMAL		

DOPPLER STUDY MITRAL

PEAK VELOCITY m/s E/A	0.65/0.59	PEAK GRADIANT MmHg	
MEAN VELOCITY m/s		MEAN GRADIANT MmHg	
MVA cm2 (PLANITMETERY)		MVA cm2 (PHT)	
MR			

AORTIC

PEAK VELOCITY m/s	1.11	PEAK GRADIANT MmHg	
MEAN VELOCITY m/s		MEAN GRADIANT MmHg	
AR			

TRICUSPID

PEAK VELOCITY m/s	0.73	PEAK GRADIANT MmHg	
MEAN VELOCITY m/s		MEAN GRADIANT MmHg	
TR		PASP mmHg	

PULMONARY

PEAK VELOCITY m/s	1.42	PEAK GRADIANT MmHg	
MEAN VELOCITY m/s		MEAN GRADIANT MmHg	
PR		RVEDP mmHg	

IMPRESSION

- **NORMAL LV SYSTOLIC & DIASTOLIC FUNCTION**
- **NO RWMA LVEF 60%**
- **NORMAL RV FUNCTION**
- **NORMAL CHAMBER DIMENSIONS**
- **NORMAL VALVULAR ECHO**
- **INTACT IAS / IVS**
- **NO THROMBUS, NO VEGETATION, NORMAL PERICARDIUM.**
- **IVC NORMAL**

CONCLUSION : FAIR LV FUNCTION.

Cardiologist

wellness partner

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CIN NO.: U85195RJ2004PTC019563

Name	: Mr. KHUSHI RAM MEENA	Registration No:	50914
Age/Gender:	32 Y/Male	Registered	: 28/Jan/2023 10:46AM
Patient ID	: 012301280033	Analysed	: 28/Jan/2023 11:15AM
BarcodeNo	: 10074594	Reported	: 28/Jan/2023 11:15AM
Referred By	: Self	Panel	: Medi Wheel (ArcoFemi Healthcare Ltd)

USG: WHOLE ABDOMEN (Male)

LIVER : **Is mild enlarged in size and shape with 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.Spleenic hilum is not dilated.

KIDNEYS : Right Kidney:-Size: 94 x 45 mm, Left Kidney:-Size: 94 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.
:No evidence of mass or stone in bladder lumen.

PROSTATE:Is normal in size, shape and echotexture,
measures: 30 x 27 x 26 mm, wt: 11 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 hepatomegaly with fatty changes

*** End Of Report ***

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

MC-5333

PATIENT NAME : KHUSHI RAM MEENA | **REF. DOCTOR :** SELF

CODE/NAME & ADDRESS : C000049066
SRL JAIPUR WELLNESS CORPORATE WALK IN
AAKRITI LABS PVT LTD. A-430, AGRASEN MARG
JAIPUR 302017
9314660100

ACCESSION NO : **0251WA001961**
PATIENT ID : KHUSM280191251
CLIENT PATIENT ID: 012301280033
ABHA NO :

AGE/SEX : 32 Years Male
DRAWN : 28/01/2023 10:46:00
RECEIVED : 28/01/2023 11:03:30
REPORTED : 28/01/2023 16:07:56

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

HAEMATOLOGY - CBC

MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE

BLOOD COUNTS,EDTA WHOLE BLOOD

HEMOGLOBIN (HB) METHOD : CYANIDE FREE DETERMINATION	14.3	13.0 - 17.0	g/dL
RED BLOOD CELL (RBC) COUNT METHOD : ELECTRICAL IMPEDANCE	4.75	4.5 - 5.5	mil/µL
WHITE BLOOD CELL (WBC) COUNT METHOD : ELECTRICAL IMPEDANCE	8.10	4.0 - 10.0	thou/µL
PLATELET COUNT METHOD : ELECTRONIC IMPEDANCE	250	150 - 410	thou/µL

RBC AND PLATELET INDICES

HEMATOCRIT (PCV) METHOD : CALCULATED PARAMETER	44.3	40 - 50	%
MEAN CORPUSCULAR VOLUME (MCV) METHOD : CALCULATED PARAMETER	93.0	83 - 101	fL
MEAN CORPUSCULAR HEMOGLOBIN (MCH) METHOD : CALCULATED PARAMETER	30.1	27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC) METHOD : CALCULATED PARAMETER	32.3	31.5 - 34.5	g/dL
RED CELL DISTRIBUTION WIDTH (RDW) METHOD : CALCULATED PARAMETER	13.0	11.6 - 14.0	%
MENTZER INDEX	19.6		
MEAN PLATELET VOLUME (MPV) METHOD : CALCULATED PARAMETER	**11.4 High**	6.8 - 10.9	fL

WBC DIFFERENTIAL COUNT

NEUTROPHILS METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY	60	40 - 80	%
LYMPHOCYTES METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY	35	20 - 40	%
MONOCYTES METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY	03	2 - 10	%
EOSINOPHILS METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY	02	1 - 6	%

Dr. Akansha Jain
Consultant Pathologist

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Rajasthan, INDIA

Patient Ref. No. 775000002221948

PATIENT NAME : KHUSHI RAM MEENA **REF. DOCTOR :** SELF

CODE/NAME & ADDRESS : C000049066 SRL JAIPUR WELLNESS CORPORATE WALK IN AAKRITI LABS PVT LTD. A-430, AGRASEN MARG JAIPUR 302017 9314660100	ACCESSION NO : **0251WA001961** PATIENT ID : KHUSM280191251 CLIENT PATIENT ID: 012301280033 ABHA NO :	AGE/SEX :32 Years Male DRAWN :28/01/2023 10:46:00 RECEIVED :28/01/2023 11:03:30 REPORTED :28/01/2023 16:07:56

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BASOPHILS	00	0 - 2	%
METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY			
ABSOLUTE NEUTROPHIL COUNT	4.86	2.0 - 7.0	thou/µL
METHOD : CALCULATED PARAMETER			
ABSOLUTE LYMPHOCYTE COUNT	2.84	1.0 - 3.0	thou/µL
METHOD : CALCULATED PARAMETER			
ABSOLUTE MONOCYTE COUNT	0.24	0.2 - 1.0	thou/µL
METHOD : CALCULATED PARAMETER			
ABSOLUTE EOSINOPHIL COUNT	0.16	0.02 - 0.50	thou/µL
METHOD : CALCULATED PARAMETER			
ABSOLUTE BASOPHIL COUNT	**0 Low**	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 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),WHOLE BLOOD			
E.S.R	11	0 - 14	mm at 1 hr

METHOD : AUTOMATED (PHOTOMETRICAL CAPILLARY STOPPED FLOW KINETIC ANALYSIS)"

Interpretation(s)
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, 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.

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Consultant Pathologist

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Patient Ref. No. 775000002221948

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

IMMUNOHAEMATOLOGY

MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP	TYPE AB		
METHOD : TUBE AGGLUTINATION			
RH TYPE	POSITIVE		
METHOD : TUBE AGGLUTINATION			

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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Rajasthan, INDIA

Accession No. 275000002221948

MC-5333

PATIENT NAME : KHUSHI RAM MEENA | **REF. DOCTOR :** SELF

CODE/NAME & ADDRESS : C000049066
SRL JAIPUR WELLNESS CORPORATE WALK IN
AAKRITI LABS PVT LTD. A-430, AGRASEN MARG
JAIPUR 302017
9314660100

ACCESSION NO : **0251WA001961**
PATIENT ID : KHUSM280191251
CLIENT PATIENT ID: 012301280033
ABHA NO :

AGE/SEX :32 Years Male
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Test Report Status **Final**	Results	Biological Reference Interval	Units

BIOCHEMISTRY

MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE

GLUCOSE FASTING,FLUORIDE PLASMA			
FBS (FASTING BLOOD SUGAR)	**101 High**	74 - 99	mg/dL
METHOD : GLUCOSE OXIDASE			
GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD			
HBA1C	5.6	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)	%
METHOD : HIGH PERFORMANCE LIQUID CHROMATOGRAPHY (HPLC)			
ESTIMATED AVERAGE GLUCOSE(EAG)	114.0	< 116.0	mg/dL
METHOD : CALCULATED PARAMETER			
GLUCOSE, POST-PRANDIAL, PLASMA			
PPBS(POST PRANDIAL BLOOD SUGAR)	**148 High**	70 - 140	mg/dL
METHOD : GLUCOSE OXIDASE			
LIPID PROFILE, SERUM			
CHOLESTEROL, TOTAL	**205 High**	< 200 Desirable 200 - 239 Borderline High >/= 240 High	mg/dL
METHOD : CHOLESTEROL OXIDASE			
TRIGLYCERIDES	**240 High**	< 150 Normal 150 - 199 Borderline High 200 - 499 High >/=500 Very High	mg/dL
METHOD : LIPASE/GPO-PAP NO CORRECTION			
HDL CHOLESTEROL	**36 Low**	< 40 Low >/=60 High	mg/dL
METHOD : DIRECT CLEARANCE METHOD			

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Lab Ref. No. 275000002221948

MC-5333

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Test Report Status **Final**	Results	Biological Reference Interval	Units
CHOLESTEROL LDL	**121 High**	< 100 Optimal 100 - 129 Near optimal/ above optimal 130 - 159 Borderline High 160 - 189 High >/= 190 Very High	mg/dL
NON HDL CHOLESTEROL	**169 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			
VERY LOW DENSITY LIPOPROTEIN	**48.0 High**	</= 30.0	mg/dL
CHOL/HDL RATIO	**5.7 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	**3.4 High**	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	

Interpretation(s)

LIVER FUNCTION PROFILE, SERUM

BILIRUBIN, TOTAL	0.47	0 - 1	mg/dL
METHOD : DIAZO WITH SULPHANILIC ACID			
BILIRUBIN, DIRECT	0.14	0.00 - 0.25	mg/dL
METHOD : DIAZO WITH SULPHANILIC ACID			
BILIRUBIN, INDIRECT	0.33	0.1 - 1.0	mg/dL
METHOD : CALCULATED PARAMETER			
TOTAL PROTEIN	**8.4 High**	6.4 - 8.2	g/dL
METHOD : BIURET REACTION, END POINT			
ALBUMIN	**4.9 High**	3.8 - 4.4	g/dL

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Test Report Status **Final**	Results	Biological Reference Interval	Units
METHOD : BROMOCRESOL GREEN			
GLOBULIN	3.5	2.0 - 4.1	g/dL
METHOD : CALCULATED PARAMETER			
ALBUMIN/GLOBULIN RATIO	1.4	1.0 - 2.1	RATIO
METHOD : CALCULATED PARAMETER			
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	30	0 - 37	U/L
METHOD : TRIS BUFFER NO P5P IFCC / SFBC 37° C			
ALANINE AMINOTRANSFERASE (ALT/SGPT)	**46 High**	0 - 40	U/L
METHOD : TRIS BUFFER NO P5P IFCC / SFBC 37° C			
ALKALINE PHOSPHATASE	57	39 - 117	U/L
METHOD : AMP OPTIMISED TO IFCC 37° C			
GAMMA GLUTAMYL TRANSFERASE (GGT)	26	11 - 50	U/L
METHOD : GAMMA GLUTAMYL-3 CARBOXY-4 NITROANILIDE (IFCC) 37° C			
LACTATE DEHYDROGENASE	352	230 - 460	U/L
BLOOD UREA NITROGEN (BUN), SERUM			
BLOOD UREA NITROGEN	8	5.0 - 18.0	mg/dL
METHOD : UREASE KINETIC			
CREATININE, SERUM			
CREATININE	0.80	0.8 - 1.3	mg/dL
METHOD : ALKALINE PICRATE NO DEPROTEINIZATION			
BUN/CREAT RATIO			
BUN/CREAT RATIO	10.00		
METHOD : CALCULATED PARAMETER			
URIC ACID, SERUM			
URIC ACID	**7.7 High**	3.4 - 7.0	mg/dL
METHOD : URICASE PEROXIDASE WITH ASCORBATE OXIDASE			
TOTAL PROTEIN, SERUM			
TOTAL PROTEIN	**8.4 High**	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	3.5	2.0 - 4.1	g/dL

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Consultant Pathologist

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MC-5333

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CODE/NAME & ADDRESS : C000049066 SRL JAIPUR WELLNESS CORPORATE WALK IN AAKRITI LABS PVT LTD. A-430, AGRASEN MARG JAIPUR 302017 9314660100	ACCESSION NO : **0251WA001961** PATIENT ID : KHUSM280191251 CLIENT PATIENT ID: 012301280033 ABHA NO :	AGE/SEX : 32 Years Male DRAWN : 28/01/2023 10:46:00 RECEIVED : 28/01/2023 11:03:30 REPORTED : 28/01/2023 16:07:56

Test Report Status **Final**	Results	Biological Reference Interval	Units
ELECTROLYTES (NA/K/CL), SERUM			
SODIUM, SERUM	141.9	137 - 145	mmol/L
METHOD : ION-SELECTIVE ELECTRODE			
POTASSIUM, SERUM	4.38	3.6 - 5.0	mmol/L
METHOD : ION-SELECTIVE ELECTRODE			
CHLORIDE, SERUM	101.5	98 - 107	mmol/L
METHOD : ION-SELECTIVE ELECTRODE			

Interpretation(s)

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

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Rajasthan, INDIA

Patient Rep. No. 775000002221948

MC-5333

PATIENT NAME : KHUSHI RAM MEENA **REF. DOCTOR :** SELF

CODE/NAME & ADDRESS : C000049066 SRL JAIPUR WELLNESS CORPORATE WALK IN AAKRITI LABS PVT LTD. A-430, AGRASEN MARG JAIPUR 302017 9314660100	ACCESSION NO : **0251WA001961** PATIENT ID : KHUSM280191251 CLIENT PATIENT ID: 012301280033 ABHA NO :	AGE/SEX : 32 Years Male DRAWN : 28/01/2023 10:46:00 RECEIVED : 28/01/2023 11:03:30 REPORTED : 28/01/2023 16:07:56

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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, 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.

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MC-5333

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CLINICAL PATH - URINALYSIS

MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE

PHYSICAL EXAMINATION, URINE			
COLOR METHOD : GROSS EXAMINATION	PALE YELLOW		
APPEARANCE METHOD : GROSS EXAMINATION	CLEAR		
CHEMICAL EXAMINATION, URINE			
PH METHOD : DOUBLE INDICATOR PRINCIPLE	6.0	4.7 - 7.5	
SPECIFIC GRAVITY METHOD : IONIC CONCENTRATION METHOD	1.010	1.003 - 1.035	
PROTEIN METHOD : PROTEIN ERROR OF INDICATORS WITH REFLECTANCE	NOT DETECTED	NOT DETECTED	
GLUCOSE METHOD : GLUCOSE OXIDASE PEROXIDASE / BENEDICTS	NOT DETECTED	NOT DETECTED	
KETONES METHOD : SODIUM NITROPRUSSIDE REACTION	NOT DETECTED	NOT DETECTED	
BLOOD METHOD : PEROCIDASE ANTI PEROXIDASE	NOT DETECTED	NOT DETECTED	
BILIRUBIN METHOD : DIPSTICK	NOT DETECTED	NOT DETECTED	
UROBILINOGEN METHOD : EHRLICH REACTION REFLECTANCE	NORMAL	NORMAL	
NITRITE METHOD : NITRATE TO NITRITE CONVERSION METHOD	NOT DETECTED	NOT DETECTED	
LEUKOCYTE ESTERASE	NOT DETECTED	NOT DETECTED	
MICROSCOPIC EXAMINATION, URINE			
RED BLOOD CELLS METHOD : MICROSCOPIC EXAMINATION	NOT DETECTED	NOT DETECTED	/HPF
PUS CELL (WBC'S) METHOD : DIPSTICK, MICROSCOPY	1-2	0-5	/HPF
EPITHELIAL CELLS METHOD : MICROSCOPIC EXAMINATION	0-1	0-5	/HPF
CASTS	NOT DETECTED		

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METHOD : MICROSCOPIC EXAMINATION			
CRYSTALS	NOT DETECTED		
METHOD : MICROSCOPIC EXAMINATION			
BACTERIA	NOT DETECTED	NOT DETECTED	
METHOD : MICROSCOPIC EXAMINATION			
YEAST	NOT DETECTED	NOT DETECTED	

Interpretation(s)

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CLINICAL PATH - STOOL ANALYSIS

MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE

PHYSICAL EXAMINATION,STOOL

COLOUR	SAMPLE NOT RECEIVED		

METHOD : GROSS EXAMINATION

Dr. Abhishek Sharma
Consultant Microbiologist

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AAKRITI LABS PVT LTD. A-430, AGRASEN MARG
JAIPUR 302017
9314660100

ACCESSION NO : **0251WA001961**
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CLIENT PATIENT ID: 012301280033
ABHA NO :

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SPECIALISED CHEMISTRY - HORMONE

MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE

THYROID PANEL, SERUM

T3	109.07	60.0 - 181.0	ng/dL
METHOD : CHEMILUMINESCENCE			
T4	**11.50 High**	4.5 - 10.9	µg/dL
METHOD : CHEMILUMINESCENCE			
TSH (ULTRASENSITIVE)	2.756	0.550 - 4.780	µIU/mL
METHOD : CHEMILUMINESCENCE			

Interpretation(s)

Triiodothyronine T3 , Thyroxine T4, and **Thyroid Stimulating Hormone TSH** are thyroid hormones which affect almost every physiological process in the body, including growth, development, metabolism, body temperature, and heart rate.
Production of T3 and its prohormone thyroxine (T4) is activated by thyroid-stimulating hormone (TSH), which is released from the pituitary gland. Elevated concentrations of T3, and T4 in the blood inhibit the production of TSH.
Excessive secretion of thyroxine in the body is hyperthyroidism, and deficient secretion is called hypothyroidism.
In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hyperthyroidism, TSH levels are low.
owidctlparowidctlparBelow mentioned are the guidelines for Pregnancy related reference ranges for Total T4, TSH & Total T3.Measurement of the serum TT3 level is a more sensitive test for the diagnosis of hyperthyroidism, and measurement of TT4 is more useful in the diagnosis of hypothyroidism.Most of the thyroid hormone in blood is bound to transport proteins. Only a very small fraction of the circulating hormone is free and biologically active. It is advisable to detect Free T3, FreeT4 along with TSH, instead of testing for albumin bound Total T3, Total T4.

Sr. No.	TSH	Total T4	FT4	Total T3	Possible Conditions
1	High	Low	Low	Low	(1) Primary Hypothyroidism (2) Chronic autoimmune Thyroiditis (3) Post Thyroidectomy (4) Post Radio-Iodine treatment
2	High	Normal	Normal	Normal	(1)Subclinical Hypothyroidism (2) Patient with insufficient thyroid hormone replacement therapy (3) In cases of Autoimmune/Hashimoto thyroiditis (4). Isolated increase in TSH levels can be due to Subclinical inflammation, drugs like amphetamines, Iodine containing drug and dopamine antagonist e.g. domperidone and other physiological reasons.
3	Normal/Low	Low	Low	Low	(1) Secondary and Tertiary Hypothyroidism
4	Low	High	High	High	(1) Primary Hyperthyroidism (Graves Disease) (2) Multinodular Goitre (3)Toxic Nodular Goitre (4) Thyroiditis (5) Over treatment of thyroid hormone (6) Drug effect e.g. Glucocorticoids, dopamine, T4 replacement therapy (7) First trimester of Pregnancy
5	Low	Normal	Normal	Normal	(1) Subclinical Hyperthyroidism
6	High	High	High	High	(1) TSH secreting pituitary adenoma (2) TRH secreting tumor
7	Low	Low	Low	Low	(1) Central Hypothyroidism (2) Euthyroid sick syndrome (3) Recent treatment for Hyperthyroidism

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8	Normal/Low	Normal	Normal	High	(1) T3 thyrotoxicosis (2) Non-Thyroidal illness
9	Low	High	High	Normal	(1) T4 Ingestion (2) Thyroiditis (3) Interfering Anti TPO antibodies

REF: 1. TIETZ Fundamentals of Clinical chemistry 2.Guidlines of the American Thyroid association during pregnancy and Postpartum, 2011.

NOTE: It is advisable to detect Free T3,FreeT4 along with TSH, instead of testing for albumin bound Total T3, Total T4.TSH is not affected by variation in thyroid - binding protein. TSH has a diurnal rhythm, with peaks at 2:00 - 4:00 a.m. And troughs at 5:00 - 6:00 p.m. With ultradian variations.

****End Of Report****

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