

Vent Rate : 98 bpm

PR Interval : 122 ms

QRS Duration: 86 ms

QT/QTc Int : 330/397 ms

P-QRS-T axis: 51.00 • 15.00 • 36.00 •

Allengers ECG (Pices)(PIS215190517)

Handwritten signature

Dr. NITIZ GOYAL
M.B.B.S., M.D.

RMC - 023319

Reported By:



Aakriti Labs


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

PATIENT NAME: MR VEER SINGH JADAV	AGE & SEX: 58Y/M
REF. by: MEDI WHEEL	DATE: 05/11/2023

USG: WHOLE ABDOMEN (Male)

- LIVER** : Is normal in size, shape and 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** : 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: 40 x 32 x 29 mm, wt: 20 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:- NORMAL STUDY


DR NEERA MEHTA
MBBS, DMRD
RMCNO.005807/14853



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www.aakritilabs.com
CIN NO.: U85195RJ2004PTC019563

PATIENT : MR. VEER SINGH JATAV	AGE / SEX: 58Y/MALE
REF. BY : MEDIWHEEL	DATE : 31.10.2023

REPORT: DIGITAL X-RAY CHEST PA VIEW

Soft tissue shadow and bony cages are normal.

Trachea is central.

Bilateral lung field and both CP angle is clear.

Domes of diaphragm are normally placed.

Transverse diameter of heart appears with normal limits.

IMPRESSION:- NO OBVIOUS ABNORMALITY DETECTED.


DR NEERA MEHTA
MBBS, DMRD



Aakriti Labs

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Tonk Road, Jaipur (Raj.) Ph.: 0141-2710661
www.aakritilabs.com
CIN NO.: U85195RJ2004PTC019563

NAME	MR VEER SINGH JATAV	AGE	58Y	SEX	MALE
REF BY	MEDI WHEEL	DATE	31/10/2023	REG NO	

ECHOCARDIOGRAM REPORT

WINDOW- POOR/ADEQUATE/GOODVALVE

MITRAL	NORMAL	TRICUSPID	NORMAL
AORTIC	NORMAL	PULMONARY	NORMAL

2D/M-MOD

IVSD mm	8.5	IVSS mm	12.9	AORTA mm	26.4
LVID mm	40.9	LVIS mm	26.7	LA mm	28.1
LVPWD mm	8.8	LVPWS mm	12.9	EF%	60%

CHAMBERS

LA	NORMAL	RA	NORMAL
LV	NORMAL	RV	NORMAL
PERICARDIUM	NORMAL		

DOPPLER STUDY MITRAL

PEAK VELOCITY m/s E/A	0.77/1.28	PEAK GRADIANT MmHg	
MEAN VELOCITY m/s		MEAN GRADIANT MmHg	
MVA cm2 (PLANIMETERY)		MVA cm2 (PHT)	
MR			

AORTIC

PEAK VELOCITY m/s	1.42	PEAK GRADIANT MmHg	
MEAN VELOCITY m/s		MEAN GRADIANT MmHg	
AR			

TRICUSPID

PEAK VELOCITY m/s	0.46	PEAK GRADIANT MmHg	
MEAN VELOCITY m/s		MEAN GRADIANT MmHg	
TR		PASP mmHg	

PULMONARY

PEAK VELOCITY m/s	0.95	PEAK GRADIANT MmHg	
MEAN VELOCITY m/s		MEAN GRADIANT MmHg	
PR		RVEDP mmHg	

IMPRESSION

- LV DIASTOLIC DYSFUNCTION GRADE-1
- NORMAL LV SYSTOLIC 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

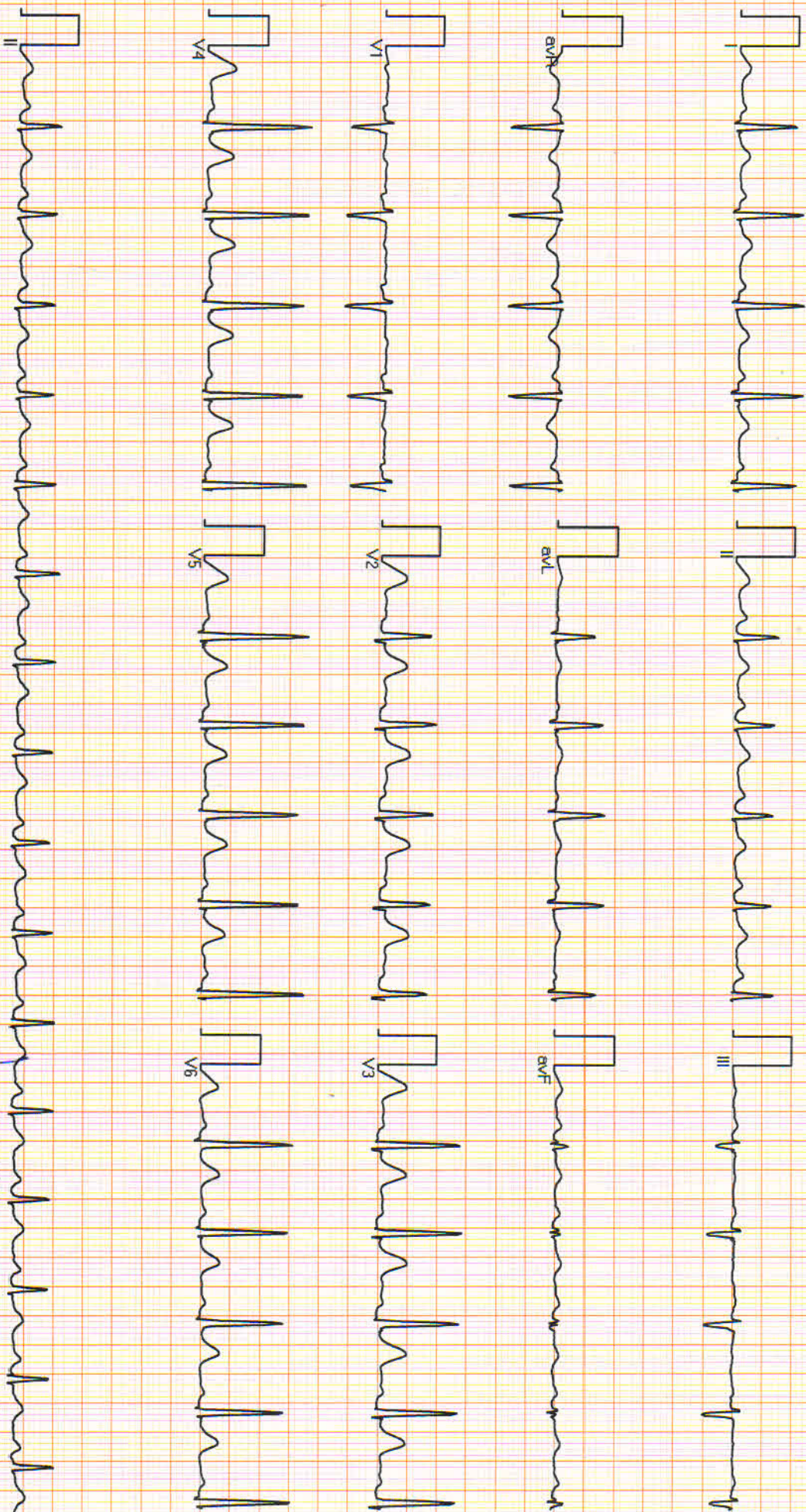
CONCLUSION : DIASTOLIC DYSFUNCTION, FAIR LV FUNCTION.


Cardiologist

AAKRITI LABS PVT.LTD JAIPUR

68853 / MR. VEER SINGH JATAV / 58 Yrs / M / Non Smoker
Heart Rate : 98 bpm / Tested On : 31-Oct-23 09:03:29 / HF 0.05 Hz - LF 100 Hz / Notch 50 Hz / Sn 1.00 Cm/mV / Sw 25 mm/s
/ Refd By: ACROFEMI HEALTH

ECG



Vent Rate : 98 bpm
PR Interval : 122 ms
QRS Duration: 86 ms
QT/QTc Int : 330/397 ms
P-QRS-T axis: 51.00° • 15.00° • 36.00°
Allengers ECG (Pscs)(PI5215190517)

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Reported By:

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



MC-5726

PATIENT NAME : VEER SINGH JATAV

REF. DOCTOR : SELF

CODE/NAME & ADDRESS : C000138404

ARCOFEMI HEALTHCARE LTD (MEDIWHEEL
F-703, F-703, LADO SARAI, MEHRAULISOUTH
WEST DELHI
NEW DELHI 110030
8800465156

ACCESSION NO : 0251WJ001893

PATIENT ID : VEERM211065251

CLIENT PATIENT ID: 012310210052

ABHA NO :

AGE/SEX : 58 Years Male

DRAWN : 21/10/2023 13:02:00

RECEIVED : 21/10/2023 13:17:58

REPORTED : 05/11/2023 15:20:38

Test Report Status **Final**

Results

Biological Reference Interval Units

HAEMATOTOLOGY - CDC

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

BLOOD COUNTS,EDTA WHOLE BLOOD

HEMOGLOBIN (HB)	13.5	13.0 - 17.0	g/dL
METHOD : CYANIDE FREE DETERMINATION			
RED BLOOD CELL (RBC) COUNT	4.63	4.5 - 5.5	mil/ μ L
METHOD : ELECTRICAL IMPEDANCE			
WHITE BLOOD CELL (WBC) COUNT	6.80	4.0 - 10.0	thou/ μ L
METHOD : ELECTRICAL IMPEDANCE			
PLATELET COUNT	74 Low	150 - 410	thou/ μ L
METHOD : ELECTRONIC IMPEDANCE			

RBC AND PLATELET INDICES

HEMATOCRIT (PCV)	40.8	40 - 50	%
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR VOLUME (MCV)	88.0	83 - 101	fL
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	29.1	27.0 - 32.0	pg
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC)	33.0	31.5 - 34.5	g/dL
METHOD : CALCULATED PARAMETER			
RED CELL DISTRIBUTION WIDTH (RDW)	13.8	11.6 - 14.0	%
METHOD : CALCULATED PARAMETER			
MENTZER INDEX	19.0		
MEAN PLATELET VOLUME (MPV)	11.3 High	6.8 - 10.9	fL
METHOD : CALCULATED PARAMETER			

WBC DIFFERENTIAL COUNT

NEUTROPHILS	61	40 - 80	%
METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY			
LYMPHOCYTES	34	20 - 40	%
METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY			
MONOCYTES	04	2 - 10	%

Dr. Akansha Jain
Consultant Pathologist



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Agilus Diagnostics Ltd.
C/O Aakriti Labs Pvt Ltd, 3, Mahatma Gandhi Marg,Gandhi Nagar Mod, Tonk Road
Jaipur, 302015
Rajasthan, India





MC-5726

PATIENT NAME : VEER SINGH JATAV**REF. DOCTOR : SELF****CODE/NAME & ADDRESS : C000138404**ARCOFEMI HEALTHCARE LTD (MEDIWHEEL
F-703, F-703, LADO SARAI, MEHRAULISOUTH
WEST DELHI
NEW DELHI 110030
8800465156**ACCESSION NO : 0251WJ001893****PATIENT ID : VEERM211065251****CLIENT PATIENT ID : 012310210052****ABHA NO :****AGE/SEX : 58 Years Male****DRAWN : 21/10/2023 13:02:00****RECEIVED : 21/10/2023 13:17:58****REPORTED : 05/11/2023 15:20:38**

Test Report Status	Final	Results	Biological Reference Interval	Units
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METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY
EOSINOPHILS

01

1 - 6

%

METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY
BASOPHILS

00

0 - 2

%

METHOD : IMPEDANCE WITH HYDRO FOCUS AND MICROSCOPY
ABSOLUTE NEUTROPHIL COUNT

4.15

2.0 - 7.0

thou/ μ L

METHOD : CALCULATED PARAMETER

ABSOLUTE LYMPHOCYTE COUNT

2.31

1.0 - 3.0

thou/ μ L

METHOD : CALCULATED PARAMETER

ABSOLUTE MONOCYTE COUNT

0.27

0.2 - 1.0

thou/ μ L

METHOD : CALCULATED PARAMETER

ABSOLUTE EOSINOPHIL COUNT

0.07

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)

1.8

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.

Dr. Akansha Jain
Consultant Pathologist

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C/O Aakriti Labs Pvt Ltd, 3, Mahatma Gandhi Marg,Gandhi Nagar Mod, Tonk Road
Jaipur, 302015
Rajasthan, India**Patient Ref. No. 775000005181322**



MC-5726

PATIENT NAME : VEER SINGH JATAV **REF. DOCTOR : SELF**

CODE/NAME & ADDRESS : C000138404 ARCOFEMI HEALTHCARE LTD (MEDIWHEEL F-703, F-703, LADO SARAI, MEHRAULISOUTH WEST DELHI NEW DELHI 110030 8800465156	ACCESSION NO : 0251WJ001893 PATIENT ID : VEERM211065251 CLIENT PATIENT ID: 012310210052 ABHA NO :	AGE/SEX : 58 Years Male DRAWN : 21/10/2023 13:02:00 RECEIVED : 21/10/2023 13:17:58 REPORTED : 05/11/2023 15:20:38
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Test Report Status	Final	Results	Biological Reference Interval	Units
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HAE MATOLOGY

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD

HBA1C	7.3 High	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)	162.8 High	< 116.0	mg/dL
METHOD : CALCULATED PARAMETER			

Dr. Akansha Jain
Consultant Pathologist



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C/O Aakriti Labs Pvt Ltd, 3, Mahatma Gandhi Marg, Gandhi Nagar Mod, Tonk Road
Jaipur, 302015
Rajasthan, India



PATIENT NAME : VEER SINGH JATAV

REF. DOCTOR : SELF

CODE/NAME & ADDRESS : C000138404

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WEST DELHI
NEW DELHI 110030
8800465156

ACCESSION NO : 0251WJ001893

PATIENT ID : VEERM211065251

CLIENT PATIENT ID: 012310210052

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AGE/SEX : 58 Years Male

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Test Report Status **Final**

Results

Biological Reference Interval Units

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

ERYTHROCYTE SEDIMENTATION RATE (ESR),WHOLE BLOOD

E.S.R 05 0 - 14 mm at 1 hr

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

Interpretation(s)

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 :

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.
4. Interference of hemoglobinopathies in HbA1c estimation is seen in

a) Homozygous hemoglobinopathy. Fructosamine is recommended for testing of HbA1c.

b) Heterozygous state detected (D10 is corrected for HbS & HbC trait.)

c) HbF > 25% on alternate paltform (Boronate affinity chromatography) is recommended for testing of HbA1c.Abnormal Hemoglobin electrophoresis (HPLC method) is recommended for detecting a hemoglobinopathy

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: 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. AACCC Press, 7th edition. Edited by S. Soldin;3. The reference for the adult reference range is "Practical Haematology by Dacie and Lewis,10th edition.

Dr. Akansha Jain
Consultant Pathologist

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Jaipur, 302015
Rajasthan, India



Patient Ref. No. 775000005181322

PATIENT NAME : VEER SINGH JATAV

REF. DOCTOR : SELF

CODE/NAME & ADDRESS : C000138404

ARCOFEMI HEALTHCARE LTD (MEDIWHEEL
F-703, F-703, LADO SARAI, MEHRAULISOUTH
WEST DELHI
NEW DELHI 110030
8800465156

ACCESSION NO : 0251WJ001893

PATIENT ID : VEERM211065251

CLIENT PATIENT ID: 012310210052

ABHA NO :

AGE/SEX : 58 Years Male

DRAWN : 21/10/2023 13:02:00

RECEIVED : 21/10/2023 13:17:58

REPORTED : 05/11/2023 15:20:38

Test Report Status Final

Results

Biological Reference Interval Units

IMMUNOHAEMATOLGY

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP

TYPE B

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.

Dr. Akansha Jain
Consultant Pathologist

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Rajasthan, India



Patient Ref. No. 775000005181322



MC-5726

PATIENT NAME : VEER SINGH JATAV

REF. DOCTOR : SELF

CODE/NAME & ADDRESS : C000138404
ARCOFEMI HEALTHCARE LTD (MEDIWHEEL
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Table with 4 columns: Test Report Status, Results, Biological Reference Interval, Units. Status is Final.

BIOCHEMISTRY

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

GLUCOSE FASTING, FLUORIDE PLASMA

FBS (FASTING BLOOD SUGAR) 148 High 74 - 99 mg/dL
METHOD : GLUCOSE OXIDASE

GLUCOSE, POST-PRANDIAL, PLASMA

PPBS(POST PRANDIAL BLOOD SUGAR) 207 High 70 - 140 mg/dL
METHOD : GLUCOSE OXIDASE

LIPID PROFILE WITH CALCULATED LDL

CHOLESTEROL, TOTAL 125 < 200 Desirable, 200 - 239 Borderline High, >/= 240 High mg/dL
METHOD : CHOLESTEROL OXIDASE

TRIGLYCERIDES 143 < 150 Normal, 150 - 199 Borderline High, 200 - 499 High, >/=500 Very High mg/dL
METHOD : LIPASE/GPO-PAP NO CORRECTION

HDL CHOLESTEROL 47 < 40 Low, >/=60 High mg/dL
METHOD : DIRECT CLEARANCE METHOD

CHOLESTEROL LDL 50 < 100 Optimal, 100 - 129 Near optimal/ above optimal, 130 - 159 Borderline High, 160 - 189 High, >/= 190 Very High mg/dL

NON HDL CHOLESTEROL 78 Desirable: Less than 130, Above Desirable: 130 - 159, Borderline High: 160 - 189, High: 190 - 219, Very high: > or = 220 mg/dL
METHOD : CALCULATED PARAMETER

Handwritten signature of Dr. Akansha Jain

Dr. Akansha Jain
Consultant Pathologist



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Test Report Status Final Results Biological Reference Interval Units

VERY LOW DENSITY LIPOPROTEIN	28.6	</= 30.0	mg/dL
CHOL/HDL RATIO	2.7 Low	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	1.1	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	

Interpretation(s)

Serum lipid profile is measured for cardiovascular risk prediction. Lipid Association of India recommends LDL-C as primary target and Non HDL-C as co-primary treatment target.

Risk Stratification for ASCVD (Atherosclerotic cardiovascular disease) by Lipid Association of India

Risk Category	
Extreme risk group	A. CAD with > 1 feature of high risk group B. CAD with > 1 feature of Very high risk group or recurrent ACS (within 1 year) despite LDL-C < or = 50 mg/dl or polyvascular disease
Very High Risk	1. Established ASCVD 2. Diabetes with 2 major risk factors or evidence of end organ damage 3. Familial Homozygous Hypercholesterolemia
High Risk	1. Three major ASCVD risk factors. 2. Diabetes with 1 major risk factor or no evidence of end organ damage. 3. CKD stage 3B or 4. 4. LDL >190 mg/dl 5. Extreme of a single risk factor. 6. Coronary Artery Calcium - CAC >300 AU. 7. Lipoprotein a >= 50mg/dl 8. Non stenotic carotid plaque
Moderate Risk	2 major ASCVD risk factors
Low Risk	0-1 major ASCVD risk factors

Major ASCVD (Atherosclerotic cardiovascular disease) Risk Factors

1. Age > or = 45 years in males and > or = 55 years in females	3. Current Cigarette smoking or tobacco use
2. Family history of premature ASCVD	4. High blood pressure
5. Low HDL	

Newer treatment goals and statin initiation thresholds based on the risk categories proposed by LAI in 2020.

Risk Group	Treatment Goals		Consider Drug Therapy	
	LDL-C (mg/dl)	Non-HDL (mg/dl)	LDL-C (mg/dl)	Non-HDL (mg/dl)
Extreme Risk Group Category A	<50 (Optional goal < OR = 30)	< 80 (Optional goal < OR = 60)	>OR = 50	>OR = 80

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Consultant Pathologist

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Patient Ref. No. 775000005181322



MC-5726

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Test Report Status Final Results Biological Reference Interval Units

Extreme Risk Group Category B	<OR = 30	<OR = 60	> 30	>60
Very High Risk	<50	<80	>OR= 50	>OR= 80
High Risk	<70	<100	>OR= 70	>OR= 100
Moderate Risk	<100	<130	>OR= 100	>OR= 130
Low Risk	<100	<130	>OR= 130*	>OR= 160

*After an adequate non-pharmacological intervention for at least 3 months.

References: Management of Dyslipidaemia for the Prevention of Stroke: Clinical Practice Recommendations from the Lipid Association of India. Current Vascular Pharmacology, 2022, 20, 134-155.**LIVER FUNCTION PROFILE, SERUM**

BILIRUBIN, TOTAL	0.88	0 - 1	mg/dL
METHOD : DIAZO WITH SULPHANILIC ACID			
BILIRUBIN, DIRECT	0.27 High	0.00 - 0.25	mg/dL
METHOD : DIAZO WITH SULPHANILIC ACID			
BILIRUBIN, INDIRECT	0.61	0.1 - 1.0	mg/dL
METHOD : CALCULATED PARAMETER			
TOTAL PROTEIN	7.8	6.4 - 8.2	g/dL
METHOD : BIURET REACTION, END POINT			
ALBUMIN	4.6 High	3.8 - 4.4	g/dL
METHOD : BROMOCRESOL GREEN			
GLOBULIN	3.2	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)	37	0 - 37	U/L
METHOD : TRIS BUFFER NO P5P IFCC / SFBC 37° C			
ALANINE AMINOTRANSFERASE (ALT/SGPT)	38	0 - 40	U/L
METHOD : TRIS BUFFER NO P5P IFCC / SFBC 37° C			
ALKALINE PHOSPHATASE	81	39 - 117	U/L
METHOD : AMP OPTIMISED TO IFCC 37° C			
GAMMA GLUTAMYL TRANSFERASE (GGT)	103 High	11 - 50	U/L
METHOD : GAMMA GLUTAMYL-3 CARBOXY-4 NITROANILIDE (IFCC) 37° C			
LACTATE DEHYDROGENASE	325	230 - 460	U/L

BLOOD UREA NITROGEN (BUN), SERUM

BLOOD UREA NITROGEN	8	5.0 - 18.0	mg/dL
METHOD : UREASE KINETIC			

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Rajasthan, India

Patient Ref. No. 775000005181322



MC-5726

PATIENT NAME : VEER SINGH JATAV**REF. DOCTOR : SELF****CODE/NAME & ADDRESS : C000138404**ARCOFEMI HEALTHCARE LTD (MEDIWHEEL
F-703, F-703, LADO SARAI, MEHRAULISOUTH
WEST DELHI
NEW DELHI 110030
8800465156**ACCESSION NO : 0251WJ001893**

PATIENT ID : VEERM211065251

CLIENT PATIENT ID: 012310210052

ABHA NO :

AGE/SEX : 58 Years Male

DRAWN : 21/10/2023 13:02:00

RECEIVED : 21/10/2023 13:17:58

REPORTED : 05/11/2023 15:20:38

Test Report Status Final**Results****Biological Reference Interval Units****CREATININE, SERUM**

CREATININE

0.83

0.8 - 1.3

mg/dL

METHOD : ALKALINE PICRATE NO DEPROTEINIZATION

BUN/CREAT RATIO

BUN/CREAT RATIO

9.64

METHOD : CALCULATED PARAMETER

URIC ACID, SERUM

URIC ACID

6.2

3.4 - 7.0

mg/dL

METHOD : URICASE PEROXIDASE WITH ASCORBATE OXIDASE

TOTAL PROTEIN, SERUM

TOTAL PROTEIN

7.8

6.4 - 8.3

g/dL

METHOD : BIURET REACTION, END POINT

ALBUMIN, SERUM

ALBUMIN

4.6 High

3.8 - 4.4

g/dL

METHOD : BROMOCRESOL GREEN

GLOBULIN

GLOBULIN

3.2

2.0 - 4.1

g/dL

ELECTROLYTES (NA/K/CL), SERUM**Dr. Akansha Jain**
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Test Report Status	Final	Results	Biological Reference Interval	Units
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SODIUM, SERUM		139.9	137 - 145	mmol/L
METHOD : ION-SELECTIVE ELECTRODE				
POTASSIUM, SERUM		3.95	3.6 - 5.0	mmol/L
METHOD : ION-SELECTIVE ELECTRODE				
CHLORIDE, SERUM		102.9	98 - 107	mmol/L
METHOD : ION-SELECTIVE ELECTRODE				

Interpretation(s)

Sodium	Potassium	Chloride
Decreased in: CCF, cirrhosis, vomiting, diarrhea, excessive sweating, salt-losing nephropathy, adrenal insufficiency, nephrotic syndrome, water intoxication, SIADH. Drugs: thiazides, diuretics, ACE inhibitors, chlorpropamide, carbamazepine, antidepressants (SSRI), antipsychotics.	Decreased in: Low potassium intake, prolonged vomiting or diarrhea, RTA types I and II, hyperaldosteronism, Cushing's syndrome, osmotic diuresis (e.g., hyperglycemia), alkalosis, familial periodic paralysis, trauma (transient). Drugs: Adrenergic agents, diuretics.	Decreased in: Vomiting, diarrhea, renal failure combined with salt deprivation, over-treatment with diuretics, chronic respiratory acidosis, diabetic ketoacidosis, excessive sweating, SIADH, salt-losing nephropathy, porphyria, expansion of extracellular fluid volume, adrenal insufficiency, hyperaldosteronism, metabolic alkalosis. Drugs: chronic laxative, corticosteroids, diuretics.
Increased in: Dehydration (excessive sweating, severe vomiting or diarrhea), diabetes mellitus, diabetes insipidus, hyperaldosteronism, inadequate water intake. Drugs: steroids, licorice, oral contraceptives.	Increased in: Massive hemolysis, severe tissue damage, rhabdomyolysis, acidosis, dehydration, renal failure, Addison's disease, RTA type IV, hyperkalemic familial periodic paralysis. Drugs: potassium salts, potassium-sparing diuretics, NSAIDs, beta-blockers, ACE inhibitors, high-dose trimethoprim-sulfamethoxazole.	Increased in: Renal failure, nephrotic syndrome, RTA, dehydration, overtreatment with saline, hyperparathyroidism, diabetes insipidus, metabolic acidosis from diarrhea (Loss of HCO3-), respiratory alkalosis, hyperadrenocorticism. Drugs: acetazolamide, androgens, hydrochlorothiazide, salicylates.
Interferences: Severe lipemia or hyperproteinemia, if sodium analysis involves a dilution step can cause spurious results. The serum sodium falls about 1.6 mEq/L for each 100 mg/dL increase in blood glucose.	Interferences: Hemolysis of sample, delayed separation of serum, prolonged fist clenching during blood drawing, and prolonged tourniquet placement. Very high WBC/PLT counts may cause spurious. Plasma potassium levels are normal.	Interferences: Test is helpful in assessing normal and increased anion gap metabolic acidosis and in distinguishing hypercalcemia due to hyperparathyroidism (high serum chloride) from that due to malignancy (Normal serum chloride)

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 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: 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.

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Patient Ref. No. 775000005181322



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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.
 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-

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.

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MC-5726

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Test Report Status	Final	Results	Biological Reference Interval	Units
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CLINICAL PATH - URINALYSIS

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

PHYSICAL EXAMINATION, URINE

COLOR	PALE YELLOW
METHOD : GROSS EXAMINATION	
APPEARANCE	CLEAR
METHOD : GROSS EXAMINATION	

CHEMICAL EXAMINATION, URINE

PH	5.5	4.7 - 7.5
METHOD : DOUBLE INDICATOR PRINCIPLE		
SPECIFIC GRAVITY	<=1.005	1.003 - 1.035
METHOD : IONIC CONCENTRATION METHOD		
PROTEIN	NOT DETECTED	NEGATIVE
METHOD : PROTEIN ERROR OF INDICATORS WITH REFLECTANCE		
GLUCOSE	DETECTED (+)	NOT DETECTED
METHOD : GLUCOSE OXIDASE PEROXIDASE / BENEDICTS		
KETONES	NOT DETECTED	NOT DETECTED
METHOD : SODIUM NITROPRUSSIDE REACTION		
BLOOD	NOT DETECTED	NEGATIVE
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)	0-1	0-5	/HPF
METHOD : DIPSTICK, MICROSCOPY			

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8800465156**ACCESSION NO : 0251WJ001893**

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EPITHELIAL CELLS		1-2	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)

The following table describes the probable conditions, in which the analytes are present in urine

Presence of	Conditions
Proteins	Inflammation or immune illnesses
Pus (White Blood Cells)	Urinary tract infection, urinary tract or kidney stone, tumors or any kind of kidney impairment
Glucose	Diabetes or kidney disease
Ketones	Diabetic ketoacidosis (DKA), starvation or thirst
Urobilinogen	Liver disease such as hepatitis or cirrhosis
Blood	Renal or genital disorders/trauma
Bilirubin	Liver disease
Erythrocytes	Urological diseases (e.g. kidney and bladder cancer, urolithiasis), urinary tract infection and glomerular diseases
Leukocytes	Urinary tract infection, glomerulonephritis, interstitial nephritis either acute or chronic, polycystic kidney disease, urolithiasis, contamination by genital secretions
Epithelial cells	Urolithiasis, bladder carcinoma or hydronephrosis, ureteric stents or bladder catheters for prolonged periods of time
Granular Casts	Low intratubular pH, high urine osmolality and sodium concentration, interaction with Bence-Jones protein
Hyaline casts	Physical stress, fever, dehydration, acute congestive heart failure, renal diseases

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MC-5726

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Calcium oxalate	Metabolic stone disease, primary or secondary hyperoxaluria, intravenous infusion of large doses of vitamin C, the use of vasodilator naftidrofuryl oxalate or the gastrointestinal lipase inhibitor orlistat, ingestion of ethylene glycol or of star fruit (Averrhoa carambola) or its juice
Uric acid	arthritis
Bacteria	Urinary infection when present in significant numbers & with pus cells.
Trichomonas vaginalis	Vaginitis, cervicitis or salpingitis

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CLINICAL PATH - STOOL ANALYSIS

MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE

PHYSICAL EXAMINATION,STOOL

COLOUR	BROWN	
CONSISTENCY	WELL FORMED	
MUCUS	NOT DETECTED	NOT DETECTED
VISIBLE BLOOD	ABSENT	ABSENT

MICROSCOPIC EXAMINATION,STOOL

CYSTS	NOT DETECTED	NOT DETECTED
OVA	NOT DETECTED	
LARVAE	NOT DETECTED	NOT DETECTED
TROPHOZOITES	NOT DETECTED	NOT DETECTED

Interpretation(s)

Stool routine analysis is only a screening test for disorders of gastrointestinal tract like infection, malabsorption, etc. The following table describes the probable conditions, in which the analytes are present in stool.

PRESENCE OF	CONDITION
Pus cells	Pus in the stool is an indication of infection
Red Blood cells	Parasitic or bacterial infection or an inflammatory bowel condition such as ulcerative colitis
Parasites	Infection of the digestive system. Stool examination for ova and parasite detects presence of parasitic infestation of gastrointestinal tract. Various forms of parasite that can be detected include cyst, trophozoite and larvae. One negative result does not rule out the possibility of parasitic infestation. Intermittent shedding of parasites warrants examinations of multiple specimens tested on consecutive days. Stool specimens for parasitic examination should be collected before initiation of antidiarrheal therapy or antiparasitic therapy. This test does not detect presence of opportunistic parasites like Cyclospora, Cryptosporidia and Isospora species. Examination of Ova and Parasite has been carried out by direct and concentration techniques.

Abhishek Sharma
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Mucus	Mucus is a protective layer that lubricates, protects & reduces damage due to bacteria or viruses.
Charcot-Leyden crystal	Parasitic diseases.
Ova & cyst	Ova & cyst indicate parasitic infestation of intestine.
Frank blood	Bleeding in the rectum or colon.
Occult blood	Occult blood indicates upper GI bleeding.
Macrophages	Macrophages in stool are an indication of infection as they are protective cells.
Epithelial cells	Epithelial cells that normally line the body surface and internal organs show up in stool when there is inflammation or infection.
Fat	Increased fat in stool maybe seen in conditions like diarrhoea or malabsorption.
pH	Normal stool pH is slightly acidic to neutral. Breast-fed babies generally have an acidic stool.

ADDITIONAL STOOL TESTS :

- Stool Culture**:- This test is done to find cause of GI infection, make decision about best treatment for GI infection & to find out if treatment for GI infection worked.
- Fecal Calprotectin**: It is a marker of intestinal inflammation. This test is done to differentiate Inflammatory Bowel Disease (IBD) from Irritable Bowel Syndrome (IBS).
- Fecal Occult Blood Test(FOBT)**: This test is done to screen for colon cancer & to evaluate possible cause of unexplained anaemia.
- Clostridium Difficile Toxin Assay**: This test is strongly recommended in healthcare associated bloody or watery diarrhoea, due to overuse of broad spectrum antibiotics which alter the normal GI flora.
- Biofire (Film Array) GI PANEL**: In patients of Diarrhoea, Dysentery, Rice watery Stool, FDA approved, Biofire Film Array Test,(Real Time Multiplex PCR) is strongly recommended as it identifies organisms, bacteria, fungi, virus parasite and other opportunistic pathogens, Vibrio cholera infections only in 3 hours. Sensitivity 96% & Specificity 99%.
- Rota Virus Immunoassay**: This test is recommended in severe gastroenteritis in infants & children associated with watery diarrhoea, vomiting & abdominal cramps. Adults are also affected. It is highly contagious in nature.

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WEST DELHI
NEW DELHI 110030
8800465156**ACCESSION NO : 0251WJ001893****PATIENT ID : VEERM211065251****CLIENT PATIENT ID : 012310210052****ABHA NO :****AGE/SEX : 58 Years Male****DRAWN : 21/10/2023 13:02:00****RECEIVED : 21/10/2023 13:17:58****REPORTED : 05/11/2023 15:20:38****Test Report Status Final****Results****Biological Reference Interval Units****SPECIALISED CHEMISTRY - HORMONE****MEDI WHEEL FULL BODY HEALTH CHECK UP ABOVE 40 MALE****THYROID PANEL, SERUM**

T3 METHOD : CHEMILUMINESCENCE	101.13	60.0 - 181.0	ng/dL
T4 METHOD : CHEMILUMINESCENCE	10.10	4.5 - 10.9	µg/dL
TSH (ULTRASENSITIVE) METHOD : CHEMILUMINESCENCE	3.259	0.550 - 4.780	µIU/mL

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.

Below 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

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C/O Aakriti Labs Pvt Ltd, 3, Mahatma Gandhi Marg, Gandhi Nagar Mod, Tonk Road
Jaipur, 302015
Rajasthan, India**Patient Ref. No. 775000005181322**



MC-5726

PATIENT NAME : VEER SINGH JATAV**REF. DOCTOR : SELF****CODE/NAME & ADDRESS : C000138404**ARCOFEMI HEALTHCARE LTD (MEDIWHEEL
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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
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. Guidelines of the American Thyroid association during pregnancy and Postpartum, 2011.

NOTE: It is advisable to detect Free T3, Free T4 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******Please visit www.agilusdiagnostics.com for related Test Information for this accession****Dr. Akansha Jain**
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