

Patient Name	Centre
Age/Gender	OP/IP No/UHID
MaxID/Lab ID	Collection Date/Time
Ref Doctor	Reporting Date/Time

**Hematology**
**Wellwise Platinum Profile**


SIN No:B2B5138603

**Complete Haemogram, Peripheral Smear and ESR,EDTA**

Date		Unit	Bio Ref Interval
	<b>04/Apr/2024</b>		
	<b>08:10AM</b>		
Haemoglobin	<b>8.3</b>	g/dl	12.0 - 15.0
Modified cyanmethemoglobin			
Packed Cell, Volume	<b>28.5</b>	%	40-50
Calculated			
Total Leucocyte Count (TLC)	10.0	10~9/L	4.0-10.0
Electrical Impedance			
RBC Count	4.75	10~12/L	3.8-4.8
Electrical Impedance			
MCV	<b>60.0</b>	fL	83-101
Electrical Impedance			
MCH	<b>17.4</b>	pg	27-32
Calculated			
MCHC	<b>29.0</b>	g/dl	31.5-34.5
Calculated			
Platelet Count	160	10~9/L	150-410
Electrical Impedance			
MPV	10.7	fL	7.8-11.2
Calculated			
RDW	<b>20.8</b>	%	11.5-14.5
Calculated			

**Differential Cell Count**

VCS / Light Microscopy

Neutrophils	68	%	40-80
Lymphocytes	20	%	20-40
Monocytes	06	%	2-10
Eosinophils	06	%	1-6

**Absolute Leukocyte Count**

Calculated from TLC &amp; DLC

Absolute Neutrophil Count	6.8	10~9/L	2.0-7.0
Absolute Lymphocyte Count	2.0	10~9/L	1.0-3.0
Absolute Monocyte Count	0.6	10~9/L	0.2-1.0
Absolute Eosinophil Count	<b>0.6</b>	10~9/L	0.02-0.5
ESR (Modified Westergren)	12	mm/hr	<=19

**Peripheral Smear Examination**

**RBC-** Anisocytosis (++) Microcytes (+++) Hypochromia (+++)  
**WBC-** Counts within normal limit

Test Performed at :1060 - Max Hospital Shalimar Bagh, Max Lab

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## Laboratory Investigation Report

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## Hematology



SIN No:B2B5138603

## Wellwise Platinum Profile

**Platelet-** Adequate**Impression-** Microcytic hypochromic anaemia**Advise-** Serum Iron, Serum Ferritin and TIBC

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*

Dr. Pooja Bhasin M.D.  
Associate Director & HOD  
Lab Service Pathology

Dr. Vijay Laxmi Sharma, MD  
Associate Director & Quality Manager

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**Laboratory Investigation Report**

Patient Name	Centre
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**Clinical Biochemistry**  
**Wellwise Platinum Profile**


SIN No:B2B5138603

**Fasting Blood Sugar (Glucose) , (FBS), Fluoride Plasma**

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
Glucose (Fasting)	102	mg/dL	74 - 99

**HbA1c (Glycated/ Glycosylated Hemoglobin) Test**  
 HPLC

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
Glycosylated Haemoglobin(Hb A1c)	5.0	%	< 5.7
Glycosylated Haemoglobin(Hb A1c) IFCC	31.13	mmol/mol	< 39.0
Average Glucose Value For the Last 3 Months	96.8	mg/dL	
Average Glucose Value For the Last 3 Months IFCC	5.36	mmol/L	

**Interpretation** The following HbA1c ranges recommended by the American Diabetes Association(ADA) may be used as an aid in the diagnosis of diabetes mellitus.

HbA1C(NGSP %)	HbA1C(IFCC mmol/mol)	Suggested Diagnosis
≥6.5	≥48	Diabetic
5.7 - 6.4	39 - 47	Pre- Diabetic
< 5.7	< 39	Non - Diabetic

HbA1C provides a useful index of average glycaemia over the preceding 6-8 weeks.

It is suggested that HbA1c is measured every 6 months in stable patients, every 3 months in patients with unstable metabolic control and every month in pregnancy. Increased Glycated hemoglobin is a reflection of Hyperglycemia.

Kindly correlate with clinical findings

**\*\*\* End Of Report \*\*\***



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Lab Service Pathology



Dr. Vijay Laxmi Sharma, MD  
Associate Director & Quality Manager

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**Laboratory Investigation Report**

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**Immunoassay**


SIN No:B2B5138603

**Wellwise Platinum Profile**
**Thyroid Profile\*, Serum**

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
Free Triiodothyronine (FT3)	3.48	pg/mL	2.6 - 4.2
CLIA			
Free Thyroxine (FT4)	0.81	ng/dL	0.58 - 1.64
CLIA			
Thyroid Stimulating Hormone	<b>7.281</b>	μIU/mL	0.38 - 5.33
CLIA			

**Comment**

Parameter	Unit	Premature (28 - 36 weeks)	Cord Blood (> 37 weeks)	Upto 2 Month	1st Trimester	2nd Trimester	3rd Trimester
FT3	Pg/mL			0.15 - 3.91	2.4 - 5.6	2.11 - 3.83	1.96 - 3.38
FT4	ng/dl			1.7 - 4.0		0.7 - 2.0	0.5 - 1.6
TSH	uIU/ml	0.7 - 27.0	2.3 - 13.2	0.5 - 10	0.05 - 3.7	0.31 - 4.35	0.41 - 5.18

**Note :** TSH levels are subject to circadian variation, reaching peak levels between 2 – 4 am and at a minimum between 6 – 10 pm. The variation is of the order of 50% - 206 %, hence time of the day has influence on the measured serum TSH concentrations.

Kindly correlate with clinical findings

**\*\*\* End Of Report \*\*\***



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Associate Director & Quality Manager

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**Clinical Biochemistry**  
**Wellwise Platinum Profile**


SIN No:B2B5138603

**Kidney Function Test (KFT) Profile with Calcium, Uric Acid, Serum**

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
Urea Urease, UV	23.4	mg/dL	17.0 - 43.0
Blood Urea Nitrogen Urease, UV	10.93	mg/dL	7.9 - 20.0
Creatinine Alkaline picrate kinetic	<b>0.56</b>	mg/dL	0.6 - 1.1
eGFR by MDRD MDRD	113.24	ml/min/1.73 m <sup>2</sup>	
eGFR by CKD EPI 2021	109.06		
Bun/Creatinine Ratio Calculated	19.52	Ratio	12:1 - 20:1
Uric Acid Uricase, Colorimetric	4.5	mg/dL	2.6 - 6.0
Calcium (Total) Arsenazo III	9.51	mg/dL	8.8 - 10.6
Sodium ISE indirect	139.9	mmol/L	136 - 146
Potassium ISE indirect	4.58	mmol/L	3.5 - 5.1
Chloride ISE indirect	105.7	mmol/L	101 - 109
Bicarbonate Enzymatic	27.0	mmol/L	21 - 31

**Ref. Range** eGFR - Estimated Glomerular Filtration Rate is calculated by MDRD equation which is most accurate for GFRs  $\leq$  60ml / min /1.73 m<sup>2</sup>.MDRD equation is used for adult population only.

<60ml / min / 1.73 m<sup>2</sup> - Chronic Kidney Disease

<15 ml / min /1.73 m<sup>2</sup> - Kidney failure

**BUN/Creatinine Ratio :-**

Increased in reduced renal perfusion (e.g. dehydration, Hypovolemic shock, Congestive Heart Failure) or Obstructive uropathy. Decreased in Acute Renal Tubular necrosis.

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Clinical Biochemistry  
Wellwise Platinum Profile

SIN No:B2B5138603

## Inorganic Phosphorus, Serum

Date		Unit	Bio Ref Interval
	04/Apr/2024		
	08:10AM		
Phosphorus(inorg) Phosphomolybdate-UV	4.59	mg/dL	2.5 - 4.5

## Interpretation

Increased in Osteolytic metastatic bone tumors, myelogenous leukemia, sarcoidosis, milk-alkali syndrome, vitamin D intoxication, healing fractures, renal failure, hyperparathyroidism, PTH resistance (Pseudohypoparathyroidism) and diabetes mellitus with ketosis.

Decreased in Osteomalacia, steatorrhea, renal tubular acidosis, growth hormone deficiency, acute alcoholism, gram-negative bacterial septicemia, hypokalemia, familial hypophosphatemic rickets, Vitamin D deficiency, severe malnutrition, malabsorption, secondary diarrhea, vomiting, nasogastric suction, primary hyperthyroidism and PTH producing tumors.

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**Clinical Biochemistry**  
**Wellwise Platinum Profile**


SIN No:B2B5138603

**Liver Function Test (LFT), Serum**

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
Total Protein	7.80	g/dL	6.6 - 8.3
Biuret			
Albumin	4.4	g/dL	3.5 - 5.2
Bromoresol Green (BCG)			
Globulin	3.4	g/dl	2.3 - 3.5
Calculated			
A.G. ratio	1.3		1.2 - 1.5
Calculated			
Bilirubin (Total) DPD	0.81	mg/dL	0.3 - 1.2
Bilirubin (Direct) Diazotization	0.13	mg/dL	0.0 - 0.2
Bilirubin (Indirect) Calculated	0.68	mg/dL	0.1 - 1.0
SGOT- Aspartate Transaminase (AST) UV without P5P	27	U/L	< 50
SGPT- Alanine Transaminase (ALT) UV without P5P	25	U/L	< 35
AST/ALT Ratio Calculated	1.08	Ratio	
Alkaline Phosphatase PNPP, AMP Buffer	147	U/L	30 - 120
GGTP (Gamma GT), Serum Enzymatic Rate	37.0	U/L	7 - 50

**Interpretation AST/ALT Ratio :-**

In Case of deranged AST and/or ALT, the AST/ALT ratio is > 2.0 in alcoholic liver damage and < 2.0 in non – alcoholic liver damage



MC-2262

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**Clinical Biochemistry**  
**Wellwise Platinum Profile**


SIN No:B2B5138603

**Lipid Profile,Serum**

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
Cholesterol	154	mg/dL	< 200
Cholesterol oxidase, esterase, peroxidase			
HDL Cholesterol	47.9	mg/dL	> 40
Direct measure, immuno inhibition			
LDL Cholesterol	97	mg/dL	< 100
Direct measure			
Triglyceride	114.0	mg/dL	< 150
Enzymatic, end point			
VLDL Cholesterol	22.8	mg/dL	< 30
Calculated			
Total Cholesterol/HDL Ratio	3.2	..	0.0-4.9
Calculated			
Non-HDL Cholesterol	106.10	mg/dL	< 130
Calculated			
HDL/LDL	0.49	Ratio	0.3 - 0.4
Calculated			

**Interpretation**

Total Cholesterol	Desirable: < 200 mg/dL Borderline High: 200-239 mg/dL High ≥ 240 mg/dL	LDL-C	Optimal: < 100 mg/dL Near Optimal/ Above Optimal: 100-129 mg/dL Borderline High: 130-159 mg/dL High: 160-189 mg/dL Very High: ≥ 190 mg/dL
HDL-C	Low HDL: < 40 mg/dL High HDL: ≥ 60 mg/dL	Triglyceride	Normal: <150 mg/dL Borderline High: 150-199 mg/dL High: 200-499 mg/dL Very High: ≥ 500 mg/dL

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Clinical Biochemistry  
Wellwise Platinum Profile

SIN No:B2B5138603

## Creatine Kinase (CPK), Serum

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
Creatine Kinase (CPK)	73 NAC activated	U/L	= 171.0

## Interpretation

CK is elevated in most myopathies such as Duchenne-muscular dystrophy, in conditions associated with muscle necrosis such as rhabdomyolysis, in diseases of the CNS such as Reyes Syndrome where a 70 fold increase in CK activity indicates the severity of the encephalopathy. CK activity rises following myocardial damage. The diagnostic sensitivity and specificity of total CK estimation for the diagnosis of an MI can be improved by determining the rate of increase of CK on serial samples obtained on admission and at 4, 8 and 12 hours thereafter. A 50% incremental increase per hour over the time period differentiates between an acute MI and non-infarction with an overall efficiency of 94%.

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Clinical Biochemistry  
Wellwise Platinum Profile

SIN No:B2B5138603

Test Name	Result	Unit	Bio Ref Interval
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**High Sensitivity CRP (HS CRP), Serum**

C-Reactive Protein, High Sensitive Immuno-Turbidimetric Test(Latex)	5.385	mg/L
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Reference Values in the table given below are recommended cardiovascular risk groups, in primary prevention settings by AHA/CDC and NACB expert panel.

Risk Level	CRP hs (mg/L)	CRP hs (mg/dL)
Low	< 1.0	< 0.10
Average	1.0 - 3.0	0.10 - 0.30
High	> 3.0	>0.30

Increase in CRP levels is non – specific, and interpretation must be undertaken in comparison with previous Hs CRP values or other cardiac risk indicators (Cholesterol, HDL etc.) Single measurement may lead to an erroneous assessment of early cardiac inflammation.

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Clinical Biochemistry  
Wellwise Platinum Profile

SIN No:B2B5138603

## CRP- C- Reactive Protein\*, Serum

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
CRP	6.035	mg/L	

Latex Particle Immunoturbidimetric

**Interpretation** This helps in detecting neonatal septicemia, meningitis and useful to assess the activity of inflammatory diseases like rheumatoid arthritis. It is increased after myocardial infarction, stress, trauma, infection, inflammation, surgery, or neoplastic proliferation. The increase with inflammation occurs within 6 -12 hours and peaks at about 48 hours.

## Ref Range :

Mg/L	Mg/dL
< 5.0	< 0.5

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*

Dr. Pooja Bhasin M.D.  
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Lab Service Pathology

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**Laboratory Investigation Report**

Patient Name	Centre
Age/Gender	OP/IP No/UHID
MaxID/Lab ID	Collection Date/Time
Ref Doctor	Reporting Date/Time

**Clinical Biochemistry**  
**Wellwise Platinum Profile**


SIN No:B2B5138603

**Apolipoproteins A1 & B,Serum**

Immunoturbidimetric

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
Apolipoprotein (A) Immunoturbidimetric	137	mg/ dL	120-190
Apolipoprotein (B) Immunoturbidimetric	86	mg/dl	55 - 130
Apo B/ Apo A1 Ratio Calculated	0.63		0.35 - 0.98

**Rheumatoid Factor(Quantitative), Serum**

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
Rheumatoid Factor Immunoturbidimetric	5.3	IU/ mL	0-12

**Interpretation** Rheumatoid factor is found in rheumatoid arthritis, Sjögren's syndrome, Scleroderma, dermatomyositis, Waldenström's disease, sarcoidosis and SLE. 75% patients with rheumatoid arthritis have RF of IgM class. Highest titers of Rheumatoid arthritis are seen in severe, active, chronic disease with vasculitis and subcutaneous nodules

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*

**Dr. Poonam. S. Das, M.D.**  
Principal Director-  
Max Lab & Blood Bank Services

**Dr. Dilip Kumar M.D.**  
Associate Director &  
Manager Quality



**Dr. Rajeev Kumar, MD**  
Associate Consultant  
Biochemistry

Test Performed at :910 - Max Hospital - Saket M S S H, Press Enclave Road, Mandir Marg, Saket, New Delhi, Delhi 110017

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**Laboratory Investigation Report**

Patient Name	Centre
Age/Gender	OP/IP No/UHID
MaxID/Lab ID	Collection Date/Time
Ref Doctor	Reporting Date/Time

**Clinical Biochemistry**  
**Wellwise Platinum Profile**


SIN No:B2B5138603

**Total Iron Binding Capacity (TIBC)\*, Serum**

Date		Unit	Bio Ref Interval
	<b>04/Apr/2024</b>		
	<b>08:10AM</b>		
Iron	<b>14</b>	µg/dL	60 - 180
TPTZ- No deproteinization			
UIBC	<b>474.9</b>	µg/dL	155 - 355
Nitroso - PSAP			
Total Iron Binding Capacity Calculated	488.9	µg/dL	215 - 535
Transferrin Saturation Calculated	<b>2.86</b>	%	17 - 37

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*

Dr. Pooja Bhasin M.D.  
Associate Director & HOD  
Lab Service Pathology

Dr. Vijay Laxmi Sharma, MD  
Associate Director & Quality Manager

Test Performed at :1060 - Max Hospital Shalimar Bagh, Max Lab

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## Laboratory Investigation Report

Patient Name	Centre
Age/Gender	OP/IP No/UHID
MaxID/Lab ID	Collection Date/Time
Ref Doctor	Reporting Date/Time

Immunoassay



SIN No:B2B5138603

Wellwise Platinum Profile

**Ferritin\*, Serum**

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
Ferritin CLIA	3.0	ng/mL	11 - 306.8

**Comment** Ferritin is a large hollow spherical protein containing iron, concentration of which roughly reflects the body iron content in many individuals. Serum ferritin concentration is a sensitive indicator of iron deficiency. Serum Ferritin concentration is increased in many disorders like infection, inflammatory disorders like rheumatoid arthritis or renal disease; common liver conditions (e.g. alcoholism, viral hepatitis B or C); heart disease, cancer. In patients with these disorders who also have iron deficiency their serum ferritin concentrations are often normal. An increase in serum ferritin concentration occurs as a result of ferritin release due to liver cell injury of diverse causes. Serum ferritin is also increased in patients with iron overload of any cause. Serum transferrin saturation is a better screening test for early iron overload than serum ferritin.

Kindly correlate with clinical findings

**\*\*\* End Of Report \*\*\***

Dr. Pooja Bhasin M.D.  
Associate Director & HOD  
Lab Service Pathology

Dr. Vijay Laxmi Sharma, MD  
Associate Director & Quality Manager

Test Performed at :1060 - Max Hospital Shalimar Bagh, Max Lab

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### Laboratory Investigation Report

Patient Name	Centre
Age/Gender	OP/IP No/UHID
MaxID/Lab ID	Collection Date/Time
Ref Doctor	Reporting Date/Time

Clinical Biochemistry



SIN No:B2B5138603

Wellwise Platinum Profile

#### LDH (Lactate Dehydrogenase) Total , Serum\*

Date		Unit	Bio Ref Interval
	04/Apr/2024		
	08:10AM		
LDH	221	IU/L	98 - 192
Enzymatic			

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*

Dr. Pooja Bhasin M.D.  
Associate Director & HOD  
Lab Service Pathology

Dr. Vijay Laxmi Sharma, MD  
Associate Director & Quality Manager

Test Performed at :1060 - Max Hospital Shalimar Bagh, Max Lab

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**Laboratory Investigation Report**

Patient Name	Centre
Age/Gender	OP/IP No/UHID
MaxID/Lab ID	Collection Date/Time
Ref Doctor	Reporting Date/Time

**Immunoassay**


SIN No:B2B5138603

**Wellwise Platinum Profile**
**Vitamin B12 (Vit- B12), (Cyanocobalamin)\*, Serum**

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
Vitamin B12 CLIA	191.0	pg/mL	120 - 914

**Interpretation**
**Note:- Vitamin B12 (Cobalamin)**

Vitamin B12 is tested for patients with GIT disease, Neurological disease, psychiatric disturbances, malnutrition, alcohol abuse.

Increased in chronic renal failure, severe CHF.

Decreased in megaloblastic anemia.

**Advise:** CBC, peripheral smear, serum folate levels, intrinsic factor antibodies (IFA), bone marrow examination, if Vit B12 deficient.

**Vitamin D, 25 - Hydroxy Test (Vit. D3)\*, Serum**

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
25 Hydroxy, Vitamin D CLIA	10.73	ng/mL	30-100

**Ref Range**

Vitamin D Status	25 (OH) Vitamin D Concentration Range (ng/ml)
Sufficiency	30-100
Insufficiency	20-29
Deficiency	<20
Potential Toxicity	>100

**Interpretation**

Vitamin D toxicity can be due to

1. Use of high doses of vitamin D for prophylaxis or treatment
2. Taking vitamin D supplements with existing health problems such as kidney disease, liver disease, tuberculosis and hyperparathyroidism

Vitamin D deficiency can be due to:

1. Inadequate exposure to sunlight,
2. Diet deficient in vitamin D
3. Malabsorption

**Advise:** Serum calcium, phosphorus and PTH



## Laboratory Investigation Report

Patient Name	Centre
Age/Gender	OP/IP No/UHID
MaxID/Lab ID	Collection Date/Time
Ref Doctor	Reporting Date/Time

Immunoassay



SIN No:B2B5138603

Wellwise Platinum Profile

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*

Dr. Pooja Bhasin M.D.  
Associate Director & HOD  
Lab Service Pathology

Dr. Vijay Laxmi Sharma, MD  
Associate Director & Quality Manager

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**Laboratory Investigation Report**

Patient Name	Centre
Age/Gender	OP/IP No/UHID
MaxID/Lab ID	Collection Date/Time
Ref Doctor	Reporting Date/Time

**Serology**


SIN No:B2B5138603

**Wellwise Platinum Profile**

Test Name	Result	Unit	Bio Ref Interval
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**Hepatitis B Surface Antigen,Serum**

CLIA

HBsAg Negative

CLIA

HBsAg Test Value 0.14 S/CO  
CLIA
**Ref. Range**

Negative	< 0.90
Borderline	0.90 - 5.0
Positive	> 5.0

**Interpretation**

- This test is used to detect hepatitis B surface antigen (HBsAg) in serum sample based on VITROS immunometric immunoassay technique and aid in the laboratory diagnosis of HBV infection.
- Viral hepatitis is a major public health problem with an estimated 257 million persistent carriers of hepatitis B virus (HBV) worldwide. Infection with HBV results in a wide spectrum of acute and chronic liver diseases that may lead to cirrhosis and hepatocellular carcinoma.
- Transmission of HBV occurs by percutaneous exposure to blood products, needle stick injury, sexual contact and perinatally from HBV-infected mothers to baby.
- Hepatitis B surface antigen (HBsAg), derived from the viral envelope, is the first antigen to appear following infection.
- Positive results should be correlated with other potential laboratory abnormalities and clinical picture.
- A negative test result does not exclude the possibility of exposure to or infection with hepatitis B virus.
- Levels of HBsAg may be undetectable both in early infection and late after infection.
- In rare cases HBsAg tests do not detect certain HBV mutant strains.
- HBs Ag disappears with recovery from clinical disease in most patients, however, it persists for years in carriers.

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*

Dr. Shakti Jain, MD

Associate Director and HOD -Microbiology

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**Laboratory Investigation Report**

Patient Name	Centre
Age/Gender	OP/IP No/UHID
MaxID/Lab ID	Collection Date/Time
Ref Doctor	Reporting Date/Time

**Serology Special**


SIN No:B2B5138603

**Wellwise Platinum Profile**

Test Name	Result	Unit	Bio Ref Interval
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**Allergy Screen-PhadiaTop/Inf, Serum**

FEIA

Allergy Screen, Phadia Top Fluoroenzyme Immunoassay	0.10	PAU/L	< 0.34
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**Comment**

ImmunoCAPPhadiatop is an in vitro qualitative and semiquantitative assay for gradeddetermination of IgE antibodies specific to inhalant allergens in human serum or plasma. It is intended for in vitro diagnostic use as an aid in the clinical diagnosis of IgE mediated allergicdisorders in conjunction with other clinical findings, and is to be used in clinical laboratories. In patients suffering from extrinsic asthma, hay fever or atopic eczema, symptoms developimmediately after exposure to specific allergens. This immediate type of allergy is a functionof a special type of serum antibodies belonging to the IgE class of immunoglobulins.

Kindly correlate with clinical findings

**\*\*\* End Of Report \*\*\***

**Dr. Poonam S. Das, M.D.**  
Principal Director-  
Max Lab & Blood Bank Services

**Dr. Bansidhar Tarai, M.D.**  
Associate Director  
Microbiology & Molecular Diagnostics



**Dr. Sonu Kumari Aggrawal, MD**  
Consultant Microbiology



**Dr Nidhi Malik, MD**  
Consultant Microbiology

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**Laboratory Investigation Report**

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MaxID/Lab ID	Collection Date/Time
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**Clinical Pathology**
**Wellwise Platinum Profile**


SIN No:B2B5138603

**Urine Routine And Microscopy**

Date	04/Apr/2024 08:10AM	Unit	Bio Ref Interval
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**Macroscopy**

Colour	Pale Yellow	Pale Yellow
Visual Observation/ Automated		
PH	6.0	..
Double Indicator		5-6
Specific Gravity	1.020	1.015 - 1.025
pKa change		
Protein	Neg	Nil
Protein-error of indicators		
Glucose.	Neg	Nil
Enzyme Reaction		
Ketones	Neg	Nil
Acetoacetic Reaction		
Blood	Neg	Nil
Benzidine Reaction		
Bilirubin	Neg	Nil
Diazo Reaction		
Urobilinogen	Normal	Normal
Ehrlichs Reaction		

**Microscopy**

Red Blood Cells (RBC)	0	/HPF	Nil
Light Microscopy/Image capture microscopy			
White Blood Cells	1	/HPF	0.0-5.0
Light Microscopy/Image capture microscopy			
Squamous Epithelial Cells	1	/HPF	
Light Microscopy/Image capture microscopy			
Cast	Nil	/LPF	Nil
Light Microscopy/Image capture microscopy			
Crystals	Nil	..	Nil
Light Microscopy/Image capture microscopy			
Bacteria	Nil	/HPF	Nil
Light Microscopy/Image capture microscopy			

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## Laboratory Investigation Report

Patient Name	Centre
Age/Gender	OP/IP No/UHID
MaxID/Lab ID	Collection Date/Time
Ref Doctor	Reporting Date/Time

Clinical Pathology



SIN No:B2B5138603

Wellwise Platinum Profile

Kindly correlate with clinical findings

\*\*\* End Of Report \*\*\*

Dr. Pooja Bhasin M.D.  
Associate Director & HOD  
Lab Service Pathology

Dr. Vijay Laxmi Sharma, MD  
Associate Director & Quality Manager

Test Performed at :1060 - Max Hospital Shalimar Bagh, Max Lab

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