





Unit

Laboratory Investigation Report

RED CROSS HOSPITAL

Patient Name

OP/IP No/UHID
Age/Gender

MaxID/Lab ID

Ref Doctor

Reporting
Date/Time

BLOOD CENTRE

Bio Ref Interval

Max-Antenatal Profile (Basic 2)

Blood Grouping and RH Factor*, EDTA

Date 31/Jan/2023

11:29AM

Blood Group A POSITIVE

Haemagglutination

Kindly correlate with clinical findings

*** End Of Report ***

Dr. Renu Aggarwal (MBBS,MD)

Consultant Pathologist

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

Clinical Biochemistry

Max-Antenatal Profile (Basic 2)

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

Date 31/Jan/2023 Unit **Bio Ref Interval**

11:29AM

83.99 74 - 99 Glucose (Fasting) mg/dl

Hexokinase

Kindly correlate with clinical findings

*** End Of Report ***

Dr. Renu Aggarwal (MBBS,MD)

Consultant Pathologist







RED CROSS HOSPITAL

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Max-Antenatal Profile (Basic 2)

Complete Haemogram, Peripheral Smear and ESR,EDTA*

Date	31/Jan/2023 11:29AM	Unit	Bio Ref Interval
Haemoglobin Modified cyanmethemoglobin	10.19	g/dl	12.0 - 15.0
Packed Cell, Volume Calculated	31.1	%	40-50
Total Leucocyte Count (TLC) Electrical Impedance	4.76	10~9/L	4.0-10.0
RBC Count Electrical Impedance	4.28	10~12/L	3.8-4.8
MCV Electrical Impedance	72.7	fL	83-101
MCH Calculated	23.8	pg	27-32
MCHC Calculated	32.8	g/dl	31.5-34.5
Platelet Count Electrical Impedance	236	10~9/L	150-410
MPV Calculated	7.95	fl	7.8-11.2
RDW Calculated	17.4	%	11.5-14.5
Differential Cell Count VCS / Light Microscopy			
Neutrophils	60	%	40-80
Lymphocytes	30	%	20-40
Monocytes	07	%	2-10
Eosinophils	03	%	1-6
Absolute Leukocyte Count Calculated from TLC & DLC			
Absolute Neutrophil Count	2.86	10~9/L	2.0-7.0
Absolute Lymphocyte Count	1.4	10~9/L	1.0-3.0
Absolute Monocyte Count	0.33	10~9/L	0.2-1.0
Absolute Eosinophil Count	0.14	10~9/L	0.02-0.5
ESR (Westergren)	30	mm/hr	<=12
Peripheral Smear			

 $Test\ Performed\ at\ : 3954\ -\ MM\ Healthcare\ Ltd.,\ A-15\ Pushpanjali,\ Vikas\ Marg\ Extn,\ Delhi$

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Conditions of Reporting: 1. The tests are carried out in the lab with the presumption that the specimen belongs to the patient name as identified in the bill/test request form. 2. The test results relate specifically to the sample received in the lab and are presumed to have been generated and transported per specific instructions given by the physicians/laboratory. 3. The reported results are for the information and interpretation by the referring doctor only. 4. Some tests are referred to other laboratories to provide a wider test menu to the customer. 5. Max Healthcare shall in no event be liable for accidental damages loss, or destruction of specimen which is not attributable to any direct and maia fide act or omission of Max Healthcare or its employees. Liability of Max Healthcare for deficiency of services, or other errors and omissions shall be limited to fee paid by the patient for the relevant laboratory services.

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RED CROSS HOSPITAL

Patient Name

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Collection
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Ref Doctor

Reporting
Date/Time

Hematology

Max-Antenatal Profile (Basic 2)

Examination

RBC: - Normocytic Normochromic **WBC:** - Counts within normal limits

Platelet: - Adequate

Kindly correlate with clinical findings

*** End Of Report ***

Dr. Renu Aggarwal (MBBS,MD) Consultant Pathologist

Test Performed at :3954 - MM Healthcare Ltd., A-15 Pushpanjali, Vikas Marg Extn, Delhi
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 Patient Name
 Centre

 Age/Gender
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 MaxID/Lab ID
 Date/Time

 Ref Doctor
 Reporting

Date/Time Serology

SIN No P C0027207

Max-Antenatal Profile (Basic 2)

Popult Unit Die Def Interval

HCV IgG Antibody (Hepatitis C Virus)

CLIA

CLIA

Toot Name

HCV, IgG Negative

HCV,IgG Test Value

0.07

Ref. Range

Negative < 0.90Borderline 0.90 - 8.0Positive > 8.0

Interpretation

This test is a screening test performed on VITROS immunodiagnostic system using immunometric technique.

- 1. Hepatitis C (HCV) is an RNA virus of Flavivirus group transmitted via blood transfusions, transplantation, injection drug users, accidental needle punctures in healthcare workers, dialysis patients and rarely from mother to infant.
- 2. 10% of new cases show sexual transmission. As compared to HAV & HBV, chronic infection with HCV occurs in 85% of infected individuals.
- 3. This test is indicator of past or present infection, but does not differentiate between Acute / Chronic / Resolved infection .HCV RNA PCR recommended in all reactive results to differentiate between past and present infection
- 4. A definitive clinical diagnosis should not be made by result of a single test only, but should be made by taking clinical history and other laboratory findings in to account.

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Test Performed at: 794 - Max Hospital - Vaishali, W-3, Sector-1, Vaishali, Ghaziabad-201012, U.P.









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

Serology

SIN No P C0027207

Max-Antenatal Profile (Basic 2)

Popult Unit Pio Pof Interval

Hepatitis B Surface Antigen, Serum*

CLIA

Toot Name

HBsAg Negative

CLIA

Ref. Range

Negative < 0.90Borderline 0.90 - 5.0Positive > 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.

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RED CROSS HOSPITAL Centre Patient Name OP/IP No/UHID

Age/Gender Collection MaxID/Lab ID Date/Time

Ref Doctor Reporting Date/Time

Serology

Max-Antenatal Profile (Basic 2)

l Init Die Def Interval

HIV I & II, Serum

CLIA

Toot Name

HIV (I and II) Negative CLIA

Interpretation

1) A Negative result implies that no Anti HIV-1 or HIV-2 antibodies have been detected in the sample by this method. This means that either the patient has not been exposed to HIV-1 or HIV-2 infection or the sample has been taken during the "WINDOW PERIOD" (before the development of detectable levels of antibodies, 2 12 weeks after exposure).

2) Positive result suggests the possibility of HIV-1 or HIV-2 infection.

To rule out false positivity, false negativity and window period, kindly perform "Confirmatory Tests" like HIV I RNA (Qualitative) Real Time PCR.

Kindly correlate with clinical findings

*** End Of Report ***

Dr. Dimple Juneja, M.D. Consultant Pathologist

Dr. Renu Aggarwal (MBBS,MD)

Consultant Pathologist

Dr. Saloni Sehgal (MBBS, MD) Principal Consultant &

Head Microbiology & Infection Control

Dr. Neera Kaushik (M Phil) Senior Consultant Microbiology

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SIN No. P. C0027207

Clinical Pathology

Max-Antenatal Profile (Basic 2)

Urine Routine And Microscopy*

Date 31/Jan/2023 Unit **Bio Ref Interval**

11:29AM

Macroscopy

Pale Yellow Deep Yellow Colour

Visual Observation/ Automated

PΗ 5-6 6.5

Double Indicator

1.015 - 1.025 1.025 Specific Gravity

pKa change

Protein Nil Trace Protein-error of indicators

Nil Nil Glucose. Enzyme Reaction

Nil Nil Ketones Acetoacetic Reaction

ABSENT Nil Blood

Benzidine Reaction

Nil Nil Bilirubin Diazo Reaction

Urobilinogen Normal Normal Ehrlichs Reaction

Nitrite

Negative Conversion of Nitrate

Microscopy

Red Blood Cells (RBC) Nil /HPF Nil

Light Microscopy/Image capture microscopy

White Blood Cells /HPF 0.0-5.0 3-4

Light Microscopy/Image capture

microscopy

/HPF Squamous Epithelial Cells 6-8

Light Microscopy/Image capture

microscopy

Cast Nil /LPF Nil

Light Microscopy/Image capture microscopy

Crystals Nil

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Nil

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Date/Time

 RED CROSS HOSPITAL
 Centre

 Patient Name
 OP/IP No/UHID

 Age/Gender
 Collection

 MaxID/Lab ID
 Date/Time

 Ref Doctor
 Reporting

Clinical Pathology

Nil

/HPF

Max-Antenatal Profile (Basic 2)

Light Microscopy/Image capture microscopy

Bacteria Nil

Light Microscopy/Image capture

microscopy

Kindly correlate with clinical findings

*** End Of Report ***

Dr. Renu Aggarwal (MBBS,MD)

Consultant Pathologist







Centre

OP/IP No/UHID

Collection Date/Time

Reporting

Date/Time

Serology Max-Antenatal Profile (Basic 2)

> Hois Die Def Interval

VDRL/RPR*, Serum

Patient Name

Age/Gender

MaxID/Lab ID

Ref Doctor

Toot Name

Non Reactive RPR(Syphilis) Slide Flocculation

Comment

Interpretation

- 1. It is a screening test for syphilis which is useful for following the progression of disease and response to therapy. Rising titers are of immense value in confirming the diagnosis.
- 2. Biological false positive reactions exhibit low titers and are seen in conditions like Viral fevers, Mycoplasma infection, Chlamydia infection, Malaria, Immunizations, Pregnancy, Autoimmune disorders & past history of Treponemal infection.
- 3. It is advisable to confirm the diagnosis by tests such as TPHA & FTA-ABS.
- 4. A definitive clinical diagnosis should not be made by result of a single test only, but should be made by taking clinical history and other laboratory findings in to account.

Kindly correlate with clinical findings

*** End Of Report ***

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Dr. Renu Aggarwal (MBBS,MD)

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Dr. Saloni Sehgal (MBBS, MD) **Principal Consultant &**

Head Microbiology & Infection Control

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 Patient Name
 Centre

 Age/Gender
 OP/IP No/UHID

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 Date/Time

 Ref Doctor
 Reporting

 Date/Time

Immunoassay

Max-Antenatal Profile (Basic 2)

Thyroid Profile*, Serum*

Date	31/Jan/2023 11:29AM	Unit	Bio Ref Interval
Free Triiodothyronine (FT3)	3.96	pg/mL	2.6 - 4.2
Free Thyroxine (FT4) CLIA	1.21	ng/dL	0.58 - 1.64
Thyroid Stimulating Hormone CLIA	0.203	μIU/mL	0.34 - 5.6

Comment

Parameter	Unit	Premature (28 - 36 weeks)	(> 37 weeks)	Upto 2 Month	Adult	1st Trimester	2nd Trimester	3rd Trimester
FT3	Pg/mL		0.15 - 3.91	2.4 - 5.6	2.6 - 4.2	2.11 - 3.83	1.96 - 3.38	1.96 - 3.38
TSH	uIU/ml	0.7 - 27.0	2.3 - 13.2	0.5 - 10	0.38 - 5.33	0.1 - 2.5	0.2 - 3.0	0.3 - 3.0

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 ***

Dr. Dimple Juneja, M.D. Consultant Pathologist

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

	Hematology Special			
	Max-Antenatal Profile (Bas	ic 2)	SIIV IVO.RCO02	.7207
Toot Name	Popult	l Init	Dia Daf Intanual	
Haemoglobin HPLC,EDTA				
Lab Number	1326/2023			
Hb F	0.4	%	0.0-1.0	
P2	3.9			
P3	4.6	%	<6.0	
HbA0 (non-glycated)	87.2	%	80-90	
Hb A2	2.7	%	2.0 - 3.5	
RBC Indices				
Haemoglobin. Non- Haemoglobincyanide	10.1	gm/dl	12-15	
RBC Count.	4.45	10~12/L	4.5-5.5	

fl

pg

83-101

27-32

11.5-14.5

Interpretation (HPLC)

Electrical Impedance

Electrical Impedance

MCV

MCH.

Calculated RDW.

Calculated

Normal Chromatogram

Recommend repeat Hemogram and HPLC after replenishing iron stores.

Please refer to comment.

Comment

(Syn: - High Performance Liquid Chromatography, Hb HPLC)

Hb HPLC is used for screening of Beta Thalassemia and detection of many other haemoglobin variants (Hemoglobin E, D Punjab, Sickle cell anaemia etc.) along with their concentration to establish zygosity.

Normal HbA2 values can be seen in Alpha Thalassemia trait, silent/ atypical Beta Thalassemia and Beta Thalassemia with iron deficiency.

A repeat Hb HPLC should be performed after replenishing iron stores in cases with microcytic hypochromic indices.

73.5

22.7

15.9

Advise: -

If there is absence of iron deficiency & persistence of microcytic hypochromic indices, then the possibility of Alpha Thalassemia trait or silent/ atypical Beta Thalassemia trait cannot be ruled out. Hb HPLC of parents and molecular sequencing of globin genes are recommended in these cases.

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Patient Name

Age/Gender





Laboratory Investigation Report

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In individuals with beta thalassemia trait and carriers of haemoglobin variants –

Hematology Special

Max-Antenatal Profile (Basic 2)

Die Def Interval

1. Familial study is recommended.

- 2. In cases of young/ pregnant females Hb HPLC testing of spouse is strongly recommended.
- 3. Genetic counselling

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. Nitin Dayal, M.D. Principal Consultant & Head, Haematopathology

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