

Patient Name	: Mr. MOHIT BAJPAI	Age/Sex	: 44 Year(s) / Male
UHID	: SHHM.97245	Order Date	: 15/06/2024 10:55
Episode	: OP		
Ref. Doctor	: self	Mobile No	: 9702031980
		DOB	: 24/01/1980
		Facility	: SEVENHILLS HOSPITAL, MUMBAI

Blood Bank

Test Name		Result				
Sample No: 00338299	A Collection Date :	15/06/24 10:56	Ack Date :	15/06/2024 11:36	Report Date :	15/06/24 12:04
BLOOD GROUPING	/ CROSS-MATCHING	BY SEMI AUTOM	ATION			
BLOOD GROUP (ABO))	'0'	ı			
Rh Type Method - Column Agglutina	ation	POS	SITIVE			
						-

REMARK: THE REPORTED RESULTS PERTAIN TO THE SAMPLE RECEIVED AT THE BLOOD CENTRE.

Interpretation:

Blood typing is used to determine an individual's blood group, to establish whether a person is blood group A, B, AB, or O and whether he or she is Rh positive or Rh negative. Blood typing has the following significance,

• Ensure compatibility between the blood type of a person who requires a transfusion of blood or blood components and the ABO and Rh type of the unit of blood that will be transfused.

• Determine compatibility between a pregnant woman and her developing baby (fetus). Rh typing is especially important during pregnancy because a mother and her fetus could be incompatible.

• Determine the blood group of potential blood donors at a collection facility.

• Determine the blood group of potential donors and recipients of organs, tissues, or bone marrow, as part of a workup for a transplant procedure.

End of Report

Dr.Pooja Vinod Mishra MD Pathology Jr Consultant Pathologist, MMC Reg No. 2017052191 RegNo: 2017/05/2191



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Biochemistry

Test Name			Resu	lt	Unit	Bio	logical Reference Interval
Sample No :	O0338299B	Collection Date :	15/06/24 10	:56 Ack Date :	15/06/2024 11:12	Report Date :	15/06/24 14:21
Blood Sug	ar FBS						
FBS Method - Hexc	okinase			79.86		mg/dl	70 - 100
GLUCOSE-	PLASMA POST	T PRANDIAL					
Glucose,Pos	st Prandial			94.5		mg/dl	70 - 140

American Diabetes Association Reference Range :

FASTING:-Normal : < 100 mg/dl Impaired fasting glucose(Prediabetes) : 100 - 126 mg/dl Diabetes : >= 126 mg/dl

Post-Prandial Blood Glucose: Non- Diabetic: Up to 140mg/dL Pre-Diabetic: 140-199 mg/dL Diabetic :>200 mg/dL

References:

Pack Insert of Bio system
 Tietz Textbook Of Clinical Chemistry And Molecular Diagnostics, 6th Ed, Editors: Rifai et al. 2018

Interpretation :-

Conditions that can result in an elevated blood glucose level include: Acromegaly, Acute stress (response to trauma, heart attack, and stroke for instance), Chronic kidney disease, Cushing syndrome, Excessive consumption of food, Hyperthyroidism, Pancreatitis.

A low level of glucose may indicate hypoglycemia, a condition characterized by a drop in blood glucose to a level where first it causes nervous system symptoms (sweating, palpitations, hunger, trembling, and anxiety), then begins to affect the brain (causing confusion, hallucinations, blurred vision, and sometimes even coma and death). A low blood glucose level (hypoglycemia) may be



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seen with:Adrenal insufficiency, Drinking excessive alcohol, Severe liver disease, Hypopituitarism, Hypothyroidism, Severe infections, Severe heart failure, Chronic kidney (renal) failure, Insulin overdose, Tumors that produce insulin (insulinomas), Starvation.

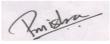
<u>ALT(SGPT) - SERUM</u>					
SGPT (Alanine Transaminase) - SERUM Method - IFCC	16.19	IU/L	0 - 45		
References : 1)Pack Insert of Bio system 2) Tietz Textbook Of Clinical Chemistry And Molecular Diagnostics, 6th Ed, Editors: Rifai et al. 2018					
Total Bilirubin - SERUM Method - Diazo	0.36	mg/dl	0 - 2		
Direct Bilirubin SERUM Method - Diazotization	0.25	mg/dl	0 - 0.4		
Indirect Bilirubin - Calculated Method - Calculated	0.11	mg/dl			
BUN-SERUM					
BUN - SERUM	23.94 ▲ (H)	mg/dl	4 - 18		

1)Pack Insert of Bio system

2) Tietz Textbook Of Clinical Chemistry And Molecular Diagnostics, 6th Ed, Editors: Rifai et al. 2018



Dr.Ritesh Kharche MD, PGD-HM Consultant Pathologist and Director of Laboratory Services RegNo: 2006/03/1680



Dr.Pooja Vinod Mishra MD Pathology Jr Consultant Pathologist, MMC Reg No. 2017052191 RegNo: 2017/05/2191



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HAEMATOLOGY

st Name			Result		Unit	Bio	ological Reference Interva
Sample No :	O0338299A	Collection Date :	15/06/24 10:56	Ack Date :	15/06/2024 11:12	Report Date :	15/06/24 12:00
COMPLETI	E BLOOD COUN	T (CBC) - EDTA	WHOLE BLOOD				
Total WBC	Count		7.02			x10^3/ul	4.00 - 10.00
Neutrophils			60.9			%	40.00 - 80.00
Lymphocyte	es		32.0			%	20.00 - 40.00
Eosinophils			2.8			%	1.00 - 6.00
Monocytes			4.2			%	2.00 - 10.00
Basophils			0.1	▼ (L)		%	1.00 - 2.00
Absolute Ne	eutrophil Count		4.28			x10^3/ul	2.00 - 7.00
Absolute Ly	mphocyte Count		2.25			x10^3/ul	0.80 - 4.00
Absolute Ec	osinophil Count		0.19			x10^3/ul	0.02 - 0.50
Absolute Mo	onocyte Count		0.30			x10^3/ul	0.12 - 1.20
Absolute Ba	asophil Count		0.00			x10^3/ul	0.00 - 0.10
RBCs				▼ (L)		x10^6/ul	4.50 - 5.50
Hemoglobir	1		13.2	.,		gm/dl	13.00 - 17.00
Hematocrit			40.4			%	40.00 - 50.00
MCV			90.6			fl	83.00 - 101.00
МСН			29.5			pg	27.00 - 32.00



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МСНС	32.6		gm/dl	31.50 - 34.50
RED CELL DISTRIBUTION WIDTH-CV (I	RDW-CV) 12.8		%	11.00 - 16.00
RED CELL DISTRIBUTION WIDTH-SD (I	RDW-SD) 43.1		fl	35.00 - 56.00
Platelet	341		x10^3/ul	150.00 - 410.00
Mean Platelet Volume (MPV)	8.8		fl	6.78 - 13.46
PLATELET DISTRIBUTION WIDTH (PDV	/) 15.8		%	9.00 - 17.00
PLATELETCRIT (PCT)	0.301 ▲ (H)		%	0.11 - 0.28

Method:-

HB Colorimetric Method. RBC/PLT Electrical Impedance Method. WBC data Flow Cytometry by Laser Method. MCV,MCH,MCHC,RDW and rest parameters - Calculated. All Abnormal Haemograms are reviewed confirmed microscopically.

NOTE: Wallach's Interpretation of Diagnostic Tests. 11th Ed, Editors: Rao LV. 2021

NOTE :-

The International Council for Standardization in Haematology (ICSH) recommends reporting of absolute counts of various WBC subsets for clinical decision making. This test has been performed on a fully automated 5 part differential cell counter which counts over 10,000 WBCs to derive differential counts. A complete blood count is a blood panel that gives information about the cells in a patient's blood, such as the cell count for each cell type and the concentrations of Hemoglobin and platelets. The cells that circulate in the bloodstream are generally divided into three types: white blood cells (leukocytes), red blood cells (erythrocytes), and platelets (thrombocytes). Abnormally high or low counts may be physiological or may indicate disease conditions, and hence need to be interpreted clinically.

End of Report



Dr.Ritesh Kharche MD, PGD-HM



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Consultant Pathologist and Director of Laboratory Services RegNo: 2006/03/1680





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HAEMATOLOGY

Test Name			Resu	lt	Unit	Bio	logical Reference Interval
Sample No :	O0338299A	Collection Date :	15/06/24 10	:56 Ack Date :	15/06/2024 11:12	Report Date :	15/06/24 13:07
ERYTHROC	CYTE SEDIMEN	<u>TATION RATE (E</u>	<u>SR)</u>				
ESR				35 ▲ (H)		mm/hr	0 - 20

Method: Westergren Method

INTERPRETATION :-

ESR is a non-specific phenomenon, its measurement is clinically useful in disorders associated with an increased production of acute-phase proteins. It provides an index of progress of the disease in rheumatoid arthritis or tuberculosis, and it is of considerable value in diagnosis of temporal arteritis and polymyalgia rheumatica. It is often used if multiple myeloma is suspected, but when the myeloma is non-secretory or light chain, a normal ESR does not exclude this diagnosis.

An elevated ESR may occur as an early feature in myocardial infarction. Although a normal ESR cannot be taken to exclude the presence of organic disease, the vast majority of acute or chronic infections and most neoplastic and degenerative diseases are associated with changes in the plasma proteins that increased ESR values.

The ESR is influenced by age, stage of the menstrual cycle and medications taken (corticosteroids, contraceptive pills). It is especially low (0–1 mm) in polycythaemia, hypofibrinogenaemia and congestive cardiac failure and when there are abnormalities of the red cells such as poikilocytosis, spherocytosis, or sickle cells. In cases of performance enhancing drug intake by athletes the ESR values are generally lower than the usual value for the individual and as a result of the increase in haemoglobin (i.e. the effect of secondary polycythaemia).

- End of Report -

Dr.Ritesh Kharche MD, PGD-HM Consultant Pathologist and Director of Laboratory Services RegNo: 2006/03/1680

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Biochemistry

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Sample No :	O0338299B	Collection Date :	15/06/24 10	:56 Ack Date :	15/06/2024 11:12	Report Date :	15/06/24 14:21
Blood Sug	ar FBS						
FBS Method - Hexc	okinase			79.86		mg/dl	70 - 100
GLUCOSE-	PLASMA POST	T PRANDIAL					
Glucose,Pos	st Prandial			94.5		mg/dl	70 - 140

American Diabetes Association Reference Range :

FASTING:-Normal : < 100 mg/dl Impaired fasting glucose(Prediabetes) : 100 - 126 mg/dl Diabetes : >= 126 mg/dl

Post-Prandial Blood Glucose: Non- Diabetic: Up to 140mg/dL Pre-Diabetic: 140-199 mg/dL Diabetic :>200 mg/dL

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Interpretation :-

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A low level of glucose may indicate hypoglycemia, a condition characterized by a drop in blood glucose to a level where first it causes nervous system symptoms (sweating, palpitations, hunger, trembling, and anxiety), then begins to affect the brain (causing confusion, hallucinations, blurred vision, and sometimes even coma and death). A low blood glucose level (hypoglycemia) may be



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seen with:Adrenal insufficiency, Drinking excessive alcohol, Severe liver disease, Hypopituitarism, Hypothyroidism, Severe infections, Severe heart failure, Chronic kidney (renal) failure, Insulin overdose, Tumors that produce insulin (insulinomas), Starvation.

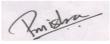
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BUN - SERUM	23.94 ▲ (H)	mg/dl	4 - 18

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Dr.Pooja Vinod Mishra MD Pathology Jr Consultant Pathologist, MMC Reg No. 2017052191 RegNo: 2017/05/2191



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Urinalysis

Test Name		Resul	t	Unit	Bio	logical Reference Interval
Sample No: 00338299D	Collection Date :	15/06/24 10:	S6 Ack Date :	15/06/2024 11:13	Report Date :	15/06/24 13:37
Physical Examination						
QUANTITY			30		ml	
Colour			Pale Yellow			
Appearance			Clear			
DEPOSIT			Absent			Absent
рН			Acidic			
Specific Gravity			1.015			
Chemical Examination						
Protein			Absent			Absent
Glucose			Absent			Absent
ketones			POSITIVE (+)			Absent
Blood			NEGATIVE			Negative
Bilirubin			Negative			
Urobilinogen			normal			Normal
NITRATE			Absent			Absent
LEUKOCYTES			Absent			Absent
Microscopic Examination						

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Pus cells		2-3		/HPF			
Epithelial Cells		3-4		/HPF			
RBC		absent		/HPF	Absent		
Cast		absent		/LPF	Absent		
Crystal		absent		/HPF	Absent		
Amorphous Mat	terials	Absent			Absent		
Yeast		Absent			Absent		
Bacteria		Absent			Absent		

End of Report



Dr.Nipa Dhorda MD Pathologist RegNo: 91821



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UHID Ref. Doctor	: self	Facility	: SEVENHILLS HOSPITAL,
Address	 601, FLAMINGO APARTMENTS, 14TH ROAD, khar west, Mumbai, Maharastra, 400052 	Mobile	MUMBAI : 9702031980

DIAGNOSTICS REPORT

X RAY CHEST PA VIEW

Both lungs are clear.

The frontal cardiac dimensions are normal.

The pleural spaces are clear.

Both hilar shadows are normal in position and density.

No diaphragmatic abnormality is seen.

The soft tissues and bony thorax are normal.

IMPRESSION: No pleuroparenchymal lesion is seen.

Kula

Dr.Bhujang Pai MBBS,MD

Consultant RegNo: 49380