



BML519909

Laboratory Investigation Report

Name	: Mr. AZAD HOSSAIN PATWARY	Ref No.	: 40616
DOB	: 27/04/1987	Sample No.	: 2504557479
Age / Gender	: 37 Y / Male	Collected	: 01/04/2025 16:45
Referred by	: DR.AMAIZAH ISHTIAQ	Registered	: 02/04/2025 09:56
Centre	: CITICARE MEDICAL CENTER	Reported	: 02/04/2025 11:49

BIOCHEMISTRY

Test	Result	Flag	Unit	Reference Range	Methodology
URIC ACID (SERUM)	7.3	H	mg/dL	3.4 - 7.0 Please note change. Source: Roche IFU.	Enzymatic colorimetric assay
CREATININE (SERUM)	12	CH	mg/dL	0.74 - 1.35	Kinetic colorimetric assay based on Jaffe method

Comments : Please correlate clinically.

INTERPRETATION NOTES :

1. Creatinine measurements are used as an aid in diagnosis and monitoring of renal disorders, Chronic Kidney disease (CKD) and in monitoring of renal dialysis and also used for the calculation of the fractional excretion of other urine analytes (e. g., albumin, α -amylase).
2. Creatinine is a break-down product of creatine phosphate in muscle, and is produced at a fairly constant rate by the body (depending on muscle mass). It is freely filtered by the glomeruli and, under normal conditions, is not reabsorbed by the tubules to any appreciable extent. A small but significant amount is also actively secreted. Its concentration is thus, inversely related to glomerular filtration rate (GFR).
3. Physiological factors affecting serum creatinine concentration include age, gender, race, muscularity, exercise, pregnancy, certain drugs, diet, dehydration and nutritional status.
4. Low serum Creatinine levels is seen in cases of low muscle mass like muscular atrophy, or aging.
5. High serum creatinine levels is seen in Acute and Chronic kidney disease, obstruction.
6. Since a rise in blood creatinine is observed only with marked damage of the nephrons, it is not suited to detect early stage kidney disease.

UREA (SERUM)	196	CH	mg/dL	12.86 - 42.86 Please note change. Source: Roche IFU	Kinetic test with urease and glutamate dehydrogenase
--------------	-----	----	-------	---	--

Comments : Please correlate clinically.

POTASSIUM (K)	5.5	H	mmol/L	3.5 - 5.1 Please note change. Source: Roche IFU.	ISE (Indirect)
---------------	-----	---	--------	--	----------------

Sample Type : Serum

End of Report

Dr. Adley Mark Fernandes
M.D (Pathology)
Pathologist

Gyoma V. Shah
Dr. Vyoma V Shah
M.D (Pathology)
Clinical Pathologist

ANJUMOL D V
ANJUMOL D V
Laboratory Technologist

This is an electronically authenticated report

Page 1 of 3

Printed on: 02/04/2025 12:39

Test result pertains only to the sample tested and to be interpreted in the light of clinical history. These tests are accredited under ISO 15189:2012 unless specified by (^). Test marked with # is performed in an accredited referral laboratory.





BML519909

Laboratory Investigation Report

Name : Mr. AZAD HOSSAIN PATWARY
DOB : 27/04/1987
Age / Gender : 37 Y / Male
Referred by : DR.AMAIZAH ISHTIAQ
Centre : CITICARE MEDICAL CENTER

Ref No. : 40616
Sample No. : 2504557479
Collected : 01/04/2025 16:45
Registered : 02/04/2025 09:56
Reported : 02/04/2025 12:37

HEMATOLOGY

Test	Result	Flag	Unit	Reference Range	Methodology
COMPLETE BLOOD COUNT (CBC)					
HEMOGLOBIN	8.4	L	g/dL	13.5 - 17.5	Photometric
RBC COUNT	2.8	L	10 ⁶ /μL	4.3 - 5.7	Electrical Impedance
HEMATOCRIT	24.3	L	%	38 - 50	Calculation
MCV	84.4		fL	82 - 98	Calculation
MCH	29.2		pg	27 - 32	Calculation
MCHC	34.6		g/dL	32 - 37	Calculation
RDW	15.1		%	11.8 - 15.6	Calculation
RDW-SD	45.1		fL		Calculation
MPV	10.0		fL	7.6 - 10.8	Calculation
PLATELET COUNT	177		10 ³ /uL	150 - 450	Electrical Impedance
PCT	0.1		%	0.01 - 9.99	Calculation
PDW	17.5		Not Applicable	0.1 - 99.9	Calculation
NUCLEATED RBC (NRBC)^	0.5		/100 WBC		VCS 360 Technology
ABSOLUTE NRBC COUNT^	0.03		10 ³ /uL		Calculation
EARLY GRANULOCYTE COUNT (EGC)^	0.3		%		VCS 360 Technology
ABSOLUTE EGC^	0.0		10 ³ /uL		Calculation
WBC COUNT	6.4		10 ³ /μL	4 - 11	Electrical Impedance
DIFFERENTIAL COUNT (DC)					
NEUTROPHILS	66		%	40 - 75	VCS 360 Technology
LYMPHOCYTES	22		%	20 - 45	VCS 360 Technology
EOSINOPHILS	7	H	%	0 - 6	VCS 360 Technology
MONOCYTES	5		%	1 - 6	VCS 360 Technology
BASOPHILS	0		%	0 - 1	VCS 360 Technology
ABSOLUTE COUNT					
ABSOLUTE NEUTROPHIL COUNT	4.3		10 ³ /uL	1.6 - 8.25	Calculation
ABSOLUTE LYMPHOCYTE COUNT	1.4		10 ³ /uL	0.8 - 4.95	Calculation
ABSOLUTE MONOCYTE COUNT	0.3		10 ³ /uL	0.04 - 0.66	Calculation
ABSOLUTE EOSINOPHIL COUNT	0.5		10 ³ /uL	0 - 0.66	Calculation
ABSOLUTE BASOPHIL COUNT	0.0		10 ³ /uL	0 - 0.11	Calculation

Dr. Adley Mark Fernandes
M.D (Pathology)
Pathologist

Gyoma V. Shah
Dr. Vyoma V Shah
M.D (Pathology)
Clinical Pathologist

Jillian Joy Garcia
Jillian Joy Garcia
Laboratory Technologist

This is an electronically authenticated report

Page 2 of 3

Printed on: 02/04/2025 12:39

Test result pertains only to the sample tested and to be interpreted in the light of clinical history. These tests are accredited under ISO 15189:2012 unless specified by (^). Test marked with # is performed in an accredited referral laboratory.





BML519909

Laboratory Investigation Report

Name	: Mr. AZAD HOSSAIN PATWARY	Ref No.	: 40616
DOB	: 27/04/1987	Sample No.	: 2504557479
Age / Gender	: 37 Y / Male	Collected	: 01/04/2025 16:45
Referred by	: DR.AMAIZAH ISHTIAQ	Registered	: 02/04/2025 09:56
Centre	: CITICARE MEDICAL CENTER	Reported	: 02/04/2025 12:37

HEMATOLOGY

Test	Result	Flag	Unit	Reference Range	Methodology
------	--------	------	------	-----------------	-------------

COMPLETE BLOOD COUNT (CBC)

INTERPRETATION NOTES :

Please note update on CBC report format, reference ranges and method(Beckman Coulter).

INTERPRETATION NOTES :

RBCs - Normocytic normochromic RBCs showing mild anisopoikilocytosis with few polychromatophilic cells and poikilocytes seen.

Advise: Iron studies, rule out Anemia of Chronic Disease.

Note: Smear reviewed by the Pathologist.

Sample Type : EDTA Whole Blood

End of Report

Dr. Adley Mark Fernandes
M.D (Pathology)
Pathologist

Gyoma V. Shah
Dr. Vyoma V Shah
M.D (Pathology)
Clinical Pathologist

Jillian Joy Garcia
Jillian Joy Garcia
Laboratory Technologist

This is an electronically authenticated report

Page 3 of 3

Printed on: 02/04/2025 12:39

Test result pertains only to the sample tested and to be interpreted in the light of clinical history. These tests are accredited under ISO 15189:2012 unless specified by (^). Test marked with # is performed in an accredited referral laboratory.

