

Phone: 0522-4062223, 9305548277, 8400888844 9415577933. 9336154100, Tollfree No.: 8688360360

E-mail: charak1984@gmail.com

CMO Reg. No. RMEE 2445133 NABLReg. No. MC-2491 Certificate No. MIS-2023-0218

Patient Name : Mr.LAKSHYA MAURYA 805840 Visit No : CHA250046801

Age/Gender Registration ON : 10 Y/M : 17/Mar/2025 10:28AM Sample Collected ON Lab No : 10144096 : 17/Mar/2025 10:53AM Referred By : Dr.VIDHYA GYAN SCHOOL Sample Received ON : 17/Mar/2025 10:53AM Refer Lab/Hosp : CREDIT CLIENT Report Generated ON : 17/Mar/2025 01:42PM

FASTING,CBC (WHOLE BLOOD),ESR,LIPID-PROFILE,PROTEIN,Albumin,GLOBULIN,AG RATIO,BILIRUBIN TDI,ALK PHOS,CALCIUN,URIC Doctor Advice

ACID, CREATININE, BUN CREATININE RATIO, BUN, NA+K+, CHLORIDE, TIBC, Iron, TRANSFERRIN SATURAT



<u>VIDHYA GYAN</u>						
Test Name	Result	Unit	Bio. Ref. Range	Method		
FSD						

8.00 **Erythrocyte Sedimentation Rate ESR** 3-13 Westergreen

Note:

- 1. Test conducted on EDTA whole blood at 37°C.
- 2. ESR readings are auto- corrected with respect to Hematocrit (PCV) values.
- 3. It indicates presence and intensity of an inflammatory process. It is a prognostic test and used to monitor the course or response to treatment of diseases like tuberculosis, acute rheumatic fever. It is also increased in multiple myeloma, hypothyroidism.

HBA1C						
Glycosylated Hemoglobin	(HbA1c)	5.1	%	4 -	5.7	HPLC (EDTA)

NOTE:-

Glycosylated Hemoglobin Test (HbA1c)is performed in this laboratoryby the Gold Standard Reference method,ie:HPLC Technology(High performance Liquid Chromatography D10) from Bio-Rad Laboratories.USA.

EXPECTED (RESULT) RANGE:

Bio system Degree of normal 4.0 - 5.7 % Normal Value (OR) Non Diabetic 5.8 - 6.4 % Pre Diabetic Stage > 6.5 % Diabetic (or) Diabetic stage 6.5 - 7.0 % Well Controlled Diabet Unsatisfactory Control 7.1 - 8.0 % > 8.0 % Poor Control and needs treatment

BLOOD UREA NITROGEN					
Blood Urea Nitrogen (BUN)	12.29	mg/dL	7-21	calculated	
RUN CREATININE PATIO					

BUN CREATININE RATIO 20.40



DR. NISHANT SHARMA DR. SHADAB **PATHOLOGIST PATHOLOGIST**

Dr. SYED SAIF AHMAD MD (MICROBIOLOGY)

Print.Date/Time: 17-03-2025 15:31:18 *Patient Identity Has Not Been Verified. Not For Medicolegal

Page 1 of 8



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	<u>VII</u>	HYA GYAN		
Test Name	Result	Unit	Bio. Ref. Range	Method
URIC ACID				
Sample Type : SERUM				
SERUM URIC ACID	5.1	mg/dL	2.40 - 5.70	Uricase,Colorimetric
SERUM CALCIUM				
CALCIUM	10.3	mg/dl	8.8 - 10.8	dapta / arsenazo III

INTERPRETATION:

⁻Calcium level is decreased in patients with hemodialysis, hypoparathyroidism (primary, secondary), vitamin D deficiency, acute pancreatitis, diabetic Keto-acidosis, sepsis, acute myocardial infarction (AMI), malabsorption, osteomalacia, renal failure, rickets.

PROTEIN		y y		
PROTEIN Serum	10.00	mg/dl	6.8 - 8.5	
SERUM ALBUMIN				
ALBUMIN	5.2	gm/dl	3.20 - 5.50	Bromcresol Green (BCG)
GLOBULIN	<u> </u>	and the same of th		
GLOBULIN	4.00	gm/dl	2.0 -3.5	calculated
AG RATIO	CLI	л D л		
AG RATIO	1.50	411	1.5 : 1	



15:31:18

⁻Calcium level is increased in patients with hyperparathyroidism, Vitamin D intoxication, metastatic bone tumor, milk-alkali syndrome, multiple myeloma, Paget's disease.



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ISE Indirect

<u>VIDHYA GYAN</u>							
Test Name	Result	Unit	Bio. Ref. Range	Method			
LIPID-PROFILE							
Cholesterol/HDL Ratio	6.11	Ratio		Calculated			
LDL / HDL RATIO	4.57	Ratio		Calculated			
			Desirable / low risk - 0.5 -3.0 Low/ Moderate risk - 3.0 6.0 Elevated / High risk - >6. Desirable / low risk - 0.5 -3.0 Low/ Moderate risk - 3.0 6.0 Elevated / High risk - > 6.0)- 0 5			

CHLORIDE
Increased In:

CHLORIDE

Renal tubular diseases, Respiratory alkalosis, Drugs: Excessive administration of certain drugs (e.g., ammonium chloride, IV saline), Retention of salt and water (e.g., corticosteroids), Some cases of hyperparathyroidism, Diabetes insipidus, dehydration.

Decreased In:

Prolonged vomiting, Chronic respiratory acidosis, Salt-losing renal diseases, Adrenocortical insufficiency, Primary aldosteronism, Burns, Chronic laxative abuse

98.00



mmol/l



Tham

98 - 107



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	<u>VIDHYA GYAN</u>						
	Test Name	Result	Unit	Bio. Ref. Range	Method		
IRON			<u> </u>		<u>. </u>		
IRON		132.00	ug/ dl	59 - 148	Ferrozine-no deproteinization		

Interpretation:

Disease	Iron	TIBC	UIBC	%Transferrin Saturation	Ferritin
		A COLOR			
Iron Deficiency	Low	High	High	Low	Low
Hemochromatosis	High	Low	Low	High	High
Chronic Illness	Low	Low	Low/Normal	Low	Normal/High
Hemolytic Anemia	High	Normal/Low	Low/Normal	High	High
Sideroblastic Anemia	Normal/High	Normal/Low	Low/Normal	H igh	High
Iron Poisoning	High	Normal	Low	High	Normal

TIBC					
TIBC		345.00	ug/ml	265 - 497	calculated
TRANSFERRIN SATURATION	1				
TRANSFERRIN SATURATION		38.26	%	22 - 45	Immunoturbidimetry

INTERPRETATION:

- Low Values in iron deficiency
- High Values in iron overload
- Raised transferrin saturation is an early indicator of Iron accumulation in Genetic Haemochromatosis.

FERRITIN				
FERRITIN	42.9	ng/mL	7 - 140	CLIA

INTERPRETATION:

Ferritin is a high-molecular weight iron containing protein that functions in the body as an iron Storage compound. Ferritin provides a more sensitive, specific and reliable measurement for determining iron deficiency at an early stage. The combined use of serum ferritin levels and mean corpuscular volume (MCV) has made differentiation between iron deficiency, beta-thalassemia trait and normal subjects possible at a very high level of accuracy. Serum ferritin measurements provide important clinical parameters for assessing the response to treatment with deferoxamine, in the treatment of thalassemia. Elevated levels are seen in malignant diseases such as leukemia, Hodgkins disease, breast cancer, head and neck cancer and ovarian cancer

LIMITATIONS:

Specimens from patients who have received preparations of mouse monoclonal antibodies for diagnosis or therapy may show either false positive or depressed values.

For diagnostic purposes the ferritin result should be used in conjunction with other data, e.g.: symptoms, results of other tests, clinical impressions, etc

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<u>VIDHYA GYAN</u>							
Test Name	Result	Unit	Bio. Ref. Range	Method			
URINE EXAMINATION REPORT							
Colour-U	STRAW		Light Yellow				
Appearance (Urine)	CLEAR		Clear				
Specific Gravity	1.015		1.005 - 1.025				
pH-Urine	Acidic (6.0)		4.5 - 8.0				
PROTEIN	Absent	mg/dl	ABSENT	Dipstick			
Glucose	Absent						
Ketones	Absent		Absent				
Bilirubin-U	Absent		Absent				
Blood-U	Ab <mark>sent</mark>		Absent				
Urobilinogen-U	0.20	EU/dL	0.2 - 1.0				
Leukocytes-U	Absent Absent		Absent				
NITRITE	A <mark>bsent</mark>		Absent				
MICROSCOPIC EXAMINATION							
Pus cells / hpf	Nil	/hpf	< 5/hpf				
Epithelial Cells	Occasional	/hpf	0 - 5				
RBC / hpf	Nil		< 3/hpf				





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<u>VIDHYA GYAN</u>							
Test Name	Result	Unit	Bio. Ref. Range	Method			
CBC (COMPLETE BLOOD COUNT)							
Hb	14.7	g/dl	11 - 15	Non Cyanide			
R.B.C. COUNT	4.90	mil/cmm	4 - 5.1	Electrical			
				Impedence			
PCV	44.5	%	31 - 43	Pulse hieght			
				detection			
MCV	90.3	fL	76 - 87	calculated			
MCH	29.8	pg	26 - 28	Calculated			
MCHC	33	g/dL	33 - 35	Calculated			
RDW	13.9	%	11 - 15	RBC histogram			
				derivation			
RETIC	0.8%	%	0.3 - 1	Microscopy			
TOTAL LEUCOCYTES COUNT	7350	/cmm	4500 - 13500	Flocytrometry			
DIFFERENTIAL LEUCOCYTE COUNT							
NEUTROPHIL	48	%	40 - 70	Flowcytrometry			
LYMPHOCYTES	45	%	25 - 55	Flowcytrometry			
EOSINOPHIL	4	%	1 - 6	Flowcytrometry			
MONOCYTE	3	%	0 - 8	Flowcytrometry			
BASOPHIL	0	%	00 - 01	Flowcytrometry			
PLATELET COUNT	344,000	/cmm	150000 - 450000	Elect Imped			
PLATELET COUNT (MANUAL)	344000	/cmm	150000 - 450000	Microscopy.			
Absolute Neutrophils Count	3,528	/cmm	2000 - 7000	Calculated			
Absolute Lymphocytes Count	3,308	/cmm	1000-3000	Calculated			
Absolute Eosinophils Count	294	/cmm	20-500	Calculated			
Absolute Monocytes Count	220	/cmm	200-1000	Calculated			
Mentzer Index	18						
Peripheral Blood Picture	:						

RBC are normocytic normochromic. WBC are within normal limits. Platelets are adequate in number.





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<u>VIDHYA GYAN</u>						
Test Name	Result	Unit	Bio. Ref. Range	Method		
FASTING						
Blood Sugar Fasting	106.1	mg/dl	70 - 110	Hexokinase		
NA.W.						
NA+K+						
SODIUM Serum	141.0	MEq/L	135 - 155	ISE Direct		
POTASSIUM Serum	3.9	MEq/L	3.5 - 5.5	ISE Direct		
SERUM CREATININE			A			
CREATININE	0.60	mg/dl	0.50 - 1.40	Alkaline picrate-		
				kinetic		
BILIRUBIN TDI						
TOTAL BILIRUBIN	0.52	mg/dl	0.4 - 1.1	Diazonium Ion		
DIRECT BILIRUBIN	0.12	mg/dL	0-0.3	DIAZOTIZATION		
BILIRUBIN (INDIRECT)	0.40	mg/dl	0.1 - 1.00	CALCULATED		
ALK PHOS			7			
ALK PHOS	538.00	U/L	129 - 417	PNPP, AMP Buffer		

INTERPRETATION:

- Alkaline phosphatase is an enzyme found in your bloodstream. ALP helps break down proteins in the body and exists in different forms, depending on where it originates. Liver is one of the main sources of ALP, but some is also made in bones, intestines, pancreas, and kidneys. In pregnant women, ALP is made in the placenta.
- Higher than normal levels of ALP in blood may indicate a problem with liver or gallbladder. This could include hepatitis (liver inflammation), cirrhosis (liver scarring), liver cancer, gallstones, or a blockage in bile ducts. High levels may also indicate an issue related to the bones such as rickets, Paget's disease, bone cancer, or an overactive parathyroid gland. In rare cases, high ALP levels can indicate heart failure, kidney cancer, other cancer, mononucleosis, or bacterial infection. Having lower than normal ALP levels in blood is rare, but can indicate malnutrition, which could be caused by celiac disease or a deficiency in certain vitamins and minerals.







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Test Name	Result	Unit	Bio. Ref. Range	Method				
LIPID-PROFILE								
TOTAL CHOLESTEROL	231.10	mg/dL	Desirable: <200 mg/dl Borderline-high: 200-23 mg/dl High:>/=240 mg/dl	CHOD-PAP				
TRIGLYCERIDES	102.00	mg/dL	Normal: <150 mg/dl Borderline-high:150 - 19 mg/dl High: 200 - 499 mg/dl					
			Very high:>/=500 mg/d					
H D L CHOLESTEROL	37.80	mg/dL	30-70 mg/dl	CHER-CHOD-PAP				
L D L CHOLESTEROL	1 <mark>72.90</mark>	mg/dL	Optimal:<100 mg/dl Near Optimal:100 - 129 mg/dl Borderline High: 130 - 15	1				
			mg/dl High: 160 - 189 mg/dl Very High:>/= 190 mg/d					
VLDL	20.40	mg/dL	10 - 40	Calculated				

*** End Of Report ***



