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

0 - 15

Patient Name : Mr.PRINCE YADAV 856197 Visit No : CHA250046758

Age/Gender Registration ON : 11 Y/M : 17/Mar/2025 10:12AM Sample Collected ON Lab No : 10144053 : 17/Mar/2025 10:48AM Referred By : Dr.VIDHYA GYAN SCHOOL Sample Received ON : 17/Mar/2025 10:48AM Refer Lab/Hosp : CREDIT CLIENT Report Generated ON : 17/Mar/2025 01:41PM

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

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



Westergreen

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

12.00

Erythrocyte Sedimentation Rate ESR

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)	5.2	%	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
7.1 - 8.0 %	Unsatisfactory Control
> 8.0 %	Poor Control and needs treatment

BLOOD UREA NITROGEN				
Blood Urea Nitrogen (BUN)	8.93	mg/dL	7-21	calculated
BUN CREATININE RATIO				

BUN CREATININE RATIO 14.23



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[Checked By]

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DR. NISHANT SHARMA DR. SHADAB
PATHOLOGIST PATHOLOGIST



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	<u>VII</u>	DHYA GYAN		
Test Name	Result	Unit	Bio. Ref. Range	Method
URIC ACID	·			
Sample Type : SERUM				
SERUM URIC ACID	4.2	mg/dL	2.40 - 5.70	Uricase,Colorimetric
SERUM CALCIUM				
CALCIUM	10.8	mg/dl	8.8 - 10.8	dapta / arsenazo III

#### INTERPRETATION:

<sup>-</sup>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		7		
PROTEIN Serum	7.80	mg/dl	6.8 - 8.5	
SERUM ALBUMIN				
ALBUMIN	4.8	gm/dl	3.20 - 5.50	Bromcresol Green (BCG)
GLOBULIN		and the same of th		
GLOBULIN	3.00	gm/dl	2.0 -3.5	calculated
AG RATIO	CLI	ADA		
AG RATIO	1.60	411	1.5 : 1	



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<sup>-</sup>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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6.0 Elevated / High risk - > 6.0

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<u>VIDHYA GYAN</u>						
Test Name	Result	Unit	Bio. Ref. Range	Method		
LIPID-PROFILE						
Cholesterol/HDL Ratio	3.03	Ratio		Calculated		
LDL / HDL RATIO	1.64	Ratio		Calculated		
			Desirable / low risk - 0.	5		
			-3.0			
			Low/ Moderate risk - 3.	0-		
			6.0			
			Elevated / High risk - >6	0.0		
			Desirable / low risk - 0.	5		
			-3.0			
			Low/ Moderate risk - 3.	0-		

CHLORIDE

CHLORIDE 99.00 mmol/l 98 - 107 ISE Indirect

#### Increased In:

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





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	<u>VIDHYA GYAN</u>							
	Test Name	Result	Unit	Bio. Ref. Range	Method			
IRON								
IRON		123.00	ug/ dl	59 - 148	Ferrozine-no deproteinization			

## **Interpretation:**

Disease	Iron	TIBC	UIBC	%Transferrin Saturation	Ferritin
		A CONTRACTOR OF THE PROPERTY O			
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	High	High
Iron Poisoning	High	Normal	Low	High	Normal

TIBC				
TIBC	385.00	ug/ml	265 - 497	calculated
		iii.		
TRANSFERRIN SATURATION				
TRANSFERRIN SATURATION	31.95	%	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	20.3	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

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

**PATHOLOGIST** 

Dr. SYED SAIF AHMAD MD (MICROBIOLOGY) Page 4 of 8



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<u>VIDHYA GYAN</u>						
Test Name	Result	Unit	Bio. Ref. Range	Method		
URINE EXAMINATION REPORT						
Colour-U	Light yellow		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	<mark>Absent</mark>		Absent			
NITRITE	A <mark>bsent</mark>		Absent			
MICROSCOPIC EXAMINATION						
Pus cells / hpf	Occasional	/hpf	< 5/hpf			
Epithelial Cells	Occasional	/hpf	0 - 5			
RBC / hpf	Nil		< 3/hpf			





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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	
CBC (COMPLETE BLOOD COUNT)					
Hb	12.2	g/dl	11 - 15	Non Cyanide	
R.B.C. COUNT	4.50	mil/cmm	4 - 5.1	Electrical	
				Impedence	
PCV	38.1	%	31 - 43	Pulse hieght	
				detection	
MCV	83.9	fL	76 - 87	calculated	
MCH	26.9	pg	26 - 28	Calculated	
MCHC	32	g/dL	33 - 35	Calculated	
RDW	14.4	%	11 - 15	RBC histogram	
				derivation	
RETIC	1.0%	%	0.3 - 1	Microscopy	
TOTAL LEUCOCYTES COUNT	5940	/cmm	4500 - 13500	Flocytrometry	
DIFFERENTIAL LEUCOCYTE COUNT					
NEUTROPHIL	65	%	40 - 70	Flowcytrometry	
LYMPHOCYTES	31	%	30 - 50	Flowcytrometry	
EOSINOPHIL	2	%	1 - 6	Flowcytrometry	
MONOCYTE	2	%	0 - 8	Flowcytrometry	
BASOPHIL	0	%	00 - 01	Flowcytrometry	
PLATELET COUNT	271,000	/cmm	150000 - 450000	Elect Imped	
PLATELET COUNT (MANUAL)	271000	/cmm	150000 - 450000	Microscopy.	
Absolute Neutrophils Count	3,861	/cmm	2000 - 7000	Calculated	
Absolute Lymphocytes Count	1,841	/cmm	1000-3000	Calculated	
Absolute Eosinophils Count	119	/cmm	20-500	Calculated	
Absolute Monocytes Count	119	/cmm	200-1000	Calculated	
Mentzer Index	19				
Peripheral Blood Picture	:				

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





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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	
FASTING					
Blood Sugar Fasting	108.7	mg/dl	70 - 110	Hexokinase	
NA+K+					
SODIUM Serum	143.0	MEq/L	135 - 155	ISE Direct	
POTASSIUM Serum	3.9	MEq/L	3.5 - 5.5	ISE Direct	
SERUM CREATININE					
CREATININE	0.70	mg/dl	0.50 - 1.40	Alkaline picrate- kinetic	
BILIRUBIN TDI					
TOTAL BILIRUBIN	0.86	mg/dl	0.4 - 1.1	Diazonium Ion	
DIRECT BILIRUBIN	0.12	mg/dL	0-0.3	DIAZOTIZATION	
BILIRUBIN (INDIRECT)	0.74	mg/dl	0.1 - 1.00	CALCULATED	
ALK PHOS			1		
ALK PHOS	368.40	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	166.70	mg/dL	Desirable: <200 mg/dl Borderline-high: 200-23 mg/dl High:>/=240 mg/dl	CHOD-PAP 9	
TRIGLYCERIDES	105.00	mg/dL	Normal: <150 mg/dl Borderline-high:150 - 19 mg/dl High: 200 - 499 mg/dl Very high:>/=500 mg/d	9 endpoint	
H D L CHOLESTEROL	55.10	mg/dL	30-70 mg/dl	CHER-CHOD-PAP	
L D L CHOLESTEROL	90.60	mg/dL	Optimal:<100 mg/dl Near Optimal:100 - 129 mg/dl Borderline High: 130 - 15 mg/dl High: 160 - 189 mg/dl Very High:>/= 190 mg/d	CO-PAP	
VLDL	21.00	mg/dL	10 - 40	Calculated	

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

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