

PR.

292/05, Tulsidas Marg, Basement Chowk, Lucknow-226 003

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

E-mail: charak1984@gmail.com

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

Patient Name : Dr.ROHAN BAJPAI Visit No : CHA250037028

Age/Gender : 36 Y/M Registration ON : 01/Mar/2025 12:57PM Lab No Sample Collected ON : 10134324 : 01/Mar/2025 01:03PM Referred By : Dr.NIRUPAM PRAKASH Sample Received ON : 01/Mar/2025 01:17PM Refer Lab/Hosp : CGHS (BILLING) Report Generated ON : 01/Mar/2025 02:45PM

CBC+ESR,FASTING,URIC ACID,CRP (Quantitative),LIPID-PROFILE,KIDNEY FUNCTION TEST - I,LFT,HBA1C (EDTA) Doctor Advice :



Westergreen

Test Name	Result	Unit	Bio. Ref. Range	Method	
CBC+ESR (COMPLETE BLOOD COUNT)					







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Test Name	Result	Unit	Bio. Ref. R	ange	Method
HBA1C					
Glycosylated Hemoglobin (HbA1c)	5.8	%	4 - 5.7	HPLC (EDTA)	<u> </u>

NOTE:-

P.R.

Glycosylated Hemoglobin Test (HbA1c) is performed in this laboratory by 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	







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Test Name	Result	Unit	Bio. Ref. Range	Method
CRP-QUANTITATIVE				
CRP-OUANTITATIVE TEST	6.4	MG/L	0.1 - 6	

Method: Immunoturbidimetric

(Method: Immunoturbidimetric on photometry system)

SUMMARY: C - reactive protien (CRP) is the best known among the acute phase protiens, a group of protien whose concentration increases in blood as a response to inflammatory disorders.CRP is normally present in low concentration in blood of healthy individuals (< 1mg/L). It is elevated up to 500 mg/L in acute inflammatory processes associated with bacterial infections, post operative conditions tissue damage already measurment of CRP represents a useful aboratory test for detection of acute infection after 6 hours reaching a peak at 48 hours.. The as well as for monitoring inflammtory proceses also in acute rheumatic & gastrointestinal disease. In recent studies it has been shows that in apparrently healthy subjects there is a direct orrelation between CRP concentrations & the risk of developing oronary heart disease (CHD).

hsCRP cut off for risk assessment as per CDC/AHA

Risk Level <1.0 Low 1.0-3.0 Average High >3.0

All reports to be clinically corelated

URIC ACID	C	HAR	AK	
Sample Type : SERUM				
SERUM URIC ACID	4.8	mg/dL	2.40 - 5.70	Uricase, Colorimetric





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Test Name	Result	Unit	Bio. Ref. Range	Method
LIPID-PROFILE				
Cholesterol/HDL Ratio	4.00	Ratio	Calculat	ed
LDL / HDL RATIO	1.92	Ratio	Calculat	red

Desirable / low risk - 0.5

-3.0

Low/ Moderate risk - 3.0-

6.0

Elevated / High risk - >6.0

Desirable / low risk - 0.5

-3.0

Low/ Moderate risk - 3.0-

6.0

Elevated / High risk - > 6.0





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CBC+ESR,FASTING,URIC ACID,CRP (Quantitative),LIPID-PROFILE,KIDNEY FUNCTION TEST - I,LFT,HBA1C (EDTA) Doctor Advice :



Test Name	Result	Unit	Bio. Ref. Range	Method
CBC+ESR (COMPLETE BLOOD COUNT)				
Hb	12.1	g/dl	12 - 15	Non Cyanide
R.B.C. COUNT	5.50	mil/cmm	3.8 - 4.8	Electrical
				Impedence
PCV	42.5	%	36 - 45	Pulse hieght
				detection
MCV	77.3	fL	80 - 96	calculated
MCH	22.0	pg	27 - 33	Calculated
MCHC	28.5	g/dL	30 - 36	Calculated
RDW	15.7	%	11 - 15	RBC histogram
				derivation
RETIC	1.5 %	%	0.5 - 2.5	Microscopy
TOTAL LEUCOCYTES COUNT	9460	/cmm	4000 - 10000	Flocytrometry
DIFFERENTIAL LEUCOCYTE COUNT				
NEUTROPHIL	73	%	40 - 75	Flowcytrometry
LYMPHOCYTE	22	%	20-40	Flowcytrometry
EOSINOPHIL	1	%	1 - 6	Flowcytrometry
MONOCYTE	4	%	2 - 10	Flowcytrometry
BASOPHIL	0	%	00 - 01	Flowcytrometry
PLATELET COUNT	369,000	/cmm	150000 - 450000	Elect Imped
PLATELET COUNT (MANUAL)	369000	/cmm	150000 - 450000	Microscopy.
Mentzer Index	14	A D 4	17	
Peripheral Blood Picture	CH			

Red blood cells are microcytic hypochromic with anisocytosis+. Platelets are adequate. No immature cells or parasite seen.







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Doctor Advice : CBC+ESR,FASTING,URIC ACID,CRP (Quantitative),LIPID-PROFILE,KIDNEY FUNCTION TEST - I,LFT,HBA1C (EDTA)

		_		
Test Name	Result	Unit	Bio. Ref. Range	Method
FASTING				
Blood Sugar Fasting	108.3	mg/dl	70 - 110	Hexokinase
LIVER FUNCTION TEST				
TOTAL BILIRUBIN	0.49	mg/dl	0.4 - 1.1	Diazonium Ion
CONJUGATED (D. Bilirubin)	0.12	mg/dL	0.00-0.30	Diazotization
UNCONJUGATED (I.D. Bilirubin)	0.37	mg/dL	0.1 - 1.0	Calculated
ALK PHOS	82.50	U/L	30 - 120	PNPP, AMP Buffer
SGPT	52.0	U/L	5 - 40	UV without P5P
SGOT	42.0	U/L	5 - 40	UV without P5P
LIBIT DECEME				
LIPID-PROFILE	1/5 00	/ 11	D 1 11 000 / II	
TOTAL CHOLESTEROL	1 <mark>65.00</mark>	mg/dL	Desirable: <200 mg/dl	CHOD-PAP
			Borderline-high: 200-23	9
			mg/dl High:>/=240 mg/dl	
TRIGLYCERIDES	223.30	mg/dL	Normal: <150 mg/dl	Serum, Enzymatic,
INGLICENDES	223.30	mg/uL	Borderline-high:150 - 19	3
			mg/dl	7 Chaponit
			High: 200 - 499 mg/dl	
			Very high:>/=500 mg/d	
H D L CHOLESTEROL	41.30	mg/dL	30-70 mg/dl	CHER-CHOD-PAP
L D L CHOLESTEROL	79.10	mg/dL	Optimal:<100 mg/dl	CO-PAP
	(H)	ARI	Near Optimal: 100 - 129)
	011/	41 47	mg/dl	
			Borderline High: 130 - 15	59
			mg/dl	
			High: 160 - 189 mg/dl	
			Very High:>/= 190 mg/c	
VLDL	44.60	mg/dL	10 - 40	Calculated









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Test Name	Result	Unit	Bio. Ref. Range	Method
KIDNEY FUNCTION TEST - I				
Sample Type : SERUM				
BLOOD UREA	16.30	mg/dl	15 - 45	Urease, UV, Serum
CREATININE	0.70	mg/dl	0.50 - 1.40	Alkaline picrate- kinetic
SODIUM Serum	142.0	MEq/L	135 - 155	ISE Direct
POTASSIUM Serum	3.9	MEq/L	3.5 - 5.5	ISE Direct

*** End Of Report ***







