

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

: CHA250046972

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.SUDARSHAN YADAV

Age/Gender : 62 Y/M Lab No : 10144267 Referred By : Dr.KGMU

· CHARAK NA

Registration ON Sample Collected ON

Visit No

: 17/Mar/2025 12:00PM 17/Mar/2025 12:02PM

Sample Received ON : 17/Mar/2025 12:12PM

Report Generated ON : 17/Mar/2025 02:23PM

. CHEST PA,25 OH vit. D,HCV,HBSAg,HIV,URIC ACID,RANDOM,CREATININE,SGPT,CRP (Quantitative),ESR,CBC (WHOLE BLOOD) Doctor Advice

Test Name Bio. Ref. Range Method Unit Result ESR

Erythrocyte Sedimentation Rate ESR

21.00

0 - 20

Westergreen

Note:

PR.

Refer Lab/Hosp

- 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.

CRP-QUANTITATIVE

CRP-OUANTITATIVE TEST

7.8

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 after 6 hours reaching a peak at 48 hours.. The measurment of CRP represents a useful aboratory test for detection of acute infection 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 >3.0 High

CHARAK

All reports to be clinically corelated

URIC ACID

Sample Type: SERUM

SERUM URIC ACID

mg/dL

2.40 - 5.70

Uricase, Colorimetric

[Checked By]



5.0

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

PATHOLOGIST

Dr. SYED SAIF AHMAD MD (MICROBIOLOGY)

Print.Date/Time: 17-03-2025 17:55:09 *Patient Identity Has Not Been Verified. Not For Medicolegal

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ECLIA

Test Name	Result	Unit	Bio. Ref. Range	Method
25 OH vit D				

25 Hydroxy Vitamin D 14.11 ng/ml

Deficiency < 10 Insufficiency 10 - 30 Sufficiency 30 - 100 Toxicity > 100

DONE BY: ELECTROCHEMILUMINESCENCE IMMUNOASSAY(Cobas e 411, Unicel Dx1600, vitros ECI)

HEPATITIS B SURFACE ANTIGEN (HBsAg)

Sample Type: SERUM

HEPATITIS B SURFACE ANTIGEN NON REACTIVE <1 - Non Reactive **CMIA**

>1 - Reactive

Note: This is only a Screening test. Confirmation of the result (Non Reactive/Reactive) should be done by performing a PCR based test.

COMMENTS:

-HBsAg is the first serological marker after infection with Hepatitis B Virus appearing one to ten weeks after exposure and two to eight weeks before the onset of clinical symptoms. HBsAg persists during the acute phase and clears late in the convalescence phase. Failure to clear HBsAg within six months indicates a chronic HBsAg carrier state. HBsAg assays are used to identify the persons infected with HBV and to prevent transmission of the virus by blood and blood products as well as to monitor the status of infected individuals in combination with other hepatitis B serological markers.
-Borderline cases must be confirmed with confirmatory neutralizing assay

LIMITATIONS:

- -Results should be used in conjunction with patient history and other hepatitis markers for diagnosis of acute and chronic infections
- -Specimens from patients who have received preparations of mouse monoclonal antibodies for diagnosis or therapy may contain human anti-mouse antibodies (HAMA) which may produce anomalous values when tested with assay kits that employs mouse monoclonal antibodies
- -Heterophilic antibodies in human serum can react with reagent immunoglobulins, interfering with in vitro immunoassays. Patients routinely exposed to animals or animal serum products can be prone to this interference and anomalous results may be observed. -Cross reactivity for specimens from individual with medical conditions (Pregnancy, HIV etc) has been observed.
- -HBsAg mutations may result in a false negative result in some HBsAg assays.
 -If HBsAg results are inconsistent with clinical evidence, additional testing is suggested to confirm the result.





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Doctor Advice : CHEST PA,25 OH vit. D,HCV,HBSAg,HIV,URIC ACID,RANDOM,CREATININE,SGPT,CRP (Quantitative),ESR,CBC (WHOLE BLOOD)

Test Name Result Unit Bio. Ref. Range Method

HIV-SEROLOGY NON REACTIVE <1.0 : NON REACTIVE >1.0 : REACTIVE

Done by: Vitros ECI (Sandwich Assay)

Note:-Elisa test is a screening method for HIV. It is known to give false Positive & Negative result.

Hence confirmation: "Western Blot" method is advised.

HEPATITIS C VIRUS (HCV) ANTIBODIES

HEPATITIS C VIRUS (HCV) ANTIBODIES NON REACTIVE

Non Reactive

(TRIO DOT ASSAY)

Note: This is only a Screening test. Confirmation of the result (Non Reactive/Reactive) should be done by performing a PCR based test.





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P.R.

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Test Name	Result	Unit	Bio. Ref. Range	Method
CBC (COMPLETE BLOOD COUNT)				
Hb	12.7	g/dl	12 - 15	Non Cyanide
R.B.C. COUNT	4.70	mil/cmm	3.8 - 4.8	Electrical
				Impedence
PCV	40.2	%	36 - 45	Pulse hieght
				detection
MCV	85.4	fL	80 - 96	calculated
MCH	27.0	pg	27 - 33	Calculated
MCHC	31.6	g/dL	30 - 36	Calculated
RDW	15	%	11 - 15	RBC histogram
	/ /			derivation
RETIC	0.5 %	%	0.5 - 2.5	Microscopy
TOTAL LEUCOCYTES COUNT	7150	/cmm	4000 - 10000	Flocytrometry
DIFFERENTIAL LEUCOCYTE COUNT	50	04	40. 75	
NEUTROPHIL	52	%	40 - 75	Flowcytrometry
LYMPHOCYTES	39	%	25 - 45	Flowcytrometry
EOSINOPHIL	4	%	1 - 6	Flowcytrometry
MONOCYTE	5	%	2 - 10	Flowcytrometry
BASOPHIL	0	%	00 - 01	Flowcytrometry
PLATELET COUNT	167,000	/cmm	150000 - 450000	Elect Imped
PLATELET COUNT (MANUAL)	167000	/cmm	150000 - 450000	Microscopy.
Absolute Neutrophils Count	3,718	/cmm	2000 - 7000	Calculated
Absolute Lymphocytes Count	2,788	/cmm	1000-3000	Calculated
Absolute Eosinophils Count	286	/cmm	20-500	Calculated
Absolute Monocytes Count	358	/cmm	200-1000	Calculated
Mentzer Index	18			
Peripheral Blood Picture	:			

Red blood cells are normocytic normochromic . Platelets are adequate. No immature cells or parasite seen.





Than



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Test Name	Result	Unit	Bio. Ref. Range	Method
BLOOD SUGAR RANDOM				
BLOOD SUGAR RANDOM	98.2	mg/dl	70 - 170	Hexokinase
SERUM CREATININE				
CREATININE	0.70	mg/dl	0.50 - 1.40	Alkaline picrate- kinetic
SGPT				
SGPT	30.0	U/L	5 - 40	UV without P5P

*** End Of Report ***





