

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

: CHA250033546

: 24/Feb/2025 08:04PM

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.DEEPAK SONI

Age/Gender : 22 Y/M Lab No : 10130842 Referred By : Dr.MANISH TANDON

: CHARAK NA Report Generated ON

Sample Collected ON 24/Feb/2025 08:08PM Sample Received ON : 24/Feb/2025 08:37PM

25/Feb/2025 10:23AM . RANDOM,NA+K+,CREATININE,LFT,CRP (Quantitative),ESR,CBC (WHOLE BLOOD),USG WHOLE ABDOMEN,DIGITAL 1

Test Name Bio. Ref. Range Method Unit Result

ESR

PR.

Erythrocyte Sedimentation Rate ESR

15.00

0 - 15

Visit No

Registration ON

Westergreen

Note:

Refer Lab/Hosp

Doctor Advice

- 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

2.6

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 High >3.0

CHARAK

All reports to be clinically corelated



DR. ADITI D AGARWAL



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Doctor Advice : RANDOM,NA+K+,CREATININE,LFT,CRP (Quantitative),ESR,CBC (WHOLE BLOOD),USG WHOLE ABDOMEN,DIGITAL 1

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Test Name	Result	Unit	Bio. Ref. Range	Method
CBC (COMPLETE BLOOD COUNT)				
Hb	14.4	g/dl	12 - 15	Non Cyanide
R.B.C. COUNT	4.60	mil/cmm	3.8 - 4.8	Electrical
				Impedence
PCV	43.5	%	36 - 45	Pulse hieght
				detection
MCV	95.0	fL	80 - 96	calculated
MCH	31.4	pg	27 - 33	Calculated
MCHC	33.1	g/dL	30 - 36	Calculated
RDW	13.2	%	11 - 15	RBC histogram
				derivation
RETIC	0.9 %	%	0.5 - 2.5	Microscopy
TOTAL LEUCOCYTES COUNT	8440	/cmm	4000 - 10000	Flocytrometry
DIFFERENTIAL LEUCOCYTE COUNT				
NEUTROPHIL	68	%	40 - 75	Flowcytrometry
LYMPHOCYTES	23	%	25 - 45	Flowcytrometry
EOSINOPHIL	5	%	1 - 6	Flowcytrometry
MONOCYTE	4	%	2 - 10	Flowcytrometry
BASOPHIL	0	%	00 - 01	Flowcytrometry
PLATELET COUNT	258,000	/cmm	150000 - 450000	Elect Imped
PLATELET COUNT (MANUAL)	258000	/cmm	150000 - 450000	Microscopy.
Absolute Neutrophils Count	5,739	/cmm	2000 - 7000	Calculated
Absolute Lymphocytes Count	1,941	/cmm	1000-3000	Calculated
Absolute Eosinophils Count	422	/cmm	20-500	Calculated
Absolute Monocytes Count	338	/cmm	200-1000	Calculated
Mentzer Index	21			
Peripheral Blood Picture	:			

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







P.R.

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				<u> </u>	
Test Name	Result	Unit	Bio. Ref. Range	Method	
BLOOD SUGAR RANDOM					
BLOOD SUGAR RANDOM	92.1	mg/dl	70 - 170	Hexokinase	
NA+K+					
SODIUM Serum	137.3	MEq/L	135 - 155	ISE Direct	
POTASSIUM Serum	4.3	MEq/L	3.5 - 5.5	ISE Direct	
SERUM CREATININE		7			
CREATININE	0.80	mg/dl	0.50 - 1.40	Alkaline picrate-	
				kinetic	
LIVER FUNCTION TEST					
TOTAL BILIRUBIN	0.40	mg/dl	0.4 - 1.1	Diazonium Ion	
CONJUGATED (D. Bilirubin)	0.10	mg/dL	0.00-0.30	Diazotization	
UNCONJUGATED (I.D. Bilirubin)	0.30	mg/dL	0.1 - 1.0	Calculated	
ALK PHOS	8 <mark>2.50</mark>	U/L	30 - 120	PNPP, AMP Buffer	
SGPT	25.2	U/L	5 - 40	UV without P5P	
SGOT	46.4	U/L	5 - 40	UV without P5P	

*** End Of Report ***

CHARAK







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ULTRASOUND STUDY OF WHOLE ABDOMEN

Excessive gaseous abdomen

- <u>Liver</u> is mildly enlarged in size (~152mm) and shows mild inhomogenous echotexture of liver parenchyma. No intrahepatic biliary radicle dilatation is seen. No space occupying lesion is seen. Hepatic veins and IVC are seen normally.
- <u>Gall bladder</u> is normal in size and shows anechoic lumen. No calculus / mass lesion is seen. GB walls are not thickened.
- CBD is normal at porta. No obstructive lesion is seen.
- Portal vein is normal at porta.
- <u>Pancreas</u> is normal in size and shows homogenous echotexture of parenchyma. PD is not dilated. No parenchymal calcification is seen. No peripancreatic collection is seen.
- <u>Spleen</u> is normal in size and shows homogenous echotexture of parenchyma. No SOL is seen.
- Few subcentimeteric mesenteric lymphnodes are seen with maintained hilum (non specific).
- No ascites is seen.
- <u>Both kidneys</u> are normal in size and position. No hydronephrosis is seen. No calculus or mass lesion is seen. Cortico-medullary differentiation is well maintained. Parenchymal thickness is normal. No scarring is seen. Right kidney measures 91 x 45 mm in size. Left kidney measures 95 x 49 mm in size.
- <u>Ureters</u> Both ureters are not dilated. UVJ are seen normally.
- <u>Urinary bladder</u> is inadequate distended.

OPINION:

PR

- MILD HEPATOMEGALY WITH MILD INHOMOGENOUS ECHOTEXTURE OF LIVER PARENCHYMA.
- FEW SUBCENTIMETERIC MESENTERIC LYMPHNODES WITH MAINTAINED HILUM (NON SPECIFIC).

(Possibility of acid peptic disease could not be ruled out).

Clinical correlation is necessary.

[DR. R. K. SINGH, MD]

Transcribed by Gausiya



PR.

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SKIAGRAM ABDOMEN (ERECT) AP VIEW

- No free gas is seen under both dome of diaphragm.
- No abnormal air fluid levels are seen.

Clinical correlation is necessary.

Transcribed By: Purvi

[DR. R. K. SINGH, MD]

*** End Of Report ***

