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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

0 - 20

Patient Name : Mr.RAMESH RAM Visit No : CHA250035932

30.00

Age/Gender : 70 Y/M Registration ON : 28/Feb/2025 09:13AM Lab No Sample Collected ON : 10133228 : 28/Feb/2025 09:17AM Referred By : Dr.NIRUPAM PRAKASH Sample Received ON : 28/Feb/2025 09:44AM Refer Lab/Hosp : CGHS (BILLING) Report Generated ON 28/Feb/2025 11:16AM

CHEST PA,ECG,NA+K+,URIC ACID,URINE COM. EXMAMINATION,PSA-TOTAL,HBA1C (EDTA),PP,FASTING,LIPID-PROFILE,LFT,KIDNEY Doctor Advice :

FUNCTION TEST - I,CBC+ESR

Erythrocyte Sedimentation Rate ESR



Westergreen

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







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FUNCTION TEST - I,CBC+ESR



Test Name	Result	Unit	Bio. Ref. Range	Method	
HBA1C					
Glycosylated Hemoglobin (HbA1c)	6.3	%	4 - 5.7	HPLC (EDTA)	

NOTE:-

PR.

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

URIC ACID				
Sample Type : SERUM				
SERUM URIC ACID	11.2	mg/dL	2.40 - 5.70	Uricase,Colorimetric
LIPID-PROFILE	CH	AD/	\K	
Cholesterol/HDL Ratio	3.29	Ratio		Calculated
LDL / HDL RATIO	1.75	Ratio		Calculated
			Desirable / low risk - ().5
			-3.0	
			Low/ Moderate risk - 3	3.0-
			6.0	
			Elevated / High risk - >	6.0
			Desirable / low risk - (0.5
			-3.0	
			Low/ Moderate risk - 3	3.0-
			6.0	
			Elevated / High risk - >	6.0



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

Dr. SYED SAIF AHMAD MD (MICROBIOLOGY)

Print.Date/Time: 28-02-2025 15:10:15 *Patient Identity Has Not Been Verified. Not For Medicolegal

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FUNCTION TEST - I,CBC+ESR

Test Name	Result	Unit	Bio. Ref. Range	Method
URINE EXAMINATION REPORT				
Colour-U	Light yellow		Light Yellow	
Appearance (Urine)	CLEAR		Clear	
Specific Gravity	1.010		1.005 - 1.025	
pH-Urine	Acidic (6.0)		4.5 - 8.0	
PROTEIN	10 mg/dl	mg/dl	ABSENT	Dipstick
Glucose	Absent			
Ketones	Absent		Absent	
Bilirubin-U	Absent		Absent	
Blood-U	Absent		Absent	
Urobilinogen-U	0.20	EU/dL	0.2 - 1.0	
Leukocytes-U	A <mark>bsent</mark>		Absent	
NITRITE	<mark>Absent</mark>		Absent	
MICROSCOPIC EXAMINATION				
Pus cells / hpf	Nil	/hpf	< 5/hpf	
Epithelial Cells	1-2	/hpf	0 - 5	
RBC / hpf	Nil		< 3/hpf	

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FUNCTION TEST - I,CBC+ESR

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Test Name	Result	Unit	Bio. Ref. Range	Method
CBC+ESR (COMPLETE BLOOD COUNT)				
Hb	12.0	g/dl	12 - 15	Non Cyanide
R.B.C. COUNT	4.10	mil/cmm	3.8 - 4.8	Electrical
				Impedence
PCV	37.8	%	36 - 45	Pulse hieght
				detection
MCV	91.5	fL	80 - 96	calculated
MCH	29.1	pg	27 - 33	Calculated
MCHC	31.7	g/dL	30 - 36	Calculated
RDW	14.7	%	11 - 15	RBC histogram
				derivation
RETIC	0.8 %	%	0.5 - 2.5	Microscopy
TOTAL LEUCOCYTES COUNT	9140	/cmm	4000 - 10000	Flocytrometry
DIFFERENTIAL LEUCOCYTE COUNT				
NEUTROPHIL	73	%	40 - 75	Flowcytrometry
LYMPHOCYTE	21	%	20-40	Flowcytrometry
EOSINOPHIL	2	%	1 - 6	Flowcytrometry
MONOCYTE	4	%	2 - 10	Flowcytrometry
BASOPHIL	0	%	00 - 01	Flowcytrometry
PLATELET COUNT	205,000	/cmm	150000 - 450000	Elect Imped
PLATELET COUNT (MANUAL)	205000	/cmm	150000 - 450000	Microscopy.
Mentzer Index	22		0.17	
Peripheral Blood Picture	GH			

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





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Doctor Advice : CHEST PA,ECG,NA+K+,URIC ACID,URINE COM. EXMAMINATION,PSA-TOTAL,HBA1C (EDTA),PP,FASTING,LIPID-PROFILE,LFT,KIDNEY

FUNCTION TEST - I,CBC+ESR



Test Name	Result	Unit		je Metho
FASTING	<u> </u>	·	<u> </u>	· .
Blood Sugar Fasting	110.6	mg/dl	70 - 110	Hexokinase
PP				
Blood Sugar PP	128.0	mg/dl	up to - 170	Hexokinase
NA+K+				
SODIUM Serum	136.0	MEq/L	135 - 155	ISE Direct
POTASSIUM Serum	5.8	MEq/L	3.5 - 5.5	ISE Direct
LIVER FUNCTION TEST				
TOTAL BILIRUBIN	0.70	mg/dl	0.4 - 1.1	Diazonium Ion
CONJUGATED (D. Bilirubin)	0.11	mg/dL	0.00-0.30	Diazotization
UNCONJUGATED (I.D. Bilirubin)	0.59	mg/dL	0.1 - 1.0	Calculated
ALK PHOS	123. <mark>90</mark>	U/L	30 - 120	PNPP, AMP Buffer
SGPT	33.0	U/L	5 - 40	UV without P5P
SGOT	40.0	U/L	5 - 40	UV without P5P
LIPID-PROFILE				
TOTAL CHOLESTEROL	199.50	mg/dL	Desirable: <200 mg/dl Borderline-high: 200-239 mg/dl High:>/=240 mg/dl	CHOD-PAP
TRIGLYCERIDES	162.90	mg/dL	Normal: <150 mg/dl	Serum, Enzymatic,
	CH	IAF	Borderline-high:150 - 19 ⁰ mg/dl High: 200 - 499 mg/dl Very high:>/=500 mg/dl	·
H D L CHOLESTEROL	60.70	mg/dL	30-70 mg/dl	CHER-CHOD-PAP
L D L CHOLESTEROL	106.30	mg/dL	Optimal:<100 mg/dl Near Optimal:100 - 129 mg/dl Borderline High: 130 - 15 mg/dl High: 160 - 189 mg/dl	9
VLDL	32.50	mg/dL	Very High:>/= 190 mg/d 10 - 40	ı Calculated
VLUL	32.30	my/uL	10 - 40	Calculated





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FUNCTION TEST - I,CBC+ESR



Test Name	Result	Unit	Bio. Ref. Rang	e Method
KIDNEY FUNCTION TEST - I				
Sample Type : SERUM				
BLOOD UREA	67.90	mg/dl	15 - 45	Urease, UV, Serum
CREATININE	1.60	mg/dl	0.50 - 1.40	Alkaline picrate- kinetic
SODIUM Serum	136.0	MEq/L	135 - 155	ISE Direct
POTASSIUM Serum	5.8	MEq/L	3.5 - 5.5	ISE Direct
PSA-TOTAL				
PROSTATE SPECIFIC ANTIGEN	0.19	ng/mL	0.2-4.0	CLIA

COMMENT: 1. Prostate specific antigen (PSA) is useful for diagnosis of disseminated CA prostate & its equential measurement is the most sensitive measure of monitoring treatment of disseminated CA prostate with its shorter half life (half life of 2.2 days only) it is superior to prostatic acis phosphatase(PAP). PSA is elevated in nearly all patients with stage D carcinoma whereas PAP is elevated in only 45 % of patient. Mild PSA elevation are also reported in some patients of BHP.

2. Blood samples should be obtained before prostate biopsy or prostatecomy or prostatic massage or digital pre rectal examination as it may result intrasient levation of PSA value for few days.

NOTE: - PSA values obtained in different types of PSA assay methods cannot be used interchangeably as the PSA value in a given sample varies with assays from different manufactures due to difference in assay methodology and reagent specificity. If in the course of monitoring a patient the assay method used for determination is changed, additional sequential testing should be carried out to confirm baseline value.

DONE BY:

Enhanced Chemiluminescence "VITROS ECI

*** End Of Report ***





DR. NISHANT SHARMA DR. SHADAB

PATHOLOGIST

Dr. SYED SAIF AHMAD

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Age/Gender : 70 Y/M **Lab No** : **10133228**

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Refer Lab/Hosp : CGHS (BILLING)

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Visit No : CHA250035932

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Sample Received ON

Report Generated ON : 28/Feb/2025 10:45AM

ECG-REPORT

RATE : 60 bpm

* RHYTHM : Normal

* P wave : -

* PR interval : -

* QRS Axis : Left axis

Duration : 140 m sec

Configuration : Normal

* ST-T Changes : None

* QT interval :

* QTc interval : Sec.

* Other :

OPINION: REGULAR RV PACED RHYTHM

(FINDING TO BE CORRELATED CLINICALLY)

[DR. PANKAJ RASTOGI, MD, DM]



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 : 28/Feb/2025 09:13AM

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Referred By : Dr.NIRUPAM PRAKASH Sample Received ON :

Refer Lab/Hosp : CGHS (BILLING) Report Generated ON : 28/Feb/2025 02:36PM

SKIAGRAM CHEST PA VIEW

• Both lung fields are clear.

- Bilateral hilar shadows are normal.
- Cardiac shadow is within normal limits.
- Both CP angles are clear.
- Pacemaker is seen on left side chest.
- Both domes of diaphragm are sharply defined.

To be correlated with previous records.

[DR. RAJESH KUMAR SHARMA, MD]

Transcribed by R R...

*** End Of Report ***

