

#### B.U.R./N.M.T/Y/ 2458 SENIOR CITIZEN

: K.N. Pemmareddy : ನಂ.19, ನಾಗಸಾಥಪುರ, ಬೆಂಗಳೂರು ಪಬ್ಲಿಕ್ ಸ್ಕೂಲ್ ಪರಪ್ಪನ ಅಗ್ರಹಾರ, ಬೆಂಗಳೂರು -560100 Name ವಿಳಾಸ

: #19, Naganathapura, Bangalore Public School, Parappana Agrahara, Bengaluru-100

ಜನ್ಮ ದಿನಾಂಕ: 06/03/1953 Date of Birth : ದೂರವಾಣಿ:

Phone: 8694986876

Address

ರಕ್ತದ ಗುಂಪು: B+ve Blood Group:



# **Prabha Eye Clinic & Research Center**

# 504, 40th Cross, 8th Block, Jayanagar, Bengaluru - 560 070. Tel.: 080-26659595, 26659090, 42659090, 46659595 Fax: 080-22446360

email:info@prabhaeyeclinic.com

web:www.prabhaeyeclinic.com

### PATIENT SUMMARY

Page 1 of 1

Patient: PEMMA REDDY - 69/Years MALE

OP Number: KA-PEC2022/346630

Address : CL

: CLUMAX

Phone

: +918694986876

### 14/03/2022

### **OPTOMETRIST FINDINGS (-13:45:22)**

**UNAIDED VISION DIST** 

6/9P RE 6/6P LE

UNAIDED VISION NEAR

N8 RE N8 LE

**COLOR VISION** 

RE Normal LE Normal

Sleeping with Contact Lens

NO

# DOCTOR ADVICE (DR.RAKSHA V - 13:47:03)

PRESENTING COMPLAINTS

clumax, h/o dm, htn since 20 years

LIDS & ADNEXA

RE: N; LE: N

**PUPIL** 

RE: RRR; LE: RRR

CORNEA

RERE: CLEAR; LE: CLEAR

LENS

RERE: SMC; LE: SMC

CONJUNCTIVA

RERE: N; LE: N

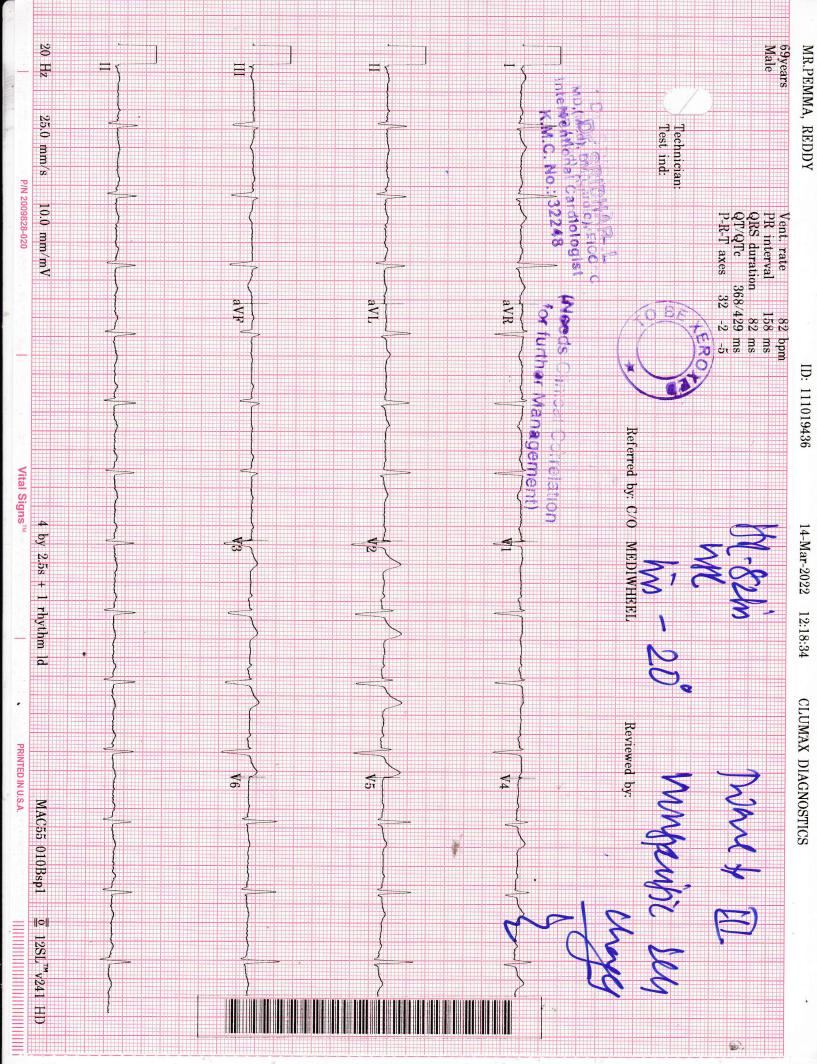
IRIS .

RERE: N; LE: N

ANTERIOR CHAMBER

RE:QUIET; LE: QUIET

Thanking you for giving us an opportunity to provide you eye care services.





Name ·	MR.PEMMA REDDY	ID	MED111019436
Age & Gender	69Y/MALE	Visit Date	14/03/2022
Ref Doctor	MediWheel		, 55/2022

# **2D ECHOCARDIOGRAPHIC STUDY**

## M mode measurement:

AORTA : 2.82 cms

LEFT ATRIUM : 4.08 cms

AVS : 1.47 cms

LEFT VENTRICLE (DIASTOLE) : 4.61 cms

(SYSTOLE) : 3.39 cms

VENTRICULAR SEPTUM (DIASTOLE) : 1.39 cms

(SYSTOLE) : 1.63 cms

POSTERIOR WALL (DIASTOLE) : 1.43 cms

(SYSTOLE) : 2.33 cms

EDV : 97 ml

 $ESV_{\bullet}$ : 47 ml

FRACTIONAL SHORTENING : 26 %

EJECTION FRACTION : 51 %

EPSS : cms

RVID : 1.96 cms

# DOPPLER MEASUREMENTS

MITRAL VALVE : 'E' -0.71m/s 'A' -1.00m/s

: 'E' -0.71m/s 'A' -1.00m/s TRIVIAL MR E/A REVERSED

AORTIC VALVE :1.35 m/s

5 m/s NO AR

TRICUSPID VALVE : PASP : 22 mmHg

TRIVIAL TR

PULMONARY VALVE

:0.84 m/s

NO PR

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Age & Gender	69Y/MALE	Visit Date	14/03/2022
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:2:

### 2D ECHOCARDIOGRAPHY FINDINGS:

Left Ventricle :

Concentric Left Ventricular Hypertrophy

Left Atrium

Dilated

Right Ventricle

Normal

Right Atrium

Normal.

Mitral valve

Normal, No mitral valve prolapse.

Aortic valve

Normal, Trileaflet

Tricuspid valve

Normal.

Pulmonary valve

Normal.

IAS

Intact.

IVS

Intact.

Pericardium

No Pericardial effusion.

# **IMPRESSION:**

- CONCENTRIC LEFT VENTRICULAR HYPERTROPHY
- > TRIVIAL MITRAL REGURGITATION
- > TRIVIAL TRICUSPID REGURGITATION. PASP 22 mmHg
- > LV DIASTOLIC DYSFUNCTION
- > ADEQUATE LV SYSTOLIC FUNCTION. EF: 51 %
- > NO CLOTS / PERICARDIAL EFFUSION / VEGETATION.

(KINDLY CORRELATE CLINICALLY AND WITH ECG)

DR.SRIDHAR.L MD,DM,FICC. CONSULTANT CARDIOLOGIST

Ls/ml Dr. SRIDHAR L

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Age & Gender	69Y/MALE	Visit Date	14/03/2022
Ref Doctor	MediWheel		

### **ABDOMINO-PELVIC ULTRASONOGRAPHY**

**LIVER** is normal in shape, size and has increased echopattern. No evidence of focal lesion or intrahepatic biliary ductal dilatation. Hepatic and portal vein radicals are normal.

GALL BLADDER show normal shape and has clear contents.
Gall bladder wall is of normal thickness.
CBD is of normal calibre.

**PANCREAS** has normal shape, size and uniform echopattern. No evidence of ductal dilatation or calcification.

**SPLEEN** show normal shape, size and echopattern.

No demonstrable Para -aortic lymphadenopathy.

**KIDNEYS** move well with respiration and have normal shape, size and echopattern. Cortico- medullary differentiations are well madeout. No evidence of calculus or hydronephrosis.

The kidney measures as follows

	Bipolar length (cms)	Parenchymal thickness (cms)
Right Kidney	9.8	1.7
Left Kidney	9.1	1.8

URINARY BLADDER show normal shape and wall thickness.

It has clear contents. No evidence of diverticula.

Prevoid: 170ml.

Postvoid: 20ml.

**PROSTATE** is enlarged in size and measures 3.8 x 3.7 x 3.8 cms, wt-29gms.

No evidence of ascites.





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# **Impression:**

- > Prostatomegaly.
- > Increased hepatic echopattern suggestive of fatty infiltration

**CONSULTANT RADIOLOGISTS:** 

DR. H. K. ANAND

DR. PRAJNA SHENOY

DR. MAHESH. M. S

DR. RADHA KRISHNA. A.

DR. HIMA BINDU.P Ms/so





Name	PEMMA REDDY	Customer ID	MED111019436
Age & Gender	69Y/M	Visit Date	Mar 14 2022 9:18AM
Ref Doctor	MediWheel		

# X - RAY CHEST PA VIEW

Bilateral lung fields appear normal.

Cardiac size is within normal limits.

Bilateral hilar regions appear normal.

Bilateral domes of diaphragm and costophrenic angles are normal.

Visualised bones and soft tissues appear normal.

Impression: Essentially normal study.

DR. H.K. ANAND

DR. SHWETHAS

DR. PRAJNA SHENOY

DR. MAHESH MS

CONSULTANT RADIOLOGISTS

: Mr. PEMMA REDDY Name

PID No. : MED111019436 Register On : 14/03/2022 9:23 AM : 922016198 SID No. Collection On : 14/03/2022 9:44 AM Age / Sex : 69 Year(s) / Male Report On : 15/03/2022 2:04 PM

**Printed On** 

: 16/03/2022 8:16 PM



: OP Ref. Dr : MediWheel

Type

Investigation  HAEMATOLOGY	Observed Value	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
Complete Blood Count With - ESR			
Haemoglobin (EDTA Blood/Spectrophotometry)	15.0	g/dL	13.5 - 18.0
Packed Cell Volume(PCV)/Haematocrit (EDTA Blood/Derived from Impedance)	48.3	%	42 - 52
RBC Count (EDTA Blood/Impedance Variation)	6.03	mill/cu.mm	4.7 - 6.0
Mean Corpuscular Volume(MCV) (EDTA Blood/Derived from Impedance)	80.0	fL	78 - 100
Mean Corpuscular Haemoglobin(MCH) (EDTA Blood/Derived from Impedance)	24.9	pg	27 - 32
Mean Corpuscular Haemoglobin concentration(MCHC) (EDTA Blood/Derived from Impedance)	31.0	g/dL	32 - 36
RDW-CV (EDTA Blood/Derived from Impedance)	17.0	%	11.5 - 16.0
RDW-SD (EDTA Blood/Derived from Impedance)	48.1	fL	39 - 46
Total Leukocyte Count (TC) (EDTA Blood/Impedance Variation)	13300	cells/cu.mm	4000 - 11000
Neutrophils (EDTA Blood/Impedance Variation & Flow Cytometry)	78.4	%	40 - 75
Lymphocytes (EDTA Blood/Impedance Variation & Flow Cytometry)	12.7	%	20 - 45
Eosinophils (EDTA Blood/Impedance Variation & Flow Cytometry)	0.9	%	01 - 06



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Investigation	Observed Value	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
Monocytes (EDTA Blood/Impedance Variation & Flow Cytometry)	7.1	%	01 - 10
Basophils (EDTA Blood/Impedance Variation & Flow Cytometry)	0.9	%	00 - 02
Absolute Neutrophil count (EDTA Blood/Impedance Variation & Flow Cytometry)	10.4	10^3 / μ1	1.5 - 6.6
Absolute Lymphocyte Count (EDTA Blood/Impedance Variation & Flow Cytometry)	1.7	10^3 / µl	1.5 - 3.5
Absolute Eosinophil Count (AEC) (EDTA Blood/Impedance Variation & Flow Cytometry)	0.10	10^3 / μl	0.04 - 0.44
Absolute Monocyte Count (EDTA Blood/Impedance Variation & Flow Cytometry)	0.9	10^3 / μl	< 1.0
Absolute Basophil count (EDTA Blood/Impedance Variation & Flow Cytometry)	0.1	10^3 / μl	< 0.2
Platelet Count (EDTA Blood/Impedance Variation)	322	10^3 / μl	150 - 450
MPV (EDTA Blood/Derived from Impedance)	8.4	fL	7.9 - 13.7
PCT (EDTA Blood/Automated Blood cell Counter)	0.271	%	0.18 - 0.28
ESR (Erythrocyte Sedimentation Rate) (EDTA Blood/Modified Westergren)	39	mm/hr	< 20



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Investigation	Observed Value	<u>Unit</u>	Biological Reference Interval
<b>BIOCHEMISTRY</b>			
Liver Function Test			
Bilirubin(Total) (Serum/Diazotized Sulfanilic Acid)	1.1	mg/dL	0.1 - 1.2
Bilirubin(Direct) (Serum/Diazotized Sulfanilic Acid)	0.2	mg/dL	0.0 - 0.3
Bilirubin(Indirect) (Serum/Derived)	0.9	mg/dL	0.1 - 1.0
Total Protein (Serum/Biuret)	7.4	gm/dL	6.0 - 8.0
Albumin (Serum/Bromocresol green)	3.8	gm/dL	3.5 - 5.2
Globulin (Serum/ <i>Derived</i> )	3.6	gm/dL	2.3 - 3.6
A : G Ratio (Serum/Derived)	1.1		1.1 - 2.2
SGOT/AST (Aspartate Aminotransferase) (Serum/IFCC Kinetic)	10	U/L	5 - 40
SGPT/ALT (Alanine Aminotransferase) (Serum/IFCC / Kinetic)	12	U/L	5 - 41
Alkaline Phosphatase (SAP) (Serum/IFCC Kinetic)	95	U/L	56 - 119
GGT(Gamma Glutamyl Transpeptidase) (Serum/SZASZ standarised IFCC)	10	U/L	< 55





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Investigation	Observed <u>Value</u>	<u>Unit</u>	<u>Biological</u> Reference Interval
<u>Lipid Profile</u>			
Cholesterol Total (Serum/Cholesterol oxidase/Peroxidase)	109	mg/dL	Optimal: < 200 Borderline: 200 - 239 High Risk: >= 240
Triglycerides (Serum/Glycerol phosphate oxidase / peroxidase)	88	mg/dL	Optimal: < 150 Borderline: 150 - 199 High: 200 - 499 Very High: >= 500

**INTERPRETATION:** The reference ranges are based on fasting condition. Triglyceride levels change drastically in response to food, increasing as much as 5 to 10 times the fasting levels, just a few hours after eating. Fasting triglyceride levels show considerable diurnal variation too. There is evidence recommending triglycerides estimation in non-fasting condition for evaluating the risk of heart disease and screening for metabolic syndrome, as non-fasting sample is more representative of the "usual" circulating level of triglycerides during most part of the day.

HDL Cholesterol (Serum/Immunoinhibition)	32	mg/dL	Optimal(Negative Risk Factor): >= 60 Borderline: 40 - 59 High Risk: < 40
LDL Cholesterol (Serum/Calculated)	59.4	mg/dL	Optimal: < 100 Above Optimal: 100 - 129 Borderline: 130 - 159 High: 160 - 189 Very High: >=190
VLDL Cholesterol (Serum/Calculated)	17.6	mg/dL	< 30
Non HDL Cholesterol (Serum/Calculated)	77.0	mg/dL	Optimal: < 130 Above Optimal: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very High: >= 220

**INTERPRETATION:** 1. Non-HDL Cholesterol is now proven to be a better cardiovascular risk marker than LDL Cholesterol. 2. It is the sum of all potentially atherogenic proteins including LDL, IDL, VLDL and chylomicrons and it is the "new bad cholesterol" and is a co-primary target for cholesterol lowering therapy.





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Investigation	Observed <u>Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
Total Cholesterol/HDL Cholesterol Ratio (Serum/Calculated)	3.4		Optimal: < 3.3 Low Risk: 3.4 - 4.4 Average Risk: 4.5 - 7.1 Moderate Risk: 7.2 - 11.0 High Risk: > 11.0
Triglyceride/HDL Cholesterol Ratio (TG/HDL) (Serum/Calculated)	2.8		Optimal: < 2.5 Mild to moderate risk: 2.5 - 5.0 High Risk: > 5.0
LDL/HDL Cholesterol Ratio (Serum/Calculated)	1.9		Optimal: 0.5 - 3.0 Borderline: 3.1 - 6.0 High Risk: > 6.0

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Investigation	<u>Observed</u> <u>Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
Glycosylated Haemoglobin (HbA1c)			
HbA1C (Whole Blood/ <i>HPLC</i> )	6.8	%	Normal: 4.5 - 5.6 Prediabetes: 5.7 - 6.4 Diabetic: >= 6.5

INTERPRETATION: If Diabetes - Good control: 6.1 - 7.0 %, Fair control: 7.1 - 8.0 %, Poor control >= 8.1 %

Estimated Average Glucose 148.46 mg/dL

(Whole Blood)

### **INTERPRETATION: Comments**

HbA1c provides an index of Average Blood Glucose levels over the past 8 - 12 weeks and is a much better indicator of long term glycemic control as compared to blood and urinary glucose determinations.

Conditions that prolong RBC life span like Iron deficiency anemia, Vitamin B12 & Folate deficiency,

hypertriglyceridemia,hyperbilirubinemia,Drugs, Alcohol, Lead Poisoning, Asplenia can give falsely elevated HbAlC values. Conditions that shorten RBC survival like acute or chronic blood loss, hemolytic anemia, Hemoglobinopathies, Splenomegaly,Vitamin E ingestion, Pregnancy, End stage Renal disease can cause falsely low HbAlc.





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Investigation	Observed Un	<u>Biological</u>
-	<u>Value</u>	Reference Interval

### **IMMUNOASSAY**

### THYROID PROFILE / TFT

T3 (Triiodothyronine) - Total 1.08 ng/mL 0.4 - 1.81

(Serum/CMIA)

### INTERPRETATION:

#### **Comment:**

Total T3 variation can be seen in other condition like pregnancy, drugs, nephrosis etc. In such cases, Free T3 is recommended as it is Metabolically active.

T4 (Thyroxine) - Total 9.50 μg/dL 4.2 - 12.0

(Serum/CMIA)

#### INTERPRETATION:

#### Comment:

Total T4 variation can be seen in other condition like pregnancy, drugs, nephrosis etc. In such cases, Free T4 is recommended as it is Metabolically active.

TSH (Thyroid Stimulating Hormone) 1.38 µIU/mL 0.35 - 5.50

(Serum/Chemiluminescent Microparticle

Immunoassay(CMIA))

#### INTERPRETATION:

Reference range for cord blood - upto 20

1 st trimester: 0.1-2.5 2 nd trimester 0.2-3.0 3 rd trimester: 0.3-3.0

(Indian Thyroid Society Guidelines)

#### **Comment:**

- 1.TSH reference range during pregnancy depends on Iodine intake, TPO status, Serum HCG concentration, race, Ethnicity and BMI.
- 2.TSH Levels are subject to circadian variation, reaching peak levels between 2-4am and at a minimum between 6-10PM. The variation can be of the order of 50%, hence time of the day has influence on the measured serum TSH concentrations.
- 3. Values&amplt,0.03 µIU/mL need to be clinically correlated due to presence of rare TSH variant in some individuals.



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Investigation	Observed Value	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
<b>CLINICAL PATHOLOGY</b>			
PHYSICAL EXAMINATION			
Colour (Urine)	Yellow		
Appearance (Urine)	Clear		Clear
Volume (Urine)	15	mL	
CHEMICAL EXAMINATION(Automated- Urineanalyser)			
pH (Urine/AUTOMATED URINANALYSER)	6.5		4.5 - 8.0
Specific Gravity (Urine)	1.010		1.002 - 1.035
Ketones (Urine)	Negative		Negative
Urobilinogen (Urine/AUTOMATED URINANALYSER)	0.2		0.2 - 1.0
Blood (Urine/AUTOMATED URINANALYSER)	Negative		Negative
Nitrite (Urine/AUTOMATED URINANALYSER)	Negative		Negative
Bilirubin (Urine/AUTOMATED URINANALYSER)	Negative		Negative
Protein (Urine)	Negative		Negative



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Glucose (Urine)



Negative

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Investigation	Observed <u>Value</u>	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
Leukocytes (Urine)  MICROSCOPY(URINE DEPOSITS)	Negative	leuco/uL	Negative
Pus Cells (Urine/Flow cytometry)	1-2	/hpf	3-5
Epithelial Cells (Urine)	1-2	/hpf	1-2
RBCs (Urine/Flow cytometry)	Nil	/hpf	2-3
Others (Urine)	Nil		Nil
Casts (Urine/Flow cytometry)	Nil	/hpf	0 - 1
Crystals (Urine)	Nil		NIL

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Investigation BIOCHEMISTRY	Observed Value	<u>Unit</u>	<u>Biological</u> <u>Reference Interval</u>
BUN / Creatinine Ratio	11		6 - 22
Glucose Fasting (FBS) (Plasma - F/GOD - POD)	102	mg/dL	Normal: < 100 Pre Diabetic: 100 - 125 Diabetic: >= 126

**INTERPRETATION:** Factors such as type, quantity and time of food intake, Physical activity, Psychological stress, and drugs can influence blood glucose level.

Glucose Fasting - Urine	+		Negative
(Urine - F)			
Glucose Postprandial (PPBS)	104	mg/dL	70 - 140
(Plasma - PP/GOD - POD)			

### INTERPRETATION:

Factors such as type, quantity and time of food intake, Physical activity, Psychological stress, and drugs can influence blood glucose level. Fasting blood glucose level may be higher than Postprandial glucose, because of physiological surge in Postprandial Insulin secretion, Insulin resistance, Exercise or Stress, Dawn Phenomenon, Somogyi Phenomenon, Anti- diabetic medication during treatment for Diabetes.

Blood Urea Nitrogen (BUN) (Serum/ <i>Urease-GLDH</i> )	11	mg/dL	7.0 - 21
Creatinine (Serum/Jaffe Kinetic)	1.0	mg/dL	0.8 - 1.3

**INTERPRETATION:** Elevated Creatinine values are encountered in increased muscle mass, severe dehydration, Pre-eclampsia, increased ingestion of cooked meat, consuming Protein/ Creatine supplements, Diabetic Ketoacidosis, prolonged fasting, renal dysfunction and drugs such as cefoxitin, cefazolin, ACE inhibitors, angiotensin II receptor antagonists, N-acetylcyteine, chemotherapeutic agent such as flucytosine etc.

Uric Acid 4.7 mg/dL 3.5 - 7.2

(Serum/Uricase/Peroxidase)





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Prostate: > 10.0

Investigation	Observed <u>Value</u>	<u>Unit</u>	<u>Biological</u> Reference Interval
<b>IMMUNOASSAY</b>			
Prostate specific antigen - Total(PSA) (Serum/Chemiluminescent Microparticle Immunoassay(CMIA))	0.874	ng/mL	Normal: 0.0 - 4.0 Inflammatory & Non Malignant conditions of Prostate & genitourinary system: 4.01 - 10.0 Suspicious of Malignant disease of

### INTERPRETATION: Analytical sensitivity: 0.008 - 100 ng/mL

PSA is a tumor marker for screening of prostate cancer. Increased levels of PSA are associated with prostate cancer and benign conditions like bacterial infection, inflammation of prostate gland and benign hypertrophy of prostate/ benign prostatic hyperplasia (BPH).

Transient elevation of PSA levels are seen following digital rectal examination, rigorous physical activity like bicycle riding, ejaculation within 24 hours.

PSA levels tend to increase in all men as they age.

Clinical Utility of PSA:

- •In the early detection of Prostate cancer.
- •As an aid in discriminating between Prostate cancer and Benign Prostatic disease.
- •To detect cancer recurrence or disease progression.



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# **IMMUNOHAEMATOLOGY**

BLOOD GROUPING AND Rh TYPING 'B' 'Negative'

 $({\rm EDTA~Blood} Agglutination)$ 



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**APPROVED BY** 

-- End of Report --