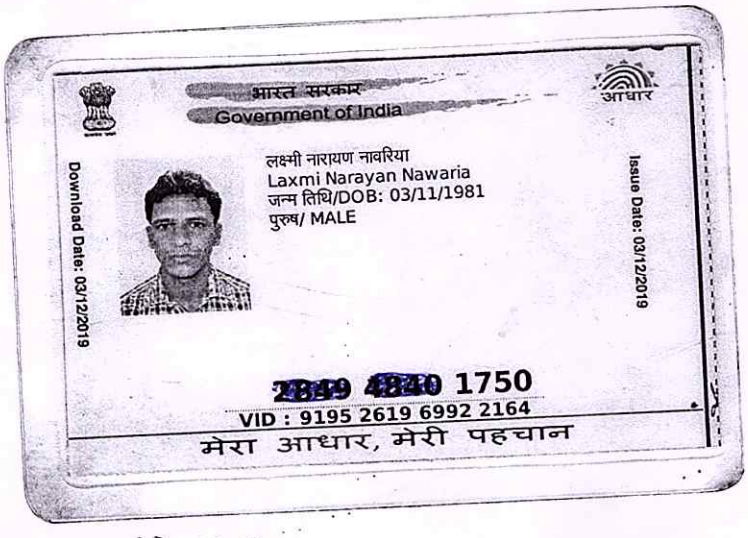


भारत सरकार
Government of India

आधार

लक्ष्मी नारायण नावरिया
Laxmi Narayan Nawaria
जन्म तिथि/DOB: 03/11/1981
पुरुष/ MALE

Download Date: 03/12/2019

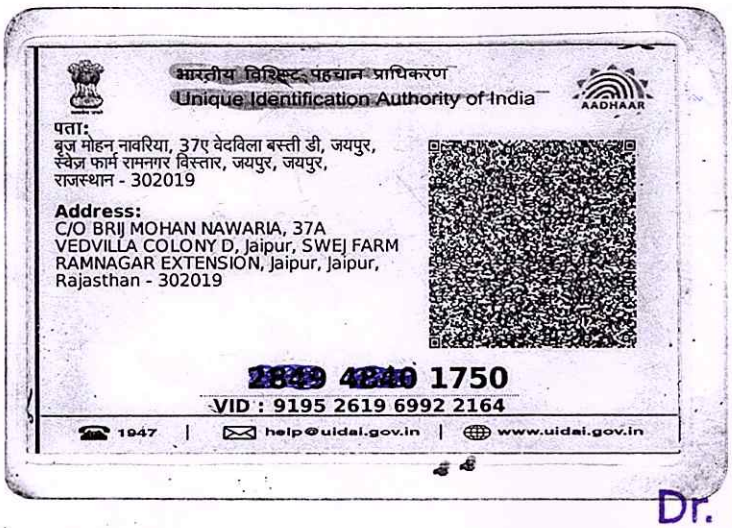
Issue Date: 03/12/2019

[redacted] 1750

VID : 9195 2619 6992 2164

मेरा आधार, मेरी पहचान

Lxmi Nawaria



भारतीय विशिष्ट पहचान प्राधिकरण
Unique Identification Authority of India

AADHAAR

पता:
बृज मोहन नावरिया, 37ए वेदविला बस्ती डी, जयपुर, स्वेज़ फार्म रामनगर विस्तार, जयपुर, जयपुर, राजस्थान - 302019

Address:
C/O BRIJ MOHAN NAWARIA, 37A VEDVILLA COLONY D, Jaipur, SWEJ FARM RAMNAGAR EXTENSION, Jaipur, Jaipur, Rajasthan - 302019

[redacted] 1750

VID : 9195 2619 6992 2164

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Dr. PIYUSH GOYAL
MBBS, DMRD (Radiologist)
RMC No. 037041

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General Physical Examination

Date of Examination: 24/08/2024

Name: Laxmi Narayan Nawaria Age: 42 DOB: 03/11/1981 Sex: Male

Referred By: Bank of Baroda

Photo ID: Adhar Card ID #: 1750

Ht: 170 (cm) Wt: 60 (Kg)

Chest (Expiration): 89 (cm) Abdomen Circumference: 80 (cm)

Blood Pressure: 90/80 mm Hg PR: 78 / min RR: 18 / min Temp: Afebrile

BMI 20.8

Eye Examination: RE-CIG, NIG, NCB
LE-CIG, NIG, NCB

Other: No

On examination he/she appears physically and mentally fit: ~~Yes~~ / No

Signature Of Examine : ---------- Name of Examinee: LAXMI NARAYAN NAWARIA

Signature Medical Examiner : ---------- Name Medical Examiner DR. PIYUSH GOYAL

PIYUSH GOYAL
(Radiologist)
RMC No.-037041

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Patient ID	1224990 Patient Mob No.9166636200	Registered On	24/08/2024 11:31:46
NAME	**Mr. LAXMI NARAYAN NAWARIA**	Collected On	24/08/2024 12:11:44
Age / Sex	Male 42 Yrs 9 Mon 22 Days	Authorized On	25/08/2024 09:21:25
Ref. By	BANK OF BARODA	Printed On	25/08/2024 09:21:35
Lab/Hosp	Mr.MEDIWHEEL		

HAEMOGARAM

HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
FULL BODY HEALTH CHECKUP ABOVE 40 MALE			
HAEMOGLOBIN (Hb)	13.8	g/dL	13.0 - 17.0
TOTAL LEUCOCYTE COUNT	7.60	/cumm	4.00 - 10.00
DIFFERENTIAL LEUCOCYTE COUNT			
NEUTROPHIL	51.6	%	40.0 - 80.0
LYMPHOCYTE	**44.5** H	%	20.0 - 40.0
EOSINOPHIL	1.1	%	1.0 - 6.0
MONOCYTE	2.8	%	2.0 - 10.0
BASOPHIL	0.0	%	0.0 - 2.0
TOTAL RED BLOOD CELL COUNT (RBC)	4.81	x10^6/uL	4.50 - 5.50
HEMATOCRIT (HCT)	42.40	%	40.00 - 50.00
MEAN CORP VOLUME (MCV)	88.0	fL	83.0 - 101.0
MEAN CORP HB (MCH)	28.8	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	32.7	g/dL	31.5 - 34.5
PLATELET COUNT	**122** L	x10^3/uL	150 - 410
RDW-CV	**15.1** H	%	11.6 - 14.0

Technologist

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HAEMATOLOGY

HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
Erythrocyte Sedimentation Rate (ESR) Methord:- Westergreen	**20** H	mm in 1st hr	00 - 15

The erythrocyte sedimentation rate (ESR or sed rate) is a relatively simple, inexpensive, non-specific test that has been used for many years to help detect inflammation associated with conditions such as infections, cancers, and autoimmune diseases.ESR is said to be a non-specific test because an elevated result often indicates the presence of inflammation but does not tell the health practitioner exactly where the inflammation is in the body or what is causing it. An ESR can be affected by other conditions besides inflammation. For this reason, the ESR is typically used in conjunction with other tests, such as C-reactive protein.ESR is used to help diagnose certain specific inflammatory diseases, including temporal arteritis, systemic vasculitis and polymyalgia rheumatica. (For more on these, read the article on Vasculitis.) A significantly elevated ESR is one of the main test results used to support the diagnosis.This test may also be used to monitor disease activity and response to therapy in both of the above diseases as well as

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(CBC): Methodology: TLC,DLC Fluorescent Flow cytometry, HB SLS method,TRBC,PCV,PLT Hydrodynamically focused Impedance. and MCH,MCV,MCHC,MENTZER INDEX are calculated. **InstrumentName:** Sysmex 6 part fully automatic analyzer XN-L,Japan

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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
FASTING BLOOD SUGAR (Plasma) Methord:- GLUCOSE OXIDASE/PEROXIDASE	99.4	mg/dl	70.0 - 115.0

Impaired glucose tolerance (IGT)	111 - 125 mg/dL
Diabetes Mellitus (DM)	> 126 mg/dL

Instrument Name: HORIBA CA60 Interpretation: Elevated glucose levels (hyperglycemia) may occur with diabetes, pancreatic neoplasm,
hyperthyroidism and adrenal cortical hyper-function as well as other disorders.Decreased glucose levels(hypoglycemia) may result from excessive insulin
therapy or various liver diseases .

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HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
GLYCOSYLATED HEMOGLOBIN (HbA1C)			
Methord:- CAPILLARY with EDTA	5.7	mg%	Non-Diabetic < 6.0 Good Control 6.0-7.0 Weak Control 7.0-8.0 Poor control > 8.0
MEAN PLASMA GLUCOSE Methord:- Calculated Parameter	117	mg/dL	68 - 125

INTERPRETATION
AS PER AMERICAN DIABETES ASSOCIATION (ADA)
Reference Group HbA1c in %
Non diabetic adults >=18 years < 5.7
At risk (Prediabetes) 5.7 - 6.4
Diagnosing Diabetes >= 6.5

CLINICAL NOTES
In vitro quantitative determination of HbA1c in whole blood is utilized in long term monitoring of glycemia.The HbA1c level correlates with the mean glucose concentration prevailing in the course of the patient's recent history (approx - 6-8 weeks) and therefore provides much more reliable information for glycemia monitoring than do determinations of blood glucose or urinary glucose. It is recommended that the determination of HbA1c be performed at intervals of 4-6 weeks during Diabetes Mellitus therapy. Results of HbA1c should be assessed in conjunction with the patient's medical history, clinical examinations and other findings.
Some of the factors that influence HbA1c and its measurement [Adapted from Gallagher et al]
1. Erythropoiesis
- Increased HbA1c: iron, vitamin B12 deficiency, decreased erythropoiesis.
- Decreased HbA1c: administration of erythropoietin, iron, vitamin B12, reticulocytosis, chronic liver disease.
2. Altered Haemoglobin: Genetic or chemical alterations in hemoglobin: hemoglobinopathies, HbF, methemoglobin, may increase or decrease HbA1c.
3. Glycation
- Increased HbA1c: alcoholism, chronic renal failure, decreased intraerythrocytic pH.
- Decreased HbA1c: certain hemoglobinopathies, increased intra-erythrocyte pH
.4. Erythrocyte destruction
- Increased HbA1c: increased erythrocyte life span: Splenectomy.
- Decreased A1c: decreased RBC life span: hemoglobinopathies, splenomegaly, rheumatoid arthritis or drugs such as antiretrovirals, ribavirin & dapsone.
5. Others
- Increased HbA1c: hyperbilirubinemia, carbamylated hemoglobin, alcoholism, large doses of aspirin, chronic opiate use,chronic renal failure
- Decreased HbA1c: hypertriglyceridemia,reticulocytosis, chronic liver disease, aspirin, vitamin C and E,splenomegaly, rheumatoid arthritis or drugs

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HAEMATOLOGY

HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
BLOOD GROUP ABO Methord - Haemagglutination reaction	"B" POSITIVE		

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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIPID PROFILE			
SERUM TOTAL CHOLESTEROL Methord - CHOLESTEROL OXIDASE/PEROXIDASE	218.00	mg/dl	Desirable <200 Borderline 200-239 High> 240
InstrumentName HORIBA **Interpretation:** Cholesterol measurements are used in the diagnosis and treatments of lipid lipoprotein metabolism disorders			
SERUM TRIGLYCERIDES Methord - GLYCEROL PHOSPHATE OXIDASE/PREOXIDASE	121.50	mg/dl	Normal <150 Borderline high 150-199 High 200-499 Very high >500
InstrumentName Randox Rx Imola **Interpretation** : Triglyceride measurements are used in the diagnosis and treatment of diseases involving lipid metabolism and various endocrine disorders e.g. diabetes mellitus, nephrosis and liver obstruction.			
DIRECT HDL CHOLESTEROL Methord - Direct clearance Method	45.80	mg/dl	MALE- 30-70 FEMALE - 30-85
Instrument Name Rx Daytona plus **Interpretation:** An inverse relationship between HDL-cholesterol (HDL-C) levels in serum and the incidence/prevalence of coronary heart disease (CHD) has been demonstrated in a number of epidemiological studies Accurate measurement of HDL-C is of vital importance when assessing patient risk from CHD. Direct measurement gives improved accuracy and reproducibility when compared to precipitation methods			
LDL CHOLESTEROL Methord - Calculated Method	**151.95** H	mg/dl	Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190
Interpretation: Accurate measurement of LDL-Cholesterol is of vital importance in therapies which focus on lipid reduction to prevent atherosclerosis or reduce its progress and to avoid plaque rupture.			
VLDL CHOLESTEROL Methord - Calculated	24.30	mg/dl	0.00 - 80.00

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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
T.CHOLESTEROL/HDL CHOLESTEROL RATIO Method:- Calculated	4.76		0.00 - 4.90
LDL / HDL CHOLESTEROL RATIO Method - Calculated	3.32		0.00 - 3.50
TOTAL LIPID Method:- CALCULATED	633.80	mg/dl	400.00 - 1000.00

1. Measurements in the same patient can show physiological& analytical variations. Three serialsamples 1 week apart are recommended for Total Cholesterol, Triglycerides, HDL& LDL Cholesterol.
2. As per NCEP guidelines, all adults above the age of 20 years should be screened for lipid status.Selective screening of children above the age of 2 years with a family history of premature cardiovascular disease or those with at least one parent with high total cholesterol is recommended
3. Low HDL levels are associated with Coronary Heart Disease due to insufficient HDL being available to participate in reverse cholesterol transport, the process by which cholesterol is eliminated fromperipheral tissues.

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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIVER PROFILE WITH GGT			
SERUM BILIRUBIN (TOTAL) Method:- DIAZOTIZED SULFANILIC	0.62	mg/dL	Infants : 0.2-8.0 mg/dL Adult - Up to - 1.2 mg/dL
SERUM BILIRUBIN (DIRECT) Method:- DIAZOTIZED SULFANILIC	0.16	mg/dL	Up to 0.40 mg/dL
SERUM BILIRUBIN (INDIRECT) Method:- Calculated	0.46	mg/dl	0.30-0.70
SGOT Method - IFCC	29.5	U/L	0.0 - 40.0
SGPT Method - IFCC	22.6	U/L	0.0 - 40.0
SERUM ALKALINE PHOSPHATASE Method - DGKC - SCE	70.00	U/L	53.00 - 141.00
SERUM GAMMA GT Method:- Szasz methodology Instrument Name Rapidex By Imola Interpretation: Elevations in GGT levels areseen earlier and more pronounced than those with other liver enzymes in cases of obstructive jaundice and metastatic neoplasms. It may reach 5 to 30 times normal levels in intra-or post-hepatic biliary obstruction. Only moderate elevations in the enzyme level (2 to 5 times normal)are observed with infectious hepatitis.	17.00	U/L	10.00 - 45.00
SERUM TOTAL PROTEIN Method:- BIURET	6.73	g/dl	6.00 - 8.40
SERUM ALBUMIN Method:- BROMOCRESOL GREEN	4.84	g/dl	3.50 - 5.50
SERUM GLOBULIN Method:- CALCULATION	**1.89** L	gm/dl	2.20 - 3.50
A/G RATIO	**2.56** H		1.30 - 2.50

Interpretation : Measurements obtained by this method are used in the diagnosis and treatment of a variety of diseases involving the liver, kidney and bone marrow as well as other metabolic or nutritional disorders.

Note :- These are group of tests that can be used to detect the presence of liver disease, distinguish among different types of liver disorders, gauge the extent of known liver damage, and monitor the response to treatment. Most liver diseases cause only mild symptoms initially, but these diseases must be detected early. Some tests are associated with functionality (e.g.,

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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval

albumin), some with cellular integrity (e.g., transaminase), and some with conditions linked to the biliary tract (gamma-glutamyl transferase and alkaline phosphatase). Conditions with elevated levels of ALT and AST include hepatitis A,B ,C ,paracetamol toxicity etc Several biochemical tests are useful in the evaluation and management of patients with hepatic dysfunction. Some or all of these measurements are also carried out (usually about twice a year for routine cases) on those individuals taking certain medications, such as anticonvulsants, to ensure that the medications are not adversely impacting the person's liver.

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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
RFT / KFT WITH ELECTROLYTES			
SERUM UREA Method:- UREASE/ GLUTAMATE DEHYDROGENASE	23.90	mg/dl	10.00 - 50.00
InstrumentName: HORIBA CA 60 **Interpretation** : Urea measurements are used in the diagnosis and treatment of certain renal and metabolic diseases.			
SERUM CREATININE Method:- JAFFE	0.69	mg/dl	Males : 0.6-1.50 mg/dl Females : 0.6 -1.40 mg/dl
Interpretation : Creatinine is measured primarily to assess kidney function and has certain advantages over the measurement of urea. The plasma level of creatinine is relatively independent of protein ingestion, water intake, rate of urine production and exercise. Depressed levels of plasma creatinine are rare and not clinically significant.			
SERUM URIC ACID Method:- URICASE/PEROXIDASE	6.60	mg/dl	2.40 - 7.00
InstrumentName:HORIBA YUMIZEN CA60 Daytona plus **Interpretation**: **Elevated Urate**:High purine diet,Alcohol• Renal insufficiency,Drugs , Polycythaemia vera, Malignancies,Hypothyroidism,Rare enzyme defects ,Downs syndrome,Metabolic syndrome, Pregnancy,Gout.			
SODIUM Method:- Ion-Selective Electrode with Serum	139.7	mmol/L	135.0 - 145.0
POTASSIUM Method:- Ion-Selective Electrode with Serum	**3.44** L	mmol/L	3.50 - 5.00
CHLORIDE Method:- Ion-Selective Electrode with Serum	103.3	mmol/L	97.0 - 107.0
SERUM CALCIUM Method:- Arsenazo III Method	10.00	mg/dL	8.80 - 10.20
InstrumentName:MISPA PLUS **Interpretation**: Serum calcium levels are believed to be controlled by parathyroid hormone and vitamin D. Increases in serum PTH or vitamin D are usually associated with hypercalcemia .Hypocalcemia may be observed in hypoparathyroidism, nephrosis and pancreatitis.			
SERUM TOTAL PROTEIN Method:- BIURET	6.73	g/dl	6.00 - 8.40
SERUM ALBUMIN Method:- BROMOCRESOL GREEN	4.84	g/dl	3.50 - 5.50

Technologist
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BIOCHEMISTRY

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
SERUM GLOBULIN Method - CALCULATION	**1.89** L	gm/dl	2.20 - 3.50
A/G RATIO	**2.56** H		1.30 - 2.50

Interpretation : Measurements obtained by this method are used in the diagnosis and treatment of a variety of diseases involving the liver, kidney and bone marrow as well as other metabolic or nutritional disorders.

INTERPRETATION

Kidney function tests are group of tests that can be used to evaluate how well the kidneys are functioning. Creatinine is a waste product that comes from protein in the diet and also comes from the normal wear and tear of muscles of the body. In blood, it is a marker of GFR .in urine, it can remove the need for 24-hourcollections for many analytes or be used as a quality assurance tool to assess the accuracy of a 24-hour collection Higher levels may be a sign that the kidneys are not working properly. As kidney disease progresses, the level of creatinine and urea in the blood increases. Certain drugs are nephrotoxic hence KFT is done before and after initiation of treatment with these drugs.

Low serum creatinine values are rare; they almost always reflect low muscle mass

Apart from renal failure Blood Urea can increase in dehydration and GI bleed

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Lab/Hosp	Mr MEDIWHEEL		

CLINICAL PATHOLOGY

CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
URINE SUGAR (FASTING) Collected Sample Received	Nil		Nil
URINE SUGAR PP Collected Sample Received	Nil		Nil

Technologist

DR.TANU RUNGTA
MD (Pathology)
RMC No. 17226

P3 HEALTH SOLUTIONS LLP

B-14, Vidhyadhar Nagar Enclave-II, Near Axis Bank
Central Spine, Vidhyadhar Nagar, Jaipur-302 023

+91 141 4824885 p3healthsolutionsllp@gmail.com

Patient ID	1224990 Patient Mob No.9166636200	Registered On	24/08/2024 11:31:46
NAME	Mr. LAXMI NARAYAN NAWARIA	Collected On	24/08/2024 12:11:44
Age / Sex	Male 42 Yrs 9 Mon 22 Days	Authorized On	25/08/2024 09:21:25
Ref. By	BANK OF BARODA	Printed On	25/08/2024 09:21:35
Lab/Hosp	Mr.MEDIWHEEL		

IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
PSA (PROSTATE SPECIFIC ANTIGEN) -TOTAL Method:- Methodology CLIA	2.190	ng/mL	0.00-4.00

CLINICAL NOTES - Prostate-specific antigen (PSA)is a 34-kD glycoprotein produced almost exclusively by the prostate gland.

PSA is normally present in the blood at very low levels. Increased levels of PSA may suggest the presence of prostate cancer.

1.Immediate PSA testing following digital rectal examination, ejaculation, prostatic massage, indwelling catheterization, ultrasonography and needle biopsy of prostate is not
recommended as they falsely elevate levels

2. PSA values regardless of levels should not be interpreted as absolute evidence of the presence or absence of disease. All values should be correlated with clinical
findings and other investigations

3. Physiological decrease in PSA level by 18% has been observed in sedentary patients either due to supine position or suspended sexual activity

Clinical Use

• An aid in the early detection of Prostate cancer when used in conjunction with Digital rectal examination in males more than 50 years of age and in those with two or more
affected first degree relatives.
• Follow up and management of Prostate cancer patients
• Detect metastatic or persistent disease in patients following surgical or medical treatment of Prostate cancer

NOTE

PSA levels can be also increased by prostatitis, irritation, benign prostatic hyperplasia (BPH), and recent ejaculation, producing a false positive result. Digital rectal examination (DRE) has been shown in several studies to produce an increase in PSA. However, the effect is clinically insignificant, since DRE causes the most substantial
increases in patients with PSA levels already elevated over 4.0 ng/mL.

Obesity has been reported to reduce serum PSA levels. Delayed early detection may partially explain worse outcomes in obese men with early prostate cancer. Aftertreatment, higher BMI also correlates to higher risk of recurrence.

Technologist

Page 1 of 1

DR.TANU RUNGTA
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IMMUNOASSAY

IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
TOTAL THYROID PROFILE			
THYROID-TRIIODOTHYRONINE T3 Methord - Chemiluminescence	1.38	ng/ml	0.69 - 2.15
THYROID - THYROXINE (T4) Methord - Chemiluminescence	5.89	ug/dl	5.20 - 12.70
TSH Methord - Chemiluminescence	4.470	µIU/mL	0.470 - 4.680

Note:
1. TSH levels are subject to circadian variation, reaching peak levels between 2 - 4.a.m. and at a minimum between 6-10 pm . The variation is of the order of 50% . hence time of the day has influence on the measured serum TSH concentrations.
2. Recommended test for T3 and T4 is unbound fraction or free levels as it is metabolically active.
3. Physiological rise in Total T3 / T4 levels is seen in pregnancy and in patients on steroid therapy.

Clinical Use
1. in infancy and early childhood

*** End of Report ***

*** End of Report ***

Technologist

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Lab/Hosp	Mr.MEDIWHEEL		

CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
Urine Routine			
PHYSICAL EXAMINATION			
COLOUR	PALE YELLOW		PALE YELLOW
APPEARANCE	Clear		Clear
CHEMICAL EXAMINATION			
REACTION(PH)	7.0		5.0 - 7.5
SPECIFIC GRAVITY	1.020		1.010 - 1.030
PROTEIN	NIL		NIL
SUGAR	NIL		NIL
BILIRUBIN	NEGATIVE		NEGATIVE
UROBILINOGEN	NORMAL		NORMAL
KETONES	NEGATIVE		NEGATIVE
NITRITE	NEGATIVE		NEGATIVE
MICROSCOPY EXAMINATION			
RBC/HPF	NIL	/HPF	NIL
WBC/HPF	2-3	/HPF	2-3
EPITHELIAL CELLS	0-2	/HPF	2-3
CRYSTALS/HPF	ABSENT		ABSENT
CAST/HPF	ABSENT		ABSENT
AMORPHOUS SEDIMENT	ABSENT		ABSENT
BACTERIAL FLORA	ABSENT		ABSENT
YEAST CELL	ABSENT		ABSENT
OTHER	ABSENT		

Technologist

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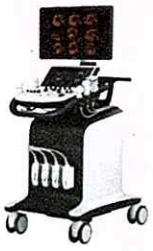


NAME:	MR. LAXMI NARAYAN NAWARIA	AGE	42 YRS/M
REF.BY	BANK OF BARODA	DATE	24/08/2024

CHEST X RAY (PA VIEW)

Bilateral lung fields appear clear.

Bilateral costo-phrenic angles appear clear.

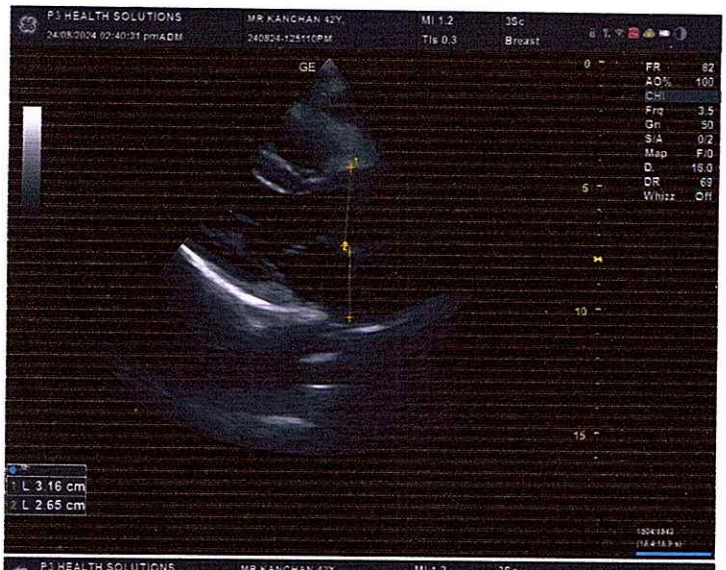
Cardiothoracic ratio is normal.

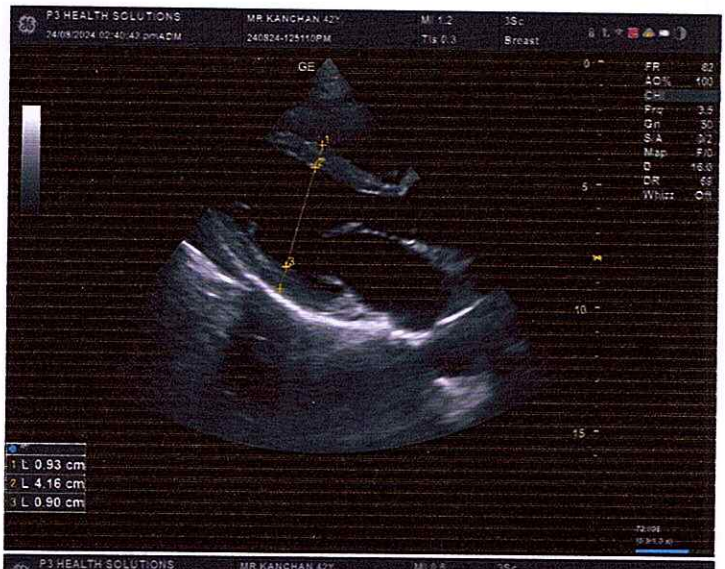
Thoracic soft tissue and skeletal system appear unremarkable.

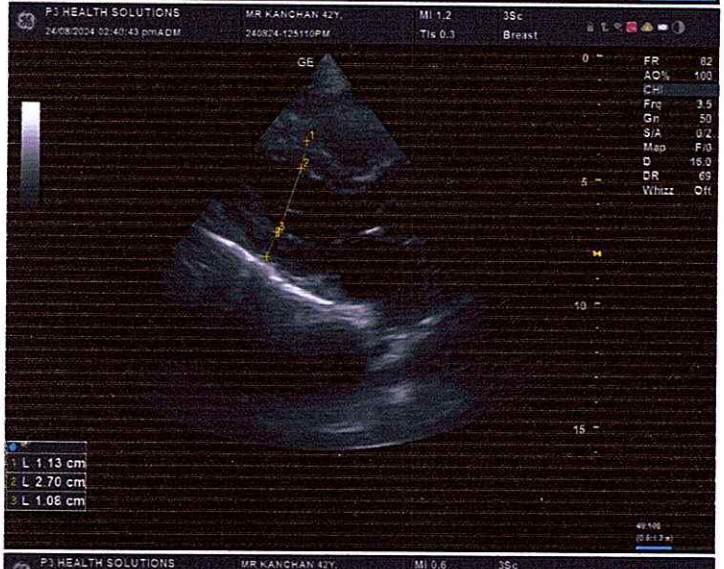
Soft tissue shadows appear normal.

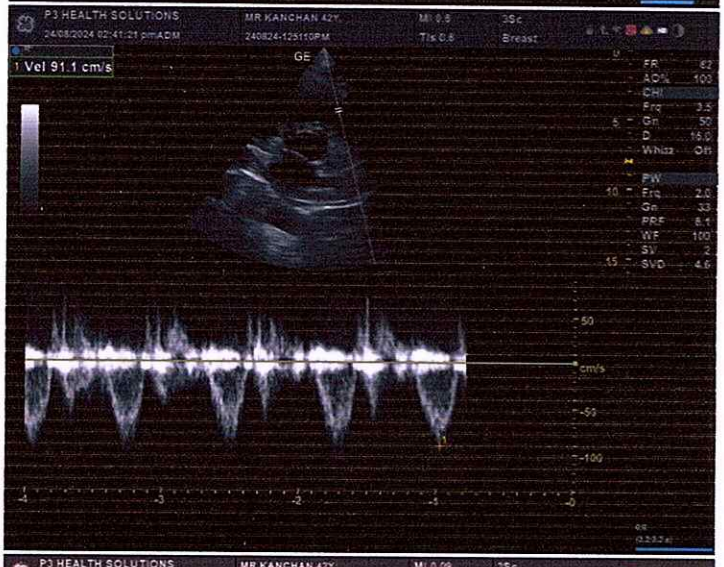
IMPRESSION: No significant abnormality is detected

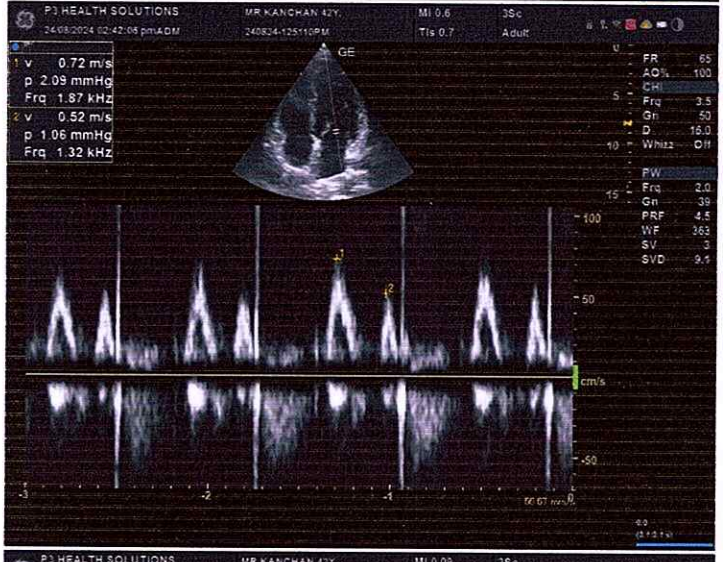
DR. ROHAN GAUR
M.B.B.S, M.D (Radiodiagnosis)
RMC no. 17887

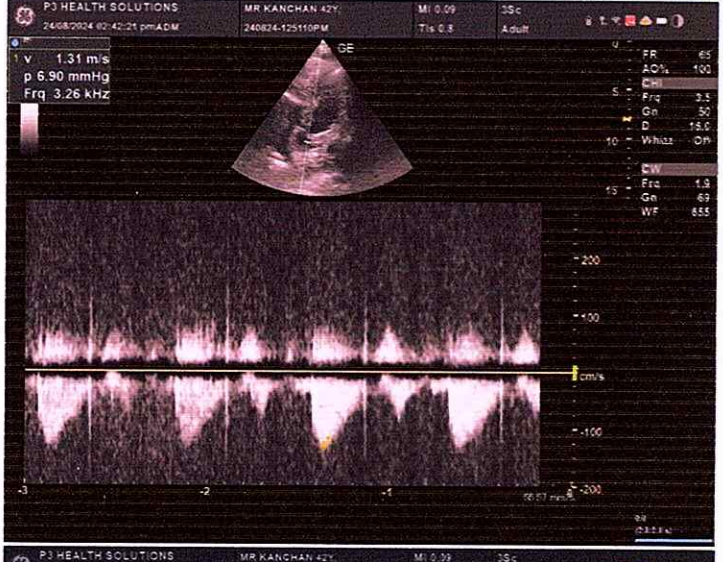

P3 HEALTH SOLUTIONS
MR KANCHAN 42Y,
240824-125110PM
Breast
GE
L 3.16 cm
L 2.65 cm

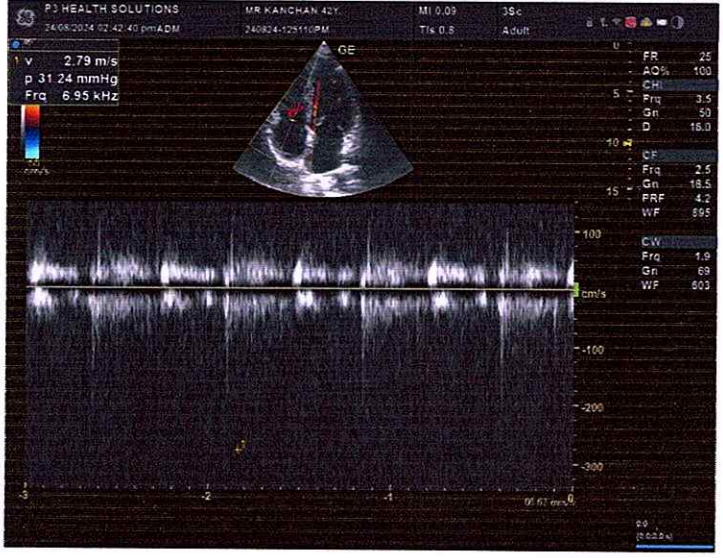

P3 HEALTH SOLUTIONS
MR KANCHAN 42Y,
240824-125110PM
Breast
GE
L 0.93 cm
L 4.16 cm
L 0.90 cm

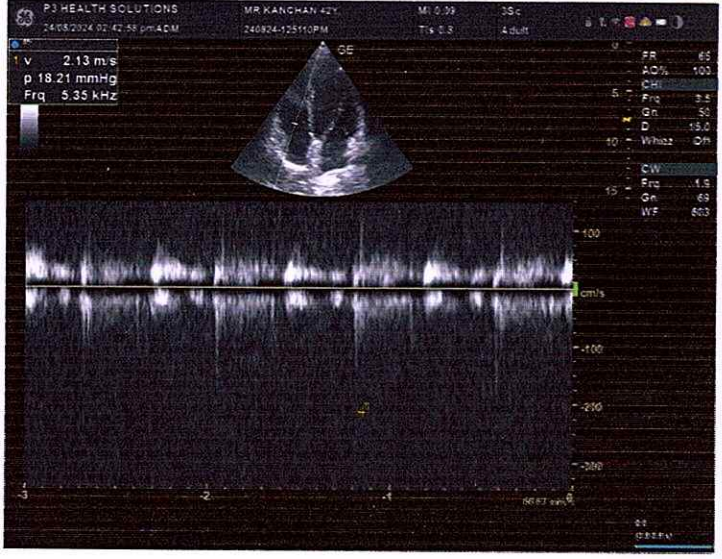

P3 HEALTH SOLUTIONS
MR KANCHAN 42Y,
240824-125110PM
Breast
GE
L 1.13 cm
L 2.70 cm
L 1.08 cm


P3 HEALTH SOLUTIONS
MR KANCHAN 42Y,
240824-125110PM
Breast
GE
Vel 91.1 cm/s
cm/s


P3 HEALTH SOLUTIONS
MR KANCHAN 42Y,
240824-125110PM
Adult
GE
v 0.72 m/s
p 2.09 mmHg
Frq 1.87 kHz
v 0.52 m/s
p 1.06 mmHg
Frq 1.32 kHz
cm/s


P3 HEALTH SOLUTIONS
MR KANCHAN 42Y,
240824-125110PM
Adult
GE
v 1.31 m/s
p 6.90 mmHg
Frq 3.26 kHz
cm/s


P3 HEALTH SOLUTIONS
MR KANCHAN 42Y,
240824-125110PM
Adult
GE
v 2.79 m/s
p 31.24 mmHg
Frq 6.95 kHz
cm/s


P3 HEALTH SOLUTIONS
MR KANCHAN 42Y,
240824-125110PM
Adult
GE
v 2.13 m/s
p 18.21 mmHg
Frq 5.35 kHz
cm/s

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MR. LAXMI NARAYAN NAWARIA	42 Y/M
Registration Date: 24/08/2024	Ref. by: BANK OF BARODA

2D-ECHOCARDIOGRAPHY M.MODE WITH DOPPLER STUDY:

FAIR TRANSTHORACIC ECHOCARIDIOGRAPHIC WINDOW MORPHOLOGY:

MITRAL VALVE	NORMAL	TRICUSPID VALVE	NORMAL
AORTIC VALVE	NORMAL	PULMONARY VALVE	NORMAL

M.MODE EXAMITATION:

AO	2.6	Cm	LA	2.7	cm	IVS-D	0.8	cm
IVS-S	1.2	cm	LVID	4.2	cm	LVSD	2.7	cm
LVPW-D	0.8	cm	LVPW-S	1.2	cm	RV		cm
RVWT		cm	EDV		Ml	LVVS		ml
LVEF	55-60%			RWMA		ABSENT		

CHAMBERS:

LA	NORMAL	RA	NORMAL
LV	NORMAL	RV	NORMAL
PERICARDIUM		NORMAL	

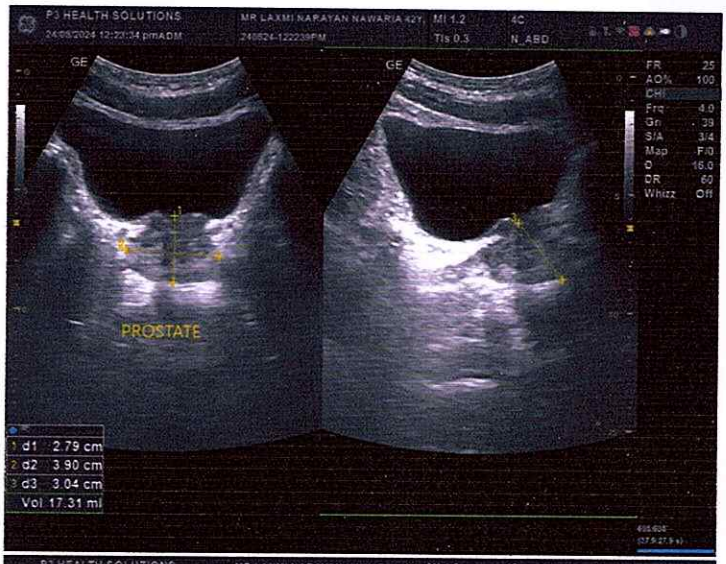
COLOUR DOPPLER:

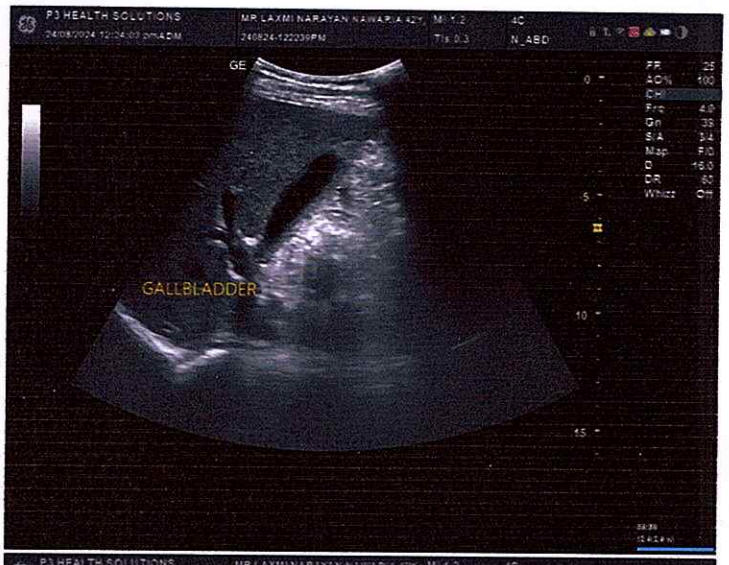
MITRAL VALVE					
E VELOCITY	0.97	m/sec	PEAK GRADIENT		Mm/hg
A VELOCITY	0.68	m/sec	MEAN GRADIENT		Mm/hg
MVA BY PHT		Cm2	MVA BY PLANIMETRY		Cm2
MITRAL REGURGITATION			ABSENT		
AORTIC VALVE					
PEAK VELOCITY	1.47	m/sec	PEAK GRADIENT		mm/hg
AR VMAX		m/sec	MEAN GRADIENT		mm/hg
AORTIC REGURGITATION			ABSENT		
TRICUSPID VALVE					
PEAK VELOCITY		m/sec	PEAK GRADIENT		mm/hg
MEAN VELOCITY		m/sec	MEAN GRADIENT		mm/hg
VMax VELOCITY					
TRICUSPID REGURGITATION			ABSENT		
PULMONARY VALVE					
PEAK VELOCITY	1.04	M/sec.	PEAK GRADIENT		Mm/hg
MEAN VALOCITY			MEAN GRADIENT		Mm/hg
PULMONARY REGURGITATION			ABSENT		

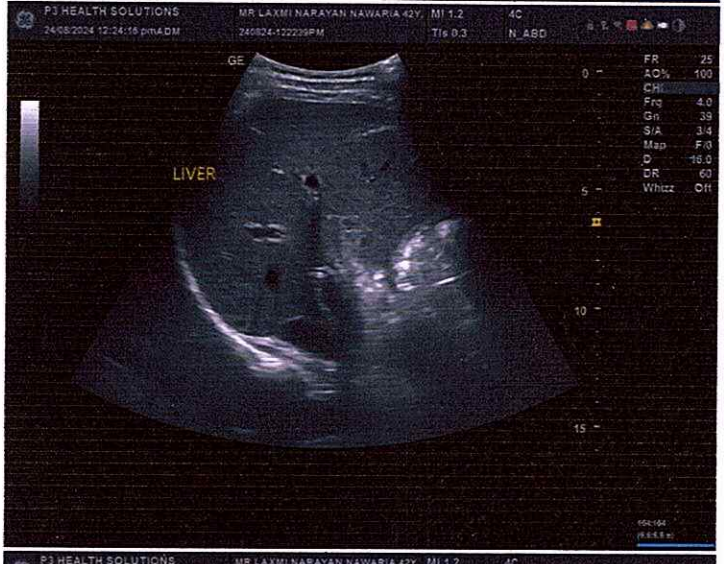
Impression—

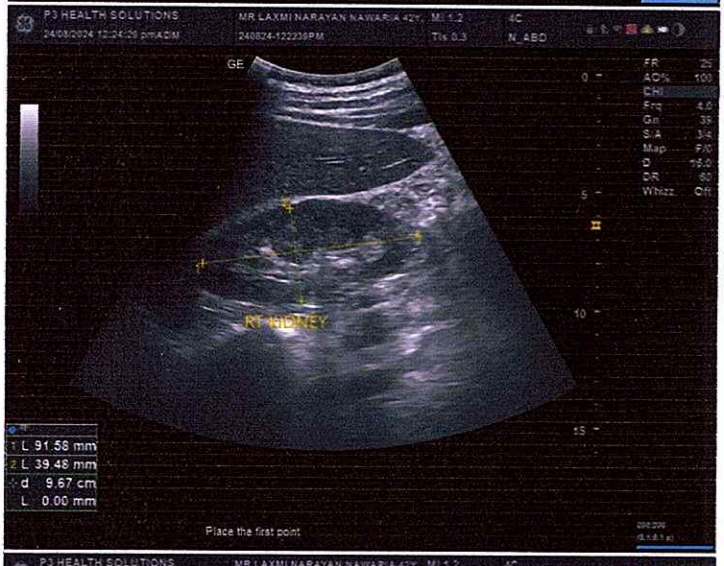
- **NORMAL LV SIZE & CONTRACTILITY.**
- **NO RWMA, LVEF 55-60%.**
- **NORMAL DIASTOLIC FUNCTION.**
- **ALL CARDIAC VALVES ARE NORMAL.**
- **NO CLOT, NO VEGETATION, NO PERICARDIAL EFFUSION.**

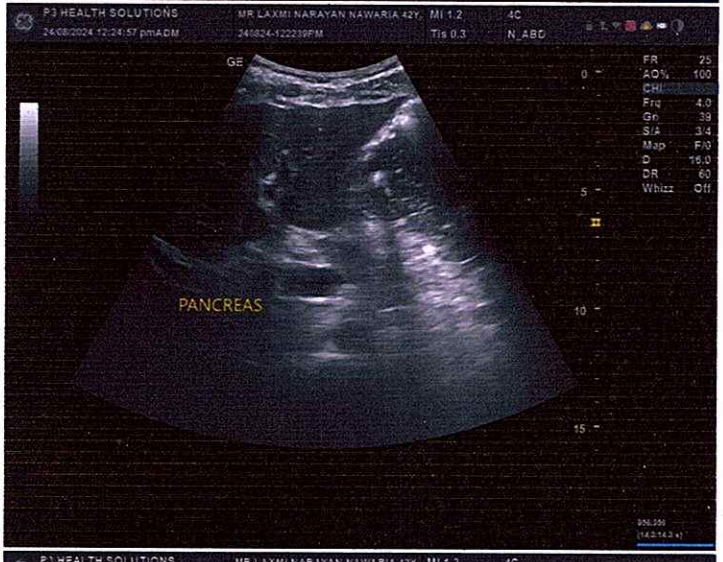
(Cardiologist)

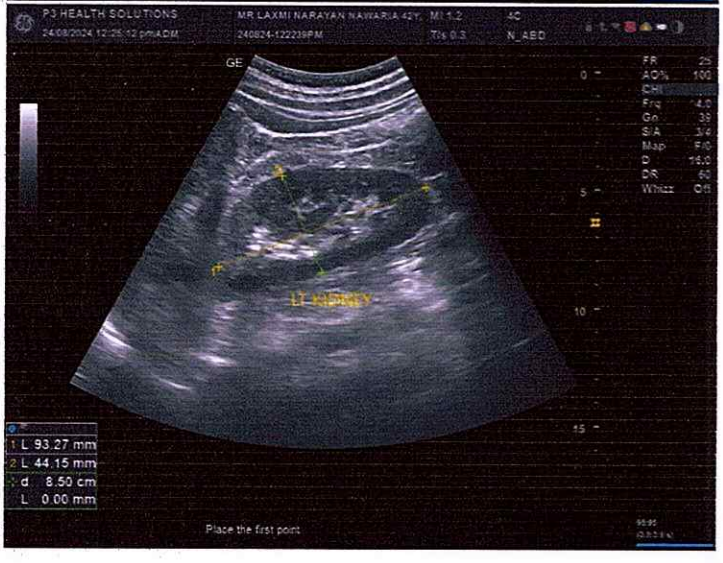

PROSTATE
d1 2.79 cm
d2 3.90 cm
d3 3.04 cm
Vol 17.31 ml

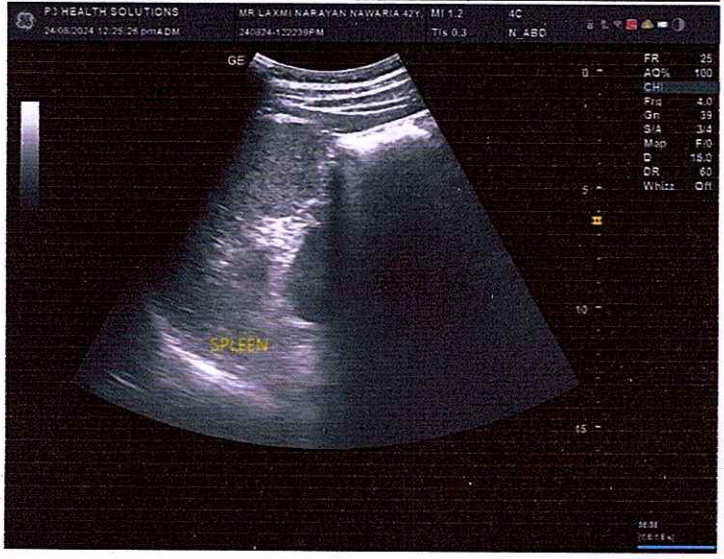

GALLBLADDER


LIVER


RT KIDNEY
L 91.58 mm
L 39.48 mm
d 9.67 cm
L 0.00 mm
Place the first point


PANCREAS


LT KIDNEY
L 93.27 mm
L 44.15 mm
d 8.50 cm
L 0.00 mm
Place the first point


SPLEEN

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MR. LAXMI NARAYAN NAWARIA	42 Y/M
Registration Date: 24/08/2024	Ref. by: BANK OF BARODA

ULTRASOUND OF WHOLE ABDOMEN

Liver is of normal size (13.0 cm). Echo-texture is normal. No focal space occupying lesion is seen within liver parenchyma. Intra hepatic biliary channels are not dilated. Portal vein diameter is normal.

Gall bladder is well distended. Wall is not thickened. No calculus or mass lesion is seen in gall bladder. Common bile duct is not dilated.

Pancreas is of normal size and contour. Echo-pattern is normal. No focal lesion is seen within pancreas.

Spleen is of normal size and shape. Echotexture is normal. No focal lesion is seen.

Kidneys are normally sited and are of normal size and shape. Cortico-medullary echoes are normal. Collecting system does not show any calculus or dilatation.
Right kidney is measuring approx. 9.1 x 3.9 cm.
Left kidney is measuring approx. 9.3 x 4.3 cm.

Urinary bladder is well distended and does not show any calculus or mass lesion.

Prostate is normal in size (17.0 cc) with normal echotexture and outline.

No enlarged nodes are visualized. No retro-peritoneal lesion is identified.
No significant free fluid is seen in pelvis.

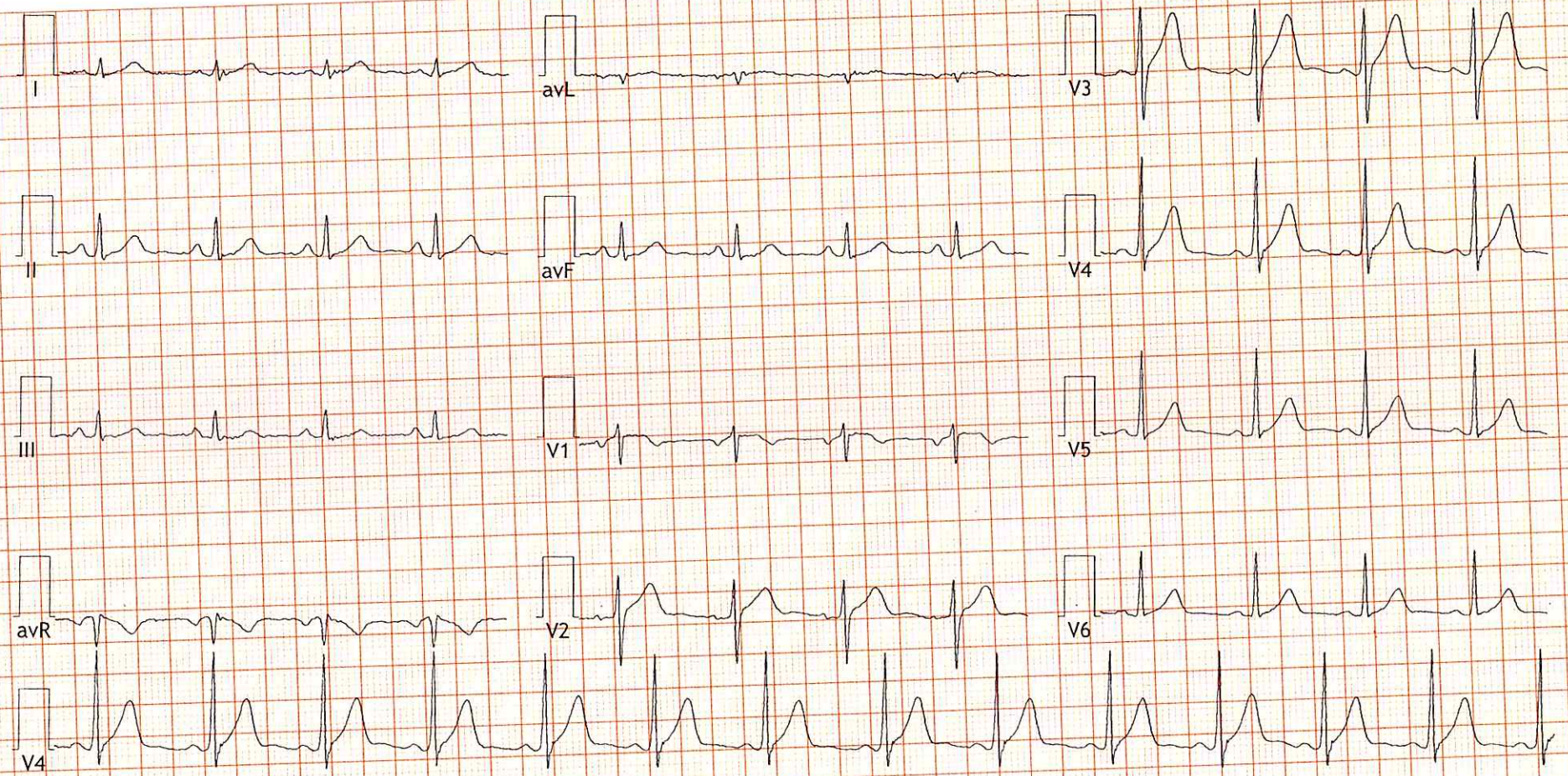
IMPRESSION:-

- **No significant abnormality is detected.**

DR. ROHAN GAUR
M.B.B.S, M.D (Radiodiagnosis)
RMC no. 17887

(P3 HEALTH SOLUTION LLP)
B-14 VIDHYDHAR NAGAR ,(JAIPUR)
1225202/Laxmi Narayan Nawaria 42Yrs-7Months/Male Kgs/ Cms BP: ___/___ mmHg HR: 80 bpm
Ref.: BANK OF BARODA Test Date: 24-Aug-2024(3:31:10 P) Notch: 50Hz 0.05Hz - 100Hz 10mm/mV 25mm/Sec

PR Interval: 118 ms
QRS Duration: 114 ms
QT/QTc: 361/419ms
P-QRS-T Axis: 75 - 72 - 46 (Deg)



TWNL

FINDINGS: Normal Sinus Rhythm
Vent Rate : 80 bpm; PR Interval : 118 ms; QRS Duration: 114 msQT/QTc Int : 361/419 ms
P-QRS-T axis: 75• 72• 46• (Deg)
Comments :

Dr. Naresh Kumar Mohanka
RMC No.: 35703
MBBS, DIP. CARDIO (ESCORTS)
D.E.M. (RCGP,UK)
Dr.NARESH MOHNKA