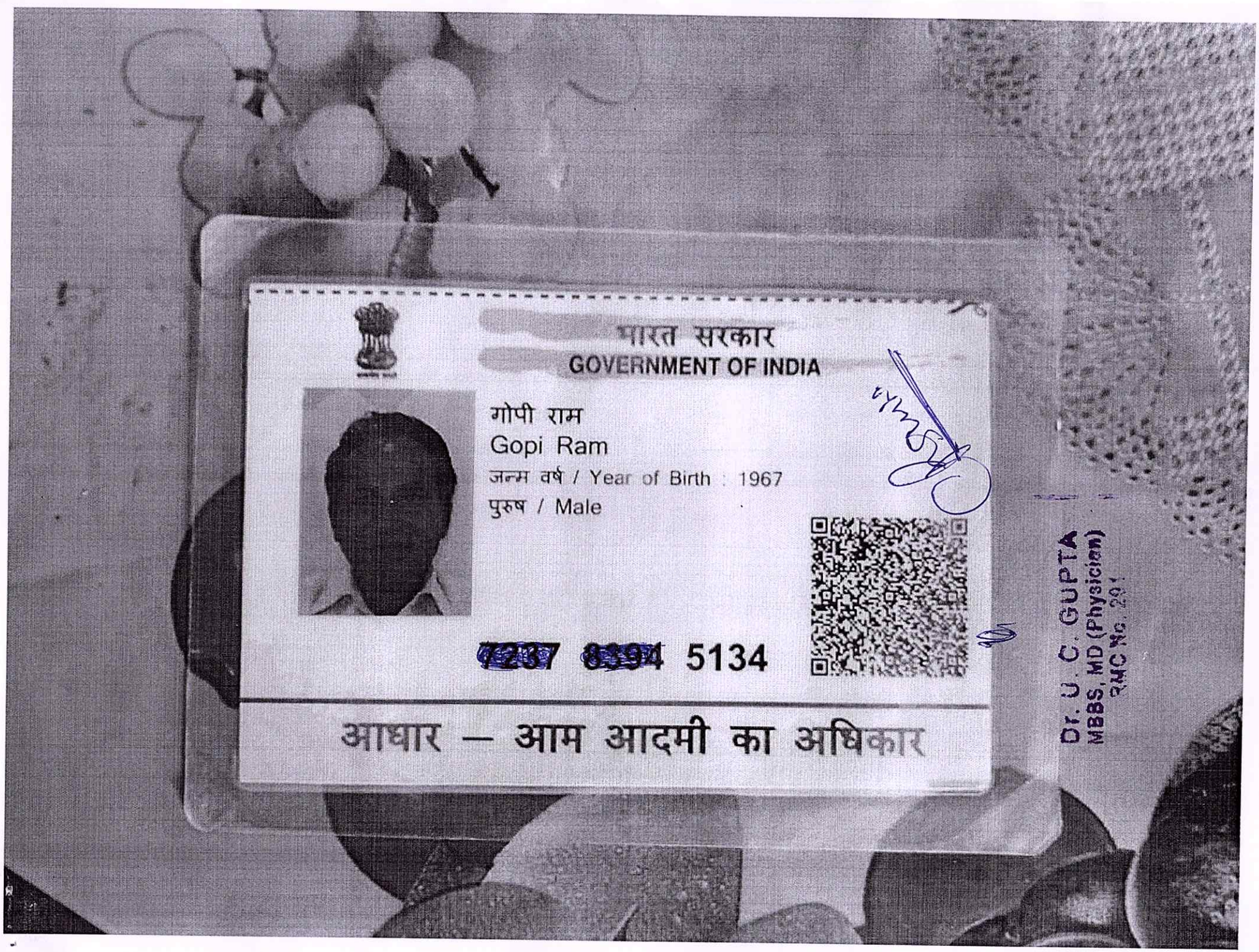
भारत सरकार
GOVERNMENT OF INDIA
गोपी राम
Gopi Ram
जन्म वर्ष / Year of Birth : 1967
पुरुष / Male
7237 8394 5134
आधार — आम आदमी का अधिकार
Dr. U. C. GUPTA
MBBS, MD (Physician)
RMC No. 20

P3 HEALTH SOLUTIONS LLP
(ASSOCIATES OF MAXCARE DIAGNOSTICS)

B-14, Vidhyadhar Enclave - II, Near Axis Bank
Central Spine, Vidhyadhar Nagar, Jaipur - 302023
+91 141 4824885 maxcarediagnostics1@gmail.com

General Physical Examination

Date of Examination: 02/12/2022

Name: GOPI RAM Age: 55 DOB: 05/06/1967 Sex: MALE

Referred By: BANK of BARODA

Photo ID: AADHAR ID #: 5134

Ht: 159 (cm) Wt: 69 (Kg)

Chest (Expiration): 98 (cm) Abdomen Circumference: 108 (cm)

Blood Pressure: 175/93 mm Hg PR: 78 / min RR: 17 / min Temp: _____

BMI 27

Eye Examination: With Glass R/E- 6/6, N6, NCB
L/E- 6/6, N, 6, NCB

Other: N/A

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

Signature Of Examine : _____ **Name of Examinee:** GOPI RAM

Signature Medical Examiner : _____ **Name Medical Examiner** DR. UC GUPTA

Dr. U. C. GUPTA
MBBS, MD (Physician)
RMC No. 291

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NAME:	MR. GOPI RAM	AGE	55 YRS/M
REF.BY	DR. BANK OF BARODA	DATE	02-12-2022

CHEST X RAY (PA VIEW)

Bilateral lung fields appear clear.

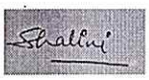
Bilateral costo-phrenic angles appear clear.

Cardiothoracic ratio is normal.

Thoracic soft tissue and skeletal system appear unremarkable.

Soft tissue shadows appear normal.

IMPRESSION: No significant abnormality is detected.



DR.SHALINI GOEL
M.B.B.S, D.N.B (Radiodiagnosis)
RMC No.: 21954

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+91 141 4824885 maxcarediagnostics1@gmail.com

MR. GOPI RAM	**55 Y/Male**
Registration Date:02/12/2022	**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	2.8	Cm	LA	2.8	cm	IVS-D	1.2	cm
IVS-S	1.4	cm	LVID	4.1	cm	LVSD	2.9	cm
LVPW-D	1.2	cm	LVPW-S	1.4	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	0.54	m/sec	PEAK GRADIENT		Mm/hg
A VELOCITY	0.83	m/sec	MEAN GRADIENT		Mm/hg
MVA BY PHT		Cm2	MVA BY PLANIMETRY		Cm2
MITRAL REGURGITATION			MILD		
AORTIC VALVE					
PEAK VELOCITY	1.50	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			MILD		
PULMONARY VALVE					
PEAK VELOCITY	0.64	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%.**
- **MILD TR/ PAH (RVSP 33 MMHG+ RAP), MILD MR.**
- **GRADE- 1 DIASTOLIC DYSFUNCTION.**
- **CONCENTRIC LVH.**
- **NO CLOT, NO VEGETATION, NO PERICARDIAL EFFUSION.**

(Cardiologist)

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Mr. GOPI RAM	**AGE- 55 Y/Male**
Registration Date: 02/12/2022	**Ref. by: DR. BANK OF BARODA**

ULTRA SOUND WHOLE ABDOMEN

Liver is of normal size (12.2 cm). Echo-texture is normal. No focal space occupying lesion is seen within liver parenchyma. Intrahepatic biliary channels are not dilated. Portal vein diameter is normal.

Gall bladder is well distended. 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 (9.5 cm) and shape. Echotexture is normal. No focal lesion is seen.

Transplanted Right kidney is noted in right iliac fossa region, measuring approx. 10.0 x 4.9 cm. Renal artery and vein appear patent with maintained cortico-medullary differentiation. Cortical thickness is normal (25-26 mm).
Right kidney is measuring approx. 6.8 x 3.4 cm. **Cortical echogenicity is increased with poor visibility of renal pyramids and sinus, loss of cortico-medullary differentiation, marginal irregularities and reduced renal length.** Collecting system does not show any dilatation.

Few simple exophytic cortical cysts are noted at inter-polar region, largest measuring 14 x 12 mm.

Left kidney is measuring approx. 7.0 x 3.3 cm. **Cortical echogenicity is increased with poor visibility of renal pyramids and sinus, loss of cortico-medullary differentiation, marginal irregularities and reduced renal length.** Collecting system does not show any dilatation.

Urinary bladder does not show any calculus or mass lesion.

Prostate is normal in size (2.8 x 4.4 x 2.9 cm volume-19-20 cc) with normal echotexture and outline.

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

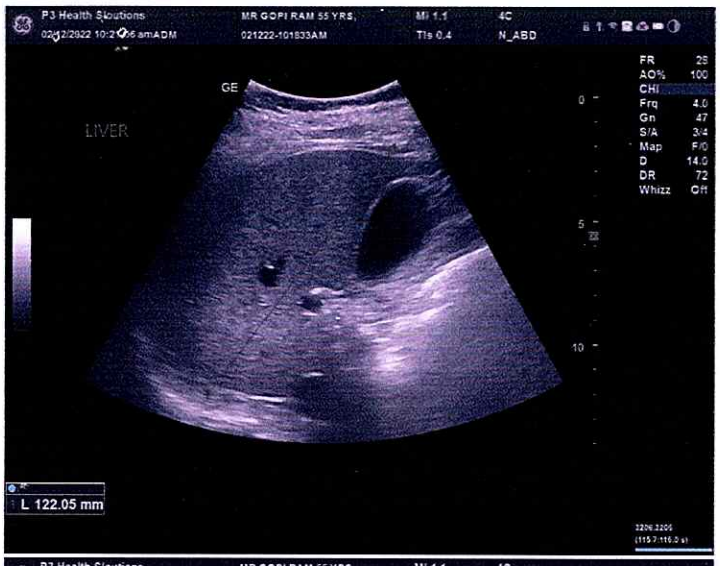
IMPRESSION:

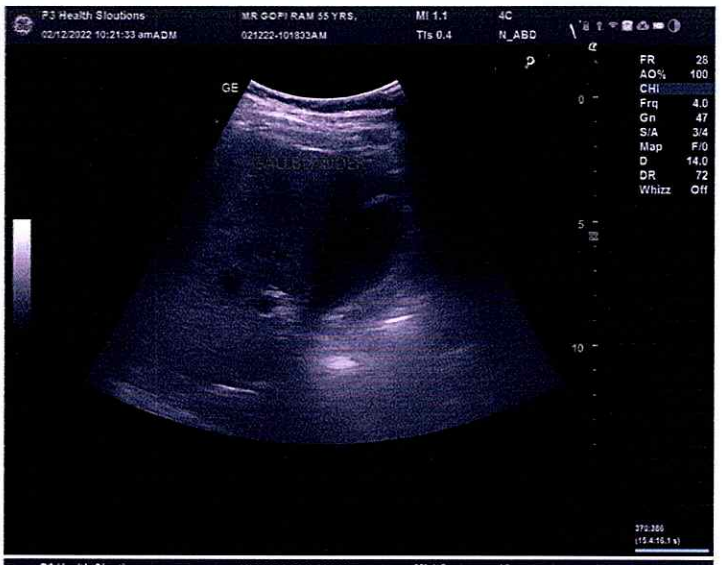
- **Bilateral grade 5 CKD with transplanted right kidney as described above.**

(Please correlate clinically)

DR.SHALINI GOEL

M.B.B.S, D.N.B (Radiodiagnosis)


P3 Health Sloutions
MR GOPI RAM 55 YRS,
MI 1.1
4C
02/12/2022 10:21:06 amADM
021222-101833AM
TIs 0.4
N_ABD
LIVER
L 122.05 mm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MI 1.1
4C
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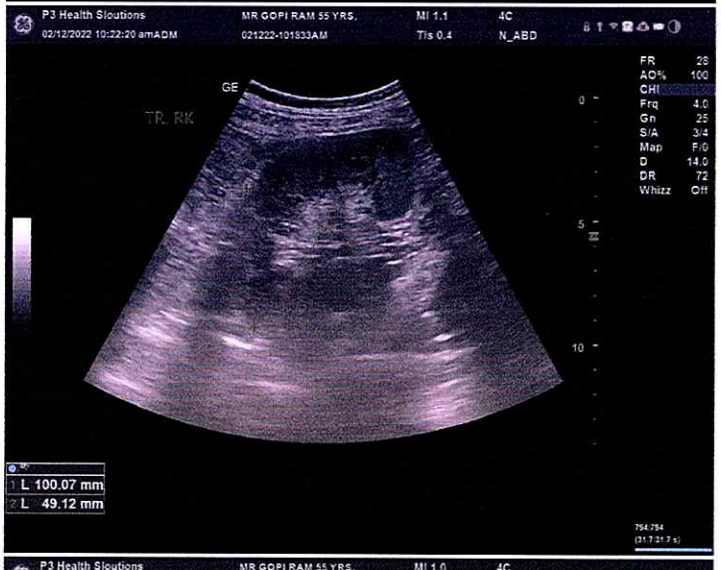
021222-101833AM
TIs 0.4
N_ABD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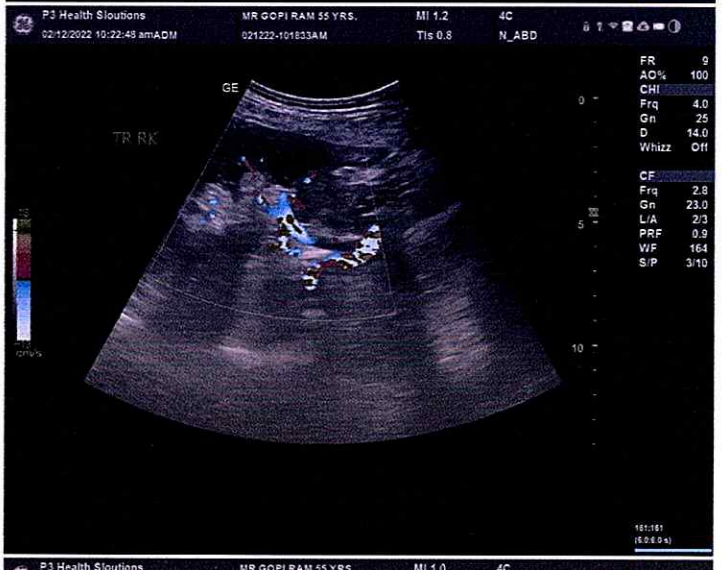
MR GOPI RAM 55 YRS,
MI 1.1
4C
02/12/2022 10:22:20 amADM
021222-101833AM
TIs 0.4
N_ABD
TR. RK
L 100.07 mm
L 49.12 mm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MI 1.2
4C
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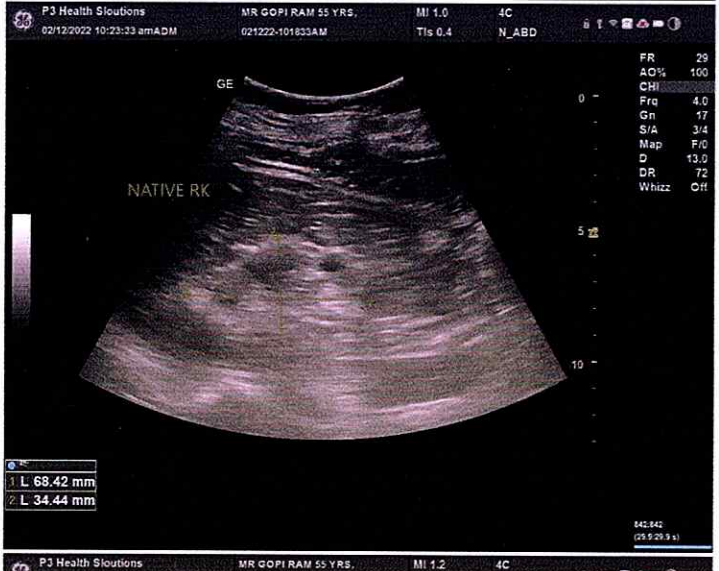
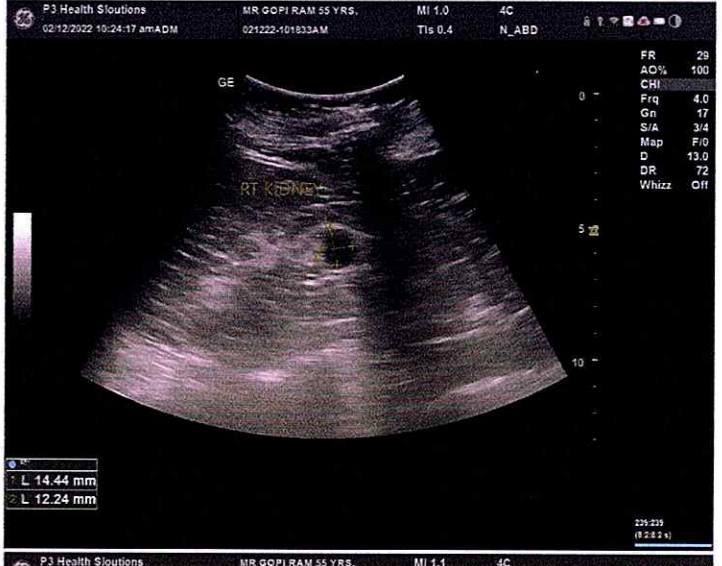
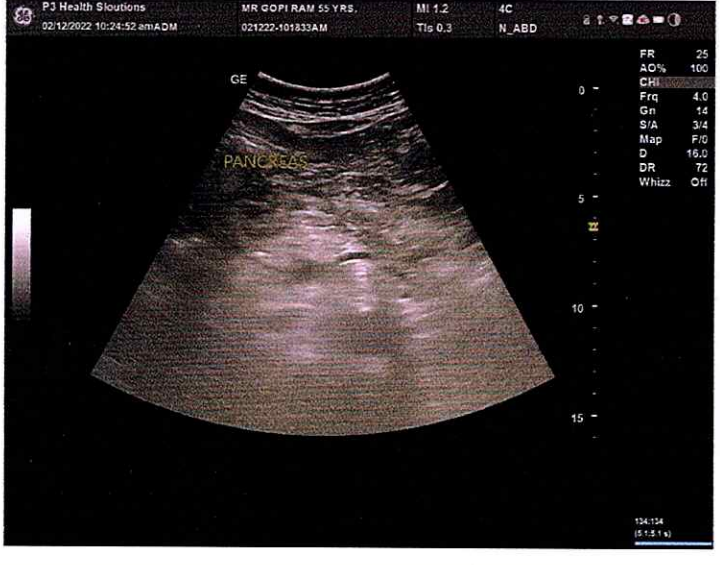
021222-101833AM
TIs 0.8
N_ABD
TR RK


P3 Health Sloutions
MR GOPI RAM 55 YRS,
MI 1.0
4C
02/12/2022 10:23:33 amADM
021222-101833AM
TIs 0.4
N_ABD
NATIVE RK
L 68.42 mm
L 34.44 mm


P3 Health Sloutions
MR GOPI RAM 55 YRS,
MI 1.0
4C
02/12/2022 10:24:17 amADM
021222-101833AM
TIs 0.4
N_ABD
RT KIDNEY
L 14.44 mm
L 12.24 mm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MR GOPI RAM 55 YRS,
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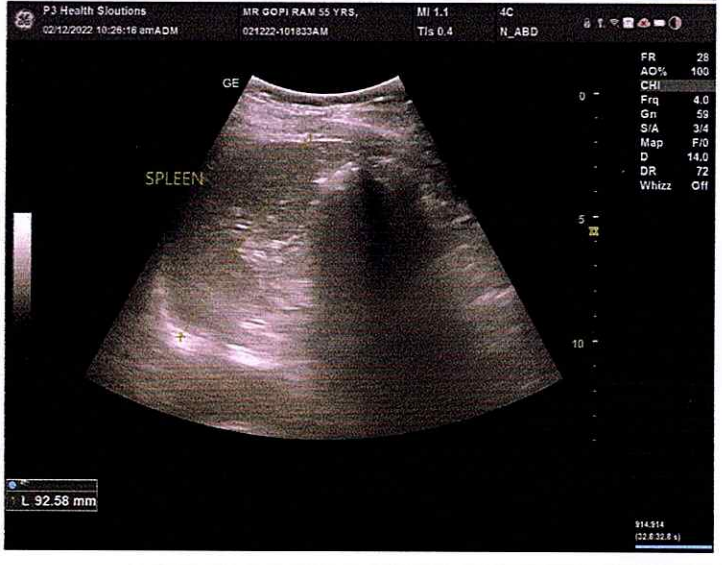
4C
02/12/2022 10:24:52 amADM
021222-101833AM
TIs 0.3
N_ABD
PANCREAS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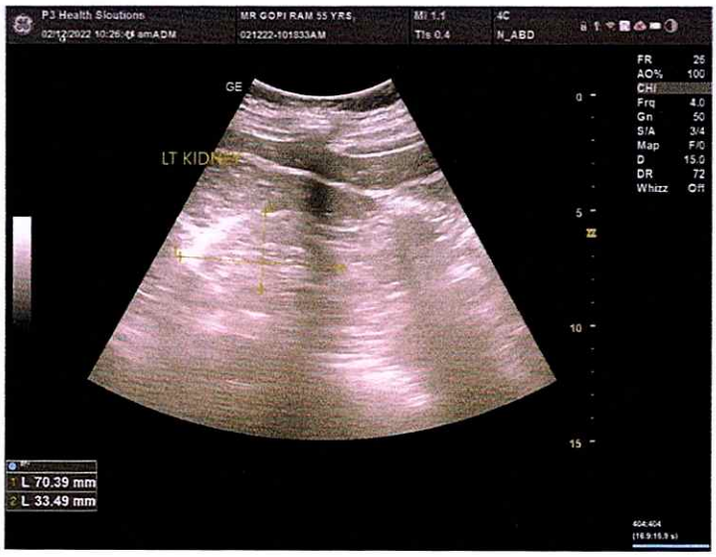
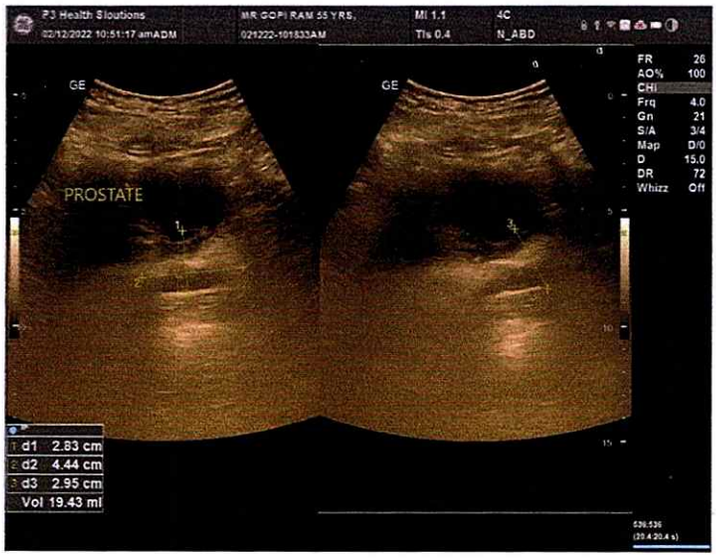
MR GOPI RAM 55 YRS,
MI 1.1
4C
02/12/2022 10:26:16 amADM
021222-101833AM
TIs 0.4
N_ABD
SPLEEN
L 92.58 mm


P3 Health Sloutions
MR GOPI RAM 55 YRS,
021222-101833AM
Mi 1.1
TIs 0.4
4C
N_ABD
GE
LT KIDNEY
FR 26
AO% 100
CHI
Frq 4.0
Gn 50
S/A 3/4
Map F/0
D 15.0
DR 72
Whizz Off
1 L 70.39 mm
2 L 33.49 mm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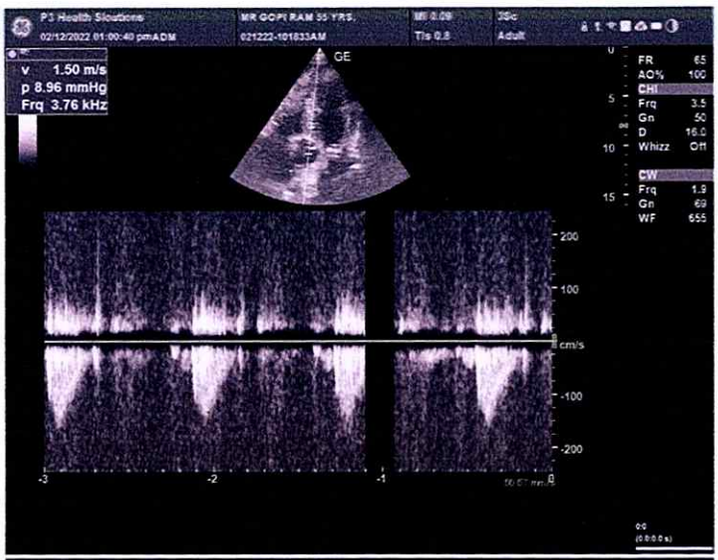
MR GOPI RAM 55 YRS,
021222-101833AM
Mi 1.1
TIs 0.4
4C
N_ABD
GE
GE
PROSTATE
FR 26
AO% 100
CHI
Frq 4.0
Gn 21
S/A 3/4
Map D/0
D 15.0
DR 72
Whizz Off
1 d1 2.83 cm
2 d2 4.44 cm
3 d3 2.95 cm
Vol 19.43 ml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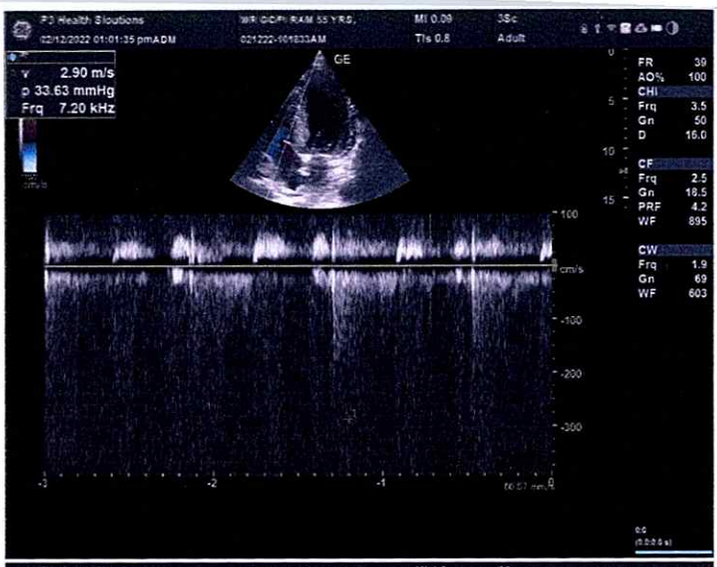
02/12/2022 01:00:40 pmADM
MR GOPI RAM 55 YRS.
021222-101833AM
MI 0.09
TIs 0.8
3Sc
Adult
v 1.50 m/s
p 8.96 mmHg
Frq 3.76 kHz
GE
FR 65
AO% 100
CHI
Frq 3.5
Gn 50
D 16.0
Whizz Off
CW
Frq 1.9
Gn 69
WF 655
200
100
cm/s
-100
-200
-3
-2
-1
0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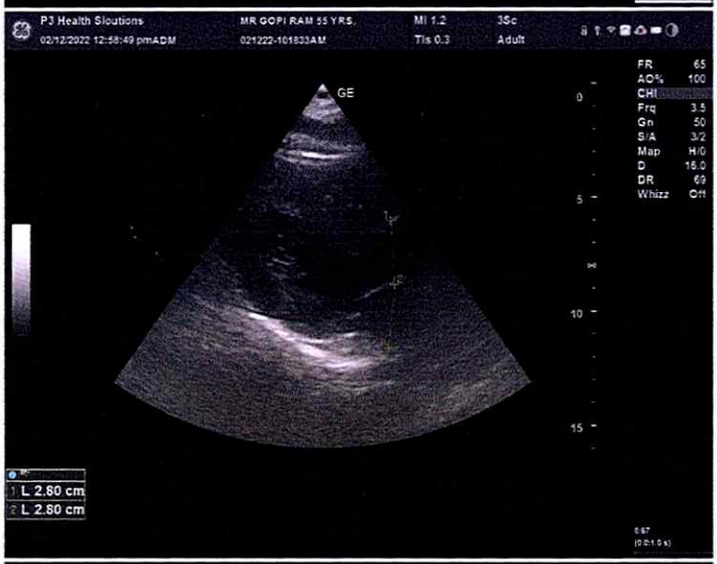
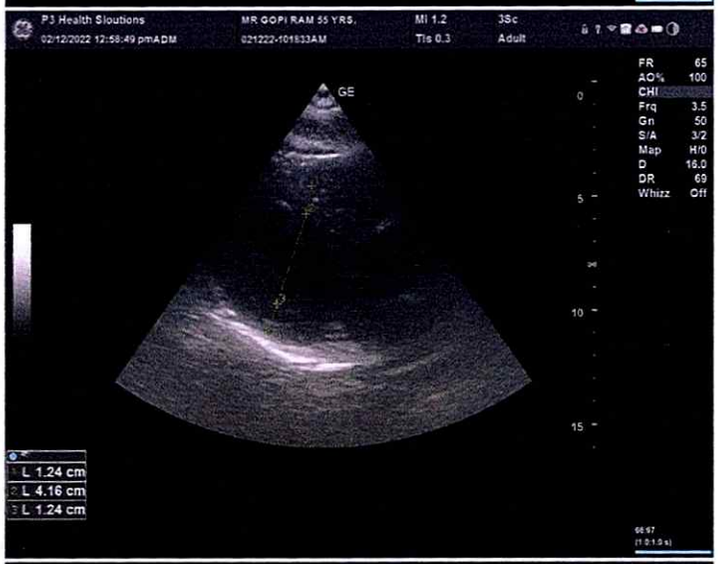
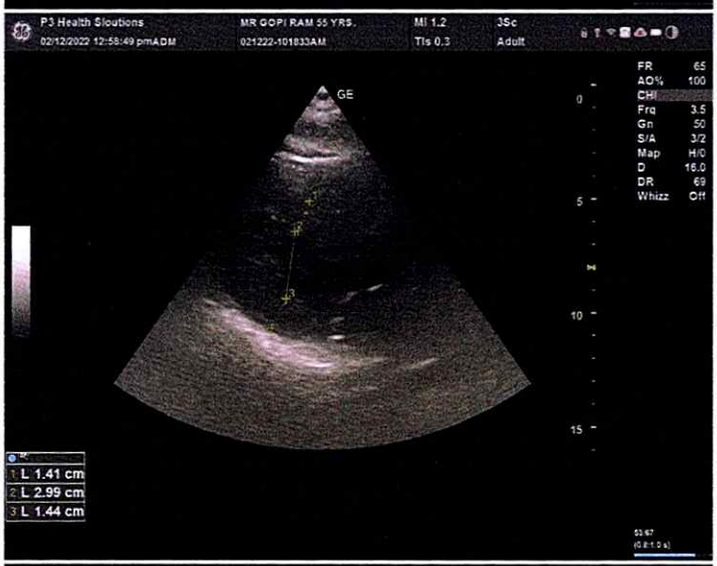
MR GOPI RAM 55 YRS.
021222-101833AM
MI 0.09
TIs 0.8
3Sc
Adult
v 2.90 m/s
p 33.63 mmHg
Frq 7.20 kHz
GE
FR 39
AO% 100
CHI
Frq 3.5
Gn 50
D 16.0
CF
Frq 2.5
Gn 18.5
PRF 4.2
WF 895
CW
Frq 1.9
Gn 69
WF 603
100
cm/s
-100
-200
-300
-3
-2
-1
0


P3 Health Sloutions
02/12/2022 12:58:49 pmADM
MR GOPI RAM 55 YRS.
021222-101833AM
MI 1.2
TIs 0.3
3Sc
Adult
GE
FR 65
AO% 100
CHI
Frq 3.5
Gn 50
S/A 3/2
Map H/0
D 16.0
DR 69
Whizz Off
1 L 2.80 cm
2 L 2.80 cm
0
5
10
15


P3 Health Sloutions
02/12/2022 12:58:49 pmADM
MR GOPI RAM 55 YRS.
021222-101833AM
MI 1.2
TIs 0.3
3Sc
Adult
GE
FR 65
AO% 100
CHI
Frq 3.5
Gn 50
S/A 3/2
Map H/0
D 16.0
DR 69
Whizz Off
1 L 1.24 cm
2 L 4.16 cm
3 L 1.24 cm
0
5
10
15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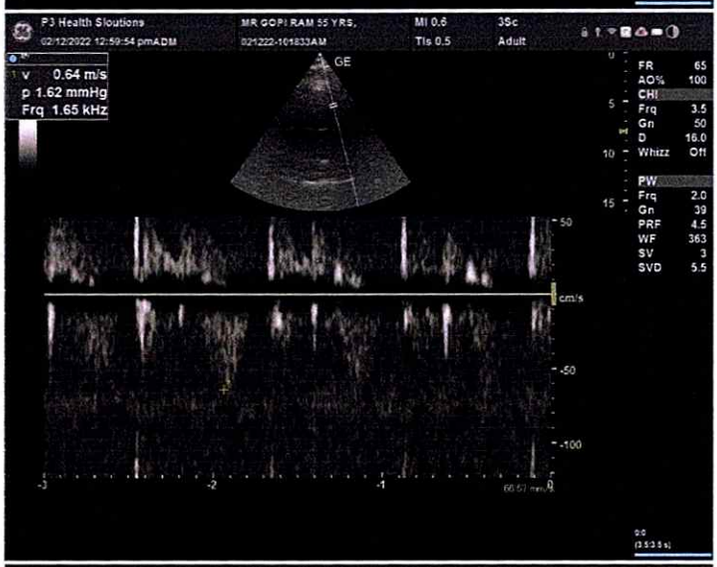
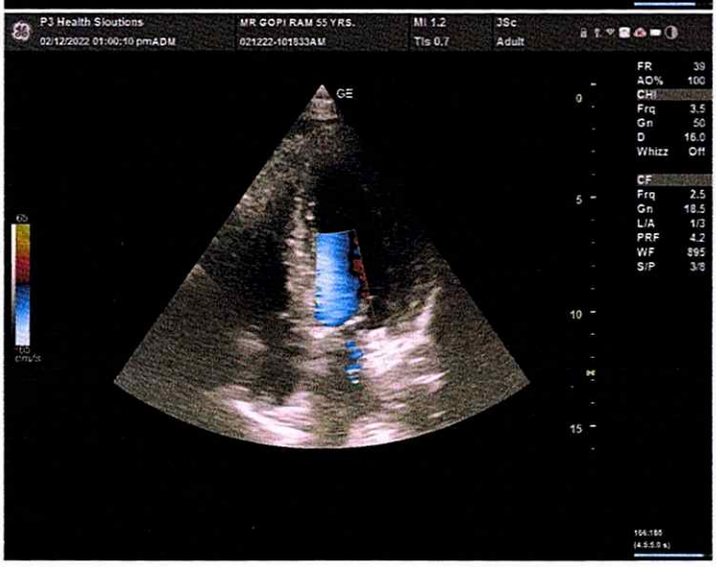
02/12/2022 12:58:49 pmADM
MR GOPI RAM 55 YRS.
021222-101833AM
MI 1.2
TIs 0.3
3Sc
Adult
GE
FR 65
AO% 100
CHI
Frq 3.5
Gn 50
S/A 3/2
Map H/0
D 16.0
DR 69
Whizz Off
1 L 1.41 cm
2 L 2.99 cm
3 L 1.44 cm
0
5
10
15


P3 Health Sloutions
02/12/2022 12:59:54 pmADM
MR GOPI RAM 55 YRS.
021222-101833AM
MI 0.6
TIs 0.5
3Sc
Adult
1 v 0.64 m/s
p 1.62 mmHg
Frq 1.65 kHz
GE
FR 65
AO% 100
CHI
Frq 3.5
Gn 50
D 16.0
Whizz Off
PW
Frq 2.0
Gn 39
PRF 4.5
WF 363
SV 3
SVD 5.5
50
cm/s
-50
-100
-3
-2
-1
0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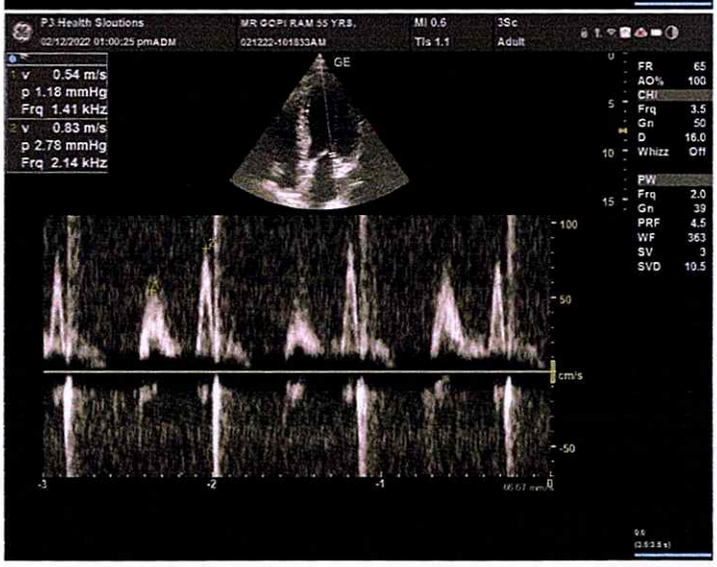
021222-101833AM
MI 1.2
TIs 0.7
3Sc
Adult
GE
FR 39
AO% 100
CHI
Frq 3.5
Gn 50
D 16.0
Whizz Off
CF
Frq 2.5
Gn 18.5
L/A 1/3
PRF 4.2
WF 895
S/P 3/8
0
5
10
15


P3 Health Sloutions
02/12/2022 01:00:25 pmADM
MR GOPI RAM 55 YRS.
021222-101833AM
MI 0.6
TIs 1.1
3Sc
Adult
1 v 0.54 m/s
p 1.18 mmHg
Frq 1.41 kHz
2 v 0.83 m/s
p 2.78 mmHg
Frq 2.14 kHz
GE
FR 65
AO% 100
CHI
Frq 3.5
Gn 50
D 16.0
Whizz Off
PW
Frq 2.0
Gn 39
PRF 4.5
WF 363
SV 3
SVD 19.5
100
50
cm/s
-50
-3
-2
-1
0

P3 HEALTH SOLUTIONS LLP
(ASSOCIATES OF MAXCARE DIAGNOSTICS)

B-14, Vidhyadhar Enclave - II, Near Axis Bank
Central Spine, Vidhyadhar Nagar, Jaipur - 302023
+91 141 4824885 maxcarediagnostics1@gmail.com

NAME :- Mr. GOPI RAM
Age :- 55 Yrs 5 Mon 29 Days
Sex :- Male

Patient ID :-12222581 Date :- 02/12/2022 09:19:00
Ref. By Doctor:-BANK OF BARODA
Lab/Hosp :-
Company :- Mr.MEDI ASSIST TPA

Final Authentication : 02/12/2022 14:21:47

HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
FULL BODY HEALTH CHECKUP ABOVE 40 MALE			
HAEMOGARAM			
HAEMOGLOBIN (Hb)	15.2	g/dL	13.0 - 17.0
TOTAL LEUCOCYTE COUNT	8.40	/cumm	4.00 - 10.00
DIFFERENTIAL LEUCOCYTE COUNT			
NEUTROPHIL	**84.0** H	%	40.0 - 80.0
LYMPHOCYTE	**11.0** L	%	20.0 - 40.0
EOSINOPHIL	2.0	%	1.0 - 6.0
MONOCYTE	3.0	%	2.0 - 10.0
BASOPHIL	0.0	%	0.0 - 2.0
TOTAL RED BLOOD CELL COUNT (RBC)	4.72	x10^6/uL	4.50 - 5.50
HEMATOCRIT (HCT)	46.40	%	40.00 - 50.00
MEAN CORP VOLUME (MCV)	98.0	fL	83.0 - 101.0
MEAN CORP HB (MCH)	**32.2** H	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	32.7	g/dL	31.5 - 34.5
PLATELET COUNT	**140** L	x10^3/uL	150 - 410
RDW-CV	**14.2** H	%	11.6 - 14.0
MENTZER INDEX	**20.76** H		0.00 - 0.00

A complete blood picture (CBP) is a kind of blood test that is done to assess a person's overall health and diagnose a wide range of health disorders like leukemia, anemia and other infections.

A complete blood count (CBC) is a complete blood test that diagnose many components and features of a persons blood which includes: -

*__Red Blood Cells (RBC),__ which carry oxygen -

*__White Blood Cells (WBC)__, which help in fighting against infections -

*__Hemoglobin__, which is the oxygen carrying protein in the red blood cells -

*__Hematocrit (HCT)__, the proportion of RBC to the fluid component, or plasma present in blood -

*__Platelets,__ which aid in blood clotting

(CBC): Methodology: TLC,TRBC,PCV,PLT Impedance method, HB Calorimetric method,. and **MCH,MCV,MCHC,MENTZER INDEX** are calculated. **InstrumentName: MINDRAY BC-3000 Plus 3** part automatic analyzer,

ADIYTA

Technologist

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

P3 HEALTH SOLUTIONS LLP
(ASSOCIATES OF MAXCARE DIAGNOSTICS)

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NAME :- Mr. GOPI RAM	Patient ID :-12222581	Date :- 02/12/2022	09:19:00
Age :- 55 Yrs 5 Mon 29 Days	Ref. By Doctor:-BANK OF BARODA		
Sex :- Male	Lab/Hosp :-		
	Company :- Mr.MEDI ASSIST TPA		

Final Authentication : 02/12/2022 14:21:47

HAEMATOLOGY

Erythrocyte Sedimentation Rate (ESR)	10	mm in 1st hr	00 - 15
Methord:- Westergreen			

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:- GOD POD	104.0	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 .

BLOOD SUGAR PP (Plasma) Methord:- GOD PAP	**179.0** H	mg/dl	70.0 - 140.0

Instrument Name: HORIBA 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	6.2	mg%	Non-Diabetic < 6.0 Good Control 6.0-7.0 Weak Control 7.0-8.0 Poor control > 8.0
MEAN PLASMA GLUCOSE	**131** H	mg/dL	68 - 125
Methord:- Calculated Parameter			

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

Note:
1.Shortened RBC life span –HbA1c test will not be accurate when a person has a condition that affects the average lifespan of red blood cells (RBCs), such as hemolytic anemia or blood loss. When the lifespan of RBCs in circulation is shortened, the A1c result is falsely low and is an unreliable measurement of a person's average glucose over time.
2.Abnormal forms of hemoglobin – The presence of some hemoglobin variants, such as hemoglobin S in sickle cell anemia, may affect certain methods for measuring A1c. In these cases, fructosamine can be used to monitor glucose control.

Advised:
1.To follow patient for glycemic control test like fructosamine or glycated albumin may be performed instead.
2.Hemoglobin HPLC screen to analyze abnormal hemoglobin variant.
estimated Average Glucose (eAG) : based on value calculated according to National Glycohemoglobin Standardization Program (NGSP) criteria.

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HAEMATOLOGY

BLOOD GROUP ABO Methord:- Haemagglutination reaction	"B" POSITIVE

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BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIPID PROFILE			
TOTAL CHOLESTEROL Methord:- CHOD-PAP methodology	211.00	mg/dl	Desirable <200 Borderline 200-239 High> 240
InstrumentName:MISPA PLUS **Interpretation:** Cholesterol measurements are used in the diagnosis and treatments of lipid lipoprotein metabolism disorders.			
TRIGLYCERIDES Methord:- GPO-TOPS methodology	57.20	mg/dl	Normal <150 Borderline high 150-199 High 200-499 Very high >500
InstrumentName:MISPA PLUS **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:- Selective inhibition Method	83.10	mg/dl	Male 35-80 Female 42-88
Instrument Name:MISPA 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	118.37	mg/dl	Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190
VLDL CHOLESTEROL Methord:- Calculated	11.44	mg/dl	0.00 - 80.00
T.CHOLESTEROL/HDL CHOLESTEROL RATIO Methord:- Calculated	2.54		0.00 - 4.90
LDL / HDL CHOLESTEROL RATIO Methord:- Calculated	1.42		0.00 - 3.50
TOTAL LIPID Methord:- CALCULATED	553.61	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.

Comments: 1- ATP III suggested the addition of Non HDL Cholesterol (Total Cholesterol – HDL Cholesterol) as an indicator of all

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BIOCHEMISTRY

atherogenic lipoproteins (mainly LDL & VLDL). The Non HDL Cholesterolis used as a secondary target of therapy in persons with triglycerides >=200 mg/dL. The goal for Non HDL Cholesterol in those with increased triglyceride is 30 mg/dL above that set for LDL Cholesterol.

2 -For calculation of CHD risk,history of smoking, any medication for hypertension & current B.P. levels are required.

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BIOCHEMISTRY

LIVER PROFILE WITH GGT			
SERUM BILIRUBIN (TOTAL) Methord:- DMSO/Diazo	0.51	mg/dL	Infants : 0.2-8.0 mg/dL Adult - Up to - 1.2 mg/dL
SERUM BILIRUBIN (DIRECT) Methord:- DMSO/Diazo	0.21	mg/dL	Up to 0.40 mg/dL
SERUM BILIRUBIN (INDIRECT) Methord:- Calculated	0.30	mg/dl	0.30-0.70
SGOT Methord:- IFCC	37.0	U/L	Men- Up to - 37.0 Female - Up to - 31.0
SGPT Methord:- IFCC	36.6	U/L	Men- Up to - 40.0 Female- Up to - 31.0
SERUM ALKALINE PHOSPHATASE Methord:- DGKC - SCE	85.20	U/L	53.00 - 141.00
SERUM GAMMA GT Methord:- Szasz methodology Instrument Name Randox Rx Imola	18.90	U/L	10.00 - 45.00
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			
SERUM TOTAL PROTEIN Methord:- Direct Biuret Reagent	6.10	g/dl	5.10 - 8.00
SERUM ALBUMIN Methord:- Bromocresol Green	4.10	g/dl	2.80 - 4.50
SERUM GLOBULIN Methord:- CALCULATION	**2.00** L	gm/dl	2.20 - 3.50
A/G RATIO	2.05		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., 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

RFT / KFT WITH ELECTROLYTES

SERUM UREA Methord:- Urease/GLDH	29.20	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 Methord:- Jaffe's Method	1.30	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	3.90	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 Methord:- ISE	**133.0** L	mmol/L	135.0 - 150.0

Interpretation: Decreased sodium - Hyponatraemia Causes include: fluid or electrolyte loss,Drugs,Oedematous states,Legionnaire's disease and other chest infections,pseudonatremia, Hyperlipidaemias and paraproteinaemias,endocrine diseases ,SIADH.

POTASSIUM Methord:- ISE	4.66	mmol/L	3.50 - 5.50

Interpretation: A. Elevated potassium (hyperkalaemia)• Artefactual,Physiological elevation,Drugs, Pathological states,Renal failure Adrenocortical insufficiency, metabolic acidoses, very high platelet or white cell counts B. Decreased potassium (hypokalaemia)Drugs, Liquoric,Diarrhoea and vomiting,Metabolic alkalosis,Corticosteroid excess, Oedematous state,Anorexia nervosa/bulimia

CHLORIDE Methord:- ISE	106.0	mmol/L	94.0 - 110.0

Interpretation: Used for Electrolyte monitoring.

SERUM CALCIUM Methord:- Colorimetric method	9.74	mg/dl	8.10 - 11.50

InstrumentName:Rx Daytona 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 Methord:- Direct Biuret Reagent	6.10	g/dl	5.10 - 8.00
SERUM ALBUMIN Methord:- Bromocresol Green	4.10	g/dl	

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BIOCHEMISTRY

SERUM GLOBULIN Methord:- CALCULATION	**2.00** L	gm/dl	2.20 - 3.50
A/G RATIO	2.05		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 bloodincreases. 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.

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CLINICAL PATHOLOGY

URINE SUGAR (FASTING) **Collected Sample Received**	Nil	Nil

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CLINICAL PATHOLOGY

STOOL ANALYSIS

PHYSICAL EXAMINATION		
COLOUR	YELLOW BROWN	
CONSISTENCY	SEMI SOLID	
MUCUS	ABSENT	
BLOOD	ABSENT	
MICROSCOPIC EXAMINATION		
RBC's	NIL	/HPF
WBC/HPF	NIL	/HPF
MACROPHAGES	ABSENT	
OVA	ABSENT	
CYSTS	ABSENT	
TROPHOZOITES	ABSENT	
CHARCOT LEYDEN CRYSTALS	ABSENT	
OTHERS	ABSENT	

Collected Sample Received

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IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
PSA (PROSTATE SPECIFIC ANTIGEN) -TOTAL Methord:- Methodology: CLIA	0.468	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.

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IMMUNOASSAY

TOTAL THYROID PROFILE			
THYROID-TRIIODOTHYRONINE T3	0.91	ng/mL	0.70 - 2.04

Methord:- ECLIA

NOTE-TSH levels are subject to circardian variation,reaching peak levels between 2-4 AM and min between 6-10 PM. The variation is the order of 50% hence time of the day has influence on the measures serum TSH concentration.Dose and time of drug intake also influence the test result. Transient increase in TSH levels or abnormal TSH levels can be seen in some non thyroidal conditions,simoultaneous measurement of TSH with free T4 is useful in evaluating differantial diagnosis

INTERPRETATION-Ultra Sensitive 4th generation assay 1.Primary hyperthyroidism is accompanied by ↑serum T3 & T4 values along with " TSH level.2.Low TSH,high FT4 and TSH receptor antibody(TRAb) +ve seen in patients with Graves disease 3.Low TSH,high FT4 and TSH receptor antibody(TRAb) -ve seen in patients with Toxic adenoma/Toxic Multinodular goiter 4.HighTSH,Low FT4 and Thyroid microsomal antibody increased seen in patients with Hashimotos thyroiditis 5.HighTSH,Low FT4 and Thyroid microsomal antibody normal seen in patients with Iodine deficiency/Congenital T4 synthesis deficiency 6.Low TSH,Low FT4 and TRH stimulation test -Delayed response seen in patients with Tertiary hypothyroidism
7.Primary hypothyroidism is accompanied by ↓ serum T3 and T4 values & 'serum TSH levels8.Normal T4 levels accompanied by ' T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis9.Normal or" T3 & 'T
10.Normal T3 & T4 along with " TSH indicate mild / Subclinical Hyperthyroidism .11.Normal T3 & " T4 along with ' TSH is seen in Hypothyroidism .12.Normal T3 & T4 levels with ' TSH indicate Mild / Subclinical Hypoth

DURING PREGNANCY - REFERENCE RANGE for TSH IN uIU/mL (As per American Thyroid Association) 1st Trimester : 0.10-2.50 uIU/mL 2nd Trimester : 0.20-3.00 uIU/mL 3rd Trimester : 0.30-3.00 uIU/mL The production, circulation, and disintegration of thyroid hormones are altered throughout the stages of pregnancy.

REMARK-Assay results should be interpreted in context to the clinical condition and associated results of other investigations. Previous treatment with corticosteroid therapy may result in lower TSH levels while thyroid hormone levels are normal. Results are invalidated if the client has undergone a radionuclide scan within 7-14 days before the test. Abnormal thyroid test findings often found in critically ill patients should be repeated after the critical nature of the condition is resolved.TSH is an important marker for the diagnosis of thyroid dysfunction.Recent studies have shown that the TSH distribution progressively shifts to a higher concentration with age ,and it is debatable whether this is due to a real change with age or an increasing proportion of unrecognized thyroid disease in the elderly. ***

THYROID - THYROXINE (T4)	6.91	µIU/ml	5.10 - 14.10

Methord:- ECLIA

NOTE-TSH levels are subject to circardian variation,reaching peak levels between 2-4 AM and min between 6-10 PM. The variation is the order of 50% hence time of the day has influence on the measures serum TSH concentration.Dose and time of drug intake also influence the test result. Transient increase in TSH levels or abnormal TSH levels can be seen in some non thyroidal conditions,simoultaneous measurement of TSH with free T4 is useful in evaluating differantial diagnosis

INTERPRETATION-Ultra Sensitive 4th generation assay 1.Primary hyperthyroidism is accompanied by ↑serum T3 & T4 values along with " TSH level.2.Low TSH,high FT4 and TSH receptor antibody(TRAb) +ve seen in patients with Graves disease 3.Low TSH,high FT4 and TSH receptor antibody(TRAb) -ve seen in patients with Toxic adenoma/Toxic Multinodular goiter 4.HighTSH,Low FT4 and Thyroid microsomal antibody increased seen in patients with Hashimotos thyroiditis 5.HighTSH,Low FT4 and Thyroid microsomal antibody normal seen in patients with Iodine deficiency/Congenital T4 synthesis deficiency 6.Low TSH,Low FT4 and TRH stimulation test -Delayed response seen in patients with Tertiary hypothyroidism
7.Primary hypothyroidism is accompanied by ↓ serum T3 and T4 values & 'serum TSH levels8.Normal T4 levels accompanied by ' T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis9.Normal or" T3 & 'T
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TSH	1.728	µIU/mL	0.350 - 5.500

Methord:- ECLIA

NOTE-TSH levels are subject to circardian variation,reaching peak levels between 2-4 AM and min between 6-10 PM. The variation is the order of 50% hence time of the day has influence on the measures serum TSH concentration.Dose and time of drug intake also influence the test result.
Transient increase in TSH levels or abnormal TSH levels can be seen in some non thyroidal conditions,simoultaneous measurement of TSH with free T4 is useful in evaluating differantial diagnosis

INTERPRETATION-Ultra Sensitive 4th generation assay
1.Primary hyperthyroidism is accompanied by ↑serum T3 & T4 values along with ↓ TSH level.
2.Low TSH,high FT4 and TSH receptor antibody(TRAb) +ve seen in patients with Graves disease

ADITYA

Technologist

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

P3 HEALTH SOLUTIONS LLP
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B-14, Vidhyadhar Enclave - II, Near Axis Bank
Central Spine, Vidhyadhar Nagar, Jaipur - 302023
+91 141 4824885 maxcarediagnostics1@gmail.com

NAME :- Mr. GOPI RAM
Age :- 55 Yrs 5 Mon 29 Days
Sex :- Male

Patient ID :-12222581 Date :- 02/12/2022 09:19:00
Ref. By Doctor:-BANK OF BARODA
Lab/Hosp :-
Company :- Mr.MEDI ASSIST TPA

Final Authentication : 02/12/2022 14:21:47

IMMUNOASSAY

3.Low TSH,high FT4 and TSH receptor antibody(TRAb) -ve seen in patients with Toxic adenoma/Toxic Multinodular goiter
4.HighTSH,Low FT4 and Thyroid microsomal antibody increased seen in patients with Hashimotos thyroiditis
5.HighTSH,Low FT4 and Thyroid microsomal antibody normal seen in patients with Iodine deficiency/Congenital T4 synthesis deficiency
6.Low TSH,Low FT4 and TRH stimulation test -Delayed response seen in patients with Tertiary hypothyroidism
7.Primary hypothyroidism is accompanied by ↓ serum T3 and T4 values & ↑serum TSH levels
8.Normal T4 levels accompanied by ↑ T3 levels and low TSH are seen in patients with T3 Thyrotoxicosis
9.Normal or↓ T3 & ↑T4 levels indicate T4 Thyrotoxicosis (problem is conversion of T4 to T3)
10.Normal T3 & T4 along with ↓ TSH indicate mild / Subclinical Hyperthyroidism .
11.Normal T3 & ↓ T4 along with ↑ TSH is seen in Hypothyroidism .
12.Normal T3 & T4 levels with ↑ TSH indicate Mild / Subclinical Hypothyroidism .
13.Slightly ↑ T3 levels may be found in pregnancy and in estrogen therapy while ↓ levels may be encountered in severe illness , malnutrition , renal failure and during therapy with drugs like propanolol.
14.Although ↑ TSH levels are nearly always indicative of Primary Hypothroidism ,rarely they can result from TSH secreting pituitary tumours.

DURING PREGNANCY - REFERENCE RANGE for TSH IN uIU/mL (As per American Thyroid Association)
1st Trimester : 0.10-2.50 uIU/mL
2nd Trimester : 0.20-3.00 uIU/mL
3rd Trimester : 0.30-3.00 uIU/mL
The production, circulation, and disintegration of thyroid hormones are altered throughout the stages of pregnancy.

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*** End of Report ***

ADIYTA

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Final Authentication : 02/12/2022 14:21:47

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)	6.0		5.0 - 7.5
SPECIFIC GRAVITY	1.010		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	2-3	/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		

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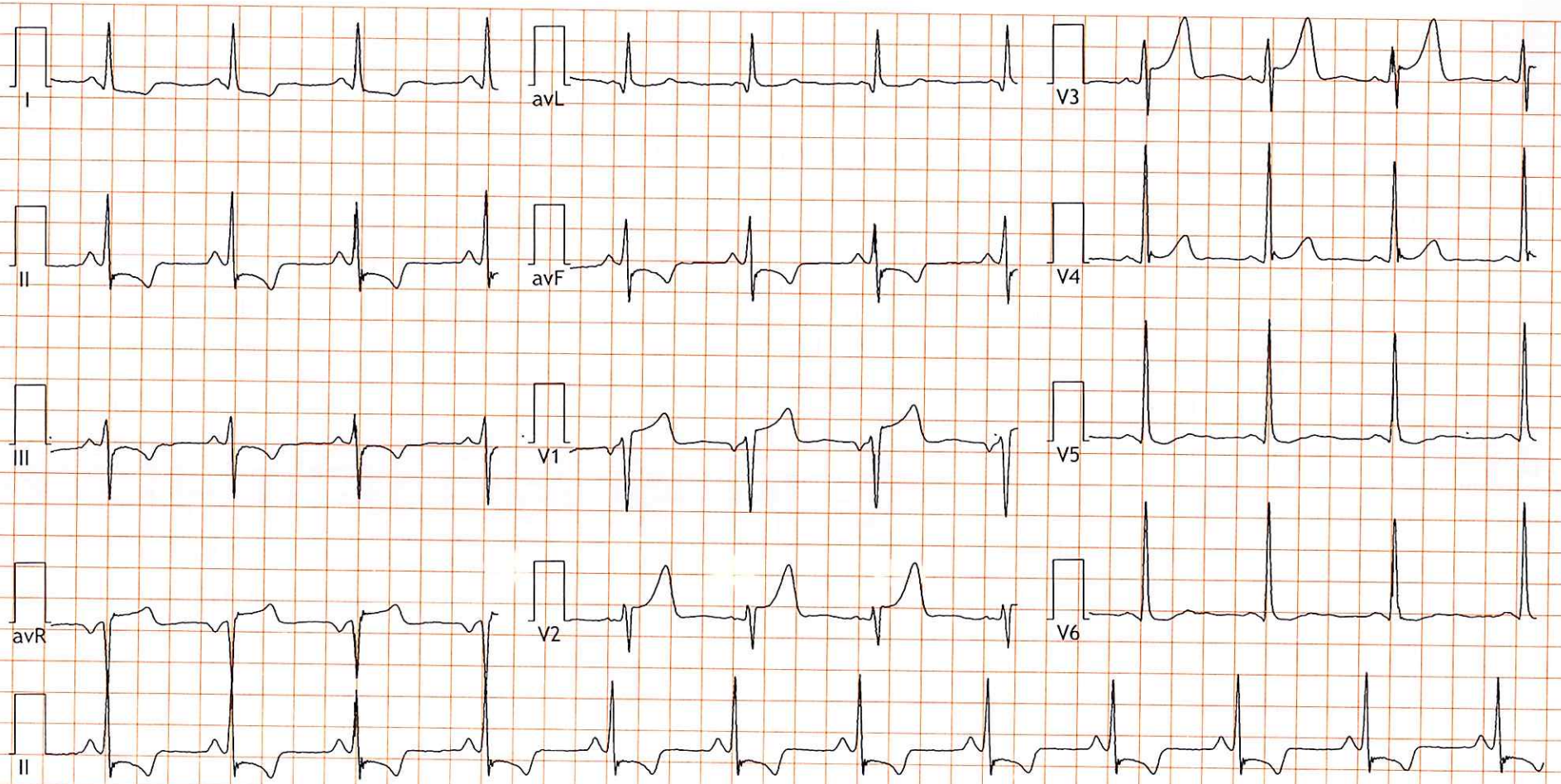
P3 HEALTH SOLUTIONS LLP
B-14, Vidhyanagar Nagar, Enclave, Phase-2, Jaipur
12229451322602/Mr gopi Ram 55Yrs-2Months/Male Kgs/ Cms BP: ___/___ mmHg **HR: 70 bpm**
Ref.: BANK OF BARODA Test Date: 02-Dec-2022(12:22:47) Notch: 50Hz 0.05Hz - 100Hz 10mm/mV 25mm/Sec

PR Interval: 118 ms
QRS Duration: 90 ms
QT/QTc: 373/405ms
P-QRS-T Axis: 55 - 6 - -124 (Deg)



FINDINGS: Abnormal ECG with Indication of Old MI (Posterior)
Vent Rate : 70 bpm; PR Interval : 118 ms; QRS Duration: 90 ms; QT/QTc Int : 373/405 ms
P-QRS-T axis: 55• 6• -124• (Deg)
Comments :

Sinus rhythm with T inversion in inferior lead with poor R progression in lead V1 V3

Dr. Naresh Kumar Mohanka
RMC No.: 35703
MBBS, DIP. CARDIO (ESCO...
D.E.M. (RCG...