

# Dr. Goyal's

## Path Lab & Imaging Centre

B-51, Ganesh Nagar, Opp. Janpath Corner, New Sanganer Road, Jaipur-302019  
Tele: 0141-2293346, 4049787, 9887049787  
Website: www.drgoyalspathlab.com | E-mail: drgoyalpiyush@gmail.com



### General Physical Examination

Date of Examination: 15/04/23

Name: MANOJ SURBHA Age: 53 Sex: M

DOB: 23/11/1969

Referred By: BOB (Medicel)

Photo ID: Adhar ID #: Adhar

Ht: 171 (cm)

Wt: 70 (Kg)

Chest (Expiration): 92 (cm)

Abdomen Circumference: 88 (cm)

Blood Pressure: 150/92 mm Hg PR: 76 / min RR: 16 / min Temp: Afebrile

BMI 30.1

Eye Examination: Dis vision 6/6, Near vision N/6 (Both eyes).  
No colour blindness.

Other: Not Significant

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

Signature Of Examinee : [Signature] Name of Examinee: \_\_\_\_\_

Signature Medical Examiner : [Signature] Name Medical Examiner \_\_\_\_\_

Dr Piyush Goyal  
M.B.B.S., D.M.R.D.  
RMC Reg No - 017906



भारत सरकार



मनोज गुप्ता  
Manoj Gupta  
जन्म तिथि / DOB: 23/11/1969  
पुरुष / MALE  
Mobile No.: 7740855705



9266 3784 5464

मेरा आधार, मेरी पहचान

*(Handwritten signature)*



भारतीय विधिबन्ध पहचान प्राधिकरण

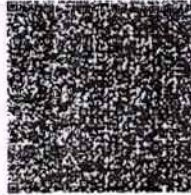
भारत सरकार

Download Date: 21/07/2018

पत्ता:  
S/O ध्रुव प्रसाद गुप्ता, ६६/८९, हीरापथ, जयपुर, जयपुर,  
राजस्थान - 302020

Address:  
S/O Dhruv Prasad Gupta, 66/81,  
heerapath, Jaipur, Jaipur, Rajasthan -  
302020

QR Code with Photograph



GENERATION DATE: 20/09/2018



1947  
1800 300 1947

help@uidai.gov.in www.uidai.gov.in

P.O. Box No. 1947,  
Bengaluru-560 001

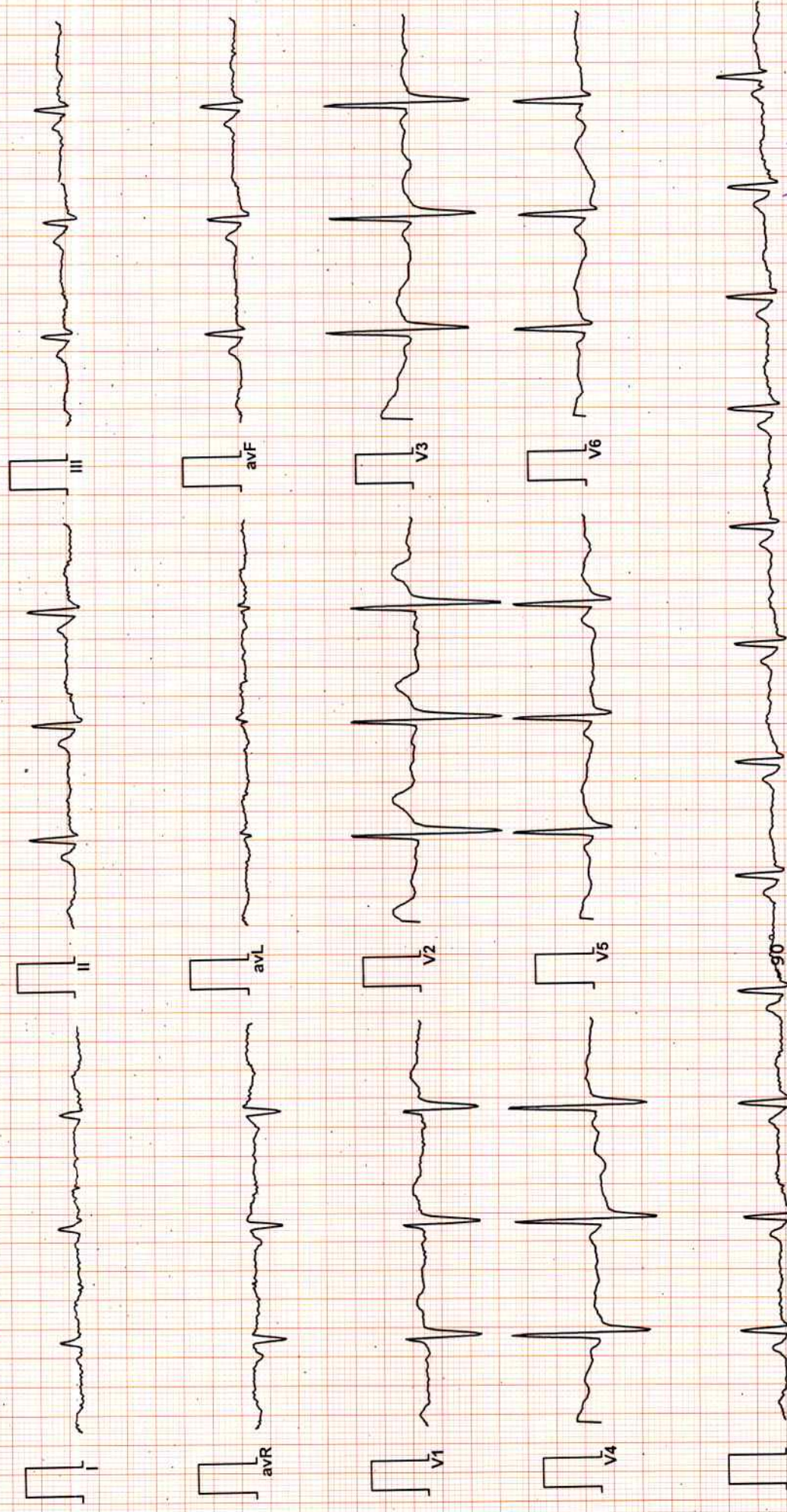
Dr Piyush Goyal  
M.B.S., D.M.R.D  
RMC Reg No -017988

ECG

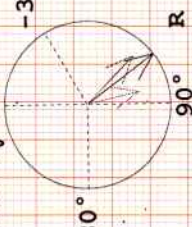
DR. GOYAL PATH LAB & IMAGING CENTER, JAIPUR

4439 / MR MANOJ GUPTA / 53 Yrs / M / Non Smoker

Heart Rate : 76 bpm / Tested On : 15-Apr-23 10:55:47 / HF 0.05 Hz - LF 35 Hz / Notch 50 Hz / Sn 1.00 Cm/mV / Sw 25 mm/s / Refd By: BOB



*sinus rhythm with e inversion in infarto lead*



Vent Rate : 76 bpm  
PR Interval : 138 ms  
QRS Duration: 108 ms  
QT/QTc Int : 390/418 ms  
P-QRS-T axis: 77.00 • 53.00 • 44.00 •

Axis  
R 53.00° T 44.00° P 77.00°

Reported By: *[Signature]*

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Date :- 15/04/2023 09:25:18	Patient ID :- 1223231
<b>NAME :- Mr. MANOJ GUPTA</b>	Ref. By Doctor:-BOB
Sex / Age :- Male 53 Yrs	Lab/Hosp :-
Company :- MediWheel	

Final Authentication : 15/04/2023 14:27:24

BOB PACKAGE ABOVE 40MALE  
 2D ECHO OPTION TMT (ADULT/CHILD)

### 2D-ECHOCARDIOGRAPHY M.MODE WITH DOPPLER STUDY:

#### FAIR TRANSTHORACIC ECHOCARDIOGRAPHIC WINDOW MORPHOLOGY:

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

#### M.MODE EXAMINATION:

AO	28	mm	LA	34	Mm	IVS-D	9	mm
IVS-S	16	mm	LVID	44	Mm	LVSD	29	mm
LVPW-D	11	mm	LVPW-S	18	Mm	RV		mm
RVWT		mm	EDV		MI	LVVS		ml
LVEF	62%		RWMA		ABSENT			

#### CHAMBERS:

LA	NORMAL	RA	NORMAL
LV	NORMAL	RV	NORMAL
PERICARDIUM		NORMAL	

#### COLOUR DOPPLER:

MITRAL VALVE					
E VELOCITY	0.63	m/sec	PEAK GRADIENT		Mm/hg
A VELOCITY	0.92	m/sec	MEAN GRADIENT		Mm/hg
MVA BY PHT		Cm2	MVA BY PLANIMETRY		Cm2
MITRAL REGURGITATION				ABSENT	
AORTIC VALVE					
PEAK VELOCITY	1.3	m/sec	PEAK GRADIENT		mm/hg
AR VMAX		m/sec	MEAN GRADIENT		mm/hg
AORTIC REGURGITATION				ABSENT	
TRICUSPID VALVE					
PEAK VELOCITY	0.38	m/sec	PEAK GRADIENT		mm/hg
MEAN VELOCITY		m/sec	MEAN GRADIENT		mm/hg
VMax VELOCITY					
TRICUSPID REGURGITATION				ABSENT	
PULMONARY VALVE					
PEAK VELOCITY	0.90	M/sec.	PEAK GRADIENT		Mm/hg
MEAN VELOCITY			MEAN GRADIENT		Mm/hg
PULMONARY REGURGITATION				ABSENT	

TABBSUM

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### Impression--

1. LV Diastolic Dysfunction Grade I.
2. Normal LV size & contractility
3. No RWMA, LVEF 62%.
4. Normal cardiac chamber.
5. Normal valve
6. No clot, no vegetation, no pericardial effusion.

  
(Cardiologist)

\*\*\* End of Report \*\*\*

TABBSUM

Page No: 2 of 2

Dr. Piyush Goyal  
M.B.B.S., D.M.R.D.  
RMC Reg No. 017996

Dr. Poonam Gupta  
MBBS, MD (Radio Diagnosis)  
RMC No. 32495

Dr. Ashish Choudhary  
MBBS, MD (Radio Diagnosis)  
Fetal Medicine Consultant  
FMF ID - 260517 | RMC No 22430

Dr. Abhishek Jain  
MBBS, DNB, (Radio-Diagnosis)  
RMC No. 21687

Transcript by.



Date :- 15/04/2023 09:25:18	Patient ID :- 1223231
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Sex / Age :- Male 53 Yrs	Lab/Hosp :-
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Final Authentication : 15/04/2023 12:56:23

BOB PACKAGE ABOVE 40MALE

**USG WHOLE ABDOMEN**

**Liver** is of normal size. 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 of normal size. 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. No focal lesion is seen. Collecting system does not show any dilatation or calculus.

**Urinary bladder** is well distended and showing smooth wall with normal thickness. Urinary bladder does not show any calculus or mass lesion.

**Prostate** is normal in size with normal echo-texture and outline.  
No enlarged nodes are visualised.No retro-peritoneal lesion is identified  
No significant free fluid is seen in peritoneal cavity.

**IMPRESSION:**

**\* Normal study**

*Needs clinical correlation for further evaluation*

\*\*\* End of Report \*\*\*



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Final Authentication : 15/04/2023 10:45:45

BOB PACKAGE ABOVE 40MALE

### X RAY CHEST PA VIEW:

Old malunited fracture of right 3rd,4th, 5th & 6th ribs seen.

Both lung fields appears clear.

Bronchovascular markings appear normal.

Trachea is in midline.

Both the hilar shadows are normal.

Both the C.P.angles is clear.

Both the domes of diaphragm are normally placed.

Heart shadows appear normal.

Unfolding of arch of aorta is seen.

(Please correlate clinically and with relevant further investigations)

\*\*\* End of Report \*\*\*

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Patient ID :-1223231



**NAME :- Mr. MANOJ GUPTA**

Ref. By Dr:- BOB

Sex / Age :- Male 53 Yrs

Lab/Hosp :-

Company :- MediWheel

Sample Type :- EDTA

Sample Collected Time 15/04/2023 09:38:53

Final Authentication : 15/04/2023 11:45:21

### HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
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BOB PACKAGE ABOVE 40MALE

GLYCOSYLATED HEMOGLOBIN (HbA1C)

6.2 H %

Non-diabetic: < 5.7  
Pre-diabetics: 5.7-6.4  
Diabetics: = 6.5 or higher  
ADA Target: 7.0  
Action suggested: > 6.5

Method:- HPLC

Instrument name: ARKRAY's ADAMS Lite HA 8380V, JAPAN.

#### Test Interpretation:

HbA1C is formed by the condensation of glucose with n-terminal valine residue of each beta chain of HbA to form an unstable schiff base. It is the major fraction, constituting approximately 80% of HbA1c. Formation of glycosylated hemoglobin (GHb) is essentially irreversible and the concentration in the blood depends on both the lifespan of the red blood cells (RBC) (120 days) and the blood glucose concentration. The GHb concentration represents the integrated values for glucose over the period of 6 to 8 weeks. GHb values are free of day to day glucose fluctuations and are unaffected by recent exercise or food ingestion. Concentration of plasma glucose concentration in GHb depends on the time interval, with more recent values providing a larger contribution than earlier values. The interpretation of GHb depends on RBC having a normal life span. Patients with hemolytic disease or other conditions with shortened RBC survival exhibit a substantial reduction of GHb. High GHb have been reported in iron deficiency anemia. GHb has been firmly established as an index of long term blood glucose concentrations and as a measure of the risk for the development of complications in patients with diabetes mellitus. The absolute risk of retinopathy and nephropathy are directly proportional to the mean of HbA1C. Genetic variants (e.g. HbS trait, HbC trait), elevated HbF and chemically modified derivatives of hemoglobin can affect the accuracy of HbA1c measurements. The effects vary depending on the specific Hb variant or derivative and the specific HbA1c method.

Ref by ADA 2020

MEAN PLASMA GLUCOSE

131 H mg/dL

Non Diabetic < 100 mg/dL  
Prediabetic 100- 125 mg/dL  
Diabetic 126 mg/dL or Higher

Method:- Calculated Parameter

AJAYSINGH  
Technologist

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**Dr. Chandrika Gupta**  
MBBS.MD ( Path )  
RMC NO. 21021/008037



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### HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
<b>HAEMOGARAM</b>			
HAEMOGLOBIN (Hb)	14.5	g/dL	13.0 - 17.0
TOTAL LEUCOCYTE COUNT	6.52	/cumm	4.00 - 10.00
<b>DIFFERENTIAL LEUCOCYTE COUNT</b>			
NEUTROPHIL	66.6	%	40.0 - 80.0
LYMPHOCYTE	28.9	%	20.0 - 40.0
EOSINOPHIL	1.5	%	1.0 - 6.0
MONOCYTE	2.7	%	2.0 - 10.0
BASOPHIL	0.3	%	0.0 - 2.0
NEUT#	4.35	10 <sup>3</sup> /uL	1.50 - 7.00
LYMPH#	1.89	10 <sup>3</sup> /uL	1.00 - 3.70
EO#	0.09	10 <sup>3</sup> /uL	0.00 - 0.40
MONO#	0.17	10 <sup>3</sup> /uL	0.00 - 0.70
BASO#	0.02	10 <sup>3</sup> /uL	0.00 - 0.10
TOTAL RED BLOOD CELL COUNT (RBC)	5.42	x10 <sup>6</sup> /uL	4.50 - 5.50
HEMATOCRIT (HCT)	43.70	%	40.00 - 50.00
MEAN CORP VOLUME (MCV)	<b>80.8</b> L	fL	83.0 - 101.0
MEAN CORP HB (MCH)	<b>26.7</b> L	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	33.1	g/dL	31.5 - 34.5
<b>PLATELET COUNT</b>			
RDW-CV	14.0	%	11.6 - 14.0
MENTZER INDEX	14.91		

The Mentzer index is used to differentiate iron deficiency anemia from beta thalassemia trait. If a CBC indicates microcytic anemia, these are two of the most likely causes, making it necessary to distinguish between them.

If the quotient of the mean corpuscular volume divided by the red blood cell count is less than 13, thalassemia is more likely. If the result is greater than 13, then iron-deficiency anemia is more likely.

AJAYSINGH  
Technologist

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Lab/Hosp :-

Company :- MediWheel

Sample Type :- EDTA

Sample Collected Time 15/04/2023 09:38:53

Final Authentication : 15/04/2023 11:45:21

### HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
Erythrocyte Sedimentation Rate (ESR)	28 H	mm/hr.	00 - 13

**(ESR) Methodology** : Measurement of ESR by cells aggregation.

**Instrument Name** : Independent form Hematocrit value by Automated Analyzer (Roller-20)

**Interpretation** : ESR test is a non-specific indicator of inflammatory disease and abnormal protein states.

The test is used to detect, follow course of a certain disease (e.g. tuberculosis, rheumatic fever, myocardial infarction)

Levels are higher in pregnancy due to hyperfibrinogenaemia.

The "3-figure ESR"  $\times > 100$  value nearly always indicates serious disease such as a serious infection, malignant paraproteinaemia or connective tissue disease.

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

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Technologist

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 Company :- MediWHEEL

Patient ID :-1223231  
 Ref. By Dr:- BOB  
 Lab/Hosp :-



Sample Type :- PLAIN/SERUM

Sample Collected Time 15/04/2023 09:38:53

Final Authentication : 15/04/2023 11:13:50

### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
<b>LIPID PROFILE</b>			
TOTAL CHOLESTEROL Method:- Enzymatic Endpoint Method	194.94	mg/dl	Desirable <200 Borderline 200-239 High > 240
TRIGLYCERIDES Method:- GPO-PAP	150.31 H	mg/dl	Normal <150 Borderline high 150-199 High 200-499 Very high >500
DIRECT HDL CHOLESTEROL Method:- Direct clearance Method	31.99	mg/dl	Low < 40 High > 60
DIRECT LDL CHOLESTEROL Method:- Direct clearance Method	137.90	mg/dl	Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190
VLDL CHOLESTEROL Method:- Calculated	30.06	mg/dl	0.00 - 80.00
T.CHOLESTEROL/HDL CHOLESTEROL RATIO Method:- Calculated	6.09 H		0.00 - 4.90
LDL / HDL CHOLESTEROL RATIO Method:- Calculated	4.31 H		0.00 - 3.50
TOTAL LIPID Method:- CALCULATED	610.27	mg/dl	400.00 - 1000.00
<p><b>TOTAL CHOLESTEROL</b> InstrumentName:Radox Rx Imola Interpretation: Cholesterol measurements are used in the diagnosis and treatments of lipid lipoprotein metabolism disorders.</p> <p><b>TRIGLYCERIDES</b> InstrumentName:Radox 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.</p> <p><b>DIRECT HDLCHOLESTERO</b> InstrumentName:Radox Rx Imola 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.</p> <p><b>DIRECT LDL-CHOLESTEROL</b> InstrumentName:Radox Rx Imola 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.</p> <p><b>TOTAL LIPID AND VLDL ARE CALCULATED</b></p>			

SURENDRAKHANGA

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Sample Type :- PLAIN/SERUM

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### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
<b>LIVER PROFILE WITH GGT</b>			
SERUM BILIRUBIN (TOTAL) Method:- Colorimetric method	1.36	mg/dl	Up to - 1.0 Cord blood <2 Premature < 6 days <16 Full-term < 6 days= 12 1month - <12 months <2 1-19 years <1.5 Adult - Up to - 1.2 Ref-(ACCP 2020)
SERUM BILIRUBIN (DIRECT) Method:- Colorimetric Method	0.37	mg/dL	Adult - Up to 0.25 Newborn - <0.6 mg/dL >- 1 month - <0.2 mg/dL
SERUM BILIRUBIN (INDIRECT) Method:- Calculated	0.99	mg/dl	0.30-0.70
SGOT Method:- IFCC	32.7	U/L	Men- Up to - 37.0 Women - Up to - 31.0
SGPT Method:- IFCC	67.0 H	U/L	Men- Up to - 40.0 Women - Up to - 31.0
SERUM ALKALINE PHOSPHATASE Method:- AMP Buffer	81.70	IU/L	30.00 - 120.00
SERUM GAMMA GT Method:- IFCC	28.70	U/L	11.00 - 50.00
SERUM TOTAL PROTEIN Method:- Biuret Reagent	7.42	g/dl	6.40 - 8.30
SERUM ALBUMIN Method:- Bromocresol Green	4.36	g/dl	3.80 - 5.00
SERUM GLOBULIN Method:- CALCULATION	3.06	gm/dl	2.20 - 3.50
A/G RATIO	1.42		1.30 - 2.50

**Total Bilirubin** Methodology: Colorimetric method InstrumentName Randox Rx Imola Interpretation An increase in bilirubin concentration in the serum occurs in toxic or infectious diseases of the liver e.g. hepatitis B or obstruction of the bile duct and in rhesus incompatible babies High levels of unconjugated bilirubin indicate that too much haemoglobin is being destroyed or that the liver is not actively treating the haemoglobin it is receiving.

**AST Aspartate Aminotransferase** Methodology: IFCC InstrumentName Randox Rx Imola Interpretation: Elevated levels of AST can signal myocardial infarction, hepatic disease, muscular dystrophy and organ damage. Although heart muscle is found to have the most activity of the enzyme, significant activity has also been seen in the brain, liver, gastric mucosa, adipose tissue and kidneys of humans.

**ALT Alanine Aminotransferase** Methodology: IFCC InstrumentName Randox Rx Imola Interpretation: The enzyme ALT has been found to be in highest concentrations in the liver, with decreasing concentrations found in kidney, heart, skeletal muscle, pancreas, spleen and lung tissue respectively. Elevated levels of the transaminases can indicate myocardial infarction, hepatic disease, muscular dystrophy and organ damage.

**Alkaline Phosphatase** Methodology: AMP Buffer InstrumentName Randox Rx Imola Interpretation: Measurements of alkaline phosphatase are of use in the diagnosis, treatment and investigation of hepatobiliary disease and in bone disease associated with increased osteoblastic activity. Alkaline phosphatase is also used in the diagnosis of parathyroid and intestinal disease.

**TOTAL PROTEIN** Methodology: Biuret Reagent InstrumentName Randox Rx Imola 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.

**ALBUMIN (ALB)** Methodology: Bromocresol Green InstrumentName Randox Rx Imola Interpretation: Albumin measurements are used in the diagnosis and treatment of numerous diseases involving primarily the liver or kidneys. Globulin & A/G ratio is calculated.

**Instrument Name Randox Rx Imola Interpretation:** Elevations in GGT levels are seen 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)

SURENDRAKHANGA

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Patient ID :-1223231

Ref. By Dr:- BOB

Lab/Hosp :-



Sample Type :- PLAIN/SERUM

Sample Collected Time 15/04/2023 09:38:53

Final Authentication : 15/04/2023 11:39:19

### IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
<b>TOTAL THYROID PROFILE</b>			
SERUM TOTAL T3 Method:- Chemiluminescence(Competitive immunoassay)	1.145	ng/ml	0.970 - 1.690
SERUM TOTAL T4 Method:- Chemiluminescence(Competitive immunoassay)	7.628	ug/dl	5.530 - 11.000
SERUM TSH ULTRA Method:- Enhanced Chemiluminescence Immunoassay	4.700	μIU/mL	0.550 - 4.780

**Interpretation:** Triiodothyronine (T3) contributes to the maintenance of the euthyroid state. A decrease in T3 concentration of up to 50% occurs in a variety of clinical situations, including acute and chronic disease. Although T3 results alone cannot be used to diagnose hypothyroidism, T3 concentration may be more sensitive than thyroxine (T4) for hyperthyroidism. Consequently, the total T3 assay can be used in conjunction with other assays to aid in the differential diagnosis of thyroid disease. T3 concentrations may be altered in some conditions, such as pregnancy, that affect the capacity of the thyroid hormone-binding proteins. Under such conditions, Free T3 can provide the best estimate of the metabolically active hormone concentration. Alternatively, T3 uptake, or T4 uptake can be used with the total T3 result to calculate the free T3 index and estimate the concentration of free T3.

**Interpretation:** The measurement of Total T4 aids in the differential diagnosis of thyroid disease. While >99.9% of T4 is protein-bound, primarily to thyroxine-binding globulin (TBG), it is the free fraction that is biologically active. In most patients, the total T4 concentration is a good indicator of thyroid status. T4 concentrations may be altered in some conditions, such as pregnancy, that affect the capacity of the thyroid hormone-binding proteins. Under such conditions, free T4 can provide the best estimate of the metabolically active hormone concentration. Alternatively, T3 uptake may be used with the total T4 result to calculate the free T4 index (FT4I) and estimate the concentration of free T4. Some drugs and some nonthyroidal patient conditions are known to alter TT4 concentrations in vivo.

**Interpretation:** TSH stimulates the production of thyroxine (T4) and triiodothyronine (T3) by the thyroid gland. The diagnosis of overt hypothyroidism by the finding of a low total T4 or free T4 concentration is readily confirmed by a raised TSH concentration. Measurement of low or undetectable TSH concentrations may assist the diagnosis of hyperthyroidism, where concentrations of T4 and T3 are elevated and TSH secretion is suppressed. These have the advantage of discriminating between the concentrations of TSH observed in thyrotoxicosis, compared with the low, but detectable, concentrations that occur in subclinical hyperthyroidism. The performance of this assay has not been established for neonatal specimens. Some drugs and some nonthyroidal patient conditions are known to alter TSH concentrations in vivo.

### INTERPRETATION

PREGNANCY	REFERENCE RANGE FOR TSH IN uIU/mL (As per American Thyroid Association)
1st Trimester	0.10-2.50
2nd Trimester	0.20-3.00
3rd Trimester	0.30-3.00

AJAYKUMAR  
Technologist

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Dr. Chandrika Gupta  
MBBS.MD ( Path )  
RMC NO. 21021/008037

# Dr. Goyal's

## Path Lab & Imaging Centre



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Website: www.drgoyalpathlab.com | E-mail: drgoyalpiyush@gmail.com

Date :- 15/04/2023 09:25:18  
**NAME :- Mr. MANOJ GUPTA**  
Sex / Age :- Male 53 Yrs  
Company :- MediWheel

Patient ID :- 1223231  
Ref. By Dr:- BOB  
Lab/Hosp :-



Sample Type :- URINE

Sample Collected Time 15/04/2023 09:38:53

Final Authentication : 15/04/2023 15:50:02

### CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
<b>Urine Routine</b>			
<b>PHYSICAL EXAMINATION</b>			
COLOUR	PALE YELLOW		PALE YELLOW
APPEARANCE	Clear		Clear
<b>CHEMICAL EXAMINATION</b>			
REACTION(PH) Method:- Reagent Strip(Double indicator blue reaction)	6.5		5.0 - 7.5
SPECIFIC GRAVITY Method:- Reagent Strip(bromthymol blue)	1.025		1.010 - 1.030
PROTEIN Method:- Reagent Strip (Sulphosalicylic acid test)	NIL		NIL
GLUCOSE Method:- Reagent Strip (Glu.Oxidase Peroxidase Benedict)	NIL		NIL
BILIRUBIN Method:- Reagent Strip (Azo-coupling reaction)	NEGATIVE		NEGATIVE
UROBILINOGEN Method:- Reagent Strip (Modified ehrlich reaction)	NORMAL		NORMAL
KETONES Method:- Reagent Strip (Sodium Nitropruside) Rothera's	NEGATIVE		NEGATIVE
NITRITE Method:- Reagent Strip (Diazotization reaction)	NEGATIVE		NEGATIVE
RBC Method:- Reagent Strip (Peroxidase like activity)	NIL		NIL
<b>MICROSCOPY EXAMINATION</b>			
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		ABSENT

VIJENDRAMEENA  
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Page No: 7 of 13



**Dr. Rashmi Bakshi**  
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Date :- 15/04/2023 09:25:18

Patient ID :-1223231



NAME :- Mr. MANOJ GUPTA

Ref. By Dr:- BOB

Sex / Age :- Male 53 Yrs

Lab/Hosp :-

Company :- MediWheel

Sample Type :- STOOL

Sample Collected Time 15/04/2023 09:38:53

Final Authentication : 15/04/2023 15:50:02

### CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
<b>STOOL ANALYSIS</b>			
<b>PHYSICAL EXAMINATION</b>			
MUCUS			
BLOOD			
<b>MICROSCOPIC EXAMINATION</b>			
RBC's		/HPF	
WBC/HPF		/HPF	
OVA			
CYSTS			
OTHERS			
Collected Sample Received			

VIJENDRAMEENA  
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Date :- 15/04/2023 09:25:18

NAME :- Mr. MANOJ GUPTA

Sex / Age :- Male 53 Yrs

Company :- MediWheel

Patient ID :-1223231

Ref. By Dr:- BOB

Lab/Hosp :-



### HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
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AHSAN, AJAYKUMAR, AJAYSINGH, BILAL, MUKESH SINGH, SURENDRAKHANGA, TABBSUM, VIJENDRAMEENA

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# Dr. Goyal's

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Date :- 15/04/2023 09:25:18  
**NAME :- Mr. MANOJ GUPTA**  
Sex / Age :- Male 53 Yrs  
Company :- MediWheel

Patient ID :-1223231  
Ref. By Dr:- BOB  
Lab/Hosp :-



Sample Type :- EDTA, URINE

Sample Collected Time 15/04/2023 09:38:53

Final Authentication : 15/04/2023 15:50:02

### HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
BLOOD GROUP ABO	"O" POSITIVE		
BLOOD GROUP ABO Methodology : Haemagglutination reaction Kit Name : Monoclonal agglutinating antibodies (Span clone).			
URINE SUGAR (FASTING) Collected Sample Received	Nil		Nil

AJAYSINGH, VIJENDRAMEENA  
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Date :- 15/04/2023 09:25:18

Patient ID :- 1223231



**NAME :- Mr. MANOJ GUPTA**

Ref. By Dr:- BOB

Sex / Age :- Male 53 Yrs

Lab/Hosp :-

Company :- MediWheel

Sample Type :- PLAIN/SERUM

Sample Collected Time 15/04/2023 09:38:53

Final Authentication : 15/04/2023 11:13:50

### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
BLOOD UREA NITROGEN (BUN)	9.7	mg/dl	0.0 - 23.0

SURENDRAKHANGA

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**Dr. Chandrika Gupta**  
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# Dr. Goyal's

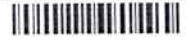
## Path Lab & Imaging Centre

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Sex / Age :- Male 53 Yrs  
Company :- MediWheel

Patient ID :-1223231  
Ref. By Dr:- BOB  
Lab/Hosp :-



Sample Type :- PLAIN/SERUM

Sample Collected Time 15/04/2023 09:38:53

Final Authentication : 15/04/2023 11:39:19

### IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
TOTAL PSA Method:- Chemiluminescence	0.463	ng/ml	0.000 - 4.000

**InstrumentName:** VITROS ECI **Interpretation :** Elevated serum PSA concentrations are found in men with prostate cancer, benign prostatic hypertrophy (BHP) or inflammatory conditions of other adjacent genitourinary tissues, but not in apparently healthy men or in men with cancers other than prostate cancer. PSA has been demonstrated to be an accurate marker for monitoring advancing clinical stage in untreated patients and for monitoring response to therapy by radical prostatectomy, radiation therapy and anti-androgen therapy. PSA is also important in determining the potential and actual effectiveness of surgery or other therapies. Progressive disease is defined by an increase of at least 25%. Sampling should be repeated within two to four weeks for additional evidence. Different assay methods cannot be used interchangeably.

\*\*\* End of Report \*\*\*

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