

भारत सरकार  
Government of India

Naresh Kumar Kumawat  
Year of Birth : 1966  
Male



4361 0298 9611

आधार - आम आदमी का अधिकार

Dr. Piyush Goyal  
M.B.P.S., D.M.F.L.  
RMC Reg. No.-017990

आधार  
Unique Identification Authority of India

Address:  
S/O: Hari Ram Kumawat, Tihawali, Tihawali, Sikar,  
Fatehpur, Rajasthan, 332307

4361 0298 9611

1947  
1800 300 1947

help@uidai.gov.in

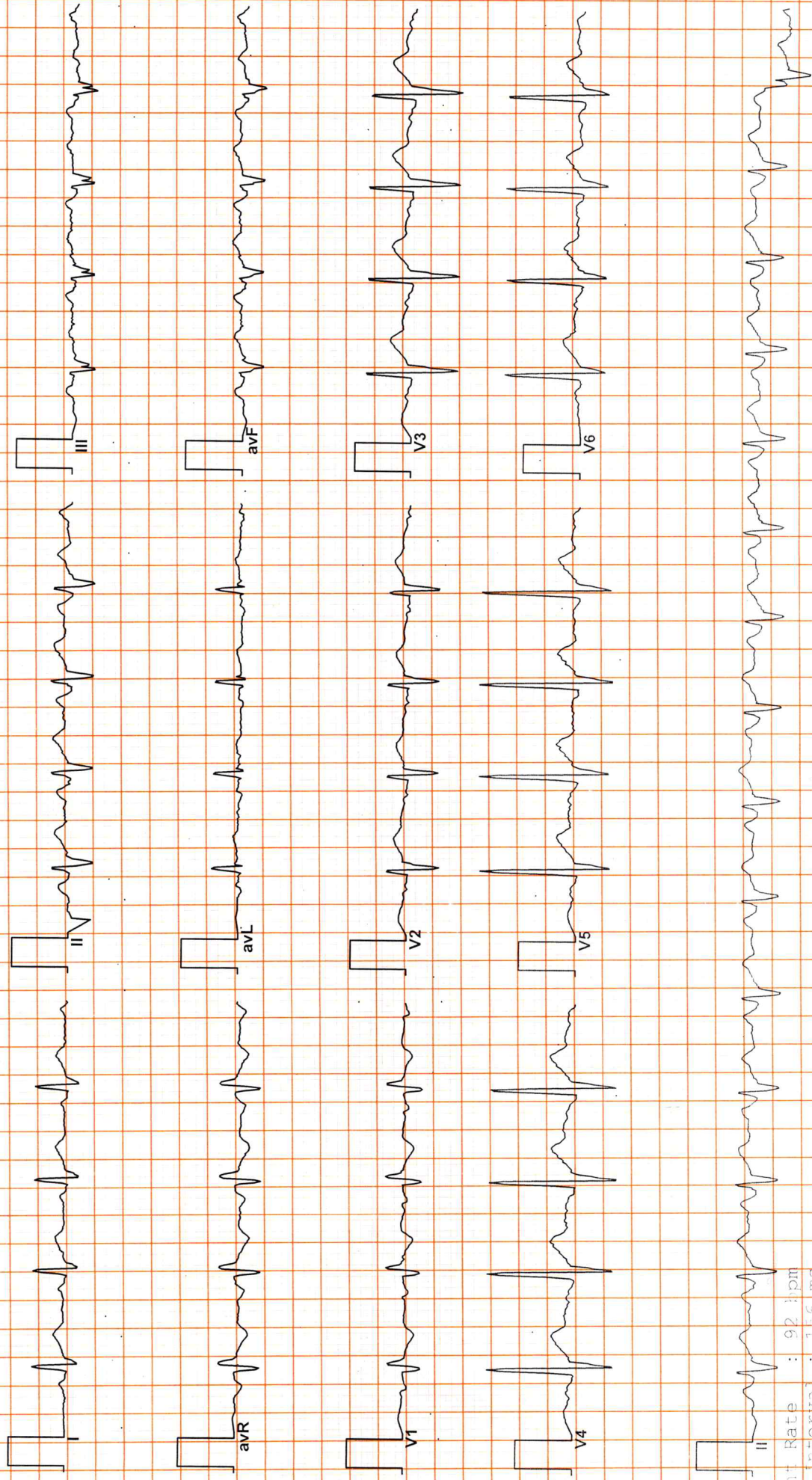
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www.uidai.gov.in

**DR. GOYALS PATH LAB & IMAGING CENTRE**

116 / MR. NARESH KUMAR KUMAWAT / 55 Yrs / M / 169Cms. / 52Kgs. / Non Smoker

Heart Rate : 92 bpm / Tested On : 28-Mar-22 14:37:22 / HF 0.05 Hz - LF 100 Hz / Notch 50 Hz / Sn 1.00 Cm/mV / Sw 25 mm/s / Refd By: BOB

**ECG**



Heart Rate : 92 bpm  
PR Interval : 156 ms  
QRS Duration : 96 ms  
QT/QTc Int : 352/408 ms  
P-QRS-T axis: 66.00 • -29.00 • 53.00 •

*[Signature]*  
DR. P. JAYARAJ  
M.B.B.S., C.I.F.I.  
REG. NO. - 17596

# Dr. Goyal's

## Path Lab & Imaging Centre



B-51, Ganesh Nagar, Opp. Janpath Corner, New Sangner Road, Jaipur-302019

Tele: 0141-2293346, 4049787, 9887049787

Website: www.dr.goyalspathlab.com | E-mail: dr.goyalpiyush@gmail.com

Date :- 28/03/2022 10:40:19

NAME :- Mr. NARESH KUMAR KUMAWAT

Sex / Age :- Male 55 Yrs 8 Mon 23 Days

Company :- MediWheel

Patient ID :- 122127992

Ref. By Dr:- BOB

Lab/Hosp :-



Sample Type :- EDTA

Sample Collected Time 28/03/2022 10:44:52

Final Authentication : 28/03/2022 12:36:10

### HAEMATOLOGY

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

**GLYCOSYLATED HEMOGLOBIN (HbA1C)**

Method:- HPLC

5.7

%

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

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

Method:- Calculated Parameter

117

mg/dL

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

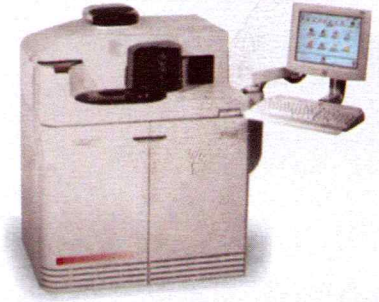
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Technologist

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### HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
<b>HAEMOGARAM</b>			
HAEMOGLOBIN (Hb)	14.9	g/dL	13.0 - 17.0
TOTAL LEUCOCYTE COUNT	9.13	/cumm	4.00 - 10.00
<b>DIFFERENTIAL LEUCOCYTE COUNT</b>			
NEUTROPHIL	61.9	%	40.0 - 80.0
LYMPHOCYTE	32.1	%	20.0 - 40.0
EOSINOPHIL	2.0	%	1.0 - 6.0
MONOCYTE	3.7	%	2.0 - 10.0
BASOPHIL	0.3	%	0.0 - 2.0
NEUT#	5.66	10 <sup>3</sup> /uL	1.50 - 7.00
LYMPH#	2.93	10 <sup>3</sup> /uL	1.00 - 3.70
EO#	0.18	10 <sup>3</sup> /uL	0.00 - 0.40
MONO#	0.33	10 <sup>3</sup> /uL	0.00 - 0.70
BASO#	0.03	10 <sup>3</sup> /uL	0.00 - 0.10
TOTAL RED BLOOD CELL COUNT (RBC)	4.79	x10 <sup>6</sup> /uL	4.50 - 5.50
HEMATOCRIT (HCT)	44.40	%	40.00 - 50.00
MEAN CORP VOLUME (MCV)	92.7	fL	83.0 - 101.0
MEAN CORP HB (MCH)	31.2	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	33.7	g/dL	31.5 - 34.5
<b>PLATELET COUNT</b>	189	x10 <sup>3</sup> /uL	150 - 410
RDW-CV	<b>14.1</b> H	%	11.6 - 14.0
MENTZER INDEX	19.35		

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.

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### HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
-----------	-------	------	-------------------------

Erythrocyte Sedimentation Rate (ESR)	38 H	mm/hr.	00 - 13
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**(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 (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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Company :- MediWheel



Sample Type :- PLAIN/SERUM

Sample Collected Time 28/03/2022 10:44:52

Final Authentication : 28/03/2022 12:22:42

### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
<b>LIPID PROFILE</b>			
TOTAL CHOLESTEROL Method:- Enzymatic Endpoint Method	182.52	mg/dl	Desirable <200 Borderline 200-239 High > 240
TRIGLYCERIDES Method:- GPO-PAP	133.31	mg/dl	Normal <150 Borderline high 150-199 High 200-499 Very high >500
VLDL CHOLESTEROL Method:- Calculated	26.66	mg/dl	0.00 - 80.00

JITENDRAKUMAWAT

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### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
DIRECT HDL CHOLESTEROL Method:- Direct clearance Method	52.50	mg/dl	Low < 40 High > 60
DIRECT LDL CHOLESTEROL Method:- Direct clearance Method	107.80	mg/dl	Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190
T.CHOLESTEROL/HDL CHOLESTEROL RATIO Method:- Calculated	3.48		0.00 - 4.90
LDL / HDL CHOLESTEROL RATIO Method:- Calculated	2.05		0.00 - 3.50
TOTAL LIPID Method:- CALCULATED	565.07	mg/dl	400.00 - 1000.00
<b>TOTAL CHOLESTEROL InstrumentName:</b> Radox Rx Imola <b>Interpretation:</b> Cholesterol measurements are used in the diagnosis and treatments of lipid lipoprotein metabolism disorders.			
<b>TRIGLYCERIDES InstrumentName:</b> Radox Rx Imola <b>Interpretation :</b> 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.			
<b>DIRECT HDLCHOLESTERO InstrumentName:</b> Radox Rx Imola <b>Interpretation:</b> 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.			
<b>DIRECT LDL-CHOLESTEROL InstrumentName:</b> Radox Rx Imola <b>Interpretation:</b> 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.			
<b>TOTAL LIPID AND VLDL ARE CALCULATED</b>			

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Sample Collected Time 28/03/2022 10:44:52

Final Authentication : 28/03/2022 12:22:42

### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
<b>LIVER PROFILE WITH GGT</b>			
SERUM BILIRUBIN (TOTAL) Method:- Colorimetric method	0.89	mg/dl	Up to - 1.0 Cord blood <2 mg/dL Premature < 6 days <16mg/dL Full-term < 6 days= 12 mg/dL 1month - <12 months <2 mg/dL 1-19 years <1.5 mg/dL Adult - Up to - 1.2 Ref-(ACCP 2020)
SGOT Method:- IFCC	31.2	U/L	Men- Up to - 37.0 Women - Up to - 31.0
SGPT Method:- IFCC	38.6	U/L	Men- Up to - 40.0 Women - Up to - 31.0
SERUM ALKALINE PHOSPHATASE Method:- AMP Buffer	88.30	IU/L	30.00 - 120.00
SERUM TOTAL PROTEIN Method:- Biuret Reagent	7.32	g/dl	6.40 - 8.30
SERUM ALBUMIN Method:- Bromocresol Green	4.50	g/dl	3.80 - 5.00
SERUM GLOBULIN Method:- CALCULATION	2.82	gm/dl	2.20 - 3.50
A/G RATIO	1.60		1.30 - 2.50

JITENDRAKUMAWAT

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### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
SERUM BILIRUBIN (DIRECT) Method:- Colorimetric Method	0.30	mg/dL	Adult - Up to 0.25 Newborn - <0.6 mg/dL >- 1 month - <0.2 mg/dL
SERUM BILIRUBIN (INDIRECT) Method:- Calculated	0.59	mg/dl	0.30-0.70
SERUM GAMMA GT Method:- IFCC	<b>187.10</b> H	U/L	11.00 - 50.00

**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) are observed with infectious hepatitis.

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Company :- MediWheel

Sample Type :- PLAIN/SERUM

Sample Collected Time 28/03/2022 10:44:52

Final Authentication : 28/03/2022 12:05:41



### IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
<b>TOTAL THYROID PROFILE</b>			
SERUM TSH ULTRA Method:- Enhanced Chemiluminescence Immunoassay	3.6720	μIU/mL	0.4001 - 4.0490

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### IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
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SERUM TOTAL T3 1.240 ng/ml 0.970 - 1.690  
**Method:- Chemiluminescence(Competitive immunoassay)**

SERUM TOTAL T4 10.800 ug/dl 5.530 - 11.000  
**Method:- Chemiluminescence(Competitive immunoassay)**

**InstrumentName:** VITROS ECI **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.

**InstrumentName:** VITROS ECI **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.

**InstrumentName:** VITROS ECI **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

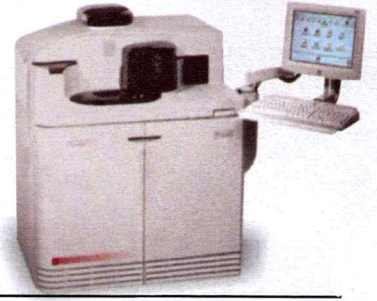
ANANDSHARMA  
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Sample Type :- URINE

Sample Collected Time 28/03/2022 10:44:52

Final Authentication : 28/03/2022 12:26:48

### CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
<b>Urine Routine</b>			
<b><u>MICROSCOPY EXAMINATION</u></b>			
RBC/HPF	NIL	/HPF	NIL
WBC/HPF	2-3	/HPF	2-3
EPITHELIAL CELLS	1-2	/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		ABSENT

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### CLINICAL PATHOLOGY

Test Name	Value	Unit	Biological Ref Interval
<b><u>PHYSICAL EXAMINATION</u></b>			
COLOUR	PALE YELLOW		PALE YELLOW
APPEARANCE	Clear		Clear
<b><u>CHEMICAL EXAMINATION</u></b>			
REACTION(PH)	5.5		5.0 - 7.5
SPECIFIC GRAVITY	1.025		1.010 - 1.030
PROTEIN	NIL		NIL
SUGAR	NIL		NIL
BILIRUBIN	NEGATIVE		NEGATIVE
UROBILINOGEN	NORMAL		NORMAL
KETONES	NEGATIVE		NEGATIVE
NITRITE	NEGATIVE		NEGATIVE

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Technologist

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Sample Type :- KOx/Na FLUORIDE-F, KOx/Na Sodium Chloride and Serum  
 Date of Report :- 28/03/2022 15:24:37 Final Authentication : 28/03/2022 16:12:34

### BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
FASTING BLOOD SUGAR (Plasma) Method:- GOD PAP	124.6 H	mg/dl	75.0 - 115.0
<b>Impaired glucose tolerance (IGT)</b>	111 - 125 mg/dL		
<b>Diabetes Mellitus (DM)</b>	> 126 mg/dL		
<p><b>Instrument Name:</b> Randox Rx Imola <b>Interpretation:</b> 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 .</p>			
BLOOD SUGAR PP (Plasma) Method:- GOD PAP	216.3 H	mg/dl	70.0 - 140.0
<p><b>Instrument Name:</b> Randox Rx Imola <b>Interpretation:</b> 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 .</p>			
SERUM CREATININE Method:- Colorimetric Method	0.96	mg/dl	Men - 0.6-1.30 Women - 0.5-1.20
SERUM URIC ACID Method:- Enzymatic colorimetric	6.07	mg/dl	Men - 3.4-7.0 Women - 2.4-5.7

JITENDRAKUMAWAT

Page No: 12 of 15



**Dr. Piyush Goyal**  
 (D.M.R.D.)  
**DR. TANURUNGTA**

# 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



Date :- 28/03/2022 10:40:19

Patient ID :- 122127992

**NAME :- Mr. NARESH KUMAR KUMAWAT**

Ref. By Dr:- BOB

Sex / Age :- Male 55 Yrs 8 Mon 23 Days

Lab/Hosp :-

Company :- MediWheel



Sample Type :- EDTA, PLAIN/SERUM, URINE, SPINE COLLECTED Time 28/03/2022 15:23:44

Final Authentication : 28/03/2022 16:50:56

### HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
BLOOD GROUP ABO	"O" POSITIVE		
<b>BLOOD GROUP ABO Methodology :</b> Haemagglutination reaction <b>Kit Name :</b> Monoclonal agglutinating antibodies (Span clone).			
URINE SUGAR (FASTING) Collected Sample Received	trace		Nil
URINE SUGAR PP Collected Sample Received	++		Nil
BLOOD UREA NITROGEN (BUN)	13.5	mg/dl	0.0 - 23.0

AJAYSINGH, JITENDRAKUMAWAT, POOJABOHRA, SAPNA  
**Technologist**

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Ref. By Dr:- BOB

Sex / Age :- Male 55 Yrs 8 Mon 23 Days

Lab/Hosp :-

Company :- MediWheel



Sample Type :- PLAIN/SERUM

Sample Collected Time 28/03/2022 10:44:52

Final Authentication : 28/03/2022 12:05:41

### IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
TOTAL PSA Method:- Chemiluminescence	0.802	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 \*\*\*

ANANDSHARMA  
Technologist

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**NAME :- Mr. NARESH KUMAR KUMAWAT**  
Sex / Age :- Male 55 Yrs 8 Mon 23 Days  
Company :- MediWheel

Patient ID :- 122127992  
Ref. By Doctor :- BOB  
Lab/Hosp :-

Final Authentication : 28/03/2022 14:00:11

BOB PACKAGE ABOVE 40MALE

### X RAY CHEST PA VIEW:

**Bronchovascular marking are prominent.**  
**Left hila is prominent.**  
**Old healed fracture of 6th posterior rib on right side.**  
Otherwise lung fields are clear.  
Trachea is in midline.  
Both the C.P.angles is clear.  
Both the domes of diaphragm are normally placed.  
Heart shadows appear normal.

(Please correlate clinically and with relevant further investigations.)

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

Page No: 1 of 1

**Dr. Piyush Goyal**  
(D.M.R.D.) BILAL

**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. Tej Prakash Gupta**  
MBBS, DMRD, UCAM  
Fetal Medicine Specialist  
RMC No 24436 FMF ID 102534

**Dr. Rathod Hetali Amrutlal**  
MBBS, M.D. (Radio-Diagnosis)  
RMC No. 17163

Transcript by

Print Copy

# Dr. Goyal's

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B-51, Ganesh Nagar, Opp. Janpath Corner, New Sanganer Road, Jaipur  
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Sex / Age :- Male 55 Yrs 8 Mon 23 Days  
Company :- MediWheel

Patient ID :- 122127992  
Ref. By Doctor:-BOB  
Lab/Hosp :-

Final Authentication : 28/03/2022 15:32:06

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 ( head & neck ) appear non homogenous in echopattern.** Rest part is obscured due to bowel gases.

**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 (16 gms) with normal echo-texture and outline.

No enlarged nodes are visualised.No retro-peritoneal lesion is identified  
Great vessels appear normal.No significant free fluid is seen in peritoneal cavity.

### IMPRESSION:

**Non homogenous echopattern of visualised pancreas (Adv: serum amylase & lipase)  
Needs clinical correlation for further evaluation**

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

Page No: 1 of 1

SAVITA

**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. Tej Prakash Gupta**  
DMRD (RADIO DIAGNOSIS)  
RMC No. 24436

**Dr. Hitesh Kumar Sharma**  
M.B.B.S., D.M.R.D.  
RMC Reg No. 27380

Transcript by.

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 Sex / Age :- Male 55 Yrs 8 Mon 23 Days Lab/Hosp :-  
 Company :- MediWheel



Sample Type :- Sample Collected Time Final Authentication : 28/03/2022 14:42:45

### ECHOCARDIOGRAPHY 2D (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	31	mm	LA	34	Mm	IVS-D	10	mm
IVS-S	11	mm	LVID	40	Mm	LVSD	28	mm
LVPW-D	7	mm	LVPW-S	11	Mm	RV		mm
RVWT		mm	EDV		MI	LVVS		ml
LVEF	60%		RWMA			ABSENT		

##### CHAMBERS:

LA	NORMAL	RA	NORMAL
LV	NORMAL	RV	NORMAL
PERICARDIUM	NORMAL		

##### COLOUR DOPPLER:

MITRAL VALVE					
E VELOCITY	0.88	m/sec	PEAK GRADIENT		Mm/hg
A VELOCITY		m/sec	MEAN GRADIENT		Mm/hg
MVA BY PHT		Cm2	MVA BY PLANIMETRY		Cm2
MITRAL REGURGITATION		ABSENT			
AORTIC VALVE					
PEAK VELOCITY	1.0	m/sec	PEAK GRADIENT		mm/hg
AR VMAX		m/sec	MEAN GRADIENT		mm/hg
AORTIC REGURGITATION		ABSENT			
TRICUSPID VALVE					
PEAK VELOCITY	0.68	m/sec	PEAK GRADIENT		mm/hg
MEAN VELOCITY		m/sec	MEAN GRADIENT		mm/hg
VMax VELOCITY					
TRICUSPID REGURGITATION		ABSENT			
PULMONARY VALVE					
PEAK VELOCITY	0.82	M/sec.	PEAK GRADIENT		Mm/hg
MEAN VELOCITY			MEAN GRADIENT		Mm/hg
PULMONARY REGURGITATION		ABSENT			

TANVI

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Sample Type :- Sample Collected Time Final Authentication : 28/03/2022 14:42:45

### Impression--

1. Normal LV size & contractility.
2. No RWMA, LVEF 60%.
3. Normal cardiac chamber.
4. Normal valve.
5. No clot, no vegetation, no pericardial effusion.

(Cardiologist)



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

TANVI

Page No: 2 of 2



# Dr. Goyal's Path Lab

Name **NARESH KUMAR**  
 Patient Id **NARES07\_07725**

Date **03/28/2022**  
 Diagnosis Dr.

