

General Physical Examination

Date of Examination: 14/05/23

Name: Kailash chand Sahu Age: 52 Sex: male

DOB: 09/06/1970

Referred By: Medi wheel

Photo ID: Adhar ID #: attached

Ht: 161 (cm)

Wt: 66 (Kg)

Chest (Expiration): 96 (cm)

Abdomen Circumference: 90 (cm)

Blood Pressure: 130/95 mm Hg PR: 79 / min RR: 16 / min Temp: Apoite

BMI 25.5

Eye Examination: Dis vision 6/6. with spec near vision N/6
No colour blindness

Other: Not significant.

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

Signature Of Examinee : [Signature]

Name of Examinee Dr. Piyush Goyal

M.B.B.S, D.M.R.D
RMC Reg No -017998

Signature Medical Examiner : _____

Name Medical Examiner _____

भारत सरकार
Government of India

कैलाश चंद साह
Kailash Chand Sahu
जन्म तिथि / DOB : 09/06/1970
पुरुष / MALE

9210 7155 4336

मेरा आधार, मेरी पहचान

कैलाश

Dr Piyush Goyal
M.B.B.S, D.M.R.D
RMQ Reg No -017953

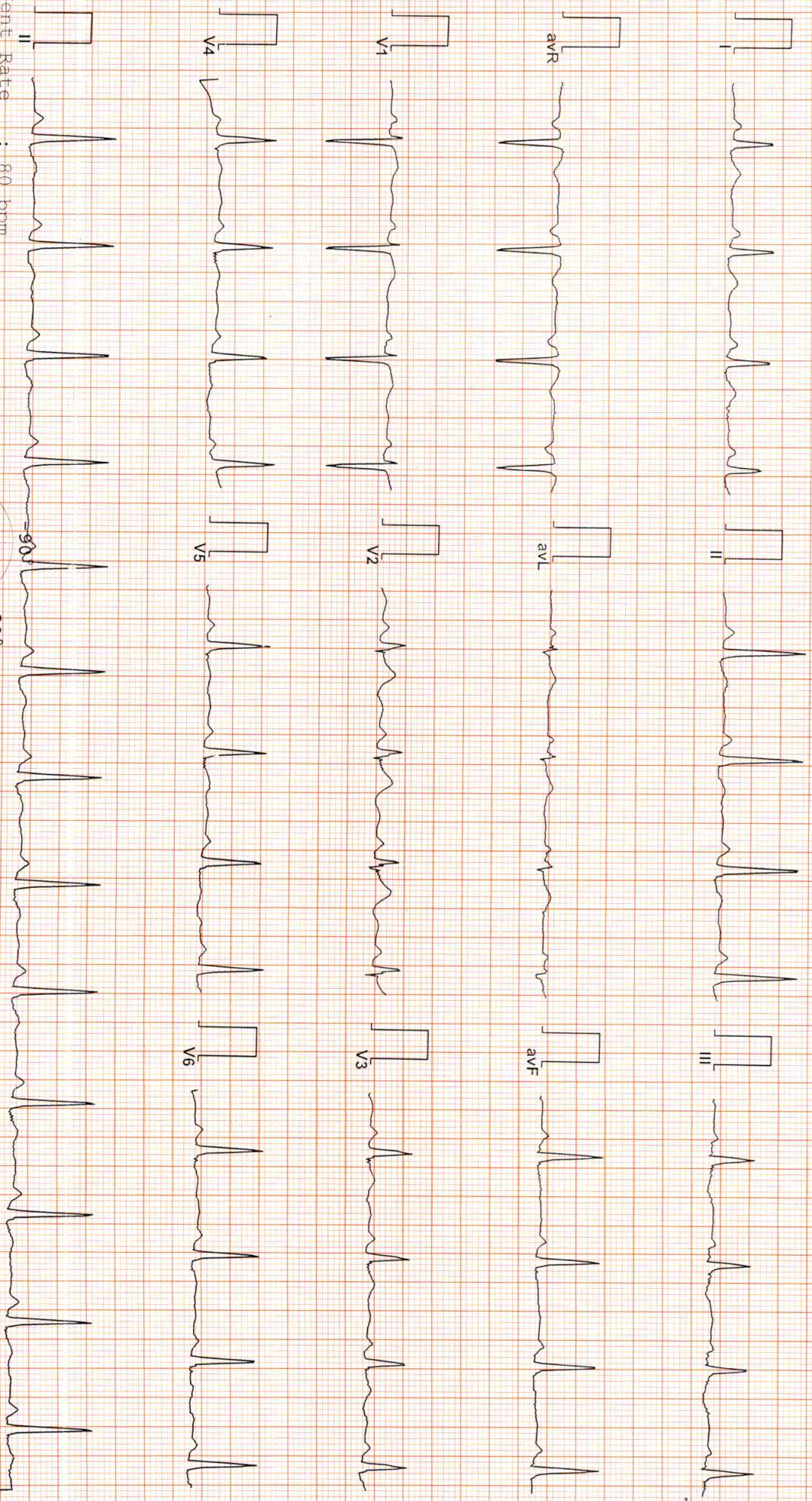
आधार
Unique Identification Authority of India

पता: S/O रामबिलास साह, 147, आदिनाथ
नगर, पालवाले बालाजी मोड, सिरसी रोड,
सिरसी, सिरसी, जयपुर, राजस्थान, 302012

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nagar, pallwale balaji mod, sirsi road, Sirsi,
Sirsi, Jaipur, Rajasthan, 302012

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Vent Rate : 80 bpm
 PR Interval : 148 ms
 QRS Duration: 76 ms
 QT/QTc Int : 326/361 ms
 P-QRS-T axis: 53.00° 61.00° 5.00°

180°
 90°
 -30°

Dr. Karan Kumar Motwani
 RMC No. 35103
 MBBS, D.P. CARDIOL (D)
 O.E.M. (RCGP-UK)

Sinus bradycardia with 1st degree AVB

Reported By: [Signature]

NAME:	KAILASH CHAND SAHU/ 1223696	AGE	52 YRS
REF.BY	BOB	DATE	14-05-2023

USG WHOLE ABDOMEN

Liver is enlarged in size (~16.3cm). Echo-texture is bright. 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. Collecting system does not show any dilatation or calculus.
Few (2-3) well defined anechoic cysts are seen in right kidney, one of them calcification & septation, largest measuring ~ 17x11mm at lower pole.

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:

- *Mild hepatomegaly with Grade I fatty changes.
 - * Right renal cortical cysts as described above (suggested follow up)
- Needs clinical correlation for further evaluation

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NAME:	MR. KAILASH CHAND SAHU	AGE	52/YRS
REF.BY	BOB	DATE	

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	27	mm	LA	33	Mm	IVS-D	8	mm
IVS-S	13	mm	LVID	42	Mm	LVSD	30	mm
LVPW-D	10	mm	LVPW-S	16	Mm	RV		mm
RVWT		mm	EDV		ml	LVVS		ml
LVEF	54%		RWMA			ABSENT		

CHAMBERS:

LA	NORMAL	RA	NORMAL
LV	NORMAL	RV	NORMAL
PERICARDIUM		NORMAL	

COLOUR DOPPLER:

MITRAL VALVE					
E VELOCITY	0.97	m/sec	PEAK GRADIENT		Mm/hg
A VELOCITY	0.81	m/sec	MEAN GRADIENT		Mm/hg
MVA BY PHT		Cm2	MVA BY PLANIMETRY		Cm2
MITRAL REGURGITATION			ABSENT		
AORTIC VALVE					
PEAK VELOCITY	1.29	m/sec	PEAK GRADIENT		mm/hg
AR VMAX		m/sec	MEAN GRADIENT		mm/hg
AORTIC REGURGITATION			ABSENT		
TRICUSPID VALVE					
PEAK VELOCITY	0.41	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		

Impression--

- Normal LV size & contractility.
- No RWMA, LVEF 54 %.
- Normal cardiac chamber.
- Normal valve.
- No clot, no vegetation, no pericardial effusion. (Cardiologist)

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 RMC Reg No. 017996

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

Dr. Ashish Choudhary
 MBBS, MD (Radio Diagnosis)
 Fetal Medicine Consultant

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

Transcript by.

FMF ID - 260517 | RMC No 22430

This report is not valid for medico-legal purpose.

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X RAY CHEST PA VIEW:

Positional rotation present.

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.

Bony cage and soft tissue shadows are normal.

Heart shadows appear normal.

Impression :- Normal Study

(Please correlate clinically and with relevant further investigations)

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MC- 5509

Date :- 14/05/2023 09:13:54

NAME :- Mr. KAILASH CHAND SHAU

Sex / Age :- Male 52 Yrs 11 Mon 5 Days

Company :- MediWheel

Patient ID :-1223696

Ref. By Dr:- BOB

Lab/Hosp :-



Sample Type :- EDTA

Sample Collected Time 14/05/2023 09:30:33

Final Authentication : 14/05/2023 14:13:27

HAEMATOLOGY

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

GLYCOSYLATED HEMOGLOBIN (HbA1C)

Method:- HPLC

6.0

%

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

126

mg/dL

Non Diabetic < 100 mg/dL

Prediabetic 100- 125 mg/dL

Diabetic 126 mg/dL or Higher

BANWARI
Technologist

Page No: 1 of 13



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Test Name	Value	Unit	Biological Ref Interval
HAEMOGARAM			
HAEMOGLOBIN (Hb)	15.8	g/dL	13.0 - 17.0
TOTAL LEUCOCYTE COUNT	6.47	/cumm	4.00 - 10.00
DIFFERENTIAL LEUCOCYTE COUNT			
NEUTROPHIL	52.1	%	40.0 - 80.0
LYMPHOCYTE	40.0	%	20.0 - 40.0
EOSINOPHIL	4.1	%	1.0 - 6.0
MONOCYTE	3.4	%	2.0 - 10.0
BASOPHIL	0.4	%	0.0 - 2.0
NEUT#	3.38	10 ³ /uL	1.50 - 7.00
LYMPH#	2.62	10 ³ /uL	1.00 - 3.70
EO#	0.23	10 ³ /uL	0.00 - 0.40
MONO#	0.21	10 ³ /uL	0.00 - 0.70
BASO#	0.03	10 ³ /uL	0.00 - 0.10
TOTAL RED BLOOD CELL COUNT (RBC)	5.50	x10 ⁶ /uL	4.50 - 5.50
HEMATOCRIT (HCT)	48.20	%	40.00 - 50.00
MEAN CORP VOLUME (MCV)	85.3	fL	83.0 - 101.0
MEAN CORP HB (MCH)	28.0	pg	27.0 - 32.0
MEAN CORP HB CONC (MCHC)	32.9	g/dL	31.5 - 34.5
PLATELET COUNT	183	x10 ³ /uL	150 - 410
RDW-CV	14.0	%	11.6 - 14.0
MENTZER INDEX	15.51		

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)	21 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

(CBC); Methodology: FLC, DLC Fluorescent Flow cytometry, HB SLS method, TRBC, PCV, PLT Hydrodynamically focused Impedance. and

of Connective tissue disease. MCH, MCV, MCHC, MENTZER INDEX are calculated. **Instrument Name:** Sysmex 6 part fully automatic analyzer XN-L, Japan

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NAME :- Mr. KAILASH CHAND SHAU

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Lab/Hosp :-

Company :- MediWheel



Sample Type :- PLAIN/SERUM

Sample Collected Time 14/05/2023 09:30:33

Final Authentication : 14/05/2023 13:27:32

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIPID PROFILE			
TOTAL CHOLESTEROL Method:- Enzymatic Endpoint Method	123.63	mg/dl	Desirable <200 Borderline 200-239 High > 240
TRIGLYCERIDES Method:- GPO-PAP	67.92	mg/dl	Normal <150 Borderline high 150-199 High 200-499 Very high >500
DIRECT HDL CHOLESTEROL Method:- Direct clearance Method	34.12	mg/dl	Low < 40 High > 60
DIRECT LDL CHOLESTEROL Method:- Direct clearance Method	78.19	mg/dl	Optimal <100 Near Optimal/above optimal 100-129 Borderline High 130-159 High 160-189 Very High > 190
VLDL CHOLESTEROL Method:- Calculated	13.58	mg/dl	0.00 - 80.00
T.CHOLESTEROL/HDL CHOLESTEROL RATIO Method:- Calculated	3.62		0.00 - 4.90
LDL / HDL CHOLESTEROL RATIO Method:- Calculated	2.29		0.00 - 3.50
TOTAL LIPID Method:- CALCULATED	366.00	L mg/dl	400.00 - 1000.00

TOTAL CHOLESTEROL **InstrumentName:**Randox Rx Imola **Interpretation:** Cholesterol measurements are used in the diagnosis and treatments of lipid lipoprotein metabolism disorders.

TRIGLYCERIDES **InstrumentName:**Randox Rx Imola **Interpretation :** Triglyceride measurements are used in the diagnosis and treatment of diseases involving lipid metabolism and various endocrine disorders e.g. diabetes mellitus, nephrosis and liver obstruction.

DIRECT HDLCHOLESTEROL **InstrumentName:**Randox 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.

DIRECT LDL-CHOLESTEROL **InstrumentName:**Randox 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.

TOTAL LIPID AND VLDL ARE CALCULATED

MUKESH SINGH

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Date :- 14/05/2023 09:13:54 Patient ID :-1223696
NAME :- Mr. KAILASH CHAND SHAU Ref. By Dr:- BOB
Sex / Age :- Male 52 Yrs 11 Mon 5 Days Lab/Hosp :-
Company :- MediWheel



Sample Type :- PLAIN/SERUM

Sample Collected Time 14/05/2023 09:30:33

Final Authentication : 14/05/2023 13:27:32

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
LIVER PROFILE WITH GGT			
SERUM BILIRUBIN (TOTAL) Method:- Colorimetric method	0.75	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	0.26	mg/dL	Adult - Up to 0.25 Newborn - <0.6 >- 1 month - <0.2
SERUM BILIRUBIN (INDIRECT) Method:- Calculated	0.49	mg/dl	0.30-0.70
SGOT Method:- IFCC	30.3	U/L	Men- Up to - 37.0 Women - Up to - 31.0
SGPT Method:- IFCC	39.3	U/L	Men- Up to - 40.0 Women - Up to - 31.0
SERUM ALKALINE PHOSPHATASE Method:- AMP Buffer	51.90	IU/L	30.00 - 120.00
SERUM GAMMA GT Method:- IFCC	24.10	U/L	11.00 - 50.00
SERUM TOTAL PROTEIN Method:- Biuret Reagent	7.07	g/dl	6.40 - 8.30
SERUM ALBUMIN Method:- Bromocresol Green	4.31	g/dl	3.80 - 5.00
SERUM GLOBULIN Method:- CALCULATION	2.76	gm/dl	2.20 - 3.50
A/G RATIO	1.56		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)

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Sex / Age :- Male 52 Yrs 11 Mon 5 Days

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Sample Type :- PLAIN/SERUM

Sample Collected Time 14/05/2023 09:30:33

Final Authentication : 14/05/2023 14:12:19

IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
TOTAL THYROID PROFILE			
SERUM TOTAL T3 Method:- Chemiluminescence(Competitive immunoassay)	1.263	ng/ml	0.970 - 1.690
SERUM TOTAL T4 Method:- Chemiluminescence(Competitive immunoassay)	8.014	ug/dl	5.530 - 11.000
SERUM TSH ULTRA Method:- Enhanced Chemiluminescence Immunoassay	1.140	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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Sex / Age :- Male 52 Yrs 11 Mon 5 Days

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Company :- MediWheel



Sample Type :- URINE

Sample Collected Time 14/05/2023 09:30:33

Final Authentication : 14/05/2023 16:57:06

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) Method:- Reagent Strip(Double indicator blue reaction)	6.5		5.0 - 7.5
SPECIFIC GRAVITY Method:- Reagent Strip(bromthymol blue)	1.020		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
MICROSCOPY EXAMINATION			
RBC/HPF	NIL	/HPF	NIL
WBC/HPF	2-3	/HPF	2-3
EPITHELIAL CELLS	0-1	/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

RAJKUMAR
Technologist

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Tele: 0141-2293346, 4049787, 9887049787
Website: www.drgoyalpathlab.com | E-mail: drgoyalpiyush@gmail.com

Date :- 14/05/2023 09:13:54 Patient ID :-1223696
NAME :- Mr. KAILASH CHAND SHAU Ref. By Dr:- BOB
Sex / Age :- Male 52 Yrs 11 Mon 5 Days Lab/Hosp :-
Company :- MediWheel



Sample Type :- STOOL Sample Collected Time 14/05/2023 09:30:33 Final Authentication : 14/05/2023 16:57:06

CLINICAL PATHOLOGY

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

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

MC- 5509

Date :- 14/05/2023 09:13:54

Patient ID :-1223696

NAME :- Mr. KAILASH CHAND SHAU

Ref. By Dr:- BOB

Sex / Age :- Male 52 Yrs 11 Mon 5 Days

Lab/Hosp :-

Company :- MediWheel



Sample Type :- KOx/Na FLUORIDE-F, KOx/Na SODIUM FLUORIDE-F, KOx/Na SODIUM FLUORIDE-F

Final Authentication : 14/05/2023 15:12:29

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
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FASTING BLOOD SUGAR (Plasma)

106.3

mg/dl

75.0 - 115.0

Method:- GOD PAP

Impaired glucose tolerance (IGT)

111 - 125 mg/dL

Diabetes Mellitus (DM)

> 126 mg/dL

Instrument Name: Randox Rx Imola **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)

114.0

mg/dl

70.0 - 140.0

Method:- GOD PAP

Instrument Name: Randox Rx Imola **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 .

SERUM CREATININE

1.02

mg/dl

Men - 0.6-1.30

Women - 0.5-1.20

Method:- Colorimetric Method

SERUM URIC ACID

7.41 H

mg/dl

Men - 3.4-7.0

Women - 2.4-5.7

Method:- Enzymatic colorimetric

MUKESH SINGH

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Date :- 14/05/2023 09:13:54

Patient ID :-1223696

NAME :- Mr. KAILASH CHAND SHAU

Ref. By Dr:- BOB

Sex / Age :- Male 52 Yrs 11 Mon 5 Days

Lab/Hosp :-

Company :- MediWheel



HAEMATOLOGY

Test Name	Value	Unit	Biological Ref Interval
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AJAYKUMAR, ANITASHARMA, BANWARI, BILAL, MUKESH SINGH, RAJKUMAR



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Date :- 14/05/2023 09:13:54 Patient ID :-1223696
NAME :- Mr. KAILASH CHAND SHAU Ref. By Dr:- BOB
Sex / Age :- Male 52 Yrs 11 Mon 5 Days Lab/Hosp :-
Company :- MediWheel



Sample Type :- EDTA, URINE, URINE-PP Sample Collected Time 14/05/2023 09:30:33 Final Authentication : 14/05/2023 16:57:06

HAEMATOLOGY

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

BANWARI, RAJKUMAR
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Date :- 14/05/2023 09:13:54

Patient ID :-1223696

NAME :- Mr. KAILASH CHAND SHAU

Ref. By Dr:- BOB

Sex / Age :- Male 52 Yrs 11 Mon 5 Days

Lab/Hosp :-

Company :- MediWheel



Sample Type :- PLAIN/SERUM

Sample Collected Time 14/05/2023 09:30:33

Final Authentication : 14/05/2023 13:27:32

BIOCHEMISTRY

Test Name	Value	Unit	Biological Ref Interval
BLOOD UREA NITROGEN (BUN)	16.2	mg/dl	0.0 - 23.0

MUKESH SINGH

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NAME :- Mr. KAILASH CHAND SHAU Ref. By Dr:- BOB
Sex / Age :- Male 52 Yrs 11 Mon 5 Days Lab/Hosp :-
Company :- MediWheel



Sample Type :- PLAIN/SERUM Sample Collected Time 14/05/2023 09:30:33 Final Authentication : 14/05/2023 14:12:19

IMMUNOASSAY

Test Name	Value	Unit	Biological Ref Interval
TOTAL PSA Method:- Chemiluminescence	0.882	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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