

Patient Name: Mr. Nitish PatraAge/Gender: 31 Y/M

 UHID/MR No.
 : CINR.0000163734
 OP Visit No
 : CINROPV220895

 Sample Collected on
 : 03-03-2024 12:24

Ref Doctor : SELF

# DEPARTMENT OF RADIOLOGY

### **ULTRASOUND - WHOLE ABDOMEN**

LIVER: Appears normal in size, shape and echopattern. No focal parenchymal lesions identified. No evidence of intra/extrahepatic biliary tree dilatation noted. Portal vein appears to be of normal size.

GALLBLADDER: Moderately distended.

: 9439451621

Emp/Auth/TPA ID

SPLEEN: Appears normal in size, shape and echopattern. No focal parenchymal lesions identified.

PANCREAS: Obscured by bowel gas. However, the visualized portion appear normal.

KIDNEYS: Both kidneys appear normal in size, shape and echopattern. Corticomedullary differentiation appears maintained. No evidence of calculi or hydronephrosis on either side.

Right kidney measures 9.8x4.7 cm.

Left kidney measures 8.9x5.5 cm.

URINARY BLADDER: Distended and appears normal. No evidence of abnormal wall thickening noted.

PROSTATE: Prostate is normal in size and echo-pattern.

No free fluid is seen.

**IMPRESSION:** 

NO SIGNIFICANT SONOGRAPHIC ABNORMALITY DETECTED.

Dr. RAMESH G
MBBS DMRD
RADIOLOGY



**Patient Name** : Mr. Nitish Patra Age/Gender : 31 Y/M

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: CINR.0000163734

**OP Visit No** Reported on : CINROPV220895

Sample Collected on

: RAD2255434

Specimen

: 03-03-2024 12:23

**Ref Doctor** 

LRN#

: SELF

Emp/Auth/TPA ID

: 9439451621

### DEPARTMENT OF RADIOLOGY

### X-RAY CHEST PA

Both lung fields and hila are normal.

No obvious active pleuro-parenchymal lesion seen.

Both costophrenic and cardiophrenic angles are clear.

Both diaphragms are normal in position and contour.

Thoracic wall and soft tissues appear normal.

# **CONCLUSION:**

No obvious abnormality seen

Dr. RAMESH G MBBS DMRD RADIOLOGY







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: ARCOFEMI HEALTHCARE LIMITED

### **DEPARTMENT OF HAEMATOLOGY**

### PERIPHERAL SMEAR, WHOLE BLOOD EDTA

RBC PREDOMINANTLY NORMOCYTIC NORMOCHROMIC.MILD MICROCYTOSIS NOTED. WBC WITHIN NORMAL LIMITS

PLATELETS ARE ADEQUATE ON SMEAR NO HEMOPARASITES SEEN

IMPRESSION: NORMOCYTIC NORMOCHROMIC BLOOD PICTURE

Page 1 of 15

DR.Aditi Parkhe MBBS,MD(PATHOLOGY) Consultant Pathologist Dr Priya Murthy M.B.B.S,M.D(Pathology) Consultant Pathologist

SIN No:BED240056680

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### **DEPARTMENT OF HAEMATOLOGY**

### ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

Test Name	Result	Unit	Bio. Ref. Range	Method
HEMOGRAM , WHOLE BLOOD EDTA				
HAEMOGLOBIN	13	g/dL	13-17	Spectrophotometer
PCV	39.40	%	40-50	Electronic pulse & Calculation
RBC COUNT	5.47	Million/cu.mm	4.5-5.5	Electrical Impedence
MCV	72	fL	83-101	Calculated
MCH	23.7	pg	27-32	Calculated
MCHC	32.9	g/dL	31.5-34.5	Calculated
R.D.W	14.9	%	11.6-14	Calculated
TOTAL LEUCOCYTE COUNT (TLC)	5,680	cells/cu.mm	4000-10000	Electrical Impedance
DIFFERENTIAL LEUCOCYTIC COUNT (	DLC)			
NEUTROPHILS	64.7	%	40-80	Electrical Impedance
LYMPHOCYTES	26.3	%	20-40	Electrical Impedance
EOSINOPHILS	1.6	%	1-6	Electrical Impedance
MONOCYTES	7.1	%	2-10	Electrical Impedance
BASOPHILS	0.3	%	<1-2	Electrical Impedance
ABSOLUTE LEUCOCYTE COUNT				
NEUTROPHILS	3674.96	Cells/cu.mm	2000-7000	Calculated
LYMPHOCYTES	1493.84	Cells/cu.mm	1000-3000	Calculated
EOSINOPHILS	90.88	Cells/cu.mm	20-500	Calculated
MONOCYTES	403.28	Cells/cu.mm	200-1000	Calculated
BASOPHILS	17.04	Cells/cu.mm	0-100	Calculated
Neutrophil lymphocyte ratio (NLR)	2.46		0.78- 3.53	Calculated
PLATELET COUNT	249000	cells/cu.mm	150000-410000	Electrical impedence
ERYTHROCYTE SEDIMENTATION RATE (ESR)	35	mm at the end of 1 hour	0-15	Modified Westegren method
PERIPHERAL SMEAR				

RBC PREDOMINANTLY NORMOCYTIC NORMOCHROMIC.MILD MICROCYTOSIS NOTED. WBC WITHIN NORMAL LIMITS

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MBBS,MD(PATHOLOGY)
Consultant Pathologist

Dr Priya Murthy M.B.B.S,M.D(Pathology) Consultant Pathologist

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### ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

PLATELETS ARE ADEQUATE ON SMEAR NO HEMOPARASITES SEEN

IMPRESSION: NORMOCYTIC NORMOCHROMIC BLOOD PICTURE

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Test Name	Result	Unit	Bio. Ref. Range	Method
BLOOD GROUP ABO AND RH FACT	OR , WHOLE BLOOD EDTA			
BLOOD GROUP TYPE	В			Microplate Hemagglutination
Rh TYPE	Positive			Microplate Hemagglutination

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### **DEPARTMENT OF BIOCHEMISTRY**

### ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

Test Name	Result	Unit	Bio. Ref. Range	Method
GLUCOSE, FASTING , NAF PLASMA	81	mg/dL	70-100	HEXOKINASE

## **Comment:**

As per American Diabetes Guidelines, 2023

Fasting Glucose Values in mg/dL	Interpretation	
70-100 mg/dL	Normal	
100-125 mg/dL	Prediabetes	
≥126 mg/dL	Diabetes	
<70 mg/dL	Hypoglycemia	

#### Note:

- 1.The diagnosis of Diabetes requires a fasting plasma glucose of > or = 126 mg/dL and/or a random / 2 hr post glucose value of > or = 200 mg/dL on at least 2 occasions
- 2. Very high glucose levels (>450 mg/dL in adults) may result in Diabetic Ketoacidosis & is considered critical.

Test Name	Result	Unit	Bio. Ref. Range	Method
HBA1C (GLYCATED HEMOGLOBIN) , V	VHOLE BLOOD EDTA			
HBA1C, GLYCATED HEMOGLOBIN	5.2	%		HPLC
ESTIMATED AVERAGE GLUCOSE (eAG)	103	mg/dL		Calculated

### **Comment:**

Reference Range as per American Diabetes Association (ADA) 2023 Guidelines:

REFERENCE GROUP	HBA1C %	
NON DIABETIC	<5.7	
PREDIABETES	5.7 – 6.4	
DIABETES	≥ 6.5	
DIABETICS		
EXCELLENT CONTROL	6 – 7	
FAIR TO GOOD CONTROL	7 – 8	
UNSATISFACTORY CONTROL	8 – 10	

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DR.SHIVARAJA SHETTY
M.B.B.S,M.D(Biochemistry)
CONSULTANT BIOCHEMIST

SIN No:EDT240025583

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### ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

POOR CONTROL

>10

**Note:** Dietary preparation or fasting is not required.

- 1. HbA1C is recommended by American Diabetes Association for Diagnosing Diabetes and monitoring Glycemic Control by American Diabetes Association guidelines 2023.
- 2. Trends in HbA1C values is a better indicator of Glycemic control than a single test.
- 3. Low HbA1C in Non-Diabetic patients are associated with Anemia (Iron Deficiency/Hemolytic), Liver Disorders, Chronic Kidney Disease. Clinical Correlation is advised in interpretation of low Values.
- 4. Falsely low HbA1c (below 4%) may be observed in patients with clinical conditions that shorten erythrocyte life span or decrease mean erythrocyte age. HbA1c may not accurately reflect glycemic control when clinical conditions that affect erythrocyte survival are present.
- 5. In cases of Interference of Hemoglobin variants in HbA1C, alternative methods (Fructosamine) estimation is recommended for Glycemic Control
  - A: HbF >25%
  - B: Homozygous Hemoglobinopathy.

(Hb Electrophoresis is recommended method for detection of Hemoglobinopathy)

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### ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

Test Name	Result	Unit	Bio. Ref. Range	Method
LIPID PROFILE , SERUM				
TOTAL CHOLESTEROL	180	mg/dL	<200	CHO-POD
TRIGLYCERIDES	71	mg/dL	<150	GPO-POD
HDL CHOLESTEROL	52	mg/dL	40-60	Enzymatic Immunoinhibition
NON-HDL CHOLESTEROL	128	mg/dL	<130	Calculated
LDL CHOLESTEROL	113.9	mg/dL	<100	Calculated
VLDL CHOLESTEROL	14.2	mg/dL	<30	Calculated
CHOL / HDL RATIO	3.46		0-4.97	Calculated

### **Comment:**

Reference Interval as per National Cholesterol Education Program (NCEP) Adult Treatment Panel III Report.

	Desirable	Borderline High	High	Very High
TOTAL CHOLESTEROL	< 200	200 - 239	≥ 240	
TRIGLYCERIDES	<150	150 - 199	200 - 499	≥ 500
LDL	Optimal < 100 Near Optimal 100-129	130 - 159	160 - 189	≥ 190
HDL	≥ 60			
NON-HDL CHOLESTEROL	Optimal <130; Above Optimal 130-159	160-189	190-219	>220

- 1. Measurements in the same patient on different days can show physiological and analytical variations.
- 2. NCEP ATP III identifies non-HDL cholesterol as a secondary target of therapy in persons with high triglycerides.
- 3. Primary prevention algorithm now includes absolute risk estimation and lower LDL Cholesterol target levels to determine eligibility of drug therapy.
- **4.** 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 from peripheral tissues.
- 5. 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.
- 6. VLDL, LDL Cholesterol Non HDL Cholesterol, CHOL/HDL RATIO, LDL/HDL RATIO are calculated parameters when Triglycerides are below 400 mg/dL. When Triglycerides are more than 400 mg/dL LDL cholesterol is a direct measurement.

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DR.SHIVARAJA SHETTY
M.B.B.S,M.D(Biochemistry)
CONSULTANT BIOCHEMIST

SIN No:SE04649048

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### **DEPARTMENT OF BIOCHEMISTRY**

ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

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## ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

Test Name	Result	Unit	Bio. Ref. Range	Method
IVER FUNCTION TEST (LFT) , SERUM				
BILIRUBIN, TOTAL	0.95	mg/dL	0.3–1.2	DPD
BILIRUBIN CONJUGATED (DIRECT)	0.18	mg/dL	<0.2	DPD
BILIRUBIN (INDIRECT)	0.77	mg/dL	0.0-1.1	Dual Wavelength
ALANINE AMINOTRANSFERASE (ALT/SGPT)	25	U/L	<50	IFCC
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	27.0	U/L	<50	IFCC
ALKALINE PHOSPHATASE	98.00	U/L	30-120	IFCC
PROTEIN, TOTAL	7.59	g/dL	6.6-8.3	Biuret
ALBUMIN	4.57	g/dL	3.5-5.2	BROMO CRESOL GREEN
GLOBULIN	3.02	g/dL	2.0-3.5	Calculated
A/G RATIO	1.51		0.9-2.0	Calculated

### **Comment:**

LFT results reflect different aspects of the health of the liver, i.e., hepatocyte integrity (AST & ALT), synthesis and secretion of bile (Bilirubin, ALP), cholestasis (ALP, GGT), protein synthesis (Albumin)

Common patterns seen:

#### 1. Hepatocellular Injury:

- AST Elevated levels can be seen. However, it is not specific to liver and can be raised in cardiac and skeletal injuries.
- ALT Elevated levels indicate hepatocellular damage. It is considered to be most specific lab test for hepatocellular injury. Values also correlate well with increasing BMI .• Disproportionate increase in AST, ALT compared with ALP. Bilirubin may be elevated.
- AST: ALT (ratio) In case of hepatocellular injury AST: ALT > 1In Alcoholic Liver Disease AST: ALT usually >2. This ratio is also seen to be increased in NAFLD, Wilsons's diseases, Cirrhosis, but the increase is usually not >2.
- 2. Cholestatic Pattern:
- ALP Disproportionate increase in ALP compared with AST, ALT.
- Bilirubin may be elevated. ALP elevation also seen in pregnancy, impacted by age and sex.
- To establish the hepatic origin correlation with GGT helps. If GGT elevated indicates hepatic cause of increased ALP.
- 3. Synthetic function impairment: Albumin- Liver disease reduces albumin levels. Correlation with PT (Prothrombin Time) helps.

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Test Name	Result	Unit	Bio. Ref. Range	Method
RENAL PROFILE/KIDNEY FUNCTION	TEST (RFT/KFT), SEF	RUM		
CREATININE	0.97	mg/dL	0.67-1.17	Jaffe's, Method
UREA	17.80	mg/dL	17-43	GLDH, Kinetic Assay
BLOOD UREA NITROGEN	8.3	mg/dL	8.0 - 23.0	Calculated
URIC ACID	7.48	mg/dL	3.5–7.2	Uricase PAP
CALCIUM	10.00	mg/dL	8.8-10.6	Arsenazo III
PHOSPHORUS, INORGANIC	3.14	mg/dL	2.5-4.5	Phosphomolybdate Complex
SODIUM	138	mmol/L	136–146	ISE (Indirect)
POTASSIUM	4.9	mmol/L	3.5–5.1	ISE (Indirect)
CHLORIDE	107	mmol/L	101–109	ISE (Indirect)
PROTEIN, TOTAL	7.59	g/dL	6.6-8.3	Biuret
ALBUMIN	4.57	g/dL	3.5-5.2	BROMO CRESOL GREEN
GLOBULIN	3.02	g/dL	2.0-3.5	Calculated
A/G RATIO	1.51		0.9-2.0	Calculated

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Test Name	Result	Unit	Bio. Ref. Range	Method
GAMMA GLUTAMYL	21.00	U/L	<55	IFCC
TRANSPEPTIDASE (GGT), SERUM				

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CONSULTANT BIOCHEMIST

SIN No:SE04649048

This test has been performed at Apollo Health & Lifestyle Ltd, RRL BANGALORE Laboratory

THIS TEST HAS BEEN PERFORMED AT APOLLO HEALTH AND LIFSTYLE LIMITED- RRL BANGALORE

Apollo Health and Lifestyle Limited (CIN-U85110TG2000PLC115819)

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APOLLO CLINICS NETWORK









: Mr.NITISH PATRA

Age/Gender

: 31 Y 0 M 8 D/M

UHID/MR No

: CINR.0000163734

Visit ID

: CINROPV220895

Ref Doctor Emp/Auth/TPA ID

: Dr.SELF : 9439451621 Collected

: 03/Mar/2024 09:34AM

Received

: 03/Mar/2024 01:52PM

Reported

Status

: 03/Mar/2024 03:57PM

: Final Report

Sponsor Name

: ARCOFEMI HEALTHCARE LIMITED

### **DEPARTMENT OF IMMUNOLOGY**

### ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

Test Name	Result	Unit	Bio. Ref. Range	Method
THYROID PROFILE TOTAL (T3, T4, TSH)	, SERUM			
TRI-IODOTHYRONINE (T3, TOTAL)	0.8	ng/mL	0.7-2.04	CLIA
THYROXINE (T4, TOTAL)	10.10	μg/dL	5.48-14.28	CLIA
THYROID STIMULATING HORMONE (TSH)	2.978	μIU/mL	0.34-5.60	CLIA

### **Comment:**

For pregnant females	Bio Ref Range for TSH in uIU/ml (As per American Thyroid Association)				
First trimester	0.1 - 2.5				
Second trimester	0.2 - 3.0				
Third trimester	0.3 – 3.0				

- 1. TSH is a glycoprotein hormone secreted by the anterior pituitary. TSH activates production of T3 (Triiodothyronine) and its prohormone T4 (Thyroxine). Increased blood level of T3 and T4 inhibit production of TSH.
- 2. TSH is elevated in primary hypothyroidism and will be low in primary hyperthyroidism. Elevated or low TSH in the context of normal free thyroxine is often referred to as sub-clinical hypo- or hyperthyroidism respectively.
- 3. Both T4 & T3 provides limited clinical information as both are highly bound to proteins in circulation and reflects mostly inactive hormone. Only a very small fraction of circulating hormone is free and biologically active.
- 4. Significant variations in TSH can occur with circadian rhythm, hormonal status, stress, sleep deprivation, medication & circulating antibodies.

TSH	Т3	T4	FT4	Conditions			
High	Low	Low	Low	Primary Hypothyroidism, Post Thyroidectomy, Chronic Autoimmune Thyroiditis			
High	N	N	N	subclinical Hypothyroidism, Autoimmune Thyroiditis, Insufficient Hormone Replacement Cherapy.			
N/Low	Low	Low	Low	econdary and Tertiary Hypothyroidism			
Low	High	High	High	Primary Hyperthyroidism, Goitre, Thyroiditis, Drug effects, Early Pregnancy			
Low	N	N	N	Subclinical Hyperthyroidism			
Low	Low	Low	Low	Central Hypothyroidism, Treatment with Hyperthyroidism			
Low	N	High	High	Thyroiditis, Interfering Antibodies			
N/Low	High	N	N	T3 Thyrotoxicosis, Non thyroidal causes			
High	High	High	High	Pituitary Adenoma; TSHoma/Thyrotropinoma			

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: Mr.NITISH PATRA

Age/Gender

: 31 Y 0 M 8 D/M

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: CINR.0000163734

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Reported Status : 03/Mar/2024 03:57PM : Final Report

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### **DEPARTMENT OF IMMUNOLOGY**

ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

Page 13 of 15



DR.SHIVARAJA SHETTY
M.B.B.S,M.D(Biochemistry)
CONSULTANT BIOCHEMIST

SIN No:SPL24037475

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APOLLO CLINICS NETWORK









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: CINR.0000163734

Visit ID

: CINROPV220895

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: 03/Mar/2024 09:34AM

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: 03/Mar/2024 02:00PM

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: 03/Mar/2024 03:22PM

Status

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Sponsor Name

: ARCOFEMI HEALTHCARE LIMITED

## **DEPARTMENT OF CLINICAL PATHOLOGY**

### ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

Test Name	Result	Unit	Bio. Ref. Range	Method
COMPLETE URINE EXAMINATION (	CUE) , URINE			
PHYSICAL EXAMINATION				
COLOUR	PALE YELLOW		PALE YELLOW	Visual
TRANSPARENCY	CLEAR		CLEAR	Visual
рН	6.5		5-7.5	DOUBLE INDICATOR
SP. GRAVITY	1.010		1.002-1.030	Bromothymol Blue
BIOCHEMICAL EXAMINATION				
URINE PROTEIN	NEGATIVE		NEGATIVE	PROTEIN ERROR OF INDICATOR
GLUCOSE	NEGATIVE		NEGATIVE	GLUCOSE OXIDASE
URINE BILIRUBIN	NEGATIVE		NEGATIVE	AZO COUPLING REACTION
URINE KETONES (RANDOM)	NEGATIVE		NEGATIVE	SODIUM NITRO PRUSSIDE
UROBILINOGEN	NORMAL		NORMAL	MODIFED EHRLICH REACTION
BLOOD	NEGATIVE		NEGATIVE	Peroxidase
NITRITE	NEGATIVE		NEGATIVE	Diazotization
LEUCOCYTE ESTERASE	NEGATIVE		NEGATIVE	LEUCOCYTE ESTERASE
CENTRIFUGED SEDIMENT WET M	OUNT AND MICROSCOPY	1		
PUS CELLS	2-3	/hpf	0-5	Microscopy
EPITHELIAL CELLS	1-2	/hpf	<10	MICROSCOPY
RBC	NIL	/hpf	0-2	MICROSCOPY
CASTS	NIL		0-2 Hyaline Cast	MICROSCOPY
CRYSTALS	ABSENT		ABSENT	MICROSCOPY

DR.Aditi Parkhe MBBS, MD(PATHOLOGY) Consultant Pathologist

Dr Priya Murthy M.B.B.S, M.D (Pathology) Consultant Pathologist

SIN No:UR2296580

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### **DEPARTMENT OF CLINICAL PATHOLOGY**

### ARCOFEMI - MEDIWHEEL - FULL BODY ANNUAL PLUS MALE - 2D ECHO - PAN INDIA - FY2324

Test Name	Result	Unit	Bio. Ref. Range	Method
URINE GLUCOSE(FASTING)	NEGATIVE		NEGATIVE	Dipstick

\*\*\* End Of Report \*\*\*

Result/s to Follow:

GLUCOSE (POST PRANDIAL) - URINE, GLUCOSE, POST PRANDIAL (PP), 2 HOURS (POST MEAL)

DR.Aditi Parkhe MBBS, MD(PATHOLOGY) Consultant Pathologist

M.B.B.S, M.D (Pathology) Consultant Pathologist

SIN No:UF010877

This test has been performed at Apollo Health & Lifestyle Ltd, RRL BANGALORE Laboratory

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