



CLIENT CODE: CA00010147 - MEDIWHEEL
CLIENT'S NAME AND ADDRESSY THOUSE HANTED

MEDIWHEEL ARCOFEMI HEALTHCARE LIMITED F701A, LADO SARAI, NEW DELHI, SOUTH DELHI, DELHI, SOUTH DELHI 110030 DELHI INDIA Cert. No. MC-2809

DDRC SRL DIAGNOSTICS

GANDHI NAGAR, KTM KERALA, INDIA Tel: 93334 93334

Email: customercare.ddrc@srl.in

PATIENT NAME: REMYA PATIENT ID: REMYF1002794036

ACCESSION NO: 4036WB001799 AGE: 44 Years SEX: Female ABHA NO:

DRAWN: RECEIVED: 10/02/2023 08:14 REPORTED: 10/02/2023 17:16

REFERRING DOCTOR: DR. MEDIWHEEL CLIENT PATIENT ID:

Test Report Status Final Results Biological Reference Interval Units

# **MEDIWHEEL HEALTH CHECKUP ABOVE 40(F)TMT**

\* TREADMILL TEST

TREADMILL TEST COMPLETED

**DENTAL CHECK UP** 

DENTAL CHECK UP COMPLETED

**OPTHAL** 

8800465156

**OPTHAL** COMPLETED

\* PHYSICAL EXAMINATION

PHYSICAL EXAMINATION COMPLETED









Units

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<u>Final</u>

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**Test Report Status** 



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MEDIWHEEL HEALTH CHECKUP ABOVE 40(F)TM	ш			
BLOOD UREA NITROGEN (BUN), SERUM				
BLOOD UREA NITROGEN * BUN/CREAT RATIO	5		Adult(<60 yrs) : 6 to 20	mg/dL
BUN/CREAT RATIO CREATININE, SERUM	8.19		5 - 15	
CREATININE GLUCOSE, POST-PRANDIAL, PLASMA	0.61		18 - 60 yrs : 0.6 - 1.1	mg/dL
GLUCOSE, POST-PRANDIAL, PLASMA	130		Diabetes Mellitus : > or = 200. Impaired Glucose tolerance/ Prediabetes : 140 - 199. Hypoglycemia : < 55.	mg/dL
LIPID PROFILE, SERUM			71 37	
CHOLESTEROL	227		Desirable : < 200 Borderline : 200-239 High : >or= 240	mg/dL
TRIGLYCERIDES	138		Normal: < 150 High: 150-199 Hypertriglyceridemia: 200-499 Very High: > 499	mg/dL
HDL CHOLESTEROL	55		General range : 40-60	mg/dL
DIRECT LDL CHOLESTEROL	169	High	Optimum : < 100 Above Optimum : 100-139 Borderline High : 130-159 High : 160-189 Very High : > or = 190	mg/dL
NON HDL CHOLESTEROL	172	High	Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL
VERY LOW DENSITY LIPOPROTEIN	27.6		< or = 30.0	mg/dL
CHOL/HDL RATIO	4.1		3.30 - 4.40	
LDL/HDL RATIO	3.1	High	0.5 - 3.0	









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#### Interpretation(s)

- 1) Cholesterol levels help assess the patient risk status and to follow the progress of patient under treatment to lower serum cholesterol concentrations.
- 2) Serum Triglyceride (TG) are a type of fat and a major source of energy for the body. Both quantity and composition of the diet impact on plasma triglyceride concentrations. Elevations in TG levels are the result of overproduction and impaired clearance. High TG are associated with increased risk for CAD (Coronary artery disease) in patients with other risk factors, such as low HDL-C, some patient groups with elevated apolipoprotein B concentrations, and patients with forms of LDL that may be particularly atherogenic.
- 3)HDL-C plays a crucial role in the initial step of reverse cholesterol transport, this considered to be the primary atheroprotective function of HDL.
- 4) LDL -C plays a key role in causing and influencing the progression of atherosclerosis and, in particular, coronary sclerosis. The majority of cholesterol stored in atherosclerotic plaques originates from LDL, thus LDL-C value is the most powerful clinical predictor.
- 5)Non HDL cholesterol: Non-HDL-C measures the cholesterol content of all atherogenic lipoproteins, including LDL hence it is a better marker of risk in both primary and secondary prevention studies. Non-HDL-C also covers, to some extent, the excess ASCVD risk imparted by the sdLDL, which is significantly more atherogenic than the normal large buoyant particles, an elevated non-HDL-C indirectly suggests greater proportion of the small, dense variety of LDL particles

Serum lipid profile is measured for cardiovascular risk prediction. Lipid Association of India recommends LDL-C as primary target and Non HDL-C as co-primary treatment target.

## Risk Stratification for ASCVD (Atherosclerotic cardiovascular disease) by Lipid Association of India

A.CAD with > 1 feature of high risk group						
B. CAD with > 1 feature of Very high risk g	group or recurrent ACS (within 1 year) despite LDL-C					
< or = 50 mg/dl or polyvascular disease	< or = 50 mg/dl or polyvascular disease					
	major risk factors or evidence of end organ damage 3.					
Familial Homozygous Hypercholesterolemi	a					
1. Three major ASCVD risk factors. 2. Diabetes with 1 major risk factor or no evidence of end						
organ damage. 3. CKD stage 3B or 4. 4. LDL >190 mg/dl 5. Extreme of a single risk factor. 6.						
Coronary Artery Calcium - CAC >300 AU. 7. Lipoprotein a >/= 50mg/dl 8. Non stenotic carotid						
plaque						
2 major ASCVD risk factors						
0-1 major ASCVD risk factors						
Major ASCVD (Atherosclerotic cardiovascular disease) Risk Factors						
1. Age > or = 45 years in males and > or = 55 years in females  3. Current Cigarette smoking or tobacco use						
2. Family history of premature ASCVD 4. High blood pressure						
	B. CAD with > 1 feature of Very high risk § < or = 50 mg/dl or polyvascular disease  1. Established ASCVD 2. Diabetes with 2 r Familial Homozygous Hypercholesterolemi  1. Three major ASCVD risk factors. 2. Dia organ damage. 3. CKD stage 3B or 4. 4. L Coronary Artery Calcium - CAC > 300 AU. plaque  2 major ASCVD risk factors  0-1 major ASCVD risk factors  erosclerotic cardiovascular disease) Risk Fa s in males and > or = 55 years in females					

Newer treatment goals and statin initiation thresholds based on the risk categories proposed by LAI in 2020.

Risk Group	Treatment Goals	Consider Drug Therapy





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Units

	LDL-C (mg/dl)	Non-HDL (mg/dl)	LDL-C (mg/dl)	Non-HDL (mg/dl)
Extreme Risk Group	<50 (Optional goal	< 80 (Optional goal	>OR = 50	>OR = 80
Category A	$\langle OR = 30 \rangle$	$\langle OR = 60 \rangle$		
Extreme Risk Group	<OR = 30	< OR = 60	> 30	>60
Category B				
Very High Risk	<50	<80	>OR= 50	>OR= 80
High Risk	<70	<100	>OR= 70	>OR= 100
Moderate Risk	<100	<130	>OR= 100	>OR= 130
Low Risk	<100	<130	>OR= 130*	>OR= 160

<sup>\*</sup>After an adequate non-pharmacological intervention for at least 3 months.

**References:** Management of Dyslipidaemia for the Prevention of Stroke: Clinical Practice Recommendations from the Lipid Association of India. Current Vascular Pharmacology, 2022, 20, 134-155.

# GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD

GLYCOSYLATED HEMOGLOBIN (HBA1C)	5.8	Normal	: 4.0 - 5.6%.	%
01.000.1	0.0	Non-diabetic level	: < 5.7%.	

Diabetic : >6.5%

Glycemic control goal

More stringent goal : < 6.5 %. General goal : < 7%. Less stringent goal : < 8%.

Glycemic targets in CKD :- If eGFR > 60 : < 7%. If eGFR < 60 : 7 - 8.5%.

### LIVER FUNCTION TEST WITH GGT

BILIRUBIN, TOTAL	0.31	General Range : < 1.1	mg/dL
BILIRUBIN, DIRECT	0.12	General Range : < 0.3	mg/dL
BILIRUBIN, INDIRECT	0.19	0.00 - 1.00	mg/dL
TOTAL PROTEIN	7.5	Ambulatory: 6.4 - 8.3 Recumbant: 6 - 7.8	g/dL
ALBUMIN	4.4	20-60yrs: 3.5 - 5.2	g/dL
GLOBULIN	3.1	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO	1.4	1.0 - 2.0	RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	24	Adults: < 33	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT)	14	Adults: < 34	U/L





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ALIVALIANE PLICEPLIATACE	70		Adult( <60,000) , 25 105	1171
ALKALINE PHOSPHATASE	72		Adult (60yrs): 35 - 105	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT)	23		Adult (female) : < 40	U/L
TOTAL PROTEIN, SERUM				
TOTAL PROTEIN	7.5		Ambulatory: 6.4 - 8.3	g/dL
			Recumbant : 6 - 7.8	
URIC ACID	F 4		Adulta . 2 4 F 7	
URIC ACID  ABO GROUP & RH TYPE, EDTA WHOLE BLOOD	5.4		Adults: 2.4-5.7	mg/dL
ABO GROUP	TYPE O			
RH TYPF	POSITIVE			
BLOOD COUNTS,EDTA WHOLE BLOOD	POSITIVE			
HEMOGLOBIN	12.7		12.0 - 15.0	g/dL
RED BLOOD CELL COUNT	4.14		3.8 - 4.8	mil/μL
WHITE BLOOD CELL COUNT	8.00		4.0 - 10.0	thou/µL
PLATELET COUNT	411	High	150 - 410	thou/µL
RBC AND PLATELET INDICES				
HEMATOCRIT	38.4		36 - 46	%
MEAN CORPUSCULAR VOL	93.0		83 - 101	fL
MEAN CORPUSCULAR HGB.	30.7		27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN	33.1		31.5 - 34.5	g/dL
CONCENTRATION	10.0		11.6.11.0	0/
RED CELL DISTRIBUTION WIDTH	10.8	Low	11.6 - 14.0	%
MENTZER INDEX WBC DIFFERENTIAL COUNT	22.5			
	FO		40 - 80	%
SEGMENTED NEUTROPHILS LYMPHOCYTES	59 39		20 - 40	%
MONOCYTES	<b>00</b>	Low	2 - 10	%
EOSINOPHILS	02		1 - 6	%
ABSOLUTE NEUTROPHIL COUNT	4.72		2.0 - 7.0	thou/µL
ABSOLUTE LYMPHOCYTE COUNT	3.12	High	1.0 - 3.0	thou/µL
ABSOLUTE MONOCYTE COUNT	0	Low	0.2 - 1.0	thou/µL
ABSOLUTE EOSINOPHIL COUNT	0.16		0.02 - 0.50	thou/µL





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NEUTROPHIL LYMPHOCYTE RATIO (NLR)	) 1.5		
<b>ERYTHROCYTE SEDIMENTATION RATE (ESR) BLOOD</b>	,WHOLE		
SEDIMENTATION RATE (ESR)	17	0 - 20	mm at 1 hr
SUGAR URINE - POST PRANDIAL			
SUGAR URINE - POST PRANDIAL	NOT DETECTED	NOT DETECTED	
THYROID PANEL, SERUM			
T3	116.33	Non-Pregnant: 60-181	ng/dL
	10.10	Pregnant Trimester-wise 1st: 81-190 2nd: 100-260 3rd: 100-260	/-
T4	10.10	3.2 - 12.6	µg/dl
TSH 3RD GENERATION	5.950	(Non Pregnant) : 0.4 - 4.2	μIU/mL
		Pregnant(Trimester wise) 1st : 0.1 - 2.5 2nd : 0.2 - 3	











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#### Interpretation(s)

**Triiodothyronine T3**, **Thyroxine T4**, and **Thyroid Stimulating Hormone TSH** are thyroid hormones which affect almost every physiological process in the body, including growth, development, metabolism, body temperature, and heart rate.

Production of T3 and its prohormone thyroxine (T4) is activated by thyroid-stimulating hormone (TSH), which is released from the pituitary gland. Elevated concentrations of T3, and T4 in the blood inhibit the production of TSH.

Excessive secretion of thyroxine in the body is hyperthyroidism, and deficient secretion is called hypothyroidism.

In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hyperthyroidism, TSH levels are low. Below mentioned are the guidelines for Pregnancy related reference ranges for Total T4, TSH & Total T3. Measurement of the serum TT3 level is a more sensitive test for the diagnosis of hyperthyroidism, and measurement of TT4 is more useful in the diagnosis of hypothyroidism. Most of the thyroid hormone in blood is bound to transport proteins. Only a very small fraction of the circulating hormone is free and biologically active. It is advisable to detect Free T3, FreeT4 along with TSH, instead of testing for albumin bound Total T3, Total T4.

Sr. No.	TSH	Total T4	FT4	Total T3	Possible Conditions
1	High	Low	Low	Low	(1) Primary Hypothyroidism (2) Chronic autoimmune Thyroiditis (3)
					Post Thyroidectomy (4) Post Radio-Iodine treatment
2	High	Normal	Normal	Normal	(1)Subclinical Hypothyroidism (2) Patient with insufficient thyroid
					hormone replacement therapy (3) In cases of Autoimmune/Hashimoto
					thyroiditis (4). Isolated increase in TSH levels can be due to Subclinical
					inflammation, drugs like amphetamines, Iodine containing drug and
					dopamine antagonist e.g. domperidone and other physiological reasons.
3	Normal/Low	Low	Low	Low	(1) Secondary and Tertiary Hypothyroidism
4	Low	High	High	High	(1) Primary Hyperthyroidism (Graves Disease) (2) Multinodular Goitre
					(3)Toxic Nodular Goitre (4) Thyroiditis (5) Over treatment of thyroid
					hormone (6) Drug effect e.g. Glucocorticoids, dopamine, T4
					replacement therapy (7) First trimester of Pregnancy
5	Low	Normal	Normal	Normal	(1) Subclinical Hyperthyroidism
6	High	High	High	High	(1) TSH secreting pituitary adenoma (2) TRH secreting tumor
7	Low	Low	Low	Low	(1) Central Hypothyroidism (2) Euthyroid sick syndrome (3) Recent
					treatment for Hyperthyroidism
8	Normal/Low	Normal	Normal	High	(1) T3 thyrotoxicosis (2) Non-Thyroidal illness
9	Low	High	High	Normal	(1) T4 Ingestion (2) Thyroiditis (3) Interfering Anti TPO antibodies

REF: 1. TIETZ Fundamentals of Clinical chemistry 2.Guidlines of the American Thyroid association during pregnancy and Postpartum, 2011.

NOTE: It is advisable to detect Free T3,FreeT4 along with TSH, instead of testing for albumin bound Total T3, Total T4.TSH is not affected by variation in thyroid - binding protein. TSH has a diurnal rhythm, with peaks at 2:00 - 4:00 a.m. And troughs at 5:00 - 6:00 p.m. With ultradian variations.

**SUGAR URINE - FASTING** 

SUGAR URINE - FASTING NOT DETECTED NOT DETECTED

PHYSICAL EXAMINATION, URINE

COLOR PALE YELLOW APPEARANCE SLIGHTLY HAZY









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* CHEMICAL EXAMIN	IATION, U	RINE			
PH			5.0	4.7 - 7.5	
SPECIFIC GRAVITY	′		1.020	1.003 - 1.035	
PROTEIN			NOT DETECTED	NOT DETECTED	
GLUCOSE			NOT DETECTED	NOT DETECTED	
KETONES			NOT DETECTED	NOT DETECTED	
BLOOD			NOT DETECTED	NOT DETECTED	
BILIRUBIN			NOT DETECTED	NOT DETECTED	
UROBILINOGEN			NORMAL	NORMAL	
NITRITE			NOT DETECTED	NOT DETECTED	
MICROSCOPIC EXAM	INATION,	URINE			
RED BLOOD CELLS	5		1 - 2	NOT DETECTED	/HPF
WBC			8-10	0-5	/HPF
EPITHELIAL CELLS			8-10	0-5	/HPF
CASTS			NOT DETECTED		
CRYSTALS			NOT DETECTED		
BACTERIA			DETECTED	NOT DETECTED	
YEAST			DETECTED	NOT DETECTED	

<b>Comments</b>

NOTE - Kindly correlate clinically.









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### Interpretation(s)

The following table describes the probable conditions, in which the analytes are present in urine

Presence of	Conditions
Proteins	Inflammation or immune illnesses
Pus (White Blood Cells)	Urinary tract infection, urinary tract or kidney stone, tumors or any kind
	of kidney impairment
Glucose	Diabetes or kidney disease
Ketones	Diabetic ketoacidosis (DKA), starvation or thirst
Urobilinogen	Liver disease such as hepatitis or cirrhosis
Blood	Renal or genital disorders/trauma
Bilirubin	Liver disease
Erythrocytes	Urological diseases (e.g. kidney and bladder cancer, urolithiasis), urinary
	tract infection and glomerular diseases
Leukocytes	Urinary tract infection, glomerulonephritis, interstitial nephritis either
	acute or chronic, polycystic kidney disease, urolithiasis, contamination by
	genital secretions
Epithelial cells	Urolithiasis, bladder carcinoma or hydronephrosis, ureteric stents or
	bladder catheters for prolonged periods of time
Granular Casts	Low intratubular pH, high urine osmolality and sodium concentration, interaction with Bence-Jones protein
Hyaline casts	Physical stress, fever, dehydration, acute congestive heart failure, renal
Tryarme casts	diseases
Calcium oxalate	Metabolic stone disease, primary or secondary hyperoxaluria, intravenous
Careram Oxarace	infusion of large doses of vitamin C, the use of vasodilator naftidrofuryl
	oxalate or the gastrointestinal lipase inhibitor orlistat, ingestion of
	ethylene glycol or of star fruit (Averrhoa carambola) or its juice
Uric acid	arthritis
Bacteria	Urinary infectionwhen present in significant numbers & with pus cells.
Trichomonas vaginalis	Vaginitis, cervicitis or salpingitis

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**GLUCOSE FASTING, FLUORIDE PLASMA** 

GLUCOSE, FASTING, PLASMA

Diabetes Mellitus : > or = 126. mg/dL

Impaired fasting Glucose/ Prediabetes: 101 - 125. Hypoglycemia: < 55.



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SEX: Female

# MEDIWHEEL HEALTH CHECKUP ABOVE 40(F)TMT

\* ECG WITH REPORT

**REPORT** 

COMPLETED

\* MAMMOGRAPHY -BOTH

**REPORT** 

COMPLETED

\* USG ABDOMEN AND PELVIS

REPORT

COMPLETED

\* CHEST X-RAY WITH REPORT

**REPORT** 

**COMPLETED** 

\*\*End Of Report\*\*

Please visit www.srlworld.com for related Test Information for this accession TEST MARKED WITH '\*' ARE OUTSIDE THE NABL ACCREDITED SCOPE OF THE LABORATORY.

**PRASEEDA S NAIR BIOCHEMIST** 

DR.KRIPA ELIZABETH JOHN **CONSULTANT PATHOLOGIST** 

K.MEERA BHAI **SENIOR BIOCHEMIST** 



