





CLIENT CODE: CA00010147 - MEDIWHEEL

CLIENT'S NAME AND ADDRESS : MEDIWHEEL ARCOFEMI HEALTHCARE LIMITED

F701A, LADO SARAI, NEW DELHI,

SOUTH DELHI, DELHI, SOUTH DELHI 110030

DELHI INDIA 8800465156 DDRC SRL DIAGNOSTICS

GANDHI NAGAR, KTM KERALA, INDIA Tel: 93334 93334

Email: customercare.ddrc@srl.in

PATIENT NAME: ATUL JOSE PHILIP PATIENT ID: ATULM1401904036

ACCESSION NO: **4036WA002673** AGE: 33 Years SEX: Male ABHA NO:

RECEIVED: 14/01/2023 08:55 14/01/2023 15:04 DRAWN: REPORTED:

REFERRING DOCTOR: DR. MEDIWHEEL CLIENT PATIENT ID:

Test Report Status Results **Biological Reference Interval Units Preliminary**

MEDIWHEEL HEALTH CHEKUP BELOW 40(M)TMT

OPTHAL

COMPLETED OPTHAL

* TREADMILL TEST

COMPLETED TREADMILL TEST

* PHYSICAL EXAMINATION

COMPLETED PHYSICAL EXAMINATION





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		_	_
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MEDIWHEEL HEALTH CHEKUP BELOW 40(M)TMT

* PUN /CDFAT RATIO			
* BUN/CREAT RATIO	0.25	5 - 15	
BUN/CREAT RATIO CREATININE, SERUM	9.25	5 - 15	
CREATININE	0.81	18 - 60 yrs : 0.9 - 1.3	mg/dL
GLUCOSE, POST-PRANDIAL, PLASMA			
GLUCOSE, POST-PRANDIAL, PLASMA	105	Diabetes Mellitus : > or = 200. Impaired Glucose tolerance/ Prediabetes : 140 - 199. Hypoglycemia : < 55.	mg/dL
GLUCOSE FASTING, FLUORIDE PLASMA			
GLUCOSE, FASTING, PLASMA	100	Diabetes Mellitus: > or = 126. Impaired fasting Glucose/ Prediabetes: 101 - 125. Hypoglycemia: < 55.	mg/dL
GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA V BLOOD	/HOLE		
GLYCOSYLATED HEMOGLOBIN (HBA1C)	5.2	Normal : 4.0 - 5.6%. 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%.	
MEAN PLASMA GLUCOSE LIPID PROFILE, SERUM	102.5	< 116.0	mg/dL
CHOLESTEROL	158	Desirable: < 200 Borderline: 200-239 High: >or= 240	mg/dL
TRIGLYCERIDES	55	Normal : < 150 High : 150-199 Hypertriglyceridemia : 200-499 Very High : > 499	mg/dL
HDL CHOLESTEROL	60	General range: 40-60	mg/dL











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DIRECT LDL CHOLESTEROL	105		Optimum : < 100 Above Optimum : 100-139 Borderline High : 130-159 High : 160-189 Very High : >or= 190	mg/dL
NON HDL CHOLESTEROL	98		Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL
CHOL/HDL RATIO	2.6	Low	3.30 - 4.40	
LDL/HDL RATIO	1.8		0.5 - 3.0	
VERY LOW DENSITY LIPOPROTEIN LIVER FUNCTION TEST WITH GGT	11.0		< or = 30.0	mg/dL
BILIRUBIN, TOTAL	1.30		General Range : < 1.1	mg/dL
BILIRUBIN, DIRECT	0.46	High	General Range : < 0.3	mg/dL
BILIRUBIN, INDIRECT	0.84		0.00 - 1.00	mg/dL
TOTAL PROTEIN	7.3		Ambulatory: 6.4 - 8.3 Recumbant: 6 - 7.8	g/dL
ALBUMIN	5.2		20-60yrs: 3.5 - 5.2	g/dL
GLOBULIN	2.1		2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO	2.5	High	1.0 - 2.0	RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	19		Adults: < 40	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT)	19		Adults: < 45	U/L
ALKALINE PHOSPHATASE	64		Adult(<60yrs): 40 - 130	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT) TOTAL PROTEIN, SERUM	13		Adult (male) : < 60	U/L
TOTAL PROTEIN	7.3		Ambulatory: 6.4 - 8.3 Recumbant: 6 - 7.8	g/dL
URIC ACID, SERUM				
URIC ACID ABO GROUP & RH TYPE, EDTA WHOLE BLOOD	5.6		Adults: 3.4-7	mg/dL
ABO GROUP RH TYPE	TYPE O POSITIVE			
BLOOD COUNTS,EDTA WHOLE BLOOD				
HEMOGLOBIN	15.3		13.0 - 17.0	g/dL
RED BLOOD CELL COUNT	5.14		4.5 - 5.5	mil/μL











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Results			Units
5.00		4.0 - 10.0	thou/µL
266		150 - 410	thou/µL
42.7		40 - 50	%
83.0		83 - 101	fL
29.8		27.0 - 32.0	pg
36.0	High	31.5 - 34.5	g/dL
12.1		11.6 - 14.0	%
16.2			
54		40 - 80	%
44	High	20 - 40	%
00	Low	2 - 10	%
02		1 - 6	%
2.7		2.0 - 7.0	thou/µL
2.2		1.0 - 3.0	thou/µL
0	Low	0.2 - 1.0	thou/µL
0.1		0.02 - 0.50	thou/µL
1.2			
HOLE			
03		0 - 14	mm at 1 hr
RESULT PENDING			
112.86		20-50 yrs : 60-181	ng/dL
9.90		3.2 - 12.6	μg/dl
0.750		18-49 yrs : 0.4 - 4.2	μIU/mL
	5.00 266 42.7 83.0 29.8 36.0 12.1 16.2 54 44 00 02 2.7 2.2 0 0.1 1.2 HOLE 03 RESULT PENDING	5.00 266 42.7 83.0 29.8 36.0 High 12.1 16.2 54 44 High 00 Low 02 2.7 2.2 0 Low 0.1 1.2 HOLE 03 RESULT PENDING	5.00









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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 hyporthyroidism, 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.

PHYSICAL EXAMINATION, URINE

COLOR PALE YELLOW

APPEARANCE CLEAR

* CHEMICAL EXAMINATION, URINE

4.8 - 7.4 PH 7.0 1.015 - 1.030 SPECIFIC GRAVITY 1.015











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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		2-3	0-5	/HPF
EPITHELIAL CELLS		NOT DETECTED	NOT DETECTED	/HPF
CASTS		NOT DETECTED		
CRYSTALS		NOT DETECTED		
BACTERIA		NOT DETECTED	NOT DETECTED	
YEAST		NOT DETECTED	NOT DETECTED	
BLOOD UREA NITRO	GEN (BUN), SERUM			
BLOOD UREA NITE	ROGEN	8	Adult(<60 yrs): 6 to 20	mg/dL
SUGAR URINE - FAS	TING			
SUGAR URINE - F	ASTING	NOT DETECTED	NOT DETECTED	
* PHYSICAL EXAMIN	ATION,STOOL	RESULT PENDING		
* CHEMICAL EXAMIN	IATION,STOOL	RESULT PENDING		
* MICROSCOPIC EXA	MINATION,STOOL	RESULT PENDING		

Interpretation(s)

- CREATININE, SERUM-Higher than normal level may be due to:

 Blockage in the urinary tract

 Kidney problems, such as kidney damage or failure, infection, or reduced blood flow
- Loss of body fluid (dehydration)Muscle problems, such as breakdown of muscle fibers
- Problems during pregnancy, such as seizures (eclampsia)), or high blood pressure caused by pregnancy (preeclampsia)

Lower than normal level may be due to:

- Myasthenia GravisMuscular dystrophy

GLUCOSE, POST-PRANDIAL, PLASMA-High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glyosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc.Additional test HbA1c GLUCOSE FASTING, FLUORIDE PLASMA-TEST DESCRIPTION

Normally, the glucose concentration in extracellular fluid is closely regulated so that a source of energy is readily available to tissues and sothat no glucose is excreted in the

Increased in

Diabetes mellitus, Cushing's syndrome (10 - 15%), chronic pancreatitis (30%). Drugs:corticosteroids,phenytoin, estrogen, thiazides.











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Decreased in

Pancreatic islet cell disease with increased insulin,insulinoma,adrenocortical insufficiency, hypopituitarism,diffuse liver disease, malignancy (adrenocortical, stomach,fibrosarcoma), infant of a diabetic mother, enzyme deficiency diseases(e.g., galactosemia),Drugs- insulin, ethanol, propranolol; sulfonylureas,tolbutamide, and other oral hypoglycemic agents

NOTE:

While random serum glucose levels correlate with home glucose monitoring results (weekly mean capillary glucose values), there is wide fluctuation within individuals. Thus,

glycosylated hemoglobin(HbA1c) levels are favored to monitor glycemic control.

High fasting glucose level in comparison to post prandial glucose level may be seen due to effect of Oral Hypoglycaemics & Insulin treatment, Renal Glyosuria, Glycaemic index & response to food consumed, Alimentary Hypoglycemia, Increased insulin response & sensitivity etc. GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD-**Used For**:

- 1. Evaluating the long-term control of blood glucose concentrations in diabetic patients.
- 2.Diagnosing diabetes.
- 3. Identifying patients at increased risk for diabetes (prediabetes).

The ADA recommends measurement of HbAIc (typically 3-4 times per year for type 1 and poorly controlled type 2 diabetic patients, and 2 times per year for well-controlled type 2 diabetic patients) to determine whether a patients metabolic control has remained continuously within the target range.

- 1.eAG (Estimated average glucose) converts percentage HbA1c to md/dl, to compare blood glucose levels.

 2. eAG gives an evaluation of blood glucose levels for the last couple of months.

 3. eAG is calculated as eAG (mg/dl) = 28.7 * HbA1c 46.7

HbA1c Estimation can get affected due to :

I.Shortened Erythrocyte survival: Any condition that shortens erythrocyte survival or decreases mean erythrocyte age (e.g. recovery from acute blood loss,hemolytic anemia) will falsely lower HbA1c test results.Fructosamine is recommended in these patients which indicates diabetes control over 15 days.

II.Vitamin C & E are reported to falsely lower test results.(possibly by inhibiting glycation of hemoglobin.
III.Iron deficiency anemia is reported to increase test results. Hypertriglyceridemia, uremia, hyperbilirubinemia, chronic alcoholism, chronic ingestion of salicylates & opiates addiction are reported to interfere with some assay methods, falsely increasing results. IV.Interference of hemoglobinopathies in HbA1c estimation is seen in

a. Homozygous hemoglobinopathy. Fructosamine is recommended for testing of HbA1c.
b. Heterozygous state detected (D10 is corrected for HbS & HbC trait.)
c. HbF > 25% on alternate paltform (Boronate affinity chromatography) is recommended for testing of HbA1c. Abnormal Hemoglobin electrophoresis (HPLC method) is recommended for detecting a hemoglobinopathy
LIPID PROFILE, SERUM-Serum cholesterol is a blood test that can provide valuable information for the risk of coronary artery disease This test can help determine your risk

Serum Triglyceride are a type of fat in the blood. When you eat, your body converts any calories it doesn'''''''t need into triglycerides, which are stored in fat cells. High triglyceride levels are associated with several factors, including being overweight, eating too many sweets or drinking too much alcohol, smoking, being sedentary, or having diabetes with elevated blood sugar levels. Analysis has proven useful in the diagnosis and treatment of patients with diabetes mellitus, nephrosis, liver obstruction, other diseases involving lipid metabolism, and various endocrine disorders. In conjunction with high density lipoprotein and total serum cholesterol, a triglyceride determination provides valuable information for the assessment of coronary heart disease risk. It is done in fasting state.

High-density lipoprotein (HDL) cholesterol. This is sometimes called the ""good"" cholesterol because it helps carry away LDL cholesterol, thus keeping arteries open and blood flowing more freely. HDL cholesterol is inversely related to the risk for cardiovascular disease. It increases following regular exercise, moderate alcohol consumption and with oral estrogen therapy. Decreased levels are associated with obesity, stress, cigarette smoking and diabetes mellitus.

SERUM LDL The small dense LDL test can be used to determine cardiovascular risk in individuals with metabolic syndrome or established/progressing coronary artery disease, individuals with triglyceride levels between 70 and 140 mg/dL, as well as individuals with a diet high in trans-fat or carbohydrates. Elevated sdLDL levels are associated with metabolic syndrome and an 'atherogenic lipoprotein profile', and are a strong, independent predictor of cardiovascular disease. Elevated levels of LDL arise from multiple sources. A major factor is sedentary lifestyle with a diet high in saturated fat. Insulin-resistance and pre-diabetes have also been implicated, as has genetic predisposition. Measurement of sdLDL allows the clinician to get a more comprehensive picture of lipid risk factors and tailor treatment accordingly. Reducing LDL levels will reduce the risk of CVD and MI.

Non HDL Cholesterol - Adult treatment panel ATP III suggested the addition of Non-HDL Cholesterol as an indicator of all atherogenic lipoproteins (mainly LDL and VLDL). NICE guidelines recommend Non-HDL Cholesterol measurement before initiating lipid lowering therapy. It has also been shown to be a better marker of risk in both primary and secondary prevention studies.

Recommendations:

Results of Lipids should always be interpreted in conjunction with the patient's medical history, clinical presentation and other findings.

NON FASTING LIPID PROFILE includes Total Cholesterol, HDL Cholesterol and calculated non-HDL Cholesterol. It does not include trialycerides and may be best used in

patients for whom fasting is difficult.

TOTAL PROTEIN, SERUM-Serum total protein, also known as total protein, is a biochemical test for measuring the total amount of protein in serum. Protein in the plasma is made up of albumin and globulin

Higher-than-normal levels may be due to: Chronic inflammation or infection, including HIV and hepatitis B or C. Multiple myeloma, Waldenstrom''''''s disease



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Lower-than-normal levels may be due to: Agammaglobulinemia, Bleeding (hemorrhage), Burns, Glomerulonephritis, Liver disease, Malabsorption, Malnutrition, Nephrotic syndrome, Protein-losing enteropathy etc

URIC ACID, SERUM-Causes of Increased levels:-Dietary(High Protein Intake,Prolonged Fasting,Rapid weight loss),Gout,Lesch nyhan syndrome,Type 2 DM,Metabolic

Causes of decreased levels-Low Zinc intake, OCP, Multiple Sclerosis

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD-Blood group is identified by antigens and antibodies present in the blood. Antigens are protein molecules found on the surface of red blood cells. Antibodies are found in plasma. To determine blood group, red cells are mixed with different antibody solutions to give A,B,O or AB.

Disclaimer: "Please note, as the results of previous ABO and Rh group (Blood Group) for pregnant women are not available, please check with the patient records for availability of the same.

The test is performed by both forward as well as reverse grouping methods.

BLOOD COUNTS,EDTA WHOLE BLOOD-The cell morphology is well preserved for 24hrs. However after 24-48 hrs a progressive increase in MCV and HCT is observed leading to a decrease in MCHC. A direct smear is recommended for an accurate differential count and for examination of RBC morphology.

RBC AND PLATELET INDICES-Mentzer index (MCV/RBC) is an automated cell-counter based calculated screen tool to differentiate cases of Iron deficiency anaemia(>13) from Beta thalassaemia trait

(<13) in patients with microcytic anaemia. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HbA2 remains the gold standard for diagnosing a case of beta thalassaemia trait.

WBC DIFFERENTIAL COUNT-The optimal threshold of 3.3 for NLR showed a prognostic possibility of clinical symptoms to change from mild to severe in COVID positive

patients. When age = 49.5 years old and NLR = 3.3, 46.1% COVID-19 patients with mild disease might become severe. By contrast, when age < 49.5 years old and NLR < 3.3, COVID-19 patients tend to show mild disease.

(Reference to - The diagnostic and predictive role of NLR, d-NLR and PLR in COVID-19 patients; A.-P. Yang, et al.; International Immunopharmacology 84 (2020) 106504

This ratio element is a calculated parameter and out of NABL scope. ERYTHROCYTE SEDIMENTATION RATE (ESR), WHOLE BLOOD-TEST DESCRIPTION:

Erythrocyte sedimentation rate (ESR) is a test that indirectly measures the degree of inflammation present in the body. The test actually measures the rate of fall (sedimentation) of erythrocytes in a sample of blood that has been placed into a tall, thin, vertical tube. Results are reported as the millimetres of clear fluid (plasma) that are present at the top portion of the tube after one hour. Nowadays fully automated instruments are available to measure ESR.

ESR is not diagnostic; it is a non-specific test that may be elevated in a number of different conditions. It provides general information about the presence of an inflammatory condition.CRP is superior to ESR because it is more sensitive and reflects a more rapid change **TEST INTERPRETATION**

Increase in: Infections, Vasculities, Inflammatory arthritis, Renal disease, Anemia, Malignancies and plasma cell dyscrasias, Acute allergy Tissue injury, Pregnancy,

Estrogen medication, Aging.
Finding a very accelerated ESR(>100 mm/hour) in patients with ill-defined symptoms directs the physician to search for a systemic disease (Paraproteinemias, Disseminated malignancies, connective tissue disease, severe infections such as bacterial endocarditis).

In pregnancy BRI in first trimester is 0-48 mm/hr(62 if anemic) and in second trimester (0-70 mm /hr(95 if anemic). ESR returns to normal 4th week post partum.

Decreased in: Polycythermia vera, Sickle cell anemia

False elevated ESR: Increased fibrinogen, Drugs(Vitamin A, Dextran etc), Hypercholesterolemia

False Decreased: Poikilocytosis, (SickleCells, spherocytes), Microcytosis, Low fibrinogen, Very high WBC counts, Drugs (Quinine,

1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition; 2. Paediatric reference intervals. AACC Press, 7th edition. Edited by S. Soldin; 3. The reference for the adult reference range is "Practical Haematology by Dacie and Lewis, 10th edition.

BLOOD UREA NITROGEN (BUN), SERUM-Causes of Increased levels include Pre renal (High protein diet, Increased protein catabolism, GI haemorrhage, Cortisol, Dehydration, CHF Renal), Renal Failure, Post Renal (Malignancy, Nephrolithiasis, Prostatism)

Causes of decreased level include Liver disease, SIADH.

SUGAR URINE - FASTING-METHOD: DIPSTICK/BENEDICT'S TEST











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GANDHI NAGAR, KTM KERALA, INDIA Tel: 93334 93334

Email: customercare.ddrc@srl.in

PATIENT NAME: ATUL JOSE PHILIP PATIENT ID: ATULM1401904036

ACCESSION NO: **4036WA002673** AGE: 33 Years SEX: Male ABHA NO:

RECEIVED: 14/01/2023 08:55 14/01/2023 15:04 DRAWN: REPORTED:

REFERRING DOCTOR: DR. MEDIWHEEL CLIENT PATIENT ID:

Test Report Status Results Units **Preliminary**

MEDIWHEEL HEALTH CHEKUP BELOW 40(M)TMT

* ECG WITH REPORT

RFPORT

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**



Scan to View Report

Page 9 Of 9



MEDICAL EXAMINATION REPORT (MER)

If the examinee is suffering from an acute life threatening situation, you may be obliged to disclose the result of the medical examination to the examinee.

ATUL JOSE PHILIP 1 Name of the examinee Mr./Mrs./Ms. (Mole/Scar/any other (specify location)): Male on right-hand small fingel 2. Mark of Identification 33: 10 03 1989 3. Age/Date of Birth Gender: (Passport/Election Card/PAN Card/Driving Licence/Company ID) 4. Photo ID Checked

PHYSICAL DETAILS:

a. Height1.6.8 (cms)	b. Weight	6.3 (Kgs)	c. Girth of A	bdomen
d. Pulse Rate 5.7 (/Min)	e. Blood Press	sure: 110 80	Systolic	Diastolic
		1 st Reading	110	80
	antimosa free and	2 nd Reading	110	80

FAMILY HISTORY:

Relation	Age if Living	Health Status	If deceased, age at the time and cause
Father	68	hood	
Mother	68	bood	
Brother(s)		•	
Sister(s)		Memory of the amployment	se you plank he/she to blef 1957 if a visit on the

HABITS & ADDICTIONS: Does the examinee consume any of the following? - NA

Tobacco in any form	Sedative	Alcohol
his her identity and the findings stated.	-JA -	and the same of th

PERSONAL HISTORY

- a. Are you presently in good health and entirely free from any mental or Physical impairment or deformity. If No, please attach details. AMAN AP
- b. Have you undergone/been advised any surgical procedure? No. 77615
- c. During the last 5 years have you been medically examined, received any advice or treatment or admitted to any hospital? X/X
- d. Have you lost or gained weight in past 12 months?

Have you ever suffered from any of the following?

- Psychological Disorders or any kind of disorders of the Nervous System?
- · Any disorders of Respiratory system?
- Any Cardiac or Circulatory Disorders?
- Enlarged glands or any form of Cancer/Tumour?
- · Any Musculoskeletal disorder?

- Any disorder of Gastrointestinal System?
- · Unexplained recurrent or persistent fever, and/or weight loss
- Have you been tested for HIV/HBsAg / HCV before? If yes attach reports
- Are you presently taking medication f any kind?

GANDHINAGAR KOTTAYAN

686008

DDRC SRL Diagnostics Private I

Corp. Office: DDRC SRL Tower, G- 131, Panampilly Nagar, Ernakulari 682 036 Ph No. 0484-2318223, 2318222, e-mail: info@ddrcsrl.com, web: www.ddrcsrl.com

Corp. Office: DDRC SRL Tower, G- 131, Panampilly Nagar, Ernakulam - 682 036, Ph No: 2310688, 231822, web: www.ddrcsrl.com

XIN

Y/N

YN

XIN

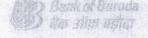
			ļ	
• Any disorders of Urinary System?		y disorder of th outh & Skin	e Eyes, Ears Nose,	Throat or
FOR FEMALE CANDIDATES ONLY				See Diag
a. Is there any history of diseases of breast/genital organs?		you have any h	istory of miscarriag	ge/ Y/N
 b. Is there any history of abnormal PAP Smear/Mammogram/USG of Pelvis or any other tests? (If yes attach reports) 	duri		, were there any co such as gestational of	
c. Do you suspect any disease of Uterus, Cervix or Ovaries?	f. Are	you now pregi	nant? If yes, how ma	any months?
CONFIDENTAIL COMMENTS FROM MEDICA	AL EXAMINE	R		HYSICAL DET
➤ Was the examinee co-operative?			Guna) 22	XIN
➤ Is there anything about the examine's health, life his/her job?	estyle that might	affect him/her	in the near future w	rith regard to
> Are there any points on which you suggest further	er information b	e obtained?		Y/N
Based on your clinical impression, please provid			nendations below;	
Nu				
	1.07			
. /				
Do you think he/she is MEDICALLY FIT or UN	NFIT for employ	ment.		
MEDICAL EXAMINER'S DECLARATION				
I hereby confirm that I have examined the above indi- above are true and correct to the best of my knowledge		fication of his/h	ner identity and the	findings stated
			TORY	218 / VOX 93
ring the last 5 years have you been medically	D 01111	ANIN 700	ned. All	(/NN)
Name & Signature of the Medical Examiner :	Dr. Nuns	ADID ZAO	The late of Violati	461
		DID ZAMAN	(P	
		MBBS No. 77615	INCA	producedore
Seal of Medical Examiner :		TCMC)	DIAGNO	lave you ever su
		abacelo le	GANDHINAGAR KOTTAYAM	· Psychological
Name & Seal of DDRC SRL Branch		18	685008	the Nervous : * Any disorder
		St. Stole	violetičii) te	

DDRC SRL Diagnostics Private Limited

Date & Time

ANDHINAGAR

Corp. Office: DDRC SRL Tower, G- 131, Panampilly Nagar, Ernakulam - 682 036 Ph No. 0484-2318223, 2318222, e-mail: info@ddrcsrl.com, web: www.ddrcsrl.com



Name

ATUL JOSE PHILIP

E C No 174532







Spite of a series of a series





मिलने पर निस्सितिशिक्त को जीटाए सहाराक महाप्रविधक (सरझा) वैक प्रीच सुदीदा, बहोदा कापोरेस सेन्टर सी - 26 जी-वर्गक, बान्दा पुल्ही कोम्प्रतीवस, मुंबई 1900,55 भगरत फोन् 91 12 6698 5196 फिक्स 91 22 2852 5747

H found, please-return to Apar, General Wanager (Security) Bank of Bandas, Baroda Corporne Centre C-25, G-Block, Sandra-Kinia Complex, Mumbai 400054 - India Paorie 41 22 5092 5196, F-91 22 2652 5747

रहर १६२ /Blood Group: O+ve प्रकार विकास विकास (Catholication Marks: A mole on the right hand small finger.



OPHTHALMOLOGY REPORT

ACCESSION NO:4036WA002673

This is to certify that I have examined

MR/MS PIVL JOSE PHILIP Aged 33 and

His / her visual standard is as follows.

Acuity of Vision

For Far

R. 616

L 6/6

For Near

R NE

L NG

Colour Vision

NORMAL

DATE: 14/01/23



OPTOMETRIST



Name: ATUL JOSE PHILIP

Age/Sex: 33 yrs/M

Accession No: 4036WA002673

Report Date: 14.01.2023 Ref.by: Mediwheel

USG ABDOMEN & PELVIS

OBSERVATIONS:

Normal in size. Shows normal parenchymal echotexture. No focal Liver:

parenchymal lesion noted. The biliary radicals appear normal. Portal

vein is normal (10 mm).

Distended (measures 4.9 x 1.3 cm). No calculus seen. No e/o of any Gall bladder:

wall thickening / edema. No e/o any pericholecystic collection.

Not dilated (6 mm). CBD:

Normal in size (10.8 cm) and echotexture. No focal lesion. Spleen:

Head (2 cm), body (1.2 cm) and tail (1.1cm) appear normal. No focal Pancreas:

lesion. No calcification or duct dilatation noted.

Right kidney length measures 10 cm. Parenchymal thickness 1.7 cm Kidneys:

Normal in position & size. Cortical echogenicity is normal. There is good cortico-medullary differentiation. No calculus or mass lesion

seen. No hydronephrosis.

Left kidney length measures 10.2 cm. Parenchymal thickness 1.7 cm

Normal in position & size. Cortical echogenicity is normal. There is good cortico-medullary differentiation. No calculus or mass lesion

seen. No hydronephrosis.

Not dilated. Ureters:

Urinary Bladder: Distended, No luminal or wall abnormality noted.

Normal in size, volume 24 cc. Shows homogenous parenchymal Prostate:

texture. No evidence of any mass lesion.

lymphadenopathy. No evidence of bowel wall evident Others:

thickening/echogenic mesentery/dilated bowel loops. Normal peristalsis

seen. No free fluid in the peritoneal cavity. No pleural effusion noted.

IMPRESSION:

No significant abnormality detected.

KOTTAYAN 686008

Dr. Deepak.V, MBBS, DMRD

Radiologist

Note: This is radiological opinion and not the final diagnosis. Ultrasound is limited by patient adiposity, bowel gas and correlate clinically and investigate further as needed.

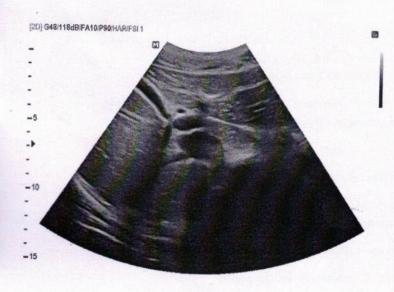
Exam 14-01-2023-0006

Accession # Exam Date Description Sonographer

14012023

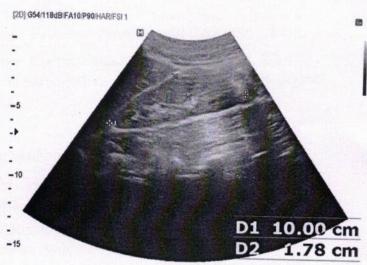
ame Birth Date Gender

Other

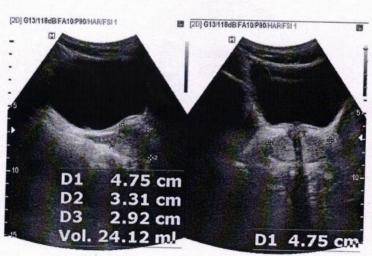














ECG REPORT

ACCESSION NO

: 4036WA002673

NAME

: ATUL JOSE PHILIP

AGE

: 33

SEX

: MALE

DATE

: 14.01.2023

COMPANY

: MEDIWHEEL

RATE

58/min

RHYTHM

Normal sinus Phythm

P. WAVE

Normal

P-R INTERVAL

- Nornal 120 ms

Q,R,S,T. WAVES

Norna

AXIS

worne

ARRHYTHMIAS

nd

QT INTERVAL

360 ms

OTHERS

Nid

GANDHINAGAR KOTTAYAM 686008

OPINION

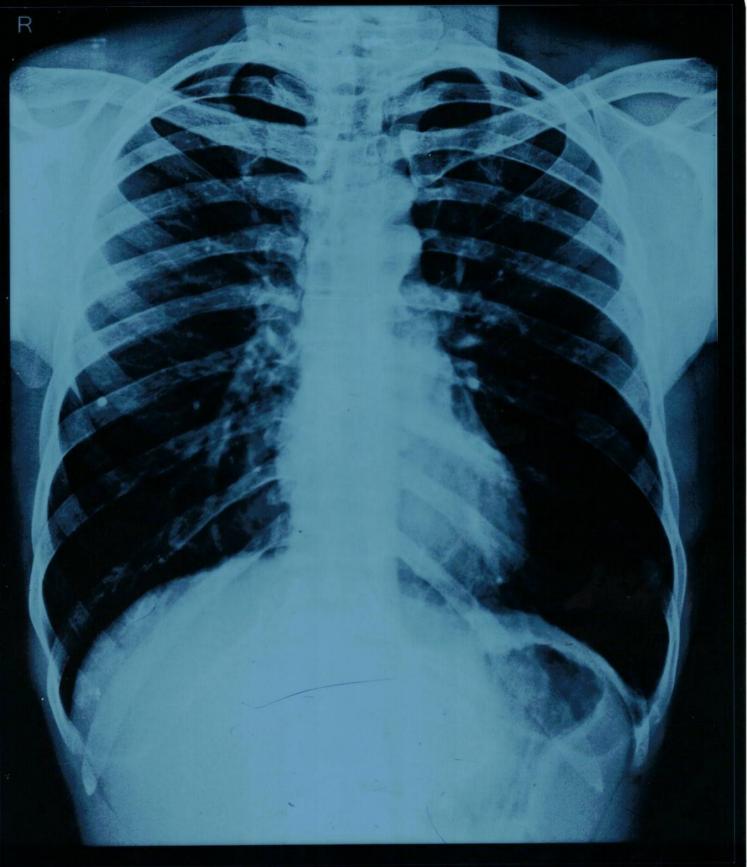
Normal ECG.

DEMUTADID ZAMAN AP MBBS

Reg. No. 77615 (TCMC)

CIN: U85190MH2006PTC161480

(Refer to "CONDITIONS OF REPORTING" Overleaf)



ATUL JOSE PHILIP 33/Y 5349 CHEST-PA 14-01-2023

DDRC SRL DIAGNOSTICS, GANDHI NAGAR, KOTTAYAM

10/11/





X - RAY CHEST - REPORT

ACCESSION NO

: 4036WA002673

NAME

: ATUL JOSE PHILIP

AGE

: 33

SEX

: MALE

DATE

: 14.01.2023

COMPANY

: MEDIWHEEL

EXPOSURE

POSITIONING

SOFT TISSUES

Normal shodows

LUNG FIELDS

Norma

HEART SHADOW

Norma

CARDIOPHRENIC ANGLE

not obliterated, Nornal

COSTOPHRENIC ANGLE

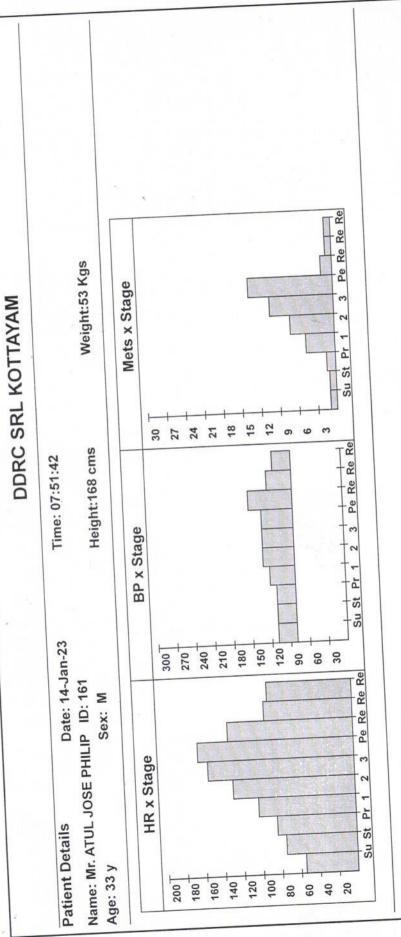
not obliterated, Normal

HILUM

OPINION

sormal chest dras

(Refer to "CONDITIONS OF REPORTING" Overleaf)



Interpretation

STRESSED UPTO 10 MTS ON BRUCE PROTOCOL AND ATTAINED 88% OF THR AT HR OF 165 BPM WITH A WORKLOAD OF 11 METS.RPP- 24750.

NORMAL HR AND BP RESPONSE.

NO ANGINA/ARRHYTHMIA.

BASELINE ECG SHOWS SINUS BRADYCARDIA. NO SIGNIFICANT ST SHIFT.

IMP:- TEST IS NEGATIVE FOR INDUCIBLE ISCHEMIA. GOOD EFFORT TOLERANCE.





Doctor:

(c) Schiller Healthcare India Pvt. Ltd. V 4.7

Summany Report edited by user) Ref. Doctor: --

DDRC SRL KOTTAYAM

Date: 14-Jan-23 Patient Details

Name: Mr. ATUL JOSE PHILIP ID: 161

Sex: M Age: 33 y

Clinical History: FOR CARDIAC EVALUATION

Medications: NIL

Test Details

Protocol: Bruce

10 m 0 s Total Exec. Time:

Max. BP: 150 / 80 mmHg

FATIGUE Test Termination Criteria:

Height:168 cms

Time: 07:51:42

Weight:53 Kgs

THR: 168 (90 % of Pr.MHR) bpm

Max. Mets: 13.50

Max. HR: 165 (88% of Pr.MHR) bpm Max. BP x HR: 24750 mmHg/min

Pr.MHR: 187 bpm

Min. BP x HR: 4400 mmHg/min

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0
-
0

Stage Name	Stage Time (min : sec)	Mets	Speed (mph)	Grade (%)	Heart Rate (bpm)	Max. BP (mm/Hg)	Max. ST Level (mm)	Max. ST Slope (mV/s)
	4 - 55	10	0	0	55	110 / 80	-5.94 V4	-5.661
Supine	25.50	0 7	0	0	75	110 / 80	-0.85 aVR	1.06 11
Standing	t	2. 4	17	10	103	120 / 80	-2.12 aVR	3.89 V3
	0 0	2 0	2.5	12	129	130 / 80	-1.06 aVR	4.95 V4
	0 0	10.0	2.4	14	155	130 / 80	-1.70 aVR	5.66 V3
	0.0	1.0.1	4.2	. 4	165	130 / 80	-1.70 aVR	5.66 V3
Peak EX	0	ς, α	1 -	0	133	150 / 80	-2.97 aVR	5.66 11
Recovery(1)	2.0	0. 1	. 0	0	94	120 / 80	-2.97 aVR	5.66 11
Recovery(2)	1.11	1.0	0	0	06	110 / 80	-1.27 aVR	3.18 V3



DDRC SRL KOTTAYAM

Exec Time: 6 m 0 s Stage Time: 3 m 0 s HR: 129 bpm

Date: 14-Jan-23

ID: 161

Mr. ATUL JOSE PHILIP (33 M)

DDRC SRL KOTTAYAM

Exec Time: 10 m 0 s Stage Time: 1 m 0 s HR: 165 bpm

(THR: 168 bpm) B.P: 130 / 80

Grade: .16 %

Date: 14-Jan-23

Mr. ATUL JOSE PHILIP (33 M)

DDRC SRL KOTTAYAM