

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.

1. Name of the examinee VIMAC Mr./Mrs./Ms. 2. Mark of Identification (Mole/\$cat/any other (specify location)): 3. Age/Date of Birth 21 - 12 - 1992 Gender: 29 F/M 4. Photo ID Checked

(Passport/Election Card/PAN Card/Driving Licence/Company ID)

PHYSICAL DETAILS:

a. Height	b. Weight	c. Girth of Ab	odomen
d. Pulse Rate	e. Blood Pressure:	Systolic	Diastolic
	1 st Reading	100	70
	2 nd Reading	16	· Aci
FAMILY HISTORY:	6 B	((0	.,0

Relation	Age if Living	Health Status	If deceased, age at the time and cause
Father	60	Leovy	
Mother	5	Ciord	· · · · · · · · · · · · · · · · · · ·
Brother(s) (2)	31,27	CLOUG	4 7
Sister(s) (1)	23	5,000	ibre,

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

Tobacco in any form	Sedative	Alcohol
Ale .	<u>N</u> 0	decenal

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. Ý/N
- b. Have you undergone/been advised any surgical procedure? Y/N

Have you ever suffered from any of the following?

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

- c. During the last 5 years have you been medically examined, received any advice or treatment or admitted to any hospital?
- d. Have you lost or gained weight in past 12 months?
 - YЛ

Y/N

Y/N

- 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 of any kind?

DDRC SRL Diagnostics Private Limited

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

Y/N

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

• Any disorders of Urinary System?

FOR FEMALE CANDIDATES ONLY

- a. Is there any history of diseases of breast/genital organs?
- b. Is there any history of abnormal PAP Smear/Mammogram/USG of Pelvis or any other tests? (If yes attach reports)
- c. Do you suspect any disease of Uterus, Cervix or **Ovaries**?

CONFIDENTAIL COMMENTS FROM MEDICAL EXAMINER

- > Was the examinee co-operative?
- > Is there anything about the examine's health, lifestyle that might affect him/her in the near future with regard to his/her job?

.....

Y/N

Y/N

- > Are there any points on which you suggest further information be obtained?
- > Based on your clinical impression, please provide your suggestions and recommendations below;

> Do you think he/she is MEDICALLY FIT or UNFIT for e ployment.

MEDICAL EXAMINER'S DECLARATION

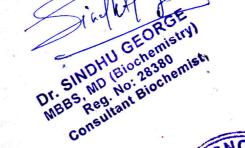
I hereby confirm that I have examined the above adividual after verification of his/her identity and the findings stated above are true and correct to the best of my knowledge.

Name & Signature of the Medical Examiner

Seal of Medical Examiner

Name & Seal of DDRC SRL Branch

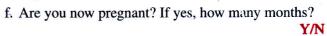
Date & Time



PI(

Any disorder of the Eyes, Ears Nose, Throat or Mouth & Skin

- d. Do you have any history of miscarriage/ abortion or MTP Y/N e. For Parous Women, were there any complication
- during pregnancy such as gestational diabetes. hypertension etc Y/N
- Y/N



Y/N

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Regd. Office: 4th Floor, Prime Square, Plot No.1, Gaiwadi Industrial Estate, S.V. Road, Goregaon (West), Mumbai - 400062.



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 Capital City,26/548/5,6,Ground Floor,Korappath Lane,Round North,Thrissur TRICHUR, 680020 KERALA, INDIA Tel : 9446425900 Email : thrissur.ddrc@srl.in

PATIENT NAME : VIMAL M T PATIENT ID : VIMAM0310934177 ACCESSION NO : 4177VJ000425 AGE : 29 Years SEX : Male ABHA NO: REPORTED : RECEIVED : 03/10/2022 16:25 03/10/2022 23:25 DRAWN : **REFERRING DOCTOR :** DR. SINDHU CLIENT PATIENT ID : **Test Report Status** Results **Biological Reference Interval** Units <u>Final</u>

MEDIWHEEL HEALTH CHEKUP BELOW 40(M)TMT

OPTHAL	
OPTHAL	ATTACHED
TREADMILL TEST	
TREADMILL TEST	COMPLETED
PHYSICAL EXAMINATION	
PHYSICAL EXAMINATION	COMPLETED







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BUN/CREAT RATIO				
BUN/CREAT RATIO	12.3		5.00 - 15.00	
CREATININE, SERUM				
CREATININE	0.89	Low	0.9 - 1.3	mg/dL
GLUCOSE, POST-PRANDIAL, PLASMA				
GLUCOSE, POST-PRANDIAL, PLASMA	96		Diabetes Mellitus : > or = 200 mg/dL. Impaired Glucose tolerance/ Prediabetes : 140 to 199 mg/dl Hypoglycemia : < 55 mg/dL.	
GLUCOSE, FASTING, PLASMA				
GLUCOSE, FASTING, PLASMA	93		Diabetes Mellitus : > or = 126 mg/dL. Impaired fasting Glucose/ Prediabetes : 101 to 125 mg/dl Hypoglycemia : < 55 mg/dL.	
GLYCOSYLATED HEMOGLOBIN, EDTA WHOLE B	LOOD			
GLYCOSYLATED HEMOGLOBIN (HBA1C)	5.4		Normal : $4.0 - 5.6$ %. Non-diabetic level : < 5.7 %. 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	108.3		< 116.0	mg/dL
CORONARY RISK PROFILE (LIPID PROFILE), S	ERUM			
CHOLESTEROL	152		Desirable: <200 BorderlineHigh : 200-239 High : > or = 240	mg/dL
TRIGLYCERIDES	45		Normal : < 150 High : 150-199 Hypertriglyceridemia : 200-499 Very High: > 499	mg/dL
HDL CHOLESTEROL	71	High	40 - 60	mg/dL







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DIRECT LDL CHOLESTEROL	79		Adult levels: Optimal < 100 Near optimal/above optimal: 1 129 Borderline high : 130-159 High : 160-189 Very high : = 190	mg/dL 00-
NON HDL CHOLESTEROL	81		Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL
CHOL/HDL RATIO	2.1	Low	3.30 - 4.40	
LDL/HDL RATIO	1.1		0.5 - 3.0	
VERY LOW DENSITY LIPOPROTEIN LIVER FUNCTION TEST WITH GGT	9.0		< or = 30.0	mg/dL
BILIRUBIN, TOTAL	0.53		< 1.1	mg/dL
BILIRUBIN, DIRECT	0.20		0.0 - 0.2	mg/dL
BILIRUBIN, INDIRECT	0.33		0.00 - 1.00	mg/dL
TOTAL PROTEIN	7.1		Ambulatory : 6.4 - 8.3 Recumbant : 6 - 7.8	g/dL
ALBUMIN	5.2		3.5 - 5.2	g/dL
GLOBULIN	1.9	Low	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO	2.7	High	1.0 - 2.0	RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	17		< 40	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT)	32		< 45	U/L
ALKALINE PHOSPHATASE	50		40 - 130	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT) TOTAL PROTEIN, SERUM	28		< 60	U/L
TOTAL PROTEIN	7.1		Ambulatory : 6.4 - 8.3 Recumbant : 6 - 7.8	g/dL
URIC ACID, SERUM				
URIC ACID Abo group & rh type, edta whole blood	5.3		3.4 - 7.0	mg/dL
ABO GROUP METHOD : GEL CARD METHOD	0			

NEGATIVE



RH TYPE





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BLOOD COUNTS				
HEMOGLOBIN	16.1		13.0 - 17.0	g/dL
RED BLOOD CELL COUNT	5.31		4.5 - 5.5	mil/µL
WHITE BLOOD CELL COUNT	5.54		4.0 - 10.0	thou/µL
PLATELET COUNT	279		150 - 410	thou/µL
RBC AND PLATELET INDICES	2,5			
HEMATOCRIT	46.0		40 - 50	%
MEAN CORPUSCULAR VOL	86.7		83 - 101	fL
MEAN CORPUSCULAR HGB.	30.3		27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION	35.0	High	31.5 - 34.5	g/dL
RED CELL DISTRIBUTION WIDTH	12.3		11.6 - 14.0	%
MEAN PLATELET VOLUME	10.6		6.8 - 10.9	fL
WBC DIFFERENTIAL COUNT - NLR				
SEGMENTED NEUTROPHILS	47		40 - 80	%
ABSOLUTE NEUTROPHIL COUNT	2.60		2.0 - 7.0	thou/µL
LYMPHOCYTES	49	High	20 - 40	%
ABSOLUTE LYMPHOCYTE COUNT	2.71		1 - 3	thou/µL
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	1.0			
EOSINOPHILS	02		1 - 6	%
ABSOLUTE EOSINOPHIL COUNT	0.11		0.02 - 0.50	thou/µL
MONOCYTES	02		2 - 10	%
ABSOLUTE MONOCYTE COUNT	0.11	Low	0.20 - 1.00	thou/µL
BASOPHILS	00		< 1 - 2	%
ABSOLUTE BASOPHIL COUNT	00	Low	0.02 - 0.10	thou/µL
ERYTHRO SEDIMENTATION RATE, BLOOD				
SEDIMENTATION RATE (ESR) STOOL: OVA & PARASITE	05		0 - 14	mm at 1 hr
COLOUR	BROWN			
CONSISTENCY	SEMI FORMED			
ODOUR	FAECAL			
MUCUS	NOT DETECTED		NOT DETECTED	
VISIBLE BLOOD	ABSENT		ABSENT	
POLYMORPHONUCLEAR LEUKOCYTES	0-1		0 - 5	/HPF







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Results

CLIENT CODE : CA00010147

DELHI INDIA

8800465156

DRAWN :

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VIMAM0310934177

Units

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<u>Final</u>

ACCESSION NO : 4177VJ000425

(main in the second se			
RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
CYSTS	NOT DETECTED	NOT DETECTED	
OVA	NOT DETECTED		
SUGAR URINE - POST PRANDIAL			
SUGAR URINE - POST PRANDIAL	NOT DETECTED	NOT DETECTED	
THYROID PANEL, SERUM			
Τ3	99.73	Male and Non-Pregnant : Pregnant Trimester-wise 1st : 81-190 2nd : 100-260 3rd : 100-260	70-204ng/dL
Τ4	7.10	3.2 - 12.6	µg/dl
TSH 3RD GENERATION	1.370	0.35 - 5.50	µIU/mL
URINE ANALYSIS			
COLOR	PALE YELLOW		
APPEARANCE	CLEAR		
SPECIFIC GRAVITY	1.010	1.003 - 1.035	
GLUCOSE	NOT DETECTED	NOT DETECTED	
KETONES	NOT DETECTED	NOT DETECTED	
BLOOD	NOT DETECTED	NOT DETECTED	
UROBILINOGEN	NORMAL	NORMAL	
EPITHELIAL CELLS	2-3	0-5	/HPF
RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
CASTS	NIL		
CRYSTALS	NIL		
BACTERIA	NOT DETECTED	NOT DETECTED	
CHEMICAL EXAMINATION, URINE			
PH	6.0	4.7 - 7.5	
PROTEIN	NOT DETECTED	NOT DETECTED	
BILIRUBIN	NOT DETECTED	NOT DETECTED	
NITRITE	NOT DETECTED	NOT DETECTED	
MICROSCOPIC EXAMINATION, URINE			<i></i>
WBC	1-2	0-5	/HPF
SERUM BLOOD UREA NITROGEN		6 20	<i>.</i>
BLOOD UREA NITROGEN	11	6 - 20	mg/dL

SEX : Male







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Test Report Status **Final** Results

Units

SUGAR URINE - FASTING

SUGAR URINE - FASTING

NOT DETECTED

SEX: Male

NOT DETECTED

Interpretation(s) CREATININE, SERUM-

DRAWN:

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)

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Lower than normal level may be due to:

Mvasthenia Gravis

Muscular dystrophy

GLUCOSE, POST-PRANDIAL, PLASMA-

ADA Guidelines for 2 h post prandial glucose levels is only after ingestion of 75grams of glucose in 300 ml water, over a period of 5 minutes.

GLUCOSE, FASTING, PLASMA-

ADA 2012 guidelines for adults as follows: Pre-diabetics: 100 - 125 mg/dL Diabetic: > or = 126 mg/dL

(Ref: Tietz 4th Edition & ADA 2012 Guidelines)

GLYCOSYLATED HEMOGLOBIN, EDTA WHOLE BLOOD-Glycosylated hemoglobin (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. Formation of GHb is essentially irreversible, and the concentration in the blood depends on both the life span of the red blood cell (average 120 days) and the blood glucose concentration. Because the rate of formation of GHb is directly proportional to the concentration of glucose in the blood,

the GHb concentration represents the integrated values for glucose over the preceding 6-8 weeks. Any condition that alters the life span of the red blood cells has the potential to alter the GHb level. Samples from patients with hemolytic anemias will exhibit decreased glycated hemoglobin values due to the shortened life span of the red cells. This effect will depend upon the severity of the anemia. Samples from patients with polycythemia or post-splenectomy may exhibit increased glycated hemoglobin values due to a somewhat longer life span of the red cells. Glycosylated hemoglobins results from patients with HbSS, HbCC, and HbSC and HbD must be interpreted with caution, given the pathological processes, including anemia,

increased red cell turnover, transfusion requirements, that adversely impact HbA1c as a marker of long-term glycemic control. In these conditions, alternative forms of testing such as glycated serum protein (fructosamine) should be considered.

"Targets should be individualized; More or less stringent glycemic goals may be appropriate for individual patients. Goals should be individualized based on duration of diabetes, age/life expectancy, comorbid conditions, known CVD or advanced microvascular complications, hypoglycemia unawareness, and individual patient considerations."

References

Tietz Textbook of Clinical Chemistry and Molecular Diagnostics, edited by Carl A Burtis, Edward R.Ashwood, David E Bruns, 4th Edition, Elsevier publication, 2006, 879-884.

2. Forsham PH. Diabetes Mellitus: A rational plan for management. Postgrad Med 1982, 71,139-154.

3. Mayer TK, Freedman ZR: Protein glycosylation in Diabetes Mellitus: A review of laboratory measurements and their clinical utility. Clin Chim Acta 1983, 127, 147-184. CORONARY RISK PROFILE (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 of the build up of plaques in your arteries that can lead to narrowed or blocked arteries throughout your body (atherosclerosis). High cholesterol levels usually don't cause any signs or symptoms, so a cholesterol test is an important tool. High cholesterol levels often are a significant risk factor for heart disease and important for diagnosis of hyperlipoproteinemia, atherosclerosis, hepatic and thyroid diseases.

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.







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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 triglycerides 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 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 syndrome.

Causes of decreased levels

- Low Zinc Intake
- OCP's Multiple Sclerosis

Nutritional tips to manage increased Uric acid levels

- Drink plenty of fluids
 Limit animal proteins
- High Fibre foods
- Vit C Intake
- Antioxidant rich foods 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-

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-

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. WBC DIFFERENTIAL COUNT - NLR-

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. ERYTHRO SEDIMENTATION RATE, BLOOD-







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REFERRING DOCTOR : DR. SINDHU		CLIENT PATIENT ID :
Test Report Status Final	Results	Units

Erythrocyte sedimentation rate (ESR) is a non - specific phenomena and is clinically useful in the diagnosis and monitoring of disorders associated with an increased production of acute phase reactants. The ESR is increased in pregnancy from about the 3rd month and returns to normal by the 4th week post partum. ESR is influenced by age, sex, menstrual cycle and drugs (eg. corticosteroids, contraceptives). It is especially low (0 -1mm) in polycythaemia, hypofibrinogenemia or congestive cardiac failure and when there are abnormalities of the red cells such as poikilocytosis, spherocytosis or sickle cells.

Reference

1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition

Paediatric reference intervals. AACC Press, 7th edition. Edited by S. Soldin
 The reference for the adult reference range is "Practical Haematology by Dacie and Lewis, 10th Edition"

SUGAR URINE - POST PRANDIAL-METHOD: DIPSTICK/BENEDICT'S TEST

THYROID PANEL, SERUM-Trilodothyronine T3, is a thyroid hormone. It affects 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.

Thyroxine T4, Thyroxine's principal function is to stimulate the metabolism of all cells and tissues in the body. Excessive secretion of thyroxine in the body is hyperthyroidism, and deficient secretion is called 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.

In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hypothyroidism, TSH levels are low. Below mentioned are the guidelines for Pregnancy related reference ranges for Total T4, TSH & Total T3

Delow mendoned	are the guidennes h	of frequency relate	a reference ranges for rotal
Levels in	TOTAL T4	TSH3G	TOTAL T3
Pregnancy	(µg/dL)	(µIU/mL)	(ng/dL)
First Trimester	6.6 - 12.4	0.1 - 2.5	81 - 190
2nd Trimester	6.6 - 15.5	0.2 - 3.0	100 - 260
3rd Trimester	6.6 - 15.5	0.3 - 3.0	100 - 260
Below mentioned	are the guidelines for	or age related refer	ence ranges for T3 and T4.
Т3		T4	

(μg/dL) 1-3 day: 8.2 - 19.9 1 Week: 6.0 - 15.9 (ng/dL) New Born: 75 - 260

NOTE: TSH concentrations in apparently normal euthyroid subjects are known to be highly skewed, with a strong tailed distribution towards higher TSH values. This is well

documented in the pediatric population including the infant age group. Kindly note: Method specific reference ranges are appearing on the report under biological reference range.

Reference:

1. Burtis C.A., Ashwood E. R. Bruns D.E. Teitz textbook of Clinical Chemistry and Molecular Diagnostics, 4th Edition.

Gowenlock A.H. Varley's Practical Clinical Biochemistry, 6th Edition.
 Behrman R.E. Kilegman R.M., Jenson H. B. Nelson Text Book of Pediatrics, 17th Edition MICROSCOPIC EXAMINATION, URINE-

Routine urine analysis assists in screening and diagnosis of various metabolic, urological, kidney and liver disorders Protein: Elevated proteins can be an early sign of kidney disease. Urinary protein excretion can also be temporarily elevated by strenuous exercise, orthostatic proteinuria, dehydration, urinary tract infections and acute illness with fever

Glucose: Uncontrolled diabetes mellitus can lead to presence of glucose in urine. Other causes include pregnancy, hormonal disturbances, liver disease and certain medications.

Ketones: Uncontrolled diabetes mellitus can lead to presence of ketones in urine. Ketones can also be seen in starvation, frequent vomiting, pregnancy and strenuous exercise.

Blood: Occult blood can occur in urine as intact erythrocytes or haemoglobin, which can occur in various urological, nephrological and bleeding disorders.

Leukocytes: An increase in leukocytes is an indication of inflammation in urinary tract or kidneys. Most common cause is bacterial urinary tract infection. Nitrite: Many bacteria give positive results when their number is high. Nitrite concentration during infection increases with length of time the urine specimen is retained in bladder prior to collection. pH: The kidneys play an important role in maintaining acid base balance of the body. Conditions of the body producing acidosis/ alkalosis or ingestion of certain type of food

can affect the pH of urine.

Specific gravity: Specific gravity gives an indication of how concentrated the urine is. Increased specific gravity is seen in conditions like dehydration, glycosuria and proteinuria while decreased specific gravity is seen in excessive fluid intake, renal failure and diabetes insipidus.

Bilirubin: In certain liver diseases such as biliary obstruction or hepatitis, bilirubin gets excreted in urine. Urobilinogen: Positive results are seen in liver diseases like hepatitis and cirrhosis and in cases of hemolytic anemia

SERUM BLOOD UREA NITROGEN-Causes of Increased levels

Pre renal

High protein diet, Increased protein catabolism, GI haemorrhage, Cortisol, Dehydration, CHF Renal
 Renal Failure

Post Renal

Malignancy, Nephrolithiasis, Prostatism

Causes of decreased levels









CLIENT'S NAME AND ADDRESS : MEDIWHEEL ARCOFEMI HEALTHCARE LIMITED F701A, LADO SARAI, NEW DELHI, SOUTH DELHI, DELHI, SOUTH DELHI 110030 DELHI INDIA 8900465156 8800465156

DDRC SRL DIAGNOSTICS Capital City, 26/548/5, 6, Ground Floor, Korappath Lane, Round North, Thrissur TRICHUR, 680020 KERALA, INDIA Tel: 9446425900 Email : thrissur.ddrc@srl.in

PATIENT NAME : VIMAL M T		PATIENT ID : VIMAM0310934177
ACCESSION NO : 4177VJ000425	AGE : 29 Years SEX : Male	ABHA NO :
DRAWN :	RECEIVED : 03/10/2022 16:25	REPORTED : 03/10/2022 23:25
REFERRING DOCTOR : DR. SINDHU		CLIENT PATIENT ID :
Test Report Status Final	Results	Units

• SIADH.

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







DELHI INDIA

8800465156

CLIENT'S NAME AND ADDRESS : MEDIWHEEL ARCOFEMI HEALTHCARE LIMITED F701A, LADO SARAI, NEW DELHI, SOUTH DELHI, DELHI, SOUTH DELHI 110030

DDRC SRL DIAGNOSTICS Capital City,26/548/5,6,Ground Floor,Korappath Lane,Round North,Thrissur TRICHUR, 680020 KERALA, INDIA Tel : 9446425900 Email : thrissur.ddrc@srl.in

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Test Report Status Final	Results	Units

MEDIWHEEL HEALTH CHEKUP BELOW 40(M)TMT

ECG WITH REPORT 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

DR.HARI SHANKAR, MBBS MD HEAD - Biochemistry & Immunology

SYALMA P THOMAS LAB TECHNICIAN

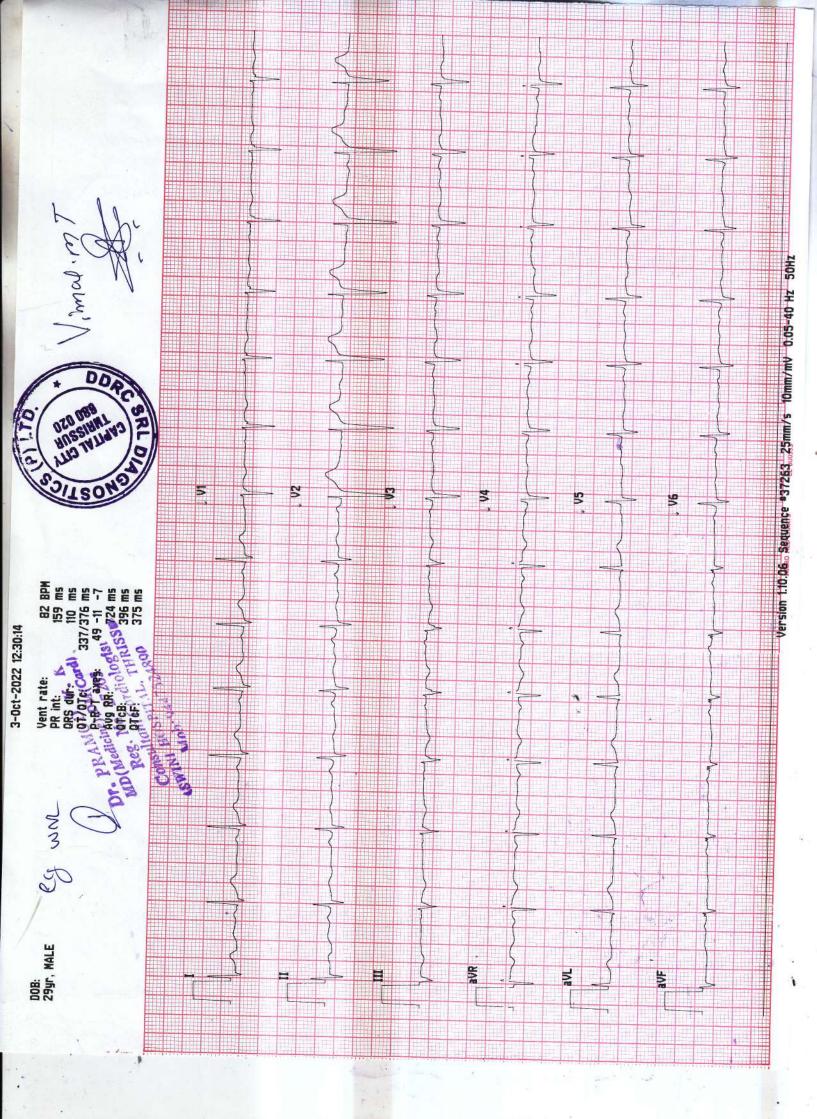
BIJI K S LAB TECHNICIAN

5

MANJU SHAJI RADIOGRAPHER









Name: MR.VIMAL M T Date: 03.10.2022 Age/Sex: 29 Y/ M AC 0425

CHEST X-RAY (PA View):

Trachea is central.

Cardiac shadow appears normal in size and configuration.

Both lung fields are clear.

Bilateral costophrenic and cardiophrenic angles are clear.

No focal consolidation, effusion, pulmonary edema, or pneumothorax.

Both hila appear normal.

Bony thorax and soft tissues are unremarkable.

IMPRESSION:

> No significant abnormality detected.



DR. JES PAULSON DMRD SULTANT RADIOLOGIST CON

Dr. Jeswin Paulson Mess, LARD Reg. No. 43581 Consultant Radiologist



Drishyam Eye Care Hospital LLP

See The World With Us



VISION CERTIFICATE

examined and results are as follows

			Right Eye			Left Eye
	Distant Vision	:	6/6			6/6
	Near vision	:	NG			NG
Contraction of the second	IOP(Intra ocular pressure)	:	16 mm tg	WNL	01 B)	14 mm Hg
	Anterior segment	2	NORMAL			NORMAL
	Fundus	:	NORMAL		2	NORMAL
	Squint	:	NORMAL			NORMAL
	Colour Vision	:	NORMAL			NORMAL

Doctor's Signature

Place : THRISSURDate : 3|10|2022 MEAC Dr. VI MBBS, DO SR. CONSUL

MBBS, DO, MMEL, FNGS (Ed) SR. CONSULTANT OPHTHALMOLOGIST Reg. No. 19178

Contact: 0487 22 222 99 www.drishyameye.com info@drishyameye.com Drishyam Eye Care Hospital LLP Opp. BSNL Office, Kovilakathumpadam, Thrissur, Kerala -680022 | Mob: +91 7025 11 11 99



Patient Name: Mr. VIMAL	Age: 29 Y	Sex: Male
Ref. Consultant:	AC No: 4177VJ002	Date: 03.10.2022
Clinical details:		

USG ABDOMEN

Liver measures 11 cm, normal in size and echotexture. No focal lesions seen. PV and CBD are normal in course and calibre. No dilatation of intrahepatic biliary radicles seen. Subphrenic spaces are normal.

Gall bladder is distended and appears normal. No calculus or mass seen.

Spleen measures 9.1 cm, normal in size and echotexture. No focal or diffuse lesions seen.

Pancreas: Head and body visualized, normal in size and echotexture. No focal lesions seen. No duct dilatation or calcification seen. Tail is obscured.

Right kidney measures 8.4 x 3.5 cm and left kidney measures 8.5 x 3.7 cm. Both kidneys are normal in size and cortical echogenicity. Cortico medullary differentiation is maintained. No calculus or dilatation of pelvicalyceal system on both sides.

Urinary bladder is partially distended and grossly appears normal.

Prostate measures 13 cc, normal in size and echotexture.

No ascites. No definite evidence of any abnormal bowel dilatation / wall thickening seen.

IMPRESSION

> No significant abnormality detected.



Thanks for your referral. Ultrasound reports need not be fully accurate. It has to be correlated clinically and with relevant investigations.

Dr. Jeswin Paulson MBBS, DMRD Reg. No. 43581 Consultant Radiologist

-tiont	Mr. VIMAL 29 M	Age/Sex	29 Years / Male
atient name	210511SU2-22-10-03-16	Visit No	1
atient ID		Visit Date	03/10/2022
eferred by	Dr. SELF	AC DE ARD MI 0.7	DDRC SRL DIAGNOSTICS
VIMAL 29 M, 2105115U2-22-10-0 PB PB D1 3.33cm 2D2 2.58cm 3D3 2.87cm Vol 12.911cm ³ Vol 12.911cm ³ Vol 12.911cm ³ Vol 12.911cm ³	ABD-NEW Har-low Gn -15 Sri II 51 CR12 Sri II 51 CR12 Sri II 51 CR12 ABD-NEW Har-low Gn -15 Sri II 51 CR12 ABD-NEW Har-low Sri II 51 CR12 Sri II 51 CR12 ABD-NEW Har-low Sri II 51 CR12 Sri II 51 CR12 ABD-NEW Har-low Sri	2-22-10-03-15 16.7cm/1.2/44Hz TIS 0.4 U.B U.B U.B U.B U.B U.B U.B U.B	DDRC SRL DIAGNOSTICS 03.10.2022 2:29:52 PM ABD-NEW Har-low 1000 C7 / M7 SHIILS / CR12 03.10.2022 2:30:42 PM ABD-NEW Har-low 1000 C7 / M7 FF2 / E3 C7 / M7 FF2 / FF3 / M7 FF2 / FF3 / M7 FF2 / FF3 / M7 FF2 /
1 D 10.97cm 2 D 8.45cm 3 D 3.50cm What 29 M. 2 D 115115U2-22-1 - 98	AC-RS/ABD MI 0.7 DDRC SRL DIAGNOSTICS 0-05-15 15.7cm/1.2/44Hz TIs 0.4 0.3.0.2022 2.32.57 PM ABD-NEW Har-kow SJORP GR -19 SRL DIAGNOSTICS CT / N7 FF2/E3 SRL DIAGNOSTICS		

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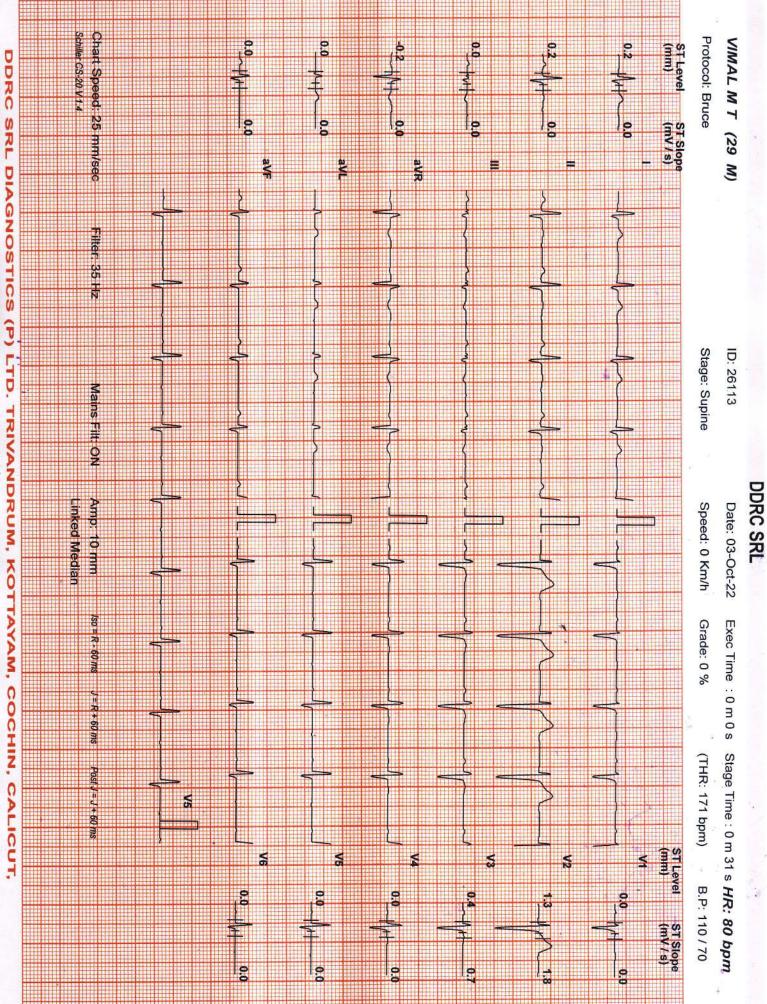


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DDRC SRL DIAGNOSTICS (P) LTD. TRIVANDRUM, KOTTAYAM, COCHIN, CALICUT,



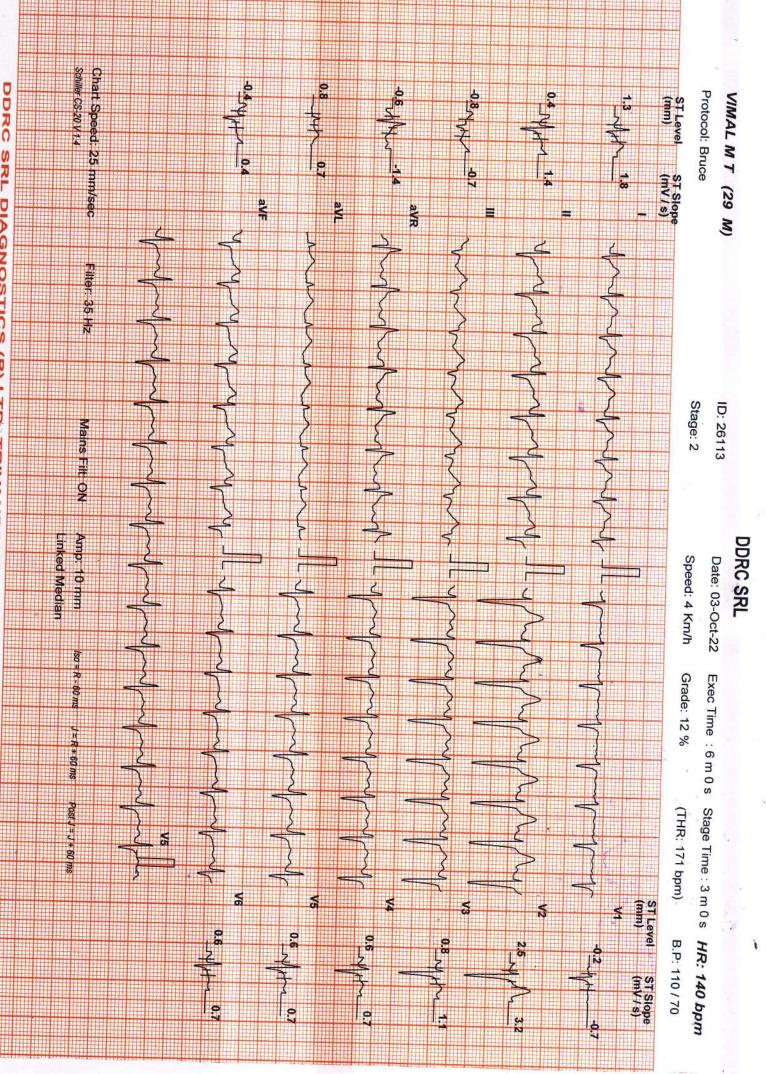


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DDRC SRL DIAGNOSTICS (P) LTD, TRIVANDRUM, KOTTAYAM, COCHIN, CALICUT,

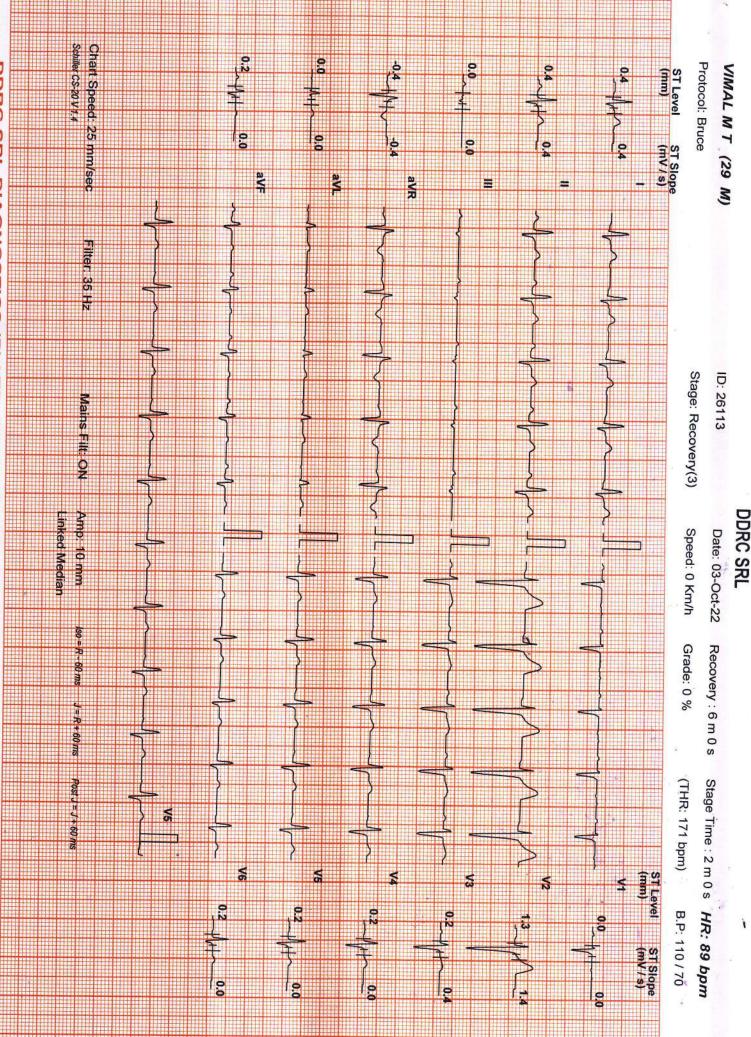
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Date: 03-Oct-22 Speed: 6.7 Km/h				And Make		MAMM		And Mark	A Martine		Stage: Peak Ex	ID: 26113
	Linked			Why Thy				Why The And	Mr The Mark		Speed: 6.7 Km	Date: 03-Oct-22
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Mains Filt ON						- Arra			Stage: Recovery(1)	ID: 26113
Amp: 10 mm s Linked Median	A A								Speed: 0 Km/h	Date: 03-Oct-22
\$\$\$ = R - 60 ms J = R + 60 ms		-l-l-				M			K VS	Recovery: 2 m 0 s
Post J = J + o0 ms		ر ۱						ST Lev (mm)		Stage Time : 2 m 0 s
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	5 Post J = J + 60 ms	150 = R60 ms	Amp: 10 mm / Linked Median	Mains Filt: ON	c Filler: 35 Hz	Chart Speed: 25 mm/sec Schiller:CS:20/V1:4
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	(THR: 171 bpm)	Grade: 0 %	Speed: 0 Km/h	Stage: Recovery(2)		Bru
m0s HR: 98 bpm	Stage Time : 2 m 0 s	Recovery : 4 m 0 s	Date: 03-Oct-22	ID: 26113	0	VIMAL M T (29 M)

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DDRC SRL

Patient Details	Date: 03-Oct-22	Time: 1:17:42 PM			
Name: VIMAL M T ID	: 26113				
Age: 29 y	Sex: M	Height: 167 cms	Weight: 64 Kgs		
Clinical History:					

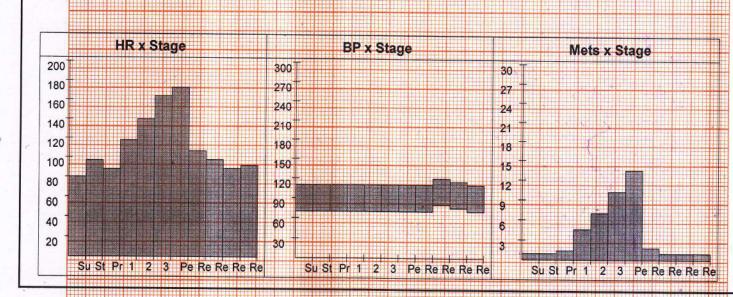
Medications:

Test Details

Protocol: Bruce	Pr.MHR: 191 bpm	THR: 171 (90 % of Pr.MHR) bpm
Total Exec. Time: 9 m 29 s	Max. HR: 173 (91% of Pr.MHR)bpm	Max. Mets: 13.50
Max. BP: 120/80 mmHg	Max. BP x HR: 20760 mmHg/min	Min. BP x HR: 5600 mmHg/min
Test Termination Criteria:		

Protocol Details

Stage Name	Stage Time (min : sec)	Mets	Speed (Km/h)	Grade (%)	Heart Rate (bpm)	Max. BP (mm/Hg)	Max. ST Level (mm)	Max. ST Slope (mV/s)
Standing	0:30	1.0	0	0	97	110 / 70	-0.42 aVR	1.77 V2
1	3:0	4.6	2.7	10	118	110/70	-0.85 aVR	2.83 V2
2	3:0	7.0	4	12	140	110 / 70	-1.27 aVR	4.25 V2
3	3:0	10.2	5.4	14	164	110/70	-1.27 aVR	5.66 V2
Peak Ex	0:29	13.5	6.7	16	173	110/70	-1.06 aVR	5.66 V2
Recovery(1)	2:0	1.8	1.6	0	107	110 / 70	-1.70 aVR	5.66 V2
Recovery(2)	2:0	1.0	0	0	98	120/80	-0.64 aVR	5.66 V2
Recovery(3)	2:0	1.0	0	0	89	115/75	-0.42 aVR	1.77 V2
Recovery(4)	0:14	1.0	0	0	92	110 / 70	-0.42 aVR	1.42 V2



DDRC SRL

Patient Details Date: 03-Oct-22 Time: 1:17:42 PM Name: VIMAL M T ID: 26113 Age: 29 y Sex: M Height: 167 cms Weight: 64 Kgs

Interpretation

Ref. Doctor: -----

(Summary Report edited by user)

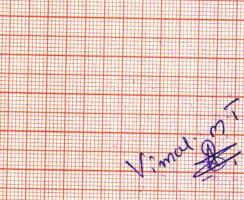
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Doctor: -----

Schiller CS-20 V 1.4

COCHIN, CALICUT, TRIVANDRUM, KOTTAYAM, LTD. 6 DIAGNOSTICS SRL DDRC