

Name Mr. ATHUL PRAMOD		Age/Sex	33Y/Male	
Ref from:	MEDI WHEEL	Date	27.08.2022	

ULTRASOUND SCAN OF ABDOMEN AND PELVIS

(With relevant image copies)

LIVER: Normal in size and **shows diffusely increased echotexture.** No e/o focal parenchymal lesions / IHBD. PV, HV & IVC are within normal limits.

GB: Normally distended, normal wall thickness. No e/o calculi/polyps/pericholecystic collections.

CBD: Normal

PANCREAS: Head and body visualized, and are of normal size and echotexture. No e/o focal/diffuse parenchymal lesions/ductal dilatation/calculi. Tail could not be visualized due to poor acoustic window.

SPLEEN: Normal in size and echotexture. Splenic vein shows normal diameter.

KIDNEYS: Both kidneys are normal in size and echotexture. No e/o calculi/hydronephrosis/focal lesions/perinephric collections.

RIGHT KIDNEY: Measures 104 x 48 mms LEFT KIDNEY: Measures 110 x 54 mms

UB: Well distended, shows normal wall thickness. No e/o calculi/ growth/

diverticulae. Both UV junctions are within normal limits.

PROSTATE: 15 cc, normal in size and echotexture.

No e/o intraperitoneal free fluid/ abdominal lymphadenopathy /mass lesion.

IMPRESSION:

- GRADE I FATTY LIVER
- > NO OTHER SONOLOGICALLY DETECTED ABNORMALITY.

Dr. P.NIYAZI NASIR

MBBS, DMRD

(Because of technical and technological limitation complete diagnosis cannot be assured on imaging sonography. Clinical correlation, consultation if required repeat imaging required in the event of controversies. This document is not for legal purposes).

Dr. P. NIYAZI NASIR. MBBS, DMRD REG. No. 41419 CONSULTANT RADIOLOGIST DDRC SRL DIAGNOSTIC (P)*LTD. KANNUR



DDRC SRL DIAGNOSTICS

KANNUR KERALA, INDIA Tel: 93334 93334

Email: customercare.ddrc@srl.in

PATIENT NAME: ATHUL PRAMOD PATIENT ID: ATHUM2708884053

ACCESSION NO: 4053VH002530 AGE: 34 Years SEX: Male

DRAWN: RECEIVED: 27/08/2022 11:56 REPORTED: 30/08/2022 15:41

REFERRING DOCTOR: SELF CLIENT PATIENT ID:

Test Report Status <u>Final</u> Results Biological Reference Interval Units

MEDIWHEEL HEALTH CHEKUP BELOW 40(M)TMT

OPTHAL

OPTHAL COMPLETED

TREADMILL TEST

TREADMILL TEST COMPLETED







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LIVER PROFILE - EXTENDED				
ASPARTATE AMINOTRANSFERASE (AST/SGOT)	25		< 40	U/L
ALANINE AMINOTRANSFERASE (ALT/SGPT)	50	High	< 45	U/L
ALKALINE PHOSPHATASE	76		40 - 130	U/L
LACTATE DEHYDROGENASE	188		135 - 225	U/L
BILIRUBIN, TOTAL	0.70		Upto 1.2	mg/dL
BILIRUBIN, DIRECT	0.21		< 0.31	mg/dL
BILIRUBIN, INDIRECT	0.49		0.00 - 0.60	mg/dL
TOTAL PROTEIN	7.3		Ambulatory: 6.4 - 8.3 Recumbant: 6 - 7.8	g/dL
ALBUMIN	4.8		3.5 - 5.2	g/dL
GLOBULIN	2.5		2.0 - 4.0	g/dL
ALBUMIN/GLOBULIN RATIO	1.9		1.0 - 2.0	Ratio
HEPATITIS B SURFACE ANTIGEN	NON REACTIVE		NON REACTIVE	
BUN/CREAT RATIO				
BUN/CREAT RATIO	21	High	5.00 - 15.00	
CREATININE, SERUM				
CREATININE	0.90		0.9 - 1.3	mg/dL
GLUCOSE, POST-PRANDIAL, PLASMA				
GLUCOSE, POST-PRANDIAL, PLASMA	146	High	Diabetes Mellitus : > or = 200 mg/dL. Impaired Glucose tolerance/ Prediabetes : 140 to 199 mg/d Hypoglycemia : < 55 mg/dL.	mg/dL L.
GLUCOSE, FASTING, PLASMA				
GLUCOSE, FASTING, PLASMA	98		Diabetes Mellitus : > or = 126 mg/dL.	mg/dL

Impaired fasting Glucose/ Prediabetes: 101 to 125 mg/dL. Hypoglycemia: < 55 mg/dL.

GLYCOSYLATED HEMOGLOBIN, EDTA WHOLE BLOOD





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GLYCOSYLATED HEMO	GLOBIN (HBA1C)	6.2	High	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%.	%
CORONARY RISK PR	OFILE (LIPID PROFILE),	SERUM			
CHOLESTEROL		234	High	< 200 Desirable 200 - 239 Borderline High >/= 240 High	mg/dL
TRIGLYCERIDES		135		< 150 Normal 150 - 199 Borderline High 200 - 499 High >/=500 Very High	mg/dL
HDL CHOLESTEROL		48		< 40 Low >/=60 High	mg/dL
DIRECT LDL CHOLEST	EROL	161	High	< 100 Optimal 100 - 129 Near or above optima 130 - 159 Borderline High 160 - 189 High >/= 190 Very High	mg/dL al
NON HDL CHOLESTER	OL	186	High	Desirable-Less than 130 Above Desirable-130-159 Borderline High-160-189 High-190-219 Very High- >or =220	mg/dL
CHOL/HDL RATIO		4.9	High	3.3 - 4.4 Low Risk 4.5 - 7.0 Average Risk 7.1 - 11.0 Moderate Risk > 11.0 High Risk	
LDL/HDL RATIO		3.4	High	0.5-3 Desirable/Low risk 3.1-6 Borderline/Moderate risk >6.0 High Risk	
VERY LOW DENSITY L	IPOPROTEIN	27		= 30</td <td>mg/dL</td>	mg/dL
LIVER FUNCTION TE	ST WITH GGT				
TOTAL PROTEIN		7.3		Ambulatory: 6.4 - 8.3 Recumbant: 6 - 7.8	g/dL
ALBUMIN		4.8		3.5 - 5.2	g/dL
GLOBULIN		2.5		2.0 - 4.0	g/dL
ALBUMIN/GLOBULIN R	ATIO	1.9		1.0 - 2.0	RATIO
ASPARTATE AMINOTRA	ANSFERASE (AST/SGOT)	25		< 40	U/L
ALANINE AMINOTRANS	SFERASE (ALT/SGPT)	50	High	< 45	U/L







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ALKALINE PHOSPHATASE	76	40 - 130	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT)	72	High < 60	U/L
URIC ACID, SERUM			
URIC ACID	5.2	3.4 - 7.0	mg/dL
ABO GROUP & RH TYPE, EDTA WHOLE BLOOD			
ABO GROUP	TYPE A		
RH TYPE	POSITIVE		
BLOOD COUNTS			
HEMOGLOBIN	15.1	13.0 - 17.0	g/dL
RED BLOOD CELL COUNT	4.81	4.5 - 5.5	mil/µL
WHITE BLOOD CELL COUNT	7.23	4.0 - 10.0	thou/µL
PLATELET COUNT	216	150 - 410	thou/µL
RBC AND PLATELET INDICES			
HEMATOCRIT	44.0	40 - 50	%
MEAN CORPUSCULAR VOL	91.6	83 - 101	fL
MEAN CORPUSCULAR HGB.	31.3	27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION	34.2	31.5 - 34.5	g/dL
MEAN PLATELET VOLUME	10.1	6.8 - 10.9	fL
WBC DIFFERENTIAL COUNT - NLR			
SEGMENTED NEUTROPHILS	69	40 - 80	%
ABSOLUTE NEUTROPHIL COUNT	4.99		thou/µL
LYMPHOCYTES	26	20 - 40	%
ABSOLUTE LYMPHOCYTE COUNT	1.81		thou/µL
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	2.8		
EOSINOPHILS	2	1 - 6	%
ABSOLUTE EOSINOPHIL COUNT	0.14		thou/µL
MONOCYTES	2	2 - 10	%
ABSOLUTE MONOCYTE COUNT	0.14		thou/µL
BASOPHILS	1	0 - 1	%
ERYTHRO SEDIMENTATION RATE, BLOOD			
SEDIMENTATION RATE (ESR)	6	0 - 14	mm at 1 hr
STOOL: OVA & PARASITE			
COLOUR	YELLOW		



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CONSISTENCY	SEMI LIQUID		
ODOUR	FAECAL		
MUCUS	NOT DETECTED	NOT DETECTED	
POLYMORPHONUCLEAR LEUKOCYTES	1-2	0 - 5	/HPF
RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
OVA	NOT DETECTED		
SUGAR URINE - POST PRANDIAL			
SUGAR URINE - POST PRANDIAL	DETECTED (TRACE)	NOT DETECTED	
URINALYSIS			
COLOR	PALE YELLOW		
APPEARANCE	CLEAR		
PH	5.0	4.7 - 7.5	
SPECIFIC GRAVITY	1.020	1.003 - 1.035	
GLUCOSE	NOT DETECTED	NOT DETECTED	
PROTEIN	NOT DETECTED	NOT DETECTED	
KETONES	NOT DETECTED	NOT DETECTED	
BILIRUBIN	NOT DETECTED	NOT DETECTED	
UROBILINOGEN	NORMAL	NORMAL	
WBC	2-3	0-5	/HPF
EPITHELIAL CELLS	0-1	0-5	/HPF
RED BLOOD CELLS	NOT DETECTED	NOT DETECTED	/HPF
CASTS	NOT DETECTED		
CRYSTALS	NOT DETECTED		
BACTERIA	NOT DETECTED	NOT DETECTED	
THYROID PANEL, SERUM			
Т3	132.10	80.00 - 200.00	ng/dL
T4	5.78	5.10 - 14.10	μg/dl
TSH 3RD GENERATION	1.540	0.4 - 4.2	μIU/mL

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



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• Problems during pregnancy, such as seizures (eclampsia)), or high blood pressure caused by pregnancy (preeclampsia)

Lower than normal level may be due to:

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Myasthenia GravisMuscular dystrophy

GLUCOSE, POST-PRANDIAL, PLASMA-ADA Guidelines for 2hr 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

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

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.
URIC ACID, SERUM-





Scan to View Report



CLIENT CODE: CA00010147 CLIENT'S NAME AND ADDRESS :

MEDIWHEEL ARCOFEMI HEALTHCARE LIMITED F701A, LADO SARAI, NEW DELHI, SOUTH DELHI, DELHI, SOUTH DELHI 110030 **DELHI INDIA** 8800465156

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Causes of Increased levels

Dietary

- High Protein Intake.
- Prolonged Fasting,Rapid weight loss.

Lesch nyhan syndrome.

Metabolic syndrome.

Causes of decreased levels

- . Low Zinc Intake
- Multiple Sclerosis

Nutritional tips to manage increased Uric acid levels

- Drink plenty of fluidsLimit 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.

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 - NLRThe 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
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.

- 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

URINALYSIS-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

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





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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

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THYROID PANEL, SERUMTriiodothyronine 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

Levels in TOTAL T4 TSH3G TOTAL T3

(µIU/mL) 0.1 - 2.5 0.2 - 3.0 0.3 - 3.0 (ng/dL) 81 - 190 100 - 260 100 - 260 Pregnancy First Trimester (µg/dL) 6.6 - 12.4 6.6 - 15.5 6.6 - 15.5 2nd Trimester 3rd Trimester Below mentioned are the guidelines for age related reference ranges for T3 and T4. T3 T4

(µg/dL) 1-3 day: 8.2 - 19.9 1 Week: 6.0 - 15.9 (na/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.

- 1. Burtis C.A., Ashwood E. R. Bruns D.E. Teitz textbook of Clinical Chemistry and Molecular Diagnostics, 4th Edition. 2. Gowenlock A.H. Varley's Practical Clinical Biochemistry, 6th Edition.
- 3. Behrman R.E. Kilegman R.M., Jenson H. B. Nelson Text Book of Pediatrics, 17th Edition





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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

JINSHA KRISHNAN **LAB TECHNICIAN**

JINISHA M

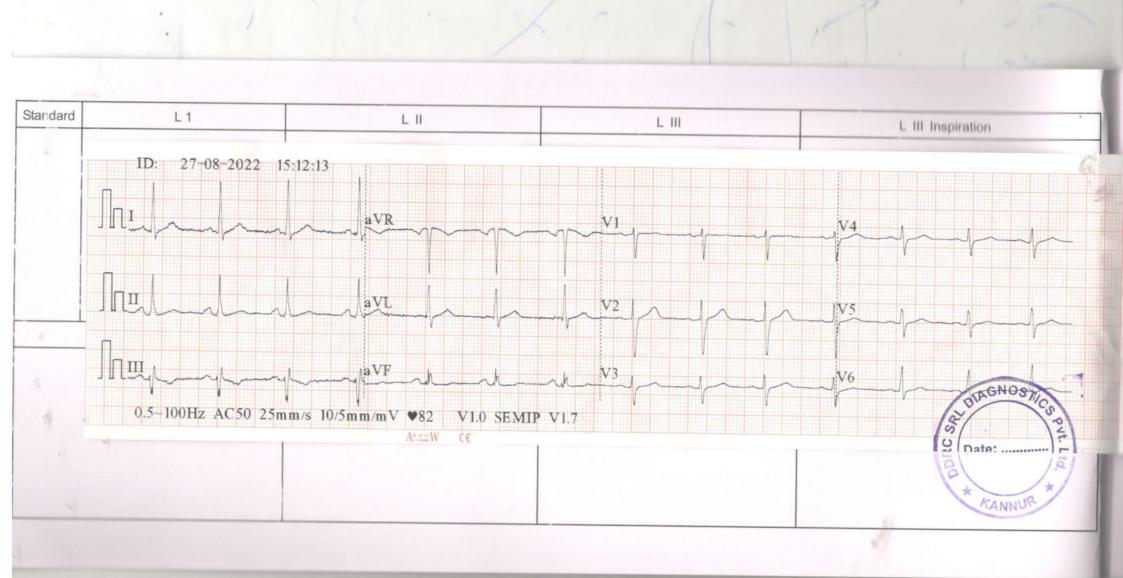
LAB TECHNICIAN

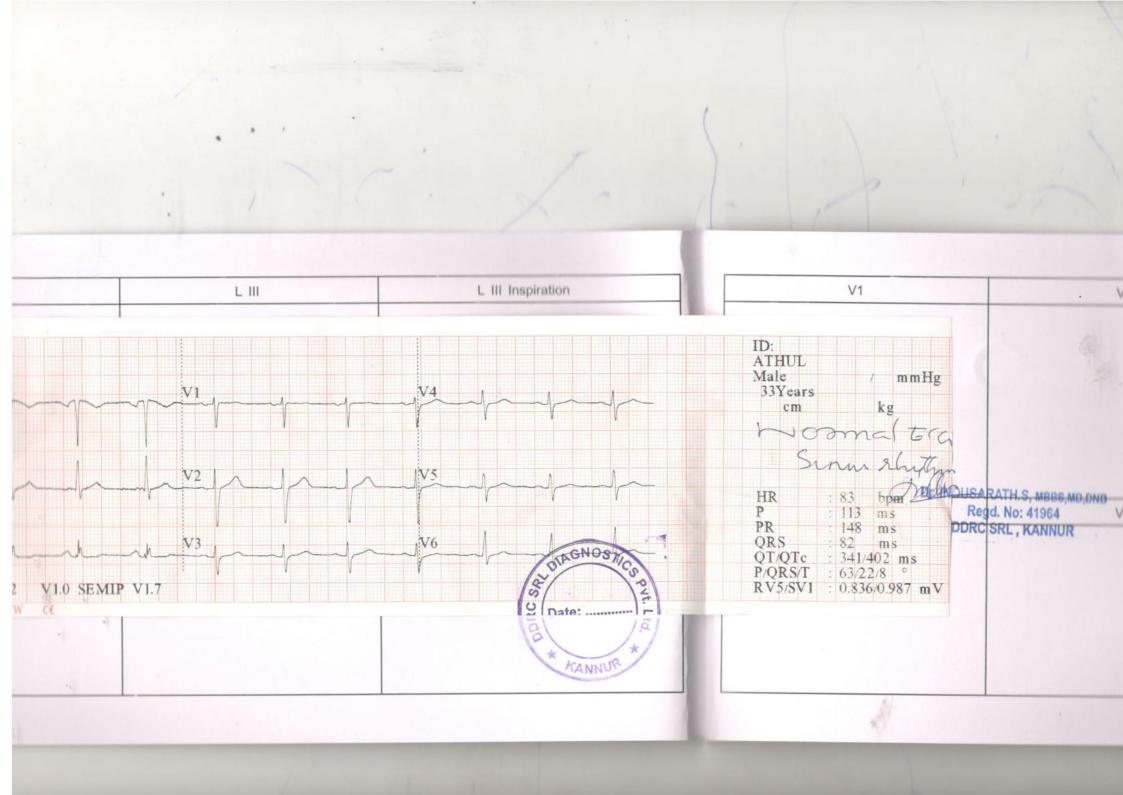
NIMISHA K LAB TECHNICIAN

JANCY T P LAB TECHNICIAN











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.

medical examination	to the examinee.					A COLOR
 Name of the e Mark of Ident Age/Date of E Photo ID Che 	ification : (M	ole/Scar/any	other (specification Card/PAN	y location		f foreary
PHYSICAL DETA	ILS:					
a. Height 16 9	(cms) b. We	eight 82	(Kgs)	с. (Girth of Abdome	en99 (cms)
d. Pulse Rate		ood Pressure			tolic I	
			1 st Reading		110	70
			2 nd Reading		1+0	70
FAMILY HISTOR	Y:					
Relation	Age if Living	Health	Status	If dec	ceased, age at th	e time and cause
Father	72	Hea	Chan			
Mother	60	'n				51, 24 51, 634
Brother(s)						
Sister(s)	35)	4 54 56			ndittan shink may on
	TIONS: Does the exam			following	2-1	Alcohol
Tobacco	o in any form	3	Sedative		MARA E 2010, TO	
The same specific of the same state of the same		10	occasional		esterially	
PERSONAL HIST	ORY					
from any menta If No, please att	ly in good health and er l or Physical impairmen ach details. gone/been advised any	t or deformi	ty. exan	nined, rece	eived any advice	ou been medically or treatment or treatment or it in past 12 months
Have you ever suffe	ered from any of the fo	ollowing?		2011	1.80	
 Psychological Disorders or any kind of disorders of the Nervous System? Any disorders of Respiratory system? 			Une	The state of the s		
Any Cardiac or	Circulatory Disorders? or any form of Cancer/Tu	Y	· Have		tested for HIV	/HBsAg / HCV Y
Any Musculosk		Y(• Are	you preser	ntly taking medi	cation of any kind?

DDRCSRL 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

 Any disorders of Urinary System? 	YN	 Any disorder of the Eyes, Ears, Nose, Throat or Mouth & Skin
FOR FEMALE CANDIDATES ONLY		Diagnostic Services
a. Is there any history of diseases of breast/genital organs?	Y/N	d. Do you have any history of miscarriage/ abortion or MTP
b. Is there any history of abnormal PAP Smear/Mammogram/USG of Pelvis or any other tests? (If yes attach reports)	Y/N	e. For Parous Women, were there any complication during pregnancy such as gestational diabetes, hypertension etc
c. Do you suspect any disease of Uterus, Cervix or Ovaries?	Y/N	f. Are you now pregnant? If yes, how many months?
CONFIDENTAIL COMMENTS FROM MEDICA	AL EXA	AMINER PARATHER LANDS
➤ Was the examinee co-operative?		The state of the s
Is there anything about the examine's health, life his/her job?	style tha	at might affect him/her in the near future with regard to
Are there any points on which you suggest further	er inform	nation be obtained?
> Based on your clinical impression, please provid	e your s	uggestions and recommendations below;
	************	Mary Co- 1 out
➤ Do you think he/she is MEDICALLY FIT or UN	FIT for	employment.
medie	ally	of to ammore subsection 1820 11 miles 2/2/11
MEDICAL EXAMINER'S DECLARATION		Treatments and pro-
I hereby confirm that I have examined the above indivabove are true and correct to the best of my knowledge		ter verification of his/her identity and the findings stated
Name & Signature of the Medical Examiner :	D	In Suranally Sound Um
Description of the Wedleth Examiner	TV	Dr. INDUSARATH.S, MBBS, MD, DMB Regd. No: 41964 DDRC SRL, KANNUR
Seal of Medical Examiner :		Touches An
		DIAGNOS . In one ment trees the seed but a
		Production of the second of th
Name & Seal of DDRC SRL Branch :		O Date Valore Carolina Salar S

DDRC SRL Diagnostics Private Limited

27.08. 22

Date & Time



ഭാരത സർക്കാർ GOVERNMENT OF INDIA

അതുൽ പമോദ് Athul Pramod



mm cura-de-Year of Birth: 1988 அவுகள் / Male

8286 7232 3665



ആധാർ – സാധാരണക്കാരന്റെ അവകാശം

7016260166





ഭാരതീയ സവിശേഷ തിരിച്ചറിയൽ അതോര UNIQUE IDENTIFICATION AUTHORITY OF INDIV

മേൽവിലാസം: S/O: കെ പി പമ്രോദ് കിഴക്കേടത്ത് ഹൌസ്, നിയര് ഓണ്ടൻ പറമ്പ ചൊവ്വ, ചൊവ്വ, കണ്ണൂർ, കേരളം, 670006

Address: S/O: K P Pramod Kizhakkedath House, Neal Onden Paramba, Chovva, Kannur, Chovva, Kerala, 670006



M help@uidai.gov.ir



OPTHALMOLOGY REPORT

TO WHOM-SO-EVER IT MAY CONCERN

This is to certify that I have examined Mr. ATHUL PRAMOD, 34 years Male on 27.08.2022 and his visual standards are as follows:

	OD	OS
UNCORRECTED DISTANCE VISUAL ACUITY	6/6	6/6
UNCORRECTED NEAR VISUAL ACUITY	N6	N6
COLOUR VISION	NORMAL	NORMAL

NOTE: NO HISTORY OF SPECS

NO RELEVANT MEDICAL HISTORY

VIMEGA .V OPTOMETRIST Date:

DATE: 27.08.2022

MEDIWHEEL ARCOFEMI CHEST P-A 27-Aug-22 01:52 PM

DDRC SRL KANNUR



INDIA'S	LEADING	DIAGNOS	TICS NE	TWORK

Name	Mr. ATHUL PRAMOD	Age/Sex	34Y/Male
Ref: By:	MEDIWHEEL	Date	27.08.2022

Thanks for referral

CHEST X-RAY - PA VIEW

Trachea is central. Carina and principal bronchi are normal.

Cardio-thoracic ratio is within normal limits.

Both lungs show normal Broncho-vascular markings. No definite focal opacities noted.

No volume loss in either hemithorax.

No definite mediastinal widening or other abnormalities noted.

CP angles, diaphragm, bony cage and soft tissue shadows - not remarkable.

IMPRESSION:

Normal X-ray chest

DR. P. NIYAZI NASIR, MBBS, DMRD

(Because of technical and technological limitation complete diagnosis cannot be assured on imaging sonography. Clinical correlation, consultation if required repeat imaging required in the event of controversies. This document is not for legal purposes).

Dr. P. NIYAZI NASIR. MB88, DMRO REG. No. 41419 CONSULTANT RADIOLOGIST DDRC SRL DIAGNOSTIC (P) LTD. KANNUR