

ECHO REPORT

Name: SRUTHY.A.S	Age/Sex:32Y/F	Date:14/01/2023
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Left Ventricle:-

	Diastole	Systole
IVS	1.06cm	1.20cm
LV	4.09cm	2.61cm
LVPW	1.13cm	1.20cm

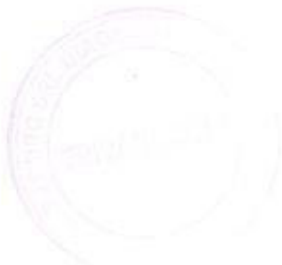
EF - 66% FS - 36%

AO	LA
3.39cm	3.67cm

PV - 1.01m/s
AV - 1.31m/s
MVE - 1.02m/s
MVA - 0.64m/s
E/A - 1.59

IMPRESSION:-

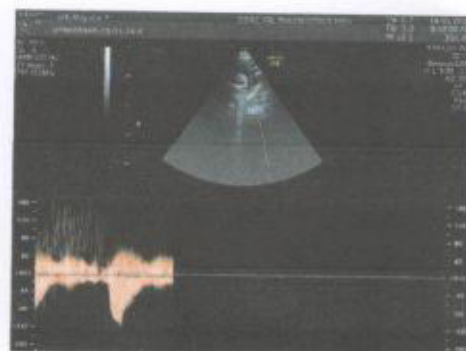
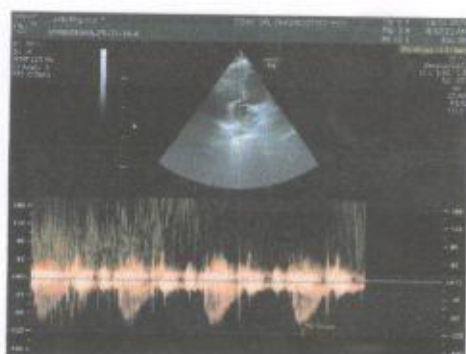
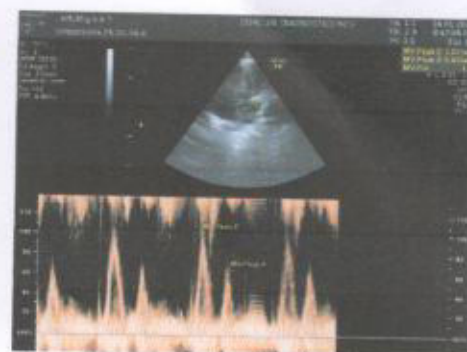
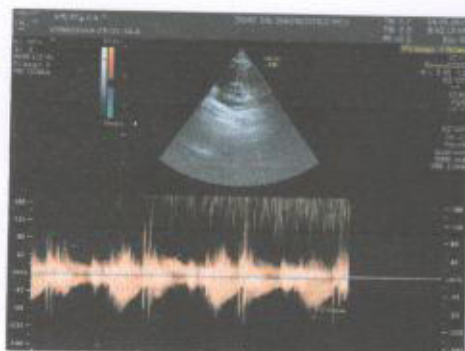
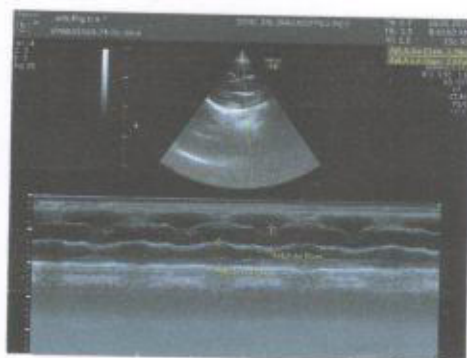
- Normal chambers dimensions
- No RWMA
- Good LV systolic function
- No diastolic dysfunction
- No AS,AR,MR,MS,TR,PAH
- No Vegetation/clot/effusion
- IAS/IVS intact



Consultant Cardiologist

Dr. SRUTHY A.S
M.D.S., Cardiology
Consultant Cardiologist
TUMS Reg No: 73538

COMPLETE IMAGING SOLUTIONS





Patient Ref. No. 66600003020027



Cert. No. MC-2812

CLIENT CODE : CA00010147 - MEDIWHEEL
ARCOFEMI HEALTHCARE LIMITED
CLIENT'S NAME AND ADDRESS :
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F701A, LADO SARAI, NEW DELHI,
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SOUTH DELHI 110030
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Tel : 93334 93334, Fax : CIN - U85190MH2006PTC161480
Email : customercare.ddrc@srl.in

PATIENT NAME : MRS SRUTHY A S PATIENT ID : MRSSF1401914182

ACCESSION NO : 4182WA006506 AGE : 32 Years SEX : Female ABHA NO :

DRAWN : RECEIVED : 14/01/2023 08:19 REPORTED : 16/01/2023 08:20

REFERRING DOCTOR : SELF CLIENT PATIENT ID :

Test Report Status	Results	Biological Reference Interval	Units
Preliminary			

MEDIWHEEL HEALTH CHECKUP BELOW 40(F)2DECHO

OPHTHAL
OPHTHAL REPORT ATTACHED
*** PHYSICAL EXAMINATION**
PHYSICAL EXAMINATION REPORT ATTACHED



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MEDIWHEEL HEALTH CHECKUP BELOW 40(F)2DECHO

* BLOOD UREA NITROGEN (BUN), SERUM

BLOOD UREA NITROGEN 10 Adult(<60 yrs) : 6 to 20 mg/dL

* BUN/CREAT RATIO

BUN/CREAT RATIO 14.5

CREATININE, SERUM

CREATININE 0.66 18 - 60 yrs : 0.6 - 1.1 mg/dL

* GLUCOSE, POST-PRANDIAL, PLASMA

GLUCOSE, POST-PRANDIAL, PLASMA 75 Diabetes Mellitus : > or = 200. mg/dL
Impaired Glucose tolerance/ Prediabetes : 140 - 199. Hypoglycemia : < 55.

GLUCOSE FASTING, FLUORIDE PLASMA

GLUCOSE, FASTING, PLASMA 101 Diabetes Mellitus : > or = 126. mg/dL
Impaired fasting Glucose/ Prediabetes : 101 - 125. Hypoglycemia : < 55.

* GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD

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 102.5 mg/dL

* LIPID PROFILE, SERUM

CHOLESTEROL 184 Desirable : < 200 mg/dL
Borderline : 200-239

TRIGLYCERIDES 46 High : >or= 240 mg/dL
Normal : < 150

HDL CHOLESTEROL 52 High : 150-199 mg/dL
Hypertriglyceridemia : 200-499
Very High : > 499
General range : 40-60





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DIRECT LDL CHOLESTEROL 124 Optimum : < 100 mg/dL
Above Optimum : 100-139
Borderline High : 130-159
High : 160-189
Very High : >or= 190

NON HDL CHOLESTEROL 132 High Desirable: Less than 130 mg/dL
Above Desirable: 130 - 159
Borderline High: 160 - 189
High: 190 - 219
Very high: > or = 220

CHOL/HDL RATIO 3.5 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 2.4 0.5 - 3.0 Desirable/Low Risk
3.1 - 6.0 Borderline/Moderate Risk
>6.0 High Risk

VERY LOW DENSITY LIPOPROTEIN 9.2 Low Desirable value : mg/dL
10 - 35

* LIVER FUNCTION TEST WITH GGT

BILIRUBIN, TOTAL 0.37 General Range : < 1.1 mg/dL

BILIRUBIN, DIRECT 0.14 General Range : < 0.3 mg/dL

BILIRUBIN, INDIRECT 0.23 0.00 - 0.60 mg/dL

TOTAL PROTEIN 7.5 Ambulatory : 6.4 - 8.3 g/dL
Recumbant : 6 - 7.8

ALBUMIN 4.6 20-60yrs : 3.5 - 5.2 g/dL

GLOBULIN 2.9 2.0 - 4.0 g/dL
Neonates -
Pre Mature:
0.29 - 1.04

ALBUMIN/GLOBULIN RATIO 1.6 General Range : 1.1 - 2.5 RATIO

ASPARTATE AMINOTRANSFERASE 10 Adults : < 33 U/L

(AST/SGOT) 8 Adults : < 34 U/L

ALANINE AMINOTRANSFERASE 104 Adult (<60yrs) : 35 - 105 U/L

(ALT/SGPT) 14 Adult (female) : < 40 U/L

ALKALINE PHOSPHATASE 14 Adult (<60yrs) : 35 - 105 U/L

GAMMA GLUTAMYL TRANSFERASE (GGT) 14 Adult (female) : < 40 U/L

TOTAL PROTEIN, SERUM

TOTAL PROTEIN 7.5 Ambulatory : 6.4 - 8.3 g/dL
Recumbant : 6 - 7.8

URIC ACID, SERUM

URIC ACID 3.8 Adults : 2.4-5.7 mg/dL





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ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP TYPE A
RH TYPE POSITIVE

METHOD : COLUMN AGGLUTINATION TECHNOLOGY

BLOOD COUNTS, EDTA WHOLE BLOOD

Table with 4 columns: Test Name, Result, Reference Range, Units. Rows include Hemoglobin, Red Blood Cell Count, White Blood Cell Count, Platelet Count.

Comments

Rechecked

RBC AND PLATELET INDICES

Table with 4 columns: Test Name, Result, Reference Range, Units. Rows include Hematocrit, Mean Corpuscular Vol, Mean Corpuscular Hgb, Mean Corpuscular Hemoglobin Concentration, Red Cell Distribution Width, Mentzer Index, Mean Platelet Volume, WBC Differential Count, Segmented Neutrophils, Lymphocytes, Monocytes, Eosinophils, Basophils, Absolute Neutrophil Count, Absolute Lymphocyte Count, Absolute Monocyte Count.



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ABSOLUTE EOSINOPHIL COUNT	0.19	0.02 - 0.50	thou/ μ L
ABSOLUTE BASOPHIL COUNT	0.0		thou/ μ L
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	1.9		
ERYTHROCYTE SEDIMENTATION RATE (ESR),WHOLE BLOOD			
SEDIMENTATION RATE (ESR)	14	0 - 20	mm at 1 hr
* SUGAR URINE - POST PRANDIAL			
SUGAR URINE - POST PRANDIAL	NOT DETECTED	NOT DETECTED	
* THYROID PANEL, SERUM			
T3	103.90	80 - 200	ng/dL
T4	7.33	5.1 - 14.1	μ g/dl
TSH 3RD GENERATION	1.970	Non-Pregnant : 0.4-4.2	μ IU/mL
		Pregnant Trimester-wise :	
		1st : 0.1 - 2.5	
		2nd : 0.2 - 3	
		3rd : 0.3 - 3	



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Interpretation(s)

Triiodothyronine T3, Thyroxine T4, and Thyroid Stimulating Hormone TSH are thyroid hormones which affect almost every physiological process in the body, including growth, development, metabolism, body temperature, and heart rate. Production of T3 and its prohormone thyroxine (T4) is activated by thyroid-stimulating hormone (TSH), which is released from the pituitary gland. Elevated concentrations of T3, and T4 in the blood inhibit the production of TSH. Excessive secretion of thyroxine in the body is hyperthyroidism, and deficient secretion is called hypothyroidism. In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hyperthyroidism, TSH levels are low. Below mentioned are the guidelines for Pregnancy related reference ranges for Total T4, TSH & Total T3. Measurement of the serum TT3 level is a more sensitive test for the diagnosis of hyperthyroidism, and measurement of TT4 is more useful in the diagnosis of hypothyroidism. Most of the thyroid hormone in blood is bound to transport proteins. Only a very small fraction of the circulating hormone is free and biologically active. It is advisable to detect Free T3, FreeT4 along with TSH, instead of testing for albumin bound Total T3, Total T4.

Table with 6 columns: Sr. No., TSH, Total T4, FT4, Total T3, Possible Conditions. Contains 9 rows of clinical correlations.

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 AMBER
APPEARANCE CLEAR

CHEMICAL EXAMINATION, URINE

PH 5.0 4.7 - 7.5
SPECIFIC GRAVITY 1.020 1.003 - 1.035



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Table with 4 columns: Test Report Status, Preliminary, Results, Units. Rows include PROTEIN, GLUCOSE, KETONES, BLOOD, BILIRUBIN, UROBILINOGEN, NITRITE, MICROSCOPIC EXAMINATION, URINE, RED BLOOD CELLS, WBC, EPITHELIAL CELLS, CASTS, CRYSTALS, REMARKS, SUGAR URINE - FASTING, PHYSICAL EXAMINATION, STOOL, CHEMICAL EXAMINATION, STOOL, MICROSCOPIC EXAMINATION, STOOL.

Interpretation(s)

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.

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 Gravis
Muscular 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 Glycosuria, 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 so that no glucose is excreted in the urine.

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; sulfonyleureas, 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 Glycosuria, 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 HbA1c (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 patient's metabolic control has remained continuously within the target range.
1. eAG (Estimated average glucose) converts percentage HbA1c to mg/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, chronic alcoholism, chronic ingestion of salicylates & opiates addition 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 platform (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 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.
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 syndrome

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, Vasculitides, 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: Polycythemia vera, Sickle cell anemia

LIMITATIONS

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

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

REFERENCE :

1. Nathan and Oski's Haematology of Infancy and Childhood, 5th edition; 2. Paediatric reference intervals. AACCC Press, 7th edition. Edited by S. Soldin; 3. 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

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



Scan to View Details



Scan to View Report



Patient Ref. No. 66600003020027



Cert. No. MC-2812

CLIENT CODE : CA00010147 - MEDIWHEEL
ARCOFEMI HEALTHCARE LIMITED
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
ASTER SQUARE BUILDING, ULLOOR,
MEDICAL COLLEGE P.O
TRIVANDRUM, 695011
KERALA, INDIA
Tel : 93334 93334, Fax : CIN - U85190MH2006PTC161480
Email : customercare.ddrc@srl.in

PATIENT NAME : MRS SRUTHY A S PATIENT ID : MRSSF1401914182

ACCESSION NO : 4182WA006506 AGE : 32 Years SEX : Female ABHA NO :

DRAWN : RECEIVED : 14/01/2023 08:19 REPORTED : 16/01/2023 08:20

REFERRING DOCTOR : SELF CLIENT PATIENT ID :

Test Report Status	Preliminary	Results	Units
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MEDIWHEEL HEALTH CHECKUP BELOW 40(F)2DECHO

* ECG WITH REPORT

REPORT

REPORT ATTACHED

* USG ABDOMEN AND PELVIS

REPORT

REPORT ATTACHED

* CHEST X-RAY WITH REPORT

REPORT

REPORT ATTACHED

* 2D - ECHO WITH COLOR DOPPLER

REPORT

REPORT ATTACHED

****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.

BABU K MATHEW
HOD -BIOCHEMISTRY

DR.VAISHALI RAJAN, MBBS
DCP(Pathology)
(Reg No - TCC 27150)
HOD - HAEMATOLOGY

DR. ASTHA YADAV, MD
Biochemistry
(Reg No - DMC/R/20690)
CONSULTANT BIOCHEMIST

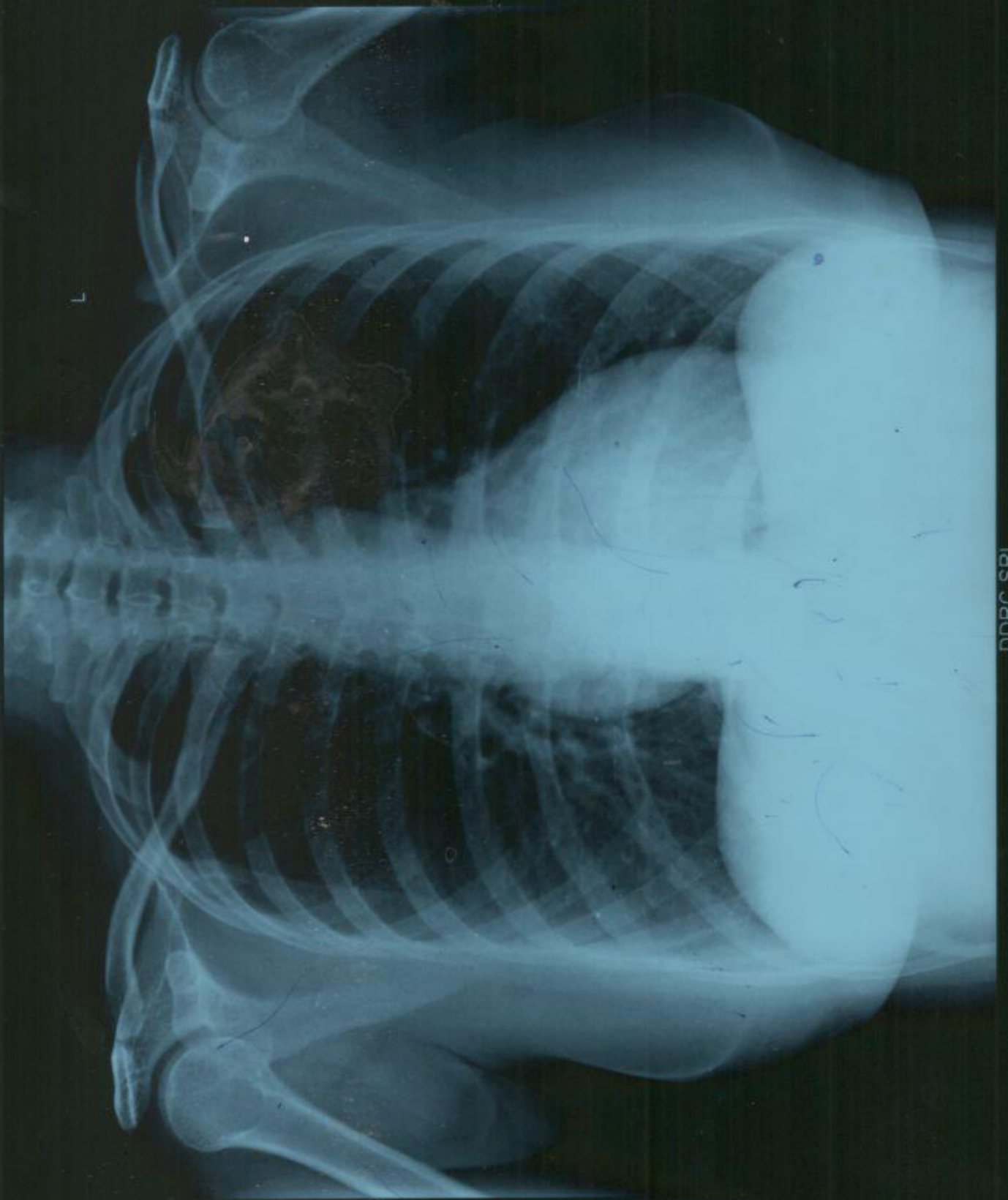
DR NISHA UNNI, MBBS,MD
(RD),DNB (Reg.No:50162)
Consultant Radiologist



Scan to View Details



Scan to View Report



L

DDRC SRL
COLTURY A. S. 2007 E. 4/4/10002 CHEST PA IMA005506

Acc no:4182WA006506	Name: Mrs. Sruthy A S	Age: 32 y	Sex:Female	Date: 14.01.23
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US SCAN WHOLE ABDOMEN (TAS + TVS)

LIVER is normal in size (14.2 cm). Margins are regular. Hepatic parenchyma shows normal echogenicity. No focal lesions seen. No dilatation of intrahepatic biliary radicles. CBD is not dilated. Portal vein is normal in caliber (9.5 mm).

GALL BLADDER is partially distended and grossly normal. No pericholecystic fluid seen.

SPLEEN is normal in size (9.8 cm) and parenchymal echotexture. No focal lesion seen.

PANCREAS Head and body visualized, appears normal in size and parenchymal echotexture. Pancreatic duct is not dilated.

RIGHT KIDNEY is normal in size (10.8 x 4.4 cm) and shows normal parenchymal echotexture. Cortico medullary differentiation is maintained. Parenchymal thickness is normal. No echogenic focus with shadowing suggestive of renal calculi seen. No dilatation of pelvicalyceal system seen. Ureter is not dilated. Perinephric spaces are normal.

LEFT KIDNEY is normal in size (10.2 x 4.8 cm) and shows normal parenchymal echotexture. Cortico medullary differentiation is maintained. Parenchymal thickness is normal. No echogenic focus with shadowing suggestive of renal calculi seen. No dilatation of pelvicalyceal system seen. Ureter is not dilated. Perinephric spaces are normal.

PARAAORTIC AREA No retroperitoneal lymphadenopathy or mass seen.

URINARY BLADDER is distended, normal in wall thickness, lumen clear.

UTERUS measures 8.5 x 4.1 x 5.4 cm, myometrial echopattern normal. No focal lesions seen. Endometrial thickness is 15.6 mm.

Right ovary appears bulky in size, vol - 10.5 cc and shows corpus luteum measuring 2.1 x 1.2 cm. Left ovary appears normal in size, vol - 6.3 cc. **Both ovaries shows multiple peripherally arranged small follicles with central echogenic stroma.** No adnexal mass seen. No fluid in pouch of Douglas. No ascites or pleural effusion.

Gaseous distension of bowel loops noted. No obvious bowel wall thickening seen sonologically.

CONCLUSION:-

- **Bilateral polycystic ovarian morphology, however evidence of ovulation noted from right ovary at present - Suggest clinical & biochemical correlation to rule out PCOS.**



Dr. Nisha Unni MD, DNB (RD)
Consultant radiologist.

Thanks for referral. Your feedback will be appreciated.

(Please bring relevant investigation reports during all visits)

Because of technical and technological limitations complete accuracy cannot be assured on imaging.

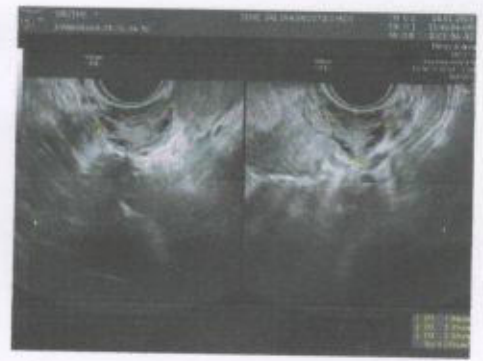
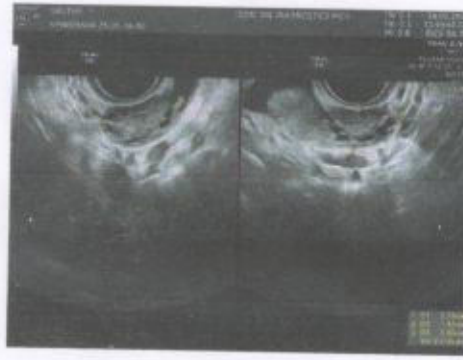
Suggested correlation with clinical findings and other relevant investigations consultations, and if required repeat

imaging recommended in the event of controversies. AR

DDRC SRL Diagnostics Private Limited

Aster Square, Medical College P.O., Trivandrum - 695 011. Ph: 0471 - 2551125. e-mail: info@ddrcsrl.com, web: www.ddrcsrl.com

Corp. Office: DDRC SRL Tower, G-131, Panampilly Nagar, Ernakulam, Kerala - 682 036. Web: www.ddrcsrl.com



V1

V2

V3

V4

ID: 006506

Diagnosis Information:

Female
32 Years
cm

kg
mmHg

MM
Bruckly. A-5

HR : 72 bpm
P : 118 ms
PR : 172 ms
QRS : 90 ms
QT/QTc : 377/413 ms
P/QRS/T : 75/61/46 °
RV5/SV1 : 1.153/0.890 mV

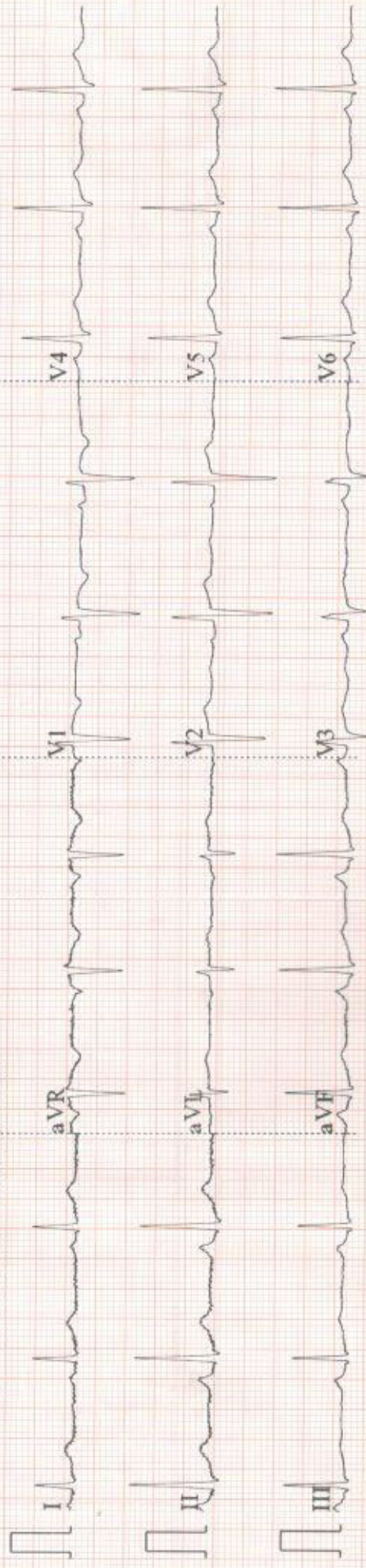
Report Confirmed by:



Standard

Standard	L 1	L II	L III	L III Inspiration

ID: 006506 14-01-2023 10:19:57 AM



0.5~35Hz AC50 25mm/s 10mm/mV ♡74 V1.0 SEMIP V1.7 DDRCSRL

A.M.W CE

TRI

Square
 akkara
 akkara
 'entre
 agappo
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 'kada
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 inkeez
 inkeez
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 vanga
 or
 xpura
 indarr



MEDICAL EXAMINATION REPORT (MER)

Bp: 110/70 mmHg

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	Mr./Mrs./Ms. <u>Soutly, A.S.</u>
2. Mark of Identification	(Mole/Scar/any other (specify location)): <u>Black mole over below lower lip</u>
3. Age/Date of Birth	<u>32 (6/6/1990)</u> , Gender: <u>F/M</u>
4. Photo ID Checked	(Passport/Election Card/PAN Card/Driving Licence/Company ID)

PHYSICAL DETAILS:

a. Height <u>159</u> (cms)	b. Weight <u>57</u> (Kgs)	c. Girth of Abdomen (cms)
d. Pulse Rate (/Min)	e. Blood Pressure: <u>110/70 mmHg</u> Systolic <u>110</u> Diastolic <u>70</u>	
	1 st Reading	
	2 nd Reading	

FAMILY HISTORY:

Relation	Age if Living	Health Status	If deceased, age at the time and cause
Father	<u>64</u>	<u>HTN</u>	
Mother	<u>57</u>	<u>HTN/DLP</u>	
Brother(s)			
Sister(s)			

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

Tobacco in any form	Sedative	Alcohol
<u>No</u>	<u>No</u>	<u>No</u>

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. Y/N
- b. Have you undergone/been advised any surgical procedure? Y/N
- c. During the last 5 years have you been medically examined, received any advice or treatment or admitted to any hospital? Y/N
- d. Have you lost or gained weight in past 12 months? 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? Y/N
- Any Cardiac or Circulatory Disorders? Y/N
- Enlarged glands or any form of Cancer/Tumour? Y/N
- Any Musculoskeletal disorder? Y/N
- Any disorder of Gastrointestinal System? Y/N
- Unexplained recurrent or persistent fever, and/or weight loss Y/N
- Have you been tested for HIV/HBsAg / HCV before? If yes attach reports Y/N
- Are you presently taking medication of any kind? Y/N

• Any disorders of Urinary System?

Y/N ✓

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



FOR FEMALE CANDIDATES ONLY

a. Is there any history of diseases of breast/genital organs?

Y/N ✓

d. Do you have any history of miscarriage/abortion or MTP

Y/N ✓

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

Y/N ✓

c. Do you suspect any disease of Uterus, Cervix or Ovaries?

Y/N ✓

f. Are you now pregnant? If yes, how many months?

Y/N ✓

CONFIDENTIAL COMMENTS FROM MEDICAL EXAMINER

- Was the examinee co-operative? Y/N ✓
- Is there anything about the examinee's health, lifestyle that might affect him/her in the near future with regard to his/her job? Y/N ✓
- Are there any points on which you suggest further information be obtained? Y/N ✓
- Based on your clinical impression, please provide your suggestions and recommendations below:

.....
.....

➤ Do you think he/she is MEDICALLY FIT or UNFIT for employment.

MEDICAL EXAMINER'S DECLARATION

I hereby confirm that I have examined the above individual 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 : **Dr. S. S. LOPEZ, MBBS**
MEDICAL OFFICER
DDRC SRL Diagnostics Ltd.

Seal of Medical Examiner : Aster Square, Medical College P.O., TVM



Name & Seal of DDRC SRL Branch

Date & Time : 17/10/2023

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

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

NAME : MRS SRUTHY A S	AGE:32/F	DATE:14/01/2023
-----------------------	----------	-----------------

CHEST X-RAY REPORT

CHEST X-RAY PA VIEW : Trachea central
 No cardiomegaly
 Normal vascularity
 No parenchymal lesion.
 Costophrenic and cardiophrenic angles clear

➤ **IMPRESSION** : Normal Chest Xray

ELECTRO CARDIOGRAM : NSR:72/minute
 No evidence of ischaemia.

➤ **IMPRESSION** : Normal Ecg.



Serin Lopez
 Dr. SERIN LOPEZ, MBBS
 MEDICAL OFFICER
 DDRC SRL Diagnostics Ltd.
 Aster Square, Medical College P.O., TVM
 Reg. No. 77656

DR SERIN LOPEZ MBBS

Reg No 77656

DDRC SRL DIAGNOSTICS LTD