



CLIENT CODE : C000138381

CLIENT'S NAME AND ADDRESS :
ACROFEMI HEALTHCARE LTD (MEDIWHEEL)
F-703, LADO SARAI, MEHRAULI
SOUTH WEST DELHI
NEW DELHI 110030
DELHI INDIA
8800465156

SRL Ltd
SRL Wellness Centre, SCO. 13, Sector 16 Market, Faridabad
FARIDABAD, 121001
Haryana, INDIA
Tel : 9111591115, Fax :
CIN - U74899PB1995PLC045956

PATIENT NAME : ANU KHANDELWAL

PATIENT ID : ANUKF10038771

ACCESSION NO : 0071VG000271 AGE : 35 Years SEX : Female

DRAWN :

RECEIVED : 09/07/2022 10:21

REPORTED : 12/07/2022 09:47

REFERRING DOCTOR : SELF

CLIENT PATIENT ID :

Test Report Status	Preliminary	Results	Biological Reference Interval	Units
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MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE**BLOOD COUNTS,EDTA WHOLE BLOOD**

HEMOGLOBIN	11.8	Low 12.0 - 15.0	g/dL
METHOD : SPECTROPHOTOMETRY			
RED BLOOD CELL COUNT	4.08	3.8 - 4.8	mil/ μ L
METHOD : IMPEDANCE			
WHITE BLOOD CELL COUNT	7.36	4.0 - 10.0	thou/ μ L
METHOD : IMPEDANCE			
PLATELET COUNT	257	150 - 410	thou/ μ L
METHOD : IMPEDANCE			

RBC AND PLATELET INDICES

HEMATOCRIT	38.6	36 - 46	%
METHOD : CALCULATED			
MEAN CORPUSCULAR VOL	94.6	83 - 101	fL
METHOD : DERIVED FROM IMPEDANCE MEASURE			
MEAN CORPUSCULAR HGB.	28.9	27.0 - 32.0	pg
METHOD : CALCULATED PARAMETER			
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION	30.5	Low 31.5 - 34.5	g/dL
METHOD : CALCULATED PARAMETER			
MENTZER INDEX	23.2		
RED CELL DISTRIBUTION WIDTH	17.0	High 11.6 - 14.0	%
METHOD : DERIVED FROM IMPEDANCE MEASURE			
MEAN PLATELET VOLUME	11.3	High 6.8 - 10.9	fL
METHOD : DERIVED FROM IMPEDANCE MEASURE			

WBC DIFFERENTIAL COUNT - NLR

SEGMENTED NEUTROPHILS	63	40 - 80	%
METHOD : DHSS FLOWCYTOMETRY			
ABSOLUTE NEUTROPHIL COUNT	4.60	2.0 - 7.0	thou/ μ L
METHOD : DHSS FLOWCYTOMETRY, CALCULATED			
LYMPHOCYTES	26	20 - 40	%
METHOD : DHSS FLOWCYTOMETRY			
ABSOLUTE LYMPHOCYTE COUNT	1.90	1 - 3	thou/ μ L
METHOD : DHSS FLOWCYTOMETRY, CALCULATED			
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	2.4		
METHOD : CALCULATED			
EOSINOPHILS	3	1 - 6	%





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METHOD : DHSS FLOWCYTOMETRY				
ABSOLUTE EOSINOPHIL COUNT		0.19	0.02 - 0.50	thou/ μ L
METHOD : DHSS FLOWCYTOMETRY, CALCULATED				
MONOCYTES		08	2 - 10	%
METHOD : DHSS FLOWCYTOMETRY				
ABSOLUTE MONOCYTE COUNT		0.59	0.20 - 1.00	thou/ μ L
METHOD : DHSS FLOWCYTOMETRY, CALCULATED				
BASOPHILS		0	0 - 2	%
METHOD : IMPEDANCE				
ABSOLUTE BASOPHIL COUNT		0.01	Low 0.02 - 0.10	thou/ μ L
METHOD : DHSS FLOWCYTOMETRY, CALCULATED				
ERYTHRO SEDIMENTATION RATE, BLOOD				
SEDIMENTATION RATE (ESR)		41	High 0 - 20	mm at 1 hr
METHOD : AUTOMATED (PHOTOMETRICAL CAPILLARY STOPPED FLOW KINETIC ANALYSIS)				
GLUCOSE, FASTING, PLASMA				
GLUCOSE, FASTING, PLASMA		80	Normal 75 - 99 Pre-diabetics: 100 - 125 Diabetic: > or = 126	mg/dL
METHOD : SPECTROPHOTOMETRY HEXOKINASE				
GLYCOSYLATED HEMOGLOBIN, EDTA WHOLE BLOOD				
GLYCOSYLATED HEMOGLOBIN (HBA1C)		5.0	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 ADA Target: 7.0 Action suggested: > 8.0	%
METHOD : CAPILLARY ELECTROPHORESIS				
MEAN PLASMA GLUCOSE		96.8	< 116	mg/dL
METHOD : CALCULATED PARAMETER				
GLUCOSE, POST-PRANDIAL, PLASMA				
GLUCOSE, POST-PRANDIAL, PLASMA		95	70 - 139	mg/dL
METHOD : SPECTROPHOTOMETRY, HEXOKINASE				
CORONARY RISK PROFILE (LIPID PROFILE), SERUM.				
CHOLESTEROL		187	Desirable cholesterol level < 200 Borderline high cholesterol 200 - 239 High cholesterol > / = 240	mg/dL
METHOD : ENZYMATIC COLORIMETRIC ASSAY				





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TRIGLYCERIDES		77	Normal: < 150 Borderline high: 150 - 199 High: 200 - 499 Very High: >/= 500	mg/dL
HDL CHOLESTEROL		51	<40 = 60	mg/dL
DIRECT LDL CHOLESTEROL		125.00	Optimal : < 100 Near optimal/above optimal : 100 - 129 Borderline high : 130 - 159 High : 160 - 189 Very high : > / = 190	mg/dL
NON HDL CHOLESTEROL		136		mg/dL
CHOL/HDL RATIO		3.7		
LDL/HDL RATIO		2.5		
VERY LOW DENSITY LIPOPROTEIN		15.4		mg/dL
LIVER FUNCTION PROFILE, SERUM				
BILIRUBIN, TOTAL		0.4	Upto 1.2	mg/dL
BILIRUBIN, DIRECT		0.2	< 0.30	mg/dL
METHOD : DIAZO METHOD				
BILIRUBIN, INDIRECT		0.20	0.1 - 1.0	mg/dL
TOTAL PROTEIN		7.2	6.0 - 8.0	g/dL
METHOD : SPECTROPHOTOMETRY, BIURET				
ALBUMIN		4.4	3.97 - 4.94	g/dL
METHOD : SPECTROPHOTOMETRY, BCP - DYE BINDING				
GLOBULIN		2.8	2.0 - 3.5	g/dL
METHOD : CALCULATED PARAMETER				
ALBUMIN/GLOBULIN RATIO		1.6		RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT)		26	< OR = 35	U/L
METHOD : SPECTROPHOTOMETRY, WITHOUT PYRIDOXAL PHOSPHATE ACTIVATION(P5P) - IFCC				
ALANINE AMINOTRANSFERASE (ALT/SGPT)		22	< OR = 35	U/L
METHOD : SPECTROPHOTOMETRY, WITHOUT PYRIDOXAL PHOSPHATE ACTIVATION(P5P) - IFCC				
ALKALINE PHOSPHATASE		90		U/L
METHOD : SPECTROPHOTOMETRY				
GAMMA GLUTAMYL TRANSFERASE (GGT)		15	0 - 40	U/L
METHOD : ENZYMATIC COLORIMETRIC ASSAY				
LACTATE DEHYDROGENASE		184	125 - 220	U/L
METHOD : SPECTROPHOTOMETRY, LACTATE TO PYRUVATE - UV-IFCC				

SERUM BLOOD UREA NITROGEN

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BLOOD UREA NITROGEN		7.6	6 - 20	mg/dL
METHOD : SPECTROPHOTOMETRY, KINETIC ENZYMATIC				
CREATININE, SERUM				
CREATININE		0.40	Low 0.5 - 0.9	mg/dL
METHOD : SPECTROPHOTOMETRIC, JAFFE'S KINETICS				
BUN/CREAT RATIO				
BUN/CREAT RATIO		21.50	High 8.0 - 15.0	
METHOD : CALCULATED PARAMETER				
URIC ACID, SERUM				
URIC ACID		3.7	2.4 - 5.7	mg/dL
METHOD : SPECTROPHOTOMETRY				
NOTE : THIS TEST WAS DEVELOPED AND ITS PERFORMANCE CHARACTERISTICS WERE DETERMINED BY HAEMATOLOGY SECTION, SRL LIMITED, MUMBAI. THIS TEST IS USED FOR CLINICAL PURPOSE AND SHOULD NOT BE REGARDED AS INVESTIGATIONAL OR FOR RESEARCH.				
TOTAL PROTEIN, SERUM				
TOTAL PROTEIN		7.2	6.0 - 8.0	g/dL
ALBUMIN, SERUM				
ALBUMIN		4.4	3.97 - 4.94	g/dL
METHOD : SPECTROPHOTOMETRY, BCP - DYE BINDING				
GLOBULIN				
GLOBULIN		2.8	2.0 - 3.5	g/dL
METHOD : CALCULATED PARAMETER				
ELECTROLYTES (NA/K/CL), SERUM				
SODIUM		140	136 - 145	mmol/L
METHOD : ISE DIRECT				
POTASSIUM		4.5	High 3.4 - 4.4	mmol/L
METHOD : ISE DIRECT				
CHLORIDE		103		mmol/L
PHYSICAL EXAMINATION, URINE				
COLOR		PALE YELLOW		
APPEARANCE		CLEAR		
SPECIFIC GRAVITY		1.020	1.003 - 1.035	

Comments

NOTE : MICROSCOPIC EXAMINATION OF URINE IS PERFORMED ON CENTRIFUGED URINARY SEDIMENT. IN NORMAL URINE SAMPLES CAST AND CRYSTALS ARE NOT DETECTED.

CHEMICAL EXAMINATION, URINE

PH 6.0 4.7 - 7.5



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PROTEIN		NOT DETECTED	NOT DETECTED	
GLUCOSE		NOT DETECTED	NOT DETECTED	
KETONES		NOT DETECTED	NOT DETECTED	
BLOOD		NOT DETECTED	NOT DETECTED	
BILIRUBIN		NOT DETECTED	NOT DETECTED	
UROBILINOGEN		NORMAL	NORMAL	
NITRITE		NOT DETECTED	NOT DETECTED	

MICROSCOPIC EXAMINATION, URINE

PUS CELL (WBC'S)	1-2	0-5	/HPF
EPITHELIAL CELLS	2-3	0-5	/HPF
ERYTHROCYTES (RBC'S)	NOT DETECTED	NOT DETECTED	/HPF
CASTS	NOT DETECTED		
CRYSTALS	NOT DETECTED		
BACTERIA	NOT DETECTED	NOT DETECTED	

METHOD : DIP STICK/MICRO SCOPY/REFLECTANCE SPECTROPHOTOMETRY

THYROID PANEL, SERUM

T3	116.0	80 - 200	ng/dL
METHOD : ELECTROCHEMILUMINESCENCE IMMUNO ASSAY			
T4	6.40	5.1 - 14.1	µg/dL
METHOD : ELECTROCHEMILUMINESCENCE IMMUNO ASSAY			
TSH 3RD GENERATION	1.840	0.27 - 4.2	µIU/mL
METHOD : ELECTROCHEMILUMINESCENCE IMMUNO ASSAY			

PAPANICOLAOU SMEAR

RESULT PENDING

LETTER

RESULT PENDING

STOOL: OVA & PARASITE

REMARK

TEST CANCELLED AS SPECIMEN NOT RECEIVED

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP

B

RH TYPE

RH-

XRAY-CHEST

>>>

BOTH THE LUNG FIELDS ARE CLEAR

>>>

BOTH THE COSTOPHRENIC AND CARIOPHRENIC ANGLES ARE CLEAR

>>>

BOTH THE HILA ARE NORMAL

>>>

CARDIAC AND AORTIC SHADOWS APPEAR NORMAL

>>>

BOTH THE DOMES OF THE DIAPHRAM ARE NORMAL



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»»	VISUALIZED BONY THORAX IS NORMAL		
IMPRESSION	NO ABNORMALITY DETECTED		
TMT OR ECHO			
TMT OR ECHO	REPORT ENCLOSED		
ECG			
ECG	WITHIN NORMAL LIMITS		
MEDICAL HISTORY			
RELEVANT PRESENT HISTORY	NOT SIGNIFICANT		
RELEVANT PAST HISTORY	PULMONARY EMBOLISM 2011 , 2018		
RELEVANT PERSONAL HISTORY	WIDOW, 1 CHILD. VEGETERIAN		
MENSTRUAL HISTORY (FOR FEMALES)	REGULAR		
LMP (FOR FEMALES)	19.06.2022		
OBSTETRIC HISTORY (FOR FEMALES)	G1P1		
LCB (FOR FEMALES)	27.02.2011		
RELEVANT FAMILY HISTORY	NOT SIGNIFICANT		
OCCUPATIONAL HISTORY	B.COM PGDC		
HISTORY OF MEDICATIONS	NOT SIGNIFICANT		
ANTHROPOMETRIC DATA & BMI			
HEIGHT IN METERS	1.41		mts
WEIGHT IN KGS.	60		Kgs
BMI	30		

BMI & Weight Status as follows: kg/sqmts
 Below 18.5: Underweight
 18.5 - 24.9: Normal
 25.0 - 29.9: Overweight
 30.0 and Above: Obese

GENERAL EXAMINATION

MENTAL / EMOTIONAL STATE	NORMAL
PHYSICAL ATTITUDE	NORMAL
GENERAL APPEARANCE / NUTRITIONAL STATUS	HEALTHY
BUILT / SKELETAL FRAMEWORK	AVERAGE
FACIAL APPEARANCE	NORMAL
SKIN	NORMAL
UPPER LIMB	NORMAL
LOWER LIMB	NORMAL
NECK	NORMAL
NECK LYMPHATICS / SALIVARY GLANDS	NOT ENLARGED OR TENDER





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THYROID GLAND		NOT ENLARGED		
CAROTID PULSATION		NORMAL		
BREAST (FOR FEMALES)		NORMAL		
TEMPERATURE		NORMAL		
PULSE		106 MIN/REGULAR, ALL PERIPHERAL PULSES WELL FELT		
RESPIRATORY RATE		NORMAL		
CARDIOVASCULAR SYSTEM				
BP		140/96 MM HG (SITTING)		mm/Hg
PERICARDIUM		NORMAL		
APEX BEAT		NORMAL		
HEART SOUNDS		S1, S2 HEARD NORMALLY		
MURMURS		ABSENT		
RESPIRATORY SYSTEM				
SIZE AND SHAPE OF CHEST		NORMAL		
MOVEMENTS OF CHEST		SYMMETRICAL		
BREATH SOUNDS INTENSITY		NORMAL		
BREATH SOUNDS QUALITY		VESICULAR (NORMAL)		
ADDED SOUNDS		ABSENT		
PER ABDOMEN				
APPEARANCE		NORMAL		
VENOUS PROMINENCE		ABSENT		
LIVER		NOT PALPABLE		
SPLEEN		NOT PALPABLE		
HERNIA		ABSENT		
CENTRAL NERVOUS SYSTEM				
HIGHER FUNCTIONS		NORMAL		
CRANIAL NERVES		NORMAL		
CEREBELLAR FUNCTIONS		NORMAL		
SENSORY SYSTEM		NORMAL		
MOTOR SYSTEM		NORMAL		
REFLEXES		NORMAL		
MUSCULOSKELETAL SYSTEM				
SPINE		NORMAL		
JOINTS		NORMAL		





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BASIC EYE EXAMINATION

CONJUNCTIVA	NORMAL
EYELIDS	NORMAL
EYE MOVEMENTS	NORMAL
CORNEA	NORMAL
DISTANT VISION RIGHT EYE WITHOUT GLASSES	6/6
DISTANT VISION LEFT EYE WITHOUT GLASSES	6/6

BASIC ENT EXAMINATION

EXTERNAL EAR CANAL	NORMAL
TYMPANIC MEMBRANE	NORMAL
NOSE	NO ABNORMALITY DETECTED
SINUSES	CLEAR
THROAT	NO ABNORMALITY DETECTED
TONSILS	NOT ENLARGED

SUMMARY

RELEVANT HISTORY	NOT SIGNIFICANT
RELEVANT NON PATHOLOGY DIAGNOSTICS	NO ABNORMALITIES DETECTED

FITNESS STATUS

FITNESS STATUS	FIT (WITH MEDICAL ADVICE) (AS PER REQUESTED PANEL OF TESTS)
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Comments

OUR PANEL OF DOCTORS.
GENERAL PHYSICIAN - DR. MUKUL GOSWAMI
CONSULTANT RADIOLOGIST - DR. D.R. CHUGH
CONSULTANT CARDIOLOGIST : DR. SANDEEP KUMAR
CONSULTANT GYNAECOLOGIST : DR. KAVITA

THIS REPORT CARRIES THE SIGNATURE OF OUR LABORATORY DIRECTOR.
THIS IS AN INVIOABLE FEATURE OF OUR LAB MANAGEMENT SOFTWARE.
HOWEVER, ALL EXAMINATION AND INVESTIGATIONS HAVE BEEN
CONDUCTED BY OUR PANEL OF DOCTORS

Interpretation(s)

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



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diagnosing a case of beta thalassaemia trait.

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-

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 polikilocytosis, spherocytosis or sickle cells.

Reference :

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

GLUCOSE, FASTING, PLASMA-

ADA 2021 guidelines for adults, after 8 hrs fasting is as follows:

Pre-diabetics: 100 - 125 mg/dL

Diabetic: > or = 126 mg/dL

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 glycosylated 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 glycosylated 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 glycosylated 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, 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.

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).





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FARIDABAD, 121001
Haryana, INDIA
Tel : 9111591115, Fax :
CIN - U74899PB1995PLC045956

PATIENT NAME : ANU KHANDELWAL

PATIENT ID : ANUKF10038771

ACCESSION NO : 0071VG000271 AGE : 35 Years SEX : Female

DRAWN : RECEIVED : 09/07/2022 10:21 REPORTED : 12/07/2022 09:47

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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.

LIVER FUNCTION PROFILE, SERUM-LIVER FUNCTION PROFILE

Bilirubin is a yellowish pigment found in bile and is a breakdown product of normal heme catabolism. Bilirubin is excreted in bile and urine, and elevated levels may give yellow discoloration in jaundice. Elevated levels result from increased bilirubin production (eg, hemolysis and ineffective erythropoiesis), decreased bilirubin excretion (eg, obstruction and hepatitis), and abnormal bilirubin metabolism (eg, hereditary and neonatal jaundice). Conjugated (direct) bilirubin is elevated more than unconjugated (indirect) bilirubin in Viral hepatitis, Drug reactions, Alcoholic liver disease. Conjugated (direct) bilirubin is also elevated more than unconjugated (indirect) bilirubin when there is some kind of blockage of the bile ducts like in Gallstones getting into the bile ducts, tumors & Scarring of the bile ducts. Increased unconjugated (indirect) bilirubin may be a result of Hemolytic or pernicious anemia, Transfusion reaction & a common metabolic condition termed Gilbert syndrome, due to low levels of the enzyme that attaches sugar molecules to bilirubin.

AST is an enzyme found in various parts of the body. AST is found in the liver, heart, skeletal muscle, kidneys, brain, and red blood cells, and it is commonly measured clinically as a marker for liver health. AST levels increase during chronic viral hepatitis, blockage of the bile duct, cirrhosis of the liver, liver cancer, kidney failure, hemolytic anemia, pancreatitis, hemochromatosis. AST levels may also increase after a heart attack or strenuous activity. ALT test measures the amount of this enzyme in the blood. ALT is found mainly in the liver, but also in smaller amounts in the kidneys, heart, muscles, and pancreas. It is commonly measured as a part of a diagnostic evaluation of hepatocellular injury, to determine liver health. AST levels increase during acute hepatitis, sometimes due to a viral infection, ischemia to the liver, chronic hepatitis, obstruction of bile ducts, cirrhosis.

ALP is a protein found in almost all body tissues. Tissues with higher amounts of ALP include the liver, bile ducts and bone. Elevated ALP levels are seen in Biliary obstruction, Osteoblastic bone tumors, osteomalacia, hepatitis, Hyperparathyroidism, Leukemia, Lymphoma, Paget's disease, Rickets, Sarcoidosis etc. Lower-than-normal ALP levels seen in Hypophosphatasia, Malnutrition, Protein deficiency, Wilson's disease. GGT is an enzyme found in cell membranes of many tissues mainly in the liver, kidney and pancreas. It is also found in other tissues including intestine, spleen, heart, brain and seminal vesicles. The highest concentration is in the kidney, but the liver is considered the source of normal enzyme activity. Serum GGT has been widely used as an index of liver dysfunction. Elevated serum GGT activity can be found in diseases of the liver, biliary system and pancreas. Conditions that increase serum GGT are obstructive liver disease, high alcohol consumption and use of enzyme-inducing drugs etc. 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. Human serum albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by: Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance, malnutrition and wasting etc

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

- 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

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.



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Tel : 9111591115, Fax :
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PATIENT ID : ANUKF10038771

ACCESSION NO : 0071VG000271 AGE : 35 Years SEX : Female

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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

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.

ALBUMIN, SERUM-

Human serum albumin is the most abundant protein in human blood plasma. It is produced in the liver. Albumin constitutes about half of the blood serum protein. Low blood albumin levels (hypoalbuminemia) can be caused by: Liver disease like cirrhosis of the liver, nephrotic syndrome, protein-losing enteropathy, Burns, hemodilution, increased vascular permeability or decreased lymphatic clearance, malnutrition and wasting etc.

ELECTROLYTES (NA/K/CL), SERUM-

Sodium levels are Increased in dehydration, Cushing's syndrome, aldosteronism & decreased in Addison's disease, hypopituitarism, liver disease. Hypokalemia (low K) is common in vomiting, diarrhea, alcoholism, folic acid deficiency and primary aldosteronism. Hyperkalemia may be seen in end-stage renal failure, hemolysis, trauma, Addison's disease, metabolic acidosis, acute starvation, dehydration, and with rapid K infusion. Chloride is increased in dehydration, renal tubular acidosis (hyperchloremia metabolic acidosis), acute renal failure, metabolic acidosis associated with prolonged diarrhea and loss of sodium bicarbonate, diabetes insipidus, adrenocortical hyperfunction, salicylate intoxication and with excessive infusion of isotonic saline or extremely high dietary intake of salt. Chloride is decreased in overhydration, chronic respiratory acidosis, salt-losing nephritis, metabolic alkalosis, congestive heart failure, Addisonian crisis, certain types of metabolic acidosis, persistent gastric secretion and prolonged vomiting,

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

THYROID PANEL, SERUM-

Triiodothyronine 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 T ₄ (µg/dL)	TSH3G (µIU/mL)	TOTAL T ₃ (ng/dL)
Pregnancy			
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 age related reference ranges for T₃ and T₄.

	T ₃ (ng/dL)	T ₄ (µg/dL)
New Born:	75 - 260	1-3 day: 8.2 - 19.9



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 CIN - U74899PB1995PLC045956

PATIENT NAME : ANU KHANDELWAL

PATIENT ID : ANUKF10038771

ACCESSION NO : 0071VG000271 **AGE :** 35 Years **SEX :** Female

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1 Week: 6.0 - 15.9

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.
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

STOOL: OVA & PARASITE-

Acute infective diarrhoea and gastroenteritis (diarrhoea with vomiting) are major causes of ill health and premature death in developing countries. Loss of water and electrolytes from the body can lead to severe dehydration which if untreated, can be rapidly fatal in young children, especially that are malnourished, hypoglycaemic, and generally in poor health.

Laboratory diagnosis of parasitic infection is mainly based on microscopic examination and the gross examination of the stool specimen. Depending on the nature of the parasite, the microscopic observations include the identification of cysts, ova, trophozoites, larvae or portions of adult structure. The two classes of parasites that cause human infection are the Protozoa and Helminths. The protozoan infections include amoebiasis mainly caused by Entamoeba histolytica and giardiasis caused by Giardia lamblia. The common helminthic parasites are Trichuris trichiura, Ascaris lumbricoides, Strongyloides stercoralis, Taenia sp. etc

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.

MEDICAL

HISTORY-*****
 THIS REPORT CARRIES THE SIGNATURE OF OUR LABORATORY DIRECTOR. THIS IS AN INVOLABLE FEATURE OF OUR LAB MANAGEMENT SOFTWARE. HOWEVER, ALL EXAMINATIONS AND INVESTIGATIONS HAVE BEEN CONDUCTED BY OUR PANEL OF DOCTORS.

FITNESS STATUS-

Conclusion on an individual's Fitness, which is commented upon mainly for Pre employment cases, is based on multi factorial findings and does not depend on any one single parameter. The final Fitness assigned to a candidate will depend on the Physician's findings and overall judgement on a case to case basis, details of the candidate's past and personal history; as well as the comprehensiveness of the diagnostic panel which has been requested for .These are then further correlated with details of the job under consideration to eventually fit the right man to the right job.

Basis the above, SRL classifies a candidate's Fitness Status into one of the following categories:

- Fit (As per requested panel of tests) – SRL Limited gives the individual a clean chit to join the organization, on the basis of the General Physical Examination and the specific test panel requested for.
- Fit (with medical advice) (As per requested panel of tests) - This indicates that although the candidate can be declared as FIT to join the job, minimal problems have been detected during the Pre- employment examination. Examples of conditions which could fall in this category could be cases of mild reversible medical abnormalities such as height weight disproportions, borderline raised Blood Pressure readings, mildly raised Blood sugar and Blood Lipid levels, Hematuria, etc. Most of these relate to sedentary lifestyles and come under the broad category of life style disorders. The idea is to caution an individual to bring about certain lifestyle changes as well as seek a Physician's consultation and counseling in order to bring back to normal the mildly deranged parameters. For all purposes the individual is FIT to join the job.
- Fitness on Hold (Temporary Unfit) (As per requested panel of tests) - Candidate's reports are kept on hold when either the diagnostic tests or the physical findings reveal the presence of a medical condition which warrants further tests, counseling and/or specialist opinion, on the basis of which a candidate can either be placed into Fit, Fit (With Medical Advice), or Unfit category. Conditions which may fall into this category could be high blood pressure, abnormal ECG, heart murmurs, abnormal vision, grossly elevated blood sugars, etc.
- Unfit (As per requested panel of tests) - An unfit report by SRL Limited clearly indicates that the individual is not suitable for the respective job profile e.g. total color blindness in color related jobs.





Patient Ref. No. 7100000302444

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MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE**ULTRASOUND ABDOMEN****ULTRASOUND ABDOMEN**

REPORT ENCLOSED

****End Of Report****Please visit www.srlworld.com for related Test Information for this accession

Dr. Arpita Roy, MD
Section Head-Hematology

Dr. Mamta Kumari, MBBS, MD
Consultant Microbiologist

Dr. Chandan Hazarika
Microbiologist

Dr. Geeta
Pathologist

CONDITIONS OF LABORATORY TESTING & REPORTING

1. It is presumed that the test sample belongs to the patient named or identified in the test requisition form.
2. All Tests are performed and reported as per the turnaround time stated in the SRL Directory of services (DOS).
3. SRL confirms that all tests have been performed or assayed with highest quality standards, clinical safety & technical integrity.
4. A requested test might not be performed if:
 - a. Specimen received is insufficient or inappropriate specimen quality is unsatisfactory
 - b. Incorrect specimen type
 - c. Request for testing is withdrawn by the ordering doctor or patient
 - d. There is a discrepancy between the label on the specimen container and the name on the test requisition form
5. The results of a laboratory test are dependent on the quality of the sample as well as the assay technology.
6. Result delays could be because of uncontrolled circumstances. e.g. assay run failure.
7. Tests parameters marked by asterisks are excluded from the "scope" of NABL accredited tests. (If laboratory is accredited).
8. Laboratory results should be correlated with clinical information to determine Final diagnosis.
9. Test results are not valid for Medico- legal purposes.
10. In case of queries or unexpected test results please call at SRL customer care (Toll free: 1800-222-000). Post proper investigation repeat analysis may be carried out.

SRL Limited

Fortis Hospital, Sector 62, Phase VIII,
Mohali 160062



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SRL DIAGNOSTICS WELLNESS CENTER
SCO 13, SECTOR 16 FARIDABAD
PHONE NO- 0129-4179185

NAME :- MS. ANU KHANDELWAL	AGE /SEX/35/YEARS/F
ACC:- 0071VG000371	DATE :- 11/06/2022

ULTRA SOUND SCAN OF WHOLE ABDOMEN

Liver: Normal in size, and shows homogeneous echotexture. No obvious focal or diffuse pathology is noted in either of the lobes. Fatty changes present in the liver of grade I
Hepatic veins appear normal

Gall bladder: ABSENT

CBD AND PORTALVEIN : normal in caliber

Pancreas : Normal in size shape and echotexture no e/o focal lesion /calcification. Pancreatic duct appears normal.

Spleen: Normal in size, shape and Echotexture. No e/o focal lesion

Both Kidneys: Both kidneys are normal in size and echotexture No e/o hydronephrosis/focal lesion

Urinary bladder: Well distended. No e/o calculi/internal echoes. Wall thickness appears normal.

UTERUS: Normal in size, Shape, position and echotexture. Endometrial cavity is central empty. No focal myometrial lesion seen.

OVARIES: Both the ovaries are normal in size, Position and echotexture
No adnexal mass is seen
There is no free fluid seen in the cul-de-sac

IMPRESSION:- WHOLE ABDOMEN REVEALS FATTY CHANGES IN THE LIVER .

Correlate with clinically findings.

Dr. D.R. CHUGH
(MBBS, DMRD)
SRL LIMITED
SCO-13, Sec 16, HUDA Market,
Faridabad-11002
Tel: 0129-4179184/85

DR. D.R CHUGH
(RADIOLOGIST)

Disclaimer:

The science of ultrasound is based upon interpretation of moving shadows of normal and abnormal tissue. This is neither complete nor accurate, hence findings should always be interpreted in to the light of clinico-pathological correlation. This a professional opinion, not a diagnosis. Not meant for medico legal purpose.

NAME :- MRS. ANU KHANDELWAL	Age/ Sex/35/Years/F
ACC:- 0071VG000271	Date :- 09/07/2022

X-RAY CHEST PA VIEW

- ❖ Both lung fields are normal.
- ❖ Both costophrenic angles are normal.
- ❖ Both domes of diaphragm are normal.
- ❖ Both hilar shadow are normal.
- ❖ Cardiac size is normal.
- ❖ Visualized soft tissues & thoracic cage are normal.
- ❖ **IMPRESSION :**

Please Correlate Clinically.

Wedge in base in the clud

Dr. D.R. CHUGH
(MBBS, DMRD)
SRL LIMITED
SCO-13, Sec-16, HUDA Market,
Faridkot-142002
Tel. 0126-4179184/85
Dr. D.R CHUGH
(RADIOLOGIST)

Disclaimer:

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2D ECHOCARDIOGRAPHY REPORT

NAME :- MS. ANU KHANDELWAL
ACC:- 0071VG000271

AGE/SEX/35/YEARS/F
DATE : 09/06/2022

OBSERVATIONS BY M- MODE & 2D ECHOCARDIOGRAPHY.

LEFT Ventricle		Ed	Es
AO	(mm)	27	
IVS	(mm)	09	12
L V ID	(mm)	28	20
Left Vent Post . Wall Thickness	(mm)	09	11
LA	(mm)	34	
LVEF		60 %	
Aortic Root Diameter		Normal	
RIGHT Ventricle		Normal	
MITRAL VALVE		Normal	
AROTIC VALVE		Normal	
TRICUSPID VALVE		Normal	
PULMONARY VALVE		Normal	
PERICARDIUM		Normal	
2D STUDY of wall motion			
RIGHT Ventricle		Normal	
LEFT Ventricle		No RWMA	

DOPLER STUDY

MITRAL	Grade 1 DRA
AORTIC	NORMAL
TRICUSPID	NORMAL
PULMONARY	NORMAL

COLOUR FLOW MAPPING

No Valvular stenosis / Trivial Mitral Regurgitation.
Trivial Tricuspid Regurgitation .

CONCLUSION

No Regional wall motion abnormality
Normal cardiac chamber dimensions.
Normal LV Systolic Function.
Grade 1 DRA
Normal RV Size and function
No Valvular stenosis./ Trivial Mitral Regurgitation.
Trivial Tricuspid Regurgitation. PASP 20 mm of Hg
No intracardiac Mass/ clot
IVC Collapsed
LVEF-60%

DR. SANDEEP KUMAR
MBBS, PGDCC, CCEBDM
GENERAL PHYSICIAN,
CONSULTANT CLINICAL CARDIOLOGY
SRL LIMITED, SCO-13, Sec-16, FARIDABAD

Dr. Sandeep Kumar
M.B.B.S., PGDCC
General Physician, Consultant Clinical
Cardiology

SRL DIAGNOSTICS WELLNESS CENTER
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PHONE NO- 0129-4179185

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Pancreas : Normal in size shape and echotexture no e/o focal lesion /calcification. Pancreatic duct appears normal.

Spleen: Normal in size, shape and Echotexture. No e/o focal lesion

Both Kidneys: Both kidneys are normal in size and echotexture No e/o hydronephrosis/focal lesion

Urinary bladder: Well distended. No e/o calculi/internal echoes. Wall thickness appears normal.

UTERUS: Normal in size, Shape, position and echotexture. Endometrial cavity is central empty. No focal myometrial lesion seen.

OVARIES: Both the ovaries are normal in size, Position and echotexture
No adnexal mass is seen
There is no free fluid seen in the cul-de-sac

IMPRESSION:- WHOLE ABDOMEN REVEALS FATTY CHANGES IN THE LIVER .

Correlate with clinically findings.

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DR. D.R CHUGH
(RADIOLOGIST)

Disclaimer:

The science of ultrasound is based upon interpretation of moving shadows of normal and abnormal tissue. This is neither complete nor accurate, hence findings should always be interpreted in to the light of clinico-pathological correlation. This a professional opinion, not a diagnosis. Not meant for medico legal purpose.

NAME :- MRS. ANU KHANDELWAL	Age/ Sex/35/Years/F
ACC:- 0071VG000271	Date :- 09/07/2022

ELECTROCARDIOGRAM

Result	Values	Nomal Rate
Rate	108	60-100b/m
Rhythm	Sin	Sinus
P Wave	0.11	Width<0.11Sec.Height<3mm
QRS complex	0.10	<0.10sec in duration
T Wave	Upright	Upright
U Wave	absent	
P R Interval	0.14	0.12 – 0.20sec.
S T segment	iso	Isoelectric

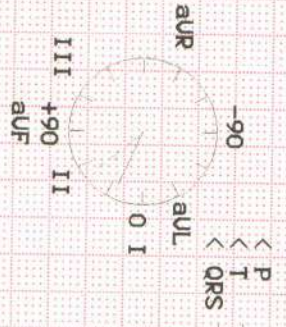
IMPRESSION : *Tachycardia*
otherwise normal

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Disclaimer:

The science of cardiology is based upon interpretation of normal and abnormal ECG graph. This is neither complete or accurate, hence findings should always be interpreted in to the light of clinico-pathological correlation. This a professional opinion, not a diagnosis. Not meant for medico legal purpose.

Measurement Results:
 QRS : 102 ms
 QT/QTcB : 346 / 467 ms
 PR : 142 ms
 P : 118 ms
 RR/PP : 548 / 540 ms
 P/QRS/T : 45 / 25 / 50 degrees
 QTd/QTcBD : 36 / 49 ms
 Sokolow : 1.1 mV
 NK : 15



Interpretation:
 sinus tachycardia
 R/S inversion area between U1 and U2
 probably abnormal ECG

Unconfirmed report.

