



CLIENT CODE : C000138396
CLIENT'S NAME AND ADDRESS :
 ACROFEMI HEALTHCARE LTD (MEDIWHEEL)
 F-703, F-703, LADO SARAI, MEHRAULI
 SOUTH WEST DELHI
 NEW DELHI 110030
 DELHI INDIA
 8800465156

SRL Ltd
 57, Cowley Brown Road, R S Puram
 COIMBATORE, 641002
 TAMILNADU, INDIA
 Tel : 9111591115, Fax : CIN - U74899PB1995PLC045956
 Email : customercare.coimbatore@srl.in

PATIENT NAME : KOUSIKA J

PATIENT ID : KOUSF310789183

ACCESSION NO : **0183WB001656** AGE : 33 Years SEX : Female

DRAWN : 24/02/2023 00:00

RECEIVED : 24/02/2023 09:39

REPORTED : 01/03/2023 17:08

REFERRING DOCTOR : DR. BANK OF BARODA

CLIENT PATIENT ID :

Test Report Status	Final	Results	Biological Reference Interval	Units
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MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE

BLOOD COUNTS,EDTA WHOLE BLOOD

HEMOGLOBIN (HB)	13.2	12.0 - 15.0	g/dL
RED BLOOD CELL (RBC) COUNT	4.61	3.8 - 4.8	mil/ μ L
WHITE BLOOD CELL (WBC) COUNT	6.90	4.0 - 10.0	thou/ μ L
PLATELET COUNT	278	150 - 410	thou/ μ L

RBC AND PLATELET INDICES

HEMATOCRIT (PCV)	41.0	36 - 46	%
MEAN CORPUSCULAR VOLUME (MCV)	89.0	83 - 101	fL
MEAN CORPUSCULAR HEMOGLOBIN (MCH)	28.7	27.0 - 32.0	pg
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION (MCHC)	32.2	31.5 - 34.5	g/dL
RED CELL DISTRIBUTION WIDTH (RDW)	12.4	11.6 - 14.0	%
MENTZER INDEX	19.3		
MEAN PLATELET VOLUME (MPV)	7.4	6.8 - 10.9	fL

WBC DIFFERENTIAL COUNT

NEUTROPHILS	50	40 - 80	%
LYMPHOCYTES	38	20 - 40	%
MONOCYTES	04	2 - 10	%
EOSINOPHILS	07	High 1 - 6	%
BASOPHILS	01	< 1 - 2	%
ABSOLUTE NEUTROPHIL COUNT	3.45	2.0 - 7.0	thou/ μ L
ABSOLUTE LYMPHOCYTE COUNT	2.62	1.0 - 3.0	thou/ μ L
ABSOLUTE MONOCYTE COUNT	0.28	0.2 - 1.0	thou/ μ L
ABSOLUTE EOSINOPHIL COUNT	0.48	0.02 - 0.50	thou/ μ L
ABSOLUTE BASOPHIL COUNT	0.07	0.02 - 0.10	thou/ μ L
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	1.3		

ERYTHROCYTE SEDIMENTATION RATE (ESR),WHOLE BLOOD

E.S.R	23	High 0 - 20	mm at 1 hr
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GLUCOSE FASTING,FLUORIDE PLASMA

FBS (FASTING BLOOD SUGAR)	106	High 74 - 99	mg/dL
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METHOD : HEXOKINASE / SPECTROPHOTOMETRY

GLYCOSYLATED HEMOGLOBIN(HBA1C), EDTA WHOLE BLOOD



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HBA1C		5.7	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 ADA Target: 7.0 Action suggested: > 8.0	%
ESTIMATED AVERAGE GLUCOSE(EAG)		116.9	High < 116.0	mg/dL
GLUCOSE, POST-PRANDIAL, PLASMA				
PPBS(POST PRANDIAL BLOOD SUGAR)		99	70 - 139	mg/dL
METHOD : HEXOKINASE / SPECTROPHOTOMETRY				
LIPID PROFILE, SERUM				
CHOLESTEROL, TOTAL		201	High < 200 Desirable 200 - 239 Borderline High >/= 240 High	mg/dL
METHOD : CHOLESTEROL OXIDASE / SPECTROPHOTOMETRY				
TRIGLYCERIDES		112	< 150 Normal 150 - 199 Borderline High 200 - 499 High >/=500 Very High	mg/dL
HDL CHOLESTEROL		50	< 40 Low >/=60 High	mg/dL
CHOLESTEROL LDL		129	High < 100 Optimal 100 - 129 Near optimal/ above optimal 130 - 159 Borderline High 160 - 189 High >/= 190 Very High	mg/dL
NON HDL CHOLESTEROL		151	High Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL
VERY LOW DENSITY LIPOPROTEIN		22.4	</= 30.0	mg/dL
CHOL/HDL RATIO		4.0	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		2.6	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	

LIVER FUNCTION PROFILE, SERUM



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BILIRUBIN, TOTAL		0.50	0.2 - 1.0	mg/dL
METHOD : DIAZOTIZED SULFANILIC ACID / SPECTROPHOTOMETRY				
BILIRUBIN, DIRECT		0.10	0.0 - 0.2	mg/dL
METHOD : DIAZOTIZED SULFANILIC ACID / SPECTROPHOTOMETRY				
BILIRUBIN, INDIRECT		0.4	0.1 - 1.0	mg/dL
TOTAL PROTEIN		6.5	6.4 - 8.2	g/dL
ALBUMIN		3.8	3.4 - 5.0	g/dL
METHOD : BCP DYE BINDING / SPECTOPHOTOMETER				
GLOBULIN		2.7	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO		1.4	1.0 - 2.1	RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT)		14	Low 15 - 37	U/L
METHOD : UV WITH PYRIDOXAL 5 PHOSPHATE / SPECTROPHOTOMETER				
ALANINE AMINOTRANSFERASE (ALT/SGPT)		20	< 34.0	U/L
METHOD : UV WITH PYRIDOXAL 5 PHOSPHATE / SPECTROPHOTOMETER				
ALKALINE PHOSPHATASE		56	30 - 120	U/L
GAMMA GLUTAMYL TRANSFERASE (GGT)		22	5 - 55	U/L
METHOD : GCNA / SPECTROPHOTOMETRY				
LACTATE DEHYDROGENASE		117	100 - 190	U/L
METHOD : LACTATE PYRUVATE UV/ L.LACTATE / SPECTOPHOTOMETER				
BLOOD UREA NITROGEN (BUN), SERUM				
BLOOD UREA NITROGEN		10	6 - 20	mg/dL
METHOD : UREASE / GLDH / SPECTROPHOTOMETRY				
CREATININE, SERUM				
CREATININE		0.58	Low 0.60 - 1.10	mg/dL
METHOD : PICRATE/ JAFFE / SPECTOPHOTOMETER				
BUN/CREAT RATIO				
BUN/CREAT RATIO		17.24	High 5.00 - 15.00	
URIC ACID, SERUM				
URIC ACID		5.6	2.6 - 6.0	mg/dL
METHOD : URICASE / CATALASE UV / SPECTROPHOTOMETRY				
TOTAL PROTEIN, SERUM				
TOTAL PROTEIN		6.5	6.4 - 8.2	g/dL
ALBUMIN, SERUM				
ALBUMIN		3.8	3.4 - 5.0	g/dL
METHOD : BCP DYE BINDING / SPECTOPHOTOMETER				
GLOBULIN				



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T3		118.90	Non-Pregnant Women 80.0 - 200.0 Pregnant Women 1st Trimester:105.0 - 230.0 2nd Trimester:129.0 - 262.0 3rd Trimester:135.0 - 262.0	ng/dL
T4		10.45	Non-Pregnant Women 5.10 - 14.10 Pregnant Women 1st Trimester: 7.33 - 14.80 2nd Trimester: 7.93 - 16.10 3rd Trimester: 6.95 - 15.70	µg/dL
TSH (ULTRASENSITIVE)		1.090	Non Pregnant Women 0.27 - 4.20 Pregnant Women 1st Trimester: 0.33 - 4.59 2nd Trimester: 0.35 - 4.10 3rd Trimester: 0.21 - 3.15	µIU/mL

LETTER

ADDITIONAL COMMUNICATION

SAMPLE NOT RECEIVED

MICROSCOPIC EXAMINATION,STOOL

REMARK

TEST CANCELLED AS SPECIMEN NOT RECEIVED

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP

TYPE B

RH TYPE

POSITIVE

XRAY-CHEST

>>>

BOTH THE LUNG FIELDS ARE CLEAR

>>>

BOTH THE COSTOPHRENIC AND CARIOPHRENIC ANGELS ARE CLEAR

>>>

BOTH THE HILA ARE NORMAL

>>>

CARDIAC AND AORTIC SHADOWS APPEAR NORMAL

>>>

BOTH THE DOMES OF THE DIAPHRAM ARE NORMAL

>>>

VISUALIZED BONY THORAX IS NORMAL

IMPRESSION

NO ABNORMALITY DETECTED

TMT OR ECHO

TMT OR ECHO

DONE

ECG

ECG

WITHIN NORMAL LIMITS

MEDICAL HISTORY

RELEVANT PRESENT HISTORY

NOT SIGNIFICANT



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RELEVANT PAST HISTORY	NOT SIGNIFICANT
RELEVANT PERSONAL HISTORY	MARRIED
MENSTRUAL HISTORY (FOR FEMALES)	REGULAR
LMP (FOR FEMALES)	01/02/2023
OBSTETRIC HISTORY (FOR FEMALES)	G1 P1 A0
LCB (FOR FEMALES)	2018
RELEVANT FAMILY HISTORY	NOT SIGNIFICANT
OCCUPATIONAL HISTORY	NOT SIGNIFICANT
HISTORY OF MEDICATIONS	NOT SIGNIFICANT

ANTHROPOMETRIC DATA & BMI

HEIGHT IN METERS	1.62	mts
WEIGHT IN KGS.	77	Kgs
BMI	29	

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	OVERWEIGHT
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
THYROID GLAND	NOT ENLARGED
CAROTID PULSATION	NORMAL
BREAST (FOR FEMALES)	NORMAL
TEMPERATURE	NORMAL
PULSE	78/MINS, REGULAR, ALL PERIPHERAL PULSES WELL FELT, NO CAROTID BRUIT
RESPIRATORY RATE	NORMAL

CARDIOVASCULAR SYSTEM



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BP		120/80 MM HG (SITTING)		mm/Hg
PERICARDIUM		NORMAL		
APEX BEAT		NORMAL		
HEART SOUNDS		NORMAL		
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		
BASIC EYE EXAMINATION				
CONJUNCTIVA		NORMAL		
EYELIDS		NORMAL		
EYE MOVEMENTS		NORMAL		
CORNEA		NORMAL		
DISTANT VISION RIGHT EYE WITHOUT GLASSES		6/6		



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DISTANT VISION LEFT EYE WITHOUT GLASSES	6/6
NEAR VISION RIGHT EYE WITHOUT GLASSES	N/6
NEAR VISION LEFT EYE WITHOUT GLASSES	N/6
COLOUR VISION	NORMAL

BASIC ENT EXAMINATION

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

BASIC DENTAL EXAMINATION

TEETH	NORMAL
GUMS	HEALTHY

SUMMARY

RELEVANT HISTORY	NOT SIGNIFICANT
RELEVANT GP EXAMINATION FINDINGS	NOT SIGNIFICANT
RELEVANT LAB INVESTIGATIONS	WITHIN NORMAL LIMITS
RELEVANT NON PATHOLOGY DIAGNOSTICS	NO ABNORMALITIES DETECTED
REMARKS / RECOMMENDATIONS	NONE

FITNESS STATUS

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

OUR PANEL OF DOCTORS :
 GENERAL PHYSICIANS - DR.S. B. PRAVEEN., M.B.B.S.,M.Sc(Psy).,F.Diab., AFIH.
 RADIOLOGIST - DR.DEBABRATA NITYARANJAN DAS, MD(RAD)., M.R. FELLOW (USA)
 GYNECOLOGIST - DR. PREMALATHA KRISHNAKUMAR. MD.,MRCOG., Dip.in Colposcopy(UK).

THIS REPORT CARRIES THE SIGNATURE OF OUR LABORATORY HEAD.
 THIS IS AN INVIOABLE FEATURE OF OUR LAB MANAGEMENT SOFTWARE.
 HOWEVER ALL EXAMINATIONS 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.



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RBC AND PLATELET INDICES-Mentzer index (MCV/RBC) is an automated cell-counter based calculated screen tool to differentiate cases of Iron deficiency anaemia(>13) from Beta thalassaemia trait (<13) in patients with microcytic anaemia. This needs to be interpreted in line with clinical correlation and suspicion. Estimation of HbA2 remains the gold standard for diagnosing a case of beta thalassaemia trait.
 WBC DIFFERENTIAL COUNT-The optimal threshold of 3.3 for NLR showed a prognostic possibility of clinical symptoms to change from mild to severe in COVID positive patients. When age = 49.5 years old and NLR = 3.3, 46.1% COVID-19 patients with mild disease might become severe. By contrast, when age < 49.5 years old and NLR < 3.3, COVID-19 patients tend to show mild disease.
 (Reference to - The diagnostic and predictive role of NLR, d-NLR and PLR in COVID-19 patients A.-P. Yang, et al. International Immunopharmacology 84 (2020) 106504 This ratio element is a calculated parameter and out of NABL scope.)
ERYTHROCYTE SEDIMENTATION RATE (ESR),WHOLE BLOOD-TEST DESCRIPTION :-
 Erythrocyte sedimentation rate (ESR) is a test that indirectly measures the degree of inflammation present in the body. The test actually measures the rate of fall (sedimentation) of erythrocytes in a sample of blood that has been placed into a tall, thin, vertical tube. Results are reported as the millimetres of clear fluid (plasma) that are present at the top portion of the tube after one hour. Nowadays fully automated instruments are available to measure ESR.

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

Increase in: Infections, Vasculities, Inflammatory arthritis, Renal disease, Anemia, Malignancies and plasma cell dyscrasias, Acute allergy Tissue injury, Pregnancy, Estrogen medication, Aging.
 Finding a very accelerated ESR(>100 mm/hour) in patients with ill-defined symptoms directs the physician to search for a systemic disease (Paraproteinemias, Disseminated malignancies, connective tissue disease, severe infections such as bacterial endocarditis).
 In pregnancy BRI in first trimester is 0-48 mm/hr(62 if anemic) and in second trimester (0-70 mm/hr(95 if anemic). ESR returns to normal 4th week post partum.
Decreased in: Polycythemia vera, Sickle cell anemia

LIMITATIONS

False elevated ESR : Increased fibrinogen, Drugs(Vitamin A, Dextran etc), Hypercholesterolemia
False Decreased : Poikilocytosis,(SickleCells,spherocytes),Microcytosis, Low fibrinogen, Very high WBC counts, Drugs(Quinine, salicylates)

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,FLUORIDE PLASMA-TEST DESCRIPTION
 Normally, the glucose concentration in extracellular fluid is closely regulated so that a source of energy is readily available to tissues and sothat no glucose is excreted in the urine.
Increased in:Diabetes mellitus, Cushing's syndrome (10 – 15%), chronic pancreatitis (30%). Drugs:corticosteroids,phenytoin, estrogen, thiazides.
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:

- Evaluating the long-term control of blood glucose concentrations in diabetic patients.
 - Diagnosing diabetes.
 - 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 patients metabolic control has remained continuously within the target range.
- eAG (Estimated average glucose) converts percentage HbA1c to mg/dl, to compare blood glucose levels.
 - eAG gives an evaluation of blood glucose levels for the last couple of months.
 - eAG is calculated as eAG (mg/dl) = 28.7 * HbA1c - 46.7

HbA1c Estimation can get affected due to :

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



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 COIMBATORE, 641002
 TAMILNADU, INDIA
 Tel : 9111591115, Fax : CIN - U74899PB1995PLC045956
 Email : customercare.coimbatore@srl.in

PATIENT NAME : KOUSIKA J PATIENT ID : **KOUSF310789183**

ACCESSION NO : **0183WB001656** AGE : 33 Years SEX : Female

DRAWN : 24/02/2023 00:00 RECEIVED : 24/02/2023 09:39 REPORTED : 01/03/2023 17:08

REFERRING DOCTOR : DR. BANK OF BARODA CLIENT PATIENT ID :

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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

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

LIVER FUNCTION PROFILE, SERUM-
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** results 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, Alcohol 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 perniciosa 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, Pagets disease, Rickets, Sarcoidosis etc. Lower-than-normal ALP levels seen in Hypophosphatasia, Malnutrition, Protein deficiency, Wilsons 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.

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, Waldenstroms 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 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

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, Muscuophy

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

TOTAL PROTEIN, SERUM- 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, Waldenstroms 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.

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 INVOLUBLE 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



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CLIENT CODE : C000138396
CLIENT'S NAME AND ADDRESS :
 ACROFEMI HEALTHCARE LTD (MEDIWHEEL)
 F-703, F-703, LADO SARAI, MEHRAULI
 SOUTH WEST DELHI
 NEW DELHI 110030
 DELHI INDIA
 8800465156

SRL Ltd
 57, Cowley Brown Road, R S Puram
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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.



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MEDI WHEEL FULL BODY HEALTH CHECKUP BELOW 40FEMALE

ULTRASOUND ABDOMEN
ULTRASOUND ABDOMEN
 NO ABNORMALITIES DETECTED

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



Dr. Karthick Prabhu R
 Consultant 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. 3. Result delays could occur due to unforeseen circumstances such as non-availability of kits / equipment breakdown / natural calamities / technical downtime or any other unforeseen event. 4. A requested test might not be performed if: i. Specimen received is insufficient or inappropriate ii. Specimen quality is unsatisfactory iii. Incorrect specimen type iv. Discrepancy between identification on specimen container label and test requisition form	5. SRL confirms that all tests have been performed or assayed with highest quality standards, clinical safety & technical integrity. 6. Laboratory results should not be interpreted in isolation; it must be correlated with clinical information and be interpreted by registered medical practitioners only to determine final diagnosis. 7. Test results may vary based on time of collection, physiological condition of the patient, current medication or nutritional and dietary changes. Please consult your doctor or call us for any clarification. 8. Test results cannot be used for Medico legal purposes. 9. In case of queries please call customer care (91115 91115) within 48 hours of the report.
<p>SRL Limited Fortis Hospital, Sector 62, Phase VIII, Mohali 160062</p>	



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