



CLIENT CODE : C000138355

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
34/2, NEW PALASIA, NEAR OM SHANTI BHAWAN CIRCLE,BEHIND
INDUSTRY HOUSE
INDORE, 452001
MADHYA PRADESH, INDIA
Tel : 9111591115,
CIN - U74899PB1995PLC045956
Email : customercare.indore@srl.in

PATIENT NAME : SHYAM MALVIYA

PATIENT ID : SHYAM1809897

ACCESSION NO : 0007VI002197 AGE : 32 Years SEX : Male

ABHA NO :

DRAWN :

RECEIVED : 10/09/2022 10:14

REPORTED : 10/09/2022 16:33

REFERRING DOCTOR : DR. ACROFEMI HEALTHCARE LTD (MEDIWHEEL)

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Test Report Status	Final	Results	Biological Reference Interval	Units
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MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE**BLOOD COUNTS,EDTA WHOLE BLOOD**

HEMOGLOBIN	17.1	High	13.0 - 17.0	g/dL
METHOD : SPECTROPHOTOMETRIC				
RED BLOOD CELL COUNT	5.35		4.5 - 5.5	mil/ μ L
METHOD : ELECTRICAL IMPEDANCE				
WHITE BLOOD CELL COUNT	5.50		4.0 - 10.0	thou/ μ L
PLATELET COUNT	310		150 - 410	thou/ μ L
METHOD : ELECTRICAL IMPEDANCE				

RBC AND PLATELET INDICES

HEMATOCRIT	50.0		40 - 50	%
METHOD : CALCULATED PARAMETER				
MEAN CORPUSCULAR VOL	93.0		83 - 101	fL
METHOD : CALCULATED PARAMETER				
MEAN CORPUSCULAR HGB.	31.9		27.0 - 32.0	pg
METHOD : CALCULATED PARAMETER				
MEAN CORPUSCULAR HEMOGLOBIN CONCENTRATION	34.1		31.5 - 34.5	g/dL
METHOD : CALCULATED PARAMETER				
MENTZER INDEX	17.4			
RED CELL DISTRIBUTION WIDTH	13.3		11.6 - 14.0	%
METHOD : CALCULATED PARAMETER				
MEAN PLATELET VOLUME	8.7		6.8 - 10.9	fL
METHOD : CALCULATED PARAMETER				

WBC DIFFERENTIAL COUNT - NLR

SEGMENTED NEUTROPHILS	48		40 - 80	%
METHOD : IMPEDENCE / MICROSCOPY				
ABSOLUTE NEUTROPHIL COUNT	2.64		2.0 - 7.0	thou/ μ L
METHOD : CALCULATED PARAMETER				
LYMPHOCYTES	40		20 - 40	%
METHOD : IMPEDENCE / MICROSCOPY				
ABSOLUTE LYMPHOCYTE COUNT	2.2		1.0 - 3.0	thou/ μ L
METHOD : CALCULATED PARAMETER				
NEUTROPHIL LYMPHOCYTE RATIO (NLR)	1.2			
METHOD : CALCULATED PARAMETER				
EOSINOPHILS	06		1 - 6	%



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METHOD : IMPEDENCE / MICROSCOPY				
ABSOLUTE EOSINOPHIL COUNT		0.33	0.02 - 0.50	thou/ μ L
METHOD : CALCULATED PARAMETER				
MONOCYTES		06	2 - 10	%
METHOD : IMPEDENCE / MICROSCOPY				
ABSOLUTE MONOCYTE COUNT		0.33	0.2 - 1.0	thou/ μ L
METHOD : CALCULATED PARAMETER				
DIFFERENTIAL COUNT PERFORMED ON:		EDTA SMEAR		

Comments

Please note that :

The Automatic analyzer used to estimate Complete Blood Counts (Blood cell Indices & counts) is "ABX PENTRA XL 80" (HORIBA); the values are correlated manually with microscopic picture.

ERYTHRO SEDIMENTATION RATE, BLOOD

SEDIMENTATION RATE (ESR)	05	0 - 14	mm at 1 hr
METHOD : WESTERGREN METHOD			

GLUCOSE, FASTING, PLASMA

GLUCOSE, FASTING, PLASMA	99	74 - 99	mg/dL
METHOD : HEXOKINASE			

GLYCOSYLATED HEMOGLOBIN, EDTA WHOLE BLOOD

GLYCOSYLATED HEMOGLOBIN (HBA1C)	5.6	Non-diabetic: < 5.7 Pre-diabetics: 5.7 - 6.4 Diabetics: > or = 6.5 ADA Target: 7.0 Action suggested: > 8.0	%
METHOD : HPLC			

MEAN PLASMA GLUCOSE	114.0	< 116.0	mg/dL
METHOD : CALCULATED PARAMETER			

GLUCOSE, POST-PRANDIAL, PLASMA

GLUCOSE, POST-PRANDIAL, PLASMA	115	Normal: < 140, Impaired Glucose Tolerance: 140-199 Diabetic > or = 200	mg/dL
METHOD : HEXOKINASE			

CORONARY RISK PROFILE, SERUM

CHOLESTEROL	170	Desirable: <200 BorderlineHigh : 200-239 High : > or = 240	mg/dL
METHOD : OXIDASE, ESTERASE, PEROXIDASE			



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TRIGLYCERIDES		95	Desirable: < 150 Borderline High: 150 - 199 High: 200 - 499 Very High : > or = 500	mg/dL
METHOD : ENZYMATIC ASSAY				
HDL CHOLESTEROL		39	Low < 40 High > or = 60	mg/dL
CHOLESTEROL LDL		112	High Adult levels: Optimal < 100 Near optimal/above optimal: 100-129 Borderline high : 130-159 High : 160-189 Very high : = 190	mg/dL
NON HDL CHOLESTEROL		131	High Desirable: Less than 130 Above Desirable: 130 - 159 Borderline High: 160 - 189 High: 190 - 219 Very high: > or = 220	mg/dL
CHOL/HDL RATIO		4.4		
LDL/HDL RATIO		2.9	0.5 - 3.0 Desirable/Low Risk 3.1 - 6.0 Borderline/Moderate Risk >6.0 High Risk	
VERY LOW DENSITY LIPOPROTEIN		19.0		mg/dL
LIVER FUNCTION PROFILE, SERUM				
BILIRUBIN, TOTAL		0.66	0.0 - 1.2	mg/dL
METHOD : JENDRASSIK AND GROFF				
BILIRUBIN, DIRECT		0.26	High 0.0 - 0.2	mg/dL
METHOD : DIAZOTIZATION				
BILIRUBIN, INDIRECT		0.40	0.00 - 1.00	mg/dL
TOTAL PROTEIN		8.2	6.4 - 8.3	g/dL
METHOD : BIURET				
ALBUMIN		5.6	High 3.50 - 5.20	g/dL
METHOD : BROMOCRESOL PURPLE				
GLOBULIN		2.6	2.0 - 4.1	g/dL
ALBUMIN/GLOBULIN RATIO		2.2	High 1.0 - 2.0	RATIO
ASPARTATE AMINOTRANSFERASE (AST/SGOT)		28	UPTO 40	U/L
METHOD : UV WITH P5P				
ALANINE AMINOTRANSFERASE (ALT/SGPT)		38	UP TO 45	U/L
METHOD : UV WITH P5P				
ALKALINE PHOSPHATASE		90	40 - 129	U/L



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METHOD : PNPP

GAMMA GLUTAMYL TRANSFERASE (GGT)	34	8 - 61	U/L
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METHOD : G-GLUTAMYL-CARBOXY-NITROANILIDE

LACTATE DEHYDROGENASE	212	135 - 225	U/L
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METHOD : ENZYMATIC LACTATE - PYRUVATE(IFCC)

SERUM BLOOD UREA NITROGEN

BLOOD UREA NITROGEN	10	6 - 20	mg/dL
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METHOD : UREASE KINETIC

CREATININE, SERUM

CREATININE	0.93	0.70 - 1.20	mg/dL
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METHOD : ALKALINE PICRATE-KINETIC

BUN/CREAT RATIO

BUN/CREAT RATIO	10.75	5.0 - 15.0	
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URIC ACID, SERUM

URIC ACID	7.4	High 3.5 - 7.2	mg/dL
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METHOD : URICASE/CATALASE UV

TOTAL PROTEIN, SERUM

TOTAL PROTEIN	8.2	6.4 - 8.3	g/dL
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METHOD : BIURET

ALBUMIN, SERUM

ALBUMIN	5.6	High 3.5 - 5.2	g/dL
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METHOD : BROMOCRESOL PURPLE

GLOBULIN

GLOBULIN	2.6	2.0 - 4.1	g/dL
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ELECTROLYTES (NA/K/CL), SERUM

SODIUM	144.1	136.0 - 146.0	mmol/L
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POTASSIUM	4.38	3.50 - 5.10	mmol/L
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CHLORIDE	103.1	98.0 - 106.0	mmol/L
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PHYSICAL EXAMINATION, URINE

COLOR	PALE YELLOW		
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METHOD : MACROSCOPY

APPEARANCE	CLEAR		
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METHOD : VISUAL

SPECIFIC GRAVITY	<=1.005	1.003 - 1.035	
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METHOD : REFLECTANCE SPECTROPHOTOMETRY

CHEMICAL EXAMINATION, URINE

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PH		5.5	4.7 - 7.5	
METHOD : PH INDICATOR AND REFLECTANCE				
PROTEIN		NOT DETECTED	NOT DETECTED	
METHOD : PROTEIN ERROR OF INDICATORS WITH REFLECTANCE				
GLUCOSE		NOT DETECTED	NOT DETECTED	
METHOD : GLUCOSE OXIDASE				
KETONES		NOT DETECTED	NOT DETECTED	
METHOD : ROTHERA'S WITH REFLECTANCE				
BLOOD		NOT DETECTED	NOT DETECTED	
METHOD : PEROXIDASE METHOD WITH REFLECTANCE				
BILIRUBIN		NOT DETECTED	NOT DETECTED	
METHOD : DIAZOTIZED WITH REFLECTANCE				
UROBILINOGEN		NORMAL	NORMAL	
METHOD : EHRlich REACTION REFLECTANCE				
NITRITE		NOT DETECTED	NOT DETECTED	
METHOD : DIAZOTIZED WITH REFLECTANCE				
LEUKOCYTE ESTERASE		NOT DETECTED	NOT DETECTED	
MICROSCOPIC EXAMINATION, URINE				
PUS CELL (WBC'S)		2-3	0-5	/HPF
METHOD : ESTERASES METHOD WITH REFLECTANCE				
EPITHELIAL CELLS		1-2	0-5	/HPF
METHOD : MICROSCOPIC EXAMINATION				
ERYTHROCYTES (RBC'S)		NOT DETECTED	NOT DETECTED	/HPF
CASTS		NOT DETECTED		
METHOD : MICROSCOPIC EXAMINATION				
CRYSTALS		NOT DETECTED		
METHOD : MICROSCOPIC EXAMINATION				
BACTERIA		NOT DETECTED	NOT DETECTED	
METHOD : MICROSCOPIC EXAMINATION				
YEAST		NOT DETECTED	NOT DETECTED	
REMARKS		Please note that all the urinary findings are confirmed manually as well.		
THYROID PANEL, SERUM				
T3		135.3	80.00 - 200.00	ng/dL
METHOD : ELECTROCHEMILUMINESCENCE IMMUNO ASSAY				
T4		8.96	5.10 - 14.10	µg/dL
METHOD : ELECTROCHEMILUMINESCENCE IMMUNO ASSAY				



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TSH 3RD GENERATION	2.540	0.270 - 4.200	µIU/mL
METHOD : ELECTROCHEMILUMINESCENCE IMMUNO ASSAY			

ABO GROUP & RH TYPE, EDTA WHOLE BLOOD

ABO GROUP TYPE O

METHOD : TUBE AGGLUTINATION

RH TYPE POSITIVE

METHOD : TUBE AGGLUTINATION

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 NEGATIVE

ECG

ECG SINUS RHYTHM, NORMAL ECG

MEDICAL HISTORY

RELEVANT PRESENT HISTORY NOT SIGNIFICANT
 RELEVANT PAST HISTORY NOT SIGNIFICANT
 RELEVANT PERSONAL HISTORY NOT SIGNIFICANT
 RELEVANT FAMILY HISTORY F/H/O HTN/ HYPOTHYROIDISM- MOTHER
 OCCUPATIONAL HISTORY NOT SIGNIFICANT
 HISTORY OF MEDICATIONS NOT SIGNIFICANT

ANTHROPOMETRIC DATA & BMI

HEIGHT IN METERS	1.72	mts
WEIGHT IN KGS.	80	Kgs
BMI	27	

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



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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		
TEMPERATURE		AFEBRILE		
PULSE		72/MIN, REGULAR, ALL PERIPHERAL PULSES WELL FELT, NO CAROTID BRUIT		
RESPIRATORY RATE		NORMAL		
CARDIOVASCULAR SYSTEM				
BP		130/80 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		



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HERNIA

NORMAL

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 WITH GLASSES

N/6, WITH GLASSES NORMAL

DISTANT VISION LEFT EYE WITH GLASSES

N/6, WITH GLASSES NORMAL

NEAR VISION RIGHT EYE WITHOUT GLASSES

6/6, WITHIN NORMAL LIMIT

NEAR VISION LEFT EYE WITHOUT GLASSES

6/6, WITHIN NORMAL LIMIT

COLOUR VISION

NORMAL

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 GP EXAMINATION FINDINGS

OVERWEIGHT

REMARKS / RECOMMENDATIONS

NONE

FITNESS STATUS

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

CLINICAL FINDINGS :-

RAISED URIC ACID.

DYSLIPIDEMIA.

OVER WEIGHT STATUS.

USG SHOWS EARLY FATTY INFILTRATION OF LIVER.

FITNESS STATUS :-

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

ADVICE : WEIGHT REDUCTION, LOW FAT& CARBOHYDRATE DIET AND REGULAR PHYSICAL EXERCISE FOR OVERWEIGHT STATUS AND DYSLIPIDEMIA.

NEED PHYSICIAN CONSULTATION FOR LIFE STYLE MODIFICATION.

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 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 poikilocytosis, 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



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 CIN - U74899PB1995PLC045956
 Email : customercare.indore@srl.in

PATIENT NAME : SHYAM MALVIYA

PATIENT ID : SHYAM1809897

ACCESSION NO : 0007VI002197 **AGE :** 32 Years **SEX :** Male

ABHA NO :

DRAWN : **RECEIVED :** 10/09/2022 10:14

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REFERRING DOCTOR : DR. ACROFEMI HEALTHCARE LTD (MEDIWHEEL)

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Test Report Status	Final	Results	Biological Reference Interval	Units
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complications in patients with diabetes mellitus. Formation of GHb is essentially irreversible, and the concentration in the blood depends on both the life span of the red blood cell (average 120 days) and the blood glucose concentration. Because the rate of formation of GHb is directly proportional to the concentration of glucose in the blood, the GHb concentration represents the integrated values for glucose over the preceding 6-8 weeks. Any condition that alters the life span of the red blood cells has the potential to alter the GHb level. Samples from patients with hemolytic anemias will exhibit decreased glycated hemoglobin values due to the shortened life span of the red cells. This effect will depend upon the severity of the anemia. Samples from patients with polycythemia or post-splenectomy may exhibit increased glycated hemoglobin values due to a somewhat longer life span of the red cells. Glycosylated hemoglobins results from patients with HbSS, HbCC, and HbSC and HbD must be interpreted with caution, given the pathological processes, including anemia, increased red cell turnover, transfusion requirements, that adversely impact HbA1c as a marker of long-term glycemic control. In these conditions, alternative forms of testing such as glycated serum protein (fructosamine) should be considered. "Targets should be individualized; More or less stringent glycemic goals may be appropriate for individual patients. Goals should be individualized based on duration of diabetes, age/life expectancy, comorbid conditions, known CVD or advanced microvascular complications, hypoglycemia unawareness, and individual patient considerations."

References

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

**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 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, 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)



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PATIENT NAME : SHYAM MALVIYA

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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.

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



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PATIENT NAME : SHYAM MALVIYA

PATIENT ID : SHYAM1809897

ACCESSION NO : 0007VI002197 **AGE :** 32 Years **SEX :** Male **ABHA NO :**

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heart rate. Production of T3 and its prohormone thyroxine (T4) is activated by thyroid-stimulating hormone (TSH), which is released from the pituitary gland. Elevated concentrations of T3, and T4 in the blood inhibit the production of TSH. Thyroxine T4, Thyroxine's principal function is to stimulate the metabolism of all cells and tissues in the body. Excessive secretion of thyroxine in the body is hyperthyroidism, and deficient secretion is called hypothyroidism. Most of the thyroid hormone in blood is bound to transport proteins. Only a very small fraction of the circulating hormone is free and biologically active.

In primary hypothyroidism, TSH levels are significantly elevated, while in secondary and tertiary hypothyroidism, TSH levels are low.

Below mentioned are the guidelines for Pregnancy related reference ranges for Total T4, TSH & Total T3

Levels in	TOTAL T4	TSH3G	TOTAL T3
Pregnancy	(µg/dL)	(µIU/mL)	(ng/dL)
First Trimester	6.6 - 12.4	0.1 - 2.5	81 - 190
2nd Trimester	6.6 - 15.5	0.2 - 3.0	100 - 260
3rd Trimester	6.6 - 15.5	0.3 - 3.0	100 - 260

Below mentioned are the guidelines for age related reference ranges for T3 and T4.

	T3	T4
	(ng/dL)	(µg/dL)
New Born:	75 - 260	1-3 day: 8.2 - 19.9
.		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

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 INVIOABLE 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.



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MEDI WHEEL FULL BODY HEALTH CHECK UP BELOW 40 MALE**ULTRASOUND ABDOMEN****ULTRASOUND ABDOMEN**

DONE

Comments

USG-

IMPRESSION- EARLY FATTY INFILTRATION OF LIVER.

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

Dr.Arpita Pasari, MD
Consultant Pathologist



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