

Patient Name	: Mrs. TULASI K	Reg. No.	: 00432512270002
Age and Sex	: 37 Yrs / Female	PCC Code	: PCL-KA-656
Referring Doctor	: Dr. Swarna Diagnostic Centre	Sample Drawn Date	: 27-Dec-2025 10:10 AM
Referring Customer	: N/A	Registration Date	: 27-Dec-2025 12:48 PM
Vial ID	: Q3933726	Report Date	: 27-Dec-2025 03:19 PM
Sample Type	: WB-EDTA	Report Status	: Final Report
Client Address	: New Mico Layout, Garve Bavi Palya, Bangalore		

HEMATOLOGY

PATHCHECK ULTIMA

Test Name	Obtained Value	Units	Bio. Ref. Intervals (Age/Gender specific)	Method
Complete Blood Count (CBC)				
Haemoglobin	12.9	g/dL	11.5-15.5	Colorimetric
RBC Count	4.7	10^12/L	4.5-5.5	Electrical Impedance
Haematocrit (HCT)	39.3	%	40-50	Calculated
MCV	83.9	fL	83-101	RBC Histogram
MCH	27.6	pg	27-32	Calculated
MCHC	32.8	g/dL	31.5-34.5	Calculated
RDW-CV	14.0	%	11.6-14.0	RBC Histogram
Platelet Count	205	10^9/L	150-410	Electrical Impedance/Microscopy
WBC count, Total	5.8	10^9/L	4.0-10.0	Impedance
Neutrophils	50.0	%	40-70	Microscopy
Neutrophil-Absolute Count	2.9	10^9/L	2.0-7.0	Calculated
Lymphocytes	41.0	%	20-40	Microscopy
Lymphocytes-Absolute Count	2.4	10^9/L	1.0-3.0	Calculated
Monocytes	6.0	%	2-10	Microscopy
Monocytes-Absolute Count	0.3	10^9/L	0.2-1.0	Calculated
Eosinophils	3.0	%	1-6	Microscopy
Eosinophils-Absolute Count	0.2	10^9/L	0.02-0.5	Calculated
Basophils	0.0	%	0-2	Microscopy
Basophils-Absolute Count	0.0	10^9/L	0.0-0.3	Calculated
Others	0.0	%	00	Microscopy
Remarks	.			

Sample is Processed on Automated CBC Analyzer

Note: Haematocrit (HCT) is derived from calculated MCV based on RBC Histogram as per Manufacturer's Manual

***Erythrocyte Sedimentation Rate 16 (ESR)**

mm in 1hr 12 or less

Westergren method

Comment:

Conditions that may be associated with a highly elevated ESR include the Hypersensitivity Vasculitis, Giant Cell Arteritis, Waldenstrom Macroglobulinemia, Polymyalgia Rheumatic, Metastatic Cancer, Chronic infection, Hyperfibrogenemia etc.



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Referring Customer	: N/A	Registration Date	: 27-Dec-2025 12:48 PM
Vial ID	: Q3933724, Q3933726, Q3933725	Report Date	: 27-Dec-2025 01:48 PM
Sample Type	: Serum, WB-EDTA , Plasma-Sodium	Report Status	: Final Report
Client Address	: New Mico Layout, Garve Bavi Palya, Bangalore		

CLINICAL BIOCHEMISTRY

PATHCHECK ULTIMA

Test Name	Obtained Value	Units	Bio. Ref. Intervals (Age/Gender specific)	Method
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*25 - Hydroxy Vitamin D- Serum	12.87	ng/mL	Deficiency - < 20 Insufficiency - 20 -30 Sufficiency - 30 - 100 Toxicity - >100	CMIA
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Comments:

- 25 OH Vitamin D is total of Vitamin D in Bone and mineral metabolism was recognized from its first identification as a factor that could cure rickets. However, Vitamin D is now recognized as a prohormone which has multiple roles in maintaining optimal health.
- Vitamin D toxicity is a recognized problem but a rare occurrence. Instead, a recent growing public health problem is Vitamin D insufficiency.

*Glycosylated Hemoglobin(GHb/HbA1c)	4.97	%	<5.7 Non diabetic, 5.7 – 6.4 Borderline diabetic,>6.4 Diabetic	Boronate Affinity HPLC
*Glycosylated Hemoglobin	30.82	mmol/mol		Calculated
*Mean Blood Glucose	95.94	mg/dL	90 - 120 : Excellent Control 121 - 150 : Good Control 151 - 180 : Average Control 181 - 210 : Action Suggested >211 :Panic Value	Calculated

Comments:

- HbA1c is an indicator of glycemic control. HbA1c represents average Glycemia over the past six to eight weeks. Glycation of Hemoglobin occurs over the entire 120 day life span of the Red Blood Cell, but within this 120 days. Clinical studies suggest that a patient in stable control will have 50% of their HbA1c formed in the month before sampling, 25% in the month before that, and the remaining 25% in months two to four.
- Mean Plasma Glucose mg/dL = $28.7 \times A1C - 46.7$. Correlation between HbA1c and Mean Plasma Glucose (MPG) is not "perfect" but rather only this means that to predict or estimate average glucose from HbA1c or vice-versa is not "perfect" but gives a good working ballpark estimate.
- Afternoon and evening results correlate more closely to HbA1c than morning results, perhaps because morning fasting glucose levels vary much more than daytime Glucose levels, which are easier to predict and control. As per IFCC recommendations 2007, HbA1c being reported as above maintaining traceability to both IFCC (nmol/mol) & NGSP (%) units.
- Reference: ADA (American Diabetic Association) Guidelines 2023.

*Glucose-Plasma -Fasting	88.0	mg/dL	Normal < 100 Pre-diabetic 100-125 Diabetic >= 126	Hexokinase
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Comments:

- Glucose is the major carbohydrate present in blood. Its oxidation in the cells is the source of energy for the body. Increased levels of Glucose are found in Diabetes Mellitus, Hyperparathyroidism, Pancreatitis and renal failure.
- Decreased levels are found in Insulinoma, Hypothyroidism, Hypopituitarism and extensive Liver disease

Criteria for diagnosis of Diabetes mellitus:

FPG >/=126 mg/dL, Fasting is defined as no caloric intake for atleast 8 hours.

Fasting hypoglycemia may be observed in persons taking certain diabetes medications, antibiotics and alcoholic beverages.

Additionally intestinal disorders, endocrine disorders, insulin surge, nature of diet, reduced food intake than usual and stress are among several other factors. Please correlate clinically.

Reference: American Diabetes association guidelines 2021.



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Sample Type	: Serum, WB-EDTA , Plasma-Sodium	Report Status	: Final Report
Client Address	: New Mico Layout, Garve Bavi Palya, Bangalore		

CLINICAL BIOCHEMISTRY

PATHCHECK ULTIMA

Test Name	Obtained Value	Units	Bio. Ref. Intervals (Age/Gender specific)	Method										
*Thyroid Stimulating Hormone (TSH)	2.06	μIU/mL	0.4-4.2	CMIA										
<table border="1"> <tr> <td align="center" colspan="2">Biological Reference Intervals : TSH(μIU/mL)</td> </tr> <tr> <td align="center" colspan="2">Pregnancy (As per American Thyroid Association)</td> </tr> <tr> <td align="center">1 Trimester</td> <td align="center">0.10-2.50</td> </tr> <tr> <td align="center">2 Trimester</td> <td align="center">0.2-3.00</td> </tr> <tr> <td align="center">3 Trimester</td> <td align="center">0.3-3.00</td> </tr> </table>					Biological Reference Intervals : TSH(μIU/mL)		Pregnancy (As per American Thyroid Association)		1 Trimester	0.10-2.50	2 Trimester	0.2-3.00	3 Trimester	0.3-3.00
Biological Reference Intervals : TSH(μIU/mL)														
Pregnancy (As per American Thyroid Association)														
1 Trimester	0.10-2.50													
2 Trimester	0.2-3.00													
3 Trimester	0.3-3.00													

Interpretation:

- Assay results should be interpreted in context to the clinical condition and associated results of other investigations.
- Previous treatment with Corticosteroid therapy may result in lower TSH levels while Thyroid hormone levels are normal.
- Results are invalidated if the client has undergone a radionuclide scan within 7-14 days before the test.
- Abnormal Thyroid test findings often found in critically ill clients should be repeated after the critical nature of the condition is resolved.
- The production, circulation, and disposal of Thyroid hormone are altered throughout the stages of pregnancy.

Electrolyte Profile - Serum

Sodium (Na)	141	mmol/L	135 - 145	ISE Direct
Potassium (K)	4.9	mmol/L	3.5 - 5.1	ISE Direct
Chloride(CL)	104	mmol/L	98 - 107	ISE Direct

Comments:-

An electrolyte panel is often part of a routine blood screening or a comprehensive metabolic panel. The test may also be used to find out if your body has a fluid imbalance or an imbalance in acid and base levels.

Electrolytes are usually measured together. But sometimes they are tested individually. Separate testing may be done to suspect a problem with a specific electrolyte.

Result rechecked and verified for abnormal cases.

*** End Of Report ***



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Sample Type	: Serum	Report Status	: Final Report
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CLINICAL BIOCHEMISTRY

PATHCHECK ULTIMA

Test Name	Obtained Value	Units	Bio. Ref. Intervals (Age/Gender specific)	Method
Liver Function Test (LFT)				
Bilirubin Total	0.50	mg/dL	0.2-1.2	Diazotization
Bilirubin Direct	0.30	mg/dL	0.1-0.5	Diazotization
Bilirubin Indirect	0.2	mg/dL	0.2 - 1.0	Calculated
Alkaline Phosphatase (ALP)	54.0	U/L	46-116	Para-Nitrophenyl-phosphate
Aspartate Aminotransferase (SGOT)	17.0	U/L	5-34	NADH w/o P-5'-P
Alanine Transaminase (ALT/SGPT)	13.0	U/L	0.0-55	NADH w/o P-5'-P
Gamma Glutamyl Transferase (GGT)	13	U/L	9-36	L-g-g-3-Carboxy-4-Nitroanilide subs
Protein Total	6.8	g/dL	6.4-8.3	Biuret
Albumin	4.0	g/dL	3.5-5.2	Brom cresol Green
Globulin	2.8	g/dL	2.5 - 3.8	Calculated
Albumin / Globulin Ratio	1.4		1.0 - 2.1	Calculated
SGOT/SGPT Ratio	1	U/U	>2.0 suggests alcoholic liver disease.	Calculated

*Liver function tests are blood tests used to help diagnose and monitor Liver disease or damage. Screen for Liver infections, such as Hepatitis, monitor possible side effects of medications. Measure the severity of a disease, particularly scarring of the Liver (Cirrhosis)

*Alanine Transaminase (ALT)- an enzyme found in the Liver that helps your body metabolize protein. When the Liver is damaged, ALT is released into the bloodstream and levels increase.

*Aspartate Transaminase (AST)- an enzyme that helps metabolize Alanine, an amino acid. Like ALT, AST is normally present in blood at low levels. An increase in AST levels may indicate Liver damage or disease or Muscle damage.

*Alkaline Phosphatase (ALP)- an enzyme in the Liver, bile ducts and bone. Higher-than-normal levels of ALP may indicate liver damage or disease, such as a blocked bile duct, or certain bone diseases.

*Albumin and Total Protein- Albumin is one of several proteins made in the Liver. Your body needs these proteins to fight infections and to perform other functions. Lower-than-normal levels of albumin and total protein might indicate Liver damage or disease

*Bilirubin- a substance produced during the normal breakdown of red blood cells. Bilirubin passes through the liver and is excreted in stool. Elevated levels of bilirubin (jaundice) might indicate liver damage or disease or certain types of anemia.

*Gamma-Glutamyltransferase (GGT)- GGT is an enzyme in the blood. Higher-than-normal levels may indicate liver or bile duct damage.

*** End Of Report ***



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CLINICAL BIOCHEMISTRY

PATHCHECK ULTIMA

Test Name	Obtained Value	Units	Bio. Ref. Intervals (Age/Gender specific)	Method
Lipid Profile				
Cholesterol Total	135	mg/dL	Adult: Desirable<200 mg/dL, Borderline: 200 – 239 mg/dL, High:>240 mg/dL	Enzymatic
Cholesterol HDL	51	mg / dL	40 - 60	Direct Homogenous
Cholesterol - LDL	75	mg/dL	<100 Optimal	Calculated
Cholesterol VLDL	9	mg/dL	7-40	Calculated
Non-HDL cholesterol	84	mg/dL	Optimal < 130	Calculated
Triglycerides	45.0	mg/dL	Normal: <150 Borderline High: 150–199 High: 200–499 Very High: >500	Glycerol Phosphate Oxidase
Cholesterol Total/Cholesterol HDL Ratio	2.6		0 - 4.0	Calculated
Cholesterol LDL/Cholesterol HDL	1.5		0 - 3.5	Calculated

COMMENTS: Therapeutic target levels of lipids as per NCEP – ATP III recommendations:

Total Cholesterol (mg/dL)	<200 - Desirable, 200-239 - Borderline High, >240 - High
HDL Cholesterol (mg/dL)	<40 - Low, >60 - High
LDL Cholesterol (mg/dL)	<100 Optimal, [Primary Target of Therapy], 100-129 - Near Optimal/Above Optimal, 130-159 - Borderline High, 160-189 - High, >190 Very High
Serum Triglycerides (mg/dL)	<150 Normal, 150-199 Borderline High, 200-499 High, >500 Very High

NCEP recommends lowering of LDL Cholesterol as the primary therapeutic target with Lipid lowering agents, however, if Triglycerides remain >200 mg/dL after LDL goal is reached, set secondary goal for non-HDL Cholesterol (total minus HDL) 30 mg/dL higher than LDL goal.

When Triglyceride level is > 400 mg/dL, Friedewald Equation is not applicable for calculation of LDL & VLDL. Hence the calculated values are not provided for such samples.

ATP III Guidelines:

Risk Category	LDL Goal	LDL Level at Which to Initiate Therapeutic Lifestyle Changes (TLC)	LDL Level at Which to Consider Drug Therapy
CHD or CHD RiskEquivalents(10-year risk >20%)	<100 mg/dL	>100 mg/dL	>130 mg/dL (100-129 mg/dL: drug optional)*
2+ Risk Factors (10-year risk <20%)	<130 mg/dL	>130 mg/dL	10-year risk 10-20%: >130 mg/dL 10-year risk <10%:>160mg/dL
0-1 Risk Factor	<160 mg/dL	>160 mg/dL	>190 mg/dL (160-189 mg/dL: LDL-lowering drug optional)

*** End Of Report ***



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Note: If the test results are alarming or unexpected, Client is advised to contact the laboratory immediately for possible consultation.

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Sample Type	: Serum	Report Status	: Final Report
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CLINICAL BIOCHEMISTRY

PATHCHECK ULTIMA

Test Name	Obtained Value	Units	Bio. Ref. Intervals (Age/Gender specific)	Method
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Iron Deficiency Profile-II

Iron	116	µg/dL	50-170	Ferene
Iron Binding Capacity - Total (TIBC)	290	µg/dL	250-450	Calculated
Transferrin	202.5	mg/dL	176 - 280	Calculated
% Iron Saturation	40	%	20 - 50	Calculated
Iron Iron found in the blood is mainly present in the hemoglobin of the RBCs. Its role in the body is mainly in the transportation. Iron is absorbed in the Small Intestine, and bound to a globulin in the plasma called Transferrin and transported to the Bone Marrow for the formation of Hemoglobin. Increased serum levels are found in Hemolytic Anemias, Hepatitis, Lead and Iron poisoning. Decreased serum levels are found in Anemias caused by Iron Deficiency due to insufficient intake or absorption of Iron, chronic blood loss, late pregnancy and Cancer				
TIBC The serum TIBC varies in disorders of Iron metabolism. In Iron-deficiency Anemia the TIBC is elevated and the transferrin saturation is lowered to 15% or less. Low serum Iron associated with low TIBC is characteristic of the anemia of chronic disorders, malignant tumors, and infections.				

*Calcium	9.40	mg/dl	8.4-10.2	Arsenazo III Complex
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Interpretation:

Category	Normal Ref. Range
Premature	6.2 mg/dL to 11.0 mg/dL
0 to 10 days	7.6 mg/dL to 10.4 mg/dL
10 days to 24 months	9.0 mg/dL to 11.0 mg/dL
Child 2 to 12 years	8.8 mg/dL to 10.8 mg/dL
Adult	8.4 mg/dL to 10.2 mg/dL
Male > 60 years	8.8 mg/dL to 10.0 mg/dL

Comments:
 * Calcium in the body is found mainly in the bones (approximately 99%). In serum, Calcium exists in a free ionised form and in bound form (with Albumin). Hence, a decrease in Albumin causes lower Calcium levels and vice-versa.
 * Calcium levels in serum depend on the Parathyroid Hormone.
 * Increased Calcium levels are found in Bone tumors, Hyperparathyroidism. decreased levels are found in Hypoparathyroidism, renal failure, Rickets.

*Vitamin - B12	276.9	pg/mL	187-833	CMIA
Comments:				
<ul style="list-style-type: none"> Vitamin B12 is essential in DNA synthesis Hematopoiesis, and Central Nervous System integrity. Its absorption depends on the presence of intrinsic factor (IF) and may be due to lack of IF secretion by gastric mucosa. Vitamin B12 deficiency frequently causes Macrocytic Anemia, Glossitis, Peripheral Neuropathy, Weakness, Hyperreflexia, Ataxia, Loss of Proprioception, poor coordination and effective behavioural changes. A significant increase in RBC MCV may be an important indicator of Vitamin B12 deficiency. 				

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CLINICAL BIOCHEMISTRY

PATHCHECK ULTIMA

Test Name	Obtained Value	Units	Bio. Ref. Intervals (Age/Gender specific)	Method
Kidney Function Test (KFT) - III				
Creatinine	0.92	mg/dL	0.57-1.11	Kinetic Alkaline picrate
Urea	22.7	mg/dL	15.0-40.0	Calculated
Uric Acid	3.6	mg/dL	2.6-6.0	Uricase
Urea / Creatinine Ratio	24.67	mg/mg	Elevated ratio: >100:1 Reduced ratio: <40:1	Calculated

Correlate Clinically.

*** End Of Report ***



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