



PO No :PO2200904675-101



Customer Name	: Mr.TAPAN GUPTA ROY	Collected Via	: TATA 1MG KOLKATA
Age/Gender	: 69/Male	Referred By	: Dr.
Lab Visit ID	: KOL756355	Collection Date	: 10/Dec/2025 01:33PM
Barcode ID/Order ID	: D26822109 / 15113992	Report Date	: 10/Dec/2025 06:37PM
Sample Type	: Serum	Report Status	: Final Report

### BIOCHEMISTRY

Test Name	Result	Unit	Bio. Ref. Interval	Method
<b>Kidney Function Test.</b>				
Blood Urea Nitrogen	<b>68</b>	mg/dL	9.0 - 23.0	Urease with GLDH
Urea	<b>145.52</b>	mg/dL	19.26 - 49.22	Calculated
Creatinine	<b>8.64</b>	mg/dL	0.7-1.3	Alkaline picrate-kinetic
Uric Acid	5.4	mg/dL	3.5-7.2	Uricase/Peroxidase
Sodium	<b>135</b>	mmol/L	136-145	Indirect ISE
Potassium	4.46	mmol/L	3.5-5.1	Indirect ISE
Chloride	100.0	mmol/L	98-107	Indirect ISE
BUN/Creatinine Ratio	7.9	Ratio	12:1 - 20:1	Calculated

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**Critical Result-(Creatinine) . Please consult your doctor immediately.**

**Advice -Kindly correlate clinically.Repeat testing may be done on a fresh sample, if clinically indicated.**

#### Comment:

**BUN** is directly related to protein intake and nitrogen metabolism and inversely related to the rate of excretion of urea. Blood urea nitrogen (BUN) levels reflect the balance between the production and excretion of urea. Increased levels are seen in renal failure (acute or chronic), urinary tract obstruction, dehydration, shock, burns, CHF, GI bleeding, nephrotoxic drugs. Decreased levels are seen in hepatic failure, nephrotic syndrome, cachexia (low-protein and high-carbohydrate diets).

**Urea** is a non-proteinous nitrogen compound formed in the liver from ammonia as an end product of protein metabolism. Urea diffuses freely into extracellular and intracellular fluid and is ultimately excreted by the kidneys. Increased levels are found in acute renal failure, chronic glomerulonephritis, congestive heart failure, decreased renal perfusion, diabetes, excessive protein ingestion, gastrointestinal (GI) bleeding, hyperalimentation, hypovolemia, ketoacidosis, muscle wasting from starvation, neoplasms, pyelonephritis, shock, urinary tract obstruction, nephrotoxic drugs. Decreased levels are seen in inadequate dietary protein, low-protein/high-carbohydrate diet, malabsorption syndromes, pregnancy, severe liver disease, certain drugs.

**Creatinine** is catabolic product of creatinine phosphate, which is excreted by filtration through the glomerulus and by tubular secretion. Creatinine clearance is an acceptable clinical measure of glomerular filtration rate (GFR). Increased levels are seen in acute/chronic renal failure, urinary tract obstruction, hypothyroidism, nephrotoxic drugs, shock, dehydration, congestive heart failure, diabetes. Decreased levels are found in muscular dystrophy.

**BUN/Creatinine ratio** (normally 12:1-20:1) is decreased in acute tubular necrosis, advanced liver disease, low protein intake, and following hemodialysis. BUN/Creatinine ratio is increased in dehydration, GI bleeding, and increased catabolism.

**Uric acid** levels show diurnal variation. The level is usually higher in the morning and lower in the evening. Increased levels are seen in starvation, strenuous exercise, malnutrition, or lead poisoning, gout, renal disorders, increased breakdown of body cells in some cancers (including leukemia, lymphoma, and multiple myeloma) or cancer treatments, hemolytic anemia, sickle cell anemia, or heart failure, pre-eclampsia, liver disease (cirrhosis), obesity, psoriasis, hypothyroidism, low blood levels of parathyroid hormone (PTH), certain drugs, foods that are very high in purines - such as organ meats, red meats, some seafood and beer. Decreased levels are seen in liver disease, Wilson's disease, Syndrome of inappropriate antidiuretic hormone (SIADH), certain drugs.

#### Calcium

Calcium	9.7	mg/dL	8.7-10.4	Arsenazo III
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This test has been performed at

**TATA 1MG KOLKATA**

Address: Plot No. Y-17, Block - EP,  
Electronic Complex, Bidhannagar, Kolkata -  
700091

Dr Sulagna Ray Pal  
M.Sc, PhD (Biochemistry)  
Consultant Biochemist

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**Comment:**

**Increased in:** Hyperparathyroidism primary and secondary, Acute and chronic renal failure, Following renal transplantation, Osteomalacia with malabsorption, Acute osteoporosis, Malignant tumours (specially of breast, lung and kidney), Drugs: Vit. D and A intoxication, Diuretics, estrogen, androgen, tamoxifen, lithium

**Decreased in:** Hypoparathyroidism, Surgical and Idiopathic, Pseudohypoparathyroidism, Chronic renal disease with uremia and phosphate retention, Malabsorption of Calcium and Vit.D, obstructive jaundice, Bone Disease ( Osteomalacia and rickets ), Drugs: Cancer chemotherapy drugs, calcitonin, loop-actives diuretics, Hypomagnesemia, Hypoalbuminemia

### Phosphorus, Serum

Phosphorus	5.00	mg/dl	2.4 - 5.1	Phosphomolybdate
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**Comment:**

Phosphate metabolism is under the regulation of PTH, Vitamin D metabolites, and Fibroblast growth factor-23. Serum phosphate concentrations are about 50% higher in infants than in adults and decline throughout childhood as a consequence of the ability of growth hormone to increase the renal phosphate threshold.

**Increased in:**

Decreased Renal filtraton (Acute or chronic renal failure) or increased reabsorption (e.g., hypoparathyroidism)  
 Increased Phosphate load (e.g., Oral or iv administration, Phosphate-containing laxatives or enemas, Vitamin D intoxication)  
 Cell Lysis (e.g., hemolysis, leukemias, chemotherapy, rhabdomyolysis)  
 Bone disease (e.g., healing fractures, multiple myeloma, Paget disease, osteolytic tumors)  
 Genetic (e.g., Hypoparathyroidism, Tumoral calcinosis)

**Decreased in:**

Intracellular Shift (e.g., Oral or intravenous Glucose, Insulin, Diabetic ketoacidosis, Respiratory alkalosis, Alcoholism, Severe burns)  
 Lowered Renal Phosphate Threshold (e.g., Primary or secondary hyperparathyroidism, Renal tubular defects)  
 Decreased Intestinal Absorption (Malabsorption syndrome, Vitamin D deficiency) or increased loss (Vomiting, Diarrhea)  
 Drugs (e.g., Salicylate, Paracetamol, Estrogens, Diuretics, Bisphosphonates, Anticonvulsants, Phosphate binding antacids, Antiviral drugs etc)

**Note:**

Because a significant diurnal variation in plasma phosphate has been reported, fasting morning specimens are recommended. Levels are influenced by dietary intake, meals, and exercise.

**\*\*\* End Of Report \*\*\***
**Disclaimer:**

1. The reported results based on laboratory investigation, are only for the purposes of diagnosis and should be clinically correlated and interpreted by the referring physician/ medical practitioner. For any queries relating to the reported results, you may write to our customer support team on care@1mg.com  
 2. It is presumed that the tests performed are, on the specimen / sample being to the patient named or identified and the verifications of particulars have been confirmed by the patient or his / her representative at the point of generation of said specimen.



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 700091



 Dr Sulagna Ray Pal  
 M.Sc, PhD (Biochemistry)  
 Consultant Biochemist


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4. The patient's details along with their results in certain cases like notifiable diseases and as per local regulatory requirements will be communicated to the assigned regulatory bodies.				
5. The patient samples can be used as part of internal quality control, test verification, data analysis purposes within the testing scope of the laboratory.				
6. This report is not valid for medico legal purposes. It is performed to facilitate medical diagnosis only.				
7. Pregnant women should seek guidance from a qualified obstetrician as test parameters may vary during pregnancy				

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**TATA 1mg | Labs**



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