

Patient Information	Specimen Information	Client / Doctor Information
Name : Mr.P SRINIVAS	Visit ID : LKOQ32523	Client Code : PUP4521
Age/Gender : 54 Y 0 M 0 D /Male	Collected : 05/Jun/2026 13:26	Client Name : PUP SARTHAK
MobileNo : 7903824101	Received : 05/Jun/2026 13:30	DIAGNOSTICS
UHID : LDAA05807088	Reported : 05/Jun/2026 14:50	Client Add. : GOLMURI
Address :	IP/OP/Barcode :	Client No. : 9939796234
	Report Status : Final Report	Ref Doctor : Dr.SELF

KEY CLINICAL INDICATORS

Test Name	Result	Ref. Range	Unit	Test Name	Result	Ref. Range	Unit
Kidney/Renal Function Tests Gold (KFT Gold /RFT Gold)				Kidney/Renal Function Tests Gold (KFT Gold /RFT Gold)			
◆ Creatinine	3.58 H	0.66-1.25	mg/dL	● Potassium	3.89	3.5-5.1	mmol/L
◆ Urea	44.7 H	19.0-43.0	mg/dL	● Chloride	103.0	98 - 107	mmol/L
◆ Blood Urea Nitrogen (BUN)	20.9 H	9.0 - 20.0	mg/dL	LIPID PROFILE			
● Uric Acid	5.81	3.7-7.7	mg/dL	◆ Total Cholesterol	210.00 H	<200	mg/dL
● Total Protein	7.28	6.3-8.2	g/dL	◆ Triglycerides	181.00 H	<150	mg/dL
◆ Sodium	136.8 L	137-145	mmol/L	◆ HDL Cholesterol	34.00 L	40-60	mg/dL
				LDL Cholesterol,	139.80 H	<100	mg/dL
				◆ Calculated			

● Round shaped values are normal ◆ Crystal shaped values are abnormal



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Lupin Diagnostics Limited, 3rd Floor,
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Santacruz (East), Mumbai-400 055, India.
CIN: U24100MH2011PLC214885



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Test Name	Result	Bio. Ref. Range	Unit	Method
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Kidney/Renal Function Tests Gold (KFT Gold /RFT Gold) , SERUM

Creatinine	3.58 H	0.66-1.25	mg/dL	Creatinine amidohydrolase
Urea	44.7 H	19.0-43.0	mg/dL	Urease
Blood Urea Nitrogen (BUN)	20.9 H	9.0 - 20.0	mg/dL	Urease
Uric Acid	5.81	3.7-7.7	mg/dL	Uricase
Total Protein	7.28	6.3-8.2	g/dL	Biuret
Albumin	4.46	3.5 - 5	g/dL	Bromocresol Green
Globulin	2.82	2-3.5	g/dL	Calculated
Albumin : Globulin Ratio (A:G Ratio)	1.58	1.1-2.5		Calculated
Calcium	9.5	8.4 - 10.2	mg/dL	Arsenazo
Sodium	136.8 L	137-145	mmol/L	Direct ISE
Potassium	3.89	3.5-5.1	mmol/L	Direct ISE
Chloride	103.0	98 - 107	mmol/L	Direct ISE

LIPID PROFILE , SERUM

Total Cholesterol	210.00 H	<200	mg/dL	CHE/CHO/POD
Triglycerides	181.00 H	<150	mg/dL	Enzymatic
HDL Cholesterol	34.00 L	40-60	mg/dL	CHE/CHO/POD
LDL Cholesterol, Calculated	139.80 H	<100	mg/dL	Calculated
VLDL	36.20 H	<30	mg/dL	Calculated
Non HDL Cholesterol	176.00 H	<130	mg/dL	Calculated
Cholesterol : HDL Ratio	6.18 H	<5.0		Calculated

Interpretation:

The National Lipid Association and the National Cholesterol Education Program (NCEP) have set the following guidelines for lipids i.e. total cholesterol (in children, ages less than 18 years and adults ages 18 years and up), triglycerides, HDL cholesterol, LDL cholesterol, and non-HDL cholesterol:

Parameter	Desirable (mg/dL)	Borderline high (mg/dL)	High (mg/dL)	Very high (mg/dL)
Total Cholesterol Adults (>18 Yrs)	<200	200-239	>240	
Total Cholesterol Children (<18 Yrs)	<170	170-199	>200	
Triglycerides	<150	150-199	200-499	>500
LDL Cholesterol	Optimal <100 Above Optimal: 100 - 129	130-159	>160	
Non-HDL Cholesterol	Optimal <130 Above Optimal: 130 - 159			
Total Cholesterol/HDL Ratio	Male<5 Female<4.50			
LDL/HDL Ratio	Male <3.60 Female <3.20			

	Desirable (mg/dL)	Borderline risk (mg/dL)	Undesirable (mg/dL)	
HDL Cholesterol	>60	40-59	<40	

This test has been performed at Lupin Diagnostics Laboratory, RLM AAROYAMM DIAGNOSTICS 4TH FLOOR DAYAL TRADE CENTRE, SHER E PUNJAB CHOWK, Sher E Punjab Chowk, SARAIKELA KHARSAWAN , 831013



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If the testing is performed in a non-fasting state, only the values for Total Cholesterol and HDL Cholesterol are considered reliable. If Total Cholesterol is >200mg/dl or HDL Cholesterol is <40 mg/dl in non-fasting sample, a follow-up fasting lipoprotein profile is recommended to accurately assess LDL Cholesterol levels for appropriate clinical management. Measurements in the same patient can show physiological and analytical variations. NCEP ATP III identifies non-HDL cholesterol as a secondary target of therapy in persons with high triglycerides				

If Values are marked with * , they are critical values.

*** End Of Report ***

B. Baidya
Dr B. BAITHA
MD Pathology
Senior Consultant Pathologist
REG NO- 32908



This test has been performed at Lupin Diagnostics Laboratory, RLM AAROYAMM DIAGNOSTICS 4TH FLOOR DAYAL TRADE CENTRE, SHER E PUNJAB CHOWK, LUDHIANA, PUNJAB, INDIA. 141005



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Conditions of Reporting

1. The contents of this report are to be forwarded to the Referring Doctor only for the purpose of assisting in his/her diagnosis and management of the patient.
2. Test results relate only to the specimen submitted.
3. Unless the collection state is specified, the current date will automatically be printed as the collection date.
4. When tests are performed on specimens received from locations outside of Lupin Diagnostics, it is presumed that the test sample belongs to the patient named or identified in the form. This authentication is done at the time the specimen is generated.
5. Observations/Technical notes about the test including its nature, its essential interpretative value, and any other relevant information regarding performance, results, and follow-up information are given below the test heading interpretation.
6. A requested test might not be performed if:
 - a. Test results will appear with the message "QTY. NOT ADEQUATE" if the sample received is insufficient for performing the requested test(s).
 - b. Test results will appear with the message "TEST NOT PERFORMED" if the specimen quality is unacceptable (Hemolysed/ Clotted/Lipemic etc.) or the specimen type is incorrect. The exact reason will appear as a comment at the end of the report.
 - c. In case where test is cancelled either on request of the patient or doctor, or for incorrect test code/test name, the response - "TEST CANCELLED" will appear against the test name and the reason will appear as a comment at the end of the report.
7. For various technical reasons, a test requested might yield "INCONCLUSIVE RESULTS". This response will appear against the test name, followed by a detailed comment at the end of the report.
8. "PROVISIONAL" status of the report signifies that the results are yet to be authenticated for some reason.
9. A repeat investigation/additional investigation for clinical samples and histopathology specimens will be possible, as per our sample retention policy.
10. Assays are performed in accordance with standard procedure on receipt. The laboratory test results depend on the sample's quality, assay technology, equipment used, method specificity, and sensitivity. All investigations have their limitations, and isolated laboratory investigations are not the final diagnosis of disease. They only help in arriving at a diagnosis in conjunction with clinical findings and relevant investigations. The reports are therefore expected to be interpreted, taking these facts into consideration.
11. We perform and report all tests as per the testing schedule. Results may be delayed for reasons beyond our control. Lupin Diagnostics will make all efforts to resolve these issues.
12. Any questions from the Referring Physician regarding this report should be directed to: customerconnect@lupindiagnosics.com
13. Lupin Diagnostics continually strives to conduct each test with clinical safety, care and technical integrity. However, considering that test are subject to factors such as reagent variations, machine breakdowns etc. beyond the control of Lupin Diagnostics, there are no implied representations or warranties given on the accuracy of the reports/results. If the results indicate an unexpected abnormality, the same should be retested.
- 14.. The reports/results are to be used to assist in diagnosing medical diseases and not for forensic applications or medico-legal purposes.
15. The total liability of Lupin Diagnostics is limited to the total amount of the invoice paid, for the services rendered.

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