

A PIONEER DIAGNOSTIC CENTRE

■ 0171-2532620, 8222896961 **■** pkrjainhealthcare@gmail.com

NAME : Mr. PARDEEP

AGE/ GENDER : 26 YRS/MALE **PATIENT ID** :1753983

COLLECTED BY REG. NO./LAB NO. : 122502120005

REFERRED BY **REGISTRATION DATE** : 12/Feb/2025 08:57 AM BARCODE NO. **COLLECTION DATE** : 12/Feb/2025 09:27AM : 12506967 CLIENT CODE. : P.K.R JAIN HEALTHCARE INSTITUTE REPORTING DATE : 12/Feb/2025 01:30PM

CLIENT ADDRESS : NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYANA

Value Test Name Unit **Biological Reference interval**

CLINICAL CHEMISTRY/BIOCHEMISTRY

LIPID PROFILE: BASIC

CHOLESTEROL TOTAL: SERUM OPTIMAL: < 200.0 287.24^H mg/dL

by CHOLESTEROL OXIDASE PAP BORDERLINE HIGH: 200.0 -

2390

HIGH CHOLESTEROL: > OR =

240.0

TRIGLYCERIDES: SERUM OPTIMAL: < 150.0 230.54^H mg/dL by GLYCEROL PHOSPHATE OXIDASE (ENZYMATIC) BORDERLINE HIGH: 150.0 -

199.0

HIGH: 200.0 - 499.0

VERY HIGH: > OR = 500.0

HDL CHOLESTEROL (DIRECT): SERUM LOW HDL: < 30.0 44.86 mg/dL

by SELECTIVE INHIBITION

BORDERLINE HIGH HDL: 30.0 -

60.0

 $HIGH\ HDL: > OR = 60.0$ LDL CHOLESTEROL: SERUM OPTIMAL: < 100.0 196.27^H mg/dL

by CALCULATED, SPECTROPHOTOMETRY ABOVE OPTIMAL: 100.0 - 129.0

BORDERLINE HIGH: 130.0 -

HIGH: 160.0 - 189.0 VERY HIGH: > OR = 190.0

 242.38^{H} mg/dL OPTIMAL: < 130.0

ABOVE OPTIMAL: 130.0 - 159.0

BORDERLINE HIGH: 160.0 -

HIGH: 190.0 - 219.0

VERY HIGH: > OR = 220.0

mg/dL VLDL CHOLESTEROL: SERUM 46.11^H 0.00 - 45.00

by CALCULATED. SPECTROPHOTOMETRY

TOTAL LIPIDS: SERUM 350.00 - 700.00 mg/dL 805.02^H by CALCULATED, SPECTROPHOTOMETRY

6.4^H CHOLESTEROL/HDL RATIO: SERUM RATIO LOW RISK: 3.30 - 4.40 by CALCULATED, SPECTROPHOTOMETRY

AVERAGE RISK: 4.50 - 7.0



DR.VINAY CHOPRA CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY & MICROBIOLOGY)

DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY)



NON HDL CHOLESTEROL: SERUM

by CALCULATED, SPECTROPHOTOMETRY





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Test Name	Value	Unit	Biological Reference interval
LDL/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	4.38 ^H	RATIO	MODERATE RISK: 7.10 - 11.0 HIGH RISK: > 11.0 LOW RISK: 0.50 - 3.0 MODERATE RISK: 3.10 - 6.0 HIGH RISK: > 6.0
TRIGLYCERIDES/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	5.14 ^H	RATIO	3.00 - 5.00

INTERPRETATION:

1. Measurements in the same patient can show physiological analytical variations. Three serial samples 1 week apart are recommended for

Total Cholesterol, Triglycerides, HDL & LDL Cholesterol.

2. As per NLA-2014 guidelines, all adults above the age of 20 years should be screened for lipid status. Selective screening of children above the age of 2 years with a family history of premature cardiovascular disease or those with at least one parent with high total cholesterol is recommended.

3. Low HDL levels are associated with increased risk for Atherosclerotic Cardiovascular disease (ASCVD) due to insufficient HDL being available

to participate in reverse cholesterol transport, the process by which cholesterol is eliminated from peripheral tissues.

4. NLA-2014 identifies Non HDL Cholesterol (an indicator of all atherogeniclipoproteins such as LDL, VLDL, IDL, Lpa, Chylomicron remnants) along with LDL-cholesterol as co- primary target for cholesterol lowering therapy. Note that major risk factors can modify treatment goals for LDL &Non

5. Additional testing for Apolipoprotein B, hsCRP,Lp(a) & LP-PLA2 should be considered among patients with moderate risk for ASCVD for risk refinement



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Test Name	Value	Unit	Biological Reference interval		
KIDNEY FUNCTION TEST (BASIC)					
UREA: SERUM by UREASE - GLUTAMATE DEHYDROGENASE (GLDH)	33.89	mg/dL	10.00 - 50.00		
CREATININE: SERUM by ENZYMATIC, SPECTROPHOTOMETERY	1.19	mg/dL	0.40 - 1.40		
BLOOD UREA NITROGEN (BUN): SERUM by CALCULATED, SPECTROPHOTOMETERY	15.84	mg/dL	7.0 - 25.0		
BLOOD UREA NITROGEN (BUN)/CREATININE RATIO: SERUM by CALCULATED, SPECTROPHOTOMETERY	13.31	RATIO	10.0 - 20.0		
UREA/CREATININE RATIO: SERUM by CALCULATED, SPECTROPHOTOMETERY	28.48	RATIO			
URIC ACID: SERUM by URICASE - OXIDASE PEROXIDASE	5.08	mg/dL	3.60 - 7.70		



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Test Name Value Unit **Biological Reference interval**

INTERPRETATION:

Normal range for a healthy person on normal diet: 12 - 20

To Differentiate between pre- and postrenal azotemia. INCREASED RATIO (>20:1) WITH NORMAL CREATININE:

1. Prerenal azotemia (BUN rises without increase in creatinine) e.g. heart failure, salt depletion, dehydration, blood loss) due to decreased glomerular filtration rate.

Ž.Catabolic states with increased tissue breakdown.

3.GI hemorrhage.

4. High protein intake.

5. Impaired renal function plus.

6. Excess protein intake or production or tissue breakdown (e.g. infection, GI bleeding, thyrotoxicosis, Cushings syndrome, high protein diet,

burns, surgery, cachexia, high fever)

7. Urine reabsorption (e.g. ureterocolostomy)
8. Reduced muscle mass (subnormal creatinine production)
9. Certain drugs (e.g. tetracycline, glucocorticoids)
INCREASED RATIO (pia (PLIN rices diegrapartic particular partic

1. Postrenal azotemia (BUN rises disproportionately more than creatinine) (e.g. obstructive uropathy).

2. Prerenal azotemia superimposed on renal disease.

DECREASED RATIO (<10:1) WITH DECREASED BUN:

1.Acute tubular necrosis.

2.Low protein diet and starvation.

3. Severe liver disease.

4. Other causes of decreased urea synthesis.

5. Repeated dialysis (urea rather than creatinine diffuses out of extracellular fluid).

6.Inherited hyperammonemias (urea is virtually absent in blood)

7.SIADH (syndrome of inappropiate antidiuretic harmone) due to tubular secretion of urea.

8. Pregnancy

DECREASED RATIO (<10:1) WITH INCREASED CREATININE:

- 1. Phenacimide therapy (accelerates conversion of creatine to creatinine).
- 2. Rhabdomyolysis (releases muscle creatinine).
- 3. Muscular patients who develop renal failure

INAPPROPIATE RATIO

1. Diabetic ketoacidosis (acetoacetate causes false increase in creatinine with certain methodologies, resulting in normal ratio when dehydration should produce an increased BUN/creatinine ratio).

2. Cephalosporin therapy (interferes with creatininé measurement).

*** End Of Report ***



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