PKR JAIN HEALTHCARE INSTITUTE NASIRPUR, Hissar Road, AMBALA CITY- (Haryana)

A PIONEER DIAGNOSTIC CENTRE

🔽 0171-2532620, 8222896961 🛛 🖾 pkrjainhealthcare@gmail.com

: 70 YRS/FEMALE	РАТ	TENT ID	: 1223485	
	REG	. NO./LAB NO.	: 122503150004	
	REG	ISTRATION DATE	: 15/Mar/2025 09:09 AM	
12507503	COL	LECTION DATE	: 15/Mar/2025 10:27AM	
: P.K.R JAIN HEALTHCARE INSTITUTE		ORTING DATE	: 15/Mar/2025 12:15PM	
NASIRPUR, HISSAR ROAD, AM	MBALA CITY - HARYAN	NA		
	Value	Unit	Biological Reference interv	
CLINIC	CAL CHEMISTRY	Y/BIOCHEMIST	'RY	
	GLUCOSE FAS	STING (F)		
F): PLASMA PEROXIDASE (GOD-POD)	114.51 ^H	mg/dL	NORMAL: < 100.0 PREDIABETIC: 100.0 - 125.	
	P.K.R JAIN HEALTHCARE INS NASIRPUR, HISSAR ROAD, AN CLINIC	REG 12507503 COL P.K.R JAIN HEALTHCARE INSTITUTE REP NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYAN Value CLINICAL CHEMISTRY GLUCOSE FAS F): PLASMA 114.51 ^H	P.K.R JAIN HEALTHCARE INSTITUTE REPORTING DATE NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYANA	

A fasting plasma glucose level below 100 mg/dl is considered normal.
 A fasting plasma glucose level between 100 - 125 mg/dl is considered as glucose intolerant or prediabetic. A fasting and post-prandial blood test (after consumption of 75 gms of glucose) is recommended for all such patients.
 A fasting plasma glucose level of above 125 mg/dl is highly suggestive of diabetic state. A repeat post-prandial is strongly recommended for all such patients.
 A fasting plasma glucose level in excess of 125 mg/dl on both occasions is confirmatory for diabetic state.





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

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

440 Dated 17.5.2012 u/s 80 G OF INCOME TAX ACT. PAN NO. AAAAP1600. **REPORT ATTRACTS THE CONDITIONS PRINTED OVERLEAF (P.T.O.)**



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NAME	: Mrs. PARVESH KUMARI					
AGE/ GENDER	: 70 YRS/FEMALE		PATIENT ID	: 1223485		
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BARCODE NO.	: 12507503		COLLECTION DATE	: 15/Mar/2025 10:27AM		
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CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYANA					
Test Name		Value	Unit	Biological Reference interval		
		LIPID PR	OFILE : BASIC			
CHOLESTEROL TO by CHOLESTEROL O>		241.42 ^H	mg/dL	OPTIMAL: < 200.0 BORDERLINE HIGH: 200.0 - 239.0 HIGH CHOLESTEROL: > OR = 240.0		
TRIGLYCERIDES: S by GLYCEROL PHOSF	ERUM PHATE OXIDASE (ENZYMATIC)	217.62 ^H	mg/dL	OPTIMAL: < 150.0 BORDERLINE HIGH: 150.0 - 199.0 HIGH: 200.0 - 499.0 VERY HIGH: > OR = 500.0		
HDL CHOLESTERO by SELECTIVE INHIBIT	L (DIRECT): SERUM Ton	40.9	mg/dL	LOW HDL: < 30.0 BORDERLINE HIGH HDL: 30.0 60.0 HIGH HDL: > OR = 60.0		
LDL CHOLESTERO by CALCULATED, SPE		157 ^H	mg/dL	OPTIMAL: < 100.0 ABOVE OPTIMAL: 100.0 - 129 BORDERLINE HIGH: 130.0 - 159.0 HIGH: 160.0 - 189.0 VERY HIGH: > OR = 190.0		
NON HDL CHOLES' by CALCULATED, SPE		200.52 ^H	mg/dL	OPTIMAL: < 130.0 ABOVE OPTIMAL: 130.0 - 159 BORDERLINE HIGH: 160.0 - 189.0 HIGH: 190.0 - 219.0 VERY HIGH: > OR = 220.0		
VLDL CHOLESTER		43.52	mg/dL	0.00 - 45.00		
TOTAL LIPIDS: SERUM by CALCULATED, SPECTROPHOTOMETRY CHOLESTEROL/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY		700.46 ^H	mg/dL	350.00 - 700.00		
		5.9 ^H	RATIO	LOW RISK: 3.30 - 4.40 AVERAGE RISK: 4.50 - 7.0 MODERATE RISK: 7.10 - 11.0 HIGH RISK: > 11.0		



TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT

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NOT VALID FOR MEDICO LEGAL PURPOSE

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Test Name	Value	Unit	Biological Reference interval
LDL/HDL RATIO: SERUM by calculated, spectrophotometry	3.84 ^H	RATIO	LOW RISK: 0.50 - 3.0 MODERATE RISK: 3.10 - 6.0 HIGH RISK: > 6.0
TRIGLYCERIDES/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	5.32 ^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 HDL.

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

*** End Of Report





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