



	Dr. Vinay Chc MD (Pathology & I Chairman & Consu	Microbiology)		m Chopra D (Pathology) ht Pathologist
NAME	: Mr. RAJBARINDER SINGH			
AGE/ GENDER	: 60 YRS/MALE	P	ATIENT ID	: 1679890
COLLECTED BY	: SURJESH	R	EG. NO./LAB NO.	: 012411230021
REFERRED BY	: CENTRAL PHOENIX CLUB (AM	(BALA CANTT)	EGISTRATION DATE	: 23/Nov/2024 09:55 AM
BARCODE NO.	: 01521295		OLLECTION DATE	: 23/Nov/2024 09:57AM
CLIENT CODE.	: KOS DIAGNOSTIC LAB		EPORTING DATE	: 23/Nov/2024 03:10PM
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, AMBALA CANTT			
Test Name		Value	Unit	Biological Reference interval
ESTIMATED AVERA	RMANCE LIQUID CHROMATOGRAPHY) IGE PLASMA GLUCOSE RMANCE LIQUID CHROMATOGRAPHY)	119.76	mg/dL	60.00 - 140.00
	AS PER AMERICAN D	DIABETES ASSOCIAT	rion (Ada):	
	REFERENCE GROUP		GLYCOSYLATED HEMOGLOGIB (HBAIC) in %	
	abetic Adults >= 18 years	<5.7		
	At Risk (Prediabetes) Diagnosing Diabetes		5.7 - 6.4 >= 6.5	
Therapeutic goals for glycemic control		Actions	Age > 19 Years f Therapy: Suggested: Age < 19 Years	< 7.0 >8.0
2.Since Hb1c reflects Ic concentration of HbAI 3.Target goals of < 7.0 patients with significal appropiate. 4.High HbA1c (>9.0 -9	ong term fluctuations in blood glucos lc. Converse is true for a diabetic prev 9 % may be beneficial in patients with nt complications of diabetes, limited l	monitoring done t e concentration, a c iously under good o short duration of c ife expectancy or e k of development	diabetic patient who has r control but now poorly con liabetes, long life expectan xtensive co-morbid condit and rapid progression of	ncy and no significant cardiovascular disease. In ions, targetting a goal of < 7.0% may not be microvascular and nerve complications

6.HbA1c results from patients with HbSS,HbSC and HbD must be interpreted with caution, given the pathological processes including anemia, increased red cell turnover, and transfusion requirement that adversely impact HbA1c as a marker of long-term gycemic control.

7.Specimens from patients with polycythemia or post-splenctomy may exhibit increse in HbA1c values due to a somewhat longer life span of the red cells.



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		hopra & Microbiology) nsultant Pathologis	Dr. Yugam Chopra MD (Pathology) t CEO & Consultant Pathologist	
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Test Name		Value	Unit	Biological Reference interval
CHOLESTEROL TOT by CHOLESTEROL OX	TAL: SERUM		STRY/BIOCHEMIST OFILE : BASIC mg/dL	OPTIMAL: < 200.0 BORDERLINE HIGH: 200.0 - 239.0 HIGH CHOLESTEROL: > OR =
FRIGLYCERIDES: SI by GLYCEROL PHOSPI	ERUM HATE OXIDASE (ENZYMATIC)	191.8 ^H	mg/dL	240.0 OPTIMAL: < 150.0 BORDERLINE HIGH: 150.0 - 199.0 HIGH: 200.0 - 499.0 VERY HIGH: > OR = 500.0
HDL CHOLESTEROL by SELECTIVE INHIBITI	L (DIRECT): SERUM ON	41.96	mg/dL	LOW HDL: < 30.0 BORDERLINE HIGH HDL: 30.0 60.0 HIGH HDL: > OR = 60.0
LDL CHOLESTEROL by CALCULATED, SPEC		49.54	mg/dL	OPTIMAL: < 100.0 ABOVE OPTIMAL: 100.0 - 129.0 BORDERLINE HIGH: 130.0 - 159.0 HIGH: 160.0 - 189.0 VERY HIGH: > OR = 190.0
NON HDL CHOLEST by CALCULATED, SPEC		87.9	mg/dL	OPTIMAL: < 130.0 ABOVE OPTIMAL: 130.0 - 159.0 BORDERLINE HIGH: 160.0 - 189.0 HIGH: 190.0 - 219.0 VERY HIGH: > OR = 220.0
VLDL CHOLESTERO		38.36	mg/dL	0.00 - 45.00
TOTAL LIPIDS: SER by CALCULATED, SPEC	UM	451.52	mg/dL	350.00 - 700.00
	L RATIO: SERUM	3.09	RATIO	LOW RISK: 3.30 - 4.40





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TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.





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Test Name		Value	Unit	Biological Reference interval			
LDL/HDL RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY		1.18	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		4.57	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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