



	Dr. Vinay Cho MD (Pathology & Chairman & Cons	Microbiology)	Dr. Yugarı MD CEO & Consultant	(Pathology)
NAME	: Mrs. MUNNI			
AGE/ GENDER	: 45 YRS/FEMALE	P	ATIENT ID	: 1576451
COLLECTED BY	: SURJESH	R	EG. NO./LAB NO.	: 012408100022
<b>REFERRED BY</b>	:	R	EGISTRATION DATE	: 10/Aug/2024 10:19 AM
BARCODE NO.	:01514819	С	OLLECTION DATE	: 10/Aug/2024 12:45PM
CLIENT CODE.	: KOS DIAGNOSTIC LAB	R	EPORTING DATE	: 10/Aug/2024 02:14PM
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, A	MBALA CANTT		Ŭ
Test Name		Value	Unit	Biological Reference interval
	GL		TOLOGY MOGLOBIN (HBA1C)	
GLYCOSYLATED HAEM		6.6 <sup>H</sup>	%	4.0 - 6.4
WHOLE BLOOD by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) ESTIMATED AVERAGE PLASMA GLUCOSE by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) INTERPRETATION:		142.72 <sup>H</sup> mg.		60.00 - 140.00
	AS PER AMERICAN DIABI			
	FERENCE GROUP	GLYCOSYLA	TED HEMOGLOGIB (HBAIC) i	n %
	etic Adults >= 18 years	<5.7		
At Risk (Prediabetes) Diagnosing Diabetes		5.7 - 6.4 >= 6.5		
Dia	gnosing Diabetes		>= 0.5 Age > 19 Years	
Therapeutic goals for glycemic control		Goals of Thera		)
		Actions Suggest		
Therapeutic		Age < 19 Years		
Therapeutic			Aye < 19 reals	

2. Since Hb1c reflects long term fluctuations in blood glucose concentration, a diabetic patient who has recently under good control may still have high concentration of HbAlc. Converse is true for a diabetic previously under good control but now poorly controlled.

3. Target goals of < 7.0 % may be beneficial in patients with short duration of diabetes, long life expectancy and no significant cardiovascular disease. In patients with significant complications of diabetes, limited life expectancy or extensive co-morbid conditions, targetting a goal of < 7.0% may not be 4.High appropiate.

HbA1c (>9.0 -9.5 %) is strongly associated with risk of development and rapid progression of microvascular and nerve complications

5.Any condition that shorten RBC life span like acute blood loss, hemolytic anemia falsely lower HbA1c results.

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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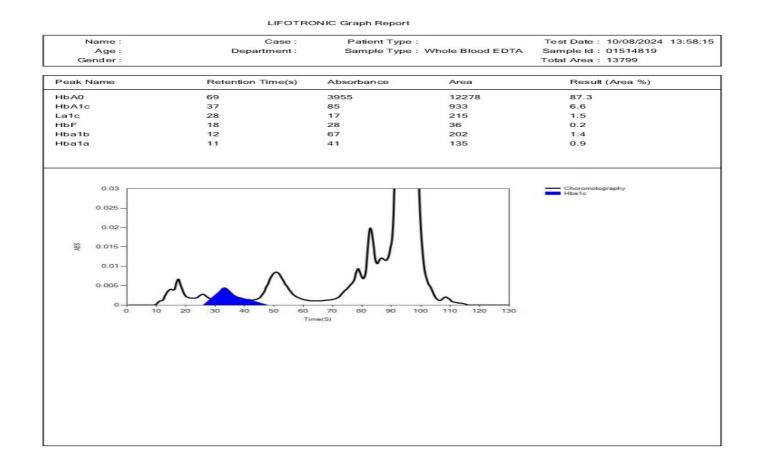
TEST PERFORMED AT KOS DIAGNOSTIC LAB. AMBALA CANTT





<b>c Lab</b> althcare)		EXCELLENCE IN HEALTHCARE & DIAGNOSTICS	
	1	Du Vuenn Channe	

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Test Name	Value	Unit	Biological Reference interval







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		<b>Chopra</b> gy & Microbiology) Consultant Pathologist		(Pathology)	
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CLIENT CODE.	: KOS DIAGNOSTIC LAB	<b>REPORTING DATE</b>		: 10/Aug/2024 02:44PM	
CLIENT ADDRESS	DRESS : 6349/1, NICHOLSON ROAD, AMBALA CANTT				
	. 0343/ 1, MCHOLSON ROP				
Test Name	. 0040/1, NICHOLSON ROP	Value	Unit	Biological Reference interval	
		Value	Unit TRY/BIOCHEMISTR	-	
	CL	Value		Y	
Test Name GLUCOSE FASTING (F	CL GLUC	Value	TRY/BIOCHEMISTR	Y	

## IN ACCORDANCE WITH AMERICAN DIABETES ASSOCIATION GUIDELINES:

**KOS Diagnostic Lab** 

(A Unit of KOS Healthcare)

 A fasting plasma glucose below 100 mg/dL and post-prandial plasma glucose level below 140 mg/dl is considered normal.
 A fasting plasma glucose level between 100 - 125 mg/dl and post-prandial plasma glucose level between 140 - 200 mg/dL is considered as glucose intolerant or pre diabetic. A fasting and post-prandial blood test (after consumption of 75 gms of glucose) is recommended for all such patients

3. A fasting plasma glucose level of above 125 mg/dL and post-prandial plasma glucose level above 200 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.



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:01514819	COL	LECTION DATE	: 10/Aug/2024 12:45PM
: KOS DIAGNOSTIC LAB	REP	ORTING DATE	: 10/Aug/2024 11:36AM
: 6349/1, NICHOLSON RO	DAD, AMBALA CANTT		
	Value	Unit	Biological Reference interval
	CHOLESTERO	.: SERUM	
: SERUM dase pap	169.76	mg/dL	OPTIMAL: < 200.0 BORDERLINE HIGH: 200.0 - 239.0 HIGH CHOLESTEROL: > OR = 240.0
	MD (Pathol Chairman 8 : Mrs. MUNNI : 45 YRS/FEMALE : SURJESH : : 01514819 : KOS DIAGNOSTIC LAB : 6349/1, NICHOLSON RO	: 45 YRS/FEMALE PAT : SURJESH REG : REG : 01514819 COL : KOS DIAGNOSTIC LAB REP : 6349/1, NICHOLSON ROAD, AMBALA CANTT Value CHOLESTEROI : SERUM 169.76	MD (Pathology & Microbiology) Chairman & Consultant Pathologist DEO & Consultant : Mrs. MUNNI : 45 YRS/FEMALE PATIENT ID : SURJESH REG. NO./LAB NO. : 01514819 COLLECTION DATE : 603 DIAGNOSTIC LAB REPORTING DATE : 6349/1, NICHOLSON ROAD, AMBALA CANTT Value Unit CHOLESTEROL: SERUM : SERUM 169.76 mg/dL

NATIONAL LIPID ASSOCIATION RECOMMENDATIONS (NLA-2014)	CHOLESTEROL IN ADULTS (mg/dL)	CHOLESTEROL IN ADULTS (mg/dL)
DESIRABLE	< 200.0	< 170.0
BORDERLINE HIGH	200.0 – 239.0	171.0 - 199.0
HIGH	>= 240.0	>= 200.0

## NOTE:

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.
 As per National Lipid association - 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.



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		URIC ACI	)	
URIC ACID: SERUM		4.61	mg/dL	2.50 - 6.80
by URICASE - OXIDAS	E PEROXIDASE		5	
<ul> <li>4.Polycythemai vera</li> <li>5.Psoriasis.</li> <li>6.Sickle cell anaemia</li> <li>(B).DUE TO DECREASE</li> <li>1.Alcohol ingestion.</li> <li>2.Thiazide diuretics.</li> <li>3.Lactic acidosis.</li> <li>4.Aspirin ingestion (li</li> <li>5.Diabetic ketoacido</li> <li>6.Renal failure due to</li> <li>DECREASED:-</li> <li>(A).DUE TO DIETARY LE</li> <li>1.Dietary deficiency of</li> <li>2.Fanconi syndrome</li> <li>3.Multiple sclerosis.</li> <li>4.Syndrome of inappi</li> <li>(B).DUE TO INCREASE</li> </ul>	D EXCREATION (BY KIDNEYS) ess than 2 grams per day ). sis or starvation. o any cause etc. DEFICIENCY of Zinc, Iron and molybdenum. & Wilsons disease. ropriate antidiuretic hormone (SIA D EXCREATION	ADH) secretion & low pu		ds and ACTH, anti-coagulants and estrogens et
		** End Of Report		as and norm, and boughting and ostrogens et





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