



	Dr. Vinay Ch MD (Pathology & Chairman & Con			(Pathology)
IAME	: Mr. RAJESH SHARMA			
AGE/ GENDER	: 63 YRS/MALE		PATIENT ID	: 1538075
COLLECTED BY	:		REG. NO./LAB NO.	: 012407040027
REFERRED BY	: CENTRAL PHOENIX CLUB (A	MBALA CANTT)	REGISTRATION DATE	: 04/Jul/2024 10:28 AM
BARCODE NO.	: 01512502	,	COLLECTION DATE	:04/Jul/2024 12:59PM
CLIENT CODE.	: KOS DIAGNOSTIC LAB		REPORTING DATE	: 04/Jul/2024 11:44AM
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD,	AMBALA CANTT		
Test Name		Value	Unit	Biological Reference interval
GLYCOSYLATED HAEMO			IATOLOGY AEMOGLOBIN (HBA1C) %	4.0 - 6.4
WHOLE BLOOD	IANCE LIQUID CHROMATOGRAPHY)	0.4		4.0 - 0.4
ESTIMATED AVERAGE F by HPLC (HIGH PERFORM INTERPRETATION:	PLASMA GLUCOSE MANCE LIQUID CHROMATOGRAPHY)	136.98	mg/dL	60.00 - 140.00
	AS PER AMERICAN DIAE	BETES ASSOCIATION	(ADA):	
REFERENCE GROUP		GLYCOSYLATED HEMOGLOGIB (HBAIC) i		n %
	etic Adults >= 18 years	<5.7		
	Risk (Prediabetes)		5.7 – 6.4	
Dia	gnosing Diabetes		>= 6.5	
		Coole of The	Age > 19 Years	
	goals for glycemic control	Goals of The Actions Sugg		
Inerapeutic	geale ist gijterine control	Actions Sugg	Age < 19 Years	
Therapeutic				

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 appropriate. 4. High

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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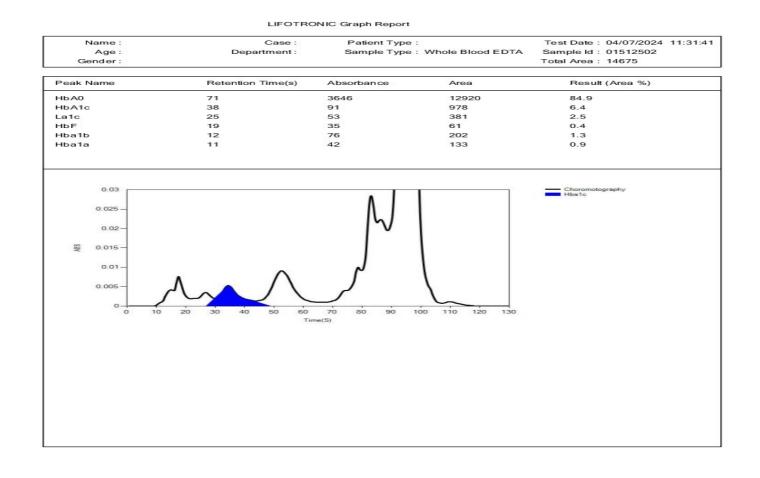


TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT





	Dr. Vinay Chopra MD (Pathology & Microbiology) Chairman & Consultant Pathologis		(Pathology)
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Test Name	Value	Unit	Biological Reference interval







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CLIENT CODE.	: KOS DIAGNOSTIC LAB		REPORTING DATE	:04/Jul/202401:52PM
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD	, AMBALA CANTT	•	
Test Name		Value	Unit	Biological Reference interval
Test Name			STRY/BIOCHEMISTR	Y
	GLUCO	NICAL CHEMIS SE FASTING (F)	STRY/BIOCHEMISTR AND POST PRANDIAL	Y (PP)
GLUCOSE FASTING (GLUCO		STRY/BIOCHEMISTR	Y

IN ACCORDANCE WITH AMERICAN DIABETES ASSOCIATION GUIDELINES:

 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.

*** End Of Report ***





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