



	MD (Pathology &	Dr. Vinay Chopra MD (Pathology & Microbiology) Chairman & Consultant Pathologist		Chopra (Pathology) Pathologist	
NAME	: Mrs. RAJBIR KAUR				
AGE/ GENDER	: 43 YRS/FEMALE	PA	ATIENT ID	: 1662350	
COLLECTED BY	: SURJESH	RI	EG. NO./LAB NO.	: 012411050084	
REFERRED BY	:	RI	EGISTRATION DATE	: 05/Nov/2024 05:20 PM	
BARCODE NO.	:01520180	CO	DLLECTION DATE	: 05/Nov/2024 05:42PM	
CLIENT CODE.	: KOS DIAGNOSTIC LAB	RI	EPORTING DATE	:05/Nov/202406:46PM	
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, A	MBALA CANTT			
Test Name		Value	Unit	Biological Reference interval	
GLYCOSYLATED HAEMOGLOBIN (HbA1c): WHOLE BLOOD by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) ESTIMATED AVERAGE PLASMA GLUCOSE by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) INTERPRETATION:		10.2 ^H 246.04 ^H	% mg/dL	4.0 - 6.4 60.00 - 140.00	
AS PER AMERICAN DIABETES ASSOCIATION (ADA):					
REFERENCE GROUP Non diabetic Adults >= 18 years		GLYC	GLYCOSYLATED HEMOGLOGIB (HBAIC) in %		
At Risk (Prediabetes)			5.7 - 6.4		
Diagnosing Diabetes			>= 6.5		
		Cashaf	Age > 19 Years	7.0	
Therapeutic goals for glycemic control			Therapy: uggested:	< 7.0 >8.0	
			Age < 19 Years		
		Goal of	therapy:	<7.5	
2.Since Hb1c reflects lo concentration of HbAl 3.Target goals of < 7.0	ong term fluctuations in blood glucos lc. Converse is true for a diabetic prev 9 % may be beneficial in patients with	e concentration, a di viously under good co short duration of di	abetic patient who has rec ontrol but now poorly cont abetes, long life expectanc	erapeutic regimen in diabetic patients. ently under good control may still have high rolled. y and no significant cardiovascular disease. In ns, targetting a goal of < 7.0% may not be	

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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.

*** End Of Report ***



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TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT