



	<b>Dr. Vinay Chopra</b> MD (Pathology & Microbiology) Chairman & Consultant Pathologist		Dr. Yugam Chopra MD (Pathology) CEO & Consultant Pathologist		
NAME	: Mrs. JAGMALO				
AGE/ GENDER	: 53 YRS/FEMALE	P	ATIENT ID	: 1700127	
COLLECTED BY	:	R	EG. NO./LAB NO.	: 012412160036	
<b>REFERRED BY</b>	:	R	EGISTRATION DATE	: 16/Dec/2024 12:20 PM	
BARCODE NO.	:01522524	C	OLLECTION DATE	: 16/Dec/2024 12:26PM	
CLIENT CODE.	: KOS DIAGNOSTIC LAB	R	EPORTING DATE	: 16/Dec/2024 02:48PM	
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, AMBALA CANTT				
Test Name		Value	Unit	<b>Biological Reference interval</b>	
GLYCOSYLATED HAEMOGLOBIN (HbA1c): WHOLE BLOOD by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) ESTIMATED AVERAGE PLASMA GLUCOSE by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) INTERPRETATION:		13.1 <sup>H</sup> 329.27 <sup>H</sup>	% mg/dL	4.0 - 6.4 60.00 - 140.00	
	AS PER AMERICAN	DIABETES ASSOCIAT			
REFERENCE GROUP			COSYLATED HEMOGLOGIB	(HBAIC) in %	
Non diabetic Adults >= 18 years			<5.7		
At Risk (Prediabetes)		_	5.7 - 6.4		
	Diagnosing Diabetes		>= 6.5 Age > 19 Years		
Therapeutic goals for glycemic control			Therapy: Suggested:	< 7.0 >8.0	
		Goal of	Age < 19 Years therapy:	<7.5	
2.Since Hb1c reflects in concentration of HbA 3.Target goals of < 7.0	ong term fluctuations in blood gluco Ic. Converse is true for a diabetic pre ) % may be beneficial in patients wit	se concentration, a d viously under good c h short duration of di	liabetic patient who has re ontrol but now poorly con iabetes, long life expectan	nerapeutic regimen in diabetic patients. cently under good control may still have high trolled. cy and no significant cardiovascular disease. In ons, 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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Page 1 of