



	Dr. Vinay Cho MD (Pathology & M Chairman & Consu	licrobiology)	Dr. Yugan MD CEO & Consultan	(Pathology)	
NAME	: Mr. SUKET DHAWAN				
AGE/ GENDER	: 61 YRS/MALE	I	PATIENT ID	: 1822379	
COLLECTED BY	: SURJESH	T	REG. NO./LAB NO.	: 012504080057	
REFERRED BY	: CENTRAL PHOENIX CLUB (AM		REGISTRATION DATE	: 08/Apr/2025 12:08 PM	
BARCODE NO.	: 01528610	(	COLLECTION DATE	:08/Apr/2025 12:17PM	
CLIENT CODE.	: KOS DIAGNOSTIC LAB	F	REPORTING DATE	: 08/Apr/2025 02:05PM	
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, AM	/IBALA CANTT			
Test Name		Value	Unit	Biological Reference	interva
GLYCOSYLATED HAEMOGLOBIN (HbA1c): WHOLE BLOOD by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) ESTIMATED AVERAGE PLASMA GLUCOSE by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY)		5.6 114.02	% mg/dL	4.0 - 6.4 60.00 - 140.00	
INTERPRETATION:					
D	AS PER AMERICAN D EFERENCE GROUP		TION (ADA): COSYLATED HEMOGLOGIE	(HPAIC) in %	
	betic Adults >= 18 years	GLY	<5.7		
	At Risk (Prediabetes)		5.7 – 6.4		
Diagnosing Diabetes			>= 6.5		
	5 5		Age > 19 Years		
			of Therapy:	< 7.0	
Therapeutic	c goals for glycemic control	Actions	Suggested:	>8.0	
merupeutit			Age < 19 Years		
merupeuti			f therapy:	<7.5	

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 appropiate.

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