



	Dr. Vinay Cho MD (Pathology & N Chairman & Consu	icrobiology) MD (Pathology)				
NAME	: Mr. TRILOK CHAND					
AGE/ GENDER	: 59 YRS/MALE		PATIENT ID	: 1630104		
COLLECTED BY	:		REG. NO./LAB NO.	: 01240930008	1	
REFERRED BY			REGISTRATION DAT	<b>FE</b> : 30/Sep/2024 04	:06 PM	
BARCODE NO.	: 01518054		COLLECTION DATE	: 30/Sep/2024 04		
CLIENT CODE.	: KOS DIAGNOSTIC LAB		REPORTING DATE	: 30/Sep/2024 04		
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, AI			. 30/ 50/ 2024 04		
Test Name		Value	Unit	Biologic	al Reference interval	
	GLYC		ATOLOGY EMOGLOBIN (HBA	.1C)		
GLYCOSYLATED HAEMOGLOBIN (HbA1c): WHOLE BLOOD		5.9	%	4.0 - 6.4	4.0 - 6.4	
by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) ESTIMATED AVERAGE PLASMA GLUCOSE by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) INTERPRETATION:		122.63	mg/d	mg/dL 60.00 - 140.00		
	AS PER AMERICAN D					
	REFERENCE GROUP	GL	GLYCOSYLATED HEMOGLOGIB (HBAIC) in %			
	abetic Adults >= 18 years	_	<5.7			
At Risk (Prediabetes)		5.7 - 6.4				
Diagnosing Diabetes		>= 6.5				
		Age > 19 Years Goals of Therapy:		ears < 7.0		
Therapeut	ic goals for glycemic control	Actions Suggested:		>8.0		
merapeut	ie gouis for grycenne control	Actions suggested: >8.0 Age < 19 Years				
		Goal of therapy:		501.3		

**KOS Diagnostic Lab** 

(A Unit of KOS Healthcare)

1.Glycosylated hemoglobin (HbA1c) test is three monthly monitoring done to assess compliace with therapeutic regimen in diabetic patients. 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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