



	bra Dr. Yugam Chop licrobiology) MD (Patholo tant Pathologist CEO & Consultant Patholog			(Pathology)		
NAME	: Mr. PREETI KUMAR					
AGE/ GENDER	DER : 34 YRS/MALE		PATIENT ID		: 1616474	
OLLECTED BY : EFERRED BY :			REG. NO./LAB NO. REGISTRATION DATE		: <b>012409170051</b> : 17/Sep/2024 05:38 PM : 17/Sep/2024 05:41PM	
BARCODE NO.	:01517152	517152 COLLECTION DATE				
CLIENT CODE.	: KOS DIAGNOSTIC LAB			-		
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, A	MBALA CANTI				-
Test Name		Value		Unit	Biological Re	ference interval
GLYCOSYLATED HAEMOGLOBIN (HbA1c): WHOLE BLOOD by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) ESTIMATED AVERAGE PLASMA GLUCOSE by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY) INTERPRETATION:		6.4 136.98	AEMOGLOBIN (HBA1C) % mg/dL		4.0 - 6.4 60.00 - 140.00	)
<u>MIERI KETAHON.</u>	AS PER AMERICAN D					
REFERENCE GROUP			GLYCOSYLATED HEMOGLOGIB (HBAIC) in %			
Non diabetic Adults >= 18 years		<5.7				
At Risk (Prediabetes)			5.7 - 6.4			
Diagnosing Diabetes		>= 6.5				
Therapeutic goals for glycemic control		Age > 19 Years   Goals of Therapy:   Actions Suggested:		< 7.0 >8.0		
		Age < 19 Years Goal of therapy:			.7.5	
		Goa	i oi therapy:		<7.5	

**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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