



	Dr. Vinay Chopra MD (Pathology & Micr Chairman & Consultar	Microbiology) MD (Pathology)		
IAME	: Mr. AMANPREET SINGH			
GE/ GENDER	: 56 YRS/MALE	PA	ATIENT ID	: 1797284
OLLECTED BY	:	RI	EG. NO./LAB NO.	: 012503190001
REFERRED BY	:	RI	EGISTRATION DATE	: 19/Mar/2025 06:47 AM
BARCODE NO.	: 01527362		DLLECTION DATE	: 19/Mar/2025 09:15AM
LIENT CODE.	: KOS DIAGNOSTIC LAB		EPORTING DATE	: 19/Mar/2025 10:58AM
LIENT ADDRESS	: 6349/1, NICHOLSON ROAD, AMB.	ALA CANTT		
Fest Name		Value	Unit	Biological Reference interval
		НАЕМАТ	TOLOGY	
	COMP	LETE BLOC	DD COUNT (CBC)	
RED BLOOD CELLS	S (RBCS) COUNT AND INDICES			
IAEMOGLOBIN (H	B)	11.6 ^L	gm/dL	12.0 - 17.0
by CALORIMETRIC RED BLOOD CELL (RBC) COUNT	3.92	Millions/c	mm 3.50 - 5.00
by HYDRO DYNAMIC F	OCUSING, ELECTRICAL IMPEDENCE		0/	10.0 51.0
PACKED CELL VOLUME (PCV) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER MEAN CORPUSCULAR VOLUME (MCV) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER		35.1 ^L	%	40.0 - 54.0
		89.6	fL	80.0 - 100.0
AEAN CORPUSCUL	AR HAEMOGLOBIN (MCH)	29.5	pg	27.0 - 34.0
	UTOMATED HEMATOLOGY ANALYZER AR HEMOGLOBIN CONC. (MCHC)	33	g/dL	32.0 - 36.0
by CALCULATED BY A	UTOMATED HEMATOLOGY ANALYZER		ç	
	UTION WIDTH (RDW-CV) UTOMATED HEMATOLOGY ANALYZER	17.3 ^H	%	11.00 - 16.00
	UTION WIDTH (RDW-SD) UTOMATED HEMATOLOGY ANALYZER	58.1 ^H	fL	35.0 - 56.0
MENTZERS INDEX	OTOMATED TIEMATOLOGT ANALIZEN	22.86	RATIO	BETA THALASSEMIA TRAIT: <
by CALCULATED				13.0 IRON DEFICIENCY ANEMIA:
				>13.0
GREEN & KING IND	DEX	39.42	RATIO	BETA THALASSEMIA TRAIT:<=
by CALCULATED				65.0 IRON DEFICIENCY ANEMIA: >
				65.0
VHITE BLOOD CE				1000 11000
DTAL LEUCOCYTE	E COUNT (TLC) (by sf cube & microscopy	3550 ^L	/cmm	4000 - 11000
	BLOOD CELLS (nRBCS)	NIL		0.00 - 20.00
	TI NEMA I ULUGY ANALYZEK			
by AUTOMATED 6 PAP	BLOOD CELLS (nRBCS) %	NIL	%	< 10 %





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TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.



:

:

NAME

AGE/ GENDER

COLLECTED BY

REFERRED BY

BARCODE NO.

CLIENT CODE.

CLIENT ADDRESS



Dr. Yugam Chopra

MD (Pathology)

:1797284

:012503190001

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Dr. Vinay Chopra MD (Pathology & Microbiology) Chairman & Consultant Pathologist CEO & Consultant Pathologist : Mr. AMANPREET SINGH **PATIENT ID** : 56 YRS/MALE REG. NO./LAB NO. **REGISTRATION DATE** :01527362 **COLLECTION DATE** : KOS DIAGNOSTIC LAB **REPORTING DATE** : 6349/1, NICHOLSON ROAD, AMBALA CANTT

Test Name	Value	Unit	Biological Reference interval
DIFFERENTIAL LEUCOCYTE COUNT (DLC)			
NEUTROPHILS by flow cytometry by SF cube & microscopy	47 ^L	%	50 - 70
LYMPHOCYTES by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	39	%	20 - 40
EOSINOPHILS by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	4	%	1 - 6
MONOCYTES by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	10	%	2 - 12
BASOPHILS by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	0	%	0 - 1
ABSOLUTE LEUKOCYTES (WBC) COUNT			
ABSOLUTE NEUTROPHIL COUNT by flow cytometry by sf cube & microscopy	1669 ^L	/cmm	2000 - 7500
ABSOLUTE LYMPHOCYTE COUNT by flow cytometry by SF cube & microscopy	1384	/cmm	800 - 4900
ABSOLUTE EOSINOPHIL COUNT by flow cytometry by sf cube & microscopy	142	/cmm	40 - 440
ABSOLUTE MONOCYTE COUNT by flow cytometry by SF cube & microscopy	355	/cmm	80 - 880
ABSOLUTE BASOPHIL COUNT by flow cytometry by SF cube & microscopy	0	/cmm	0 - 110
ABSOLUTE IMMATURE GRANULOCYTE COUNT by flow cytometry by sf cube & microscopy	0	/cmm	0.0 - 999.0
PLATELETS AND OTHER PLATELET PREDICTIVE	MARKERS.		
PLATELET COUNT (PLT) by hydro dynamic focusing, electrical impedence	52000 ^L	/cmm	150000 - 450000
PLATELETCRIT (PCT) by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	0.07 ^L	%	0.10 - 0.36
MEAN PLATELET VOLUME (MPV) by hydro dynamic focusing, electrical impedence	14 ^H	fL	6.50 - 12.0
PLATELET LARGE CELL COUNT (P-LCC) by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	28000 ^L	/cmm	30000 - 90000
PLATELET LARGE CELL RATIO (P-LCR) by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	54.5 ^H	%	11.0 - 45.0
PLATELET DISTRIBUTION WIDTH (PDW) by hydro dynamic focusing, electrical impedence	16.8	%	15.0 - 17.0



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Test Name	Value	Unit	Biological Reference interval

NOTE: TEST CONDUCTED ON EDTA WHOLE BLOOD

RECHECKED.

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Test Name		Value	Unit	Biological Reference interva	
WHOLE BLOOD	EMOGLOBIN (HbA1c):	6	MOGLOBIN (HBA1) %	4.0 - 6.4	
ESTIMATED AVERAGE PLASMA GLUCOSE by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY)		125.5	mg/dL	60.00 - 140.00	
INTERPRETATION:					
	AS PER AMERICAN D	ABETES ASSOCIATI	ION (ADA):		
1	REFERENCE GROUP	GLYCOSYLATED HEMOGLOGIB (HBAIC) in %			
	abetic Adults >= 18 years	1	<5.7		
At Risk (Prediabetes)		5.7 - 6.4			
D	iagnosing Diabetes		>= 6.5		
			Age > 19 Years		
Therapeutic goals for glycemic control			Therapy:	< 7.0	
		Actions Suggested:		>8.0	
			Age < 19 Years therapy:	<7.5	
1		Gudi Ul	uiciapy.	<1.J	

KOS Diagnostic Lab

(A Unit of KOS Healthcare)

COMMENTS

TEST PERFORMED AT KOS DIAGNOSTIC LAB. AMBALA CANTT

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 appropriate.

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.



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		& Microbiology) onsultant Pathologist	MD CEO & Consultant	(Pathology) : Pathologist
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Test Name		Value	Unit	Biological Reference interval
	CLINI	CAL CHEMISTRY	/BIOCHEMIST	'RY
			FINC (F)	
		GLUCOSE FAS	I ING (F)	

IN ACCORDANCE WITH AMERICAN DIABETES ASSOCIATION GUIDELINES:

A fasting plasma glucose level below 100 mg/dl is considered normal.
 A fasting plasma glucose level between 100 - 125 mg/dl is considered as glucose intolerant or prediabetic. A fasting and post-prandial blood test (after consumption of 75 gms of glucose) is recommended for all such patients.

test (after consumption of 75 gms of glucose) is recommended for all such patients. 3. A fasting plasma glucose level of above 125 mg/dl is highly suggestive of diabetic state. A repeat post-prandial is strongly recommended for all such patients. A fasting plasma glucose level in excess of 125 mg/dl on both occasions is confirmatory for diabetic state.



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Test Name		Value	Unit	Biological Reference interval
	G	LUCOSE POS	Г PRANDIAL (PP)	
	ANDIAL (PP): PLASMA e - peroxidase (god-pod)	252.11 ^H	mg/dL	NORMAL: < 140.00 PREDIABETIC: 140.0 - 200.0 DIABETIC: > 0R = 200.0

KOS Diagnostic Lab (A Unit of KOS Healthcare)

TEST PERFORMED AT KOS DIAGNOSTIC LAB. AMBALA CANTT

INTERPRETATION IN ACCORDANCE WITH AMERICAN DIABETES ASSOCIATION GUIDELINES: 1. A post-prandial plasma glucose level below 140 mg/dl is considered normal. 2. A post-prandial glucose level between 140 - 200 mg/dl is considered as glucose intolerant or prediabetic. A fasting and post-prandial blood test (after consumption of 75 gms of glucose) is recommended for all such patients. 3. A post-prandial plasma glucose level of above 200 mg/dl is highly suggestive of diabetic state. A repeat post-prandial is strongly recommended for all such patients. A fasting plasma glucose level in excess of 125 mg/dl on both occasions is confirmatory for diabetic state.

*** End Of Report ***





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