



	Dr. Vinay Chopra MD (Pathology & Micr Chairman & Consultar	obiology)		(Pathology)
NAME	: Mr. PANKAJ			
AGE/ GENDER	: 43 YRS/MALE		PATIENT ID	: 1684569
COLLECTED BY	:		REG. NO./LAB NO.	: 042411280003
REFERRED BY	:		REGISTRATION DATE	: 28/Nov/2024 10:13 AM
BARCODE NO.	: A1259983		COLLECTION DATE	: 28/Nov/2024 03:39PM
CLIENT CODE.	: KOS DIAGNOSTIC SHAHBAD		REPORTING DATE	: 28/Nov/2024 03:48PM
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, AMB/	ALA CANTT		
Test Name		Value	Unit	Biological Reference interval
		HAEM	ATOLOGY	
	COMP	PLETE BLO	DOD COUNT (CBC)	
RED BLOOD CELLS	(RBCS) COUNT AND INDICES			
HAEMOGLOBIN (H	B)	16.6	gm/dL	12.0 - 17.0
by CALORIMETRIC RED BLOOD CELL (RBC) COUNT	5.23 ^H	Millions	/cmm 3.50 - 5.00
by HYDRO DYNAMIC F	OCUSING, ELECTRICAL IMPEDENCE			
PACKED CELL VOLU	JME (PCV) UTOMATED HEMATOLOGY ANALYZER	51.2	%	40.0 - 54.0
MEAN CORPUSCUL	AR VOLUME (MCV)	97.8	fL	80.0 - 100.0
	UTOMATED HEMATOLOGY ANALYZER AR HAEMOGLOBIN (MCH)	31.6	pg	27.0 - 34.0
by CALCULATED BY A	UTOMATED HEMATOLOGY ANALYZER			
	AR HEMOGLOBIN CONC. (MCHC) UTOMATED HEMATOLOGY ANALYZER	32.4	g/dL	32.0 - 36.0
	UTION WIDTH (RDW-CV) UTOMATED HEMATOLOGY ANALYZER	14.6	%	11.00 - 16.00
•	UTION WIDTH (RDW-SD)	53.2	fL	35.0 - 56.0
by CALCULATED BY A MENTZERS INDEX	UTOMATED HEMATOLOGY ANALYZER	10 7	RATIO	
by CALCULATED		18.7	KATIO	BETA THALASSEMIA TRAIT: < 13.0
				IRON DEFICIENCY ANEMIA:
GREEN & KING IND	ЪЕХ	27.18	RATIO	>13.0 BETA THALASSEMIA TRAIT:<=
by CALCULATED	<u></u>	£7.10	101110	65.0
				IRON DEFICIENCY ANEMIA: > 65.0
WHITE BLOOD CE	LLS (WBCS)			0010
TOTAL LEUCOCYTE		8380	/cmm	4000 - 11000
by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY		NIL		0.00 - 20.00
	LOOD CELLS (nRBCS)			0.00 20.00
NUCLEATED RED B	LOOD CELLS (NRBCS) <i>it hematology analyzer</i> LOOD CELLS (NRBCS) %	NIL	%	< 10 %

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TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.





Dr. Vinay Chopra

MD (Pathology & Microbiology) Chairman & Consultant Pathologist



Dr. Yugam Chopra MD (Pathology) CEO & Consultant Pathologist

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Test Name	Value	Unit	Biological Reference interval
DIFFERENTIAL LEUCOCYTE COUNT (DLC)			
NEUTROPHILS by flow cytometry by SF cube & microscopy	54	%	50 - 70
LYMPHOCYTES by flow cytometry by SF cube & microscopy	38	%	20 - 40
EOSINOPHILS by flow cytometry by SF cube & microscopy	2	%	1 - 6
MONOCYTES by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	6	%	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	4525	/cmm	2000 - 7500
ABSOLUTE LYMPHOCYTE COUNT by flow cytometry by sf cube & microscopy	3184	/cmm	800 - 4900
ABSOLUTE EOSINOPHIL COUNT by flow cytometry by sf cube & microscopy	168	/cmm	40 - 440
ABSOLUTE MONOCYTE COUNT by flow cytometry by sf cube & microscopy	503	/cmm	80 - 880
PLATELETS AND OTHER PLATELET PREDICTIVE	MARKERS.		
PLATELET COUNT (PLT) by hydro dynamic focusing, electrical impedence	318000	/cmm	150000 - 450000
PLATELETCRIT (PCT) by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	0.3	%	0.10 - 0.36
MEAN PLATELET VOLUME (MPV) by hydro dynamic focusing, electrical impedence	10	fL	6.50 - 12.0
PLATELET LARGE CELL COUNT (P-LCC) by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	73000	/cmm	30000 - 90000
PLATELET LARGE CELL RATIO (P-LCR) by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	22.9	%	11.0 - 45.0
PLATELET DISTRIBUTION WIDTH (PDW) by hydro dynamic focusing, electrical impedence NOTE: TEST CONDUCTED ON EDTA WHOLE BLOOD	16.2	%	15.0 - 17.0



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CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, AM	IBALA CANTT		
Test Name		Value	Unit	Biological Reference interva
	GLYCOS	YLATED HA	AEMOGLOBIN (HBA10	C)
GLYCOSYLATED HAEMOGLOBIN (HbA1c): WHOLE BLOOD by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY)		5.6	%	4.0 - 6.4
ESTIMATED AVERAGE PLASMA GLUCOSE by HPLC (HIGH PERFORMANCE LIQUID CHROMATOGRAPHY)		114.02	mg/dL	60.00 - 140.00
by HPLC (HIGH PERFO	RMANCE LIQUID CHROMATOGRAPHY)			
by HPLC (HIGH PERFO		ABETES ASSOCI	ATION (ADA):	
by HPLC (HIGH PERFO INTERPRETATION:	AS PER AMERICAN DI		ATION (ADA): LYCOSYLATED HEMOGLOGIB	(HBAIC) in %
by HPLC (HIGH PERFO INTERPRETATION:	AS PER AMERICAN DI			(HBAIC) in %
by HPLC (HIGH PERFO INTERPRETATION: Non di	AS PER AMERICAN DI REFERENCE GROUP		LYCOSYLATED HEMOGLOGIB	(HBAIC) in %
by HPLC (HIGH PERFO INTERPRETATION: Non di A	AS PER AMERICAN DI REFERENCE GROUP abetic Adults >= 18 years		LYCOSYLATED HEMOGLOGIB <5.7	(HBAIC) in %
by HPLC (HIGH PERFO INTERPRETATION: Non di A	AS PER AMERICAN DI REFERENCE GROUP abetic Adults >= 18 years t Risk (Prediabetes)	G	<5.7 5.7 - 6.4 >= 6.5 Age > 19 Years	
by HPLC (HIGH PERFO INTERPRETATION: Non di A D	AS PER AMERICAN DI REFERENCE GROUP abetic Adults >= 18 years t Risk (Prediabetes) iagnosing Diabetes	G	<5.7 5.7 - 6.4 >= 6.5 Age > 19 Years s of Therapy:	< 7.0
by HPLC (HIGH PERFO INTERPRETATION: Non di A D	AS PER AMERICAN DI REFERENCE GROUP abetic Adults >= 18 years t Risk (Prediabetes)	G	Action State State <t< td=""><td></td></t<>	
by HPLC (HIGH PERFO INTERPRETATION: Non di A D	AS PER AMERICAN DI REFERENCE GROUP abetic Adults >= 18 years t Risk (Prediabetes) iagnosing Diabetes	Goals Goals Actior	<5.7 5.7 - 6.4 >= 6.5 Age > 19 Years s of Therapy:	< 7.0

COMMENTS:

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.

*** End Of Report ***





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