



	MD (Pathology & Micr	Dr. Vinay Chopra MD (Pathology & Microbiology) Chairman & Consultant Pathologist		Dr. Yugam Chopra MD (Pathology) CEO & Consultant Pathologist	
IAME	: Mr. RAMAN JAIN				
GE/ GENDER	: 53 YRS/MALE	PA	TIENT ID	: 1661796	
COLLECTED BY	: SURJESH	RE	G. NO./LAB NO.	: 012411050069	
REFERRED BY	:		GISTRATION DATE	: 05/Nov/2024 01:49 PM	
SARCODE NO.	: 01520165		LLECTION DATE	: 05/Nov/2024 01:57PM	
CLIENT CODE. CLIENT ADDRESS	: KOS DIAGNOSTIC LAB : 6349/1, NICHOLSON ROAD, AMB/		PORTING DATE	: 05/Nov/2024 02:17PM	
fest Name		Value	Unit	Biological Reference interval	
		НАЕМАТ	OLOGY		
	COMP	LETE BLOO	D COUNT (CBC)		
ED BLOOD CELLS	S (RBCS) COUNT AND INDICES				
AEMOGLOBIN (H	B)	14.8	gm/dL	12.0 - 17.0	
ED BLOOD CELL (RBC) COUNT	5.45 ^H	Millions/c	mm 3.50 - 5.00	
PACKED CELL VOLUME (PCV)		46.1	%	40.0 - 54.0	
by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER MEAN CORPUSCULAR VOLUME (MCV) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER		84.6	fL	80.0 - 100.0	
	AR HAEMOGLOBIN (MCH) UTOMATED HEMATOLOGY ANALYZER	27.2	pg	27.0 - 34.0	
IEAN CORPUSCUL	AR HEMOGLOBIN CONC. (MCHC) UTOMATED HEMATOLOGY ANALYZER	32.2	g/dL	32.0 - 36.0	
RED CELL DISTRIBUTION WIDTH (RDW-CV)		14.1	%	11.00 - 16.00	
ED CELL DISTRIB	UTION WIDTH (RDW-SD) UTOMATED HEMATOLOGY ANALYZER	44.4	fL	35.0 - 56.0	
MENTZERS INDEX by CALCULATED		15.52	RATIO	BETA THALASSEMIA TRAIT: < 13.0 IRON DEFICIENCY ANEMIA: >13.0	
GREEN & KING INE by CALCULATED		21.92	RATIO	BETA THALASSEMIA TRAIT:<= 65.0 IRON DEFICIENCY ANEMIA: > 65.0	
VHITE BLOOD CE					
FOTAL LEUCOCYTE COUNT (TLC) by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY		8530	/cmm	4000 - 11000	
UCLEATED RED B	BLOOD CELLS (nRBCS) RT HEMATOLOGY ANALYZER	NIL		0.00 - 20.00	
	LOOD CELLS (nRBCS) %	NIL	%	< 10 %	





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TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.





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Dr. Vinay Chopra



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Test Name	Value	Unit	Biological Reference interval
DIFFERENTIAL LEUCOCYTE COUNT (DLC)			
NEUTROPHILS by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	63	%	50 - 70
LYMPHOCYTES by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	29	%	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	5374	/cmm	2000 - 7500
ABSOLUTE LYMPHOCYTE COUNT by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	2474	/cmm	800 - 4900
ABSOLUTE EOSINOPHIL COUNT by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	171	/cmm	40 - 440
ABSOLUTE MONOCYTE COUNT by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	512	/cmm	80 - 880
PLATELETS AND OTHER PLATELET PREDICTIVE	MARKERS.		
PLATELET COUNT (PLT) by hydro dynamic focusing, electrical impedence	369000	/cmm	150000 - 450000
PLATELETCRIT (PCT) by hydro dynamic focusing, electrical impedence	0.44 ^H	%	0.10 - 0.36
MEAN PLATELET VOLUME (MPV) by hydro dynamic focusing, electrical impedence	12	fL	6.50 - 12.0
PLATELET LARGE CELL COUNT (P-LCC) by hydro dynamic focusing, electrical impedence	146000 ^H	/cmm	30000 - 90000
PLATELET LARGE CELL RATIO (P-LCR) by hydro dynamic focusing, electrical impedence	39.5	%	11.0 - 45.0
PLATELET DISTRIBUTION WIDTH (PDW) by hydro dynamic focusing, electrical impedence NOTE: TEST CONDUCTED ON EDTA WHOLE BLOOD	16.1	%	15.0 - 17.0



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CLIENT ADDRESS	: 6349/1, NICHOLSON F	COAD, AMBALA CANT	Т	
Test Name		Value	Unit	Biological Reference interval
		IMMUNOPATH	HOLOGY/SEROLOGY	
	DENGUE F	EVER COMBO SCREI	ENING - (NS1 ANTIGEN, Ig	G AND IgM)
DENGUE NS1 ANTIGEN - SCREENING by ICT (IMMUNOCHROMATOGRAPHY)		NEGATIVE (-ve)		NEGATIVE (-ve)
DENGUE ANTIBODY Ig		NEGATIVE (-ve)		NEGATIVE (-ve)
DENGUE ANTIBODY Ig by ICT (IMMUNOCHROMAT		NEGATIVE (-ve)		NEGATIVE (-ve)

INTERPRETATION:-

7. This is a solid phase immunochromatographic ELISA test for the qualitative detection of the specific IgG and IgM antibodies against the Dengue virus.

KOS Diagnostic Lab (A Unit of KOS Healthcare)

2. The IgM antibodies take a minimum of 5-10 days in primary infection and 4-5 days in secondary infections to test positive and hence are suitable for the diagnosis of dengue fever only when the fever is approximately one week old.

3. The IgG antibodies develop at least two weeks after exposure to primary infection and subsequently remain positive for the rest of the life. A positive result is incapable of differentiating a current infection from a past infection.

4. The Dengue NS-1 antigen test is most suited for early diagnosis (within the first week of exposure).





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Test Name		Value	Unit	Biological Reference interval
		WIDAL SLIDE A	GGLUTINATION TEST	
SALMONELLA TYPHI O by SLIDE AGGLUTINATION		1:20	TITRE	1:80
SALMONELLA TYPHI H by slide agglutination		1:20	TITRE	1:160
SALMONELLA PARATYPHI AH by SLIDE AGGLUTINATION		NIL	TITRE	1:160
SALMONELLA PARATYPHI BH		NIL	TITRE	1:160

SALMONELLA PARATYPHI BH by SLIDE AGGLUTINATION

INTERPRETATION:

1. Titres of 1:80 or more for "O" agglutinin is considered significant.

2. Titres of 1:160 or more for "H" agglutinin is considered significant.

LIMITATIONS:

1.Agglutinins usually appear by 5th to 6th day of illness of enteric fever, hence a negative result in early stage is inconclusive. The titre then rises till 3rd or 4th week, after which it declines gradually.

2.Lower titres may be found in normal individuals.

3.A single positive result has less significance than the rising agglutination titre, since demonstration of rising titre four or more in 1st and 3rd week is considered as a definite evidence of infection.

4.A simultaneous rise in H agglutinins is suggestive of paratyphoid infection.

NOTE:

1. Individuals with prior infection or immunization with TAB vaccine may develop an ANAMNESTIC RESPONSE (False-Positive) during an unrelated fever i.e High titres of antibodies to various antigens. This may be differentiated by repitition of the test after a week.

2. The anamnestic response shows only a transient rise, while in enteric fever rise is sustained.

3.H agglutinins tend to persist for many months after vaccination but O agglutinins tend to disappear sooner i.e within 6 months. Therefore rise in Oagglutinins indicate recent infection.

*** End Of Report ***





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