

TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.



	Dr. Vinay Chopra MD (Pathology & Micr Chairman & Consultan	obiology)	M	m Chopra D (Pathology) nt Pathologist	
NAME : N	Mrs. YOGITA				
AGE/ GENDER : 4	19 YRS/FEMALE		PATIENT ID	: 1774466	
COLLECTED BY :			REG. NO./LAB NO.	:0125030	10031
REFERRED BY :			REGISTRATION DATE	:01/Mar/20	025 12:29 PM
	01526290		COLLECTION DATE	:01/Mar/20	025 12:30PM
	KOS DIAGNOSTIC LAB		REPORTING DATE	:01/Mar/20	025 12:43PM
CLIENT ADDRESS : 6	3349/1, NICHOLSON ROAD, AMBA	ALA CANT'I			
Test Name		Value	Unit	Bi	ological Reference interval
		HAEM	ATOLOGY		
	COMP		OOD COUNT (CBC)		
RED BLOOD CELLS (R	BCS) COUNT AND INDICES				
HAEMOGLOBIN (HB) by CALORIMETRIC		11.4 ^L	gm/dL	12	2.0 - 16.0
RED BLOOD CELL (RBC	C) COUNT ISING, ELECTRICAL IMPEDENCE	4.65	Millions	s/cmm 3.	50 - 5.00
PACKED CELL VOLUME		36.2 ^L	%	37	7.0 - 50.0
MEAN CORPUSCULAR		77.8 ^L	fL	80	0.0 - 100.0
	HAEMOGLOBIN (MCH) MATED HEMATOLOGY ANALYZER	24.5 ^L	pg	27	7.0 - 34.0
MEAN CORPUSCULAR	HEMOGLOBIN CONC. (MCHC) MATED HEMATOLOGY ANALYZER	31.5 ^L	g/dL	32	2.0 - 36.0
•	MATED HEMATOLOGY ANALYZER	16.1 ^H	%	11	1.00 - 16.00
RED CELL DISTRIBUTI by calculated by auto	ON WIDTH (RDW-SD) mated hematology analyzer	46.9	fL	35	5.0 - 56.0
MENTZERS INDEX by CALCULATED		16.73	RATIO	13 IR	ETA THALASSEMIA TRAIT: < 3.0 20N DEFICIENCY ANEMIA: 13.0
GREEN & KING INDEX by CALCULATED		26.92	RATIO	65 IR	ETA THALASSEMIA TRAIT:<= 5.0 20N DEFICIENCY ANEMIA: > 5.0
WHITE BLOOD CELLS	(WBCS)				
TOTAL LEUCOCYTE CO by FLOW CYTOMETRY BY		9190	/cmm	40	000 - 11000
NUCLEATED RED BLO by AUTOMATED 6 PART H	OD CELLS (nRBCS)	NIL		0.0	00 - 20.00
NUCLEATED RED BLO		NIL	%	<	10 %





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

NAME	: Mrs. YOGITA		
AGE/ GENDER	: 49 YRS/FEMALE	PATIENT ID	: 1774466
COLLECTED BY	:	REG. NO./LAB NO.	: 012503010031
REFERRED BY	:	REGISTRATION DATE	: 01/Mar/2025 12:29 PM
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Dr. Vinay Chopra

Test Name	Value	Unit	Biological Reference interval
DIFFERENTIAL LEUCOCYTE COUNT (DLC)			
NEUTROPHILS by flow cytometry by SF cube & microscopy	56	%	50 - 70
LYMPHOCYTES by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	37	%	20 - 40
EOSINOPHILS by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	2	%	1 - 6
MONOCYTES by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	5	%	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	5146	/cmm	2000 - 7500
ABSOLUTE LYMPHOCYTE COUNT by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	3400	/cmm	800 - 4900
ABSOLUTE EOSINOPHIL COUNT by flow cytometry by SF cube & microscopy	184	/cmm	40 - 440
ABSOLUTE MONOCYTE COUNT by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	460	/cmm	80 - 880
ABSOLUTE BASOPHIL COUNT by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	0	/cmm	0 - 110
PLATELETS AND OTHER PLATELET PREDICTIVE	MARKERS.		
PLATELET COUNT (PLT) by hydro dynamic focusing, electrical impedence	303000	/cmm	150000 - 450000
PLATELETCRIT (PCT) by hydro dynamic focusing, electrical impedence	0.32	%	0.10 - 0.36
MEAN PLATELET VOLUME (MPV) by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	11	fL	6.50 - 12.0
PLATELET LARGE CELL COUNT (P-LCC) by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	92000 ^H	/cmm	30000 - 90000
PLATELET LARGE CELL RATIO (P-LCR) by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	30.3	%	11.0 - 45.0
PLATELET DISTRIBUTION WIDTH (PDW) by hydro dynamic focusing, electrical impedence NOTE: TEST CONDUCTED ON EDTA WHOLE BLOOD	16	%	15.0 - 17.0





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Test Name	V	alue Unit	Biological Reference interval



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TITRE

1:160

		hopra & Microbiology) onsultant Pathologist	Dr. Yugam MD (I CEO & Consultant F	Pathology)
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Test Name		Value	Unit	Biological Reference interva
	IM	MUNOPATHOLO	GY/SEROLOGY	
	W	IDAL SLIDE AGGLU	TINATION TEST	
SALMONELLA TYP by slide agglutina		NIL	TITRE	1:80
SALMONELLA TYP by SLIDE AGGLUTINA		NIL	TITRE	1:160
SALMONELLA PAR		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.

NIL

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:

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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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