

A PIONEER DIAGNOSTIC CENTRE

■ 0171-2532620, 8222896961 ■ pkrjainhealthcare@gmail.com

NAME : Mrs. REKHA RANI

AGE/ GENDER : 38 YRS/FEMALE **PATIENT ID** : 1669380

COLLECTED BY REG. NO./LAB NO. : 122411120013

REFERRED BY **REGISTRATION DATE** : 12/Nov/2024 10:44 AM BARCODE NO. : 12505616 **COLLECTION DATE** : 12/Nov/2024 11:22AM CLIENT CODE. : P.K.R JAIN HEALTHCARE INSTITUTE REPORTING DATE : 12/Nov/2024 12:23PM

CLIENT ADDRESS : NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYANA

Value Unit **Biological Reference interval Test Name**

HAEMATOLOGY COMPLETE BLOOD COUNT (CBC)

RED BLOOD CELLS (RBCS) COUNT AND INDICES

HAEMOGLOBIN (HB) by CALORIMETRIC	11.9 ^L	gm/dL	12.0 - 16.0
RED BLOOD CELL (RBC) COUNT by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	4.12	Millions/cmm	3.50 - 5.00
PACKED CELL VOLUME (PCV) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER	34.3 ^L	%	37.0 - 50.0
MEAN CORPUSCULAR VOLUME (MCV) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER	83.1	fL	80.0 - 100.0
MEAN CORPUSCULAR HAEMOGLOBIN (MCH) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER	28.9	pg	27.0 - 34.0
MEAN CORPUSCULAR HEMOGLOBIN CONC. (MCHC) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER	34.7	g/dL	32.0 - 36.0
RED CELL DISTRIBUTION WIDTH (RDW-CV) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER	15.6	%	11.00 - 16.00
RED CELL DISTRIBUTION WIDTH (RDW-SD) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER	49.6	fL	35.0 - 56.0
MENTZERS INDEX by CALCULATED	20.17	RATIO	BETA THALASSEMIA TRAIT: < 13.0 IRON DEFICIENCY ANEMIA: >13.0
GREEN & KING INDEX by CALCULATED	31.48	RATIO	BETA THALASSEMIA TRAIT:<= 65.0 IRON DEFICIENCY ANEMIA: > 65.0
WHITE BLOOD CELLS (WBCS)			
TOTAL LEUCOCYTE COUNT (TLC) by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	14450 ^H	/cmm	4000 - 11000
DIFFERENTIAL LEUCOCYTE COUNT (DLC)			
NEUTROPHILS by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	60	%	50 - 70
LYMPHOCYTES	31	%	20 - 40



CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY & MICROBIOLOGY)

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Test Name	Value	Unit	Biological Reference interval				
by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY							
EOSINOPHILS	2	%	1 - 6				
by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY							
MONOCYTES	7	%	2 - 12				
by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	0	0/	0 1				
BASOPHILS by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	0	%	0 - 1				
ABSOLUTE LEUKOCYTES (WBC) COUNT							
ABSOLUTE NEUTROPHIL COUNT	8670 ^H	/cmm	2000 - 7500				
by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	8070-	/ CIIIII	2000 1000				
ABSOLUTE LYMPHOCYTE COUNT	4480	/cmm	800 - 4900				
by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY							
ABSOLUTE EOSINOPHIL COUNT	289	/cmm	40 - 440				
by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY			00.000				
ABSOLUTE MONOCYTE COUNT by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY	1012 ^H	/cmm	80 - 880				
ABSOLUTE BASOPHIL COUNT	0	/cmm	0 - 110				
by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY		7 011111	0 110				
PLATELETS AND OTHER PLATELET PREDICTIVE MARKERS.							
PLATELET COUNT (PLT)	439000	/cmm	150000 - 450000				
by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE							
PLATELETCRIT (PCT)	0.36	%	0.10 - 0.36				
by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE		ĆŢ.	0.50 10.0				
MEAN PLATELET VOLUME (MPV) by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	8	fL	6.50 - 12.0				
PLATELET LARGE CELL COUNT (P-LCC)	69000	/cmm	30000 - 90000				
by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE	03000	/ CIIIII	30000 - 30000				
PLATELET LARGE CELL RATIO (P-LCR)	15.6	%	11.0 - 45.0				
by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE							
PLATELET DISTRIBUTION WIDTH (PDW)	15.7	%	15.0 - 17.0				
by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE							
NOTE: TEST CONDUCTED ON EDTA WHOLE BLOOD							



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Value Unit **Biological Reference interval Test Name**

CLINICAL CHEMISTRY/BIOCHEMISTRY

CHOLESTEROL: SERUM

CHOLESTEROL TOTAL: SERUM OPTIMAL: < 200.0 239.15^{H} mg/dL

by CHOLESTEROL OXIDASE PAP BORDERLINE HIGH: 200.0 -

239.0

HIGH CHOLESTEROL: > OR =

240.0

INTERPRETATION:

NATIONAL LIPID ASSOCIATION RECOMMENDATIONS (NLA-2014)	CHOLESTEROL IN ADULTS (mg/dL)	CHOLESTEROL IN ADULTS (mg/dL)	
DESIRABLE	< 200.0	< 170.0	
BORDERLINE HIGH	200.0 - 239. 0	171.0 – 199.0	
HIGH	>= 240.0	>= 200.0	

1. Measurements in the same patient can show physiological & analytical variations. Three serial samples 1 week apart are recommended for Total Cholesterol, Triglycerides, HDL & LDL Cholesterol.

2. As per National Lipid association - 2014 guidelines, all adults above the age of 20 years should be screened for lipid status. Selective screening of children above the age of 2 years with a family history of premature cardiovascular disease or those with at least one parent with high total cholesterol is recommended.



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Value Unit **Biological Reference interval Test Name**

IMMUNOPATHOLOGY/SEROLOGY WIDAL SLIDE AGGLUTINATION TEST

SALMONELLA TYPHI O by SLIDE AGGLUTINATION	1:80	TITRE	1:80
SALMONELLA TYPHI H by SLIDE AGGLUTINATION	1:40	TITRE	1:160
SALMONELLA PARATYPHI AH by SLIDE AGGLUTINATION	NIL	TITRE	1:160
SALMONELLA PARATYPHI BH	NIL	TITRE	1:160

INTERPRETATION:

CLIENT CODE.

- 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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