PKR JAIN HEALTHCARE INSTITUTE NASIRPUR, Hissar Road, AMBALA CITY- (Haryana) A PIONEER DIAGNOSTIC CENTRE

【 0171-2532620, 8222896961 🛛 🖾 pkrjainhealthcare@gmail.com

NAME	: Miss. SHIVANI			
AGE/ GENDER	: 24 YRS/FEMALE		PATIENT ID	: 1681282
COLLECTED BY	:		REG. NO./LAB NO.	: 122411250014
REFERRED BY	:		REGISTRATION DATE	: 25/Nov/2024 10:54 AM
BARCODE NO.	: 12505833		COLLECTION DATE	: 25/Nov/2024 11:08AM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITU	TE	REPORTING DATE	: 25/Nov/2024 12:59PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBAL	A CITY - HA	ARYANA	
Test Name		Value	Unit	Biological Reference interval
		HAEM	IATOLOGY	
	СОМР	PLETE BI	LOOD COUNT (CBC)	
RED BLOOD CELLS	(RBCS) COUNT AND INDICES			
HAEMOGLOBIN (HI by Calorimetric		13.4	gm/dL	12.0 - 16.0
RED BLOOD CELL (I	RBC) COUNT	4.41	Millions/	cmm 3.50 - 5.00
PACKED CELL VOLUME (PCV) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER		40.1	%	37.0 - 50.0
MEAN CORPUSCULAR VOLUME (MCV) by calculated by automated hematology analyzer		91.1	KR fL	80.0 - 100.0
MEAN CORPUSCULAR HAEMOGLOBIN (MCH) by calculated by automated hematology analyzer		30.5	pg	27.0 - 34.0
MEAN CORPUSCULAR HEMOGLOBIN CONC. (MCHC) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER		33.5	g/dL	32.0 - 36.0
RED CELL DISTRIBUTION WIDTH (RDW-CV) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER		13.9	%	11.00 - 16.00
RED CELL DISTRIBUTION WIDTH (RDW-SD) by CALCULATED BY AUTOMATED HEMATOLOGY ANALYZER		47.5	fL	35.0 - 56.0
MENTZERS INDEX by CALCULATED		20.66	RATIO	BETA THALASSEMIA TRAIT: < 13.0 IRON DEFICIENCY ANEMIA: >13.0
GREEN & KING INDEX by CALCULATED		28.82	RATIO	BETA THALASSEMIA TRAIT:< 65.0 IRON DEFICIENCY ANEMIA: > 65.0
WHITE BLOOD CEI	LS (WBCS)			00.0
TOTAL LEUCOCYTE COUNT (TLC) by flow cytometry by sf cube & microscopy		9480	/cmm	4000 - 11000
DIFFERENTIAL LE	<u>UCOCYTE COUNT (DLC)</u>			
NEUTROPHILS by flow cytometry	BY SF CUBE & MICROSCOPY	71 ^H	%	50 - 70
LYMPHOCYTES		23	%	20 - 40

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DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST

NOT VALID FOR MEDICO LEGAL PURPOSE



: Miss. SHIVANI

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Test Name		Value	Unit	Biological Reference interval
by FLOW CYTOMETR	RY BY SF CUBE & MICROSCOPY			
EOSINOPHILS			%	1 - 6
MONOCYTES by FLOW CYTOMETR	MONOCYTES by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY		%	2 - 12
BASOPHILS		0	%	0 - 1
	RY BY SF CUBE & MICROSCOPY OCYTES (WBC) COUNT			
ABSOLUTE NEUTH	ABSOLUTE NEUTROPHIL COUNT by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY		/cmm	2000 - 7500
ABSOLUTE LYMPHOCYTE COUNT by FLOW CYTOMETRY BY SF CUBE & MICROSCOPY		2180	/cmm	800 - 4900
ABSOLUTE EOSIN		0 ^L	/cmm	40 - 440
ABSOLUTE MONO		569	/cmm	80 - 880
ABSOLUTE BASOF		0	/cmm	0 - 110
-	OTHER PLATELET PREDICTIVE	MARKERS.		
PLATELET COUNT by HYDRO DYNAMIC	C (PLT) FOCUSING, ELECTRICAL IMPEDENCE	353000	/cmm	150000 - 450000
PLATELETCRIT (P by HYDRO DYNAMIC	CT) FOCUSING, ELECTRICAL IMPEDENCE	0.32	%	0.10 - 0.36
MEAN PLATELET	VOLUME (MPV) FOCUSING, ELECTRICAL IMPEDENCE	9	fL	6.50 - 12.0
	CELL COUNT (P-LCC)	70000	/cmm	30000 - 90000
	CELL RATIO (P-LCR)	19.9	%	11.0 - 45.0
PLATELET DISTRIBUTION WIDTH (PDW)		15.8	%	15.0 - 17.0



NAME



by HYDRO DYNAMIC FOCUSING, ELECTRICAL IMPEDENCE NOTE: TEST CONDUCTED ON EDTA WHOLE BLOOD

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Test Name	V	alue Unit	Biological Reference interval
	ERYTHROCYT	E SEDIMENTATION RAT	ГЕ (ESR)
by RED CELL AGGREG INTERPRETATION: 1. ESR is a non-specif	GATION BY CAPILLARY PHOTOMETRY	ndicates the presence of inflan	/1st hr 0 - 20
2. An ESR can be affe as C-reactive protein	cted by other conditions besides inflamn be used to monitor disease activity and r	nation. For this reason, the ESR	the above diseases as well as some others, such as
(polycythaemia), sigr	n with conditions that inhibit the normal nificantly high white blood cell count (leu e cell anaemia) also lower the ESR.	sedimentation of red blood ce acocytosis) , and some protein	ells, such as a high red blood cell count abnormalities. Some changes in red cell shape (sucl
1. ESR and C - reactiv 2. Generally, ESR doe 3. CRP is not affected 4. If the ESR is elevat 5. Women tend to ha 6. Drugs such as dext	e protein (C-RP) are both markers of infla as not change as rapidly as does CRP, eith by as many other factors as is ESR, makir ed, it is typically a result of two types of ve a higher ESR, and menstruation and pr ran, methyldopa, oral contraceptives, pe id quinine may decrease it	her at the start of inflammation ng it a better marker of inflamm proteins, globulins or fibrinoge regnancy can cause temporary	nation. en.



TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT

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Test Name		Value	Unit	Biological Reference interva
		CAL CHEMICT	DV /DIOCHEMICT	N1/
	CLINI			RY
	CLINI		RY/BIOCHEMIST	ĸy
	CLINI	GLUCOSE R		ĸy
GLUCOSE RANDOM (F				KY NORMAL: < 140.00 PREDIABETIC: 140.0 - 200.0 DIABETIC: > 0R = 200.0

(after consumption of 75 gms of glucose) is recommended for all such patients. 3. A random 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.





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TITRE

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1:160

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Test Name	Va	lue Unit	Biological Reference interval		
	IMMUNOP	ATHOLOGY/SEROLOG	Y		
	WIDAL SLI	DE AGGLUTINATION TEST			
SALMONELLA TYPHI O 1:80 by SLIDE AGGLUTINATION		80 TITRE	1:80		
SALMONELLA TYP by SLIDE AGGLUTINA		40 TITRE	1:160		

SALMONELLA PARATYPHI AH

by SLIDE AGGLUTINATION SALMONELLA PARATYPHI BH

by SLIDE AGGLUTINATION

INTERPRETATION:

LIMITATIONS:

TEST PERFORMED AT KOS DIAGNOSTIC LAB. AMBALA CANTI

week is considered as a definite evidence of infection. 4.A simultaneous rise in H agglutinins is suggestive of paratyphoid infection.

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.

till 3rd or 4th week, after which it declines gradually. 2.Lower titres may be found in normal individuals.

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.

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

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

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.

1:20

1:20

*** End Of Report ***





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