TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.

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

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

NAME	: Mr. BIPIN			
AGE/ GENDER	: 21 YRS/MALE		PATIENT ID	: 1749658
COLLECTED BY	:		REG. NO./LAB NO.	: 122502080020
REFERRED BY	:		REGISTRATION DATE	: 08/Feb/2025 11:42 AM
BARCODE NO.	: 12506908		COLLECTION DATE	: 08/Feb/2025 12:33PM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITU	ΤЕ	REPORTING DATE	: 08/Feb/2025 02:42PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBAL	A CITY - HA	ARYANA	
Test Name		Value	Unit	Biological Reference interval
		HAEM	ATOLOGY	
	СОМР	LETE BI	OOD COUNT (CBC)	
RED BLOOD CELLS	(RBCS) COUNT AND INDICES			
HAEMOGLOBIN (H)	B)	17.1 ^H	gm/dL	12.0 - 17.0
RED BLOOD CELL (RBC) COUNT OCUSING, ELECTRICAL IMPEDENCE	5.69 ^H	Millions/	cmm 3.50 - 5.00
PACKED CELL VOLU		49.2	%	40.0 - 54.0
MEAN CORPUSCUL		86.6	KR fl	80.0 - 100.0
MEAN CORPUSCUL	AR HAEMOGLOBIN (MCH) UTOMATED HEMATOLOGY ANALYZER	30	pg	27.0 - 34.0
MEAN CORPUSCUL	AR HEMOGLOBIN CONC. (MCHC) UTOMATED HEMATOLOGY ANALYZER	34.7	g/dL	32.0 - 36.0
	UTION WIDTH (RDW-CV) UTOMATED HEMATOLOGY ANALYZER	12.9	%	11.00 - 16.00
	UTION WIDTH (RDW-SD) utomated hematology analyzer	42	fL	35.0 - 56.0
MENTZERS INDEX by CALCULATED		15.22	RATIO	BETA THALASSEMIA TRAIT: < 13.0 IRON DEFICIENCY ANEMIA:
GREEN & KING IND by CALCULATED	DEX	19.6	RATIO	>13.0 BETA THALASSEMIA TRAIT:< 65.0 IRON DEFICIENCY ANEMIA: > 65.0
WHITE BLOOD CE	LLS (WBCS)			
TOTAL LEUCOCYTE	COUNT (TLC) y by sf cube & microscopy	7710	/cmm	4000 - 11000
DIFFERENTIAL LE	<u>UCOCYTE COUNT (DLC)</u>			
NEUTROPHILS by FLOW CYTOMETRY	Y BY SF CUBE & MICROSCOPY	70	%	50 - 70
LYMPHOCYTES		22	%	20 - 40

DR.VINAY CHOPRA CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY & MICROBIOLOGY) MBBS , MD (PATHOLOGY)

DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST

440 Dated 17.5.2012 u/s 80 G OF INCOME TAX ACT. PAN NO. AAAAP1600. **REPORT ATTRACTS THE CONDITIONS PRINTED OVERLEAF (P.T.O.)**



PKR JAIN HEALTHCARE INSTITUTE NASIRPUR, Hissar Road, AMBALA CITY- (Haryana)

A PIONEER DIAGNOSTIC CENTRE

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

NAME	: Mr. BIPIN			
AGE/ GENDER	: 21 YRS/MALE		PATIENT ID	: 1749658
COLLECTED BY	:		REG. NO./LAB NO.	: 122502080020
REFERRED BY	:		REGISTRATION DATE	: 08/Feb/2025 11:42 AM
BARCODE NO.	: 12506908		COLLECTION DATE	: 08/Feb/2025 12:33PM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTIT	TUTE	REPORTING DATE	: 08/Feb/2025 02:42PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMB	ALA CITY - H	ARYANA	
Test Name		Value	Unit	Biological Reference interval
by FLOW CYTOMETR	Y BY SF CUBE & MICROSCOPY			
EOSINOPHILS	Y BY SF CUBE & MICROSCOPY	1	%	1 - 6
MONOCYTES		7	%	2 - 12
•	Y BY SF CUBE & MICROSCOPY	0		
BASOPHILS by FLOW CYTOMETR	Y BY SF CUBE & MICROSCOPY	0	%	0 - 1
	CYTES (WBC) COUNT			
ABSOLUTE NEUTR		5397	/cmm	2000 - 7500
by FLOW CYTOMETR ABSOLUTE LYMPH	Y BY SF CUBE & MICROSCOPY	1696	/cmm	800 - 4900
	Y BY SF CUBE & MICROSCOPY	1090		800 - 4900
ABSOLUTE EOSING	OPHIL COUNT Y BY SF CUBE & MICROSCOPY	77	/cmm	40 - 440
ABSOLUTE MONOC		540	/cmm	80 - 880
by FLOW CYTOMETR	Y BY SF CUBE & MICROSCOPY			
ABSOLUTE BASOP	HIL COUNT y by sf cube & microscopy	0	/cmm	0 - 110
-	OTHER PLATELET PREDICTIVE	MARKERS.		
PLATELET COUNT		86000 ^L	/cmm	150000 - 450000
	FOCUSING, ELECTRICAL IMPEDENCE	0.10	%	0.10 0.00
PLATELETCRIT (PC by HYDRO DYNAMIC F	51) FOCUSING, ELECTRICAL IMPEDENCE	0.13	%	0.10 - 0.36
MEAN PLATELET V		15 ^H	fL	6.50 - 12.0
-	FOCUSING, ELECTRICAL IMPEDENCE CELL COUNT (P-LCC)	50000	/cmm	30000 - 90000
	FOCUSING, ELECTRICAL IMPEDENCE	50000	/ CIIIII	30000 - 30000
PLATELET LARGE	CELL RATIO (P-LCR) FOCUSING, ELECTRICAL IMPEDENCE	58.3 ^H	%	11.0 - 45.0
PLATELET DISTRI	BUTION WIDTH (PDW)	16.5	%	15.0 - 17.0
-	FOCUSING, ELECTRICAL IMPEDENCE			
NOTE: TEST CONDU	ICTED ON EDTA WHOLE BLOOD			



DR.VINAY CHOPRA CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY & MICROBIOLOGY) MBBS , MD (PATHOLOGY)

DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST

440 Dated 17.5.2012 u/s 80 G OF INCOME TAX ACT. PAN NO. AAAAP1600. **REPORT ATTRACTS THE CONDITIONS PRINTED OVERLEAF (P.T.O.)**



PKR JAIN HEALTHCARE INSTITUTE NASIRPUR, Hissar Road, AMBALA CITY- (Haryana)

NASIRPUR, Hissar Road, AMBALA CITY- (Haryana) A PIONEER DIAGNOSTIC CENTRE

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

NAME	: Mr. BIPIN			
AGE/ GENDER	: 21 YRS/MALE	РАТ	TENT ID	: 1749658
COLLECTED BY	:	REG	. NO./LAB NO.	: 122502080020
REFERRED BY	:	REG	ISTRATION DATE	: 08/Feb/2025 11:42 AM
BARCODE NO.	: 12506908	COL	LECTION DATE	: 08/Feb/2025 12:33PM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INST	ITUTE REP	ORTING DATE	: 08/Feb/2025 07:09PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AM	BALA CITY - HARYAN	NA	
Test Name		Value	Unit	Biological Reference interval
	IMM	UNOPATHOL	GY/SEROLOGY	1
	· · · · ·	C-REACTIVE PRO	JIEIN (URP)	
SERUM by NEPHLOMETRY	EIN (CRP) QUANTITATIVE:	0.35	mg/L	0.0 - 6.0
SERUM by NEPHLOMETRY INTERPRETATION: 1. C-reactive protein	EIN (CRP) QUANTITATIVE:	0.35 acute-phase reactant	mg/L	
SERUM by NEPHLOMETRY INTERPRETATION: 1. C-reactive protein 2. CRP levels can incr	EIN (CRP) QUANTITATIVE:	0.35 acute-phase reactant	mg/L	0.0 - 6.0 n, inflammation, surgery, or neoplastic
SERUM by NEPHLOMETRY INTERPRETATION: 1. C-reactive protein 2. CRP levels can incr proliferation. 3. CRP levels (Quanti	EIN (CRP) QUANTITATIVE: (CRP) is one of the most sensitive rease dramatically (100-fold or mo	0.35 acute-phase reactant ore) after severe trau tivity of inflammatory	mg/L ts for inflammation. ma, bacterial infection	

4. As compared to ESR, CRP shows an earlier rise in inflammatory disorders which begins in 4-6 hrs, the intensity of the rise being higher than ESR and the recovery being earlier than ESR. Unlike ESR, CRP levels are not influenced by hematologic conditions like Anemia, Polycythemia etc., 5. Elevated values are consistent with an acute inflammatory process. NOTE:

1. Elevated C-reactive protein (CRP) values are nonspecific and should not be interpreted without a complete clinical history.

2. Oral contraceptives may increase CRP levels.





DR.VINAY CHOPRA CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY & MICROBIOLOGY)

DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST MBBS , MD (PATHOLOGY)

440 Dated 17.5.2012 u/s 80 G OF INCOME TAX ACT. PAN NO. AAAAP1600, REPORT ATTRACTS THE CONDITIONS PRINTED OVERLEAF (P.T.O.)



PKR JAIN HEALTHCARE INSTITUTE NASIRPUR, Hissar Road, AMBALA CITY- (Haryana)

A PIONEER DIAGNOSTIC CENTRE

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

	: Mr. BIPIN			
AGE/ GENDER	: 21 YRS/MALE		PATIENT ID	: 1749658
COLLECTED BY	:		REG. NO./LAB NO.	: 122502080020
REFERRED BY	:		REGISTRATION DATE	: 08/Feb/2025 11:42 AM
BARCODE NO.	: 12506908		COLLECTION DATE	: 08/Feb/2025 12:33PM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INST	ITUTE	REPORTING DATE	: 08/Feb/2025 07:09PM
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AM	BALA CITY - HA	RYANA	
Test Name		Value	Unit	Biological Reference interval
	RHEUMATOII) FACTOR (F	RA): QUANTITATIVE	- SERUM
RHEUMATOID (RA) SERUM by nephlometry) FACTOR QUANTITATIVE:	1.39	IU/mL	NEGATIVE: < 18.0 BORDERLINE: 18.0 - 25.0 POSITIVE: > 25.0
 Rheumatoid factor Over 75% of patier useful although it ma Inflammatory Marl The titer of RF corr 	rs (RF) are antibodies that are direct nts with rheumatoid arthritis (RA) ay not be etiologically related to RA kers such as ESR & C-Reactive prot relates poorly with disease activity	have an IgM an A. ein (CRP) are no , but those patie	tibody to IgG immunoglobu prmal in about 60 % of patie ents with high titers tend to	Ilin. This autoantibody (RF) is diagnostically
 Rheumatoid factor Over 75% of patier useful although it ma Inflammatory Marl The titer of RF corr The test is useful f Rheumatoid Arthir nembrane lining (syn The disease spreda The diagnosis of R. measurement of RA factor is not spe Non rheumatoid and RA factor is not spe Non rheumatoid and A patients with variou 	rs (RF) are antibodies that are direct nts with rheumatoid arthritis (RA) ay not be etiologically related to RA kers such as ESR & C-Reactive prot relates poorly with disease activity for diagnosis and prognosis of rheue RTIS: ritis is a systemic autoimmune dise novium) joints which ledas to prop as from small to large joints, with a A is primarily based on clinical, ra actor. TIVE): - ceific for Rheumatoid arthiritis, as it nor rheumatoid arthritis (RA) population on control and 8% of nonrheum us nonrheumatoid diseases, characta	have an IgM an A. ein (CRP) are no , but those patie umatoid arthriti ease that is mul greassive joint de greatest damage diological & imr <i>is often present i</i> <i>tions are not clea</i> <i>hatoid patients h</i> <i>erized by chronic</i>	tibody to IgG immunoglobu ormal in about 60 % of patie ents with high titers tend to is. ti-functional in origin and i estruction and in most case e in early phase. nunological features. The n in healthy individuals with o arly separate with regard to have a positive titer). inflammation may have posi-	Ilin. This autoantibody (RF) is diagnostically ents with positive RA. have more severe disease course. s characterized by chronic inflammation of t is to disability and reduction of quality life. host frequent serological test is the ther autoimmune diseases and chronic infection the presence of rheumatoid factor (RF) (15% of sitive tests for RF. These diseases include system
 Over 75% of patier useful although it ma Inflammatory Marl The titer of RF corr The test is useful f RHEUMATOID ARTHIR Rheumatoid Arthir membrane lining (syr The disease spreda The diagnosis of R, measurement of RA fa CAUTION (FALSE POS) RA factor is not spe Non rheumatoid an RA patients have a no Patients with variou, Anti-CCP have been specific (98%) than RA Upto 30 % of patier 	rs (RF) are antibodies that are direct nts with rheumatoid arthritis (RA) ay not be etiologically related to RA kers such as ESR & C-Reactive prot relates poorly with disease activity for diagnosis and prognosis of rheu RTIS: ritis is a systemic autoimmune dise novium) joints which ledas to prog as from small to large joints, with of A is primarily based on clinical, ra actor. TIVE):- ecific for Rheumatoid arthiritis, as it for rheumatoid arthritis (RA) population on reactive titer and 8% of nonrheum us nonrheumatoid diseases, character polymyositis, tuberculosis, syphilis, n discovered in joints of patients with	have an IgM and A. but those patie umatoid arthriti ease that is mul gressive joint de greatest damage diological & imm is often present tions are not clea hatoid patients h erized by chronic viral hepatitis, in h RA, but not in c arthiritis also sho	tibody to IgG immunoglobu ormal in about 60 % of patie ents with high titers tend to is. ti-functional in origin and i estruction and in most case e in early phase. nunological features. The n in healthy individuals with o arly separate with regard to have a positive titer). inflammation may have po- nfectious mononucleosis, an other form of joint disease. A pow Anti-CCP antibodies.	Ilin. This autoantibody (RF) is diagnostically ents with positive RA. have more severe disease course. s characterized by chronic inflammation of the sto disability and reduction of quality life. host frequent serological test is the ther autoimmune diseases and chronic infection the presence of rheumatoid factor (RF) (15% of sitive tests for RF. These diseases include system d influenza. nti-CCP2 is HIGHLY SENSITIVE (71%) & more





DR.VINAY CHOPRA CONSULTANT PATHOLOGIST MBBS, MD (PATHOLOGY & MICROBIOLOGY) MBBS , MD (PATHOLOGY)

DR.YUGAM CHOPRA CONSULTANT PATHOLOGIST

TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.

440 Dated 17.5.2012 u/s 80 G OF INCOME TAX ACT. PAN NO. AAAAP1600. **REPORT ATTRACTS THE CONDITIONS PRINTED OVERLEAF (P.T.O.)**

