



		Chopra v & Microbiology) onsultant Pathologi		(Pathology)	
NAME	: Mr. DEEPAK				
AGE/ GENDER	: 53 YRS/MALE		PATIENT ID	: 1572587	
COLLECTED BY	: SURJESH		REG. NO./LAB NO.	: 012408060057	
REFERRED BY	:		<b>REGISTRATION DATE</b>	: 06/Aug/2024 02:48 PM	
BARCODE NO.	: 01514605		COLLECTION DATE	: 06/Aug/2024 03:43PM	
CLIENT CODE.	: KOS DIAGNOSTIC LAB		<b>REPORTING DATE</b>	: 06/Aug/2024 05:04PM	
CLIENT ADDRESS	: 6349/1, NICHOLSON ROA	D, AMBALA CANTI	ſ		
Test Name		Value	Unit	Biological Reference interval	
		MMUNOPATH	IOLOGY/SEROLOGY		
	HEP	ATITIS C VIRUS	(HCV) ANTIBODY: TOT	AL	
HEPATITIS C ANTIBODY (HCV) TOTAL: SERUM by CMIA (CHEMILUMINESCENT MICROPARTICLE IMMUNOAS		0.06 DASSAY)	S/CO	NEGATIVE: < 1.00 POSITIVE: > 1.00	
HEPATITIS C ANTIBC	DDY (HCV) TOTAL	NON - RE	ACTIVE		
RESULT					
by CMIA (CHEMILUMIN INTERPRETATION:-	IESCENT MICROPARTICLE IMMUN	DASSAY)			
RESULT (INDEX)			REMARKS		
< 1.00			NON - REACTIVE/NOT - DETECTED		
> =1.00			REACTIVE/ASYMPTOMATIC/INFECTIVE STATE/CARRIER STATE.		
needle punctures in compared to HAV & I	healthcare workers, dialysis pa	atients and rarely fi CV occurs in 85 % o	rom mother to infant. 10 % f infected individuals. In hig	ntation, injection drug abusers, accidental of new cases show sexual transmission. As h risk population, the predictive value of Ant	

2. Routine screening of low and high prevelance population including blood donors.

NOTE:

1. False positive results are seen in Auto-immune disease, Rheumatoid Factor, HYpergammaglobulinemia, Paraproteinemia, Passive antibody transfer, Anti-idiotypes and Anti-superoxide dismutase.

2. False negative results are seen in early Acute infection, Immunosuppression and Immuno-incompetence.

3. HCV-RNĂ PCR recommended in all reactive results to differentiate between past and present infection.





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BARCODE NO.	: 01514605	C	OLLECTION DATE	: 06/Aug/2024 03:43PM
CLIENT CODE.	: KOS DIAGNOSTIC LAB	R	EPORTING DATE	: 07/Aug/2024 08:19AM
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, A	AMBALA CANTT		
Test Name		Value	Unit	Biological Reference interval
	ANTI NU	JCLEAR ANTIBO	DY/FACTOR (ANA/A	NF)
	IBODIES (ANA): SERUM	0.5	INDEX VA	LUE NEGATIVE: < 1.0 BORDERLINE: 1.0 - 1.20

## **INTERPRETATION:-**

1.For diagnostic purposes, ANA value should be used as an adjuvant to other clinical and laboratory data available.

2. Measurement of antinuclear antibodies (ANAs) in serum is the most commonly performed screening test for patients suspected of having a systemic rheumatic disease, also referred to as connective tissue disease.

3.ANAs occur in patients with a variety of autoimmune diseases, both systemic and organ-specific. They are particularly common in the systemic rheumatic diseases, which include lupus erythematosus (LE), discoid LE, drug-induced LE, mixed connective tissue disease, Sjogren syndrome scleroderma (systemic sclerosis), CREST (calcinosis, Raynaud's phenomenon, esophageal dysmotility, sclerodactyly, telangiectasia) syndrome, polymyositis/dermatomyositis, and rheumatoid arthritis. NOTE:

1. The diagnosis of a systemic rheumatic disease is based primarily on the presence of compatible clinical signs and symptoms.

The results of tests for autoantibodies including ANA and specific autoantibodies are ancillary. Additional diagnostic criteria include consistent histopathology or specific radiographic findings. Although individual systemic rheumatic diseases are relatively uncommon, a great many patients present with clinical findings that are compatible with a systemic rheumatic disease ANA screening may be useful for ruling out the disease.

2.Secondary, disease specific auto antibodies maybe ordered for patients who are screen positive as ancillary aids for the diagnosis of specific auto-immune disorders.



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<b>REFERRED BY</b>	:				
BARCODE NO.	:01514605		COLLECTION DATE	: 06/Aug/2024 03:43PM	
CLIENT CODE.	: KOS DIAGNOSTIC LAB		<b>REPORTING DATE</b>	: 07/Aug/2024 11:39AM	
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAI	), AMBALA CANTT			
Test Name		Value	Unit	Biological Reference interval	
	HEP	ATITIS E VIRUS (	(HEV) ANTIBODY: TOT	AL	
HEPATITIS E ANTBODY (HEV) TOTAL 1.58 <sup>H</sup> 2UANTITATIVE by ELISA (ENZYME LINKED IMMUNOASSAY)		1.58 <sup>H</sup>	AI	< 0.90	
HEPATITIS E ANTBODY (HEV) TOTAL		REACTIVE		NON - REACTIVE	
RESULT by ELISA (ENZYME LI INTERPRETATION:	INKED IMMUNOASSAY)				
NEGATIVE			AI	< 0.90	
EQUIVOCAL			AI	0.90 - 1.10	
	POSITIVE		AI	>1.10	

1.Hepatitis E virus is a positive-sense single-stranded RNA icosahedral virus. 2.It usuallsy causes a self limiting hepatitis which results in complete remission.

3. Occasional cases of fulminant hepatic necrosis are known to be associated with the infection. Transmission is mainly feco-oral.

4. The average incubation period for the infection is 3-8 weeks from the time of exposure.

5.IgM antibodies become detectable in the serum prior to the onset of clinically identifiable disease and if detected, they are indicative of a recent infection.



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TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.





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CLIENT ADDRESS	: 6349/1, NICHOLSON ROA	D, AMBALA CANTT		
Test Name		Value	Unit	<b>Biological Reference interval</b>
	НЕРА	TITIS B SURFACE ANT	TIGEN (HBsAg) UL	TRA
	HEPATITIS B SURFACE ANTIGEN (HBsAg): SERUM			
SERUM		0.17	S/CO	NEGATIVE: < 1.0 POSITIVE: > 1.0
SERUM by CMIA (CHEMILUMII HEPATITIS B SURFAC RESULT	NESCENT MICROPARTICLE IMMUNG CE ANTIGEN (HBSAg)	DASSAY) NON REACTIVE	S/CO	
SERUM by CMIA (CHEMILUMII HEPATITIS B SURFA) RESULT by CMIA (CHEMILUMII	NESCENT MICROPARTICLE IMMUNO	DASSAY) NON REACTIVE	S/CO	
SERUM by CMIA (CHEMILUMII HEPATITIS B SURFA) RESULT by CMIA (CHEMILUMII INTERPRETATION:	NESCENT MICROPARTICLE IMMUNG CE ANTIGEN (HBSAg)	DASSAY) NON REACTIVE	S/CO REMARKS	
SERUM by CMIA (CHEMILUMII HEPATITIS B SURFA) RESULT by CMIA (CHEMILUMII INTERPRETATION: RESUI	NESCENT MICROPARTICLE IMMUNG CE ANTIGEN (HBSAg) NESCENT MICROPARTICLE IMMUNG	DASSAY) NON REACTIVE		

Hepatitis B Virus (HBV) is a member of the Hepadna virus family causing infection of the liver with extremely variable clinical features. Hepatitis B is transmitted primarily by body fluids especially serum and also spread effectively sexually and from mother to baby. In most individuals HBV hepatitis is self limiting, but 1-2 % normal adolescent and adults develop Chronic Hepatitis. Frequency of chronic HBV infection is 5-10% in immunocompromised patients and 80 % neonates. The initial serological marker of acute infection is HBsAg which typically appears 2-3 months after infection and disappears 12-20 weeks after onset of symtoms. Persistence of HBsAg for more than 6 months indicates carrier state or Chronic Liver disease.





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BARCODE NO.			COLLECTION DATE	:06/Aug/202403:43PM	
CLIENT CODE.	: KOS DIAGNOSTIC LAB		<b>REPORTING DATE</b>	: 07/Aug/2024 11:45AM	
CLIENT ADDRESS	: 6349/1, NICHOLSON ROA	D, AMBALA CANTT			
Test Name		Value	Unit	Biological Reference interval	
	HEP	ATITIS A VIRUS (	HAV) ANTIBODY: TOT	AL	
HEPATITIS A ANTIBODY (HAV) TOTAL QUANTITATIVE by Elisa (ENZYME LINKED IMMUNOASSAY)		1.674 <sup>H</sup>	AI	< 0.90	
HEPATITIS A ANTIBODY (HAV) TOTAL		REACTIVE		NON - REACTIVE	
RESULT by ELISA (ENZYME LI <u>INTERPRETATION</u>	INKED IMMUNOASSAY)				
	HEPATIT	TIS A VIRUS (HAV) TO	TAL ANTIBODIES		
	NON REACTIVE		< 0.90		
	EQUIVOCAL		0.90 - 1.10		

KOS Diagnostic Lab (A Unit of KOS Healthcare)

 POSITIVE
 >1.10

 1.Hepatitis A virus is a non-enveloped RNA virus that is classified as picorna virus. It usually causes a self limiting hepatitis which results in complete remission.

2. Occasional cases of fulminant hepatic necrosis are known to be associated with the infection. Transmission is mainly oro-faecal.

3. The incubation period is between 15-50 days from the time of exposure.

4.IgM antibody is only present in the blood following an acute hepatitis A infection and is a fairly reliable marker of a recent infection. It is detectable from one to two weeks after the initial infection and persists for up to 14 weeks after exposure.

\* \* \* End Of Report \* \* \*





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