



	Dr. Vinay Cl MD (Pathology Chairman & Co		Dr. Yugam Chopra MD (Pathology) CEO & Consultant Pathologist		
NAME : I	Mr. MANMEET				
AGE/ GENDER : 3	32 YRS/MALE	P	ATIENT ID	: 1593932	
OLLECTED BY :		R	EG. NO./LAB NO.	: 012408280032	
REFERRED BY			REGISTRATION DATE	: 28/Aug/2024 10:48 AM	
	1515007			0	
	01515867		OLLECTION DATE	: 28/Aug/2024 10:49AM	
	KOS DIAGNOSTIC LAB		EPORTING DATE	: 28/Aug/2024 11:48AM	
CLIENT ADDRESS : 6	3349/1, NICHOLSON ROAD	, AMBALA CANTT			
Test Name		Value	Unit	Biological Reference interval	
SGOT/AST: SERUM by IFCC, WITHOUT PYRIDO)XAL PHOSPHATE	33.9	U/L	7.00 - 45.00	
SGPT/ALT: SERUM		67.8 ^H	U/L	0.00 - 49.00	
by IFCC, WITHOUT PYRID	DXAL PHOSPHATE	0.5			
	OPHOTOMETRY	0.5			
by CALCULATED, SPECTR					
by CALCULATED, SPECTR INTERPRETATION NOTE:- To be correlated in USE:- Differential diagno: INCREASED:-	n individuals having SGOT ar sis of diseases of hepatobil	nd SGPT values highe ary system and pan	creas.	Range.	
by CALCULATED, SPECTR INTERPRETATION NOTE:- To be correlated in JSE:- Differential diagno: NCREASED:- DRUG HEPATOTOXICITY	n individuals having SGOT ar	nd SGPT values highe	> 2		
by CALCULATED, SPECTR INTERPRETATION NOTE:- To be correlated in JSE:- Differential diagnos NCREASED:- DRUG HEPATOTOXICITY ALCOHOLIC HEPATITIS	n individuals having SGOT ar	nd SGPT values highe ary system and pan	creas. > 2 > 2 (Highly Sugges		
by CALCULATED, SPECTR INTERPRETATION VOTE:- To be correlated in JSE:- Differential diagno: NCREASED:- DRUG HEPATOTOXICITY ALCOHOLIC HEPATITIS CIRRHOSIS	n individuals having SGOT ar sis of diseases of hepatobil	nd SGPT values highe ary system and pan	> 2 > 2 (Highly Sugges 1.4 - 2.0		
by CALCULATED, SPECTR INTERPRETATION NOTE:- To be correlated in USE:- Differential diagno: INCREASED:- DRUG HEPATOTOXICITY ALCOHOLIC HEPATITIS CIRRHOSIS INTRAHEPATIC CHOLEST	n individuals having SGOT ar sis of diseases of hepatobil	iary system and pan	> 2 > 2 (Highly Sugges 1.4 - 2.0 > 1.5	stive)	
INTERPRETATION NOTE:- To be correlated in USE:- Differential diagno: INCREASED:- DRUG HEPATOTOXICITY ALCOHOLIC HEPATITIS CIRRHOSIS INTRAHEPATIC CHOLEST	n individuals having SGOT ar sis of diseases of hepatobil	iary system and pan	> 2 > 2 (Highly Sugges 1.4 - 2.0	stive)	
by CALCULATED, SPECTR INTERPRETATION NOTE:- To be correlated in USE:- Differential diagno: INCREASED:- DRUG HEPATOTOXICITY ALCOHOLIC HEPATITIS CIRRHOSIS INTRAHEPATIC CHOLEST HEPATOCELLULAR CARCI DECREASED:- 1. Acute Hepatitis due to	n individuals having SGOT ar sis of diseases of hepatobil ATIS INOMA & CHRONIC HEPATITI virus, drugs, toxins (with AS	ST increased 3 to 10	> 2 > 2 (Highly Sugges 1.4 - 2.0 > 1.5 > 1.3 (Slightly Inc.)	stive)	
by CALCULATED, SPECTR INTERPRETATION NOTE:- To be correlated in USE:- Differential diagno: INCREASED:- DRUG HEPATOTOXICITY ALCOHOLIC HEPATITIS CIRRHOSIS INTRAHEPATIC CHOLEST HEPATOCELLULAR CARCI DECREASED:- 1. Acute Hepatitis due to	n individuals having SGOT ar sis of diseases of hepatobil ATIS INOMA & CHRONIC HEPATITI	ST increased 3 to 10	> 2 > 2 (Highly Sugges 1.4 - 2.0 > 1.5 > 1.3 (Slightly Inc.)	stive)	

NORMAL	< 0.65		
GOOD PROGNOSTIC SIGN	0.3 - 0.6		
POOR PROGNOSTIC SIGN	1.2 - 1.6		







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

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







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Test Name		Value	Unit	Biological Reference interval
		URIC ACIE		
JRIC ACID: SERUM		8.5 ^H	mg/dL	3.60 - 7.70
by URICASE - OXIDAS	SE PEROXIDASE	0.5	J	
3.Cytolytic treatment 4.Polycythemai vera 5.Psoriasis. 6.Sickle cell anaemia (B).DUE TO DECREASE 1.Alcohol ingestion. 2.Thiazide diuretics. 3.Lactic acidosis. 4.Aspirin ingestion (lu 5.Diabetic ketoacido: 6.Renal failure due to DECREASED:- (A).DUE TO DIETARY E 1.Dietary deficiency of 2.Fanconi syndrome 3.Multiple sclerosis.	urines (organ meats,legumes,and t of malignancies especially leuke & myeloid metaplasia. etc. D EXCREATION (BY KIDNEYS) ess than 2 grams per day). sis or starvation. o any cause etc. DEFICIENCY of Zinc, Iron and molybdenum. & Wilsons disease.	emais & lymphomas.		
(B). DUE TO INCREASE				ids and ACTH, anti-coagulants and estrogens et
		** End Of Report		





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

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

 KOS Central Lab: 6349/1, Nicholson Road, Ambala Cantt -133 001, Haryana

 KOS Molecular Lab: IInd Floor, Parry Hotel, Staff Road, Opp. GPO, Ambala Cantt -133 001, Haryana

 0171-2643898, +91 99910 43898
 care@koshealthcare.com
 www.koshealthcare.com



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