

NAME : MASTER HARJEET SINGH Ref. By : . SELF Address : plot no. 182 TP nagar korba 19 Years / Male Reg No. : 18636

Reg. Date : 30/07/2022 11:47AM

Collected At : MedZone Center

### **INVESTIGATION REPORT**

### **CLINICAL BIOCHEMISTRY**

TEST	RESULT	<u>UNIT</u>	BIOLOGICAL REF RANGE	TEST METHOD
TSH (Thyroid Stimulating Hormone	)			
Sample Type	: SERUM			
TSH (Thyroid Stimulating Hormone)	: 3.31	µIU/mL	0.37 - 4.8 : Adults 0.46 - 8.1 : 1mon–5 Yrs 0.52 -16.0 : 1 – 30 Days	Fully Automated Roche E411 (ECL)

Thyroid-stimulating hormone (TSH, thyrotropin) is a glycoprotein having a molecular weight of approx. 30,000 daltons and consisting of two subunits. The beta-subunit carries the TSH-specific immunological and biological information, whereas the alpha-chain carries species-specific information and has an identical amino acid sequence to the alpha-chains of LH, FSH and hCG. TSH is formed in specific basophil cells of the anterior pituitary and is subject to a circardian secretion sequence. The hypophyseal release of TSH (thyrotropic hormone) is the central regulating mechanism for the biological action of thyroid hormones. TSH has a stimulating action in all stages of thyroid hormone formation and secretion; it also has a proliferative effect. The determination of TSH serves as the initial test in thyroid diagnostics. Even very slight changes in the concentrations of the free thyroid hormones bring about much greater opposite changes in the TSH level. Accordingly, TSH is a very sensitive and specific parameter for assessing thyroid function and is particularly suitable for early detection or exclusion of disorders in the central regulating circuit between the hypothalamus, pituitary and thyroid. Roche Cobas TSH employs monoclonal antibodies specifically directed against human TSH. The antibodies labeled with ruthenium complex\* consist of a chimeric construct from human and mouse-specific components. As a result, interfering effects due to HAMA (human anti-mouse antibodies) are largely eliminated.

METHOD: One-step sandwich and competitive FEIA

INSTRUMENT: TOSHO AIA-360 JAPAN



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### **CLINICAL BIOCHEMISTRY**

TEST	RESULT	<u>UNIT</u>	BIOLOGICAL REF RANGE	TEST METHOD
LFT (Liver Function Test)				
Sample Type	: SERUM			
Bilirubin Total	: 0.75	mg/dl	Adults : 0.1 - 1.2 New born : 0.1 - 12.6	Diazoted Sulfanilic
Bilirubin Direct	: 0.32	mg/dl	Upto 0.4	Diazoted Sulfanilic
Bilirubin Indirect	: 0.43	mg/dl	0.3 - 1.0	
Aspartate Amino Transferase (SGOT)	: 20.5	U/L	Upto 41	IFCC without pyridoxal phosphate
Alanine Amino Transferase (SGPT)	: 33.1	U/L	Upto 40	IFCC without pyridoxal phosphate
Alkaline Phosphatase	: 100.9	U/L	1 month to 9 yrs : 82 - 383 10 yrs to 15 yrs : 42 - 390 16 yrs to 18 yrs : 52 - 171 Adults : 53 - 141	Diethanolamine buffer
Serum Protein	: 7.6	gm/dl	6.0 - 8.3	Biuret
Serum Albumin	: 4.7	gm/dl	3.5 - 5.2	Bromocresol green
Serum Globulin	: 2.9	gm/dl	2.5 - 3.5	
Alb/Glo Ratio	: 1.62		1-2	

LFT: Liver Function tests are a measurement of blood components that provide a lead to the existence, the extent and the type of liver damage.

BILIRUBIN: Bilirubin levels may rise due to hemolysis, failure of conjugating mechanism in the liver, obstruction in the biliary system.

ALKALINE PHOSPHATASE: \*Increase in ALP activity is an index of cholestasis, a blockage of bile flow. \*Increase may also occur in infiltrative diseases of the liver and cirrhosis

TRANSAMINASES (AST & ALT): \*The serum transaminases activities are a measure of the integrity of liver cells. \*They are elevated in acute damage to hepatocytes irrespective of etiology. \*The causes include – hepatitis, toxic injury, drug overdose, shock, severe hypoxia.

SERUM TOTAL PROTEINS: A decrease in serum total proteins indicates a decrease in the liver's synthetic capacity and thus indicates the severity of the liver disease.

METHOD: Spectrophotometry

INSTRUMENT: BS-400 Fully Automated Chemistry Analyser

#### **RENAL FUNCTION TEST**

Sample Type	: SERUM					
Blood urea	: 17.4	mg/dl	10-40	Urease UV		
Serum Creatinine	: 0.84	mg/dl	0.6-1.4	Alkaline Picrate		
Blood Urea Nitrogen	: 8.13	mg/dl	7-21			
Serum Sodium	: 143	mmol/L	136-145	ISE		
Serum Potassium	: 5.09	mmol/L	3.5-5.1	ISE		
chloride	: 106.3	Meq/L	96-106			



# ADVANCE DIAGNOSTICS CENTRE C1-C2/17A, NEAR NIHARIKA TALKIES KORBA- 495677 PH-09228333 MOBILE-9300888178

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Address	: plot no.	182 TP nagar korba			Collected	d At: MedZor	ne Center
			End Of Report	_			
Sample Regist	ered On	: 30/07/2022 11:47/	M			1.	
Sample Receiv	ved On	: 30/07/2022 05:30	M Home	Collec	tion	an.	
Report Release	ed On	: 31/07/2022 11:11/	Μ			Dr VANDANA	CHANDANI
Sample Barc	ode:		Checkec	l By:go	opal	2	



## **ADVANCE DIAGNOSTICS CENTRE** C1-C2/17A, NEAR NIHARIKA TALKIES

KORBA- 495677 PH-09228333 MOBILE-9300888178

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### **INVESTIGATION REPORT**

### HAEMATOLOGY

TEST	<b>RESULT</b>	<u>UNIT</u>	<b>BIOLOGICAL REF RANGE</b>	TEST METHOD
CBP (Complete Blood Picture)				
Sample Type	: WB - EDTA			
Haemoglobin	: 14.5	gm%	12.0 - 18.0	
Total Erythrocyte Count	: 5.28	M/cmm	4.0 - 6.2	Cell Counter
Hemotocrit (PCV)	: 44.6	Vol %	35.0 - 50.0	
Mean Corpuscular Volume	: 84.5	fL	80 - 100	
Mean Corpuscular Hemoglobin	: 27.5	PG	26 - 34	
MCHC	: 32.5	g/L	31 - 35	
RDW	: 12.9	%	11.5 - 14.5	
Total Leucocyte Count.	: 4400	/cumm	4000 - 11000	
DIFFERENTIAL COUNT :				
Neutrophils	: 48	%	40 - 75	
Lymphocytes.	: 39	%	20 - 40	Cell Counter
Monocytes.	: 06	%	2 - 10	Cell Counter
Eosinophils	: 07	%	1 - 6	Cell Counter
Basophils	: 0	%	0 - 1	Cell Counter
Platelet Count	: 183000	/cmm	150000 - 450000	

# ESR (Erythrocyte Sedimentation Rate)

Sample Type	: PLASMA -Na Citrate			
ESR (Erythrocyte Sedimentation Rate)	: 07	mm/hr	0 - 15 :1st Hour	Sedimentation me

# AEC (Absolute Eosinophil Count)

Sample Type	: WB - EDTA	VB - EDTA				
AEC (Absolute Eosinophil Count)	: 308	cells / cmm 40 - 440	Cell counter			

	End Of Report	
: 30/07/2022 11:47AM		1
: 30/07/2022 05:30PM	Home Collection	Dhr.
: 31/07/2022 11:11AM		Dr. VANDANA CHANDANI
	Checked By:gopal	
	: 30/07/2022 11:47AM : 30/07/2022 05:30PM : 31/07/2022 11:11AM	End Of Report : 30/07/2022 11:47AM : 30/07/2022 05:30PM Home Collection : 31/07/2022 11:11AM 



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# INVESTIGATION REPORT

#### IMMUNOLOGY/SEROLOGY

TEST	RESULT	<u>UNIT</u>	BIOLOGICAL REF RANGE	TEST METHOD
Hepatitis C Virus (HCV Antibody)				
Sample Type	: SERUM			
Hepatitis C Virus (HCV Antibody)	: 0.03	s/co	< 1.0 : NEGATIVE > 1.0 : POSITIVE	ELISA

Hepatitis C virus (HCV) is now know to be the major cause of parenterally transmitted non-A, non-B hepatitis. Until the virus was was characterized, diagnosis was made by exclusion of all other known causes of hepatitis. Antibody to HCV is found in over 80% of patients with well documented non-A non-B hepatitis. The Worldwide prevalence of HCV is 0.2 to 2% in blood donors and up to 80% in intravenous- drug users. Seroepidemiologic studies show that the seropevalence of HCV infection in India varies between 0.3% to 11.3%. In a large percentage of HCV cases transmission by transfusion and other parenteral means such as sharing of neediest, occupational exposure to blood and hemdialysis. However in case of half of HCV infections, the route of transmission is unknown. HCV establishes a chronic infection in 50 to 80% of cases. Chronic infection is often asymptomatic even in the presence of liver damage discernible on biopsy. Chronic HCV is characterized by fluctuating alanine aminotransferase (ALT or SGPT) levels and recognizable changes in liver histology. Chronic infection can lead to cirrhosis and hepatocellular carcinoma.

Method: ELISA

INSTRUMENT: Rayto **RT – 2100c** Microplate Reader & Washer

<u>Hepatitis B Surf</u>	ace antigen (HE	BsA <u>g</u> )			Method : E	
Sample Type		: SERUM				
Hepatitis B Surface ant	igen (HBsAg)	: 0.06	s/co	< 1.0	: NEGATIVE	ELECTROCHEMILU
Method : ELISA				> 1.0	: POSITIVE	MI
		End Of	Report			
Sample Registered On	: 30/07/2022 11	:47AM				
Sample Received On	: 30/07/2022 05	:30PM	Home Collec	ction		pri.
Report Released On	: 05/08/2022 12	:31PM			Dr VA	
Sample Barcode :			Checked By:g	opal		