



SHREE VEERABHADRESHWARA DIAGNOSTICS ಶ್ರೀ ವೀರಭದ್ರೇಶ್ವರ ಡಯಾಗ್ನೋಸ್ಟ್ರಿಕ್ಸ್

Near 5 road circle opposite Marikamba Tyres and Rayappa Hulekal School Sirsi (U.K) 581 401 Ph:7892637060 / 9845753744 Email id:shreevd45@gmail.com

NAME: Manju Poojari

AGE: 60Years

SEX:Male

REF BY: Govt. Dialysis Sirsi

DATE:15/06/2024

REF NO:502

Test	Result	Normal Range	Unit
Haemoglobin	:8.0	12.0 - 16.0(F)	gm/dl
		14.0-18.0(M)	gm/dL
W.B.C Count	: 5,900	4,000 – 10,000	cells/cmm
DIFFERENTIAL COUNTS Neutrophils	: 64	40 - 75	%
Lymphocytes	:30	20 - 45	%
Eosinophils	: 04	01 - 06	%
Monocytes	:02	0 - 4	%
Basophils	:00	0 - 1	%
Instrument : Five part different	ial Cell Counter (Minds	rau RC-1800)	
Blood Urea Method: UREASE-GLDH Method	: 97	10 – 40	mg/dl
Creatinine Method:Modified Jaffes Method	:12.1	0.7 – 1.4(M) 0.6 – 1.0 (F)	mg/dl mg/dl
ELECTROLYTES			
Sodium	: 145	135 – 145	mEq/L
Potassium	: 5.1	3.5 – 5.5	mEq/L

Lab Technician



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TEST	RESULT	NORMAL RANGE	UNIT
S.G.P.T Method IFCC Method	: 28	0-40	IU/L
S.G.O.T Method IFCC Method	: 25	0-40	IU/L
Albumin Method : BCG method	: 3.8	3.5-5.5	gm/dl

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Ph:7892637060 / 9845753744 Email id:shreevd45@gmail.com

Patient Name: MR. MANJU POOJARY

Age / Gender: 60 years / Male

Patient ID: 13767

Client: Veerabadreswra Laboratory Sirsi-Ul217

Referral: Dr. GOVT DIALYSIS

Collection Time: Jun 16, 2024, 08:53 a.m. Receiving Time: Jun 16, 2024, 09:07 a.m. Reporting Time: Jun 16, 2024, 09:53 a.m.

Sample ID:

241680025

	THE RESIDENCE OF THE PROPERTY	241680025		
Test Description	Value(s)	Reference Range	Unit(s)	
PHOSPHORUS SERUM				
Phosphorus (Inorganic)* Method : Phosphomolybdate	5.2	2.3 – 5.0	mg/dL	

END OF REPORT

Verified By: Ashritha

Dr. Árjun C.P. MBBS, MD KMC Reg. No. 89655 Consultant Pathologist



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Test Description Value(s) Reference Range Unit(s)

HIV 1 & 2 Antibodies (ELISA)

HIV 1 & 2 Antibodies

Method : ELISA

Non Reactive: 0.10 >1.0 Reactive

<1.0 Non reactive

Internetation

1. A non-reactive result implies that no anti HIV I or Anti II antibodies have been detected in the sample by this method. This means that either the patient has not been exposed to HIV or HIV II infection or the sample has tested during the window phase (before the development of detectable levels of antibodies).

2. A provisionally reactive or borderline reactive result suggests the possibility of HIV I / HIV II infection and confirmatory diagnosis should be done by WESTERN BLOT tests.

END OF REPORT

Verified By: Chandrakala

Dr Madhumati MBBS MD Microbiology

Consultant Microbiologis



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Test Description

Method: ELISA

Value(s)

Reference Range

Unit(s)

HEPATITIS B SURFACE ANTIGEN (HBSAg) (ELISA)

HEPATITIS B SURFACE ANTIGEN (HBSAg)

Non Reactive: 0.11 <1.0 Non Reactive

>1.0 Reactive

Relatik:

· All Reactive results must be confirmed by Neutralizing confirmatory test or by HBV DNA detection assay.

- 1. Reactive test result indicates presence of Hepatitis B Surface Antigen. It cannot differentiate between the stages of Hepatitis B viral
- 2. Non-Reactive test result indicates absence of Hepatitis B Surface Antigen.
- 3. False positive results may be observed in patients receiving mouse monoclonal antibodies, on heparin therapy, on biotin supplements for diagnosis or therapy, presence of heterophilic antibodies in serum or after HBV vaccination for transient period of time.
- 4. False negative reaction may be due to processing of sample collected early in the course of disease or presence of mutant forms of HBsAg.
- 5. For monitoring HBsAg levels, Quantitative HBsAg assay is recommended

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Test Description	Value(s)	Reference Range	Unit(s)
HCV ANTIBODIES -ELISA HCV ANTIBODIES	Non Reactive: 0.18	>1.0 reactive	S/Co
Method : ELISA		<1.0 Non reactive	

Contients:

Hepatitis C Virus was identified in 1989 as the main aetiological agent of non-A, non-B hepatitis (NANBH) accounting for greater than 90% of posttransfusion hepatitis cases. HCV is a spherical virus of about 30-60 nm in diameter with single positive stranded RNA and is related to the family flaviviridae. It is considered to be the major cause of acute chronic hepatitis, liver cirrhosis and hepatocellular carcinoma throughout the world. Antibodies to HCV can be detected throughout virtually the total infection period. Therefore, the use of highly sensitive antibody assays is the primary approach in serodiagnosis of HCV infection. The diagnosis of hepatitis C can be easily made by finding elevated serum ALT levels and presence of anti HCV in serum/plasma.

Specimens with Sample cut of OD values 1.00 are considered reactive. This is an Antibody detection test and results might depend on immune response of the individual. Patients with auto-immune liver diseases may show false reactive results. HCV Antibodies might take 2 weeks to 5 months to appear after acquiring HCV infection. This antibody may never become detectable in 5–10% of patients with acute hepatitis C, and levels of anti-HCV may rarely become undetectable after recovery. In patients with chronic hepatitis C, anti-HCV is detectable in >95% of cases.

END OF REPORT

Verified By: Chandrakala

Dr Madhumati MBBS MD Microbiology 67106

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