

## Sagepath Labs Pvt. Ltd.

Lab Address:- # Plot No. 564 , 1st floor , Buddhanagar , Near Sai Baba Temple Peerzadiguda Boduppal Hyderabad, Telangana. ICMR Reg .No. SAPALAPVLHT (Covid -19)

LABORATORY TEST REPORT

Name	: Mrs. PADMA					
Sample ID	: A0787950					
Age/Gender	: 45 Years/Female	Reg. No	: 0312410250023			
Referred by	: Dr. Chakrapani	SPP Code	: SPL-CV-172			
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 25-Oct-2024 01:19 PM			
Primary Sample	: Whole Blood	Received On	: 25-Oct-2024 04:10 PM			
Sample Tested In	: Serum	Reported On	: 25-Oct-2024 07:24 PM			
Client Address	: Kimtee colony ,Gokul Nagar,Tarnaka	Report Status	: Final Report			

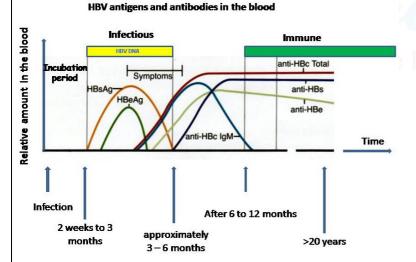
	IMMUNOLOGY & SEROLOGY			
Test Name	Results	Units	Biological Reference Interval	
Hepatitis B Surface Antigen (HBsAg)	0.27	S/Co	<1.00 :Negative >1.00 :Positive	

#### Interpretation:

ITDOSE INFOSYSTEMS PVT. LTD.

- Negative result implies that antibodies to HBsAg have not been detected in the sample. This means the patient has either not been exposed to HBsAg infection or the sample has been tested during the "window phase" i.e. before the development of detectable levels of antibodies. Hence a Non-Reactive result does not exclude the possibility of exposure or infection with HBsAg.
- Positive result implies that antibodies to HBsAg have been detected in the sample.

Hepatitis B Virus (HBV) is a member of the Hepadna virus family causing infections 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 adolescents and adults develop Chronic Hepatitis. Frequency of chronic HBV infection is 5-10% in immunocompromised patients and 80% in 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 symptoms. Persistence of HBsAg for more than six months indicates development of carrier state or Chronic liver disease.



Note:

1. All Reactive results are tested additionally by Specific antibody Neutralization assay . For further confirmation Molecular assays are recommended For diagnostic purposes, results should be used in conjunction with clinical history and other hepatitis markers for Acute or Chronic infection

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



DR. RUTURAJ MANIKLAL KOLHAPURE MD, MICROBIOLOGIST

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LABORATORY TEST REPORT

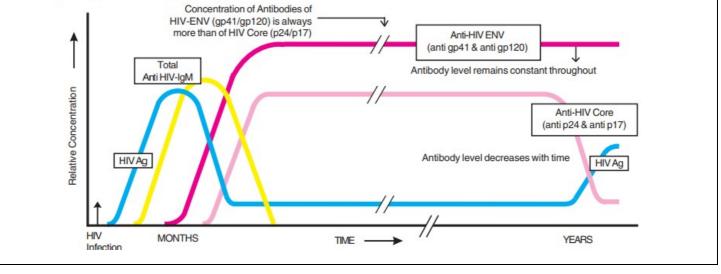
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Sample ID	: A0787950		
Age/Gender	: 45 Years/Female	Reg. No	: 0312410250023
Referred by	: Dr. Chakrapani	SPP Code	: SPL-CV-172
Referring Customer	: V CARE MEDICAL DIAGNOSTICS	Collected On	: 25-Oct-2024 01:19 PM
Primary Sample	: Whole Blood	Received On	: 25-Oct-2024 04:10 PM
Sample Tested In	: Serum	Reported On	: 25-Oct-2024 06:47 PM
Client Address	: Kimtee colony ,Gokul Nagar,Tarnaka	Report Status	: Final Report

	IMMUNOLOGY & SEROLOGY			
Test Name	Results	Units	Biological Reference Interval	
HIV (1& 2) Antibody	0.46	S/Co	< 1.00 : Negative > 1.00 : Positive	

### Interpretation

ITDOSE INFOSYSTEMS PVT. LTD.

- Non Reactive result implies that antibodies to HIV 1 / 2 have not been detected in the sample. This means the patient has either not been exposed to HIV 1 / 2 infection or the sample has been tested during the "window phase" i.e. before the development of detectable levels of antibodies. Hence a Non Reactive result does not exclude the possibility of exposure or infection with HIV 1 / 2.
- Pre and Post test counseling to be done by the concerned referring doctor. The sensitivity and specificity of this test has been determined by National HIV Reference Centers of Govt. of India and WHO collaborating Centers, using various other test panels."
- Reactive samples by ELISA Method are confirmed by 2 other supplemental tests for confirm of HIV infection as per NACO guidelines.
- All patients' reports inderminate should be repeated with a second sample taken 14-28 days. In case the serological results continue to be inderminate the sample should be subject to western blot for confirmation.



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



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