Name : DUMMY : DMY25G01 Lab No.

: DR. DUMMY DUMMY Ref Bv

: 25/2/2025 10:31:00AM Collected Reported

A/c Status

Collected at : PRODUCTION TEST COLLECTION CENTRE

SECTOR - 18, BLOCK-E ROHINI

DELHI 110085

: 25 Years Age : Female Gender

: 25/2/2025 10:38:32AM

Report Status : Interim

Processed at LPL-NATIONAL REFERENCE LAB

> National Reference laboratory, Block E, Sector 18, Rohini, New Delhi -110085

Test Report

Test Name Results Units Bio. Ref. Interval

HIV-2 RNA QUANTITATIVE REAL TIME PCR (Real-time reverse transcriptase PCR)				
HIV-2 RNA	Not Detected			
HIV-2 RNA Quantitative	50.00	IU/mL		
HIV-2 RNA Quantitative	2	Log IU/ml		

Interpretation

	Log IU/mL	Result in IU/ml	REMARKS	Conclusion
	<2	<100	Indicates absence of HIV-2 RNA or it is present below the detection limit of the assay	Negative
	≥2	≥100	Indicates presence of HIV-2 RNA	Positive
	Indetermina	ate	Inhibitors detected in the sample provided	Repeat testing on fresh sample is recommended

^{**}Linear reporting range of this assay is 100 IU/ml to 1 x 109 IU/ml (2.00-9.00 log IU/ml).

Note

- 1. Positive Result: Indicates the presence of HIV-2 RNA in the sample, detected by targeting the Long Terminal Repeat (LTR) region of the HIV-2 genome.
- 2. Negative Result: Indicates the absence of HIV-2 RNA in the sample.
- 3. False-Positive Results: Can occur due to non-specific amplification, contamination, or cross-reactivity with other sequences, leading to the incorrect detection of HIV-2 RNA.
- 4. False-Negative Results: May arise from suboptimal sample collection, low viral load below the detection limit, mutations in the targeted LTR region of HIV-2 RNA, or the presence of inhibitors, causing under quantification or failure to detect the virus.
- 5. Results should always be evaluated in conjunction with the patient's clinical presentation and risk factors.



Name : DUMMY : DMY25G01 Lab No.

: DR. DUMMY DUMMY Ref By

: 25/2/2025 10:31:00AM Collected

A/c Status

Collected at : PRODUCTION TEST COLLECTION CENTRE

SECTOR - 18, BLOCK-E ROHINI

DELHI 110085

: 25 Years Age Gender : Female

: 25/2/2025 10:38:32AM Reported

Report Status : Interim

: LPL-NATIONAL REFERENCE LAB Processed at

> National Reference laboratory, Block E, Sector 18, Rohini, New Delhi -110085

Test Report

Test Name Results Units Bio. Ref. Interval

Usage:

The quantitative HIV-2 viral load assay is used to monitor HIV-2 RNA levels in plasma from patients known to be infected with HIV-2

Dr.Anirudh Bharat Kumar Gupta MD, Microbiology

Senior Consultant Microbiologist NRL - Dr Lal PathLabs Ltd

Dr Shailender Kumar Ph.D. Biotechnology

Research Scientist

Dr Shalabh Malik MD, Microbiology

Technical Director - Microbiology. Infectious Disease Molecular & Serology, Clinical Pathology NRL - Dr Lal PathLabs Ltd

Result/s to follow:

HIV-2 RNA QUANTITATIVE REAL TIME PCR

IMPORTANT INSTRUCTIONS

•Test results released pertain to the specimen submitted.•All test results are dependent on the quality of the sample received by the Laboratory •Laboratory investigations are only a tool to facilitate in arriving at a diagnosis and should be clinically correlated by the Referring Physician .•Report delivery may be delayed due to unforeseen circumstances. Inconvenience is regretted. • Certain tests may require further testing at additional cost for derivation of exact value. Kindly submit request within 72 hours post reporting. Test results may show interlaboratory variations. The Courts/Forum at Delhi shall have exclusive jurisdiction in all disputes/claims concerning the test(s) & or results of test(s). • Test results are not valid for medico legal purposes. This is computer generated medical diagnostic report that has been validated by Authorized Medical Practitioner /Doctor. The report does not need physical signature.

(#) Sample drawn from outside source.

If Test results are alarming or unexpected, client is advised to contact the Customer Care immediately for possible remedial action.

Tel: +91-11-49885050,Fax: - +91-11-2788-2134, E-mail: lalpathlabs@lalpathlabs.com

National Reference lab, Delhi, a CAP (7171001) Accredited, ISO 9001:2015 (FS60411) & ISO 27001:2013 (616691) Certified laboratory.

