

# HIV-1 Quantitative RT-PCR Detection Kit

REF HI-01-20T | 20 Rxns |  -20°C 

Expiry Date - See box label for information

 Please read this IFU carefully before starting

## INTENDED USE

HIV-1 Quantitative RT-PCR Detection Kit is an *in-vitro* RT-PCR quantitative assay for the detection and quantification of HIV-1 (Human Immunodeficiency virus) RNA extracted from human plasma, serum, whole blood along with human endogenous control. The intended user of the product is medical/research professionals in laboratory, healthcare, etc.

## PRINCIPLE OF THE TEST

HIV-1 Quantitative RT-PCR Detection Kit works on one-step, Real-Time polymerase chain reaction (RT-PCR), where the enzymes provided in the master mix converts RNA into cDNA and amplifies with the help of target –specific primers and detection happens through the fluorescent probes bearing reporter and quencher. The intensity of fluorescence is measured at every cycle of reaction with a Real-time PCR thermal cycler data collection unit and the standards provided in the kit will help in quantification of the HIV-1.

## ANALYTICAL SPECIFICATION

HIV-1 Quantitative RT-PCR Detection kit standards are traceable to 4th WHO International Standard for HIV-1 RNA (NIBSC code: 16/194). Limit of detection of HIV-1 is 20 IU/mL (11.6 copies/mL) for ≥95% of the time. The linear range of the HIV-1 Quantitative RT-PCR kit has been observed to detect viral load from 10<sup>7</sup> IU/mL to 20 IU/mL.

## COMPATIBLE INSTRUMENTS

Compatible with Real-Time PCR, instruments like, BIORAD-CFX96, THERMO-QS5, QIAGEN-ROTOR-GENE Q and other instruments which supports CY5 (649 nm - 670 nm) and Texas Red (596 nm - 615 nm)

## PRECAUTIONS

- Treat all the specimens as potentially infectious.
- Wear protective disposable powder-free gloves, a laboratory coat and eye protection when handling specimens.
- Store positive and/or potentially positive material separated from all other components of the kit.
- Keep separate area for master mix and template preparation and work under biosafety cabinets.
- Use aerosol barrier pipette tips and frequently change the gloves.

## STORAGE AND HANDLING

- Store all HIV-1 Quantitative RT-PCR Detection Kit reagents at -20°C.
- Do not repeatedly freeze-thaw reagents more than 5 times it leads to reduced assay sensitivity. Thaw the reagents only on ice or at 4°C.
- Kit components are stable through the end of the expiration date indicated on the box when stored at -20°C. Shelf Life - 12 Months from date of manufacture.

## MATERIALS REQUIRED BUT NOT PROVIDED

**Consumables**  
PCR Plates/tubes  
PCR plate covers/tube caps

## KIT CONTENTS

| Sl.No | Components                     | Volume     | International Units (IU/μL) |
|-------|--------------------------------|------------|-----------------------------|
| 1.    | Master Mix                     | 2 X 750 μL | -                           |
| 2.    | Primer and Probe mix           | 150 μL     | -                           |
| 3.    | Nuclease Free Water            | 1000 μL    | -                           |
| 4.    | Positive Test Control          | 100 μL     | -                           |
| 5.    | Quantitation Standard 1 (QS 1) | 750 μL     | 10,000                      |
| 6.    | Quantitation Standard 2 (QS 2) | 750 μL     | 1000                        |
| 7.    | Quantitation Standard 3 (QS 3) | 750 μL     | 100                         |
| 8.    | Quantitation Standard 4 (QS 4) | 750 μL     | 10                          |
| 9.    | Quantitation Standard 5 (QS 5) | 750 μL     | 1                           |

Table 1:- Kit Content

## FLUORESCENT PROBE DETAILS

| Target                 | Reporter      | Quencher |
|------------------------|---------------|----------|
| HIV-1 Gag protein Gene | Cy5           | BHQ2     |
| Human Beta-Actin       | TEXAS RED/ROX | BHQ2     |

Table 2:- Target, Reporter and Quencher

**\*Note:** In some instruments where the BHQ option is unavailable, please set the quencher to "NONE."

## REACTION MIXTURE 35 μL

| Reagents            | 1 Rxn  | 20 Rxns | 50 Rxns | 100 Rxns |
|---------------------|--------|---------|---------|----------|
| Master Mix          | 15 μL  | 300 μL  | 750 μL  | 1500 μL  |
| Primer & Probe Mix  | 1.5 μL | 30 μL   | 75 μL   | 150 μL   |
| Nuclease Free Water | 3.5 μL | 70 μL   | 175 μL  | 350 μL   |
| Total               | 20 μL  | 400 μL  | 1000 μL | 2000 μL  |

Use 15 μL of the test RNA / Standards per reaction

Table 3:- Reaction Mixture

## REACTION SET-UP

- Thaw all components of the kit on ice, mix gently using vortex and spindown the contents for 5 secs and use it immediately.
- Calculate the number of reactions for each experiment including all controls with one excess reaction volume in the reaction cocktail to accommodate pipetting errors (eg: number of reaction (n) including controls are 10 add 1 extra reaction during the preparation n+1).
- Prepare the reaction mix in a 1.5/2 mL tube for the calculated number of samples referring the table no. 3, in Master Mix Preparation room.
- Assay should be run along with positive controls and negative controls.
- Refer to table no. 3, to prepare the reaction mixture of 35 μL.
- Calculate the number of the samples to be tested along with the controls and multiply with 1 reaction volume extra to get the final reaction mixture.
- Spin down the tubes and dispense 20 μL reaction mix in each tube strips or 96 well plate.
- Before moving to template adding area, add 15 μL of nuclease free water in NTC well.
- Carefully add 15 μL RNA samples kept on ice in the designated wells in template addition room.
- Add 15 μL of Standards (QS) in the designated wells.
- Add 2 μL of PTC in a separate hood and make up the volume by adding 13 μL of nuclease free water.
- Seal the plate carefully, briefly spin down and use any qRT-PCR instrument which complies with the dyes specified in the kit insert.

## THERMAL CYCLING CONDITION

| STEPS                 | TEMP (°C) | TIME   | QUANTITATION | CYCLE |
|-----------------------|-----------|--------|--------------|-------|
| Reverse transcriptase | 45        | 20 min | Off          | 1     |
|                       | 95        | 2 min  | Off          | 1     |
| PCR and Quantitation  | 95        | 15 sec | Off          | 45    |
|                       | 58        | 30 sec | On           |       |

Table 4:- Thermal Cycling Condition

## STANDARD CURVE CONTROL PARAMETER

| Control Parameters         | Valid Value  |
|----------------------------|--------------|
| Slope                      | -3.00 / 3.74 |
| PCR Efficiency             | 85% / 115%   |
| R square (R <sup>2</sup> ) | 0.95-1       |

Table 5:- Curve Control Parameter

## SAMPLE TYPE

- HIV-1 Quantitative RT-PCR Detection Kit developed by NeoDx Biotech Labs Pvt Ltd is a robust and convenient molecular diagnostic device for the Quantitation of HIV-1 RNA in human plasma, serum or whole blood RNA extracts.

- Always use freshly extracted patient RNA, or the RNA samples stored following proper guidelines, to get the optimal results.

## RESULT VALIDATION

| HIV-1 genes                       | HIV-1 Assay result                                                                                                                            |
|-----------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------|
| HIV-1 RNA > 20 IU/mL              | The result is within the acceptance range. The detection probability if HIV-1 RNA is >95%. The Positive test result is statistically ensured. |
| HIV-1 RNA < 20 IU/mL              | The result is outside the acceptance range. The reproducibility of the positive result is not assured.                                        |
| No amplification for HIV-1 target | No HIV-1 RNA was detected.                                                                                                                    |

Table 6:- Result Validation

## DATA ANALYSIS

- After completion of the run, analyze the data as per the RT-PCR manufacturer instructions.
- Analysis should be performed separately for each target using a manual threshold settings.
- In case internal control has not worked for a sample re-do the test with 2 or more dilutions.
- Negative results do not exclude infection and should not be used as the sole basis for the treatment.
- The quantitation standards are defined as IU/μL. The following equation has to be applied to convert the values determined using the standard curve into IU/ mL of sample material.

$$\text{Result (IU/mL)} = \frac{\text{Elution Volume (μL)} \times \text{Result (IU/μL)}}{\text{Sample Volume (mL)}}$$

Sample Volume is the volume of sample used for RNA extraction.

## QUANTITATIVE STANDARD IN IU/μL

| Standards                      | International Units (IU/μL) |
|--------------------------------|-----------------------------|
| Quantitation Standard 1 (QS 1) | 10000                       |
| Quantitation Standard 2 (QS 2) | 1000                        |
| Quantitation Standard 3 (QS 3) | 100                         |
| Quantitation Standard 4 (QS 4) | 10                          |
| Quantitation Standard 5 (QS 5) | 1                           |

Table 7:- Quantitative Standards

## WARNING

### Positive control showed no amplification

Inappropriate storage of reagents

- Store the reagents at recommended temperature for their optimal performance.
- Avoid repetitive freezing and thawing.
- Check the expiry of reagents.

### Negative controls are positive

Causes - Cross-Contamination

- Follow good laboratory practices to avoid contamination issues.
- Use a new batch of reagents and repeat the experiment.

### Abnormal plot and/or low ΔRn values in amplification curve

The baseline was set improperly (some samples have CT values lower than the baseline value)

- Switch from manual to automatic baseline, or move the baseline stop value to a lower CT (2 cycles before the amplification curve for the sample crosses the threshold).

### An amplification signal is detected in the early cycles

Dilute the sample to increase the CT value.

## REFERENCES

1.Kibirige, C.N., Manak, M., King, D. et al. Development of a sensitive, quantitative assay with broad subtype specificity for detection of total HIV-1 nucleic acids in plasma and PBMC. Sci Rep 12, 1550 (2022). <https://doi.org/10.1038/s41598-021-03016-1>

| SYMBOL                                                                                | MEANING                              |
|---------------------------------------------------------------------------------------|--------------------------------------|
|    | For Research Use Only                |
|    | Product Code                         |
|    | Lot Number                           |
|    | Store at -20°C                       |
|    | Contains sufficient for <n> reaction |
|    | Manufacturer                         |
|    | Date of Manufacture (MMM, YYYY)      |
|   | Expiry Date (MMM, YYYY)              |
|  | Instruction for use                  |

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