

**REF** **800304 NeuMoDx™ HIV-1 Calibrators**

**R only**

CAUTION: For US Export Only

**IVD** For *in vitro* diagnostic use with the NeuMoDx 288 and NeuMoDx 96 Molecular System

For insert updates, go to: [www.qiagen.com/neumodx-ifu](http://www.qiagen.com/neumodx-ifu)

For detailed instructions, refer to the NeuMoDx 288 Molecular System Operator's Manual; P/N 40600108 [REF 500100]



For detailed instructions, refer to the NeuMoDx 96 Molecular System Operator's Manual; P/N 40600317 [REF 500200] or P/N 40600655 [REF 500201]

See also the NeuMoDx HIV-1 Quant Test Strip Instructions for Use; P/N 40600412

### INTENDED USE

The NeuMoDx HIV-1 Calibrators are a component of the NeuMoDx HIV-1 Quant Assay, an *in vitro* diagnostic nucleic acid amplification test intended for the detection and quantitation of human immunodeficiency virus type 1 (HIV-1) RNA in human plasma. As implemented on the fully automated NeuMoDx 288 Molecular System or NeuMoDx 96 Molecular System (NeuMoDx System(s)), the NeuMoDx HIV-1 Calibrators are used to establish a calibration coefficient associated with the standard curve of a particular lot of the NeuMoDx HIV-1 Quant Test Strip, allowing for the accurate quantitation of HIV-1 RNA in human plasma specimens. The HIV-1 target in these calibrators are traceable to the WHO 3<sup>rd</sup> HIV-1 International Standard.

### SUMMARY AND EXPLANATION

The NeuMoDx HIV-1 Calibrators are provided in three paired sets of low positive and high positive calibrators. One low positive and one high positive calibrator (1 set) is processed every 90 days or with every new lot of NeuMoDx HIV-1 Quant Test Strips to establish a valid calibration of the NeuMoDx HIV-1 Quant Assay. The HIV-1 target in the calibrators is a non-infectious, replication-defective mammalian recombinant virus containing HIV-1 genome sequences. The recombinant virus is diluted in Basematrix 53 Diluent (Basematrix) (Seracare Life Sciences, Inc., Milford, MA, USA) at final concentrations of 3.0 log<sub>10</sub> IU/mL and 5.0 log<sub>10</sub> IU/mL for the low and high calibrator, respectively.

The NeuMoDx HIV-1 Quant Assay combines automated RNA extraction, amplification, and detection by real-time reverse transcription PCR to enable the quantitative detection of HIV-1 RNA in human plasma specimens. The results attained from processing the NeuMoDx HIV-1 Calibrators are applied to the stored standard curve and used to generate a calibration coefficient, which is used to automatically adjust the standard curve for slight variations across systems or test strip lots. Use of both the standard curve and the system/lot specific calibration coefficient allows for the accurate quantitation of the HIV-1 RNA in the human clinical samples.

In addition, the traceability of these calibrators to the WHO 3<sup>rd</sup> HIV-1 International Standard enables laboratories to ensure that results obtained from the NeuMoDx HIV-1 Quant Assay are consistent across reagent lots, systems, and operators.

### PRINCIPLES OF THE PROCEDURE

The NeuMoDx HIV-1 Calibrators have been formulated to mimic natural human plasma specimens containing HIV-1 RNA. The encapsulated target material used in these calibrators allows for the verification of efficacious nucleic acid extraction and real-time PCR amplification and detection, enabling calibration of the entire testing process. One set of calibrators is processed every 90 days or with a change in NeuMoDx System, software, or NeuMoDx HIV-1 Quant Test Strip lot. The NeuMoDx System will automatically process each calibrator in triplicate. Such routine processing of the NeuMoDx HIV-1 Calibrators enables laboratories to ensure accuracy of test results for human clinical specimens processed within the validity period. These calibrators are processed in a manner identical to the processing of the human clinical specimens intended for quantitative HIV-1 testing.

Software on the NeuMoDx System automatically alerts the operator when a calibration is required. During processing, criteria for acceptance of the calibrator are automatically verified by the NeuMoDx System software. If fewer than two of the calibrator replicates is valid, the software automatically invalidates the run. Samples in an invalidated run must be retested using a new set of calibrators and controls.

Upon successful processing of the NeuMoDx HIV-1 Calibrators, the system software automatically records the validity of the processed calibrators for a period of 90 days, unless there is a change to the system that causes the validity period to expire. The NeuMoDx System software will automatically notify the user to process new calibrators when the previously processed calibrator validity period has expired and will not allow the processing of patient samples until a new period is established.

### REAGENTS & CONSUMABLES

#### Material Provided

REF	Contents	Tests per unit	Total tests per kit
800304	<b>NeuMoDx HIV-1 Calibrators</b> <i>Single use sets of HIV-1 High and Low Calibrators to establish validity of standard curve</i> <i>(1 vial of each level = 1 set)</i>	1 set	3

**Materials Required but Available Separately**



REF	Contents
300500	<b>NeuMoDx HIV-1 Quant Test Strip</b> <i>Dried PCR reagents containing HIV-1 specific TaqMan<sup>®</sup> probes and primers &amp; SPC2 specific TaqMan probe and primers</i>
100200	<b>NeuMoDx Extraction Plate</b> <i>Dried paramagnetic particles, lytic enzyme, and sample process controls</i>
900301	<b>NeuMoDx HIV-1 External Controls</b> <i>Single use sets of HIV-1 Positive and Negative External Controls to establish daily validity of NeuMoDx HIV-1 Quant Assay</i>
400600	<b>NeuMoDx Lysis Buffer 3</b>
400100	<b>NeuMoDx Wash Reagent</b>
400200	<b>NeuMoDx Release Reagent</b>
100100	<b>NeuMoDx Cartridge</b>
235903	<b>Hamilton CO-RE/CO-RE II Tips (300 µL) with Filters</b>
235905	<b>Hamilton CO-RE/CO-RE II Tips (1000 µL) with Filters</b>

**Instrumentation Required**

NeuMoDx 288 Molecular System [REF 500100] or NeuMoDx 96 Molecular System [REF 500200 or 500201]



**WARNINGS & PRECAUTIONS**

- The NeuMoDx HIV-1 Calibrators are for *in vitro* diagnostic use only with the NeuMoDx HIV-1 Quant Test Strip, as implemented on the NeuMoDx System.
  - Do not use NeuMoDx HIV-1 Calibrators after the listed expiration date.
  - Do not use the NeuMoDx HIV-1 Calibrators if the packaging is damaged or the contents are not frozen upon arrival.
-   
  


  - NeuMoDx HIV-1 Calibrators contain defibrinated human plasma that is negative for HBV DNA, HCV RNA, HIV-1 RNA, Human Parvovirus B19 DNA, and HAV RNA using nucleic acid amplification methods, and non-reactive for HBsAg and antibodies to HIV-1 and HIV-2, HCV, HTLV I and HTLV II, HBs, and HBe using FDA licensed test methods. This does not ensure the absence of these or other human pathogens. Follow universal precautions when handling.
  - Always handle specimens as if they are infectious and in accordance with safe laboratory procedures such as those described in Biosafety in Microbiological and Biomedical Laboratories<sup>1</sup> and in CLSI Document M29-A3.<sup>2</sup>
  - Do not pipette by mouth. Do not smoke, drink, or eat in areas where specimens or reagents are being handled.
  - Dispose of unused reagents and waste in accordance with country, federal, provincial, state and local regulations.
  - Clean, powder-free, nitrile gloves should be worn when handling all NeuMoDx reagents and consumables.
  - Wash hands thoroughly after performing the test.
  - Safety Data Sheets (SDS) are provided for each reagent (as applicable) at [www.qiagen.com/safety](http://www.qiagen.com/safety)
  - When working with chemicals, always wear a suitable lab coat, disposable gloves, and protective goggles. For more information, please consult the appropriate safety data sheets (SDS).

**Emergency information**

CHEMTREC

Outside USA & Canada +1 703-527-3887

**Disposal**

Dispose of as hazardous waste in compliance with local and national regulations. This also applies to unused products. Follow recommendations in the Safety Data Sheet (SDS).

**PRODUCT STORAGE, HANDLING & STABILITY**

- The NeuMoDx HIV-1 Calibrators are shipped with dry ice to maintain a frozen state; do not use if contents are not frozen upon receipt.
- It is recommended that the NeuMoDx HIV-1 Calibrators be stored at -15 °C to -20 °C to ensure stability.



- Calibrator vials are intended for single use only. Thawed calibrators may be stored at 4 °C for no longer than 24 hours.
- Refreezing after a first thaw is not recommended.
- Although the NeuMoDx HIV-1 Calibrators are non-infectious, any unused material should be discarded after use as biohazard waste to reduce risk of contamination by the target nucleic acid contained.
- Discard any calibrators that appear cloudy or contain large precipitates after thawing.

### INSTRUCTIONS FOR USE

1. NeuMoDx HIV-1 Calibrators must be processed under the following scenarios:
  - a. Validity of previously established calibration has expired (past 90 days)
  - b. Calibration validity has not been established on the NeuMoDx System(s)
  - c. Calibration validity has not been established for a given lot of NeuMoDx HIV-1 Quant Test Strips
  - d. The NeuMoDx System software has been modified
2. If a valid calibration does not exist, the NeuMoDx System will prompt the user to process calibrators (and external controls) before sample results can be reported.
3. If calibrators are required, process the NeuMoDx HIV-1 Calibrators (1 high calibrator and 1 low calibrator):

NeuMoDx HIV-1 Calibrator	Label Color Scheme
High Calibrator (HCHIV)	Green
Low Calibrator (LCHIV)	Blue

4. Retrieve a set of NeuMoDx HIV-1 Calibrators from freezer and place at room temperature (15-30 °C) until completely thawed.
5. Vortex gently to ensure homogeneity.
6. Load the calibrator vials into a standard 32-tube Specimen Tube Carrier, and ensure caps are removed from all tubes.
7. Place the Specimen Tube Carrier on the Autoloader shelf and use the touchscreen to load carrier onto the NeuMoDx System worktable.
8. The NeuMoDx System will recognize the barcode and start processing the specimen tubes unless reagents or consumables required for testing are not available.
9. To generate valid results, at least 2 out of the 3 replicates must yield results within predefined parameters. The low calibrator nominal target is 3.0 log<sub>10</sub> IU/mL and the high calibrator nominal target is 5.0 log<sub>10</sub> IU/mL.

NeuMoDx HIV-1 External Calibrator	HIV-1 Result
High Calibrator (HCHIV)	2/3 Calibrators Valid
Low Calibrator (LCHIV)	2/3 Calibrators Valid

10. Discrepant result handling for external calibrators should be performed as follows:
  - a. If one or both the calibrators fails the validity check, repeat processing of the failed calibrator(s) using new vial(s). In the event one calibrator fails validity, it is possible to only repeat the failed calibrator as the NeuMoDx System does not require the user to run both calibrators.
  - b. If problem persists, contact NeuMoDx Molecular, Inc.
11. External controls [REF 900301] must be processed *after* calibrator validity has been established and prior to obtaining test results from samples.

### LIMITATIONS

1. The NeuMoDx HIV-1 Calibrators can only be used in conjunction with the NeuMoDx HIV-1 Quant Test Strips on the NeuMoDx System.
2. A valid calibration of the NeuMoDx HIV-1 Quant Test Strip using the NeuMoDx HIV-1 Calibrators [REF 800304] is required *before* the NeuMoDx HIV-1 External Controls [REF 900301] can be processed.
3. Erroneous results could occur from improper handling, storage, or other technical error.
4. Operation of the NeuMoDx System is limited to use by personnel trained on the use of the NeuMoDx System.

### REFERENCES

1. Centers for Disease Control and Prevention Biosafety in Microbiological and Biomedical Laboratories, 6th edition. HHS Publication HHS Publication No. (CDC) 300859, Revised June 2020
2. Clinical and Laboratory Standards Institute (CLSI). Protection of Laboratory Workers from Occupationally Acquired Infections; Approved Guideline – Fourth Edition. CLSI document M29-A4; May 2014.

---

### TRADEMARKS


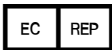








NeuMoDx™ is a trademark of NeuMoDx Molecular, Inc.

TaqMan® is a registered trademark of Roche Molecular Systems, Inc.

All other product names, trademarks, and registered trademarks that may appear in this document are property of their respective owners.

### SYMBOLS

The following symbols may appear in the instructions for use or on the packaging and labeling:

SYMBOL	MEANING
<b>R only</b>	Prescription use only
	Manufacturer
<b>IVD</b>	<i>In vitro</i> diagnostic medical device
	Authorized representative in the European Community
<b>REF</b>	Catalog number
<b>LOT</b>	Batch code
	Use-by date
	Temperature limit
	Humidity limitation
	Do not re-use
	Contains sufficient for <n> tests
	Consult instructions for use
	Caution
	Biological risks
<b>CE</b>	CE Mark



NeuMoDx Molecular, Inc.  
1250 Eisenhower Place  
Ann Arbor, MI 48108, USA



Emergo Europe B.V.  
Westervoortsedijk 60  
6827 AT Arnhem  
The Netherlands



Technical support/Vigilance reporting: [support.qiagen.com](http://support.qiagen.com)

Patent: [www.neumodx.com/patents](http://www.neumodx.com/patents)