

Certificate of Analysis for NR-59585

Genomic RNA from Influenza A Virus, A/Macha/O1453/2021 (H1N1)

Catalog No. NR-59585

Product Description:

Genomic RNA was isolated from a preparation of cell lysate and supernatant from Madin-Darby canine kidney SIAT-1 (MDCK-SIAT1) cells infected with influenza A virus, A/Macha/O1453/2021 (H1N1) using QIAamp® Viral RNA Mini Kit (Qiagen® 52906). The viral genomic RNA is in a background of cellular nucleic acid and carrier RNA.

Lot: 70063753 Manufacturing Date: 12JAN2024

TEST	SPECIFICATIONS	RESULTS
Genotypic Analysis Sequencing of species-specific region (~ 920 nucleotides)	Consistent with source virus	Consistent with source virus ¹
Functional Activity by RT-PCR Amplification ² Matrix gene	~ 1000 base pair amplicon	~ 1000 base pair amplicon
Estimated Concentration (post-dilution) by Qubit® Measurement (Viral, Cellular and Carrier) ³	Report results	2.1 ng per 100 µL (0.0207 µg/mL)
Estimated Amount per Vial ³	Report results	2.1 ng
Genome Copy Number Using BioRad QX200 Droplet Digital PCR (ddPCR™) System (Post vial; 12 replicates)	Report results	1.1 × 10 ⁸ NDU per mL ⁴
Virus Inactivation 10% of total yield inoculated on MDCK-SIAT1 cells and evaluated for cytopathic effect and HA after serial passage ⁵	No viable virus detected	No viable virus detected

¹Sequence information for influenza A virus A/Macha/O1453/2021 (H1N1) is not available in the NCBI database; nucleotide sequence obtained for NR-59585 lot 70063753 is identical to the source virus.

/Sonia Bjorum Brower/ Sonia Bjorum Brower

<u>11 APR 2024</u>

Technical Manager or designee, ATCC Federal Solutions

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²Amplified using iTaq[™] Universal SYBR Green One-step Kit (Bio-Rad[®] 172-5151) with 5 μL of NR-59585 in a 50 μL reaction

³Measurement is determined pre-vial prior to dilution due to the limit of detection of the quantification method

⁴NDU; NAAT-detectable units

⁵Use of the QIAamp® Viral RNA Mini Kit has been demonstrated to consistently inactivate 100% of influenza A viruses as shown by the absence of cytopathic effect (CPE) and HA after plating the entire extract on virus-susceptible cells for two passages.