SUPPORTING INFECTIOUS DISEASE RESEARCH

## Genomic RNA from Influenza B Virus, B/Baltimore/JH002/2021

### Catalog No. NR-59588

#### **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 B virus, B/Baltimore/JH002/2021 using QIAamp<sup>®</sup> Viral RNA Mini Kit (Qiagen<sup>®</sup> 52906). The viral genomic RNA is in a background of cellular nucleic acid and carrier RNA.

### Lot: 70063757

# Manufacturing Date: 01NOV2023

TEST	SPECIFICATIONS	RESULTS
Genotypic Analysis Sequencing of species-specific region Hemagglutinin gene (~ 710 nucleotides)	Consistent with source virus	Consistent with source virus <sup>1</sup>
Functional Activity by RT-PCR Amplification <sup>2</sup> Hemagglutinin gene	~ 1000 base pair amplicon	~ 1000 base pair amplicon
Estimated Concentration (post-dilution) by RiboGreen <sup>®</sup> Measurement (Viral, Cellular and Carrier) <sup>3</sup>	Report results	5.3 ng per 100 μL (0.0531 μg/mL)
Estimated Amount per Vial <sup>3</sup>	Report results	5.3 ng
Genome Copy Number Using BioRad QX200 Droplet Digital PCR (ddPCR™) System (Post vial; 12 replicates)	Report results	1.7 × 10 <sup>8</sup> NDU/mL <sup>4</sup>
Virus Inactivation 10% of total yield inoculated on MDCK-SIAT1 cells and evaluated for cytopathic effect and HA after serial passage <sup>5</sup>	No viable virus detected	No viable virus detected

<sup>1</sup>Sequence information for influenza B virus B/Baltimore/JH002/2021 is not available in the NCBI database; nucleotide sequence obtained for NR-59588 lot 70063757 is identical to the source virus.

<sup>2</sup>Amplified using iTaq™ Universal SYBR Green One-step Kit (Bio-Rad<sup>®</sup> 172-5151) with 5 μL of NR-59588 in a 50 μL reaction

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

<sup>4</sup>NDU; NAAT-detectable units

<sup>5</sup>Use of the QIAamp<sup>®</sup> 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.

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