

**Genomic RNA from Yellow Fever Virus, 17D**

**Catalog No. NR-2869**

**Product Description:** Genomic RNA was isolated from a preparation of cell lysate and supernatant from *Cercopithecus aethiops* kidney epithelial (Vero)<sup>1</sup> cells infected with yellow fever virus (YFV), 17D.

**Lot<sup>2</sup>: 64315467**

**Manufacturing Date: 17JUN2016**

TEST	SPECIFICATIONS	RESULTS
<b>Sequencing of Species-Specific Region</b> Polyprotein gene NS5 region (1093 nucleotides)	Consistent with YFV, 17D	99% identity with YFV, 17D (GenBank: X03700)
<b>Functional Activity by RT-PCR Amplification<sup>3</sup></b>	~ 1100 bp amplicon	~ 1100 bp amplicon (Figure 1)
<b>Total RNA Content by RiboGreen<sup>®</sup> Measurement (Viral, Cellular and Carrier)</b>	Report results	28.6 ng per 100 µL
<b>Virus Inactivation</b> 10% of total yield inoculated on Vero cells and evaluated for cytopathic effect <sup>4</sup>	No viable virus detected	No viable virus detected

<sup>1</sup>Vero cells: ATCC<sup>®</sup> CCL-81™

<sup>2</sup>Nucleic acid was extracted from a preparation of YFV, 17D (BEI Resources NR-116, Lot 7496109), using a QIAamp<sup>®</sup> Viral RNA Mini Kit (Qiagen 52906).

<sup>3</sup>Reverse transcription was performed using an iScript™ cDNA Synthesis Kit (Bio-Rad 1708891) with 10 µL of NR-2869 in a 20 µL reaction; PCR was performed using iTaq™ DNA Polymerase (Bio-Rad 1708870) with 10 µL of cDNA in a 50 µL reaction.

<sup>4</sup>Use of the QIAamp<sup>®</sup> Viral RNA Mini Kit has been demonstrated to consistently inactivate 100% of yellow fever virus as shown by the absence of cytopathic effect (CPE) after plating the entire extract on virus-susceptible cells.

**Date:** 29 NOV 2016

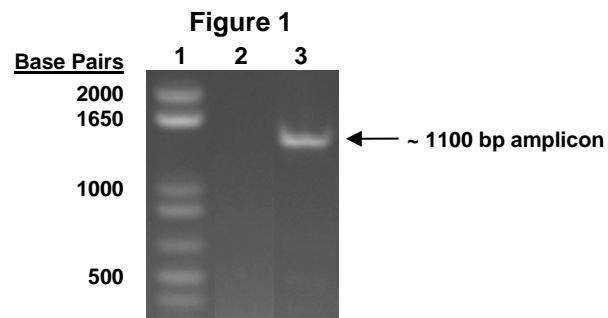
**Signature:**   
BEI Resources Authentication

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Lane 1: Invitrogen™ TrackIt™ 1 Kb Plus DNA Ladder  
Lane 2: Negative control  
Lane 3: NR-2869