

## Certificate of Analysis for NR-46501

## Naegleria fowleri, Strain CDC:V615

## Catalog No. NR-46501

This reagent is the tangible property of the U.S. Government.

**Product Description:** *Naegleria fowleri (N. fowleri)*, strain CDC:V615 is a clinical isolate collected in 2009 from the cerebral spinal fluid of a 22-year-old male the United States.

Lot<sup>1,2</sup>: 64357374 Manufacturing Date: 14JUN2016

TEST	SPECIFICATIONS	RESULTS
Genotyping Sequencing of Internal Transcribed Spacer 1 (ITS 1) and 5.8S ribosomal RNA gene (~ 540 base pairs)	Consistent with N. fowleri	Consistent with <i>N. fowleri</i> , genotype I <sup>3</sup>
Functional Activity by PCR Amplification <sup>4</sup> ITS 1, 5.8S ribosomal RNA gene	~ 600 base pair amplicon	~ 600 base pair amplicon
Viable Cell Count by Hemacytometry (pre-freeze)	> 10 <sup>6</sup> cells/mL	7 × 10 <sup>6</sup> cells/mL
Viability (post-freeze) <sup>5</sup>	Growth	Growth
Sterility (21-day incubation)		
Harpo's HTYE broth <sup>6</sup> , 37°C and 26°C, aerobic	No growth	No growth
Trypticase soy broth, 37°C and 26°C, aerobic	No growth	No growth
Sabouraud broth, 37°C and 26°C, aerobic	No growth	No growth
DMEM with 10% FBS, 37°C, aerobic	No growth	No growth
Sheep blood agar, 37°C, aerobic	No growth	No growth
Sheep blood agar, 37°C, anaerobic	No growth	No growth
Thioglycollate broth, 37°C, anaerobic	No growth	No growth

<sup>&</sup>lt;sup>1</sup>NR-46501 was produced by cultivation of the deposited material in modified PYNFH medium (ATCC<sup>®</sup> medium 1034) supplemented with 10% heat-inactivated fetal bovine serum for 4 days at 35°C in an aerobic atmosphere until peak density was reached.

Date: 03 FEB 2017 Signature:

**BEI Resources Authentication** 

ATCC®, on behalf of BEI Resources, hereby represents and warrants that the material provided under this certificate has been subjected to the tests and procedures specified and that the results described, along with any other data provided in this certificate, are true and accurate to the best of ATCC®'s knowledge.

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<sup>&</sup>lt;sup>2</sup>Quality control testing completed on post-freeze material unless specified as pre-freeze.

<sup>&</sup>lt;sup>3</sup>For genotyping details refer to Zhou, L., et al. "Genetic Variations in the Internal Transcribed Spacer and Mitochondrial Small Subunit rRNA Gene of *Naegleria* Spp." J. Eukaryot. Microbiol. 50 (2003): 522-526. PubMed: 14736150.

<sup>&</sup>lt;sup>4</sup>PCR amplification was performed using the NF-ITS-F1 and NT-ITS-F2 primer set as described in Zhou, L., et al. "Genetic Variations in the Internal Transcribed Spacer and Mitochondrial Small Subunit rRNA Gene of *Naegleria* Spp." <u>J. Eukaryot. Microbiol.</u> 50 (2003): 522-526. PubMed: 14736150.

<sup>&</sup>lt;sup>5</sup>Viable cells were observed after 1 day at 25°C in an aerobic atmosphere in Modified M199 medium supplemented with 10% heat-inactivated fetal bovine serum and 10 μg/mL hemin.

<sup>&</sup>lt;sup>6</sup>Atlas, Ronald M. Handbook of Microbiological Media. 3rd ed. Ed. Lawrence C. Parks. Boca Raton: CRC Press, 2004, p. 798.