

## **Certificate of Analysis for NR-33655**

## Acanthamoeba sp., Strain CDC:V025

Catalog No. NR-33655

**Product Description:** Acanthamoeba sp., strain CDC:V025 was isolated from the contact lens of a man in Louisiana.

Lot<sup>1</sup>: 60732092 Manufacturing Date: 16FEB2012

TEST	SPECIFICATIONS	RESULTS
Genotyping Sequencing of 18S ribosomal RNA gene (~ 440 bp)	Consistent with Acanthamoeba sp.	Consistent with <i>Acanthamoeba</i> sp.
Functional Activity by PCR Amplification <sup>2</sup> 18S ribosomal RNA gene (amplicon ASA.S1)	423 bp to 551 bp amplicon	~ 450 bp amplicon
Viable Cell Count by Hemacytometry (pre-freeze)	> 10 <sup>6</sup> cells/mL	6.4 x 10 <sup>6</sup> cells/mL
Viability (post-freeze) <sup>3</sup>	Growth	Growth
Sterility (21-day incubation) Harpo's HTYE broth <sup>4</sup> , 37°C and 26°C, aerobic Trypticase soy broth, 37°C and 26°C, aerobic Sabouraud broth, 37°C and 26°C, aerobic Brain heart infusion, 37°C and 26°C, aerobic Sheep blood agar, 37°C, aerobic Sheep blood agar, 37°C, anaerobic Thioglycollate broth, 37°C, anaerobic	No growth	No growth

NR-33655 was produced by cultivation of *Acanthamoeba* sp., strain CDC:V025 in PYG medium (<u>ATCC medium 712</u>) for 6 days at 25°C in an aerobic atmosphere and preserved.

**Date:** 10 AUG 2012 **Signature:** 

**Title:** Technical Manager, BEI Authentication or designee

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<sup>&</sup>lt;sup>2</sup>PCR amplification was performed using the JDP1 and JDP2 primer set as described [Schroeder, J. M. et al. "Use of Subgenic 18S Ribosomal DNA PCR and Sequencing for Genus and Genotype Identification of Acanthamoebae from Humans with Keratitis and from Sewage Sludge." <u>J. Clin. Microbiol.</u> 39 (2001): 1903-1911. PubMed: 11326011].

<sup>&</sup>lt;sup>3</sup>Viable cells were observed after 6 days under cultivation conditions.

<sup>&</sup>lt;sup>4</sup>Atlas, Ronald M. Handbook of Microbiological Media. 3rd ed. Ed. Lawrence C. Parks. Boca Raton: CRC Press, 2004, p. 798.