

Certificate of Analysis for NR-46470

Acanthamoeba sp., Strain CDC:V411

Catalog No. NR-46470

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

Product Description: *Acanthamoeba* sp., strain CDC:V411 is a clinical isolate collected in 1998 from the brain tissue of a male patient in Spain.

Lot^{1,2}: 64463925 Manufacturing Date: 05AUG2016

TEST	SPECIFICATIONS	RESULTS
Cellular Morphology	Report results	Adherent and non-adherent
Genotyping Sequencing of 18S ribosomal RNA gene (~ 350 base pairs)	Consistent with Acanthamoeba sp.	Consistent with <i>Acanthamoeba</i> sp. (genotype T4)
Functional Activity by PCR Amplification ³ 18S ribosomal RNA gene (amplicon ASA.S1)	423 to 551 base pair amplicon	~ 450 base pair amplicon
Viable Cell Count by Hemocytometry (pre-freeze)	> 10 ⁶ cells/mL	3.9 × 10 ⁶ cells/mL
Viability ⁴	Growth	Growth
Sterility (21-day incubation) Harpo's HTYE broth ⁵ , 37°C and 26°C, aerobic Tryptic Soy broth, 37°C and 26°C, aerobic Sabouraud Dextrose broth, 37°C and 26°C, aerobic DMEM with 10% FBS, 37°C, aerobic Sheep Blood agar, 37°C, aerobic Sheep Blood agar, 37°C, anaerobic Thioglycollate broth, 37°C, anaerobic	No growth	No growth

¹NR-46470 was produced by cultivation of the deposited material in Peptone Yeast Glucose medium (PYG; ATCC® medium 712) for 7 days at 25°C in an aerobic atmosphere to produce this lot.

Date: 06 MAR 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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²Quality control testing completed on post-freeze material unless specified as pre-freeze.

³PCR amplification was performed using the JDP1 and JDP2 primer set (JDP1: 5'-GGCCCAGATCGTTTACCGTGAA-3' and JDP2: 5'-TCTCACAAGCTGCTAGGGAGTCA-3') as described in 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." J. Clin. Microbiol. 39 (2001): 1903-1911. PubMed: 11326011.

⁴Viable cells were observed after 1 day at 25°C in an aerobic atmosphere in PYG medium.

⁵Atlas, Ronald M. <u>Handbook of Microbiological Media</u>. 3rd ed. Ed. Lawrence C. Parks. Boca Raton: CRC Press, 2004, p. 798.