

Certificate of Analysis for NR-46475

Acanthamoeba sp., Strain CDC:V540

Catalog No. NR-46475

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

Product Description: *Acanthamoeba* sp., strain CDC:V540 is a clinical isolate collected in 2003 from the frozen brain tissue of a 60-year-old female patient in Massachusetts, USA.

Lot^{1,2}: 2186 Manufacturing Date: 04NOV2016

TEST	SPECIFICATIONS	RESULTS
Cellular Morphology	Report results	Adherent and non-adherent, refractile
Genotyping Sequencing of 18S ribosomal RNA gene (~ 430 base pairs)	Consistent with Acanthamoeba sp.	Consistent with <i>Acanthamoeba</i> sp. (genotype T1)
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	6.7 × 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-46475 was produced by cultivation of the deposited material in Peptone Yeast Glucose (PYG) medium (ATCC® medium 712) for 8 days at 30°C in an aerobic atmosphere to produce this lot.

Date: 28 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.

ATCC® is a trademark of the American Type Culture Collection.

You are authorized to use this product for research use only. It is not intended for human use.

BEI Resources

www.beiresources.org

E-mail: contact@beiresources.org

Tel: 800-359-7370 Fax: 703-365-2898

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'-GGCCCAGATCGTTACCGTGAA-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 30°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.