b|e|i RESOURCES

SUPPORTING INFECTIOUS DISEASE RESEARCH

Certificate of Analysis for NR-10272

Toxoplasma gondii, GT1-FUDR3.3

Catalog No. NR-10272

Product Description: *Toxoplasma gondii* (*T. gondii*), GT1-FUDR3.3 is a virulent Type I parental strain that was used in a genetic cross with the nonvirulent genotype Type III parental strain CTG.11 ARA-SNF (also referred to as CTG.11 ARA-SYN, CTG.11 ARA-A^R/SNF^R and CEP.11 ARA-A^R/SNF^R).

Lot[1,2]: 59534843 **Manufacturing Date: 28OCT2010**

TEST	SPECIFICATIONS	RESULTS
Genotyping[2-4]		
AK16 locus (*Hinf*I digestion)	Consistent with parental Type I strain	Consistent with parental Type I strain
L358 locus (*Hae*III digestion)	Consistent with parental Type I strain	Consistent with parental Type I strain
Drug Susceptibility[2,5]		
Sinefungin (SNF)	Sensitive	Sensitive
Adenine arabinose (Ara-A)	Sensitive	Sensitive
Viable Cell Count by Hemacytometry (pre-freeze)	> 10^6 cells/mL	4.8 x 10^7 cells/mL
Viability (post-freeze)[6]	Growth	Growth
Sterility (21-day incubation)		
Harpo's HTYE broth[7], 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
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
DMEM with 10% FBS, 37°C and 5% CO_2	No growth	No growth
Mycoplasma Contamination		
DNA Detection by PCR	None detected	None detected

[1]NR-10272 lot 59534843 was produced by cultivation of NR-10272 lot 28792126 in human foreskin fibroblast cells (ATCC® CRL-1634™) with cell cultivation medium for parasites (ATCC medium 2222: adjusted to contain 10% heat-inactivated fetal bovine serum). The culture was propagated in 95% air, 5% CO_2 for 6 days at 37°C, until lysis of the host cell monolayer was reached.

[2]The Genotyping and Drug Susceptibility tests were performed on the source culture (NR-10272 lot 28792126) only and were not repeated for NR-10272 lot 59534843.

[3]PCR amplification was performed separately for the two loci AK16 and L358. Where appropriate, samples were subjected to restriction enzyme digestion typing by agarose gel electrophoresis.

[4]Primer sequences, annealing temperatures, and conditions for restriction enzyme digestion may be obtained at the *Toxoplasma* Genome Map website (Toxoplasma Genome Map).

[5]Sinefungin was used at a concentration of 2.7 x 10^{-7} M and ara-A was used at a concentration of 1.3 x 10^{-4} M, as described (Sibley, L. D., et al. "Generation of a Restriction Fragment Length Polymorphism Linkage Map for *Toxoplasma gondii*." Genetics 132 (1992): 1003-1015. PubMed: 1360931).

[6]Viable cells and signs of infection were seen after 6 days under cultivation conditions at 37°C.

[7]Atlas, Ronald M. Handbook of Microbiological Media. 3rd ed. Ed. Lawrence C. Parks. Boca Raton: CRC Press, 2004, p. 798.

Date: 12 APR 2011 **Signature:**

Title: Technical Manager, BEI Authentication or designee

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