

Certificate of Analysis for NR-10159

Toxoplasma gondii, Clone CL19

Catalog No. NR-10159

Product Description: *Toxoplasma gondii*, clone CL19 is a recombinant F1 clone selected from progeny of two parallel genetic crosses between a Type II parental strain [ME49 (clone B7)] and a Type III parental strain (CTG ARA-SYN).

Lot¹: 58270251 Manufacturing Date: 30JUL2008

TEST	SPECIFICATIONS	RESULTS
Genotyping ²		
850 locus (Sfa NI digestion) ³	Consistent with parental Type III strain	Consistent with parental Type III strain
SAG1 locus ⁴	Consistent with parental Type III strain	Consistent with parental Type III strain
Drug susceptibility ⁵		
Sinefungin	Resistant	Resistant
Ara-A	Susceptible	Susceptible
Viable Cell Count by Hemacytometry (pre-freeze)	> 10 ⁶ cells/mL	3.6 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

¹NR-10159 was produced by cultivation of the deposited material 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% CO₂ for 19 days at 37°C, until lysis of the host cell monolayer was reached.

Date: 15 OCT 2009 **Signature:** Signature on File

Title: Technical Manager, BEI Authentication or designee

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²PCR amplification was performed separately for the two loci 850 and SAG1. Where appropriate, samples were subjected to restriction enzyme digestion typing by agarose gel electrophoresis.

³Primer sequences, annealing temperatures, and conditions for restriction enzyme digestion may be obtained at the *Toxoplasma* Genome Map website (http://toxomap.wustl.edu/Toxo Genetic Map Table.html). ⁴Primer sequences and conditions for PCR are available upon request.

⁵Sinefungin 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.*" <u>Genetics</u> 132 (1992): 1003-1015. PubMed: 1360931.)

⁶Incubated under cultivation conditions for 15 days at 37°C.

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