

Certificate of Analysis for NR-10260

Toxoplasma gondii, Clone c285-13

Catalog No. NR-10260

Product Description: *Toxoplasma gondii*, c285-13 is a virulent recombinant F1 clone selected from progeny of two parallel genetic crosses between a highly virulent Type I parental strain, GT1-FUDR3.3 and a less virulent Type III parental strain, CTG.11 ARA-SNF.

Lot¹: 59095937 Manufacturing Date: 16FEB2010

TEST	SPECIFICATIONS	RESULTS
Genotyping ^{2,3}		
AK16 locus (<i>Hinf</i> l digestion) L358 locus (<i>Hae</i> III digestion)	Consistent with parental Type III strain Consistent with parental Type III strain	Consistent with parental Type III strain Consistent with parental Type III strain
Drug susceptibility ⁴ Sinefungin (SNF) Adenine arabinose (Ara-A)	Sensitive Sensitive	Sensitive Sensitive
Viable Cell Count by Hemacytometry (pre-freeze)	> 10 ⁶ cells/mL	2.8 x 10 ⁷ cells/mL
Viability (post-freeze) ⁵	Growth	Growth
Sterility (21-day incubation) Harpo's HTYE broth ⁶ , 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
Mycoplasma Contamination DNA Detection by PCR	None detected	None detected

¹NR-10260 was produced by cultivation of the deposited material in human foreskin fibroblast cells (ATCC[®] CRL-1634™) with cell cultivation medium for parasites (<u>ATCC medium 2222:</u> adjusted to contain 10% heat-inactivated fetal bovine serum). The culture was propagated in 95% air, 5% CO₂ for 4 days at 37°C, until lysis of the host cell monolayer was reached.

Date: 24 JUN 2010 **Signature:** Signature on File

Title: Technical Manager, BEI Authentication or designee

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²PCR 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.

³Primer sequences, annealing temperatures, and conditions for restriction enzyme digestion may be obtained at the *Toxoplasma* Genome Map website (<u>Toxoplasma Genome Map</u>).

⁴Sinefungin 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).

⁵Viable cells and signs of infection were seen after 6 days under cultivation conditions at 37°C.

⁶Atlas, Ronald M. Handbook of Microbiological Media. 3rd ed. Ed. Lawrence C. Parks. Boca Raton: CRC Press, 2004, p. 798.