SUPPORTING INFECTIOUS DISEASE RESEARCH

### Toxoplasma gondii, Strain SF8

#### Catalog No. NR-49181

**Product Description:** *Toxoplasma gondii* (*T. gondii*), strain SF8 is a recombinant F1 clone selected from progeny of a genetic cross between a sinfungin-resistant line of the highly virulent Type I GT-1 strain (GT1-SNF<sup>R</sup>) and a 5-fluoro-2'-deoxyuridine-resistant line of the non-virulent Type 2 ME49 strain (ME49 FUDR<sup>R</sup>).

# Lot<sup>1,2</sup>: 63626656

## Manufacturing Date: 27JUL2015

TEST	SPECIFICATIONS	RESULTS		
Cell Morphology	Report results	Refractile and crescent shaped		
PCR Assay of Extracted DNA <sup>3,4</sup> AK56 locus	~ 520 base pair amplicon	~ 520 base pair amplicon		
<b>Genotypic Analysis</b> <sup>3,4</sup> Sequencing of AK56 locus (~ 480 base pairs) AK56 locus ( <i>Mfe</i> l digestion)	Consistent with <i>T. gondii</i> Consistent with parental Type I strain	Consistent with <i>T. gondii</i> (Figure 1) Consistent with parental Type I strain		
Viable Cell Count by Hemacytometry (pre-freeze)	> 10 <sup>6</sup> cells/mL	$3.7 \times 10^7$ cells/mL		
Viability (post-freeze) <sup>5</sup>	Viable parasites	Viable parasites		
Sterility (21-day incubation) Harpo's HTYE broth <sup>6</sup> , 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 No growth No growth No growth No growth No growth	No growth No growth No growth No growth No growth No growth No growth		
Mycoplasma Contamination DNA Detection by PCR	None detected	None detected		

<sup>1</sup>NR-49181 was produced by cultivation of the deposited material in human foreskin fibroblast cells (ATCC<sup>®</sup> CRL-1634<sup>™</sup>) with cell cultivation medium for parasites (ATCC<sup>®</sup> medium 2222: adjusted to contain 10% heat-inactivated fetal bovine serum). The culture was propagated 7 days at 37°C in an aerobic atmosphere with 5% CO<sub>2</sub> until lysis of the host cell monolayer was reached.

<sup>2</sup>Quality control testing completed on post-freeze material unless specified as pre-freeze.

<sup>3</sup>PCR amplification of the AK56 locus was performed. Samples were subjected to restriction enzyme digestion typing by agarose gel electrophoresis. <sup>4</sup>Primer sequences, annealing temperatures, and conditions for restriction enzyme digestion may be obtained at the *Toxoplasma* Genome Map website (<u>Toxoplasma Genome Map</u>).

<sup>5</sup>Viable cells and signs of infection were seen after 9 days under cultivation conditions at 37°C.

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

#### Figure 1: AK56 (Chromosome II) Amplicon Sequence

TATTAGGTTT	TTCCGTGTTT	TCGCGGAGTC	GTCTGAGCTC	GGCACTCGCT	GCTTTCCAAA	ATCTCGTTTC	AACGTATCGC
GGCGCCGTCA	CCGCGCGCAA	TCCACTGTGA	TGCATGATTC	TGTTTCTAAA	AACTGCGCAT	TTTAGCCGGC	TCGTTTTTGC
ATACGTTTGG	ACCATAAAAC	CTCGTATTGT	TGAAGAAGAA	TGCAATTGGT	GTCTGTGCTG	ATCACCGTAT	GAAAATCGGC
GTGTCTCGCC	CCCTGCCGTG	TGCGCGTCCG	CTTTTTGCGA	CCCCGGTACA	CCCGTTTTTT	GTGGTCAGCG	AGGAACGCAC
TTTTGCTGTT	ATTGTTCACT	TTTCAGCGTA	ACACTGACCC	CTTTCATCGT	GGCAGGAAAC	GAACTCTCAG	CAAGAATTTT
CGAGCACTAC	TGCGTCGCAG	CAGCCTAGTG	GGGTGGACAC	GCATGTGCAG	GACGGACAGA	AACTGCAAGC	TTGTTCCGCA

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# **Certificate of Analysis for NR-49181**

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Date: 26 JAN 2016

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