SUPPORTING INFECTIOUS DISEASE RESEARCH

Toxoplasma gondii, Strain C7B1

Catalog No. NR-49175

Product Description: *Toxoplasma gondii* (*T. gondii*), strain C7B1 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^R) and a 5-fluoro-2'-deoxyuridine-resistant line of the non-virulent Type 2 ME49 strain (ME49 FUDR^R).

Lot^{1,2}: 63500702

Manufacturing Date: 22MAY2015

TEST	SPECIFICATIONS	RESULTS
Cell Morphology	Report results	Refractile and crescent shaped
PCR Assay of Extracted DNA ^{3,4} AK56 locus	~ 520 base pair amplicon	~ 520 base pair amplicon
Genotypic Analysis ^{3,4} Sequencing of AK56 locus (~ 500 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 ⁶ cells/mL	2.6×10^7 cells/mL
Viability (post-freeze) ⁵	Viable parasites	Viable parasites (Figure 2)
Sterility (21-day incubation) Harpo's HTYE broth ⁶ , 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

¹Quality control testing completed on post-freeze material unless specified as pre-freeze.

²NR-49175 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 3 days at 37°C in an aerobic atmosphere with 5% CO₂ until lysis of the host cell monolayer was reached.

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

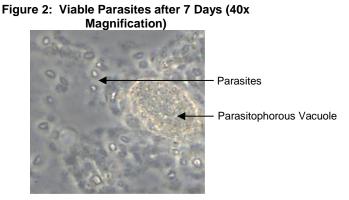
⁵Viable cells and signs of infection were seen after 1 day at 37°C in an aerobic atmosphere with 5% CO₂.

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

Figure 1: AK56 Locus (Chromosome II) Amplicon Sequence

CACTGCTTTT ATTAGGTTTT TCCGTGTTTT CGCGGAGTCG TCTGAGCTCG GCACTCGCTG CTTTCCAAAA TCTCGTTTCA ACGTATCGCG GCGCCGTCAC CGCGCGCAAT CCACTGTGAT GCATGATTCT GTTTCTAAAA ACTGCGCATT TTAGCCGGCT CGTTTTTGCA TACGTTTGGA CCATAAAACC TCGTATTGTT GAAGAAGAAT GCAATTGGTG TCTGTGCTGA TCACCGTATG AAAATCGGCG TGTCTCGCCC CCTGCCGTGT GCGCGTCCGC TTTTTGCGAC CCCGGTACAC CCGTTTTTTG TGGTCAGCGA GGAACGCACT TTTGCTGTTA TTGTTCACTT TTCAGCGTAA CACTGACCCC TTTCATCGTG GCAGGAAACG AACTCTCAGC AAGAATTTC GAGCACTACT GCGTCGCAGC AGCCTAGTGG GGTGGACACG CATGTGCAGG ACGGACAGAA ACTGCAAGCT TGTTCCGCAG GCTAAAACTC G **d**|**e**|**i** resources

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Date: 08 JAN 2016

Signature: (

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