

Certificate of Analysis for NR-49323

Clostridium difficile, Isolate 20110740

Catalog No. NR-49323

Product Description: Clostridium difficile (C. difficile; also referred to as Peptoclostridium difficile), isolate 20110740 was obtained from the stool of an older adult female patient with a community-associated (CA) C. difficile infection in northestern USA, in 2011. Isolate 20110740 was deposited as PCR ribotype 018, North American pulsed-field gel electrophoresis type 12 (NAP12), containing tcdA, tcdB and tcdC of the PaLoc operon. This isolate is reported to be negative for the C. difficile binary toxin (CDT).²

Lot¹: 63719914 Manufacturing Date: 30SEP2015

TEST	SPECIFICATIONS	RESULTS
Phenotypic Analysis		
Cellular morphology	Gram-positive rods	Gram-positive rods
Colony morphology ²	Report results	Irregular, flat, undulate, rough and gray (Figure 1)
Hemolysis ²	Non-hemolytic	Non-hemolytic
Motility (wet mount)	Report results	Motile
Biochemical tests:		
Esculin hydrolysis ³	Positive	Positive
Gelatin hydrolysis ³	Positive	Positive
VITEK® MS (MALDI-TOF)	Consistent with C. difficile	Consistent with <i>C. difficile</i> (99.9%)
Genotypic Analysis Sequencing of 16S ribosomal RNA gene (~ 1360 base pairs)	≥ 99% sequence identity to C. difficile type strain	99.1% sequence identity to AB075770 (ATCC [®] 9689™)
PCR Assay of Extracted DNA Presence of <i>C. difficile</i> -specific genes ⁴		
Triose phosphate isomerase (<i>tpi</i>) Presence of toxin genes ^{4,5}	~ 230 base pair amplicon	~ 230 base pair amplicon
cdtB	No amplicon	No amplicon
tcdA (wild type)	~ 370 base pair amplicon	~ 370 base pair amplicon
tcdA (partial deletion)	No amplicon	No amplicon
tcdB	~ 160 base pair amplicon	~ 160 base pair amplicon
Purity (post-freeze)		
Anaerobic growth ⁶	Growth consistent with expected morphology	Growth consistent with expected morphology
Aerobic growth ⁷	No growth	No growth
Viability (post-freeze) ²	Growth	Growth

NR-49323 was produced by inoculation of the deposited material into Modified Reinforced Clostridial medium and incubated for 1 day at 37°C in an anaerobic atmosphere (< 0.5% O₂; Remel™ Anaero Pack-Anaero™ R681001). The material from the initial growth was passaged in Modified Reinforced Clostridial medium for 1 day at 37°C in an anaerobic atmosphere to produce this lot.

BEI Resources

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²2 days at 37°C in an anaerobic atmosphere on Tryptic Soy agar with 5% defibrinated sheep blood

³Tests were assessed after 7 days at 37°C in an anaerobic atmosphere. The gelatin tube was placed at 4°C for one hour prior to result determination.

⁴Lemee, L., et al. "Multiplex PCR Targeting *tpi* (Triose Phosphate Isomerase), *tcdA* (Toxin A), and *tcdB* (Toxin B) Genes for Toxigenic Culture of *Clostridium difficile*." J. Clin. Microbiol. 42 (2004): 5710-5714. PubMed: 15583303.

⁵Antikainen, J., et al. "Detection of Virulence Genes of Clostridium difficile by Multiplex PCR." APMIS. 117 (2009): 607-613. PubMed: 19664132.

⁶Purity of this lot was assessed for 7 days at 37°C in an anaerobic atmosphere on Tryptic Soy agar with 5% defibrinated sheep blood.

⁷Purity of this lot was assessed for 7 days at 37°C in an aerobic atmosphere with 5% CO₂ on Tryptic Soy agar with 5% defibrinated sheep blood.



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Figure 1: Colony Morphology



Date: 22 FEB 2016

Signature:

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