

Certificate of Analysis for NR-49297

Clostridium difficile, Isolate 20120613

Catalog No. NR-49297

Product Description: Clostridium difficile (C. difficile; also referred to as Peptoclostridium difficile), isolate 20120613 was obtained from the stool of an elderly male patient with a healthcare-associated (HA) C. difficile infection in southern USA in 2011. Isolate 20120613 was deposited as PCR ribotype 014, North American pulsed-field gel electrophoresis type 4 (NAP4), containing tcdA and tcdB of the PaLoc operon. This isolate is reported to be negative for the C. difficile binary toxin (CDT).

Lot¹: 63950685 Manufacturing Date: 08APR2016

TEST	SPECIFICATIONS	RESULTS
Phenotypic Analysis		
Cellular morphology	Gram-positive rods	Gram-positive rods
Colony morphology ²	Report results	Irregular, flat, lobate, translucent and gray (Figure 1)
Hemolysis ²	Report results	Non-hemolytic
Motility (wet mount)	Report results	Motile
Biochemical tests:	·	
Esculin hydrolysis ³	Positive	Positive
Gelatin hydrolysis ⁴	Positive	Positive
VITEK® MS (MALDI-TOF)	C. difficile	C. difficile (99.9%) ⁵
Genotypic Analysis		
Sequencing of 16S ribosomal RNA gene (~ 1440 base pairs)	≥ 99% sequence identity to C. difficile type strain (GenBank: CP011968.1)	99.9% sequence identity to C. difficile type strain (GenBank: CP011968.1)
PCR Assay of Extracted DNA		
Presence of C. difficile-specific genes ⁶		
Triose phosphate isomerase (tpi)	~ 230 base pair amplicon	~ 230 base pair amplicon
Presence of toxin genes ^{6,7}	' '	·
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 ⁸	Growth consistent with expected morphology	Growth consistent with expected morphology
Aerobic growth ⁹	No growth	No growth
Viability (post-freeze) ²	Growth	Growth

The deposited material was inoculated into Modified Reinforced Clostridial medium, which was passaged three times at 37°C in an anaerobic atmosphere (< 0.5% O₂; Remel™ Anaero Pack-Anaero™ R681001) and preserved in 10% glycerol. NR-49297 was produced by inoculation of the preserved material into Modified Reinforced Clostridial medium and incubated for 2 days at 37°C in an anaerobic atmosphere. The material from the initial growth was passaged once in Modified Reinforced Clostridial medium for 1 day at 37°C in an anaerobic atmosphere to produce this lot.

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

³Esculin hydrolysis was assessed after 7 days at 37°C in an anaerobic atmosphere.

⁴Gelatin hydrolysis was assessed after 2 days at 37°C in an anaerobic atmosphere. The gelatin tube was placed at 4°C for one hour prior to result determination.

⁵Percent probabilities above 90% indicate a close match to the typical biochemical pattern for the given organism, with a percent probability of 99% being a perfect match between the test reaction pattern and the unique biochemical pattern of the given organism or organism group. For additional information, please refer to O'Hara, C.M. and J. M. Miller. "Evaluation of the VITEK 2 ID-GNB Assay for Identification of Members of the Family Enterobacteriaceae and Other Nonenteric Gram-Negative Bacilli and Comparison with the VITEK GNI+ Card." J. Clin. Microbiol. 41 (2003): 2096-2101. PubMed: 12734254.

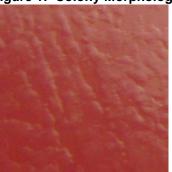
⁶Lemee, L., et al. "Multiplex PCR Targeting *tpi* (Triose Phosphate Isomerase), *tcdA* (Toxin A), and *tcdB* (Toxin B) Genes for Toxigenic Culture of *Clostridium difficile*." J. Clin. Microbiol. 42 (2004): 5710-5714. PubMed: 15583303.



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⁷Antikainen, J., et al. "Detection of Virulence Genes of *Clostridium difficile* by Multiplex PCR." <u>APMIS.</u> 117 (2009): 607-613. PubMed: 19664132. ⁸Purity of this lot was assessed for 3 days at 37°C in an anaerobic atmosphere on Tryptic Soy agar with 5% defibrinated sheep blood. ⁹Purity 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.

Figure 1: Colony Morphology



Date: 03 NOV 2016

Signature:

BEI Resources Authentication

ATCC®, on behalf of BEI Resources, hereby represents and warrants that the material provided under this certificate has been subjected to the tests and procedures specified and that the results described, along with any other data provided in this certificate, are true and accurate to the best of ATCC®'s knowledge.

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