

**Genomic DNA from *Clostridium botulinum*, Strain Walls 8G (VPI 4404)**

**Catalog No. NR-2713**

**Product Description:** Genomic DNA was isolated from a preparation of *Clostridium botulinum* (*C. botulinum*), strain Walls 8G (VPI 4404, BL 4851), serotype F (proteolytic).

**Lot<sup>1</sup>: 63360140**

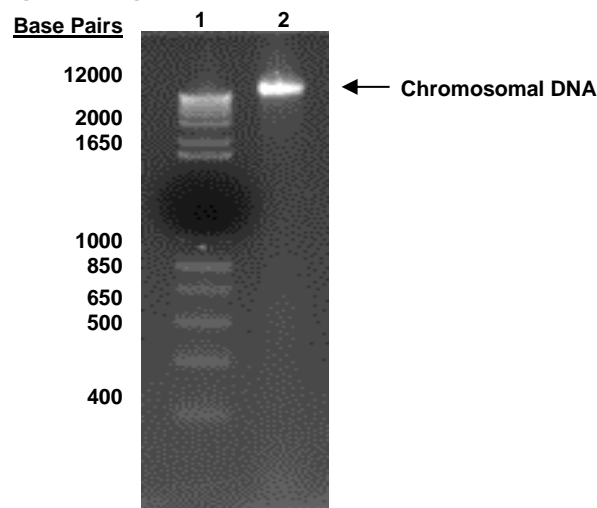
**Manufacturing Date: 15JUL2015**

TEST	SPECIFICATIONS	RESULTS
<b>Genotypic Analysis</b> Sequencing of 16S ribosomal RNA gene (~ 1290 base pairs)	≥ 99% sequence identity to <i>C. botulinum</i> , strain Walls 8G (GenBank: LBFH01000001)	99.8% sequence identity to <i>C. botulinum</i> , strain Walls 8G (GenBank: LBFH01000001)
<b>Agarose Gel Electrophoresis</b>	High molecular weight chromosomal DNA	High molecular weight chromosomal DNA (Figure 1)
<b>Concentration by PicoGreen<sup>®</sup> Measurement</b>	0.7 to 1.5 µg in 25 to 100 µL	1.2 µg in 81 µL per vial (15 µg/mL)
<b>Functional Activity by PCR Amplification</b> 16S ribosomal RNA gene	~ 1500 base pair amplicon	~ 1500 base pair amplicon
<b>OD<sub>260</sub>/OD<sub>280</sub> Ratio</b>	1.7 to 2.1	2.0
<b>Bacterial Inactivation</b> 10% of total yield plated on agar for 7 days <sup>2</sup>	No viable bacteria detected	No viable bacteria detected

<sup>1</sup>The bacterial preparation used for extraction of genomic DNA was produced from BEI Resources NR-272 lot 7364651. Genomic DNA was extracted using proprietary technology, and is provided in TE buffer (10 mM Tris-HCl, 1 mM EDTA, pH ~ 8.0).

<sup>2</sup>An extraction procedure was used that has been shown to consistently inactivate 100% of Gram-positive and Gram-negative bacteria.

**Figure 1: Agarose Gel Electrophoresis**



Lane 1: Invitrogen™ TrackIt 1 Kb Plus DNA Ladder™  
Lane 2: 200 ng of NR-2713

/Heather Couch/  
Heather Couch

21 AUG 2018

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