SUPPORTING INFECTIOUS DISEASE RESEARCH

# Genomic DNA from Bacillus megaterium, Strain CIP 66.20

# Catalog No. NR-52272

## **Product Description:**

Genomic DNA was extracted from a preparation of *Bacillus megaterium (B. megaterium)*, Strain CIP 66.20. The bacterial preparation used for extraction of genomic DNA was produced by culture of BEI Resources NR-52259 lot 70033090. Genomic DNA was extracted using proprietary technology and is provided in TE buffer (10 mM Tris-HCI and 1 mM EDTA, pH 8).

### Lot: 70033334

# Manufacturing Date: 30MAR2020

TEST	SPECIFICATIONS	RESULTS
Genotypic Analysis Digital DNA-DNA hybridization (dDDH) <sup>1</sup>	≥ 70% dDDH value for identity to <i>B. megaterium</i>	<i>B. megaterium</i> (92.1%) <sup>2</sup>
Agarose Gel Electrophoresis	High molecular weight chromosomal DNA	High molecular weight chromosomal DNA (Figure 1)
Concentration by PicoGreen <sup>®</sup> Measurement	0.7 to 1.5 $\mu g$ in 25 to 100 $\mu L$ per vial	1.1 μg in 34 μL per vial (33.1 μg per mL)
Amount per Vial	0.7 to 1.5 µg	1.1 µg
Functional Activity by PCR Amplification 16S ribosomal RNA gene	~ 1500 base pair amplicon	~ 1500 base pair amplicon
OD <sub>260</sub> /OD <sub>280</sub> Ratio	1.7 to 2.1	1.8
<b>Bacterial Inactivation</b> 100% and 10% of total yield from different pellets plated on agar for 14 days <sup>3,4</sup>	No viable bacteria detected	No viable bacteria detected

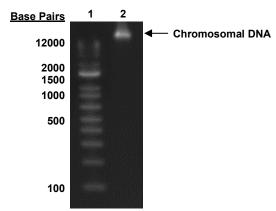
<sup>1</sup>Relatedness between bacterial strains has traditionally been determined using DDH. For additional information, refer to Auch, A. F., et al. "Digital DNA-DNA Hybridization for Microbial Species Delineation by Means of Genome-to-Genome Sequence Comparison." <u>Stand. Genomic Sci.</u> 2 (2010): 117-134. PubMed: 21304684. dDDH analysis was performed using the Type (Strain) Genome Server.

<sup>2</sup>The whole genome of *B. megaterium*, Strain CIP 66.20 was sequenced using the Illumina<sup>®</sup> MiSeq<sup>®</sup> system. *De novo* contig sequences were generated using Unicycler v0.4.8-beta.

<sup>3</sup>14 days under propagation conditions

<sup>4</sup>An extraction procedure was used that has been shown to consistently inactivate 100% of Gram-negative and Gram-positive bacteria.

### Figure 1: Agarose Gel Electrophoresis





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# **Certificate of Analysis for NR-52272**

### /Heather Couch/ Heather Couch

08 JUL 2021

Program Manager or designee, ATCC Federal Solutions

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