

**Genomic DNA from *Borrelia burgdorferi*, Strain B31 (Clone 5A1)**

**Catalog No. NR-56541**

**Product Description:**

Genomic DNA was extracted from a preparation of *Borrelia burgdorferi* (*B. burgdorferi*), strain B31 (clone 5A1). The bacterial preparation used for extraction of genomic DNA was produced by culture of BEI Resources NR-13251 lot 70021457. Genomic DNA was extracted using proprietary technology and is provided in TE buffer (10 mM Tris-HCl, 1 mM EDTA, pH 8.0).

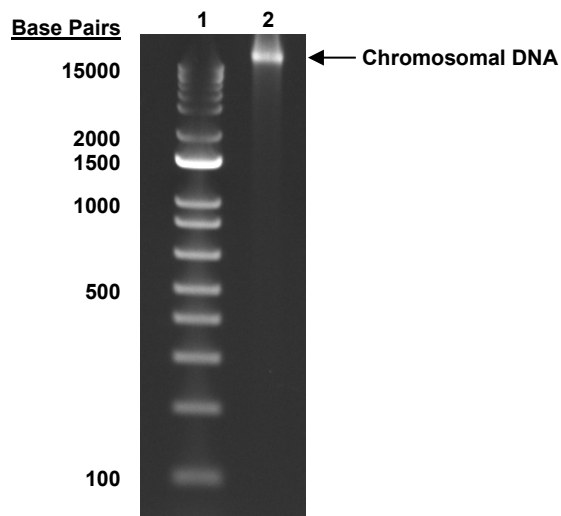
**Lot: 70050993**

**Manufacturing Date: 05APR2022**

TEST	SPECIFICATIONS	RESULTS
<b>Genotypic Analysis</b> Sequencing of 16S ribosomal RNA gene (~ 1440 base pairs)	≥ 99% sequence identity to <i>B. burgdorferi</i> , strain B31 (GenBank: AE000783.1)	99.9% sequence identity to <i>B. burgdorferi</i> , strain B31 (GenBank: AE000783.1)
<b>Agarose Gel Electrophoresis</b>	High molecular weight chromosomal DNA	High molecular weight chromosomal DNA (Figure 1)
<b>Concentration by PicoGreen® Measurement</b>	0.7 to 1.5 µg in 25 to 100 µL per vial	1 µg in 28 µL per vial (35 µg per mL)
<b>Amount per Vial</b>	0.7 to 1.5 µg	1 µg
<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	1.8
<b>Bacterial Inactivation<sup>1</sup></b> 10% of total yield inoculated in Revised Barbour- Stoenner-Kelly medium incubated for 14 days at 30°C in a microaerophilic atmosphere	No viable bacteria detected	No viable bacteria detected

<sup>1</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-56541

/Sonia Bjorum Brower/  
 Sonia Bjorum Brower

Lead Technical Writer or designee, ATCC Federal Solutions

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