SUPPORTING INFECTIOUS DISEASE RESEARCH

Genomic DNA from Mycobacterium microti, Strain NLA000015496

Catalog No. NR-49664

Product Description: Genomic DNA was extracted from a preparation of *Mycobacterium microti* (*M. microti*), strain NLA000015496. *M. microti*, strain NLA000015496 was isolated from a vole in the United Kingdom.

Lot^{1,2}: 63954396

Manufacturing Date: 31MAR2016

TEST	SPECIFICATIONS	RESULTS
Genotypic Analysis Sequencing of 16S ribosomal RNA gene (~ 780 base pairs)	≥ 99% sequence identity to <i>M. microti</i> type strain (GenBank: AF480605.1)	100% sequence identity to <i>M. microti</i> type strain (GenBank: AF480605.1) ³
Agarose Gel Electrophoresis	High molecular weight chromosomal DNA	High molecular weight chromosomal DNA (Figure 1)
Concentration by PicoGreen® Measurement	0.7 to 1.5 μg in 25 to 100 μL	0.8 μg in 141 μL per vial ⁴ (5.9 μg/mL)
Amount per vial	0.7 to 1.5 μg	0.8 µg
Functional Activity by PCR Amplification 16S ribosomal RNA gene	~ 1500 base pair amplicon	~ 1500 base pair amplicon
OD ₂₆₀ /OD ₂₈₀ Ratio	1.7 to 2.1	1.9
Bacterial Inactivation 10% of total yield plated on agar ^{5,6}	No viable bacteria detected	No viable bacteria detected

¹The bacterial preparation used for extraction of genomic DNA was produced from the deposited material. Genomic DNA was extracted using proprietary technology.

²NR-49664 lot 63954396 was vialed in TE buffer (10 mM Tris-HCl, 1 mM EDTA, pH ~ 8.0).

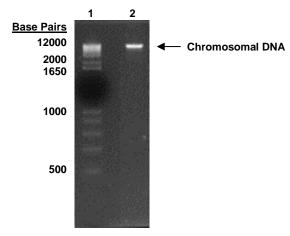
³Also consistent with other *Mycobacterium* species

⁴The volume of genomic DNA per vial is above the current specification, but does not negatively impact the final product.

⁵30 days at 37°C in an aerobic atmosphere with 5% CO₂ on Middlebrook 7H10 agar with OADC enrichment

⁶An extraction procedure was used that has been shown to consistently inactivate 100% of Gram-positive and Gram-negative bacteria.

Figure 1: Agarose Gel Electrophoresis





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