

**Genomic DNA from *Streptococcus pneumoniae*, Strain TCH8431**

**Catalog No. HM-145D**

**Product Description:** Genomic DNA was obtained from a preparation of *Streptococcus pneumoniae* (*S. pneumoniae*), strain TCH8431.

**Lot<sup>1</sup>: 59673066**

**Manufacturing Date: 17JAN2011**

TEST	SPECIFICATIONS	RESULTS
<b>Sequencing of 16S Ribosomal RNA Gene (~ 1470 base pairs)</b>	≥ 99% identical to depositor's sequence Consistent with <i>S. pneumoniae</i>	≥ 99% identical to depositor's sequence Consistent with <i>S. pneumoniae</i>
<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 per vial	0.93 µg in 33 µL per vial (28 µg/mL)
<b>Functional Activity by PCR Amplification</b> 16S ribosomal RNA gene	~ 1500 bp amplicon	~ 1500 bp amplicon
<b>OD<sub>260</sub>/OD<sub>280</sub> Ratio</b>	1.7 to 2.0	2.0
<b>Bacterial Inactivation</b> 10% of total yield plated on Tryptic Soy Agar with 5% sheep blood <sup>2,3</sup>	No viable bacteria detected	No viable bacteria detected

<sup>1</sup>The bacterial preparation used for extraction of genomic DNA was produced by Brain Heart Infusion Broth culture of the deposited material. After incubation for 24 hours at 37°C in an aerobic atmosphere with 5% CO<sub>2</sub>, genomic DNA was extracted using proprietary technology.

<sup>2</sup>7 days at 37°C in an aerobic atmosphere with 5% CO<sub>2</sub>

<sup>3</sup>An extraction procedure was used that has been shown to consistently inactivate 100% of Gram-positive and Gram-negative bacteria.

**Date:** 19 APR 2011

**Signature:**



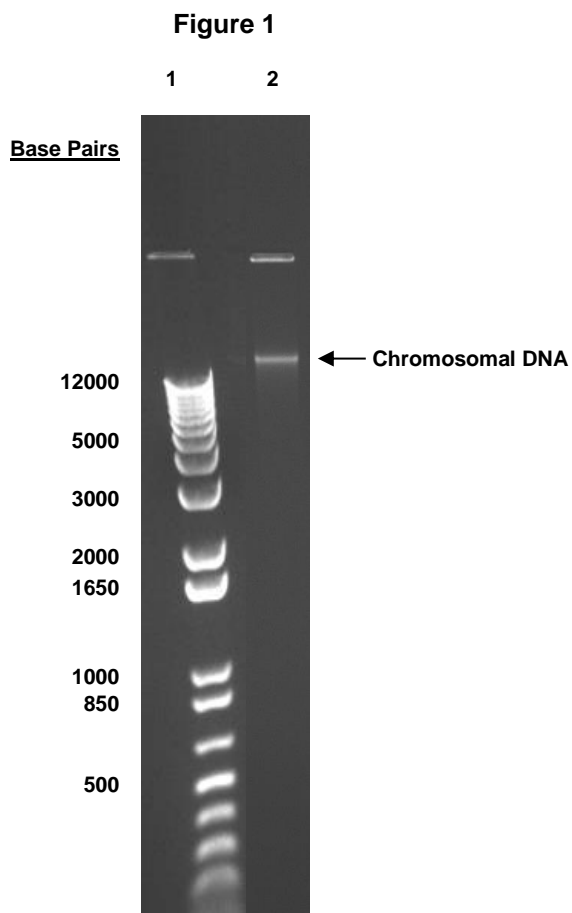
**Title:** Technical Manager, BEI Authentication or designee

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Lane 1: Invitrogen™ TrackIt 1 Kb Plus DNA Ladder™  
Lane 2: 200 ng of HM-145D