

Certificate of Analysis for NR-47416

Staphylococcus aureus subsp. aureus, Strain JE2, Transposon Mutant NE873 (SAUSA300_1991)

Catalog No. NR-47416

Product Description:

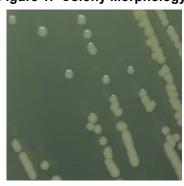
Staphylococcus aureus (S. aureus) subsp. aureus, transposon mutant NE873 was derived from S. aureus subsp. aureus, strain JE2. Mutagenesis occurred through the use of the *mariner*-based transposon *bursa aurealis* resulting in an erythromycin-resistant deletion strain of JE2. S. aureus subsp. aureus, transposon mutant NE873 was created by disruption of SAUSA300_1991, which encodes for accessory gene regulator protein C. Strain JE2 is a plasmid-cured derivative of strain LAC that was isolated in 2002 from a skin and soft tissue infection of an inmate in the Los Angeles County Jail in California, USA. NR-47416 was produced by inoculation of BEI Resources seed lot 62691023 into Tryptic Soy broth with 5 µg per mL erythromycin and incubated for 1 day at 37°C in an aerobic atmosphere. Broth inoculum was added to Tryptic Soy agar with 5 µg per mL erythromycin kolles, which were grown for 2 days at 37°C in an aerobic atmosphere to produce this lot.

Lot: 70048310 Manufacturing Date: 05NOV2021

TEST	SPECIFICATIONS	RESULTS
Phenotypic Analysis		
Cellular morphology	Gram-positive cocci	Gram-positive cocci
Colony morphology 1 day at 37°C in an aerobic atmosphere on Tryptic Soy agar with 5 μg per mL erythromycin	Report results	Circular, convex, entire, smooth and cream (Figure 1)
Motility (wet mount)	Report results	Non-motile
Confirmation of Transposon Insertion ¹	Resistant to erythromycin	Resistant to erythromycin
Purity (post-freeze) 7 days at 37°C in an aerobic atmosphere with and without 5% CO ₂ on Tryptic Soy agar with 5% defibrinated sheep blood	Growth consistent with expected colony morphology	Growth consistent with expected colony morphology
Viability (post-freeze) 1 day at 37°C in an aerobic atmosphere on Tryptic Soy agar with 5 µg per mL erythromycin	Growth	Growth

¹Prior to initiating work, it is recommended that the presence and location of the transposon is confirmed. Gene specific primers should be paired with either the "Upstream" primer (5'-CTCGATTCTATTAACAAGGG-3') for transposons in the "plus" orientation or the "Buster" primer (5'-GCTTTTTCTAAATGTTTTTTAAGTAAATCAAGTAC-3') for transposons in the "minus" orientation. For additional information, refer to Fey, P. D., et al. "A Genetic Resource for Rapid and Comprehensive Phenotype Screening of Nonessential *Staphylococcus aureus* Genes." mBio. 4 (2013): e00537-12. PubMed: 23404398.

Figure 1: Colony Morphology



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Program Manager or designee, ATCC Federal Solutions

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