SUPPORTING INFECTIOUS DISEASE RESEARCH

Borrelia burgdorferi, Signature-Tagged Mutagenesis Library Clone T08TC701 (Gene BB_0002)

Catalog No. NR-25134

Product Description:

Borrelia burgdorferi (B. burgdorferi), strain B31 5A18NP1 STM library clone T08TC701 was produced by signaturetagged mutagenesis (STM) of the BB_0002 gene. NR-25134 was produced by inoculation of the deposited material into Revised Barbour-Stoenner-Kelly medium supplemented with 200 µg per mL kanamycin and 40 µg per mL gentamicin and grown for 10 days at 32°C in a microaerophilic atmosphere to produce this lot. Quality control testing was completed under propagation conditions unless otherwise noted.

Lot: 70045494

Manufacturing Date: 16JUL2021

| TEST | SPECIFICATIONS | RESULTS |
|---|----------------------------|---|
| Phenotypic Analysis | | |
| Cellular morphology | Spirochete | Spirochete |
| Motility (wet mount) | Report results | Motile |
| Purity 11 days at 37°C in an aerobic atmosphere with 5% CO ₂ in Tryptic Soy agar with 5% defibrinated sheep blood | No growth | No growth |
| Viability (post-freeze) | | |
| Visual observation | Growth | Growth |
| LIVE/DEAD [®] <i>Bac</i> Light™ Bacterial Viability | Green fluorescence visible | Green fluorescence visible ¹ |

¹Determined after 11 days at 32°C in a microaerophilic atmosphere in Revised Barbour-Stoenner-Kelly broth supplemented with 200 µg per mL kanamycin and 40 µg per mL gentamicin with LIVE/DEAD[®] *Bac*Light[™] Bacterial Viability Kit, 1000× magnification (Invitrogen[™] L7007). Cells with a compromised membrane that are dead or dying will stain red, while cells with an intact membrane will stain green.

Figure 1: LIVE/DEAD[®] BacLight[™] Bacterial Viability



/Heather Couch/ Heather Couch

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Program Manager or designee, ATCC Federal Solutions

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