

***Borrelia burgdorferi*, Signature-Tagged Mutagenesis Library Clone T04TC201 (Gene IR_BB_S37-BB_S38)**

Catalog No. NR-22763

Product Description: *Borrelia burgdorferi* (*B. burgdorferi*), strain B31 5A18NP1 STM library clone T04TC201 was produced by signature-tagged mutagenesis (STM) of the intergenic region between the BB_S37 and BB_S38 genes.

Lot¹: 70021445

Manufacturing Date: 25JAN2019

| TEST | SPECIFICATIONS | RESULTS |
|--|--------------------------------------|---|
| Phenotypic Analysis Cellular morphology ² Motility (wet mount) | Spirochete Report results | Spirochete Motile |
| Purity (post-freeze)³ | No growth observed | No growth observed |
| Viability (post-freeze) Visual observation LIVE/DEAD [®] BacLight [™] Bacterial Viability | Growth Green fluorescence visible | Growth ² Green fluorescence visible (Figure 1) ⁴ |

¹NR-22763 was produced by inoculation of the deposited material into Revised Barbour-Stoenner-Kelly medium supplemented with 200 µg/mL kanamycin and 40 µg/mL gentamicin and grown for 10 days at 32°C in a microaerophilic atmosphere to produce this lot.

²8 days at 32°C in a microaerophilic atmosphere in Revised Barbour-Stoenner-Kelly broth supplemented with 200 µg/mL kanamycin and 40 µg/mL gentamicin

³Purity of this lot was assessed for 8 days at 37°C in an aerobic atmosphere with 5% CO₂ on Tryptic Soy agar with 5% defibrinated sheep blood.

⁴Determined with LIVE/DEAD[®] BacLight[™] Bacterial Viability Kit, 100x magnification (Invitrogen[™] L7007) after a 8-day incubation at 32°C in a microaerophilic atmosphere in Revised Barbour-Stoenner-Kelly broth supplemented with 200 µg/mL kanamycin and 40 µg/mL gentamicin. 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

04 MAR 2019

Program Manager or designee, ATCC Federal Solutions

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