

Certificate of Analysis for NR-51672

Borrelia recurrentis, Strain PBek

Catalog No. NR-51672

Product Description:

Borrelia recurrentis (B. recurrentis), strain PBek was isolated in 2004 from the blood of a human with louse-borne relapsing fever from a traveler returning from Ethiopia. NR-51672 lot 70027333 was produced by inoculation of the deposited material into Revised Barbour-Stoenner-Kelly broth and grown for 6 days at 33°C in a microaerophilic atmosphere (6-16% O₂ and 2-10% CO₂; BD GasPak™ EZ Campy). Broth inoculum was added to Revised Barbour-Stoenner-Kelly broth and grown for 7 days at 33°C in a microaerophilic atmosphere to produce this lot.

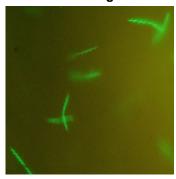
Lot: 70027333 Manufacturing Date: 31OCT2019

TEST	SPECIFICATIONS	RESULTS
Phenotypic Analysis		
Cellular morphology	Spirochete	Spirochete
7 days at 33°C in a microaerophilic atmosphere in		
Revised Barbour-Stoenner-Kelly broth		
Motility (wet mount)	Report results	Motile
Genotypic Analysis		
Sequencing of 16S ribosomal RNA (rRNA) gene	≥ 99% sequence identity to	100% sequence identity to
(1350 base pairs)	B. recurrentis, strain A1	B. recurrentis, strain A1
	(GenBank: CP000993.1)	(GenBank: CP000993.1) ¹
Purity		
7 days at 33°C in a microaerophilic atmosphere in Tryptic	Growth consistent with colony	No growth
Soy agar with 5% defibrinated sheep blood	morphology or no growth	
7 days at 37°C in an aerobic atmosphere with 5% CO ₂	No growth	No growth
in Tryptic Soy agar with 5% defibrinated sheep blood		
Viability (post-freeze)		
Visual observation	Growth	Growth
7 days at 33°C in a microaerophilic atmosphere in		
Revised Barbour-Stoenner-Kelly broth		
LIVE/DEAD [®] <i>Bac</i> Light™ Bacterial Viability	Green fluorescence visible	Green fluorescence visible
		(Figure 1) ²

¹Also consistent with other *Borrelia* species. *B. recurrentis* and *B. duttonii* cannot be differentiated by sequencing of the 16S rRNA gene (Marosevic, D., et al. "First Insights in the Variability of *Borrelia recurrentis* Genomes." <u>PLoS Negl. Trop. Dis.</u> 11 (2017): e0005865. PubMed: 28902847.).

²Determined after 7 days at 33°C in a microaerophilic atmosphere in Revised Barbour-Stoenner-Kelly broth with LIVE/DEAD[®] *Bac*Light™ Bacterial Viability Kit, 1000× magnification (Invitrogen™ L34856). 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



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/Heather Couch/

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Program Manager or designee, ATCC Federal Solutions

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