SUPPORTING INFECTIOUS DISEASE RESEARCH

Borrelia recurrentis, Strain PAbJ

Catalog No. NR-51674

For research use only. Not for human use.

Contributor:

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Manufacturer:

BEI Resources

Product Description:

Bacteria Classification: Spirochaetaceae, Borrelia Species: Borrelia recurrentis Strain: PAbJ Original Source: Parrelia recurrentia (P. recurrenti

- <u>Original Source</u>: *Borrelia recurrentis (B. recurrentis)*, strain PAbJ was isolated in Germany in 2015 from the blood of a human with louse-borne relapsing fever who originated in Somalia and migrated to Germany.^{1,2}
- <u>Comments</u>: *B. recurrentis*, strain PAbJ is reported to be multi-locus sequence type (MLST) ST-669.² The complete genome of *B. recurrentis*, strain PAbJ has been sequenced (BioProject: <u>PRJNA378726</u>).²

B. recurrentis is a large, loosely coiled, motile spirochete transmitted by a single known vector, Pediculus humanus (body louse).^{2,3,4} It is the causative agent of louse-borne relapsing fever (LBRF), an epidemic disease with a distinctive relapsing phenomenon endemic to the Horn of Africa that is re-emerging in Europe, primarily along refugee migration routes.^{2,3} B. recurrentis has the most simple genome of all Borrelia spp., composed of one linear chromosome, seven linear plasmids and 990 protein-coding genes.^{2,3} Genomic analysis indicates that *B. recurrentis* and *B. duttoni*, which causes tick-borne relapsing fever, are nearly identical and cannot be differentiated by sequencing of the 16S ribosomal RNA gene.^{2,3} Approximately 30 genes or gene families are missing or damaged in B. recurrentis, including the RecA and RadA proteins involved in DNA double-strand break repair, suggesting that B. recurrentis is a decaying strain of B. duttonii.^{2,3,4}

Material Provided:

Each vial contains approximately 0.5 mL of bacterial culture in Revised Barbour-Stoenner-Kelly broth supplemented with 15% glycerol.

<u>Note</u>: If homogeneity is required for your intended use, please purify prior to initiating work.

Packaging/Storage:

NR-51674 was packaged aseptically in cryovials. The product is provided frozen and should be stored at -60°C or colder immediately upon arrival. For long-term storage, the vapor phase of a liquid nitrogen freezer is recommended. Freeze-thaw cycles should be avoided.

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Growth Conditions:

<u>Media</u>:

Revised Barbour-Stoenner-Kelly broth or equivalent (Appendix I)

Note: Medium should be prepared fresh before each use.

Incubation:

Temperature: 32°C to 34°C

Atmosphere: Aerobic with 5% CO₂

Propagation:

<u>Note</u>: It is recommended that NR-51674 be cultured in 24-well plates until growth is established from the frozen vial.

- 1. Place the frozen vial in a 35°C to 37°C water bath and thaw for approximately 2 to 3 minutes. Immerse the vial just enough to cover the frozen material. Do not agitate the vial. Do not leave the vial in the water bath after it is thawed.
- Immediately after thawing, aseptically transfer the contents of the vial to 2 wells of a 24-well plate containing 1.5 mL fresh Revised Barbour-Stoenner-Kelly medium per well.
- Incubate the plate at 32°C to 34°C. Do not shake culture during growth. It may take up to 21 days for the culture to establish from the frozen state.

<u>Note</u>: NR-51674 should be subcultured during the log phase of growth, as viability of the culture may decrease quickly.

Maintenance:

- Monitor growth of the culture by live/dead staining every 3 to 6 days. When the culture has reached the log phase, transfer approximately 2 mL into to a T-25 tissue culture flask containing 8 mL fresh Revised Barbour-Stoenner-Kelly medium.
- 2. Incubate the plate at 32°C to 34°C.
- 3. Transfer the culture every 3 to 21 days as described in Maintenance steps 1 and 2. The transfer interval will depend on the size of the inoculum and the quality of the medium. This should be determined by performing live/dead staining every 3 to 6 days. Do not allow the culture to overgrow. Viability of the culture may be affected soon after reaching peak density.

Citation:

Acknowledgment for publications should read "The following reagent was obtained through BEI Resources, NIAID, NIH: *Borrelia recurrentis*, Strain PAbJ, NR-51674."

Biosafety Level: 2

Appropriate safety procedures should always be used with this material. Laboratory safety is discussed in the following publication: U.S. Department of Health and Human Services, Public Health Service, Centers for Disease Control and Prevention, and National Institutes of Health. <u>Biosafety in Microbiological and Biomedical Laboratories</u>. 5th ed. Washington, DC: U.S. Government Printing Office, 2009; see www.cdc.gov/biosafety/publications/bmbl5/index.htm.

Disclaimers:

You are authorized to use this product for research use only. It is not intended for human use.

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References:

- 1. Fingerle, V., Personal Communication.
- Marosevic, D., et al. "First Insights in the Variability of Borrelia recurrentis Genomes." <u>PLoS Negl. Trop. Dis.</u> 11 (2017): e0005865. PubMed: 28902847.
- Warrell, D. A. "Louse-Borne Relapsing Fever (*Borrelia recurrentis* Infection)." <u>Epidemiol. Infect.</u> 147 (2019): e106. PubMed: 30869050.
- Lescot, M., et al. "The Genome of *Borrelia recurrentis*, the Agent of Louse-Borne Relapsing Fever, is a Degraded Subset of Tick-Borne *Borrelia duttonii*." <u>PLoS Genet.</u> 4 (2008): e1000185. PubMed: 18787695.

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APPENDIX I: REVISED BSK MEDIUM (ATCC® MEDIUM: 1914)

1. Prepare the Revised BSK medium directly before each use following the recipe below by dissolving each component one at a time in distilled water:

| HEPES | 5.64 g |
|----------------------------------|--------|
| Neopeptone | 4.7 g |
| Sodium citrate | 0.7 g |
| Glucose | 5.64 g |
| NaHCO ₃ | 2.0 g |
| TC-Yeastolate | 2.0 g |
| Sodium pyruvate | 0.75 g |
| N-acetylglucosamine | 0.37 g |
| Bovine serum albumin, fraction V | 47.0 g |
| Distilled water | 840 mL |

- 2. Adjust the pH of the base medium to 7.5 using 1 N HCl or 1 N NaOH and filter-sterilize using a 0.22 µm filter.
- 3. Aseptically add the next two components to the base medium:

CMRL 1066 Medium, 10× (w/o Glutamine and NaHCO3)100.0 mLHeat-inactivated rabbit serum60.0 mL

- 4. Mix well and aseptically dispense into appropriate vessels. The medium may be stored in aliquots of 50 mL in freezer-safe vessels and stored frozen at -20°C until use. Once thawed, each aliquot should be kept at 2°C to 8°C and used within one month.
- 5. Adjust the pH of the complete medium to 7.5 to 7.6, as needed, using sterile solutions of 1 N HCl or 1 N NaOH, before use.

<u>Note</u>: Medium should be prepared fresh directly before each use or immediately aliquoted and frozen at -20°C until needed. Once thawed, each aliquot should be kept at 2°C to 8°C and used within one month.