

SUPPORTING INFECTIOUS DISEASE RESEARCH

Product Information Sheet for NR-50050

Genomic DNA from *Burkholderia mallei*, Strain China 7 (NBL 7)

Catalog No. NR-50050

For research use only. Not for human use.

Contributor:

ATCC[®]

Manufacturer:

BEI Resources

Product Description:

Genomic DNA was extracted from a preparation of *Burkholderia mallei* (*B. mallei*), strain China 7 (NBL 7) (BEI Resources NR-23 which was derived from ATCC[®] 23344[™]). Genome variability upon passage has been reported to be a feature of *B. mallei*, strain China 7 (ATCC[®] 23344[™]). Genomic DNA from BEI Resources NR-4071 (a preparation of *B. mallei* strain China 7 that was derived from ATCC[®] 23344[™] via several passages by different individuals prior to its deposit at BEI Resources) is available as BEI Resources NR-9318.

B. mallei, strain China 7 was isolated from postmortem cultures of knee fluid, skin pustules, and blood of a Chinese soldier who died in Burma (1944) from a glanders-melioidosis type of infection. The complete genomic sequence of *B. mallei*, strain China 7 has been determined (GenBank: CP000010 and CP000011).²

NR-50050 has been qualified for PCR applications by amplification of approximately 1500 base pairs of the 16S ribosomal RNA gene.

Material Provided:

Each vial of NR-50050 contains 0.7 μ g to 1.5 μ g of bacterial genomic DNA in 10 mM Tris-HCl, pH 8 - 8.5. The vial should be centrifuged prior to opening.

Packaging/Storage:

NR-50050 was packaged aseptically in screw-capped plastic cryovials. The product is provided frozen and should be stored at -20°C or colder immediately upon arrival. For long-term storage, the product should be stored at -80°C. Freezethaw cycles should be minimized. Note: NR-50050 is not provided in EDTA; for long-term storage, EDTA may be added to a final concentration of 0.1 mM to 1 mM.

Citation:

Acknowledgment for publications should read "The following reagent was obtained through BEI Resources, NIAID, NIH: Genomic DNA from *Burkholderia mallei*, Strain China 7 (NBL 7), NR-50050."

Biosafety Level: 1

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:

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References:

- Romero, C. M., et al. "Genome Sequence Alterations Detected upon Passage of *Burkholderia mallei* ATCC 23344 in Culture and in Mammalian Hosts." <u>BMC Genomics</u> 7 (2006): 228-238. PubMed: 16953889.
- Nierman, W. C., et al. "Structural Flexibility in the Burkholderia mallei Genome." <u>Proc. Natl. Acad. Sci.</u> USA 101 (2004): 14246-14251. <u>PubMed</u>: 15377793.
- Bauernfeind, A., et al. "Molecular Procedure for Rapid Detection of Burkholderia mallei and Burkholderia pseudomallei." J. Clin. Microbiol. 36 (1998): 2737-2741.

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PubMed: 9705426.

- Godoy, D., et al. "Multilocus Sequence Typing and Evolutionary Relationships Among the Causative Agents of Melioidosis and Glanders, *Burkholderia pseudomallei* and *Burkholderia mallei*." J. Clin. Microbiol. 41 (2003): 2068-2079. PubMed: 12734250.
- Gee, J. E., et al. "Use of 16S rRNA Gene Sequencing for Rapid Identification and Differentiation of Burkholderia pseudomallei and B. mallei." J. Clin. Microbiol. 41 (2003): 4647-4654. PubMed: 14532197.
- Ong, C., et al. "Patterns of Large-Scale Genomic Variation in Virulent and Avirulent *Burkholderia* Species." <u>Genome Res.</u> 14 (2004): 2295-2307. PubMed: 15520292.

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