SUPPORTING INFECTIOUS DISEASE RESEARCH

Genomic DNA from *Yersinia pestis*, Strain A1122

Catalog No. NR-3040

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For research use only. Not for human use.

Contributor:

Centers for Disease Control and Prevention, Division of Vector-Borne Infectious Diseases, Fort Collins, Colorado, USA

Manufacturer:

BEI Resources

Product Description:

Genomic DNA was isolated from a preparation of Yersinia pestis (Y. pestis), strain A1122.

Y. pestis A1122 was isolated from a California ground squirrel *(Spermophilus beecheyi)* in California in 1939.¹ It contains the pMT1/pFra and the pPCP1/pPla plasmids, but lacks the pCD1/pYV plasmid that is essential for virulence as well as the unstable *pgm* locus.²⁻⁴ The complete genome and plasmid sequences are available (GenBank: <u>CP002956</u>, <u>CP002957</u> and <u>CP002958</u>).

The presence of the pMT1/pFra and the pPCP1/pPla plasmids and the absence of the pCD1/pYV plasmid and the *pgm* locus in NR-3040 have been confirmed by Next Generation Sequencing. NR-3040 has been qualified for PCR applications by amplification of approximately 1500 base pairs of the 16S ribosomal RNA gene.

Material Provided:

Each vial of lot 63815634 contains approximately 0.7 μ g to 1.5 μ g of bacterial genomic DNA in TE buffer (10 mM Tris-HCl, 1 mM EDTA, pH ~ 8.0). Each vial of lot 7513127 contains approximately 5 μ g of bacterial genomic DNA in TE buffer (10 mM Tris-HCl, 1 mM EDTA, pH ~ 7.4). The concentration, expressed as μ g per μ L, is shown on the Certificate of Analysis. The vial should be centrifuged prior to opening.

Packaging/Storage:

NR-3040 was packaged aseptically in screw-capped plastic cryovials. The product is provided frozen on dry ice and should be stored at -20°C or colder immediately upon arrival. Freeze-thaw cycles should be minimized.

Citation:

Acknowledgment for publications should read "The following reagent was obtained through BEI Resources, NIAID, NIH: Genomic DNA from *Yersinia pestis*, Strain A1122, NR-3040."

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:

- Huang, X. Z., M. C. Chu, D. M. Engelthaler, and L. E. Lindler. "Genotyping of a Homogeneous Group of *Yersinia pestis* Strains Isolated in the United States." <u>J.</u> <u>Clin. Microbiol.</u> 40 (2002): 1164-1173. PubMed: 11923326.
- Hinchcliffe, S. J., et al. "Application of DNA Microarrays to Study the Evolutionary Genomics of *Yersinia pestis* and *Yersinia pseudotuberculosis*." <u>Genome Res.</u> 13 (2003): 2018-2029. PubMed: 12952873.

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- Chu, M. C., X. Q. Dong, X. Zhou, and C. F. Garon. "A Cryptic 19-Kilobase Plasmid Associated with U.S. Isolates of Yersinia pestis: A Dimer of the 9.5-Kilobase Plasmid." <u>Am. J. Trop. Med. Hyg.</u> 59 (1998): 679-686. PubMed: 9840581.
- Parkhill, J., et al. "Genome Sequence of Yersinia pestis, the Causative Agent of Plague." <u>Nature</u> 413 (2001): 523-527. PubMed: 11586360.
- Chu, M. C. <u>Laboratory Manual of Plague Diagnostic</u> <u>Tests</u>. Centers for Disease Control and Prevention, Atlanta, 2000.

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