

Product Information Sheet for NR-2561

Yersinia pestis F1-V Fusion Protein, Monomer-Enriched Antigen, Recombinant from Escherichia coli

Catalog No. NR-2561

This reagent is the tangible property of the U.S. Government.

For research use only. Not for human use.

Contributor:

National Institutes of Allergy and Infectious Diseases, National Institutes of Health

Product Description:

Recombinant Yersinia pestis (Y. pestis) F1-V fusion protein, monomer-enriched antigen was purified from Escherichia coli and depleted of DNA and endotoxin. Originally developed by the U.S. Army Medical Research Institute of Infectious Disease (USAMRIID), F1-V is a fusion protein consisting of the Fraction 1 (F1) capsular protein and the virulenceassociated (V) regulatory protein from *Y. pestis* (GenPept: AAY23169).^{2,3}

Material Provided:

Each vial contains approximately 1.5 mg of recombinant F1-V fusion protein in 20 mM L-arginine, 10 mM sodium chloride and 1 mM L-cysteine. The concentration and postvialing pH are shown on the Certificate of Analysis for each

Packaging/Storage:

NR-2561 was packaged in glass serum vials. It is provided frozen and should be stored at -70°C or colder immediately upon arrival. Thawed material should be held at 2°C to 8°C and used within 8 hours.

Functional Activity:

NR-2561 was demonstrated to be functionally active based on its reactivity with antibodies to both the F1 and V proteins. NR-2561 is protective in a Y. pestis lethal challenge murine model.

Citation:

Acknowledgment for publications should read "The following reagent was obtained through the NIH Biodefense and Emerging Infections Research Resources Repository, NIAID, NIH: Yersinia pestis F1-V Fusion Protein, Monomer-Enriched Antigen, Recombinant from Escherichia coli, NR-2561."

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. Biosafety in Microbiological and Biomedical Laboratories. 5th ed. Washington, DC: U.S. Government Printing Office, 2007; see www.cdc.gov/od/ohs/biosfty/bmbl5/bmbl5toc.htm.

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References:

- 1. Goodin, J. L., et al. "Purification and Protective Efficacy of Monomeric and Modified Yersinia pestis Capsular F1-V Antigen Fusion Proteins for Vaccination against Plague." Protein Expr. Purif. 53 (2007): 63-79. PubMed: 17293124.
- 2. Powell, B. S., et al. "Design and Testing for a Nontagged F1-V Fusion Protein as Vaccine Antigen against Bubonic and Pneumonic Plague." Biotechnol. Prog. 21 (2005): 1490-1510. PubMed: 16209555.
- 3. Heath, D. G., et al. "Protection against Experimental Bubonic and Pneumonic Plague by a Recombinant Capsular F1-V Antigen Fusion Protein Vaccine." Vaccine 16 (1998): 1131-1137. PubMed: 9682370.
- 4. Glynn, A., et al. "Protection against Aerosolized Yersinia pestis Challenge following Homologous and Heterologous

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- Prime-Boost with Recombinant Plague Antigens." Infect. Immun. 73 (2005): 5256–5261. PubMed: 16041052.
- Jones, T., et al. "Intranasal Protollin/F1-V Vaccine Elicits Respiratory and Serum Antibody Responses and Protects Mice against Lethal Aerosolized Plague Infection." <u>Vaccine</u> 24 (2006): 1625–1632. PubMed: 16243411.
- Santi, L., et al. "Protection Conferred by Recombinant Yersinia pestis Antigens Produced by a Rapid and Highly Scalable Plant Expression System." <u>Proc. Natl. Acad. Sci.</u> <u>U.S.A.</u> 103 (2006): 861–866. PubMed: 16410352.

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