

SUPPORTING INFECTIOUS DISEASE RESEARCH

# **Product Information Sheet for NR-4488**

Monoclonal Anti-Bacillus anthracis Protective Antigen, Clone PA 2II 14B7-1-1 (purified IgG, produced in vitro)

## Catalog No. NR-4488

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

# For research use only. Not for human use.

#### **Contributor:**

United States Army Medical Research Institute of Infectious Diseases, Fort Detrick, Maryland

#### Manufacturer:

**BEI Resources** 

#### **Product Description:**

Antibody Class: IgG1k

Monoclonal antibody prepared against recombinant protective antigen (PA) from *Bacillus anthracis* (*B. anthracis*)<sup>1-3</sup> was purified from hybridoma clone PA 2II 14B7-1-1<sup>4-6</sup> supernatant by protein G affinity chromatography. The B cell hybridoma was generated by the fusion of Sp2/0-Ag14 myeloma cells with splenocytes from BALB/c mice immunized with purified recombinant protein.

#### **Material Provided:**

Each vial of NR-4488 contains approximately 100  $\mu$ L of purified monoclonal antibody in PBS. The concentration, expressed as mg per mL, is shown on the Certificate of Analysis.

#### Packaging/Storage:

NR-4488 was packaged aseptically in screw-capped plastic vials and is provided frozen on dry ice. The product should be stored at -20°C or colder immediately upon arrival. Freezethaw cycles should be avoided.

## **Functional Activity:**

NR-4488 reacts with *B. anthracis* protective antigen in western blot assays.

### Citation:

Acknowledgment for publications should read "The following reagent was obtained through BEI Resources, NIAID, NIH: Monoclonal Anti-Bacillus anthracis Protective Antigen, Clone PA 2II 14B7-1-1 (purified IgG, produced *in vitro*), NR-4488."

## **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, 2009; see www.cdc.gov/biosafety/publications/bmbl5/index.htm.

#### **Disclaimers:**

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## References:

- 1. S. H. Leppla, personal communication.
- Leppla, S. H. "Production and Purification of Anthrax Toxin." Methods Enzymol. 165 (1988): 103-116. PubMed: 3148094.
- Leppla, S. H. "Purification and Characterization of Adenylyl Cyclase from *Bacillus anthracis*." <u>Methods</u> <u>Enzymol.</u> 195 (1991): 153-168. PubMed: 1903483.
- Little, S. F., et al. "Characterization of Lethal Factor Binding and Cell Receptor Binding Domains of Protective Antigen of Bacillus anthracis using Monoclonal Antibodies." <u>Microbiology</u> 142 (1996): 707-715. PubMed: 8868446.
- Little, S. F. and J. R. Lowe. "Location of Receptor-Binding Region of Protective Antigen from Bacillus anthracis." <u>Biochem. Biophys. Res. Commun.</u> 180 (1991): 531-537. PubMed: 1953724.

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 Little, S. F., S. H. Leppla and E. Cora. "Production and Characterization of Monoclonal Antibodies to Protective Antigen Component of *Bacillus anthracis* Toxin." <u>Infect.</u> <u>Immun.</u> 56 (1988): 1807-1813. PubMed: 3384478.

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