

Product Information Sheet for NR-50173

SUPPORTING INFECTIOUS DISEASE RESEARCH

N7 Neuraminidase (NA) Protein with N-Terminal Histidine Tag from Influenza Virus, A/harbor seal/Germany/1/2014 (H10N7), Recombinant from Baculovirus

Catalog No. NR-50173

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

For research use only. Not for human use.

Contributor:

Florian Krammer, Ph.D., Departments of Medicine and Microbiology, Icahn School of Medicine at Mount Sinai, One Gustave L. Levey Place, New York, New York, USA, provided under government contract (CEIRS)

Manufacturer:

BEI Resources

Product Description:

A recombinant form of the N7 neuraminidase (NA) protein from influenza A virus, A/harbor seal/Germany/1/2014 (H10N7)¹ containing an N-terminal histidine tag was produced in Sf9 insect cells using a baculovirus expression vector system and was purified by nickel affinity chromatography. The predicted ectodomain coding region of the NA gene was fused to a synthetic gene segment encoding an N-terminal six histidine tag followed by a tetramerization domain from vasodilator-stimulated phosphoprotein (VASP) and a thrombin cleavage site.^{2,3}

Material Provided:

Each vial contains 50 μg to 100 μg of purified recombinant NA protein in 50 mM Na₂HPO₄, pH 8, with 500 mM NaCl and 5 mM EDTA. The protein content in μg and the concentration, expressed as $\mu g/mL$, are shown on the Certificate of Analysis.

Packaging/Storage:

Purified recombinant NA protein was packaged aseptically, in screw-capped plastic cryovials. This product is provided on ice bricks and should be stored at -20°C immediately upon arrival.

Functional Activity:

NR-50173 was demonstrated to be functionally active based on its ability to cleave the fluorogenic substrate 2'-(4-methylumbelliferyl)-α-D-N-acetylneuraminic acid (4-MUNANA).⁴

Citation:

Acknowledgment for publications should read "The following reagent was obtained through BEI Resources, NIAID, NIH: N7 Neuraminidase (NA) Protein with N-Terminal Histidine Tag from Influenza Virus, A/harbor seal/Germany/1/2014 (H10N7), Recombinant from Baculovirus, NR-50173."

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:

- Bodewes, R., et al. "Avian Influenza A(H10N7) Virus-Associated Mass Deaths Among Harbor Seals." <u>Emerg.</u> <u>Infect. Dis.</u> 21 (2015): 720-722. PubMed: 25811303.
- Kühnel, K., et al. "The VASP Tetramerization Domain is a Right-Handed Coiled Coil Based on a 15-Residue Repeat." <u>Proc. Natl. Acad. Sci. USA</u> 101 (2004): 17027-17032. PubMed: 15569942.
- Margine, I., P. Palese, and F. Krammer. "Expression of Functional Recombinant Hemagglutinin and Neuraminidase Proteins from the Novel H7N9 Influenza

BEI Resources www.beiresources.org E-mail: contact@beiresources.org

Tel: 800-359-7370 Fax: 703-365-2898



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- Virus Using the Baculovirus Expression System." <u>J. Vis. Exp.</u> 6 (2013): e51112. PubMed: 24300384.
- Wetherall, N. T., et al. "Evaluation of Neuraminidase Enzyme Assays Using Different Substrates to Measure Susceptibility of Influenza Virus Clinical Isolates to Neuraminidase Inhibitors: Report of the Neuraminidase Inhibitor Susceptibility Network." J. Clin. Microbiol. 41 (2003): 742-750. PubMed: 12574276.

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BEI Resources www.beiresources.org E-mail: contact@beiresources.org
Tel: 800-359-7370

Fax: 703-365-2898