

Product Information Sheet for NR-45106

Monoclonal Anti-Murine Coronavirus Nucleocapsid (N) Protein, Clone 1.16.1 (produced *in vitro*)

Catalog No. NR-45106

For research use only. Not for human use.

Contributor:

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Manufacturer:

BEI Resources

Product Description:

Antibody Class: IgG1κ

Mouse monoclonal antibody prepared against the nucleocapsid (N) protein of murine coronavirus (formerly known as mouse hepatitis virus), strain JHM was purified from clone 1.16.1 hybridoma supernatant by protein G affinity chromatography. The B cell hybridoma was generated by the fusion of P3X63Ag8.653 mouse myeloma cells with immunized mouse splenocytes.¹

Coronaviral N protein is a phosphoprotein with both structural and regulatory functions.^{2,3} It plays a primary role in packaging the RNA genome, and is also involved in viral RNA synthesis, translation and modulation of host cell metabolism.^{4,5,6,7,8} These multifunctional properties make N protein an attractive antiviral target.

Material Provided:

Each vial of NR-45106 contains approximately 100 μ L of purified monoclonal antibody in PBS. The concentration, expressed as mg per mL, is shown on the Certificate of Analysis.

Packaging/Storage:

NR-45106 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. Freeze-thaw cycles should be avoided.

Functional Activity:

NR-45106 is reactive with NCTC clone 1469 cells infected with recombinant murine coronavirus icA59 (BEI Resources NR-43000) in indirect immunofluorescence assays. The antibody is also reported to react with all strains of murine coronavirus examined to date, to function in immunocytochemistry, immunohistochemistry, immunoprecipitation and western blot assays, and to be non-neutralizing.

Citation:

Acknowledgment for publications should read "The following reagent was obtained through BEI Resources, NIAID, NIH: Monoclonal Anti-Murine Coronavirus Nucleocapsid (N) Protein, Clone 1.16.1 (produced *in vitro*), NR-45106."

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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- Calvo, E., et al. "Phosphorylation and Subcellular Localization of Transmissible Gastroenteritis Virus Nucleocapsid Protein in Infected Cells." <u>J. Gen. Virol.</u> 86 (2005): 2255-2267. PubMed: 16033973.
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- Baric, R. S., et al. "Interactions Between Coronavirus Nucleocapsid Protein and Viral RNAs: Implications for Viral Transcription." <u>J. Virol.</u> 62 (1988): 4280-4287. PubMed: 2845140.
- Stohlman, S. A., et al. "Specific Interaction Between Coronavirus Leader RNA and Nucleocapsid Protein." <u>J. Virol.</u> 62 (1988): 4288-4295. PubMed: 2845141.
- Nelson, G. W., S. A. Stohlman and S. M. Tahara. "High Affinity Interaction Between Nucleocapsid Protein and Leader/Intergenic Sequence of Mouse Hepatitis Virus RNA." <u>J. Gen. Virol.</u> 81 (2000): 181-188. PubMed: 10640556.
- Eléouët, J. F., et al. "The Viral Nucleocapsid Protein of Transmissible Gastroenteritis Coronavirus (TGEV) is Cleaved by Caspase-6 and -7 during TGEV-Induced Apoptosis." <u>J. Virol.</u> 74 (2000): 3975-3983. PubMed: 10756009.
- 9. Leibowitz, J. L., Personal Communication.

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