RD SYSTEMS a biotechne brand

MERS-CoV Nucleocapsid Antibody

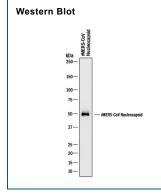
Monoclonal Mouse IgG_{2B} Clone # 1038729 Catalog Number: MAB10729

DESCRIPTION	
Species Reactivity	MERS-CoV
Specificity	Detects Mouse MERS-CoV Nucleocapsid in direct ELISAs.
Source	Monoclonal Mouse IgG _{2B} Clone # 1038729
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	<i>Spodoptera frugiperda</i> insect ovarian cell line <i>SI</i> /21-derived mouse MERS-CoV Nucleocapsid Met1-Thr411 Accession # YP_007188586.1
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 µm filtered solution in PBS.

APPLICATIONS

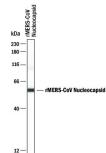
Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.			
	Recommended Concentration	Sample	
Western Blot	1 µg/mL	Recombinant MERS-CoV Nucleocapsid protein	
Simple Western	25 µg/mL	Recombinant MERS-CoV nucleocapsid	

DATA



Detection of MERS-CoV MERS-CoV Nucleocapsid by Western Blot. Western blot shows recombinant MERS-CoV Nucleocapsid protein. PVDF membrane was probed with 1 µg/mL of Mouse Anti-Mouse MERS-CoV Nucleocapsid Monoclonal Antibody (Catalog # MAB10729) followed by HRP-conjugated Anti-Mouse IgG Secondary Antibody (Catalog # HAF018). A specific band was detected for MERS-CoV Nucleocapsid at approximately 49 kDa (as indicated). This experiment was conducted under reducing conditions and using Western Blot Buffer Group 1.

Simple Western



Detection of Nucleocapsid by Simple WesternTM. Simple Western lane view shows recombinant MERS-CoV nucleocapsid, loaded at 0.2 mg/mL. A specific band was detected for Nucleocapsid at approximately 57 kDa (as indicated) using 25 µg/mL of Mouse Anti-MERS-CoV Nucleocapsid Monoclonal Antibody (Catalog # MAB10729) . This experiment was conducted under reducing conditions and using the 12-230 kDa separation system.

PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.5 mg/mL in sterile PBS.	
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C	
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. • 12 months from date of receipt, -20 to -70 °C as supplied. • 1 month, 2 to 8 °C under sterile conditions after reconstitution. • 6 months, -20 to -70 °C under sterile conditions after reconstitution.	

BACKGROUND

MERS-CoV, which causes the Middles East Respiratory Syndrome (MERS), belongs to a family of viruses known as coronaviruses that are commonly comprised of four structural proteins: Spike protein (S), Envelope protein (E), Membrane protein (M), and Nucleocapsid protein (N) (1). While the S, E and M proteins build up the viral envelope, the N protein is involved transcription, replication and packaging of the viral RNA genome into a helical ribonucleocapsid (RNP) (1, 2). The MERS-CoV N protein is a ~45 kDa protein composed of two independent structural domains connected by a linker region. The N-terminal region contains an Intrinsically Disordered Region (3) and an RNA binding domain (4), the linker region interacts with the M protein and the C-terminal region contains a self-association domain (1, 2). The MERS-CoV N protein shares 46.3% and 4.5% amino acid sequence identity with SARS-CoV-1 and SARS-CoV-2 N protein, respectively. MERS-CoV N proteins have been shown to inhibit Type I Interferon(IFN) production(1). In addition, the N protein is an abundant protein during coronavirus infection and displays high immunogenic activity, making it a promising therapeutic target (5-7).

References:

- 1. Li, Y. et al. (2019) Engineering. 5:940.
- 2. Hurst, K. R. et al. (2009) J. Virol. 83:7221.
- 3. Wang, Y. et al. (2015) Acta. Crystallogr. F. Struct. Biol. Commun. 71:977.
- 4. Papageorgiou, N. et al. (2016) Acta. Crystallogr. D. Struct. Biol. 72:192.
- 5. Che, X. Y. et al. (2004) J. Clin. Microbiol. 42:2629.
- 6. Guan, M. et al. (2004) Clin. Diagn. Lab. Immunol. 11:287.
- 7. Chang, C-K. *et al.*(2016) Drug Discov. Today. **21**:562.

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Global bio-techne.com info@bio-techne.com techsupport@bio-techne.com TEL +1 612 379 2956 USA TEL 800 343 7475 Canada TEL 855 668 8722 China TEL +86 (21) 52380373 Europe | Middle East | Africa TEL +44 (0)1235 529449