

Influenza A NP (9G8): sc-101352

BACKGROUND

Influenza A viruses are negative sense, single-stranded, segmented RNA viruses which are hosted by birds but may infect several species of mammals. All known subtypes are endemic in birds. Influenza A subtypes are classified based on the combination of the virus coat glycoproteins hemagglutinin (HA) and neuraminidase (NA) subtypes. There are 16 different HA antigens (H1-H16) and 9 different NA antigens (N1-N9) for Influenza A. The extent of infection into host organisms is determined by HA, which interacts with cell surface proteins containing oligosaccharides with terminal sialyl residues. Influenza A nucleoprotein (NP) associates with its RNA genome and is present in eight separate segments of ribonucleoprotein (RNP), each of which has to be present for successful replication.

REFERENCES

- Green, N., et al. 1982. Immunogenic structure of the influenza virus hemagglutinin. *Cell* 28: 477-487.
- Gething, M.J., et al. 1986. Expression of wild-type and mutant forms of influenza hemagglutinin: the role of folding in intracellular transport. *Cell* 46: 939-950.
- Webster, R.G., et al. 1987. Influenza virus A pathogenicity: the pivotal role of hemagglutinin. *Cell* 50: 665-666.

SOURCE

Influenza A NP (9G8) is a mouse monoclonal antibody raised against recombinant Influenza A NP.

PRODUCT

Each vial contains 50 µg IgG_{2a} kappa light chain in 500 µl of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

Influenza A NP (9G8) is recommended for detection of nucleoprotein (NP) of Influenza A virus origin by Western Blotting (starting dilution to be determined by researcher, dilution range 1:100-1:5000), immunofluorescence (starting dilution to be determined by researcher, dilution range 1:50-1:2500) and solid phase ELISA (starting dilution to be determined by researcher, dilution range 1:100-1:5000); non cross-reactive with Influenza B or other respiratory viruses.

Molecular Weight of Influenza A NP: 56 kDa.

RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgGκ BP-HRP: sc-516102 or m-IgGκ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker™ Molecular Weight Standards: sc-2035, UltraCruz® Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use m-IgGκ BP-FITC: sc-516140 or m-IgGκ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

STORAGE

Store at 4° C, ****DO NOT FREEZE****. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

SELECT PRODUCT CITATIONS

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- Gao, S., et al. 2015. Interaction of NS2 with AIMP2 facilitates the switch from ubiquitination to SUMOylation of M1 in Influenza A Virus-infected cells. *J. Virol.* 89: 300-311.
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- Lejal, N., et al. 2018. Turning off NADPH oxidase-2 by impeding p67phox activation in infected mouse macrophages reduced viral entry and inflammation. *Biochim. Biophys. Acta* 1862: 1263-1275.
- Phuong, N.H., et al. 2018. Development and characterization of monoclonal antibodies against nucleoprotein for diagnosis of Influenza A virus. *J. Microbiol. Biotechnol.* 28: 809-815.
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- Zheng, W., et al. 2019. Naproxen exhibits broad anti-influenza virus activity in mice by impeding viral nucleoprotein nuclear export. *Cell Rep.* 27: 1875-1885.e5.
- Chen, Y., et al. 2020. Synthesis of a hexavalent betulinic acid derivative as a hemagglutinin-targeted influenza virus entry inhibitor. *Mol. Pharm.* 17: 2546-2554.
- Sharma, A., et al. 2020. Influenza A virus nucleoprotein activates the JNK stress-signaling pathway for viral replication by sequestering host filamin A protein. *Front. Microbiol.* 11: 581867.
- Farre, C., et al. 2020. Specific and sensitive detection of Influenza A virus using a biotin-coated nanoparticle enhanced immunomagnetic assay. *Anal. Bioanal. Chem.* E-published.

RESEARCH USE

For research use only, not for use in diagnostic procedures.