

# Recombinant SARS-CoV-2 AY.2 Spike (GCN4-IZ) His-tag

Catalog Number: 10923-CV

## DESCRIPTION

Source

Human embryonic kidney cell, HEK293-derived sars-cov-2 Spike protein

SARS-CoV-2 AY.2 Spike (Val16-Lys1211)(Thr19Arg, Val70Phe, Gly142Asp, Glu156Gly, Phe157 del, Arg158 del, Arg222Val, Lys417Asn, Leu452Arg, Thr478Lys, Asp614Gly, Pro681Arg, Asp950Asn) (Arg682Ser, Arg685Ser, Lys986Pro, Val987Pro) Accession # YP\_009724390.1

GCN4-IZ

6-His tag

N-terminus C-terminus

N-terminal Sequence Val16 Analysis **Predicted Molecular** 

138 kDa

Mass

SPECIFICATIONS	
SDS-PAGE	142-170 kDa, under reducing conditions.
Activity	Measured by its binding ability in a functional ELISA with Recombinant Human ACE-2 His-tag (Catalog # 933-ZN).
Endotoxin Level	<0.10 EU per 1 µg of the protein by the LAL method.
Purity	>95%, by SDS-PAGE under reducing conditions and visualized by silver stain.
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details.

## PREPARATION AND STORAGE

Reconstitution Reconstitute at 500  $\mu g/mL$  in PBS

Shipping The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.

## 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
- 3 months, -20 to -70 °C under sterile conditions after reconstitution.

## SDS-PAGE



Recombinant SARS-CoV-2 AY.2 Spike (GCN4-IZ) His-tag Protein SDS-PAGE. 2 µg/lane of Recombinant SARS-CoV-2 AY.2 Spike (GCN4-IZ) His-tag Protein (Catalog # 10923-CV) was resolved with SDS-PAGE under reducing (R) and non-reducing (NR) conditions and visualized by Coomassie® Blue staining, showing bands at 142-170 kDa.

Rev. 10/26/2021 Page 1 of 2





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## BACKGROUND

SARS-CoV-2, which causes the global pandemic coronavirus disease 2019 (Covid-19), belongs to a family of viruses known as coronaviruses that also include MERS and SARS-CoV-1. Coronaviruses are commonly comprised of four structural proteins: Spike protein (S), Envelope protein (E), Membrane protein (M) and Nucleocapsid protein (N) (1). The SARS-CoV-2 S protein is a glycoprotein that mediates membrane fusion and viral entry. The S protein is homotrimeric, with each ~180-kDa monomer consisting of two subunits, S1 and S2 (2). In SARS-CoV-2, as with most coronaviruses, proteolytic cleavage of the S protein into S1 and S2 subunits is required for activation. The S1 subunit is focused on attachment of the protein to the host receptor while the S2 subunit is involved with cell fusion (3-5). A metallopeptidase, angiotensin-converting enzyme 2 (ACE2), has been identified as a functional receptor for SARS-CoV-2 through interaction with a receptor binding domain (RBD) located at the C-terminus of S1 subunit (6,7). The S protein of SARS-CoV-2 shares 75% and 29% amino acid (aa) sequence identity with the S protein of SARS-CoV-1 and MERS, respectively. AY.2 is a sublineage of SARS-CoV-2 delta plus variant which carries the aa substitution Thr19Arg, Val70Phe, Gly142Asp, Glu156Gly, Phe157 del, Arg158 del, Arg222Val, Lys417Asn, Leu452Arg, Thr478Lys, Asp614Gly, Pro681Arg, and Asp950Asn in the Spike was first identified in India and quickly spread to US through England and Japan. It is reported to partially escape neutralization of antibodies and vaccines (8).

### References:

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- 7. Wong, S.K. et al. (2004) J. Biol. Chem. 279:3197.
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