

Product Data Sheet

SARS-CoV-2 S glycoprotein (V483A, HEK293, His)

| Cat. No.: | HY-P72042 | |
|-------------------|--------------------------|--|
| Synonyms: | E2 Peplomer protein | |
| Species: | Virus | |
| Source: | HEK293 | |
| Accession: | PODTC2 (R319-F541,V483A) | |
| Gene ID: | 43740568 | |
| Molecular Weight: | Approximately 27.8 kDa | |

| DDODEDTIEC | | | | | |
|----------------------|---|--------------------------------|-------------------|------------|--|
| PROPERTIES | | | | | |
| AA Sequence | | | | | |
| · | RVQPTESIVR | FPNITNLCPF | GEVFNATRFA | SVYAWNRKRI | |
| | SNCVADYSVL | Y N S A S F S T F K | CYGVSPTKLN | DLCFTNVYAD | |
| | SFVIRGDEVR | QIAPGQTGKI | ADYNYKLPDD | FTGCVIAWNS | |
| | N N L D S K V G G N | YNYLYRLFRK | SNLKPFERDI | STEIYQAGST | |
| | PCNGAEGFNC | YFPLQSYGFQ | PTNGVGYQPY | RVVVLSFELL | |
| | НАРАТУССРК | KSTNLVKNKC | VNF | | |
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| Appearance | Lyophilized powder. | | | | |
| E | | | | | |
| Formulation | Lyophilized from a 0.2 μ m solution of 20 mM Tris-HCl, 0.5 M NaCl, 6% Trehalose, pH 8.0. | | | | |
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| Endotoxin Level | <1 EU/µg, determined by LAL method. | | | | |
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| Reconsititution | It is not recommended to reconstitute to a concentration less than 100 μ g/mL in ddH ₂ O. | | | | |
| | | | | | |
| Storage & Stability | Stored at -20°C for 2 years. After reconstitution, it is stable at 4°C for 1 week or -20°C for longer (with carrier protein | | | | |
| | recommended to freeze a | iliquots at -20°C or -80°C for | extended storage. | | |
| | | | | | |
| Shipping | Room temperature in continental US;may vary elsewhere. | | | | |

DESCRIPTION

Background

The spike (S) glycoprotein of coronaviruses is known to be essential in the binding of the virus to the host cell at the advent of the infection process. The viral pathogen responsible, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), binds to the host receptor through its spike (S) glycoprotein, which mediates membrane fusion and viral entry^{[1][2]}. The spike (S) glycoprotein of coronaviruses is known to be essential in the binding of the virus to the host cell at the advent of the infection process. The viral pathogen responsible, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), binds to the host receptor through its spike (S) glycoprotein, which mediates membrane fusion and viral entry^{[1][2]}. The spike (S) glycoprotein of coronaviruses is known to be essential in the binding of the virus to the host cell at the advent of the host receptor through its spike (S) glycoprotein, which mediates membrane fusion and viral entry^{[1][2]}. The spike (S) glycoprotein of coronaviruses is known to be essential in the binding of the virus to the host cell at the advent of the infection process. The viral pathogen responsible, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), binds to the host receptor through its spike (S) glycoprotein, which mediates membrane fusion and viral entry^{[1][2]}. The spike (S) glycoprotein of coronaviruses is known to be essential in the binding of the virus to the host cell at the advent of the infection process. The viral pathogen responsible, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), binds to the host receptor through its spike (S) glycoprotein, which mediates membrane fusion and viral entry^{[1][2]}.

REFERENCES

[1]. Shen S, et al. Expression, glycosylation, and modification of the spike (S) glycoprotein of SARS CoV. Methods Mol Biol. 2007;379:127-135.

[2]. Wang S, et al. AXL is a candidate receptor for SARS-CoV-2 that promotes infection of pulmonary and bronchial epithelial cells. Cell Res. 2021;31(2):126-140.

Caution: Product has not been fully validated for medical applications. For research use only.

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