Recombinant SARS-CoV-2 NSP8 Protein, His-tagged

Cat. No. HUM-363 **Lot. No.** (See product label)

SPECIFICATION

Product Overview	SARS-CoV-2 NSP8 protein, His-tag is a recombinant full-length viral protein expressed in E. coli. SARS-CoV-2, previously known as the 2019 Novel Coronavirus (2019-nCoV), causes the pandemic COVID-19 disease.
Tag	His
Background	In December 2019 a novel coronavirus, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), formerly known as the 2019 novel coronavirus (2019-nCoV) was identified in Wuhan, China, causing a world-wide pandemic (Wu et al., 2020). Three coronaviruses, SARS-CoV, MERS-CoV, and SARS-CoV-2 have been identified as being a highly pathogenic for humans, and there is currently no effective antiviral treatment. Therefore, studies are focused on rapid development of vaccines and antiviral drugs to prevent and treat coronavirus infection. There are several potential strategies to pharmacologically fight against the disease (COVID-19), including vaccines, monoclonal antibodies, oligonucleotide-based therapies, peptides, interferon therapies, and small-molecule drugs (Dömling & Gao, 2020). Coronavirus RNA genome encodes a complex replication machinery consisting of 16 viral non-structural proteins (NSPs). NSP7 and NSP8 stabilize regions of NSP12 (RNA-dependent RNA polymerase; RdRp) involved in RNA binding and are essential for a highly active NSP12 polymerase complex. NSP7 has been suggested to confer RNA-binding properties to the trimeric polymerase complex (Subissi et al., 2014). NSP8 has been shown to act as an oligo(U)-templated polyadenylyltransferase and a robust (mono/oligo) adenylate transferase (Tvarogova et al., 2019). The NSP7/NSP8/NSP12 polymerase complex associates with NSP14, which is required

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	to safeguard coronavirus replication fidelity (Subissi et al., 2014). Specific mutations in NSP7 and NSP8 may have a role in stabilization of the replication/transcription complex (Reshamwala et al., 2021). Amino acid substitutions in RdRp (P323L) have been shown to cause conformational changes near the NSP8 binding site that may affect virus replication and transcription (Alosaimi et al., 2021).
Purity	>90% purity.
Formulation	Presented as liquid in 520 mM Tris, pH 7.5, 300mM NaCl, 200mM imidazole, 1 mM DTT, 10% glycerol.
Notes	This product is intended for research and manufacturing uses only. It is not a diagnostic device. The user assumes all responsibility for care, custody and control of the material, including its disposal, in accordance with all regulations.
Type	Recombinant
ClassID 1	Infectious Disease

GENE INFORMATION

Synonyms	SARS-CoV-2 NSP8

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