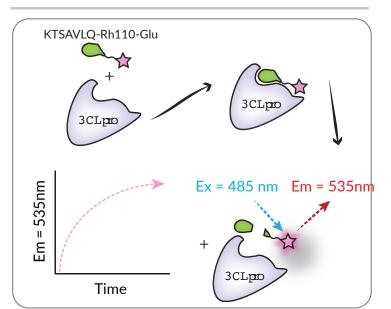
# His<sub>10</sub>-SARS-CoV-2 Mpro (3CLpro)

Cat. No. SBB-DE0129 Lot. No. 220300129

### His<sub>10</sub>-SARS-CoV-2 Mpro (3CL-Pro)

The human SARS-CoV-2 coronavirus harbors two proteases Papain Like Protease PLpro and 3CLpro (Chymotrypsin like Protease) or Mpro (Main-Protease) which is a C30-type cysteine protease. The viral genome encodes more than 20 proteins, with 3CLpro located within the non-structural protein 5 (nsp5) section of the viral polypeptide that cleaves together with PLpro polyproteins (PP1A and PP1AB) into individual functional components.

3CLpro recognizes the peptide sequence LQ[S/A/G] where it cleaves c-terminal to the amino acid glutamine (use product SBB-PS0130, KTSAVLQ-Rh110-Glu as universal substrate). The protease 3CLpro is a potential drug target for coronavirus infections due to its essential role in processing the polyproteins that are translated from the viral RNA. The X-ray structures of the unliganded SARS-CoV-2 protease 3CLpro and its complex with an  $\alpha$ -ketoamide inhibitor provides a basis for design of  $\alpha$ -ketoamide inhibitors. This SARS Coronavirus recombinant 3CLpro is N-terminally His<sub>10</sub>-tagged and expressed in *E.coli*.





#### **Product Information**

Quantity: 50 µg Molecular Weight: 35.6 kDa

Concentration: 80 µM, 2.8 mg/mL

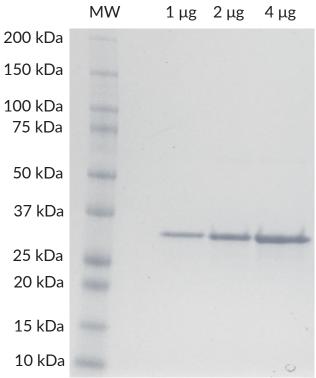
Purity: >98% by SDS-PAGE

Storage Buffer: 50 mM HEPES pH 7.5, 100 mM NaCl, 1 mM TCEP

Storage: -80C, Avoid multiple freeze / thaw

Usage: Working concentrations of this enzyme range from 10 to 100 nM using KTSAVLQ-Rh110 (SBB-PS0130) as substrate.

## **Quality Control and Performance Data**



 $His_{10}$ -Mpro SDS-PAGE. From left to right, increasing amunts of  $His_{10}$ -Mpro loaded onto a 4-20% SDS-PAGE gel, stained with coomassie brillant blue. Purity is >98%.

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www.southbaybio.com

Contact: info@southbaybio.com

5941 Optical Ct, Suite 229 San Jose, CA 95138 USA

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#### References

1) Fran Robson et al. "Coronavirus RNA Proofreading: Molecular Basis and Therapeutic Targeting". Molecular Cell, VOLUME 79, ISSUE 5, P710-727, SEPTEMBER 03, 2020, https://doi.org/10.1016/j.molcel.2020.07.027

2) Linlin Zhang et al. "Crystal structure of SARS-CoV-2 main protease provides a basis for design of improved  $\alpha$ -ketoamide inhibitors". SCIENCE • 20 Mar 2020 • Vol 368, Issue 6489 • pp. 409-412 • DOI: 10.1126/science. abb3405

3) Kiemer, L., Lund, O., Brunak, S. et al. Coronavirus 3CLproproteinase cleavage sites: Possible relevance to SARS virus pathology. BMC Bioinformatics 5, 72 (2004). https://doi.org/10.1186/1471-2105-5-72

4) Chen YW, Yiu CPB and Wong KY. Prediction of the SARS-CoV-2 (2019-nCoV) 3C-like protease (3CLpro) structure: virtual screening reveals velpatasvir, ledipasvir, and other drug repurposing candidates [version 2; peer review: 3 approved]. F1000Research 2020, 9:129 (https://doi.org/10.12688/f1000research.22457.2)

5) Ocain TD et al. "alpha-Keto amide inhibitors of aminopeptidases". Journal of Medicinal Chemistry. (February 1992). 35 (3): 451–6. doi:10.1021/jm00081a005.

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