## SARS-CoV2-2 lota B.1.526 Variant Recombinant Spike Trimer His Tag

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## #2656

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| Applications          | Detection   | Clonality         | Isotype    |
|-----------------------|---|-------------------|------------|
| ELISA                 | Anti-SARS-CoV-2 mAb   | SARS-CoV-2        | SARS-CoV-2 |
| Format:               | His tag   |                   |            |
| Cross Reactivity:     | Predicted to work with mouse, rat and other homologues.   |                   |            |
| Formulation:          | 1X PBS, 0.025% NaN3   |                   |            |
| Preparation:          | His tag purification  |                   |            |
| Reactivity:           | Other   |                   |            |
| Recommended<br>Usage: | Control antigen for SARS-CoV-2 (COVID-19) diagnostic assays. Contains C-terminal His tag used for purification.   |                   |            |
| Immunogen:            | N/A   |                   |            |
| Description:          | N/A   |                   |            |
| References:           | 1. Walls A.C. et al., 2019 Cell, 176: 1026-103<br>2. Tang T. et al., 2020, Antiviral Res., 178:10<br>3. Jiang S et al., 2022, Trends Immunol, 41:3<br>4. Bosch BJ, et al., 2003, J Virol 77:8801-88 | 1479.<br>355-359. |            |



Concentration-response curves for binding of CoV2 spike protein lota variant to human ACE2 in cell-free ELISA-type assays. Microtiter wells were coated with 100 uL of each spike trimer at 2 ug/mL in PBS at 4?C overnight. The wells were washed with PBS and blocked with 200 µL of 1% BSA/PBS. ACE2-Fc was serially diluted from 2 µg/mL in 1% BSA/PBS. The blocker was discarded, and the wells were incubated with 100 µL of serially diluted ACE2-Fc at 37?C for 1 hour. The wells were washed with PBS and the bound ACE2-Fc was detected with 100 µL of Peroxidase AffiniPure Goat Anti-Human IgG, Fc? fragment specific (Jackson Immuno Research 109-035-098)(1:5,000 in 1% BSA/PBS) at 37?C for 1 hour. The wells were washed with PBS and the wells were developed with 100 µL of MB/E Ultra Sensitive, Blue, Horseradish Peroxidase Substrate (EMD Millipore ES022-500ml) at RT for 5 min. The reaction was stopped with 100 µL of 0.6N H2SO4 and the signals were read at 450 nm