## SARS-CoV-2 BA.2.86.1 Omicron Variant Recombinant Spike Trimer His Tag

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Isotype

Clonality

**Catalog:** #2788-203 **Store at:** 2-8<sup>o</sup>C

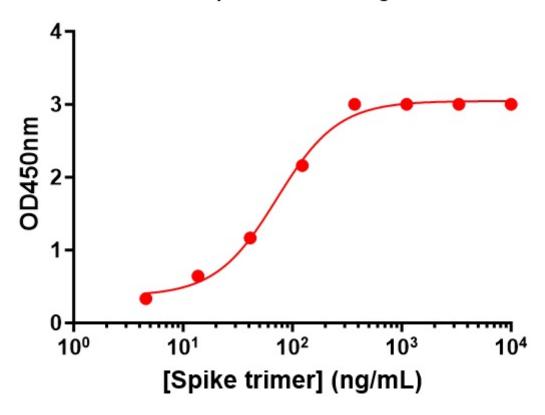
**Detection** 

For Research Use Only. Not For Use In Diagnostic Procedures.

**Applications** 

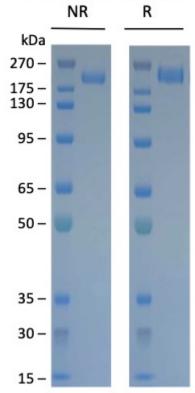
ELISA	Anti-SARS-CoV-2 mAb	SARS-CoV-2	Other
Format:	Unconjugated		
Cross Reactivity:	Predicted to work with mouse, rat and other homologues.		
Formulation:	PBS, 0.025% NaN3		
Preparation:	His tag purification		
Reactivity:	Other		
Recommended Usage:	Control antigen for SARS-CoV-2 (COVID-19) diagnostic assays. Contains C-terminal His tag used for purification.		
Immunogen:	N/A		
Description:			
References:	1. Walls A.C. et al., 2019 Cell, 176: 102 2. Tang T. et al., 2020, Antiviral Res., 1 3. Jiang S et al., 2022, Trends Immunol 4. Bosch BJ, et al., 2003, J Virol 77:880	.78:10479. l, 41:355-359.	

## BA.2.86.1 spike trimer binding to ACE2



Concentration response curves for binding of CoV2 spike protein to human ACE2 in cell-free ELISA-type assays. Microtiter wells were coated with 100 uL of ACE2-Fc at 2 ug/mL in PBS at 4C overnight. The wells were washed with PBS and blocked with 200  $\mu$ L of 1% BSA/PBS at RT for 1 hour. Spike trimer was serially diluted from 10  $\mu$ g/mL in 1% BSA/PBS. The blocker was discarded, and the wells were incubated with 100  $\mu$ L of serially diluted spike trimer at RT for 1 hour. The wells were washed with PBS and the bound spike trimer was detected with 100  $\mu$ L of Peroxidase AffiniPure Rabbit Anti-His Tag (Jackson Immuno Research 300-035-240)(1:5,000 in 1% BSA/PBS) at RT for 1 hour. The wells were washed with PBS and the wells were developed with 100  $\mu$ L of MB/E Ultra-Sensitive, Blue, Horseradish Peroxidase Substrate (EMD Millipore ES022-500ml) at RT for 5 min. The reaction was stopped with 100  $\mu$ L of 0.6N H2SO4 and the signals were read at 450 nm using a plate reader (Biotek).

SARS-CoV2 BA.2.86.1 Recombinant Spike Trimer Protein His Tag



SARS-CoV-2 Spike Trimer, His Tag BA.2.86.1 Omicron on SDS-PAGE under non-reducing (NR) and reducing (R) conditions. The gel was stained with Coomassie G-250 per manufacturer instructions. The purity of the protein is greater than 95% and it is between 175 and 270 kDa.