## SARS-CoV-2 XBB.1/XBB.1.9 Omicron Variant recombinant Spike Trimer His Tag

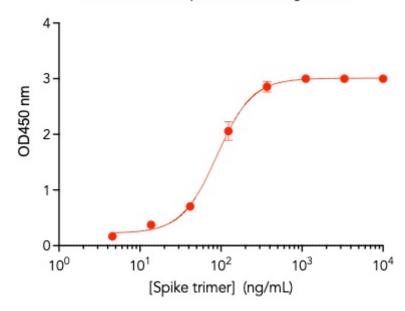
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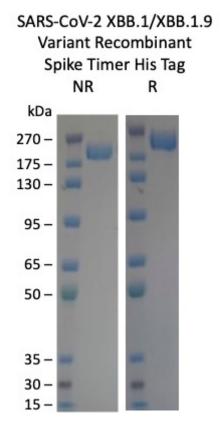
For Research Use Only. Not For Use In Diagnostic Procedures.

<b>Applications</b> ELISA	<b>Detection</b> Anti-SARS-CoV-2 mAb	<b>Clonality</b> SARS-CoV-2	<b>Isotype</b> SARS-CoV-2
		57 IN S COV 2	37113 607 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 Usage:	Control antigen for SARS-CoV-2 (COVID-19) diagnostic assays. Contains C-terminal His tag used for purification.		
Immunogen:	N/A		
Description:			
References:	<ol> <li>Walls A.C. et al., 2019 Cell, 176: 1026-1039.</li> <li>Tang T. et al., 2020, Antiviral Res., 178:10479.</li> <li>Jiang S et al., 2022, Trends Immunol, 41:355-359.</li> <li>Bosch BJ, et al., 2003, J Virol 77:8801-8811.</li> </ol>		

XBB.1/XBB.1.9 spike trimer binding to ACE2



Concentration-response curves for binding of CoV2 spike protein XBB.1 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  $\mu$ L of 1% BSA/PBS. ACE2-Fc was serially diluted from 2  $\mu$ g/mL in 1% BSA/PBS. The blocker was discarded, and the wells were incubated with 100  $\mu$ L 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  $\mu$ L 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  $\mu$ 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-CoV-2 Spike Trimer, His Tag (XBB.1/XBB1.9 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.