

Catalog: #2656

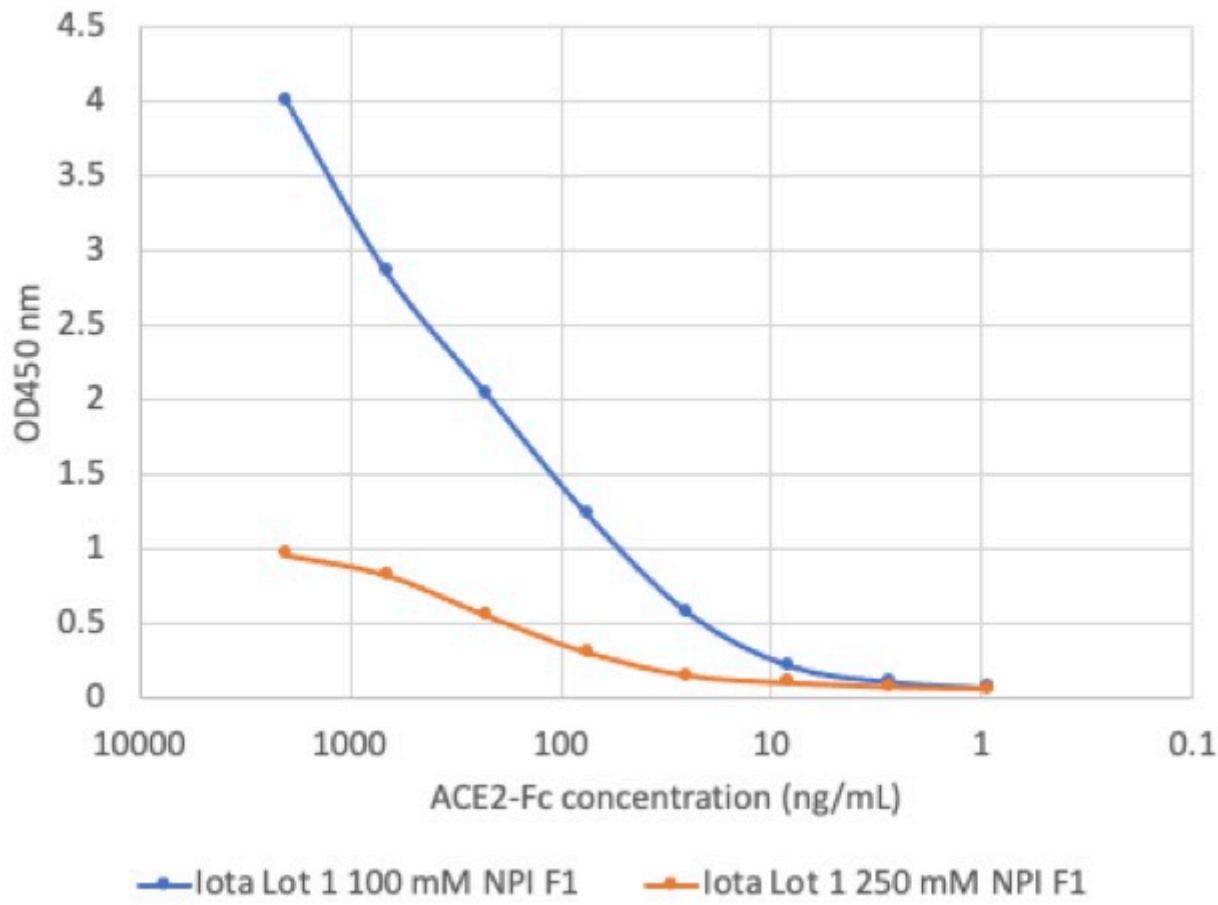
Store at: 2-8°C

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

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 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 style="list-style-type: none"> 1. Walls A.C. et al., 2019 Cell, 176: 1026-1039. 2. Tang T. et al., 2020, Antiviral Res., 178:10479. 3. Jiang S et al., 2022, Trends Immunol, 41:355-359. 4. Bosch BJ, et al., 2003, J Virol 77:8801-8811.

Binding of ACE2-Fc to Iota variant fractions (2 µg/mL)



Concentration-response curves for binding of CoV2 spike protein Iota variant to human ACE2 in cell-free ELISA-type assays. Microtiter wells were coated with 100 µL of each spike trimer at 2 µg/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 \square 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 using a plate reader (Biotek).