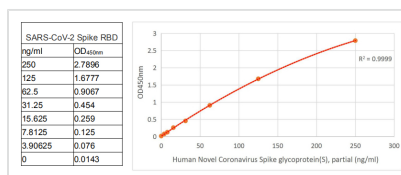




# SARS-CoV-2 Spike RBD Antibody Pair 1

<b>Product Code</b>	CSB-EAP33245
<b>Uniprot No.</b>	P0DTC2
<b>Immunogen</b>	Llama with human IgG1 Fc
<b>Species Reactivity</b>	Human Novel Coronavirus (SARS-CoV-2/ 2019-nCoV)
<b>Tested Applications</b>	S-ELISA
<b>Form</b>	Liquid
<b>Product Type</b>	Antibody Pairs
<b>Immunogen Species</b>	Human Novel Coronavirus (SARS-CoV-2/ 2019-nCoV)
<b>Protein Names</b>	Human Novel Coronavirus Spike glycoprotein (S)
<b>Notes</b>	We recommend using the capture antibody at a concentration of 1ug/ml and the detection antibody at a concentration of 0.42ug/ml. Optimal dilutions should be determined experimentally by the researcher.

## Image



CSB-EAP33245 is a solid phase sandwich Enzyme Linked-Immuno-Sorbent Assay (Sandwich ELISA). An antibody specific for SARS-CoV-2 Spike RBD has been pre-coated onto the microwells. The SARS-CoV-2 Spike RBD protein in samples is captured by the coated antibody after incubation. Following extensive washing, another antibody HRP conjugated specific for SARS-CoV-2 Spike RBD is added to detect the captured SARS-CoV-2 Spike RBD protein. Followed by Tetramethylbenzidine (TMB) reagent. Solution containing sulfuric acid is used to stop color development and the color intensity which is proportional to the quantity of bound protein is measurable at 450nm.

<b>Host</b>	Capture: Llama with human IgG1 Fc Detection: Mouse with human IgG1 Fc
<b>Components</b>	Capture: CSB-EAP33245C Detection: CSB-EAP33245B(HRP) Reagents are sufficient for at least 5 x 96 well plates using recommended protocol.
<b>Storage-Buffer</b>	Capture: 50% Glycerol, 0.01M PBS, PH 7.4 Detection: 50% Glycerol, 0.01M PBS, PH 7.4