

Purified anti-SARS-CoV-2 S Protein S1 Antibody

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| Catalog# / Size | 945101 / 25 µg 945102 / 100 µg |
| Clone | A20103O |
| Regulatory Status | RUO |
| Other Names | S1, Spike Protein |
| Isotype | Rat IgG2a, κ |
| Description | <p>SARS-CoV-2 is a respiratory virus which causes coronavirus disease 2019 (COVID-19). The coronavirus spike (S) glycoprotein is a class I viral fusion protein on the outer envelope of the virion that plays a critical role in viral infection by recognizing host cell receptors and mediating fusion of the viral and cellular membranes. The S glycoprotein is synthesized as a precursor protein consisting of ~1,300 amino acids that is then cleaved into an amino (N)-terminal S1 subunit (~700 amino acids) and a carboxyl (C)-terminal S2 subunit (~600 amino acids). Three S1/S2 heterodimers assemble to form a trimer spike protruding from the viral envelope. The S1 subunit contains a receptor-binding domain (RBD) that can specifically bind to angiotensin-converting enzyme 2 (ACE2), the receptor on target cells. Triggered by receptor binding, proteolytic processing and/or acidic pH in the cellular compartments, the class I viral fusion protein undergoes a transition from a metastable pre-fusion state to a stable post-fusion state during infection, in which the receptor-binding subunit is cleaved, and the fusion subunit undergoes large-scale conformational rearrangements to expose the hydrophobic fusion peptide, induce the formation of a six-helix bundle, and bring the viral and cellular membranes close for fusion. The trimeric SARS coronavirus (SARS-CoV-2) S glycoprotein consisting of three S1-S2 heterodimers binds the cellular receptor angiotensin-converting enzyme 2 (ACE2) and mediates fusion of the viral and cellular membranes through a pre- to post-fusion conformation transition.</p> |

Product Details

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| Verified Reactivity | SARS-CoV-2 |
| Antibody Type | Monoclonal |
| Host Species | Rat |
| Immunogen | Partial recombinant SARS-CoV-2 S protein corresponding to S1 subunit |
| Formulation | Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide |
| Preparation | The antibody was purified by affinity chromatography. |
| Concentration | 0.5 mg/mL |
| Storage & Handling | The antibody solution should be stored undiluted between 2°C and 8°C. |
| Application | WB - Quality tested Direct ELISA, FC - Verified |
| Recommended Usage | Each lot of this antibody is quality control tested by western blotting. For western blotting, the suggested use of this reagent is 1.0 µg/mL. For Direct ELISA, a concentration of 114.9 ng/mL is recommended. For flow cytometric staining, the suggested use of this reagent is ≤ 0.5 µg per million cells in 100 µL volume. It is recommended that the reagent be titrated for optimal performance for each application. |
| Application Notes | BioLegend offers multiple clones that recognize SARS-CoV-2 S protein S1. Clone A20103H displayed the strongest performance for western blot of all clones validated for the application. |
| Product Citations | 1. Mamedov T, <i>et al.</i> 2021. Viruses. 13:. PubMed |
| RRID | AB_2890875 (BioLegend Cat. No. 945101) AB_2890875 (BioLegend Cat. No. 945102) |

Antigen Details

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| Structure | Spike glycoprotein is a homotrimer. Each monomer consists of 1,273 amino acids with a theoretical molecular weight of 141 kD, and consists of S1 and S2 subunits. |
| Distribution | Viral envelope protein, host cell membrane, host cell endoplasmic reticulum-Golgi intermediate compartment membrane |
| Function | Mediates fusion between virus and host cell membranes |
| Interaction | ACE2 |
| Ligand/Receptor | ACE2 |
| Biology Area | COVID-19 |
| Antigen References | <ol style="list-style-type: none">1. Walls AC, <i>et al.</i> 2020. <i>Cell</i>. 181(2):281-292.2. Yan R, <i>et al.</i> 2020. <i>Science</i>. 367 (6485):1444-1448.3. Wrapp D, <i>et al.</i> 2020. <i>Science</i>. 367 (6483):1260-1263.4. Shang J, <i>et al.</i> 2020. <i>PNAS</i>. 117(21):11727-11734 |
| Gene ID | 43740568 |

Related Protocols

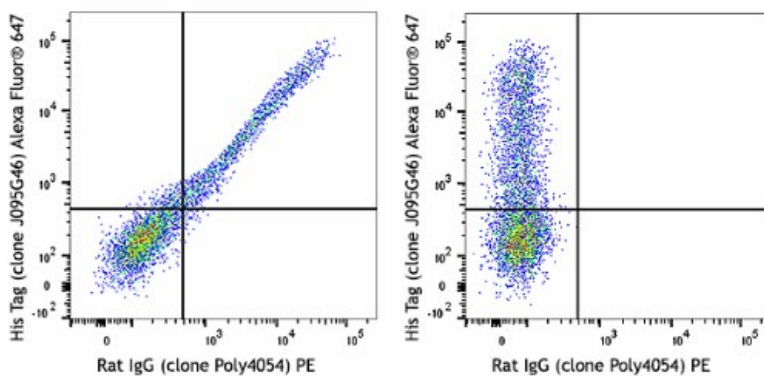
[Cell Surface Flow Cytometry Staining Protocol](#)

[Western Blotting Protocol](#)

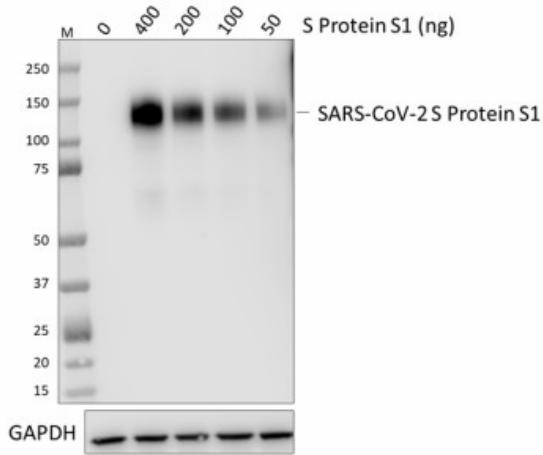
Other Formats

Purified anti-SARS-CoV-2 S Protein S1 Antibody

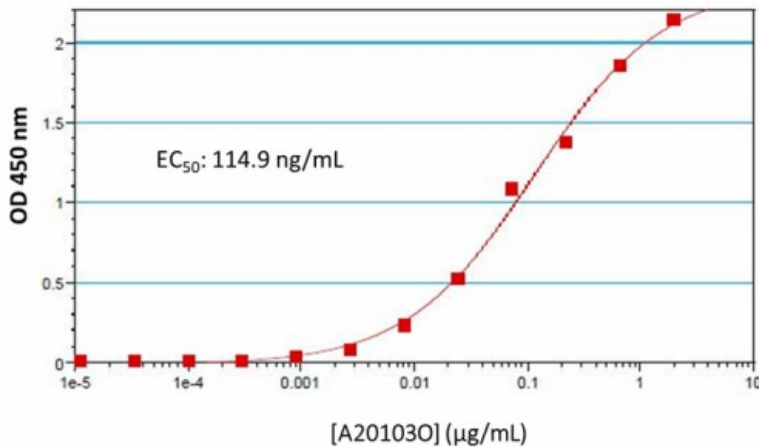
Product Data



His-tagged SARS-CoV-2 S protein S1 transfected CHO cells were stained with anti-S1 (clone A20103O) purified (left) or rat IgG2a, κ isotype control (clone RTK2758) purified (right) followed by anti-rat IgG PE (clone Poly4054) and Alexa Fluor® 647 anti-His Tag (clone J095G46).



Whole cell extracts (15 μ g total protein) from HeLa cells mixed with the indicated amount of His-tagged recombinant SARS-CoV-2 S Protein S1 (Cat. No. 792906) were resolved by 4-12% Bis-Tris gel electrophoresis, transferred to a PVDF membrane, and probed with 1.0 μ g/mL (1:500 dilution) purified anti-SARS-CoV-2 S Protein S1 antibody (clone A20103O) overnight at 4°C. Proteins were visualized by chemiluminescence detection using HRP goat anti-human IgG antibody at a 1:3000 dilution. Direct-Blot™ HRP anti-GAPDH antibody (Cat. No. 607904) was used as a loading control at a 1:25000 dilution (lower). Lane M: Molecular weight marker.



200 ng of recombinant SARS-CoV-2 S protein S1 + S2 (Cat. No. 793704) was coated onto a Costar™ 96-well high binding assay plate and incubated with a dilution series of purified anti-SARS-CoV-2 S protein S1 antibody (clone A20103O). Bound antibodies were detected with HRP goat anti-mouse IgG antibody (Cat. No. 405306) followed by TMB substrate solution. Absorbance was measured at 450 nm.

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