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SARS-CoV-2 Spike S1 monoclonal antibody (S71)

ANT0091 100 μg

Description Coronavirus disease 2019 is a newly emerging infectious disease currently spreading across the world. It

is caused by a novel coronavirus, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). The spike (S) protein of SARS-CoV-2, which plays a key role in the receptor recognition and cell membrane fusion process, is composed of two subunits, S1 and S2. The S1 subunit contains a receptor-binding domain that recognizes and binds to the host receptor angiotensin-converting enzyme 2, while the S2 subunit mediates viral cell membrane fusion by forming a six-helical bundle via the two-heptad repeat domain. The specific antibodies against different subunits of spike (S) protein of SARS-CoV-2 are excellent

research tools for the research against coronavirus

Product type Monoclonal antibody

ImmunogenSpike1 (S)SourceMouse

Reacts with Conformational SARS-CoV-2 Spike 1

Specificity Anti SARS CoV-2 Spike1 monoclonal antibody detects full-length Spike1 also in the glycosylated form. The

antibody does not cross react with SARS CoV-2 Spike2 and SARS CoV-2 Spike1 RBD domain.

Tested applications WB, ELISA.

Recommended dilutions Recommended starting dilutions can vary lot-to-lot.

Consult the product information label in the package for lot specific values.

Note: When using any primary antibody or fluorescence-labelled secondary antibody for the first time, titrate out the antibody to determine which dilution allows the strongest specific signal with the lowest

background for your sample.

Purity Mouse monoclonal immunoglobulins IgG2b subclass were purified by protein A affinity chromatography

and stabilized with 0.05% of glycerol.

Form Liquid. Supplied in 100mM sodium citrate, 50mM Tris and 0.05% v/v glycerol. Neutral pH.

Storage instructions Shipped at -20°C When stored at -20°C, the antibody is stable for 12 months.

Note: Avoid repeated freezing and thawing cycles. It is recommended aliquoting the product upon

arrival.

ReferencesClinical and Analytical Performance of an Automated Serological Test That Identifies S1/S2-Neutralizing

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