ABclonal®

SARS-CoV-2 Spike Rabbit pAb

Catalog No.: A20137 1 Publications

Basic Information

Observed MW

110kDa/180kDa

Calculated MW

141kDa

Category

Polyclonal Antibody

Applications

WB,IF/ICC,IP,ELISA,DB

Cross-Reactivity

SARS-CoV-2

Background

Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) is an enveloped, positive-sense, single-stranded RNA virus that causes coronavirus disease 2019 (COVID-19). Virus particles include the RNA genetic material and structural proteins needed for invasion of host cells. Once inside the cell the infecting RNA is used to encode structural proteins that make up virus particles, nonstructural proteins that direct virus assembly, transcription, replication and host control and accessory proteins whose function has not been determined.~ The structural proteins of SARS-CoV-2 include the envelope protein (E), spike or surface glycoprotein (S), membrane protein (M) and the nucleocapsid protein (N). The spike glycoprotein is found on the outside of the virus particle and gives coronavirus viruses their crown-like appearance. This glycoprotein mediates attachment of the virus particle and entry into the host cell. S protein is an important target for vaccine development, antibody therapies and diagnostic antigen-based tests.

Recommended Dilutions

WB 1:500 - 1:1000

DB 1:500 - 1:2000

IF/ICC 1:50 - 1:200

IP 0.5μg-4μg antibody for

200µg-400µg extracts

of whole cells

ELISA Recommended starting

concentration is 1 µg/mL. Please optimize the concentration based on your specific

assay requirements.

Contact

www.abclonal.com

Immunogen Information

Gene ID43740568

Swiss Prot
PODTC2

Immunogen

Synthetic peptide. This information is considered to be commercially sensitive.

Synonyms

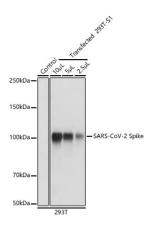
spike glycoprotein; SARS-CoV-2 Spike

Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

Storage

Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.05% proclin300,50% glycerol,pH7.3.



Western blot analysis of extracts of 293T and transfected 293T-S1 (His-tag), using SARS-CoV-2 Spike Rabbit pAb (A20137) at 1:1000 dilution.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000

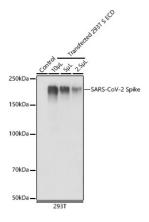
dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 1s.



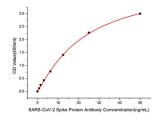
Western blot analysis of extracts of 293T and transfected 293T-S ECD (His-tag), using SARS-CoV-2 Spike Rabbit pAb (A20137) at 1:1000 dilution.

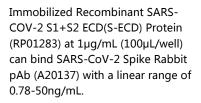
Lysates/proteins: 25µg per lane.

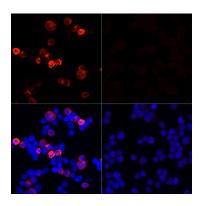
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 1s.

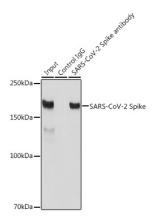






Immunofluorescence analysis of 293T cells transfected with SARS-CoV-2 Spike protein and untreated 293T cells use SARS-CoV-2 Spike Rabbit pAb (A20137) at dilution of 1:100 (40x lens). Secondary antibody: Cy3-conjugated Goat anti-Rabbit IgG (H+L) (AS007) at 1:500 dilution. Blue: DAPI for nuclear staining.

Validation Data



Immunoprecipitation analysis of 300 μ g extracts of 293T cells using 3 μ g SARS-CoV-2 Spike antibody (A20137). Western blot was performed from the immunoprecipitate using SARS-CoV-2 Spike antibody (A20137) at a dilution of 1:1000.