

Product Datasheet

SARS-CoV-2 (COVID-19) Spike Antibody (orb1239976)

Catalog Number	orb1239976
Category	Antibodies
Description	SARS-CoV-2 (COVID-19) Spike Antibody
Target	S
Clonality	Polyclonal
Species/Host	Rabbit
Isotype	IgG
Conjugation	Unconjugated
Reactivity	Virus
Form/Appearance	Liquid
Concentration	1 mg/mL
Buffer/Preservatives	SARS-CoV-2 (COVID-19) Spike Antibody is supplied in PBS containing 0.02% sodium azide.
Purification	SARS-CoV-2 (COVID-19) Spike Antibody is affinity chromatography purified via peptide column.
Immunogen	Anti-SARS-CoV-2 (COVID-19) Spike antibody (orb1239976) was raised against a peptide corresponding to 20 amino acids near the carboxy terminus of SARS-CoV-2 (COVID-19) Spike glycoprotein. The immunogen is located within the last 50 amino acids of SARS-CoV-2 (COVID-19) Spike protein.

Biorbyt Ltd.

7 Signet Court, Swann Road
Cambridge
CB5 8LA
United Kingdom

Email: info@biorbyt.com, support@biorbyt.com

Phone: [+44 \(0\)1223 859353](tel:+44(0)1223859353) | Fax: [+1 \(415\) 651-8558](tel:+1(415)651-8558)

Biorbyt LLC

68 TW Alexander Drive
Research Triangle Park
Durham
NC 27713-2847
United States

Email: info@biorbyt.com, support@biorbyt.com

Phone: [+1 \(415\) 906-5211](tel:+1(415)906-5211) | Fax: [+1 \(415\) 651-8558](tel:+1(415)651-8558)

UniProt ID**P0DTC2****Tested applications**

ELISA, IF, IHC, WB

Application notes

WB: 1 µg/mL; IF: 1 µg/mL. IHC: 0.2 µg/mL; Antibody validated: Immunofluorescence and Western blot in human samples. Immunohistochemistry and Immunofluorescence in COVID-19 patient samples. It will detect 4 ng of free peptide at 1 µg/mL. The immunogen for this is within the last 50 aa of the spike protein - a peptide corresponding to 20 amino acids near the carboxy terminus of SARS-CoV-2 (COVID-19) Spike glycoprotein. The Extracellular domain (ECD) is from aa 1 to 1208 (full length 1273aa). Therefore, this antibody detects the transmembrane and cytoplasm domains at the C terminus, but does not detect the ECD (which is the region expressed in many commercially available spike proteins). This antibody can be used for the detection of full length spike protein and spike protein in COVID-19 patient samples. All other applications and species not yet tested.

Antibody Type

Primary Antibody

Modifications

None

Storage

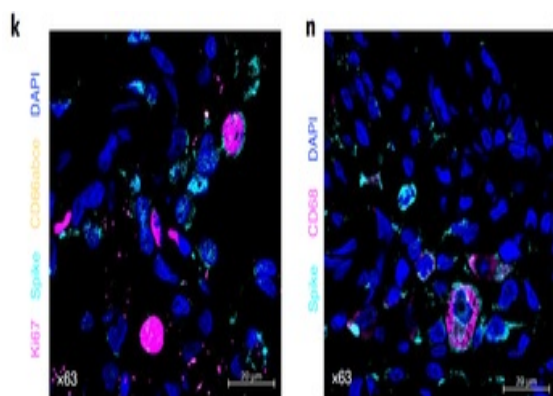
Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Note

For research use only

NCBI**QHD43416****Expiration Date**

12 months from date of receipt.



Immunofluorescent Validation of orb1239976 in SARS-CoV-2 Infected Lung Tissue (Singh et al., Nature Microbiology, 2021). Multilabel confocal immunofluorescence microscopy of formalin-fixed paraffin-embedded lung sections from rhesus macaques infected with SARS-CoV-2. SARS-CoV-2 spike-specific antibodies, orb1239976 (k, n) (turquoise) ; Ki67 (magenta) and neutrophil marker CD66abce (yellow) (k) ; pan-macrophage marker CD68 (magenta) (n) and DAPI (blue).

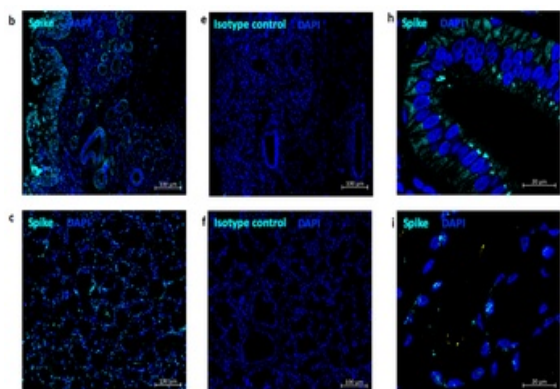
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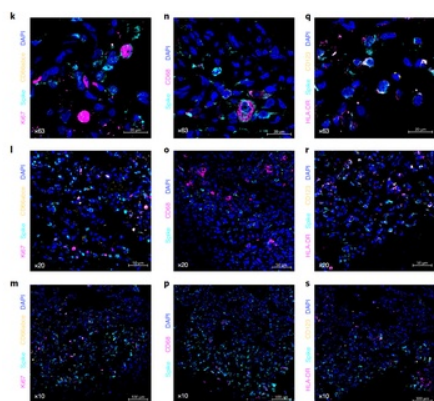
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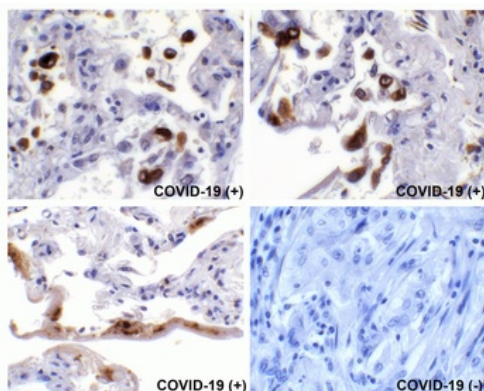
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Immunofluorescent Validation of orb1239976 in SARS-CoV-2 Infected Nose and Tonsil (Singh et al., Nature Microbiology, 2021). Multi-label confocal immunofluorescence microscopy of nasal epithelium (20X-b, 63Xh) and tonsil (20X-c, 63X-i) from rhesus macaques infected with SARS-CoV-2 with SARS-CoV-2 spike-specific antibodies, orb1239976 (turquoise), DAPI (blue). Rabbit IgG isotype control antibody was used to stain the tissues to rule out any non-specific staining (e, f).



Immunofluorescent Validation of orb1239976 in SARS-CoV-2 Infected Lung Tissue (Singh et al., Nature Microbiology, 2021). Multilabel confocal immunofluorescence microscopy of formalin-fixed paraffin-embedded lung sections from rhesus macaques infected with SARS-CoV-2. SARS-CoV-2 spike-specific antibodies, orb1239976 (turquoise) ; Ki67 (magenta) and neutrophil marker CD66abce (yellow) (k-m) ; pan-macrophage marker CD68 (magenta) (n-p) ; HLA-DR (magenta) and pDC marker CD123 (yellow) (q-s) and DAPI (blue).



Immunohistochemistry Validation of SARS-CoV-2 (COVID-19) Spike in COVID-19 Patient Lung. Immunohistochemical analysis of paraffin-embedded COVID-19 patient lung tissue using anti-SARS-CoV-2 (COVID-19) Spike S2 antibody (orb1239976, 0.5 µg/mL). Tissue was fixed with formaldehyde and blocked with 10% serum for 1 h at RT; antigen retrieval was by heat mediation with a citrate buffer (pH6). Samples were incubated with primary antibody overnight at 4 °C. A goat anti-rabbit IgG H&L (HRP) at 1/250 was used as secondary. Counter stained with Hematoxylin. Strong spike protein signal was observed in macrophages and airway epithelium of COVID-19 patient lung, but not in non-COVID-19 patient lung.

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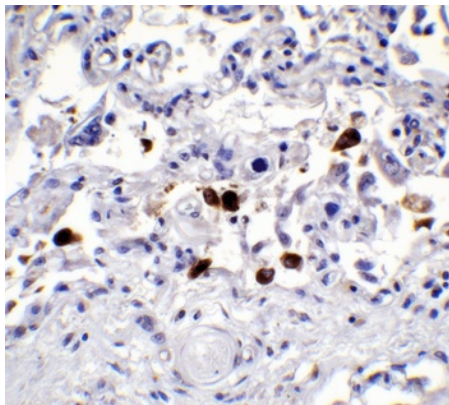
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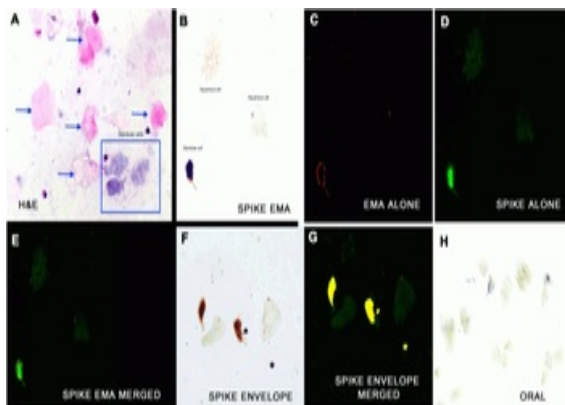
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Immunohistochemistry Validation of SARS-CoV-2 (COVID-19) Spike in COVID-19 Patient Lung. Immunohistochemical analysis of paraffin-embedded COVID-19 patient lung tissue using anti-SARS-CoV-2 (COVID-19) Spike S2 antibody (orb1239976, 0.5 $\mu\text{g}/\text{mL}$). Tissue was fixed with formaldehyde and blocked with 10% serum for 1 h at RT; antigen retrieval was by heat mediation with a citrate buffer (pH6). Samples were incubated with primary antibody overnight at 4 $^{\circ}\text{C}$. A goat anti-rabbit IgG H&L (HRP) at 1/250 was used as secondary. Counter stained with Hematoxylin. Strong spike protein signal was observed in macrophages of COVID-19 patient lung.



IHC/IF Validation in COVID-19 Patient Sample. (Nuovo et al., 2020) Detection of SARS-CoV-2 proteins in nasopharyngeal swab cell preparations. B. An intense signal for covid-19 spike protein tested by SARS-CoV-2 spike antibodies (orb1239976) was observed in the glandular cells. F-H. Co-expression of spike detected by spike antibodies (orb1239976, 0.2 $\mu\text{g}/\text{mL}$) and envelope proteins detected by envelope antibodies (orb1239971, 2 $\mu\text{g}/\text{mL}$) of SARS-CoV-2 (F) documented localization of each protein to glandular cells (G, yellow). No signal was seen in oral swabs of positive cases (H). Both the spike and envelope protein detected by anti-spike antibodies (orb1239976) and anti-envelope antibodies (orb1239971) produced a signal in the nasopharyngeal swabs of the three cases and no signal was evident in the nasopharyngeal swabs of the seven controls.

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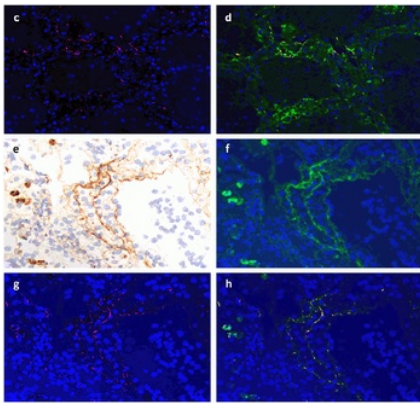
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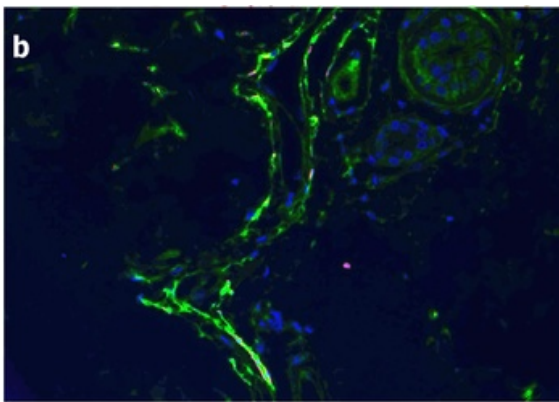
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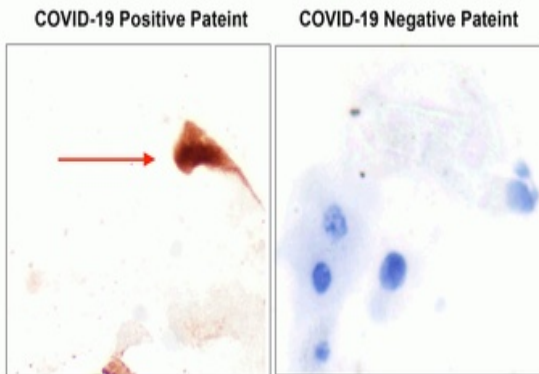
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IF Validation of SARS-CoV2 Spike in COVID-19 Patient Lung. (Magro et al., 2020) SARS-CoV2 spike protein (red, panel C) detected detected by anti-spike antibodies (orb1239976, 0.2 µg/mL) colocalized with C4d (green in panel d, merged in yellow). Spike protein (red, panel g) was also colocalized with C5b-9 (green in panel f&h, merged in yellow).



IF Validation of SARS-CoV2 Spike in COVID-19 Patient Skin. (Magro et al., 2020) C4d is highlighted green while COVID-19 spike protein detected by anti-spike antibodies (orb1239976, 0.2 µg/mL) shows a red staining pattern; a yellow signal is discernible indicative of co-localization of C4d and viral protein within the microvasculature.



IHC Validation of SARS-CoV2 Spike in the Nasopharyngeal Swab Sample of the COVID-19 Patient. Strong spike signal was detected by anti-spike antibodies (orb1239976, 0.2 µg/mL) in the nasopharyngeal swab sample of the COVID-19 patient and no spike signal was observed in the sample of the COVID-19 negative patient. COVID-19 cases were confirmed by PCR.

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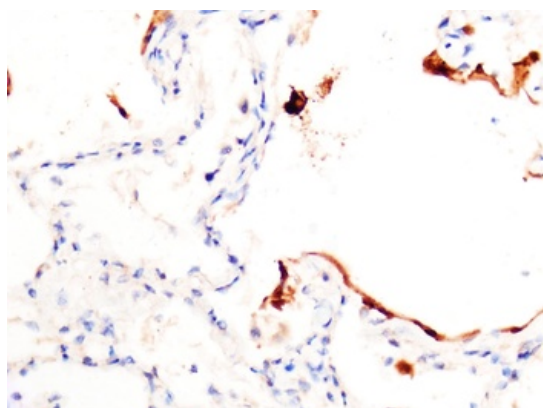
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Cambridge
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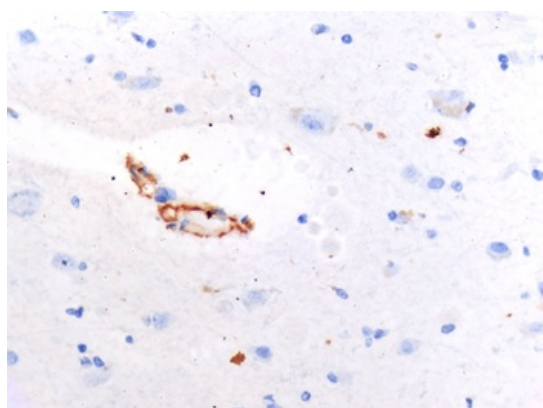
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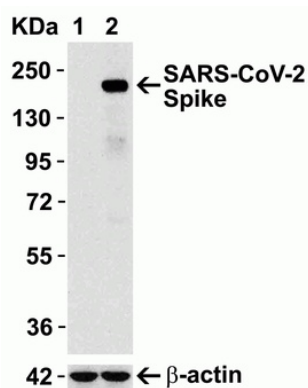
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IHC Validation of SARS-CoV2 Spike in COVID-19 Patient Lung. Strong spike signal was detected by anti-spike antibodies (orb1239976, 0.2 µg/mL) in the lung of the COVID-19 patient confirmed by PCR.



IHC Validation of SARS-CoV2 Spike in COVID-19 Patient Brain. Spike protein was detected by anti-spike antibodies (orb1239976, 0.2 µg/mL) in the brain of the COVID-19 patient confirmed by PCR.



Overexpression Validation in Spike Transfected 293 Cells. Loading: 15 µg per lane of 293 cell lysate. Antibodies: SARS-CoV-2 (COVID-19) Spike, orb1239976 (1 µg/mL), 1h incubation at RT in 5% NFDM/TBST. Secondary: Goat anti-rabbit IgG HRP conjugate at 1:10000 dilution. Lane 1: WT 293 cells and Lane 2: SARS-CoV-2 Spike overexpressed 293 cells.

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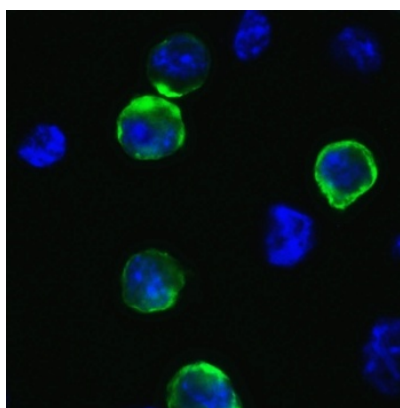
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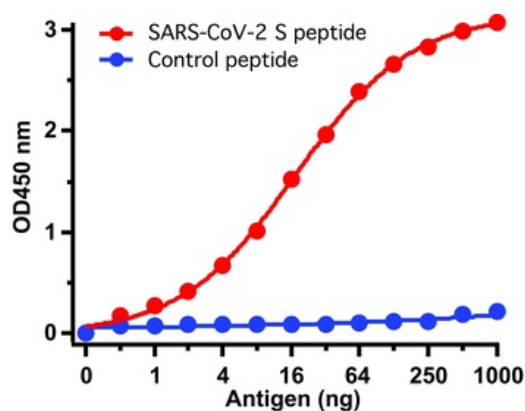
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Research Triangle Park
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Immunofluorescence Validation of SARS-CoV-2 (COVID-19) Spike in 293T Cells. Immunofluorescent analysis of 4% paraformaldehyde-fixed 293T cells labeling SARS-CoV-2 (COVID-19) Spike with orb1239976 at 1 $\mu\text{g}/\text{mL}$, followed by goat anti-rabbit IgG secondary antibody at 1/500 dilution (green) and DAPI staining (blue).



ELISA Test. Antibodies: SARS-CoV-2 (COVID-19) Spike antibody, orb1239976 (1 $\mu\text{g}/\text{mL}$). A direct ELISA was performed using immunogen or control peptide as coating antigen and the anti-SARS-CoV-2 (COVID-19) Spike antibody as the capture antibody. Secondary: Goat anti-rabbit IgG HRP conjugate at 1:20000 dilution. Detection range is from 0.5 ng/mL to 1000 ng/mL .

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Cambridge
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