## **Product Sheet**





RBD

S1

S2

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## **SARS-CoV-2 spike protein**

Catalogue no.: Q104c Clone name: MCV-4C3

**Product:** VHH directed against SARS-CoV-2 spike protein

Target: The disease COVID-19 is caused by the virus SARS-CoV-2 and responsible for the

global pandemic starting in 2020. SARS-CoV-2 is a spherical-shaped positive-strand RNA virus. The SARS-CoV-2 spike protein (S protein) is a homotrimeric transmembrane glycoprotein that is one of the major protein complexes on the virus and which plays an important role in infection into host cells. Each spike protein monomer is a 140 kDa protein with an N-terminal S1 domain, a membrane-proximal S2 domain, a transmembrane domain, and a C-terminal domain. Via the receptor binding domain (RBD) within the S1 domain, the spike proteins bind to Angiotensin-Converting Enzyme 2 (ACE2) receptors on host cells, which is then followed by fusion of the virus with the membrane. By interfering with the interaction of the RBD with ACE2, infection can be blocked.

S1 and RBD in particular, is an interesting therapeutic target for COVID-19.<sup>3</sup>

Recombinant monoclonal VHH (Llama glama), purified from S.cerevisiae using affinity chromatography. Immunization with and phage-display

selection on recombinant protein using total elution.

**Specificity:** SARS-CoV-2 spike protein domain S1.

**Formulation:** 0.2 μm filtered solution in PBS.

**Mol. Weight:** 14.1 kDa **Ext. Coeff. (ε):** 21555 M<sup>-1</sup> cm<sup>-1</sup>

**A<sub>280</sub> at 1g/L:** 1.5

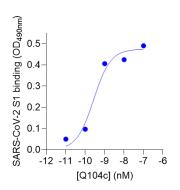
**Storage:** Shipped on blue ice. Store at 4°C or -20°C (aliquots). Addition of 0.02%

sodiumazide is optional.

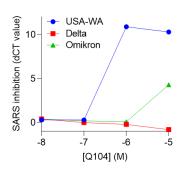
**Applications:** ELISA, viral neutralisation.

**Examples:** 

Source:



Binding of Q104c to recombinant SARS-CoV-2 spike protein in ELISA.



Inhibition of SARS-CoV-2 infection by Q104.

## **References:**

- 1 Sharma et al., (2021) Viruses. 13(2):202
- 2 Khailany et al., (2020) Gene Reports. 100682
- 3 Walls et al., (2020) Cell. 180:281-292
- 4 Salvatori et al., (2020) J Transl Med 18:222