Validation data for Spike-S1-Fc

https://www.invivogen.com/sars2-spike-s1-proteins

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Version 20109-NJ

Spike-S1-Fc is a soluble SARS-CoV-2 fusion protein generated by fusing the full-length Spike S1 subunit (V16-R685) to a C-terminal human IgG1-Fc tag with a TEV (Tobacco Etch Virus) sequence linker. This fusion protein has a molecular weight of ~124kDa on a SDS PAGE gel (Figure 1). The recognition of the Spike-S1-Fc protein by an Anti-SARS-CoV-Spike human IgM (clone CR3022) has been verified by ELISA (Figure 2).

Spike-S1-Fc purity analysis by SDS PAGE

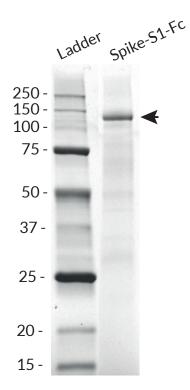


Figure 1: SDS PAGE analysis of the SARS-CoV-2 Spike-S1-Fc protein. 2 μg of the fusion protein was loaded onto a 12% Mini-PROTEAN® TGX Stain-Free™ Precast Gel (Bio-Rad). Detection was performed as per manufacturer's instructions.

Recognition of Spike-S1-Fc by an Anti-SARS-CoV-Spike (CR3022) human IgM

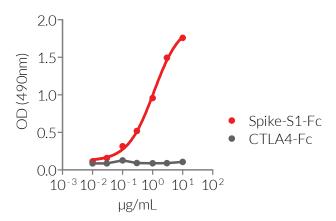


Figure 2: ELISA detection of Spike-S1-Fc fusion protein with an Anti-SARS-CoV spike human IgM (CR3022). Anti-SARS-CoV-Spike hIgM antibody (5 µg/ml) was coated onto ELISA plates overnight. Following this, a 3-fold serial dilution of Spike-S1-Fc (red curve) or control protein (CTLA4-Fc; grey curve) were added and incubated for 1 hour. Binding was detected using a HRP-labelled anti-His antibody (1/1000 dilution) and the HRP substrate OPD (o-phenylenediamine dihydrochloride). Absorbance was read at 490 nm.

