

# **Product Datasheet**

# Human ACE2/ACEH Protein. Fc Tag (orb1945965)

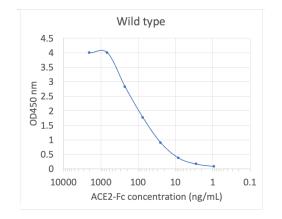
Catalog Number	orb1945965
Description	Human ACE2/ACEH Protein. Fc Tag
Reactivity	Human
Tested Applications	ELISA
Preservatives	1X PBS, 0.09% NaN3
Concentration	0.5 mg/ml
Storage	2-8°C
Tag	Fc tag
Note	For research use only
lsotype	Other
Clone Number	ACE2
Dilution Range	For SARS-CoV-2 (COVID-19) diagnostic assays.
Expiration Date	6 months from date of receipt.

### **Biorbyt Ltd.**

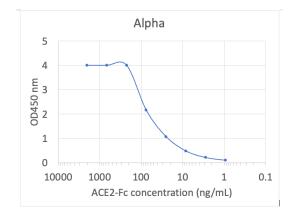
7 Signet Court, Swann's Road, Cambridge, CB5 8LA, United Kingdom Email: <u>info@biorbyt.com</u>, <u>support@biorbyt.com</u> Phone: +44 (0) 1223 859-353 | Fax: +1 (415) 651-8558

### **Biorbyt LLC.**





Microtiter wells were coated with 100 uL of each spike trimer at 2 ug/mL in PBS at 4?C overnight. The wells were washed with PBS and blocked with 200  $\mu$ L of 1% BSA/PBS. ACE2-Fc was serially diluted from 2  $\mu$ g/mL in 1% BSA/PBS. The blocker was discarded, and the wells were incubated with 100  $\mu$ L of serially diluted ACE2-Fc at 37?C for 1 hour. The wells were washed with PBS and the bound ACE2-Fc was detected with 100  $\mu$ L of Peroxidase AffiniPure Goat Anti-Human IgG, Fc? fragment specific (1:5, 000 in 1% BSA/PBS) at 37?C for 1 hour. The wells were washed with PBS and the wells were developed with 100  $\mu$ L of MB/E Ultra Sensitive, Blue, Horseradish Peroxidase Substrate at RT for 5 min. The reaction was stopped with 100  $\mu$ L of 0.6N H2SO4 and the signals were read at 450 nm using a plate reader.



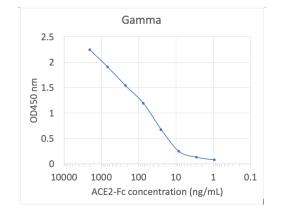
Microtiter wells were coated with 100 uL of each spike trimer at 2 ug/mL in PBS at 4?C overnight. The wells were washed with PBS and blocked with 200  $\mu$ L of 1% BSA/PBS. ACE2-Fc was serially diluted from 2  $\mu$ g/mL in 1% BSA/PBS. The blocker was discarded, and the wells were incubated with 100  $\mu$ L of serially diluted ACE2-Fc at 37?C for 1 hour. The wells were washed with PBS and the bound ACE2-Fc was detected with 100  $\mu$ L of Peroxidase AffiniPure Goat Anti-Human IgG, Fc $\gamma$  fragment specific (1:5, 000 in 1% BSA/PBS) at 37?C for 1 hour. The wells were washed with PDS and the wells were developed with 100  $\mu$ L of MB/E Ultra Sensitive, Blue, Horseradish Peroxidase Substrate at RT for 5 min. The reaction was stopped with 100  $\mu$ L of 0.6N H2SO4 and the signals were read at 450 nm using a plate reader.

#### **Biorbyt Ltd.**

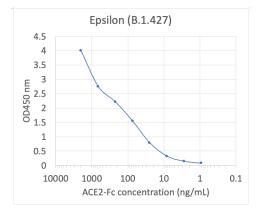
7 Signet Court, Swann's Road, Cambridge, CB5 8LA, United Kingdom Email: <u>info@biorbyt.com</u>, <u>support@biorbyt.com</u> Phone: +44 (0) 1223 859-353 | Fax: +1 (415) 651-8558

#### **Biorbyt LLC.**





Microtiter wells were coated with 100 uL of each spike trimer at 2 ug/mL in PBS at 4?C overnight. The wells were washed with PBS and blocked with 200  $\mu$ L of 1% BSA/PBS. ACE2-Fc was serially diluted from 2  $\mu$ g/mL in 1% BSA/PBS. The blocker was discarded, and the wells were incubated with 100  $\mu$ L of serially diluted ACE2-Fc at 37?C for 1 hour. The wells were washed with PBS and the bound ACE2-Fc was detected with 100  $\mu$ L of Peroxidase AffiniPure Goat Anti-Human IgG, Fc? fragment specific (1:5, 000 in 1% BSA/PBS) at 37?C for 1 hour. The wells were washed with PDS and the wells were developed with 100  $\mu$ L of MB/E Ultra Sensitive, Blue, Horseradish Peroxidase Substrate at RT for 5 min. The reaction was stopped with 100  $\mu$ L of 0.6N H2SO4 and the signals were read at 450 nm using a plate reader.



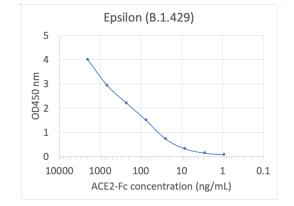
Microtiter wells were coated with 100 uL of each spike trimer at 2 ug/mL in PBS at 4?C overnight. The wells were washed with PBS and blocked with 200  $\mu$ L of 1% BSA/PBS. ACE2-Fc was serially diluted from 2  $\mu$ g/mL in 1% BSA/PBS. The blocker was discarded, and the wells were incubated with 100  $\mu$ L of serially diluted ACE2-Fc at 37?C for 1 hour. The wells were washed with PBS and the bound ACE2-Fc was detected with 100  $\mu$ L of Peroxidase AffiniPure Goat Anti-Human IgG, Fc? fragment specific (1:5, 000 in 1% BSA/PBS) at 37?C for 1 hour. The wells were washed with PDS and the wells were developed with 100  $\mu$ L of MB/E Ultra Sensitive, Blue, Horseradish Peroxidase Substrate at RT for 5 min. The reaction was stopped with 100  $\mu$ L of 0.6N H2SO4 and the signals were read at 450 nm using a plate reader.

#### **Biorbyt Ltd.**

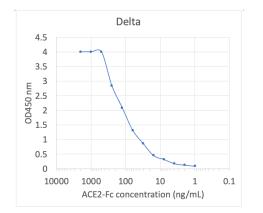
7 Signet Court, Swann's Road, Cambridge, CB5 8LA, United Kingdom Email: <u>info@biorbyt.com</u>, <u>support@biorbyt.com</u> Phone: +44 (0) 1223 859-353 | Fax: +1 (415) 651-8558

#### **Biorbyt LLC.**





Microtiter wells were coated with 100 uL of each spike trimer at 2 ug/mL in PBS at 4?C overnight. The wells were washed with PBS and blocked with 200  $\mu$ L of 1% BSA/PBS. ACE2-Fc was serially diluted from 2  $\mu$ g/mL in 1% BSA/PBS. The blocker was discarded, and the wells were incubated with 100  $\mu$ L of serially diluted ACE2-Fc at 37?C for 1 hour. The wells were washed with PBS and the bound ACE2-Fc was detected with 100  $\mu$ L of Peroxidase AffiniPure Goat Anti-Human IgG, Fc? fragment specific (1:5, 000 in 1% BSA/PBS) at 37?C for 1 hour. The wells were washed with PDS and the wells were developed with 100  $\mu$ L of MB/E Ultra Sensitive, Blue, Horseradish Peroxidase Substrate at RT for 5 min. The reaction was stopped with 100  $\mu$ L of 0.6N H2SO4 and the signals were read at 450 nm using a plate reader.



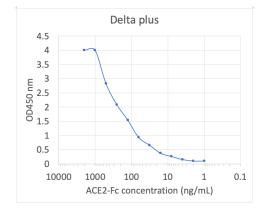
Microtiter wells were coated with 100 uL of each spike trimer at 2 ug/mL in PBS at 4?C overnight. The wells were washed with PBS and blocked with 200  $\mu$ L of 1% BSA/PBS. ACE2-Fc was serially diluted from 2  $\mu$ g/mL in 1% BSA/PBS. The blocker was discarded, and the wells were incubated with 100  $\mu$ L of serially diluted ACE2-Fc at 37?C for 1 hour. The wells were washed with PBS and the bound ACE2-Fc was detected with 100  $\mu$ L of Peroxidase AffiniPure Goat Anti-Human IgG, Fc? fragment specific (1:5, 000 in 1% BSA/PBS) at 37?C for 1 hour. The wells were washed with PDS and the wells were developed with 100  $\mu$ L of MB/E Ultra Sensitive, Blue, Horseradish Peroxidase Substrate at RT for 5 min. The reaction was stopped with 100  $\mu$ L of 0.6N H2SO4 and the signals were read at 450 nm using a plate reader.

#### **Biorbyt Ltd.**

7 Signet Court, Swann's Road, Cambridge, CB5 8LA, United Kingdom Email: <u>info@biorbyt.com</u>, <u>support@biorbyt.com</u> Phone: +44 (0) 1223 859-353 | Fax: +1 (415) 651-8558

#### **Biorbyt LLC.**





Microtiter wells were coated with 100 uL of each spike trimer at 2 ug/mL in PBS at 4?C overnight. The wells were washed with PBS and blocked with 200  $\mu$ L of 1% BSA/PBS. ACE2-Fc was serially diluted from 2  $\mu$ g/mL in 1% BSA/PBS. The blocker was discarded, and the wells were incubated with 100  $\mu$ L of serially diluted ACE2-Fc at 37?C for 1 hour. The wells were washed with PBS and the bound ACE2-Fc was detected with 100  $\mu$ L of Peroxidase AffiniPure Goat Anti-Human IgG, Fc? fragment specific (1:5, 000 in 1% BSA/PBS) at 37?C for 1 hour. The wells were washed with PDS and the wells were developed with 100  $\mu$ L of MB/E Ultra Sensitive, Blue, Horseradish Peroxidase Substrate at RT for 5 min. The reaction was stopped with 100  $\mu$ L of 0.6N H2SO4 and the signals were read at 450 nm using a plate reader.

#### **Biorbyt Ltd.**

7 Signet Court, Swann's Road, Cambridge, CB5 8LA, United Kingdom Email: <u>info@biorbyt.com</u>, <u>support@biorbyt.com</u> Phone: +44 (0) 1223 859-353 | Fax: +1 (415) 651-8558

#### **Biorbyt LLC.**