

A

gRBD-Fc
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 DLCTFTNVTADSFVIRGDEVQRQIAPGQGTGIADYNYKLPDNFTGCVIAWNSNLDKSVGGNYNYLRFKRSNLRKPF
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 TVLQDMLNGKEYKCKVKNALPAPEIKETISAKCGPRPEQVTLPPSRDELTKNQVSLTCLVGGFTYPSDIAVEWES
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gRBD-foldon
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NAP-gRBD
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 RDISTEIQAGSTPCNGVEGFNCYFPLQSYGFPQPTNGVGYQYRVVVLSENLTAAPATVCGPGSSGGSGPGGIIP
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gRBD-ferritin
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 RDISTEIQAGSTPCNGVEGFNCYFPLQSYGFPQPTNGVGYQYRVVVLSENLTAAPATVCGPGSSGGSGPGGIIP
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gRBD-m13
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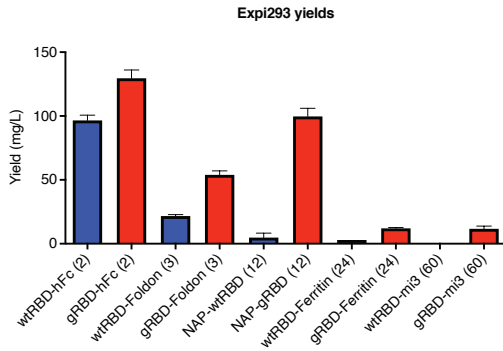
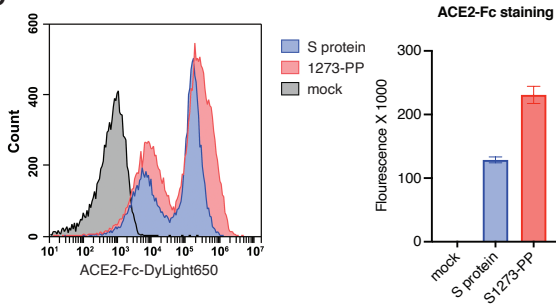
B**C**

Fig. S4. Multivalent gRBD fusion proteins express more efficiently than their wtRBD counterparts. (A) Complete sequences of the multivalent gRBD fusion constructs used in these studies. Green indicated signal peptide; black, linker residues; blue, multivalent carrier protein; red, gRBD; purple, affinity tag. (B) Yields of purified wtRBD and gRBD multimers expressed from the CMV/R vector in Expi293 cells. Values reflect a minimum of two independent transfections. Error bars indicate s.d. (C) Flow cytometry of 293T cells transiently transfected with SARS-CoV2 S protein in the pCAGGS vector, or S1273-PP in the CMVR vector. Expression was measured by staining with ACE2-Fc-DyLight650. Right panel indicates mean fluorescence intensity as determine from the histogram shown in the left panel.