Supplementary Information 4 Supplementary Shelf-life Stability Data

SDS-PAGE for RBD-I53-50 nanoparticle



The integrity of samples after incubation at four temperatures over a 28 day (D) study was analyzed by SDS-PAGE. Molecular weights of the standard are noted in kDa. Each sample was analyzed +/-reducing agent (DTT).

35-40°C

22-27°C

hACE2-Fc binding for RBD-I53-50 nanoparticle



hACE2-Fc binding of antigen incubated at four different temperatures for 28 days (D) was analyzed by bio-layer Interferometry (BLI). Protein A biosensors loaded with hACE2-Fc were incubated with immunogen (association, x = 590-889 s) and then buffer (dissociation, x = 890-1190 s).

CR3022 binding for RBD-I53-50 nanoparticle



CR3022 IgG binding of antigen incubated at four different temperatures for 28 days (D) was analyzed by bio-layer Interferometry (BLI). Protein A biosensors loaded with CR3022 were incubated with immunogen (association, x = 590-889 s) and then buffer (dissociation, x = 890-1190 s).

nsEM for RBD-I53-50 nanoparticle



Representative negative stain electron micrographs for each sample at days (D) 1 and 28 following incubation at four temperatures. Scale bar, 50 nm.

Dynamic light scattering for RBD-I53-50 nanoparticle



SDS-PAGE for Rpk4-I53-50 nanoparticle



The integrity of samples after incubation at four temperatures over a 28 day (D) study was analyzed by SDS-PAGE. Molecular weights of the standard are noted in kDa. Each sample was analyzed +/- reducing agent (DTT).

2-8°C

22 - 27

°C

-80°C

35-40°C

hACE2-Fc binding for Rpk4-I53-50 nanoparticle



hACE2-Fc binding of antigen incubated at four different temperatures for 28 days (D) was analyzed by bio-layer Interferometry (BLI). Protein A biosensors loaded with hACE2-Fc were incubated with immunogen (association, x = 590-889 s) and then buffer (dissociation, x = 890-1190 s).

CR3022 binding for Rpk4-I53-50 nanoparticle



CR3022 IgG binding of antigen incubated at four different temperatures for 28 days (D) was analyzed by bio-layer Interferometry (BLI). Protein A biosensors loaded with CR3022 were incubated with immunogen (association, x = 590-889 s) and then buffer (dissociation, x = 890-1190 s).

nsEM for Rpk4-I53-50 nanoparticle



Representative negative stain electron micrographs for each sample at days (D) 1 and 28 following incubation at four temperatures. Scale bar, 50 nm.

Dynamic light scattering for Rpk4-I53-50 nanoparticle



SDS-PAGE for Rpk9-I53-50 nanoparticle



hACE2-Fc binding for Rpk9-I53-50 nanoparticle



hACE2-Fc binding of antigen incubated at four different temperatures for 28 days (D) was analyzed by bio-layer Interferometry (BLI). Protein A biosensors loaded with hACE2-Fc were incubated with immunogen (association, x = 590-889 s) and then buffer (dissociation, x = 890-1190 s).

CR3022 binding for Rpk9-I53-50 nanoparticle



CR3022 IgG binding of antigen incubated at four different temperatures for 28 days (D) was analyzed by bio-layer Interferometry (BLI). Protein A biosensors loaded with CR3022 were incubated with immunogen (association, x = 590-889 s) and then buffer (dissociation, x = 890-1190 s).

nsEM for Rpk9-I53-50 nanoparticle



Representative negative stain electron micrographs for each sample at days (D) 1 and 28 following incubation at four temperatures. Scale bar, 50 nm.

Dynamic light scattering for Rpk9-I53-50 nanoparticle

