SARS-CoV-2 spike trimer vaccine expressed in *Nicotiana benthamiana* adjuvanted with Alum elicits protective immune responses in mice

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**Summary**
The ongoing coronavirus disease 2019 (COVID-19) pandemic has spurred rapid development of vaccines as part of the public health response. However, the general strategy used to construct recombinant trimeric severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) spike (S) proteins in mammalian cells is not completely adaptive to molecular farming. Therefore, we generated several constructs of recombinant S proteins for high expression in *Nicotiana benthamiana*. Intramuscular injection of *N. benthamiana*-expressed S protein vaccine (NSctVac) into Balb/c mice elicited both humoral and cellular immune responses, and booster doses increased neutralizing antibody titres. In human angiotensin-converting enzyme knock-in mice, two doses of NSctVac induced anti-S and neutralizing antibodies, which cross-neutralized Alpha, Beta, Delta and Omicron variants. Survival rates after lethal challenge with SARS-CoV-2 were up to 80%, without significant body weight loss, and viral titres in lung tissue fell rapidly, with no infectious virus detectable at 7-day post-infection. Thus, plant-derived NSctVac could be a candidate COVID-19 vaccine.

**Introduction**
Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) is an enveloped virus that contains a single-stranded RNA genome of 29 903 bp, encoding 16 non-structural proteins, eight accessory proteins and four structural proteins (Mittal et al., 2020; Shang et al., 2020; Yan et al., 2020). Coronaviruses infect a wide range of hosts, including humans. The host specificity is determined by the spike (S) protein, a surface glycoprotein that forms trimers. The S protein can be divided into two functional subunits, S1 and S2, which are separated by proteolytic cleavage by furin or furin-like host proteases (Ord et al., 2020; Peacock et al., 2021). Each of these subunits comprises multiple functional domains. The S1 subunit contains the receptor-binding domain (RBD), which is responsible for binding to a specific protein (or receptor) on the surface of the host cell. Thus, the S1 subunit determines host range and cell tropism (Hossain et al., 2022; Yuan et al., 2021). The S2 subunit, which also contains multiple domains (i.e. a fusion peptide and two heptad repeats), undergoes a dramatic conformational change to produce a so-called trimer of hairpins that supports fusion between the viral and host cell membranes, followed by entry of the virus into the host cell (Berger and Schaffitzel, 2020; Jackson et al., 2022; Xia et al., 2020).

During the ongoing global coronavirus disease 2019 (COVID-19) pandemic, the imperative is to prevent the spread of SARS-CoV-2. Thus, development and production of large amounts of vaccines that are effective against SARS-CoV-2 is necessary. Various approaches have been used to develop such vaccines. Currently, several vaccines (including two mRNA-based vaccines: mRNA-1273; Corbett et al., 2020) and BNT162b2 (Vogel et al., 2021) are in use to protect individuals from infection by SARS-CoV-2. These vaccines induce stronger immune responses, with high neutralizing antibody titres than other types of vaccines in most cases (Francis et al., 2022). However, mRNA-based vaccines appear to cause more side effects than conventional vaccines (Castells and Phillips, 2021). There is also a dispute about the potential risk of reverse transcription of mRNA (Aldén et al., 2022; Zhang et al., 2021). The older platforms (such as protein subunit vaccines) are well accepted by authorities and the lay public compared with novel mRNA platforms. Virus-derived vaccines were also introduced in the field (Bos et al., 2020; Watanabe et al., 2021; Wu et al., 2020), although their protective efficacy is slightly less than that of mRNA-based vaccines because they induce weaker immune responses (Francis et al., 2022). Recently, recombinant protein-based vaccines NVX-CoV2373 (Tian et al., 2021), ZF2001 (Ai et al., 2022) and CoVLP (Ward et al., 2021) were introduced into the field. In contrast to mRNA-based vaccines, these recombinant protein-based vaccines have fewer and weaker side effects and are free from concerns about transgenic contamination. However, the disadvantage of protein-based vaccines is that they take a longer time to develop, and
downstream production is more demanding than DNA or RNA-based vaccines.

Here, we aimed to develop and produce new vaccines against SARS-CoV-2 using recombinant proteins produced in N. benthamiana. Recombinant proteins for vaccine development can be produced using multiple approaches. Although proteins can be obtained from inactivated/attenuated viruses, this is not easy because virus amplification requires appropriate cell lines and secure facilities for virus production. Instead, recombinant technology can be used to produce proteins using particle genes obtained from a virus. Traditionally, animal cells or bacteria are used to produce recombinant proteins (Brindha and Kuroda, 2022; Esposito et al., 2020; Gobeil et al., 2021; Li et al., 2020). Recently, however, plants have been used for this purpose because they are easy to grow and production can be scaled up (D'Aoust et al., 2010; Royal et al., 2021; Song et al., 2021). Indeed, using influenza as a model virus, plants have proved to be an excellent system for rapid production of large amounts of recombinant proteins for use as vaccine candidates. Indeed, about 100 million doses of recombinant hemagglutinin (HA) protein were prepared from N. benthamiana within 6 weeks via Agrobacterium-mediated transformation (D'Aoust et al., 2010). Notably, a plant-based VLP vaccine for SARS-CoV-2 was approved by the Canadian government at the beginning of 2022 (Pillet et al., 2022; Ward et al., 2021). The current COVID-19 situation requires rapid, large-scale production of vaccines to meet the global demand. Here, we describe the use of a plant-based system to generate subunit vaccine candidates for use against SARS-CoV-2.

SARS-CoV-2 surface proteins are a good candidate for vaccine development. This is because all individuals that have recovered from COVID-19 generated anti-S1 and anti-RBD antibodies. However, only a small fraction of these antibodies block the binding of the RBD to the human angiotensin-converting enzyme 2 (hACE2 receptor; Chen et al., 2020). The transient and dynamic conformational states of the S protein mean that there is likely a very limited window during which host B cells are exposed to immunogenic epitopes within the RBD (Pallesci et al., 2017). The S protein of SARS-CoV-2 also induces the production of neutralizing antibodies, including antibodies specific for the RBD.

In this study, we used the full-length S protein lacking the C-terminal transmembrane domain (TMD) and cytosolic tail domain (TMD-C) and asked whether it can be used as a vaccine candidate. We provide evidence that the plant-produced near full-length S protein is highly immunogenic in mice and induced production of antibodies that protect against SARS-CoV-2.

Results

Expression and purification of SARS-CoV-2 S protein from N. benthamiana

To identify a vaccine candidate for SARS-CoV-2, we decided to take the recombinant vaccine approach. The S protein expressed on the surface of SARS-CoV-2 is an excellent vaccine candidate compared with its subunit domain RBD or S1, since the S trimer exhibited stronger immunogenicity than that of recombinant S1 protein and RBD protein (Liu et al., 2021). Notably, full-length S protein contains conserved T cell epitopes in the S2 domain (Grifoni et al., 2021), which may contribute to conferring a certain degree of protection against variants. However, production as a recombinant protein is rather challenging due to its large size, trimeric structure, extensive N-glycosylations and structural instability. Therefore, to produce a recombinant S protein in N. benthamiana, we considered various aspects such as expression level, trimerization, structural stability, N-glycosylation and production of a prefusion form. First, to increase expression, we designed a full-length S protein lacking the TMD and C-terminal cytosolic tail domain (S(TMD-C)). Then, we compared the trimerization motifs of the Foldon domain (FD) of bacteriophage T4 fibritin with those of mouse Coronin 1A (mCor1). The FD is a common trimer-inducing domain used for the production of various trimeric recombinant proteins in mammalian cells, including HA (Lu et al., 2014; Santiago et al., 2012; Wei et al., 2008) and S proteins (Esposito et al., 2020; Gobeil et al., 2021). mCor1 is used to induce trimerization of HA (Song et al., 2021). The mode of trimerization between the FD and mCor1 is different, from which the FD domain contains a short N-terminal α-helix structure and a longer C-terminal flexible structure while mCor1 is a complete α-helix structure; therefore, their effect on expression of the fusion protein was unknown. To produce S protein with N-glycosylations, we used the leader sequence of Arabidopsis BiP (ER-localized chaperon) to induce localization in the endoplasmic reticulum (ER; Figure 1a). Additionally, five histidine residues (His5) and an HDEL motif (used as a purification tag and ER retention signal, respectively) were added to the C terminus of the FD or mCor1 to yield the final constructs BiP:SA(TMD-C):FD: His5:HDEL (Sbi) and BiP:SA(TMD-C):mCor1:His5:HDEL (Scor), respectively. First, we produced these proteins in N. benthamiana using the transient expression method (Song et al., 2022). S1α and Sα bands were detected at approximately 180 kDa by Western blot analysis using anti-His antibody (Figure 1c). The apparent size of these proteins was larger than the calculated molecular weight, but similar to that reported in recent studies (Cai et al., 2020; Hsieh et al., 2020). This was due to N-glycosylation since S proteins were expressed in the ER. Consistent with this idea, treatment of purified S proteins with PNGase F caused a shift in migration patterns (Figure 3c). The two trimeric forms, Sα and Sα, showed differences in expression level: the amount of Sα was about 1.8-fold higher than that of Sα according to the result of Western blot analysis (Figure 1b). Therefore, we used mCor1 as a trimerization motif in the following experiments. Next, we introduced a 3P mutation (A942P/K986P/V987P) into the S2 subunit of the Sα construct. These three proline substitutioins stabilize the S protein and increase expression in mammalian cells (Pallesci et al., 2017; Hsieh et al., 2020). Finally, we mutated the furin cleavage site to QQAQ to maintain the prefusion form. As a result, Scor-3P and Scor-3PQQQ were expressed at higher levels than Scor (Figure 1b).

Next, we examined conditions that promote higher-level expression of Scor. A previous study showed that treatment with lipoic acid and ascorbic acid (both antioxidants capable of scavenging reactive oxygen species) along with heat shock treatment led to higher expression in N. benthamiana (Norkunas et al., 2018; Wang et al., 2004; Zhao et al., 2017). Of these, heat shock treatment for 30 min at 1-day post-infection (dpi) greatly increased expression of Scor (Figure 5a and b), suggesting that chaperones induced by heat shock play a critical role in high-level expression of the S protein in N. benthamiana. This result prompted us to examine the effect of chaperones in the ER, that is, does co-expression of ER chaperones lead to higher expression of Scor? A recent study showed that human calreticulin (CRT)
facilitates expression of HIV gp140 in plants (Margolin et al., 2020). We co-infiltrated *N. benthamiana* with a mixture of *Agrobacteria* harbouring *Sct* and CRT. Indeed, co-expression of CRT increased expression of *Sct* by 3.51-fold at the ratios of *Sct* to CRT ranging from 2 : 1 to 5 : 1 (Figure 2a–c). CRT was expressed at the same level regardless of the ratio, indicating that CRT is highly expressed in *N. benthamiana* (Figure 2b). However, heat shock treatment did not further enhance expression of *Sct* in the presence of CRT (Figure 2b). Next, we examined the effect of CRT on expression of *Sct-3P*. These results indicate that co-expression...
of CRT contributes to the folding process of S protein in *N. benthamiana* in a way similar to heat shock treatment (Margolin *et al.*, 2020). However, CRT did not enhance the expression of Sct-3P (Figure 2d), suggesting that the 3P mutation contributes to the folding process which is supported by co-expressed CRT in *N. benthamiana*.

We examined trimerization of Sct at the biochemical level. As a control, we generated a monomeric S protein construct, BiP:SA (TMD-C):His5:HDEL (Sm) lacking mCor1 (Figure 1a). Sm was expressed at the same level as Sct (Figure 1c). Next, we purified both Sct and Sm by Ni²⁺-NTA affinity column chromatography. The eluates were further purified by size exclusion chromatography. Sct eluted earlier than Sm, at a position corresponding to approximately 600 kDa, confirming trimer formation (Figure 1d). To confirm trimerization by mCor1, we generated BiP:SA(TMD-C):mCor1:His5:HDEL (S2ct) and BiP:SA(TMD-C):His5:HDEL (S2m) using the S2 subunit without the TMD and cytosolic domain (Figure S2A). S2ct and S2m were detected at positions corresponding to 83 and 80 kDa, respectively (Figure S2B). Moreover, S2ct also eluted earlier than S2m, consistent with the idea that mCor1 induces trimerization of the S protein (Figure S2C).

As a first step towards vaccine development using S proteins produced in *N. benthamiana*, we used a pull-down assay to examine whether plant-produced Sct and Sm bind to ACE2, the human receptor for SARS-CoV-2. HA-tagged human ACE2 (hACE2-HA) was expressed in *N. benthamiana* together with Sct or Sm. Total protein extracts were prepared and pulled down with Ni²⁺-NTA beads. The pull-down results were analysed by Western blotting using an anti-HA antibody. Sct-HA was detected in the pull-down from Sct and Sm (Figure 1e), indicating that both Sct and Sm co-expressed in plant cells interact with hACE2.

**Purification of SARS-CoV-2 S and S2 from *Nicotiana benthamiana***

Next, we optimized the conditions required to purify Sct and S2ct from protein extracts of *N. benthamiana*. First, we purified Sct and S2ct using Ni²⁺-NTA affinity column chromatography. Here, we first established the purification protocol using total protein extracts prepared from *N. benthamiana* that had been infiltrated with *Agrobacterium* harbouring Sct and P38. These proteins contained six histidine residues at the C terminus. Total soluble protein extracts were prepared in TBS buffer (100 mM NaCl, 25 mM Tris–HCl, pH 8, 0.1% Tween 20 and 0.5 mM protease inhibitor cocktail) and applied to the Ni²⁺-NTA affinity column. The eluates were subjected to further purification by size exclusion chromatography. These samples were analysed by SDS-PAGE and stained with CBB (Figure 3a, Figure S3A). Sct recombinant proteins were purified using Ni²⁺-NTA affinity column chromatography followed by SEC as established above in Figure 3. Both proteins were diluted to a final concentration of 0.5–0.7 μg/μL (Purity > 95%) for use in animal experiments. Before using Sct for immunization, we also examined the stability of Sct. E-tubes containing 5 μg of purified Sct were kept at 4°C up to 5 days, which were stored at −80°C or used for SDS-PAGE analysis. Plant-produced Sct can be stored at 4°C for 2 days without loss of quality (Figure 3d).

Sct and S2ct produced from *N. benthamiana* form trimers

The behaviour of Sct upon size exclusion chromatography agreed with the expectation of trimer formation. To confirm this, we examined the morphology of the proteins by electron microscopy (EM) after negative staining. Sct and S2ct obtained from a Ni²⁺-NTA affinity column (Figure 3a, Figure S2D) were further purified.
by size exclusion chromatography (SEC), the peak fractions were collected, and the proteins were negatively stained. The negative EM images yielded both side and top views of the trimeric S protein particles (Figure 3b, Figure S2E), similar to earlier studies (Cai et al., 2020; Hsieh et al., 2020). These results confirmed that Sct and S2ct are produced as trimers in N. benthamiana.
Figure 2  Co-expression of human calreticulin greatly enhances the expression of S$_{ct}$ but not S$_{ct}$-3P. (a) Schematic representation of the recombinant CRT construct. BiP, the ER targeting leader sequence of Arabidopsis BiP1; CRT, human CRT from aa positions 18–414; HA, the small hemagglutinin tag; KDEL, the ER retention motif; 35S, CaMV 35S promoter; HSP, Arabidopsis HSP terminator. (b, d) Effect of CRT co-expression on S$_{ct}$ or S$_{ct}$-3P expression. Agrobacteria harbouring S$_{ct}$, S$_{ct}$-3P, or CRT were cultured in LB medium and resuspended in infiltration buffer at a concentration of OD$_{600}$ = 0.8. S$_{ct}$ or S$_{ct}$-3P-harbouring Agrobacterium resuspensions were mixed with CRT-harbouring Agrobacterium resuspension at ratios from 1 : 0 to 5 : 1 and used for syringe infiltration. Leaf tissues were harvested at 4 DPI, and total soluble protein extracts were prepared for Western blot analysis with anti-His and anti-HA antibodies. Immunoblot membranes were stained with CBB. HS, heat shock treatment by placing the infiltrated plants in a 37 °C incubator for 30 min at 1 DPI. Red arrow, S$_{ct}$; blue arrow, CRT-HA on CBB-stained PVDF membrane. (c) Quantification of the S$_{ct}$ expression levels upon co-expression of CRT. The signal intensity of S$_{ct}$ upon co-expression of CRT at ratios from 2 : 1 to 5 : 1 was compared with that of S$_{ct}$ alone (1 : 0), with or without heat shock, and presented as relative values.

Figure 3  Plant-produced recombinant S$_{ct}$ glycoprotein form trimers. (a) SDS/PAGE analysis of purified proteins. S$_{ct}$ was purified using Ni$^{2+}$-NTA affinity resin and size exclusion chromatography, and separated by SDS/PAGE, followed by CBB staining. (b) Negative EM analysis of S$_{ct}$. Scale bars in the magnified pictures indicate 10 nm. (c) Deglycosylation of purified S$_{ct}$. 2 µg of purified S$_{ct}$ was treated by PNGase F (+) and separated by SDS/PAGE with untreated S$_{ct}$ (−) as a control. (d) Stability of purified S$_{ct}$. E-tubes containing 5 µg of purified S$_{ct}$ were placed at 4 °C. Samples were taken out on different days and analysed by SDS/PAGE. The band density was quantified. Black arrow indicates the S$_{ct}$. Statistical analysis was carried out using Student’s t-test. Error bar, standard error (n = 3).
Figure 4 Humoral and cellular immune responses of NS\textsubscript{ct}Vac in mice. (a) Schematic presentation of the mice immunization experiment. Five Balb/c mice per group were injected intramuscularly with NS\textsubscript{ct}Vac, with or without alum hydroxide, and then boosted twice (with an interval of 2 weeks). Mice sera were collected before and after immunization to determine humoral immune responses. (b) The end-point titre of IgG antibodies was tested in ELISA plates coated with recombinant S antigen. (c) The titre of neutralizing antibodies (NAbs) was determined in a 50% plaque reduction neutralizing test (PRNT\textsubscript{50}) at 27 and 42 days post-initial immunization. (d) Cross-neutralizing activity against three variant strains was determined in a PRNT using sera collected at 42-day post-initial immunization. (e) Cytotoxic T cell responses of splenocytes were evaluated 14 days after the 2nd boost. Secretion of IFN-\ensuremath{\gamma} and granzyme B was measured in an ELISPOT assay after stimulation with recombinant SARS-CoV-2S protein. Data were expressed as the mean ± SEM. Experiments were conducted independently and in duplicate ($n=5$ per group/experiment). The statistical significance of differences between groups was calculated by two-way ANOVA with Tukey’s multiple comparisons test ($*P < 0.05$, $**P < 0.01$, $***P < 0.001$). Dotted lines reflect the assay limit of quantitation. See also Figure S3.
NSctVac induces humoral and cellular immune responses in mice

Immune responses to the N. benthamiana-expressed S\textsubscript{ct} vaccine (NSctVac) candidate were evaluated in 8-week-old female Balb/c mice (n = 5). Mice received an intramuscular injection of various doses (1, 5, 15 and 30 \(\mu\)g) of NSctVac with or without alum hydroxide as an adjuvant. Mice were boosted twice (with an interval of 2 weeks) after the initial immunization, and sera were collected 1 day before each immunization (Figure 4a). To evaluate S-specific antibodies, we used ELISA plates coated with S protein, the S1 subunit, or the S2 subunit of SARS-CoV-2. On Day 13 post-prime, S antigen-binding antibody responses were detected in mice immunized with low to high doses of NSctVac. Antibody responses were increased in a dose-dependent manner 2 weeks after the first booster immunization in immunized mice. At 2 weeks after the second boost immunization, antibody levels were elevated in all immunized groups, with the high-dose group (30 \(\mu\)g plus alum hydroxide) reaching saturated end-point titres (Figure 4b). Interestingly, although both S1 and S2 subunit-specific antibodies were induced after the initial immunization, the level of S2-specific antibodies was higher than that of S1-specific antibodies (Figure 5S). Moreover, we observed S1- and S2-specific antibody responses after vaccination with inactivated virus, which were comparable with the antibody responses induced by the high dose of NSctVac. In line with NSctVac vaccination, we also found that vaccination with inactivated virus elicited higher titres of S2-specific antibodies (Figure 5G). In contrast, the placebo group immunized with phosphate saline (PBS) did not produce detectable antibodies.

To determine the neutralizing capacity of sera from immunized mice, we next measured neutralizing antibody (NAb) titres against SARS-CoV-2 S clade strain (KCDC003) in a PRNT assay (we used the same sera used for the ELISAs). After 2 weeks of the first booster shot, neutralizing antibodies were induced only by the high-dose (30 \(\mu\)g) vaccine, with a reciprocal mean titre of ~400 (88–1113). However, after the second booster shot, NAbS were also induced in the low- (1 \(\mu\)g) and medium-dose (5 and 15 \(\mu\)g) groups adjuvanted with alum hydroxide. In the high-dose (30 \(\mu\)g) group, the mean NAb titre reached 2050 (1342–3537), and the deviation between individuals was smaller than in the other groups (Figure 4C). In addition, we evaluated the ability of antibodies elicited by NSctVac (in mouse sera collected 42-day post-initial immunization) to neutralize the early variant GV (GSAID classification) and two WHO-announced variants of concern (VoC; Alpha, B.1.17 and Beta, B.1.351). Neutralizing titres against these strains were similar to those against the S clade; however, some individuals showed lower titres against the Beta variant than against the prototype S clade strain, although the difference was not statistically significant (Figure 4D).

To evaluate antibody-dependent enhancement (ADE) of infection by S-specific antibodies induced by NSctVac, we cultured SARS-CoV-2 virus in J774A.1 cells bearing mouse FcRRI\(\alpha\) and exposed the cells to serially diluted sera from immunized mice. We then measured the titre of progeny virus in a plaque assay. Since J774A.1 cells can be naturally infected with the SARS-CoV-2 virus, we observed neither virus infection nor ADE (Figure S7). Cellular immune responses elicited by NSctVac vaccination were evaluated in an ELISPOT assay with mouse splenocytes. Fourteen days after the second booster, we sacrificed mice and collected splenocytes, which were then cultured with recombinant S antigen as a stimulator. Subsequently, intracellular interferon \(\gamma\) (IFN-\(\gamma\)), granzyme B and interleukin 4 (IL-4) induction were measured. Compared with the placebo group, the NSctVac-immunized group showed increased secretion of IFN-\(\gamma\) and IL-4. Although IFN-\(\gamma\) was induced to a lesser extent than IL-4, functional granzyme B secretion (indicative of a cytotoxic T cell response) was significantly higher than in the placebo group (Figure 4E).

NSctVac protects hACE2 transgenic mice from SARS-CoV-2 infection

Finally, we evaluated the protective efficacy of NSctVac in K18-hACE2 transgenic mice expressing hACE2. Six-week-old K18-ACE2 transgenic mice (n = 14) received an intramuscular injection of NSctVac (30 \(\mu\)g) plus alum hydroxide, followed by a booster 2 weeks later (Figure 5A). Sera were collected 2 weeks after the prime and booster vaccinations and neutralizing antibody production was measured. On Day 13 post-prime, S-specific antibodies were induced in more than half of the mice, reaching a mean titre of 4514 at 13 days after the booster shot (Figure 5B). In addition, S1- and S2-specific antibody responses were similar to those seen in Balb/c mice. NAb titres were elicited after 13-day post-prime, with a mean titre of 76 (35–156), fivefold lower than in Balb/c mice, at 13-day post-boost (Figure 5C). Next, we used mouse sera collected 27-day post-initial immunization to determine the spectrum of cross-neutralization against four recently circulating VoC strains (Alpha, B.1.17; Beta, B.1.351; Delta, B.1.617.2; Omicron, B.1.529). The neutralizing antibody titres were lower than those against the prototype strain (S clade), with mean titres of 30.5, 28.6, 46.9 and 39.8 against Alpha, Beta, Delta and Omicron, respectively. Thus, the titres of Nabs against Delta and Omicron were higher than those against the Alpha and Beta variants (Figure 5D).

Subsequently, we infected NSctVac-immunized K18-hACE2 mice with \(5 \times 10^4\) pfu of SARS-CoV-2 (KCDC003) via the intranasal route on Day 14 post-booster injection. Bodyweight changes and survival were observed for 14 days. We found that no significant body weight loss was observed in the vaccinated mice. By contrast, body weight in the placebo control group fell gradually down until Day 8, and only one mouse survived and recovered (Figure 5F). For the purposes of survival analysis, subjects who lost 20% or more of their body weight were considered dead in accordance with the regulation of the Institutional Animal Care and Use Committee of the Korea Centers for Disease Prevention and Control. In the NSctVac-immunized group, only one mouse was terminated due to body weight loss (on Day 7 post-infection); the remaining mice survived for 2 weeks (a survival rate of 80%). However, four mice in the placebo group succumbed on Day 9 post-infection, and only one survived for 2 weeks (a survival rate of 20%; Figure 5F). To determine the viral titre in lung tissue, three mice were euthanized and tissues were collected at 3-, 5- and 7-day post-challenge. The viral load in the lungs of the NSctVac-immunized group was lower than that in the placebo group on Day 5 post-challenge. On Day 7 post-infection, no live virus was detected in vaccinated mice, whereas the placebo group still retained virus in the lung tissue (Figure 5E). Histopathological examination of lung tissues showed that SARS-CoV-2 incubation induced extensive inflammation at 3, 5 and 7 post-infection in the placebo group. Whereas, mice immunized with NSctVac showed minimal lesions (Figure 5G). Taken together, these data suggest that vaccination
with NSctVac induces protection in K18-hACE2 mice challenged with a high dose of virus.

**Discussion**

Expression level is a key aspect of recombinant protein production. Therefore, we explored various conditions. In general, membrane proteins have limitations with respect to high-level expression. We used a full-length protein lacking the TMD and the C-terminal domain. Another consideration was to produce a trimeric recombinant S protein to mimic that expressed by the native virus. Of the two trimerization motifs, FD and mCor1, we found that mCor1 was better for the expression level. We also examined the effect of double proline substitution (2P) in the S2 subunit that prevents fusogenic conformational change of MERS-CoV S (Pallesen et al., 2017), thereby stabilizing the prefusion state of S glycoproteins, which in turn leads to an increase in the final production yield. 2P substitution was used for the current SARS-CoV-2 vaccines. Moreover, a Hexa proline substitution (6P) increases the final yield in mammalian cells (Hsieh et al., 2020). The 3P substitution in the S2 subunit increased expression of the S protein in *N. benthamiana*. This suggests that the stability of the S protein is important for high-level expression in plants. Consistent with this idea, heat shock treatment and co-expression of CRT greatly enhanced the expression level. However, CRT and 3P did not show any additive effect in plants. Besides, a furin cleavage site mutation from RRAR to QQAQ was included in the subunit vaccine NVX-CoV2373 to prevent cleavage of the S protein into its S1 and S2 subunits. However, Sct expressed in plants was not subjected to furin cleavage at the furin site. This may be due to localization of the S protein in the ER. Another possibility is that plants may not contain furin homologues since no furin cleavage activity was observed in plants (Mamedov et al., 2019; Wilbers et al., 2016).

Protein purification, which is thought to account for up to 80% of total production costs (Buyel, 2015; Wilken and Nikolov, 2012), is the most challenging step during recombinant protein production. Initially, we used the same buffer conditions as those for S protein purification from animal cells, but found that they were not suitable for purification of recombinant Sct proteins from *N. benthamiana* extracts. Sct proteins purified from *N. benthamiana* were highly aggregated and exhibited a smeared background on the SDS-PAGE (Figure S4A), which was not removed after gel infiltration (Figure S4B). Plant extracts contain high levels of polyphenolic compounds and pigments, which can interfere with protein purification by interacting with proteins or binding to the affinity resin. Therefore, we optimized the extraction buffer with respect to the type and concentration of salt, buffer pH and detergents. Upon pretreatment of ground leaf tissues with PVPP and AC, two polyphenol adsorbents that bind to polyphenols and pigments, recombinant Sct protein no longer produced high molecular weight aggregates or a smeared background during purification. PVPP is used to remove polyphenols during beer and juice manufacturing (Cimini et al., 2014; Youn et al., 2004). Polyphenol is a kind of polymer containing several to dozens of –OH groups that bind to proline-rich segments of proteins. Also, polyphenolic compounds can interact with resins such as Ni²⁺-NTA, thereby interfering with specific binding of the His-tag to the resin. Finally, we set up the purification protocol in such a way that L-proline, PVPP and AC were mixed with ground tissue powder before adding extraction buffer; this eliminated the polyphenolic compounds and pigments.

The immunogenicity and protective efficacy of NSctVac were evaluated in Balb/c and K18-hACE2 mice. Antibody responses to NSctVac in mice were comparable with those induced by plant-derived recombinant protein antigen vaccines published by others (Mamedov et al., 2021; Tran et al., 2021). Notably, we found that alum hydroxide adjuvant increased humoral immunity in all vaccinated groups, but not cellular immunity (Figure 4e). These results are consistent with a previous report (Hogenesch, 2012). Since NSctVac comprises the full-size trimeric S antigen, we performed ELISAs with three different coating antigens to detect S1 and S2 subunit-specific antibodies. Unexpectedly, S2-specific antibodies were induced at relatively high concentrations (Figure 5). Similarly, we found that S2-specific antibody levels in sera from mice infected with inactivated virus were similar to those in vaccinated mice (Figure 56). As such, it is plausible that the intact S antigen elicits a high S2 antibody response, which is consistent with a recent study describing high levels of anti-S2 antibodies in serum from COVID-19 convalescent subjects (Nguyen-Contant et al., 2020). The structurally conserved SARS-CoV-2 S2 subunit shows >88% sequence homology among Beta coronaviruses, including previous SARS-CoVs; therefore, the functional S2 fusion peptide region might confer cross-neutralizing potency against other coronaviruses (Walls et al., 2020). High levels of S2-specific antibodies might have cross-reactive potential as we observed no significant ADE infection in this study (Figure 57).

In line with IgG generation, NAbs were induced 2 weeks after the primary vaccination, and titres increased following the booster
shots (Figure 4c). Due to the limited amount of serum collected from each Balb/c mouse, we only conducted PRNT against the GV (GISAID classification), Alpha (B.1.1.7) and Beta (B.1.351) variants available at that time. Antibody-mediated neutralization activity against the Beta variant was partially reduced; however, there was no significant difference in the neutralizing activity of the other variants (Figure 4d). In addition, neutralizing titres against Delta (B.1.617.2.1) and Omicron (B.1.1.529) were somewhat decreased in K18-hACE2 mice serum after the booster shot (Figure 5d). This finding is consistent with those of recent clinical studies showing that two-dose vaccination with the currently used mRNA vaccines provides limited protection against infection by the Omicron variant (Sarah A. Buchanan, 2022). However, a third booster dose of the current mRNA vaccines increases vaccine efficacy and neutralizing antibody titres against the Omicron variant, which may lower the risk of breakthrough infection (Andrews et al., 2022; Nemet et al., 2022; Pajon et al., 2022; Simone I. Richardson et al., 2022). As human CD8+ T cell epitopes are located throughout the SARS-CoV-2 S gene (Grifoni et al., 2021), a full-length S recombinant vaccine may stimulate T cell responses to a greater extent than a partial subunit vaccine such as an RBD targeted vaccine. Antigen-specific T cell responses to N505 Vac were determined after ex vivo stimulation of splenocytes. The number of IFN-γ- and granzyme B-secreting cells after vaccination was S- and 15-fold higher, respectively, than in the placebo group, implying cross-presentation of recombinant N505 Vac vaccine to antigen-presenting cells (Heath and Carbone, 2001; Joffre et al., 2012). The protective efficacy of N505 Vac was evaluated by challenging hACE2 knock-in mice, which showed severe illness and mortality after intranasal infection with SARS-CoV-2, with live virus (Oladunni et al., 2020). Despite the finding that N505 Vac-vaccinated K18-hACE2 mice have a weaker antibody response than Balb/c mice, there was a protective effect against challenge infection. As previously described, host T cells provide a complementary protective effect against SARS-CoV-2 infection, particularly when the humoral immune response is insufficient (Zhuang et al., 2021). Thus, T cell immunity contributes to protection from infection and ameliorates disease severity.

Although this study demonstrates the immunogenic and protective effects of a plant-derived vaccine in murine models, there are several limitations. First, although we measured overall IFN-γ secretion in splenocytes by ELISPOT, in-depth analysis of CD4+ and CD8+ T cell responses should be undertaken to fully identify the protective mechanism. Second, N505 Vac vaccination induced large amounts of S2-specific antibodies; however, the role of these antibodies remains to be addressed. Since the S2 subunit is conserved across coronaviruses (Shah et al., 2021), it is plausible to examine cross-reactivity among coronaviruses. Third, a 2-week vaccination interval may not be optimal. Since insufficient B cell maturation is attributed to a short vaccine interval, high levels of neutralizing antibody induction and affinity maturation may not occur (Rossier et al., 2022).

Regarding the current situation of variants pandemics, SARS-CoV-2 is constantly mutating and breaking through the pre-established immunity. Fast updating and producing the vaccines for new variants economically is crucial for controlling the waves of the pandemic worldwide. Plant-based vaccine manufacturing has various advantages (e.g. rapid production, cost-effectiveness and easy scalability) over mammalian systems; thus, it can be used to rapidly supply cheap vaccines to fight emerging infectious disease pandemics (Dhama et al., 2020). Currently, there have been several attempts to develop COVID-19 vaccines using plant-derived subunits and VLP platforms (ClinicalTrials.gov, 2022; Maharjan and Choe, 2021; Mamedov et al., 2021; Siriwat-tanonan et al., 2021) or plant-derived products to prophylactic protection against COVID-19 in general social settings (Daniell et al., 2022). Of these, a VLP vaccine from Medicago Inc. is the only licensed vaccine to date (HealthCanada, 2022). In all these efforts, one crucial aspect is the yield of proteins produced in N. benthamiana. In this study, we used an ordinary binary vector with a strong promoter and terminator and obtained 106 µg/g of fresh weight. For real commercialization, higher levels of expression would be more preferable. In fact, many high-performing vectors based on RNA or DNA viruses have been developed that can be used for commercialization of vaccine produced using S<sub>ct</sub> in the future.

In summary, immunization with a SARS-CoV-2 S trimer vaccine expressed in N. benthamiana induces robust humoral and cellular immune responses in mice, which confer protection against lethal challenge. Immunization with the prefusion-stabilized trimeric S induced broad-spectrum antibodies in a manner similar to natural viral infection. Thus, we suggest that plant-derived vaccines will be a useful platform in the near future and that N<sub>ct</sub>Vac has potential as another plant vaccine candidate in addition to VLP.

Materials and methods
Cells and viruses
Vero-E6 (CRL-1586) cells were purchased from the American Type Culture Collection (Manassas, VA) and maintained in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% FBS, 100 U/mL of penicillin and 100 µg/mL of streptomycin (Gibco, NY). The SARS-CoV-2 viruses [BetaCoV/Korea/KCDC03/2020, hCoV-19/Korea/KDCA23857/2020 (GV), hCoV-19/Korea/KDCA51463/2021 (Alpha) and hCoV-19/Korea/KDCAS1463/2021 (Omicron)] were obtained from the National Culture Collection for Pathogens at the Korea National Institute of Health (KNIH). Viruses were propagated in Vero-E6 cells, and the virus stocks were stored at −70 °C until use. All experiments using viruses, including those in mice, were performed in the animal biosafety level 3 (ABSL-3) laboratory facilities at KNIH.

Design and construction of the SARS-CoV-2 S and S2 vaccine candidates
The ectopic region of the S protein from amino acid positions 16 to 1213 [S(TMD-CT)] was codon-optimized and chemically synthesized (Gene Universal, Newark). BamHI1:S:S/(TMD-CT):L: SpeI was PCR amplified from template Sa/(TMD-CT) using primers F-S and R-S. BamHI1:S2:SpeI was PCR amplified from template Sa/(TMD-CT) using primers F-S2 and R-S. The site-specific mutations were introduced by two-step PCRs. The primers used for the A942P mutation were F-S, R-A942P, F-A942P and R-S; those used for the K986P/V987P mutations were F-S, R-K986P/V987P, F-K986P/V987P and R-S, and those used for the QOQA mutation were F-S and R-QOQA, F-QOQA and R-S. The trimerization motifs of the Foldon domain (FD; GYIPEAPRDGFY) or mouse Coronin 1A (mCor1; VSLLEEDVRLNAILQKLQERLDRLETVQAK), together with additional flanking regions (Spe1:Linker:F: hS5:HDEL:Xhol and Spe1: Linker:mCor1:Hs5:HDEL:Xhol, respectively), were chemically synthesized (Gene Universal, Newark). BamHI1:S:S/(TMD-CT):L: SpeI was PCR amplified from template Sa/(TMD-CT) using primers F-S and R-S. BamHI1:S2:SpeI was PCR amplified from template Sa/(TMD-CT) using primers F-S2 and R-S. The site-specific insertions were introduced by two-step PCRs. The primers used for the A942P mutation were F-S, R-A942P, F-A942P and R-S; those used for the K986P/V987P mutations were F-S, R-K986P/V987P, F-K986P/V987P and R-S, and those used for the QOQA mutation were F-S and R-QOQA, F-QOQA and R-S. The trimerization motifs of the Foldon domain (FD; GYIPEAPRDGFY) or mouse Coronin 1A (mCor1; VSLLEEDVRLNAILQKLQERLDRLETVQAK), together with additional flanking regions (Spe1:Linker:F: hS5:HDEL:Xhol and Spe1: Linker:mCor1:Hs5:HDEL:Xhol, respectively), were chemically synthesized.
synthesized. The PCR fragment BamH1::Sa(TMD-CT)::L::Spe1 was ligated to Spe1::Linker::FD::His5::HDEL::Xhol or Spe1::Linker::mcOr1::His5::HDEL::Xhol and ligated downstream of the leader sequence of Arabidopsis BiP in the pTEX1 vector harbouring the MacT promoter and the RD29B terminator (Song et al., 2021) using restriction endonuclease sites BamHI and Xhol to give BiP::Sa (TMD-CT)::FD::His5::HDEL and BiP::Sa::mcOr1::His5::HDEL, respectively. Sa(TMD-CT)::FD::His5::HDEL and BiP::Sa::mcOr1::His5::HDEL was replaced with BamH1::S2::Spe1 to yield BiP::S2::mcOr1::His5:: HDEL. The sequences of the primers are listed in the Table S1.

Expression and purification of recombinant S proteins from N. benthamiana

The expression vectors were transformed into Agrobacterium strain GV3101 and transiently expressed in 4–5-week-old N. benthamiana plant leaves using Agroinfiltration. The infiltrated leaves, the vacuum-mediated Agroinfiltration method was used to examine expression. To obtain a large amount of infiltrated leaves, the vacuum-mediated Agroinfiltration method was used and plant leaves were harvested at 3, 5, and 7 days post-infiltration (dpi). To examine expression of recombinant S and S2 proteins, infiltrated leaves were ground and mixed with protein extraction buffer (300 mM NaCl, 50 mM Tris–HCl, pH 8.5, 1 mM EDTA, 0.1% Triton X-100 and 1× protease inhibitor cocktail) at a ratio of 5:1 (V/V). The extracts were separated on 7.5% SDS-PAGE gels and analysed by Western blotting with a mouse anti-His antibody (15 000 dilution). The blot was developed with the enhanced chemiluminescence kit (Amersham Pharmacia Biotech, Piscataway, NJ), and images were captured using the LAS3000 system (Fujifilm, Tokyo, Japan).

To purify recombinant proteins, Ni²⁺-NTA affinity chromatography was performed according to the manufacturer’s protocol. Briefly, total protein extracts were prepared using 3 volumes of extraction buffer (100 mM NaCl, 25 mM Tris–HCl, pH 8, 2 mM CaCl₂, 0.1% Tween 20, 0.5% L-proline, 0.5% AC, 0.5% PVPP and 0.5× protease inhibitor cocktail). The mixture was shaken for 20 min in a cold room, followed by additional shaking for 10 min after addition of 2 mM Na+-phytate. The mixture was subjected to centrifugation at 18 000 g (3×10 min), and the supernatant was collected. Total protein extracts were loaded onto an Ni²⁺-NTA affinity column, followed by washing three times. Proteins were eluted using a total of 15 mL buffer containing 400 mM imidazole. The eluates were concentrated and loaded onto a Superose 6 10/300 increase chromatography was performed according to the manufacturer’s protocols. The sequences of the primers are listed in the Table S1.

Enzymatic deglycosylation of purified Sct proteins

Purified Sct protein was subjected to treatment with PNGase F (NEB #P0701) according to a previous study (Song et al., 2021). Briefly, 2 μg of Sct was denatured by boiling for 10 min. The denatured protein (10 μL) was chilled on ice and mixed with 2 μL GlycoBuffer (10×), 2 μL 10% NP-40, 5 μL H₂O and 1 μL PNGase F and incubated at 37 °C for 1 h. The samples were separated by SDS-PAGE, followed by CBB staining.

Negative staining and EM

Carbon-coated copper grids (Electron Microscopy Sciences) were glow-discharged at 15 mA for 30 s using a Pelco-Easiglow™ glow discharge cleaning system (Ted Pella). Purified Sct and S2ct spike trimer samples (0.01 mg/mL) were loaded on the glow-discharged carbon-coated copper grids for 1 min. Uranyl acetate (30 μg, 2%, w/vol) was used to stain the samples on the grid for 2 min. The stained grid was then blotted with filter paper, dried and stored for TEM analysis. TEM analysis was performed using a JEOL JEM-1011 TEM (JEOL USA, Inc., Peabody, MA). Images were obtained at a magnification of x150 000.

Immunization of Balb/c mice

Specific pathogen-free female BALB/c mice aged 6–8 weeks (Orient Bio, Sungnam, Korea) were used to investigate humoral and cellular immune responses. Mice (n = 5/group) were immunized with various doses (1, 5, 15 and 30 μg) of N. benthamiana-derived trimer S vaccine (N.Sct-Vac) via the intramuscular route, with or without 50 μL of 2% alum hydroxide adjuvant (Alhydro® gel; Invivogen). Two boosters were given post-initial immunization (at an interval of 14 days). Sera were collected at 1 day before each immunization to detect SARS-CoV-2 spike-specific IgG and neutralizing antibodies. Spleens were collected at Day 42 after initial immunization to assess cellular immunity by ELISPOT (R&D Systems, MN). All animal experiments were authorized by the Institutional Animal Care and Use Committee of the Korea Centers for Disease Prevention and Control (Protocol approval No. KDCD-075-19-2A), and all experiments were performed according to the guidelines of this committee.

Human ACE2 knock-in mice immunization and challenge

For the virus challenge experiments, 14 female hACE2 knock-in transgenic mice (B6.Cg-Tg(K18-hACE2)2Prlmn/J; aged 6–8 weeks; Jackson Laboratory, ME) were immunized with N.Sct-Vac (30 μg) plus alum hydroxide via the intramuscular route. Mice were boosted 2 weeks after the initial immunization. On Day 28 post-primarying, mice were anaesthetized by intra-peritoneal injection (200 mg/kg) and inoculated intranasally with 5 × 10⁴ pfu SARS-CoV-2 virus (BetaCoV/Korea/KCDC03/2020). Three mice were euthanized humanely, and lungs were collected to determine the viral load in a plaque assay (Cho et al., 2020) and histopathology on Days 3, 5 and 7 post-infection. The remaining mice (n = 5) were monitored continuously for 14 days to assess body weight changes and survival.

Enzyme-linked immunosorbent assay (ELISA)

Recombinant full-length SARS-CoV-2 spike (S), subunit 1 (S1) and subunit 2 (S2) proteins were expressed in Escherichia coli (BL21-DE3) and purified using an Ni²⁺-NTA column (Thermo Fisher). To detect spike-specific antibodies, 96-well plates were coated with purified recombinant S protein (0.5 μg/well) in PBS and incubated at 4°C overnight. After incubation, plates were washed three times with PBST (0.05% Tween 20 in PBS) and blocked with 5% skim milk in PBS. Serially diluted (twofold in blocking buffer, starting at 1:50) mouse serum was added to the plates for 1 h at room temperature. After washing with PBST, rabbit anti-mouse IgG-HRP (Southern Biotech, AL; diluted 1:1000 in blocking buffer) was added for 1 h. The colour signal was developed using o-phenylenediamine dihydrochloride (OPD) substrate solution (0.5 mg/mL OPD in 0.05 M phosphate citrate buffer, pH 5.0;
Sigma), and the reaction was stopped by adding 100 µL of 0.5 M sulphuric acid. Absorbance at an optical density (OD) of 490 nm was measured using an EPOCH microplate reader (BioTek, VT). Serum titres were determined as the largest reciprocal of the endpoint dilution that was greater than twice the background value.

Plaque reduction neutralization assay (PRNT)

Vero-E6 cells (200 000 cells/well) were seeded in 12-well plates and incubated for 16 h until 90% confluent. Serum samples were inactivated at 56 °C for 30 min and serially diluted (twofold diluted, starting at 1 : 10) in DMEM (Gibco) containing 2% FBS (Gibco) and 1% penicillin-streptomycin (P/S; 10 000 U/mL; Gibco). The diluted sera were mixed with the same volume of virus to generate a mixture containing ~80 PFU/200 µL of virus and then incubated at 37 °C for 1 h. The virus-serum mixture was added to Vero-E6 cell monolayers at 37 °C for 1 h. The plates were fixed with 4% paraformaldehyde (Biosesang, Korea) and stained with 1% crystal violet (Sigma-Aldrich, St. Louis, MO, USA). The 50% neutralization titre (PRNT50) was calculated using the Reed and Muench method (Reed and Muench, 1938).

Enzyme-linked immunosorbent spot assay (ELISPOT)

On Day 14 after the second boost, Balb/c mice spleens were isolated and the tissues were ground and filtered through a 70 µm nylon mesh cell strainer (Corning) to yield splenocytes. After red blood cells (RBC) were lysed using RBC lysis buffer, assays were performed using ELISPot kits (R&D Systems). Briefly, monoclonal antibodies specific for mouse IFN-γ, IL-4 or Granzyme B were pre-coated onto a PVDF-backed microplate and blocked with blocking buffer (10% FBS in RPMI). Splenocytes (1 × 10^6) were seeded into each well and stimulated with either DMSO (vehicle control) or various dilutions of the S protein. After 48 h of incubation at 37 °C/5% CO2, biotinylated anti-mouse IFN-γ, IL-4 or Granzyme B antibodies were added to each well at room temperature for 2 h. Finally, the spots were developed using a BCP/NBT substrate. The number of spot-forming cells on the plates was counted using an automated CTL Analyzer system (Cellular Technology, Cleveland, OH).

ADE assay

The murine monocyte-macrophage cell line (J774A.1) was seeded in 24-well plates at a density of 1.5 × 10^5 cells per well. Serially diluted mice sera were mixed with SARS-CoV-2 virus at a multiplicity of infection of 0.01 and incubated for 1 h. The virus/serum mixtures were added to the cells and incubated at 37 °C for 48 h. Cells and supernatants were harvested by centrifugation at 200 g for 5 min, and the virus titre in Vero-E6 cells was measured by plaque assay.

Statistical analysis

Details regarding statistical analysis are provided in the figure legends. The end-point IgG titre is presented as the mean titre along with the standard error of the mean (SEM) and was calculated by assigning a titre of 10 to samples with no detectable IgG antibodies at the starting dilution (titre <50). The statistical significance of differences between control and vaccination groups was assessed using two-way ANOVA with Tukey’s multiple comparison test. Data plotting and statistical analysis were performed using GraphPad Prism 8 (GraphPad Software, San Diego, CA). A value of P < 0.05 was considered significant.

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Conflict of interest

The authors declare no conflicts of interest.

Author contributions

J.H.C. and I.H. designed and supervised the experiments and analysed the data. J.H.C., S.I.S., H.K. and E.Y.J. wrote the manuscript. E.-J.S designed a codon-optimized S protein gene. S.I.S. and H.P.D. conducted S antigen expression and purification experiments. H.J. performed the EM experiment. M.R.I.K. made the CRT and 3P mutations. H.K. and E.Y.J. conducted the animal experiments, ADE, and ELISPOT. Y.J.L. and J.H.N. performed PRNT. J.H.N. and S.R.K. expressed recombinant proteins and conducted ELISAs.

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Supporting information

Additional supporting information may be found online in the Supporting Information section at the end of the article.

Figure S1 Optimizing expression of recombinant S protein constructs in N. benthamiana.

Figure S2 Expression of recombinant S2A and S2B proteins in N. benthamiana.

Figure S3 Expression of S1 in the presence of CRT and its purification by Ni²⁺-NTA.

Figure S4 Pretreatment with PPVP and activated charcoal removes high molecular weight aggregates and the smeared background.

Figure S5 SARS-CoV-2 S1- and S2-specific antibody responses after plant-produced NSctVac vaccination of mice.

Figure S6 SARS-CoV-2 S-specific antibody responses after vaccination of Balb/c mice with inactivated virus.

Figure S7 Evaluation of antibody-dependent enhancement (ADE) after NSctVac vaccination.

Table S1 The nucleotide sequences of primers used in this study.