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A defective viral genome strategy elicits broad protective immunity against respiratory viruses

Graphical Abstract

Highlights
- eTIP1 is a novel tool to protect against viral infections, including SARS-CoV-2 strains
- Intranasal eTIP1 delivery elicits innate antiviral responses in the respiratory tract
- eTIP1 provides pre- and post-exposure protection against respiratory viral infections
- eTIP1 treatment boosts generation of protective antibodies against pathogenic virus

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In brief
A defective viral genome derived from poliovirus induces type I interferon-mediated prophylactic and therapeutic effects against respiratory viruses, including SARS-CoV-2 and influenza, in mouse infection models.
A defective viral genome strategy elicits broad protective immunity against respiratory viruses

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SUMMARY

RNA viruses generate defective viral genomes (DVGs) that can interfere with replication of the parental wild-type virus. To examine their therapeutic potential, we created a DVG by deleting the capsid-coding region of poliovirus. Strikingly, intraperitoneal or intranasal administration of this genome, which we termed eTIP1, elicits an antiviral response, inhibits replication, and protects mice from several RNA viruses, including enteroviruses, influenza, and SARS-CoV-2. While eTIP1 replication following intranasal administration is limited to the nasal cavity, its antiviral action extends non-cell-autonomously to the lungs. eTIP1 broad-spectrum antiviral effects are mediated by both local and distal type I interferon responses. Importantly, while a single eTIP1 dose protects animals from SARS-CoV-2 infection, it also stimulates production of SARS-CoV-2 neutralizing antibodies that afford long-lasting protection from SARS-CoV-2 reinfection. Thus, eTIP1 is a safe and effective broad-spectrum antiviral generating short- and long-term protection against SARS-CoV-2 and other respiratory infections in animal models.

INTRODUCTION

RNA viruses pose a continuing worldwide health threat. Each new epidemic, from HIV to influenza to dengue and Zika to SARS-CoV-2, highlights an urgent need for effective antiviral drugs and interventions (Meganck and Baric, 2021). The highly contagious coronavirus disease 2019 (COVID-19) (Zhu et al., 2020) has caused more than 200 million cases and nearly 5 million deaths, and the numbers are still increasing (WHO, 2021). Despite development of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) vaccines, the virus is still circulating (Cohen et al., 2021; Deng et al., 2021; Guo et al., 2021; Kang et al., 2021; Abdool Karim and de Oliveira, 2021; Rappazzo et al., 2021; Sun et al., 2021; Zhou et al., 2021). A few antivirals and monoclonal antibodies are available for SARS-CoV-2 therapy (Canedo-Marroquin et al., 2020; Gonçalves et al., 2021; Lundstrom, 2020), but the high mutation rate of the RNA virus is likely to give rise to resistant variants. Clearly, novel therapeutics with a low risk of drug resistance are essential to combat COVID-19 and future epidemics. Unfortunately, developing broad-spectrum antiviral strategies is extremely challenging (Geller et al., 2012; Meganck and Baric, 2021; Tse et al., 2020).

One attractive strategy would be to harness the effective antiviral defenses of the host. Viral infections elicit a range of responses that prevent or attenuate most infections. Indeed, even for viruses such as influenza virus or SARS-CoV-2, most infections are asymptomatic (Cohen et al., 2021; Lutrick et al., 2021; Zhang et al., 2021). However, harnessing these responses is difficult. First, most viruses have mechanisms to inactivate or dampen innate immunity, often at multiple steps of the pathway (Kikkert, 2020). Second, direct interventions to induce these responses, such as intravenous administration of interferon, have undesirable side effects, likely because they override the delicate regulatory balance that keeps beneficial innate immunity from damaging tissues (Guo et al., 2021; Nelemans and Kikkert, 2019).
The innate immune system evolved tightly regulated mechanisms to protect from viral infection and prevent damage to the host. The challenge is to harness these beneficial responses without triggering detrimental side-effects. Our strategy was based on an unexpected and remarkable observation from early epidemiological studies of the Sabin poliovirus (PV) vaccine. Immunization with attenuated Sabin PV protected from PV infection and also reduced influenza virus morbidity by almost fourfold. It even accelerated healing of genital lesions caused by herpes simplex virus. These benefits were never observed upon immunization with an inactivated PV virus (e.g., Salk vaccine). Importantly, the Sabin vaccine causes none of the severe side-effects of interferon administration (Chumakov et al., 2020, 1991). These puzzling observations raise the hypothesis that a non-pathogenic virus or virus-like entity could safely stimulate the host innate antiviral responses.

To test this hypothesis, we developed an engineered virus-like entity, based on a defective viral genome (DVG), and tested its potency as a broad-spectrum antiviral. Due to the high error rate of their RNA-dependent RNA-polymerases, RNA viruses naturally generate genome deletions with various degrees of viability (Crotty and Andino, 2002; Crotty et al., 2001; Domingo et al., 1996, 2002; Peersen, 2017; Pfeiffer and Kirkegaard, 2003). These DVGs are maintained by co-infection with the parental virus (Dimmock and Easton, 2014, 2015; Easton et al., 2011; Goff et al., 2012; Huang and Baltimore, 1970; Kim et al., 1998; Kitamura et al., 1981; McClure et al., 1980; Perrault, 1981; Perrault and Semler, 1979; Shirogane et al., 2019; Vignuzzi and López, 2019), but lack critical portions of the viral genome required for replication. eTIP1 carries a large region (yellow) encodes the enzymatic machinery of the engineered DVG genome, herein called eTIP1.

We engineered a PV-derived DVG that can be delivered by lipid nanoparticles and is highly effective at preventing replication of respiratory viruses, including rhinovirus, influenza virus and SARS-CoV-2 in cell and animal models. The DVG blocks viral replication by inducing an antiviral state in the respiratory tract. It can be administered intranasally, as pre- or post-exposure prophylaxis, to protect mice from pathogenic viruses without detrimental side-effects. Indeed, eTIP1 (enteroviral therapeutic interfering particle) infectious particles protect even...
when administered 24–48 h post-infection with SARS-CoV-2, PV, coxsackievirus B3 (CBV3), and influenza virus. Importantly, eTIP1 reduces virus load by several orders of magnitude and also enables generation of neutralizing antibodies against the challenging virus. This enhanced antibody response provides long-term protection from reinfection, lasting weeks after the initial intervention. We suggest that our approach is an effective, non-invasive, broad-spectrum strategy to block viral infections, including SARS-CoV-2.

RESULTS

Engineering a defective poliovirus genome as a broad-spectrum antiviral

We engineered a DVG for PV type 1 (PV1) by replacing the entire P1 region, which encodes structural proteins, with GFP (Figure 1B). eTIP1 infectious particles were produced using a packaging HeLa cell line that stably expresses the PV1 capsid protein precursor P1 (HelaS3/P1) (Figure 1C). Transfection of HeLaS3/P1 with in vitro transcribed eTIP1 RNA generated eTIP1 infectious particles that were amplified by repeated infection of HeLaS3/P1 (Figure 1C). eTIP1 particles can infect cells, as determined by expression of GFP and immunofluorescence (EM) with polio-3A antibody (Figures 1E and S1A). Importantly, eTIP1s cannot spread from cell to cell without a WT PV1 acting as a helper virus (Dimmock and Easton, 2015; Perneault and Semler, 1979; Shirogane et al., 2021a; Vignuzzi and López, 2019).

To test the therapeutic potential of eTIP1s, we tested if it could block replication of PV1 and related enteroviruses of clinical importance. To this end, we infected cells with enteroviruses with and without eTIP1s at a ratio of 1:20 (eTIP1 multiplicity of infection [moi] = 1–5). eTIP1s effectively blocked replication of PV1 and other enteroviruses, including enterovirus EV-D68, EV-A71, CVB3, and rhinoviruses (Figures 1F and S1B). Virus replication was inhibited 10- to 1,000-fold, depending on the virus and cell line (Figure 1F). To determine if eTIP1 antiviral activity is restricted to the genus of its parental virus, we next examined its activity against an unrelated enterovirus. eTIP1 co-infection at a ratio of 1:50 inhibited influenza A (A/PR8) and SARS-CoV-2 by 100- to 1,000-fold (Figure 1F). Antiviral activity was even stronger when eTIP1 was administered 5 h before infection with PV1, A/PR8, or SARS-CoV-2 (Figure 1G). These results suggest that the inhibitory activity of eTIP1 does not rely on direct competition for enteroviral proteins, such as the viral capsid or other viral or host factors required for enterovirus replication.

eTIP1 prevents lethal infection in mice

Given the broad-spectrum inhibitory effects of eTIP1s in cell culture, we examined their ability to prevent disease in mice infected with several pathogenic viruses. After intraperitoneal (IP) inoculation of susceptible mice, PV1 replicates and accumulates to high titers in diverse tissues, ultimately reaching the central nervous system (CNS) to cause paralysis and death (Ida-Hosonuma et al., 2005; Ohka et al., 2007; Pfeiffer, 2010; Xiao et al., 2017). Similarly, after intranasal (IN) inoculation, PV1 reaches the CNS rapidly through the olfactory nerve, causing severe disease (Nagata et al., 2004).

We infected mice by the IP route with high doses of PV1 (10⁷ plaque-forming units, pfu) with or without co-inoculation with eTIP1s (Figure 2A, i, IP). Under these conditions, eTIP1 significantly attenuated disease and protected 80%–90% of mice from lethal infection. Importantly, co-inoculation of eTIP1s activated by UV irradiation did not protect mice from PV1, indicating that the eTIP1s must be replication competent to exert its antiviral properties. This experiment ruled out that protection was conferred by a contaminant introduced in eTIP1 production. eTIP1s also protected animals from disease (weight loss) and death after infection with a non-polio enterovirus, CVB3 (Figures S2A and S2B). Next, we determined if eTIP1 protects animals from respiratory infection. IN co-inoculation of PV1 and eTIP1s protected from death (Figure 2A, ii) and reduced PV1 viral loads in spleen and brain over 500-fold (Figure S1C).

To determine if eTIP1-induced protection is long lasting, we administered a single IN dose of eTIP1 and challenged the mice with pathogenic PV1 48 h later. While animals inoculated with PV1 alone succumbed to infection, pre-exposure administration of eTIP1 protected 90% of the animals from lethal disease (Figure 2B, i). Post-exposure administration of eTIP1 24 and 48 h after PV1 infection also elicited significant protection (Figure 2B, ii). The protective effects were lost if treatment is initiated 72 h after PV1 infection (not shown). These results indicate that both pre- and post-exposure prophylaxis with eTIP1 protects against IN PV1 infection, preventing severe disease and death. In addition, eTIP1 protected from infection with A/PR8 and CVB3 when administered 24 h (A/PR8 and CVB3 (Figure 2C, ii, and Figure S1D) or 48 h (CVB3) after the viral challenge (Figure 2C, iii). This indicates that eTIP1 can protect from respiratory infections after a single IN dose.

Intranasal inoculation of eTIP1 lipid nanoparticles and its replication tissue distribution

Our finding that eTIP1s interfere with viral infection in mice supports their use as an antiviral. However, their applicability may be limited by practical considerations such as a requirement for the PV receptor (PVR) to enter target cells and neutralization in vaccinated individuals. We thus tested whether eTIP1 DVG could be delivered by synthetic nanostructured lipid/RNA complexes (herein lipid nanoparticles, or LNPs). LNPs deliver RNA via endocytosis, which is unaffected by pre-existing immunity, and so may be administered as multiple consecutive doses (Figure 3A). We chose a cationic lipid formulation that binds well to the phosphate backbone of nucleic acids, and is easy to prepare, non-toxic, and extensively characterized (Kranz et al., 2016; Scheideler et al., 2020). We find that LNPs protect RNA molecules from hydrolysis and enable eTIP1 delivery to initiate replication at mucosal repatory surfaces. We confirmed that LNPs deliver synthetic eTIP1 RNA in cell cultures (Figure S3A).

Then, we compared the site of eTIP1 replication when delivered to mice via IN administration of particles or LNPs. At 24 h post-inoculation, immunohistochemistry of mouse heads and lungs
detected eTIP1 replication in epithelial cells within the ethmoid turbinates of the upper respiratory nasal cavity (Figure 3B). eTIP1 replication was limited to those tissues, not detected in other regions of the respiratory tract, and observed for only 12–24 h post-inoculation.

We next tested if eTIP1 can be mobilized from the site of inoculation to other tissues by a co-infecting help virus. We co-inoculated highly susceptible mice (IFNAR−/−) intramuscularly with 200 pfu of PV1 with eTIP1s (50,000 IUs). The high concentration of the initial inoculum at the site of injection increases the probability PV1 and eTIP1 co-infection, enabling trans-encapsidation of eTIP1s by PV1 structural proteins. Using IFNAR−/− mice increases susceptibility to PV infection and the rapid spread of the virus to diverse tissues. We examined PV1 and eTIP1 replication in muscle, spleen, and spinal cord by RT-qPCR. eTIP1 RNA accumulated at the site of inoculation on days 1 and 3 and decayed by day 6 (Figure S3B). eTIP1 RNA was barely or not detected in spleen or spinal cord, indicating that the eTIP1 does not spread beyond the site of inoculation even with PV1 (Figure S3B). IFNAR−/− mice inoculated only with eTIP1s showed no sign of distress and survived inoculation for several weeks (Figure S3C). Thus, eTIP1 replication is restricted to the site of initial replication (i.e., muscle [IM] or nasal turbinates [IN]).
Figure 3. eTIP1 protects against PV through a type I IFN response

(A) Schematic representation of the eTIP1 biological particles and eTIP1 RNA complex with LNPs. Animals were analyzed 24 h after IN intranasal inoculation by immune histochemistry (IHC) or by RNA-seq transcriptome profiling of the lung gene expression. (B) eTIP1 site of replication within the respiratory tract. Mice were inoculated IN with 30 μg of eTIP1 RNA or mock (PBS with empty LNPs) or infected with 6 × 10^6 infectious units of eTIP1. Heads of inoculated animals were analyzed 24 h post-inoculation by IHC. Heads and lungs were collected and fixed in 4% paraformaldehyde (PFA), embedded in paraffin wax, and cut into 5-μm sections. eTIP1s and eTIP1 RNA were stained using PV antibody VPg (3B protein). VPg (3B) (red), ACTUB (green), nuclear (blue). eTIP1 replication was restricted to the upper respiratory nasal cavity. We observed no replication in the lungs.

(C) C57BL6 TgPVR mice were infected with 6 × 10^6 IU eTIP1 particles or PBS (mock) (i), or animals were inoculated IN with 30 μg eTIP1 LNP or mock (empty LNP). Lung tissues were collected 24 (eTIP1 LNP) or 48 h (eTIP1 particles) post-inoculation, and mRNA was isolated from these tissues and examined by RNA-seq. Volcano plot shows pairwise comparisons of mRNA levels in infected versus mock-infected lung tissues and represented as a volcano plot of the genes with significant changes in expression, compared to the mock-treated group (false discovery rate, q-value < 0.05). n = 3 on two experimental replicates.

(D) Pairwise comparison of eTIP1 RNA/LNP versus eTIP1 particle in lung of infected animal. Red dots represent type I IFN genes.

(E) eTIP1 fails to protect against PV1 (IN in mice lacking IFNAR /C0 /C0). IFNAR /C0 mice were infected with 5 × 10^4 pfu PV1 alone or co-infected with mixed PV1 + eTIP1 at a ratio of 1:20 by IN route. Black line represents PV1 alone. Red dash line represents co-infected mixed PV1 + eTIP1 group (n = 7-10). Data were collected from two independent experiments. The comparison of survival curves was performed by log-rank (Mantel-Cox) test. ns, not significant.

(F) RT-qPCR validation of upregulated genes in lungs of animals treated with eTIP1. K18-hACE2 mice were transfected with 30 μg of eTIP1 RNA or infused with 6 × 10^6 IU of eTIP1 particles or PV1. As control, we mock-infected animals (PBS with empty LNPs, lipofectamine 2000) for 24 h. Lungs were collected, and total RNA was extracted with Trizol reagents. RT-qPCR were performed to qualify the IFN-induced genes MX1 and ISG56 (IFIT1), n = 3, normalized to GAPDH. Unpaired Student’s t tests. *p < 0.05; **p < 0.01; ***p < 0.001.
Role of interferon in eTIP1-mediated antiviral protection

We next sought to determine the mechanism by which eTIP1 induces the antiviral protective effect. We considered two possible models. In one, DVGs, whose shorter genomes may provide a replication advantage, outcompete the full-length viral genome for cellular resources and encapsidation by structural proteins, thus impairing propagation of parental virus to other cells in the tissue (Shirogane et al., 2021b; Vignuzzi and López, 2019). In the second, DVGs induce an innate response that cross-protects from other viral infections. Self-replicating RNAs, particularly DVGs that form cytosolic double-stranded RNA (dsRNA) intermediates, activate pattern recognition receptors, and trigger innate immune responses that lead to production of interferon (IFN)-stimulated genes (ISGs) (Brennan and Bowie, 2010; Finlay and McFadden, 2006; Kikkert, 2020; Narayanan and Makino, 2009; Vignuzzi and López, 2019). In this way, DVGs may induce a systemic antiviral state that interferes with replication of WT virus. In addition, DVGs may cause cells to lose the integrity of their plasma membrane and release damage-associated molecules that recruit various types of circulating leukocytes to the site (Preissner et al., 2020). While the first option requires co-infection of DVG and WT virus for interference to occur, the second model is consistent with DVGs impairing WT virus replication in a non-cell-autonomous manner. Our data are consistent with the second mechanism.

To test this hypothesis, we performed unbiased transcriptome profiling in lungs of eTIP1- or mock (PBS)-treated mice (Figure 3A). Whole lungs were harvested at 1 or 2 days post-IN administration of either 6 × 10^6 IU of eTIP1s or 30 μg of eTIP1 RNA in LNPs. RNA sequencing (RNA-seq) analysis showed that both methods induced a common set of bona fide type I IFN-responsive genes (Schoggins and Rice, 2011; Schoggins et al., 2011, 2014; Xiao et al., 2017), including Ifit1-3, Eif2ak2 (PKR), Irf7, Ifg15, Rasd2, Mx1, Adar, and a number of Oas paralogs (Figures 3C). Type I ISGs were similarly induced regardless of method (Figures 3D and S4B). This was confirmed by RT-qPCR analysis of two ISGs, Mx1 and Ifi1, in lungs of mice inoculated with eTIP1s or LNPs (Figure 2F). These genes were also induced at 24 h post-infection with PV1, albeit to lower levels (Figure 2F). This suggests that the IFN response is stronger without the capsid coding region of PV1. Foot-and-mouth disease virus capsid protein VP3 inhibits IFN signaling (Li et al., 2016a, 2016b). Thus, IN delivery of eTIP1 as biological particles or LNPs stimulates a more robust innate immune response than WT PV.

These experiments suggest that eTIP1 induces an IFN-mediated antiviral state. Next, we directly tested whether eTIP1-mediated antiviral protection requires systemic innate immune stimulation of an IFN response. We tested the capacity of eTIP1 to protect animals lacking the IFN-α/β receptor (IFNAR−/−) against PV1 infection. Under conditions in which eTIP1 elicits a robust protective activity in immunocompetent WT mice (Figure 2), the antiviral protection is completely lost in animals lacking the IFN-α/β receptor (IFNAR−/−; Figures 3E and S4B). This indicates that IFN-I responses are important for the eTIP1-mediated protection from IP and IN infection with pathogenic PV1.

Pre- and post-exposure eTIP1 treatment protects from SARS-CoV-2

Next, we determined if eTIP1 protects animals from SARS-CoV-2 infection. At 24 h after a single IN dose of eTIP1 LNPs, we infected K18-hACE2 mice with 6 × 10^6 pfu SARS-CoV-2 (D614G variant). We measured the effects of eTIP1 on SARS-CoV-2 replication in clinically relevant tissues (i.e., lungs and brains) collected on days 3 and 6 post-infection. SARS-CoV-2 replication was detected by plaque assay and RT-qPCR, as well as immunohistochemistry with antibodies against the nucleocapsid proteins (NPs) and spike proteins (SPs). As expected, mice pre-treated with control LNPs had significant SARS-CoV-2 titers and widespread NP and SP immunoreactivity throughout the lung and brain (Figures 4A–4C). Pretreatment with eTIP1 LNPs lowered SARS-CoV-2 titers and viral RNA copies by 2–3 logs (Figure 4A) and reduced immunoreactivity in lungs and brains (Figures 4B and 4C). Inoculation of LNPs carrying eTIP1 inactivated by UV, a replication-incompetent eTIP1 RNA carrying a large deletion of the 3’ of the genome (truncated PvuII), or poly IC did not protect mice from SARS-CoV-2 infection (Figure SSA), indicating that eTIP1s must be replication competent to protect from SARS-CoV-2 infection, as shown earlier (Figure 2A).

We also tested if eTIP1s protect against other variants of concern or interest (VOCs or VOIs), including B.1.1.7 (Alpha), B.1.617.2 (Delta), and B.1.427/B.1.429 (Epsilon). Inoculation of eTIP1 24 h before infection reduced virus titers in lungs by two orders of magnitude (Figure 4A, ii, eTIP1-prophylactic). Treatment with eTIP1s 24 h post-exposure to SARS-CoV-2 also elicited protection (Figure 4A, ii, eTIP1-therapeutic). The protective effects of eTIP1s were lost when administered 48 h after infection, but repeating eTIP1 treatment at both 24 and 48 h post-infection increased its efficiency (Figure 5A). These results indicate that pre- and post-exposure eTIP1 prophylaxis protects from IN infection with several circulating SARS-CoV-2 variants.

Weight loss is a sensitive measure of animal distress. SARS-CoV-2-infected mice lost weight due to disease progression. Strikingly, a single IN eTIP1 dose prevented weight loss in SARS-CoV-2-infected mice. In fact, eTIP1-treated animals maintained their body weight, similar to mock-infected controls (Figure 5B). This confirmed that eTIP1s reduce SARS-CoV-2 replication and prevent disease symptoms without additional distress. Next, we wondered if eTIP1s protect lungs from SARS-CoV-2 infection-induced inflammation. Lung sections were analyzed after staining with hematoxylin and eosin (H&E) and scored on tissue pathology (Meyerholz and Beck, 2020). SARS-CoV-2-infected mice had higher histopathology scores than mock-infected mice (Figure 5C, i and ii). The major histopathology findings in infected mice were proteinaceous debris in the alveolar space, neutrophils in the interstitial space, and alveolar septal thickening. These are consistent with previous studies that detected signs of lung injury, including interstitial pneumonia, inflammatory cell infiltrates, and alveolar septal thickening (Dinnon et al., 2020; Gu et al., 2020). By histopathology analysis, treatment with eTIP1 LNPs reduced lung inflammation after exposure to SARS-CoV-2 (histopathology score of 1/16 compared to empty-LNP histopathology score of 5.4/16) at day 3 post-infection (Figure 5C, i and ii, SARS-CoV-2/eTIP1). No peribronchial inflammation was noted in the lungs when animals were treated with eTIP1s alone,
indicating that eTIP1 replication does not cause persistent inflammation (Figure 5C, i and ii, eTIP1). Thus, a single IN dose of eTIP1s reduces in vivo replication of SARS-CoV-2 by orders of magnitude and prevents lung inflammation and SARS-CoV-2 infection-related weight loss (Jiang et al., 2020; Rappazzo et al., 2021; Sefik et al., 2021; Sun et al., 2020).

Although SARS-CoV-2 replication is strongly inhibited, H&E staining showed infiltration of lymphoid cells into the lung (Figure 5C), which may be linked to an eTIP1-mediated antiviral state. To determine which cells are recruited to the lung upon eTIP1 treatment, we performed immune cell profiling in lung tissues at 24 h after IN inoculation with eTIP1 LNPs (Figure S5). We also examined immune cell profiles 3 days post-SARS-CoV-2 infection with or without eTIP1 pre-exposure prophylaxis. No differences were observed in lung immune cell populations 24 h after inoculation with eTIP1 (Figure 5D, compare mock with eTIP1/LNPs). In the context of SARS-CoV-2 infection, eTIP1 treatment recruited different lymphoid cells into the lung, including eosinophils, plasmacytoid dendritic cells, and neutrophils, which are poorly recruited by SARS-CoV-2 infection alone (Figure 5D). Eosinophils can be recruited to the airways in response to influenza and mycobacterium tuberculosis (TB) infection in a type I IFN-dependent fashion (Bohrer et al., 2021; Samarasinghe et al., 2014). eTIP1s reduced the recruitment of CD11b+/CD11c+/Ly6C+ monocyte to the lungs of SARS-CoV-2-infected animals (Figure 5D), which may be associated with the pro-inflammatory response to SARS-CoV-2 infection. No other changes in immune cell populations were seen after eTIP1 pre-exposure prophylaxis (Figures 5D and S6). These data indicate that eTIP1 treatment modulates the recruitment of immune cells into the lung during SARS-CoV-2 infection.

eTIP1 treatment induces long-lasting protection against SARS-CoV-2 reinfection

A critical question is whether eTIP1-mediated protection from SARS-CoV-2 prevents the development of adaptive immunity
against future reinfections. Immunity could last at least 5–6 months after infection (Huang et al., 2021). Thus, reinfection with SARS-CoV-2 may occur in fewer than 1% of individuals who tested positive for SARS-CoV-2, at least within 3–6 months after infection. However, the natural protection is variable and unreliable, especially among older people (Collier et al., 2021).

To test if adaptive immunity against SARS-CoV-2 is induced in animals protected by eTIP1s, mice were treated with eTIP1s or control empty LNPs, and 24 h post-treatment, mice were infected with 10^5, 10^4, or 10^3 pfu of SARS-CoV-2 (B.1.1.7 variant). Inoculation of eTIP1s reduced virus titers in lungs by three orders of magnitude more than controls (Figure 6B, for 10^5 pfu). By RT-qPCR, we found that eTIP1 treatment effectively reduced SARS-CoV-2 RNA load in lungs 24-fold compared to SARS-CoV-2 infection without eTIP1 treatment. However, we detected low levels of viral RNA in lungs (Figure 6B). At 3 weeks after infection, we measured neutralizing antibody titers. Antibody titers and seroconversion rate correlated with the initial SARS-CoV-2 concentration used to infect the mice (Figure 6C). Thus, mice infected with 10^5 pfu had a more effective induction of neutralizing antibodies than those infected with 10^4 or 10^3 pfu. Strikingly, antibody titers were similar in eTIP1-treated and untreated animals (Figure 6C). We next challenged animals intranasally with 10^5 pfu of SARS-CoV-2 (B.1.1.7) and measured virus loads in lung tissue 3 days post-reinfection (Figure 6D). Mice initially infected with 10^5 pfu of SARS-CoV-2 and treated with eTIP1 were well protected, but low doses of SARS-CoV-2 in the initial infection did not reliably generate protective immunity. Mice infected with high doses of SARS-CoV-2 were more susceptible to reinfection. Perhaps higher doses of SARS-CoV-2 suppress effective induction of adaptive immunity, enabling re-infection. Additional investigations will be necessary to fully understand this observation.

Nevertheless, our findings indicate that eTIP1s replicate in the initial infected cells at the site of administration, without spreading to other cells (Figure 3B). Given that only replication-competent...
eTIP1s elicit protective responses (Figures 2A and S5), we hypothesize that intermediates of replication, such as dsRNA, are detected by pattern recognition receptors, leading to synthesis of type I IFN and induction of potent innate responses. Because the PV1 protease and other non-structural viral proteins in the eTIP1 induce profound rearrangements of cellular organelles and pathways, which may generate additional signals that further activate innate responses (Figure 6E). This results in the recruitment of leukocytes into the tissue and promotes an antiviral response that protects local and distal sites (non-cell-autonomous, distal protection, is effective prophylactically and therapeutically, and can boost neutralizing antibody production.

Mucosal challenges with RNA viruses achieve sterilizing immunity with no adverse effects involving short- or long-term immune dysfunction (Ascough et al., 2019; Jangra et al., 2021; Kikkert, 2020). Our data suggest that modulation of natural innate antiviral immunity is a primary component of eTIP1 antiviral activity. Importantly, a single intranasal dose prevents progression to severe viral disease longer than a single dose of small-molecule antivirals (Chen et al., 2021; Gu et al., 2020; Wang et al., 2020). We conclude that IN delivery of eTIP1 nanoparticles safely and potently stimulates host innate immunity that broadly protect from infection by reducing viral loads and preventing disease. While the antiviral response elicited by eTIP1 confers short-term protection as pre- and post-exposure prophylaxis, it does not interfere with long-term adaptive immunity (Figure 6C) that protects from reinfection (Figure 6D). We propose that eTIP1 has a compelling clinical potential in the treatment of respiratory viral disease.

**DISCUSSION**

Here, we provide proof of principle that DVGs can be used as pre- and post-exposure prophylaxis against diverse viral pathogens. We show that eTIP1s can be administered IN to combat respiratory infections from enterovirus to SARS-CoV-2, without...
side effects. Because eTIP1 IN inoculation offers protection from 48 h before to 24 h after exposure, it provides an effective therapeutic window, comparable to small-molecule antivirals. Inoculation of eTIP1 provides rapid protection against ongoing infection, but also promotes long-term protection by generating longer-lasting adaptive immunity.

Our study provides a mechanistic rationale for decades-old observations that immunization with live attenuated PV cross-protects against other viruses and DVGs modulate the course of infection (Easton et al., 2011; Genover and López, 2019; Huang and Baltimore, 1970; Levi et al., 2021; Perrault and Semler, 1979; Rezelj et al., 2021; Semler et al., 1978; Vignuzzi and López, 2019). Our results are consistent with research indicating that DVG protection depends on induction of innate responses (Vignuzzi and López, 2019). For example, influenza lacking the NS1 protein was studied as a potential vaccine and antiviral (Vignuzzi and López, 2019). Our results are consistent with research indicating that DVG protection depends on induction of innate responses (Vignuzzi and López, 2019).

In the last 2 years, the challenges of antiviral drug development have become clearer than ever. The mutational plasticity of RNA viruses can render most drug and even antibody therapies ineffective. Vaccinations, the only available tool to prevent viral infection, harness the natural defenses of the body. No similar approach is available for prophylaxis or treatment. We propose that eTIP1s provide an alternative strategy with potential benefits over conventional pharmaceuticals. While viruses can become resistant to currently available antivirals (Deng et al., 2021; Gonçalves et al., 2021; Abdool Karim and de Oliveira, 2021; Lundstrom, 2020; Zhou et al., 2021), eTIP1s induce a multicomponent antiviral response, in the form of diverse ISGs, that is unlikely to result in resistance by viral mutation. Mutation of the innate response by eTIP1 replication may enhance development of adaptive immunity, thus providing long-term protection.

Our eTIP1 design has several safety features. First, the PV backbone is non-pathogenic and can be administered non-invasively. Modification of the eTIP1 genome, based on improvements in the live-attenuated PV vaccine (Van Damme et al., 2019; Konopka-Anstadt et al., 2020; Konz et al., 2021; Yeh et al., 2020), may enhance their safety. Second, eTIP1 replication is limited to a few cells near the site of inoculation and is detectable by 24 h (not shown). It elicits a non-cell-autonomous protective immune response without disease symptoms by recruiting immune cells that circulate throughout the respiratory tract (Figures 5D and S6). Third, because eTIP1 replicates and self-amplifies in the few cells it enters, it requires less RNA to induce full non-cell-autonomous antiviral protection. eTIP1s seem to induce a more potent responses than PV1 (Figure 3F), suggesting that the capsid proteins modulate IFN responses. Finally, the nanoparticle formulation of the eTIP1 circumvents concerns that pre-existing immunity will prevent repeated administration.

eTIP1 protective action also induces a balanced and non-detrimental antiviral state. Such a state is not seen with WT viruses, likely because all viruses evolved mechanisms to suppress antiviral responses. In addition, there was no weight loss or signs of distress in the animals treated with eTIP1s, and even those co-infected with SARS-CoV-2 showed no signs of disease. Animals infected with SARS-CoV-2 can be protected by IN administration of IFN-I (Bessiére et al., 2021; Hoagland et al., 2021). However, antiviral treatments relying on administration of IFNs or dsRNA mimics (e.g., poly(I:C) or 5′triphosphate dsRNA; Caskey et al., 2011; Fitzpatrick et al., 2020; Furio et al., 2009; Gilfoy and Mason, 2007) cause side effects (e.g., fever, headache, fatigue, arthralgia, and myalgia; Channappa-var et al., 2016). We propose that eTIP1s achieve balanced regulation of innate responses and avoid the detrimental effects of IFN treatment.

One IN dose of eTIP1s provide a powerful prophylactic and therapeutic weapon in the COVID-19 pandemic and future respiratory diseases, including influenza and common cold, and other enterovirus diseases. In SARS-CoV-2 infection in mice, eTIP1s appear to thwart a proinflammatory response by SARS-CoV-2, thereby preventing damage to the lung and brain (Figure 5). The genes induced by eTIP1 are not proinflammatory but antiviral (Figures 3). Thus, eTIP1s block SARS-CoV-2 replication, an important first step in controlling disease, and redirect the host response from inflammatory to antiviral, which should increase the protection from disease. We also found that eTIP1s protect animals already infected with SARS-CoV-2 from disease and recruit immune cells, including eosinophils, plasmacytoid dendritic cells, and neutrophils (Figure 5). eTIP1 treatment reduced recruitment of myeloid subset CD11b+/CD11c+/Ly6C+ monocyte into the lung in infection (Figure 5D), possibly limiting production of proinflammatory cytokines. Thus, eTIP1s mediate antiviral immunity and prevent inflammation. Surprisingly, when eTIP1s are administered 24 h post-infection with SARS-CoV-2, they protect mice from infection and increase high-titer protective neutralizing antibodies (Figures 6C and 6D). We speculate that the IFN-dependent antiviral state produced by eTIP1s enhances the presentation of even small amounts of SARS-CoV-2 antigens to the adaptive immune system. eTIP1s act as an antiviral and a vaccine adjuvant, leading to short- and long-term protection.

SARS-CoV-2 infections result in heterogeneous outcomes, ranging from no symptoms to severe disease and death. The reasons are unclear. The main hypothesis is that a dysregulated immune response, probably early on, leads to systemic hyperinflammation (cytokine storm) that may drive acute respiratory distress syndrome (ARDS) and multi-organ damage in severe disease (Darif et al., 2021; Que et al., 2021; Zhou et al., 2020). Important questions are whether SARS-CoV-2 infection induces long-lasting protective immunity and if an initial infection protects from recurrent disease. In macaques, SARS-CoV-2 infection results in protective immunity when the animals are challenged soon after resolution of the primary infection (Bao et al., 2020; Chandra et al., 2020; Deng et al., 2020). However, reinfection in humans has been reported a few months after initial infection, challenging the idea of long-lasting protective immunity (Cohen et al., 2021; Gupta et al., 2020; Murillo-Zamora et al., 2021). Thus, dysregulation of the immune response by SARS-CoV-2 may impair long-lasting immunity after infection. Our data suggest that eTIP1 co-infection circumvents this
limitation, eTIP1s modulate the immune response to SARS-CoV-2 infection by activating IFN-I responses (Figures 3 and 5D); this seems to counteract the suppressive effects of SARS-CoV-2, allowing production of neutralizing antibodies even if viral antigens are scarce.

We observed that differences in virus replication between eTIP1-treated and untreated animals do not change the ability of infected animals to generate protective neutralizing antibodies (Figures 6C and 6D). This suggests that the initial SARS-CoV-2 replication produces sufficient antigenic proteins to elicit protective adaptive immunity, even without SARS-CoV-2 replication and spread and that eTIP1s act as an efficient therapy against SARS-CoV-2 infection and as an adjuvant, effectively turning the infecting virus into a vaccine. We conclude that eTIP1 prevents severe disease in the short term and facilitates long-lasting adaptive immunity. This strategy may provide a solution to emerging SARS-CoV-2 variants.

Our data indicate that eTIP1s are an effective antiviral approach to harness natural immunity to fight infection in a balanced, safe, and controlled manner, utilizing the regulatory circuits that evolved to ensure protection from disease without causing self-afflicted damage (Rahim et al., 2020; Vignuzzi and López, 2019). DVG-based antiviral therapy may offer greater benefits and safety than conventional therapies, given its broad-spectrum capacity that may prevent rapid emergence of resistant variants and its short and long-lasting protective activity. These properties may contribute to fighting COVID-19 and other emerging or re-emerging viral threats.

Limitations of the study
Effective protection by eTIP1s is limited to 24–48 h before and after infection. This may reduce the usefulness of the approach to infections with a short incubation time before onset of symptoms. Our study is a proof of principle in small-animal models of infection, and more studies will be needed to understand its full therapeutic potential, given that COVID-19 characteristics are not fully replicated in mice. While we provide evidence that co-infection with a helper virus does not mobilize eTIP1 to other tissues, a more extensive trial is needed to establish the safety of this approach in the context of co-circulating enteroviruses. A particularly important question is if eTIP1 treatment can enhance disease through the induction of a “cytokine storm” (Fajgenbaum and June, 2020; Findlay et al., 2015; Yuan et al., 2021). The mechanisms and dynamics of the induction of protective responses are limited by the number of time points and type of samples analyzed. To optimize the efficacy and safety of this strategy, additional data with higher degrees of temporal and anatomic resolution are needed.

STAR METHODS

Detailed methods are provided in the online version of this paper and include the following:

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AUTHOR CONTRIBUTIONS

Y.X. designed and carried out the experiments, participated in data analysis, and wrote the manuscript. Y.S. cloned eTIP1 cDNA plasmid and generated the stable packaging cell line. G.D. developed nanostructured lipid/eTIP1 RNA complexes, C.E.G. contributed animal work and RNA-seq libraries preparation. Y.X. and P.V.L. carried out SARS-CoV-2 experiments. A.N., Y.X., and D.T. produced and purified eTIP1 particles and LNPs. Y.X. and C.T.W. carried out and analyzed the immunohistochemistry studies on tissue sections. W.S. carried out droplet RT-qPCR for gene expression analyses from tissues. Y.X., R. Aviner, and W.L. carried out RNA-seq analyses. A.C. carried out EM negative stain studies. W.Z. and J.D.E. contributed to flow cytometry analyses of lung tissues. R. Andino, J.F., R.N., J.D.E., S.B., and P.K.J. conceived and directed the project and/or wrote the manuscript.

DECLARATION OF INTERESTS

Y.X., R.N., and R. Andino have submitted a patent application. Provisional patent application: recombinant enteroviruses and uses thereof. eTIP1. US Provisional Patent Filed 7/2020. The application was accorded serial
no, 63/047,398. D.T. and R.N. are shareholders and employees of Aleph Therapeutics, Inc. E.B. is a shareholder and employee of Pine Biotech Inc.

INCLUSION AND DIVERSITY

We worked to ensure sex balance in the selection of non-human subjects. One or more of the authors of this paper self-identifies as a member of the LGBTQ+ community. One or more of the authors of this paper received support from a program designed to increase minority representation in science. While citing references scientifically relevant for this work, we also actively worked to promote gender balance in our reference list.

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### STAR★METHODS

#### KEY RESOURCES TABLE

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Antibodies**      |        |            |
| CD45                | Biolegend | Cat#103128, RRID#AB_493715 |
| CD11c               | Biolegend | Cat#117324, RRID#AB_830649 |
| Singlec-F           | BD Biosciences | Cat#552126, RRID: AB_394341 |
| Ly6G                | Biolegend | Cat#127618, RRID: AB_1877261 |
| Ly6C                | Biolegend | Cat#128036, RRID: AB_2562353 |
| CD11b               | Biolegend | Cat#101242, RRID: AB_2563310 |
| CD103               | Biolegend | Cat#121414, RRID: AB_1227502 |
| MHC-2               | Biolegend | Cat#107632, RRID: AB_2650896 |
| CD45R               | Biolegend | Cat#103210, RRID: AB_312995 |
| CD317               | Biolegend | Cat#127012, RRID: AB_1953287 |
| eFluor 506 Fix Viability | eBioscience | Cat#65-0866-14 |
| NK1.1               | Biolegend | Cat#156508, RRID: AB_2876526 |
| CD19                | Biolegend | Cat#115508, RRID: AB_313643 |
| CD3                 | Biolegend | Cat#100214, RRID: AB_493645 |
| GammaDeltaTCR       | Biolegend | Cat#118129, RRID: AB_2563356 |
| CD4                 | Biolegend | Cat#100412, RRID: AB_312697 |
| CD8a                | Biolegend | Cat#100722, RRID: AB_312761 |
| Mouse Fc block      | BD PharMingen | Cat#535141, RRID: AB_394656 |
| BD brilliant stain buffer | BD Biosciences | Cat#563794, RRID: AB_2869761 |
| anti-acetylated α Tubulin (ACTUB) | Santa Cruz | Cat#sc-23950, clone: 6-11B-1 |
| anti-SARS-CoV-2 nucleocapsid(N) | GeneTex | Cat#GTX135361, RRID: AB_2887484 |
| anti-SARS-CoV-2 spike(S) | GeneTex | Cat#GTX632604, RRID: AB_2864418 |
| Polio-3A antibody   | This paper | N/A |
| Polio-3B-Vpg antibody | This paper | N/A |
| **Bacterial and virus strains** |        |            |
| SARS-CoV-2, USA-WA1/2020 | BEI | Cat No: NR-52281 |
| SARS-CoV-2, D614G    | A Gift from Sara Sunshine, (UCSF) | N/A |
| SARS-CoV-2, B1.4.27  | A Gift from Charles Chiu lab (UCSF) | N/A |
| SARS-CoV-2, B1.1.7   | This paper | N/A |
| SARS-CoV-2, B.1.617.2 | This paper | N/A |
| PV1                 | This paper | N/A |
| CVB3                | This paper | N/A |
| EV-71               | This paper | N/A |
| EV-D68              | This paper | N/A |
| Rhinovirus 16, 1B   | This paper | N/A |
| Influenza A H1N1(PR8 strain) | A gift from the Professor Christopher Byron Brooke (University of Illinois) | N/A |
| **Chemicals, peptides, and recombinant proteins** |        |            |
| Dulbecco’s Modified Eagle’s Medium | Sigma-Aldrich | Cat#D5796 |
| Bovine albumin fraction V, 7.5% | GIBCO | Cat#15260037 |
| Formaldehyde solution | Sigma | Cat#252549 |
| Crystal violet | Sigma | Cat#C0775 |
| RPMI-1640           | GIBCO™ | Cat#21875034 |
| DMEM high glucose/F12 medium | UCSF CCF facility | N/A |

(Continued on next page)
| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| Fetal bovine serum (FBS) | Sigma | Cat# F4135 |
| Penicillin-Streptomycin | GIBCO™ |Cat#10378016 |
| Lipofectamine-2000 | Thermo Fisher Scientific |Cat#11668019 |
| DNase I | Sigma-Aldrich |Cat#A1049201 |
| ACK lysis buffer | Thermo Fisher |Cat#11668019 |
| Collagen D | Worthington Biochemical |Cat#LS004210 |
| TPCk trypsin | Worthington Biochemical |Cat# LS003741 |
| DAKO Target Retrieval Solution, Ph9.0 | DAKO Agilent |Cat# S236784-2 |
| Zeocin™ Selection Reagent | GIBCO™ |Cat# R25001 |
| Gentamicin | Sigma |Cat# G1397 |

Critical commercial assays

- KAPA Stranded RNA-Seq Kit (24 rxn) Roche Cat#KK8400
- KAPA Library Quantification Kit Roche Cat# KK4844
- KAPA SYBR FAST Bio-Rad Roche Cat#KK4608
- AMPure XP magnetic beads Beckman Coulter Cat#A63880
- T7 RiboMAX™ Express Large Scale RNA Production System Promega Cat# P1320

Deposited data

- Raw tissue RNaseq This paper SRA, BioProjects #PRJNA781226

Experimental models: Cell lines

- HeLaS3 ATCC CCL-2.2
- Calu-3 ATCC HTB-55
- HeLaH1 ATCC CRL-1958
- RD ATCC CCL-136
- Vero-E6 ATCC 1586
- MDCK ATCC CCL-34
- A549-Ace2 a gift form Peter Jackson lab (Stanford university) N/A
- Packaging cell lines (HeLaS3/P1) This paper N/A
- Vero-TMPRSS2 a gift form Whelan lab (Washington University) N/A

Experimental models: Organisms/strains

- K18-hACE2 mice, (B6. Cg-Tg(K18-ACE2) 2Prmnr/J, Heminzygous) The Jackson laboratory https://www.jax.org/strain/034860 stock number: 034860
- C57BL6TgPVR a gift form Dr. Satoshi Koike of Tokyo Metropolitan Institute for Neuroscience (“TOKYO”) N/A
- C57BL6TgPVR IFNAR-/- a gift form Dr. Satoshi Koike of Tokyo Metropolitan Institute for Neuroscience (“TOKYO”) N/A

Oligonucleotides

- The primers and probe for PV1 genomes Integrated DNA Technologies 5’-CCACATACAGACGATCCCTAC-3’, 5’-CTGCCCAATGTTGAGGTAAT-3’, and 5’-6-FAM-TCTGCCCTGCACTCTCTCCAG CTT-3’-BHQ1.
- The primers and probe for eTIP1 genomes Integrated DNA Technologies 5’-GACAGCGAAGGCAATCCA-3’, 5’-CCA TGTTAGTGCTCCATTT-3’, and 5’-HEX-ACGGAAGAG/ZE/ TGTTACCAGGCAC-3’-IABkFQ.
- NEXTFlex RNA-Seq Barcodes BIOO Scientific CAT#512914

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RESOURCE AVAILABILITY

Lead contact
Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Raul Andino (raul.andino@ucsf.edu).

Materials availability
All unique/stable reagents generated in this study are available from the Lead Contact without restriction.

Data and code availability
Raw tissue RNaseq data are publicly available. Raw tissue RNaseq data have been submitted to NCBI Sequence Read Archive Bio-Projects #PRJNA781226.

Any additional information required to reanalyze the data reported in this paper is available from the lead contact Raul Andino (raul.andino@ucsf.edu).

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Cells lines
Cells used for this study include HeLaS3 cells, Calu-3 cells, A549-Ace2, HelaS3/P1 cells, HeLa H1 cells, RD cells, Vero-E6 cells, MDCK cells, and Vero-TMPRSS2 cells.

HeLaS3 cells (ATCC, CCL-2.2), Calu-3 cells (ATCC, HTB-55), A549-Ace2 cells were cultured in DMEM high glucose/F12 medium supplemented with 10% fetal bovine serum (Sigma) and 1x penicillin/streptomycin/glutamine (100xPSG, Gibco). Packaging cell line: HelaS3 cells stable overexpression poliovirus P1 gene (HelaS3/P1) were cultured in DMEM high glucose/F12 medium supplemented with 10% fetal bovine serum (Sigma) and 1x penicillin/streptomycin/glutamine (100xPSG, Gibco) plus 0.015% Zeocin (Invitrogen). HeLa H1 (ATCC, CRL-1958) cells or RD cells (ATCC, CCL-136) were cultured in DMEM/H21 medium supplemented with 10% fetal bovine serum (Sigma) and 1x penicillin/streptomycin/glutamine (100xPSG, Gibco), African green monkey kidney Vero-E6 cell line (ATCC#1586) was maintained in Minimum Essential Medium (MEM, Gibco) supplemented with 10% fetal bovine serum (FBS, Gibco), 1% Penicillin-Streptomycin-Glutamine (Gibco) at 37 °C in a humidified 5% CO2 incubator. A549-Ace2 cells are stable expression under-CMV promoter, a gift form Peter Jackson lab (Stanford university). MDCK(ATCC, CCL-34) was maintained in Minimum Essential Medium (MEM, Gibco) supplemented with 8.5% fetal bovine serum (FBS, Gibco), 1% Penicillin-Streptomycin-Glutamine (Gibco), Vero-TMPRSS2...
cells, a gift from Whelan lab (Washington University), were maintained in DMEM high glucose/F12 medium supplemented with 1x sodium pyruvate, 1x penicillin-streptomycin-glutamine and 10% FBS.

**Plasmids**

Plasmids with cDNA were used in this study. The Mahoney strain of poliovirus Type 1 (PV1) was used as wild type PV1 in this study. Defective interference genomes (eTIP1) was used as eTIP1. (See Methods details).

**Viruses**

PV1, PV3, CVB3, EV-A71, EV-D68, Rhinovirus A16, and Rhinovirus 1B viruses are Andino lab stocks. Influenza A virus strain A/PR/8/34 (H1N1) is a gift from the Professor Christopher Byron Brooke (University of Illinois). A clinical isolates: SARS-CoV-2 (USA-WA1/2020, BEI Cat No: NR-52281), SARS-CoV-2 strain(D614G) is a gift from Sara Sunshine (UCSF). SARS-CoV-2, B1.4.27 strain is a gift from Charles Chiu lab (UCSF), SARS-CoV-2, B1.1.7, SARS-CoV-2, B.1.617.2 are Andino lab stocks (See Method details).

**Mice strains**

Mice strains used in this study include K18-hACE2 mice, C57BL6TgPVR and IFNAR from Charles Chiu lab (UCSF), SARS-CoV-2, B1.1.7, SARS-CoV-2, B.1.617.2 are Andino lab stocks. SARS-CoV-2, B1.4.27 strain is a gift from Sara Sunshine (UCSF). SARS-CoV-2 strain(D614G) is a gift from the Professor Christopher Byron Brooke (University of Illinois). A clinical isolates: SARS-CoV-2 (USA-WA1/2020, BEI Cat No: NR-52281), SARS-CoV-2 strain(D614G) is a gift from Sara Sunshine (UCSF). SARS-CoV-2, B1.4.27 strain is a gift from Charles Chiu lab (UCSF), SARS-CoV-2, B1.1.7, SARS-CoV-2, B.1.617.2 are Andino lab stocks (See Method details).

**METHOD DETAILS**

**In vitro transcription (IVT) RNAs, transfection and eTIP1 production**

To generate viruses and eTIP1 particle, T7 polymerase was used to generate in vitro transcribed (IVT) viral RNA derived from corresponding linearized prib (+) Xpa1 Mahoney or eTIP1 plasmid by Apa1. The resulting 10 μg IVT RNA of PV1 were electroporated into 8x10^6 HeLaS3 cells. And IVT RNAs of eTIP1 were electroporated into 8x10^6 packing cells line. The detail methods as below. Monolayer of HeLaS3 or Packaging cells was trypsinized and washed three times in D-PBS. Cells were resuspended in D-PBS and the number of cells were counted on a hemo-cytometer, adjusting the concentration to 10^7 cells per ml. 800 μL of cells and 10 μg IVT RNAs were transfer into a chilled 4-mm electroporation cuvette and incubated 20 min on ice. Cells were electroporated with IVT RNAs (voltage = 200 V, capacitance = 1000 μF) using Gene Pulser I (Bio-Rad) and recovered in 8 mL pre-warmed medium (Burrill et al., 2013a, 2013b). Viruses and eTIP1 were harvested at around 24 h (or total CPE) to generate P0 virus or eTIP1 stocks. P0 virus stock were amplified once in cultured HeLaS3 in 2% serum media at M.O.I = 0.1 with virus alone or co-infected with mixed virus +eTIP1 (at ratio: 1:10, 1:20, 1:50, 1:100) medium with 10^4 cells per well one day before performing the TCID50.

**Titration of virus and eTIP1 samples**

Monolayers of HeLaS3 cells in 6-well plates were infected with 250 μL of serially diluted virus samples then incubated at 37°C for about 30 min, then 1% agarose overlay were added on the top. For titer eTIP, HeLaS3 cells were grown in 48-wells plate. On the following day, 100ul, 10-folds serially diluted eTIP1 samples were added to each well. Then incubation for 1 h, 400ul regular medium was added into each well. At 8-9 h post-infection, GFP-positive cells were counted as the eTIP1 titers and represent IU/mL. To measure the EV-D68 infected samples, TCID50 were performed on RD cells. RD cells were seed to 96 wells plate in 2% FBS DMEM/H21 medium with 10^4 cells per well one day before performing the TCID50.

**Design of primers and Taqman probes (Droplet PCR)**

Primers and Taqman probes for droplet digital PCR assay were designed with PrimerQuest Tool (Integrated DNA Technologies). The primers and probe for PV1 genomes are 5'-CCACATAACAGGATCCATAC-3', 5'-GTCCGCCAGTGTGATGATAT-3', and 5'-6-FAM-TCTGCTGTCACTCTCCAGCTTT-3'-BHQ1. The primers and probe for eTIP1 genomes are 5'-GACAGGGAAGCCACTCA-3', 5'-CCATGGTCGTACGCCTCATC-3', and 5'-HEX-ACGAAAGAG/ZEN/TCGGTACCAAGGCGC-3'-IABkFQ.

Droplet digital PCR assay. 2 μL of serially diluted cDNA samples was mixed with 10 μL of 2x ddPCR super-mix for probes (Bio-Rad), 1 μL of 20x PV1 or eTIP1 primers/probe, 1 μL of 20x eTIP1 primers/probe, and 6 μL of nuclease-free water. 20 μL reaction mix of each sample was dispensed into the droplet generator cartridge, followed by droplet production with QX100 droplet generator (Bio-Rad). Then PCR was performed on a thermal cycler using the following parameters: 1 cycle of 10 min at 95°C. 60°C of 1 min for 40 cycles. Then the PCR product were read and calculated by QX100 droplet reader.

**Virus growth curve of PV1 or other wildtype viruses and co-infected replication kinetics in cell culture models**

2.5 x 10^6 HelaS3 cells were seeded in 24-well plates. On the following day, cells were washed twice with PBS and were infected with virus in 200 μL, 2% serum media at M.O.I = 0.1 with virus alone or co-infected with mixed virus +eTIP1 (at ratio: 1:10, 1:20, 1:50, 1:100)
or with eTIP1 alone at MOI is 5 (three replicate wells were used for each virus at each time point). Following an h incubation at 37°C, each well was washed twice with PBS, and cells were covered with fresh complete media. At each indicated time point, the corresponding plate was frozen at −80°C. Following three freeze-thaw cycles of the plates, standard plaque assays were performed on monolayer HeLaS3 cells grown in a 6-wells plate (~10^6 HeLaS3 cells per well). Or TCID50 for EV-D68 on RD cells.

**Purification of the eTIP1 particles**

Packaging cell lines (HeLaS3/P1) generating eTIP1 (500 ml) was harvested with 0.5% NP-40, and the sample was stored at −80°C. For virus purification, the sample was subjected to three freeze-thaw cycles. For virus precipitation, PEG 8000 was added to a final concentration of 10% and stored overnight at 4°C. The precipitated sample was pelleted by spinning at 3,500 g for 1 h. The pellet was suspended in 10 mL EB-buffer (50mM Tris pH 8.0, 300 mM NaCl, 5 mM MgCl2, 0.5% NP-40) and centrifuged at 3,500 g for 30 min at 4°C to remove cell debris and insoluble materials. The soluble fraction containing eTIP1 in the supernatant was overlayed on a 2 mL 30% sucrose cushion in EB-buffer at 105,000 g for 3 h at 4°C. The precipitated sample was pelleted by spinning at 3,500 g for 10 min at 4°C to remove insoluble material. The soluble fraction containing eTIP1 was then laid on the top of a 15%–45% sucrose gradient in EB-buffer and centrifuged at 105,000 g for 3 h at 4°C. Fractions of 1 mL size from top of the gradient was collected containing eTIP1. Two fractions from top were pooled together, and sucrose in the sample was removed using a spin desalting column (Zeba; Pierce) and buffer exchanged with PBS. eTIP1 in PBS were then concentrated using Amicon ultra device with 100 kDa MWCO. Purity and integrity of the eTIP1 were tested by SDS-PAGE, silver staining. Negative stain and electron-microscopy were used on the particle. Fractiion#5-6 is combined, titered and used to inoculate into mice.

**Infection of susceptible mice**

We followed protocols approved by the UCSF Institutional Animal Care and Use Committee (IACUC) for the mouse studies. In these experiments, 5 to 6-weeks-old C57BL6 TgPVR or 6 to 8-weeks-old C57BL6 TgPVR interferon α/β receptor knockout (IFNAR−/−) both male and female mice were used and infected under anesthesia. C57BL6 TgPVR and IFNAR−/− were kindly provided by Professor Julie Pfieffer of the University of Texas Southwestern Medical Center, and originally were generated by Dr. Satoshi Koike of Tokyo Metropolitan Institute for Neuroscience (“TOKYO”) (Ida-Hosonuma et al., 2005). For mice survival studies, mice were injected by intraperitoneal injection (IP, 100 µL per mouse), by intranasal injection (IN, 20-50ul per mouse) with serial dilutions of each virus, respectively (10 mice per group) (Nagata et al., 2004). If UV- treatment was involved, then the eTIP1 were UV-treatment for 2 h. For the influenza H1N1(PR8 strain) experiment, the mouse will weight daily. Mice were monitored twice daily for the onset of paralysis and were euthanized when death was imminent.

For protection study on poliovirus, we injected C57BL6 TgPVR mice with viral supernatant 10⁷ pfu of PV1 alone or with eTIP1 ratio at 1:1, 1:10 per mouse by IP route or 1:20 by I.N. route (PV1 is 3x10⁵ pfu per mouse), For tissue distribution studies, we injected C57BL6 TgPVR mice (3 to 5 mice per group) with 3x10⁵ pfu of PV1 virus per mouse by intranasal(IN) route. Half of the organs were collected from infected mice and homogenized in 1 mL serum-free media. Viral supernatants were collected from the tissue homogenates, following three freeze-thaw cycles, and centrifuged at 12,000 g for 30 min at 4°C. Regular plaque assays were performed on HeLaS3 cells to titer viral supernatants from tissues. For protection study on CVB3 virus, we injected Tg21 mice with viral supernatant 10⁷ pfu of PV1 alone or with eTIP1 ratio at 1:100, 1:10 per mouse by IP route 24 h or 48 h post-infection, animals were weighed and checked daily.

For the pre-treat experiments, we inoculated C57BL6 TgPVR mice with 6x10⁶ IU eTIP1 particles in PBS by intranasally. 24 h and 48 h inoculation, 3x10⁵ pfu PV1 or 10⁶ pfu influenza(A/PR8) virus was inoculated into Tg21 mice. For the therapeutics experiments on poliovirus, we inoculated C57BL6 TgPVR mice, 3x10⁵ pfu PV1 by intranasally at day 0, then inoculated with 6x10⁵ IU eTIP1 in PBS by intranasally daily from day 1 to day 5. For the therapeutics experiments on flu, 10⁵ pfu influenza(A/PR8) by intranasally at day 0, then inoculated with 30ug eTIP1 RNA at 24- or 48 h post-infection.

For tissue distribution studies, The IFNAR−/− mice were inoculated by I.M. route with 200 pfu PV1 alone or with eTIP (PV1: eTIP1 at ratio = 1:5000) per mouse (3 mice per group). Mice were euthanized with CO2, muscle, spleen, spinal cord were collected and at 1, 3, 6 days post infection. The tissues were homogenized in 1 mL Trizol reagents (Ambion). Total RNAs were extracted and treated with Dnase1(NEB). RT-qPCR were formed as droplet qPCR section.

**SARS-CoV-2 virus propagation and infection**

SARS-CoV-2 cell culture and animals works were performed in the Biosafety level 3 (BSL-3), under the guidance and protocols approved by UCSF Biosafety and Animal research committees. African green monkey kidney Vero-E6 cell line (ATCC® C1586) and Calu-3 cells (ATCC® HTB-55) was obtained from American Type Culture Collection (ATCC® C1586) and maintained in Minimum Essential Medium (MEM, GIBCO Invitrogen) supplemented with 10% fetal bovine serum (FBS, GIBCO Invitrogen), 1% Penicillin-Streptomycin-Glutamine (GIBCO Invitrogen) at 37°C in a humidified 5% CO2 incubator. A clinical isolate of SARS-CoV-2 (USA-WA1/2020, BEI Cat No: NR-52281 with D614G) or 2 or variants of concern (V.O.C), was propagated in Vero E6 cells and A549-ACE2 cells. Viral titer was quantified with plaque assay. All the infections in the context of SARS-CoV-2 were performed at biosafety level-3 (BSL-3).
For prophylactic and coinfection experiments, ~70% monolayers of Calu-3 cells (1 × 10^5 cells/well in 24-well plates) were pre-treated with eTIP1 particles with MOI = 5 for 5 h (pretreatment), then infected with SARS-CoV-2 (MOI = 0.1) for 1 h at 37 °C, the virus mixture was removed, cells were further cultured with medium. At indication time-point 16, 24, 36, 48 hpi (h post infection), supernatants were collected, and viral titers of supernatant were detected by plaque assay.

For co-infection experiment, Calu-3 cells were infected with SARS-CoV-2 (MOI = 0.1) alone or co-infected with eTIP1 particles at different ratios (1:1, 1:10, 1:50) for 1 h at 37 °C, the virus mixture was removed, cells were further cultured with medium. At indicated time-points, 24, 36, 48 hpi (h post infection), supernatants were collected, and viral titers of supernatant were measured with plaque assay on Vero-E6 cells.

**Plaque assay for SARS-CoV-2**

For SARS-CoV-2 plaque assays, 80% Confluent monolayers of Vero E6 cells for (D614G) or Vero-TMPRSS2 (for V.O.C. B1.1.7 or B1.4.27), grown in 6-well plates were incubated with the serial dilutions of virus samples (250 μl/well) at 37 °C for 1 h. Next, the cells were overlayed with 1% agarose (Invitrogen) prepared with MEM supplemented containing 2% fetal bovine serum. Three days later, cells were fixed with 4% formaldehyde for 2 h, the overlay was discarded, and samples were stained with crystal violet dye.

**Mouse experiments for SARS-CoV-2**

K18-hACE2 mice (Winkler et al., 2021) (The Jackson laboratory, https://www.jax.org/strain/034860, stock number: 034860, B6. Cg-Tg(K18-ACE2)2Prlm/J, Hemizygous). The K18-hACE2 mice were inbred and housed in UCSF animal facility. All the experiments we performed, the mice were under anesthesia and at the BSL3 level. For prophylactic protection experiments, 30 μg eTIP1 RNA with lipofectamine-2000 were inoculated into mice intranasally, 18-20 h later, K18-hACE2 mice were anesthetized with isofluorance and inoculated with 6×10^3 pfu of SARS-COV-2 intranasally, mice were monitored daily, and weight was measured at indicated time-points. For therapeutic experiments for eTIP1, K18-hACE2 mice were anesthetized and infected with SARS-CoV-2 or variants of concern (V.O.C), then the eTIP1 RNA were delivered with lipofectamine-2000 at 24 hour, 48 hour, 24 and 48 h. For tissue distribution, mouse will be sacrificed at indication time-points, the tissues will be collected and homogenized with 1ml 2% FBS MEM medium with gentleMACS - C tubes (Miltenyi Biotec Catlog# 130-093-237), plaque assays were performed for titration of the virus. For RNA extraction, the 100mg tissues were homogenized in 1ml trizol reagents (Ambion) with gentleMACS - M tube (Miltenyi Biotec, Catlog# 130-093-236), RNA were treated with DNase1, 1mg total RNA were used to make cDNA by lscript (Bio-Rad). DNase1 treated total RNA, then poly A beads purification (Bio Scientific), then the RNASeq libraries will be prepared with the KAPA biosystem (KAPA Stranded RNA-Seq Library Preparation Kit).

**Examining production of neutralizing antibodies**

SBackspace6 to 8 weeks old C57BL6 TgPVR mice were inoculated with 30 μg eTIP1 RNA with lipofectamine-2000 or empty lipofectamine-2000 were inoculated into mice intranasally, 20-24 h later, mice were anesthetized with isofluorance and inoculated with 10^2, 10^3, 10^4 pfu of SARS-COV-2 (B1.1.7 strain) intranasally, for detect neutralizing antibody, blood were collected at 21days post-infection, sera were collected, then the neutralization antibody titer were determined by PRINT assay on Vero-TMPRSS2 cells.

For viral loads experiments on B.1.1.7 (Figure 6), C57B/6j Mice were sacrificed, and lungs were collected at 3 days post-infection, weighted and homogenized with 1ml 2% FBS MEM medium with gentleMACS - C tubes (Miltenyi Biotec Catlog# 130-093-236), RNA were treated with DNase1, 1mg total RNA were used to make cDNA by Iscript (Bio-Rad), DNase1 treated total RNA, then poly A beads purification (Bio Scientific), then the RNASeq libraries will be prepared with the KAPA biosystem (KAPA Stranded RNA-Seq Library Preparation Kit).

**Hematoxylin Eosin (H&E), Immunofluorescence (IF) staining on tissue section and imaging**

For pathology and immuno-fluorescence, mouse tissues were collected and fixed in the 4% PFA, then the tissues were embedding with paraffin and wax, then processed and the tissue samples will cut at 5 μm sections. At indication time-point 16, 24, 36, 48 hpi (h post infection), supernatants were collected, and viral titers of supernatant were measured with plaque assay on Vero-E6 cells.

For pathological and immuno-fluorescence, mouse tissues were collected and fixed in the 4% PFA, then the tissues were embedding with paraffin and wax, then processed and the tissue samples will cut at 5 μm, and H&E staining were performed at the Gladstone Histology and light core.

Deparaffinization, rehydration, and HIER were performed on an a ST4020 small linear stainer (Leica). For deparaffinization, slides were baked at 70 °C for 1–1.5 h, followed by rehydration in descending concentrations of ethanol (100% twice, 95% twice, 80%, 70%, ddH2O2O twice; each step for 30 s). Washes were performed using a Leica ST4020 Linear Stainer (Leica Biosystems, Wetzlar, Germany) programmed to three dips per wash for 30 s each. H&E staining were performed. For I.F., HIER was performed in a Lab VisionTM PT module (Thermo Fisher) using Dako Target Retrieval Solution, pH 9 (S236784-2, DAKO Agilent) at 97 °C for 10 min and cooled down to 65 °C. After further cooling to room temperature for 30 min, slides were washed for 10 min three times in Tris-Buffered Saline (TBS), containing 0.1% Tween 20 (Cell Marque; TBST). Sections were then blocked in 5% normal donkey serum in TBST at room temperature for 1 h, followed by incubation with primary antibodies in the blocking solution. After one overnight incubation of primary antibodies in 4 °C, sections were washed three times with TBST and stained with the appropriate secondary antibodies in PBS with 3% bovine serum albumin, 0.4% saponin, and 0.02% sodium azide at room temperature for 1 h. Following this, sections were washed three times with TBST and mounted with ProLong Gold Antifade mounting medium with DAPI (Invitrogen). The primary antibodies and final titrations used were mouse anti-acetylated α Tubulin (ACTUB) (1:300; Santa Cruz sc-23950), rabbit anti-SARS-CoV-2 nucleocapsid(N) (1:1000; GeneTex GTX135361), and mouse anti-SARS-CoV-2 spike(S)
Mouse lung histological analysis
Paraffin-embedded lung tissue blocks for mouse lungs were cut into 5 μm sections. Sections were stained with hematoxylin and eosin (H&E) and analyzed. Digital light microscopic scans of whole lung processed in toto were examined by an experienced veterinary pathologist. Hematoxylin Eosin (H&E) stained sections of lung from K18 hACE2 mice were examined by implementing a semi-quantitative, 5-point grading scheme (0 - within normal limits, 1 - mild, 2 - moderate, 3 - marked, 4 - severe), that considers four different histopathological parameters: 1) perivascular inflammation 2) bronchial or bronchiolar epithelial degeneration or necrosis 3) bronchial or bronchiolar inflammation and 4) alveolar inflammation. These changes were absent (grade 0) in lungs from vehicle and eTIP1 treated uninfected mice from groups that were utilized for this assessment (Meyerholz and Beck, 2020; White et al., 2021).

Flowcytometry analysis of the immune cells profiling from lung
For the prophylactic experiments for poliovirus challenging, we inoculated 6 weeks old Tg21 PVR mice with or without 30ug eTIP1 RNA with lipofectamine 2000 by intranasally. 24 h inoculation. Then mice were infected with SARS-CoV-2 intranasally, please refer to the SARS-CoV-2 mouse section. Then mice were euthanized with CO2 and perfusion with PBS. The full lungs were removed, washed twice with PBS and RPMI-1640(GIBCO). Then we cut the whole lungs as small pieces put with 4ml digestion buffer (RPMI-1640+10mg/mL Collagen D +10mg/mL DNase1, 5% FBS) for 30mins, then the tissues were minced with 10ml syringe, and pass through with 70μm cell-strainer. Cells then spin it down and wash twice with D-PBS at 650 g at 4°C for 5mins. The red cells were lysis with ACK buffer (Thermo Fisher, Cat#A1049201) for 2mins. Cells then spin it down and wash twice with D-PBS at 650 g at 4°C for 5mins. Collagen D (Worthington Biochemical, Cat# LS004210), DNase1(Sigma-Aldrich, Cat#11284932001).

Cells were stained with trypan blue and counted. 10⁶ Cells will used for full antibodies panel staining. 10⁵ cells were used for live/dead staining for 15mins (eFluor 506 Fix Viability, eBioscience, Cat#65-0866-14), single antibody or unstaining cells or all antibodies. Mouse Fc block (BD PharMingen, Cat#553141) were diluted in D-PBS. Antibodies were diluted in 1:100 in BD brilliant stain buffer (BD Biosciences, Cat#563794). After all steps staining were completed, cells were fixed with 2% PFA for 30mins, then the tissues were minced with 10ml syringe, and pass through with 70μm cell-strainer. Cells then spin it down and wash twice with D-PBS at 650 g at 4°C for 5mins. Collagen D (Worthington Biochemical, Cat# LS004210), DNase1(Sigma-Aldrich, Cat#11284932001).

Samples were resuspended in 300ul FACS buffer (D-PBS +0.2% BSA +2mMEDTA). Samples were run in the BD Aries 3 and analyzed with Flowjo software.

Antibody panels (key resources table) and gating method are described in detail in Figure S2.

mRNaseq libraries preparation and analysis
Mice tissues were collected and homogenized in 1 mL Trizol reagents (Ambion). Total RNAs were extracted, 1mg treated with DNase1(NEB). Then the mRNAs were purified by polyA beads. mRNaseq libraries were prepared by following the instruction with KAPA Biosciences. Then mRNaseq libraries were pooled and sequenced by Illumina HiSeq 4000 with single read in the UCSF core facility (Center for Advanced Technology, https://uccore.org/ucsf-center-for-advanced-technology/).

Differential gene and transcript expression analysis of mRNA-seq experiments with TopHat-Cufflinks-Cuffdiff pipeline (Trapnell et al., 2012). The Figures were plot by R, ggplot2 package. Annotation of type I ISGs was based on Liu et al. (PMID 22371602) for IFNα and Gene Ontology (GO) Biological Process “Response to interferon-beta” for IFNβ. For direct comparisons between eTIP1 particles and LNPs, transcripts were filtered for statistical significance in both datasets. Fisher enrichment analyses was performed in Perseus version 1.6.7.0, using GO annotations and the above type I ISG list. Analyses were performed separately for statistically significant transcripts that are induced (log2(FC)>0) or depleted (log2(FC)<0), and the results were filtered for intersection > 9 and enrichment > 5.

QUANTIFICATION AND STATISTICAL ANALYSIS
Data were presented as mean ± SD. Statistical analysis was performed using GraphPad Prism (GraphPad Software). Statistical significance was calculated using a two-tailed Student’s t test and p value < 0.05 was considered significant. Significance is noted with asterisks as described in the figure legends. Animal experiments were not blinded or randomized, and no animals or samples were removed as outliers from the analysis. The 50% Lethal dose (LD50) and Survival curves were compared with Log-rank (Mantel-Cox) test methods performed using GraphPad Prism (GraphPad Software). p value < 0.05 was considered significant.
Figure S1. Replication of eTIP1 and doses-dependent effect on EV-D68 replication, related to Figure 1
(A) Immunofluorescence (IF) for eTIP1 particles on infected lung cell type, Calu-3 cells at moi = 0.1. At 5 and 24 h post-infection, cells were fixed with 4% PFA and the IF were performed with polio-3A antibody (red color) (STAR Methods).
(B) eTIP1 was to infect RD cells at different multiplicity of infection (moi) ranging from 10 to 0.1, and coinfected with EV-D68 eTIP1 (moi = 0.1). eTIP1 inhibits replication on EV-D68, implicated in outbreaks of severe respiratory illness in the US in a dose-dependent manner.
Figure S2. eTIP1 protects respiratory tract and enterovirus infection and protect from disease cause by coxsackievirus B3, poliovirus and influenza virus (AIV), related to Figure 2.

(A) Intraperitoneal inoculation in immune-competent C57BL6 TgPVR mice with $10^5$ P.F.U. coxsackievirus B3 (CVB3) or co-infected eTIP1 at a ratio of 1:100 ($n = 7-9$). Survival curve. Black line represents CVB3 alone. Red line are mice infected with a mix of CVB3 + eTIP1 group. The statistical analysis of survival curves was carried out by log-rank (Mantel-Cox) test. Significance is noted with asterisks; ns, not significant.

(B) Weight lost, experiment was carried out as in Fig.S2A. blue line represents CVB3 alone, red line are mice infected with a mix of CVB3 + eTIP1 group, black line represents mock(un-infected) control.

(C) Virus loads in spleen and brain tissues of PV1-infected (black) or PV1 + eTIP1 co-infected intranasally (IN) at a ratio of 1:20. Tissues were collected at indicated times, homogenized and tittered by plaque assay. $n = 3-5$. Data was analyzed using unpaired Student’s t tests. Significance is noted with asterisks as follow: **p < 0.001; and ***p < 0.0001. Two independent experiments.

(D) Weight lost, red solid line represents influenza virus (PR8) alone, red dotted line are mice infected with a mix of PR8 + eTIP1 group, black line represents mock(un-infected) control ($n = 5-7$). Significance is noted with asterisks as follow: *p < 0.05, **p < 0.001; and ***p < 0.0001. Two independent experiments.
Figure S3. Delivery of eTIP1 using lipid nanoparticles (LNPs) and safety characterization of eTIP1 inoculated intramuscularly in highly susceptible mice, related to Figure 3

(A) eTIP1 RNA transfection and expression. Schematic representation of eTIP1 RNA/LNP complex. eTIP1 RNA expression in cell culture model. 2μg eTIP1 RNAs were transfected into HelaS3 cells with lipofectamine 2000, then immunofluorescence (IF) staining with the poliovirus 3A antibody at 8 h post-transfection. Poliovirus 3A protein staining (Red), eTIP1 RNA (green), the nuclear (blue).

(B) eTIP1 replication is restricted to the site of inoculation (leg muscle) even in the presence of WT polio helper virus. 6 to 8-weeks-old C57BL6 PVR interferon α/β receptor knockout (IFNAR⁻/⁻) mice were infected with 200 P.F.U. wildtype poliovirus or co-infected with mixed PV1+ eTIP1 at ratio 1:5000 by intramuscular (I.M.) route. eTIP1 inhibits wildtype virus spread into central neuron system (CNS). However, eTIP1 replication is limited to the site of inoculation (muscle) but not spread and replicates in spleen and spinal cord. RNA genome copies for eTIP1 and PV1 by digital droplet RT-qPCR. y axis represents RNA genome copies per 1 mg total RNA. Black line with square represents as PV1 genome copies in wildtype virus alone group. Red line with square represents as PV1 genome copies in co-infection group. (n = 3). Two tails multiple-t tests. Significance is noted with asterisks as follow: *p < 0.05; and ***p < 0.001.

(C) Survival curves of mice inoculated intramuscularly with eTIP1. 6 to 8-weeks-old C57BL6 PVR interferon α/β receptor knockout (IFNAR⁻/⁻) mice were infected with 2 x 10⁸ P.F.U of encapsidated biological particles eTIP1.
Figure S4. RNA-seq analysis of mice treated with eTIP1/LNP and the role of Type I IFN, related to Figure 3

(A) Gene Ontology (GO) annotations with manual validation were used to calculate an average fold change from mock for all host mRNA induced during eTIP1 inoculation. ANOVA p value indicated for each.

(B) eTIP1 fails to protect against poliovirus (PV1) intraperitoneally (iP) in mice lacking a type I interferon response (IFNAR\(^{-/-}\)). IFNAR\(^{-/-}\) mice were infected with 5\(\times\)10\(^4\) pfu PV1 alone or co-infected with mixed PV1 + eTIP1 at a ratio of 1:10 by iP route. Black line represents PV1 alone. Red dash line represents co-infected mixed PV1 + eTIP1 group (n = 7-8). Data presented was collected from two independent experiments. The comparison of survival curves was performed by log-rank (Mantel-Cox) test. ns, not significant.
Figure S5. Only replication-competent eTIP1 protects against SARS-CoV-2, delivery of none-replicative RNA in LNP complex are not effective, related to Figures 4 and 5

(A) eTIP1 RNA/LNP and controls, e.g., UV-inactivated eTIP1 RNA, replication incompetent eTIP1 with a large deletion (~2kbps) of the most-3' region of the eTIP1 genome, or poly IC, were inoculated into K18-hACE2 mice by the intranasal route, and at 24 h mice were infected with 10^4 P.F.U. SARS-CoV-2 D614G. B by IN route (n = 4-5). Virus titers in lungs were determined by plaque assays 3 days after infection.

(B) K18-hACE2 mice were infected by the intranasal route with 10^4 P.F.U. SARS-CoV-2 Delta variant by IN route (n = 4-5). Mice were inoculated with eTIP1 RNA/LNP (30µg) or mock (empty LNPs) 72 h post-infection. Virus titers in lungs at 5 days post infection were determined by plaque assays. Direct animal observation suggested that eTIP1 inoculation during ongoing infection did not enhance signs of disease stress or augmented lethality of SARS-CoV-2.
Figure S6. Gating of the flow cytometry to identify subset immune cells from the lung of eTIP1 inoculated mice, related to Figure 5.

(A) Mice were perfused with PBS, lungs were collected, dissociated, red cells were lysed, single cells from each lung were obtained. Cells were counted, stained with a flow cytometry antibody panels. Cells were then fixed with 1% PFA overnight, then washed two times with DPBS. Samples resuspend in FACS buffer and analyzed FACS machine. 200K events were collected from each sample. Data were analyzed by flowjo by sequential gating. For the first antibody panel (Myeloid Cells), from single and live cells: CD45+ cells represent as immune-cells. CD45+/SiglecF+ is eosinophils (EOS). CD45+/SiglecF+/CD11C+ is alveolar macrophage (AM). CD45+/SiglecF-/CD317+/CD45R+ is pDC. CD45+/SiglecF-/CD45R-/LY6G+ is neutrophils (Neut). CD45+/SiglecF-/CD45R-/LY6G-/LY6C+/MHC2+ is LY6C+ positive monocytes (LY6C+ Mono). CD45+/SiglecF-/CD45R-/LY6G-/LY6C+/MHC2-/CD11b+ is LY6C+ CD11b+ positive monocytes (LY6C+ CD11b+ Mono). CD45+/SiglecF+/CD45R+/LY6G-/LY6C+/MHC2-/CD11b+ positive monocytes (LY6C+ CD11b+ CD11C+Mono).
CD45+/SiglecF-/CD45R-/LY6G-/LY6C+/MHC2+/CD11b+/−/CD11C+ is LY6C+ CD11b+ CD11C+ positive monocytes (LY6C+ CD11b+ CD11C-Mono), CD45+/SiglecF-/CD45R-/LY6G-/LY6C+/MHC2+/CD11+ is MHC2+ CD11+ dendritic cells (DCs), CD45+/SiglecF-/CD45R-/LY6G-/LY6C+/MHC2+/CD11+/CD103+-/Cd11b+ is LY6C+/MHC2+ CD11+/CD103+ /Cd11b dendritic cells (DCs).

For the 2nd antibody panel (Lymphocytes Cell gating), from single and live cells: CD45+ cells represent as immune-cells. NK1.1+ positive cells is NK cells. NK1.1−/CD19+ cells is B cells, NK1.1−/CD19−/gd TCR+ cells is Gammadelta TCR cells, NK1.1−/CD19−/gd TCR−/CD3+/CD4+ is CD4+ T cells, NK1.1−/CD19−/gd TCR−/CD3+/CD8+ is CD8+ T cells.

Then cells numbers were calculated, data represented as cell counts group. Unpaired Student’s t tests. *p < 0.05 as significant. **p < 0.01, ***p < 0.001. ns, no significant.

(B) Immune cell profiling (flow cytometry) of lung of mice infected intranasally with eTIP1 RNA/LNPs or mock infected. Data are represented as the number of CD45+ specific immune cells per 200,000 total cells obtained from mouse lungs (n = 3-8). Data are from two independent experiments. Unpaired Student’s t tests. Significance is noted with asterisks as follow: n.s., not significant.