

Engineering protease-resistant peptides to inhibit human parainfluenza viral respiratory infection

Victor K. Outlaw^{1†}, Ross W. Cheloha^{1†}, Eric M. Jurgens^{2†}, Francesca T. Bovier^{2,3,4}, Yun Zhu^{2,4,5}, Dale F. Kreitler¹, Olivia Harder⁶, Stefan Niewiesk⁶, Matteo Porotto^{2,3,4}, Samuel H. Gellman^{1*}, Anne Moscona^{2,4,7,8,*}

¹ Department of Chemistry, University of Wisconsin, Madison, Wisconsin, 53706, United States

² Department of Pediatrics, Columbia University Vagelos College of Physicians & Surgeons, New York, New York, 10032, United States

³ Department of Experimental Medicine, University of Campania 'Luigi Vanvitelli', Italy

⁴ Center for Host-Pathogen Interaction, Columbia University Vagelos College of Physicians & Surgeons, New York, New York, 10032, United States

⁵ Beijing Pediatric Research Institute, Beijing Children's Hospital, Capital Medical University, Beijing 100045, China.

⁶ Department of Veterinary Biosciences, College of Veterinary Medicine, Ohio State University, Columbus, USA.

⁷ Department of Microbiology & Immunology, Columbia University Vagelos College of Physicians & Surgeons, New York, New York, 10032, United States

⁸ Department of Physiology & Cellular Biophysics, Columbia University Vagelos College of Physicians & Surgeons, New York, New York, 10032, United States

ABSTRACT: The lower respiratory tract infections affecting children worldwide are in large part caused by the parainfluenza viruses (HPIVs), particularly HPIV3, along with human metapneumovirus and respiratory syncytial virus, enveloped negative-strand RNA viruses. There are no vaccines for these important human pathogens, and existing treatments have limited or no efficacy. Infection by HPIV is initiated by viral glycoprotein-mediated fusion between viral and host cell membranes. A viral fusion protein (F), once activated in proximity to a target cell, undergoes a series of conformational changes that first extend the trimer subunits to allow insertion of the hydrophobic domains into the target cell membrane, and then refold the trimer into a stable postfusion state, driving the merger of the viral and host cell membranes. Lipopeptides derived from the C-terminal heptad repeat (HRC) domain of HPIV3 F inhibit infection by interfering with the structural transitions of the trimeric F assembly. Clinical application of this strategy, however, requires improving the *in vivo* stability of antiviral peptides. We show that the HRC peptide backbone can be modified via partial replacement of α -amino acid residues with β -amino acid residues to generate α/β -peptides that retain antiviral activity but are poor protease substrates. Relative to a conventional α -lipopeptide, our best α/β -lipopeptide exhibits improved persistence *in vivo* and improved anti-HPIV3 antiviral activity in animals.

Introduction

Human parainfluenza viruses (HPIV1-4) are paramyxoviruses that cause human respiratory diseases, including 30-40% of childhood croup, bronchitis, bronchiolitis, and pneumonia. Infants, children, and immune-compromised individuals are particularly vulnerable to HPIV infection^{1,2}, and no FDA-approved vaccines or targeted antiviral therapeutics are currently available for any of the HPIV serotypes. Vaccines for parainfluenza have long been under study, and a limited number of vaccines are in Phase I clinical trials^{3,4}; however, no vaccine is close to FDA approval, and even this eventual development will not eliminate HPIV3 disease in the foreseeable future. While corticosteroids have decreased hospitalizations for HPIV1-associated croup⁵, HPIV2 and HPIV3 infections remain untreatable, and HPIV3 is responsible for more hospitalizations than HPIV1 and 2 combined.⁶

As for other paramyxoviruses, HPIV3 infection is initiated by the fusion of viral and host cell membranes, mediated by attachment (HN for HPIV3; H or G for related paramyxoviruses) and fusion (F) glycoproteins, that form the fusion complex required for entry.⁷⁻¹⁷ HPIV3 HN attaches to the host cell by engaging a surface receptor. The homotrimeric HPIV3 F, initially anchored to the viral envelope by transmembrane segments and adjacent C-terminal heptad repeat (HRC) domains in a metastable prefusion conformation, is then activated by the receptor-engaged HN and undergoes a sequence of structural transitions. First, N-terminal heptad repeat (HRN) domains within F extend, and fusion peptide segments insert into the host cell membrane to form a transient prehairpin intermediate. Subsequent refolding of F causes the HRN and HRC domains to form a highly stable six-helix bundle (6HB), a process that is coupled to membrane fusion.

the appearance of strong CD signals that are consistent with formation of helical co-assemblies (Figure 2A).^{33,36-38}

Variable-temperature CD measurements can be used to assess the stability of 6HB assemblies.²⁷ The co-assembly formed between HPIV3 HRN and α -VI exhibited an apparent thermal denaturation temperature ($T_{m,app}$; mid-point of the denaturation process) of 88 °C (Figure 2B). All α/β -peptides formed co-assemblies with HPIV3 HRN that denatured at lower temperatures relative to the HPIV3 HRN+ α -VI co-assembly (Table 1). Co-assemblies with α/β -peptides containing β residues throughout the entire sequence (α/β -VI-1, α/β -VI-2, α/β -VI-3 and α/β -VI-4) were less stable (lower $T_{m,app}$) than co-assemblies of α/β -peptides lacking β residues near the N-terminus (α/β -VI-5, α/β -VI-6, α/β -VI-7 and α/β -VI-8). Among this latter group, the α/β -peptides that contained cyclic residues (α/β -VI-6 and α/β -VI-8) formed more stable assemblies than did the analogues that contained only flexible β^3 residues (α/β -5 and α/β -7). This trend is consistent with past observations regarding the stabilizing effects of replacing β^3 residues with cyclic β residues. Peptide α/β -VI-8, which contains six cyclically-constrained β residues and two β^3 -hGlu (β^3E) residues at heptad positions *c* and *f* within the C-terminal segment, formed the most stable co-assembly with HPIV3 HRN and was selected for further characterization.

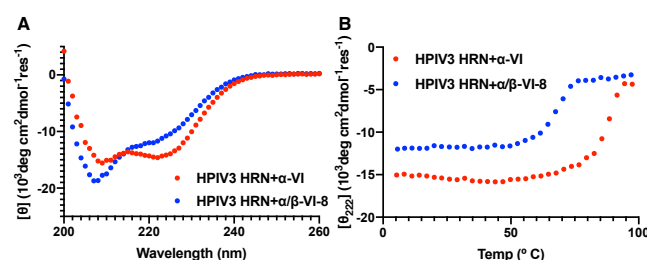


Figure 2. Circular dichroism studies of HPIV3 HRN + VI variant co-assemblies. (A) Circular dichroism (CD) spectra of 1:1 mixtures of α -VI (red) and α/β -VI-8 (blue) with HPIV3 HRN at 50 μ M in PBS. (B) Temperature-dependent denaturation of co-assemblies formed by 1:1 mixtures of α -VI (red) and α/β -VI-8 (blue) with HPIV3 HRN at 50 μ M in PBS.

Table 1. Apparent melting temperatures ($T_{m,app}$) of co-assemblies formed between VI variants and HPIV3 HRN

Inhibitor	$T_{m,app}$ (°C)
α -VI	88
α/β -VI-1	52
α/β -VI-2	51
α/β -VI-3	50
α/β -VI-4	48
α/β -VI-5	56
α/β -VI-6	60
α/β -VI-7	53
α/β -VI-8	67

Crystal structure of α/β -VI-8 co-assembled with the HRN domain of HPIV3 F. The crystal structure of α/β -VI-8 + HPIV3 HRN was determined (Figure 3B) and shows a 6HB assembly similar to

that observed within the HPIV3 F protein in the post-fusion conformation.⁴² Three copies of α/β -VI-8 are arrayed around the core formed by three HRN peptides. Each α/β -VI-8 molecule displays an α -helix-like conformation spanning residues 457–484 (numbering based on the HPIV3 F sequence); each turn of this helix, however, contains one more carbon atom relative to an authentic α -helical turn. The extended conformation of the N-terminal eight residues of α/β -VI-8 matches the corresponding segment of full-length F in the post-fusion state⁴² and of α -VI + HPIV3 HRN.²⁷ All eight β residues of α/β -VI-8 align along one side of the helix, and none of the β residue side chains contacts the HRN core (Figure 3C-F). Thus, the α/β -VI-8 + HPIV3 HRN co-crystal structure shows that our design goal of sequestering β residues away from the HRC-HRN interface was achieved.

Inhibition of HPIV3 F-mediated cell-cell fusion by α/β analogues of α -VI. We evaluated inhibition of fusion mediated by HPIV3 F co-expressed with HPIV3 HN using a cell–cell fusion assay based on β -galactosidase complementation (Supplemental Figure 1).^{43,44} While we anticipated that maximum inhibitory potency in this assay would require lipid conjugation of HRC-based peptides, in preliminary studies we compared the eight α/β -peptide analogues of α -VI (Figure 1A) without lipid modification. The trends for cell–cell fusion inhibitory potency were similar to those observed in the variable-temperature CD studies. The α/β -peptides containing β residues throughout the entire sequence failed to block cell–cell fusion (not shown). Among the four α/β -peptides lacking β residues in the N-terminal segment, the two containing cyclic residues (α/β -VI-6 and α/β -VI-8) were the most potent inhibitors. Neither, however, matched α -VI as an inhibitor of F-mediated fusion. The α/β -peptides containing only β^3 residues (α/β -VI-5 and α/β -VI-7) were less effective than the analogues containing cyclic β residues. We focused on α/β -VI-8 for subsequent studies involving lipid-conjugated peptides.

We evaluated derivatives of α -VI and α/β -VI-8 bearing a C-terminal cholesterol appendage (Figure 4A) as inhibitors of F-mediated cell–cell fusion.⁴⁵ Each 36-residue sequence was extended at the C-terminus by the segment Gly-Ser-Gly-Ser-Gly-Cys. Reaction of the Cys thiol with a bromoacetyl derivative of cholesterol containing a 4-unit PEG spacer (PEG₄) generated the lipopeptides. The PEG₄ segment provides a flexible linker between the peptide inhibitor and the membrane-embedded cholesterol moiety and enhances inhibitory activity in HPIV3 HRC-derived peptides.^{23,25} The resulting molecules were designated “ α -VI-PEG₄-Chol” and “ α/β -VI-8-PEG₄-Chol”.

α/β -VI-8-PEG₄-Chol exhibits enhanced resistance to protease-mediated degradation. The specific sites at which α -VI and α/β -VI-8 may be proteolytically cleaved in vivo are unknown; we evaluated relative degradation in the presence of the aggressive protease trypsin to compare α -VI-PEG₄-Chol and α/β -VI-8-PEG₄-Chol (Figure 4B). The cholesterol-conjugated derivative of α/β -VI-8 was stable under conditions that rapidly caused degradation of the analogous α -lipopeptide. After one-hour incubation with trypsin at 0 °C, nearly all α -VI-PEG₄-Chol was degraded. After incubation with trypsin at 37 °C, α -VI-PEG₄-Chol could not be detected. However, α/β -VI-8-PEG₄-Chol remained intact under these conditions, which demonstrates the substantial inhibition of proteolysis provided by periodic α -to- β replacement.

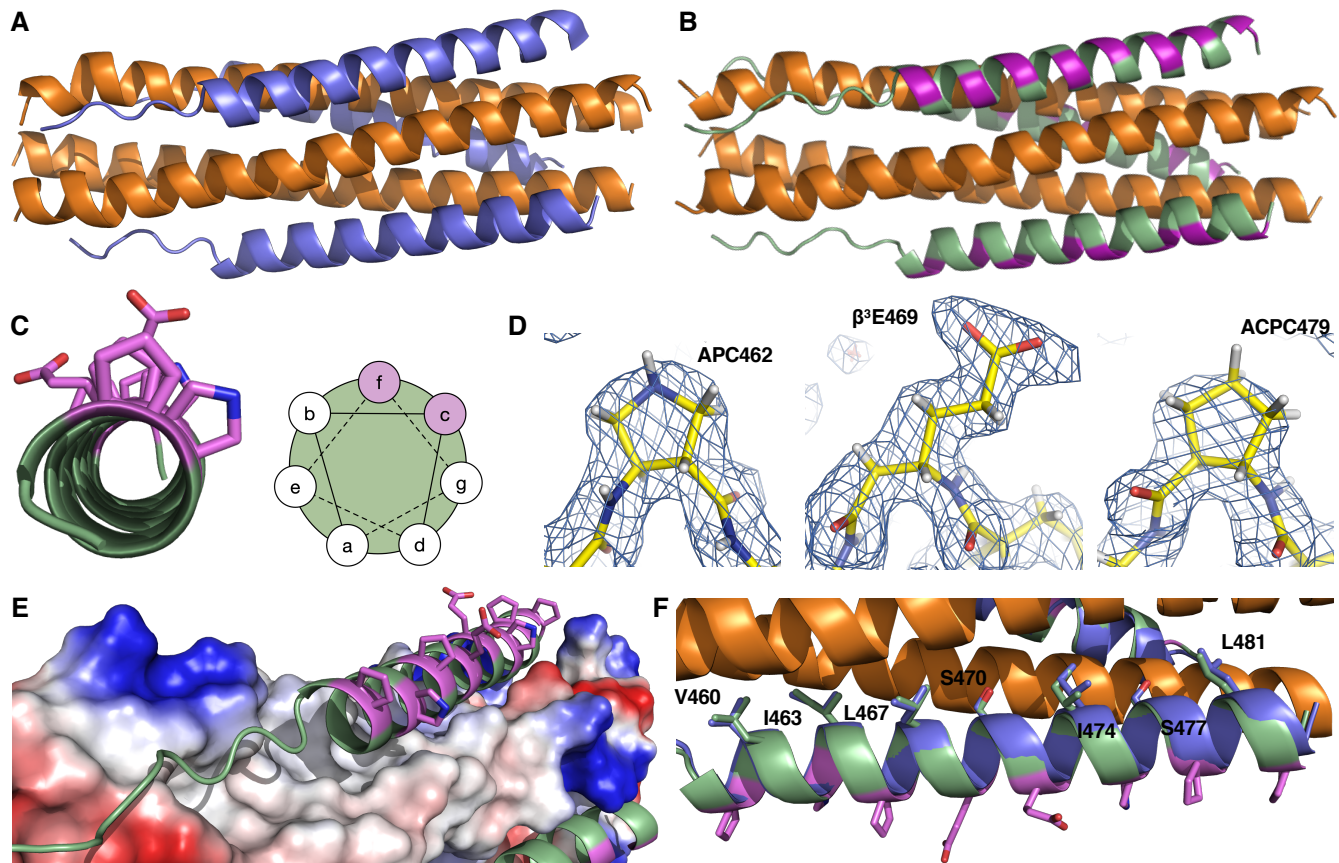


Figure 3. Crystal structure of α/β -VI-8 bound to the HRN domain of HPIV3. (A) The 6HB co-assembly formed by HPIV3 HRN (orange) and α -VI (blue) (PDB 6NYX). (B) The 6HB co-assembly formed by HPIV3 (orange) and α/β -VI-8 (green, β residues depicted as violet) (PDB 6VJO). (C) Helix-wheel diagram showing orientation of β residues (violet) at c and f positions of the heptad repeat as designed. (D) Electron-density maps of representative β residues APC462 (left), β^3 E469 (middle), and ACPC479 (right). (E) α/β -VI-8 binds within a hydrophobic groove on the HPIV3 HRN trimer (shown as map of negative (red), neutral (gray), or positive (blue) electrostatic potential). (F) Overlay of α -VI (blue) and α/β -VI-8 (green, β residues depicted as violet). Conformations of interfacial amino acid residues were conserved.

α/β -VI-8-PEG₄-Chol potently inhibits HPIV3 infection in monolayer cultured cells. We assessed the efficacy of α/β -VI-8-PEG₄-Chol at inhibiting infection by HPIV3 in cell culture (Figure 4C). We used a well-characterized HPIV3 clinical isolate virus ("CI") because we have shown that the HN-F glycoprotein fusion complexes of circulating viruses differ significantly from the HN and F of standard laboratory-grown viruses.⁴⁶⁻⁵¹ The fusion complex in laboratory culture-adapted viruses is more readily activated to promote fusion and less sensitive to inhibition by HRC peptides compared to the fusion complex in authentic circulating HPIV3, making it essential to assess inhibitors using authentic circulating HPIV3.⁴⁶⁻⁵¹ α/β -VI-8-PEG₄-Chol potently inhibited infection by HPIV3 CI, with potency similar to that of α -VI-PEG₄-Chol. Both cholesterol-conjugated peptides achieved 100% inhibition at 0.1 nM (Fig. 4C), while the corresponding unconjugated peptides displayed IC₅₀ for fusion inhibition of ~10 nM. This dramatic difference in potency between unconjugated and lipid-conjugated peptides is consistent with previous observations.²³⁻²⁶

α/β -VI-8-PEG₄-Chol potently inhibits HPIV3 infection in human airway epithelium. Human airway epithelium (HAE) cultures have proven to be an authentic model of human lung, reflecting the cell environment and selective pressure of the natural tissue^{46,48}, and the efficacy with which fusion inhibitors block viral infection in HAE seems to correlate with *in vivo* efficacy.^{23,46} We compared the antiviral effects of α -VI-PEG₄-Chol and α/β -VI-8-PEG₄-Chol in

HAE. A single dose of 1 μ M (final concentration) of either α -VI-PEG₄-Chol or α/β -VI-8-PEG₄-Chol decreased the viral titer of HPIV3 CI-eGFP by 98% at day 1 (Figure 4D). At day 2, α -VI-PEG₄-Chol treatment reduced titer by 80% while α/β -VI-8-PEG₄-Chol reduced titer by close to 98% (data not shown).

Biodistribution: α/β -VI-8-PEG₄-Chol reaches higher concentrations than does α -VI-PEG₄-Chol in serum and lung tissue. The utility of a fusion-inhibitory peptide as an antiviral agent depends on availability at the site of infection. We assessed biodistribution of α -VI-PEG₄-Chol and α/β -VI-8-PEG₄-Chol in cotton rats after either intranasal or subcutaneous administration. α/β -VI-8-PEG₄-Chol was present at higher concentration than was α -VI-PEG₄-Chol in the lung 24 hours after either intranasal or subcutaneous administration (Figure 5A). A substantially higher concentration of α/β -VI-8-PEG₄-Chol was achieved in the lung upon intranasal administration relative to subcutaneous administration. The α/β -lipopeptide also demonstrated a longer serum half-life than the α -lipopeptide after either administration route (Figure 5B). The concentration difference in the lung, particularly after intranasal administration, indicates a major advantage for the backbone-modified peptide, since persistence at the relevant tissue site should improve efficacy *in vivo*.

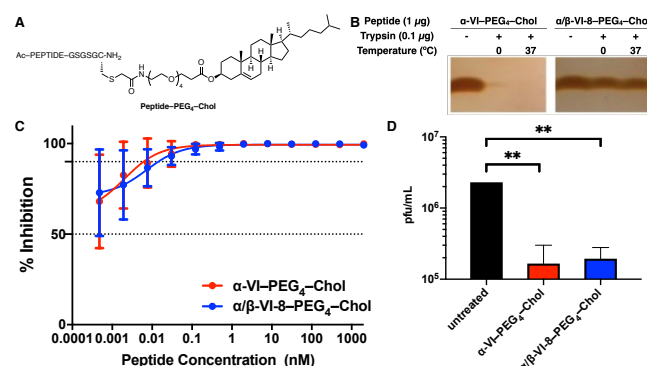


Figure 4. Proteolytic resistance and antiviral efficacy of α/β -VI-8-PEG₄-Chol. (A) Structure of cholesterol-conjugated peptides. (B) Degradation of α -VI-PEG₄-Chol (left) and α/β -VI-8-PEG₄-Chol (right) by incubation with trypsin at 0 and 37 °C. (C) Inhibition of HPIV3 infection in cell monolayers by α -VI-PEG₄-Chol (red) or α/β -VI-8-PEG₄-Chol (blue). Percent inhibition was calculated as the ratio of plaque forming units in the presence of a specific concentration of inhibitor and the plaque forming units in the absence of inhibitor. Each point represents the mean from three separate experiments $\pm$ SEM with the curve representing a three-parameter dose-response model. (D) Inhibition of HPIV3 infection in human airway epithelial (HAE) cultures by α -VI-PEG₄-Chol (red) or α/β -VI-8-PEG₄-Chol (blue). HAE cultures were treated with α -VI-PEG₄-Chol or α/β -VI-8-PEG₄-Chol at 1 μ M and infected with 5000 pfu/well of hPIV3 CI-eGFP. After 2 h, the peptides were removed, and supernatants were collected 1 day post infection. Infectious viruses released were quantified by titration. Data are depicted as the mean from three separate experiments $\pm$ SD. Data were analyzed by Student's two-tailed unpaired *t* test (***P* < 0.01).

Inhibition of HPIV3 infection *in vivo* by α/β -VI-8-PEG₄-Chol. To assess *in vivo* efficacy of the α/β -lipopeptide, cotton rats were administered 2 mg/kg of either α -VI-PEG₄-Chol or α/β -VI-8-PEG₄-Chol peptides intranasally, at 24 and 12 hours before infection and then infected with 10⁶ pfu of HPIV3 CI; the experimental scheme is shown in Figure 5C (top). Three days after infection, lung viral titers were measured. Figure 5C (bottom) shows viral titer in lipopeptide-treated animals compared to untreated animals. α/β -VI-8-PEG₄-Chol was superior to α -VI-PEG₄-Chol, with the α/β -lipopeptide reducing viral titer by over 10-fold relative to no treatment (*P* < 0.0001 using Tukey's multiple comparison test). α/β -VI-8-PEG₄-Chol reduced viral titer by 0.5 log more than the α -VI-PEG₄-Chol (*P* < 0.01). When the lower concentration of 0.4 mg/kg of peptide was administered 24 and 12 hours before infection, a less than 10-fold but nonetheless significant reduction in viral titer was achieved (Supplementary Figure 2). A comparable reduction in humans would be clinically significant.⁵²

Discussion

The results presented above suggest that modification of an HRC-derived peptide via periodic α -to- β replacement at positions selected to avoid the HRC-HRN interface in the 6HB assembly results in maintenance of fusion-inhibitory activity and inhibition of viral entry, while hindering proteolytic degradation. The resistance to proteolysis leads to enhanced persistence and antiviral performance *in vivo* for an α/β -lipopeptide relative to a comparable α -lipopeptide. This study represents, to our knowledge, the first *in vivo* evaluation of an α/β -peptide inhibitor of viral infection and lays the foundation

for a new type of agent to address HPIV3 infections in a prophylactic and/or therapeutic manner.

Because entry is the essential first step of infection by enveloped viruses, the inhibition of fusion protein structural transitions is an attractive basis for antiviral therapy.⁸ The advance we have achieved in developing α/β -VI-8-PEG₄-Chol builds upon earlier work with α -lipopeptide entry inhibitors in which we showed that specific features of amino acid sequence, identity of the lipid moiety, linker length, and solubility are all important for anti-viral efficacy.²⁰⁻²⁷ Our observations suggest that further improvements in α/β -peptide activity may be possible by pursuing two strategies. (1) Variable-temperature CD data suggest that the 6HB assembly formed by α/β -VI-8 with the HPIV3 HRN is less stable than the 6HB assembly formed by α -VI and the HPIV3 HRN. We predict that modifications to α/β -VI-8 that lead to enhanced 6HB stability will result in improved antiviral potency. (2) Although α/β -VI-8-PEG₄-Chol is highly resistant to degradation by trypsin, it is possible that the N-terminal segment of a residues in this α/β -lipopeptide or the C-terminal α residue segment used to connect the cholesterol moiety is susceptible to cleavage by other proteases in the lung. Incorporation of one or two β residues in these segments could therefore improve lipopeptide persistence *in vivo*. In this regard, we note recent results indicating that several positions in the N-terminal segment of an HRC peptide related to α -VI tolerate α -to- β substitution without loss of potency for inhibiting HPIV3 infection.⁵³

The favorable performance of our lead α/β -lipopeptide in the HAE system (Figure 4D) is promising in terms of our long-term aim to develop backbone-modified peptides for clinical use. We previously showed that the fusion complex formed by HN and F for HPIV3 differs in important ways between clinical isolate (CI) viruses (*i.e.*, those that can grow in humans) and laboratory-adapted strains.⁴⁶⁻⁵¹ Viruses bearing the fusion complex of clinical strains grow efficiently in HAE and *in vivo*, but fail to grow on immortalized cells because of the profound specificity of the CI viral fusion machinery for the authentic host.^{46-48,54} We posit that it is critical to assess antiviral agents, such as those introduced here, in the context of the viral fusion complexes that exist in human infections (*i.e.*, to assess efficacy of inhibiting viral infection in authentic lung tissues).

In the cotton rat experiment (Figure 5C), the superiority of α/β -VI-8-PEG₄-Chol over the α -lipopeptide, α -VI-PEG₄-Chol, must be attributed to the resistance of the α/β -lipopeptide to proteolysis (Figure 4B), which we propose to be the source of the significantly enhanced persistence of the α/β -lipopeptide relative to the α -lipopeptide *in vivo* (Figure 5A-B). The efficacy of the α/β -lipopeptide *in vivo* is especially striking given that the six-helix bundle assembly formed by α/β -VI-8 with HPIV3 HRN is significantly less stable than the assembly formed by α -VI, and α/β -VI-8 is less effective in inhibiting F-mediated cell-cell fusion than is α -VI (Supplementary Figure 1).

Two major factors augment the potential for clinical application of our α/β -lipopeptide inhibitors. First, the ability to administer these inhibitors intranasally offers a tractable and practical route of delivery. In fact we recently showed for another respiratory virus (SARS-CoV-2) that daily intranasal peptide administration completely blocked infection of naïve animals by infected animals.⁵⁵ These results lend support to the feasibility of using intranasal anti-HPIV3 peptides to prevent infection of vulnerable individuals (*e.g.*, during immune compromise, stem cell or organ transplant, etc.). Second, α/β -lipopeptides are chemically stable and should tolerate storage at room temperature for long periods, which means that no

cold chain would be required. These factors suggest α/β -lipopeptides could be useful even outside of advanced health care settings,

which is important given the global distribution of HPIV3 and the impact of this pathogen in developing societies.

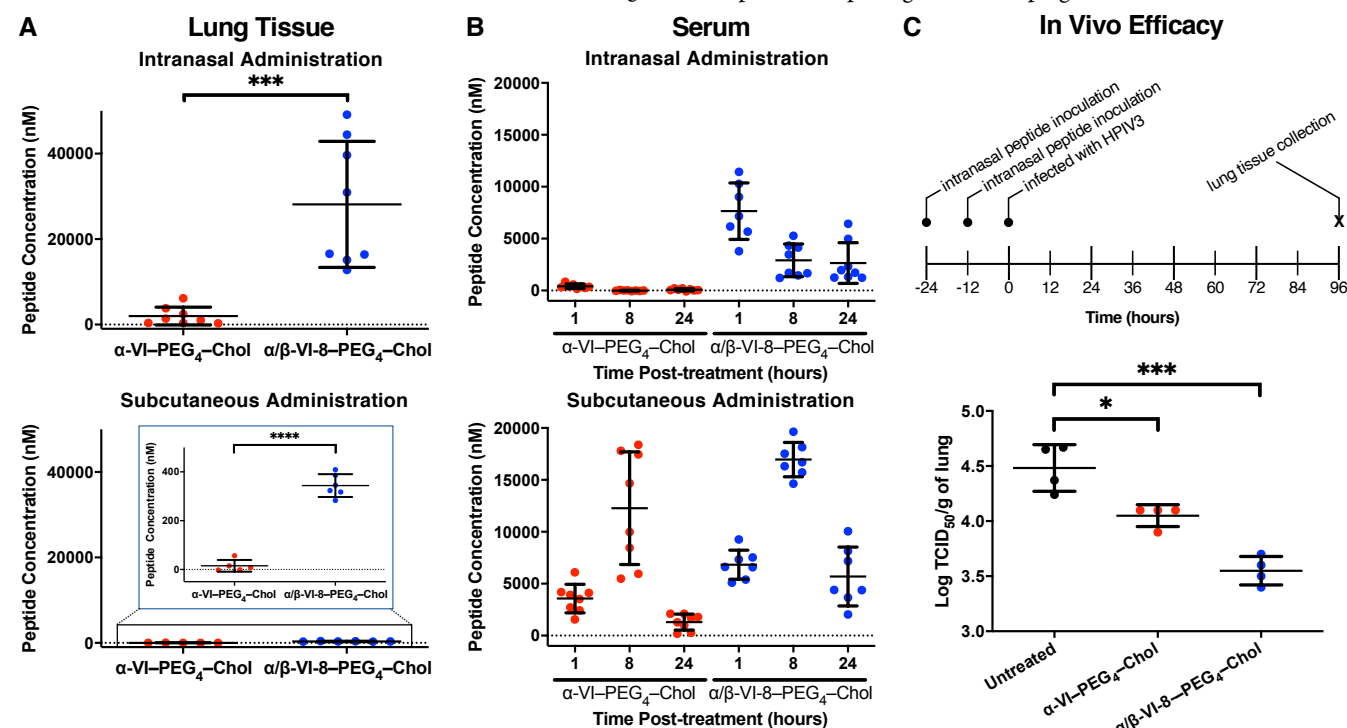


Figure 5. Biodistribution and efficacy of α/β -VI-8-PEG₄-Chol *in vivo*. (A) Biodistribution of α/β -VI-8-PEG₄-Chol in cotton rat lung tissue. Peptide concentration in the lung was determined by sandwich ELISA 24 h after intranasal (top) or subcutaneous (bottom) administration of lipopeptide. Data are depicted as the mean from eight (intranasal) or six (subcutaneous) separate experiments $\pm$ SD. Data were analyzed by Student's two-tailed unpaired *t* test (***P* < 0.001 and *****P* < 0.0001). (B) Biodistribution of α/β -VI-8-PEG₄-Chol in serum. Peptide concentration in serum was determined by sandwich ELISA 24 h after intranasal (top) or subcutaneous (bottom) administration of lipopeptide. Data are depicted as the mean from eight separate experiments $\pm$ SD. (C) *In vivo* efficacy of α/β -VI-8-PEG₄-Chol against HPIV3 infection in cotton rats with (top) schematic of experimental design and (bottom) data depicted as the mean from four separate animals $\pm$ SD. Data were analyzed by Tukey's multiple comparison test (**P* < 0.01, *****P* < 0.0001).

Materials and Methods

Cells. 293T (human kidney epithelial), CV-1 and Vero E6 cells were obtained from ATCC and were grown in Dulbecco's modified Eagle's medium (DMEM) (Gibco) supplemented with 10% fetal bovine serum and antibiotics at 37 °C and 5% CO₂. All cells tested negative for mycoplasma (MycoAlert™ Mycoplasma Detection Kit (Lonza)).

Peptide synthesis. All peptides were produced by standard Fmoc-based solid-phase methods. The cholesterol moiety was attached to the peptide via chemoselective reaction between the thiol group of an extra cysteine residue, added C-terminally to the sequence, and a bromoacetyl derivative of cholesterol, as previously described.^{23,24,56}

Structure Determination. Co-crystallization screening was performed using a 1:1 mixture of α/β -VI-8 and HPIV3 HRN. Crystal growth was promoted by vapor diffusion using hanging drop methods. Conditions that generated crystal hits were optimized for concentrations of peptide, salt, and polyethylene glycol (PEG), as well as pH, until crystals of sufficient quality for x-ray analysis were obtained. The optimized crystals diffracted to 2.0 Å. The structure was solved by molecular replacement using a truncated dimer of HPIV3 HRN+VIQKI (PDB: 6NRO) as a search model.²⁷ Specifically, chains A and B from 6NRO were truncated to include residues 153–173 (chain A; numbers correspond to full-length F) and 460–480

(chain B), and all hydrogen atoms and side chains were removed prior to molecular replacement.

Viruses. HPIV3 clinical isolate virus (CI) was obtained from the Clinical Microbiology Laboratories at New York Presbyterian Hospital and grown in human airway epithelium (HAE) at an air-liquid interface for only one passage prior to use in these experiments. Recombinant CI virus used in the cotton rat experiments was generated as described.⁴⁸

Antiviral activity against live HPIV3. Peptide activity against HPIV3 was determined by plaque reduction assays in infected cell monolayers.⁴⁶ Data were expressed as mean $\pm$ standard deviation (s.d.) (*n*=3 separate experiments).

HAE cultures. The EpiAirway AIR-100 system (MatTek Corporation) consists of normal human-derived tracheo/bronchial epithelial cells that have been cultured to form a pseudostratified, highly differentiated mucociliary epithelium closely resembling that of epithelial tissue *in vivo*. Upon receipt from the manufacturer, HAE cultures were transferred to 6-well plates (containing 0.9 mL medium per well) with the apical surface remaining exposed to air and incubated at 37 °C in 5% CO₂.

Peptide efficacy assessment in HAE cultures. HAE cultures were infected by applying 100 μ L of EpiAirway medium containing 5,000 plaque forming units (PFU) of HPIV3 with or without peptide (1 μ M final concentration) to the apical surface for 90 min at 37 °C. The medium containing the inoculum was removed, and

cultures were placed at 37 °C and fed each day with 0.9 mL medium via the basolateral surface. Viruses were harvested by adding 200 μ L medium per well to the HAE cultures' apical surface and allowing the cultures to equilibrate for 30 min at 37 °C. The suspension was then collected, and viral titers were determined as previously described.⁴⁶ Viral collection was performed sequentially with the same wells of cells on each day post infection.

Protease sensitivity of HPIV3 derived peptides. For the trypsin digestion, 1 μ g of each peptide was treated with the indicated amount of trypsin in 10 μ L of PBS. Peptide solutions were then incubated at 0 °C or 37 °C for 1 hour. Following incubation, 10 μ L of Laemmli's SDS reducing buffer was added to each solution. Samples were boiled for 10 min at 99 °C then run on a 4-20% Tris Glycine gel at 120 V. The gel was allowed to fix overnight in 0.0125% glutaraldehyde in PBS. Gels were stained using a Pierce® Silver Stain kit (Cat# 24600).

Infection of cotton rats, peptide treatment, and virus titer determination. Inbred cotton rats (*Sigmodon hispidus*) were purchased from Envigo, Inc. (Indianapolis). Both female and male cotton rats at the age of 5 to 7 weeks were used for this study. Group size was based on power analysis to determine a difference in treatment of at least one log₁₀ in viral titer after treatment with the α -VI peptide.

For intranasal (i.n.) infection, 10⁶ TCID₅₀ of recombinant CI HPIV3⁴⁸ was inoculated in phosphate-buffered saline to isofluorane anesthetized cotton rats in a volume of 100 μ L i.n. as previously described^{46,57}. To evaluate the effect of HRC peptides, animals were inoculated i.n. with peptide (2 mg/kg or 0.4 mg/kg in 100 μ L of water) 24 h and 12 h before infection. Four days after infection, the animals were asphyxiated using CO₂, and their lungs were collected and weighed. Lung tissue was minced with scissors and dounced with a glass homogenizer. Serial 10-fold dilutions of supernatant fluids were assessed for the presence of infectious virus in 48-well plates as described.⁴⁸ The TCID₅₀ was calculated as described previously. The animal experiments were approved by the Institutional Animal Care and Use Committee of The Ohio State University.

Biodistribution study methods. Animal experiment: For biodistribution experiments in cotton rats, the animals received the indicated peptides (2 mg/kg) either i.n. or s.q. in 100 μ L of diluent. At 1 h and 8 h post administration, blood was collected retroorbitally in EDTA tubes from all animals, centrifuged for 10 min at 2000 RPM, and plasma was transferred into new tubes and stored at -20 °C for use in ELISA. At 24 h after administration, the animals were euthanized, blood was collected by intracardiac puncture in EDTA vacutainer tubes, and sera were conserved at -20 °C until their use in ELISA. Organs from each animal were collected and conserved at -80 °C. The tissues were separated in two: one half frozen on dry ice and conserved at -80 °C for ELISA.

Enzyme-linked immunosorbent assay (ELISA) for determination of peptide concentration. ELISA was performed from frozen samples as previously described.²⁸ Each organ was weighed and mixed in PBS (1:1, w/vol) using an ultra turrax homogenizer. Samples were then treated with acetonitrile/1% TFA (1:4, vol/vol) for 1 h on a rotor at 4 °C and then centrifuged for 10min at 8000 rpm. Maxisorp 96 well plates (Nunc) were coated overnight with 20 μ g/mL affinity-purified Ab (anti α -VI and anti α / β -VI-8 (custom made by GeneScript)) in carbonate/bicarbonate buffer (pH 7.4) at 4 °C. Plates were washed 3x with PBS, then blocked with PBS / 3% BSA for 2 hours and placed on ice. Serial dilutions of organ and serum samples

were prepared, and the blocking buffer was replaced by the serial dilutions of each sample added in 3% PBS-BSA (pH 7.4). The samples were incubated at RT for 2.5 hours. The plates were washed 4x with 150 μ L of PBST (0.05%), and 100 μ L / well of purified rabbit biotin-conjugated and biotin-conjugated anti- α / β -VI-8 (horse radish peroxidase [HRP]-conjugated) at 1:500 in PBS BSA (3%) were added to the plates and incubated at RT for 2 h. After washing the plates 4x with PBST and 2x with PBS, streptavidin HRP (1:1000 in PBS) was added, and the plates were incubated for 30 min at RT, washed 4x with PBST, and 3x with PBS before addition of 1-Step™ Ultra TMB-ELISA (ThermoFisher #34028). The plates were covered and monitored for development, 50 μ L of 2 M sulfuric acid was added to each well to stop the reaction, and results were recorded as absorbance at 450 nm on the Infinity PRO1000 (Tecan). Standard curves were established for each peptide using 1:2 serial dilutions from 40 nM to 0 nM (using the same ELISA conditions as for the test samples), and the detection limit was determined to be 0.15 nM.

Statistical Analysis. Data are described as mean $\pm$ s.d. unless otherwise stated. The statistical analysis performed on *in vivo* experiments for comparison of HPIV3 viral titers in cotton rat lungs were based on Tukey's multiple comparison test. All the data and statistical tests were performed in Prism 5 (GraphPad Software, La Jolla, CA).

ASSOCIATED CONTENT

Supporting Information

Supporting Information is available free of charge on the ACS Publications website at DOI:

Additional figures, method descriptions, and characterization data for all synthesized peptides (PDF)

AUTHOR INFORMATION

Corresponding Author

* am939@cumc.columbia.edu, gellman@chem.wisc.edu

Author Contributions

†These authors contributed equally.

ACKNOWLEDGMENT

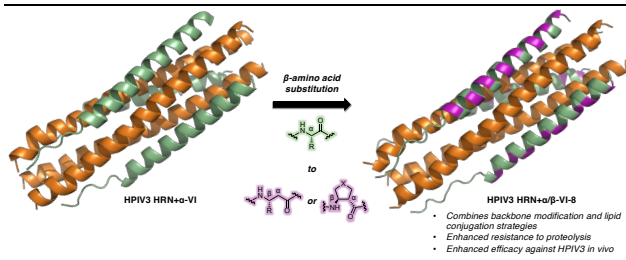
This work was supported by NIH grants R01AI114736 to A.M and R01 GM056414 to S.H.G., and by the Sharon Golub fund. V.K.O. was supported in part by NIH F32 GM122263. R.W.C. was supported in part by NIH T32 GM008349.

REFERENCES

- (1) Loughlin, G. M. & Moscona, A. The cell biology of acute childhood respiratory disease: therapeutic implications. *Pediatr Clin North Am* **53**, 929-959 (2006).
- (2) Nair, H. *et al.* Global burden of acute lower respiratory infections due to respiratory syncytial virus in young children: a systematic review and meta-analysis. *Lancet* **375**, 1545-1555, doi:10.1016/S0140-6736(10)60206-1 (2010).
- (3) Schmidt, A. C. *et al.* Progress in the development of human parainfluenza virus vaccines. *Expert Rev Respir Med* **5**, 515-526, doi:10.1586/ers.11.32 (2011).
- (4) Anderson, A. J., Snelling, T. L., Moore, H. C. & Blyth, C. C. Advances in Vaccines to Prevent Viral Respiratory Illnesses in Children. *Paediatr Drugs* **19**, 523-531, doi:10.1007/s40272-017-0257-x (2017).
- (5) Bjornson, C. L. & Johnson, D. W. Croup. *Lancet* **371**, 329-339, doi:10.1016/S0140-6736(08)60170-1 (2008).

- (6) Weinberg, G. A. *et al.* Parainfluenza virus infection of young children: estimates of the population-based burden of hospitalization. *J Pediatr* **154**, 694-699, doi:10.1016/j.jpeds.2008.11.034 (2009).
- (7) Yin, H. S., Wen, X., Paterson, R. G., Lamb, R. A. & Jardetzky, T. S. Structure of the parainfluenza virus 5 F protein in its metastable, prefusion conformation. *Nature* **439**, 38-44 (2006).
- (8) Harrison, S. C. Viral membrane fusion. *Nat Struct Mol Biol* **15**, 690-698 (2008).
- (9) Chang, A. & Dutch, R. E. Paramyxovirus fusion and entry: multiple paths to a common end. *Viruses* **4**, 613-636, doi:10.3390/v4040613 (2012).
- (10) Lamb, R. A., Paterson, R. G. & Jardetzky, T. S. Paramyxovirus membrane fusion: lessons from the F and HN atomic structures. *Virology* **344**, 30-37 (2006).
- (11) Plemper, R. K., Brindley, M. A. & Iorio, R. M. Structural and mechanistic studies of measles virus illuminate paramyxovirus entry. *PLoS pathogens* **7**, e1002058, doi:10.1371/journal.ppat.1002058 (2011).
- (12) White, J. M., Delos, S. E., Brecher, M. & Schornberg, K. Structures and mechanisms of viral membrane fusion proteins: multiple variations on a common theme. *Critical reviews in biochemistry and molecular biology* **43**, 189-219 (2008).
- (13) Sapir, A., Avinoam, O., Podbilewicz, B. & Chernomordik, L. V. Viral and developmental cell fusion mechanisms: conservation and divergence. *Dev Cell* **14**, 11-21, doi:S1534-5807(07)00485-6 [pii]10.1016/j.devcel.2007.12.008 (2008).
- (14) Smith, E. C., Popa, A., Chang, A., Masante, C. & Dutch, R. E. Viral entry mechanisms: the increasing diversity of paramyxovirus entry. *Febs J* **276**, 7217-7227, doi:10.1111/j.1742-4658.2009.07401.x (2009).
- (15) Lee, B. & Ataman, Z. A. Modes of paramyxovirus fusion: a Henipavirus perspective. *Trends Microbiol* **19**, 389-399, doi:10.1016/j.tim.2011.03.005 S0966-842X(11)00053-9 [pii] (2011).
- (16) Moscona, A. & Peluso, R. W. Fusion properties of cells persistently infected with human parainfluenza virus type 3: participation of hemagglutinin-neuraminidase in membrane fusion. *J Virol* **65**, 2773-2777 (1991).
- (17) Porotto, M., Murrell, M., Greengard, O. & Moscona, A. Triggering of human parainfluenza virus 3 fusion protein (F) by the hemagglutinin-neuraminidase (HN) protein: an HN mutation diminishes the rate of F activation and fusion. *J Virol* **77**, 3647-3654 (2003).
- (18) Outlaw, V. K. *et al.* Dual Inhibition of Human Parainfluenza Type 3 and Respiratory Syncytial Virus Infectivity with a Single Agent. *J Am Chem Soc*, doi:10.1021/jacs.9b04615 (2019).
- (19) Lambert, D. M. *et al.* Peptides from conserved regions of paramyxovirus fusion (F) proteins are potent inhibitors of viral fusion. *Proc Natl Acad Sci U S A* **93**, 2186-2191 (1996).
- (20) Porotto, M. *et al.* Inhibition of hendra virus fusion. *J Virol* **80**, 9837-9849, doi:10.1128/JVI.00736-06 (2006).
- (21) Porotto, M. *et al.* Molecular determinants of antiviral potency of paramyxovirus entry inhibitors. *J Virol* **81**, 10567-10574, doi:10.1128/JVI.01181-07 (2007).
- (22) Porotto, M. *et al.* Kinetic dependence of paramyxovirus entry inhibition. *J Virol* **83**, 6947-6951, doi:10.1128/JVI.00416-09 (2009).
- (23) Porotto, M. *et al.* Inhibition of Nipah virus infection in vivo: targeting an early stage of paramyxovirus fusion activation during viral entry. *PLoS Pathog* **6**, e1001168, doi:10.1371/journal.ppat.1001168 (2010).
- (24) Porotto, M. *et al.* Viral entry inhibitors targeted to the membrane site of action. *J Virol* **84**, 6760-6768, doi:10.1128/JVI.00135-10 (2010).
- (25) Mathieu, C. *et al.* Broad spectrum antiviral activity for paramyxoviruses is modulated by biophysical properties of fusion inhibitory peptides. *Sci Rep* **7**, 43610, doi:10.1038/srep43610 (2017).
- (26) Mathieu, C., Porotto, M., Figueira, T. N., Horvat, B. & Moscona, A. Fusion Inhibitory Lipopeptides Engineered for Prophylaxis of Nipah Virus in Primates. *J Infect Dis* **218**, 218-227, doi:10.1093/infdis/jiy152 (2018).
- (27) Outlaw, V. K. *et al.* Dual Inhibition of Human Parainfluenza Type 3 and Respiratory Syncytial Virus Infectivity with a Single Agent. *J Am Chem Soc* **141**, 12648-12656, doi:10.1021/jacs.9b04615 (2019).
- (28) Welsch, J. C. *et al.* Fatal measles virus infection prevented by brain-penetrant fusion inhibitors. *J Virol* **87**, 13785-13794, doi:10.1128/JVI.02436-13 (2013).
- (29) Mathieu, C. *et al.* Prevention of measles virus infection by intranasal delivery of fusion inhibitor peptides. *J Virol* **89**, 1143-1155, doi:10.1128/JVI.02417-14 (2015).
- (30) Figueira, T. N. *et al.* In Vivo Efficacy of Measles Virus Fusion Protein-Derived Peptides Is Modulated by the Properties of Self-Assembly and Membrane Residence. *J Virol* **91**, doi:10.1128/JVI.01554-16 (2017).
- (31) Figueira, T. N. *et al.* Quantitative analysis of molecular partition towards lipid membranes using surface plasmon resonance. *Sci Rep* **7**, 45647, doi:10.1038/srep45647 (2017).
- (32) Diao, L. & Meibohm, B. Pharmacokinetics and pharmacokinetic-pharmacodynamic correlations of therapeutic peptides. *Clin Pharmacokinet* **52**, 855-868, doi:10.1007/s40262-013-0079-0 (2013).
- (33) Horne, W. S. *et al.* Structural and biological mimicry of protein surface recognition by alpha/beta-peptide foldamers. *Proceedings of the National Academy of Sciences of the United States of America* **106**, 14751-14756, doi:10.1073/pnas.0902663106 (2009).
- (34) Johnson, L. M. *et al.* Enhancement of alpha-helix mimicry by an alpha/beta-peptide foldamer via incorporation of a dense ionic side-chain array. *Journal of the American Chemical Society* **134**, 7317-7320, doi:10.1021/ja302428d (2012).
- (35) Johnson, L. M., Horne, W. S. & Gellman, S. H. Broad distribution of energetically important contacts across an extended protein interface. *Journal of the American Chemical Society* **133**, 10038-10041, doi:10.1021/ja203358t (2011).
- (36) Horne, W. S., Price, J. L., Keck, J. L. & Gellman, S. H. Helix bundle quaternary structure from alpha/beta-peptide foldamers. *Journal of the American Chemical Society* **129**, 4178-4180, doi:10.1021/ja070396f (2007).
- (37) Horne, W. S., Price, J. L. & Gellman, S. H. Interplay among side chain sequence, backbone composition, and residue rigidification in polypeptide folding and assembly. *Proceedings of the National Academy of Sciences of the United States of America* **105**, 9151-9156, doi:10.1073/pnas.0801135105 (2008).
- (38) Price, J. L., Horne, W. S. & Gellman, S. H. Structural consequences of beta-amino acid preorganization in a self-assembling alpha/beta-peptide: fundamental studies of foldameric helix bundles. *Journal of the American Chemical Society* **132**, 12378-12387, doi:10.1021/ja103543s (2010).
- (39) Horne, W. S., Boersma, M. D., Windsor, M. A. & Gellman, S. H. Sequence-based design of alpha/beta-peptide foldamers that mimic BH3 domains. *Angewandte Chemie* **47**, 2853-2856, doi:10.1002/anie.200705315 (2008).
- (40) Lee, E. F. *et al.* Structural basis of Bcl-xL recognition by a BH3-mimetic alpha/beta-peptide generated by sequence-based design. *Chembiochem: a European journal of chemical biology* **12**, 2025-2032, doi:10.1002/cbic.201100314 (2011).
- (41) Boersma, M. D. *et al.* Evaluation of diverse alpha/beta-backbone patterns for functional alpha-helix mimicry: analogues of the Bim BH3 domain. *Journal of the American Chemical Society* **134**, 315-323, doi:10.1021/ja207148m (2012).
- (42) Yin, H. S., Paterson, R. G., Wen, X., Lamb, R. A. & Jardetzky, T. S. Structure of the uncleaved ectodomain of the paramyxovirus (hPIV3) fusion protein. *Proc Natl Acad Sci U S A* **102**, 9288-9293, doi:10.1073/pnas.0503989102 (2005).
- (43) Porotto, M., Fornabaio, M., Kellogg, G. E. & Moscona, A. A second receptor binding site on human parainfluenza virus type 3 hemagglutinin-neuraminidase contributes to activation of the fusion mechanism. *J Virol* **81**, 3216-3228, doi:10.1128/JVI.02617-06 (2007).
- (44) Palermo, L. M., Porotto, M., Greengard, O. & Moscona, A. Fusion promotion by a paramyxovirus hemagglutinin-neuraminidase protein: pH modulation of receptor avidity of binding sites I and II. *J Virol* **81**, 9152-9161 (2007).
- (45) Pessi, A. *et al.* A general strategy to endow natural fusion-protein-derived peptides with potent antiviral activity. *PLoS One* **7**, e36833, doi:10.1371/journal.pone.0036833 (2012).
- (46) Palmer, S. G. *et al.* Circulating clinical strains of human parainfluenza virus reveal viral entry requirements for in vivo infection. *J Virol* **88**, 13495-13502, doi:10.1128/JVI.01965-14 (2014).

- (47) Palermo, L. M. *et al.* Features of Circulating Parainfluenza Virus Required for Growth in Human Airway. *MBio* **7**, e00235, doi:10.1128/mBio.00235-16 (2016).
- (48) Iketani, S. *et al.* Viral Entry Properties Required for Fitness in Humans Are Lost through Rapid Genomic Change during Viral Isolation. *MBio* **9**, doi:10.1128/mBio.00898-18 (2018).
- (49) Palmer, S. G. *et al.* Adaptation of human parainfluenza virus to airway epithelium reveals fusion properties required for growth in host tissue. *MBio* **3**, doi:10.1128/mBio.00137-12 (2012).
- (50) Palermo, L. M. *et al.* Human parainfluenza virus infection of the airway epithelium: viral hemagglutinin-neuraminidase regulates fusion protein activation and modulates infectivity. *J Virol* **83**, 6900-6908, doi:10.1128/JVI.00475-09 (2009).
- (51) Xu, R. *et al.* Interaction between the hemagglutinin-neuraminidase and fusion glycoproteins of human parainfluenza virus type III regulates viral growth in vivo. *MBio* **4**, e00803-00813, doi:10.1128/mBio.00803-13 (2013).
- (52) Uusitupa, E., Waris, M. & Heikkinen, T. Association of Viral Load With Disease Severity in Outpatient Children With Respiratory Syncytial Virus Infection. *J Infect Dis* **222**, 298-304, doi:10.1093/infdis/jiaa076 (2020).
- (53) Outlaw, V. K. *et al.* Effects of single alpha-to-beta residue replacements on recognition of an extended segment in a viral fusion protein. *ACS Infect Dis*, doi:10.1021/acsinfecdis.0c00385 (2020).
- (54) Porotto, M. *et al.* Authentic Modeling of Human Respiratory Virus Infection in Human Pluripotent Stem Cell-Derived Lung Organoids. *MBio* **10**, doi:10.1128/mBio.00723-19 (2019).
- (55) de Vries, R. D. *et al.* Intranasal fusion inhibitory lipopeptide prevents direct contact SARS-CoV-2 transmission in ferrets. *bioRxiv*, doi:10.1101/2020.11.04.361154 (2020).
- (56) Ingallinella, P. *et al.* Addition of a cholesterol group to an HIV-1 peptide fusion inhibitor dramatically increases its antiviral potency. *Proc Natl Acad Sci U S A* **106**, S801-S806 (2009).
- (57) Moscona, A. *et al.* A Recombinant Sialidase Fusion Protein Effectively Inhibits Human Parainfluenza Viral Infection In Vitro and In Vivo. *J Infect Dis* (2010).



Insert Table of Contents artwork here