Rhinovirus Reduces the Severity of Subsequent Respiratory Viral Infections by Interferon-Dependent and -Independent Mechanisms

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ABSTRACT Coinfection by heterologous viruses in the respiratory tract is common and can alter disease severity compared to infection by individual virus strains. We previously found that inoculation of mice with rhinovirus (RV) 2 days before inoculation with a lethal dose of influenza A virus [A/Puerto Rico/8/34 (H1N1) (PR8)] provides complete protection against mortality. Here, we extended that finding to a second lethal respiratory virus, pneumonia virus of mice (PVM), and analyzed potential mechanisms of RV-induced protection. RV completely prevented mortality and weight loss associated with PVM infection. Major changes in host gene expression upon PVM infection were delayed compared to PR8. RV induced earlier recruitment of inflammatory cells, which were reduced at later times in RV-inoculated mice. Findings common to both virus pairs included the upregulated expression of mucin-associated genes and dampening of inflammation-related genes in mice that were inoculated with RV before lethal virus infection. However, type I interferon (IFN) signaling was required for RV-mediated protection against PR8 but not PVM. IFN signaling had minor effects on PR8 replication and contributed to controlling neutrophilic inflammation and hemorrhagic lung pathology in RV/PR8-infected mice. These findings, combined with differences in virus replication levels and disease severity, suggest that the suppression of inflammation in RV/PVM-infected mice may be due to an early, IFN-independent suppression of viral replication, while that in RV/PR8-infected mice may be due to IFN-dependent modulation of immune responses. Thus, a mild upper respiratory viral infection can reduce the severity of a subsequent severe viral infection in the lungs through virus-dependent mechanisms.

IMPORTANCE Respiratory viruses from diverse families cocirculate in human populations and are frequently detected within the same host. Although clinical studies suggest that infection by multiple different respiratory viruses may alter disease severity, animal models in which we can control the doses, timing, and strains of coinfecting viruses are critical to understanding how coinfection affects disease severity. Here, we compared gene expression and immune cell recruitment between two pairs of viruses (RV/PR8 and RV/PVM) inoculated sequentially in mice, both of which result in reduced severity compared to lethal infection by PR8 or PVM alone. Reduced disease severity was associated with suppression of inflammatory responses in the lungs. However, differences in disease kinetics and host and viral gene expression suggest that protection by coinfection with RV may be due to distinct molecular mechanisms. Indeed, we found that antiviral cytokine signaling was required for RV-mediated protection against lethal infection by PR8 but not PVM.
The detection of more than one virus in respiratory samples is quite common, especially among pediatric patients (1–4). There are differences in the outcomes of coinfection, whether it results in increased, decreased, or no effect on disease severity, that likely reflect different virus pairings, patient populations, and study criteria. For example, coinfection with influenza B virus was found to increase the severity of seasonal influenza A virus, while other virus pairings did not reach statistical significance (2). Another study found increased rates of hospitalization, but not other measures of clinical severity, associated with viral coinfections (1). In contrast, Martin et al. found that patients with one virus detected had an increased risk of severe disease compared to those with multiple viruses detected, and some virus pairings were associated with lower viral loads in coinfected patients (5). Coinfection by non-severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) viruses has been detected in coronavirus disease 2019 (COVID-19) patients, but the impact on disease severity is not well understood (6–8). Despite differences among studies, it is clear that coinfecting viruses have the potential to alter pathogenesis and disease outcomes. While clinical studies illuminate this potential, model systems in which the virus pairs, doses, and timing of coinfection can be controlled are critical to understanding how coinfection alters pathogenesis in the respiratory tract. Therefore, we developed a mouse model using pairwise combinations of respiratory viruses from different families for this purpose (9).

We previously found that inoculation of mice with a mild respiratory virus (rhinovirus strain 1B [RV1B] or mouse hepatitis virus strain 1 [MHV-1]) attenuates the severity of subsequent infection by influenza A virus [strain A/Puerto Rico/8/34 (H1N1) (PR8)] (9). Although coinfection does not prevent PR8 replication, it leads to faster viral clearance and resolution of pulmonary inflammation. Protection from severe disease by viral coinfection has also been demonstrated by other groups. Similar to our study, Hua et al. found that nasal-restricted infection by MHV-1 protects mice from lethal infection by PR8 and mouse-adapted severe acute respiratory syndrome coronavirus (SARS-CoV) (10). Furthermore, they found that protection is associated with enhanced macrophage recruitment to the lungs and upregulation of SARS-CoV-specific CD4+ and CD8+ T cell responses (10). Inhibition or delay of viral shedding also occurs in ferrets during sequential inoculations with antigenically similar or dissimilar strains of influenza A and B viruses (11). A 2009 pandemic influenza A [A(H1N1)pdm09] virus prevents subsequent infection of ferrets by human respiratory syncytial virus (RSV) when the viruses are given within a short time frame (12). In contrast, RSV does not prevent the replication of A(H1N1)pdm09 in ferrets but reduces morbidity as determined by weight loss (12). Inhibition of RSV by prior inoculation with influenza A virus has also been demonstrated in mice (13). Influenza A virus was recently shown to enhance infection of mice by SARS-CoV-2 through increased viral loads and severity of lung histopathology (14). Those authors further showed that enhancement of SARS-CoV-2 infection in cell lines by preinfection with influenza A virus was due to increased expression of the angiotensin converting enzyme 2 (ACE2) receptor, and infection was not enhanced by other respiratory viruses (14).

In this study, we aimed to evaluate whether RV-mediated disease attenuation was specific to PR8 or generalizable to other respiratory viral infections. We found that RV reduced the severity of an additional lethal respiratory virus, pneumonia virus of mice (PVM). PVM is a relative of human RSV in the family Pneumoviridae, genus Orthopneumovirus. Infection of mice with PVM results in severe disease that shares clinical features of the most severe infections by RSV, including infection of the bronchiolar epithelium, granulocytic inflammation, and pulmonary edema (15–17). Lethality upon infection of mice with PR8 or PVM is mediated largely by dysregulated inflammatory responses rather than overt damage due to viral replication (18–21). Interestingly, there are differences in the types of immune responses that mediate protection against these two viruses. Type I and III interferon (IFN)
signaling and alveolar macrophages are protective in PR8- but not PVM-infected mice (18, 20, 22–25). Conversely, plasmacytoid dendritic cells are required for protection in PVM- but not PR8-infected mice (26, 27). Interleukin-6 (IL-6) limits the severity of influenza A virus infection in mice, while it exacerbates disease in PVM-infected mice (28–30). Multiple mechanisms reduce the severity of both PR8 and PVM infections, including mucin production and inhibition of inflammatory responses (31–37). Based on these complex differences in immunity to infection with PR8 and PVM, we used a global gene expression approach combined with flow cytometry to evaluate potential mechanisms whereby preinoculation with RV reduces the severity of PR8 and PVM.

RESULTS

Inoculation of mice with RV reduces the severity of PVM infection. We previously showed that inoculation of mice with RV completely prevented mortality of a lethal infection by influenza A virus strain PR8 (9). Protection against lethal PR8 infection was most effective when RV was given 2 days before PR8, but significant disease attenuation was also seen when mice were inoculated with RV and PR8 concurrently. To determine if RV would attenuate disease by a second lethal respiratory viral pathogen, we inoculated mice with medium (mock) or RV 2 days before, simultaneously with, or 2 days after inoculation with PVM. Mice were monitored daily for mortality, weight loss, and clinical signs of disease for 14 days after PVM inoculation.

Infection with PVM alone (mock/PVM) was 100% lethal, with all mice succumbing to infection or reaching humane endpoints by day 8 (Fig. 1A). PVM-infected mice had rapid weight loss (Fig. 1B) and exhibited clinical signs of disease, including ruffled fur, hunched posture, labored breathing, and lethargy. Mice that were inoculated with RV 2 days before (RV/PVM) or simultaneously with (RV+PVM) PVM were completely protected from mortality (Fig. 1A), weight loss (Fig. 1B), and clinical signs of disease. However, mice that received RV 2 days after PVM (PVM/RV) had disease severity equivalent to that of mock/PVM-infected mice. Although inoculation with RV reduced the severity of both PR8 (9) and PVM (Fig. 1), protection was more effective against PVM. Importantly, complete protection against morbidity in addition to mortality was seen when RV was given 2 days before or concurrently with PVM (Fig. 1). In contrast, RV prevented mortality, but not morbidity, when given 2 days before PR8, and inoculation of RV and PR8 concurrently was less effective at reducing disease severity (9).

Infections by PR8 and PVM induce different gene expression signatures in mouse lungs over time. To determine potential mechanisms of protection mediated by RV against PR8 and PVM, we undertook a comprehensive transcriptome analysis of mouse lungs (Fig. 2). Mice were inoculated with RV 2 days before PR8 or PVM, and RNA isolated from the right lobes was analyzed on days 0, 2, 4, and 6 after PR8 or PVM inoculation. Single-virus-infected mice were mock inoculated 2 days before PR8 or PVM. Weight loss was monitored daily to test for consistency with our previous studies. RV-mediated protection against PR8 was not evident by 6 days postinfection (see Fig. S1 in the supplemental material), although both the mock/PR8 and RV/PR8 groups experienced weight loss at a rate similar to that in our previous study (9). In contrast, complete protection against weight loss was evident in RV/PVM-inoculated mice (Fig. S1).

Lung mRNA from 3 mice per infection group and time point was processed and sequenced on the illumina platform (RNA-seq). Of the 14.6 million to 55.3 million (mean, 36.4 million) reads obtained for each sample, 83 to 91.2% were uniquely mapped to the mm10 mouse genome (RefSeq accession number GCA_000001635.2). The expression levels of 24,243 and 24,421 genes were measured using Cufflinks and HTSeq, respectively. We conservatively included differentially expressed genes (DEGs) by requiring that they be identified by Cuffdiff, EdgeR, and DESeq2 as having a log2 fold change value of at least 0.5 and a P value of less than 0.05.

A principal-component analysis (PCA) was done with the 500 most highly variable genes to visualize variation across infection groups, time points, and replicates. For the most part, replicate samples clustered together (Fig. 3A). Most variation in host gene expression was explained by the time elapsed since viral infection (Fig. 3A). However,
differences between viruses were also apparent. Infection by PVM caused fewer gene expression changes at early time points than in PR8-infected mice, as seen by mock/PVM and RV/PVM day 2 samples clustering closely with mock and RV day 0 samples, respectively (Fig. 3A, blue and purple points). By day 4, the PVM-infected mice had very different gene expression profiles than those of mock-inoculated mice on day 0. This delayed response to PVM infection has been observed in other studies (18, 38). In contrast, PR8 infection dramatically altered gene expression in mice by day 2 (Fig. 3A, light green points). Early changes in host gene expression induced by PR8, but not PVM, corresponded to the earlier detection of PR8-specific transcripts (Fig. 4). Moreover, the RV/PVM and mock/PVM samples were very different from each other by day 4 (Fig. 3A, brown points). The RV/PR8 and mock/PR8 groups had similar gene expression signatures until day 6 (Fig. 3A, orange points). This corresponds to the similar disease kinetics in the mock/PR8 and RV/PR8 groups early during infection (9) (Fig. S1).

We next calculated differential gene expression levels in samples from RV-inoculated mice compared to mock-inoculated mice on day 0 (Fig. 3B). Similar to the PCA, gene expression was significantly changed 2 days after inoculation with RV, and the response was markedly skewed toward upregulation. DEGs identified in all infection groups versus mock were compared between single-virus-infected and RV-treated mice at each time point (Fig. 3C). Gene expression changed over time, as samples taken at 2 days postinfection were most similar to those of mock-inoculated controls, and samples taken at 6 days postinfection were most different (Fig. 3C and D). The
transition of point clustering in Fig. 3C from around the origin on day 2 to spread along the 1:1 line on day 6 illustrates this delayed gene expression response to viral infection. Genes that were differentially expressed between single-virus-infected and RV-treated mice also displayed time-dependent clustering shifts in both the PVM and PR8 experiments (Fig. 3C). At 2 days postinfection, the expression values of DEGs indicated that the samples from the RV/PR8 and RV/PVM groups were more different from mock than the samples from single-virus-infected mice (points are spread along the y axis). At 6 days postinfection, DEGs differed more between mock-inoculated and single-virus-infected mice than between mock- and RV-treated mice (points are spread along the x axis). This dampening of DEGs in the RV/PR8 and RV/PVM groups suggests that infection by a lethal virus is ablated not only in virulence but also in the host gene expression response.

A summary of the numbers of DEGs across comparisons of all virus-infected to mock-inoculated mice and between single-virus-infected and RV/PR8- or RV/PVM-treated mice is shown in Fig. 3D, and lists of these DEGs are included in Table S1. Again, the numbers of DEGs reflect the delayed response to PVM infection, dampened gene expression in RV/PR8- and RV/PVM-infected mice, and relative similarity in gene expression in mock/PR8- and RV/PR8-infected mice early in infection.

Viral gene expression. Sequencing reads that did not map to the mouse genome were aligned with viral genomes. An insignificant number of reads mapped to the RV genome. This was expected because the abundance of positive-sense RV RNA and viral titers peak 24 h after inoculation of BALB/c mice (39). The earliest time that we analyzed was 48 h (day 0) after inoculation, at which point RV RNA would likely represent a very low proportion of the total RNA. We confirmed this by quantifying infectious RV in homogenized lung tissue and bronchoalveolar lavage fluid (BALF) 24 h and 48 h after RV inoculation, days −1 and 0, respectively. In agreement with published studies and our RNA-seq results, low levels of RV were detected on day −1 (10^2 to 10^3 50%...
FIG 3 Patterns of mouse gene expression in lungs upon infection with and without RV pretreatment. (A) Principal-component analysis of RNA-seq data showing that the greatest variance in gene expression is primarily due to time since infection. The largest distances in single-virus-infected versus RV-treated groups are mock/PVM versus RV/PVM on day 4 and mock/PR8 versus RV/PR8 on day 6. (B) Volcano plot showing gene expression changes between mock-inoculated and RV-inoculated mice at 2 days (Continued on next page)
tissue culture infective doses [TCID_{50}] per ml of BALF or g of lung tissue) but not day 0. Thus, we cannot conclude that RV is replicating in mouse lungs despite having a significant impact on host gene expression.

Many reads mapped to the PR8 and PVM genomes from the mice infected with these viruses (Fig. 4). Regression analyses were performed to identify significant changes over time (Table S3). Pretreatment with RV did not prevent PR8-specific gene expression, but the numbers of reads that mapped to PR8 were lower in RV/PR8-infected mice at all time points (Fig. 4A). When this analysis was expanded to additional animals assayed by reverse transcription-quantitative PCR (RT-qPCR) for the hemagglutinin (HA) gene, RV/PR8-infected mice had significantly lower viral gene expression levels than mock/PR8-infected mice only on day 6 (Fig. 4B), which was also the time point that was the most dramatically different in RNA-seq reads. Similarly, we previously showed that infectious PR8 titers in the lungs were equivalent in mock/PR8- and RV/PR8-infected mice on days 2 and 4 after PR8 inoculation (9). However, inoculation with RV led to the earlier clearance of PR8 (by day 7), corresponding to the dramatically lower PR8 mRNA levels seen on day 6 in the present study. These findings confirm that infection with RV does not prevent subsequent infection by PR8; rather, it reduces viral gene expression, specifically late during infection. Our sequencing protocol captured only polyadenylated RNAs; thus, the PR8 reads represent viral mRNA and not complementary (cRNA) or genomic (vRNA) viral RNAs, while RT-qPCR was performed on total RNA. Based on individual gene mapping, the PR8 reads predominantly mapped to the nucleoprotein (NP) and HA mRNAs (Fig. S1), which are known to be expressed at high levels during infection (40–42).

PVM gene expression was highest on day 4 but was robustly suppressed in RV-inoculated mice (Fig. 4). This trend was confirmed by RT-qPCR quantification of the PVM small hydrophobic protein (SH) gene (Fig. 4B) and quantification of infectious virus (see Fig. 7E, below). This suggests that RV may limit PVM pneumonia early, which corresponds to the more effective prevention of weight loss in RV/PVM (Fig. 1) than in RV/PR8 (9)-infected mice (Fig. S1). All viral genes were detected in mice infected by PVM alone (mock/PVM) on day 4, with the genes that express the attachment (G), nucleoprotein (N), nonstructural 2 (NS2), fusion (F), phosphoprotein (P), and matrix (M) proteins being present at the highest levels (Fig. S1). This does not strictly follow the gradient of mRNA levels corresponding to gene order that is expected from pneumoviruses (43), which likely reflects posttranscriptional differences in mRNA stability and the heterogeneous nature of collecting cells at different stages of the virus replication cycle.

**RV induces innate immune responses prior to secondary viral infection.** We compared gene expression levels in lung tissues from mock- versus RV-inoculated mice 2 days after inoculation (RV, day 0). Of the 24,421 genes compared, we identified 330 DEGs, of which only 3 were downregulated (Fig. 3B and D and Table S1). To get a functional picture of the RV-treated lung on day 0, we identified enriched gene ontology (GO) terms and KEGG pathways in the 330 DEGs. The most highly enriched terms on this list suggest that changes in the regulation of the cell cycle or cell division were occurring in RV- compared to mock-inoculated mice; nearly all of the top 50 most enriched terms involved chromosome remodeling and mitosis (Table S2), suggesting an RV-induced increase in cellular proliferation. This list of enriched terms was quite different from the processes that were differentially regulated in PR8- or PVM-infected mice. Upregulation of cell division-associated genes could be occurring in epithelial cells in conjunction with repair, immune cells recruited to the lungs, or both. In addition to cell division, multiple immune response-related GO terms related to type I IFN postinfection. Only DEGs identified by EdgeR, DESeq2, and Cufflinks were considered significant (red points). (C) Scatterplots showing the log_{2}, fold change values of all genes compared to mock-inoculated mice on day 0. Genes upregulated in RV-treated compared to single-virus-infected mice are in green. Downregulated genes are in purple. (D) Numbers of DEGs in all pairwise comparisons. The numbers of genes upregulated in the infected (versus mock-infected) or RV-treated (versus mock- or single-virus-infected) mice are shown in green. Downregulated genes are shown in purple. See Table S1 in the supplemental material for lists of gene names and log, fold change values.
signaling, chemokine signaling, and immune cell chemotaxis were enriched in RV-treated mice (Table S2). RV induced the upregulation of several chemokine genes, including those that recruit monocytes, neutrophils, NK cells, T cells, B cells, and eosinophils (Ccl-2, -3, -6, -7, -8, -12, -17, and -22 and Cxcl-1,-3,-5,-9,-10, and -13).

**Host gene expression changes in RV/PR8- and RV/PVM-infected mice.** We next identified DEGs specific to mice exposed to two viruses (RV/PR8 or RV/PVM) versus mock-inoculated mice that were not differentially expressed in single-virus-infected (mock/PR8 or mock/PVM) versus mock-inoculated mice at the same time points. The expression data for these three gene sets (unique to RV/PR8, unique to RV/PVM, and...
shared by RV/PR8 and RV/PVM) are provided in Fig. S2 to S4. Genes with shared upregulation in RV-inoculated animals, regardless of the time point and second virus, included those in the mucin biosynthesis pathway, major histocompatibility complex (MHC) class II genes, and immunomodulatory genes. Genes that had higher upregulation in the single-virus infections were predominantly increased late (days 4 and/or 6) and included genes associated with inflammation (Angpt4) or pulmonary fibrosis (Fosl2, Pappa, and Sphk1). These genes were common to both PR8 and PVM infections and likely result from excessive inflammation and tissue damage within the infected lungs. Additional genes associated with inflammation and fibrosis (Nkbia and Mmp9) or stress responses (Hspb8 and Nupr1) had lower expression levels in RV/PVM- than in mock/PVM-infected mice. Finally, a set of genes was upregulated in both mock/PVM- and RV/PVM-infected mice but to a higher level in mock/PVM-infected mice (S100a9, Prss22, Fga, and Krt17). These genes are also largely involved in inflammation and tissue damage and repair processes.

Genes involved in goblet cell metaplasia and mucin production were specifically increased in both RV/PVM- and RV/PR8-infected mice compared to the single-virus-infected mice at various time points. These included the major gel-forming airway mucins (Muc5ac and Muc5b), a disulfide isomerase (Agtr2) required for mucin folding and polymerization, and a chloride channel regulator (Clica1) required for the proper hydration of mucus. Additional ion channels (Scl6a20a), aquaporins (Aqp9), and mucus-associated proteins (Itln1) also had increased expression in the lungs of RV-treated mice. In addition to the mucin-related genes shared with RV/PR8-infected mice, RV/PVM-infected mice had increased expression of a transcription factor (FoxA3) that promotes goblet cell metaplasia and mucus production.

Based on the importance of type I IFN and inflammatory signaling pathways in the pathogenesis of viral infections, we generated heat maps demonstrating relative gene expression levels across all groups for genes in the hallmark interferon alpha response and inflammatory response gene sets from MSigDB (44, 45). Overall, IFN response genes had higher expression levels in RV/PR8- than in mock/PR8-infected mice on day 2 and lower expression levels by day 6 (Fig. 5A and Fig. S5). This pattern corresponds to our previous study that demonstrated increased expression of IFN-β only on day 2 in RV/PR8- compared to mock/PR8-infected mice (9) and also the lower PR8 gene expression levels in RV/PR8-infected mice (Fig. 4). In contrast, mock/PVM- and RV/PVM-infected mice had delayed upregulation of IFN response genes, and mock/PVM-infected mice had dramatically higher expression levels of IFN response genes than RV/PVM-infected mice on day 4 (Fig. 5B and Fig. S5). This pattern was validated by RT-qPCR analysis of IFN-β expression (see Fig. 7D) and corresponds to the overall delayed PVM-induced gene expression and reduced levels of PVM RNA in RV/PVM-infected mice (Fig. 4). Inoculation with RV induced the expression of a small subset of IFN response genes (Il17, Ifi27, Lamp3, Cd74, Ifi30, and Lpar6) on day 0, which was maintained in RV/PVM-infected mice on day 2 (Fig. 5B). While more variation in gene expression patterns was observed in the inflammatory response gene set, many genes followed the same trends as those seen for IFN response genes, i.e., a largely dampened response in RV-treated mice, especially at later time points (Fig. 5C and D and Fig. S6). By day 6, mock/PR8-infected mice had strongly up- and downregulated expression of inflammatory response genes, while RV/PR8-infected mice had muted changes in the expression of these genes (Fig. 5C). A large subset of inflammatory response genes had patterns similar to those of IFN response genes in mock/PVM- and RV/PVM-infected mice, including delayed expression and a muted response in RV-treated mice (Fig. 5D). This subset includes important inducers of inflammatory responses, such as Nkbia, Rela, and Tlr3. Other subsets of genes (e.g., Ccl17 and Ccl22) had early upregulation in RV-inoculated mice on day 0, which was maintained on day 2 after infection with PVM, and reduced in RV/PVM-infected mice on days 4 and 6. In contrast to IFN response genes, subsets of inflammatory response genes had higher expression levels in RV/PVM- than in mock/PVM-infected mice throughout the course of infection. This suggests that despite low virus levels (Fig. 4), no clinical signs
Expression of MSigDB hallmark interferon alpha response and inflammatory response gene sets. (A and B) Heat maps of hallmark interferon alpha response genes showing the relative expression (Z-scores) of genes for the PR8 (A) or PVM (B) infection RNA-seq time course. (C and D) Heat maps of hallmark inflammatory response genes showing the relative expression (Z-scores) of genes for the PR8 (C) or PVM (D) infection RNA-seq time course. All heat maps represent DESeq2-normalized counts, where each row represents an individual gene. The colors (blue < white < red) represent the Z-scores, and a more intense color indicates a lower (blue) or higher (red) relative expression level of that gene under that condition. These heat maps with the gene names included can be found in Fig. S5 and S6 in the supplemental material.
of disease (Fig. 1 and Fig. S1), and largely muted host gene expression, the expression of some host genes is increased in RV/PVM-infected mice.

**Flow cytometry analysis.** Immune cells recruited to the lungs are affected by and contribute to the gene expression signatures seen in whole lung tissues. To analyze differences in immune cell recruitment to the lungs of PR8- and PVM-infected mice with and without RV pretreatment, we quantified innate immune cells in the left lobes by flow cytometry at the same time points as those for our gene expression analyses (Fig. 6) and performed regression analyses to identify significant changes over time (Table S3). Total cell counts from mock/PVM- and RV/PVM-infected mice were fairly consistent across all time points, while total cells in mock/PR8- and RV/PR8-infected mice increased by day 6 (Fig. S7). CD11b was used to differentiate leukocytes from resident lung cell populations. CD11b+ cells accounted for the increase in mock/PR8- and RV/PR8-infected lungs (Fig. 6A). In contrast, while CD11b+ cells increased in the lungs of mice infected with mock/PVM, RV/PVM-infected mice had reduced recruitment of these cells on days 4 and 6 (Fig. 6E). Neutrophils were gated based on high expression levels of CD11b and Ly6G, and the remaining cells were identified as alveolar macrophages (CD64+/SiglecF−) and interstitial macrophages (CD11b+/CD64+/SiglecF−) (Fig. S7) (46, 47).

The proportions of neutrophils and interstitial macrophages followed the same trends as those of total CD11b+ cells in PVM-infected mice, with lower proportions of these cells on days 4 and 6 in RV/PVM-infected mice (Fig. 6F and G). In contrast, interstitial macrophages were increased in RV/PVM- compared to mock/PVM-infected mice early in infection. This indicates that pretreatment with RV stimulates the early recruitment of CD11b+ cells, specifically interstitial macrophages, while limiting the recruitment of inflammatory cells later in infection. PR8-infected mice had similar trends; however, the differences between the mock/PR8 and RV/PR8 groups were less dramatic (Fig. 6B and C). Neutrophil numbers were suppressed in RV/PR8-infected mice compared to mock/PR8-infected mice throughout the time course (Fig. 6B). The interstitial macrophage proportions in mock/PR8- and RV/PR8-infected mice increased over time similarly to the total CD11b+ populations (Fig. 6C). The lower proportions of neutrophils and interstitial macrophages at later time points in RV-treated mice corresponded to mRNA levels for chemokines. This was predominantly the case for the neutrophil chemokines Cxcl1 and Cxcl2 and the macrophage chemokines Ccl2 and Ccl7. Chemokine mRNA levels were generally lower in RV/PR8-infected mice on day 6 and RV/ PVM-infected mice on days 4 and 6 than in mock/PR8- and mock/PVM-infected mice, respectively (Fig. S6; Table S1).

There were no clear trends in alveolar macrophage numbers in mock/PR8- and RV/ PR8-infected mice (Fig. 6D), although their proportions were significantly higher in RV/ PVM-infected mice than in mock/PVM-infected mice on days 4 and 6 (Fig. 6H). This is likely due to the depletion of alveolar macrophages by PVM infection of these cells (48). A separate flow cytometry antibody (Ab) panel was used to quantify CD4+ and CD8+ T cells in the lungs. T cell numbers in mock/PR8- and RV/PR8-infected mice increased over time, but the differences between the groups were not significant (Fig. 6I and J). RV/PVM-infected mice had modest yet significantly high numbers of CD4+ and CD8+ T cells compared to mock/PVM-infected mice on day 6 (Fig. 6K and L).

**IFNAR signaling is required for RV-mediated protection against PR8 but not PVM.** To determine whether signaling by type I IFNs is required for RV-mediated reduction in disease severity, we treated mice with an IFN-α/β receptor 1 (IFNAR-1)-blocking Ab on day −2 (with RV inoculation) and day 0 (with PR8 or PVM inoculation). Mock/ PR8-infected mice had similar weight loss and mortality with anti-IFNAR and control antibody (Ctl Ab) treatments (Fig. 7A). In contrast, treatment with anti-IFNAR Ab completely abrogated the dampening effects of RV on the morbidity and mortality from PR8 (Fig. 7A). Thus, IFNAR signaling is required for the reduced disease severity observed in RV/PR8-infected mice. Mice that were inoculated with RV alone did not experience weight loss, clinical signs of infection, or mortality over 14 days when treated with either anti-IFNAR Ab or Ctl Ab (data not shown). Interestingly, there were no significant differences in weight loss and mortality between anti-IFNAR- and Ctl Ab-treated RV/PVM-infected mice (Fig. 7B). Thus, type I IFN signaling is not required for RV-mediated protection against lethal PVM infection.
FIG 6 Flow cytometry analysis of innate immune cell populations in the lungs upon viral infection and RV treatment. Mice were inoculated with mock or RV on day –2 and PR8 (A to D, I, and J) or PVM (E to G) on day 0. 

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To verify that IFNAR signaling was effectively inhibited in anti-IFNAR Ab-treated mice, we analyzed the expression of an IFN-induced gene, \textit{Mx1}, in mock/PVM- and RV/PVM-infected mice treated with anti-IFNAR Ab or Ctl Ab. Significant inhibition of \textit{Mx1} expression was seen through the course of anti-IFNAR Ab treatment (days -1 through 2), with recovery of IFNAR signaling on day 4 after PVM infection (Fig. 7C). In agreement with the weight loss and mortality data, inhibition of IFNAR signaling also had no significant effect on the viral load of PVM in mock/PVM- or RV/PVM-infected mice (Fig. 7E). \textit{Mx1} expression was significantly inhibited in anti-IFNAR Ab-treated mice infected with mock/PR8 and RV/PR8 on day 4 (Fig. 7F), confirming effective inhibition of type I IFN signaling.

**IFN-dependent protection in RV/PR8-infected mice is associated with reduced viral spread, neutrophilic inflammation, and histopathology in the lungs.** Type I IFN signaling stimulates multiple downstream responses, including the expression of antiviral genes, activation of innate and adaptive responses, and modulation of inflammatory responses. To better understand the role of IFNAR signaling in limiting disease severity in RV/PR8-infected mice, we analyzed PR8 replication, immune cell recruitment, and histopathology in RV/PR8-infected mice treated with anti-IFNAR Ab or Ctl Ab. Anti-IFNAR Ab-treated mice had higher levels of PR8 in the lungs on days 2 and 6 than Ctl Ab-treated mice, although the differences were not statistically significant (Fig. 8A). Similarly, PR8 antigen was more widespread in the lungs of anti-IFNAR Ab-treated mice on days 2 and 6 postinfection (Fig. 8B). However, IFNAR signaling did not completely prevent PR8 replication in RV/PR8-infected mice. This is in agreement with the similar levels of PR8 RNA (Fig. 4), viral loads, and antigen (9) in RV/PR8- compared to mock/PR8-infected mice.

Mice treated with anti-IFNAR Ab had a slightly higher percentage of neutrophils in the airways than mice treated with a Ctl Ab (Fig. 8C); however, the difference was not statistically significant. Overall, levels of inflammation were similar in the lungs of RV/PR8-infected mice treated with anti-IFNAR Ab and Ctl Ab (Fig. 8D). However, immunohistochemistry (IHC) staining for a neutrophil-specific protein, Ly6G, indicated that IFNAR signaling limits neutrophilic inflammation in RV/PR8-infected lungs (Fig. 8E). Enhanced neutrophilic inflammation in the lung parenchyma of anti-IFNAR Ab-treated mice was accompanied by increased infiltration of red blood cells into the alveoli (pulmonary hemorrhage) on day 6 (Fig. 8D). Altogether, these results suggest that type I IFN signaling plays a role in reducing, but not eliminating, viral replication and curtailing both neutrophil spread within the lungs and pulmonary hemorrhage in RV/PR8-infected mice.

To evaluate RV-induced recruitment of cells in the airways of mice, we quantified macrophages, neutrophils, and lymphocytes in BALF from mice that received mock or RV with or without anti-IFNAR Ab treatment on day 2 after inoculation (day 0 of the study). Inoculation with RV did not increase the overall cell counts in BALF but dramatically changed the composition from predominantly macrophages to neutrophils (Fig. 9). Inhibition of IFNAR signaling did not alter cellular recruitment in response to RV infection.

As described above, both RV/PR8- and RV/PVM-infected mice had increased expression levels of genes in the mucin biosynthesis pathway compared to mock/PR8- and mock/PVM-infected mice. \textit{Muc5ac} and \textit{Clca1} had increased expression in RV-treated mice at all time points (Fig. 10A). As Muc5ac is an important gel-forming airway mucin involved in the protection against respiratory viral infections, we confirmed its upregulation by RV treatment by Western blotting (Fig. 10B). The production of Muc5ac in the lungs was only slightly reduced by anti-IFNAR Ab treatment in RV-inoculated mice (Fig. 10B).
FIG 7  IFNAR signaling is required for RV-mediated protection in RV/PR8- but not RV/PVM-infected mice. Mice were treated with anti-IFNAR or an isotype control (Ctl) antibody (Ab) intranasally with viral inocula (mock or RV on day 2 and PR8 [A and F] or PVM [B to E] on day 0). (A and B) Animal weights and survival were monitored in five mice per group for 14 days after inoculation with PR8 (A) or PVM (B). Significant differences in survival between IFNAR Ab- and Ctl Ab-treated mice were determined using survival curve analysis by a log rank Mantel-Cox test. *, P < 0.05; ns, not significant (P > 0.05). (C) Expression of the IFN-induced gene Mx1 was monitored to evaluate the effectiveness of IFNAR Ab treatment in mock/PVM- and RV/PVM-infected mice. Data shown are means and standard errors for 4 to 5 mice per group and are representative of results from two replicate assays. (D) IFN-β RNA was quantified by RT-qPCR in 5 animals per group in mock/PVM- and RV/PVM-infected mice without antibody treatment. Data shown are means with standard errors and are representative of results from two assays. (E) PVM titers in BALF were quantified by TCID₅₀ assays. Data shown are from individual animals, with the geometric means and standard deviations indicated. (F) Expression of Mx1 was analyzed by RT-qPCR to evaluate the effectiveness of IFNAR Ab treatment in mock/PR8- and RV/PR8-infected mice. Data shown are means and standard errors for 4 to 5 mice per group and are representative of results from two replicate assays. Statistical significance between groups within each time point was determined by unpaired t tests corrected for multiple comparisons by the Holm-Sidak method. **, P < 0.01; ***, P < 0.001.
DISCUSSION

Previously, we found that inoculation of mice with a mild respiratory viral pathogen, RV or the murine coronavirus MHV-1, 2 days before PR8 provided significant protection against PR8-mediated disease (9). In this study, we expanded these results to show that RV-mediated protection was not specific to PR8 but also provided significant disease protection against a respiratory virus from another viral family, PVM. This is in agreement with other studies showing heterologous immunity by respiratory viral pathogens in mouse models (10–13). Furthermore, RV has been shown to inhibit the replication of influenza A virus and SARS-CoV-2 in cultured airway epithelial cells (49, 50). Interference by respiratory viruses within a host may contribute to reduced disease severity and alter population-level viral dynamics. Coinfection by rhinoviruses has been associated with reduced severity of the 2009 pandemic influenza A virus strain despite similar H1N1 viral loads (51). Conversely, infection with influenza viruses has also been found to reduce the severity of rhinovirus infections (52). Epidemiological studies have suggested that cocirculating viruses, especially rhinoviruses, can inhibit population-level dynamics of other respiratory viruses (49, 51, 53–55). In 2009, the introduction of the new pandemic H1N1 strain delayed the circulation of other seasonal respiratory viruses (56). A large study of children with community-acquired pneumonia found the incidence of rhinoviruses to be high; however, rhinoviruses were found less frequently in combination with other RNA viruses, including influenza virus and RSV, than would be expected if coinfection were random (3). Similar findings have been reported in adult populations (49). Another large, multicenter study of infants observed significant interference between RSV and RV (57). A prospective study of children found that infection with RV did not affect the likelihood of having an influenza virus infection the following week, but influenza virus infection decreased the chances of having RV the following week (58). In a study of RSV and influenza cases across seven seasons in Israel, Drori et al. found that when the peak of RSV cases coincided with the influenza peak, the percentages of RSV-positive cases were lower than when the RSV cases peaked prior to influenza (13). While these studies suggest an important role of respiratory viral coinfection in viral pathogenesis and epidemiology, animal models are necessary to understand the interactions between coinfecting viruses and their host that may contribute to these observations.

Despite the commonality of RV reducing the disease severity of two heterologous lethal viruses in our mouse model, there are differences between the virus combinations in disease kinetics, viral replication, and mechanisms of protection. Inoculation of mice with RV provided more effective protection against PVM than PR8. RV/PVM-infected mice had few to no signs of disease and significantly limited PVM gene expression. In contrast, coinfection by RV prevented mortality, but not morbidity, associated with PR8 infection and reduced viral gene expression but did not prevent infection by PR8 (9). Furthermore, RV given concurrently with PVM was as effective as when it was given 2 days before PVM. In contrast, we previously showed that RV was less effective at reducing the severity of PR8 when given concurrently and even exacerbated disease when it was given 2 days after PR8 (9). We also observed differences in the kinetics of gene expression in response to these virus pairs. Host and viral gene expression changes in response to PVM were delayed compared to PR8, thereby giving a larger window for RV-mediated protection. Thus, RV may be inducing antiviral mechanisms that are more effective against PVM, or different mechanisms may be responsible for inhibiting PVM infection and mediating effective clearance of PR8. In support of the latter, we found that IFNAR signaling was required for RV-mediated reduction of disease severity during PR8, but not PVM, infection.

While we cannot conclude that RV replicated in the airways or lungs, intranasal inoculation of mice with a high dose of RV resulted in the dramatic upregulation of host gene expression prior to inoculation with the second, lethal virus. Despite the expression of several chemokine and chemokine receptor genes, we did not observe a dramatic increase in immune cells in the lungs of RV-inoculated mice on day 0. Our
FIG 8 IFNAR-dependent protection in RV/PR8-infected mice is associated with reduced viral spread, neutrophilic inflammation, and histopathology in the lungs. Mice were inoculated with RV 2 days prior to PR8 (on day 0) and control (Ctl) or IFNAR antibodies were given with viral inoculations. (A) PR8 was quantified in homogenized lungs by TCID_{50} assays. Data shown are from five individual mice per group, with geometric means and standard deviations (Continued on next page)
flow cytometry panel was limited to focus on inflammatory cells (neutrophils and macrophages) and T cells and likely missed other cell types that could be recruited by RV early in infection. Furthermore, by using whole lungs for our assays, we may miss populations of cells that are small, but significant, in the airways. Indeed, other studies have found that neutrophils and lymphocytes are increased in the airways of RV-infected mice 2 days after inoculation (39, 59), which we also detected by analyzing BALF. While the chemokine signaling genes that we identified in RV-inoculated mice on day 0 were also increased in single-virus-infected mice later, the early upregulation of immune cell recruitment in RV-treated mice may contribute to the early control of infection and reduced disease severity.

In contrast to the early recruitment of neutrophils, RV-treated mice had reduced numbers of neutrophils and interstitial macrophages later during infection, which could be a result of reduced viral infection and/or a direct downregulation of the inflammatory response. Other studies have shown that rhinoviruses inhibit macrophage responses to bacterial infection and Toll-like receptor (TLR) stimulation (60, 61). This downregulation leads to reduced neutrophil recruitment and activation and results in enhanced bacterial infection. While suppression of TLR signaling is detrimental during a subsequent bacterial infection, it can be protective in the case of respiratory viral infections for which inflammatory responses contribute to disease pathology. Blocking signaling from multiple TLRs (TLR2, TLR4, TLR7, and TLR9) through inhibiting the adaptor protein TIRAP reduces the severity of lethal PR8 infection in mice (62). Inhibition of TIRAP reduces PR8-induced cytokine production by macrophages and is likely protective by limiting inflammatory responses (62). Similarly, inhibition of TLR2 and TLR4 signaling during PR8 infection reduces inflammatory cytokine responses and disease severity (63).

Inflammatory responses, particularly by granulocytes, are also associated with disease pathogenesis of PVM infection, independent of viral replication (34, 35). Priming with probiotic

FIG 9

Inoculation of mice with RV results in IFNAR-independent recruitment of neutrophils to the airways. Cytospins from BALF were stained with Hema 3 to identify airway macrophages, neutrophils, and lymphocytes. At least 300 total cells were counted from duplicate cytospins for each sample to determine the percentage of each cell type. Total cells were counted in the BALF samples prior to cytospin preparation. Data shown are means and standard errors from 5 mice per group. Statistical significance between groups was determined using unpaired t tests corrected for multiple comparisons using the Holm-Sidak method. ***, P < 0.001.

FIG 8 (Continued)

indicated. Statistical significance between Ctl Ab- and IFNAR Ab-treated mice was determined using unpaired t tests corrected for multiple comparisons using the Holm-Sidak method and was not significant (P > 0.05). Viral antigen in lung tissues was visualized by IHC using antibody against PR8 HA protein followed by alkaline phosphatase with the immPACT red substrate. Cytospins from BALF were stained with Hema 3 to identify airway neutrophils. At least 300 total cells were counted from duplicate cytospins from each animal, and the percentage of neutrophils was calculated. Data shown are from individual mice, with means and standard errors indicated. Statistical significance between Ctl Ab- and IFNAR Ab-treated mice was determined using unpaired t tests corrected for multiple comparisons using the Holm-Sidak method and was not significant (P > 0.05). Histopathology in lung tissues was visualized by hematoxylin and eosin staining on day 6 after PR8 infection. Arrows indicate examples of pulmonary hemorrhage. Neutrophils in lung tissues were visualized by IHC using antibody against Ly6G followed by horseradish peroxidase and the diaminobenzidine (immPACT DAB, brown) substrate. Images in panels B, D, and E are representative of results from two mice per group and time point.
bacteria or parasitic infections can reduce the severity of PVM, which corresponds to reduced pulmonary inflammation. When given intranasally on days −14 and −7 prior to or on days 1 and 2 after PVM inoculation, *Lactobacillus* spp. prevent lethal viral infection (28, 64, 65). Bacterial priming is associated with reduced production of proinflammatory cytokines and chemokines and inflammatory cell recruitment to the lungs, especially neutrophils (28, 64, 65).

FIG 10 Mucin-related genes have increased expression in RV-inoculated mice, and IFNAR inhibition does not prevent muc5ac production. (A) Normalized read count data for the mucin 5ac (Muc5ac) and chloride channel accessory 1 (Clca1) genes, which are required for mucus production and hydration, respectively. (B) Immunoblotting of Muc5ac protein and quantification relative to tubulin from RV-treated mouse lungs with and without IFNAR inhibition. Statistical significance was determined using unpaired t tests with *P* values indicated. Western blot is representative of two blots including five animals per group.
Similarly, chronic schistosomiasis protects mice from lethal PVM infection, which corresponds to the upregulation of mucus secretion pathways (32).

We found that RV-treated mice had increased expression levels of several genes in the mucin production pathway. Muc5ac, the predominant gel-forming mucin in the airways of humans and mice, is produced by goblet cells in the airway epithelium, and Clca1 is a chloride channel regulator that is needed for proper mucus hydration. The forkhead transcription factor FoxA3 induces the expression of multiple genes in the mucus production pathway that we found to be concurrently upregulated, including Muc5ac, Muc5b, Agr2, and Itln1 (66). Chen et al. demonstrated increased expression of FoxA3 and Muc5ac and goblet cell metaplasia in the airway epithelium of mice 3 days after infection by RV (66), and additional studies have reported the upregulation of Muc5ac mRNA and protein in the lungs and airways of RV-infected mice (39, 66, 67).

Influenza A virus strains, including PR8, also induce the expression of Muc5ac and mucous secretion (68–70). While stimulation of mucus production can promote pathology in chronic airway diseases such as asthma and chronic obstructive pulmonary disease (COPD), acute mucus production in hosts with healthy airways promotes innate defense against pathogen invasion. The overexpression of Muc5ac in mice has been shown to reduce PR8 infection and neutrophil recruitment (33). Muc5ac also promotes neutrophil transmigration and recruitment (71); thus, the reduction in neutrophils seen by Ehre et al. may be a consequence of lower levels of viral replication. Muc5ac has also been shown to reduce the severity of PR8 and PVM infections in a parasite coinfection model (32). The production of Muc5ac is induced by multiple inflammatory mediators, including IL-4, IL-13, epidermal growth factor (EGF), and transforming growth factor α (TGF-α), and is enhanced by additional signaling molecules through mitogen-activated protein kinase (MAPK)-dependent pathways (72). Muc5ac is also upregulated by IFN-β in human bronchial and alveolar epithelial cells and mouse lung tissues and is dependent on signaling through aryl hydrocarbon receptor (AhR) (73). The upregulation of Muc5ac by RV was not prevented by the inhibition of IFNAR signaling, and thus, Muc5ac is likely not sufficient to prevent PR8 lethality. Future studies will address the potential role of mucins in RV-mediated protection against PVM disease.

We found dramatic differences in the kinetics of IFN response gene expression in PR8- versus PVM-infected mice, which corresponded to differences in the requirement for IFNAR signaling in protection against PR8 but not PVM. Although PVM encodes two nonstructural proteins with potent antagonist activity against type I IFN, treatment of mice with type I IFN prior to PVM infection is protective (22). It is possible that the IFN response to RV is not robust enough to account for protection against PVM, or multiple, redundant mechanisms may contribute to protection, for example, type I and type III IFNs (22).

Type I IFN signaling is important for immediate antiviral defense mechanisms as well as orchestrating the correct balance of immune cells responding to infection. Others have shown that type I IFN signaling is critical for orchestrating monocye responses to PR8 infection, which in turn limits neutrophilic inflammation (20). We observed increased numbers of neutrophils in BALF and throughout lung tissues in RV/PR8-coinfected mice that were treated with an IFNAR-inhibiting antibody. These findings demonstrate that type I IFN can provide protection during respiratory viral infections that is independent of direct inhibition of viral replication and correlates with reduced neutrophilic inflammation and tissue damage. Type I IFN responses are also important in the activation of T cell-mediated immune responses (74–76). While they did not reach statistical significance, the numbers of CD4+ and CD8+ T cells in the lungs were modestly increased in RV/PR8- compared to mock/PR8-infected mice on day 6. Future studies to evaluate potential functional differences in lung-resident and recruited T cell subsets and their roles in protection will be critical for a complete understanding of the type I IFN-dependent mechanisms whereby RV reduces the severity of PR8 infection.
MATERIALS AND METHODS

Ethics statement. All mouse procedures were approved by the University of Idaho (UI) Institutional Animal Care and Use Committee, in compliance with the NIH Guide for the Care and Use of Laboratory Animals (77). Female 6- to 7-week-old BALB/c mice were ordered from InvivoGen/Envigo and were allowed to acclimate for 10 days prior to experimentation. All mice were housed in the UI Laboratory Animal Research Facility under 12-h light/dark cycles, received food and water ad libitum, and were monitored daily for any signs of distress. Mice were humanely euthanized if they reached endpoints including more than 25% weight loss and/or severe clinical signs of disease.

Virus infections. Viruses used in this study include PVM strain 15 (ATCC VR-25), RV1B strain B632 (ATCC VR-1645), and influenza A virus strain A/Puerto Rico/8/34 (H1N1) (catalog number NR-3169; BEI Resources). Viruses were grown and titrated by TCID$_{50}$ assays in BHK21 (PVM), HeLa (RV1B), and MDCK (PR8) cell lines.

Mice were anesthetized using isoflurane during intranasal inoculation. RV/PVM timing experiments were performed using groups of seven mice. Mice were either mock inoculated (phosphate-buffered saline [PBS]–2% fetal bovine serum [FBS]) or inoculated with 7.6 × 10$^6$ TCID$_{50}$ of RV1B 2 days before, simultaneously with, or 2 days after 1.0 × 10$^6$ TCID$_{50}$ of PVM. Mice were then monitored daily for mortality, weight loss, and clinical signs of disease (ruffled fur, hunched posture, lethargy, and labored breathing). Clinical signs in these four categories were scored on a scale of 0 to 3 (0, none; 3, severe). Humane euthanasia was performed when mice lost more than 25% of their starting weight or exhibited severe clinical signs of disease. Survival and weight loss data were analyzed with Prism 8.0 software (GraphPad) using a log rank Mantel-Cox test and Student’s t test corrected for multiple comparisons using the Holm-Sidak method, respectively.

RNA-seq and flow cytometry experiments were performed using groups of five mice. Mice were either mock inoculated (PBS–2% FBS) or inoculated with 7.6 × 10$^6$ TCID$_{50}$ RV1B intranasally on day −2. On day 0, mice were euthanized and 50 TCID$_{50}$ of PR8 or 1.0 × 10$^6$ TCID$_{50}$ of PVM. Mice were euthanized on days 0, 2, 4, and 6 to collect lungs for analyses. Left lobes were used for flow cytometry analysis, and right lobes were placed in RNAlater for RNA-seq analysis (see below). We consistently see infection in the right and left lobes upon intranasal inoculation of PR8 or PVM in 50-μl volumes (data not shown).

Signaling by interferon alpha receptor 1 (IFNAR1) was inhibited by treating mice with 0.05 mg anti-IFNAR1 antibody (clone MAR1-5A3; Bio X Cell) intranasally on days −2 and 0 with virus or mock inoculations. Antibody of the same isotype (mouse IgG1, clone MOPC-21; Bio X Cell) was used as a negative control. Mice were monitored and weighed daily as described above, and groups of 5 mice were euthanized at predetermined time points to quantify viral replication, IFN-induced gene expression, and cellular infiltration.

RNA-seq analysis. RNA was extracted from mouse lung tissue according to the RNeasy Plus with genomic DNA (gDNA) removal protocol (Qiagen) and quantified using an HS RNA kit and fragment analyzer (Advanced Analytical). The three samples with the highest RNA quality from each group were used for RNA-seq analysis. Stranded RNA libraries were prepared from 4 μg RNA of each sample by UI’s iBEST Genomics Resources Core according to Kapa’s stranded mRNA-seq (catalog number KB8420) library preparation protocol with capture of polyadenylated mRNAs. Libraries were tagged with unique ligation adaptors (BioOScientific), amplified, quantified (Qubit and AATI fragment analyzer), and pooled at an equimolar ratio. The pooled library was split, pooled with other libraries, and sequenced across 5 lanes of an Illumina HiSeq4000 100-bp paired-end platform run at the University of Oregon’s Genomics and Cell Characterization Core. Paired-end reads (100 bp) were quality trimmed and filtered using Trimomatic v0.36 (ILLUMINACLIP:2:20:10, HEADCROP:10, SLIDINGWINDOW:4:15, and MINLEN:36) and mapped to the mouse genome (GRCm38/mm10, downloaded from the UCSC database) using TopHat v2.1.1 (r 300) (78, 79). The TopHat alignments were the starting points for two methods of quantifying read counts and four methods of analyzing differential expression (DE). Data from the Cufflinks v2.1.1 package (Cufflinks, Cuffmerge, and Cuffdiff) were compared to those from DE pipelines using HTSeq v0.8.0 (htseq-count -m intersection-nonempty -s reverse -t exon) followed by DESeq2 v1.18.1 and EdgeR 3.18.1 (80, 81). DESeq2 and EdgeR were run within the SARTools wrapper (82). An adjusted P value cutoff of 0.05 was used to determine significance.

(i) Analysis of viral gene expression. Reads that did not map to the mouse genome were extracted and deinterleaved using the BBmap reformat script (http://sourceforge.net/projects/bbmap/) to give paired reads for mapping. These reads were aligned to the RV1B (GenBank accession number D00239), PR8 (GenBank accession numbers LC120388 to LC120395), and PVM (GenBank accession number AY729016) genomes using TopHat v2.1.1, and gene coverages were counted using HTSeq v0.8.0 (see the parameters described above). Statistical significance was determined using a generalized linear model in R.

(ii) DEG function analysis. The DEG lists for each treatment comparison were searched for differentially enriched gene ontology (GO) terms and KEGG pathways using CompGO (83). CompGO utilizes gene expression level data (as opposed to just gene names) to identify potential functions that differ between treatments. We ran CompGO on DEG lists from each method (Cufflinks, EdgeR, and DESeq2) and combinations of gene sets to anecdotally look for consistency across methods.

RT-qPCR. RNA was extracted from lung tissues using TRIzol (Invitrogen) or RNeasy Plus (Qiagen) reagents according to the manufacturers’ protocols. cDNA synthesis was performed using SuperScript IV Vilo master mix (Invitrogen), and qPCR was done with PowerUp SYBR green (Applied Biosystems) using a StepOnePlus real-time instrument (Applied Biosystems). Published primer sets were used to quantify PR8 HA (84), PVM SH (85), IFN-β (86), Mx1 (74), glyceraldehyde-3-phosphate dehydrogenase (GAPDH) (87), and β-actin (88). Mx1 and IFN-β threshold cycle (C$_t$) values were normalized to GAPDH, and the
resulting ΔCt values were normalized to those from mock-inoculated mice and expressed as 2−ΔΔCt values. Ct values of viral genes were normalized to β-actin and expressed as 2−ΔCt values.

Flow cytometry. The left lung lobe of each sample was dissociated in RPMI 1640 medium containing type IV collagenase (MP Biomedicals) and DNase I (Sigma) using gentleMACS C tubes according to the manufacturer’s recommendations (Miltenyi Biotec). Cells were filtered through a 70-μm strainer, and red blood cells were lysed before blocking Fc receptors using anti-CD16/CD32 (BioLegend). Cell-specific proteins were labeled with CD11b-Alexa Fluor 488 (eBioscience), Ly6G-allophycocyanin (APC) (eBioscience), CD64-phycocerythrin (PE) (BioLegend), or S IgG-F-peridinin chlorophyll protein (PerCP)-Cy5.5 (BD Biosciences) antibody or the appropriate isotype control antibodies. T cell subsets were stained with antibodies against CD3-PerCP-Cy5.5 (eBioscience), CD8-APC (eBioscience), and CD4-Alexa Fluor 488 (BioLegend). Stained cells were incubated in BD stabilizing fixative (BD Biosciences) and analyzed using a FACSARia cytometer (BD Biosciences). Results were analyzed using FlowJo software (TreeStar), and gating was performed based on fluorescence-minus-one controls. The gating strategy used to identify neutrophils, alveolar macrophages, and interstitial macrophages is shown in Fig. S7 in the supplemental material. T cells were gated as CD3+/CD4+ T helper cells and CD3+CD8+ cytotoxic T cells.

Quantification of viral loads. Infectious virus was quantified in homogenized lungs or BALF samples by TCID50 assays in HeLa (RV), MDCK (PR8), and BHK21 (PVM) cell lines. We have previously found that the presence of RV in samples does not interfere with the titration of PR8 or PVM in MDCK cells or BHK21 cells, respectively (9).

Analysis of BALF cells. BALF was collected by tracheal cannulation and lavage with 1-ml PBS twice per sample. Collected cells were counted and spun onto microscope slides in duplicate samples, which were stained with Hema 3 staining solutions (Fisher Scientific). Gridded coverslips and microscopy were used to count at least 300 neutrophils, macrophages, and lymphocytes per sample, and these counts were used to calculate the percentage of each cell type.

Histology and immunohistochemistry. Lung tissues from two animals per treatment group and time point were processed and stained as previously described (9). Antibodies specific for PR8 hemagglutinin protein (catalog number RR-3148; BEI Resources) and Ly6G (clone 1A8; Bio X Cell) were detected with immPACT vector red and diaminobenzidine immPACT DAB, respectively (Vector Laboratories).

Western blotting. Whole-lung homogenates were separated by low-speed centrifugation, and the pellets were lysed in radioimmunoprecipitation assay (RIPA) buffer. Muc5ac-specific antibody (clone 45.1; Novus) (89) was used to probe Western blots as described previously (90). An alpha-tubulin-specific antibody (clone DM1A; Abcam) was used as a protein loading control. Blots were imaged on an Amersham 600 imager, and densitometry was performed using Bio-Rad software.

Statistical analyses. We analyzed flow cytometry (Fig. 6) and viral read count (Fig. 4) data resulting from our experiments to identify time-varying differences between mice infected with PR8 or PVM alone and those pretreated with RV. To this end, we used negative binomial regression on each response variable with an explanatory model that had a main effect of days postinfection, a main effect of treatment, and the interaction between the two main effects. Response variables were the numbers of CD11b cells, neutrophils, interstitial macrophages, alveolar macrophages, CD4+ T cells, CD8+ T cells, and viral read counts; all response variables were normalized versus the total cell count, except for the viral RNA read count, which was normalized against the total RNA read count. Due to a priori visual investigation of our data, it seemed that some of our response variables might be better fit with a quadratic time term. Because of this, we fit an alternative model that included an orthogonal polynomial of degree 2 for time. We assessed whether the quadratic model was better than the linear model using a likelihood ratio test and chose the quadratic model if it offered a significant improvement in fit over the simpler linear model. The significance of treatment and time was determined using type I analysis of variance (ANOVA). To detect differences between treatments at a given time, we also performed post hoc pairwise comparisons of the modeled means at our observational time points (days 0, 2, 4, and 6) using the emmeans package in R. Table S3 provides information on the analyses, including which time model was used and the significance of treatment and time. Significant contrasts are shown in individual figures.

 Survival curve, weight loss, viral titer, RT-qPCR, and BALF cell count data were analyzed with Prism 8.0 software (GraphPad) using the methods described in each figure legend.

Data availability. Raw Illumina reads have been deposited in the NCBI SRA database under accession numbers SRR7060116 to SRR706162 under BioProject accession number PRJNA453386.

SUPPLEMENTAL MATERIAL
Supplemental material is available online only.

FIG S1, PDF file, 0.1 MB.
FIG S2, TIF file, 0.7 MB.
FIG S3, TIF file, 0.6 MB.
FIG S4, TIF file, 0.9 MB.
FIG S5, TIF file, 1.3 MB.
FIG S6, TIF file, 1 MB.
FIG S7, TIF file, 1.2 MB.
TABLE S1, XLSX file, 1.2 MB.
TABLE S2, XLSX file, 2.4 MB.
TABLE S3, XLSX file, 0.01 MB.
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