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Rapid Short Communication

Type I interferon receptor signalling deficiency results in dysregulated innate immune responses to SARS-CoV-2 in mice

Patricia P. Ogger1, Minerva Garcia Martín1, Christina Michalaki1, Jie Zhou2, Jonathan C. Brown2, Yue Du3, Kamran M. Miah3, Omar Habib3, Stephen C. Hyde3, Deborah R. Gill3, Wendy S. Barclay2 and Cecilia Johansson1

1 Section of Respiratory Infections, National Heart and Lung Institute, Imperial College London, St Mary’s Campus, London, UK
2 Department of Infectious Disease, Imperial College London, St Mary’s Campus, London, UK
3 Radcliffe Department of Medicine (NDCLS), University of Oxford, Oxford, UK

SARS-CoV-2 is a newly emerged coronavirus, causing the global pandemic of respiratory coronavirus disease (COVID-19). The type I interferon (IFN) pathway is of particular importance for anti-viral defense and recent studies identified that type I IFNs drive early inflammatory responses to SARS-CoV-2. Here, we use a mouse model of SARS-CoV-2 infection, facilitating viral entry by intranasal recombinant Adeno-Associated Virus (rAAV) transduction of hACE2 in wildtype (WT) and type I IFN receptor-1 deficient (Ifnar1–/–) mice, to study the role of type I IFN signalling and innate immune responses during SARS-CoV-2 infection. Our data show that type I IFN signalling is essential for inducing anti-viral effector responses to SARS-CoV-2, control of virus replication, and to prevent enhanced disease. Furthermore, hACE2-Ifnar1–/– mice had increased gene expression of the chemokine Cxcl1 and airway infiltration of neutrophils as well as reduced and delayed production of monocyte-recruiting chemokine CCL2. hACE2-Ifnar1–/– mice showed altered recruitment of inflammatory myeloid cells to the lung upon SARS-CoV-2 infection, with a shift from Ly6C+ to Ly6C– expressing cells. Together, our findings suggest that type I IFN signalling deficiency results in a dysregulated innate immune response to SARS-CoV-2 infection.

Keywords: innate immune response · type I IFN · in vivo · SARS-CoV-2 · myeloid cells

Introduction

The ongoing coronavirus disease (COVID-19) pandemic caused by severe acute respiratory syndrome-coronavirus 2 (SARS-CoV-2) has resulted in over 400 million cases in the first two years of the pandemic. The estimated fatality rate lies between 1–2%, however this is considerably higher for elderly patients over 80 years of age (~10%) and nursing home residents (>20%) [1]. Type I interferons (IFNs) are one of the first responses elicited against viral infection and they induce anti-viral defense mechanisms by binding to the IFN-α/β receptor (IFNAR) and signalling through...
the JAK-STAT pathway. This induces expression of IFN stimulated genes (ISG), resulting in expression of anti-viral effector proteins that restrict viral replication and activation of immune cells via induction of chemokine and cytokine production [2], including CXCL10 and CCL2 [3]. Early in vivo studies using human bronchial epithelial cell lines infected with SARS-CoV-2 showed decreased production of type I and III IFNs coupled with low anti-viral defense signals and a pro-inflammatory environment compared to infection with influenza A virus (IAV) [4]. Furthermore, in severe and critically ill COVID-19 patients, an impaired type I IFN response has been observed, resulting in decreased viral clearance [5]. A lack of an efficient type I IFN response in these patients is in part due to inborn errors of type I IFN immunity [6] or circulating auto-antibodies neutralising type I IFNs [7]. Also, a recent animal study has identified that type I IFN signalling is required for the recruitment of pro-inflammatory cells into the lungs following SARS-CoV-2 infection [8]. These findings highlight the importance of functional type I IFN responses for anti-viral defenses against SARS-CoV-2.

Using a mouse model of SARS-CoV-2 infection, facilitated by intranasal recombinant Adeno-Associated Virus (rAAV) induced expression of human angiotensin-converting enzyme 2 (hACE2), this study investigated the dynamics of innate immune responses to infection with SARS-CoV-2 in the context of type I IFN receptor signalling impairment. Overall, the data show that type I IFN receptor signalling is essential to induce anti-viral responses and control viral replication and disease severity during SARS-CoV-2 infection. Furthermore, type I IFN receptor-deficient mice show dysregulated innate immune responses to SARS-CoV-2 infection, marked by increased neutrophil recruitment into the airways and delayed recruitment of myeloid inflammatory cells to the lungs.

Results and discussion

To investigate the dynamics of innate, anti-viral immune responses to SARS-CoV-2 infection in the context of type I IFN-signalling deficiency, 8–12-week-old C57BL/6 wildtype (WT) or interferon alpha receptor-1−/− (Ifnar1−/−) mice were intranasally transduced with rAAV9 containing either hACE2 or eGFP (control) as published recently [9], followed by intranasal infection with 2×10^6 PFU SARS-CoV-2 (D614G, first wave isolate) 20 days later and study endpoints were at 2, 4 and 8 days post-infection (d.p.i.) (Figure 1A). Gene expression analysis of hACE2 in lung tissue 20 days post administration of rAAV9 (before infection) showed similar expression in WT and Ifnar1−/− mice relative to Gapdh (Figure 1B). Cryo-sectioning of lung tissue 20 days post administration of rAAV9-eGFP (before infection) furthermore showed a similar distribution of eGFP in WT and Ifnar1−/− mice (Supp. Figure 1A), suggesting similar rAAV transduction efficacy in both groups of mice. Upon infection with SARS-CoV-2, hACE2-Ifnar1−/− mice showed increased viral loads, peaking at 6 d.p.i., compared with hACE2-WT mice (Figure 1C). Furthermore, plaque assays on Vero cells overexpressing SARS-CoV-2 binding receptors ACE2 and transmembrane protease serine 2 precursor TMPRSS2 (VAT cells) showed that IFNAR1-deficiency results in significantly higher viral loads at days 2 and 4 post-infection with SARS-CoV-2 compared to hACE2-WT mice (Figure 1D). Gene expression of SARS-CoV-2 nucleocapsid (N) and envelope (E) genes was significantly higher in hACE2-Ifnar1−/− mice compared with hACE2-WT mice from 2 d.p.i. (Figure 1B). Since AAV-eGFP transduced Ifnar1−/− mice did not become infected with SARS-CoV-2 and did overall not differ significantly from AAV-eGFP transduced WT mice (Supp. Figure 2), this group was not included in all experiments to reduce animal numbers. Together, these data show increased viral load in hACE2 expressing IFNAR1-deficient mice upon SARS-CoV-2 infection, measured both by plaque assay and N and E gene expression (Figure 1D-E and Supp. Figure 3A-B). Previous studies have shown similar trends [8], although differences between WT and Ifnar1−/− are more pronounced in the model used here. This could be due to several factors: here, Vero cells overexpressing hACE2 and TMPRSS2 were used for plaque assays, allowing for better viral replication, while the isolate used for infection was hCoV-19/England/IC19/2020, which harbors the D614G spike mutation as opposed to USA-WA1/2020, harboring viral replication [10]. Together, these factors might explain why we detected higher viral loads and more prominent differences emerged between hACE2-WT and hACE2-Ifnar1−/− mice.

To investigate the impact of impaired type I IFN signalling during SARS-CoV-2 infection, we first assessed type I IFN expression upon infection. At 2 d.p.i. both IFN-α and IFN-β were significantly increased in BAL fluid of hACE2-WT compared to eGFP-WT mice (Supp. Figure 1B). In the hACE2-Ifnar1−/− group, expression of IFN-α was significantly lower, while IFN-β levels were higher compared to the hACE2-WT group. We next investigated ISG expression after SARS-CoV-2 infection, since in Ifnar1−/− mice limited amounts of type I IFNs can be produced but cannot signal for downstream ISG induction. Chemokine Cxcl10 and anti-viral effectors Mr1, Oas1 and Viperin were quantified. The expression of these ISGs was increased in hACE2-WT mice upon infection with SARS-CoV-2 at 2 d.p.i., however in hACE2-Ifnar1−/− mice, ISG expression was significantly reduced, but not completely absent (Figure 1F and Supp. Figure 3C-F). These results suggest an initial increase of IFN-β in hACE2-Ifnar1−/− mice, which may be due to higher viral titers, but that is not translated into ISG expression due to the lack of signalling through the IFNAR1. We therefore investigated the expression of type III IFNs, IFN-λ-3, which can contribute to ISG expression. IFN-λ expression was induced upon infection in hACE2-WT mice at 2 d.p.i. while remaining at baseline levels in the hACE2-Ifnar1−/− mice (Supp. Figure 1C). Furthermore, Ifng gene expression was significantly increased in hACE2-Ifnar1−/− mice later during the infection, by 4 d.p.i. (Supp. Figure 4A). This correlated with CD3+ T cell recruitment to the airways (Supp. Figure 4B-D). Our data suggest that limited levels of type I or III IFNs are produced early during infection in the Ifnar1−/− mice resulting in some ISG expression but overall, type I IFN receptor signalling is the main driver for inducing cell intrinsic anti-viral responses.

We next assessed the gene expression of inflammatory mediators and found that expression of the chemokine Cxcl1, which is
Increased viral load, weight loss, and lower expression of ISGs in Ifnar1−/− mice during infection with SARS-CoV-2. (A) Recombinant Adeno-associated virus (rAAV) containing human angiotensin-converting enzyme 2 (hACE2) or eGFP genes was administered intranasally to Ifnar1−/− or wildtype (WT) mice (1 × 10^{11} Dnase resistant gene copies/mouse). 20 days later mice were intranasally infected with SARS-CoV-2 (D614G, 2 × 10^6 PFU/mouse). Lungs and bronchoalveolar lavage (BAL) were harvested at 2, 4, and 8 days post-infection (d.p.i.). (B) Expression of hACE2 in lung tissue relative to Gapdh, measured by RT-PCR before infection (d20 post-transduction with rAAV). WT/rAAV-eGFP n = 3, Ifnar1−/−/rAAV-eGFP n = 4, WT/rAAV-hACE2 n = 4, Ifnar1−/−/rAAV-hACE2 n = 5. (C) Weight loss post-infection with SARS-CoV-2. (D) Viral load measured by plaque assay on Vero cells overexpressing hACE2 and TMPRESS2. (E) Expression of SARS-CoV-2 N gene (nucleocapsid phosphoprotein) and E gene (envelope protein) in lung tissue relative to Gapdh, measured by RT-PCR. (F) Gene expression analysis of IFN stimulated genes (ISG) Cxcl10, Mx1, Oas1, and Viperin measured by RT-PCR, relative to expression of Gapdh or total copy numbers normalised to expression of Gapdh (Oas1 and Viperin). Data are shown as mean ± SEM. B) WT/rAAV-eGFP n = 3, WT/rAAV-hACE2 n = 4, Ifnar1−/−/rAAV-hACE2 n = 5 C – F) Two independent experiments per time point, data pooled, n = 6–8 per group. One Way ANOVA + Tukey’s multiple comparison test per time point; * indicates significant difference between WT rAAV-hACE2 and Ifnar1−/− rAAV-hACE2, *P < 0.05, **P < 0.01, ***P < 0.005, ****P < 0.001.

not dependent on type I IFN receptor signalling [3], was increased in hACE2-Ifnar1−/− mice at 2 and 4 d.p.i. compared to hACE2-WT mice (Figure 2A and Supp. Figure 3G). As CXCL1 plays an essential role in early host immune responses by recruiting neutrophils [11], we next analysed the infiltration of neutrophils (gated as live, CD45+, Ly6 G+, Supp. Figure 5A) into the airways at 2 d.p.i. In line with highly increased gene expression of Cxcl1, neutrophil recruitment to the airways (BAL) was significantly increased in hACE2-Ifnar1−/− mice at 2 d.p.i., both proportional of leukocytes (CD45+ cells) and in total numbers (Figure 2B and C). This was recapitulated in lung tissue with increased proportions of neutrophils in type I IFN signalling-impaired mice at 2 d.p.i. (Figure 2D and Supp. Figure 3H), decreasing over time. Taken together, these findings suggest that during SARS-CoV-2 infection, type I IFN receptor signalling deficiency results in increased neutrophil recruitment via CXCL1, thereby contributing to a pro-inflammatory environment. Indeed, Cxcl1 is also increased in Ifnar1−/− mice during influenza A virus infection with secondary pneumococcal infection [12], but decreased during RSV infection in mice [13], highlighting a pathogen-specific CXCL1 response. Furthermore, since we show similar trends for viral load and neutrophil recruitment upon SARS-CoV-2 infection (both significantly increased in hACE2-Ifnar1−/− mice), which is not detected in other respiratory viral infections such as respiratory syncytial virus (RSV) [13, 14], it will be important to further investigate the link between neutrophil recruitment and viral load in this model.
Figure 2. Increased neutrophil recruitment to airways in IFNAR1-deficient mice during SARS-CoV-2 infection. (A) Gene expression of Cxcl1 in lung tissue relative to Gapdh at 2, 4, and 8 d.p.i. with SARS-CoV-2 (D614G), measured by RT-PCR. (B) Proportions of live, CD45+, and total numbers of neutrophils in BAL at 2 d.p.i. (C) Representative flow cytometry plots of lung cells gated on live, CD45+Ly6G+. (D) Proportions of live, CD45+ and total numbers of neutrophils in lung tissue at 2, 4, and 8 d.p.i. Data are shown as mean ± SEM. Two independent experiments per time point, data pooled, n = 6–8 per group. One Way ANOVA + Tukey’s multiple comparison test per time point; * indicates significant difference between hACE2-WT and hACE2-Ifnar1−/−, *P < 0.05, **P < 0.01, ***P < 0.005, ****P < 0.001.

Since monocyte recruitment to the airways and lungs is key to early host responses to viral infection, we next investigated the expression of monocyte recruiting chemokine CCL2 and the recruitment of inflammatory myeloid cells. CCL2 protein expression was increased in BAL fluid of hACE2-WT mice at 2 d.p.i. with SARS-CoV-2 (Figure 3A and Supp. Figure 3I). However, in hACE2-Ifnar1−/− mice CCL2 expression was significantly lower at 2 d.p.i., peaking at 4 d.p.i. at lower levels than in IFNAR1-sufficient mice (Figure 3A). These findings are in line with a report identifying early CCR2 signalling essential to restrict viral burden in a mouse model of SARS-CoV-2 infection [15]. The recruitment of CD64+CD11b+ inflammatory myeloid cells to the lung followed similar kinetics, as in hACE2-WT proportions were highest at 2 d.p.i. and subsequently decreased, while in IFNAR1-deficient mice proportions and total numbers of CD64+CD11b+ inflammatory myeloid cells increased between 2 and 4 d.p.i. and were highest at 8 d.p.i. (Figure 3B and Supp. Figure 3J and 5A). We next assessed expression of the monocyte/macrophage differentiation antigen Ly6C within this population, since previous studies reported the infiltration of CD64+CD11b+Ly6C+ inflammatory myeloid cells into the lung during SARS-CoV-2 infection [8, 15]. This showed highly increased proportions of CD64+CD11b+Ly6C+ in hACE2-WT but not in IFNAR1-deficient mice at 2 d.p.i. in the BAL (Supp. Figure 5B-C) and lung (Figure 3C-D and Supp. Figure 3K), suggesting type I IFN receptor dependency for recruitment. However, as we have previously shown that Ly6C is gradually downregulated on monocytes during response to respiratory viral infection [16], we also analysed CD64+CD11b+Ly6C− cells. The presence of CD64+CD11b+Ly6C− inflammatory myeloid cells in the airways was not type I IFN signalling dependent, since both proportions and total numbers were significantly increased in hACE2-Ifnar1−/− mice at 4 and 8 d.p.i. (Figure 3C and E), while at 2 d.p.i. in the airways no significant differences emerged (Supp. Figure 5C). This accounts for the delayed emergence of inflammatory myeloid cells in the lung during type I IFN signalling impairment shown in Figure 3B and overall indicates altered recruitment dynamics of inflammatory myeloid cells. Taking these data together, our model recapitulates the deficiency of type I interferon responses seen in severe SARS-CoV-2 infection, which in patients is marked by decreased IFN-α, type I IFN activity and ISG score, as well as neutrophilia and increased CCL2 [5]. Our data suggest that the lack of type I IFN receptor signalling results in dysregulated innate immune responses in the lung during SARS-CoV-2 infection.

Concluding Remarks

In summary, using a mouse model of SARS-CoV-2 infection we show that type I IFN receptor signalling is essential for inducing
Figure 3. Type I interferon signaling deficiency results in dysregulated inflammatory myeloid cell recruitment during SARS-CoV-2 infection. (A) Protein expression of CCL2 in BAL fluid at 2, 4, and 8 d.p.i. measured by ELISA. (B) Proportions of live, CD45+ and total numbers of CD64+CD11b+ inflammatory myeloid cells in lung tissue at 2, 4, and 8 d.p.i. (C) Representative flow cytometry plots of lung cells gated on live, CD45+Ly6G−SigF−CD11b+. (D) Proportions of live, CD45+ and total numbers of CD64+CD11b+ Ly6C+ inflammatory myeloid cells in lung tissue at 2, 4, and 8 d.p.i. (E) Proportions of live, CD45+, and total numbers of CD64+CD11b+ Ly6C− inflammatory myeloid cells in lung tissue at 2, 4, and 8 d.p.i. Data are shown as mean ± SEM. Two independent experiments per time point, data pooled, n = 6–8 per group. One Way ANOVA + Tukey’s multiple comparison test per time point; * indicates significant difference between hACE2-WT and hACE2-Ifnar1−/−, **P < 0.05, ***P < 0.01, ****P < 0.005, *****P < 0.001. Dotted line = limit of detection.

Our data indicate that type I IFN signalling-deficient mice express increased levels of Cxcl1 in the lung and present with increased infiltration of neutrophils to the airways compared to WT controls. Furthermore, we found reduced and delayed production of CCL2 and altered recruitment of inflammatory myeloid cells during IFNAR1-deficiency. This, together with an increased viral burden is associated with more severe disease in type I IFN signalling-deficient mice. The data shown here will be valuable for better understanding how impaired type I IFN signalling drives SARS-CoV-2 pathology and disease severity, which is highly relevant considering the large contribution of impaired type I IFN responses on life-threatening SARS-CoV-2 infections [6, 7] and deaths [17] and for the development of type I IFN-based treatment options for COVID-19 in vulnerable populations. To conclude, our findings show that type I IFN receptor deficiency results in dysregulated innate immune responses to SARS-CoV-2 infection in the rAAV-hACE2 mouse model.

Materials and methods

Mice

C57BL/6 mice were purchased from Charles River UK Inc. Ifnar1−/− mice on a C57BL/6 background were bred in-house. All mice were bred and maintained in pathogen-free conditions and 8–12-week-old mice were used for experiments. All animal experiments were reviewed and approved by the Animal Welfare and Ethical Review Board (AWERB) at Imperial College London and approved by the UK Home Office in accordance with the Animals
Act 1986 (Scientific Procedures) and ARRIVE guidelines. Both male and female mice were used for experiments after excluding sex bias in preliminary experiments. All experiments were performed twice, independently, per time point.

**rAAV vector production**

The production, purification, and titration of rAAV2/9-eGFP or hACE2 vectors were performed as previously described [9]. Briefly, the respective rAAV vector was produced by polyethyleneimine (PEI, Polysciences)-based triple transfection of human embryonic kidney (HEK) 293T/17 cells (ATCC, CRL-11268). The AAV plasmids transfected included the Adenovirus helper plasmid (pAdDeltaF6), AAV Rep-Cap pAAV2/9 plasmid, and the transgene plasmid. The transgene plasmid containing eGFP or hACE2 was engineered to include a lung-optimized hCEFI (human Cytomegalovirus enhancer/elongation factor 1 alpha) promoter [18], Woodchuck Hepatitis Virus Post-transcriptional Regulatory Element (WPRE) [19] and mir142-3pT [20]. rAAV particles were concentrated and formulated into PBS using 100 kDa Ultra centrifugal filters (Amicon, Merck) after iodixanol gradient centrifugation. Physical titre (DNase-resistant genome copies, DRGC/mL) was determined by quantitative polymerase chain reaction (qPCR) analysis with primers and a probe against WPRE [21]. Purity of vectors was confirmed by analysing 20 μl of diluted vector on 4–12% SDS polyacrylamide gels, where total protein was visualised using Coomassie stain according to the manufacturer’s protocols (Life Technologies).

**hACE2 transduction**

For transduction, WT or Ifnar1−/− mice were lightly anesthetised and instilled i.n. with 1×1011 DNase Resistant Gene Copies (DRGC) rAAV9-eGFP or rAAV9-hACE2 in 100 μl PBS. hACE2 gene expression in whole lung homogenate was assessed at day 20 post instillation by relative quantification to Gapdh using primers and probes for hACE2 listed in the key resource table.

**Cryosectioning and native eGFP detection**

Mice were sacrificed 20 days post instillation of AAV-eGFP or PBS and lungs were removed after inflation with 4% PFA. After 24-hour fixation in 4% PFA, lungs were inflated with 30% sucrose and submerged in 30% sucrose for 24 hours. Lungs were subsequently inflated with 1:1 cryo embedding matrix (OCT)/30% sucrose and individual lobes were submerged in OCT/30% sucrose in plastic molds and frozen at -80°C. Left lungs were cryosectioned to produce 7 μm thick sections, mounted using DAPI-supplemented mounting media with coverslip, and eGFP expression was detected by fluorescent microscopy using the EVOS FL Auto 2 system (Thermo Scientific).

**Virus and infections**

First wave SARS-CoV-2 (D614G, isolate of hCoV-19/England/IC19/2020) was grown in African green monkey kidney cells overexpressing human ACE2 and TMPRSS2 (Vero-ACE2-TMPRSS2; VAT cells) [22]. For infection 20 days post-transduction with rAAVs, mice were lightly anesthetised and instilled i.n. with 2×10⁶ plaque-forming units (PFU) of SARS-CoV-2 in 100 μl volume. SARS-CoV-2 titer was assessed in lungs at 2, 4, and 8 d.p.i. using a plaque assay. In brief, serial dilutions of lung homogenate in serum-free Dulbecco’s Modified Eagle Medium (DMEM, containing 1% non-essential amino acids (NEAA), 100U/ml Penicillin, and 100 μg/ml Streptomycin) were performed and inoculated onto VAT cells for 1 h at 37°C. The inoculum was then removed and replaced with overlay medium (1x MEM, 0.2% w/v BSA, 0.16% w/v NaHCO₃, 10 mM HEPES, 2 mM L-Glutamine, 100 U/ml penicillin, 100 μg/ml streptomycin and 0.84% agarose). Plates were incubated for 3 days at 37°C before overlay was removed and cells were stained for 1 h at room temperature in 2x crystal violet solution. Virus plaques were counted and multiplied by the dilution factor to calculate titer as PFU/ml.

**Isolation of lung cells**

Mice were sacrificed at 0.75, 2, 4, and 8 d.p.i. and lungs were perfused with PBS. To obtain lung leukocytes, lung lobes were cut into smaller pieces and incubated in complete DMEM (dMEM, supplemented with 10% fetal bovine serum, 2mM L-glutamine, 100 U/ml penicillin, and 100 μg/ml streptomycin), 1mg/ml Collagenase D (Roche) and 30 μg/ml DNase I (Invitrogen) for 1h at 37°C and then mashed through a 100-μm filter (BD). Red blood cells were lysed using Ammonium-Chloride-Potassium buffer.

**BAL cell processing**

BAL was collected by flushing the lungs three times with 1 ml PBS supplemented with 5 mM EDTA (Life Technologies). BAL cells and supernatant were separated by centrifugation and BAL supernatants were exposed to UV light for 2 min to inactivate SARS-CoV-2. Red blood cells were lysed using Ammonium-Chloride-Potassium buffer.

**Flow cytometry**

After red blood cell lysis, lung and BAL cells were incubated for 30 min with fixable live-dead Aqua dye (Invitrogen), followed by fixation for 30 minutes with 4% paraformaldehyde (PFA) to inactivate virus. Cells were then incubated for 20 min with a purified rat IgG₂b anti-mouse CD16/CD32 receptor antibody (BD) to block Fc binding, followed by staining with fluorochrome-conjugated antibodies against CD45 (30-F11, BV605), CD26 (H194-112, BV711),...
RNA isolation and quantitative RT-PCR

Lung tissue was homogenised in TRIzol and RNA extraction performed according to manufacturer’s instructions. After the chloroform step, the aqueous phase containing RNA was further processed using the RNeasy Mini Kit (QIAGEN) according to manufacturer’s instructions. 2 μg RNA was reverse transcribed using a High-Capacity RNA-to-cDNA kit (Applied Biosystems) according to manufacturer’s instructions including DNase digestion. To quantify mRNA levels in lung tissue, quantitative RT-PCR reactions for Gapdh, Ifnl1, Ifn2, Viperin and Cxcl10 were performed using primers and probes as previously described [23]. Analysis was performed using the Quantitect Probe PCR Master Mix (QIAGEN) and the 7500 Fast real-time PCR System (Applied Biosystems). For absolute quantification, the exact number of copies of the gene of interest was calculated using a plasmid DNA standard curve, and for relative quantification, the exact number of copies of the gene of interest was calculated using a plasmid DNA standard curve, and the results were normalised to levels of Gapdh. Analysis was performed using the 7500 Fast System SDS Software (Applied Biosystems).

Statistical analysis

Statistical analysis was performed using Prism 9.2 (Graph-Pad Software). One-way ANOVA with Tukey’s post hoc test was used to compare multiple groups. Data are expressed as mean ± SEM, and for all tests a value of P < 0.05 was considered significant. *P < 0.05, **P < 0.01, ***P < 0.005, ****P < 0.001.

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Author contribution: P.P.O. designed, performed, and analysed the experiments and wrote the paper. M.G.M. and C.M. performed specific experiments and reviewed the paper. Y.D., D.R.G., S.C.H, O.H., and K.M.M. designed, manufactured and provided rAAV9-specific experiments and reviewed the paper. J.Z., J.C.B. and W.S.B. provided advice and the SARS-CoV-2 WT isolate (D614G) and reviewed the paper. C.J. supervised the project, designed the experiments, and wrote the paper.

Ethics approval: All animal experiments were reviewed and approved by the Animal Welfare and Ethical Review Board (AWERB) at Imperial College London and approved by the UK Home Office in accordance with the Animals (Scientific Procedures) Act 1986 Amendment Regulations (PPL P3AFF00DD Johansson).

Data availability statement: The data that support the findings of this study are available from the corresponding author upon reasonable request.

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References

1 O’Driscoll, M., Ribeiro Dos Santos, G., Wang, L., Cummings, D. A. T., Azman, A. S., Paireau, J., Fontanet, A. et al., Age-specific mortality and immunity patterns of SARS-CoV-2. Nature 2021. 590: 140–145.
Plante, J. A., Johansson, C., Vanderheiden, A., Goritzka, M., Hadjadj, J., Du, Y., Zhang, Q., Platanias, L. C., Y., Belperio, J. A. et al.: Alpha/Beta Interferon Receptor Signaling Amplifies influenza bacterial pneumonia in mice. Against SARS-CoV-2 infection. Drives Development of COVID-19. Møller, R. 2021. M. et al. Mucosal Immunology, CCR2 Signaling Restricts SARS-CoV-2 Infection. Early Proinflammatory Cytokine Production in the Lung during Respirations. Impaired type I interferon activity and inflammatory responses in severe COVID-19 patients. Science 2020. 370: eabd4570.

Bastard, P., Rosen, L. B., Zhang, Q., Michailidis, E., Hoffmann, H. H., Zhang, Y., Dorgham, K. et al.: Autoantibodies against type I IFNs in patients with life-threatening COVID-19. Science 2020. 370: eabd4585.

Isaebow, B., Song, E., Mao, T., Liu, P., Meir, A., Liu, F., Alfájarro, M. M. et al.: Mouse model of SARS-CoV-2 reveals inflammatory role of type i interferon signaling. J Exp Med 2020. 217: e20201241.

Du, Y., Miáh, K. M., Habib, O., Meyer-Berg, H., Conway, C. C., Viegas, M. A., Dean, R. et al.: Lung directed antibody gene transfer confers protection against SARS-CoV-2 infection. Thorax 2022;thoraxjnl-2021-217650.

Plante, J. A., Liu, Y., Liu, J., Xia, H., Johnson, B. A., Lokugamage, K. G., Lokugamage, K. G. et al.: Spike mutation D614G alters SARS-CoV-2 fitness. Nature 2021. 592: 116–121.

Johansson, C., Kirsebom, F. C. M.: Neutrophils in respiratory viral infections. Mucosal Immunology 2021. 14: 815–827.

Shahangian, A., Chow, E. K., Tian, X., Kang, J. R., Chaffari, A., Liu, S. Y., Belperio, J. A. et al.: Type I IFNs mediate development of postinfluenza bacterial pneumonia in mice. J Clin Invest 2009. 119: 1910–1920.

Goritzka, M., Durant, L. R., Pereira, C., Salek-Ardakani, S., Openshaw, P. J. M. and Johansson, C.: Alpha/Beta Interferon Receptor Signaling Amplifies Early Proinflammatory Cytokine Production in the Lung during Respiratory Syncytial Virus Infection. J Virol 2014. 88: 6128–6136.

Goritzka, M., Pereira, C., Makris, S., Durant, L. R. and Johansson, C.: T cell responses are elicited against Respiratory Syncytial Virus in the absence of signalling through TLRs, RLRs and IL-1/IL-18R. Sci Rep 2015. 5: 18533.

Vanderheiden, A., Thomas, J., Soung, A. L., Davis-Gardner, M. E., Floyd, K., Jin, F., Cowan, D. A. et al.: CCR2 Signaling Restricts SARS-CoV-2 Infection. mbio 2021. 12: e02749-21.

Goritzka, M., Makris, S., Kausar, F., Durant, L. R., Pereira, C., Kumagai, Y., Culley, F. J. et al.: Alveolar macrophage-derived type I interferons orchestrate innate immunity to RSV through recruitment of antiviral monocytes. J Exp Med 2015. 212: 699–714.

Bastard, P., Gervais, A., Voyer, T. Le, Rovain, J., Philippot, Q., Manry, J., Michailidis, E. et al.: Autoantibodies neutralizing type I IFNs are present in ~4% of uninfected individuals over 70 years old and account for ~20% of COVID-19 deaths. Science Immunology 2021. 6: eabd4340.

Hyde, S. C., Pringle, I. A., Abdullah, S., Lawton, A. E., Davies, L. A., Varathalingam, A. et al.: CpG-free plasmids confer reduced inflammation and sustained pulmonary gene expression. Nat Biotechnol 2008. 26: 549–551.

Zanta-Boussif, M. A., Charrier, S., Pringle, M., Brice-Ouze, A., Martin, S., Opolon, P., Thrasher, A. J., Hope, T. J. et al.: Validation of a mutated PRE sequence allowing high and sustained transgene expression while abrogating WHV-X protein synthesis: Application to the gene therapy of WAS. Gene Ther 2009. 16: 605–619.

Brown, B. D., Venneri, M. A., Zingale, A., Sér, L. S. and Naldini, L.: Endogenous microRNA regulation suppresses transgene expression in hematopoietic lineages and enables stable gene transfer. Nat Med 2006. 12: 585–591.

Meyer-Berg, H., Zhou Yang, L., Pilar de Lucas, M., Zambrano, A., Hyde, S. C. and Gill, D. R.: Identification of AAV serotypes for lung gene therapy in human embryonic stem cell-derived lung organoids. Stem Cell Research and Therapy 2020. 11: 448.

Rihn, S. J., Merits, A., Bakshi, S., Turnbull, M. L., Wickenhagen, A., Alexander, A. J. T. et al.: A plasmid DNA-launched SARS-CoV-2 reverse genetics system and coronavirus toolkit for COVID-19 research. PLoS Biol 2021. 19: e3001091.

Kirsebom, F. C. M., Kausar, F., Nuriev, R., Makris, S. and Johansson, C.: Neutrophil recruitment and activation are differentially dependent on MyD88/TRIF and MAVS signaling during RSV infection. Mucosal Immunology 2019. 12: 1244–1255.

Full correspondence: Prof. Cecilia Johansson, St Mary's Campus, National Heart and Lung Institute, Imperial College London, W2 1PG, UK.
Email: c.johansson@imperial.ac.uk

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