Corticosteroid suppression of antiviral immunity increases bacterial loads and mucus production in COPD exacerbations

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Inhaled corticosteroids (ICS) have limited efficacy in reducing chronic obstructive pulmonary disease (COPD) exacerbations and increase pneumonia risk, through unknown mechanisms. Rhinoviruses precipitate most exacerbations and increase susceptibility to secondary bacterial infections. Here, we show that the ICS fluticasone propionate (FP) impairs innate and acquired antiviral immune responses leading to delayed virus clearance and previously unrecognised adverse effects of enhanced mucus, impaired antimicrobial peptide secretion and increased pulmonary bacterial load during virus-induced exacerbations. Exogenous interferon-β reverses these effects. FP suppression of interferon may occur through inhibition of TLR3- and RIG-I virus-sensing pathways. Mice deficient in the type I interferon-α/β receptor (IFNAR1−/−) have suppressed antimicrobial peptide and enhanced mucin responses to rhinovirus infection. This study identifies type I interferon as a central regulator of antibacterial immunity and mucus production. Suppression of interferon by ICS during virus-induced COPD exacerbations likely mediates pneumonia risk and raises suggestion that inhaled interferon-β therapy may protect.
cute exacerbations of airway diseases are a major cause of morbidity and mortality. Rhinoviruses (RVs) are the most frequent cause of exacerbations, and experimental challenge studies in asthma and chronic obstructive pulmonary disease (COPD) confirm a causal role between RV infection and exacerbation. The innate immune response to RV involves production of type I and III interferons (IFNs) by airway epithelial and immune cells. IFN responses to virus infection are impaired in asthma and COPD and are related to increased exacerbation severity.

Inhaled corticosteroids (ICS) are frequently prescribed therapies in airway diseases but only modestly reduce COPD exacerbation frequency. Clinical trials reported increased pneumonia risk with ICS use in COPD, although the mechanisms underlying this effect are unknown. More severe RV infection increases frequency of secondary bacterial infections in COPD, suggesting that the increased pneumonia risk may be a consequence of increased severity of initial virus infection.

Emerging evidence suggests that corticosteroids may impair innate antiviral immune responses, but studies have assessed the effects of ICS on host antiviral and antibacterial responses specifically in COPD, a disease associated with insensitivity to corticosteroids. The precise mechanisms, including the role of type I IFN, underlying adverse effects have not been elucidated.

Here, using models of rhinovirus infection and rhinovirus-induced COPD exacerbation, we demonstrate that the commonly used ICS fluticasone propionate (FP) impairs innate and acquired antiviral immune responses leading to delayed virus clearance, mucus hypersecretion and increased lung bacterial loads. Our studies highlight a central role for type I IFN in regulating anti-microbial immunity and mucus secretion during virus-induced exacerbations. The adverse effects associated with immune suppression by ICS provide a mechanism to explain the increased risk of pneumonia associated with their use in COPD.

Results

FP affects transcription factors regulating immunity to RV. To determine the efficacy and duration of action of ICS in a mouse model of FP treatment, we assessed glucocorticoid receptor (GR) activation by measuring nuclear GR-DNA binding. Administration of 20 μg FP, a dose similar to that employed in previous animal studies, significantly increased GR activation, while a tenfold lower dose did not (Fig. 1a). RV infection activates immune-regulating transcription factors, including nuclear factor kappa-B (NFκB), required for proinflammatory responses to RV infection and IFN regulatory factor-3 (IRF-3), required for antiviral responses. We assessed the effect of FP administration on RV activation of these transcription factors in mouse lung. FP suppressed lung NFκB p65 activation in RV-infected mice at 8 h post infection (Fig. 1c). For IRF-3, activation by RV occurred at 2 h post infection and significant suppression was also observed in FP-treated mice (Fig. 1c).

ICS suppress innate immunity and impair virus control. We next assessed the effect of FP on innate antiviral immune responses to RV infection in mice. FP suppressed RV induction of lung IFNβ and IFNL2/3 mRNAs (~75% and ~60% reductions, respectively, Fig. 1d) and IFN-α, IFN-β and IFN-λ2/3 BAL proteins (~85%, ~80% and ~50% reductions, respectively, Fig. 1e). FP also suppressed RV induction of the interferon-stimulated genes (ISGs) 2′-5′-oligoadenylate synthetase (2′-5′-OAS), protein kinase-R (PKR) and viperin and CXCL10/IP-10 protein in BAL (~75%, ~45%, ~60% and ~80% reductions, respectively, Fig. 1f).

Type I IFN receptor signalling is required for airway natural killer (NK) cell responses to RVs. Reduced numbers of total and activated (CD69 expressing) NK cells were observed in RV-infected mice treated with FP (Fig. 1g). Consistent with the suppression of innate antiviral responses, FP treatment led to delayed virus clearance with increased RV RNA copies in FP-treated mice at 8, 24 and 48 h post infection (maximal effect size threefold at 8 and 48 h) and increased levels of infectious virus at 24 h (Fig. 1h).

To confirm that other ICS compounds apart from FP can also impair antiviral immunity, we evaluated the effect of another commonly used agent, budesonide, on RV infection in mice. Administration of 20 μg of budesonide similarly suppressed RV induction of lung IFNβ and IFNL2/3 and increased RV RNA (Supplementary Fig. 1a-d).

FP impairs adaptive immune responses to RV infection. We next determined the effects of FP on adaptive immune responses to RV infection (Fig. 2a). Total and activated numbers of both CD4+ and CD8+ T cells were suppressed in FP-treated mice (Fig. 2b, c) as were serum levels of RV-specific IgG1 and IgG2a antibodies and neutralising antibody (Fig. 2d).

FP suppresses virus-induced airway inflammation. FP administration prior to RV infection (Supplementary Fig. 2a) suppressed cellular airway inflammation with reduced numbers of neutrophils, lymphocytes and macrophages observed in BAL (Supplementary Fig. 2b). FP also suppressed RV induction of BAL neutrophil chemokines CXCL1/KC and CXCL2/MIP-2 and the lymphocyte chemokine CCL5/RANTES (Supplementary Fig. 2c).

FP enhances mucin production in RV-infected mice. We evaluated the effect of FP on BAL fluid concentrations of the mucins MUC5AC and MUC5B, the major glycoprotein constituents of airway mucus. RV infection alone increased airway production of both mucin glycoproteins and both were significantly further increased by FP treatment (~85% for MUC5AC and ~80% for MUC5B, Fig. 2e).

FP impairs antibacterial responses in RV-infected mice. To further evaluate adverse effects associated with FP administration during RV infection, we assessed lung bacterial loads by measurement of bacterial 16S rRNA. FP increased bacterial loads (~2.5-fold) in RV-infected mice (Fig. 2f). FP also suppressed RV induction of secretory leukoprotease inhibitor (SLPI) protein (~50% reduction, Fig. 2f), an antimicrobial peptide (AMP) implicated in protection against secondary bacterial infection following RV infection in COPD. FP administration also suppressed RV induction of proinflammatory cytokines IL-6 and TNF (Supplementary Fig. 2d) and the AMP pentraxin-3, but had no effect on RV induction of other AMPs α-defensin 1, β-defensin 2 or mannose-binding lectin-2 (Supplementary Fig. 2e). The AMPs surfactant protein-D and lysozyme were not significantly induced by RV infection nor suppressed by FP in mice (Supplementary Fig. 2f).

FP suppresses TLR3- and RIG-I-mediated IFN responses. To investigate mechanisms whereby FP suppresses innate and acquired immune responses, we next assessed which pattern recognition receptor (PRR) and IFN signalling pathways were inhibited by FP in human BEAS-2B bronchial epithelial cells. As observed in mice in vivo, FP treatment of BEAS-2B cells in vitro suppressed RV induction of IFNβ and IFNL2/3 (Fig. 3a). Double-stranded RNA intermediates form during RV replication and
Fluticasone propionate suppresses innate antiviral immune responses and delays virus clearance in mice. C57BL/6 mice were treated intranasally with fluticasone propionate (20 or 2 μg) or vehicle DMSO control. Glucocorticoid receptor activation in lung tissue was assessed by measuring nuclear DNA binding by ELISA. d IFN-α, IFN-β and IFN-λ2/3 proteins in bronchoalveolar lavage (BAL) at 24 h post infection were measured by ELISA. e Interferon-stimulated genes Z-5′ OAS, PKR and Viperin mRNAs in lung tissue at 8 h post infection were measured by quantitative PCR. f BAL cells were stained for CD3 and the NK cell marker NK1.1 and analysed by flow cytometry. BAL CD3− NK1.1− cell numbers and BAL CD3+ NK1.1+ cell numbers were enumerated at day 2 post infection. h Rhinovirus RNA copies in lung tissue were measured by Taqman quantitative PCR and infectious virus in lung tissue was measured by titration in HeLa cells. Data represents mean (±SEM) of five–eight mice per treatment group, representative of at least two independent experiments. Data were analysed by one- or two-way ANOVA with Bonferroni post test. ns non-significant. *p < 0.05, **p < 0.01, ***p < 0.001
activate the PRRs TLR3, RIG-I and MDA-5.21,23. FP treatment markedly suppressed induction of IFNβ and IFNλ2/3 by TLR3 (Fig. 3b) and RIG-I agonists (Fig. 3c) but had no effect on MDA-5 agonist (Fig. 3d).

TLR3 induces IFN expression via the adaptor molecule TIR-domain-containing adaptor inducing IFN-β (TRIF), while MDA-5 and RIG-I signal via mitochondrial antiviral signalling protein (MAVS)22,24. We therefore transfection BEAS-2B cells with IFNβ promoter–reporter constructs and stimulated using plasmids encoding constitutively active TRIF (ΔTRIF) or MAVS (ΔMAVS). Transfection with ΔTRIF or ΔMAVS significantly increased IFNβ promoter activity (Fig. 3e, f). Consistent with our finding that FP suppressed TLR3- and RIG-I-mediated IFN induction, treatment with FP suppressed both ΔTRIF- and ΔMAVS-induced IFNβ promoter activity (Fig. 3e, f).

Secondary amplification of virus-induced IFN release involves type I IFN acting in an autocrine manner via the IFN alpha/beta receptor (IFNAR) to stimulate further induction of IFNs and ISGs.20 To assess effects of FP on IFN signalling, we stimulated BEAS-2B cells with recombinant IFN-β and found no effect of FP on IFN-β induction of ISGs 2′−5′-OAS, viperin and myxovirus resistance A (Mx1) (Fig. 3g), thereby demonstrating that FP suppression of IFN is restricted to effects on initial virus-sensing pathways and not secondary amplification via IFNAR. IFN signalling via IFNAR induces phosphorylation of STAT1/2. Therefore, to further confirm this finding, immunodetection of STAT1 phosphorylated at Y701 (pY701) and STAT2 phosphorylated at Y690 (pY690) was investigated in BEAS-2B cells stimulated with recombinant IFN-β. FP had no effect on IFN-β induction of either STAT1 or STAT2 phosphorylation (Fig. 3h).

We additionally confirmed these in vitro findings by demonstrating that FP administration had no effect on lung ISG induction following recombinant IFN-β administration in mice (Fig. 3i).

IFNβ reduces virus load and reverses mucin augmentation by FP. To demonstrate that suppression of IFN by FP is functionally related to increased virus load and mucin production, we administered recombinant IFN-β in combination with FP treatment and RV infection (Fig. 4a). For clarity, selected experimental groups are in Fig. 4 and the full data set in Supplementary Fig. 3. IFN-β reversed FP-mediated suppression of lung 2′−5′-OAS and

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**Fig. 2** Fluticasone propionate suppresses acquired responses and augments mucin production and bacterial loads in mice. a C57BL/6 mice were treated intranasally with fluticasone propionate (20 μg) or vehicle DMSO control and challenged intranasally with rhinovirus (RV)-A1 or UV-inactivated RV-A1. b, c BAL cells were stained with antibodies specific for CD3, CD4, CD8 and CD69 and analysed by flow cytometry. b BAL CD3+ CD4+ T cell numbers and BAL CD3+ CD4+ CD69+ T cell numbers at day 7 post infection. c BAL CD3+ CD8+ T cell numbers and BAL CD3+ CD8+ CD69+ T cell numbers at day 2 post infection were evaluated. d Peripheral blood was harvested at day 14 post infection. RV-specific IgG1 and RV-specific IgG2A were quantified in serum by ELISA. Sera were assayed for their ability to prevent cytopathic effect caused by the same RV serotype used for in vivo challenge. Cytopathic effect was quantified by crystal violet staining. Top dotted line: serum only, (uninfected) controls. Bottom dotted line: virus infected (no serum) control. e MUC5AC and MUC5B proteins in BAL at day 7 post infection were measured by ELISA. f Bacterial copy number of 16S rRNA in lung tissue at 96 h was measured by quantitative PCR. Secretory leucocyte protease inhibitor (SLPI) protein in BAL was measured by ELISA. In all figures except d (right), data represents mean (±SEM) of five–eight mice per group, representative of at least two independent experiments. Data were analysed by one- or two-way ANOVA with Bonferroni post test. ns non-significant; *p < 0.05, **p < 0.01, ***p < 0.001. In (d) (right), data points represent sera pooled from five mice per group, representative of two independent experiments.
viperin expression (Fig. 4b) and of BAL CXCL10/IP-10 and IFN-α2/3 protein concentrations (Fig. 4c). Compared to FP-treated RV-infected mice, administration of IFN-β following FP treatment and infection significantly reduced virus loads to levels observed in untreated virus-infected mice (Fig. 4d). IFN-β treatment additionally prevented FP upregulation of BAL MUC5AC glycoprotein levels (Fig. 4e).

IFNβ reverses FP impairment of antibacterial immunity. Secondary bacterial infection in RV-induced COPD exacerbations is mediated by virus-induced neutrophil elastase-mediated degradation of AMPs, including SLPI14. We therefore investigated whether IFN-β could prevent this occurring. IFN-β treatment further enhanced FP suppression of BAL neutrophil numbers and neutrophil elastase levels (Fig. 4f), reversed FP suppression of BAL SLPI levels and prevented FP augmentation of bacterial load (Fig. 4g) in RV-infected mice. However, IFN-β treatment had no effect on FP suppression of antibacterial mediators IL-6, TNF or pentaxin-3 (Supplementary Fig. 4a–c).

**IFNAR1**−/− mice have altered mucin and SLPI responses to RV. To confirm suppression of IFN is functionally related to

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**Fig. 3** Fluticasone propionate suppresses TLR3 and RIG-I-induced interferon responses but does not block type I IFN signalling. a–d BEAS-2B cells were treated with fluticasone propionate at 1 and 10 nM concentrations and stimulated with rhinovirus-A1 or receptor-specific agonists. Cell lysates were collected at 24 h post stimulation. a IFNβ and IFNα2/3 mRNA expression following RV-A1 stimulation was measured by quantitative PCR. b IFNβ and IFNα2/3 mRNA expression following RIG-I agonist (transfected 5′ppp-dsRNA) stimulation was measured by quantitative PCR. Transfected dsRNA lacking the 5′ triphosphate was used as a ‘RIG-I control’. d IFNβ and IFNα2/3 mRNA expression following MDA-5 agonist (transfected HMW Poly(I:C) directly pre-coupled to the transfection reagent LyoVec) stimulation was measured by quantitative PCR. LyoVec without Poly(I:C) (LyoVec control) was used as a control for transfection. e, f BEAS-2B cells were transfected with interferon-β promoter–reporter constructs together with ΔTRIF, ΔMAVS or vector control (pUNO) and then treated with fluticasone propionate (FP) 1 or 10 nM or medium control 3 h later. Cells were collected 24 h later and relative light units (RLU) were determined. e IFNβ promoter activity following transfection with ΔTRIF. f IFNβ promoter activity following transfection with ΔMAVS. g, h BEAS-2B cells were treated with fluticasone propionate at 1 and 10 nM concentrations and stimulated with recombinant IFN-β. g Z2′-S′ OAS, viperin and Mxi1 mRNA expression at 24 h post stimulation was measured by quantitative PCR. h Cell extracts were collected at 1 h post stimulation and analysed by western blotting with antibodies to pSTAT1 Y701, STAT1, pSTAT1 Y690 and STAT2. a–g data represent mean (±SEM) comprising three independent experiments, analysed by one-way ANOVA with Bonferroni post test. ns non-significant. *p < 0.05, **p < 0.01, ***p < 0.001. In h, data shown are representative of results from three independent experiments. i C57BL/6 mice were treated intranasally with fluticasone propionate (20 μg) or vehicle DMSO control and additionally with recombinant 104 units IFN-β. Interferon-stimulated genes Z2′-S′ OAS, and Viperin mRNAs in lung tissue at 8 h post infection were measured by quantitative PCR. Data represents mean (±SEM) of five mice per treatment group, representative of at least two independent experiments. Data were analysed by one-way ANOVA with Bonferroni post test. ns non-significant. *p < 0.05, **p < 0.01, ***p < 0.001
enhanced mucin and impaired antibacterial immunity, we evaluated MUC5AC and SLPI expression in RV-infected mice deficient in IFNAR signalling (IFNAR1−/−) (Fig. 4h). BAL MUC5AC protein concentrations in RV-infected IFNAR1−/− mice were increased (Fig. 4i). To confirm that IFNAR signalling is also important for SLPI production during RV infection, we measured BAL SLPI protein in RV-infected IFNAR1−/− mice and observed reduced concentrations (Fig. 4j).

IFNβ does not restore acquired responses nor reduce MUC5B. We observed no effect of IFN-β treatment on FP suppression of adaptive immune responses with no difference in BAL CD4+ or CD8+ T cell numbers or serum-neutralizing antibody levels in RV-infected mice treated with FP and IFN-β versus FP alone (Supplementary Fig. 4d). In contrast to the effect observed on MUC5AC (Fig. 4e), recombinant IFN-β also had no effect on FP enhancement of BAL MUC5B protein levels (Supplementary Fig. 4e).

FP suppresses IFN and enhances mucus secretion in COPD cells. We next investigated whether similar adverse effects of FP occurred in cells from COPD patients. We evaluated the effects of FP in airway epithelial cells (AECs) sampled bronchoscopically from nine patients with GOLD stage 3 disease, cultured at an air–liquid interface. The quality of cultures was confirmed by the presence of ciliated epithelium, mucus-producing cells, pseudostratified structure and assessment of trans-epithelial electrical resistance (Supplementary Fig. 5a). Demographic/clinical characteristics of the patients are in Table 1. FP significantly suppressed early RV induction of IFNB and IFNA1 and IFNA2/3 (Fig. 5a–c and Supplementary Fig. 6a). FP also suppressed RV induction of 2′-5′-OAS, PKR and viperin (Fig. 5d–f) and additionally suppressed IFN-β, IFN-Α1/3 and CXCL10/IP-10 proteins in supernatants of COPD AECs (Fig. 5g–i).

In keeping with our finding that FP administration augmented mucin production in response to RV infection in mice, we also observed an increase in mucus staining associated with FP treatment in RV-infected human COPD AECs in n = 8 subjects for whom sample was available for histological analysis (Fig. 5j).

FP impairs IFN responses in a COPD exacerbation mouse model. We investigated the effects of FP on RV-induced exacerbation of COPD-like disease in vivo in a mouse model of RV-exacerbated elastase-induced emphysema (Fig. 6a) in which many of the features of human virus-induced COPD exacerbations are recapitulated20. FP suppressed RV induction of IFN-β...
and IFN-λ2/3 proteins in BAL of elastase-treated mice (Fig. 6b) and increased virus load at days 1 and 4 post infection (Fig. 6c).

FP suppressed RV induction of BAL total cells and lymphocytes but had no significant effect on neutrophils (Fig. 6d). FP also suppressed RV induction of the lymphocyte chemokines CCL5/RANTES and CXCL10/IP-10 in BAL (Fig. 6e) and additionally suppressed RV induction of IL-6, TNF and SLPI (Fig. 6f, g). FP also enhanced RV induction of BAL MUC5AC protein (Fig. 6h) and mucus staining in lung sections (Fig. 6i), but there was no significant effect on BAL MUC5B protein levels (Fig. 6h).
Adverse effects of ICS use during human COPD exacerbation.

To confirm the clinical relevance of our findings, we measured antiviral responses in sputum samples from a cohort of 40 patients with COPD, in which 18 episodes of virus-induced COPD exacerbations occurred (rhinovirus \( RRV = 11 \), coronavirus \( RRV = 4 \), RSV \( RRV = 2 \), influenza \( RRV = 1 \)). Samples were assessed in 36 patients who had sufficient sample for evaluation at clinical stability (baseline) and in exacerbating patients at exacerbation onset and 2 weeks during exacerbation (Fig. 7a). Patients were stratified according to current use \(( n = 7 \) ) or non-use \(( n = 11 \) ) of ICS, baseline characteristics of these subgroups are in Table 2. ICS users had suppressed baseline sputum cell \( IFN \beta \) expression versus ICS non-users with no significant differences observed for \( IFN \lambda 1 \) and \( IFN \lambda 2/3 \) (Supplementary Fig. 7a).

In keeping with prior findings in vitro and in animal models, ICS users had suppressed sputum cell \( IFN \beta \) and \( IFN \lambda 2/3 \) expression at exacerbation onset (Fig. 7b). Evaluation of mRNA expression as fold-change from baseline also demonstrated significant suppression in ICS users for \( IFN \lambda 2/3 \) (Supplementary Fig. 7b). As current smoking can affect antiviral responses, we also carried out a sub-analysis excluding current smokers. In this subgroup, ICS users had significantly suppressed sputum cell \( IFN \lambda 2/3 \) (Supplementary Fig. 7c).

Sputum cell expression of \( 2'5'-OAS \) and \( viperin \) at 2 weeks was also significantly suppressed in ICS users versus non-users with no difference observed for \( Mx1 \) (Fig. 7c). ICS users also had suppressed sputum supernatant \( CXCL10/IP-10 \) protein concentrations at exacerbation onset (Fig. 7d) and enhanced sputum \( MUC5AC \) glycoprotein concentrations at 2 weeks (Fig. 7e).

Secondary bacterial infection occurs following experimental RV infection in COPD subjects who were not taking ICS, with peak bacterial load occurring at day 15 post RV infection.
determine if ICS use was associated with increased secondary bacterial infection in COPD, we evaluated bacterial load by 16S qPCR at the similar time point of 2 weeks post-exacerbation onset and found a significant increase in ICS users versus non-users (Fig. 7f). In keeping with effects of FP in mice, ICS users also had suppressed sputum cell SLPI expression at exacerbation (Fig. 7f).

ICS use during virus-induced COPD exacerbation was also associated with a greater maximum percentage decline from baseline in post-bronchodilator peak expiratory flow in ICS users versus ICS non-users (Fig. 7g).

**Correlation of IFNs with MUC5AC or SLPI during exacerbation.** Consistent with our findings that type I IFN negatively regulates mucin production during RV infection in mice, we observed that peak sputum cell expression of IFNβ, IFNα2/3 and peak sputum concentrations of CXCL10/IP-10 protein negatively correlated with sputum MUC5AC during virus-induced COPD exacerbation (Fig. 7h). Similarly, in keeping with our findings in mice that interferon responses are also involved in SLPI production during virus infection, we found that peak sputum cell expression of IFNβ and viperin positively correlate with SLPI expression (Fig. 7i).

**Discussion**

Using models of RV infection we have shown that FP, a commonly used ICS in man, suppresses critical antiviral immune responses, leading to impaired lung virus control and previously unrecognised adverse effects of mucin enhancement and impairment of antibacterial immunity, effects that were ameliorated by recombinant IFN-β administration and thus related to...
IFN suppression associated with FP administration. We importantly show that these adverse effects of FP also occur during COPD exacerbations, thus confirming relevance and importance in the context of human disease.

In primary RV infection models, FP suppressed induction of type I and type III IFNs by RV and delayed virus clearance in vivo. These data extend observations from previous studies which reported suppressed IFN responses to virus infection associated with corticosteroid administration in peripheral blood mononuclear cells, fibroblasts and airway epithelial cells. An intact type I IFN response is important for effective lung virus control and we have previously reported that IFN-β, thereby suggesting that inhibitory effects of FP may be confined to initial virus-sensing pathways and not through effects on autocrine amplification via IFNAR. We have previously shown that induction of IFN by RV infection occurs initially via the endosomal PRR TLR3 and later via the inducible cytoplasmic helicases RIG-I and MDA-5. In the current study, FP suppressed TLR3-mediated IFN expression, suggesting that corticosteroid inhibition of IFN production occurs through an early effect on the initial endosomal sensing of RVs. In keeping with this finding, we observed that FP suppressed IFNβ promoter activity induced by TRIF, the adaptor molecule utilised by TLR3 to stimulate type I IFN expression. We additionally evaluated the effect of FP administration on stimulation of airway epithelial cells by agonists for the cytosolic PRRs downstream of TLR3 and surprisingly found a differential effect with suppression of RIG-I-induced, but no effect on MDA-5-induced, IFN expression. Additionally, FP suppressed IFNβ promoter activity induced by MAVS, the adapter molecule that is utilised by both MDA-5 and RIG-I to stimulate IFN expression. The reason for the discrepancy between the effect of FP on pathways mediated by the cytoplasmic receptors RIG-I and MDA-5 (which share many similarities) is unclear. It is possible that differences exist between how the adaptor molecule MAVS is utilised by both MDA-5 and RIG-I to stimulate IFN expression. Nonetheless, our data suggest that FP may exert its inhibitory effects on IFN induction through inhibition of the TLR3 virus-sensing pathway. This is consistent with a study which reported that FP suppressed poly(I-C) induction of CXCL10/IP-10 in BEAS-2B cells. However, it should be noted that, although the selectivity of agonists used is supported by previous studies, we cannot exclude that ligands employed in our studies do not trigger additional PRRs to those stated. Further evidence that IFN affect IFN signalling was shown in our mouse model, where FP administration suppressed activation of lung IFR-3, a signalling intermediate of the innate response. However, it should be noted that any nuclei obtained using our methods will be from multiple cell types, so it is possible that these data may be explained by effects of ICS on cellular composition.

Adaptive immune responses are important for both virus clearance and prevention of reinfecion and we observed that FP administration prior to RV infection in mice led to suppressed

| Table 2 Characteristics of patients with COPD presenting with virus-positive exacerbations, stratified according to current ICS use |
|-----------------------------------------------|
|                                | ICS users (n = 7) | ICS non-users (n = 11) | p Value |
| Age (median (IQR))     | 68 (63–74)       | 65 (45–71)             | 0.71    |
| Male                   | 4 (57.1%)        | 7 (63.6%)              | 1.0     |
| GOLD stage (median (IQR)) | 2 (1.5–2)       | 2 (2–2.5)              | 0.46    |
| Current smoker         | 3 (42.8%)        | 2 (18.2%)              | 0.33    |
| Pack-year history (median (IQR)) | 40 (35–47.5) | 50 (47.5–53)           | 0.12    |
| Body mass index (median (IQR)) | 27.4 (26.1–28.4) | 28.0 (23.1–32.9) | 1.0     |
| Ischaemic heart disease | 0 (0%)           | 1 (9.1%)               | 1.0     |
| Diabetes mellitus      | 1 (14.3%)        | 1 (9.1%)               | 1.0     |
| Cerebrovascular disease | 1 (14.3%)        | 0 (0%)                 | 0.39    |
| Prior annual exacerbation frequency (median (IQR)) | 3 (0.5–3.5) | 2 (1–2)                | 0.67    |
| Prior influenza vaccination | 6 (85.7%)     | 8 (72.7%)              | 1.0     |
| Beta agonist inhaler use | 5 (71.4%)       | 8 (72.7%)              | 1.0     |
| Anti-muscarinic inhaler use | 4 (57.1%)     | 7 (63.6%)              | 1.0     |
| Prednisolone initiated at exacerbation | 2 (28.6%) | 1 (9.1%)               | 0.53    |
| Antibiotics initiated at exacerbation  | 4 (57.1%)       | 4 (36.4%)              | 0.63    |

Categorical variables were compared using the χ² test. Continuous variables were compared using the Mann–Whitney U-test.
cellular adaptive responses and impaired production of RV-specific IgG and neutralising antibodies. Our finding that FP impairs production of antibody responses to RV infection in the mouse model suggests that use of these commonly prescribed inhalers during COPD exacerbations could impair natural protective responses against future RV infections and thus limit long-term efficacy of ICS in reduction of exacerbations. This is supported by a previous study which reported that patients with exacerbation-prone COPD have lower RV-specific IgG1 antibody levels during stable state than non-exacerbating patients, suggesting that deficient humoral immune responses to RV might be an important determinant of exacerbation risk.

RV induction of the major airway mucins MUC5AC and MUC5B was also enhanced by FP administration in the mouse model. Impaired mucociliary clearance with mucus hypersecretion is a feature of stable disease in COPD and likely contributes to airflow limitation, accelerated lung function decline and increased morbidity. Additionally, lung MUC5AC over-expression has been reported in cases of fatal asthma and is thus associated with adverse outcomes. RV infections can stimulate mucus production from airway epithelium and thereby potentiate sputum production during acute exacerbations, and our finding that FP administration diminishes control of RV infection leading to increased mucin expression further suggests that use of these therapies may worsen mucus hypersecretion, leading to enhanced and/or prolonged symptoms and delayed recovery from exacerbation.

Administration of recombinant IFN-β in combination with FP and RV infection in mice reconstituted suppressed innate responses and improved virus clearance, suggesting that FP suppression of IFN responses plays a major mechanistic role in the adverse effects on virus control. RV-infected mice given steroid and then treated with IFN-β also had reduced MUC5AC expression compared with FP-treated infected mice, which further supports the hypothesis that suppression of IFN underlies the enhanced MUC5AC production associated with FP administration. We additionally observed that IFNAR1−/− mice which have diminished production of type I IFN in response to RV infection, also have increased BAL MUC5AC, providing further evidence that type I IFN negatively regulates mucin production. Similar observations have been previously reported in IFNAR1−/− mice infected with Cryptococcus neoformans.

The association between ICS and pneumonia risk in COPD was first reported in 2007 and has now been confirmed in numerous studies. Based on this perceived risk, some clinicians are now questioning the uncritical use of these widely prescribed therapies in COPD with ongoing evidence of practice change. Remarkably, an underlying mechanism for this effect has never been identified. We have previously reported that experimental RV challenge induces secondary bacterial infection in COPD, through virus-induced suppression of the AMPs SLPI and elafin. In the current studies, in addition to suppression of IFN, FP also suppressed RV induction of a number of airway antibacterial responses, including SLPI, accompanied by increases in bacterial load. Recombinant IFN-β administration reversed suppression of SLPI and reduced increased bacterial loads associated with FP administration without affecting suppression of other antibacterial responses. Combined, these data suggest that FP may potentially increase risk of virus-induced secondary bacterial pneumonia through suppression of SLPI and that recombinant IFN therapy may prevent increased bacterial loads following RV infection. We speculate that this could be one mechanism explaining the increased pneumonia risk associated with ICS use in COPD, and is consistent with previous data showing that higher RV loads are associated with increased risk of secondary bacterial infection in COPD and that ICS-related pneumonia episodes typically follow protracted symptomatic exacerbations.

Some studies have previously suggested that corticosteroids may enhance SLPI production by bronchial epithelium in vitro, but these studies were carried out in models of stable rather than exacerbated disease and conflicting data exists on the effects of corticosteroids in other cell types, such as THP-1 monocytes, where suppression rather than enhancement has been observed. During virus infection, we clearly demonstrate in a range of models that FP has a suppressive effect on airway SLPI production. Since STAT1 has previously been shown to regulate SLPI expression, FP likely impairs SLPI production during RV infection by suppressing PRR-induced IFN production thereby reducing subsequent IFNAR-mediated STAT1 activation. This mechanism is supported by our finding that IFNAR1−/− mice also had reduced RV induction of SLPI. Additional evidence for the importance of deficient SLPI being an important mechanism for bacterial infection in COPD is provided by studies that have shown reduced airway SLPI in frequent, notably, of whom a greater proportion was taking ICS) versus infrequent COPD exacerbators and in patients with positive bacterial culture at exacerbation.

Collectively, our studies using recombinant IFN administration in mice and RV infection in the IFNAR1−/− mouse strain identify IFN as a central regulator of antibacterial responses and mucus production during RV infection. In a clinical trial in asthma patients taking ICS who developed a cold, IFN-β therapy resulted in a trend towards reduced virus loads, with clinical benefit in patients with moderate/severe disease. Our finding that IFN-β administration in the mouse model improved virus control, augmented antibacterial immunity and reversed the enhanced MUC5AC expression associated with FP treatment, adds further weight to the potential benefit of inhaled interferon as a treatment for virus-induced exacerbations of airborne disease. Thomas et al. reported that administration of IFN-α and IFN-λ may similarly ameliorate ICS impairment of lung influenza virus control in mice, although inhaled IFN-β is the therapy that is in clinical development and therefore of greatest current interest. Inhaled IFN-β has previously been shown to be safe in patients with asthma, and dedicated studies are required to investigate its utility in COPD.

It should be noted that the overall exacerbation reduction associated with ICS use in COPD outweighs the potential risk of pneumonia. Our data showing that FP suppresses virus-induced airway inflammation supports the beneficial effects associated with use of these therapies. It is plausible that the adverse effects of ICS on host-defence reported in the current study might occur more frequently in the context of more severe infection, leading to unfavourable skewing of the risk/benefit ratio. Future development of selective steroids that retain potentially beneficial anti-inflammatory effects without having adverse effects on antiviral immunity could provide more effective approaches.

We additionally confirmed that the adverse effects of FP identified in our primary infection models also occur in models of COPD exacerbation. Evaluation of whether these detrimental effects occur in COPD is important because COPD is reported to be a relatively steroid-resistant disease and therefore it is feasible that steroid-induced adverse effects may not be observed in this context. In primary COPD AECS, FP suppressed RV induction of IFN responses and also enhanced mucus production. We confirmed these findings in vivo using a mouse model of COPD exacerbation, to similarly demonstrate that FP suppresses RV-induced IFN responses leading to delayed virus clearance and mucus hypersecretion in vivo. Thus, FP administration in both in vitro and in vivo COPD models was associated...
with suppressed IFN responses, confirming that these adverse effects still occur in steroid-resistant disease models, and the increased virus load and mucus production observed in vivo could impact adversely on exacerbation severity. The elastase mouse model of COPD reproduces many of the immunopathological features of human COPD exacerbation, but it is well recognised that existing models cannot completely recapitulate the complexities of human COPD, which occurs in genetically susceptible individuals following prolonged cigarette smoke exposure. Elastase administration represents a synthetic model of COPD and further animal studies using alternative modelling approaches, such as smoke exposure or genetic manipulation, are required to confirm our findings.

We confirmed the direct relevance of our findings to use of ICS in a clinical setting through analysis of sputum samples from patients with COPD presenting with virus-associated exacerbations. ICS use was associated with suppressed IFN responses at exacerbation with a concomitant increase in MUC5AC levels. In keeping with our findings in the mouse RV infection model and in support of our hypothesis that ICS-mediated suppression of SLPI promotes increased pneumonia risk in COPD, we also observed suppressed sputum SLPI mRNA and increased bacterial loads in ICS users. These data suggest that our prior in vitro and in vivo mouse model findings could be applicable in the context of human COPD exacerbations and provide preliminary clinical evidence that use of ICS may have adverse effects when used in exacerbating patients. Further studies using larger numbers of COPD subjects will be needed to confirm these data. A negative correlation between interferon responses and MUC5AC concentrations and positive correlation between interferon responses and SLPI expression was also observed, further supporting our conclusion that type I IFN plays a central role in regulation of mucus secretion and antimicrobial immunity during exacerbations.

In summary, these studies demonstrate that inhaled corticosteroid use suppresses antiviral immune responses, accompanied by adverse effects on mucin production and antibacterial immunity during virus-induced COPD exacerbations. The role of recombinant interferon as an adjunctive therapy to negate the adverse effects of ICS in COPD warrants further exploration. A schematic representation showing the proposed adverse consequences of inhaled corticosteroid use during exacerbations and potential beneficial effects of interferon therapy are shown in Fig. 8. Future development of novel selective steroids that retain beneficial anti-inflammatory effects but lack adverse effects on antimicrobial immunity and mucin production could be more effective for treatment/prevention of COPD exacerbations and thus be of greater benefit in clinical practice.

**Methods**

**Cells and viruses.** BEAS-2B cells (European Collection of cell cultures) were cultured in RPMI 1640 medium with 10% foetal calf serum (FCS). Rhinovirus serotype A1 was propagated in Ohio HeLa cells (European Collection of cell cultures) by standard methods and inactivated by exposure to UV light at 1200 mJ/cm² for 30 min. For in vivo use in mouse models, virus was purified using precipitation in 7% polyethylene glycol 6000 (Sigma-Aldrich), 0.5 M NaCl, followed by filtration using a 0.2 μm syringe filter and buffer exchange and concentration in an Amicon Ultra-15 centrifugal filter unit (100 KDa nominal molecular weight limit) (Millipore, USA)33,59.

**Virus infection and treatment with FP and receptor agonists.** BEAS-2B cells were cultured to 80% confluence in 12-well plates. Cells were treated with 1 or 10 nM fluticasone propionate (Sigma-Aldrich) or diluent alone (mock) for 1 h. Cultures were subsequently stimulated with 0.2 ml of RV-A1 (MOI 2) or 5 μg/ml TLR3 agonist (poly(I:C); Invitrogen), 250 ng/ml MDA-5 agonist (transfected poly(I:C); high MW form which is specific for MDA-5, as previously described59, Invivogen).
membrane was carefully cut from the inserts and collected into RLT buffer. The cultures and stored for protein expression analyses and half of the transwell samples were collected at 24, 48, 72 and 96 h post infection. At each time point, apical media was removed from the transwell inserts and replaced with BEBM minimal at 35 °C, after which, infection media was replaced with 500 μl of PBS on ice. Cells were then incubated at 37 °C for 3 h with 5% CO2. Three hours after transfections, cells were treated with FP at 1 and 10 nM concentrations, as described above and lysates were collected 24 h post transfection. Relative light units were assessed using a Dual Luciferase kit (Promega).

Treatment and infection of primary bronchial epithelial cells. Primary bronchial epithelial cells were obtained from patients with GOLD stage 3 COPD bronchoscopically. All subjects gave informed written consent and the study protocol was approved by the Hunter New England Human Research Ethics Committee (05/08/10/03). Primary cells were grown in complete BEGM (Lonza) with growth factor supplements in submerged monolayer culture and then seeded at 2 × 10^5 cells/cm² in a 12-well plate (Corning) with ALL-IN-1 medium comprised of 50% BEBM/50% DMEM containing 0.1% hydrocortisone, 0.1% bovine insulin, 0.1% epinephrine, 0.1% transferrin, 0.4% bovine pituitary extract (all from Lonza) and ethanolamine (final concentration 80 μM), MgSO4 (final concentration 0.3 mM), MgCl2 (final concentration 0.4 mM), bovine serum albumin (final concentration 0.5 mg/ml), L-glutamine (final concentration 250 μM), transferrin (final concentration 30 mg/ml) and 2% penicillin/streptomycin in a humidified 5% CO2, 35 °C incubator. Cells were grown to 80% confluence before the transwell experiments were performed. The same day after the transwell setup, cells were infected with 0.5–1 μg/ml of rh-EGF with a final concentration of 250 μg/ml, all-trans retinoic acid (30 ng/ml) and 2% penicillin/streptomycin with 10 ng/ml of rh-EGF and incubated for 24 h for histological cross-sections to confirm differentiation and staining with periodic acid-Schiff (PAS) to assess mucus production.

Rhinovirus infection and treatment of mice. In vivo protocols: Female mice (8–10 weeks of age) on a C57BL/6 background were used for all animal studies. Mice were purchased from Charles River Laboratories and housed in individually ventilated cages under specific pathogen-free conditions. All animal work was performed under the authority of the UK Home Office outlined in the Animals (Scientific Procedures) Act 1986 after ethical review by Imperial College London Animal Welfare and Ethical Review Body (project licence PPL 70/7234).

FP powder (Sigma-Aldrich) was resuspended at a concentration of 357 μg/ml in dimethyl sulfoxide (DMSO; Sigma-Aldrich) and then diluted 1:1000 in PBS. Mice were treated intranasally under light isoflurane anesthesia with 50 μl of FP solution (equal to 20 μg FP dose) or vehicle DMSO diluted 1:1000 in PBS as control. One hour after FP or vehicle administration, mice were infected intranasally with 50 μl of FP solution (5 × 10^7 TCID50) or UV-inactivated RV-A1 control (as determined by plaque assay experiments with LTR priming). For the FP challenge, mice were additionally treated intranasally with 50 μl of PBS containing 10^5 units of recombinant IFN-β (R&D systems).

Flucitasone administration in COPD exacerbation mouse model. Mice were treated intranasally under light isoflurane anesthesia with 1.2 units of porcine pancreatic elastase (Merck) to induce emphysematous lung changes. Ten days following elastase treatment, mice were treated intranasally with the same volume PBS or Flucitasone solution (prepared as detailed above), 1 h prior to intranasal infection with 50 μl of RV-A1 (2.5 × 10^5 TCID50, a 50% dose reduction from the primary RV infection model) or UV-inactivated RV-A1 control, as detailed above.

Naturally occurring human COPD exacerbation cohort. A cohort of 40 COPD subjects was recruited to a study investigating naturally occurring exacerbations between the period November 2012–March 2013. Subjects were all grades of COPD exacerbation as confirmed by spirometry, were recruited and all treatments were approved. All subjects gave informed written consent and the study protocol was approved by the East London Research Ethics Committee (Protocol number 11/LO/0029). All subjects had an initial baseline visit during clinical stability for medical assessment, peak expiratory flow rate measurements, spirometry (forced expiratory volume in 1 s (FEV1), forced vital capacity (FVC) and clinical sample collection.

Sputum induction and processing: Sputum was induced using pre-medication with 200 μg salbutamol via metered dose inhaler and large volume spacer. Four per cent saline was administered using a DeVilbiss UltraNeB99 ultrasonic nebuliser until an adequate sputum sample was obtained. Sputum was processed within 2 h of induction. Sputum plug was selected from saliva by macroscopic inspection of the sample. An aliquot was selected and stored unprocessed at –80 °C. The remaining sample was weighed and 0.1% dithiothreitol (DTT) was added in the ratio 4 ml DTT to 1 g sputum. The mixture was then agitated and filtered. The same volume PBS was added, the filtrate centrifuged and the supernatant aliquotted and stored at –80 °C.

Virus detection: Viruses were detected in sputum using polymerase chain reaction. The following viruses were investigated: adenovirus, coronavirus, human bocavirus, human metapneumovirus, influenza (A/H1, A/H3, B), parainfluenza 1–3, picornavirus and respiratory syncytial virus. RES primers interacted with QIAamp viral RNA mini kit (Qiagen). For picornaviruses, 5 μl RNA was converted to cDNA by addition to a reaction mixture containing 1.55 μl OL27 (Invirotogen), 1 μl 0.1 M DTT, 0.2 μl dNTPs (Invitrogen) and 0.25 μl (10,000U) reverse transcriptase (Invitrogen). A single round of PCR of 32 cycles was used to detect picornaviruses in this cDNA. Differentiation of RVs from enteroviruses was achieved by restriction enzyme digestion of the PCR product.

Protein expression: For all other viruses, conversion to cDNA was achieved with 18 μl RNA added to a reaction mixture containing 2.5 μl of random hexamer primers (0.5 μg/μl, Promega) and 6.5 μl of UHQ (nuclease-free water, Promega). This mixture was polymerised for 40 cycles at 70 °C for 10 min. The final volume was adjusted to 70 °C on ice with 6 μl of UHQ, 5 μl 0.1 M DTT, 1.25 μl dNTPs and 2 μl (200U) reverse transcriptase (Qiagen) were added following incubation at 37 °C for 1 h to yield cDNA. An aliquot of 4 μl of this cDNA was then used for each RT-PCR in the panel, performed using a 2720 Thermal Cycler (Applied Biosystems). Viruses: Subjects were split into two groups: Subjects had respiratory symptoms at the time of the study team who were judged to be clinically stable and were followed up for a minimum of 6 months. Subjects reported to the study team when they developed symptoms of an acute exacerbation defined using the East London cohort criteria and were seen within 48 h of symptom onset. Sputum samples were collected at the onset of exacerbation and at 2 weeks after exacerbation.

Cell assays. In mouse models, BAL was performed by cannulation through the trachea and lavage with 1.5 ml sterile PBS. Cells were pelletted by centrifugation, resuspended in ACK buffer to lyse red cells, washed in PBS and resuspended in RPMI medium with 10% FBS. Cells were stained with Quik Diff (Reagena) for differential counting. For flow cytometry analysis, lung leukocytes were obtained from BAL and red cells were lysed with ACK buffer. BAL cells were stained with Live/Dead fixable dead cell stain (Life Technologies, Carlbad, CA), incubated with anti-mouse CD16/CD32 (FC block; BD Biosciences) and then incubated in a 1:1000 dilution of primary antibody. Secondary antibodies used for analysis of cell surface staining are shown in Supplementary Fig. 8.

Protein assays. Cytokine protein levels in mouse BAL, human sputum supernatants or cell supernatants from in vitro experiments were assayed using commercial ‘duoSet’ enzyme-linked immunosorbent assay kits (R&D Systems, Abingdon, UK). MUCSAC and MUC5B proteins in BAL were measured after adsorption to a 96-well plate by allowing samples to evaporate at 37 °C overnight. Plates were washed three times between steps. Plates were blocked for 2 h with 5% BSA in PBS. plates were then incubated following RV-A1 infection. Detection of MUCSAC, the antibody detection was used was biotinylated anti-MUC5B (Thermo Scientific) at 400 ng/ml. For the MUC5B assay, detection antibody was mouse anti-MUC5B clone E6 (MUC5B). Bound anti-MUC5B was detected with peroxidase-conjugated goat anti-mouse IgG (Sigma-Aldrich). Standard curves for each ELISA were generated by serial 1:2 dilutions of supernatants taken from ionomin-stimulated H929 cells (MUCSAC) and BAL supernatants previously taken from ovalbumin-induced hyper-allergic mice (MUCSB).

Western blotting was used to evaluate proteins in BEAS-2B cell extracts. Cells were lysed in radioimmunoprecipitation (RIPA) buffer (Sigma-Aldrich), supplemented with protease (Roche) and phosphate inhibitors (Sigma-Aldrich). Protein content was measured by the bicinchoninic acid assay (Thermo Scientific). Equal amounts of protein were loaded onto 4–12% Bis-Tris SDS-PAGE gels and transferred onto polyvinylidene difluoride (PVDF) membranes (both Life Technologies). Membranes were blocked with 5% RSA in Tris-buffered saline for 1 h at room temperature. The following primary antibodies were used: rabbit anti-pSTAT1 Y701 (1:1000 diluted; New England Biolabs), anti-pSTAT2 Y690 (1:1000, Life Technologies).
New England Biolabs), anti-STAT1 (1:1000 diluted; New England Biolabs) and anti-STAT2 (1:1000 diluted; New England Biolabs). Primary antibodies were incubated overnight at 4 °C with shaking and secondary antibody conjugated to horseradish peroxidase (1:5000 diluted; Jackson Immunoresearch) was incubated for 1 h at room temperature. Data were acquired using a Fusion FX7 image analyser (Vilber Lourmat). Uncropped western blot images are shown in Supplementary Fig 9.

For measurement of serum antibodies in the mouse model, peripheral blood was collected from the carotid arteries and serum RV-specific IgGs were measured by in-house enzyme-linked immunosorbent assay. Ninety-six-well plates were coated with purified RV1B, as used for in vivo infection, and incubated at 4 °C overnight. Plates were blocked with 5% milk in PBS-0.05% Tween 20 at room temperature. An aliquot of 50 µl per well of serum diluted 1:50 in 5% milk/PBS was then added with further 2 h incubation at room temperature. IgG detection antibodies were biotinylated rat anti-mouse IgG1 (0.5 µg/ml clone A85-1 BD Biosciences) and IgG2a (0.5 µg/ml clone R19-15; BD Biosciences).

Neutralisation assays. Neutralisation of RV was measured in Ohio HeLa cells (UK Health Protection Agency General Cell Collection) grown until ~90% confluency on 96-well plates. Pooled sera for each given treatment group were incubated with purified RV-A1 or RV-A1 control at room temperature with shaking for 1 h and then added to HeLa cells with further incubation at 37 °C for 48–72 h. Protection from cytopathic effect was measured by crystal violet cell viability assay. Cells were stained with 0.1% (w/v) crystal violet (Sigma-Aldrich) at room temperature for 10 min. Cells were then washed with water, air-dried and crystal violet was solubilised with 1% sodium dodecyl sulphate (SDS) solution (w/v). Absorbance was measured by a Spectramax Plus plate reader at 560 nm.

RNA extraction and cDNA synthesis and quantitative PCR. Total RNA was extracted from BEAS-2B cells, primary airway epithelial cells or sputum cells (RNAeasy kit, Qiagen) and 2 µg was used for cDNA synthesis (Omniscript RT kit). For mouse models, total RNA was extracted from the right upper lobe of mouse lung and placed in RNA later, prior to RNA extraction and cDNA synthesis (as detailed above). Quantitative PCR was carried out using previously described specific primers and probes for each gene of interest23,33, and normalised to 18S rRNA33. Reactions were analysed using ABI 7500 Fast Real-time PCR system (Applied Biosystems, Carlsbad, CA).

DNA extraction and bacterial 16S quantitative PCR. DNA extraction from human sputum and mouse lung tissue was performed using the FastDNA Spin Kit for Soil (Qbiogene, Santa Ana, USA), as per the manufacturer’s instructions. Bead-beating was performed for two cycles of 30 s at 6800 rpm (Precellys, Bertin Technologies). Total 16S bacterial loads were measured using quantitative PCR, performed in triplicate using the Via7 Real-Time PCR system (Life Technologies). In addition to template, each PCR reaction mix contained 2 µl of forward primer 520F (10 µm); 5′-AYTGGYDTAAAGNG-3′; 0.2 µl reverse primer 802R (10 µm); 5′-TACNVGGGTATCAATTACC-3′; 5 µl SYBR Fast Universal master mix (Kapa Biosystems) and 3.6 µl H2O. PCR cycling conditions were: 1 cycle of 90 °C for 3 min, followed by 40 cycles of 95 °C (20 s), 50 °C (30 s), 72 °C (30 s) and melt conditions of 1 cycle of 95 °C (15 s). For creation of the standard curve, a 1:10 dilution series (10−3–10−9 copies) of partial 16S rRNA gene of Vibrio natriegens DSMZ 759 (Deutsche Sammlung von Mikroorganismen, Braunschweig, Germany) from positions 27 to 1492 of the Escherichia coli reference cloned using TOPO TA cloning kit; (Invitrogen, ThermoFisher Scientific) was used.

Statistical analyses. For mouse experiments, animals were studied in group sizes of five-eight mice per experimental condition and data are presented as mean ± SEM, representative of at least two combined independent experiments. Data were analysed using one- or two-way ANOVA with significant differences between groups Bonferroni’s multiple comparison test. In vitro BEAS-2B experiments were performed three-five times and data are expressed as mean ± SEM. Data were analysed by one- or two-way ANOVA test with differences between groups assessed by Bonferroni’s multiple comparison test. In vitro primary airway epithelial cell experiments were performed with a group size of n = 9 patients and individual data points are shown. Data were analysed using one-way ANOVA with significant differences between groups assessed by Bonferroni’s multiple comparison test. Data from the human COPD cohort were analysed by using the Mann–Whitney U-test or Kruskal–Wallis test with Dunn’s post test comparison to predict protein levels or mRNA expression between ICS users and non-users.

Correlations between data sets were examined using Spearman’s rank correlation coefficient. All statistics were performed using GraphPad Prism 6 software. Differences were considered significant when p < 0.05.

Data availability. The data supporting the findings of the study are available in this article and its Supplementary Information files, or from the corresponding author on request.

Received: 11 September 2017 Accepted: 10 May 2018

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Acknowledgements
This work was supported by a Wellcome Trust Clinical Research Training Fellowship to A.S. (grant number WT1096382AIA), a pump priming grant from the British Lung Foundation to A.S. (grant number PRPRG15-9), a research grant from the British Medical Association to A.S. (grant ref: HC Roscoe 2015 grant), a National Institute for Health Research (NIHR) Senior Investigator Award to S.L.J., the NIHR Clinical Lecturer funding scheme (PM) and funding from the Imperial College and NIHR Biomaterial Research Centre (BRC) scheme.

Author contributions
A.S., N.G., M.R.E., P.M., N.W.B. and S.L.J. conceived and designed the experiments. A.S., N.G., M.R.E., P.M., N.W.B. and S.L.J. performed experimental work and analysed data. A.S., N.G., J.L.G., Y.M.C., A.M., J.D.P., M.T., R.P.W., L.J.F., J.A., J.Z., M.-B.T.-T., M.A.C., S.-L.L., P.C.V., P.S.P., K.S.N., A.T.R., P.L.J., M.R.E., P.M. and N.W.B. contributed to writing and critical review of the manuscript.

Additional information
Supplementary Information accompanies this paper at https://doi.org/10.1038/s41467-018-04574-1.

Competing interests: S.L.J. has personally received consultancy fees from Myelo Therapeutics GmbH, Concert Pharmaceuticals, Bayer, and Sanofi Pasteura, and Avargen, be and his institution received consultancy fees from Synargen, Novartis, Boehringer Ingelheim, Chiesi, GlassmthKline, and Centocor. S.L.J. is an inventor on patents on the use of inhaled interferons for treatment of exacerbations of airway diseases. M.A.C. was employed by Chiesi Pharmaceuticals from January 2015 to December 2018.

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