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INTRODUCTION

Randomized controlled trials have shown that blood eosinophil counts are a biomarker that predict the effects of inhaled corticosteroids (ICS) in chronic obstructive pulmonary disease (COPD) patients at increased exacerbation risk.\(^1\)\(^,\)\(^2\) The Global initiative for the management of Obstructive Lung Disease (GOLD) report recommends the use of blood eosinophil measurements to...

The relationship between airway immunoglobulin activity and eosinophils in COPD

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Abstract

In chronic obstructive pulmonary disease (COPD), the effects of inhaled corticosteroids are predicted by blood eosinophil counts. We previously briefly reported increased immunoglobulin (Ig)A and IgM levels in bronchoalveolar lavage (BAL) of COPD patients with higher (eosinophil\(^{\text{high}}\)) compared to lower (eosinophil\(^{\text{low}}\)) blood eosinophils (>250/μL versus <150/μL), suggesting differences in adaptive immune function. An inverse relationship exists between eosinophil counts and airway pathogenic bacteria levels. The mechanistic reasons for these associations between eosinophils, corticosteroids and pathogenic bacteria are unclear. IgA, IgM and IgG levels were assessed in BAL, bronchial biopsies and epithelium collected from eosinophil\(^{\text{high}}\) (\(n = 20\)) and eosinophil\(^{\text{low}}\) (\(n = 21\)) patients. Bronchial B-cell numbers were measured by immunohistochemistry. B-cell activity was assessed in bronchial samples and following exposure to BAL from eosinophil\(^{\text{high}}\) and eosinophil\(^{\text{low}}\) patients. BAL levels of non-typeable Haemophilus influenza (NTHi)-specific immunoglobulins were quantified. Results showed airway expression of IgA, IgG1 and IgM were lower in eosinophil\(^{\text{low}}\) compared to eosinophil\(^{\text{high}}\) patients, with lower levels of NTHi-specific IgA and IgM. Bronchial B-cell numbers were similar in both groups, but B-cell activity was lower in eosinophil\(^{\text{low}}\) patients. In conclusion, COPD eosinophil\(^{\text{low}}\) patients show differences in adaptive immune function compared to COPD eosinophil\(^{\text{high}}\) patients. These differences may cause different microbiomes in these COPD phenotypes.

KEYWORDS

B-lymphocytes, haemophilus influenza, immunoglobulin A, immunoglobulin G, immunoglobulin M

1 INTRODUCTION

Randomized controlled trials have shown that blood eosinophil counts are a biomarker that predict the effects of inhaled corticosteroids (ICS) in chronic obstructive pulmonary disease (COPD) patients at increased exacerbation risk.\(^1\)\(^,\)\(^2\) The Global initiative for the management of Obstructive Lung Disease (GOLD) report recommends the use of blood eosinophil measurements to...
help guide the use of ICS containing combination treatments in COPD patients.  

The mechanistic reasons for the increased ICS response in COPD patients with higher blood eosinophil counts are unclear. We previously performed a bronchoscopy study in 21 blood eosinophil low (<150 eosinophils/µl; eosinophillow) and 20 blood eosinophil high (>250 eosinophils/µl; eosinophilhigh) COPD patients to investigate biological differences associated with eosinophil counts.  

We reported higher lung eosinophil numbers, thicker reticular basement membrane and differences in the levels of various inflammatory mediators including IL-5, IL-13, CCL24 and CCL26 in eosinophilhigh compared to eosinophillow patients.  

Such findings are also observed in patients with asthma, providing insights into potential reasons for differential ICS responses associated with blood eosinophil counts in COPD.  

Our previous report noted higher IgA and IgM levels in the bronchoalveolar lavage (BAL) of COPD eosinophilhigh compared to eosinophillow patients, suggesting differences in adaptive immunity between these groups. An inverse association has been reported between sputum eosinophil counts and the levels of pathogenic bacteria in the airways.  

Additionally, COPD patients with blood eosinophil counts <100 cells/µl are more likely to have chronic bacterial airway infection. Overall, these findings support the concept that COPD patients with lower eosinophil levels are more susceptible to bacterial airway infection.  

Furthermore, COPD or asthma patients with raised blood and sputum eosinophils demonstrate greater presence of bacterial phylum Bacteroidetes, whereas low sputum eosinophil levels were associated with lower bacterial diversity and increased Proteobacteria, specifically the Haemophilus genus. These altered microbiome profiles associated with different eosinophil counts may cause distinct airway inflammation profiles in the airway which respond differently to anti-inflammatory drugs.  

Here, we report a further analysis using samples from our bronchoscopy study that compared COPD eosinophilhigh with eosinophillow patients; we focus on differences in adaptive immune function that may cause altered immunity against bacteria. We investigated IgA, IgM and IgG levels using different samples and techniques, and measured B-cell activation and bacterial opsonization.

2 | METHODS

2.1 | Patient recruitment and sample collection

21 eosinophillow (blood count < 150/µl) and 20 eosinophilhigh (>250/µl) COPD patients were recruited for bronchoscopy. Cut-offs were chosen as they identified upper and lower tertiles for blood eosinophils counts in COPD patients previously studied at the Medicines Evaluation Unit, Manchester, UK. As fluctuations in blood eosinophil numbers in COPD patients are typically minor, particularly at lower blood eosinophil counts, the exclusion of the middle tertile helped ensure the eosinophilhigh and eosinophillow patients were distinct populations. Initial results and clinical characteristics were reported previously (Table 1). Patients with COPD >40 years old with >10 pack-year smoking history and post-bronchodilator forced expiratory volume in 1 second (FEV1)/forced vital capacity (FVC) ratio <0.7 were recruited.  

Patients with a previous diagnosis of childhood or adult asthma, or those with atopy demonstrated by a positive skin prick test against house dust mite extract or cat dander or grass pollen allergens (Alk-Abello, Hørsholm Denmark) were excluded. Patients receiving oral corticosteroids or antibiotics within 6 weeks of recruitment were excluded. Bronchial biopsies, bronchoalveolar lavage and bronchial brush samples were collected; some patients were not able to tolerate collection of all samples. B cells were isolated from the blood of a healthy non-smoking volunteer. Sample collection was approved by the local research ethics committee (NRES Committee North West – Greater Manchester South; REC Ref: 06/Q1403/156). All subjects provided written informed consent.

2.2 | Gene expression analysis

Bronchial epithelial brushings were collected from the lower lobes (n = 20 eosinophilhigh; n = 17 eosinophillow), and blood was collected into PaxGene RNA tubes (BD biosciences, Wokingham, UK). Gene expression of the immunoglobulin genes IGHA1, IGHA2, IGHM, JCHAIN, IGHG1, IGHG2, IGHG3, IGHG4 and polymeric immunoglobulin receptor (PIGR) were assessed and genes associated with plasma cell activity, identified by literature review. These genes are listed in Table 2, including a functional description. Detailed methods are in the online data supplement.

2.3 | BAL Immunoglobulin assessment

BAL was successfully collected from n = 15 eosinophilhigh and n = 15 eosinophillow. Levels of IgG1 and IgG2 in BAL fluid were determined by ELISA (ThermoFisher). Secretory IgA levels were assessed by ELISA (Demeditec). IgA and IgM levels have been previously assessed in the same BAL samples by multiplex assay (Myriad RBM).  

Levels of immunoglobulins were normalized to patient-specific BAL urea concentrations, measured by colorimetric assay (Biovision Inc). The use of urea for normalization is not universally accepted, so for completeness, we also present non-normalized results in the online supplement.

2.4 | Bronchial biopsy Immunohistochemistry

Formalin-fixed, paraffin-embedded bronchial biopsies were assessed for levels of B cells and plasma cells, along with expression of IgA2, IgG1, IgG2 and IgM by immunohistochemistry. Detailed methods are in the online data supplement.
2.5 | In vitro B-cell treatment

B cells from the blood of a healthy non-smoking volunteer were exposed to BAL fluid from eosinophil\textsuperscript{high} and eosinophil\textsuperscript{low} patients. Healthy donor B cells were preferred over COPD B cells, which may be dysfunctional giving data that is difficult to interpret. The effect of BAL on X-box-binding protein 1 (XBP1) gene expression was assessed by quantitative PCR. Detailed methods are in the online data supplement.

2.6 | NTHi-specific immunoglobulin binding assay

Methodology for immunoglobulin-specific opsonization of non-typeable haemophilus influenza (NTHi) was adapted from Staples et al.\textsuperscript{19} Detailed methods are in the online data supplement.

2.7 | Statistical analysis

All results were assessed using Graphpad Prism version 7.04. Comparisons between eosinophil\textsuperscript{high} and eosinophil\textsuperscript{low} were by \textit{t} test or Mann–Whitney test, with distribution of data assessed by D’Agostino & Pearson normality test.

3 | RESULTS

3.1 | Subjects

Clinical characteristics for the 21 eosinophil\textsuperscript{low} and 20 eosinophil\textsuperscript{high} COPD patients are summarized in Table 1. Lung function, proportion of patients using ICS, symptoms and exacerbation rates were similar.
No history of immunodeficiency was reported for any patient, and blood immunoglobulin gene expression levels were similar in eosinophil high and eosinophil low patients, as were serum IgA and IgM levels (Table S1). Blood basophil numbers were higher in eosinophil high compared with eosinophil low patients.

### 3.2 | IgA and IgM BAL protein and epithelial gene expression

Previously, we briefly reported that COPD eosinophil high patients had higher absolute BAL IgA and IgM protein levels, compared with eosinophil low patients (Figure S1A,B). BAL recovery rates vary between patients, causing variable dilution effects. We now present BAL immunoglobulin results normalized to patient-specific BAL urea levels (Figure 1A); BAL IgA and IgM protein levels remained higher in eosinophil high compared to eosinophil low patients. Immunoglobulin gene expression in epithelial brushings from the same patients (Figure 1B) demonstrated higher expression of genes for the IgA and IgM heavy chains, and the J chain that enables IgA and IgM polymORIZATION, in the eosinophil high COPD patients. Secretory IgA levels in BAL from eosinophil high patients were higher compared with eosinophil low patients (Figure 1A; Figure S1C).

### 3.3 | IgG BAL protein and epithelial gene expression

Bronchial epithelial gene expression levels of IGHG1 and IGHG2 were greater in COPD eosinophil high patients (Figure 1B). The mean levels of IGHG3 and IGHG4 gene expression were also higher in the eosinophil high group, but did not reach statistical significance (Figure 1B). We then measured IgG1 and IgG2 levels in BAL; the levels of these immunoglobulins were higher in BAL from eosinophil high compared to eosinophil low patients (median IgG1: 4340 vs 1772 ng/ml respectively, P = .0003; median IgG2: 488 vs 133 ng/ml respectively, Table S1).
P = .0186; Figure 1A with urea normalized results and Figure S1D,E with non-normalized results).

3.4 | Immunoglobulin immunohistochemistry

IgA was predominantly found on the apical surface of the bronchial epithelium (Figure 2A). The percentage of the epithelium immunoreactive for IgA was significantly higher in the eosinophil

high patients compared with the eosinophil

low patients (Figure 2E).

Remodelled airways in COPD patients have limited apical IgA expression.20 This has been associated with a localized reduction in polymeric immunoglobulin receptor (pIgR) protein and gene expression, resulting in flawed IgA epithelial translocation. PIGR expression in epithelial brush samples was similar between the groups (eosinophil

high: 15.78 ± 0.55; eosinophil

low: 15.50 ± 0.57; P = .129), suggesting that the observed differences in apical IgA staining are unlikely due to differences in translocation.

The majority of bronchial biopsies were negative for epithelial IgM expression. Expression was observed in the submucosal bronchial glands (Figure 2B). Approximately 40% of biopsies contained submucosal glands making comparisons between groups difficult with small sample sizes.

IgG1 and IgG2 expression was identified in both the epithelium and subepithelium (Figure 2C,D). IgG1 subepithelium expression was significantly higher in eosinophil

high patients (Figure 2F), while subepithelial IgG2 expression was significantly higher in the eosinophil

low patients (Figure 2G); epithelial expression of IgG1 and IgG2 showed similar trends (Figure S2; P = .08 and P = .05, respectively).

3.5 | B-cell numbers and activity

Differences in immunoglobulin levels may be due to different B cells or plasma cell numbers. We assessed levels of CD19+ B cells and CD27+ plasma cells in bronchial biopsies by immunohistochemistry (Figure 3A,B). B cells and plasma cell numbers were similar between the eosinophil

high and eosinophil

low groups (Figure 3C,D). Similar gene expression of CD19 and CD27 in bronchial epithelial brush samples supported these findings (Figure 3E,F).

We treated healthy B cells with BAL fluid from eosinophil

high and eosinophil

low patients to assess B-cell activation. Cells treated with eosinophil

high BAL expressed higher levels of XBP1, a key regulator of plasma cell immunoglobulin production (Figure 4). We compared bronchial epithelial gene expression of mediators that influence plasma cell activity (Table 2). Levels of NOS2 and the lymphotoxins LTA and LTβ were significantly lower in eosinophil

low patients.

3.6 | NTHi opsonization

Reduced immunoglobulin levels in eosinophil

low COPD patients may confer reduced anti-bacterial activity. Immunoglobulin-specific NTHi opsonization experiments were carried out using BAL fluid. The results showed that approximately twice as many bacteria were opsonized with IgA and IgM using BAL fluid from eosinophil

high compared to eosinophil

low patients (Figure 5), indicating lower levels of NTHi-specific IgA and IgM antibodies in eosinophil

low patients. Levels of NTHi-specific IgG1 and IgG2 opsonization were similar between groups.

4 | DISCUSSION

We have demonstrated higher IgA, secretory IgA, IgM and IgG1 levels in COPD eosinophil

high compared with eosinophil

low patients. These results were consistent when immunoglobulins were measured by 3 different methods; BAL protein measurement, bronchial epithelial gene expression and bronchial mucosal immunohistochemistry. Furthermore, BAL supernatants from COPD eosinophil

high patients.

FIGURE 2 Immunohistochemical analysis of immunoglobulin expression in bronchial biopsies from eosinophil

high and eosinophil

low patients. Bronchial biopsies were formalin-fixed, paraffin-embedded. Tissue sections were stained for IgA2 (A), IgM (B), IgG1 (C) and IgG2 (D). Images represent typical staining for each immunoglobulin, with arrows identifying examples of positive areas; size bars illustrate 100μm. Comparisons of epithelial IgA2 (E), and subepithelial IgG1 (F) and IgG2 (G) between eosinophil

high and eosinophil

low were by Mann-Whitney test: *P < .05. Box and whisker plots represent median, interquartile range and range.
caused greater NTHi opsonization and B-cell activation. Overall, these findings indicate reduced anti-proteobacterial immunity in COPD eosinophil-low patients.

Pneumonia risk is greater in COPD patients with lower blood eosinophil counts. Lower airway eosinophil numbers are associated with increased presence of colonizing bacteria in stable COPD patients, and ICS use was associated with an increase in bacterial load only in patients with lower eosinophils. Overall, these previous findings suggest that COPD patients with lower eosinophil counts have a different susceptibility to bacterial infection. Our data showing lower immunoglobulin levels and activity in COPD eosinophil-low patients provide a potential mechanistic explanation for these observations. The risk of pneumonia in COPD patients is associated with certain clinical characteristics, such as previous pneumonia events, older age and lower body mass index. The use of ICS in such individuals may increase pneumonia risk further. Overall, there appear to be complex interactions between clinical characteristics, blood eosinophil counts and ICS use that influence pneumonia risk in COPD patients.

COPD and asthma patients with lower airway eosinophils have greater levels of proteobacteria, including Haemophilus. NTHi is found in the airways of COPD patients, both at stable state and during exacerbation. NTHi strains isolated from COPD patients have enriched virulence factors, including genes associated with iron uptake, immune evasion, adherence and oxidative stress counter-measures. In addition to these mechanisms that can potentiate or prolong NTHi activity, we now show lower antibody-mediated immunity against NTHi in COPD eosinophil-low patients.

IgA is the predominant immunoglobulin found in the human lung, with 80% of mucosal plasma cells being IgA+. BAL IgA levels from mild/moderate COPD patients are higher compared with controls. However, Du et al reported lower BAL IgA levels in severe/very severe patients compared with controls. We show lower BAL IgA levels in COPD patients with lower eosinophil levels. IgA is produced by subepithelial plasma cells before being transported through the epithelial layer in a complex with polymeric immunoglobulin receptor (pIgR). Released secretory IgA (sIgA) coats the mucosal surface, providing protection against microbes. The remodelled epithelium in COPD patients has reduced pIgR expression, causing decreased mucosal IgA. Areas of decreased IgA show localized increases in inflammatory cells and submucosal invasion by both bacteria and viruses. We observed that eosinophil-low COPD patients had lower mucosal IgA expression, which was not due to altered epithelial IgA translocation, as PIGR expression was similar between groups. Our findings may be due to reduced B-cell activity in the

![Figure 3](image-url)
eosinophil<sub>low</sub> patients. Lower BAL IgA levels in COPD have previously been associated with increased NTHi colonization, supporting a link between eosinophils, immunoglobulins and microbiome.

Eosinophil<sub>low</sub> patients also had lower levels of NTHi-specific IgA, which induced lower levels of bacterial opsonization. Staples et al showed reduced bacterial opsonization, following BAL supernatant exposure, results in lower levels of NTHi phagocytosis; this mechanism may contribute to the association between a low airway eosinophils phenotype and increased bacterial colonization.

IgG1, IgG2 and IgG3 levels are elevated in BAL from COPD patients compared with controls, but IgG1 and NTHi-specific IgG1 levels are lower in patients colonized by NTHi; IgG1 is the most abundant form of IgG. There were lower BAL levels of total IgG1 in COPD eosinophil<sub>low</sub> patients compared with eosinophil<sub>high</sub> patients, but we did not show a difference between groups for NTHi-specific IgG1 antibodies. Further work is required to see if eosinophil<sub>low</sub> patients have impaired IgG responses against other bacterial pathogens, such as Moraxella catarrhalis and Streptococcus pneumoniae. Although IGHG2 expression was lower in eosinophil<sub>low</sub> patients, higher protein levels were retained in the subepithelium. This may be due to IgG2 being actively retained within the tissue to combat subepithelial bacterial colonization associated with lower apical IgA expression.

Lymphoid follicles act as focal centres for antigen presentation and adaptive immune responses; increased numbers of B-cell rich lymphoid follicles are associated with COPD severity. IgM<sup><sub>+</sub></sub> plasma cells are the predominant B-cell found in airway lymphoid follicles. BAL IgM levels were reported to be higher in COPD patients compared to controls. We have shown lower levels of IgM BAL protein and epithelial gene expression in eosinophil<sub>low</sub> compared with eosinophil<sub>high</sub> COPD patients. It is probable that these epithelial gene expression results are due to bronchial brush samples contain a mixture of epithelial and other cell types including B cells.

We were unable to determine if the distribution of tissue IgM differed between the eosinophil<sub>high</sub> and eosinophil<sub>low</sub> groups. Immunohistochemical analysis showed that IgM was predominantly found in mucosal glands. Previous work has shown that immunoglobulins produced in glands do not distribute across the airway mucosal surface, but are localized to mucous plugs. Like IgA, we showed that eosinophil<sub>low</sub> COPD patients had reduced levels of NTHi-specific IgM in BAL compared with eosinophil<sub>high</sub> patients.

B-cell and plasma cell numbers are increased in both the large and small airways of COPD patients compared to controls. A recent study reported that lymphoid follicles in severe COPD patients have increased IgA<sup><sub>+</sub></sub> plasma cells. We found no differences
in B-cell or plasma cell numbers in biopsies from eosinophil\textsuperscript{high} and eosinophil\textsuperscript{low} patients. However, BAL fluid from eosinophil\textsuperscript{low} patients induced lower levels of XBP1 expression than BAL from eosinophil\textsuperscript{high} patients. The transcription factor Xbp1 is required for plasma cell differentiation and is essential for high levels of antibody secretion,\textsuperscript{35} and our findings suggest that the airways of eosinophil\textsuperscript{low} patients may be less primed to stimulate immunoglobulin production. In smoke-exposed mice, efficient clearance of NTHI has been shown to be B-cell dependent and relates to levels of NTHI-specific IgA.\textsuperscript{36}

The epithelial gene expression of NOS2 and lymphotoxins LTA and LTB was lower in the eosinophil\textsuperscript{low} group; these findings are discussed further in the online supplement.

We previously reported higher BAL IL-5 levels in eosinophil\textsuperscript{high} patients.\textsuperscript{3} This cytokine is important for eosinophil maturation, activation and survival, and stimulates IgA production by plasma cells in mice,\textsuperscript{37} although this has not been shown in humans. A Phase 2a placebo-controlled, randomized clinical trial in COPD patients with sputum eosinophilia showed that the anti-IL-5 receptor monoclonal antibody, benralizumab, caused near-complete depletion of blood and sputum eosinophils.\textsuperscript{38} Phase 3 clinical trials with benralizumab did not show efficacy in the primary analysis of exacerbation rate reduction,\textsuperscript{39} but pre-specified subgroup analysis demonstrated that benralizumab reduced exacerbation rates in patients with blood eosinophil levels ≥220 cells/µL, ≥3 exacerbations in the prior year and using inhaled triple therapy.\textsuperscript{39} This suggests that increased eosinophil activity has a mechanistic involvement in disease pathophysiology in a subgroup of COPD patients with certain clinical characteristics plus higher blood eosinophil counts.

The Phase III COPD clinical programs conducted with the anti-eosinophil therapies benralizumab and mepolizumab showed no evidence of increased exacerbations or bacterial infections compared to placebo,\textsuperscript{40,41} supporting that IL-5 and/or eosinophils are not directly responsible for the higher immunoglobulin levels seen in our eosinophil\textsuperscript{high} COPD patients. Furthermore, in the Phase II COPD trial, sputum microbiome analysis demonstrated no increase in airway bacterial load with benralizumab treatment, but rather a reduction in Streptococcus pneumonia levels, with no effect on Haemophilus influenzae.\textsuperscript{38} The authors proposed that Streptococcus pneumonia reduction may be due to macrophages being redirected from eosinophil efferocytosis towards bacterial phagocytosis after eosinophil depletion. Additionally, eosinophils have no direct anti-bacterial activity against Streptococcus pneumonia and Haemophilus influenzae.\textsuperscript{38} Overall, the eosinophil itself appears to have little direct anti-bacterial activity, but here we show that eosinophil\textsuperscript{low} COPD patients display differences in B-cell activity and immunoglobulin levels that may cause differences in microbiome.

We have recently shown higher airway IL-13 levels in the same group of eosinophil\textsuperscript{high} (versus eosinophil\textsuperscript{low}) patients in this report.\textsuperscript{4} IL-13 promotes humoral responses.\textsuperscript{42} It is possible that the T2 environment present in the airways of eosinophil\textsuperscript{high} COPD patients facilitates better anti-bacterial defence through mechanisms such as IL-13-stimulated B-cell activation.

Hyperesinophilic syndrome commonly involves the lungs (63% of patients).\textsuperscript{43} The airways from chronic eosinophilic pneumonia (CEP) patients contain many more eosinophils compared to eosinophil\textsuperscript{high} COPD patients; for example, median BAL eosinophil percentage in CEP is 52% and 0.75% in eosinophil\textsuperscript{high} COPD.\textsuperscript{3,44} However, like eosinophil\textsuperscript{high} COPD, CEP patients also have elevated airway type 2 inflammation\textsuperscript{44} and increased BAL levels of IgA, IgM and IgG.\textsuperscript{45}

We did not include healthy controls, as it has already been reported that immunoglobulin expression levels are higher in mild COPD patients compared with controls,\textsuperscript{39} but more severe patients have lower secretory IgA levels.\textsuperscript{27} We cannot state if the immunoglobulin levels observed here are elevated or decreased compared to normal levels, but can state that immunoglobulin levels are higher in eosinophil\textsuperscript{high} relative to the eosinophil\textsuperscript{low} patients. There is growing evidence that low eosinophil numbers are associated with bacterial colonization.\textsuperscript{6-8,10} A limitation is that we did not assess bacteriology in the current study, preventing comparisons between immunoglobulin levels and bacterial load. We had few severe COPD patients, as these individuals are practically more difficult to bronchoscope for research. The sample size of 41 subjects was modest, but sufficient to observe consistent biological differences between groups.

In conclusion, we have shown differences in adaptive immunity in COPD eosinophil\textsuperscript{low} patients compared with COPD eosinophil\textsuperscript{high} patients, associated with altered anti-proteobacterial immunity. These results can explain differences in the airway microbiome associated with eosinophil counts that have been reported.\textsuperscript{6-10}

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**CONFLICT OF INTEREST**

DS reports personal fees from AstraZeneca, personal fees from Boehringer Ingelheim, personal fees from Chiesi, personal fees from Cipla, personal fees from Genentech, personal fees from GlaxoSmithKline, personal fees from Glenmark, personal fees from Gossamerbio, personal fees from Menarini, personal fees from Mundipharma, personal fees from Novartis, personal fees from Peptinnovate, personal fees from Pfizer, personal fees from Pulmatrix, personal fees from Theravance, personal fees from Verona. SS, T-HP and PN are employees of AstraZeneca and also hold stock in AstraZeneca. T. Southworth, AH, UK, JL, JD and T. Scott have no conflicts to report.

**AUTHOR CONTRIBUTION**

Thomas Southworth: Conceptualization (lead); Formal analysis (lead); Funding acquisition (supporting); Investigation (equal); Methodology (equal); Project administration (lead); Resources (lead); Supervision (lead); Visualization (lead); Writing-original
draft (lead); Writing-review & editing (equal). Andrew Higham: Conceptualization (supporting); Formal analysis (supporting); Investigation (equal); Methodology (equal); Project administration (supporting); Supervision (supporting); Visualization (supporting); Writing-original draft (supporting); Writing-review & editing (equal). Umme Kolsum: Conceptualization (supporting); Formal analysis (supporting); Funding acquisition (supporting); Investigation (equal); Methodology (supporting); Project administration (supporting); Resources (lead); Supervision (supporting); Visualization (supporting); Writing-review & editing (equal). Jian Li: Formal analysis (supporting); Investigation (equal); Methodology (equal); Visualization (supporting); Writing-original draft (supporting); Writing-review & editing (equal). Josiah Dungwa: Formal analysis (supporting); Investigation (equal); Methodology (supporting); Visualization (supporting); Writing-review & editing (equal). Sriman Sridhar: Formal analysis (supporting); Investigation (equal); Methodology (equal); Writing-original draft (supporting); Writing-review & editing (equal). Paul Newbold: Conceptualization (lead); Funding acquisition (lead); Investigation (equal); Project administration (supporting); Supervision (supporting); Writing-review & editing (equal). Dave Singh: Conceptualization (lead); Funding acquisition (lead); Project administration (supporting); Resources (lead); Supervision (lead); Writing-original draft (lead); Writing-review & editing (equal).

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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SUPPORTING INFORMATION
Additional supporting information may be found online in the Supporting Information section.

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