Prosurvival activity for airway neutrophils in severe asthma

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ABSTRACT

Background Airway neutrophilia is a recognised feature of chronic severe asthma, but the mechanisms that underline this phenomenon are unknown. Evidence for factors present in airway secretions that prolong neutrophil survival has been sought and it has been hypothesised that these might be augmented in neutrophilic asthma.

Methods Non-smoking subjects with severe asthma (SA) or mild asthma (MA) and healthy control subjects (HC) underwent sputum induction. The SA group was subdivided into subjects with neutrophil counts above (SA-high) and those within the normal range (SA-low). Apoptotic neutrophils were enumerated in the cellular phase while the fluid phase was assessed for its ability to prolong the in vitro survival of blood-derived neutrophils using morphometric and flow cytometric analyses.

Results There was a significant difference between all four subject groups with respect to the percentage of apoptotic sputum neutrophils (Kruskal–Wallis, p=0.042). Cuzick test showed a highly significant (p=0.008) trend towards decreasing numbers of apoptotic neutrophils across the four groups with increasing asthma severity and neutrophil count. The sputum antiapoptotic activity was also different between the groups (p=0.039), with a highly significant (p=0.005) decreasing trend across the four groups. The survival effect could not be inhibited by blocking selective chemotaxin receptors, neutralising neutrophil survival factors, inhibiting phosphatidylinositol-3-kinase (using LY294002) or with pertussis toxin pretreatment. Similarly, it could not be explained by lipopolysaccharide contamination or by the presence of inhaled corticosteroids in sputum.

Conclusions These data demonstrate the capacity of as yet unidentified factor(s) in the sputum of subjects with asthma to delay human neutrophil apoptosis and extend their lifespan as a potential mechanism contributing to unresolving airways neutrophilia in severe asthma.

INTRODUCTION

Accumulating evidence points to an important role for neutrophils during acute exacerbations of asthma and in severe forms of the disease1–3 where raised neutrophil numbers have been observed in bronchoalveolar lavage (BAL), induced sputum and both bronchial and transbronchial biopsies.1 4 5 While increased concentrations of neutrophil-active mediators interleukin 8 (IL-8),6 7 leukotriene B4 (LTB4),7 8 granulocyte–macrophage colony-stimulating factor (GM-CSF)9 and tumour necrosis factor α (TNFα)10 have been detected in BAL and plasma of patients with severe asthma, a clear mechanistic association between these mediators and airways neutrophilia has not yet been demonstrated.

Theoretically, the observed neutrophilia in the airways of those with asthma could be due to increased chemotactic activity produced in the inflamed airways or longer survival due to locally produced factors, such as TNFα and GM-CSF, which delay their apoptosis.11 It is increasingly recognised that diverse phenotypes exist in asthma.12 13 It is also known that there is marked variability in airway cytokine/chemokine/growth factor profiles, expression of receptors and signalling pathways,4 14–17 all of which could represent different phenotypes. In a subgroup of subjects with severe asthma, partial or complete steroid resistance is seen.2 Paradoxically, corticosteroid treatment itself could contribute to airway neutrophilia in poorly controlled asthma by promoting neutrophil survival.18–20 While the exact role of neutrophils in asthma is still unknown, an extended lifespan and/or failed clearance of apoptotic neutrophils could enhance the proinflammatory potential of these cells to cause lung tissue damage and to impede the resolution of asthmatic inflammation.21

In order to elucidate the mechanisms that lead to airways neutrophilia in severe asthma, we have conducted a cross-sectional study of subjects with mild asthma treated with β2-agonists alone and those with severe asthma on high dose inhaled (ICS) and oral corticosteroids (OCS), using healthy individuals as control subjects. We have hypothesised that raised neutrophil numbers in more severe asthma are the result of their prolonged survival promoted by factors released by inflammatory and structural cells of the airways. In order to test this hypothesis, we first quantified the extent of in vivo apoptosis by enumerating sputum neutrophils that display characteristic morphological features of apoptosis. Further evidence of increased survival in asthmatic airways was then sought by studying whether factors present in the fluid phase of sputum, which samples the epithelial lining fluid (ELF), could alter the proportions of neutrophils that become apoptotic in neutrophil cultures. Any effect of residual corticosteroids in sputum samples was tested by pretreating neutrophils with the glucocorticoid (GC) receptor antagonist, mifepristone (RU486), added to neutrophil survival assays.22 The mechanisms underlying any prononeutrophil survival effect of the sputum fluid phase were then studied using selective blockers for mediator pathways known to delay neutrophil apoptosis.

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1 Supplementary methods and a figure are published online only. To view these files please visit the journal online (http://thorax.bmj.com).

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METHODS

Subjects
Non-smoking volunteers, classified as severe atopic asthmatics (SA; n=15), mild atopic asthmatics (MA; n=11) or healthy control subjects (HC (n=12) using established GINA (Global Initiative for Asthma) guidelines, were enrolled. The SA subjects were further divided into two subgroups according to the percentage of neutrophils in sputum using the 65% cut-off for normal neutrophil counts reported by Belda and co-workers\textsuperscript{23} to define SA with high counts (SA-high) and SA with low counts (SA-low) (for criteria see the online supplement). The study was approved by the Southampton and South West Hampshire Ethics committee and all subjects were recruited using advertisements in the media, either directly or via a departmental database.

Sputum induction and processing
Sputum induction was conducted using guidelines of the ERS (European Respiratory Society) Task Force on induced sputum\textsuperscript{24} with 4.5% hypertonic saline as described previously.\textsuperscript{25}

Sputum cell counts and quantification of apoptotic neutrophils
Immunocytochemistry was applied to sputum cell phase cyto- spins to identify neutrophils\textsuperscript{26} using mouse monoclonal anti-neutrophil elastase antibody, followed by secondary rabbit antimouse bionylated immunoglobulin G (IgG), streptavidin—AP (alkaline phosphatase) and Fast Red, and counterstained with haematoxylin. Apoptotic neutrophils were defined as those positively immunostained cells that also exhibited cell shrinkage, nuclear condensation and fragmentation, plasma membrane ruffling and blebbing, features known to be characteristic of apoptotic cells and widely used for their identification.\textsuperscript{21}

In vitro assessment of antiapoptotic activity of the sputum fluid phase
Neutrophils were obtained from normal healthy donors, purified to >98% purity using standard methods known not to activate neutrophils\textsuperscript{27} and resuspended at 4×10^6 cells/ml in RPMI 1640 with 20% fetal calf serum (FCS) plus 2%L-glutamine, sodium pyruvate supplemented with 50 U/ml streptomycin, and penicillin G, and incubated at 37°C in 5% CO2. For all subjects, experiments were carried out in this medium in the presence or absence of sputum fluid phase (final concentration 1:20, shown in preliminary experiments to be optimal in prolonging neutrophil survival), or lipopolysaccharide (LPS) (10 ng/ml), a known antiapoptotic factor for neutrophils, as a positive control. Experiments were conducted in parallel in the presence of the GC receptor antagonist, mifepristone (1 μM) in order to control for any residual corticosteroids present in sputum. In separate culture wells, the effect of mifepristone was controlled by testing its ability to abrogate the inhibitory effects of dexamethasone (100 nM). Dexamethasone and mifepristone were dissolved in ethanol (methasone (100 nM). Dexamethasone and mifepristone were control for any residual corticosteroids present in sputum. In 300 g for 10 min at 4°C, washed twice and resuspended with 10% heat-inactivated FCS/phosphate-buffered saline (PBS). Cytocentrifuged preparations were fixed in methanol and stained with Diff-Quik: this allowed blind counting of slides and for apoptosis to be scored morphometrically as those cells with decreased size and nuclear and cytoplasmic condensation (figure S1; online supplement). Numbers of apoptotic cells were reported as percentages of total neutrophil counts. In subsequent inhibition experiments, in which inhibitors of various mediators and signalling pathways were used to block the effect of sputum on in vitro apoptosis, the extent of apoptosis was assessed by flow cytometry using fluorescein isothiocyanate (FITC)-labelled Annexin V (BD Biosciences, Oxford, UK) and a FACS Calibur with Cell Quest software (BD Biosciences) by counting 10,000 events per sample. We have previously shown a strong (r=0.70) and highly significant (p<0.0001) correlation between the extent of neutrophil apoptosis measured using standard morphological criteria and the extent quantified by flow cytometry.\textsuperscript{27} For further details of all the experimental methods see the online supplement.

Statistics
All statistical analyses were performed using Prism version 5 for Windows (GraphPad Software, San Diego, California, USA) except the Cuzick test\textsuperscript{28} which was performed using STATA, Version 11.0 (StataCorp, College Station, Texas, USA). Sputum neutrophil numbers and neutrophil survival in in vitro assays were not normally distributed. These data are therefore expressed as median, IQR and range from the indicated numbers of experiments. Differences in these indices among the four groups of patients were assessed by the Kruskal–Wallis test. Where this returned a significant difference, the Cuzick non-parametric test for trend across the four groups (HC, MA, SA-low and SA-high) was performed.\textsuperscript{29} The effects of drugs on the antiapoptotic activity of sputum supernatants on neutrophils in vitro were assessed using the paired Wilcoxon signed-rank test. A probability of 0.05 was defined as significant.

RESULTS

Subject details
Subjects’ clinical characteristics and their sputum neutrophil and eosinophil counts are shown in table 1. Forced expiratory volume in 1 s (FEV1) was significantly lower in the SA-high and SA-low groups than in the MA or HC groups (all p<0.0001). Analysis of sputum revealed that the percentage of neutrophils in the SA-high group was significantly higher than in the SA-low, MA and HC groups (all p<0.0001).

Neutrophil apoptosis in sputum
The percentage of apoptotic neutrophils counted in the sputum cell phase was significantly different between the four subject groups as assessed by the Kruskal–Wallis test (p=0.042). Trend

Table 1 Subject characteristics

<table>
<thead>
<tr>
<th>Variables</th>
<th>HC (n = 12)</th>
<th>MA (n = 11)</th>
<th>SA-high (n = 6)</th>
<th>SA-low (n = 9)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>43±15.5</td>
<td>36±10.4</td>
<td>45±7.1</td>
<td>45±11.5</td>
</tr>
<tr>
<td>Gender (M:F)</td>
<td>9:3</td>
<td>7:4</td>
<td>4:2</td>
<td>5:4</td>
</tr>
<tr>
<td>Duration of asthma (years)</td>
<td>NA</td>
<td>20.8±12.3</td>
<td>33.3±12.5</td>
<td>35.4±16.9</td>
</tr>
<tr>
<td>FEV1 (% of predicted)</td>
<td>99:7.2</td>
<td>95:12.8</td>
<td>62±6.8*</td>
<td>72±18.8*</td>
</tr>
<tr>
<td>Sputum neutrophils (% of total inflammatory cells)</td>
<td>50.6±17.2</td>
<td>36.4±16.9*</td>
<td>81.1±9.9*</td>
<td>38.5±15.1</td>
</tr>
<tr>
<td>Sputum eosinophils (% of total inflammatory cells)</td>
<td>0.2±0.4</td>
<td>2.0±1.8*</td>
<td>2.6±2.6*</td>
<td>13.4±17.7*</td>
</tr>
</tbody>
</table>

Data are expressed as mean and SEM. The SA-high and SA-low groups were defined according to neutrophil counts >65% and <65% of total inflammatory cell counts, respectively.

For comparison of data, subjects' age, gender, duration of asthma, FEV1 and sputum cell counts.

\*p<0.0001 versus MA and HC.
\| p<0.0001 versus MA, SA-low and HC.
\t p=0.043 versus HC.
\f p=0.0012 versus HC.
\t p=0.0009 versus HC.

F, female; FEV1, forced expiratory volume in 1 s; HC, healthy controls; M, male; MA, mild asthma; NA, not applicable; SA, severe asthma.
analysis using the Cuzick test showed a highly significant decreasing trend in the ranked score of apoptotic neutrophils across the four groups with increasing asthma severity and neutrophil counts (Cuzick test for trend, p = 0.008) (figure 1).

Effect of sputum supernatant on neutrophil apoptosis
All sputum samples contained significant prosurvival activity, as demonstrated by their ability to reduce the degree of constitutive neutrophil apoptosis in vitro (median of 56.90%, range of 19.70–85.30%). However, the activity was significantly different between the four groups (p = 0.039, Kruskal–Wallis test) and analysis by the Cuzick test showed a highly significant (p = 0.005) decreasing trend in the ranked score of antiapoptotic activity across the four groups with increasing asthma severity and neutrophil counts (figure 2).

Effect of mifepristone on neutrophilic apoptosis
Mifepristone (1 μM), an antagonist of the GC receptor,22 had no significant effect (p = 0.608) on constitutive neutrophil apoptosis alone (n = 6; data not shown). In contrast, when used at 1 μM it completely inhibited the apoptosis-delaying effect of dexamethasone (100 nM) in all experiments (n = 7; data not shown). Mifepristone had no effect on the degree of neutrophil apoptosis when co-culturing sputum supernatants from SA, MA and HC in the absence or presence of mifepristone (1 μM) (figure 5), excluding the possibility that the prosurvival effects of sputum from SA were mediated by residual GC present in sputum.

Effects of inhibiting GM-CSF, TNFα, LTB4 and IL-8 on neutrophil survival in vitro
In order to identify factors responsible for the prosurvival effect observed in the sputum from SA subjects, neutralising monoclonal antibodies (mAbs) against TNFα (100 μg/ml) and GM-CSF (100 μg/ml), as well as the selective antagonists for the IL-8 receptor, CXCR2 (SB-225002, 100 nM), and the LTB4 receptor, BLT1 (CP-105696, 10 μM), were applied in neutrophil cultures in the presence of sputum fluid phase. Neither the spontaneous apoptosis in buffer control (figure 4A) nor the antiapoptotic effects of sputum (figure 4B) were significantly (p = 0.989 and p = 0.998, respectively) inhibited by any of the selective receptor antagonists or neutralising mAbs in isolation or in combination.

Study of the roles of signalling pathways and LPS in neutrophil survival mediated by sputum
To elucidate the signalling pathways involved in the antiapoptotic effects mediated by sputum from SA subjects,
Figure 4 Neutralisation/blockade of sputum-mediated neutrophil survival. Application of the BLT1 antagonist, CP-105696 (10 μM), or the CXCR2 antagonist, SB-225002 (100 nM), along with the antitumour necrosis factor antibody (anti TNF-α Ab) (100 μg/ml) or antigranulocyte macrophage colony-stimulating factor (GM-CSF) Ab (100 μg/ml), or with their isotype-matched immunoglobulin G (IgG) controls (100 μg/ml), does not affect spontaneous, control apoptosis (A) or sputum fluid phase-induced neutrophil survival (B). Data are shown as mean±SD of the percentages of apoptotic neutrophils from different non-atopic healthy donors and sputum from subjects with severe asthma with high sputum neutrophil counts.

Figure 5 Sputum-induced neutrophil survival effect is not regulated by lipopolysaccharide (LPS), phosphatidylinositol-3-kinase (PI3K) or Gi/o-type G-proteins. Neutrophils (4×10^6/ml) were incubated in full RPMI culture medium alone, or with sputum from five subjects with severe asthma, applying sputum alone (Untreated) or sputum in the presence of the lipopolysaccharide (LPS)-neutralising agent, polymyxin B (Poly B; 5 μg/ml), or using neutrophils pretreated with the PI3K inhibitor, LY294002 (30 μM), or the Gi/o blocker, pertussis toxin (PTx; 100 ng/ml, 2 h). Data are the mean±SD values of the percentages of apoptotic neutrophils.

**DISCUSSION**

This study has demonstrated the presence of prominent anti-apoptotic activity for neutrophils in sputum samples in both healthy subjects and patients with asthma. Although a number of studies have investigated the characteristics of neutrophil apoptosis in induced sputum from subjects with asthma in vitro, this is the first investigation of the relationship between sputum neutrophil apoptosis and the prosurvival effect of the fluid phase of sputum on human neutrophils suggesting that prosurvival factors in the ELF are important for regulating the extent of airway neutrophilia in both health and disease. Of relevance to the pathogenesis of severe asthma, the anti-apoptotic activity was highest in patients with increased asthma severity that was associated with neutrophilia (p=0.005 for trend). However, extensive analysis of sputum factors that could be responsible for this phenomenon using selective inhibitors and blocking antibodies failed to attribute this activity to any of the proinflammatory cytokines, chemokines and lipid mediators that we studied.

Severe asthma has been a focus of investigation of many studies, but the mechanisms underlying its pathogenesis remain poorly understood. It is widely believed that severe asthma is not a single disease entity but consists of several pathologically distinct phenotypes. Patients with severe asthma have persistent granulocytic inflammation and markedly differ in their airway cytokine/chemokine profile and their receptors when compared with those with mild or moderate asthma. Significant advances have been made using sputum eosinophilia and neutrophilia as biomarkers of eosinophilic and neutrophilic forms of asthma, respectively, although the exact pathogenic role of these granulocytes remains unclear. Thus, eosinophilic asthma is usually viewed as being responsive to corticosteroids; indeed, studies have shown that corticosteroid treatment tailored on the basis of sputum eosinophil counts is superior to treatment guided by symptom control alone. Eosinophilia in patients with severe asthma is also associated with enhanced mast cell activation, T lymphocyte infiltration and production of profibrotic factors. Consistent with the latter, sputum eosinophilia has been associated with persistent airflow limitation, possibly because eosinophils are a major source of transforming growth factor β (TGFβ), a key profibrotic factor. In contrast, persistent neutrophilia is a feature of airway inflammation in patients with non-eosinophilic forms of asthma that are poorly responsive to corticosteroids. The mechanisms underlying the accumulation of these two inflammatory cell phenotypes are unclear, although eosinophilia is largely explained by an overproduction of T helper (Th) type 2 cytokines that can be found in both atopic and non-atopic forms of asthma. It is widely recognised that neutrophilia is a consistent feature in asthma exacerbations in which context it
could be a beneficial response to viral infection. In the current study, we excluded bacterial infection/colonisation (ie, all sputum samples were tested by routine bacterial culture methods). However, we did not look for evidence of persistent respiratory viral or Chlamyphila (formerly Chlamydia) infection, pathogens which have been associated with corticosteroid resistance (albeit in guinea-pigs) and greater decline in lung function in those with severe asthma.

Neutrophils are key effector cells in innate immunity in the defence against bacterial, fungal and viral infections, so their increased presence in severe asthma does not necessarily imply that they cause harm. Inflammatory neutrophils can contribute to airway inflammation by virtue of their capacity to release cytotoxic proteases and reactive oxygen species. Such processes are considered to be a critical determinant in the development of acute respiratory distress syndrome and chronic obstructive pulmonary disease where they are associated with the decline in lung function and small airways dysfunction. It has been proposed that the neutrophilia seen in asthma is due to corticosteroid treatment as these agents directly promote neutrophil survival by inhibiting their apoptosis. In our study, we were unable to inhibit the prosurvival effect of sputum by using the GC receptor antagonist, mifepristone. While we cannot exclude an in vivo effect of corticosteroids, which could explain the reduced number of apoptotic neutrophils in severe neutrophilic asthma, this therapeutic in vivo effect cannot explain the observation of an ex vivo effect of factors present in the sputum fluid phase on blood-derived neutrophils.

Increases in concentrations of neutrophil-active factors, IL-8, LTβs, GM-CSF and TNFα, have been reported in both blood and airway secretions from subjects with asthma and we have recently shown these to be involved in promoting neutrophil chemotaxis mediated by the bronchial epithelium conditioned with epidermal growth factor. We postulated that these proinflammatory mediators might mediate the observed antiapoptotic activity of sputum, but this possibility was excluded by the ineffectiveness of specific neutralising/blocking agents for these neutrophil-active factors. We further investigated the intracellular signalling that might be involved in the observed survival-promoting activity and initially focused on PI3K which is known to modulate the survival responses of neutrophils. We found that the inhibition of apoptosis caused by the sputum fluid phase was not affected by the PI3K inhibitor, LY294002. We then sought to investigate whether the prosurvival activity is initiated by coupling of the receptor to a PTx-sensitive G-protein, such as Gi or Go, and found that pretreatment with PTx (100 ng/ml) for 2 h, optimally known to inhibit neutrophil functions mediated via these G-proteins, failed to block prosurvival effects induced by sputum. It was further speculated that the sputum might contain residual or contaminating levels of LPS which can also delay neutrophil apoptosis. However, we found no effects of the inactivator of endotoxins, polymyxin B, arguing against the possibility that the prosurvival effects of sputum are mediated by contaminating LPS.

One possible explanation for the observed antiapoptotic effects of sputum is ATP released from dying airway cells. This premise is based on several lines of evidence that ATP has prosurvival activity for neutrophils and is released from dying cells, notably in chronic inflammatory conditions of the airways such as cystic fibrosis. Neutrophils themselves also release ATP, particularly under conditions of hypoxia, and ATP could act in an autocrine or paracrine manner to enhance neutrophil longevity at sites of inflammation. It would be interesting to determine, through a blocking approach, whether selective purinergic receptor antagonism is sufficient to prevent sputum-induced neutrophil survival. A further possible explanation for the observed antiapoptotic effects of sputum is that neutrophil survival may be prolonged because of respiratory viral infections and atypical bacterial pathogens such as Mycoplasma and Chlamyphila pneumoniae which are known to cause exacerbations of asthma. However, our study did not look at whether these respiratory pathogens are present in the sputum of those with severe asthma or whether interaction with these respiratory pathogens alters the bioactivity of sputum to modulate neutrophil survival programmes, a possibility that warrants further investigation. It is important to note, however, that none of the patients had had an exacerbation of asthma for at least 6 weeks.

One limitation of this study is the use of neutrophils from healthy donors, which raises the question of whether survival of neutrophils from patients with asthma might be different. Patients with asthma could not serve as donors of blood neutrophils since their blood often contains high eosinophil counts and there are no methods of neutrophil purification which can selectively deplete these cells. It would also have been desirable to study the interaction between the sputum fluid phase and neutrophils from sputum itself and from the same (autologous) donors, but this would have been technically difficult because of the large numbers of eosinophils that are also seen in neutrophil-enriched samples. This study should, therefore, be seen as a demonstration of increased proneutrophil survival activity in the ELF, leaving some uncertainty as to how exactly this heightened neutrophil apoptosis-resistant effect in the airways of those with severe asthma interacts with the neutrophils of the same patients.

In summary, this study has shown, for the first time, the presence of a prominent prosurvival activity in the airways of both healthy individuals and patients with asthma, the levels of which increase relative to the neutrophil counts and are highest in patients with severe neutrophilic asthma. Further investigation is required to fully elucidate the survival pathways regulated by these sputum-derived factors and how they contribute towards the apoptosis-resistant phenotype of neutrophils present in the airways of subjects with severe asthma.

**REFERENCES**


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