Effect of inhaled steroids on airway hyperresponsiveness, sputum eosinophils, and exhaled nitric oxide levels in patients with asthma

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Abstract

Background—Airway hyperresponsiveness, induced sputum eosinophils, and exhaled nitric oxide (NO) levels have all been proposed as non-invasive markers for monitoring airway inflammation in patients with asthma. The aim of this study was to compare the changes in each of these markers following treatment with inhaled glucocorticosteroids in a single study.

Methods—In a randomised, double blind, placebo controlled, parallel study 25 patients with mild asthma (19–34 years, forced expiratory volume in one second (FEV₁) >75% predicted, concentration of histamine provoking a fall in FEV₁ of 20% or more (PC₂₀) <4 mg/ml) inhaled fluticasone propionate (500 µg twice daily) for four weeks. PC₂₀ to histamine, sputum eosinophil numbers, and exhaled NO levels were determined at weeks 0, 2, and 4, and two weeks after completing treatment. Sputum was induced by inhalation of hypertonic (4.5%) saline and eosinophil counts were expressed as percentage nonsquamous cells. Exhaled NO levels (ppb) were measured by chemiluminescence.

Results—In the steroid treated group there was a significant increase in PC₂₀, decrease in sputum eosinophils, and decrease in exhaled NO levels compared with baseline at weeks 2 and 4 of treatment. Subsequently, each of these variables showed significant worsening during the two week washout period compared with week 4. These changes were significantly different from those in the placebo group, except for the changes in sputum eosinophils and exhaled NO levels during the washout period. There were no significant correlations between the changes in the three markers in either group at any time.

Conclusions—Treatment of asthmatic subjects with inhaled steroids for four weeks leads to improvements in airway hyperresponsiveness to histamine, eosinophil counts in induced sputum, and exhaled nitric oxide levels. The results suggest that these markers may provide different information when monitoring anti-inflammatory treatment in asthma.

Keywords: airway hyperresponsiveness; asthma; fluticasone propionate; inhaled corticosteroids; nitric oxide; induced sputum

Asthma is an inflammatory disease of the airways associated with airway hyperresponsiveness to various bronchoconstrictor stimuli such as histamine. The accompanying inflammation is characterised by the presence of inflammatory cells such as T lymphocytes, neutrophils and eosinophils and their cytokines in the airway mucosa, as demonstrated in bronchial biopsy specimens. The current treatment of asthmatic patients is based on the belief that reducing airway inflammation is essential, and that control of such inflammation can be indirectly assessed by optimising symptoms and lung function. However, monitoring airway inflammation more closely by measurement of non-invasive and sensitive markers of inflammation, such as airway hyperresponsiveness, sputum eosinophils, or exhaled NO levels, may provide additional information for assessing asthma control.

Inhaled glucocorticosteroids are currently the most effective treatment for asthma, not only reducing symptoms and airway hyperresponsiveness but also leading to an improvement in airway inflammation. However, recent evidence has suggested that such treatment often provides only partial suppression of airway inflammation, as shown by persisting eosinophilic inflammation in the bronchial (sub)mucosa after long term inhaled steroid treatment in some patients. Among the non-invasive techniques, hypertonic saline induced sputum has been shown to be a reliable method for measuring eosinophilic airways inflammation. The number of eosinophils in sputum is associated with asthma severity and decreases following treatment with inhaled steroids. In addition, nitric oxide levels in exhaled air have also been proposed as a marker for disease severity in asthma. Indeed, inhaled glucocorticosteroids decrease the levels of exhaled NO in patients with asthma in a dose dependent way.

Although the effects of inhaled steroids on sputum eosinophils and exhaled NO have been well established, comparative analysis is required before any of these markers can be recommended in the monitoring of asthma treatment. In the present study we investigated treatment induced changes in airway hyperresponsiveness, sputum eosinophils, and exhaled NO levels in asthma. To that end we performed histamine challenge, induced sputum, and exhaled NO measurements before, during, and after four weeks of treatment with...
Table 1 Characteristics of the subjects

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<th>PC20 (mg/ml)</th>
<th>Eosinophils (%)</th>
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| Placebo group | | | | | | |
| 13          | M   | 23          | 23.3 (2.4)† | 96.2 (9.0)† | 0.91 (1.62)* | 2.85 (2.46)† | 6.30 (3.34)† |
| 14          | F   | 21          | 82           | 0.11         | 21.2          | 13.41    |
| 15          | M   | 29          | 80           | 0.14         | 24.6          | 12.05    |
| 16          | M   | 34          | 83           | 0.30         | 0.0           | 4.17     |
| 17          | M   | 21          | 98           | 0.46         | 1.6           | 13.40    |
| 18          | M   | 28          | 80           | 0.54         | NA            | 14.08    |
| 19          | M   | 25          | 86           | 0.73         | 0.0           | 3.26     |
| 20          | M   | 24          | 98           | 0.77         | 1.8           | 4.22     |
| 21          | F   | 24          | 106          | 0.89         | 3.2           | 2.48     |
| 22          | M   | 28          | 90           | 1.00         | 1.2           | 3.36     |
| 23          | F   | 28          | 106          | 1.20         | NA            | 5.05     |
| 24          | F   | 25          | 98           | 1.51         | 0.0           | 5.82     |
| 25          | M   | 19          | 97           | 1.70         | 0.4           | 9.26     |
| 26          | M   | 17          | 95.6 (10.6)† | 0.52 (1.38)* | 4.91 (8.98)* | 7.47 (4.37)* |

Table 2 Airway hyperresponsiveness, sputum eosinophils and exhaled nitric oxide (NO) levels during and after steroid and placebo treatment

| Steroid group | | | | | | |
| PC20 (mg/ml) | 0.91 (1.62) | 3.19 (1.54)* | 3.67 (1.05)* | 0.93 (1.50)† | 6.30 (3.34)† |
| Eosinophils (%) | 0.85 (2.46) | 0.68 (0.94)* | 0.44 (0.56)* | 8.14 (7.50)* |
| NO (ppb) | 6.30 (3.34) | 1.52 (1.28)* | 1.43 (0.86)* | 5.22 (4.20)† |

| Placebo group | | | | | | |
| PC20 (mg/ml) | 0.52 (1.38) | 0.64 (1.21) | 0.59 (1.86) | 0.66 (1.26) |
| Eosinophils (%) | 4.91 (8.98) | 4.62 (6.04) | 5.74 (5.25) | 8.69 (9.87) |
| NO (ppb) | 7.47 (4.37) | 7.27 (5.38) | 5.97 (3.42) | 7.14 (5.59) |

FEV1 = forced expiratory volume in one second; PC20 = provocative concentration of histamine causing a 20% fall in FEV1; NO = exhaled nitric oxide; NA = not applicable.

†p<0.03 compared with week 4.

| p<0.01 compared with baseline.

Methods

SUBJECTS
Twenty five non-smoking atopic patients (16 men, age range 19–34 years) with mild persistent asthma volunteered to participate in the study (Table 1). Symptoms of episodic chest tightness and wheezing were treated by on-demand usage of inhaled salbutamol alone, which was discontinued at least eight hours before the measurements. Two weeks before the study all subjects were free from symptoms of respiratory tract infection. Atopy was indicated by a positive skin prick test (>3 mm weal) to one or more of 10 common airborne allergen extracts (Vividagnost, ALK, The Netherlands). The forced expiratory volume in one second (FEV1) was more than 75% of the predicted value and all subjects were responsive to inhaled histamine (provocative concentration causing a fall in FEV1 of 20% or more (PC20) of <4 mg/ml). The study was approved by the medical ethics committee of the Leiden University Medical Center and written informed consent was obtained from all volunteers.

DESIGN OF STUDY
The study was of a randomised, double blind, placebo controlled, parallel design. During screening the selection criteria were checked for all subjects. Before entering the treatment period baseline values of PC20 histamine and percentage eosinophils in induced sputum were determined. These two measurements were carried out on two separate days with a 2–4 day interval between them. Prior to histamine challenge and sputum induction, baseline values of FEV1 and exhaled NO were recorded. This sequence of measurements was used at all time points during the study.

Immediately after the second baseline visit the subjects were treated with inhaled fluticasone propionate (500 µg twice daily) or placebo for a period of four weeks. The measurement of PC20 histamine, sputum eosinophils, FEV1 and exhaled NO were repeated during the treatment period (at weeks 2 and 4) and during the washout period at two weeks after the treatment period.

HISTAMINE CHALLENGE
Histamine challenges were performed according to a standardised methodology. Histamine-di-phosphate (Sigma Chemicals, St Louis, MO, USA) in phosphate buffered saline (PBS) was stored at 4°C and administered at room temperature. Doubling concentrations between 0.06 and 16 mg/ml were used. The aerosols were generated by a DeVilbiss 646 nebuliser (output 0.13 ml/min), connected to an inspiratory and expiratory valve box with an aspiratory aerosol filter (Pall Ultipur BB50T). Each dose was inhaled through the mouth by tidal breathing for two minutes at five minute intervals with the nose clipped.

The airway responses to the inhaled aerosols were measured using FEV1, recorded by a dry rolling seal spirometer (Morgan Spiroflow, Morgan UK) and monitored on-line by a personal computer with a special software program. Before each test FEV1 was measured in triplicate for calculation of mean baseline levels. The airway response was recorded at 30 and 90 seconds after each dose. After each inhalation the lowest technically satisfactory FEV1 value was applied in the analysis to calculate the percentage fall in FEV1 from baseline. The test was discontinued if FEV1 decreased by 20% or more. The PC20 was calculated by log-linear interpolation of the final two data points.

SPUTUM INDUCTION
Sputum was induced and processed by the so-called full sample method according to a protocol that has been validated in our laboratory. Hypertonic saline aerosols (NaCl 4.5%) were generated at room temperature by a DeVilbiss Ultraneb 2000 ultrasonic nebuliser with a calibrated particle size (MMAD 4.5 µm) at maximal output (2.5 ml/min). The aerosols were administered to the subjects through a 100 cm long tube with an internal diameter of 22 mm...
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administered if needed. FEV₁ was measured and salbutamol was
injected into a clean plastic container by coughing. After testing,
water, and to expectorate sputum into a clean container.
started coughing, they were asked to blow their nose, to rinse their mouth and throat with water, and to expectorate sputum into a clean plastic container by coughing. After testing, FEV₁ was measured and salbutamol was administered if needed.

**SPUTUM PROCESSING AND CELL DIFFERENTIAL COUNTS**

The volume of the induced sputum samples was determined and mixed with an equal volume of 0.1% sputolysin (dithiotreitol, Calbiochem, USA). To ensure complete homogenisation the samples were placed in a shaking water bath at 37°C for 15 minutes and then gently mixed. The homogenised sputum was centrifuged (350 × 10³ g) for 10 minutes at room temperature. The cell pellet was resuspended in PBS to a final volume of 2–5 ml, then filtered through a gauze (pore size approximately 1 mm) to remove clumps. Total cell counts were performed in a haemacytometer (Tumson, Zoetermeer, The Netherlands). The sample was then diluted with PBS to a final concentration of 0.3 × 10⁶ cells/ml which was used for preparation of the cytocentrifuge slides (1500 rpm, three minutes, 50 µl/slide; Shandon 3, Life Sciences International, Veldhoven, The Netherlands).

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**DIFFERENTIAL COUNTS**

Differential counts of eosinophils, neutrophils, lymphocytes, macrophages, epithelial and squamous cells were performed on Diff-Quik stained cytocenters by a qualified cytopathologist. To correct for the variable salivary contamination, differential leucocyte and cylindrical epithelial cell counts were expressed as a percentage of 250 nucleated cells, excluding squamous cells. For each sample differential cell counts were performed twice by the same observer and the mean values were used in the analysis. A sputum sample was considered adequate when the percentage squamous cells was less than 80%. The reproducibility of the sputum cell counts obtained by this method has been shown to be satisfactory. To ensure a blind analysis of the sputum samples all cytocentrifuge slides were coded before analysis by an investigator who was not involved in the counting.

**EXHALED NO**

Exhaled NO levels were measured by a chemiluminescence analyser (Sievers NOA 270B) according to a standardised procedure which has previously been used by our laboratory. The subjects were connected to a closed system to avoid contamination of the measurements with ambient NO. Pressurised air with a low NO concentration (<1 ppb) was administered through a 150 litre reservoir connected to the inspiratory side of a Hans-Rudolph three way valve. The subjects performed a slow vital capacity manoeuvre with a constant expiratory flow of 10 l/min against an expiratory resistance of 3–4 cm H₂O. The expiratory NO concentration was sampled continuously from the centre of the mouthpiece at a flow rate of 440 ml/min and the average concentration (in parts per billion; ppb) was determined for a period of 10 seconds. Baseline values of exhaled NO were obtained from the mean values of the two NO measurements recorded before histamine challenge and sputum induction because their reproducibility was good (intraclass correlation coefficient, R > 0.92).

**ANALYSIS OF DATA**

PC₂₀ was log transformed before statistical analysis and expressed as geometric mean (SD) doubling doses. Based on their close to normal distribution, the percentage of eosinophils in sputum and the levels of exhaled NO were expressed as mean (SD). To test for differences between and within the treatment groups in general, multivariate analysis of variance (MANOVA) was applied for FEV₁ and log PC₂₀ whilst the Kruskal-Wallis test was used.
for sputum eosinophils and exhaled NO. The changes in PC20 (expressed in doubling doses), sputum eosinophils, and exhaled NO levels within each treatment group were analysed using the Student’s paired t test whilst changes in PC20, sputum eosinophils, and exhaled NO levels between both groups were tested using the Student’s unpaired t test, providing the 95% confidence intervals (95% CI). Finally, Pearson correlation analysis was used to examine the relationship between the changes in PC20, sputum eosinophils, and exhaled levels of NO. The results were considered significant if the p value was <0.05. All statistical analyses were performed using the SPSS program.

Results
Three of the subjects dropped out during the washout period between weeks 4 and 6 because of a history of respiratory tract infection (nos 5 and 6) or because they were taking an antihistamine (no. 11). Three subjects (nos 9, 18 and 23) did not produce adequate sputum at baseline, whilst subjects 21 and 7 were not able to produce sputum at week 2 and week 4, respectively. These time points were handled as missing data.

LUNG FUNCTION AND HISTAMINE CHALLENGE
At baseline there were no significant differences in FEV1 and PC20 between the groups (p>0.19; table 1). During the study there were no significant changes in FEV1 in the two groups (p>0.96, MANOVA). In the placebo group there were no significant changes in PC20 (p = 0.92, MANOVA) while in the steroid treated group PC20 increased significantly at week 4 compared with baseline values (mean change 2.01 (95% CI 0.683 to 2.090); p = 0.001; fig 1). After a two week washout period PC20 decreased again compared with week 4 by –1.75 (–1.831 to –0.582) doubling doses (p = 0.002; table 2, fig 1). These changes were significantly different from the changes in the placebo group (p<0.003; table 3).

SPUTUM EOSINOPHILS
The mean (SD) percentage of squamous cells in this study was 33.4 (17.6)%. Baseline sputum eosinophils were not significantly different in the two groups (p = 0.31; table 1). There were no significant changes in sputum eosinophils within the placebo group (p = 0.85, MANOVA), but in the steroid treated group a significant decrease in sputum eosinophils was observed compared with baseline values (mean change at week 4 –2.46 (95% CI –4.260 to –0.660)%; p = 0.01) with a subsequent worsening in the washout period compared with week 4 (mean change 6.13 (95% CI 0.804 to 11.459)%; p = 0.03; table 2, fig 2). The changes in sputum eosinophils were not significantly different between the two groups when baseline values were compared with week 4, or week 4 values were compared with those in the washout period (table 3).

EXHALED NO
At baseline exhaled NO levels were not significantly different in the two groups (p = 0.55; table 1). During the study there were no significant changes in exhaled NO levels in the placebo group (p = 0.54, MANOVA; table 2) but in the steroid treated group the levels of exhaled NO decreased significantly at week 4 compared with baseline values with a mean change of –4.88 (95% CI –6.862 to –2.892) ppb (p < 0.001), with a subsequent increase during the washout period compared with

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Within the steroid group there were no significant correlations between the changes in PC_{20}, sputum eosinophils, and exhaled NO at any time point (Pearson’s r <0.56, p >0.15; figs 4–6).

**Discussion**

The results of this study indicate that four weeks of treatment with inhaled steroids leads to improvements in airway hyperresponsiveness, sputum eosinophils, and levels of exhaled NO in patients with mild atopic asthma. In addition, it appears that the improvements in these markers are lost two weeks after cessation of treatment. This suggests that each of these markers may be useful for monitoring patients with asthma, even though there might be small differences between the markers in their earliest response to anti-inflammatory treatment.

To our knowledge this is the first study to compare the treatment induced changes in airway hyperresponsiveness to histamine, eosinophil counts in induced sputum, and exhaled NO levels in a group of asthmatic patients. Our study confirms and extends the results of others who have shown the beneficial effect of glucocorticosteroids on each of these markers separately. Like Kraan et al, we found an improvement of two doubling doses in airway hyperresponsiveness after four weeks of treatment with inhaled steroids. Furthermore, our findings are in agreement with those of Keatings et al and Kharitonov et al who demonstrated a decrease in sputum eosinophils and exhaled NO levels in patients, respectively, after treatment with inhaled steroids.

Although cross sectional relationships between airway hyperresponsiveness, sputum eosinophils, and exhaled NO levels in patients with asthma have been reported previously, only limited data are available on the comparison of within subject changes in these markers during treatment follow up. Our results are in agreement with those of Baraldi et al who also failed to find a correlation between steroid induced changes in PD_{20} and sputum eosinophils. The absence of such relationships may reflect the partially distinct pathophysiological backgrounds of these markers and might indicate the possible independent complementary clinical information during anti-inflammatory therapy.

We do not believe that our data were influenced by measurement errors since we used validated and reproducible methods. All subjects in the study were carefully selected as non-smokers with stable atopic asthma who had not used inhaled steroids for at least one month prior to the study. We chose a relatively high dose of inhaled steroid to ensure an optimal anti-inflammatory effect. To avoid carryover effects the histamine challenge for determination of PC_{20} and the sputum induction were separated by 2–4 days. Furthermore, exhaled NO levels on these two days appeared to be highly reproducible. Our inability to show a significant improvement in lung function following steroid treatment may be due to the normal baseline levels of FEV_{1} in our study (77–111% of the predicted value).

How can the present findings be interpreted? Firstly, corticosteroids are likely to decrease the percentage of eosinophils in the sputum by reducing the release and subsequent effects of cytokines such as interleukin 5 (IL-5) and granulocyte-macrophage colony-stimulating factor (GM-CSF) on eosinophil infiltration and survival. Secondly, the steroid induced reduction in exhaled NO levels can be explained by the inhibition of inducible NO synthase (iNOS) expression directly and/or indirectly by reduction in the levels of stimulatory cytokines, for instance in epithelial cells. Finally, the improvement in the physiological marker PC_{20} is likely to be due to effects of steroids on the presence and activity of multiple (infiltrative and resident) cells. Hence, it may not be surprising that the steroid induced changes in the three markers were not significantly correlated with each other. It would appear that early improvement of
eosinophils in sputum in response to steroid treatment is somewhat out of phase with the other two markers. However, we believe that this has few implications, given the consistency in the changes between the markers after four weeks of treatment.

What are the clinical implications of the present findings? Treatment according to the current guidelines is based on minimising symptoms and optimising lung function. However, frequently this fails to provide complete suppression of airway inflammation. It has been postulated that persistent airway inflammation in asthma leads to airway remodelling and an irreversible loss of lung function. This may require the use of more direct markers for monitoring airway inflammation. Indeed, a recent study by Sont et al showed that the adjustment of long term inhaled steroid treatment, additionally guided by the level of airway hyperresponsiveness, leads to a significantly better clinical, as well as histological, outcome than treatment guided by symptoms and lung function alone.

Based on the present data, it is now necessary to determine in long term prospective trials whether monitoring sputum eosinophils and/or exhaled NO levels can provide similar benefits in the management of asthma.

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References


