Effect of regular inhaled salbutamol on airway responsiveness and airway inflammation in rhinitic non-asthmatic subjects

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Abstract

Background – Regular, inhaled β agonists may increase airway responsiveness in asthmatic subjects. The mechanism is not known but may be via an increase in airway inflammation. A study was undertaken to examine the effect of regular inhaled salbutamol on airway responsiveness to methacholine and hypertonic saline, on the maximal response plateau to methacholine, and on inflammatory cells in induced sputum in rhinitic non-asthmatic subjects.

Methods – Thirty subjects with a baseline maximal response plateau of >15% fall in forced expiratory volume in one second (FEV1) entered a randomised, placebo controlled, parallel trial consisting of two weeks run in, four weeks of treatment, and two weeks washout. Methacholine challenges were performed at the beginning of the run in period, before treatment, after treatment, and after washout. Hypertonic saline challenges were performed before and after treatment and induced sputum samples were collected for differential cell counting.

Results – There was no change in airway responsiveness, maximal response plateau to methacholine, or in induced sputum eosinophils or mast cells. The maximum fall in FEV1, after hypertonic saline increased in the salbutamol group (median change 6.0%, interquartile range (IQR) 11.0) but did not change in the placebo group (median change 1.3%, IQR 5.5).

Conclusions – Regular inhaled salbutamol for four weeks increases airway responsiveness to hypertonic saline but does not alter airway responsiveness to methacholine or cells in induced sputum in non-asthmatic individuals with rhinitis. The relevance of these findings to asthmatic subjects has not been established.

Keywords: airway responsiveness, regular salbutamol, rhinitic non-asthmatic subjects, sputum cells.

Recent studies have suggested that regular β agonist administration may increase airway responsiveness to histamine and methacholine in asthmatic subjects.1-3 The mechanism by which this occurs is not known, but one possibility may be via an increase in airway inflammation. Persistent bronchodilatation induced by regular β agonist administration may allow an increased allergen load to reach the airways.4 Alternatively, tachyphylaxis of β receptors present on mast cells may reduce their stabilising effect and allow inflammatory mediator release in response to a lesser stimulus than is normally required.5 6

Methacholine and histamine act directly on airway smooth muscle to cause bronchoconstriction which is related to thickening of the airway wall, smooth muscle contraction, and lung elastic recoil.7 Hypertonic saline is believed to cause bronchoconstriction indirectly by the release of inflammatory mediators from airway mast cells and possibly airway nerves,8 and may therefore be a marker of inflammatory processes. Inhalation of hypertonic saline is also the basis of techniques which induce sputum for cytological examination.9

Although the sensitivity of the airways is a useful indicator of the degree of airway responsiveness, the maximal extent of airway narrowing has been recognised as an important measurement in recent years. Maximal airway narrowing can be assessed during bronchial challenge by increasing the dose of inhaled stimulus until the forced expiratory volume in one second (FEV1) ceases to decrease in response to further provocation, producing a plateau.10 It has been suggested that the maximal response plateau may be an index of greater clinical relevance than airway sensitivity as it is a reflection of the potential severity of an exacerbation of asthma.11

The aim of this study was to determine if regular treatment with a β agonist (salbutamol) increases airway sensitivity to direct and indirect stimuli, increases the maximal response plateau, and increases the degree of inflammation of the airways, as assessed by inflammatory cells and mediator concentration in induced sputum. Non-asthmatic subjects with rhinitis with mildly increased maximal response plateaux at baseline were selected to provide subjects with a mild abnormality of the airway that did not require treatment.

Methods

Subjects

Subjects with symptoms of rhinitis who had not used β agonist or corticosteroid medications in the previous 12 months were recruited by advertisement. At screening a medical history was taken and high dose methacholine chal-
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Subject characteristics were assessed by skin prick test. Chemicals, Sydney, Australia) was administered upon completion of the procedure. Subjects were required to cease salbutamol or placebo treatment at least six hours before methacholine challenge.

**LUNG FUNCTION MEASUREMENTS**

Spirometric tests were performed on a Vitalograph dry spirometer (Vitalograph Co, Buckingham, UK). Each measurement was repeated until two curves were obtained in which the FEV₁ fell by 20% from the baseline value or when the longest hypertonlic saline exposure was completed.

**HIGH DOSE METHACHOLINE CHALLENGE**

Methacholine chloride (ACIC Inc, Ontario, Canada) was administered using a DeVilbiss No. 646 nebuliser (DeVilbiss Co, Somerset, Pennsylvania, USA) regulated by a breath activated dosimeter (Rosenthal French, Baltimore, Maryland, USA) and attached to compressed oxygen at 138 kPa pressure. The dosimeter was set to allow nebulisation for 0.6 seconds per inhalation which produced a mean (SE) output of 0.01 (0.0011) ml per inhalation. A control dose of 0.9% sterile saline was administered at the commencement of each challenge, followed by methacholine chloride in concentrations of 3, 6, 25, 50, 100 and 200 mg/ml. Five inhalations, from slightly below functional residual capacity to total lung capacity, were taken of each solution. An additional five inhalations were taken of the 6 mg/ml solution and an additional 10 inhalations were taken of the 200 mg/ml solution. This method delivered methacholine chloride in cumulative, approximately doubling, doses ranging from 0.15 μmol to 199 μmol.

The FEV₁ was measured 60 seconds after the final inhalation of each solution. FVC was recorded at baseline, after the control dose, when a 20% reduction in FEV₁ occurred, and after administration of the final dose. The procedure was terminated when a plateau had been established, when the highest dose had been administered, or when the FEV₁ had fallen by 60% of the control FEV₁ value. Inhaled salbutamol, 200 μg, was administered upon completion of the procedure. Subjects were required to cease salbutamol or placebo treatment at least six hours before methacholine challenge.

**HYPERTONIC SALINE CHALLENGE**

Challenges with hypertonic saline were undertaken on the day following methacholine challenge. A solution of 4.5% saline (Ajax Chemicals, Sydney, Australia) was administered via an Ulthaneb 2000 ultrasonic nebuliser (DeVilbiss Co, Somerset, Pennsylvania, USA) for durations of 30 seconds, one minute, two minutes, four minutes, and two periods of eight minutes. The canister containing the saline solution was weighed before and after the challenge, and the rate of output of the nebuliser in ml/min was determined. The test was stopped when the FEV₁ fell by 20% from the baseline value or when the longest hypertonic saline exposure was completed.
Sputum induction

Sputum was induced during the hypertonic saline challenge. Subjects were asked to produce sputum by forced coughing after the four and eight minute periods of inhaling hypertonic saline. In subjects who failed to produce sputum during the standard challenge an additional eight minute period of inhaled 4.5% saline was given after administration of 200 µg of inhaled salbutamol.

Differential cell count

At least four sputum plugs were selected from the sputum samples from each subject; the plugs were smeared onto glass slides and allowed to air dry. When possible, plugs were selected from sputum produced late in the procedure. All samples were smeared within two hours of production. Two slides from each subject were fixed in absolute methanol for 10 minutes, then stained in undiluted May-Grünwald stain for 10 minutes, briefly rinsed in tap water, and counterstained in Giemsa stain diluted 1:10 in Giemsa buffer (pH 6.8). Two slides were also fixed in Carnoy’s solution for 10 minutes, dehydrated in 70% alcohol for one minute, rinsed in tap water, and stained in 0.5% toluidine blue stain (pH 0.5) for 10 minutes.

Differential cell counts for neutrophils, macrophages, lymphocytes, bronchial epithelial cells, and eosinophils were performed on slides stained with May-Grünwald-Giemsa. The best slide from each subject at each saline challenge was selected and scanned on low power (40 ×) for a region of at least 400 cells which were able to be counted continuously. Cells were counted on high power (100 ×) across the whole width of the slide in a direction perpendicular to the direction of smearing until a total of 400 cells had been reached. Mast cell counts were performed on the slides stained with toluidine blue by the same procedure, except that a total of 1500 cells was counted.

Cell counting was performed blind to the subjects’ identities. Results for each cell type were expressed as a percentage of the total cells counted. Intraobserver repeatability of differential cell counts was assessed by recounting 10 slides selected at random. Interobserver repeatability was assessed by a second reader counting the same 10 slides.

Eosinophil cationic protein analysis

To the remaining sputum sample an equal volume of 1% dithiotreitol was added. This was vortexed and placed in a shaking water bath for 30 minutes. The sample was then centrifuged at 700 g for five minutes at 4°C and the supernatant removed and stored at 20°C until eosinophil cationic protein (ECP) assay using an RIA-ECP kit (Pharmacia).

Data analysis

Dose-response curves to methacholine and hypertonic saline were constructed by plotting the percentage fall in FEV₁ from the control FEV₁ value against the log dose of methacholine or hypertonic saline. A maximal response plateau to methacholine was established when the decrease in FEV₁ between two or more of the final doses was less than 5% of the control FEV₁ value and the level of the plateau was defined as the mean of the fall in FEV₁ at these doses. By this definition of the plateau the maximal percentage fall in FEV₁ to hypertonic saline was also calculated. Dose response ratios (DRRs) to methacholine and to hypertonic saline were calculated from the first point of the plateau as the percentage fall in FEV₁ divided by the cumulative dose of methacholine or hypertonic saline.

The maximal response plateau to methacholine, FEV₁, FVC, cytological, and ECP data are reported as arithmetic means with 95% confidence intervals (95% CI); dose response ratios to methacholine are reported as geometric means with 95% CIs and the change in DRR is reported in doubling dose units. Two-tailed paired t tests were used to analyse changes within a treatment group for these data, and two tailed unpaired t tests were used to analyse changes between the two treatment groups. Maximal falls in FEV₁ and DRR to hypertonic saline are reported as medians with interquartile ranges (IQR); the Wilcoxon signed rank sign test was used to analyse changes within a treatment group, and the Mann-Whitney U test was used to analyse changes between treatment groups. The distribution of categorical data was analysed using Fisher’s exact test. Correlations between data were examined by least square linear regression analysis. Significance for all tests was accepted at the 95% level. Repeatability of the maximal response plateau and the DRR to methacholine were analysed by the method of Peat et al. and repeatability of cell counts was assessed with analysis of variance by calculation of the coefficient of reliability (R).

Results

subjects

Of the 60 volunteers screened for the study, 33 had mildly elevated maximal response plateaux to methacholine and were eligible to participate in the trial. Of these 33 subjects, one withdrew because of illness and one because of non-compliance. The data obtained from one subject were not included in the analysis because of poor repeatability of baseline spirometric values. The baseline characteristics of the salbutamol and placebo subject groups are shown in table 1. All but four subjects were lifetime non-smokers, and none of the subjects had smoked for 10 years prior to the study. None had current symptoms of asthma or any significant chronic illness. All but two subjects completed the six assessment visits of the trial. Data from these two subjects have been included in the analysis, however, as they completed the post-treatment high dose methacholine challenge. Twelve subjects (five salbutamol, seven placebo) reported symptoms of respiratory tract infections during the trial, but there was no consistent effect of these episodes on any measurement. Seven subjects (four sal-
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Table 1 Baseline characteristics of subject groups at the initial screening visit

<table>
<thead>
<tr>
<th></th>
<th>Salbutamol (n = 16)</th>
<th>Placebo (n = 14)</th>
</tr>
</thead>
<tbody>
<tr>
<td>M:F</td>
<td>6:10</td>
<td>4:10</td>
</tr>
<tr>
<td>Age (years)</td>
<td>40.9 (34.0 to 47.9)</td>
<td>31.7 (25.3 to 38.2)</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>169.5 (164.8 to 174.2)</td>
<td>171.5 (167.3 to 175.7)</td>
</tr>
<tr>
<td>FEV1 (% predicted)</td>
<td>104.9 (98.2 to 111.5)</td>
<td>105.6 (95.8 to 115.5)</td>
</tr>
<tr>
<td>Number atopic</td>
<td>15</td>
<td>12</td>
</tr>
<tr>
<td>Methacholine DRR</td>
<td>1.2 (0.7 to 2.2)</td>
<td>0.9 (0.3 to 1.6)</td>
</tr>
<tr>
<td>Methacholine plateau (% fall FEV1/μmol)</td>
<td>3.8 (11.0)</td>
<td>1.7 (3.0)</td>
</tr>
<tr>
<td>Hypertonic saline DRR</td>
<td>0.24 (0.60)</td>
<td>0.08 (0.14)</td>
</tr>
<tr>
<td>Hypertonic saline plateau (% fall FEV1)</td>
<td>28.3 (22.8 to 33.8)</td>
<td>25.8 (20.1 to 31.5)</td>
</tr>
</tbody>
</table>

FEV1 = forced expiratory volume in one second; DRR = dose-response ratio.

Values for age, height, FEV1 (% predicted), and methacholine plateau are mean and 95% confidence intervals. Values for methacholine dose-response ratio (DRR) are geometric means and 95% confidence intervals. Values for hypertonic saline challenge, dose-response ratio and plateau are median and interquartile range.

butamol, three placebo) reported past treatment with anti-asthma or present treatment with anti-rhinitis medication, but there was no consistent effect on any measurement.

Compliance with treatment was checked by weighing the canisters from the returned metered dose inhalers. The weight loss from the 28 canisters returned showed that 25 subjects took at least six inhalations per day, while three subjects took only five inhalations per day. These three subjects were all in the placebo group. Two subjects who failed to return their canisters gave a verbal assurance of compliance. Exclusion of the data from these five subjects did not alter the results of any statistical analysis.

MAXIMAL RESPONSE PLATEAU

There was no difference in mean baseline FEV1 or FVC values at any assessment. The repeatability of the maximal response plateau was ±7.2% for the entire group and did not differ significantly between active (±7.5%) and placebo (±7.1%) groups. Treatment with salbutamol had no effect on the maximal response plateau to methacholine. Figure 1 shows mean values for the maximal response plateau at screening, before and after treatment, and at the follow up assessment. The mean change in the maximal response plateau before and after treatment was a fall of 3.1% (−1.8 to 7.95) in the salbutamol group and a fall of 0.6% (−3.61 to 4.8) in the placebo group.

DOSE RESPONSE RATIO TO METHACHOLINE

Treatment with salbutamol had no significant effect on the DRR to methacholine. Figure 2 shows the mean DRRs of the active and placebo groups at the four high dose methacholine challenges completed during the study. The mean change in DRR before and after treatment was 0.79% fall/μmol (0.42 to 1.51) in the salbutamol group and 0.66% fall/μmol (0.37 to 1.17) in the placebo group (p = 0.65). The repeatability of the DRR to methacholine at the two pretreatment measurements was ±2.1 doubling doses (DD) for the entire group and did not differ significantly between the active group (±2.4 DD) and placebo group (±2.0 DD). There was a significant correlation between the DRR to methacholine and the maximal response plateau (r = 0.73, p < 0.001).

HYPERTONIC SALINE CHALLENGE

Figure 3 shows the individual changes in maximal percentage fall in FEV1 to hypertonic saline in the active and placebo groups. All but two subjects had a plateau on their dose response curve to saline. The treatment groups differed significantly both in median change in maximum percentage fall in FEV1 (salbutamol 6.0% (IQR 11.0), placebo 1.3 (5.5%), p = 0.04) and change in DRR (salbutamol 0.45%/ml (IQR 1.04), placebo 0.11 (0.29)%/ml, p = 0.03). In the active group the maximum percentage fall in FEV1 increased from a median
Table 2  Mean (95% confidence interval) sputum differential cell counts before and after treatment with salbutamol as a percentage of total cells counted

<table>
<thead>
<tr>
<th>Group</th>
<th>Before treatment</th>
<th>After treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Salbutamol group</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bronchial epithelial cells</td>
<td>5.1 (0 to 10.5)</td>
<td>8.6 (0.3 to 16.9)</td>
</tr>
<tr>
<td>Macrophages</td>
<td>44.8 (33.0 to 56.9)</td>
<td>47.4 (34.9 to 59.9)</td>
</tr>
<tr>
<td>Neutrophils</td>
<td>43.0 (29.0 to 56.9)</td>
<td>37.1 (18.9 to 55.3)</td>
</tr>
<tr>
<td>Lymphocytes</td>
<td>3.0 (2.0 to 3.9)</td>
<td>3.4 (2.0 to 4.7)</td>
</tr>
<tr>
<td>Eosinophils</td>
<td>4.4 (0 to 10.2)</td>
<td>3.7 (0.3 to 7.1)</td>
</tr>
<tr>
<td>Mast cells</td>
<td>0.09 (0 to 0.17)</td>
<td>0.22 (0 to 0.5)</td>
</tr>
<tr>
<td><strong>Placebo group</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bronchial epithelial cells</td>
<td>6.3 (0 to 14.3)</td>
<td>2.7 (1.1 to 4.2)</td>
</tr>
<tr>
<td>Macrophages</td>
<td>48.0 (37.8 to 58.2)</td>
<td>57.9 (47.5 to 68.3)</td>
</tr>
<tr>
<td>Neutrophils</td>
<td>41.0 (29.2 to 52.9)</td>
<td>33.3 (22.8 to 43.8)</td>
</tr>
<tr>
<td>Lymphocytes</td>
<td>2.6 (1.9 to 3.3)</td>
<td>3.4 (2.1 to 4.8)</td>
</tr>
<tr>
<td>Eosinophils</td>
<td>2.2 (0.5 to 3.8)</td>
<td>2.8 (0.0 to 6.9)</td>
</tr>
<tr>
<td>Mast cells</td>
<td>0.16 (0 to 0.4)</td>
<td>0.13 (0 to 0.4)</td>
</tr>
</tbody>
</table>

**Sputum cytology**

Twenty two subjects (79%) were able to produce sputum during both saline challenges, and 24 (85%) were able to produce sputum during at least one saline challenge. Only the data from the 22 subjects (11 in each group) who produced samples both before and after treatment were included in the analysis. For the entire subject group the mean differential eosinophil count at baseline was 3.3% of total cells counted (95% CI 0.5 to 6.0) and the mean mast cell count was 0.12% (95% CI 0.0 to 0.3). There was no significant difference in mean cell counts of the active and placebo groups either at baseline or after treatment (table 2). The mean change between the cell counts before and after treatment did not differ significantly between groups for either eosinophils (active −3.74 (−9.8 to 2.5); placebo −3.0 (−6.3 to 0.2); p = 0.83) or mast cells (active 0.1 (−0.1 to 0.4); placebo −0.1 (−0.2 to 0.4); p = 0.13). Exclusion of the non-atopic subjects did not alter the results. Intraobserver and interobserver repeatability of cell counts was very good (coefficient of reliability >0.8) or good (coefficient of reliability 0.61–0.8) for all cell types, except for the intraobserver counts of lymphocytes which were moderately repeatable (R = 0.44).

There was no relationship at baseline or after treatment between differential sputum eosinophils or mast cell counts and the DRR or the maximal response plateau to methacholine, or the maximal percentage fall in FEV1, or DRR to hypertonic saline. The eosinophil and mast cell counts at baseline did not predict change in DRR or maximal percentage fall in FEV1, to hypertonic saline or methacholine after treatment, and there was no relationship between individual changes in eosinophil or mast cell count and individual changes in maximal response plateau or responsiveness to hypertonic saline.

**ECP analysis**

Eosinophil cationic protein was measured in paired sputum samples from the 14 subjects (eight in the salbutamol group and six in the placebo group) for whom sufficient sputum remained after four sputum plugs were removed for cell counting. There was no significant difference between the groups in the ECP levels before and after treatment or the mean change in ECP after the treatment period (active −11.7 ng/ml (−45.1 to 21.8); placebo 0.32 ng/ml (−9.8 to 10.4); p = 0.49). There was a tendency for sputum levels of ECP to fall in the salbutamol group (30.4 ng/ml (95% CI 0.7 to 60.0) before treatment, 19.9 ng/ml (1.1 to 38.5) after treatment), but this did not achieve statistical significance. There was no change in the sputum levels of ECP in the control group (6.5 ng/ml (−3.1 to 16.3) before treatment, 6.9 ng/ml (4.0 to 9.7) after treatment).

**Discussion**

This study has shown that salbutamol, inhaled regularly for four weeks, did not increase airway sensitivity or maximal response plateau to methacholine in rhinitic non-asthmatic subjects with mildly increased maximal response plateaux. Salbutamol use was associated with a small but significant increase in airway responsiveness to hypertonic saline. There was no detectable change in inflammatory cells present in induced sputum.

Previous studies1−3 which have suggested that regular β agonists cause an increase in airway hyperresponsiveness have only examined asthmatic subjects, so it is not clear whether such effects are peculiar to the asthmatic state. Non-asthmatic rhinitic subjects were recruited for this study for several reasons. Firstly, we wished to exclude subjects who were taking regular β agonist medication to avoid any confounding effects of persistent β receptor stimulation. Secondly, we wished to include subjects with a maximal response plateau to inhaled methacholine of more than 15% fall in FEV1. This criterion was based upon the results of an open trial, previously undertaken in our laboratory, which showed a significant increase in plateau after regular β agonist use only in non-asthmatic subjects with a baseline plateau of more than 15%. Many subjects with rhinitis have mild airway hyperresponsiveness,18 and we expected that some of our patients would have an increased maximal response plateau; 55% of screened subjects met the criterion of a maximal response plateau of more than 15%. In asthmatic individuals it is not usually possible to measure a maximal response plateau during methacholine challenge. Thirdly, subjects with rhinitis are likely to be atopic. If β agonists alter airway sensitivity by increasing the extent of allergic inflammation of the airways, as has been proposed,1 this change might be most...
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evident in atopic individuals. Of the 30 subjects recruited in this study 29 had rhinitis and 27 were atopic.

At baseline the active group was slightly more responsive to methacholine and to hypertonic saline than the control group, although this difference was not significant. Four subjects had responses to methacholine in the hyperresponsive range (DRR >5% fall in FEV₁/µmol methacholine) and two of them experienced a fall in FEV₁ of more than 20% during the hypertonic saline challenge. A response of this magnitude is usually seen only in asthmatic individuals, although these subjects did not report symptoms of asthma and were not being treated for asthma. Despite double blind randomisation, two of the three non-atopic subjects (including the non-rhinitic subject) were assigned to the control group, and three of the four subjects with airway hyperresponsiveness (including the two subjects hyperresponsive to hypertonic saline) were assigned to the active group. Use of a crossover study design may have overcome this problem, but this approach was rejected because, at the time the study was planned, we were unable to determine the length of the washout period between treatments required to reverse any effect of β agonists. Exclusion of the data from the hyperresponsive subjects did not substantially alter the results.

Regular inhaled salbutamol was associated with a small but significant increase in sensitivity and a maximal percentage fall in FEV₁ to hypertonic saline but had no effect on sensitivity or the maximal response plateau to methacholine. While methacholine constricts airway smooth muscle directly, hypertonic saline acts indirectly via release of mediators from airway inflammatory cells. The results of this study suggest that regular salbutamol affects the indirect, but not the direct, component of airway responsiveness. However, regular salbutamol inhalation was not associated with a change in eosinophil numbers or ECP levels in bronchoalveolar lavage fluid. An increase in mast cell activity is a possible explanation for the increased airway responsiveness to hypertonic saline, which is believed to induce bronchoconstriction by stimulating mast cell degranulation. Beta receptors are present on mast cells and exert a stabilising effect on these cells. Researchers have suggested that increased mast cell activity results from tachyphylaxis of the β receptor on circulating human leucocytes and, in animal models, agonist-induced tachyphylaxis of β receptors on airway mast cells has been demonstrated. Studies of regular β agonist use in asthmatic subjects have also suggested that increased mast cell activity, resulting from tachyphylaxis of the mast cell β receptor, is responsible for an increase in airway responsiveness. Three studies have shown an increase in both the early and late asthmatic response to inhaled allergen following regular β agonist use. Another study reached a similar conclusion by comparing the broncho-protective effect of terbutaline to challenge with methacholine and with AMP (a mast cell stimulus) after seven days of treatment with terbutaline. Tachyphylaxis of β receptors modifies the release of mediators from mast cells, then the results of the present study, in which both the sensitivity and the level of the plateau to hypertonic saline increased, suggest that both the sensitivity and the level of the plateau to hypertonic saline challenge. Following treatment with salbutamol one of these subjects experienced a substantial increase in the maximal percentage
fall to hypertonic saline; the other subject showed a large increase in both the maximal percentage fall to hypertonic saline and maximal response plateau to methacholine, and reported wheezing and breathlessness during the treatment period.

In summary, this study has shown that regular inhaled salbutamol increases airway responsiveness to inhaled hypertonic saline in non-asthmatic individuals. This change is consistent with an effect on mast cell activity, although the mechanism is not known.

The authors thank Ms Deanne Anson for her important contribution to this work in preparing the sputum samples for ECP analysis and performing the assay. We are also grateful to Ms Lorna Gutierrez and Dr Peter Gibson for their expert advice on the identification of cells in sputum.

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