Failure of ipratropium bromide to modify the diurnal variation of asthma in asthmatic children

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ABSTRACT Thirty one children with asthma were given 40 μg of ipratropium bromide and identical placebo by inhalation three times a day in a double blind, randomised crossover study to test the ability of an anticholinergic drug to modify the diurnal variation in airway calibre and bronchial reactivity. Subjects measured peak expiratory flow rate approximately eight hourly, before and after inhaled salbutamol, for four week periods. Paired t tests and cosinor analysis were used to assess the diurnal variation in airway calibre from the peak expiratory flow rate recorded before salbutamol and to assess the diurnal variation in bronchodilator responsiveness from the increase in peak expiratory flow rate after salbutamol. Maintenance treatment with ipratropium bromide 40 μg three times daily reduced the provocative dose of histamine which caused a 20% fall in FEV₁ (geometric mean PD₂₀ = 0.78 v 0.49 mg/ml, p < 0.05), despite an eight to 12 hour gap between the last dose of ipratropium and histamine challenge. It did not, however, diminish the diurnal variation in airway calibre (mean amplitude = 12.7 v 10.1) or in bronchodilator responsiveness (mean amplitude = 62.4 v 63.5). There was no improvement in the clinical state of subjects while they were taking ipratropium bromide.

A diurnal variation in airway calibre, as judged by peak expiratory flow (PEF), has been described in adults with asthma¹ and children with asthma.² Three The mechanisms underlying this rhythm are unknown. Diurnal variation in airway sensitivity to histamine³ and to acetylcholine⁴ has also been described and could not be explained by variation in baseline airway calibre. The mechanisms underlying this diurnal variation in airway sensitivity are not known. Bronchoconstriction after inhalation of histamine is thought to be mediated in part by vagal reflexes.⁵ Thus the diurnal variation in airway sensitivity to histamine, and other substances, may also be mediated in part through the vagus.

Nocturnal exacerbations of asthma have been attributed to an exaggeration of the diurnal variation in airway calibre,¹ but the heightened airway sensitivity seen in the early hours of the morning⁴ ⁶ may also be a contributing factor. If nocturnal asthma is related to the diurnal variation in airway calibre or in bronchial reactivity, or both, a treatment regimen that would abolish or diminish these variations might be able to decrease some of the morbidity and mortality associated with nocturnal asthma.⁷ ⁸ In recent studies anticholinergic drugs have been shown to increase early morning PEF more than evening PEF⁹ and to decrease morning dipping,¹⁰ which suggests that these agents may be able to diminish the diurnal variation in airway calibre.

This study was performed to test the ability of the anticholinergic agent ipratropium bromide in clinically acceptable doses to modify the diurnal variations in airway calibre and in bronchodilator responsiveness. Thirty one children with asthma were given ipratropium bromide and placebo in a double blind crossover study. The diurnal variation in airway calibre was assessed from PEF recordings and the diurnal variation in bronchodilator responsiveness assessed from the increase in PEF after salbutamol.
Thirty one children with asthma (21 male, 10 female), aged from 8 to 18 years (mean 12.0 years), were studied. The study consisted of four periods of four weeks: a baseline period and two treatment periods separated by a washout period. Each child kept a diary card recording a daily symptom score, medications taken, and PEF (Wright mini peak flow meter) approximately eight hourly, before and 10 minutes after inhaling salbutamol from an aerosol. PEF was measured on waking in the morning, at 1600 hours, and at bed time, as suggested by Hetzel and Clark.1 At each test time the children carried out three measurements and recorded the highest PEF on the diary card. Children were maintained on their usual medication and carried out normal activities during the study. Medications used were: salbutamol, two puffs from a metered dose aerosol (200 µg) or 0.03 ml/kg up to a maximum of 1·0 ml of a 0·5% solution by wet nebulisation; a 12 hour slow release theophylline preparation at a dose of 20 mg/kg per day to a maximum dose of 500 mg twice daily or as directed by blood concentrations; and inhaled beclomethasone dipropionate by a metered dose aerosol, 50 µg per puff, two to four puffs three or four times daily.

Each day the children were required to give a score between zero and three for each of the following: amount of wheeze during the night; amount of cough during the night; amount of wheeze during the day; and limitation of activity during the day. Symptom scores were added to give a daily total for each child. Daily symptom scores could range from zero, for an asymptomatic day, up to 12 for a day with a pronounced cough, wheeze, and limitation of activity. Daily symptom scores were added to give a symptom score for each period for each child. The study treatment consisted of two puffs of either ipratropium bromide (40 µg) or placebo three times a day from identical metered dose aerosols labelled A or B. The children were randomly allocated to receive either aerosol A or B first.

Histamine challenge tests were performed, following the protocol of Yan et al.,11 at the same time of day, at the completion of each study period. Subjects did not use salbutamol or treatment aerosols for between eight and 12 hours before the challenge test. Other medications were continued, but as subjects were taking the same treatment regimen throughout the study this should not contribute to any differences found between the challenge tests. Dose-response curves were plotted and the cumulative dose of histamine causing a 20% fall in FEV1 (PD20) calculated.

The study protocol was approved by the hospital's ethics committee, and informed consent was obtained from each subject and at least one parent.

DATA ANALYSIS
All statistical tests were performed on natural log transformed PD20 data. Wilcoxon's matched pairs signed ranks test was used to compare PD20 values between study periods. Dunnett's paired t test12 was used to test differences between afternoon, morning, and evening PEF values for each child within each period, afternoon PEF being considered as “control” and morning and evening PEF as two “treatments.”

The amplitude of the diurnal variation in PEF was assessed by cosinor analysis, using the modified technique reported by Hetzel and Clark.1 PEF was analysed against time using the equation:

$$PEF = Co + a \cos(2\pi t/24) + b \sin(2\pi t/24),$$

where Co = mean value or intercept; t = time PEF was measured. Coefficients a and b were determined and used to calculate the “peak to trough” amplitude (A = 2√(a^2 + b^2)). Amplitude was expressed as a percentage of the subject’s mean PEF for the period under study, and compared between treatment periods by paired t test.

The increase in PEF (l/min) after salbutamol inhalation was calculated at each test time.Bronchodilator responsiveness was analysed as the absolute increase in PEF as a function of time by cosinor analysis, as described above. Amplitude, expressed as a percentage of the subject’s mean increase in PEF after salbutamol inhalation for the period under study, was compared between treatment periods by paired t test. Dunnett’s paired t test12 was used to compare the increase in PEF after salbutamol at each test time on data for each child.

Symptom scores and excess bronchodilator mediation were compared between treatment periods by paired t tests. The null hypothesis, that there were no differences between ipratropium bromide and placebo, was rejected at p < 0.05.

Results
Treatment with ipratropium bromide caused no change in mean PEF (table 1). Eight children had higher morning PEF values while taking ipratropium bromide, but another eight had higher morning PEF values while taking placebo.

HISTAMINE CHALLENGE
Treatment with ipratropium bromide was associated with significantly higher mean PD20 values (geometric mean 0.78 v0.49 mg/ml, p < 0.01) (figure).

DIURNAL VARIATION IN AIRWAY CALIBRE
Seventy four per cent (23 of 31) of subjects showed
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Table 1  Group mean peak flow rates

<table>
<thead>
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<th>Mean (SD)</th>
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<tr>
<td>Baseline</td>
<td></td>
</tr>
<tr>
<td>Morning</td>
<td>309 (82)</td>
</tr>
<tr>
<td>Afternoon</td>
<td>332 (79)</td>
</tr>
<tr>
<td>Evening</td>
<td>320 (81)</td>
</tr>
<tr>
<td>Ipratropium bromide</td>
<td></td>
</tr>
<tr>
<td>Morning</td>
<td>305 (94)</td>
</tr>
<tr>
<td>Afternoon</td>
<td>325 (89)</td>
</tr>
<tr>
<td>Evening</td>
<td>325 (90)</td>
</tr>
<tr>
<td>Placebo</td>
<td></td>
</tr>
<tr>
<td>Morning</td>
<td>309 (92)</td>
</tr>
<tr>
<td>Afternoon</td>
<td>325 (85)</td>
</tr>
<tr>
<td>Evening</td>
<td>325 (87)</td>
</tr>
</tbody>
</table>

Significant diurnal variation (Dunnett's paired t test) during the baseline period. The mean (SD) amplitude of the diurnal variation in PEF was 23.1% (13.7%). Treatment with ipratropium bromide did not affect the diurnal variation in PEF. Table 1 shows group mean PEF data for each treatment period. Seventy seven per cent (24 of 31) of subjects had significant diurnal variations while taking ipratropium bromide compared with 68% (21 of 31) taking placebo. When the eight children who had higher morning PEF recordings were examined separately there was no relation between the increase in PEF and the diurnal variation in PEF. There was no difference between the mean (SD) amplitude of the diurnal variation in PEF on ipratropium bromide (mean 22.9% (13.7)) and placebo (mean 22.1%, (16.2)).

Diurnal variation in bronchodilator responsiveness

Twenty one children inhaled salbutamol in constant dosage, regularly three times a day. Fifty seven per cent (12 of 21) had significantly greater bronchodilatation in the morning during both ipratropium bromide and placebo periods (p < 0.05). Treatment with ipratropium bromide did not alter the amplitude of the diurnal variation in bronchodilator responsiveness (mean amplitude = 63.5% (range 13–138%) during placebo period; 62.4% (range 12–145%) during ipratropium bromide period). The absolute increases in PEF after salbutamol were not altered by treatment with ipratropium bromide. Mean (SD) increases in PEF (l/min) for the morning, afternoon, and evening were 36 (19.7), 22 (9.7), and 29 (13.5) respectively for the placebo period, and 34 (19.2), 22 (11.9), and 29 (16.2) respectively for the ipratropium bromide period.

Diary cards

There were no differences in symptom scores or additional doses of salbutamol during the baseline, ipratropium bromide, or placebo periods (table 2). No child was able to reduce maintenance treatment while taking ipratropium bromide.

Discussion

Treatment with ipratropium bromide (40 µg) caused no change in mean PEF values in the 31 children who took part in this study, and it failed to modify the diurnal variation in airway calibre or in bronchodilator responsiveness to salbutamol. Nor did it reduce symptom scores or decrease the use of additional doses of salbutamol.

Table 2  Diary card information

<table>
<thead>
<tr>
<th></th>
<th>Symptom scores</th>
<th>Additional doses of salbutamol</th>
</tr>
</thead>
<tbody>
<tr>
<td>Baseline</td>
<td>35 (34-2)</td>
<td>14 (22-1)</td>
</tr>
<tr>
<td>Ipratropium bromide</td>
<td>37 (46-7)</td>
<td>19 (35-3)</td>
</tr>
<tr>
<td>Placebo</td>
<td>32 (51-0)</td>
<td>13 (29-1)</td>
</tr>
</tbody>
</table>

Results are group means (SD) of the total reported for each study period by each subject.

Results of histamine challenge tests (ln PD_{20}) after the ipratropium bromide and placebo periods for each subject; p < 0.05.
In previous reports it has been suggested that ipratropium bromide may decrease the diurnal variation in PEF. The results of this study do not support this suggestion. The reasons for the difference between the results of this study and those in previous reports are not obvious. Mann and Hiller, who also studied children, used the same dosage regimen of ipratropium bromide as we did, and measurements were also made three times a day. They did not examine the diurnal variation in PEF directly but they found significantly higher mean PEF values in the morning with ipratropium bromide. Evening PEF recordings were also higher with ipratropium bromide but the increase was not significant. In the present study the effect of ipratropium bromide was assessed in each subject by comparing morning and afternoon PEF values each day using Dunnett’s paired t-test. Mann and Hiller appear to have used a paired t-test on the group as a whole, comparing only the mean value of morning and evening PEF for each subject. In the present study eight children had significantly higher morning PEF values when taking ipratropium but only four of these had lower amplitudes of the diurnal variation in PEF while taking ipratropium. This suggests that simply comparing mean morning and evening (or afternoon) PEF values on and off a treatment for a group as a whole may not be the best way to evaluate the effect of that medication on the diurnal variation in PEF.

Maintenance treatment with clinically acceptable doses of ipratropium bromide did not result in clinical improvement in the children in this study. Most reported symptoms of asthma during the study and most required additional doses of salbutamol. Ipratropium bromide treatment did not reduce symptom scores or result in the use of fewer additional doses of salbutamol. These results, coupled with the failure of ipratropium bromide to decrease the amplitude of the diurnal variation in airway calibre or in bronchodilator responsiveness, do not support the use of ipratropium bromide as an additional treatment for children with asthma. The subjects in this study were clinically stable, and it is possible that a different result may have been found in children with unstable asthma.

The failure of ipratropium bromide to modify the amplitudes of the diurnal variations in airway calibre or in bronchodilator responsiveness suggests that these rhythms are not vagally mediated. Ipratropium bromide can cause vagal efferent blockade in man. A dose of 40 μg three times daily has been reported to cause bronchodilatation in children and to blunt the response to experimental bronchoconstriction. The bronchodilator effect of ipratropium bromide peaks one hour after inhalation and virtually disappears within six hours. The failure of ipratropium bromide to diminish the diurnal variation in airway calibre and in bronchodilator responsiveness may be because its anticholinergic action does not last long enough to be effective in an eight-hourly regimen. The small but significant increase in PD20 seen after treatment with ipratropium bromide, despite an eight to 12-hour gap between the last dose of ipratropium bromide and the histamine challenge test, suggests, however, that the anticholinergic action of ipratropium bromide may last longer than its detectable bronchodilator action.

References