Comparison of nebulised salbutamol and ipratropium bromide with salbutamol alone in the treatment of chronic obstructive pulmonary disease

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Abstract

Background – Patients admitted with acute exacerbation of chronic obstructive pulmonary disease (COPD) are often prescribed ipratropium bromide in combination with a β_2 agonist such as salbutamol. Studies have not shown any benefit in adding ipratropium bromide to salbutamol in acute exacerbations of COPD, but these studies have only assessed patients for 60–90 minutes and short term studies may not predict long term clinical response. Combination therapy with the two drugs was compared with salbutamol alone in the treatment of acute exacerbations of COPD during a hospital admission.

Methods – Seventy patients admitted to hospital with an acute exacerbation of COPD were randomly allocated to receive either nebulised salbutamol 5 mg and ipratropium bromide 500 μ g, or nebulised salbutamol 5 mg alone (all four times a day) on admission. All other treatment was prescribed at the discretion of the attending physician. Length of stay in hospital and spirometric values on days 1, 3, 7, 14, and discharge were assessed. Patients completed a subjective symptom score each day.

Results – There was no difference between the two groups in the mean (SD) length of stay (salbutamol 10.5 (4.7) days, salbutamol + ipratropium bromide 11.8 (4.4) days; 95% CI –1.02 to 3.62). There was no difference in spirometric values on days 1, 3, 7, 14, or discharge between the two groups. The subjective improvement was similar with both treatments.

Conclusions – The routine addition of nebulised ipratropium bromide to salbutamol appears to be of no benefit in the treatment of acute exacerbations of COPD.

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Keywords: chronic obstructive pulmonary disease, β_2 agonist, ipratropium bromide.

Patients admitted to hospital with an acute exacerbation of chronic obstructive pulmonary disease (COPD) are usually treated with a high dose β_2 agonist as part of their therapy. It is now commonplace for patients to receive, in addition, nebulised ipratropium bromide with

obvious additional cost. This is logical pharmacologically as ipratropium has a different mode of action, inhibiting vagally-mediated bronchomotor tone.¹ There is evidence that adding ipratropium bromide to a nebulised β_2 agonist is valuable in the long term management of COPD^{2–4} and in acute severe asthma.⁵ There have been no studies, however, on the place of ipratropium bromide in the management of acute exacerbations of COPD. Additionally, it has been known for some years that atropine-like drugs may be more effective in remissions of airways obstruction than in relapse.^{6,7} We have therefore compared nebulised salbutamol with nebulised salbutamol plus ipratropium bromide (combination therapy) in the treatment of acute exacerbations of COPD in a randomised trial.

Methods

Patients admitted as emergencies to acute medical units with a diagnosis of an acute exacerbation of COPD who were not taking regular nebulised bronchodilators at home were eligible for the study. All were aged over 45 years and had a smoking history of more than 10 pack years. All had a forced expiratory volume in one second (FEV₁) of <65% predicted when well, and a history of exertional dyspnoea resulting from respiratory disease for over three years. The diagnosis of non-asthmatic COPD was made previously by a consultant respiratory physician. Exclusion criteria included a history suggestive of asthma (childhood respiratory disease, atopy, night time wheezing) and a peripheral eosinophilia of >10%. Patients with a >20% (at least 200 ml) reversibility of FEV₁ to 400 μ g of inhaled salbutamol on the day of discharge were also excluded.

On admission patients were randomised to receive either nebulised salbutamol 5 mg four times daily or salbutamol 5 mg plus ipratropium bromide 500 μ g four times daily. The drugs were administered by an air driven nebuliser (Bard Inspiron Mini-Neb) at a flow rate of 8 l/min until the chamber was dry. The combination therapy was given as a mixture. All other medication was prescribed at the attending physician's discretion.

Spirometric values (best of three attempts using a Vitalograph dry wedge spirometer by one of two experienced operators) were measured before the 18.00 hours nebuliser treat-

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How is your shortness of breath today compared with yesterday?	
Better	+ 1
Same	0
Worse	- 1
Please put the appropriate number next to each day.	
Day 1	
2	
3	
4	
5	
6	
7	
... etc	

Figure 1 Subjective symptom assessment.

ment on days 1, 3, 7, and then weekly and on the day of discharge. Thus, patients admitted before 18.00 hours had spirometric parameters measured on the day of admission whilst those admitted after this time were measured at 18.00 hours the following day (all classified as day 1). In all cases the time of admission was noted. A simple subjective symptom score (fig 1) was recorded daily. The patients were asked whether they felt better, worse, or the same as the previous day.

The change in FEV₁ and forced vital capacity (FVC) from day 1 to days 3, 7, 14, and day of discharge was calculated. The differences between each treatment group and the symptom scores were analysed by the Student's unpaired *t* test. The sample size in this study would detect a 280 ml change in FEV₁ and 500 ml change in FVC with 80% power and 95% confidence.

Verbal consent was obtained from all patients. The study was approved by the clinical research ethics committees of the two health districts involved. Patients were blind to the medication

they received, and the doctors who performed the spirometric measurements were not aware of the treatment group of each patient.

Results

Seventy patients entered the study. Three withdrew (one from the salbutamol group and two from the combination group), two were excluded as they had reversibility of >20% (one from each group), and one patient from the salbutamol group was withdrawn as the reason for admission was newly diagnosed bronchial carcinoma. Two patients were withdrawn due to side effects of the nebulised drugs; one from the salbutamol only group developed chest pain and one in the combination therapy group developed wheezing which resolved on discontinuation of ipratropium. The remaining 62 patients were suitable for analysis. Three patients died during the course of their admission, one in the salbutamol group and two receiving combination therapy. Two patients prescribed ipratropium developed urinary obstruction but this did not necessitate their withdrawal from the study. Table 1 shows the baseline characteristics of the two treatment groups. The patients were well matched for all the parameters (Student's unpaired *t* test). Day 1 spirometric values were always measured at 18.00 hours and therefore at a variable time from admission, but always within 24 hours. There was, however, no difference in the time of admission between the two groups (table 1).

The length of hospital stay and duration of nebuliser therapy was similar in both groups. There was no significant difference in the quantity of intravenous hydrocortisone, aminophylline and oral antibiotics received by the two groups (table 2).

Patient progress was monitored by FEV₁ and FVC (figs 2 and 3) and the change in these parameters between days 1, 3, 7, 14, and discharge was calculated. There was no significant

Table 1 Mean (SD) baseline characteristics of the two treatment groups

	S (n=33)	S+IB (n=29)	p	95% CI of difference
Age (years)	70.4 (9.1)	67.8 (6.7)	NS	-6.71 to 1.51
Height (cm)	163.8 (9.7)	165.0 (9.6)	NS	-3.71 to 6.11
Weight (kg)	65.0 (16.9)	65.3 (15.7)	NS	-8.02 to 8.62
Years breathless	10.6 (11.3)	16.5 (12.9)	NS	-0.25 to 12.04
Smoking (pack years)	48.8 (32.3)	55.0 (32.1)	NS	-10.20 to 22.60
Pao ₂ * (kPa)	8.5 (2.4)	7.8 (1.6)	NS	-1.75 to 0.35
Paco ₂ * (kPa)	5.4 (1.3)	5.9 (1.5)	NS	-0.21 to 1.21
Time of admission	13.40 (5.0)	12.50 (5.0)	NS	-3.38 to 1.71
FEV ₁ day 1 (l)	0.77 (0.34)	0.78 (0.41)	NS	-0.18 to 0.20
FVC day 1 (l)	1.47 (0.65)	1.55 (0.71)	NS	-0.27 to 0.43

S=salbutamol alone; S+IB=combination therapy; Pao₂, Paco₂=arterial oxygen and carbon dioxide tensions; FEV₁=forced expiratory volume in one second; FVC=forced vital capacity.

* On admission.

Table 2 Mean (SD) length of hospital stay and drug therapy of the two treatment groups

	S (n=33)	S+IB (n=29)	p	95% CI of difference
Days to discharge	10.5 (4.7)	11.8 (4.4)	NS	-1.02 to 3.62
Days on nebulisers	8.5 (4.2)	8.2 (3.6)	NS	-2.30 to 1.70
Days on iv aminophylline	0.1 (0.5)	0.6 (1.4)	NS	-0.02 to 1.02
Days on iv steroids	0.1 (0.3)	0.6 (1.4)	0.05	-0.005 to 1.0
Days on antibiotics	5.8 (4.3)	5.7 (3.3)	NS	-2.07 to 1.87

S=salbutamol alone; S+IB=combination therapy.

Table 3 Mean (SD) changes in forced expiratory volume in one second (FEV₁) and forced vital capacity (FVC) over time

	S (n=33)	S+IB (n=29)	p	95% CI of difference
FEV ₁				
Day 3 - Day 1	0.17 (0.28)	0.05 (0.24)	NS	-0.25 to 1.34
Day 7 - Day 1	0.21 (0.42)	0.15 (0.26)	NS	-0.24 to 0.12
Day 14 - Day 1	0.06 (0.11)†	0.26 (0.29)*	NS	-0.02 to 0.42
Day of discharge - Day 1	0.23 (0.32)	0.15 (0.32)	NS	-0.24 to 0.08
FVC				
Day 3 - Day 1	0.25 (0.42)	0.04 (0.41)	0.05	-0.42 to 0.001
Day 7 - Day 1	0.39 (0.52)	0.17 (0.45)	NS	-0.47 to 0.03
Day 14 - Day 1	0.33 (0.23)†	0.62 (0.51)*	NS	-0.10 to 0.68
Day of discharge - Day 1	0.56 (0.47)	0.42 (0.61)	NS	-0.41 to 0.13

* 10 patients, † 9 patients.

S=salbutamol alone, S+IB=combination therapy.

Table 4 Mean (SD) subjective improvement over hospital stay

	S (n=33)	S+IB (n=29)	p	95% CI of difference
Number of days better	4.5 (1.8)	4.8 (2.4)	NS	-0.77 to 1.37
Number of days same	3.9 (2.8)	4.7 (2.7)	NS	-0.60 to 2.20
Number of days worse	1.2 (1.4)	1.3 (1.5)	NS	-0.64 to 0.84

S=salbutamol alone; S+IB=combination therapy.

difference between the two groups (table 3), and no difference was observed in the subjective improvement between the two groups (table 4).

Figure 2 Mean (SE) FEV₁ over admission.

Figure 3 Mean (SE) FVC over admission.

Discussion

The combination of salbutamol and ipratropium bromide appeared to give no additional benefit compared with salbutamol alone during the routine inpatient treatment of an acute exacerbation of COPD. No differences were observed in spirometric values, subjective symptom scores, duration of hospital stay, or number of days on a nebuliser between the two groups. This is in contrast to findings in stable COPD although results are not consistent. Many trials in stable outpatients with COPD have shown combination therapy to be beneficial,⁸⁻¹² although some suggest that any improvement is small.^{9,13,14} Furthermore, some workers have shown that ipratropium bromide alone is as effective as either salbutamol alone or combination therapy.¹⁵⁻¹⁷

There are far fewer studies on acute exacerbations of COPD. In a study by O'Driscoll *et al*² combination therapy provided no additional benefit to single agent therapy in the first hour. An earlier study¹⁸ found that either fenoterol or ipratropium bromide was as effective as the two agents combined in producing bronchodilation over a 90 minute period. These studies have assessed patients with acute exacerbations of COPD over 60-90 minutes, but short term reversibility alone cannot be relied upon to predict longer term clinical response.¹⁹⁻²¹ In assessing therapy it is important to monitor patients with acute exacerbations of COPD throughout a hospital admission and, to our knowledge, this is the first study which compares the use of salbutamol alone with combination therapy for this duration.

It is possible that this study did not have a sufficiently large sample size to detect a small improvement in spirometric values by the addition of ipratropium bromide. This study would detect a difference in mean FEV₁ of 280 ml at each point with 80% power and we feel that a difference of less than this magnitude is of doubtful clinical significance. We consider that the dose of ipratropium bromide used was sufficient because a previous study has shown that 500 µg is in excess of that required to

achieve maximal acute reversibility in stable COPD.²²

As in other studies, we considered changes in FEV₁ and FVC to be more objective than serial peak flow measurements in measuring bronchodilation. It is well recognised that there is only limited correlation between spirometric values and subjective sensations of dyspnoea and exercise tolerance in COPD.²³⁻²⁵ There are a number of validated multidimensional questionnaires to assess wellbeing in patients with COPD.²⁶ These, however, have mainly been used in patients with stable COPD and the few questionnaires designed to respond to rapid changes in disease activity are time consuming to complete.²⁷ We therefore confined ourselves to a single measure of subjective benefit similar to that used in other studies assessing nebuliser therapy.²⁸

Although ipratropium bromide is generally well tolerated and safe, adverse (anticholinergic) effects do occur. The addition of ipratropium bromide to a β_2 agonist in the treatment of acute COPD will therefore increase the possibility of side effects as well as the cost of therapy.

We conclude that the addition of ipratropium to salbutamol confers no benefit in the routine management of hospital inpatients with acute exacerbations of COPD. We consider that these results apply only to acute exacerbations and the situation with respect to long term domiciliary treatment is different. In the treatment of an acute exacerbation of COPD it would seem reasonable to advise adding nebulised ipratropium bromide to a β_2 agonist only when a patient is not progressing satisfactorily. This is now our unit policy.

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- 1 Gross NJ, Skorodin MS. Anticholinergic, antimuscarinic bronchodilators. *Am Rev Respir Dis* 1984;129:856-70.
- 2 O'Driscoll BR, Horsley MG, Taylor RJ, Chambers DK, Bernstein A. Nebulised salbutamol with and without ipratropium bromide in acute airflow obstruction. *Lancet* 1989;i:1418-20.
- 3 Morrison JFJ, Jones PC, Muers MFM. Assessing physiological benefit from domiciliary nebulised bronchodilators in severe airflow limitation. *Eur Respir J* 1992;5:424-9.
- 4 Goldman JM, Teale C, Muers MFM. Simplifying the assessment of patients with chronic airflow limitation for home nebuliser therapy. *Respir Med* 1992;86:33-8.

- 5 Ward MF, Fentem PH, Roderick Smith WH, Davies D. Ipratropium bromide in acute asthma. *BMJ* 1981;282:598-600.
- 6 Altounyan REC. Variation of drug action on airway obstruction in man. *Thorax* 1964;19:406-15.
- 7 Teale C, Morrison JFJ, Muers MFM, Pearson SB. Response to nebulised ipratropium bromide and terbutaline in acute severe asthma. *Respir Med* 1992;86:215-8.
- 8 Brown IG, Chan CS, Kelly CA, Dent AG, Zimmerman PV. Assessment of the clinical usefulness of nebulised ipratropium bromide in patients with chronic airflow limitation. *Thorax* 1984;39:272-6.
- 9 Hughes JA, Tobin MJ, Bellamy D, Hutchinson DCS. Effects of ipratropium bromide and fenoterol aerosols in pulmonary emphysema. *Thorax* 1982;37:667-70.
- 10 Gross NJ, Skorodin MS. Role of the parasympathetic system in airway obstruction due to emphysema. *N Engl J Med* 1984;311:421-5.
- 11 Crane J, Gamble S, Purdie G. Comparison of a fenoterol/ipratropium combination with salbutamol form metered dose inhalers in subjects with chronic partially reversible airways obstruction. *NZ Med J* 1987;100:385-7.
- 12 Rudolf M. Combination therapy - a review of clinical studies. *Postgrad Med J* 1984;60:(Suppl) 9-12.
- 13 Duvent - is logic enough? *Drug Ther Bull* 1985;23:3-4.
- 14 Connolly CK, Chan NS. Salbutamol and ipratropium in partially reversible airway obstruction. *Br J Dis Chest* 1987;81:55-61.
- 15 Braun SR, McKenzie WJ, Copeland C, Knight L, Ellersieck M. A comparison of the effect of ipratropium and albuterol in the treatment of chronic obstructive airways disease. *Arch Intern Med* 1989;149:544-7.
- 16 Easton PA, Jadue C, Dhingra S, Anthonisen NR. A comparison of the bronchodilating effects of a beta-2 adrenergic agent (albuterol) and an anticholinergic agent (ipratropium bromide), given by aerosol alone or in sequence. *N Engl J Med* 1986;315:735-9.
- 17 Karpel JP. Bronchodilator response to anticholinergic and beta-adrenergic agents in acute and stable COPD. *Chest* 1991;99:871-6.
- 18 Rebuck AS, Chapman KR, Abboud R, Pare PD, Kreisman H, Wolkove N, et al. Nebulized airways disease in the emergency room. *Am J Med* 1987;82:59-64.
- 19 Pratter MR, Irwin RS. Predicting response to bronchodilator therapy in chronic obstructive pulmonary disease. *Arch Intern Med* 1988;148:1909-10.
- 20 Taylor DR, Buick B, Kinney C, Lowry RC, McDevitt DG. The efficacy of orally administered theophylline, inhaled salbutamol, and a combination of the two as chronic therapy in the management of chronic bronchitis with reversible airflow obstruction. *Am Rev Respir Dis* 1985;131:747-51.
- 21 Teale C, Morrison JFJ, Jones PC, Muers MFM. Reversibility tests in chronic obstructive airways disease: their predictive value with reference to benefit from domiciliary nebuliser therapy. *Respir Med* 1991;85:281-4.
- 22 Jenkins CR, Chow CM, Fisher BL, Marlin GE. Comparison of ipratropium bromide and salbutamol by aerosolized solution. *Aust NZ J Med* 1981;11:513-6.
- 23 Mahler DA, Weinberg DH, Wells CK, Feinstein AR. The measurement of dyspnea. *Chest* 1985;85:751-8.
- 24 Pineda H, Haas F, Axen K, Haas A. Accuracy of pulmonary function tests in predicting exercise tolerance in chronic obstructive pulmonary disease. *Chest* 1984;86:564-7.
- 25 Levy SF. Bronchodilators in COPD. To the max. *Chest* 1991;99:793.
- 26 Guyatt GH, Berman LB, Townsend M, Pugsley SO, Chambers LW. A measure of quality life for clinical trials in chronic lung disease. *Thorax* 1987;42:773-8.
- 27 Jones PW, Quirk FH, Baveystock CM, Littlejohns P. A self-complete measure of health status for chronic airflow limitation. The St George's Respiratory Questionnaire. *Am Rev Respir Dis* 1992;145:1321-7.
- 28 O'Driscoll BR, Kay EA, Taylor RJ, Weatherby H, Chetty MCP, Bernstein A. A longterm prospective assessment of home nebuliser treatment. *Respir Med* 1992;86:317-25.