Comparison of almitrine bismesylate and medroxyprogesterone acetate on oxygenation during wakefulness and sleep in patients with chronic obstructive lung disease

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Abstract

The effects of almitrine bismesylate and medroxyprogesterone acetate on oxygenation during wakefulness and sleep were compared in six patients with chronic obstructive lung disease and carbon dioxide retention. Patients received 1.5 mg/kg almitrine (a peripheral chemoreceptor stimulant), 100 mg of medroxyprogesterone (a central respiratory stimulant), or matched placebo daily for 15 days in random order in a crossover trial. When subjects were awake almitrine increased the ventilatory response to hypoxia and increased arterial oxygen tension (PaO_2) to a greater extent than medroxyprogesterone, whereas medroxyprogesterone augmented the ventilatory response to hypercapnia and decreased arterial carbon dioxide tension (PaCO_2) to a greater extent than almitrine. Neither drug influenced sleep architecture significantly, except that medroxyprogesterone increased the number of arousals. Almitrine had a more favourable effect than placebo on oxygenation as estimated from arterial oxygen saturation (SaO_2) during the different stages of sleep, the number of episodes of hypoxaemia, and the amount of time that SaO_2 was below 80%. The only change with medroxyprogesterone by comparison with placebo was a decrease in the number of hypoxaemic episodes. It is concluded that both active drugs improved blood gases during wakefulness, but that 1.5 mg/kg of almitrine is superior to 100 mg of medroxyprogesterone in improving SaO_2 during sleep.

During the last few years there has been renewed interest in the use of respiratory stimulants to alleviate hypoxaemia in patients with chronic obstructive pulmonary disease during wakefulness and sleep. Some of the factors responsible for this renewed interest are the recognition that normoxaemic patients with chronic obstructive pulmonary disease may have hypoxaemic episodes during sleep¹; the development of respiratory stimulants for long term oral use with few side effects; the reluctance of some patients to have long term oxygen therapy; and the various practical problems associated with oxygen therapy.

The most commonly used respiratory stimulants are medroxyprogesterone acetate and almitrine bismesylate. Although the two drugs act by different mechanisms, both have been shown to improve blood gases during wakefulness and sleep.²⁻⁴ Medroxyprogesterone, which is considered to be a central chemoreceptor stimulant, increases alveolar ventilation during wakefulness and sleep.³ Almitrine, a peripheral chemoreceptor stimulant, increases alveolar ventilation and improves the ventilation-perfusion ($\dot{V}/\dot{Q}$) ratio.⁵

The aim of this study was to compare the effects of medroxyprogesterone and almitrine on oxygenation during wakefulness and sleep in patients with chronic obstructive lung disease and carbon dioxide retention. This is, we believe, the first direct comparison of these agents in the same individuals.

Methods

PATIENTS

We studied six non-obese men (mean age 62 (SD 6) years) with chronic obstructive lung disease. Their respiratory function and blood gas values are shown in table 1. All were clinically stable for more than four weeks before and throughout the study. None was receiving long term domiciliary oxygen therapy. They continued their usual medications (inhaled β_2 sympathomimetics, theophylline derivatives, diuretics, digoxin) in unchanged dosage throughout the study.

STUDY DESIGN

The study was designed as a placebo controlled, crossover trial. After a selection night to exclude patients with the sleep apnoea syndrome and for purposes of familiarisation, subjects received in random order 1.5 mg/kg

Table 1 Mean (SD) values for respiratory function measures and arterial blood gases

Pulmonary function (% pred)	Arterial blood gases
FEV ₁ 35 (11)	PaO_2 51 (3) mm Hg
VC 51 (4)	PaCO_2 50 (4) mm Hg
RV 180 (18)	pH 7.41 (0.02)
MMER 13 (5)	SaO_2 83 (3) %

Conversion to SI units: 1 mm Hg $\approx$ 0.133 kPa.

FEV₁—forced expiratory volume in one second; VC—vital capacity; RV—residual volume; MMER—maximum mid expiratory flow (25–75% VC); PaO_2 —arterial oxygen tension; PaCO_2 —arterial carbon dioxide tension; SaO_2 —arterial oxygen saturation. 1 mm Hg $\approx$ 0.133 kPa.

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almitrine a day divided into two doses, 100 mg of medroxyprogesterone a day divided into two doses, or matched placebo for 15 days. Drugs and placebo were administered at 0800 and 2000 hours. An acclimatisation night in the sleep laboratory preceded every sleep study. Each period of study was separated by a four week washout period.

The protocol was approved by the ethical committee of the department of internal medicine and each patient gave written, informed consent.

AWAKE STUDY

The study procedures were carried out in the morning, one hour after administration of the drugs.

Arterial blood samples were taken at rest with the patient in the supine posture through a small cannula inserted in the radial artery. Blood was analysed for arterial oxygen tension (P_{aO_2}) and arterial carbon dioxide tension (P_{aCO_2}) by electrodes (Radiometer, Copenhagen). The alveolar-arterial oxygen difference ($A-aDO_2$) was computed according to the simplified formula

$$P_{aO_2} = P_{iO_2} - 1.25 \cdot P_{aCO_2}.$$

Spirometry was performed with a water sealed Godart spirometer. Lung volumes were measured by the helium dilution method.⁶

The ventilatory response to hypercapnia was estimated by the rebreathing method of Read.⁷ The ventilatory response to hypoxia was assessed by the use of progressive hypoxia with progressive hypercapnia.⁸

SLEEP STUDY

All patients underwent a whole night polysomnographic study with simultaneous recording of the electroencephalogram, elec-

tro-oculogram, and submental electromyogram.⁹ Air flow was detected by continuously analysing for carbon dioxide (Godart, mark III) at the nose and mouth through a common catheter. Rib cage and abdominal motion was monitored by respiratory inductance plethysmography with two bands (Respirace Jacket, Ambulatory Monitoring, White Plains, New York). Oxygen saturation was monitored by a Hewlett Packard (47201A) ear oximeter. All variables were recorded on an eight channel recorder (Alval) running at 15 mm/s.

Baseline SAO_2 when the patient was awake was defined as the mean SAO_2 over a period of 10 minutes before sleep; a hypoxaemic episode was defined as a fall in SAO_2 of 10% or more below baseline SAO_2 lasting for at least one minute.

STATISTICAL ANALYSIS

The data were analysed by comparing the values obtained after administration of placebo, medroxyprogesterone, and almitrine with the paired *t* test. Values of *p* of 0.05 or less were considered significant.

Results

AWAKE STUDY

Both medroxyprogesterone and almitrine had a favourable effect on arterial blood gas tensions during wakefulness without any change in maximum expiratory flow or lung volumes.

Almitrine had the greater effect on P_{aO_2} and this was significantly different from that seen with medroxyprogesterone (table 2). On the other hand, medroxyprogesterone caused the greater reduction in P_{aCO_2} and this was significantly greater than the reduction seen after almitrine.

There was an increase in the calculated alveolar-arterial oxygen difference ($A-aDO_2$) after medroxyprogesterone (table 2).

The ventilatory response to hypercapnia was increased after medroxyprogesterone and almost unchanged after almitrine administration (table 2). The ventilatory response to hypoxia increased greatly after almitrine and to a lesser degree after medroxyprogesterone administration.

SLEEP STUDY

There were no differences in the percentage of each sleep stage seen on the different study nights except that the number of arousals was increased after medroxyprogesterone (*p* < 0.001, table 3). Sleep architecture was unchanged, as judged by the duration of sleep stages expressed as a percentage of sleep period time (figure).

Mean SAO_2 was increased during REM and non-REM sleep after medroxyprogesterone and to a greater extent after almitrine. SAO_2 was significantly higher after almitrine than after placebo (table 4). The lowest SAO_2 recorded during sleep, the time for which SAO_2 fell below 80%, and the number of hypoxaemic episodes were less after administration of the active drugs, almitrine causing a greater

Table 2 Mean (SD) values for blood gas tensions, alveolar-arterial oxygen difference ($A-aDO_2$), and ventilatory response to hypoxia and hypercapnia after administration of placebo, medroxyprogesterone, and almitrine in the six patients

	Placebo	Medroxyprogesterone	Almitrine	<i>p</i> *
P_{aO_2} (mm Hg)	53 (2)	56 (3)	58 (2)	<0.05
P_{aCO_2} (mm Hg)	52 (4)	44 (2)	47 (2)	<0.01
$A-aDO_2$ (mm Hg)	31 (3)	38 (4)	33 (4)	<0.05
Hypercapnic ventilatory response (l/min/mm Hg)	0.93 (0.5)	1.19 (0.6)	0.94 (0.5)	<0.05
Hypoxic ventilatory response $\Delta \dot{V}_{E_{79}}$ (l/min)	6.67 (5.1)	7.43 (4.2)	9.68 (4.2)	<0.05

*Comparison between medroxyprogesterone and almitrine; the paired *t* test was used with the absolute values.

Conversion to SI units: 1 mm Hg $\approx$ 0.133 kPa.

$\Delta \dot{V}_{E_{79}}$ —change in minute ventilation when SAO_2 is 79%.

Other abbreviations as in table 1.

Table 3 Data on sleep quality (mean (SD) values)

	Placebo	Medroxyprogesterone	Almitrine	<i>p</i> *
Time in bed (min)	308 (49)	324 (35)	316 (29)	NS
Sleep latency (min)	24 (11)	17 (13)	21 (14)	NS
Sleep period time (min)	285 (24)	307 (31)	295 (18)	NS
Total sleep time (min)	218 (15)	251 (38)	229 (22)	NS
Sleep efficiency index (%)	71 (6)	77 (12)	76 (9)	NS
Arousals/hour	17 (4)	25 (3)	18 (4)	<0.001*

*Comparison between placebo and medroxyprogesterone; the paired *t* test was used.

Mean (SD) duration of sleep stages expressed as percentages of sleep period time after placebo □, medroxyprogesterone ▨, and almitrine ▤ administration.

improvement in oxygen saturation than medroxyprogesterone (table 5).

We calculated the probable mean fall in P_{aO_2} (ΔP_{aO_2}) from P_{aO_2} values when the patients were awake, using mean SaO_2 during stage 2 and the oxyhaemoglobin dissociation curve and assuming that pH values, body temperature, and P_{50} (half oxygenation tension of haemoglobin) were unchanged during stage 2 sleep (table 6). The calculated ΔP_{aO_2} was lower during almitrine treatment than with medroxyprogesterone or placebo, though the difference was not significant.

Discussion

Patients with chronic obstructive pulmonary disease and carbon dioxide retention usually have normal respiratory drive and normal minute ventilation.¹⁰ Carbon dioxide retention is caused by insufficient respiratory effort and an abnormal breathing pattern in the presence of serious disturbances of the distribution of lung perfusion.¹¹ Respiratory stimulants with a beneficial effect on the pattern of breathing may

have a place in therapeutic management of these patients.

Non-apnoeic oxyhaemoglobin desaturation is known to occur in patients with chronic obstructive lung disease, mainly in association with REM sleep, and this may have deleterious consequences for the prognosis of these patients. Low P_{aO_2} and high P_{aCO_2} during wakefulness are predictors of deterioration of hypoxaemia during sleep.¹ The principal causative mechanisms for nocturnal desaturation are alveolar hypoventilation and an increase in the degree of $\dot{V}/\dot{Q}$ disturbance.^{12,13}

Although the exact mechanism of respiratory stimulation caused by medroxyprogesterone is still obscure, it is believed to operate directly on the respiratory centre after crossing the blood-brain barrier.¹⁴ Medroxyprogesterone increases respiratory drive,¹¹ the ventilatory and neuromuscular response to hypercapnia,¹⁵ and the ventilatory response to hypoxia,¹⁶ and it restores the impaired load compensation in patients with chronic obstructive lung disease.¹⁷ When the patients react favourably to medroxyprogesterone, an increase in their tidal volume and mean inspiratory flow is observed that persists during sleep.³ According to Skatrud,¹⁸ all our patients can be defined as "correctors" as they showed a greater than 5 mm Hg (0.67 kPa) decrease in P_{aCO_2} . The remarkable increase in A-a $\dot{V}O_2$ in our patients after medroxyprogesterone shows that the increase in ventilation is not accompanied by an appropriate increase in their P_{aO_2} . To interpret this we would need detailed information about pattern of breathing, cardiac output, and $\dot{V}/\dot{Q}$ distribution.

On the other hand, almitrine stimulates peripheral chemoreceptors,¹⁹ increases the ventilatory response to hypoxaemia²⁰ and alveolar ventilation,²¹ and has a beneficial effect on the ventilation-perfusion relationship.²² The changes in the pattern of breathing seen with almitrine consist of a faster, deeper, and shorter inspiration followed by a longer expiration without any change in breathing frequency.²³ The significant improvement in arterial blood gas tensions without a significant change in A-a $\dot{V}O_2$ after almitrine suggests some normalisation of $\dot{V}/\dot{Q}$ matching in conjunction with the increase in ventilation (otherwise A-a $\dot{V}O_2$ would be expected to have fallen). Almitrine has a positive effect on the distribution of ventilation²¹ in addition to redistributing blood flow to areas with better ventilation.²⁴

Both medroxyprogesterone and almitrine have been shown to improve blood oxygenation during wakefulness and sleep. Our comparison of the two shows that almitrine in the doses we used increases P_{aO_2} during wakefulness more than does medroxyprogesterone. Furthermore, almitrine significantly improved oxygenation during sleep by comparison with placebo, whereas the improvement after medroxyprogesterone administration was not significant. The calculated change in arterial oxygen pressure (ΔP_{aO_2}) after almitrine during stage 2 sleep suggests that the improvement in oxygenation with almitrine is not due only to the higher baseline SaO_2 during wakefulness.

Table 4 Mean (SD) values for arterial oxygen saturation (%) during the different sleep stages in the six patients

	Placebo	Medroxyprogesterone	Almitrine	Placebo-almitrine
Awake	85 (2)*	88 (2)*	89 (2)	<0.001
Stage 1	77 (3)	81 (6)	83 (3)	<0.01
Stage 2	76 (3)	79 (5)	82 (4)	<0.01
Stage 3 + 4	73 (8)	76 (8)	79 (5)	<0.05
REM (n = 5)	61 (9)	66 (9)	68 (11)	<0.01

*p < 0.01 (paired t test).

Table 5 Arterial oxygen saturation (SaO_2) during sleep after administration of placebo, medroxyprogesterone and almitrine

	Placebo	Medroxyprogesterone	Almitrine
Lowest SaO_2 during sleep (%)	41 (19)	45 (23)	55 (13)
Duration (min) SaO_2 < 80%	122 (35)**	85 (48)	62 (22)**
Total No of hypoxaemic episodes	28 (7)*†	19 (8)*	16 (5)†

*p < 0.05, **p < 0.01, †p < 0.001 (paired t test).

Table 6 Mean (SD) arterial oxygen saturation (SaO_2) during wakefulness and stage 2 sleep, the corresponding arterial oxygen tension (PaO_2) calculated from the oxyhaemoglobin dissociation curve, and the fall in PaO_2 (ΔPaO_2) between wakefulness and sleep after placebo, medroxyprogesterone, and almitrine administration in the six patients

	Placebo		Medroxyprogesterone		Almitrine	
	Awake	Sleep stage 2	Awake	Sleep stage 2	Awake	Sleep stage 2
SaO_2 (%)	85 (2)	76 (3)	88 (2)	79 (5)	89 (2)	82 (4)
PaO_2 (mm Hg)	52 (2)	41 (4)**†	56 (3)	44 (8)**	58 (2)	48 (6)†
ΔPaO_2 (mm Hg)		12.1 (4.5)*		11.1 (5.0)		9.7 (6.2)*

*0.1 > p > 0.05, **p < 0.05, †p < 0.01 (paired t test). 1 mm Hg $\approx$ 0.133 kPa.

This is in agreement with the finding of Catterall *et al.*,²⁵ who showed that the level of arterial oxygenation when the patient is awake is not the sole determinant of the degree of nocturnal hypoxaemia.

The basic sleep architecture was not affected significantly by either drug. It is remarkable that the number of arousals increased after medroxyprogesterone administration. Sleep disturbance in patients with chronic obstructive lung disease is multifactorial.²⁶ Hypoxaemia and hypercapnia are two probable causes.²⁷ The large rise in the number of arousals after medroxyprogesterone was possibly the result of the combined augmentation of the chemosensitivity to hypercapnia (mainly) and to hypoxia that was documented during wakefulness.

Although, according to Voelkel,²⁸ "the elusive mechanisms of alveolar hypoxia induced constriction of lung vessels are still one of the vexing problems of respiratory physiology," this mechanism has been claimed to originate, in part at least, from peripheral chemoreceptors²⁹ and in part from the lung.³⁰ Almitrine is known to stimulate peripheral chemoreceptors directly and experimental studies have shown that it causes pulmonary vasoconstriction, producing a local action on the lung.³¹ Almitrine probably enhances the rapid changes in perfusion distribution that follow the uneven changes in the distribution of ventilation during hypoventilation episodes, thus preventing further $\dot{V}/\dot{Q}$ inequalities and securing a higher PaO_2 during sleep.

We conclude that both almitrine and medroxyprogesterone can improve blood gases during wakefulness in patients with chronic obstructive lung disease and carbon dioxide retention. In the doses used, however, almitrine seemed somewhat more effective in improving nocturnal oxygen saturation.

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