Induction of sleep apnoea with negative pressure ventilation in patients with chronic obstructive lung disease

Robert D Levy, Manuel G Cosio, Laurie Gibbons, Peter T Macklem, James G Martin

Abstract

Background Negative pressure ventilation provides intermittent non-invasive ventilatory assistance for patients with advanced chronic obstructive lung disease. Upper airway obstruction during sleep, a reported complication of the technique, may, however, limit its clinical applicability.

Methods The effects of nocturnal negative pressure ventilation on ventilation and on indices of sleep quality were investigated in five patients with severe chronic obstructive lung disease (mean (SE) FEV₁, 31% (3%) predicted) who had completed three months of nightly negative pressure ventilation. Subjects underwent overnight polysomnography on consecutive nights, the first night serving as a control and negative pressure ventilation being provided on the second night. Ventilators were adjusted to result in maximum suppression of the peak phasic electromyogram signal from the diaphragm.

Results Negative pressure ventilation resulted in substantial increases in episodes of obstructive apnoea and hypopnoea (mean (SE)/h 59-3 (19-8) v 3-2 (1-3) on control nights). Most obstructive events, however, were associated with under 3% oxygen desaturation, and the lowest recorded values for overnight oxygen saturation were similar on the two study nights. Negative pressure ventilation was also associated with significant increases in the frequencies of movement arousals and changes in sleep stage.

Conclusions Negative pressure ventilation applied during sleep to patients with advanced chronic obstructive lung disease may result in the development of recurrent episodes of apnoea and hypopnoea as well as altered sleep quality, which could limit its clinical applicability.

Negative pressure ventilation has been found to be effective for the provision of intermittent non-invasive assisted ventilation in patients with chronic hypercapnic respiratory failure resulting from neuromuscular disorders and thoracic deformities. Some recent studies have suggested that intermittent application of negative pressure ventilation may result in improvements in ventilatory performance in patients with severe chronic obstructive lung disease, possibly as a result of providing rest for chronically fatigued respiratory muscles.

Several investigators have reported upper airway obstruction to be a complication of negative pressure ventilation in both normal subjects and in patients with neuromuscular disorders. To determine whether this phenomenon could limit the applicability of the technique, we examined the effects of negative pressure ventilation on breathing during sleep and sleep quality in patients with severe chronic obstructive lung disease.

Methods

SUBJECTS

We studied five subjects with severe, stable chronic obstructive lung disease. They were part of a large randomised, prospective clinical trial of ventilatory muscle rest. Details of the study design, protocols for selecting patients, and application of intermittent assisted ventilation have been reported elsewhere. The study protocol was approved by an ethics committee of the Montreal Chest Hospital Centre, McGill University, and informed consent was obtained from the subjects.

Subjects had just completed three months of intermittent assisted ventilation, during which they had been asked to apply the negative pressure ventilation for four to eight hours daily. They had been given the option of undergoing assisted ventilation during waking or sleep periods, or both. Most subjects in the larger trial of ventilatory muscle rest had chosen to use the equipment during the day, stating that use during sleep resulted in limitation of mobility, musculoskeletal discomfort, or a choking sensation. The five subjects enrolled in the present study were the first five to complete the trial who had chosen to use the equipment during sleep and agreed to participate in the study protocol.

NEGATIVE PRESSURE VENTILATION

Negative pressure ventilation was applied with a pressure cycled Thompson Maxivent ventilator (Thompson Respiration Products Inc, Boulder, Colorado) connected by a hose to an airtight pneumosuit (New Tech Medical Specialties, Palisades Park, New Jersey). Subjects were constrained by the equipment to remain in the supine or semi-recumbent position.
To assess the activity of the inspiratory muscles, the electromyogram (EMG) of the diaphragm was recorded with surface electrodes placed over the right sixth and seventh intercostal spaces near the costal margin. The EMG signals were processed by a moving time averager (CWE Inc, Ardmore, Pennsylvania) with a time constant of 100 ms after band pass filtering (0.03–10 KHz) and rectification. Diaphragm EMG signals recorded from surface electrodes have been shown to provide a reasonable approximation to those recorded from a bipolar oesophageal electrode.

The optimal ventilator settings—that is, those that resulted in the maximum reduction of spontaneous inspiratory efforts—were determined at the beginning of the study and were not changed subsequently. The ventilation frequency was set to equal the subject's spontaneous respiratory rate, and the inspiratory time: total time per breath ratio was fixed at 0.4. The peak negative pressure in the pneumosuit was gradually increased until a maximum reduction in the phasic amplitude of the diaphragm EMG signal was observed.

SLEEP STUDIES
Subjects underwent two overnight polysomnographic studies on consecutive nights at the end of the three months of intermittent negative pressure ventilation. The first night served as a control night, during which subjects were fully instrumented and slept in the negative pressure ventilation set up while breathing spontaneously—although the ventilator cycled, it was not connected to the suit. On the second night negative pressure ventilation was applied with the ventilator settings determined at the beginning of the study.

Sleep was staged in 30 second epochs according to standard criteria. An episode of apnoea was defined as at least 10 seconds without airflow and of hypopnoea as 10 seconds with at least a 50% reduction in airflow as measured by thermocouples. Episodes of apnoea and hypopnoea had to be associated with movement arousals in order to be scored as such. Apnoeas and hypopnoeas episodes were divided into those associated with at least a 3% decrement in oxygen saturation (Sao2), and those with less than a 3% reduction. On control nights obstructive apnoea was an episode in which respiratory effort was present, as indicated by thoracoabdominal movements measured by respiratory inductive plethysmography (Respitrace, AM1 Inc, Ardsley, New York) as well as phasic EMG activity of the diaphragm. When no such effort was evident apnoea was considered to be of central origin. Mixed apnoeas contained periods of both central and obstructive apnoea. As few episodes were observed on control nights, mixed and obstructive events were combined and called obstructive. With negative pressure ventilation apnoea was by definition obstructive as respiratory “effort” was provided by the ventilator.

ANALYSIS
The significance of differences in means between control and negative pressure ventilation nights was determined by Wilcoxon's signed rank test. p Values below 0.05 were considered to be statistically significant. All data are reported as means with standard errors of the mean (SE) in parentheses.

Results
SUBJECTS’ CHARACTERISTICS
Five subjects (three male, two female), aged 48–75 years, participated in the study. None was obese (82–103% of predicted ideal body weight). The participants did not have histories suggesting sleep apnoea (excessive snoring, witnessed apnoea, daytime hypersonnolence). They had severe chronic obstructive lung disease with forced expiratory volume in one second (FEV1) 31% (3%) predicted, forced vital capacity (FVC) 48% (6%) predicted, arterial oxygen tension (Pao2) 8.4 (0.4) kPa, arterial carbon dioxide tension (Paco2) 5.3 (0.4) kPa, and pH 7.4 (0.01).

All subjects had completed three months of daily intermittent negative pressure ventilation at home, and had chosen to use it during sleep. Internal clocks in the ventilators indicated a mean of 3.8 (1.3) hours of use nightly.

VENTILATORY INDICES
The ventilator rates ranged from 16 to 24 breaths/min, and the peak negative pressures in the pneumosuits from −15 to −30 cm H2O. These settings resulted in reductions in the
peak phasic amplitude of the diaphragm EMGs ranging from 64% to 92% of those recorded during spontaneous breathing when the subjects were awake.

**SLEEP STUDIES**

Subjects had more disrupted sleep on negative pressure ventilation nights than on control nights, as indicated by more movement arousals and changes in sleep stage (table 1). Furthermore, the sleep architecture (figure) was altered, with a reduction in rapid eye movement (REM) sleep from 15-0% (1-1%) of total sleep time on control nights to 2-0% (1-5%) with negative pressure ventilation (p < 0.05), and an increase in stage 1 sleep (54-2% (3-9%) e 60-7% (6-0%); p < 0-05). Whereas all subjects reached all sleep stages on control nights, only three had REM sleep with assisted ventilation. Despite the polysomnographic findings, three of the five subjects claimed to benefit from more restful sleep with negative pressure ventilation, and none had subjective complaints of excessive daytime somnolence after three months of nightly use.

Hypopnoea associated with a less than 3% decrement in oxygen saturation was substantially more frequent on negative pressure ventilation nights than on control nights (table 2). Furthermore, although the overall mean (SE) frequency of apnoea and hypopnoea increased from 3-2 (1-3) to 59-3 (19-8) events/hour (p < 0-05) on negative pressure ventilation nights, only a minority (10-4 (7-1)) were associated with oxygen desaturation of 3% or more (table 2). The onset of obstructive apnoea and hypopnoea was associated with levels of inspiratory diaphragm EMG activity similar to those of the baseline before the onset of obstruction. The phasic activation increased with the progression of the apnoea or hypopnoea, reaching a maximum with the last obstructed breath.

On the control nights all episodes of obstructive apnoea and hypopnoea occurred during stages 1 and 2 and REM sleep. With negative pressure ventilation 6% of the events occurred during stage 3 or 4. No relationship was evident for the group between the peak negative pressure applied and the apnoea-hypopnoea index. Furthermore, there was no association between the degree of suppression of the phasic diaphragm EMG and the frequency of obstructive events.

The baseline oxygen saturation, recorded when subjects were awake and supine in the negative pressure ventilation set up, was similar on the two study nights (table 2). Despite the increased frequency of disordered breathing events on assisted ventilation nights, the lowest recorded overnight oxygen saturation was similar on control (81-4% (6-0%)) and negative pressure ventilation (85-6% (2-3%)) nights (table 2). The mean duration of episodes of apnoea and hypopnoea was also similar on control and negative pressure ventilation nights.

**Discussion**

Although the therapeutic role of ventilatory muscle rest in patients with chronic obstructive lung disease remains uncertain, preliminary data suggest that intermittent negative pressure ventilation may offer substantial benefit in such patients.5-7 Upper airway obstruction has long been recognised to be a complication of negative pressure ventilation, and has been observed during sleep in normal subjects8 as well as in patients.9-13 The purpose of the present study was to define the extent of the problem in a group of patients for whom negative pressure ventilation was being evaluated as a means of providing intermittent ventilatory muscle assistance.

Our findings show that negative pressure ventilation during sleep results in the induction of intermittent upper airway obstruction as well as altered sleep characteristics in patients with severe chronic obstructive lung disease. In so far as very few patients in the larger trial of ventilatory muscle rest chose to sleep with negative pressure ventilation, the results for the sample studied with the current protocol are not likely to be representative of the findings in a randomly selected population. As our subjects decided to apply the apparatus during sleep, they may well have been those who were least affected in terms of upper airway obstruction.

The mechanism of the upper airway obstruc-

<table>
<thead>
<tr>
<th>No of episodes of:</th>
<th>Control</th>
<th>Negative pressure ventilation</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Apnoea/h sleep</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Central</td>
<td>1-2 (0-6)</td>
<td>1-0 (NS)</td>
<td>&lt;0-05</td>
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<tr>
<td>Obstructive</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;3% desaturation</td>
<td>0-2 (1-0)</td>
<td>4-3 (1-9)</td>
<td>NS</td>
</tr>
<tr>
<td>≥3% desaturation</td>
<td>0-0 (0-0)</td>
<td>0-7 (0-3)</td>
<td>NS</td>
</tr>
<tr>
<td>Hypopnoea/h sleep</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;3% desaturation</td>
<td>1-4 (0-9)</td>
<td>44-6 (21-1)</td>
<td>&lt;0-05</td>
</tr>
<tr>
<td>≥3% desaturation</td>
<td>0-6 (1-3)</td>
<td>9-7 (7-3)</td>
<td>NS</td>
</tr>
<tr>
<td>Obstructive apnoea and hypopnoea/h</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>&lt;3% desaturation</td>
<td>1-6 (1-0)</td>
<td>49-0 (20-4)</td>
<td>&lt;0-05</td>
</tr>
<tr>
<td>≥3% desaturation</td>
<td>1-6 (1-3)</td>
<td>10-4 (7-0)</td>
<td>&lt;0-05</td>
</tr>
<tr>
<td>Total</td>
<td>3-2 (1-3)</td>
<td>59-3 (19-8)</td>
<td>NS</td>
</tr>
<tr>
<td>Apnoea or hypopnoea duration (%)</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Baseline SaO₂ (%) (awake, supine)</td>
<td>90-1 (2-1)</td>
<td>91-2 (1-0)</td>
<td>NS</td>
</tr>
<tr>
<td>Lowest overnight SaO₂ (%)</td>
<td>81-4 (6-0)</td>
<td>85-6 (2-3)</td>
<td>NS</td>
</tr>
</tbody>
</table>

*All episodes of apnoea with negative pressure ventilation were by definition obstructive (see text for details).
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