Respiratory muscle activity and oxygenation during sleep in patients with muscle weakness

J.E.S. White*, M.J. Drinnan**, A.J. Smithson**, C.J. Griffiths**, G.J. Gibson*


ABSTRACT: Patients with respiratory muscle weakness show nocturnal hypoventilation, with oxygen desaturation particularly during rapid eye movement (REM) sleep, but evidence in individuals with isolated bilateral diaphragmatic paresis (BDP) is conflicting. The effect of sleep on relative activity of the different respiratory muscles of such patients and, consequently, the precise mechanisms causing desaturation have not been clarified.

We have studied eight patients, four with generalized muscle weakness and four with isolated BDP during nocturnal sleep with measurements including oxygen saturation and surface electromyographic (EMG) activity of various respiratory muscle groups.

Nocturnal oxygenation correlated inversely with postural fall in vital capacity, an index of diaphragmatic strength. During REM sleep, hypopnoea and desaturation occurred particularly during periods of rapid eye movements (phasic REM sleep). In most subjects, such events were "central" in type and associated with marked suppression of intercostal muscle activity, but two subjects had recurrent desaturation due to "obstructive" hypopnoea and/or apnoea. Expiratory activity of the external oblique muscle was present whilst awake and during non-rapid eye movement (NREM) sleep in seven of the eight subjects in the semirecumbent posture. This probably represents an "accessory inspiratory" effect, which aids passive caudal diaphragmatic motion as the abdominal muscles relax at the onset of inspiration. Expiratory abdominal muscle activity was suppressed in phasic REM sleep, suggesting that loss of this "accessory inspiratory" effect may contribute to "central" hypopnoea.

We conclude that, in patients with muscle weakness, nocturnal oxygenation correlates with diaphragmatic strength. During sleep, episodes of oxygen desaturation are predominantly due to reduced muscle activity in association with eye movements during REM sleep resulting in central hypopnoea, though obstructive events may also occur.


It is well recognized that patients with respiratory muscle weakness are at risk of hypoxaemia and hypercapnia during sleep, especially rapid eye movement (REM) sleep [1, 2]. The likely mechanism is the normal reduction in respiratory drive associated with sleep acting on muscles with reduced capacity due to disease. Data in normal subjects during REM sleep suggest differential suppression of various muscle groups, with relative preservation of activity of the diaphragm [3–5]. It might, therefore, be expected that patients with the most severely impaired diaphragmatic function would be the most vulnerable during REM sleep. LAROCHE et al. [6] studied patients with isolated bilateral diaphragmatic paresis (BDP) and suggested that this was not the case, and that significant nocturnal hypoxaemia was seen only in the presence of generalized respiratory muscle weakness.

Hypopnoea and desaturation might also result if the activity of the upper airway muscles were disproportionately suppressed, leading to narrowing or occlusion of the upper airway. In a recent study of patients with Duchenne's muscular dystrophy, it was inferred that apnoeas were predominantly obstructive [7]. Furthermore, the frequency of obstructive apnoea or hypopnoea in patients with muscle weakness may be underestimated if reliance is placed on conventional sleep measurements, such as chest wall motion [8].

In awake subjects with severe weakness or paralysis of the diaphragm, the abdominal muscles may have an important role as accessory inspiratory muscles [9], but such an effect is dependent on gravity and it is not known whether it is relevant during sleep.

We have, therefore, performed detailed investigation of respiratory muscle activity during REM and non-rapid eye movement (NREM) sleep in patients with either...
generalized muscle weakness or isolated BDP, in order to assess the contribution of the different muscle groups to hypopnoea (hypventilation) and consequent hypoxaemia. In particular, we have concentrated on: 1) the comparative severity of hypoxaemia associated with generalized weakness and BDP alone; 2) the respective contributions of "obstructive" and "central" hypopnoea and their mechanisms; 3) the activity of the abdominal muscles as possible accessory inspiratory muscles during sleep.

**Subjects and methods**

Eight subjects were studied, four (Nos. 1–4) had generalized muscle disease and four (Nos. 5–8) had isolated BDP with normal maximal expiratory pressures measured at the mouth and no evidence of weakness of other skeletal muscles. Their diagnoses, characteristics and pulmonary function are shown in table 1. All except one (No. 1) showed paradoxical abdominal motion in the posteroanterior dimension both awake and asleep when semi-recumbent. No subject was taking any drugs which might have affected respiration. Subjects were recruited to the study on the basis of willingness to participate and were referred to the study on the basis of willingness to participate and might have affected respiration. Subjects were recruited to the study on the basis of willingness to participate and were referred to the study on the basis of willingness to participate and might have affected respiration. 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### Table 1. Subject characteristics

<table>
<thead>
<tr>
<th>Sub. No.</th>
<th>Sex</th>
<th>Age (yrs)</th>
<th>Diagnosis</th>
<th>BMI (kg/m²)</th>
<th>FEV1 (%)</th>
<th>VC (%)</th>
<th>ΔVC (%)</th>
<th>FRC (%)</th>
<th>Pdi,sniff (cm H₂O)</th>
<th>Pmax (cm H₂O)</th>
<th>Pmax (cm H₂O)</th>
<th>Awake* (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>F</td>
<td>36</td>
<td>Mitochondrial myopathy</td>
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<td>62</td>
<td>61</td>
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<td>Unclassified myopathy</td>
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<td>74</td>
<td>94</td>
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<tr>
<td>3</td>
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<td>49</td>
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<td>94</td>
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<td>-30</td>
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<td>91</td>
</tr>
<tr>
<td>4</td>
<td>M</td>
<td>68</td>
<td>Unclassified myopathy</td>
<td>26.6</td>
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<td>61</td>
<td>39</td>
<td>108</td>
<td>7</td>
<td>-18</td>
<td>75</td>
<td>97</td>
</tr>
<tr>
<td>5</td>
<td>M</td>
<td>71</td>
<td>Idiopathic BDP</td>
<td>26.2</td>
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<td>63</td>
<td>47</td>
<td>139</td>
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<td>80</td>
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<tr>
<td>6</td>
<td>M</td>
<td>57</td>
<td>Idiopathic BDP</td>
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<td>63</td>
<td>61</td>
<td>83</td>
<td>55</td>
<td>5</td>
<td>-78</td>
<td>124</td>
<td>94</td>
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<tr>
<td>7</td>
<td>M</td>
<td>63</td>
<td>BDP in association with diabetes mellitus surgery</td>
<td>36.6</td>
<td>41</td>
<td>51</td>
<td>81</td>
<td>149</td>
<td>8</td>
<td>-16</td>
<td>132</td>
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<tr>
<td>Mean</td>
<td></td>
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<td></td>
<td>31.7</td>
<td>58</td>
<td>57</td>
<td>53</td>
<td>107</td>
<td>15</td>
<td>-39</td>
<td>82</td>
<td>94</td>
</tr>
</tbody>
</table>

* Sub: subject; M: male; F: female; BDP: bilateral diaphragmatic paresis. BMI: body mass index (weight/height²); FEV1: forced expiratory volume in one second; VC: vital capacity; ΔVC: percentage fall in VC from erect to supine; FRC: functional residual capacity; Pmax: maximal inspiratory pressure measured at the mouth at TLC; Pdi,sniff: maximal transdiaphragmatic pressure during a maximal sniff from FRC. *: in sleeping (semi-recumbent) posture.
in the constant preferred posture were transferred to a personal computer using an analogue to digital converter (Dash-16, Metrabyte, Taunton, MA, USA). A total of 28 periods of NREM and 16 periods of REM sleep (ranges 3–6 and 1–5 per subject, respectively) were captured from the six subjects who had at least one epoch of REM sleep. These periods included mean (sd) numbers of 65 (15) and 73 (20) breaths, respectively. Analysis of EMG activity was performed using a specifically developed software package which calculated the peak level of EMG activity for each muscle for each breath, and for the captured section as a whole. Results were expressed as ratios between sleep stages.

Apnoea was defined as cessation of airflow and hypopnoea as a reduction in ribcage excursion of at least 50%, both for at least 10 s. Events were further classified as "central" if there was either an absence of, or an overall reduction in, inspiratory muscle activity and "obstructive" if the event was accompanied by persisting or increasing muscle activity with, at termination, a large burst of activity in EMGge, all in the presence of snoring. Since in most cases posteroanterior ribcage and abdominal motion was continually out of phase due to diaphragmatic weakness, this conventional criterion for the development of upper airway obstruction could not be used. An episode of desaturation was defined as a fall in SaO2 of at least 4%.

Oxygenation

Table 2 shows oxygenation data for all the subjects for the whole study and by sleep stage. Only one subject (No. 1) had a relatively normal overnight saturation profile; it was noteworthy that she had no REM sleep. Otherwise, the nadir SaO2 of the subjects was markedly reduced below normal. Recurrent desaturations were recorded in all except subject No. 1. They occurred most frequently, and generally were most severe, during REM sleep. However desaturations were also seen in NREM sleep in five subjects (Nos. 2, 4–6 and 8). In the six subjects who had at least one epoch of REM sleep, mean and nadir SaO2 during REM were lower than during NREM sleep (p<0.02) (table 2).

Daytime pulmonary function and nocturnal oxygenation

There was a significant negative correlation between overnight oxygenation (whether expressed as mean SaO2 or nadir SaO2) and the fall in vital capacity when changing from erect to supine posture (ΔVC) (fig. 1a and b) (r=-0.73 and -0.78, respectively; both p<0.05). Other measures of lung volume, e.g. functional residual capacity (FRC), measurements of maximum static respiratory pressures and transdiaphragmatic pressure during a maximal sniff (Pdi,sniff) showed no significant correlations with nocturnal oxygenation.

Muscle activity - wakefulness and NREM sleep

Phasic inspiratory activity in EMGge was seen in all subjects during sleep. During wakefulness, this was obscured in most cases by high levels of tonic activity. All subjects showed inspiratory activity in EMGint during wakefulness and NREM sleep. Inspiratory activity in EMGdi during wakefulness and NREM sleep was recorded in four subjects (Nos. 1, 2, 5 and 6). In the others (Nos. 3, 4, 7 and 8), apparent expiratory activity was present, but since this corresponded completely in timing and pattern with activity in EMGexob it was considered to be contaminant.

Results

Sleep

The duration and quality of sleep varied considerably between individuals. Subjects Nos 1 and 4 had no REM sleep. Three subjects (Nos. 4, 5 and 8) had no slow wave (stage 3–4) sleep. Subject No. 4 slept particularly poorly, spending more than 60% of the study time awake. He had recurrent arousals related to hypopnoea, resulting in gross sleep fragmentation.

Table 2. – Oxygenation data

<table>
<thead>
<tr>
<th>Subject</th>
<th>Total sleep time</th>
<th>NREM sleep</th>
<th>REM sleep</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean SaO2 %</td>
<td>Nadir SaO2 %</td>
<td>Desat. rate n·h⁻¹</td>
</tr>
<tr>
<td>1</td>
<td>97</td>
<td>94</td>
<td>0</td>
</tr>
<tr>
<td>2</td>
<td>93</td>
<td>68</td>
<td>13</td>
</tr>
<tr>
<td>3</td>
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<td>62</td>
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</tr>
<tr>
<td>8</td>
<td>72</td>
<td>50</td>
<td>21</td>
</tr>
<tr>
<td>Mean</td>
<td>90</td>
<td>69</td>
<td>14</td>
</tr>
</tbody>
</table>

Desat rate: desaturation rate; SaO2: arterial oxygen saturation; REM: rapid eye movement; NREM: non-REM.
Activity of EMGexob which was always expiratory in timing, was detected in seven of the eight subjects during most of the study period whilst awake and in NREM sleep. In two of these subjects (Nos 6 and 8) expiratory activity in EMGexob during NREM sleep was at times of considerable magnitude; such bursts of activity were accompanied by reduced abdominal paradox (fig. 2). No factors could be identified which were associated with these episodes of increased EMGexob activity: in particular, they were not related to changes in posture. Little activity was seen in EMGra in any of the subjects; where present it was expiratory in timing and mirrored the pattern of EMGexob.

Muscle activity - REM sleep

Six of the eight subjects (Nos. 2, 3 and 5–8) had at least one epoch of REM sleep. During pREM, intercostal muscle activity was significantly less than during NREM sleep (p<0.001) (fig. 3). Differences in genioglossus activity were more variable, with two subjects (Nos. 2 and 3) showing a small increase during pREM.

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**Fig. 1.** Relationship between nocturnal: a) mean oxygen saturation (mean $\text{Sa}_2\text{O}_2$); and b) nadir oxygen saturation (nadir $\text{Sa}_2\text{O}_2$) and fall in vital capacity on lying supine ($\Delta VC$) for patients with isolated bilateral diaphragmatic weakness (●) and generalized respiratory muscle weakness (○). Both show a significant negative correlation ($r$=0.73 and $r$=0.78, respectively; both $p<0.05$).

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**Fig. 2.** Recordings from subject No. 6 during wakefulness (left hand panel) and non-rapid eye movement (NREM) sleep (right hand panel). The recordings are in order, from the top: oxygen saturation ($\text{Sa}_2\text{O}_2$); airflow ($V'$); posteroanterior ribcage motion (RCPA); postero-anterior abdominal motion (ABPA); electro-oculogram (EOG); genioglossus group electromyographic (EMG) activity (EMGge); intercostal EMG (EMGint); diaphragmatic EMG (EMGdi); external oblique EMG (EMGexob). Note inspiratory activity in all three muscle groups and marked expiratory activity of EMGexob in NREM sleep. During wakefulness there is paradoxical abdominal motion throughout inspiration, but paradox only in late inspiration during NREM sleep.

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**Fig. 3.** Ratios of average peak EMG activity (error bar=1 SD) during phasic REM and NREM sleep recorded from genioglossus, intercostal and external oblique muscle groups in six subjects. (Nos. 1 and 4 had no pREM sleep). Intercostal activity was significantly less in pREM than in NREM sleep ($p<0.02$). Subject No. 2 showed no activity in EMGexob. REM: rapid eye movement; pREM: phasic REM; NREM: non-REM. For further abbreviations see legend to figure 2. : genioglossus; : intercostal; : external oblique.
sleep and the other subjects showing a reduction. Expiratory activity of EMGexob was present both during NREM and pREM in five subjects: its magnitude was significantly less in pREM sleep (p<0.05) (fig. 3). Of the three subjects (Nos. 2, 5 and 6) who showed inspiratory activity of EMGdi during pREM sleep, this was similar to the activity in NREM sleep in two subjects (Nos. 2 and 6) and less in the other.

Mechanisms of hypoxaemia and hypopnoea

No episodes of desaturation or periods of hypopnoea were recorded in subject No. 1. The numbers of hypopnoeas and apnoeas of different types for the other subjects are shown in table 3. Events during REM sleep were generally more prolonged and accompanied by more severe desaturation than during NREM sleep. Subject No. 4 (who had no REM sleep) showed a fall in SaO₂ every time he entered light sleep, due to "central" hypopnoea associated with a reduction in EMGint activity, followed at termination of each event by an arousal. Two subjects (Nos. 5 and 6) had periodic desaturation during both NREM and REM sleep. During NREM sleep, the desaturations were associated with "obstructive" events, with approximately equal numbers of apnoeas and hypopnoeas in one subject (No. 5), and predominantly hypopnoeas in the other (No. 6). During REM sleep, subject No. 5 again had predominantly "obstructive" apnoeas, although the mechanism was sometimes more complex, as illustrated in figure 4 where a long obstructive apnoea encompasses a brief "central" component due to reduced intercostal muscle activity associated with phasic eye movements. During such "obstructive" events, there was an increase in abdominal motion, which remained paradoxical throughout the study. Desaturation during REM sleep in all subjects other than subject No. 5 was due to "central" hypopnoea. These episodes during REM sleep were closely associated with phasic eye movements (fig. 5).

Discussion

Background. Few studies of the activity of the inspiratory muscles during sleep have been reported even in normal subjects. In general, they show a reduction in
intercostal muscle activity during REM compared with NREM sleep, usually, but not exclusively in association with eye movements. In healthy individuals, however, activity of the diaphragm is generally preserved or even increased in REM compared with NREM sleep [3–5]. If a similar pattern were to occur in REM sleep in patients with weakness of the respiratory muscles, particularly the diaphragm, profound consequences might be expected.

A number of studies of nocturnal ventilation in subjects with muscle weakness have been reported. Newsom Davis et al. [14] studied eight patients with severe BDP, seven of whom had generalized neuromuscular disorders. Blood gas values during sleep showed a marked deterioration. Skatrud et al. [1] reported the case of a man with BDP due to limb girdle dystrophy, who had alveolar hypoventilation when awake which worsened during sleep. They inferred from reduced chest wall movement that the numerous repetitive desaturations seen during REM sleep resulted from a deterioration of intercostal muscle activity, but no direct measurements were made. Bye et al. [2] showed markedly reduced accessory muscle activity in REM sleep in one patient with BDP secondary to polio. Nocturnal studies of patients with myotonic dystrophy have shown that periodic desaturation can result from either ”central” or ”obstructive” events [15], and a recent study in patients with Duchenne’s muscular dystrophy found predominantly obstructive events during sleep [7]. A further study of patients with Duchenne’s muscular dystrophy [8] noted that ”obstructive” events may be misinterpreted as ”central” if conventional sleep recordings are made in patients with muscle weakness unable to generate large inspiratory pressures.

The above studies investigated patients with BDP in the context of generalized muscle disease and there have been fewer studies on patients with isolated BDP. Stradling and Warley [16] reported one such patient who had normal daytime blood gases but recurrent episodes of oxygen desaturation during sleep, which they deduced were due to ”central apnoeas”. Kreitzer et al. [17] described a non-obese patient with apparently isolated BDP, who presented in respiratory failure. They noted that respiratory failure was worse in the supine than the erect posture and that there was further deterioration during sleep, but the mechanism was not studied. By contrast, La roche et al. [6] studying six patients thought to have isolated BDP concluded that they did not have marked nocturnal hypoventilation. Three of their patients were said to have brief episodes of mild oxygen desaturation, but detailed information was not given.

We report the detailed recordings of respiratory muscle activity during REM and NREM sleep in patients with generalized muscle weakness and isolated BDP in order to assess the contributions of the different muscle groups to hypopnoea or apnoea and consequent hypoxaemia. In particular, we have concentrated on: 1) the severity of hypoxaemia associated with BDP alone; 2) the respective contributions of ”obstructive” and ”central” events and their mechanisms; and 3) the role of abdominal muscles as potential accessory inspiratory muscles during sleep.

**Technical considerations.** Despite an acclimatization night to minimize the ”first night effect” our subjects generally slept poorly. We did, however, obtain good quality recordings during REM sleep in six of the eight subjects. The data available for analysis were also limited by restricting measurements to those obtained in a constant posture.

Surface EMG recordings have the disadvantage of reduced specificity compared with those obtained using needle electrodes. The electrode placements were, however, chosen to reflect EMG activity of the relevant muscles with least contamination from other muscles. Moreover, in the case of genioglossus, previous work has shown EMG activity from transmandibular electrodes to be qualitatively similar to signals from intramuscular electrodes [18]. We found that contamination of EMGs by abdominal muscle activity was easily differentiated from that of the diaphragm itself by its expiratory timing. Since the abdominal muscle most commonly active in healthy awake subjects during stimulated breathing is transversus abdominis [19], this muscle may also have been active in our patients. It is, however, unlikely that the expiratory EMG activity that we observed originated from transversus abdominis, since this is the abdominal muscle least accessible to surface recordings.

**Oxygenation.** Nocturnal oxygenation was reduced compared with normal subjects, with the most severe desaturation occurring during REM sleep (table 2). Desaturation was seen in both groups of subjects and was not confined to those with generalized respiratory muscle weakness. On the contrary, figure 1 suggests that the degree of diaphragmatic weakness is an important determinant of desaturation both in generalized muscle weakness and isolated BDP. This conclusion is not incompatible with the results of La roche et al. [6]; although their subjects showed only mild nocturnal desaturation they had less awake hypoxaemia than those in the present study. The postural change in VC, however, is dependent not only on diaphragmatic strength but also on other factors, such as obesity, which is common in patients with muscle weakness [20] and was present in some of the subjects in this study. A large ∆VC is likely to be associated with a large reduction in FRC which might contribute to more severe desaturation; no direct measurements of absolute lung volume during sleep are, however, available in patients with respiratory muscle weakness. Bye et al. [2] reported a similar relationship between ∆VC and nocturnal oxygenation in neuromuscular disease. The importance of lung volume for the severity of oxygen desaturation during sleep was demonstrated by the correlation between expiratory reserve volume (ERV) and severity of desaturation in patients with obstructive apnoea [21]. It is likely that lung volume and, thus, ”oxygen reserve” is an important determinant of the severity of desaturation, and that the more severe the diaphragmatic weakness the smaller that reserve will be. Hence, the degree of functional diaphragmatic impairment is an important determinant of the severity of overnight hypoxaemia.
Mechanisms of desaturation. In all but two subjects (Nos. 5 and 6) events were characterized by hypopnoea rather than complete apnoea. Oxygen desaturation during these events was sometimes less than 4%; equally, some episodes of hypoventilation lasted less than 10 s and, therefore, were not scored as hypopnoea and yet resulted in desaturation of ≥4%. Consequently, small discrepancies were seen between the numbers of respiratory events and desaturations as defined for the purposes of this study. We classified as “central” those events where an overall reduction in inspiratory EMG activity was observed; “obstructive” events were those where muscle activity was maintained or increased during hypopnoea/apnoea with evidence of arousal and a large increase in EMG at event termination in the presence of snoring. The predominant type of event (i.e. whether “central” or “obstructive”) was generally consistent within an individual, but varied between subjects. Occasional events were more complex, as shown by the example in figure 4. The commonest mechanism of desaturation during pREM sleep was “central” hypopnoea associated with suppression of activity of the intercostal muscles, supporting the conclusions of other studies in patients with muscle weakness where “central” hypopnoea due to reduced intercostal muscle activity during REM sleep has been inferred as the mechanism of hypoventilation and desaturation [1, 16]. Obstructive hypopneas and/or apnoeas were seen in two subjects, but in one these were confined to NREM sleep.

The degree to which the activity of any particular muscle group was altered during pREM sleep varied between individuals (fig. 3). Genioglossus activity was more variably affected than that of the intercostal muscles. Clearly, differential suppression of inspiratory muscle groups has implications for the mechanism of the event. In general, an important determinant of whether an apnoea or hypopnoea is “central” or “obstructive” is the balance of activity between the upper airway dilator muscles and the inspiratory “pump” muscles. Clearly, other factors, such as upper airway anatomy and compliance, also play an important part in determining the extent to which the upper airway narrows or occludes (and thus whether an event appears to be “obstructive”).

REM sleep is not an homogeneous state but consists of phasic REM (pREM) sleep, characterized by rapid eye movements and myoclonic twitches, and tonic REM (tREM) sleep, in which there is muscle atonia and desynchronized EEG [22]. In healthy subjects, reductions in ventilation during REM sleep are of variable extent and duration and occur particularly in relation to periods of rapid eye movement. i.e. pREM [23–25]. Figures 1–4 illustrate the important associations between phasic eye movements during REM sleep and reduced activity of the inspiratory muscles in these patients, as has also been shown in normal subjects [5] and patients with COPD [26, 27].

Abdominal muscles. We observed phasic expiratory activity of the external oblique muscle in seven of the eight subjects when semirecumbent during both wakefulness and NREM sleep, even though this was not continuous in all subjects. These results provide direct support for the conclusions of Newsom Davis et al. [14], who inferred such activity from measurements of abdominal EMG. Expiratory activity of the abdominal muscles produces diaphragmatic motion in the cranial direction; their relaxation at the start of inspiration together with the effect of gravity produces passive descent of the diaphragm, which facilitates inspiration. This “inspiratory accessory” activity of the abdominal muscles will only occur if the subjects are at least partially upright, as was the case in this study. Two subjects (Nos. 6 (fig. 2) and 8), intermittently showed pronounced EMG exob activity accompanied by a change in abdominal motion. In the presence of such activity, the PA abdominal dimension, which previously declined during inspiration (abdominal paradox), showed an increase in early inspiration with a reduction only towards the end of inspiration. The initial increase is compatible with passive diaphragmatic descent at the start of inspiration, consequent on relaxation of the abdominal wall muscles. It was also noteworthy that expiratory activity of EMG exob was reduced or absent during “central” hypopnoeas (fig. 5), suggesting that suppression of this activity and the consequent loss of passive caudal motion of the diaphragm may also contribute significantly to the hypopnoea and consequent hypoxaemia.

In conclusion, our results show that in patients with respiratory muscle weakness the severity of nocturnal hypoxaemia is related to the postural fall in vital capacity, suggesting that the degree of impairment of diaphragmatic function is an important determinant of nocturnal oxygenation, irrespective of involvement of other respiratory muscles. Secondly, hypopnoea or apnoea during sleep in weak patients may be either “obstructive” or “central” depending on the pattern of suppression of the different groups of respiratory muscles. In REM sleep, events occur particularly during periods of phasic eye movements. Finally, expiratory activity of the external oblique muscles persists during NREM sleep in semirecumbent subjects; this probably aids early inspiratory descent of the weak or paralysed diaphragm and suppression of this activity during phasic REM sleep may contribute to “central” hypopnoea in some individuals.

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References


