

Domiciliary nasal respiratory support - first experiences in Malta

S. Montefort*, R. Camilleri*, A. Galea Debono*

ABSTRACT: Nasal respiratory support is a non-invasive alternative to conventional assisted ventilation with endotracheal intubation, or the more cumbersome negative pressure ventilators. The two main types of this relatively new therapy are nasal intermittent positive pressure ventilation [NIPPV] and nasal continuous positive airway pressure [NCPAP] respiratory support, which are mostly used in chronic hypoventilatory states and obstructive sleep apnoea [OSA] respectively. We have introduced these two types of respiratory support to five patients suffering from neuromuscular disorders and twenty-four patients with OSA with marked improvement in the quality of life of all patients concerned. Our experiences with these patients should hopefully lead to further development in the diagnostic and therapeutic facilities in this field in Malta.

*Department of Medicine, St. Luke's Hospital, Guardamangia

Correspondence: S. Montefort, Department of Medicine, St. Luke's Hospital, Guardamangia

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Introduction

Until recently, patients with ventilatory problems needing long-term artificial assisted ventilation, initially required endotracheal intubation and later a permanent tracheostomy or the cumbersome 'cuirass'. Since the advent of nasal respiratory support, physicians can treat these patients without having to resort to such invasive measures, thereby improving the patient's quality of life markedly.

Nasal intermittent positive pressure ventilation [NIPPV] and nasal continuous positive airway pressure [NCPAP] are the two most widely used modes of this type of non-invasive respiratory support, and when used in the appropriate cases, can transform the patient's lifestyle in the short term^{1,2} and improve the prognosis of his ventilatory condition together with its secondary effects, in the long term^{3,4}.

Nasal respiratory support has been used in Malta in the last two years and we have succeeded in starting a good number of patients on this type of treatment from which they are still benefiting today.

The start of nasal respiratory support in Malta

Up to two years ago the only Maltese patient utilising domiciliary NIPPV was a young girl suffering from Ondine's curse secondary to damage of her respiratory centre by severe pertussis infection. She had initially been treated in an 'iron lung' in the intensive care unit at St. Luke's Hospital, after unsuccessful attempts at weaning her off invasive ventilation. Following transfer to the UK, she was started on an NIPPV ventilator which she subsequently used every night because of nocturnal hypoventilation. She forms part of our series of patients described below. In November 1994 a fourteen year old male suffering from Duchenne muscular dystrophy, who

was admitted to hospital 'in extremis', was the first patient to be started on NIPPV at St. Luke's Hospital. He had been complaining of morning headaches for the weeks preceding admission and his parents had noted that he was becoming more drowsy and cyanosed as time went by. Ultimately he lapsed into unconsciousness on the day of admission when his arterial blood gases revealed a P_aCO_2 of 94.2mm Hg and a P_aO_2 of 65mm Hg. Within two hours of commencing NIPPV he had regained consciousness and his arterial blood gases had improved markedly [P_aCO_2 62mm Hg, P_aO_2 82mm Hg]. This young man is still alive today and is using NIPPV nightly to compensate for nocturnal hypoventilation and daytime hypoxia and hypercapnia. The success of these two cases prompted the application of nasal respiratory support for other such patients and also for other indications.

Nasal intermittent positive pressure ventilation [NIPPV]

Nocturnal hypoventilation is the major indication for this type of ventilation in which the airway pressure changes phasically throughout the cycle, delivering the entire tidal volume and then allowing passive exhalation. NIPPV is thought to help these patients by resting fatigued muscle⁵, increasing lung compliance⁶, and restoring some of the blunted CO_2 sensitivity of the respiratory centre⁷. Like other types of nasal respiratory support, it is administered through a small ventilator/blower which may either be volume or pressure-dependent. The positive pressure is instituted through a tight-fitting nasal mask which forms a seal by being held to the face by an appropriate type of head-gear⁸ (See Fig 1). This assembly transmits the pressure or volume generated by the ventilator to the airways. The development of silicone masks have made these much

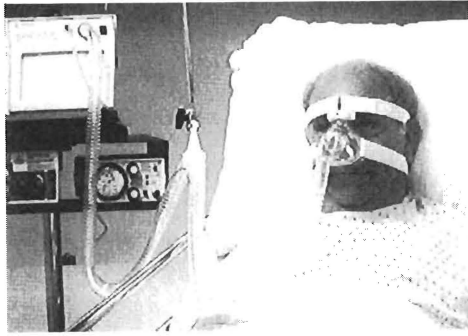


Fig. 1 - A patient with the nasal mask and head gear used in nasal support

more comfortable while creating better seals. Improvements are being made continuously such as full face masks for patients who have mouth-leaks and small nasal cushion masks for those who find the bigger masks claustrophobic⁹.

Nasal continuous positive airway pressure [NCPAP]

With NCPAP positive airway pressure is maintained, relative to atmospheric pressure, throughout the cycle¹⁰. NCPAP was introduced by Dr. Colin Sullivan in Australia as a means of splinting the upper airways during sleep in patients with obstructive sleep apnoea [OSA]¹¹. Nasal bi-level positive airway pressure [NBIPAP] is a modification of NCPAP in that the positive pressure is decreased slightly during the expiratory phase of the cycle so as to make the whole process more comfortable for the patient thus, presumably, increasing the compliance rate.

The commonest symptoms of obstructive sleep apnoea are excessive daytime somnolence and irregular pattern of snoring, which latter, in the majority of cases, is due to repeated collapse of the oropharyngeal musculature during sleep. This results in complete or partial obstruction of the subject's airway accompanied by a reduction in blood oxygen saturation and followed by waking to breathe. These repeated episodes lead to sleep fragmentation with the deeper stages of sleep [rapid eye movement stages] being the most affected. It should be distinguished from the much less common central sleep apnoea which is a neurological disturbance resulting in a cessation of respiratory effort in sleep. OSA has come to the medical forefront in the last few years as the commonest type of sleep-related breathing disorder. Many sufferers are unaware of the cause of their excessive daytime sleepiness and pathological tendency to fall asleep, sometimes even while driving their car, leading to fatal or near-fatal accidents¹². Their sleeping partner complains about their loud snoring interrupted by long breathing pauses and excessive body-movements. This often leads to marital disharmony, mental depression¹³, lack of libido and impotence¹⁴. OSA adversely affects the quality of life of the subject and can decrease the life expectancy of a 50 year old sufferer by 50%¹⁵. The effects of OSA on the cardiovascular system include life-threatening arrhythmias¹⁶, systemic hypertension¹⁷ and myocardial ischaemia¹⁸, while the recurrent hypoxic episodes can also lead to pulmonary vascular disorders¹⁹.

The diagnosis is made from a suggestive history and a sleep-study (polysomnogram)²⁰ in order to differentiate OSA from other sleep-related disorders and simple snoring^{21,22}. During polysomnography a number of the following parameters are monitored: electroencephalogram (EEG), electrocardiogram (ECG), ocular and genohyoid electromyogram (EMG), respiratory effort and air flow, oxygen saturation, apnoea/hypopnoea time and frequency, body position, limb activity and snoring²³. An important analysis is the apnoea index which is the number of breathing pauses lasting 10 seconds or longer per hour of sleep. An apnoea index of more than 5 is considered pathological by some investigators but others feel that 10 - 15/hr is a more specific measure for diagnosis²⁴. Oxygen desaturations of >4% of baseline during hypopnoeas (a reduction in airflow associated with a fall in oxygen saturation and an arousal from sleep)²⁵ also points towards a diagnosis of OSA. Difficulties in defining the diagnosis²⁶ may explain the differences in prevalence rates between studies but the range is between 0.5 - 5% of the population^{24,27}.

This paper describes how we screened, diagnosed and treated the first Maltese patients with OSA and nocturnal hypoventilation using nasal ventilation. The future of this modality of treatment in Malta is then discussed.

Method

1) NIPPV Subjects

Since 1994, four patients suffering from neuromuscular conditions with secondary nocturnal hypoventilation were started on NIPPV. Three presented in acute Type II respiratory failure. A young girl was started on NIPPV for Ondine's curse following severe pertussis affecting her respiratory centre, and continued to be followed up. (Table 1)

Ventilators

A bulky Bromptonpac ventilator (Pneumopac Ltd., Luton) was initially used on two of these patients and later changed to a pressure-dependent NIPPV ventilator (Thomas Respiratory Systems UK) weighing around 5 kgs. Two of the other patients are also on such a ventilator and all these four patients have had their inspiratory pressures set at around 10 cms H₂O.

Table 1 - Patients on NIPPV

Subject	Sex	Age (yrs)	Condition leading to res. failure
1	F	16	Central Res. Hypoventilation
2	M	14	Duchenne Muscular Dystrophy
3	F	42	Limb-Girdle Muscular Dystrophy
4	F	37	Limb-Girdle Muscular Dystrophy
5	M	17	Duchenne Muscular Dystrophy

2) NCPAP

Subjects

Twenty-four patients (21 male and 3 female) with a mean age \pm SEM 43.4 \pm 2.9 yrs (range 19 - 66 yrs) all suffering from obstructive sleep apnoea were treated on NCPAP. All had a neck size of more than 17 inches in circumference (range 17 - 19.5 inches). Most were above the ideal weight for their height except two, who had an element of micrognathia. They all came from different walks of life but the majority (14 out of 24) were of professional or executive status.

Diagnosis of obstructive sleep apnoea

These twenty-four patients were diagnosed using a combination of history, clinical examination, blood investigations and a partial sleep study.

History

The help of the patient's sleeping partner was sought in taking a history where we specifically enquired about cardinal features such as excessive snoring with apnoeic spells, nocturnal choking sensation, nocturia, reduced libido, morning headaches, increased limb movement in sleep, lack of daytime concentration and general well-being. A modification of the Epworth sleepiness scale²⁸ was used to try and grade the patients' daytime sleepiness in specific situations such as when driving, reading, watching TV or talking to someone. Seventeen of the patients were asked to put a score of 1 to 10 on a number of these symptoms before and after starting to use NCPAP so that we could grade any improvement the patient might experience. Enquiry was also made about alcohol or sedative ingestion, smoking, sleep hygiene, shift work, attempted weight reduction, nasal blockage and symptoms suggestive of hypothyroidism.

Clinical examination

During the clinical examination emphasis was put on nasal patency, the state of their soft palate and uvula, presence of micrognathia, neck size, blood pressure and any clinical signs suggestive of hypothyroidism and acromegaly.

Blood investigations

Apart from the routine blood tests the patients were screened for hypothyroidism and acromegaly in cases where these conditions were suspected.

Partial sleep studies

These studies were carried out at the coronary care unit at St. Luke's Hospital, when a monitor bed was available for the night in question. Using these monitors, continuous percutaneous oxygen saturation, ECG, blood pressure, respiratory rate and apnoea frequency were studied. The nurses on night duty and one of the investigators (SM) observed the patient from time to time during the night to try and observe the

snoring pattern, apnoeas, limb movements and body position during these events. In all we screened 38 patients, 24 of which met most of the criteria we could monitor for OSA. When possible or in cases where the initial study revealed heart rhythm problems, we carried out another study during the first night on the machine.

NCPAP machines

In order to try and maximise the compliance rate we opted to use NBIPAP on most of the patients so as to make this treatment as comfortable as possible. In fact 11 of our patients are on Respironics BIPAPS machines with the pressures used ranging between 10 - 18 cms H₂O for the inspiratory positive airway pressure and 5 - 14 cms H₂O for the expiratory positive airway pressure. The machine was set on spontaneous mode so that the patient would trigger off the machine and not encounter undue resistance on expiration. Once adequate experience had been gained, it was decided to start the next patients on NCPAP (a mix of Respironics Remstar, DeVilbiss, Healthdyne and ResMed machines), which proved to be sufficient for the rest of the patients and considerably less expensive than the NBIPAP machines.

Statistical analysis

The symptom scores before and after NCPAP were compared using the Wilcoxon paired rank test. A p value of <0.05 was taken to represent a significant association or difference.

Results

NIPPV

All five patients have noted a very marked subjective improvement in their quality of life since initiation of treatment and this was confirmed by their relatives. One of the young men is back attending school while another is now much more responsive and in good spirits than before treatment. The latter initially had some difficulty with increased oral secretions on using the ventilator but this has decreased since being started on inhaled Ipratropium Bromide² inhalations nocte through a spacer device. He is now also having some swallowing difficulties which are probably secondary to his muscular dystrophy. In the past, these patients would have succumbed to chest problems before encountering these difficulties. Similar improvement was noted in the other two female patients, both of which have returned to their jobs. All these patients are very compliant with treatment and are on the ventilator for at least six hours every night. Some even use it when experiencing problems in breathing during the day. At first some, especially the younger ones, felt claustrophobic with the nasal mask, but soon, all got used to it.

The girl with the Ondine's curse was recently sent to the UK for a full sleep study and told that she could now stop using the NIPPV on a regular basis.

NCPAP

Of the 38 patients screened, 24 had enough evidence for a diagnosis of OSA. None of the patients screened

had any endocrine or major nasal problem to explain their complaints, while those who imbibed alcohol with their evening meal (four) or used sleeping-pills (three) stopped doing so. These measures, however, did not improve their sleep-related breathing disorder. All the overweight subjects had tried to lose weight but only two managed to lose a substantial amount and even these did not improve enough to render treatment with nasal CPAP unnecessary. There was no great difference in the sleepiness scale between the patients who ended up on nasal respiratory support and those who did not. However there was a marked difference before and after starting nasal ventilation in how sleepy the patient felt during the day (mean score before vs mean score after) (7.8 vs 1.6) and during driving (6.9 vs 0) (Fig 2). This improvement was also noted for nocturnal choking sensation (4.4 vs 0), concentration (5.1 vs 8.2) and general well-being (4.2 vs 8.8). The snoring in the treated group was noted to be as loud as in the group who were subsequently labelled as simple snorers. However, the snoring was more irregular, reached a crescendo more often and was interspersed with apnoeic spells. Much to the relief and appreciation of the patients' sleeping partners the patients on nasal NCPAP stopped snoring almost completely (9.3 vs 0.5). All these changes reached statistical significance ($p < 0.05$). These symptom scores were not carried out by our seven most recent patients as it was too early to grade any improvement in concentration and well-being but they too improved in the rest of the symptoms.

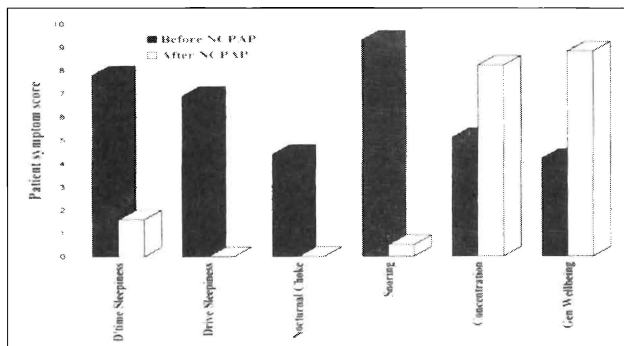


Fig. 2 - Histogram showing patient scores for various obstructive sleep apnoea symptoms before and after NCPAP

The partial sleep studies which were considered positive showed repeated oxygen desaturations which had to be at least 4% of baseline to be considered significant. In some cases these oxygen saturations dipped down even to the level of 55%. Four of our patients developed very worrying arrhythmias (one had episodes of complete heart block, another had pauses of severe bradycardia (Fig 3) and the other two had runs of fast atrial fibrillation together with multiple ventricular ectopics) during some of the oxygen desaturations. Before inserting permanent pacemakers, it was decided to start nasal respiratory support urgently and this corrected the arrhythmias together with the oxygen desaturations in the two patients with bradyarrhythmias. The other two patients, who happen to be two of our recently treated patients, had fewer runs of atrial fibrillation than before during their first night on NCPAP. It is planned to repeat their sleep study in a few

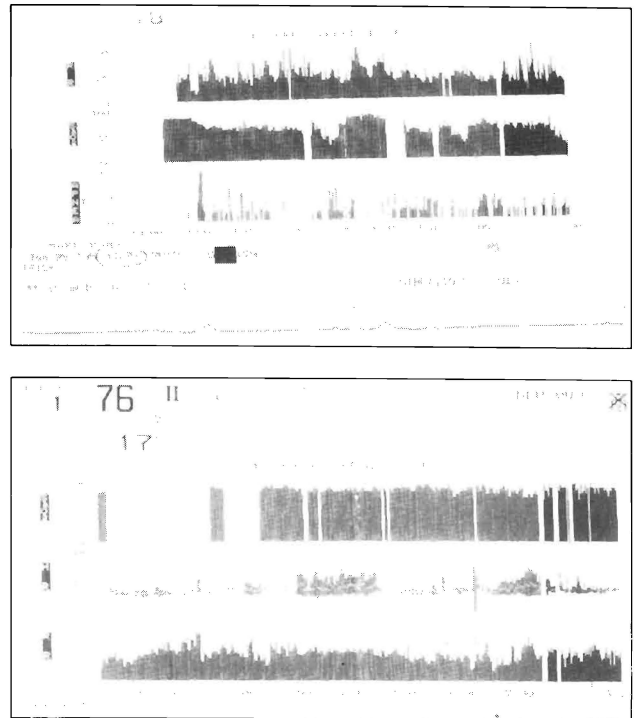


Fig. 3 - Nocturnal oximetry and ECG readings in a patient suffering from OSA
 a) before NCPAP showing desaturations and arrhythmia
 b) while on NCPAP showing correction of abnormal parameters

weeks time to see whether the arrhythmogenicity of their myocardium will have decreased following the use of NCPAP for this period of time.

The patients on whom NCPAP or NBIPAP was instituted, were all admitted to hospital for their first night on the ventilator. All but two accepted the nasal mask very well. One of these two patients who continued to find difficulties was given a nasal bubble-cushion mask (Monarch mask; Respironics, USA) and has adapted to this very well. The other patient has decided to postpone regular use of NCPAP for some time. Once he decides to initiate treatment we plan to use the Monarch mask again so as to increase the chances of acceptance. The pressure settings of the machine were adjusted according to the build of the patient²⁹ as we did not have the facility to titrate airway pressures in our partial sleep studies. During this first night we decreased the pressures gradually till the patient either started snoring or breathing irregularly and then increased again to just above the breakthrough pressure level. All the patients, without exception, commented favourably about how much better they felt on waking up after that first night and their subjective improvement continued to increase over the first few nights. Subsequent improvement decreased when patients defaulted treatment for as little as 2-3 nights. The most common complaints were airway dryness, nasal bridge soreness and facial marks from the mask, breathing warm air on hot nights and the aesthetics of the whole process.

Discussion

In this paper we have demonstrated the introduction of

a novel mode of treatment for patients with nocturnal hypoventilation secondary to neuromuscular conditions and for OSA sufferers in Malta. Nasal respiratory support has improved the quality of life in all these patients, indeed for three patients with neuromuscular disease, this proved to be life-saving. This improvement in quality of life is the short term result of the treatment^{2,30}. The long term benefits which have been shown to occur in major studies, will hopefully be experienced by our patients as well.

In the past, most patients suffering from muscular dystrophy, would die prematurely from respiratory problems - usually due to acute or acute on chronic respiratory failure precipitated by a chest infection. Preceding this, these patients usually develop cor pulmonale from hypoventilation due to their muscular weakness and thoracic skeletal malformations. Now, these acute and chronic chest problems can all be helped by NIPPV as demonstrated in this series of patients. Similar patients presenting in acute respiratory failure usually present a difficult dilemma to the admitting physician as whether to intubate and ventilate, knowing that their chances of being weaned off are slim³¹. Despite a poor prognosis, these patients would occupy a precious ITU bed for a long time. Nasal ventilation offers a useful alternative to full ventilatory support. Indeed, this was successful in the three acute situations encountered. In the chronic situation, all these patients felt better during their waking hours as their hypercapnic and hypoxic state was ameliorated.

NIPPV can also be used in other cases of chronic respiratory failure such as multiple sclerosis, central hypoventilation syndrome, motor neurone disease and chest wall deformities, but its use in chronic obstructive airway disease is still controversial³². The current position is that NIPPV seems to benefit only those cases of COAD who have high daytime P_aCO_2 ($>56\text{mmHg}$)³³ and may also be considered in acute exacerbations³⁴ in patients who either refuse intubation or who would not otherwise be considered ideal for endotracheal intubation and ventilation.

NCPAP and NBIPAP, the other mode of nasal respiratory support utilised in this series of patients, also proved to be of major benefit to the patients involved. OSA is a debilitating condition with implications and negative effects on the patient's personal, social, professional and physical lifestyle. Again the long-term effects of this condition can also lead to an untimely death due to cardiac arrhythmias and ischaemia together with the complications encountered in chronic respiratory failure and systemic hypertension^{16,18}. The real life caricature of an overweight man who dozes off easily and snores loudly presents a serious medical problem. OSA is little known among lay people and doctors alike, and must be 'publicised' so as to be recognised and treated more often. The number of patients presenting to us is now increasing partly as a result of the positive feedback from successfully treated patients. Our patients have all experienced a great improvement in their quality of life and this, in turn, is improving their compliance to treatment.

The fact that accidents such as motor-vehicle collisions can affect third parties, makes the situation even more worrisome. The condition is much more common than one expects with 2 - 4% of the American

population thought to suffer from various grades of OSA²⁴. The fact that the average Maltese person is rather overweight makes it likely that the prevalence of OSA in our country is at least just as high.

Great changes are being made in the field of development of NCPAP machines as these ventilators are now becoming more 'user-friendly', comfortable, lighter and thus more portable. Features such as ramping, which allows a slow build-up of pressure after sleep onset, demand CPAP and AutoCPAP in which the machine senses when the patient does not require CPAP and switches off, have become available in the last couple of years. All these are designed to increase patient compliance³⁵. Options such as drastic surgery³⁶ as in uvulopalatopharyngoplasty (UVPP)³⁷ have not been very successful, while newer laser surgery is still unproven. Other studies have looked at tracheostomies and oral appliances³⁸ but these are not very popular. In children, where sleep apnoea is not encountered, surgery in the form of adenoidotomy tends to correct the majority of OSA cases³⁹. NCPAP is only required in children with craniofacial anomalies, trisomy²¹ and skeletal dysplasia syndromes, who have not responded fully to surgery⁴⁰.

In our opinion, the setting up of a sleep laboratory is badly needed at our hospital as evidenced by the above results. The condition is common and treatable, but the current facilities are not sensitive or specific enough for us to be sure that we are detecting all cases of OSA amongst the patients we screen. At the present time we are only able to monitor few of the criteria required for a diagnosis and even these are not being sampled frequently enough⁴¹. We are also currently unable to titrate NCPAP pressures and have to use a rather crude method to decide pressure settings for different patients. The institution of a sleep laboratory does not entail alot of expense apart from the initial expenditure and the training of a sleep technician. If space is a problem there are new polysomnograms available on the market which the patient can take home with him for monitoring in his usual nocturnal surroundings⁴². Another facility that would obviously be required is a sleep clinic where these patients would be seen, screened and followed up. Such a clinic would also be able to carry out research and try these modes of therapy for new indications such as acute left ventricular failure⁴³.

In conclusion we have reviewed the first series of patients which has been initiated on respiratory support for different conditions in Malta. We have discussed the positive impact of this treatment on these patients and their condition and demonstrated that this is a field which is worth pursuing. The development of domiciliary respiratory support can offer significant benefits to patients with respiratory insufficiency and should be pursued by clinicians and the Health Division, alike.

References

1. Lamphere J, Roehrs T, Wittig R, Zorick F, Conway WA, Roth T. Recovery of alertness after CPAP in apnoea. *Chest* 189; 96:1364 - 7.
2. Dardarian SS, Bridenbaugh RH, Rajagopal KR. Neuropsychologic symptoms in obstructive sleep apnoea improve after treatment with nasal continuous positive airway pressure. *Chest* 1988; 94:1023-7.
3. Sawicka EH, Loih L, Branthwaite MA. Domiciliary

- ventilatory support: an analysis of outcome. *Thorax* 1983; 43:31-5.
4. Partinen M, Guilleminault C. Mortality of patients with obstructive sleep apnoea: a follow-up study. *Chest* 1988; 94:1200-4.
 5. Roussos C. Function and fatigue of respiratory muscles. *Chest* 1985; 88: 1245-1315.
 6. Hoepfner VH, Cockcroft DW, Dosman JA, Cotton DJ. Nighttime ventilation improves respiratory failure in secondary kyphoscoliosis. *Am Rev Respir Dis* 1984; 129:240-3.
 7. Goldstein RS, Molotiu N, Skrastins R, Long S, De Rosie J et al. Reversal of sleep-induced hypoventilation by nocturnal negative pressure ventilation in patients with restrictive ventilatory impairment. *Am Rev Respir Dis* 1987; 135:1049-55.
 8. Sanders MH, Moore SE, Eveslage J. CPAP via nasal mask: a treatment for occlusive sleep apnoea. *Chest* 1983; 83:144-5.
 9. Mayer LS, Kerby GR, Whitman RA. Evaluation of a new nasal device for administration of continuous positive airway pressure for obstructive sleep apnoea. *Am Rev Respir Dis* 1989; 139:A114.
 10. American Thoracic Society Statement: Indications and standards for use of nasal continuous positive airway pressure (CPAP) in sleep apnoea syndromes. *Am Rev Respir Dis* 1994; 150:1738-45.
 11. Sullivan CE, Issa FG, Berthoin-Jones, Eves L. Reversal of obstructive sleep apnoea by continuous positive airway pressure applied through the nares. *Lancet* 1981; 1:862-5.
 12. Findley LJ, Weiss JW, Jabour ER. Drivers with untreated sleep apnoea. A cause of death and serious injury. *Arch Intern Med* 1991; 151:1451-2.
 13. Berrettini WH. Paranoid psychosis and sleep apnoea syndrome. *Am J Psych* 1980; 137:493-4.
 14. Santamaria JD, Prior JC, Fleetham JA. Reversible reproductive dysfunction in men with obstructive sleep apnoea. *Clin Endocrinol Oxf* 1988; 28:461-70.
 15. He J, Kryger MH, Zorick FJ, Conway W, Roth T. Mortality and apnoea index in obstructive sleep apnoea. Experience in 385 male patients. *Chest* 1988; 94:9-14.
 16. Guilleminault C, Connolly SJ, Winkle RA. Cardiac arrhythmias and conduction disturbances during sleep in 400 patients with sleep apnoea syndrome. *Am J Cardiol* 1983; 52:490-4.
 17. Hirschowitz M, Karacan I, Gurakar A, Williams RL. Hypertension, erectile dysfunction and occult sleep apnoea. *Sleep* 1989; 12:223-32.
 18. Partinen M, Guilleminault C. Daytime sleepiness and vascular morbidity at seven-year follow-up in obstructive sleep apnoea patients. *Chest* 1990; 97:27-32.
 19. Kreiger J, Sforza E, Apprill M, Lampart E, Weitzenblum E, Ratomaharo J. Pulmonary hypertension, hypoxaemia and hypercapnia in obstructive sleep apnoea patients. *Chest* 1989; 96:729-37.
 20. Indications and standards for cardiopulmonary sleep studies. *Am Rev Respir Dis* 1989; 139:559-68.
 21. Hillerdal G, Hetta J, Lindholm CE, Hulcrantz E, Boman G. Symptoms in heavy snorers with and without obstructive sleep apnoea. *Acta Otolaryngeal Stockh* 1991; 111:574-81.
 22. Viner S, Szalai JP, Hoffstein V. Are history and physical examination a good screening test for sleep apnoea? *Ann Intern Med* 1991; 115:356-9.
 23. Stradling JR. The sleep study, recording and analysis in: *Handbook of sleep-related breathing disorders*. Stradling J.R. (ed) Oxford Med Publ UK 1993; 87-116.
 24. Young T, Palta M, Dempsey J, Skatrud J, Weber S, Badr S. The occurrence of sleep-disordered breathing among middle-aged adults. *N Eng J Med* 1993; 328:1230-5.
 25. Stradling JR, Warley ARH, Sharpley A, Apps M, Calverley PMA et al. Oximetry versus polysomnography in the diagnosis of sleep disorders. *J Amb Mon* 1989; 2:197-201.
 26. Luchsinger J, Garshick E, Schaul N, Hackshaw R, Schurf SM. Criteria for defining sleep disordered breathing events are non-uniform. *Sleep Res* 1990; 19:370 (abst).
 27. Stradling JR, Crosby J. Prevalence of sleep apnoea in 1001 men aged years 35-65 in: Horne J. (ed) *Sleep '90*. Bochum Pontenagel Press 1990; 170-3.
 28. Johns MW. A new method for measuring daytime sleepiness: the Epworth sleepiness scale. *Sleep* 1991; 14:540-5.
 29. Taylor L, Santiago S, Williams A. Correlation of the level of nasal CPAP with body weight. *Am Rev Respir Dis* 1991; 143 (No 4, pt 2): A591 (Abst).
 30. Rajagopal KR, Bennett LL, Dillard TA. Overnight nasal CPAP improves hypersomnolence in sleep apnoea. *Chest* 1986; 90:172-6.
 31. Rogers RM, Weiler C, Ruppenthal B. Impact of the respiratory intensive care unit on survival of patients with acute respiratory failure. *Chest* 1972; 62:94-7.
 32. Strumpf DA, Millman RP, Carlisle CC, Grattan LM, Rayn SM et al. Nocturnal positive pressure ventilation via nasal mask in patients with severe chronic obstructive pulmonary disease. *Am Rev Respir Dis* 1991; 144:1234-9.
 33. Scano G, Gigliotti F, Duranti R, Spinelli A, Gorini M, Schiavina M. Changes in ventilatory muscle function with negative pressure ventilation in patients with severe COPD. *Chest* 1990; 97:322-7.
 34. Brochard L, Isabey D, Piquet J. Reversal of acute exacerbations of chronic obstructive lung disease by inspiratory assistance with a face mask. *N Eng J Med* 1990; 323:1523-30.
 35. Kribbs NB, Pack AI, Kline LR, Smith PL, Schwartz AR et al. Objective measurement of patterns of nasal CPAP use by patients with obstructive sleep apnoea. *Am Rev Respir Dis* 1993; 147:887-95.
 36. Harmon JD, Morgan W, Chaudhary B. Sleep Apnoea: Morbidity and mortality of surgical treatment. *South Med J* 1989; 82:161-4.
 37. Guilleminault C. Treatments in obstructive sleep apnoea in: *Obstructive Sleep Apnoea syndrome*. Guilleminault C, Partinen M (eds) New York Raven Press. 1990:99-118.
 38. Ferguson KA, Onon T, Lowe A, Keenan SP, Fleetham JA. A randomised crossover study of an oral appliance vs nasal positive airway pressure in the treatment of mild-moderate obstructive sleep apnoea. *Chest* 1996; 109:1269-75.
 39. Croft CB, Brockbank MJ, Wright A, Swanston AR. Obstructive sleep apnoea in children undergoing routine tonsillectomy and adenoidectomy. *Clin Otolaryngol* 1990; 15:307-14.
 40. Ryan CF, Lowe AA, Fleetham JA. Nasal continuous positive airway pressure therapy for obstructive sleep apnoea in Hallermann-Streiff syndrome. *Clin Pediatr Phila* 1990; 29:122-4.
 41. Varley AR, Mitchell JH, Stradling JR. Evaluation of the Ohmeda 3700 pulse oximeter. *Thorax* 1987; 42:892-6.
 42. Gyulay S, Gould D, Sawyer B, Pond D, Mant A et al. Evaluation of a microprocessor-based portable home monitoring system to measure breathing during sleep. *Sleep* 1987; 10:130-42.
 43. Hoffman B, Weite T. Non-invasive bilevel positive airway pressure ventilation for the treatment of the acute left ventricular failure. *Eur Resp J* 1996; 9(Supp 23): 330s (abst).

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