ORIGINAL ARTICLE

Oronasal masks require higher levels of positive airway pressure than nasal masks to treat obstructive sleep apnea

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Abstract

Purpose The purpose of this study was to compare the therapeutic pressure determined by an automated CPAP device (AutoCPAP) during the titration period, between nasal and oronasal mask and the residual apnea-hypopnea index (AHI) on a subsequent poligraphy performed with the established therapeutic CPAP.

Methods As a retrospective study, 109 subjects with moderate and severe obstructive sleep apnea-hypopnea (apnea-hypopnea index \geq 15 events/h) were studied. CPAP titration was performed using an auto-titrating device.

Results There was significant difference in the mean pressure delivered with autoCPAP between the group of patients using the nasal mask (mean 10.0 cmH₂O \pm 2.0 SD) and the group

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Prima Medicina Interna, Spedali Civili, Piazzale Spedali Civili, 1, 25123 Brescia, Italy e-mail: luciano.corda@spedalicivili.brescia.it which used the oronasal mask (mean 11.2 cmH₂O±2.1) (p<0.05). Residual apneas were lower when using a nasal mask: average AHI of 2.6±2.5 compared to 4.5±4.0 using an oronasal mask (p<0.05). On multivariate analysis, the only independent predictor of the level of therapeutic pressure of CPAP was the type of mask used (r=0.245, p 0.008).

Conclusions Therapeutic CPAP level for OSAH is higher when administered via oronasal mask, leaving more residual events. These findings suggest that nasal mask should be the first choice for OSAH treatment.

Keywords Oronasal mask \cdot Nasal mask \cdot CPAP \cdot OSAH

Abbreviations

SD	Standard deviation
OSAH	Obstructive Sleep Apnea-Hypopnea
AHI	Apnea–Hypopnoea Index
AutoCPAP	automated CPAP device
BMI	Body Mass Index
ESS	Epworth Sleepiness scale
CPAP	Continuous Positive Airway Pressure

Introduction

Obstructive Sleep Apnea-Hypopnea (OSAH) is a sleeprelated breathing disorder characterized by full or partial occlusion of upper airways during sleep that can lead to repeated oxyhemoglobin desaturations, inspiratory efforts, and sleep fragmentation.

Many studies have shown that untreated OSAH is associated with an increased risk of cardiovascular and cerebrovascular diseases [1–6].

Presently, the main treatment for moderate-to-severe OSAH is continuous positive airway pressure (CPAP), which acts as a pneumatic splint by raising the intraluminal (transmural) pressure above a critical pressure of the upper airway (Pcrit) [7], keeping the upper airway open and preventing obstructive respiratory events [8].

CPAP is delivered to the patient via a hose and a mask. There are many different mask models, generally classified as the ones which only cover nose (nasal mask), nose and mouth (oronasal mask), or entire face [9].

CPAP is virtually always effective in OSAH treatment, but its primary limitation is the low compliance rate [10–12]. Yaggi et al. found that only 58 % of patients with OSAH met minimum criteria for required compliance with CPAP (at least 4 h per night for at least five nights a week) [1].

The reasons for poor compliance are mask discomfort; concomitant pathologic condition such as depression or claustrophobia; side effects due to CPAP therapy such as mucosal dryness, air leaks, nasal congestion, or obstruction; and skin irritation. Mask discomfort plays an important role in poor compliance which is related to the type of mask that the patient (or the physician) chooses on initial CPAP titration session [13]. Although clinicians generally consider that the choice of the interface is crucial for the success of CPAP treatment, so far, only very few studies have specifically looked at the impact of different mask types on CPAP efficacy, and a 2006 Cochrane meta-analysis [14] was not able to establish the optimum form of CPAP delivery interface due to lack of data about use of different masks in OSAH patients.

The aim of this study was to evaluate, in a subgroup of moderate-to-severe OSAH patients, the possible effect of nasal and oronasal masks on the therapeutic pressure determined by an automated CPAP device [15] (AutoCPAP) during the titration period and the residual apnea-hypopnea index (AHI) on a subsequent poligraphy performed with the established therapeutic CPAP.

Materials and methods

This retrospective study was conducted in the Sleep Laboratories of the Respiratory Medicine Department at Spedali Civili Hospital, Brescia, Italy and M. Mellini Hospital, Chiari, Italy.

Subjects suspected for having OSAH were referred to our laboratories from May 2012 to June 2013 and were evaluated with all-night sleep study (Siesta Compumedics Polysomnographs) at baseline. For all patients, the following data were recorded: age, sex, Body Mass Index (BMI), neck and abdominal circumference, Epworth Sleepiness Scale (ESS) [16], Apnea-Hypopnea Index (AHI), and average and minimum oxyhemoglobin saturation (SaO₂ %).

Respiratory events were classified according to the AASM criteria [17]: an apnea was defined as a >90 % decrease in peak nasal pressure than baseline, lasting at least 10 s. Hypopnea was defined by a >30 % decrease than baseline nasal pressure, lasting at least 10 s and associated with a \geq 4 % oxyhemoglobin saturation drop. Arterial oxygen saturation was continuously measured by a finger tip infrared pulsoximetry and one ECG derivation was also monitored.

On the basis of clinical history and symptoms, patients with an AHI≥15 were considered eligible for CPAP treatment and were enrolled in the study. Exclusion criteria were the presence of comorbidities such as heart failure, COPD, obesity hypoventilation syndrome, history of nasal or paranasal disease, and anatomical problems precluding the use of a nasal or oronasal mask.

All patients included in the study underwent a home titration period of three to four nights using Philips-Respironics System One REMstar AutoCPAP with heated humidification. Before the first titration night, patients wore different masks models, choosing the more comfortable ones.

Patients were divided into two groups according to their choice between nasal (Philips-Respironics ComfortGel and EasyLife, Fisher & Paykel FlexiFit) and oronasal (Philips-Respironics ComfortGel Full and ComfortFull, ResMed

	All patients $(n=109)$	Nasal mask (<i>n</i> =67)	Oronasal mask $(n=42)$	Two-tailed <i>p</i> value
Male (n)	85	50	35	ns
Age (years)	58.3±12.9	57±12.8	60±13.1	ns
BMI (Kg/m ²)	33.9±6.3	33.7±6.9	34.3±5.2	ns
ESS	$10{\pm}4.9$	$10{\pm}4.6$	$10{\pm}5.2$	ns
Baseline AHI (events/h)	41.1±20.5	45.3±18.5	50.1±23.3	ns
Residual AHI (events/h)	3.3±3.3	$2.6{\pm}2.5$	4.5±4.0	< 0.05
Baseline mean SaO ₂	91±3.9	91 ±3.5	90±4.3	ns
Baseline minimum SaO2 %	74±10.5	75±10.5	72±10.5	ns
Mean SaO ₂ % on CPAP	94±1.8	94±1.6	94±2.0	ns
Minimum SaO2 % on CPAP	87±5.2	87±5.1	85.4±5.4	ns
CPAP pressure	10.5±2.1	10.0±2.0	11.2 ± 2.1	< 0.05

Table 1Demographic and descriptive data of the patients

Mirage Quattro) masks. The most appropriate mask size was applied in each case to minimize leakage. The patients wore the mask connected with AutoCPAP for 2 h in the afternoon adaptation session.

After the home titration period, 90th percentile (therapeutic) AutoCPAP pressure and average unintentional leak were downloaded from AutoCPAP.

Patients underwent a second polisomnographic study on therapeutic CPAP pressure and residual AHI was calculated.

The study was approved by the local Ethic Committee (Spedali Civili, Brescia, Italy; no number available).

Statistical analysis

Continuous variables were compared using two-tailed, unpaired *t* tests for variables with normally distributed data and Mann–Whitney test for variables with non-normally distributed data. Fisher's exact test was used to compare categorical data between different groups of subjects. Relationships among variables were analyzed using linear regression where appropriate. A two-tailed *p* value of <0.05 was considered statistically significant. The variables for which *p* was less than 0.05 according to univariate analysis were entered into a multivariate model to identify the independent variables that influence the CPAP pressure. Data are expressed as mean±SD. Statistical analysis was performed using SPSS 18.0 (SPSS; Chicago, IL) and Graph Pad Prism 5.0 (MacKiev Software, Boston, MA).

Results

One hundred and nine patients were included in this retrospective study. Table 1 shows the demographic and descriptive data of patients taken as a whole group and divided between patients who chose nasal and oronasal masks. Nasal masks were preferred in 67 cases (61.5 % of total), oronasal masks in 42 cases (38.5 % of total).

There were no differences between the two groups in terms of gender distribution, age, BMI, baseline AHI, mean and minimum $SaO_2\%$ both at baseline, and on CPAP polisomnography. Patients who chose oronasal masks had higher residual AHI at

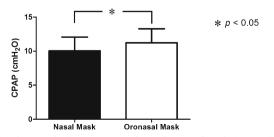


Fig. 1 Therapeutic CPAP pressure for each type of mask (mean±SD)

 Table 2
 Significant variables correlated with therapeutic CPAP pressure on univariate analysis

	Correlation coefficient (<i>r</i>)	Two-tailed p value
Gender		ns
Age (years)		ns
BMI (Kg/m ²)	0.22	< 0.05
ESS		ns
Baseline AHI (events/h)	0.22	< 0.05
Baseline mean SaO ₂		ns
Baseline minimum SaO2 %		ns
Type of mask	0.26	< 0.05

polisomnoghraphy on CPAP despite a significantly higher therapeutic pressure determined during titration session (Fig. 1).

Considering the therapeutic CPAP as dependent variable, univariate analysis revealed a significant relationship with BMI, baseline AHI, and mask type. Age, gender, ESS, and mean or minimum SaO₂ were not related to the therapeutic CPAP (see Table 2). The multivariate analysis which included the three significant variables described above showed that the only significant independent correlate of the therapeutic CPAP was the type of mask used (r=0.245, p 0.008) (see Table 3).

Discussion

Continuous positive airway pressure is the most effective treatment for OSAH [18] but the patient's compliance with CPAP treatment is surprisingly low [11, 19]. Among different reasons of discontinuation, one of the most relevant is the mask discomfort.

Due to the limited number of studies available comparing various interface types, there are no specific recommendations on the initial choice of interface. The physician, however, should evaluate both the most effective mask and patient comfort.

Oronasal masks are frequently used because of presumptive excessive mouth leak while using nasal masks, but some studies showed that the number of residual respiratory events was significantly higher with oronasal masks than with nasal masks [20, 21].

 Table 3
 Significant variables correlated with therapeutic CPAP pressure on multivariate analysis

	Correlation coefficient (r)	Two-tailed p value
BMI (Kg/m ²)		ns
Baseline AHI (events/h)		ns
Type of mask	0.245	0.008

In two studies comparing oronasal and nasal masks on auto-titrating CPAP, Teo et al. and Bakker et al. found no significant differences on the final pressure of CPAP between the two types of mask [20, 21]. However, these studies were limited by small samples size.

In contrast, we showed in this retrospective study that CPAP delivered through a nasal mask is effective for moderate to severe OSAH requiring significantly lower pressures than oronasal mask. Moreover, residual apneas decrease using a nasal mask.

In previous studies [22–25], higher comfort was reported with nasal mask. Accordingly, also in this work, nasal mask was preferred in 61.5 % of cases.

Wearing oronasal masks is associated with an increase in upper airway resistance [26] due to the posterior displacement of the tongue and this might be the cause for higher residual respiratory events than those observed during CPAP treatment. In line with the increase in upper airway resistance induced by oronasal masks, we and others found that oronasal masks were associated with higher therapeutic pressure than nasal ones. We speculate that a higher CPAP pressure is required to counteract the posterior displacement of the mandibula induced by an oronasal mask with increasing risk of unintentional leaks. This may contribute to further reduce CPAP compliance.

For these reasons, in patients with sleep apnea, it seems to be useful to propose a nasal mask from the beginning to improve adherence to CPAP and treatment efficacy.

Conclusions

In summary, the choice of the interface between the patient and the ventilator appears crucial. Since a higher level of therapeutic CPAP is required, on average, with the oronasal mask, the first attempt should be always made with the patient wearing a nasal mask. This could decrease patients' discomfort and consequently increase the CPAP compliance.

Conflict of interest The authors have no conflict of interest to disclose. No financial support, off label, or investigational use received or performed.

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