The association between dysphagia and OSA

Disfagia e OSA

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SUMMARY

Objective. The aim of our study was to investigate the presence of dysphagia in patients with Obstructive Sleep Apnoea (OSA) and to correlate swallowing impairment with hypnologic and anatomic parameters.

Methods. The study population includes 36 patients suffering from OSA. Patients were divided into two groups using the presence of dysphagia as a distinctive parameter. Group 1 included 27 OSA patients without signs of dysphagia and Group 2 included 9 OSA patients with signs of dysphagia.

Results. The age of patients in Group 2 was higher compared with the age of patients in Group 1. Analysis of Continuous Positive Airway Pressure (CPAP), obtained in the titration phase, showed that OSA patients with signs of dysphagia required a higher level of CPAP pressure than those who were not affected by swallowing abnormalities (12.6 ± 1 vs 10.5 ± 1.9 p = 0.003). No other differences in anthropometric, hypnologic, or arterial blood gas values were found between the two groups.

Conclusions. In clinical practice, all OSA patients should undergo a complete ENT exam, including assessment of swallowing, before CPAP therapy is started. This may predict the need for higher CPAP pressure settings to resolve apnoea episodes in the presence of dysphagia as well as guide the choice of CPAP interfaces (orofacial vs. nasal) in these patients.

KEY WORDS: obstructive sleep apnoea syndrome, dysphagia, swallowing impairment, CPAP

RIASSUNTO

Obiettivo. Il nostro studio ha avuto come obiettivo la ricerca nei pazienti OSA di evidenze di disfagia e l’associazione della presenza di disfagia alla anatomia delle prime vie aero-digestive e ai parametri ipnologici ed emogasanalitici di questi pazienti.

Metodi. Tutti i pazienti sono stati sottoposti a emogasanalisi, spirometria, rinomanometria, videofibrorinolaringoscopia, valutazione fibroendoscopica della deglutizione (FEES), polisomnografia notturna, studio di titolazione della pressione di lavoro di CPAP, somministrazione del questionario “SWAL-QOL”.

Risultati. Il 25% della popolazione studiata ha presentato segni subclinici di disfagia. Il gruppo dei pazienti senza disfagia era formato da soggetti più giovani rispetto al gruppo dei pazienti disfagici. Il distretto orofaringeo è risultato essere il sito di ostruzione più frequente per entrambi i gruppi. Nello studio di titolazione della pressione di lavoro di CPAP, i pazienti disfagici necessitano di valori di pressione più alti.

Conclusioni. Un paziente OSA su 4 ha presentato segni di disfagia. La disfagia è associata a valori più elevati di pressione allo studio di titolazione della CPAP. I pazienti OSA hanno un rischio più elevato di presentare disturbi di deglutizione con l’avanzare dell’età rispetto alla popolazione generale.

PAROLE CHIAVE: sindrome delle apnee ostruttive nel sonno, disfagia, disturbi di deglutizione, CPAP

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Introduction

Obstructive Sleep Apnoea (OSA) is characterised by repeated episodes of upper airway occlusion, associated with oxyhaemoglobin desaturations and brief awakenings from sleep (arousals) 1. Due to sleep fragmentation, signs and symptoms of OSA include snoring, abrupt awakenings accompanied by gasping or choking and excessive daytime sleepiness. This latter condition inevitably leads to an increase in cardiovascular risk (risk of arterial hypertension, myocardial infarction and cerebral ischaemia) 2 and worsening of the subject’s quality of life, as well as an obvious increase in the frequency of road or occupational accidents 3. The disruption of the oro-nasal airflow lasts, by definition, for at least 10 seconds and can be complete (apnoea) or partial (hypopnoea) 4.

The severity of OSA is scored according to the number of apnoeas or hypopnoeas, recorded during the study, per hour of sleep (apnoea/hypopnoea index - AHI). OSA is therefore classified as mild (AHI > 5, but < 15 per hour), moderate (AHI > 15, but < 30 per hour) and severe (AHI > 30 per hour) 5. During nocturnal respiratory events, the size of the upper airway depends on the balance between forces that would collapse the airway (such as negative intraluminal pressure and increased tissue-extraluminal pressure) and those that maintain airway (contraction of pharyngeal dilator muscles). The most frequent sites of pharyngeal collapse are the soft palate, lateral pharyngeal walls, palatine tonsils and base of the tongue. The larynx can also be involved as a site of obstruction.

During sleep, the compensatory tonic input to the motor neurons of the pharyngeal dilator muscle is lost, causing collapse of the pharyngeal airway until chemoreceptor stimuli activate the neuromuscular compensatory functions that maintain airway patency. The use of CPAP or surgical treatments of OSA reduce collapse in the upper airway (contraction of pharyngeal dilator muscles). The exclusion criteria included age > 75 years and < 45 years, daytime respiratory failure, Overlap Syndrome (Chronic Obstructive Pulmonary Disease and OSA), obesity hypoventilation syndrome (OHS), neuromuscular disorders, neurological diseases, gastroenterological diseases, craniofacial malformations, decompensated clinical and psychiatric illnesses and patients who had used CPAP therapy in the preceding three months.

The complications of OSA may include cognitive impairment, depression, nocturnal oesophageal reflux and nocturia 10. Some studies have shown that patients with OSA may be affected by swallowing dysfunctions, which reflects the abnormal function of nerves and muscles in the suprapharynx 11. The aim of the present study was to investigate the presence of oropharyngeal dysphagia in patients with OSA and to correlate swallowing impairment with hypnologic, anatomic and arterial blood gas values.

Materials and methods

Thirty-six OSA patients from the Sleep Disorder Centre of “Department of Respiratory Disease” were recruited and assessed at “ENT Clinic”, University of Bari. Of the 36 patients enrolled, 26 were males and 10 females (respectively, 72.2% and 27.8%). The exclusion criteria included age > 75 years and < 45 years, daytime respiratory failure, Overlap Syndrome (Chronic Obstructive Pulmonary Disease and OSA), obesity hypoventilation syndrome (OHS), neuromuscular disorders, neurological diseases, gastroenterological diseases, craniofacial malformations, decompensated clinical and psychiatric illnesses and patients who had used CPAP therapy in the preceding three months.

Participants were approached and informed about the study objectives and significance, before confirming their availability to take part in the present research. All patients underwent the following tests:

- recent and remote pathological anamnensis, family history and pharmacological anamnensis. Patients were also interrogated about dietary and other habits, comorbidities and general symptoms of sleep disorders.
- Body Mass Index (BMI);
- Epworth Sleepiness Scale (ESS);
- arterial blood gas analysis;
- spirometry;
- full-night cardiorespiratory polygraphy recording (Nox-T3, Nox Medical Inc. Reykjavik, Iceland). The following data were collected: snoring sound, body position, activity, heart rate, nasal pressure, oxyhaemoglobin saturation, and thoracic and abdominal movements. The hypnological parameters obtained were evaluated by a physician experienced in sleep breathing disorders;
- clinical examination to assess the anatomical characteristics of the upper airways [Friedman tongue posi-
tion staging system (FTP), tonsil hypertrophy grading scale)\(^1\);

- rhinomanometry;
- fiber optic nasopharyngolaryngoscopy of the upper airways, during which a modified Muller maneuver at oropharyngeal and hypopharyngeal levels was carried out according to the nose oropharynx hypopharynx and larynx (NOHL) classification\(^13\);
- fiber optic endoscopic evaluation of swallowing (FEES) using boluses of different texture: liquid, semisolid and solid bolus. We analysed the anatomical structures and the functional aspects of the swallowing events, according to the following criteria: (a) premature oral leakage, if the bolus entered the pharynx without eliciting the swallowing reflex; (b) laryngeal penetration if the bolus penetrated the laryngeal vestibule but not below the vocal cords; (c) tracheal aspiration if the bolus passed below the vocal cords; lastly, (d) pharyngeal stasis if the bolus remained in the valleculae and/or pyriform sinuses after the swallowing sequence, using the Yale Pharyngeal Residue Severity Rating Scale\(^14\). The Yale Pharyngeal Residue Severity Rating Scale is a standardised, anatomically defined, image-based assessment of post-swallow pharyngeal residue, based on a 5-grade rating of the amount of residue in two locations, the vallecula and the pyriform sinus as observed during fiberoptic endoscopic evaluation of swallowing;
- the Italian version of SWAL-QOL questionnaire on quality of life in dysphagia\(^15\);
- CPAP titration study.

The results are presented as percentages for categorical variables and as mean ± standard deviation (SD) for continuous variables. The proportions were compared using the Pearson \(\chi^2\) test. The means were compared using the two-tailed Student’s t-test. A p value < 0.05 was considered as significant.

### Results

The clinical and demographic characteristics of the 36 patients in this study are presented in Table I. Oropharyngeal examination revealed that 53% of subjects (n = 19) had grade I, 31% had grade II (n = 11) and 16 (n = 6) had grade III tonsil size.

According to the Friedman tongue position staging system, 11% of the cases (n = 4) were classified as FTP stage I, 17% (n = 6) as stage 2, 39% (n = 14) as stage 3 and 33% (n = 12) as stage 4.

The results of FEES demonstrated that a post-swallow residue with solid bolus texture was present in 25% of patients (n = 9). According to the Yale Pharyngeal Residue Severity Rating Scale, none/trace grade of severity of vallecular residue was observed in 2 subjects (22.2%) mild grade of severity of vallecular residue was observed in 3 (33.3%) and moderate grade of severity in 4 (44.4%) subjects, but no participant showed severe vallecular residue.

As for the pyriform sinus location, 3 (33.3%) respondents showed none/trace, and 4 patients (44.4%) had mild severity of residue. While the remaining 2 (22.2%) subjects had moderate severity of residue, no individuals with severe residue in the pyriform sinuses were observed.

In the group of patients who presented bolus residue after swallowing, 8 patients had severe OSA (88.9%) and 1 patient had moderate OSA (11.1%).

No cases of laryngeal penetration or tracheal aspiration were observed. We used the NOHL classification system and found that the oropharyngeal level was the most frequent obstruction site (average value 2.5 ± 1.13) followed by the hypopharynx level (average value = 1.62 ± 1.04), the larynx (average value 0.26 ± 0.57) and nasal level (average value 0.18 ± 0.39) (Fig. 1).

Using the presence of dysphagia as a distinctive parameter, the cohort of patients was divided into two groups. Group I included 27 patients with OSA without signs of dysphagia and Group 2 included 9 patients with diagnosis of OSA and signs of dysphagia.

### Table I. Demographic and clinical characteristics of the cohort.

| Patients (n) | 36 |
|-------------|----|
| Age (years, Mean ± SD) | 60.7 ± 11.1 |
| Male gender n (%) | 26 (72.2%) |
| BMI (Mean ± SD) | 33.6 ± 6.9 |
| ESS (Mean ± SD) | 7 ± 5 |
| OSA severity n (%) | | |
| Mild | 3 (8%) |
| Moderate | 6 (17%) |
| Severe | 27 (75%) |
| CPAP Pressure (Mean ± SD) | 11.1 ± 1.9 |
| Tonsil volume n (%) | | |
| Grade 1 | 19 (53%) |
| Grade 2 | 11 (31%) |
| Grade 3 | 6 (16%) |
| Friedman tongue position staging system n (%) | | |
| Grade 1 | 4 (11%) |
| Grade 2 | 6 (17%) |
| Grade 3 | 14 (39%) |
| Grade 4 | 12 (33%) |
| Post-swallowing stasis at FEES n (%) | 9 (25%) |

SD: standard deviation; BMI: Body Mass Index; ESS: Epworth Sleepiness Scale; OSA: Obstructive Sleep Apnoea Syndrome; CPAP: continuous positive airway pressure; FEES: Fibre-optic endoscopic evaluation of swallowing.
Anthropometric, hypnologic and arterial blood gas values were compared between the two groups: Patients in Group 2 (dysphagia) were older than patients in Group 1 (no dysphagia). Statistically significant differences were not detected in other items (Tab. II).

No significant differences were found between the two groups with respect to the grading of tonsil size, Friedman tongue position staging system, sites of upper airway collapse, or nasal resistances (Tab. III).

There was also no significant difference between the SWAL-QOL scores of patients with and without signs of dysphagia (Tab. IV).

Finally, we analysed the CPAP pressures obtained in the titration phase in the 2 groups: OSA patients with signs of dysphagia required higher level of CPAP pressure than OSA patients not affected by swallowing abnormalities (p = 0.003) (Tab. V).

Discussion

Although our sample size is not large, the OSA patients enrolled (average age 60.7 years) showed a prevalence of dysphagia of 25%. The recent literature has already highlighted a prevalence of swallowing disorders between 16% and 78% in the population affected by OSA under the age of 75 years, which is consistent with our result. Furthermore, our data show that the perception of swallowing abnormalities among the same type of patients appears to have little-to-no significance.

Our patients completed autonomously the Italian version of the SWAL-QOL test before FEES examination: the SWAL-QOL scores of patients with and without signs of dysphagia (Group 1 and Group 2, respectively) showed no significant difference between the two groups, suggesting that patients who suffer from OSA may present subclinical manifestations of abnormal swallowing. Schindler et al. enrolled 72 patients (54 men and 18 women); all patients underwent fibre optic evaluation of swallowing using boluses of different textures and each patient also completed the SWAL-QOL questionnaire: 64 patients (89%) presented spillage, 20 (28%) presented piecemeal deglutition, 26 patients (36%) presented penetration and 30% presented retention. Comparing the SWAL-QOL scales in OSA patients with and without signs of dysphagia, they found a small difference in only two spheres (general burden and symptoms) of the SWAL-QOL questionnaire. These studies show that the evaluation of dysphagia cannot be performed exclusively using self-assessment questionnaires, but...
that performing instrumental investigations is crucial. In particular, 41 heavy snorers and 15 non-snoring volunteers were video-radiographically examined by Jaghagen et al. with regards to the oral and oropharyngeal phases of swallowing. The most common finding of abnormal pharyngeal swallowing function in snorers was repeated premature bolus leakage that intermittently continued down to the swallowing reflex was elicited (51%), followed by pre-swallowing function in snorers was repeated premature bolus leakage. The most common finding of abnormal pharyngeal swallowing dysfunction remained asymptomatic. Recent study shows that more than 30% of OSA patients seem to have a poor muscle responsiveness, as measured by genioglossus electromyography. The success of therapy in this type of OSA patients with dysphagia should include careful follow-up to ensure a good compliance of the patient, despite the high CPAP pressure setting. This may also help improve the patient acceptance of CPAP therapy.

Moreover, the treatment of OSA with CPAP seems to be able to reverse the endoscopic findings of swallowing dysfunction and to improve the quality of life, according to the criteria measured by the SWAL-QOL questionnaire, as demonstrated in the study by Caparroz et al. on 70 adult patients with moderate or severe OSA. The study analysed the signs and symptoms of oropharyngeal dysphagia using FEES and SWAL-QOL in 18 patients with signs of dysphagia, of whom 12 were treated with CPAP for 3 months. The results revealed that the treatment of OSA with CPAP was able to reverse the endoscopic findings of dysphagia and improve quality of life, as measured by the SWAL-QOL test. Moreover, assessing the presence of dysphagia in OSA patients plays a fundamental role in prevention of one of the most frequent complications in swallowing deficiency: inhalation pneumonia. The incidence of inhalation pneumonia in the elderly rises with increasing age and Cabre et al. showed that, in a population of 134 patients, aged over 75 years, hospitalised for pneumonia, 55.2% presented signs of oropharyngeal aspiration. Dysphagia can therefore be considered a poor prognostic factor in terms of mortality but OSA alone, per se, seems to carry a greater risk of developing pneumonia. Chiner et al. demonstrated that the presence of OSA is associated with a greater risk of developing community-acquired pneumonia, and AH1 correlates with the severity of infection. This connection between OSA and the pulmonary infections provide strong evidence of the need to follow OSA patients with dysphagia, for which the infective risk is certainly greater.

**Table IV. The SWAL-QOL scores.**

| Group | Burden | Eating desire | Eating duration | Symptoms | Food Selection | Communication | Fear | Mental health | Social functioning | Fatigue | Sleep |
|-------|--------|---------------|-----------------|----------|----------------|---------------|------|---------------|-------------------|----------|-------|
| Group 1 | 9 ± 1.6 | 14.7 ± 3.8 | 9.3 ± 1.3 | 61 ± 6.9 | 8.4 ± 2.4 | 8.9 ± 2 | 17.7 ± 3.3 | 22.6 ± 4.5 | 22.8 ± 4 | 10.4± 3.3 | 7.3 ± 2.4 |
| Group 2 | 8.7 ± 2.3 | 13.4 ± 2.2 | 8.7 ± 2.1 | 59 ± 8.5 | 8.3 ± 2.3 | 9.3 ± 1.1 | 17.6 ± 3.5 | 22.8 ± 3.4 | 22.9 ± 2.4 | 8.4 ± 4.4 | 6.8 ± 2.7 |
| p     | Ns     | Ns            | Ns              | Ns       | Ns              | Ns            | Ns   | Ns            | Ns                 | Ns       | Ns    |

**SWAL-QOL questionnaire dysphagia-related quality of life concepts (Food selection, burden, mental health, social functioning, fear, eating duration, eating desire, communication) and SWAL-QOL general quality of life concepts (sleep and fatigue). Group 1: patients with OSA without dysphagia. Group 2: patients with OSA and signs of dysphagia. Ns: Not statistically significant.**

**Table V. CPAP titration.**

| Group 1 | Group 2 | p     |
|---------|---------|-------|
| CPAP titration (cmH2O) | 10.5 ± 1.9 | 12.6 ± 1 | 0.003 |

**CPAP: continuous positive airway pressure. Group 1: patients with OSA without dysphagia. Group 2: patients with OSA and signs of dysphagia.**
Conclusions
Our data demonstrate that all OSA patients should undergo a complete ENT exam before starting CPAP therapy, which includes assessment of swallowing. The importance of this recommendation lies in both the choice of the CPAP mask (orofacial vs nasal) and in the need to predict the need of high CPAP pressure setting to resolve apnoea episodes in the presence of dysphagia with the aim of obtaining satisfactory rates of adherence and tolerance to therapy. Indeed, it is acknowledged that pressures that are too high are associated with compliance issues in long-term CPAP therapy. However, the age of patients belonging to the dysphagic group was higher compared with the non-dysphagic group, and therefore age acts as a confounding factor in predicting the use of higher CPAP titration pressures. Our data also highlight that it is possible to identify a “new category” of OSA patients, “dysphagic OSA”, who need strict follow-up considering that they need higher levels of positive airway pressure in PAP therapy and they might present a higher risk of developing pneumonia compared with non-dysphagic OSA patients. Finally, our study is in line with the recent literature in otorhinolaryngology in confirming the opportunity to consider OSA as a complex pathology that requires a multidisciplinary approach for diagnosis and follow-up, in order to provide personalised and effective long-term treatment.

Conflict of interest statement
The authors declare no conflict of interest.

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Authors’ contributions
Conceptualization: LM, GC, GC, MLF, OR; Data curation: LM, GC, VDL, GC, MLF; Formal analysis: LM, GC, VDL; Writing – original draft: LM, GC, VDL, MLF; Writing – review and editing: LM, GC, VDL, GC, CS, VDN, NAAQ, GEC, OR, MLF.

Ethical consideration
The study was approved by the Institutional Review Board of teaching hospital Policlinico of Bari (protocol number 6750). The research was conducted ethically, with all study procedures being performed in accordance with the requirements of the World Medical Association’s Declaration of Helsinki.

Written informed consent was obtained from each patient for study participation and data publication.

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