Cone-beam CT analysis of patients with obstructive sleep apnea compared to normal controls

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ABSTRACT

Purpose: To evaluate the upper airway dimensions of obstructive sleep apnea (OSA) and control subjects using a cone-beam computed tomography (CBCT) unit commonly applied in clinical practice in order to assess airway dimensions in the same fashion as that routinely employed in a clinical setting.

Materials and Methods: This was a retrospective analysis utilizing existing CBCT scans to evaluate the dimensions of the upper airway in OSA and control subjects. The CBCT data of sixteen OSA and sixteen control subjects were compared. The average area, average volume, total volume, and total length of the upper airway were computed. Width and anterior-posterior (AP) measurements were obtained on the smallest axial slice.

Results: OSA subjects had a significantly smaller average airway area, average airway volume, total airway volume, and mean airway width. OSA subjects had a significantly larger airway length measurement. The mean A-P distance was not significantly different between groups.

Conclusion: OSA subjects have a smaller upper airway compared to controls with the exception of airway length. The lack of a significant difference in the mean A-P distance may indicate that patient position during imaging (upright vs. supine) can affect this measurement. Comparison of this study with a future prospective study design will allow for validation of these results. (Imaging Sci Dent 2016; 46: 9-16)

KEY WORDS: Sleep Apnea, Obstructive; Cone-Beam Computed Tomography; Sleep Apnea Syndromes

Introduction

The National Commission on Sleep Disorders estimates that minimal sleep-disordered breathing affects 7 to 18 million people in the United States, while 2 to 4 million Americans have moderate to severe disease. Obstructive sleep apnea (OSA) is characterized by repetitive episodes of pharyngeal collapse, specifically of the upper airway, in association with an increase in resistance to airflow during sleep.1-3 OSA is diagnosed when there are sleep-related clinical symptoms in the presence of at least five obstructive respiratory events per hour of sleep.4,5 Alternatively, OSA is diagnosed in the absence of sleep-related clinical symptoms when there are ≥15 obstructive respiratory events per hour of sleep.4,5 Obstructive respiratory events during sleep are reported according to the apnea-hypopnea index (AHI) or respiratory disturbance index (RDI).4 Mild OSA is defined as a RDI≥5 and <15 while moderate and severe disease are defined as a RDI≥15.4 OSA is classified as severe when the RDI is >30.4
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Table 1. Structural and non-structural risk factors for obstructive sleep apnea

| Structural risk factors | Non-structural risk factors |
|-------------------------|-----------------------------|
| Innate anatomic variations (facial elongation, posterior facial compression) | Obesity |
| Retrognathia and micrognathia | Central fat distribution |
| Mandibular hypoplasia | Male sex |
| Brachycephalic head form (associated with an increased AHI in whites but not in African Americans) | Age |
| Inferior displacement of the hyoid | Postmenopausal state |
| Adenotonsillar hypertrophy, particularly in children and young adults | Alcohol use |
| Pierre Robin, Down, Marfan, and Prader-Willi Syndromes | Sedative use |
| High, arched palate (particularly in women) | Smoking |
| | Habitual snoring with daytime somnolence |
| | Supine sleep position |
| | Rapid eye movement (REM) sleep |

hypertrophy, tongue volume, class II skeletal profile, and morphological deviations of the cervical spine), sleep-related factors, and central nervous system control over ventilation.²⁻⁶ Structural and non-structural risk factors for OSA are summarized in Table 1.

Despite the high incidence of OSA, it is estimated that up to 93% of women and 82% of men may have undiagnosed moderate to severe disease.⁵ Therefore, it is not unlikely that a patient with undiagnosed OSA may present for a routine visit in a dental office. Unfortunately, OSA is associated with multiple comorbidities including hypertension, obesity, gastroesophageal reflux, impotence, depression, and higher rates of cardiovascular and cerebrovascular morbidity and mortality.¹²,¹⁰,¹¹ Therefore, recognition of and screening for OSA during a routine dental visit is of vital importance.

Cone-beam computed tomography (CBCT) is a commonly used three-dimensional imaging technique introduced to dentistry in 1998.¹² The Next Generation iCAT CBCT unit (Imaging Sciences International, Hatfield, PA, USA), which was used in this study, is capable of acquiring quality images at a radiation dose equivalent to approximately one-half of the dose associated with conventional two-dimensional imaging (i.e. a full mouth series of intraoral radiographs).¹³ When utilizing a large field of view (FOV) protocol the upper airway is visible within the CBCT volume and thus CBCT is a useful diagnostic tool for evaluation of the airway. Assessment of the upper airway in individuals with OSA is essential, as they have reportedly smaller upper airways than individuals without OSA.⁸,¹⁴ For example, using CBCT, Enciso and colleagues found that the presence and severity of OSA is associated with a narrow lateral dimension of the airway.¹⁵ Similarly, using medical CT, Mayer and colleagues reported a decrease in the transverse width of the oropharynx in OSA subjects.¹⁶ Furthermore, evaluation of the upper airway is essential due to the reported increase in the frequency of airway collapse in individuals with narrower and longer airways.²

Due to the high incidence of OSA and widespread use of CBCT, it is possible that dental patients at risk for OSA (due to narrower and longer airways) may be identified on CBCT examinations originally acquired for other diagnostic purposes. Most of the commercial dental CBCT units currently in use acquire imaging with the patient in an upright position. For this reason, there is a need to evaluate the morphology of the airway of OSA subjects in the same manner. Unfortunately, numerous articles on the dimensions of OSA subjects have evaluated these patients in a supine position, which is not commonly applied in clinical dentistry. It has also been suggested that CBCT imaging of patients in the supine position is not appropriate because it does not simulate conditions during sleep.¹⁷

The purpose of this retrospective study was to evaluate the upper airway dimensions of OSA and control subjects using a CBCT unit, commonly utilized in clinical practice, which employs an upright patient position during image acquisition.

Materials and Methods

This was a retrospective study of CBCT imaging. All CBCT examinations were performed with the iCAT Next Generation unit (Imaging Sciences International, Hatfield, PA, USA). During image acquisition with this unit, the patient is positioned upright and seated throughout exposure. Inclusion criteria for the study subjects consisted of individuals with OSA and individuals without OSA (control group). The gender and age of the control group and OSA group subjects were matched as closely as possible. The mean age was 43.3 years in the OSA group (range 21-68) and 44.6 years in the control group (range 28-72).
The presence of OSA was confirmed with a sleep study (polysomnography (PSG) in 11 cases; home sleep test (HST) in 5 cases). All OSA subjects were diagnosed with moderate to severe OSA. In order for the CBCT scan to be included in the study, the size of the field of view needed to cover the nasopharynx and oropharynx and appropriate tongue position had to be attained. Tongue positioning plays an important role in the size of the airway, as evidenced by a decrease in airway size due to a retrodisplacement of the tongue base observed by Camacho and colleagues in supine patients. Therefore, CBCT studies displaying a posterior positioning of the tongue, as evidenced by air space between the tongue and hard palate, were not included in the study. Additional exclusion criteria consisted of edentulous patients, in whom mandibular jaw position and subsequent tongue position may be affected when the prosthesis is removed from the mouth during image acquisition, malocclusions (for example significant mandibular retrognathia or prognathia), and control subjects not matched to the age and gender of the OSA group. Application of the inclusion and exclusion criteria resulted in a total sample size of 32 (16 subjects + 16 controls). The study was approved by our University Institutional Review Board (study identification number Pro00000894).

**Radiographic Analysis**

The CBCT images were exported as DICOM (.dcm) files and then imported into the software program Analyze.

**Table 2.** Gender, age and linear measurements of width and anterior-posterior (A-P) distance in subjects with obstructive sleep apnea

| Subject identification code | Age | Gender | ± A-P (mm) | ± width (mm) |
|-----------------------------|-----|--------|------------|-------------|
| 10                          | 51  | M      | 3.6        | 16.2        |
| 15                          | 41  | F      | 8.4        | 24          |
| 11                          | 68  | M      | 4.8        | 24.6        |
| 17                          | 47  | M      | 6          | 10.2        |
| 18                          | 42  | M      | 11.4       | 25.2        |
| 19                          | 41  | M      | 7.2        | 12.6        |
| 25                          | 29  | M      | 4.2        | 14.4        |
| 26                          | 59  | F      | 1.8        | 16.8        |
| 27                          | 38  | M      | 6.6        | 3           |
| 72                          | 44  | M      | 4.8        | 15          |
| 34                          | 36  | M      | 5.4        | 10.2        |
| 44                          | 54  | M      | 4.8        | 15.6        |
| 46                          | 41  | M      | 3          | 9.6         |
| 55                          | 45  | M      | 5.4        | 17.4        |
| 57                          | 36  | M      | 4.2        | 27.6        |
| 58                          | 21  | F      | 4.2        | 10.2        |

*Obstructive sleep apnea subjects, ± Value for smallest axial slice

**Table 3.** Gender, age, and linear measurements of width and anterior-posterior (A-P) distance for control subjects

| Subject identification code | Age | Gender | ± A-P (mm) | ± width (mm) |
|-----------------------------|-----|--------|------------|-------------|
| 12                          | 54  | M      | 4.8        | 24          |
| 13                          | 35  | F      | 6.6        | 24.6        |
| 14                          | 50  | F      | 7.8        | 21          |
| 20                          | 55  | M      | 6          | 30.6        |
| 21                          | 36  | M      | 9.6        | 27.6        |
| 22                          | 35  | M      | 5.4        | 17.4        |
| 36                          | 37  | M      | 7.2        | 28.2        |
| 31                          | 65  | M      | 10.2       | 28.2        |
| 32                          | 45  | M      | 7.8        | 14.4        |
| 35                          | 54  | F      | 7.2        | 20.4        |
| 37                          | 37  | M      | 3.6        | 14.7        |
| 43                          | 33  | F      | 7.2        | 28.8        |
| 86                          | 28  | M      | 6          | 27          |
| 50                          | 48  | F      | 3          | 20.4        |
| 83                          | 29  | M      | 3.6        | 30          |
| 66                          | 72  | F      | 3          | 20.1        |

*Control subjects, ± Value for smallest axial slice

**Fig. 1.** Red area indicates the segmented airway with three-dimensional reconstruction.
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10.0 (AnalyzeDirect, Overland Park, KS, USA) for analysis of the airway. A coding system was used to anonymize the data by assigning a random two-digit number to each study. The upper airway affected by OSA is frequently defined as the soft tissue region bounded by the nasopharynx superiorly and the epiglottis inferiorly. Therefore, the oropharynx from hard palate to epiglottis was isolated from the data for analysis by manual segmentation of each axial slice from the surrounding soft tissue using thresholding, or setting the upper and lower grey level values of the area of interest (upper airway). The same individual performed all segmentation. Figure 1 demonstrates the segmented airway with three-dimensional reconstruction. Once the airway was isolated, the software computed the area (mm²) and volume (mm³) of each axial slice for the entire isolated portion of the airway. These area and volume measurements were used to identify the smallest axial slice, and width and anterior-posterior (A-P) dimension measurements were then carried out on the smallest axial slice in each CBCT study (Fig. 2). The average of multiple measurements was recorded as the value representing the width and AP dimension. All linear measurements were acquired by the same individual. The total number of axial slices segmented from the hard palate to the epiglottis were used to calculate the airway length.

Statistics

The Shapiro-Wilk test was used to confirm the assumption of normality in each group (OSA and control). If the normality assumption was valid in both groups, the unequal variance t-test was used to compare the means of the two groups. If the normality assumption was violated in either group, the Wilcoxon rank-sum test was used to compare the groups. All analyses were performed using SAS 9.4 (SAS Institute, Cary, NC, USA). All statistical tests were two-tailed and were performed using a significance level of 0.05. Data were summarized using mean ± standard deviation (S.D.).

| Variable                  | Group     | Sample Size | Mean ± SD       | Minimum | Maximum | P-value |
|---------------------------|-----------|-------------|-----------------|---------|---------|---------|
| A-P Distance (mm)         | Control   | 16          | 6.19 ± 2.20     | 3.00    | 10.20   | p > 0.05|
|                           | OSA       | 16          | 5.36 ± 2.27     | 1.80    | 11.40   |         |
| Width (mm)                | Control   | 16          | 28.59 ± 5.35    | 14.40   | 30.60   | p < 0.05|
|                           | OSA       | 16          | 15.79 ± 6.76    | 3.00    | 27.60   |         |
| Average airway area       | Control   | 16          | 253.23 ± 76.21  | 137.60  | 369.00  | p < 0.05|
|                           | OSA       | 16          | 159.29 ± 74.11  | 64.50   | 363.10  |         |
| Average airway volume     | Control   | 16          | 151.94 ± 45.73  | 82.50   | 221.40  | p < 0.05|
|                           | OSA       | 16          | 95.58 ± 44.45   | 38.70   | 217.80  |         |
| Total airway volume       | Control   | 16          | 13229.58 ± 4499.83 | 6343.70 | 21032.20 | p < 0.05|
|                           | OSA       | 16          | 8977.43 ± 3360.29 | 3597.00 | 16337.80 |         |
| Airway length (mm)        | Control   | 16          | 51.75 ± 5.92    | 41.40   | 65.40   | p < 0.05|
|                           | OSA       | 16          | 57.68 ± 6.85    | 45.00   | 72.00   |         |
Results

The Shapiro-Wilk test indicated that the assumption of normality was reasonable for A-P distance, width (lateral measurement), total airway volume, and airway length in both groups (p > 0.10 for all tests). The normality assumption was violated in the OSA group for average airway area (p = 0.0032) and average airway volume (p = 0.0032).

The data for the width, A-P distance, area, volume, and airway length in each group are summarized in Tables 2-4. The unequal variance t-test indicated that the mean width was significantly smaller in the OSA group than in the control group (p = 0.0011); however, the difference between groups in terms of mean A-P distance was not significant (p = 0.3049). The Wilcoxon rank-sum test indicated that both average airway area and average airway volume were significantly smaller in the OSA group than in the control group (p = 0.0014 for both comparisons). The unequal variance t-test indicated that the mean total airway volume was significantly smaller in the OSA group than in the control group (p = 0.0053), and that the mean airway length was significantly larger in the OSA group than in the control group (p = 0.0139). Figure 3 demonstrates the difference in upper airway width between an OSA subject and a control subject.

Discussion

The A-P distance and the width of the minimum surface area of the oropharynx are commonly used to evaluate the upper airway. A recent systematic review revealed that the most common measurements of the airway used to evaluate OSA subjects with CBCT included total volume and minimum cross-sectional area, followed by area and lateral and anterior-posterior linear measurements.
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Accordingly, in our study we evaluated the width (lateral) and A-P distance of the smallest axial cross sectional slice, the total airway volume, and the average area and volume of the airway. Additionally, we evaluated the length of the upper airway because OSA subjects have reportedly longer airways and there is an increase in the frequency of airway collapse in those with narrower and longer airways. 2

Enciso and colleagues determined using CBCT that the likelihood of OSA is 3.9 times higher when the lateral dimension of the oropharynx measures less than 17 mm. 3 The mean measurement of airway width in our subjects was 15.79 mm. Hora et al., 2007 assessed the airway in OSA patients and controls using MRI, and found the airway width to be smaller in OSA patients and in fact determined that the width of the airway at the retroglossal level was an independent predictor for OSA. 18 In our study the width of the smallest cross sectional slice of the oropharynx was significantly smaller in OSA subjects than controls, which is consistent with these published studies.

Conversely, our findings differed from those of previous studies when the anterior-posterior distance of the airway was considered. Ogawa and colleagues found that the A-P dimension was smaller in OSA subjects compared with controls, 3 and Schwab and colleagues found that the A-P dimension was significantly smaller in OSA subjects than controls when measuring the retropalatal region of the airway. 8 The differences in airway measurements in the latter study were maintained after controlling for sex, ethnicity, age, craniofacial size, and parapharyngeal fat. 8 Therefore, the smaller the retropalatal airway volume and area, the higher the risk of developing OSA. 5 In addition to smaller airway measurements, OSA subjects have been found to have a significantly longer airway relative to controls. 14 Smaller airways in OSA subjects can also be demonstrated in the treatment outcomes for OSA subjects. For example, Abramson and colleagues found that a cohort of OSA subjects that underwent maxillomandibular advancement and genial tubercle advancement showed an increase in airway volume post-surgery. 14 The majority of the patients in this study reported no symptoms after surgery and their postoperative respiratory disturbance index (RDI) at 6 months was statistically significantly lower than preoperative values. 14 Interestingly, the only airway value assessed that decreased post-surgery was airway length, although this still remained significantly greater than in control subjects. 14 Therefore, the findings in our study are consistent with the published literature.

Strengths of our study include confirmation of OSA with a sleep study (PSG in 11 cases; HST in 5 cases), the close approximation of the OSA and control subject groups for age and gender, and the criteria used for tongue positioning during CBCT scanning. The exclusion of subjects with a posterior tongue position ensured that the A-P distance measurement was not affected by the narrowing of the airway that can occur when the tongue, and consequently the soft palate, are positioned posteriorly. However, due to the retrospective design of our study and the stringent inclusion and exclusion criteria, our sample size was limited to 32, which could be considered a potential weakness of the study. Nevertheless, the sample size in our study is consistent with numerous other radiographic studies evaluating OSA. Korayem and colleagues used a sample size of 10 cases and 10 controls, with significance set at 5% and power at 70%, to evaluate OSA. 8 In a comparison of the upper airway dimensions between upright and
supine positions in OSA subjects, Tsuiki et al. in 2003 used a sample size of 15. Ogawa and colleagues used CBCT for radiographic analysis of a sample size of 10 OSA subjects and 10 controls, finding that the upper airway in OSA subjects presented with a smaller minimum cross-section area when compared to control subjects. Likewise, in two separate studies analyzing airway dimensions in surgically treated OSA subjects, sample sizes of 11 and 20 were used. Thus, the sample size used in our study is in agreement with the published literature.

The retrospective design of the study could also be viewed as a potential weakness, due to the limitations on certain information imposed by this type of study. For example, body mass index (BMI) was not available for control subjects and therefore this statistic was not available for inclusion in the analysis. Additionally, the control group was selected based upon the absence of a positive history of OSA and no report of signs or symptoms of OSA by the subject. Verification of the absence of OSA requires a sleep evaluation, which was not available due to the retrospective design of the study. A prospective study design would allow better control of the weaknesses of the current study. Therefore, this retrospective study will be followed by a prospective study design to analyze airway dimensions in OSA subjects and controls, with access to BMI and additional statistics such as blood pressure, which will allow better control over sample size and patient positioning during radiographic examination.

In conclusion, our study has demonstrated that OSA subjects have a narrower airway width, a smaller total mean airway volume, a smaller average area and volume, and a larger airway length than control subjects. Moreover, the results of our study suggest that the orientation of the patient during image acquisition (upright vs. supine) may affect the A-P distance measurement of the upper airway. A prospective study design is needed to confirm that the results of our study suggest that the orientation of the patient during image acquisition (upright vs. supine) may affect the A-P distance measurement of the upper airway. A prospective study design is needed to confirm that the orientation of the patient during image acquisition (upright vs. supine) may affect the A-P distance measurement of the upper airway.

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