Cephalometric measures correlate with polysomnography parameters in individuals with midface deficiency

Renato Fortes Bittar1, Silvio Eduardo Duailibi1,2, Gabriela Pereira Ribeiro Prado2, Lydia Masako Ferreira1,2 & Max Domingues Pereira1,2

To determine the association between cephalometric measurements and polysomnographic parameters in Brazilian patients with midface deficiency. This was a primary, clinical, observational, longitudinal, retrospective, analytical, and single-center study. Forty-eight patients with midface deficiency were divided into two groups as follows: those who underwent surgically assisted rapid palatal expansion (SARME) and those who received maxillary advancement (MA). Pre- and post-operative cephalometric and polysomnography measurements were obtained. Pearson’s correlation was used to verify the presence of any significant associations between PSG scores and cephalometric measurements. Associations between BMI (Body Mass Index) and AHI (Apnea Hypopnea Index) as well as arousals were observed. In the SARME group, associations between AHI and SNA, UAS and MP-H, arousals and SNA, and Co-A and MP-H were noted. Associations between AHI and Co-A, PoOr-A and MP-H, arousals and UAS, and between minimum saturation of O2 and SNA, SNB, and Co-A were observed in the MA group. This study demonstrates the alterations in the middle third of the face that were related to sleep disturbance. In addition, it shows the associations between the polysomnographic parameters and the cephalometric representations corresponding to the analyzed deformities and transverse or anteroposterior maxillary deficiencies.

Abbreviations

AHI Apnea Hypopnea Index
BMI Body Mass Index
DFD Dentofacial deformities
ICC Intraclass correlation coefficient
LAS Lower airspace
MA Maxillary advancement
MP-H Hyoid bone height
OSA Obstructive sleep apnea
OSAS Obstructive sleep apnea syndrome
SARME Surgically assisted rapid maxillary expansion
SED Snoring episode duration
SNA Sella-nasion-point A
SNB Sella-nasion-point B
UAS Upper airspace

Deformities of the craniofacial skeleton may be congenital or acquired in nature. Those that affect the mandible, maxilla or dentition are called dentofacial deformities (DFD)1. The degree of the DFD may vary, ranging from slight asymmetries without functional and/or aesthetic repercussions to significant changes that affect the entire functioning of the stomatognathic system, and sometimes, the respiratory system2. There is a close relationship between DFD and sleep-disordered breathing, and it is

1Postgraduate Program in Translational Surgery, Federal University of Sao Paulo, Rua Botucatu 740, 2nd floor, Vila Clementino, São Paulo, SP CEP 04023-900, Brazil. 2Division of Plastic Surgery, Department of Surgery, Federal University of Sao Paulo, São Paulo, SP, Brazil. *email: maxdp@terra.com.br
estimated that several individuals with DFD who are candidates for orthognathic surgery experience obstructive sleep apnea syndrome (OSAS). Deformities of the lower third of the face are more-commonly associated with OSA. However, the association of midface facial DFD with OSA was initially proposed in 1984; this prompted an interest in studying the relationship between maxillary deformities and OSAS14–16. The deficiency of the middle third of the face comprises three-dimensional changes that generate changes in the height, width, and anteroposterior relationship of the maxilla. These changes occur in both syndromic and non-syndromic individuals with repercussions on dental occlusion, phonation, and nasal breathing16. Changes in the anteroposterior and transverse dimensions of the maxilla cause narrowing of the nasal cavities and alterations in the base of the skull, which lead to a reduction in the nasopharynx airspace. Retro positioned maxilla can encroach the pharyngeal airway and the narrow transverse dimension increases the airflow resistance.

Initially used as an exam for the diagnosis of craniofacial deformities, cephalometry is currently used in sleep medicine to correlate craniofacial changes with the presence of OSAS. Following the association between craniofacial changes and the etiology of upper airway obstruction, several studies attempted to determine the anatomical abnormalities that might be most closely correlated; mandibular deficiency, cranial base hyperflexion, and transverse and sagittal deficiencies of the maxilla and mandible were found to be the most important factors. The hyoid bone, which provides attachment to the muscles of the floor of the mouth, is found at a lower position in patients with OSAS which leads to an increment in the MP-H (Mandibular plane to Hyoid) distance.

Thus, cephalometric measurements have been studied to demonstrate a correlation with the diagnosis and severity of OSAS, such as the presence of a value above the normal distance between the posterior nasal spine and the distal end of the uvula (PNS-P) and the distance between the mandibular plane and the hyoid bone (MP-H), both correlating with altered apnea and the hypopnea index (AHI). About 86% of OSAS patients are estimated to have reduced retrolingual airspace. Cephalometry has been used to evaluate the success of the OSAS treatment, and some studies have suggested that SNA (angle formed between the point A, N and S), lower facial height, and palate thickness may act as variables. Therefore, considering the high prevalence of individuals with DFDs, particularly, sagittal and transverse maxillary deficiencies, in the population, and the fact that this condition is often associated with the presence of OSAS, this study aimed to determine the association between cephalometric measurements and the polysomnographic parameters in Brazilian patients with midface deficiency.

Materials and methods
Study design. This is a primary, clinical, observational, longitudinal, retrospective, analytical, and single-center study.

Ethical considerations. The study was submitted and approved by the Research Ethics Committee of the Federal University of São Paulo (UNIFESP; approval number, 0178/2017).

Declarations. All the phases of the study strictly followed the Declarations of Helsinki for experiments involving humans. Informed consent was obtained from all the participants in this study and also obtained from a parent and/or legal guardian of under 18 years patient.

Patient selection. Medical records and exams of patients operated in the cranio-maxillofacial surgery department of the Plastic Surgery Discipline of the Federal University of São Paulo (UNIFESP), from 2002 to 2018.

Inclusion criteria. Patients with transverse or sagittal maxillary deficiency, with or without OSAS, who underwent surgically assisted rapid palatal expansion (SARME) or maxillary advancement (MA) were included in this study.

Criteria for non-inclusion. The following patients were excluded from the study: those with any syndromes affecting cranio-maxillofacial development; those with incomplete cephalometric, polysomnographic and operative documentation; and those undergoing any other surgical modality concomitantly.

Sample. After selecting the medical records, 48 patients were divided into two groups according to the type of surgery performed: Group 1, patients who underwent SARME (n = 25; Table 1) and Group 2, those who underwent MA (n = 23; Table 2). SARME was indicated when the transverse maxillary discrepancy was severe in sagittal dimension and beyond the scope of orthodontic camouflage. The severity of the discrepancy was assessed clinically and radiographically (cephalometry).

Lateral cephalograms were performed in all the patients at two different periods: T1 (preoperative) and T2 (6 months after surgery). Teleradiographs were scanned at a resolution of 400 dpi (dots per inch), filed in the JPEG (Joint Photographic Experts Group) format, and edited using the public domain ImageJ software (Wayne Rasband’s ImageJ) developed by the National Institute of Health.

The following cephalometric landmarks were defined: Sella (S), point located at the geometric center of the Sella turcica image; Nasion (N), point located in the most anterior region of the frontal-nasal suture, meeting the profile lines of the glabella and nasal bones; Orbitale (Or), the lowest point of orbit contour; Porion (Po), point located on the upper and outer margin of the external ear canal; Subspinatus or point A (A), point located at the deepest curve of the alveolar profile of the maxilla; Supramentonian or point B (B), deepest point of anterior mandibular concavity; Menton (Me), most anterior and inferior point of the bony chin; Gonion (Go), the lowest
and posterior point in the gonial region at the mandibular angle; Hyoid (H), the most anterosuperior point of the body of the hyoid bone; Condylum (Co), upper and posterior point of the mandibular condyle; and Tongue base (T1), most posterior point of the tongue base.

Subsequently, the following angular and linear measurements were taken: SNA, the angle formed between points S-N-A; SNB, the angle formed between points S-N-B; effective jaw length, measured between Co-A points; PoOr-A, linear measurement between the line joining Po-Or points and its orthogonal plotted to point A; NPerp-A, distance from point A to a line that passes at point N and perpendicular to the PoOr plane; Upper airspace (UAS), measurement drawn from the posterior wall of the soft palate to the region of the smallest cross-section of the posterior pharyngeal wall; Lower airspace (IAS), measured from the base of the tongue (T1) and the region of the greatest narrowing with the posterior pharyngeal wall; and Hyoid bone height (MP-H), orthogonal distance from the mandibular plane (Go-Me) and the H point (Fig. 1).

Overnight type 1 polysomnography was performed in all patients before the surgery (T1) and postoperatively (T2) at Sleep Institute Clinic, Sao Paulo – Brazil. The patients were accommodated in a dark and quiet room with a comfortable bed. Electroencephalogram (EEG), submental and tibial electromyography, right and left electro-oculogram, electrocardiography, airflow (oral and nasal), respiratory effort (chest and abdomen movements), and oxymoglobin saturation (SatO2%) were recorded with finger pulse oximetry. A microphone was used to detect snoring and lying positions. Apnea is considered as complete cessation of breathing for ≥ 10 secs and hypopnea is considered as an abnormal respiratory event for ≥ 10 s with ≥ 30% reduction in airflow with a fall of ≥ 4% of SPO2. AHI consisted of the total number of apnea or hypopneas per hour of sleep.

Statistical analysis. All reference points were plotted at three different times with 15 days difference by the same examiner to evaluate intraobserver agreement through the intraclass correlation coefficient (ICC). The average of the values of three measurements was used for each measure. The null hypothesis was that the measurements were not concordant. Pearson’s correlation test was performed to assess the relationship between the cephalometric and polysomnographic measurements for each type of surgery. Scatter plots were used for visual analysis in the case of moderate, strong, and very strong correlations. The classification proposed by DEVORE was used to estimate the power of the correlation as follows: 0–0.19, very poor correlation; 0.20–0.39, weak correlation; 0.40–0.69, moderate correlation; 0.70–0.89, strong correlation; and 0.90–1.00, very strong correlation.

| Patient | Age | BMI | Mov | AHI | Arousal | MinSatO2 | TST sat < 90% |
|---------|-----|-----|-----|-----|---------|----------|---------------|
| 1       | 31  | 24.51| 8.7 | 3   | 0       | 89       | 3.2           |
| 2       | 26  | 27.1 | 8   | 0.9 | 0       | 94       | 0             |
| 3       | 24  | 28.52| 9.2 | 0.2 | 12      | 69       | 0.3           |
| 4       | 24  | 17.19| 8.7 | 1.3 | 0       | 89       | 0.1           |
| 5       | 46  | 34.15| 8.5 | 13.4| 0       | 85       | 0.4           |
| 6       | 33  | 22.59| 7.8 | 0.4 | 14      | 95       | 0             |
| 7       | 37  | 25.91| 7   | 18.3| 6.9     | 61       | 6.1           |
| 8       | 40  | 29   | 9   | 8.1 | 25      | 91       | 0             |
| 9       | 30  | 27.94| 8.3 | 3   | 0       | 86       | 3.2           |
| 10      | 47  | 36.73| 8   | 13.2| 15      | 71       | 3.1           |
| 11      | 37  | 21.3 | 8.2 | 0   | 0       | 96       | 0             |
| 12      | 26  | 29   | 7   | 0.1 | 8       | 95       | 0             |
| 13      | 16  | 28.9 | 7.5 | 0.3 | 12      | 92       | 0             |
| 14      | 26  | 22.84| 11.3| 1.2 | 0       | 92       | 0             |
| 15      | 29  | 18.38| 8.4 | 0.4 | 0       | 93       | 0             |
| 16      | 32  | 25.2 | 9   | 2   | 9       | 92       | 0             |
| 17      | 34  | 22   | 9   | 1   | 9.9     | 97       | 0             |
| 18      | 19  | 27.6 | 7.7 | 2.6 | 7       | 95       | 0             |
| 19      | 30  | 22.72| 10.2| 0.2 | 0       | 93       | 0             |
| 20      | 29  | 38.06| 11.2| 44.8| 42.7    | 85       | 4.3           |
| 21      | 34  | 24.38| 8.8 | 6.5 | 0       | 67       | 3.8           |
| 22      | 32  | 29.06| 7.4 | 0   | 0       | 91       | 0             |
| 23      | 26  | 22.1 | 7.2 | 0   | 4.4     | 93       | 0             |
| 24      | 25  | 30.1 | 10.1| 20.7| 6       | 76       | 1.3           |
| 25      | 25  | 25.97| 7.4 | 0.3 | 2.7     | 93       | 0             |
| Average | 30.32| 24.51| 8.54| 3.68| 6.98    | 87.2     | 1.03          |
| SD      | 7.35| 5.06 | 1.18| 10.13| 9.87    | 10.14    | 1.77          |

Table 1. Characteristics of patients selected from the SARME group. Age: years; BMI: kg/m²; Mov: Millimeters; AHI: number per hour; Arousal: number per hour; MinSatO2: percentage; TST sat < 90%: percentage of the total time of sleep; SD: standard deviation.
Table 2. Characteristics of the patients in the MA group. Age: years; BMI: kg/m²; Mov. Millimeters; AHI: number per hour; Arousal: number per hour; MinSatO2: percentage; TST sat < 90%: percentage of the total time of sleep; SD: standard deviation.

| Patient | Age | BMI | Mov | AHI | Arousal | minSatO2 | TST sat < 90% |
|---------|-----|-----|-----|-----|---------|----------|---------------|
| 1       | 45  | 28  | 5   | 10.7| 17      | 90       | 0             |
| 2       | 19  | 24.22| 7   | 0.2 | 25      | 97       | 0             |
| 3       | 30  | 29.05| 6   | 2.4 | 2.4     | 95       | 0             |
| 4       | 18  | 22.4 | 5   | 0.6 | 18      | 94       | 0             |
| 5       | 39  | 26.08| 6.5 | 2.1 | 13      | 86       | 3.2           |
| 6       | 20  | 26.22| 10  | 0.2 | 0.4     | 95       | 0             |
| 7       | 34  | 25.01| 4   | 0.5 | 4.7     | 93       | 0             |
| 8       | 24  | 27.2 | 9   | 0   | 7.5     | 94       | 0             |
| 9       | 37  | 27.82| 8.5 | 2.2 | 7.8     | 85       | 1.2           |
| 10      | 47  | 26.4 | 6   | 0   | 5.7     | 92       | 0             |
| 11      | 34  | 25.4 | 10  | 18  | 45      | 91       | 0             |
| 12      | 23  | 19.8 | 9   | 0   | 15      | 98       | 0             |
| 13      | 31  | 29.06| 7   | 11.2| 22.2    | 84       | 0.5           |
| 14      | 44  | 27.1 | 7.5 | 4   | 21.2    | 95       | 0             |
| 15      | 37  | 25.07| 5   | 0   | 2.5     | 97       | 0             |
| 16      | 31  | 28.04| 6   | 11.4| 13.1    | 88       | 2.5           |
| 17      | 31  | 27.4 | 8   | 0.5 | 0       | 91       | 0             |
| 18      | 25  | 28.01| 8   | 0   | 6.9     | 92       | 0             |
| 19      | 40  | 31.22| 5   | 0   | 7.9     | 94       | 0             |
| 20      | 24  | 22.49| 9   | 1.7 | 8.2     | 93       | 0             |
| 21      | 24  | 25.4 | 8   | 0   | 4.8     | 94       | 0             |
| 22      | 21  | 23.85| 5   | 0   | 0       | 97       | 0             |
| 23      | 49  | 24.02| 7   | 0   | 11.2    | 97       | 0             |
| Average | 31.3| 26.05| 7.02| 2.86| 11.28   | 92.7     | 0.32          |
| SD      | 9.79| 2.56 | 1.75| 4.96| 10.34   | 3.94     | 0.85          |

Figure 1. S (Sella); N (Nasion); Or (Orbital); Po (Porion); A (Point A); B (Point B); Me (Mentus); Go (Gonius); H (Hyoid); Co (Condylum); T1 (Tongue base).
Initially, the association between the BMI and age of the entire sample (n = 48) with polysomnographic parameters was analyzed. Then, the analysis of the associations in each group was performed. For significant correlations, the mediation effect in multiple linear regression was tested, analyzing the interaction of BMI and age in the association.

Research involving human participants and/or animals. The study was submitted and approved by the Research Ethics Committee of the Federal University of São Paulo (UNIFESP; approval number, 0178/2017). All the phases of the study strictly followed the Declarations of Helsinki for experiments involving humans.

Informed consent. Informed consent was obtained from all the participants in this study. The informed consent was previously analyzed and approved by the Research Ethics Committee of the Federal University of São Paulo (UNIFESP).

Results
The values of the cephalometric and polysomnographic measurements at T1 and T2 in patients who underwent SARME and MA are shown in supplementary online tables.

### Intraclass correlation coefficient (ICC)

| Parameter | SNA | SNB | Co-A | PoOr-A | NPerp-A | UAS | IAS | MP-H |
|-----------|-----|-----|------|--------|---------|-----|-----|------|
| **SARME** |     |     |      |        |         |     |     |      |
| T1        | 0.999 | 0.969 | 0.999 | 1.000  | 1.000   | 0.999| 0.999| 0.999 |
| T2        | 0.986 | 0.99 | 0.993 | 0.999  | 1.000   | 0.998| 0.999| 0.998 |
| **MA**    |     |     |      |        |         |     |     |      |
| T1        | 0.999 | 0.999 | 1.000 | 1.000  | 1.000   | 0.996| 0.997| 0.998 |
| T2        | 0.992 | 0.992 | 0.997 | 0.998  | 0.999   | 0.995| 0.996| 0.997 |

Table 3. Intraclass correlation coefficient (ICC). T1: preoperative; T2: postoperative.

### Pearson’s correlation

| Entire sample | BMI (p-value) | AGE (p-value) |
|---------------|---------------|---------------|
| AHI           | 0.54 (p = 0.999) | 0.19 (p = 0.174) |
| Arousals      | 0.37 (p = 0.995) | 0.1 (p = 0.478) |
| Min SatO2     | − 0.34 (p = 0.008) | − 0.20 (p = 0.08) |
| TST SatO2 < 90% | 0.35 (p = 0.992) | 0.2 (p = 0.74) |

Table 4. Pearson’s correlation (entire sample) between BMI and AGE with AHI, arousals, minimum O2 saturation and total sleep time with < 90% O2 saturation. N = 48; AHI: Apnea Hypopnea Index; Min SatO2: minimum saturation of O2; TST SatO2 < 90%: total sleep time with saturation of O2 below 90%.

| SARME | BMI (p-value) | AGE (p-value) |
|-------|---------------|---------------|
| AHI   | 0.62 (p = 0.0008) | 0.24 (p = 0.235) |
| Arousals | 0.66 (p = 0.0003) | 0.08 (p = 0.69) |
| Min SatO2 | − 0.34 (p = 0.04) | − 0.29 (p = 0.07) |
| TST SatO2 < 90% | 0.4 (p = 0.04) | 0.31 (p = 0.235) |

Table 5. Pearson’s correlation (SARME) between BMI and AGE with AHI, arousals, minimum O2 saturation and total sleep time < 90% O2 saturation. N = 25; AHI: Apnea Hypopnea Index; Min SatO2: minimum saturation of O2; TST SatO2 < 90%: total sleep time with saturation of O2 below 90%.
correlated with preoperative SNA (r = 0.499/p = 0.011), UAS (r = −0.488/p = 0.013), and MPH (r = 0.573/p = 0.03)
and preoperative arousals were significantly correlated with preoperative SNA (r = 0.410/p = 0.042), Co-A (r = 0.491/p = 0.013), and MPH (r = 0.507/p = 0.010) in the SARME group and SNA (r = −0.415/p = 0.049), SNB (r = −0.468/p = 0.024), and CoA (r = −0.414/p = 0.049). (Online Supplementary Figures 8–14).

The mediation regression was shown to be significant with p values < 0.05 between the association SNB/MinSatO2 and MP-H/AHI in the SARME group when moderated by BMI, with p-values equal to 0.026 and 0.025, respectively.

For the MA group, the mediation regression showed an interaction of the BMI with the SNB/MinSatO2 (p = 0.042), PoOr-A (p = 0.006), and MPH (p = 0.021) preoperative associations with preoperative UAS (r = −0.440/p = 0.035); and minimum preoperative O2 saturation was significantly correlated with preoperative SNA (r = −0.415/p = 0.049), SNB (r = −0.468/p = 0.024), and CoA (r = −0.414/p = 0.049). (Online Supplementary Figures 8–14).

The mediation regression was shown to be significant with p values < 0.05 between the association SNB/arousals and MP-H/AHI in the SARME group when moderated by BMI, with p-values equal to 0.026 and 0.025, respectively.

Significant correlations were observed preoperatively in patients who underwent the MA surgical procedure (Table 8). The significant correlations were as follows: preoperative AHI was significantly correlated with preoperative PoOr-A (r = 0.424/p = 0.044), PoOr-A (r = 0.491/p = 0.013), and MPH (r = 0.507/p = 0.010). (Online Supplementary Figures 2–7).

Significant correlations were observed preoperatively in patients who underwent the MA surgical procedure (Table 8). The significant correlations were as follows: preoperative AHI was significantly correlated with preoperative PoOr-A (r = 0.424/p = 0.044), PoOr-A (r = 0.491/p = 0.013), and MPH (r = 0.507/p = 0.010). (Online Supplementary Figures 2–7).

There was no interaction between the AGE variable and the significant associations.

Discussion

This study was conducted to explore the association between the cephalometric measures and polysomnographic parameters in individuals with midface deficiency. Two polysomnographic parameters—pre-operative AHI and arousals were found to be associated with different cephalometric measures. Preoperative AHI was significantly correlated with preoperative SNA (r = 0.499/p = 0.011), UAS (r = −0.488/p = 0.013), and MPH (r = 0.573/p = 0.03) and preoperative arousals were significantly correlated with preoperative SNA (r = 0.410/p = 0.042), Co-A (r = 0.491/p = 0.013), and MPH (r = 0.507/p = 0.010). (Online Supplementary Figures 2–7).

There was no interaction between the AGE variable and the significant associations.

Table 6. Pearson’s correlation (MA) between BMI and AGE with AHI, arousals, minimum O2 saturation and total sleep time with <90% O2 saturation. N = 23; AHI: Apnea Hypopnea Index; Min SatO2: minimum saturation of O2; TST SatO2 < 90%: total sleep time with saturation of O2 below 90%.

| MA | BMI (p-value) | AGE (p-value) |
|----|--------------|---------------|
| AHI | 0.25 (p = 0.233) | 0.22 (p = 0.301) |
| Arousals | −0.10 (p = 0.310) | 0.1 (p = 0.632) |
| MinSatO2 | −0.49 (p = 0.008) | −0.25 (p = 0.121) |
| TST SatO2 < 90% | 0.18 (p = 0.04) | 0.16 (p = 0.301) |

Table 7. Pearson’s correlations between AHI, Arousals, MinSatO2, TST SatO2 < 90% and T1 and T2 cephalometric measures. SARME. Online Supplementary Figures: a-Figure 2; b-Figure 3; c-Figure 4; d-Figure 5; e-Figure 6; f-Figure 7.

| T1 | SNA | SNB | Co-A | PoOr-A | NPerp-A | UAS | IAS | MP-H |
|----|-----|-----|------|--------|---------|-----|-----|------|
| AHI | 0.499a | 0.221 | 0.290 | −0.148 | −0.365 | −0.488b | 0.092 | 0.573a |
| Arousals | 0.410b | 0.228 | 0.491c | 0.039 | −0.176 | −0.180 | 0.069 | 0.507d |
| MinSatO2 | −0.175 | 0.209 | 0.122 | 0.093 | 0.299 | 0.177 | −0.069 | −0.320 |
| TST SatO2 < 90% | 0.284 | 0.045 | 0.208 | 0.043 | −0.266 | −0.349 | 0.114 | 0.356 |

| T2 | SNA | SNB | Co-A | PoOr-A | NPerp-A | UAS | IAS | MP-H |
|----|-----|-----|------|--------|---------|-----|-----|------|
| AHI | 0.378 | 0.313 | 0.199 | −0.162 | −0.204 | −0.366 | −0.242 | 0.111 |
| Arousals | 0.060 | 0.016 | 0.246 | 0.262 | −0.196 | −0.252 | −0.147 | 0.234 |
| MinSatO2 | −0.283 | −0.173 | −0.335 | 0.263 | 0.248 | 0.284 | 0.350 | −0.049 |
| TST SatO2 < 90% | 0.258 | 0.315 | 0.105 | −0.247 | −0.144 | −0.323 | −0.272 | 0.033 |

Cephalometry does not have the technical resources to provide the data necessary to reach a diagnosis of OSAS because it involves the identification and quantification of both apnea and hypopnea. However, the identification of craniofacial changes detectable by cephalometry may help in the propaedeutic arsenal, providing
additional information about individuals more prone to OSAS, thus helping to rationalize the use of complementary exams.

Polysomnography is considered as the gold standard for the diagnosis of OSAS, however, this is a complex, uncomfortable and expensive procedure and hence not routinely performed before orthognathic surgery for the correction of dentofacial deformities, especially in the developing countries. Furthermore, polysomnography may have the "first-night" effect (i.e., the result of the first exam), which may differ significantly from the results of subsequent exams after the patient gets adapted to it. A high number of individuals with DFDs who sought surgical treatment and who, despite the presence of the symptoms of SED, had not yet been diagnosed with OSAS. This indicates that the association between DFDs and "silent" OSAS is frequent, but underdiagnosed.

In the present study, patients were evaluated for the presence of DFD with surgical indications. None of the patients underwent SARME or MA for the treatment for OSAS. Basal polysomnography was performed in all patients as part of the UNIFESP craniomaxillofacial surgery outpatient care protocol. The distribution of demographic characteristics between the groups was homogeneous, with the SARME group presenting with a mean age of 30.32 years and a BMI of 26.45 kg/m², while the MA group had a mean age of 31.30 years and a BMI of 26.05 kg/m².

Table 9. Mediation effect in multiple linear regressions of the significant associations. SQ: sums of the squares; DF: degrees of freedom; MQ: means of squares; Z: Z-score.

Table 8. Pearson's correlations between AHI, Arousals, MinSatO2, TSTSatO2 < 90% and T1 and T2 cephalometric measures. MA. Online Supplementary Figures: a-Figure 8; b-Figure 9; c-Figure 10; d-Figure 11; e-Figure 12; f-Figure 13; g-Figure 14.
Age is also defined as a risk factor for OSAS. However, in this study, the correlation obtained was not significant, perhaps because the overall mean age (30.79 years) was low and uniform between the two groups. Furthermore, AHI was positively associated with SNA (r = 0.499/p = 0.011) and MP-H (r = 0.573/p = 0.03) and negatively associated with UAS (r = −0.488/p = 0.013) in the preoperative evaluation, with no significant associations with T2 measurements in the SARME group. This finding is in agreement with the classic study by Guilleminault et al.4, and that by Armalaite et al.27. In addition, the association between AHI and MP-H has been reported by Borges et al.35, Silva et al.28 and Ryu et al.29.

The correlation between MP-H, UAS, and AHI would translate into a more caudally located larynx, which has been associated with a risk factor for OSAS, whereas a higher UAS would translate into a larger upper airway caliber and therefore, would be expected. Lower inspiratory collapse during sleep, with lower AHI. However, a more anterior maxilla would be expected to provide larger retropalatal airspace and, consequently, a smaller AHI. This finding was not verified in the current study. Unlike the study by Parhiz et al.30, the average initial SNA in the present study was 82.94, and reached 81.78 postoperatively. Brettos et al.31 observed maxillary displacement in the group that underwent SARME with Haas but not in the group that underwent SARM with Hyrax. This correlation, although significant, is not as evident in the scatter plot, and should be considered with caveats.

Arousal index was positively correlated with SNA (r = 0.410/p = 0.042), Co-A (r = 0.491/p = 0.013), and MPH (r = 0.050/p = 0.010), preoperatively. Similarly, as with AHI, the translation of MP-H into the lower larynx predisposes to obstructive events with increased AHI, and consequently, to arousals. A positive correlation between SNA would not be expected, translating a more anterior maxilla and a greater number of arousals. Another unexpected positive correlation was between Co-A, effective maxillary length, and a higher rate of arousals. The validity of such correlations may be questioned because the variation in the average Co-A was insignificant between T1 and T2 (82.70–82.12). Nevertheless, the scatter plots in this study did not suggest a significant correlation. Such associations may have occurred in isolation in this sample, but the fact that the anteroposterior dimension of the maxilla may be correlated with obstructive events and arousals in patients with transverse maxillary deficiency warrants further evaluations. One might speculate that this enlarged anteroposterior dimension could result in a more constricted maxilla thus, justifying the obstructive events and arousals.

Changes in the lateral wall of the oropharynx and the narrowing of its caliber have already been suggested by Cahali et al.32 as a possible predisposing factor for OSAS, having served as the basis for the development of the lateral pharyngoplasty technique. Such changes in transverse length could also be associated with the development of obstructive sleep events in more cranially located anatomical sites, such as nasal cavities and nasopharynx.

Significant correlations were observed preoperatively in patients belonging to the MA group. AHI was positively correlated with CoA (r = 0.424/p = 0.044), PoOr-A (r = 0.551/p = 0.006), and MPH (r = 0.479/p = 0.021). In this case, an increase in the distance from point A to the Frankurt line (an increase in the vertical dimension of the maxilla) was observed, which may translate into dolichocephalic individuals and might be a predisposing factor for obstructive events, as suggested by Kubota et al.32. As mentioned previously, an increased MP-H, representing a lower-located larynx, was associated with obstructive events in patients with decreased anteroposterior maxillary dimensions. However, Co-A, which represents the effective maxillary length, was positively correlated with AHI (mean value, 83.46–88.93). Such a correlation was unexpected because its increase following surgery should have been positively correlated with the postoperative values since there was a slight increase in AHI between T1 and T2, but the persistence of this correlation was not observed. The trend line in the scatter plot is influenced by an outlier and that the other group together without such a clear correlation.

There was a negative correlation between the arousals index and the UAS (r = −0.488/p = 0.013); therefore, the higher the index the lower the number of arousals per hour. Similar to the SARME group, a larger UAS provides a clearer airway without the propensity for collapse. Such an association might be expected with AHI, a fact that was not verified in this study.

The mean maxillary advancement was 7.02 mm, which was higher than that presented by Hellak et al.33. Furthermore, the average increase in UAS was 27.05% in the MA group; this is lower than that (37.70%) reported by Hernández-Alfaro et al.34. Likewise, Rosario et al.35 observed a significant increase in UAS volume, but question the sustainability of this increase in the long term.

Despite the use of modern technological imaging resources, the delimitation of anatomical points can prove challenging and may lead to inaccuracies in the method. Some anatomical landmarks such as Porion, Condylum, and Gonion have low reliability in the inter-observer analysis, regardless of the degree of experience of the examiner36. Although cephalometric measurements have good reproducibility, measurement errors may occur due to difficulties in identifying certain anatomical landmarks. The values of ICC for all the cephalometric variables in this study are above 0.96 and any value above 0.9 is considered to have excellent reliability37.

In the present study, the Sella (S), Nasion (N), Supramentonian (B), Hyoid (H), and Mento (M) were relatively easy to identify. However, the other demarcated points may present with relative difficulties due to the presence of overlapping images and the rotation of the patient’s head. Nonetheless, despite the limitations concerning the identification of the anatomical landmarks and the potential risk of variations in the interobserver measurements, cephalometry is a highly relevant method used in medical and dental practice because it is easily acquired, has low exposure to ionizing radiation, and is cost-effective. Furthermore, this method has gained importance in detecting the skeletal changes related to various conditions, such as OSAS, and its use has extended to the detection of areas of upper airway narrowing.

Finally, negative correlations were observed between SNA, SNB, Co-A, and the minimum O2 saturation; however, they were not evident in the scatter plots. The significant correlations obtained were of moderate degree; however, MP-H may prove to be a promising cephalometric measure that can be used to demonstrate an association with OSAS, as demonstrated by Hoekema et al.38. Further studies that associate such measures with validated questionnaires for OSAS or excessive daytime sleepiness (Berlin or Epworth, respectively) can be sought as a way to increase the predictive value for OSAS. Moreover, studies that, through multiple regression,
seek mathematical equations that determine the predictive value of surgeries or the predictive value for OSAS detection, as performed by Kim et al. Could provide relevant information.

Conclusions
The findings of this study showed that BMI was correlated with AHI and arousals. In the SARMEM group, correlations between BMI and BMI AHI, BMI and BMI MP-H, arousals and BMI, and BMI AHI and BMI BMI H were observed. In the MA group, correlations between BMI and BMI Co-A, BMI and BMI MP-H, arousals and BMI, and between the minimum saturation of BMI O2 and BMI SNA, BMI SNB, and BMI Co-A were observed.

Received: 21 May 2020; Accepted: 8 March 2021
Published online: 12 April 2021

References
1. Sato, F. R. et al. Prevalence and treatment of dentofacial deformities on a multiethnic population: A retrospective study. Oral Maxillofac. Surg. 18(2), 173–179 (2014).
2. Trench, J. D. A. & de Araújo, R. P. C. Deformidades dentofaciais: Características miofuncionais orofaciais. Revista CEFAC. 17, 1202–1214 (2015).
3. Posnick, J. C. et al. “Silent” sleep apnea in dentofacial deformities and prevalence of daytime sleepiness after orthognathic and intranasal surgery. J. Oral Maxillofac. Surg. 76(4), 833–843 (2018).
4. Guilleminault, C., Riley, R. & Powell, N. Obstructive sleep apnea and abnormal cephalmatic measurements. Implications for treatment. Chest 86(5), 793–794 (1984).
5. Cistulli, P. A. Craniofacial abnormalities in obstructive sleep apnoea: implications for treatment. Respiratory 1(3), 167–174 (1996).
6. Menon, S. et al. Enlargement of the pharynx resulting from surgically assisted rapid maxillary expansion. J. Oral Maxillofac. Surg. 74(2), 369–379 (2016).
7. Menon, S. et al. Correction of midface deficiency using intra-oral distraction device. J. Maxillofac. Oral Surg. 9(1), 57–59 (2010).
8. Menon, S., Manerikar, R. & Sinha, R. Surgical management of transverse maxillary deficiency in adults. J. Maxillofac. Oral Surg. 9(3), 241–246 (2010).
9. Yu, X. et al. Cephalometric analysis in obese and nonobese patients with obstructive sleep apnea syndrome. Chest 124(1), 212–218 (2003).
10. Akpinar, M. E. et al. The comparison of cephalmatic characteristics in nonobese obstructive sleep apnea subjects and primary snorers cephalmatic measures in nonobese OSA and primary snorers. Eur. Arch. Orothognal Surg. 268(7), 1053–1059 (2011).
11. Kim, Y. K. et al. Influencing factors on the effect of mandibular advancement device in obstructive sleep apnea patients: Analysis on cephalmatic and polysomnographic parameters. Sleep Breath. 18(2), 305–311 (2014).
12. Ng, A. T. et al. Cephalometry and prediction of oral appliance treatment outcome. Sleep Breath. 16(1), 47–58 (2012).
13. Ronchi, P. et al. Maxillomandibular advancement in obstructive sleep apnea syndrome patients: A retrospective study on the sagittal cephalmatic variables. J. Oral Maxillofac. Res. 4(2), e5 (2013).
14. Devore, J. In Probabilidad e estadística: para engenharia e ciências (Pioneira Thompson Learning, 2006).
15. Hersh, H. G., Stewart, B. L. & Warren, D. W. Changes in nasal airway resistance associated with rapid maxillary expansion. Sleep Med. 20(2), 501–508 (2016).
16. Vinha, P. P. et al. Effects of surgically assisted rapid maxillary expansion on obstructive sleep apnea and daytime sleepiness. Sleep Breath. 20(2), 501–508 (2016).
17. Bonetti, G. A. et al. Case report on the efficacy of transverse expansion in severe obstructive sleep apnea syndrome. J. Maxillofac. Oral Surg. 11, 23 (2015).
18. Yu, X. et al. The usefulness of cephalmatic measurement as a diagnostic tool for obstructive sleep apnea syndrome: A retrospective study. Oral Surg. Oral Med. Oral Pathol. Oral Radiol. 119(1), 20–31 (2015).
19. Parhiz, A. et al. Lateral cephalmatic changes after SARPE. Int. J. Oral Maxillofac. Surg. 40(7), 662–671 (2011).
20. Bretos, J. I. et al. Sagittal and vertical maxillary effects after surgically assisted rapid maxillary expansion (SARME) using Haas and Hyrax expanders. J. Craniofac. Surg. 18(6), 1322–1326 (2007).
21. Cahal, M. B. Laterale pharyngoplasty: A new treatment for obstructive sleep apnea hypopnea syndrome. Laryngoscope. 113(11), 1961–1968 (2003).
22. Hellak, A. F. et al. Influence of maxillary advancement surgery on skeletal and soft-tissue changes in the nose—A retrospective cone beam computed tomography study. Head Face Med. 11, 23 (2015).
23. Cistulli, P. A. Craniofacial abnormalities in obstructive sleep apnoea: implications for treatment. Respiratory 1(3), 167–174 (1996).
24. Guilleminault, C., Riley, R. & Powell, N. Obstructive sleep apnea and abnormal cephalmatic measurements. Implications for treatment. Chest 86(5), 793–794 (1984).
25. Posnick, J. C. et al. “Silent” sleep apnea in dentofacial deformities and prevalence of daytime sleepiness after orthognathic and intranasal surgery. J. Oral Maxillofac. Surg. 76(4), 833–843 (2018).
26. Dentomaxillofac. Radiol. 45(1), 20150085 (2016).
27. Silva, V. G. et al. Correlation between cephalmatic data and severity of sleep apnea. Braz. J. Oral Maxillofac. Surg. 80(3), 191–195 (2014).
28. Ryu, H. H. et al. The usefulness of cephalmatic measurement as a diagnostic tool for obstructive sleep apnea syndrome: A retrospective study. Oral Surg. Oral Med. Oral Pathol. Oral Radiol. 119(1), 20–31 (2015).
29. Parhiz, A. et al. Lateral cephalmatic changes after SARPE. Int. J. Oral Maxillofac. Surg. 40(7), 662–671 (2011).
30. Bretos, J. I. et al. Sagittal and vertical maxillary effects after surgically assisted rapid maxillary expansion (SARME) using Haas and Hyrax expanders. J. Craniofac. Surg. 18(6), 1322–1326 (2007).
31. Cahal, M. B. Laterale pharyngoplasty: A new treatment for obstructive sleep apnea hypopnea syndrome. Laryngoscope. 113(11), 1961–1968 (2003).
32. Hellak, A. F. et al. Influence of maxillary advancement surgery on skeletal and soft-tissue changes in the nose—A retrospective cone beam computed tomography study. Head Face Med. 11, 23 (2015).
33. Hernández-Alfaro, F., Guitarro-Martínez, R. & Mareque-Bueno, J. Effect of mono- and bimaxillary advancement on pharyngeal airway volume: Cone-beam computed tomography evaluation. J. Oral Maxillofac. Surg. 69(1), 835–840 (2011).
34. Rosário, H. D. et al. Surgical maxillary advancement increases upper airway volume in skeletal class III patients: A cone beam computed tomography-based study. J. Clin. Sleep Med. 12(11), 1527–1533 (2016).
36. Miloro, M. et al. Is there consistency in cephalometric landmark identification amongst oral and maxillofacial surgeons? Int. J. Oral Maxillofac. Surg. 43(4), 445–453 (2014).
37. Koo, T. K. & Li, M. Y. A guideline of selecting and reporting intraclass correlation coefficients for reliability research. J. Chiropr. Med. 15(2), 155–163 (2016).
38. Hoekema, A. et al. Craniofacial morphology and obstructive sleep apnoea: A cephalometric analysis. J. Oral Rehabil. 30(7), 690–696 (2003).
39. Kim, S. J. et al. Cephalometric predictors of therapeutic response to multilevel surgery in patients with obstructive sleep apnea. J. Oral Maxillofac. Surg. 70(6), 1404–1412 (2012).

Author contributions
R.F.B. and M.D.P. conceived of the presented idea and developed the theory and performed the computations. M.D.P. and G.P.P. verified the analytical methods. S.E.D. and L.M.F. supervised the findings of this work. All authors discussed the results and contributed to the final manuscript.

Competing interests
The authors declare no competing interests.

Additional information
Supplementary Information The online version contains supplementary material available at https://doi.org/10.1038/s41598-021-85935-7.
Correspondence and requests for materials should be addressed to M.D.P.
Reprints and permissions information is available at www.nature.com/reprints.
Publisher’s note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2021