Assessment of Periodontal Biotype in a Young Chinese Population using Different Measurement Methods

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Periodontal biotype is used to describe the morphological characteristics of periodontal tissues and is closely related to periodontal health and prognosis of many dental treatments. This study was undertaken to explore the periodontal biotype distribution in a young Chinese population and to evaluate the accuracy of different methods for gingival thickness (GT) measurement. A total of 372 teeth from 31 periodontally healthy subjects were included. GT was measured simultaneously by probe transparency, transgingival probing and cone-beam computed tomography (CBCT). Some other anatomic parameters, including crown width/crown length ratio, attached gingival width, labial bone thickness and papilla volume were recorded for periodontal biotype classification. As found by probe transparency, the gingivae of 222 teeth (59.68%) were thick, while those of 150 teeth (40.32%) were thin. The mean GT of included subjects was 1.03 ± 0.31 mm as measured by transgingival probing and 1.03 ± 0.24 mm as measured by CBCT. Four groups were identified by cluster analysis. Thick-flap biotype, average-scalloped biotype, average-flap biotype and thin-scalloped biotype comprised 137 teeth (36.83%), 96 teeth (25.81%), 39 teeth (10.48%) and 100 teeth (26.88%), respectively. These results demonstrate that the most common periodontal biotype in this young Chinese population was the thick-flap type with low aesthetic risk.

Clinical appearance of gingiva differs from subject to subject and even among different teeth. It is closely related to periodontal health and the prognosis of many dental treatments. To describe the variation in gingival contour, “gingival biotype” was first proposed by Ochsenbein and Ross1. Then, Seibert and Lindhe introduced the term “periodontal biotype”, which classified the gingival contour into two categories, the “thick” and “thin” biotype, based on simple visual appearance of gingivae2. In 1997, Muller included some new parameters, such as tooth shape and gingival width, in his analysis of periodontal biotype3. De Rouck developed a new method for the classification of gingival biotype based on the following four clinical parameters: gingival thickness (GT), crown width/crown length ratio (CW/CL), gingival width and papilla height (PH). Then, three biotypes were identified: the thin-scalloped, thick-scalloped and thick-flat biotype.4 Unfortunately, until now, there has been no unified criterion for periodontal biotype classification, although GT is undoubtedly the most important indicator of all for evaluation.

Many invasive and non-invasive methods have been employed to assess periodontal biotype. However, the most accurate method for periodontal biotype assessment and GT measurement has not yet been confirmed. Simple visual inspection was the first method developed to evaluate periodontal biotype, but it was not considered to be a reliable method by many researchers5,6. Transgingival probing is a traditional invasive method with limited application in clinic. Instead, another evaluation that is based on the transparency of the periodontal probe through the gingival margin is widely used and is taken as a simple method with excellent repeatability. Ultrasonic measurement and cone beam computerized tomography (CBCT) are also non-invasive methods, but special devices are needed for these assessments. Moreover, it was reported that the accuracy of GT measurement by ultrasonic instruments was worse than that by direct puncture7. Comparisons of different measurement methods based on the same sample are limited.
Some anatomic parameters, such as CW/CL, attached gingival width (AGW), PH, bone thickness (BT) and tooth site, were reported to be related to GT. Different results of GT assessment and periodontal biotype exploration in different studies were also due to the included samples with different age, sex and racial characteristics. Until recently, most research on periodontal biotype has been about Caucasians. In contrast, explorations in Chinese populations are limited.

The aim of the present study was to explore the periodontal biotype distribution in a young Chinese population and to evaluate the accuracy of different methods for GT measurement, including probe transparency, transgingival probing and CBCT.

Method and Materials

Subjects. This study was performed in compliance with the revised Helsinki Declaration and was approved by the Ethical Committee of Nanjing Medical University (Permit Number: 20160206). Written informed consent was obtained from all recruits.

A total of 31 periodontally healthy students in the College of Stomatology, Nanjing Medical University (aged between 18 and 27 years, mean age 22.2 years, 15 males and 16 females) were enrolled in this study from October 2016 to February 2017. Inclusion criteria were: (1) 18–30 years old; (2) no site with gingival index (GI) ≥1, between 18 and 27 years, mean age 22.2 years, 15 males and 16 females) were enrolled in this study from October 2016 to February 2017. Inclusion criteria were: (1) 18–30 years old; (2) no site with gingival index (GI) ≥1, probing depth (PD) >3 mm or clinical attachment loss (CAL) ≥1 mm and no radiographic evidence of alveolar bone loss; and (3) no malocclusion, crowding, missing or supernumerary anterior teeth. Exclusion criteria were: (1) crown restorations or fillings in anterior teeth; (2) previous orthodontic treatment; (3) previous periodontal surgery, including gingivectomy, flap surgery, guided tissue regeneration or periodontal plastic surgery; (4) systemic disease; (5) pregnancy and lactation; (6) use of any drugs that might lead to gingival enlargement during the past 6 months; (7) smoking; and (8) bruxism. All volunteers received oral hygiene instructions and a full-mouth supragingival scaling 1 week before clinical examination to control any possible gingival inflammation.

Clinical measurement. A total of 372 anterior teeth from 31 subjects, including upper and inferior anterior teeth, were involved in this study. Clinical periodontal parameters, including plaque index (PLI), GI, PD and CAL were recorded at six sites (mesial labial, midlabial, distal labial, mesial palatal, midpalatal and distal palatal) per tooth. All measurements were performed by a calibrated examiner.

For transgingival probing, local anesthesia (4% articaine) was applied over the gingiva of anterior teeth. To avoid mucous swelling due to local anesthesia, clinical measurements were taken 20 min after anesthesia. GT by transgingival probing (GTp) was determined at the level of the cementoenamel junction (CEJ) by piercing midfacial gingiva with a #15 endodontic K-file (MANI, Tochigi, Japan), which had a rubber stop. Gingivae thicker than 0.8 mm were defined as thick, while others were categorized as thin.

The following anatomic parameters were measured by a Williams probe (Hu-Friedy, IL, USA) to the nearest 0.5 mm:

1. CW/CL. Crown length was recorded as the distance between the incisal edge of the tooth and the free gingival margin, or if discernible, the CEJ. Crown width was the distance between the approximal tooth surfaces, which was measured at the border between the middle and the cervical third of the crown.

2. AGW was evaluated at the midfacial point by subtracting PD from keratinized gingival width, which was the distance from the free gingival margin to the mucogingival junction.

3. PH was defined as the distance from the tip of the papilla to a line connecting the midfacial soft tissue margin of two adjacent teeth.

Moreover, the visibility of a probe inserted through the gingival margin at the midfacial site of the tooth was also determined to assess periodontal biotype. If the outline of the probe could be seen through the gingiva, it was defined as thin. Otherwise, it was regarded as thick.

Radiographic measurement. To avoid the interference of surrounding soft tissues (lip and tongue), acrylic plates of maxillary and mandibular arcs extending beyond the mucogingival junction were made, and zinc oxide–eugenol cement was loaded into it. Then, the plates were allowed to set intraorally during the process of CBCT examination.

CBCT scanning was performed using NewTom 5G (version FP, Verona, Italy). Exposure was performed at 110 kV in 4–5 mA for 3.6 s. Reconstructed images were generated, and digital measurements were taken using Mimics software (Materialise NV, Leuven, Belgium) with an accuracy of 0.01 mm. The following parameters were recorded: (1) GT in CBCT scanning (GT CT) was measured at the level of CEJ; (2) labial bone thickness (BT) was assessed at the midpoint of the root. The above-mentioned thickness measurements were conducted perpendicular to the midfacial surface of the tooth, as shown in Fig. 1; and (3) after 3D imaging reconstruction, gingival papilla volume (PV), which was defined by 14 points, was calculated automatically by Mimics software (Fig. 2).

Intra-examiner repeatability. All measurements were performed by one examiner (YMS). 10 volunteers randomly extracted from the total samples were reevaluated after 1 month by the same clinician.

Statistical analysis. Statistical analysis of clinical and radiologic parameters, including GT, CW/CL, AGW, PH and BT, was performed using Student’s t-test. Spearman’s correlation coefficient was used to analyze the consistency of GT obtained by transgingival probing and CBCT, as well as the correlations of CW/CL, AGW, PH, BT, PV and GT. Agreement of probe transparency and GT measurements was analyzed by kappa test. Hierarchical cluster analysis (HCA) based on Euclidian distances of four parameters, including GT CT, CW/CL, AGW and PV, was conducted to develop periodontal tissues typology more objectively. Before the parameters were put into...
HCA, they were standardized in order to ensure the same weight in the analysis. Differences in morphometric variables of the resultant clusters were analyzed by ANOVA, and the LSD test was used to compare differences between groups.

**Data availability.** The data generated and analyzed in this present study are available from the corresponding author upon reasonable request.

**Results**

**Intra-examiner accuracy control.** The accuracy and repeatability of the measurements were repeatedly evaluated in 10 volunteers. Pearson correlation coefficients were 0.92 ($p < 0.001$), 0.84 ($p < 0.001$), 0.94 ($p < 0.001$), 0.85 ($p < 0.001$), 0.76 ($p < 0.001$) and 0.72 ($p < 0.001$) for CW/CL, AGW, PH, GT, GT, and BT, respectively. Probe transparency also proved to be highly reproducible, with 86% agreement between duplicate measurements ($k = 0.80$, $p < 0.05$).
GT P-BT, GT CT-BT and BT-PV (p < 0.05 compared with homonym tooth in maxillary.

Table 2. Effects of tooth position on clinical and radiographic parameters (mean ± SD). CW/CL crown width/crown length ratio, AGW attached gingival width, PH papilla height, GT p gingival thickness measured by transgingival probing, GT CT gingival thickness measured by CBCT, BT bone thickness, and PV papilla volume. *p < 0.05 compared with maxillary teeth.

Table 3. Distribution of thick/thin gingiva assessed by different methods. GT p or GT CT ≥ 0.8 mm was defined as thick gingiva, while the remainder were categorized as thin.

Table 4. Correlation analysis was carried out to explore the factors closely related to periodontal biotype. A strong correlation between CW/CL and PV was revealed (p < 0.01, r = 0.554), while moderate correlations between CW/CL-AGW, CW/CL-GT p, AGW-GT CT, CT CT-GT p, PV-GT p, GT CT-PV and PV-PH were also found (p < 0.01, 0.3 < r < 0.5). In addition, weak correlations existed between CW/CL-PH, CW/CL-GT CT, CW/CL-BT, AGW-BT, AGW-PV, PH-GT p, PH-BT, GT p-BT, GT CT-BT and BT-PV (p < 0.01, 0 < r ≤ 0.3). There was no significant correlation between AGW-PH and PH-GT CT (p > 0.05) (Table 4).

Cluster analysis. By HCA, four clusters of periodontal biotype were identified based on GT p CW/CL, AGW and BT. Cluster A (thick-flap type), cluster B1 (average-scalloped type), cluster B2 (average-flap type) and cluster C (thin-scalloped type) included 137 teeth (36.83%), 96 teeth (25.81%), 39 teeth (10.48%) and 100 teeth (26.88%), respectively. Representative photographs of these four periodontal biotypes are presented in Fig. 3.
Among the four groups, the GT of cluster A was the thickest (p < 0.001), while that of cluster C was the thinnest (p < 0.001). No significant differences in GT were found between cluster B2 and B1 (p > 0.05). The CW/CL of cluster B2 was significantly larger than those of the other groups (p < 0.001), which represented a quarter tooth shape. Similarly, the PVs of clusters A and B2 were much greater than those of clusters B1 and C (p < 0.001). Furthermore, the AGWs of clusters B1 and C were significantly smaller than those of the other two groups (p < 0.001) (Table 4).

**Discussion**

Periodontal biotype is closely related to the prognosis of many dental treatments, especially in the anterior aesthetic area. After implantation, periodontal therapy or orthodontic treatment, severe gingival recession may develop more easily in patients with a thin biotype, while deep periodontal pockets are more likely to form in persons with a thick biotype. Therefore, it is important to identify periodontal biotype accurately in order to predict prognosis and avoid unexpected complications.

In this present study, the GT in a young Chinese population was evaluated by different methods (probe transparency, transgingival probing and CBCT) and the periodontal biotype distribution was explored by cluster analysis. Moreover, PV was first measured by CBCT and its relationship with periodontal biotype was analyzed at the same time.

The influences of some factors, such as the position and shape of the tooth, race, gender and age, on periodontal biotype have been confirmed by some researchers. Until now, most studies on periodontal biotype have been in Caucasians. In Stein’s study, 60 central maxillary incisors from 60 periodontally healthy Caucasian were involved, and the mean GT was 1.25 mm. In Nikiforidou’s research, 42 periodontally healthy subjects were recruited, and the mean GT of 186 maxillary anterior teeth was found to be 1.20 mm. Le’s study included 84 maxillary teeth from

| CWCL | AGW | PH | GT<br> | GT<br> | BT | PV
|------|-----|----|-------|-------|----|----|
| 0.328* | 0.161* | 0.161* | 0.286* | 0.183* | 0.554* |
| 0.384* | 0.308* | 0.308* | 0.275* | 0.263* |
| 0.182* | 0.182* | 0.182* | 0.182* | 0.182* |
| 0.196* | 0.196* | 0.196* | 0.196* | 0.196* |
| 0.264* | 0.264* | 0.264* | 0.264* | 0.264* |

*<p>0.001.

Table 4. Correlation analysis of clinical and radiographic parameters (Spearman’s rho). CW/CL crown width/crown length ratio, AGW attached gingival width, PH papilla height, GT gingival thickness measured by transgingival probing, GT gingival thickness measured by CBCT, BT bone thickness, and PV papilla volume.

Figure 3. Representative photographs of the four periodontal biotypes. 11 in (a) 11 in (b) 11 in (c) and 22 in (d) were thick-flap, average-scalloped, average-flap and thin-scalloped biotype respectively.
Table 5. Clinical parameters per cluster (mean ± SD). CW/CL crown width/crown length ratio, AGW attached gingival width, PH papilla height, GT, gingival thickness measured by transgingival probing, GT, gingival thickness measured by CBCT, BT bone thickness, and PV papilla volume. \*p < 0.001 in comparison with cluster B1, \*p < 0.001 in comparison with cluster B2, \*p < 0.001 in comparison with cluster C.

| Parameter       | Cluster A | Cluster B1 | Cluster B2 | Cluster C | Total  |
|-----------------|-----------|------------|------------|-----------|--------|
| n               | 137       | 96         | 39         | 100       | 372    |
| CW/CL (mm)      | 0.84 ± 0.10± 0.13 | 0.72 ± 0.07± 0.13 | 0.92 ± 0.10± 0.13 | 0.65 ± 0.09± 0.13 | 0.77 ± 0.13± 0.13 |
| AGW (mm)        | 6.00 ± 1.17± 0.13 | 4.31 ± 0.84± 0.13 | 7.21 ± 1.20± 0.13 | 4.82 ± 1.10± 0.13 | 5.37 ± 1.42± 0.13 |
| PH (mm)         | 4.56 ± 0.89± 0.13 | 4.39 ± 0.75± 0.13 | 4.67 ± 1.09± 0.13 | 4.00 ± 0.72± 0.13 | 4.34 ± 0.84± 0.13 |
| GT, (mm)        | 1.27 ± 0.20± 0.13 | 1.02 ± 0.26± 0.13 | 1.01 ± 0.26± 0.13 | 0.71 ± 0.17± 0.13 | 1.03 ± 0.31± 0.13 |
| CT, GT (mm)     | 1.13 ± 0.26± 0.13 | 0.99 ± 0.22± 0.13 | 1.07 ± 0.26± 0.13 | 0.93 ± 0.19± 0.13 | 1.03 ± 0.24± 0.13 |
| BT (mm)         | 0.72 ± 0.25± 0.13 | 0.65 ± 0.20± 0.13 | 0.65 ± 0.19± 0.13 | 0.63 ± 0.15± 0.13 | 0.67 ± 0.21± 0.13 |
| PV (mm³)        | 89.98 ± 26.24± 0.13 | 69.20 ± 20.24± 0.13 | 90.24 ± 23.65± 0.13 | 55.36 ± 18.18± 0.13 | 75.21 ± 25.27± 0.13 |

14 periodontally healthy Chinese individuals, and revealed that subjects with thin biotypes were only 9% of that population. In addition, in most studies, only maxillary anterior teeth were involved. Differences in GT between maxillary and mandibular anterior teeth were reported by La Rocca and Müller. Although the aesthetic appearance of mandibular anterior teeth does exist, studies concerning mandibular teeth are still limited. Therefore, both maxillary and mandible teeth were included in this study and the sample size was expanded to 372 teeth.

In the past ten years, periodontal biotype distribution was explored in numerous studies. Among them, the periodontal biotypes of subjects, but not their teeth, were confirmed by most researchers. However, the effects of tooth sites on GT were revealed by Müller and Vandana. Therefore, differences in periodontal biotype/GT from tooth to tooth in the same subject do exist. It is imprecise to classify a mandibular tooth with thin gingiva as a thick biotype based on the data from a maxillary tooth with thick gingiva. De Rouck noticed that the results of transgingival probing in two maxillary incisors might be different. In this present study, differences in GT, AGW, PH and BT between maxillary and mandibular teeth were also observed, which further confirmed that the periodontal biotype of upper anterior teeth might not represent the biotype of all anterior teeth, let alone the whole dentition. To avoid unsatisfactory prognosis, it is still very important to explore the biotype of inferior teeth accurately before some dental treatments, such as orthodontic treatment and implantation.

Many invasive and non-invasive methods have been employed to assess periodontal biotype, such as transgingival probing, probe transparency and CBCT. Transgingival probing is an invasive direct measurement performed by inserting an injection needle, periodontal probe or endodontic file through gingiva under local anesthesia. Infusion of the anesthetic agent, angulations of probing and distortion of tissues might affect the precision of this measurement. In addition, excessive puncture strength might lead to the penetration of periosteum and even lamina dura. Probe transparency is a non-invasive method based on the transparency of a periodontal probe through the gingival margin. A clinical trial involving 100 periodontally healthy subjects indicated that probe transparency was highly reproducible, with 85% agreement between duplicate recordings. However, it is also a subjective method, and its accuracy depends on the examiner’s experience. Stein found that the prognostic value of probe transparency for GT and BT was limited. For convenience of measurement, crown length was recorded as the distance between incisal edge of the teeth and free gingival margin in this study. However, it is important to point out that, to some degree, clinical crown length (from incisal edge to free gingival margin) is different from anatomical crown length (from incisal edge to CEJ), which might affect the results. Moreover, Nikiforidou indicated that GT at the CEJ was related to labial BT at a point 3 mm apical to the CEJ. However, La Rocca pointed out that BT was related to AGW, but not to GT. Our study indicated that the correlations between BT at the midpoint of the root and some other parameters (GT, CW/CL, AGW and PV) were weak. In addition, there were no differences in BT among the four groups created by cluster analysis, which suggested that BT might be less associated with periodontal biotypes.
Gingival papilla in the anterior aesthetic area is not only an important biological barrier to protect the teeth and surrounding tissues but also an artificial factor in aesthetics. During treatment in the aesthetic area, attention needs to be paid not only to the health of gingivae but also to their beauty and harmony. The appearance of interdental papilla, especially its height, is important for aesthetics in implantation. De Rouck reported that the mean PH in Caucasians was 3.96 mm. Cao found that the mean height and width of papilla were 3.67 mm and 4.35 mm in Chinese, which was different from our results. Compared with the height and width of papilla, PV is a parameter in three-dimensional space and may represent the quantity of soft tissue more accurately. It is almost impossible to measure PV without CBCT scanning and three-dimensional reconstruction. Therefore, until now, there have been no studies exploring PV. Our study first found that PV was highly correlated with CW/CL and GT, which implied the underlying relationship between PV and the appearance of tooth and periodontal soft tissues. In this present study, smaller PV was found to be more common in teeth characterized by average-scalloped and thin-scalloped biotypes. Black triangles caused by the destruction of papilla are a major cosmetic deficiency for patients who expect aesthetically pleasing restorative or orthodontic treatments. Some researchers found that black triangles developed more easily in patients with scalloped teeth. It is unclear whether PV is a risk factor for black triangles, and the relationship between PV and black triangles needs to be further explored.

Exploration of periodontal biotype distribution by cluster analysis was undertaken by some researchers. K-mean clustering algorithm, which classified samples according to categories specified by investigators, was employed by Müller, De Rouck and Gobbato, who artificially classified the included subjects into three periodontal biotypes. Nikiforidou found four periodontal types by HCA. During this statistical analysis, numbers of categories were decided by the characteristics of the data themselves, not artificially imposed by researchers. When a small sample is analyzed, the results by HCA may be scattered and difficult to generalize, while k-mean clustering may be efficient and easy to derive results. However, when larger samples are analyzed, HCA can reflect the characteristics of the data themselves as much as possible. Therefore, owing to the quantity of teeth included in this study, HCA was employed.

In papers that employed cluster analysis, different variables were analyzed in different studies, and their weights were also different. In general, GT and the shape of teeth have been the most important and frequently employed parameters to distinguish periodontal biotypes. Gobbato even paid so much attention to tooth morphology that he did not consider GT and BT as parameters for cluster analysis. Some other variables, including AGW, keratinized gingiva width and PH, were also taken into consideration by some studies. Due to the strong correlations of PV-CW/CL and PV-GT, we included PV in cluster analysis for the first time.

As a result of the variety of included samples, measurement means and statistical methods, different categories and proportional distribution were obtained in different papers. In Stein's study, 46.7% of Caucasians were classified as having the thin biotype, and the others were classified as having the thick biotype. De Rouck included 100 Caucasians and thin-scalloped, thick-flat and thick-scalloped biotypes were found to be 37%, 29% and 34% of the sample, respectively. Let's study identified that the thin, compromised and thick biotypes in Chinese individuals were 9%, 50% and 41%, respectively. Four periodontal biotypes were reported by Nikiforidou: thin, average, mixed and thick type, which comprised 31.9%, 31.3%, 16.6% and 20.2% of the sample, respectively. In our results, the most and the least common types were thick-flat (36.8%) and average-flat type (10.5%), while the rest were average-scalloped and thin-scalloped type (each approximately 1/4 of the total samples). Our present study also revealed an effectively equal distribution of slender and square teeth in a young Chinese population (slender 52.69% and square 47.31%). Moreover, there were more slender teeth in subjects with average/thin gingival and more square teeth in persons with thick/average gingiva.

In summary, a possible diversity of periodontal biotypes beyond thick and thin might exist in the young Chinese population. The most common periodontal biotype was the thick-flat biotype, which displayed a square tooth, average AGW, thick BT and large gingival papilla. Meanwhile, the second-largest biotype, the thin-scalloped biotype, which displayed a slender tooth, narrow AGW, thin BT and small gingival papilla, was the one with the highest aesthetic risk. In clinical practice, more attention should be paid to these teeth, especially in the aesthetic zone.

References
1. Ochsenbein, C. & Ross, S. A reevaluation of osseous surgery. *Dent Clin North Am.* 13, 87–102 (1969).
2. Seibert, J. & Lindhe, J. Esthetics and periodontal therapy. In *Textbook of Clinical Periodontology* (ed. Lindhe, J.) 477–514 (Munkgaard, 1989).
3. Müller, H. P. & Eger, T. Gingival phenotypes in young male adults. *J Clin Periodontol.* 24, 65–71 (1997).
4. De Rouck, T., Eghbali, R., Collys, K., De Bruyn, H. & Cosyn, J. The gingival biotype revisited: transparency of the periodontal probe through the gingival margin as a method to discriminate thin from thick gingiva. *J Clin Periodontol.* 36, 428–433 (2009).
5. Cuny-Houchmand, M. et al. Gingival biotype assessment: visual inspection relevance and maxillary versus mandibular comparison. *Open Dent J.* 7, 1–6 (2013).
6. Eghbali, A., De Rouck, T., De Bruyn, H. & Cosyn, J. The gingival biotype assessed by experienced and inexperienced clinicians. *J Clin Periodontol.* 36, 958–963 (2009).
7. Savitha, B. & Vandana, K. L. Comparative assessment of gingival thickness using transgingival probing and ultrasonographic method. *Indian J Dent Res.* 16, 135–139 (2005).
8. Fischer, K. R. et al. On the relationship between gingival biotypes and supraeruptional gingival height, crown form and papilla height. *Clin Oral Implants Res.* 25, 894–898 (2014).
9. Fischer, K. R., Richter, T., Kebschull, M., Petersen, N. & Fickl, S. On the relationship between gingival biotypes and gingival thickness in young Caucasians. *Clin Oral Implants Res.* 26, 865–869 (2015).
10. Kim, J. H., Cho, Y. J., Lee, J. Y., Kim, S. J. & Choi, J. J. An analysis on the factors responsible for relative position of interproximal papilla in healthy subjects. *J Periodontal Implant Sci.* 43, 160–167 (2013).
11. Stein, J. M. et al. The gingival biotype: measurement of soft and hard tissue dimensions - a radiographic morphometric study. *J Clin Periodontol.* 40, 1132–1139 (2013).
12. Nikiforidou, M. et al. Classification of periodontal biotypes with the use of CBCT. A cross-sectional study. *Clin Oral Investig.* 20, 2061–2071 (2016).
13. Frost, N. A., Mealey, B. L., Jones, A. A. & Huynh-Ba, G. Periodontal biotype: Gingival thickness as it relates to probe visibility and buccal plate thickness. J Periodontol. 86, 1141–1149 (2015).
14. Lee, A., Fu, J. H. & Wang, H. L. Soft tissue biotype affects implant success. Implant Dent. 20, 38–47 (2011).
15. Le, D., Zhang, H., Hu, W. J. & Liu, D. G. Preliminary study on gingival biotype by periodontal probing by periodontal probing. Zhonghua Kou Qiang Yi Xue Za Zhi. 47, 81–84 (2012).
16. La Rocca, A. P. et al. Anterior maxillary and mandibular biotype: relationship between gingival thickness and width with respect to underlying bone thickness. Implant Dent. 21, 507–515 (2012).
17. Müller, H. P., Schaller, N., Eger, T. & Heinecke, A. Thickness of masticatory mucosa. J Clin Periodontol. 27, 431–436 (2000).
18. Lazos, J. P., Senn, L. F. & Brunotto, M. N. Characterization of maxillary central incisor: novel crown-root relationships. Clin Oral Investig. 18, 1561–1567 (2014).
19. Monteverchi, M., Checchi, V., Piana, L. & Cecchi, L. Variables Affecting the Gingival Embrasure Space in Aesthetically Important Regions: Differences between Central and Lateral Papillae. Open Dent J 5, 126–135 (2011).
20. Stellini, E., Comuzzi, L., Mazzocco, F., Parente, N. & Gobbato, L. Relationships between different tooth shapes and patient's periodontal phenotype. J Periodontal Res. 48, 657–662 (2013).
21. Vandana, K. L. & Savitha, B. Thickness of gingiva in association with age, gender and dental arch location. J Clin Periodontol. 32, 828–830 (2005).
22. Joss-Vassalli, L., Grebenstein, C., Topouzelis, N., Sculean, A. & Katsaros, C. Orthodontic therapy and gingival recession: a systematic review. Orthod Craniofac Res. 13, 127–141 (2010).
23. Malhotra, R., Grover, V., Bhardwaj, A. & Mohindra, K. Analysis of the gingival biotype based on the measurement of the dentopapillary complex. J Indian Soc Periodontol. 18, 43–47 (2014).
24. Gobbato, L., Tsukiyama, T., Levi, P. A., Griffin, T. J. & Weisgold, A. S. Analysis of the shapes of maxillary central incisors in a Caucasian population. Int J Periodontics Restorative Dent. 32, 69–78 (2012).
25. Müller, H. P., Heinecke, A., Schaller, N. & Eger, T. Masticatory mucosa in subjects with different periodontal phenotypes. J Clin Periodontol. 27, 621–626 (2000).
26. Silva, J. N. N. et al. Influence of lip retraction on the cone beam computed tomography assessment of bone and gingival tissues of the anterior maxilla. Oral Surg Oral Med Oral Pathol Oral Radiol. 123, 714–720 (2017).
27. Kim, Y. J. et al. New method of assessing the relationship between buccal bone thickness and gingival thickness. J Periodontal Implant Sci. 46, 372–381 (2016).
28. Mallikarjun, S. et al. Comparative evaluation of soft and hard tissue dimensions in the anterior maxilla using radiovisography and cone beam computed tomography: A pilot study. J Indian Soc Periodontol. 20, 174–177 (2016).
29. Zetu, L. & Wang, H. L. Management of inter-dental/inter-implant papilla. J Clin Periodontol. 32, 831–839 (2005).
30. Thoma, D. S., Buranawat, B., Hämmerle, C. H., Held, U. & Jung, R. E. Efficacy of soft tissue augmentation around dental implants and in partially edentulous areas: a systematic review. J Clin Periodontol. 41, 77–91 (2014).
31. Cao, J., Hu, W. J., Zhang, H., Liu, D. G. & Le, D. Preliminary study on measuring interdental papilla height and thickness of the underlying bone thickness. Implant Dent. 21, 507–515 (2012).
32. Chow, Y. C., Eber, R. M., Tsao, Y. P., Shotwell, J. L. & Wang, H. L. Factors associated with the appearance of gingival papillae. J Clin Periodontol. 37, 719–727 (2010).

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Author Contributions

Y.S. conceived and designed the experiments. Y.M.S., L.L.Y. and J.Y.G. performed the experiments. D.M.W. analyzed the data. Y.M.S. and L.L.Y. contributed to the drafting. W.L. prepared figures and tables. Y.S. revised the paper and supervised the whole study. All authors approved the final manuscript as submitted and agree to be accountable for the work.

Additional Information

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