Accuracy of periodontal probe visibility in the assessment of gingival thickness

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ABSTRACT

Purpose: The present study was undertaken to examine whether periodontal probe visibility (PV) accurately reflects gingival thickness (GT) and to identify factors affecting PV using cluster and multivariate analyses.

Methods: The clinical characteristics of the maxillary central incisors (n=90 subjects) were examined. Clinical photographs, sex, PV, probing depth, gingival width, papilla height, GT as measured with an ultrasonic device, and the ratio of crown width to crown length were recorded. Multivariate analysis, using multinomial baseline-category logistic regression, was used to identify factors predictive of PV. Cluster analysis was used to identify gingival biotypes.

Results: In the multivariate analysis, sex was the only significant predictor of PV (odds ratio, 6.48). Two clusters of subjects were created based on morphometric parameters. The mean GT among cluster A subjects was significantly lower than that among cluster B subjects (P=0.015). No significant difference was found between cluster A and B subjects in terms of PV score (P=0.583).

Conclusions: Periodontal PV was not associated with GT as measured directly using an ultrasonic device. Sex was a highly significant predictor of periodontal PV. GT was found to be correlated with morphological characteristics of the periodontium.

Keywords: Cluster analysis; Gingiva; Periodontal index; Periodontium; Phenotype; Ultrasonics

INTRODUCTION

The clinical appearance of healthy periodontal tissue is variable. Ochsenbein and Ross [1] first classified the gingival anatomy as being of a flat or pronounced scalloped biotype, suggesting that a flat gingiva was associated with a square tooth form and that a pronounced scalloped gingiva was associated with a tapered tooth form. Gingival biotype is related to gingival thickness (GT); thin gingivae is ≤1.5 mm thick, whereas thick gingivae are ≥2 mm thick [2]. Later, Seibert and Lindhe [3] proposed the existence of 2 distinct gingival biotypes, which are based on morphological characteristics of the periodontium: the scalloped-thin biotype and the flat-thick biotype.
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Conflict of Interest
No potential conflict of interest relevant to this article was reported.

Esthetics has become a central concern for those undergoing dental procedures; therefore, understanding how the periodontium responds to therapy is critical to achieving acceptable outcomes for patients. The thin biotype has been reported to be associated with a relatively high prevalence of gingival recession in studies of periodontal surgery for conventional or implant prostheses. Consequently, it would be clinically helpful to identify a patient’s gingival biotype and GT before that patient undergoes dental treatment in highly visible areas of the mouth [4].

In the past, GT was measured directly by inserting a sterile syringe needle marked with an endodontic gauge into the gum or by using calipers after tooth extraction [5]. Understandably, due to the invasiveness of these methods, direct measurements of GT were not commonly performed. Kan et al. [6] suggested an alternative, non-invasive method for estimating GT, namely determination of the gingival biotype. Using this method, the gingival biotype is categorized as either thin or thick according to the visibility of the underlying periodontal probe through the gingival sulcus. This non-invasive method was widely accepted as the easiest and most straightforward means of determining the gingival biotype, which is used as a proxy for GT. In addition, another direct measurement method was introduced, involving an ultrasonic device and the pulse echo principle [7]. In this method, ultrasonic pulses are transmitted through the sound-permeable mucosa and reflected at the surface of hard tissue (i.e., the alveolar bone or tooth). The time delay between pulse transmission and the arrival of the echo enables the thickness of the masticatory mucosa to be estimated at a resolution of 0.1 mm, and the results are displayed [7-10]. The reliability and validity of measurements of mucosal thickness in different parts of the mouth have been found to be acceptable [10-12]. This ultrasonic device is not currently commercially available and is therefore rarely used; nevertheless, it does enable the non-invasive and accurate measurement of GT.

Few reports have been published regarding whether measured GT is associated with probe visibility (PV) as assessed using a periodontal probe [13]. Although gingival biotype classification is closely related to the morphological features of the tooth and periodontium [14], most clinicians prefer to use a periodontal probe to assess GT.

The primary purpose of this study was to assess whether periodontal PV can be used to accurately evaluate the GT around the maxillary central incisors. The secondary aim was to identify the periodontal biotype and related factors using cluster analysis based on various morphological characteristics.

MATERIALS AND METHODS

This observational, cross-sectional study of the relationship between PV and GT was approved by the Institutional Review Board (IRB) of Asan Medical Center and was conducted at the Department of Periodontics at Asan Medical Center (IRB No. 2012-0119). All participants provided written informed consent in accordance with the Declaration of Helsinki.

Participants
The study included 90 Koreans who were 16–40 years of age and had all of their maxillary anterior teeth, with a probing depth (PD) of ≤4 mm. The exclusion criteria were: 1) crown restorations or fillings involving the incisal edge on both maxillary central incisors, 2)
attrition or abrasion of the maxillary central incisors, and 3) use of medications with any
known effect on the periodontal soft tissues in the previous year (for example, non-steroidal
anti-inflammatory drugs, antibiotics, immunosuppressants, and calcium channel blockers).

Sample size calculation
The minimal sample size required to ensure adequate power was calculated using a method
adopted in a previous study involving the same ultrasonic device used in this study (Krupp
SDM®, Austenal Medizintechnik, Cologne, Germany) [15]. The standard deviations of the
thin and thick groups were 0.25 and 0.30, respectively. An alpha level of 0.05 and a power
of 0.9 were used to calculate the sample size needed to detect the difference between the
groups. The required minimum sample size was 32 subjects, and the target sample size was
set at 90 subjects for cluster analysis.

Clinical examination
Seven clinical parameters for both maxillary central incisors were systematically recorded by
a single examiner (J. P.) (Figure 1).

1) Clinical photographs of the maxillary anterior teeth (frontal view) were obtained and
stored for the record.
2) PV was evaluated by probing the mid-facial sulcus. We determined whether the outline
of the underlying probe (CPU 15UNC, Hu-Friedy, Chicago, IL, USA) at the base of the
sulcus was visible through the gingival margin [6]. Each subject was assigned a PV score
as follows [14]:
   0: visible at both central incisors
   1: visible at 1 central incisor
   2: not visible at either central incisor
3) PD was measured to the nearest 0.5 mm at the mid-facial point for both central incisors.
4) Gingival width (GW) was measured to the nearest 0.5 mm at the mid-facial point for
both central incisors using a periodontal probe. This parameter was defined as the
distance from the free gingival margin to the mucogingival junction. The average of the
values obtained from both central incisors was used in the analysis.
5) Papilla height (PH) was assessed at the mesial aspect of both central incisors using the
same periodontal probe that was used to measure GW. The distance from the top of the

![Figure 1. Measurement of the clinical morphometric characteristics of the maxillary central incisors. PD: probing depth, PV: probe visibility, PH: papilla height, CW: crown width, CL: crown length.](https://doi.org/10.5051/jpis.2003880194)
papilla to a line connecting the mid-facial soft-tissue margin of the 2 central incisors was measured to the nearest 0.5 mm [5].

6) GT was measured with an ultrasonic device at the marginal gingiva of both central incisors, 2 mm apical to the mid-facial gingival margin. The measurement was performed 6 times per site; the highest and lowest values were excluded, and the average of the 4 remaining values was taken and used in the analysis, in accordance with previous studies [10].

7) The ratio of crown width to crown length (CW/CL) for both central incisors was assessed according to the methods described by Olsson and Lindhe [16]. CL was measured from the incisal edge of the crown to the free gingival margin or, if discernible, the cemento-enamel junction at the mid-facial point. Values were recorded to the nearest 0.1 mm using a caliper. The crown was divided into 3 equal portions along its length, and the CW was measured at the junction of the middle and cervical regions.

**Intra-examiner repeatability**
PT and GT were re-examined by the same clinician after 2 weeks for 1 randomly-selected subject out of every 10 volunteers (n=9 in total). The intra-examiner reliability for GT as measured with the ultrasonic device was assessed using the intraclass correlation coefficient, and repeated measurements of PV were evaluated using the Cohen kappa coefficient.

**Statistical analysis**
For all continuous variables, means and standard deviations were calculated for each subject. The frequency distribution of GT was analyzed by means of percentile agreement. Differences in the PV score and GT between male and female participants were tested using analysis of variance (ANOVA), the χ² test, or the independent-samples t-test. The capacity of PV to estimate GT was assessed via comparison with direct measurements made using an ultrasonic device, and this capacity was reported as the area under the receiver operating characteristic curve (ROC-AUC).

Multivariate analysis using multinomial baseline-category logistic regression was performed to investigate the association between the PV score and the following clinical explanatory variables: sex, GT, PD, PH, GW, and CW/CL.

Hierarchical clustering, widely applied in cluster analysis, was used to classify the gingival biotype of each patient. Several algorithms are available for merging clusters, of which the method described by Ward [17] was chosen. Agglomerative hierarchical cluster analysis was performed to identify groups with similar values for PD, PH, GW, and CW/CL. Before analysis, all parameters were standardized so as not to affect the squared Euclidean distance. Each case was initially treated as a separate cluster (i.e., equal numbers of clusters and cases), and cases were then combined sequentially into new clusters. This process was repeated until only 1 cluster remained. The similarity measure (squared Euclidean distance) and clustering algorithm (Ward) are both commonly used and known to be efficient at hierarchical clustering. Semi-partial R square, pseudo-F statistics, and pseudo-t² statistics were used to determine the optimal number of clusters.

Clusters were compared with regard to PD, PH, GW, and CW/CL. Differences between clusters were assessed using ANOVA or the χ² test. Statistical analyses were performed using SAS 9.2 software (SAS Institute Inc., Cary, NC, USA). A P value <0.05 was considered to indicate statistical significance.
RESULTS

Study subjects
In total, 90 periodontally healthy participants (46 male and 44 female) were recruited for the study. The mean age was 29.7 years (range, 16–40 years).

Intra-examiner repeatability
The reliability of the silent drive motor measurements was consistently very high (intraclass correlation test, \( r=0.939 \)). The repeatability of the PV assessment was assessed using the Cohen kappa statistic. The reproducibility was verified as substantial, with a kappa value of 0.61 (\( r=0.026 \)).

Frequency distribution of PV score
With regard to stratification by PV score, 21 subjects (23.3%) had a score of 0, 21 subjects (23.3%) had a score of 1, and 48 subjects (53.3%) had a score of 2 (Table 1).

The distribution of men and women was not equal between the PV groups (Pearson \( \chi^2 \) test, \( r=0.003 \)). More than half of the men (65%) had a PV score of 2 compared with 41% of women; in addition, 39% of women had a PV score of 0 compared with only 8.7% of men.

Relationships between PV score and morphological parameters
The baseline characteristics of 5 clinical parameters of the study subjects, grouped by PV score, are presented in Table 2. The group with a PV score of 0 had relatively low GT, PH, and GW values and a relatively high CW/CL. However, none of the morphological variables (PD, PH, GW, or CW/CL) were significantly related to the PV score (\( r>0.05 \)).

Relationships between PV score and GT
The mean GT values in the PV=0, PV=1, and PV=2 groups, respectively, were 1.33 mm, 1.42 mm, and 1.48 mm (total mean, 1.43 mm). No significant difference in GT was found among groups (1-way ANOVA, \( r=0.152 \)) (Table 2). The mean GT for men was 1.50 mm, while that for women was 1.38 mm; the difference between these values was not statistically significant (Student \( t \)-test, \( r=0.138 \)).

Table 1. Frequency distribution of PV score for 90 subjects

| Variable | PV score 0 | PV score 1 | PV score 2 |
|----------|-----------|-----------|-----------|
| Male     | 4 (8.7)   | 12 (26.1) | 30 (65.2) |
| Female   | 17 (38.6) | 9 (20.5)  | 18 (40.9) |
| Total    | 21 (23.3) | 21 (23.3) | 48 (53.3) |

Values are presented as number (%).
PV: probe visibility.

Table 2. Baseline morphological characteristics, grouped by PV score

| Variable | PV score 0 | PV score 1 | PV score 2 |
|----------|-----------|-----------|-----------|
| GT (mm)  | 1.33±0.29 | 1.42±0.21 | 1.48±0.33 |
| PD (mm)  | 1.43±0.42 | 1.43±0.36 | 1.34±0.43 |
| PH (mm)  | 4.36±0.88 | 4.60±0.70 | 4.68±0.84 |
| GW (mm)  | 4.71±1.71 | 5.18±0.83 | 5.14±1.11 |
| CW/CL (%)| 72.20±10.40 | 70.04±11.86 | 70.62±8.25 |

Data are shown as mean±standard deviation.
PV: probe visibility, PV score 0: probe visible at both central incisors, PV score 1: probe visible at one central incisor, PV score 2: probe visible at neither central incisor, GT: gingival thickness, PD: probing depth, PH: papilla height, GW: gingival width, CW/CL: ratio of crown width to crown length × 100 (%).
The ROC-AUC for PV as measured using an ultrasonic device was 0.63; a cut-off value of 1.38 yielded 66.7% sensitivity and 59.0% specificity (Figure 2). The low ROC-AUC value indicates the presence of a weak inter-relationship between PV and GT; therefore, the derived cut-off value is barely acceptable.

**Multivariate analysis**

Multivariate analysis was used to model the dependent variable PV score, with sex, GT, PD, PH, GW, and CW/CL as explanatory variables (Table 3). Sex was the only highly significant predictor of PV score (odds ratio, 6.48; 95% CI, 1.71–24.56; \( P < 0.05 \)).

**Table 3. Multivariate multinomial logistic regression analysis**

| Variable | PV score | OR | 95% CI       | \( P \) value |
|----------|----------|----|--------------|---------------|
| Sex      | 2 (ref)  | 1  |              |               |
|          | 1        | 1.268 | 0.427–3.767  | 0.669         |
|          | 0        | 6.480  | 1.710–24.556 | 0.006         |
| GT       | 2 (ref)  | 1  |              |               |
|          | 1        | 0.443  | 0.054–3.631  | 0.448         |
|          | 0        | 0.120  | 0.011–1.377  | 0.089         |
| PD       | 2 (ref)  | 1  |              |               |
|          | 1        | 1.583  | 0.422–5.939  | 0.496         |
|          | 0        | 1.007  | 0.214–4.737  | 0.993         |
| PH       | 2 (ref)  | 1  |              |               |
|          | 1        | 0.842  | 0.391–1.814  | 0.661         |
|          | 0        | 0.377  | 0.150–0.947  | 0.038         |
| GW       | 2 (ref)  | 1  |              |               |
|          | 1        | 1.106  | 0.642–1.907  | 0.716         |
|          | 0        | 0.777  | 0.447–1.348  | 0.369         |
| CW/CL    | 2 (ref)  | 1  |              |               |
|          | 1        | 0.988  | 0.926–1.054  | 0.719         |
|          | 0        | 1.010  | 0.941–1.085  | 0.775         |

Baseline category indicated by (ref). PV: probe visibility, OR: odds ratio, CI: confidence interval, PV score 0: probe visible at both central incisors, PV score 1: probe visible at 1 central incisor, PV score 2: probe visible at neither central incisor, ref: reference, GT: gingival thickness, PD: probing depth, PH: papilla height, GW: gingival width, CW/CL: ratio of crown width to crown length \( \times 100 \) (%).

*Sex was the only significant predictor of PV.*
Cluster analysis

Cluster analysis was based on the morphometric parameters of PD, PH, GW, and CW/CL. Two clusters were created as a result of the clustering procedures. Table 4 presents descriptive features of each cluster. Cluster A comprised 50 participants (28 male and 22 female), whereas cluster B consisted of 40 participants (18 male and 22 female).

Clusters A and B differed significantly in terms of the mean values of PD, PH, GW, and CW/CL. Compared with cluster B, cluster A subjects had a more slender tooth form (CW/CL, 65.52% vs. 77.51%), greater PH (4.96 mm vs. 4.11 mm), shallower PD (1.21 mm vs. 1.61 mm), and narrower GW (4.57 mm vs. 5.66 mm). The morphological characteristics of cluster A subjects corresponded to those of what was previously known as the scalloped biotype, whereas those of cluster B subjects corresponded to the flat biotype.

The mean GT of the cluster A (1.36±0.29 mm) subjects was significantly lower than that of the cluster B subjects (1.52±0.29 mm) (Student t-test, P=0.015). However, the PV score was statistically similar between cluster A and B subjects (Student t-test, P=0.583).

DISCUSSION

The present study indicates that the biotype around the maxillary central incisors as assessed by periodontal PV does not accurately reflect GT. Sex was the only significant predictor of periodontal PV. Two morphological periodontal biotypes were identified (scalloped and flat), and these biotypes had different GT values as measured directly using an ultrasonic device.

The finding that periodontal PV does not accurately reflect GT is at odds with results from previous studies in which gingival biotype was determined using this method. Kan et al. [13] reported that the reliability of a periodontal probe was comparable with that of direct measurements using calipers after tooth extraction, and they suggested gingival biotype classification by millimeters of GT. Differences in measurement procedures, probe color, and study design, along with the absence of a supporting tooth, may explain the discrepancy between our results and those of Kan et al. [13]

The finding that sex was the only significant predictor of periodontal PV is supported by De Rouck et al. [14] Factors including a lighter skin color may explain the greater transparency.

Table 4. Morphological characteristics of clusters A and B

| Variable     | Cluster   | P value  |
|--------------|-----------|----------|
| PD           | A (n=50)  | 1.21±0.37| B (n=40)  | 1.61±0.35| <0.000 |
| PH           | 4.96±0.74 |          | 4.11±0.66 |          | <0.000 |
| GW           | 4.57±0.91 |          | 5.66±1.30 |          | <0.000 |
| CW/CL        | 65.52±8.33|          | 77.51±6.47|          | <0.000 |
| GT           | 1.36±0.29 |          | 1.52±0.29 |          | 0.015  |
| PV score     |           |          | 0.583    |
| 2            | 29 (60.42)|          | 19 (39.58)|          |
| 1            | 11 (52.38)|          | 10 (47.62)|          |
| 0            | 10 (47.62)|          | 11 (52.38)|          |

Data are shown as mean±standard deviation or number (%).

PD: probing depth, PH: papilla height, GW: gingival width, CW/CL: ratio of crown width to crown length × 100 (%), GT: gingival thickness, PV: probe visibility, PV score 0: probe visible at both central incisors, PV score 1: probe visible at 1 central incisor, PV score 2: probe visible at neither central incisor.
of gum tissue in women than in men. Skin color is affected by the quantity and structure of melanin pigments, blood vessel distribution, and non-melanin pigments such as hemoglobin. The skin color of normal Asian subjects as measured by spectrophotometry, as well as melanin and erythema indexes, are significantly higher in men than in women [18]. Hemoglobin is red, and its concentration is thought to contribute significantly to gingival transparency. Feiner et al. [19] reported that ethnicity and sex were predictive of errors in pulse oximetry, primarily because women tend to have lower hemoglobin levels than men; however, it was not possible to separate the contributions of sex from those of low hemoglobin. From an optical perspective, this may explain our finding of greater PV in female subjects.

Our finding of 2 different morphological groups, scalloped and flat, is consistent with existing knowledge of periodontal biotypes. De Rouck et al. [14] previously reported that 3 clusters (thin-scalloped, thick-scalloped, and thick-flat) encompassed 2 morphologically different groups, the subgroups of which are determined by PV. Interestingly, this study showed that GT, as measured directly using an ultrasonic device, differs significantly between morphological clusters, albeit by the very small amount of 0.2 mm. Therefore, it is not unreasonable to hypothesize that an association exists between GT and tooth morphology. Cluster analysis based on GT (as measured using an ultrasonic device), GW, and CW/CL in 42 subjects resulted in 3 clusters: normal thickness-normal crown ratio, thick-quadratic tooth, and an unknown combination [7]. The third group was subsequently revised according to GW. This finding supports the proposed hypothesis of an association between GT and tooth morphology. However, in the current study, it was found that periodontal PV is unrelated to patient group clustered by morphological characteristics, suggesting that PV does not always correspond to gingival biotype.

Several groups have measured GT around the maxillary central incisors using an ultrasonic device [7,11,15,20,21]. The mean GT in the current study (1.43 mm) was greater than that reported previously; for example, 1.28 mm was reported in a study conducted in Germany [7], and 1.2 mm was reported in a Korean population [20]. However, our measured value is comparable to that acquired previously using a syringe needle (1.46 mm) [5]. Variations in pressure applied to the gingiva when taking measurements, in the angle formed between the rounded gingival surface and the measuring probe, and in the amount of water used as a medium are all potential sources of variability that may account for the discrepancies between studies.

Recently, gingival biotypes have been assessed radiographically using cone-beam computed tomography (CBCT) scans [22,23]; the results were used to evaluate whether any association exists between periodontal treatment outcome and anatomical characteristics [24,25]. However, these studies also acknowledged the limitations and paucity of data regarding the accuracy of the various techniques used to measure tissue thickness, including CBCT.

The present study does have some limitations. First, GT may vary within subjects at different sites in the mouth, as indicated by the high proportion of subjects with a PV score of 1. This implies that the gingival biotype determined at 1 specific site is not necessarily representative of the overall gingival biotype. Second, our periodontal PV observations may be affected by relatively weak intra-examiner measurement repeatability. For a number of subjects, PV was difficult to determine because the probe was only recognizable in the coronal part of the sulcus. PV was determined when the outline of the probe, including the apex, could be
identified; therefore, gingival biotype determination using a periodontal probe is affected by each examiner’s subjective decision-making [26]. To counteract this, Cook et al. [27] limited the classification of PV into just 2 groups: thin and thick/average.

In conclusion, periodontal PV is not related to GT as measured directly using an ultrasonic device. Sex was the only highly significant predictor of periodontal PV. The presence of a significant correlation between GT and morphological characteristics of the periodontium was confirmed. Clinically, measuring the more accurate GT instead of PV and considering the relevant morphological features can enable the proper determination of gingival biotype.

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