Use of host- and bacteria-derived salivary markers in detection of periodontitis: A cumulative approach

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Abstract. In the present study, we propose a novel diagnostic approach, using 3 different salivary markers, representing periodontal pathogen burden, inflammation, and tissue degradation, for detecting periodontitis. The salivary concentrations of Porphyromonas gingivalis, interleukin-1β, and matrix metalloproteinase-8, available from salivary specimens of 165 subjects (84 subjects with advanced periodontitis and 81 controls), were calculated together to obtain a cumulative risk score (CRS). In the calculation of CRS, the concentrations of each marker were divided into tertiles, and cumulative sub-score per each subject were calculated by the multiplication of the tertile values. Three CRS groups, indicating the lowest, medium, or highest risk, were formed with the cumulative sub-scores. Logistic regression analysis and ROC curves were performed to study the association of CRS with periodontitis. The results indicate that CRS, calculated from the 3 salivary biomarkers, is associated with advanced periodontitis more strongly than any of the markers individually. CRS offers a novel, non-invasive model for advanced periodontitis risk categorization that is especially useful in large population surveys where a periodontal examination is not feasible.

Keywords: IL-1β, MMP-8, Porphyromonas gingivalis, risk score, saliva

1. Introduction

Periodontitis forms a group of infection-induced inflammatory diseases, initiated by imbalance between pathogenic microorganisms in subgingival biofilm, affecting tooth-supporting soft tissues and alveolar bone [1]. The traditional diagnosis of periodontitis is based on clinical and radiographic assessments of periodontal tissues. However, when assessing clinically the periodontal health status in large-scale health surveys, a full periodontal examination turns to be laborious and expensive, requiring trained dental professionals [2].

The most desirable goal in health care promotion and delivery would be to monitor the periodontal health status and the onset, progression, and treatment outcome of periodontal diseases through non-invasive means. As an easily collected and non-invasive specimen, saliva has been used as a diagnostic fluid in medicine [3, 4]. Elevated levels of bacteria, bacterial products (lipopolysaccharide; LPS, enzymes), host-originated enzymes, cytokines, and bone turnover biomarkers have been found in saliva of periodontitis patients in relation to their controls [3, 4].
During the onset of periodontitis, markers of inflammation, such as interleukin (IL)-1β, IL-6, and tumor necrosis factor (TNF)-α, are released from cells of the junctional epithelium. In the course of the disease, a number of enzymes, such as matrix metalloproteinase (MMP)-8, MMP-9, and MMP-13, are produced and activated by host cells in a cascade leading to degradation of tooth-supporting soft and bone tissues [1,5,6]. As a result of bone degradation, pyridinoline cross-linked carboxyterminal telopeptide (ICTP) and osteocalcin are released into periodontal tissues and, eventually, into saliva [3,7–11]. Periodontal pathogens initiate the inflammatory conditions; hence, their simple carriage or presence in high numbers in saliva have been associated with periodontitis [12–18].

Various oral bacteria, their by-products, cytokines, host enzymes, and biomarkers of bone turnover have been analyzed from saliva aiming to find a valuable marker in distinguishing the subjects with periodontitis and periodontally healthy subjects [3,4]. When searching a potential salivary biomarker for detecting periodontitis, a common approach in cross-sectional studies is to look for a statistical difference ($p < 0.05$) between periodontitis and control subjects and to determine a meaningful threshold for the tested marker. So far, this aim has not been fulfilled. Many inconsistencies in the current literature are generally attributed to the heterogeneity of detection techniques as well as local and systemic factors that may affect the detection level of examined markers in saliva.

Our hypothesis is that the increase or the decrease in bacterial burden, inflammatory response, and tissue destruction do not occur simultaneously but consecutively at different sites of periodontal tissues (Fig. 1). Bacterial LPS and enzymes, and cytokines and enzymes produced by host cells work as the major determinants of this cascade. When bone resorption is at its highest level due to the increased stimulation of osteoclastogenesis by inflammatory cytokines and neutrophilic enzymes, the same response suppresses the level of bacterial burden. This means that a bacteria-derived biomarker can be present at a low level, while bone resorption is at its highest level. Also, a biomarker responsible for bone degradation may fail to be connected to the disease at a phase when the microbial attack is at its highest level.

In the present study, we used clinical, microbiological, and inflammatory data available from our recent studies [7,13,14] and propose a novel statistical approach for diagnosing periodontitis to be used, in particular, in public health surveys or in large cross-sectional studies. In this approach, the salivary concentrations of selected biomarkers, representing periodontal pathogen burden, inflammation, and tissue degradation, have been combined to obtain a cumulative risk score (CRS).
2. Materials and methods

2.1. Data collection

The present study population is a subsample of the nationally representative population of the “Health 2000 Health Examination Survey”. A total of 8,028 subjects, aged ≥30 years and steadily living in Finland, participated in the health examination in the field, divided in 5 districts, where data on their general and oral health were collected. Periodontal measurements from 4 sites per tooth included probing pocket depths (PPD) and bleeding on probing. Each tooth was recorded as having PPD < 4 mm, PPD ≥ 4 mm, or PPD ≥ 6 mm. Paraffin-stimulated whole saliva samples were collected from 1,294 of the Health 2000 subjects in the southern district, and frozen at −70°C until further used [12]. Information about data collection methods can be found in the report of the National Public Health Institute (http://www.terveys2000.fi/julkaisut/oralHealth.pdf). All protocols had been approved by the institutional ethics committees.

Salivary data used in the present analysis were obtained from a subpopulation of 165 subjects, divided into 2 extreme groups on the basis of their periodontal status; 84 advanced periodontitis subjects with at least 14 teeth with PPD of ≥4 mm (mean age 49.6 years) and 81 control subjects with no pocket teeth (mean age 47.9 years). All subjects had at least 20 teeth. The detailed information on the techniques used for the salivary analyses of periodontal pathogen burden and inflammatory and tissue destruction markers can be found elsewhere [7,13,14]. Briefly, frozen salivary samples were thawed and centrifuged, and the supernatants were aliquoted. The salivary concentrations of the 3 markers used in the present study were detected as follows: IL-1β with a commercial ELISA kit (Amersham, GE Healthcare, Buckinghamshire, UK), MMP-8 with a time-resolved immunofluorometric assay (IFMA), and Porphyromonas gingivalis with a quantitative real-time PCR assay [7,13,14].

2.2. Selection of the biomarkers for calculating CRS

On the basis of the association with periodontitis, we previously concluded that the salivary concentrations of IL-1β, MMP-8, tissue inhibitor of matrix metalloproteinase (TIMP) -1, and ICTP, and the presence of pathogenic species in saliva could potentially be used as markers of periodontitis (Table 1) [7,13,14]. From the examined host- and bacteria-derived markers, P. gingivalis, IL-1β, and MMP-8 were selected as the most potential markers of periodontal pathogen burden, inflammation, and tissue destruction, respectively, for present analyses.

2.3. Calculation of cumulative risk scores (CRS)

For the calculation of CRS, the salivary concentrations of P. gingivalis, IL-1β, and MMP-8, which were separately analyzed in our previous studies [7,13,14], were statistically re-analyzed with a novel approach as follows.

The salivary concentrations of each marker, originally collected as continuous data, were divided into tertiles 1–3 (Table 2). A cumulative sub-score per each subject was calculated by the multiplication of the corresponding tertile values of P. gingivalis, IL-1β, and MMP-8. According to this calculation, the subject’s cumulative sub-score can be 1, 2, 3, 4, 8, 9, 12, 18, or 27. Furthermore, based on these cumulative sub-scores, 3 CRS groups were formed as follows:

- **CRS I**: Lowest risk (the cumulative sub-scores of 1, 2, 3)
- **CRS II**: Medium risk (the cumulative sub-scores of 4, 8, 9)
- **CRS III**: Highest risk (the cumulative sub-scores of 12, 18, 27)
Table 2
Tertile values of *P. gingivalis*, IL-1β, and MMP-8 were calculated from the salivary concentrations of each marker that were originally collected as continuous data.

| Tertile value | *P. gingivalis* (GE/ml) | IL-1β (pg/ml) | MMP-8 (ng/ml) |
|--------------|-------------------------|---------------|---------------|
|              | minimum | maximum | minimum | maximum | minimum | maximum |
| 1            | 0.00  | 387.2    | 35.14  | 411.9    | 2.8  | 315.8 |
| 2            | 434.5 | 18,764.4 | 419.8  | 685.1    | 318.0 | 780.6 |
| 3            | 19,980 | 51,373,537.5 | 716.9  | 1260.1   | 785.3 | 2,576.6 |

CRS III: Highest risk (the cumulative sub-scores of 12, 18, 27)

For example, a subject with salivary concentrations of 550 GE/ml for *P. gingivalis*, 600 pg/ml for IL-1β, and 300 ng/ml for MMP-8 will have tertile values of 2, 2, and 1, correspondingly (Table 2). When these tertile values are multiplied (2 × 2 × 1), a cumulative sub-score will be 4, and the subject will be categorized as being at medium risk of having periodontitis (CRS II).

### 2.4. Statistical analysis

The SPSS statistical program (Version 15.0; SPSS Inc., Chicago, IL, USA) was used to analyze the data. An ANOVA test was used to compare the biomarker concentrations between the periodontitis and control groups. A logistic regression analysis and receiver operating characteristics (ROC) curves were performed to study the association of *P. gingivalis*, IL-1β, MMP-8, and CRS as tertiles with periodontitis.

### 3. Results

For the calculation of CRS, the salivary concentrations of *P. gingivalis*, IL-1β, and MMP-8 were selected due to their high prevalence in the population and significantly different levels between periodontitis and control subjects. The reasons to exclude the other markers (Table 1) were as follows: the low prevalence of *Aggregatibacter actinomycetemcomitans*, *Treponema denticola*, and *Prevotella intermedia* among the study population, the weak statistical significance of ICTP values between the periodontitis and control groups and the lacking diagnostic contribution of *Tannerella forsythia* and TIMP-1 to the present model.

Table 3 presents the odds ratios and area under curve (AUC) values with 95% confidence intervals separately for each of the 3 selected biomarkers and cumulatively as CRS. Each marker had an association with periodontitis at a different level. The association increased considerably by using CRS, which had 2 to 3 times higher odds ratios in comparison with single biomarkers. The increase in the diagnostic accuracy was also demonstrated in the ROC analyses, where the AUC value increased from 0.694–0.710 up to 0.766.

### 4. Discussion

The present study proposes a novel periodontitis risk categorization model, taking into account periodontal pathogen, inflammation, and tissue degradation markers in saliva. The strength of this new approach is that when one of the 3 biomarkers, e.g., the amount of periodontopathogenic bacteria, is in remission due to the strong host response, the concentration of other 2 biomarkers, associated with inflammation and tissue degradation, will be exacerbated, but CRS will still identify the subjects with the disease. This will diminish the number of false negatives which are often obtained in single-marker analyses.

Even after 3 decades of work, there is not yet a clear consensus on the strength of associations between salivary biomarkers and periodontal diseases. A good example of this controversy is the detection of pathogenic bacteria in saliva and its association with periodontitis. According to Paju et al. [15], who analyzed the saliva samples of 1198 adult dentate subjects, the number of pathogenic species rather than a certain periodontal pathogen is associated with periodontitis. In contrast, Leblebicioglu et al. [16] concluded that the presence of periodontopathogenic pathogens in saliva correlates with signs of gingivitis but not with alveolar bone loss (periodontitis). Interestingly, in a study by Sawamoto et al. [17], salivary *A. actinomycetemcomitans* levels proved to be higher in periodontally healthy subjects than in periodontitis patients. Hamlet et al. [18] suggested that the deepened pocket depth, a clinical marker of periodontitis, is associated with the presence of *P. gingivalis* and *A. actinomycetemcomitans* but not with that of *P. intermedia*. However, in a more recent study by K{"o}n{"o}nen et al. [12], the number of teeth with pocket depths of $\geq$ 4 mm was associated with the salivary detection rates...
Table 3

| Risk group  | No. of periodontitis/ control subjects | OR (95% CI)       | AUC (95% CI)       |
|-------------|---------------------------------------|-------------------|-------------------|
| P. gingivalis | III 43/12                             | 8.01 (3.39–18.89) | 0.710 (0.631–0.789) |
|             | II 24/31                              | 1.7 (0.7–3.7)     | 0.694 (0.613–0.775) |
|             | I 17/38                               |                   | 1                 |
| IL-1β       | III 38/17                             | 6.54 (2.8–15.06)  | 0.694 (0.613–0.775) |
|             | II 32/23                              | 4.07 (1.81–9.15)  |                   |
|             | I 14/41                               | 1                 |                   |
| MMP-8       | III 41/14                             | 7.81 (3.34–18.24) | 0.710 (0.631–0.789) |
|             | II 28/27                              | 2.76 (1.24–6.12)  |                   |
|             | I 15/40                               | 1                 |                   |
| CRS         | III 38/7                              | 19.9 (7.11–55.65) | 0.766 (0.694–0.839) |
|             | II 34/30                              | 4.7 (1.26–12.51)  |                   |
|             | I 12/44                               | 1                 |                   |

of *P. gingivalis, P. intermedia, T. denticola*, and *A. actinomycetemcomitans*. Such differences in the results can also be observed in studies where various cytokines have been analyzed from saliva. For example, salivary IL-1β, a proinflammatory cytokine predominantly secreted by neutrophils and macrophages, has been either associated with periodontitis [3,14], or not [19]. Similarly, higher levels of TNF-α have been found in salivary samples of periodontitis subjects than in those of periodontally healthy controls [20], while other researchers have failed to show such a difference [14, 21]. Usually, these discrepancies have been explained by differences in the study population, such as the age range, percentage of smoking subjects, or number of study subjects. For example, cigarette smoke inhibits the enzyme activity due to the interaction with aldehydes in smoke and host-derived enzymes [22]. Also, the detection method used has influence on the outcome; when analyzing MMP-8 from saliva, mouthrinse, or gingival crevicular fluid, IFMA results in a more reliable distinction of periodontitis subjects from their controls than by ELISA with different antibodies [7,23, 24].

It is possible that the episodic character of periodontitis plays a central role in the level of salivary biomarkers. Chronic periodontitis, the most prevalent form of destructive periodontal disease, has dynamic states of exacerbation and remission [25,26]. It is usually related to a polymicrobial infection, *T. denticola, P. gingivalis,* and *T. forsythia* being considered the major players [27]. With the increase in bacterial numbers and their virulence factors, the local inflammatory reaction in periodontal tissues activates the immune system [28]. The initial inflammatory response aims to limit bacterial invasion; hence, it promotes the infiltration of neutrophils and macrophages to the site of infection. This infiltration is achieved by the secretion of proinflammatory cytokines from epithelial cells and fibroblasts. After their migration to inflamed tissues, leukocytes suppress the bacterial invasion by their oxygen-dependent or -independent mechanisms. Subsequently, T- and B-cells emerge to the sites of infection and secrete immunoglobulins as an antigen-specific response [29,30]. While the host defense suppresses the level of infection by eliminating pathogens, the increased expression of proinflammatory cytokines, such as IL-1, -6, -11, and -17, and TNF-α, stimulates alveolar bone resorption. These cytokines can induce osteoclastogenesis by increasing the expression of Receptor Activator for Nuclear Factor-κB Ligand (RANKL) and by decreasing the osteoprotegerin (OPG) production in osteoblasts/stromal cells [31,32]. Matrix metalloproteinases, especially MMP-8 and -13, can contribute to the resorption of alveolar bone. With decreasing bacterial challenge, anti-inflammatory cytokines, such as IL-4, -10, -12, -13, and -18, and lipid mediators inhibit the inflammation in periodontal tissues and bone resorption [32].

Salivary diagnostics in periodontology is not anymore trying to find a silver bullet in detection of periodontitis [33]. Therefore, the presence of multiple periodontopathogenic bacteria [13,15], combinations and proportions of host-derived biomarkers [7], or combinations of host-derived biomarkers with periodontal pathogens [34] have been introduced as potential approaches for newly developed point-of-care tests in diagnosing oral and systemic diseases [6,23,24]. In above-mentioned studies, a combinational value has
been calculated as probabilities in a logistic regression model, whereas the present CRS was calculated by multiplying the tertile values of each marker. One practical improvement of the present proposal is to define universal reference ranges (high, medium, low) for the salivary levels of each biomarker. In that way CRS would be comparable between different study populations.

In the present model, the selection of the 3 salivary markers, *P. gingivalis*, IL-1β, and MMP-8, can be taken as an example. The major obstacle in salivary diagnostics is that several markers of periodontitis, such as TNF-α, IL-6, RANKL, ICTP, or OPG, are found at very low levels in saliva where they are highly diluted unlike in gingival crevicular fluid. This fact limits the comparisons between periodontitis and control subjects. With the improvement of salivary detection techniques, for example, with new proteomic analyses, various bacterial, inflammatory, or tissue degradation markers can be applied in the current model. In the present study, however, the selection of the 3 disease markers is still unique, since *P. gingivalis* can efficiently process and activate IL-1β and MMP-8, thus inducing the bacteria-host destruction cascade in periodontitis [34–36]. *P. gingivalis* can directly stimulate collagen degradation via activation of pro-MMP-8 by cleaving the “prodomain” with their gingipains; however, in the pathogenesis of periodontitis, the significance of this effect is not yet clarified.

As saliva is a pooled fluid of the whole mouth, the present approach is not suitable for gingival crevicular fluid. It is notable that salivary diagnostics basically aims to detect subjects with periodontitis but not to localize the disease in the dentition. Therefore, our present approach uses multiplication to result in a cumulative effect of several individual markers, even when present at low levels in saliva. The strength of CRS is in its potential in public health surveys or large cross-sectional studies with different age groups. In health care promotion and delivery in periodontics, the demonstrated potential of saliva enables to monitor the periodontal health status through non-invasive means.

5. Conclusion

The use of the 3 salivary biomarkers, representing periodontal pathogen burden, inflammation, and tissue degradation, and calculating a cumulative risk score, based on these markers, reveal periodontitis more accurately than analyzing any of the salivary biomarkers alone.

Acknowledgements

The authors declare that they have no conflicts of interests. The present study is part of the Health 2000 Health Examination survey, organized by the National Institute for Health and Welfare of Finland, and financially supported by the Finnish Dental Association and the Finnish Dental Society. These series of studies were supported by grants from the IADR and the Orion, Paulo, Yrjö Jahnsson (grant 5639 for U. K. G) and Finnish Cultural Foundations (for U. K. G and grant 130408 for T. S.), the Academy of Finland (grant 78443 for E. K. and grant 118391 for P. J. P), the Sigrid Juselius Foundation (for P. J. P) and the Helsinki University Central Hospital Research Foundation. Timo Sorsa is an inventor of US-patents 5652227, 5736341, 5866432, and 6143476.

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