The periodontium of periodontitis patients contains citrullinated proteins which may play a role in ACPA (anti-citrullinated protein antibody) formation

Nesse, Willem; Westra, Johanna; van der Wal, Jacqueline E.; Abbas, Frank; Nicholas, Anthony P.; Vissink, Arjan; Brouwer, Elisabeth; Westra J., [No Value]

Published in: JOURNAL OF CLINICAL PERIODONTOLOGY

DOI: 10.1111/j.1600-051X.2012.01885.x

IMPORTANT NOTE: You are advised to consult the publisher’s version (publisher’s PDF) if you wish to cite from it. Please check the document version below.

Document Version
Publisher’s PDF, also known as Version of record

Publication date:
2012

Link to publication in University of Groningen/UMCG research database

Citation for published version (APA):

Copyright
Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

Take-down policy
If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Downloaded from the University of Groningen/UMCG research database (Pure): http://www.rug.nl/research/portal. For technical reasons the number of authors shown on this cover page is limited to 10 maximum.
The periodontium of periodontitis patients contains citrullinated proteins which may play a role in ACPA (anti-citrullinated protein antibody) formation


Abstract

Aim: To determine the presence and location (stroma versus epithelium) of citrullinated proteins in periodontitis tissue as compared to non-periodontitis tissue and synovial tissue of RA patients.

Materials & Methods: Periodontitis, healthy periodontal and RA-affected synovial tissue samples were collected in addition to buccal swabs. These samples were stained for the presence of citrullinated proteins using polyclonal (Ab5612) and monoclonal (F95) antibodies. Furthermore, Western blotting with F95 was performed on lysates prepared from periodontal and synovial tissues.

Results: In periodontitis stroma, increased citrullinated protein presence (80%) was observed compared with control stroma (33%), the latter was associated with inflammation of non-periodontitis origin. Periodontal epithelium always stained positive for Ab5612. Noteworthy, only periodontitis-affected epithelium stained positive for F95. All buccal mucosal swabs and 3 of 4 synovial tissue samples stained positive for both Ab5612 and F95. Western blotting with F95 showed presence of similar citrullinated proteins in both periodontitis and RA-affected synovial tissue.

Conclusion: Within the periodontal stroma, citrullination is an inflammation-dependent process. In periodontal epithelium, citrullination is a physiological process. Additional citrullinated proteins are formed in periodontitis, apparently similar to those formed in RA-affected synovial tissue. Periodontitis induced citrullination may play a role in the aetiology of rheumatoid arthritis.

Willem Nesse¹, Johanna Westra², Jacqueline E. van der Wal³, Frank Abbas⁴, Anthony P. Nicholas⁵, Arjan Vissink¹ and Elisabeth Brouwer²

¹Department of Oral and Maxillofacial Surgery, University of Groningen and University Medical Center Groningen, Groningen, The Netherlands; ²Department of Rheumatology and Clinical Immunology, University Medical Center Groningen, University of Groningen, Groningen, The Netherlands; ³Department of Pathology and Medical Biology, University Medical Center Groningen, University of Groningen, Groningen, The Netherlands; ⁴Center for Dentistry and Oral Hygiene, University Medical Center Groningen, University of Groningen, Groningen, The Netherlands; ⁵Department of Neurology, University of Alabama, Birmingham, AL, USA

Key words: ACPA; citrullinated proteins; periodontitis; rheumatoid arthritis

Accepted for publication 18 March 2012

Conflict of interest and source of funding statement

The authors declare that they have no conflict of interest. Funding has mostly been made available from the authors’ institutions. Part of this research was funded by a BOOA research grant provided by the Dutch Association of Oral and Maxillofacial Surgeons.
known bacterium to contain a peptidyl arginine deiminase (PAD) enzyme (Wegner et al. 2010). The formation of citrullinated proteins is catalysed by PAD enzymes. Therefore, PAD formed by *P. gingivalis* (PPAD) may cause citrullinated protein formation, which may in turn elicit an antibody response to these proteins, i.e. lead to ACPA formation. Furthermore, citrullinated *z*-enolase peptide 1, is considered to be a major auto-antigen in RA. This immunodominant peptide showed 82% homology with citrullinated enolase from *P. gingivalis*. Thus, ACPA response to synovial citrullinated *z*-enolase may be induced by *P. gingivalis* produced citrullinated enolase in the periodontium (Lundberg et al. 2008). Moreover, antibody titres to *P. gingivalis* are correlated with ACPA titres of RA patients (Mikuls et al. 2009). This implies that *P. gingivalis*, either through PAD, citrullinated enolase or otherwise, contributes to inducing ACPA formation. A third line of evidence implicating periodontitis as an inducer of ACPA formation, stems from the fact that auto-antibody formation commonly occurs in periodontitis. Strikingly, the auto-antibodies produced in periodontitis mainly target collagen (Berglundh & Donati 2005). Furthermore, citrullination is known for its ability to break immunological tolerance (Lundberg et al. 2005), and ACPAs are known for recognizing two or more different citrullinated peptides (van de Stadt et al. 2010). Therefore, an auto-antibody (ACPA) response against citrullinated collagen initiated in the periodontium, might cross react with citrullinated collagen in the synovium.

In summary, ACPAs are thought to play an important role in the aetiology of RA and periodontitis and might, for more than one good reason, be involved in the initiation of ACPA formation (Fig. 1). This study was designed to assess the presence and localization (stromal versus epithelial) of citrullinated proteins in periodontitis tissue compared versus epithelial) of citrullinated proteins in periodontitis tissue compared with non-periodontitis tissue. Although auto-antibody formation is known to occur in periodontitis (Berglundh & Donati 2005), the presence of citrullinated proteins is a pre-requisite for the formation of ACPA.

### Material and Methods

#### Patients and tissue samples

Patients with chronic persistent periodontitis, not responding to initial therapy, are routinely subjected to surgical excision of the inflamed part of their periodontium. Periodontal tissue samples of 15 randomly selected periodontitis patients, without any other known medical condition (including RA), undergoing such surgical excision were obtained. Information on age, gender, smoking (pack years) and periodontitis severity was obtained from patients’ dental records, in which these data were routinely recorded (Table 1). Clinically non-inflamed periodontal tissue samples (no bleeding on probing, no periodontal pockets ≥4 mm) were obtained from six patients undergoing prophylactic removal of impacted third molars (wisdom teeth). Synovial tissue samples of four RA patients, previously obtained during joint replacing surgery, and buccal mucosal cell swabs of three healthy donors known to be positive for anti-perinuclear factor staining, were used as positive control tissues for citrullinated protein staining. Informed consent was obtained from all patients and the institutional review board approved this study.

For Western blotting, periodontal and synovial tissue samples of an additional five, randomly selected, periodontitis patients without any other known diseases, three RA patients, and one RA patient with periodontitis, were used. These patients were not the same as those who provided the tissue samples used for immunohistochemistry.

#### Processing the tissue samples

After excision, periodontal tissue samples of periodontitis patients were symmetrically dissected. Immediately thereafter, half of the sample was frozen using liquid nitrogen, the other half was put in formaldehyde and later embedded in paraffin. Cryostat sections were fixed in acetone and endogenous peroxidase was blocked with 1% hydrogen peroxide (H2O2) in phosphate buffered saline (PBS). Then cryostat sections were incubated with either of two primary
antibodies: (1) anti-citrullinated protein rabbit polyclonal antibody (Ab5612; Millipore, MA, USA) diluted 1:100 in PBS + 1% bovine serum albumin (BSA), and (2) mouse anti-citrullinated protein IgM monoclonal antibody (F95; kindly provided by A.P.Nicholas, University of Alabama, Birmingham, AL, USA) diluted 1:1000 in PBS. These two antibodies were selected for their suitability to be used in immunohistochemistry and are the only two currently available antibodies appropriate for immunohistochemical staining. It is possible that F95, which is raised against a deca-citrullinated peptide, only reacts when multiple citrulline residues are present, whereas AB5612 can also react with single citrulline residues, as this antibody is raised against citrullin-glutaraldehyde BSA.

Prior to blocking endogenous peroxidase with 1% H₂O₂, the cryo-sections were blocked with 10% pool serum in PBS when F95 was used. Finally, detection of primary antibodies was performed using respectively (1) a goat anti-Rabbit (Dako, Glostrup, Denmark) diluted 1:50 in PBS + 1% BSA for Ab5612 and (2) a goat anti-Mouse IgM-HRP (Southern Biotech, AL, USA) diluted 1:500 in PBS + 1% AB serum for F95, followed by using a DAB kit (K3467, Dako).

Cryostat sections of six clinically non-inflamed periodontal tissue samples and four synovial tissue samples of RA patients were stained for citrullinated proteins in a similar manner as periodontal tissue samples from periodontitis patients. Likewise, the buccal mucosal swabs, fixed in acetone, were stained. All tissue samples were haematoxylin and eosin (HE) stained. As control for false positive staining, all tissues were additionally stained using control primary antibodies [a mouse IgM isotype control for F95 (Southern Biotech, clone 11E10, 1:1000 in PBS) and rabbit IgG control for Ab5612 (Southern Biotech, nr. 0111-01, 1:100 in PBS + 1% BSA)], followed by identical detection procedures. Finally, the presence of citrullinated proteins in periodontitis tissue and synovium was analysed qualitatively, i.e. either present or absent.

To obtain information on cell types involved in the inflammatory periodontitis lesions (Table 1), paraffin sections of periodontitis tissues were stained for CD3 (Monosan, Uden, the Netherlands, clone PS1, 1:20; T-cells), CD20 (Dako, clone L-26, 1:200; B-cells), CD138 (IQ Products, Groningen, the Netherlands, clone B-A38, 1:80; plasma cells), CD68 (Dako, clone KP1, 1:100; macrophages), CD1a (Immunotech, Marseille, France, clone O10, R.T.U.; dendritic cells), CD31 (Dako, clone JC 70A, 1:40; endothelial cells) and toluidine blue (mast cells). Staining with antibodies was performed using standard methods as described by the manufacturers.
Table 1. Citrullinated protein presence (+) versus absence (−) in periodontitis patients and controls

<table>
<thead>
<tr>
<th>Variable</th>
<th>Periodontitis Ab5612 + n = 8</th>
<th>Periodontitis Ab5612 − n = 7</th>
<th>p-value</th>
<th>Periodontitis F95 + n = 10</th>
<th>Periodontitis F95 − n = 5</th>
<th>p-value</th>
<th>Periodontitis Either + n = T2</th>
<th>Periodontitis Both − n = 3</th>
<th>p-value</th>
<th>Controls Either weak + n = 2</th>
<th>Controls Both − n = n = 4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age in years*</td>
<td>48.9 (11.4)</td>
<td>36.6 (12.9)</td>
<td>0.55*</td>
<td>55.0 (12.4)</td>
<td>43.2 (16.3)</td>
<td>0.14*</td>
<td>52.3 (14.0)</td>
<td>46.2 (18.5)</td>
<td>0.53*</td>
<td>28.5 (5.5)*</td>
<td>28 (8.6)*</td>
</tr>
<tr>
<td>PPD in mm†</td>
<td>6.6 (1.2)</td>
<td>6.4 (1.2)</td>
<td>0.67*</td>
<td>6.7 (6.2)</td>
<td>6.2 (0.8)</td>
<td>0.50*</td>
<td>6.6 (1.2)</td>
<td>6.0 (0.9)</td>
<td>0.43*</td>
<td>≤3 mm</td>
<td>≤3 mm</td>
</tr>
<tr>
<td>Female sex†</td>
<td>6 (63%)</td>
<td>5 (71%)</td>
<td>0.71†</td>
<td>5 (50%)</td>
<td>5 (100%)</td>
<td>0.053†</td>
<td>7 (58%)</td>
<td>3 (100%)</td>
<td>0.51†</td>
<td>2 (100%)†</td>
<td>2 (50%)†</td>
</tr>
<tr>
<td>Currently smoking¶</td>
<td>4 (50%)</td>
<td>2 (29%)</td>
<td>0.40†</td>
<td>4 (40%)</td>
<td>2 (40%)</td>
<td>1.0†</td>
<td>5 (42%)</td>
<td>1 (33%)</td>
<td>1.0§</td>
<td>1 (50%)§</td>
<td>1 (25%)§</td>
</tr>
<tr>
<td>Packyears</td>
<td>0 (0-23)</td>
<td>0 (0-23)</td>
<td>0.71†</td>
<td>10.5 (15.1)</td>
<td>12.6 (18.3)</td>
<td>0.95†</td>
<td>0 (0-15)</td>
<td>23 (0-40)</td>
<td>0.24‡</td>
<td>Not assessed</td>
<td></td>
</tr>
<tr>
<td>CD3 + cells/field</td>
<td>33 (11-45)</td>
<td>14 (8-45)</td>
<td>0.44‡</td>
<td>31 (12-43)</td>
<td>12 (8-51)</td>
<td>0.55‡</td>
<td>31 (7-39)</td>
<td>11.5 (9-68)</td>
<td>0.55‡</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CD20 + cells/field</td>
<td>9 (0-34)</td>
<td>0 (0-18)</td>
<td>0.23‡</td>
<td>1 (0-20)</td>
<td>12 (0-26)</td>
<td>0.84‡</td>
<td>7 (0-35)</td>
<td>0 (0-13.5)</td>
<td>0.84‡</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CD138 + cells/field</td>
<td>7 (2-27)</td>
<td>24 (5-100)</td>
<td>0.33‡</td>
<td>7 (3-62)</td>
<td>24 (5-66)</td>
<td>0.55‡</td>
<td>7 (3-62)</td>
<td>24 (5-66)</td>
<td>0.55‡</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CD68 + cells/field</td>
<td>3 (0-4)</td>
<td>0 (0-2)</td>
<td>0.13‡</td>
<td>2 (0-5)</td>
<td>0 (0-2)</td>
<td>0.12‡</td>
<td>2 (0-4.5)</td>
<td>0 (0-0)</td>
<td>0.12‡</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CD1a + cells/field</td>
<td>3 (0-6)</td>
<td>1 (0-4)</td>
<td>0.38‡</td>
<td>3 (0-5)</td>
<td>0 (0-4)</td>
<td>0.53‡</td>
<td>2.5 (0-5.5)</td>
<td>0 (0-4)</td>
<td>0.53‡</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CD31 + vessels/field</td>
<td>10 (9-14)</td>
<td>9 (6-15)</td>
<td>0.41‡</td>
<td>10 (6-14)</td>
<td>8 (6-16)</td>
<td>0.34‡</td>
<td>9 (6.5-13)</td>
<td>5.5 (5.5-22.5)</td>
<td>0.34‡</td>
<td></td>
<td></td>
</tr>
<tr>
<td>TBlue cells/field</td>
<td>9 (8-13)</td>
<td>6 (4-24)</td>
<td>0.34‡</td>
<td>10 (8-16)</td>
<td>9 (8-12)</td>
<td>0.35‡</td>
<td>10 (8.5-14.5)</td>
<td>9 (6.5-14)</td>
<td>0.35‡</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*Variables presented as mean (standard deviation), differences tested for significance using Student’s t-test.  †Variables presented as median (interquartile range), differences tested for significance using Mann–Whitney U-test.  ‡Variables presented as numbers (percentages), differences tested for significance using Chi-Square test.  §Differences tested for significance using Fisher’s Exact Test.  NB No p-values were calculated within the control group (participants without periodontitis) because numbers were either identical or very small (1 or 2).
Results

In stroma, citrullinated proteins were detected in 53% \((n = 8)\), 67% \((n = 10)\) and 80% \((n = 12)\) of periodontitis tissue samples using, respectively, Ab5612, F95, or either antibody (Fig. 2a–d, Fig. 3a–d). Of six control periodontal tissue samples, two tissue samples were from persons who had complained of pain and swelling in the previous weeks, although not clinically evident at the time of surgery. Citrullinated proteins were detected in stroma of these two controls only (Figs 2e and 3e citrulline negative control tissue, Figs 2f and 3f citrulline positive control tissue). In one of these two controls this was apparently associated with an inflammatory infiltrate. Thus, citrullinated proteins were detected in stroma of 33% \((2 \text{ in } 6)\) of control tissue samples.

Stroma of only 3 of 15 periodontitis patients was negative for citrullinated proteins, i.e. neither AB5612 nor F95 stained positive. For these three patients, there were no macrophages (CD68) per visible field (Table 1). In contrast, in citrulline positive stroma \((n = 12)\) median number of macrophages per visual field was 2.0 with an interquartile range of 0–4.5 \((p = 0.011)\). This difference was not significant after correcting significance level \(z\) with Bonferroni–Holm correction for multiple comparisons, probably due to small numbers. In Fig. 4, pictures are shown of a citrulline negative and a citrulline positive periodontal tissue to visualize inflammation. Percentage of cell counts in citrulline negative and citrulline positive are given in Fig. 4C.

Likewise, there were no statistically significant differences in age, gender, smoking (current and pack years) and periodontitis severity between periodontitis patients whose stroma contained citrullinated proteins (using Ab5612, F95, or either antibody) and those whose periodontium did not.

In epithelium, citrullinated proteins were detected in periodontitis and control tissue samples whenever using Ab5612. In contrast, when using F95, epithelium stained positive in periodontitis tissue samples only. Citrullinated proteins were detected in three and four of four synovial tissue samples of RA patients using Ab5612 or F95 respectively. The buccal mucosal cell swabs were positive for both antibodies (Fig. 5, only F95 shown).

Western blotting with F95 resulted in a variety of similar bands of citrullinated proteins in all periodontal tissues and in RA-affected synovial tissue with a prominent band around 100 kDa (Fig. 6). Testing reactivity of F95 on citrullinated peptides and their native equivalents by ELISA techniques as described by van de Stadt et al., demonstrated that F95 reacted primarily with citrullinated peptides and not with the native peptides (data not shown) (van de Stadt et al. 2011).

Discussion

To the best of our knowledge, this was the first study on the presence and location (stromal versus epithelial) of citrullinated proteins in periodontal tissue (Nesse et al. 2009) and is the first full article on this issue. Citrullinated protein presence was observed in 80% of periodontitis-affected stroma compared with 33% of control periodontal stroma (33%). In control periodontal stroma, citrullinated protein presence appeared to be associated with inflammation of non-periodontitis
origin. Recently, our results have been confirmed by others in an abstract, i.e. showing citrullinated protein presence in 80% of peri-odontitis patients and 40% of healthy controls (Yucel-Lindberg et al. 2010). Thus, citrullinated protein formation within the periodontal stroma appears to be an inflammation-depended process. This is again in agreement with previous studies showing that the formation of citrullinated proteins in other sites of the human body is inflammation-driven (Makrygiannakis et al. 2006).

In contrast with stroma, periodontal epithelium, whether inflamed or not, always stained positive for Ab5612. Citrullinated protein detection in periodontal epithelium samples could have been expected. In fact, the discovery of ACPA in 1964, then named anti-perinuclear factor for perinuclear staining pattern, was made on buccal epithelial (i.e. mucosal) cells (Nienhuis & Mandema 1964). Filaggrin, a keratin binding protein abundantly present in epithelium, is known to bind ACPA (Schellekens et al. 1998). Citrullinated proteins have also been shown to be present in the epidermis (Tsuji et al. 2003). It has even been posed that citrullination plays an essential role in epithelial differentiation through integration and disintegration of keratin filaments. In other words, citrullinated proteins naturally occur in epithelial tissues. However, periodontitis leads to the formation of additional types of citrullinated proteins (targeted by F95) in addition to the types of citrullinated proteins naturally present in non-inflamed periodontal epithelium (targeted by Ab5612 only).

Regarding the different type of citrullinated proteins present in periodontitis tissue compared with healthy periodontal tissue, it is striking that a variety of similar bands of citrullinated proteins (targeted by F95) were found in both synovium of RA and periodontitis tissue, with a strong band around 100 kDa. This may be due to the fact that the periodontium contains many of the cellular and structural components of a joint. The periodontium contain collagens, proteoglycans and hyaluronic acid, major constituents of cartilage (Bartold & Narayanan 2006). Many of the major auto-antigens targeted by ACPA in RA can be found within the periodontium. The periodontal epithelium contains keratin and filaggrin, and periodontal fibroblasts express vimentin. Citrullinated α-enolase is considered to be another major autoantigen in RA with which P. gingivalis produced citrullinated α-enolase shares, a largely identical immunodominant epitope (Lundberg et al. 2008, 2010, Wegner et al. 2010). The similar bands of citrullinated proteins in periodontitis and RA-affected synovial tissue may point out that similar proteins might get citrullinated in periodontitis and RA-affected synovial tissue. When these identical proteins are degraded by the activity of matrix metalloproteases and subsequently phagocytized by macrophages, as is the case in both RA and periodontitis, an auto-antibody response may be initiated in the periodontium targeting the same components in joints.

In this respect, another striking finding is the fact that the number of macrophages in stroma of periodontitis tissue samples without detectable citrullinated proteins (n = 3) is consistently zero, whereas their numbers are small, but almost consistently higher in stroma of periodontitis tissue containing citrullinated proteins. The presence of macrophages thus appears to be associated with citrullinated proteins in periodontium of periodontitis patients, synovium of RA patients and healthy periodontal tissue using mouse anti-citrullinated protein monoclonal antibody F95, magnification 100x. No false positive staining for citrullinated proteins was shown using a mouse IgM isotype control followed by secondary antibody staining in synovium of RA patient (a) and periodontium of periodontitis patient (c). Brown staining indicates the presence of citrullinated proteins using mouse anti-citrulline antibody (F95) followed by secondary antibody staining in synovium of RA patient (b) and periodontium of periodontitis patient (d). Healthy periodontal tissue stained negative for citrulline using F95 (e) or positive in the case of presence of inflammation (f).
nated protein formation in periodontitis stroma, and macrophages may play a role in initiation of ACPA formation as antigen presenting cells. Gemmel et al. (2003) described that a predominance of B cells was found in moderate and extensive infiltrates of periodontal lesions, which can be regarded as a measure for inflammation. In our study the mean number of B cells was also increased in citrulline positive cells, however not significantly raised, probably due to small numbers.

No significant differences were found with regards to any of the other investigated parameters between citrullinated protein positive and negative periodontitis stroma (i.e. age, severity of periodontitis, gender, and smoking). However this may be due to the small number of patients.

The question might rise why the buccal mucosa cells we used as a positive control in our study stained positive for antiperinuclear factor (Brouwer et al. 2006, Gyorgy et al. 2006). Thus, the presence of citrullinated proteins targeted by Ab5612 and F95 in buccal mucosal cells of healthy donors is most likely physiological and unrelated to ACPA associated aetiology of RA. Although this study is the first to show citrullinated protein presence in periodontitis tissue, ACPA presence and local ACPA formation in periodontitis tissue and peripheral blood were not assessed. Neither was P. gingivalis presence assessed. It is not possible to stain for PPAD, as no antibodies are available to this antigen and staining for human PAD4 showed positive neutrophils in the tissue (data not shown). Thus, the most important focus of future studies should be on potential ACPA formation within periodontitis tissue. We are currently investigating presence of ACPA in serum of patients with severe periodontitis. Other factors related to ACPA formation in periodontitis should also be assessed in studies with a larger sample size, i.e. smoking habits, the types of citrullinated proteins present as well as the specific role of P. gingivalis, the presence of HLA DR B1 shared epitope alleles and the role of auto-antibody producing plasma cells and collagen auto-immunity.

In conclusion, citrullinated protein formation within the periodontal stroma appears to be an inflammation-dependent process. In periodontal epithelium, inflammation may lead to the formation of different types of citrullinated proteins.


© 2012 John Wiley & Sons A/S


**Clinical Relevance**

**Scientific rationale for the study:** Anti-citrullinated protein antibodies (ACPAs) may play a role in the aetiology of rheumatoid arthritis (RA). Periodontitis contains all of the ingredients necessary to induce ACPA formation. However, whether citrullinated proteins are present in periodontitis tissue currently remains unknown.

**Principal findings:** Periodontitis leads to the formation of additional types of citrullinated proteins which appear to be similar to those formed in RA-affected synovial tissue.

**Practical implications:** In the presence of a certain genetic predisposition and smoking, citrullinated proteins formed in association with periodontitis may play a role in ACPA formation. Periodontitis might contribute to RA development.