Cytokines modulate routes of collagen breakdown: review with special emphasis on mechanisms of collagen degradation in the periodontium and the burst hypothesis of periodontal disease progression
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Review with special emphasis on mechanisms of collagen degradation in the periodontium and the burst hypothesis of periodontal disease progression


Abstract. In this paper, we review recent work on collagen degradation. 2 main routes of breakdown are described and their relevance during healthy and inflammatory conditions of the periodontium is discussed. Special attention is paid to the possible rôle of cytokines, in particular interleukin 1 (IL-1) and transforming growth factor β (TGF-β), on the modulation of collagen phagocytosis and metalloproteinase production. IL-1 has been shown to have a dual function in collagen digestion. It inhibits the intracellular phagocytic pathway, but at the same time, it strongly promotes extracellular digestion by inducing the release of collagenolytic enzymes like collagenase. TGF-β has an opposite effect on both pathways and antagonizes IL-1. Collagenase is released in an inactive form, and a considerable fraction of the proenzyme may become incorporated in the extracellular matrix. This reservoir of latent enzyme can be activated (for instance by plasmin), leading to a sudden and extensive breakdown of the collagenous fibre meshwork. It is suggested that this phenomenon may also take place during progressive periodontitis and could explain an episodic nature of collagenolysis, clinically resulting in bursts of attachment loss (burst hypothesis).

The Intracellular Route

Under non-pathological conditions phagocytosis and intracellular digestion of collagen fibrils (Fig. 2) is a process observed at a high level in dynamic soft connective tissues such as gingiva and periodontal ligament (Ten Cate & Freeman 1974, Beertsen & Everts 1977, Beertsen et al. 1978, Sodek & Overall 1988). Of all connective tissues studied so far it appears that the periodontal ligament has the fastest turnover of collagenous proteins. In rodents the half-life of collagen in the ligament is in order of 1–5 days, which is considerably faster than that in skin (Sodek 1977). This rapid renewal is considered to be important to allow positional adaptation of teeth during function (Sodek, 1989). It is intriguing how the periodontal ligament is able to provide firm attachment, while at the same time allowing the tooth to undergo processes like eruption and spatial adjustments in the dental arch. Obviously, the metabolism of collagens in the periodontal ligament has to be an accurately controlled balance between synthesis and degradation (Beertsen & Everts 1977, Beertsen et al. 1978).

The breakdown of collagenous proteins occurs via two different pathways: an intracellular and an extracellular route (Fig. 1: Everts et al. (1989), Everts & Beertsen (1992), Murphy & Reynolds (1993), Birkedal-Hansen (1993a), Birkedal-Hansen et al. (1993)).
The two major routes of collagen breakdown in soft connective tissue: the intracellular and extracellular pathways. (A) The intracellular pathway primarily occurring during normal turnover and remodeling. (A1) Cytoplasmic protrusions surround a collagen fibril, thereby segregating it from the rest of the extracellular matrix and forming a phagosome. (A2) Subsequently, after fusion of a lysosomal vacuole with the phagosome, the proteolytic lysosomal enzymes, such as cysteine proteinases, degrade the fibril intracellularly. (B) The extracellular pathway occurring during excessive breakdown of collagen, e.g., during inflammation or uterus involution. (B1) Following production, procollagenase is released in the extracellular environment where it may be activated (e.g., by plasmin). (B2) Activated collagenase degrades collagenous proteins (in combination with other proteinases such as gelatinase) or it may be inhibited by a tissue inhibitor of metalloproteinases (TIMP). Both pathways are likely to be modulated by cytokines like IL-1α and TGF-β.

Éverts et al. (1989). In spite of its putative physiological importance (Éverts & Beertsen 1992), little is known about the mechanisms involved in the regulation of the phagocytic pathway. Recently, however, evidence was obtained that cytokines may modulate the intracellular pathway of collagen breakdown (van der Zee et al. 1995a). While TGF-β enhanced collagen phagocytosis, IL-1α inhibited the process. In combination these cytokines proved to antagonize each other (Table 1).

### The Extracellular Route

In pathological situations, such as during periodontal disease, the delicate balance between synthesis and degradation is disturbed. Microbial products may trigger a host response which induces production and release of cytokines and proteolytic enzymes by both inflammatory and resident cells (Page 1991, Genco 1992, Birkedal-Hansen 1993a). Matrix metalloproteinase-1 (MMP-1), or interstitial collagenase, is the best known representative of this group of enzymes. It has the unique capacity to cleave the interstitial collagens types I and III at a single locus of the triple helical body of the collagen molecule which is relatively resistant to other proteinases (Harris & Cartwright 1977) thus producing characteristic N-terminal 3/4 and C-terminal 1/4 fragments.

The action of collagenase is controlled at least at three distinct levels involving production, activation and inhibition (Fig. 3). First, the enzyme is synthesized and secreted in an inactive proform. Second, the enzymes are activated for instance by autoactivation, by plasmin, by activators such as gelatinase or by other proteinases (Harris & Cartwright 1977) and catabolized by other proteinases such as TIMP.

### Table 1. Effect of cytokines on collagenase release and collagen phagocytosis by rabbit periosteal explants (derived from van der Zee et al., 1994, 1995a).

<table>
<thead>
<tr>
<th></th>
<th>EGF</th>
<th>IL-1α</th>
<th>TGF-β</th>
<th>IL-1α + EGF</th>
<th>IL-1α + TGF-β</th>
</tr>
</thead>
<tbody>
<tr>
<td>Collagenase release</td>
<td>no effect</td>
<td>stimulation</td>
<td>inhibition</td>
<td>synergistic increase</td>
<td>antagonism</td>
</tr>
<tr>
<td>Phagocytosis</td>
<td>no effect</td>
<td>inhibition</td>
<td>stimulation</td>
<td>(strong) inhibition</td>
<td>antagonism</td>
</tr>
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</table>
Cytokines, collagenolysis and the 'burst hypothesis'

Inhibitors of collagenolytic activity

Fig. 3. Possible mechanism resulting in a burst of periodontal breakdown. Under the influence of cytokines (e.g., IL-1) a high level of procollagenase is produced and stored in the extracellular matrix (ECM) of the periodontal soft connective tissues. Activation of this enzyme reservoir (e.g., by plasmin) results in a high level of active enzyme that causes a rapid breakdown of collagen fibres, clinically resulting in a burst of loss of attachment. Tissue inhibitors of metalloproteinases (TIMPs) that may also be induced by cytokines (e.g., TGF-β) inhibit the collagenolytic activity and stop further progression of the disease until a newly build-up enzymatic reservoir becomes activated and a new burst occurs.

Is there a Relationship between the Intra- and Extracellular Pathways of Collagen Degradation?

Although the relationship between the intracellular and extracellular pathways of collagen breakdown is presently unknown, it is generally taken that, prior to phagocytosis, partial extracellular proteolysis has to occur by collagenase or other MMPs (Harris & Krane 1974, Murphy & Reynolds 1985, Laurent 1987). One may wonder, however, whether collagenase is indeed involved in this process. In this respect it is interesting to note that it has not been possible to demonstrate the presence of collagenase in healthy periodontal ligament, being a tissue with one of the highest physiological collagen degradation rates (Overall et al. 1987). Neither was it possible to influence the level of collagen phagocytosis by blocking collagenase activity with anti-collagenase antibodies or TIMP in cultured periodontal tissue (Everts et al. 1989).

Recently, even a negative correlation was found under the influence of cytokines between the production of procollagenase (collagenase-mediated extracellular pathway) and the level of collagen phagocytosis (intracellular pathway) in rabbit periostea ex vivo (Table 1). This may be a first indication of a relationship between the two collagenolytic pathways and it may emphasize a role of cytokines in balancing the relative contribution of each route (see also Everts et al. 1996).

Table 2. List of abbreviations

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Description</th>
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<tbody>
<tr>
<td>EGF</td>
<td>epidermal growth factor</td>
</tr>
<tr>
<td>FGF</td>
<td>fibroblast growth factor</td>
</tr>
<tr>
<td>IL-1</td>
<td>interleukin 1</td>
</tr>
<tr>
<td>MMP</td>
<td>matrix metalloproteinase</td>
</tr>
<tr>
<td>PDGF</td>
<td>platelet-derived growth factor</td>
</tr>
<tr>
<td>TGF</td>
<td>transforming growth factor</td>
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<tr>
<td>TIMP</td>
<td>tissue inhibitor of metalloproteinases</td>
</tr>
<tr>
<td>TNF</td>
<td>tumor necrosis factor</td>
</tr>
</tbody>
</table>

Table 2

Collagenase and Periodontal Disease

Collagenase that is found in case of periodontitis is derived from the host and not from periodontal bacteria (Krystaltaskyj & Sodek 1987). The presence of two distinct types of collagenases has been described: the 57/52 kD (MMP-1) and the 75 kD form (MMP-8). The former enzyme is produced and released as an inactive proenzyme by many periodontal cell types including fibroblasts but also macrophages, endothelial cells, epithelial cells, Langerhans cells, and osteoblasts (Birkedal-Hansen 1993a). This form is also known as interstitial collagenase or 'fibroblast-type' collagenase. MMP-8 is derived from neutrophils where it is stored intracellularly in granules ('neutrophil-collagenase') (Sorsa et al. 1992).

Although there is no direct evidence for a causal relationship between metalloproteinases and periodontal tissue destruction, the involvement of collagenase in collagen degradation during chronic inflammatory periodontal disease is highly suggestive and based on a vast amount of studies. MMP-8 is detected at high levels in gingival crevicular fluid and saliva during gingivitis or periodontitis whereas it is undetectable in healthy individuals (Sorsa et al. 1992, Ingman et al. 1993). Moreover, a positive correlation has been shown between the amount of this enzyme and the level of disease activity (Krystaltaskyj et al. 1986) and a negative correlation between the enzyme level and the intensity of periodontal treatment (Golub et al. 1976, Kowashi et al., 1979 Lamster et al. 1985, Larivée et al. 1986, Villela et al. 1987, Hakkarainen et al. 1988). Furthermore, in extracts or homogenates of diseased periodontal tissues MMP-1 is abundantly present in contrast to healthy specimens and, as with MMP-8, a positive correlation was found between the presence of the enzyme and the severity of inflammation (Overall et al. 1987, Sorsa et al. 1988, Robinson et al. 1992). In addition, MMP-1 has been immunolocalized in inflamed but not in healthy periodontal tissue (Woolley & Davies 1981). More recently it was shown that mRNA expression for collagenase and TIMP is enhanced in diseased gingiva (Nomura et al. 1993, Tonetti et al. 1993). Following initial periodontal treatment however, Meikle et al. (1994) could not find detectable levels of MMP-1 in fresh gingival biopsies.

The relative contribution of MMP-8 and MMP-1 in the pathogenesis of periodontitis is still not entirely clear and needs further investigation. It has been shown that most if not all collagenase...
in gingival crevicular fluid or saliva of patients with adult periodontitis is derived from neutrophils (MMP-8), without detectable levels of MMP-1 (Sorsa et al. 1988, 1992, Sodek & Overall 1992). On the other hand, immunolocalization of MMP-1 in inflamed periodontal tissues showed a strong labeling and association between the presence of the enzyme and resident connective tissue cells (Woolley & Davies 1981). These data may suggest that, following production during inflammation, most MMP-1 remains in the gingival tissue, whereas the vast majority of released MMP-8 finds its way to the pocket.

Cytokines Mediate Collagenase Production

During the last 2 decades numerous biological effectors have been elucidated which participate in the regulation/modulation of MMP-mediated collagen degradation. As mentioned above, cytokines, being polypeptides which mediate cell metabolic processes, appear to play a crucial role in this process. Also periodontitis is strongly associated with the presence of cytokines, such as IL-1α and IL-1β and indeed many cell types in the periodontium have the capacity to produce these compounds (Mundy 1991, Birkedal-Hansen 1993b). In inflamed gingiva, however, macrophages were found to be the prime IL-1 mRNA expressing cells (Matsu ki et al. 1993). Although the rate of IL-1α and IL-1β expression was similar, in gingival crevicular fluid IL-1α was the predominant active cytokine (Matsu ki et al. 1993). A positive correlation was found between the presence and activity of periodontal disease and the level of this cytokine in tissue extracts (Honig et al. 1989, Jandinski et al. 1991, Stash enko et al. 1991, Matsuki et al. 1993) or gingival crevicular fluid (Charon et al. 1982).

The role of cytokines in the pathogenesis of periodontal disease is probably rather complex. There is ample in vitro evidence, however, suggesting that these compounds are involved in the regulation of metalloproteinase production and activity which eventually results in loss of periodontal attachment. First, it was shown that fibroblasts of periodontal tissues produce MMP-1 as well as inhibitor molecules in vitro (Pet tigrew et al. 1980, Heath et al. 1982) their production being mediated by cytokines. In particular IL-1 proved to be extremely potent in inducing proMMP-1 production by many cell types, including fibroblasts of the gingiva and periodontal ligament (Lark et al. 1990, Richards & Rutherford 1988). On the other hand, TGF-β, a cytokine known for its wound healing and repair-stimulating activities (Roberts & Sporn 1993), has a downregulating effect on MMP-1 expression, synthesis, and release, and moreover appears to neutralize the activity by stimulating the production of TIMP (Overall et al. 1989).

Studies on the effects of combinations of various cytokines are of interest, since they may act in a synergistic or antagonistic fashion (Lynch et al. 1987, Chandrasekhar & Harvey 1988, Andrews et al. 1989, MacNaul et al. 1990, Circolo et al. 1991, Tingström et al. 1992). Recently, it was demonstrated that IL-1α in combination with EGF synergistically enhances the production of procollagenase by periodontal tissue explants up to 100-fold (Van der Zee et al. 1993). TGF-β, on the other hand, induces an inhibition of procollagenase release (Overall et al. 1989, Van der Zee et al. 1995a). When IL-1α and TGF-β are added in combination an antagonistic effect is observed (Chandrasekhar & Harvey 1988, Andrews et al. 1989, Van der Zee et al. 1995a, b). Members of the TGF-β superfamily (TGF-βs and bone morphogenetic proteins) appear to be essentially anabolic growth factors promoting deposition of stroma (Roberts et al. 1986, Overall et al. 1989, Roberts & Sporn 1993) whereas cytokines like IL-1 and TNFα are generally proinflammatory mediators inducing transcription of MMP genes which eventually results in resorption (Dayer et al. 1985, McCachren et al. 1989, Saito et al. 1990, Tatakis 1993).

Another process mediated by cytokines like IL-1α, EGF and TGF-β is the resorption of bone (Raisz et al. 1980, Gowen et al. 1983, Gowen & Mundy 1986, Lorenzo et al. 1988, Pleilschifer et al. 1988, Tatakis 1993). Although bone resorption should probably be considered as a distinct degradative pathway with a central role played by the osteoclast (Birkedal-Hansen, 1993a), collagenase also appears to be involved at least in some steps of this process like breakdown of the collagen fringe prior to and following osteoclastic bone resorption (Everts et al. 1994) as well as in the actual osteoclast mediated bone resorption (Delaisse et al. 1993, Hill et al. 1994, 1995). Lorenzo et al. (1988) demonstrated that bone resorption of rat long bone cultures is enhanced by EGF or IL-1α and that a combination of both cytokines resulted in additional effects. Recently, a comparable effect of cytokines in rabbit calvariae was found together with a concomitant increase in the release of the metalloproteinases, collagenase and gelatinase, and tissue inhibitor of metalloproteinases (TIMP), suggesting that MMP-mediated steps in bone resorption are also modulated by cytokines (Van der Zee et al. in preparation).

Cytokine-induced Reservoir of Latent Collagenase in Extracellular Matrix: a Possible Mechanism for a "Burst" of Periodontal Breakdown

Several studies have suggested that periodontitis has a cyclic behavior being characterized by relatively short periods of exacerbation during which progressive loss of attachment occurs followed by periods of remission (Goodson et al. 1982, Scransky et al. 1984, Goodson 1992, Cohen 1993, Machtet et al. 1993a, b). Understanding of the pathogenesis of periodontitis is important for an optimal treatment and prevention of the disease. Therefore, it is of considerable interest to know when, how and why progression of periodontitis occurs. Periods of progressive loss are likely to involve significant proteolysis during a relatively short time-interval. Such a proteolytic burst can be explained by assuming that a sudden increase occurs in the synthesis and/or release of catabolic enzymes (e.g. MMP-1/MMP-8). Alternatively, a similar excess proteolytic activity could occur if these proteolytic enzymes are stored at high concentrations in the extracellular matrix and suddenly become activated. In support of this latter mechanism are recent in vitro data demonstrating that a vast amount of procollagenase can be incorporated into the extracellular matrix of cultured soft connective tissue (periosteum) without resulting in degradation (Van der Zee et al. 1994). Activation of this matrix-stored enzyme fraction by plasmin (considered to be a putative physiological activator), resulted in a rapid breakdown of the bulk of the collagenous proteins present (70%) (Van der Zee et al. 1996). The activation process may be regarded as a rate-limiting step.
an explanation for the episodic nature of collagenolysis resulting in bursts of attachment loss ('burst hypothesis'). Other cytokines, like TGF-β, may counterbalance these effects during phases of remission or healing, and contribute to restoration of a state of equilibrium.

**Zusammenfassung**

Zytokine beeinflussen die Wege des Kollagenabbau. Übersichtsartikel mit besonderer Betonung der Mechanismen des Kollagenabbaus im Parodont und der Aushbruch-Hypothesse bei progredienter Parodontalkrankheit


**Résumé**

Voies modulées des cytokines de la destruction du collagène. Revue insistant sur les mécanismes de la dégradation du collagène dans le parodonte et sur l'hypothèse de flambées de progression dans la maladie parodontale

 Sous les conditions physiologiques, en état normal, le collagène est détruit principalement via la voie intracellulaire. Sous des conditions pathologiques (par exemple l'inflammation) des cytokines comme l'interleukine-1 sont libérées et peuvent d'une part induire la production et la libération d'enzymes collagénolytiques et d'autre part inhiber la phagocytose du collagène. Suite à la libération de procollagènes une importante fraction du proenzyme est incorporée dans la matrice extra-cellulaire. Sous les conditions inflammatoires un tel réservoir d'enzymes latents pourrait être activé, entraînant une destruction rapide et importante du collagène. Si ce phénomène prend place durant la parodontite et il peut en partie du moins apporter une explication à la nature épisodique de la collagènesé produisant par flammées de perte d'attache. D'autres cytokines tel le facteur transformant la croissance-β, peuvent contrebalance ces effets durant les phases de remission ou de guérison, et contribuent à la restauration d'un état d'équilibre.

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