Quebecol Shows Potential to Alleviate Periodontal Tissue Damage and Promote Bone Formation in In Vitro Models

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**ABSTRACT:** Quebecol is a polyphenolic compound initially isolated from Canadian maple syrup in 2011. Recently, our group demonstrated in a macrophage model that quebecol inhibits the secretion of pro-inflammatory cytokines and reduces the activation of the NF-kB transcription factor. In this study, we further explored the therapeutic potential of quebecol against periodontal disease, an inflammatory disorder of bacterial origin affecting tooth-supporting tissues. More specifically, the effects of this natural compound on matrix metalloproteinase (MMP) activity and macrophage secretion, as well as on the mineralization activity of osteoblasts (bone-forming cells), were investigated. Results showed that exposing lipopolysaccharide (LPS)-treated macrophages to quebecol led to a significant decrease in the secretion of MMP-8 and MMP-9. In addition, quebecol dose dependently inhibited the catalytic activity of MMP-9. Quebecol also enhanced the mineralization activity of osteoblasts. This study brought forward additional evidence to support the potential of quebecol as a nutraceutical agent against periodontitis.

**INTRODUCTION**

Periodontitis, a destructive form of periodontal disease that causes permanent damage to the periodontium, is a widespread pathology that currently ranks as the sixth most common chronic inflammatory disorder in the world.1 As such, it remains to this day a worldwide public health concern with major economic and social burdens. Periodontitis results from an uncontrolled inflammatory response of mucosal and immune cells to specific Gram-negative strictly anaerobic bacterial species colonizing the subgingival sites as a biofilm.2,3 Evolution of this pathology leads to the progressive destruction of periodontal connective tissue and alveolar bone that may cause tooth loss.4 This pathogenic process involves the expression and secretion by resident and infiltrating cells of the periodontium of a wide variety of pro-inflammatory mediators and matrix metalloproteinases (MMPs) that modulate periodontal tissue destruction and perpetuate the pathology.5−6

MMPs, a family of host-derived calcium-dependent zinc-containing proteolytic enzymes, are involved in the degradation of extracellular matrix (ECM) components, such as collagen, fibronectin, laminin, and elastin, during both normal and pathological conditions.7 MMPs are produced by the major cell types found in periodontal tissue, including epithelial cells, fibroblasts, neutrophils, and macrophages.5,8 In healthy periodontal tissues, their expression is quite low and their activity is regulated by endogenous molecular agents called tissue inhibitors of metalloproteinases (TIMPs).5,6 However, overexpression of MMPs has been associated with the pathological tissue destruction process observed during periodontitis.5,6,8,9 Therefore, the excessive production and activity of MMPs has been proposed as a potential target for pharmacological intervention in periodontitis.5,10,11

MMPs also markedly contribute to the alveolar bone loss occurring during periodontitis, which is a process triggered by osteoclasts and attributed to the inflammatory nature of the disease.12 Osteoblasts, which are cells involved in the process of bone formation, can restore normal bone homeostasis.13,14 These cells are derived from mesenchymal cells and secrete the organic matrix (mostly collagen) and induce calcification during the formation of new bones.13,14 Targeting osteoblast activity may therefore represent a relevant therapeutic strategy against periodontitis.

Quebecol (ethyl 2,3,3-tris(4-hydroxy-3-methoxyphenyl)-propan-1-ol; compound 1; Figure 1a) is a polyphenolic compound initially isolated from Canadian maple syrup in 2011.15 It is formed during the heating process required to transform the sap of Acer saccharum into syrup. In vitro bioassays brought evidence that quebecol may represent a promising chemopreventive or chemotherapeutic agent.16−18

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Recently, we developed an efficient synthesis protocol for quebecol (Figure 1a)\textsuperscript{19} and reported its in vitro anti-inflammatory activity.\textsuperscript{20} More precisely, we showed that treatment of lipopolysaccharide (LPS)-stimulated human macrophages with quebecol leads to an inhibition of the secretion of two pro-inflammatory cytokines (IL-6 and TNF-\(\alpha\)), while a reduction of the activation of the NF-\(\kappa\)B transcription factor was also observed.\textsuperscript{20} In parallel to our work on quebecol itself,\textsuperscript{20} related compounds (precursors, isomers, and substructures) were synthesized and used in structure–activity relationship studies.\textsuperscript{20,21} In brief, these studies showed that quebecol (Figure 1a) and its diarylmethane substructure (compound 3; Figure 1b) exhibit anti-inflammatory activity.\textsuperscript{20,21} Results also demonstrated that the substitution pattern on the aromatic rings of these active structures (compounds 4 and 5; Figure 1b) led to a slightly enhanced activity but higher cytotoxicity.\textsuperscript{21} It was concluded that the natural product 1 was still the most promising anti-inflammatory compound based on biological activity and toxicity arguments.

Encouraged by the above results, we wished to further explore the therapeutic potential of quebecol against periodontitis by focusing on the two specific therapeutic targets discussed above: MMPs and osteoblasts. First, we investigated the ability of quebecol to attenuate the secretion of MMPs by LPS-stimulated macrophages and to inhibit the catalytic activity of MMP-9. Second, we assessed the effects of quebecol on the mineralization activity of osteoblasts.

\section*{RESULTS}

\subsection*{Quebecol and MMP Secretion by Macrophages.}

The effect of quebecol on the in vitro regulation of MMPs was investigated by assessing its capacity to impair their secretion in a macrophage model, which represents a leukocyte type recognized as an important producer of MMPs in the context of periodontitis. To this end, we assessed its effect on the basal and LPS-induced secretion of MMP-3, MMP-8, and MMP-9. These three specific MMPs were selected on the basis of their known association with active periodontitis.\textsuperscript{5,6,8–10} It is also worth mentioning that these three MMPs were those secreted in the highest amount by the U937 macrophages used in this study.

The secretion of MMP-3, MMP-8, and MMP-9 by macrophages kept under normal conditions or stimulated with \textit{Aggregatibacter actinomycetemcomitans} LPS is reported in Figure 2. MMP-9 was found to be produced in significantly greater amounts than MMP-3 and MMP-8. The stimulation of macrophages with LPS did not significantly modify the secretion of MMP-3 and MMP-9, while it increased the secretion of MMP-8 by 20.8\% (\(p<0.001\)). More specifically, LPS-stimulated macrophages secreted 650 ng/mL, 3.109 pg/mL, and 2.354 pg/mL of MMP-9, MMP-8, and MMP-3, respectively.

We first investigated the effect of quebecol on MMP secretion by macrophages kept under normal conditions (without LPS stimulation). When the cells were exposed to the highest concentration (500 \(\mu\)M) of quebecol, the secretion of MMP-3, MMP-8, and MMP-9 significantly decreased by 15.9, 34.4, and 26.8\% (\(p<0.001\)) (Figure 3). Quebecol at concentrations \(\leq\) 125 \(\mu\)M did not reduce the secretion of any of the MMPs in a significant manner.

Then, the same analysis was performed using macrophages that were stimulated with \textit{A. actinomycetemcomitans} LPS in a way to simulate an inflammatory condition. As shown in Figure 3, while only a slight inhibition of MMP-3 secretion by quebecol was observed for LPS-stimulated macrophages, significant dose-dependent reductions in the secretion of MMP-8 and MMP-9 were caused by the presence of quebecol even at the lowest concentration tested (31.25 \(\mu\)M), with
inhibition of 15.2 and 21.4%, respectively. Complete tabulated results can be found in the electronic Supporting Information.

Overall, a difference regarding the inhibitory activity of quebecol on the secretion of MMP-8, MMP-9 and to a lesser extent MMP-3 by macrophages kept under normal conditions or stimulated with *A. actinomycetemcomitans* LPS was observed. Such a selectivity would be advantageous regarding a potential clinical use since this natural compound would have a minimal impact on healthy gingival tissues.

**Quebecol and MMP-9 Activity.** Given the interesting results obtained regarding the effect of quebecol on MMP-9 secretion and that MMP-9 was produced in high amounts by macrophages, we selected this MMP to investigate the effect of quebecol on its catalytic activity.22 As reported in Table 1, quebecol reduced the degradation of gelatin by MMP-9 in a dose-dependent manner. More specifically, the presence of quebecol at 250 and 500 μM inhibited the activity of MMP-9 by 42 and 92.6%, respectively. It is interesting to note that while the secretion of this MMP by macrophages remained similar between these two concentrations of quebecol under an inflammatory context, an important difference with respect to MMP-9 activity was observed.

**Quebecol and Osteoblast Mineralization Activity.** An important process involving osteoblasts during bone formation is their mineralization activity. It was thus of interest to investigate the effect of quebecol on the mineralization activity of SaOS-2 cells by monitoring the formation of extracellular mineral deposits with a staining agent (Alizarin Red S) that selectively binds to calcium salts. As shown in Figure 4, the presence of quebecol promoted the formation of calcium deposits that appear as red mineralized nodules. More specifically, quebecol, from a concentration of 12.5 μg/mL (corresponding to 29.3 μM), dose dependently promoted the mineralization activity of osteoblasts (Figure 5). A marked increase in mineralization activity was observed at the highest concentrations tested. For instance, the presence of quebecol at 100 μg/mL (corresponding to 234 μM) increased the mineralization activity by 2.4-fold.

**DISCUSSION**

Polyphenols are compounds that possess an aromatic ring with at least one hydroxyl group and show high heterogeneity in their structure, from simple molecules to complex polymers.23 Due to their antimicrobial, antioxidant, and immunomodulatory properties, there has been growing research interest in polyphenols with respect to oral health benefits.24−26 A process-derived polyphenolic compound isolated from maple syrup, named quebecol after the Canadian province of Quebec (the world’s leading producer of maple-derived products), has been the topic of several studies.15−21 Based on our recent investigations reporting the anti-inflammatory properties of quebecol,20 the present study further investigated its potential

![Graph](https://example.com/graph.png)  
**Figure 3.** Inhibitory effect of quebecol on the secretion of MMP-3, MMP-8, and MMP-9 by macrophages stimulated with *A. actinomycetemcomitans* LPS or not. Mean ± standard deviations of triplicate assays. *p < 0.001, #p < 0.05.

| concentration (μM) | relative activity (%) |
|-------------------|----------------------|
| 0                 | 100 ± 2              |
| 62.5              | 92.2 ± 0.6           |
| 125               | 82 ± 1               |
| 250               | 58 ± 3               |
| 500               | 7.4 ± 0.6            |

*Significantly different from control (no quebecol) at p < 0.001.*
as a prophylactic and therapeutic agent against periodontitis, a widespread chronic inflammatory disease of bacterial origin that affects the tooth-supporting tissue.

The continuous, high production of MMPs by infiltrating and resident cells of the periodontium triggered by periodontopathogens contributes to connective tissue destruction and alveolar bone resorption.\(^6,8,9\) In fact, it is well known that MMP levels and activities are significantly higher in the gingival tissue and gingival crevicular fluid of periodontitis-affected individuals than in healthy subjects.\(^6,8,9\) Therefore, MMP inhibitors may have interesting potential for use as an effective therapeutic approach for treating periodontitis patients.\(^10,11\) Consequently, quebecol was tested for its effect on MMP secretion in a macrophage model, as macrophages are potent MMP producers and constitute an important part of the inflammatory infiltrate in active periodontal lesions.\(^27,28\) In the present study, we showed that \textit{A. actinomycetemcomitans} LPS acts as an inducer of the secretion of MMP-8 by macrophages, while it has no effect on the secretion of MMP-3 and MMP-9. Our results showed that quebecol caused a significant inhibition of the secretion of MMP-8 (50.9\%) and MMP-9 (34.8\%) when the cells were treated with this phenolic compound (250 \(\mu\)M) in an inflammatory context (stimulation with LPS). MMP-8 (collagenase-2) and MMP-9 (gelatinase B) are present in high concentrations in inflamed periodontitis sites and are strongly associated with the progression of periodontitis.\(^29\) Periodontal therapy (scaling and root planing) has been shown to significantly reduce salivary MMP-8 and MMP-9 levels,\(^30,31\) suggesting that these two MMPs are potential markers for monitoring periodontal disease activity. Consequently, the ability of quebecol to attenuate the secretion of these two MMPs may contribute to reduced periodontal tissue destruction. Although the exact inhibitory mechanism of action of quebecol was not investigated, it is likely related to the previously reported capacity of quebecol to impair NF-\(\kappa\)B activation in macrophages.\(^20\) Indeed, previous studies have shown that the expression of MMPs by macrophages involves the NF-\(\kappa\)B signaling pathway.\(^32,33\) Moreover, La et al.\(^34\) reported that cranberry proanthocyanidins reduce MMP-9 secretion in macrophages through the inhibition of NF-\(\kappa\)B activation.

To complement the results of the impact of quebecol on MMP secretion by macrophages, we assessed its effect on MMP-9 activity. This analysis revealed that the polyphenolic compound exerts a marked dose-dependent inhibitory effect on the capacity of MMP-9 to degrade its natural substrate (gelatin). Although the mechanism of action remains to be elucidated, it can be hypothesized that quebecol, with its four phenol groups, acts as a ligand on MMP-9 that either deactivates the zinc ion on the catalytic cycle or blocks the access of the substrate to the active site of the enzyme.

Since bone formation is instrumental to counterbalance tissue destruction in the context of periodontitis, we also studied the effect of quebecol in an \textit{in vitro} osteoblast model. While they secrete collagen and proteins during bone formation, osteoblasts also trigger calcium and mineral deposition, leading to the formation of hydroxyapatite on the newly formed bone matrix.\(^13,14\) We demonstrated that the mineralization activity of osteoblasts was positively affected by quebecol in a dose-dependent fashion, the most important increase in the formation of extracellular calcium deposits being observed at the highest concentrations tested. It is also noteworthy to mention that results obtained in a parallel study led by our group strongly suggest that the enhanced mineralization activity caused by quebecol is not due to an effect of this compound on the population of osteoblast cells (see the Supporting Information). In accordance with our results, polyphenols from green tea are also known to promote osteoblastic mineralization.\(^35\)
In conclusion, when taken together, the results of our present and previous studies point toward the potential use of quebecol as a promising multitarget therapeutic agent against periodontitis and, most likely, other inflammatory disorders. Indeed, at nontoxic concentrations, quebecol was found to inhibit the secretion of MMP-8 and MMP-9 in a macrophage model (stimulated with LPS), to inhibit the catalytic activity of MMP-9, and to stimulate osteoblast mineralization activity, in addition to impairing NF-κB activation and IL-6 secretion in macrophages. In this regard, studies on the effects provided by the use of oral hygiene products (mouth rinses and chewing gums) or slow mineralization activity, in addition to impairing NF-κB catalytic activity of MMP-9, and to stimulate osteoblast differentiation into osteoblasts, which represent the second contributing factor to periodontitis. Studies on the effect of quebecol on alveolar bone osteoclasts are also underway.

### METHODS

**Synthesis of Quebecol.** Quebecol was synthesized and characterized according to a previously reported protocol.

**Preparation of Monocyte-Derived Macrophages.** U937 human monocytes (ATCC CRL-1593.2) were purchased from the American Type Culture Collection (Manassas, VA) and were cultivated under the following conditions: RPMI-1640 + 10% heat-inactivated fetal bovine serum (FBS) + 100 μg/mL of penicillin G/streptomycin, 37 °C, and 5% CO₂ atmosphere. In a way to induce their differentiation into adherent macrophage-like cells, monocytes (2.5 × 10⁵ cells/mL) were incubated for 48 h in RPMI-10% FBS medium containing 100 ng/mL of phorbol myristic acid (PMA, Sigma-Aldrich Canada Ltd., Oakville, ON, Canada) and then harvested by scraping. After centrifugation (1200g for 5 min), the adherent macrophage-like cells were washed, suspended in RPMI-1% FBS (concentration of 1 × 10⁶ cells/mL), and seeded into wells of 12-well microplates (1 × 10⁶ cells/well), which were incubated for 24 h at 37 °C in a 5% CO₂ atmosphere.

**Determination of MMP-3, MMP-8, and MMP-9 Secretion by Macrophages.** The aforementioned macrophages were treated with twofold serial dilutions of quebecol (500–31.25 μM in culture medium) for 2 h prior to stimulation with LPS (1 μg/mL) prepared from A. actinomycescomitans ATCC 29522 according to a previously described protocol. A series of macrophages treated with quebecol but not stimulated with LPS was also prepared. After a 24 h incubation (37 °C, 5% CO₂ atmosphere), the culture supernatants were collected and stored at −20 °C until analyzed. Cells incubated in a culture medium without quebecol and/or LPS were used as negative controls. BAY-11-7082 (Selleckchem, Houston, TX) at a concentration of 5 μg/mL was used as a positive inhibitory control. Quantification of MMP-3, MMP-8, and MMP-9 in cell culture supernatants was performed with enzyme-linked immunosorbent assay (ELISA) kits (R & D Systems, Minneapolis, MN) according to the manufacturer's protocols. Absorbance was recorded at 450 nm using a microplate reader (Bio-Rad Laboratories, Mississauga, ON, Canada) with a wavelength correction set at 550 nm. Sensitivities of the commercial ELISA kits were 0.156, 0.058, and 0.313 ng/mL for MMP-3, MMP-8, and MMP-9, respectively. The concentration of each MMP (active + latent forms) was determined in triplicate assays from two independent experiments. **Determination of MMP-9 Activity.** Human recombinant MMP-9 was purchased from Calbiochem (San Diego, CA) and prepared at a concentration of 20 μg/mL in 5 mM Tris-HCl (pH 7.5), with 10% glycerol, 0.1 mM CaCl₂, and 0.005% Brij 35. Activation of MMP-9 was performed in 50 mM Tris-HCl (pH 7.5) with 0.5 mM p-aminophenylmercuric acetate for 1 h at 37 °C. Activated MMP-9 was diluted to a final concentration of 10 μg/mL in the reaction buffer (50 mM Tris-HCl, 5 mM CaCl₂, 200 mM NaCl, and 0.2% Brij 35) and was treated with twofold serial dilutions of quebecol (500–62.5 μM) or without (control). After 15 min of incubation at 25 °C, the fluorogenic substrate fluorescein isothiocyanate-labeled DQ gelatin (Molecular Probes, Eugene, OR) was added at a final concentration of 50 μg/mL. The assay mixtures were incubated in the dark at 37 °C for 2 h before the fluorescence was measured with a Synergy 2 microplate reader (BioTek Instruments, Winooski, VT), using an excitation wavelength of 490 nm and an emission wavelength of 520 nm. Results were reported as % of activity (mean ± standard deviations of triplicate assays from two independent experiments) from a comparison with the control assay without quebecol (set as 100%). **Mineralization Activity of Osteoblasts.** Human osteogenic sarcoma SaOS-2 cells, ATCC HTB-85, were purchased from ATCC and routinely cultured at 37 °C in an atmosphere of 5% CO₂ in McCoy’s 5A medium containing 2 mM L-glutamine and supplemented with 15% heat-inactivated FBS and 100 μg/mL of penicillin G-streptomycin. For the mineralization assay, SaOS-2 cells were cultivated in the above culture medium supplemented with the mineralization activation cocktail (MAC) composed of 10 mM β-glycerophosphate and 0.2 mM ascorbic acid. The cells were seeded at 1 × 10⁵ cells/well in a 96-well microplate and incubated for 24 h at 37 °C in an atmosphere of 5% CO₂. The medium was replaced with a fresh medium containing MAC and twofold serial dilutions of quebecol (100–1,5625 μg/mL). After 5 days of incubation, cells in the wells were fixed with 4% paraformaldehyde and stained with Alizarin Red S (Sciencell Research Laboratories, Carlsbad, CA), according to the manufacturer’s protocol. Cells were observed with an inverted Olympus FSX100 microscope (Olympus, Tokyo, Japan). Concentration (in nM) of bound Alizarin Red S was then determined by measuring the absorbance at 405 nm and using a calibration curve. Untreated cells (incubated without quebecol) were used as a control. The mineralization assays were performed in triplicate in two independent experiments. **Statistical Analysis.** Unless indicated otherwise, all analyses were performed in triplicate in two independent experiments. The data is expressed as mean ± standard deviations (SD). Statistical analyses were performed using a one-way or two-way analysis of variance with a post hoc Bonferroni multiple comparison test (GraphPad Software Inc., La Jolla, CA). All results were considered statistically significant at p < 0.001 or p < 0.05.
Complete tabulated results for the secretion of MMP-3, MMP-8, and MMP-9; proliferation of osteoblast study (PDF)

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Notes
The authors declare no competing financial interest.

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