# Essential oils: a Chemotherapeutic Option in Periodontics

Asquino, Natalia\*, García, Ma. Victoria\*\*, Mayol, Magdalena\*, Andrade, Ernesto\*\*, Bueno Rossy, Luis Alexandro\*\*\*

## Abstract

Periodontal diseases are a group of pathologies that affect the tissues that support teeth. The efficient control of daily oral biofilm can prevent these diseases. Most individuals have difficulty in developing good oral hygiene habits and use only mechanical elements. This has led to the development of chemical adjuvants so that patients can maximize biofilm control. Essential oils (EOs) are effective and more efficient at controlling supragingival plaque and inflammation compared to a placebo and to cetylpyridinium chloride. Nevertheless, EOs were similar in their antiinflammatory effectiveness and less efficient in plaque control than chlorhexidine, causing fewer adverse effects. Current evidence suggests that chlorhexidine remains the first choice for short-term oral health care and that essential oils are best indicated for long-term treatments.

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<sup>\*</sup> Teaching Assistant, Periodontics Department, Universidad de la República. Uruguay.

<sup>\*\*</sup> Assistant Professor, Periodontics Department, Universidad de la República. Uruguay.

<sup>\*\*\*</sup> Professor, Periodontics Department, Universidad de la República. Uruguay. Director of Postgraduate Degree in Periodontics, Universidad de la República. Uruguay. Specialist in Implant Dentistry, University of Guarulhos. Brazil.

## Introduction

Periodontal diseases are a group of pathologies that affect the tissues that support teeth. Gingival diseases involve only the gingival margin, but if the attached gingiva is also involved, the condition is called Periodontitis. Clinical attachment loss, pathological pockets and bone resorption are typical signs of periodontal disease. Lack of treatment may lead to tooth loss (1). Periodontal disease has high prevalence among adults and lower social classes (2-5) and is currently one of the main public health concerns (6).

It includes pathologies with different clinical manifestations, as similar diagnosis may include furcal involvement, gingival recession, tooth mobility, mucogingival complications, etc. These diseases differ in their etiology, natural history and response to therapy, but their pathogenesis shows similar events, which can be modified by genetic factors and/or risk factors (7).

Dental biofilm and its byproducts are considered precipitating factors of periodontal disease. The link between dental biofilm and the prevalence and severity of these pathologies has been well established (8). Longitudinal studies have shown that these conditions can be successfully treated by removing bacterial deposits, removing tartar and teaching oral hygiene practices (9, 10).

Gingivitis can be prevented by controlling oral biofilm daily, frequently and effectively (11). Therefore, treating and eliminating gingivitis would be the most efficient way to prevent periodontitis. However, several studies show how difficult it is to help people develop good oral hygiene habits using only mechanical elements (12-15). Hence the need to supplement dental plaque removal using mechanical elements with chemical (chemotherapeutic) adjuvants (15, 16).

Ideally, the goal of periodontal therapy is to

reduce the number of periodontopathogenic species that cause periodontal disease and to keep their numbers low. Therefore, the infection is treated by reducing the microbial load and/or by modifying the subgingival habitat (17).

Currently available chemotherapeutic colutory formulations include: Triclosan/ Copolymer, Cetylpyridinium Chloride (CPC), Chlorhexidine (CHX) Digluconate, or a fixed combination of essential oils as active ingredients.

#### Nature of essential oils

Essential oils (EOs) are organic compounds made up of various constituents obtained from plants for specific purposes. The formula includes four active ingredients: Eucalyptol 0.092%, Menthol 0.042%, Methyl Salicylate 0.060%, Thymol 0.064% (18).

#### Action mechanisms

EOs have proven to be effective at controlling inflammation and supragingival biofilm. They are also safe for patient use (19-21). They have the ability to alter the cell surface of specific microorganisms and eliminate their enzymatic activity (22). They can also inhibit the endotoxins of Gramnegative pathogens (23). In vitro and in vivo studies have shown how EOs can penetrate the dental biofilm and have a bactericidal effect (24, 25).

#### Anti-inflammatory and antiplaque action

Phenolic compounds show anti-inflammatory action and inhibit prostaglandin synthesis, act as scavengers of oxygen free-radicals, thus affecting leukocyte activity. Studies conducted on animal cells show that phenolic compounds commonly used in these chemical formulations (thymol, menthol, eucalyptol) inhibited neutrophil chemotaxis and superoxide generation in neutrophils (concentration-dependent), and they also led to the elimination or scavenging of freeradicals and the inhibition of prostaglandins. These aromatic compounds have a free phenolic hydroxyl group which accounts for the anti-inflammatory action described above (26, 27).

#### Essential oils versus placebo

An alcohol-based mouthwash with EOs has been used as a chemotherapeutic agent, showing relevant clinical reductions of supragingival plaque and inflammation both in short (28) and long (29) term studies.

In a 6-month randomized controlled clinical trial, EO mouthwashes showed a reduction of up to 70% in oral biofilm and a reduction of up to 36% in gingivitis compared to a control group (5% hydroalcoholic solution) among patients with mild to moderate levels of plaque and inflammation (P<0.001) (30). Likewise, in 15-day models, EO colutories showed a reduction in oral biofilm greater than 21% (P<0.001) and a 12% greater reduction of inflammation than the control group (5% hydroalcoholic solution) on patients with mild or moderate gingival disease (P<0.001) (31).

Alcohol-free EO mouthwash: In a 15-day evolution clinical trial, the group using the test product had a mean Plaque Index (PI) lower than that of the control group, with a 23.9% reduction (P<0.001). Additionally, when the dependent variable was the Modified Gingival Index (MGI), there was a 10.4% reduction. Regarding Bleeding on Probing (BOP) Index, the secondary efficacy variable, the control group showed a 53.8% reduction in the ratio of bleeding areas (p<0.001) (32). Alcohol-free mouthwash with EOs led to a significant reduction in dental plaque (31.6%) and gingival inflammation (24%) compared to a negative control (33) 6 months after the beginning of the trial (P<0.001).

**Essential oils vs Chlorhexidine (CHX):** In a recent systematic review, EOs were considered an alternative to CHX to control gingival diseases.

Colutories containing CHX were more effective regarding dental plaque values, but no significant difference was found in gingival inflammation. The most probable explanation is that the mouthwash with CHX has an antiplaque effect, while EOs have a direct anti-inflammatory effect. Chlorhexidine is significantly superior at reducing bacterial deposits when compared to EOs, which is not the case for the long-term reduction of gingival inflammation. Secondary effects (stains and tartar) were more serious in CHX users compared to EOs users (34).

**Essential oils vs Cetylpyridinium Chloride** (**CPC**): Long-term studies have shown that the group using EOs had a reduction in the plaque index which was 56.2% higher than that of the group treated with 0.05% CPC (P<0.001). Regarding the MGI, the changes showed a reduction greater than 32.4% compared to the CPC group (P<0.001) (30, 33).

An EO formulation with zinc chloride (as anti-tartar agent), sodium fluoride and acidulated phosphate (for cavity control) was compared to a negative control and a CPCbased mouthwash. After three months, the EO-based mouthwash was more effective at reducing biofilm when compared to the negative control and to the CPC. These results remained the same for six months. Both regarding dental plaque and inflammation after 3 and 6 months, the EO mouthwash was better than the negative control and CPC (18).

#### Essential oils and oral cancer

The link between EO-based colutories and the risk of developing oral cancer has been controversial for decades, ever since the initial remarks made by Weaver et al. (35).

Daily alcohol consumption (ethanol) has

proven to be a risk factor for oral cancer.

A number of mouthwashes contain ethanol in concentrations that range between 5% and 27%. Ethanol improves product solubility, stabilization and conservation, thus modifying its taste and increasing antiplaque properties. In a 18% - 27% concentration, ethanol improves the antibacterial action of EOs (high penetration achieved in 30 seconds) (36).

Currently it is not possible to conclude that there is a statistically significant link between the use of alcohol-based mouthwash and the risk of oral cancer. Likewise, there is no significant indication that the risk increases with daily use, as evidence is mostly retrospective.

### Indications

Hass and col. (41) conducted a systematic review of the efficacy in biofilm and gingival inflammation control in patients wearing fixed orthodontic appliances. These authors concluded that after 6 months, plaque and gingivitis levels decreased up to 50% among these patients when used as adjuvants to mechanical therapy. Regarding implant therapy, the systematic use of essential oils twice a day, in a 3-month follow-up period, led to a significant reduction in the plaque and tissue inflammation percentages when compared to the placebo (hydroalcoholic solution) (42).Patients undergoing periodontal maintenance therapy would also benefit from this mouthwash as an adjuvant to mechanical therapy (34).

# Conclusions

Chlorhexidine is the most effective antiplaque agent, which makes it the first therapeutic short-term option. Considering their adverse effects, mouthwashes with essential oils seem to be a reliable alternative for long-term use.

# References

- 1. Armitage GC. Development of a classification system for periodontal diseases and conditions. Ann Periodontol Am Acad Periodontol. 1999; 79:1-6.
- Albandar JM, Brunelle JA, Kingman A. Destructive periodontal disease in adults 30 years of age and older in the United States, 1988-1994. J Periodontol. 1999; 70:13-29.
- 3. Susin C, Dalla Vecchia CF, Oppermann R V, Haugejorden O, Albandar JM. Periodontal attachment loss in an urban population of Brazilian adults: effect of demographic, behavioral, and environmental risk indicators. J Periodontol 2004; 75(7):1033-41.
- Susin C, Oppermann R V, Haugejorden O, Albandar JM. Tooth loss and associated risk indicators in an adult urban population from south Brazil. Acta Odontol Scand. 2005; 63(2):85-93.
- Haas AN, Gaio EJ, Oppermann RV, Rösing CK, Albandar JM SC. Pattern and rate of progression of periodontal attachment loss in an urban population of South Brazil: a 5-years population-based prospective study. J Clin Periodontol. 2012; 39(1):1-9.
- 6. Tonetti MS, Chapple ILC. Biological approaches to the development of novel periodontal therapies--consensus of the Seventh European Workshop on Periodontology. J Clin Periodontol. 2011; 114-8.
- 7. Page RC, Offenbacher S, Schroeder HE, Seymour GJ, Kornman KS. Advances in the pathogenesis of periodontitis: summary of developments, clinical

implications and future directions. Periodontol 2000 1997; 14:216–48.

- Moore WE, Holdeman LV, Smibert RM, Cato EP, Burmeister JA, Palcanis KG, et al Bacteriology of experimental gingivitis in young adult humans. Infect Immun 1982; 38:651-67.
- Claffey, N., Loos, B., Gantes, B., Martin, M., Heins, P. & Egelberg. The relative effects of therapy and periodontal disease on loss of probing attachment after root debridment. J Clin Periodontol 1988; 15:163–69.
- van der Weijden, G. A. & Timmerman, M. F.A systematic review on the clinical efficacy of subgingival debridement in the treatment of chronic periodontitis. J Clin Periodontol 2000; 29:(3) 55–71.
- 11. Löe H, Theilade E JB. Experimental gingivitis in man. J Periodontol. 1965; 36:177-87.
- 12. Lang NP, Cumming BR LH. Toothbrushing frequency as it relates to plaque development and gingival health. J Periodontol. 1973; 44(7): 396–405.
- 13. Hioe KPKJ, Van Der Weijden GA. The effectiveness of self-performed mechanical plaque control with triclosan containing dentifrices. Int J Dent Hyg. 2005; 3:192-204.
- 14. Oppermann RV, Haas AN, Villoria GEM, Primo LG, Serra-Negra JM, Ferreira EF e, et al. Proposal for the teaching of the chemical control of supragingival biofilm. Braz Oral Res. 2010; 24:(1)33– 6. http://dx.doi.org/10.1590/S1806-83242010000500006.
- Rode SDM, Gimenez X, Montoya VC, Gómez M, Blanc SL De, Medina M, et al. Daily biofilm control and oral health: consensus on the epidemiological challenge - Latin American Advisory Panel. Braz Oral Res. 2012; 26(1):133– 43. http://dx.doi.org/10.1590/S1806-

83242012000700020

- 16. Johnson NW. Hygiene and health: the value of antiplaque agents in promoting oral health. Int Dent J. 1993; 43:375-86.
- 17. Teles RP, Haffajee AD, Socransky SS. Microbiological goals of periodontal therapy. Periodontol 2000 2006: 180-218.
- 18. Cortelli SC, Cortelli JR, Wu M-M, Simmons K, Charles CA. Comparative antiplaque and antigingivitis efficacy of a multipurpose essential oil-containing mouthrinse cetylpyridinium and а mouthrinse: chloride-containing А trial. 6-month randomized clinical Quintessence Int. 2012; 43:82-94.
- 19. Overholser CD, Meiller TF, DePaola LG, Minah GE, Niehaus C. Comparative effects of 2 chemotherapeutic mouthrinses on the development of supragingival dental plaque and gingivitis. J Clin Periodontol. 1990; 17:575-9.
- 20. Sharma N, Charles CH, Lynch MC, Qaqish J, McGuire JA, Galustians JG, et al. Adjunctive benefit of an essential oilcontaining mouthrinse in reducing plaque and gingivitis in patients who brush and floss regularly: a six-month study. J Am Dent Assoc 2004; 135 496–504.
- 21. Stoeken J, Paraskevas S, Van der Weijden G. The long-term effect of a mouthrinse containing essential oils on dental plaque and gingivitis: a systematic review. J Periodontol. 2007; 7(78):1218–28.
- 22. Kubert D, Rubin M, Barnett ML, Vincent JW. Antiseptic mouthrinseinduced microbial cell surface alterations. Am J Dent. 1993; 6:277-9.
- 23. Fine DH, Furgang D, Lieb R, Korik I, Vincent JW, Barnett ML. Effects of sublethal exposure to an antiseptic mouthrinse on representative plaque bacteria. J Clin Periodontol. 1996; 23:444-51.

- 24. Charles CH, Pan PC, Sturdivant L, Vincent JW. In Vivo Antimicrobial Activity of an Essential Oil-Containing Mouthrinse on Interproximal Plaque Bacteria. J Clin Dent. 2000; 11:94-7.
- 25. Ouhayoun J-P. Penetrating the plaque biofilm: impact of essential oil mouthwash. J Clin Periodontol. 2003; 30(5):10–2.
- 26. Dewhirst FE. Structure-activity relationships for inhibition of prostaglandin cyclooxygenase by phenolic compounds. Prostaglandins 1980; 20:209-22.
- 27. Azuma Y, Ozasa N, Ueda Y, Takagi N. Pharmacological studies on the antiinflammatory action of phenolic compounds. J Dent Res 1986; 65:53-6.
- 28. Lusk S, Bowers G, Tow H, Watson W, Moffitt W. Effects of an oral rinse on experimental gingivitis plaque formation, and formed plaque. J Am Soc Prev Dent. 1974; 4(4):31–3.
- 29. Gunsolley JC. A meta-analysis of six-month studies of antiplaque and antigingivitis agents. J Am Dent Assoc 1939. 2006; 137:1649-57.
- 30. Sharma NC, Araujo MWB, Wu MM, Qaqish J, Charles CH. Superiority of an essential oil mouthrinse when compared with a 0.05% cetylpyridinium chloride containing mouthrinse: a six-month study. Int Dent J. 2010; 60:175-80.
- 31. Amini P, Araujo M, Wu M, Charles C, Sharma N. Comparative antiplaque and antigingivitis efficacy of three antiseptics mouthrinses: a two week randomized clinical trial. Braz Oral Res. 2009; 23:319-25. http://dx.doi.org/10.1590/ S1806-83242009000300016
- 32. Charles CA, Amini P, Gallob J, Shang H, McGuire JA, Costa R. Antiplaque and antigingivitis efficacy of an alcohol-free essential-oil containing mouthrinse: A

2-week clinical trial. Am J Dent. 2012; 25:195-8.

- 33. Cortelli SC, Cortelli JR, Shang H, Mcguire JA, Charles CA. Long-term management of plaque and gingivitis using an alcoholfree essential oil containing mouthrinse: A 6month randomized clinical trial. Am J Dent. 2013; 26:149-55.
- 34. Van Leeuwen MPC, Slot DE, Van der Weijden GA. Essential oils compared to chlorhexidine with respect to plaque and parameters of gingival inflammation: a systematic review. J Periodontol. 2011; 82:174-94.
- 35. Weaver A, Fleming SM, Smith DB. Mouthwash and oral cancer: carcinogen or coincidence? J Oral Surg. 1979; 37(4):250–3.
- 36. Gandini S, Negri E, Boffetta P, La Vecchia C, Boyle P. Mouthwash and oral cancer risk - Quantitative meta-analysis of epidemiologic studies. Ann Agric Environ Med. 2012; 19:173-80.
- 37. Elmore J, Horwitz R. Oral cancer and mouthwash use - evaluation of the epidemiologic evidence. Otolaryngol Neck Surg. 1995; 113(3):253–61.
- 38. Shapiro S, Castellana J V, Sprafka JM. Alcohol-containing mouthwashes and oropharyngeal cancer: a spurious association due to underascertainment of confounders? Am J Epidemiol. 1996; 144:1091–5.
- 39. Cole P, Rodu B, Mathisen A. Alcoholcontaining mouthwash and oropharyngeal cancer: a review of the epidemiology. J Am Dent Assoc. 2003; 134:1079–87.
- 40. Vecchia CL. Mouthwash and oral cancer risk: An update. Oral Oncology. 2009; 45:198–200.
- Haas A, Mendes C, Andrade AK, Escobar EC, Almeida ER, Costa FO, Cortelli JR, Cortelli SV, Rode SM, Pedrazzi V, Oppermann RV. Mouthwashes for

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the control of supragingival biofilm and gingivitis in orthodontic patients: evidence-based recommendations for clinicians. Braz Oral Res [Internet]. 2014; 28 (Spec Iss 1):1-8. [Cited: 2016 Apr]. Available from: http://dx.doi. org/10.1590/1807-3107-BOR-2014. vol28.0021.

42. Ciancio, F. Lauciello, O. Shibly, M. et al. The Effect of an Antiseptic Mouthrinse on Implant Maintenance: Plaque and Peri-Implant Gingival Tissues. J Periodontol. 1995: Nov; 962-65.

Natalia Asquino: natalia.asquino@gmail.com