

## Essential oils, their therapeutic properties, and implication in dentistry: A review

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### Abstract

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#### Background:

Antibacterial treatments currently used for treatment cause several side effects, and bacterial resistance to the antibiotics is also increasing. Therefore, there is need to find better alternatives. Essential oils (EOs) have been used for treatment of various ailments since ancient times and have gained popularity over the years. Safety and efficacy of EOs have been proved by several clinical trials. This review gives an overview on the EOs, their uses, and adverse effects.

#### Materials and Methods:

A literature search was performed in the PubMed for clinical trial studies and review articles on EOs published up to February 2015. The search was performed during March 2015. The following keywords were used: “Lavender essential oil,” “cinnamon oil,” “clove oil,” “eucalyptus oil,” “peppermint oil,” “lemon EOs,” and “tea tree oil.”

#### Results:

Total 70 relevant articles were found in PubMed database. After screening of abstracts, 52 articles were selected to be included in the present review.

#### Conclusion:

On the basis of the available information, it can be concluded that EOs have the potential to be developed as preventive or therapeutic agents for various oral diseases, but further clinical trials are required to establish their safety and efficacy.

**Keywords:** *Alternative medicine, essential oils, oral pathogens*

## INTRODUCTION

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According to the World Oral Health Report, despite great improvements in oral health in several countries, oral health problems still persist, particularly among underprivileged groups in both developing and developed countries.[1] Dental caries and periodontal diseases are identified as the most important among oral health problems globally. Oral diseases adversely affect the general health too. Quality of life and the working capacity of an individual are also affected.[2]

The antibacterial agents that are currently used for treatment of oral health problems are reported to cause several side effects such as diarrhea, vomiting, etc., Increasing bacterial resistance to the drugs is also a major concern. Because of the adverse effects, increasing bacterial resistance, and high cost associated with the standard therapeutic procedure, there is a need to explore new therapeutic agents and conduct further clinical research on traditional medicines obtained from various plant sources.

Many traditionally used medicines for treating infections have been studied again, and clinical trials are being done to establish their efficacy and possible side effects. One of these natural medicines is essential oils (EOs).[3,4] In the recent years, there has been an increased interest toward EOs.

Approximately 3000 Eos are known till now.[5] EOs are one of the plant extracts that have been used for treatment of various medical and dental problems since ancient times. These are secondary metabolites produced by various medicinal plants and possess antibacterial, antifungal, and antioxidant properties. [6,7,8]

The purpose of this systematic review is to analyze the published data related to the EOs. A number of studies have been conducted to prove the therapeutic properties of various EOs, but very few reviews have been published on their implication in dental treatment. The review gives an overview on the EOs, their therapeutic properties, and adverse effects.

## MATERIALS AND METHODS

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To identify relevant literature, an electronic search was performed on PubMed database.

Titles and abstracts were screened. Only articles related to lavender oil, eucalyptus oil, clove oil, cinnamon oil, and lemon EOs have been included in this review. Studies related to several other EOs were excluded. Total 52 articles found relevant were selected for this review.

### EOs and their composition

EOs are secondary metabolites of plants whose constituents are basically a complex mixture of terpenic hydrocarbons, especially monoterpenes and sesquiterpenes, and oxygenated derivatives such as aldehydes, ketones, epoxides, alcohols, and esters.[9] EOs greatly differ in their compositions. Even the composition of EOs extracted from the plants of same species differ in different geographic locations.[10] Composition also depends on the maturity of the plant from which the EOs are extracted.[10,11]

### Mechanism of action

The mechanisms of action of EOs are dependent on their chemical composition and the location of one or more functional groups on the molecules present in them.[12]

Membrane damage is proposed to be the main mechanism of action.[13] Solubility of EOs in the phospholipid bilayer of cell membranes seems to have an important role in their antimicrobial activity. Clove oil has reported to reduce the quantity of ergosterol which is found specifically in fungal cell membrane.[14] Terpenoids in EOs have been found to interfere with the enzymatic reactions of energy metabolism.[15]

Essential oils that have potential to be used in oral disease prevention and treatment are discussed subsequently.

### Lavender oil

**Composition** Major components found are linalool, linalyl acetate, 1,8-cineole, B-ocimene, terpinen-4-ol, l-fenchone, camphor, and viridiflorol.[10,16] However, the relative level of each of these constituents varies in different species. Lavender oil, obtained from the flowers of *Lavandula angustifolia* (Family: Lamiaceae) by steam distillation, is chiefly composed of linalyl acetate (3,7-dimethyl-1,6-octadien-3-yl acetate), linalool (3,7-dimethylocta-1,6-dien-3-ol), lavandulol, 1,8-cineole, lavandulyl acetate, and camphor.

The activity of linalool reflects that of the whole oil, indicating that linalool may be the active component of lavender oil.[13]

### Therapeutic properties

- Antimicrobial activity: EOs extracted from *Lavandula stoechas* L. exhibit good antimicrobial activities against most of the bacteria, filamentous fungi, and yeasts. In the study of Benabdelkader *et al.*, minimum inhibitory concentrations were found to be ranging from 0.16 to 11.90 mg/ml.[10] It also shows antipseudomonal activity[16]
- *In vitro* study on the antibacterial activity of the EO of *Lavandula coronopifolia* against antibiotic-resistant bacteria suggested its bactericidal effect[17]
- Anxiolytic: Lavender EO is reported to reduce stress, anxiety, and improve mood when inhaled or orally administered.[18,19] It is not very effective in cases of high anxiety[20]
- Antifungal: EOs of *Lavandula luisieri* show an inhibitory effect on yeast, dermatophyte, and *Aspergillus* strains.[21] *Lavandula viridis* is reported to have fungicidal effect. *Cryptococcus neoformans* is the most sensitive fungus, followed by *Candida* species.

### Eucalyptus oil

**Composition** The main component is 1,8-cineole followed by cryptone,  $\alpha$ -pinene, *p*-cymene,  $\alpha$ -terpineol, trans-pinocarveol, phellandral, cuminal, globulol, limonene, aromadendrene, spathulenol, and terpinene-4-ol.[22]

### Therapeutic properties

- Antimicrobial effect: Antimicrobial activity was found to be related to the synergic effects between major and minor components rather than the concentration of a single component.[22] EO of the leaves of *Eucalyptus globulus* has antimicrobial activity against Gram-negative bacteria (*Escherichia coli*) as well as Gram-positive bacteria (*Staphylococcus aureus*).[23] Studies done on eight eucalyptus species show that *Eucalyptus odorata* oil possesses strong cytotoxic effect and also antibacterial effect against *S. aureus*, *Haemophilus influenzae*, *Staphylococcus pyogenes*, and *Staphylococcus pneumoniae*. *Eucalyptus bicostata* and *Eucalyptus astringens* showed antibacterial effects[22]
- Anti-inflammatory effect: Immunoregulatory agent: The study of Serafino *et al.* demonstrates that eucalyptus EO can stimulate the innate cell-mediated immune response suggesting its use as adjuvant in immunosuppression, in infectious disease, as well as in tumor chemotherapy.[24]

### Peppermint oil

Peppermint (*Mentha piperita*) oil is one of the most popular and widely used EOs. In the EO from *M. piperita*, menthol is identified as the major compound, followed by menthyl acetate and menthofuran.[25]

### Therapeutic properties

- Antibacterial: Peppermint oil shows an inhibitory effect on the proliferation of staphylococci[26]
- Antifungal: Studies show that EOs exhibit fungistatic and fungicidal activities against both the standard and clinical strains of *Candida* species at concentrations ranging from 0.5 to 8 µL/mL. EOs exhibit similar antifungal effect against the azole-resistant and azole-susceptible strains[25]
- Antibiofilm: Biofilm inhibition in fungal strains helps to decrease pathogenesis and drug resistance. Studies show that EO inhibits the biofilm formation of *Candida albicans* completely up to 2 µl/ml in a dose-dependent manner.[25]

**Melaleuca alternifolia (Myrtaceae)** It is also known as Tea Tree Oil (TTO). Its composition shows terpinen-4-ol,  $\gamma$ -terpinene, *p*-cymene,  $\alpha$ -terpinene, 1,8-cineole,  $\alpha$ -terpineol, and  $\alpha$ -pinene.[27]

### Effects

- Antibacterial: In a clinical trial, the melaleuca gel was found to possess an inhibitory effect on various bacterial colonies and dental biofilm.[28] It shows strong antibacterial action against oral pathogens[29]
- Antifungal activity: *Melaleuca alternifolia* possesses antimycotic activity, terpinen-4-ol being its most effective component.[30]

### Lemon EO

**Composition** Mostly, it contains almost exclusively terpenes and oxygenated terpenes.[31]

Therapeutic activity shows antifungal potential against three *Candida* species (*C. albicans*, *Candida tropicalis*, and *Candida glabrata*). Lemon EO is suggested to be used as an effective remedy against candidiasis caused by *C. albicans*. [31,32]

### Clove oil

Main constituents found in the clove bud oil are the phenylpropanoids eugenol, eugenyl acetate, carvacrol, thymol, cinnamaldehyde,  $\beta$ -caryophyllene, and 2-heptanone, when analyzed by gas chromatography. [33,34]

**Medicinal properties** Eugenol is well-known for its therapeutic properties and is widely used in dentistry.

- Antioxidant: When tested against *tert*-butylated hydroxytoluene, EO exhibited a very strong radical scavenging activity[33]
- Antifungal: It possesses antifungal activity.[33] Clove oil and its main content eugenol also reduce the quantity of ergosterol, which is a specific component of fungal cell membrane. Germ tube formation by *C. albicans* is also inhibited[14]
- Antibacterial: It was found to possess inhibitory effect on multi-resistant *Staphylococcus* spp.[34]

### Cinnamon oil

#### Composition

The volatile oils obtained from the bark, leaf, and root barks vary significantly in chemical composition. Three of the main components of the EOs obtained from the bark of *Cinnamomum zeylanicum* are *trans*-cinnamaldehyde, eugenol, and linalool, which represent 82.5% of the total composition. Cinnamaldehyde is the major constituent of cinnamon EO, and studies show that it is the most active component too.[35]

### Medicinal properties

Antimicrobial effect: Inhibitory effect on the growth of various isolates of bacteria including Gram-positive, Gram-negative, and fungi.[36]

Antimutagenic: It has antimutagenic potential against spontaneous mutations in human cells.[37] Furthermore, the study of Cabello *et al.* performed in animals shows that oral administration of cinnamaldehyde (CA) exerts significant anti-melanoma activity.[38]

Besides these activities, studies suggest that *Cinnamomum zeylanicum* (CZ) has antiparasitic, antioxidant, and free radical scavenging properties.[39]

### Implications in dental practice

Potential implications of EOs have been described below and the information is consolidated in [Table 1](#).

**Lavender oil** It can be used in dental clinics to reduce patients' anxiety. It is found to be useful as an anxiolytic agent when used in waiting area.[18,20] The study performed by Zabirunnisa showed statistically significant reduction in anxiety scores when the fragrance of lavender oil was used at the reception area. It is also helpful during surgical procedures, as it has been shown to reduce the pain of needle insertion.[19]

**Eucalyptus oil** It shows an inhibitory effect on oral pathogens like *Lactobacillus acidophilus*, which makes this suitable to be used as an anticariogenic agent.[44]

### Peppermint oil

Eugenol oil is used widely in dentistry. It is active against oral pathogens associated with dental caries and periodontal disease.[45] Studies done on five EOs (TTO, lavender oil, thyme oil, peppermint oil, and eugenol oil) against four common oral pathogens (*S. aureus*, *Enterococcus faecalis*, *E. coli*, and *C. albicans*) showed significant inhibitory effect of eugenol oil, peppermint oil, and TTO. Among them, eugenol oil showed antimicrobial activity at the lowest concentration level.[41]

TTO and some of its individual components, specifically terpinen-4-ol, exhibit strong antimicrobial efficacy against fungal biofilms. TTO can be a solution for the increasing resistance of *C. albicans* to established antifungal drugs. It can be used to treat oral candidiasis[42] and is suitable for use in prophylactic oral hygiene products. The study performed by Ramage *et al.* shows that it is more appropriate and safe to use terpinen-4-ol, the major component of TTO, than TTO itself.[46]

### Cinnamon oil

A Phase I clinical trial conducted on cinnamon EO concluded that it is safe to be used in healthy patients with dentures for the treatment of oral candidiasis.[40]

### Lemon EO

Lemon EO is suggested to be used as an effective remedy against candidiasis caused by *C. albicans*.[32]

### Combination of EOs

Combining EOs and antibiotics can reduce antibiotic resistance in multidrug-resistant bacteria. Peppermint, cinnamon bark, and lavender EOs were found to be antibiotic resistance-modifying agents, when used in combination with piperacillin.[43]

### Studies not supporting the use of EOs

Several studies support the benefits of EOs, but some studies raise questions about their efficacy.

A study in which 0.2% chlorhexidine rinse and an EO mouth rinse were compared for their efficacy showed that EOs are effective only for very short duration, i.e., 2–3 h, and concluded that use of chlorhexidine is preferable over EOs.[47]

A study done on EOs to measure their efficacy when used as a coolant concluded that there was no benefit over water during ultrasonic root debridement for the treatment of chronic periodontitis.[48]

### Adverse effects caused by EOs

Natural medicines are not always free of side effects. Adverse effects are also reported with EOs. In the study of Millet *et al.*, commercial preparations of essences of sage, hyssop, thuja, and cedar have been reported to cause neurotoxicity and human intoxication, of which tonic–clonic convulsions formed the major symptom.[49]

According to a review by Posadzki *et al.*, mild to severe adverse effects including fatality can be caused by EOs like lavender, peppermint, TTO, and ylang-ylang when used in aromatherapy. Most common adverse effect among them was dermatitis.[22]

Toxicological tests are often lacking for traditional medicines. Therefore, further clinical trials are required to exclude the possibility of side effect and poisoning.

### Limitations

Only seven EOs that are found to be used commonly are included in this review. The review is subjected to publication bias as it is written on the basis of published literature. Only English language articles were referred. Article search was performed only in one database, PubMed.

## CONCLUSIONS

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As described in this review, there is considerable evidence that EOs have potential to be developed as preventive or therapeutic agents for various oral diseases. Although several other potential uses of EOs have been described[50] and many claims of therapeutic efficacy have been validated adequately by either *in vitro* testing or *in vivo* clinical trials, still there is need for conducting further research to establish the safety and efficacy of these EOs before including them in clinical practice. If used properly, they may prove very useful in dental therapy and may contribute in improving the quality of dental treatments.

In particular, clinical trials that confirm the therapeutic potential of EOs *in vivo* and address issues such as adverse effects, toxicity, and their interaction with other drug molecules would be of great value.

### Footnotes

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**Conflict of Interest:** None declared.

## REFERENCES

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1. Petersen PE. The World Oral Health Report 2003: Continuous improvement of oral health in the 21st century-the approach of the WHO Global Oral Health Programme. *Community Dent Oral Epidemiol.* 2003;31(Suppl 1):3–23. [PubMed: 15015736]
2. Palombo EA. Traditional medicinal plant extracts and natural products with activity against oral bacteria: Potential application in the prevention and treatment of oral diseases. *Evid Based Complement Alternat Med* 2011. 2011 680354. [PMCID: PMC3145422] [PubMed: 19596745]
3. Cowan MM. Plant products as antimicrobial agents. *Clin Microbiol Rev.* 1999;12:564–82. [PMCID: PMC88925] [PubMed: 10515903]
4. Kalembe D, Kunicka A. Antibacterial and antifungal properties of essential oils. *Curr Med Chem.* 2003;10:813–29. [PubMed: 12678685]
5. Thosar N, Basak S, Bahadure RN, Rajurkar M. Antimicrobial efficacy of five essential oils against oral pathogens: An *in vitro* study. *Eur J Dent.* 2013;7(Suppl 1):S71–7. [PMCID: PMC4054083] [PubMed: 24966732]
6. Baratta MT, Dorman HJ, Deans SG, Figueiredo AC, Barroso JG, Ruberto G. Antimicrobial and antioxidant properties of some commercial essential oils. *Flavour Fragr J.* 1998;13:235–44.
7. Hammer KA, Carson CF, Riley TV. Antimicrobial activity of essential oils and other plant extracts. *J Appl Microbiol.* 1999;86:985–90. [PubMed: 10438227]
8. Guleria S, Tiku AK, Koul A, Gupta S, Singh G, Razdan VK. Antioxidant and antimicrobial properties of the essential oil and extracts of *Zanthoxylum alatum* grown in north-western Himalaya. *Scientific World Journal* 2013. 2013 790580. [PMCID: PMC3679694] [PubMed: 23781160]
9. Rehman SU, Ahmad MM, Kazmi ZH, Raza MS. Physico-chemical variations in essential oils of *Citrus reticulata*. *J Food Sci Technol.* 2007;44:353–6.
10. Benabdelkader T, Zitouni A, Guitton Y, Jullien F, Maitre D, Casabianca H, et al. Essential oils from wild populations of Algerian *Lavandula stoechas* L.: Composition, chemical variability, and *in vitro* biological properties. *Chem Biodivers.* 2011;8:937–53. [PubMed: 21560242]
11. Kiran CR, Chakka AK, Amma KP, Menon AN, Kumar MM, Venugopalan VV. Influence of cultivar and maturity at harvest on the essential oil composition, oleoresin and [6]-gingerol contents in fresh ginger from northeast India. *J Agric Food Chem.* 2013;61:4145–54. [PubMed: 23570262]
12. Dorman HJ, Deans SG. Antimicrobial agents from plants: Antibacterial activity of plant volatile oils. *J Appl Microbiol.* 2000;88:308–16. [PubMed: 10736000]
13. Prashar A, Locke IC, Evans CS. Cytotoxicity of lavender oil and its major components to human skin cells. *Cell Prolif.* 2004;37:221–9. [PMCID: PMC6496511] [PubMed: 15144499]
14. Pinto E, Vale-Silva L, Cavaleiro C, Salgueiro L. Antifungal activity of the clove essential oil from *Syzygium aromaticum* on *Candida*, *Aspergillus* and dermatophyte species. *J Med Microbiol.* 2009;58:1454–62. [PubMed: 19589904]
15. Knobloch K, Pauli A, Iberl B, Weis N, Weigand H. Antibacterial activity and antifungal properties of essential oil components. *J Essent Oils Res.* 1988;1:119–28.
16. Végh A, Bencsik T, Molnár P, Böszörményi A, Lemberkovics E, Kovács K, et al. Composition and antipseudomonal effect of essential oils isolated from different lavender species. *Nat Prod Commun.* 2012;7:1393–6. [PubMed: 23157020]

17. Ait Said L, Zahlane K, Ghalbane I, El Messoussi S, Romane A, Cavaleiro C, et al. Chemical composition and antibacterial activity of *Lavandula coronopifolia* essential oil against antibiotic-resistant bacteria. *Nat Prod Res.* 2015;29:582–5. [PubMed: 25174508]
18. Lehrner J, Marwinski G, Lehr S, Jöhren P, Deecke L. Ambient odors of orange and lavender reduce anxiety and improve mood in a dental office. *Physiol Behav.* 2005;86:92–5. [PubMed: 16095639]
19. Kim S, Kim HJ, Yeo JS, Hong SJ, Lee JM, Jeon Y. The effect of lavender oil on stress, bispectral index values, and needle insertion pain in volunteers. *J Altern Complement Med.* 2011;17:823–6. [PubMed: 21854199]
20. Bradley BF, Brown SL, Chu S, Lea RW. Effects of orally administered lavender essential oil on responses to anxiety-provoking film clips. *Hum Psychopharmacol.* 2009;24:319–30. [PubMed: 19382124]
21. Zuzarte M, Gonçalves MJ, Cruz MT, Cavaleiro C, Canhoto J, Vaz S, et al. *Lavandula luisieri* essential oil as a source of antifungal drugs. *Food Chem.* 2012;135:1505–10. [PubMed: 22953886]
22. Posadzki P, Alotaibi A, Ernst E. Adverse effects of aromatherapy: A systematic review of case reports and case series. *Int J Risk Saf Med.* 2012;24:147–61. [PubMed: 22936057]
23. Bachir RG, Benali M. Antibacterial activity of the essential oils from the leaves of *Eucalyptus globulus* against *Escherichia coli* and *Staphylococcus aureus*. *Asian Pac J Trop Biomed.* 2012;2:739–42. [PMCID: PMC3609378] [PubMed: 23570005]
24. Yap PS, Lim SH, Hu CP, Yiap BC. Combination of essential oils and antibiotics reduce antibiotic resistance in plasmid-conferred multidrug resistant bacteria. *Phytomedicine.* 2013;20:710–3. [PubMed: 23537749]
25. Saharkhiz MJ, Motamedi M, Zomorodian K, Pakshir K, Miri R, Hemyari K. Chemical composition, antifungal and antibiofilm activities of the essential oil of *Mentha piperita* L. *ISRN Pharm* 2012. 2012 718645. [PMCID: PMC3532871] [PubMed: 23304561]
26. Witkowska D, Sowinska J. The effectiveness of peppermint and thyme essential oil mist in reducing bacterial contamination in broiler houses. *Poult Sci.* 2013;92:2834–43. [PubMed: 24135585]
27. Pereira TS, de Sant’anna JR, Silva EL, Pinheiro AL, de Castro-Prado MA. *In vitro* genotoxicity of *Melaleuca alternifolia* essential oil in human lymphocytes. *J Ethnopharmacol.* 2014;151:852–7. [PubMed: 24315850]
28. Santamaria M, Jr, Petermann KD, Vedovello SA, Degan V, Lucato A, Franzini CM. Antimicrobial effect of *Melaleuca alternifolia* dental gel in orthodontic patients. *Am J Orthod Dentofacial Orthop.* 2014;145:198–202. [PubMed: 24485734]
29. Takarada K, Kimizuka R, Takahashi N, Honma K, Okuda K, Kato T. A comparison of the antibacterial efficacies of essential oils against oral pathogens. *Oral Microbiol Immunol.* 2004;19:61–4. [PubMed: 14678476]
30. Terzi V, Morcia C, Faccioli P, Valè G, Tacconi G, Malnati M. *In vitro* antifungal activity of the tea tree (*Melaleuca alternifolia*) essential oil and its major components against plant pathogens. *Lett Appl Microbiol.* 2007;44:613–8. [PubMed: 17576222]
31. Trombetta D, Castelli F, Sarpietro MG, Venuti V, Cristani M, Daniele C, et al. Mechanisms of antibacterial action of three monoterpenes. *Antimicrob Agents Chemother.* 2005;49:2474–8. [PMCID: PMC1140516] [PubMed: 15917549]

32. Białoń M, Krzyśko-Łupicka T, Koszałkowska M, Wieczorek PP. The influence of chemical composition of commercial lemon essential oils on the growth of *Candida* strains. *Mycopathologia*. 2014;177:29–39. [PMCID: PMC3915084] [PubMed: 24436010]
33. Chaieb K, Zmantar T, Ksouri R, Hajlaoui H, Mahdouani K, Abdelly C, et al. Antioxidant properties of the essential oil of *Eugenia caryophyllata* and its antifungal activity against a large number of clinical *Candida* species. *Mycoses*. 2007;50:403–6. [PubMed: 17714361]
34. Chaieb K, Hajlaoui H, Zmantar T, Kahla-Nakbi AB, Rouabhia M, Mahdouani K, et al. The chemical composition and biological activity of clove essential oil, *Eugenia caryophyllata* (*Syzygium aromaticum* L. Myrtaceae): A short review. *Phytother Res*. 2007;21:501–6. [PubMed: 17380552]
35. Naveed R, Hussain I, Tawab A, Tariq M, Rahman M, Hameed S, et al. Antimicrobial activity of the bioactive components of essential oils from Pakistani spices against *Salmonella* and other multi-drug resistant bacteria. *BMC Complement Altern Med*. 2013;13:265. [PMCID: PMC3853939] [PubMed: 24119438]
36. Ooi LS, Li Y, Kam SL, Wang H, Wong EY, Ooi VE. Antimicrobial activities of cinnamon oil and cinnamaldehyde from the Chinese medicinal herb *Cinnamomum cassia* Blume. *Am J Chin Med*. 2006;34:511–22. [PubMed: 16710900]
37. King AA, Shaughnessy DT, Mure K, Leszczynska J, Ward WO, Umbach DM, et al. Antimutagenicity of cinnamaldehyde and vanillin in human cells: Global gene expression and possible role of DNA damage and repair. *Mutat Res*. 2007;616:60–9. [PMCID: PMC1955325] [PubMed: 17178418]
38. Oliveira Jde A, da Silva IC, Trindade LA, Lima EO, Carlo HL, Cavalcanti AL, et al. Safety and tolerability of essential oil from *Cinnamomum zeylanicum* blume leaves with action on oral candidosis and its effect on the physical properties of the acrylic resin. *Evid Based Complement Alternat Med* 2014. 2014 325670. [PMCID: PMC4276295] [PubMed: 25574178]
39. Ramage G, Milligan S, Lappin DF, Sherry L, Sweeney P, Williams C, et al. Antifungal, cytotoxic, and immunomodulatory properties of tea tree oil and its derivative components: Potential role in management of oral candidosis in cancer patients. *Front Microbiol*. 2012;3:220. [PMCID: PMC3376416] [PubMed: 22719736]
40. Ranasinghe P, Pigera S, Premakumara GA, Galappaththy P, Constantine GR, Katulanda P. Medicinal properties of 'true' cinnamon (*Cinnamomum zeylanicum*): A systematic review. *BMC Complement Altern Med*. 2013;13:275. [PMCID: PMC3854496] [PubMed: 24148965]
41. Elaissi A, Rouis Z, Salem NA, Mabrouk S, ben Salem Y, Salah KB, et al. Chemical composition of 8 eucalyptus species' essential oils and the evaluation of their antibacterial, antifungal and antiviral activities. *BMC Complement Altern Med*. 2012;12:81. [PMCID: PMC3475086] [PubMed: 22742534]
42. Ishnava KB, Chauhan JB, Barad MB. Anticariogenic and phytochemical evaluation of *Eucalyptus globules* Labill. *Saudi J Biol Sci*. 2013;20:69–74. [PMCID: PMC3730900] [PubMed: 23961222]
43. Cai L, Wu CD. Compounds from *Syzygium aromaticum* possessing growth inhibitory activity against oral pathogens. *J Nat Prod*. 1996;59:987–90. [PubMed: 8904847]
44. Serafino A, Sinibaldi Vallebona P, Andreola F, Zonfrillo M, Mercuri L, Federici M, et al. Stimulatory effect of *Eucalyptus essential* oil on innate cell-mediated immune response. *BMC Immunol*. 2008;9:17. [PMCID: PMC2374764] [PubMed: 18423004]
45. van de Braak SA, Leijten GC. Rotterdam: CBI Centre for the Promotion of Imports from Developing Countries; 1994. Essential Oils and Oleoresins: A Survey in the Netherlands and Other Major Markets in the European Union; p. 116.

46. Jandourek A, Vaishampayan JK, Vazquez JA. Efficacy of melaleuca oral solution for the treatment of fluconazole refractory oral candidiasis in AIDS patients. *AIDS*. 1998;12:1033–7. [PubMed: 9662200]
47. Cabello CM, Bair WB 3rd, Lamore SD, Ley S, Bause AS, Azimian S, et al. The cinnamon-derived Michael acceptor cinnamic aldehyde impairs melanoma cell proliferation, invasiveness, and tumor growth. *Free Radic Biol Med*. 2009;46:220–31. [PMCID: PMC2650023] [PubMed: 19000754]
48. Malhotra S, Yeltiwar RK. Evaluation of two mouth rinses in reduction of oral malodor using a spectrophotometric technique. *J Indian Soc Periodontol*. 2011;15:250–4. [PMCID: PMC3200021] [PubMed: 22028512]
49. Millet Y, Jouglard J, Steinmetz MD, Tognetti P, Joanny P, Arditti J. Toxicity of some essential plant oils. Clinical and experimental study. *Clin Toxicol*. 1981;18:1485–98. [PubMed: 7333081]
50. Dagli N, Dagli R. Possible use of essential oils in dentistry. *J Int Oral Health*. 2014;6:i–ii. [PMCID: PMC4109163] [PubMed: 25083049]

## **Figures and Tables**

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**Table 1**

Essential oils and their potential implications in dentistry

<b>Name of EO</b>	<b>Potential implications in dentistry</b>
Lavender EO	As an anxiolytic in dental clinics Reduces pain of needle insertion <sup>[19]</sup>
Eucalyptus EO	Anticariogenic agent
Peppermint EO	Antimicrobial activity Use in oral hygiene products
Cinnamon EO	In treating oral candidiasis <sup>[40]</sup>
Lemon EO	In treating candidiasis <sup>[32]</sup>
Eugenol EO	Shows antimicrobial activity against several oral pathogens <sup>[41]</sup>
Tea Tree Oil	Oral candidiasis <sup>[42]</sup> Suitable for use in prophylactic oral hygiene products
Combination of EOs	Antibiotic resistance-modifying agent <sup>[43]</sup>

*EO=Essential oil*