Use of essential oils to inhibit *Alicyclobacillus acidoterrestris*: a short overview of the literature

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Essential oils (EOs) are promising and friendly antimicrobials for the prolongation of the shelf life of many foods. They have been extensively used to inhibit spoiling and pathogenic microorganisms of many kinds of products like fruit juices and acidic drinks. Therefore, they could be used successfully to control the germination of spores of *Alicyclobacillus acidoterrestris*, that finds in these products an optimal environment for growth. This paper reports a brief overview of the literature available, focusing on the effects of EOs toward alicyclobacilli.

**Keywords:** *Alicyclobacillus acidoterrestris*, essential oils, spore inhibition, active compounds, friendly compounds

**ESSENTIAL OILS: DEFINITION AND PERSPECTIVES**

**DEFINITION AND COMPOSITION**

Essential oils (EOs), also called ethereal or volatile oils, are complex mixture extracted from various part of plants (Burt, 2004; Speranza and Corbo, 2010). Apart from a great variability amongst the different oils, EOs are soluble in lipids (i.e., are oils) and possess some compounds, acting as antimicrobials, perfumes, or flavoring (i.e., they contain an *essentia*, as suggested by Paracelsus von Hohenheim in sixteenth century; Speranza and Corbo, 2010). Actually, ca. 3,000 different EOs are known, but only few hundreds are commonly used as food-grade additives and for the production of soap, perfumes, and cosmetics (Burt, 2004; Speranza and Corbo, 2010).

Essential oils can be produced by distillation, expression, enfleurage, and fermentation, being the hydro-distillation the most used system, and virtually they can be extracted form every part of a plant, i.e., from leaves (e.g., basil, oregano, eucalyptus, lemon grass, rosemary, peppar mint), from flowers (orange, rose, marjoram, clove), peel (lemon, bergamot, orange, tangerine), seeds (anise, cumin).

Most of the commercially distributed EOs have been studied and characterized and the results published by some International Organizations (European Pharmacopoeia, International Organization for Standardization, World Health Organization, Council of Europe). Chemically, EOs are very complex mixture of 60 or more molecules; however, they contain one or more compounds at high levels and the others in trace. The major component of an EOs is usually labeled “active compound” (Speranza and Corbo, 2010).

**BIOACTIVITY**

There many literature reports on the use of EOs as antimicrobials, both in laboratory media and in foods (Burt, 2004; Bevilacqua et al., 2008c; Speranza and Corbo, 2010); due to the extreme variability in chemical structure, composition, and source, different modes of action have been postulated (Burt, 2004; Speranza and Corbo, 2010):

1. Due to their solubility in lipids, EOs pass through the cell wall and disrupt membrane, causing its permeabilization, the loss of ions and the reduction of membrane potential, the collapse of the proton pump, and the depletion of the ATP pool (Dorman and Deans, 2000; Walsh et al., 2003).
2. EOs can coagulate the cytoplasm and cause damages to lipids and proteins (Jerkovic et al., 2001).
3. In eukaryotic cells, EOs provoke depolarization of the mitochondrial membranes by decreasing the membrane potential, influence ionic \(Ca^{2+}\) cycling and other ionic channels, and reduce the pH gradient, affecting (as in bacteria) the proton pump and the ATP pool. (Armstrong, 2006).

What are the most susceptible microorganisms to EOs? Due to the extreme variability of compounds and the various methods proposed in the literature for the evaluation of their antimicrobial
ESSENTIAL OILS AS FOOD-GR ADE ADDITIVES

Many EOs and their active compounds are in the list EAFUS (everything added to food in the United States) of the Food and Drug Administration [FDA; Center for Food Safety and Applied Nutrition (CFSAN), 2011; www.cfsan.fda.gov]. As an example, the status of thymol can be reported: thyme EO and thyme are considered generally recognized as safe (21 CFR 180.10 and 180.20), while thymol is labeled as a synthetic food additive (21 CFR 172.515). According to Annex II of Council Regulation 2377/90, the Committee of Experts on Flavoring Substances of the European Commission (EC) has registered thymol in the flavorings in foodstuff list. The same status has been recognized for carvacrol, cineole, cinnamaldehyde, citral, p-cymene, eugenol, limonene, and menthol (Speranza and Corbo, 2010).

WHY USE ESSENTIAL OILS TO INHIBIT ALICYCLOBACILLUS SPP.

The use of EOs for the inhibition and/or the control of spore germination was postulated firstly by Chaibi et al. (1997), in the light of a renewed interest in 1990s toward natural antimicrobials and low-impact preservation technologies. Although it is not clear how EOs damage spores, the use of spices, and plant extract was proposed for the inhibition of different spore-former microorganisms (Burt, 2004; Tajkarami et al., 2010). For example, it is well known that many oils are effective against Bacillus spp. and Sofia et al. (2007), based on the results of a disk diffusion assay, proposed the following bioactivity scale (from the strongest oil to the less effective one): clove > mustard > cinnamon > garlic > ginger > mint. Other authors used EOs against Clostridium spp.: Davidson and Naidu (2000) reported the bioactivity of clove oil toward C. perfringens, while Friedman (2007) proposed the extract of black tea for the inactivation of the neurotoxin of C. botulinum. Both Davidson and Naidu (2000) and Friedman (2007) referred to antimicrobial assays performed through different protocols (disk diffusion assay, micro-dilution method).

From these studies to the application of EOs against an emerging spoiling microorganism (Alicyclobacillus acidoterrestris) there is a short distance. Alicyclobacillus spp. have been recognized as an increasing threat in food industry, due to its thermo-resistance and thermo-acidophilic behavior (Bevilacqua et al., 2008c). Silva and Gibbs (2004) proposed A. acidoterrestris as a target to design thermal treatments for fruit juices, since it appeared more heat resistant than other spoiling microorganisms. Heat resistance of spores is greatly variable and relies upon various elements (the strain, the pH and kind of medium, the conditions attained throughout sporulation); Silva and Gibbs (2004) reported that D-value of spores at 95°C in juices could range from 0.06 to 5.3 min and z-value varied from 7.2 to 12.9°C.

Apart from strain variability, it has been reported that alicyclobacilli can survive thermal treatments used by juice producers (90–95°C from second to minutes); a possibility could be the use of a more severe processing, thus affecting the sensorial quality of juices, in terms of color, odor, and nutrient content (Bevilacqua and Corbo, 2010a).

A second possibility could be the use of a friendly approach, i.e., a chemical or a physical treatment able to inactivate spore without or with low effects on the sensorial quality of juices. This approach, based on the green consumerism philosophy (Bevilacqua et al., 2010b), has been proposed successfully for alicyclobacilli, through the use of high pressure processing and high pressure homogenization, ultrasound, microwave an natural antimicrobials (Bevilacqua et al., 2008c).

The use of natural antimicrobials (e.g., lysozyme, nisin, other bacteriocins, and essential oils) has gained an increased interest for the low-impact of these compounds on human health and environment. Essential oils appear as a promising way, as many of them are extracted from fruits and their use in fruit-based products can be friendly for the characteristics of raw material.

USE OF ESSENTIAL OILS AGAINST ALICYCLOBACILLUS ACIDOTERRESTRIS: ELEMENTS ACTING ON THEIR BIOACTIVITY

Despite the increased and renewed interest toward the antimicrobial activity of EOs, only few papers focused on their effectiveness against alicyclobacilli, namely against A. acidoterrestris. The first report was by Takahashi et al. (2004), who proposed the use of leaf extracts and flavonoids from Eucalyptus spp. and found a minimum inhibitory concentration of extracts ranging from 7.8 g/l (E. globosa) to 31 g/l (E. saligna). Then, this topic was addressed exclusively by the research group of Applied Microbiology of University of Foggia; therefore, in the following sheets there is a brief overview of their results, with the aim to give an insight into EOs effect along with the factors that can enhance or limit their bioactivity.

CHEMICAL STRUCTURE AND SECONDARY GROUPS

Bevilacqua et al. (2008a) used cinnamaldehyde, eugenol, and limonene (0.05–0.5 g/l; Figure 1 for the chemical structure) against the spores of two different strains of A. acidoterrestris (c8 and γ4, isolated respectively form soil and spoiled pear juice; Sini-gaglia et al., 2003; Bevilacqua et al., 2006) and modeled the data as inhibition index, i.e., as percentage of absorbance at 420 nm referred to the control. Cinnamaldehyde was the most effective
compound and a concentration of 500 ppm inhibited completely spore germination for 13 days; otherwise a lower amount of this compound (100 ppm), inhibited the microbial target by 96–97 and 58–70% after 8 and 13 days, respectively.

Eugenol appeared as less effective than cinnamaldehyde, as it inhibited significantly spore germination only at the highest amounts (500 ppm). The stronger effect of cinnamaldehyde was later confirmed by Bevilacqua et al. (2010c), who combined cinnamaldehyde and eugenol to inhibit and/or control the germination of a cocktail of *A. acidoterrestris* strains and found that 160 ppm of eugenol prolonged the lag phase of the microbial target by 1.5 days, whereas this parameter increased by 4–4.5 days with 80 ppm of cinnamaldehyde added. Bevilacqua et al. (2008a) studied also the bioactivity of limonene, but this oil was not effective in inhibiting spore germination.

The antimicrobial activity of the natural compounds seems to be related to the phenolic rings, but the type of alkyl group has been found to influence the antibacterial effectiveness (Burt, 2004). The results of Bevilacqua et al. (2008a) confirmed this idea: in fact, the phenolic ring is the major part of all the compounds (limonene, eugenol, and cinnamaldehyde). The difference amongst the various antimicrobials is the secondary group, linked to phenol.

**SOURCE OF THE OIL**

The composition of EOs is quite variable and depends upon the climate, the location of plants, as well as the method of oil extraction; moreover, the extraction of an oil from different parts of the same plant could result in a strong variability and in a different bioactivity (Burt, 2004).

Bevilacqua et al. (data not published) studied the antimicrobial activity of three essential oils, extracted from various part of citrus and/or lemon, i.e., neroli, lemon extract, and biocitro (a complex citrus extract); the bioactivity of this compound was tested against two different strains of *A. acidoterrestris*: the isolate DSMZ 2498, purchased from a Public Collection, and the wild strain c8, isolated from soil (Bevilacqua et al., 2006).

Neroli (extracted from *Citrus aurantium* var. *amara* or Bigaradia) is a plant oil similar in scent to bergamot; it is produced from the blossom of the bitter orange tree and contains α-pinene, camphene, β-pinene, α-terpinene, neryl acetate, farnesol, geraniol, linalool, nerolidol, linalyl acetate, methyl anthranilate, and indole.

Biocitro is a complex oil purchased from Quinabara (Probena, Spain) and extracted from citrus; the producer reports the following composition for the oil: ascorbic acid and ascorbates (vitamin C), linked with citrus bioflavonoids, 4.0–7.20%; hydrated glycerin linked with other traces of citrus polyphenols, carbohydrates, bio-flavoproteins, pectin, citrus sugars, citric acid, 30.80–36.60%; water, 6.00–11.00%; stabilizer and inert carrier, 50.00%. Chemical analyses revealed that the concentration of ascorbic acid and citrus bioflavonoid is ca. 56,000 ppm; amongst the bioflavonoids, naringin is 6,500 ppm minimum. Finally, the content of limonene is 30,000–50,000 ppm (Bevilacqua et al., 2010d).

Lemon extract (or oil) shows a quite variable composition. Generally it is extracted from the fresh fruit peel by cold expression. The main chemical components of lemon oil are α-pinene, camphene, β-pinene, sabine, myrcene, α-terpinene, linalool, β-bisabolene, limonene, trans-β-bergamotene, nerol, and neral. This extract was used by several authors (Conte et al., 2007, 2009) as a natural antimicrobials toward the spoiling microflora of fruit and vegetables, showing some promising properties.

Neroli, lemon extract, and citrus extract were used to inhibit the spores of different strains of *A. acidoterrestris* and a micro-dilution method in a laboratory medium (Malt Extract broth, acidified to pH 4.5), combined with the traditional plate count, was used to point out oil effectiveness (Bevilacqua et al., data not published). Neroli was able to inhibit spore germination only at high concentrations (>500 ppm), not compatible with foods, as the odor of this extract was too strong; on the other hand both biocitro (citrus extract) and lemon extract showed some interesting results, as they were able to reduce spore concentration by 1–2 log cfu/ml after 1 day at 20–80 ppm; this variability in MIC value relied upon the strains, as some isolate appeared more resistant than others.

Besides being effective against *A. acidoterrestris*, Bioctiro was also active against spoilage microflora of juices (*Bacillus coagulans*, *Lactobacillus brevis* and *L. plantarum*, *Saccharomyces bayanus*, *Pichia membranifaciens*, and *Rhodotorula bacarum*) as reported by Bevilacqua et al. (2010d).

Why the oils showed a different bioactivity? Many and different ideas could be suggested; however, the main difference between neroli and biocitro/lemon extract is the source: neroli is generally extracted from flowers of orange tree, otherwise the source of citrus and lemon extract is citrus peel. Thus it could suggested that
both the kind of plant and oil source (i.e., part of the plan from which the oil is extracted) play a fundamental role for the antimicrobial activity, as reported also by Speranza and Corbo (2010).

**pH OF THE MEDIUM**

A factor playing a fundamental role in the bioactivity of EOs is the pH of the medium; this idea was supported by some papers (Bevilacqua et al., 2008b, 2009b, 2010c,e), addressing this topic and reporting the combinations of cinnamaldehyde (the active compound of cinnamon oil) and eugenol (the main component of clove oil), both in laboratory media and in juices. The use of acidic pH enhanced the effect of active compounds, due to its effect on the partition equilibrium and hydrophobicity; it is generally accepted that EOs constituents are most effective at acidic pHs (Burt, 2004; Ait-Ouazzou et al., 2011); however, that conclusion is mainly based on studies carried out to evaluate their capacity to prevent microbial growth. Regarding bactericidal activity, results varied among studies: Friedman et al. (2004) did not find any influence of acidity when evaluating carvacrol or linalool in apple juice, while Rivas et al. (2010) recovered a higher susceptibility of *E. coli* to carvacrol or thymol when decreasing the pH to 4. The trend experienced by *A. acidoterrestris* seemed to confirm this effect, although more data are required to postulate a general hypothesis.

**COMBINATION OF EOs WITH PHYSICAL HURDLES**

A promising way for the inhibition of alicyclobacilli is the use of the combined approach, i.e., a multi-target preservation based on the theory of multiple hurdles (Bevilacqua et al., 2008c). For example, it is possible to combine cinnamonaldehyde with a heat treatment: the combination of the antimicrobial at 97 ppm with a thermal treatment at 90°C for 8 min, conducted at pH 4.3, assured the maximal combination of the antimicrobial at 97 ppm with a thermal treatment or homogenization, respectively. This effect was related to a phenomenon due to the isolation source (Bevilacqua et al., 2008a) and to a probable difference in the hydrophobic behavior of the strains (i.e., the different distribution of the fatty acids in the membrane). The more hydrophobic an organism behaves, the more sensitive it will be to a hydrophobic antimicrobial agents, as the active compound of EOs. This idea, however, is only a speculation and needs to be confirmed by molecular and chemical analyses on the genes involved in the susceptibility to EOs constituents and fatty acids distribution in the membrane.

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An interesting effect clearly revealed by the cited authors is that it is not convenient to increase processing temperature indefinitely, because a thermal degradation of cinnamaldehyde could occur at a thermal treatment of 12 min/90°C.

**STRAIN VARIABILITY**

Finally, a major role in the bioactivity of EOs is played by strain variability; this effect was postulated by Bevilacqua et al. (2008a) and Bevilacqua et al. (2009a), with reference to the bioactivity of cinnamonaldehyde and eugenol alone or combined with a thermal treatment or homogenization, respectively. This effect was related to a phenomenon due to the isolation source (Bevilacqua et al., 2008a) and to a probable difference in the hydrophobic behavior of the strains (i.e., the different distribution of the fatty acids in the membrane). The more hydrophobic an organism behaves, the more sensitive it will be to a hydrophobic antimicrobial agents, as the active compound of EOs. This idea, however, is only a speculation and needs to be confirmed by molecular and chemical analyses on the genes involved in the susceptibility to EOs constituents and fatty acids distribution in the membrane.

**CONCLUSION AND FUTURE PERSPECTIVES**

The use of EOs against *A. acidoterrestris* appears as a promising way; however, the research is at the beginning. The reports available in the literature show some interesting directions, but further investigations are required, in order to understand some critical issues:

1. the real effect of EOs against spore;
2. the influence of the source of oil and strain variability on the bioactivity of EOs and their active compounds;
3. the impact of EOs on food shelf life for a prolonged contact time (i.e., months or years);
4. the ability of EOs to control post-contamination and spore recovery;
5. the existence of additive effects with some physical hurdles;
6. the correlation of the sensitivity toward EOs with the membrane.

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