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(54) **Title:** AN ANTI BACTERIAL COMPOSITION

(57) **Abstract:** The present invention relates to a method of disinfecting a surface and to an antibacterial composition. Essential oils, which are used as antibacterial actives, are also known for their strong odour; using high amounts of these gives a strong smell to the product that is not always appreciated by the consumer. It is therefore an object of the invention to provide an antibacterial composition, having good anti-bacterial properties, at very low levels of essential oil actives. The present inventors have achieved this using a synergistic combination of select anti-bacterial actives, specific polymers and select hydrotropes.



AN ANTIBACTERIAL COMPOSITION

Technical Field

The present invention relates to a method of disinfecting a surface and to an
5 antibacterial composition. It particularly relates to an antibacterial composition for
disinfecting various substrates like external surface of the human or animal body like
skin and hair, the oral cavity, and hard surfaces in households or other indoor and
outdoor environs.

10 Background and Prior Art

Sanitizing and disinfecting compositions comprising chlorine and nascent oxygen
based bleaching agents are known. Such compositions require rather long contact time
to provide efficacious antibacterial action. In practice, users, in particular children, do
not spend long time in cleaning and as a result, cleaning with such compositions does
15 not provide adequate prevention from surface or topical infection or adequate
protection against diseases. The user, in spite of cleaning hands, is likely to have skin
with relatively inadequate bacterial removal and may cause contamination of further
animate and/or inanimate surfaces and this leads to spreading of pathogens and
consequent diseases. Further, many antibacterial actives in addition to abrasives are
20 included in oral care compositions like dentifrices but these actives generally require
several minutes if not hours before effective antibacterial action is achieved.

Similarly in the area of hard surface cleaning e.g. cleaning of floors, table tops or
utensils, the antibacterial actives in the compositions are in contact with the substrate
25 for less than a few minutes after which the surface is either wiped off or rinsed with
water. These short time scales of cleaning action are ineffective in providing the
desired benefit since most known antibacterials commonly used in such products take
several hours to provide the desired kill of microbes.

30 The present applicants in pursuit of solving this problem have disclosed in
WO2010046238 a combination of essential oil actives, thymol and terpineol that
interact synergistically to provide anti-bacterial activity in very fast times, in many cases
as low as 15 seconds or lesser.

Many essential oils actives are relatively expensive ingredients. Additionally, essential oils are also known for their strong odour; using high amounts of these gives a strong smell to the product that is not always appreciated by the consumer.

5

Accordingly it remains to be desired to prepare anti-bacterial compositions having a high anti-bacterial effect, even with a low dosage of anti-bacterial essential oil actives.

The present applicants in pursuit of this objective, have, in W01 1151 172 disclosed a
10 combination of two essential oil actives along with a polymer of a select class to provide synergistic antibacterial action at low concentration of the essential oil actives.

Further, in W01 1151 171 the present applicants have disclosed a combination of two polymers along with an essential oil active to provide synergistic antibacterial action at
15 low concentration of the essential oil actives. It has been found that from the polymers and essential oil actives chosen, a single polymer and a single essential oil active in combination do not provide the synergistic activity. It has now been found that if only one essential oil active and one polymer is used, the enhanced effect is seen only when a hydrotrope of the present invention is included.

20

In the paper published in the Letters in Applied Microbiology 2002, 34, 168-172 titled "Chitosan potentiates the antimicrobial action of sodium benzoate on spoilage yeasts" the synergistic activity against three yeasts has been demonstrated using a combination of sodium benzoate and chitosan glutamate. There is no indication that
25 this combination can be useful against bacteria.

The present applicants have been working on further improving this technology and in an effort to find solutions to the problem of achieving high antibacterial efficacy at even further lower concentrations of essential oil actives, have arrived at the present invention. They have found that a combination of specific essential oil actives along
30 with a polymer of a select class in the presence of specific hydrotropes are able to achieve the high antibacterial efficacy at even lower concentration of the actives which is not achieved by each of the ingredients singly or as binary combinations.

It is therefore an object of the invention to provide an antibacterial composition, having good anti-bacterial properties, at very low levels of essential oil actives.

SUMMARY OF THE INVENTION

- 5 According to the first aspect of the invention there is provided an antibacterial composition comprising
- (a) 0.01 to 10% by weight of an essential oil active selected from the group consisting of eugenol, thymol, geraniol, terpineol, and mixtures thereof;
 - (b) a polymer selected from the group consisting of polymers of vinyl alcohol,
10 chitosan, and mixtures thereof; and
 - (c) a hydrotrope selected from the group consisting of sodium benzoate, sodium toluene sulphonate, sodium cumene sulphonate, sodium xylene sulphonate, sodium salicylate, sodium acetate, and mixtures thereof.
- 15 Another aspect of the present invention provides for a method for providing an anti-bacterial effect to a substrate comprising the steps of:
- (a) applying a composition of the first aspect to the substrate, and
 - (b) waiting for at least 15 seconds.
- 20 According to yet another aspect, the present invention provides use of a composition comprising 0.01 to 10% by weight of an essential oil active selected from the group consisting of eugenol, thymol, geraniol, terpineol and mixtures thereof; a polymer selected from the group consisting of chitosan, polymers of vinyl alcohol, , and mixtures thereof; and a hydrotrope selected from the group consisting of sodium benzoate,
25 sodium toluene sulphonate, sodium cumene sulphonate, sodium xylene sulphonate, sodium salicylate, or sodium acetate and mixtures thereof; for fast reduction in bacterial count.

DETAILED DESCRIPTION OF THE INVENTION

These and other aspects, features and advantages will become apparent to those of ordinary skill in the art from a reading of the following detailed description and the
5 appended claims. For the avoidance of doubt, any feature of one aspect of the present invention may be utilised in any other aspect of the invention. The word "comprising" is intended to mean "including" but not necessarily "consisting of" or "composed of." In other words, the listed steps or options need not be exhaustive. It is noted that the examples given in the description below are intended to clarify the invention and are
10 not intended to limit the invention to those examples per se. Similarly, all percentages are weight/weight percentages unless otherwise indicated.

Except in the operating and comparative examples, or where otherwise explicitly indicated, all numbers in this description indicating amounts of material or conditions of
15 reaction, physical properties of materials and/or use are to be understood as modified by the word "about". Unless specified otherwise, numerical ranges expressed in the format "from x to y" are understood to include x and y. When for a specific feature multiple preferred ranges are described in the format "from x to y", it is understood that all ranges combining the different endpoints are also contemplated.

20

By an antibacterial composition as used herein, is meant to include a composition for cleaning and disinfecting topical areas e.g. skin and/or hair of mammals, especially humans. Such a composition may be generally classified as leave-on or rinse off, and includes any product applied to a human body for also improving appearance,
25 cleansing, odor control or general aesthetics. It is more preferably a rinse off product. The composition of the present invention may be in the form of a liquid but may also be modified to include a lotion, cream, foam or gel, or toner, or applied with an implement or via a face mask, pad or patch. "Skin" as used herein is meant to include skin on the face and body (e.g., neck, chest, back, arms, underarms, hands, legs, buttocks and
30 scalp). The composition of the invention is also of relevance to applications on any other keratinous substrates of the human body other than skin e.g. hair where products may be formulated with specific aim of providing disinfection and cleaning.

By anti-septic liquid is meant a composition, usually in transparent form which may be coloured or may be substantially colourless that is used to disinfect various animate and inanimate surfaces. The transparent form is usually in micro-emulsion format where oils are dispersed in water using surfactants where the oil droplet size is so small as to be smaller than that which causes diffraction of light thereby providing a substantially transparent appearance to the composition. The anti-septic liquid is usually used after dilution with water usually in the weight ratio of 1:1 to 1:200 preferably in a ratio of 1:10 to 1:80. The diluted solution may be transparent but is preferably turbid or hazy. This liquid is usually used to disinfect surface of human or animal body especially but not necessarily when the skin is wounded. The antiseptic liquid is also used to disinfect fabric especially where it is believed to be high in microorganisms e.g. linen in hospitals, clinics, nappies, and undergarments. The antiseptic liquid may also be used for cleaning floors and other surfaces in homes e.g. in kitchens and bathrooms and in certain public places where cleanliness and disinfection are highly sought after.

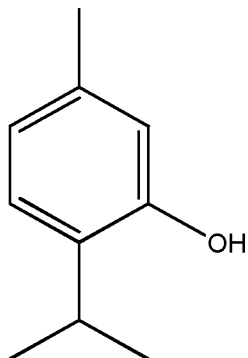
The composition of the invention comprises a selected essential oil active, a selected polymer and a selected hydrotrope. This composition is especially useful since the concentration of essential oil antibacterial actives that needs to be used is low. However, the essential oil actives may also be used at higher concentrations e.g from 0.01 to 10% by weight of the composition.

The composition preferably comprises 0.01 to 5%, more preferably 0.01 to 1%, further more preferably 0.01 to 0.5% by weight of essential oil active. The essential oil active is selected from the group consisting of thymol, terpineol eugenol, geraniol, and mixtures thereof. More preferably, the essential oil active is selected from the group consisting of thymol, terpineol, eugenol, and mixtures thereof. Further more preferably, the essential active is a mixture of thymol and terpineol. Even further more preferred is a mixture of thymol, terpineol and eugenol.

30

Thymol

The structure of thymol is given below:

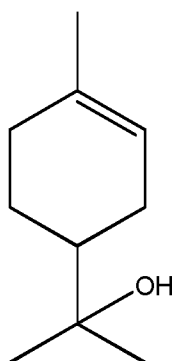


- 5 The composition of the invention comprises preferably 0.01 to 5%, more preferably 0.01 to 1%, further more preferably 0.01 to 0.4%, by weight thymol. Thymol may be added to the composition in purified form. Alternatively, thyme oil or thyme extract comprising thymol may be added to the composition, while ensuring that thymol is present in the desired concentration in the composition of the present invention.
- 10 Thyme oil or thyme extract is obtained from the thyme plant. Thyme plant refers to a plant belonging be genus *Thymus* and includes but is not limited to the following species: *Thymus vulgaris*, *Thymus zygis*, *Thymus saturoides*, *Thymus mastichina*, *Thymus broussonetti*, *Thymus maroccanus*, *Thymus pallidus*, *Thymus algeriensis*, *Thymus serpyllum*, *Thymus pulegoide*, and *Thymus citriodorus* .

15

Terpineol

The structure of a terpineol compound is given below:



The terpeneol is preferably selected from alpha-terpineol, beta-terpineol, gamma-terpineol or mixtures thereof. It is particularly preferred that the terpeneol is alpha-terpineol. Terpeneol may be added to the antibacterial composition in purified form. Alternatively pine oil comprising terpeneol may be added to the antibacterial

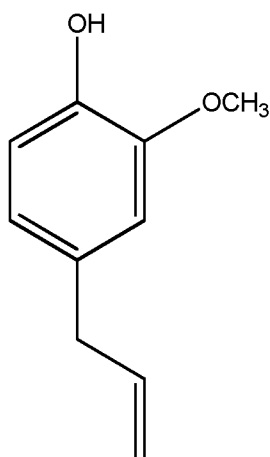
5 composition while ensuring that terpeneol is present in the desired concentration in the composition of the present invention. The composition preferably comprises 0.01 to 5%, more preferably 0.02 to 5%, further more preferably 0.03 to 1%, and even more preferably 0.04 to 0.6% by weight terpeneol.

10 Eugenol

Eugenol is an allyl chain-substituted guaiacol. It is generally extracted from certain spices like clove or cinnamon. Eugenol has been used as a perfumery component, in preparing flavors, as an antiseptic or as a local anesthetic. The composition of the invention preferably comprises 0.005 to 5%, preferably 0.02 to 1%, more preferably

15 0.03 to 0.4%, by weight eugenol.

Eugenol has the structure:



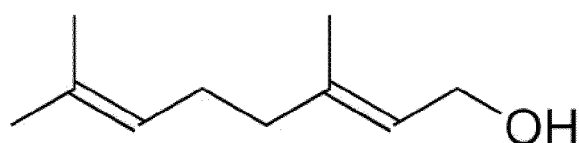
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The composition may preferably comprise a combination of thymol and terpeneol in any of the preferred concentrations as specified above for thymol and terpeneol, respectively. For instance, the composition may preferably comprise a mixture of 0.01 to 0.6% thymol and 0.02 to 1.5% terpeneol by weight of the composition. The

composition of the present invention most preferably comprises a mixture of 0.01 to 0.4% eugenol, 0.01 to 0.6% thymol, and 0.02 to 1.5% terpineol by weight of the composition.

5 Geraniol

The structure of the geraniol compound is given below:



- 10 Geraniol is a monoterpenoid and an aliphatic alcohol. It is the primary component of rose oil, palmrosa oil and citronella oil. It has a rose- -like odor and is commonly used in perfumes. The composition of the invention preferably comprises 0.005 to 5%, preferably 0.02 to 1%, more preferably 0.03 to 0.4%, by weight geraniol.
- 15 The composition of the invention comprises a polymer. The polymer is selected from the group consisting of polymers of vinyl alcohol, chitosan and mixtures thereof. The polymer is preferably present in 0.001 to 25%, more preferably in 0.1 to 1.0 and most preferably in 0.1 to 0.2 % by weight of the composition.
- 20 The composition comprises a hydrotrope that is selected from the group consisting of sodium benzoate, sodium toluene sulphonate, sodium cumene sulphonate, sodium xylene sulphonate, sodium salicylate, sodium acetate, and mixtures thereof. The more preferred hydrotropes are sodium benzoate, sodium acetate and sodium salicylate. The hydrotrope is preferably present in 0.2 to 20%, more preferably 0.2 to 8 %, further
- 25 more preferably 1 to 4 % by weight of the composition.

Without wishing to be bound by theory, it is believed that very low concentration of the essential oil actives can be used since the polymer and the hydrotrope interact synergistically through different mechanisms e.g. the polymer ensuring that there is

30 entrapment of actives thereby increasing local concentration and the hydrotrope enhances the entrapment through formation of a complex with the essential oil actives

thereby providing an antibacterial composition that gives the desired action using minimal amount of actives, while ensuring that sensory issues of off odour, irritation are minimized, while ensuring affordability in terms of low cost are provided to the consumer.

5

The antibacterial composition of the invention may be used to develop various personal care and household care products. Examples include personal care compositions in the form of creams, lotions and gels which may provide various other benefits like

10 moisturization, suncreening and skin lightening benefits, malodour control and antiperspirancy. The antibacterial benefits are afforded by the composition of the invention in very fast times e.g. in less than 5 minutes, often in less than one minutes, in some cases in less than 30 seconds and in certain other cases in less than 15 seconds. This fast acting antibacterial composition is especially suited for

15 incorporation in wash-off products e.g. soaps in the form of bars, liquids and gels. These products may be used for personal cleansing e.g. as personal wash soap bars, body wash liquids, shower gels, hand wash liquids, gels and lotions, and as face wash products. It is preferred that the composition of the invention is formulated to have a pH of 3 to 11 preferably 3.5 to 8.0 where the efficacy of the synergistic interaction

20 between the antibacterial actives, the polymer and the hydrotrope is seen to be maximum. The composition may also be used for providing disinfection benefits other personal substrates like hair and the oral cavity. Thus the composition may be used to formulate shampoos, conditions and mouthwashes.

25 The composition of the invention may be used for cleaning hard surfaces and thus may be formulated as a floor cleaner, toilet cleaner or a gel or emulsion for cleaning surfaces in the kitchen like table tops, utensils, crockery or as an oven cleaner.

Particularly preferred carriers for formulating the composition of the invention in the

30 various products mentioned above are water or oil/solvent, more preferred carrier being a mixture of water and oil. In most of the envisaged applications like personal care/washing, oral care and hard surface cleaning, the antibacterial composition may be formulated in an aqueous base (water being the carrier) e.g. products in gel format

or in purely oil/solvent base e.g. products in anhydrous stick form or propellant containing products. However, most preferred product format has an emulsion base (water and oil being the carriers) e.g. soap products in liquid, solid, lotion or semisolid form for hand wash, face wash, body wash, or shaving applications; toothpaste/
5 dentifrices for oral care applications or products for hard surface cleaning in bars or liquids form.

The antibacterial composition preferably comprises 1 to 80% surfactant. In general, the surfactants may be chosen from the surfactants described in well known textbooks like
10 "Surface Active Agents" Vol. 1, by Schwartz & Perry, Interscience 1949, Vol. 2 by Schwartz, Perry & Berch, Interscience 1958, and/or the current edition of "McCutcheon's Emulsifiers and Detergents" published by Manufacturing Confectioners Company or in "Tenside-Taschenbuch", H. Stache, 2nd Edn., Carl Hauser Verlag, 1981. Any type of surfactant, i.e. anionic, cationic, nonionic, zwitterionic or amphoteric
15 can be used.

A particularly preferred surfactant is soap. Soap is a suitable surfactant for personal washing applications of the antibacterial composition of the invention. The soap is preferably C8-C24 soap, more preferably C10-C20 soap and most preferably C12-C16
20 soap.

The antibacterial composition of the invention is useful in hard surface cleaning applications. In such applications preferred surfactants are nonionic surfactants, such as C8-C22, preferably C8-C16 fatty alcohol ethoxylates, comprising between 1 and 8
25 ethylene oxide groups when the product is in the liquid form. When the product is in the solid form for hard surface cleaning applications, surfactants are preferably selected from primary alkyl sulphate, secondary alkyl sulphonates, alkyl benzene sulphonates, or ethoxylated alkyl sulphates. The composition may further comprise an anionic surfactant, such as alkyl ether sulphate preferably those having between 1 and
30 3 ethylene oxide groups, either from natural or synthetic source and/or sulphonic acid. Especially preferred are sodium lauryl ether sulphates. Alkyl polyglucoside may also be present in the composition, preferably those having a carbon chain length between C6 and C16. Suitable surfactant concentrations in liquid forms of hard surface cleaning

application are generally from about 0.5 to 10%, preferably from 1 to 5 % by weight of the composition. In solid compositions, surfactant is preferably present in 5 to 40%, preferably from 10 to 30% by weight of the composition.

- 5 The antibacterial composition of the invention is useful in oral care compositions e.g. in a dentifrice/ toothpaste or oral rinse product. In such applications, preferred surfactants are anionic, nonionic or amphoteric in nature, preferably anionic or amphoteric. Anionic surfactant is preferably an alkali metal alkyl sulphate, more preferably a sodium lauryl sulphate (SLS). Mixtures of anionic surfactants may also be employed. The
- 10 amphoteric surfactant is preferably a betaine, more preferably an alkylamidopropyl betaine (wherein the alkyl group is a linear C10-C18 chain), and most preferably is cocoamidopropyl betaine (CAPB). Mixtures of amphoteric surfactants may also be employed. Suitable surfactant concentrations in oral care application are generally from about 2% to about 15%, preferably from about 2.2% to about 10%, more
- 15 preferably from about 2.5 to about 5% by weight of the total composition.

Thus, in a highly preferred aspect, the antibacterial compositions include soap, alkyl sulphate or linear alkyl benzene sulphonate as the surfactants.

- 20 The composition may further comprise various additional ingredients known to a person skilled in the art. Such additional ingredients include but are not limited to: perfumes, pigments, preservative, emollients, sunscreens, emulsifiers, gelling agents, or thickening agents.
- 25 According to one aspect water is a preferred carrier. When water is present, it is preferably present in at least 1%, more preferably at least 2%, further more preferably at least 5% by weight of the composition. Water is preferably present in 10 to 99% in most formulations prepared with the composition of the invention. Most compositions have water from 55 to 99% by weight of the composition.

30

According to another aspect, inorganic particulate material is also a suitable carrier. When inorganic particulate material is the carrier, the antibacterial composition is in a solid form. Preferably the inorganic particulate material is talc. When the inorganic

particulate material is talc, the solid antibacterial composition is particularly useful as a talcum powder for application on face or body.

According to one preferred aspect, the invention provides for non-therapeutic benefits.

5

Thus, according to yet another aspect of the invention there is provided use of a composition comprising 0.01 to 10% by weight an essential oil active selected from eugenol, thymol, geraniol, or terpineol; a polymer selected from the group of consisting of polymers of vinyl alcohol, chitosan and mixtures thereof; and a hydrotrope selected
10 from sodium benzoate, sodium toluene sulphonate, sodium cumene sulphonate, sodium xylene sulphonate, sodium salicylate, or sodium acetate for fast reduction in bacterial count. The desired reduction in bacterial count is preferably achieved in less than 5 minutes, more preferably less than 2 minutes, even more preferably less than 1 minute, still more preferably less than 30 seconds and even more preferably within 15
15 seconds.

The invention also provides for therapeutic benefits.

Another aspect of the present invention relates to a method for providing an anti-
20 bacterial effect to a substrate of interest comprising the steps of (a) applying a composition of the first aspect to the substrate, and (b) waiting for at least 15 seconds.

The method preferably comprises the step of wiping or rinsing the composition from the substrate after the waiting step.

25

The invention will now be illustrated with the help of the following non-limiting examples.

30

EXAMPLES

Examples 1-6: Compositions to demonstrate the synergistic interaction of the ingredients of the invention

Compositions as shown in Table - 1 were prepared and the compositions were tested for antibacterial efficacy in a 15 seconds contact test, using the following protocol:

Protocol: Contact kill Assay (15 second contact kill)

- 5 The test bacteria *E.coli* ATCC 10536 was grown overnight in TSB broth (Difco - 30gpl) at 37°C for 16hrs. 2ml of this was sub-cultured in 40ml of fresh TSB broth and allowed to grow for 4 hours at 37°C. Then the culture was processed by spinning at 4000 rpm for 5 minutes, washed twice and the cells were then collected. The cell density was
- 10 adjusted at 620 nm to get the final count of 10^8 cfu/ml (0.8 OD). The test solutions/formulations were prepared and kept for 3 hours for maturation.
- 9 ml of the test solution was taken in a sample container and 1 ml of processed culture was added to it. After 15 seconds of contact time, 1 ml of the above mixture was immediately neutralized in D/E broth (Difco - 39gpl). Serial dilution was done in D/E
- 15 broth and plated on TSA (Difco - 40gpl) in duplicates. In case of the control, 1 ml of test culture was added to 9 ml of saline and was serially diluted and plated on TSA. After solidification, the plates were incubated at 37°C for 48 hrs. The residual colonies were counted after 48 hours incubation and efficacy was calculated by comparing with
- 20 control. The reduction in the amount of bacteria is depicted as log reduction of bacteria.

Table - 1

Ingredient	Example 1 weight%	Example 2 weight%	Example 3 weight%	Example 4 weight%	Example 5 weight%	Example 6 weight%
Thymol	0.02	0.02	0.02	0.02	0.02	0.02
Terpineol	0.05	0.05	0.05	0.05	0.05	0.05
PVA	-	-	0.125	0.125	-	-
Chitosan	-	-	-	-	0.125	0.125
Sodium benzoate	-	2.0	-	2.0	-	2.0
Water	To 100	To 100	To 100	To 100	To 100	To 100
Log reduction	No reduction	0.3	5.0	7.2	3.2	7.2

PVA is poly vinyl alcohol

- 5 The data in Table - 1 indicates that in the compositions as per the invention (Examples 4 and 6), the ingredients interact synergistically as compared to compositions comprising individual ingredients or binary combinations. (Examples 1 to 3 and 5).

Examples 7 to 11: Examples demonstrating synergy with other types of polymers and

10 hydrotropes

Several more samples were prepared and the log reduction afforded by these samples were measured and the data is summarized in Table - 2.

Table - 2

Ingredient	Example 1 weight%	Example 7 weight%	Example 8 weight%	Example 9 weight%	Example 10 weight%	Example 11 weight%
Thymol	0.02	0.02	-	0.02	0.02	0.02
Terpineol	0.05	0.05	-	0.05	0.05	0.05
PVA	-	-	-	-	-	-
Chitosan	-	-	0.125	0.125	0.125	0.125
Sodium cumene sulphonate	-	-	-	-	-	2.0
Sodium salicylate	-	2.0	-	-	2.0	
Water	To 100	To 100	To 100	To 100	To 100	To 100
Log reduction	No reduction	4.1	0.1	4.5	7.2	7.2

The data in Table - 2 indicates that compositions as per the invention (Examples 10 and 11) give vastly improved antibacterial efficacy as compared to compositions 5 outside the invention (Examples 1 and Examples 7 to 9) and the data indicates synergistic interaction for the examples of the invention.

Examples 12 to 16: Synergistic interaction when only one essential oil active is used

Several more samples were prepared using geraniol as the essential oil active along 10 with the other ingredients of the invention and the log reduction afforded by these samples were measured and the data is summarized in Table - 3.

Table - 3

Ingredient	Example 12 weight%	Example 13 weight%	Example 14 weight%	Example 15 weight%	Example 16 weight%
Geraniol	0.08	-	0.08	-	0.05
PVA	-	-	-	0.125	0.125
Sodium Benzoate	-	2.0	2.0	-	2.0
Log reduction	0.2	0.1	4.1	0.2	7.2

The data in Table -3 indicates the synergistic interaction between geraniol (an essential oil active) a hydrotrope and a polymer provides enhanced antibacterial efficacy (Example 16) as compared to compositions outside the invention (Examples 12 to 15).

5 Examples 17 to 19: Synergistic interaction when eugenol is used as the essential oil active

Several more samples were prepared using eugenol as the essential oil active along with the other ingredients of the invention and the log reduction afforded by these samples were measured and the data is summarized in Table - 4.

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Table - 4

Ingredient	Example 17 weight%	Example 18 weight%	Example 19 weight%
Eugenol	0.25	0.25	0.25
PVA	-	-	0.125
Sodium Benzoate	-	2.0	2.0
Log reduction	0.3	2.9	7.3

The data in Table -4 indicates synergistic interaction between eugenol (an essential oil active in combination) a hydrotrope and a polymer provides enhanced antibacterial efficacy (Example 19) as compared to compositions outside the invention (Examples 17 and 18).

The invention thus provides for a composition that exhibits vastly improved antibacterial efficacy by synergistic interaction of the ingredients and this is achieved using very low concentration of the essential oil actives.

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Claims

- 1 An antibacterial composition comprising
 - (a) 0.01 to 10% by weight of an essential oil active selected from the group consisting of eugenol, thymol, geraniol, terpineol, and mixtures thereof;
 - (b) a polymer selected from the group consisting of chitosan and polymers of vinyl alcohol and mixtures thereof; and
 - (c) a hydrotrope selected from the group consisting of sodium benzoate, sodium toluene sulphonate, sodium cumene sulphonate, sodium xylene sulphonate, sodium salicylate, sodium acetate, and mixtures thereof.
- 2 A composition as claimed in any claim 1 comprising 0.001 to 25% of said polymer by weight of the composition.
- 3 A composition as claimed in claim 1 or 2 wherein said essential oil active is selected from the group consisting of thymol, terpineol, eugenol and mixtures thereof.
- 4 A composition as claimed in claim 3 wherein said essential oil active comprises thymol and terpineol.
- 5 A composition as claimed in any one of the preceding claims wherein said hydrotrope is selected from the group consisting of sodium benzoate, sodium acetate and sodium salicylate.
- 6 A composition as claimed in any one of the preceding claims comprising 0.2 to 20% of said hydrotrope.
- 7 A composition as claimed in any one of the preceding claims comprising 55 to 99% of water.
- 8 A composition as claimed in any one of the preceding claims, wherein the pH of the composition is between 3.0 and 11.0.

- 9 A method for providing an anti-bacterial effect to a substrate comprising the steps of:
- (a) applying a composition according to any one of preceding claims to the substrate, and
 - (b) waiting for at least 15 seconds.
- 10 A method as claimed in claim 9, wherein the composition is wiped or rinsed from the substrate after step 'b'.
- 11 Use of a composition comprising 0.01 to 10% by weight of an essential oil active selected from the group consisting of eugenol, thymol, geraniol, terpineol and mixtures thereof; a polymer selected from the group consisting of chitosan and polymers of vinyl alcohol, and mixtures thereof; and a hydrotrope selected from the group consisting of sodium benzoate, sodium toluene sulphonate, sodium cumene sulphonate, sodium xylene sulphonate, sodium salicylate, or sodium acetate and mixtures thereof; for reduction in bacterial count in less than 5 minutes.

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2012/073005

A. CLASSIFICATION OF SUBJECT MATTER		
INV.	A61K8/34	A61K8/368
	A61Q19/10	A61Q17/00
ADD.		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols)		
A61K A61Q		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)		
EPO-Internal , WPI Data, CHEM ABS Data		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	EP 0 916 720 A1 (PROCTER & GAMBLE [US]) 19 May 1999 (1999-05-19) paragraphs [0001] , [0004] , [0008] , [0022] , [0024] , [0026] - [0036] , [0039] , [0060] , [0063] claim 1	1-11
A	W0 2011/039630 A1 (YISSUM RES DEV CO [IL]; TOUITOU ELKA [IL]) 7 April 2011 (2011-04-07) paragraphs [0009] , [0010] , [0016] example 10	1-11
----- -/- .		
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input checked="" type="checkbox"/> See patent family annex.		
* Special categories of cited documents : "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family		
Date of the actual completion of the international search		Date of mailing of the international search report
8 March 2013		25/03/2013
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer Lenzen, Achim

INTERNATIONAL SEARCH REPORT

International application No

PCT/EP2012/073005

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>DATABASE WPI Secti on Ch, Week 1992 Thomson Sci enti fic, London, GB; Class A96, AN 1992-103097 XP002682076, "Mi xt. for mfr. of tooth-paste - contai ns addi ti onal mi neral Lami nari a extract, thymol and sodi um benzoate as anti septi cs, and polyvi nyl -pyrrol i done", & su 1 644 963 AI 30 Apr i l 1991 (1991-04-30) abstract</p> <p>-----</p>	1-11
A	<p>US 2008/253976 AI (SCOTT DOUGLAS CRAIG [US] ET AL) 16 October 2008 (2008-10-16) paragraph [0010] exampl e IV</p> <p>-----</p>	1-11
Y	<p>SAGOO SK ET AL: "Chi tosan potenti ates the antimi crobi al acti on of sodi um benzoate on spoi lage yeasts", LETTERS IN APPLI ED MICROBIOLOGY, WI LEY-BLACKWELL PUBLISHING LTD, GB, vol . 34, no. 3, 1 March 2002 (2002-03-01) , pages 168-172 , XP009162064, ISSN: 0266-8254, DOI : 10. 1046/J . 1472-765X. 2002 . 01067 . X [retri eved on 2002-03-01] page 168, right col umn, lines 21-24</p> <p>-----</p>	1-11
Y	<p>w0 2011/151171 AI (UNI LEVER NV [NL] ; UNI LEVER PLC [GB] ; UNI LEVER HINDUSTAN [IN] ; BARNE SA) 8 December 2011 (2011-12-08) page 2, line 30 - line 32 claim 1</p> <p>-----</p>	1-11
A	<p>w0 2010/046238 AI (UNI LEVER NV [NL] ; UNI LEVER PLC [GB] ; UNI LEVER HINDUSTAN [IN] ; CHAKRABO) 29 Apr i l 2010 (2010-04-29) cited in the appl icati on exampl es 1-33 claim 1</p> <p>-----</p>	1-11
A	<p>w0 2011/036048 AI (UNI LEVER NV [NL] ; UNI LEVER PLC [GB] ; UNI LEVER HINDUSTAN [IN] ; MEDEPALL) 31 March 2011 (2011-03-31) page 2, line 23 - line 27 exampl es 1-17 claim 1</p> <p>-----</p> <p style="text-align: center;">-/--</p>	1-11

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2012/073005

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>WO 2006/053458 AI (GIVAUDAN SA [CH] ; NATSCH ANDREAS [CH]) 26 May 2006 (2006-05-26) page 17, line 18 - line 22 claim 1</p> <p style="text-align: center;">-----</p>	1-11
Y	<p>LINA WANG ET AL: "Synergi sti c Antimi crobi al Acti vities of Natural Essenti al Oils with Chi tosan Fi lms" , JOURNAL OF AGRICULTURAL AND FOOD CHEMISTRY, vol . 59, no. 23 , 29 October 2011 (2011-10-29) , pages 12411-12419 , XP055051544, ISSN: 0021-8561 , DOI : 10. 1021/j f 203165k abstract</p> <p style="text-align: center;">-----</p>	1-11

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/EP2012/073005

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0916720	AI	19-05-1999	AR 017410 AI 05-09-2001
		AU 9756998 A 07-06-1999	
		EP 0916720 AI 19-05-1999	
		EP 1032630 AI 06-09-2000	
		JP 2001523755 A 27-11-2001	
		WO 9925800 AI 27-05-1999	

WO 2011039630	AI	07-04-2011	CA 2775929 AI 07-04-2011
		CN 102638973 A 15-08-2012	
		EP 2488016 AI 22-08-2012	
		US 2012328548 AI 27-12-2012	
		WO 2011039630 AI 07-04-2011	

SU 1644963	AI	30-04-1991	NONE

US 2008253976	AI	16-10-2008	AU 2008238839 AI 23-10 -2008
		CA 2682797 AI 23-10 -2008	
		CN 101677924 A 24-03 -2010	
		EP 2144591 A2 20-01 -2010	
		JP 2010523551 A 15-07 -2010	
		RU 2009136476 A 27-05 -2011	
		US 2008253976 AI 16-10 -2008	
		WO 2008126057 A2 23-10 -2008	

WO 2011151171	AI	08-12-2011	AR 081553 AI 03-10-2012
		CN 102905683 A 30-01-2013	
		WO 2011151171 AI 08-12-2011	

WO 2010046238	AI	29-04-2010	AR 073903 AI 09--12 -2010
		AU 2009306592 AI 29--04 -2010	
		CA 2739843 AI 29--04 -2010	
		CN 102186341 A 14--09 -2011	
		EA 201100656 AI 31--10 -2011	
		EP 2348838 AI 03--08 -2011	
		JP 2012505851 A 08--03 -2012	
		KR 20110081198 A 13--07 -2011	
		US 2011223114 AI 15--09 -2011	
		WO 2010046238 AI 29--04 -2010	

WO 2011036048	AI	31-03-2011	AR 078310 AI 26--10 -2011
		AU 2010297406 AI 05--04 -2012	
		CA 2801143 AI 31--03 -2011	
		CN 102510723 A 20--06 -2012	
		EA 201200531 AI 28--09 -2012	
		EP 2480090 AI 01--08 -2012	
		US 2012276022 AI 01--11 -2012	
		WO 2011036048 AI 31--03 -2011	

WO 2006053458	AI	26-05-2006	BR PI0517765 A 21--10 -2008
		CN 101076315 A 21--11 -2007	
		EP 1853214 AI 14--11 -2007	
		JP 2008520593 A 19--06 -2008	
		US 2008118591 AI 22--05 -2008	
		WO 2006053458 AI 26--05 -2006	
