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(54) Title: ANTIMICROBIAL COMPOSITION

(57) **Abstract**: The present invention relates to an antimicrobial composition and a method for disinfection involving the antimicrobial composition. It particularly relates to an antimicrobial composition for personal cleaning, oral care or hard surface cleaning applications. It was found that compositions comprising thymol, selected cyclic monoterpene alcohols and a carrier provide synergistic antimicrobial action. In a preferred aspect the composition also comprises 1 to 80 %-wt of one or more surfactants.



ANTIMICROBIAL COMPOSITION

FIELD OF THE INVENTION

The present invention relates to an antimicrobial composition and a method for
5 disinfection involving the antimicrobial composition. It particularly relates to an
antimicrobial composition for personal cleaning, oral care or hard surface cleaning
applications.

BACKGROUND TO THE INVENTION

10 Sanitising and disinfecting soap or cleaning compositions are of great benefit to
individuals, since proper use generally may reduce the number of germs and
pathogens the individual is exposed to. Thus, such compositions may for instance play
an important role in reducing the occurrence and spread of infectious diseases.

Compounds displaying antimicrobial, disinfecting or sanitising activity are
15 generally known, for example from "*Disinfection, Sterilisation and Preservation*", ed. S
S Block, 5th edition, Lippincott, Williams and Wilkins, Philadelphia, 2001.

Sanitising and disinfecting soap compositions comprising chlorine-based
antimicrobial agents such as triclosan are known. Such compositions require a rather
long contact time to provide efficacious antimicrobial action. In practice, users, in
20 particular children, do not spend a long time on cleansing and as a result cleaning with
such compositions does not provide adequate prevention from surface or topical
infection or adequate protection against diseases. The user, in spite of cleaning his
hands, is generally likely to end up with relatively inadequate bacterial removal from his
skin. Therefore, he may cause contamination of further animate and/or inanimate
25 surfaces and contribute to the spreading of pathogens and consequent diseases.
Users in general and children in particular who wash contaminated hands before meals
with slow-acting antimicrobial compositions for relatively short time are at risk of
contracting diseases.

Similarly in the area of hard surface cleaning, e.g. cleaning of floors, table tops or
30 utensils, the antimicrobial actives in the compositions are in contact with the substrate
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 antimicrobials commonly used in such products take
many minutes to hours to provide the desired kill of microbes.

Therefore, there is a need of providing a composition that –upon application– provides relatively more efficacious antimicrobial action during a relatively short cleaning period, preferably about 30 seconds or less.

A particular problem of thymol is that its presence in a formulation is generally well-perceptible due to its olfactory properties. Although the latter may – at least to some extent – be appreciated in certain fragrance compositions, it is considered too intense by some users when applied at concentrations efficacious in rapid disinfection. Additionally, a lower concentration of odoriferous compounds including thymol or the availability of antimicrobial compounds that are less or not odoriferous allows greater flexibility to the manufacturer in providing alternative scents to his composition at lower doses. Hence there is a need to provide alternative antimicrobial compositions and methods that preferably require lower concentrations of thymol and/or have a more acceptable sensory profile.

WO 2010/046238 A1 describes an effective antimicrobial composition which provides rapid kill of pathogenic bacteria and which comprises 0.01 to 5% by weight of thymol, 0.01 to 5% by weight of terpineol and a carrier. WO 2010/046238 A1 also discloses a method of disinfecting a surface including the step of applying the above composition to the surface.

WO 2000/21364 discloses a composition to repel or kill insects, fungi, nematodes and bacteria, comprising an essential oil or a component thereof, derived from at least one plant species in the Family *Labiatae* and *Umbellifera*. It also discloses that the major active components of such essential oils are carvacrol, thymol, cymene and anethole.

WO 2011/045557 A1 describes compositions useful for treating, preventing or ameliorating the symptoms of microbial or viral infections of the upper respiratory tract. The compositions comprising effective amounts of extracts of: (a) *Coridothymus capitatus*; (b) *Origanum dictamnus*; and (c) at least one of *Salvia fruticosa* or *Salvia pomifera*.

A. Pérez-Vásquez *et al* [J. Pharm. Pharmacol. vol 63, pp.579-586 (2011)] describe the constituents of the essential oil of *Hofmeisteria schaffneri* as determined by gas chromatography and mass spectroscopy. They also disclose the antibacterial effects of the essential oil.

Y.H. Byun and S. Shin [Natural Product Sciences, vol 14(2), pp. 138-142 (2008)] describe the constituents of the essential oils of *Chrysanthemum* species as

determined by gas chromatography and mass spectroscopy. They also disclose the antibacterial effects of the essential oils.

A. Jaafari *et al.* [Revista Brasileira de Farmacognosia, vol. 17(4), pp 477-491 (2007)] describe the constituents of the essential oils of different varieties of Moroccan
5 thyme species as determined by gas chromatography and mass spectroscopy. They also describe the cytotoxicity of the essential oils.

FR 2 697 133 discloses biocidal and/or biostatic compositions, comprising mono-oxygenated sesquiterpenes of general formula $C_{15}H_xO$, wherein x is between 20 and 26 and aromatic compounds.

10 Despite the general availability of antimicrobial compounds and compositions, there remains a continuous need to find alternative antimicrobial compositions and active compounds that are suitable for use in such compositions. In particular, alternative compositions providing fast antimicrobial action remain highly desirable in view of current consumer habits. Such alternatives may reduce the dependency on
15 current raw materials. Moreover, in the field of antimicrobials, the availability of alternatives may reduce the risk of development of microbial resistance or insensitivity to particular antimicrobial compounds.

In addition, there is a continued need to reduce the total amount of active ingredients required in such an antimicrobial composition. This need may for instance
20 be driven by the desire for cost-efficiency, because such compositions are particularly relevant to developing countries. Moreover, reducing the amounts may also be beneficial for environmental reasons.

In view of the above-observed problems and drawbacks of the prior art, it is an object of the present invention to provide alternative antimicrobial compositions.

25 It is a particular object of the invention to provide such compositions, requiring a lower dose of antimicrobial compounds.

Similarly, it is an object of the present invention to provide an antimicrobial composition in which the olfactory contribution of the antimicrobially active compounds is reduced or in which the active compound contributes to providing a consumer-
30 acceptable or even consumer-appreciated scent.

It is another particular object of the invention to provide an antimicrobial composition that contributes to reducing the required contact time in a method for disinfection of a surface.

In particular, it is an object of the invention to provide an antimicrobial composition which gives improved disinfection during cleansing of surfaces of the human body, such as the skin and the oral cavity.

It is yet another object of the present invention to provide an alternative method
5 for sanitising and/or disinfecting, in particular of surfaces.

It is a further object of the invention to provide a method for disinfection with a reduced disinfection time. More specifically, it is an object of the invention to provide a method, wherein the disinfection time of the method is less than 300 seconds, preferably less than 120 seconds, more preferably less than 60 seconds, and even
10 more preferably less than 15 seconds.

In particular, it is an object of the invention to provide a method for disinfection that gives improved disinfection during cleansing of surfaces, in particular hard surfaces, or surfaces of the human body, such as the skin and the oral cavity.

15 SUMMARY OF THE INVENTION

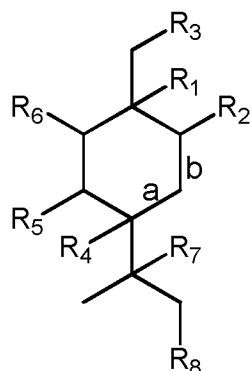
We have now found that one or more of the above objects are met by the present invention. Thus, we have found that compositions comprising selected cyclic monoterpene alcohols and thymol provide synergistic antimicrobial action. Such compositions provide similar or more efficacious anti-microbial action, at similar or
20 lower concentrations when compared to thymol and alpha-terpineol. In particular, we found that combinations of cyclic monoterpene alcohols and thymol according to this invention are capable of very fast antimicrobial action. For instance, we found that complete microbial inactivation could be effected with compositions according to the present invention after a contact time of only 15 seconds.

25 By virtue of the enhanced antimicrobial efficacy of the selected cyclic monoterpene alcohols together with thymol, the reduction of the required amount of thymol advantageously also contributes to a reduction of the olfactory disadvantages associated with the presence of larger amounts of thymol.

30 Accordingly, in a first aspect the invention provides an antimicrobial composition comprising:

- i. 0.001 to 5% by weight of thymol,
- ii. 0.001 to 5% by weight of one or more cyclic monoterpene alcohols, and
- iii. a carrier;

35 wherein the one or more cyclic monoterpene alcohols have the following structure



wherein

- one of the bonds a and b is a double bond, the other being a single bond;
- one of the substituents R₁ to R₈ is a hydroxy, the others being hydrogen;

5 subject to the proviso that

- the double bond and the hydroxy are not attached to the same carbon atom; and
- R₄ is absent when bond a is the double bond;

, and wherein the carrier comprises water, ethanol, propanol, isopropanol, ethylene glycol, propylene glycol, diethylene glycol, or mixtures thereof.

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According to a second aspect of the invention, there is provided a method of disinfecting a surface comprising the steps of

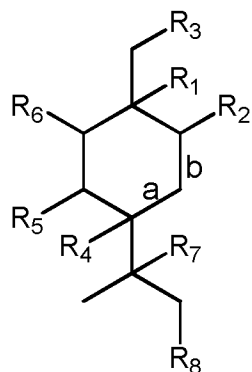
a. applying on to the surface a composition comprising

i. 0.001 to 5% by weight of thymol,

15 ii. 0.001 to 5% by weight of one or more cyclic monoterpene alcohols, and

iii. a carrier;

wherein the one or more cyclic monoterpene alcohols have the following structure



20

wherein

- one of the bonds a and b is a double bond, the other being a single bond;
- one of the substituents R₁ to R₈ is a hydroxy, the others being hydrogen;

subject to the proviso that

- the double bond and the hydroxy are not attached to the same carbon atom;
and

- R_4 is absent when bond a is the double bond; and

5 b. removing the composition from the surface.

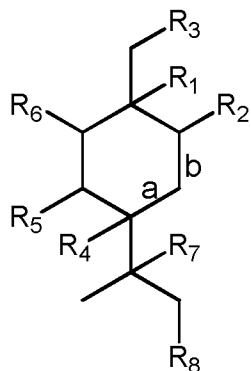
It is particularly preferred that the composition used in the method according to the second aspect of the invention is a composition according to the first aspect of the invention.

10

In a third aspect, the invention provides the use of a composition for improved hygiene, wherein the composition comprises

- 0.001 to 5% by weight of thymol,
- 0.001 to 5% by weight of one or more cyclic monoterpene alcohols, and
- 15 iii. a carrier;

wherein the one or more cyclic monoterpene alcohols have the following structure



wherein

20 • one of the bonds a and b is a double bond, the other being a single bond;

- one of the substituents R_1 to R_8 is a hydroxy, the others being hydrogen;

subject to the proviso that

- the double bond and the hydroxy are not attached to the same carbon atom;
and

25 • R_4 is absent when bond a is the double bond.

DETAILED DESCRIPTION OF THE INVENTION

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." Thus, the term
5 "comprising" is meant not to be limiting to any subsequently stated elements but rather to optionally also encompass non-specified elements of major or minor functional importance. In other words, the listed steps or options need not be exhaustive. Whenever the words "including" or "having" are used, these terms are meant to be equivalent to "comprising" as defined above. It is noted that the examples given in the
10 description below are intended to clarify the invention and are not intended to limit the invention to those examples per se.

Except in the examples, or where otherwise explicitly indicated, all numbers in this description indicating amounts of material or conditions of reaction, physical properties of materials and/or use are to be understood as modified by the word
15 "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.

Throughout this description, the term disinfection refers to reduction of the
20 number of viable microorganisms in a given medium or on a given surface by physical or chemical means. Typically, disinfection involves the destruction or inactivation of said microorganisms. Both animate and inanimate media and surfaces are contemplated.

The term "microbicide" refers to a compound capable of killing, inhibiting the
25 growth of or controlling the growth of microorganisms at a locus; microbicides include bactericides, fungicides and algacides. The term "microorganism" includes, for example, fungi (including yeast and mould), bacteria and algae.)

The antimicrobial composition comprises thymol, one or more cyclic
monoterpene alcohols, and a carrier. Various components of the antimicrobial
30 composition are described below.

The compositions of the present invention are preferred for non-therapeutic use, and more particularly preferred for use in cleaning surfaces of human body including skin, hair or oral cavity or for hard surface cleaning applications.

35 *Cyclic monoterpene alcohols*

The antimicrobial composition according to the invention comprises 0.001 to 5% by weight of one or more cyclic monoterpene alcohols. The composition comprises preferably 0.005 to 4.5 wt-%, more preferably 0.01 to 4 wt-%, even more preferably 0.02 to 3 wt-%, yet more preferably 0.03 to 2 wt-%, still more preferably 0.04 to 1 wt-%, 5 even more preferably 0.05 to 0.75 wt-% and still more preferably 0.1 to 0.5 wt-% of the one or more cyclic monoterpene alcohols. In compositions intended to be diluted before application, the minimum preferred concentrations of the one or more cyclic monoterpene alcohols can be higher. For example, when washing hands with water and a composition according to the invention, the lather produced, typically is a 50 wt% 10 dilution of the original composition. Similarly, in body wash situations, soap bars or soap liquids are typically diluted until about 8 wt% soap in water, corresponding to an approximately tenfold dilution of the product. Therefore, compositions according to the invention intended for dilution upon use preferably comprise 0.05 to 4.5 wt-%, more preferably 0.1 to 4 wt-%, even more preferably 0.2 to 3 wt-%, still more preferably 0.4 15 to 1 wt-%, and still more preferably 0.5 to 1 wt-% of the one or more cyclic monoterpene alcohols. Thus, the concentration of the one or more cyclic monoterpene alcohols in the antimicrobial composition is preferably such that, when the composition is diluted or dissolved with a suitable medium during use, the concentration in the diluted or dissolved mixture is still sufficient to be antimicrobially efficacious.

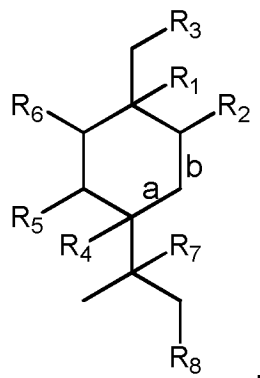
20 The cyclic monoterpene alcohol can be a single compound or can be a mixture of the cyclic monoterpene alcohols as detailed below. In certain preferred embodiments, mixtures of cyclic monoterpene alcohols are preferred, since such mixtures may show increased antimicrobial activity against a wider range of microbes. On the other hand, for reasons including e.g. control over the formulation, it is preferred 25 that in case the composition according to the invention comprises a mixture of such cyclic monoterpene alcohols, the mixture preferably comprises at least 30%, more preferably at least 50%, even more preferably at least 70% and still more preferably at least 90% by weight of one cyclic monoterpene alcohol with respect to the total weight of the cyclic monoterpene alcohols.

30 At concentration ranges of the cyclic monoterpene alcohols below their lower concentration limits, the desired fast acting antimicrobial kinetics in combination with thymol would not be met. At concentrations higher than the higher preferred concentrations of the cyclic monoterpene alcohols, when in combination with thymol, while the kinetics of action would not be compromised, the present inventors have 35 found that unlike in therapeutic/pesticidal/herbicidal applications where sensorial

aspects are not critical, in the present invention, which preferably relates to personal cleaning, oral care or hard surface cleaning applications, the product is in contact with hands, mouth or other body parts, the sensorial aspects including smell and skin feel would be compromised.

5

The one or more cyclic monoterpene alcohols have the following structure



wherein

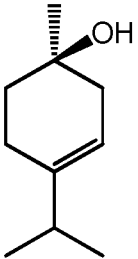
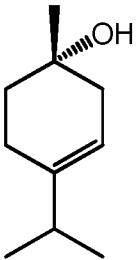
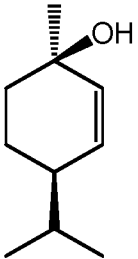
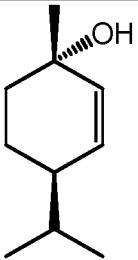
- one of the bonds a and b is a double bond, the other being a single bond;
 - 10 • one of the substituents R₁ to R₈ is a hydroxy, the others being hydrogen; subject to the proviso that
 - the double bond and the hydroxy are not attached to the same carbon atom; and
 - R₄ is absent when bond a is the double bond.
- 15 Thus, the cyclic monoterpene alcohols are selected from the group consisting of 4-isopropyl-1-methylcyclohex-3-enols, (4-isopropylcyclohex-3-en-1-yl)methanols, 2-(4-methylcyclohex-1-en-1-yl)propan-1-ols, 2-(4-methylcyclohex-1-en-1-yl)propan-2-ols, 2-isopropyl-5-methylcyclohex-2-enols, 3-isopropyl-6-methylcyclohex-2-enols, 3-isopropyl-6-methylcyclohex-3-enols, 4-isopropyl-1-methylcyclohex-2-enols, (4-isopropylcyclohex-2-en-1-yl)methanols, 1-isopropyl-4-methylcyclohex-2-enols, 2-(4-methylcyclohex-2-en-1-yl)propan-1-ols, 2-(4-methylcyclohex-2-en-1-yl)propan-2-ols, 2-isopropyl-5-methylcyclohex-3-enols, and 5-isopropyl-2-methylcyclohex-3-enols.

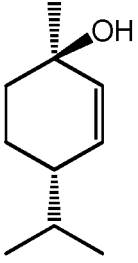
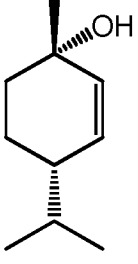
All stereoisomers of the cyclic monoterpene alcohols according to the invention are contemplated. For example, if a cyclic monoterpene alcohol according to the invention comprises exactly one stereocentre (for example 4-isopropyl-1-methylcyclohex-3-enol), both enantiomers are contemplated. Analogously, if a cyclic monoterpene alcohol according to the invention comprises for instance exactly two stereocentres (for instance 4-isopropyl-1-methylcyclohex-2-enol), there are two sets of

two enantiomers, whereby the members of one set are diastereomers with respect to the members of the other set. All four of these stereoisomers are contemplated. Thus, compositions comprising enantiomerically pure compounds, racemic mixtures and other mixtures of different stereoisomers are equally preferable, unless specified
5 otherwise in the below.

By means of a non-limiting example, the stereoisomers of 4-isopropyl-1-methylcyclohex-3-enol and 4-isopropyl-1-methylcyclohex-2-enol are provided in Table 1.

Table 1

(<i>R</i>)-4-isopropyl-1-methylcyclohex-3-enol	
(<i>S</i>)-4-isopropyl-1-methylcyclohex-3-enol	
(1 <i>R</i> ,4 <i>R</i>)-4-isopropyl-1-methylcyclohex-2-enol	
(1 <i>S</i> ,4 <i>R</i>)-4-isopropyl-1-methylcyclohex-2-enol	

(1 <i>R</i> ,4 <i>S</i>)-4-isopropyl-1-methylcyclohex-2-enol	
(1 <i>S</i> ,4 <i>S</i>)-4-isopropyl-1-methylcyclohex-2-enol	

In case the carrier or the dissolution medium during later application is water-based, it can be advantageous if the one or more cyclic monoterpene alcohols are sufficiently water-soluble. The cyclic monoterpene alcohols are sufficiently water-soluble if they
5 are soluble to at least the minimum concentration required in the antimicrobial composition according to the invention.

Preferably, the one or more cyclic monoterpene alcohols are selected from the group consisting of 4-isopropyl-1-methylcyclohex-3-enols, 3-isopropyl-6-methylcyclohex-2-enols, and 4-isopropyl-1-methylcyclohex-2-enols. More preferably,
10 the one or more cyclic monoterpene alcohols are selected from the group consisting of 4-isopropyl-1-methylcyclohex-3-enols and 4-isopropyl-1-methylcyclohex-2-enols. Even more preferably, the cyclic monoterpene alcohol is 4-isopropyl-1-methylcyclohex-3-enol.

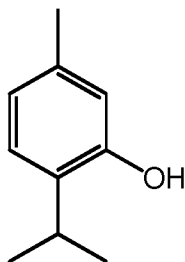
Mixtures of preferred cyclic monoterpene alcohols are also preferred.

15 Without wishing to be bound by theory, it is believed that the synergistic mode of antimicrobial action of the one or more cyclic monoterpene alcohols according to the present invention in combination with thymol is similar for the different respective cyclic monoterpene alcohols.

Suitable cyclic monoterpene alcohols according to the present invention may be
20 commercially sourced, obtained from natural sources or obtained via synthetic chemical methods. Such methods are generally well-known to the person skilled in the art.

Thymol

Thymol has the following structure:



Thymol is also known as 2-isopropyl-5-methylphenol.

5

The antimicrobial composition according to the invention comprises 0.001 to 5% by weight of thymol. The composition comprises preferably 0.005 to 4.5 wt-%, more preferably 0.01 to 4 wt-%, even more preferably 0.02 to 3 wt-%, yet more preferably 0.03 to 2 wt-%, still more preferably 0.04 to 1 wt-%, still more preferably 0.05 to 0.75 wt-% and even more preferably 0.1 to 0.5 wt-% of thymol. In compositions intended to be diluted before application, the minimum preferred concentrations of the thymol can be higher, for the same reasons as for the cyclic monoterpene alcohols. Therefore, compositions according to the invention intended for dilution upon use preferably comprise 0.05 to 4.5 wt-%, more preferably 0.1 to 4 wt-%, even more preferably 0.2 to 3 wt-%, still more preferably 0.4 to 1 wt-%, and still more preferably 0.5 to 1 wt-% of the thymol. Any of the concentrations ranges for the thymol is preferably combined with any of the concentration ranges for the one or more cyclic monoterpene alcohols specified above. Therefore, the antimicrobial composition according to the invention for example comprises:

- 20 a. 0.01 to 0.4% by weight of thymol;
 b. 0.05 to 1% by weight of the one or more cyclic monoterpene alcohols.

The preferred concentrations ranges of the thymol are important for the same reasons as the preferred concentration ranges of the one or more cyclic monoterpene alcohols in meeting the desired fast acting antimicrobial kinetics while not being sensorially unpleasant when used in products for personal cleaning, oral care or hard surface cleaning applications.

Thymol may be added to the antimicrobial composition in purified form. However, thymol is compound that is naturally present in many plant species. Therefore, alternatively, plant oils or plant extracts derived from such thymol-containing plant species can be added to the antimicrobial composition, while ensuring that thymol

30

is present in the desired concentration in the composition of the present invention. For example, thyme oil or thyme extract naturally comprise thymol. Thyme oil or thyme extract is obtained from the thyme plant. Thyme plant refers to a plant belonging to the genus *Thymus* and includes but is not limited to the following species:

- 5 *vulgaris*, *Thymus zygis*, *Thymus saturoides*, *Thymus mastichina*, *Thymus broussonetti*, *Thymus maroccanus*, *Thymus pallidus*, *Thymus algeriensis*, *Thymus serpyllum*, *Thymus pulegoide*, and *Thymus citriodorus*.

Carrier

- 10 The antimicrobial composition according to the invention comprises a carrier. The carrier is preferably selected from the group consisting of water, oil, solvent, inorganic particulate material, starch, air and mixtures thereof. The carrier of the composition according to the first aspect of the invention comprises water, ethanol, propanol, isopropanol, ethylene glycol, propylene glycol, diethylene glycol, or mixtures thereof.
- 15 The carrier is preferably from 0.1 to 99% by weight of the composition. The antimicrobial composition may be in form of a solid, liquid, gel, paste or soft solid and the carrier may be selected by a person skilled in the art depending on the format of the antimicrobial composition.

Examples of inorganic particulate materials include clay, talc, calcite, dolomite, silica, and aluminosilicate. Examples of oils include mineral oils, oils of biological origin (e.g. vegetable oils), and petroleum-derived oils and waxes. The oils of biological origin are preferably triglyceride-based. Preferably, the carrier oil is not a perfume oil. Thus, the carrier oil preferably does not substantially contribute to the odour of the composition, more preferably it does not contribute to that odour. Examples of solvents

25 include alcohols, ethers and acetone. The starch may be natural starch obtained from food grains or may be a modified starch.

In certain preferred embodiments, suitable solvents include, for example, water; glycols, including ethylene glycol, propylene glycol, diethylene glycol, dipropylene glycol, polyethylene glycol, and polypropylene glycol; glycol ethers; alcohols, such as

30 methanol, ethanol, propanol, phenethyl alcohol and phenoxypropanol; ketones, including acetone and methyl ethyl ketone; esters, including ethyl acetate, butyl acetate, triacetyl citrate, and glycerol triacetate; carbonates, including propylene carbonate and dimethyl carbonate; and mixtures thereof. It is preferred that the solvent is selected from water, glycols, glycol ethers, esters and mixtures thereof. In certain

35 preferred embodiments, suitable solid carriers include, for example, cyclodextrin,

silicas, diatomaceous earth, waxes, cellulosic materials, alkali and alkaline earth (e.g., sodium, magnesium, potassium) metal salts (e.g., chloride, nitrate, bromide, sulfate) and charcoal.

Air can for instance be used as a carrier when the cyclic monoterpene alcohols
5 according to the invention and/or the thymol are atomised or otherwise dispersed as a fine mist.

Particularly preferred carriers are water or oil/solvent and even more preferred is a carrier that is a mixture of water and oil. Thus, in many of the envisaged applications like personal care/washing, oral care and hard surface cleaning, the
10 antimicrobial composition may be formulated with either an aqueous base or a oil/solvent base. Compositions with an aqueous base (water being the carrier), may also for instance be products in gel format. Compositions with a purely oil/solvent base may for instance be products in anhydrous stick form or propellant-containing products.

Thus, the antimicrobial composition may for instance, preferably be an
15 antimicrobial anhydrous stick personal care composition on a purely oil/solvent base wherein the composition has a water content of less than 0.01% by weight, and wherein the composition preferably is free of water. Alternatively, the antimicrobial composition may for instance, preferably be an antimicrobial propellant-drivable personal care composition, also comprising a propellant. Air can also be used as
20 propellant, for instance in the form of compressed or liquefied air.

However, the most preferred product format has an emulsion base (water and/or oil being the carrier) or is capable of forming an emulsion upon dilution, e.g. soap products in liquid, solid, lotion or semisolid form for hand wash, face wash, body wash, or shaving applications; toothpaste/ dentifrices for oral care applications or
25 products for hard surface cleaning in bars or liquids form. If the product comprises an emulsion base, it preferably also comprises one or more surfactants as described below.

Surfactants

30 The antimicrobial composition according to the invention preferably comprises from 1 to 80% by weight of one or more surfactants. Surfactants may for instance advantageously contribute to the cleaning efficacy or the formulation stability of a composition.

Thus, the antimicrobial composition according to the invention preferably
35 comprises

- a. 0.001 to 5% by weight of thymol,
- b. 0.001 to 5% by weight of the one or more cyclic monoterpene alcohols according to the invention,
- c. a carrier, and
- 5 d. from 1 to 80% by weight of one or more surfactants.

It is particularly preferred that the antimicrobial composition comprises from 1 to 80% by weight of one or more surfactants in combination with the one or more cyclic monoterpene alcohols, and the thymol at their more preferred concentrations as
10 specified above.

In general, the surfactants may be chosen from the surfactants described in well-known textbooks like "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
15 Manufacturing Confectioners Company or in "Tenside-Taschenbuch", H. Stache, 2nd Edn., Carl Hauser Verlag, 1981; "Handbook of Industrial Surfactants" (4th Edn.) by Michael Ash and Irene Ash; Synapse Information Resources, 2008. Any type of surfactant, i.e. anionic, cationic, nonionic, zwitterionic or amphoteric can be used. Preferably, the one or more surfactants are anionic, nonionic, or a combination of
20 anionic and nonionic surfactants. More preferably, the one or more surfactants are anionic.

A particularly preferred surfactant is soap. Soap is a suitable surfactant for personal washing applications of the antimicrobial composition of the invention. The soap is preferably C₈-C₂₄ soap, more preferably a C₁₀-C₂₀ soap and most preferably
25 C₁₂-C₁₆ soap. The soap may or may not have one or more carbon-carbon double bonds or triple bonds. The cation of the soap can for instance be an alkali metal, alkaline earth metal or ammonium. Preferably, the cation of the soap is selected from sodium, potassium or ammonium. More preferably the cation of the soap is sodium or potassium.

30 *The soap may be obtained by saponifying a fat and/or a fatty acid. The fats or oils may be fats or oils generally used in soap manufacture, such as tallow, tallow stearines, palm oil, palm stearines, soya bean oil, fish oil, castor oil, rice bran oil, sunflower oil, coconut oil, babassu oil, palm kernel oil, and others. In the above process the fatty acids are derived from oils/fats selected from coconut, rice bran,*
35 *groundnut, tallow, palm, palm kernel, cotton seed, soyabean, castor etc. The fatty acid*

soaps can also be synthetically prepared (e.g. by the oxidation of petroleum or by the hydrogenation of carbon monoxide by the Fischer-Tropsch process). Resin acids, such as those present in tall oil, may be used. Naphthenic acids are also suitable.

Tallow fatty acids can be derived from various animal sources. Other similar
5 mixtures, such as those from palm oil and those derived from various animal tallow and lard are also included.

A typical fatty acid blend consists of 5 to 30%-wt coconut fatty acids and 70 to 95%-wt fatty acids ex hardened rice bran oil. Fatty acids derived from other suitable oils/fats such as groundnut, soybean, tallow, palm, palm kernel, etc. may also be used
10 in other desired proportions. The soap, when present in solid forms of the present invention, is preferably present in an amount of 30 to 80%, more preferably from 50 to 80%, and even more preferably 55 to 75% by weight of the composition. The soap, when present in liquid forms of the composition is preferably present in 0.5 to 20%, more preferably from 1 to 10% by weight of the composition.

15 Other preferred surfactants are fatty acid glycinates and fatty amphocarboxylates. The fatty acid glycinates are fatty acid amides or salts of glycine, including for example sodium cocoyl glycinate. The fatty amphocarboxylates are amphoteric surfactants including for example sodium lauroamphoacetate (i.e. sodium 2-[1-(2-hydroxyethyl)-2-undecyl-4,5-dihydroimidazol-1-ium-1-yl]acetate). Yet another
20 example of suitable surfactants are derivatives of isethionates, including acylisethionates.

The antimicrobial composition of the invention is also 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
25 between 1 and 8 ethylene oxide groups when the product is in the liquid form. When the product for hard surface cleaning applications is in the solid form, surfactants are preferably selected from primary alkyl sulphates, secondary alkyl sulphonates, alkyl benzene sulphonates, ethoxylated alkyl sulphates, or alcohol ethoxylate nonionic surfactants. The composition may further comprise an anionic surfactant, such as alkyl
30 ether sulphate preferably those having between 1 and 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. Other classes of useful surfactants include cationic surfactants, such as long chain quaternary
35 ammonium compounds and amphoteric surfactants such as betaines and alkyl

dimethyl amine oxides. 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 antimicrobial composition of the invention is useful in oral care compositions e.g. in a dentifrice/ toothpaste or an oral rinse product. In such applications, preferred surfactants are anionic, nonionic or amphoteric in nature, preferably anionic or amphoteric. The anionic surfactant is preferably an alkali metal alkyl sulphate, more preferably a sodium lauryl sulphate (SLS). Mixtures of anionic
10 surfactants may also be employed. The 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
15 2.2% to about 10%, more preferably from about 2.5 to about 5% by weight of the total composition.

Thus, it is highly preferred that the antimicrobial compositions include soap, alkyl sulphate or linear alkyl benzene sulphonate as the surfactants. More preferably, the one or more surfactants are selected from the group consisting of soaps, alkyl
20 sulphates and linear alkyl benzene sulphonates.

Liquid and solid compositions

The antimicrobial composition may be in form of a solid, a liquid, a gel or a paste. A person skilled in the art can prepare compositions in various formats by choosing one
25 or more carrier materials and/or surfactant. The antimicrobial compositions of the present invention are useful for cleansing and care, in particular for skin cleansing and skin care. It is envisaged that the antimicrobial composition can be used as a leave-on product or a wash-off product, preferably a wash-off product. The antimicrobial composition of the present invention can also be used for cleansing and care of hard
30 surfaces such as glass, metal, plastic and the like.

A particularly preferred carrier is water. When water is the carrier, both liquid and solid compositions are possible. Different amounts of water may be preferred depending on the product format. 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. When water is the carrier, a preferred liquid antimicrobial composition according to the invention comprises:

- a. 0.01 to 5% by weight of thymol,
- b. 0.05 to 5% by weight of the one or more cyclic monoterpene alcohols,
- 5 c. 10 to 99.9% by weight of water, and;
- d. 1 to 30% by weight of the one or more surfactants.

The liquid antimicrobial composition is useful for skin cleansing, in particular for hand wash or a face wash.

10

When water is the carrier, a preferred solid antimicrobial composition according to the invention comprises:

- a. 0.05 to 5% by weight of the thymol,
- b. 0.05 to 5% by weight of the one or more cyclic monoterpene alcohols,
- 15 c. 5 to 30% by weight of water, and;
- d. 30 to 80% by weight of surfactant.

The solid antimicrobial composition is preferably in form of a shaped solid, more preferably a bar. The solid antimicrobial composition is particularly useful for skin cleansing in particular for hand wash or a face wash.

- 20 Such a bar-shaped solid antimicrobial composition may for instance be a soap bar. Soap bar compositions are well-known and may be similar to the following non-limiting example composition, comprising 75.6 wt-% of anhydrous sodium soap, 1.0 wt-% of glycerine, 0.5 wt-% of sodium carbonate, 0.2 wt-% of EHDP (ethane-1-hydroxy-1,1-disphosphonate) acid, 0.04 wt-% of EDTA (ethylenediaminetetraacetic acid)
- 25 tetrasodium salt, 8.5 wt-% of hydrated magnesium silicate (Talc), 0.7 wt-% of sodium chloride, 0.05 wt-% of dyes, 0.75 wt-% perfume, 0.05 to 10 wt-% of antimicrobial agents including the cyclic monoterpene alcohols and the thymol according to the present invention, and water up to 100 wt-%.

- Alternatively, inorganic particulate material is also a suitable carrier. When
- 30 inorganic particulate material is the carrier, the antimicrobial composition is in a solid form. Preferably the inorganic particulate material is talc. When the inorganic particulate material is talc, the solid antimicrobial composition is particularly useful as a talcum powder for application on face or body.

- According to another alternative, a solvent different from water is a preferred
- 35 carrier. Although any solvent can be used, alcohol is a preferred solvent. Short chain

alcohols –in particular ethanol, propanol, and isopropanol– are particularly preferred as carrier for an antimicrobial wipe or an antimicrobial hand sanitiser composition.

Solvents like ethanol and isopropanol generally show antimicrobial efficacy themselves. However, they are also volatile and may readily evaporate during
5 application of the composition. Thus, their levels on the surface that is treated might even reduce until below the minimum level required for antimicrobial action, before the minimum period needed for disinfection has passed. In contrast, the thymol and the one or more cyclic monoterpene alcohols according to the present invention are much less volatile and may therefore yield prolonged antimicrobial action after applying them
10 to the skin.

Additional ingredients

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:
15 perfumes, pigments, preservative, emollients, sunscreens, emulsifiers, gelling agents, thickening agents, humectants (e.g. glycerine, sorbitol), sequestrants (e.g. EDTA) or polymers (e.g. cellulose derivatives for structuring such as methyl cellulose)

Both thymol and some of the cyclic monoterpene alcohols according to the invention may contribute to the olfactory properties of the composition. Although some
20 of these compounds might be applied for instance in perfume compositions, the present invention is directed towards antimicrobial compositions. Therefore, the composition is preferably not a perfume composition, although other perfume components can be present. Here, a perfume composition is defined as a composition comprising a plurality of olfactory components, wherein the composition is solely
25 intended to provide a harmonious scent.

Synergistic effect of the invention

The inventors have surprisingly found that while one or more of the cyclic monoterpene alcohols according to the present invention alone or thymol alone do not individually
30 provide the fast antimicrobial kinetic action, a combination of one or more cyclic monoterpene alcohols and thymol at the selective concentrations provides a synergistic antimicrobial action which is especially important in a wash off processes where the contact time of the antimicrobial actives with the surface is low, i.e. of the order of less than 5 minutes, preferably less than 2 minutes, further more preferably less than a
35 minute and in many cases less than 15 seconds.

Synergistic combinations of cyclic monoterpene alcohols and thymol

The antimicrobial action of two or more active compounds is considered additive if the combined action merely results from the addition of the effects the individual

5 components would have in isolation. In contrast, the antimicrobial action of two or more active compounds is considered to be synergistic if the combined effect of the two or more compounds is stronger than expected based on the assumption of additivity. Without wishing to be bound by theory, it is believed that the antimicrobial action of the one compound may be enhanced by the action of the other compound and vice versa.

10 Such enhancement may for instance originate from cooperative interplay between the mechanisms of antimicrobial action at the molecular level. Such enhanced antimicrobial action may manifest itself for instance by the fact that lower concentrations of active compounds are required to obtain complete microbial kill, or alternatively, that the same extent of microbial kill is arrived at in a shorter time.

15 Whether an antimicrobial composition comprising two or more active compounds is capable of synergistic antimicrobial action may for instance be determined following the procedures and using the criteria as outlined in Example 1 below. Typically, evidence of synergistic antimicrobial action may be provided at concentrations below the minimum biocidal concentrations of each of the components when taken individually.

20 However, it is generally believed that synergistic action can still occur when the concentration of one or more of the active compounds is raised above its minimum biocidal concentration (when taken individually).

The antimicrobial composition according to the present invention preferably comprises the one or more cyclic monoterpene alcohols and thymol according to the

25 invention at concentrations at which they are capable of synergistic antimicrobial action. Thus, the concentrations of the one or more cyclic monoterpene alcohols and of the thymol in the antimicrobial composition are preferably such that, when the composition is diluted or dissolved with a suitable medium during use, (e.g. when washing hands with water and a composition according to the invention) the

30 concentration in the diluted or dissolved mixture is still sufficient to be antimicrobially efficacious. That is, to be capable of synergistic antimicrobial action, the concentrations of the one or more cyclic monoterpene alcohols and the thymol in the composition ($C_{\text{comp,alc}}$ and $C_{\text{comp,thymol}}$ respectively) are preferably such that upon application, at a given concentration of the one or more cyclic monoterpene alcohols in the application

35 medium ($C_{\text{med,alc}}$), the thymol is available at at least a minimum medium concentration

($C_{\text{med,thymol}}$) or vice versa (i.e. such that at a given $C_{\text{med,thymol}}$, a minimum $C_{\text{med,alc}}$ is available, sufficient to provide synergistic antimicrobial action). Here, the application medium denotes the medium in which the antimicrobial action desirably takes place. For example, in personal care applications like hand-washing, the composition may be a solid soap bar. In that case, C_{comp} refers to the concentration of the component in the soap bar, whereas C_{med} refers to the concentration in the lather. The minimum and optimum concentrations may for instance be determined by a protocol as described for Example 1 or by one of the standards as detailed below. It is generally preferred that the concentrations of the one or more cyclic monoterpene alcohols and the thymol in the composition according to the invention are equal to or higher than the optimal concentrations in the application medium, because in many typical applications, the composition is either used pure or is diluted to form the application medium.

Surprisingly, the synergy between the cyclic monoterpene alcohols and thymol in compositions according to the invention, occurs over a wide range of concentrations and concentration ratios. Depending on factors including the type of antimicrobial composition, its intended application (for instance a hard surface cleaner, a skin cleanser, or a hand sanitiser) different concentration ranges and ratios will be preferred.

Thus, when for instance antimicrobial action against *E. Coli* is desired, the data of Example 1 may be used to determine preferable medium concentrations C_{med} . For example, in case complete microbial inactivation is desired in the particular medium of Example 1 and if the cyclic monoterpene alcohol is 4-isopropyl-1-methylcyclohex-3-enol, and $C_{\text{med,alc}}$ is selected as 0.112 % (w/v), $C_{\text{med,thymol}}$ preferably is at least 0.025 % (w/v) and vice versa.

Alternatively, the desired antimicrobial effect may be obtained by selecting a ratio of the respective concentrations of the one or more cyclic monoterpene alcohols and the thymol. In view of the above-described considerations regarding the intended antimicrobial efficacy and other considerations, including for instance sensory properties, solubility, economic considerations, a concentration ratio of thymol to cyclic monoterpene alcohols larger than one is preferred in some applications, whereas a concentration ratio of thymol to the cyclic monoterpene alcohols smaller than one is preferred in others, whereby the concentrations are expressed in %-wt.

Thus, in case a concentration ratio of thymol to cyclic monoterpene alcohols smaller than one is desired, then the antimicrobial composition according to the invention preferably comprises the one or more cyclic monoterpene alcohols and

thymol in a concentration ratio (thymol : cyclic monoterpene alcohols) of between 1:2 and 1:12, wherein the concentration is expressed as weight percent.

A further additional advantage of the present invention is that it is observed that treatment of a surface with a composition according to the invention comprising one or
5 more cyclic monoterpene alcohols and thymol, surprisingly enables continued protection of the surface against growth of microbes for a substantial period of time thereafter.

Effect of including surfactant

10 Favourably, compositions suitable in wash-off processes as described above include a surfactant for the cleaning action. To the further surprise of the inventors, while the surfactant alone does not provide the fast antimicrobial kill at the concentration present in wash off processes, it provides for further improvement in extent of reduction in
15 viable microbial counts on the surface in the short period of time when surfaces are washed with a composition comprising one or more cyclic monoterpene alcohols, thymol and additionally surfactant. Thus, while surfactant is generally known to be responsible for washing off dirt and also antimicrobial actives used in the composition, in the present invention, it provides a highly useful additional benefit in that it enhances the reduction of viable microbial count in a composition comprising a combination of
20 cyclic monoterpene alcohols and thymol alone.

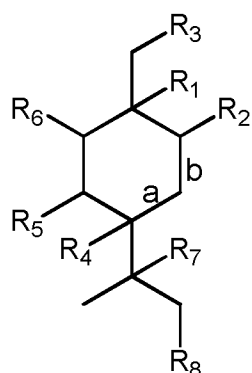
However, it was surprisingly found that certain surfactants may reduce the activity of antimicrobial agents. This may happen for instance with cocoyl glycinate and lauroamphoacetate. Generally, surfactants are required in cleaning compositions to obtain good cleaning results. Since it is *inter alia* an object of this invention to provide
25 antimicrobial cleaning compositions, it therefore also is an object of this invention to provide cyclic monoterpene alcohols capable of enhanced antimicrobial action upon combination with thymol in the presence of such surfactants. It was found that in particular 4-isopropyl-1-methyl-cyclohex-3-enol shows enhanced antimicrobial action in combination with thymol. Therefore the composition according to the invention
30 preferably comprises 4-isopropyl-1-methyl-cyclohex-3-enol. More specifically, it is preferred that in case the composition according to the invention comprises a surfactant selected from cocoyl glycinate and lauroamphoacetate, the cyclic monoterpene alcohols comprise 4-isopropyl-1-methyl-cyclohex-3-enol.

Method according to the invention

According to the second aspect, the invention relates to a method of disinfecting a surface comprising the steps of

- a. applying on to the surface a composition comprising
- i. 0.001 to 5% by weight of thymol,
 - ii. 0.001 to 5% by weight of one or more cyclic monoterpene alcohols, and
 - iii. a carrier;

wherein the one or more cyclic monoterpene alcohols have the following structure



wherein

- one of the bonds a and b is a double bond, the other being a single bond;
 - one of the substituents R_1 to R_8 is a hydroxy, the others being hydrogen;
- subject to the proviso that

- the double bond and the hydroxy are not attached to the same carbon atom;
- and
- R_4 is absent when bond a is the double bond; and

- b. removing the composition from the surface.

Preferably, the surface is skin. Thus, for example, a surface like the hands, face, body, or the oral cavity is contacted with the composition of the invention. If the surface is a surface of a human or animal body, the method preferably is a non-therapeutic method of disinfecting a surface. Alternatively, the surface is any hard surface. Typically, such hard surfaces are surfaces that commonly require cleaning and preferably also require sanitisation or disinfection. Such surfaces may be found in many household or industrial environments, and may include for example kitchen and bathroom surfaces, table tops, floors, walls, windows, utensils, cutlery, and crockery. Such surfaces may

be made from many different materials, including for instance plastics, wood, metal, ceramics, glass, concrete, marble, and painted surfaces.

The composition may be applied to the surface by any suitable means known to the skilled person. For instance, a suitable means may be pouring, dropping, spraying
5 or wiping in case of liquid compositions.

The composition that is applied in the method according to the second aspect of the invention is described above. It is particularly preferred that the composition is a composition according to the first aspect of the invention.

Preferably, the method includes diluting or dissolving the composition with a
10 suitable solvent, preferably water, before or whilst applying the composition to the surface. Such dissolving is preferred in particular in case the composition is a solid composition. Alternatively, solid compositions may also be directly spread, rubbed, or sprayed, e.g. in the form of a powder.

The method according to the second aspect of the present invention also
15 includes the step of removing the composition from the surface. Here, removing the composition also encompasses partially removing the composition, because traces of the composition may remain on the surface. In many typical situations, including washing of the skin or hard-surface cleaning, it is acceptable or sometimes even desirable if part of the composition – in particular certain active ingredients – remains
20 on the surface. Therefore, step b preferably involves removing at least 5%, more preferably at least 10%, even more preferably at least 25%, still more preferably at least 50% and yet more preferably at least 75% of the composition by weight. Preferably, the step of removing the composition comprises rinsing the surface with a suitable solvent or wiping the surface with a suitable wipe, more preferably, this step
25 consists of rinsing the surface with a suitable solvent or wiping the surface with a suitable wipe. Alternatively, the removal step can also include evaporation of part of the composition, for example when the composition comprises volatile components, e.g. solvents.

A suitable medium for rinsing the surface is water but it could also be for
30 example a mixture of water and alcohol. It is then rinsed preferably with sufficient amounts of water after a pre-determined period of time to remove any visible or sensory residue of the composition. Alternatively, an alcohol wipe or a water/alcohol impregnated wipe may be used to wipe the surface to be visibly free of the anti-microbial composition. The step of removing the composition (e.g. by rinsing or wiping
35 the surface) is preferably started 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 yet more preferably less than 20 seconds after commencement of the step of applying the composition on the surface, because of the surprisingly fast antimicrobial action of the compositions according to the present invention. Even
5 though partial microbial kill may be almost instantaneous upon application of the composition according to the invention, it is preferred that the step of removing the composition from the surface is started out at least 5 seconds, preferably at least 10 seconds, more preferably at least 15 seconds after commencement of the step of applying the composition on the surface, in order to effect optimal antimicrobial action.
10 Combinations of these times into time intervals are preferred too. Therefore, it is particularly preferred that the step of removing the composition from the surface (i.e. step b) is started between 2 minutes and 5 seconds, more preferably between 1 minute and 10 seconds, even more preferably between 30 and 10 seconds and still more preferably between 20 and 15 seconds after commencement of the step of applying the
15 composition on the surface (i.e. step a).

Disinfection time

These times between applying the composition and rinsing or wiping are preferably related to the disinfection time, in order to ensure optimal cleansing and sanitising of
20 the surface. Therefore, the invention preferably relates to a method, wherein the disinfection time T of said method is less than 300 seconds, preferably less than 120 seconds, more preferably less than 60 seconds, and even more preferably less than 15 seconds; wherein T is defined as the time that elapses from the moment of adding the composition to a microbial culture until the number of microbes per unit volume of the
25 culture is reduced by a factor of 100 000; and wherein the initial number of microbes preferably exceeds about 100 000 000 microbes per millilitre and wherein the composition is preferably a liquid composition.

The disinfecting action of the method (as may be expressed in terms of the disinfection time T) is preferably determined according to the protocol of Example 1 as
30 described hereinafter. This test relates to a standardised test environment in which the microbial culture is kept in suspension. A similarly suitable test is the standard suspension method described in European Standard EN1276, with the proviso that the disinfection time is adapted to suit the above criteria as will be clear to a person skilled in the art. Alternatively, one of the test methods as described in WO 2010/046238 may
35 for instance be applied to establish the disinfecting action.

Such test methods may preferably also be used by the skilled person to determine the optimal concentrations of the one or more cyclic monoterpene alcohols and the thymol in an antimicrobial composition according to the present invention.

Alternatively, since the method is directed towards surface disinfection, the disinfection time may also be determined by test methods involving a surface. Therefore, the invention preferably relates to a method according to the present invention, wherein the surface disinfection time T2 of said method is less than 60 seconds, preferably less than 15 seconds, wherein T2 is defined as the time starting from the moment of applying the composition to the surface to be disinfected after which the number of microbes per unit area is reduced by a factor of 10000 (i.e. a 4 log reduction), wherein the initial number of microbes preferably exceeds 10^3 , more preferably 10^5 , and even more preferably 10^7 microbes per square centimeter. Such tests may for instance be performed as described in WO 2010/046238, or as described in European Standards EN 13697:2001 and EN 1500:1997.

Use according to the invention

The invention preferably provides for non-therapeutic benefits. Thus, for instance, the invention relates to use of an antimicrobial composition according to the present invention for faster reduction in viable microbial count.

Thus, according to the third aspect of the invention, there is provided use of a composition according to the invention for improved hygiene. Such use relates for example to use of an antimicrobial composition comprising the one or more cyclic monoterpene alcohols, thymol and a carrier, for reduction in viable microbial count, preferably fast reduction of viable microbial count. Thus, such use preferably is use in a method for disinfection. Fast reduction in viable microbial count therefore preferably relates to use for disinfection whereby the disinfection time is less than 300 seconds, preferably less than 120 seconds, more preferably less than 60 seconds, and even more preferably less than 15 seconds. Here, the disinfection is preferably defined similar to the disinfection times T and T2 as described above.

Thus, there is provided use of a composition according to the invention for improved hygiene of surfaces of the human body. Such surfaces include e.g. skin, hands and the oral cavity. According to a preferred aspect, the invention relates to use of a composition according to the invention for improved hand hygiene. According to another preferred aspect, the invention relates to use of a composition according to the invention for improved oral hygiene.

The microbicide compositions of the present invention can be used to inhibit the growth of microorganisms by introducing a microbicidally effective amount of the compositions onto, into or at a locus subject to attack. For instance, in the field of institutional and industrial applications, suitable loci include, for example: industrial

5 process water including electrocoat deposition systems, cooling towers and air washers; gas scrubbers; wastewater treatment; ornamental fountains; reverse osmosis filtration; ultrafiltration; ballast water; evaporative condensers and heat exchangers; pulp and paper processing fluids and additives; mineral slurries; starch; plastics; emulsions; dispersions; paints; latices; coatings, such as varnishes; construction

10 products, such as mastics, caulks, and sealants; construction adhesives, such as ceramic adhesives, carpet backing adhesives, and laminating adhesives; industrial or consumer adhesives; photographic chemicals; printing fluids; household and institutional products used in restaurants, healthcare facilities, schools, food processing facilities and farms including, cleaners, sanitizers and disinfectants, wipes, soaps,

15 detergents, floor polishes and laundry rinse water; cosmetics; toiletries; shampoos; metalworking fluids; conveyor lubricants; hydraulic fluids; leather and leather processing products; textiles; textile and textile processing products; wood and wood processing products, such as plywood, chipboard, wallboard, flakeboard, laminated beams, oriented strandboard, hardboard, and particleboard; oil and gas processing

20 fluids such as injection fluids, fracture fluids, drilling muds and produced water; fuel transportation and storage systems; agriculture adjuvant preservation; surfactant preservation; medical devices; diagnostic reagent preservation; food preservation, such as plastic or paper food wrap; food, beverage, and industrial process pasteurizers; toilet bowls; recreational water; pools; and spas.

25 Preferably, the microbicidal compositions of the present invention are used to inhibit the growth of microorganisms at a locus selected from one or more of mineral slurries, pulp and paper processing fluids and additives, starch, emulsions, dispersions, paints, latices, coatings, construction adhesives (such as ceramic adhesives), carpet backing adhesives, photographic chemicals, printing fluids, household and institutional

30 products such as cleaners, sanitizers, disinfectants, wipes, cosmetics, toiletries, shampoos, soaps, detergents, floor polishes, laundry rinse water, metal working fluids, textile products, wood and wood products, agriculture adjuvant preservation, surfactant preservation, diagnostic reagent preservation, food preservation, food, beverage, and industrial process pasteurizers and oil and gas processing fluids.

Fields of use

The composition according to the invention can in view of the above be applied for disinfection, reduction in viable microbial count or improved hygiene, especially at a surface. In preferred embodiments, the composition is particularly suited for
5 application to the skin. For example, a surface like the hands, face, body, or the oral cavity can suitably be contacted with the composition of the invention. In other preferred embodiments, the surface is any hard surface. Typically, such hard surfaces are surfaces that commonly require cleaning and often also require sanitisation or disinfection. Such surfaces can be found in many household or industrial
10 environments, and can include for example kitchen and bathroom surfaces, table tops, floors, walls, windows, utensils, cutlery, and crockery. Such surfaces can be made from many different materials, for instance plastics, wood, metal, ceramics, glass, concrete, marble, and painted surfaces. In other preferred embodiments, the compositions can be used for such disinfection, reduction in viable microbial count or
15 improved hygiene at loci other than the surfaces as described hereinbefore.

In preferred embodiments, the invention relates to compositions according to the invention for use as or incorporation in home care products and personal care products. More preferably, this embodiment of the invention relates to a composition according to the invention which is a home care product or a personal care product.

20 A "home care product" is a product for the treatment, cleaning, caring or conditioning of the home or any of its contents. The foregoing includes, but is not limited to, compositions, products, or combinations thereof relating to or having use or application in the treatment, cleaning, cleansing, caring or conditioning of surfaces, furniture and atmosphere of the home and household contents, such as clothes, fabrics
25 and/or cloth fibres and the manufacture of all of the foregoing products. A "personal care product" is a product for the treatment, cleaning, caring or conditioning of the person. The foregoing includes, but is not limited to, chemicals, compositions, products, or combinations thereof relating to or having use or application in the treatment, cleaning, cleansing or conditioning of the person (including in particular the
30 skin, hair and oral cavity), and the manufacture of all the foregoing. Home care products and personal care products are for example products marketed under mass market brands, non-limiting examples being soap bars, deodorants, shampoos, and home surface sanitisers/disinfectants.

Another preferred embodiment of the invention relates to compositions
35 according to the invention for use as or incorporation in industrial and/or institutional

products. More preferably, this embodiment of the invention relates to a composition according to the invention which is an industrial and/or an institutional product.

Industrial and institutional products are for example products being marketed under professional brands, non-limiting examples being for industrial, institutional, janitorial,

5 and medical cleaning, cleaning-in-place, food services, veterinary, and agricultural products. Industrial and/or institutional products also include products for cleaning of the person (such as hand sanitisers) for medical offices, hospitals and/or other institutions.

In another preferred embodiment, the invention also relates to a method or use
10 according to the invention involving home care products or personal care products. For example, the method according to the invention – which comprises application of a composition according to the invention in step a – can be a method wherein that composition is a composition for use as or incorporation in home care products and personal care products as described hereinabove. Similarly, in another preferred
15 embodiment, the invention also relates to a method or use according to the invention involving industrial and/or institutional products. For example, the method according to the invention – which comprises application of a composition according to the invention in step a – can be a method wherein that composition is a composition for use as or incorporation in industrial and/or institutional products as described hereinabove.

20 Products and/or methods for use in the home care or personal care field are generally distinct from products and/or methods for use in the industrial and/or institutional field. Thus, for example, a product that is marketed as a home or personal care product will generally not be marketed as a product for industrial and/or institutional use and vice versa. Therefore, certain embodiments of the present
25 invention, when carried forth into practice, will relate to the one field, but not the other.

EXAMPLES

The invention is illustrated by the following non-limiting examples.

30 **Example 1: Assessment of antimicrobial efficacy**

Materials

Cyclic monoterpene alcohols were purchased as fine chemicals from suppliers such as Sigma Aldrich, Alfa Aesar and TCI Fine Chemicals. Thus, 4-isopropyl-1-methylcyclohex-3-enol was obtained as a mixture of isomers with CAS no. [586-82-3].

35 Thymol was purchased as 99.5% pure fine chemical grade (ex Sigma Aldrich).

A terpineol mixture comprising ca 88 wt-% of (S)-alpha-terpineol, and 12 wt-% of gamma-terpineol was purchased from Sigma Aldrich and is referred to below as alpha-terpineol or terpineol unless specified otherwise.

5 General method for assessment of antimicrobial synergy

The efficacies of antimicrobial agents can be usefully compared by determining the Minimum Biocidal Concentration (MBC). The MBC is defined as the lowest absolute concentration of the particular active that provides complete kill (zero bacterial growth).

- 10 The differing behaviours of inhibitory antimicrobials in isolation and mixtures have been widely explored using the concept of the Fractional Concentration and Fractional Inhibitory Concentration (FIC). See for instance JRW Lambert and R Lambert, *J. Appl. Microbiol* **95**, 734 (2003); T. Jadavji, CG Prober and R Cheung, *Antimicrobial Agents and Chemotherapy* **26**, 91 (1984), and WO 2004/006876. These parameters can be
15 defined as follows:

$$\text{FC (component a)} = \frac{\text{Concentration of component a tested in the mixture}}{\text{MIC (component a tested as a single active)}}$$

$$\text{FIC (component a)} = \frac{\text{MIC (component a tested in the mixture)}}{\text{MIC (component a tested as a single active)}}$$

By analogy the Fractional Biocidal Concentration (FBC) is given by:

20

$$\text{FBC (component a)} = \frac{\text{MBC (component a tested in the mixture)}}{\text{MBC (component a tested as a single active)}}$$

- The interactions between antimicrobials can be additive, synergistic or possibly antagonistic depending on whether the efficacy of the combination is equivalent to, greater than or less than that obtained for the same total concentration of the individual
25 components when tested alone.

These relationships can be expressed mathematically by summing the fractional MBC values for all the components present in the mixture to give the "fractional biocidal index":

$$\Sigma\text{FBC} = \text{FBC}_{(\text{component 1})} + \text{FBC}_{(\text{component 2})} + \text{FBC}_{(\text{component 3})} + \dots \text{ etc}$$

such that

$\Sigma\text{FBC} \geq 1$ corresponds to additive or antagonistic bactericidal activity

5 $\Sigma\text{FBC} < 1$ corresponds to synergistic bactericidal activity

Experimental method

Antimicrobial efficacy is tested against a representative pathogenic bacterial organism, Gram negative *Escherichia coli*. Concentrations of actives are expressed in terms of
10 the percentage weight/volume (%w/v) throughout Example 1.

Bacterial stock

An overnight culture of *Escherichia coli* (10536 strain) was prepared in 50 ml total volume of TSB broth, grown for ca. 18 hrs at 37°C and shaken at 150 rpm. 1 ml of this
15 overnight *E. coli* culture was transferred to 50 ml of fresh TSB broth and incubated at 37°C at 150 rpm for ca. 4 hours. This culture was separated into equal volumes and centrifuged at 4000 rpm for 15 minutes, washed with sterile saline (0.85% NaCl), centrifuged once more and re-suspended in saline to give a final concentration of 0.8 OD₆₂₀ equivalent to about 10⁸ cells per millilitre for this particular organism. Here,
20 OD₆₂₀ indicates the absorbance of a sample in a cuvette of 1.0 cm path length at a wavelength of 620 nm. This bacterial stock was used for assaying against antimicrobial actives (in triplicate).

Protocol

25 The following assay describes the testing of 8 materials using 6 dilutions across half of a 96-well micro titre plate (MTP). Using this approach it is possible to assay 16 actives (without replicates) with one full dilution plate, replicating this set up in two halves of the plate columns, 1-6 and 7-12.

30 1M solutions of the test actives were prepared in dimethylsulphoxide (DMSO). Stock solutions of the actives at 1.11 times the desired final concentration were prepared by diluting the DMSO solutions in water, so that for example a 0.89% w/v solution was prepared for a desired "in test" concentration of 0.8% w/v in order to allow for the further dilution of the active when the bacterial suspension is added (dilution from 270µl
35 to 300 µl), as described below.

Aliquots (270 µl) of the materials at 1.11 times the final concentration were dispensed into the wells of the MTP along one column (A1-H1). This MTP was labelled as the "Screening plate".

5

In another MTP, labelled as the "Dilution plate", 270µl of D/E neutralising solution from DIFCO Composition was added to column 1. The composition of the neutralising solution was as follows: pancreatic digest of casein, 5.0g/L; Yeast Extract, 2.5 g/L; Dextrose, 10 g/L, sodium thioglycollate, 1.0 g/L, sodium thiosulphate, 6.0 g/L; sodium
10 bisulphite, 2.5 g/L; Polysorbate 80, 5.0 g/L; lecithin 7.0 g/L; bromocresol purple, 0.02 g/L with a pH in the range 7.6±0.2.

270µl of tryptone diluent solution was added to all the remaining wells of the Dilution MTP (columns 2-6).

15

Bacterial stock (30 µl) was then added to the prepared 270 µl of the solution of actives in the Screening Plate and mixed, using a multichannel pipette with 8 tips to aspirate and dispense the same volume of bacterial stock in parallel to 8 wells in rows A-H. After a contact time of 15 seconds, the mixtures were quenched by transferring 30µl
20 volumes of the mixtures into the 270µl D/E neutralising solution in the prepared dilution plate, using aspiration to mix. After exactly 5 minutes in the D/E neutralising solution, 30µl volumes were transferred from column 1 to column 2 of the Dilution MTP and mixed, before transferring further 30µl volumes from column 2 into column 3. This process was repeated serially diluting the bacteria across the plate to column 6.

25

30µl volumes from each well in the Dilution MTP were transferred onto pre-labelled segment of Tryptone Soya Agar (TSA) plates starting from the lowest bacterial concentration (highest dilution, column 6) to the highest bacterial concentration (column 1). The TSA plates were allowed to stand for ca. 2 hours so that the 30µl
30 inocula spots could dry and the plates were then inverted and incubated overnight at 37°C before enumerating the bacterial colonies at the labelled dilutions to determine the effects of the actives on bacterial growth.

Calculation of results

Mean bacterial survival numbers N_{MBS} (expressed in Log CFU/ml) are obtained by first determining the segment of the TSA plate where the number of bacterial colonies is countable. From the colony number in this segment, N_{MBS} is calculated by the formula:

$$N_{\text{MBS}} = \log\{ N_{\text{col}} \cdot 10^{\text{DF}} \cdot 100 / 3 \}$$

Here, N_{col} is the colony count, and DF is the dilution factor taken from the MTP-well corresponding to the TSA plate segment (i.e. DF may range from 1 for the quench, to 6 for the highest dilution). The factor 100/3 is a conversion factor from the volume of the inocula spot to one millilitre.

Every assay test was performed in triplicate. The reported mean bacterial survival results are the average of such a triplet, the error is the corresponding standard deviation.

Thus, a value of N_{MBS} of about 7 corresponds to a count of about 3 colonies from the fifth dilution well, i.e. with DF = 5. Such a count of about 7 is generally observed when bacteria are exposed to non-biocidal materials. In case no surviving colonies are observed in any segment of the TSA plate, this is interpreted as complete kill and a value of $N_{\text{MBS}}=0$ is reported.

Validation

All test results were validated by running every test assay in parallel with four control experiments on the same MTP. All control experiments are executed exactly according to the above protocol, but with the following active ingredients:

- A 0.025 %(w/v) thymol
- B 0.15 %(w/v) alpha-terpineol
- C 0.025 %(w/v) thymol + 0.15 %(w/v) alpha-terpineol
- D no active component

The control experiments A, B and D validate a test assay by not showing bacterial kill, whereas control experiment C, comprising a synergistic combination of thymol and alpha-terpineol according to WO 2010/046238 A1 validates a test assay by showing complete bacterial kill.

A reference experiment according to the above protocol, but without active component, showed that DMSO does not affect bacterial growth at the concentrations present in the test solutions in this protocol (<5 % (w/v)), as can be seen in Table 2.

Table 2

DMSO in water (% w/v)	Mean bacterial survival [log CFU/ml]	Standard deviation
4.5	8.2	0.1
3.6	8.4	0.2
2.7	8.2	0.1
1.8	8.5	0.2
0.9	8.6	0.1
0.0	8.5	0.1

Results

The above method was applied to assess the antibacterial efficacy of the a cyclic
5 monoterpene alcohol according to the invention. Table 3 shows the antibacterial
activities of a selected cyclic monoterpene alcohol, and selected comparative
compounds both alone and in conjunction with thymol.

Table 3

Example	Thymol concentration C_{thymol}(% w/v)	Concentration of terpeneol / alcohol / comparative compound C_{alc}(% w/v)	N_{MBS}^a	S.D.^b
1:1*	0.075	0	0	0
1:2*	0.05	0	0	0
1:3*	0.025	0	> 7	0.1
1:4*	0	0.5% alpha-terpineol	0	0
1:5*	0	0.4% alpha-terpineol	0	0
1:6*	0	0.3% alpha-terpineol	7	0.2
1:7*	0	0.15% alpha-terpineol	7	0.2
1:8*	0	0.3% 4-isopropyl-1-methyl- cyclohex-3-enol	0	0
1:9*	0	0.15% 4-isopropyl-1-methyl- cyclohex-3-enol	7.3	0.2
1:10	0.025% thymol	0.112% 4-isopropyl-1-methyl- cyclohex-3-enol	0	0
1:11*	0.025% thymol	0.15% (+/-)-limonene	7.4	0.2

1:12*	0.025% thymol	0.15% (+)-p-menth-1-ene	7.1	0.3
1:13*	0.075% thymol	0.15% (+/-)-menthol	7.6	0.1
* Examples (1:1) to (1:9) and (1:11) to (1:13) are comparative examples ^a N _{MBS} in [log CFU/ml] ^b S.D. = standard deviation				

Comparative examples

Determination of the parameter Σ FBC, which is used as measure of the synergistic antimicrobial action of compositions according to the present invention, requires

5 determination of the Minimum Biocidal Concentrations (MBCs) of the relevant actives first. As described above, the MBC for an active can be defined as the lowest concentration of the active that provides zero bacterial survival in the particular medium. Data for examples (1:1) to (1:3) demonstrate that the MBC value for thymol is 0.05% (w/v). For alpha-terpineol, compositions (1:4) to (1:7), show that the

10 MBC is 0.4% w/v. The same analysis has been carried out for 4-isopropyl-1-methyl-cyclohex-3-enol and is summarised in Table 4 below. These MBC values constitute the upper boundaries to the respective MBCs in the particular medium used in these examples.

15 **Table 4:** Minimum biocidal concentrations of antimicrobial components

Component	MBC (% w/v)
thymol	0.05
alpha-terpineol	0.4
4-isopropyl-1-methyl-cyclohex-3-enol	0.3

It is clear from the data in Table 4 that 4-isopropyl-1-methyl-cyclohex-3-enol is antimicrobially more efficacious than terpineol and can be used at lower concentrations.

20

Synergistic interactions

The tested combinations of the selected cyclic monoterpene alcohol with thymol provide complete bacterial kill in example (1:10). Using the MBC values listed in Table 4 above, the fractional MBC values for the components present in these mixtures and

25 the experimental Σ FBC of the compositions can be calculated in order to discriminate

between combinations providing evidence of synergistic effects, as opposed to additive biocidal effects. The results from this analysis are given in Table 5.

Table 5: Extent of synergistic interactions between binary compound mixtures for
5 compositions providing complete bacterial kill

cyclic monoterpene alcohol				thymol		Σ FBC	Evidence of Synergy ^c
compound	Ex.	MBC %(w/v)	FBC ^a	MBC %(w/v)	FBC ^b		
4-isopropyl-1-methyl- cyclohex-3-enol	1:10	0.3	0.37	0.05	0.5	0.87	Yes
^a FBC of cyclic monoterpene alcohol: C_{alc}/MBC_{alc} ^b FBC of thymol: $C_{thymol} / MBC_{thymol}$ ^c Criterion for synergy: (Σ FBC <1)							

For Example (1:10), the Σ FBC value is below 1, thus providing evidence for synergistic interactions, according to the set criteria. Therefore, these examples show how the
10 antimicrobial efficacy of thymol and a cyclic monoterpene alcohol according to invention is enhanced when they are applied together. Such synergies allow for reductions in the concentrations of the antimicrobials required to achieve complete kill.

Comparative examples

15 The comparative examples (1:11) to (1:13) show that compositions comprising thymol and several compounds that are not compounds according to the present invention at concentrations comparable to those in the examples according to the invention do not give rise to fast antimicrobial action.

20 **Example 2: Automated assessment of efficacy in surfactant base**

Sample preparation

In these examples, the efficacy of combinations of thymol and 4-isopropyl-1-methyl-
cyclohex-3-enol were tested in a surfactant cleansing formulation comprising 2.85%
sodium cocoyl glycinate and 1.85% sodium lauroamphoacetate. This corresponds to a
25 50% in use dilution with water of a typical neat formulation containing 5.7% cocoyl
glycinate and 3.7% % sodium lauroamphoacetate during hand washing. Solutions
were prepared such that the concentrations of the surfactant components and test

actives were 1.1 times the final desired concentration in order to allow for dilution with the bacterial inoculum in the test. The solutions were manually adjusted to pH 10.0 by dropwise addition of sodium hydroxide solution, as measured with a pH meter at ambient temperature. Solutions of the thymol and/or 4-isopropyl-1-methyl-cyclohex-3-enol were prepared at a maximum of 24 hours before testing. The same thymol and 4-isopropyl-1-methyl-cyclohex-3-enol were tested as in Example 1.

Test methodology

The efficacy of the combinations of the present invention was determined against the same bacterium as in Example 1, *Escherichia coli* (*E. coli* – ATCC #10536), at a concentration of approximately 1×10^8 bacteria per mL.

Tests were conducted using standard microtiter plate assays using an automated liquid handling system. 270µl of the surfactant test solution was pipetted into each well of the microtitre plate (Nunc F Gamma Irradiated 96F untreated microtitre plates of clear polystyrene) and 30µl of the bacterial suspension was then added. After exactly 15 seconds of bacterial exposure, a 30µl volume of bacterial cells was withdrawn and transferred to 270µl of D/E quench solution. After 5 minutes in the D/E quench, the optical density (OD) was measured for each plate in turn at two specific wavelengths (450nm and 590nm). These provide a dual check of antimicrobial activity, as the OD₄₅₀ reading is specific for the yellow colour of D/E quench when bacterial growth is observed, whereas OD₅₉₀ is specific for the initial purple colour of the D/E quench which is retained if no bacterial growth is observed. After the time zero OD measurements, plates were then incubated at 37°C overnight (16 hours) before repeating the OD measurements. Delta OD values were calculated by subtracting the OD values at 16 hours from the initial value at time zero from those at time = 16 hours. Bacterial growth is observed as an increase in OD₄₅₀ and a decrease in OD₅₉₀. To identify antibacterially efficacious systems (those preventing appreciable bacterial growth after incubation), the following threshold changes in OD readings have been adopted: if (1). OD₄₅₀ increases by less than 0.2 absorbance unit (AU) on incubation and (2). OD₅₉₀ decreases by less than 0.35 AU on incubation. Conversely, where OD₄₅₀ increases by more than 0.1 AU and OD₅₉₀ decreases by more than 0.1 AU after incubation, corresponding to a colour shift from purple to yellow, the test system allows bacterial growth and is not deemed efficacious. Four replicate measurements in the same plate have been made for each test system. The number of replicate wells

showing either bacterial growth or no growth is also readily assessed by eye by following the colour change. Thymol and terpineol were tested both alone and in combination for comparison purposes.

- 5 Dose responses for individual components and binary mixtures of actives at a fixed concentration ratio were generated by sequential dilution of liquors with further surfactant solution to obtain a series of endpoints ranging from 0.2 to 0.1% of the thymol and 0.5% to 0.25% of the 4-isopropyl-1-methyl-cyclohex-3-enol. In each case, binary mixtures were assessed in the weight to weight ratio of thymol to 4-isopropyl-1-
- 10 methyl-cyclohex-3-enol of 1:2.5.

Table 6: Antibacterial activities of 4-isopropyl-1-methyl-cyclohex-3-enol alone, and in combination with thymol in model surfactant solution.

Ex.	C _{thymol} ^(a) (% w/v)	C _{alc} ^(b) (% w/v)	DeltaOD (450 nm) ^(c)		DeltaOD (590 nm) ^(d)		N _{rep} ^(e) out of 4
			Mean	S.D. ^(f)	Mean	S.D. ^(f)	
2:1*	0	0	-0.60	0.02	0.64	0.02	4
2:2*	0.2%	0	-0.54	0.02	0.64	0.02	4
2:3*	0.175%	0	-0.54	0.02	0.56	0.02	4
2:4*	0.15%	0	-0.57	0.01	0.55	0.01	4
2:5*	0.125%	0	-0.58	0.01	0.55	0.01	4
2:6*	0.1%	0	-0.58	0.00	0.54	0.02	4
2:7*	0.075%	0	-0.59	0.01	0.54	0.01	4
2:8*	0.05%	0	-0.58	0.03	0.55	0.01	4
2:9*	0.025%	0	-0.55	0.01	0.65	0.02	4
2:10*	0	0.5%	-0.41	0.01	0.66	0.02	4
2:11*	0	0.4%	-0.44	0.02	0.59	0.02	4
2:12*	0	0.35%	-0.48	0.01	0.56	0.01	4
2:13*	0	0.3%	-0.51	0.02	0.56	0.01	4
2:14*	0	0.25%	-0.52	0.02	0.56	0.01	4
2:15	0.2%	0.5%	0.19	0.03	0.20	0.03	0
2:16	0.175%	0.4375%	0.15	0.02	0.16	0.01	0
2:17	0.15%	0.375%	0.12	0.02	0.14	0.01	0
	0.125%	0.3125%	-0.22	0.33	0.31	0.25	2

Ex.	C _{thymol} ^(a) (% w/v)	C _{alc} ^(b) (% w/v)	DeltaOD (450 nm) ^(c)		DeltaOD (590 nm) ^(d)		N _{rep} ^(e) out of 4
			Mean	S.D. ^(f)	Mean	S.D. ^(f)	
	0.1%	0.25%	-0.51	0.01	0.54	0.02	4
* Examples marked with an asterisk (*) are comparative examples (a) Concentration of thymol (b) Concentration of 4-isopropyl-1-methyl-cyclohex-3-enol (c) DeltaOD (450 nm) = OD₄₅₀ (time = 16 hours) - OD₄₅₀ (time zero) (d) DeltaOD (590 nm) = OD₅₉₀ (time = 16 hours) - OD₅₉₀ (time zero) (e) N_{rep} = No. of replicates showing growth (out of 4) (f) S.D. = standard deviation							

Table 7: Minimum biocidal concentrations of antimicrobial components in 2.85% sodium cocoyl glycinate + 1.85 % sodium lauroamphoacetate solution at pH 10

Component	MBC (% w/v)
Thymol	> 0.2
4-isopropyl-1-methyl-cyclohex-3-enol	> 0.5

5 Results

The surfactants used are not themselves antimicrobially active against *E. coli* at the concentrations employed as shown by the results of Ex. (2:1) in Table 6. Thus, any antimicrobial efficacy can be ascribed to the 4-isopropyl-1-methyl-cyclohex-3-enol and/or thymol. Table 7 presents MBC-values determined similarly as described for

10 Example 1. Thymol and 4-isopropyl-1-methyl-cyclohex-3-enol have an MBC higher than the highest tested concentrations, in the presence of the specified surfactants.

The results of Tables 6 and 7 demonstrate that 4-isopropyl-1-methyl-cyclohex-3-enol show 15 second bactericidal efficacy (complete kill in all 4 or 3 replicates, respectively) against *E. coli* when tested in combination with thymol at concentrations

15 lower than their MBC in the same surfactant formulation (comprising cocoyl glycinate and lauroamphoacetate).

Thus, it was found that cyclic monoterpene alcohols according to the invention and in particular 4-isopropyl-1-methyl-cyclohex-3-enol show enhanced antimicrobial action in combination with thymol in the presence of surfactant, in particular cocoyl

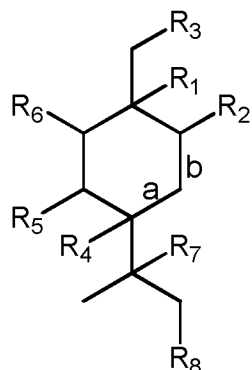
20 glycinate and lauroamphoacetate.

CLAIMS

1. An antimicrobial composition comprising:

- i. 0.001 to 5% by weight of thymol,
- ii. 0.01 to 5% by weight of one or more cyclic monoterpene alcohols, and
- iii. a carrier;

wherein the one or more cyclic monoterpene alcohols have the following structure



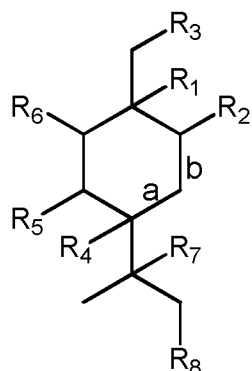
wherein

- one of the bonds a and b is a double bond, the other being a single bond;
 - one of the substituents R_1 to R_8 is a hydroxy, the others being hydrogen;
- subject to the proviso that
- the double bond and the hydroxy are not attached to the same carbon atom;
 - and
 - R_4 is absent when bond a is the double bond;

and wherein the carrier comprises water, ethanol, propanol, isopropanol, ethylene glycol, propylene glycol, diethylene glycol, or mixtures thereof.

2. An antimicrobial composition according to claim 1, wherein the one or more cyclic monoterpene alcohols are selected from the group consisting of
 - 4-isopropyl-1-methylcyclohex-3-enols,
 - 3-isopropyl-6-methylcyclohex-2-enols, and
 - 4-isopropyl-1-methylcyclohex-2-enols.
3. An antimicrobial composition according to claim 2, wherein the one or more cyclic monoterpene alcohols are selected from the group consisting of
 - 4-isopropyl-1-methylcyclohex-3-enols and
 - 4-isopropyl-1-methylcyclohex-2-enols.

4. An antimicrobial composition according to claim 3, wherein the cyclic monoterpene alcohol is 4-isopropyl-1-methylcyclohex-3-enol.
5. An antimicrobial composition according to any one of claims 1 to 4 wherein the thymol is present at a concentration of 0.01 to 0.4% by weight, and the one or more cyclic monoterpene alcohols are present at a concentration of 0.05 to 1% by weight.
6. An antimicrobial composition according to any one of claims 1 to 5 comprising from 1 to 80% by weight of one or more surfactants.
7. An antimicrobial composition according to claim 6 wherein the one or more surfactants are selected from anionic, non-ionic, and a combination of anionic and non-ionic surfactants.
8. A solid antimicrobial composition according to any one of claims 6 to 8 comprising:
 - a. 0.05 to 5% by weight of thymol,
 - b. 0.05 to 5% by weight of the one or more cyclic monoterpene alcohols,
 - c. 5 to 30% by weight of water, and;
 - d. 30 to 80% by weight of the one or more surfactants.
9. A method of disinfecting a surface comprising the steps of
 - c. applying on to the surface a composition comprising
 - i. 0.001 to 5% by weight of thymol,
 - ii. 0.001 to 5% by weight of one or more cyclic monoterpene alcohols, and
 - iii. a carrier;wherein the one or more cyclic monoterpene alcohols have the following structure



wherein

- one of the bonds a and b is a double bond, the other being a single bond;
 - one of the substituents R₁ to R₈ is a hydroxy, the others being hydrogen;
- subject to the proviso that
- the double bond and the hydroxy are not attached to the same carbon atom;
- and

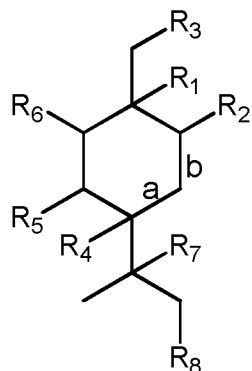
- R₄ is absent when bond a is the double bond; and
- d. removing the composition from the surface.

10. A method according to claim 9 wherein the composition is a composition according to any one of claims 1 to 8.
11. A method according to claim 9 or 10, wherein the surface is a skin surface.
12. A method according to any one of claims 9 to 11, wherein step (b) is started between 2 minutes and 5 seconds, more preferably between 1 minute and 10 seconds, even more preferably between 30 and 10 seconds and still more preferably between 20 and 15 seconds after commencement of step (a).
13. A method according to any one of claims 9 to 12, wherein the disinfection time T of said method is less than 300 seconds, preferably less than 60 seconds, and more preferably less than 15 seconds; wherein T is defined as the time that elapses from the moment of adding the composition to a microbial culture until the number of microbes per unit volume of the culture is reduced by a factor of 100 000; and wherein the initial number of microbes preferably exceeds about 100 000 000 microbes per millilitre and wherein the composition is preferably a liquid composition.

14. Use of a composition for improved hand hygiene, whereby the composition comprises

- i. 0.001 to 5% by weight of thymol,
- ii. 0.001 to 5% by weight of one or more cyclic monoterpene alcohols, and
- iii. a carrier;

wherein the one or more cyclic monoterpene alcohols have the following structure



wherein

- one of the bonds a and b is a double bond, the other being a single bond;
 - one of the substituents R₁ to R₈ is a hydroxy, the others being hydrogen;
- subject to the proviso that
- the double bond and the hydroxy are not attached to the same carbon atom;
- and
- R₄ is absent when bond a is the double bond.

15. Use of a as specified in claim 14 for improved oral hygiene.

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2013/061717

A. CLASSIFICATION OF SUBJECT MATTER

INV. A01P1/00 A01N31/08 A01N31/06 A01N31/04 A61K8/34
A61K31/05

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)

A01N A61K

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, CHEM ABS Data, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	ARACELI PÉREZ-VÁSQUEZ ET AL: "Antimicrobial activity and chemical composition of the essential oil of Hofmeisteria schaffneri", JOURNAL OF PHARMACY AND PHARMACOLOGY, vol. 63, no. 4, 1 March 2011 (2011-03-01), pages 579-586, XP055037535, ISSN: 0022-3573, DOI: 10.1111/j.2042-7158.2010.01243.x cited in the application	1-5
Y	table 2 ----- -/--	6-15

☒ Further documents are listed in the continuation of Box C.

☒ 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

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"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

11 October 2013

Date of mailing of the international search report

18/10/2013

Name and mailing address of the ISA/

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INTERNATIONAL SEARCH REPORT

International application No

PCT/EP2013/061717

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	BYUN Y H ET AL: "Analysis of composition and activity of essential oil from Chrysanthemum zawadskii var. latilobum and C. indicum against antibiotic-resistant pathogenic bacteria", NATURAL PRODUCT SCIENCES, KOREAN SOCIETY OF PHARMACOGNOSY, SEOUL, KR, vol. 14, no. 2, 1 January 2008 (2008-01-01), pages 138-142, XP008102334, ISSN: 1226-3907 cited in the application	1-5
Y	table 1	6-15
X	Abdeslam Jaafari ET AL: "Chemical composition and antitumor activity of different wild varieties of Moroccan thyme", Revista Brasileira de Farmacognosia, 2007, 17(4),, 1 December 2007 (2007-12-01), pages 477-491, XP055027766, DOI: 10.1590/S0102-695X2007000400002 Retrieved from the Internet: URL: http://www.scielo.br/pdf/rbfar/v17n4/a02v17n4.pdf [retrieved on 2012-05-22] cited in the application	1-5
Y	tables 1-4	6-15
A	FR 2 697 133 A1 (TRANSBIOTECH [FR]; TERROM GERARD) 29 April 1994 (1994-04-29) cited in the application pages 5-6,8; claims 1,2,6,13	1-15
Y	WO 2010/046238 A1 (UNILEVER NV [NL]; UNILEVER PLC [GB]; UNILEVER HINDUSTAN [IN]; CHAKRABO) 29 April 2010 (2010-04-29) cited in the application claims 1-14; table 1	6-15

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/EP2013/061717

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