

(19) **DANMARK**



Patent- og
Varemærkestyrelsen

(12)

Oversættelse af europæisk patentskrift

(10) **DK/EP 3427718 T3**

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- (51) Int.Cl.: **A 61 K 8/06 (2006.01)** **A 61 K 8/02 (2006.01)** **A 61 K 8/34 (2006.01)**
A 61 K 8/362 (2006.01) **A 61 K 8/368 (2006.01)** **A 61 K 8/46 (2006.01)**
A 61 K 8/92 (2006.01) **A 61 Q 11/00 (2006.01)** **A 61 Q 17/00 (2006.01)**
- (45) Oversættelsen bekendtgjort den: **2022-08-15**
- (80) Dato for Den Europæiske Patentmyndigheds bekendtgørelse om meddelelse af patentet: **2022-07-06**
- (86) Europæisk ansøgning nr.: **18184912.6**
- (86) Europæisk indleveringsdag: **2011-06-30**
- (87) Den europæiske ansøgnings publiceringsdag: **2019-01-16**
- (30) Prioritet: **2010-06-30 US 82792710**
- (62) Stamansøgningsnr: **11172281.5**
- (84) Designerede stater: **AL AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HR HU IE IS IT LI LT LU LV MC MK MT NL NO PL PT RO RS SE SI SK SM TR**
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- (54) Benævnelse: **IKKE-ALKOHOLHOLDIG, BIOAKTIV MUNDSKYL MED ÆTERISK OLIE**
- (56) Fremdragne publikationer:
EP-A2- 0 338 978
EP-A2- 2 401 999
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DESCRIPTION

Field of the Invention

[0001] The invention relates generally to mouth rinses for the prevention and elimination of bad breath as well as for the reduction of oral microorganisms responsible for the development of dental plaque and tooth decay. In particular, the present invention relates to a non-alcohol or reduced alcohol mouth rinses effective at preventing the above-mentioned problems.

Background of the Invention

[0002] Mouth rinse or mouthwash compositions have been used by people for many years for the prevention of bad breath and for the elimination of bacteria and other oral microorganisms that are responsible not only for bad breath but also tooth decay, plaque and gum diseases such as gingivitis and periodontitis. To this end, antiseptic mouthwashes in the past have been designed to clean the oral cavity, provide fresh breath and kill these pathogenic microbes.

[0003] Leading antiseptic mouth rinses have traditionally contained alcohol (i.e., ethanol) at fairly high levels, ranging from approximately 20% up to about 30% by volume, based on the total mouthwash volume (hereinafter referred to as "% v/v"). Alcohol is used both as a vehicle and as a solvent in which the active ingredients, and additives such as astringents, fluorides, color additives, flavor oils, and the like, can be dissolved and then dispersed into solution. Alcohol also provides a preservative role for the mouth rinse during storage and use, and enhances the flavor oil organoleptic cues.

[0004] However, the use of high levels of alcohol may sometimes be found unacceptable by some mouthwash users. Senior citizens have also complained about problems related to gargling with such mouth rinses, and chronic exposure has been found to result in a feeling of gum "burn" resulting from the high concentrations of alcohol. It has also been reported that alcoholic mouth rinses can result in an unpleasant "dry mouth" sensation.

[0005] On the other hand, reducing the levels of alcohol in these mouth rinse compositions can have significant disadvantages. It has been found that lower alcohol concentrations result not only in a loss in the solubility of the actives and other ingredients in the composition, but also in a noticeable decrease in the ability of the composition to kill the oral microorganisms responsible for bad breath, plaque and gum disease. This loss in antimicrobial activity is not only due to the reduction of alcohol as a vehicle, but also to the reduced bioavailability of the solubilized actives.

[0006] Thymol, for example, is a well-known antiseptic compound, also known as an essential oil, which is utilized for its antimicrobial activity in a variety of mouthwash or mouth rinse preparations. In particular, thymol can be utilized in oral hygiene compositions such as mouth rinses in sufficient quantities to provide desired beneficial therapeutic effects. Mouthwashes with thymol are well-known, and have been used by millions of people for over one hundred years. They have been proven effective in killing microbes in the oral cavity that are responsible for plaque, gingivitis and bad breath. Thymol, together with other essential oils such as methyl salicylate, menthol and eucalyptol, comprise the active component in some antiseptic mouth rinses. These oils achieve good efficacy although present in small amounts. Without being restricted to any specific theory, it is now believed that the efficacy and taste of antiseptic mouth rinses may be due to the improved dispersion or dissolution of the oils and bioavailability after such dispersion or dissolution of

these four active ingredients.

[0007] The above described dispersion or dissolution is also important from an aesthetic point of view since a clear mouth rinse solutions are certainly more preferred by consumers than cloudy, turbid or otherwise heterogeneous ones. Obviously then, there is a substantial need for the development of a reduced and/or no alcohol mouth rinses which provide improved dispersion or dissolution of the essential oils yet maintain the bioavailability of the essential oils for preventing bad breath, killing oral microbes and reducing or eliminating plaque.

[0008] US 5,817,295 describes a substantially alcohol-free oral mouthwash composition which is said to be effective in eliminating the bacteria and other oral microflora responsible for the production of plaque, periodontitis, gingivitis, gum disease and bad breath. US 5,290,541 describes oral compositions which are said to be effective against plaque and gingivitis and which contain a noncationic water insoluble antibacterial agent. WO 96/16633 describes an antimicrobial mouthwash composition that is said to be useful in the prevention and reduction of bad breath, plaque and related gum diseases.

Summary of the Invention

[0009] The present invention is defined in the claims. The present invention provides an antimicrobial mouth rinse composition, comprising: an oil phase comprising at least one antimicrobial essential oil; a solvent system comprising a polyol solvent and a sugar alcohol solvent; an alkylsulfate surfactant; a buffer system and water, wherein the composition is free of C₂-C₄ monohydric alcohols.

[0010] The reduced alcohol or non-alcohol, antimicrobial mouth rinse composition is a microemulsion comprising micelles having an aggregate size of less than about 200 nm.

[0011] The reduced alcohol or non-alcohol, antimicrobial mouth rinse composition of the present invention preferably exhibits a high level of antimicrobial activity as measured by an M-factor greater than 0.5 (or about 0.5), optionally greater than 1.0 (or about 1.0) optionally, greater than 2.0 (or about 2.0), or optionally greater than 3.0 (or about 3.0) where "M-factor" equals the log RLU value of water used as the negative control minus the log RLU value of the mouth rinse composition being tested. In addition, the oral mouth rinse compositions of this invention are clear (to the unaided human eye) and aesthetically appealing products.

[0012] In further embodiments, the reduced alcohol or non-alcohol, antimicrobial mouth rinse composition of the present invention comprises (a) an antimicrobially effective amount of one or more antimicrobial essential oils; (b) a co-solvent of a glycol and a polyol; (c) an alkylsulfate surfactant; and (d) water.

Detailed Description of the Invention

[0013] The compositions of the present invention can comprise, consist of, or consist essentially of the essential elements and limitations of the invention described herein, as well any of the additional or optional ingredients, components, or limitations described herein. The term "comprising" (and its grammatical variations) as used herein is used in the inclusive sense of "having" or "including" and not in the exclusive sense of "consisting only of."

[0014] The terms "a" and "the" as used herein are understood to encompass the plural as well as the

singular.

[0015] The citation of any document is not to be construed as an admission that it is prior art with response to the present invention.

[0016] The reduced alcohol or non-alcohol mouthwash and mouth rinse compositions described herein provide an antimicrobially effective amount of one or more antimicrobial essential oils towards oral microorganisms responsible for oral malodor and the build-up of plaque and calculus and the resulting tooth and gum diseases that may follow.

[0017] The phrase "antimicrobially effective amount" means the concentration or quantity or level of the compound of the present invention that can attain a particular medical end in having toxic activity for oral microorganisms.

[0018] The phrase "orally acceptable" means that the carrier is suitable for application to the surfaces of the oral cavity or ingestion by a living organism including, but not limited to, mammals and humans without undue toxicity, incompatibility, instability, allergic response, and the like.

[0019] All percentages, parts and ratios are based upon the total weight of the composition of the present invention, unless otherwise specified. All such weights as they pertain to the listed ingredients are based on the level of the particular ingredient described and, therefore, do not include carriers or by-products that may be included in commercially available materials, unless otherwise specified.

[0020] The phrase "reduced alcohol" or "reduced level of alcohol" indicates that the liquid compositions are essentially free of C₂-C₄ monohydric alcohol, which means the liquid compositions or mouth rinses contain an amount of a C₂-C₄ monohydric alcohol up to 10% v/v (or about 10% v/v), optionally, up to 5% v/v (or about 5% v/v), optionally, up to 1.0% v/v (or about 1.0% v/v), optionally up to 0.1% v/v (or about 0.1% v/v) by volume of the total composition. The compositions of the present invention are free of C₂-C₄ monohydric alcohols.

[0021] The term "sterile water", as used herein, means sterile water for irrigation/injection U.S.P. The USP designation means that the sterile water for irrigation/injection is the subject of an official monograph in the current (as of the filing date of this application) US Pharmacopeia.

[0022] Unless otherwise specified, the term "oil(s)" or "oily" component means any hydrophobic, water immiscible component, including but not limited to, essential oils (such as menthol, thymol, eucalyptol and methyl salicylate), flavor oils and hydrophobic compounds such as vitamin E, vitamin E acetate, apigenin and triclosan.

Oil Phase

[0023] The compositions of the present invention comprise an oil phase comprising at least one oil. The oil phase of the present invention comprises at least one antimicrobial essential oil.

Antimicrobial Essential Oils

[0024] The enhanced antimicrobial efficacy of the non-alcohol mouth rinse compositions herein may be

attributed to the presence of minor amounts of one or more antimicrobial or bioactive essential oils (i.e. thymol, eucalyptol, menthol and methyl salicylate).

[0025] Thymol, $[(CH_3)_2CHC_6H_3(CH_3)OH]$, also known as isopropyl-m-cresol], is only slightly soluble in water but is soluble in alcohol, and its presence is one of the reasons alcohol was necessary in the well-established, high alcohol commercial mouth rinses. Methyl salicylate, $[C_6H_4OHCOOCH_3]$, also known as wintergreen oil], additionally provides flavoring to the together with its antimicrobial function. Eucalyptol ($C_{10}H_{18}O$, also known as cineol) is a terpene ether and provides a cooling, spicy taste. Eucalyptol may be used in place of thymol in the same amount if desired. Menthol ($CH_3C_6H_3(C_3H_7)OH$), also known as hexahydrothymol) is also only slightly soluble in alcohol, and is fairly volatile. Menthol, in addition to any antiseptic properties, provides a cooling, tingling sensation.

[0026] The essential oils may be used in amounts effective to provide antimicrobial activity in the oral cavity. In specific embodiments, the total amount of essential oils present in the disclosed compositions can be from 0.001% (or about 0.001%) to 0.35% (or about 0.35%) w/v, or optionally from 0.16% (or about 0.16%) to 0.28% (or about 0.28%) w/v of the composition.

[0027] The compositions of the present invention may contain thymol and additionally eucalyptol, menthol, or methyl salicylate, or mixtures thereof. Optionally, the composition contains all four of these essential oils.

[0028] Thymol may be employed in amounts of from 0.001% (or about 0.001%) to 0.25% (or about 0.25%) w/v, or optionally from 0.04% (or about 0.04%) to 0.07% (or about 0.07%) w/v of the composition. Eucalyptol may be employed in amounts of from 0.001% (or about 0.001%) to 0.11% (or about 0.11%) w/v, or optionally from 0.085% (or about 0.085%) to 0.10% (or about 0.10%) w/v of the composition. Menthol may be employed in amounts of from 0.001% (or about 0.001%) to 0.25% (or about 0.25%) w/v, or optionally from 0.035% (or about 0.035%) to 0.05% (or about 0.05%) w/v of the composition. Methyl salicylate may be employed in amounts of from 0.001% (or about 0.001%) to 0.08% (or about 0.08%) w/v, or optionally from 0.04% (or about 0.04%) to 0.07% (or about 0.07%) w/v of the composition.

[0029] The carrier for the essential oils (the active ingredients) may typically be a water-alcohol mixture, generally water-ethanol. In the past, some antiseptic oral mouth rinse compositions, required ethanol levels of up to about 27% v/v. These high levels were necessary to assist the actives in providing the necessary antimicrobial functionality as well as providing a clear, aesthetically attractive liquid medium. Merely reducing the alcohol levels, without the addition of other formulation components, results in a cloudy, less efficacious product.

[0030] Without being bound to any theory, it is believed that in these high alcohol level oral compositions, the alcohol solubilizes the antimicrobial essential oils and in so doing acts by an active enhancement mechanism. The antimicrobial essential oils are more readily dispersed throughout the solution and remain free or unbound to attack pathogenic microbes throughout the oral cavity. Reducing the alcohol levels was believed to adversely affect this enhancement mechanism. In accordance with the present invention, however, it was surprisingly and unexpectedly found that the level of alcohol can be reduced or eliminated without sacrificing antimicrobial efficacy or clarity if the mouth rinse composition contains a solvent system and alkyl sulfate surfactant as taught herein.

[0031] The total amount of oil phase present in the disclosed compositions of the present invention should preferably not exceed 1.35% w/v (or about 1.35% w/v) of the mouth rinse composition. Optionally, the total oil phase, can be present in an amount of from 0.04% (or about 0.04%) to 1.35% (or about 1.35%) w/v, or optionally from 0.10% (or about 0.10%) to 0.4% (or about 0.4%) w/v of the mouth rinse composition.

Solvent System

[0032] The mouth rinse compositions of the present invention also include a solvent system comprising at least one polyol solvent and at least one sugar alcohol.

Polyol Solvent

[0033] Polyol or polyhydric alcohol solvents used in the solvent system of the present invention are selected from the group consisting of polyhydric alkanes (such as propylene glycol, glycerin, butylene glycol, hexylene glycol, 1,3-propanediol); polyhydric alkane esters (dipropylene glycol, ethoxydiglycol); polyalkene glycols (such as polyethylene glycol, polypropylene glycol) and mixtures thereof. The polyol solvent may be present in an amount of from 1.0% (or about 1.0%) to 30.0% (or about 30.0%) w/v, or optionally from 5.0% (or about 5.0%) to 15.0% (or about 15.0%) w/v of the composition.

Sugar Alcohol Solvent

[0034] The sugar alcohol solvent(s) may be selected from those multi-hydroxy-functional compounds that are conventionally used in oral and ingestible products. The sugar alcohol (s) should preferably be non-metabolized and non-fermentable sugar alcohol (s). The sugar alcohols are selected from the groups consisting of xylitol, sorbitol, mannitol, maltitol, inositol, allitol, altritol, dulcitol, galactitol, glucitol, hexitol, iditol, pentitol, ribitol, erythritol and mixtures thereof. Optionally, the sugar alcohol is selected from the group consisting of sorbitol and xylitol or mixtures thereof. Optionally, the sugar alcohol is sorbitol.

[0035] The total amount of sugar alcohol (s) which are added to effectively aid in the dispersion or dissolution of the active ingredients should not exceed 17% w/v of the composition. Optionally, total amount of sugar alcohol should not exceed 10% w/v (or about 10% w/v) of the composition. The sugar alcohol can be in an amount of from 1.0% (or about 1.0%) to 17.0% (or about 17.0%) w/v, or optionally from 5.0% (or about 5.0%) to 15.0% (or about 15.0%) w/v of the composition.

[0036] The total amount of the solvent system which is added to effectively aid in the dissolution or dispersion of the active ingredients should preferably not exceed 47% w/v (or about 47% w/v) of the composition. Optionally, total amount of solvent system should not exceed 20% w/v (or about 20% w/v) of the composition. The solvent system can be in an amount of from 2% (or about 2%) to 47% (or about 47%) w/v, or optionally from 10% (or about 10%) to 20% (or about 20%) w/v of the composition.

[0037] The ratio of the sugar alcohol to the polyol solvent in the composition may preferably be from 10:1 (or about 10:10) to 1:10 (or about 1:10), optionally from 5:1 (or about 5:1) to 1:5 (or about 1:5), optionally 1:1 (or about 1:1) by weight.

Alkyl Sulfate Surfactant

[0038] The mouth rinse compositions of the present invention also contain at least one alkyl sulfate surfactant. Suitable alkyl sulfate surfactants include, but are not limited to sulfated C₈ to C₁₈, optionally sulfated C₁₀ to C₁₆ even numbered carbon chain length alcohols neutralized with a suitable basic salt such

as sodium carbonate or sodium hydroxide and mixtures thereof such that the alkyl sulfate surfactant has an even numbered C₈ to C₁₈, optionally C₁₀ to C₁₆, chain length. The alkyl sulfate may be selected from the group consisting of sodium lauryl sulfate, hexadecyl sulfate and mixtures thereof. Commercially available mixtures of alkyl sulfates may be used. A typical percentage breakdown of alkyl sulfates by alkyl chain length in commercially available sodium lauryl sulfate (SLS) is as follows:

Alkyl Chain Length	Component Percentage in SLS
C ₁₂	> 60%
C ₁₄	20% - 35%
C ₁₆	< 10%
C ₁₀	< 1%
C ₁₈	< 1%

[0039] Suitable commercially available mixtures include Stepanol WA-100 NF USP, (Stepan, Northfield, IL), Texapon K12 G PH, (Texapon, Cognis, Germany) and mixtures thereof.

[0040] The amount of the alkyl sulfate surfactant added to the composition can be from 0.05% (or about 0.05%) to 2.0% (or about 2.0%) w/v, or optionally from 0.1% (or about 0.1%) to 0.5% (or about 0.5%) w/v of the composition.

[0041] Additional surfactants may be added with the alkylsulfate surfactant to aid in solubilization of the essential oils provided such surfactants do not affect the bioavailability of the essential oils. Suitable examples include additional anionic surfactants, nonionic surfactants, amphoteric surfactants and mixtures thereof. However, the total surfactant concentration (including the alkyl sulfate surfactant alone or in combination with other surfactants) of the mouth rinses of the present invention should preferably be less than or equal to 2% (or about 2%), optionally, the total surfactant concentration should be less than or equal to 1.5% (or about 1.5%), optionally, the total surfactant concentration should be less than or equal to 1.0% (or about 1.0%), optionally, the total surfactant concentration should be less than or equal to 0.5% (or about 0.5%) by weight of the composition.

[0042] Anionic surfactants useful herein include, but are not limited to, sarcosine type surfactants or sarcosinates; taurates such as sodium methyl cocoyl taurate; sodium lauryl sulfoacetate; sodium lauroyl isethionate; sodium laureth carboxylate; sodium dodecyl benzenesulfonate and mixtures thereof. Many suitable anionic surfactants are disclosed in U.S. Pat. No. 3,959, 458, to Agricola, et al., herein incorporated by reference in its entirety.

[0043] Nonionic surfactants which can be used in the compositions of the present invention include, but are not limited to, compounds produced by the condensation of alkylene oxide groups (hydrophilic in nature) with an organic hydrophobic compound which may be aliphatic or alkyl-aromatic in nature. Examples of suitable nonionic surfactants include, but are not limited to, alkyl polyglucosides; ethoxylated hydrogenated castor oils available commercially for example under the trade name CRODURET (Croda Inc., Edison, NJ), and/or; fatty alcohol ethoxylates; polyethylene oxide condensates of alkyl phenols; products derived from the condensation of ethylene oxide with the reaction product of propylene oxide and ethylene diamine; ethylene oxide condensates of aliphatic alcohols; long chain tertiary amine oxides; long chain tertiary phosphine oxides; long chain dialkyl sulfoxides; and mixtures thereof.

[0044] The amphoteric surfactants useful in the present invention include, but are not limited to, derivatives of aliphatic secondary and tertiary amines in which the aliphatic radical can be a straight chain or branched

and wherein one of the aliphatic substituents contains from about 8 to about 18 carbon atoms and one contains an anionic water-solubilizing group, e.g., carboxylate, sulfonate, sulfate, phosphate, or phosphonate. Examples of suitable amphoteric surfactants include, but are not limited alkylimino-dipropionates, alkylamphoglycinates (mono or di), alkylamphopropionates (mono or di), alkylamphoacetates (mono or di), N-alkyl β -aminopropionic acids, alkylpolyamino carboxylates, phosphorylated imidazolines, alkyl betaines, alkylamido betaines, alkylamidopropyl betaines, alkyl sultaines, alkylamido sultaines, and mixtures thereof. The amphoteric surfactant may be selected from the group consisting of alkylamidopropyl betaines, amphoacetates such as sodium lauroamphoacetate and mixtures thereof. Mixtures of any of the above mentioned surfactants can also be employed. A more detailed discussion of anionic, nonionic and amphoteric surfactants can be found in U.S. Pat. Nos. 7,087,650 to Lennon; 7,084,104 to Martin et al.; 5,190,747 to Sekiguchi et al.; and 4,051,234, Gieske, et al.,

[0045] The additional surfactant to be added the mouth rinses of the present invention with the alkyl sulfate surfactant may be selected from the group consisting of taurates. Optionally, the additional surfactant is selected from the group consisting of sodium methyl lauryl taurate, sodium methyl oleoyl taurate, sodium methyl cocoyl taurate and mixtures thereof. The additional surfactant may be sodium methyl cocoyl taurate.

[0046] The ratio of the solvent system to the total amount of surfactant in the composition may be from 360:1 (or about 360:1) to 10:1 (or about 10:1), optionally from 100:1 (or about 100:1) to 20:1 (or about 20:1) by weight.

[0047] The ratio of the oil phase to the solvent system to the total amount of surfactant (or, the total surfactant concentration including the alkyl sulfate surfactant) by weight may be 1:200:1.5 (or about 1:200:1.5) or, optionally, 1:60:1.5 (or about 1:60:1.5) by weight.

Aqueous Phase

[0048] An aqueous phase comprising water is added to the oil phase of the present compositions to form oil-in-water or water-in-oil dispersions, micro emulsions or emulsions. The aqueous phase may comprise from about 60% to about 95%, or optionally from about 75% to about 93%, by weight of the composition.

[0049] Alternatively, the mouth rinse compositions of the present invention may be formulated in a dry powder, chewing gum, semi-solid, solid or liquid concentrate form. In such embodiments, for example, water is added to q.s. as necessary in the case of liquid concentrates or powdered formulations, or water may be removed using standard evaporation procedures known in the art to produce a composition in dry powder form. Evaporated, or freeze dried forms are advantageous for storage and shipping.

Micelle Size

[0050] The mouth rinse compositions of the present invention comprise colloidal aggregates of amphipathic molecules called micelles. The "aggregate size" of the micelles of the compositions of the present invention is equivalent to the "Z-average micelle dH" of the compositions of the present invention, which is equivalent to the "Z-average aggregate micelle size" of the compositions of the present invention. The micelles of the present invention have an aggregate size of less than 200 nm (or about 200 nm), optionally less than 100 nm (or about 100 nm), optionally less than 50 nm (or about 50 nm), or optionally less than 25 nm (or about 25). In order to measure the aggregate micelle size of the compositions of the present invention, samples were filtered and analyzed using a Zetasizer Nano ZS DLS instrument (Malvern Instruments, Inc., Southborough, MA) operating at 25.0 °C. The values of micelle dH and the micelle size distribution were

calculated using the Dispersion Technology Software (DTS) v4.10 package (Malvern Instruments Inc., Southborough, MA), which calculates the Z-average micelle dH according to the ISO13321 test method.

Optional Ingredients

Insoluble Particulates

[0051] The oral care compositions of the present invention may optionally comprise a safe and effective amount of a water insoluble particulate. The water insoluble particulate can be an abrasive particle (such as a dentally acceptable abrasive) or non-abrasive particulate.

[0052] Optional, dentally acceptable abrasives include, but are not limited to, water insoluble calcium salts such as calcium carbonate, and various calcium phosphates, alumina, silica, synthetic resins and mixtures thereof. Suitable dentally acceptable abrasives may generally be defined as those having a radioactive dentine abrasion value (RDA) of from about 30 to about 250 at the concentrations used in the compositions of the present invention. The abrasives may be non-crystalline, hydrated, silica abrasives, particularly in the form of precipitated silica or milled silica gels available commercially, for example, under the trade names ZEODENT (J. M. Huber Corporation, Edison, NJ), and SYLODENT (W.R. Grace & Co., New York, NY), respectively. The compositions comprise from about 1% to about 20%, or, optionally, from about 5% to about 10% by weight of the abrasive.

[0053] Alternatively, the insoluble particulate is a non-abrasive particulate which is visible to the unaided eye and stable in the compositions of the present invention.

[0054] The non-abrasive particulate can be of any size, shape, or color, according to the desired characteristic of the product. The non-abrasive particulates will typically have the shape of a small round or substantially round ball or sphere, however, platelet or rod-shaped configurations are also contemplated herein. Generally, a non-abrasive particulate has an average diameter of from about 50 μm to about 5000 μm , optionally from about 100 μm to about 3000 μm , or optionally from about 300 μm to about 1000 μm . By the terms "stable" and/or "stability", it is meant that the abrasive or non-abrasive particulates are not disintegrated, agglomerated, or separated under normal shelf conditions. The terms "stable" and/or "stability" may further mean that the compositions of present invention contain no visible or minimally visible (to the unaided eye) signs of sedimentation of the insoluble particulates after 8 weeks, optionally 26 weeks, optionally 52 weeks, at room temperature.

[0055] The non-abrasive particulates herein are typically incorporated in the present compositions at levels of from about 0.01% to about 25%, optionally, from about 0.01% to about 5%, or optionally, from about 0.05% to about 3%, by weight of the composition.

[0056] The non-abrasive particulate herein will typically comprise a structural material and/or, optionally, an encompassed material.

[0057] The structural material provides a certain strength to the non-abrasive particulates so that they retain their distinctively detectable structure in the compositions of the present invention under normal shelf conditions. In one embodiment, the structural material further can be broken and disintegrated with very little shear on the teeth, tongue or oral mucosa upon use.

[0058] The non-abrasive particulates can be solid or liquid, filled or un-filled, as long as they are stable in the compositions of the present invention. The structural material used for making the non-abrasive particulates varies depending on the compatibility with other components, as well as material, if any, to be encompassed in the non-abrasive particulates. Exemplary materials for making the non-abrasive particulates herein include: polysaccharide and saccharide derivatives such as crystalline cellulose, cellulose acetate, cellulose acetate butyrate, cellulose acetate phthalate, cellulose nitrate, ethylcellulose, hydroxypropylcellulose, hydroxypropylmethylcellulose, hydroxypropylmethylcellulose phthalate, methyl cellulose, sodium carboxymethylcellulose, gum acacia (gum arabic), agar, agarose, maltodextrin, sodium alginate, calcium alginate, dextran, starch, galactose, glucosamine, cyclodextrin, chitin, amylose, amylopectin, glycogen, laminaran, lichenan, curdlan, inulin, levan, pectin, mannan, xylan, alginic acid, arabic acid, glucomannan, agarose, agarpectin, prophyran, carrageen, fucoidan, glycosaminoglycan, hyaluronic acid, chondroitin, peptidoglycan, lipopolysaccharide, guar gum, starch, and starch derivatives; oligosaccharides such as sucrose, lactose, maltose, uronic acid, muramic acid, cellobiose, isomaltose, planteose, melezitose, gentianose, maltotriose, stachyose, glucoside and polyglucoside; monosaccharides such as glucose, fructose, and mannose; synthetic polymers such as acrylic polymers and copolymers including polyacrylamide, poly(alkyl cyanoacrylate), and poly(ethylene-vinyl acetate), and carboxyvinyl polymer, polyamide, poly(methyl vinyl ether-maleic anhydride), poly(adipyl-L-lysine), polycarbonate, polyterephthalamide, polyvinyl acetate phthalate, poly(terephthaloyl-L-lysine), polyarylsulfone, poly(methylmethacrylate), poly(ϵ -caprolactone), polyvinylpyrrolidone, polydimethylsiloxane, polyoxyethylene, polyester, polyglycolic acid, polylactic acid, polyglutamic acid, polylysine, polystyrene, poly(styrene-acrylonitrile), polyimide, and poly(vinyl alcohol); and other material such as fat, fatty acid, fatty alcohol, milk solids, molasses, gelatin, gluten, albumin, shellac, caseinate, bees wax, carnauba wax, spermaceti wax, hydrogenated tallow, glycerol monopalmitate, glycerol dipalmitate, hydrogenated castor oil, glycerol monostearate, glycerol distearate, glycerol tristearate, 12-hydroxystearyl alcohol, protein, and protein derivatives; and mixtures thereof. Components herein may be described in other sections as useful components for the present composition. The components as described in this section may form the structure of the non-abrasive particulates so as to not be substantially dissolved or dispersed from the particulates and into the compositions of the present invention under normal shelf conditions.

[0059] In other embodiments, The structural material herein may comprise components selected from the group consisting of polysaccharides and their derivatives, saccharides and their derivatives, oligosaccharides, monosaccharides, and mixtures thereof, or optionally, comprises components are having various degrees of water solubility. In some embodiments, the structural material comprises lactose, cellulose, and hydroxypropyl methylcellulose.

[0060] Suitable non-abrasive particulates also include organogel particles as described in detail in U.S. Pat. No. 6,797,683 . Non-abrasive particulates that are organogel particles typically comprise a structural material selected from waxes (e.g., beeswax, paraffin, water-insoluble wax, carbon-based wax, silicone wax, microcrystalline wax, etc.), triglycerides, acid triglycerides, polymers, fluoroalkyl (meth)acrylate polymers and copolymers, acrylate polymers, ethylene/acrylate copolymers, polyethylene, polypropylene polymers and copolymers, fatty acids, fatty alcohols, fatty acid esters, fatty acid ethers, fatty acid amides, alkylene polyhydric alcohols, fatty acid amide of an alkanolamine, glyceryl monostearate, (aryl-substituted)sugars, dibenzyl sorbitol (or mannitol, rabbitol, etc.), condensates and precondensates of lower monohydric alcohols, trihydric alcohols, lower polyglycols, propylene/ethylene polycondensates, and the like. Optionally, structural material for non-abrasive particulates that are organogel particles include beeswax, carnauba wax, low molecular weight ethylene homopolymers (e.g. Polywax 500, Polywax 1000, or Polywax 2000 polyethylene materials available from Baker Petrolite Corp.), or paraffin wax.

[0061] The non-abrasive particulates herein may encompass, contain, or be filled with an encompassed material. Such encompassed material can be water soluble or water insoluble. Suitable encompassed

materials include benefit agents as described herein such as: oral care actives, vitamins, pigments, dyes, antimicrobial agents, chelating agents, optical brighteners, flavors, perfumes, humectants, minerals, and mixtures thereof. The encompassed materials herein are substantially retained within the non-abrasive particulates, and are substantially not dissolved from the particulates and into the compositions of the present composition under normal shelf conditions.

[0062] Particularly useful commercially available non-abrasive particulates herein are those with tradenames Unisphere and Unicerin available from Induchem AG (Switzerland), and Confetti Dermal Essentials available from United-Guardian Inc. (NY, USA). Unisphere and Unicerin particles are made of microcrystalline cellulose, hydroxypropyl cellulose, lactose, vitamins, pigments, and proteins. Upon use, the Unisphere and Unicerin particles can be disintegrated with very little shear and with practically no resistance, and readily disperse in the compositions of the present invention.

[0063] Suitable non-abrasive particulates for incorporation in the present compositions are described in detail in U.S. Pat. No. 6,797,683 (organogel particles); U.S. Pat. No. 6,045,813 (rupturable beads); U.S. Pat. Publ. 2004/0047822 A1 (visible capsules); and U.S. Pat. No. 6,106,815 (capsulated or particulated oily substances), .

[0064] The abrasive and/or nonabrasive particles may have a density different or, optionally, substantially different from the carrier in which these particles are formulated.

Flavors or Flavorants

[0065] Flavors or flavorants may also optionally be added to further modify or magnify the taste of the mouth rinse, or reduce or mask the sharp "bite" or "burn" of ingredients such as thymol. Suitable flavors include, but are not limited to, oil of anise, anethole, benzyl alcohol, spearmint oil, citrus oils, vanillin and the like may be incorporated. The amount of flavor oil added to the composition can be from 0.001% (or about 0.001%) to 1.0% (or about 1.0%) w/v, or optionally from 0.1% (or about 0.10%) to 0.30% (or about 0.30%) w/v of the composition.

[0066] The particular flavors or flavorants, and other taste-improving ingredients, employed will vary depending upon the particular taste and feel desired. Those skilled in the art can select and customize these types of ingredients to provide the desired results.

Buffers

[0067] Additional conventional components may be added as in mouthwashes and mouth rinses of the prior art. Whereas some alcohol containing mouth rinses have a pH of about 7.0, reduction of the alcohol level may require the addition of acidic preservatives, such as sorbic acid or benzoic acid, which reduce pH levels. Buffer systems are then necessary to control the pH of the composition at optimal levels. This is generally accomplished through the addition of a weak acid and its salt or a weak base and its salt. The present invention comprises useful systems selected from the group consisting of sodium benzoate and benzoic acid in amounts of from 0.01% (or about 0.01% w/v) to 1.0% w/v (or about 1.0% w/v) of the composition, and sodium citrate and citric acid in amounts of from 0.001% (or about 0.001% w/v) to 1.0% w/v (or about 1.0% w/v) of the composition, phosphoric acid and sodium/potassium phosphate of amounts from 0.01% (or about 0.01%) to 1.0% (or about 1.0%) by weight of the composition. The buffers may be incorporated in amounts that maintain the pH at levels of from 3.0 (or about 3.0) to 8.0 (or about 8.0), optionally from 3.5 (or about 3.5) to 6.5 (or about 6.5), optionally from 3.5 (or about 3.5) to 5.0 (or about

5.0). Without being limited by any theory, it is believed that these pH levels provide the essential oils with an environment that also maximizes their antimicrobial activity and promotes stability.

Fluoride Releasing Compounds

[0068] Fluoride providing compounds may optionally be present in the mouth rinse compositions of this invention. These compounds may be slightly water soluble or may be fully water soluble and are characterized by their ability to release fluoride ions or fluoride containing ions in water. Typical fluoride providing compounds are inorganic fluoride salts such as soluble alkali metal, alkaline earth metal, and heavy metal salts, for example, sodium fluoride, potassium fluoride, ammonium fluoride, cupric fluoride, zinc fluoride, stannic fluoride, stannous fluoride, barium fluoride, sodium hexafluorosilicate, ammonium hexafluorosilicate, sodium fluorozirconate, sodium monofluorophosphate, aluminum mono- and difluorophosphate and fluorinated sodium calcium pyrophosphate. Amine fluorides, such as N'-octadecyltrimethylenediamine-N,N,N'-tris(2-ethanol)-dihydrofluoride and 9-octadecenylamine-hydrofluoride, may also be used.

[0069] The fluoride providing compound may generally be present in an amount sufficient to release up to 0.15% (or about 0.15%), optionally 0.001% (or about 0.001%) to 0.1% (or about 0.1%), optionally from 0.001% (or about 0.001%) to 0.05% (or about 0.05%) fluoride by weight of the composition.

Zinc Salts

[0070] Zinc salts such as zinc chloride, zinc acetate or zinc citrate may optionally be added as an astringent for an "antiseptic cleaning" feeling, as a breath protection enhancer or as anticalculus agent in an amount of from 0.0025% w/v (or about 0.0025% w/v) to 0.1% w/v (or about 0.1% w/v) of the composition.

Sensitivity Reducing Agents

[0071] Sensitivity reducing agents, namely potassium salts of nitrate and oxalate in an amount from 0.1% (or about 0.1%) to 5.0% (or about 5.0%) w/v of the composition may be incorporated into the present invention. Other potassium releasing compounds are feasible (e.g. KCl). High concentrations of calcium phosphates may also provide some added sensitivity relief. These agents are believed to work by either forming an occlusive surface mineral deposit on the tooth surface or through providing potassium to the nerves within the teeth to depolarize the nerves. A more detailed discussion of suitable sensitivity reducing can be found in US 20060013778 to Hodosh and US Pat. 6,416,745 to Markowitz et al.,

Anticalculus Agents

[0072] Compounds with anti-calculus benefits (e.g. polyphosphates, phosphonates, various carboxylates, polyaspartic acid, inositol phosphate etc.) may optionally be incorporated into the present invention. Also useful as an anticalculus agent the anionic polymeric polycarboxylates. Such materials are well known in the art, being employed in the form of their free acids or partially or preferably fully neutralized water soluble alkali metal (e.g. potassium and preferably sodium) or ammonium salts. Preferred are 1:4 to 4:1 by weight copolymers of maleic anhydride or acid with another polymerizable ethylenically unsaturated monomer, preferably methyl vinyl ether (methoxyethylene) having a molecular weight (M.W.) of about 30,000 to about

1,000,000. These copolymers are available for example as Gantrez AN 139 (M.W. 500,000), AN 119 (M.W. 250,000) and preferably S-97 Pharmaceutical Grade (M.W. 70,000), of GAF Chemicals Corporation.

Additional Ingredients

[0073] Although the mouth rinse compositions of the present invention may be formulated to be substantially clear and/or colorless to the unaided eye, acceptably approved food dyes are preferably used to provide a pleasing color to the compositions of the invention. These may be selected from, but not limited to, the long list of acceptable food dyes. Suitable dyes for this purpose include FD&C yellow #5, FD&C yellow #10, FD&C blue #1 and FD&C green #3. These are added in conventional amounts, typically in individual amounts of from 0.00001% w/v (or about 0.00001% w/v) to 0.0008% w/v (or about 0.0008% w/v), optionally from 0.00035% w/v (or about 0.00035% w/v) to 0.0005% w/v (or about 0.0005% w/v) of the composition.

[0074] Other conventional ingredients may be used in the mouth rinse compositions of this invention, including those known and used in the art. Examples of such ingredients include thickeners, suspending agents and softeners. Thickeners and suspending agents useful in the compositions of the present invention can be found in US Pat. No. 5,328,682 to Pullen et al., These may be incorporated in amounts of from 0.1% w/v (about 0.1% w/v) to 0.6% w/v (or about 0.6% w/v), optionally from 0.1% w/v (about 0.1% w/v) to 0.5% w/v (or about 0.5% w/v) of the composition.

[0075] A more detailed description of useful oral care actives and/or inactive ingredients and further examples thereof can be found in US Patents 6,682,722 to Majeti et al. and 6,121,315 to Nair et al.

[0076] The compositions of the present invention are preferably free of or essentially free of bioavailability affecting compounds. As used herein, "bioavailability affecting compound", means compounds that negatively affect the bioavailability of any incorporated essential oils such as by binding the essential oils or otherwise inactivating the essential oils. "Essentially free" as used with respect to bioavailability affecting compounds is defined as formulations having less than 5% (or about 5%), optionally, less than 3% (or about 3%), optionally, less than 1% (or about 1%), or optionally less than 0.1% (or about 0.1%), or optionally, less than 0.01% (or about 0.01%), by weight (w/v) of the total composition of a bioavailability affecting compound. The bioavailability affecting compound may include, but is not limited to, polyethylene oxide/polypropylene oxide block copolymers such as poloxamers; cyclodextrins; polysorbates such as Tweens; and mixtures thereof.

Methods of Practicing the Present Invention

[0077] The invention illustratively disclosed herein may be practiced in the absence of any component, ingredient, or step which is not specifically disclosed herein.

[0078] The compositions of the present invention may be applied to teeth and/or soft surfaces of the oral cavity for at least two consecutive applications, optionally, at least (or greater than) 3 (or about 3) or optionally, at least (or greater than) 5 (or about 5) consecutive applications.

[0079] When applied to teeth and/or soft surfaces of the oral cavity, The composition may allowed to remain in contact with the teeth and/or soft surfaces of the oral cavity for at least (or greater than) 10 (or about 10) seconds, optionally at least (or greater than) 20 (or about 20) seconds, optionally at least (or greater than) than 30 (or about 30) seconds, optionally (at least or greater than) 50 (or about 50) seconds, or optionally at

least (or greater than) 60 (or about 60) seconds.

[0080] Various embodiments of the invention have been set forth above. Each embodiment is provided by way of explanation of the invention, not limitation of the invention. In fact, it will be apparent to those skilled in the art that various modifications and variations can be made in the present invention without departing from the scope of the invention. For instance, features illustrated or described as part of one embodiment, can be used on another embodiment to yield a still further embodiment. Thus, it is intended that the present invention cover such modifications and variations as come within the scope of the appended claims and their equivalents.

EXAMPLES

[0081] The following examples are illustrative only and should not be construed as limiting the invention in any way. Those skilled in the art will appreciate that variations are possible which are within the scope of the appended claims.

Example 1: Solubilization of Essential Oils with Cyclodextrins (Comparative Example)

[0082] Four formulations were prepared and tested for mixed-species biofilm kill analysis using a flow cell model. A 24-hour salivary biofilm was grown on a polystyrene peg plate (96 pegs, N=24 per test group) in an aerobic incubator set at a temperature of 30.5°C. The growth media was BM media*. The pegs were subsequently treated for thirty seconds with 185-μl of each of the four formulations as well as positive and negative controls. Treatments occurred twice daily for a total of five treatments. The positive control was a commercially available essential oil mouth rinse. The negative control was sterile water.

[0083] After treatment, viable bacteria remaining on the substrate were removed by sonication using a Misonix XL-2000 Ultrasonic processor (Qsonica, LLC, Newtown, CT) for a total of 20 seconds using power level 6. During sonication, the sonic energy is pulsed by turning on sonication for 3 seconds and then off for 1 second until all 20 seconds have elapsed.

[0084] Using a Celsis Rapid Detection RapiScreen kit (Celsis International PLC, Chicago), the bacteria was lysed with Celsis Luminex for 10 seconds and then the adenosine triphosphate (ATP) released from the lysed bacteria was measured at ambient temperature with a Berthold Centro LB 960 microplate luminometer using the bioluminescence marker Celsis LuminATE.

[0085] Data was reported in log RLU (relative light units) where decreasing log RLUs indicates fewer viable bacteria remaining on the substrate and in M-factor units where M-factor is the difference between the log RLU values of the compound tested and the negative control.

***BM media, made as follows:**

[0086]

- Proteose Peptone: 10-grams/liter (Bacto)
- Trypticase Peptone (Tryptone): 5-grams/liter (Bacto)
- Yeast Extract: 5-grams/liter (DIFCO)

- Potassium Chloride: 2.5-grams/liter (SIGMA CAS# 7447-40-7)
- Cysteine Hydrochloride: 0.5-grams/liter
- Add purified water to dry ingredients and stir until dissolved. QS to appropriate volume with water. Sterilize by autoclaving at 121°C for 20 minutes.
- After autoclaving add: 1.0ml Hemin/liter (SIGMA 16009-13-5)
- Add 0.2mL Menadione (Vitamin K) stock solution/liter (SIGMA CAS# 57414-02-5)

[0087] The four formulations, as well as results of the mixed-species biofilm kill tests, are shown on Table 1. Final formulations were adjusted to pH 4.2 with 0.1M NaOH or 0.1M HCl if necessary.

Table 1

Ingredients	Formulations					
	1A (% w/w)	1B (% w/w)	1C (% w/w)	1D (% w/w)	Positive Control (%w/w)	Negative Control (%w/w)
Purified Water USP	QS	QS	QS	QS	QS	100
Hydroxypropyl Cyclodextrin	2.00	2.00	0.00	2.00	--	--
Poloxamer 407	--	--	--	--	0.25	--
Thymol	0.06	0.06	--	--	0.0636	--
Menthol	0.04	0.04	--	--	0.0420	--
Eucalyptol	0.09	0.09	--	--	0.0922	--
Methyl Salicylate	0.06	0.06	--	--	0.060	--
Glycerin	5.00	5.00	5.00	5.00	--	--
Sorbitol (70% solution)	5.00	5.00	5.00	5.00	20.0	-
Benzoic Acid	--	--	--	--	0.12	--
Citric Acid	0.10	0.10	0.10	0.10	--	--
NaOH	0.25	0.25	0.25	0.25	--	--
Ethanol - 200 proof	--	20.00	20.00	--	21.6 (v/v)	--
Flavor	--	--	--	--	0.085	--
Sweetener	--	--	--	--	0.117	--
Color	--	--	--	--	0.0005	--
Total	100.00	100.00	100.00	100.00	100.00	--
log RLU	7.50	8.00	8.09	7.94	5.67	7.54
M-factor	0.04	-0.46	-0.55	-0.40	1.87	0

[0088] The M-factors for formulations 1A through 1D were less than 0.1. This data indicates that the essential oils, though solubilized in the cyclodextrin formulas, are not sufficiently bioavailable to kill bacterial species in a biofilm. The commercially available alcohol containing essential oil mouth rinse provided a relatively high M-factor with M-factor equal to 1.87.

Example 2: Various Surfactants

[0089] Seven propylene-glycol based mouth rinse formulations were prepared using various surfactants that are approved for use in oral care products and tested using an *in-vitro* single species *S. mutans* biofilm model. *S. mutans* was grown in Brain Heart Infusion broth for 24 hours at 33°C and then transferred to Jordan's media with 3.5% sucrose (Jordan, H.V. et al. (1960) J Dent. Res. 39:116-123) and 190 µL was added to each well of a 96 well plate (N=32 per test formulation). The plate was then incubated at 33°C for 20 hours. After 20 hours, the spent media was removed and 190 µL of fresh Jordan's media with 3.5% sucrose was added to each well and two hours later, the media was removed and the *S. mutans* biofilm remaining in each well was exposed to 200 µL of the test formulation for 30 seconds. The positive control was a commercially available essential oil mouth rinse. The negative control was sterile water.

[0090] After treatment, remaining biofilm was neutralized and removed by sonication using a Misonix XL-2000 Ultrasonic processor (Qsonica, LLC, Newtown, CT) for a total of 20 seconds using power level 7. During sonication, the sonic energy was pulsed by turning on sonication for 3 seconds and then off for 1 second until all 20 seconds have elapsed.

[0091] Using a Celsis Rapid Detection RapiScreen kit (Celsis International PLC, Chicago), the bacteria was lysed with Celsis Luminex for 10 seconds and then the adenosine triphosphate (ATP) released from the lysed bacteria was measured at ambient temperature with a Berthold Centro LB 960 microplate luminometer using the bioluminescence marker Celsis LuminATE.

[0092] Data was reported in log RLU (relative light units) where decreasing log RLUs indicates fewer viable bacteria remaining on the substrate and in M-factor units where M-factor was the difference between the log RLU values of the compound tested and the negative control.

[0093] The seven formulations, as well as results of the *S. mutans* biofilm kill tests, in log RLU and M-Factor units, are shown on Table 2. Final formulations were adjusted to pH 4.2 with 0.1M NaOH or 0.1M HCl if necessary. A typical M-factor for a commercially available alcohol containing essential oil mouth rinse is about 2.1 (log RLU of 5.8) in this model.

Table 2

Ingredients	Formulations							
	2A* (% w/w)	2B* (% w/w)	2C* (% w/w)	2D* (% w/w)	2E* (% w/w)	2F (% w/w)	2G* (% w/w)	Negative Control (% w/w)
Propylene glycol	10.0	10.0	10.0	10.0	10.0	10.0	10.0	100
Benzoic Acid	0.30	0.30	0.30	0.30	0.30	0.30	0.30	---
Menthol	0.042	0.042	0.042	0.042	0.042	0.042	0.042	---
Methyl salicylate	0.060	0.060	0.060	0.060	0.060	0.060	0.060	---
Thymol	0.064	0.064	0.064	0.064	0.064	0.064	0.064	---
Eucalyptol	0.092	0.092	0.092	0.092	0.092	0.092	0.092	---
Tween 20	---	2.00	---	---	---	---	---	---
Sodium Lauryl Sulfate	0.30	---	1.00	---	---	0.30	0.15	---
Cocamidopropyl betaine	---	---	---	1.00	---	---	---	---
Hexadecyl								

Ingredients	Formulations							
	2A* (% w/w)	2B* (% w/w)	2C* (% w/w)	2D* (% w/w)	2E* (% w/w)	2F (% w/w)	2G* (% w/w)	Negative Control (% w/w)
sulfate	---	---	---	---	0.30	---	0.15	---
Sorbitol (70% solution)	---	---	---	---	---	10.00	---	---
Sodium Benzoate	0.30	0.30	0.30	0.30	0.30	0.30	0.30	---
Purified Water USP	QS	QS	QS	QS	QS	QS	QS	---
TOTAL	100.0	100.0	100.0	100.0	100.0	100.0	100.0	-
log RLU	5.63	8.02	5.30	7.73	6.50	4.52	4.22	7.90
M-factor	2.27	-0.12	2.60	0.17	1.40	3.38	3.68	0
Z-average Aggregate Micelle size (nm)	6.1	12.9	16.1	22.1	Not available	20.1	11.2	N/A
*Formulations 2A-2E and 2G are provided as comparative formulations. Table 2 shows biocidal activity ranges significantly depending on the identity of the surfactant, with alkyl sulfates and alkyl sulfate combinations displaying the highest activity (M-factor greater than 2.0.) Additionally, it was observed that sorbitol increases bioactivity as well, as otherwise identical formulation 2A without sorbitol provided M-factor of 2.27 while formulation 2F with sorbitol provided a much greater M-factor of 3.68.								

Example 3: Optimization of Solvent Level in a Low Alcohol Mouth Rinse

[0094] To optimize solvent (ethanol, sorbitol, propylene glycol) levels for low-alcohol essential oil mouth rinse formulations, a central composite design was used for these three factors. Fifteen formulations were prepared and tested using the single-species *S. mutans* model described in Example 2, and log RLU was monitored to evaluate bioactivity. Sterile water was used as the negative control.

[0095] The fifteen formulations, as well as results of the *S. mutans* biofilm kill tests, are shown on Tables 3A and 3B. Final formulations were adjusted to pH 4.2 with 0.1M NaOH or 0.1M HCl if necessary. A typical M-factor for a commercially available alcohol containing essential oil mouth rinse is about 2.1 (log RLU of 5.8) in this model.

Table 3A

Ingredients	Formulations							
	3A* (% w/w)	3B* (% w/w)	3C* (% w/w)	3D* (% w/w)	3E* (% w/w)	3F* (% w/w)	3G* (% w/w)	3H* (% w/w)
Propylene glycol	2.82	2.82	5.50	5.50	5.50	8.17	8.17	5.50
Cremophor (PEG Castor Oil)	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Ethanol	2.02	7.97	10.00	5.00	5.00	2.03	2.02	5.00
Benzoic	0.08	0.08	0.08	0.0	0.08	0.08	0.08	0.08

Ingredients	Formulations							
	3A* (% w/w)	3B* (% w/w)	3C* (% w/w)	3D* (% w/w)	3E* (% w/w)	3F* (% w/w)	3G* (% w/w)	3H* (% w/w)
Acid								
Menthol	0.042	0.042	0.042	0.042	0.042	0.042	0.042	0.042
Methyl salicylate	0.060	0.060	0.060	0.060	0.060	0.060	0.060	0.060
Thymol	0.064	0.064	0.064	0.064	0.064	0.064	0.064	0.064
Eucalyptol	0.092	0.092	0.092	0.092	0.092	0.092	0.092	0.092
Sodium Lauryl Sulfate	0.30	0.30	0.30	0.30	0.30	0.30	0.30	0.30
Sorbitol (70% solution)	15.94	4.05	10.00	10.00	---	4.05	15.94	20.00
Sodium Benzoate	0.07	0.07	0.07	0.07	0.07	0.07	0.07	0.07
Flavor	0.30	0.30	0.30	0.30	0.30	0.30	0.30	0.30
Saccharin	0.12	0.12	0.12	0.12	0.12	0.12	0.12	0.12
FD&C Green #3	0.0005	0.0005	0.0005	0.0005	0.0005	0.0005	0.0005	0.0005
Purified Water USP	QS	QS	QS	QS	QS	QS	QS	QS
TOTAL	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
log RLU	5.08	4.91	4.38	4.44	4.77	4.67	4.64	4.86
M-factor	2.82	2.99	3.52	3.46	3.13	3.23	3.26	3.04

Table 3B

Ingredients	Formulations							
	3I* (% w/w)	3J* (% w/w)	3K* (% w/w)	3L (% w/w)	3M* (% w/w)	3N* (% w/w)	3O* (% w/w)	Negative Control (% w/w)
Propylene glycol	2.82	8.18	8.18	5.50	2.82	1.00	10.00	100
Cremophor (PEG Castor Oil)	0.1	0.1	0.1	0.1	0.1	0.1	0.1	--
Ethanol	2.02	7.97	7.97	-	7.97	5.00	5.00	--
Benzoic Acid	0.08	0.08	0.08	0.08	0.08	0.08	0.08	--
Menthol	0.042	0.042	0.042	0.042	0.042	0.042	0.042	--
Methyl salicylate	0.060	0.060	0.060	0.060	0.060	0.060	0.060	--
Thymol	0.064	0.064	0.064	0.064	0.064	0.064	0.064	--
Eucalyptol	0.092	0.092	0.092	0.092	0.092	0.092	0.092	--
Sodium Lauryl								--

Ingredients	Formulations							
	3I* (% w/w)	3J* (% w/w)	3K* (% w/w)	3L (% w/w)	3M* (% w/w)	3N* (% w/w)	3O* (% w/w)	Negative Control (% w/w)
Sulfate	0.30	0.30	0.30	0.30	0.30	0.30	0.30	
Sorbitol (70% solution)	4.05	4.05	15.95	10.00	15.94	10.00	10.00	--
Sodium Benzoate	0.07	0.07	0.07	0.07	0.07	0.07	0.07	--
Flavor	0.30	0.30	0.30	0.30	0.30	0.30	0.30	--
Saccharin	0.12	0.12	0.12	0.12	0.12	0.12	0.12	-
FD&C Green #3	0.0005	0.0005	0.0005	0.0005	0.0005	0.0005	0.0005	--
Purified Water USP	QS	QS	QS	QS	QS	QS	QS	--
TOTAL	100.0	100.0	100.0	100.0	100.0	100.0	100.0	-
log RLU	4.56	4.67	4.45	4.58	4.91	4.70	4.62	7.9
M-factor	3.34	3.29	3.45	3.32	2.99	3.2	3.28	0

*Formulations 3A-3K and 3M-3N are provided as comparative formulations. As illustrated in Tables 3A and 3B, at a level of 0.3%, sodium lauryl sulfate and a total oil phase content of about 0.6% w/w, the propylene glycol (or polyol solvent) concentration is optimum between 4% and 10% w/w and sorbitol (or sugar alcohol solvent) is optimum between 6% and 14% w/w when added as a 70% sorbitol solution. Ethanol concentration did not have a large impact on efficacy in this system.

Example 4: Optimization of Solvent Levels in a Non-alcohol Mouth Rinse with a Two-factor Central Composite Design

[0096] To optimize solvent (sorbitol, propylene glycol) levels for non-alcohol essential oil mouth rinse formulations, a central composite design was used for these two factors. Nine formulations were prepared and tested using the single-species *S.mutans* model described in Example 2, and log RLU was monitored to evaluate bioactivity. Sterile water was used as the negative control.

[0097] The nine formulations, as well as results of the *S. mutans* biofilm kill tests, are shown on Table 4. Final formulations were adjusted to pH 4.2 with 0.1M NaOH or 0.1M HCl if necessary. A typical M-factor for a commercially available alcohol containing essential oil mouth rinse is about 2.1 (log RLU of 5.8) in this model.

Table 4

[illegible]

Ingredients	Formulations									
	4A (% w/w)	4B (% w/w)	4C (% w/w)	4D* (% w/w)	4E (% w/w)	4F (% w/w)	4G (% w/w)	4H (% w/w)	4I (% w/w)	Negative Control (% w/w)
Methyl salicylate	0.060	0.060	0.060	0.060	0.060	0.060	0.060	0.060	0.060	
Thymol	0.065	0.065	0.065	0.065	0.065	0.065	0.065	0.065	0.065	
Eucalyptol	0.093	0.093	0.093	0.093	0.093	0.093	0.093	0.093	0.093	
Flavor	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	
Sorbitol (70% solution)	3.66	21.34	25.00	-	21.34	12.50	12.50	3.66	12.50	
Sodium Lauryl Sulfate	0.35	0.35	0.35	0.35	0.35	0.35	0.35	0.35	0.35	
Purified Water USP	QS	QS	QS	QS	QS	QS	QS	QS	QS	
Benzoic Acid	0.086	0.086	0.086	0.086	0.086	0.086	0.086	0.086	0.086	
Sodium Benzoate	0.077	0.077	0.077	0.077	0.077	0.077	0.077	0.077	0.077	
Saccharin	0.06	0.06	0.06	0.06	0.06	0.06	0.06	0.06	0.06	
Sucralose	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	
FD&C Green #3	0.0005	0.0005	0.0005	0.0005	0.0005	0.0005	0.0005	0.0005	0.0005	
TOTAL	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	
log RLU	5.96	5.44	5.57	5.69	5.77	5.27	5.38	5.8	5.87	7.94
M-factor	1.98	2.50	2.37	2.25	2.17	2.67	2.56	2.14	2.07	0
*Formulation 4D is provided as a comparative formulation. As illustrated in Table 4, at a level of 0.35% w/w sodium lauryl sulfate, and a total oil phase content of about 0.36% w/w, the propylene glycol (or polyol solvent) concentration is optimum between 5 and 13% w/w while sorbitol (or sugar alcohol solvent) is optimum between 10 and 25% w/w when added as a 70% sorbitol solution. Solvent optimums can vary depending on surfactant concentration and flavor choice.										

Example 5: Evaluation of Other Glycol Solvents

[0098] Two formulations were prepared and tested using the single-species *S.mutans* model described in Example 2, with log RLU monitored to evaluate bioactivity. Sterile water was used as the negative control.

[0099] The purpose was to compare non-alcohol essential oil formulations containing propylene glycol to non-alcohol essential oil formulations containing other glycol solvents, such as 1,3-propanediol.

[0100] The formulations, as well as results of the *S. mutans* biofilm kill tests, are shown on Table 5. Final formulations were adjusted to pH 4.2 with 0.1M NaOH or 0.1M HCl if necessary. A typical M-factor for a

commercially available alcohol containing essential oil mouth rinse is about 2.1 (log RLU of 5.8) in this model.

Table 5

Ingredients	Formulations		
	5A (% w/w)	5B (% w/w)	Negative Control (% w/w)
Purified Water USP	QS	QS	100
Sorbitol (70% solution)	10.00	10.00	--
1,3-propanediol	7.00	--	--
Propylene glycol	--	7.00	--
Sodium lauryl sulfate	0.30	0.30	--
Sodium Saccharin	0.12	0.12	--
Flavor	0.10	0.10	--
Benzoic Acid	0.10	0.10	--
Sodium Benzoate	0.05	0.05	--
Eucalyptol	0.093	0.093	--
Thymol	0.065	0.065	--
Methyl Salicylate	0.060	0.060	--
Menthol	0.042	0.042	--
FD&C Green #3	0.0005	0.0005	--
TOTAL	100.00	100.00	--
log RLU	4.35	4.42	7.94
M-factor	3.59	3.52	0

[0101] The RLU values for ATP from oral biofilms treated with formulation 5A (1,3-propanediol as the polyol solvent) and formulation 5B (propylene glycol as the polyol solvent) are very comparable (M-factors equaling 3.59 and 3.52, respectively), indicating that propylene glycol can potentially be substituted with other polyol solvents (i.e., 1,3-propanediol) and still retain bioactivity. A typical M-factor for a commercially available alcohol containing essential oil mouth rinse is about 2.1 (log RLU of 5.8) in this model.

Example 6: Effect of poloxamer

[0102] Four formulations were prepared and tested for mixed-species biofilm kill using the single species model discussed in Example 2. The negative control was sterile water.

[0103] The four formulations, as well as results of the single species biofilm kill tests, are shown on Table 6. Final formulations were adjusted to pH 4.2 with 0.1M NaOH or 0.1M HCl if necessary. A typical M-factor for a commercially available alcohol containing essential oil mouth rinse is about 2.1 (log RLU of 5.8) in this model.

Table 6

Ingredients	Formulations				
	6A (% w/w)	6B (% w/w)	6C (% w/w)	6D (% w/w)	Negative Control (% w/w)
Propylene glycol	7.00	7.00	7.00	7.00	100

Ingredients	Formulations				
	6A (% w/w)	6B (% w/w)	6C (% w/w)	6D (% w/w)	Negative Control (% w/w)
Menthol	0.042	0.042	0.042	0.042	--
Methyl salicylate	0.060	0.060	0.060	0.060	--
Thymol	0.065	0.065	0.065	0.065	--
Eucalyptol	0.093	0.093	0.093	0.093	--
Flavor	0.100	0.100	0.100	0.100	--
Sorbitol	10.00	10.00	10.00	10.00	--
Sodium lauryl sulfate	0.35	0.20	0.20	0.20	--
Poloxamer 407	0.00	0.20	0.50	0.75	--
Purified Water USP	QS	QS	QS	QS	--
Benzoic Acid	0.086	0.086	0.086	0.086	--
Sodium Benzoate	0.077	0.077	0.077	0.077	--
Saccharin	0.060	0.060	0.060	0.060	--
Sucralose	0.01	0.01	0.01	0.01	--
FD&C Green #3	0.0005	0.0005	0.0005	0.0005	----
TOTAL	100.0	100.0	100.0	100.0	
log RLU	6.25	6.60	7.20	7.52	7.94
M-factor	1.69	1.34	0.74	0.42	0

[0104] The log RLU for water was 7.94. The table shows that as the amount of Poloxamer 407 added to the formulation was increased, the log RLUs (relative light units) increased. Increasing log RLUs indicates more viable bacteria remaining on the substrate. So, compositions free of Poloxamer are more effective on single species biofilm. The M-factors for all the formulations containing Poloxamer 407 (formulations 6B-6D) were less than 1.50.

Example 7: Evaluation of Glycerin as a glycol solvent

[0105] Six formulations were prepared and tested using the single-species *S. mutans* model described in Example 2, with log RLU monitored to evaluate bioactivity. Sterile water was used as the negative control.

[0106] The purpose was to compare non-alcohol essential oil formulations containing propylene glycol to non-alcohol essential oil formulations containing glycerin or glycerin/propylene glycol combinations.

[0107] The formulations, as well as results of the *S. mutans* biofilm kill tests, are shown on Table 7. Final formulations were adjusted to pH 4.2 with 0.1M NaOH or 0.1M HCl if necessary. A typical M-factor for a commercially available alcohol containing essential oil mouth rinse is about 2.1 (log RLU of 5.8) in this model.

Table 7

Ingredients	Formulations						
	7A* (% w/w)	7B* (% w/w)	7C* (% w/w)	7D* (% w/w)	7E* (% w/w)	7F (% w/w)	Negative Control (% w/w)
Propylene glycol	10.000	5.000	--	5.000	--	--	--
Glycerin	--	5.000	10.000	10.000	20.000	5.000	--
Sorbitol	--	--	--	--	--	10.000	--
Benzoic Acid	0.100	0.100	0.100	0.100	0.100	0.100	--
Menthol	0.042	0.042	0.042	0.042	0.042	0.042	--
Methyl salicylate	0.060	0.060	0.060	0.060	0.060	0.060	--
Thymol	0.064	0.064	0.064	0.064	0.064	0.064	--
Eucalyptol	0.092	0.092	0.092	0.092	0.092	0.092	--
Sodium lauryl sulfate	0.350	0.350	0.350	0.350	0.350	0.350	--
Sodium Benzoate	0.080	0.080	0.080	0.080	0.080	0.080	--
Purified Water USP	QS	QS	QS	QS	QS	QS	100
TOTAL	100	100	100	100	100	100	--
log RLU	6.20	6.65	6.30	6.32	6.46	6.12	7.9
M-factor	1.7	1.25	1.6	1.58	1.44	1.78	0
*Formulations 7A-7E are provided as comparative formulations.							

[0108] The log RLU values based on ATP from oral biofilms treated with both formulations are very comparable, indicating that propylene glycol can potentially be substituted with other glycol solvents and bioactivity will be maintained. The log RLU values based on ATP from oral biofilms treated with formulation 7A (propylene glycol) and formulation 7C (glycerin) are comparable (M-factors equaling 1.7 and 1.6, respectively), indicating that propylene glycol can potentially be substituted with other polyol solvents (i.e., glycerin) and still retain bioactivity.

Example 8: Combination of SLS with other surfactants

[0109] Three formulations were prepared and tested for mixed-species biofilm kill using a flow-through model, as shown in Example 1. Sterile water was used as the negative control.

[0110] The three formulations, as well as results of the mixed-species biofilm kill tests, are shown on Table 8. Final formulations were adjusted to pH 4.2 with 0.1M NaOH or 0.1M HCl if necessary. A typical M-factor for a commercially available alcohol containing essential oil mouth rinse is about 1.8 (log RLU of 5.7) in this model.

Table 8

Ingredients	Formulations			
	8A (% w/w)	8B (% w/w)	8C (% w/w)	Negative Control (% w/w)
Propylene glycol	7.00	0.00	0.00	--
Glycerin	0.00	10.00	10.00	--
Menthol	0.042	0.042	0.042	--
Methyl salicylate	0.060	0.060	0.060	--
Thymol	0.064	0.064	0.064	--
Eucalyptol	0.092	0.092	0.092	--
Flavor	0.100	0.100	0.100	--
Sorbitol	10.00	10.00	10.00	--
Sodium lauryl sulfate	0.35	0.15	0.00	--
Sodium methyl cocoyl taurate	0.00	0.15	0.30	--
Purified Water USP	QS	QS	QS	100
Benzoic Acid	0.086	0.086	0.086	--
Sodium Benzoate	0.077	0.077	0.077	--
FD&C Green #3	0.0005	0.0005	0.0005	--
TOTAL	100.0	100.0	100.0	--
log RLU	5.47	5.18	6.29	7.5
M-factor	2.03	2.32	1.21	0

[0111] The log RLU values for ATP from oral biofilms treated with sodium lauryl sulfate and sodium lauryl sulfate/sodium methyl cocoyl taurate formulations are very comparable, indicating that the sodium lauryl sulfate level can be reduced when an additional surfactant (e.g, sodium methyl cocoyl taurate) is added to the formulation (as indicated by M-factors of 2.03 and 2.32, respectively) and bioactivity will be maintained. If no alkyl sulfate surfactant is present, a loss of bioactivity is observed (as indicated by the lower M-factor of 1.21).

REFERENCES CITED IN THE DESCRIPTION

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- **JORDAN, H.V. et al.**J Dent. Res., 1960, vol. 39, 116-123 [0089]

Patentkrav**1.** Antimikrobiel mundskylssammensætning omfattende:

- a. polyolopløsningsmiddel valgt fra gruppen bestående af polyvalente alkaner, polyvalente alkanestere, polyalkenglycoler og blandinger deraf;
- 5 b. sukkeralkoholopløsningsmiddel valgt fra gruppen bestående af sorbitol, xylitol, mannitol, maltitol, inositol, allitol, altritol, dulcitol, galactitol, glucitol, hexitol, iditol, pentitol, ribitol, erythritol og blandinger deraf, hvor den samlede mængde af sukkeralkoholopløsningsmiddel ikke overstiger 17% vægt/volumen af sammensætningen;
- 10 c. alkylsulfat-overfladeaktivt middel, og eventuelt et yderligere overfladeaktivt middel valgt fra gruppen bestående af anioniske overfladeaktive midler, non-ioniske overfladeaktive midler, amfoteriske overfladeaktive midler og blandinger deraf;
- 15 d. buffersystem valgt fra gruppen bestående af natriumbenzoat og benzoesyre i mængder på fra 0,01% til 1,0% vægt/volumen af sammensætningen; og natriumcitrat og citronsyre i mængder på fra 0,001% til 1,0% vægt/volumen af sammensætningen; og phosphorsyre og natrium-/kaliumphosphat i mængder fra 0,01 til 1,0 vægt-% af sammensætningen;
- 20 e. en eller flere antimikrobielle æteriske olier; og
f. vand;

hvor nævnte sammensætning er fri for monovalente C₂-C₄-alkoholer.

2. Sammensætningen ifølge krav 1, hvor nævnte polyolopløsningsmiddel omfatter
25 propylenglycol.

3. Sammensætningen ifølge et hvilket som helst foregående krav, hvor nævnte sukkeralkoholopløsningsmiddel omfatter sorbitol.

30 **4.** Sammensætningen ifølge et hvilket som helst foregående krav, hvor nævnte alkylsulfat-overfladeaktivt middel omfatter natriumlaurylsulfat.

5. Sammensætningen ifølge et hvilket som helst foregående krav, hvor nævnte buffersystem omfatter natriumbenzoat og benzoesyre.

6. Sammensætningen ifølge et hvilket som helst foregående krav, hvor nævnte
5 ene eller flere antimikrobielle æteriske olier er valgt fra gruppen bestående af menthol, thymol, eucalyptol, methylsalicylat og kombinationer af to eller flere deraf.