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(54) Title: ANTIVIRAL DISINFECTANTS AND APPLICATIONS

(57) Abstract: This invention provides a method of inactivating non-enveloped virus particles. The method includes the step of contacting the virus with a virucidally-enhanced alcoholic composition that includes an alcohol, and an enhancer selected from the group consisting of cationic oligomers and polymers, proton donors, chaotropic agents, and mixtures thereof.



WO 2007/095008 A3

ANTIVIRAL METHOD

RELATED APPLICATIONS

[01] This application is a continuation-in-part of U.S. Patent Application Serial No. 11/499,227, filed on August 7, 2006, which claims priority from U.S. Provisional Patent Application Serial No. 60/771,744, filed on February 9, 2006, both of which are hereby incorporated by reference.

TECHNICAL FIELD

[01] The present invention relates to a method for inactivating non-enveloped viruses. The invention provides a method for producing a topical virucidal effect on mammalian skin against non-enveloped virus. A method for enhancing the efficacy of alcohol against non-enveloped viruses is also provided.

BACKGROUND OF THE INVENTION

[02] Skin disinfectants containing one or more lower alcohols are widely known. Disinfectants containing at least about 50 weight percent alcohol exhibit antibacterial efficacy, however the antiviral efficacy of these alcohol disinfectants depends upon the type of virus.

[03] Pathogenic viruses can be classified into two general types with respect to the viral structure: enveloped viruses and non-enveloped viruses. Some well known enveloped viruses include herpes virus, influenza virus; paramyxovirus, respiratory syncytial virus, corona virus, HIV, hepatitis B virus, hepatitis C virus, SARS-CoV, and toga virus. Non-enveloped viruses, sometimes referred to as "naked" viruses, include the families *Picornaviridae*, *Reoviridae*, *Caliciviridae*, *Adenoviridae* and *Parvoviridae*. Members of these families include rhinovirus, poliovirus, adenovirus, hepatitis A virus, norovirus, papillomavirus, and rotavirus.

[04] It is known in the art that "enveloped" viruses are relatively sensitive and, thus, can be inactivated by commonly used disinfectants. In contrast, non-enveloped viruses are substantially more resistant to conventional disinfectants and are more environmentally stable than enveloped viruses. Although a number of non-enveloped viruses can be inactivated with

relatively high concentrations of formaldehyde, the use of formaldehyde is undesirable because of its toxicity.

[05] The antiviral efficacy of acid-containing disinfectants, and of disinfectants having an acidic pH, depends upon the type of virus. A few non-enveloped viruses, namely
5 rhinovirus, feline calicivirus, and canine calicivirus, are believed to be at least somewhat affected by acid. See *Virus Taxonomy: VIIIth Report of the International Committee On Taxonomy of Viruses*, Elsevier Science & Technology Books, ISBN 0122499514, 2005, which is hereby incorporated by reference in its entirety. At least one reference suggests that a pH of less than 5 will provide efficacy against rhinovirus, and other acid labile viruses.

10 [06] However, many non-enveloped viruses are known to be stable at an acid pH. These include Hepatitis A, Poliovirus, Coxsackievirus, Echovirus, Enterovirus, Adenovirus, Rotavirus, Parvovirus, Papillomavirus, and Norovirus. Thus, while acid-containing disinfectants have been reported to have some antiviral efficacy against, for example, rhinovirus, they have insufficient efficacy against other non-enveloped viruses. That is, the
15 efficacy of these acidic disinfectants is narrow and limited.

[07] U.S. Pat. No. 6,080,417 teaches a hand disinfectant that contains from 50 to 60 volume percent lower alcohol, a C₃₋₅ diol, and a synergist selected from hydrogen peroxide, alkane sulfonates, and salts of thiocyanic acid.

[08] U.S. Pat. No. 6,034,133 teaches a hand lotion containing a C₁₋₆ alcohol, malic acid, and citric acid that, when applied frequently, is asserted to prevent hand-to-hand
20 transmission of rhinoviruses. The lotion was applied to finger pads and dried. A viral suspension was applied to the same finger pads and allowed to dry for ten to fifteen minutes. The finger pads were rinsed, and a viral titration determined that the rhinovirus had been eradicated.

25 [09] U.S. Pat. No. 5,043,357 teaches virucidal composition containing at least 70 weight percent ethanol and/or propanol, and from 1-5 weight percent of a short-chain organic acid. The virucidal composition is stated to have broad spectrum antiviral efficacy after periods of treatment of at least 1 to 2 minutes. The skin to be disinfected must first be treated to remove skin fats before the antiviral composition is applied.

30 [10] U.S. Pub. App. No. 2002/0165278 A1 teaches a method for inactivating viruses comprising contacting the virus with a virucidally effective amount of a composition consisting essentially of a dilute aqueous solution of from 0.2 to 13 volume percent C₁₋₃

monohydroxy alcohol or a C₂₋₄ diol, and a sufficient amount of acid to adjust the pH to below 4.6. At these relatively low levels of alcohol, this composition would not be expected to have rapid antibacterial efficacy.

[11] U.S. Pub. App. No. 2005/105070 A1 teaches an aqueous antimicrobial composition stated to have antiviral efficacy against rhinovirus, rotavirus, coronavirus, and respiratory syncytial virus. The composition includes up to 70 % of an organic acid and up to 40 % of a specific short-chain anionic surfactant having at least one of a large hydrophilic head group, a branched alkyl chain, or an unsaturated alkyl chain. The composition was tested for antiviral efficacy for periods of from 1 to 10 minutes. These relatively high levels of acid and anionic surfactant would be expected to be irritating to the skin, and would not be suitable for leave-on type antiviral products.

[12] U.S. Pub. App. No. 2004/101726 A1 teaches a composition comprising from 10 to 30 volume % alcohol, from 10 to 30 volume % of a long-chain alkyl polyamine, and a halogen, such as iodine. The composition is stated to have antiviral efficacy, and was tested against poliovirus for periods of from 5 to 60 minutes. No testing of other non-enveloped viruses was reported. Also, there was no indication of contact periods of less than 5 minutes.

[13] International Pub. App. No. WO 2001/28340 teaches an antimicrobial composition stated to have antiviral efficacy, although no test data was reported. The composition comprises a dicarboxylic acid, a metal salt, and a dermatologically acceptable carrier. Suitable metal salts include those of metals of Group I, II, IIIA, IV, VIB, VIII, rare earth compounds, and combinations thereof.

[14] None of the aforementioned publications teaches methods that have broad, fast efficacy against non-enveloped viruses. Each is either limited in its spectrum of antiviral activity or requires long contact times. Therefore, it would be desirable to have a method that achieves a high level of inactivation of non-enveloped virus particles in a short amount of time. A need continues to exist for a method for rapidly inactivating most, if not all, viruses. Furthermore, a need exists for alcoholic compositions that have bactericidal and virucidal efficacy and may be used topically against a broad spectrum of enveloped and non-enveloped viruses. In addition, there is a need for an antiviral composition that does not require toxic, regulated, or sensitizing components.

[15] The European Committee for Standardization developed an antiviral test method denominated EN 14476:2005 and entitled "Virucidal Quantitative Suspension Test for

Chemical Disinfectants and Antiseptics Used in Human Medicine.” This Standard sets forth protocols by which hygienic handrubs and handwashes are to be tested for efficacy against poliovirus and adenovirus. There is a need for antiviral compositions that provide efficacy against poliovirus and adenovirus when tested according to the EN 14476:2005.

5

SUMMARY OF THE INVENTION

[16] This invention provides a method of inactivating non-enveloped virus particles, the method comprising contacting non-enveloped virus particles with a virucidally-enhanced alcoholic composition comprising a C₁₋₆ alcohol, and an efficacy-enhancing amount of one or
10 more enhancers selected from the group consisting of cationic oligomers and polymers, proton donors, chaotropic agents, and mixtures thereof, with the proviso that when the alcoholic composition comprises a proton donor, the composition further comprises a synergistic amount of a cationic oligomer or polymer.

[17] The invention further provides a method of producing a topical virucidal effect on
15 mammalian skin against non-enveloped virus by applying a virucidally-enhanced alcoholic composition comprising a C₁₋₆ alcohol, and an efficacy-enhancing amount of one or more enhancers selected from the group consisting of cationic oligomers and polymers, proton donors, chaotropic agents, and mixtures thereof, with the proviso that when the alcoholic composition comprises a proton donor, the composition further comprises a synergistic
20 amount of a cationic oligomer or polymer.

[18] The invention still further provides a method of enhancing the efficacy of a C₁₋₆ alcohol against non-enveloped virus in a topical application to a surface, the method comprising combining said C₁₋₆ alcohol with an efficacy-enhancing amount of an enhancer selected from the group consisting of cationic oligomers and polymers, proton donors,
25 chaotropic agents, and mixtures thereof, to form an antiviral composition, with the proviso that where the antiviral composition comprises a proton donor, the composition further comprises a synergistic amount of a cationic oligomer or polymer.

[19] The invention further provides a virucidally-enhanced alcoholic composition comprising a C₁₋₆ alcohol; and an efficacy-enhancing amount of an enhancer selected from the
30 group consisting of cationic oligomers and polymers, proton donors, chaotropic agents, and mixtures thereof, with the proviso that where the alcoholic composition comprises a proton donor, the composition further comprises a synergistic amount of a cationic oligomer or

polymer, wherein said virucidal composition exhibits an efficacy against non-enveloped viruses that is higher than the efficacy of the same composition but not comprising said enhancer.

5 DETAILED DESCRIPTION OF ILLUSTRATIVE EMBODIMENTS

[20] The present invention provides a method of inactivating non-enveloped virus particles. In one embodiment, the antiviral method has rapid antiviral efficacy against non-enveloped viruses including members of the families *Picornaviridae*, *Reoviridae*, *Caliciviridae*, *Adenoviridae* and *Parvoviridae*. More specifically, in certain embodiments, the antiviral method has rapid antiviral efficacy against non-enveloped viruses such as rhinovirus, poliovirus, adenovirus, norovirus, papillomavirus, feline calicivirus, hepatitis A virus, parvovirus, and rotavirus. In one or more embodiments, the antiviral method has rapid antiviral efficacy against adenovirus, norovirus, papillomavirus, feline calicivirus, hepatitis A virus, parvovirus, and rotavirus. Advantageously, the antiviral method has rapid antiviral efficacy against papillomavirus, feline calicivirus, hepatitis A virus, and parvovirus.

[21] In certain embodiments, the antiviral method of the present invention is also effective in killing gram negative and gram positive bacteria, fungi, parasites, and enveloped viruses. More specifically, in certain embodiments the antiviral method has rapid antibacterial efficacy against gram positive bacteria such as *Staphylococcus*, and against gram negative bacteria such as *Escherichia coli*. In these or other embodiments, the present method has rapid efficacy against fungi such as *Aspergillus*. In one or more embodiments, the present method has efficacy against enveloped viruses such as herpes and influenza.

[22] The antiviral method includes contacting the virus with an antiviral composition. The physical form of the antiviral composition is not particularly limited, and in one or more embodiments, the composition may be presented as a liquid that is poured, pumped, sprayed, or otherwise dispensed, a gel, an aerosol, or a foam, including both aerosol and non-aerosol foams. The antiviral composition may be employed on a wide variety of surfaces or substrates, including skin, porous, and non-porous surfaces. In one or more embodiments, the antiviral composition may be presented as a wipe, *i.e.* a tissue or cloth that is wiped over a surface. In general, the antiviral composition includes an alcohol, and an enhancer selected from cationic oligomers or polymers, proton donors, chaotropic agents, and mixtures thereof.

[23] Advantageously, the method of the present invention has antiviral efficacy over a wide range of temperatures, including ambient temperatures of from about 25 to about 35 °C. In one embodiment, the antiviral composition is brought into contact with the virus particles, and greater than 1 log reduction is achieved in less than 60 seconds, in another embodiment greater than 2 log reduction is achieved, and in yet another embodiment, greater than 3 log reduction is achieved in less than 60 seconds. In another embodiment, greater than 3.5 log reduction is achieved in less than 60 seconds, and in yet another embodiment, greater than 4 log reduction is achieved in less than 60 seconds. In one or more embodiments, the virus is completely inactivated to the limits of detection of the test method within about 60 seconds. In certain embodiments, the antiviral composition is brought into contact with the virus particles, and greater than 1 log reduction is achieved in less than 30 seconds, in another embodiment greater than 2 log reduction is achieved, and in yet another embodiment, greater than 3 log reduction is achieved in less than 30 seconds, in another embodiment, greater than 3.5 log reduction is achieved in less than 30 seconds, and in yet another embodiment, greater than 4 log reduction is achieved in less than 30 seconds. In one or more embodiments, the virus is completely inactivated to the limits of detection of the test method within about 30 seconds.

[24] The antiviral composition exhibits efficacy against MS2, a non-enveloped bacteriophage that is sometimes employed in tests to indicate efficacy against non-enveloped viruses. In one embodiment, the antiviral composition is brought into contact with the non-enveloped bacteriophage MS2, and greater than 1 log reduction is achieved in less than 60 seconds, in another embodiment greater than 2 log reduction is achieved, and in yet another embodiment, greater than 3 log reduction is achieved in less than 60 seconds. In another embodiment, greater than 3.5 log reduction of MS2 virus is achieved in less than 60 seconds. In yet another embodiment, greater than 4 log reduction of MS2 is achieved in less than 60 seconds. In one or more embodiments, the virus is completely inactivated to the limits of detection of the test method within about 60 seconds. In certain embodiments, the antiviral composition is brought into contact with the virus particles, and greater than 1 log reduction is achieved in less than 30 seconds, in another embodiment greater than 2 log reduction is achieved, and in yet another embodiment, greater than 3 log reduction of MS2 is achieved in less than 30 seconds. In another embodiment, greater than 3.5 log reduction of MS2 is achieved in less than 30 seconds. In yet another embodiment, greater than 4 log reduction of

MS2 is achieved in less than 30 seconds. In one or more embodiments, the virus is completely inactivated to the limits of detection of the test method within about 30 seconds.

[25] In another embodiment, the antiviral composition is brought into contact with a mammalian virus, such as adenovirus, and greater than 1 log reduction is achieved in less than 5 60 seconds, in another embodiment greater than 2 log reduction is achieved, and in yet another embodiment, greater than 3 log reduction is achieved in less than 60 seconds. In another embodiment, greater than 3.5 log reduction is achieved in less than 60 seconds. In yet another embodiment, greater than 4 log reduction is achieved in less than 60 seconds. In one or more embodiments, the virus is completely inactivated to the limits of detection of the test method within about 60 seconds. In certain embodiments, the antiviral composition is brought into contact with the adenovirus particles, and greater than 1 log reduction is achieved in less than 30 seconds, in another embodiment greater than 2 log reduction is achieved, and in yet another embodiment, greater than 3 log reduction is achieved in less than 30 seconds. In another embodiment, greater than 3.5 log reduction is achieved in less than 30 15 seconds. In yet another embodiment, greater than 4 log reduction is achieved in less than 30 seconds. In one or more embodiments, the virus is completely inactivated to the limits of detection of the test method within about 30 seconds.

[26] In one embodiment, the methods of bringing the antiviral composition into contact with a virus on human skin includes applying an amount of the composition to the skin, and allowing the composition to remain in contact with the skin for a suitable amount of 20 time. In other embodiments, the composition may be spread over the surface of the skin, rubbed in, or rinsed off, allowed to dry via evaporation, or wiped off.

[27] Advantageously, the antiviral composition of the present invention exhibits enhanced efficacy against non-enveloped viruses, when compared to the efficacy of alcohol. Whereas C₁₋₆ alcohols have little efficacy against non-enveloped virus, the efficacy may be enhanced by combining the C₁₋₆ alcohol with an efficacy-enhancing amount of an enhancer, to form an antiviral composition. In one or more embodiments, the antiviral composition exhibits an increased efficacy against non-enveloped viruses when compared to a composition containing an equivalent amount of C₁₋₆ alcohol. In certain embodiments, a synergistic effect 25 is seen. In other words, the efficacy of the antiviral composition against non-enveloped virus is greater than the sum of the efficacies of equivalent amounts of the individual components. 30

[28] Therefore, the present invention provides a virucidally-enhanced alcoholic composition comprising alcohol, and an enhancer. In one embodiment, the alcohol is a lower alkanol, i.e. an alcohol containing 1 to 6 carbon atoms. Typically, these alcohols have antimicrobial properties. Examples of lower alkanols include, but are not limited to, methanol, ethanol, propanol, butanol, pentanol, hexanol, and isomers and mixtures thereof. In one embodiment, the alcohol comprises ethanol, propanol, or butanol, or isomers or mixtures thereof. In another embodiment, the alcohol comprises ethanol.

[29] Generally, the antiviral composition comprises an amount of alcohol of at least about 50 percent by weight. In embodiments where rapid antimicrobial efficacy is not a requirement, the amount of alcohol may be reduced. In one embodiment, the antiviral composition comprises at least about 60 weight percent alcohol, in another embodiment, the antiviral composition comprises at least about 65 weight percent alcohol, in yet another embodiment, the antiviral composition comprises at least about 70 weight percent alcohol, and in still yet another embodiment, the antiviral composition comprises at least about 78 weight percent alcohol, based upon the total weight of antiviral composition. More or less alcohol may be required in certain instances, depending particularly on other ingredients and/or the amounts thereof employed in the composition. In certain embodiments, the antiviral composition comprises from about 50 weight percent to about 98 weight percent alcohol, in other embodiments, the antiviral composition comprises from about 60 weight percent to about 95 weight percent of alcohol, in yet other embodiments, the antiviral composition comprises from about 65 weight percent to about 90 weight percent of alcohol, and in still other embodiments, the antiviral composition comprises from about 70 weight percent to about 85 weight percent of alcohol, based upon the total weight of the antiviral composition.

[30] It has been found that, in certain embodiments, a cationic oligomer or polymer enhances the antiviral efficacy of alcoholic compositions against non-enveloped viruses. Cationic oligomers or polymers include, but are not necessarily limited to, cationic polysaccharides, cationic copolymers of saccharides and synthetic cationic monomers, and synthetic cationic oligomer or polymers. Synthetic cationic oligomers or polymers include cationic polyalkylene imines, cationic ethoxy polyalkylene imines, cationic poly[N-[3-(dialkylammonio)alkyl] N'[3-(alkyleneoxyalkylene dialkylammonio)alkyl]urea dichloride], vinyl caprolactam/VP/dialkylaminoalkyl alkylate copolymers, and polyquaternium polymers.

[31] Examples of cationic oligomers or polymers include chitosan, copolymers of isophorone diisocyanate and PEG-15 cocamine, vinyl caprolactam/VP/dimethylaminoethyl methacrylate copolymer, polyquaternium-4/hydroxypropyl starch copolymer, butylmethacrylate-(2-dimethylaminoethyl)methacrylate-methylmethacrylate-copolymer, guar
 5 hydroxypropyl trimonium chloride and dilinoleyl amidopropyl dimethylammonium chloride hydroxypropyl copolymer. Examples of polyquaterniums include those listed in Table 1, below, including the INCI name and technical name.

Table 1

INCI Name	Technical Name
Polyquaternium	
-X	
-2	Bis(2-chloroethyl)ether, polym. w. N,N'-bis[3-(dimethylamino)propyl]urea
-4	Hydroxyethylcellulose Dimethyldiallylammoinum Chloride Copolymer
-5	Copolymer of acrylamide and beta-methacrylyloxyethyl trimethyl ammonium methosulfate
-6	Polydimethyldiallyl Ammonium Chloride
-7	Dimethyldiallyl Ammonium Chloride & Acrylamide Copolymer
-9	Polydimethyaminoethyl methacrylate quaternized with Methyl Bromide
-10	Hydroxyethylcellulose reacted with trimethyl ammonium substituted epoxide
-11	PVP N,N-Dimethyl Aminoethyl Methacrylic Acid Copolymer Diethyl Sulfate Soln
-14	Ethanaminium, N,N,N-Trimethyl-2-[(2-methyl-1-oxo-2-propenyl)oxy]-, Methyl Sulfate Homopolymer
-15	Acrylamide-Dimethylaminoethyl Methacrylate Methyl Chloride Copolymer
-16	3-Methyl-1-Vinylimidazolium Chloride-1-Vinyl-2-Pyrrolidinone Chloride
-17	Quat salt made from Adipic acid & diethylaminopropylamine & dichloroether
-18	Quat salt prepared by the reaction of adipic acid and dimethylaminopropylamine, reacted with dichloroethyl ether
-19	Quat ammonium salt prepared by the reaction of polyvinyl alcohol with 2,3-epoxypropylamine
-20	Quat ammonium salt prepared by the reaction of polyvinyl octadecyl ether with 2,3-epoxypropylamine
-22	Acrylic Acid-Diallyldimethylammonium Chloride (DADMAC) Polymer
-24	Polyquat ammonium salt of hydroxyethyl cellulose reacted with lauryl dimethyl ammonium substituted epoxide
-27	Block Copolymer of Polyquaternium-2 and 17

INCI Name	Technical Name
Polyquaternium	
-X	
-28	Vinylpyrrolidone/Methacrylamidopropyltrimethylammonium Chloride Copolymer
-29	Propoxylated Chitosan quaternized with epichlorhydrin
-30	Ethanaminium, N-Carboxymethyl)-N,N-Dimethyl-2-((2-Methyl-1-Oxo-2-Propenyl)Oxy)-, Inner Salt, Polymer with Methyl 2-Methyl-2-Propenoate
-31	2-propane nitrile reaction product w/ N,N-dimethylpropanediamine, Sulfate
-32	Acrylamide-Dimethylaminoethyl Methacrylate Methyl Chloride (DMAEMA) Copolymer
-37	Trimethylaminoethyl Methacrylate Chloride Polymer
-39	Acrylic Acid (AA), Polymer w/ Acrylamide & Diallyldimethylammonium Chloride(DADMAC)
-42	Polyoxyethylene (dimethyliminio)ethylene-(dimethyliminio)ethylene dichloride
-43	Copolymer of Acrylamide, acrylamidopropyltrimonium chloride, amidopropylacrylamide & DMAPA Monomers
-44	Polyquat ammonium salt of vinylpyrrolidone & quaternized imidazoline monomers
-46	Quat ammonium salt of vinylcaprolactum, vinylpyrrolidone & methylvinylimidazolium
-47	Quat ammonium chloride- acrylic acid, methyl acrylate & methacrylamidopropyltrimonium Chloride
-48	Copolymer of methacryloyl ethyl betaine, 2-hydroxyethylmethacrylate & methacryloylethyltrimethylammonium chloride
-51	3,5,8-Triox-4-Phosphaundec-10-en-1-aminium, 4-Hydroxy-N,N,N,10-Tetramethyl-9-Oxo, Inner Salt, 4-Oxide, Polymer with Butyl 2-Methyl-2-Propenoate
-53	Acrylic Acid (AA)/Acrylamide/Methacrylamidopropyltrimonium Chloride (MAPTAC) Copolymer
-54	Polymeric quaternary ammonium salt prepared by the reaction of aspartic acid and C6-18 alkylamine with dimethylaminopropylamine and sodium chloroacetate
-55	1-Dodecanaminium, N,N-Dimethyl-N-[3-[(2-Methyl-1-Oxo-2-Propenyl)AminoPropyl]-, Chloride, Polymer with N-[3-(Dimethylamino)Propyl]-2-Methyl-2-Propenamide and 1-Ethenyl-2-Pyrrolidinone
-56	Polymeric quaternary ammonium salt prepared by the reaction of aspartic acid and C6-18 alkylamine with dimethylaminopropylamine and sodium chloroacetate.
-57	Polymeric quaternary ammonium salt consisting of Castor Isostearate Succinate (q.v.) and Ricinoleamidopropyltrimonium Chloride (q.v.) monomers
-58	2-Propenoic Acid, Methyl Ester, Polymer with 2,2-Bis[(2-

INCI Name	Technical Name
Polyquaternium -X	Propenyloxy)Methyl]-1-Butanol and Diethenylbenzene, Reaction Products with N,N-Dimethyl-1,3-Propanediamine, Chloromethane-Quaternized
-59	Polyquaternium polyester
-60	9-Octadecenoic Acid, 12-Hydroxy-, [(2-Hydroxyethyl)Imino]Di-2,1-Ethanediy Ester, Polymer with 5-Isocyanato-1-(Isocyanatomethyl)-1,3,3-Trimethylcyclohexane, Compd. with Diethyl Sulfate
-62	Polymeric quaternary ammonium salt prepared by the reaction of butyl methacrylate, polyethylene glycol methyl ether methacrylate, ethylene glycol dimethacrylate and 2-methacryloyethyl trimonium chloride with 2,2'-azobis(2-methyl propionamidine) dihydrochloride
-63	Copolymer of acrylamide, acrylic acid and ethyltrimonium chloride acrylate
-65	Polymeric quaternary ammonium salt consisting of 2-methacryloyloxyethylphosphorylcholine, butyl methacrylate and sodium methacrylate monomers
-68	Quaternized copolymers of vinylpyrrolidone (VP), methacrylamide(MAM) vinylimidazole(VI) & quaternized vinylimidazole (QVI)
-69	Polymeric quaternary ammonium salt containing vinyl caprolactam, vinylpyrrolidone, dimethylaminopropyl methacrylamide (DMAPA), and methoacryloylaminopropyl lauryldimonium chloride
-70	
-71	
-72	
-73	
-74	
-75	

[32] In one or more embodiments, the polyquaternium polymer includes polyquaternium-2, polyquaternium-4, polyquaternium-5, polyquaternium-6, polyquaternium-7, polyquaternium-10, polyquaternium-11, polyquaternium-16, polyquaternium-22, polyquaternium-24, polyquaternium-28, polyquaternium-32, polyquaternium-37, polyquaternium-39, polyquaternium-42, polyquaternium-43, polyquaternium-44, polyquaternium-46, polyquaternium-47, polyquaternium-51, polyquaternium-53, polyquaternium-55, polyquaternium-57, polyquaternium-58, polyquaternium-59, polyquaternium-60, polyquaternium-63, polyquaternium-64, polyquaternium-65, polyquaternium-68, or mixtures thereof.

[33] In one embodiment, the polyquaternium polymer includes polyquaternium-2, polyquaternium-4, polyquaternium-6, polyquaternium-7, polyquaternium-11, polyquaternium-16, polyquaternium-22, polyquaternium-28, polyquaternium-32, polyquaternium-37, polyquaternium-39, polyquaternium-42, polyquaternium-47, 5 polyquaternium-51, polyquaternium-53, polyquaternium-55, polyquaternium-58, or mixtures thereof. In another embodiment, the polyquaternium polymer includes polyquaternium-37.

[34] In certain embodiments, the cationic oligomer or polymer is characterized by a charge density that may be determined by methods known in the art, such as colloidal titration. In one embodiment, the charge density of the cationic oligomer or polymer is at 10 least about 0.1 meq/g, in another embodiment at least about 2.5 meq/g, and in yet another embodiment, at least about 5 meq/g.

[35] Advantageously, it has been found that antiviral compositions comprising alcohol and an efficacy-enhancing amount of cationic oligomer or polymer have increased efficacy against a broad spectrum of non-enveloped viruses, when compared to antiviral compositions 15 comprising alcohol without cationic oligomer or polymer. In certain embodiments, cationic oligomers or polymers that exhibit no efficacy on their own against non-enveloped viruses, provide an enhanced efficacy when combined with alcohol according to the present invention.

[36] In one embodiment, an efficacy-enhancing amount of cationic oligomer or polymer is at least about 0.02 percent by weight, based upon the total weight of the antiviral 20 composition, in another embodiment at least about 0.05, and in yet another embodiment at least about 0.1 percent by weight, based upon the total weight of the antiviral composition. Generally, an efficacy-enhancing amount of cationic oligomer or polymer is from about 0.02 to about 20 percent by weight, based upon the total weight of the antiviral composition. In one embodiment, the cationic oligomer or polymer is present in an amount of from about 0.1 25 to about 10 weight percent, in another embodiment, the cationic oligomer or polymer is present in an amount of from about 0.25 to about 5 percent by weight, and in yet another embodiment, from about 0.4 to about 1 percent by weight, based upon the total weight of the antiviral composition. In certain embodiments, the amount of cationic oligomer or polymer may affect the viscosity of the antiviral composition, as well as other aesthetic qualities. 30 Nevertheless, it will be understood that greater amounts of cationic oligomer or polymer can be employed, if desired, and are expected to perform at least equally as well, in terms of antiviral efficacy.

[37] The cationic oligomer or polymer may be supplied in the form of a dry powder, or as an emulsion or liquid mixture. In one embodiment, the cationic oligomer or polymer is added to the antiviral composition as a solid. In another embodiment, the cationic oligomer or polymer is added to the antiviral composition as a solution or emulsion. In other words, the cationic oligomer or polymer may be premixed with a carrier, and optionally one or more other ingredients, to form a cationic oligomer or polymer solution or emulsion, with the proviso that the carrier does not deleteriously affect the antiviral properties of the composition. More specifically, a carrier deleteriously affects the antiviral properties of the composition when it decreases the log reduction by more than a *de minimus* amount. By *de minimus* is meant a decrease of less than about 0.5 log reduction.

[38] Examples of carriers include water, alcohol, or blends of water and another carrier such as glycols, ketones, linear and/or cyclic hydrocarbons, triglycerides, carbonates, silicones, alkenes, esters such as acetates, benzoates, fatty esters, glyceryl esters, ethers, amides, polyethylene glycols, PEG/PPG copolymers, inorganic salt solutions such as saline, and mixtures thereof. It will be understood that, when the cationic oligomer or polymer is premixed to form a cationic oligomer or polymer solution or emulsion, the amount of solution or emulsion that is added to the antiviral composition is selected so that the amount of cationic oligomer or polymer falls within the ranges set forth hereinabove.

[39] In certain embodiments, the antiviral composition further includes a proton donor. Proton donors include Arrhenius acids, Bronsted-Lowry acids and Lewis acids. Strong or weak acids may be used.

[40] Examples of acids include mineral acids and organic acids. Mineral acids include, without limitation, hydrochloric acid, nitric acid, phosphoric acid, phosphonic acid, boric acid, and sulfuric acid. Organic acids include sulfonic acids, organophosphorus acids, carboxylic acids such as benzoic acids, propionic acids, phthalic acids, butyric acids, acetic acids, amino acids, and other substituted and unsubstituted organic acids.

[41] Examples of organic acids include adipic acid, benzene 1,3,5 tricarboxylic acid, chlorosuccinic acid, choline chloride, *cis*-aconitic acid, citramalic acid, citric acid, cyclobutane 1,1,3,3 tetracarboxylic acid, cyclohexane 1,2,4,5 tetracarboxylic acid, cyclopentane 1,2,3,4 tetracarboxylic acid, diglycolic acid, fumaric acid, glutamic acid, glutaric acid, glyoxylic acid, isocitric acid, ketomalononic acid, lactic acid, maleic acid, malic acid, malonic acid, nitrilotriacetic acid, oxalacetic acid, oxalic acid, phytic acid, p-

toluenesulfonic acid, salicylic acid, succinic acid, tartaric acid, tartronic acid, tetrahydrofuran 2,3,4,5 tetracarboxylic acid, tricarballic acid, versene acids, 3-hydroxyglutaric acid, 2-hydroxypropane 1,3 dicarboxylic acid, glyceric acid, furan 2,5 dicarboxylic acid, 3,4-dihydroxyfuran-2,5 dicarboxylic acid, 3,4-dihydroxytetrahydrofuran-2,5-dicarboxylic acid, 2-oxo-glutaric acid, *dl*-glyceric acid, and 2,5 furandicarboxylic acid.

[42] In certain embodiments, the proton donor includes a hydroxy carboxylic acid, and in one embodiment, the hydroxy acid includes two or more carboxylic acid groups. In one or more embodiments, the hydroxy carboxylic acid includes alpha-hydroxy acids and beta-hydroxy acids. Examples of alpha-hydroxy acids having two or more carboxylic acid groups include tartaric acid, malic acid, citric acid, and isocitric acid. Examples of other alpha-hydroxy carboxylic acids include lactic acid, tartronic acid, and malonic acid. In one embodiment, the proton donor includes citric acid, lactic acid, malic acid, tartaric acid, salicylic acid, oxalic acid, or mixtures thereof. In one embodiment, the proton donor includes citric acid.

[43] It has been found that, in certain embodiments, a proton donor enhances the antiviral efficacy of alcoholic solutions against non-enveloped viruses. In one or more embodiments, proton donors that exhibit moderate or no efficacy on their own against non-enveloped viruses, provide an enhanced efficacy when present in the antiviral composition of the present invention.

[44] In one or more embodiments, a synergistic enhancement of antiviral efficacy may be achieved by contacting non-enveloped virus particles with a virucidally-enhanced alcoholic composition comprising a C₁₋₆ alcohol, an efficacy-enhancing amount of a proton donor, and a synergistic amount of a cationic oligomer or polymer. The minimum amount of cationic oligomer or polymer that corresponds to a synergistic amount is at least about 0.02 percent by weight, based upon the total weight of the antiviral composition, in another embodiment at least about 0.05, and in yet another embodiment at least about 0.1 percent by weight, based upon the total weight of the antiviral composition.

[45] The amount of proton donor is not particularly limited, so long as it is at least an efficacy-enhancing amount. The minimum amount of proton donor that corresponds to an efficacy-enhancing amount can be determined by comparing the log reduction of virus achieved by a composition comprising an alcohol to a composition comprising an alcohol and a given amount of proton donor. The amount of proton donor below which no difference in

log reduction is seen is an efficacy-enhancing amount. In certain embodiments, for example when efficacy against MS2 virus is desired, the minimum efficacy-enhancing amount of proton donor is about 0.01 percent by weight, based upon the total weight of the antiviral composition. In another embodiment, for example when efficacy against feline calicivirus is desired, the minimum efficacy-enhancing amount of proton donor is about 0.04 percent by weight, based upon the total weight of the antiviral composition.

[46] In one embodiment, the proton donor is added in an amount of from about 0.01 to about 1 weight percent, based upon the total weight of the antiviral composition. In another embodiment, the amount of proton donor is from about 0.015 to about 0.5 weight percent, and in yet another embodiment, from about 0.03 to about 0.3 weight percent, based upon the total weight of the antiviral composition. It will be understood that greater levels of proton donor can be used, if desired, and are expected to perform at least equally as well.

[47] In one embodiment, the proton donor is added to the antiviral composition as a solution or emulsion. In other words, the proton donor may be premixed with a carrier, and optionally one or more other ingredients, to form a proton donor solution or emulsion, with the proviso that the carrier does not deleteriously affect the antiviral properties of the composition. Examples of carriers include water, alcohol, any of the blends described above as carriers for the cationic oligomer or polymer, and mixtures thereof. It will be understood that, when the proton donor is premixed to form a proton donor solution or emulsion, the amount of solution or emulsion that is added to the antiviral composition is selected so that the amount of proton donor falls within the ranges set forth hereinabove.

[48] In one or more embodiments, the virucidally-enhanced alcoholic composition comprises alcohol, a cationic oligomer or polymer, and a synergistic amount of a zinc or copper compound. Synergistic zinc or copper compounds include those where the zinc or copper is present in the compound as an ion (*e.g.* has an oxidation state of I or II). In one or more embodiments, the copper or zinc compound is soluble in water and/or hydroalcoholic compositions.

[49] Examples of efficacy-enhancing zinc compounds include aluminum zinc oxide, ammonium silver zinc aluminum silicate, ethylene/zinc acrylate copolymer, lactobacillus/milk/calcium/phosphorus/magnesium/zinc ferment, lactobacillus/milk/manganese/zinc ferment lysate, luminescent zinc sulfide, magnesium/aluminum/zinc/hydroxide/carbonate, porphyridium/zinc ferment,

saccharomyces/zinc ferment, saccharomyces/zinc/iron /germanium/copper /magnesium /silicon ferment, saccharomyces/zinc/magnesium/calcium/germanium/selenium ferment, silicon /titanium/cerium/zinc oxides, sodium zinc cetyl phosphate, sodium zinc histidine dithiooctanamide, zinc acetate, zinc acetylmethionate, zinc adenosine triphosphate, zinc
 5 ascorbate, zinc aspartate, zinc borate, zinc borosilicate, zinc carbonate, zinc carbonate hydroxide, zinc cerium oxide, zinc chloride, zinc citrate, zinc coceth sulfate, zinc coco-sulfate, zinc cysteinate, zinc dibutyldithiocarbamate, zinc DNA, zinc formaldehyde sulfoxylate, zinc glucoheptonate, zinc gluconate, zinc glutamate, zinc glycinate, zinc glycyrrhetinate, zinc hexametaphosphate, zinc hydrolyzed collagen, zinc lactate, zinc laurate,
 10 zinc magnesium aspartate, zinc myristate, zinc neodecanoate, zinc oxide, zinc palmitate, zinc PCA, zinc pentadecene tricarboxylate, zinc peroxide, zinc phenolsulfonate, zinc picolinate, zinc pyrithione, zinc ricinoleate, zinc rosinatate, zinc salicylate, zinc silicates, zinc stearate, zinc sulfate, zinc sulfide, zinc thiosalicylate, zinc undecylenate, zinc undecylenoyl hydrolyzed wheat protein, and zinc zeolite.

15 **[50]** Examples of efficacy-enhancing copper compounds include copper sulfate, copper citrate, copper oxylate, copper usnate, copper acetate, copper chloride, copper carbonate, alanine/histidine/lysine polypeptide copper HCl, bis(tripeptide-1) copper acetate, chlorophyllin-copper complex, copper acetylmethionate, copper acetyl tyrosinate methylsilano, copper adenosine triphosphate, copper aspartate, copper chlorophyll, copper
 20 DNA, copper gluconate, copper PCA, copper PCA methylsilanol, copper picolinate, copper powder, copper sulfate, copper tripeptide-1, disodium EDTA-copper, saccharomyces/copper ferment, saccharomyces/copper ferment lysate filtrate, saccharomyces/zinc/iron/germanium/copper/magnesium/silicon ferment, and silver copper zeolite.

25 **[51]** It has been found that, in certain embodiments, a copper or zinc compound enhances the antiviral efficacy of alcoholic solutions against non-enveloped viruses. In one or more embodiments, copper or zinc compounds that exhibit moderate or no efficacy on their own against non-enveloped viruses, provide an enhanced efficacy when present in the antiviral composition of the present invention.

30 **[52]** In one or more embodiments, a synergistic enhancement of antiviral efficacy may be achieved by contacting non-enveloped virus particles with a virucidally-enhanced

alcoholic composition comprising a C₁₋₆ alcohol, an efficacy-enhancing amount of a cationic oligomer or polymer, and a synergistic amount of a copper or zinc compound.

[53] The amount of copper or zinc compound is not particularly limited, so long as it is at least a synergistic amount. The minimum amount of copper or zinc compound that corresponds to a synergistic amount can be determined by comparing the log reduction of virus achieved by a composition comprising an alcohol and a cationic oligomer or polymer to a composition comprising an alcohol and a given amount of copper or zinc compound. The amount of copper or zinc compound below which no difference in log reduction is seen is a synergistic amount.

[54] In certain embodiments, the minimum synergistic amount of copper or zinc compound is that which will provide an effective amount of copper or zinc ion to the antiviral composition. In one or more embodiments, an effective amount of copper or zinc ion is at least about 1 part per million (ppm) by weight, based upon the total weight of the antiviral composition, in other embodiments, at least about 10 ppm, and in yet other embodiments, at least about 30 ppm by weight, based upon the total weight of the antiviral composition. One of ordinary skill in the art will be able to determine the molecular weight of a particular copper or zinc compound and calculate a synergistic amount (*i.e.* the amount necessary to deliver the desired parts per million of copper or zinc ion to the antiviral composition).

[55] In one or more embodiments, the minimum synergistic amount of copper or zinc compound is about 0.01 percent by weight, based upon the total weight of the antiviral composition. In certain embodiments, a synergistic amount of copper or zinc compound is at least about 0.03 percent by weight, and in other embodiments, at least about 0.05 percent by weight, based upon the total weight of the antiviral composition. The synergistic amount may vary depending upon which copper or zinc compound is selected and upon which virus is to be inactivated.

[56] In one embodiment, the copper or zinc compound is added in an amount of from about 0.01 to about 1 weight percent, based upon the total weight of the antiviral composition. In another embodiment, the amount of copper or zinc compound is from about 0.03 to about 0.5 weight percent, and in yet another embodiment, from about 0.05 to about 0.1 weight percent, based upon the total weight of the antiviral composition. It will be understood that greater levels of copper or zinc compound can be used, if desired, and are expected to perform at least equally as well.

[57] The copper or zinc compound may be added to the antiviral composition in any appropriate form, for example as a solid or liquid. In one or more embodiments, the copper or zinc compound is added as a powder that dissolves or is dispersed in the antiviral composition. In other embodiments, the copper or zinc compound is added to the antiviral composition as a solution or emulsion. In other words, the copper or zinc compound may be premixed with a carrier, and optionally one or more other ingredients, to form a copper or zinc compound solution or emulsion, with the proviso that the carrier does not deleteriously affect the antiviral properties of the composition. Examples of carriers include water, alcohol, any of the blends described above as carriers for the cationic oligomer or polymer, and mixtures thereof. It will be understood that, when the copper or zinc compound is premixed to form a copper or zinc compound solution or emulsion, the amount of solution or emulsion that is added to the antiviral composition is selected so that the amount of copper or zinc compound falls within the ranges set forth hereinabove.

[58] In one or more embodiments where the antiviral composition includes an efficacy enhancing copper or zinc compound, the amount of acid is limited. In one embodiment, the amount of acid is less than about 0.05 percent by weight, in another embodiment, less than about 0.01 percent by weight, and in yet another embodiment, less than about 0.005 weight percent, based upon the total weight of the antiviral composition. In another embodiment, the antiviral composition is devoid of acid.

[59] In certain embodiments, the antiviral composition includes a chaotropic agent. Chaotropic agents include agents that disrupt molecular structure, particularly molecular structure formed by nonbonding forces such as hydrogen bonding, Van der Waals interaction, and hydrophobic effect. Chaotropic agents are well known in the field of biochemistry and include, but are not limited to, urea, thiourea, guanidine-HCl, guanidine thiocyanate, aminoguanidine bicarbonate, guanidine carbonate, guanidine phosphate, and aminoguanidine-HCl. Although it is known in the art that heat may act as a chaotropic agent, for purposes of this specification, the term chaotropic agent refers to a substance other than heat. This should not be interpreted to exclude the presence of heat from the method of the present invention, because as stated hereinbelow, the method of the present invention operates over a wide range of temperatures.

[60] In one embodiment, the chaotropic agent comprises urea. The chaotropic agent may be supplied in the form of a dry powder, or as an emulsion or liquid mixture, and can

optionally include a carrier such as those described above for the cationic oligomer or polymer.

[61] It has been found that, in certain embodiments, the presence of a chaotropic agent enhances the antiviral efficacy of alcoholic solutions against non-enveloped viruses. Advantageously, a synergistic antiviral effect is observed when the chaotropic agent is combined with alcohol and a cationic oligomer or polymer. Without wishing to be bound by theory, it is believed that the chaotropic agent may enhance the antiviral efficacy of the alcoholic composition by disrupting the proteins of the virus capsid. In certain embodiments, chaotropic agents that exhibit no efficacy on their own against non-enveloped viruses, provide an enhanced efficacy when combined with alcohol according to the present invention. In contrast to views expressed in the prior art, where concentrations of about 6-8 M are advocated for chaotropic agents in order to denature proteins, it has surprisingly been found that the antiviral method of the present invention provides good antiviral efficacy at much lower concentrations of chaotrope.

[62] The amount of chaotropic agent is not particularly limited, so long as it is at least an efficacy-enhancing amount. The minimum amount of chaotropic agent that corresponds to an efficacy-enhancing amount can be determined by comparing the log reduction of virus achieved by a composition comprising an alcohol to a composition comprising an alcohol and a given amount of chaotropic agent. The amount of chaotropic agent below which no difference in log reduction is seen is an efficacy-enhancing amount.

[63] In one embodiment, the chaotropic agent is added in an amount of from about 0.25 to about 20 weight percent, based upon the total weight of the antiviral composition. In another embodiment, the amount of chaotropic agent is from about 1 to about 15 weight percent, and in yet another embodiment, from about 4 to about 12 weight percent, based upon the total weight of the antiviral composition. It will be understood that greater levels of chaotropic agent can be used, if desired, and are expected to perform equally as well.

[64] As described hereinabove, the antiviral composition of this invention includes an alcohol, and an enhancer selected from cationic oligomers or polymers, proton donors and chaotropic agents. The composition can further comprise a wide range of optional ingredients, with the proviso that they do not deleteriously affect the antiviral efficacy of the composition. By deleterious is meant that the decrease in the log reduction is not de minimus, or in other words, the log reduction does not decrease by more than about 0.5. The CTFA

International Cosmetic Ingredient Dictionary and Handbook, Eleventh Edition 2005, and the 2004 CTFA International Buyer's Guide, both of which are incorporated by reference herein in their entirety, describe a wide variety of non-limiting cosmetic and pharmaceutical ingredients commonly used in the skin care industry, that are suitable for use in the compositions of the present invention. Nonlimiting examples of functional classes of ingredients are described at page 537 of the Handbook. Examples of these functional classes include: abrasives, anti-acne agents, anticaking agents, antioxidants, binders, biological additives, bulking agents, chelating agents, chemical additives; colorants, cosmetic astringents, cosmetic biocides, denaturants, drug astringents, emulsifiers, external analgesics, film formers, fragrance components, humectants, opacifying agents, plasticizers, preservatives (sometimes referred to as antimicrobials), propellants, reducing agents, skin bleaching agents, skin-conditioning agents (emollient, miscellaneous, and occlusive), skin protectants, solvents, surfactants, foam boosters, hydrotropes, solubilizing agents, suspending agents (nonsurfactant), sunscreen agents, ultraviolet light absorbers, detackifiers, and viscosity increasing agents (aqueous and nonaqueous). Examples of other functional classes of materials useful herein that are well known to one of ordinary skill in the art include solubilizing agents, sequestrants, and keratolytics, topical active ingredients, and the like. In one embodiment, the antiviral composition further comprises glycerin.

[65] Foaming surfactants may be included, with the proviso that they will not deleteriously affect the antiviral efficacy of the composition. The foaming surfactant contributes foaming properties to the alcoholic composition, and may include anionic, cationic, nonionic, zwitterionic, or amphoteric surfactants and their associated salts. In one embodiment, the foaming surfactant includes a fluorosurfactant, a siloxane polymer surfactant, or a combination thereof. Fluorosurfactants include compounds that contain at least one fluorine atom. Examples of fluorosurfactants include perfluoroalkylethyl phosphates, perfluoroalkylethyl betaines, fluoroaliphatic amine oxides, fluoroaliphatic sodium sulfosuccinates, fluoroaliphatic stearate esters, fluoroaliphatic phosphate esters, fluoroaliphatic quaternaries, fluoroaliphatic polyoxyethylenes, and the like, and mixtures thereof.

[66] Examples of fluorosurfactants include perfluoroalkylethyl phosphates, perfluoroalkylethyl betaines, fluoroaliphatic amine oxides, fluoroaliphatic sodium sulfosuccinates, fluoroaliphatic phosphate esters, and fluoroaliphatic quaternaries. Specific

examples of fluorosurfactants include DEA-C8-18 perfluoroalkylethyl phosphate, TEA-C8-18 perfluoroalkylethyl phosphate, NH₄-C8-18 perfluoroalkylethyl phosphate, and C8-18 perfluoroalkylethyl betaine.

[67] Siloxane polymer surfactants may be generally characterized by containing one or more Si-O-Si linkages in the polymer backbone. The siloxane polymer surfactant may or may not include a fluorine atom. Therefore, some foaming surfactants may be classified as both fluorosurfactants and siloxane polymer surfactants. Siloxane polymer surfactants include organopolysiloxane dimethicone polyols, silicone carbinol fluids, silicone polyethers, alkylmethyl siloxanes, amodimethicones, trisiloxane ethoxylates, dimethiconols, quaternized silicone surfactants, polysilicones, silicone crosspolymers, and silicone waxes.

[68] Examples of siloxane polymer surfactants include dimethicone PEG-7 undecylenate, PEG-10 dimethicone, PEG-8 dimethicone, PEG-12 dimethicone, perfluorononylethyl carboxydecal PEG 10, PEG-20/PPG-23 dimethicone, PEG-11 methyl ether dimethicone, bis-PEG/PPG-20/20 dimethicone, silicone quats, PEG-9 dimethicone, PPG-12 dimethicone, fluoro PEG-8 dimethicone, PEG 23/PPG 6 dimethicone, PEG 20/PPG 23 dimethicone, PEG 17 dimethicone, PEG5/PPG3 methicone, bis PEG20 dimethicone, PEG/PPG20/15 dimethicone copolyol and sulfosuccinate blends, PEG-8 dimethicone\dimmer acid blends, PEG-8 dimethicone\fatty acid blends, PEG-8 dimethicone\cold pressed vegetable oil\polyquaternium blends, random block polymers and mixtures thereof.

[69] The amount of foaming surfactant is not particularly limited, so long as an effective amount to produce foaming is present. In certain embodiments, the effective amount to produce foaming may vary, depending upon the amount of alcohol and other ingredients that are present. In one or more embodiments, the alcoholic composition includes at least about 0.002 wt. % of foaming surfactant, based upon the total weight of the alcoholic composition. In another embodiment, the alcoholic composition includes at least about 0.01 wt. % of foaming surfactant, based upon the total weight of the alcoholic composition. In yet another embodiment, the alcoholic composition includes at least about 0.05 wt. % of foaming surfactant, based upon the total weight of the alcoholic composition.

[70] Foamable alcoholic compositions are described in co-pending U.S. Patent Application Serial No. 11/438,664, which is hereby incorporated by reference in its entirety.

[71] In certain embodiments, alcohol is the only active antimicrobial or preservative ingredient introduced into the composition. Any antimicrobial or preservative ingredient

other than alcohol may be referred to as an auxiliary antimicrobial agent. In one embodiment, the amount of auxiliary antimicrobial agent is less than about 0.1 percent by weight, in another embodiment, less than about 0.05 percent by weight, based upon the total weight of the antiviral composition. In another embodiment, the antiviral composition is devoid of auxiliary antimicrobial agents.

[72] It is envisioned that, in other embodiments, auxiliary antimicrobial agents could be included, with the proviso that the antimicrobial ingredient does not deleteriously affect the antiviral properties of the composition. Examples of auxiliary antimicrobial agents include, but are not limited to, triclosan, also known as 5-chloro-2(2,4-dichlorophenoxy) phenol (PCMX) and available from Ciba-Geigy Corporation under the tradename IRGASAN®; chloroxylonol, also known as 4-chloro-3,5-xylonol, available from Nipa Laboratories, Inc. under the tradenames NIPACIDE® MX or PX; hexetidine, also known as 5-amino-1,3-bis(2-ethylhexyl)-5-methyl-hexahydropyrimidine; chlorhexidine salts including chlorhexidine gluconate and the salts of N,N"-Bis(4-chlorophenyl)-3,12-diimino-2,4,11,14-tetraazatetradecanediimide amide; 2-bromo-2-nitropropane-1,3-diol, benzalkonium chloride; cetylpyridinium chloride; alkylbenzyltrimethylammonium chlorides; iodine; phenol, bisphenol, diphenyl ether, phenol derivatives, povidone-iodine including polyvinylpyrrolidone-iodine; parabens; hydantoins and derivatives thereof, including 2,4-imidazolidinedione and derivatives of 2,4-imidazolidinedione as well as dimethylol-5,5-dimethylhydantoin (also known as DMDM hydantoin or glydant); phenoxyethanol; cis isomer of 1-(3-chloroallyl)-3,5,6-triaza-1-azoniaadamantane chloride, also known as quaternium-15 and available from Dow Chemical Company under the tradename DOWCIL™ 2000; diazolidinyl urea; benzethonium chloride; methylbenzethonium chloride; glyceryl laurate, transition metal compounds such as silver, copper, magnesium, zinc compounds, hydrogen peroxide, chlorine dioxide, anilides, bisguanidines, and mixtures thereof. When used, the auxiliary antimicrobial agents are present in amounts of from about 0.1 to about 1 percent by weight, based upon the total weight of the antiviral composition.

[73] In certain embodiments, the combination of alcohol and enhancer is the virucidally active ingredient, and the amount of other virucidally active materials is limited. In one embodiment, the amount of auxiliary virucidally active materials is less than about 0.1 percent by weight, in another embodiment less than about 0.05 percent by weight, and in

another embodiment, less than about 0.02 percent by weight, based upon the total weight of the antiviral composition. In another embodiment, the antiviral composition is devoid of auxiliary virucidally active material.

[74] It is envisioned that, in other embodiments, auxiliary antiviral agents could be included, with the proviso that the antiviral ingredient does not deleteriously affect the antiviral properties of the composition according to the present invention. Examples of auxiliary antivirals include botanicals such as rosmarinic acid, tetrahydrocurcuminoids, oleuropen, oleanolic acid, aspalathus linearis extract, white tea, red tea, green tea extract, neem oil limonoids, coleus oil, licorice extract, burnet, ginger & cinnamon extracts, alpha-glucan oligosaccharide, perilla ocymoides leaf powder, camphor, camellia oleifera leaf extract, ginger, menthol, eucalyptus, capillisil hc, hydroxyprolisilane cn, sandalwood oil/resin, calendula oil, rosemary oil, lime/orange oils, and hop acids.

[75] Advantageously, certain ingredients that have been designated in the prior art as critical to achieving rapid antiviral efficacy can be limited in the antiviral composition of the present invention. For example, zinc compounds are not necessary, and can be limited, if desired, to less than about 0.5 percent by weight, or in another embodiment to less than about 0.1 percent by weight, based upon the total weight of the disinfecting composition. In another embodiment, the disinfecting composition is devoid of organic salts of zinc. Zinc compounds that may be so limited include any of those listed hereinabove. Specific zinc compounds that may be so limited include those having a counterion selected from gluconate, acetate, chloride, acetylacetonate, bromide citrate, formate, glycerophosphate, iodide, lactate, nitrate, salicylate, sulfate, pyrrithione, and tartrate.

[76] In certain embodiments, the amount of metal salts in the composition is limited. For example, in one embodiment, the virucidally enhanced composition comprises alcohol, a cationic oligomer or polymer, and a proton donor, and the amount of metal salt is limited. In one embodiment, the amount of metal salts is less than about 0.05 percent by weight, in another embodiment, less than about 0.01 percent by weight, and in yet another embodiment, less than about 0.001 weight percent, based upon the total weight of the antiviral composition. In another embodiment, the antiviral composition is devoid of metal salts.

[77] In certain embodiments, the amount of iodine in the composition is limited. In one embodiment, the amount of iodine is less than about 1 percent by weight, in another embodiment, less than about 0.1 percent by weight, and in yet another embodiment, less than

about 0.01 percent by weight, based upon the total weight of the antiviral composition. In another embodiment, the antiviral composition is devoid of iodine.

[78] In these or other embodiments, the amount of inorganic salts, aluminum compounds, zirconium compounds, or aluminum-zirconium complexes may be limited. In one or more embodiments, the amount of inorganic salts, aluminum compounds, zirconium compounds, or aluminum-zirconium complexes is less than about 0.05 percent by weight, based upon the total weight of the antiviral composition.

[79] In certain embodiments, the amount of fatty acid may be limited. In these embodiments, the amount of fatty acid may be less than about 1 percent by weight, in another embodiment less than about 0.1 percent by weight, in yet another embodiment, less than about 0.05 percent by weight, and in still yet another embodiment, less than about 0.01 percent by weight, based upon the total weight of the antiviral composition. In another embodiment, the antiviral composition is devoid of fatty acid. In these or other embodiments, the amount of fatty ester may be limited. In these embodiments, the amount of fatty ester may be less than about 1 percent by weight, in another embodiment less than about 0.1 percent by weight, in yet another embodiment, less than about 0.05 percent by weight, and in still yet another embodiment, less than about 0.01 percent by weight, based upon the total weight of the antiviral composition. In another embodiment, the antiviral composition is devoid of fatty ester. In these or yet other embodiments, the amount of fatty ether may be limited. In these embodiments, the amount of fatty ether may be less than about 1 percent by weight, in another embodiment less than about 0.1 percent by weight, in yet another embodiment, less than about 0.05 percent by weight, and in still yet another embodiment, less than about 0.01 percent by weight, based upon the total weight of the antiviral composition. In another embodiment, the antiviral composition is devoid of fatty ether.

[80] In general, the fatty acids, fatty esters, and fatty ethers that may optionally be limited include those that are claimed in the literature to have antimicrobial properties. Examples of these antimicrobial fatty compounds include (C6-C14) alkyl carboxylic acids, (C6-C14) alkyl carboxylate ester carboxylic acids, (C8-C22) mono- or polyunsaturated carboxylic acids, (C7-C12)saturated fatty acid esters of polyhydric alcohols, (C8-C22)unsaturated fatty acid esters of polyhydric alcohols, (C7-C22)saturated fatty ethers of polyhydric alcohols, (C8-C22)unsaturated fatty ethers of polyhydric alcohols, and alkoxyated derivatives thereof.

[81] Indeed, any component other than the alcohol and enhancer is not necessary to achieve antimicrobial or antiviral efficacy and can optionally be limited to less than about 0.5 percent by weight, if desired to less than about 0.1 percent by weight, if desired to less than about 0.01 percent by weight, or if desired to less than about 0.001 percent by weight, based upon the total weight of the antiviral composition.

[82] In one or more embodiments, the balance of the alcoholic composition includes water or other suitable solvent. The antiviral composition may be prepared by simply mixing the components together. In one embodiment, where the cationic oligomer or polymer is obtained as a solid powder, the antiviral composition is prepared by a method comprising dispersing the cationic oligomer or polymer in water, adding alcohol with slow to moderate agitation, and then adding other ingredients as desired, and mixing until the mixture is homogeneous.

[83] As stated hereinabove, the antiviral composition of the present invention may be embodied in a variety of forms, including as a liquid, gel, or foam. Surprisingly, it has been found that the viscosity of the liquid antiviral composition does not affect the disinfecting efficacy of the composition. For example, in one or more embodiments of the present invention, the same amount of log reduction is achieved with a liquid antiviral composition having a viscosity of 5 centipoise (cPs) and a disinfecting composition having a viscosity of about 2000 cPs. Thus it will be understood that the viscosity of the antiviral composition of the present invention is not limited.

[84] It will also be understood that the viscosity of the antiviral composition may be affected by the relative amounts of ingredients. For example, a decrease in the relative amount of certain polyquaternium polymers may result in a lower viscosity. Also, the type of polyquaternium polymer can affect the viscosity of the antiviral composition. For example, when a non-thickening cationic oligomer or polymer, such as polyquaternium-22, is employed, the amount of cationic oligomer or polymer may not substantially affect the viscosity of the antiviral composition.

[85] In one embodiment, where the antiviral composition is a liquid, the viscosity is from about 0 cPs to about 5000 cPs, in another embodiment, from about 50 to about 500 cPs, and in another embodiment, from about 100 to about 400 cPs, as measured by Brookfield RV Viscometer using RV and/or LV Spindles at 22 °C +/- 3 °C.

[86] Surprisingly, it has been found that the antiviral composition may provide antiviral efficacy over a wide range of pH. Antiviral efficacy may be achieved at a pH of from 0 to about 14. More specifically, in one or more embodiments of the present invention, 3 log reduction or greater against non-enveloped viruses is achieved with antiviral compositions having a pH of greater than about 2.5, in other embodiments greater than about 3, in yet other embodiments greater than about 3.5, in other embodiments greater than about 4, in still yet other embodiments greater than about 4.5, and in still other embodiments, greater than about 5. In certain embodiments, 3 log reduction or greater against non-enveloped viruses is achieved with antiviral compositions having a pH of from about 4.5 to about 9, in other embodiments from about 5 to about 8.5, and in yet other embodiments from about 5.5 to about 7.5.

[87] In order to demonstrate the practice of the present invention, the following examples have been prepared and tested. The examples should not, however, be viewed as limiting the scope of the invention. The claims will serve to define the invention.

15

EXAMPLES

[*Bacteriophage propagation*]

[88] MS2 (obtained from ATCC) was grown to high titres on *E. coli* ATCC 15597. An exponentially growing culture of *E. coli* in LB broth supplemented with 2 mM CaCl₂ was divided into 200 microliter aliquots and inoculated with 200 microliters of serially diluted phage stock. The mixtures were added to 2.5 ml molten soft (0.7%) MS agar held at 44 °C and immediately poured over the surface of an LB agar plate. After 16 hours incubation at 37 °C, phage were harvested from plates demonstrating complete lysis of the *E. coli* lawn. To harvest the phage, 10 mL of sterile SM buffer was added to the surface of the plate and the soft agar was broken with a bent sterile glass rod. The broken agar was centrifuged for 10 minutes at 5000 G to remove debris and the supernatant containing purified phage was treated with chloroform and stored for up to 2 months at 4 °C. Prior to use, phage suspensions were allowed to equilibrate to room temperature.

[*Bacteriophage titre*]

[89] Infectious particles were counted by using a soft agar overlay technique. Molten, soft (0.7%) MS agar was dispensed in 2.5 ml aliquots in glass bottles and held at 44 °C. Phage-containing solutions were serially diluted in SM buffer at 20 °C and 0.1 ml added, together with 0.1 ml exponential culture of *E. coli* ATCC 15597 to the molten agar. The contents were gently mixed and poured over the surface of a nutrient agar plate. Plaques were countable after 24 hours incubation at 37 °C and results expressed as plaque forming units per milliliter (pfu ml⁻¹).

[*Virucidal suspension tests with MS2*]

[90] Suspension tests with MS2 were performed essentially as follows. Typically, 100 µl phage was added to 9.9 ml of antiviral composition. After the desired contact time at 25 °C, 0.1 ml suspension was neutralized by dilution into 9.9 ml D.E. broth. Further 10-fold serial dilutions were prepared in D.E. broth. The remaining active phage was quantified by infecting *E. coli* and using the soft agar overlay method as described above.

[*Virucidal suspension tests with mammalian viruses*]

[91] Virucidal suspension tests with mammalian viruses were performed using a modification of the Standard Test Method for Efficacy of Virucidal Agents Intended for Special Applications (ASTM E1052). Viral strains and indicator cells lines were as follows: Rhinovirus type 37, ATCC VR-1147 grown on MRC-5 human embryonic lung cells; Feline

calicivirus Strain F-9, ATCC VR-782 grown on CRFK feline kidney cells, Adenovirus type 2, ATCC VR-846 grown on A-549 human lung carcinoma cells; Rotavirus WA, ATCC VR-2018, grown on MA-104 rhesus monkey kidney cells; Herpes Simplex Type 1 Strain F(1), ATCC VR-733 grown on rabbit kidney cells (RK) from ViroMed Laboratories; Hepatitis A
5 Virus Strain HM-175 was grown on Fetal Rhesus monkey kidney cells (FRhK-4) from AppTec Laboratory Services; Canine Parvovirus Strain Cornell, ATCC VR-72017, was grown on A-72 canine tumor cells from ViroMed Laboratories. A 4.5 ml aliquot of each test substance was dispensed into separate sterile 15 ml conical tubes and each was mixed with a 0.5 ml aliquot of the stock virus suspension. The mixtures were vortex mixed for 10 seconds
10 and held the remainder of the 30 second exposure time at 33 ± 2 ° C. Immediately following the exposure period, a 0.1 ml aliquot was removed from each tube and the mixtures were titered by 10-fold serial dilutions and assayed for the presence of virus by infecting indicator cell lines. Cytopathic effect (CPE) was used in each case to indicate infection and TCID50 values were calculated by the method of Spearman Karber. Virus controls, neutralization
15 controls, and cytotoxicity controls were also performed.

[Preparation and testing of antiviral compositions]

[92] Example 1 - 95 % ethanol was mixed with water to form a 78 % by weight ethanol mixture.

[93] Example 2 - was prepared as described for Example 1, except that 1.25 wt. % of
20 1 M citric acid in water was added, with stirring, to form a homogeneous mixture.

[94] Example 3 - Powdered Synthalen CR (polyquaternium-37) was added to water in a flask, and mixed until a smooth gel was formed. 78 % ethanol was added to the flask, with stirring, to form a homogeneous mixture.

[95] Example 4 - Powdered Synthalen CR (polyquaternium-37) was added to water in
25 a flask, and mixed until a smooth gel was formed. 78 % ethanol was added to the flask, with stirring, to form a homogeneous mixture. 1.25 wt. % of 1 M citric acid in water was added, with mixing.

[96] The antiviral efficacy of Examples 1-4 were tested as described above for MS2, and the results are shown in Table 2.

Table 2

EXAMPLE	COMPOSITION	LOG REDUCTION, MS2 ¹
1	78% ethanol	0.2
2	78% ethanol + 0.25% citric acid	0.7
3	78% ethanol + 0.4% polyquaternium-37	0.9
4	78% ethanol + 0.25% citric acid + 0.4% polyquaternium-37	4.3

¹60 seconds at 25 °CExamples 5-13

- 5 **[97]** Example 5 was prepared by mixing 95 % ethanol with water to form a 70 % by weight ethanol mixture. Example 6 was prepared by dissolving urea in water to form a 10 wt. % mixture. Example 7 was prepared as for Example 5, except that urea was also added. Example 8 was prepared as for Example 7, except that polyquaternium-37 was also added. The pH of Example 8 was about 5.5. Example 9 was prepared as for Example 5, except that
- 10 polyquaternium-22 was also added. Example 10 was prepared as for Example 9, except that urea was also added. The pH of Example 10 was about 4.9. Example 11 was prepared as for Example 5, except that guanidine HCl was also added. The pH of Example 11 was about 7.6. Example 12 was prepared as for Example 11, except that polyquaternium-22 was also added. The pH of Example 12 was about 6.2. Example 13 was prepared as for Example 12. The pH
- 15 of Example 13 was about 5.8. The antiviral efficacy of Examples 5-13 were tested as described above for MS2, and the results are shown in Table 3.

Table 3

EXAMPLE	COMPOSITION	LOG REDUCTION MS2 ¹
5	70% ethanol	0
6	10 % urea in water	0
7	70% ethanol + 10% urea	0.9
8	70% ethanol + 10% urea + 0.4% polyquaternium-37	≥6.1
9	70% ethanol + 1% polyquaternium-22	0.7
10	70% ethanol + 10% urea + 0.4% polyquaternium-22	6.1
11	70% ethanol + 10% guanidine HCl	2.7
12	70% ethanol + 10% guanidine HCl + 0.4% polyquaternium-22	5.5
13	70% ethanol + 10% aminoguanidine HCl + 0.4% polyquaternium-22	5.8

¹60 seconds at 25 °C

Examples 14-15

[98] Example 14 was prepared as described for Example 1, and Example 15 was prepared as described for Example 4. The efficacy of Examples 14 and 15 against feline calicivirus was tested by using a modification of the Standard Test Method for Efficacy of Virucidal Agents Intended for Special Applications (ASTM E1052). The samples were tested by *in-vitro* virucidal suspension assay. The F-9 strain of Feline Calicivirus stock virus was obtained from the American Type Culture Collection, Manassas, VA (ATCC VR-782). A suspension of virus was exposed to the sample. At a pre-determined exposure time, an aliquot was removed, neutralized by serial dilution, and assayed for the presence of virus by infecting CRFK cells and measuring CPE as described hereinabove. Positive virus controls, cytotoxicity controls, and neutralization controls were assayed in parallel. Log reduction was calculated, and the results are shown in Table 4.

Table 4

EXAMPLE	COMPOSITION	LOG REDUCTION, FELINE CALICIVIRUS ¹
14	78% ethanol	3.4
15	78% ethanol + 0.25% citric acid + 0.4% polyquaternium-37	≥4.7

¹30 seconds at 33 °CExamples 16-17

[99] Example 16 was prepared as described for Example 2, and Example 17 was prepared as described for Example 4. The efficacy of Examples 16 and 17 against adenovirus type 2 was tested by using a modification of ASTM E1052. The samples were tested by *in-vitro* virucidal suspension assay. The Adenoid 6 strain of Adenovirus type 2 stock virus was obtained from the American Type Culture Collection, Manassas, VA (ATCC VR-846). A suspension of virus was exposed to the sample. At a pre-determined exposure time, an aliquot was removed, neutralized by serial dilution, and assayed for the presence of virus. Positive virus controls, cytotoxicity controls, and neutralization controls were assayed in parallel. Log reduction was calculated, and the results are shown in Table 5.

Table 5

EXAMPLE	COMPOSITION	LOG REDUCTION, ADENOVIRUS ¹
16	78% ethanol + 0.25% citric acid	1.3
17	78% ethanol + 0.25% citric acid + 0.4% polyquaternium-37	≥5.0

¹30 seconds at 33 °C

5

Examples 18-20

[100] Example 18 was prepared as described for Example 4, except that the concentration of ethanol was 70 % by weight. Example 19 was prepared as described for Example 4. Example 20 was prepared as described for Example 4, except that tartaric acid was used instead of citric acid. The mixtures were tested for efficacy against five different viruses, and the results are shown in Table 6.

[101] The efficacy of Examples 18-20 against rhinovirus type 37 was tested by using a modification of ASTM E1052. The samples were tested by *in-vitro* virucidal suspension assay. The 151-1 strain of Rhinovirus type 37 stock virus was obtained from the American Type Culture Collection, Manassas, VA (ATCC VR-1147). A suspension of virus was exposed to the sample. At a pre-determined exposure time, an aliquot was removed, neutralized by serial dilution, and assayed for the presence of virus by infecting MRC-5 cells and measuring CPE as described hereinabove. Positive virus controls, cytotoxicity controls, and neutralization controls were assayed in parallel.

[102] The efficacy of Examples 18-20 against rotovirus was tested by using a modification of ASTM E1052. The samples were tested by *in-vitro* virucidal suspension assay. The WA stock virus was obtained from the American Type Culture Collection, Manassas, VA (ATCC VR-2018). A suspension of virus was exposed to the sample. At a pre-determined exposure time, an aliquot was removed, neutralized by serial dilution, and assayed for the presence of virus by infecting MA-104 cells and measuring CPE as described hereinabove. Positive virus controls, cytotoxicity controls, and neutralization controls were assayed in parallel.

TABLE 6

EX.	COMPOSITION	MS2 ¹	FELINE CALICIVIRUS ²	ADENOVIRUS ³	ROTAVIRUS ⁴	RHINOVIRUS ⁵
18	70% ethanol + 0.25% citric acid + 0.4% polyquaternium-37	2.4	≥4.7	≥5.0	≥3.8	≥3.3
19	78% ethanol + 0.25% citric acid + 0.4% polyquaternium-37	3.7	≥4.7	≥5.0	≥3.8	≥3.3
20	78% ethanol + 0.25% tartaric acid + 0.4% polyquaternium-37	4.4	≥4.7	≥5.0	≥3.8	≥3.3

¹60 seconds at 25 °C; average of replicates; ²⁻⁵30 seconds at 33 °C

Examples 21-22

[103] Example 21 was prepared by mixing 95 % ethanol with water to form a 78 % by weight ethanol mixture. Example 22 was prepared as for Example 21, except that polyquaternium-37 was also added. The efficacy of Examples 21-22 against hepatitis A virus was tested by using a modification of ASTM E1052. The samples were tested by *in-vitro* virucidal suspension assay. The HM-175 strain of Hepatitis A virus (HAV) stock virus was obtained from AppTec Laboratory Services, Camden, N.J. A suspension of virus was exposed to the sample. At a pre-determined exposure time, an aliquot was removed, neutralized by serial dilution, and assayed for the presence of virus by infecting FRhK-4 cells and measuring CPE as described hereinabove. Positive virus controls, cytotoxicity controls, and neutralization controls were assayed in parallel. Results are shown in Table 7.

Table 7

EXAMPLE	COMPOSITION	LOG REDUCTION, HEPATITIS A VIRUS ¹
21	78% ethanol	1.25
22	78% ethanol + 1% polyquaternium-37	3.0

¹60 seconds at 25 °CExamples 23-24

[104] Example 23 was prepared as for Example 18. Example 24 represents an antibacterial hand sanitizer composition similar to a product currently commercially available, the label of which is marked with U.S. Pat. No. 6,080,417. The efficacy of Examples 23-24 against Canine parvovirus was tested by using a modification of ASTM E1052. The samples were tested by *in-vitro* virucidal suspension assay. The virus tested was Strain Cornell, ATCC VR-2017, cell line A-72 canine tumor cells, ATCC CRL-1542. A suspension of virus was exposed to the sample. At a pre-determined exposure time, an aliquot was removed, neutralized by serial dilution, and assayed for the presence of virus by infecting CRFK cells and measuring CPE as described hereinabove. Positive virus controls, cytotoxicity controls, and neutralization controls were assayed in parallel. Results are shown in Table 8.

Table 8

EXAMPLE	COMPOSITION	LOG REDUCTION, CANINE PARVOVIRUS
23	70% ethanol + 0.25% citric acid + 0.4% polyquaternium-37	1.0
24	Manorapid Synergy	0

30 seconds at 33 °C

Examples 25-26

- 5 [105] Examples 25 – 26 represent antibacterial hand sanitizer compositions similar to products currently commercially available. The compositions were formulated as shown in Table 9, and tested for efficacy against MS2.

Table 9

EXAMPLE	COMPOSITION	LOG REDUCTION, MS2 ¹
25	62% ethanol in carbomer gel	0
26	Manorapid Synergy	0.8

10

¹60 seconds at 25 °C

- [106] Fingerpad in vivo testing of Examples 19 and 23 was performed according to ASTM E 1838 – 96, “Standard Test Method for Determining the Virus-Eliminating Effectiveness of Liquid Hygienic Handwash Agents Using the Fingerpads of Adult
- 15 Volunteers.” The efficacy of the compositions was tested against feline calicivirus and rotavirus, and the results are shown in Table 10.

Table 10

EXAMPLE	COMPOSITION	LOG REDUCTION, FELINE CALICIVIRUS ¹	LOG REDUCTION, ROTAVIRUS ¹
Example 23	62% ethanol in carbomer gel	0.6	2.5
Example 19	78% ethanol + 0.25% citric acid + 0.4% polyquaternium-37	1.6	3.0

¹log₁₀ reduction at 15 seconds

Examples 25-26

[107] The efficacy of Examples 25-26 against herpes virus (an enveloped virus) was tested by *in-vitro* virucidal suspension assay. (Herpes Simplex Type 1 Strain F(1), ATCC VR-733 grown on rabbit kidney cells (RK) from ViroMed Laboratories) A suspension of virus was exposed to the sample. At a pre-determined exposure time, an aliquot was removed, neutralized by serial dilution, and assayed for the presence of virus by infecting RK cells and measuring CPE as described hereinabove. Positive virus controls, cytotoxicity controls, and neutralization controls were assayed in parallel. Results are shown in Table 11.

Table 11

EXAMPLE	COMPOSITION	LOG REDUCTION HERPES VIRUS ¹
25	62% ethanol in carbomer gel	≥5.5
26	62 % ethanol + 1.5 % polyquaternium-37	≥4.5

¹60 seconds at room temperature

[Virucidal suspension tests with adenovirus and poliovirus according to EN 14476:2005]

[108] Virucidal suspension tests with mammalian viruses were performed using European Standard 14476:2005.

[109] The adenovirus viral strain used was Adenovirus Type 5, strain Adenoid 75, ATCC VR-5 obtained from the Institute of Medical Virology, Hannover Medical School, Hannover Germany. Adenovirus was grown on A549 human lung epithelial carcinoma cells also procured from Institute of Medical Virology, Hannover Medical School, Hannover Germany.

[110] The poliovirus viral strain was Poliovirus Type 1, LSc-2ab (Chiron-Behring) obtained from Eurovir, Luckenwalde, Germany. Poliovirus was grown on buffalo green monkey kidney cells obtained from Institut für angewandte Zellkultur, München, Germany.

[111] A 0.1 ml aliquot of the stock virus suspension virus was added to 0.1 ml of phosphate buffered saline and vortex mixed. A 0.8 ml aliquot of test substance was added to the tube, vortex mixed and held for the remainder of the exposure time in a water bath at 20±1°C. Immediately following the exposure period (ranging from 30 seconds to 5 minutes), the test mixture was neutralized via 10-8 dilution and assayed for the presence of virus by

infecting the indicator cell lines. The infectivity was determined through measurement of the cytopathic effect ten days after infection. Calculation of the virus concentration was carried out by the Spearman-Kärber method to determine log₁₀TCID₅₀/mL. Experimental controls included a 0.7% formaldehyde solution, virus controls and neutralization controls.

5 *[Preparation and testing of antiviral compositions]*

[112] Example 27 – was prepared as described for Example 5.

[113] Example 28 – was prepared by adding copper gluconate powder to water to form a solution. Ethanol was added, with stirring, to form a homogeneous mixture having the composition shown in Table 12.

10 [114] Example 29 – powdered Synthalen CR (polyquaternium-37) was added to water in a flask, and mixed until a smooth gel was formed. 70% ethanol was added to the flask, with stirring, to form a homogeneous mixture.

[115] Example 30 – was prepared as described for Example 29, except that a sufficient amount of a solution of copper gluconate in water was added, with stirring, to form a
15 homogeneous mixture having the composition shown in Table 12.

[116] The antiviral efficacy of Examples 27-30 was tested as described above for EN 14476:2005, and the results, in terms of log reduction, are shown in Table 12.

Table 12

EXAMPLE	COMPOSITION	ADENOVIRUS		POLIOVIRUS	
		30 SEC	1 MIN	30 SEC	1 MIN
27	70 % ethanol	>5.69	>5.69	0.25	0.75
28	70 % ethanol + 0.08 % Cu gluconate	2.37	3.87	0.50	0.50
29	70 % ethanol + 0.4 % polyquaternium-37	>4.81	>4.81	0.00	0.00
30	70 % ethanol + 0.4 % polyquaternium-37 + 0.08 % Cu gluconate	>5.00	>5.00	0.50	>4.00

20 [117] Thus, it should be evident that the present invention provides a method for inactivating virus. In certain embodiments, a virucidal composition comprising alcohol, a cationic oligomer or polymer, and an enhancer exhibits an efficacy against non-enveloped viruses that is higher than the efficacy of the same composition but not comprising the enhancer. In one embodiment, the virucidal composition exhibits an efficacy against non-
25 enveloped viruses that is at least about 0.5 log reduction higher than the efficacy of the same composition but not comprising the enhancer. In another embodiment, the composition exhibits an efficacy against non-enveloped viruses that is at least about 1 log reduction higher

than the efficacy of the same composition but not comprising the enhancer.

[118] The antiviral composition is highly efficacious for household cleaning applications (e.g., hard surfaces like floors, countertops, tubs, dishes and softer cloth materials like clothing, sponges, paper towels, etc.), personal care applications (e. g. lotions, shower
5 gels, soaps, hand sanitizers, shampoos, wipes) and industrial and hospital applications (e. g., disinfection of instruments, surfaces, medical devices, gloves). This composition is efficacious for rapidly sanitizing or de-germing surfaces that are infected or contaminated with Gram negative bacteria, fungi, parasites, Gram positive bacteria, enveloped viruses, and non-enveloped viruses. The efficacy of alcoholic compositions comprising a C1-6 alcohol, an
10 acid, and a cationic oligomer or polymer against resident and transient flora is described in co-pending U.S. Provisional Patent Application Serial No. 60/771,784, which is hereby incorporated by reference in its entirety.

[119] Various modifications and alterations that do not depart from the scope and spirit of this invention will become apparent to those skilled in the art. This invention is not to be
15 duly limited to the illustrative embodiments set forth herein.

What is claimed is:

1. A method of inactivating non-enveloped virus particles, the method comprising:
 contacting non-enveloped virus particles with a virucidally-enhanced alcoholic composition comprising at least about 50 wt. % of a C₁₋₆ alcohol, and from about 0.02 to about 20 percent by weight of one or more enhancers selected from the group consisting of cationic oligomers and polymers, from about 0.01 to about 1 percent by weight of a zinc or copper compound, and less than about 0.05 percent by weight acid, based upon the total weight of the alcoholic composition, wherein said cationic oligomer or polymer comprises a polyquaternium polymer, a cationic copolymer of saccharide and a synthetic cationic monomer, a cationic polyalkylene imine, a cationic ethoxy polyalkylene imine, a cationic poly[N-[3-(dialkylammonio)alkyl] N'[3-(alkyleneoxyalkylene dialkylammonio)alkyl]urea dichloride], a vinyl caprolactam/VP/dialkylaminoalkyl alkylate copolymer, or a mixture thereof.

2. A method of producing a topical virucidal effect on mammalian skin against non-enveloped virus by applying a virucidally-enhanced alcoholic composition comprising at least about 50 wt. % of a C₁₋₆ alcohol, and from about 0.02 to about 20 percent by weight of one or more enhancers selected from the group consisting of cationic oligomers and polymers, from about 0.01 to about 1 percent by weight of a zinc or copper compound, and less than about 0.05 percent by weight acid, based upon the total weight of the alcoholic composition, wherein said cationic oligomer or polymer comprises a polyquaternium polymer, a cationic copolymer of saccharide and a synthetic cationic monomer, a cationic polyalkylene imine, a cationic ethoxy polyalkylene imine, a cationic poly[N-[3-(dialkylammonio)alkyl] N'[3-(alkyleneoxyalkylene dialkylammonio)alkyl]urea dichloride], a vinyl caprolactam/VP/dialkylaminoalkyl alkylate copolymer, or a mixture thereof.

3. A method of enhancing the efficacy of a C₁₋₆ alcohol against non-enveloped virus in a topical application to a surface, the method comprising:
 combining said C₁₋₆ alcohol with from about 0.02 to about 20 percent by

weight of one or more enhancers selected from the group consisting of cationic oligomers and polymers, from about 0.01 to about 1 percent by weight of a zinc or copper compound, and with less than about 0.05 percent by weight acid, based upon the total weight of the alcoholic composition, to form an antiviral composition that includes at least about 50 wt. % of said alcohol, wherein said cationic oligomer or polymer comprises a polyquaternium polymer, a cationic copolymer of saccharide and a synthetic cationic monomer, a cationic polyalkylene imine, a cationic ethoxy polyalkylene imine, a cationic poly[N-[3-(dialkylammonio)alkyl] N'[3-(alkyleneoxyalkylene dialkylammonio)alkyl]urea dichloride], a vinyl caprolactam/VP/dialkylaminoalkyl alkylate copolymer, or a mixture thereof.

4. A virucidally-enhanced alcoholic composition comprising:

at least about 50 wt. % of C₁₋₆ alcohol; from about 0.02 to about 20 percent by weight of one or more enhancers selected from the group consisting of cationic oligomers and polymers; from about 0.01 to about 1 percent by weight of a zinc or copper compound; and less than about 0.05 percent by weight acid, based upon the total weight of the alcoholic composition, wherein said cationic oligomer or polymer comprises a polyquaternium polymer, a cationic copolymer of saccharide and a synthetic cationic monomer, a cationic polyalkylene imine, a cationic ethoxy polyalkylene imine, a cationic poly[N-[3-(dialkylammonio)alkyl] N'[3-(alkyleneoxyalkylene dialkylammonio)alkyl]urea dichloride], a vinyl caprolactam/VP/dialkylaminoalkyl alkylate copolymer, or a mixture thereof, and wherein said virucidally-enhanced composition exhibits an efficacy against non-enveloped viruses that is higher than the efficacy of the same composition but not comprising said enhancer.

5. The method of any of claims 1-3, or the composition of claim 4, wherein said method is operative to inactivate enveloped viruses or microbes including gram positive bacteria, gram negative bacteria, fungi, or parasites, and wherein said method further comprises:

contacting enveloped virus or microbes with said composition.

6. The method of any of claims 1-3, or the composition of claim 4, wherein said cationic oligomer or polymer is selected from the group consisting of polyquaternium-2, polyquaternium-4, polyquaternium-5, polyquaternium-6, polyquaternium-7, polyquaternium-10, polyquaternium-11, polyquaternium-16, polyquaternium-22, polyquaternium-24, polyquaternium-28, polyquaternium-32, polyquaternium-37, polyquaternium-39, polyquaternium-42, polyquaternium-43, polyquaternium-44, polyquaternium-46, polyquaternium-47, polyquaternium-51, polyquaternium-53, polyquaternium-55, polyquaternium-57, polyquaternium-58, polyquaternium-59, polyquaternium-60, polyquaternium-63, polyquaternium-64, polyquaternium-65, polyquaternium-68, and mixtures thereof.

7. The method of any of claims 1-3, or the composition of claim 4, wherein said copper compound is copper gluconate.

8. The method or the composition of claim 7, wherein said enhancer is polyquaternium-37.

9. A method of inactivating non-enveloped virus particles, the method comprising:
contacting non-enveloped virus particles with a virucidally-enhanced alcoholic composition comprising at least about 50 wt. % of a C₁₋₆ alcohol, and from about 0.25 to about 20 percent by weight of an enhancer, based upon the total weight of the alcoholic composition, wherein said enhancer is a chaotropic agent.

10. The method of claim 9, wherein said chaotropic agent comprises urea, thiourea, guanidine HCl, guanidine thiocyanate, aminoguanidine HCl, aminoguanidine bicarbonate, guanidine carbonate, guanidine phosphate, or mixtures thereof.

11. The method of any of the preceding claims, wherein said method exhibits an increased log reduction against said non-enveloped virus particles, when compared to the

log reduction of a composition comprising the same amount of said C₁₋₆ alcohol, and less than about 0.02 wt. % of said enhancer, based upon the total weight of the composition.

12. The method or composition of any of the preceeding claims, wherein said non-enveloped virus particles are selected from members of the families *Picornaviridae*, *Reoviridae*, *Caliciviridae*, *Adenoviridae* and *Parvoviridae*.

13. The method or composition of any of the preceeding claims, wherein said non-enveloped virus particles are selected from adenovirus, feline calicivirus, norovirus, papillomavirus, poliovirus, rhinovirus, hepatitis A virus, parvovirus, and rotavirus.