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(71) Applicant: MICROBAN PRODUCTS COMPANY

[US/US]; 11400 Vanstory Drive, Huntersville, NC 28078 (US).

(72) Inventors: LAN, Tian; 13232 Willow Breeze Lane,

Huntersville, NC 28078 (US). ONG, Ivan, W.; 6508 Tall

Oaks Trail, Charlotte, NC 28210 (US). SLOAN, Gina,

Parise; 218 Wilson Farm Road, Statesville, NC 28625 (US).

WELCH, Karen, Terry; 5038 Century Drive, Kannapolis,

NC 28081 (US).

(74) Agent: JACKSON, Susan, S. et al.; Nelson Mullins Riley

& Scarborough LLP, Bank of America Corporate Center,

42nd Floor, 100 North Tryon Street, Charlotte, NC 28202

(US).

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(54) Title: TOUCH SCREEN CLEANING AND PROTECTANT COMPOSITION

(57) Abstract: A cleaning and protectant composition and a method for using the composition on an article such as a touch screen or screen protector for a mobile device, computer, or other electronic device. The composition comprises a polymeric film former. The polymeric film former is an oxazoline homopolymer or an extended or a modified polymer based on an oxazoline homopolymer. The polymeric film former forms a retention coating to retain components within the composition for an extended period of time. The composition is also present in a kit with an application device.



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TOUCH SCREEN CLEANING AND PROTECTANT COMPOSITION

CROSS-REFERENCE TO RELATED APPLICATION

The present application claims priority from U.S. provisional patent application
5 62/340,310, filed on May 23, 2016, and U.S. utility patent application 15/599,774, filed on
May 19, 2017, incorporated by reference herein in their entirety.

FIELD OF THE INVENTION

The present invention relates to a cleaning and protectant composition, more
particularly to a cleaning and protectant composition suitable for use on a touch screen or
10 screen protector of a mobile device, computer, or other electronic device.

BACKGROUND OF THE INVENTION

The number of mobile devices, computers, and other devices having touch screens
has exploded. These screens can have lots of bacteria and germs from repeated use and
touch without cleaning. Cleaning products to date that are available for this purpose are
15 typically general glass cleaners. Most of these glass cleaners are alcohol-based liquid
products. Such products are only limited to cleaning dirt or contamination off of a screen
surface without leaving a protection film. A touch screen can be re-contaminated quickly
and has to be cleaned frequently to keep it clean.

Other polymer and water-based products suffer from a number of problems
20 including, but not limited to, reduction of the transparency and clearness of the screen, a
short-life cycle before reapplication, product build-up on the screen negatively affecting
transparency, and difficulty in removal.

Thus, there is a need for a product that overcomes the problems set forth above and
is suitable for use on a touch screen or screen protector or on other articles where such
25 problems are also of concern.

SUMMARY OF THE INVENTION

The present invention relates to a cleaning and protectant composition suitable for use on an article such as a touch screen or screen protector for a mobile device, computer, or other device.

5 The present invention also relates to a cleaning and protectant composition suitable for use on an article such as a clear plastic substrate or coating material such as food wrap.

In an embodiment of the invention, the composition comprises a carrier and a polymeric film former, wherein the polymeric film former is an oxazoline homopolymer or an extended or a modified polymer based on an oxazoline homopolymer.

10 In an embodiment of the invention, the polymeric film former forms a retention coating to retain components within the composition for an extended period of time.

In an embodiment of the invention, an article is provided. The article comprises a surface having been treated with a composition comprised of a carrier and a polymeric film former, wherein the polymeric film former is an oxazoline homopolymer or an
15 extended or a modified polymer based on an oxazoline homopolymer. The article is, for example, a touch screen, a screen protector, a wipe, food wrap, among others.

In an embodiment of the invention, a method of using a composition is provided. The method comprises obtaining a composition comprised of a carrier and a polymeric film former, wherein the polymeric film former is an oxazoline homopolymer or an
20 extended or a modified polymer based on an oxazoline homopolymer; and applying the composition to an article.

In an embodiment of the invention, a kit is provided. The kit comprises an application device and a composition comprised of a carrier and a polymeric film former, wherein the polymeric film former is an oxazoline homopolymer or an extended or a

modified polymer based on an oxazoline homopolymer. The application device is used to apply the composition to the article, for example, to the touch screen or screen protector.

Further areas of applicability of the present invention will become apparent from the detailed description provided hereinafter. It should be understood that the detailed description and specific examples, while indicating the preferred embodiments of the invention, are intended for purposes of illustration only and are not intended to limit the scope of the invention.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

The following description of the embodiments of the present invention is merely exemplary in nature and is in no way intended to limit the invention, its application, or uses. The present invention has broad potential application and utility, which is contemplated to be adaptable across a wide range of industries. The following description is provided herein solely by way of example for purposes of providing an enabling disclosure of the invention, but does not limit the scope or substance of the invention.

As used herein, the terms "microbe" or "microbial" should be interpreted to refer to any of the microscopic organisms studied by microbiologists or found in the use environment of a treated article. Such organisms include, but are not limited to, bacteria and fungi as well as other single-celled organisms such as mold, mildew and algae. Viral particles and other infectious agents are also included in the term microbe.

"Antimicrobial" further should be understood to encompass both microbicidal and microbistatic properties. That is, the term comprehends microbe killing, leading to a reduction in number of microbes, as well as a retarding effect of microbial growth, wherein numbers may remain more or less constant (but nonetheless allowing for slight increase/decrease).

For ease of discussion, this description uses the term antimicrobial to denote a broad spectrum activity (e.g. against bacteria and fungi). When speaking of efficacy against a particular microorganism or taxonomic rank, the more focused term will be used (e.g. antifungal to denote efficacy against fungal growth in particular).

5 Using the above example, it should be understood that efficacy against fungi does not in any way preclude the possibility that the same antimicrobial composition may demonstrate efficacy against another class of microbes.

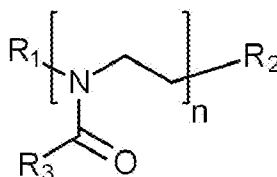
For example, discussion of the strong bacterial efficacy demonstrated by a disclosed embodiment should not be read to exclude that embodiment from also
10 demonstrating antifungal activity. This method of presentation should not be interpreted as limiting the scope of the invention in any way.

Further, the term "or" as used in this disclosure and the appended claims is intended to mean an inclusive "or" rather than an exclusive "or." That is, unless specified otherwise, or clear from the context, the phrase "X employs A or B" is intended to mean
15 any of the natural inclusive permutations. That is, the phrase "X employs A or B" is satisfied by any of the following instances: X employs A; X employs B; or X employs both A and B. In addition, the articles "a" and "an" as used in this application and the appended claims should generally be construed to mean "one or more" unless specified otherwise or clear from the context to be directed to a singular form. Throughout the
20 specification and claims, the following terms take at least the meanings explicitly associated herein, unless the context dictates otherwise. The meanings identified below do not necessarily limit the terms, but merely provided illustrative examples for the terms. The meaning of "a," "an," and "the" may include plural references, and the meaning of "in" may include "in" and "on." The phrase "in one embodiment," as used herein does not
25 necessarily refer to the same embodiment, although it may.

In an embodiment of the invention, a composition is provided that is suitable for use as a cleaner and a protectant on an article. Examples of articles include, but are not limited to, a touch screen and/or screen protector of a mobile device, computer, or other electronic device, a wipe, food wrap, a plastic substrate, and a plastic coating.

5 The composition comprises a polymeric film former. In a preferred embodiment of the invention, the polymeric film former forms a retention coating to retain components within the composition for an extended period of time. Examples of such components include, but are not limited to, colorants, dyes, pigments, fragrance, biocidal agents, and other components.

10 In an embodiment of the invention, the polymeric film former is an oxazoline homopolymer. As another feature of the invention, the oxazoline homopolymer has the following structure:



wherein

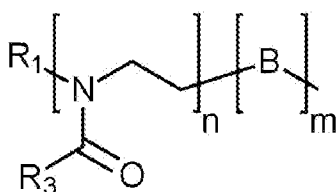
15 R_1 and R_2 are end groups determined by the polymerization techniques used to synthesize oxazoline homopolymer. R_1 and R_2 are independently selected and include, but are not limited to, hydrogen, alkyl, alkenyl, alkoxy, alkylamino, alkynyl, allyl, amino, anilino, aryl, benzyl, carboxyl, carboxyalkyl, carboxyalkenyl, cyano, glycosyl, halo, hydroxyl, oxazolinium mesylate, oxazolinium tosylate, oxazolinium triflate, silyl
 20 oxazolinium, phenolic, polyalkoxy, quaternary ammonium, thiol, or thioether groups. Alternatively, R_2 could include a macrocyclic structure formed during synthesis as a consequence of intramolecular attack.

For example, R_1 is a methyl group and R_2 is oxazolinium tosylate if methyl tosylate is used as the initiator in the cationic initiated polymerization of oxazoline.

R_3 is an end group determined by the type of oxazoline used in the preparation of the polymeric film former of this invention. R_3 includes, but is not limited to, hydrogen, alkyl, alkenyl, alkoxy, aryl, benzyl, hydroxyalkyl, or perfluoroalkyl. For example, R_3 is an ethyl group if ethyloxazoline is the monomer used to prepare the polymeric film former for the present invention.

n is the degree of oxazoline polymerization in the homopolymer. n is in a range of 1 to 1,000,000. Preferably, n is in a range of 500 to 250,000; most preferably, n is in a range of 2500 to 100,000.

Similar to oxazoline homopolymer, extended or modified polymers with some variations based on the oxazoline homopolymer are also suitable for the present invention. The techniques and options for performing chemical or molecular structure variations or modifications to oxazoline should be familiar to those skilled in the art. A class of extended or modified polymers based on oxazoline homopolymer can be represented with the following molecular structure:

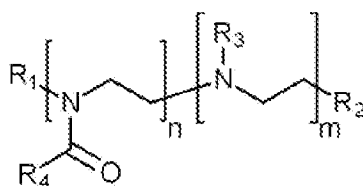


wherein

B is additional monomer repeating unit linked to oxazoline in a copolymer. The types of arrangement of the repeating units between B and oxazoline in the copolymer can include, but are not limited to, block, alternating, periodic, or combinations thereof. There is no limitation as to the types of B that can be used to copolymerize with or modify the oxazoline of the present invention.

n is the degree of polymerization for an oxazoline repeating unit; n in the copolymer is in a range of 1 to 1,000,000 and the degree of polymerization for B repeating unit in the copolymer m is in a range of 0 to 500,000 at the same time. Preferably, n is in a range of 500 to 250,000 and m is in a range of 20 to 10,000; and most preferably, n is in a range of 2500 to 100,000 and m is in a range of 50 to 5,000. In addition to linking B to ethyloxazoline through copolymerization, B could also be linked to oxazoline as an end group in a cationic polymerization by using B as a cationic initiator if B itself is already a quaternary ammonium compound.

Not intended to be all inclusive, B can be, for example, ethyleneimine with the following molecular structure:



wherein

R₁ and R₂ end groups have the same definition as those outlined for oxazoline homopolymer.

R₃ includes, but is not limited to, hydrogen, alkyl, alkenyl, alkoxy, aryl, benzyl, hydroxyalkyl, or perfluoroalkyl.

R₄ includes, but is not limited to, hydrogen, alkyl, alkenyl, alkoxy, aryl, benzyl, hydroxyalkyl, or perfluoroalkyl.

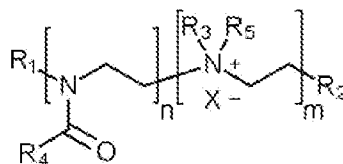
m is in a range of 0 to 500,000; preferably, in a range of 20 to 10,000; and most preferably, in a range of 50 to 5,000.

n is in a range of 1 to 1,000,000; preferably, 500 to 250,000; most preferably, in a range of 2500 to 100,000.

The synthesis of oxazoline and ethyleneimine copolymer can be phased into two steps, for example. In a first step, a cationic ring opening polymerization technique can be used to make polyoxazoline homopolymer. In a second step, the polyoxazoline made in the first step can be hydrolyzed to convert part of polyoxazoline repeating units into polyethyleneimine. Alternatively, oxazoline-ethyleneimine copolymer can be made with the appropriate respective monomers, an oxazoline and an aziridine. The result would be a cationic polymer having the above structure.

The degree of polymerization for oxazoline repeating unit n in the copolymer is in a range of 1 to 1,000,000 and the degree of polymerization for ethyleneimine repeating unit in the copolymer m is in a range of 0 to 500,000 at the same time. Preferably, n is in a range of 500 to 250,000 and m is in a range of 20 to 10,000, and most preferably n is in a range of 2500 to 100,000 and m is in a range of 50 to 5,000.

Alternatively, the nitrogen in the ethyleneimine repeating unit could be further quaternized to generate the following cationic copolymer:



15

Any quaternization technique that is familiar to those skilled in the art could be used to quaternize the polymer of this example. R_1 , R_2 , R_3 and R_4 have the same meaning as those designated in the above oxazoline-ethyleneimine copolymer. R_5 includes, but is not limited to, a hydrogen, methyl, ethyl, propyl, or other types of alkyl group. The corresponding anion X^- is a halogen, sulfonate, sulfate, phosphonate, phosphate, carbonate/bicarbonate, hydroxy, or carboxylate.

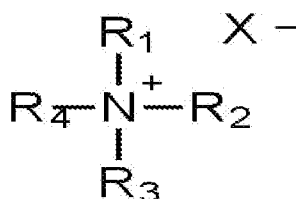
20

The ranges for n and m are also the same as those described in oxazoline-ethyleneimine copolymer.

Examples of commercially available polyethyloxazolines include, but are not limited to, Aquazol 500 from Polymer Chemistry Innovations, Inc.

In an embodiment of the invention, the composition optionally comprises a biocidal agent. A non-exhaustive list of biocidal compounds includes, but is not limited to, triclosan, zinc pyrithione, metal salts and oxides, phenols, botanicals, halogens, peroxides, heterocyclic antimicrobials, aldehydes, and alcohols.

In a preferred embodiment of the invention, the biocidal agent is a quaternary ammonium compound (QAC) with the following molecular structure:



10 wherein

R_1 , R_2 , R_3 , and R_4 are independently selected and include, but are not limited to, alkyl, alkoxy, or aryl, either with or without heteroatoms, or saturated or non-saturated. Some or all of the functional groups may be the same.

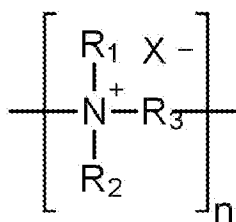
The corresponding anion X^- includes, but is not limited to, a halogen, sulfonate, sulfate, phosphonate, phosphate, carbonate/bicarbonate, hydroxy, or carboxylate.

QACs include, but are not limited to, n-alkyl dimethyl benzyl ammonium chloride, di-n-octyl dimethyl ammonium chloride, dodecyl dimethyl ammonium chloride, n-alkyl dimethyl benzyl ammonium saccharinate, and 3-(trimethoxysilyl) propyldimethyloctadecyl ammonium chloride.

20 Combinations of monomeric QACs are preferred to be used for the invention. A specific example of QAC combination is N-alkyl dimethyl benzyl ammonium chloride (40%); N-octyl decyl dimethyl ammonium chloride (30%); di-n-decyl dimethyl ammonium chloride (15%); and di-n-dioctyl dimethyl ammonium chloride (15%). The

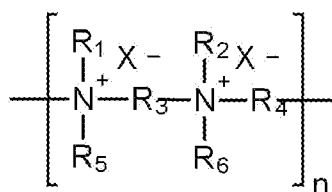
percentage is the weight percentage of individual QAC based on the total weight of blended QACs composition.

Polymeric version of the QACs with the following structures can also be used for the invention.



5

or



10 wherein

R_1 , R_2 , R_5 , and R_6 , independently, include, but are not limited to, hydrogen, methyl, ethyl, propyl or other longer carbon alkyl groups.

R_3 and R_4 are independently selected and include, but are not limited to, methylene, ethylene, propylene or other longer alkylene linking groups.

15 n is the degree of polymerization; n is an integer in a range of from 2 to 10,000.

Examples of cationic polymers with the above structure, include but are not limited to, polyamines derived from dimethylamine and epichlorohydrin such as Superfloc C-572 commercially available from Kemira Chemicals.

20 Still another polymeric QAC suitable for the invention is polydiallyldimethylammonium chloride or polyDADMAC.

Yet another class of QACs useful for the present invention are those chemical compounds with biguanide moiety in the molecule. Examples of this class of cationic antimicrobials include, but are not limited to, PHMB and chlorhexidine.

Examples of commercially available quaternary ammonium compounds include, but are not limited to, Bardac 205M and 208M from Lonza, and BTC885 from Stepan Company.

5 The composition of the present invention not only provides an initial cleaning action but also leaves a protection film on an article such as a touch screen for continuous protection of the screen from re-contamination.

In an embodiment of the invention, the composition comprises a cleaning and soil resistant agent.

10 Optionally, the composition comprises a fragrance, a colorant, or a combination thereof. The fragrance, colorant, or combination thereof can be added to the composition for aesthetic appeal, for example.

The composition may further comprise other components including, but not limited to, a surfactant, a stain resistant agent, an oil and/or water repellent, and a carrier such as water or a solvent.

15 Examples of solvents include, but are not limited to, water and low molecular weight alcohols such as C1 to C8 alkanols. Specific examples include, but are not limited to, ethanol, isopropyl alcohol, butanol, pentanol, and combinations thereof.

Another class of solvents for use in the invention includes alkylene glycol ether. Examples include, but are not limited to, ethylene glycol monopropyl ether, ethylene glycol monobutyl ether, ethylene glycol monohexyl ether, ethylene glycol monohexyl ether, diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, diethylene glycol monobutyl ether, diethylene glycol monohexyl ether, triethylene glycol monomethyl ether, triethylene glycol monoethyl ether, triethylene glycol monobutyl ether, propylene glycol methyl ether, propylene glycol methyl ether acetate, propylene glycol n-butyl ether, dipropylene glycol n-butyl ether, dipropylene glycol methyl ether, dipropylene

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glycol methyl ether acetate, propylene glycol n-propyl ether, dipropylene glycol n-propyl ether, and tripropylene glycol methyl ether.

Another class of solvents for use in the invention is based on terpenes and their derivatives such as terpene alcohols, terpene esters, terpene ethers, or terpene aldehydes.

5 Examples of solvents, include but are not limited to, pine oil, lemon oil, limonene, pinene, cymene, myrcene, fenchone, borneol, nopol, cineole, ionone and the like.

In an embodiment of the invention, the composition is in the form of a liquid. The liquid composition can be packaged and used in a spray bottle such as trigger spray or aerosol. In a spray application, the composition can be sprayed onto a contaminated or
10 dirty screen and then wiped out clean and dry. As the result of such a cleaning action, the screen surface is cleaned and, subsequently, a clear dry film is deposited on the screen for continuous protection for an extended period of time.

In an embodiment, a wipe impregnated or otherwise treated with the liquid composition of the present invention can be used. The wipe having the liquid composition
15 thereon can be used to wipe a contaminated surface such as a touch screen or screen protector (such as back and forth a couple of times) until the surface comes clean. After the wiping process, a film is automatically deposited onto the screen, and once it is dry, it provides continuous protection to the screen for an extended period of time.

The composition of the invention not only provides initial cleaning activity but
20 also leaves a protection film on a screen immediately after cleaning. It keeps screen clean, sanitized, and pleasant for an extended period of time without frequent cleaning.

In an embodiment of the invention, a kit is provided. The kit comprises an application device and the composition described herein comprised of a carrier and a polymeric film former, wherein the polymeric film former is an oxazoline homopolymer
25 or an extended or a modified polymer based on an oxazoline homopolymer. The

application device is any device used to apply the composition to a surface of the touch screen or other article. The application device is preferably selected from the group consisting of a spray bottle, sponge, a wipe, and a combination thereof.

In an embodiment of the invention, a method of using is provided. The method
5 comprises obtaining the composition described herein comprised of a carrier and a polymeric film former, wherein the polymeric film former is an oxazoline homopolymer or an extended or a modified polymer based on an oxazoline homopolymer; and applying the composition to an article. The composition may be applied with the application device.

10 Alternatively, the composition may be applied by other application techniques and methods including, but not limited to, industrial coating techniques. Examples of industrial coating techniques include, but are not limited to, slot coating, curtain coating, dip coating, roll coating, kiss coating, spray coating, and knife coating.

Among the benefits of the composition of the present invention are that it provides
15 aesthetic enhancement, increases the useful life of a touch screen, does not affect the transparency of a touch screen, and is easily removable, among other benefits. The use of the polymeric film former creates a retention coating for coloration and aroma effects as well as provides finger print protection and protection from scratching.

Examples

20 The following examples illustrate liquid formulations made in accordance with various aspects of the present invention. The testing results on these formulations demonstrate the desired residual sanitizing or disinfecting performance once being applied onto surfaces and dried.

Formulations were tested for residual efficacy using the EPA 01-1A protocol.
25 Briefly, bacteria were added to a glass slide and allowed to dry on the surface. The

formulation was then sprayed onto the surface and dried to form a transparent film. Once a film had formed, the glass slide was exposed to alternating wet and dry cycles using the Gardner wear tester as described in the protocol. In between each cycle the slide was re-inoculated with bacteria. After the appropriate number of wear and re-inoculations (48 passes and 11 re-inoculations for healthcare formulation and 24 passes 5 re-inoculation for homecare formulation) the slide was exposed to bacteria for the indicated time frame (i.e. 5 minutes) followed by recovery in an appropriate neutralizing solution.

Examples

The following formulation in the example uses alcohol as the major carrier in order to provide fast drying property to the liquid formulations.

Table 1

Components	HE1 (wt %)	HE2 (wt %)	HE3 (wt%)
Water	balance	balance	balance
Ethanol	70	70	0
2-Propanol	0	0	70
Polyethyloxazoline	2	2	2
Quaternary ammonium compound	0.8	1.2	1.2
Wetting agent/Surfactant	0.1	0.1	0.1

The residual efficacy testing was conducted using EP01-1A protocol and the results are listed in the following Table.

Table 2

Formulation	EP01-1A (average log reduction bacterial)
HE1	3.53
HE2	5.50
HE3	4.50

These formulations show excellent residual efficacy result based on EP01-1A test.

The ASTM E 1153 test protocol was also followed to assess the initial biocidal property of HE2. Test results are presented in the following Table.

Table 3

Initial Efficacy	Time	Method	
Bacterial	3 log reduction	Complete kill (<10 CFU/PFU)	
<i>Klebsiella pneumoniae</i>	30 seconds	1 minute	ASTM E 1153
<i>Pseudomonas aeruginosa</i>	30 seconds	30 seconds	ASTM E 1153
<i>Staphylococcus aureus</i>	30 seconds	30 seconds	ASTM E 1153
<i>MRSA</i>	30 seconds	30 seconds	ASTM E 1153
<i>VRE</i>	30 seconds	30 seconds	ASTM E 1153
<i>Enterobacter aerogenes</i>	30 seconds	30 seconds	ASTM E 1153
<i>Enterococcus faecalis</i>	30 seconds	1 minute	ASTM E 1153
Fungal			
<i>Aspergillus niger</i>	1 minute	5 minutes	ASTM E 1153
<i>Tricophyton mentagrophytes</i>	1 minute	5 minutes	ASTM E 1153
Viral			
H1N1 (envelope)	30 seconds	30 seconds	ASTM E 1053
MS2 (Non-enveloped)	30 seconds	5 minutes	ASTM E 1053
Residual Efficacy	Time frame of exposure	Log reduction	Method
<i>Pseudomonas aeruginosa</i>	5 minutes	>3	EPA 01-1A
<i>Enterobacter aerogenes</i>	5 minutes	>3	EPA 01-1A
<i>Staphylococcus aureus</i>	5 minutes	>3	EPA 01-1A

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These data clearly demonstrate that sample surfaces treated with the exemplary liquid formulation disclosed herein possess a demonstrable biocidal activity at the indicated time frame.

It will therefore be readily understood by those persons skilled in the art that the present invention is susceptible of broad utility and application. Many embodiments and

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adaptations of the present invention other than those herein described, as well as many variations, modifications and equivalent arrangements, will be apparent from or reasonably suggested by the present invention and the foregoing description thereof, without departing from the substance or scope of the present invention. Accordingly,

5 while the present invention has been described herein in detail in relation to its preferred embodiment, it is to be understood that this disclosure is only illustrative and exemplary of the present invention and is made merely for purposes of providing a full and enabling disclosure of the invention. The foregoing disclosure is not intended or to be construed to limit the present invention or otherwise to exclude any such other embodiments,

10 adaptations, variations, modifications and equivalent arrangements.

What is claimed is:

1. An article comprising:
a surface having been treated with a composition comprising:
a polymeric film former, wherein the polymeric film former is an oxazoline
5 homopolymer or an extended or a modified polymer based on an oxazoline homopolymer,
and
a carrier.
2. The article according to claim 1, wherein the composition is in a form of a liquid.
10
3. The article according to claim 1, wherein the composition further comprises a
biocidal agent.
4. The article according to claim 3, the biocidal agent is a quaternary ammonium
15 compound or a combination of quaternary ammonium compounds.
5. The article according to claim 1, wherein the composition further comprises a
cleaning and soil resistant agent.
- 20 6. The article according to claim 1, wherein the carrier is selected from the group
consisting of water, low molecular weight alcohol, alkylene glycol ether, terpenes and
their derivatives, other solvent, and a combination thereof.
7. The article according to claim 1, wherein the carrier is present in an amount of at
25 least 70 weight percent.

8. The article according to claim 1, wherein the article is selected from the group consisting of a touch screen, screen protector, wipe, food wrap, plastic substrate, plastic coating, and a combination thereof.

5

9. A method of using a composition, the method comprising:

obtaining a composition comprised of a polymeric film former, wherein the polymeric film former is an oxazoline homopolymer or an extended or a modified polymer based on an oxazoline homopolymer, and a carrier; and

10 applying the composition to an article.

10. The method of using according to claim 9, wherein the composition is in a form of a liquid.

15 11. The method of using according to claim 9, wherein the composition further comprises a biocidal agent.

12. The method of using according to claim 11, wherein the biocidal agent is a quaternary ammonium compound or a combination of quaternary ammonium compounds.

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13. The method of using according to claim 9, wherein the composition further comprises a cleaning and soil resistant agent.

14. The method of using according to claim 9, wherein the carrier is selected from the group consisting of water, low molecular weight alcohol, alkylene glycol ether, terpenes and their derivatives, other solvent, and a combination thereof.

5 15. The method of using according to claim 9, wherein the carrier is present in an amount of at least 70 weight percent.

16. The method of using according to claim 9, wherein the article is selected from the group consisting of a touch screen, screen protector, wipe, food wrap, plastic substrate,
10 plastic coating, and a combination thereof.

17. A kit comprising:

a composition comprised of a carrier and a polymeric film former, wherein the polymeric film former is an oxazoline homopolymer or an extended or a modified polymer
15 based on an oxazoline homopolymer; and

an an application device.

18. The kit according to claim 17, wherein the application device is selected from the group consisting of a spray bottle, sponge, a wipe, and a combination thereof.

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19. The kit according to claim 17, wherein the composition further comprises a quaternary ammonium compound or a combination of quaternary ammonium compounds.

20. The kit according to claim 17, wherein the composition further comprises a
25 cleaning and soil resistant agent.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US17/33750

A. CLASSIFICATION OF SUBJECT MATTER
 IPC - A61K 31/522; A01N 25/02, 25/10, 33/02, 33/12 (2017.01)
 CPC - A61K 31/522; A01N 25/02, 25/04, 25/10, 33/02, 33/12

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

See Search History document

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

See Search History document

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

See Search History document

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X -- Y	US 4,481,167 A (GINTER, SP et al.) 6 November 1984; abstract; column 1, lines 50-65; column 6, lines 37-46; column 7, lines 1-45; column 9, lines 30-60; column 10, lines 44-54	1-3, 6-7, 9-11, 14-15 -- 3-5, 8, 11-13, 16-20
Y	WO 2007/070801 A2 (ALLAC-CEM, INC.) 21 June 2007; abstract; page 5, lines 30-40; page 25, lines 15-30; page 36, lines 5-10; page 55, lines 10-35; page 61, lines 5-10, 20-35	3-4, 8, 11-12, 16-20
Y	US 2,897,182 A (DE BENNEVILLE, PL et al.) 28 July 1959; column 11, lines 50-65	5, 13, 20
Y	US 6,110,387 A (CHOUDHURY, P et al.) 29 August 2000; abstract	3, 11
A	US 4,912,154 A (ARORA, KS et al.) 27 March 1990; entire document	1-20
A	WO 2008/127416 A2 (MASSACHUSETTS INSTITUTE OF TECHNOLOGY) 23 October 2008; entire document	1-20
A	CN 101473825 A (JIALONG SUN) 8 July 2009; entire document	1-20

Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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"E" earlier application or patent but published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
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"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search 20 July 2017 (20.07.2017)	Date of mailing of the international search report 14 AUG 2017
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