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(54) **METHOD FOR IMPROVING DYE STABILITY IN COLORED ACIDIC RINSE-AID FORMULATIONS**

VERFAHREN ZUR ERHÖHUNG DER FARBSTOFFSTABILITÄT IN GEFÄRBTEN SAUREN
KLARSPÜLMITTELN

PROCEDE SERVANT A AMELIORER LA STABILITE D'UN COLORANT DE FORMULATIONS
D'ADJUVANT DE RIN AGE ACIDE ET COLORE

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(73) Proprietor: **THE PROCTER & GAMBLE COMPANY**
Cincinnati, Ohio 45202 (US)

(72) Inventors:
• **FOLEY, Peter, Robert**
Cininatti, OH 45209 (US)
• **SONG, Brian, X.**
West Chester, OH 45069 (US)

• **PIERONI, Lucio**
Cincinnati, OH 45202 (US)

(74) Representative: **Brooks, Maxim Courtney et al**
Procter & Gamble Technical Centres Limited,
Whitley Road,
Longbenton
Newcastle upon Tyne NE12 9TS (GB)

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• **SCHICK, MARTIN : "NONIONIC
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NEW YORK**

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DescriptionTECHNICAL FIELD

5 **[0001]** The present invention relates to acidic rinse aid formulations used in automatic dishwashing. More particularly, the invention relates to a method for improving stability of a dye used in a colored aqueous acidic rinse aid composition to ambient and ultra-violet light radiation of a machine dishwasher rinse aid formulation.

BACKGROUND OF THE INVENTION

10 **[0002]** The use of rinse aids in commercial and institutional machine dishwashers and also in household automatic dishwashers, is well known. In automatic dishwashing, during the rinse cycle, a final rinse of fresh water serves to displace pre-final rinse water and its attendant detergent and soil residues. Rinse aid formulations are aqueous solutions containing a low foam nonionic surfactant. During the rinse cycle, the rinse aid is injected into the final fresh water
15 rinse at a concentration of about 100 to about 500 ppm. The surfactant in the rinse water lowers the surface tension of the rinse water and improves the wetting action of the rinse water.

[0003] The inventors of the present invention have previously discovered that it is possible to formulate an aqueous rinse aid containing a low foam nonionic surfactant and a low molecular weight neutralized, partially neutralized, or non-neutralized polyacrylate polymer without the use of a compatibilizing high molecular weight polymer or a high cloud point co-surfactant, thus providing an acidic, stable aqueous rinse aid dispersion which reduces spotting and
20 filming while improving sheeting action and drainage.

[0004] One major problem, however, associated with acidic, stable rinse aid dispersions is that when a dye is added in the composition to impart a pleasing color, especially a typical blue or green dye, such as a FD&C #1 Blue dye for example, there is a tendency for such rinse aid compositions to change color upon exposure to ambient and UV light,
25 for example sunlight. Frequently, if not inevitably, acidic rinse aid products will be subjected to UV light (e.g., sunlight) repeatedly and over extended periods of time prior to use by the consumer. The formulations of typical automatic dishwashing (ADW) rinse aid products are specifically tailored to a color that has been proven to be aesthetically pleasing to consumers. Any change in the target formulation color, especially when clear or translucent bottle or other packaging container is used, can negatively impact the consumer's perception of the quality of the product.

[0005] Typically, in the past, colored rinse aid compositions have been formulated at or near neutral pH, since it has been generally heretofore understood that non-acidic rinse aid compositions will not reduce the dye stability to ambient and ultraviolet light radiation. The inventors of the present invention have addressed this dye stability issue in the present invention and provided an effective solution to this problem. Before the present invention, there has been no method or composition known heretofore that addresses this dye stability problem in acidic rinse aid compositions.
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[0006] Accordingly, it has thus been desirable to have a colored acidic rinse aid product which is not significantly altered by UV light. Further, despite the above-mentioned disclosures in the art, it has been desirable to have an acidic rinse aid product which has superior performance in addition to a packaging appearance. It is extremely desirable that the packaging appearance (e.g., color) is maintained until the consumer has disposed of the product. The present invention overcomes the problems as set forth above.
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BACKGROUND ART

[0007] U.S. Patent No. 5,254,282, issued on Oct. 19, 1993 to Fusiak, discloses acidified paint removal formulations which are stabilized against color formation by inclusion of a chelating-agent; and
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U.S. Patent No. 5,206,006, issued on April 27, 1993 to Frontini et al. discloses a composition of the trinder type for detecting hydrogen peroxide evolved in a solution, which contains a chelating agent.

SUMMARY OF THE INVENTION

50 **[0008]** The invention meets the needs above by providing a method and composition for improving stability of a dye used in a colored aqueous acidic rinse aid composition to ambient and ultra-violet light radiation.

[0009] In one aspect of the present invention, the method for improving stability of a dye used in a colored aqueous acidic rinse aid composition to ambient and ultra-violet light radiation includes the step of adding a chelant selected from the group consisting of amino carboxylates, amino phosphonates, polyfunctionally-substituted aromatic chelants, and mixtures thereof to an acidic rinse aid composition. The chelant is present in an amount of at least 1 ppm by weight of the rinse aid composition.
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[0010] In another aspect of the present invention, a colored acidic rinse aid composition exhibiting improved stability of a dye used therein to ambient and ultra-violet light radiation includes a low foaming nonionic surfactant present in

a range of from about 2% to about 80% by weight of the rinse aid composition. The composition also includes a hydrotrope present in a range of from about 0.5% to about 20% by weight of the rinse aid composition. The composition further includes a polyacrylate polymer present in a range of from about 0.1% to about 15% by weight of the rinse aid composition. The polymer has a weight average molecular weight in a range of from about 1000 to about 50,000. The composition still further includes a chelant selected from the group consisting of amino carboxylates, amino phosphonates, polyfunctionally-substituted aromatic chelants, and mixtures thereof. The chelant is present in an amount of at least 1 ppm by weight of the rinse aid composition.

DETAILED DESCRIPTION OF THE INVENTION

[0011] In the preferred embodiment of the present invention, the method includes the step of adding a chelant selected from the group consisting of amino carboxylates, amino phosphonates, polyfunctionally-substituted aromatic chelants, and mixtures thereof to an acidic rinse aid composition. The chelant is present in an amount of at least 1 ppm, preferably at least 50 ppm, more preferably at least 100 ppm and most preferably at least 150 ppm by weight of the rinse aid composition.

Chelants

[0012] The acidic rinse aid composition includes chelants selected from the group consisting of amino carboxylates, amino phosphonates, polyfunctionally-substituted aromatic chelants, and mixtures thereof. Other useful chelants include iron and/or manganese chelants. Amino carboxylates useful as optional chelants include ethylenediaminetetraacetates, N-hydroxyethylethylenediaminetriacetates, nitrilotriacetates, ethylenediamine tetrapropionates, triethylene-tetra-amine-hexacetates, diethylene-triaminepentaacetates, and ethanoldiglycines, alkali metal, ammonium, and substituted ammonium salts therein and mixtures therein.

[0013] Polyfunctionally-substituted aromatic chelants include dihydroxydisulfobenzenes such as 1,2-dihydroxy-3,5-disulfobenzene, and ethylenediamine disuccinate ("EDDS").

[0014] In the preferred embodiment, the chelant is selected from the group consisting of ethylene diamine tetra acetic acid carboxylate, diethylene triamine penta acetic acid carboxylate, and mixtures thereof. In another preferred embodiment, the chelants are selected from polyphosphonate, ethylene diamine tetra acetic acid carboxylate (EDTA) and diethylene triamine penta acetic acid carboxylate (DTPA). Most preferably, the chelant is ethylene triamine penta phosphoric acid.

Dyes

[0015] In the preferred embodiment of the present invention, the dye is selected from the group consisting of Pontamine Blue (Copper Phthalocyanine Dyestuff), and Liquitant Patent Blue (polymeric colorant) and FD&C Blue #1 (having the molecular formula $C_{37}H_{34}N_2Na_2O_9S_3$), and mixtures thereof. Preferably, the dye has a color selected from the group consisting of blue, green, or mixtures thereof. It has been found that the blue and green dyes typically used in rinse aid formulations, such as the FD&C #1, are unstable to UV light in acidic conditions unless the rinse aid composition is formulated with a chelant according to the present invention. It has been found that even the so-called "acid stable" dyes, such as Acid Blue 80, fade quickly when used in acidic rinse aids, i.e., the dye is colorless after an exposure of 1 to 2 days under UV light radiation, unless the rinse aid is formulated according to the present invention, with a chelant. Desirably, the dyes are added in an amount in a range of from 0.25% of a 1% solution in water to about 0.5% of a 1% solution in water.

Dye-Chelant Systems

[0016] In the preferred embodiment the dye is selected from the group consisting of copper phthalocyanine dyestuff, FD&C #1 colors, polymeric colorants, and mixtures thereof, and the chelant is selected from the group consisting of amino carboxylates, amino phosphonates, polyfunctionally-substituted aromatic chelants, and mixtures thereof. It has been found that these "dye-chelant" systems, optimally yield the most effective dye color stability to ambient and UV light radiation, when such dyes are used in acidic rinse aids.

[0017] The most preferred chelants are selected from polyphosphonate ethylene diamine tetra acetic acid carboxylate (EDTA) and diethylene triamine penta acetic acid carboxylate (DTPA) in combination with the most preferred dyes selected from blue dyes such as Pontamine Blue (Copper Phthalocyanine Dyestuff), and Liquitant Patent Blue (polymeric colorant) and FD&C Blue #1 (for example, an FD&C Blue #1 having the molecular formula $C_{37}H_{34}N_2Na_2O_9S_3$). These dye-chelant systems discovered by the inventors, when formulated in an acidic rinse aid dispersion, all form ambient and UV light resistant, color stable systems.

[0018] The especially preferred Dye-Chelant systems discovered by the inventors are listed below:

Liquitant Patent Blue in combination with DTPA;
FD&C #1 in combination with DTPA; and
Liquitant Patent Blue in combination with DTPA.

Carboxylic acids

[0019] In the preferred embodiment, the dye stability is further enhanced in the presence of carboxylic acids. Especially preferred are polycarboxylic acids, such as Sokolan PA13PN manufactured by BASF, Acusol 480, and Acusol 480N, both manufactured by Rohm & Haas. These polycarboxylic acids provide multiple benefits, as they help stabilize the dye and deliver filming performance benefits in the rinse cycle of automatic dishwashing. Desirably, the polycarboxylic acid is present in an amount in a range of from 0.25% to 10% by weight of the acidic rinse aid composition, and preferably, in an amount in a range of from 1% to 6% by weight.

[0020] In another embodiment of the present invention, the colored acidic rinse aid composition exhibiting improved stability of a dye used therein to ambient and ultra-violet light radiation includes a low foaming nonionic surfactant desirably present in a range of from 2% to 80% by weight of the rinse aid composition. The composition also includes a hydrotrope desirably present in a range of from 0.5% to 20% by weight of the rinse aid composition. The composition further includes a polyacrylate polymer desirably present in a range of from 0.1% to 15% by weight of the rinse aid composition. The polymer has a weight average molecular weight in a range of from 1000 to 50,000. The composition still further includes a chelant. The chelant is present in an amount of at least 1 ppm by weight of the rinse aid composition.

Low Foaming Nonionic Surfactant

[0021] The nonionic surfactants useful in the acidic rinse aid dispersion may be any known low foaming nonionic surfactant used in machine dishwashing applications. Typical suitable nonionic surfactants include the following commercially available materials: Triton R™ CF-10 (an alkylaryl polyether) and Triton DF-16 (a modified polyalkoxylated alcohol) manufactured by Rohm and Haas Company; Plurafac LF404™, Plurafac LF400™ (mixed linear alcohol alkoxylates), and Pluronic R™ L-62 (a polyoxyethylene-polyoxypropylene block copolymer), all manufactured by BASF Wyandotte Corporation, and BASF RA30 surfactant, manufactured by BASF Corporation. The rinse aid formulation of the invention may contain one or a mixture of such low foaming nonionic surfactants.

[0022] Essentially any nonionic surfactants' useful for deterative purposes can be included in the rinse aid compositions. Exemplary, non-limiting classes of useful nonionic surfactants are listed below.

(i) Nonionic polyhydroxy fatty acid amide surfactant

[0023] Polyhydroxy fatty acid amides suitable for use herein are those having the structural formula R_2CONR_1Z wherein: R_1 is H, C_1 - C_4 hydrocarbyl, 2-hydroxy ethyl, 2-hydroxy propyl, or a mixture thereof, preferable C_1 - C_4 alkyl, more preferably C_1 or C_2 alkyl, most preferably C_1 alkyl (i.e., methyl); and R_2 is a C_5 - C_{31} hydrocarbyl, preferably straight-chain C_5 - C_{19} alkyl or alkenyl, more preferably straight-chain C_9 - C_{17} alkyl or alkenyl, most preferably straight-chain C_{11} - C_{17} alkyl or alkenyl, or mixture thereof; and Z is a polyhydroxyhydrocarbyl having a linear hydrocarbyl chain with at least 3 hydroxyls directly connected to the chain, or an alkoxylated derivative (preferably ethoxylated or propoxylated) thereof. Z preferably will be derived from a reducing sugar in a reductive amination reaction; more preferably Z is a glycityl.

(ii) Nonionic condensates of alkyl phenols

[0024] The polyethylene, polypropylene, and polybutylene oxide condensates of alkyl phenols are suitable for use herein. In general, the polyethylene oxide condensates are preferred. These compounds include the condensation products of alkyl phenols having an alkyl group containing from 6 to 18 carbon atoms in either a straight chain or branched chain configuration with the alkylene oxide.

(iii) Nonionic ethoxylated alcohol surfactant

[0025] The alkyl ethoxylate condensation products of aliphatic alcohols with from about 1 to about 25 moles of ethylene oxide are suitable for use herein. The alkyl chain of the aliphatic alcohol can either be straight or branched, primary or secondary, and generally contains from 6 to 22 carbon atoms. Particularly preferred are the condensation

products of alcohols having an alkyl group containing from 8 to 20 carbon atoms with from 2 to 10 moles of ethylene oxide per mole of alcohol.

(iv) Nonionic ethoxylated/propoxylated fatty alcohol surfactant

[0026] The ethoxylated C₆-C₁₈ fatty alcohols and C₆-C₁₈ mixed ethoxylated/propoxylated fatty alcohols are highly preferred surfactants for use herein, particularly where water soluble. Preferably the ethoxylated fatty alcohols are the C₁₀-C₁₈ ethoxylated fatty alcohols with a degree of ethoxylation of from 3 to 50, most preferably these are the C₁₂-C₁₈ ethoxylated fatty alcohols with a degree of ethoxylation from 3 to 40. Preferably the mixed ethoxylated/propoxylated fatty alcohols have an alkyl chain length of from 10 to 18 carbon atoms, a degree of ethoxylation of from 3 to 30 and a degree of propoxylation of from 1 to 10.

(v) Nonionic EO/PO condensates with propylene glycol

[0027] The condensation products of ethylene oxide with a hydrophobic base formed by the condensation of propylene oxide with propylene glycol are suitable for use herein. The hydrophobic portion of these compounds preferably has a molecular weight of from 1500 to 1800 and exhibits water insolubility. Examples of compounds of this type include certain of the commercially-available Pluronic™ surfactants, marketed by BASF.

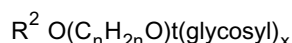
(vi) Nonionic EO condensation products with propylene oxide/ethylene diamine adducts

[0028] The condensation products of ethylene oxide with the product resulting from the reaction of propylene oxide and ethylenediamine are suitable for use herein. The hydrophobic moiety of these products consists of the reaction product of ethylenediamine and excess propylene oxide, and generally has a molecular weight of from 2500 to 3000. Examples of this type of nonionic surfactant include certain of the commercially available Tetronic™ compounds, marketed by BASF.

(vii) Nonionic alkylpolysaccharide surfactant

[0029] Suitable alkylpolysaccharides for use herein are disclosed in U.S. Pat. No. 4,565,647, Llenado, issued Jan. 21, 1986, having a hydrophobic group containing from 6 to 30 carbon atoms, preferably from 10 to 16 carbon atoms and a polysaccharide, e.g., a polyglycoside, hydrophilic group containing from 1.3 to 10, preferably from 1.3 to 3, most preferably from 1.3 to 2.7 saccharide units. Any reducing saccharide containing 5 or 6 carbon atoms can be used, e.g., glucose, galactose and galactosyl moieties can be substituted for the glucosyl moieties. (Optionally the hydrophobic group is attached at the 2-, 3-, 4-, etc. positions thus giving a glucose or galactose as opposed to a glucoside or galactoside.) The intersaccharide bonds can be, e.g., between the one position of the additional saccharide units and the 2-, 3-, 4-, and/or 6- positions on the preceding saccharide units.

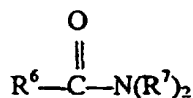
[0030] The preferred alkylpolyglycosides have the formula:



wherein R² is selected from the group consisting of alkyl, alkylphenyl, hydroxyalkyl, hydroxyalkylphenyl, and mixtures thereof in which the alkyl groups contain from 10 to 18, preferably from 12 to 14, carbon atoms; n is 2 or 3, preferably from 1.3 to 3, most preferably from 1.3 to 2.7. The glycosyl is preferably derived from glucose.

(viii) Nonionic fatty acid amide surfactant

[0031] Fatty acid amide surfactants suitable for use herein are those having the formula:



wherein R⁶ is an alkyl group containing from 7 to 21, preferably from 9 to 17 carbon atoms and each R⁷ is selected from the group consisting of hydrogen, C₁-C₄ alkyl, C₁-C₄ hydroxyalkyl, and -(C₂ H₄ O)_x H, where x is in the range of

from 1 to 3.

[0032] The hydrotrope useful in the present rinse aid dispersion is selected from the group consisting of sodium cumene sulfonate, sodium xylene sulfonate, sodium toluenesulfonate, dioctyl sodium sulfosuccinate, alkyl naphthalene sulfonate and dihexyl sodium sulfosuccinate. Preferably, the hydrotrope is sodium cumene sulfonate.

Polyacrylate polymer

[0033] The low molecular weight polyacrylate polymer useful in the present acidic rinse aid dispersion is selected from the group consisting of polyacrylate homopolymer, polyacrylate copolymer, polyacrylate terpolymer, and mixtures thereof. Preferably, the polyacrylate polymer is a partially neutralized or non-neutralized polyacrylate polymer having a pH in a range of from 2.0 to 4.0. In the preferred embodiment, the polyacrylate polymer is desirably at least 75% non-neutralized. Alternatively, the polyacrylate polymer is non-neutralized. Still alternatively, a neutralized polyacrylate polymer, which is more than 25% neutralized may also be used, but an acid must be added in an amount sufficient to bring the pH value of the resultant rinse aid dispersion within a range of from 2 to 6.

[0034] Typically the polyacrylate polymer is a homopolymer of acrylic or methacrylic acid, or a copolymer formed from at least 50 weight percent acrylic acid. The polyacrylate polymer is a low molecular weight polymer or its alkali metal or ammonium salt, having a weight average molecular weight ranging from 1,000 to 50,000.

Acid

[0035] When adding an acid to the rinse aid dispersion, any acid may be used although organic acids are more desirable and the preferred organic acid is citric acid.

[0036] It has been discovered that even in the absence of a compatibilizing high molecular weight polymer, but with the addition of a hydrotrope and in acidic conditions, the stable rinse aid dispersion does not phase separate upon storage for a reasonable time or under actual use conditions and permits each component to effectively reduce spotting and filming and improve sheeting action without interfering with the foaming and defoaming action of the surfactant.

[0037] In the preferred embodiment, the rinse aid formulation of the invention is an aqueous dispersion at a pH of from 2 to 6. Preferably, the pH ranges from 3 to 5.

[0038] The rinse aid dispersion of the present invention is preferably prepared by stirring the desired amount of the surfactant into an aqueous solution of the low molecular weight non-neutralized or only partially neutralized (no more than 25% neutralized) polyacrylic acid followed by the gradual addition of the hydrotrope and if necessary, citric acid to bring the dispersion to within the desired pH range.

[0039] The formulation may also contain other additives including sequestants such as NTA, EDTA, or sodium citrate and water miscible solvents such as ethanol, isopropanol and propylene glycol. Ethanol is the preferred solvent, present desirably in a range of from 0.1% to 10% by weight of the rinse aid dispersion and preferably in a range of from 2% to 8% by weight.

Hydrotrope

[0040] In the preferred embodiment, the hydrotrope is selected from the group consisting of sodium cumene sulfonate, sodium xylene sulfonate, sodium toluenesulfonate, dioctyl sodium sulfosuccinate, alkyl naphthalene sulfonate and dihexyl sodium sulfosuccinate. Preferably, the hydrotrope is sodium cumene sulfonate, and the partially-neutralized polyacrylate polymer is a polyacrylate copolymer which is about 20% neutralized. In the preferred embodiment, the stable aqueous rinse aid dispersion is essentially free of a high molecular weight compatibilizing polymer and is also free of an additional nonionic surfactant having a cloud point of at least 70 degrees C.

[0041] The following examples are intended to illustrate the invention.

EXAMPLE 1

[0042] A rinse aid dispersion according to the present invention is made as follows, from the following composition, by weight percent:

Low foam nonionic surfactant	20.0%
Sodium cumene sulfonate hydrotrope	7.0%
Polyacrylate copolymer (20% neutralized)	4.5%
Ethanol	6.0%
Chelant (DTPA)	0.015%

(continued)

Liquitant Patent Blue	0.25%
Water	62.235%

This dye has ambient and UV light radiation stability and its color is unchanged after exposure to UV radiation for 1-2 days.

EXAMPLE 2

[0043] Another rinse aid dispersion according to the present invention is made as follows, from the following composition, by weight percent:

Low foam nonionic surfactant	35.0%
Sodium cumene sulfonate hydrotrope	4.0%
Polyacrylate copolymer (20% neutralized)	4.0%
Citric acid	1.6%
Ethanol	6.0%
Chelant (polyphosphonate)	0.015%
FD&C Blue #1	0.25%
Water	49.135%

This dye has ambient and UV light radiation stability and its color is unchanged after exposure to UV radiation for 1-2 days.

EXAMPLE 3

[0044] Another rinse aid dispersion according to the present invention is made as follows, from the following composition, by weight percent:

Low foam nonionic surfactant	12.0%
Sodium cumene sulfonate hydrotrope	4.0%
Polyacrylate copolymer (20% neutralized)	1.0%
Citric acid	1.6%
Ethanol	5.0%
Chelant (polyphosphonate)	0.015%
FD&C Blue #1	0.25%
Water	76.135%

This dye has ambient and UV light radiation stability and its color is unchanged after exposure to UV radiation for 1-2 days.

[0045] The above rinse aid formulations are prepared by the following procedure: appropriate amounts of: a low foaming nonionic surfactant (Plurafac LF 404™), a 45% active sodium cumene sulfonate hydrotrope (45% active SCS), a 50% active polyacrylate copolymer (Acusol 480™, made by Rohm & Haas), a 50% active citric acid, ethanol and deionized water are added to a beaker and stirred sequentially. Also added is a chelant in an amount of 0.015 weight% (150 ppm by weight) and a color dye as indicated. The resulting mixture obtained is a blue or green colored, single-phase acidic dispersion and is stable, with no phase separation occurring after the resulting mixture is kept undisturbed in a closed container for a period of 2 weeks at a temperature of 50 degrees C. Further, the resulting mixture exhibits no more than about a 30% dye discoloration when exposed to ambient light for 6 months, and no more than about a 30% dye discoloration when exposed to UV light for 2 days.

Claims

1. A method for improving stability of a dye used in a colored aqueous acidic rinse aid composition to ambient and

ultra-violet light radiation, **characterized by** the steps of:

adding a chelant selected from the group consisting of amino carboxylates, amino phosphonates, polyfunctionally-substituted aromatic chelants, and mixtures thereof to an acidic rinse aid composition, said chelant being present in an amount of at least 1 ppm by weight of said rinse aid composition.

2. A method according to claim 1, wherein said chelant is present in an amount of at least 50 ppm by weight of said rinse aid composition.

3. A method according to claims 1-2, wherein said chelant is selected from the group consisting of ethylene diamine tetra acetic acid carboxylate, diethylene triamine penta acetic acid carboxylate, and mixtures thereof.

4. A method according to claims 1-3, wherein said chelant is ethylene triamine penta phosphoric acid.

5. A method according to claims 1-4, wherein said dye is selected from the group consisting of copper phthalocyanine dyestuff, FD&C #1 color, polymeric colorants, and mixtures thereof.

6. A colored acidic rinse aid composition exhibiting improved stability of a dye used therein to ambient and ultra-violet light radiation, **characterized by**:

a low foaming nonionic surfactant present in a range of from 2% to 80% by weight of said rinse aid composition; a hydrotrope present in a range of from 0.5% to 20% by weight of said rinse aid composition; a polyacrylate polymer present in a range of from 0.1% to 15% by weight of said rinse aid composition, said polymer having a weight average molecular weight in a range of from 1000 to 50,000; and a chelant selected from the group consisting of amino carboxylates, amino phosphonates, polyfunctionally-substituted aromatic chelants, and mixtures thereof said chelant being present in an amount of at least 1 ppm by weight of said rinse aid composition.

7. An acidic rinse aid composition according to claim 6 wherein said rinse aid has a pH in a range of from 2 to 7.

8. An acidic rinse aid composition according to claim 6 including an acid present in a range of from 0.1% to 20% by weight of said rinse aid composition.

Patentansprüche

1. Verfahren zur Verbesserung der Stabilität eines in einer gefärbten, wässrigen, sauren Klarspülzusammensetzung verwendeten Farbstoffs gegenüber Umgebungs- und Ultraviolettlichtstrahlung, **gekennzeichnet durch** die Schritte:

Zugeben eines Komplexbildners, gewählt aus der Gruppe, bestehend aus Aminocarboxylaten, Aminophosphonaten, polyfunktionell substituierten aromatischen Komplexbildnern und Mischungen hiervon, zu einer sauren Klarspülzusammensetzung, wobei der Komplexbildner in einer Menge von mindestens 1 Gew.-ppm der Klarspülzusammensetzung vorliegt.

2. Verfahren nach Anspruch 1, wobei der Komplexbildner in einer Menge von mindestens 50 Gew.-ppm der Klarspülzusammensetzung vorliegt.

3. Verfahren nach den Ansprüchen 1-2, wobei der Komplexbildner aus der Gruppe gewählt ist, bestehend aus Ethylendiamintetraessigsäurecarboxylat, Diethylentriaminpentaessigsäurecarboxylat und Mischungen hiervon.

4. Verfahren nach den Ansprüchen 1-3, wobei der Komplexbildner Ethylentriaminpentaphosphorsäure ist.

5. Verfahren nach den Ansprüchen 1-4, wobei der Farbstoff aus der Gruppe gewählt ist, bestehend aus Kupferphthalocyanin-Farbstoff, FD&C#1 Farbe, polymeren Färbemitteln und Mischungen hiervon.

6. Gefärbte, saure Klarspülzusammensetzung, welche eine verbesserte Stabilität eines darin verwendeten Farbstoffs gegenüber Umgebungs- und Ultraviolettlichtstrahlung zeigt, **gekennzeichnet durch**

ein gering schäumendes, nichtionisches Tensid, das im Bereich von 2 bis 80 Gew.-% der Klarspülzusammensetzung vorliegt;

ein Hydrotrop, das im Bereich von 0,5 bis 20 Gew.-% der Klarspülzusammensetzung vorliegt;

ein Polyacrylatpolymer, das im Bereich von 0,1 bis 15 Gew.-% der Klarspülzusammensetzung vorliegt, wobei das Polymer ein gewichtsmittleres Molekulargewicht im Bereich von 1.000 bis 50.000 aufweist; und

einen Komplexbildner, gewählt aus der Gruppe, bestehend aus Aminocarboxylaten, Aminophosphonaten, polyfunktionell substituierten aromatischen Komplexbildnern und Mischungen hiervon, wobei der Komplexbildner in einer Menge von mindestens 1 Gew.-ppm der Klarspülzusammensetzung vorliegt.

7. Saure Klarspülzusammensetzung nach Anspruch 6, wobei der Klarspüler einen pH im Bereich von 2 bis 7 aufweist.

8. Saure Klarspülzusammensetzung nach Anspruch 6, einschließlich einer Säure, die im Bereich von 0.1 bis 20 Gew.-% der Klarspülzusammensetzung vorliegt.

Revendications

1. Procédé pour améliorer la stabilité d'un colorant utilisé dans une composition d'adjuvant de rinçage acide aqueuse colorée, vis-à-vis du rayonnement de la lumière ambiante et ultraviolette, **caractérisé par** les étapes consistant à:

ajouter un agent chélatant choisi dans le groupe constitué par les aminocarboxylates, les aminophosphonates, les agents chélatants aromatiques à substitution polyfonctionnelle, et leurs mélanges, à une composition d'adjuvant de rinçage acide, ledit agent chélatant étant présent en une quantité d'au moins 1 ppm en poids de ladite composition d'adjuvant de rinçage.

2. Procédé selon la revendication 1, dans lequel ledit agent chélatant est présent en une quantité d'au moins 50 ppm en poids de ladite composition d'adjuvant de rinçage.

3. Procédé selon les revendications 1-2, dans lequel ledit agent chélatant est choisi dans le groupe constitué par un carboxylate d'acide éthylènediaminetétraacétique, un carboxylate d'acide diéthylènetriaminepentaacétique, et leurs mélanges.

4. Procédé selon les revendications 1-3, dans lequel ledit agent chélatant est l'acide éthylènetriaminepentaphosphorique.

5. Procédé selon les revendications 1-4, dans lequel ledit colorant est choisi dans le groupe constitué par la matière colorante phtalocyanine de cuivre, le colorant FD&C n° 1, les colorants polymères et leurs mélanges.

6. Composition d'adjuvant de rinçage acide colorée présentant une meilleure stabilité d'un colorant utilisé dans ladite composition, vis-à-vis du rayonnement de la lumière ambiante et ultraviolette, **caractérisée par**:

un tensioactif non ionique faiblement moussant, présent dans un intervalle de 2% à 80% en poids de ladite composition d'adjuvant de rinçage;

un hydrotrope présent dans un intervalle de 0,5% à 20% en poids de ladite composition d'adjuvant de rinçage;

un polymère de type polyacrylate présent dans un intervalle de 0,1% à 15% en poids de ladite composition d'adjuvant de rinçage, ledit polymère ayant une masse moléculaire moyenne en poids comprise dans un intervalle de 1000 à 50 000; et

un agent chélatant choisi dans le groupe constitué par les aminocarboxylates, les aminophosphonates, les agents chélatants aromatiques à substitution polyfonctionnelle, et leurs mélanges, ledit agent chélatant étant présent en une quantité d'au moins 1 ppm en poids de ladite composition d'adjuvant de rinçage.

7. Composition d'adjuvant de rinçage acide selon la revendication 6, dans laquelle ledit adjuvant de rinçage a un pH compris dans un intervalle de 2 à 7.

8. Composition d'adjuvant de rinçage acide selon la revendication 6, incluant un acide présent dans un intervalle de 0,1% à 20% en poids de ladite composition d'adjuvant de rinçage.