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(54) CLEANING COMPOSITIONS INCLUDING ENZYMES AND ALKOXYLATED PHENOL

(57) Cleaning compositions that include a galactanase enzyme and alkoxylated phenol, in particular an alkoxylated polyaryl phenol and/or an alkoxylated polyalkyl phenol. Methods of making and using such cleaning compositions. Use of a galactanase enzyme and an alkoxylated phenol.

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Description

REFERENCE TO A SEQUENCE LISTING

⁵ **[0001]** This application contains a Sequence Listing in computer readable form, which is incorporated herein by reference.

FIELD OF THE INVENTION

¹⁰ **[0002]** The present invention relates to cleaning compositions that include a galactanase enzyme and alkoxylated phenol, in particular an alkoxylated polyaryl phenol and/or an alkoxylated polyalkyl phenol. The present invention also relates to methods of making and using such cleaning compositions. The present invention also relates to the use of a galactanase and an alkoxylated phenol.

15 BACKGROUND OF THE INVENTION

[0003] The laundry detergent formulator is constantly aiming to improve the performance of detergent compositions. Many detergent components may provide stain removal benefits, but some stains are particularly resistant to removal. Even when a soil is lifted from a fabric, the lifted material may redeposit onto the fabric during the wash cycle, impacting

whiteness and/or color contrast between white and colored regions on the same fabric.
 [0004] There is a need for improved cleaning compositions that provide whiteness and/or color contrast benefits.

SUMMARY OF THE INVENTION

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²⁵ **[0005]** The present invention relates to a cleaning composition that includes: an endo-beta-1,6-galactanase enzyme; and an alkoxylated phenol compound that is selected from the group consisting of an alkoxylated polyaryl phenol compound, and mixtures thereof.

[0006] The present invention also relates to a method of cleaning a surface, preferably a textile, where the method includes mixing the cleaning composition as described herein with water to form an aqueous liquor and contacting a surface, preferably a textile, with the aqueous liquor in a laundering step.

[0007] The present invention also relates to the use of an endo-beta-1,6-galactanase enzyme and an alkoxylated polyaryl phenol and/or an alkoxylated polyalkyl phenol in a cleaning composition to enhance the stain-removal, whiteness, and/or for malodor reduction.

35 DETAILED DESCRIPTION OF THE INVENTION

[0008] The present invention relates to cleaning compositions that include a galactanase enzyme and an alkoxylated (polyaryl/polyalkyl) phenol. Without wishing to be bound by theory, it is believed that the galactanase enzymes are active on soils including soils of microbial origin such as extracellular DNA materials that adhere to, and cause other soils to

- 40 adhere to, fabric. It is further believed that the alkoxylated phenols described herein work synergistically with the galactanase enzymes to assist in the removal of extracellular DNA-bound soils and prevent redeposition of the reaction products arising from the action of the galactanase enzymes. In doing so, it is believed that the activity of the enzyme on the textile surface is enhanced by reduced interference or inhibition from redeposited soils. [0009] The components of the compositions and processes of the present invention are described in more detail below.
- [0009] The components of the compositions and processes of the present invention are described in more detail below.
 [0010] As used herein, the articles "a" and "an" when used in a claim, are understood to mean one or more of what is claimed or described. As used herein, the terms "include," "includes," and "including" are meant to be non-limiting. The compositions of the present invention can comprise, consist essentially of, or consist of, the components of the present invention.
- [0011] The terms "substantially free of" or "substantially free from" may be used herein. This means that the indicated material is at the very minimum not deliberately added to the composition to form part of it, or, preferably, is not present at analytically detectable levels. It is meant to include compositions whereby the indicated material is present only as an impurity in one of the other materials deliberately included. The indicated material may be present, if at all, at a level of less than 1%, or less than 0.1 %, or less than 0.01 %, or even 0%, by weight of the composition.
- [0012] Unless otherwise noted, all component or composition levels are in reference to the active portion of that ⁵⁵ component or composition, and are exclusive of impurities, for example, residual solvents or by-products, which may be present in commercially available sources of such components or compositions.

[0013] All temperatures herein are in degrees Celsius (°C) unless otherwise indicated. Unless otherwise specified, all measurements herein are conducted at 20°C and under the atmospheric pressure.

[0014] In all embodiments of the present invention, all percentages are by weight of the total composition, unless specifically stated otherwise. All ratios are weight ratios, unless specifically stated otherwise.

[0015] It should be understood that every maximum numerical limitation given throughout this specification includes every lower numerical limitation, as if such lower numerical limitations were expressly written herein. Every minimum

⁵ numerical limitation given throughout this specification will include every higher numerical limitation, as if such higher numerical limitations were expressly written herein. Every numerical range given throughout this specification will include every narrower numerical range that falls within such broader numerical range, as if such narrower numerical ranges were all expressly written herein.

[0016] As used herein, the term "alkoxy" is intended to include C1-C8 alkoxy and C1-C8 alkoxy derivatives of polyols having repeating units such as butylene oxide, glycidol oxide, ethylene oxide or propylene oxide.

[0017] As used herein, unless otherwise specified, the terms "alkyl" and "alkyl capped" are intended to include C1-C18 alkyl groups, or even C1-C6 alkyl groups.

[0018] As used herein, unless otherwise specified, the term "aryl" is intended to include C3-12 aryl groups.

[0019] As used herein, unless otherwise specified, the term "arylalkyl" and "alkaryl" are equivalent and are each intended to include groups comprising an alkyl moiety bound to an aromatic moiety, typically having C1-C18 alkyl groups and, in one aspect, C1-C6 alkyl groups.

[0020] The terms "ethylene oxide," "propylene oxide" and "butylene oxide" may be shown herein by their typical designation of "EO," "PO" and "BO," respectively.

 $[0021] \quad As used herein, the term "cleaning and/or treatment composition" includes, unless otherwise indicated, granular,$

- 20 powder, liquid, gel, paste, unit dose, bar form and/or flake type washing agents and/or fabric treatment compositions. [0022] As used herein, "cellulosic substrates" are intended to include any substrate which comprises cellulose, either 100% by weight cellulose or at least 20% by weight, or at least 30% by weight or at least 40% or at least 50% by weight or even at least 60% by weight cellulose. Cellulose may be found in wood, cotton, linen, jute, and hemp. Cellulosic substrates may be in the form of powders, fibers, pulp and articles formed from powders, fibers and pulp. Cellulosic
- ²⁵ fibers, include, without limitation, cotton, rayon (regenerated cellulose), acetate (cellulose acetate), triacetate (cellulose triacetate), and mixtures thereof. Typically cellulosic substrates comprise cotton. Articles formed from cellulosic fibers include textile articles such as fabrics. Articles formed from pulp include paper.

[0023] As used herein, the term "maximum extinction coefficient" is intended to describe the molar extinction coefficient at the wavelength of maximum absorption (also referred to herein as the maximum wavelength), in the range of 400 nanometers to 750 nanometers.

[0024] As used herein "average molecular weight" is reported as a weight average molecular weight, as determined by its molecular weight distribution; as a consequence of their manufacturing process, polymers disclosed herein may contain a distribution of repeating units in their polymeric moiety.

[0025] As used herein the term "variant" refers to a polypeptide that contains an amino acid sequence that differs from a wild type or reference sequence. A variant polypeptide can differ from the wild type or reference sequence due to a deletion, insertion, or substitution of a nucleotide(s) relative to said reference or wild type nucleotide sequence. The reference or wild type sequence can be a full-length native polypeptide sequence or any other fragment of a full- length polypeptide sequence. A polypeptide variant generally has at least about 70% amino acid sequence identity with the reference sequence, but may include 75% amino acid sequence identity within the reference sequence, 80% amino

- 40 acid sequence identity within the reference sequence, 85% amino acid sequence identity with the reference sequence, 86% amino acid sequence identity with the reference sequence, 87% amino acid sequence identity with the reference sequence, 88% amino acid sequence identity with the reference sequence, 89% amino acid sequence identity with the reference sequence, 90% amino acid sequence identity with the reference sequence, 91% amino acid sequence identity with the reference sequence, 92% amino acid sequence identity with the reference sequence, 93% amino acid sequence
- ⁴⁵ identity with the reference sequence, 94% amino acid sequence identity with the reference sequence, 95% amino acid sequence identity with the reference sequence, 96% amino acid sequence identity with the reference sequence, 97% amino acid sequence identity with the reference sequence, 98% amino acid sequence identity with the reference sequence, 98% amino acid sequence identity with the reference sequence, 98% amino acid sequence identity with the reference sequence, 98% amino acid sequence identity with the reference sequence, 98% amino acid sequence identity with the reference sequence or 99% amino acid sequence identity with the reference sequence.
- ⁵⁰ **[0026]** As used herein, the term "solid" includes granular, powder, bar and tablet product forms.
 - [0027] As used herein, the term "fluid" includes liquid, gel, paste, and gas product forms.

Cleaning Composition

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⁵⁵ **[0028]** The present disclosure relates to cleaning and/or treatment compositions. The cleaning composition may be selected from the group of light duty liquid detergents compositions, heavy duty liquid detergent compositions, solid, for example particulate/powder or "dry" cleaning compositions, hard surface cleaning compositions, detergent gels commonly used for laundry, bleaching compositions, laundry additives, fabric enhancer compositions, shampoos, body

washes, other personal care compositions, and mixtures thereof. The cleaning composition may be a hard surface cleaning composition (such as a dishwashing composition) or a laundry composition (such as a heavy duty liquid or solid detergent composition).

[0029] The cleaning compositions may be in any suitable form. The composition can be selected from a liquid, solid,

- ⁵ or combination thereof. As used herein, "liquid" includes free-flowing liquids, as well as pastes, gels, foams and mousses. Non-limiting examples of liquids include light duty and heavy duty liquid detergent compositions, fabric enhancers, detergent gels commonly used for laundry, bleach and laundry additives. Gases, e.g., suspended bubbles, or solids, e.g. particles, may be included within the liquids. A "solid" as used herein includes, but is not limited to, powders, agglomerates, and mixtures thereof. Non-limiting examples of solids include: granules, micro-capsules, beads, noodles,
- and pearlised balls. Solid compositions may provide a technical benefit including, but not limited to, through-the-wash benefits, pre-treatment benefits, and/or aesthetic effects.
 [0030] The cleaning composition may be in the form of a unitized dose article, such as a tablet or in the form of a pouch. Such pouches typically include a water-soluble film, such as a polyvinyl alcohol water-soluble film, that at least partially encapsulates a composition. Suitable films are available from MonoSol, LLC (Indiana, USA). The composition
- 15 can be encapsulated in a single or multi-compartment pouch. A multi-compartment pouch may have at least two, at least three, or at least four compartments. A multi-compartmented pouch may include compartments that are side-by-side and/or superposed. The composition contained in the pouch may be liquid, solid (such as powders), or combinations thereof.
- 20 Galactanase Enzyme

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[0031] The endo-beta-1,6-galactanase enzyme is an extracellular polymer-degrading enzyme. The term "endo-beta-1,6-galactanase" or "a polypeptide having endo-beta-1,6-galactanase activity" means a endo-beta-1,6-galactanase activity (EC 3.2.1.164) that catalyzes the hydrolytic cleavage of 1,6-3-D-galactooligosaccharides with a degree of polym-

²⁵ erization (DP) higher than 3, and their acidic derivatives with 4-O-methylglucosyluronate or glucosyluronate groups at the non-reducing terminals.

[0032] For purposes of the present disclosure, endo-beta-1,6-galactanase activity is determined according to the procedure described in WO 2015185689 in Assay I. Suitable examples from class EC 3.2.1.164 are described in WO 2015185689, such as the mature polypeptide SEQ ID NO: 2 described therein. Preferably the galactanase enzyme is selected from Glycoside Hydrolase (GH) Family 30.

[0033] Preferably, the endo-beta-1,6-galactanase comprises a microbial enzyme. The endo-beta-1,6-galactanase may be fungal or bacterial in origin. Bacterial endo-beta-1,6-galactanase may be most preferred. Fungal endo-beta-1,6-galactanase may be most preferred.

[0034] A bacterial endo-beta-1,6-galactanase is obtainable from *Streptomyces*, for example *Streptomyces davawensis*. A preferred endo-beta-1,6-galactanase is obtainable from *Streptomyces davawensis* JCM 4913 defined in SEQ ID NO: 1 herein, or a variant thereof, for example having at least 40% or 50% or 60% or 70% or 75% or 80% or 85% or

90% or 95%, 96%, 97%, 98%, 99% or 100% identity thereto.
[0035] Other bacterial endo-beta-1,6-galactanase include those encoded by the DNA sequences of *Streptomyces avermitilis* MA-4680 with amino acid sequence defined in SEQ ID NO: 2 herein, or a variant thereof, for example having

- 40 at least 40% or 50% or 60% or 70% or 75% or 80% or 85% or 90% or 95%, 96%, 97%, 98%, 99% or 100% identity thereto. [0036] A fungal endo-beta-1,6-galactanase is obtainable from *Trichoderma*, for example *Trichoderma harzianum*. A preferred endo-beta-1,6-galactanase is obtainable from *Trichoderma harzianum* defined in SEQ ID NO: 3 herein, or a variant thereof, for example having at least 40% or 50% or 60% or 70% or 75% or 80% or 85% or 90% or 95%, 96%, 97%, 98%, 99% or 100% identical thereto.
- ⁴⁵ **[0037]** Other fungal endo-beta-1,6-galactanases include those encoded by the DNA sequences of *Ceratocystis fimbriata* f. sp. Platani, *Muscodor strobelii* WG-2009a, *Oculimacula yallundae, Trichoderma viride* GD36A, *Thermomyces stellatus, Myceliophthora thermophilia.*

[0038] Preferably the galactanase has an amino acid sequence having at least 60%, or at least 80%, or at least 90% or at least 95% identity with the amino acid sequence shown in SEQ ID NO:1, SEQ ID NO:2 or SEQ ID NO:3. Preferably the galactanase is an isolated galactanase.

[0039] Preferably the galactanase enzyme is present in a laundering aqueous liquor in an amount of from 0.01ppm to 1000 ppm of the galactanase enzyme, or from 0.05 or from 0.1ppm to 750 or 500ppm.

[0040] The galactanase or composition comprising galactanase may also give rise to biofilm-disrupting effects.

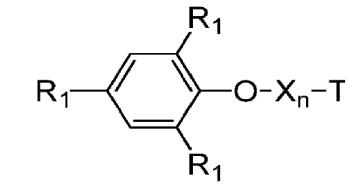
⁵⁵ Alkoxylated phenol compound

[0041] The cleaning compositions of the present invention include an alkoxylated phenol compound. The alkoxylated phenol compound may be selected from the group consisting of an alkoxylated polyaryl phenol compound, an alkoxylated

polyalkyl phenol compound, and mixtures thereof. The alkoxylated phenol compound may be an alkoxylated polyaryl phenol compound. The alkoxylated phenol compound may be an alkoxylated polyalkyl phenol compound.

[0042] The alkoxylated phenol compound may be present in the cleaning composition at a level of from about 0.2% to about 10%, or from about 0.5% to about 5%, by weight of the cleaning composition.

- [0043] The alkoxylated phenol compound preferably has a weight average molecular weight between 280 and 2880.
- [0044] The alkoxylated phenol compound may have the following structure:



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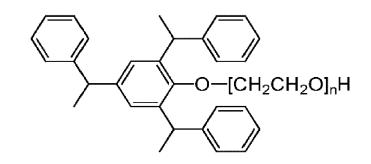
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wherein each R_1 is independently selected from linear of branched C3-C15 alkyl groups and aryl groups, X is selected from ethoxy or propoxy groups, n is from 2 to 70, T is selected from H, SO₃⁻, COO⁻ and PO₃²⁻, preferably H and SO₃⁻. Each R_1 may be the same.

[0045] The alkoxylated polyaryl or alkoxylated polyalkyl phenol compound is preferably selected from groups (i) to (v):

(i) Uncharged alkoxylated tristyrylphenols of the following structure:



wherein n is selected from 2 to 70, more preferably n is selected from 10 to 54, most preferably n = 16 or 20; (ii) Anionic alkoxylated tristyrylphenols of the following structure



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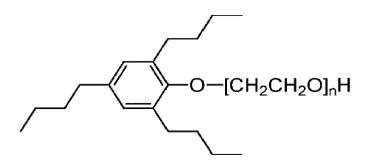
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wherein R is selected from SO₃⁻, COO- and PO₃²⁻, preferably selected from SO₃⁻ and COO⁻, wherein n is selected from 2 to 54;

CH₂CH₂O]_nR

(iii) Uncharged alkoxylated tri(n-butyl)phenols of the following structure:



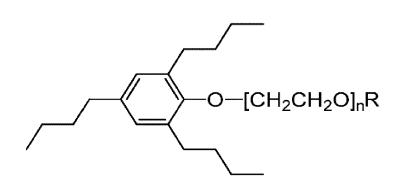
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wherein n is selected from 2 to 50; (iv) Anionic alkoxylated tri(n-butyl)phenols of the following structure:



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wherein R is selected from SO₃⁻, COO⁻ and PO₃²⁻, preferably selected from SO₃⁻ and COO⁻, wherein n is selected from 6 to 50; and

(v) mixtures thereof.

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[0046] Such compounds are available from industrial suppliers, for example Solvay under the Soprophor trade name, from Clariant under the Emulsogen trade name, Aoki Oil Industrial Co. under the Blaunon trade name, from Stepan under the Makon trade name, and from TOTO Chemical Industry Co. under the Sorpol trade name.

35 Adjuncts

[0047] The cleaning compositions described herein may optionally include other adjunct components, for example selected from surfactants, fabric shading dyes, fabric care benefit agent; additional enzyme; deposition aid; rheology modifier; builder; chelant; bleach; bleaching agent; bleach precursor; bleach booster; bleach activator, bleach catalyst; perfume and/or perfume microcapsules; perfume loaded zeolite; starch encapsulated accord; polyglycerol esters; whitening agent; pearlescent agent; enzyme stabilizing systems; scavenging agents including fixing agents for anionic dyes,

- complexing agents for anionic surfactants, and mixtures thereof; optical brighteners or fluorescers; polymer including but not limited to soil release polymer and/or soil suspension polymer; dispersants; antifoam agents; non-aqueous solvent; fatty acid; suds suppressors, e.g., silicone suds suppressors; cationic starches; scum dispersants; substantive
 dves; colorants; opacifier; antioxidant; hvdrotropes such as toluenesulfonates, cumenesulfonates and naphthalenesul-
- ⁴⁵ dyes; colorants; opacifier; antioxidant; hydrotropes such as toluenesulfonates, cumenesulfonates and naphthalenesulfonates; color speckles; colored beads, spheres or extrudates; clay softening agents; anti-bacterial agents. Additionally or alternatively, the compositions may comprise surfactants, and/or solvent systems. Quaternary ammonium compounds may be present, particularly in fabric enhancer compositions, such as fabric softeners, and comprise quaternary ammonium cations that are positively charged polyatomic ions of the structure NR₄⁺, where R is an alkyl group or an aryl group.

Additional Enzymes

[0048] Preferably the composition of the invention comprises additional enzymes, for example selected from lipases, amylases, proteases, nucleases, pectate lyases, cellulases, cutinases, and mixtures thereof. The cleaning compositions preferably comprise one or more additional enzymes from the group selected from nucleases. The cleaning compositions preferably comprises one or more additional enzymes selected from the group amylases, lipases, proteases, pectate lyases, cellulases, cellulases, cutinases, and mixtures thereof. The cleaning compositions preferably comprises one or more additional enzymes selected from the group amylases, lipases, proteases, pectate lyases, cellulases, cutinases, and mixtures thereof. Preferably, the cleaning compositions comprises one or more additional enzymes for the group and lyases.

tional enzymes selected from amylases and proteases and mixtures thereof. Preferably the cleaning compositions comprise one or more additional enzymes selected from lipases. The compositions may also comprise hemicellulases, peroxidases, xylanases, pectinases, keratinases, reductases, oxidases, phenoloxidases, lipoxygenases, ligninases, pullulanases, tannases, pentosanases, malanases, β -glucanases, arabinosidases, hyaluronidase, chondroitinase, lac-

⁵ case and mixtures thereof. When present in the composition, the aforementioned additional enzymes may be present at levels from about 0.0001% to about 2%, from about 0.0001 % to about 1% or even from about 0.001 % to about 0.5% enzyme protein by weight of the composition. Preferably the or each additional enzyme is present in the laundering aqueous liquor in an amount of from 0.01ppm to 1000 ppm of the active enzyme protein, or from 0.05 or from 0.1ppm to 750 or 500ppm.

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Nucleases

[0049] Preferably the composition additionally comprises a nuclease enzyme. The nuclease enzyme is an enzyme capable of cleaving the phosphodiester bonds between the nucleotide subunits of nucleic acids. Suitable nuclease enzymes may be deoxyribonuclease or ribonuclease enzyme or a functional fragment thereof. By functional fragment or part is meant the portion of the nuclease enzyme that catalyzes the cleavage of phosphodiester linkages in the DNA backbone and so is a region of said nuclease protein that retains catalytic activity. Thus it includes truncated, but functional versions, of the enzyme and/or variants and/or derivatives and/or homologues whose functionality is maintained.

- [0050] Preferably the nuclease enzyme is a deoxyribonuclease, preferably selected from any of the classes E.C. 3.1.21.x, where x=1, 2, 3, 4, 5, 6, 7, 8 or 9, E.C. 3.1.22.y where y=1, 2, 4 or 5, E.C. 3.1.30.z where z= 1 or 2, E.C. 3.1.31.1 and mixtures thereof. Nuclease enzymes from class E.C. 3.1.21.x and especially where x=1 are particularly preferred. Nucleases in class E.C. 3.1.22.y cleave at the 5' hydroxyl to liberate 3' phosphomonoesters. Enzymes in class E.C. 3.1.30.z may be preferred as they act on both DNA and RNA and liberate 5'-phosphomonoesters. Suitable examples from class E.C. 3.1.31.2 are described in US2012/0135498A, such as SEQ ID NO:3 therein. Such enzymes are com-
- ²⁵ mercially available as DENARASE® enzyme from c-LECTA. Nuclease enzymes from class E.C. 3.1.31.1 produce 3'phosphomonoesters.

[0051] Preferably, the nuclease enzyme comprises a microbial enzyme. The nuclease enzyme may be fungal or bacterial in origin. Bacterial nucleases may be most preferred. Fungal nucleases may be most preferred.

- [0052] The microbial nuclease is obtainable from *Bacillus*, such as a *Bacillus licheniformis* or *Bacillus subtilis* bacterial nucleases. A preferred nuclease is obtainable from *Bacillus licheniformis*, preferably from strain EI-34-6. A preferred deoxyribonuclease is a variant of *Bacillus licheniformis*, from strain EI-34-6 nucB deoxyribonuclease defined in SEQ ID NO:4 herein, or variant thereof, for example having at least 70% or 75% or 80% or 85% or 90% or 95%, 96%, 97%, 98%, 99% or 100% identical thereto. Other suitable nucleases are defined in SEQ ID NO:5 herein, or variant thereof, for example having at least 70% or 95%, 96%, 97%, 98%, 99% or 100% identical thereto.
- Other suitable nucleases are defined in SEQ ID NO: 6 herein, or variant thereof, for example having at least 70% or 75% or 80% or 85% or 90% or 95%, 96%, 97%, 98%, 99% or 100% identical thereto.
 [0053] A fungal nuclease is obtainable from *Aspergillus*, for example *Aspergillus oryzae*. A preferred nuclease is obtainable from *Aspergillus oryzae* defined in SEQ ID NO: 7 herein, or variant thereof, for example having at least 60% or 70% or75% or 80% or 85% or 90% or 95%, 96%, 97%, 98%, 99% or 100% identical thereto.
- 40 [0054] Another suitable fungal nuclease is obtainable from *Trichoderma*, for example *Trichoderma harzianum*. A preferred nuclease is obtainable from *Trichoderma harzianum* defined in SEQ ID NO: 8 herein, or variant thereof, for example having at least 60% or 70% or 75% or 80% or 85% or 90% or 95%, 96%, 97%, 98%, 99% or 100% identical thereto. [0055] Other fungal nucleases include those encoded by the DNA sequences of *Aspergillus oryzae* RIB40, *Aspergillus oryzae* 3.042, *Aspergillus flavus* NRRL3357, *Aspergillus parasiticus* SU-1, *Aspergillus nomius* NRRL13137, *Trichoderma*
- ⁴⁵ reesei QM6a, Trichoderma virens Gv29-8, Oidiodendron maius Zn, Metarhizium guizhouense ARSEF 977, Metarhizium majus ARSEF 297, Metarhizium robertsii ARSEF 23, Metarhizium acridum CQMa 102, Metarhizium brunneum ARSEF 3297, Metarhizium anisopliae, Colletotrichum fioriniae PJ7, Colletotrichum sublineola, Trichoderma atroviride IMI 206040, Tolypocladium ophioglossoides CBS 100239, Beauveria bassiana ARSEF 2860, Colletotrichum higginsianum, Hirsutella minnesotensis 3608, Scedosporium apiospermum, Phaeomoniella chlamydospora, Fusarium verticillioides
- 50 7600, Fusarium oxysporum f. sp. cubense race 4, Colletotrichum graminicola M1.001, Fusarium oxysporum FOSC 3a, Fusarium avenaceum, Fusarium langsethiae, Grosmannia clavigera kw1407, Claviceps purpurea 20.1, Verticillium longisporum, Fusarium oxysporum f. sp. cubense race 1, Magnaporthe oryzae 70-15, Beauveria bassiana D1-5, Fusarium pseudograminearum CS3096, Neonectria ditissima, Magnaporthiopsis poae ATCC 64411, Cordyceps militaris CM01, Marssonina brunnea f. sp. 'multigermtubi' MB_m1, Diaporthe ampelina, Metarhizium album ARSEF 1941, Colletotrichum
- ⁵⁵ gloeosporioides Nara gc5, Madurella mycetomatis, Metarhizium brunneum ARSEF 3297, Verticillium alfalfae VaMs.102, Gaeumannomyces graminis var. tritici R3-111a-1, Nectria haematococca mpVI 77-13-4, Verticillium longisporum, Verticillium dahliae VdLs.17, Torrubiella hemipterigena, Verticillium longisporum, Verticillium dahliae VdLs.17, Botrytis cinerea B05.10, Chaetomium globosum CBS 148.51, Metarhizium anisopliae, Stemphylium lycopersici, Sclerotinia bore-

alis F-4157, Metarhizium robertsii ARSEF 23, Myceliophthora thermophila ATCC 42464, Phaeosphaeria nodorum SN15, Phialophora attae, Ustilaginoidea virens, Diplodia seriata, Ophiostoma piceae UAMH 11346, Pseudogymnoascus pannorum VKM F-4515 (FW-2607), Bipolaris oryzae ATCC 44560, Metarhizium guizhouense ARSEF 977, Chaetomium thermophilum var. thermophilum DSM 1495, Pestalotiopsis fici W106-1, Bipolaris zeicola 26-R-13, Setosphaeria turcica

5 Et28A, Arthroderma otae CBS 113480 and Pyrenophora tritici-repentis Pt-1C-BFP.

[0056] Preferably the nuclease is an isolated nuclease.

[0057] Preferably the nuclease enzyme is present in the laundering aqueous liquor in an amount of from 0.01ppm to 1000 ppm of the nuclease enzyme, or from 0.05 or from 0.1ppm to 750 or 500ppm.

10 Acetylglucosaminidases.

[0058] Preferably the composition comprises an acetylglucosaminidase enzyme, preferably a β -N-acetylglucosaminidase enzyme from E.C. 3.2.1.52, preferably an enzyme having at least 70%, or at least 75% or at least 80% or at least 85% or at least 90% or at least 95% or at least 96% or at least 97% or at least 98% or at least 99% or at least or 100%

15 identity to SEQ ID NO: 9.

Mannanases

- [0059] Preferably the composition comprises a mannanase enzyme. The term "mannanase" means a polypeptide 20 having mannan endo-1,4- beta-mannosidase activity (EC 3.2.1.78) from the glycoside hydrolase family 26 that catalyzes the hydrolysis of 1,4-3-D-mannosidic linkages in mannans, galactomannans and glucomannans. Alternative names of mannan endo-1,4-beta-mannosidase are 1,4-3-D-mannan mannanohydrolase; endo-1,4-3-mannanase; endo-β-1,4mannase; β-mannanase B; 3-1,4-mannan 4-mannanohydrolase; endo-3-mannanase; and β-D-mannanase. Preferred mannanases are members of the glycoside hydrolase family 26.
- 25 [0060] For purposes of the present disclosure, mannanase activity may be determined using the Reducing End Assay as described in the experimental section of WO 2015040159. Suitable examples from class EC 3.2.1.78 are described in WO 2015040159, such as the mature polypeptide SEQ ID NO: 2 described therein. [0061] Preferred mannanases are variants having at least 60%, at least 65%, at least 70%, at least 75%, at least 80%,
- at least 81 %, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 30 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99% or 100% sequence identity to the mature polypeptide SEQ ID NO: 10 from Ascobolus stictoideus; Preferred mannanases are variants having at least 81 %, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91 %, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99% or 100% sequence identity to the mature polypeptide 35

SEQ ID NO: 11 from Chaetomium virescens. [0062] Preferred mannanases are variants having at least 75%, at least 76%, at least 77%, at least 78%, at least 79%, at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99% or 100% sequence identity to the mature polypeptide SEQ ID NO: 12 from Preussia 40 aemulans.

[0063] Preferred mannanases are variants having at least at least 65%, at least 66%, at least 67%, at least 68%, at least 69%, at least 70%, at least 71%, at least 72%, at least 73%, at least 74%, at least 75%, at least 76%, at least 77%, at least 78%, at least 79%, at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91 %, at least 92%, at least 93%, at least 94%, at

45 least 95%, at least 96%, at least 97%, at least 98%, at least 99% or 100% sequence identity to the mature polypeptide SEQ ID NO: 13 from Yunnania penicillata. [0064] Preferred mannanases are variants having at least at least 75%, at least 76%, at least 77%, at least 78%, at least 79%, at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least

- 50 96%, at least 97%, at least 98%, at least 99% or 100% sequence identity to the mature polypeptide SEQ ID NO: 14 from *Myrothecium roridum*. Preferably the mannanase is an isolated mannanase. [0065] Preferably the mannanase enzyme is present in the cleaning compositions in an amount from 0.001 to 1 wt% based on active protein in the composition, or from 0.005 to 0.5 wt% or from 0.01 to 0.25 wt%. Preferably the mannanase enzyme is present in the laundering aqueous liquor in an amount of from 0.01ppm to 1000 ppm of the mannanase
- 55 enzyme, or from 0.05 or from 0.1 ppm to 750 or 500ppm. The compositions of the invention comprising both galactanase and mannanase may be particularly effective against sticky soils and for improved cleaning. It is believed the two enzymes function together in a complementary way.

Further Glycosyl Hydrolases

[0066] The composition may comprise a glycosyl hydrolase selected from GH family 39 and GH family 114 and mixtures thereof, for example as described in co-pending applications having applicants reference numbers CM4645FM and CM4646 FM, respectively.

Proteases.

[0067] Preferably the composition comprises one or more proteases. Suitable proteases include metalloproteases and serine proteases, including neutral or alkaline microbial serine proteases, such as subtilisins (EC 3.4.21.62). Suitable proteases include those of animal, vegetable or microbial origin. In one aspect, such suitable protease may be of microbial origin. The suitable proteases include chemically or genetically modified mutants of the aforementioned suitable proteases. In one aspect, the suitable protease may be a serine protease, such as an alkaline microbial protease or/and a trypsin-type protease. Examples of suitable neutral or alkaline proteases include:

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(a) subtilisins (EC 3.4.21.62), preferably those derived from *Bacillus sp.*, such as *B. lentus*, *B. alkalophilus*, *B. subtilis*, *B. amyloliquefaciens*, *B. pumilus* and *B.* gibsonii and *B. akibaii* described in WO2004067737, WO2015091989, WO2015091990, WO2015024739, WO2015143360, US 6,312,936 B1, US 5,679,630, US 4,760,025, US7,262,042 and WO09/021867, DE102006022216A1, DE102006022224A1, WO2015089447, WO2015089467, WO201508947, WO201508948, WO20150898, WO2015089, WO2015089, WO20150898, WO2015089, WO20150898, WO20150

- WO2016066756, WO2016066757, WO2016069557, WO2016069563, WO2016069569.
 (b) trypsin-type or chymotrypsin-type proteases, such as trypsin (e.g., of porcine or bovine origin), including the *Fusarium* protease described in WO 89/06270 and the chymotrypsin proteases derived from *Cellumonas* described in WO 05/052161 and WO 05/052146.
- (c) metalloproteases, preferably those derived from *Bacillus amyloliquefaciens* described in WO 07/044993A2; from
 Bacillus, Brevibacillus, Thermoactinomyces, Geobacillus, Paenibacillus, Lysinibacillus or *Streptomyces spp.* Described in WO2014194032, WO2014194054 and WO2014194117; from *Kribella alluminosa* described in WO2015193488; and from *Streptomyces* and *Lysobacter* described in WO2016075078.

(d) protease having at least 90% identity to the subtilase from Bacillus sp. TY145, NCIMB 40339, described in WO92/17577 (Novozymes A/S), including the variants of this Bacillus sp TY145 subtilase described in WO2015024739, and WO2016066757.

[0068] Preferred proteases include those derived from Bacillus gibsonii or Bacillus Lentus.

[0069] Suitable commercially available protease enzymes include those sold under the trade names Alcalase®, Savinase®, Primase®, Durazym®, Polarzyme®, Kannase®, Liquanase®, Liquanase Ultra®, Savinase Ultra®, Ovozyme®, Neutrase®, Everlase® and Esperase® by Novozymes A/S (Denmark), those sold under the tradename Maxatase®,

- Neutrase®, Everlase® and Esperase® by Novozymes A/S (Denmark), those sold under the tradename Maxatase®, Maxacal®, Maxapem®, Properase®, Purafect®, Purafect Prime®, Purafect Ox®, FN3®, FN4®, Excellase® and Purafect OXP® by Genencor International, those sold under the tradename Opticlean® and Optimase® by Solvay Enzymes, those available from Henkel/ Kemira, namely BLAP (sequence shown in Figure 29 of US 5,352,604 with the following mutations S99D + S101 R + S103A + V104I + G159S, hereinafter referred to as BLAP), BLAP R (BLAP with S3T + V4I
- + V199M + V205I + L217D), BLAP X (BLAP with S3T + V4I + V205I) and BLAP F49 (BLAP with S3T + V4I + A194P + V199M + V205I + L217D) all from Henkel/Kemira; and KAP (*Bacillus alkalophilus* subtilisin with mutations A230V + S256G + S259N) from Kao, or as disclosed in WO2009/149144, WO2009/149145, WO2010/56653, WO2010/56640, WO2011/07211 US2011/0237487, WO2011/140316, WO2012/151480, EP2510092, EP2566960 OR EP2705145.
- 45 Amylases

[0070] Preferably the composition may comprise an amylase. Suitable alpha-amylases include those of bacterial or fungal origin. Chemically or genetically modified mutants (variants) are included. A preferred alkaline alpha-amylase is derived from a strain of Bacillus, such as Bacillus licheniformis, Bacillus amyloliquefaciens, Bacillus stearothermophilus, Bacillus subtilis, or other Bacillus sp., such as Bacillus sp. NCIB 12289, NCIB 12512, NCIB 12513, DSM 9375 (USP 7,153,818) DSM 12368, DSMZ no. 12649, KSM AP1378 (WO 97/00324), KSM K36 or KSM K38 (EP 1,022,334). Preferred amylases include:

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(a) the variants described in WO 94/02597, WO 94/18314, WO96/23874 and WO 97/43424, especially the variants with substitutions in one or more of the following positions versus the enzyme listed as SEQ ID No. 2 in WO 96/23874: 15, 23, 105, 106, 124, 128, 133, 154, 156, 181, 188, 190, 197, 202, 208, 209, 243, 264, 304, 305, 391, 408, and 444.
(b) the variants described in USP 5,856,164 and WO99/23211, WO 96/23873, WO00/60060 and WO 06/002643, especially the variants with one or more substitutions in the following positions versus the AA560 enzyme listed as

SEQ ID No. 12 in WO 06/002643:

26, 30, 33, 82, 37, 106, 118, 128, 133, 149, 150, 160, 178, 182, 186, 193, 203, 214, 231, 256, 257, 258, 269, 270, 272, 283, 295, 296, 298, 299, 303, 304, 305, 311, 314, 315, 318, 319, 339, 345, 361, 378, 383, 419, 421, 437, 441, 444, 445, 446, 447, 450, 461, 471, 482, 484, preferably that also contain the deletions of D183* and G184*.

(c) variants exhibiting at least 90% identity with SEQ ID No. 4 in WO06/002643, the wild-type enzyme from Bacillus SP722, especially variants with deletions in the 183 and 184 positions and variants described in WO 00/60060, which is incorporated herein by reference.

(d) variants exhibiting at least 95% identity with the wild-type enzyme from Bacillus sp.707 (SEQ ID NO:7 in US 6,093, 562), especially those comprising one or more of the following mutations M202, M208, S255, R172, and/or M261. Preferably said amylase comprises one or more of M202L, M202V, M202S, M202T, M202I, M202Q, M202W, S255N and/or R172Q. Particularly preferred are those comprising the M202L or M202T mutations.

(e) variants described in WO 09/149130, preferably those exhibiting at least 90% identity with SEQ ID NO: 1 or SEQ ID NO:2 in WO 09/149130, the wild-type enzyme from Geobacillus Stearophermophilus or a truncated version thereof;

(f) variants as described in EP2540825 and EP2357220, EP2534233; (g) variants as described in WO2009100102 and WO2010115028.

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[0071] Suitable commercially available alpha-amylases include DURAMYL®, LIQUEZYME®, TERMAMYL®, TER-MAMYL ULTRA®, NATALASE®, SUPRAMYL®, STAINZYME®, STAINZYME PLUS®, FUNGAMYL® and BAN® (Novozymes A/S, Bagsvaerd, Denmark), KEMZYM® AT 9000 Biozym Biotech Trading GmbH Wehlistrasse 27b A-1200 Wien Austria, RAPIDASE®, PURASTAR®, ENZYSIZE®, OPTISIZE HT PLUS®, POWERASE® and PURASTAR OX-

²⁵ AM® (Genencor International Inc., Palo Alto, California) and KAM® (Kao, 14-10 Nihonbashi Kayabacho, 1-chome, Chuoku Tokyo 103-8210, Japan). In one aspect, suitable amylases include NATALASE®, STAINZYME® and STAINZYME PLUS® and mixtures thereof.

Lipases

[0072] Preferably the composition comprises one or more lipases, including "first cycle lipases" such as those described in U.S. Patent 6,939,702 B1 and US PA 2009/0217464. Preferred lipases are first-wash lipases. In one embodiment of the invention the composition comprises a first wash lipase. First wash lipases includes a lipase which is a polypeptide having an amino acid sequence which: (a) has at least 90% identity with the wild-type lipase derived from *Humicola*

- ³⁵ *lanuginosa* strain DSM 4109; (b) compared to said wild-type lipase, comprises a substitution of an electrically neutral or negatively charged amino acid at the surface of the three-dimensional structure within 15A of E1 or Q249 with a positively charged amino acid; and (c) comprises a peptide addition at the C-terminal; and/or (d) comprises a peptide addition at the N-terminal and/or (e) meets the following limitations: i) comprises a negative amino acid in position E210 of said wild-type lipase; ii) comprises a negatively charged amino acid so positions 90-101
- 40 of said wild-type lipase; and iii) comprises a neutral or negative amino acid at a position corresponding to N94 or said wild-type lipase and/or has a negative or neutral net electric charge in the region corresponding to positions 90-101 of said wild-type lipase. Preferred are variants of the wild-type lipase from *Thermomyces lanuginosus* comprising one or more of the T231R and N233R mutations. The wild-type sequence is the 269 amino acids (amino acids 23 - 291) of the Swissprot accession number Swiss-Prot 059952 (derived from *Thermomyces lanuginosus* (*Humicola lanuginosa*)). Pre-
- ⁴⁵ ferred lipases include those sold under the tradenames Lipex® and Lipolex® and Lipoclean®. Other suitable lipases include those described in European Patent Application No. 12001034.3 or EP2623586.

Endoglucanases

⁵⁰ **[0073]** Other preferred enzymes include microbial-derived endoglucanases exhibiting endo-beta-1,4-glucanase activity (E.C. 3.2.1.4), including a bacterial polypeptide endogenous to a member of the genus Bacillus which has a sequence of at least 90%, 94%, 97% and even 99% identity to the amino acid sequence SEQ ID NO:2 in US7,141,403B2) and mixtures thereof. Suitable endoglucanases are sold under the tradenames Celluclean® and Whitezyme® (Novozymes A/S, Bagsvaerd, Denmark).

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Pectate Lyases

[0074] Other preferred enzymes include pectate lyases sold under the tradenames Pectawash®, Pectaway®, Xpect®

and mannanases sold under the tradenames Mannaway® (all from Novozymes A/S, Bagsvaerd, Denmark), and Purabrite® (Genencor International Inc., Palo Alto, California).

Cleaning Cellulase

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[0075] The cleaning composition described herein may additionally comprise a cleaning cellulase. The cellulase may be an endoglucanase. The cellulase may have endo beta 1,4-glucanase activity and a structure which does not comprise a class A Carbohydrate Binding Module (CBM). A class A CBM is defined according to A. B. Boraston et al. Biochemical Journal 2004, Volume 382 (part 3) pages 769-781. In particular, the cellulase does not comprise a class A CBM from families 1, 2a, 3, 5 and 10.

[0076] The cellulase may be a glycosyl hydrolase having enzymatic activity towards amorphous cellulose substrates, wherein the glycosyl hydrolase is selected from GH families 5, 7, 12, 16, 44 or 74. Preferably, the cellulase is a glycosyl hydrolase selected from GH family 5. A preferred cellulase is Celluclean, supplied by Novozymes. This preferred cellulase is described in more detail in WO2002/099091. The glycosyl hydrolase (GH) family definition is described in more detail

¹⁵ in Biochem J. 1991, v280, 309-316. Another preferred cellulase is a glycosyl hydrolase having enzymatic activity towards both xyloglucan and amorphous cellulose substrates, wherein the glycosyl hydrolase is selected from GH families 5, 12, 44 or 74. Preferably, the glycosyl hydrolase selected from GH family 44.

[0077] For purposes of the present invention, the degree of identity between two amino acid sequences is determined using the Needleman-Wunsch algorithm (Needleman and Wunsch, 1970, J. Mol. Biol. 48: 443-453) as implemented in the Needle program of the EMBOSS package (EMBOSS: The European Molecular Biology Open Software Suite, Rice et al., 2000, Trends in Genetics 16: 276-277), preferably version 3.0.0 or later. The optional parameters used are gap open penalty of 10, gap extension penalty of 0.5, and the EBLOSUM62 (EMBOSS version of BLOSUM62) substitution matrix. The output of Needle labeled "longest identity" (obtained using the -nobrief option) is used as the percent identity

- and is calculated as follows: (Identical Residues x 100)/(Length of Alignment Total Number of Gaps in Alignment).
 [0078] Suitable cleaning cellulase glycosyl hydrolases are selected from the group consisting of: GH family 44 glycosyl hydrolases from *Paenibacillus polyxyma* (wild-type) such as XYG1006 described in WO 01/062903 or are variants thereof; GH family 12 glycosyl hydrolases from *Bacillus licheniformis* (wild-type) such as Seq. No. ID: 1 described in WO 99/02663 or are variants thereof; GH family 5 glycosyl hydrolases from *Paenibacillus* (wild type) or variants thereof; GH family 5 glycosyl hydrolases from *Paenibacillus* (wild type) such as XYG1034 and XYG 1022described in
- 30 WO 01/064853 or variants thereof; GH family 74 glycosyl hydrolases from *Jonesia sp.* (wild type) such as XYG1020 described in WO 2002/077242 or variants thereof; and GH family 74 glycosyl hydrolases from *Trichoderma Reesei* (wild type), such as the enzyme described in more detail in Sequence ID no. 2 of WO03/089598, or variants thereof. [0079] Preferred glycosyl hydrolases are selected from the group consisting of: GH family 44 glycosyl hydrolases from *Paenibacillus polyxyma* (wild-type) such as XYG1006 or are variants thereof.
- ³⁵ **[0080]** Typically, the cellulase modifies the fabric surface during the laundering process so as to improve the removal of soils adhered to the fabric after the laundering process during wearing and usage of the fabric, in subsequent wash cycles. Preferably, the cellulase modifies the fabric surface during the laundering process so as to improve the removal of soils adhered to the fabric after the laundering process during wearing and usage of the fabric, in the subsequent two, or even three wash cycles.
- 40 [0081] Typically, the cellulase is used at a concentration of 0.005ppm to 1.0ppm in the aqueous liquor during the first laundering process. Preferably, the cellulase is used at a concentration of 0.02ppm to 0.5ppm in the aqueous liquor during the first laundering process.

Surfactant system

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[0082] The cleaning composition may comprise a surfactant system. The cleaning composition may comprise from about 1% to about 80%, or from 1% to about 60%, preferably from about 5% to about 50% more preferably from about 8% to about 40%, by weight of the cleaning composition, of a surfactant system.

[0083] Surfactants suitable for use in the surfactant system may be derived from natural and/or renewable sources.

- ⁵⁰ **[0084]** The surfactant system may comprise an anionic surfactant, more preferably an anionic surfactant selected from the group consisting of, alkyl benzene sulfonate, alkyl sulfate, alkyl alkoxy sulfate, especially alkyl ethoxy sulfate, paraffin sulfonate and mixtures thereof, alkyl benzene sulfonates are particularly preferred. The surfactant system may further comprise a surfactant selected from the group consisting of nonionic surfactant, cationic surfactant, amphoteric surfactant, zwitterionic surfactant, and mixtures thereof. The surfactant system preferably comprises a nonionic sur-
- ⁵⁵ factant, for example an ethoxylated nonionic surfactant. The surfactant system may comprise an amphoteric surfactant, for example an amine oxide surfactant, such as an alkyl dimethyl amine oxide. The surfactant system may comprise a zwitterionic surfactant, such as a betaine.

[0085] The most preferred surfactant system for the detergent composition of the present invention comprises from

1% to 40%, preferably 6% to 35%, more preferably 8% to 30% weight of the total composition of an anionic surfactant, preferably comprising an alkyl benzene sulphonate. The preferred surfactant system may optionally in addition comprise an alkyl alkoxy sulfate surfactant, more preferably an alkyl ethoxy sulfate, optionally combined with 0.5% to 15%, preferably from 1% to 12%, more preferably from 2% to 10% by weight of the composition of amphoteric and/or zwitterionic

5 surfactant, more preferably an amphoteric and even more preferably an amine oxide surfactant, especially an alkyl dimethyl amine oxide.

[0086] Preferably the composition further comprises a nonionic surfactant, especially an alcohol alkoxylate in particular an alcohol ethoxylate nonionic surfactant. Most preferably the surfactant system comprises an anionic and a nonionic surfactant, preferably the weight ratio of the anionic to nonionic surfactant is from 25:1 to 1:2.

Anionic surfactant

[0087] Anionic surfactants may be in salt form or acid form, typically in the form of a water-soluble sodium, potassium, ammonium, magnesium or mono-, di- or tri- C2-C3 alkanolammonium salt, with the sodium cation being the usual one chosen.

Sulfonate Surfactant

[0088] Suitable anionic sulfonate surfactants for use herein include water-soluble salts of C8-C18 alkyl or hydroxyalkyl 20 sulfonates; C11-C18 alkyl benzene sulfonates (LAS), modified alkylbenzene sulfonate (MLAS) as discussed in WO 99/05243, WO 99/05242, WO 99/05244, WO 99/05082, WO 99/05084, WO 99/05241, WO 99/07656, WO 00/23549, and WO 00/23548; methyl ester sulfonate (MES); and alpha-olefin sulfonate (AOS). Those also include the paraffin sulfonates may be monosulfonates and/or disulfonates, obtained by sulfonating paraffins of 10 to 20 carbon atoms. The sulfonate surfactant may also include the alkyl glyceryl sulfonate surfactants.

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Sulfated anionic surfactant

[0089] Preferably the sulfated anionic surfactant is alkoxylated, more preferably, an alkoxylated branched sulfated anionic surfactant having an alkoxylation degree of from about 0.2 to about 4, even more preferably from about 0.3 to 30 about 3, even more preferably from about 0.4 to about 1.5 and especially from about 0.4 to about 1. Preferably, the alkoxy group is ethoxy. When the sulfated anionic surfactant is a mixture of sulfated anionic surfactants, the alkoxylation degree is the weight average alkoxylation degree of all the components of the mixture (weight average alkoxylation degree). In the weight average alkoxylation degree calculation the weight of sulfated anionic surfactant components not having alkoxylated groups should also be included.

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Weight average alkoxylation degree = (x1 * alkoxylation degree of surfactant 1 + x2 *

alkoxylation degree of surfactant 2 +) / (x1 + x2 +)

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wherein x1, x2, ... are the weights in grams of each sulfated anionic surfactant of the mixture and alkoxylation degree is the number of alkoxy groups in each sulfated anionic surfactant.

[0090] Preferably, the branching group is an alkyl. Typically, the alkyl is selected from methyl, ethyl, propyl, butyl, pentyl, cyclic alkyl groups and mixtures thereof. Single or multiple alkyl branches could be present on the main hydrocarbyl chain of the starting alcohol(s) used to produce the sulfated anionic surfactant used in the detergent of the invention. Most preferably the branched sulfated anionic surfactant is selected from alkyl sulfates, alkyl ethoxy sulfates, and mixtures thereof.

[0091] The branched sulfated anionic surfactant can be a single anionic surfactant or a mixture of anionic surfactants. In the case of a single surfactant the percentage of branching refers to the weight percentage of the hydrocarbyl chains that are branched in the original alcohol from which the surfactant is derived.

[0092] In the case of a surfactant mixture the percentage of branching is the weight average and it is defined according to the following formula:

Weight average of branching (%)= [(x1 * wt% branched alcohol 1 in alcohol 1 + x2 *55

wt% branched alcohol 2 in alcohol 2 +) / (x1 + x2 +)] * 100

wherein x1, x2, ... are the weight in grams of each alcohol in the total alcohol mixture of the alcohols which were used as starting material for the anionic surfactant for the detergent of the invention. In the weight average branching degree calculation the weight of anionic surfactant components not having branched groups should also be included.

[0093] Suitable sulfate surfactants for use herein include water-soluble salts of C8-C18 alkyl or hydroxyalkyl, sulfate and/or ether sulfate. Suitable counterions include alkali metal cation or ammonium or substituted ammonium, but preferably sodium.

[0094] The sulfate surfactants may be selected from C8-C18 primary, branched chain and random alkyl sulfates (AS); C8-C18 secondary (2,3) alkyl sulfates; C8-C18 alkyl alkoxy sulfates (AExS) wherein preferably x is from 1-30 in which the alkoxy group could be selected from ethoxy, propoxy, butoxy or even higher alkoxy groups and mixtures thereof.

- [0095] Alkyl sulfates and alkyl alkoxy sulfates are commercially available with a variety of chain lengths, ethoxylation and branching degrees. Commercially available sulfates include, those based on Neodol alcohols ex the Shell company, Lial Isalchem and Safol ex the Sasol company, natural alcohols ex The Procter & Gamble Chemicals company.
 [0096] Preferred alkyl sulfates are those in which the anionic surfactant is an alkyl ethoxy sulfate with a degree of
- ethoxylation of from about 0.2 to about 3, more preferably from about 0.3 to about 2, even more preferably from about
 0.4 to about 1.5, and especially from about 0.4 to about 1. They are also preferred anionic surfactant having a level of branching of from about 5% to about 40%, even more preferably from about 10% to 35% and especially from about 20% to 30%.

Nonionic surfactant

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[0097] Preferably the surfactant system comprises a nonionic surfactant, in an amount of from 0.1% to 40%, preferably 0.2% to 20%, most preferably 0.5% to 10% by weight of the composition. Suitable nonionic surfactants include the condensation products of aliphatic alcohols with from 1 to 25 moles of ethylene oxide. The alkyl chain of the aliphatic alcohol can either be straight or branched, primary or secondary, and generally contains from 8 to 22 carbon atoms.

- Particularly preferred are the condensation products of alcohols having an alkyl group containing from 10 to 18 carbon atoms, preferably from 10 to 15 carbon atoms with from 2 to 18 moles, preferably 2 to 15, more preferably 5-12 of ethylene oxide per mole of alcohol. Highly preferred nonionic surfactants are the condensation products of guerbet alcohols with from 2 to 18 moles, preferably 2 to 15, more preferably 5-12 of ethylene oxide per mole of alcohol. **100981** Other suitable non-ionic surfactants for use berein include fatty alcohol polyglycol ethers, alkylpolyglycolycol ethers.
- [0098] Other suitable non-ionic surfactants for use herein include fatty alcohol polyglycol ethers, alkylpolyglucosides and fatty acid glucamides.

Amphoteric surfactant

- [0099] The surfactant system may include amphoteric surfactant, such as amine oxide. Preferred amine oxides are alkyl dimethyl amine oxide or alkyl amido propyl dimethyl amine oxide, more preferably alkyl dimethyl amine oxide and especially coco dimethyl amino oxide. Amine oxide may have a linear or mid-branched alkyl moiety. Typical linear amine oxides include water-soluble amine oxides containing one R1 C8-18 alkyl moiety and 2 R2 and R3 moieties selected from the group consisting of C1-3 alkyl groups and C1-3 hydroxyalkyl groups. Preferably amine oxide is characterized by the formula R1 N(R2)(R3) 0 wherein R1 is a C8-18 alkyl and R2 and R3 are selected from the group consisting of
- 40 methyl, ethyl, propyl, isopropyl, 2-hydroxethyl, 2-hydroxypropyl and 3-hydroxypropyl. The linear amine oxide surfactants in particular may include linear C10-C18 alkyl dimethyl amine oxides and linear C8-C12 alkoxy ethyl dihydroxy ethyl amine oxides. Preferred amine oxides include linear C10, linear C10-C12, and linear C12-C14 alkyl dimethyl amine oxides. As used herein "mid-branched" means that the amine oxide has one alkyl moiety having n1 carbon atoms with one alkyl branch on the alkyl moiety having n2 carbon atoms. The alkyl branch is located on the α carbon from the
- ⁴⁵ nitrogen on the alkyl moiety. This type of branching for the amine oxide is also known in the art as an internal amine oxide. The total sum of n1 and n2 is from 10 to 24 carbon atoms, preferably from 12 to 20, and more preferably from 10 to 16. The number of carbon atoms for the one alkyl moiety (n1) should be approximately the same number of carbon atoms as the one alkyl branch (n2) such that the one alkyl moiety and the one alkyl branch are symmetric. As used herein "symmetric" means that |n1 n2| is less than or equal to 5, preferably 4, most preferably from 0 to 4 carbon atoms
- in at least 50 wt%, more preferably at least 75 wt% to 100 wt% of the mid-branched amine oxides for use herein.
 [0100] The amine oxide may further comprise two moieties, independently selected from a C1-3 alkyl, a C1-3 hydroxy-alkyl group, or a polyethylene oxide group containing an average of from about 1 to about 3 ethylene oxide groups. Preferably the two moieties are selected from a C1-3 alkyl, more preferably both are selected as a C 1 alkyl.

55 Zwitterionic surfactant

[0101] Other suitable surfactants include betaines, such as alkyl betaines, alkylamidobetaine, amidazoliniumbetaine, sulfobetaine (INCI Sultaines) as well as the Phosphobetaine and preferably meets formula (I):

$$R^{1}-[CO-X (CH_{2})_{n}]_{x}-N^{+}(R^{2})(R_{3})-(CH_{2})_{m}-[CH(OH)-CH_{2}]_{y}-Y-$$
(I)

wherein 5

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- R¹ is a saturated or unsaturated C6-22 alkyl residue, preferably C8-18 alkyl residue, in particular a saturated C10-16 alkyl residue, for example a saturated C12-14 alkyl residue;
- X is NH, NR⁴ with C1-4 Alkyl residue R⁴, O or S,
- n a number from 1 to 10, preferably 2 to 5, in particular 3,
- x 0 or 1, preferably 1,

R², R³ are independently a C1-4 alkyl residue, potentially hydroxy substituted such as a hydroxyethyl, preferably a methyl.

m a number from 1 to 4, in particular 1, 2 or 3,

y 0 or 1 and

Y is COO, SO3, OPO(OR⁵)O or P(O)(OR⁵)O, whereby R⁵ is a hydrogen atom H or a C1-4 alkyl residue.

[0102] Preferred betaines are the alkyl betaines of the formula (Ia), the alkyl amido propyl betaine of the formula (Ib), the Sulfo betaines of the formula (Ic) and the Amido sulfobetaine of the formula (Id);

in which R¹1 as the same meaning as in formula I. Particularly preferred betaines are the Carbobetaine [wherein Y⁻ =COO⁻], in particular the Carbobetaine of the formula (Ia) and (Ib), more preferred are the Alkylamidobetaine of the formula (Ib).

[0103] Examples of suitable betaines and sulfobetaine are the following [designated in accordance with INCI]: Almondamidopropyl of betaines, Apricotam idopropyl betaines, Avocadamidopropyl of betaines, Babassuamidopropyl of betaines, Behenam idopropyl betaines, Behenyl of betaines, betaines, Canolam idopropyl betaines, Capryl/Capram idopropyl betaines, Carnitine, Cetyl of betaines, Cocamidoethyl of betaines, Cocam idopropyl betaines, Cocam idopropyl

- ³⁵ Hydroxysultaine, Coco betaines, Coco Hydroxysultaine, Coco/Oleam idopropyl betaines, Coco Sultaine, Decyl of betaines, Dihydroxyethyl Oleyl Glycinate, Dihydroxyethyl Soy Glycinate, Dihydroxyethyl Stearyl Glycinate, Dihydroxyethyl Tallow Glycinate, Dimethicone Propyl of PG-betaines, Erucam idopropyl Hydroxysultaine, Hydrogenated Tallow of betaines, Isostearam idopropyl betaines, Lauram idopropyl betaines, Lauryl of betaines, Lauryl Hydroxysultaine, Lauryl Sultaine, Milkam idopropyl betaines, Minkamidopropyl of betaines, Myristam idopropyl betaines, Myristyl of betaines,
- 40 Oleam idopropyl betaines, Oleam idopropyl Hydroxysultaine, Oleyl of betaines, Olivamidopropyl of betaines, Palmam idopropyl betaines, Palm itam idopropyl betaines, Palmitoyl Carnitine, Palm Kernelam idopropyl betaines, Polytetrafluoroethylene Acetoxypropyl of betaines, Ricinoleam idopropyl betaines, Sesam idopropyl betaines, Soyam idopropyl betaines, Stearam idopropyl betaines, Stearyl of betaines, Tallowam idopropyl betaines, Tallowam idopropyl Hydroxysultaine, Tallow of betaines, Tallow Dihydroxyethyl of betaines, Undecylenam idopropyl betaines and Wheat Germam
- ⁴⁵ idopropyl betaines. A preferred betaine is, for example, Cocoamidopropylbetaine.

Fatty Acid

[0104] Especially when in liquid form, preferably, the detergent composition comprises between 1.5% and 20%, more preferably between 2% and 15%, even more preferably between 3% and 10%, most preferably between 4% and 8% by weight of the liquid detergent composition of soap, preferably a fatty acid salt, more preferably an amine neutralized fatty acid salt, wherein preferably the amine is an alkanolamine more preferably selected from monoethanolamine, diethanolamine, triethanolamine or a mixture thereof, more preferably monoethanolamine.

55 Perfume

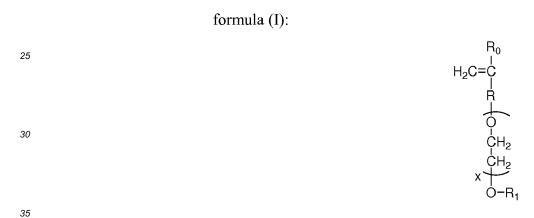
[0105] Preferred compositions of the invention comprise perfume. Typically the composition comprises a perfume that comprises one or more perfume raw materials, selected from the group as described in WO08/87497. However, any

perfume useful in a detergent may be used. A preferred method of incorporating perfume into the compositions of the invention is via an encapsulated perfume particle comprising either a water-soluble hydroxylic compound or melamineformaldehyde or modified polyvinyl alcohol. In one aspect the encapsulate comprises (a) an at least partially watersoluble solid matrix comprising one or more water-soluble hydroxylic compounds, preferably starch; and (b) a perfume

5 oil encapsulated by the solid matrix. In a further aspect the perfume may be pre-complexed with a polyamine, preferably a polyethylenimine so as to form a Schiff base.

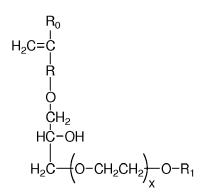
Polymers

- 10 [0106] The detergent composition may comprise one or more polymers for example for cleaning and/or care. Examples are optionally modified carboxymethylcellulose, poly (ethylene glycol), poly(vinyl alcohol), polycarboxylates such as polyacrylates, maleic/acrylic acid copolymers and lauryl methacrylate/acrylic acid co-polymers and carboxylate polymers. [0107] Suitable carboxylate polymers include maleate/acrylate random copolymer or polyacrylate homopolymer. The carboxylate polymer may be a polyacrylate homopolymer having a molecular weight of from 4.000 Da to 9,000 Da, or
- 15 from 6,000 Da to 9,000 Da. Other suitable carboxylate polymers are co-polymers of maleic acid and acrylic acid, and may have a molecular weight in the range of from 4,000 Da to 90,000 Da. [0108] Other suitable carboxylate polymers are co-polymers comprising: (i) from 50 to less than 98 wt% structural units derived from one or more monomers comprising carboxyl groups; (ii) from 1 to less than 49 wt% structural units derived from one or more monomers comprising sulfonate moieties; and (iii) from 1 to 49 wt% structural units derived
- 20 from one or more types of monomers selected from ether bond-containing monomers represented by formulas (I) and (II):



wherein in formula (I), R₀ represents a hydrogen atom or CH₃ group, R represents a CH₂ group, CH₂CH₂ group or single bond, X represents a number 0-5 provided X represents a number 1-5 when R is a single bond, and R1 is a hydrogen atom or C1 to C20 organic group;

40	formula (II)



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in formula (II), R0 represents a hydrogen atom or CH3 group, R represents a CH2 group, CH2CH2 group or single bond, X represents a number 0-5, and R_1 is a hydrogen atom or C1 to C20 organic group.

[0109] The composition may comprise one or more amphiphilic cleaning polymers such as the compound having the

following general structure: $bis((C_2H_5O)(C_2H_4O)_n)(CH_3)-N^+-C_xH_{2x}-N^+-(CH_3)-bis((C_2H_5O)(C_2H_4O)n)$, wherein n = from 20 to 30, and x = from 3 to 8, or sulphated or sulphonated variants thereof. In one aspect, this polymer is sulphated or sulphonated to provide a zwitterionic soil suspension polymer.

- [0110] The composition preferably comprises amphiphilic alkoxylated grease cleaning polymers which have balanced
- ⁵ hydrophilic and properties such that they remove grease particles from fabrics and surfaces. Preferred amphiphilic alkoxylated grease cleaning polymers comprise a core structure and a plurality of alkoxylate groups attached to that core structure. These may comprise alkoxylated polyalkylenimines, preferably having an inner polyethylene oxide block and an outer polypropylene oxide block. Typically these may be incorporated into the compositions of the invention in amounts of from 0.005 to 10 wt%, generally from 0.5 to 8 wt%.
- 10 [0111] Alkoxylated polycarboxylates such as those prepared from polyacrylates are useful herein to provide additional grease removal performance. Such materials are described in WO 91/08281 and PCT 90/01815. Chemically, these materials comprise polyacrylates having one ethoxy side-chain per every 7-8 acrylate units. The side-chains are of the formula -(CH₂CH₂O)_m (CH₂)_nCH₃ wherein m is 2-3 and n is 6-12. The side-chains are ester-linked to the polyacrylate "backbone" to provide a "comb" polymer type structure. The molecular weight can vary, but is typically in the range of
- about 2000 to about 50,000. Such alkoxylated polycarboxylates can comprise from about 0.05% to about 10%, by weight, of the compositions herein.
 [0112] The composition may comprise polyethylene glycol polymers and these may be particularly preferred in com-

positions comprising mixed surfactant systems. Suitable polyethylene glycol polymers include random graft co-polymers comprising: (i) hydrophilic backbone comprising polyethylene glycol; and (ii) side chain(s) selected from the group

- ²⁰ consisting of: C4-C25 alkyl group, polypropylene, polybutylene, vinyl ester of a saturated C1-C6 mono-carboxylic acid, C1-C6 alkyl ester of acrylic or methacrylic acid, and mixtures thereof. Suitable polyethylene glycol polymers have a polyethylene glycol backbone with random grafted polyvinyl acetate side chains. The average molecular weight of the polyethylene glycol backbone can be in the range of from 2,000 Da to 20,000 Da, or from 4,000 Da to 8,000 Da. The molecular weight ratio of the polyethylene glycol backbone to the polyvinyl acetate side chains can be in the range of
- from 1:1 to 1:5, or from 1:1.2 to 1:2. The average number of graft sites per ethylene oxide units can be less than 1, or less than 0.8, the average number of graft sites per ethylene oxide units can be in the range of from 0.5 to 0.9, or the average number of graft sites per ethylene oxide units can be in the range of from 0.1 to 0.5, or from 0.2 to 0.4. A suitable polyethylene glycol polymer is Sokalan HP22.
- [0113] Typically these polymers when present are each incorporated into the compositions of the invention in amounts from 0.005 to 10 wt%, more usually from 0.05 to 8 wt%.
- **[0114]** Preferably the composition comprises one or more carboxylate polymer, such as a maleate/acrylate random copolymer or polyacrylate homopolymer. In one aspect, the carboxylate polymer is a polyacrylate homopolymer having a molecular weight of from 4,000 Da to 9,000 Da, or from 6,000 Da to 9,000 Da. Typically these are incorporated into the compositions of the invention in amounts from 0.005 to 10 wt%, or from 0.05 to 8 wt%.
- ³⁵ [0115] Preferably the composition comprises one or more soil release polymers. [0116] Suitable soil release polymers are polyester soil release polymers such as Repel-o-tex polymers, including Repel-o-tex SF, SF-2 and SRP6 supplied by Rhodia. Other suitable soil release polymers include Texcare polymers, including Texcare SRA100, SRA300, SRN100, SRN170, SRN240, SRN260, SRN300 and SRN325 supplied by Clariant. Other suitable soil release polymers are Marloquest polymers, such as Marloquest SL supplied by Sasol.
- 40 [0117] Preferably the composition comprises one or more cellulosic polymer, including those selected from alkyl cellulose, alkyl alkoxyalkyl cellulose, carboxyalkyl cellulose, alkyl carboxyalkyl cellulose. Preferred cellulosic polymers are selected from the group comprising carboxymethyl cellulose, methyl cellulose, methyl hydroxyethyl cellulose, methyl carboxymethyl cellulose, and mixures thereof. In one aspect, the carboxymethyl cellulose has a degree of carboxymethyl substitution from 0.5 to 0.9 and a molecular weight from 100,000 Da to 300,000 Da.
- ⁴⁵ **[0118]** The composition preferably comprises a cationically-modified polysaccharide polymer. Preferably, the cationic polysaccharide polymer is selected from cationically modified hydroxyethyl cellulose, cationically modified hydroxypropyl cellulose, cationically and hydrophobically modified hydroxyethyl cellulose, cationically and hydrophobically modified hydroxyethyl cellulose, cationically and hydrophobically modified hydroxyethyl cellulose, cationically cellulose, cationically and hydrophobically modified hydroxyethyl cellulose, cationically and hydrophobically modified hydroxyethyl cellulose, cationically and hydroxyethyl cellulose, cationically and hydroxyethyl cellulose, cationically modified hydroxyethyl cellulose, cationically and hydroxyethy

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Amines

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[0119] The cleaning compositions described herein may contain an amine. The cleaning compositions may include from about 0.1% to about 10%, or from about 0.2% to about 5%, or from about 0.5% to about 4%, or from about 0.1 % to about 2%, by weight of the composition, of an amine. The amine can be subjected to protonation depending on the pH of the cleaning medium in which it is used. Non-limiting examples of amines include, but are not limited to, etheramines, cyclic amines, polyamines, oligoamines (e.g., triamines, diamines, pentamines, tetraamines), or combinations thereof. The compositions described herein may comprise an amine selected from the

group consisting of oligoamines, etheramines, cyclic amines, and combinations thereof. In some aspects, the amine is not an alkanolamine. In some aspects, the amine is not a polyalkyleneimine. Examples of suitable oligoamines include tetraethylenepentamine, triethylenetetraamine, diethylenetriamine, and mixtures thereof. Etheramines and cyclic amines may be particularly preferred.

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Fabric Shading Dye

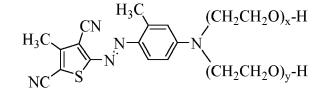
[0120] The composition may comprise a fabric shading agent. Suitable fabric shading agents include dyes, dye-clay conjugates, and pigments. Suitable dyes include small molecule dyes and polymeric dyes. Suitable small molecule dyes include small molecule dyes selected from the group consisting of dyes falling into the Colour Index (C.I.) classifications of Direct Blue, Direct Red, Direct Violet, Acid Blue, Acid Red, Acid Violet, Basic Blue, Basic Violet and Basic Red, or mixtures thereof. Preferered dyes include alkoxylated azothiophenes, Solvent Violet 13, Acid Violet 50 and Direct Violet 9. Particularly preferred dyes are polymeric dyes, particularly comprising polyalkoxy, most preferably polyethoxy groups, for example:

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wherein the index values x and y are independently selected from 1 to 10.

25 Dye Transfer Inhibitors

[0121] Suitable dye transfer inhibitors include polyamine N-oxide polymers, copolymers of N-vinylpyrrolidone and N-vinylimidazole, polyvinylpyrrolidone, polyvinyloxazolidone, polyvinylimidazole and mixtures thereof. Preferred are poly(vinyl pyrrolidone), poly(vinylpyridine betaine), poly(vinylpyridine N-oxide), poly(vinyl pyrrolidone-vinyl imidazole) and mixtures thereof. Suitable commercially available dye transfer inhibitors include PVP-K15 and K30 (Ashland), Sokalan® HP165, HP50, HP53, HP59, HP56K, HP56, HP66 (BASF), Chromabond® S-400, S403E and S-100 (Ashland).

Chelant

³⁵ [0122] The composition may comprise chelant for example selected from phosphonic, sulphonic, succinic and acetic chelants or mixtures thereof. Suitable examples include HEDP, DTPA, EDTA, MGDA, GLDA, EDDS and 4,5-dihydroxy-1,3-benzenedisulfonic acids and salts thereof.

Encapsulated Benefit Agent

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[0123] The composition may further comprise an encapsulated benefit agent. The encapsulated benefit may comprise a shell surrounding a core. The core may comprise a benefit agent. The benefit agent may comprise perfume raw materials.

[0124] The shell may comprise a material selected from the group consisting of aminoplast copolymer, an acrylic, an acrylate, and mixtures thereof. The aminoplast copolymer may be melamine-formaldehyde, urea-formaldehyde, crosslinked melamine formaldehyde, or mixtures thereof.

[0125] The shell may be coated with one or more materials, such as a polymer, that aids in the deposition and/or retention of the perfume microcapsule on the site that is treated with the composition disclosed herein. The polymer may be a cationic polymer selected from the group consisting of polysaccharides, cationically modified starch, cationically

⁵⁰ modified guar, polysiloxanes, poly diallyl dimethyl ammonium halides, copolymers of poly diallyl dimethyl ammonium chloride and vinyl pyrrolidone, acrylamides, imidazoles, imidazolinium halides, imidazolium halides, poly vinyl amine, copolymers of poly vinyl amine and N-vinyl formamide, and mixtures thereof.

[0126] The core may comprise a benefit agent. Suitable benefit agents include a material selected from the group consisting of perfume raw materials, silicone oils, waxes, hydrocarbons, higher fatty acids, essential oils, lipids, skin coolants, vitamins, sunscreens, antioxidants, glycerine, catalysts, bleach particles, silicon dioxide particles, malodor reducing agents, odor-controlling materials, chelating agents, antistatic agents, softening agents, insect and moth repelling agents, colorants, antioxidants, chelants, bodying agents, drape and form control agents, smoothness agents, wrinkle control agents, sanitization agents, disinfecting agents, germ control agents, mold control agents, mildew control

agents, antiviral agents, drying agents, stain resistance agents, soil release agents, fabric refreshing agents and freshness extending agents, chlorine bleach odor control agents, dye fixatives, dye transfer inhibitors, color maintenance agents, optical brighteners, color restoration/rejuvenation agents, anti-fading agents, whiteness enhancers, anti-abrasion agents, wear resistance agents, fabric integrity agents, anti-wear agents, anti-pilling agents, defoamers, anti-foaming agents,

- ⁵ UV protection agents, sun fade inhibitors, anti-allergenic agents, enzymes, water proofing agents, fabric comfort agents, shrinkage resistance agents, stretch resistance agents, stretch recovery agents, skin care agents, glycerin, and natural actives, antibacterial actives, antiperspirant actives, cationic polymers, dyes and mixtures thereof. The benefit agent may comprise perfume raw materials.
- [0127] The composition may comprise, based on total composition weight, from about 0.01% to about 10%, or from about 0.1% to about 5%, or from about 0.2% to about 1%, of encapsulated benefit agent. The encapsulated benefit agent may be friable and/or have a mean particle size of from about 10 microns to about 500 microns or from about 20 microns to about 200 microns.
 - [0128] Suitable encapsulated benefit agents may be obtained from Encapsys, LLC, of Appleton, Wisconsin USA.
 - [0129] Formaldehyde scavengers may also be used in or with such encapsulated benefit agents.
- ¹⁵ **[0130]** In a further preferred aspect of the invention, the composition is preferably liquid and comprises particulate benefit agents such as the encapsulated benefit agents mentioned above. The combination of the galactanase enzyme in addition to the plant fiber and particulate benefit agent has been found to provide the additional benefit of enhanced deposition of the particulate benefit agent. Thus, the present invention also provides a method of enhancing deposition of a particulate benefit agent comprising contacting a textile with an aqueous liquor comprising a composition defined
- ²⁰ herein, comprising a galactanase enzyme and a plant fiber and in addition a particulate benefit agent in a textile treatment step, preferably a laundering step, and optionally rinsing and drying the textile. In a preferred method, the aqueous liquor is an aqueous wash liquor. In a preferred method, the particulate benefit agent comprises an encapsulated perfume particle, most preferably comprising a shell which comprises a material selected from the group consisting of polymers or copolymers comprising acrylic acid and/or acrylates, and mixtures thereof.
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Methods of Making the Composition

Methods of Making the Composition

- ³⁰ **[0131]** The present invention relates to methods of making the compositions described herein. The compositions of the invention may be solid (for example granules or tablets) or liquid form. Preferably the compositions are in liquid form. They may be made by any process chosen by the formulator, including by a batch process, a continuous loop process, or combinations thereof.
- **[0132]** When in the form of a liquid, the compositions of the invention may be aqueous (typically above 2 wt% or even above 5 or 10 wt% total water, up to 90 or up to 80wt% or 70 wt% total water) or non-aqueous (typically below 2 wt% total water content). Typically the compositions of the invention will be in the form of an aqueous solution or uniform dispersion or suspension of optical brightener, DTI and optional additional adjunct materials, some of which may normally be in solid form, that have been combined with the normally liquid components of the composition, such as the liquid alcohol ethoxylate nonionic, the aqueous liquid carrier, and any other normally liquid optional ingredients. Such a solution,
- ⁴⁰ dispersion or suspension will be acceptably phase stable. When in the form of a liquid, the detergents of the invention preferably have viscosity from 1 to 1500 centipoises (1-1500 mPa*s), more preferably from 100 to 1000 centipoises (100-1000 mPa*s), and most preferably from 200 to 500 centipoises (200-500 mPa*s) at 20s-1 and 21°C. Viscosity can be determined by conventional methods. Viscosity may be measured using an AR 550 rheometer from TA instruments using a plate steel spindle at 40 mm diameter and a gap size of 500 µm. The high shear viscosity at 20s-1 and low shear
- ⁴⁵ viscosity at 0.05-1 can be obtained from a logarithmic shear rate sweep from 0.1-1 to 25-1 in 3 minutes time at 21C. The preferred rheology described therein may be achieved using internal existing structuring with detergent ingredients or by employing an external rheology modifier. More preferably the detergents, such as detergent liquid compositions have a high shear rate viscosity of from about 100 centipoise to 1500 centipoise, more preferably from 100 to 1000 cps. Unit Dose detergents, such as detergent liquid compositions have high shear rate viscosity of from 400 to 1000cps.
- ⁵⁰ Detergents such as laundry softening compositions typically have high shear rate viscosity of from 10 to 1000, more preferably from 10 to 800 cps, most preferably from 10 to 500 cps. Hand dishwashing compositions have high shear rate viscosity of from 300 to 4000 cps, more preferably 300 to 1000 cps.

[0133] The cleaning and/or treatment compositions in the form of a liquid herein can be prepared by combining the components thereof in any convenient order and by mixing, e.g., agitating, the resulting component combination to form a phase stable liquid detergent composition. In a process for preparing such compositions, a liquid matrix is formed containing at least a major proportion, or even substantially all, of the liquid components, e.g., nonionic surfactant, the non-surface active liquid carriers and other optional liquid components, with the liquid components being thoroughly admixed by imparting shear agitation to this liquid combination. For example, rapid stirring with a mechanical stirrer may

usefully be employed. While shear agitation is maintained, substantially all of any anionic surfactants and the solid form ingredients can be added. Agitation of the mixture is continued, and if necessary, can be increased at this point to form a solution or a uniform dispersion of insoluble solid phase particulates within the liquid phase. After some or all of the solid-form materials have been added to this agitated mixture, particles of any enzyme material to be included, e.g.,

- ⁵ enzyme granulates, are incorporated. As a variation of the composition preparation procedure hereinbefore described, one or more of the solid components may be added to the agitated mixture as a solution or slurry of particles premixed with a minor portion of one or more of the liquid components. After addition of all of the composition components, agitation of the mixture is continued for a period of time sufficient to form compositions having the requisite viscosity and phase stability characteristics. Frequently this will involve agitation for a period of from about 30 to 60 minutes.
- ¹⁰ **[0134]** The adjunct ingredients in the compositions of this invention may be incorporated into the composition as the product of the synthesis generating such components, either with or without an intermediate purification step. Where there is no purification step, commonly the mixture used will comprise the desired component or mixtures thereof (and percentages given herein relate to the weight percent of the component itself unless otherwise specified) and in addition unreacted starting materials and impurities formed from side reactions and/or incomplete reaction. For example, for an
- ¹⁵ ethoxylated or substituted component, the mixture will likely comprise different degrees of ethoxylation/substitution.

Method of Use

- [0135] The present invention relates to methods of using the cleaning compositions of the present invention to clean a surface, such as a textile. In general, the method includes mixing the cleaning composition as described herein with water to form an aqueous liquor and contacting a surface, preferably a textile, with the aqueous liquor in a laundering step. The target surface may include a greasy soil such as a body soil. The compositions herein, typically prepared as hereinbefore described, can be used to form aqueous washing/treatment solutions for use in the laundering/treatment of fabrics and/or hard surfaces. Generally, an effective amount of such a composition is added to water, for example in
- ²⁵ a conventional fabric automatic washing machine, to form such aqueous liquor laundering solutions. The aqueous liquor so formed is then contacted, typically under agitation, with the fabrics to be laundered/treated therewith. An effective amount of the cleaning composition herein added to water to form aqueous liquors for washing can comprise amounts sufficient to form from about 500 to 25,000 ppm, or from 500 to 15,000 ppm of composition in aqueousliquor, or from about 1,000 to 3,000 ppm of the cleaning compositions herein will be provided in aqueous liquor.
- ³⁰ [0136] Typically, the aqueous liquor is formed by contacting the detergent with wash water in such an amount so that the concentration of the cleaning composition in the aqueous liquor is from above 0.1 g/l to 5g/l, or from 1g/l, and to 4.5g/l, or to 4.0g/l, or to 3.5g/l, or to 3.0g/l, or to 2.5g/l, or even to 2.0g/l, or even to 1.5g/l. The method of laundering fabric or textile may be carried out in a top-loading or front-loading automatic washing machine, or can be used in a hand-wash laundry application. In these applications, the aqueous liquor formed and concentration of laundry detergent composition in the aqueous liquor is that of the main wash cycle. Any input of water during any optional rinsing step(s)
- [0137] The aqueous liquor may comprise 40 litres or less of water, or 30 litres or less, or 20 litres or less, or 10 litres or less, or 8 litres or less, or even 6 litres or less of water. The wash liquor may comprise from above 0 to 15 litres, or from 2 litres, and to 12 litres, or even to 8 litres of water. Typically from 0.01kg to 2kg of fabric per litre of aqueous liquor
- is dosed into said aqueous liquor. Typically from 0.01kg, or from 0.05kg, or from 0.07kg, or from 0.10kg, or from 0.15kg, or from 0.20kg, or from 0.25kg fabric per litre of aqueous liquor is dosed into said aqueous liquor. Optionally, 50g or less, or 45g or less, or 40g or less, or 35g or less, or 30g or less, or 25g or less, or 20g or less, or even 15g or less, or even 10g or less of the composition is contacted to water to form the aqueous liquor. When the wash solvent is water, employed at concentrations of from about 500 ppm to about 15,000 ppm in solution. When the wash solvent is water,
- ⁴⁵ the water temperature typically ranges from about 5 °C to about 90 °C and, when the situs comprises a fabric, the water to fabric ratio is typically from about 1:1 to about 30:1. Typically the aqueous liquor comprising the detergent of the invention has a pH of from 3 to 11.5.

[0138] In one aspect, such method comprises the steps of optionally washing and/or rinsing said surface or fabric, contacting said surface or fabric with any composition disclosed in this specification then optionally washing and/or rinsing said surface or fabric is disclosed, with an optional drying step.

- ⁵⁰ rinsing said surface or fabric is disclosed, with an optional drying step. [0139] Drying of such surfaces or fabrics may be accomplished by any one of the common means employed either in domestic or industrial settings: machine drying or open-air drying. The fabric may comprise any fabric capable of being laundered in normal consumer or institutional use conditions, and the invention is particularly suitable for synthetic textiles such as polyester and nylon and especially for treatment of mixed fabrics and/or fibres comprising synthetic and
- ⁵⁵ cellulosic fabrics and/or fibres. As examples of synthetic fabrics are polyester, nylon, these may be present in mixtures with cellulosic fibres, for example, polycotton fabrics. The solution typically has a pH of from 7 to 11, more usually 8 to 10.5. The compositions are typically employed at concentrations from 500 ppm to 5,000 ppm in solution. The water temperatures typically range from about 5 °C to about 90 °C. The water to fabric ratio is typically from about 1:1 to about

30:1.

Use of an Alkoxylated Phenol

5 [0140] The present invention further relates to a use of an alkoxylated polyaryl phenol and/or an alkoxylated polyalkyl phenol in a cleaning composition to enhance the stain-removal, whiteness, and/or malodor-reducing benefits of a galactanase enzyme.

EXAMPLES

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[0141] The following are illustrative examples of cleaning compositions according to the present invention and are not intended to be limiting.

Examples 1-7

[0142] Heavy Duty Liquid laundry detergent compositions.

	Ingradiante	1	2	3	4	5	6	7
20	Ingredients			C	% weight			
	AE _{1.8} S	6.77	5.16	1.36	1.30	-	-	-
	AE ₃ S	-	-	-	-	0.45	-	-
	LAS	0.86	2.06	2.72	0.68	0.95	1.56	3.55
25	HSAS	1.85	2.63	1.02	-	-	-	-
	AE9	6.32	9.85	10.20	7.92			
	AE8							35.45
30	AE7					8.40	12.44	
	C ₁₂₋₁₄ dimethyl Amine Oxide	0.30	0.73	0.23	0.37	-	-	-
	C ₁₂₋₁₈ Fatty Acid	0.80	1.90	0.60	0.99	1.20	-	15.00
	Citric Acid	2.50	3.96	1.88	1.98	0.90	2.50	0.60
35	Optical Brightener 1	1.00	0.80	0.10	0.30	0.05	0.50	0.001
	Optical Brightener 3	0.001	0.05	0.01	0.20	0.50	-	1.00
	Sodium formate	1.60	0.09	1.20	0.04	1.60	1.20	0.20
40	DTI 1	0.32	0.05	-	0.60	0.10	0.60	0.01
	DTI 2	0.32	0.10	0.60	0.60	0.05	0.40	0.20
	Sodium hydroxide	2.30	3.80	1.70	1.90	1.70	2.50	2.30
	Monoethanolamine	1.40	1.49	1.00	0.70	-	-	-
45	Diethylene glycol	5.50	-	4.10	-	-	-	-
	Chelant 1	0.15	0.15	0.11	0.07	0.50	0.11	0.80
	4-formyl-phenylboronic acid	-	-	-	-	0.05	0.02	0.01
50	Sodium tetraborate	1.43	1.50	1.10	0.75	-	1.07	-
	Ethanol	1.54	1.77	1.15	0.89	-	3.00	7.00
	Polymer 1	0.10	-	-	-	-	-	2.00
	Polymer 2	0.30	0.33	0.23	0.17	-	-	-
55	Polymer 3	-	-	-	-	-	-	0.80
	Polymer 4	0.80	0.81	0.60	0.40	1.00	1.00	-

	(continued)									
	Ingredients	1	2	3	4	5	6	7		
5	ingredients	% weight								
0	1,2-Propanediol	-	6.60	-	3.30	0.50	2.00	8.00		
	Structurant	0.10	-	-	-	-	-	0.10		
	Perfume	1.60	1.10	1.00	0.80	0.90	1.50	1.60		
10	Perfume encapsulate	0.10	0.05	0.01	0.02	0.10	0.05	0.10		
	Protease	0.80	0.60	0.70	0.90	0.70	0.60	1.50		
	Mannanase	0.07	0.05	-	0.06	0.04	0.045	-		
15	Amylase 1	0.30	-	0.30	0.10	-	0.40	0.10		
	Amylase 2	-	0.20	0.10	0.15	0.07	-	0.10		
	Xyloglucannase	0.20	0.10	-	-	0.05	0.05	0.20		
	Lipase	0.40	0.20	0.30	0.10	0.20	-	-		
20	Polishing enzyme	-	0.04	-	-	-	0.004	-		
	Galactanase	0.05	0.03	0.01	0.03	0.03	0.003	0.003		
	Dispersin B	-	-	-	0.05	0.03	0.001	0.001		
25	Acid Violet 50	0.05	-	-	-	-	-	0.005		
	Direct Violet 9	-	-	-	-	-	0.05	-		
30	Violet DD	-	0.035	0.02	0.037	0.04	-	-		
	Alkoxylated polyaryl/polyalkyl phenol	0.8	1.0	1.2	0.9	0.5	2.2	3.1		
	Water, dyes & minors				Balance					
	рН				8.2					

(continued)

Based on total cleaning and/or treatment composition weight. Enzyme levels are reported as raw material.

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Examples 8 to 18: Unit Dose Compositions.

[0143] These examples provide various formulations for unit dose laundry detergents. Compositions 8 to 12 comprise a single unit dose compartment. The film used to encapsulate the compositions is a polyvinyl-alcohol-based film.

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In an adjointe	8	9	10	11	12
Ingredients			% weight		
LAS	19.09	16.76	8.59	6.56	3.44
AE3S	1.91	0.74	0.18	0.46	0.07
AE7	14.00	17.50	26.33	28.08	31.59
Citric Acid	0.6	0.6	0.6	0.6	0.6
C12-15 Fatty Acid	14.8	14.8	14.8	14.8	14.8
Polymer 3	4.0	4.0	4.0	4.0	4.0
Chelant 2	1.2	1.2	1.2	1.2	1.2
Optical Brightener 1	0.20	0.25	0.01	0.01	0.50
Optical Brightener 2	0.20	-	0.25	0.03	0.01
Optical Brightener 3	0.18	0.09	0.30	0.01	-

EP 3	3 330	352	A1
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	Ingredients	8	9	10	11	12	
5	ingredients	% weight			:		
0	DTI 1	0.10	-	0.20	0.01	0.05	
	DTI 2	-	0.10	0.20	0.25	0.05	
	Glycerol	6.1	6.1	6.1	6.1	6.1	
10	Monoethanol amine	8.0	8.0	8.0	8.0	8.0	
	Tri-isopropanol amine	-	-	2.0	-	-	
	Tri-ethanol amine	-	2.0	-	-	-	
15	Cumene sulfonate	-	-	-	-	2.0	
15	Protease	0.80	0.60	0.07	1.00	1.50	
	Mannanase	0.07	0.05	0.05	-	-	
	Amylase 1	0.20	0.11	0.30	0.50	0.05	
20	Amylase 2	0.11	0.20	0.10	-	0.50	
	Polishing enzyme	0.005	0.05	-	-	-	
	Galactanase	0.005	0.05	0.005	0.010	0.005	
25	Dispersin B	0.010	0.05	-	0.005	-	
20	Cyclohexyl dimethanol	-	-	-	2.0	-	
	Acid violet 50	0.03	0.02				
	Violet DD			0.01	0.05	0.02	
30	Structurant	0.14	0.14	0.14	0.14	0.14	
	Perfume	1.9	1.9	1.9	1.9	1.9	
	Alkoxylated polyaryl/polyalkyl phenol	1.0	1.7	1.9	0.9	0.8	
35	Water and miscellaneous			To 100%			
	рН			7.5-8.2			

(continued)

[0144] Based on total cleaning and/or treatment composition weight. Enzyme levels are reported as raw material. [0145] In the following examples the unit dose has three compartments, but similar compositions can be made with 40 two, four or five compartments. The film used to encapsulate the compartments is polyvinyl alcohol.

	Base compositions Ingredients	13	14	15	16
			% we	eight	
45	HLAS	26.82	16.35	7.50	3.34
	AE7	17.88	16.35	22.50	30.06
	Citric Acid	0.5	0.7	0.6	0.5
50	C12-15 Fatty acid		6.0	11.0	13.0
	Polymer 1	2.9	0.1	-	-
	Polymer 3	1.1	5.1	2.5	4.2
	Cationic cellulose polymer	-	-	0.3	0.5
55	Polymer 6	-	1.5	0.3	0.2
	Chelant 2	1.1	2.0	0.6	1.5

	(contin	(continued)			
	Bass compositions in gradients	13	14	15	16
ī	Base compositions Ingredients		% we	eight	
	Optical Brightener 1	0.20	0.25	0.01	0.005
	Optical Brightener 3	0.18	0.09	0.30	0.005
	DTI 1	0.1	-	0.2	-
	DTI 2	-	0.1	0.2	-
	Glycerol	5.3	5.0	5.0	4.2
	Monoethanolamine	10.0	8.1	8.4	7.6
	Polyethylene glycol	-	-	2.5	3.0
	Potassium sulfite	0.2	0.3	0.5	0.7
	Protease	0.80	0.60	0.40	0.80
	Amylase 1	0.20	0.20	0.200	0.30
	Polishing enzyme	-	-	0.005	0.005
	Galactanase	0.05	0.010	0.005	0.005
	Dispersin B	-	0.010	0.010	0.010
	MgCl ₂	0.2	0.2	0.1	0.3
	Structurant	0.2	0.1	0.2	0.2
	Acid Violet 50	0.04	0.03	0.05	0.03
	Perfume / encapsulates	0.10	0.30	0.01	0.05
	Alkoxylated polyaryl/polyalkyl phenol	1.1	1.8	0.9	1.2
	Solvents and misc.		To 1	00%	
	рН		7.0-	-8.2	

(continued)

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Finishing compositions	17 18					
Compartment	А	В	С	А	В	С
Volume of each compartment	40 ml	5 ml	5 ml	40 ml	5 ml	5 m
Ingredients		Act	ive mate	erial in Wt.	%	
Perfume	1.6	1.6	1.6	1.6	1.6	1.
Violet DD	0	0.006	0	0	0.004	
TiO2	-	-	0.1	-		0.
Sodium Sulfite	0.4	0.4	0.4	0.3	0.3	0.
Polymer 5	-			2	-	
Hydrogenated castor oil	0.14	0.14	0.14	0.14	0.14	0.1
Base Composition 13, 14, 15 or 16	Add to 100%					

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[0146] Based on total cleaning and/or treatment composition weight, enzyme levels are reported as raw material.

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Examples 19 to 24

[0147] Granular laundry detergent compositions for hand washing or washing machines, typically top-loading washing

machines.

	In mus dis ut	19	20	21	22	23	24		
5	Ingredient	% weight							
	LAS	11.33	10.81	7.04	4.20	3.92	2.29		
	Quaternary ammonium	0.70	0.20	1.00	0.60	-	-		
	AE3S	0.51	0.49	0.32	-	0.08	0.10		
10	AE7	8.36	11.50	12.54	11.20	16.00	21.51		
	Sodium Tripolyphosphate	5.0	-	4.0	9.0	2.0	-		
	Zeolite A	-	1.0	-	1.0	4.0	1.0		
15	Sodium silicate 1.6R	7.0	5.0	2.0	3.0	3.0	5.0		
	Sodium carbonate	20.0	17.0	23.0	14.0	14.0	16.0		
	Polyacrylate MW 4500	1.0	0.6	1.0	1.0	1.5	1.0		
	Polymer 6	0.1	0.2	-	-	0.1	-		
20	Carboxymethyl cellulose	1.0	0.3	1.0	1.0	1.0	1.0		
	Acid Violet 50	0.05	-	0.02	-	0.04	-		
	Violet DD	-	0.03	-	0.03	-	0.03		
25	Protease 2	0.10	0.10	0.10	0.10	-	0.10		
	Amylase	0.03	-	0.03	0.03	0.03	0.03		
	Lipase	0.03	0.07	0.30	0.10	0.07	0.40		
	Polishing enzyme	0.002	-	0.05	-	0.02	-		
30	Galactanase	0.001	0.001	0.01	0.05	0.002	0.02		
	Dispersin B	0.001	0.001	0.05	-	0.001	-		
	Optical Brightener 1	0.200	0.001	0.300	0.650	0.050	0.001		
35	Optical Brightener 2	0.060	-	0.650	0.180	0.200	0.060		
	Optical Brightener 3	0.100	0.060	0.050	-	0.030	0.300		
	Chelant 1	0.60	0.80	0.60	0.25	0.60	0.60		
10	DTI 1	0.32	0.15	0.15	-	0.10	0.10		
40	DTI 2	0.32	0.15	0.30	0.30	0.10	0.20		
	Sodium Percarbonate	-	5.2	0.1	-	-	-		
	Sodium Perborate	4.4	-	3.85	2.09	0.78	3.63		
45	Nonanoyloxybenzensulfonate	1.9	0.0	1.66	0.0	0.33	0.75		
	Tetraacetylehtylenediamine	0.58	1.2	0.51	0.0	0.015	0.28		
	Photobleach	0.0030	0.0	0.0012	0.0030	0.0021	-		
50	S-ACMC	0.1	0.0	0.0	0.0	0.06	0.0		
00	Alkoxylated polyaryl/ polyalkyl phenol	1.9	1.1	2.0	1.8	2.2	1.2		
	Sulfate/Moisture			Bala	ince				

55 Examples 25-30

[0148] Granular laundry detergent compositions typically for front-loading automatic washing machines.

	Ingredient	25	26	27	28	29	30
5	LAS	6.08	5.05	4.27	3.24	2.30	1.09
	AE3S	-	0.90	0.21	0.18	-	0.06
	AS	0.34	-	-	-	-	-
10	AE7	4.28	5.95	6.72	7.98	9.20	10.35
	Quaternary ammonium	0.5	-	-	0.3	-	-
	Crystalline layered silicate	4.1	-	4.8	-	-	-
	Zeolite A	5.0	-	2.0	-	2.0	2.0
15	Citric acid	3.0	4.0	3.0	4.0	2.5	3.0
	Sodium carbonate	11.0	17.0	12.0	15.0	18.0	18.0
	Sodium silicate 2R	0.08	-	0.11	-	-	-
20	Optical Brightener 1	-	0.25	0.05	0.01	0.10	0.02
	Optical Brightener 2	-	-	0.25	0.20	0.01	0.08
	Optical Brightener 3	-	0.06	0.04	0.15	-	0.05
	DTI 1	0.08	-	0.04	-	0.10	0.01
25	DTI 2	0.08	-	0.04	0.10	0.10	0.02
	Soil release agent	0.75	0.72	0.71	0.72	-	-
	Acrylic /maleic acid copolymer	1.1	3.7	1.0	3.7	2.6	3.8
30	Carboxymethyl cellulose	0.2	1.4	0.2	1.4	1.0	0.5
	Protease 3	0.20	0.20	0.30	0.15	0.12	0.13
	Amylase 3	0.20	0.15	0.20	0.30	0.15	0.15
	Lipase	0.05	0.15	0.10	-	-	-
35	Amylase 2	0.03	0.07	-	-	0.05	0.05
	Cellulase 2	-	-	-	-	0.10	0.10
	Polishing enzyme	0.003	0.005	0.020	-	-	-
40	Galactanase	0.002	0.010	0.020	0.020	0.010	0.003
	Dispersin B	0.002	0.010	0.020	0.020	-	0.002
	Tetraacetylehtylenediamine	3.6	4.0	3.6	4.0	2.2	1.4
45	Sodium percabonate	13.0	13.2	13.0	13.2	16.0	14.0
45	Chelant 3	-	0.2	-	0.2	-	0.2
	Chelant 2	0.2	-	0.2	-	0.2	0.2
	MgSO ₄	-	0.42	-	0.42	-	0.4
50	Perfume	0.5	0.6	0.5	0.6	0.6	0.6
	Suds suppressor agglomerate	0.05	0.10	0.05	0.10	0.06	0.05
	Soap	0.45	0.45	0.45	0.45	-	-
55	Acid Violet 50	0.04	-	0.05	-	0.04	-
55	Violet DD	-	0.04	-	0.05	-	0.04
	S-ACMC	0.01	0.01	-	0.01	-	-

(continued)

Ingredient	25	26	27	28	29	30						
ingredient	% weight											
Direct Violet 9 (active)	-	-	0.0001	0.0001	-	-						
Alkoxylated polyaryl/ polyalkyl phenol	1.1	1.1	0.81	1.6	2.3	0.6						
Sulfate/ Water & Miscellaneous			Bala	ance								

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Examples 31-37: Heavy Duty Liquid laundry detergent compositions.

[0149]

15	Ingredients	31	32	33	34	35	36	37
	ingredients			C	% weight			
	AE _{1.8} S	6.77	5.16	1.36	1.30	-	-	-
20	AE ₃ S	-	-	-	-	0.45	-	-
	LAS	0.86	2.06	2.72	0.68	0.95	1.56	3.55
	HSAS	1.85	2.63	1.02	-	-	-	-
	AE9	6.32	9.85	10.20	7.92			
25	AE8							35.45
	AE7					8.40	12.44	
	C ₁₂₋₁₄ dimethyl Amine Oxide	0.30	0.73	0.23	0.37	-	-	-
30	C ₁₂₋₁₈ Fatty Acid	0.80	1.90	0.60	0.99	1.20	-	15.00
	Citric Acid	2.50	3.96	1.88	1.98	0.90	2.50	0.60
	Optical Brightener 1	1.00	0.80	0.10	0.30	0.05	0.50	0.001
	Optical Brightener 3	0.001	0.05	0.01	0.20	0.50	-	1.00
35	Sodium formate	1.60	0.09	1.20	0.04	1.60	1.20	0.20
	DTI 1	0.32	0.05	-	0.60	0.10	0.60	0.01
	DTI 2	0.32	0.10	0.60	0.60	0.05	0.40	0.20
40	Sodium hydroxide	2.30	3.80	1.70	1.90	1.70	2.50	2.30
	Monoethanolamine	1.40	1.49	1.00	0.70	-	-	-
	Diethylene glycol	5.50	-	4.10	-	-	-	-
	Chelant 1	0.15	0.15	0.11	0.07	0.50	0.11	0.80
45	4-formyl-phenylboronic acid	-	-	-	-	0.05	0.02	0.01
	Sodium tetraborate	1.43	1.50	1.10	0.75	-	1.07	-
	Ethanol	1.54	1.77	1.15	0.89	-	3.00	7.00
50	Polymer 1	0.10	-	-	-	-	-	2.00
	Polymer 2	0.30	0.33	0.23	0.17	-	-	-
	Polymer 3	-	-	-	-	-	-	0.80
	Polymer 4	0.80	0.81	0.60	0.40	1.00	1.00	-
55	1,2-Propanediol	-	6.60	-	3.30	0.50	2.00	8.00
	Structurant	0.10	-	-	-	-	-	0.10

		(00)	nunuea)					
	Ingredients	31	32	33	34	35	36	37
5	ingreatents			(% weight			
5	Perfume	1.60	1.10	1.00	0.80	0.90	1.50	1.60
	Perfume encapsulate	0.10	0.05	0.01	0.02	0.10	0.05	0.10
	Protease	0.80	0.60	0.70	0.90	0.70	0.60	1.50
10	Galactanase of any of SEQ ID Nos: 1-3	0.07	0.05	0.045	0.06	0.04	0.045	0.10
	Amylase 1	0.30	-	0.30	0.10	-	0.40	0.10
	Amylase 2	-	0.20	0.10	0.15	0.07	-	0.10
15	Xyloglucanase	0.20	0.10	-	-	0.05	0.05	0.20
15	Lipase	0.40	0.20	0.30	0.10	0.20	-	-
	Polishing enzyme	-	0.04	-	-	-	0.004	-
	Nuclease	0.05	0.03	0.01	0.03	0.03	0.003	0.003
20	Dispersin B	-	-	-	0.05	0.03	0.001	0.001
	Acid Violet 50	0.05	-	-	-	-	-	0.005
	Direct Violet 9	-	-	-	-	-	0.05	-
25	Violet DD	-	0.035	0.02	0.037	0.04	-	_
	Water insoluble plant fiber	0.2	0.1	0.3	0.25	1.2	1.5	0.25
	Dye control agent	-	0.3	-	0.5	-	0.3	-
	Alkoxylated polyaryl/ polyalkyl phenol	0.8	1.1	1.2	1.5	0.85	1.1	3.1
30	Water, dyes & minors				Balance			
	рН				8.2			

(continued)

[0150] Based on total cleaning and/or treatment composition weight. Unless indicated otherwise, enzyme levels are reported as raw material.

40	AE1.8S AE3S AE7 AE8 AE9 Alkoxylated polyaryl/ polyalkyl phenol	is C ₁₂₋₁₅ alkyl ethoxy sulfate with an average degree of ethoxylation of 1.8 is C ₁₂₋₁₅ alkyl ethoxy sulfate with an av degree of ethoxylation of 3 is C ₁₂₋₁₃ alcohol ethoxylate, with an average degree of ethoxylation of 7 is C ₁₂₋₁₃ alcohol ethoxylate, with an average degree of ethoxylation of 8 is C ₁₂₋₁₃ alcohol ethoxylate, with an average degree of ethoxylation of 9 is alkoxylated polyaryl/polyalkyl phenol in accordance with the invention, for example Emulsogen® TS160, Hostapal® BV conc., Sapogenat® T110 or Sa- pogenat® T139, all from Clariant
45	Amylase 1 Amylase 2 Amylase 3	is Stainzyme®, 15 mg active/g is Natalase®, 29 mg active/g is Stainzyme Plus®, 20 mg active/g,
50	AS Cellulase 2	is C ₁₂₋₁₄ alkylsulfate is Celluclean™, 15.6 mg active/g
50	Xyloglucanase Chelant 1 Chelant 2	is Whitezyme®, 20mg active/g is diethylene triamine pentaacetic acid is 1-hydroxyethane 1,1-diphosphonic acid
55	Chelant 3 Dispersin B DTI 1 DTI 2 Dye control agent	is sodium salt of ethylenediamine-N,N'-disuccinic acid, (S,S) isomer (EDDS) is a glycoside hydrolase, reported as 1000mg active/g is poly(4-vinylpyridine-1-oxide) (such as Chromabond S-403E®), is poly(1-vinylpyrrolidone-co-1-vinylimidazole) (such as Sokalan HP56®). Dye control agent in accordance with the invention, for example Suparex®

	HSAS LAS	O.IN (M1), Nylofixan® P (M2), Nylofixan® PM (M3), or Nylofixan® HF (M4) is mid-branched alkyl sulfate as disclosed in US 6,020,303 and US6,060,443 is linear alkylbenzenesulfonate having an average aliphatic carbon chain length C_{9} - C_{15} (HLAS is acid form).
5	Galactanase	is SEQ ID NO: 1, 2 and/or 3, as active protein.
	Lipase	is Lipex®, 18 mg active/g
	Mannanase	is Mannaway®, 25 mg active/g
	Optical Brightener 1	is disodium 4,4'-bis{[4-anilino-6-morpholino-s-triazin-2-yl]-amino}-2,2'-stilben- edisulfonate
10	Optical Brightener 2	is disodium 4,4'-bis-(2-sulfostyryl)biphenyl (sodium salt)
	Optical Brightener 3	is Optiblanc SPL10® from 3V Sigma
	Perfume encapsulate	is a core-shell melamine formaldehyde perfume microcapsules.
	Photobleach	is a sulfonated zinc phthalocyanine
	Polishing enzyme	is Para-nitrobenzyl esterase, reported as 1000mg active/g
15	Polyetheramine	as described in present disclosure.
	Polymer 1	is $bis((C_2H_5O)(C_2H_4O)n)(CH_3)-N^+-C_xH_{2x}-N^+-(CH_3)-bis((C_2H_5O)(C_2H_4O)n)$, wherein n = 20-30,x = 3 to 8 or sulphated or sulfonated variants thereof
	Polymer 2	is ethoxylated (EO ₁₅) tetraethylene pentamine
	Polymer 3	is ethoxylated polyethylenimine
20	Polymer 4	is ethoxylated hexamethylene diamine
	Polymer 5	is Acusol 305, provided by Rohm&Haas
	Polymer 6	is a polyethylene glycol polymer grafted with vinyl acetate side chains, provided by BASF.
	Protease	is Purafect Prime®, 40.6 mg active/g
25	Protease 2	is Savinase®, 32.89 mg active/g
	Protease 3	is Purafect®, 84 mg active/g
	Quaternary ammonium	is C ₁₂₋₁₄ Dimethylhydroxyethyl ammonium chloride
	S-ACMC	is Reactive Blue 19 Azo-CM-Cellulose provided by Megazyme
	Soil release agent	is Repel-o-tex® SF2
30	Structurant	is Hydrogenated Castor Oil
	Violet DD	is a thiophene azo dye provided by Milliken

[0151] The dimensions and values disclosed herein are not to be understood as being strictly limited to the exact numerical values recited. Instead, unless otherwise specified, each such dimension is intended to mean both the recited value and a functionally equivalent range surrounding that value. For example, a dimension disclosed as "40 mm" is intended to mean "about 40 mm."

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SEQUENCE LISTING

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	Gly Trp Gly Thr Ser Leu Ala Trp Trp Gly Asn Val Phe Gly Thr Arg 20 25 30
25	Asp Asp Phe Ala Asp Leu Phe Phe Thr Thr Lys Ser Val Thr Tyr Asn 35 40 45
30	Gly Thr Ser Leu Pro Gly Leu Gly Leu Asn Ile Ala Arg Tyr Asn Leu 50 55 60
35	Gly Ala Cys Ser Trp Asn Ala Val Asn Gly Glu Thr Met Val Lys Ser 65 70 75 80
	Pro Asn Ile Pro Ala Phe Lys Gln Ile Glu Gly Phe Trp Gln Asp Trp 85 90 95
40	Asn Asn Glu Asp Pro Thr Ser Ser Ala Trp Asp Trp Thr Ala Asp Ala 100 105 110
45	Thr Gln Arg Ala Met Leu Val Lys Ala Thr Gln Arg Gly Ala Val Thr 115 120 125
50	Glu Leu Phe Ala Asn Ser Pro Met Trp Trp Met Cys Tyr Asn His Asn 130 135 140
	Pro Ser Gly Ala Ala Asp Gly Gly Asn Asn Leu Gln Thr Trp Asn Tyr 145 150 155 160
55	Arg Gln His Ala Ser His Leu Ala Ala Val Ala Leu Tyr Ala Arg Thr 165 170 175

	Asn	Trp	Gly	Val 180	Asn	Phe	Ala	Thr	Val 185	Asp	Pro	Phe	Asn	Glu 190	Pro	Ala
5	Ser	Ser	Trp 195	Trp	Thr	Ala	Ser	Gly 200	Thr	Gln	Glu	Gly	Cys 205	His	Leu	Asp
10	Pro	Ala 210	Val	Gln	Ala	Ala	Val 215	Leu	Pro	Tyr	Met	Arg 220	Ser	Glu	Leu	Asp
	Lys 225	Arg	Gly	Leu	Thr	Gly 230	Val	Arg	Ile	Ser	Ala 235	Ser	Asp	Glu	Thr	Asn 240
15	Tyr	Asp	Thr	Ala	Arg 245	Ser	Thr	Trp	Ser	Ser 250	Phe	Gly	Ser	Ala	Thr 255	Lys
20	Ala	Leu	Val	Ser 260	Gln	Val	Asn	Val	His 265	Gly	Tyr	Gln	Gly	Thr 270	Gly	Gly
25	Arg	Arg	As p 275	Leu	Leu	Tyr	Thr	Asp 280	Val	Val	Thr	Thr	Ser 285	Gly	Lys	Lys
	Leu	Trp 290	Asn	Ser	Glu	Thr	Gly 295	Asp	Ser	Asp	Gly	Thr 300	Gly	Leu	Ser	Met
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	Asn Gln Arg Ala Met Leu Gln Lys Ala Lys Ala Asn Gly Ala Asn Ilo 115 120 125	Э
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	Cys Thr Ile Glu 35	Arg Ser Gly Al 40	a Asp Lys Arg Arg	Gln Glu Ser Leu 45
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25	Thr Trp P 145	ne Gln	Ile Thr 150	Lys Phe	Thr Gly	Ala Ala 155	Gly Pro	Tyr Cys 160
	Lys Ala L	_	Ser Asn 165	Asp Lys	Ser Val 170	Cys Asp	Lys Asn	Lys Asn 175
30	Ile Ala G	Ly Asp 180	Trp Gly	Phe Asp	Pro Ala 185	Lys Trp	Ala Tyr 190	Gln Tyr
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Gln Val Asn Asn Ala Cys Glu Ser Gln Ser Gly Thr Trp Ile Ser Pro Tyr Asp Asn Ala Ser Phe Thr Asn Ala Ser Ser Leu Asp Ile Asp His Met Val Pro Leu Lys Asn Ala Trp Ile Ser Gly Ala Ser Ser Trp Thr Thr Ala Gln Arg Glu Ala Leu Ala Asn Asp Val Ser Arg Pro Gln Leu Trp Ala Val Ser Ala Ser Ala Asn Arg Ser Lys Gly Asp Arg Ser Pro Asp Gln Trp Lys Pro Pro Leu Thr Ser Phe Tyr Cys Thr Tyr Ala Lys Ser Trp Ile Asp Val Lys Ser Phe Tyr Lys Leu Thr Ile Thr Ser Ala Glu Lys Thr Ala Leu Ser Ser Met Leu Asp Thr Cys <210> 9 <211> 361 <212> PRT <213> Aggregatibacter actinomycetemcomitans <400> 9 Asn Cys Cys Val Lys Gly Asn Ser Ile Tyr Pro Gln Lys Thr Ser Thr Lys Gln Thr Gly Leu Met Leu Asp Ile Ala Arg His Phe Tyr Ser Pro Glu Val Ile Lys Ser Phe Ile Asp Thr Ile Ser Leu Ser Gly Gly Asn Phe Leu His Leu His Phe Ser Asp His Glu Asn Tyr Ala Ile Glu Ser His Leu Leu Asn Gln Arg Ala Glu Asn Ala Val Gln Gly Lys Asp Gly Ile Tyr Ile Asn Pro Tyr Thr Gly Lys Pro Phe Leu Ser Tyr Arg Gln

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Claims

- 1. A cleaning composition comprising:
- a endo-beta-1,6-galactanase enzyme; and
 an alkoxylated phenol compound that is selected from the group consisting of an alkoxylated polyaryl phenol
 compound, an alkoxylated polyalkyl phenol compound, and mixtures thereof.
 - A cleaning composition according to claim 1, wherein the enzyme has an amino acid sequence having at least 60%, or at least 80%, or at least 90% or at least 95% identity with the amino acid sequence shown in SEQ ID NO:1, SEQ ID NO:2 or SEQ ID NO:3.
 - **3.** A cleaning composition according to claims 1 and 2, wherein the galactanase enzyme is selected from Glycoside Hydrolase Family 30.
- 35

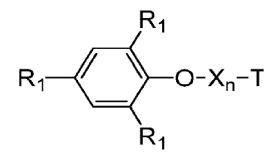
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- **4.** A cleaning composition according to any preceding claim, wherein the galactanase enzyme is obtainable from *Streptomyces davawensis, Trichoderma harzianum, Streptomyces avermitilis,* or a mixture thereof.
- **5.** A cleaning composition according to any preceding claim, wherein the composition further comprises a β-N-acetylglucosaminidase enzyme from E.C. 3.2.1.52, preferably an enzyme having at least 70% identity to SEQ ID NO:9.
- 6. A cleaning composition according to any preceding claim, wherein the alkoxylated phenol compound has a structure according to Formula I:
- 45

40

Formula I

50



55

wherein

each R1 is independently selected from linear of branched C3-C15 alkyl groups and C3-C15 aryl groups, X is selected from ethoxy or propoxy groups, n is from 2 to 70, and T is selected from H, SO₃⁻, COO⁻ and PO₃²⁻, preferably H and SO₃⁻.

- 5
- 7. A cleaning composition according to claim 6, wherein each R_1 is the same.
- 8. A cleaning composition according to any preceding claim, wherein the alkoxylated phenol is selected from the group consisting of:
- 10

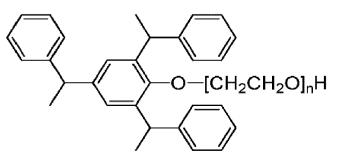
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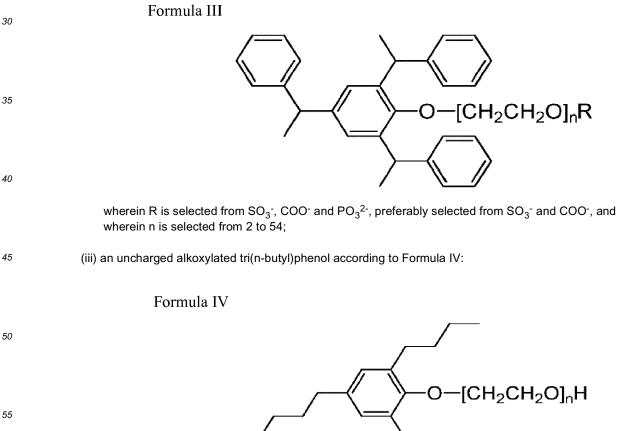
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(i) an uncharged alkoxylated tristyrylphenols according to Formula II:

Formula II

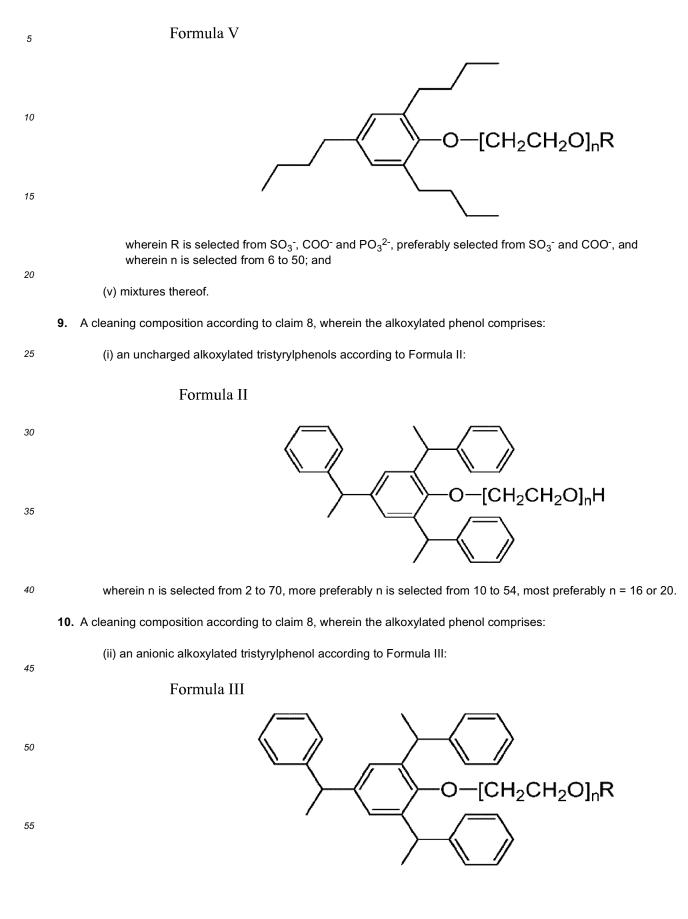


wherein n is selected from 2 to 70, more preferably n is selected from 10 to 54, most preferably n = 16 or 20; (ii) an anionic alkoxylated tristyrylphenol according to Formula III:



wherein n is selected from 2 to 50;

(iv) an anionic alkoxylated tri(n-butyl)phenol of the following structure according to Formula V:



wherein R is selected from SO₃⁻, COO⁻ and PO₃²⁻, preferably selected from SO₃⁻ and COO⁻, and wherein n is selected from 2 to 54.

- **11.** A cleaning composition according to claim 8, wherein the alkoxylated phenol comprises:
 - (iii) an uncharged alkoxylated tri(n-butyl)phenol according to Formula IV:

	Formula IV
10	
15	O_[CH₂CH₂O] _n H
20	wherein n is selected from 2 to 50.
	12. A cleaning composition according to claim 8, wherein the alkoxylated phenol comprises:
25	(iv) an anionic alkoxylated tri(n-butyl)phenol of the following structure according to Formula V:
	Formula V
30	
35	- $ -$
40	
	wherein R is selected from SO ₃ ⁻ , COO ⁻ and PO ₃ ²⁻ , preferably selected from SO ₃ ⁻ and COO ⁻ , and wherein n is selected from 6 to 50.

- **13.** A cleaning composition according to any preceding claim, wherein the alkoxylated phenol compound is an alkoxylated polyaryl phenol compound or an alkoxylated polyalkyl phenol compound.
 - **14.** A method of cleaning a surface, preferably a textile, comprising mixing the cleaning composition according to any preceding claim with water to form an aqueous liquor and contacting a surface, preferably a textile, with the aqueous liquor in a laundering step.

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15. The use of an endo-beta-1,6-galactanase enzyme and an alkoxylated polyaryl phenol and/or an alkoxylated polyalkyl phenol in a cleaning composition to enhance the stain-removal, whiteness, and/or malodor-reducing benefits.



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EP 3 330 352 A1

EUROPEAN SEARCH REPORT

Application Number EP 17 20 4768

		DOCUMENTS CONSIDE	ERED TO BE RELEVANT		
	Category	Citation of document with in of relevant passa		Relevant to claim	CLASSIFICATION OF THE APPLICATION (IPC)
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15	Y	WO 2016/110379 A1 (UUNILEVER NV [NL]; CO [US]) 14 July 2016 * claims; examples	ONOPCO INC DBĀ UNILEVER (2016-07-14)	1-15	
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30					TECHNICAL FIELDS SEARCHED (IPC) C11D
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1	The present search report has been drawn up for all claims				
(1) 07		Place of search Munich	Date of completion of the search 11 January 2018	Ver	^{Examiner} nier, Frédéric
0d) 28:00 55 55	Munich CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with anothe document of the same category A : technological background O : non-written disclosure P : intermediate document		E : earlier patent doct after the filing date er D : document cited in L : document cited for	T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons	
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11-01-2018

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