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(54) Titre : COMPOSITION DE DETERGENT POUR LAVE-VAISSELLE
(54) Title: AUTOMATIC DISHWASHING DETERGENT COMPOSITION

(57) Abrégé/Abstract:

A phosphate-free automatic dishwashing cleaning composition comprising: a) from 10% to 50% by weight of the composition of an organic complexing agent system comprising at least 10% by weight of the composition of methyl glycine diacetic acid; b) an amylase; and c) a protease wherein the protease is a variant having at least 90% identity with the amino acid sequence of SEQ ID NO:1 and comprising at least one amino acid substitution (using the SEQ ID NO:1 numbering) selected from the group consisting of S39E, S99R, S126A-D127E-F128G, N242D and mixtures thereof; or d) a protease wherein the protease is a variant having at least 90% identity with the amino acid sequence of SEQ ID NO:2 and comprising at least one amino acid substitution (using the SEQ ID NO:2 numbering) selected from the group consisting of X39E, X126A-X127E-X128G, X242D and mixtures thereof.

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(54) Title: AUTOMATIC DISHWASHING DETERGENT COMPOSITION

(57) Abstract: A phosphate-free automatic dishwashing cleaning composition comprising: a) from 10% to 50% by weight of the composition of an organic complexing agent system comprising at least 10% by weight of the composition of methyl glycine diacetic acid; b) an amylase; and c) a protease wherein the protease is a variant having at least 90% identity with the amino acid sequence of SEQ ID NO:1 and comprising at least one amino acid substitution (using the SEQ ID NO:1 numbering) selected from the group consisting of S39E, S99R, S126A-D127E-F128G, N242D and mixtures thereof; or d) a protease wherein the protease is a variant having at least 90% identity with the amino acid sequence of SEQ ID NO:2 and comprising at least one amino acid substitution (using the SEQ ID NO:2 numbering) selected from the group consisting of X39E, X126A-X127E-X128G, X242D and mixtures thereof.



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AUTOMATIC DISHWASHING DETERGENT COMPOSITION

FIELD OF THE INVENTION

The present invention is in the field of detergents. In particular, it relates to a phosphate-free automatic dishwashing detergent composition comprising a complexing agent system, an α -amylase and a specific protease. The composition provides improved cleaning under a plurality of conditions versus compositions comprising conventional proteases.

BACKGROUND OF INVENTION

There is a permanent desire to improve the performance of automatic dishwashing compositions and their environmental profile.

Due to environmental concerns phosphate is increasingly being replaced by biodegradable complexing agents. These complexing agents are usually strong builders and can negatively affect the performance of enzymes, in particular the strong complexing agents can negatively affect proteases by extracting the structural calcium metal ions of proteases. The proteases can be affected in product and/or in-use. For the toughest items, consumers would usually select hot, long automatic dishwashing cycles. In these cycles typically it is difficult to balance the need for excellent shine with the need for excellent performance driven by the stresses that these cycles create on enzymes. These stresses can be further exacerbated by high levels of bleach.

Automatic dishwashing compositions can be designed to have optimum performance under certain in-use conditions, for example a composition can be designed to have optimum performance in a hard water cycle, however a composition that has optimum performance in hard water might not have optimum performance in a soft water cycle and vice versa.

The object of the present invention is to provide a phosphate-free automatic dishwashing composition that provides better cleaning when used in hard water and preferably under different water hardnesses and levels of soil in the dishwashing load.

SUMMARY

Certain exemplary embodiments provide a phosphate-free automatic dishwashing cleaning composition comprising: a) from 10% to 50% by weight of the composition of an organic complexing agent system comprising at least 10% by weight of the composition of methyl glycine

diacetic acid; b) an amylase; and c) a protease wherein the protease is a variant having at least 90% identity with the amino acid sequence of SEQ ID NO:1 and comprising mutations S039E-S099R-S126A-D127E-F128G.

According to the first aspect of the present invention, there is provided a phosphate-free automatic dishwashing cleaning composition. The composition comprises a complexing agent system and a specific protease. The composition presents improved performance even when hard water is used in the automatic dishwashing process and the dishwashing program used is hot (the temperature is above 50°C) and long (the main wash is longer than 25 mins) and the dishware is heavily soiled. Even under these very stressed conditions the composition of the invention provides very good shine and cleaning, including very good removal of protein and starch based soils.

According to the second aspect of the invention there is provided a method of automatic dishwashing using hard water. According to the third aspect of the invention, there is provided the use of the composition of the invention for automatic dishwashing using hard water.

The elements of the composition of the invention described in connexion with the first aspect of the invention apply *mutatis mutandis* to the other aspects of the invention.

DETAILED DESCRIPTION OF THE INVENTION

The present invention encompasses an automatic dishwashing cleaning composition. The composition is phosphate-free and comprises a complexing agent system comprising a strong complexing agent, a protease and an amylase. The composition successfully provides excellent shine along with good protein and starch soil removal, including good removal of mixed protein/starch soils. The composition provides good cleaning, in particular on egg, baked cheese and starch. The invention also encompasses methods of automatic dishwashing with hard water.

By “hard” water is herein meant water having a hardness of more than 12 (205.7 ppm), preferably more than 15 (257.1 ppm) and more preferably more than 19 gpg (325.6 ppm). Grain per gallon (gpg) is a unit of water hardness defined as 1 grain (64.8 milligrams) of calcium carbonate dissolved in 1 US gallon of water (3.785412 L). It translates into 17.1 parts per million (ppm).

By “hot” cycle is herein understood a dishwashing program in which the main cycle is performed at a temperature above 50°C, preferably above 55°C.

By “long” cycle is herein understood a dishwashing program in which the main cycle has a duration of at least 25, preferably at least 30 and more preferably at least 35 minutes.

The composition of the invention comprises a variant protease, the variant proteases have a defined percentage of identity with respect to a reference protease (proteases of SEQ ID NO: 1 to 10).

The protease of the composition of the invention is herein sometimes referred to as “the protease of the invention”. The proteases having any of sequences ID NO:1 to 10 are herein sometimes referred to as “the reference protease”.

The relatedness between two amino acid sequences or between two nucleotide sequences is described by the parameter “sequence identity”.

The term “variant” means a protease comprising a mutation, i.e., a substitution, insertion, and/or deletion, at one or more (e.g., several) positions relative to the reference protease. A substitution means replacement of the amino acid occupying a position with a different amino acid; a deletion means removal of the amino acid occupying a position; and an insertion means adding an amino acid adjacent to and immediately following the amino acid occupying a position. The variants of the present invention have at least 60%, preferably at least 70%, more preferably at least 75% and especially 90% identity with the reference protease.

In preferred embodiments, the variant presents at least 90%, more preferably at least 92% identity with the protease of SEQ ID NO: 1. SEQ ID NO: 1 corresponds to *B. gibsonii*-clade subtilisin Bgi02446. In other embodiments, the variant presents at least 90%, more preferably at least 92% identity with the protease of SEQ ID NO: 2. SEQ ID NO: 2 corresponds to *B. lentus* subtilisin. In other embodiments, the variant presents at least 90%, more preferably at least 92% identity with one of the proteases of sequences SEQ ID NO: 3-10.

The term “wild-type” protease means a protease expressed by a naturally occurring microorganism, such as a bacterium, yeast, or filamentous fungus found in nature.

Enzyme related terminology

Nomenclature for amino acid modifications

In describing enzyme variants herein, the following nomenclature is used for ease of reference:

Original amino acid(s):position(s):substituted amino acid(s).

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According to this nomenclature, for instance the substitution of glutamic acid for glycine in position 195 is shown as G195E. A deletion of glycine in the same position is shown as G195*, and insertion of an additional amino acid residue such as lysine is shown as G195GK. Where a specific enzyme contains a "deletion" in comparison with other enzyme and an insertion is made in such a position this is indicated as *36D for insertion of an aspartic acid in position 36. Multiple mutations are separated by pluses, i.e.: S99G+V102N, representing mutations in positions 99 and 102 substituting serine and valine for glycine and asparagine, respectively. Where the amino acid in a position (*e.g.* 102) may be substituted by another amino acid selected from a group of amino acids, *e.g.* the group consisting of N and I, this will be indicated by V102N, I.

15 In all cases, the accepted IUPAC single letter or triple letter amino acid abbreviation is employed.

Protease Amino Acid Numbering

The numbering used in this patent is versus the sequences shown and not the BPN' numbering.

20 Amino acid identity

The relatedness between two amino acid sequences is described by the parameter "identity". For purposes of the present invention, the alignment of two amino acid sequences is determined by using the Needle program from the EMBOSS package (<http://emboss.org>) version 2.8.0. The Needle program implements the global alignment algorithm described in Needleman, S. B. and Wunsch, C. D. (1970) J. Mol. Biol. 48, 443-453. The substitution matrix used is BLOSUM62, gap opening penalty is 10, and gap extension penalty is 0.5.

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The degree of identity between an amino acid sequence of an enzyme used herein ("invention sequence") and a different amino acid sequence ("foreign sequence") is calculated as the number of exact matches in an alignment of the two sequences, divided by the length of the "invention sequence" or the length of the "foreign sequence", whichever is the shortest. The result is expressed in percent identity. An exact match occurs when the "invention sequence" and the

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“foreign sequence” have identical amino acid residues in the same positions of the overlap. The length of a sequence is the number of amino acid residues in the sequence.

The term “succinate based compound” and “succinic acid based compound” are used interchangeably herein.

5 As used herein, articles such as “a” and “an” when used in a claim, are understood to mean one or more of what is claimed or described.

Unless otherwise noted, all component or composition levels are in reference to the active portion of that 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
10 compositions.

All percentages and ratios are calculated by weight unless otherwise indicated. All percentages and ratios are calculated based on the total composition unless otherwise indicated.

The protease of the invention

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The variants of the present invention have at least 90%, more preferably at least 92% identity with the protease of SEQ ID NO: 1 or with the protease of SEQ ID NO: 2 or with one of the proteases of sequences ID NO: 3-10.

The protease of the invention is selected from:

- a) a variant having at least 90%, preferably at least 92% identity with the amino acid sequence of SEQ ID NO:1 and comprising at least one amino acid substitution (using the SEQ ID NO:1 numbering) selected from the group consisting of S39E, S99R, S126A-D127E-F128G, N242D and mixtures thereof; and
- b) a variant having at least 90%, preferably at least 92% identity with the amino acid sequence of SEQ ID NO:2 and comprising at least one amino acid substitution (using the SEQ ID NO:2 numbering) selected from the group consisting of X39E, X126A-X127E-X128G, X242D and mixtures thereof.

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Preferably, the variant comprises at least two, more preferably at least three amino acid substitutions (using the SEQ ID NO:1 numbering) selected from the group consisting of S39E, S99R, S126A-D127E-F128G and N242D, and preferably the variant comprises the following substitutions S39E, S99R, S126A-D127E-F128G and N242D.

Preferably, the variant further comprises at least one amino acid substitution (using the SEQ ID NO:1 numbering) selected from P54T, X114Q and X114C, preferably T114Q.

5 Preferably, the variant further comprises at least one and more preferably at least two and more preferably three amino acid substitution(s) (using the SEQ ID NO:1 numbering) selected from the group consisting of N74D; I80V, R, Y; N85S, C, D, I, K; E87D, C and M211L.

10 Preferably, the variant comprises at least 90% identity with the amino acid sequence of SEQ ID NO:1 and said variant further comprises at least one substitution (using the SEQ ID NO:1 numbering), preferably at least two substitutions and specially at least three substitutions selected from the group consisting of T3V, T9R, A15T, V66A, N74D, N85R, N97NE, N97AD, N97D/G, S99G/M, S101A, V102E/I, N116V/R, S126L, D127Q, F128A, G157S, Y161A, R164S, T188P, V199I, Q200C/E/I/K/T/V/W/L, Y203W, N212D, M216S/F, Q239R and T249R.

15 Especially preferred variants for use in the composition of the invention are selected from the group consisting of variants having at least 90%, more preferably at least 92% identity with the amino acid sequence SEQ ID:1 and comprising substitutions (using the SEQ ID NO:1 numbering) selected from the group consisting of:

- 1) T056Y-S099R-S126A-D127E-F128G
- 20 2) S039E-I080V-S099R-S126A-D127E-F128G-M211L
- 3) S039E-P054T-S099R-S126A-D127E-F128G-M211L
- 4) S039E-I043V-S099R-S126A-D127E-F128G-M211L
- 5) S039E-N042R-S099R-S126A-D127E-F128G
- 6) S039E-I080V-S099R-S126A-D127E-F128G
- 25 7) S039E-S099R-S126A-D127E-F128G-M211L
- 8) S039E-N085S-S099R-S126A-D127E-F128G-M211L
- 9) S039E-T056Y-S099R-S126A-D127E-F128G-M211L
- 10) S039E-A047V-S099R-S126A-D127E-F128G-M211L
- 11) S039E-S099R-S126A-D127E-F128G-Y203W
- 30 12) A037T-S039E-S099R-S126A-D127E-F128G-M211L
- 13) S039E-S099R-S126A-D127E-F128G-V199I
- 14) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G

- 15) S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 16) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G
- 5 17) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G
- 18) S039E-I043V-S099R-S126A-D127E-F128G
- 19) S039E-S099R-S126A-D127E-F128G-N253D
- 20) T009E-S039E-S099R-S126A-D127E-F128G
- 10 21) S039E-S099R-S126A-D127E-F128G-S255W
- 22) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-S099R-S126A-D127E-F128G-N242D
- 23) A037T-S039E-I043V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 15 24) S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 25) S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 26) S039E-T056Y-S099R-S126A-D127E-F128G
- 20 27) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G
- 28) A037T-S039E-I043V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 29) A037T-S039E-I043V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 25 30) A037T-S039E-I043V-A047V-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 31) A037T-S039E-I043V-A047V-P054T-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 30 32) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-N242D
- 33) A037T-S039E-I043V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D

- 34) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-N242D
- 35) S039E-N085S-S099R-S126A-D127E-F128G
- 5 36) A037T-S039E-I043V-A047V-P054T-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 37) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-S099R-T114Q-S126A-D127E-F128G-N242D
- 38) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-N242D
- 10 39) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S126A-D127E-F128G-N242D
- 40) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-F128G-N242D
- 41) A037T-S039E-I043V-A047V-T056Y-I080V-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 15 42) S039E-S099R-T114Q-S126A-D127E-F128G
- 43) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-N242D
- 44) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 20 45) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-D127E-F128G-N242D
- 46) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-S099R-T114Q-S126A-D127E-F128G-N242D
- 25 47) S039E-A047V-S099R-S126A-D127E-F128G
- 48) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-S099R-S126A-D127E-F128G-N242D
- 49) A037T-S039E-I043V-A047V-T056Y-S099R-T114Q-S126A-D127E-F128G
- 50) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-E087D-S099R-S126A-D127E-F128G-N242D
- 30 51) S039E-P054T-S099R-S126A-D127E-F128G
- 52) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D

- 53) A037T-S039E-I043V-A047V-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 54) A037T-S039E-I043V-A047V-T056Y-I080V-E087D-S099R-S126A-D127E-F128G-N242D
- 5 55) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-D127E-F128G-N242D
- 56) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-D127E-F128G-N242D
- 57) S039E-E087D-S099R-S126A-D127E-F128G
- 10 58) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-D127E-F128G-N242D
- 59) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-F128G-N242D
- 60) A037T-S039E-S099R-S126A-D127E-F128G
- 15 61) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 62) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 63) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-F128G-N242D
- 20 64) A037T-S039E-I043V-A047V-T056Y-S099R-S126A-D127E-F128G
- 65) A037T-S039E-I043V-A047V-P054T-T056Y-S099R-T114Q-S126A-D127E-F128G
- 66) A037T-S039E-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 25 67) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-F128G-N242D
- 68) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-T114Q-S126A-D127E-F128G-N242D
- 30 69) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S126A-D127E-F128G-N242D
- 70) A037T-S039E-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 71) S039E-S099R-S126A-D127E-F128G

- 72) A037T-S039E-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 73) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-T114Q-S126A-D127E-F128G-N242D
- 5 74) A037T-S039E-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 75) S039E-S099R-S126A-D127E-F128G-Q256E
- 76) S039E-S099R-S126A-D127E-F128G-N242D
- 77) S039E-S099R-S126A-D127E-F128G-Q200L
- 10 78) A037T-S039E-I043V-A047V-P054T-T056Y-S099R-T114Q-S126A-D127E-F128G-N242D
- 79) A037T-S039E-I043V-A047V-P054T-T056Y-S099R-S126A-D127E-F128G-N242D
- 80) S039E-N074D-I080V-S099R-S126A-D127E-F128G
- 15 81) A037T-S039E-I043V-A047V-P054T-T056Y-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 82) A037T-S039E-I043V-A047V-T056Y-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 83) A037T-S039E-I043V-A047V
- 20 84) A037T-S039E-I043V-A047V-N242D
- 85) S039E-P054T-N074D-S099R-S126A-D127E-F128G
- 86) S039E-A047V-N074D-S099R-S126A-D127E-F128G
- 87) A037T-S039E-I043V-A047V-P054T-T056Y-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 25 88) S039E-N074D-S099R-T114Q-S126A-D127E-F128G
- 89) S039E-N074D-N085S-S099R-S126A-D127E-F128G
- 90) A037T-S039E-I043V-A047V-T056Y-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 91) S039E-N074D-E087D-S099R-S126A-D127E-F128G
- 30 92) S039E-I043V-N074D-S099R-S126A-D127E-F128G
- 93) S039E-T056Y-N074D-S099R-S126A-D127E-F128G
- 94) S039E-N074D-S099R-S126A-D127E-F128G-N242D

Especially preferred variants for use in the composition of the invention are selected from the group consisting of variants having at least 90%, more preferably at least 92% identity with the amino acid sequence SEQ ID:1 and comprising substitutions (using the SEQ ID NO:1 numbering) selected from the group consisting of:

- 5 (a) A37T-S39E-I43V-A47V-T56Y-I80V-N85S-E87D-S99R-T114Q-S126D-D127S-F128A-N242D;
- (b) A37T-S39E-I43V-A47V-T56Y-I80V-N85S-E87D-T114Q-S126G-D127T-F128E-N242D;
- (c) A37T-S39E-I43V-A47V-T56Y-I80V-N85S-E87D-S99R-T114Q-D127A-F128C-N242D;
- 10 (d) A37T-S39E-I43V-A47V-P54T-T56Y-I80V-N85S-E87D-S99E-T114Q-S126T-F128A-N242D;
- (e) A37T-S39E-I43V-A47V-P54T-T56Y-I80V-N85S-E87D-S99R-T114Q-S126V-D127A-F128T-N242D;
- 15 (f) A37T-S39E-I43V-A47V-P54T-T56Y-I80V-N85S-E87D-S99R-T114Q-F128E-N242D;
- (g) A37T-S39E-I43V-A47V-T56Y-I80V-N85S-E87D-S99E-T114Q-D127E-F128G-N242D;
- (h) A37T-S39E-I43V-A47V-P54T-T56Y-I80V-N85S-E87D-S99R-T114Q-S126T-D127E-F128A-N242D;
- 20 (i) A37T-S39E-I43V-A47V-T56Y-I80V-N85S-E87D-T114Q-D127P-F128E-N242D;
- (j) A37T-S39E-I43V-A47V-P54T-T56Y-I80V-N85S-E87D-S99R-T114Q-S126G-D127E-F128D-N242D;
- 25 (k) A37T-S39E-I43V-A47V-T56Y-I80V-N85S-E87D-S99R-T114Q-F128E-N242D;
- (l) A37T-S39E-I43V-A47V-T56Y-I80V-N85S-E87D-S99R-T114Q-S126T-D127E-F128E-N242D;
- (m) A37T-S39E-I43V-A47V-P54T-T56Y-I80V-N85S-E87D-S99R-T114Q-D127G-F128P-N242D; and
- 30 (n) A37T-S39E-I43V-A47V-P54T-T56Y-I80V-N85S-E87D-S99R-T114Q-D127G-F128E-N242D.

Preferably, the variant has at least 90% identity with SEQ ID NO:2 and comprises at least two, more preferably at least three amino acid substitutions (using the SEQ ID NO:2 numbering)

selected from the group consisting of X39E, X126A-X127E-X128G and X242D, preferably P39E, S126A-P127E-S128G and N242D and preferably the variant comprises the following substitutions P39E, S126A-P127E-S128G and N242D.

- 5 Preferably, the variant further comprises at least one amino acid substitution (using the SEQ ID NO:2 numbering) selected from P54T, N114Q and N114C.

Preferably, the variant further comprises at least one and more preferably at least two and more preferably three amino acid substitution(s) (using the SEQ ID NO:2 numbering) selected from the group consisting of N74D; L80V, R, Y; S85S, C, D, I, K; and E87D, C.

- Preferably, the variant has at least 90% identity with the amino acid sequence of a parent protease said parent protease having the amino acid sequence of SEQ ID NO:2 and said variant further
10 comprises at least one substitution (using the SEQ ID NO:2 numbering), preferably at least two substitutions and specially at least three substitutions selected from the group consisting of S3V, S9R, A15T, V66A, N74D, S85N/R, S97SE, S97AD, S97D/G, S99G/M, S101A, V102E/I, G116V/R, S126F, P127Q, S128A, G157S, Y161A, R164S, A188P, V199I, Q200C/E/I/K/T/V/W/L, Y203W, N212D, M216S/F, Q239R and T249R.

15

Especially preferred variants for use in the composition of the invention are selected from the group consisting of variants having at least 90%, more preferably at least 92% identity with the amino acid sequence SEQ ID NO:2 and comprising substitutions (using the SEQ ID NO:2 numbering) selected from the group consisting of:

20

- 1) T056Y-S099R-S126A-P127E-S128G
- 2) P039E-L080V-S099R-S126A-P127E-S128G-M211L
- 3) P039E-P054T-S099R-S126A-P127E-S128G-M211L
- 4) P039E-I043V-S099R-S126A-P127E-S128G-M211L
- 25 5) P039E-N042R-S099R-S126A-P127E-S128G
- 6) P039E-L080V-S099R-S126A-P127E-S128G
- 7) P039E-S099R-S126A-P127E-S128G-M211L
- 8) P039E--S099R-S126A-P127E-S128G-M211L
- 9) P039E-T056Y-S099R-S126A-P127E-S128G-M211L
- 30 10) P039E-A047V-S099R-S126A-P127E-S128G-M211L

- 11) P039E-S099R-S126A-P127E-S128G-Y203W
- 12) P039E-S099R-S126A-P127E-S128G-M211L
- 13) P039E-S099R-S126A-P127E-S128G-V199I
- 14) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S099R-N114Q-S126A-P127E-
5 S128G
- 15) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S099R-S126A-P127E-S128G-
N242D
- 16) P039E-I043V-A047V-T056Y-L080V--E087D-S099R-N114Q-S126A-P127E-S128G
- 17) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S099R-S126A-P127E-S128G
- 18) P039E-I043V-S099R-S126A-P127E-S128G
- 19) P039E-S099R-S126A-P127E-S128G-N253D
- 20) T009E-P039E-S099R-S126A-P127E-S128G
- 21) P039E-S099R-S126A-P127E-S128G-S255W
- 22) P039E-I043V-A047V-P054T-T056Y-L080V--S099R-S126A-P127E-S128G-N242D
- 23) P039E-I043V-P054T-T056Y-L080V--E087D-S099R-S126A-P127E-S128G-N242D
- 24) P039E-I043V-A047V-T056Y-L080V--E087D-S099R-S126A-P127E-S128G-N242D
- 25) P039E-I043V-A047V-T056Y-L080V--E087D-S099R-N114Q-S126A-P127E-S128G-
N242D
- 26) P039E-T056Y-S099R-S126A-P127E-S128G
- 27) P039E-I043V-A047V-T056Y-L080V--E087D-S099R-S126A-P127E-S128G
- 28) P039E-I043V-P054T-T056Y-L080V--E087D-S099R-N114Q-S126A-P127E-S128G-
N242D
- 29) P039E-I043V-T056Y-L080V--E087D-S099R-N114Q-S126A-P127E-S128G-N242D
- 30) P039E-I043V-A047V-L080V--E087D-S099R-N114Q-S126A-P127E-S128G-N242D
- 31) P039E-I043V-A047V-P054T-L080V--E087D-S099R-S126A-P127E-S128G-N242D
- 32) P039E-I043V-A047V-T056Y-L080V--E087D-S099R-S126A-P127E-N242D
- 33) P039E-I043V-T056Y-L080V--E087D-S099R-S126A-P127E-S128G-N242D
- 34) P039E-I043V-A047V-T056Y-L080V--E087D-S099R-N114Q-S126A-P127E-N242D
- 35) P039E--S099R-S126A-P127E-S128G
- 36) P039E-I043V-A047V-P054T-L080V--E087D-S099R-N114Q-S126A-P127E-S128G-
N242D
- 37) P039E-I043V-A047V-P054T-T056Y-L080V--S099R-N114Q-S126A-P127E-S128G-
N242D

- 38) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S099R-N114Q-S126A-P127E-N242D
- 39) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S126A-P127E-S128G-N242D
- 40) P039E-I043V-A047V-T056Y-L080V--E087D-S099R-S126A-S128G-N242D
- 5 41) P039E-I043V-A047V-T056Y-L080V-E087D-S099R-N114Q-S126A-P127E-S128G-N242D
- 42) P039E-S099R-N114Q-S126A-P127E-S128G
- 43) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S099R-S126A-P127E-N242D
- 44) P039E-I043V-A047V-T056Y-L080V--E087D-S099R-S126A-P127E-S128G-N242D
- 10 45) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S099R-N114Q-P127E-S128G-N242D
- 46) P039E-I043V-A047V-T056Y-L080V--S099R-N114Q-S126A-P127E-S128G-N242D
- 47) P039E-A047V-S099R-S126A-P127E-S128G
- 48) P039E-I043V-A047V-T056Y-L080V--S099R-S126A-P127E-S128G-N242D
- 15 49) P039E-I043V-A047V-T056Y-S099R-N114Q-S126A-P127E-S128G
- 50) P039E-I043V-A047V-P054T-T056Y-L080V-E087D-S099R-S126A-P127E-S128G-N242D
- 51) P039E-P054T-S099R-S126A-P127E-S128G
- 52) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S099R-N114Q-S126A-P127E-S128G-N242D
- 20 53) P039E-I043V-A047V-L080V--E087D-S099R-S126A-P127E-S128G-N242D
- 54) P039E-I043V-A047V-T056Y-L080V-E087D-S099R-S126A-P127E-S128G-N242D
- 55) P039E-I043V-A047V-T056Y-L080V--E087D-S099R-N114Q-P127E-S128G-N242D
- 56) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S099R-P127E-S128G-N242D
- 25 57) P039E-E087D-S099R-S126A-P127E-S128G
- 58) P039E-I043V-A047V-T056Y-L080V--E087D-S099R-P127E-S128G-N242D
- 59) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S099R-N114Q-S126A-S128G-N242D
- 60) P039E-S099R-S126A-P127E-S128G
- 30 61) P039E-I043V-A047V-P054T-T056Y-L080V-E087D-S099R-N114Q-S126A-P127E-S128G-N242D
- 62) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S099R-S126A-P127E-S128G-N242D
- 63) P039E-I043V-A047V-T056Y-L080V--E087D-S099R-N114Q-S126A-S128G-N242D

- 64) P039E-I043V-A047V-T056Y-S099R-S126A-P127E-S128G
- 65) P039E-I043V-A047V-P054T-T056Y-S099R-N114Q-S126A-P127E-S128G
- 66) P039E-A047V-P054T-T056Y-L080V--E087D-S099R-S126A-P127E-S128G-N242D
- 67) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-S099R-S126A-S128G-N242D
- 5 68) P039E-I043V-A047V-P054T-T056Y-L080V--E087D-N114Q-S126A-P127E-S128G-N242D
- 69) P039E-I043V-A047V-T056Y-L080V--E087D-S126A-P127E-S128G-N242D
- 70) P039E-A047V-T056Y-L080V--E087D-S099R-N114Q-S126A-P127E-S128G-N242D
- 71) P039E-S099R-S126A-P127E-S128G
- 10 72) P039E-A047V-P054T-T056Y-L080V--E087D-S099R-N114Q-S126A-P127E-S128G-N242D
- 73) P039E-I043V-A047V-T056Y-L080V--E087D-N114Q-S126A-P127E-S128G-N242D
- 74) P039E-A047V-T056Y-L080V--E087D-S099R-S126A-P127E-S128G-N242D
- 75) P039E-S099R-S126A-P127E-S128G-Q256E
- 15 76) P039E-S099R-S126A-P127E-S128G-N242D
- 77) P039E-S099R-S126A-P127E-S128G-Q200L
- 78) P039E-I043V-A047V-P054T-T056Y-S099R-N114Q-S126A-P127E-S128G-N242D
- 79) P039E-I043V-A047V-P054T-T056Y-S099R-S126A-P127E-S128G-N242D
- 80) P039E-N074D-L080V-S099R-S126A-P127E-S128G
- 20 81) P039E-I043V-A047V-P054T-T056Y--E087D-S099R-S126A-P127E-S128G-N242D
- 82) P039E-I043V-A047V-T056Y--E087D-S099R-N114Q-S126A-P127E-S128G-N242D
- 83) P039E-I043V-A047V
- 84) P039E-I043V-A047V-N242D
- 85) P039E-P054T-N074D-S099R-S126A-P127E-S128G
- 25 86) P039E-A047V-N074D-S099R-S126A-P127E-S128G
- 87) P039E-I043V-A047V-P054T-T056Y--E087D-S099R-N114Q-S126A-P127E-S128G-N242D
- 88) P039E-N074D-S099R-N114Q-S126A-P127E-S128G
- 89) P039E-N074D--S099R-S126A-P127E-S128G
- 30 90) P039E-I043V-A047V-T056Y--E087D-S099R-S126A-P127E-S128G-N242D
- 91) P039E-N074D-E087D-S099R-S126A-P127E-S128G
- 92) P039E-I043V-N074D-S099R-S126A-P127E-S128G
- 93) P039E-T056Y-N074D-S099R-S126A-P127E-S128G
- 94) P039E-N074D-S099R-S126A-P127E-S128G-N242D

Especially preferred variants for use in the composition of the invention are selected from the group consisting of variants having at least 90%, more preferably at least 92% identity with the amino acid sequence SEQ ID NO:2 and comprising substitutions (using the SEQ ID NO:2 numbering) selected from the group consisting of:

- 5 (i) P39E-I43V-A47V-T56Y-L80V-E87D-S99R-N114Q -S126D-P127S-S128A-N242D;
- (ii) P39E-I43V-A47V-T56Y-L80V-E87D- N114Q -S126G-P127T-S128E-N242D;
- (iii) P39E-I43V-A47V-T56Y-L80V- E87D-S99R-N114Q -P127A-S128C-N242D;
- (iv) P39E-I43V-A47V-P54T-T56Y-L80V- E87D-S99E-N114Q -S126T-P127D-S128A-N242D;
- 10 (v) P39E-I43V-A47V-P54T-T56Y-L80V- E87D-S99R-N114Q -S126V-P127A-S128T-N242D;
- (vi) P39E-I43V-A47V-P54T-T56Y-L80V-E87D-S99R-N114Q --S128E-N242D;
- (vii) P39E-I43V-A47V-T56Y-L80V- E87D-S99E-N114Q -P127E-S128G-N242D;
- (viii) P39E-I43V-A47V-P54T-T56Y-L80V-E87D-S99R-N114Q -S126T-P127E-S128A-
- 15 N242D;
- (ix) P39E-I43V-A47V-T56Y-L80V-E87D-S128E-N242D;
- (x) P39E-I43V-A47V-P54T-T56Y-L80V-E87D-S99R-N114Q -S126G-P127E-S128D-N242D;
- (xi) P39E-I43V-A47V-T56Y-L80V- E87D-S99R-N114Q -P127D-S128E-N242D;
- 20 (xii) P39E-I43V-A47V-T56Y-L80V- E87D-S99R-N114Q -S126T-P127E-S128E-N242D;
- (xiii) P39E-I43V-A47V-P54T-T56Y-L80V- E87D-S99R-N114Q -S126A-P127E-S128G-N242D;
- (xiv) P39E-I43V-A47V-P54T-T56Y-L80V-E87D-S99R-N114Q -P127G-S128P-N242D;
- (xv) P39E-I43V-A47V-P54T-T56Y-L80V- E87D-S99R-N114Q -P127G-S128E-N242D;
- 25 (xvi) P39E-I43V-A47V-P54T-T56Y-L80V- E87D-S99R-N114Q -S126A-P127D-S128G-N242D; and/or
- (xvii) P39E-I43V-A47V-P54T-T56Y-L80V- E87D-S99E-N114Q -S126T-P127D-S128A-N242D

- 30 Preferably, the variant has 90% identity with a sequence selected from the group consisting of SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7; SEQ ID NO:8, SEQ ID NO:9 and SEQ ID NO:10 and comprises at least two, more preferably at least three amino acid substitutions (using the SEQ ID NO:2 numbering) selected from the group consisting of

X39E, X126A-X127E-X128G and X242D, and preferably the variant comprises the following substitutions X39E, X126A-X127E-X128G and X242D.

5 Preferably, the variant further comprises at least one amino acid substitution (using the SEQ ID NO:2 numbering) selected from X54T, X114Q and X114C.

10 Preferably, the variant further comprises at least one and more preferably at least two and more preferably three amino acid substitution(s) (using the SEQ ID NO:2 numbering) selected from the group consisting of X74D; X80V, R, Y; X85S, C, D, I, K; and X87D, C.

15 Preferred variants have a sequence having at least 90% identity with a sequence selected from the group consisting of SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7; SEQ ID NO:8, SEQ ID NO:9 and SEQ ID NO:10. The substitutions listed above (1)-94) and (i) to (xvii)) for SEQ ID NO:2 applies *mutatis mutandis* to SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7; SEQ ID NO:8, SEQ ID NO:9 and SEQ ID NO:10, regardless of the amino acid residue present in these sequences at the positions listed for SEQ ID NO:2. It is understood that SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7; SEQ ID NO:8, SEQ ID NO:9 and SEQ ID NO:10 might not have the same amino acid residues as SEQ ID NO:2 in the same positions, but the variants comprise the same substitutions.

20 For example:

T056Y-S099R-S126A-P127E-S128G with respect to SEQ ID No:2 becomes X056Y-X099R-X126A-X127E-X128G with respect to any of SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7; SEQ ID NO:8, SEQ ID NO:9 and SEQ ID NO:10.

25 The protease of the invention performs very well in phosphate-free compositions even when the compositions are used in soft water.

Preferred levels of protease in the composition of the invention include from about 0.04 to about 5 mg, more preferably from about 0.05 to about 2 mg of active protease per gram of the composition.

30 Automatic dishwashing cleaning composition

The automatic dishwashing cleaning composition can be in any physical form. It can be a loose powder, a gel or presented in unit dose form. Preferably it is in unit dose form, unit dose forms

include pressed tablets and water-soluble packs. The automatic dishwashing cleaning composition of the invention is preferably presented in unit-dose form and it can be in any physical form including solid, liquid and gel form. The composition of the invention is very well suited to be presented in the form of a multi-compartment pack, more in particular a multi-compartment pack

5 comprising compartments with compositions in different physical forms, for example a compartment comprising a composition in solid form and another compartment comprising a composition in liquid form. The composition is preferably enveloped by a water-soluble film such as polyvinyl alcohol. Especially preferred are compositions in unit dose form wrapped in a polyvinyl alcohol film having a thickness of less than 100 μm , preferably from 20 to 90 μm . The

10 detergent composition of the invention weighs from about 8 to about 25 grams, preferably from about 10 to about 20 grams. This weight range fits comfortably in a dishwasher dispenser. Even though this range amounts to a low amount of detergent, the detergent has been formulated in a way that provides all the benefits mentioned herein above.

15 The composition is preferably phosphate free. By “phosphate-free” is herein understood that the composition comprises less than 1%, preferably less than 0.1% by weight of the composition of phosphate.

The composition of the invention is phosphate-free and comprises a complexing agent system

20 comprising a high level of a strong complexing agent.

Complexing agent system

For the purpose of this invention, a “complexing agent” is a compound capable of binding

25 polyvalent ions such as calcium, magnesium, lead, copper, zinc, cadmium, mercury, manganese, iron, aluminium and other cationic polyvalent ions to form a water-soluble complex. The complexing agent has a logarithmic stability constant ($[\log K]$) for Ca^{2+} of at least 3. The stability constant, $\log K$, is measured in a solution of ionic strength of 0.1, at a temperature of 25° C.

The composition of the invention comprises from 10% to 50% by weight of the composition of a

30 complexing agent system comprising at least 10% by weight of the composition of methyl glycine diacetic acid (MGDA). The complexing system may additionally comprise a complexing agent selected from the group consisting of citric acid, glutamic-N,N-diacetic acid (GLDA), iminodisuccinic acid (IDS), carboxy methyl inulin, L-Aspartic acid N, N-diacetic acid tetrasodium

salt (ASDA) and mixtures thereof. For the purpose of this invention, the term “acid”, when referring to complexing agents, includes the acid and salts thereof.

5 In a preferred embodiment, the composition comprises at least 15%, more preferably from 20% to 40% by weight of the composition of MGDA, more preferably the tri-sodium salt of MGDA. Compositions comprising this high level of MGDA perform well in hard water and also in long and/or hot cycles.

The complexing agent system of the invention can further comprise citric acid.

10

Dispersant polymer

A dispersant polymer can be used in any suitable amount from about 0.1 to about 20%, preferably from 0.2 to about 15%, more preferably from 0.3 to % by weight of the composition.

15 The dispersant polymer is capable to suspend calcium or calcium carbonate in an automatic dishwashing process.

The dispersant polymer has a calcium binding capacity within the range between 30 to 250 mg of Ca/g of dispersant polymer, preferably between 35 to 200 mg of Ca/g of dispersant polymer, more preferably 40 to 150 mg of Ca/g of dispersant polymer at 25°C. In order to determine if a polymer is a dispersant polymer within the meaning of the invention, the following calcium binding-
20 capacity determination is conducted in accordance with the following instructions:

Calcium binding capacity test method

The calcium binding capacity referred to herein is determined via titration using a pH/ion meter, such as the Meettler Toledo SevenMulti™ bench top meter and a PerfectION™ comb Ca combination electrode. To measure the binding capacity a heating and stirring device suitable for
25 beakers or tergotometer pots is set to 25 °C, and the ion electrode with meter are calibrated according to the manufacturer’s instructions. The standard concentrations for the electrode calibration should bracket the test concentration and should be measured at 25 °C. A stock solution of 1000 mg/g of Ca is prepared by adding 3.67 g of CaCl₂·2H₂O into 1 L of deionised water, then dilutions are carried out to prepare three working solutions of 100 mL each, respectively
30 comprising 100 mg/g, 10 mg/g, and 1 mg/g concentrations of Calcium. The 100 mg Ca/g working solution is used as the initial concentration during the titration, which is conducted at 25 °C. The

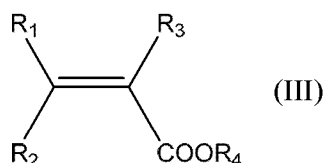
ionic strength of each working solution is adjusted by adding 2.5 g/L of NaCl to each. The 100 mL of 100 mg Ca/g working solution is heated and stirred until it reaches 25 °C. The initial reading of Calcium ion concentration is conducted at when the solution reaches 25 °C using the ion electrode. Then the test polymer is added incrementally to the calcium working solution (at 0.01 g/L intervals) and measured after 5 minutes of agitation following each incremental addition. The titration is stopped when the solution reaches 1 mg/g of Calcium. The titration procedure is repeated using the remaining two calcium concentration working solutions. The binding capacity of the test polymer is calculated as the linear slope of the calcium concentrations measured against the grams/L of test polymer that was added.

- 10 The dispersant polymer preferably bears a negative net charge when dissolved in an aqueous solution with a pH greater than 6.

The dispersant polymer can bear also sulfonated carboxylic esters or amides, in order to increase the negative charge at lower pH and improve their dispersing properties in hard water. The preferred dispersant polymers are sulfonated / carboxylated polymers, i.e., polymer comprising both sulfonated and carboxylated monomers.

Preferably, the dispersant polymers are sulfonated derivatives of polycarboxylic acids and may comprise two, three, four or more different monomer units. The preferred copolymers contain:

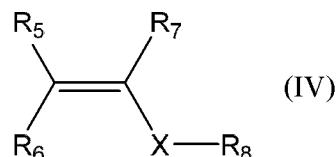
At least one structural unit derived from a carboxylic acid monomer having the general formula (III):



- 20 wherein R₁ to R₃ are independently selected from hydrogen, methyl, linear or branched saturated alkyl groups having from 2 to 12 carbon atoms, linear or branched mono or polyunsaturated alkenyl groups having from 2 to 12 carbon atoms, alkyl or alkenyl groups as aforementioned substituted with –NH₂ or –OH, or –COOH, or COOR₄, where R₄ is selected from hydrogen, alkali metal, or a linear or branched, saturated or unsaturated alkyl or alkenyl group with 2 to 12 carbons;
- 25 Preferred carboxylic acid monomers include one or more of the following: acrylic acid, maleic acid, maleic anhydride, itaconic acid, citraconic acid, 2-phenylacrylic acid, cinnamic acid, crotonic

acid, fumaric acid, methacrylic acid, 2-ethylacrylic acid, methylenemalononic acid, or sorbic acid. Acrylic and methacrylic acids being more preferred.

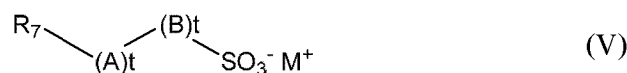
Optionally, one or more structural units derived from at least one nonionic monomer having the general formula (IV):

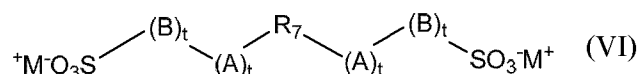


- 5 Wherein R₅ to R₇ are independently selected from hydrogen, methyl, phenyl or hydroxyalkyl groups containing 1 to 6 carbon atoms, and can be part of a cyclic structure, X is an optionally present spacer group which is selected from -CH₂-, -COO-, -CONH- or -CONR₈-, and R₈ is selected from linear or branched, saturated alkyl radicals having 1 to 22 carbon atoms or unsaturated, preferably aromatic, radicals having from 6 to 22 carbon atoms.
- 10 Preferred non-ionic monomers include one or more of the following: butene, isobutene, pentene, 2-methylpent-1-ene, 3-methylpent-1-ene, 2,4,4-trimethylpent-1-ene, 2,4,4-trimethylpent-2-ene, cyclopentene, methylcyclopentene, 2-methyl-3-methyl-cyclopentene, hexene, 2,3-dimethylhex-1-ene, 2,4-dimethylhex-1-ene, 2,5-dimethylhex-1-ene, 3,5-dimethylhex-1-ene, 4,4-dimethylhex-1-ene, cyclohexene, methylcyclohexene, cycloheptene, alpha olefins having 10 or more carbon
- 15 atoms such as, dec-1-ene, dodec-1-ene, hexadec-1-ene, octadec-1-ene and docos-1-ene, preferred aromatic monomers are styrene, alpha methylstyrene, 3-methylstyrene, 4-dodecylstyrene, 2-ethyl-4-benzylstyrene, 4-cyclohexylstyrene, 4-propylstyrol, 1-vinylnaphtalene, 2-vinylnaphtalene; preferred carboxylic ester monomers are methyl (meth)acrylate, ethyl (meth)acrylate, propyl (meth)acrylate, t-butyl (meth)acrylate, pentyl (meth)acrylate, hexyl (meth)acrylate, 2-ethylhexyl
- 20 (meth)acrylate, octyl (meth)acrylate, lauryl (meth)acrylate, stearyl (meth)acrylate and behenyl (meth)acrylate; preferred amides are N-methyl acrylamide, N-ethyl acrylamide, N-t-butyl acrylamide, N-2-ethylhexyl acrylamide, N-octyl acrylamide, N-lauryl acrylamide, N-stearyl acrylamide, N-behenyl acrylamide.

and at least one structural unit derived from at least one sulfonic acid monomer having the general

25 formula (V) and (VI):





wherein R₇ is a group comprising at least one sp² bond, A is O, N, P, S, an amido or ester linkage, B is a mono- or polycyclic aromatic group or an aliphatic group, each t is independently 0 or 1, and M⁺ is a cation. In one aspect, R₇ is a C₂ to C₆ alkene. In another aspect, R₇ is ethene, butene or propene.

- 5 Preferred sulfonated monomers include one or more of the following: 1-acrylamido-1-propanesulfonic acid, 2-acrylamido-2-propanesulfonic acid, 2-acrylamido-2-methyl-1-propanesulfonic acid, 2-methacrylamido-2-methyl-1-propanesulfonic acid, 3-methacrylamido-2-hydroxy-propanesulfonic acid, allylsulfonic acid, methallylsulfonic acid, allyloxybenzenesulfonic acid, methallyloxybenzenesulfonic acid, 2-hydroxy-3-(2-propenyloxy) propanesulfonic acid, 2-methyl-2-propen-1-sulfonic acid, styrenesulfonic acid, vinylsulfonic acid, 3-sulfopropyl, 3-sulfo-
10 propylmethacrylate, sulfomethacrylamide, sulfomethylmethacrylamide and mixtures of said acids or their water-soluble salts.

Preferably, the polymer comprises the following levels of monomers: from about 40 to about 90%, preferably from about 60 to about 90% by weight of the polymer of one or more carboxylic acid
15 monomer; from about 5 to about 50%, preferably from about 10 to about 40% by weight of the polymer of one or more sulfonic acid monomer; and optionally from about 1% to about 30%, preferably from about 2 to about 20% by weight of the polymer of one or more non-ionic monomer. An especially preferred polymer comprises about 70% to about 80% by weight of the polymer of at least one carboxylic acid monomer and from about 20% to about 30% by weight of
20 the polymer of at least one sulfonic acid monomer.

In the polymers, all or some of the carboxylic or sulfonic acid groups can be present in neutralized form, i.e. the acidic hydrogen atom of the carboxylic and/or sulfonic acid group in some or all acid groups can be replaced with metal ions, preferably alkali metal ions and in particular with sodium ions.

- 25 The carboxylic acid is preferably (meth)acrylic acid. The sulfonic acid monomer is preferably 2-acrylamido-2-propanesulfonic acid (AMPS).

Preferred commercial available polymers include: AlcosperseTM 240, AquatreatTM AR 540 and Aquatreat MPS supplied by Alco Chemical; AcumerTM 3100, Acumer 2000, AcusolTM 587G and

Acusol 588G supplied by Rohm & Haas; Goodrich K-798, K-775 and K-797 supplied by BF Goodrich; and ACP 1042 supplied by ISP technologies Inc. Particularly preferred polymers are Acusol 587G and Acusol 588G supplied by Rohm & Haas.

Suitable dispersant polymers include anionic carboxylic polymer of low molecular weight. They
5 can be homopolymers or copolymers with a weight average molecular weight of less than or equal to about 200,000 g/mol, or less than or equal to about 75,000 g/mol, or less than or equal to about 50,000 g/mol, or from about 3,000 to about 50,000 g/mol, preferably from about 5,000 to about 45,000 g/mol. The dispersant polymer may be a low molecular weight homopolymer of polyacrylate, with an average molecular weight of from 1,000 to 20,000, particularly from 2,000
10 to 10,000, and particularly preferably from 3,000 to 5,000.

The dispersant polymer may be a copolymer of acrylic with methacrylic acid, acrylic and/or methacrylic with maleic acid, and acrylic and/or methacrylic with fumaric acid, with a molecular weight of less than 70,000. Their molecular weight ranges from 2,000 to 80,000 and more preferably from 20,000 to 50,000 and in particular 30,000 to 40,000 g/mol. and a ratio of
15 (meth)acrylate to maleate or fumarate segments of from 30:1 to 1:2.

The dispersant polymer may be a copolymer of acrylamide and acrylate having a molecular weight of from 3,000 to 100,000, alternatively from 4,000 to 20,000, and an acrylamide content of less than 50%, alternatively less than 20%, by weight of the dispersant polymer can also be used. Alternatively, such dispersant polymer may have a molecular weight of from 4,000 to 20,000 and
20 an acrylamide content of from 0% to 15%, by weight of the polymer.

Dispersant polymers suitable herein also include itaconic acid homopolymers and copolymers. Alternatively, the dispersant polymer can be selected from the group consisting of alkoxylated polyalkyleneimines, alkoxylated polycarboxylates, polyethylene glycols, styrene co-polymers, cellulose sulfate esters, carboxylated polysaccharides, amphiphilic graft copolymers and mixtures
25 thereof.

Bleaching system

The composition of the invention preferably comprises a bleaching system comprising a high level
30 of bleach, preferably percarbonate in combination with a bleach activator or a bleach catalyst or both. Preferably the bleach activator is TAED and the bleach catalyst is a manganese bleach catalyst.

Bleach

The composition of the invention preferably comprises from about 10 to about 20%, more preferably from about 12 to about 18% of bleach, preferably percarbonate, by weight of the composition.

- 5 Inorganic and organic bleaches are suitable for use herein. Inorganic bleaches include perhydrate salts such as perborate, percarbonate, perphosphate, persulfate and persilicate salts. The inorganic perhydrate salts are normally the alkali metal salts. The inorganic perhydrate salt may be included as the crystalline solid without additional protection. Alternatively, the salt can be coated. Suitable coatings include sodium sulphate, sodium carbonate, sodium silicate and mixtures thereof. Said
10 coatings can be applied as a mixture applied to the surface or sequentially in layers.

Alkali metal percarbonates, particularly sodium percarbonate is the preferred bleach for use herein. The percarbonate is most preferably incorporated into the products in a coated form which provides in-product stability.

Potassium peroxymonopersulfate is another inorganic perhydrate salt of utility herein.

- 15 Typical organic bleaches are organic peroxyacids, especially dodecanediperoxoic acid, tetradecanediperoxoic acid, and hexadecanediperoxoic acid. Mono- and diperazelaic acid, mono- and diperbrassylic acid are also suitable herein. Diacyl and Tetraacylperoxides, for instance dibenzoyl peroxide and dilauroyl peroxide, are other organic peroxides that can be used in the context of this invention.

- 20 Further typical organic bleaches include the peroxyacids, particular examples being the alkylperoxy acids and the arylperoxy acids. Preferred representatives are (a) peroxybenzoic acid and its ring-substituted derivatives, such as alkylperoxybenzoic acids, but also peroxy- α -naphthoic acid and magnesium monoperphthalate, (b) the aliphatic or substituted aliphatic peroxy acids, such as peroxyauric acid, peroxysearic acid, ϵ -phthalimidoperoxycaproic
25 acid[phthaloininoperoxylhexanoic acid (PAP)], o-carboxybenzamidoperoxycaproic acid, N-nonenylamidoperadipic acid and N-nonenylamidopersuccinates, and (c) aliphatic and araliphatic peroxydicarboxylic acids, such as 1,12-diperoxycarboxylic acid, 1,9-diperoxyazelaic acid, diperoxysebacic acid, diperoxybrassylic acid, the diperoxyphthalic acids, 2-decyldiperoxybutane-1,4-dioic acid, N,N-terephthaloyldi(6-aminopercaproic acid).

30

Bleach Activators

Bleach activators are typically organic peracid precursors that enhance the bleaching action in the course of cleaning at temperatures of 60° C and below. Bleach activators suitable for use herein

include compounds which, under perhydrolysis conditions, give aliphatic peroxycarboxylic acids having preferably from 1 to 12 carbon atoms, in particular from 2 to 10 carbon atoms, and/or optionally substituted perbenzoic acid. Suitable substances bear O-acyl and/or N-acyl groups of the number of carbon atoms specified and/or optionally substituted benzoyl groups. Preference is

5 given to polyacylated alkylenediamines, in particular tetraacetylenediamine (TAED), acylated triazine derivatives, in particular 1,5-diacetyl-2,4-dioxohexahydro-1,3,5-triazine (DADHT), acylated glycolurils, in particular tetraacetyl glycoluril (TAGU), N-acylimides, in particular N-nonanoylsuccinimide (NOSI), acylated phenolsulfonates, in particular n-nonanoyl-

10 or isononanoyloxybenzenesulfonate (n- or iso-NOBS), decanoyloxybenzoic acid (DOBA), carboxylic anhydrides, in particular phthalic anhydride, acylated polyhydric alcohols, in particular triacetin, ethylene glycol diacetate and 2,5-diacetoxy-2,5-dihydrofuran and also triethylacetyl citrate (TEAC). If present the composition of the invention comprises from 0.01 to 5, preferably from 0.2 to 2% by weight of the composition of bleach activator, preferably TAED.

15 Bleach Catalyst

The composition herein preferably contains a bleach catalyst, preferably a metal containing bleach catalyst. More preferably the metal containing bleach catalyst is a transition metal containing bleach catalyst, especially a manganese or cobalt-containing bleach catalyst.

20 Bleach catalysts preferred for use herein include manganese triazacyclononane and related complexes; Co, Cu, Mn and Fe bispyridylamine and related complexes; and pentamine acetate cobalt(III) and related complexes. Especially preferred bleach catalyst for use herein are 1,4,7-trimethyl-1,4,7-triazacyclononane (Me-TACN) and 1,2, 4,7- tetramethyl-1,4,7-triazacyclononane (Me/Me-TACN).

25 Preferably the composition of the invention comprises from 0.001 to 0.5, more preferably from 0.002 to 0.05%, more preferably from 0.005 to 0.075% of bleach catalyst by weight of the composition. Preferably the bleach catalyst is a manganese bleach catalyst.

Inorganic builder

30

The composition of the invention preferably comprises an inorganic builder. Suitable inorganic builders are selected from the group consisting of carbonate, silicate and mixtures thereof. Especially preferred for use herein is sodium carbonate. Preferably the composition of the

invention comprises from 5 to 60%, more preferably from 10 to 50% and especially from 15 to 45% of sodium carbonate by weight of the composition.

Surfactant

5

Surfactants suitable for use herein include non-ionic surfactants, preferably the compositions are free of any other surfactants. Traditionally, non-ionic surfactants have been used in automatic dishwashing for surface modification purposes in particular for sheeting to avoid filming and spotting and to improve shine. It has been found that non-ionic surfactants can also contribute to prevent redeposition of soils.

10

Preferably the composition of the invention comprises a non-ionic surfactant or a non-ionic surfactant system, more preferably the non-ionic surfactant or a non-ionic surfactant system has a phase inversion temperature, as measured at a concentration of 1% in distilled water, between 40 and 70°C, preferably between 45 and 65°C. By a “non-ionic surfactant system” is meant herein a mixture of two or more non-ionic surfactants. Preferred for use herein are non-ionic surfactant systems. They seem to have improved cleaning and finishing properties and better stability in product than single non-ionic surfactants.

15

Phase inversion temperature is the temperature below which a surfactant, or a mixture thereof, partitions preferentially into the water phase as oil-swollen micelles and above which it partitions preferentially into the oil phase as water swollen inverted micelles. Phase inversion temperature can be determined visually by identifying at which temperature cloudiness occurs.

20

The phase inversion temperature of a non-ionic surfactant or system can be determined as follows: a solution containing 1% of the corresponding surfactant or mixture by weight of the solution in distilled water is prepared. The solution is stirred gently before phase inversion temperature analysis to ensure that the process occurs in chemical equilibrium. The phase inversion temperature is taken in a thermostable bath by immersing the solutions in 75 mm sealed glass test tube. To ensure the absence of leakage, the test tube is weighed before and after phase inversion temperature measurement. The temperature is gradually increased at a rate of less than 1°C per minute, until the temperature reaches a few degrees below the pre-estimated phase inversion temperature. Phase inversion temperature is determined visually at the first sign of turbidity.

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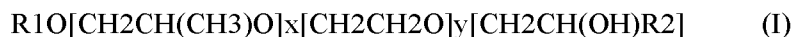
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Suitable nonionic surfactants include: i) ethoxylated non-ionic surfactants prepared by the reaction of a monohydroxy alkanol or alkylphenol with 6 to 20 carbon atoms with preferably at least 12 moles particularly preferred at least 16 moles, and still more preferred at least 20 moles of ethylene oxide per mole of alcohol or alkylphenol; ii) alcohol alkoxylated surfactants having a from 6 to 20

carbon atoms and at least one ethoxy and propoxy group. Preferred for use herein are mixtures of surfactants i) and ii).

Another suitable non-ionic surfactants are epoxy-capped poly(oxyalkylated) alcohols represented by the formula:

5



wherein R₁ is a linear or branched, aliphatic hydrocarbon radical having from 4 to 18 carbon atoms; R₂ is a linear or branched aliphatic hydrocarbon radical having from 2 to 26 carbon atoms;
 10 x is an integer having an average value of from 0.5 to 1.5, more preferably about 1; and y is an integer having a value of at least 15, more preferably at least 20.

Preferably, the surfactant of formula I, at least about 10 carbon atoms in the terminal epoxide unit [CH₂CH(OH)R₂]. Suitable surfactants of formula I, according to the present invention, are Olin Corporation's POLY-TERGENT® SLF-18B nonionic surfactants, as described, for example, in
 15 WO 94/22800, published October 13, 1994 by Olin Corporation.

Enzymes

Other proteases

20 The composition of the invention can comprise a protease in addition to the protease of the invention. A mixture of two or more proteases can contribute to an enhanced cleaning across a broader temperature, cycle duration, and/or substrate range, and provide superior shine benefits, especially when used in conjunction with an anti-redeposition agent and/or a sulfonated polymer.

25 Suitable proteases for use in combination with the variant proteases of the invention 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
 30 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:

(a) subtilisins (EC 3.4.21.62), especially those derived from *Bacillus*, such as *Bacillus* sp., *B. lentus*, *B. alkalophilus*, *B. subtilis*, *B. amyloliquefaciens*, *B. pumilus*, *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, DE102006022216A1, DE102006022224A1, WO2015089447, WO2015089441, 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 *Cellulomonas* described in WO 05/052161 and WO 05/052146.

(c) metalloproteases, especially those derived from *Bacillus amyloliquefaciens* described in WO07/044993A2; 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.

Especially preferred additional proteases for the detergent of the invention are polypeptides demonstrating at least 90%, preferably at least 95%, more preferably at least 98%, even more preferably at least 99% and especially 100% identity with the wild-type enzyme from *Bacillus lentus*, comprising mutations in one or more, preferably two or more and more preferably three or more of the following positions, using the BPN' numbering system and amino acid abbreviations as illustrated in WO00/37627: V68A, N76D, N87S, S99D, S99SD, S99A, S101G, S101M, S103A, V104N/I, G118V, G118R, S128L, P129Q, S130A, Y167A, R170S, A194P, V205I, Q206L/D/E, Y209W and/or M222S.

Most preferably the additional protease is selected from the group of proteases comprising the below mutations (BPN' numbering system) versus either the PB92 wild-type (SEQ ID NO:2 in WO 08/010925) or the subtilisin 309 wild-type (sequence as per PB92 backbone, except comprising a natural variation of N87S).

(i) G118V + S128L + P129Q + S130A

(ii) S101M + G118V + S128L + P129Q + S130A

5 (iii) N76D + N87R + G118R + S128L + P129Q + S130A + S188D + N248R

(iv) N76D + N87R + G118R + S128L + P129Q + S130A + S188D + V244R

(v) N76D + N87R + G118R + S128L + P129Q + S130A

10

(vi) V68A + N87S + S101G + V104N

(vii) S99AD

15 Suitable commercially available additional protease enzymes include those sold under the trade names Alcalase®, Savinase®, Primase®, Durazym®, Polarzyme®, Kannase®, Liquanase®, Liquanase Ultra®, Savinase Ultra®, Ovozyme®, Neutrase®, Everlase®, Coronase®, Blaze®, Blaze Ultra® and Esperase® by Novozymes A/S (Denmark); those sold under the tradename Maxatase®, Maxacal®, Maxapem®, Properase®, Purafect®, Purafect Prime®, Purafect Ox®,
 20 FN3®, FN4®, Excellase®, Ultimase® and Purafect OXP® by Dupont; those sold under the tradename Opticlean® and Optimase® by Solvay Enzymes; and 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)
 25 and BLAP F49 (BLAP with S3T + V4I + A194P + V199M + V205I + L217D); and KAP (Bacillus alkalophilus subtilisin with mutations A230V + S256G + S259N) from Kao.

Especially preferred for use herein in combination with the variant protease of the invention are commercial proteases selected from the group consisting of Properase®, Blaze®, Ultimase®,
 30 Everlase®, Savinase®, Excellase®, Blaze Ultra®, BLAP and BLAP variants.

Preferred levels of protease in the product of the invention include from about 0.05 to about 10, more preferably from about 0.5 to about 7 and especially from about 1 to about 6 mg of active protease/g of composition.

Amylases

Preferably the composition of the invention 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
 5 *Bacillus licheniformis*, *Bacillus amyloliquefaciens*, *Bacillus stearothermophilus*, *Bacillus subtilis*, or other *Bacillus* sp., such as *Bacillus* sp. NCBI 12289, NCBI 12512, NCBI 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:

- 10 (a) variants described in WO 96/23873, WO00/60060, WO06/002643 and WO2017/192657, especially the variants with one or more substitutions in the following positions versus the AA560 enzyme listed as SEQ ID NO. 12 in WO06/002643:

26, 30, 33, 82, 37, 106, 118, 128, 133, 149, 150, 160, 178, 182, 186, 193, 202, 214, 231, 246, 256,
 15 257, 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*.

- (b) 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
 20 variants described in WO 00/60060, WO2011/100410 and WO2013/003659.

- (c) 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
 25 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.

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

(e) variants exhibiting at least 89% identity with SEQ ID NO:1 in WO2016091688, especially those comprising deletions at positions H183+G184 and additionally one or more mutations at positions 405, 421, 422 and/or 428.

5 (f) variants exhibiting at least 60% amino acid sequence identity with the "PcuAmyl α -amylase" from *Paenibacillus curdlanolyticus* YK9 (SEQ ID NO:3 in WO2014099523).

(g) variants exhibiting at least 60% amino acid sequence identity with the "CspAmy2 amylase" from *Cytophaga* sp. (SEQ ID NO:1 in WO2014164777).

10

(h) variants exhibiting at least 85% identity with AmyE from *Bacillus subtilis* (SEQ ID NO:1 in WO2009149271).

15 (i) variants exhibiting at least 90% identity with the wild-type amylase from *Bacillus* sp. KSM-K38 with accession number AB051102.

(j) variants exhibiting at least 80% identity with the mature amino acid sequence of AAI10 from *Bacillus* sp (SEQ ID NO:7 in WO2016180748)

20 (k) variants exhibiting at least 80% identity with the mature amino acid sequence of *Alicyclobacillus* sp. amylase (SEQ ID NO:8 in WO2016180748)

25 Preferably the amylase is an engineered enzyme, wherein one or more of the amino acids prone to bleach oxidation have been substituted by an amino acid less prone to oxidation. In particular it is preferred that methionine residues are substituted with any other amino acid. In particular it is preferred that the methionine most prone to oxidation is substituted. Preferably the methionine in a position equivalent to 202 in the AA560 enzyme listed as SEQ ID NO. 12 in WO06/002643 is substituted. Preferably, the methionine at this position is substituted with threonine or leucine, preferably leucine.

30

Suitable commercially available alpha-amylases include DURAMYL®, LIQUEZYME®, TERMAMYL®, TERMAMYL ULTRA®, NATALASE®, SUPRAMYL®, STAINZYME®, STAINZYME PLUS®, FUNGAMYL®, ATLANTIC®, INTENSA® and BAN® (Novozymes A/S, Bagsvaerd, Denmark), KEMZYM® AT 9000 Biozym Biotech Trading GmbH Wehlstrasse

27b A-1200 Wien Austria, RAPIDASE®, PURASTAR®, ENZYSE®, OPTISIZE HT PLUS®, POWERASE®, PREFERENZ S® series (including PREFERENZ S1000® and PREFERENZ S2000® and PURASTAR OXAM® (DuPont., Palo Alto, California) and KAM® (Kao, 14-10 Nihonbashi Kayabacho, 1-chome, Chuo-ku Tokyo 103-8210, Japan). In one aspect, suitable
 5 amylases include ATLANTIC®, STAINZYME®, POWERASE®, INTENSA® and STAINZYME PLUS® and mixtures thereof.

Preferably, the product of the invention comprises at least 0.01 mg, preferably from about 0.05 to about 10, more preferably from about 0.1 to about 6, especially from about 0.2 to about 5 mg of
 10 active amylase/ g of composition.

Preferably, the protease and/or amylase of the composition of the invention are in the form of granulates, the granulates comprise more than 29% of sodium sulfate by weight of the granulate and/or the sodium sulfate and the active enzyme (protease and/or amylase) are in a weight ratio of
 15 between 3:1 and 100:1 or preferably between 4:1 and 30:1 or more preferably between 5:1 and 20:1.

Crystal growth inhibitor

20 Crystal growth inhibitors are materials that can bind to calcium carbonate crystals and prevent further growth of species such as aragonite and calcite.
 Examples of effective crystal growth inhibitors include phosphonates, polyphosphonates, inulin derivatives, polyitaconic acid homopolymers and cyclic polycarboxylates.

25 Suitable crystal growth inhibitors may be selected from the group comprising HEDP (1-hydroxyethylidene 1,1-diphosphonic acid), carboxymethylinulin (CMI), tricarballic acid and cyclic carboxylates. For the purposes of this invention the term carboxylate covers both the anionic form and the protonated carboxylic acid form.

Cyclic carboxylates contain at least two, preferably three or preferably at least four
 30 carboxylate groups and the cyclic structure is based on either a mono- or bi-cyclic alkane or a heterocycle. Suitable cyclic structures include cyclopropane, cyclobutane, cyclohexane or cyclopentane or cycloheptane, bicyclo-heptane or bicyclo-octane and/or tetrahydrofuran. One preferred crystal growth inhibitor is cyclopentane tetracarboxylate.

Cyclic carboxylates having at least 75%, preferably 100% of the carboxylate groups on the same side, or in the “cis” position of the 3D-structure of the cycle are preferred for use herein.

It is preferred that the two carboxylate groups, which are on the same side of the cycle are in directly neighbouring or “ortho” positions.

5 Preferred crystal growth inhibitors include HEDP, tricarballic acid, tetrahydrofuranetracarboxylic acid (THFTCA) and cyclopentanetracarboxylic acid (CPTCA). The THFTCA is preferably in the 2c,3t,4t,5c-configuration, and the CPTCA in the cis,cis,cis,cis-configuration. Especially preferred crystal growth inhibitor for use herein is HEDP.

Also, preferred for use herein are partially decarboxylated polyitaconic acid homopolymers,
10 preferably having a level of decarboxylation is in the range of 50 mole % to 90 mole %. Especially preferred polymer for use herein is Itaconix TSI® provided by Itaconix.

The crystal growth inhibitors are present preferably in a quantity from about 0.01 to about 10 %, particularly from about 0.02 to about 5 % and in particular, from 0.05 to 3 % by weight of the
15 composition.

Metal Care Agents

Metal care agents may prevent or reduce the tarnishing, corrosion or oxidation of metals, including
20 aluminium, stainless steel and non-ferrous metals, such as silver and copper. Preferably the composition of the invention comprises from 0.1 to 5%, more preferably from 0.2 to 4% and especially from 0.3 to 3% by weight of the product of a metal care agent, preferably the metal care agent is benzo triazole (BTA).

25 Glass Care Agents

Glass care agents protect the appearance of glass items during the dishwashing process. Preferably the composition of the invention comprises from 0.1 to 5%, more preferably from 0.2 to 4% and specially from 0.3 to 3% by weight of the composition of a metal care agent, preferably the glass
30 care agent is a zinc containing material, specially hydrozincite. Other suitable glass care agents are polyethyleneimine (PEI). A particularly preferred PEI is Lupasol® FG, supplied by BASF.

The automatic dishwashing composition of the invention preferably has a pH as measured in 1% weight/volume aqueous solution in distilled water at 20°C of from about 9 to about 12, more preferably from about 10 to less than about 11.5 and especially from about 10.5 to about 11.5.

The automatic dishwashing composition of the invention preferably has a reserve alkalinity of
 5 from about 10 to about 20, more preferably from about 12 to about 18 at a pH of 9.5 as measured in NaOH with 100 grams of product at 20°C.

A preferred automatic dishwashing composition of the invention comprises:

- 10 i) from 10 to 20% by weight of the composition of bleach, preferably sodium percarbonate;
- ii) TAED;
- iii) optionally but preferably from 5 to 50% by weight of the composition of an inorganic builder, preferably sodium carbonate;
- 15 iv) optionally but preferably from 2 to 10% by weight of the composition of a non-ionic surfactant;
- v) other optional ingredients include: a crystal growth inhibitor, preferably HEDP, and glass care agents.

A preferred automatic dishwashing composition of the invention comprises:

- 20 i) from 10 to 20% by weight of the composition of bleach, preferably sodium percarbonate;
- ii) a manganese bleach catalyst and optionally TAED;
- iii) optionally but preferably from 5 to 50% by weight of the composition of an inorganic builder, preferably sodium carbonate;
- 25 iv) optionally but preferably from 2 to 10% by weight of the composition of a non-ionic surfactant;
- v) optionally but preferably a glass care agent.

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.

30 For example, a dimension disclosed as “40 mm” is intended to mean “about 40 mm.”

EXAMPLES

Egg yolk removal by automatic dishwashing compositions comprising variant proteases were compared with the same compositions comprising the parent protease.

- 5 The compositions displayed in Table 1 were used. 3 g of each composition were dissolved in a litre of deionized water to produce a cleaning solution having a pH of 11. The corresponding protease was added to the cleaning solution at a level between 0.25-0.75ppm.

Table 1: Automatic Dishwashing Compositions

Ingredients (active weight %)	ADW Formula A	ADW Formula B
Solid ingredients		
Sodium carbonate	41.7	41.7
Sodium sulphate	0.00	2.03
MGDA	21.0	10.1
Sodium citrate	0.00	10.1
TAED	1.68	1.68
Sodium percarbonate	12.6	12.6
Sulfonated polymer	2.5	2.5
Bleach catalyst	1.2	1.2
Amylase	0.11	0.11
Liquid ingredients		
Lutensol TO7	19.3	19.3

Table 1: Automatic Dish Washing (ADW) Compositions

10

Amylase	Stainzyme [®] Plus supplied by Novozymes
TAED	Tetraacetylenediamine
MGDA	Three-sodium methyl glycine diacetate supplied by BASF
Bleach catalyst	MnTACN (Manganese 1,4,7-Triazacyclononane)
15 Sulfonated polymer	Acusol 588 supplied by Dow Chemicals
Lutensol TO7	Nonionic surfactant supplied by BASF

Cleaning Performance Method: Egg Yolk cleaning performance using PAS-38

The cleaning performance of the proteases variants listed in Table 2 was tested relative to the parent using Automatic Dishwashing Compositions A and B (see Table 1), as measured by the stain removal on egg yolk microswatches (PAS-38, Center for Testmaterials BV, Vlaardingen, Netherlands). The egg swatch stains were pre-sized to fit the microtiter plate (MTPs); standard
 5 96 well plate. The stain removal of the PAS-38 egg swatches was measured post wash versus a reference.

The MTPs were filled prior to protease addition with 3 g/l of detergent and the detergent and deionized water.

After incubating the PAS-38 swatches for 30°C min at 50°C, absorbance was read at 405 nm
 10 with a SpectraMax plate reader. Absorbance results were obtained by subtracting the value for a blank control (containing no protease) from each sample value (hereinafter “blank subtracted absorbance”). For each condition and variant, a performance index (PI) was calculated by dividing the blank subtracted absorbance by that of the parent protease at the same concentration. The value for the parent protease was determined from a standard curve of the parent protease
 15 which was included in the test and which was fitted to a Langmuir fit.

Protease Enzyme Activity Method 2: AAPF Assay

The protease activity of parent and subtilisin variants thereof was tested by measuring hydrolysis of N-suc-AAPF-pNA. The reagent solutions used for the AAPF hydrolysis assay were: 100 mM Tris/HCl pH 8.6, containing 0.005% TWEEN®-80 (Tris dilution buffer); 100 mM Tris buffer
 20 pH 8.6, containing 10 mM CaCl₂ and 0.005% TWEEN®-80 (Tris/Ca buffer); and 160 mM suc-AAPF-pNA in DMSO (suc-AAPF-pNA stock solution) (Sigma: S-7388). A substrate working solution was prepared by adding 1 mL suc-AAPF-pNA stock solution to 100 mL Tris/Ca buffer and mixed well. An enzyme sample was added to a MTP (Greiner 781101) containing 1 mg/suc-AAPF-pNA working solution and assayed for activity at 405 nm over 3 min with a SpectraMax
 25 plate reader in kinetic mode at room temperature (RT). The absorbance of a blank containing no protease was subtracted from each sample reading. The protease activity was expressed as mOD·min⁻¹.

Protease Enzyme Stability - Stability Assay

30 The stability of the variants described herein was measured by diluting the variants in stress buffer and measuring the proteolytic activity of the variants before and after a heat incubation step of 5 minutes at 56°C using the AAPF assay described above. Stability was measured in Tris-EDTA

(50mM Tris pH 9; 5 mM EDTA; 0.005% Tween 80) buffered condition. % Residual activities were calculated by taking a ratio of the stressed to unstressed activity and multiplying by 100.

Protease Data Table 2

5

		Stability	PAS-38 stain Cleaning	
Variants	Substitutions with respect to SEQ ID No:1	TRIS-EDTA	ADW formula A	ADW formula B
SEQ ID No:1	none	1	1.0	1.0
WALBSP-07063	T056Y-S099R-S126A-D127E-F128G	9	1.7	2.6
WALBSP-07078	S039E-I080V-S099R-S126A-D127E-F128G-M211L	21	2.2	ND
WALBSP-07137	S039E-P054T-S099R-S126A-D127E-F128G-M211L	28	2.3	ND
WALBSP-07117	S039E-I043V-S099R-S126A-D127E-F128G-M211L	28	2.3	3.7
WALBSP-07119	S039E-N042R-S099R-S126A-D127E-F128G	28	2.3	3.9
WALBSP-07052	S039E-I080V-S099R-S126A-D127E-F128G	29	2.1	2.8
WALBSP-07097	S039E-S099R-S126A-D127E-F128G-M211L	29	2.4	ND
WALBSP-07088	S039E-N085S-S099R-S126A-D127E-F128G-M211L	30	2.3	3.6
WALBSP-07147	S039E-T056Y-S099R-S126A-D127E-F128G-M211L	31	2.0	3.8

WALBSP-07127	S039E-A047V-S099R-S126A-D127E-F128G-M211L	31	2.2	3.9
WALBSP-07149	S039E-S099R-S126A-D127E-F128G-Y203W	32	2.0	ND
WALBSP-07107	A037T-S039E-S099R-S126A-D127E-F128G-M211L	32	2.3	ND
WALBSP-07129	S039E-S099R-S126A-D127E-F128G-V199I	34	2.3	4.3
WALBSP-07058	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G	35	2.0	2.8
WALBSP-07050	S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D	35	2.0	2.9
WALBSP-07040	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G	36	2.0	3.1
WALBSP-07014	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G	37	3.6	3.9
WALBSP-07012	S039E-I043V-S099R-S126A-D127E-F128G	38	2.2	2.7

WALBSP-07080	S039E-S099R-S126A-D127E-F128G-N253D	38	1.7	ND
WALBSP-07109	T009E-S039E-S099R-S126A-D127E-F128G	38	2.1	3.4
WALBSP-07090	S039E-S099R-S126A-D127E-F128G-S255W	38	2.4	3.8
WALBSP-07041	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-S099R-S126A-D127E-F128G-N242D	38	2.0	2.9
WALBSP-07001	A037T-S039E-I043V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D	38	1.9	3.0
WALBSP-07024	S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D	39	2.1	2.5
WALBSP-07068	S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D	39	1.8	2.8
WALBSP-07042	S039E-T056Y-S099R-S126A-D127E-F128G	39	2.1	2.6
WALBSP-07065	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G	39	2.1	2.7
WALBSP-07027	A037T-S039E-I043V-P054T-T056Y-I080V-	39	1.9	3.0

	N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D			
WALBSP-07019	A037T-S039E-I043V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D	39	2.1	2.9
WALBSP-07029	A037T-S039E-I043V-A047V-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D	39	1.9	3.0
WALBSP-07011	A037T-S039E-I043V-A047V-P054T-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D	39	2.0	2.8
WALBSP-07055	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-N242D	39	2.0	2.6
WALBSP-07054	A037T-S039E-I043V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D	39	2.0	2.7
WALBSP-07030	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-N242D	39	2.0	3.0
WALBSP-07062	S039E-N085S-S099R-S126A-D127E-F128G	40	2.0	2.7

WALBSP-07047	A037T-S039E-I043V-A047V-P054T-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D	40	1.9	2.8
WALBSP-06998	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-S099R-T114Q-S126A-D127E-F128G-N242D	40	1.9	2.7
WALBSP-07048	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-N242D	40	2.0	3.0
WALBSP-07051	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S126A-D127E-F128G-N242D	40	0.7	0.8
WALBSP-07045	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-F128G-N242D	40	2.1	2.6
WALBSP-07049	A037T-S039E-I043V-A047V-T056Y-I080V-E087D-S099R-T114Q-S126A-D127E-F128G-N242D	41	2.0	2.9
WALBSP-07003	S039E-S099R-T114Q-S126A-D127E-F128G	41	2.1	2.8
WALBSP-07004	A037T-S039E-I043V-A047V-P054T-T056Y-	41	1.9	2.6

	I080V-N085S-E087D-S099R-S126A-D127E-N242D			
WALBSP-07000	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D	41	1.9	2.7
WALBSP-07028	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-D127E-F128G-N242D	41	2.1	2.7
WALBSP-07059	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-S099R-T114Q-S126A-D127E-F128G-N242D	41	2.0	3.0
WALBSP-07022	S039E-A047V-S099R-S126A-D127E-F128G	41	2.1	2.5
WALBSP-07015	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-S099R-S126A-D127E-F128G-N242D	41	2.0	2.7
WALBSP-07033	A037T-S039E-I043V-A047V-T056Y-S099R-T114Q-S126A-D127E-F128G	41	2.1	2.6
WALBSP-07031	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-E087D-S099R-S126A-D127E-F128G-N242D	41	2.0	2.8

WALBSP-07032	S039E-P054T-S099R-S126A-D127E-F128G	41	3.6	4.6
WALBSP-04815	A37T-S39E-I43V-A47V-P54T-T56Y-I80V-N85S-E87D-S99R-T114Q-S126A-D127E-F128G-N242D	36	2.0	2.6
WALBSP-07064	A037T-S039E-I043V-A047V-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D	42	2.0	2.4
WALBSP-07005	A037T-S039E-I043V-A047V-T056Y-I080V-E087D-S099R-S126A-D127E-F128G-N242D	42	1.9	2.5
WALBSP-07010	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-D127E-F128G-N242D	42	2.1	2.9
WALBSP-07061	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-D127E-F128G-N242D	42	2.0	2.9
WALBSP-07072	S039E-E087D-S099R-S126A-D127E-F128G	42	2.0	2.8
WALBSP-07035	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-D127E-F128G-N242D	42	2.1	2.7
WALBSP-07038	A037T-S039E-I043V-A047V-P054T-T056Y-	43	1.9	2.8

	I080V-N085S-E087D-S099R-T114Q-S126A-F128G-N242D			
WALBSP-07002	A037T-S039E-S099R-S126A-D127E-F128G	43	3.8	5.5
WALBSP-07067	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-E087D-S099R-T114Q-S126A-D127E-F128G-N242D	43	1.8	2.9
WALBSP-07018	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D	43	1.9	2.8
WALBSP-07020	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-F128G-N242D	43	2.0	3.1
WALBSP-07043	A037T-S039E-I043V-A047V-T056Y-S099R-S126A-D127E-F128G	44	2.0	2.7
WALBSP-07023	A037T-S039E-I043V-A047V-P054T-T056Y-S099R-T114Q-S126A-D127E-F128G	44	2.1	2.7
WALBSP-07070	A037T-S039E-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D	44	2.0	2.9

WALBSP-07071	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-F128G-N242D	44	1.9	3.0
WALBSP-07008	A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-T114Q-S126A-D127E-F128G-N242D	44	0.7	0.8
WALBSP-07025	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S126A-D127E-F128G-N242D	45	0.8	0.7
WALBSP-07009	A037T-S039E-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D	45	2.1	3.2
WALBSP-07075	S039E-S099R-S126A-D127E-F128G	45	1.9	2.8
WALBSP-07017	A037T-S039E-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D	45	2.0	2.8
WALBSP-07069	A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-T114Q-S126A-D127E-F128G-N242D	46	0.8	1.0
WALBSP-07044	A037T-S039E-A047V-T056Y-I080V-N085S-	46	2.1	2.7

	E087D-S099R-S126A-D127E-F128G-N242D			
WALBSP-07100	S039E-S099R-S126A-D127E-F128G-Q256E	47	2.3	3.8
WALBSP-07013	S039E-S099R-S126A-D127E-F128G-N242D	47	2.1	2.6
WALBSP-07139	S039E-S099R-S126A-D127E-F128G-Q200L	48	2.2	3.7
WALBSP-07077	A037T-S039E-I043V-A047V-P054T-T056Y-S099R-T114Q-S126A-D127E-F128G-N242D	51	2.2	3.6
WALBSP-07087	A037T-S039E-I043V-A047V-P054T-T056Y-S099R-S126A-D127E-F128G-N242D	51	2.1	3.6
WALBSP-07138	S039E-N074D-I080V-S099R-S126A-D127E-F128G	53	2.3	3.8
WALBSP-07021	A037T-S039E-I043V-A047V-P054T-T056Y-N085S-E087D-S099R-S126A-D127E-F128G-N242D	54	1.9	2.9
WALBSP-07039	A037T-S039E-I043V-A047V-T056Y-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D	54	2.0	3.0
WALBSP-07053	A037T-S039E-I043V-A047V	54	0.8	0.7
WALBSP-07073	A037T-S039E-I043V-A047V-N242D	54	0.7	0.9

WALBSP-07118	S039E-P054T-N074D-S099R-S126A-D127E-F128G	55	2.3	4.2
WALBSP-07108	S039E-A047V-N074D-S099R-S126A-D127E-F128G	55	2.3	3.3
WALBSP-07057	A037T-S039E-I043V-A047V-P054T-T056Y-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D	55	2.0	2.7
WALBSP-07089	S039E-N074D-S099R-T114Q-S126A-D127E-F128G	55	2.5	3.8
WALBSP-07148	S039E-N074D-N085S-S099R-S126A-D127E-F128G	56	2.3	3.4
WALBSP-07074	A037T-S039E-I043V-A047V-T056Y-N085S-E087D-S099R-S126A-D127E-F128G-N242D	57	1.8	2.5
WALBSP-07079	S039E-N074D-E087D-S099R-S126A-D127E-F128G	57	2.3	ND
WALBSP-07098	S039E-I043V-N074D-S099R-S126A-D127E-F128G	57	ND	3.5
WALBSP-07128	S039E-T056Y-N074D-S099R-S126A-D127E-F128G	58	2.2	3.6
WALBSP-07099	S039E-N074D-S099R-S126A-D127E-F128G-N242D	77	2.2	3.8

As it can be seen from Table 2 the variants of the composition of the invention are considerable better than the parent protease for the removal of egg stains.

CLAIMS

1. A phosphate-free automatic dishwashing cleaning composition comprising:
 - a) from 10% to 50% by weight of the composition of an organic complexing agent system comprising at least 10% by weight of the composition of methyl glycine diacetic acid;
 - b) an amylase; and
 - c) a protease wherein the protease is a variant having at least 90% identity with the amino acid sequence of SEQ ID NO:1 and comprising mutations S039E-S099R-S126A-D127E-F128G.
2. The cleaning composition according to claim 1 wherein the protease is: a variant having at least 90% identity with the amino acid sequence of SEQ ID NO:1 and said variant additionally comprising at least one substitution based upon SEQ ID NO:1 numbering comprising one or more of P54T, X114Q or X114C.
3. The cleaning composition according to claim 2, wherein said variant comprises one or both substitutions P54T and T114Q.
4. The composition according to any one of claims 1 to 3 wherein the protease is: a variant having at least 90% identity with the amino acid sequence of SEQ ID NO:1 and said variant additionally comprising at least one substitution based upon SEQ ID NO:1 numbering comprising one or more of N74D; I80V, R, Y; N85S, C, D, I, K; E87D, C or M211L.
5. The composition according to any one claims 1 to 4 wherein the protease is: a variant having at least 90% identity with the amino acid sequence of SEQ ID NO:1 and said variant additionally comprising at least one substitution based upon SEQ ID NO:1 numbering comprising one or more of T3V, T9R, A15T, V66A, N74D, N85R, N97NE, N97AD, N97D/G, S101A, V102E/I, N116V/R, G157S, Y161A, R164S, T188P, V199I, Q200C/E/I/K/T/V/W/L, Y203W, N212D, M216S/F, Q239R or T249R.

6. The composition according to claim 1 wherein the protease substitutions based upon SEQ ID NO:1 comprise:

- 1) S039E-I080V-S099R-S126A-D127E-F128G-M211L
- 2) S039E-P054T-S099R-S126A-D127E-F128G-M211L
- 5 3) S039E-I043V-S099R-S126A-D127E-F128G-M211L
- 4) S039E-N042R-S099R-S126A-D127E-F128G
- 5) S039E-I080V-S099R-S126A-D127E-F128G
- 6) S039E-S099R-S126A-D127E-F128G-M211L
- 7) S039E-N085S-S099R-S126A-D127E-F128G-M211L
- 10 8) S039E-T056Y-S099R-S126A-D127E-F128G-M211L
- 9) S039E-A047V-S099R-S126A-D127E-F128G-M211L
- 10) S039E-S099R-S126A-D127E-F128G-Y203W
- 11) A037T-S039E-S099R-S126A-D127E-F128G-M211L
- 12) S039E-S099R-S126A-D127E-F128G-V199I
- 15 13) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G
- 14) S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 15) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-
- 20 D127E-F128G
- 16) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G
- 17) S039E-I043V-S099R-S126A-D127E-F128G
- 18) S039E-S099R-S126A-D127E-F128G-N253D
- 25 19) T009E-S039E-S099R-S126A-D127E-F128G
- 20) S039E-S099R-S126A-D127E-F128G-S255W
- 21) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-S099R-S126A-D127E-F128G-N242D
- 22) A037T-S039E-I043V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-
- 30 F128G-N242D
- 23) S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 24) S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D

- 25) S039E-T056Y-S099R-S126A-D127E-F128G
- 26) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G
- 27) A037T-S039E-I043V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 5 28) A037T-S039E-I043V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 29) A037T-S039E-I043V-A047V-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 10 30) A037T-S039E-I043V-A047V-P054T-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 31) A037T-S039E-I043V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 32) S039E-N085S-S099R-S126A-D127E-F128G
- 15 33) A037T-S039E-I043V-A047V-P054T-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 34) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-S099R-T114Q-S126A-D127E-F128G-N242D
- 35) A037T-S039E-I043V-A047V-T056Y-I080V-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 20 36) S039E-S099R-T114Q-S126A-D127E-F128G
- 37) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 38) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-S099R-T114Q-S126A-D127E-F128G-N242D
- 25 39) S039E-A047V-S099R-S126A-D127E-F128G
- 40) A037T-S039E-I043V-A047V-T056Y-I080V-N085S-S099R-S126A-D127E-F128G-N242D
- 41) A037T-S039E-I043V-A047V-T056Y-S099R-T114Q-S126A-D127E-F128G
- 30 42) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-E087D-S099R-S126A-D127E-F128G-N242D
- 43) S039E-P054T-S099R-S126A-D127E-F128G
- 44) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D

- 45) A037T-S039E-I043V-A047V-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 46) A037T-S039E-I043V-A047V-T056Y-I080V-E087D-S099R-S126A-D127E-F128G-N242D
- 5 47) S039E-E087D-S099R-S126A-D127E-F128G
- 48) A037T-S039E-S099R-S126A-D127E-F128G
- 49) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 50) A037T-S039E-I043V-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 10 51) A037T-S039E-I043V-A047V-T056Y-S099R-S126A-D127E-F128G
- 52) A037T-S039E-I043V-A047V-P054T-T056Y-S099R-T114Q-S126A-D127E-F128G
- 53) A037T-S039E-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 15 54) A037T-S039E-A047V-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 55) S039E-S099R-S126A-D127E-F128G
- 56) A037T-S039E-A047V-P054T-T056Y-I080V-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 20 57) A037T-S039E-A047V-T056Y-I080V-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 58) S039E-S099R-S126A-D127E-F128G-Q256E
- 59) S039E-S099R-S126A-D127E-F128G-N242D
- 60) S039E-S099R-S126A-D127E-F128G-Q200L
- 25 61) A037T-S039E-I043V-A047V-P054T-T056Y-S099R-T114Q-S126A-D127E-F128G-N242D
- 62) A037T-S039E-I043V-A047V-P054T-T056Y-S099R-S126A-D127E-F128G-N242D
- 63) S039E-N074D-I080V-S099R-S126A-D127E-F128G
- 64) A037T-S039E-I043V-A047V-P054T-T056Y-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 30 65) A037T-S039E-I043V-A047V-T056Y-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 66) S039E-P054T-N074D-S099R-S126A-D127E-F128G
- 67) S039E-A047V-N074D-S099R-S126A-D127E-F128G

- 68) A037T-S039E-I043V-A047V-P054T-T056Y-N085S-E087D-S099R-T114Q-S126A-D127E-F128G-N242D
- 69) S039E-N074D-S099R-T114Q-S126A-D127E-F128G
- 70) S039E-N074D-N085S-S099R-S126A-D127E-F128G
- 5 71) A037T-S039E-I043V-A047V-T056Y-N085S-E087D-S099R-S126A-D127E-F128G-N242D
- 72) S039E-N074D-E087D-S099R-S126A-D127E-F128G
- 73) S039E-I043V-N074D-S099R-S126A-D127E-F128G
- 74) S039E-T056Y-N074D-S099R-S126A-D127E-F128G or
- 10 75) S039E-N074D-S099R-S126A-D127E-F128G-N242D
7. The composition according to any one of claims 1 to 6 wherein the complexing agent system comprises a complexing agent that is citric acid, methyl glycine diacetic acid, glutamic-N,N-diacetic acid, iminodisuccinic acid, or carboxy methyl inulin or a mixture thereof.
- 15 8. The composition according to claim 7 wherein the complexing agent system comprises a complexing agent of a salt of methyl glycine diacetic acid.
9. The composition according to any one of claims 1 to 8 comprising a bleaching system comprising at least 10% by weight of the composition of bleach, and optionally a bleach activator and/or a bleach catalyst.
10. The composition according to claim 9 wherein the bleaching system comprises percarbonate.
11. The composition according to claim 9 or 10 wherein the bleaching system comprises a bleach activator.
12. The composition according to any one of claims 9 to 11 wherein the bleaching system comprises a bleach catalyst.
13. The composition according to claim 12 wherein the bleaching system comprises a manganese bleach catalyst.

14. The composition according to any one of claims 1 to 13 wherein the composition comprises a dispersant polymer.
15. The composition according to claim 14 wherein the dispersant polymer comprises a polymer comprising both sulfonated and carboxylated monomers.
16. The composition according to any one of claims 1 to 15 comprising:
- i) a bleaching system comprising at least 10% by weight of the composition of percarbonate and optionally a bleach activator and/or a bleach catalyst;
 - ii) a non-ionic surfactant;
 - iii) a dispersant polymer;
 - 5 iv) an amylase; and
 - v) optionally a glass care agent.
17. The composition according to claim 16, wherein the composition comprises the glass care agent.
- 10
18. A method of washing soiled dishware in a dishwasher in the presence of hard water comprising the steps of:
- i) providing the soiled dishware;
 - ii) treating the dishware with a cleaning composition according to any one of claims 1
 - 15 to 17; and
 - iii) optionally rinsing the dishware.
19. Use of a composition according to any one of claims 1 to 16 for automatic dishwashing in hard water.