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(54) **ALPHA-AMYLASE VARIANTS DERIVED FROM THE ALPHA AMYLASE OF CYTOPHAGA SP.
AMYLASEI(CSPAMY2).**

AUS DER ALPHA-AMYLASE-VON CYTOPHAGA-SP. AMYLASEI(CSPAMY2) ABGELEITETE
ALPHA-AMYLASE-VARIANTEN.

VARIANTS D'ALPHA-AMYLASE DÉRIVÉS DE L'ALPHA-AMYLASE DE CYTOPHAGA SP.
AMYLASEI (CSPAMY2).

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• **DECLERCK N ET AL: "Probing structural
determinants specifying high thermostability in
Bacillus licheniformis alpha-amylase",
JOURNAL OF MOLECULAR BIOLOGY,
ACADEMIC PRESS, UNITED KINGDOM, vol. 301,
no. 4, 25 August 2000 (2000-08-25), pages
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Description**FIELD OF THE INVENTION**

5 [0001] Disclosed are compositions and methods relating to variant α -amylases. The variant α -amylases are useful, for example, for starch liquefaction and saccharification, cleaning starchy stains, textile desizing, baking, and brewing.

BACKGROUND

10 [0002] Starch consists of a mixture of amylose (15-30% w/w) and amylopectin (70-85% w/w). Amylose consists of linear chains of α -1,4-linked glucose units having a molecular weight (MW) from about 60,000 to about 800,000. Amylopectin is a branched polymer containing α -1,6 branch points every 24-30 glucose units; its MW may be as high as 100 million.

15 [0003] Sugars from starch, in the form of concentrated dextrose syrups, are currently produced by an enzyme catalyzed process involving: (1) liquefaction (or viscosity reduction) of solid starch with an α -amylase into dextrins having an average degree of polymerization of about 7-10, and (2) saccharification of the resulting liquefied starch (*i.e.* starch hydrolysate) with amyloglucosidase (also called glucoamylase or GA). The resulting syrup has a high glucose content. Much of the glucose syrup that is commercially produced is subsequently enzymatically isomerized to a dextrose/fructose mixture known as isosyrup. The resulting syrup also may be fermented with microorganisms, such as yeast, to produce commercial products including ethanol, citric acid, lactic acid, succinic acid, itaconic acid, monosodium glutamate, gluconates, lysine, other organic acids, other amino acids, and other biochemicals, for example. Fermentation and saccharification can be conducted simultaneously (*i.e.*, an SSF process) to achieve greater economy and efficiency.

20 [0004] α -amylases hydrolyze starch, glycogen, and related polysaccharides by cleaving internal α -1,4-glucosidic bonds at random. α -amylases, particularly from *Bacilli*, have been used for a variety of different purposes, including starch liquefaction and saccharification, textile desizing, starch modification in the paper and pulp industry, brewing, baking, production of syrups for the food industry, production of feedstocks for fermentation processes, and in animal feed to increase digestability. These enzymes can also be used to remove starchy soils and stains during dishwashing and laundry washing.

25 [0005] WO11/80352A1 discloses alpha-amylases, nucleic acids encoding the alpha- amylases and methods of producing and using the alpha- amylases. US7498158B2 discloses variants (mutants) of parent Xermamyl-like alpha-amylases, which have alpha-amylase activity and which exhibit altered properties relative to the parent alpha-amylase. US2011/0033882A1 discloses Variants of *B. licheniformis* alpha-amylase. Declerck et al (J. Mol. Biol. 301, 1041-1057) disclose methods of probing structural determinants specifying high thermostability in bacillus licheniformis alpha-amylase. WO03/14358A2 discloses alpha-amylases which are derived from α -amylases of the bacteria species *Bacillus amyloliquefaciens* and *Bacillus licheniformis* for use in washing and cleaning.

SUMMARY

30 [0006] The present invention provides variant α -amylase polypeptides derived from a parental α -amylase polypeptide, comprising at least one combinable mutation at a productive amino acid position; wherein: (i) the or each combinable mutation is a mutation that improves at least one desirable enzymatic and biochemical property of the variant α -amylase compared to the parental α -amylase, while not significantly decreasing either expression, activity, or stability of the variant α -amylase, compared to the parental α -amylase, (ii) the or each productive position is an amino acid position that can be substituted with a plurality of different amino acid residues, each of which substitutions result in a variant α -amylase that meets the requirements of (i), and (iii) the or each combinable mutation is listed in Lists A, B, C, D, E, or F, or in Table D, which uses SEQ ID NO: 1 for numbering, and wherein the variant α -amylase has at least 60%, at least 70%, at least 80% or at least 90% amino acid sequence identity to the amino acid sequence of SEQ ID NO: 1 or SEQ ID NO: 2, wherein the variant amylase includes a combinable mutation at the productive position corresponding to position 203 of SEQ ID NO: 1 which is substitution to Tyr.

35 [0007] In some embodiments, the combinable mutation which is substitution to Tyr at position 203 SEQ ID NO: 1 for numbering produces a variant amylase wherein the minimum performance indices (PI) relative to the parental amylase for (i) protein expression, (ii) activity, and (iii) detergent stability or thermostability are greater than or equal to 0.9, and the PI for any one of (i), (ii), or (iii) that is greater than or equal to 1.0, whereby the PI for the variant is calculated by comparing the performance or stability of the variant (measured value) and the parental amylase (theoretical value) at the same protein concentration.

40 [0008] In some embodiments, the combinable mutation which is substitution to Tyr at position 203 SEQ ID NO: 1 for numbering produces a variant amylase wherein the minimum performance indices (PI) relative to the parental amylase for (i) protein expression, (ii) activity, and (iii) detergent stability or thermostability are greater than or equal to 0.8, and

the PI for any one of (i), (ii), or (iii) that is greater than or equal to 1.2, whereby the PI for the variant is calculated by comparing the performance or stability of the variant (measured value) and the parental amylase (theoretical value) at the same protein concentration.

[0009] In some embodiments, the combinable mutation which is substitution to Tyr at position 203 SEQ ID NO: 1 for numbering produces a variant amylase wherein the minimum performance indices (PI) relative to the parental amylase for (i) protein expression, (ii) activity, and (iii) detergent stability or thermostability are greater than or equal to 0.5, and the PI for any one of (i), (ii), or (iii) that is greater than or equal to 1.5, whereby the PI for the variant is calculated by comparing the performance or stability of the variant (measured value) and the parental amylase (theoretical value) at the same protein concentration.

[0010] In some embodiments, the or each combinable mutation has a suitability score of +++, +++, or +++, referring to Table C.

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[0012] In some embodiments, the or each combinable mutation has a suitability score of +++, referring to Table C.

[0013] In some embodiments, the or each combinable mutation has a productivity score of 1 or 2, referring to Table B.

[0014] In some embodiments, the variant amylase has a plurality of combinable mutations.

[0015] In some embodiments, the variant amylase further comprises a deletion corresponding to a residue selected from the group consisting of Arg-178, Gly-179, Thr-180, and Gly-181, using SEQ ID NO: 1 for numbering.

[0016] In some embodiments, the variant amylase further comprises deletions corresponding to residues Arg-178 and Gly-179, using SEQ ID NO: 1 for numbering.

[0017] In some embodiments, the variant amylase further comprising a surfactant, wherein the composition is optionally a detergent composition, e.g. a laundry detergent or a laundry detergent additive.

[0018] The present invention also provides methods of saccharifying a composition comprising starch to produce a composition comprising glucose, the methods comprising:

- (i) contacting the solution comprising starch with effective amount of the variant amylase of any of the claims 1-11; and
- (ii) saccharifying the solution comprising starch to produce the composition comprising glucose; wherein the variant amylase catalyzes the saccharification of the starch solution to glucose. The composition comprising starch may comprise liquefied starch, gelatinized starch, or granular starch.

[0019] In some embodiments, saccharification is conducted at a temperature range of about 30°C to about 75°C, optionally wherein the temperature range is 47°C-74°C; and/or wherein saccharification is conducted over a pH range of pH 2.0-7.5, optionally wherein the pH range is pH 3.5-5.5, further optionally wherein the pH range is pH 3.5-4.5.

[0020] In some embodiments, the methods of the invention comprise the step of fermenting the glucose composition to produce an end of fermentation (EOF) product.

[0021] Further compositions and methods that relate to variant amylase polypeptides, and methods of use, thereof are also disclosed herein.

BRIEF DESCRIPTION OF THE SEQUENCES

[0022]

SEQ ID NO: 1 sets forth the amino acid sequence of the mature form of *Cytophaga* sp. amylase (CspAmy2).

SEQ ID NO: 2 sets forth the amino acid sequence of the mature form of a variant *Cytophaga* sp. amylase (CspAmy2-v1) having deletions of both Arginine 178 and Glycine 179.

SEQ ID NO: 3 sets forth the amino acid sequence of the immature/precursor form of the variant form of *Cytophaga* sp. amylase (CspAmy2-v1) having a signal peptide.

SEQ ID NO: 4 sets forth the nucleotide sequence of a synthetic DNA fragment (CspAmy2-v1 DNA) encoding CspAmy2-v1 amylase (SEQ ID NO: 2).

DETAILED DESCRIPTION

[0023] Described are compositions and methods relating to variant amylase enzymes. The variants were discovered by a combination of experimental approaches, as detailed in the appended Examples. The approaches include the use of site evaluation libraries (SELs) and structure-based analysis. Exemplary applications for the variant amylase enzymes are for starch liquefaction and saccharification, for cleaning starchy stains in laundry, dishwashing, and other applications, for textile processing (e.g., desizing), in animal feed for improving digestibility, and for baking and brewing. These and other aspects of the compositions and methods are described in detail, below. The scope of the invention is defined

by the claims.

[0024] Prior to describing the various aspects and embodiments of the present compositions and methods, the following definitions and abbreviations are described.

1. Definitions and Abbreviations

[0025] In accordance with this detailed description, the following abbreviations and definitions apply. Note that the singular forms "a," "an," and "the" include plural referents unless the context clearly dictates otherwise. Thus, for example, reference to "an enzyme" includes a plurality of such enzymes, and reference to "the dosage" includes reference to one or more dosages and equivalents thereof known to those skilled in the art, and so forth.

[0026] The present document is organized into a number of sections for ease of reading; however, the reader will appreciate that statements made in one section may apply to other sections. In this manner, the headings used for different sections of the disclosure should not be construed as limiting.

[0027] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art. The following terms are provided below.

1.1. Abbreviations and Acronyms

[0028] The following abbreviations/acronyms have the following meanings unless otherwise specified:

ABTS	2,2-azino-bis-3-ethylbenzothiazoline-6-sulfonic acid
AE or AEO	alcohol ethoxylate
AES or AEOS	alcohol ethoxysulfate
AkAA	<i>Aspergillus kawachii</i> α -amylase
AnGA	<i>Aspergillus niger</i> glucoamylase
AOS	α -olefinsulfonate
AS	alkyl sulfate
cDNA	complementary DNA
CMC	carboxymethylcellulose
DE	dextrose equivalent
DNA	deoxyribonucleic acid
DP _n	degree of saccharide polymerization having n subunits
ds or DS	dry solids
DTMPA	diethylenetriaminepentaacetic acid
EC	Enzyme Commission
EDTA	ethylenediaminetetraacetic acid
EO	ethylene oxide (polymer fragment)
EOF	End of Fermentation
GA	glucoamylase
GAU/g ds	glucoamylase activity unit/gram dry solids
HFCS	high fructose corn syrup
HgGA	<i>Humicola grisea</i> glucoamylase
IPTG	isopropyl β -D-thiogalactoside
IRS	insoluble residual starch
kDa	kiloDalton
LAS	linear alkylbenzenesulfonate
LAT, BLA	<i>B. licheniformis</i> amylase
MW	molecular weight
MWU	modified Wohlgemuth unit; 1.6×10^{-5} mg/MWU = unit of activity
NCBI	National Center for Biotechnology Information
NOBS	nonanoyloxybenzenesulfonate
NTA	nitroacetic acid
OxAm	Purastar HPAM 5000L (Danisco US Inc.)
PAHBAH	p-hydroxybenzoic acid hydrazide
PEG	polyethyleneglycol
pI	isoelectric point
PI	performance index
ppm	parts per million, e.g., μ g protein per gram dry solid

	PVA	poly(vinyl alcohol)
	PVP	poly(vinylpyrrolidone)
	RCF	relative centrifugal/centripetal force (<i>i.e.</i> , x gravity)
	RNA	ribonucleic acid
5	SAS	alkanesulfonate
	SDS-PAGE	sodium dodecyl sulfate polyacrylamide gel electrophoresis
	SSF	simultaneous saccharification and fermentation
	SSU/g solid	soluble starch unit/gram dry solids
	sp.	species
10	TAED	tetraacetylenediamine
	T _m	melting temperature
	TrGA	<i>Trichoderma reesei</i> glucoamylase
	w/v	weight/volume
	w/w	weight/weight
15	v/v	volume/volume
	wt%	weight percent
	°C	degrees Centigrade
	H ₂ O	water
	dH ₂ O or DI	deionized water
20	dIH ₂ O	deionized water, Milli-Q filtration
	g or gm	grams
	μg	micrograms
	mg	milligrams
	kg	kilograms
25	μL and μl	microliters
	mL and ml	milliliters
	mm	millimeters
	μm	micrometer
	M	molar
30	mM	millimolar
	μM	micromolar
	U	units
	sec	seconds
	min(s)	minute/minutes
35	hr(s)	hour/hours
	DO	dissolved oxygen
	Ncm	Newton centimeter
	ETOH	ethanol
	eq.	equivalents
40	N	normal
	uPWA	variant α-amylase derived from <i>Pyrococcus woesei</i>
	PWA	α-amylase from <i>Pyrococcus woesei</i>
	MWCO	molecular weight cut-off
	SSRL	Stanford Synchrotron Radiation Lightsource
45	PDB	Protein Database
	CAZy	Carbohydrate-Active Enzymes database
	Tris-HCl	tris(hydroxymethyl)aminomethane hydrochloride
	HEPES	4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid

50 1.2. Definitions

[0029] The terms "amylase" or "amylolytic enzyme" refer to an enzyme that is, among other things, capable of catalyzing the degradation of starch. α-Amylases are hydrolases that cleave the α-D-(1→4) O-glycosidic linkages in starch. Generally, α-amylases (EC 3.2.1.1; α-D-(1→4)-glucan glucanohydrolase) are defined as endo-acting enzymes cleaving α-D-(1→4) O-glycosidic linkages within the starch molecule in a random fashion yielding polysaccharides containing three or more (1-4)-α-linked D-glucose units. In contrast, the exo-acting amylolytic enzymes, such as β-amylases (EC 3.2.1.2; α-D-(1→4)-glucan maltohydrolase) and some product-specific amylases like maltogenic α-amylase (EC 3.2.1.133) cleave the polysaccharide molecule from the non-reducing end of the substrate. β-amylases, α-glucosidases (EC

3.2.1.20; α -D-glucoside glucohydrolase), glucoamylase (EC 3.2.1.3; α -D-(1 $\rightarrow$ 4)-glucan glucohydrolase), and product-specific amylases like the maltotetraosidases (EC 3.2.1.60) and the maltohexaosidases (EC 3.2.1.98) can produce malto-oligosaccharides of a specific length or enriched syrups of specific maltooligosaccharides.

[0030] "Enzyme units" herein refer to the amount of product formed per time under the specified conditions of the assay. For example, a "glucoamylase activity unit" (GAU) is defined as the amount of enzyme that produces 1 g of glucose per hour from soluble starch substrate (4% DS) at 60°C, pH 4.2. A "soluble starch unit" (SSU) is the amount of enzyme that produces 1 mg of glucose per minute from soluble starch substrate (4% DS) at pH 4.5, 50°C. DS refers to "dry solids."

[0031] The term "starch" refers to any material comprised of the complex polysaccharide carbohydrates of plants, comprised of amylose and amylopectin with the formula $(C_6H_{10}O_5)_x$, wherein X can be any number. The term includes plant-based materials such as grains, cereal, grasses, tubers and roots, and more specifically materials obtained from wheat, barley, corn, rye, rice, sorghum, brans, cassava, millet, milo, potato, sweet potato, and tapioca. The term "starch" includes granular starch. The term "granular starch" refers to raw, *i.e.*, uncooked starch, *e.g.*, starch that has not been subject to gelatinization.

[0032] The terms, "wild-type," "parental," or "reference," with respect to a polypeptide, refer to a naturally-occurring polypeptide that does not include a man-made substitution, insertion, or deletion at one or more amino acid positions. Similarly, the terms "wild-type," "parental," or "reference," with respect to a polynucleotide, refer to a naturally-occurring polynucleotide that does not include a man-made nucleoside change. However, note that a polynucleotide encoding a wild-type, parental, or reference polypeptide is not limited to a naturally-occurring polynucleotide, and encompasses any polynucleotide encoding the wild-type, parental, or reference polypeptide.

[0033] Reference to the wild-type polypeptide is understood to include the mature form of the polypeptide. A "mature" polypeptide or variant, thereof, is one in which a signal sequence is absent, for example, cleaved from an immature form of the polypeptide during or following expression of the polypeptide.

[0034] The term "variant," with respect to a polypeptide, refers to a polypeptide that differs from a specified wild-type, parental, or reference polypeptide in that it includes one or more naturally-occurring or man-made substitutions, insertions, or deletions of an amino acid. Similarly, the term "variant," with respect to a polynucleotide, refers to a polynucleotide that differs in nucleotide sequence from a specified wild-type, parental, or reference polynucleotide. The identity of the wild-type, parental, or reference polypeptide or polynucleotide will be apparent from context.

[0035] In the case of the present α -amylases, "activity" refers to α -amylase activity, which can be measured as described, herein.

[0036] The term "recombinant," when used in reference to a subject cell, nucleic acid, protein or vector, indicates that the subject has been modified from its native state. Thus, for example, recombinant cells express genes that are not found within the native (non-recombinant) form of the cell, or express native genes at different levels or under different conditions than found in nature. Recombinant nucleic acids differ from a native sequence by one or more nucleotides and/or are operably linked to heterologous sequences, *e.g.*, a heterologous promoter in an expression vector. Recombinant proteins may differ from a native sequence by one or more amino acids and/or are fused with heterologous sequences. A vector comprising a nucleic acid encoding an amylase is a recombinant vector.

[0037] "Combinatorial variants" are variants comprising two or more mutations, *e.g.*, 2, 3, 4, 5, 6, 7, 8, 9, 10, etc., substitutions, deletions, and/or insertions.

[0038] "Combinable mutations" are mutations at any amino acid position that can be used to make combinatorial variants. Combinable mutations improve at least one desired property of the molecule (in this case, an amylase), while not significantly decreasing either expression, activity, or stability. Combinable mutations can be grouped as follows:

Group A: A mutation that produces a variant wherein the minimum performance indices (PI) relative to a defined parental protein for: (i) protein expression, (ii) CS-28 microswatch activity at pH 8 (25°C) or pH10 (50°C), or activity in Ceralpha assay, or activity in corn flour or corn starch application assays, and (iii) detergent stability or thermostability at pH 5.0 or pH 5.7 is greater than or equal to 0.9, and in addition have a PI for any one of these tests that is greater than or equal to 1.0.

Group B: A mutation that produces a variant wherein the minimum performance indices (PI) relative to a defined parental protein for: (i) protein expression, (ii) CS-28 microswatch activity at pH 8 (25°C) or pH10 (50°C), or activity in Ceralpha assay, or activity in corn flour or corn starch application assays, and (iii) detergent stability or thermostability at pH 5.0 or pH 5.7 is greater than or equal to 0.8, and in addition have a PI for any one of these tests that is greater than or equal to 1.2.

Group C: A mutation that produces a variant wherein the minimum performance indices (PI) relative to a defined parental protein for: (i) protein expression, (ii) CS-28 microswatch activity at pH 8 (25°C) or pH10 (50°C), or activity in Ceralpha assay, or activity in corn flour or corn starch application assays, and (iii) detergent stability or thermostability at pH 5.0 or pH 5.7 is greater than or equal to 0.5, and in addition have a PI for any one of these tests that is greater than or equal to 1.5.

[0039] The properties and profiles of combinable mutations are summarized in the following Table.

Table A. Performance properties and profiles for each group of combinable mutations

Group	Performance Index (PI)						
	Expression	Cleaning (pH 8 or 10)	Stability (detergent)	Activity (corn flour or corn starch)	Activity in Ceralpha assay	Thermostability (pH 5.0 or pH 5.7)	Minimum PI in one or more tests
A	≥ 0.9	≥ 0.9	≥ 0.9	≥ 0.9	≥ 0.9	≥ 0.9	X ≥ 1.0
B	≥ 0.8	≥ 0.8	≥ 0.8	≥ 0.8	≥ 0.8	≥ 0.8	X ≥ 1.2
C	≥ 0.5	≥ 0.5	≥ 0.5	≥ 0.5	≥ 0.5	≥ 0.5	X ≥ 1.5

[0040] Preferred combinable mutations are at "productive positions," as described, below. In the case of the present amylases, "activity" refers to amylase activity, which can be measured as described, herein.

[0041] "Productive positions" are amino acid positions that are tolerant to substitution with different amino acid residues, wherein the resulting variants meet a set of performance criteria for combinability, as set forth above. Productive positions can be assigned a Productivity Score as follows: Positions where less than 15% of the substitutions at a given position fall within groups A, B, or C are given a Productivity Score of "1." Positions where less than 30%, but greater than, or equal to 15% of the substitutions at a given position fall within groups A, B, or C are given a Productivity Score of "2." Positions where less than 50%, but greater than, or equal to 30% of the substitutions at a given position fall within groups A, B, or C are given a Productivity Score of "3." Positions where 50% or more of the substitutions at a given position fall within groups A, B, or C are given a Productivity Score of "4." Productivity scores are described in the following Table:

Table B. Performance criteria associated with productivity scores

Productivity Score	Performance criteria for combinability
1	15% of the substitutions at a given position fall within groups A, B, or C.
2	Positions where less than 30%, but greater than, or equal to 15% of the substitutions at a given position fall within groups A, B, or C.
3	50%, but greater than, or equal to 30% of the substitutions at a given position fall within groups A, B, or C.
4	50% or more of the substitutions at a given position fall within groups A, B, or C.

[0042] Preferred productive positions are combinable mutations.

[0043] Suitability score refers to the ability of one or more combinable mutations to be used to make combinatorial variants, based on the performance criteria for combinability (*i.e.*, A, B, and C, as set forth, above) in which each of the mutations fall. A higher suitability score indicates a mutation or mutations that are more suitable for use in making combinatorial variants. Suitability scores are described in the following Table:

Table C. Definitions of suitability scores

Substitutions Occur in Group(s)	Suitability Score
A, B and C	+++++
A and B	++++
A or (B and C)	+++
B	++
C	+

[0044] The terms "recovered," "isolated," and "separated," refer to a compound, protein (polypeptides), cell, nucleic acid, amino acid, or other specified material or component that is removed from at least one other material or component with which it is naturally associated as found in nature. An "isolated" polypeptides, thereof, includes, but is not limited to, a culture broth containing secreted polypeptide expressed in a heterologous host cell.

[0045] The term "purified" refers to material (e.g., an isolated polypeptide or polynucleotide) that is in a relatively pure state, e.g., at least about 90% pure, at least about 95% pure, at least about 98% pure, or even at least about 99% pure.

[0046] The term "enriched" refers to material (e.g., an isolated polypeptide or polynucleotide) that is in about 50% pure, at least about 60% pure, at least about 70% pure, or even at least about 70% pure.

[0047] The terms "thermostable" and "thermostability," with reference to an enzyme, refer to the ability of the enzyme to retain activity after exposure to an elevated temperature. The thermostability of an enzyme, such as an amylase enzyme, is measured by its half-life ($t_{1/2}$) given in minutes, hours, or days, during which half the enzyme activity is lost under defined conditions. The half-life may be calculated by measuring residual α -amylase activity following exposure to (i.e., challenge by) an elevated temperature.

[0048] A "pH range," with reference to an enzyme, refers to the range of pH values under which the enzyme exhibits catalytic activity.

[0049] The terms "pH stable" and "pH stability," with reference to an enzyme, relate to the ability of the enzyme to retain activity over a wide range of pH values for a predetermined period of time (e.g., 15 min., 30 min., 1 hour).

[0050] The term "amino acid sequence" is synonymous with the terms "polypeptide," "protein," and "peptide," and are used interchangeably. Where such amino acid sequences exhibit activity, they may be referred to as an "enzyme." The conventional one-letter or three-letter codes for amino acid residues are used, with amino acid sequences being presented in the standard amino-to-carboxy terminal orientation (i.e., N→C).

[0051] The term "nucleic acid" encompasses DNA, RNA, heteroduplexes, and synthetic molecules capable of encoding a polypeptide. Nucleic acids may be single stranded or double stranded, and may be chemical modifications. The terms "nucleic acid" and "polynucleotide" are used interchangeably. Because the genetic code is degenerate, more than one codon may be used to encode a particular amino acid, and the present compositions and methods encompass nucleotide sequences that encode a particular amino acid sequence. Unless otherwise indicated, nucleic acid sequences are presented in 5'-to-3' orientation.

[0052] "Hybridization" refers to the process by which one strand of nucleic acid forms a duplex with, i.e., base pairs with, a complementary strand, as occurs during blot hybridization techniques and PCR techniques. Stringent hybridization conditions are exemplified by hybridization under the following conditions: 65°C and 0.1X SSC (where 1X SSC = 0.15 M NaCl, 0.015 M Na₃ citrate, pH 7.0). Hybridized, duplex nucleic acids are characterized by a melting temperature (T_m), where one-half of the hybridized nucleic acids are unpaired with the complementary strand. Mismatched nucleotides within the duplex lower the T_m . A nucleic acid encoding a variant α -amylase may have a T_m reduced by 1°C - 3°C or more compared to a duplex formed between the nucleotide of SEQ ID NO: 2 and its identical complement.

[0053] A "synthetic" molecule is produced by *in vitro* chemical or enzymatic synthesis rather than by an organism.

[0054] The terms "transformed," "stably transformed," and "transgenic," used with reference to a cell means that the cell contains a non-native (e.g., heterologous) nucleic acid sequence integrated into its genome or carried as an episome that is maintained through multiple generations.

[0055] The term "introduced" in the context of inserting a nucleic acid sequence into a cell, means "transfection", "transformation" or "transduction," as known in the art.

[0056] A "host strain" or "host cell" is an organism into which an expression vector, phage, virus, or other DNA construct, including a polynucleotide encoding a polypeptide of interest (e.g., an amylase) has been introduced. Exemplary host strains are microorganism cells (e.g., bacteria, filamentous fungi, and yeast) capable of expressing the polypeptide of interest and/or fermenting saccharides. The term "host cell" includes protoplasts created from cells.

[0057] The term "heterologous" with reference to a polynucleotide or protein refers to a polynucleotide or protein that does not naturally occur in a host cell.

[0058] The term "endogenous" with reference to a polynucleotide or protein refers to a polynucleotide or protein that occurs naturally in the host cell.

[0059] The term "expression" refers to the process by which a polypeptide is produced based on a nucleic acid sequence. The process includes both transcription and translation.

[0060] A "selective marker" or "selectable marker" refers to a gene capable of being expressed in a host to facilitate selection of host cells carrying the gene. Examples of selectable markers include but are not limited to antimicrobials (e.g., hygromycin, bleomycin, or chloramphenicol) and/or genes that confer a metabolic advantage, such as a nutritional advantage on the host cell.

[0061] A "vector" refers to a polynucleotide sequence designed to introduce nucleic acids into one or more cell types. Vectors include cloning vectors, expression vectors, shuttle vectors, plasmids, phage particles, cassettes and the like.

[0062] An "expression vector" refers to a DNA construct comprising a DNA sequence encoding a polypeptide of interest, which coding sequence is operably linked to a suitable control sequence capable of effecting expression of the DNA in a suitable host. Such control sequences may include a promoter to effect transcription, an optional operator sequence to control transcription, a sequence encoding suitable ribosome binding sites on the mRNA, enhancers and sequences which control termination of transcription and translation.

[0063] The term "operably linked" means that specified components are in a relationship (including but not limited to

juxtaposition) permitting them to function in an intended manner. For example, a regulatory sequence is operably linked to a coding sequence such that expression of the coding sequence is under control of the regulatory sequences.

[0064] A "signal sequence" is a sequence of amino acids attached to the N-terminal portion of a protein, which facilitates the secretion of the protein outside the cell. The mature form of an extracellular protein lacks the signal sequence, which is cleaved off during the secretion process.

[0065] "Biologically active" refer to a sequence having a specified biological activity, such an enzymatic activity.

[0066] The term "specific activity" refers to the number of moles of substrate that can be converted to product by an enzyme or enzyme preparation per unit time under specific conditions. Specific activity is generally expressed as units (U)/mg of protein.

[0067] As used herein, "water hardness" is a measure of the minerals (e.g., calcium and magnesium) present in water.

[0068] A "swatch" is a piece of material such as a fabric that has a stain applied thereto. The material can be, for example, fabrics made of cotton, polyester or mixtures of natural and synthetic fibers. The swatch can further be paper, such as filter paper or nitrocellulose, or a piece of a hard material such as ceramic, metal, or glass. For amylases, the stain is starch based, but can include blood, milk, ink, grass, tea, wine, spinach, gravy, chocolate, egg, cheese, clay, pigment, oil, or mixtures of these compounds.

[0069] A "smaller swatch" is a section of the swatch that has been cut with a single hole punch device, or has been cut with a custom manufactured 96-hole punch device, where the pattern of the multi-hole punch is matched to standard 96-well microtiter plates, or the section has been otherwise removed from the swatch. The swatch can be of textile, paper, metal, or other suitable material. The smaller swatch can have the stain affixed either before or after it is placed into the well of a 24-, 48- or 96-well microtiter plate. The smaller swatch can also be made by applying a stain to a small piece of material. For example, the smaller swatch can be a stained piece of fabric 5/8" or 0.25" in diameter. The custom manufactured punch is designed in such a manner that it delivers 96 swatches simultaneously to all wells of a 96-well plate. The device allows delivery of more than one swatch per well by simply loading the same 96-well plate multiple times. Multi-hole punch devices can be conceived of to deliver simultaneously swatches to any format plate, including but not limited to 24-well, 48-well, and 96-well plates. In another conceivable method, the soiled test platform can be a bead made of metal, plastic, glass, ceramic, or another suitable material that is coated with the soil substrate. The one or more coated beads are then placed into wells of 96-, 48-, or 24-well plates or larger formats, containing suitable buffer and enzyme.

[0070] "A cultured cell material comprising an amylase" or similar language, refers to a cell lysate or supernatant (including media) that includes an amylase as a component. The cell material may be from a heterologous host that is grown in culture for the purpose of producing the amylase.

[0071] "Percent sequence identity" means that a particular sequence has at least a certain percentage of amino acid residues identical to those in a specified reference sequence, when aligned using the CLUSTAL W algorithm with default parameters. See Thompson et al. (1994) Nucleic Acids Res. 22:4673-4680. Default parameters for the CLUSTAL W algorithm are:

Gap opening penalty:	10.0
Gap extension penalty:	0.05
Protein weight matrix:	BLOSUM series
DNA weight matrix:	IUB
Delay divergent sequences %:	40
Gap separation distance:	8
DNA transitions weight:	0.50
List hydrophilic residues:	GPSNDQEKR
Use negative matrix:	OFF
Toggle Residue specific penalties:	ON
Toggle hydrophilic penalties:	ON
Toggle end gap separation penalty	OFF.

[0072] Deletions are counted as non-identical residues, compared to a reference sequence. Deletions occurring at either termini are included. For example, a variant with five amino acid deletions of the C-terminus of the mature CspAmy2 polypeptide of SEQ ID NO: 1 would have a percent sequence identity of 99% ($612 / 617$ identical residues $\times 100$, rounded to the nearest whole number) relative to the mature polypeptide. Such a variant would be encompassed by a variant having "at least 99% sequence identity" to a mature amylase polypeptide.

[0073] "Fused" polypeptide sequences are connected, i.e., operably linked, via a peptide bond between two subject polypeptide sequences.

[0074] The term "filamentous fungi" refers to all filamentous forms of the subdivision Eumycotina, particularly Pezizomycotina species.

[0075] The term "degree of polymerization" (DP) refers to the number (n) of anhydro-glucopyranose units in a given saccharide. Examples of DP1 are the monosaccharides glucose and fructose. Examples of DP2 are the disaccharides maltose and sucrose. The term "DE," or "dextrose equivalent," is defined as the percentage of reducing sugar, *i.e.*, D-glucose, as a fraction of total carbohydrate in a syrup.

[0076] The term "dry solids content" (ds) refers to the total solids of a slurry in a dry weight percent basis. The term "slurry" refers to an aqueous mixture containing insoluble solids.

[0077] The phrase "simultaneous saccharification and fermentation (SSF)" refers to a process in the production of biochemicals in which a microbial organism, such as an ethanologenic microorganism, and at least one enzyme, such as an amylase, are present during the same process step. SSF includes the contemporaneous hydrolysis of starch substrates (granular, liquefied, or solubilized) to saccharides, including glucose, and the fermentation of the saccharides into alcohol or other biochemical or biomaterial in the same reactor vessel.

[0078] An "ethanologenic microorganism" refers to a microorganism with the ability to convert a sugar or oligosaccharide to ethanol.

[0079] The term "fermented beverage" refers to any beverage produced by a method comprising a fermentation process, such as a microbial fermentation, *e.g.*, a bacterial and/or fungal fermentation. "Beer" is an example of such a fermented beverage, and the term "beer" is meant to comprise any fermented wort produced by fermentation/brewing of a starch-containing plant material. Often, beer is produced exclusively from malt or adjunct, or any combination of malt and adjunct. Examples of beers include: full malted beer, beer brewed under the "Reinheitsgebot," ale, India pale ale, lager, pilsner, bitter, Happoshu (second beer), third beer, dry beer, near beer, light beer, low alcohol beer, low calorie beer, porter, bock, dopplebock, stout, porter, malt liquor, non-alcoholic beer, non-alcoholic malt liquor and the like, but also alternative cereal and malt beverages such as fruit flavored malt beverages, *e.g.*, citrus flavored, such as lemon-, orange-, lime-, or berry-flavored malt beverages, liquor flavored malt beverages, *e.g.*, vodka-, rum-, or tequila-flavored malt liquor, or coffee flavored malt beverages, such as caffeine-flavored malt liquor, and the like.

[0080] The term "malt" refers to any malted cereal grain, such as malted barley or wheat.

[0081] The term "adjunct" refers to any starch and/or sugar containing plant material that is not malt, such as barley or wheat malt. Examples of adjuncts include common corn grits, refined corn grits, brewer's milled yeast, rice, sorghum, refined corn starch, barley, barley starch, dehusked barley, wheat, wheat starch, torrefied cereal, cereal flakes, rye, oats, potato, tapioca, cassava and syrups, such as corn syrup, sugar cane syrup, inverted sugar syrup, barley and/or wheat syrups, and the like.

[0082] The term "mash" refers to an aqueous slurry of any starch and/or sugar containing plant material, such as grist, *e.g.*, comprising crushed barley malt, crushed barley, and/or other adjunct or a combination thereof, mixed with water later to be separated into wort and spent grains.

[0083] The term "wort" refers to the unfermented liquor run-off following extracting the grist during mashing.

[0084] "Iodine-positive starch" or "IPS" refers to (1) amylose that is not hydrolyzed after liquefaction and saccharification, or (2) a retrograded starch polymer. When saccharified starch or saccharide liquor is tested with iodine, the high DPn amylose or the retrograded starch polymer binds iodine and produces a characteristic blue color. The saccharide liquor is thus termed "iodine-positive saccharide," "blue saccharide," or "blue sac."

[0085] The terms "retrograded starch" or "starch retrogradation" refer to changes that occur spontaneously in a starch paste or gel on ageing.

[0086] The term "about" refers to $\pm 15\%$ to the referenced value.

2. α -Amylase variants

[0087] An aspect of the present compositions and methods is variant amylase enzymes which are exclusively as defined in the appended claims and which have been discovered using a combination of experimental approaches, including the use of site evaluation libraries (SELs) and structure-based analysis.

2.1 α -Amylase variants based on an SEL library of *Cytophaga* sp. α -amylase

[0088] An α -amylase from *Cytophaga* sp. (herein, CspAmy2 amylase") has been previously described (Jeang, C-L et al. (2002) Applied and Environmental Microbiology, 68:3651-54). The amino acid sequence of the mature CspAmy2 amylase polypeptide is shown, below (SEQ ID NO: 1):

AATNGTMMQY FEWYVPNDGQ QWNRLRTDAP YLSSVGITAV WTPPAYKGTG
 QADVGYGPDY LYDLGEFNQK GTVRTKYGTK GELKSAVNLT HSNIGQVYGD
 5 VVMNHKAGAD YTENVTADEV NPSNRNQETS GEYNIQAWTG FNFPGRGTTY
 SNFKWQWFHF DGTDWDQSRS LSRIFKFRGT GKAWDWEVSS ENGNYDYLMY
 ADIDYDHPDV VNEMKKWGVW YANVGLDGY RLDVAVKHIF SFLKDWVDNA
 10 RAATGKEMFT VGEYWQNDLG ALNNYLAKVN YNQSLFDAPL HYNFYAASTG
 GGYDMRNIL NNTLVASNPT KAVTLVENHD TQPGQSLEST VQPFKPLAY
 AFILTRSGGY PSVFYGDYMG TKGTTTREIP ALKSKIEPLL KARKDYAYGT
 QRDYIDNPDV IGWTREGDST KAKSGLATVI TDGPGGSKRM YVGTSNAGEI
 15 WYDLTGNRD KITIGSDGYA TFPVNGGSVS VWVQQ

[0089] In SEQ ID NO: 1, Arginine 178 and Glycine 179 are underlined. A variant of the *Cytophaga* sp. α -amylase having a deletion of both Arginine 178 and Glycine 179 (herein, CspAmy2-v1) has also been described (Shiau, R-J. et al. (2003) Applied and Environmental Microbiology, 69:2383-85). The amino acid sequence of the mature CspAmy2-v1 amylase polypeptide is shown, below, as SEQ ID NO: 2:

AATNGTMMQY FEWYVPNDGQ QWNRLRTDAP YLSSVGITAV WTPPAYKGTG
 QADVGYGPDY LYDLGEFNQK GTVRTKYGTK GELKSAVNLT HSNIGQVYGD
 25 VVMNHKAGAD YTENVTADEV NPSNRNQETS GEYNIQAWTG FNFPGRGTTY
 SNFKWQWFHF DGTDWDQSRS LSRIFKFTGK AWDWEVSS ENGYDYLMYAD
 IDYDHPDVVN EMKKWGVWYA NEVGLDGYRL DAVKHKIFSF LKDWVDNARA
 30 ATGKEMFTVG EYWQNDLGAL NNYLAKVNYN QSLFDAPLHY NFYAASTGGG
 YYDMRNILNN TLVASNPTKA VTLVENHDTQ PGQSLESTVQ PWFKPLAYAF
 ILTRSGGYPS VFYGDYMGTK GTTTREIPAL KSKIEPLLKA RKDYAYGTQR
 35 DYIDNPDVIG WTREGDSTKA KSGLATVITD GPGGSKRMVY GTSNAGEIYW
 DLTGNRTDKI TIGSDGYATF PVNGGSVSVW VQQ

[0090] α -amylase variants that include combinable mutations were identified by making a site evaluation library (SEL) based on CspAmy2-v1 and testing the resulting variants for corn flour and corn starch hydrolysis, thermostability assay, cleaning performance, detergent stability, activity in a standardized amylases assay, and expression levels, the detailed procedures for which are described in the Examples or otherwise known. Each variant was assayed for the different enzymatic and biochemical properties, and characterized by a performance index (PI) value, which compared the relative performance of the variant to CspAmy2-v1 amylase for each performance criteria. A PI that is greater than 1 (*i.e.*, PI>1) indicated improved performance by a variant as compared to CspAmy2-v1, while a PI of 1 (*i.e.*, PI=1) indicated a variant that performed the same as the CspAmy2-v1, and a PI that is less than 1 (*i.e.*, PI<1) indicated a variant that performed worse than the CspAmy2-v1. PI values were then used to identify combinable mutations and productive positions.

[0091] Combinable mutations are mutations at any amino acid position that improve at least one desired property of the molecule, while not significantly decreasing expression, activity, or stability. Combinable mutations are assigned to one of three Groups (*i.e.*, A, B, or C), as set forth, herein. Preferred combinable mutations are at productive positions. Productive positions are amino acid positions that are tolerant to substitution with different amino acid residues, wherein the resulting variants meet a set of performance criteria for combinability, as set forth herein.

[0092] Combinable mutations and productive positions are not to be confused with previously-identified, single-site mutations, some of which have subsequently been found by trial and error to work in combination with other mutations. Previously-identified, single-site mutations are invariably "winners" with respect to improving any one performance or stability feature. While this makes them attractive mutations to include in variant amylases, these "winners" tend to adversely affect other performance or stability features of the variants, which often requires making additional mutations

to correct the defects.

[0093] In contrast, combinable mutations may be only incrementally beneficial in improving any one performance or stability feature of an variant amylase. However, they are carefully selected to be minimally detrimental to other desired performance or stability features, making them well suited for use in combination with other combinable mutations to construct variant amylases having desired improved enzymatic and biochemical properties without being crippled in others, resulting in robust variants having a good balance of performance, stability, and expression potential.

[0094] Further based on measured enzymatic and biochemical properties of the variant amylases, the suitability scores of the different mutations for making combinatorial variants were determined. The suitability score refers to the ability of one or more combinable mutations to be used to make combinatorial variants, based on the performance criteria for combinability (*i.e.*, A, B, and C, as set forth, above), in which each of the mutations fall.

[0095] The suitability scores of individual substitutions in CspAmy2-v1 are shown in Table D. The position numbering is based on the amino acid sequence of the mature CspAmy2 polypeptide (SEQ ID NO: 1). Wild-type residues at the indicated positions are given a suitability score of +++. Substitutions more likely to be combinable with other mutations are given a suitability score of +++++, or even ++++++. In general, preferred suitability scores are +++, +++++, or ++++++, +++++ or ++++++, or even ++++++.

Table D: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)*	(++++)	(+++++)
1		I	A	EGKNQRTVY	
2			AEGHKNPQ RSY		
3			TADFGMPQ RS		
4	T	Q	N		
5			G		ACDEFHIKL MNPQRSTVY
7			MI		
8			MF		
11			F	Y	
15			VN	ILS	CT
20			QE		
21			QLW	T	
23	DF		NSWY	AEHKMQTV	
26			RK		
27			TDEFGHIKL MNQSWY	R	
28			DAEN		
30			PHT	ACFLRWY	DEGKNS
31		M	YEKR	FHW	
33			SDEGHNQR	K	
35			VHIMN		

(continued)

Table D: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)*	(++++)	(+++++)
38			TS	DN	
39			AS		
40			VI		
42			TACILMV		
45			APS		
46			YFMT		
48		A	G		
49	M	CGHY	TS	ADEFIKLNV	
50			SEK	D	
51			QS		
52			AFGHIKNQS TW		
54	RS	G	VLQ	CDIN	AET
57	K		G		
58			P	C	
68			NACEFMSY		
70	NW	R	K		
71			GAN		
72	H		TGS		
73			VT		
75	C		T		
81			GIPTV	ADFHKLMNSY	EQRW
82			EQ		
83			LF		
84			KIQV		
85			SACDEHKM NQR	L	
87		I	V		
88			NADEQRT		H
89			TCDEHMQ RSV		
92	L		SAMRV		
93	DT		NM		
94			GN		

(continued)

Table D: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)*	(++++)	(+++++)
96	I		Q		
97			VI		
98	W		YF		
101			VI		
103			MV	I	
104	D		N		
106	AIV		K		
107	G		A		
108	KR		GAS		
109			AP		
111		M	YW	ACDGHKLNQRST V	F
112			TFGIVW	ACDELMPQRY	
113	Y		ED		
114			NG		
115		A	V	IM	
116			T	ACDEGHIKLMNS VW	
117		C	AS		
118	Q	LM	VACFIKNS	R	
119	S		E		
120			V	C	
121		S	NKR		
122		T	P	AKQR	
123			S	ACEGHKNQTY	R
124			ND		
126	Y		N		
127		CY	QAM	HIKRTV	E
128			ES	GKVY	I
129		AFG	TQY	HIKLRSV	
130		AIRV	ST	GH	
131		FM	GT	AHIKNW	PQ
132			ENS	ACGHILMRTVY	PQ
133		FN	YE	ADHLTV	K
134			NCFY	DGHMPQST	

(continued)

Table D: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)*	(++++)	(+++++)
135	HR	M	IV		
136	W		QY	AFGHIK	NT
137		V	A		
138	PR	GV	W	AFKLMQSTY	DH
140	M	D	GCEFHILTV Y	S	KN
141	H		F	W	
142		CV	NEY	FGHIKLQRW	ADMST
144	DQ	F	PCHM	AGKLNRY	IT
145	LWY	IMV	GEQT	AHKNRS	
147			GCLV	EHKNQR	I
148			TAGHIKLSW Y		
149			TACDEHIKM NRSWY	L	
150	DPQ	GM	Y	FHW	I
151			SD		
152		F	NW	ACEGHLMRSTV	DQ
153			F	H	WY
154			KAEHNRST	F	
156	FGILMV		QDSTY	H	K
157	N		W		
158	P		FADEIRY	LNSV	CGHTW
160	C		F		
163	F	CMNQ V	T	DLS	
164		F	DN		
165	P	H	WF		
166	ACGKM		D		
167			Q	ACDGHKNRSTVY	
168		C	SRW	DEGIKMNTV	L
169	EQW		RDMT	CHKLY	A
170			S	CGHNRT	

(continued)

Table D: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)*	(++++)	(+++++)
171			L	FGIMRVWY	HKNQS
172			SACDEHKNR T		
173			RK		W
174		L	I		
175	AD	S	FLMV	HWY	
176	L		K		
177	G		FH		
182	H		K		
183			AE	K	R
187			E	V	P
189	A		SC	D	
190			S		P
191			EACLMNT	I	
192	S	M	NR		FHY
193			GACFIKST	HRV	
195		D	Y		
198	CG		L		A
200	EL		Y		
201	L		A		
203	HNQ	A	I	CV	FLMY
206	I	CY	DM		AQ
207		AKR	HF	DEMSY	N
208			PAEHKLNQR ST	D	
209		CIK	D	G	
210		S	V		
211		FS	V	CEIMNT	DLQ
212			NY	ACDEHILMV	GQRST
215		LY	KRT	EFMNQ	
216			KR		
219			VEL	I	
222		T	A		
225			VL		
226		C	GKQ	E	

(continued)

Table D: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)*	(++++)	(+++++)
227			LY		
232			L	R	V
235		LT	VA	C	
238	M		I	LQR	P
239	M	Q	K	C	
240		D	F		
241		T	SFR	P	ACDEGHIKL NQVWY
242	E	Y	FV		I
243			LAST	IM	
244		Y	KT	ACHMNQRS	
245			DE		
246			WF		
248	A		DQ	N	
249			NKR	ACDEFGMST	QW
250			AS		
251	ADKLMN QST		R		
252			ACDFHMR		
253			ACDELMNQST	RV	
254		F	TILM		
256			KR		
257	ACGRVY	FHK	E	LQS	
258			MCI	FLY	
259	P		F		
260	Y		T	S	AL
262			G	A	
266	EK	HLN	Q	CFGMRW	ADIST
267			ND		
269	A	V	LI		
270		CIQV	G	ADFHKPRSWY	
271			ACS		
273			NDEILMQS	HK	
275			YF		

(continued)

Table D: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)*	(++++)	(+++++)
276			LM		
277		D	ACNW	FHILMQTY	EK
280			NCY	ADGHKLQT	E
281		G	YA	D	
283			QT	HV	
284	EF		SHY	KMRW	
285			LA		
286	L		F	M	
288			A	V	
295			YLQ	H	
296		FIMQS T	ADHKR	E	
299			TEI	R	
300		Q	GAKL	R	
301			GT	AFHKMQRSY	
302		S	G		
303		F	Y	IW	ARTV
307			RQS		
308			NV	ADGLQTY	CEFHMR
310			LDET		
311		E	NDHKQSV		
312			NDG		
313		AS	T		
316			AKQS		
317	W	A	SDGLT	CEMQY	HK
318			NFILVW	ACDEKMQRSY	H
320	PV	M	TCGKY	AHNQRW	DE
321			KHR		
325			LFIMV		
327			ED		
335	G		Q		
336			SAD		
339	Q		S		
342			QEL		
343	AMW		P		

(continued)

Table D: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)*	(++++)	(+++++)
348			LCGS	Q	
349	W		AS	G	
357		Y	SACDELMN V	Q	
358			GDEQS		
360	CEHV		YI	F	L
362	EQT		S	AC	IV
363			VL	I	
368			MLWY		
372			KADHMNR		
374			TAKNPQS		
375			TS		
376		H	TAGNQSY	K	
377	ILMV	C	R	H	ADGKSTY
378			E	Q	
384		D	SEGN	HP	
385			KAE		
388		L	PCDKRS	I	
390			LC	I	
391			KFTVY	EL	
392			AG		
394	E		KHM		
395		IKS	DC	EFMQRWY	
396			YCFK		
397			AS		
400			T	ACDEMNQRSWY	HKL
401			QM		
402			RFKLSTVWY	Q	
403			DS		
404			YW		
405			IL		
407			NACDEGHQ S		
408	S	Y	PEHMQR	KVW	

(continued)

Table D: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)*	(++++)	(+++++)
409		N	D		
410		E	VIKLMSY		
414			TAS		
416		MT	E	CFHKLNRSV	DGQ
418			DACEFGHIK LMNQRSTW Y		
419			SAEGKPQRT VY		
420			TFI	ACDEGHPQRSVW Y	L
421			KACFGHILR SWY	N	
422			A	EIKLMNPQRSTV WY	CD
423	P		KV	ACEFGMNQWY	DS
424			SA		
426			LC		
427			A	C	GS
429			VC	LM	
430		C	IL	EM	
431		D	TA		
433	HS	A	GN		
434		M	PNR	DQS	
435		I	GACEQRSTY	LVN	
437			SADEFHKLQ Y		
441			YCKLNQRS W		
442		L	VAIT		

(continued)

Table D: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)*	(++++)	(+++++)
444			TAEFHIKLM NPQVY		
445			SACDHMRT VW		
447			AGKQRSTV		
448			GEL	AFHKNQTY	DW
449			EQ		
450			IDLP	ACFHKQRTV	N
452			YAILMSVW		
454			LAEFHKS	CIMQTVY	
455			TIV	ACLMS	
456			G	ACDEFHKL MNRS TWY	
458			RDEFSW	ACIN	MY
459			TACDFGLSV W		
460			DEHN		
461			KADGILMNP SY		
462			IV		
463			TCEFIKLMN VY		
464			IV		
465			GAMNPQ		
466			SCFN	ADEHILMRVWY	KT
467			DN		
469			YCFISV	L	
470			AG		
471	K		TADEFGHIN PQW		
474			VC		

(continued)

Table D: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)*	(++++)	(+++++)
475			NFIP	ADGHKLMSTV	
476	DE		G		ACHKNPQRS TVY
477	P		G		ADEHIKQRST VY
479			VCHW		
480			SG		
481			VACIL		
482			WY		
483			VI		
484		AHK	Q		
485			QFR	ADEHIKLMNPTV Y	

* The first listed amino acid residue is the wild-type residue.

[0096] While evaluating mutations based on suitability score represents one refined aspect of the present compositions and methods, the identification of productive positions, which are tolerant to substitution with different amino acid residues, represents a number of significant embodiments.

[0097] Each productive position identified in the following lists with specified criteria, and each substitution identified in parenthesis following the numerical position identifier in each of these lists, represents a mutation, identified by experimental data, that either directly contributes to the performance of an amylase variant, or is determined to be combinable with other mutations to produce a amylase variant with improved performance.

[0098] The productive positions in CspAmy2-v1 that fall within the previously described Productivity Scores of "1," "2," "3," and "4," and the substitutions within those positions that are combinable, are listed below. The position numbering is based on mature CspAmy2 polypeptide (SEQ ID NO: 1).

LIST A:

[0099]

1(A,E,G,I,K,N,Q,R,T,V,Y); 2(A,E,G,H,K,N,P,Q,R,S,Y); 3(T,A,D,F,G,M,P,Q,R,S); 4(N,Q,T);
 5(G,A,C,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,Y); 7(M,I); 8(M,F); 11(F,Y); 15(V,C,I,L,N,S,T);
 20(Q,E); 21(Q,L,T,W); 23(N,A,D,E,F,H,K,M,Q,S,T,V,W,Y); 26(R,K);
 27(T,D,E,F,G,H,I,K,L,M,N,Q,R,S,W,Y); 28(D,A,E,N);
 30(P,A,C,D,E,F,G,H,K,L,N,R,S,T,W,Y); 31(Y,E,F,H,K,M,R,W); 33(S,D,E,G,H,K,N,Q,R);
 35(V,H,I,M,N); 38(T,D,N,S); 39(A,S); 40(V,I); 42(T,A,C,I,L,M,V); 45(A,P,S); 46(Y,F,M,T);
 48(G,A); 49(T,A,C,D,E,F,G,H,I,K,L,M,N,S,V,Y); 50(S,D,E,K); 51(Q,S);
 52(A,F,G,H,I,K,N,Q,S,T,W); 54(V,A,C,D,E,G,I,L,N,Q,R,S,T); 57(G,K); 58(P,C);
 68(N,A,C,E,F,M,S,Y); 70(K,N,R,W); 71(G,A,N); 72(T,G,H,S); 73(V,T); 75(T,C);
 81(G,A,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,W,Y); 82(E,Q); 83(L,F); 84(K,I,Q,V);
 85(S,A,C,D,E,H,K,L,M,N,Q,R); 87(V,I); 88(N,A,D,E,H,Q,R,T); 89(T,C,D,E,H,M,N,Q,R,S,V);
 92(S,A,L,M,R,V); 93(N,D,M,T); 94(G,N); 96(Q,I); 97(V,I); 98(Y,F,W); 101(V,I); 103(M,I,V);
 104(N,D); 106(K,A,I,V); 107(A,G); 108(G,A,K,R,S); 109(A,P);

111(Y,A,C,D,F,G,H,K,L,M,N,Q,R,S,T,V,W); 112(T,A,C,D,E,F,G,I,L,M,P,Q,R,V,W,Y);
 113(E,D,Y); 114(N,G); 115(V,A,I,M); 116(T,A,C,D,E,G,H,I,K,L,M,N,S,V,W); 117(A,C,S);
 118(V,A,C,F,I,K,L,M,N,Q,R,S); 119(E,S); 120(V,C); 121(N,K,R,S); 122(P,A,K,Q,R,T);
 123(S,A,C,E,G,H,K,N,Q,R,T,Y); 124(N,D); 126(N,Y); 127(Q,A,C,E,H,I,K,M,R,T,V,Y);
 5 128(E,G,I,K,S,V,Y); 129(T,A,F,G,H,I,K,L,Q,R,S,V,Y); 130(S,A,G,H,I,R,T,V);
 131(G,A,F,H,I,K,M,N,P,Q,T,W); 132(E,A,C,G,H,I,L,M,N,P,Q,R,S,T,V,Y);
 133(Y,A,D,E,F,H,K,L,N,T,V); 134(N,C,D,F,G,H,M,P,Q,S,T,Y); 135(I,H,M,R,V);
 136(Q,A,F,G,H,I,K,N,T,W,Y); 137(A,V); 138(W,A,D,F,G,H,K,L,M,P,Q,R,S,T,V,Y);
 140(G,C,D,E,F,H,I,K,L,M,N,S,T,V,Y); 141(F,H,W);
 10 142(N,A,C,D,E,F,G,H,I,K,L,M,Q,R,S,T,V,W,Y); 144(P,A,C,D,F,G,H,I,K,L,M,N,Q,R,T,Y);
 145(G,A,E,H,I,K,L,M,N,Q,R,S,T,V,W,Y); 147(G,C,E,H,I,K,L,N,Q,R,V);
 148(T,A,G,H,I,K,L,S,W,Y); 149(T,A,C,D,E,H,I,K,L,M,N,R,S,W,Y);
 150(Y,D,F,G,H,I,M,P,Q,W); 151(S,D); 152(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,W);
 153(F,H,W,Y); 154(K,A,E,F,H,N,R,S,T); 156(Q,D,F,G,H,I,K,L,M,S,T,V,Y); 157(W,N);
 15 158(F,A,C,D,E,G,H,I,L,N,P,R,S,T,V,W,Y); 160(F,C); 163(T,C,D,F,L,M,N,Q,S,V); 164(D,F,N);
 165(W,F,H,P); 166(D,A,C,G,K,M); 167(Q,A,C,D,G,H,K,N,R,S,T,V,Y);
 168(S,C,D,E,G,I,K,L,M,N,R,T,V,W); 169(R,A,C,D,E,H,K,L,M,Q,T,W,Y);
 170(S,C,G,H,N,R,T); 171(L,F,G,H,I,K,M,N,Q,R,S,V,W,Y); 172(S,A,C,D,E,H,K,N,R,T);
 173(R,K,W); 174(I,L); 175(F,A,D,H,L,M,S,V,W,Y); 176(K,L); 177(F,G,H); 182(K,H);
 20 183(A,E,K,R); 187(E,P,V); 189(S,A,C,D); 190(S,P); 191(E,A,C,I,L,M,N,T);
 192(N,F,H,M,R,S,Y); 193(G,A,C,F,H,I,K,R,S,T,V); 195(Y,D); 198(L,A,C,G); 200(Y,E,L);
 201(A,L); 203(I,A,C,F,H,L,M,N,Q,V,Y); 206(D,A,C,I,M,Q,Y); 207(H,A,D,E,F,K,M,N,R,S,Y);
 208(P,A,D,E,H,K,L,N,Q,R,S,T); 209(D,C,G,I,K); 210(V,S); 211(V,C,D,E,F,I,L,M,N,Q,S,T);
 212(N,A,C,D,E,G,H,I,L,M,Q,R,S,T,V,Y); 215(K,E,F,L,M,N,Q,R,T,Y); 216(K,R); 219(V,E,I,L);
 25 222(A,T); 225(V,L); 226(G,C,E,K,Q); 227(L,Y); 232(L,R,V); 235(V,A,C,L,T);
 238(I,L,M,P,Q,R); 239(K,C,M,Q); 240(F,D); 241(S,A,C,D,E,F,G,H,I,K,L,N,P,Q,R,T,V,W,Y);
 242(F,E,I,V,Y); 243(L,A,I,M,S,T); 244(K,A,C,H,M,N,Q,R,S,T,Y); 245(D,E); 246(W,F);
 248(D,A,N,Q); 249(N,A,C,D,E,F,G,K,M,Q,R,S,T,W); 250(A,S); 251(R,A,D,K,L,M,N,Q,S,T);
 252(A,C,D,F,H,M,R); 253(A,C,D,E,L,M,N,Q,R,S,T,V); 254(T,F,I,L,M); 256(K,R);
 30 257(E,A,C,F,G,H,K,L,Q,R,S,V,Y); 258(M,C,F,I,L,Y); 259(F,P); 260(T,A,L,S,Y); 262(G,A);
 266(Q,A,C,D,E,F,G,H,I,K,L,M,N,R,S,T,W); 267(N,D); 269(L,A,I,V);
 270(G,A,C,D,F,H,I,K,P,Q,R,S,V,W,Y); 271(A,C,S); 273(N,D,E,H,I,K,L,M,Q,S); 275(Y,F);
 276(L,M); 277(A,C,D,E,F,H,I,K,L,M,N,Q,T,W,Y); 280(N,A,C,D,E,G,H,K,L,Q,T,Y);
 281(Y,A,D,G); 283(Q,H,T,V); 284(S,E,F,H,K,M,R,W,Y); 285(L,A); 286(F,L,M); 288(A,V);
 35 295(Y,H,L,Q); 296(A,D,E,F,H,I,K,M,Q,R,S,T); 299(T,E,I,R); 300(G,A,K,L,Q,R);
 301(G,A,F,H,K,M,Q,R,S,T,Y); 302(G,S); 303(Y,A,F,I,R,T,V,W); 307(R,Q,S);
 308(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,Y); 310(L,D,E,T); 311(N,D,E,H,K,Q,S,V); 312(N,D,G);
 313(T,A,S); 316(A,K,Q,S); 317(S,A,C,D,E,G,H,K,L,M,Q,T,W,Y);
 318(N,A,C,D,E,F,H,I,K,L,M,Q,R,S,V,W,Y); 320(T,A,C,D,E,G,H,K,M,N,P,Q,R,V,W,Y);
 40 321(K,H,R); 325(L,F,I,M,V); 327(E,D); 335(Q,G); 336(S,A,D); 339(S,Q); 342(Q,E,L);
 343(P,A,M,W); 348(L,C,G,Q,S); 349(A,G,S,W); 357(S,A,C,D,E,L,M,N,Q,V,Y);
 358(G,D,E,Q,S); 360(Y,C,E,F,H,I,L,V); 362(S,A,C,E,I,Q,T,V); 363(V,I,L); 368(M,L,W,Y);
 372(K,A,D,H,M,N,R); 374(T,A,K,N,P,Q,S); 375(T,S); 376(T,A,G,H,K,N,Q,S,Y);
 377(R,A,C,D,G,H,I,K,L,M,S,T,V,Y); 378(E,Q); 384(S,D,E,G,H,N,P); 385(K,A,E);
 45 388(P,C,D,I,K,L,R,S); 390(L,C,I); 391(K,E,F,L,T,V,Y); 392(A,G); 394(K,E,H,M);
 395(D,C,E,F,I,K,M,Q,R,S,W,Y); 396(Y,C,F,K); 397(A,S);
 400(T,A,C,D,E,H,K,L,M,N,Q,R,S,V,W,Y); 401(Q,M); 402(R,F,K,L,Q,S,T,V,W,Y); 403(D,S);
 404(Y,W); 405(I,L); 407(N,A,C,D,E,G,H,Q,S); 408(P,E,H,K,M,Q,R,S,V,W,Y); 409(D,N);
 410(V,E,I,K,L,M,R,S,Y); 414(T,A,S); 416(E,C,D,F,G,H,K,L,M,N,Q,R,S,T,V);
 50 418(D,A,C,E,F,G,H,I,K,L,M,N,Q,R,S,T,W,Y); 419(S,A,E,G,K,P,Q,R,T,V,Y);
 420(T,A,C,D,E,F,G,H,I,L,P,Q,R,S,V,W,Y); 421(K,A,C,F,G,H,I,L,N,R,S,W,Y);
 422(A,C,D,E,I,K,L,M,N,P,Q,R,S,T,V,W,Y); 423(K,A,C,D,E,F,G,M,N,P,Q,S,V,W,Y);
 424(S,A); 426(L,C); 427(A,C,G,S); 429(V,C,L,M); 430(I,C,E,L,M); 431(T,A,D);
 433(G,A,H,N,S); 434(P,D,M,N,Q,R,S); 435(G,A,C,E,I,L,N,Q,R,S,T,V,Y);
 55 437(S,A,D,E,F,H,K,L,Q,Y); 441(Y,C,K,L,N,Q,R,S,W); 442(V,A,I,L,T);
 444(T,A,E,F,H,I,K,L,M,N,P,Q,V,Y); 445(S,A,C,D,H,M,R,T,V,W); 447(A,G,K,Q,R,S,T,V);
 448(G,A,D,E,F,H,K,L,N,Q,T,W,Y); 449(E,Q); 450(I,A,C,D,F,H,K,L,N,P,Q,R,T,V);
 452(Y,A,I,L,M,S,V,W); 454(L,A,C,E,F,H,I,K,M,Q,S,T,V,Y); 455(T,A,C,I,L,M,S,V);

456(G,A,C,D,E,F,H,K,L,M,N,R,S,T,W,Y); 458(R,A,C,D,E,F,I,M,N,S,W,Y);
 459(T,A,C,D,F,G,L,S,V,W); 460(DE,H,N); 461(K,A,D,G,I,L,M,N,P,S,Y); 462(I,V);
 463(T,C,E,F,I,K,L,M,N,V,Y); 464(I,V); 465(G,A,M,N,P,Q);
 466(S,A,C,D,E,F,H,I,K,L,M,N,R,T,V,W,Y); 467(D,N); 469(Y,C,F,I,L,S,V); 470(A,G);
 471(T,A,D,E,F,G,H,I,K,L,M,N,P,Q,W); 474(V,C); 475(N,A,D,F,G,H,I,K,L,M,P,S,T,V);
 476(G,A,C,D,E,H,K,N,P,Q,R,S,T,V,Y); 477(G,A,D,E,H,I,K,P,Q,R,S,T,V,Y); 479(V,C,H,W);
 480(S,G); 481(VA,C,I,L); 482(W,Y); 483(V,I); 484(Q,A,H,K); and
 485(Q,A,D,E,F,H,I,K,L,M,N,P,R,T,V,Y)

[0100] The productive positions in CspAmy2-v1 that fall within the previously described Productivity Scores of "2," "3," and "4," and the substitutions within those positions that are combinable, are listed below. The position numbering is based on mature CspAmy2 protein listed in SEQ ID NO: 1.

LIST B:

[0101]

1(A,E,G,I,K,N,Q,R,T,V,Y); 2(A,E,G,H,K,N,P,Q,R,S,Y); 3(T,A,D,F,G,M,P,Q,R,S);
 5(G,A,C,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,Y); 15(V,C,I,L,N,S,T); 21(Q,L,T,W);
 23(N,A,D,E,F,H,K,M,Q,S,T,V,W,Y); 27(T,D,E,F,G,H,I,K,L,M,N,Q,R,S,W,Y); 28(D,A,E,N);
 30(P,A,C,D,E,F,G,H,K,L,N,R,S,T,W,Y); 31(Y,E,F,H,K,M,R,W); 33(S,D,E,G,H,K,N,Q,R);
 35(V,H,I,M,N); 38(T,D,N,S); 42(T,A,C,I,L,M,V); 46(Y,F,M,T);
 49(T,A,C,D,E,F,G,H,I,K,L,M,N,S,V,Y); 50(S,D,E,K); 52(A,F,G,H,I,K,N,Q,S,T,W);
 54(V,A,C,D,E,G,I,L,N,Q,R,S,T); 68(N,A,C,E,F,M,S,Y); 70(K,N,R,W); 72(T,G,H,S);
 81(G,A,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,W,Y); 84(K,I,Q,V); 85(S,A,C,D,E,H,K,L,M,N,Q,R);
 88(N,A,D,E,H,Q,R,T); 89(T,C,D,E,H,M,N,Q,R,S,V); 92(S,A,L,M,R,V); 93(N,D,M,T);
 106(K,A,I,V); 108(G,A,K,R,S); 111(Y,A,C,D,F,G,H,K,L,M,N,Q,R,S,T,V,W);
 112(T,A,C,D,E,F,G,I,L,M,P,Q,R,V,W,Y); 115(V,A,I,M);
 116(T,A,C,D,E,G,H,I,K,L,M,N,S,V,W); 118(V,A,C,F,I,K,L,M,N,Q,R,S); 121(N,K,R,S);
 122(P,A,K,Q,R,T); 123(S,A,C,E,G,H,K,N,Q,R,T,Y); 127(Q,A,C,E,H,I,K,M,R,T,V,Y);
 128(E,G,I,K,S,V,Y); 129(T,A,F,G,H,I,K,L,Q,R,S,V,Y); 130(S,A,G,H,I,R,T,V);
 131(G,A,F,H,I,K,M,N,P,Q,T,W); 132(E,A,C,G,H,I,L,M,N,P,Q,R,S,T,V,Y);
 133(Y,A,D,E,F,H,K,L,N,T,V); 134(N,C,D,F,G,H,M,P,Q,S,T,Y); 135(I,H,M,R,V);
 136(Q,A,F,G,H,I,K,N,T,W,Y); 138(W,A,D,F,G,H,K,L,M,P,Q,R,S,T,V,Y);
 140(G,C,D,E,F,H,I,K,L,M,N,S,T,V,Y); 142(N,A,C,D,E,F,G,H,I,K,L,M,Q,R,S,T,V,W,Y);
 144(P,A,C,D,F,G,H,I,K,L,M,N,Q,R,T,Y); 145(G,A,E,H,I,K,L,M,N,Q,R,S,T,V,W,Y);
 147(G,C,E,H,I,K,L,N,Q,R,V); 148(T,A,G,H,I,K,L,S,W,Y);
 149(T,A,C,D,E,H,I,K,L,M,N,R,S,W,Y); 150(Y,D,F,G,H,I,M,P,Q,W);
 152(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,W); 153(F,H,W,Y); 154(K,A,E,F,H,N,R,S,T);
 156(Q,D,F,G,H,I,K,L,M,S,T,V,Y); 158(F,A,C,D,E,G,H,I,L,N,P,R,S,T,V,W,Y);
 163(T,C,D,F,L,M,N,Q,S,V); 165(W,F,H,P); 166(D,A,C,G,K,M);
 167(Q,A,C,D,G,H,K,N,R,S,T,V,Y); 168(S,C,D,E,G,I,K,L,M,N,R,T,V,W);
 169(R,A,C,D,E,H,K,L,M,Q,T,W,Y); 170(S,C,G,H,N,R,T);
 171(L,F,G,H,I,K,M,N,Q,R,S,V,W,Y); 172(S,A,C,D,E,H,K,N,R,T);
 175(F,A,D,H,L,M,S,V,W,Y); 183(A,E,K,R); 189(S,A,C,D); 191(E,A,C,I,L,M,N,T);
 192(N,F,H,M,R,S,Y); 193(G,A,C,F,H,I,K,R,S,T,V); 198(L,A,C,G);
 203(I,A,C,F,H,L,M,N,Q,V,Y); 206(D,A,C,I,M,Q,Y); 207(H,A,D,E,F,K,M,N,R,S,Y);
 208(P,A,D,E,H,K,L,N,Q,R,S,T); 209(D,C,G,I,K); 211(V,C,D,E,F,I,L,M,N,Q,S,T);
 212(N,A,C,D,E,G,H,I,L,M,Q,R,S,T,V,Y); 215(K,E,F,L,M,N,Q,R,T,Y); 219(V,E,I,L);
 226(G,C,E,K,Q); 235(V,A,C,L,T); 238(I,L,M,P,Q,R); 239(K,C,M,Q);
 241(S,A,C,D,E,F,G,H,I,K,L,N,P,Q,R,T,V,W,Y); 242(F,E,I,V,Y); 243(L,A,I,M,S,T);
 244(K,A,C,H,M,N,Q,R,S,T,Y); 248(D,A,N,Q); 249(N,A,C,D,E,F,G,K,M,Q,R,S,T,W);
 251(R,A,D,K,L,M,N,Q,S,T); 252(A,C,D,F,H,M,R); 253(A,C,D,E,L,M,N,Q,R,S,T,V);
 254(T,F,I,L,M); 257(E,A,C,F,G,H,K,L,Q,R,S,V,Y); 258(M,C,F,I,L,Y); 260(T,A,L,S,Y);
 266(Q,A,C,D,E,F,G,H,I,K,L,M,N,R,S,T,W); 269(L,A,I,V);
 270(G,A,C,D,F,H,I,K,P,Q,R,S,V,W,Y); 273(N,D,E,H,I,K,L,M,Q,S);
 277(A,C,D,E,F,H,I,K,L,M,N,Q,T,W,Y); 280(N,A,C,D,E,G,H,K,L,Q,T,Y); 281(Y,A,D,G);
 283(Q,H,T,V); 284(S,E,F,H,K,M,R,W,Y); 295(Y,H,L,Q); 296(A,D,E,F,H,I,K,M,Q,R,S,T);

299(T,E,I,R); 300(G,A,K,L,Q,R); 301(G,A,F,H,K,M,Q,R,S,T,Y); 303(Y,A,F,I,R,T,V,W);
 308(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,Y); 310(L,D,E,T); 311(N,D,E,H,K,Q,S,V);
 316(A,K,Q,S); 317(S,A,C,D,E,G,H,K,L,M,Q,T,W,Y);
 318(N,A,C,D,E,F,H,I,K,L,M,Q,R,S,V,W,Y); 320(T,A,C,D,E,G,H,K,M,N,P,Q,R,V,W,Y);
 5 321(K,H,R); 325(L,F,I,M,V); 343(P,A,M,W); 348(L,C,G,Q,S); 349(A,G,S,W);
 357(S,A,C,D,E,L,M,N,Q,V,Y); 358(G,D,E,Q,S); 360(Y,C,E,F,H,I,L,V); 362(S,A,C,E,I,Q,T,V);
 368(M,L,W,Y); 372(K,A,D,H,M,N,R); 374(T,A,K,N,P,Q,S); 376(T,A,G,H,K,N,Q,S,Y);
 377(R,A,C,D,G,H,I,K,L,M,S,T,V,Y); 384(S,D,E,G,H,N,P); 385(K,A,E); 388(P,C,D,I,K,L,R,S);
 391(K,E,F,L,T,V,Y); 394(K,E,H,M); 395(D,C,E,F,I,K,M,Q,R,S,W,Y); 396(Y,C,F,K);
 10 400(T,A,C,D,E,H,K,L,M,N,Q,R,S,V,W,Y); 402(R,F,K,L,Q,S,T,V,W,Y);
 407(N,A,C,D,E,G,H,Q,S); 408(P,E,H,K,M,Q,R,S,V,W,Y); 410(V,E,I,K,L,M,R,S,Y);
 416(E,C,D,F,G,H,K,L,M,N,Q,R,S,T,V,Y); 418(D,A,C,E,F,G,H,I,K,L,M,N,Q,R,S,T,W,Y);
 419(S,A,E,G,K,P,Q,R,T,V,Y); 420(T,A,C,D,E,F,G,H,I,L,P,Q,R,S,V,W,Y);
 421(K,A,C,F,G,H,I,L,N,R,S,W,Y); 422(A,C,D,E,I,K,L,M,N,P,Q,R,S,T,V,W,Y);
 15 423(K,A,C,D,E,F,G,M,N,P,Q,S,V,W,Y); 427(A,C,G,S); 429(V,C,L,M); 430(I,C,E,L,M);
 433(G,A,H,N,S); 434(P,D,M,N,Q,R,S); 435(G,A,C,E,I,L,N,Q,R,S,T,V,Y);
 437(S,A,D,E,F,H,K,L,Q,Y); 441(Y,C,K,L,N,Q,R,S,W); 442(V,A,I,L,T);
 444(T,A,E,F,H,I,K,L,M,N,P,Q,V,Y); 445(S,A,C,D,H,M,R,T,V,W); 447(A,G,K,Q,R,S,T,V);
 448(G,A,D,E,F,H,K,L,N,Q,T,W,Y); 450(I,A,C,D,F,H,K,L,N,P,Q,R,T,V);
 20 452(Y,A,I,L,M,S,V,W); 454(L,A,C,E,F,H,I,K,M,Q,S,T,V,Y); 455(T,A,C,I,L,M,S,V);
 456(G,A,C,D,E,F,H,K,L,M,N,R,S,T,W,Y); 458(R,A,C,D,E,F,I,M,N,S,W,Y);
 459(T,A,C,D,F,G,L,S,V,W); 460(DE,H,N); 461(K,A,D,G,I,L,M,N,P,S,Y);
 463(T,C,E,F,I,K,L,M,N,V,Y); 465(G,A,M,N,P,Q); 466(S,A,C,D,E,F,H,I,K,L,M,N,R,T,V,W,Y);
 469(Y,C,F,I,L,S,V); 471(T,A,D,E,F,G,H,I,K,N,P,Q,W); 475(N,A,D,F,G,H,I,K,L,M,P,S,T,V);
 25 476(G,A,C,D,E,H,K,N,P,Q,R,S,T,V,Y); 477(G,A,D,E,H,I,K,P,Q,R,S,T,V,Y); 479(V,C,H,W);
 481(VA,C,I,L); 484(Q,A,H,K); and 485(Q,A,D,E,F,H,I,K,L,M,N,P,R,T,V,Y)

[0102] The productive positions in CspAmy2-v1 that fall within the previously described Productivity Scores of "3" and "4," and the substitutions within those positions that are combinable, are listed below. The position numbering is based on mature CspAmy2 protein listed in SEQ ID NO: 1.

LIST C:

[0103]

1(A,E,G,I,K,N,Q,R,T,V,Y); 2(A,E,G,H,K,N,P,Q,R,S,Y); 3(T,A,D,F,G,M,P,Q,R,S);
 5(G,A,C,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,Y); 15(V,C,I,L,N,S,T);
 23(N,A,D,E,F,H,K,M,Q,S,T,V,W,Y); 27(T,D,E,F,G,H,I,K,L,M,N,Q,R,S,W,Y);
 30(P,A,C,D,E,F,G,H,K,L,N,R,S,T,W,Y); 31(Y,E,F,H,K,M,R,W); 33(S,D,E,G,H,K,N,Q,R);
 40 42(T,A,C,I,L,M,V); 49(T,A,C,D,E,F,G,H,I,K,L,M,N,S,V,Y); 52(A,F,G,H,I,K,N,Q,S,T,W);
 54(V,A,C,D,E,G,I,L,N,Q,R,S,T); 68(N,A,C,E,F,M,S,Y);
 81(G,A,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,W,Y); 85(S,A,C,D,E,H,K,L,M,N,Q,R);
 88(N,A,D,E,H,Q,R,T); 89(T,C,D,E,H,M,N,Q,R,S,V); 92(S,A,L,M,R,V);
 111(Y,A,C,D,F,G,H,K,L,M,N,Q,R,S,T,V,W); 112(T,A,C,D,E,F,G,I,L,M,P,Q,R,V,W,Y);
 45 116(T,A,C,D,E,G,H,I,K,L,M,N,S,V,W); 118(V,A,C,F,I,K,L,M,N,Q,R,S); 122(P,A,K,Q,R,T);
 123(S,A,C,E,G,H,K,N,Q,R,T,Y); 127(Q,A,C,E,H,I,K,M,R,T,V,Y); 128(E,G,I,K,S,V,Y);
 129(T,A,F,G,H,I,K,L,Q,R,S,V,Y); 130(S,A,G,H,I,R,T,V); 131(G,A,F,H,I,K,M,N,P,Q,T,W);
 132(E,A,C,G,H,I,L,M,N,P,Q,R,S,T,V,Y); 133(Y,A,D,E,F,H,K,L,N,T,V);
 134(N,C,D,F,G,H,M,P,Q,S,T,Y); 136(Q,A,F,G,H,I,K,N,T,W,Y);
 50 138(W,A,D,F,G,H,K,L,M,P,Q,R,S,T,V,Y); 140(G,C,D,E,F,H,I,K,L,M,N,S,T,V,Y);
 142(N,A,C,D,E,F,G,H,I,K,L,M,Q,R,S,T,V,W,Y); 144(P,A,C,D,F,G,H,I,K,L,M,N,Q,R,T,Y);
 145(G,A,E,H,I,K,L,M,N,Q,R,S,T,V,W,Y); 147(G,C,E,H,I,K,L,N,Q,R,V);
 148(T,A,G,H,I,K,L,S,W,Y); 149(T,A,C,D,E,H,I,K,L,M,N,R,S,W,Y);
 150(Y,D,F,G,H,I,M,P,Q,W); 152(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,W);
 55 154(K,A,E,F,H,N,R,S,T); 156(Q,D,F,G,H,I,K,L,M,S,T,V,Y);
 158(F,A,C,D,E,G,H,I,L,N,P,R,S,T,V,W,Y); 163(T,C,D,F,L,M,N,Q,S,V); 166(D,A,C,G,K,M);
 167(Q,A,C,D,G,H,K,N,R,S,T,V,Y); 168(S,C,D,E,G,I,K,L,M,N,R,T,V,W);
 169(R,A,C,D,E,H,K,L,M,Q,T,W,Y); 170(S,C,G,H,N,R,T);

171(L,F,G,H,I,K,M,N,Q,R,S,V,W,Y); 172(S,A,C,D,E,H,K,N,R,T);
 175(F,A,D,H,L,M,S,V,W,Y); 191(E,A,C,I,L,M,N,T); 192(N,F,H,M,R,S,Y);
 193(G,A,C,F,H,I,K,R,S,T,V); 203(I,A,C,F,H,L,M,N,Q,V,Y); 206(D,A,C,I,M,Q,Y);
 207(H,A,D,E,F,K,M,N,R,S,Y); 208(P,A,D,E,H,K,L,N,Q,R,S,T); 209(D,C,G,I,K);
 5 211(V,C,D,E,F,I,L,M,N,Q,S,T); 212(N,A,C,D,E,G,H,I,L,M,Q,R,S,T,V,Y);
 215(K,E,F,L,M,N,Q,R,T,Y); 241(S,A,C,D,E,F,G,H,I,K,L,N,P,Q,R,T,V,W,Y);
 244(K,A,C,H,M,N,Q,R,S,T,Y); 249(N,A,C,D,E,F,G,K,M,Q,R,S,T,W);
 251(R,A,D,K,L,M,N,Q,S,T); 252(A,C,D,F,H,M,R); 253(A,C,D,E,L,M,N,Q,R,S,T,V);
 257(E,A,C,F,G,H,K,L,Q,R,S,V,Y); 258(M,C,F,I,L,Y);
 10 266(Q,A,C,D,E,F,G,H,I,K,L,M,N,R,S,T,W); 270(G,A,C,D,F,H,I,K,P,Q,R,S,V,W,Y);
 273(N,D,E,H,I,K,L,M,Q,S); 277(A,C,D,E,F,H,I,K,L,M,N,Q,T,W,Y);
 280(N,A,C,D,E,G,H,K,L,Q,T,Y); 284(S,E,F,H,K,M,R,W,Y); 296(A,D,E,F,H,I,K,M,Q,R,S,T);
 301(G,A,F,H,K,M,Q,R,S,T,Y); 303(Y,A,F,I,R,T,V,W);
 308(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,Y); 311(N,D,E,H,K,Q,S,V);
 15 317(S,A,C,D,E,G,H,K,L,M,Q,T,W,Y); 318(N,A,C,D,E,F,H,I,K,L,M,Q,R,S,V,W,Y);
 320(T,A,C,D,E,G,H,K,M,N,P,Q,R,V,W,Y); 357(S,A,C,D,E,L,M,N,Q,V,Y);
 360(Y,C,E,H,I,L,V); 362(S,A,C,E,I,Q,T,V); 372(K,A,D,H,M,N,R); 374(T,A,K,N,P,Q,S);
 376(T,A,G,H,K,N,Q,S,Y); 377(R,A,C,D,G,H,I,K,L,M,S,T,V,Y); 384(S,D,E,G,H,N,P);
 388(P,C,D,I,K,L,R,S); 391(K,E,F,L,T,V,Y); 395(D,C,E,F,I,K,M,Q,R,S,W,Y); 396(Y,C,F,K);
 20 400(T,A,C,D,E,H,K,L,M,N,Q,R,S,V,W,Y); 402(R,F,K,L,Q,S,T,V,W,Y);
 407(N,A,C,D,E,G,H,Q,S); 408(P,E,H,K,M,Q,R,S,V,W,Y); 410(V,E,I,K,L,M,R,S,Y);
 416(E,C,D,F,G,H,K,L,M,N,Q,R,S,T,V); 418(D,A,C,E,F,G,H,I,K,L,M,N,Q,R,S,T,W,Y);
 419(S,A,E,G,K,P,Q,R,T,V,Y); 420(T,A,C,D,E,F,G,H,I,L,P,Q,R,S,V,W,Y);
 421(K,A,C,F,G,H,I,L,N,R,S,W,Y); 422(A,C,D,E,I,K,L,M,N,P,Q,R,S,T,V,W,Y);
 25 423(K,A,C,D,E,F,G,M,N,P,Q,S,V,W,Y); 434(P,D,M,N,Q,R,S);
 435(G,A,C,E,I,L,N,Q,R,S,T,V,Y); 437(S,A,D,E,F,H,K,L,Q,Y); 441(Y,C,K,L,N,Q,R,S,W);
 444(T,A,E,F,H,I,K,L,M,N,P,Q,V,Y); 445(S,A,C,D,H,M,R,T,V,W); 447(A,G,K,Q,R,S,T,V);
 448(G,A,D,E,F,H,K,L,N,Q,T,W,Y); 450(I,A,C,D,F,H,K,L,N,P,Q,R,T,V);
 452(Y,A,I,L,M,S,V,W); 454(L,A,C,E,F,H,I,K,M,Q,S,T,V,Y); 455(T,A,C,I,L,M,S,V);
 30 456(G,A,C,D,E,F,H,K,L,M,N,R,S,T,W,Y); 458(R,A,C,D,E,F,I,M,N,S,W,Y);
 459(T,A,C,D,F,G,L,S,V,W); 461(K,A,D,G,I,L,M,N,P,S,Y); 463(T,C,E,F,I,K,L,M,N,V,Y);
 465(G,A,M,N,P,Q); 466(S,A,C,D,E,F,H,I,K,L,M,N,R,T,V,W,Y); 469(Y,C,F,I,L,S,V);
 471(T,A,D,E,F,G,H,I,K,N,P,Q,W); 475(N,A,D,F,G,H,I,K,L,M,P,S,T,V);
 476(G,A,C,D,E,H,K,N,P,Q,R,S,T,V,Y); 477(G,A,D,E,H,I,K,P,Q,R,S,T,V,Y); and
 35 485(Q,A,D,E,F,H,I,K,L,M,N,P,R,T,V,Y)

[0104] The productive positions in CspAmy2-v1 that fall within the previously described Productivity Scores of "4," and the substitutions within those positions that are combinable, are listed below. The position numbering is based on mature CspAmy2 protein listed in SEQ ID NO: 1.

LIST D:

[0105]

1(A,E,G,I,K,N,Q,R,T,V,Y); 2(A,E,G,H,K,N,P,Q,R,S,Y); 3(T,A,D,F,G,M,P,Q,R,S);
 5(G,A,C,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,Y); 23(N,A,D,E,F,H,K,M,Q,S,T,V,W,Y);
 27(T,D,E,F,G,H,I,K,L,M,N,Q,R,S,W,Y); 30(P,A,C,D,E,F,G,H,K,L,N,R,S,T,W,Y);
 49(T,A,C,D,E,F,G,H,I,K,L,M,N,S,V,Y); 52(A,F,G,H,I,K,N,Q,S,T,W);
 54(V,A,C,D,E,G,I,L,N,Q,R,S,T); 68(N,A,C,E,F,M,S,Y);
 81(G,A,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,W,Y); 85(S,A,C,D,E,H,K,L,M,N,Q,R);
 89(T,C,D,E,H,M,N,Q,R,S,V); 111(Y,A,C,D,F,G,H,K,L,M,N,Q,R,S,T,V,W);
 112(T,A,C,D,E,F,G,I,L,M,P,Q,R,V,W,Y); 116(T,A,C,D,E,G,H,I,K,L,M,N,S,V,W);
 118(V,A,C,F,I,K,L,M,N,Q,R,S); 123(S,A,C,E,G,H,K,N,Q,R,T,Y);
 127(Q,A,C,E,H,I,K,M,R,T,V,Y); 128(E,G,I,K,S,V,Y); 129(T,A,F,G,H,I,K,L,Q,R,S,V,Y);
 131(G,A,F,H,I,K,M,N,P,Q,T,W); 132(E,A,C,G,H,I,L,M,N,P,Q,R,S,T,V,Y);
 133(Y,A,D,E,F,H,K,L,N,T,V); 134(N,C,D,F,G,H,M,P,Q,S,T,Y); 136(Q,A,F,G,H,I,K,N,T,W,Y);
 138(W,A,D,F,G,H,K,L,M,P,Q,R,S,T,V,Y); 140(G,C,D,E,F,H,I,K,L,M,N,S,T,V,Y);
 142(N,A,C,D,E,F,G,H,I,K,L,M,Q,R,S,T,V,W,Y); 144(P,A,C,D,F,G,H,I,K,L,M,N,Q,R,T,Y);

145(G,A,E,H,I,K,L,M,N,Q,R,S,T,V,W,Y); 147(G,C,E,H,I,K,L,N,Q,R,V);
 148(T,A,G,H,I,K,L,S,W,Y); 149(T,A,C,D,E,H,I,K,L,M,N,R,S,W,Y);
 150(Y,D,F,G,H,I,M,P,Q,W); 152(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,W);
 156(Q,D,F,G,H,I,K,L,M,S,T,V,Y); 158(F,A,C,D,E,G,H,I,L,N,P,R,S,T,V,W,Y);
 5 163(T,C,D,F,L,M,N,Q,S,V); 167(Q,A,C,D,G,H,K,N,R,S,T,V,Y);
 168(S,C,D,E,G,I,K,L,M,N,R,T,V,W); 169(R,A,C,D,E,H,K,L,M,Q,T,W,Y);
 171(L,F,G,H,I,K,M,N,Q,R,S,V,W,Y); 172(S,A,C,D,E,H,K,N,R,T);
 193(G,A,C,F,H,I,K,R,S,T,V); 203(I,A,C,F,H,L,M,N,Q,V,Y); 207(H,A,D,E,F,K,M,N,R,S,Y);
 208(P,A,D,E,H,K,L,N,Q,R,S,T); 211(V,C,D,E,F,I,L,M,N,Q,S,T);
 10 212(N,A,C,D,E,G,H,I,L,M,Q,R,S,T,V,Y); 215(K,E,F,L,M,N,Q,R,T,Y);
 241(S,A,C,D,E,F,G,H,I,K,L,N,P,Q,R,T,V,W,Y); 244(K,A,C,H,M,N,Q,R,S,T,Y);
 249(N,A,C,D,E,F,G,K,M,Q,R,S,T,W); 251(R,A,D,K,L,M,N,Q,S,T);
 253(A,C,D,E,L,M,N,Q,R,S,T,V); 257(E,A,C,F,G,H,K,L,Q,R,S,V,Y);
 266(Q,A,C,D,E,F,G,H,I,K,L,M,N,R,S,T,W); 270(G,A,C,D,F,H,I,K,P,Q,R,S,V,W,Y);
 15 273(N,D,E,H,I,K,L,M,Q,S); 277(A,C,D,E,F,H,I,K,L,M,N,Q,T,W,Y);
 280(N,A,C,D,E,G,H,K,L,Q,T,Y); 284(S,E,F,H,K,M,R,W,Y); 296(A,D,E,F,H,I,K,M,Q,R,S,T);
 301(G,A,F,H,K,M,Q,R,S,T,Y); 308(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,Y);
 317(S,A,C,D,E,G,H,K,L,M,Q,T,W,Y); 318(N,A,C,D,E,F,H,I,K,L,M,Q,R,S,V,W,Y);
 320(T,A,C,D,E,G,H,K,M,N,P,Q,R,V,W,Y); 357(S,A,C,D,E,L,M,N,Q,V,Y);
 20 360(Y,C,E,F,H,I,L,V); 377(R,A,C,D,G,H,I,K,L,M,S,T,V,Y); 384(S,D,E,G,H,N,P);
 388(P,C,D,I,K,L,R,S); 395(D,C,E,F,I,K,M,Q,R,S,W,Y);
 400(T,A,C,D,E,H,K,L,M,N,Q,R,S,V,W,Y); 408(P,E,H,K,M,Q,R,S,V,W,Y);
 416(E,C,D,F,G,H,K,L,M,N,Q,R,S,T,V); 418(D,A,C,E,F,G,H,I,K,L,M,N,Q,R,S,T,W,Y);
 419(S,A,E,G,K,P,Q,R,T,V,Y); 420(T,A,C,D,E,F,G,H,I,L,P,Q,R,S,V,W,Y);
 25 421(K,A,C,F,G,H,I,L,N,R,S,W,Y); 422(A,C,D,E,I,K,L,M,N,P,Q,R,S,T,V,W,Y);
 423(K,A,C,D,E,F,G,M,N,P,Q,S,V,W,Y); 434(P,D,M,N,Q,R,S);
 435(G,A,C,E,I,L,N,Q,R,S,T,V,Y); 437(S,A,D,E,F,H,K,L,Q,Y);
 444(T,A,E,F,H,I,K,L,M,N,P,Q,V,Y); 445(S,A,C,D,H,M,R,T,V,W);
 448(G,A,D,E,F,H,K,L,N,Q,T,W,Y); 450(I,A,C,D,F,H,K,L,N,P,Q,R,T,V);
 30 454(L,A,C,E,F,H,I,K,M,Q,S,T,V,Y); 456(G,A,C,D,E,F,H,K,L,M,N,R,S,T,W,Y);
 458(R,A,C,D,E,F,I,M,N,S,W,Y); 459(T,A,C,D,F,G,L,S,V,W); 461(K,A,D,G,I,L,M,N,P,S,Y);
 463(T,C,E,F,I,K,L,M,N,V,Y); 466(S,A,C,D,E,F,H,I,K,L,M,N,R,T,V,W,Y);
 471(T,A,D,E,F,G,H,I,K,N,P,Q,W); 475(N,A,D,F,G,H,I,K,L,M,P,S,T,V);
 476(G,A,C,D,E,H,K,N,P,Q,R,S,T,V,Y); 477(G,A,D,E,H,I,K,P,Q,R,S,T,V,Y); and
 35 485(Q,A,D,E,F,H,I,K,L,M,N,P,R,T,V,Y)

[0106] The productive positions in CspAmy2-v1 suitable for charge or hydrophobicity modifications are listed below. The position numbering is based on mature CspAmy2 protein listed in SEQ ID NO: 1.

40 LIST E:

[0107] 1, 5, 15, 23, 30, 31, 49, 68, 111, 112, 116, 123, 127, 128, 129, 131, 132, 134, 140, 142, 144, 147, 150, 152, 153, 168, 170, 171, 183, 187, 192, 203, 207, 209, 211, 212, 232, 241, 243, 244, 253, 266, 277, 280, 300, 301, 308, 320, 357, 362, 377, 388, 400, 402, 408, 416, 420, 423, 448, 450, 454, 455, 456, 458, 466, 475, and 485

45 **[0108]** The productive positions and the substitutions within those positions in CspAmy2-v1 suitable for charge or hydrophobicity modifications are listed below. The position numbering is based on mature CspAmy2 protein listed in SEQ ID NO: 1.

50 LIST F:

[0109]

1(A,K,V,Y); 5(G,F,V); 15(V,S); 23(N,K); 30(P,E,K,L,W); 31(Y,F,K,W); 49(T,I); 68(N,Y);
 111(Y,D,Q,S,T,W); 112(T,I,W); 116(T,L); 123(S,K); 127(Q,I); 128(E,I,V); 129(T,I);
 55 131(G,H,K); 132(E,G,H,I,M,P,R,T,V,Y); 134(N,D,F,M,Y); 140(G,E,F,H,K); 142(N,D,I,R);
 144(P,G,K,L); 147(G,C,E,H,L,R,V); 150(Y,W); 152(N,D,E,L,R); 153(F,H,Y); 168(S,L);
 170(S,R); 171(L,H,N,Q,R); 183(A,K); 187(E,P); 192(N,F,Y); 203(I,C); 207(H,E,F); 209(D,G);
 211(V,D,E,N,Q); 212(N,D,E); 232(L,R); 241(S,D,E,I,L,W,Y); 243(L,S); 244(K,C); 253(A,R);

266(Q,I,R,W); 277(A,F,L,Y); 280(N,L); 300(G,R); 301(G,H); 308(N,D,E,F,L,R,V,Y);
 320(T,E,W); 357(S,E); 362(S,I,V); 377(R,G,H); 388(P,K); 400(T,E,K,L,W,Y); 402(R,F);
 408(P,E,R,W); 416(E,C,F,G,H,K,L,N,R,S,V); 420(T,D,E,I,L,R,W);
 423(K,E,F,G,M,N,Q,S,V,Y); 448(G,F,H,K,W); 450(I,K,R,T); 454(L,C); 455(T,L,M);
 456(G,F,W); 458(R,A,C,D,E,F,I,S); 466(S,I,L); 475(N,D); and 485(Q,E,K,L,Y,Y)

[0110] Although the foregoing mutations were identified using an SEL library based on CspAmy2-v1 (SEQ ID NO: 2), it is known that many bacterial (and other) α -amylases share the same fold, and often share significant amino acid sequence identity, and often benefit from the same mutations. In the present case, corresponding amino acid positions in other α -amylases can readily be identified by amino acid sequence alignment with CspAmy2 (SEQ ID NO: 1) or CspAmy2-v1 (SEQ ID NO: 2), using Clustal W with default parameters. α -amylases in which the foregoing mutations are likely to produce a performance benefit include those having a similar fold and/or having 60% or greater amino acid sequence identity to any of the well-known *Bacillus* amylases (e.g., from *B. licheniformis*, *B. stearothermophilus*, and *B. amyloliquifaciens*), Carbohydrate-Active Enzymes database (CAZy) Family 13 amylases, or any amylase that has heretofore been referred to by the descriptive term, "Termamyl-like." The reader will appreciate that where an α -amylase naturally has a mutation listed above (i.e., where the wild-type α -amylase already comprised a residue identified as a mutation), then that particular mutation does not apply to that α -amylase. However, other described mutations may work in combination with the naturally occurring residue at that position.

[0111] In some embodiments, the present α -amylase variants have at least one combinable mutation at a productive position corresponding to the combinable mutations at productive positions described, above, in Lists A, B, C, D, E, and/or F, and/or a combinable mutation as described in Table D (which use SEQ ID NO: 1 for numbering) and a defined degree of amino acid sequence homology/identity to SEQ ID NO: 1 or SEQ ID NO: 2, for example, at least 60%, at least 65%, at least 70%, at least 75%, at least 76%, at least 77%, at least 78%, at least 79%, at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or even at least 99% amino acid sequence homology/identity, and have at least combinable mutation as set forth above. In some embodiments, the suitability score of the at least one mutation is +++, +++, or +++++. In some embodiments, the suitability score of the at least one mutation is +++, or +++++. In some embodiments, the suitability score of the at least one mutation is +++++. In some embodiments, the variants have a plurality (e.g., 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, or more) combinable mutations.

[0112] In some embodiments, the present α -amylase variants have at least one combinable mutation at a productive position corresponding to the combinable mutations at productive positions described, above, in Lists A, B, C, D, E, and/or F, and/or a combinable mutation as described in Table D (which use SEQ ID NO: 1 for numbering) and are derived from a parental amylase having a defined degree of amino acid sequence homology/identity to SEQ ID NO: 1 or SEQ ID NO: 2, for example, at least 60%, at least 65%, at least 70%, at least 75%, at least 76%, at least 77%, at least 78%, at least 79%, at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or even at least 99% amino acid sequence homology/identity. In some embodiments, the suitability score of the at least one mutation is +++, +++, or +++++. In some embodiments, the suitability score of the at least one mutation is +++, or +++++. In some embodiments, the suitability score of the at least one mutation is +++++. In some embodiments, the variants have, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, or more combinable mutations.

[0113] In some embodiments, the present α -amylase variants have a mutation in the calcium binding loop based on the work of Suzuki et al. (1989) J. Biol. Chem. 264:18933-938. Exemplary mutations include a deletion or substitution at one or more residues corresponding to Arg-178, Gly-179, Thr-180, or Gly-181 in SEQ ID NO: 1. In particular embodiments, the mutation corresponds to the deletion of Arg-178 and Gly-179 or Thr-180 and Gly-181 (using SEQ ID NO: 1 numbering). Homologous residues in other amylases can be determined by structural alignment, or by primary structure alignment, as illustrated by Figure 6. In some embodiments, the mutation(s) in the calcium binding loop are combined with any of the aforementioned combinable mutations and mutations at productive positions.

2.2 Additional mutations

[0114] In some embodiments, in addition to one or more of the mutations described above (e.g., in Sections 2.1, 2.2, and 2.3), the present amylases further include one or more mutations that provide a further performance or stability benefit. Exemplary performance benefits include but are not limited to increased hydrolysis of a starch substrate, increased grain, cereal or other starch substrate liquifaction performance, increased cleaning performance, increased thermal stability, increased storage stability, increased solubility, an altered pH profile, decreased calcium dependence, increased

specific activity, modified substrate specificity, modified substrate binding, modified pH-dependent activity, modified pH-dependent stability, increased oxidative stability, and increased expression. In some cases, the performance benefit is realized at a relatively low temperature. In some cases, the performance benefit is realized at relatively high temperature.

[0115] Furthermore, the present amylases may include any number of conservative amino acid substitutions. Exemplary conservative amino acid substitutions are listed in the Table E

Table E. Conservative amino acid substitutions

<i>For Amino Acid</i>	<i>Code</i>	<i>Replace with any of</i>
Alanine	A	D-Ala, Gly, beta-Ala, L-Cys, D-Cys
Arginine	R	D-Arg, Lys, D-Lys, homo-Arg, D-homo-Arg, Met, Ile, D-Met, D-Ile, Orn, D-Orn
Asparagine	N	D-Asn, Asp, D-Asp, Glu, D-Glu, Gln, D-Gln
Aspartic Acid	D	D-Asp, D-Asn, Asn, Glu, D-Glu, Gln, D-Gln
Cysteine	C	D-Cys, S-Me-Cys, Met, D-Met, Thr, D-Thr
Glutamine	Q	D-Gln, Asn, D-Asn, Glu, D-Glu, Asp, D-Asp
Glutamic Acid	E	D-Glu, D-Asp, Asp, Asn, D-Asn, Gln, D-Gln
Glycine	G	Ala, D-Ala, Pro, D-Pro, b-Ala, Acp
Isoleucine	I	D-Ile, Val, D-Val, Leu, D-Leu, Met, D-Met
Leucine	L	D-Leu, Val, D-Val, Leu, D-Leu, Met, D-Met
Lysine	K	D-Lys, Arg, D-Arg, homo-Arg, D-homo-Arg, Met, D-Met, Ile, D-Ile, Orn, D-Orn
Methionine	M	D-Met, S-Me-Cys, Ile, D-Ile, Leu, D-Leu, Val, D-Val
Phenylalanine	F	D-Phe, Tyr, D-Thr, L-Dopa, His, D-His, Trp, D-Trp, Trans-3,4, or 5-phenylproline, cis-3,4, or 5-phenylproline
Proline	P	D-Pro, L-I-thioazolidine-4- carboxylic acid, D-or L-1-oxazolidine-4-carboxylic acid
Serine	C	D-Ser, Thr, D-Thr, allo-Thr, Met, D-Met, Met(O), D-Met(O), L-Cys, D-Cys
Threonine	T	D-Thr, Ser, D-Ser, allo-Thr, Met, D-Met, Met(O), D-Met(O), Val, D-Val
Tyrosine	Y	D-Tyr, Phe, D-Phe, L-Dopa, His, D-His
Valine	V	D-Val, Leu, D-Leu, Ile, D-Ile, Met, D-Met

[0116] The reader will appreciate that some of the above mentioned conservative mutations can be produced by genetic manipulation, while others are produced by introducing synthetic amino acids into a polypeptide by genetic or other means.

[0117] The present amylase may be "precursor," "immature," or "full-length," in which case they include a signal sequence, or "mature," in which case they lack a signal sequence. Mature forms of the polypeptides are generally the most useful. Unless otherwise noted, the amino acid residue numbering used herein refers to the mature forms of the respective amylase polypeptides. The present amylase polypeptides may also be truncated to remove the N or C-termini, so long as the resulting polypeptides retain amylase activity.

[0118] The present amylase may be a "chimeric" or "hybrid" polypeptide, in that it includes at least a portion of a first amylase polypeptide, and at least a portion of a second amylase polypeptide (such chimeric amylases have recently been "rediscovered" as domain-swap amylases). The present amylases may further include heterologous signal sequence, an epitope to allow tracking or purification, or the like. Exemplary heterologous signal sequences are from *B. licheniformis* amylase (LAT), *B. subtilis* (AmyE or AprE), and *Streptomyces* CeiA.

2.3. Nucleotides encoding variant amylase polypeptides

[0119] In another aspect, nucleic acids encoding a variant amylase polypeptide are provided as defined in the appended claims. The nucleic acid may encode a particular amylase polypeptide, or an amylase having a specified degree of amino acid sequence identity to the particular amylase.

[0120] In one example, the nucleic acid encodes an amylase having at least 60%, at least 65%, at least 70%, at least 75%, at least 76%, at least 77%, at least 78%, at least 79%, at least 80%, at least 81%, at least 82%, at least 83%, at

least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or even at least 99% homology/identity to SEQ ID NO: 1 or SEQ ID NO: 2 (excluding the portion of the nucleic acid that encodes the signal sequence). It will be appreciated that due to the degeneracy of the genetic code, a plurality of nucleic acids may encode the same polypeptide.

[0121] In another example, the nucleic acid hybridizes under stringent or very stringent conditions to a nucleic acid encoding (or complementary to a nucleic acid encoding) an amylase having at least 60%, at least 65%, at least 70%, at least 75%, at least 76%, at least 77%, at least 78%, at least 79%, at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or even at least 99% homology/identity to SEQ ID NO: 1 or SEQ ID NO: 2 (excluding the portion of the nucleic acid that encodes the signal sequence). Such hybridization conditions are described herein but are also well known in the art.

[0122] Nucleic acids may encode a "full-length" ("fl" or "FL") amylase, which includes a signal sequence, only the mature form of an amylase, which lacks the signal sequence, or a truncated form of an amylase, which lacks the N or C-terminus of the mature form.

[0123] A nucleic acid that encodes a α -amylase can be operably linked to various promoters and regulators in a vector suitable for expressing the α -amylase in host cells. Exemplary promoters are from *B. licheniformis* amylase (LAT), *B. subtilis* (AmyE or AprE), and *Streptomyces* CeiA. Such a nucleic acid can also be linked to other coding sequences, e.g., to encode a chimeric polypeptide.

3. Production of Variant Amylases

[0124] The present variant amylases can be produced in host cells, for example, by secretion or intracellular expression. A cultured cell material (e.g., a whole-cell broth) comprising a variant amylase can be obtained following secretion of the variant amylase into the cell medium. Optionally, the variant amylase can be isolated from the host cells, or even isolated from the cell broth, depending on the desired purity of the final variant amylase. A gene encoding a variant amylase can be cloned and expressed according to methods well known in the art. Suitable host cells include bacterial, fungal (including yeast and filamentous fungi), and plant cells (including algae). Particularly useful host cells include *Aspergillus niger*, *Aspergillus oryzae* or *Trichoderma reesei*. Other host cells include bacterial cells, e.g., *Bacillus subtilis* or *B. licheniformis*, as well as *Streptomyces*.

[0125] The host cell further may express a nucleic acid encoding a homologous or heterologous glucoamylase, i.e., a glucoamylase that is not the same species as the host cell, or one or more other enzymes. The glucoamylase may be a variant glucoamylase, such as one of the glucoamylase variants disclosed in U.S. Patent No. 8,058,033 (Danisco US Inc.), for example. Additionally, the host may express one or more accessory enzymes, proteins, peptides. These may benefit liquefaction, saccharification, fermentation, SSF, etc processes. Furthermore, the host cell may produce biochemicals in addition to enzymes used to digest the various feedstock(s). Such host cells may be useful for fermentation or simultaneous saccharification and fermentation processes to reduce or eliminate the need to add enzymes.

3.1. Vectors

[0126] A DNA construct comprising a nucleic acid encoding variant amylases can be constructed to be expressed in a host cell. Representative nucleic acids that encode variant amylases include SEQ ID NO: 4. Because of the well-known degeneracy in the genetic code, variant polynucleotides that encode an identical amino acid sequence can be designed and made with routine skill. It is also well known in the art to optimize codon use for a particular host cell. Nucleic acids encoding variant amylases can be incorporated into a vector. Vectors can be transferred to a host cell using well-known transformation techniques, such as those disclosed below.

[0127] The vector may be any vector that can be transformed into and replicated within a host cell. For example, a vector comprising a nucleic acid encoding a variant amylase can be transformed and replicated in a bacterial host cell as a means of propagating and amplifying the vector. The vector also may be transformed into an expression host, so that the encoding nucleic acids can be expressed as a functional amylase. Host cells that serve as expression hosts can include filamentous fungi, for example. The Fungal Genetics Stock Center (FGSC) Catalogue of Strains lists suitable vectors for expression in fungal host cells. See FGSC, Catalogue of Strains, University of Missouri, at www.fgsc.net (last modified January 17, 2007). A representative vector is pJG153, a promoterless Cre expression vector that can be replicated in a bacterial host. See Harrison et al. (June 2011) Applied Environ. Microbiol. 77: 3916-22. pJG153 can be modified with routine skill to comprise and express a nucleic acid encoding an amylase variant.

[0128] A nucleic acid encoding a variant amylase can be operably linked to a suitable promoter, which allows transcription in the host cell. The promoter may be any DNA sequence that shows transcriptional activity in the host cell of choice and may be derived from genes encoding proteins either homologous or heterologous to the host cell. Exemplary

promoters for directing the transcription of the DNA sequence encoding a variant amylase, especially in a bacterial host, are the promoter of the lac operon of *E. coli*, the *Streptomyces coelicolor* agarase gene *dagA* or *celA* promoters, the promoters of the *Bacillus licheniformis* α -amylase gene (*amyL*), the promoters of the *Bacillus stearothermophilus* maltogenic amylase gene (*amyM*), the promoters of the *Bacillus amyloliquefaciens* α -amylase (*amyQ*), the promoters of the *Bacillus subtilis* *xylA* and *xylB* genes etc. For transcription in a fungal host, examples of useful promoters are those derived from the gene encoding *Aspergillus oryzae* TAKA amylase, *Rhizomucor miehei* aspartic proteinase, *Aspergillus niger* neutral α -amylase, *A. niger* acid stable α -amylase, *A. niger* glucoamylase, *Rhizomucor miehei* lipase, *A. oryzae* alkaline protease, *A. oryzae* triose phosphate isomerase, or *A. nidulans* acetamidase. When a gene encoding an amylase is expressed in a bacterial species such as *E. coli*, a suitable promoter can be selected, for example, from a bacteriophage promoter including a T7 promoter and a phage lambda promoter. Examples of suitable promoters for the expression in a yeast species include but are not limited to the Gal 1 and Gal 10 promoters of *Saccharomyces cerevisiae* and the *Pichia pastoris* AOX1 or AOX2 promoters. *cbh1* is an endogenous, inducible promoter from *T. reesei*. See Liu et al. (2008) "Improved heterologous gene expression in *Trichoderma reesei* by cellobiohydrolase I gene (*cbh1*) promoter optimization," *Acta Biochim. Biophys. Sin* (Shanghai) 40(2): 158-65.

[0129] The coding sequence can be operably linked to a signal sequence. The DNA encoding the signal sequence may be the DNA sequence naturally associated with the amylase gene to be expressed or from a different Genus or species. A signal sequence and a promoter sequence comprising a DNA construct or vector can be introduced into a fungal host cell and can be derived from the same source. For example, the signal sequence is the *cbh1* signal sequence that is operably linked to a *cbh1* promoter.

[0130] An expression vector may also comprise a suitable transcription terminator and, in eukaryotes, polyadenylation sequences operably linked to the DNA sequence encoding a variant amylase. Termination and polyadenylation sequences may suitably be derived from the same sources as the promoter.

[0131] The vector may further comprise a DNA sequence enabling the vector to replicate in the host cell. Examples of such sequences are the origins of replication of plasmids pUC19, pACYC177, pUB110, pE194, pAMB1, and pIJ702.

[0132] The vector may also comprise a selectable marker, e.g., a gene the product of which complements a defect in the isolated host cell, such as the *dal* genes from *B. subtilis* or *B. licheniformis*, or a gene that confers antibiotic resistance such as, e.g., ampicillin, kanamycin, chloramphenicol or tetracycline resistance. Furthermore, the vector may comprise *Aspergillus* selection markers such as *amdS*, *argB*, *niaD* and *xxsC*, a marker giving rise to hygromycin resistance, or the selection may be accomplished by co-transformation, such as known in the art. See e.g., International PCT Application WO 91/17243.

[0133] Intracellular expression may be advantageous in some respects, e.g., when using certain bacteria or fungi as host cells to produce large amounts of amylase for subsequent enrichment or purification. Extracellular secretion of amylase into the culture medium can also be used to make a cultured cell material comprising the isolated amylase.

[0134] The expression vector typically includes the components of a cloning vector, such as, for example, an element that permits autonomous replication of the vector in the selected host organism and one or more phenotypically detectable markers for selection purposes. The expression vector normally comprises control nucleotide sequences such as a promoter, operator, ribosome binding site, translation initiation signal and optionally, a repressor gene or one or more activator genes. Additionally, the expression vector may comprise a sequence coding for an amino acid sequence capable of targeting the amylase to a host cell organelle such as a peroxisome, or to a particular host cell compartment. Such a targeting sequence includes but is not limited to the sequence, SKL. For expression under the direction of control sequences, the nucleic acid sequence of the amylase is operably linked to the control sequences in proper manner with respect to expression.

[0135] The procedures used to ligate the DNA construct encoding an amylase, the promoter, terminator and other elements, respectively, and to insert them into suitable vectors containing the information necessary for replication, are well known to persons skilled in the art (see, e.g., Sambrook et al., MOLECULAR CLONING: A LABORATORY MANUAL, 2nd ed., Cold Spring Harbor, 1989, and 3rd ed., 2001).

3.2. Transformation and Culture of Host Cells

[0136] An isolated cell, either comprising a DNA construct or an expression vector, is advantageously used as a host cell in the recombinant production of an amylase. The cell may be transformed with the DNA construct encoding the enzyme, conveniently by integrating the DNA construct (in one or more copies) in the host chromosome. This integration is generally considered to be an advantage, as the DNA sequence is more likely to be stably maintained in the cell. Integration of the DNA constructs into the host chromosome may be performed according to conventional methods, e.g., by homologous or heterologous recombination. Alternatively, the cell may be transformed with an expression vector as described above in connection with the different types of host cells.

[0137] Examples of suitable bacterial host organisms are Gram positive bacterial species such as *Bacillaceae* including *Bacillus subtilis*, *Bacillus licheniformis*, *Bacillus lentus*, *Bacillus brevis*, *Geobacillus* (formerly *Bacillus*) *stearothermophil-*

us, *Bacillus alkalophilus*, *Bacillus amyloliquefaciens*, *Bacillus coagulans*, *Bacillus lautus*, *Bacillus megaterium*, and *Bacillus thuringiensis*; *Streptomyces* species such as *Streptomyces murinus*; lactic acid bacterial species including *Lactococcus* sp. such as *Lactococcus lactis*; *Lactobacillus* sp. including *Lactobacillus reuteri*; *Leuconostoc* sp.; *Pediococcus* sp.; and *Streptococcus* sp. Alternatively, strains of a Gram negative bacterial species belonging to *Enterobacteriaceae* including *E. coli*, or to *Pseudomonadaceae* can be selected as the host organism.

[0138] A suitable yeast host organism can be selected from the biotechnologically relevant yeasts species such as but not limited to yeast species such as *Pichia* sp., *Hansenula* sp., or *Kluyveromyces*, *Yarrowinia*, *Schizosaccharomyces* species or a species of *Saccharomyces*, including *Saccharomyces cerevisiae* or a species belonging to *Schizosaccharomyces* such as, for example, *S. pombe* species. A strain of the methylotrophic yeast species, *Pichia pastoris*, can be used as the host organism. Alternatively, the host organism can be a *Hansenula* species. Suitable host organisms among filamentous fungi include species of *Aspergillus*, e.g., *Aspergillus niger*, *Aspergillus oryzae*, *Aspergillus tubigensis*, *Aspergillus awamori*, or *Aspergillus nidulans*. Alternatively, strains of a *Fusarium* species, e.g., *Fusarium oxysporum* or of a *Rhizomucor* species such as *Rhizomucor miehei* can be used as the host organism. Other suitable strains include *Thermomyces* and *Mucor* species. In addition, *Trichoderma* sp. can be used as a host. A suitable procedure for transformation of *Aspergillus* host cells includes, for example, that described in EP 238023. An amylase expressed by a fungal host cell can be glycosylated, i.e., will comprise a glycosyl moiety. The glycosylation pattern can be the same or different as present in the wild-type amylase. The type and/or degree of glycosylation may impart changes in enzymatic and/or biochemical properties.

[0139] It is advantageous to delete genes from expression hosts, where the gene deficiency can be cured by the transformed expression vector. Known methods may be used to obtain a fungal host cell having one or more inactivated genes. Gene inactivation may be accomplished by complete or partial deletion, by insertional inactivation or by any other means that renders a gene nonfunctional for its intended purpose, such that the gene is prevented from expression of a functional protein. Any gene from a *Trichoderma* sp. or other filamentous fungal host that has been cloned can be deleted, for example, *cbh1*, *cbh2*, *egl1*, and *egl2* genes. Gene deletion may be accomplished by inserting a form of the desired gene to be inactivated into a plasmid by methods known in the art.

[0140] Introduction of a DNA construct or vector into a host cell includes techniques such as transformation; electroporation; nuclear microinjection; transduction; transfection, e.g., lipofection mediated and DEAE-Dextrin mediated transfection; incubation with calcium phosphate DNA precipitate; high velocity bombardment with DNA-coated microprojectiles; and protoplast fusion. General transformation techniques are known in the art. See, e.g., Sambrook *et al.* (2001), *supra*. The expression of heterologous protein in *Trichoderma* is described, for example, in U.S. Patent No. 6,022,725. Reference is also made to Cao *et al.* (2000) Science 9:991-1001 for transformation of *Aspergillus* strains. Genetically stable transformants can be constructed with vector systems whereby the nucleic acid encoding an amylase is stably integrated into a host cell chromosome. Transformants are then selected and purified by known techniques.

[0141] The preparation of *Trichoderma* sp. for transformation, for example, may involve the preparation of protoplasts from fungal mycelia. See Campbell *et al.* (1989) Curr. Genet. 16: 53-56. The mycelia can be obtained from germinated vegetative spores. The mycelia are treated with an enzyme that digests the cell wall, resulting in protoplasts. The protoplasts are protected by the presence of an osmotic stabilizer in the suspending medium. These stabilizers include sorbitol, mannitol, potassium chloride, magnesium sulfate, and the like. Usually the concentration of these stabilizers varies between 0.8 M and 1.2 M, e.g., a 1.2 M solution of sorbitol can be used in the suspension medium.

[0142] Uptake of DNA into the host *Trichoderma* sp. strain depends upon the calcium ion concentration. Generally, between about 10-50 mM CaCl_2 is used in an uptake solution. Additional suitable compounds include a buffering system, such as TE buffer (10 mM Tris, pH 7.4; 1 mM EDTA) or 10 mM MOPS, pH 6.0 and polyethylene glycol. The polyethylene glycol is believed to fuse the cell membranes, thus permitting the contents of the medium to be delivered into the cytoplasm of the *Trichoderma* sp. strain. This fusion frequently leaves multiple copies of the plasmid DNA integrated into the host chromosome.

[0143] Usually transformation of *Trichoderma* sp. uses protoplasts or cells that have been subjected to a permeability treatment, typically at a density of 10^5 to 10^7 /mL, particularly 2×10^6 /mL. A volume of 100 μL of these protoplasts or cells in an appropriate solution (e.g., 1.2 M sorbitol and 50 mM CaCl_2) may be mixed with the desired DNA. Generally, a high concentration of PEG is added to the uptake solution. From 0.1 to 1 volume of 25% PEG 4000 can be added to the protoplast suspension; however, it is useful to add about 0.25 volumes to the protoplast suspension. Additives, such as dimethyl sulfoxide, heparin, spermidine, potassium chloride and the like, may also be added to the uptake solution to facilitate transformation. Similar procedures are available for other fungal host cells. See, e.g., U.S. Patent No. 6,022,725.

3.3. Expression

[0144] A method of producing an amylase may comprise cultivating a host cell as described above under conditions conducive to the production of the enzyme and recovering the enzyme from the cells and/or culture medium.

[0145] The medium used to cultivate the cells may be any conventional medium suitable for growing the host cell in

question and obtaining expression of an amylase. Suitable media and media components are available from commercial suppliers or may be prepared according to published recipes (e.g., as described in catalogues of the American Type Culture Collection).

[0146] An enzyme secreted from the host cells can be used in a whole broth preparation. In the present methods, the preparation of a spent whole fermentation broth of a recombinant microorganism can be achieved using any cultivation method known in the art resulting in the expression of an α -amylase. Fermentation may, therefore, be understood as comprising shake flask cultivation, small- or large-scale fermentation (including continuous, batch, fed-batch, or solid state fermentations) in laboratory or industrial fermenters performed in a suitable medium and under conditions allowing the amylase to be expressed or isolated. The term "spent whole fermentation broth" is defined herein as unfractionated contents of fermentation material that includes culture medium, extracellular proteins (e.g., enzymes), and cellular biomass. It is understood that the term "spent whole fermentation broth" also encompasses cellular biomass that has been lysed or permeabilized using methods well known in the art.

[0147] An enzyme secreted from the host cells may conveniently be recovered from the culture medium by well-known procedures, including separating the cells from the medium by centrifugation or filtration, and precipitating proteinaceous components of the medium by means of a salt such as ammonium sulfate, followed by the use of chromatographic procedures such as ion exchange chromatography, affinity chromatography, or the like.

[0148] The polynucleotide encoding an amylase in a vector can be operably linked to a control sequence that is capable of providing for the expression of the coding sequence by the host cell, i.e. the vector is an expression vector. The control sequences may be modified, for example by the addition of further transcriptional regulatory elements to make the level of transcription directed by the control sequences more responsive to transcriptional modulators. The control sequences may in particular comprise promoters.

[0149] Host cells may be cultured under suitable conditions that allow expression of an amylase. Expression of the enzymes may be constitutive such that they are continually produced, or inducible, requiring a stimulus to initiate expression. In the case of inducible expression, protein production can be initiated when required by, for example, addition of an inducer substance to the culture medium, for example dexamethasone or IPTG or Sophorose. Polypeptides can also be produced recombinantly in an *in vitro* cell-free system, such as the TNT™ (Promega) rabbit reticulocyte system.

[0150] An expression host also can be cultured in the appropriate medium for the host, under aerobic conditions. Shaking or a combination of agitation and aeration can be provided, with production occurring at the appropriate temperature for that host, e.g., from about 25°C to about 75°C (e.g., 30°C to 45°C), depending on the needs of the host and production of the desired variant amylase. Culturing can occur from about 12 to about 100 hours or greater (and any hour value there between, e.g., from 24 to 72 hours). Typically, the culture broth is at a pH of about 4.0 to about 8.0, again depending on the culture conditions needed for the host relative to production of an amylase.

3.4. Identification of Amylase Activity

[0151] To evaluate the expression of an amylase in a host cell, assays can measure the expressed protein, corresponding mRNA, or α -amylase activity. For example, suitable assays include Northern blotting, reverse transcriptase polymerase chain reaction, and *in situ* hybridization, using an appropriately labeled hybridizing probe. Suitable assays also include measuring amylase activity in a sample, for example, by assays directly measuring reducing sugars such as glucose in the culture media. For example, glucose concentration may be determined using glucose reagent kit No. 15-UV (Sigma Chemical Co.) or an instrument, such as Technicon Autoanalyzer. α -Amylase activity also may be measured by any known method, such as the PAHBAH or ABTS assays, described below.

3.5. Methods for Enriching and Purifying Variants Amylases

[0152] Fermentation, separation, and concentration techniques are well known in the art and conventional methods can be used in order to prepare a concentrated a variant α -amylase polypeptide-containing solution.

[0153] After fermentation, a fermentation broth is obtained, the microbial cells and various suspended solids, including residual raw fermentation materials, are removed by conventional separation techniques in order to obtain an amylase solution. Filtration, centrifugation, microfiltration, rotary vacuum drum filtration, ultrafiltration, centrifugation followed by ultrafiltration, extraction, or chromatography, or the like, are generally used.

[0154] It is desirable to concentrate a variant α -amylase polypeptide-containing solution in order to optimize recovery. Use of unconcentrated solutions requires increased incubation time in order to collect the enriched or purified enzyme precipitate.

[0155] The enzyme containing solution is concentrated using conventional concentration techniques until the desired enzyme level is obtained. Concentration of the enzyme containing solution may be achieved by any of the techniques discussed herein. Exemplary methods of enrichment and purification include but are not limited to rotary vacuum filtration and/or ultrafiltration.

[0156] The enzyme solution is concentrated into a concentrated enzyme solution until the enzyme activity of the concentrated variant α -amylase polypeptide-containing solution is at a desired level.

[0157] Concentration may be performed using, e.g., a precipitation agent, such as a metal halide precipitation agent. Metal halide precipitation agents include but are not limited to alkali metal chlorides, alkali metal bromides and blends of two or more of these metal halides. Exemplary metal halides include sodium chloride, potassium chloride, sodium bromide, potassium bromide and blends of two or more of these metal halides. The metal halide precipitation agent, sodium chloride, can also be used as a preservative.

[0158] The metal halide precipitation agent is used in an amount effective to precipitate an amylase. The selection of at least an effective amount and an optimum amount of metal halide effective to cause precipitation of the enzyme, as well as the conditions of the precipitation for maximum recovery including incubation time, pH, temperature and concentration of enzyme, will be readily apparent to one of ordinary skill in the art, after routine testing.

[0159] Generally, at least about 5% w/v (weight/volume) to about 25% w/v of metal halide is added to the concentrated enzyme solution, and usually at least 8% w/v. Generally, no more than about 25% w/v of metal halide is added to the concentrated enzyme solution and usually no more than about 20% w/v. The optimal concentration of the metal halide precipitation agent will depend, among others, on the nature of the specific variant α -amylase polypeptide and on its concentration in the concentrated enzyme solution.

[0160] Another alternative way to precipitate the enzyme is to use organic compounds. Exemplary organic compound precipitating agents include: 4-hydroxybenzoic acid, alkali metal salts of 4-hydroxybenzoic acid, alkyl esters of 4-hydroxybenzoic acid, and blends of two or more of these organic compounds. The addition of the organic compound precipitation agents can take place prior to, simultaneously with or subsequent to the addition of the metal halide precipitation agent, and the addition of both precipitation agents, organic compound and metal halide, may be carried out sequentially or simultaneously.

[0161] Generally, the organic precipitation agents are selected from the group consisting of alkali metal salts of 4-hydroxybenzoic acid, such as sodium or potassium salts, and linear or branched alkyl esters of 4-hydroxybenzoic acid, wherein the alkyl group contains from 1 to 12 carbon atoms, and blends of two or more of these organic compounds. The organic compound precipitation agents can be, for example, linear or branched alkyl esters of 4-hydroxybenzoic acid, wherein the alkyl group contains from 1 to 10 carbon atoms, and blends of two or more of these organic compounds. Exemplary organic compounds are linear alkyl esters of 4-hydroxybenzoic acid, wherein the alkyl group contains from 1 to 6 carbon atoms, and blends of two or more of these organic compounds. Methyl esters of 4-hydroxybenzoic acid, propyl esters of 4-hydroxybenzoic acid, butyl ester of 4-hydroxybenzoic acid, ethyl ester of 4-hydroxybenzoic acid and blends of two or more of these organic compounds can also be used. Additional organic compounds also include but are not limited to 4-hydroxybenzoic acid methyl ester (named methyl PARABEN), 4-hydroxybenzoic acid propyl ester (named propyl PARABEN), which also are both amylase preservative agents. For further descriptions, see, e.g., U.S. Patent No. 5,281,526.

[0162] Addition of the organic compound precipitation agent provides the advantage of high flexibility of the precipitation conditions with respect to pH, temperature, variant amylase concentration, precipitation agent concentration, and time of incubation.

[0163] The organic compound precipitation agent is used in an amount effective to improve precipitation of the enzyme by means of the metal halide precipitation agent. The selection of at least an effective amount and an optimum amount of organic compound precipitation agent, as well as the conditions of the precipitation for maximum recovery including incubation time, pH, temperature and concentration of enzyme, will be readily apparent to one of ordinary skill in the art, in light of the present disclosure, after routine testing.

[0164] Generally, at least about 0.01% w/v of organic compound precipitation agent is added to the concentrated enzyme solution and usually at least about 0.02% w/v. Generally, no more than about 0.3% w/v of organic compound precipitation agent is added to the concentrated enzyme solution and usually no more than about 0.2% w/v.

[0165] The concentrated polypeptide solution, containing the metal halide precipitation agent, and the organic compound precipitation agent, can be adjusted to a pH, which will, of necessity, depend on the enzyme to be enriched or purified. Generally, the pH is adjusted at a level near the isoelectric point of the amylase. The pH can be adjusted at a pH in a range from about 2.5 pH units below the isoelectric point (pI) up to about 2.5 pH units above the isoelectric point.

[0166] The incubation time necessary to obtain an enriched or purified enzyme precipitate depends on the nature of the specific enzyme, the concentration of enzyme, and the specific precipitation agent(s) and its (their) concentration. Generally, the time effective to precipitate the enzyme is between about 1 to about 30 hours; usually it does not exceed about 25 hours. In the presence of the organic compound precipitation agent, the time of incubation can still be reduced to less about 10 hours and in most cases even about 6 hours.

[0167] Generally, the temperature during incubation is between about 4°C and about 50°C. Usually, the method is carried out at a temperature between about 10°C and about 45°C (e.g., between about 20°C and about 40°C). The optimal temperature for inducing precipitation varies according to the solution conditions and the enzyme or precipitation agent(s) used.

[0168] The overall recovery of enriched or purified enzyme precipitate, and the efficiency with which the process is conducted, is improved by agitating the solution comprising the enzyme, the added metal halide and the added organic compound. The agitation step is done both during addition of the metal halide and the organic compound, and during the subsequent incubation period. Suitable agitation methods include mechanical stirring or shaking, vigorous aeration, or any similar technique.

[0169] After the incubation period, the enriched or purified enzyme is then separated from the dissociated pigment and other impurities and collected by conventional separation techniques, such as filtration, centrifugation, microfiltration, rotary vacuum filtration, ultrafiltration, press filtration, cross membrane microfiltration, cross flow membrane microfiltration, or the like. Further enrichment or purification of the enzyme precipitate can be obtained by washing the precipitate with water. For example, the enriched or purified enzyme precipitate is washed with water containing the metal halide precipitation agent, or with water containing the metal halide and the organic compound precipitation agents.

[0170] During fermentation, a variant α -amylase polypeptide accumulates in the culture broth. For the isolation, enrichment, or purification of the desired variant α -amylase, the culture broth is centrifuged or filtered to eliminate cells, and the resulting cell-free liquid is used for enzyme enrichment or purification. In one embodiment, the cell-free broth is subjected to salting out using ammonium sulfate at about 70% saturation; the 70% saturation-precipitation fraction is then dissolved in a buffer and applied to a column such as a Sephadex G-100 column, and eluted to recover the enzyme-active fraction. For further enrichment or purification, a conventional procedure such as ion exchange chromatography may be used.

[0171] Enriched or purified enzymes are useful for laundry and cleaning applications. For example, they can be used in laundry detergents and spot removers. They can be made into a final product that is either liquid (solution, slurry) or solid (granular, powder).

[0172] A more specific example of enrichment or purification, is described in Sumitani et al. (2000) "New type of starch-binding domain: the direct repeat motif in the C-terminal region of *Bacillus* sp. 195 α -amylase contributes to starch binding and raw starch degrading," *Biochem. J.* 350: 477-484, and is briefly summarized here. The enzyme obtained from 4 liters of a *Streptomyces lividans* TK24 culture supernatant was treated with $(\text{NH}_4)_2\text{SO}_4$ at 80% saturation. The precipitate was recovered by centrifugation at $10,000 \times g$ (20 min. and 4°C) and re-dissolved in 20 mM Tris/HCl buffer (pH 7.0) containing 5 mM CaCl_2 . The solubilized precipitate was then dialyzed against the same buffer. The dialyzed sample was then applied to a Sephacryl S-200 column, which had previously been equilibrated with 20 mM Tris/HCl buffer, (pH 7.0), 5 mM CaCl_2 , and eluted at a linear flow rate of 7 mL/hr with the same buffer. Fractions from the column were collected and assessed for activity as judged by enzyme assay and SDS-PAGE. The protein was further purified as follows. A Toyopearl HW55 column (Tosoh Bioscience, Montgomeryville, PA; Cat. No. 19812) was equilibrated with 20 mM Tris/HCl buffer (pH 7.0) containing 5 mM CaCl_2 and 1.5 M $(\text{NH}_4)_2\text{SO}_4$. The enzyme was eluted with a linear gradient of 1.5 to 0 M $(\text{NH}_4)_2\text{SO}_4$ in 20 mM Tris/HCL buffer, pH 7.0 containing 5 mM CaCl_2 . The active fractions were collected, and the enzyme precipitated with $(\text{NH}_4)_2\text{SO}_4$ at 80% saturation. The precipitate was recovered, re-dissolved, and dialyzed as described above. The dialyzed sample was then applied to a Mono Q HR5/5 column (Amersham Pharmacia; Cat. No. 17-5167-01) previously equilibrated with 20 mM Tris/HCl buffer (pH 7.0) containing 5 mM CaCl_2 , at a flow rate of 60 mL/hour. The active fractions are collected and added to a 1.5 M $(\text{NH}_4)_2\text{SO}_4$ solution. The active enzyme fractions were re-chromatographed on a Toyopearl HW55 column, as before, to yield a homogeneous enzyme as determined by SDS-PAGE. See Sumitani et al. (2000) *Biochem. J.* 350: 477-484, for general discussion of the method and variations thereon.

[0173] For production scale recovery, variant α -amylase polypeptides can be enriched or partially purified as generally described above by removing cells via flocculation with polymers. Alternatively, the enzyme can be enriched or purified by microfiltration followed by concentration by ultrafiltration using available membranes and equipment. However, for some applications, the enzyme does not need to be enriched or purified, and whole broth culture can be lysed and used without further treatment. The enzyme can then be processed, for example, into granules.

4. Compositions and Uses of Variant Amylases

[0174] Variants amylases are useful for a variety of industrial applications. For example, variant amylases are useful in a starch conversion process, particularly in a saccharification process of a starch that has undergone liquefaction. The desired end-product may be any product that may be produced by the enzymatic conversion of the starch substrate. For example, the desired product may be a syrup rich in glucose and maltose, which can be used in other processes, such as the preparation of HFCS, or which can be converted into a number of other useful products, such as ascorbic acid intermediates (e.g., gluconate; 2-keto-L-gulonic acid; 5-keto-gluconate; and 2,5-diketogluconate); 1,3-propanediol; aromatic amino acids (e.g., tyrosine, phenylalanine and tryptophan); organic acids (e.g., lactate, pyruvate, succinate, isocitrate, and oxaloacetate); amino acids (e.g., serine and glycine); antibiotics; antimicrobials; enzymes; vitamins; and hormones.

[0175] The starch conversion process may be a precursor to, or simultaneous with, a fermentation process designed

to produce alcohol for fuel or drinking (*i.e.*, potable alcohol). One skilled in the art is aware of various fermentation conditions that may be used in the production of these end-products. Variant amylases are also useful in compositions and methods of food preparation. These various uses of variant amylases are described in more detail below.

4.1. Preparation of Starch Substrates

[0176] Those of general skill in the art are well aware of available methods that may be used to prepare starch substrates for use in the processes disclosed herein. For example, a useful starch substrate may be obtained from tubers, roots, stems, legumes, cereals or whole grain. More specifically, the granular starch may be obtained from corn, cobs, wheat, barley, rye, triticale, milo, sago, millet, cassava, tapioca, sorghum, rice, peas, bean, banana, or potatoes. Corn contains about 60-68% starch; barley contains about 55-65% starch; millet contains about 75-80% starch; wheat contains about 60-65% starch; and polished rice contains 70-72% starch. Specifically contemplated starch substrates are corn starch and wheat starch. The starch from a grain may be ground or whole and includes corn solids, such as kernels, bran and/or cobs. The starch may also be highly refined raw starch or feedstock from starch refinery processes. Various starches also are commercially available. For example, corn starch is available from Cerestar, Sigma, and Katayama Chemical Industry Co. (Japan); wheat starch is available from Sigma; sweet potato starch is available from Wako Pure Chemical Industry Co. (Japan); and potato starch is available from Nakaari Chemical Pharmaceutical Co. (Japan).

[0177] The starch substrate can be a crude starch from milled whole grain, which contains non-starch fractions, *e.g.*, germ residues and fibers. Milling may comprise either wet milling or dry milling or grinding. In wet milling, whole grain is soaked in water or dilute acid to separate the grain into its component parts, *e.g.*, starch, protein, germ, oil, kernel fibers. Wet milling efficiently separates the germ and meal (*i.e.*, starch granules and protein) and is especially suitable for production of syrups. In dry milling or grinding, whole kernels are ground into a fine powder and often processed without fractionating the grain into its component parts. In some cases, oils from the kernels are recovered. Dry ground grain thus will comprise significant amounts of non-starch carbohydrate compounds, in addition to starch. Dry grinding of the starch substrate can be used for production of ethanol and other biochemicals. The starch to be processed may be a highly refined starch quality, for example, at least 90%, at least 95%, at least 97%, or at least 99.5% pure.

4.2. Gelatinization and Liquefaction of Starch

[0178] As used herein, the term "liquefaction" or "liquefy" means a process by which starch is converted to less viscous and shorter chain dextrans. Generally, this process involves gelatinization of starch simultaneously with or followed by the addition of an α -amylase, although additional liquefaction-inducing enzymes optionally may be added. In some embodiments, the starch substrate prepared as described above is slurried with water. The starch slurry may contain starch as a weight percent of dry solids of about 10-55%, about 20-45%, about 30-45%, about 30-40%, or about 30-35%. α -Amylase (EC 3.2.1.1) may be added to the slurry, with a metering pump, for example. The α -amylase typically used for this application is a thermally stable, bacterial α -amylase, such as a *Geobacillus stearothermophilus* α -amylase. The α -amylase is usually supplied, for example, at about 1500 units per kg dry matter of starch. To optimize α -amylase stability and activity, the pH of the slurry typically is adjusted to about pH 5.5-6.5 and about 1 mM of calcium (about 40 ppm free calcium ions) can also be added. *Geobacillus stearothermophilus* variants or other α -amylases may require different conditions. Bacterial α -amylase remaining in the slurry following liquefaction may be deactivated via a number of methods, including lowering the pH in a subsequent reaction step or by removing calcium from the slurry in cases where the enzyme is dependent upon calcium.

[0179] The slurry of starch plus the α -amylase may be pumped continuously through a jet cooker, which is steam heated to 105°C. Gelatinization occurs rapidly under these conditions, and the enzymatic activity, combined with the significant shear forces, begins the hydrolysis of the starch substrate. The residence time in the jet cooker is brief. The partly gelatinized starch may be passed into a series of holding tubes maintained at 105-110°C and held for 5-8 min. to complete the gelatinization process ("primary liquefaction"). Hydrolysis to the required DE is completed in holding tanks at 85-95°C or higher temperatures for about 1 to 2 hours ("secondary liquefaction"). These tanks may contain baffles to discourage back mixing. As used herein, the term "minutes of secondary liquefaction" refers to the time that has elapsed from the start of secondary liquefaction to the time that the Dextrose Equivalent (DE) is measured. The slurry is then allowed to cool to room temperature. This cooling step can be 30 minutes to 180 minutes, *e.g.* 90 minutes to 120 minutes. The liquefied starch typically is in the form of a slurry having a dry solids content (w/w) of about 10-50%; about 10-45%; about 15-40%; about 20-40%; about 25-40%; or about 25-35%.

[0180] Liquefaction with variant amylases advantageously can be conducted at low pH, eliminating the requirement to adjust the pH to about pH 5.5-6.5. Variant amylases can be used for liquefaction at a pH range of 2 to 7, *e.g.*, pH 3.0 - 7.5, pH 4.0 - 6.0, or pH 4.5 - 5.8. Variant amylases can maintain liquefying activity at a temperature range of about 85°C - 95°C, *e.g.*, 85°C, 90°C, or 95°C. For example, liquefaction can be conducted with 800 μ g an amylase in a solution of 25% DS corn starch for 10 min at pH 5.8 and 85°C, or pH 4.5 and 95°C, for example. Liquefying activity can be

assayed using any of a number of known viscosity assays in the art.

4.3. Saccharification

[0181] The liquefied starch can be saccharified into a syrup rich in lower DP (e.g., DP1 + DP2) saccharides, using variant amylases, optionally in the presence of another enzyme(s). The exact composition of the products of saccharification depends on the combination of enzymes used, as well as the type of granular starch processed. Advantageously, the syrup obtainable using the provided variant amylases may contain a weight percent of DP2 of the total oligosaccharides in the saccharified starch exceeding 30%, e.g., 45% - 65% or 55% - 65%. The weight percent of (DP1 + DP2) in the saccharified starch may exceed about 70%, e.g., 75% - 85% or 80% - 85%. The present amylases also produce a relatively high yield of glucose, e.g., DP1 > 20%, in the syrup product.

[0182] Whereas liquefaction is generally run as a continuous process, saccharification is often conducted as a batch process. Saccharification typically is most effective at temperatures of about 60-65°C and a pH of about 4.0-4.5, e.g., pH 4.3, necessitating cooling and adjusting the pH of the liquefied starch. Saccharification may be performed, for example, at a temperature between about 40°C, about 50°C, or about 55°C to about 60°C or about 65°C. Saccharification is normally conducted in stirred tanks, which may take several hours to fill or empty. Enzymes typically are added either at a fixed ratio to dried solids as the tanks are filled or added as a single dose at the commencement of the filling stage. A saccharification reaction to make a syrup typically is run over about 24-72 hours, for example, 24-48 hours. When a maximum or desired DE has been attained, the reaction is stopped by heating to 85°C for 5 min., for example. Further incubation will result in a lower DE, eventually to about 90 DE, as accumulated glucose re-polymerizes to isomaltose and/or other reversion products via an enzymatic reversion reaction and/or with the approach of thermodynamic equilibrium. When using an amylase, saccharification optimally is conducted at a temperature range of about 30°C to about 75°C, e.g., 45°C - 75°C or 47°C - 74°C. The saccharifying may be conducted over a pH range of about pH 3 to about pH 7, e.g., pH 3.0 - pH 7.5, pH 3.5 - pH 5.5, pH 3.5, pH 3.8, or pH 4.5.

[0183] An amylase may be added to the slurry in the form of a composition. Amylase can be added to a slurry of a granular starch substrate in an amount of about 0.6 - 10 ppm ds, e.g., 2 ppm ds. An amylase can be added as a whole broth, clarified, enriched, partially purified, or purified enzyme. The specific activity of the amylase may be about 300 U/mg of enzyme, for example, measured with the PAHBAH assay. The amylase also can be added as a whole broth product.

[0184] An amylase may be added to the slurry as an isolated enzyme solution. For example, an amylase can be added in the form of a cultured cell material produced by host cells expressing an amylase. An amylase may also be secreted by a host cell into the reaction medium during the fermentation or SSF process, such that the enzyme is provided continuously into the reaction. The host cell producing and secreting amylase may also express an additional enzyme, such as a glucoamylase. For example, U.S. Patent No. 5,422,267 discloses the use of a glucoamylase in yeast for production of alcoholic beverages. For example, a host cell, e.g., *Trichoderma reesei* or *Aspergillus niger*, may be engineered to co-express an amylase and a glucoamylase, e.g., HgGA, TrGA, or a TrGA variant, during saccharification. The host cell can be genetically modified so as not to express its endogenous glucoamylase and/or other enzymes, proteins or other materials. The host cell can be engineered to express a broad spectrum of various saccharolytic enzymes. For example, the recombinant yeast host cell can comprise nucleic acids encoding a glucoamylase, an alpha-glucosidase, an enzyme that utilizes pentose sugar, an α -amylase, a pullulanase, an isoamylase, and/or an isopullulanase. See, e.g., WO 2011/153516 A2.

4.4. Isomerization

[0185] The soluble starch hydrolysate produced by treatment with amylase can be converted into high fructose starch-based syrup (HFSS), such as high fructose corn syrup (HFCS). This conversion can be achieved using a glucose isomerase, particularly a glucose isomerase immobilized on a solid support. The pH is increased to about 6.0 to about 8.0, e.g., pH 7.5 (depending on the isomerase), and Ca^{2+} is removed by ion exchange. Suitable isomerases include Sweetzyme®, IT (Novozymes A/S); G-zyme® IMGI, and G-zyme® G993, Ketomax®, G-zyme® G993, G-zyme® G993 liquid, and GenSweet® IGI. Following isomerization, the mixture typically contains about 40-45% fructose, e.g., 42% fructose.

4.5. Fermentation

[0186] The soluble starch hydrolysate, particularly a glucose rich syrup, can be fermented by contacting the starch hydrolysate with a fermenting organism typically at a temperature around 32°C, such as from 30°C to 35°C for alcohol-producing yeast. The temperature and pH of the fermentation will depend upon the fermenting organism. EOF products include metabolites, such as citric acid, lactic acid, succinic acid, monosodium glutamate, gluconic acid, sodium gluco-

nate, calcium gluconate, potassium gluconate, itaconic acid and other carboxylic acids, glucono delta-lactone, sodium erythorbate, lysine and other amino acids, omega 3 fatty acid, butanol, isoprene, 1,3-propanediol and other biomaterials.

[0187] Ethanologenic microorganisms include yeast, such as *Saccharomyces cerevisiae* and bacteria, e.g., *Zymomonas mobilis*, expressing alcohol dehydrogenase and pyruvate decarboxylase. The ethanologenic microorganism can express xylose reductase and xylitol dehydrogenase, which convert xylose to xylulose. Improved strains of ethanologenic microorganisms, which can withstand higher temperatures, for example, are known in the art and can be used. See Liu et al. (2011) Sheng Wu Gong Cheng Xue Bao 27(7): 1049-56. Commercial sources of yeast include ETHANOL RED® (LeSaffre); Thermosacc® (Lallemand); RED STAR® (Red Star); FERMIOL® (DSM Specialties); and SUPER-START® (Alltech). Microorganisms that produce other metabolites, such as citric acid and lactic acid, by fermentation are also known in the art. See, e.g., Papagianni (2007) "Advances in citric acid fermentation by *Aspergillus niger*: biochemical aspects, membrane transport and modeling," Biotechnol. Adv. 25(3): 244-63; John et al. (2009) "Direct lactic acid fermentation: focus on simultaneous saccharification and lactic acid production," Biotechnol. Adv. 27(2): 145-52.

[0188] The saccharification and fermentation processes may be carried out as an SSF process. Fermentation may comprise subsequent enrichment, purification, and recovery of ethanol, for example. During the fermentation, the ethanol content of the broth or "beer" may reach about 8-18% v/v, e.g., 14-15% v/v. The broth may be distilled to produce enriched, e.g., 96% pure, solutions of ethanol. Further, CO₂ generated by fermentation may be collected with a CO₂ scrubber, compressed, and marketed for other uses, e.g., carbonating beverage or dry ice production. Solid waste from the fermentation process may be used as protein-rich products, e.g., livestock feed.

[0189] As mentioned above, an SSF process can be conducted with fungal cells that express and secrete amylase continuously throughout SSF. The fungal cells expressing amylase also can be the fermenting microorganism, e.g., an ethanologenic microorganism. Ethanol production thus can be carried out using a fungal cell that expresses sufficient amylase so that less or no enzyme has to be added exogenously. The fungal host cell can be from an appropriately engineered fungal strain. Fungal host cells that express and secrete other enzymes, in addition to amylase, also can be used. Such cells may express glucoamylase and/or a pullulanase, phytase, *alpha*-glucosidase, isoamylase, beta-amylase cellulase, xylanase, other hemicellulases, protease, *beta*-glucosidase, pectinase, esterase, redox enzymes, transferase, or other enzyme.

[0190] A variation on this process is a "fed-batch fermentation" system, where the substrate is added in increments as the fermentation progresses. Fed-batch systems are useful when catabolite repression may inhibit the metabolism of the cells and where it is desirable to have limited amounts of substrate in the medium. The actual substrate concentration in fed-batch systems is estimated by the changes of measurable factors such as pH, dissolved oxygen and the partial pressure of waste gases, such as CO₂. Batch and fed-batch fermentations are common and well known in the art.

[0191] Continuous fermentation is an open system where a defined fermentation medium is added continuously to a bioreactor, and an equal amount of conditioned medium is removed simultaneously for processing. Continuous fermentation generally maintains the cultures at a constant high density where cells are primarily in log phase growth. Continuous fermentation permits modulation of cell growth and/or product concentration. For example, a limiting nutrient such as the carbon source or nitrogen source is maintained at a fixed rate and all other parameters are allowed to moderate. Because growth is maintained at a steady state, cell loss due to medium being drawn off should be balanced against the cell growth rate in the fermentation. Methods of optimizing continuous fermentation processes and maximizing the rate of product formation are well known in the art of industrial microbiology.

4.6. Compositions Comprising Variants Amylases

[0192] Variant amylases may be combined with a glucoamylase (EC 3.2.1.3), e.g., a *Trichoderma* glucoamylase or variant thereof. An exemplary glucoamylase is *Trichoderma reesei* glucoamylase (TrGA) and variants thereof that possess superior specific activity and thermal stability. See U.S. Published Applications Nos. 2006/0094080, 2007/0004018, and 2007/0015266 (Danisco US Inc.). Suitable variants of TrGA include those with glucoamylase activity and at least 80%, at least 90%, or at least 95% sequence identity to wild-type TrGA. Variant amylases advantageously increase the yield of glucose produced in a saccharification process catalyzed by TrGA.

[0193] Alternatively, the glucoamylase may be another glucoamylase derived from plants (including algae), fungi, or bacteria. For example, the glucoamylases may be *Aspergillus niger* G1 or G2 glucoamylase or its variants (e.g., Boel et al. (1984) EMBO J. 3: 1097-1102; WO 92/00381; WO 00/04136 (Novo Nordisk A/S)); and A. *awamori* glucoamylase (e.g., WO 84/02921 (Cetus Corp.)). Other contemplated *Aspergillus* glucoamylase include variants with enhanced thermal stability, e.g., G137A and G139A (Chen et al. (1996) Prot. Eng. 9: 499-505); D257E and D293E/Q (Chen et al. (1995) Prot. Eng. 8: 575-582); N182 (Chen et al. (1994) Biochem. J. 301: 275-281); A246C (Fierobe et al. (1996) Biochemistry, 35: 8698-8704); and variants with Pro residues in positions A435 and S436 (Li et al. (1997) Protein Eng. 10: 1199-1204). Other contemplated glucoamylases include *Talaromyces* glucoamylases, in particular derived from *T. emersonii* (e.g., WO 99/28448 (Novo Nordisk A/S), *T. leycettanus* (e.g., U.S. Patent No. RE 32,153 (CPC International, Inc.)), *T. duponti*,

or *T. thermophilus* (e.g., U.S. Patent No. 4,587,215). Contemplated bacterial glucoamylases include glucoamylases from the genus *Clostridium*, in particular *C. thermoamylolyticum* (e.g., EP 135,138 (CPC International, Inc.) and *C. thermohydrosulfuricum* (e.g., WO 86/01831 (Michigan Biotechnology Institute)). Suitable glucoamylases include the glucoamylases derived from *Aspergillus oryzae*, such as a glucoamylase shown in SEQ ID NO:2 in WO 00/04136 (Novo Nordisk A/S). Also suitable are commercial glucoamylases, such as AMG 200L; AMG 300 L; SAN™ SUPER and AMG™ E (Novozymes); OPTIDEX® 300 and OPTIDEX L-400 (Danisco US Inc.); AMIGASE™ and AMIGASE™ PLUS (DSM); G-ZYME® G900 (Enzyme Bio-Systems); and G-ZYME® G990 ZR (*A. niger* glucoamylase with a low protease content). Still other suitable glucoamylases include *Aspergillus fumigatus* glucoamylase, *Talaromyces* glucoamylase, *Thielavia* glucoamylase, *Trametes* glucoamylase, *Thermomyces* glucoamylase, *Athelia* glucoamylase, or *Humicola* glucoamylase (e.g., HgGA). Glucoamylases typically are added in an amount of about 0.1 - 2 glucoamylase units (GAU)/g ds, e.g., about 0.16 GAU/g ds, 0.23 GAU/g ds, or 0.33 GAU/g ds.

[0194] Other suitable enzymes that can be used with amylase include a phytase, protease, pullulanase, β -amylase, isoamylase, a different α -amylase, alpha-glucosidase, cellulase, xylanase, other hemicellulases, beta-glucosidase, transferase, pectinase, lipase, cutinase, esterase, redox enzymes, or a combination thereof. For example, a debranching enzyme, such as an isoamylase (EC 3.2.1.68), may be added in effective amounts well known to the person skilled in the art. A pullulanase (EC 3.2.1.41), e.g., Promozyme®, is also suitable. Pullulanase typically is added at 100 U/kg ds. Further suitable enzymes include proteases, such as fungal and bacterial proteases. Fungal proteases include those obtained from *Aspergillus*, such as *A. niger*, *A. awamori*, *A. oryzae*; *Mucor* (e.g., *M. miehei*); *Rhizopus*; and *Trichoderma*.

[0195] β -Amylases (EC 3.2.1.2) are exo-acting maltogenic amylases, which catalyze the hydrolysis of 1,4- α -glucosidic linkages into amylopectin and related glucose polymers, thereby releasing maltose. β -Amylases have been isolated from various plants and microorganisms. See Fogarty et al. (1979) in PROGRESS IN INDUSTRIAL MICROBIOLOGY, Vol. 15, pp. 112-115. These β -Amylases have optimum temperatures in the range from 40°C to 65°C and optimum pH in the range from about 4.5 to about 7.0. Contemplated β -amylases include, but are not limited to, β -amylases from barley Spezyme® BBA 1500, Spezyme® DBA, Optimalt™ ME, Optimalt™ BBA (Danisco US Inc.); and Novozym™ WBA (Novozymes A/S).

[0196] Compositions comprising the present amylases may be aqueous or non-aqueous formulations, granules, powders, gels, slurries, pastes, etc., which may further comprise any one or more of the additional enzymes listed, herein, along with buffers, salts, preservatives, water, co-solvents, surfactants, and the like. Such compositions may work in combination with endogenous enzymes or other ingredients already present in a slurry, water bath, washing machine, food or drink product, etc, for example, endogenous plant (including algal) enzymes, residual enzymes from a prior processing step, and the like.

5. Compositions and Methods for Baking and Food Preparation

[0197] The present disclosure also relates to a "food composition," including but not limited to a food product, animal feed and/or food/feed additives, comprising an amylase, and methods for preparing such a food composition comprising mixing variant amylase with one or more food ingredients, or uses thereof.

[0198] Furthermore, the present disclosure relates to the use of an amylase in the preparation of a food composition, wherein the food composition is baked subsequent to the addition of the polypeptide of the invention. As used herein the term "baking composition" means any composition and/or additive prepared in the process of providing a baked food product, including but not limited to bakers flour, a dough, a baking additive and/or a baked product. The food composition or additive may be liquid or solid.

[0199] As used herein, the term "flour" means milled or ground cereal grain. The term "flour" also may mean Sago or tuber products that have been ground or mashed. In some embodiments, flour may also contain components in addition to the milled or mashed cereal or plant matter. An example of an additional component, although not intended to be limiting, is a leavening agent. Cereal grains include wheat, oat, rye, and barley. Tuber products include tapioca flour, cassava flour, and custard powder. The term "flour" also includes ground corn flour, maize-meal, rice flour, whole-meal flour, self-rising flour, tapioca flour, cassava flour, ground rice, enriched flower, and custard powder.

[0200] For the commercial and home use of flour for baking and food production, it is important to maintain an appropriate level of α -amylase activity in the flour. A level of activity that is too high may result in a product that is sticky and/or doughy and therefore unmarketable. Flour with insufficient α -amylase activity may not contain enough sugar for proper yeast function, resulting in dry, crumbly bread, or baked products. Accordingly, an amylase, by itself or in combination with another α -amylase(s), may be added to the flour to augment the level of endogenous α -amylase activity in flour.

[0201] An amylase can further be added alone or in a combination with other amylases to prevent or retard staling, i.e., crumb firming of baked products. The amount of anti-staling amylase will typically be in the range of 0.01-10 mg of enzyme protein per kg of flour, e.g., 0.5 mg/kg ds. Additional anti-staling amylases that can be used in combination with an amylase include an endo-amylase, e.g., a bacterial endo-amylase from *Bacillus*. The additional amylase can be another maltogenic α -amylase (EC 3.2.1.133), e.g., from *Bacillus*. Novamyl® is an exemplary maltogenic α -amylase

from *B. stearrowthermophilus* strain NCIB 11837 and is described in Christophersen et al. (1997) Starch 50: 39-45. Other examples of anti-staling endo-amylases include bacterial α -amylases derived from *Bacillus*, such as *B. licheniformis* or *B. amyloliquefaciens*. The anti-staling amylase may be an exo-amylase, such as β -amylase, e.g., from plant sources, such as soy bean, or from microbial sources, such as *Bacillus*.

[0202] The baking composition comprising an amylase further can comprise a phospholipase or enzyme with phospholipase activity. An enzyme with phospholipase activity has an activity that can be measured in Lipase Units (LU). The phospholipase may have A_1 or A_2 activity to remove fatty acid from the phospholipids, forming a lysophospholipid. It may or may not have lipase activity, i.e., activity on triglyceride substrates. The phospholipase typically has a temperature optimum in the range of 30-90°C., e.g., 30-70°C. The added phospholipases can be of animal origin, for example, from pancreas, e.g., bovine or porcine pancreas, snake venom or bee venom. Alternatively, the phospholipase may be of microbial origin, e.g., from filamentous fungi, yeast or bacteria, for example.

[0203] The phospholipase is added in an amount that improves the softness of the bread during the initial period after baking, particularly the first 24 hours. The amount of phospholipase will typically be in the range of 0.01-10 mg of enzyme protein per kg of flour, e.g., 0.1-5 mg/kg. That is, phospholipase activity generally will be in the range of 20-1000 LU/kg of flour, where a Lipase Unit is defined as the amount of enzyme required to release 1 μ mol butyric acid per minute at 30°C, pH 7.0, with gum arabic as emulsifier and tributyrin as substrate.

[0204] Compositions of dough generally comprise wheat meal or wheat flour and/or other types of meal, flour or starch such as corn flour, cornstarch, rye meal, rye flour, oat flour, oatmeal, soy flour, sorghum meal, sorghum flour, potato meal, potato flour or potato starch. The dough may be fresh, frozen or par-baked. The dough can be a leavened dough or a dough to be subjected to leavening. The dough may be leavened in various ways, such as by adding chemical leavening agents, e.g., sodium bicarbonate or by adding a leaven, i.e., fermenting dough. Dough also may be leavened by adding a suitable yeast culture, such as a culture of *Saccharomyces cerevisiae* (baker's yeast), e.g., a commercially available strain of *S. cerevisiae*.

[0205] The dough may also comprise other conventional dough ingredients, e.g., proteins, such as milk powder, gluten, and soy; eggs (e.g., whole eggs, egg yolks or egg whites); an oxidant, such as ascorbic acid, potassium bromate, potassium iodate, azodicarbonamide (ADA) or ammonium persulfate; an amino acid such as L-cysteine; a sugar; or a salt, such as sodium chloride, calcium acetate, sodium sulfate or calcium sulfate. The dough further may comprise fat, e.g., triglyceride, such as granulated fat or shortening. The dough further may comprise an emulsifier such as mono- or diglycerides, diacetyl tartaric acid esters of mono- or diglycerides, sugar esters of fatty acids, polyglycerol esters of fatty acids, lactic acid esters of monoglycerides, acetic acid esters of monoglycerides, polyoxyethylene stearates, or lysol-ecithin. In particular, the dough can be made without addition of emulsifiers.

[0206] The dough product may be any processed dough product, including fried, deep fried, roasted, baked, steamed and boiled doughs, such as steamed bread and rice cakes. In one embodiment, the food product is a bakery product. Typical bakery (baked) products include bread - such as loaves, rolls, buns, bagels, pizza bases etc. pastry, pretzels, tortillas, cakes, cookies, biscuits, crackers etc.

[0207] Optionally, an additional enzyme may be used together with the anti-staling amylase and the phospholipase. The additional enzyme may be a second amylase, such as an amyloglucosidase, a β -amylase, a cyclodextrin glucanotransferase, or the additional enzyme may be a peptidase, in particular an exopeptidase, a transglutaminase, a lipase, a cellulase, a xylanase, a protease, a protein disulfide isomerase, e.g., a protein disulfide isomerase as disclosed in WO 95/00636, for example, a glycosyltransferase, a branching enzyme (1,4- α -glucan branching enzyme), a 4- α -glucanotransferase (dextrin glycosyltransferase) or an oxidoreductase, e.g., a peroxidase, a laccase, a glucose oxidase, a pyranose oxidase, a lipooxygenase, an L-amino acid oxidase or a carbohydrate oxidase. The additional enzyme(s) may be of any origin, including mammalian and plant, and particularly of microbial (bacterial, yeast or fungal) origin and may be obtained by techniques conventionally used in the art.

[0208] The xylanase is typically of microbial origin, e.g., derived from a bacterium or fungus, such as a strain of *Aspergillus*. Xylanases include Pentopan® and Novozym 384®, for example, which are commercially available xylanase preparations produced from *Trichoderma reesei*. The amyloglucosidase may be an *A. niger* amyloglucosidase (such as AMG®). Other useful amylase products include Grindamyl® A 1000 or A 5000 (Grindsted Products, Denmark) and Amylase® H or Amylase® P (DSM). The glucose oxidase may be a fungal glucose oxidase, in particular an *Aspergillus niger* glucose oxidase (such as Gluzyne®). An exemplary protease is Neutrase®.

[0209] The process may be used for any kind of baked product prepared from dough, either of a soft or a crisp character, either of a white, light or dark type. Examples are bread, particularly white, whole-meal or rye bread, typically in the form of loaves or rolls, such as, but not limited to, French baguette-type bread, pita bread, tortillas, cakes, pancakes, biscuits, cookies, pie crusts, crisp bread, steamed bread, pizza and the like.

[0210] An amylase may be used in a pre-mix, comprising flour together with an anti-staling amylase, a phospholipase, and/or a phospholipid. The pre-mix may contain other dough-improving and/or bread-improving additives, e.g., any of the additives, including enzymes, mentioned above. An amylase can be a component of an enzyme preparation comprising an anti-staling amylase and a phospholipase, for use as a baking additive.

[0211] The enzyme preparation is optionally in the form of a granulate or agglomerated powder. The preparation can have a narrow particle size distribution with more than 95% (by weight) of the particles in the range from 25 to 500 μm . Granulates and agglomerated powders may be prepared by conventional methods, e.g., by spraying an amylase onto a carrier in a fluid-bed granulator. The carrier may consist of particulate cores having a suitable particle size. The carrier

may be soluble or insoluble, e.g., a salt (such as NaCl or sodium sulfate), a sugar (such as sucrose or lactose), a sugar alcohol (such as sorbitol), starch, rice, corn grits, or soy.

[0212] Enveloped particles, i.e., α -amylase particles, can comprise an amylase. To prepare enveloped α -amylase particles, the enzyme is contacted with a food grade lipid in sufficient quantity to suspend all of the α -amylase particles. Food grade lipids, as used herein, may be any naturally organic compound that is insoluble in water but is soluble in non-polar organic solvents such as hydrocarbon or diethyl ether. Suitable food grade lipids include, but are not limited to, triglycerides either in the form of fats or oils that are either saturated or unsaturated. Examples of fatty acids and combinations thereof which make up the saturated triglycerides include, but are not limited to, butyric (derived from milk fat), palmitic (derived from animal and plant fat), and/or stearic (derived from animal and plant fat). Examples of fatty acids and combinations thereof which make up the unsaturated triglycerides include, but are not limited to, palmitoleic (derived from animal and plant fat), oleic (derived from animal and plant fat), linoleic (derived from plant oils), and/or linolenic (derived from linseed oil). Other suitable food grade lipids include, but are not limited to, monoglycerides and diglycerides derived from the triglycerides discussed above, phospholipids and glycolipids.

[0213] The food grade lipid, particularly in the liquid form, is contacted with a powdered form of the α -amylase particles in such a fashion that the lipid material covers at least a portion of the surface of at least a majority, e.g., 100% of the α -amylase particles. Thus, each α -amylase particle is individually enveloped in a lipid. For example, all or substantially all of the α -amylase particles are provided with a thin, continuous, enveloping film of lipid. This can be accomplished by first pouring a quantity of lipid into a container, and then slurring the α -amylase particles so that the lipid thoroughly wets the surface of each α -amylase particle. After a short period of stirring, the enveloped α -amylase particles, carrying a substantial amount of the lipids on their surfaces, are recovered. The thickness of the coating so applied to the particles of α -amylase can be controlled by selection of the type of lipid used and by repeating the operation in order to build up a thicker film, when desired.

[0214] The storing, handling and incorporation of the loaded delivery vehicle can be accomplished by means of a packaged mix. The packaged mix can comprise the enveloped α -amylase. However, the packaged mix may further contain additional ingredients as required by the manufacturer or baker. After the enveloped α -amylase has been incorporated into the dough, the baker continues through the normal production process for that product.

[0215] The advantages of enveloping the α -amylase particles are two-fold. First, the food grade lipid protects the enzyme from thermal denaturation during the baking process for those enzymes that are heat labile. Consequently, while the α -amylase is stabilized and protected during the proving and baking stages, it is released from the protective coating in the final baked good product, where it hydrolyzes the glucosidic linkages in polyglucans. The loaded delivery vehicle also provides a sustained release of the active enzyme into the baked good. That is, following the baking process, active α -amylase is continually released from the protective coating at a rate that counteracts, and therefore reduces the rate of, staling mechanisms.

[0216] In general, the amount of lipid applied to the α -amylase particles can vary from a few percent of the total weight of the α -amylase to many times that weight, depending upon the nature of the lipid, the manner in which it is applied to the α -amylase particles, the composition of the dough mixture to be treated, and the severity of the dough-mixing operation involved.

[0217] The loaded delivery vehicle, i.e., the lipid-enveloped enzyme, is added to the ingredients used to prepare a baked good in an effective amount to extend the shelf-life of the baked good. The baker computes the amount of enveloped α -amylase, prepared as discussed above, that will be required to achieve the desired anti-staling effect. The amount of the enveloped α -amylase required is calculated based on the concentration of enzyme enveloped and on the proportion of α -amylase to flour specified. A wide range of concentrations has been found to be effective, although, as has been discussed, observable improvements in anti-staling do not correspond linearly with the α -amylase concentration, but above certain minimal levels, large increases in α -amylase concentration produce little additional improvement. The α -amylase concentration actually used in a particular bakery production could be much higher than the minimum necessary to provide the baker with some insurance against inadvertent under-measurement errors by the baker. The lower limit of enzyme concentration is determined by the minimum anti-staling effect the baker wishes to achieve.

[0218] A method of preparing a baked good may comprise: a) preparing lipid-coated α -amylase particles, where substantially all of the α -amylase particles are coated; b) mixing a dough containing flour; c) adding the lipid-coated α -amylase to the dough before the mixing is complete and terminating the mixing before the lipid coating is removed from the α -amylase; d) proofing the dough; and e) baking the dough to provide the baked good, where the α -amylase is inactive during the mixing, proofing and baking stages and is active in the baked good.

[0219] The enveloped α -amylase can be added to the dough during the mix cycle, e.g., near the end of the mix cycle. The enveloped α -amylase is added at a point in the mixing stage that allows sufficient distribution of the enveloped α -

amylase throughout the dough; however, the mixing stage is terminated before the protective coating becomes stripped from the α -amylase particle(s). Depending on the type and volume of dough, and mixer action and speed, anywhere from one to six minutes or more might be required to mix the enveloped α -amylase into the dough, but two to four minutes is average. Thus, several variables may determine the precise procedure. First, the quantity of enveloped α -amylase should have a total volume sufficient to allow the enveloped α -amylase to be spread throughout the dough mix. If the preparation of enveloped α -amylase is highly concentrated, additional oil may need to be added to the pre-mix before the enveloped α -amylase is added to the dough. Recipes and production processes may require specific modifications; however, good results generally can be achieved when 25% of the oil specified in a bread dough formula is held out of the dough and is used as a carrier for a concentrated enveloped α -amylase when added near the end of the mix cycle. In bread or other baked goods, particularly those having a low fat content, e.g., French-style breads, an enveloped α -amylase mixture of approximately 1% of the dry flour weight is sufficient to admix the enveloped α -amylase properly with the dough. The range of suitable percentages is wide and depends on the formula, finished product, and production methodology requirements of the individual baker. Second, the enveloped α -amylase suspension should be added to the mix with sufficient time for complete mixture into the dough, but not for such a time that excessive mechanical action strips the protective lipid coating from the enveloped α -amylase particles.

[0220] In a further aspect, the food composition is an oil, meat, lard, composition comprising an amylase. In this context the term "[oil/meat/lard] composition" means any composition, based on, made from and/or containing oil, meat or lard, respectively. Another aspect relates to a method of preparing an oil or meat or lard composition and/or additive comprising an amylase, comprising mixing the polypeptide of the invention with a oil/meat/lard composition and/or additive ingredients.

[0221] In a further aspect, the food composition is an animal feed composition, animal feed additive and/or pet food comprising an amylase and variants thereof. The present disclosure further relates to a method for preparing such an animal feed composition, animal feed additive composition and/or pet food comprising mixing an amylase and variants thereof with one or more animal feed ingredients and/or animal feed additive ingredients and/or pet food ingredients. Furthermore, the present disclosure relates to the use of an amylase in the preparation of an animal feed composition and/or animal feed additive composition and/or pet food.

[0222] The term "animal" includes all non-ruminant and ruminant animals. In a particular embodiment, the animal is a non-ruminant animal, such as a horse and a mono-gastric animal. Examples of mono-gastric animals include, but are not limited to, pigs and swine, such as piglets, growing pigs, sows; poultry such as turkeys, ducks, chicken, broiler chicks, layers; fish such as salmon, trout, tilapia, catfish and carps; and crustaceans such as shrimps and prawns. In a further embodiment the animal is a ruminant animal including, but not limited to, cattle, young calves, goats, sheep, giraffes, bison, moose, elk, yaks, water buffalo, deer, camels, alpacas, llamas, antelope, pronghorn and nilgai.

[0223] In the present context, it is intended that the term "pet food" is understood to mean a food for a household animal such as, but not limited to dogs, cats, gerbils, hamsters, chinchillas, fancy rats, guinea pigs; avian pets, such as canaries, parakeets, and parrots; reptile pets, such as turtles, lizards and snakes; and aquatic pets, such as tropical fish and frogs.

[0224] The terms "animal feed composition," "feedstuff" and "fodder" are used interchangeably and may comprise one or more feed materials selected from the group comprising a) cereals, such as small grains (e.g., wheat, barley, rye, oats and combinations thereof) and/or large grains such as maize or sorghum; b) by products from cereals, such as corn gluten meal, Distillers Dried Grain Solubles (DDGS) (particularly corn based Distillers Dried Grain Solubles (cDDGS), wheat bran, wheat middlings, wheat shorts, rice bran, rice hulls, oat hulls, palm kernel, and citrus pulp; c) protein obtained from sources such as soya, sunflower, peanut, lupin, peas, fava beans, cotton, canola, fish meal, dried plasma protein, meat and bone meal, potato protein, whey, copra, sesame; d) oils and fats obtained from vegetable and animal sources; e) minerals and vitamins.

6. Textile Desizing Compositions and Use

[0225] Also contemplated are compositions and methods of treating fabrics (e.g., to desize a textile) using a variant α -amylase as defined in the appended claims. Fabric-treating methods are well known in the art (see, e.g., U.S. Patent No.6,077,316). For example, the feel and appearance of a fabric can be improved by a method comprising contacting the fabric with an amylase in a solution. The fabric can be treated with the solution under pressure.

[0226] An amylase can be applied during or after the weaving of a textile, or during the desizing stage, or one or more additional fabric processing steps. During the weaving of textiles, the threads are exposed to considerable mechanical strain. Prior to weaving on mechanical looms, warp yarns are often coated with sizing starch or starch derivatives to increase their tensile strength and to prevent breaking. An amylase can be applied during or after the weaving to remove these sizing starch or starch derivatives. After weaving, an amylase can be used to remove the size coating before further processing the fabric to ensure a homogeneous and wash-proof result.

[0227] An amylase can be used alone or with other desizing chemical reagents and/or desizing enzymes to desize

fabrics, including cotton-containing fabrics, as detergent additives, e.g., in aqueous compositions. An amylase also can be used in compositions and methods for producing a stonewashed look on indigo-dyed denim fabric and garments. For the manufacture of clothes, the fabric can be cut and sewn into clothes or garments, which are afterwards finished. In particular, for the manufacture of denim jeans, different enzymatic finishing methods have been developed. The finishing of denim garment normally is initiated with an enzymatic desizing step, during which garments are subjected to the action of amylolytic enzymes to provide softness to the fabric and make the cotton more accessible to the subsequent enzymatic finishing steps. An amylase can be used in methods of finishing denim garments (e.g., a "bio-stoning process"), enzymatic desizing and providing softness to fabrics, and/or finishing process.

7. Cleaning Compositions

[0228] An aspect of the present compositions and methods is a cleaning composition that includes a variant α -amylase as defined in the appended claims. as a component. An amylase polypeptide can be used as a component in detergent compositions for hand washing, laundry washing, dishwashing, and other hard-surface cleaning.

7.1. Overview

[0229] Preferably, an amylase is incorporated into detergents at or near a concentration conventionally used for amylase in detergents. For example, an amylase polypeptide may be added in amount corresponding to 0.00001 - 1 mg (calculated as pure enzyme protein) of amylase per liter of wash/dishwash liquor. Exemplary formulations are provided herein, as exemplified by the following:

[0230] An amylase polypeptide may be a component of a detergent composition, as the only enzyme or with other enzymes including other amylolytic enzymes. As such, it may be included in the detergent composition in the form of a non-dusting granulate, a stabilized liquid, or a protected enzyme. Non-dusting granulates may be produced, e.g., as disclosed in U.S. Patent Nos. 4,106,991 and 4,661,452 and may optionally be coated by methods known in the art. Examples of waxy coating materials are poly(ethylene oxide) products (polyethyleneglycol, PEG) with mean molar weights of 1,000 to 20,000; ethoxylated nonylphenols having from 16 to 50 ethylene oxide units; ethoxylated fatty alcohols in which the alcohol contains from 12 to 20 carbon atoms and in which there are 15 to 80 ethylene oxide units; fatty alcohols; fatty acids; and mono- and di- and triglycerides of fatty acids. Examples of film-forming coating materials suitable for application by fluid bed techniques are given in, for example, GB 1483591. Liquid enzyme preparations may, for instance, be stabilized by adding a polyol such as propylene glycol, a sugar or sugar alcohol, lactic acid or boric acid according to established methods. Other enzyme stabilizers are known in the art. Protected enzymes may be prepared according to the method disclosed in for example EP 238 216. Polyols have long been recognized as stabilizers of proteins, as well as improving protein solubility.

[0231] The detergent composition may be in any useful form, e.g., as powders, granules, pastes, or liquid. A liquid detergent may be aqueous, typically containing up to about 70% of water and 0% to about 30% of organic solvent. It may also be in the form of a compact gel type containing only about 30% water.

[0232] The detergent composition comprises one or more surfactants, each of which may be anionic, nonionic, cationic, or zwitterionic. The detergent will usually contain 0% to about 50% of anionic surfactant, such as linear alkylbenzenesulfonate (LAS); α -olefinsulfonate (AOS); alkyl sulfate (fatty alcohol sulfate) (AS); alcohol ethoxysulfate (AEOS or AES); secondary alkanesulfonates (SAS); α -sulfo fatty acid methyl esters; alkyl- or alkenylsuccinic acid; or soap. The composition may also contain 0% to about 40% of nonionic surfactant such as alcohol ethoxylate (AEO or AE), carboxylated alcohol ethoxylates, nonylphenol ethoxylate, alkylpolyglycoside, alkylidimethylamineoxide, ethoxylated fatty acid monoethanolamide, fatty acid monoethanolamide, or polyhydroxy alkyl fatty acid amide (as described for example in WO 92/06154).

[0233] The detergent composition may additionally comprise one or more other enzymes, such as proteases, another amylolytic enzyme, cutinase, lipase, cellulase, pectate lyase, perhydrolase, xylanase, peroxidase, and/or laccase in any combination.

[0234] The detergent may contain about 1% to about 65% of a detergent builder or complexing agent such as zeolite, diphosphate, triphosphate, phosphonate, citrate, nitrilotriacetic acid (NTA), ethylenediaminetetraacetic acid (EDTA), diethylenetriaminepentaacetic acid (DTMPA), alkyl- or alkenylsuccinic acid, soluble silicates or layered silicates (e.g., SKS-6 from Hoechst). The detergent may also be unbuilt, i.e. essentially free of detergent builder. The enzymes can be used in any composition compatible with the stability of the enzyme. Enzymes generally can be protected against deleterious components by known forms of encapsulation, for example, by granulation or sequestration in hydro gels. Enzymes, and specifically amylases, either with or without starch binding domains, can be used in a variety of compositions including laundry and dishwashing applications, surface cleaners, as well as in compositions for ethanol production from starch or biomass.

[0235] The detergent may comprise one or more polymers. Examples include carboxymethylcellulose (CMC), poly(vi-

nylpyrrolidone) (PVP), polyethyleneglycol (PEG), poly(vinyl alcohol) (PVA), polycarboxylates such as polyacrylates, maleic/acrylic acid copolymers and lauryl methacrylate/acrylic acid copolymers.

[0236] The detergent may contain a bleaching system, which may comprise a H₂O₂ source such as perborate or percarbonate, which may be combined with a peracid-forming bleach activator such as tetraacetythylenediamine (TAED) or nonanoyloxybenzenesulfonate (NOBS). Alternatively, the bleaching system may comprise peroxyacids (e.g., the amide, imide, or sulfone type peroxyacids). The bleaching system can also be an enzymatic bleaching system, for example, perhydrolase, such as that described in International PCT Application WO 2005/056783.

[0237] The enzymes of the detergent composition may be stabilized using conventional stabilizing agents, e.g., a polyol such as propylene glycol or glycerol; a sugar or sugar alcohol; lactic acid; boric acid or a boric acid derivative such as, e.g., an aromatic borate ester; and the composition may be formulated as described in, e.g., WO 92/19709 and WO 92/19708.

[0238] The detergent may also contain other conventional detergent ingredients such as e.g., fabric conditioners including clays, foam boosters, suds suppressors, anti-corrosion agents, soil-suspending agents, anti-soil redeposition agents, dyes, bactericides, tarnish inhibitors, optical brighteners, or perfumes.

[0239] The pH (measured in aqueous solution at use concentration) is usually neutral or alkaline, e.g., pH about 7.0 to about 11.0.

[0240] Particular forms of detergent compositions for inclusion of the present α -amylase are described below.

7.2. Heavy Duty Liquid (HDL) laundry detergent composition

[0241] Exemplary HDL laundry detergent compositions includes a deterative surfactant (10%-40% wt/wt), including an anionic deterative surfactant (selected from a group of linear or branched or random chain, substituted or unsubstituted alkyl sulphates, alkyl sulphonates, alkyl alkoxyated sulphate, alkyl phosphates, alkyl phosphonates, alkyl carboxylates, and/or mixtures thereof), and optionally non-ionic surfactant (selected from a group of linear or branched or random chain, substituted or unsubstituted alkyl alkoxyated alcohol, for example a C₈-C₁₈ alkyl ethoxyated alcohol and/or C₆-C₁₂ alkyl phenol alkoxyates), wherein the weight ratio of anionic deterative surfactant (with a hydrophilic index (Hlc) of from 6.0 to 9) to non-ionic deterative surfactant is greater than 1: 1. Suitable deterative surfactants also include cationic deterative surfactants (selected from a group of alkyl pyridinium compounds, alkyl quarternary ammonium compounds, alkyl quarternary phosphonium compounds, alkyl ternary sulphonium compounds, and/or mixtures thereof); zwitterionic and/or amphoteric deterative surfactants (selected from a group of alkanolamine sulfo-betaines); ampholytic surfactants; semi-polar non-ionic surfactants and mixtures thereof.

[0242] The composition may optionally include, a surfactancy boosting polymer consisting of amphiphilic alkoxyated grease cleaning polymers (selected from a group of alkoxyated polymers having branched hydrophilic and hydrophobic properties, such as alkoxyated polyalkylenimines in the range of 0.05wt%- 10wt%) and/or random graft polymers (typically comprising of hydrophilic backbone comprising monomers selected from the group consisting of: unsaturated C₁-C₆ carboxylic acids, ethers, alcohols, aldehydes, ketones, esters, sugar units, alkoxy units, maleic anhydride, saturated polyalcohols such as glycerol, and mixtures thereof; and hydrophobic side chain(s) selected from the group consisting of: C₄-C₂₅ alkyl group, polypropylene, polybutylene, vinyl ester of a saturated C₁-C₆ mono-carboxylic acid, C₁-C₆ alkyl ester of acrylic or methacrylic acid, and mixtures thereof).

[0243] The composition may include additional polymers such as soil release polymers (include anionically end-capped polyesters, for example SRP1, polymers comprising at least one monomer unit selected from saccharide, dicarboxylic acid, polyol and combinations thereof, in random or block configuration, ethylene terephthalate-based polymers and co-polymers thereof in random or block configuration, for example Repel-o-tex SF, SF-2 and SRP6, Texcare SRA100, SRA300, SRN100, SRN170, SRN240, SRN300 and SRN325, Marloquest SL), antiredeposition polymers (0.1 wt% to 10wt%, include carboxylate polymers, such as polymers comprising at least one monomer selected from acrylic acid, maleic acid (or maleic anhydride), fumaric acid, itaconic acid, aconitic acid, mesaconic acid, citraconic acid, methylenemalononic acid, and any mixture thereof, vinylpyrrolidone homopolymer, and/or polyethylene glycol, molecular weight in the range of from 500 to 100,000 Da); cellulosic polymer (including those selected from alkyl cellulose, alkyl alkoxyalkyl cellulose, carboxyalkyl cellulose, alkyl carboxyalkyl cellulose examples of which include carboxymethyl cellulose, methyl cellulose, methyl hydroxyethyl cellulose, methyl carboxymethyl cellulose, and mixures thereof) and polymeric carboxylate (such as maleate/acrylate random copolymer or polyacrylate homopolymer).

[0244] The composition may further include saturated or unsaturated fatty acid, preferably saturated or unsaturated C₁₂-C₂₄ fatty acid (0 wt% to 10 wt%); deposition aids (examples for which include polysaccharides, preferably cellulosic polymers, poly diallyl dimethyl ammonium halides (DADMAC), and co-polymers of DAD MAC with vinyl pyrrolidone, acrylamides, imidazoles, imidazolinium halides, and mixtures thereof, in random or block configuration, cationic guar gum, cationic cellulose such as cationic hydroxyethyl cellulose, cationic starch, cationic polyacrylamides, and mixtures thereof).

[0245] The composition may further include dye transfer inhibiting agents, examples of which include manganese

phthalocyanine, peroxidases, polyvinylpyrrolidone polymers, polyamine N-oxide polymers, copolymers of N-vinylpyrrolidone and N-vinylimidazole, polyvinylloxazolidones and polyvinylimidazoles and/or mixtures thereof; chelating agents, examples of which include ethylene-diamine-tetraacetic acid (EDTA), diethylene triamine penta methylene phosphonic acid (DTPMP), hydroxy-ethane diphosphonic acid (HEDP), ethylenediamine N,N'-disuccinic acid (EDDS), methyl glycine diacetic acid (MGDA), diethylene triamine penta acetic acid (DTPA), propylene diamine tetracetic acid (PDT A), 2-hydroxypyridine-N-oxide (HPNO), or methyl glycine diacetic acid (MGDA), glutamic acid N,N-diacetic acid (N,N-dicarboxymethyl glutamic acid tetrasodium salt (GLDA), nitrilotriacetic acid (NTA), 4,5-dihydroxy-m-benzenedisulfonic acid, citric acid and any salts thereof, N-hydroxyethylethylenediaminetri-acetic acid (HEDTA), triethylenetetraaminehexaacetic acid (TTHA), N-hydroxyethyliminodiacetic acid (HEIDA), dihydroxyethylglycine (DHEG), ethylenediaminetetrapropionic acid (EDTP), and derivatives thereof.

[0246] The composition preferably included enzymes (generally about 0.01 wt% active enzyme to 0.03wt% active enzyme) selected from proteases, amylases, lipases, cellulases, choline oxidases, peroxidases/oxidases, pectate lyases, mannanases, cutinases, laccases, phospholipases, lysophospholipases, acyltransferases, perhydrolases, arylesterases, and any mixture thereof. The composition may include an enzyme stabilizer (examples of which include polyols such as propylene glycol or glycerol, sugar or sugar alcohol, lactic acid, reversible protease inhibitor, boric acid, or a boric acid derivative, e.g., an aromatic borate ester, or a phenyl boronic acid derivative such as 4-formylphenyl boronic acid).

[0247] The composition optionally include silicone or fatty-acid based suds suppressors; heuing dyes, calcium and magnesium cations, visual signaling ingredients, anti-foam (0.001wt% to about 4.0wt%), and/or structurant/thickener (0.01wt% to 5wt%, selected from the group consisting of diglycerides and triglycerides, ethylene glycol distearate, microcrystalline cellulose, cellulose based materials, microfiber cellulose, biopolymers, xanthan gum, gellan gum, and mixtures thereof).

[0248] The composition can be any liquid form, for example a liquid or gel form, or any combination thereof. The composition may be in any unit dose form, for example a pouch.

7.3. Heavy Duty Dry/Solid (HDD) laundry detergent composition

[0249] Exemplary HDD laundry detergent compositions includes a deterative surfactant, including anionic deterative surfactants (e.g., linear or branched or random chain, substituted or unsubstituted alkyl sulphates, alkyl sulphonates, alkyl alkoxyated sulphate, alkyl phosphates, alkyl phosphonates, alkyl carboxylates and/or mixtures thereof), non-ionic deterative surfactant (e.g., linear or branched or random chain, substituted or unsubstituted C₈-C₁₈ alkyl ethoxylates, and/or C₆-C₁₂ alkyl phenol alkoxyates), cationic deterative surfactants (e.g., alkyl pyridinium compounds, alkyl quaternary ammonium compounds, alkyl quaternary phosphonium compounds, alkyl ternary sulphonium compounds, and mixtures thereof), zwitterionic and/or amphoteric deterative surfactants (e.g., alkanolamine sulfo-betaines), ampholytic surfactants, semi-polar non-ionic surfactants, and mixtures thereof; builders including phosphate free builders (for example zeolite builders examples which include zeolite A, zeolite X, zeolite P and zeolite MAP in the range of 0wt% to less than 10wt%), phosphate builders (for example sodium tri-polyphosphate in the range of 0wt% to less than 10wt%), citric acid, citrate salts and nitrilotriacetic acid, silicate salt (e.g., sodium or potassium silicate or sodium meta-silicate in the range of 0wt% to less than 10wt%, or layered silicate (SKS-6)); carbonate salt (e.g., sodium carbonate and/or sodium bicarbonate in the range of 0wt% to less than 80wt%); and bleaching agents including photobleaches (e.g., sulfonated zinc phthalocyanines, sulfonated aluminum phthalocyanines, xanthenes dyes, and mixtures thereof) hydrophobic or hydrophilic bleach activators (e.g., dodecanoyl oxybenzene sulfonate, decanoyl oxybenzene sulfonate, decanoyl oxybenzoic acid or salts thereof, 3,5,5-trimethy hexanoyl oxybenzene sulfonate, tetraacetyl ethylene diamine-TAED, nonanoyloxybenzene sulfonate-NOBS, nitrile quats, and mixtures thereof), sources of hydrogen peroxide (e.g., inorganic perhydrate salts examples of which include mono or tetra hydrate sodium salt of perborate, percarbonate, persulfate, perphosphate, or persilicate), preformed hydrophilic and/or hydrophobic peracids (e.g., percarboxylic acids and salts, percarbonic acids and salts, perimidic acids and salts, peroxymonosulfuric acids and salts, and mixtures thereof), and/or bleach catalysts (e.g., imine bleach boosters (examples of which include iminium cations and polyions), iminium zwitterions, modified amines, modified amine oxides, N-sulphonyl imines, N-phosphonyl imines, N-acyl imines, thiadiazole dioxides, perfluoroimines, cyclic sugar ketones, and mixtures thereof, and metal-containing bleach catalysts (e.g., copper, iron, titanium, ruthenium, tungsten, molybdenum, or manganese cations along with an auxiliary metal cations such as zinc or aluminum and a sequester such as ethylenediaminetetraacetic acid, ethylenediaminetetra(methylenephosphonic acid), and watersoluble salts thereof).

[0250] The composition preferably includes enzymes, e.g., proteases, amylases, lipases, cellulases, choline oxidases, peroxidases/oxidases, pectate lyases, mannanases, cutinases, laccases, phospholipases, lysophospholipases, acyltransferase, perhydrolase, arylesterase, and any mixture thereof.

[0251] The composition may optionally include additional detergent ingredients including perfume microcapsules, starch encapsulated perfume accord, hueing agents, additional polymers, including fabric integrity and cationic polymers, dye-lock ingredients, fabric-softening agents, brighteners (for example C.I. Fluorescent brighteners), flocculating agents,

chelating agents, alkoxyated polyamines, fabric deposition aids, and/or cyclodextrin.

7.4. Automatic dishwashing (ADW) detergent composition

5 **[0252]** Exemplary ADW detergent composition includes non-ionic surfactants, including ethoxylated non-ionic surfactants, alcohol alkoxyated surfactants, epoxy-capped poly(oxyalkylated) alcohols, or amine oxide surfactants present in amounts from 0 to 10% by weight; builders in the range of 5-60% including phosphate builders (e.g., mono-phosphates, di-phosphates, tri-polyphosphates, other oligomeric-polyphosphates, sodium tripolyphosphate-STPP) and phosphate-free builders (e.g., amino acid-based compounds including methylglycine-diacetic acid (MGDA) and salts and derivatives thereof, glutamic-N,N-diacetic acid (GLDA) and salts and derivatives thereof, iminodisuccinic acid (IDS) and salts and derivatives thereof, carboxy methyl inulin and salts and derivatives thereof, nitrilotriacetic acid (NTA), diethylene triamine penta acetic acid (DTPA), B-alaninediacetic acid (B-ADA) and their salts, homopolymers and copolymers of poly-carboxylic acids and their partially or completely neutralized salts, monomeric polycarboxylic acids and hydroxycarboxylic acids and their salts in the range of 0.5% to 50% by weight; sulfonated/carboxylated polymers in the range of about 0.1
10 % to about 50% by weight to provide dimensional stability; drying aids in the range of about 0.1% to about 10% by weight (e.g., polyesters, especially anionic polyesters, optionally together with further monomers with 3 to 6 functionalities - typically acid, alcohol or ester functionalities which are conducive to polycondensation, polycarbonate-, polyurethane- and/or polyurea-polyorganosiloxane compounds or precursor compounds, thereof, particularly of the reactive cyclic carbonate and urea type); silicates in the range from about 1% to about 20% by weight (including sodium or potassium silicates for example sodium disilicate, sodium meta-silicate and crystalline phyllosilicates); inorganic bleach (e.g., perhydrate salts such as perborate, percarbonate, perphosphate, persulfate and persilicate salts) and organic bleach (e.g., organic peroxyacids, including diacyl and tetraacylperoxides, especially diperoxydodecanedioc acid, diperoxytetradecanedioc acid, and diperoxyhexadecanedioc acid); bleach activators (i.e., organic peracid precursors in the range from about 0.1% to about 10% by weight); bleach catalysts (e.g., manganese triazacyclononane and related complexes, Co,
25 Cu, Mn, and Fe bispyridylamine and related complexes, and pentamine acetate cobalt(III) and related complexes); metal care agents in the range from about 0.1% to 5% by weight (e.g., benzotriazoles, metal salts and complexes, and/or silicates); enzymes in the range from about 0.01 to 5.0 mg of active enzyme per gram of automatic dishwashing detergent composition (e.g., proteases, amylases, lipases, cellulases, choline oxidases, peroxidases/oxidases, pectate lyases, mannanases, cutinases, laccases, phospholipases, lysophospholipases, acyltransferase, perhydrolase, arylesterase, and mixtures thereof); and enzyme stabilizer components (e.g., oligosaccharides, polysaccharides, and inorganic divalent metal salts).

7.5. Additional detergent compositions

35 **[0253]** Additional exemplary detergent formulations to which the present amylase can be added are described, below, in the numbered paragraphs.

1) A detergent composition formulated as a granulate having a bulk density of at least 600 g/L comprising linear alkylbenzenesulfonate (calculated as acid) about 7% to about 12%; alcohol ethoxysulfate (e.g., C₁₂₋₁₈ alcohol, 1-2
40 ethylene oxide (EO)) or alkyl sulfate (e.g., C₁₆₋₁₈) about 1% to about 4%; alcohol ethoxylate (e.g., C₁₄₋₁₅ alcohol, 7 EO) about 5% to about 9%; sodium carbonate (e.g., Na₂CO₃) about 14% to about 20%; soluble silicate (e.g., Na₂O, 2SiO₂) about 2 to about 6%; zeolite (e.g., NaAlSiO₄) about 15% to about 22%; sodium sulfate (e.g., Na₂SO₄) 0% to about 6%; sodium citrate/citric acid (e.g., C₆H₅Na₃O₇/C₆H₈O₇) about 0% to about 15%; sodium perborate (e.g., NaBO₃H₂O) about 11% to about 18%; TAED about 2% to about 6%; carboxymethylcellulose (CMC) and 0% to about 2%; polymers (e.g., maleic/acrylic acid, copolymer, PVP, PEG) 0-3%; enzymes (calculated as pure enzyme)
45 0.0001-0.1% protein; and minor ingredients (e.g., suds suppressors, perfumes, optical brightener, photobleach) 0-5%.

2) A detergent composition formulated as a granulate having a bulk density of at least 600 g/L comprising linear alkylbenzenesulfonate (calculated as acid) about 6% to about 11%; alcohol ethoxysulfate (e.g., C₁₂₋₁₈ alcohol, 1-2
50 EO) or alkyl sulfate (e.g., C₁₆₋₁₈) about 1% to about 3%; alcohol ethoxylate (e.g., C₁₄₋₁₅ alcohol, 7 EO) about 5% to about 9%; sodium carbonate (e.g., Na₂CO₃) about 15% to about 21%; soluble silicate (e.g., Na₂O, 2SiO₂) about 1% to about 4%; zeolite (e.g., NaAlSiO₄) about 24% to about 34%; sodium sulfate (e.g., Na₂SO₄) about 4% to about 10%; sodium citrate/citric acid (e.g., C₆H₅Na₃O₇/C₆H₈O₇) 0% to about 15%; carboxymethylcellulose (CMC) 0% to about 2%; polymers (e.g., maleic/acrylic acid copolymer, PVP, PEG) 1-6%; enzymes (calculated as pure enzyme
55 protein) 0.0001-0.1%; minor ingredients (e.g., suds suppressors, perfume) 0-5%.

3) A detergent composition formulated as a granulate having a bulk density of at least 600 g/L comprising linear

alkylbenzenesulfonate (calculated as acid) about 5% to about 9%; alcohol ethoxylate (e.g., C₁₂₋₁₅ alcohol, 7 EO) about 7% to about 14%; Soap as fatty acid (e.g., C₁₆₋₂₂ fatty acid) about 1 to about 3%; sodium carbonate (as Na₂CO₃) about 10% to about 17%; soluble silicate (e.g., Na₂O, 2SiO₂) about 3% to about 9%; zeolite (as NaAlSiO₄) about 23% to about 33%; sodium sulfate (e.g., Na₂SO₄) 0% to about 4%; sodium perborate (e.g., NaBO₃H₂O) about 8% to about 16%; TAED about 2% to about 8%; phosphonate (e.g., EDTMPA) 0% to about 1%; carboxymethylcellulose (CMC) 0% to about 2%; polymers (e.g., maleic/acrylic acid copolymer, PVP, PEG) 0-3%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; minor ingredients (e.g., suds suppressors, perfume, optical brightener) 0-5%.

4) A detergent composition formulated as a granulate having a bulk density of at least 600 g/L comprising linear alkylbenzenesulfonate (calculated as acid) about 8% to about 12%; alcohol ethoxylate (e.g., C₁₂₋₁₅ alcohol, 7 EO) about 10% to about 25%; sodium carbonate (as Na₂CO₃) about 14% to about 22%; soluble silicate (e.g., Na₂O, 2SiO₂) about 1% to about 5%; zeolite (e.g., NaAlSiO₄) about 25% to about 35%; sodium sulfate (e.g., Na₂SO₄) 0% to about 10%; carboxymethylcellulose (CMC) 0% to about 2%; polymers (e.g., maleic/acrylic acid copolymer, PVP, PEG) 1-3%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; and minor ingredients (e.g., suds suppressors, perfume) 0-5%.

5) An aqueous liquid detergent composition comprising linear alkylbenzenesulfonate (calculated as acid) about 15% to about 21%; alcohol ethoxylate (e.g., C₁₂₋₁₅ alcohol, 7 EO or C₁₂₋₁₅ alcohol, 5 EO) about 12% to about 18%; soap as fatty acid (e.g., oleic acid) about 3% to about 13%; alkenylsuccinic acid (C₁₂₋₁₄) 0% to about 13%; aminoethanol about 8% to about 18%; citric acid about 2% to about 8%; phosphonate 0% to about 3%; polymers (e.g., PVP, PEG) 0% to about 3%; borate (e.g., B₄O₇) 0% to about 2%; ethanol 0% to about 3%; propylene glycol about 8% to about 14%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; and minor ingredients (e.g., dispersants, suds suppressors, perfume, optical brightener) 0-5%.

6) An aqueous structured liquid detergent composition comprising linear alkylbenzenesulfonate (calculated as acid) about 15% to about 21%; alcohol ethoxylate (e.g., C₁₂₋₁₅ alcohol, 7 EO, or C₁₂₋₁₅ alcohol, 5 EO) 3-9%; soap as fatty acid (e.g., oleic acid) about 3% to about 10%; zeolite (as NaAlSiO₄) about 14% to about 22%; potassium citrate about 9% to about 18%; borate (e.g., B₄O₇) 0% to about 2%; carboxymethylcellulose (CMC) 0% to about 2%; polymers (e.g., PEG, PVP) 0% to about 3%; anchoring polymers such as, e.g., lauryl methacrylate/acrylic acid copolymer; molar ratio 25:1, MW 3800) 0% to about 3%; glycerol 0% to about 5%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; and minor ingredients (e.g., dispersants, suds suppressors, perfume, optical brighteners) 0-5%.

7) A detergent composition formulated as a granulate having a bulk density of at least 600 g/L comprising fatty alcohol sulfate about 5% to about 10%; ethoxylated fatty acid monoethanolamide about 3% to about 9%; soap as fatty acid 0-3%; sodium carbonate (e.g., Na₂CO₃) about 5% to about 10%; Soluble silicate (e.g., Na₂O, 2SiO₂) about 1% to about 4%; zeolite (e.g., NaAlSiO₄) about 20% to about 40%; Sodium sulfate (e.g., Na₂SO₄) about 2% to about 8%; sodium perborate (e.g., NaBO₃H₂O) about 12% to about 18%; TAED about 2% to about 7%; polymers (e.g., maleic/acrylic acid copolymer, PEG) about 1% to about 5%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; and minor ingredients (e.g., optical brightener, suds suppressors, perfume) 0-5%.

8) A detergent composition formulated as a granulate comprising linear alkylbenzenesulfonate (calculated as acid) about 8% to about 14%; ethoxylated fatty acid monoethanolamide about 5% to about 11%; soap as fatty acid 0% to about 3%; sodium carbonate (e.g., Na₂CO₃) about 4% to about 10%; soluble silicate (Na₂O, 2SiO₂) about 1% to about 4%; zeolite (e.g., NaAlSiO₄) about 30% to about 50%; sodium sulfate (e.g., Na₂SO₄) about 3% to about 11%; sodium citrate (e.g., C₆H₅Na₃O₇) about 5% to about 12%; polymers (e.g., PVP, maleic/acrylic acid copolymer, PEG) about 1% to about 5%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; and minor ingredients (e.g., suds suppressors, perfume) 0-5%.

9) A detergent composition formulated as a granulate comprising linear alkylbenzenesulfonate (calculated as acid) about 6% to about 12%; nonionic surfactant about 1% to about 4%; soap as fatty acid about 2% to about 6%; sodium carbonate (e.g., Na₂CO₃) about 14% to about 22%; zeolite (e.g., NaAlSiO₄) about 18% to about 32%; sodium sulfate (e.g., Na₂SO₄) about 5% to about 20%; sodium citrate (e.g., C₆H₅Na₃O₇) about 3% to about 8%; sodium perborate (e.g., NaBO₃H₂O) about 4% to about 9%; bleach activator (e.g., NOBS or TAED) about 1% to about 5%; carboxymethylcellulose (CMC) 0% to about 2%; polymers (e.g., polycarboxylate or PEG) about 1% to about 5%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; and minor ingredients (e.g., optical brightener, perfume) 0-5%.

10) An aqueous liquid detergent composition comprising linear alkylbenzenesulfonate (calculated as acid) about 15% to about 23%; alcohol ethoxysulfate (e.g., C₁₂₋₁₅ alcohol, 2-3 EO) about 8% to about 15%; alcohol ethoxylate (e.g., C₁₂₋₁₅ alcohol, 7 EO, or C₁₂₋₁₅ alcohol, 5 EO) about 3% to about 9%; soap as fatty acid (e.g., lauric acid) 0% to about 3%; aminoethanol about 1% to about 5%; sodium citrate about 5% to about 10%; hydrotrope (e.g., sodium toluenesulfonate) about 2% to about 6%; borate (e.g., B₄O₇) 0% to about 2%; carboxymethylcellulose 0% to about 1%; ethanol about 1% to about 3%; propylene glycol about 2% to about 5%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; and minor ingredients (e.g., polymers, dispersants, perfume, optical brighteners) 0-5%.

11) An aqueous liquid detergent composition comprising linear alkylbenzenesulfonate (calculated as acid) about 20% to about 32%; alcohol ethoxylate (e.g., C₁₂₋₁₅ alcohol, 7 EO, or C₁₂₋₁₅ alcohol, 5 EO) 6-12%; aminoethanol about 2% to about 6%; citric acid about 8% to about 14%; borate (e.g., B₄O₇) about 1% to about 3%; polymer (e.g., maleic/acrylic acid copolymer, anchoring polymer such as, e.g., lauryl methacrylate/acrylic acid copolymer) 0% to about 3%; glycerol about 3% to about 8%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; and minor ingredients (e.g., hydrotropes, dispersants, perfume, optical brighteners) 0-5%.

12) A detergent composition formulated as a granulate having a bulk density of at least 600 g/L comprising anionic surfactant (linear alkylbenzenesulfonate, alkyl sulfate, α -olefinsulfonate, α -sulfo fatty acid methyl esters, alkanesulfonates, soap) about 25% to about 40%; nonionic surfactant (e.g., alcohol ethoxylate) about 1% to about 10%; sodium carbonate (e.g., Na₂CO₃) about 8% to about 25%; soluble silicates (e.g., Na₂O, 2SiO₂) about 5% to about 15%; sodium sulfate (e.g., Na₂SO₄) 0% to about 5%; zeolite (NaAlSiO₄) about 15% to about 28%; sodium perborate (e.g., NaBO₃·4H₂O) 0% to about 20%; bleach activator (TAED or NOBS) about 0% to about 5%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; minor ingredients (e.g., perfume, optical brighteners) 0-3%.

13) Detergent compositions as described in compositions 1)-12) *supra*, wherein all or part of the linear alkylbenzenesulfonate is replaced by (C₁₂-C₁₈) alkyl sulfate.

14) A detergent composition formulated as a granulate having a bulk density of at least 600 g/L comprising (C₁₂-C₁₈) alkyl sulfate about 9% to about 15%; alcohol ethoxylate about 3% to about 6%; polyhydroxy alkyl fatty acid amide about 1% to about 5%; zeolite (e.g., NaAlSiO₄) about 10% to about 20%; layered disilicate (e.g., SK56 from Hoechst) about 10% to about 20%; sodium carbonate (e.g., Na₂CO₃) about 3% to about 12%; soluble silicate (e.g., Na₂O, 2SiO₂) 0% to about 6%; sodium citrate about 4% to about 8%; sodium percarbonate about 13% to about 22%; TAED about 3% to about 8%; polymers (e.g., polycarboxylates and PVP) 0% to about 5%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; and minor ingredients (e.g., optical brightener, photobleach, perfume, suds suppressors) 0-5%.

15) A detergent composition formulated as a granulate having a bulk density of at least 600 g/L comprising (C₁₂-C₁₈) alkyl sulfate about 4% to about 8%; alcohol ethoxylate about 11% to about 15%; soap about 1% to about 4%; zeolite MAP or zeolite A about 35% to about 45%; sodium carbonate (as Na₂CO₃) about 2% to about 8%; soluble silicate (e.g., Na₂O, 2SiO₂) 0% to about 4%; sodium percarbonate about 13% to about 22%; TAED 1-8%; carboxymethylcellulose (CMC) 0% to about 3%; polymers (e.g., polycarboxylates and PVP) 0% to about 3%; enzymes (calculated as pure enzyme protein) 0.0001-0.1%; and minor ingredients (e.g., optical brightener, phosphonate, perfume) 0-3%.

16) Detergent formulations as described in 1)-15) *supra*, which contain a stabilized or encapsulated peracid, either as an additional component or as a substitute for already specified bleach systems.

17) Detergent compositions as described *supra* in 1), 3), 7), 9), and 12), wherein perborate is replaced by percarbonate.

18) Detergent compositions as described *supra* in 1), 3), 7), 9), 12), 14), and 15), which additionally contain a manganese catalyst. The manganese catalyst for example is one of the compounds described in "Efficient manganese catalysts for low-temperature bleaching," Nature 369: 637-639 (1994).

19) Detergent composition formulated as a non-aqueous detergent liquid comprising a liquid nonionic surfactant such as, e.g., linear alkoxyated primary alcohol, a builder system (e.g., phosphate), an enzyme(s), and alkali. The detergent may also comprise anionic surfactant and/or a bleach system.

[0254] As above, the present amylase polypeptide may be incorporated at a concentration conventionally employed in detergents. It is at present contemplated that, in the detergent composition, the enzyme may be added in an amount

corresponding to 0.00001-1.0 mg (calculated as pure enzyme protein) of amylase polypeptide per liter of wash liquor.

[0255] The detergent composition may also contain other conventional detergent ingredients, e.g., deflocculant material, filler material, foam depressors, anti-corrosion agents, soil-suspending agents, sequestering agents, anti-soil redeposition agents, dehydrating agents, dyes, bactericides, fluorescers, thickeners, and perfumes.

[0256] The detergent composition may be formulated as a hand (manual) or machine (automatic) laundry detergent composition, including a laundry additive composition suitable for pre-treatment of stained fabrics and a rinse added fabric softener composition, or be formulated as a detergent composition for use in general household hard surface cleaning operations, or be formulated for manual or automatic dishwashing operations.

[0257] Any of the cleaning compositions described, herein, may include any number of additional enzymes. In general the enzyme(s) should be compatible with the selected detergent, (e.g., with respect to pH-optimum, compatibility with other enzymatic and non-enzymatic ingredients, and the like), and the enzyme(s) should be present in effective amounts. The following enzymes are provided as examples.

[0258] *Proteases*: Suitable proteases include those of animal, vegetable or microbial origin. Chemically modified or protein engineered mutants are included, as well as naturally processed proteins. The protease may be a serine protease or a metalloprotease, an alkaline microbial protease, a trypsin-like protease, or a chymotrypsin-like protease. Examples of alkaline proteases are subtilisins, especially those derived from *Bacillus*, e.g., subtilisin Novo, subtilisin Carlsberg, subtilisin 309, subtilisin 147, and subtilisin 168 (see, e.g., WO 89/06279). Examples of trypsin-like proteases are trypsin (e.g., of porcine or bovine origin), and *Fusarium* proteases (see, e.g., WO 89/06270 and WO 94/25583). Examples of useful proteases also include but are not limited to the variants described in WO 92/19729, WO 98/20115, WO 98/20116, and WO 98/34946. Commercially available protease enzymes include but are not limited to: ALCALASE®, SAVINASE®, PRIMASE™, DURALASE™, ESPERASE®, KANNASE™, and BLAZE™ (Novo Nordisk A/S and Novozymes A/S); MAX-ATASE®, MAXACAL™, MAXAPEM™, PROPERASE®, PURAFECT®, PURAFECT OXP™, FN2™, and FN3™ (Danisco US Inc.). Other exemplary proteases include NprE from *Bacillus amyloliquifaciens* and ASP from *Cellulomonas* sp. strain 69B4.

[0259] *Lipases*: Suitable lipases include those of bacterial or fungal origin. Chemically modified, proteolytically modified, or protein engineered mutants are included. Examples of useful lipases include but are not limited to lipases from *Humicola* (synonym *Thermomyces*), e.g., from *H. lanuginosa* (*T. lanuginosus*) (see e.g., EP 258068 and EP 305216), from *H. insolens* (see e.g., WO 96/13580); a *Pseudomonas* lipase (e.g., from *P. alcaligenes* or *P. pseudoalcaligenes*; see, e.g., EP 218 272), *P. cepacia* (see e.g., EP 331 376), *P. stutzeri* (see e.g., GB 1,372,034), *P. fluorescens*, *Pseudomonas* sp. strain SD 705 (see e.g., WO 95/06720 and WO 96/27002), *P. wisconsinensis* (see e.g., WO 96/12012); a *Bacillus* lipase (e.g., from *B. subtilis*; see e.g., Dartois et al. Biochemica et Biophysica Acta, 1131: 253-360 (1993)), *B. stearothermophilus* (see e.g., JP 64/744992), or *B. pumilus* (see e.g., WO 91/16422). Additional lipase variants contemplated for use in the formulations include those described for example in: WO 92/05249, WO 94/01541, WO 95/35381, WO 96/00292, WO 95/30744, WO 94/25578, WO 95/14783, WO 95/22615, WO 97/04079, WO 97/07202, EP 407225, and EP 260105. Some commercially available lipase enzymes include LIPOLASE® and LIPOLASE ULTRA™ (Novo Nordisk A/S and Novozymes A/S).

[0260] *Polyesterases*: Suitable polyesterases can be included in the composition, such as those described in, for example, WO 01/34899, WO 01/14629, and US6933140.

[0261] *Amylases*: The compositions can be combined with other amylases, such as non-production enhanced amylase. These can include commercially available amylases, such as but not limited to STAINZYME®, NATALASE®, DURAMYL®, TERMAMYL®, FUNGAMYL® and BAN™ (Novo Nordisk A/S and Novozymes A/S); RAPIDASE®, POWERASE®, and PURASTAR® (from Danisco US Inc.).

[0262] *Cellulases*: Cellulases can be added to the compositions. Suitable cellulases include those of bacterial or fungal origin. Chemically modified or protein engineered mutants are included. Suitable cellulases include cellulases from the genera *Bacillus*, *Pseudomonas*, *Humicola*, *Fusarium*, *Thielavia*, *Acremonium*, e.g., the fungal cellulases produced from *Humicola insolens*, *Myceliophthora thermophila* and *Fusarium oxysporum* disclosed for example in U.S. Patent Nos. 4,435,307; 5,648,263; 5,691,178; 5,776,757; and WO 89/09259. Exemplary cellulases contemplated for use are those having color care benefit for the textile. Examples of such cellulases are cellulases described in for example EP 0495257, EP 0531372, WO 96/11262, WO 96/29397, and WO 98/08940. Other examples are cellulase variants, such as those described in WO 94/07998; WO 98/12307; WO 95/24471; PCT/DK98/00299; EP 531315; U.S. Patent Nos. 5,457,046; 5,686,593; and 5,763,254. Commercially available cellulases include CELLUZYME® and CAREZYME® (Novo Nordisk A/S and Novozymes A/S); CLAZINASE® and PURADAX HA® (Danisco US Inc.); and KAC-500(B)™ (Kao Corporation).

[0263] *Peroxidases/Oxidases*: Suitable peroxidases/oxidases contemplated for use in the compositions include those of plant, bacterial or fungal origin. Chemically modified or protein engineered mutants are included. Examples of useful peroxidases include peroxidases from *Coprinus*, e.g., from *C. cinereus*, and variants thereof as those described in WO 93/24618, WO 95/10602, and WO 98/15257. Commercially available peroxidases include for example GUARDZYME™ (Novo Nordisk A/S and Novozymes A/S).

[0264] The detergent composition can also comprise 2,6-β-D-fructan hydrolase, which is effective for removal/cleaning

of biofilm present on household and/or industrial textile/laundry.

[0265] The detergent enzyme(s) may be included in a detergent composition by adding separate additives containing one or more enzymes, or by adding a combined additive comprising all of these enzymes. A detergent additive, *i.e.* a separate additive or a combined additive, can be formulated *e.g.*, as a granulate, a liquid, a slurry, and the like. Exemplary detergent additive formulations include but are not limited to granulates, in particular non-dusting granulates, liquids, in particular stabilized liquids or slurries.

[0266] Non-dusting granulates may be produced, *e.g.*, as disclosed in U.S. Patent Nos. 4,106,991 and 4,661,452 and may optionally be coated by methods known in the art. Examples of waxy coating materials are poly(ethylene oxide) products (*e.g.*, polyethyleneglycol, PEG) with mean molar weights of 1,000 to 20,000; ethoxylated nonylphenols having from 16 to 50 ethylene oxide units; ethoxylated fatty alcohols in which the alcohol contains from 12 to 20 carbon atoms and in which there are 15 to 80 ethylene oxide units; fatty alcohols; fatty acids; and mono- and di- and triglycerides of fatty acids. Examples of film-forming coating materials suitable for application by fluid bed techniques are given in, for example, GB 1483591. Liquid enzyme preparations may, for instance, be stabilized by adding a polyol such as propylene glycol, a sugar or sugar alcohol, lactic acid or boric acid according to established methods. Protected enzymes may be prepared according to the method disclosed in EP 238,216.

[0267] The detergent composition may be in any convenient form, *e.g.*, a bar, a tablet, a powder, a granule, a paste, or a liquid. A liquid detergent may be aqueous, typically containing up to about 70% water, and 0% to about 30% organic solvent. Compact detergent gels containing about 30% or less water are also contemplated. The detergent composition can optionally comprise one or more surfactants, which may be non-ionic, including semi-polar and/or anionic and/or cationic and/or zwitterionic. The surfactants can be present in a wide range, from about 0.1% to about 60% by weight.

[0268] When included therein the detergent will typically contain from about 1% to about 40% of an anionic surfactant, such as linear alkylbenzenesulfonate, α -olefinsulfonate, alkyl sulfate (fatty alcohol sulfate), alcohol ethoxysulfate, secondary alkanesulfonate, α -sulfo fatty acid methyl ester, alkyl- or alkenylsuccinic acid, or soap.

[0269] When included therein, the detergent will usually contain from about 0.2% to about 40% of a non-ionic surfactant such as alcohol ethoxylate, nonylphenol ethoxylate, alkylpolyglycoside, alkyltrimethylamineoxide, ethoxylated fatty acid monoethanolamide, fatty acid monoethanolamide, polyhydroxy alkyl fatty acid amide, or N-acyl-N-alkyl derivatives of glucosamine ("glucamides").

[0270] The detergent may contain 0% to about 65% of a detergent builder or complexing agent such as zeolite, diphosphate, triphosphate, phosphonate, carbonate, citrate, nitrilotriacetic acid, ethylenediaminetetraacetic acid (EDTA), diethylenetriaminepentaacetic acid, alkyl- or alkenylsuccinic acid, soluble silicates or layered silicates (*e.g.*, SKS-6 from Hoechst).

[0271] The detergent may comprise one or more polymers. Exemplary polymers include carboxymethylcellulose (CMC), poly(vinylpyrrolidone) (PVP), poly(ethylene glycol) (PEG), poly(vinyl alcohol) (PVA), poly(vinylpyridine-N-oxide), poly(vinylimidazole), polycarboxylates *e.g.*, polyacrylates, maleic/acrylic acid copolymers), and lauryl methacrylate/acrylic acid copolymers.

[0272] The enzyme(s) of the detergent composition may be stabilized using conventional stabilizing agents, *e.g.*, as polyol (*e.g.*, propylene glycol or glycerol), a sugar or sugar alcohol, lactic acid, boric acid, or a boric acid derivative (*e.g.*, an aromatic borate ester), or a phenyl boronic acid derivative (*e.g.*, 4-formylphenyl boronic acid). The composition may be formulated as described in WO 92/19709 and WO 92/19708.

[0273] It is contemplated that in the detergent compositions, in particular the enzyme variants, may be added in an amount corresponding to about 0.01 to about 100 mg of enzyme protein per liter of wash liquor (*e.g.*, about 0.05 to about 5.0 mg of enzyme protein per liter of wash liquor or 0.1 to about 1.0 mg of enzyme protein per liter of wash liquor).

[0274] Yet additional exemplary detergent formulations to which the present amylase can be added (or is in some cases identified as a component) are listed in the following Tables:

HDL Detergent Composition	
Ingredient	wt%
Enzyme (s) (Protease + Lipase + Amylase)	3
Linear alkyl benzene sulphonic acid (HLAS)	10
C12-14 alkyl ethoxylated alcohol having an average degree of ethoxylation of 9 (AE9)	2
C12-14 alkyl ethoxylated sulphonic acid having an average degree of ethoxylation of 3 (HAES)	23
C16-17 alkyl mid chain branched alkyl sulphate	4
Amine oxide	1
C12-18 fatty acid	2

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(continued)

HDL Detergent Composition	
Ingredient	wt%
PE20 polymer	3
Polyethylene imine polymer	3
Chelant	1.4
FW A 15 Brightener	0.4
p-glycol (solvent)	8
DEG (solvent)	0.5
Ethanol	3
Monoethanolamine	6
Water	26
NaOH	0.3
Perfume	1
Silicone suds suppressor	0.06
Violet DD dye	0.01
Other dyes	0.03
Hydrogenated castor oil (structurant/thickener)	0.1
Mica	0.2
Calcium formate	0.1
Sodium formate	0.2
Miscellaneous	to 100

HDD Detergent Compositions				
Ingredient	Composition A	Composition B	Composition C	Composition D
Enzyme (Lipase + other enzymes)	0.8 wt%	0.8 wt%	0.8 wt%	0.8 wt%
Linear alkyl benzene sulphonate	9 wt%	9 wt%	12 wt%	8 wt%
Alkyl ethoxylated sulphate having an average degree of ethoxylation of from 0.5 to 3	3 wt%	2 wt%	1 wt%	2 wt%
Cationic deterative surfactant	0.5 wt%	0.5 wt%	0.5 wt%	0.5 wt%
Sodium sulphate	55 wt%	55 wt%	55 wt%	55 wt%
Sodium carbonate	8 wt%	10 wt%	5 wt%	8 wt%
Glycerol carbonate	9 wt%	12 wt%	8 wt%	10 wt%
Oxaziridiniyum-based bleach catalyst	0.005wt%	0.005wt%	0.005wt%	0.005wt%
Sodium silicate	3 wt%	0 wt%	3 wt%	0 wt%
Carboxylate polymer	2wt%	2wt%	2wt%	2wt%
Brightener	0.02 wt%	0.02 wt%	0.02 wt%	0.02 wt%
Cellulosic polymer	0.3 wt%	0.3 wt%	0.3 wt%	0.3 wt%
Misc & Moisture	to 100 wt%	to 100 wt%	to 100 wt%	to 100 wt%

	HDD Detergent Compositions						
	Ingredient	1 (wt%)	2 (wt%)	3 (wt%)	4 (wt%)	5 (wt%)	6 (wt%)
5	Sodium linear alky lbenzenesulfonate with average aliphatic chain length C11-12	10.3	10.7	14	17	12.2	8.3
	Sodium lauryl sulfate	0	3.5	0	1.4	1.2	0
	Sodium C12-14 alcohol ethoxy-3-sulfate	0	0	0.8	0	0	3
10	C13-15 oxo alcohol ethoxylate with average 7 moles of ethoxylation (Lutensol® A07)	1.57	0	0	0	1.2	0
	C10-Guerbet (2-propylheptan-l-ol) alcohol ethoxylate with average 7 moles of ethoxylation (Lutensol® XP70)	0	1.5	0	0	1.2	0
15	C16-18 alcohol ethoxylate with average 7 moles of ethoxylation	0	0.5	0	0	0.3	0
	C12-18 alcohol ethoxylate with average 5 moles of ethoxylation	0	0.3	0	0	0	0
20	C12-14 alkyl	0	0	0.7	0.54	0.1	1
	hydroxyethyl dimethyl ammonium chloride (Praepagen® HY)						
25	Sodium tripolyphosphate	0	0	0.6	0	1	0
	Zeolite A (builder)	2.7	3.4	0	0	0.5	1.6
	Citric Acid	1.8	2	0	1.4	0	2
30	Sodium citrate	0	1.9	0	0	0	0
	Sodium bicarbonate	29	35	36.7	34	53	22
	Sodium sesquicarbonate dihydrate	0	0	1.2	0	0	0
35	Sodium carbonate	1.2	0	1.9	0	0	0
	Sodium polyacrylate (MW 4000, Sokalan PA25 CL)	0	0	1	0	0	0
	Sodium polyacrylate (MW 8000, Sokalan PA30 CL)	1.45	1.6	0	0.97	1	0
40	Sodium polyacrylate/maleate copolymer MW 70,000, 70:30 ratio, Sokalan® CPS	0	0	0.3	0	0	3
	Polyethylene glycol/vinyl acetate random graft copolymer	0	0	0.8	1	1	0
45	Carboxymethyl cellulose (Finnfix® GDA)	1	0.9	0	0	0	0
	Carboxymethyl cellulose (Finnfix® V)	0	0	0	0.3	1.1	0.92
	Hydrophobically modified carboxymethyl cellulose (Finnfix® SH-1)	0	0	0.5	0	0	0
50	C. I. Fluorescent Brightener 260	0.1	0.13	0.1	0.03	0.05	0.18
	C. I. Fluorescent Brightener 351 (Tinopal® CBS)	0	0.06	0.08	0	0	0
55	Diethylenetriamine pentaacetic acid	0	0	0.2	0.1	0.2	0
	Tetrasodium S,S-ethylenediamine disuccinate	0	0	0	0.3	0	0.3

(continued)

HDD Detergent Compositions						
Ingredient	1 (wt%)	2 (wt%)	3 (wt%)	4 (wt%)	5 (wt%)	6 (wt%)
Diethylenetriamine penta (methylene phosphonic acid), heptasodium salt	0	0.2	0	0	0	0
1-Hydroxyethane-1,1-diphosphonic acid	0.1	0.2	0.3	0	0.2	0.4
2-Phosphonobutane	0	0	0	0.4	0	0
1,2,4- tricarboxylic acid (Bayhibit® AM)						
MgSO ₄	0	0	0	0.8	0	0.4
Sodium percarbonate	9	12	7	6	8	9
Propylene glycol diacetate	7	10	10.8	0	0	0
Triethylene glycol diacetate	0	0	0	5	7	3.9
Oxaziridinium-based bleach booster	0.03	0	0.03	0.02	0.05	0.02
Protease 1	4.3	3.3	6.3	5.7	3.3	0
Protease 2	0	0	0	0	0	2.2
Amylase	2.2	1.51	1	2.2	1.9	3.3
Lipase	0	0	3.6	0	0	2.7
Endoglucanase 1	0	0	5.3	3.3	0	0
Endoglucanase 2	2.1	1.3	0	0	0	2.4
Mannanase	1.3	1.54	1.3	0	1.2	1.9
Perhydrolase 1	2	0	1.8	0	2.1	1.9
Perhydrolase 2	0	4.1	0	2.3	0	0
Direct Violet 9	0	0	0.0003	0.0004	0	0
Solvent Violet 13	0	0	0.002	0	0	0
Texcare® SRA300F	0.3	1.2	0	1	0.33	0.3
Dye lock	0.02	0.02	0	0	0	0
(Tinolux® BMC)	0	0	0	0	0	0.0015
C.I. Food Red 14	0	0	0.001	0	0	0.001
Suds suppressor granule	0.2	0.2	0	0	0.3	0
Moisture	7	6.3	8.9	9.1	4.3	4.6
Perfume	0.2	0.3	0.4	0.3	0.2	0.3
Sodium sulfate	Balance to 100%	Balance to 100%	Balance to 100%	Balance to 100%	Balance to 100%	Balance to 100%

Automatic Dishwashing (ADW) Detergent Compositions				
Formulation	1	2	3	4
Ingredient	Level %wt	Level %wt	Level %wt	Level %wt
Solid ADW detergent composition				
STPP	35	0	0	56
Carbonate	24	45	40	18.5

(continued)

Automatic Dishwashing (ADW) Detergent Compositions				
Formulation	1	2	3	4
Ingredient	Level %wt	Level %wt	Level %wt	Level %wt
Solid ADW detergent composition				
Methylglycine diacetic acid (83% active)	0	15	20	0
Silicate	7	7	7	1.5
TEAD (Tetraacety lethy lenediamine)	0.5	0.5	0.5	3.8
Zinc carbonate	0.5	0.5	0.5	0
SLF18	1.5	1.5	1.5	0
Plurafac LF224				0.6
Penta Amine Acetato-cobalt(III) nitrate (1 % active)	0.5	0.5	0.5	0.6
Percarbonate	15	15	15	11
Sulphonated polymer	10	4	3	5.1
Amylase (14.4mg/g active)	1.3	1.8	1.5	0.7
Processing aids, perfume and sodium sulphate	To balance	To balance	To balance	To balance
Liquid automatic dishwashing detergent composition				
Dipropylene glycol	45	45	45	25
SLF18	45	45	45	0
Neodol1-9	3	3	3	2.6
Lutensol T07				30
Plurafac LF224				32.4
Amine Oxide				3.6
Glycerine	2	2	2	4
Processing aids and Dyes	To balance	To balance	To balance	To balance
Second Liquid automatic dishwashing detergent composition (part of three compartment unit dose)				

HDL Detergent Compositions					
Compound	Formulations				
	I	II	III	IV	V
LAS	24	32	6	3	6
NaC ₁₆ -C ₁₇ HSAS	-	-	-	5	-
C ₁₂ -C ₁₅ AE _{1.8} S	-	-	8	7	5
C ₈ -C ₁₀ propyl dimethyl amine	2	2	2	2	1
C ₁₂ -C ₁₄ alkyl dimethyl amine oxide	-	-	-	-	2
C ₁₂ -C ₁₅ AS alkyl sulphate	-	-	17	-	8
C12-C14 alkyl N-methyl glucamide (CFAA) surfactant	-	5	4	4	3
C ₁₂ -C ₁₄ Fatty alcohol ethoxylate	12	6	1	1	1

(continued)

HDL Detergent Compositions					
Compound	Formulations				
C ₁₂ -C ₁₈ Fatty acid	3	-	4	2	3
Citric acid (anhydrous)	4.5	5	3	2	1
DETPMP	-	-	1	1	0.5
Monoethanolamine	5	5	5	5	2
Sodium hydroxide	-	-	2.5	1	1.5
1 N HCl aqueous solution	#1	#1	-	-	-
Propanediol	12.7	14.5	13.1	10	8
Ethanol	1.8	2.4	4.7	5.4	1
DTPA	0.5	0.4	0.3	0.4	0.5
Pectin Lyase	-	-	-	0.005	-
Amylase	0.001	0.002	-	-	-
Cellulase	-	-	0.0002	-	0.0001
Lipase	0.1	-	0.1	-	0.1
Metalloprotease 1 (optional)	0.05	0.3	-	0.5	0.2
Metalloprotease 2	-	-	0.08	-	-
Protease A (optional)	-	-	-	-	0.1
Aldose Oxidase	-	-	0.3	-	0.003
ZnCl ₂	0.1	0.05	0.05	0.05	0.02
Ca formate	0.05	0.07	0.05	0.06	0.07
DETBCHD	-	-	0.02	0.01	-
SRP1 (anionically end capped polyesters)	0.5	0.5	-	0.3	0.3
Boric acid	-	-	-	-	2.4
Sodium xylene sulfonate	-	-	3	-	-
Sodium cumene sulfonate	-	-	-	0.3	0.5
DC 3225C	1	1	1	1	1
2-butyl-octanol	0.03	0.04	0.04	0.03	0.03
Brightener 1	0.12	0.1	0.18	0.08	0.1
Balance to 100% perfume / dye and/or water					
#1: Add 1N HCl aq. soln to adjust the neat pH of the formula in the range from about 3 to about 5. The pH of Examples above (I)-(II) is about 5 to about 7, and of (III)-(V) is about 7.5 to about 8.5.					

HDL Detergent Compositions						
Compound	Formulations					
	I	II	III	IV	V	VI
LAS	11.5	11.5	9	-	4	-
C ₁₂ -C ₁₅ AE _{2.85} S	-	-	3	18	-	16

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(continued)

HDL Detergent Compositions						
Compound	Formulations					
	I	II	III	IV	V	VI
C ₁₄ -C ₁₅ E _{2.5} S	11.5	11.5	3	-	16	-
C ₁₂ -C ₁₃ E ₉	-	-	3	2	2	1
C ₁₂ -C ₁₃ E ₇	3.2	3.2	-	-	-	-
C12-C14 alkyl N-methyl glucamide (CFAA) surfactant	-	-	-	5	-	3
TPKFA (C12-C14 topped whole cut fatty acids)	2	2	-	2	0.5	2
Citric Acid (Anhydrous)	3.2	3.2	0.5	1.2	2	1.2
Ca formate	0.1	0.1	0.06	0.1	-	-
Na formate	0.5	0.5	0.06	0.1	0.05	0.05
ZnC12	0.1	0.05	0.06	0.03	0.05	0.05
Sodium Cumene Sulfonate	4	4	1	3	1.2	-
Borate	0.6	0.6	1.5	-	-	-
Sodium Hydroxide	6	6	2	3.5	4	3
Ethanol	2	2	1	4	4	3
1,2 Propanediol	3	3	2	8	8	5
Monoethanolamine	3	3	1.5	1	2.5	1
TEPAE (tetraethylene pentaamine ethoxylate)	2	2	-	1	1	1
Metalloprotease 1 (optional)	0.03	0.05	-	0.03	-	0.02
Metalloprotease 2	-	-	0.01	-	0.08	-
Protease A (optional)	-	-	0.01	-	-	-
Lipase	-	-	-	0.002	-	-
Amylase	-	-	-	-	0.002	-
Cellulase	-	-	-	-	-	0.0001
Pectin Lyase	0.005	0.005	-		-	-
Aldose Oxidase	0.05	-	-	0.05	-	0.02
Galactose oxidase	-	0.04				
pentaamine acetate cobalt (III) salt PAAC	0.03	0.03	0.02	-	-	-
DETBCHD	-	-	-	0.02	0.01	-
SRP1 (anionically end capped polyesters)	0.2	0.2	-	0.1	-	-
DTPA	-	-	-	0.3	-	-
polyvinyl pyridine-N-Oxide (PVNO)	-	-	-	0.3	-	0.2
Brightener 1	0.2	0.2	0.07	0.1	-	-
Silicone antifoam	0.04	0.04	0.02	0.1	0.1	0.1
Balance to 100% perfume/dye and/or water						

Liquid Hand Dishwashing (Hand Dish Liquid) Detergent Compositions						
Compound	Formulations					
	I	II	III	IV	V	VI
C ₁₂ -C ₁₅ AE _{1.8} S	30	28	25	-	15	10
LAS	-	-	-	5	15	12
Paraffin Sulfonate	-	-	-	20	-	-
C ₁₀ -C ₁₈ Alkyl Dimethyl Amine Oxide	5	3	7	-	-	-
Betaine	3	-	1	3	1	-
C ₁₂ poly-hydroxy fatty acid amide	-	-	-	3	-	1
C ₁₄ poly-OH fatty acid amide	-	1.5	-	-	-	-
C ₁₁ E ₉	2	-	4	-	-	20
DTPA	-	-	-	-	0.2	-
Tri-sodium Citrate dihydrate (builder)	0.25	-	-	0.7	-	-
Diamine (Dimethyl aminopropyl amine; 1,6-hexane diamine; 1,3-propane diamine; 2-methyl-1,5-pentane diamine; 1,3-pentanediamine; 1-methyl-diaminopropane)	1	5	7	1	5	7
MgCl ₂	0.25	-	-	1	-	-
Metalloprotease 1 (optional)	0.02	0.01	-	0.01	-	0.05
Metalloprotease 2	-	-	0.03	-	0.02	-
Protease A (optional)	-	0.01	-	-	-	-
Amylase	0.001	-	-	0.002	-	0.001
Aldose Oxidase	0.03	-	0.02	-	0.05	-
Sodim Cumene Sulfonate	-	-	-	2	1.5	3
pentaamine acetate cobalt (III) salt	0.01	0.01	0.02	-	-	-
DETBCHD	-	-	-	0.01	0.02	0.01
Balance to 100% perfume / dye and/or water						
The pH of Examples (I)-(VI) is about 8 to about 11.						

Liquid Automatic Dish Washing Detergent Compositions					
Compound	Formulations				
	I	II	III	IV	V
STPP (sodium tripoly phosphate)	16.00	16.00	18.00	16.00	16.00
Potassium Sulfate	-	10.00	8.00	-	10.00
1,2 propanediol	6.00	0.50	2.00	6.00	0.50
Boric Acid	-	-	-	4.00	3.00
CaCl ₂ dihydrate	0.04	0.04	0.04	0.04	0.04
Nonionic surfactant	0.50	0.50	0.50	0.50	0.50
Metalloprotease 1 (optional)	0.10	0.03	-	0.03	-
Metalloprotease 2	-	-	0.05	-	0.06
Protease B (optional)	-	-	-	0.01	-

(continued)

Liquid Automatic Dish Washing Detergent Compositions					
Compound	Formulations				
	I	II	III	IV	V
Amylase	0.02	-	0.02	0.02	-
Aldose Oxidase	-	0.15	0.02	-	0.01
Galactose Oxidase	-	-	0.01	-	0.01
pentaamine acetate cobalt (III) salt PAAC (bleach catalyst)	0.01	-	-	0.01	-
DETBCHD	-	0.01	-	-	0.01
Balance to 100% perfume / dye and/or water					

Granular and/or Tablet Detergent Compositions					
Compound	Formulations				
	I	II	III	IV	V
C ₁₄ -C ₁₅ AS or TAS (sodium tallow alkyl sulfate)	8	5	3	3	3
LAS	8	-	8	-	7
C ₁₂ -C ₁₅ AE ₃ S	0.5	2	1	-	-
C ₁₂ -C ₁₅ E ₅ or E ₃	2	-	5	2	2
QAS (quarternary ammonium salt)	-	-	-	1	1
Zeolite A	20	18	11	-	10
SKS-6 (dry add) (layered silicate)	-	-	9	-	-
MA/AA (acrylate/maleate copolymer)	2	2	2	-	-
AA (polyacrylate polymer)	-	-	-	-	4
3Na Citrate 2H ₂ O	-	2	-	-	-
Citric Acid (Anhydrous)	2	-	1.5	2	-
DTPA	0.2	0.2	-	-	-
EDDS	-	-	0.5	0.1	-
HEDP	-	-	0.2	0.1	-
PB1 (sodium perborate monohydrate)	3	4.8	-	-	4
Percarbonate	-	-	3.8	5.2	-
NOBS	1.9	-	-	-	-
NACA OBS	-	-	2	-	-
TAED	0.5	2	2	5	1
BB1 (3-(3,4-Dihydroisoquinolinium)propane sulfonate (DIPS))	0.06	-	0.34	-	0.14
BB2 3-(3,4-Dihydroisoquinolinium)-decane- 2-sulfate	-	0.14	-	0.2	-
Anhydrous sodium carbonate	15	18	-	15	15
Sulfate	5	12	5	17	3
Silicate	-	1	-	-	8

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(continued)

Granular and/or Tablet Detergent Compositions					
Compound	Formulations				
	I	II	III	IV	V
Metalloprotease 1 (optional)	0.03	-	0.1	0.06	-
Metalloprotease 2	-	0.05	-	-	0.1
Protease B (optional)	-	0.01	-	-	-
Protease C (optional)	-	-	-	0.01	-
Lipase	-	0.008	-	-	-
Amylase	0.001	-	-	-	0.001
Cellulase	-	0.0014	-	-	-
Pectin Lyase	0.001	0.001	0.001	0.001	0.001
Aldose Oxidase	0.03	-	0.05	-	-
pentaamine acetate cobalt (III) salt PAAC	-	0.01	-	-	0.05
Balance to 100% Moisture and/or Minors*					
* Perfume, dye, brightener / SRP1 / Na carboxymethylcellulose/ photobleach / MgSO ₄ / PVPVI/ suds suppressor /high molecular PEG/clay.					

High Density Automatic Dish Washing Detergent Compositions						
Compound	Formulations					
	I	II	III	IV	V	VI
STPP (sodium tripoly phosphate)	-	45	45	-	-	40
3Na Citrate 2H ₂ O	17	-	-	50	40.2	-
Na Carbonate	17.5	14	20	-	8	33.6
Bicarbonate	-	-	-	26	-	-
Silicate	15	15	8	-	25	3.6
Metasilicate	2.5	4.5	4.5	-	-	-
PB1 (sodium perborate monohydrate)	-	-	4.5	-	-	-
PB4 (sodium perborate tetrahydrate)	-	-	-	5	-	-
Percarbonate	-	-	-	-	-	4.8
BB1 (3-(3,4-Dihydroisoquinolinium)propane sulfonate (DIPS))	-	0.1	0.1	-	0.5	-
BB2 3-(3,4-Dihydroisoquinolinium)-decane- 2-sulfate	0.2	0.05	-	0.1	-	0.6
Nonionic detergent	2	1.5	1.5	3	1.9	5.9
HEDP	1	-	-	-	-	-
DETPMP	0.6	-	-	-	-	-
pentaamine acetate cobalt (III) salt PAAC	0.03	0.05	0.02	-	-	-
Paraffin oil Winog 70	0.5	0.4	0.4	0.6	-	-
Metalloprotease 1 (optional)	0.072	0.053	-	0.026	-	0.01
Metalloprotease 2	-	-	0.053	-	0.059	-

(continued)

High Density Automatic Dish Washing Detergent Compositions						
Compound	Formulations					
	I	II	III	IV	V	VI
Protease B (optional)	-	-	-	-	-	0.01
Amylase	0.012	-	0.012	-	0.021	0.006
Lipase	-	0.001	-	0.005	-	-
Pectin Lyase	0.001	0.001	0.001	-	-	-
Aldose Oxidase	0.05	0.05	0.03	0.01	0.02	0.01
BTA (benzotriazole)	0.3	0.2	0.2	0.3	0.3	0.3
Polycarboxylate	6	-	-	-	4	0.9
Perfume	0.2	0.1	0.1	0.2	0.2	0.2
Balance to 100% Moisture and/or Minors*						
*Brightener / dye / SRP1 / Na carboxymethylcellulose/ photobleach / MgSO ₄ / PVPVI/ suds suppressor /high molecular PEG/clay. The pH of Examples (I) through (VI) is from about 9.6 to about 11.3.						

Tablet Detergent Compositions								
Compound	Formulations							
	I	II	III	IV	V	VI	VII	VIII
STPP (sodium tripoly phosphate)	-	48.8	44.7	38.2	-	42.4	46.1	46
3Na Citrate 2H ₂ O	20	-	-	-	35.9	-	-	-
Na Carbonate	20	5	14	15.4	8	23	20	-
Silicate	15	14.8	15	12.6	23.4	2.9	4.3	4.2
Lipase	0.001	-	0.01	-	0.02	-	-	-
Protease B	0.01	-	-	-	-	-	-	-
Protease C	-	-	-	-	-	0.01	-	-
Metalloprotease 1 (optional)	0.01	0.08	-	0.04	-	0.023	-	0.05
Metalloprotease 2	-	-	0.05	-	0.052	-	0.023	-
Amylase	0.012	0.012	0.012	-	0.015	-	0.017	0.002
Pectin Lyase	0.005	-	-	0.002	-	-	-	-
Aldose Oxidase	-	0.03	-	0.02	0.02	-	0.03	-
PB1 (sodium perborate monohydrate)	-	-	3.8	-	7.8	-	-	4.5
Percarbonate	6	-	-	6	-	5	-	-
BB1 (3-(3,4-Dihydroisoquinolinium)propane sulfonate (DIPS))	0.2	-	0.5	-	0.3	0.2	-	-
BB2 3-(3,4-Dihydroisoquinolinium)-decane-2-sulfate	-	0.2	-	0.5	-	-	0.1	0.2

(continued)

Tablet Detergent Compositions								
Compound	Formulations							
	I	II	III	IV	V	VI	VII	VIII
Nonionic surfactant	1.5	2	2	2.2	1	4.2	4	6.5
pentaamine acetate cobalt (III) salt PAAC	0.01	0.01	0.02	-	-	-	-	-
DETBCHD	-	-	-	0.02	0.02	-	-	-
TAED	-	-	-	-	-	2.1	-	1.6
HEDP	1	-	-	0.9	-	0.4	0.2	-
DETPMP	0.7	-	-	-	-	-	-	-
Paraffin oil Winog 70	0.4	0.5	0.5	0.5	-	-	0.5	-
BTA (benzotriazole)	0.2	0.3	0.3	0.3	0.3	0.3	0.3	-
Polycarboxylate	4	-	-	-	4.9	0.6	0.8	-
PEG 400-30,000	-	-	-	-	-	2	-	2
Glycerol	-	-	-	-	-	0.4	-	0.5
Perfume	-	-	-	0.05	0.2	0.2	0.2	0.2
Balance to 100% Moisture and/or Minors*								
*Brightener / SRP1 / Na carboxymethylcellulose/ photobleach / MgSO ₄ / PVPVI/ suds suppressor /high molecular PEG/clay. The pH of Examples (I) through (VII) is from about 10 to about 11.5; pH of (VIII) is from 8-10. The tablet weight of Examples (I) through (VIII) is from about 20 grams to about 30 grams.								

Liquid Hard Surface Detergent Compositions								
Compound	Formulations							
	I	II	III	IV	V	VI	VII	
C ₉ -C ₁₁ E ₅	2.4	1.9	2.5	2.5	2.5	2.4	2.5	
C ₁₂ -C ₁₄ E ₅	3.6	2.9	2.5	2.5	2.5	3.6	2.5	
C ₇ -C ₉ E ₆	-	-	-	-	8	-	-	
C ₁₂ -C ₁₄ E ₂₁	1	0.8	4	2	2	1	2	
LAS	-	-	-	0.8	0.8	-	0.8	
Sodim Cumene Sulfonate	1.5	2.6	-	1.5	1.5	1.5	1.5	
Isachem [®] AS (branched alcohol alkyl sulfate)	0.6	0.6	-	-	-	0.6	-	
Na ₂ CO ₃	0.6	0.13	0.6	0.1	0.2	0.6	0.2	
3Na Citrate 2H ₂ O	0.5	0.56	0.5	0.6	0.75	0.5	0.75	
NaOH	0.3	0.33	0.3	0.3	0.5	0.3	0.5	
Fatty Acid	0.6	0.13	0.6	0.1	0.4	0.6	0.4	
2-butyl octanol	0.3	0.3	-	0.3	0.3	0.3	0.3	
PEG DME-2000 [®]	0.4	-	0.3	0.35	0.5	-	-	
PVP (vinylpyrrolidone homopolymer)	0.3	0.4	0.6	0.3	0.5	-	-	

(continued)

Liquid Hard Surface Detergent Compositions							
Compound		Formulations					
5	MME PEG (2000)®	-	-	-	-	0.5	0.5
	Jeffamine® ED-2001 (capped polyethylene glycol)	-	0.4	-	-	0.5	-
	pentaamine acetate cobalt (III) salt PAAC	-	-	-	0.03	0.03	0.03
10	DETBCHD	0.03	0.05	0.05	-	-	-
	Metalloprotease 1 (optional)	0.07	-	0.08	0.03	-	0.01
	Metalloprotease 2	-	0.05	-	-	0.06	-
15	Protease B (optional)	-	-	-	-	-	0.01
	Amylase	0.12	0.01	0.01	-	0.02	-
	Lipase	-	0.001	-	0.005	-	0.005
	Pectin Lyase	0.001	-	0.001	-	-	-
20	ZnCl ₂	0.02	0.01	0.03	0.05	0.1	0.05
	Calcium Formate	0.03	0.03	0.01	-	-	-
	PB1 (sodium perborate monohydrate)	-	4.6	-	3.8	-	-
25	Aldose Oxidase	0.05	-	0.03	-	0.02	0.02
Balance to 100% perfume / dye and/or water							
The pH of Examples (I) through (VII) is from about 7.4 to about 9.5.							

HDL Detergent Compositions				
Ingredient	Composition			
	(wt% of composition)			
35	1	2	3	4
C ₁₂₋₁₅ Alkylethoxy(1.8)sulfate	14.7	11.6		16.31
C _{11.8} Alkylbenzene sulfonate	4.3	11.6	8.3	7.73
40 C ₁₆₋₁₇ Branched alkyl sulfate	1.7	1.29		3.09
C ₁₂₋₁₄ Alkyl-9-ethoxylate	0.9	1.07		1.31
C ₁₂ dimethylamine oxide	0.6	0.64		1.03
Citric acid	3.5	0.65	3	0.66
45 C ₁₂₋₁₈ fatty acid	1.5	2.32	3.6	1.52
Sodium Borate (Borax)	2.5	2.46	1.2	2.53
Sodium C ₁₂₋₁₄ alkyl ethoxy 3 sulfate			2.9	
50 C ₁₄₋₁₅ alkyl 7-ethoxylate			4.2	
C ₁₂₋₁₄ Alkyl-7-ethoxylate			1.7	
Ca formate	0.09	0.09		0.09
55 A compound having the following general structure: bis((C ₂ H ₅ O)(C ₂ H ₄ O) _n)(CH ₃)-N ⁺ -C _x H _{2x} -N ⁺ -(CH ₃)-bis((C ₂ H ₅ O)(C ₂ H ₄ O) _n), wherein n = from 20 to 30, and x = from 3 to 8, or sulphated or sulphonated variants thereof			1.2	

(continued)

HDL Detergent Compositions				
Ingredient	Composition			
	(wt% of composition)			
	1	2	3	4
Random graft co-polymer ¹		1.46	0.5	
Ethoxylated Polyethylenimine ²	1.5	1.29		1.44
Diethylene triamine pentaacetic acid	0.34	0.64		0.34
Diethylene triamine penta(methylene phosphonic acid)			0.3	
Tinopal AMS-GX		0.06		
Tinopal CBS-X	0.2	0.17		0.29
Amphiphilic alkoxyated grease cleaning polymer ³	1.28	1	0.4	1.93
Ethanol	2	1.58	1.6	5.4
Propylene Glycol	3.9	3.59	1.3	4.3
Diethylene glycol	1.05	1.54		1.15
Polyethylene glycol	0.06	0.04		0.1
Monoethanolamine	3.05	2.41	0.4	1.26
NaOH	2.44	1.8		3.01
Sodium Cumene Sulphonate			1	
Sodium Formate		0.11		0.09
Water, Aesthetics (Dyes, perfumes) and Minors (Enzymes, solvents, structurants)	balance	balance	balance	balance
1 Random graft copolymer is a polyvinyl acetate grafted				
polyethylene oxide copolymer having a polyethylene oxide backbone and multiple polyvinyl acetate side chains. The molecular weight of the polyethylene oxide backbone is about 6000 and the weight ratio of the polyethylene oxide to polyvinyl acetate is about 40 to 60 and no more than 1 grafting point per 50 ethylene oxide units.				
2 Polyethylenimine (MW = 600) with 20 ethoxylate groups per -NH.				
3 Amphiphilic alkoxyated grease cleaning polymer is a polyethylenimine (MW = 600) with 24 ethoxylate groups per -NH and 16 propoxylate groups per -NH.				

Light-Duty Liquid Dishwashing Detergent Compositions				
Composition	1	2	3	4
Linear Alkylbenzene Sulfonate (1)	-		-	-
Alkyl Ethoxy Sulfate (2)	18%	17%	17%	18%
Paraffin Sulfonate (C15)	-	-	-	-
CAP= coco amido propyl Betaine	-	-	9%	5%
Nonionic (3)	-	-	1%	-
Amine Oxide (4)	6%	5.50%	-	4%

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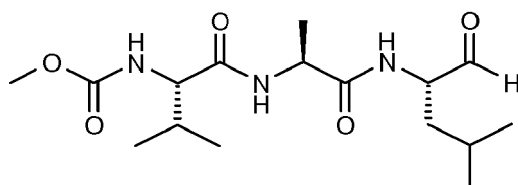
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Light-Duty Liquid Dishwashing Detergent Compositions				
Composition	1	2	3	4
Alkylpolyglucoside				4%
Alcohol (5)	-	-	5%	7%
Pura= polypropyleneglycol	1%	0.80%	-	-
Citrate	-	-	0.30%	0.60%
Salt (6)	1.20%	1.00%	-	0.50%
SCS= sodium cumene sulfonate	-	-	0.80%	-
glycerol	15%	5%	3%	-
Na-lactate	-	-	-	5%
cationic polymer (7)	0.10%	0.10%	0.30%	0.20%
Present amylase	0.0075	0.005	0.0025	0.03
Glycol distearate from Euperlan® Cognis	0.4	0	0.4	0
Hydrogenated Castor Oil Thixcin® Elementis	0	0.1	0	0.1
Mica (BASF Mearlin superfine)	0	0.05	0	0.05
Minors*	Balance to 100% with water			
pH	9	9	6	6
Optional Minors*: dyes, opacifier, perfumes, preservatives, hydrotropes, processing aids, and/or stabilizers.				
(1) Linear Alkylbenzene Sulfonate: LAS: C11.4				
(2) Alkyl Ethoxy Sulfate: AExS :				
(3) Nonionic: AlkylEthoxylate				
(4) Di-methyl coco alkyl amine oxide				
(5) Alcohol: Ethanol				
(6) Salt: NaCl				
(7) cationically modified hydroxyethyl cellulose (Polyquaternium-10 - UCARE LR-400 ex Amerchol).				

Liquid laundry detergent compositions suitable for front-loading automatic washing machines										
Ingredient	Composition									
	(wt% of composition)									
	1	2	3	4	5	6	7	8		
Alkylbenzene sulfonic acid	7	11	4.5	1.2	1.5	12.5	5.2	4		
Sodium C ₁₂₋₁₄ alkyl ethoxy 3 sulfate	2.3	3.5	4.5	4.5	7	18	1.8	2		
C ₁₄₋₁₅ alkyl 8-ethoxylate	5	8	2.5	2.6	4.5	4	3.7	2		
C ₁₂ alkyl dimethyl amine oxide	-	-	0.2	-	-	-	-	-		
C ₁₂₋₁₄ alkyl hydroxyethyl dimethyl ammonium chloride	-	-	-	0.5	-	-	-	-		
C ₁₂₋₁₈ Fatty acid	2.6	4	4	2.6	2.8	11	2.6	1.5		
Citric acid	2.6	3	1.5	2	2.5	3.5	2.6	2		
Protease *	0.05	0.03	0.04	0.03	0.04	0.03	0.03	0.02		
Amylase	0.1	0.2	0.15	-	0.05	0.5	0.1	0.2		
Mannanase	0.05	0.1	0.05	-	-	0.1	0.04	-		
Random graft co-polymer ¹	1	0.2	1	0.4	0.5	2.7	0.3	1		
A compound having the following general structure: bis((C ₂ H ₅ O)(C ₂ H ₄ O) _n)(CH ₃) ₂ N ⁺ -C _x H _{2x} -N ⁺ -(CH ₃) ₂ -bis((C ₂ H ₅ O)(C ₂ H ₄ O) _n), wherein n = from 20 to 30, and x = from 3 to 8, or sulphated or sulphonated variants thereof	0.4	2	0.4	0.6	1.5	1.8	0.7	0.3		
Ethoxylated Polyethylenimine ²	-	-	-	-	-	0.5	-	-		
Amphiphilic alkoxyated grease cleaning polymer ³	0.1	0.2	0.1	0.2	0.3	0.3	0.2	0.3		
Diethoxylated poly (1,2 propylene terephthalate)	-	-	-	-	-	-	0.3	-		
Diethylenetriaminopenta(methylenephosphonic) acid	0.2	0.3	-	-	0.2	-	0.2	0.3		
Hydroxyethane diphosphonic acid	-	-	0.45	-	-	1.5	-	0.1		
FWA (fluorescent whitening agent)	0.1	0.2	0.1	-	-	0.2	0.05	0.1		
Solvents (1,2 propanediol, ethanol),	3	4	1.5	1.5	2	4.3	2	1.5		
Hydrogenated castor oil derivative	0.4	0.4	0.3	0.1	0.3	-	0.4	0.5		
Boric acid	1.5	2.5		1.5	1.5	0.5	1.5	1.5		

(continued)

Liquid laundry detergent compositions suitable for front-loading automatic washing machines									
	Composition								
Ingredient	(wt% of composition)								
Na formate	-	-	-	1	-	-	-	-	-
Reversible protease inhibitor ⁴	-	-	0.002	-	-	-	-	-	-
Perfume	0.5	0.7	0.5	0.5	0.8	1.5	0.5	0.8	0.8
Perfume MicroCapsules slurry (30%am)	0.2	0.3	0.7	0.2	0.05	0.4	0.9	0.7	0.7
Ethoxylated thiophene Hueing Dye ⁵	0.005	0.007	0.01	0.008	0.008	0.007	0.007	0.008	0.008
Buffers (sodium hydroxide, Monoethanolamine)	To pH 8.2								
Water and minors (antifoam, aesthetics)	To 100%								
¹ Random graft copolymer is a polyvinyl acetate grafted polyethylene oxide copolymer having a polyethylene oxide backbone and multiple polyvinyl acetate side chains. The molecular weight of the polyethylene oxide backbone is about 6000 and the weight ratio of the polyethylene oxide to polyvinyl acetate is about 40 to 60 and no more than 1 grafting point per 50 ethylene oxide units.									
² Polyethylenimine (MW = 600) with 20 ethoxylate groups per -NH.									
³ Amphiphilic alkoxyated grease cleaning polymer is a polyethylenimine (MW = 600) with 24 ethoxylate groups per -NH and 16 propoxylate groups per -NH									
⁵ Ethoxylated thiophene Hueing Dye is as described in US 7,208,459 B2.									
* Remark: all enzyme levels expressed as % enzyme raw material, except for protease which is expressed as % of active protein added to the product..									
⁴ Reversible Protease inhibitor of structure shown below the Table.									



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Liquid laundry detergent compositions suitable for top-loading automatic washing machines										
Ingredient	Composition									
	(wt% of composition)									
	1	2	3	4	5	6	7	8		
C ₁₂₋₁₅ Alkylethoxy(1.8)sulfate	20.1	15.1	20	15.1	13.7	16.7	10	9.9		
C _{11.8} Alkylbenzene sulfonate	2.7	2	1	2	5.5	5.6	3	3.9		
C ₁₆₋₁₇ Branched alkyl sulfate	6.5	4.9		4.9	3	9	2			
C ₁₂₋₁₄ Alkyl-9-ethoxylate	0.8	0.8	0.8	0.8	8	1.5	0.3	11.5		
C ₁₂ dimethylamine oxide			0.9							
Citric acid	3.8	3.8	3.8	3.8	3.5	3.5	2	2.1		
C ₁₂₋₁₈ fatty acid	2	1.5	2	1.5	4.5	2.3		0.9		
Protease*	0.1	0.2	0.1	0.1	0.1	0.1	0.1	0.1		
Amylase 1	0.7	0.3	0.6	0.3	0.6	0.4				
Amylase 2								1.1		
Mannanase	0.1					0.1				
Pectate Lyase	0.1					0.2				
Borax	3	3			2	3	3	3.3		
Na & Ca formate	0.2	0.2		0.2	0.2		0.7			
A compound having the following general structure: bis((C ₂ H ₅ O)(C ₂ H ₄ O) _n)(C H ₃)-N ⁺ -C _x H _{2x} -N ⁺ -(CH ₃)-bis((C ₂ H ₅ O)(C ₂ H ₄ O) _n), wherein n = from 20 to 30, and x = from 3 to 8, or sulphated or sulphonated variants thereof	1.6	1.6	3	1.6	2	1.6	1.3	1.2		
	0.4	0.2	1	0.5	0.6	1	0.8	1		
Random graft co-polymer ¹										
Diethylene triamine pentaacetic acid	0.4	0.4	0.4	0.4	0.2	0.3	0.8			
Tinopal AMS-GX (brightener)	0.2	0.2	0.2	0.2	0.2	0.3	0.1			
Tinopal CBS-X (brightener)						0.1				
Amphiphilic alkoxyated grease cleaning polymer ³	1	1.3	1.3	1.4	1	1.1	1	1		

(continued)

Liquid laundry detergent compositions suitable for top-loading automatic washing machines									
Ingredient	Composition								
	(wt% of composition)								
1	2	3	4	5	6	7	8		
Texcare 240N (Clariant)			1						
Ethanol	2.6	2.6	2.6	1.8	3	1.3			
Propylene Glycol	4.6	4.6	4.6	3	4	2.5			
Diethylene glycol	3	3	3	3	2.7	3.6			
Polyethylene glycol	0.2	0.2	0.2	0.1	0.3	0.1	1.4		
Monoethanolamine	2.7	2.7	2.7	4.7	3.3	1.7	0.4		
Triethanolamine							0.9		
NaOH	to pH 8.3	to pH 8.3	to pH 8.3	to pH 8.3	to pH 8.3	to pH 8.3	to pH 8.5		
Suds suppressor									
Dye	0.01	0.01	0.01	0.01	0.01	0.01	0		
Perfume	0.5	0.5	0.5	0.7	0.7	0.8	0.6		
Perfume MicroCapsules slurry (30%am)	0.2	0.5	0.2	0.3	0.3	0.9	1		
Ethoxylated thiophene Hueing Dye ⁵	0.003	0.002	0.002	0.005	0.002	0.004	0.003		
Water	Balance	Balance	Balance	Balance	Balance	Balance	Balance		
¹ Random graft copolymer is a polyvinyl acetate grafted polyethylene oxide copolymer having a polyethylene oxide backbone and multiple polyvinyl acetate side chains. The molecular weight of the polyethylene oxide backbone is about 6000 and the weight ratio of the polyethylene oxide to polyvinyl acetate is about 40 to 60 and no more than 1 grafting point per 50 ethylene oxide units.									
³ Amphiphilic alkoxylated grease cleaning polymer is a polyethylenimine (MW = 600) with 24 ethoxylate groups per -NH and 16 propoxylate groups per -NH									
⁵ Ethoxylated thiophene Hueing Dye is as described in US 7,208,459 B2.									
*Remark: all enzyme levels expressed as % enzyme raw material, except for protease which is expressed as % of active protein added to the product..									

Granular detergent compositions						
Component	1	2	3	4	5	6
Linear alkylbenzenesulfonate with aliphatic carbon chain length C ₁₁ -C ₁₂	15	12	20	10	12	13
Other surfactants	1.6	1.2	1.9	3.2	0.5	1.2
Phosphate builder(s)	2	3	4			
Zeolite		1		1	4	1
Silicate	4	5	2	3	3	5
Sodium Carbonate	2	5	5	4	0	3
Polyacrylate (MW 4500)	1	0.6	1	1	1.5	1
Carboxymethyl cellulose (Finnfix BDA ex CPKelco)	1	-	0.3	-	1.1	-
Cellulase	0.23	0.17	0.5	0.2	0.2	0.6
Protease	0.23	0.17	0.5	0.2	0.2	0.6
Amylase	0.23	0.17	0.5	0.2	0.2	0.6
Fluorescent Brightener(s)	0.16	0.06	0.16	0.18	0.16	0.16
Diethylenetriamine pentaacetic acid or Ethylene diamine tetraacetic acid	0.6		0.6	0.25	0.6	0.6
MgSO ₄	1	1	1	0.5	1	1
Bleach(es) and Bleach activator(s)	6.88		6.12	2.09	1.17	4.66
Ethoxylated thiophene Hueing Dye ⁵	0.002	0.001	0.003	0.003	-	-
Direct Violet 9 ex Ciba Specialty Chemicals				0.0006	0.0004	0.0006
Sulfate/Citric Acid/ Sodium Bicarbonate/Moisture/ perfume	Balance to 100%					
⁵ Ethoxylated thiophene Hueing Dye is as described in US 7,208,459 B2.						

Granular Laundry Detergent Compositions and Their Components						
Component	Detergent Compositions					
	1	2	3	4	5	6
Linear alkylbenzenesulfonate with aliphatic carbon chain length C ₁₁ -C ₁₂	15	12	20	10	12	13
Other surfactants	1.6	1.2	1.9	3.2	0.5	1.2
Phosphate builder(s)	2	3	4			
Zeolite		1		1	4	1
Silicate	4	5	2	3	3	5
Sodium Carbonate	2	5	5	4	0	3
Polyacrylate (MW 4500)	1	0.6	1	1	1.5	1
Carboxymethyl cellulose	1	-	0.3	-	1.1	-
Cellulase (15.6mg/g)	0.23	0.17	0.5	0.2	0.2	0.6
Protease	0.23	0.17	0.05	0.2	0.03	0.1
Amylase (14mg/g)	0.23	0.17	0.5	0.2	0.2	0.6

(continued)

Granular Laundry Detergent Compositions and Their Components						
Component	Detergent Compositions					
	1	2	3	4	5	6
Mannanase (4mg/g)	0.1			0.1		0.1
Lipase (18.6mg/g)	0.2		0.1		0.3	
Fluorescent Brightener(s)	0.16	0.06	0.16	0.18	0.16	0.16
Diethylenetriamine pentaacetic acid or Ethylene diamine tetraacetic acid	0.6		0.6	0.25	0.6	0.6
MgSO ₄	1	1	1	0.5	1	1
Bleach(es) and Bleach activator(s)	6.88		6.12	2.09	1.17	4.66
Ethoxylated thiophene Hueing Dye ⁵	0.002	0.001	0.003	0.003	-	-
Direct Violet 9 ex Ciba Specialty Chemicals				0.0006	0.0004	0.0006
Sulfate/Citric Acid/ Sodium Bicarbonate/ Moisture/ perfume	Balance to 100%					
5Ethoxylated thiophene Hueing Dye is as described in US 7,208,459 B2.						

Granular Laundry Detergent Compositions and Their Components					
Component	Detergent Composition				
	7	8	9	10	11
Surfactants					
C ₁₆₋₁₇ Branched alkyl sulfate	3.55	15.8			
C ₁₂₋₁₄ alkyl sulphate			1.5		
Sodium linear alkylbenzenesulfonate with aliphatic chain length C ₁₁ -C ₁₂	9.6		10.6	7.5	9
Sodium C _{14/15} alcohol ethoxy - 3 - sulfate	1.15			2.88	
Sodium C _{14/15} alkyl sulphate	2.37				
C _{14/15} alcohol ethoxylate with average 7 moles of ethoxylation				1.17	1
mono-C ₈₋₁₀ alkyl mono-hydroxyethyl di-methyl quaternary ammonium chloride					0.45
Di methyl hydroxyl ethyl lauryl ammonium chloride			0.18		
Zeolite A	13.9	4.7	0.01	2.9	1.8
Sodium Silicate 1.6.ratio	4	0.2		4	4
Sodium Silicate 2.35.ratio			8		
Citric Acid				2.5	1.4
Sodium tripolyphosphate			5		
Sodium Carbonate	24.1	30	16.9	24.4	21
Nonanoyloxybenzenesuphionate	5.78	2.81	0.96		
Oxaziridinium-based bleach booster				0.03	0.017
Tetrasodium S,S,-ethylenediaminedisuccinate				0.2	

(continued)

Granular Laundry Detergent Compositions and Their Components					
Component	Detergent Composition				
	7	8	9	10	11
Diethylenetriamine penta (methylene phosphonic acid), heptasodium salt	0.61				0.33
Hydroxyethane dimethylene phosphonic acid				0.29	0.45
Ethylene diamine tetraacetate			0.27		
MgSO ₄			0.47	0.5994	0.782
Sodium Percarbonate	7	4.4		15.9	19.1
Tetra Acetyl Ethylene Diamine				3.3	4.6
Sodium Perborate Monohydrate			1.2		
Carboxymethyl cellulose	0.1		0.17	1.69	0.23
(e.g., Finnfix BDA ex CPKelco)					
Sodium Acrylic acid/maleic acid co-polymer (70/30)	0.0236	3.8		2	2.5
Sodium polyacrylate (Sokalan PA30 CL)	4		0.84		
Terephthalate polymer				0.23	
Polyethylene glycol/vinyl acetate random graft co polymer			0.89	0.89	0.91
Photobleach- zinc phthalocyanine tetrasulfonate			0.005	0.001	0.002
C.I. Fluorescent Brightener 260	0.11	0.15	0.04	0.23	0.15
C.I. Fluorescent Brightener 351			0.1		
(Tinopal® CBS)					
Suds suppressor granule		0.25		0.07	0.04
Hydrophobically modified carboxy methyl cellulose (Finnifix® SH-1)			0.019	0.028	
Bentonite			8.35		
Miscellaneous (Dyes, perfumes, process aids, moisture and sodium sulphate)	Balance	Balance	Balance	Balance	Balance

Unit Dose Detergent Compositions					
Ingredients	1	2	3	4	5
Alkylbenzene sulfonic acid C 11-13, 23.5% 2-phenyl isomer	14.5	14.5	14.5	14.5	14.5
C ₁₂₋₁₄ alkyl ethoxy 3 sulfate	7.5	7.5	7.5	7.5	7.5
C ₁₂₋₁₄ alkyl 7-ethoxylate	13	13	13	13	13
Citric Acid	0.6	0.6	0.6	0.6	0.6
Fatty Acid	14.8	14.8	14.8	14.8	14.8
Enzymes (as % raw material not active)	1.7	1.7	1.7	1.7	1.7
Present amylase (as % active)	0.05	0.1	0.02	0.03	0.03
Ethoxylated Polyethylenimine ¹	4	4	4	4	4

(continued)

Unit Dose Detergent Compositions					
Ingredients	1	2	3	4	5
Series 1 GG36 protease (as % active)	0.02	0	0.01	0.02	0.03
Hydroxyethane diphosphonic acid	1.2	1.2	1.2	1.2	1.2
Brightener	0.3	0.3	0.3	0.3	0.3
P-diol	15.8	13.8	13.8	13.8	13.8
Glycerol	6.1	6.1	6.1	6.1	6.1
MEA (monoethanolamide) brightener stabilizer	8	8	8	8	8
TIPA (triisopropanolamine)	-	-	2	-	-
TEA (triethanolamine)	-	2	-	-	-
Cumene sulphonate	-	-	-	-	2
cyclohexyl dimethanol	-	-	-	2	-
Water	10	10	10	10	10
Structurant	0.14	0.14	0.14	0.14	0.14
Perfume	1.9	1.9	1.9	1.9	1.9
Buffers (monoethanolamine)	To pH 8.0				
Solvents (1,2 propanediol, cthanol)	To 100%				
¹Polyethylenimine (MW = 600) with 20 ethoxylate groups per -NH.					
Multiple Compartment Unit Dose Detergent Compositions					
Base Composition 1		%			
Ingredients					
Glycerol (min 99)		5.3			
1,2-propanediol		10			
Citric Acid		0.5			
Monoethanolamine		10			
Caustic soda		-			
Dequest 2010		1.1			
Potassium sulfite		0.2			
Nonionic Marlipal C24EO7		20.1			
HLAS (surfactant)		24.6			
Optical brightener FWA49		0.2			
C12-15 Fatty acid		16.4			
Polymer Lutensit Z96		2.9			
Polyethyleneimine ethoxylate PEI600 E20		1.1			
MgCl2		0.2			
Solvents (1,2 propanediol, ethanol)		To 100%			

Multi-compartment formulations							
Composition	1				2		
Compartment	A	B	C		A	B	C
Volume of each compartment	40 ml	5 ml	5 ml		40 ml	5 ml	5 ml
Active material in Wt. %							
Perfume	1.6	1.6	1.6		1.6	1.6	1.6
Dyes	< 0.01	< 0.01	< 0.01		< 0.01	< 0.01	< 0.01
TiO ₂	0.1	-	-		-	0.1	-
Sodium Sulfite	0.4	0.4	0.4		0.3	0.3	0.3
Acusol 305, Rohm&Haas	1.2				2	-	-
Hydrogenated castor oil	0.14	0.14	0.14		0.14	0.14	0.14
Base Composition 1	Add to 100%	Add to 100%	Add to 100%		Add to 100%	Add to 100%	Add to 100%

Phosphate-Free Detergent: IEC-60436 WFK Type B (pH=10.4 in 3g/l)	
Component	Wt %
Sodium citrate dehydrate	30
Maleic acid/ Acrylic acid copolymer sodium Salt SOKALAN® CP5 BASF	12
Sodium perborate monohydrate	5
TAED	2
Sodium disilicate: Prottil A (Cognis)	25
Linear fatty alcohol ethoxylate	2
Sodium carbonate anhydrous	add to 100

Phosphate-Containing Detergent: IEC-60436 WFK Type C (pH=10.5 in 3 g/l)	
Component	Wt %
Sodium tripolyphosphate	23
Sodium citrate dehydrate	22.3
Maleic acid/ Acrylic acid copolymer sodium salt	4
Sodium perborate monohydrate	6
TAED	2
Sodium disilicate: Prottil A (Cognis)	5
Linear fatty alcohol ethoxylate	2
Sodium carbonate anhydrous	add to 100

Liquid laundry detergent compositions suitable for top-loading automatic washing machines (1 &2) and front loading washing machines (3).

5	Ingredient	Composition		
		(wt% of composition)		
		1	2	3
10	C ₁₂₋₁₅ Alkylethoxy(1.8)sulfate	14.7	11.6	
	C _{11.8} Alkylbenzene sulfonate	4.3	11.6	8.3
	C ₁₆₋₁₇ Branched alkyl sulfate	1.7	1.29	
15	C ₁₂₋₁₄ Alkyl -9-ethoxylate	0.9	1.07	
	C ₁₂ dimethylamine oxide	0.6	0.64	
	Citric acid	3.5	0.65	3
20	C ₁₂₋₁₈ fatty acid	1.5	2.32	3.6
	Sodium Borate (Borax)	2.5	2.46	1.2
	Sodium C ₁₂₋₁₄ alkyl ethoxy 3 sulfate			2.9
25	C ₁₄₋₁₅ alkyl 7-ethoxylate			4.2
	C ₁₂₋₁₄ Alkyl -7-ethoxylate			1.7
	Ca formate	0.09	0.09	
30	A compound having the following general structure: bis((C ₂ H ₅ O)(C ₂ H ₄ O) n)(CH ₃)-N ⁺ -C _x H _{2x} -N ⁺ -(CH ₃)-bis((C ₂ H ₅ O)(C ₂ H ₄ O)n), wherein n = from 20 to 30, and x = from 3 to 8, or sulphated or sulphonated variants thereof			1.2
	Random graft co-polymer ¹		1.46	0.5
	Ethoxylated Polyethylenimine ²	1.5	1.29	
35	Diethylene triamine pentaacetic acid	0.34	0.64	
	Diethylene triamine penta(methylene phosphonic acid)			0.3
	Tinopal AMS-GX		0.06	
40	Tinopal CBS-X	0.2	0.17	
	Amphiphilic alkoxyated grease cleaning polymer ³	1.28	1	0.4
	Ethanol	2	1.58	1.6
45	Propylene Glycol	3.9	3.59	1.3
	Diethylene glycol	1.05	1.54	
	Polyethylene glycol	0.06	0.04	
50	Monoethanolamine	3.05	2.41	0.4
	NaOH	2.44	1.8	
	Sodium Cumene Sulphonate			1
	Sodium Formate		0.11	

(continued)

Liquid laundry detergent compositions suitable for top-loading automatic washing machines (1 & 2) and front loading washing machines (3).

	Ingredient	Composition		
		(wt% of composition)		
		1	2	3
5				
10	Water, Aesthetics (Dyes, perfumes) and Minors (Enzymes, solvents, structurants)	balance	balance	balance
15	¹ Random graft copolymer is a polyvinyl acetate grafted polyethylene oxide copolymer having a polyethylene oxide backbone and multiple polyvinyl acetate side chains. The molecular weight of the polyethylene oxide backbone is about 6000 and the weight ratio of the polyethylene oxide to polyvinyl acetate is about 40 to 60 and no more than 1 grafting point per 50 ethylene oxide units. ² Polyethylenimine (MW = 600) with 20 ethoxylate groups per -NH. ³ Amphiphilic alkoxyated grease cleaning polymer is a polyethylenimine (MW = 600) with 24 ethoxylate groups per -NH and 16 propoxylate groups per -NH			

Granular laundry detergent compositions suitable for top-loading automatic washing machines (1-3) and front loading washing machines (4-5). The present amylase is separately added to these formulations.

	Ingredients	1	2	3	4	5
25	C ₁₆₋₁₇ Branched alkyl sulfate	3.55				
	C ₁₂₋₁₄ alkyl sulphate			1.5		
30	Sodium linear alkylbenzenesulfonate with aliphatic chain length C ₁₁ -C ₁₂	9.6	15.8	10.6	7.5	9
	Sodium C _{14/15} alcohol ethoxy - 3 - sulfate	1.15			2.88	
	Sodium C _{14/15} alkyl sulphate	2.37				
35	C _{14/15} alcohol ethoxylate with average 7 moles of ethoxylation				1.17	1
	mono-C ₈₋₁₀ alkyl mono-hydroxyethyl di-methyl quaternary ammonium chloride					0.45
40	Di methyl hydroxyl ethyl lauryl ammonium chloride			0.18		
	Zeolite A	13.9	4.7	0.01	2.9	1.8
	Sodium Silicate 1.6.ratio	4	0.2		4	4
	Sodium Silicate 2.35.ratio			8		
45	Citric Acid				2.5	1.4
	Sodium tripolyphosphate			5		
	Sodium Carbonate	24.1	30	16.9	24.4	21
50	Nonanoyloxybenzenesuphphonate	5.78	2.81	0.96		
	Oxaziridinium-based bleach booster				0.03	0.017
	Tetrasodium S, S,-ethylenediaminedisuccinate				0.2	
55	Diethylenetriamine penta (methylene phosphonic acid), heptasodium salt	0.61				0.33
	Hydroxyethane dimethylene phosphonic acid				0.29	0.45

(continued)

Granular laundry detergent compositions suitable for top-loading automatic washing machines (1-3) and front loading washing machines (4-5). The present amylase is separately added to these formulations.

Ingredients	1	2	3	4	5
Ethylene diamine tetraacetate			0.27		
MgSO ₄			0.47	0.5994	0.782
Sodium Percarbonate	7	4.4		15.9	19.1
Tetra Acetyl Ethylene Diamine				3.3	4.6
Sodium Perborate Monohydrate			1.2		
Carboxymethyl cellulose (e.g. Finnfix BDA ex CPKelco)	0.1		0.17	1.69	0.23
Sodium Acrylic acid/maleic acid co-polymer (70/30)	0.0236	3.8		2	2.5
Sodium polyacrylate (Sokalan PA30 CL)	4		0.84		
Terephthalate polymer				0.23	
Polyethylene glycol/vinyl acetate random graft co polymer			0.89	0.89	0.91
Photobleach- zinc phthalocyanine tetrasulfonate			0.005	0.001	0.002
C.I.Fluorescent Brightener 260	0.11	0.15	0.04	0.23	0.15
C.I.Fluorescent Brightener 351 (Tinopal [®] CBS)			0.1		
Suds suppressor granule		0.25		0.07	0.04
Hyrdrophobically modified carboxy methyl cellulose (Finnifix [®] SH-1)			0.019	0.028	
Bentonite			8.35		
Miscellaneous (Dyes, perfumes, process aids, moisture and sodium sulphate)	Balance	Balance	Balance	Balance	Balance

Granular Laundry Detergent Compositions and Their Components. The present amylase is separately added to these formulations.														
	Detergent Composition													
Component Sur- factants	A	B	C	D	E	F	G	H	I	J	K	L	M	N
C ₁₀ Nonionic				0.1843			0.1142	0.2894	0.1885	0.1846	0.1885	0.1979	0.1979	0.1979
C ₁₆₋₁₇ Branched alkyl sulfate	3.53	3.53	3.53											
C ₁₂₋₁₄ alkyl sulphate														
Sodium linear alkylbenzenesulfonate with aliphatic chain length C ₁₁ -C ₁₂														
Sodium C _{14/15} alcohol ethoxv - 3 - sulfate	8.98	8.98	8.98	13.58	14.75	12.94	15.69	9.01	8.42	9.51	8.92	8.92	11.5	11.5
Sodium C _{14/15} alkyl sulphate	1.28	1.28	1.28								1.62	1.62	1.125	1.125
	2.36	2.36	2.36											
C _{12/14} alcohol ethoxylate with average 7 moles of ethoxylation						2.9								
C _{12/14} alcohol ethoxylate with average 3 moles of ethoxylation									2.44					
C _{14/15} alcohol ethoxylate with average 7 moles of ethoxylation								0.97	1.17	0.97	1	1	1.5	1.5
mono-C ₈₋₁₀ alkyl mono- hydroxyethyl di-methyl quaternary ammonium chloride								0.45						
Dimethyl hydroxyl ethyl lauryl ammonium chloride				0.1803			0.195			0.45				
Zeolite A	15.31	15.31	15.31		4.47	2.01	0.39	1.83	2.58	0.59	1.63	1.63	2	2

(continued)

Granular Laundry Detergent Compositions and Their Components. The present amylase is separately added to these formulations.														
Detergent Composition														
Component Sur-	A	B	C	D	E	F	G	H	I	J	K	L	M	N
factants														
Bentonite				8.35										
Sodium Silicate 1.6.ratio					0.16			4.53	5.62	4.53	4.75	4.75	4.75	4.75
Sodium Silicate 2.0.ratio	3.72	3.72	3.72	8.41			10.1						0.06	0.06
Sodium Silicate 2.35.ratio						7.05								
Citric Acid				0.0066				1.4	1.84	1	1.1	1.1	1.1	1.1
Sodium tripolyphosphate				5.06			5.73							
Sodium Carbonate	26.1	26.18	26.1	15.9	29	12.65	15.93	21	27.31	20.2	23.3	23.3	23.3	23.3
Nonanoyl oxybenzene suphfonate	5.78	5.78	5.78	1.17	1.86		1.73							
Oxaziridinium-based bleach booster	0.037	0.037	0.037					0.0168	0.0333	0.024	0.021	0.021	0.015	0.015
Tetrasodium S, S,-ethylene diaminedisuccinate											0.26	0.26	0.26	0.26
Diethylenetriamine penta (methylene phosphonic acid), heptasodium salt	0.62	0.62	0.62					0.327		0.3272				
Hydroxyethane dimethylene phosphonic acid								0.45	0.2911	0.45	0.47	0.47	0.47	0.47
Ethylene diamine tetraacetate				0.2701			0.28		0.1957					

(continued)

Granular Laundry Detergent Compositions and Their Components. The present amylase is separately added to these formulations.															
	Detergent Composition														
Component Sur- factants	A	B	C	D	E	F	G	H	I	J	K	L	M	N	
MgSO ₄	0.056	0.056	0.056	0.47			0.54	0.79	0.6494	0.793	0.83	0.83	0.82	0.82	
Sodium Percarbonate		7.06	7.06		3.64			19.1	15.85	22.5	19.35	19.35	19.35	19.35	
Tetra Acetyl Ethylene Diamine								4.554	3.71	5.24	4.51	4.51	4.51	4.51	
Sodium Perborate Monohydrate				1.47			5.55								
Carboxymethyl cellulose (e.g. Finnfix BDA ex CPKelco)	0.38	0.38	0.38	0.173		0.62	0.21	0.23	1.07	0.2622	1.01	1.01	1.01	1.01	
Sodium Acrylic acid/ maleic acid co-polymer (70/30)	3.79	3.78	3.79		3.64	0.4	2.61	2.5	2	1.75	1.84	1.84	1.84	1.84	
Sodium polyacrylate (Sokalan PA30 CL)	3.78	3.78	3.78	0.842				0.0055	0.011	0.008	0.007	0.007	0.005	0.005	
Terephthalate polymer									0.231		0.179	0.179	0.179	0.179	
Polyethylene glycol/ vinyl acetate random graft co polymer				0.89		0.55	1.4	0.911	0.8924	0.911	0.96	0.96	0.96	0.96	
Photobleach- zinc phthalocyanine tetrasulfonate															
C.I.Fluorescent Brightener 260	0.1125	0.1125	0.1125	0.043	0.15	0.1174	0.048	0.1455	0.2252	0.1455	0.153	0.153	0.171	0.171	
C.I.Fluorescent Brightener 351 (Tinopal ® CBS)				0.0952			0.1049								

(continued)

Granular Laundry Detergent Compositions and Their Components. The present amylase is separately added to these formulations.														
	Detergent Composition													
Component Sur- factants	A	B	C	D	E	F	G	H	I	J	K	L	M	N
Suds suppressor granule	0.015	0.015	0.015		0.031			0.04	0.0658	0.04	0.042	0.042	0.042	0.042
Hydrophobically modified carboxy methyl cellulose (Finnifix ® SH-1)														
Bentonite														
Miscellaneous (Dyes, perfumes, process aids, moisture and sodium sulphate)	Balance	Balance	Balance	Balance	Balance	Balance	Balance	Balance	Balance	Balance	Balance	Balance	Balance	Balance

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Dishwashing Detergent Gel Compositions					
Ingredients	1	2	3	4	5
	(wt%)	(wt%)	(wt%)	(wt%)	(wt%)
Polytergent ® SLF-18	1	1.3	0.8	1	0.9
Sodium Benzoate (33% active)	0.61	0.61	0.61	0.6	0.6
Xanthan gum	1	0.8	1.2	1	1.1
Sodium Sulphate	10	10	10	8	10
Perfume	0.03	0.05	0.03	0.06	0.1
Sodium Silicate					2
Citric Acid (50% active)	12.5		12		
GLDA		7		8	
Protease 1 (44 mg active/g)	0.7		0.3		
4-Formyl-Phenyl BoronicAcid			0.05		
Protease 2 (10 mg/g) encapsulated		2		0.6	
Protease 3 (48 mg active/g)					0.5
Protease 4 (123 mg active/g)					
Ethanol				0.3	
Potassium Hydroxide (45% active)	14.6	14.6	14.6	14	
Calcium Chloride (25% active)	1.8	1.8	1.8	1.1	0.4
Dye	0.05	0.05	0.05	0.05	0.02
Proxcel GXL ™ (19% active)	0.05	0.05	0.05	0.05	0.05
Acusol ™ 8209	0.34	0.34	0.3	0.35	0.3
Acusol ™ 425N (50% active)	3	3	3.5	2.5	2
Amylases (25 mg/g active)	0.2	0.5	0.4	0.3	0.1
Water & other adjunct ingredients	Balance to 100%	Balance to 100%	Balance to 100%	Balance to 100%	Balance to 100%

Powder Automatic Dishwashing Compositions	
Composition 1	
Ingredients	Wt%
Nonionic surfactant	0.4-2.5%

(continued)

Powder Automatic Dishwashing Compositions	
Composition 1	
Ingredients	Wt%
Sodium metasilicate	0-20%
Sodium disilicate	0-20%
Sodium triphosphate	0-40%
Sodium carbonate	0-20%
Sodium perborate	2-9%
Tetraacetyl ethylene diamine (TAED)	1-4%
Sodium sulfate	5-33%
Enzymes	0.0001-0.1%
Composition 2	
Ingredients	Wt%
Nonionic surfactant (e.g. alcohol ethoxylate)	1-2%
Sodium disilicate	2-30%
Sodium carbonate	10-50%
Sodium phosphonate	0-5%
Trisodium citrate dehydrate	9-30%
Nitrilotrisodium acetate (NTA)	0-20%
Sodium perborate monohydrate	5-10%
Tetraacetyl ethylene diamine (TAED)	1-2%
Polyacrylate polymer (e.g. maleic acid/acrylic acid copolymer)	6-25%
Enzymes	0.0001-0.1%
Perfume	0.1-0.5%
Water	5--10
Composition 3	
Ingredients	Wt%
Nonionic surfactant	0.5-2.0%
Sodium disilicate	25-40%
Sodium citrate	30-55%
Sodium carbonate	0-29%
Sodium bicarbonate	0-20%
Sodium perborate monohydrate	0-15%
Tetraacetyl ethylene diamine (TAED)	0-6%
Maleic acid/acrylic acid copolymer	0-5%
Clay	1-3%
Polyamino acids	0-20%
Sodium polyacrylate	0-8%
Enzymes	0.0001-0.1 %

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Powder Automatic Dishwashing Compositions	
Composition 4	
Ingredients	Wt%
Nonionic surfactant	1-2%
Zeolite MAP	0-42%
Sodium disilicate	0-34%
Sodium citrate	0-12%
Sodium carbonate	0-20%
Sodium perborate monohydrate	7-15%
Tetraacetyl ethylene diamine (TAED)	0-3%
Polymer	0-4%
Maleic acid/acrylic acid copolymer	0-5%
Organic phosphonate	0-4%
Clay	1-2%
Enzymes	0.0001-0.1%
Sodium sulfate	Balance
Composition 5	
Ingredients	Wt%
Nonionic surfactant	1-7%
Sodium disilicate	18-30%
Trisodium citrate	10-24%
Sodium carbonate	12-20%
Monopersulfate (2 KHSO ₅ ·KHSO ₄ ·K ₂ SO ₄)	15-21%
Bleach stabilizer	0.1-2%
Maleic acid/acrylic acid copolymer	0-6%
Diethylene triamine pentaacetate, pentasodium salt	0-2.5%
Enzymes	0.0001-0.1%
Sodium sulfate, water	Balance

Powder and Liquid Dishwashing Composition with Cleaning Surfactant System	
Ingredients	Wt%
Nonionic surfactant	0-1.5%
Octadecyl dimethylamine N-oxide dihydrate	0-5%
80:20 wt C18/C16 blend of octadecyl dimethylamine N-oxide dihydrate and hexadecyldimethyl amine Noxide dehydrate	0-4%
70:30 wt C18/C16 blend of octadecyl bis (hydroxyethyl)amine N-oxide anhydrous and hexadecyl bis (hydroxyethyl)amine N-oxide anhydrous	0-5%
C13-C15 alkyl ethoxysulfate with an average degree of ethoxylation of 3	0-10%
C12-C15 alkyl ethoxysulfate with an average degree of ethoxylation of 3	0-5%

(continued)

Powder and Liquid Dishwashing Composition with Cleaning Surfactant System		
	Ingredients	Wt%
5	C13-C1S ethoxylated alcohol with an average degree of ethoxylation of 12	0-5%
	A blend of C 12-C 1S ethoxylated alcohols with an average degree of ethoxylation of 9	0-6.5%
	A blend of C 13-C 1S ethoxylated alcohols with an average degree of ethoxylation of 30	0-4%
10	Sodium disilicate	0-33%
	Sodium tripolyphosphate	0-46%
	Sodium citrate	0-28%
15	Citric acid	0-29%
	Sodium carbonate	0-20%
	Sodium perborate monohydrate	0-11.5%
	Tetraacetyl ethylene diamine (TAED)	0-4%
20	Maleic acid/acrylic acid copolymer	0-7.5%
	Sodium sulfate	0-12.5%
	Enzymes	0.0001-0.1 %

Non-Aqueous Liquid Automatic Dishwashing Composition		
	Ingredients	Wt%
30	Liquid nonionic surfactant (e.g. alcohol ethoxylates)	2.0-10.0%
	Alkali metal silicate	3.0-15.0%
	Alkali metal phosphate	0-40.0%
	Liquid carrier selected from higher glycols, polyglycols, polyoxides, glycol ethers	25.0-45.0%
35	Stabilizer (e.g. a partial ester of phosphoric acid and a C16-C18 alkanol)	0.5-7.0%
	Foam suppressor (e.g. silicone)	0-1.5%
	Enzymes	0.0001-0.1 %

Non-Aqueous Liquid Dishwashing Composition		
	Ingredients	Wt%
45	Liquid nonionic surfactant (e.g. alcohol ethoxylates)	2.0-10.0%
	Sodium silicate	3.0-15.0%
	Alkali metal carbonate	7.0-20.0%
	Sodium citrate	0.0-1.5%
50	Stabilizing system (e.g. mixtures of finely divided silicone and low molecular weight dialkyl polyglycol ethers)	0.5-7.0%
	Low molecule weight polyacrylate polymer	5.0-15.0%
55	Clay gel thickener (e.g. bentonite)	0.0-10.0%
	Hydroxypropyl cellulose polymer	0.0-0.6%
	Enzymes	0.0001-0.1 %

(continued)

Non-Aqueous Liquid Dishwashing Composition	
Ingredients	Wt%
Liquid carrier selected from higher lycols, polyglycols, polyoxides and glycol ethers	Balance

Thixotropic Liquid Automatic Dishwashing Composition	
Ingredients	Wt%
C 12-C 14 fatty acid	0-0.5%
Block co-polymer surfactant	1.5-15.0%
Sodium citrate	0-12%
Sodium tripolyphosphate	0-15%
Sodium carbonate	0-8%
Aluminium tristearate	0-0.1%
Sodium cumene sulfonate	0-1.7%
Polyacrylate thickener	1.32-2.5%
Sodium polyacrylate	2.4-6.0%
Boric acid	0-4.0%
Sodium formate	0-0.45%
Calcium formate	0-0.2%
Sodium n-decyldiphenyl oxide disulfonate	0-4.0%
Monoethanol amine (MEA)	0-1.86%
Sodium hydroxide (50%)	1.9-9.3%
1,2-Propanediol	0-9.4%
Enzymes	0.0001-0.1%
Suds suppressor, dye, perfumes, water	Balance

Liquid Automatic Dishwashing Composition	
Ingredients	Wt%
Alcohol ethoxylate	0-20%
Fatty acid ester sulfonate	0-30%
Sodium dodecyl sulfate	0-20%
Alkyl polyglycoside	0-21%
Oleic acid	0-10%
Sodium disilicate monohydrate	0-33%
Sodium citrate dihydrate	0-33%
Sodium stearate	0-2.5%
Sodium perborate monohydrate	0-13%
Tetraacetyl ethylene diamine (TAED)	0-8%
Maleic acid/acrylic acid copolymer	4-8%

(continued)

Liquid Automatic Dishwashing Composition	
Ingredients	Wt%
Enzymes	0.0001-0.1%

Liquid Automatic Dishwashing Composition Containing Protected Bleach Particles	
Ingredients	Wt%
Sodium silicate	5-10%
Tetrapotassium pyrophosphate	0-25%
Sodium triphosphate	0-2%
Potassium carbonate	4-8%
Protected bleach particles, e.g. chlorine	5-10%
Polymeric thickener	0.7-1.5%
Potassium hydroxide	0-2%
Enzymes	0.0001-0.1 %
Water	Balance

Compound	Composition of Model Detergent A:		Composition of Model Detergent B:	
	Amount g/ 100 g	% active ingredient	Amount g/ 100 g	% active ingredient
Surfactants				
Na-LAS (92%) (Nacconol90G) (anionic) (linear alkylbenzene sulfonate)	10.87	10	10.87	10
STEOL CS-370E (70%) (anionic), CH ₃ (CH ₂) _m -(OCH ₂ CH ₂) ₃ --OS ₀₃ -, where m~ 11-13	7.14	5	7.14	5
Bio-soft N25-7 (99.5%) (non-ionic),: CH ₃ (CH ₂) _m -(OCH ₂ CH ₂) _h --OH, where and m~ 11-14	5	5	5	5
Oleic acid (fatty acid)	2	2	2	2
Solvents				
H ₂ O	62	65	62	65
Ethanol	0.5	0.5	0.5	0.5
STS (sodium p-toluene sulfonate (40%»	3.75	1.5	3.75	1.5
Mono propylene glycol	2	2	2	2
Builder				
Tri-sodium-citrate	4	4	0	0
Diethylene triamine penta acetic acid (DTPA)	0	0	1.5	1.5
Triethanolamine (TEA)	0.5	0.5	0.5	0.5
Stabilizer				
Boric Acid	1.5	1.5	1.5	1.5

(continued)

Compound	Composition of Model Detergent A:		Composition of Model Detergent B:	
	Amount g/ 100 g	% active ingredient	Amount g/ 100 g	% active ingredient
Surfactants				
Minors				
10N NaOH (for adjustment to pH 8.5)	0.8	0.8	0.8	0.8

Liquid Detergent and Cleaning Agent Compositions								
Ingredients	E1	E2	E3	C1	C2	C3	C4	C5
Gellan gum	0.2	0.2	0.15	0.15				
Xanthan gum			0.15		0.15	0.5	0.2	
Polyacrylate (Carbopol Aqua 30)	0.4	0.4					0.6	0.6
C ₁₂₋₁₄ -fatty alcohol with 7 EO	22	10	10	10	10	10	10	10
C ₉₋₁₃ -alkyl benzenesulfonate, Na salt		10	10	10	10	10	10	10
C ₁₂₋₁₄ -alkylpolyglycoside	1							
Citric acid	1.6	3	3	3	3	3	3	3
Dequest [®] 2010 Hydroxyethylidene-1,1-diphosphonic acid, tetrasodium salt (from Solutia)	0.5	1	1	1	1	1	1	1
Sodium lauryl ether sulfate with 2 EO	10	5	5	5	5	5	5	5
Monoethanolamine	3	3	3	3	3	3	3	3
C ₁₂₋₁₈ -fatty acid	7.5	7.5	7.5	7.5	7.5	7.5	7.5	7.5
Propylene glycol		6.5	6.5	6.5	6.5	6.5	6.5	6.5
Sodium cumene sulfonate		2	2	2	2	2	2	2
Enzymes, dyes, stabilizers	+	+	+	+	+	+	+	+
Microcapsules with about 2000 µm diameter	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
Water	To 100	To 100	To 100	To 100	To 100	To 100	To 100	To 100
Flow limit (Pas)	0.58	1.16	1.16	no	no	no	yes	no

All purpose Alkaline detergent Compositions (all-purpose. glass. kitchen) Hard surface cleaning detergent composition				
Composition [% by wt.]	E1	E2	E3	E4
Fatty alcohol ethoxylate C12-7EO	1	3	5	0.5
Alkylbenzenesulfonic acid Na salt	3	1	2	4
Octyl sulfate	3	2	2	2
Sodium carbonate	1.5	0.5	1.0	1.5
Citric acid	0.5	0.5	0.5	0.5

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All purpose Alkaline detergent Compositions (all-purpose. glass. kitchen) Hard surface cleaning detergent composition

Composition [% by wt.]	E1	E2	E3	E4
Fatty acid	0.5	0.5	0.5	1.0
Ethanol	5	3	5	3
Perfume	0.2	0.2	0.2	0.2
Water	To 100	To 100	To 100	To 100

Acidic Detergent Compositions (bath, toilet)

Composition [% by wt.]	E5	E6	E7	E8
Fatty alcohol ether sulfate C12-2EO sodium salt	2	3	5	2
Ethanol	3	3	3	3
Citric acid	3	10	3	10
Thickener xanthan Kelzan ASX -T		0.05		0.05
Perfume	0.1	0.1	0.1	0.1
Water	To 100	To 100	To 100	To 100

Cleaning Paste Composition

Composition [% by wt.]	E9
C 12 Fatty alcohol sulfate	20
C16-18 Fatty alcohol ethoxylate 25 EO	20
C 12-18 Fatty acid monoethanolamide	10
Sodium sulfate	40
Sodium carbonate	5
Cellulose	4.899
Dye	0.001
Perfume	0.1

Self Foaming Cleaning Powder Composition

Composition [% by wt.]	E10
C 12 Fatty alcohol sulfate	2
Sodium sulfate	37.899
Sodium carbonate	25
Citric Acid	35
Dye	0.001
Perfume	0.1

Compositions of a Clear Aqueous Detergent and Cleaning Agent having a flow limit						
Ingredients	V1	E1	E2	E3	E4	E5
1,2 Propane diol	8	0	2	6	4	2
Dipropylene glycol	0	8	6	2	4	2
Polyacrylate (Carbopol Aqua 30)	3	3	3	3	3	
Polyacrylate (Polygel W301)	-	-	-	-	-	1.8
C ₁₂₋₁₄ -fatty alcohol with 7 EO	10	10	10	10	10	10
C ₉₋₁₃ -alkyl benzenesulfonate, Na salt	10	10	10	10	10	-
Citric Acid	3	3	3	3	3	2
Dequest® 2010 Hydroxyethylidene-1,1-diphosphonic acid, tetrasodium salt (ex Solutia)	1	1	1	1	1	-
Dequest® 2066 Diethylene triamine penta (methylenephosphonic acid) hepta Na salt (ex Solutia)	-	-	-	-	-	0.7
Sodium lauryl ether sulfate with 2 EO	10	10	10	10	10	5
Monoethanolamine	3	3	3	3	3	2
C ₁₂₋₁₈ -fatty acid Na salt	5.5	5.5	5.5	5.5	5.5	5.5
Enzymes, dyes, stabilizers	+	+	+	+	+	+
Microcapsules with about 2000 µm diameter	0.5	0.5	0.5	0.5	0.5	0.5
Water	To 100	To 100	To 100	To 100	To 100	To 100
Flow limit (Pas)	0.4	0.6	0.6	0.8	1.0	0.6
Appearance	Cloudy	Clear	Clear	Clear	Clear	Clear

Liquid Laundry Detergent	
Ingredients	Wt%
ABS (alkyl benzenesulphonate)	10
FAEOS	5
C _{12/14} 7EO	10
C _{12/18} Fatty Acid	5
Glycerol	5
Sodium citrate	3
Protease/Amylase/Cellulase	1
Tinopal® DMS-X (optical brightener manufactured by Ciba)	0.2
Water	To 100

Granular Laundry Detergent	
Ingredients	Wt%
ABS (alkyl benzenesulphonate)	11
C _{13/15} 7EO	3

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Granular Laundry Detergent	
Ingredients	Wt%
Sodium carbonate	20
Sodium hydrogencarbonate	5
Sodium sulphate	25
Sodium silicate	5
Sodium percarbonate	13
TAED	5
Sodium polyacrylate	4.5
Enzymes (protease, amylase, and cellulose)	3.5
Water	To 100

Aqueous Liquid Washing Product Formulations (without-FWM1 and with-FWM2 0.5% hyperbranched polyesteramide)		
Formulation	FWM1	FWM2
C ₁₂₋₁₄ -fatty alcohol with 2 EO	5	5
LAS	10	10
C ₁₂₋₁₈ -fatty alcohol with 7 EO	10	10
C ₁₂₋₁₈ soap	8	8
Citrate	4	4
1,2-propanediol	5	5
Hybrane® SIP 2100 (manufactured by DSM)		0.5

Liquid Laundry Detergent Compositions			
Detergent Composition	Wt%		
	E1	E2	E3
C ₁₂₋₁₄ fatty alcohol with 7 EO	5	4	10
C ₉₋₁₃ alkylbenzene sulfonate, Na salt	10	10	10
Sodium lauryl ether sulfate with 2 EO	-	-	8
Active substance (specific polycarbonate-, polyurethane-, and/or polyureapolyorganosiloxane compounds or precursor compounds thereof of the reactive cyclic carbonate and urea type	1	1	1
Polyacrylate thickener	-	-	1
Sodium percarbonate	15	18	-
TAED	3	3	-
C ₁₂₋₁₈ fatty acid, Na salt	1	1.5	7.5
PVA/Maleic acid copolymer	4.5	2	-
Citric acid, Na salt	2.5	-	2
Phosphonic acid, Na salt	0.5	0.5	1

(continued)

Liquid Laundry Detergent Compositions				
Detergent Composition		Wt%		
Sodium carbonate	10	20	-	
Propane diol	-	-	6.5	
Zeolite A	25	25	-	
Boric Acid Sodium salt	-	-	1.2	
Silicone defoamer	2.5	1.3	0.1	
Enzymes (protease, amylase, cellulase)	+	+	+	
Colorant	+	+	+	
Perfume	0.5	0.2	0.8	
Water	-	-	To 100	
Sodium sulfate	-	To 100	-	
Sodium bicarbonate	To 100	-	-	

Example formulations of preferred phosphate-free automatic dishwashing agents				
Ingredient	Formulation 1 (wt%)	Formulation 2 (wt%)	Formulation 3 (wt%)	Formulation 4 (wt%)
Citrate	5 to 60	10 to 55	15 to 50	15 to 50
Sodium percarbonate	1 to 20	2 to 15	4 to 10	4 to 10
Bleach catalyst	0.01 to 3	0.02 to 2	0.02 to 2	0.02 to 1
Copolymer ¹	0.1 to 30	0.5 to 25	1.0 to 20	1.0 to 20
Nonionic surfactant ²	1 to 10	2 to 8	2 to 8	3 to 6
Misc	To 100	To 100	To 100	To 100

Example formulations of preferred phosphate-free automatic dishwashing agents				
Ingredient	Formulation 5 (wt%)	Formulation 6 (wt%)	Formulation 7 (wt%)	Formulation 8 (wt%)
Citrate	5 to 60	10 to 55	15 to 50	15 to 50
Sodium percarbonate	1 to 20	2 to 15	4 to 10	4 to 10
Phosphonate	2 to 8	2 to 8	2 to 8	2 to 8
Copolymer ¹	0.1 to 30	0.5 to 25	1.0 to 20	1.0 to 20
Nonionic surfactant ²	1 to 10	2 to 8	2 to 8	3 to 6
Misc	To 100	To 100	To 100	To 100

Example formulations of preferred phosphate-free automatic dishwashing agents				
Ingredient	Formulation 9 (wt%)	Formulation 10 (wt%)	Formulation 11 (wt%)	Formulation 12 (wt%)
Citrate	5 to 60	10 to 55	15 to 50	15 to 50
Sodium percarbonate	1 to 20	2 to 15	4 to 10	4 to 10
Enzyme	0.1 to 6	0.2 to 5	0.4 to 5	0.4 to 5
Copolymer ¹	0.1 to 30	0.5 to 25	1.0 to 20	1.0 to 20
Nonionic surfactant ²	1 to 10	2 to 8	2 to 8	3 to 6
Misc	To 100	To 100	To 100	To 100

Example formulations of preferred phosphate-free automatic dishwashing agents				
Ingredient	Formulation 13 (wt%)	Formulation 14 (wt%)	Formulation 15 (wt%)	Formulation 16 (wt%)
Citrate	5 to 60	10 to 55	15 to 50	15 to 50
Carbonate/hydrogen carbonate	2 to 40	2 to 40	2 to 40	2 to 40
Silicate	0 to 15	0 to 15	0 to 15	0.1 to 10
Phosphonate	0 to 14	0 to 14	0 to 14	2 to 8
Sodium percarbonate	1 to 20	2 to 15	4 to 10	4 to 10
Bleach catalyst	0.01 to 3	0.02 to 2	0.02 to 2	0.02 to 1
Copolymer ¹	0.1 to 30	0.5 to 25	1.0 to 20	1.0 to 20
Nonionic surfactant ²	1 to 10	2 to 8	2 to 8	3 to 6
Enzyme	0.1 to 6	0.2 to 5	0.4 to 5	0.4 to 5
Misc	To 100	To 100	To 100	To 100

¹Copolymer comprising i) monomers from the group of mono- or polyunsaturated carboxylic acids ii) monomers of the general formula $R^1(R^2)C=C(R^3)-X-R^4$, in which R^1 to R^3 mutually independently denote -H, -CH₃ or -C₂H₅, X denotes an optionally present spacer group which is selected from -CH₂-, -C(O)O- and -C(O)-NH-, and R^4 denotes a straight chain or branched saturated alkyl residue with 2 to 22 carbon atoms or denotes an unsaturated, preferably aromatic residue with 6 to 22 carbon atoms iii) optionally further monomers

²Nonionic surfactant of the general formula $R^1-CH(OH)CH_2O-(AO)_w-(A'O)_x-(A''O)_y-(A'''O)_z-R_2$, in which R^1 denotes a straight-chain or branched, saturated or mono- or polyunsaturated C₆-24 alkyl or alkenyl residue; R_2 denotes a linear or branched hydrocarbon residue with 2 to 26 carbon atoms; A, A', A'' and A''' mutually independently denote a residue from the group comprising ---CH₂CH₂-, -CH₂CH₂---CH₂-, ---CH₂CH₂--CH(CH₃), CH₂-CH₂-CH₂CH₂-, -CH₂-CH-(CH₃)-CH₂-, -CH₂-CH(CH₂-CH₃), w, x, y and z denote values between 0.5 and 120, wherein x, y and/or z may also be 0.

Composition of phosphate-free automatic dishwashing detergents		
Raw material	V1	E1
Citrate	23	23
MGDA	8	8
Copolymer ¹	12	12

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(continued)

Composition of phosphate-free automatic dishwashing detergents		
Raw material	V1	E1
HEDP	2	2
Soda	28	28
Sodium percarbonate	10	10
TAED	2.4	2.4
Protease	2	2
Amylase	1.8	1.8
Non-ionic surfactant ²	5	-
Non-ionic surfactant ³	-	5
Misc	To 100	To 100

Textile Washing Agent	
Ingredient	wt % pure substance
Xanthan	0.3-0.5
Anti foaming agent	0.2-0.4
Glycerol	6-7
Ethanol	0.3-0.5
FAEOS	4-7
Non ionic surfactant (FAEO, APG among others)	24-28
Boric acid	1
Sodium citrate dihydrate	1-2
Soda	2-4
Coconut fatty acids	14-16
HEDP	0.5
PVP	0-0.4
Optical brightener	0-0.05
Dye	0-0.001
Perfume	0-2
Water demineralized	remainder

Example detergent compositions for application to a substrate					
Ingredients	Weight Percent (actives %)				
	D1	D2	D3	D4	D5
Sodium dodecyl benzene sulfonate	26.09	17.30	15.60	17.70	16.70
Sodium alkyl C ₁₄₋₁₅ / 7EO ether sulfate	13.80	-	-	-	-
Linear alcohol ethoxylate C ₁₄₋₁₅ / 7EO	13.44	5.4	14.6	5.5	5.2
Polyethylene glycol PEG 75	2	1.4	1.3	1.4	1.4

(continued)

Example detergent compositions for application to a substrate					
	Weight Percent (actives %)				
Ingredients	D1	D2	D3	D4	D5
Polyoxyethylene (100) stearyl ether	21.99	15.6	14.1	15.9	15.1
Sodium silicate SiO ₂ /Na ₂ O ratio 1.6-1.8	3.72	16.6	15	17	16
Sodium Silicate (Britesil [®] C24)	7	-	-	-	-
Sodium Carbonate	-	6.5	5.9	6.7	6.3
Sodium tetraborate decahydrate	-	11.9	10.8	12.2	11.5
Sodium polyacrylate ~4500 MW	-	1.8	1.7	-	5.2
EDTA-tetrasodium salt	-	0.1	0.1	0.1	0.1
Optical brightener (Tinopal [®] CBS-X)	0.15	0.1	0.09	0.1	0.1
Dyes and fragrances	0.9	0.9	0.81	1.01	0.91
Water	10.92	22.10	19.90	22.4	21.5

Example fabric conditioning compositions for application to a substrate					
	Weight Percent (actives %)				
Ingredients	FS1	FS2	FS3	FS4	FS5
Di-(hydrogenated tallow) dimethyl ammonium methyl sulfate	33.6	33.2	44.4	22.2	33.2
Unsaturated trialkylglycerides	16.8	16.6	22.2	11.1	16.6
Hydrogenated tallow fatty acid	16.8	16.6	22.2	11.1	16.6
C ₁₂₋₁₈ coco fatty acid	11.2	11.1	-	11.1	-
C ₁₂₋₁₈ fatty alcohol ethoxylate (7EO)	11.2	11.1	-	-	16.6
Fragrance oil	10.4	11.4	11.2	11.2	17

Exemplary Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Citrate	12-50	15-40	12-50	15-40
Dicarboxylic acid	1-18	1-18	2-16	4-12
Phosphate	-	-	-	-
Bleaching Agent	-	-	-	-
Misc	To 100	To 100	To 100	To 100

Additional Exemplary Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Citrate	12-50	15-40	12-50	15-40
Dicarboxylic acid	1-18	1-18	2-16	4-16
Carbonate	5-50	10-40	5-50	10-40
Phosphate	-	-	-	-
Bleaching Agent	-	-	-	-
Misc	To 100	To 100	To 100	To 100

Additional Exemplary Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Citrate	12-50	15-40	12-50	15-40
Dicarboxylic acid	1-18	1-18	2-16	4-12
Carbonate	5-50	10-30	5-50	10-30
Phosphonate	1-8	1-8	1.2-6	1.2-6
Phosphate	-	-	-	-
Bleaching Agent	-	-	-	-
Misc	To 100	To 100	To 100	To 100

Preferred Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Citrate	12-50	15-40	12-50	15-40
Dicarboxylic acid	1-18	1-18	2-16	4-12
Carbonate	0-50	0-30	0-30	0-30
Phosphonate	0-8	0-8	0-8	0-8
Phosphate	-	-	-	-
Bleaching Agent	-	-	-	-
Misc	To 100	To 100	To 100	To 100

Additional Preferred Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Citrate	12-50	15-40	12-50	15-40
Maleic acid	1-18	1-18	2-16	4-12
Carbonate	5-50	10-30	5-50	10-30

(continued)

Additional Preferred Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Phosphonate	1-8	1-8	1.2-6	1.2-6
Phosphate	-	-	-	-
Bleaching Agent	-	-	-	-
Misc	To 100	To 100	To 100	To 100

Preferred Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Citrate	12-50	15-40	12-50	15-40
Dicarboxylic acid	1-18	1-18	2-16	4-12
Carbonate	0-50	0-30	0-30	0-30
Phosphonate	0-8	0-8	0-8	0-8
Non-ionic surfactant	0.1-15	0.1-15	0.5-8	0.5-8
Phosphate	-	-	-	-
Bleaching Agent	-	-	-	-
Misc	To 100	To 100	To 100	To 100

Additional Preferred Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Citrate	12-50	15-40	12-50	15-40
Maleic acid	1-18	1-18	2-16	4-12
Carbonate	5-50	10-30	5-50	10-30
Phosphonate	1-8	1-8	1.2-6	1.2-6
Non-ionic surfactant	0.1-15	0.1-15	0.5-8	0.5-8
Phosphate	-	-	-	-
Bleaching Agent	-	-	-	-
Misc	To 100	To 100	To 100	To 100

Preferred Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Citrate	12-50	15-40	12-50	15-40
Dicarboxylic acid	1-18	1-18	2-16	4-12

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(continued)

Preferred Automatic Dishwashing Agents				
Ingredient	Wt %			
Carbonate	0-50	0-30	0-30	0-30
Phosphonate	0-8	0-8	0-8	0-8
Sulfo copolymer	0-20	0-20	0-20	0-20
Non-ionic surfactant	0-15	0-15	0-8	0-8
Enzyme preparations	0.1-12	0.1-12	0.5-8	0.5-8
Phosphate	-	-	-	-
Bleaching Agent	-	-	-	-
Misc	To 100	To 100	To 100	To 100

Additional Preferred Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Citrate	12-50	15-40	12-50	15-40
Maleic acid	1-18	1-18	2-16	4-12
Carbonate	5-50	10-30	5-50	10-30
Phosphonate	1-8	1-8	1.2-6	1.2-6
Sulfo copolymer	0-20	0-20	0-20	0-20
Non-ionic surfactant	0.1-15	0.1-15	0.5-8	0.5-8
Enzyme preparations	0.1-12	0.1-12	0.5-8	0.5-8
Phosphate	-	-	-	-
Bleaching Agent	-	-	-	-
Misc	To 100	To 100	To 100	To 100

Preferred Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Citrate	12-50	15-40	12-50	15-40
Dicarboxylic acid	1-18	1-18	2-16	4-12
Carbonate	0-50	0-30	0-30	0-30
Phosphonate	0-8	0-8	0-8	0-8
Sulfo copolymer	0-20	0-20	0-20	0-20
Non-ionic surfactant	0-15	0-15	0-8	0-8
Enzyme preparations	0-12	0-12	0-8	0-8
Organic Solvent	0.1-15	0.5-8	0.1-15	0.5-8
Phosphate	-	-	-	-
Bleaching Agent	-	-	-	-

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(continued)

Preferred Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Misc	To 100	To 100	To 100	To 100

Additional Preferred Automatic Dishwashing Agents				
Ingredient	Wt %			
	Formula 1	Formula 2	Formula 3	Formula 4
Citrate	12-50	15-40	12-50	15-40
Dicarboxylic acid	1-18	1-18	2-16	4-12
Carbonate	5-50	10-30	5-50	10-30
Phosphonate	1-8	1-8	1.2-6	1.2-6
Sulfo copolymer	0-20	0-20	0-20	0-20
Non-ionic surfactant	0.1-15	0.1-15	0.5-8	0.5-8
Enzyme preparations	0.1-12	0.1-12	0.5-8	0.5-8
Organic Solvent	0.1-15	0.5-8	0.1-15	0.5-8
Phosphate	-	-	-	-
Bleaching Agent	-	-	-	-
Misc	To 100	To 100	To 100	To 100

Automatic Dishwashing Agents		
Ingredient	Wt %	
	C 1	E 1
Sodium citrate	9	9
Potassium hydroxide	7	7
Sodium carbonate	14	14
Maleic acid	-	1
Sulfo polymer	4.2	4.2
HEDP	1.5	1.5
Non-ionic surfactant	2	2
Protease preparation	2	2
Amylase preparation	0.8	0.8
Alkanolamine	1.5	1.5
Thickener	2	2
Water, misc	To 100	To 100

Manual Dishwashing Agents							
Ingredient	Wt %						
	Invention 1	Invention 2	Invention 3	Invention 4	Invention 5	Invention 6	Invention 7
Fatty alcohol ether sulfate	10	13.33	12	12	13.3	13.3	13.3
Cocamidopropylbetaine	2.5	3.33	3.1	3.1	3	3	3
Sce. Alkanesulfonate	2.5	3.33	2.9	2.9	3.7	3.7	3.7
Fatty alcohol ethoxylate	9	6	-	-	-	-	-
Sodium chloride	24	24	22	24	20	24	20
Ethanol	-	-	2	2	2.5	2.5	4
Perfume	0.2	0.3	0.3	0.3	0.3	0.3	0.3
Colorant	0.2	0.2	0.2	0.2	0.2	0.2	0.2
Water	51.60	49.51	57.5	55.5	57	53	55.5

Antibacterially active detergent/cleaning agent						
Ingredient	V1	E1	E2	E3	E4	E5
C ₁₂₋₁₈ fatty alcohol with 7EO	12	12	12	5	5	-
N-cocoalkyl N, N dimethylamine oxide	1.95	1.95	1.95	2	2	-
Esterquat (N-methyl-N-(2 hydroxyethyl)-N-N-(ditallowacyloxy ethyl) ammonium methosulfate	-	-	-	-	-	15
AgNO ₃ ·H ₂ O	0.0043	0.0043	0.0043	0.004	0.004	0.004
C14 fatty acid	5	5	-	-	-	-
Farnesol	0.02	0.02	0.02	0.02	0.02	0.02
Coco Fatty acid	2.5	2.5	2.5	12	-	-
Citric Acid	-	-	-	1.0	0.1	-
H ₂ O ₂	-	0.5	0.035	2	5	0.5
NaOH	0.35	0.35	0.35	1.9	-	-
NH ₄ OH	0.04	0.04	0.04	0.06	-	-
2-Propanol	-	-	-	-	-	1.67
MgCl ₂ x 6H ₂ O	-	-	-	-	-	0.01
Perfume A	1.00	1.00	1.00	1.00	1.00	0.75
Water	To 100	To 100	To 100	To 100	To 100	To 100
pH	8.5	8.5	8.5	8.5	5.5	2.6

Detergent containing anti-grey agent	
Ingredients	M1 (wt%)
C ₉₋₁₃ alkylbenzenesulfonate sodium salt	10
Sodium lauryl ether sulfate with 2EO	5
C ₁₂₋₁₈ fatty alcohol with 7EO	10
C ₁₂₋₁₄ alkyl polyglycoside	2

(continued)

Detergent containing anti-grey agent	
Ingredients	M1 (wt%)
C ₁₂₋₁₈ fatty acid sodium salt	8
Glycerol	5
Trisodium citrate	1
Polyacrylate	2
Active ingredient (anti-grey agent-a polycarbonate-, polyurethane-, and/or polyurea-polyorganosiloxane compound or a precursor compound use in the production thereof)	1
Enzyme, dye, optical brightener	+
Water	To 100

Example detergent compositions for application to a substrate					
Ingredients	Weight Percent (actives %)				
	D1	D2	D3	D4	D5
Sodium dodecyl benzene sulfonate	26.09	17.30	15.60	17.70	27.00
Sodium alkyl C ₁₄₋₁₅ /7EO ether sulfate	13.80				14.00
Linear alcohol ethoxylate C ₁₄₋₁₅ /7EO	13.44	5.40	14.60	5.50	14.00
Linear alcohol ethoxylate C ₁₂₋₂₀ /7EO					23.00
Polyethylene Glycol PEG-75	2.00	1.40	1.30	1.40	2.00
Polyoxyethylene (100) stearyl ether	21.99	15.60	14.10	15.90	
Sodium Silicate SiO ₂ /Na ₂ O ratio 1.6-1.8	3.72	16.60	15.00	17.00	
Sodium Silicate (Britesil® C24)	7.00				11.00
Sodium Carbonate		6.50	5.90	6.70	
Sodium tetraborate decahydrate		11.90	10.80	12.20	
Sodium polyacrylate -4,500 MW		1.80	1.70		
EDTA - tetrasodium salt		0.10	0.10	0.10	
Optical brightener (Tinopal® CBS-X)	0.15	0.10	0.09	0.10	0.20
Dyes and fragrances	0.90	0.90	0.81	1.01	0.35
Water	10.92	22.10	19.90	22.40	9.55

Example enzyme containing compositions for application to a substrate					
Ingredients	Weight Percent (actives %)				
	E1	E2	E3	E4	E5
Polyethylene Glycol PEG-75	98.60	99.10			
Fatty acid based matrix 1			98.9		99.10
Fatty acid based matrix 2				98.80	
Protease	0.10	0.10	0.12	0.10	0.10
Mannanase	0.02		0.02	0.02	

(continued)

Example enzyme containing compositions for application to a substrate					
Ingredients	Weight Percent (actives %)				
	E1	E2	E3	E4	E5
Amylase	0.12	0.25	0.1	0.12	0.25
Cellulase	0.08		0.1	0.08	
Lipase	0.08			0.08	
Pectate Lyase				0.05	
Enzyme Stabilizers	1.00	0.55	0.75	0.75	0.55
Fatty acid based matrix 1 is comprised of 20 wt. % of the sodium salt of coconut fatty acid, 50 wt. % of non polymeric polyols (sorbitol, glycerin, propylene glycol, sucrose and glucose), 15 wt.% of anionic and nonionic surfactants, and 15 wt. % of water.					
Fatty acid based matrix 2 is comprised of 20 wt.% of the sodium salt of stearic acid, 3 wt.% of the sodium salt of lauric acid, 3 wt.% of the sodium salt of myristic acid, 50 wt.% of non polymeric polyols (sorbitol, glycerin, and propylene glycol), 2 wt.% of lauric acid, 2 wt.% of stearic acid, 10 wt.% of anionic surfactant, and 10 wt.% of water.					

Table 1 Detergent Composition	
Ingredients	(% by weight)
Soap (saturated C ₁₂₋₂₄ fatty acid soaps and oleic acid soap)	5.42
Sodium C ₁₂₋₁₄ alkyl benzenesulfonate	22.67
Sodium C ₁₄₋₁₆ fatty alcohol sulfate	4.59
C ₁₂₋₁₈ fatty alcohol.5EO	0.81
Sodium carbonate	4.55
Zeolite A	29.86
Sodium silicate	8.00
Acrylic acid/maleic acid copolymer	16.16
Opt. brightener	0.45
Phosphonate	2.30
NaOH, 50%	0.63
Water	3.88
Other salts	0.68
Table 2	
Detergent composition	59.5%
Coated bleaching agent (Na percarbonate)	23.3%
Coated bleach activator (TAED)	7%
Citric acid monohydrate	10.2%

Particulate detergent composition	
Ingredient	% wt
sodium dodecylbenzenesulphonate	8.5

(continued)

Particulate detergent composition	
Ingredient	% wt
c12-C15 primary alcohol, condensed with 7 moles of ethylene oxide	4
sodium-hardened rapeseed oil soap	1.5
sodium triphosphate	33
sodium carbonate	5
sodium silicate	6
sodium sulphate	20
water	9
fluorescers, soil-suspending agents, dyes, perfumes	minor amounts
sodium perborate	12
tetraacetyl ethylene diamine (TAED) (granules)	2
proteolytic enzyme (Savinase ex.Novo)	0.4

Detergent composition A

9 % anionic detergent
1 % nonionic detergent
21.5 % sodium tripolyphosphate
7 % sodium perborate
0.6 % Savinase (a proteolytic enzyme)
balance sodium sulphate + minor ingredients

Detergent composition B

9 % anionic detergent
4 % nonionic detergent
28% zeolite
4.5% nitrilotriacetate
5.5% sodium perborate
3.5% tetraacetylenediamine
0.5% Savinase
balance sodium sulphate + minor ingredients

Detergent composition C

5 % anionic detergent
4 % nonionic detergent
1 % soap
30 % zeolite

(continued)

Detergent composition C

3. % copolymer of acrylic acid with maleic anhydride

7.5% sodium perborate

3 % tetraacetylenediamine

balance sodium sulphate + minor ingredients

Detergent composition D

8 % anionic synthetic detergent

4 % nonionic synthetic detergent

4 % soap

35. % sodium carbonate

20 % powdered calcite

6 % sodium perborate

2 % tetraacetylenediamine

0.5% Savinase

balance sodium sulphate + minor ingredients

Laundry detergent composition

Ingredients	Parts by weight
Sodium dodecyl benzene sulphonate	8.5
C12-C15 primary alcohol, condensed with 7 moles of ethylene oxide	4
Sodium-hardened rapeseed oil soap	1.5
Sodium triphosphate	33
Sodium carbonate	5
Sodium silicate	6
Sodium sulphate	20
Water	9
Fluorescers, soil-suspending agents, dyes, perfumes	minor amount
Sodium perborate	12
Tetraacetyl ethylene diamine (TAED) (granules)	2
Proteolytic enzyme (Savinase ex NOVO)	0.4

Laundry detergent compositions

	A	B	C	D
sodium dodecylbenzene sulphonate	9	9	9	9
C13-C15 linear primary alcohol, condensed with 7 moles of ethylene oxide (e.g. Synperonic A7)	1	4	4	1

(continued)

Laundry detergent compositions				
	A	B	C	D
C13-C15 linear primary alcohol, condensed with 3 moles of ethylene oxide (e.g. Synperonic A3)	3	0	0	3
sodium tripolyphosphate	23	23	0	0
zeolite type 4A	0	0	24	24
copolymer of acrylic acid with maleic anhydride			4	4
sodium polyacrylate	2	2	0	0
alkaline silicate	5	5		
fluorescer	0.25	0.25	0.16	0.16
EDTA	0.15	0.15	0.18	0.18
SCMC	0.5	0.5	0.55	0.55
salt	2	2		
sodium sulphate	26.8	26.8	22.31	22.31
sodium carbonate	0	0	10.3	10.3
moisture	10	10	11	11
TAED	3	3	3.3	3.3
sodium perborate monohydrate	10	10	8	8
calcium Dequest® 2047	0.7	0.7	0.3	0.3
foam depressor	3	3	2.5	2.5
perfume	0.2	0.2	0	0
alkaline protease (Savinase (A) 6T)	0.4	0.4	0.4	0.4

Detergent composition				
Ingredients	Ex. 1	Ex. 2	Ex.3	Ex.4
Material	Level (parts as is)	Level (parts as is)	Level (parts as is)	Level (parts as is)
Glycerol	3.17	3.17	3.17	3.17
MPG	5.7	5.7	5.7	5.7
NaOH	2.13	2.13	2.13	2.13
TEA	2.05	2.05	2.05	2.05
Neodol 25-7	12.74	12.74	12.74	12.74
F-Dye	0.18	0.18	0.18	0.18
Citric Acid	1.71	1.71	1.71	1.71
LAS (as LAS Acid)	8.49	8.49	8.49	8.49
Fatty acid	3.03	3.03	3.03	3.03
Empigen BB	1.5	1.5	1.5	1.5
SLES	4.24	4.24	4.24	4.24

(continued)

Detergent composition				
Ingredients	Ex. 1	Ex. 2	Ex.3	Ex.4
Material	Level (parts as is)	Level (parts as is)	Level (parts as is)	Level (parts as is)
Dequest 2066	0.875	0.875	0.875	0.875
Patent Blue	0.00036	0.00036	0.00036	0.00036
Acid Yellow	0.00005	0.00005	0.00005	0.00005
Opacifier	0.0512	0.0512	0.0512	0.0512
Perfume	0.734	0.734	0.734	0.734
Borax	10	10	10	10
Savinase	2.362	2.362	2.362	2.362
Stainzyme	0.945	0.945	0.945	0.945
Soap	3.03	3.03	3.03	3.03
EPEI 20E0 (ex Nippon Shokubai) polyethyleneimine having a weight average molecular weight of about 600, and wherein the polyethyleneimine has been modified by alkoxylation with an average 20 ethylene oxide moieties	5.5	5.5	5.5	9
Lipex® (ex Novozymes)	3	3	3	3
Texcare SRN170 (ex Clariant) soil release polymer	0	7.5	0	0
Sokolan CP5 (ex BASF) Soil-release polymer	0	0	20	0

7.6. Methods of Assessing Amylase Activity in Detergent Compositions

[0275] Numerous α -amylase cleaning assays are known in the art, including swatch and micro-swatch assays. The appended Examples describe only a few such assays.

[0276] In order to further illustrate the compositions and methods, and advantages thereof, the following specific examples are given with the understanding that they are illustrative rather than limiting.

8. Brewing Compositions

[0277] The present variant amylase may be a component of a brewing composition used in a process of brewing, *i.e.*, making a fermented malt beverage. Non-fermentable carbohydrates form the majority of the dissolved solids in the final beer. This residue remains because of the inability of malt amylases to hydrolyze the alpha-1,6-linkages of the starch. The non-fermentable carbohydrates contribute about 50 calories per 12 ounces of beer. an amylase, in combination with a glucoamylase and optionally a pullulanase and/or isoamylase, assist in converting the starch into dextrins and fermentable sugars, lowering the residual non-fermentable carbohydrates in the final beer.

[0278] The principal raw materials used in making these beverages are water, hops and malt. In addition, adjuncts such as common corn grits, refined corn grits, brewer's milled yeast, rice, sorghum, refined corn starch, barley, barley starch, dehusked barley, wheat, wheat starch, torrifed cereal, cereal flakes, rye, oats, potato, tapioca, and syrups, such as corn syrup, sugar cane syrup, inverted sugar syrup, barley and/or wheat syrups, and the like may be used as a source of starch.

[0279] For a number of reasons, the malt, which is produced principally from selected varieties of barley, has the greatest effect on the overall character and quality of the beer. First, the malt is the primary flavoring agent in beer. Second, the malt provides the major portion of the fermentable sugar. Third, the malt provides the proteins, which will contribute to the body and foam character of the beer. Fourth, the malt provides the necessary enzymatic activity during mashing. Hops also contribute significantly to beer quality, including flavoring. In particular, hops (or hops constituents) add desirable bittering substances to the beer. In addition, the hops act as protein precipitants, establish preservative agents and aid in foam formation and stabilization.

[0280] Grains, such as barley, oats, wheat, as well as plant components, such as corn, hops, and rice, also are used for brewing, both in industry and for home brewing. The components used in brewing may be unmalted or may be malted, *i.e.*, partially germinated, resulting in an increase in the levels of enzymes, including α -amylase. For successful brewing, adequate levels of α -amylase enzyme activity are necessary to ensure the appropriate levels of sugars for fermentation.

an amylase, by itself or in combination with another α -amylase(s), accordingly may be added to the components used for brewing.

[0281] As used herein, the term "stock" means grains and plant components that are crushed or broken. For example, barley used in beer production is a grain that has been coarsely ground or crushed to yield a consistency appropriate for producing a mash for fermentation. As used herein, the term "stock" includes any of the aforementioned types of plants and grains in crushed or coarsely ground forms. The methods described herein may be used to determine α -amylase activity levels in both flours and stock.

[0282] Processes for making beer are well known in the art. See, *e.g.*, Wolfgang Kunze (2004) "Technology Brewing and Malting," Research and Teaching Institute of Brewing, Berlin (VLB) 3rd edition. Briefly, the process involves: (a) preparing a mash, (b) filtering the mash to prepare a wort, and (c) fermenting the wort to obtain a fermented beverage, such as beer. Typically, milled or crushed malt is mixed with water and held for a period of time under controlled temperatures to permit the enzymes present in the malt to convert the starch present in the malt into fermentable sugars. The mash is then transferred to a mash filter where the liquid is separated from the grain residue. This sweet liquid is called "wort," and the left over grain residue is called "spent grain." The mash is typically subjected to an extraction, which involves adding water to the mash in order to recover the residual soluble extract from the spent grain. The wort is then boiled vigorously to sterilize the wort and help develop the color, flavor and odor. Hops are added at some point during the boiling. The wort is cooled and transferred to a fermentor.

[0283] The wort is then contacted in a fermentor with yeast. The fermentor may be chilled to stop fermentation. The yeast flocculates and is removed. Finally, the beer is cooled and stored for a period of time, during which the beer clarifies and its flavor develops, and any material that might impair the appearance, flavor and shelf life of the beer settles out. The beer usually contains from about 2% to about 10% v/v alcohol, although beer with a higher alcohol content, *e.g.*, 18% v/v, may be obtained. Prior to packaging, the beer is carbonated and, optionally, filtered and pasteurized.

[0284] The brewing composition comprising an amylase, in combination with a glucoamylase and optionally a pullulanase and/or isoamylase, may be added to the mash of step (a) above, *i.e.*, during the preparation of the mash. Alternatively, or in addition, the brewing composition may be added to the mash of step (b) above, *i.e.*, during the filtration of the mash. Alternatively, or in addition, the brewing composition may be added to the wort of step (c) above, *i.e.*, during the fermenting of the wort.

[0285] A fermented beverage, such as a beer, can be produced by one of the methods above. The fermented beverage can be a beer, such as full malted beer, beer brewed under the "Reinheitsgebot," ale, IPA, lager, bitter, Happoshu (second beer), third beer, dry beer, near beer, light beer, low alcohol beer, low calorie beer, porter, bock beer, stout, malt liquor, non-alcoholic beer, non-alcoholic malt liquor and the like, but also alternative cereal and malt beverages such as fruit flavored malt beverages, *e.g.*, citrus flavored, such as lemon-, orange-, lime-, or berry-flavored malt beverages, liquor flavored malt beverages, *e.g.*, vodka-, rum-, or tequila-flavored malt liquor, or coffee flavored malt beverages, such as caffeine-flavored malt liquor, and the like.

9. Reduction of Iodine-Positive Starch

[0286] Variant amylases may reduce the iodine-positive starch (IPS), when used in a method of liquefaction and/or saccharification. One source of IPS is from amylose that escapes hydrolysis and/or from retrograded starch polymer. Starch retrogradation occurs spontaneously in a starch paste, or gel on ageing, because of the tendency of starch molecules to bind to one another followed by an increase in crystallinity. Solutions of low concentration become increasingly cloudy due to the progressive association of starch molecules into larger articles. Spontaneous precipitation takes place and the precipitated starch appears to be reverting to its original condition of cold-water insolubility. Pastes of higher concentration on cooling set to a gel, which on ageing becomes steadily firmer due to the increasing association of the starch molecules. This arises because of the strong tendency for hydrogen bond formation between hydroxy groups on adjacent starch molecules. See J.A. Radley, ed., STARCH AND ITS DERIVATIVES 194-201 (Chapman and Hall, London (1968)).

[0287] The presence of IPS in saccharide liquor negatively affects final product quality and represents a major issue with downstream processing. IPS plugs or slows filtration system, and fouls the carbon columns used for purification. When IPS reaches sufficiently high levels, it may leak through the carbon columns and decrease production efficiency. Additionally, it may result in hazy final product upon storage, which is unacceptable for final product quality. The amount of IPS can be reduced by isolating the saccharification tank and blending the contents back. IPS nevertheless will accumulate in carbon columns and filter systems, among other things. The use of variant amylases is expected to improve overall process performance by reducing the amount of IPS.

EXAMPLES

Example 1

Assays

[0288] Various assays used herein are set forth, below, for ease in reading. Any deviations from the protocols in later Examples are indicated in the relevant sections. In these experiments, a spectrophotometer was used to measure the absorbance of the products formed after the completion of the reactions. All assays were performed with culture supernatants treated with chelex beads.

A. Chelex bead treatment of culture supernatants

[0289] 96-well microtiter plates (MTPs) containing growing cultures were removed from incubators and EnzyScreen lids were replaced with disposable plastic sealers (Nunc cat. # 236366; Rochester, NY, USA). Cells were separated from culture supernatant via centrifugation (1118 RCF, 5 minutes). 150 μ L supernatant was removed from each well and transferred to filter plates (Millipore Multiscreen HTS, Billerica, MA, USA) containing Chelex beads prepared as described below. Plates were shaken vigorously for 5 minutes and supernatant from 3 replicate growth plates were collected into a single deep-well microtiter plate (Axygen, PDW-11-C) using a vacuum manifold device. Plates containing supernatants were sealed and stored at 4°C.

[0290] Chelex-100 beads, 200-400 mesh (BioRad, Hercules, CA, USA) were washed twice with 2 bed-volumes of 1 M HCl followed by 5 bed-volumes of ultrapure water on a sintered glass filter apparatus. 2 bed-volumes of 1 M KOH were used to wash the beads followed by another 5 bed-volume wash with ultrapure water. Filtered beads were transferred to a beaker and suspended with enough ultrapure water to produce slurry capable of mixing. The pH of the slurry was adjusted to 8-8.5 using HCl. The liquid was removed and the beads were dried using a sintered glass filter. A slurry of beads (40% w/v) was prepared in ultra pure water and its pH was adjusted to 8.0 using KOH/HCl. A slurry having a constant consistency was maintained by vigorous mixing. A bubble paddle reservoir device (V&P Scientific, San Diego, CA, USA) was used to transfer 100 μ L of slurry to all wells of filter plates. Liquid was removed using a vacuum manifold device.

B. Protein Determination Assay

[0291] Protein determination assays were performed using chelex bead-treated culture supernatant from cultures grown in 96-well micro-titer plates (MTPs) over 3 days at 37°C with shaking at 300 rpm and 80% humidity. A fresh 96-well round-bottom MTP containing 25 μ L supernatant per well was used for the High Performance Liquid Chromatography (HPLC) protein determination method. Supernatants were diluted four fold into 25 mM sodium acetate pH 5.5, and 10 μ L of each diluted sample was analyzed. An Agilent 1200 (Hewlett Packard) HPLC equipped with a Poroshell 300SB-C8 (Agilent Technologies Santa Clara, CA, USA) column was used. Sample was bound to the column using 25 mM sodium acetate pH 5.5 and eluted over a gradient up to 70% acetonitrile. Absorbance was measured at 220 nm, integrated using ChemStation software (Agilent Technologies) and the protein concentration of samples was determined based on a standard curve of purified CspAmy2-v1 protein.

C. Ceralpha α -Amylase Activity Assay

[0292] The Ceralpha α -amylase assay was performed using the Ceralpha Kit (Megazyme, Wicklow, Ireland). The assay involves incubating culture supernatant with a substrate mixture under defined conditions, and the reaction is terminated (and color developed) by the addition of borate buffer (200 mM Boric acid/NaOH buffer, pH 10). The substrate is a mixture of the defined oligosaccharide "nonreducing-end blocked *p*-nitrophenyl maltoheptaoside" (BNPNG7) and excess levels of α -glucosidase (which has no action on the native substrate due to the presence of the "blocking group"). On hydrolysis of the oligosaccharide by *endo*acting α -amylase, the excess quantities of α -glucosidase present in the mixture give instantaneous and quantitative hydrolysis of the *p*-nitrophenyl maltosaccharide fragment to glucose and free *p*-nitrophenol. The absorbance at 405 nm was measured, which relates directly to the level of amylase in the sample analyzed.

[0293] The equipment used for this assay included a Biomek FX Robot (Beckman Coulter Brea, CA, USA); a SpectraMAX MTP Reader (type 340-Molecular Devices, Sunnyvale, CA, USA) and iEMS incubator/shaker (Thermo Scientific, Rockford, IL, USA). The reagent and solutions used were:

- 1) *p*-nitrophenyl maltoheptaoside (BNPNG7) substrate (Megazyme Ceralpha HR kit);

- 2) 50 mM Malate buffer, 0.005% TWEEN® 80, pH 5.6 or 50 mM MOPS, 0.005% TWEEN® 80, pH 7 (dilution buffers); and
 3) 200 mM Boric acid / NaOH buffer, pH 10 (STOP buffer).

[0294] A vial containing 54.5 mg BPNPG7 substrate was dissolved in 10 mL of MilliQ water and then diluted into 30 mL of dilution buffer to make up 40 mL of the working substrate (1.36 mg/mL). The amylase samples (fermentation supernatant) were diluted 40X with dilution buffer. The assay was performed by adding 5 µL of diluted amylase solution into the wells of a MTP followed by the addition of 55 µL of diluted BPNPG7 working substrate solution. The solutions were mixed and the MTP was sealed with a plate seal and placed in an incubator/shaker (iEMS- Thermo Scientific) for 4 minutes at 25°C. The reaction was terminated by adding 70 µL STOP buffer and the absorbance was read at wavelength 400 nm in an MTP-Reader. A non-enzyme control was used to correct for background absorbance values.

D. Thermostability Assay

[0295] The thermostability of CspAmy2-v1 and variants was measured by determining the amylase activity using the Ceralpha α-amylase assay. The equipment used for this assay included a Biomek FX Robot (Beckman Coulter); a SpectraMAX MTP Reader (type 340-Molecular Devices), a Tetrad2DNA Engine PCR machine (Biorad), and iEMS incubator/shaker (Thermo Scientific). The reagent solutions used were (* not in all assays):

1) Heat stress buffers

- a) 50 mM KOAc pH 4.5 (5 ppm CaCl₂, 50 ppm NaCl)*,
- b) 50 mM KOAc pH 5.0 (10 ppm CaCl₂, 10 mM NaCl)
- c) 50 mM KOAc pH 5.7 (5 ppm CaCl₂, 50 ppm NaCl),
- d) 50 mM KOAc pH 5.7 (no salt condition)*,

2) p-nitrophenyl maltoheptaoside (BPNPG7) substrate (Megazyme Ceralpha HR kit):

3) 50 mM Malate buffer, 0.005% TWEEN® 80, pH 5.6 (dilution buffer); and

4) 200 mM Boric acid / NaOH, pH 10 (STOP buffer).

5) Amylase culture supernatant: 1:10 master dilution enzyme plates were diluted 1:10 in each of the four heat stress buffers in a PCR plate

[0296] 5 µL of the diluted enzyme samples were added to a 96-well PCR plate containing 55 µL of diluted BPNPG7 working substrate solution and the initial amylase activity of the samples was determined using the Ceralpha α-amylase assay as described in Section C. The samples were subjected to heat stress for 3-6 minutes in a PCR thermocycler as follows: Buffers (a) 50°C, (b) 59°-60°C, (c) 65°-70°C, and (d) 65°C. The heat stressed samples were cooled immediately to room temperature and 5 µL aliquots were assayed for amylase activity using the Ceralpha α-amylase assay as described in Section C. For each variant, the ratio of the initial and residual amylase activities was used to calculate the thermostability as follows: $\text{Thermostability} = [t_{\text{residual value}}] / [t_{\text{initial value}}]$, so the heat stability activity ratio was calculated based on enzyme activity after heat incubation divided by enzyme activity before heat incubation. For each sample (variants) the performance index (PI) is calculated. The performance index for thermostability stability is determined by comparing the thermostability of the variant enzyme with that of the similarly treated CspAmy2-v1 enzyme (SEQ ID NO: 3).

E. Starch Hydrolysis Assays (Corn Flour and Corn Starch Application assays)

[0297] Starch hydrolysis of corn flour and corn starch were used to measure specific activity of CspAmy2-v1 and variants. Activity was measured as reducing ends generated by the enzymatic breakdown of corn flour or corn starch. The reducing ends generated during the incubation with either substrate were quantified using a PAHBAH (p-hydroxybenzoic acid hydrazide) assay. The equipment used for the assay included a Biomek FX Robot (Beckman Coulter); a SpectraMAX MTP Reader (type 340-Molecular Devices), a Tetrad2DNA Engine PCR machine (Biorad), and iEMS incubator/shaker (Thermo Scientific), and a Bubble Paddle Reservoir.

[0298] Azure Farms Organic Corn Flour (Norco, CA) was ground to a fine powder using a consumer coffee grinder and then sifted to obtain a < 250 micron fraction. The sifted corn flour was washed extensively with MilliQ water by repeated suspension and centrifugation. Cargill Farms Organic Corn Starch material was also washed extensively with MilliQ water by repeated suspension and centrifugation.

[0299] Both corn flour and corn starch washed fractions were suspended in MilliQ water containing 0.005% sodium azide as 20% (w/w) stock solutions. The stock solutions were further diluted with a 20X stock buffer solution to 10.9% w/v corn flour and corn starch solutions (final buffer concentration: 55 mM KOAc, pH 5).

[0300] 55 μL of the diluted corn flour and corn starch substrates were added to PCR microtiter plates along with 5 μL of 1:10 diluted enzyme samples using a bubble paddle reservoir. The plates were sealed and placed at 83°C for 5 minutes followed by a ramp down to 45°C. The starch hydrolysis reaction was terminated by addition of 70 μL 0.1 N NaOH. The plates were sealed and centrifuged for 3 minutes at 1610 RCF. The starch hydrolysis reaction products from

both reactions were analyzed by the PAHBAH assay as described below.
[0301] *PAHBAH assay:* Aliquots of 80 μL of 0.5 N NaOH were added to all wells of an empty PCR plate (a "PAHBAH reaction plate"), followed by 20 μL of PAHBAH reagent (5% w/v p-hydroxybenzoic acid hydrazide (Sigma # H9882, St. Louis, MO), dissolved in 0.5 N HCl). The solutions were mixed by pipetting up and down. 20 μL of the starch hydrolysis reaction supernatants were added to each well of the PAHBAH reaction plate. The plates were sealed and placed in a thermocycler, programmed for 2 minutes at 95°C to develop color, and then cooled to 20°C. Samples of 80 μL of the developed PAHBAH reaction mixtures were transferred to a fresh plate, and absorbance was measured at 450 nm in a spectrophotometer.

F. CS-28 Rice Starch Microswatch Assay

[0302] The principle of this amylase assay is the liberation of an orange-dye due to the hydrolysis of rice starch incorporated in a cotton microswatch. The absorbance at 488 nm of the wash liquid is measured and this relates to the level of amylase activity in the sample analyzed at the desired conditions (pH, temperature, and buffer).

[0303] The equipment used for this assay included a Biomek FX Robot (Beckman Coulter), a SpectraMAX MTP Reader (type 340-Molecular Devices) and iEMS incubator/shaker (Thermo Scientific). The reagent and solutions used were:

- 1) CS-28 Microswatches (rice starch, colored);
- 2) 10 mM HEPES, 2 mM CaCl_2 , 0.005% TWEEN 80 buffer, pH 8.0, conductivity 1 mS/cm;
- 3) 25 mM CAPS, 2 mM CaCl_2 , 0.005% TWEEN 80 buffer, pH 10.0; conductivity 5 mS/cm (adjusted with 5M NaCl); and
- 4) 10 mM NaCl, 0.1 mM CaCl_2 , 0.005% TWEEN 80.
- 5) 50 mM MOPS pH7.15, 0.1 mM CaCl_2 .

[0304] CS-28 microswatches of 5.5 mm circular diameter were provided by the Center for Testmaterials (CFT, Vlaardingen, The Netherlands). Two microswatches were placed in each well of a 96-well Corning 9017 flat bottomed polystyrene MTP. The culture supernatants were diluted eight fold in 50 mM MOPS pH7.15, 0.1 mM CaCl_2 , and subsequently in 10 mM NaCl, 0.1 mM CaCl_2 , 0.005% TWEEN®80 solution to approximately 1 ppm, final enzyme concentration.

[0305] The incubator/shaker was set at the desired temperature, 25°C (ambient temperature) or 50°C. 174 μL or 177 μL of either HEPES or CAPS buffer, respectively, was added to each well of microswatch containing MTP and subsequently 6 μL or 3 μL of diluted enzyme solution was added to each well resulting in a total volume of 180 μL /well. The MTP was sealed with a plate seal and placed in the iEMS incubator/shaker and incubated for 15 minutes at 1150 rpm at 25°C for cleaning at pH 8, low conductivity (1 mS/cm), or 15 minutes at 1150 rpm at 50°C for cleaning at pH 10, high conductivity (5 mS/cm). Following incubation under the appropriate conditions, 100 μL of solution from each well was transferred to a new MTP, and the absorbance at 488 nm was measured using a MTP-spectrophotometer. Controls containing two microswatches and buffer but no enzyme were included for subtraction of background cleaning performance.

[0306] Each absorbance value was corrected by subtracting the blank (obtained after incubation of microswatches in the absence of enzyme), and the resulting absorbance provided a measure of the hydrolytic activity. A performance index (PI) was calculated for each sample.

[0307] For calculation of the wash performance indices (PI), the Langmuir equation was used to fit the data based on the CspAmy2-v1 enzyme (SEQ ID NO: 3) control. Using the protein concentration of the variants, the expected performance based on the curve-fit was calculated. The observed performance was divided by the calculated performance. This value was then divided by the performance of the CspAmy2-v1 enzyme (SEQ ID NO: 3).

G. Detergent Stability Assay

[0308] The stability of the reference amylase (CspAmy2-v1 enzyme (SEQ ID NO: 3)) and variants thereof was determined by measuring their activity after incubation under defined conditions, in the presence of a 10% detergent mixture (commercially purchased Persil Color Gel detergent, Henkel (Düsseldorf, Germany), purchased in 2011). The detergent was heat-inactivated before use, and the initial and residual amylase activities were determined using the Ceralpha α -amylase assay as described in section C, above.

[0309] The equipment used for this assay included a Biomek FX Robot (Beckman Coulter); a SpectraMAX MTP Reader (type 340-Molecular Devices), a Tetrad2DNA Engine PCR machine (Biorad), and iEMS incubator/shaker (Thermo Scientific). The reagent solutions used were:

- 1) p-nitrophenyl maltoheptaoside (BPNPG7) substrate (Megazyme Ceralpha HR kit);
- 2) Liquid detergent (Persil color gel, enzyme inactivated by heating for 2 hrs at 80°C);
- 3) 50 mM MOPS, 0.1 mM CaCl₂, 0.005% TWEEN®80 buffer, pH 7 (dilution buffer);
- 4) 10% detergent solution diluted in dilution buffer;
- 5) 200 mM Boric acid / NaOH buffer, pH 10 (STOP buffer)
- 6) Amylase culture supernatants diluted eight fold in 50 mM MOPS pH7.15, 0.1 mM CaCl₂ containing 0-100 µg/mL protein

[0310] 85 µL of a 10% detergent solution was added to a 96-well PCR plate and mixed with 15 µL of the diluted culture supernatant. A sample from the PCR plate was diluted 3X in dilution buffer and a 5 µL aliquot of this dilution was used to determine initial amylase activity. The PCR plate was incubated in a Tetrad PCR block at 80.5°C for 5 minutes. After incubation, detergent-enzyme mix was diluted 3X in dilution buffer and residual activity was measured. Initial (t_{initial}) and residual (t_{residual}) amylase activity was determined by the Ceralpha α -amylase assay as described above in Section C using a 5 µL sample.

[0311] For each variant, the ratio of the residual and initial amylase activities was used to calculate the detergent stability as follows: Detergent stability = [t_{residual} value] / [t_{initial} value].

[0312] For each sample (variants) the performance index (PI) was calculated. The performance index for detergent stability is determined by comparing the detergent stability of the variant enzyme with that of the similarly treated CspAmy2-v1 enzyme (SEQ ID NO: 3).

H. Performance index

[0313] The performance index (PI) compares the performance or stability of the variant and the parent enzyme (CspAmy2-v1) at the same protein concentration. In addition, the theoretical values can be calculated, using the parameters of the Langmuir equation of the standard enzyme. A performance index (PI) that is greater than 1 (PI>1) indicates improved performance by a variant as compared to the parent (e.g., CspAmy2-v1, SEQ ID NO: 3), while a PI of 1 (PI=1) identifies a variant that performs the same as the parent, and a PI that is less than 1 (PI<1) identifies a variant that performs worse than the parent.

Example 2

Generation of *B. subtilis* strains expressing CspAmy2-v1 and variants thereof

[0314] In this example, the construction of *Bacillus subtilis* strains expressing CspAmy2-v1 amylase and variants, thereof, are described. CspAmy2-v1 is a variant of CspAmy2 wild type (CspAmy2 wt) amylase having a deletion of both Arginine 178 and Glycine 179. CspAmy2 wt is an amylase from a *Cytophaga* sp., for which the nucleotide sequence was described by Chii-Ling et al. (2002) Appl. Environ. Microbiol. 68(7): 3651-3654. The CspAmy2-v1 amylase was described as having increased thermostability over the CspAmy2 wt amylase by Rong-Jen et al. (2003) Appl. Environ. Microbiol. 69(4): 2383-2385.

[0315] A synthetic DNA fragment (SEQ ID NO: 4, herein referred to as "CspAmy2-v1 DNA") encoding CspAmy2-v1 (SEQ ID NO: 2) amylase was produced by GeneArt AG (Regensburg, Germany) and served as template DNA for the construction of *Bacillus subtilis* strains expressing CspAmy2-v1 amylase and variants, thereof.

[0316] SEQ ID NO:4 includes a codon-modified nucleotide sequence encoding the mature form of CspAmy2-v1 amylase adjacent to a sequence encoding the LAT signal peptide (underlined):

ATGAAACAACAAAAACGGCTTTACGCCCCGATTGCTGACGCTGTTATTTGCGCTCATC
TTCTTGCTGCCTCATTCTGCAGCTAGCGCAGCAGCGACAAACGGAACAATGATGCA
 5 GTATTTTCGAGTGGTATGTACCTAACGACGGCCAGCAATGGAACAGACTGAGAACAG
 ATGCCCCTTACTTGTCATCTGTTGGTATTACAGCAGTATGGACACCGCCGGCTTATA
 AGGGCACGTCTCAAGCAGATGTGGGGGTACGGCCCGTACGATCTGTATGATTTAGGC
 10 GAGTTTAATCAAAAAGGTACAGTCAGAACGAAGTATGGCACAAAAGGAGAACTTA
 AATCTGCTGTTAACACGCTGCATTCAAATGGAATCCAAGTGTATGGTGATGTCGTGA
 TGAATCATAAAGCAGGTGCTGATTATACAGAAAACGTAACGGCGGTGGAGGTGAAT
 15 CCGTCTAATAGAAATCAGGAAACGAGCGGCGAATATAATATTCAGGCATGGACAGG
 CTTCAACTTTCCGGGCAGAGGAACAACGTATTCTAACTTCAAATGGCAGTGGTTCCA
 TTTTGATGGAACGGATTGGGACCAGAGCAGAAGCCTCTCTAGAATCTTCAAATTCA
 20 CGGGAAAGGCGTGGGACTGGGAGGTTTCTTCAGAAAACGGAAATTATGACTATCTG
 ATGTACGCGGACATTGATTATGACCATCCGGATGTCGTGAATGAAATGAAAAAGTG
 GGGCGTCTGGTATGCCAACGAAGTTGGGTAGATGGATACAGACTTGACGCGGTCA
 25 AACATATTAAATTTAGCTTTCTCAAAGACTGGGTGGATAACGCAAGAGCAGCGACG
 GGAAAAGAAATGTTTACGGTTGGCGAATATTGGCAAAATGATTTAGGGGCCCTGAA
 TAACTACCTGGCAAAGGTAAATTACAACCAATCTCTTTTTGATGCGCCGTTGCATTA
 30 CAACTTTTACGCTGCCTCAACAGGGGGTGGATATTACGATATGAGAAATATTCTTAA
 TAACACGTTAGTCGCAAGCAATCCGACAAAGGCTGTTACGTTAGTTGAGAATCATG
 ACACACAGCCTGGACAATCACTGGAATCAACAGTCCAACCGTGGTTTAAACCGTTA
 35 GCCTACGCGTTTATTCTCACGAGAAGCGGAGGCTATCCTTCTGTATTTTATGGAGAT
 ATGTACGGTACAAAAGGAACGACAACAAGAGAGATCCCTGCTCTTAAATCTAAAAT
 40 CGAACCTTTGCTTAAGGCTAGAAAAGACTATGCTTATGGAACACAGAGAGACTATA
 TTGATAACCCGGATGTCATTGGCTGGACGAGAGAAGGGGACTCAACGAAAGCCAA
 GAGCGGTCTGGCCACAGTGATTACAGATGGGCCGGGCGGTTCAAAAAGAATGTATG
 45 TTGGCACGAGCAATGCGGGTGAAATCTGGTATGATTTGACAGGGAATAGAACAGAT
 AAAATCACGATTGGAAGCGATGGCTATGCAACATTTCTGTCAATGGGGGCTCAGT
 TTCAGTATGGGTGCAGCAA

50 **[0317]** The precursor form of the CspAmy2-v1 polypeptide produced from the pHPLT02-CspAmy2-v1 vector is shown, below, as SEQ ID NO: 3. The LAT signal peptide is underlined:

MKQQKRLYARLLTLLFALIFLLPHSAASAAATNGTMMQYFEWYVPNDGQQWNRLRTD
 APYLSSVGITAVWTPPAYKGTSQADVGYGPYDLYDLGEFNQKGTVRTKYGTKGELKS
 5 AVNTLHSNGIQVYGDVVMNHKAGADYTENVTAVEVNPSNRNQETSGEYNIQAWTGFN
 FPGRGTTYSNFKWQWFHFDGTDWDQSRSLSRIFKFTGKAWDWEVSSENGNYDYLMY
 ADIDYDHPDVVNEMKKWGVWYANEVGLDGYRLDAVKHIKFSFLKDWVDNARAATG
 10 KEMFTVGEYWQNDLGALNNYLAKVNYNQSLFDAPLHYNFYAASTGGGYDDMRNINL
 NTLVASNPTKAVTLVENHDTQPGQSLESTVQPWFKPLAYAFILTRSGGYPSVIFYGDMY
 GTKGTTTREPALKSKIEPLLKARKDYAYGTQRDYIDNPDVIGWTREGDSTKAKSGLAT
 15 VITDGPGGSKRMYVGTSNAGEIWDLTGNRTDKITIGSDGYATFPVNGGSVSVWVQQ

[0318] The mature form of the CspAmy2-v1 polypeptide produced from the pHPLT02-CspAmy2-v1 vector is shown,
 below, as SEQ ID NO: 2.

AATNGTMMQYFEWYVPNDGQQWNRLRTDAPYLSSVGITAVWTPPAYKGTSQADVGY
 GPYDLYDLGEFNQKGTVRTKYGTKGELKSAVNTLHSNGIQVYGDVVMNHKAGADYT
 25 ENVTAVEVNPSNRNQETSGEYNIQAWTGFNFPGRGTTYSNFKWQWFHFDGTDWDQSR
 SLSRIFKFTGKAWDWEVSSENGNYDYLMYADIDYDHPDVVNEMKKWGVWYANEVGL
 DGYRLDAVKHIKFSFLKDWVDNARAATGKEMFTVGEYWQNDLGALNNYLAKVNYN
 30 QSLFDAPLHYNFYAASTGGGYDDMRNINLNTLVASNPTKAVTLVENHDTQPGQSLEST
 VQPWFKPLAYAFILTRSGGYPSVIFYGDMYGTKGTTTREPALKSKIEPLLKARKDYAYG
 TQRDYIDNPDVIGWTREGDSTKAKSGLATVITDGPGGSKRMYVGTSNAGEIWDLTGN
 35 RTDKITIGSDGYATFPVNGGSVSVWVQQ

[0319] To express CspAmy2-v1, the *CspAmy2-v1* DNA fragment was cloned into the pHPLT02 vector, a modified
 version of the pHPLT vector (Solingen et al.(2001) *Extremophiles* 5:333-341) by GeneArt and fused in-frame to the
 40 *AmyL* (LAT) signal peptide using the unique *NheI* and *XhoI* restriction sites, resulting in plasmid pHPLT02-CspAmy2-
 v1. The pHPLT expression vector contains the *B. licheniformis* LAT promoter (Plat) and additional elements from pUB110
 (McKenzie et al. (1986) *Plasmid*, 15: 93-103) including a replicase gene (reppUB), a neomycin/kanamycin resistance
 gene (neo) and a bleomycin resistance marker (bleo). Site-directed mutagenesis (Stratagene) was used to change the
 45 nucleotides 5'-TCA-3' of Serine 28 of the *AmyL* signal peptide to nucleotides 5'-AGC-3' in order to introduce the unique
NheI restriction site.

[0320] A suitable *B. subtilis* strain was transformed with pHPLT02-CspAmy2-v1 plasmid DNA using a method known
 in the art (WO 02/14490). The *B. subtilis* transformants were selected on agar plates containing heart infusion agar
 (Difco, Catalog No. 244400, Lawrence, KS, USA and 10 mg/L neomycin sulfate (Sigma, Catalog No. N-1876; contains
 732 µg neomycin per mg, St. Louis, Missouri, USA). Selective growth of *B. subtilis* transformants harboring the pHPLT02-
 50 CspAmy2-v1 plasmid was performed in shake flasks at 37°C for ~65h in MBD medium (enriched semi-defined medium
 based on MOPs buffer, with urea as major nitrogen source, glucose as the main carbon source, and supplemented with
 1% soytone for robust cell growth) containing 5 mM CaCl₂ and 10 ppm neomycin. Growth resulted in the production of
 secreted CspAmy2-v1 amylase with starch hydrolyzing activity.

Example 3

Generation of CspAmy2-v1 Site Evaluation Libraries

[0321] The construction of a CspAmy2-v1 site evaluation library (SEL) was performed by GeneArt using their technology platform for gene optimization, gene synthesis, and library generation (see, e.g., European Patent Nos. 0 200 362 and 0 201 184, US Patent Nos. 4,683,195, 4,683,202, and 6,472,184, and international patent application number WO 2004/059556A3). The pHPLT02-CspAmy2-v1 plasmid DNA served as template to produce SELs at each of the sites in the mature region of CspAmy2-v1 protein (SEQ ID NO: 3). GeneArt was commissioned to create the SELs at the positions using their standard protocols. The corresponding codons for each site were each substituted with codons for at least 10 (out of a possible 19) different amino acids. The codon-mutagenized pHPLT02-CspAmy2-v1 mixtures were used to transform competent *B. subtilis* cells as known in the art (WO 2002/014490) to generate the CspAmy2-v1 SELs. Transformation mixtures were plated on HI-agar plates (Heart Infusion agar) containing 10 mg/L neomycin sulfate. For each library, single bacterial colonies were picked and grown in TSB (tryptone and soy-based broth) liquid medium with 10 mg/ml neomycin selection for subsequent DNA isolation and gene sequence analysis. Sequence analysis data revealed a maximum of 19 CspAmy2-v1 mature variants per library. To generate CspAmy2-v1 and variant enzyme samples for biochemical characterization, selective growth of the variants was performed in 96 well MTPs at 37°C for ~65 hours with 70% humidity in MBD medium. A total of 7870 out of the 9139 possible variants were obtained for 478 out of the 481 positions mutagenized.

Example 4

Identification of Combinable and Productive Mutations

[0322] Performance index (PI) values were determined for all the CspAmy2-v1 amylase variants tested using the assays described in Example 1: Corn flour and corn starch application assays, thermostability assay (at pH 5.0 and pH 5.7), CS-28 microswatch assay (at pH 8 and pH 10), detergent stability assay, Ceralpha activity assay, and protein determination. Productive positions are described as those positions within a molecule that are most useful for making combinatorial variants exhibiting an improved characteristic, where each production position allows for at least one combinable mutation. Combinable mutations are mutations at any amino acid position that can be used to make combinatorial variants. Combinable mutations improve at least one desired property of the molecule, while not significantly decreasing expression, activity, or stability. Combinable mutations can be grouped as follows:

Group A: A mutation that produces a variant wherein the minimum performance indices (PI) relative to a defined parental protein for: (i) protein expression, (ii) CS-28 microswatch activity at pH 8 (25°C) or pH10 (50°C), or activity in Ceralpha assay, or activity in corn flour or corn starch application assays, and (iii) detergent stability or thermostability at pH 5.0 or pH 5.7 is greater than or equal to 0.9, and in addition have a PI for any one of these tests that is greater than or equal to 1.0.

Group B: A mutation that produces a variant wherein the minimum performance indices (PI) relative to a defined parental protein for: (i) protein expression, (ii) CS-28 microswatch activity at pH 8 (25°C) or pH10 (50°C), or activity in Ceralpha assay, or activity in corn flour or corn starch application assays, and (iii) detergent stability or thermostability at pH 5.0 or pH 5.7 is greater than or equal to 0.8, and in addition have a PI for any one of these tests that is greater than or equal to 1.2.

Group C: A mutation that produces a variant wherein the minimum performance indices (PI) relative to a defined parental protein for: (i) protein expression, (ii) CS-28 microswatch activity at pH 8 (25°C) or pH10 (50°C), or activity in Ceralpha assay, or activity in corn flour or corn starch application assays, and (iii) detergent stability or thermostability at pH 5.0 or pH 5.7 is greater than or equal to 0.5, and in addition have a PI for any one of these tests that is greater than or equal to 1.5.

[0323] The properties of combinable mutations are summarized in Table 4.1.

Table 4.1. Properties for each group of combinable mutations

Group	Performance Index (PI)						
	Expression	Cleaning (pH 8 or 10)	Stability (detergent)	Activity (corn flour or corn starch)	Activity in Ceralpha assay	Thermostability (pH 5.0 or pH 5.7)	Minimum PI in one or more tests
A	≥ 0.9	≥ 0.9	≥ 0.9	≥ 0.9	≥ 0.9	≥ 0.9	X ≥ 1.0
B	≥ 0.8	≥ 0.8	≥ 0.8	≥ 0.8	≥ 0.8	≥ 0.8	X ≥ 1.2
C	≥ 0.5	≥ 0.5	≥ 0.5	≥ 0.5	≥ 0.5	≥ 0.5	X ≥ 1.5

[0324] Preferred combinable mutations are at "productive positions," as described, below. In the case of the present amylases, "activity" refers to α -amylase activity, which can be measured as described, herein.

[0325] Productive positions are amino acid positions that are tolerant to substitution with different amino acid residues, wherein the resulting variants meet a set of performance criteria for combinability, as set forth above. Productive positions can be assigned a Productivity Score as follows: Positions where less than 15% of the substitutions at a given position fall within groups A, B, or C are given a Productivity Score of "1". Positions where less than 30%, but greater than, or equal to 15% of the substitutions at a given position fall within groups A, B, or C are given a Productivity Score of "2". Positions where less than 50%, but greater than, or equal to 30% of the substitutions at a given position fall within groups A, B, or C are given a Productivity Score of "3". Positions where 50% or more of the substitutions at a given position fall within groups A, B, or C are given a Productivity Score of "4".

[0326] Suitability score refers to the ability of one or more combinable mutations to be used to make combinatorial variants, based on the performance criteria for combinability, (*i.e.*, A, B, and C, as set forth, above) in which each of the mutations fall. A higher suitability score indicates a mutation or mutations that are more suitable for use in making combinatorial variants. Suitability scores are described in Table 4.2.

Table 4.2. Definitions of suitability scores

Substitutions Occur in Group(s)	Suitability Score
A, B and C	+++++
A and B	++++
A or (B and C)	+++
B	++
C	+

[0327] Suitability scores of individual substitutions in CspAmy2-v1 are shown in Table 4.3. For each CspAmy2-v1 protein position, variants are listed according to the suitability score they received (+, ++, +++, +++++, or ++++++). Position numbering is based on the mature CspAmy2 polypeptide (SEQ ID NO: 1).

Table 4.3: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)	(++++)	(+++++)
1		I	A	EGKNQRTVY	
2			AEGHKNPQ RSY		
3			TADFGMPQ RS		
4	T	Q	N		

(continued)

Table 4.3: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE					
	(+)	(++)	(+++)	WT AA 1ST	(++++)	(+++++)
						ACDEFHIKL MNPQRSTVY
5			G			
7			MI			
8			MF			
11			F	Y		
15			VN	ILS		CT
20			QE			
21			QLW	T		
23	DF		NSWY	AEHKMQTV		
26			RK			
			TDEFGHIKL MNQSWY	R		
27						
28			DAEN			
30			PHT	ACFLRWY		DEGKNS
31		M	YEKR	FHW		
33			SDEGHNQR	K		
35			VHIMN			
38			TS	DN		
39			AS			
40			VI			
42			TACILMV			
45			APS			
46			YFMT			
48		A	G			
49	M	CGHY	TS	ADEFIKLNV		
50			SEK	D		
51			QS			
			AFGHIKNQS TW			
52						
54	RS	G	VLQ	CDIN		AET
57	K		G			
58			P	C		
68			NACEFMSY			
70	NW	R	K			

(continued)

Table 4.3: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)	(++++)	(+++++)
71			GAN		
72	H		TGS		
73			VT		
75	C		T		
81			GIPTV	ADFHKLMNSY	EQRW
82			EQ		
83			LF		
84			KIQV		
85			SACDEHKM NQR	L	
87		I	V		
88			NADEQRT		H
89			TCDEHMQN RSV		
92	L		SAMRV		
93	DT		NM		
94			GN		
96	I		Q		
97			VI		
98	W		YF		
101			VI		
103			MV	I	
104	D		N		
106	AIV		K		
107	G		A		
108	KR		GAS		
109			AP		
111		M	YW	ACDGHKLNQRST V	F
112			TFGIWV	ACDELMPQRY	
113	Y		ED		
114			NG		
115		A	V	IM	

(continued)

Table 4.3: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)	(++++)	(+++++)
116			T	ACDEGHIKLMNS VW	
117		C	AS		
118	Q	LM	VACFIKNS	R	
119	S		E		
120			V	C	
121		S	NKR		
122		T	P	AKQR	
123			S	ACEGHKNQTY	R
124			ND		
126	Y		N		
127		CY	QAM	HIKRTV	E
128			ES	GKVY	I
129		AFG	TQY	HIKLRSV	
130		AIRV	ST	GH	
131		FM	GT	AHIKNW	PQ
132			ENS	ACGHILMRTVY	PQ
133		FN	YE	ADHLTV	K
134			NCFY	DGHMPQST	
135	HR	M	IV		
136	W		QY	AFGHIK	NT
137		V	A		
138	PR	GV	W	AFKLMQSTY	DH
140	M	D	GCEFHLTV Y	S	KN
141	H		F	W	
142		CV	NEY	FGHIKLQRW	ADMST
144	DQ	F	PCHM	AGKLNRY	IT
145	LWY	IMV	GEQT	AHKNRS	
147			GCLV	EHKNQR	I
148			TAGHIKLSW Y		

(continued)

Table 4.3: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)	(++++)	(+++++)
149			TACDEHIKM NRSWY	L	
150	DPQ	GM	Y	FHW	I
151			SD		
152		F	NW	ACEGHLMRSTV	DQ
153			F	H	WY
154			KAENHRST	F	
156	FGILMV		QDSTY	H	K
157	N		W		
158	P		FADEIRY	LNSV	CGHTW
160	C		F		
163	F	CMNQ V	T	DLS	
164		F	DN		
165	P	H	WF		
166	ACGKM		D		
167			Q	ACDGHKNRSTVY	
168		C	SRW	DEGIKMNTV	L
169	EQW		RDMT	CHKLY	A
170			S	CGHNRT	
171			L	FGIMRVWY	HKNQS
172			SACDEHKNR T		
173			RK		w
174		L	I		
175	AD	S	FLMV	HWY	
176	L		K		
177	G		FH		
182	H		K		
183			AE	K	R
187			E	V	P
189	A		SC	D	
190			S		P
191			EACLMNT	I	
192	S	M	NR		FHY
193			GACFIKST	HRV	

(continued)

Table 4.3: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)	(++++)	(+++++)
195		D	Y		
198	CG		L		A
200	EL		Y		
201	L		A		
203	HNQ	A	I	CV	FLMY
206	I	CY	DM		AQ
207		AKR	HF	DEMSY	N
208			PAEHKLNQR ST	D	
209		CIK	D	G	
210		S	V		
211		FS	V	CEIMNT	DLQ
212			NY	ACDEHILMV	GQRST
215		LY	KRT	EFMNQ	
216			KR		
219			VEL	I	
222		T	A		
225			VL		
226		C	GKQ	E	
227			LY		
232			L	R	V
235		LT	VA	C	
238	M		I	LQR	P
239	M	Q	K	C	
240		D	F		
241		T	SFR	P	ACDEGHIKL NQVWY
242	E	Y	FV		I
243			LAST	IM	
244		Y	KT	ACHMNQRS	
245			DE		
246			WF		
248	A		DQ	N	
249			NKR	ACDEFGMST	QW

(continued)

Table 4.3: Suitability scores of individual substitutions in CspAmy2-v1

	VARIANTS SUITABILITY SCORE					
POS	(+)	(++)	(+++)	WT AA 1ST	(++++)	(+++++)
250			AS			
251	ADKLMN					
	QST		R			
252			ACDFHMR			
253			ACDELMNQ			
			ST	RV		
254		F	TILM			
256			KR			
257	ACGRVY	FHK	E	LQS		
258			MCI	FLY		
259	P		F			
260	Y		T	S	AL	
262			G	A		
266	EK	HLN	Q	CFGMRW	ADIST	
267			ND			
269	A	V	LI			
270		CIQV	G	ADFHKPRSWY		
271			ACS			
273			NDEILMQS	HK		
275			YF			
276			LM			
277		D	ACNW	FHILMQTY	EK	
280			NCY	ADGHKLQT	E	
281		G	YA	D		
283			QT	HV		
284	EF		SHY	KMRW		
285			LA			
286	L		F	M		
288			A	V		
295			YLQ	H		
296		FIMQS				
		T	ADHKR	E		
299			TEI	R		
300		Q	GAKL	R		

(continued)

Table 4.3: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)	(++++)	(+++++)
301			GT	AFHKMQRSY	
302		S	G		
303		F	Y	IW	ARTV
307			RQS		
308			NV	ADGLQTY	CEFHMR
310			LDET		
311		E	NDHKQSV		
312			NDG		
313		AS	T		
316			AKQS		
317	W	A	SDGLT	CEMQY	HK
318			NFILVW	ACDEKMQRSY	H
320	PV	M	TCGKY	AHNQRW	DE
321			KHR		
325			LFIMV		
327			ED		
335	G		Q		
336			SAD		
339	Q		S		
342			QEL		
343	AMW		P		
348			LCGS	Q	
349	W		AS	G	
357		Y	SACDELMN V	Q	
358			GDEQS		
360	CEHV		YI	F	L
362	EQT		S	AC	IV
363			VL	I	
368			MLWY		
372			KADHMNR		
374			TAKNPQS		
375			TS		
376		H	TAGNQSY	K	
377	ILMV	C	R	H	ADGKSTY

(continued)

Table 4.3: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)	(++++)	(+++++)
378			E	Q	
384		D	SEGN	HP	
385			KAE		
388		L	PCDKRS	I	
390			LC	I	
391			KFTVY	EL	
392			AG		
394	E		KHM		
395		IKS	DC	EFMQRWY	
396			YCFK		
397			AS		
400			T	ACDEMNRQSWY	HKLV
401			QM		
402			RFKLSTVWY	Q	
403			DS		
404			YW		
405			IL		
407			NACDEGHQ S		
408	S	Y	PEHMQR	KVW	
409		N	D		
410		E	VIKLMRSY		
414			TAS		
416		MT	E	CFHKLNRSV	DGQ
418			DACEFGHIK LMNQRSTW Y		
419			SAEGKPQRT VY		
420			TFI	ACDEGHPQRSVW Y	L
421				N	

(continued)

Table 4.3: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)	(++++)	(+++++)
422			A	EIKLMNPQRSTV WY	CD
423	P		KV	ACEFGMNQWY	DS
424			SA		
426			LC		
427			A	C	GS
429			VC	LM	
430		C	IL	EM	
431		D	TA		
433	HS	A	GN		
434		M	PNR	DQS	
435		I	GACEQRSTY	LVN	
437			SADEFHKLQ Y		
441			YCKLNQRS W		
442		L	VAIT		
444			TAEFHIKLM NPQVY		
445			SACDHMRT VW		
447			AGKQRSTV		
448			GEL	AFHKNQTY	DW
449			EQ		
450			IDLP	ACFHKQRTV	N
452			YAILMSVW		
454			LAEFHKS	CIMQTVY	
455			TIV	ACLMS	
456			G	ACDEFHKL MNRS TWY	
458			RDEFSW	ACIN	MY

(continued)

Table 4.3: Suitability scores of individual substitutions in CspAmy2-v1

POS	VARIANTS SUITABILITY SCORE				
	(+)	(++)	(+++)	(++++)	(+++++)
459			TACDFGLSV W		
460			DEHN		
461			KADGILMNP SY		
462			IV		
463			TCEFIKLMN VY		
464			IV		
465			GAMNPQ		
466			SCFN	ADEHILMRVWY	KT
467			DN		
469			YCFISV	L	
470			AG		
471	K		TADEFGHIN PQW		
474			VC		
475			NFIP	ADGHKLMSTV	
476	DE		G		ACHKNPQRS TVY
477	P		G		ADEHIKQRST VY
479			VCHW		
480			SG		
481			VACIL		
482			WY		
483			VI		
484		AHK	Q		
485			QFR	ADEHIKLMNPTV Y	

[0328] The productive positions in CspAmy2-v1 that fall within the previously described Productivity Scores of "1," "2," "3," and "4," and the substitutions within those positions that are combinable, are listed below. Each position identified in the list, and each substitution identified in parenthesis following the numerical position identifier, represents a mutation that has been determined, based on experimental data, to contribute to the performance of an amylase variant, particularly a variant comprising more than one of the described mutations. The position numbering is based on mature CspAmy2 polypeptide (SEQ ID NO: 1).

[0329] The productive positions in CspAmy2-v1 that fall within the previously described Productivity Scores of "1," "2," "3," and "4," and the substitutions within those positions that are combinable, are listed below. The position numbering is based on mature CspAmy2 polypeptide (SEQ ID NO: 1).

LIST A:

[0330]

1(A,E,G,I,K,N,Q,R,T,V,Y); 2(A,E,G,H,K,N,P,Q,R,S,Y); 3(T,A,D,F,G,M,P,Q,R,S); 4(N,Q,T);
 5(5(G,A,C,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,Y); 7(M,I); 8(M,F); 11(F,Y); 15(V,C,I,L,N,S,T);
 20(Q,E); 21(Q,L,T,W); 23(N,A,D,E,F,H,K,M,Q,S,T,V,W,Y); 26(R,K);
 27(T,D,E,F,G,H,I,K,L,M,N,Q,R,S,W,Y); 28(D,A,E,N);
 30(P,A,C,D,E,F,G,H,K,L,N,R,S,T,W,Y); 31(Y,E,F,H,K,M,R,W); 33(S,D,E,G,H,K,N,Q,R);
 35(V,H,I,M,N); 38(T,D,N,S); 39(A,S); 40(V,I); 42(T,A,C,I,L,M,V); 45(A,P,S); 46(Y,F,M,T);
 48(G,A); 49(T,A,C,D,E,F,G,H,I,K,L,M,N,S,V,Y); 50(S,D,E,K); 51(Q,S);
 52(A,F,G,H,I,K,N,Q,S,T,W); 54(V,A,C,D,E,G,I,L,N,Q,R,S,T); 57(G,K); 58(P,C);
 68(N,A,C,E,F,M,S,Y); 70(K,N,R,W); 71(G,A,N); 72(T,G,H,S); 73(V,T); 75(T,C);
 81(G,A,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,W,Y); 82(E,Q); 83(L,F); 84(K,I,Q,V);
 85(S,A,C,D,E,H,K,L,M,N,Q,R); 87(V,I); 88(N,A,D,E,H,Q,R,T); 89(T,C,D,E,H,M,N,Q,R,S,V);
 92(S,A,L,M,R,V); 93(N,D,M,T); 94(G,N); 96(Q,I); 97(V,I); 98(Y,F,W); 101(V,I); 103(M,I,V);
 104(N,D); 106(K,A,I,V); 107(A,G); 108(G,A,K,R,S); 109(A,P);
 111(Y,A,C,D,F,G,H,K,L,M,N,Q,R,S,T,V,W); 112(T,A,C,D,E,F,G,I,L,M,P,Q,R,V,W,Y);
 113(E,D,Y); 114(N,G); 115(V,A,I,M); 116(T,A,C,D,E,G,H,I,K,L,M,N,S,V,W); 117(A,C,S);
 118(V,A,C,F,I,K,L,M,N,Q,R,S); 119(E,S); 120(V,C); 121(N,K,R,S); 122(P,A,K,Q,R,T);
 123(S,A,C,E,G,H,K,N,Q,R,T,Y); 124(N,D); 126(N,Y); 127(Q,A,C,E,H,I,K,M,R,T,V,Y);
 128(E,G,I,K,S,V,Y); 129(T,A,F,G,H,I,K,L,Q,R,S,V,Y); 130(S,A,G,H,I,R,T,V);
 131(G,A,F,H,I,K,M,N,P,Q,T,W); 132(E,A,C,G,H,I,L,M,N,P,Q,R,S,T,V,Y);
 133(Y,A,D,E,F,H,K,L,N,T,V); 134(N,C,D,F,G,H,M,P,Q,S,T,Y); 135(I,H,M,R,V);
 136(Q,A,F,G,H,I,K,N,T,W,Y); 137(A,V); 138(W,A,D,F,G,H,K,L,M,P,Q,R,S,T,V,Y);
 140(G,C,D,E,F,H,I,K,L,M,N,S,T,V,Y); 141(F,H,W);
 142(N,A,C,D,E,F,G,H,I,K,L,M,Q,R,S,T,V,W,Y); 144(P,A,C,D,F,G,H,I,K,L,M,N,Q,R,T,Y);
 145(G,A,E,H,I,K,L,M,N,Q,R,S,T,V,W,Y); 147(G,C,E,H,I,K,L,N,Q,R,V);
 148(T,A,G,H,I,K,L,S,W,Y); 149(T,A,C,D,E,H,I,K,L,M,N,R,S,W,Y);
 150(Y,D,F,G,H,I,M,P,Q,W); 151(S,D); 152(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,W);
 153(F,H,W,Y); 154(K,A,E,F,H,N,R,S,T); 156(Q,D,F,G,H,I,K,L,M,S,T,V,Y); 157(W,N);
 158(F,A,C,D,E,G,H,I,L,N,P,R,S,T,V,W,Y); 160(F,C); 163(T,C,D,F,L,M,N,Q,S,V); 164(D,F,N);
 165(W,F,H,P); 166(D,A,C,G,K,M); 167(Q,A,C,D,G,H,K,N,R,S,T,V,Y);
 168(S,C,D,E,G,I,K,L,M,N,R,T,V,W); 169(R,A,C,D,E,H,K,L,M,Q,T,W,Y);
 170(S,C,G,H,N,R,T); 171(L,F,G,H,I,K,M,N,Q,R,S,V,W,Y); 172(S,A,C,D,E,H,K,N,R,T);
 173(R,K,W); 174(I,L); 175(F,A,D,H,L,M,S,V,W,Y); 176(K,L); 177(F,G,H); 182(K,H);
 183(A,E,K,R); 187(E,P,V); 189(S,A,C,D); 190(S,P); 191(E,A,C,I,L,M,N,T);
 192(N,F,H,M,R,S,Y); 193(G,A,C,F,H,I,K,R,S,T,V); 195(Y,D); 198(L,A,C,G); 200(Y,E,L);
 201(A,L); 203(I,A,C,F,H,L,M,N,Q,V,Y); 206(D,A,C,I,M,Q,Y); 207(H,A,D,E,F,K,M,N,R,S,Y);
 208(P,A,D,E,H,K,L,N,Q,R,S,T); 209(D,C,G,I,K); 210(V,S); 211(V,C,D,E,F,I,L,M,N,Q,S,T);
 212(N,A,C,D,E,G,H,I,L,M,Q,R,ST,V,Y); 215(K,E,F,L,M,N,Q,R,T,Y); 216(K,R); 219(V,E,I,L);
 222(A,T); 225(V,L); 226(G,C,E,K,Q); 227(L,Y); 232(L,R,V); 235(V,A,C,L,T);
 238(I,L,M,P,Q,R); 239(K,C,M,Q); 240(F,D); 241(S,A,C,D,E,F,G,H,I,K,L,N,P,Q,R,T,V,W,Y);
 242(F,E,I,V,Y); 243(L,A,I,M,S,T); 244(K,A,C,H,M,N,Q,R,S,T,Y); 245(D,E); 246(W,F);
 248(D,A,N,Q); 249(N,A,C,D,E,F,G,K,M,Q,R,S,T,W); 250(A,S); 251(R,A,D,K,L,M,N,Q,S,T);
 252(A,C,D,F,H,M,R); 253(A,C,D,E,L,M,N,Q,R,S,T,V); 254(T,F,I,L,M); 256(K,R);
 257(E,A,C,F,G,H,K,L,Q,R,S,V,Y); 258(M,C,F,I,L,Y); 259(F,P); 260(T,A,L,S,Y); 262(G,A);
 266(Q,A,C,D,E,F,G,H,I,K,L,M,N,R,S,T,W); 267(N,D); 269(L,A,I,V);

270(G,A,C,D,F,H,I,K,P,Q,R,S,V,W,Y); 271(A,C,S); 273(N,D,E,H,I,K,L,M,Q,S); 275(Y,F);
 276(L,M); 277(A,C,D,E,F,H,I,K,L,M,N,Q,T,W,Y); 280(N,A,C,D,E,G,H,K,L,Q,T,Y);
 281(Y,A,D,G); 283(Q,H,T,V); 284(S,E,F,H,K,M,R,W,Y); 285(L,A); 286(F,L,M); 288(A,V);
 295(Y,H,L,Q); 296(A,D,E,F,H,I,K,M,Q,R,S,T); 299(T,E,I,R); 300(G,A,K,L,Q,R);
 5 301(G,A,F,H,K,M,Q,R,S,T,Y); 302(G,S); 303(Y,A,F,I,R,T,V,W); 307(R,Q,S);
 308(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,Y); 310(L,D,E,T); 311(N,D,E,H,K,Q,S,V); 312(N,D,G);
 313(T,A,S); 316(A,K,Q,S); 317(S,A,C,D,E,G,H,K,L,M,Q,T,W,Y);
 318(N,A,C,D,E,F,H,I,K,L,M,Q,R,S,V,W,Y); 320(T,A,C,D,E,G,H,K,M,N,P,Q,R,V,W,Y);
 321(K,H,R); 325(L,F,I,M,V); 327(E,D); 335(Q,G); 336(S,A,D); 339(S,Q); 342(Q,E,L);
 10 343(P,A,M,W); 348(L,C,G,Q,S); 349(A,G,S,W); 357(S,A,C,D,E,L,M,N,Q,V,Y);
 358(G,D,E,Q,S); 360(Y,C,E,F,H,I,L,V); 362(S,A,C,E,I,Q,T,V); 363(V,I,L); 368(M,L,W,Y);
 372(K,A,D,H,M,N,R); 374(T,A,K,N,P,Q,S); 375(T,S); 376(T,A,G,H,K,N,Q,S,Y);
 377(R,A,C,D,G,H,I,K,L,M,S,T,V,Y); 378(E,Q); 384(S,D,E,G,H,N,P); 385(K,A,E);
 388(P,C,D,I,K,L,R,S); 390(L,C,I); 391(K,E,F,L,T,V,Y); 392(A,G); 394(K,E,H,M);
 15 395(D,C,E,F,I,K,M,Q,R,S,W,Y); 396(Y,C,F,K); 397(A,S);
 400(T,A,C,D,E,H,K,L,M,N,Q,R,S,V,W,Y); 401(Q,M); 402(R,F,K,L,Q,S,T,V,W,Y); 403(D,S);
 404(Y,W); 405(I,L); 407(N,A,C,D,E,G,H,Q,S); 408(P,E,H,K,M,Q,R,S,V,W,Y); 409(D,N);
 410(V,E,I,K,L,M,R,S,Y); 414(T,A,S); 416(E,C,D,F,G,H,K,L,M,N,Q,R,S,T,V);
 418(D,A,C,E,F,G,H,I,K,L,M,N,Q,R,S,T,W,Y); 419(S,A,E,G,K,P,Q,R,T,V,Y);
 20 420(T,A,C,D,E,F,G,H,I,L,P,Q,R,S,V,W,Y); 421(K,A,C,F,G,H,I,L,N,R,S,W,Y);
 422(A,C,D,E,I,K,L,M,N,P,Q,R,S,T,V,W,Y); 423(K,A,C,D,E,F,G,M,N,P,Q,S,V,W,Y);
 424(S,A); 426(L,C); 427(A,C,G,S); 429(V,C,L,M); 430(I,C,E,L,M); 431(T,A,D);
 433(G,A,H,N,S); 434(P,D,M,N,Q,R,S); 435(G,A,C,E,I,L,N,Q,R,S,T,V,Y);
 437(S,A,D,E,F,H,K,L,Q,Y); 441(Y,C,K,L,N,Q,R,S,W); 442(V,A,I,L,T);
 25 444(T,A,E,F,H,I,K,L,M,N,P,Q,V,Y); 445(S,A,C,D,H,M,R,T,V,W); 447(A,G,K,Q,R,S,T,V);
 448(G,A,D,E,F,H,K,L,N,Q,T,W,Y); 449(E,Q); 450(I,A,C,D,F,H,K,L,N,P,Q,R,T,V);
 452(Y,A,I,L,M,S,V,W); 454(L,A,C,E,F,H,I,K,M,Q,S,T,V,Y); 455(T,A,C,I,L,M,S,V);
 456(G,A,C,D,E,F,H,K,L,M,N,R,S,T,W,Y); 458(R,A,C,D,E,F,I,M,N,S,W,Y);
 459(T,A,C,D,F,G,L,S,V,W); 460(DE,H,N); 461(K,A,D,G,I,L,M,N,P,S,Y); 462(I,V);
 30 463(T,C,E,F,I,K,L,M,N,V,Y); 464(I,V); 465(G,A,M,N,P,Q);
 466(S,A,C,D,E,F,H,I,K,L,M,N,R,T,V,W,Y); 467(D,N); 469(Y,C,F,I,L,S,V); 470(A,G);
 471(T,A,D,E,F,G,H,I,K,N,P,Q,W); 474(V,C); 475(N,A,D,F,G,H,I,K,L,M,P,S,T,V);
 476(G,A,C,D,E,H,K,N,P,Q,R,S,T,V,Y); 477(G,A,D,E,H,I,K,P,Q,R,S,T,V,Y); 479(V,C,H,W);
 480(S,G); 481(VA,C,I,L); 482(W,Y); 483(V,I); 484(Q,A,H,K); and
 35 485(Q,A,D,E,F ,H,I,K,L,M,N,P,R,T,V,Y)

[0331] The productive positions in CspAmy2-v1 that fall within the previously described Productivity Scores of "2," "3," and "4," and the substitutions within those positions that are combinable, are listed below. The position numbering is based on mature CspAmy2 protein listed in SEQ ID NO: 1.

LIST B:

[0332]

1(A,E,G,I,K,N,Q,R,T,V,Y); 2(A,E,G,H,K,N,P,Q,R,S,Y); 3(T,A,D,F,G,M,P,Q,R,S);
 5(G,A,C,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,Y); 15(V,C,I,L,N,S,T); 21(Q,L,T,W);
 23(N,A,D,E,F,H,K,M,Q,S,T,V,W,Y); 27(T,D,E,F,G,H,I,K,L,M,N,Q,R,S,W,Y); 28(D,A,E,N);
 30(P,A,C,D,E,F,G,H,K,L,N,R,S,T,W,Y); 31(Y,E,F,H,K,M,R,W); 33(S,D,E,G,H,K,N,Q,R);
 35(V,H,I,M,N); 38(T,D,N,S); 42(T,A,C,I,L,M,V); 46(Y,F,M,T);
 50 49(T,A,C,D,E,F,G,H,I,K,L,M,N,S,V,Y); 50(S,D,E,K); 52(A,F,G,H,I,K,N,Q,S,T,W);
 54(V,A,C,D,E,G,I,L,N,Q,R,S,T); 68(N,A,C,E,F,M,S,Y); 70(K,N,R,W); 72(T,G,H,S);
 81(G,A,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,W,Y); 84(K,I,Q,V); 85(S,A,C,D,E,H,K,L,M,N,Q,R);
 88(N,A,D,E,H,Q,R,T); 89(T,C,D,E,H,M,N,Q,R,S,V); 92(S,A,L,M,R,V); 93(N,D,M,T);
 106(K,A,I,V); 108(G,A,K,R,S); 111(Y,A,C,D,F,G,H,K,L,M,N,Q,R,S,T,V,W);
 55 112(T,A,C,D,E,F,G,I,L,M,P,Q,R,V,W,Y); 115(V,A,I,M);
 116(T,A,C,D,E,G,H,I,K,L,M,N,S,V,W); 118(V,A,C,F,I,K,L,M,N,Q,R,S); 121(N,K,R,S);
 122(P,A,K,Q,R,T); 123(S,A,C,E,G,H,K,N,Q,R,T,Y); 127(Q,A,C,E,H,I,K,M,R,T,V,Y);
 128(E,G,I,K,S,V,Y); 129(T,A,F,G,H,I,K,L,Q,R,S,V,Y); 130(S,A,G,H,I,R,T,V);

131(G,A,F,H,I,K,M,N,P,Q,T,W); 132(E,A,C,G,H,I,L,M,N,P,Q,R,S,T,V,Y);
 133(Y,A,D,E,F,H,K,L,N,T,V); 134(N,C,D,F,G,H,M,P,Q,S,T,Y); 135(I,H,M,R,V);
 136(Q,A,F,G,H,I,K,N,T,W,Y); 138(W,A,D,F,G,H,K,L,M,P,Q,R,S,T,V,Y);
 140(G,C,D,E,F,H,I,K,L,M,N,S,T,V,Y); 142(N,A,C,D,E,F,G,H,I,K,L,M,Q,R,S,T,V,W,Y);
 144(P,A,C,D,F,G,H,I,K,L,M,N,Q,R,T,Y); 145(G,A,E,H,I,K,L,M,N,Q,R,S,T,V,W,Y);
 147(G,C,E,H,I,K,L,N,Q,R,V); 148(T,A,G,H,I,K,L,S,W,Y);
 149(T,A,C,D,E,H,I,K,L,M,N,R,S,W,Y); 150(Y,D,F,G,H,I,M,P,Q,W);
 152(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,W); 153(F,H,W,Y); 154(K,A,E,F,H,N,R,S,T);
 156(Q,D,F,G,H,I,K,L,M,S,T,V,Y); 158(F,A,C,D,E,G,H,I,L,N,P,R,S,T,V,W,Y);
 163(T,C,D,F,L,M,N,Q,S,V); 165(W,F,H,P); 166(D,A,C,G,K,M);
 167(Q,A,C,D,G,H,K,N,R,S,T,V,Y); 168(S,C,D,E,G,I,K,L,M,N,R,T,V,W);
 169(R,A,C,D,E,H,K,L,M,Q,T,W,Y); 170(S,C,G,H,N,R,T);
 171(L,F,G,H,I,K,M,N,Q,R,S,V,W,Y); 172(S,A,C,D,E,H,K,N,R,T);
 175(F,A,D,H,L,M,S,V,W,Y); 183(A,E,K,R); 189(S,A,C,D); 191(E,A,C,I,L,M,N,T);
 192(N,F,H,M,R,S,Y); 193(G,A,C,F,H,I,K,R,S,T,V); 198(L,A,C,G);
 203(I,A,C,F,H,L,M,N,Q,V,Y); 206(D,A,C,I,M,Q,Y); 207(H,A,D,E,F,K,M,N,R,S,Y);
 208(P,A,D,E,H,K,L,N,Q,R,S,T); 209(D,C,G,I,K); 211(V,C,D,E,F,I,L,M,N,Q,S,T);
 212(N,A,C,D,E,G,H,I,L,M,Q,R,S,T,V,Y); 215(K,E,F,L,M,N,Q,R,T,Y); 219(V,E,I,L);
 226(G,C,E,K,Q); 235(V,A,C,L,T); 238(I,L,M,P,Q,R); 239(K,C,M,Q);
 241(S,A,C,D,E,F,G,H,I,K,L,N,P,Q,R,T,V,W,Y); 242(F,E,I,V,Y); 243(L,A,I,M,S,T);
 244(K,A,C,H,M,N,Q,R,S,T,Y); 248(D,A,N,Q); 249(N,A,C,D,E,F,G,K,M,Q,R,S,T,W);
 251(R,A,D,K,L,M,N,Q,S,T); 252(A,C,D,F,H,M,R); 253(A,C,D,E,L,M,N,Q,R,S,T,V);
 254(T,F,I,L,M); 257(E,A,C,F,G,H,K,L,Q,R,S,V,Y); 258(M,C,F,I,L,Y); 260(T,A,L,S,Y);
 266(Q,A,C,D,E,F,G,H,I,K,L,M,N,R,S,T,W); 269(L,A,I,V);
 270(G,A,C,D,F,H,I,K,P,Q,R,S,V,W,Y); 273(N,D,E,H,I,K,L,M,Q,S);
 277(A,C,D,E,F,H,I,K,L,M,N,Q,T,W,Y); 280(N,A,C,D,E,G,H,K,L,Q,T,Y); 281(Y,A,D,G);
 283(Q,H,T,V); 284(S,E,F,H,K,M,R,W,Y); 295(Y,H,L,Q); 296(A,D,E,F,H,I,K,M,Q,R,S,T);
 299(T,E,I,R); 300(G,A,K,L,Q,R); 301(G,A,F,H,K,M,Q,R,S,T,Y); 303(Y,A,F,I,R,T,V,W);
 308(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,Y); 310(L,D,E,T); 311(N,D,E,H,K,Q,S,V);
 316(A,K,Q,S); 317(S,A,C,D,E,G,H,K,L,M,Q,T,W,Y);
 318(N,A,C,D,E,F,H,I,K,L,M,Q,R,S,V,W,Y); 320(T,A,C,D,E,G,H,K,M,N,P,Q,R,V,W,Y);
 321(K,H,R); 325(L,F,I,M,V); 343(P,A,M,W); 348(L,C,G,Q,S); 349(A,G,S,W);
 357(S,A,C,D,E,L,M,N,Q,V,Y); 358(G,D,E,Q,S); 360(Y,C,E,F,H,I,L,V); 362(S,A,C,E,I,Q,T,V);
 368(M,L,W,Y); 372(K,A,D,H,M,N,R); 374(T,A,K,N,P,Q,S); 376(T,A,G,H,K,N,Q,S,Y);
 377(R,A,C,D,G,H,I,K,L,M,S,T,V,Y); 384(S,D,E,G,H,N,P); 385(K,A,E); 388(P,C,D,I,K,L,R,S);
 391(K,E,F,L,T,V,Y); 394(K,E,H,M); 395(D,C,E,F,I,K,M,Q,R,S,W,Y); 396(Y,C,F,K);
 400(T,A,C,D,E,H,K,L,M,N,Q,R,S,V,W,Y); 402(R,F,K,L,Q,S,T,V,W,Y);
 407(N,A,C,D,E,G,H,Q,S); 408(P,E,H,K,M,Q,R,S,V,W,Y); 410(V,E,I,K,L,M,R,S,Y);
 416(E,C,D,F,G,H,K,L,M,N,Q,R,S,T,V); 418(D,A,C,E,F,G,H,I,K,L,M,N,Q,R,S,T,W,Y);
 419(S,A,E,G,K,P,Q,R,T,V,Y); 420(T,A,C,D,E,F,G,H,I,L,P,Q,R,S,V,W,Y);
 421(K,A,C,F,G,H,I,L,N,R,S,W,Y); 422(A,C,D,E,I,K,L,M,N,P,Q,R,S,T,V,W,Y);
 423(K,A,C,D,E,F,G,M,N,P,Q,S,V,W,Y); 427(A,C,G,S); 429(V,C,L,M); 430(I,C,E,L,M);
 433(G,A,H,N,S); 434(P,D,M,N,Q,R,S); 435(G,A,C,E,I,L,N,Q,R,S,T,V,Y);
 437(S,A,D,E,F,H,K,L,Q,Y); 441(Y,C,K,L,N,Q,R,S,W); 442(V,A,I,L,T);
 444(T,A,E,F,H,I,K,L,M,N,P,Q,V,Y); 445(S,A,C,D,H,M,R,T,V,W); 447(A,G,K,Q,R,S,T,V);
 448(G,A,D,E,F,H,K,L,N,Q,T,W,Y); 450(I,A,C,D,F,H,K,L,N,P,Q,R,T,V);
 452(Y,A,I,L,M,S,V,W); 454(L,A,C,E,F,H,I,K,M,Q,S,T,V,Y); 455(T,A,C,I,L,M,S,V);
 456(G,A,C,D,E,F,H,K,L,M,N,R,S,T,W,Y); 458(R,A,C,D,E,F,I,M,N,S,W,Y);
 459(T,A,C,D,F,G,L,S,V,W); 460(DE,H,N); 461(K,A,D,G,I,L,M,N,P,S,Y);
 463(T,C,E,F,I,K,L,M,N,V,Y); 465(G,A,M,N,P,Q); 466(S,A,C,D,E,F,H,I,K,L,M,N,R,T,V,W,Y);
 469(Y,C,F,I,L,S,V); 471(T,A,D,E,F,G,H,I,K,N,P,Q,W); 475(N,A,D,F,G,H,I,K,L,M,P,S,T,V);
 476(G,A,C,D,E,H,K,N,P,Q,R,S,T,V,Y); 477(G,A,D,E,H,I,K,P,Q,R,S,T,V,Y); 479(V,C,H,W);
 481(VA,C,I,L); 484(Q,A,H,K); and 485(Q,A,D,E,F,H,I,K,L,M,N,P,R,T,V,Y)

[0333] The productive positions in CspAmy2-v1 that fall within the previously described Productivity Scores of "3" and "4," and the substitutions within those positions that are combinable, are listed below. The position numbering is based on mature CspAmy2 protein listed in SEQ ID NO: 1.

LIST C:

[0334]

- 5 1(A,E,G,I,K,N,Q,R,T,V,Y); 2(A,E,G,H,K,N,P,Q,R,S,Y); 3(T,A,D,F,G,M,P,Q,R,S);
 5(G,A,C,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,Y); 15(V,C,I,L,N,S,T);
 23(N,A,D,E,F,H,K,M,Q,S,T,V,W,Y); 27(T,D,E,F,G,H,I,K,L,M,N,Q,R,S,W,Y);
 30(P,A,C,D,E,F,G,H,K,L,N,R,S,T,W,Y); 31(Y,E,F,H,K,M,R,W); 33(S,D,E,G,H,K,N,Q,R);
 42(T,A,C,I,L,M,V); 49(T,A,C,D,E,F,G,H,I,K,L,M,N,S,V,Y); 52(A,F,G,H,I,K,N,Q,S,T,W);
 10 54(V,A,C,D,E,G,I,L,N,Q,R,S,T); 68(N,A,C,E,F,M,S,Y);
 81(G,A,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,W,Y); 85(S,A,C,D,E,H,K,L,M,N,Q,R);
 88(N,A,D,E,H,Q,R,T); 89(T,C,D,E,H,M,N,Q,R,S,V); 92(S,A,L,M,R,V);
 111(Y,A,C,D,F,G,H,K,L,M,N,Q,R,S,T,V,W); 112(T,A,C,D,E,F,G,I,L,M,P,Q,R,V,W,Y);
 116(T,A,C,D,E,G,H,I,K,L,M,N,S,V,W); 118(V,A,C,F,I,K,L,M,N,Q,R,S); 122(P,A,K,Q,R,T);
 15 123(S,A,C,E,G,H,K,N,Q,R,T,Y); 127(Q,A,C,E,H,I,K,M,R,T,V,Y); 128(E,G,I,K,S,V,Y);
 129(T,A,F,G,H,I,K,L,Q,R,S,V,Y); 130(S,A,G,H,I,R,T,V); 131(G,A,F,H,I,K,M,N,P,Q,T,W);
 132(E,A,C,G,H,I,L,M,N,P,Q,R,S,T,V,Y); 133(Y,A,D,E,F,H,K,L,N,T,V);
 134(N,C,D,F,G,H,M,P,Q,S,T,Y); 136(Q,A,F,G,H,I,K,N,T,W,Y);
 138(W,A,D,F,G,H,K,L,M,P,Q,R,S,T,V,Y); 140(G,C,D,E,F,H,I,K,L,M,N,S,T,V,Y);
 20 142(N,A,C,D,E,F,G,H,I,K,L,M,Q,R,S,T,V,W,Y); 144(P,A,C,D,F,G,H,I,K,L,M,N,Q,R,T,Y);
 145(G,A,E,H,I,K,L,M,N,Q,R,S,T,V,W,Y); 147(G,C,E,H,I,K,L,N,Q,R,V);
 148(T,A,G,H,I,K,L,S,W,Y); 149(T,A,C,D,E,H,I,K,L,M,N,R,S,W,Y);
 150(Y,D,F,G,H,I,M,P,Q,W); 152(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,W);
 154(K,A,E,F,H,N,R,S,T); 156(Q,D,F,G,H,I,K,L,M,S,T,V,Y);
 25 158(F,A,C,D,E,G,H,I,L,N,P,R,S,T,V,W,Y); 163(T,C,D,F,L,M,N,Q,S,V); 166(D,A,C,G,K,M);
 167(Q,A,C,D,G,H,K,N,R,S,T,V,Y); 168(S,C,D,E,G,I,K,L,M,N,R,T,V,W);
 169(R,A,C,D,E,H,K,L,M,Q,T,W,Y); 170(S,C,G,H,N,R,T);
 171(L,F,G,H,I,K,M,N,Q,R,S,V,W,Y); 172(S,A,C,D,E,H,K,N,R,T);
 175(F,A,D,H,L,M,S,V,W,Y); 191(E,A,C,I,L,M,N,T); 192(N,F,H,M,R,S,Y);
 30 193(G,A,C,F,H,I,K,R,S,T,V); 203(I,A,C,F,H,L,M,N,Q,V,Y); 206(D,A,C,I,M,Q,Y);
 207(H,A,D,E,F,K,M,N,R,S,Y); 208(P,A,D,E,H,K,L,N,Q,R,S,T); 209(D,C,G,I,K);
 211(V,C,D,E,F,I,L,M,N,Q,S,T); 212(N,A,C,D,E,G,H,I,L,M,Q,R,S,T,V,Y);
 215(K,E,F,L,M,N,Q,R,T,Y); 241(S,A,C,D,E,F,G,H,I,K,L,N,P,Q,R,T,V,W,Y);
 244(K,A,C,H,M,N,Q,R,S,T,Y); 249(N,A,C,D,E,F,G,K,M,Q,R,S,T,W);
 35 251(R,A,D,K,L,M,N,Q,S,T); 252(A,C,D,F,H,M,R); 253(A,C,D,E,L,M,N,Q,R,S,T,V);
 257(E,A,C,F,G,H,K,L,Q,R,S,V,Y); 258(M,C,F,I,L,Y);
 266(Q,A,C,D,E,F,G,H,I,K,L,M,N,R,S,T,W); 270(G,A,C,D,F,H,I,K,P,Q,R,S,V,W,Y);
 273(N,D,E,H,I,K,L,M,Q,S); 277(A,C,D,E,F,H,I,K,L,M,N,Q,T,W,Y);
 280(N,A,C,D,E,G,H,K,L,Q,T,Y); 284(S,E,F,H,K,M,R,W,Y); 296(A,D,E,F,H,I,K,M,Q,R,S,T);
 40 301(G,A,F,H,K,M,Q,R,S,T,Y); 303(Y,A,F,I,R,T,V,W);
 308(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,Y); 311(N,D,E,H,K,Q,S,V);
 317(S,A,C,D,E,G,H,K,L,M,Q,T,W,Y); 318(N,A,C,D,E,F,H,I,K,L,M,Q,R,S,V,W,Y);
 320(T,A,C,D,E,G,H,K,M,N,P,Q,R,V,W,Y); 357(S,A,C,D,E,L,M,N,Q,V,Y);
 360(Y,C,E,F,H,I,L,V); 362(S,A,C,E,I,Q,T,V); 372(K,A,D,H,M,N,R); 374(T,A,K,N,P,Q,S);
 45 376(T,A,G,H,K,N,Q,S,Y); 377(R,A,C,D,G,H,I,K,L,M,S,T,V,Y); 384(S,D,E,G,H,N,P);
 388(P,C,D,I,K,L,R,S); 391(K,E,F,L,T,V,Y); 395(D,C,E,F,I,K,M,Q,R,S,W,Y); 396(Y,C,F,K);
 400(T,A,C,D,E,H,K,L,M,N,Q,R,S,V,W,Y); 402(R,F,K,L,Q,S,T,V,W,Y);
 407(N,A,C,D,E,G,H,Q,S); 408(P,E,H,K,M,Q,R,S,V,W,Y); 410(V,E,I,K,L,M,R,S,Y);
 416(E,C,D,F,G,H,K,L,M,N,Q,R,S,T,V); 418(D,A,C,E,F,G,H,I,K,L,M,N,Q,R,S,T,W,Y);
 50 419(S,A,E,G,K,P,Q,R,T,V,Y); 420(T,A,C,D,E,F,G,H,I,L,P,Q,R,S,V,W,Y);
 421(K,A,C,F,G,H,I,L,N,R,S,W,Y); 422(A,C,D,E,I,K,L,M,N,P,Q,R,S,T,V,W,Y);
 423(K,A,C,D,E,F,G,M,N,P,Q,S,V,W,Y); 434(P,D,M,N,Q,R,S);
 435(G,A,C,E,I,L,N,Q,R,S,T,V,Y); 437(S,A,D,E,F,H,K,L,Q,Y); 441(Y,C,K,L,N,Q,R,S,W);
 444(T,A,E,F,H,I,K,L,M,N,P,Q,V,Y); 445(S,A,C,D,H,M,R,T,V,W); 447(A,G,K,Q,R,S,T,V);
 55 448(G,A,D,E,F,H,K,L,N,Q,T,W,Y); 450(I,A,C,D,F,H,K,L,N,P,Q,R,T,V);
 452(Y,A,I,L,M,S,V,W); 454(L,A,C,E,F,H,I,K,M,Q,S,T,V,Y); 455(T,A,C,I,L,M,S,V);
 456(G,A,C,D,E,F,H,K,L,M,N,R,S,T,W,Y); 458(R,A,C,D,E,F,I,M,N,S,W,Y);
 459(T,A,C,D,F,G,L,S,V,W); 461(K,A,D,G,I,L,M,N,P,S,Y); 463(T,C,E,F,I,K,L,M,N,V,Y);

465(G,A,M,N,P,Q); 466(S,A,C,D,E,F,H,I,K,L,M,N,R,T,V,W,Y); 469(Y,C,F,I,L,S,V);
 471(T,A,D,E,F,G,H,I,K,N,P,Q,W); 475(N,A,D,F,G,H,I,K,L,M,P,S,T,V);
 476(G,A,C,D,E,H,K,N,P,Q,R,S,T,V,Y); 477(G,A,D,E,H,I,K,P,Q,R,S,T,V,Y); and
 485(Q,A,D,E,F,H,I,K,L,M,N,P,R,T,V,Y)

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[0335] The productive positions in CspAmy2-v1 that fall within the previously described Productivity Scores of "4," and the substitutions within those positions that are combinable, are listed below. The position numbering is based on mature CspAmy2 protein listed in SEQ ID NO: 1.

10 LIST D:

[0336]

1(A,E,G,I,K,N,Q,R,T,V,Y); 2(A,E,G,H,K,N,P,Q,R,S,Y); 3(T,A,D,F,G,M,P,Q,R,S);
 5(G,A,C,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,Y); 23(N,A,D,E,F,H,K,M,Q,S,T,V,W,Y);
 27(T,D,E,F,G,H,I,K,L,M,N,Q,R,S,W,Y); 30(P,A,C,D,E,F,G,H,K,L,N,R,S,T,W,Y);
 49(T,A,C,D,E,F,G,H,I,K,L,M,N,S,V,Y); 52(A,F,G,H,I,K,N,Q,S,T,W);
 54(V,A,C,D,E,G,I,L,N,Q,R,S,T); 68(N,A,C,E,F,M,S,Y);
 81(G,A,D,E,F,H,I,K,L,M,N,P,Q,R,S,T,V,W,Y); 85(S,A,C,D,E,H,K,L,M,N,Q,R);
 89(T,C,D,E,H,M,N,Q,R,S,V); 111(Y,A,C,D,F,G,H,K,L,M,N,Q,R,S,T,V,W);
 112(T,A,C,D,E,F,G,I,L,M,P,Q,R,V,W,Y); 116(T,A,C,D,E,G,H,I,K,L,M,N,S,V,W);
 118(V,A,C,F,I,K,L,M,N,Q,R,S); 123(S,A,C,E,G,H,K,N,Q,R,T,Y);
 127(Q,A,C,E,H,I,K,M,R,T,V,Y); 128(E,G,I,K,S,V,Y); 129(T,A,F,G,H,I,K,L,Q,R,S,V,Y);
 131(G,A,F,H,I,K,M,N,P,Q,T,W); 132(E,A,C,G,H,I,L,M,N,P,Q,R,S,T,V,Y);
 133(Y,A,D,E,F,H,K,L,N,T,V); 134(N,C,D,F,G,H,M,P,Q,S,T,Y); 136(Q,A,F,G,H,I,K,N,T,W,Y);
 138(W,A,D,F,G,H,K,L,M,P,Q,R,S,T,V,Y); 140(G,C,D,E,F,H,I,K,L,M,N,S,T,V,Y);
 142(N,A,C,D,E,F,G,H,I,K,L,M,Q,R,S,T,V,W,Y); 144(P,A,C,D,F,G,H,I,K,L,M,N,Q,R,T,Y);
 145(G,A,E,H,I,K,L,M,N,Q,R,S,T,V,W,Y); 147(G,C,E,H,I,K,L,N,Q,R,V);
 148(T,A,G,H,I,K,L,S,W,Y); 149(T,A,C,D,E,H,I,K,L,M,N,R,S,W,Y);
 150(Y,D,F,G,H,I,M,P,Q,W); 152(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,W);
 156(Q,D,F,G,H,I,K,L,M,S,T,V,Y); 158(F,A,C,D,E,G,H,I,L,N,P,R,S,T,V,W,Y);
 163(T,C,D,F,L,M,N,Q,S,V); 167(Q,A,C,D,G,H,K,N,R,S,T,V,Y);
 168(S,C,D,E,G,I,K,L,M,N,R,T,V,W); 169(R,A,C,D,E,H,K,L,M,Q,T,W,Y);
 171(L,F,G,H,I,K,M,N,Q,R,S,V,W,Y); 172(S,A,C,D,E,H,K,N,R,T);
 193(G,A,C,F,H,I,K,R,S,T,V); 203(I,A,C,F,H,L,M,N,Q,V,Y); 207(H,A,D,E,F,K,M,N,R,S,Y);
 208(P,A,D,E,H,K,L,N,Q,R,S,T); 211(V,C,D,E,F,I,L,M,N,Q,S,T);
 212(N,A,C,D,E,G,H,I,L,M,Q,R,S,T,V,Y); 215(K,E,F,L,M,N,Q,R,T,Y);
 241(S,A,C,D,E,F,G,H,I,K,L,N,P,Q,R,T,V,W,Y); 244(K,A,C,H,M,N,Q,R,S,T,Y);
 249(N,A,C,D,E,F,G,K,M,Q,R,S,T,W); 251(R,A,D,K,L,M,N,Q,S,T);
 253(A,C,D,E,L,M,N,Q,R,S,T,V); 257(E,A,C,F,G,H,K,L,Q,R,S,V,Y);
 266(Q,A,C,D,E,F,G,H,I,K,L,M,N,R,S,T,W); 270(G,A,C,D,F,H,I,K,P,Q,R,S,V,W,Y);
 273(N,D,E,H,I,K,L,M,Q,S); 277(A,C,D,E,F,H,I,K,L,M,N,Q,T,W,Y);
 280(N,A,C,D,E,G,H,K,L,Q,T,Y); 284(S,E,F,H,K,M,R,W,Y); 296(A,D,E,F,H,I,K,M,Q,R,S,T);
 301(G,A,F,H,K,M,Q,R,S,T,Y); 308(N,A,C,D,E,F,G,H,L,M,Q,R,S,T,V,Y);
 317(S,A,C,D,E,G,H,K,L,M,Q,T,W,Y); 318(N,A,C,D,E,F,H,I,K,L,M,Q,R,S,V,W,Y);
 320(T,A,C,D,E,G,H,K,M,N,P,Q,R,V,W,Y); 357(S,A,C,D,E,L,M,N,Q,V,Y);
 360(Y,C,E,F,H,I,L,V); 377(R,A,C,D,G,H,I,K,L,M,S,T,V,Y); 384(S,D,E,G,H,N,P);
 388(P,C,D,I,K,L,R,S); 395(D,C,E,F,I,K,M,Q,R,S,W,Y);
 400(T,A,C,D,E,H,K,L,M,N,Q,R,S,V,W,Y); 408(P,E,H,K,M,Q,R,S,V,W,Y);
 416(E,C,D,F,G,H,K,L,M,N,Q,R,S,T,V); 418(D,A,C,E,F,G,H,I,K,L,M,N,Q,R,S,T,W,Y);
 419(S,A,E,G,K,P,Q,R,T,V,Y); 420(T,A,C,D,E,F,G,H,I,L,P,Q,R,S,V,W,Y);
 421(K,A,C,F,G,H,I,L,N,R,S,W,Y); 422(A,C,D,E,I,K,L,M,N,P,Q,R,S,T,V,W,Y);
 423(K,A,C,D,E,F,G,M,N,P,Q,S,V,W,Y); 434(P,D,M,N,Q,R,S);
 435(G,A,C,E,I,L,N,Q,R,S,T,V,Y); 437(S,A,D,E,F,H,K,L,Q,Y);
 444(T,A,E,F,H,I,K,L,M,N,P,Q,V,Y); 445(S,A,C,D,H,M,R,T,V,W);
 448(G,A,D,E,F,H,K,L,N,Q,T,W,Y); 450(I,A,C,D,F,H,K,L,N,P,Q,R,T,V);
 454(L,A,C,E,F,H,I,K,M,Q,S,T,V,Y); 456(G,A,C,D,E,F,H,K,L,M,N,R,S,T,W,Y);
 458(R,A,C,D,E,F,I,M,N,S,W,Y); 459(T,A,C,D,F,G,L,S,V,W); 461(K,A,D,G,I,L,M,N,P,S,Y);

55

463(T,C,E,F,I,K,L,M,N,V,Y); 466(S,A,C,D,E,F,H,I,K,L,M,N,R,T,V,W,Y);
 471(T,A,D,E,F,G,H,I,K,N,P,Q,W); 475(N,A,D,F,G,H,I,K,L,M,P,S,T,V);
 476(G,A,C,D,E,H,K,N,P,Q,R,S,T,V,Y); 477(G,A,D,E,H,I,K,P,Q,R,S,T,V,Y); and
 485(Q,A,D,E,F,H,I,K,L,M,N,P,R,T,V,Y)

[0337] The productive positions in CspAmy2-v1 suitable for charge or hydrophobicity modifications are listed below. The position numbering is based on mature CspAmy2 protein listed in SEQ ID NO: 1.

LIST E:

[0338] 1, 5, 15, 23, 30, 31, 49, 68, 111, 112, 116, 123, 127, 128, 129, 131, 132, 134, 140, 142, 144, 147, 150, 152, 153, 168, 170, 171, 183, 187, 192, 203, 207, 209, 211, 212, 232, 241, 243, 244, 253, 266, 277, 280, 300, 301, 308, 320, 357, 362, 377, 388, 400, 402, 408, 416, 420, 423, 448, 450, 454, 455, 456, 458, 466, 475, and 485

[0339] The productive positions and the substitutions within those positions in CspAmy2-v1 suitable for charge or hydrophobicity modifications are listed below. The position numbering is based on mature CspAmy2 protein listed in SEQ ID NO: 1

LIST F:

[0340] 1(A,K,V,Y); 5(G,F,V); 15(V,S); 23(N,K); 30(P,E,K,L,W); 31(Y,F,K,W); 49(T,I); 68(N,Y); 111(Y,D,Q,S,T,W); 112(T,I,W); 116(T,L); 123(S,K); 127(Q,I); 128(E,I,V); 129(T,I); 131(G,H,K); 132(E,G,H,I,M,P,R,T,V,Y); 134(N,D,F,M,Y); 140(G,E,F,H,K); 142(N,D,I,R); 144(P,G,K,L); 147(G,C,E,H,L,R,V); 150(Y,W); 152(N,D,E,L,R); 153(F,H,Y); 168(S,L); 170(S,R); 171(L,H,N,Q,R); 183(A,K); 187(E,P); 192(N,F,Y); 203(I,C); 207(H,E,F); 209(D,G); 211(V,D,E,N,Q); 212(N,D,E); 232(L,R); 241(S,D,E,I,L,W,Y); 243(L,S); 244(K,C); 253(A,R); 266(Q,I,R,W); 277(A,F,L,Y); 280(N,L); 300(G,R); 301(G,H); 308(N,D,E,F,L,R,V,Y); 320(T,E,W); 357(S,E); 362(S,I,V); 377(R,G,H); 388(P,K); 400(T,E,K,L,W,Y); 402(R,F); 408(P,E,R,W); 416(E,C,F,G,H,K,L,N,R,S,V); 420(T,D,E,I,L,R,W); 423(K,E,F,G,M,N,Q,S,V,Y); 448(G,F,H,K,W); 450(I,K,R,T); 454(L,C); 455(T,L,M); 456(G,F,W); 458(R,A,C,D,E,F,I,S); 466(S,I,L); 475(N,D); and 485(Q,E,K,L,Y,Y)

Claims

1. A variant α -amylase polypeptide derived from a parental α -amylase polypeptide, comprising at least one combinable mutation at a productive amino acid position; wherein:

- (i) the or each combinable mutation is a mutation that improves at least one desirable enzymatic and biochemical property of the variant α -amylase compared to the parental α -amylase, while not significantly decreasing either expression, activity, or stability of the variant α -amylase, compared to the parental α -amylase,
- (ii) the or each productive position is an amino acid position that can be substituted with a plurality of different amino acid residues, each of which substitutions result in a variant α -amylase that meets the requirements of (i), and
- (iii) the or each combinable mutation is listed in Lists A, B, C, D, E, or F, or in Table D, which uses SEQ ID NO: 1 for numbering, and

wherein the variant α -amylase has at least 60%, at least 70%, at least 80% or at least 90% amino acid sequence identity to the amino acid sequence of SEQ ID NO: 1 or SEQ ID NO: 2, wherein the variant amylase includes a combinable mutation at the productive position corresponding to position 203 of SEQ ID NO: 1 which is substitution to Tyr.

2. The variant amylase of claim 1, wherein the combinable mutation which is substitution to Tyr at position 203 SEQ ID NO: 1 for numbering produces a variant amylase wherein the minimum performance indices (PI) relative to the parental amylase for (i) protein expression, (ii) activity, and (iii) detergent stability or thermostability are greater than or equal to 0.9, and the PI for any one of (i), (ii), or (iii) that is greater than or equal to 1.0, whereby the PI for the variant is calculated by comparing the performance or stability of the variant (measured value) and the parental amylase (theoretical value) at the same protein concentration.

3. The variant amylase of claim 1, wherein the combinable mutation which is substitution to Tyr at position 203 SEQ ID NO: 1 for numbering produces a variant amylase wherein the minimum performance indices (PI) relative to the

parental amylase for (i) protein expression, (ii) activity, and (iii) detergent stability or thermostability are greater than or equal to 0.8, and the PI for any one of (i), (ii), or (iii) that is greater than or equal to 1.2, whereby the PI for the variant is calculated by comparing the performance or stability of the variant (measured value) and the parental amylase (theoretical value) at the same protein concentration.

4. The variant amylase of claim 1, wherein the combinable mutation which is substitution to Tyr at position 203 SEQ ID NO: 1 for numbering produces a variant amylase wherein the minimum performance indices (PI) relative to the parental amylase for (i) protein expression, (ii) activity, and (iii) detergent stability or thermostability are greater than or equal to 0.5, and the PI for any one of (i), (ii), or (iii) that is greater than or equal to 1.5, whereby the PI for the variant is calculated by comparing the performance or stability of the variant (measured value) and the parental amylase (theoretical value) at the same protein concentration.
5. The variant amylase of any of the preceding claims, wherein the or each combinable mutation has a suitability score of +++, +++, or +++, referring to Table C.
6. The variant amylase of any of the preceding claims, wherein the or each combinable mutation has a suitability score of +++, or +++, referring to Table C.
7. The variant amylase of any of the preceding claims, wherein the or each combinable mutation has a suitability score of +++, referring to Table C.
8. The variant amylase of any of the preceding claims, wherein the or each combinable mutation has a productivity score of 1 or 2, referring to Table B.
9. The variant amylase of any of the preceding claims, having a plurality of combinable mutations.
10. The variant amylase of any of the preceding claims, further comprising a deletion corresponding to a residue selected from the group consisting of Arg-178, Gly-179, Thr-180, and Gly-181, using SEQ ID NO: 1 for numbering.
11. The variant amylase of any of the preceding claims, further comprising deletions corresponding to residues Arg-178 and Gly-179, using SEQ ID NO: 1 for numbering.
12. A composition comprising the variant amylase of any of claims 1-11, further comprising a surfactant, wherein the composition is optionally a detergent composition, e.g. a laundry detergent or a laundry detergent additive.
13. A method of saccharifying a composition comprising starch to produce a composition comprising glucose, wherein the method comprises:
 - (i) contacting the solution comprising starch with effective amount of the variant amylase of any of the claims 1-11; and
 - (ii) saccharifying the solution comprising starch to produce the composition comprising glucose; wherein the variant amylase catalyzes the saccharification of the starch solution to glucose,
 optionally wherein the composition comprising starch comprises liquefied starch, gelatinized starch, or granular starch.
14. The method of claim 13, wherein saccharification is conducted at a temperature range of about 30°C to about 75°C, optionally wherein the temperature range is 47°C-74°C; and/or wherein saccharification is conducted over a pH range of pH 2.0-7.5, optionally wherein the pH range is pH 3.5-5.5, further optionally wherein the pH range is pH 3.5-4.5.
15. The method of claim 13 or 14, further comprising fermenting the glucose composition to produce an end of fermentation (EOF) product.

Patentansprüche

1. α -Amylase-Polypeptid-Variante abgeleitet von einem parentalen α -Amylase-Polypeptid, umfassend mindestens

eine kombinierbare Mutation an einer produktiven Aminosäureposition; wobei:

(i) die oder jede kombinierbare Mutation eine Mutation ist, die mindestens eine wünschenswerte enzymatische und biochemische Eigenschaft der α -Amylase-Variante im Vergleich zur parentalen α -Amylase verbessert, während sie weder Expression, Aktivität noch Stabilität der α -Amylase-Variante im Vergleich zur parentalen α -Amylase signifikant verringert,

(ii) die oder jede produktive Position eine Aminosäureposition ist, die mit einer Vielzahl von verschiedenen Aminosäureresten substituiert werden kann, wobei jede der Substitutionen in einer α -Amylase-Variante resultiert, die die Anforderungen von (i) erfüllt, und

(iii) die oder jede kombinierbare Mutation in den Listen A, B, C, D, E oder F aufgelistet ist, oder in Tabelle D, die SEQ ID NO: 1 zur Nummerierung verwendet, und

wobei die α -Amylase-Variante mindestens 60 %, mindestens 70 %, mindestens 80 % oder mindestens 90 % Aminosäure-Sequenzidentität mit der Aminosäuresequenz von SEQ ID NO: 1 oder SEQ ID NO: 2 hat, wobei die Amylase-Variante eine kombinierbare Mutation an der produktiven Position enthält, die Position 203 von SEQ ID NO: 1 entspricht, die Substitution zu Tyr ist.

2. Amylase-Variante nach Anspruch 1, wobei die kombinierbare Mutation, die Substitution zu Tyr an Position 203 SEQ ID NO: 1 zur Nummerierung ist, eine Amylase-Variante erzeugt, wobei die minimalen Leistungsindizes (PI) in Bezug auf die parentale Amylase für (i) Proteinexpression, (ii) Aktivität und (iii) Detergens-Stabilität oder Thermostabilität größer als oder gleich 0,9 sind, und der PI für jedes von (i), (ii) oder (iii), das größer als oder gleich 1,0 ist, wodurch der PI für die Variante durch Vergleichen der Leistung oder Stabilität der Variante (gemessener Wert) und der parentalen Amylase (theoretischer Wert) bei der gleichen Proteinkonzentration berechnet wird.

3. Amylase-Variante nach Anspruch 1, wobei die kombinierbare Mutation, die Substitution zu Tyr an Position 203 SEQ ID NO: 1 zur Nummerierung ist, eine Amylase-Variante erzeugt, wobei die minimalen Leistungsindizes (PI) in Bezug auf die parentale Amylase für (i) Proteinexpression, (ii) Aktivität und (iii) Detergens-Stabilität oder Thermostabilität größer als oder gleich 0,8 sind, und der PI für jedes von (i), (ii) oder (iii), das größer als oder gleich 1,2 ist, wodurch der PI für die Variante durch Vergleichen der Leistung oder Stabilität der Variante (gemessener Wert) und der parentalen Amylase (theoretischer Wert) bei der gleichen Proteinkonzentration berechnet wird.

4. Amylase-Variante nach Anspruch 1, wobei die kombinierbare Mutation, die Substitution zu Tyr an Position 203 SEQ ID NO: 1 zur Nummerierung ist, eine Amylase-Variante erzeugt, wobei die minimalen Leistungsindizes (PI) in Bezug auf die parentale Amylase für (i) Proteinexpression, (ii) Aktivität und (iii) Detergens-Stabilität oder Thermostabilität größer als oder gleich 0,5 sind, und der PI für jedes von (i), (ii) oder (iii), das größer als oder gleich 1,5 ist, wodurch der PI für die Variante durch Vergleichen der Leistung oder Stabilität der Variante (gemessener Wert) und der parentalen Amylase (theoretischer Wert) bei der gleichen Proteinkonzentration berechnet wird.

5. Amylase-Variante nach einem der vorherigen Ansprüche, wobei die oder jede kombinierbare Mutation eine Eigenschaftsbewertung von +++, +++, oder +++++ unter Bezugnahme auf Tabelle C hat.

6. Amylase-Variante nach einem der vorherigen Ansprüche, wobei die oder jede kombinierbare Mutation eine Eigenschaftsbewertung von ++++ oder +++++ unter Bezugnahme auf Tabelle C hat.

7. Amylase-Variante nach einem der vorherigen Ansprüche, wobei die oder jede kombinierbare Mutation eine Eigenschaftsbewertung von +++++ unter Bezugnahme auf Tabelle C hat.

8. Amylase-Variante nach einem der vorherigen Ansprüche, wobei die oder jede kombinierbare Mutation eine Produktivitätsbewertung von 1 oder 2 unter Bezugnahme auf Tabelle B hat.

9. Amylase-Variante nach einem der vorherigen Ansprüche mit einer Vielzahl von kombinierbaren Mutationen.

10. Amylase-Variante nach einem der vorherigen Ansprüche, ferner umfassend eine Deletion entsprechend einem Rest, ausgewählt aus der Gruppe bestehend aus Arg-178, Gly-179, Thr-180 und Gly-181, unter Verwendung von SEQ ID NO: 1 zur Nummerierung.

11. Amylase-Variante nach einem der vorherigen Ansprüche, ferner umfassend Deletionen entsprechend Resten Arg-178 und Gly-179, unter Verwendung von SEQ ID NO: 1 zur Nummerierung.

12. Zusammensetzung, umfassend die Amylase-Variante nach einem der Ansprüche 1-11, ferner umfassend ein Benetzungsmittel, wobei die Zusammensetzung optional eine Detergens-Zusammensetzung ist, z. B. ein Wäscherei-Detergens oder ein Wäscherei-Detergensadditiv.

13. Verfahren zum Saccharifizieren einer Zusammensetzung, umfassend Stärke, um eine Zusammensetzung umfassend Glucose zu erzeugen, wobei das Verfahren umfasst:

(i) Inkontaktbringen der Lösung umfassend Stärke mit einer wirksamen Menge der Amylase-Variante nach einem der Ansprüche 1-11; und

(ii) Saccharifizieren der Lösung umfassend Stärke, um die Zusammensetzung umfassend Glucose zu erzeugen; wobei die Amylase-Variante die Saccharifizierung der Stärkelösung zu Glucose katalysiert,

optional wobei die Zusammensetzung umfassend Stärke verflüssigte Stärke, gelatinierte Stärke oder granulare Stärke umfasst.

14. Verfahren nach Anspruch 13, wobei die Saccharifizierung in einem Temperaturbereich von ca. 30 °C bis ca. 75 °C durchgeführt wird, optional wobei der Temperaturbereich 47 °C - 74 °C ist; und/oder wobei die Saccharifizierung über einen pH-Bereich von pH 2,0-7,5 durchgeführt wird, optional wobei der pH-Bereich pH 3,5-5,5 ist, ferner optional wobei der pH-Bereich pH 3,5-4,5 ist.

15. Verfahren nach Anspruch 13 oder 14, ferner umfassend ein Fermentieren der Glucose-Zusammensetzung, um ein Produkt am Ende der Fermentation (end of fermentation, EOF) zu erzeugen.

Revendications

1. Polypeptide variant d' α -amylase dérivé d'un polypeptide d' α -amylase parentale, comprenant au moins une mutation pouvant être combinée à une position d'acide aminé productive ; dans lequel :

(i) la ou chaque mutation pouvant être combinée est une mutation qui améliore au moins une propriété enzymatique et biochimique souhaitable du variant d' α -amylase par comparaison à l' α -amylase parentale, tout en ne réduisant pas significativement l'expression, l'activité, ou la stabilité du variant d' α -amylase, par comparaison à l' α -amylase parentale,

(ii) la ou chaque position productive est une position d'acide aminé qui peut être substituée avec une pluralité de résidus d'acides aminés différents, chacune des substitutions donnant lieu à un variant d' α -amylase qui satisfait les exigences de (i), et

(iii) la ou chaque mutation pouvant être combinée est listée dans les Listes A, B, C, D, E, ou F, ou dans le Tableau D, qui utilise SEQ ID NO: 1 pour la numérotation, et

où le variant d' α -amylase présente au moins 60 %, au moins 70 %, au moins 80 % ou au moins 90 % d'identité de séquence d'acides aminés avec la séquence d'acides aminés de SEQ ID NO: 1 ou SEQ ID NO: 2,

où le variant d'amylase comprend une mutation pouvant être combinée à la position productive correspondant à la position 203 de SEQ ID NO: 1 qui est une substitution à Tyr.

2. Variant d'amylase selon la revendication 1, dans lequel la mutation pouvant être combinée qui est une substitution à Tyr à la position 203 de SEQ ID NO: 1 pour la numérotation produit un variant d'amylase dans lequel les indices de performance (IP) minimum par rapport à l'amylase parentale pour (i) l'expression protéique, (ii) l'activité, et (iii) la stabilité aux détergents ou la thermostabilité sont supérieurs ou égaux à 0,9, et l'IP pour l'un quelconque de (i), (ii), ou (iii) qui est supérieur ou égal à 1,0, moyennant quoi l'IP pour le variant est calculé en comparant la performance ou la stabilité du variant (valeur mesurée) et de l'amylase parentale (valeur théorique) à la même concentration protéique.

3. Variant d'amylase selon la revendication 1, dans lequel la mutation pouvant être combinée qui est une substitution à Tyr à la position 203 de SEQ ID NO: 1 pour la numérotation produit un variant d'amylase dans lequel les indices de performance (IP) minimum par rapport à l'amylase parentale pour (i) l'expression protéique, (ii) l'activité, et (iii) la stabilité aux détergents ou la thermostabilité sont supérieurs ou égaux à 0,8, et l'IP pour l'un quelconque de (i), (ii), ou (iii) qui est supérieur ou égal à 1,2,

moyennant quoi l'IP pour le variant est calculé en comparant la performance ou la stabilité du variant (valeur mesurée) et de l'amylase parentale (valeur théorique) à la même concentration protéique.

4. Variant d'amylase selon la revendication 1, dans lequel la mutation pouvant être combinée qui est une substitution à Tyr à la position 203 de SEQ ID NO: 1 pour la numérotation produit un variant d'amylase dans lequel les indices de performance (IP) minimum par rapport à l'amylase parentale pour (i) l'expression protéique, (ii) l'activité, et (iii) la stabilité aux détergents ou la thermostabilité sont supérieurs ou égaux à 0,5, et l'IP pour l'un quelconque de (i), (ii), ou (iii) qui est supérieur ou égal à 1,5, moyennant quoi l'IP pour le variant est calculé en comparant la performance ou la stabilité du variant (valeur mesurée) et de l'amylase parentale (valeur théorique) à la même concentration protéique.
5. Variant d'amylase selon l'une quelconque des revendications précédentes, dans lequel la ou chaque mutation pouvant être combinée possède un score d'adéquation de +++, +++, ou +++, en se référant au Tableau C.
6. Variant d'amylase selon l'une quelconque des revendications précédentes, dans lequel la ou chaque mutation pouvant être combinée possède un score d'adéquation de +++, ou +++, en se référant au Tableau C.
7. Variant d'amylase selon l'une quelconque des revendications précédentes, dans lequel la ou chaque mutation pouvant être combinée possède un score d'adéquation +++, en se référant au Tableau C.
8. Variant d'amylase selon l'une quelconque des revendications précédentes, dans lequel la ou chaque mutation pouvant être combinée possède un score de productivité de 1 ou 2, en se référant au Tableau B.
9. Variant d'amylase selon l'une quelconque des revendications précédentes, possédant une pluralité de mutations pouvant être combinées.
10. Variant d'amylase selon l'une quelconque des revendications précédentes, comprenant en outre une délétion correspondant à un résidu sélectionné dans le groupe consistant en Arg-178, Gly-179, Thr-180, et Gly-181, en utilisant SEQ ID NO: 1 pour la numérotation.
11. Variant d'amylase selon l'une quelconque des revendications précédentes, comprenant en outre des délétions correspondant aux résidus Arg-178 et Gly-179, en utilisant SEQ ID NO: 1 pour la numérotation.
12. Composition comprenant le variant d'amylase selon l'une quelconque des revendications 1 à 11, comprenant en outre un tensioactif, où la composition est facultativement une composition détergente, par exemple un détergent de blanchisserie ou un additif de détergent de blanchisserie.
13. Procédé de saccharification d'une composition comprenant de l'amidon afin de produire une composition comprenant du glucose, où le procédé comprend :
 - (i) la mise en contact de la solution comprenant de l'amidon avec une quantité efficace du variant d'amylase selon l'une quelconque des revendications 1 à 11 ; et
 - (ii) la saccharification de la solution comprenant de l'amidon afin de produire la composition comprenant du glucose ; où le variant d'amylase catalyse la saccharification de la solution d'amidon en glucose,
 facultativement où la composition comprenant de l'amidon comprend de l'amidon liquéfié, de l'amidon gélatinisé, ou de l'amidon granulaire.
14. Procédé selon la revendication 13, dans lequel la saccharification est réalisée dans une plage de températures d'environ 30 °C à environ 75 °C, facultativement dans lequel la plage de températures est de 47 °C-74 °C ; et/ou dans lequel la saccharification est réalisée sur une plage de pH d'un pH 2,0-7,5, facultativement dans lequel la plage de pH est un pH 3,5-5,5, facultativement en outre dans lequel la plage de pH est un pH 3,5-4,5.
15. Procédé selon la revendication 13 ou 14, comprenant en outre la fermentation de la composition de glucose afin de produire un produit de fin de fermentation (EOF).

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