



## Production of steviol glycosides in recombinant hosts

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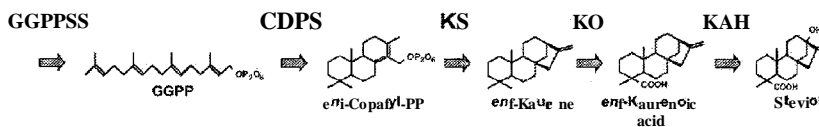
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(54) Title: PRODUCTION OF STEVIOL GLYCOSIDES IN RECOMBINANT HOSTS

Figure 1



(57) Abstract: The invention relates to recombinant microorganisms and methods for producing steviol glycosides and steviol glycoside precursors.



## PRODUCTION OF STEVIOL GLYCOSIDES IN RECOMBINANT HOSTS

### BACKGROUND OF THE INVENTION

#### Field of the Invention

**[0001]** This disclosure relates to recombinant production of steviol glycosides and steviol glycoside precursors in recombinant hosts. In particular, this disclosure relates to production of steviol glycosides comprising steviol-1 3-O-glucoside (13-SMG), steviol-1,2-bioside, steviol-1,3-bioside, steviol-1 9-O-glucoside (19-SMG), stevioside, 1,3-stevioside, rubusoside, Rebaudioside A (RebA), Rebaudioside B (RebB), Rebaudioside C (RebC), Rebaudioside D (RebD), Rebaudioside E (RebE), Rebaudioside F (RebF), Rebaudioside M (RebM), Rebaudioside Q (RebQ), Rebaudioside I (RebI), dulcoside A, or isomers thereof in recombinant hosts.

#### Description of Related Art

**[0001]** Sweeteners are well known as ingredients used most commonly in the food, beverage, or confectionary industries. The sweetener can either be incorporated into a final food product during production or for stand-alone use, when appropriately diluted, as a tabletop sweetener or an at-home replacement for sugars in baking. Sweeteners include natural sweeteners such as sucrose, high fructose corn syrup, molasses, maple syrup, and honey and artificial sweeteners such as aspartame, saccharine, and sucralose. Stevia extract is a natural sweetener that can be isolated and extracted from a perennial shrub, *Stevia rebaudiana*. Stevia is commonly grown in South America and Asia for commercial production of stevia extract. Stevia extract, purified to various degrees, is used commercially as a high intensity sweetener in foods and in blends or alone as a tabletop sweetener.

**[0002]** Chemical structures for several steviol glycosides are shown in Figure 1, including the diterpene steviol and various steviol glycosides. Extracts of the Stevia plant generally comprise steviol glycosides that contribute to the sweet flavor, although the amount of each steviol glycoside often varies, *inter alia*, among different production batches.

**[0002]** As recovery and purification of steviol glycosides from the Stevia plant have proven to be labor intensive and inefficient, there remains a need for a recombinant production system that can accumulate high yields of desired steviol glycosides, such as RebD and RebM. There

also remains a need for improved production of steviol glycosides in recombinant hosts for commercial uses.

### SUMMARY OF THE INVENTION

**[0003]** it is against the above background that the present invention provides certain advantages and advancements over the prior art.

**[0004]** Although this invention disclosed herein is not limited to specific advantages or functionalities, the invention provides a recombinant host comprising one or more of:

- (a) a gene encoding an ent-kaurene oxidase (KO) polypeptide;
- (b) a gene encoding a cytochrome P450 reductase (CPR) polypeptide; and/or
- (c) a gene encoding an ent-kaurenoic acid hydroxylase (KAH) polypeptide;

wherein at least one of the genes is a recombinant gene; and

wherein the recombinant host is capable of producing a steviol glycoside precursor.

**[0005]** The invention also provides a recombinant host comprising:

- (a) a gene encoding a geranylgeranyl diphosphate synthase (GGPPS) polypeptide;
- (b) a gene encoding an ent-copalyl diphosphate synthase (CDPS) polypeptide;
- (c) a gene encoding an ent-kaurene synthase (KS) polypeptide
- (d) a gene encoding an ent-kaurene oxidase (KO) polypeptide;
- (e) a gene encoding a cytochrome P450 reductase (CPR) polypeptide; and
- (f) a gene encoding an ent-kaurenoic acid hydroxylase (KAH) polypeptide;

wherein at least one of the genes is a recombinant gene; and

wherein the recombinant host is capable of producing stevioi.

**[0006]** In one aspect of the recombinant hosts disclosed herein,

- (a) the KO polypeptide comprises a KO polypeptide having at least 60% identity to an amino acid sequence set forth in SEQ ID NO:72 or SEQ ID NO:75; 65% identity to an amino acid sequence set forth in SEQ ID NO:54; at least 70% identity to an amino acid sequence set forth in SEQ ID NO: 70, SEQ ID NO:71 , or SEQ ID NO:79; at least 40% identity to an amino acid sequence set forth in SEQ

ID NO:77; or at least 50% identity to an amino acid sequence set forth in SEQ ID NO:78;

- (b) the CPR polypeptide comprises a CPR polypeptide having at least 70% identity to an amino acid sequences set forth in SEQ ID NO:69, SEQ ID NO:74, SEQ ID NO:76, or SEQ ID NO:87; at least 80% identity to an amino acid sequence set forth in SEQ ID NO:73; at least 85% identity to an amino acid sequence set forth in SEQ ID NO:22; at least 65% identity to an amino acid sequence set forth in SEQ ID NO:28; or at least 50% identity to an amino acid sequence set forth in SEQ ID NO:98; and/or
- (c) the KAH polypeptide comprises a KAH polypeptide having at least 40% identity to an amino acid sequence set forth in SEQ ID NO:82; at least 50% identity to an amino acid sequence set forth in SEQ ID NO:91; or at least 60% identity to an amino acid sequence set forth in SEQ ID NO:68.

**[0007]** The invention further provides a recombinant host comprising one or more of:

- (a) a gene encoding a KO polypeptide having at least 60% identity to an amino acid sequence set forth in SEQ ID NO:75;
- (b) a gene encoding a KAH polypeptide having at least 40% identity to an amino acid sequence set forth in SEQ ID NO:82; and/or
- (c) a gene encoding a CPR polypeptide having at least 50% identity to an amino acid sequence set forth in SEQ ID NO:98;

wherein at least one of the genes is a recombinant gene; and

wherein the recombinant host is capable of producing a steviol glycoside precursor.

**[0008]** The invention further provides a recombinant host comprising one or more of:

- (a) a gene encoding a KO polypeptide having at least 70% identity to an amino acid sequence set forth in SEQ ID NO:70;
- (b) a gene encoding a KAH polypeptide having at least 40% identity to an amino acid sequence set forth in SEQ ID NO:82; and/or
- (c) a gene encoding a CPR polypeptide having at least 50% identity to an amino acid sequence set forth in SEQ ID NO:98;

wherein at least one of the genes is a recombinant gene; and

wherein the recombinant host is capable of producing a steviol glycoside precursor.

**[0009]** In one aspect of the recombinant hosts disclosed herein, the host further comprises a gene encoding a KO polypeptide having at least 65% identity to an amino acid sequence set forth in SEQ ID NO:54.

**[0010]** In another aspect of the recombinant hosts disclosed herein, the recombinant host further comprises a gene encoding a KAH polypeptide having at least 60% identity to an amino acid sequence set forth in SEQ ID NO:68.

**[0011]** In another aspect of the recombinant hosts disclosed herein, the recombinant host further comprises a gene encoding a KO polypeptide having at least 70% identity to an amino acid sequence set forth in SEQ ID NO:79,

**[0012]** In one aspect of the recombinant hosts disclosed herein, the host further comprises one or more of:

- (a) a gene encoding a geranylgeranyl diphosphate synthase (GGPPS) polypeptide;
  - (b) a gene encoding an ent-copalyl diphosphate synthase (CDPS) polypeptide; and/or
  - (c) a gene encoding an ent-kaurene synthase (KS) polypeptide;
- wherein at least one of the genes is a recombinant gene; and

wherein the recombinant host is capable of producing a steviol glycoside precursor.

**[0013]** In some aspects of the recombinant hosts disclosed herein,

- (a) the GGPPS polypeptide comprises a polypeptide having at least 70% identity to an amino acid sequence set forth in SEQ ID NO:49;
- (b) the CDPS polypeptide comprises a polypeptide having at least 70% identity to an amino acid sequence set forth in SEQ ID NO:37; and/or
- (c) the KS polypeptide comprises a polypeptide having at least 40% identity to an amino acid sequence set forth in SEQ ID NO:6.

**[0014]** In one aspect of the recombinant hosts disclosed herein, the recombinant host further comprises a gene encoding an endoplasmic reticulum membrane polypeptide.

**[0015]** In another aspect of the recombinant hosts disclosed herein, the endoplasmic reticulum membrane polypeptide comprises an Inheritance of cortical ER protein 2 (ICE2)

polypeptide having at least 50% identity to the amino acid sequence set forth in SEQ ID NO:1 14.

**[0016]** In one aspect of the recombinant host disclosed herein, the KO polypeptide is a fusion construct.

**[0017]** In another aspect, the fusion construct comprises a polypeptide having at least 60% identity to an amino acid sequence set forth in SEQ ID NO:118 or SEQ ID NO:120.

**[0018]** In another aspect, the fusion construct has at least 50% identity to an amino acid sequence set forth in SEQ ID NO:100, SEQ ID NO:102, SEQ ID NO:104, SEQ ID NO:106, SEQ ID NO:108, SEQ ID NO:1 10, or SEQ ID NO:1 12.

**[0019]** in one aspect of the recombinant hosts disclosed herein, the host further comprises one or more of:

- (a) a gene encoding a UGT85C polypeptide;
- (b) a gene encoding a UGT76G polypeptide;
- (c) a gene encoding a UGT74G1 polypeptide;
- (d) a gene encoding a UGT91 D2 functional homolog polypeptide; and/or
- (e) a gene encoding an EUGT11 polypeptide;

wherein at least one of the genes is a recombinant gene; and

wherein the host is capable of producing a steviol glycoside.

**[0020]** In some aspects of the recombinant hosts disclosed herein,

- (a) the UGT85C2 polypeptide comprises a polypeptide having at least 55% identity to an amino acid sequence set forth in SEQ ID NO:30;
- (b) the UGT76G1 polypeptide comprises a polypeptide having at least 50% identity to an amino acid sequence set forth in SEQ ID NO:83;
- (c) the UGT74G1 polypeptide comprises a polypeptide having at least 55% identity to an amino acid sequence set forth in SEQ ID NO:29;
- (d) the UGT91D2 functional homolog polypeptide comprises a UGT91D2 polypeptide having 90% or greater identity to the amino acid sequence set forth in SEQ ID NO:84 or a UGT91D2e-b polypeptide having 90% or greater identity to the amino acid sequence set forth in SEQ ID NO:88; and/or

- (e) the EUGT1 1 polypeptide comprises a polypeptide having at least 65% identity to an amino acid sequence set forth in SEQ ID NO:86.

[0021] In some aspects, the recombinant hosts disclosed herein comprise a plant cell, a mammalian cell, an insect cell, a fungal cell, or a bacterial cell.

[0022] In one aspect, the bacterial cell comprises *Escherichia* bacteria cells, for example, *Escherichia coli* cells; *Lactobacillus* bacteria cells; *Lactococcus* bacteria cells; *Cornibacterium* bacteria cells; *Acetobacter* bacteria cells; *Acinetobacter* bacteria cells; or *Pseudomonas* bacterial cells.

[0023] In one aspect, the fungal cell comprises a yeast cell.

[0024] In one aspect, the yeast cell is a cell from *Saccharomyces cerevisiae*, *Schizosaccharomyces pombe*, *Yarrowia lipolytica*, *Candida glabrata*, *Ashbya gossypii*, *Cyberlindnera jadinii*, *Pichia pastoris*, *Kluyveromyces lactis*, *Hansenula polymorpha*, *Candida boidinii*, *Arxula adenivorans*, *Xanthophyllomyces dendrorhous*, or *Candida albicans* species.

[0025] In one aspect, the yeast cell is a *Saccharomycete*.

[0026] In one aspect, the yeast cell is a cell from the *Saccharomyces cerevisiae* species.

[0027] The invention further provides a method of producing a steviol glycoside or a steviol glycoside precursor, comprising:

- (a) growing a recombinant host disclosed herein in a culture medium, under conditions in which any of the genes disclosed herein are expressed;

wherein the steviol glycoside or the steviol glycoside precursor is synthesized by said host; and/or

- (b) optionally quantifying the steviol glycoside or the steviol glycoside precursor; and/or

- (c) optionally isolating the steviol glycoside or the steviol glycoside precursor.

[0028] In some aspects, the steviol glycoside comprises steviol-1,3-O-glucoside (13-SMG), steviol-1,2-bioside, steviol-1,3-bioside, steviol-1,9-O-glucoside (19-SMG), stevioside, 1,3-stevioside, rubusoside, Rebaudioside A (RebA), Rebaudioside B (RebB), Rebaudioside C (RebC), Rebaudioside D (RebD), Rebaudioside E (RebE), Rebaudioside F (RebF), Rebaudioside M (RebM), Rebaudioside Q (RebQ), Rebaudioside I (Rebi), dulcoside A, di-



glycosylated steviol, tri-glycosylated steviol, tetra-glycosylated steviol, penta-glycosylated steviol, hexa-glycosylated steviol, hepta-glycosylated steviol, or isomers thereof.

**[0029]** In some aspects, the steviol glycoside or steviol glycoside precursor produced by the recombinant hosts or methods disclosed herein accumulates to a detectable concentration when cultured under said conditions.

[0030] In some aspects, the steviol glycoside or steviol glycoside precursor produced by the recombinant hosts or methods disclosed herein has an undetectable concentration of stevia plant-derived contaminants.

**[0031]** In some aspects, the steviol glycoside or steviol glycoside precursor produced by the recombinant hosts or methods disclosed herein has a steviol glycoside composition enriched for RebD or RebM relative to the steviol glycoside composition of a wild-type *Stevia* plant.

**[0032]** These and other features and advantages of the present invention will be more fully understood from the following detailed description taken together with the accompanying claims. It is noted that the scope of the claims is defined by the recitations therein and not by the specific discussion of features and advantages set forth in the present description.

#### BRIEF DESCRIPTION OF THE DRAWINGS

**[0033]** The following detailed description of the embodiments of the present invention can be best understood when read in conjunction with the following drawings, where like structure is indicated with like reference numerals and in which:

**[0034]** Figure 1 shows a schematic of the engineered biosynthetic pathway for producing steviol in yeast from geranylgeranyl diphosphate using geranylgeranyl diphosphate synthase (GGPPS), ent-copalyl diphosphate synthase (CDPS), ent-kaurene synthase (KS), ent-kaurene oxidase (KO), and ent-kaurenoic acid hydroxylase (KAH) polypeptides.

**[0035]** Figure 2 shows representative steviol glycoside glycosylation reactions catalyzed by suitable uridine S'-diphospho (UDP) glycosyl transferases (UGT) enzymes and chemical structures for several steviol glycoside compounds.

**[0036]** Figure 3 shows Rebaudioside B (RebB) production in a steviol glycoside-producing *S. cerevisiae* strain individually expressing *S. rebaudiana* K01 (SrK01) encoded by the nucleotide sequence set forth in SEQ ID NO:59, the KO encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:55, or the KO encoded by the nucleotide sequence

set forth in SEQ ID NO:56. RebB production was measured by liquid chromatography-mass spectrometry (LC-MS) analysis as  $\mu\text{M}/\text{OD}_{600}$  of individual cultures. See Example 3.

[0037] Figure 4 shows production of ent-kaurenoic acid in steviol glycoside-producing *S. cerevisiae* strains individually expressing SrKOl encoded by the nucleotide sequence set forth in SEQ ID NO:59, the KO encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:55, or the KO encoded by the nucleotide sequence set forth in SEQ ID NO:56, as measured by LC-MS analysis of culture samples. Ent-kaurenoic acid levels were calculated as the Area under Curve (AUC) of LC-MS peaks corresponding to ent-kaurenoic acid. See Example 3.

[0038] Figure 5 shows production of total (extracellular plus intracellular) steviol glycosides in a steviol glycoside-producing *S. cerevisiae* strain overexpressing *S. rebaudiana* KAHel (SrKAHel; encoded by the nucleotide sequence set forth in SEQ ID NO:18) or in a steviol glycoside-producing *S. cerevisiae* strain co-expressing SrKAHel (encoded by the nucleotide sequence set forth in SEQ ID NO:18) and a KO encoded by the nucleotide sequences set forth in any one of SEQ ID NOs: 55-60, compared to a control strain that does not overexpress SrKAHel or express a KO encoded by the nucleotide sequence set forth in any one of SEQ ID NOs: 55-60. Production of total steviol glycosides was quantified by comparison to a standard curve. Values plotted on the y-axis in  $\mu\text{M}$  are an average of three biological replicates. See Example 4.

[0039] Figure 6 shows production of Rebaudioside A (RebA), Rebaudioside D (RebD), and Rebaudioside M (RebM) in a steviol glycoside-producing *S. cerevisiae* strain overexpressing SrKAHel (encoded by the nucleotide sequence set forth in SEQ ID NO:18) and further expressing either the KO encoded by the nucleotide sequence set forth in SEQ ID NO:56 or the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65. Production of RebA + RebD + RebM was measured in  $\mu\text{M}$ . See Example 4.

[0040] Figure 7 shows production of glycosylated ent-kaurenoic acid in a steviol glycoside-producing *S. cerevisiae* strain overexpressing SrKAHel (encoded by the nucleotide sequence set forth in SEQ ID NO:18) or in a steviol glycoside-producing strain coexpressing SrKAHel (encoded by the nucleotide sequence set forth in SEQ ID NO:18) and a KO encoded by the nucleotide sequences set forth in any one of SEQ ID NOs: 55-60). Values were calculated as the AUC of LC-MS peaks corresponding to glycosylated ent-kaurenoic acid and as an average of three biological replicates. See Example 4.

**[0041]** Figure 8 shows production of glycosylated ent-kaurenol in a steviol glycoside-producing *S. cerevisiae* strain overexpressing SrKAHel (encoded by the nucleotide sequence set forth in SEQ ID NO:18) or in a steviol glycoside-producing *S. cerevisiae* strain co-expressing SrKAHel (encoded by the nucleotide sequence set forth in SEQ ID NO:18) and a KO encoded by the nucleotide sequence set forth in SEQ ID NOs: 55-60). Values plotted on the y-axis were calculated as the AUC of LC-MS peaks corresponding to glycosylated ent-kaurenol. See Example 4.

**[0042]** Figure 9 shows Rebaudioside <sub>IVf</sub> (RebM) production in a steviol glycoside-producing *S. cerevisiae* strain expressing CPR1 (encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:61) or CPR7 (encoded by the nucleotide sequence set forth in SEQ ID NO:23). Values plotted on the y-axis were measured in  $\mu\text{M}$ . See Example 5.

**[0043]** Figure 10 shows Rebaudioside M (RebM) production in a steviol glycoside-producing *S. cerevisiae* strain overexpressing SrKAHel (encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:18) and further expressing CPR4497 encoded by the nucleotide sequence set forth in SEQ ID NO:62. Values plotted on the y-axis indicate  $\mu\text{M}$  concentration of RebM. See Example 5.

**[0044]** Figure 11A shows an LC-MS chromatogram of a steviol-1 3-O-glucoside (13-SMG) standard. Figure 11B shows production of 13-SMG by a steviol glycoside-producing *S. cerevisiae* strain expressing the KAH encoded by the nucleotide sequence set forth in SEQ ID NO:80 (amino acid sequence set forth in SEQ ID NO:82). See Example 7.

**[0045]** Figure 12 shows steviol-1 3-O-glucoside (13-SMG) and Rebaudioside B (RebB) production in a steviol glycoside-producing *S. cerevisiae* strain co-expressing a KO and a CPR. The KO was selected from SrKOl (encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:59), the KO encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:63, or the KO encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:64. The cytochrome P450 reductase (CPR) polypeptide was selected from the CPR encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:66 or the CPR encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:67. Values displayed on the y-axis are  $\mu\text{M}$  concentrations of the indicated steviol glycosides. See Example 6.

**[0046]** Figure 13 shows production of steviol-1 3-O-glucoside (13-SMG) and rubusoside in a steviol glycoside-producing *S. cerevisiae* strain expressing SrKAHel (encoded by the

nucleotide sequence set forth in SEQ ID NO: 18), the KAH encoded by the nucleotide sequence set forth in SEQ ID NO:80, or the KAH encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:81. Values displayed in the y-axis are  $\mu\text{M}$  concentrations of 13-SMG and rubusoside, averaged over eight biological replicates and normalized to  $\text{OD}_{600}$  measured using a plate reader. Error bars are  $\pm$  the respective standard deviation. See Example 7.

**[0047]** Figure 14 shows cytochrome P450 reductase (CPR) polypeptide activity on cytochrome c upon incubation with microsomal protein prepared from *S. cerevisiae* strains expressing SrKAHel (encoded by the nucleotide sequence set forth in SEQ ID NO:18) alone or in combination with CPR1 (encoded by the nucleotide sequence set forth in SEQ ID NO:61) or CPR12 (encoded by the nucleotide sequence set forth in SEQ ID NO:97). Results are shown in U/mg as an average of two biological replicates. See Example 9.

**[0048]** Figure 15A shows steviol accumulation upon 30 min incubation of ent-kaurenoic acid with microsomal protein prepared from *S. cerevisiae* strains expressing SrKAHel (encoded by the nucleotide sequence set forth in SEQ ID NO:18) alone or in combination with CPR1 (encoded by the nucleotide sequence set forth in SEQ ID NO:61) or CPR12 (encoded by the nucleotide sequence set forth in SEQ ID NO:97). Results are shown in AUC as an average of three biological replicates. Control reactions comprised the microsomal protein described above, but these were not incubated for 30 min prior to measurement of steviol accumulation. Figure 15B shows levels of ent-kaurenoic acid following 30 min incubation of ent-kaurenoic acid with microsomal protein prepared from *S. cerevisiae* strains expressing SrKAHel (encoded by the nucleotide sequence set forth in SEQ ID NO:18) alone or in combination with CPR1 (encoded by the nucleotide sequence set forth in SEQ ID NO:61) or CPR12 (encoded by the nucleotide sequence set forth in SEQ ID NO:97). Results are shown in  $\mu\text{M}$  as an average of three biological replicates. Control reactions comprised the microsomal protein described above but were not incubated for 30 min prior to measurement of ent-kaurenoic acid levels. See Example 9.

**[0049]** Figure 16 shows steviol-13-O-glucoside (13-SMG), 1,2-bioside, Rebaudioside B (RebB), ent-kaurenoic acid, and ent-kaurene levels accumulated by a steviol glycoside-producing *S. cerevisiae* strain expressing SrKOi (SEQ ID NO:59, SEQ ID NO:79), a KO encoded by the nucleotide sequence set forth in SEQ ID NO:65, or a fusion construct between either SrKOi or the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 and the NADPH-dependent P450 oxidoreductase domain of CYP102A1 (referred to herein as the "BMR domain"). Figure 16A shows levels of 13-SMG, 1,2-bioside, and RebB measured by LC-MS for

a steviol glycoside-producing *S. cerevisiae* strain expressing SrKOI (SEQ ID NO:59, SEQ ID NO:79), a fusion construct of SrKOI and BMR (SEQ ID NO:99, SEQ ID NO:100), a fusion construct of SrKOI and BMR W1046A (SEQ ID NO:101, SEQ ID NO:102), a fusion construct of truncated SrKOI and BMR (SEQ ID NO:103, SEQ ID NO:104), a fusion construct of truncated SrKOI and BMR W1046A (SEQ ID NO:105, SEQ ID NO:106), or a control plasmid. Figure 16B shows levels of ent-kaurenoic acid and ent-kaurene measured by LC-UV for a steviol glycoside-producing *S. cerevisiae* strain expressing SrKOI (SEQ ID NO:59, SEQ ID NO:79), a fusion construct of SrKOI and BMR (SEQ ID NO:99, SEQ ID NO:100), a fusion construct of SrKOI and BMR W1046A (SEQ ID NO:101, SEQ ID NO:102), a fusion construct of truncated SrKOI and BMR (SEQ ID NO:103, SEQ ID NO:104), a fusion construct of truncated SrKOI and BMR W1046A (SEQ ID NO:105, SEQ ID NO:106), or a control plasmid. Figure 16C shows levels of 13-SMG, 1,2-bioside, and RebB measured by LC-MS for a steviol glycoside-producing *S. cerevisiae* strain expressing the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65, a fusion construct of the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 and BMR (SEQ ID NO:107, SEQ ID NO:108), a fusion construct of the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 and BMR W1046A (SEQ ID NO:109, SEQ ID NO:110), a fusion construct of a truncated KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 and BMR W1046A (SEQ ID NO:111, SEQ ID NO:112), or a plasmid control. Figure 16D shows levels of ent-kaurenoic acid or ent-kaurene accumulated by a steviol glycoside-producing *S. cerevisiae* strain expressing the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65, a fusion construct of the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 and BMR (SEQ ID NO:107, SEQ ID NO:108), a fusion construct of the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 and BMR W1046A (SEQ ID NO:109, SEQ ID NO:110), a fusion construct of a truncated KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 and BMR W1046A (SEQ ID NO:111, SEQ ID NO:112), or a plasmid control. See Example 10.

## DETAILED DESCRIPTION OF THE INVENTION

**[0050]** Before describing the present invention in detail, a number of terms will be defined. As used herein, the singular forms "a," "an," and "the" include plural referents unless the context clearly dictates otherwise. For example, reference to a "nucleic acid" means one or more nucleic acids.

[0051] It is noted that terms like "preferably," "commonly," and "typically" are not utilized herein to limit the scope of the claimed invention or to imply that certain features are critical, essential, or even important to the structure or function of the claimed invention. Rather, these terms are merely intended to highlight alternative or additional features that can or cannot be utilized in a particular embodiment of the present invention.

[0052] For the purposes of describing and defining the present invention it is noted that the term "substantially" is utilized herein to represent the inherent degree of uncertainty that can be attributed to any quantitative comparison, value, measurement, or other representation. The term "substantially" is also utilized herein to represent the degree by which a quantitative representation can vary from a stated reference without resulting in a change in the basic function of the subject matter at issue.

[0053] Methods well known to those skilled in the art can be used to construct genetic expression constructs and recombinant cells according to this invention. These methods include *in vitro* recombinant DNA techniques, synthetic techniques, *in vivo* recombination techniques, and polymerase chain reaction (PCR) techniques. See, for example, techniques as described in Green & Sambrook, 2012, MOLECULAR CLONING: A LABORATORY MANUAL, Fourth Edition, Cold Spring Harbor Laboratory, New York; Ausubei *et al.*, 1989, CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, Greene Publishing Associates and Wiley Interscience, New York, and PCR Protocols: A Guide to Methods and Applications (Innis *et al.*, 1990, Academic Press, San Diego, CA).

[0054] As used herein, the terms "polynucleotide", "nucleotide", "oligonucleotide", and "nucleic acid" can be used interchangeably to refer to nucleic acid comprising DNA, RNA, derivatives thereof, or combinations thereof.

[0055] As used herein, the terms "microorganism," "microorganism host," "microorganism host cell," "recombinant host," and "recombinant host cell" can be used interchangeably. As used herein, the term "recombinant host" is intended to refer to a host, the genome of which has been augmented by at least one DNA sequence. Such DNA sequences include but are not limited to genes that are not naturally present, DNA sequences that are not normally transcribed into RNA or translated into a protein ("expressed"), and other genes or DNA sequences which one desires to introduce into a host. It will be appreciated that typically the genome of a recombinant host described herein is augmented through stable introduction of one or more recombinant genes. Generally, introduced DNA is not originally resident in the host that is the recipient of the DNA, but it is within the scope of this disclosure to isolate a DNA segment from

a given host, and to subsequently introduce one or more additional copies of that DNA into the same host, e.g., to enhance production of the product of a gene or alter the expression pattern of a gene. In some instances, the introduced DNA will modify or even replace an endogenous gene or DNA sequence by, e.g., homologous recombination or site-directed mutagenesis. Suitable recombinant hosts include microorganisms.

[0056] As used herein, the term "recombinant gene" refers to a gene or DNA sequence that is introduced into a recipient host, regardless of whether the same or a similar gene or DNA sequence may already be present in such a host. "Introduced," or "augmented" in this context, is known in the art to mean introduced or augmented by the hand of man. Thus, a recombinant gene can be a DNA sequence from another species or can be a DNA sequence that originated from or is present in the same species but has been incorporated into a host by recombinant methods to form a recombinant host. It will be appreciated that a recombinant gene that is introduced into a host can be identical to a DNA sequence that is normally present in the host being transformed, and is introduced to provide one or more additional copies of the DNA to thereby permit overexpression or modified expression of the gene product of that DNA. In some aspects, said recombinant genes are encoded by cDNA. In other embodiments, recombinant genes are synthetic and/or codon-optimized for expression in *S. cerevisiae*.

[0057] As used herein, the term "engineered biosynthetic pathway" refers to a biosynthetic pathway that occurs in a recombinant host, as described herein. In some aspects, one or more steps of the biosynthetic pathway do not naturally occur in an unmodified host. In some embodiments, a heterologous version of a gene is introduced into a host that comprises an endogenous version of the gene.

[0058] As used herein, the term "endogenous" gene refers to a gene that originates from and is produced or synthesized within a particular organism, tissue, or cell. In some embodiments, the endogenous gene is a yeast gene. In some embodiments, the gene is endogenous to *S. cerevisiae*, including, but not limited to *S. cerevisiae* strain S288C. In some embodiments, an endogenous yeast gene is overexpressed. As used herein, the term "overexpress" is used to refer to the expression of a gene in an organism at levels higher than the level of gene expression in a wild type organism. See, e.g., Prelich, 2012, *Genetics* 190:841-54. In some embodiments, an endogenous yeast gene is deleted. See, e.g., Giaever & Nislow, 2014, *Genetics* 197(2):451-65. As used herein, the terms "deletion," "deleted," "knockout," and "knocked out" can be used interchangeably to refer to an endogenous gene that

has been manipulated to no longer be expressed in an organism, including, but not limited to, *S. cerevisiae*.

[0059] As used herein, the terms "heterologous sequence" and "heterologous coding sequence" are used to describe a sequence derived from a species other than the recombinant host, in some embodiments, the recombinant host is an *S. cerevisiae* cell, and a heterologous sequence is derived from an organism other than *S. cerevisiae*. A heterologous coding sequence, for example, can be from a prokaryotic microorganism, a eukaryotic microorganism, a plant, an animal, an insect, or a fungus different than the recombinant host expressing the heterologous sequence. In some embodiments, a coding sequence is a sequence that is native to the host.

[0060] A "selectable marker" can be one of any number of genes that complement host cell auxotrophy, provide antibiotic resistance, or result in a color change. Linearized DNA fragments of the gene replacement vector then are introduced into the cells using methods well known in the art (see below). Integration of the linear fragments into the genome and the disruption of the gene can be determined based on the selection marker and can be verified by, for example, PGR or Southern blot analysis. Subsequent to its use in selection, a selectable marker can be removed from the genome of the host cell by, e.g., Cre-LoxP systems (see, e.g., Gossen *et al.*, 2002, *Ann. Rev. Genetics* 36:153-173 and U.S. 2006/0014264). Alternatively, a gene replacement vector can be constructed in such a way as to include a portion of the gene to be disrupted, where the portion is devoid of any endogenous gene promoter sequence and encodes none, or an inactive fragment of, the coding sequence of the gene.

[0061] As used herein, the terms "variant" and "mutant" are used to describe a protein sequence that has been modified at one or more amino acids, compared to the wild-type sequence of a particular protein.

[0062] As used herein, the term "inactive fragment" is a fragment of the gene that encodes a protein having, e.g., less than about 10% (e.g., less than about 9%, less than about 8%, less than about 7%, less than about 6%, less than about 5%, less than about 4%, less than about 3%, less than about 2%, less than about 1%, or 0%) of the activity of the protein produced from the full-length coding sequence of the gene. Such a portion of a gene is inserted in a vector in such a way that no known promoter sequence is operably linked to the gene sequence, but that a stop codon and a transcription termination sequence are operably linked to the portion of the gene sequence. This vector can be subsequently linearized in the portion of the gene sequence



and transformed into a cell. By way of single homologous recombination, this linearized vector is then integrated in the endogenous counterpart of the gene with inactivation thereof.

**[0063]** As used herein, the term "steviol glycoside" refers to Rebaudioside A (RebA) (CAS # 58543-16-1), Rebaudioside B (RebB) (CAS # 58543-17-2), Rebaudioside C (RebC) (CAS # 63550-99-2), Rebaudioside D (RebD) (CAS # 63279-13-0), Rebaudioside E (RebE) (CAS # 63279-14-1), Rebaudioside F (RebF) (CAS # 438045-89-7), Rebaudioside M (RebM) (CAS # 1220616-44-3), Rubusoside (CAS # 63849-39-4), Dulcoside A (CAS # 64432-06-0), Rebaudioside I (RebI) (MassBank Record: FU000332), Rebaudioside Q (RebQ), 1,2-Stevioside (CAS # 57817-89-7), 1,3-Stevioside (RebG), 1,2-bioside (MassBank Record: FU000299), 1,3-bioside, Steviol-13-O-glucoside (13-SMG), Steviol-19-O-glucoside (19-SMG), a tri-glucosylated steviol glycoside, a tetra-glucosylated steviol glycoside, a penta-glucosylated steviol glycoside, a hexa-glucosylated steviol glycoside, a hepta-glucosylated steviol glycoside, and isomers thereof. See Figure 2; see also, Steviol Glycosides Chemical and Technical Assessment 69th JECFA, 2007, prepared by Harriet Wallin, Food Agric. Org.

**[0064]** As used herein, the terms "steviol glycoside precursor" and "steviol glycoside precursor compound" are used to refer to intermediate compounds in the steviol glycoside biosynthetic pathway. Steviol glycoside precursors include, but are not limited to, geranylgeranyl diphosphate (GGPP), ent-copalyl-diphosphate, ent-kaurene, ent-kaurenol, ent-kaurenai, ent-kaurenoic acid, and steviol. See Figure 1. In some embodiments, steviol glycoside precursors are themselves steviol glycoside compounds. For example, 19-SMG, rubusoside, stevioside, and RebE are steviol glycoside precursors of RebM. See Figure 2. Steviol glycosides and/or steviol glycoside precursors can be produced *in vivo* (*i.e.*, in a recombinant host), *in vitro* (*i.e.*, enzymatically), or by whole cell bioconversion. As used herein, the terms "produce" and "accumulate" can be used interchangeably to describe synthesis of steviol glycosides and steviol glycoside precursors *in vivo*, *in vitro*, or by whole cell bioconversion.

**[0065]** As used herein, the term "di-glycosylated steviol" can be used to refer to a steviol molecule comprising two sugar moieties, such as glucose or N-acetylglucosamine (GlcNAc). Non-limiting examples of di-glycosylated steviol molecules include steviol-1,3-bioside, steviol-1,2-bioside, rubusoside, a steviol molecule comprising two glucose moieties, a steviol molecule comprising one glucose moiety and one GlcNAc moiety, and isomers thereof.

**[0066]** As used herein, the term "tri-glycosylated steviol" can be used to refer to a steviol molecule comprising three sugar moieties, such as glucose or GlcNAc. Non-limiting examples

of tri-glycosylated steviol molecules include RebB, RebG, stevioside, a steviol molecule comprising two glucose moieties and one GlcNAc moiety, and isomers thereof.

[0067] As used herein, the term "tetra-glycosylated steviol" can be used to refer to a steviol molecule comprising four sugar moieties, such as glucose or GlcNAc. Non-limiting examples of tetra-glycosylated steviol molecules include RebA, RebE, RebQ, a steviol molecule comprising four glucose moieties, a steviol molecule comprising three glucose moieties and one GlcNAc moiety, and isomers thereof.

[0068] As used herein, the term "penta-glycosylated steviol" can be used to refer to a steviol molecule comprising five sugar moieties, such as glucose or GlcNAc. Non-limiting examples of penta-glycosylated steviol molecules include RebD, a steviol molecule comprising five glucose moieties, a steviol molecule comprising four glucose moieties and one GlcNAc moiety, and isomers thereof.

[0069] As used herein, the term "hexa-glycosylated steviol" can be used to refer to a steviol molecule comprising six sugar moieties, such as glucose or GlcNAc. Non-limiting examples of hexa-glycosylated steviol molecules include RebM, a steviol molecule comprising six glucose moieties, a steviol molecule comprising five glucose moieties and one GlcNAc moiety, and isomers thereof.

[0070] As used herein, the term "hepta-glycosylated steviol" can be used to refer to a steviol molecule comprising seven sugar moieties, such as glucose or GlcNAc. Non-limiting examples of hepta-glycosylated steviol molecules include a steviol molecule comprising seven glucose moieties and isomers thereof.

[0071] As used herein, the term "glycosylated ent-kaurenoic acid" can be used to refer to an ent-kaurenoic acid molecule comprising sugar moieties, such as glucose or GlcNAc. Non-limiting examples of glycosylated ent-kaurenoic acid molecules include ent-kaurenoic acid molecule comprising two glucose moieties and one GlcNAc moiety, an ent-kaurenoic acid molecule comprising three glucose moieties, an ent-kaurenoic acid molecule comprising one glucose moiety and one GlcNAc moiety, an ent-kaurenoic acid molecule comprising two glucose moieties, and isomers thereof.

[0072] As used herein, the term "glycosylated ent-kaurenol" can be used to refer to an ent-kaurenol molecule comprising sugar moieties, such as glucose or GlcNAc. Non-limiting examples of glycosylated ent-kaurenol molecules include an ent-kaurenol molecule comprising three glucose moieties, an ent-kaurenol molecule comprising one glucose moiety and one

GlcNAc moiety, an ent-kaureno! molecule comprising two glucose moieties, and isomers thereof.

[0073] Recombinant steviol glycoside-producing *Saccharomyces cerevisiae* (*S. cerevisiae*) strains are described in WO 2011/153378, WO 2013/022989, WO 2014/122227, and WO 2014/122328. Methods of producing steviol glycosides in recombinant hosts, by whole cell bio-conversion, and *in vitro* are also described in WO 2011/153378, WO 2013/022989, WO 2014/122227, and WO 2014/122328.

[0074] In some embodiments, steviol glycosides and/or steviol glycoside precursors are produced *in vivo* through expression of one or more enzymes involved in the steviol glycoside biosynthetic pathway in a recombinant host. For example, a steviol-producing recombinant host expressing one or more of a gene encoding a GGPPS polypeptide, a gene encoding a CDPS polypeptide, a gene encoding a KS polypeptide, a gene encoding a KO polypeptide, a gene encoding a KAH polypeptide, a gene encoding a CPR polypeptide, and a gene encoding a UGT polypeptide can produce a steviol glycoside and/or steviol glycoside precursors *in vivo*. See, e.g., Figures 1 and 2. The skilled worker will appreciate that one or more of these genes can be endogenous to the host provided that at least one (and in some embodiments, all) of these genes is a recombinant gene introduced into the recombinant host.

[0075] In another example, a recombinant host expressing a gene encoding a GGPPS polypeptide, a gene encoding a CDPS polypeptide, a gene encoding a KS polypeptide, a gene encoding a KO polypeptide, a gene encoding a KAH polypeptide, and a gene encoding a CPR polypeptide can produce steviol *in vivo*. See, e.g., Figures 1. The skilled worker will appreciate that one or more of these genes can be endogenous to the host provided that at least one (and in some embodiments, all) of these genes is a recombinant gene introduced into the recombinant host.

[0076] In another example, a steviol-producing recombinant host expressing a gene encoding a GGPPS polypeptide, a gene encoding a CDPS polypeptide, a gene encoding a KS polypeptide, a gene encoding a KO polypeptide, a gene encoding a KAH polypeptide, a gene encoding a CPR polypeptide, and one or more of a gene encoding a UGT polypeptide can produce a steviol glycoside *in vivo*. See, e.g., Figures 1 and 2. The skilled worker will appreciate that one or more of these genes can be endogenous to the host provided that at least one (and in some embodiments, all) of these genes is a recombinant gene introduced into the recombinant host.

**[0077]** Non-limiting examples of KS polypeptides are set forth in SEQ ID NOs:1-4 and SEQ ID NO:6. Non-limiting examples of KO polypeptides are set forth in SEQ ID NOs:7-10, 54, 70-72, 75, and 77-79. Non-limiting examples of KAH polypeptides are set forth in SEQ ID NOs:13-17, 68, 82, and 91. Non-limiting examples of CPR polypeptides are set forth in SEQ ID NOs:20-22, 28, 69, 73, 74, 76, 87, and 98. Non-limiting examples of CDPS polypeptides are set forth in SEQ ID NOs:33-39. Non-limiting examples of CDPS-KS polypeptides are set forth in SEQ ID NOs:40-42. Non-limiting examples of GGPPS polypeptides are set forth in SEQ ID NOs:43-50.

**[0078]** In some embodiments, a recombinant host comprises a nucleic acid encoding a UGT85C2 polypeptide (SEQ ID NO:32), a nucleic acid encoding a UGT76G1 polypeptide (SEQ ID NO:83), a nucleic acid encoding a UGT74G1 polypeptide (SEQ ID NO:29), a nucleic acid encoding a UGT91D2 polypeptide, and/or a nucleic acid encoding a EUGT1 1 polypeptide (SEQ ID NO:86). In some aspects, the UGT91D2 polypeptide can be a UGT91D2e polypeptide (SEQ ID NO:84) or a UGT91D2e-b polypeptide (SEQ ID NO:88). The skilled worker will appreciate that expression of these genes may be necessary to produce a particular steviol glycoside but that one or more of these genes can be endogenous to the host provided that at least one (and in some embodiments, all) of these genes is a recombinant gene introduced into the recombinant host. In a particular embodiment, a steviol-producing recombinant microorganism comprises exogenous nucleic acids encoding UGT85C2, UGT76G1, or UGT91D2 polypeptides. In another particular embodiment, a steviol-producing recombinant microorganism comprises exogenous nucleic acids encoding UGT85C2, UGT76G1, UGT74G1, and UGT91D2 polypeptides. In yet another particular embodiment, a steviol-producing recombinant microorganism comprises exogenous nucleic acids encoding UGT85C2, UGT76G1, UGT74G1, and EUGT11 polypeptides. In yet another particular embodiment, a steviol-producing recombinant microorganism comprises the exogenous nucleic acids encoding UGT85C2, UGT76G1, UGT74G1, UGT91D2 (including *inter alia* 91D2e, 91D2m, 91D2e-b, and functional homologs thereof), and EUGT1 1 polypeptides.

**[0079]** In certain embodiments, the steviol glycoside is RebA, RebB, RebD, and/or RebM. RebA can be synthesized in a steviol-producing recombinant microorganism expressing UGT85C2, UGT76G1, UGT74G1, and UGT91D2. RebB can be synthesized in a steviol-producing recombinant microorganism expressing UGT85C2, UGT76G1, and UGT91D2. RebD can be synthesized in a steviol-producing recombinant microorganism expressing UGT85C2, UGT76G1, UGT74G1, and UGT91D2 and/or EUGT1 1. RebM can be synthesized in a steviol-

producing recombinant microorganism expressing UGT85C2, UGT76G1, UGT74G1, and UGT91 D2 and/or EUGT1 1 (see Figure 2).

[0080] In some embodiments, steviol glycosides and/or steviol glycoside precursors are produced through contact of a steviol glycoside precursor with one or more enzymes involved in the steviol glycoside pathway *in vitro*. For example, contacting steviol with a UGT polypeptide can result in production of a steviol glycoside *in vitro*. In some embodiments, a steviol glycoside precursor is produced through contact of an upstream steviol glycoside precursor with one or more enzymes involved in the steviol glycoside pathway *in vitro*. For example, contacting ent-kaurenoic acid with a KAH enzyme can result in production of steviol *in vitro*.

[0081] In some embodiments, a steviol glycoside or steviol glycoside precursor is produced by whole cell bioconversion. For whole cell bioconversion to occur, a host cell expressing one or more enzymes involved in the steviol glycoside pathway takes up and modifies a steviol glycoside precursor in the cell; following modification *in vivo*, a steviol glycoside remains in the cell and/or is excreted into the culture medium. For example, a host cell expressing a gene encoding a UGT polypeptide can take up steviol and glycosylate steviol in the cell; following glycosylation *in vivo*, a steviol glycoside can be excreted into the culture medium. In some embodiments, the cell is permeabilized to take up a substrate to be modified or to excrete a modified product.

[0082] In some embodiments, steviol, one or more steviol glycoside precursors, and/or one or more steviol glycosides are produced by co-culturing of two or more hosts. In some embodiments, one or more hosts, each expressing one or more enzymes involved in the steviol glycoside pathway, produce steviol, one or more steviol glycoside precursors, and/or one or more steviol glycosides. For example, a host comprising a GGPPS, a CDPS, a KO, a KS, a KAH, and/or a CPR and a host comprising one or more UGTs produce one or more steviol glycosides.

[0083] In some embodiments, a steviol glycoside or steviol glycoside precursor composition produced *in vivo*, *in vitro*, or by whole cell bioconversion comprises less contaminants than a stevia extract from, *inter alia*, a stevia plant. Contaminants include plant-derived compounds that contribute to off-flavors. Potential contaminants include pigments, lipids, proteins, phenolics, saccharides, spathulenol and other sesquiterpenes, labdane diterpenes, monoterpenes, decanoic acid, 8,11,14-eicosatrienoic acid, 2-methyloctadecane, pentacosane, octacosane, tetracosane, octadecanol, stigmasterol,  $\beta$ -sitosterol,  $\alpha$ -amyrin,  $\beta$ -amyrin, lupeol,  $\beta$ -

amryin acetate, pentacyclic triterpenes, centauredin, quercitin, epi-alpha-cadinoi, carophyllenes and derivatives, beta-pinene, beta-sitosterol, and gibberellin.

**[0084]** As used herein, the terms "detectable amount," "detectable concentration," "measurable amount," and "measurable concentration" refer to a level of steviol glycosides measured in AUC,  $\mu\text{M}/\text{OD}_{600}$ , mg/L,  $\mu\text{M}$ , or mM. Steviol glycoside production (*i.e.*, total, supernatant, and/or intracellular steviol glycoside levels) can be detected and/or analyzed by techniques generally available to one skilled in the art, for example, but not limited to, liquid chromatography-mass spectrometry (LC-MS), thin layer chromatography (TLC), high-performance liquid chromatography (HPLC), ultraviolet-visible spectroscopy/ spectrophotometry (UV-Vis), mass spectrometry (MS), and nuclear magnetic resonance spectroscopy (NMR).

**[0085]** As used herein, the term "undetectable concentration" refers to a level of a compound that is too low to be measured and/or analyzed by techniques such as TLC, HPLC, UV-Vis, MS, or NMR. In some embodiments, a compound of an "undetectable concentration" is not present in a steviol glycoside or steviol glycoside precursor composition.

**[0086]** As used herein, the terms "or" and "and/or" is utilized to describe multiple components in combination or exclusive of one another. For example, "x, y, and/or z" can refer to "x" alone, "y" alone, "z" alone, "x, y, and z," "(x and y) or z," "x or (y and z)," or "x or y or z." In some embodiments, "and/or" is used to refer to the exogenous nucleic acids that a recombinant cell comprises, wherein a recombinant cell comprises one or more exogenous nucleic acids selected from a group. In some embodiments, "and/or" is used to refer to production of steviol glycosides and/or steviol glycoside precursors. In some embodiments, "and/or" is used to refer to production of steviol glycosides, wherein one or more steviol glycosides are produced. In some embodiments, "and/or" is used to refer to production of steviol glycosides, wherein one or more steviol glycosides are produced through one or more of the following steps: culturing a recombinant microorganism, synthesizing one or more steviol glycosides in a recombinant microorganism, and/or isolating one or more steviol glycosides.

**[0087]** In some embodiments, the nucleotide sequence of a nucleic acid encoding a KO polypeptide is set forth in SEQ ID NO: 55, SEQ ID NO:56, SEQ ID NO:57, SEQ ID NO:58, SEQ ID NO:59, or SEQ ID NO:60, SEQ ID NO:63, SEQ ID NO:64, or SEQ ID NO:65. In some aspects, the nucleic acid encoding the KO polypeptide has at least 70% identity to the nucleotide sequence set forth in SEQ ID NO:55, SEQ ID NO:57, SEQ ID NO:59 or SEQ ID NO:60, at least 80% identity to the nucleotide sequence set forth in SEQ ID NO:56 or SEQ ID NO:58, at least 95% identity to the nucleotide sequence set forth in SEQ ID NO:63, or at least

75% identity to the nucleotide sequence set forth in SEQ ID NO:64 or SEQ ID NO:65. In some embodiments, the amino acid sequence of a KO enzyme is set forth in SEQ ID NO:54, SEQ ID NO:70, SEQ ID NO:71, SEQ ID NO:72, SEQ ID NO:75, SEQ ID NO:77, SEQ ID NO:78, OR SEQ ID NO:79. In some embodiments, a host cell comprises one or more copies of one or more nucleic acids encoding a KO polypeptide.

[0088] In some embodiments, expression of a KO gene set forth in SEQ ID NO:55 or SEQ ID NO:56 in a RebB-producing *S. cerevisiae* strain results in higher production of RebB compared to expression of SrKOl (SEQ ID NO:59, SEQ ID NO:79) in a RebB-producing *S. cerevisiae* strain. See Example 3.

[0089] In some embodiments, expression of a KO gene set forth in SEQ ID NO:55, SEQ ID NO:56, or SEQ ID NO:57 in an *S. cerevisiae* strain capable of producing RebB with a functional KO results in production of ent-kaurenoic acid. See Example 3.

[0090] As used herein, the terms "ent-kaurenoic acid hydroxylase" and "steviol synthase" can be used interchangeably and be abbreviated "KAH." In some embodiments, the nucleotide sequence of a nucleic acid encoding a KAH enzyme is set forth in SEQ ID NO:18, SEQ ID NO:80, SEQ ID NO:81, SEQ ID NO:90, or SEQ ID NO:96. In some aspects, the nucleic acid encoding the KAH polypeptide has at least 75% identity to a nucleotide sequence set forth in SEQ ID NO:80; or at least 70% identity to a nucleotide sequence set forth in SEQ ID NO:18, SEQ ID NO:81, SEQ ID NO:90, or SEQ ID NO:96. In some embodiments, the amino acid sequence of a KAH enzyme is set forth in SEQ ID NO:68, SEQ ID NO:82, or SEQ ID NO:91. In some embodiments, a host cell comprises one or more copies of one or more nucleic acids encoding a KAH enzyme.

[0091] In some embodiments, one or more copies of SrKAH1 (SEQ ID NO:18, SEQ ID NO:68) are expressed in an *S. cerevisiae* strain. For example, in some embodiments, two copies of SrKAH1 (SEQ ID NO:18, SEQ ID NO:68) are expressed in an *S. cerevisiae* strain.

[0092] In some embodiments, the nucleotide sequence of a nucleic acid encoding a KAH enzyme is set forth in SEQ ID NO:80. The nucleic acid of SEQ ID NO:80 encodes a KAH with an amino acid sequence set forth in SEQ ID NO:82. A version of SEQ ID NO:80 codon-optimized for expression in *S. cerevisiae* is set forth in SEQ ID NO:81. In some embodiments, a host cell comprises one or more copies of one or more nucleic acids encoding a KAH enzyme. See Example 7.

[0093] In some embodiments, SrKAHel (SEQ ID NO:18, SEQ ID NO:68) and either the KAH encoded by the nucleotide sequence set forth in SEQ ID NO:80 or the KAH encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:81 are co-expressed in a steviol glycoside-producing *S. cerevisiae* strain. In some embodiments, co-expression of SrKAHel (SEQ ID NO:18, SEQ ID NO:68) and either the KAH encoded by the nucleotide sequence set forth in SEQ ID NO:80 or the KAH encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:81 in a steviol glycoside-producing strain results in higher production of steviol glycosides compared to a control steviol glycoside-producing strain or a steviol glycoside producing strain overexpressing SrKAHel. See Example 7 and Table 6. In some aspects, overexpressing SrKAHel results in production of 85.5  $\mu\text{M}$  13-SMG, expression of SrKAHel and the KAH encoded by the nucleotide set forth in SEQ ID NO:80 results in production of 153.8  $\mu\text{M}$  13-SMG, and expression of SrKAHel and the KAH encoded by the nucleotide set forth in SEQ ID NO:81 results in production of 130.5  $\mu\text{M}$  13-SMG.

[0094] In some embodiments, a KO gene is expressed in a steviol glycoside-producing *S. cerevisiae* strain that further overexpresses SrKAHel (SEQ ID NO:18, SEQ ID NO:68). In some embodiments, expression of a KO gene of SEQ ID NO:55, SEQ ID NO:56, SEQ ID NO:57, SEQ ID NO:58, SEQ ID NO:59, or SEQ ID NO:60, SEQ ID NO:65 in a steviol glycoside-producing *S. cerevisiae* strain overexpressing SrKAHel results in higher expression of steviol glycosides compared to a control steviol-glycoside producing strain or a steviol glycoside-producing strain overexpressing SrKAHel (SEQ ID NO:18, SEQ ID NO:68). See Example 4.

[0095] In some embodiments, expression of a KO gene of SEQ ID NO:55, SEQ ID NO:56, SEQ ID NO:57, or SEQ ID NO:60 in a steviol glycoside-producing *S. cerevisiae* strain overexpressing SrKAHel (SEQ ID NO:18, SEQ ID NO:68) results in higher levels of glycosylated ent-kaurenoic acid compared to a control *S. cerevisiae* strain. See Example 4.

[0096] In some embodiments, expression of a KO gene of SEQ ID NO:55, SEQ ID NO:56, SEQ ID NO:57, SEQ ID NO:59, or SEQ ID NO:60 in a steviol glycoside-producing *S. cerevisiae* strain overexpressing SrKAHel (SEQ ID NO:18, SEQ ID NO:68) results in improved metabolic conversion of a glycosylated ent-kaurenol intermediate compound relative to a control *S. cerevisiae* strain or a steviol glycoside-producing *S. cerevisiae* strain overexpressing SrKAHel (SEQ ID NO:18, SEQ ID NO:68). See Example 4.

[0097] In some embodiments, a KAH is a *Prunus* KAH, such as a *Prunus avium*, *Prunus mume*, or *Prunus persica* KAH. In some embodiments, a KAH is a KAH of the CYP72A219 or CYP71A219-like family. In some embodiments, the nucleotide sequence of a nucleic acid



encoding a KAH enzyme is set forth in SEQ ID NO:90 or SEQ ID NO:96. The nucleic acids of SEQ ID NO:90 and SEQ ID NO:96 encode a KAH from *Prunus avium* with an amino acid sequence set forth in SEQ ID NO:91. In some embodiments, a KAH polypeptide is a polypeptide with an amino acid sequence set forth in SEQ ID NO:92, SEQ ID NO:93, SEQ ID NO:94, or SEQ ID NO:95. In some embodiments, a KAH polypeptide is a KAH polypeptide with at least 50% sequence identity to an amino acid sequence set forth in SEQ ID NO:91, SEQ ID NO:92, SEQ ID NO:93, SEQ ID NO:94, or SEQ ID NO:95. In some embodiments, expression of a gene encoding a polypeptide having at least 50% sequence identity to an amino acid sequence set forth in SEQ ID NO:91, SEQ ID NO:92, SEQ ID NO:93, SEQ ID NO:94, or SEQ ID NO:95 in a recombinant host results in production of a steviol glycoside or steviol glycoside precursor, such as 13-SMG and/or rubusoside. See Example 8.

**[0098]** In some embodiments, the nucleotide sequence of the nucleic acid encoding a CPR enzyme is set forth in SEQ ID NO:23, SEQ ID NO:51, SEQ ID NO:61, SEQ ID NO:62, SEQ ID NO:66, SEQ ID NO:67, or SEQ ID NO:97. In some aspects, the nucleic acid encoding the CPR polypeptide has at least 75% identity to the nucleotide sequence set forth in SEQ ID NO:23, SEQ ID NO:61, or SEQ ID NO:62, or at least 70% identity to the nucleotide sequence set forth in SEQ ID NO:24, SEQ ID NO:66, SEQ ID NO:67, SEQ ID NO:51, or SEQ ID NO:97. In some embodiments, the amino acid sequence of the CPR enzyme is set forth in SEQ ID NO:22, SEQ ID NO:28, SEQ ID NO:69, SEQ ID NO:73, SEQ ID NO:74, or SEQ ID NO:76, SEQ ID NO:87, or SEQ ID NO:98. In some embodiments, a host cell comprises one or more copies of one or more nucleic acids encoding a CPR enzyme.

**[0099]** In a non-limiting example, SrKAHel is activated by the *S. cerevisiae* CPR encoded by gene NCP1 (YHR042W). Enhanced activation of the KAH encoded by SrKAHel is observed when the *Arabidopsis thaliana* CPR encoded by the gene ATR2 (SEQ ID NO:51) or the *S. rebaudiana* CPR encoded by the genes CPR7 (SEQ ID NO:23) or CPR8 (SEQ ID NO:24, SEQ ID NO:28) are co-expressed in a recombinant cell. Amino acid sequences of the *A. thaliana* polypeptides ATR1 and ATR2 are set forth in SEQ ID NO:25 and SEQ ID NO:26, respectively. The *S. rebaudiana* polypeptides CPR7 and CPR8 are set forth in SEQ ID NO:27 and SEQ ID NO:28, respectively.

**[00100]** In some embodiments, expression of CPR1 (SEQ ID NO:61, SEQ ID NO:76) or of CPR7 in the steviol glycoside-producing *S. cerevisiae* strain co-expressing *S. rebaudiana* CPR8 (SEQ ID NO:24, SEQ ID NO:28) and *A. thaliana* ATR2 (SEQ ID NO:51) results in higher levels of RebM compared to a control steviol glycoside-producing *S. cerevisiae* strain expressing *S.*

*rebaudiana* CPR8 (SEQ ID NO:24, SEQ ID NO:28) and *A. thaliana* ATR2 (SEQ ID NO:51). In some embodiments, expression of the CPR set forth in SEQ ID NO:62 in a steviol glycoside-producing *S. cerevisiae* strain overexpressing SrKAHel (SEQ ID NO:18, SEQ ID NO:68) results in higher levels of RebM compared to a steviol glycoside-producing *S. cerevisiae* strain that does not express the nucleic acid set forth in SEQ ID NO:62 or overexpress SrKAHel. See Example 5.

**[00101]** In some embodiments, co-expression of SrKOl (SEQ ID NO:59, SEQ ID NO:79) and a CPR gene of SEQ ID NO:66 or SEQ ID NO:77 in a RebB-producing strain results in higher production of 13-SMG and RebB than co-expression of a KO gene of SEQ ID NO:63 or SEQ ID NO:64 and a CPR gene of SEQ ID NO:66 or SEQ ID NO:77. See Example 6.

**[00102]** In some embodiments, CPR1 (SEQ ID NO:61, SEQ ID NO:76) or CPR12 (SEQ ID NO:97, SEQ ID NO:98) activates cytochrome c. In some embodiments, CPR1 (SEQ ID NO:61, SEQ ID NO:76) or CPR12 (SEQ ID NO:97, SEQ ID NO:98) in the presence of SrKAHel (SEQ ID NO:18, SEQ ID NO:68) activate cytochrome c. In some embodiments, CPR1 (SEQ ID NO:61, SEQ ID NO:76) or CPR12 (SEQ ID NO:97, SEQ ID NO:98) regulate conversion of ent-kaurenoic acid to steviol. In some embodiments, CPR1 (SEQ ID NO:61, SEQ ID NO:76) or CPR12 (SEQ ID NO:97, SEQ ID NO:98) in combination with SrKAHel (SEQ ID NO:18, SEQ ID NO:68) convert ent-kaurenoic acid to steviol. In some embodiments, steviol production is detected upon incubation of ent-kaurenoic acid with microsomal protein prepared from *S. cerevisiae* strains expressing CPR1 (SEQ ID NO:61, SEQ ID NO:76) or CPR12 (SEQ ID NO:97, SEQ ID NO:98) in combination with SrKAHel (SEQ ID NO:18, SEQ ID NO:68). In some embodiments, expression of CPR1 (SEQ ID NO:61, SEQ ID NO:76) or CPR12 (SEQ ID NO:97, SEQ ID NO:98) in a recombinant host results in production of a steviol glycoside or steviol glycoside precursor. See Example 9.

**[00103]** In some embodiments, a steviol glycoside-producing strain expresses a fusion construct comprising a KO and the NADPH-dependent P450 oxidoreductase domain of CYP102A1, referred to herein as "BMR." The codon-optimized nucleotide sequence encoding the BMR polypeptide is set forth in SEQ ID NO:117; the BMR amino acid sequence is set forth in SEQ ID NO:118. In some embodiments, BMR is a mutant BMR, including, but not limited to a BMR W1046A mutant (SEQ ID NO:119, SEQ ID NO:120). The BMR mutant can be specific for NADH. In some embodiments, the KO-BMR fusion construct comprises a linker (SEQ ID NO:121, SEQ ID NO:122). In some embodiments, the KO of the fusion construct is SrKOl (SEQ ID NO:59, SEQ ID NO:79) or the KO encoded by the nucleotide sequence set forth in

SEQ ID NO:65 (corresponding to the amino acid sequence set forth in SEQ ID NO:75). In some embodiments, the KO of the fusion construct is a truncated KO. Exemplary KO-BMR fusion constructs are set forth in SEQ ID NOs:99-112. See Example 10.

**[00104]** In some embodiments, expression of SrK01-BMR fusion constructs (SEQ ID NOs:99-106) in a steviol glycoside-producing strain results in an increase in ent-kaurenoic acid, 13-SMG, and RebB levels, compared to expression of SrK01 (SEQ ID NO:59, SEQ ID NO:79) in a steviol glycoside-producing strain. In some embodiments, expression of a fusion construct (SEQ ID NO:107, SEQ ID NO:108) in a steviol glycoside-producing strain results in greater conversion of ent-kaurene to ent-kaurenoic acid and greater conversion of ent-kaurenoic acid to 13-SMG, compared to expression of the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 in a steviol glycoside-producing strain. In some embodiments, expression of a fusion construct comprising the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 and the W1046A mutant BMR (SEQ ID NO:109, SEQ ID NO:110) results in increased ent-kaurenoic acid levels. See Figure 16 (B and D) and Example 10.

**[00105]** In some embodiments, a steviol glycoside-producing strain comprises inheritance of cortical ER protein 2 (ICE2; SEQ ID NO:113, SEQ ID NO:114). ICE2 is also referred to as YIL090W. In some aspects, ICE2 is overexpressed. ICE2 can be expressed in a strain comprising CPR1 (SEQ ID NO:61, SEQ ID NO:76) and/or CPR12 (SEQ ID NO:97, SEQ ID NO:98). In some embodiments, a steviol glycoside-producing strain comprises two copies of ICE2. In some embodiments, expression of ICE2 increases ent-kaurene metabolism (resulting in decreased accumulation of ent-kaurene, ent-kaurenol, ent-kaurenal, and ent-kaurenol glycosides), resulting in increased accumulation of steviol glycosides, compared to a control strain. See Table 10 and Example 11.

**[00106]** In some embodiments, expression of the KO encoded by nucleotide sequence set forth in SEQ ID NO:56 in a steviol glycoside-producing strain cultivated by fermentation results in a lower accumulation of ent-kaurene compounds, compared to a control steviol glycoside-producing strain. In some aspects, higher levels of ent-kaurenoic acid and steviol glycosides result, as compared to a control strain. In some embodiments, expression of the KAH encoded by nucleotide sequence set forth in SEQ ID NO:80, the KO encoded by nucleotide sequence set forth in SEQ ID NO:56, and the KO encoded by nucleotide sequence set forth in SEQ ID NO:65 in a steviol glycoside-producing strain cultivated by fermentation results in decreased accumulation of ent-kaurene, ent-kaurenol, ent-kaurenal, ent-kaurenol glycosides, ent-kaurenoic acid, and ent-kaurenoic acid glycosides and increased production of steviol glycosides, as

compared to a control strain. In some embodiments, expression of CPR12 (SEQ ID NO:97, SEQ ID NO:98), the KAH encoded by nucleotide sequence set forth in SEQ ID NO:80, and the KO encoded by nucleotide sequence set forth in SEQ ID NO:56 cultivated by fermentation results in decreased ent-kaurene, ent-kaurenol, ent-kaurenal, ent-kaurenol glycosides, ent-kaurenoic acid, and ent-kaurenoic acid glycosides accumulation and higher levels of steviol glycosides, as compared to a control strain. See Table 12 and Example 12.

### **Functional Homologs**

**[00107]** Functional homologs of the polypeptides described above are also suitable for use in producing steviol glycosides in a recombinant host. A functional homolog is a polypeptide that has sequence similarity to a reference polypeptide, and that carries out one or more of the biochemical or physiological function(s) of the reference polypeptide. A functional homolog and the reference polypeptide can be a natural occurring polypeptide, and the sequence similarity can be due to convergent or divergent evolutionary events. As such, functional homologs are sometimes designated in the literature as homologs, or orthologs, or paralogs. Variants of a naturally occurring functional homolog, such as polypeptides encoded by mutants of a wild type coding sequence, can themselves be functional homologs. Functional homologs can also be created via site-directed mutagenesis of the coding sequence for a polypeptide, or by combining domains from the coding sequences for different naturally-occurring polypeptides ("domain swapping"). Techniques for modifying genes encoding functional polypeptides described herein are known and include, *inter alia*, directed evolution techniques, site-directed mutagenesis techniques and random mutagenesis techniques, and can be useful to increase specific activity of a polypeptide, alter substrate specificity, alter expression levels, alter subcellular location, or modify polypeptide-polypeptide interactions in a desired manner. Such modified polypeptides are considered functional homologs. The term "functional homolog" is sometimes applied to the nucleic acid that encodes a functionally homologous polypeptide.

**[00108]** Functional homologs can be identified by analysis of nucleotide and polypeptide sequence alignments. For example, performing a query on a database of nucleotide or polypeptide sequences can identify homologs of steviol glycoside biosynthesis polypeptides. Sequence analysis can involve BLAST, Reciprocal BLAST, or PSI-BLAST analysis of non-redundant databases using a KO, KAH, or CPR amino acid sequence as the reference sequence. Amino acid sequence is, in some instances, deduced from the nucleotide sequence. Those polypeptides in the database that have greater than 40% sequence identity are candidates for further evaluation for suitability as a steviol glycoside biosynthesis polypeptide.

Amino acid sequence similarity allows for conservative amino acid substitutions, such as substitution of one hydrophobic residue for another or substitution of one polar residue for another. If desired, manual inspection of such candidates can be carried out in order to narrow the number of candidates to be further evaluated. Manual inspection can be performed by selecting those candidates that appear to have domains present in steviol glycoside biosynthesis polypeptides, e.g., conserved functional domains. In some embodiments, nucleic acids and polypeptides are identified from transcriptome data based on expression levels rather than by using BLAST analysis.

[00109] Conserved regions can be identified by locating a region within the primary amino acid sequence of a steviol glycoside biosynthesis polypeptide that is a repeated sequence, forms some secondary structure (e.g., helices and beta sheets), establishes positively or negatively charged domains, or represents a protein motif or domain. See, e.g., the Pfam web site describing consensus sequences for a variety of protein motifs and domains on the World Wide Web at [sanger.ac.uk/Software/Pfam/](http://sanger.ac.uk/Software/Pfam/) and [pfam.janelia.org/](http://pfam.janelia.org/). The information included at the Pfam database is described in Sonnhammer *et al.*, *Nucl. Acids Res.*, 26:320-322 (1998); Sonnhammer *et al.*, *Proteins*, 28:405-420 (1997); and Bateman *et al.*, *Nucl. Acids Res.*, 27:260-262 (1999). Conserved regions also can be determined by aligning sequences of the same or related polypeptides from closely related species. Closely related species preferably are from the same family. In some embodiments, alignment of sequences from two different species is adequate to identify such homologs.

[00110] Typically, polypeptides that exhibit at least about 40% amino acid sequence identity are useful to identify conserved regions. Conserved regions of related polypeptides exhibit at least 45% amino acid sequence identity (e.g., at least 50%, at least 60%, at least 70%, at least 80%, or at least 90% amino acid sequence identity). In some embodiments, a conserved region exhibits at least 92%, 94%, 96%, 98%, or 99% amino acid sequence identity.

[00111] For example, polypeptides suitable for producing steviol in a recombinant host include functional homologs of KO, KAH, and CPR.

[00112] Methods to modify the substrate specificity of, for example, KO, KAH, or CPR, are known to those skilled in the art, and include without limitation site-directed/rational mutagenesis approaches, random directed evolution approaches and combinations in which random mutagenesis/saturation techniques are performed near the active site of the enzyme. For example see Osmani *et al.*, 2009, *Phytochemistry* 70: 325-347.

**[00113]** A candidate sequence typically has a length that is from 80% to 200% of the length of the reference sequence, e.g., 82, 85, 87, 89, 90, 93, 95, 97, 99, 100, 105, 110, 115, 120, 130, 140, 150, 160, 170, 180, 190, or 200% of the length of the reference sequence. A functional homolog polypeptide typically has a length that is from 95% to 105% of the length of the reference sequence, e.g., 90, 93, 95, 97, 99, 100, 105, 110, 115, or 120% of the length of the reference sequence, or any range between. A% identity for any candidate nucleic acid or polypeptide relative to a reference nucleic acid or polypeptide can be determined as follows. A reference sequence (e.g., a nucleic acid sequence or an amino acid sequence described herein) is aligned to one or more candidate sequences using the computer program ClustalW (version 1.83, default parameters), which allows alignments of nucleic acid or polypeptide sequences to be carried out across their entire length (global alignment). Chenna *et al.*, 2003, *Nucleic Acids Res.* 31(13):3497-500.

**[00114]** ClustalW calculates the best match between a reference and one or more candidate sequences, and aligns them so that identities, similarities and differences can be determined. Gaps of one or more residues can be inserted into a reference sequence, a candidate sequence, or both, to maximize sequence alignments. For fast pairwise alignment of nucleic acid sequences, the following default parameters are used: word size: 2; window size: 4; scoring method: % age; number of top diagonals: 4; and gap penalty: 5. For multiple alignment of nucleic acid sequences, the following parameters are used: gap opening penalty: 10.0; gap extension penalty: 5.0; and weight transitions: yes. For fast pairwise alignment of protein sequences, the following parameters are used: word size: 1; window size: 5; scoring method: % age; number of top diagonals: 5; gap penalty: 3. For multiple alignment of protein sequences, the following parameters are used: weight matrix: blosum; gap opening penalty: 10.0; gap extension penalty: 0.05; hydrophilic gaps: on; hydrophilic residues: Gly, Pro, Ser, Asn, Asp, Gin, Glu, Arg, and Lys; residue-specific gap penalties: on. The ClustalW output is a sequence alignment that reflects the relationship between sequences. ClustalW can be run, for example, at the Baylor College of Medicine Search Launcher site on the World Wide Web ([searchlauncher.bcm.tmc.edu/multi-align/multi-align.html](http://searchlauncher.bcm.tmc.edu/multi-align/multi-align.html)) and at the European Bioinformatics Institute site on the World Wide Web ([ebi.ac.uk/clustalw](http://ebi.ac.uk/clustalw)).

**[00115]** To determine % identity of a candidate nucleic acid or amino acid sequence to a reference sequence, the sequences are aligned using ClustalW, the number of identical matches in the alignment is divided by the length of the reference sequence, and the result is multiplied by 100. It is noted that the % identity value can be rounded to the nearest tenth. For

example, 78.11, 78.12, 78.13, and 78.14 are rounded down to 78.1, while 78.15, 78.16, 78.17, 78.18, and 78.19 are rounded up to 78.2.

**[001 16]** It will be appreciated that functional KO, KAH, or CPR proteins can include additional amino acids that are not involved in the enzymatic activities carried out by the enzymes. In some embodiments, KO, KAH, or CPR proteins are fusion proteins. The terms "chimera," "fusion polypeptide," "fusion protein," "fusion enzyme," "fusion construct," "chimeric protein," "chimeric polypeptide," "chimeric construct," and "chimeric enzyme" can be used interchangeably herein to refer to proteins engineered through the joining of two or more genes that code for different proteins. In some embodiments, a nucleic acid sequence encoding a KO, KAH, or CPR polypeptide can include a tag sequence that encodes a "tag" designed to facilitate subsequent manipulation (*e.g.*, to facilitate purification or detection), secretion, or localization of the encoded polypeptide. Tag sequences can be inserted in the nucleic acid sequence encoding the polypeptide such that the encoded tag is located at either the carboxyl or amino terminus of the polypeptide. Non-limiting examples of encoded tags include green fluorescent protein (GFP), human influenza hemagglutinin (HA), glutathione S transferase (GST), polyhistidine-tag (HIS tag), and Flag™ tag (Kodak, New Haven, CT). Other examples of tags include a chloroplast transit peptide, a mitochondrial transit peptide, an amyloplast peptide, signal peptide, or a secretion tag.

**[001 17]** In some embodiments, a fusion protein is a protein altered by domain swapping. As used herein, the term "domain swapping" is used to describe the process of replacing a domain of a first protein with a domain of a second protein. In some embodiments, the domain of the first protein and the domain of the second protein are functionally identical or functionally similar. In some embodiments, the structure and/or sequence of the domain of the second protein differs from the structure and/or sequence of the domain of the first protein. In some embodiments, a KO polypeptide is altered by domain swapping. See Example 10.

#### **Steviol and Steviol Glycoside Biosynthesis Nucleic Acids**

**[001 18]** A recombinant gene encoding a polypeptide described herein comprises the coding sequence for that polypeptide, operably linked in sense orientation to one or more regulatory regions suitable for expressing the polypeptide. Because many microorganisms are capable of expressing multiple gene products from a polycistronic mRNA, multiple polypeptides can be expressed under the control of a single regulatory region for those microorganisms, if desired. A coding sequence and a regulatory region are considered to be operably linked when the regulatory region and coding sequence are positioned so that the regulatory region is effective

for regulating transcription or translation of the sequence. Typically, the translation initiation site of the translational reading frame of the coding sequence is positioned between one and about fifty nucleotides downstream of the regulatory region for a monocistronic gene.

[00119] In many cases, the coding sequence for a polypeptide described herein is identified in a species other than the recombinant host, *i.e.*, is a heterologous nucleic acid. Thus, if the recombinant host is a microorganism, the coding sequence can be from other prokaryotic or eukaryotic microorganisms, from plants or from animals. In some case, however, the coding sequence is a sequence that is native to the host and is being reintroduced into that organism. A native sequence can often be distinguished from the naturally occurring sequence by the presence of non-natural sequences linked to the exogenous nucleic acid, *e.g.*, non-native regulatory sequences flanking a native sequence in a recombinant nucleic acid construct. In addition, stably transformed exogenous nucleic acids typically are integrated at positions other than the position where the native sequence is found. "Regulatory region" refers to a nucleic acid having nucleotide sequences that influence transcription or translation initiation and rate, and stability and/or mobility of a transcription or translation product. Regulatory regions include, without limitation, promoter sequences, enhancer sequences, response elements, protein recognition sites, inducible elements, protein binding sequences, 5' and 3' untranslated regions (UTRs), transcriptional start sites, termination sequences, polyadenylation sequences, introns, and combinations thereof. A regulatory region typically comprises at least a core (basal) promoter. A regulatory region also may include at least one control element, such as an enhancer sequence, an upstream element or an upstream activation region (UAR). A regulatory region is operably linked to a coding sequence by positioning the regulatory region and the coding sequence so that the regulatory region is effective for regulating transcription or translation of the sequence. For example, to operably link a coding sequence and a promoter sequence, the translation initiation site of the translational reading frame of the coding sequence is typically positioned between one and about fifty nucleotides downstream of the promoter. A regulatory region can, however, be positioned as much as about 5,000 nucleotides upstream of the translation initiation site, or about 2,000 nucleotides upstream of the transcription start site.

[00120] The choice of regulatory regions to be included depends upon several factors, including, but not limited to, efficiency, selectability, inducibility, desired expression level, and preferential expression during certain culture stages. It is a routine matter for one of skill in the art to modulate the expression of a coding sequence by appropriately selecting and positioning regulatory regions relative to the coding sequence. It will be understood that more than one



regulatory region may be present, e.g., introns, enhancers, upstream activation regions, transcription terminators, and inducible elements.

[00121] One or more genes can be combined in a recombinant nucleic acid construct in "modules" useful for a discrete aspect of steviol and/or steviol glycoside production. Combining a plurality of genes in a module, particularly a polycistronic module, facilitates the use of the module in a variety of species. For example, a steviol biosynthesis gene cluster, or a UGT gene cluster, can be combined in a polycistronic module such that, after insertion of a suitable regulatory region, the module can be introduced into a wide variety of species. As another example, a UGT gene cluster can be combined such that each UGT coding sequence is operably linked to a separate regulatory region, to form a UGT module. Such a module can be used in those species for which monocistronic expression is necessary or desirable. In addition to genes useful for steviol or steviol glycoside production, a recombinant construct typically also contains an origin of replication, and one or more selectable markers for maintenance of the construct in appropriate species.

[00122] It will be appreciated that because of the degeneracy of the genetic code, a number of nucleic acids can encode a particular polypeptide; *i.e.*, for many amino acids, there is more than one nucleotide triplet that serves as the codon for the amino acid. Thus, codons in the coding sequence for a given polypeptide can be modified such that optimal expression in a particular host is obtained, using appropriate codon bias tables for that host (e.g., microorganism). As isolated nucleic acids, these modified sequences can exist as purified molecules and can be incorporated into a vector or a virus for use in constructing modules for recombinant nucleic acid constructs.

[0003] In some cases, it is desirable to inhibit one or more functions of an endogenous polypeptide in order to divert metabolic intermediates towards steviol or steviol glycoside biosynthesis. For example, it may be desirable to downregulate synthesis of sterols in a yeast strain in order to further increase steviol or steviol glycoside production, e.g., by downregulating squalene epoxidase. As another example, it may be desirable to inhibit degradative functions of certain endogenous gene products, e.g., glycohydrolases that remove glucose moieties from secondary metabolites or phosphatases as discussed herein. In such cases, a nucleic acid that overexpresses the polypeptide or gene product may be included in a recombinant construct that is transformed into the strain. Alternatively, mutagenesis can be used to generate mutants in genes for which it is desired to increase or enhance function.

### **Host Microorganisms**

[00123] Recombinant hosts can be used to express polypeptides for the producing steviol glycosides, including mammalian, insect, plant, and algal cells. A number of prokaryotes and eukaryotes are also suitable for use in constructing the recombinant microorganisms described herein, e.g., gram-negative bacteria, yeast, and fungi. A species and strain selected for use as a steviol glycoside production strain is first analyzed to determine which production genes are endogenous to the strain and which genes are not present. Genes for which an endogenous counterpart is not present in the strain are advantageously assembled in one or more recombinant constructs, which are then transformed into the strain in order to supply the missing function(s).

[00124] Typically, the recombinant microorganism is grown in a fermenter at a defined temperature(s) for a desired period of time. The constructed and genetically engineered microorganisms provided by the invention can be cultivated using conventional fermentation processes, including, *inter alia*, chemostat, batch, fed-batch cultivations, semi-continuous fermentations such as draw and fill, continuous perfusion fermentation, and continuous perfusion cell culture. Depending on the particular microorganism used in the method, other recombinant genes such as isopentenyl biosynthesis genes and terpene synthase and cyclase genes may also be present and expressed. Levels of substrates and intermediates, e.g., isopentenyl diphosphate, dimethylallyl diphosphate, GGPP, ent-kaurene and ent-kaurenoic acid, can be determined by extracting samples from culture media for analysis according to published methods.

[00125] Carbon sources of use in the instant method include any molecule that can be metabolized by the recombinant host cell to facilitate growth and/or production of the steviol glycosides. Examples of suitable carbon sources include, but are not limited to, sucrose (e.g., as found in molasses), fructose, xylose, ethanol, glycerol, glucose, cellulose, starch, cellobiose or other glucose-comprising polymer. In embodiments employing yeast as a host, for example, carbon sources such as sucrose, fructose, xylose, ethanol, glycerol, and glucose are suitable. The carbon source can be provided to the host organism throughout the cultivation period or alternatively, the organism can be grown for a period of time in the presence of another energy source, e.g., protein, and then provided with a source of carbon only during the fed-batch phase.

[00126] After the recombinant microorganism has been grown in culture for the desired period of time, steviol and/or one or more steviol glycosides can then be recovered from the culture using various techniques known in the art. In some embodiments, a permeabilizing

agent can be added to aid the feedstock entering into the host and product getting out. For example, a crude lysate of the cultured microorganism can be centrifuged to obtain a supernatant. The resulting supernatant can then be applied to a chromatography column, e.g., a C-18 column, and washed with water to remove hydrophilic compounds, followed by elution of the compound(s) of interest with a solvent such as methanol. The compound(s) can then be further purified by preparative HPLC. See also, WO 2009/140394.

[00127] It will be appreciated that the various genes and modules discussed herein can be present in two or more recombinant hosts rather than a single host. When a plurality of recombinant hosts is used, they can be grown in a mixed culture to accumulate steviol and/or steviol glycosides.

[00128] Alternatively, the two or more hosts each can be grown in a separate culture medium and the product of the first culture medium, e.g., steviol, can be introduced into second culture medium to be converted into a subsequent intermediate, or into an end product such as, for example, RebA. The product produced by the second, or final host is then recovered. It will also be appreciated that in some embodiments, a recombinant host is grown using nutrient sources other than a culture medium and utilizing a system other than a fermenter.

[00129] Exemplary prokaryotic and eukaryotic species are described in more detail below. However, it will be appreciated that other species can be suitable. For example, suitable species can be in a genus such as *Agaricus*, *Aspergillus*, *Bacillus*, *Candida*, *Corynebacterium*, *Eremothecium*, *Escherichia*, *Fusarium/Gibberella*, *Kluyveromyces*, *Laetiporus*, *Lentinus*, *Phaffia*, *Phanerochaete*, *Pichia*, *Physcomitrella*, *Rhodoturula*, *Saccharomyces*, *Schizosaccharomyces*, *Sphaceloma*, *Xanthophyllomyces* or *Yarrowia*. Exemplary species from such genera include *Lentinus tigrinus*, *Laetiporus sulphureus*, *Phanerochaete chrysosporium*, *Pichia pastoris*, *Cyberlindnera jadinii*, *Physcomitrella patens*, *Rhodoturula glutinis*, *Rhodoturula mucilaginoso*, *Phaffia rhodozyma*, *Xanthophyllomyces dendrorhous*, *Fusarium fujikuroi/Gibberella fujikuroi*, *Candida utilis*, *Candida glabrata*, *Candida albicans*, and *Yarrowia lipolytica*.

[00130] In some embodiments, a microorganism can be a prokaryote such as *Escherichia* bacteria cells, for example, *Escherichia coli* cells; *Lactobacillus* bacteria cells; *Lactococcus* bacteria cells; *Cornebacterium* bacteria cells; *Acetobacter* bacteria cells; *Acinetobacter* bacteria cells; or *Pseudomonas* bacterial cells.

[00131] In some embodiments, a microorganism can be an Ascomycete such as *Gibberella fujikuroi*, *Kluyveromyces lactis*, *Schizosaccharomyces pombe*, *Aspergillus niger*, *Yarrowia lipolytica*, *Ashbya gossypii*, or *S. cerevisiae*.

[00132] in some embodiments, a microorganism can be an algal cell such as *Blakeslea trispora*, *Dunaliella salina*, *Haematococcus pluvialis*, *Chlorella sp.*, *Undaria pinnatifida*, *Sargassum*, *Laminaria japonica*, *Scenedesmus almeriensis* species.

[00133] In some embodiments, a microorganism can be a cyanobacterial cell such as *Blakeslea trispora*, *Dunaliella salina*, *Haematococcus pluvialis*, *Chlorella sp.*, *Undaria pinnatifida*, *Sargassum*, *Laminaria japonica*, *Scenedesmus almeriensis*.

#### Saccharomyces spp.

[00134] *Saccharomyces* is a widely used chassis organism in synthetic biology, and can be used as the recombinant microorganism platform. For example, there are libraries of mutants, plasmids, detailed computer models of metabolism and other information available for *S. cerevisiae*, allowing for rational design of various modules to enhance product yield. Methods are known for making recombinant microorganisms.

#### Aspergillus spp.

[00135] *Aspergillus* species such as *A. oryzae*, *A. niger* and *A. sojae* are widely used microorganisms in food production and can also be used as the recombinant microorganism platform. Nucleotide sequences are available for genomes of *A. nidulans*, *A. fumigatus*, *A. oryzae*, *A. clavatus*, *A. flavus*, *A. niger*, and *A. terreus*, allowing rational design and modification of endogenous pathways to enhance flux and increase product yield. Metabolic models have been developed for *Aspergillus*, as well as transcriptomic studies and proteomics studies. *A. niger* is cultured for the industrial production of a number of food ingredients such as citric acid and gluconic acid, and thus species such as *A. niger* are generally suitable for producing steviol glycosides.

#### E. coli

[00136] *E. coli*, another widely used platform organism in synthetic biology, can also be used as the recombinant microorganism platform. Similar to *Saccharomyces*, there are libraries of mutants, plasmids, detailed computer models of metabolism and other information available for *E. coli*, allowing for rational design of various modules to enhance product yield. Methods

similar to those described above for *Saccharomyces* can be used to make recombinant *E. coli* microorganisms.

*Agaricus, Gibberella, and Phanerochaete* spp.

[00137] *Agaricus, Gibberella, and Phanerochaete* spp. can be useful because they are known to produce large amounts of isoprenoids in culture. Thus, the terpene precursors for producing large amounts of steviol glycosides are already produced by endogenous genes. Thus, modules comprising recombinant genes for steviol glycoside biosynthesis polypeptides can be introduced into species from such genera without the necessity of introducing mevalonate or MEP pathway genes.

*Arxuia adenivorans (Blastobotrys adenivorans)*

[00138] *Arxuia adenivorans* is dimorphic yeast (it grows as budding yeast like the baker's yeast up to a temperature of 42°C, above this threshold it grows in a filamentous form) with unusual biochemical characteristics. It can grow on a wide range of substrates and can assimilate nitrate. It has successfully been applied to the generation of strains that can produce natural plastics or the development of a biosensor for estrogens in environmental samples.

*Yarrowia lipolytica*

[00139] *Yarrowia lipolytica* is dimorphic yeast (see *Arxuia adenivorans*) and belongs to the family Hemiascomycetes. The entire genome of *Yarrowia lipolytica* is known. *Yarrowia* species is aerobic and considered to be non-pathogenic. *Yarrowia* is efficient in using hydrophobic substrates (e.g. alkanes, fatty acids, oils) and can grow on sugars. It has a high potential for industrial applications and is an oleaginous microorganism. *Yarrowia lipolytica* can accumulate lipid content to approximately 40% of its dry cell weight and is a model organism for lipid accumulation and remobilization. See e.g., Nicaud, 2012, *Yeast* 29(10):409-18; Beopoulos et al., 2009, *Biochimie* 91(6):692-6; Bankar et al., 2009, *Appl Microbiol Biotechnol.* 84(5):847-65.

*Rhodotorula* sp.

[00140] *Rhodotorula* is unicellular, pigmented yeast. The oleaginous red yeast, *Rhodotorula glutinis*, has been shown to produce lipids and carotenoids from crude glycerol (Saenge et al., 2011, *Process Biochemistry* 46(1):210-8). *Rhodotorula toruloides* strains have been shown to be an efficient fed-batch fermentation system for improved biomass and lipid productivity (Li et al., 2007, *Enzyme and Microbial Technology* 41:312-7).

*Rhodospordium toruioides*

[00141] *Rhodospordium toruioides* is oleaginous yeast and useful for engineering lipid-production pathways (See e.g. Zhu *et al.*, 2013, *Nature Commun.* 3:1 112; Ageitos *et al.*, 2011, *Applied Microbiology and Biotechnology* 90(4):1219-27).

*Candida boidinii*

[00142] *Candida boidinii* is methylotrophic yeast (it can grow on methanol). Like other methylotrophic species such as *Hansenula polymorpha* and *Pichia pastoris*, it provides an excellent platform for producing heterologous proteins. Yields in a multigram range of a secreted foreign protein have been reported. A computational method, IPRO, recently predicted mutations that experimentally switched the cofactor specificity of *Candida boidinii* xylose reductase from NADPH to NADH. See, e.g., Mattanovich *et al.*, 2012, *Methods Mol Biol.* 824:329-58; Khoury *et al.*, 2009, *Protein Sci.* 18(10):2125-38.

*Hansenula polymorpha* (*Pichia anousta*)

[00143] *Hansenula polymorpha* is methylotrophic yeast (see *Candida boidinii*). It can furthermore grow on a wide range of other substrates; it is thermo-tolerant and can assimilate nitrate (see also *Kluyveromyces lactis*). It has been applied to producing hepatitis B vaccines, insulin and interferon alpha-2a for the treatment of hepatitis C, furthermore to a range of technical enzymes. See, e.g., Xu *et al.*, 2014, *Virology* 509(2):403-9.

*Kluyveromyces lactis*

[00144] *Kluyveromyces lactis* is yeast regularly applied to the production of kefir. It can grow on several sugars, most importantly on lactose which is present in milk and whey. It has successfully been applied among others for producing chymosin (an enzyme that is usually present in the stomach of calves) for producing cheese. Production takes place in fermenters on a 40,000 L scale. See, e.g., van Ooyen *et al.*, 2006, *FEMS Yeast Res.* 6(3):381-92.

*Pichia pastoris*

[00145] *Pichia pastoris* is methylotrophic yeast (see *Candida boidinii* and *Hansenula polymorpha*). It provides an efficient platform for producing foreign proteins. Platform elements are available as a kit and it is worldwide used in academia for producing proteins. Strains have been engineered that can produce complex human N-glycan (yeast glycans are similar but not identical to those found in humans). See, e.g., Piirainen *et al.*, 2014, *N Biotechnol.* 31(6):532-7.

*Physcomitrella* spp.

[00146] *Physcomitrella* mosses, when grown in suspension culture, have characteristics similar to yeast or other fungal cultures. This genera can be used for producing plant secondary metabolites, which can be difficult to produce in other types of cells.

### **Steviol Glycoside Compositions**

[00147] Steviol glycosides do not necessarily have equivalent performance in different food systems. It is therefore desirable to have the ability to direct the synthesis to steviol glycoside compositions of choice. Recombinant hosts described herein can produce compositions that are selectively enriched for specific steviol glycosides (e.g., RebD or RebM) and have a consistent taste profile. As used herein, the term "enriched" is used to describe a steviol glycoside composition with an increased proportion of a particular steviol glycoside, compared to a steviol glycoside composition (extract) from a stevia plant. Thus, the recombinant hosts described herein can facilitate the production of compositions that are tailored to meet the sweetening profile desired for a given food product and that have a proportion of each steviol glycoside that is consistent from batch to batch. In some embodiments, hosts described herein do not produce or produce a reduced amount of undesired plant by-products found in *Stevia* extracts. Thus, steviol glycoside compositions produced by the recombinant hosts described herein are distinguishable from compositions derived from *Stevia* plants.

[00148] The amount of an individual steviol glycoside (e.g., RebA, RebB, RebD, or RebM) accumulated can be from about 1 to about 7,000 mg/L, e.g., about 1 to about 10 mg/L, about 3 to about 10 mg/L, about 5 to about 20 mg/L, about 10 to about 50 mg/L, about 10 to about 100 mg/L, about 25 to about 500 mg/L, about 100 to about 1,500 mg/L, or about 200 to about 1,000 mg/L, at least about 1,000 mg/L, at least about 1,200 mg/L, at least about at least 1,400 mg/L, at least about 1,600 mg/L, at least about 1,800 mg/L, at least about 2,800 mg/L, or at least about 7,000 mg/L. In some aspects, the amount of an individual steviol glycoside can exceed 7,000 mg/L. The amount of a combination of steviol glycosides (e.g., RebA, RebB, RebD, or RebM) accumulated can be from about 1 mg/L to about 7,000 mg/L, e.g., about 200 to about 1,500, at least about 2,000 mg/L, at least about 3,000 mg/L, at least about 4,000 mg/L, at least about 5,000 mg/L, at least about 6,000 mg/L, or at least about 7,000 mg/L. In some aspects, the amount of a combination of steviol glycosides can exceed 7,000 mg/L. In general, longer culture times will lead to greater amounts of product. Thus, the recombinant microorganism can be cultured for from 1 day to 7 days, from 1 day to 5 days, from 3 days to 5 days, about 3 days, about 4 days, or about 5 days.

[00149] It will be appreciated that the various genes and modules discussed herein can be present in two or more recombinant microorganisms rather than a single microorganism. When a plurality of recombinant microorganisms is used, they can be grown in a mixed culture to produce steviol and/or steviol glycosides. For example, a first microorganism can comprise one or more biosynthesis genes for producing a steviol glycoside precursor, while a second microorganism comprises steviol glycoside biosynthesis genes. The product produced by the second, or final microorganism is then recovered. It will also be appreciated that in some embodiments, a recombinant microorganism is grown using nutrient sources other than a culture medium and utilizing a system other than a fermenter.

[00150] Alternatively, the two or more microorganisms each can be grown in a separate culture medium and the product of the first culture medium, e.g., steviol, can be introduced into second culture medium to be converted into a subsequent intermediate, or into an end product such as RebA. The product produced by the second, or final microorganism is then recovered. It will also be appreciated that in some embodiments, a recombinant microorganism is grown using nutrient sources other than a culture medium and utilizing a system other than a fermenter.

[00151] Steviol glycosides and compositions obtained by the methods disclosed herein can be used to make food products, dietary supplements and sweetener compositions. See, e.g., WO 2011/153378, WO 2013/022989, WO 2014/122227, and WO 2014/122328.

[00152] For example, substantially pure steviol or steviol glycoside such as RebM or RebD can be included in food products such as ice cream, carbonated beverages, fruit juices, yogurts, baked goods, chewing gums, hard and soft candies, and sauces. Substantially pure steviol or steviol glycoside can also be included in non-food products such as pharmaceutical products, medicinal products, dietary supplements and nutritional supplements. Substantially pure steviol or steviol glycosides may also be included in animal feed products for both the agriculture industry and the companion animal industry. Alternatively, a mixture of steviol and/or steviol glycosides can be made by culturing recombinant microorganisms separately, each producing a specific steviol or steviol glycoside, recovering the steviol or steviol glycoside in substantially pure form from each microorganism and then combining the compounds to obtain a mixture comprising each compound in the desired proportion. The recombinant microorganisms described herein permit more precise and consistent mixtures to be obtained compared to current Stevia products.



**[00153]** In another alternative, a substantially pure steviol or steviol glycoside can be incorporated into a food product along with other sweeteners, e.g. saccharin, dextrose, sucrose, fructose, erythritol, aspartame, sucralose, monatin, or acesulfame potassium. The weight ratio of steviol or steviol glycoside relative to other sweeteners can be varied as desired to achieve a satisfactory taste in the final food product. See, e.g., U.S. 2007/0128311. In some embodiments, the steviol or steviol glycoside may be provided with a flavor (e.g., citrus) as a flavor modulator.

**[00154]** Compositions produced by a recombinant microorganism described herein can be incorporated into food products. For example, a steviol glycoside composition produced by a recombinant microorganism can be incorporated into a food product in an amount ranging from about 20 mg steviol glycoside/kg food product to about 1800 mg steviol glycoside/kg food product on a dry weight basis, depending on the type of steviol glycoside and food product. For example, a steviol glycoside composition produced by a recombinant microorganism can be incorporated into a dessert, cold confectionary (e.g., ice cream), dairy product (e.g., yogurt), or beverage (e.g., a carbonated beverage) such that the food product has a maximum of 500 mg steviol glycoside/kg food on a dry weight basis. A steviol glycoside composition produced by a recombinant microorganism can be incorporated into a baked good (e.g., a biscuit) such that the food product has a maximum of 300 mg steviol glycoside/kg food on a dry weight basis. A steviol glycoside composition produced by a recombinant microorganism can be incorporated into a sauce (e.g., chocolate syrup) or vegetable product (e.g., pickles) such that the food product has a maximum of 1000 mg steviol glycoside/kg food on a dry weight basis. A steviol glycoside composition produced by a recombinant microorganism can be incorporated into a bread such that the food product has a maximum of 160 mg steviol glycoside/kg food on a dry weight basis. A steviol glycoside composition produced by a recombinant microorganism, plant, or plant cell can be incorporated into a hard or soft candy such that the food product has a maximum of 1600 mg steviol glycoside/kg food on a dry weight basis. A steviol glycoside composition produced by a recombinant microorganism, plant, or plant cell can be incorporated into a processed fruit product (e.g., fruit juices, fruit filling, jams, and jellies) such that the food product has a maximum of 1000 mg steviol glycoside/kg food on a dry weight basis. In some embodiments, a steviol glycoside composition produced herein is a component of a pharmaceutical composition. See, e.g., Steviol Glycosides Chemical and Technical Assessment 69th JECFA, 2007, prepared by Harriet Wailin, Food Agric. Org.; EFSA Panel on Food Additives and Nutrient Sources added to Food (ANS), "Scientific Opinion on the safety of steviol glycosides for the proposed uses as a food additive," 2010, *EFSA Journal* 8(4):1537;

U.S. Food and Drug Administration GRAS Notice 323; U.S Food and Drug Administration GRAS Notice Notice 329; WO 201 1/037959; WO 2010/146463; WO 201 1/046423; and WO 2011/056834.

[00155] For example, such a steviol glycoside composition can have from 90-99 weight % RebA and an undetectable amount of stevia plant-derived contaminants, and be incorporated into a food product at from 25-1600 mg/kg, *e.g.*, 100-500 mg/kg, 25-100 mg/kg, 250-1000 mg/kg, 50-500 mg/kg or 500-1 000 mg/kg on a dry weight basis.

[00156] Such a steviol glycoside composition can be a RebB-enriched composition having greater than 3 weight % RebB and be incorporated into the food product such that the amount of RebB in the product is from 25-1600 mg/kg, *e.g.*, 100-500 mg/kg, 25-100 mg/kg, 250-1000 mg/kg, 50-500 mg/kg or 500-1 000 mg/kg on a dry weight basis. Typically, the RebB-enriched composition has an undetectable amount of stevia plant-derived contaminants.

[00157] Such a steviol glycoside composition can be a RebD-enriched composition having greater than 3 weight % RebD and be incorporated into the food product such that the amount of RebD in the product is from 25-1600 mg/kg, *e.g.*, 100-500 mg/kg, 25-100 mg/kg, 250-1000 mg/kg, 50-500 mg/kg or 500-1000 mg/kg on a dry weight basis. Typically, the RebD-enriched composition has an undetectable amount of stevia plant-derived contaminants.

[00158] Such a steviol glycoside composition can be a RebE-enriched composition having greater than 3 weight % RebE and be incorporated into the food product such that the amount of RebE in the product is from 25-1600 mg/kg, *e.g.*, 100-500 mg/kg, 25-100 mg/kg, 250-1000 mg/kg, 50-500 mg/kg or 500-1000 mg/kg on a dry weight basis. Typically, the RebE-enriched composition has an undetectable amount of stevia plant-derived contaminants.

[00159] Such a steviol glycoside composition can be a RebM-enriched composition having greater than 3 weight % RebM and be incorporated into the food product such that the amount of RebM in the product is from 25-1600 mg/kg, *e.g.*, 100-500 mg/kg, 25-100 mg/kg, 250-1000 mg/kg, 50-500 mg/kg or 500-1000 mg/kg on a dry weight basis. Typically, the RebM-enriched composition has an undetectable amount of stevia plant-derived contaminants.

[00160] In some embodiments, a substantially pure steviol or steviol glycoside is incorporated into a tabletop sweetener or "cup-for-cup" product. Such products typically are diluted to the appropriate sweetness level with one or more bulking agents, *e.g.*, maltodextrins, known to those skilled in the art. Steviol glycoside compositions enriched for RebA, RebB, RebD, RebE, or RebM, can be package in a sachet, for example, at from 10,000 to 30,000 mg

steviol glycoside/kg product on a dry weight basis, for tabletop use. In some embodiments, a steviol glycoside produced *in vitro*, *in vivo*, or by whole cell byconversion

**[00161]** The invention will be further described in the following examples, which do not limit the scope of the invention described in the claims.

### EXAMPLES

**[00162]** The Examples that follow are illustrative of specific embodiments of the invention, and various uses thereof. They are set forth for explanatory purposes only, and are not to be taken as limiting the invention.

#### Example 1. LC-MS Analytical Procedures

**[00163]** Three LC-MS procedures were used herein. In the first method used for Examples 2-6, LC-MS analyses were performed using an Ultimate 3000 UPLC system (Dionex) fitted with a Waters Acquity UPLC @BEH shield RP18 column (2.1 x 50 mm, 1.7  $\mu$ m particles, 130 Å pore size) connected to a TSQ Quantum Access (ThermoFisher Scientific) triple quadrupole mass spectrometer with a heated electrospray ion (HESI) source. Elution was carried out using a mobile phase of eluent B (MeCN with 0.1% formic acid) and eluent A (water with 0.1% formic acid) by increasing the gradient from 25% to 47% B from min 0.0 to 4.0, increasing 47% to 100% B from min 4.0 to 5.0, and holding 100% B from min 5.0 to 6.5. The flow rate was 0.4 mL/min and the column temperature 35°C. Steviol glycosides were detected using SIM (Single Ion Monitoring) with the following m/z-traces.

**Table 1A: LC-MS analytical information for Steviol Glycosides.**

Description	Exact Mass	m/z trace (Da)	compound (typical $t_R$ in min)
Steviol + 1 Glucose	[M+H] <sup>+</sup> 481.2796 [M+Na] <sup>+</sup> 503.2615	481.2± 0.5 503.1± 0.5	19-SMG (2.29), 13-SMG (3.5)
Steviol + 2 Glucose	[M+Na] <sup>+</sup> 665.3149	665± 0.5	Rubusoside (2.52) Steviol-1,2-bioside (2.92) Steviol-1,3-bioside (2.28)
Steviol + 3 Glucose	[M+Na] <sup>+</sup> 827.3677	827.4 ± 0.5	1,2-Stevioside (2.01) 1,3-Stevioside (2.39) Rebaudioside B (2.88)
Steviol + 4 Glucose	[M+Na] <sup>+</sup> 989.4200	989.4 ± 0.5	Rebaudioside A (2.0)
Steviol + 5 Glucose	[M+Na] <sup>+</sup> 1151.4728	1151.4 ± 0.5	Rebaudioside D (1.1)
Steviol +	[M+Na] <sup>+</sup> 1313.5257	1313.5 ± 0.5	Rebaudioside M (1.3)

Description	Exact Mass	m/z trace (Da)	compound (typical $t_R$ in min)
6 Glucose			

[00164] in the second method used for Examples 7, 8, and 10, LC-MS analyses were performed on Waters ACQUITY UPLC (Waters Corporation, Milford, MA) with coupled to a Waters ACQUITY ESI (electrospray ionization)-TQD triple quadrupole mass spectrometer. Compound separation was achieved on Waters ACQUITY UPLC® BEH C18 column (2.1 x 50 mm, 1.7  $\mu$ m particles, 130 Å pore size) equipped with ACQUITY UPLC BEH C18 VanGuard pre-column (130 Å, 1.7  $\mu$ m, 2.1 mm X 5 mm) by using a gradient of the two mobile phases: A (Water with 0.1% formic acid) and B (Acetonitrile with 0.1% formic acid) increasing B from 20% to 50% between 0.3 to 2.0 min up to 100% at 2.01 min, holding to 100% for 0.6 min, and re-equilibrating for 0.6 min. The flow rate was 0.6 mL/min, and the column temperature was 55°C. The MS acquisition was in negative ion-mode using SIM mode (Single Ion Monitoring). Steviol glycoside quantification was done by comparison with authentic standards.

Table 1B: MS analytical information for Steviol Glycosides.

Compound	m/z trace (Da)	Retention time (min)
RebE	965.42	1.06
RebD	1127.48	1.09
RebM	1289.53	1.15
RebA	965.42	1.43
1,3-Stevioside	803.37	1.60
Rubusoside	641.32	1.67
RebB	803.37	1.76
1,2-bioside	641.32	1.77
13-SMG	479.26	2.04

[00165] in the third method used for Example 9, LC-MS analyses were performed on Waters ACQUITY UPLC (Waters Corporation, Milford, MA) using a Waters Acquity UPLC® BEH C18 column (2.1 x 50 mm, 1.7  $\mu$ m particles, 130 Å) coupled to a Waters single quadrupole mass spectrometer (SQD), equipped with an ESI and operated in negative mode. Compound separation was achieved by a gradient of the two mobile phases: A (water with 0.1% formic acid) and B (acetonitrile with 0.1% formic acid) by increasing from 60% to 100% B between 0.3 to 2.5 min, holding 100% B for 0.1 min, and re-equilibrating for 0.2 min. The flow rate was 0.6 mL/min, and the column temperature was set at 55°C. Steviol or ent-kaurenoic acid was

monitored using SIM (Single Ion Monitoring) and quantified by comparing with authentic standards.

**Table 1C: MS analytical information for steviol and ent-kaurenoic acid.**

Compound	m/z trace (Da)	Retention time (min)
Steviol	317.21	0.61
Ent-kaurenoic acid	301.001	1.46

**Example 2. Construction of Steviol Glycoside-Producing and RebB-Producing Yeast Strains**

**[00166]** Steviol glycoside-producing *S. cerevisiae* strains were constructed as described in WO 201 1/153378, WO 2013/022989, WO 2014/122227, and WO 2014/122328. For example, a yeast strain comprising a recombinant gene encoding a *Synechococcus sp.* GGPPS (SEQ ID NO:49) polypeptide, a recombinant gene encoding a truncated *Zea mays* CDPS (SEQ ID NO:37) polypeptide, a recombinant gene encoding an *A. thaliana* KS (SEQ ID NO:6) polypeptide, a recombinant gene encoding an *S. rebaudiana* KO (SEQ ID NO:59, SEQ ID NO:79) polypeptide, a recombinant gene encoding an *A. thaliana* ATR2 (SEQ ID NO:51, SEQ ID NO:87) polypeptide, a recombinant gene encoding an *O. sativa* EUGT1 1 (SEQ ID NO:86) polypeptide, a recombinant gene encoding an SrKAHel (SEQ ID NO:18, SEQ ID NO:68) polypeptide, a recombinant gene encoding an *S. rebaudiana* CPR8 (SEQ ID NO:24, SEQ ID NO:28) polypeptide, a recombinant gene encoding an *S. rebaudiana* UGT85C2 (SEQ ID NO:30) polypeptide, a recombinant gene encoding an *S. rebaudiana* UGT74G1 (SEQ ID NO:29) polypeptide, a recombinant gene encoding an *S. rebaudiana* UGT76G1 (SEQ ID NO:2) polypeptide, and a recombinant gene encoding an *S. rebaudiana* UGT91D2 variant, UGT91D2e-b (SEQ ID NO:88), polypeptide accumulated steviol glycosides.

**[00167]** The UGT91D2e-b variant of UGT91D2 (SEQ ID NO:5 from PCT/US201 2/050021) includes a substitution of a methionine for leucine at position 211 and a substitution of an alanine for valine at position 286. Additional variants can include variants (except T144S, M152L, L213F, S364P, and G384C variants) described in Table 14 and Example 11 of the PCT/US201 2/050021. GeneArt codon-optimized sequence encoding a *S. rebaudiana* UGT91D2e-b with the amino acid modifications L21 1M and V286A (SEQ ID NO:88 for amino acid sequence; codon optimized nucleotide sequence is set forth in SEQ ID NO:89) and

expressed from the native yeast TDH3 promoter and followed by the native yeast CYC1 terminator.

**[00168]** Cells were grown in Synthetic Complete (SC) medium at 30°C for 5 days with shaking (400 rpm for deep wells and 200 rpm for 15 ml\_ Falcon growth tubes) prior to harvest. Culture samples (without cell removal) were heated in the presence of DMSO for detection of total glycoside levels with LC-MS. The strain accumulated total amounts of RebD of over 2500 mg/L, total amounts of RebM of over 2500 mg/L, and total amounts of RebA of over 700 mg/L. See WO 2014/122227.

**[00169]** A separate *S. cerevisiae* strain was constructed to accumulate RebB. This strain comprised a recombinant gene encoding a *Synechococcus sp.* GGPPS (SEQ ID NO:49) polypeptide, a recombinant gene encoding a truncated *Z. mays* CDPS (SEQ ID NO:37) polypeptide, a recombinant gene encoding an *A. thaliana* KS (SEQ ID NO:6) polypeptide, a recombinant gene encoding an *S. rebaudiana* KO (SEQ ID NO:59, SEQ ID NO:79) polypeptide, a recombinant gene encoding an *A. thaliana* ATR2 (SEQ ID NO:51, SEQ ID NO:87) polypeptide, a recombinant gene encoding an *O. sativa* EUGT1 1 (SEQ ID NO:86) polypeptide, a recombinant gene encoding an SrKAHei (SEQ ID NO:18, SEQ ID NO:68) polypeptide, a recombinant gene encoding an *S. rebaudiana* CPR8 (SEQ ID NO:24, SEQ ID NO:28) polypeptide, a recombinant gene encoding an *S. rebaudiana* UGT85C2 (SEQ ID NO:30) polypeptide, a recombinant gene encoding an *S. rebaudiana* UGT76G1 (SEQ ID NO:2) polypeptide, and a recombinant gene encoding an *S. rebaudiana* UGT91D2 variant, UGT91D2e-b (SEQ ID NO:88), polypeptide accumulated steviol glycosides.

### **Example 3. Steviol Glycoside Production in Yeast Strains Expressing KO Genes**

**[00170]** To determine whether increased levels of ent-kaurenoic acid improve steviol glycoside production, the activity of KO genes from various species were analyzed. Putative KO genes were identified using the NCBI Basic Local Alignment Sequence Search Tool (BLAST). Genes encoding KO polypeptides were cloned and expressed the RebB-producing *S. cerevisiae* strain described in Example 2, which was modified to lack KO genes. Thus, RebB was only accumulated upon expression of a functional KO.

**[00171]** Two KO polypeptides identified by the amino acid sequences set forth in SEQ ID NO:54 (nucleotide sequence set forth in SEQ ID NO:55) and SEQ ID NO:75 (nucleotide sequences set forth in SEQ ID NO:56) were found to accumulate higher levels of RebB than

SrKOl (nucleotide sequence set forth in SEQ ID NO:59, amino acid sequences set forth in SEQ ID NO:79) in the RebB-producing strain. RebB levels ( $\mu\text{M}/00_{60}$ ) are shown in Figure 3.

**[00172]** Expression of genes (SEQ ID NO:55 or SEQ ID NO:56) encoding KO polypeptides in an *S. cerevisiae* steviol glycoside-producing strain also resulted in accumulation of ent-kaurenoic acid (Figure 4). Expression of a gene encoding a codon-optimized KO polypeptide (SEQ ID NO:57) and a gene encoding the KO polypeptide set forth in SEQ ID NO:70 also resulted in accumulation of ent-kaurenoic acid. However, expression of SrKOl (SEQ ID NO:59, SEQ ID NO:79) did not result in measurable levels of ent-kaurenoic acid. Thus, the KO polypeptides encoded by nucleotide sequences set forth in SEQ ID NOs: 55-57 more efficiently converted ent-kaurene, ent-kaurenol, and/or ent-kaurenal to ent-kaurenoic acid in *S. cerevisiae*, as compared to the SrKOl polypeptide encoded by nucleotide sequence set forth in SEQ ID NO:59.

**Example 4. Steviol Glycoside Production in Yeast Strains Expressing KO Genes and Further Overexpressing SrKAHel**

**[00173]** Cloned KO genes were individually expressed in a steviol glycoside-producing *S. cerevisiae* strain. The *S. cerevisiae* strain described in Example 2, which expresses SrKOl (SEQ ID NO:59, SEQ ID NO:79), was modified to comprise overexpress SrKAHel (SEQ ID NO:18, SEQ ID NO:68). The coding sequences of the KO genes tested, as well as their corresponding amino acid sequences, are set forth in Table 2. The sequences set forth in SEQ ID NOs: 55, 57, 58, 59, and 60 were codon-optimized for expression in *S. cerevisiae*.

**Table 2: KO Genes Expressed in Steviol Glycoside-Producing *S. cerevisiae* strain that Further Overexpresses SrKAHel.**

KO Nucleotide Sequence	Corresponding KO Amino Acid Sequence
SEQ ID NO:55	SEQ ID NO:54
SEQ ID NO:56	SEQ ID NO:75
SEQ ID NO:57	SEQ ID NO:70
SEQ ID NO:58	SEQ ID NO:71
SEQ ID NO:59	SEQ ID NO:79
SEQ ID NO:60	SEQ ID NO:72

**[00174]** *S. cerevisiae* strains co-expressing any of the heterologous nucleic acids encoding a KO enzyme of Table 2 and further overexpressing SrKAHel (SEQ ID NO:18, SEQ ID NO:68)

accumulated higher levels of steviol glycosides than the control *S. cerevisiae* strain (not expressing a KO of Table 2) or a steviol glycoside-producing *S. cerevisiae* strain only overexpressing SrKAHel , as shown in Figure 5. A steviol glycoside-producing *S. cerevisiae* strain expressing a codon-optimized version of SEQ ID NO:56, identified herein as SEQ ID NO:65, and overexpressing SrKAHel accumulated higher levels of steviol glycosides (RebA, RebD, and RebM) than the steviol glycoside-producing *S. cerevisiae* strain co-expressing the nucleic acid set forth in SEQ ID NO:56 and SrKAHel (Figure 6).

**[00175]** Additionally, *S. cerevisiae* strains co-expressing a nucleic acid set forth in SEQ ID NO:55, SEQ ID NO:56, SEQ ID NO:57, or SEQ ID NO:60 and further overexpressing SrKAHel accumulated higher levels of glycosylated ent-kaurenoic acid than the control *S. cerevisiae* strain not expressing a KO of Table 2 (Figure 7).

**[00176]** As well, *S. cerevisiae* strains co-expressing a nucleic acid set forth in SEQ ID NO:55, SEQ ID NO:56, SEQ ID NO:57, SEQ ID NO:59, or SEQ ID NO:60 and further overexpressing SrKAHel demonstrated improved metabolic conversion of intermediate compound, ent-kaurenol, which, in turn, resulted in reduced accumulation of glycosylated ent-kaurenol, relative to the control *S. cerevisiae* strain not expressing a KO of Table 2 or the steviol glycoside-producing *S. cerevisiae* strain only overexpressing SrKAHel, as shown in Figure 8. The control *S. cerevisiae* strain and the steviol glycoside-producing *S. cerevisiae* strain only overexpressing SrKAHel each accumulated higher levels of glycosylated ent-kaurenol than did *S. cerevisiae* strains expressing a nucleic acid set forth in SEQ ID NO:55, SEQ ID NO:56, SEQ ID NO:57, SEQ ID NO:59, or SEQ ID NO:60 and further overexpressing SrKAHel.

**Example 5. Steviol Glycoside Production in Yeast Strains Expressing CPR Genes**

**[00177]** Cloned CPR genes were individually expressed in a steviol glycoside-producing *S. cerevisiae* strain. The steviol glycoside-producing *S. cerevisiae* strain described in Example 2, which expresses *S. rebaudiana* CPR8 (SEQ ID NO:24, SEQ ID NO:28) and *A. thaliana* ATR2 (SEQ ID NO:51), was modified to co-express a nucleic acid encoding a CPR of Table 3. The coding sequences of the CPR genes tested, as well as their corresponding amino acid sequences, are set forth in Table 3.

**Table 3: CPR Genes Tested in Combination with CPR8 and ATR2.**

Gene	Nucleotide Sequence	Amino Acid Sequence
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<i>S. rebaudiana</i> CPR1	SEQ ID NO:61	SEQ ID NO:76
<i>S. rebaudiana</i> CPR7	SEQ ID NO:23	SEQ ID NO:69
CPR4497	SEQ ID NO:62	SEQ ID NO:74

[00178] As shown in Figure 9, expression of CPR1 (SEQ ID NO:61, SEQ ID NO:76) or of CPR7 (SEQ ID NO:23, SEQ ID NO:69) in the steviol glycoside-producing *S. cerevisiae* strain already expressing *S. rebaudiana* CPR8 (SEQ ID NO:24, SEQ ID NO:28) and *A. thaliana* ATR2 (SEQ ID NO:51) resulted in higher levels of RebM than those accumulated by the control steviol glycoside-producing *S. cerevisiae* strain not expressing CPR1 or CPR7. As well, a steviol glycoside-producing *S. cerevisiae* strain expressing the nucleic acid set forth in SEQ ID NO:62 and overexpressing SrKAHel (SEQ ID NO:18, SEQ ID NO:68) accumulated higher levels of RebM than those accumulated by the control steviol glycoside-producing *S. cerevisiae* strain that only overexpressed SrKAHel (Figure 10).

**Example 6. Steviol Glycoside Production in Yeast Strains Co-Expressing KO and CPR Genes**

[00179] Steviol glycoside production was tested in the RebB-producing *S. cerevisiae* strain described in Example 2, which was modified to co-express a KO gene of Table 4 and a CPR of Table 5.

Table 4: KO Genes Tested in Combination with CPR Genes.

Gene	Nucleotide Sequence	Amino Acid Sequence
SrKO1	SEQ ID NO:59	SEQ ID NO:79
Codon-optimized KO	SEQ ID NO:63	SEQ ID NO:77
Codon-optimized KO	SEQ ID NO:64	SEQ ID NO:78

Table 5: CPR Genes Tested in Combination with KO Genes.

Nucleotide Sequence	Amino Acid Sequence
SEQ ID NO:66	SEQ ID NO:73
SEQ ID NO:67	SEQ ID NO:22

[00180] As shown in Figure 12, co-expression of SrKO1 (SEQ ID NO:59, SEQ ID NO:79) and either of the CPR genes of Table 5 in the RebB-producing strain resulted in higher production of 13-SMG and RebB than co-expression of a nucleic acid set forth in SEQ ID NO:63 or SEQ ID NO:64 and either of the cytochrome P450 genes of Table 5.

**Example 7. Steviol Glycoside Production in Yeast Strains Expressing KAH Genes**

**[00181]** Candidate KAH enzymes were cloned and expressed in an *S. cerevisiae* strain engineered to accumulate 13-SMG. The 13-SMG-producing *S. cerevisiae* strain comprised a recombinant gene encoding a *Synechococcus sp.* GGPPS7 polypeptide (SEQ ID NO:49), a recombinant gene encoding a truncated *Z. mays* CDPS polypeptide (SEQ ID NO:37), a recombinant gene encoding an *A. thaliana* KS polypeptide (SEQ ID NO:6), SrKOH (SEQ ID NO:59, SEQ ID NO:79), CPR8 (SEQ ID NO:24, SEQ ID NO:28), the KO encoded by the nucleotide sequence set forth in SEQ ID NO:56 (amino acid sequence set forth in SEQ ID NO:75), and UGT85C2 (SEQ ID NO:30) chromosomally integrated in separate expression cassettes (Figure 11B). The strain lacked SrKAHel (SEQ ID NO:18, SEQ ID NO:68); thus, 13-SMG was only accumulated upon transformation of the *S. cerevisiae* strain with a functional KAH (Figure 11B).

**[00182]** Transformants were grown in SC-URA medium for 4 days and extracted with 1:1 with DMSO at 80°C for 10 min. The extracts were analyzed by LC-MS (method 2 of Example 1). *S. cerevisiae* transformed with the nucleic acid set forth in SEQ ID NO:80 accumulated 13-SMG (Figure 11B). Thus, the protein encoded by SEQ ID NO:80, set forth in SEQ ID NO:82, is a KAH.

**[00183]** The KAH encoded by the nucleotide sequence set forth in SEQ ID NO:80 was codon-optimized for expression in yeast (SEQ ID NO:81) and expressed in the above-described 13-SMG-producing *S. cerevisiae* strain. Similar to expression of SrKAHel (SEQ ID NO:18) or the KAH encoded by the nucleotide sequence set forth in SEQ ID NO:80, expression of the codon-optimized nucleotide sequence set forth in SEQ ID NO:81 resulted in production of 13-SMG plus rubusoside (Figure 13).

**[00184]** The KAHs encoded by the nucleotide sequence set forth in SEQ ID NO:80 and the codon-optimized nucleotide sequence set forth in SEQ ID NO:81 were also individually expressed in a steviol glycoside-producing strain, as described in Example 2, which expresses SrKAHel. Production of 13-SMG was increased upon overexpression of SrKAHel (SEQ ID NO: 18), of the KAH encoded by the nucleotide sequence set forth in SEQ ID NO:80, or of the KAH encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:81, as compared to a control strain not expressing the KAH encoded by the nucleotide sequence set forth in SEQ ID NO:80, the KAH encoded by the codon-optimized nucleotide sequence set forth

in SEQ ID NO:81, or overexpressing SrKAHe1. See Table 6. Expression of either the KAH encoded by the nucleotide sequence set forth in SEQ ID NO:80 or the KAH encoded by the codon-optimized nucleotide sequence set forth in SEQ ID NO:81 resulted in higher steviol glycoside production (13-SMG + 1,2-bioside + rubusoside + RebB + RebA + RebD + RebM) than either the control strain or the *S. cerevisiae* strain overexpressing SrKAHe1 (SEQ ID NO: 18). See Table 6.

**Table 6: Quantification of Steviol Glycosides Accumulated by Yeast Expressing KAH Genes.**

	Control ( $\mu\text{M}$ )	Overexpression of SrKAHe1 (encoded by the nucleotide set forth in SEQ ID NO:18) ( $\mu\text{M}$ )	SrKAHe1 + KAH (encoded by the nucleotide set forth in SEQ ID NO:80) ( $\mu\text{M}$ )	SrKAHe1 + KAH (encoded by the nucleotide sequence set forth in SEQ ID NO:81) ( $\mu\text{M}$ )
<b>13-SMG</b>	67.6	85.5	153.8	130.5
<b>Steviol-1,2-bioside</b>	0.4	0.3	0.4	0.4
<b>Rubusoside</b>	1.2	1.0	1.4	1.1
<b>RebB</b>	8.6	7.6	9.6	9.6
<b>RebA</b>	30.7	26.0	26.8	28.7
<b>RebD</b>	36.2	27.6	32.9	36.5
<b>RebM</b>	138.3	118.9	100.0	90.3
<b>Sum</b>	282.7	266.2	324.0	296.7

**Example 8. Steviol Glycoside Production in Yeast Strain Expressing KAH Gene of the CYP72A219 family**

[001 85] A nucleic acid of SEQ ID NO:90, which was codon-optimized for expression in *S. cerevisiae* and encodes the polypeptide of SEQ ID NO:91, was cloned and expressed in an *S. cerevisiae* strain described in Example 7, which was engineered to accumulate 13-SMG. The 13-SMG-producing *S. cerevisiae* strain comprised a recombinant gene encoding a *Synechococcus sp.* GGPPS7 polypeptide (SEQ ID NO:49), a recombinant gene encoding a truncated *Z. mays* CDPS polypeptide (SEQ ID NO:37), a recombinant gene encoding an *A. thaliana* KS polypeptide (SEQ ID NO:6), SrKOI (SEQ ID NO:59, SEQ ID NO:79), CPR8 (SEQ ID NO:24, SEQ ID NO:28), the KO encoded by the nucleotide sequence set forth in SEQ ID NO:56 (amino acid sequence set forth in SEQ ID NO:75), and UGT85C2 (SEQ ID NO:30) chromosomally integrated in separate expression cassettes.

**[00186]** Transformants were grown in SC-URA medium for 4 days and extracted 1:1 with DMSO at 80°C for 10 min. The extracts were analyzed by LC-MS (method 2 of Example 1). *S. cerevisiae* transformed with the nucleic acid set forth in SEQ ID NO:90 accumulated 13-SMG as well as rubusoside (Table 7). Thus, the protein encoded by the nucleic acid sequence of SEQ ID NO:90, set forth in SEQ ID NO:91, is a KAH.

**Table 7: Quantification of Steviol Glycosides Accumulated by Yeast Expressing the KAH encoded by the Nucleotide Sequence Set Forth in SEQ ID NO:90 (Amino Acid Sequence Set Forth in SEQ ID NO:91).**

	13-SMG ( $\mu\text{M}$ )	Rubusoside ( $\mu\text{M}$ )
KAH (encoded by the nucleotide sequence set forth in SEQ ID NO:90)	4.3 $\pm$ 0.1	0.2 $\pm$ 0.0

**Example 9. Determination of CPR1 and CPR12 Activity**

**[00187]** Activity of CPR1 and CPR12 were measured using an *in vitro* microsomal assay. Microsomes were prepared by a modified version of the method taught by Pompon *et al.*, "Yeast expression of animal and plant P450s in optimized redox environments," *Methods Enzymol.* 272:51-64 (1996). *S. cerevisiae* cells were sedimented for 10 min at 4°C. The pellets were washed with 10 mL TEK buffer (50 mM Tris-HCl (pH 7.5), 1 mM EDTA, 100 mM KCl.) The cells were sedimented again for 10 min at 4°C, and the pellets were resuspended in 1-3 mL of TES2 buffer (50 mM Tris-HCl (pH 7.5) 1 mM EDTA, 600 mM sorbitol). Glass beads (425-600 microns) were added to the samples, and the cells were broken vigorously by shaking and vortexing for 5 min at 4°C. The supernatant was collected, and the beads were washed several times with TES2 buffer. The washes were combined with the supernatant, and the samples were centrifuged for 15 min at 4°C to remove unbroken cells and glass beads. Samples were then ultracentrifuged for 1 h at 4°C. The pellets were washed twice with TES buffer (50 mM Tris-HCl (pH 7.5), 1 mM EDTA, 600 mM sorbitol, 1% (w/V) BSA, 5 mM DTT), and once with TEG buffer (50 mM Tris-HCl (pH 7.5), 1 mM EDTA, 30% (V/V) glycerol). The samples were resuspended in 1-3 mL TEG, and the pellets were homogenized.

**[00188]** Wild-type control microsomal protein was prepared as described above from wild-type *S. cerevisiae* cells that did not comprise a heterologous KAH or CPR. Microsomal protein

was also prepared from *S. cerevisiae* cells expressing i) SrKAHel (SEQ ID NO:18, SEQ ID NO:68), ii) SrKAHel (SEQ ID NO:18, SEQ ID NO:68) and CPR1 (SEQ ID NO:61, SEQ ID NO:76), or iii) SrKAHel (SEQ ID NO:18, SEQ ID NO:68) and CPR12 (SEQ ID NO:97, SEQ ID NO:98) from a genetic construct integrated at the chromosome level. Microsomal protein from a steviol glycoside-producing strain was prepared from *S. cerevisiae* cells expressing the genes described in Example 2 and additionally comprising codon-optimized CPR1 from *S. rebaudiana* (SEQ ID NO:61 corresponding to amino acid sequence SEQ ID NO:76) as well as the KO encoded by SEQ ID NO:75).

**[00189]** CPR1 and CPR12 activities were first determined using a cytochrome C reductase assay kit (Sigma-Aldrich; CY0100-1KT) to measure the ability of CPR1 or CPR12 to reduce cytochrome C in the presence of NADPH *in vitro*. Reduction of cytochrome C resulted in an increase in absorbance at 550 nm, which could be quantified spectrophotometrically. Working solution was prepared by adding 9 mg cytochrome C to 20 mL assay buffer, and solution was stored at 25°C until use. NADPH was diluted in H<sub>2</sub>O to a concentration of 0.85 mg/mL. Final reaction volumes were 1.1 mL (950 μL working solution (0.43 mg cytochrome C), 28 μL enzyme dilution buffer, 100 μL NADPH solution (0.085 mg NADPH), 20 μL cytochrome C oxidase inhibitor, 2 μL microsomal protein.) Blank samples did not comprise microsomal protein and were prepared with 950 μL working solution (0.43 mg cytochrome C), 30 μL enzyme dilution buffer, 100 μL NADPH solution (0.085 mg NADPH), and 20 μL cytochrome C oxidase inhibitor. The spectrophotometer was blanked with all components added to the reactions except for NADPH. The enzymatic reactions were initiated by addition of NADPH, the samples were thoroughly mixed by pipetting, and absorbance was measured at 550 nm for 70 s with 10 s intervals between reads. Two independent rate measurements were taken for each microsomal preparation, and rates were averaged for calculation of specific activity. After the reactions were completed, results were normalized to protein concentration, which was measured using a standard BCA assay (Thermo Scientific).

**[00190]** Units/mL was calculated using the following equation, where  $\Delta A_{550}/\text{min}$  represents the change in absorbance at 550 nm during the absorbance reading period, 1.1 represents the reaction volume in mL, and 21.1 represents the extinction coefficient for reduced cytochrome c:

$$\text{Units/mL} = (\Delta A_{550}/\text{min} \times \text{dilution factor} \times 1.1) / (21.1 \times \text{enzyme volume})$$

**[00191]** The units/mL value of each sample was divided by its respective microsomal protein concentrations to calculate CPR activity in units/mg. Figure 14 shows the activity measurements of the i) SrKAHel (SEQ ID NO:18, SEQ ID NO:68), ii) SrKAHel (SEQ ID NO:18,

SEQ ID NO:68) and CPR1 (SEQ ID NO:61, SEQ ID NO:76), and iii) SrKAHel (SEQ ID NO:18, SEQ ID NO:68) and CPR12 (SEQ ID NO:97, SEQ ID NO:98) microsomal samples.

**[00192]** The microsomal preparation from the wild-type control showed only minimal CPR activity, reflecting the low activity of native NCP1 (YHR042W). Likewise, the microsomal preparation from a yeast strain overexpressing KAHel did not demonstrate an increase in CPR activity. In contrast, microsomal preparation from strains expressing SrKAHel (SEQ ID NO:18, SEQ ID NO:68) and CPR1 (SEQ ID NO:61, SEQ ID NO:76) or SrKAHel (SEQ ID NO:18, SEQ ID NO:68) and CPR12 (SEQ ID NO:97, SEQ ID NO:98) demonstrated high CPR activity, with 7- and 14-fold higher activity, respectively, compared to the negative control (Figure 14).

**[00193]** In a separate experiment, formation of steviol and consumption of ent-kaurenoic acid in microsomes, as prepared above, were measured. 33  $\mu$ M ent-kaurenoic acid, 10 mM NADPH, and 10  $\mu$ L of microsomal protein in 50 mM phosphate buffer (pH 7.5) were incubated for 30 min at 30°C in a total reaction volume of 100  $\mu$ L. Control reactions were extracted immediately after addition of all the reaction components, which were mixed on ice and aliquoted prior to incubation. Steviol and ent-kaurenoic acid levels were quantified using the second LC-MS procedure described in Example 1. For steviol quantification, the microsomal reactions were extracted with DMSO (1:1) at 80°C for 10 min and submitted for LC-MS analysis after centrifugation. For ent-kaurenoic acid quantification the microsomes reactions were extracted with acetonitrile 1:4 (20% microsomal reaction and 80% acetonitrile) at 80°C for 10 min and after centrifugation submitted for LC-MS analysis. The AUC values obtained for the ent-kaurenoic acid measurements were converted to concentrations using a standard curve.

**[00194]** As shown in Figure 15A, microsomal protein prepared from an *S. cerevisiae* strain expressing SrKAHel (SEQ ID NO:18, SEQ ID NO:68) and either CPR1 (SEQ ID NO:61, SEQ ID NO:76) or CPR12 (SEQ ID NO:97, SEQ ID NO:98) converted ent-kaurenoic acid to steviol during the 30 minute incubation period. The steviol level shown in Figure 15A for the steviol-glycoside-producing strain control (extracted immediately with no 30 min incubation period) corresponds to steviol that was accumulated by the strain prior to microsomal preparation and that had co-purified with the microsomes. As shown in Figure 15B, ent-kaurenoic acid levels decreased upon incubation with microsomal protein prepared from *S. cerevisiae* strains expressing SrKAHel (SEQ ID NO:18, SEQ ID NO:68) alone or in combination with CPR1 (SEQ ID NO:61, SEQ ID NO:76) or CPR12 (SEQ ID NO:97, SEQ ID NO:98). The increased ent-kaurenoic acid levels shown in Figure 15B for the steviol glycoside-producing strain microsomal sample incubated for 30 min corresponds to ent-kaurenoic acid that was accumulated by the

strain prior to microsomal preparation and to ent-kaurenoic acid accumulated from ent-kaurene that had co-purified with the microsomes. The levels of ent-kaurenoic acid shown in Figure 15B were corrected for the dilution factor used.

**Example 10. Steviol Glycoside Production in *S. cerevisiae* strains comprising Fusion Constructs between a KO and a P450 Reductase Domain**

**[00195]** CYP102A1 (also referred to as P450<sub>BM3</sub>; SEQ ID NO:115, SEQ ID NO:116) is a catalytically self-sufficient soluble enzyme from *Bacillus megatarium*. See, e.g., Whitehouse *et al*, 2012, Chem Soc Rev. 41(3):1218-60. Two domains are present in the CYP102A1 polypeptide chain: a P450 heme domain (BMP) and an NADPH-dependent P450 oxidoreductase domain (BMR). CYP102A1 utilizes nearly 100% of the reducing power of NADPH to produce a monooxygenated product. See, e.g., Yuan *et al*, 2009, *Biochemistry* 48(38):9140-6.

**[00196]** The BMR domain of CYP102A1 ("BMR"; codon-optimized nucleotide sequence set forth in SEQ ID NO:117, SEQ ID NO:118) was fused to SrKOi (SEQ ID NO:59, SEQ ID NO:79) or a KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 (amino acid sequence set forth in SEQ ID NO:75) with a linker (SEQ ID NO:121, SEQ ID NO:122), as described in Dodhia *et al*, 2006, J Biol Inorg Chem. 11(7):903-16. A wild-type version of the BMR domain of CYP102A1, as well as a W1046A mutant of the BMR domain (SEQ ID NO:119, SEQ ID NO:120), which has been found to switch the cofactor specificity of CYP102A1 from NADPH to NADH, were used. See, Girvan *et al.*, 2011, Arch Biochem Biophys. 507(1):75-85. SrKOi (SEQ ID NO:59, SEQ ID NO:79) and the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 were also truncated prior to fusion with the BMR domain of CYP102A1; these truncations were predicted by bioinformatics to result in loss of membrane anchors of the KO genes and in cytosolic versions of the KO-BMR fusion constructs. The KO-BMR fusion constructs analyzed are shown in Table 8.

**Table 8: KO-BMR fusion constructs and sequences.**

Fusion Construct	Codon-Optimized Nucleotide Sequence	Amino Acid Sequence
SrKO1-BMR	SEQ ID NO:99	SEQ ID NO:100
SrKO1-BMR W1046A mutant	SEQ ID NO:101	SEQ ID NO:102
Truncated SrKO1-BMR	SEQ ID NO:103	SEQ ID NO:104

Truncated SrKO1-BMR W1046A mutant	SEQ ID NO:105	SEQ ID NO:106
KO ( <i>encoded by nucleotide sequence set forth in SEQ ID NO:65</i> )-BMR	SEQ ID NO:107	SEQ ID NO:108
KO ( <i>encoded by nucleotide sequence set forth in SEQ ID NO:65</i> )-BMR W1046A mutant	SEQ ID NO:109	SEQ ID NO:110
Truncated KO ( <i>encoded by nucleotide sequence set forth in SEQ ID NO:65</i> )-BMR W1046A mutant	SEQ ID NO:111	SEQ ID NO:112

[00197] The KO-BMR fusion constructs were cloned and transformed in the RebB-producing strain described in Example 2, which was modified to not comprise any additional KO genes. Thus, steviol glycosides, including 13-SMG, 1,2-bioside, and RebB, were only accumulated upon expression of a functional KO. Three scrapes (1  $\mu\text{L}$  loop of cells) from each transformation plate were resuspended in 200  $\mu\text{L}$  nanopure  $\text{H}_2\text{O}$ . 70  $\mu\text{L}$  were then transferred to 1 mL SC-URA in a 96 deep well plate and incubated at 30°C for 5 days at 400 rpm. Biological triplicates were analyzed by LC-MS (method 2 of Example 1) to measure 13-SMG, 1,2-bioside, and RebB levels, and single samples were analyzed by LC-UV to measure ent-kaurene and ent-kaurenoic acid levels.

[00198] For LC-MS, 50  $\mu\text{L}$  samples were mixed with 50  $\mu\text{L}$  100% DMSO and heated to 80°C for 10 min. Subsequently, the samples were spun down at 4000 RCF for 10 min, and 85  $\mu\text{L}$  of the resulting supernatant was transferred to an LC-MS plate. The LC-MS results were normalized by  $\text{OD}_{600}$  of individual cultures, which was measured by a Wallac, 2104 EnVision (Perkin Elmer) plate reader.

[00199] LC-UV was conducted with an Agilent 1290 instrument comprising a variable wavelength detector (VWD), a thermostatted column compartment (TCC), an autosampler, an autosampler cooling unit, and a binary pump and using SB-C18 rapid resolution high definition (RRHD) 2.1 mm x 300 mm, 1.8  $\mu\text{m}$  analytical columns (two 150 mm columns in series; column temperature of 65°C). Steviol glycosides and steviol glycoside precursors were separated by a reversed phase C18 column followed by detection by UV absorbance at 210 nm. Quantification of steviol glycosides was done by comparing the peak area of each analyte to standards of RebA and applying a correction factor for species with differing molar



absorptivities. Quantification of steviol glycoside precursors (such as kaurenoic acid, kaurenal, kaurenol, ent-kaurene, and geranylgeraniol) was done by comparing the peak area of each analyte to standards of kaurenoic acid and applying a correction factor for species with differing molar absorptivities. For LC-UV, 0.5 rriL cultures were spun down, the supernatant was removed, and the wet weight of the pellets was calculated. The LC-UV results were normalized by pellet wet weight.

**[00200]** As shown in Figures 16B and 16D, the *S. cerevisiae* strain transformed with empty plasmid accumulated ent-kaurene. Transformation with a plasmid comprising SrKOI (SEQ ID NO:59, SEQ ID NO:79) or with a plasmid comprising the KO gene having the nucleotide sequence set forth in SEQ ID NO:65 resulted in accumulation of 13-SMG, 1,2-bioside, and RebB (Figures 16A and 186C).

**[00201]** Expression of full-length SrKOI -BMR fusion constructs (wild type or W1046A mutant BMR; SEQ ID NOs:99-102), resulted in an increase in ent-kaurenoic acid, 13-SMG, and RebB, compared to expression of SrKOI (SEQ ID NO:59, SEQ ID NO:79). See Figures 16A and 16B. Expression of truncated SrKOI -BMR fusion constructs (wild type or W1046A mutant BMR; SEQ ID NOs:103-106) resulted in an increase in ent-kaurenoic acid, compared to expression of SrKOI (SEQ ID NO:59, SEQ ID NO:79) (Figure 16B). Although the truncated SrKOI -BMR fusion constructs also increased steviol glycoside production, glycosylation activity was higher for the full-length SrKOI-BMR fusion constructs than for the truncated SrKOI -BMR fusion constructs (Figure 16A).

**[00202]** Expression of a fusion construct comprising the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 and the wild type BMR (SEQ ID NO:107, SEQ ID NO:108) resulted in greater conversion of ent-kaurenoic acid to 13-SMG, compared to the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 (Figure 16C). Expression of a fusion construct comprising the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 and the W1046A mutant BMR (SEQ ID NO:109, SEQ ID NO:110) resulted in decreases in ent-kaurenoic acid levels but glycosylation activity similar to that of the KO encoded by the nucleotide sequence set forth in SEQ ID NO:65 (Figure 16C).

#### **Example 11. Evaluation of Steviol Glycoside Pathway in *S. cerevisiae* Strain Comprising ICE2**

**[00203]** ICE2 is an endoplasmic reticulum (ER) membrane protein involved in mechanisms such as ER zinc homeostasis and cytochrome P450 stability and/or activity. See, e.g., Estrada de Martin *et al.*, 2005, *J Cell Sci.* 118(Pt 1):65-77 and Emmerstorfer *et al.*, 2015, *Biotechnol J.* 10(4):623-35. ICE2 (SEQ ID NO:113, SEQ ID NO:114) was cloned and overexpressed in a steviol glycoside-producing *S. cerevisiae* strain comprising a recombinant gene encoding a *Synechococcus sp.* GGPPS polypeptide (SEQ ID NO:49), a recombinant gene encoding a truncated *Z. mays* CDPS polypeptide (SEQ ID NO:37), a recombinant gene encoding an *A. thaliana* KS polypeptide (SEQ ID NO:6), a recombinant gene encoding a recombinant *S. rebaudiana* KO polypeptide (SEQ ID NO:59, SEQ ID NO:79), a recombinant gene encoding an *A. thaliana* ATR2 polypeptide (SEQ ID NO:51, SEQ ID NO:87), a recombinant gene encoding an SrKAHel (SEQ ID NO:18, SEQ ID NO:68) polypeptide, a recombinant gene encoding an *S. rebaudiana* CPR8 polypeptide (SEQ ID NO:24, SEQ ID NO:28), a recombinant KAH gene encoded by the nucleotide sequence set forth in SEQ ID NO:81 (corresponding to the amino acid sequence set forth in SEQ ID NO:82), a recombinant KO gene encoded by the nucleotide sequence set forth in SEQ ID NO:56 (corresponding to the amino acid sequence set forth in SEQ ID NO:75), a recombinant KO gene encoded by the nucleotide sequence set forth in SEQ ID NO:65 (corresponding to the amino acid sequence set forth in SEQ ID NO:75), a recombinant gene encoding a UGT76G1 (SEQ ID NO:83) polypeptide, a recombinant gene encoding an *S. rebaudiana* UGT85C2 polypeptide (SEQ ID NO:30), a recombinant gene encoding an *S. rebaudiana* UGT74G1 polypeptide (SEQ ID NO:29), a recombinant gene encoding an EUGT11 (SEQ ID NO:86) polypeptide, a recombinant gene encoding a UGT91 D2e (SEQ ID NO:84) polypeptide, and a recombinant gene encoding a CPR1 (SEQ ID NO:61, SEQ ID NO:76) polypeptide. Overexpression was performed by integration using the USER cloning system; see, e.g., Nour-Eldin *et al.*, 2010, *Methods Mol Biol.* 643:185-200. Table 9 shows additional recombinant genes (ICE2 and/or CPR12) expressed in the above-described strain. The control strain did not comprise recombinant genes encoding ICE2 (SEQ ID NO:113, SEQ ID NO:114) or CPR12 (SEQ ID NO:97, SEQ ID NO:98) polypeptides.

**Table 9: ICE2 steviol glycoside-producing strains.**

Strain	Sequences
ICE2 "strain A"	ICE2 (SEQ ID NO:113, SEQ ID NO:114) Overexpressed CPR1 (SEQ ID NO:61, SEQ ID NO:76)
ICE2 "strain B"	ICE2 (SEQ ID NO:113, SEQ ID NO:114) (2 copies)

ICE2 "strain C"	ICE2 (SEQ ID NO:113, SEQ ID NO:114) CPR12 (SEQ ID NO:97, SEQ ID NO:98)
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**[00204]** Fed-batch fermentation was carried out aerobically in 2 L fermenters at 30°C with an approximate 16 h growth phase in minimal medium comprising glucose, ammonium sulfate, trace metals, vitamins, salts, and buffer followed by an approximate 110 h feeding phase with a glucose-comprising defined feed medium. A pH near 6.0 and glucose-limiting conditions were maintained. Whole culture samples (without cell removal) were analysed by the LC-UV method of Example 10 to determine levels of steviol glycosides and steviol pathway intermediates.

**[00205]** The following values were calculated based upon the measured levels of steviol glycosides and steviol glycoside precursors. "Total Flux" was calculated as a sum (in g/L RebD equivalents) of measured RebA, RebB, RebD, RebE, RebM, 13-SMG, rubusoside, steviol-1,2-bioside, di-glycosylated steviol, tri-glycosylated steviol, tetra-glycosylated steviol, penta-glycosylated steviol, hexa-glycosylated steviol, hepta-glycosylated steviol, copalol, ent-kaurenoic acid, glycosylated ent-kaurenoic acid, glycosylated ent-kaurenol, ent-kaurenal, geranylgeraniol, ent-kaurenal, and ent-kaurene levels. "Pre-steviol glycoside/flux" was calculated as  $((\text{"total flux"} - (\text{geranylgeraniol} + \text{copalol} + \text{ent-kaurene} + \text{glycosylated ent-kaurenol} + \text{ent-kaurenol} + \text{ent-kaurenal} + \text{ent-kaurenoic acid} + \text{glycosylated ent-kaurenoic acid})) / \text{"total flux"})$ . "KAH step/flux" was calculated as  $((\text{ent-kaurenoic acid} + \text{glycosylated ent-kaurenoic acid}) / \text{"total flux"})$ . "KO step/flux" was calculated as  $((\text{ent-kaurene} + \text{glycosylated ent-kaurenol} + \text{ent-kaurenol} + \text{ent-kaurenal}) / \text{"total flux"})$ .

**[00206]** The pre-steviol glycoside/flux, KO step/flux, and KAH step/flux values are shown in Table 10 below. Decreased amounts of ent-kaurene, ent-kaurenol, ent-kaurenal, glycosylated ent-kaurenol and increased amounts of ent-kaurenoic acid and glycosylated ent-kaurenoic acid were observed in the strains comprising ICE2, as compared to the control steviol glycoside-producing strain. These effects were stronger in the presence of CPR1 and/or CPR12 (Table 10). Overexpression of two copies of ICE2 (ICE2 strain B) resulted decreased ent-kaurene, ent-kaurenol, ent-kaurenal, and ent-kaurenol glycoside levels and increased steviol glycoside levels, compared to the control strain, ICE2 strain A, or ICE2 strain C (Table 10). Steviol glycoside levels increased most in the steviol glycoside-producing strain comprising two copies of ICE2. Thus, ICE2 was found to improve cytochrome P450 function.

**Table 10: Pre-steviol glycoside/flux, KO step/flux, and KAH step/flux values for steviol glycoside-producing strains comprising ICE2.**

Strain	Pre-Steviol Glycoside/Flux	KO step/Flux	KAH step/Flux
ICE2 "strain A"	0.38	0.36	0.22
ICE2 "strain B"	0.43	0.42	0.10
ICE2 "strain C"	0.39	0.38	0.19
Control	0.41	0.48	0.08

**Example 12. Steviol Glycoside Production by Fermentation of *S. cerevisiae* strain comprising CPR1 and CPR12**

[00207] Steviol glycoside-producing *S. cerevisiae* strains comprising a recombinant gene encoding a *Synechococcus sp.* GGPPS polypeptide (SEQ ID NO:49), a recombinant gene encoding a truncated *Z. mays* CDPS polypeptide (SEQ ID NO:37), a recombinant gene encoding an *A. thaliana* KS polypeptide (SEQ ID NO:6), a recombinant gene encoding a recombinant *S. rebaudiana* KO polypeptide (SEQ ID NO:59, SEQ ID NO:79), a recombinant gene encoding an *A. thaliana* ATR2 polypeptide (SEQ ID NO:51, SEQ ID NO:87), a recombinant gene encoding an SrKAHei (SEQ ID NO:18, SEQ ID NO:68) polypeptide, a recombinant gene encoding an *S. rebaudiana* CPR8 polypeptide (SEQ ID NO:24, SEQ ID NO:28), a recombinant gene encoding a CPR1 (SEQ ID NO:61, SEQ ID NO:76) polypeptide, a recombinant gene encoding an SrKAHei (SEQ ID NO:18, SEQ ID NO:68) polypeptide, a recombinant KO gene encoded by the nucleotide sequence set forth in SEQ ID NO:56 (corresponding to the amino acid sequence set forth in SEQ ID NO:75), a recombinant gene encoding a UGT76G1 (SEQ ID NO:83) polypeptide, a recombinant gene encoding an *S. rebaudiana* UGT85C2 (SEQ ID NO:30) polypeptide, a recombinant gene encoding an *S. rebaudiana* UGT74G1 (SEQ ID NO:29) polypeptide, a recombinant gene encoding a UGT91D2e-b polypeptide (SEQ ID NO:88), and a recombinant gene encoding an EUGT1 1 (SEQ ID NO:86) polypeptide, as well as the recombinant genes shown in Table 11, which were genomically integrated into the strains, were cultivated by fermentation. Levels of steviol glycosides and steviol glycoside precursors were measured by LC-UV as described in Example 11. The pre-KO/flux, pre-KAH/flux, pre-steviol glycoside/flux values were calculated as described in Example 11.

**Table 11: Recombinant genes also expressed in steviol glycoside-producing *S. cerevisiae* strain in Example 12.**

Strain	Genes

Example 12, Strain A	KO encoded by nucleotide sequence set forth in SEQ ID NO:56 (corresponding to amino acid sequence set forth in SEQ ID NO:75)
Example 12, Strain B	KAH encoded by nucleotide sequence set forth in SEQ ID NO:80 (corresponding to amino acid sequence set forth in SEQ ID NO:82)  KO encoded by nucleotide sequence set forth in SEQ ID NO:56 (corresponding to amino acid sequence set forth in SEQ ID NO:75)  KO encoded by nucleotide sequence set forth in SEQ ID NO:65 (corresponding to amino acid sequence set forth in SEQ ID NO:75)
Example 12, Strain C	CPR12 (SEQ ID NO:97, SEQ ID NO:98)  KAH encoded by nucleotide sequence set forth in SEQ ID NO:80 (corresponding to amino acid sequence set forth in SEQ ID NO:82)  KO encoded by nucleotide sequence set forth in SEQ ID NO:56 (corresponding to amino acid sequence set forth in SEQ ID NO:75)

[00208] The pre-steviol glycoside/flux, KO step/flux, and KAH step/flux values are shown in Table 12 below. In the strain comprising the KO encoded by nucleotide sequence set forth in SEQ ID NO:56 (strain A), lower accumulation of ent-kaurene, ent-kaurenol, ent-kaurnal, and ent-kaurenol glycosides resulted. Higher levels of ent-kaurenoic acid and steviol glycosides were also measured, as compared to the control strain. In the strain comprising the KAH encoded by nucleotide sequence set forth in SEQ ID NO:80, the KO encoded by nucleotide sequence set forth in SEQ ID NO:56 (corresponding to amino acid sequence set forth in SEQ ID NO:75), and the KO encoded by nucleotide sequence set forth in SEQ ID NO:65 (strain B), ent-kaurene, ent-kaurenol, ent-kaurenal, ent-kaurenol glycosides, and ent-kaurenoic acid accumulation decreased and accumulation of steviol glycosides increased, as compared to the control strain. In the strain comprising CPR12 (SEQ ID NO:97, SEQ ID NO:98), the KAH encoded by nucleotide sequence set forth in SEQ ID NO:80, and the KO encoded by nucleotide sequence set forth in SEQ ID NO:56 (strain C), ent-kaurenol, ent-kaurenal, ent-kaurenol glycosides, and ent-kaurenoic acid accumulation decreased and accumulation of steviol glycosides increased, as compared to the control. See Table 12. Thus, CPR12 was found to be a reductase protein that improves KAH and/or KO activity.

**Table 12. Pre-steviol glycoside/flux, KO step/flux, and KAH step/flux values for steviol glycoside-producing strains of Example 12.**

Strain	Pre-Steviol Glycoside/Flux	KO step/Flux	KAH step/Flux
Example 12, Strain A	0.48	0.28	0.22
Example 12, Strain B	0.64	0.18	0.12
Example 12, Strain C	0.55	0.24	0.12
Control	0.40	0.43	0.17

**[00209]** Having described the invention in detail and by reference to specific embodiments thereof, it will be apparent that modifications and variations are possible without departing from the scope of the invention defined in the appended claims. More specifically, although some aspects of the present invention are identified herein as particularly advantageous, it is contemplated that the present invention is not necessarily limited to these particular aspects of the invention.

**Table 13. Sequences disclosed herein.**

**SEQ ID NO:1**

```

MNLSLCIASP LLTKSNRPAA LSAIHTASTS HGGQTNPTNL IIDTKKERIQ KQFKNVEISV      60
SSYDTAWVAM VSPNSPKSP CFPECLNWLI NNQLNDGSWG LVNHHTHNNH PLLKDSLST      120
LACIVALKRW NVGEDQINKG LSFIESNLAS ATEKSQPSPI GFDIIFPGLL EYAKNLDINL      180
LSKQTDfSLM LHKRELEQKR CHSNEMDGYL AYISEGLGNL YDWNMVKKYQ MKNGSVFNsp      240
SATAAAFINH QNPGCLNYLN SLLDKFGNAV PTVYPHDLFI RLSMVDTIER LGISHHFRVE      300
IKNVLDETYR CWVERDEQIF MDVVTALAF RLLRINGYEV SPDPLAEITN ELALKDEYAA      360
LETYHASHIL YQEDLSSGKQ ILKSADFLKE IISTDSNRLS KLIHKEVENA LKFPINTGLE      420
RINTRRNIQL YNVNTRILK TTYHSSNISN TDYLRLAVED FYTCQSIYRE ELKGLERWVV      480
ENKLDQLKFA RQKTAYCYFS VAATLSSPEL SDARISWAKN GILTTVVDDF FDIGGTIDEL      540
TNLIQCVEKW NVDVDKDCCS EHVRLFLAL KDAICWIGDE AFKWQARDVT SHVIQWLEL      600
MNSMLREAIW TRDAYVPTLN EYMENAYVSF ALGPVVKPAI YFVGPKLSEE IVESSEYHNL      660
FKLMSTQGRL LNDIHSFKRE FKEGKLNVA LHLNNGESGK VEEVVEEMM MMIKNKRKEL      720
MKLIFEENGs IVPRACTDAF WNMCHVLNFF YANDDGFTGN TILDTVKDII YNPLVLVNEN      780
EEQR                                                                 784
    
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**SEQ ID NO:2**

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MNLSLCIASP LLTKSSRPTA LSAIHTASTS HGGQTNPTNL IIDTKKERIQ KLFKNVEISV      60
SSYDTAWVAM VSPNSPKSP CFPECLNWLI NNQLNDGSWG LVNHHTHNNH PLLKDSLST      120
LACIVALKRW NVGEDQINKG LSFIESNLAS ATDKSQPSPI GFDIIFPGLL EYAKNLDINL      180
LSKQTDfSLM LHKRELEQKR CHSNEIDGYL AYISEGLGNL YDWNMVKKYQ MKNGSVFNsp      240
SATAAAFINH QNPGCLNYLN SLLDKFGNAV PTVYPLDLYI RLSMVDTIER LGISHHFRVE      300
IKNVLDETYR CWVERDEQIF MDVVTALAF RLLRIHGYKV SPDQLAEITN ELAFKDEYAA      360
LETYHASQIL YQEDLSSGKQ ILKSADFLKG IISTDSNRLS KLIHKEVENA LKFPINTGLE      420
    
```

RINTRRNIQL	YNVDNTRILK	TTYHSSNISN	TYYLRLAVED	FYTCQSIYRE	ELKGLERWV	480
QNKLDQLKFA	RQKTAYCYFS	VAATLSSPEL	SDARISWAKN	GILTTVVDDF	FDIGGTIDEL	540
TNLIQCVEKW	NVDVDKDCCS	EHVRILFLAL	KDAICWIGDE	AFKWQARDVT	SHVIQTWLEL	600
MNSMLREAIW	TRDAYVPTLN	EYMENAYVSF	ALGPIVKPAI	YFVGPKLSEE	IVESSEYHNL	660
FKLMSTQGRL	LNDIHSFKRE	FKEGKLNVA	LHLSNGESGK	VEEEVVEEMM	MMIKNRKREL	720
MKLI FEENG	IVPRACKDAF	WNMCHVLNFF	YANDDGTGN	TILDTVKDII	YNPLVLVNE	780
EEQR						784

SEQ ID NO:3

MAMPVKLTPA	SLSLKAVCCR	FSSGGHALRF	GSSLPWRRRT	PTQRSTSSST	TRPAAEVSSG	60
KSKQHDQEAS	EATIROQLQL	VDVLENMGIS	RHFAAEIKCI	LDRTYRSWLQ	RHEEIMLDTM	120
TCAMAFRILR	LNGYNVSSDE	LYHVVEASGL	HNSLGGYLN	TRTLLELHKA	STVSISEDES	180
ILDSIGRSR	TLLREQLESG	GALRKPSLFK	EVEHALDGGP	YTTLDRLHHR	WNIEFNIE	240
QHMLETPYLS	NQHTSRDILA	LSIRDFSSSQ	FTYQQELQHL	ESWVKECRLD	QLQFARQKLA	300
YFYLSAAGTM	FSEPLSDART	LWAKNGVLT	IVDDFFDVAG	SKEELENLVM	LVEMWDEHHK	360
VEFYSEQVEI	EFSDNATCA	QLGEKASLVQ	DRSITKHLVE	IWLDDLKSM	TEVEWRLSKY	420
VPTKEYMIN	ASLIFGLGPI	VLPALYFVGP	KISESIVKDP	EYDELFLKMS	TCGRLLNDVQ	480
TFEREYNEGK	LNSVSLVLH	GGPMSISDAK	RKLQKPIDTC	RRDLSLVL	EESVVPKCK	540
ELFWKMCKVC	YFFYSTTDGF	SSQVERAKEV	DAVINEPLKL	QGSHTLVSDV		590

SEQ ID NO:4

MSCIRPWFCP	SSISATLTD	ASKLVTGEFK	TTSLNFGTK	ERIKKMFDKI	ELSVSSYDTA	60
WVAMVPSDC	PETPCFPECT	KWILENQLGD	GSWSLPHGNP	LLVKDALSS	LACILALKRW	120
GIGEEQINKG	LRFIELNSAS	VTDNEQHKPI	GFDIIFPGMI	EYAKDLNL	PLKPTDINS	180
LHRRALELTS	GGGNLEGR	AYLAYVSEGI	GKLQWEMAM	KYQRKNGSLF	NSPSTAAAF	240
IHIQDAECLH	YIRSLQKFG	NAVPTIYPLD	IYARLSMVDA	LERLIGDRHF	RKERKVLDE	300
TYRFWLQGEE	EIFSDNATCA	LAFRILRLNG	YDVSLEDHFS	NSLGGYKDS	GAALELYRAL	360
QLSYPDESLL	EKQNSRTSYF	LKQGLSNVSL	CGDRLRKNII	GEVHDALNFP	DHANLQRLAI	420
RRRIKHYATD	DTRILKTSYR	CSTIGNQDFL	KLAVEDFNIC	QSIQREEFKH	IERWVVERRL	480
DKLKFARQKE	AYCYFSAAT	LFAPELSDAR	MSWAKNGVLT	TVVDDFFDVG	GSEELVNL	540
ELIERWDVNG	SADFCSEVE	IIYSAIHSTI	SEIGDKSFGW	QGRDVKSHVI	KIWLDDLKSM	600
LTEAQWSSNK	SVPTLDEYMT	TAHVSFALGP	IVLPALYFVG	PKLSEEVAGH	PELLNLYKVM	660
STCGRLLNDW	RSEFKRESEEG	KLNAISLYMI	HSGGASTEET	TIEHFKGLID	SQRRQLQLV	720
LQEKDSIIPR	PCKDLFWNMI	KLHHTFYMKD	DGFTSNEMRN	VVKAIINEPI	SLDEL	775

SEQ ID NO:5

cgctcagtc	caaggcta	tcgctcgag	ttgctacgac	gccgtttcg	ttgcttctg	60
ttctttatg	tctatcaacc	ttcgctcctc	cggttggtcg	tctccgatct	cagctacttt	120
ggaacgagga	ttggactcag	aagtacagac	aagagctaac	aatgtgagct	ttgagcaaac	180
aaaggagaag	attaggaaga	tggtggagaa	agtgagctt	tctgtttcgg	cctacgatac	240
tagttgggta	gcaatgggtc	catcaccgag	ctcccaaaat	gctccacttt	tcccacagtg	300
tgtagaatg	ttattggata	atcaacatga	agatggatct	tggggacttg	ataaccatga	360
ccatcaatct	cttaagaagg	atgtgttatc	atctacactg	cgtagtatcc	tcgctgtaaa	420
gaagtgggga	attggtgaaa	gacaaataaa	caagggtctc	cagtttattg	agctgaattc	480
tgcattagtc	actgatgaaa	ccatacagaa	accaacaggg	ttgatatta	tatttctctg	540
gatgattaaa	tatgctagag	atttgaatct	gacgattcca	ttgggctcag	aagtgggtga	600
tgacatgata	cgaaaaagag	atctggatct	taaatgtgat	agtgaaaagt	tttcaaagg	660
aagagaagca	tatctggcct	atgttttaga	ggggacaaga	aacctaaaag	attgggattt	720
gatagtcaaa	tatcaaagga	aaaatgggtc	actgtttgat	tctccagcca	caacagcagc	780
tgctttact	cagtttgga	atgatggtg	tctccggtat	ctctgttctc	tccttcagaa	840
attcgaggct	gcagttcctt	cagtttatcc	atltgatcaa	tatgcacgcc	ttagtataat	900
tgctactcct	gaaagcttag	gaattgatag	agatttcaaa	accgaaatca	aaagcatatt	960
ggatgaaacc	tatagatatt	ggcttcgtgg	ggatgaagaa	atatgtttgg	acttggccac	1020
ttgtgctttg	gctttccgat	tattgcttgc	tcatggctat	gatgtgtcct	acgatccgct	1080
aaaaccattt	gcagaagaat	ctggtttctc	tgatactttg	gaaggatatg	ttaagaatac	1140
gttttctgtg	ttagaattat	ttaaggctgc	tcaaagttat	ccacatgaat	cagctttgaa	1200
gaaagcagtg	tgttggacta	aacaatatct	ggagatggaa	ttgtccagct	gggttaagac	1260
ctctgttcga	gataaatacc	tcaagaaaga	ggtcagggat	gctcttgctt	ttccctccta	1320
tgcaagccta	gaaagatcag	atcacaggag	aaaaatactc	aatggttctg	ctgtggaaaa	1380
caccagagtt	acaaaaacct	catatcgttt	gcacaatatt	tgacacctcg	atatcctgaa	1440
gttagctgtg	gatgacttca	atctctgcca	gtccatacac	cgtgaagaaa	tggaacgtct	1500
tgataggtgg	attgtggaga	atagattgca	ggaactgaaa	tttgccagac	agaagctggc	1560

ttactgttat	ttctctgggg	ctgcaacttt	atthttctcca	gaactatctg	atgctcgtat	1620
atcgtgggccc	aaaggtggag	tactttacaac	ggttgtagac	gacttctttg	atggttgagg	1680
gtccaaagaa	gaactggaaa	acctcataca	cttggctgaa	aagtgggatt	tgaacgggtg	1740
tcctgagtag	agctcagaac	atggttagat	catattctca	gttctaaggg	acaccattct	1800
cgaaacagga	gacaaagcat	tcacctatca	aggacgcaat	gtgacacacc	acattgtgaa	1860
aatthgggtg	gatctgctca	agtctatggt	gagagaagcc	gagtgggtcca	gtgacaagtc	1920
aacaccaagc	ttggaggatt	acatggaaaa	tgcgtagata	tcatttgcac	taggaccaat	1980
tgtcctoccca	gctacctatc	tgatcggacc	tcactttcca	gagaagacag	tcgatagcca	2040
ccaatataat	cagctctaca	agctcgtgag	cactatgggt	cgtcttctaa	atgacataca	2100
aggttttaag	agagaaagcg	cggaagggaa	gctgaatgcg	gtttcattgc	acatgaaaca	2160
cgagagagac	aatcgacagc	aagaagtgat	catagaatcg	atgaaaagtt	tagcagagag	2220
aaagagggaa	gaattgcata	agctagtgtt	ggaggagaaa	ggaagtgtgg	ttccaaggga	2280
atgcaaagaa	gcgttcttga	aaatgagcaa	agtgttgaa	ttattttaca	ggaaggacga	2340
tggattcaca	tcaaatgatc	tgatgagtct	tgtaaatca	gtgatctacg	agcctgttag	2400
cttacagaaa	gaatctttaa	cttgatccaa	gttgatctgg	caggtaaac	cagtaaatga	2460
aaataagact	ttggtcttct	tctttgttgc	ttcagaacaa	gaagag		2506

SEQ ID NO:6

MSINLRSSGC	SSPISATLER	GLDSEVQTRA	NNVSFEQTK	KIRKMLEKVE	LSVSAYDTSW	60
VAMVPSPPSSQ	NAPLFPQCVK	WLLDNQHEDG	SWGLDNHDHQ	SLKKDVLSS	LASILALKKW	120
GIGERQINKG	LQFIELNSAL	VTDETIQKPT	GFDIIFPGMI	KYARDLNLTI	PLGSEVVDDM	180
IRKRDLDLKC	DSEKFSKGRE	AYLAYVLEGT	RNLKDWDLIV	KYQRKNGSLF	DSPATTAAAF	240
TQFGNDGCLR	YLCSLLQKFE	AAVPSVYPPD	QYARLSIIVT	LESLGIDRDF	KTEIKSILDE	300
TYRWLRGDE	EICLDLATCA	LAFRLLLAHG	YDVSYDPLKP	FAEESGFSDT	LEGYVKNTFS	360
VLELFKAAQS	YPHESALKKQ	CCWTKQYLEM	ELSSVVKTSV	RDKYLKKEVE	DALAFPSYAS	420
LERSDHRRKI	LNGSAVENTR	VTKTSYRLHN	ICTSDILKLA	VDDFNFCQSI	HREEMERLDR	480
WIVENRLQEL	KFARQKLAYC	YFSGAATLFS	PELSDARISW	AKGGVLTTVV	DDFFDVGGSK	540
EELNLIHLV	EKWDLNGVPE	YSSEHVEIIF	SVLRDTILET	GDKAFYQGR	NVTHHIVKIV	600
LDLLKSMRL	AEWSSDKSTP	SLEDYMENAY	ISFALGPVIV	PATYLVGPP	PEKTVDSHQY	660
NQLYKLVSTM	GRLLNDIQGF	KRESABGKLN	AVSLHMKHER	DNRSKVEVIE	SMKGLAERKR	720
EELHKLVL	KGSVPRECK	EAFLRMSKVL	NLFYRKDDGF	TSNDLMSLVK	SVIYEPVSLQ	780
KESLT						785

SEQ ID NO:7

MDAVTGLLTV	PATAITIGGT	AVALAVALIF	WYLKSYTSAR	RSQSNHLPRV	PEVPGVPLLG	60
NLLQLKEKPK	YMTFTRWAAT	YGPIYSIKTG	ATSMVVSSN	EIAKEALVTR	FQSISTRNLS	120
KALKVLTADK	TMVAMSDYDD	YHKTVKRHIL	TAVLGPNAQK	KHRIHRDIMM	DNISTQLHEF	180
VKNPEQEEV	DLRKIFQSEL	FGLAMRQALG	KDVESLYVED	LKITMNRDEI	FQVLVDPMM	240
GAIIDVDRDF	FPYLKWVPNK	KFENTIQQMY	IRREAVMKS	IKEHKRIAS	GEKLSYIDY	300
LLSEAQTLTD	QQLMSSLWEP	IIESDRTMV	TTEWAMYELA	KNPKLQDRLY	RDIKSVCGSE	360
KITTEHLSDL	PYITAIHFET	LRRHSPVPII	PLRHVHEDTV	LGGEYHVPAGT	ELAVNIYGCN	420
MDKNVWENPE	EWNPERFMKE	NETIDFQKTM	AFGGGKRVCA	GSLQALLTAS	IGIGRMVQEF	480
EWKLDKMTQE	EVNTIGLTTQ	MLRPLRAIIC	PRI			513

SEQ ID NO:8

MAFFSMISIL	LGFVISSFI	IFFFKLLSF	SRKNMSEVST	LPSVPPVPGF	PVIGNLLQLK	60
EKKPHKTFTR	WSEIYGPIYS	IKMGSSSLIV	LNSTETAKEA	MVTRFSSIST	RKLSNALTIV	120
TCDKSMVATS	DYDDPHKLVK	RCLLNGLLGA	NAQKRKRHYR	DALIENVSSK	LHAHARDHPQ	180
EPVNFRAIFE	HELFGVALKQ	AFGKDVESIY	VKELGVTLK	DEIFKVLVHD	MMEGAIDVDW	240
RDFFPYLKWI	PNKSFEARIQ	QKHRRRLAVM	NALIQRDLKQ	NGESDSDDCY	LNFLMSEAKT	300
LTKEQIAILV	HETIETADT	TLVTEWAIY	ELAKHPSVQD	RLCKBIQNV	GGEKFKEEQL	360
SQVPYLNQVF	HETLRKYS	PLVPIRYAHE	DTQIGGYHVP	AGSEIAINIY	GCNMDKKRWE	420
RPEDWWPERF	LDDGKYETSD	LHKTMAFGAG	KRVCAGALQA	SLMAGIAIGR	LVQEFWKLRL	480
DGEENVDTY	GLTSQKLYPL	MAIINPRRS				509

SEQ ID NO:9

MSKSNSMNST	SHETLFQQLV	LGLDRMPLMD	VHWLIYVAFG	AWLCSYVIHV	LSSSSTVKVP	60
VGYRSVFEP	TWLLRLRFVW	EGGSIIGQGY	NKFKDSIFQV	RKLGTDIVII	PPNYIDEVRK	120
LSQDKTRSV	PFINDFAGQY	TRGMVFLQSD	LQNRVIQQL	TPKLVSLTKV	MKEELDYALT	180
KEMPEMKNDE	WVEVDISSIM	VRLISRISAR	VFLGPEHCRN	QEWLTTTAEY	SESLEFITGFI	240



LRVVPILRP	FIAPLLPSYR	TLLRNVSAGR	RVIGDIIRSQ	QGDGNEDILS	WMRDAATGEE	300
KQIDNIAQRM	LILSLASIHT	TAMTMTHAMY	DLCACPEYIE	PLRDEVKSVV	GASGWDKTAL	360
NRPHKLDSEFL	KESQRFNPVF	LLTFNRIYHQ	SMTLSDGTNI	PSGTRIAVPS	HAMLQDSAHV	420
PGPTPTEFD	GFRYSKIRSD	SNYAQKYLFS	MTDSSNMAFG	YGKYACPGRF	YASNEMKLTL	480
AILLQFEFK	LPDGKGRPRN	ITIDSDMIPD	PRARLCVRKR	SLRDE		525

SEQ ID NO:10

MEDPTVLYAC	LAIAVATFV	RWYRDPLRSI	PTVGGSDLPI	LSYIGALRWT	RRGREILQEG	60
YDGYRGSTFK	IAMLDRWIVI	ANGPKLADEV	RRRPDEELNF	MDGLGAFVQT	KYTLGEAIHN	120
DPYHVDIIRE	KLTRGLPAVL	PDVIEELTLA	VRQYIPTEGD	EWVSVNCSKA	ARDIVARASN	180
RVFVGLPACR	NQGYLDLAI	FTLSVVKDRA	IINMFPELLK	PIVGRVVGNA	TRNVRRVAVPF	240
VAPLVEERRR	LMEEYGEDWS	EKPNDMLQWI	MDEAASRDSS	VKAIARLLM	VNFAAIHTSS	300
NTITHALYHL	AEMPETLQPL	REEIEPLVKE	EGWTKAAMGK	MWWLDSFLRE	SQRYNGINIV	360
SLTRMADKDI	TLSDGTFLPK	GTLVAVPAYS	THRDDAVYAD	ALVFDPRFVS	RMRAREGEGT	420
KHQFVNTSVE	YVFPFHGKHA	CPGRFFAANE	LKAMLAYIVL	NYDVKLPGDG	KRPLNMYWGP	480
TVLPAPAGQV	LFRKRQVSL					499

SEQ ID NO:11

aaacaaagaa	tgattcaagt	tctaacaccg	atccttctct	tcctcatttt	cttcgttttc	60
tggaaggttt	acaagcacca	gaaaacccaa	atcaatcttc	caccgggaag	cttcggatgg	120
ccatttctgg	gcaaaactct	ggcactccta	cgtgcaggtt	gggactcaga	gccggagaga	180
ttgttctgtg	aacggatcaa	gaaacacgga	agtctcttag	tgtttaagac	gtcgttggtt	240
ggcgaccgtt	ttgcggtggt	gtgtggacct	gccggaaaca	agttcctggt	ctgcaacgag	300
aaacagctgg	tggcgtcgtg	gtggccggtt	ccggtgagga	agcttttcgg	caagtctctg	360
ctcacgattc	gtgggtgatga	agctaagtgg	atgaggaaga	tgttgttata	gtatctcggt	420
cctgatgctt	tgcgaactca	ttatgccgtc	accatggagc	tcgtcacccg	tcggcatatc	480
gacgttcatt	ggcgagggaa	ggaagaggtg	aacgtattcc	aaaccgttaa	gttatatgcc	540
tttgagcttg	catgtcgttt	attcatgaac	ctagacgacc	caaaccacat	tgcaaaaactc	600
ggttccttgt	tcaacatttt	cttgaaaggc	atcattgagc	ttccaatcga	cgtcccaggg	660
acacgatttt	atagctccaa	aaaagcagca	gcagctatca	ggattgaact	aaaaaaattg	720
attaaagcaa	gaaaactgga	actgaaagaa	gggaagggat	catcttcaca	agacctctta	780
tcacatttgc	ttacatctcc	agatgaaaaat	ggtatgtttc	taaccgaaga	agagattgta	840
gacaacatct	tgttactact	ctttgctggg	catgatacct	cggctctttc	aatcactttg	900
ctcatgaaga	ctcttgccga	acattctgat	gtttatgaca	aggtgttaaa	agagcaacta	960
gagatatcga	agacgaaaga	agcatgggag	tccctgaaat	gggaggacat	acaaaagatg	1020
aaatactcct	ggagtgattt	atgtgaagtc	atgagactaa	atccacctgt	tataggaacc	1080
tatagagagg	cccttggtga	tattgattat	gcgggttata	ccatccccaa	aggatggaag	1140
ctgcaactga	gtgctgtatc	gacacaaagg	gacgaggcta	actttgaaga	cgtaaacactg	1200
tttgaccat	cacggtttga	aggcgcagga	ccgactccat	tcacctttgt	tccgtttgga	1260
ggggggccta	gaatgtgttt	agggaaagaa	tttgctcgat	tggaagtact	tgcgtttcct	1320
cacaatattg	tcaccaattt	caaatgggac	ctgttgatac	ctgatgagaa	aatagaatat	1380
gatcccatgg	ctacccagc	aaaggggctt	ccaatctgtc	ttcatcccca	tcaagtttga	1440
ttacttcaag	catgaatcag	tgatgtgaag	gtaaaccata	atggatctta	ttggtagtta	1500
cagattatgt	gtttttatgg	catgaagaag	ttatgataaa	taaaattgtg	ttattctaca	1560
acttatgtaa	tttgctcctg	taagtaactg	aatctattaa	tgttttatgt	gacatgaaac	1620
ataaatgtat	aattagtaaa	ttttctgctc	aaaaaaaaaa	aaaaaaaaaa	aaaaaaaaaa	1678

SEQ ID NO:12

MIQVLTPIIL	FLIFFVFWKV	YKHQTKKINL	PPGSFGWPF	GETLALLRAG	WDSEPERFVR	60
ERIKKHGSPL	VFKTSLFGDR	FAVLCGPAGN	KFLFCNENKL	VASWWPVPVR	KLFGKSLTI	120
RGDEAKWMRK	MLLSYLGPA	FATHYAVTMD	VVTRRHIDVH	WRGKEEVNVE	QTVKLYAFEL	180
ACRLFMNLDD	PNHIAKLGSL	FNIFLKGIEE	LPIDVPGTRF	YSSKKAATAI	RIELKLIKA	240
RKLELKEGKA	SSSQDLLSHL	LTSPDENGME	LTEEEIVDNI	LLLLFAGHDT	SALSITLLMK	300
TLGEHSDVYD	KVLKEQLEIS	KTKEAWESLK	WEDIQMKYS	WSVICEVMRL	NPPVIGTYRE	360
ALVDIDYAGY	TIPKGWKLHW	SAVSTQRDEA	NFEDVTRFDP	SRFEGAGPTP	FTFVFPGGGP	420
RMCLGKEFAR	LEVLAFLHNI	VTNFKWDLII	PDEKIEYDPM	ATPAKGLPIR	LHPHQV	476

SEQ ID NO:13

MGLFPLEDSY	ALVFEGLAIT	LALYLLSFI	YKTSKKTCTP	PKASGEHPIT	GHLNLLSGSS	60
GLPHLALASL	ADRCGPIFTI	RLGIRRVLVV	SNWEIAKEIF	TTHDLIVSNR	PKYLAAKILG	120
FNYVVSFSFAP	YGPYVVGIRK	IIATKLMSSS	RLQKQLQFVRV	FELENSMKSI	RESWKEKKDE	180
EGKVLVEMKK	WFWELNMNIV	LRTVAGKQYT	GTVDDADAKR	ISELFREWFH	YTGRFVVGDA	240
FPFLGWLDLG	GKKTMEIVA	SRLDSMVSKW	LDEHRKKQAN	DDKKEDMDFM	DIMISMTEAN	300
SPLEGYGTDT	IIKTTCTMLI	VSGVDTTSIV	LTWALSLLN	NRDTLKKAE	ELDMCVGKGR	360
QVNESDLVNL	IYLEAVLKEA	LRLYPAAFGL	GPRAFLEDCT	VAGYRIPKGT	CLLINMWKLH	420
RDPNIWSDPC	EFKPERFLTP	NQKDVDVIGM	DFELIPFGAG	RRYCPGTRLA	LQMLHIVLAT	480
LLQNFEMSTP	NDAPVDMTAS	VGMTNAKASP	LEVLLSPRVK	WS		522

SEQ ID NO:14

MIQVLTPIILL	FLIFFVFWKV	YKHQKTKINL	PPGSFGWPFLL	GETLALLRAG	WDSEPERFVR	60
ERIKKHGSPL	VFKTSLFQDR	FAVLGCPAGN	KFLFCNENKL	VASWWPVPVR	KLFGKSLITI	120
RGDEAKWMRK	MLLSYLGPA	FATHYAVTMD	VVTRRHIDVH	WRGKEEVNMF	QTVKLYAFEL	180
ACRLFMNLDD	PNHIAKLGSL	FNIFLKGIIE	LPIDVPGTRF	YSSKKAATAI	RIELKKLIKA	240
RKLELKEGKA	SSSQDLLSHL	LTSPEDEMGF	LTEEEIVDNI	LLLLFAGHDT	SALSITLLMK	300
TLGEHSDVYD	KVLKEQLEIS	KTKEAWESLK	WEDIQMKYS	WSVICEVMRL	NPPVIGTYRE	360
ALVDIDYAGY	TIPKGWKLHW	SAVSTQDEA	NFEDVTRFDP	SRFEGAGPTP	FTFVFPFGGGP	420
RMCLGKEFAR	LEVLAFLHNI	VTNFKWDLI	PDEKIEYDPM	ATPAKGLPIR	LPHQV	476

SEQ ID NO:15

MESLVVHTVN	AIWCIVIVGI	FSVGYHVYGR	AVVEQWRMRR	SLKLQGVKGP	PPSIFNGNVS	60
EMQRIQSEAK	HCSGDNIIISH	DYSSSLFPHF	DHWRKQYGR	YTYSTGLKQH	LYINHPEMVK	120
ELSQTNTLNL	GRITHITKRL	NPILGNGIIT	SNGPHWAHQ	RIIAYEFTHD	KIKGMVGLMV	180
ESAMPMLNKW	EEMVKRGGEM	GCDIRVDEDL	KDVSADVIK	ACFGSSFSKG	KAIFSMIRDL	240
LTAITKRSVL	FRFNGFTDMV	FGSKKHGDVD	IDALEMELES	SIWETVKERE	IECKDTHKKD	300
LMQLILEGAM	RSCDGNLWDK	SAYRRFVVDN	CKSIYFAGHD	STAVSVSWCL	MLLALNPSWQ	360
VKIRDEILSS	KVNGIPDAES	IPNLKTVTMV	IQETMRLYPP	APIVGREASK	DIRLGLLVVP	420
KGVCIWTLIP	ALHRDPEIWG	PDANDFKPER	FSEGISKACK	YPQSYIPFGL	GPRTCVGKNF	480
GMMEVKVLVS	LIVSKFSFTL	SPTYQHSPSH	KLLVEPQHG	VIRVV		525

SEQ ID NO:16

MYFLLQYLN	TTVGVFATLF	LSYCLLWRS	RAGNKIAP	AAAAWPIIGH	LHLLAGGSHQ	60
LPHITLGNMA	DKYGPVFTIR	IGLHRAVVVS	SWEMAKECST	ANDQVSSSRP	ELLASKLLGY	120
NYAMFGFSPY	GSYWREMRKI	ISLELLNSR	LELLKDVRAS	EVVTSIKELY	KLWAEKKNES	180
GLVSVEMKQW	FGDLTLNVIL	RMVAGKRYFS	ASDASENKQA	QRCRRVREF	FHLSGLFVVA	240
DAIPFLGWLD	WGRHEKTLK	TAIEMDSIAQ	EWLEEHRRR	DSGDDNSTQD	FMDVMQSVLD	300
GKNLGGYDAD	TINKATCLTL	ISGGSPTTVV	SLTWALSIVL	NNRDTLKKQA	EELDIQVQGE	360
RLVNEQDISK	LVYLQAIVKE	TLRLYPPGPL	GGLRQFTEDC	TLGGYHVSKG	TRLIMNLSKI	420
QKDPRIWSDP	TEFQPERFLT	THKDVDPGRK	HFEFIPFGAG	RRACPGITFG	LQVLHLTLAS	480
FLHAFEFSTP	SNEQVMRES	LGLTNMKSTP	LEVLLSPRLS	SCSLYN		526

SEQ ID NO:17

MEPNFYLSLL	LLFVTFISLS	LFFIFYKQKS	PLNLPPGKMG	YPIIGESLEF	LSTGWKGHPE	60
KFIFDRMRKY	SSELEKTSIV	GESTVCCGA	ASNKFLFSNE	NKLVTAWWD	SVNKIFPTTS	120
LDSNLKEESI	KMRKLLPQFF	KPEALQRYVG	VMDVIAQRHF	VTHWDNKNEI	TVYPLAKRYT	180
FLLACRLFMS	VEDENHVAKF	SDPFQLIAG	IISLPIDLP	TPFNKAIKAS	NFIRKELIKI	240
IKQRRVDLAE	GTASPTQDIL	SHMLLTSDEN	GKSMNELNIA	DKILGLLIG	HDTASVACTF	300
LVKYLGELEPH	IYDKVYQEQM	EIAKSKPAGE	LLNWDLLKMM	KYSWNVACEV	MRLSPPLQGG	360
FREAITDFMF	NGFSIPKQWK	LYWSANSTHK	NAECFFMPEK	FDPTRFEGNG	PAPYTFVFPFG	420
GGPRMCPGKE	YARLEILVEM	HNLVKRKFKE	KVIPDEKIIV	DPFPIPAKDL	PIRLYPHKA	479

SEQ ID NO:18

atggaagcct	cttaccctata	catttctatt	ttgcttttac	tggcatcata	cctgttcacc	60
actcaactta	gaaggaagag	cgctaactta	ccaccaaccg	tggttccatc	aataccaatc	120
attggacact	tatacttact	caaaaagcct	ctttatagaa	ctttagcaaa	aattgcccgt	180
aagtacggac	caactactgca	attacaactc	ggctacagac	gtgttctggt	gatttctcctca	240
ccatcagcag	cagaagagtg	ctttaccaat	aacgatgtaa	tcttcgcaaa	tagacctaac	300

acattgtttg	gcaaaatagt	gggtggaaca	tcccttgcca	gtttatccta	cggcgatcaa	360
tgccgtaatc	taaggagagt	agcttctatc	gaaatcctat	cagttcatag	gttgaacgaa	420
tttcatgata	tcagagtggg	tgagaacaga	ttgttaatta	gaaaacttag	aagttcatct	480
tctcctgtta	ctcttataac	agtcttttat	gctctaacat	tgaacgtcat	tatgagaatg	540
atctctggca	aaagatattt	cgacagtggg	gatagagaat	tggaggagga	aggtaagaga	600
tttcgagaaa	tcttagacga	aacgttgctt	ctagccggtg	cttctaagt	tggcgactac	660
ttaccaatat	tgaactgggt	gggagttaag	tctctgaaa	agaaattgat	cgctttgcag	720
aaaaagagag	atgacttttt	ccagggtttg	attgaacag	ttagaaaatc	tcgtggtgct	780
aaagtaggca	aaggtagaaa	aacgatgac	gaactcttat	tatctttgca	agagtcagaa	840
cctgagtact	atacagatgc	tatgataaga	tcttttgctc	taggtctgct	ggctgcaggt	900
agtgataact	cagcgggcac	tatggaatgg	gccatgagct	tactggtcaa	tcaccacat	960
gtattgaaga	aagctcaagc	tgaaatcgat	agagttatcg	gtaataacag	attgattgac	1020
gagtcagaca	ttggaaatat	cccttacatc	gggtgtatta	tcaatgaaac	tctaagactc	1080
tatccagcag	ggccagatgc	gttcccacat	gaaagttctg	ccgactgctg	tatttccggt	1140
tacaatatac	ctagaggtac	aatgttaatc	gtaaaccaat	ggcgattca	tcacgatcct	1200
aaagtctggg	atgatcctga	aacctttaa	cctgaaagat	ttcaaggatt	agaaggaact	1260
agagatggtt	tcaaaccttat	gccattcggg	tctgggagaa	gaggatgtcc	aggatgaagg	1320
ttggcaataa	ggctgttagg	gatgacacta	ggctcagtga	tccaatgttt	tgattgggag	1380
agagtaggag	atgagatggt	tgacatgaca	gaaggtttgg	gtgtcacact	tcctaaggcc	1440
gttccattag	ttgccaaatg	taagccacgt	tccgaaatga	ctaactctct	atccgaactt	1500
taa						1503

SEQ ID NO:19

MEASYLYISI	LLLLASYLFT	TQLRRKSANL	PPTVFPSIPI	IGHLYLLKKP	LYRTLAKIAA	60
KYGPILQLQL	GYRRVLVISS	PSAABECFTN	NDVIFANRPK	TLFGKIVGGT	SLGSLSYGDQ	120
WRNLRRVASI	EILSVHRLNE	FHDIRVDENR	LLIRKLRSSS	SPVTLITVYF	ALTLNVIMRM	180
ISGKRYFDSG	DRELEEEGKR	FREILDETL	LAGASVNGDY	LPILNWLGVK	SLEKLIALQ	240
KKRDDFFQGL	IEQVRKSRGA	KVGKGRKTM	ELLSLQESE	PEYYTDAMIR	SFVLGLLAAG	300
SDTSAGTMEW	AMSLLVNHPH	VLKKAQAEID	RVIGNRLID	ESDIGNIPYI	GCIINETLRL	360
YPAGPLLFPH	ESSADCVISG	YNIPRGTMLI	VNQWAIHHPD	KVWDDPETFK	PERFQGLEGT	420
RDGFKLMPFG	SRRGCPGEG	LAIRLLGMTL	GSVIQCFDWE	RVGDEMVDMT	EGLGVTLPKA	480
VPLVAKCKPR	SEMTNLLSEL					500

SEQ ID NO:20

MQSDSVKVSP	FDLVSAMNG	KAMEKLNASE	SEDPTLPAL	KMLVENRELL	TLFTTSFAVL	60
IGCLVFLMWR	RSSSKKLVDQ	PVPQVIVVKK	KEKESEVDDG	KKKVSIFYGT	QTGTAEGFAK	120
ALVEEAKVRY	EKTSFKVIDL	DDYAADDDEY	EELKKESIA	FFFLATYGDG	EPTDNAANFY	180
KWFTEGDDKG	EWLKKLQYGV	FGLGNRQYEH	FNKIAIVVDD	KLTEMGAKRL	VPVGLGDDDQ	240
CIEDDFTAWK	ELVWPELDQL	LRDEDDTSVT	TPYTAAVLEY	RVVYHDKPAD	SYAEDQHTN	300
GHVVDHAQHP	SRSNVAFKKE	LHTSQSDRSC	THLEFDISHT	GLSYETGDHV	GVYSENLSV	360
VDEALKLLGL	SPDITYFSVHA	DKEDGTPIGG	ASLPPFPFPC	TLRDALTRYA	DVLSSPKKVA	420
LLALAAHASD	PSEADRLKFL	ASPAGKDEYA	QWIVANQRSL	LEVMSQFSPA	KPPLGVFFAA	480
VAPRLQPRYY	SISSSPKMSP	NRHVTCALV	YETTPAGRIH	RGLCSTWMKN	AVPLTESPDC	540
SQASIFVRTS	NFRLPVPDPK	PVIMIGPGTG	LAPFRGFLOE	RLALKESGTE	LGSSIFFFGC	600
RNRKVDFIYE	DELNNFVETG	ALSELIVAFS	REGTAKEYVQ	HKMSQKASDI	WKLLSEGAYL	660
YVCGDAKGMA	KDVHRTLHTI	VQEQQSLDSS	KAELYVKNLQ	MSGRYLRD VW		710

SEQ ID NO:21

MTSALYASDL	FKQLKSIMGT	DSLSDDVVLV	IATTSALVA	GFVLLWKKT	TADRSGELKP	60
LMIPKSLMAK	DEDDDLGLS	GKTRVSIFFG	TQTGTAEGFA	KALSEEIKAR	YEKAAVKVID	120
LDDYAADDQ	YEEKLKKETL	AFFCVATYGD	GEPTDNAARF	YKWFTEENER	DIKLQQLAYG	180
VFALGNRQYE	HFNKIGIVLD	EELCKKGAKR	LIEVGLGDDD	QSIEDDFNAW	KESLWSELDK	240
LLKDEDDKSV	ATPYTAVIPE	YRVVTHDPRF	TQOKSMESNV	ANGNTTIDIH	HPCRVDVAVQ	300
KELHTHESDR	SCIHFLEFDS	RTGITYETGD	HVGVYAENHV	EIVEEAGKLL	GHSIDLVSFI	360
HADKEDGSPL	ESAVPPPFG	PCTLGTGLAR	YADLLNPRK	SALVALAAYA	TEPSEAEKLN	420
HLTSPDGKDE	YSQWIVASQR	SILEVMAAFP	SAKPPLVVFF	AAIAPRLQPR	YYSISSPRL	480
APSRVHVTS	LVYGPPTGR	IHKGVCSTWM	KNAPVPAEKSH	ECSGAPIFIR	ASNFKLPSNP	540
STPIVMVGP	TGLAPFRGFL	QERMALKEDE	EELGSSLLFF	GCRNRQMDFI	YEDELNNFVD	600
QGVISELIMA	FSREGAQKEY	VQHMMKEAA	QWDLIKEEG	YLYVCGDAKG	MARDVHRTLH	660
TIVQEQEGVS	SSEAEAIVKK	LQTEGRYLDR	VW			692

SEQ ID NO:22

MAELDTLIDIV	VLGVIFLGTV	AYFTKGKLGW	VTKDPYANGF	AAGGASKPGR	TRNIVEAMEE	60
SGKNCVVFYG	SQTGTAEDYA	SRLAKEGKSR	FGLNTMIADL	EDYDFDNLDT	VPSDNIVMFV	120
LATYGEGETP	DNAVDFYEFT	TGEDASFNEG	NDPPLGNLNY	VAFGLGNNTY	EHYNSMVRNV	180
NKALEKLGAH	RIGEAGEGDD	GAGTMEEDFL	AWKDPMWEAL	AKKMGLEERE	AVYEPIFAIN	240
ERDDLTPKAN	EVYLGEPNKL	HLEGTAKGPF	NSHNPIYAPI	AESYELFSAK	DRNCLHMEID	300
ISGSNLKYET	GDHIAIWPTN	PGEEVNKFLD	ILDLSGKQHS	VVTVKALEPT	AKVFFPNPTT	360
YDAILRYHLE	ICAPVSRQFV	STLAAFAPND	DIKAEMNRLG	SDKDYFHEKT	GPHYNYIARF	420
LASVSKGEKW	TKIPFSAFIE	GLTKLQPRYY	SISSSSLVQP	KKISITAVVE	SQIQPRDDP	480
FRGVATNYLF	ALKQKQNGDP	NPAPFGQSYE	LTGPRNKYDG	IHVPHVHRHS	NFKLPSPDPGK	540
PIIMIGPGTG	VAPFRGFVQE	RAKQARDGVE	VGKTLFFFGC	RKSTEDFMYQ	KEWQEYKEAL	600
GDKFEMITAF	SREGSKKVVY	QHRLKERSKE	VSDLLSQKAY	FYVCGDAAHM	AREVNTVLAQ	660
IIAEGRGVSE	AKGEEIVKNM	RSANQYQVCS	DFVTLHCKET	TYANSELQED	VWS	713

SEQ ID NO:23

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gcgatgatgt	tcgaaattcg	tgatctgttg	ctgattttga	ctacgtcagt	tgctgttttg	180
gtcggatggt	tcggtgtttt	gggtgtggaag	agatcgtccg	ggaagaagtc	cggcaaggaa	240
ttggagccgc	cgaagatcgt	tgtgccgaag	aggcggctgg	agcaggaggt	tgatgatggt	300
aagaagaagg	ttacgatatt	cttcggaaca	caaactggaa	cggtggaagg	tttcgctaag	360
gcacttttgc	aagaagcgaa	agcgcgatat	gaaaagcgag	cgtttaaggt	gattgatttg	420
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gcaccgcggt	tacagcctcg	ttactactct	atcttctcct	ccccaaagt	ggaaccaaac	1500
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ctttctgagc	ttgatgttgc	tttctcccgc	gatggcccga	cgaaagaata	cggtcaacat	1920
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SEQ ID NO:24

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aaagttacgg	ttttctctcg	caccocaaact	ggaacagctg	aaggcttcgc	taaggcactt	360
gttgaggaag	ctaaaagctcg	atatgaaaag	gctgtcttta	aagtaattga	tttgatgat	420
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ggtgatgcca	aaggcatggc	caaagatgta	catcgaaccc	tccacacaat	tgtgcaagaa	2040
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agatacctcc	gtgacgtttg	gtaa				2124

SEQ ID NO:25

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LMIPKSLMAK	DEDDDLDLGS	GKTRVSIFFG	TQTGTAEFGA	KALSEEIKAR	YEKAAVKVID	120
LDYYAADDQ	YEEKLKKETL	AFFCVATYGD	GEPTDNAARE	YKWFTEENER	DIKLQQLAYG	180
VFALGNRQYE	HFNKIGIVLD	EELCKKGAKR	LIEVGLGDDD	QSIEDDFNAW	KESLWSELDK	240
LLKDEDDKSV	ATPYTAVIPE	YRVVTHDPRF	TTQKSMESNV	ANGNTTIDIH	HPCRVDDVAVQ	300
KELHTHESDR	SCIHLEFDIS	RTGITYETGD	HVGVYAENHV	EIVEEAGKLL	GHSLDLVFSI	360
HADKEDGSPL	ESAVPPPPFG	PCTLGTGLAR	YADLLNPPRK	SALVALAAYA	TEPSEAEKLL	420
HLTSPDGKDE	YSQWIVASQR	SLEEVMAAFP	SAKPLGVVFF	AAIAPRLQPR	YYSISSPRL	480
APSRVHVTSA	LVYGPTPTGR	IHKGVCSTWM	KNAVPAEKSH	ECSGAPIFIR	ASNFKLPSNP	540
STPIVMGPG	TGLAPFRGFL	QERMALKEDG	EELGSSLLFF	GCRNRQMDFI	YEDELNNFVD	600
QGVISELIMA	FSREGAQKEY	VQHKMMEKAA	QVWDLIKEEG	YLYVCGDAKG	MARDVHRTLH	660
TIVQEQEGVS	SSEAEAI VKK	LQTEGRYL RD	VW			692

SEQ ID NO:26

MSSSSSSSTS	MIDLMAAIK	GEPVIVSDPA	NASAYESVAA	ELSSMLIENR	QFAMIVTTSI	60
AVLIGCI VML	VWRRSGSGNS	KRVEPLKPLV	IKPREEEIDD	GRKKVTIFFG	TQTGTAEFGA	120
KALGEEAKAR	YEKTRFKIVD	LDYYAADDDE	YEEKLKKEDV	AFFFLATYGD	GEPTDNAARE	180
YKWFTEGNDR	GEWLKNLKYG	VFGLGNRQYE	HFNKVAKVVD	DILVEQGAQR	LVQVGLGDDD	240
QCIEDDFTAW	REALWPELDT	ILREEGDTAV	ATPYTAAVLE	YRVSIHDSER	AKFNDITLAN	300
GNGYTVFDAQ	HPYKANVAVK	RELHTPESDR	SCIHLEFDIA	GSGLTMKLG	HVGVLCDNLS	360
ETVDEALRLL	DMSPTDYFSL	HAEKEDGTPI	SSSLPPPPFP	CNLRALTRY	ACLLSSPKKS	420
ALVALAAHAS	DPTAEARLKH	LASPAGKDEY	SKWVVESQRS	LLEVMAEFPS	AKPPLGVFFA	480
GVAPRLQPRF	YSTSSSPKIA	ETRIHVTCAL	VYEKMPTGRI	HKGVCSTWMK	NAVPEKSEK	540
LFLGRPIFVR	QSNFKLPSDS	KVPIIMIGPG	TGLAPFRGFL	QERLALVESG	VELGPSVLEFF	600
GCRNRMDFI	YEEELQRFVE	SGALAELSVA	FSREGPTKEY	VQHKMMDKAS	DIWNMISQGA	660
YLYVCGDAKG	MARDVHRS LH	TIAQEQQSMD	STKAEGFVKV	LQTSGRYL RD	VW	712

SEQ ID NO:27

MQSESEVAST	IDLMTAVLKD	TVIDTANASD	NGDSKMPPAL	AMMFEIRDLL	LILTTSVAVL	60
VGCFVVLVWK	RSSGKKSKE	LEPPKIVVPK	RRLEQEVDDG	KKKVTIFFGT	QGTAEFGFAK	120
ALFEEAKARY	EKAFAFKVIDL	DDYAADLDEY	AEKLLKETYA	FFFLATYGDG	EPTDNAAKFY	180
KWFTEGDEKG	VWLQKLQYGV	FGLGNRQYEH	FNKIGIVVDD	GLTEQGA KRI	VPVGLGDDDQ	240
SIEDDFS AWK	ELVWPEL DLL	LRDEDDKAAA	TPYTA AIPEY	RVVFHDKPDA	FSDDHTQTNG	300

HAVHDAQHPC	RSNVAVKKEL	HTPESDRSCT	HLEFDISHTG	LSYETGDHVG	VYCENLIEVV	360
EEAGKLLGLS	TDTYFSLHID	NEDGSPGGP	SLQPPFPCT	LRKALTNYAD	LLSSPKKSTL	420
LALAAHASDP	TEADRLRFLA	SREGKDEYAE	WVVANQRSLL	EVMEAFPSAR	PPLGVFFAAV	480
APRLQPRYS	ISSSPKMEPN	RIHVTCALVY	EKTPAGRIHK	GICSTWMKNA	VPLTESQDCS	540
WAPIFVRTSN	FRLPIDPKVP	VIMIGPGTGL	APFRGFLQER	LALKESGTEL	GSSILFFGCR	600
NRKVDIYIEN	ELNNFVENGA	LSELDVAFSR	DGPTKEYVQH	KMTQKASEIW	NMLSEGAYLY	660
VCGDAKGMAK	DVHRTLHTIV	QEQQSLDSSK	AELYVKNLQM	SGRYLRDVM		709

SEQ ID NO:28

MQNSSVKISP	LDLVTALFSG	KVLDTSNASE	SGESAMLPIT	AMIMENRELL	MILTTSVAVL	60
IGCVVVLVWR	RSSTKKSAL	PPVIVPKRV	QEEVDDGKK	KVTVFFGTQT	GTAEGFAKAL	120
VEEAKARYEK	AVFKVIDLDD	YAADDDEYEE	KLKKESLAFF	FLATYGDGEP	TDNAARFYKW	180
FTEGDAKGEW	LNKLOYGVFG	LGNRQYEHFN	KIAKVVDGGL	VEQGAKRLVP	VGLGDDDDQCI	240
EDDFTAWKEL	VWPELDQLLR	DEDDTTVATP	YTAAVAEYRV	VFHEKPDALS	EDYSYTNNGHA	300
VHDAQHPCRS	NVAVKKELHS	PESDRSCTHL	EFDISNTGLS	YETGDHVGTV	CENLSEVVND	360
AERLVGLPPD	TYSSIHTDSE	DGSPLGGASL	PPFPCTLR	KALTCYADVL	SSPKKSALLA	420
LAHAHTDPSE	ADRLKFLASP	AGKDEYSQWI	VASQRSLEEV	MEAFPSAKPS	LGVFFASVAP	480
RLQPRYSIS	SSPKMAPDRI	HVTCALVYEK	TPAGRIHKGV	CSTWMKNAVP	MTESQDCSWA	540
PIYVRTSNFR	LPSDPKVPVI	MIGPGTGLAP	FRGFLQERLA	LKEAGTDLGL	SILFFGCRNR	600
KVDFIYENEL	NNFVETGALS	ELIVAFSREG	PTKEYVQHKM	SEKASDIWNL	LSEGAYLYVC	660
GDARGMKAVD	HRTLHTIVQE	QGSLLDSSKAE	LYVKNLQMSG	RYLRDVM		707

SEQ ID NO:29

MAEQKIKKKS	PHVLLIPFPL	QGHINFFIQF	GKRLISKGVK	TTLVTTIHTL	NSTLNHSNTT	60
TTSIEIQAIS	DGCEGGFMS	AGESYLETFK	QVGSKSLADL	IKKLQSEGTT	IDAIYDSMT	120
EWVLDVAIEF	GIDGGSFFTQ	ACVVNSLYYH	VHKGLISLPL	GETVSVPGFP	VLQRWETPLI	180
LQNHQIQSP	WSQMLFGQFA	NIDQARWVFT	NSFYKLEEV	IEWTRKIWNL	KVIGPTLPSM	240
YLDKRLDDDK	DNGFNLYKAN	HHECMNWLDD	KPKESVVYVA	FGSLVKHGPE	QVEEITRALI	300
DSDVNFVWLI	KHKEEGKLE	NLSEVIKTGK	GLIVAWCKQL	DVLAHESVGC	FVTHCGFNST	360
LEAISLGPVP	VAMPQFSDQT	TNAKLLDEIL	GVGVRVKADE	NGIVRRGNLA	SCIKMIMEEE	420
RGVIRKNAV	KWKDLAKVAV	HEGSSDNDI	VEFVSELIKA			460

SEQ ID NO:30

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CLDGAPGFRF	ETIPDGVSHS	PEASIPIRES	LLRSIETNFL	DRFIDLVTKL	PDPPTCIISD	120
GFLSVFTIDA	AKKLGIPVMM	YWTLAACGFM	GFYHIHSLIE	KGFAPLKDAS	YLTNGYLDTV	180
IDWVPGMEGI	RLKDFPLDWS	TDLNDKVLMF	TTEAPQRSHK	VSHHI PHTFD	ELEPSIKTL	240
SLRYNHIYTI	GPLQLLLDQI	PEEKQTGIT	SLHGYSLVKE	EPECFQWLQS	KEPNSVVYVN	300
FGSTTVMSLE	DMTEFGWGLA	NSNHYFLWII	RSNLVIGENA	VLPPELEEHI	KKRGFIASWC	360
SQEKVLKHP	VGGFLTHCGW	GSTIESLSAG	VPMICWPYSW	DQLTNCRYIC	KEWEVGLEMG	420
TKVKRDEVKR	LVQELMGE	HKMRNKAKDW	KEKARIAIAP	NGSSSLNIDK	MVKEITVLAR	480
						481

SEQ ID NO:31

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cccgaagaga	aaaagcaaac	tggaaattacg	agtctccatg	gatacagttt	agtaaaagaa	840
gaaccagagt	gtttccagtg	gcttcagttc	aaagaaccaa	atctccgtcg	ttatgtaaat	900
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tcacaagaaa	aggtcttgaa	gcacccttcg	gttggagggt	tcttgactca	ttgtgggtgg	1140
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SEQ ID NO:32

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ataactttcg	tgaataccga	cttcatccat	aatcaatttc	tggaatctag	tggccctcat	180
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attgactggg	taccaggata	ggaaggata	agacttaaag	atctcctttt	ggatgggtct	600
acagacctta	atgataaagt	attgatgttt	actacagaag	ctccacaaag	atctcataag	660
gtttcacatc	atatctttca	cacctttgat	gaattggaac	catcaatcat	caaaaccttg	720
tctctaagat	acaatcataat	ctacactatt	ggtccattac	aattacttct	agatcaaaat	780
cctgaagaga	aaaagcaaac	tggtattaca	tccttacacg	gctactcttt	agtgaagag	840
gaaccagaat	gttttcaatg	gctacaaaagt	aaagagccta	atctctgtgt	ctacgtcaac	900
ttcggaagta	caacagtc	gtccttggaa	gatatgactg	aatttgggtg	gggccttgct	960
aattcaaatc	attactttct	atggattatc	aggccaatt	tggaatagg	ggaaaacgcc	1020
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acaaaggtta	aacgtgatga	agtgaaaaga	ttggttcagg	agttgatggg	ggaaggtggc	1320
cacaagatga	gaaacaaggc	caaagattgg	aaggaaaaag	ccagaattgc	tattgctcct	1380
aacgggtcat	cctctctaaa	cattgataag	atggtcaaag	agattacagt	cttagccaga	1440
aactaa						1446

SEQ ID NO:33

MKTGFISPAT	VFHHRISPAT	TFRHLSPAT	TNSTGIVALR	DINFRCKAVS	KEYSDLLQKD	60
EASFTKWDD	KVKDHLDTNK	NLYPNDEIKE	FVESVKAMFG	SMNDGEINVS	AYDTAWVALV	120
QDVVDSGSPQ	FPSSLEWIAN	NQLSDGSWGD	HLLFSAHDRI	INTLACVIAL	TSWNVHPSKC	180
EKGLNFLREN	ICKLEDENAE	HMPIGFEVTF	PSLIDIAKKL	NIEVPEDTPA	LKEIYARRDI	240
KLTKIPMEVL	HKVPTTLHS	LEGMPDLEWE	KLLKLQCKDG	SPLFSPSSTA	FALMQTKDEK	300
CLQYLNTIVT	KFNGGVPNVY	PVDLFEHIWV	VDRLQRLGIA	RYFKSEIKDC	VEYINKYWTK	360
NGICWARNT	VQDIDDTAMG	FRVLRAHGYD	VTPDVFRQFE	KDGKFCVCFAG	QSTQAVTGMF	420
NVYRASQLMF	PGERILEDK	KFSYNYLKEK	QSTNELLDKW	IIAKDLPGEV	GYALDIPWYA	480
SLPRLETRY	LEQYGGEDDV	WIGKTLYRMG	YVSNNTYLEM	AKLDYNNYVA	VLQLEWYTIQ	540
QWYVDIGIEK	FESDNIKSVL	VSYYLAAASI	FEPERSKERI	AWAKTTILVD	KITSIFDSSQ	600
SSKEDITAFI	DKFRNKSSSK	KHSINGEPWH	EVMVALKKT	HGFALDALMT	HSQDIHPQLH	660
QAWEMWLTKL	QDGVDTAEL	MVQMINMTAG	RWVSKELLTH	PQYQLSTVT	NSVCHDITKL	720
HMFKENSTTV	DSKVQELVQL	VESDTPDDL	QDMKQFTLV	MKTFYKAWC	DPNTINDHIS	780
KVFEIVI						787

SEQ ID NO:34

MPDAHDAAPP	QIRQRTLVE	ATQLLTSAE	DAWGEVSVSE	YETARLVAHA	TWLGGHATRV	60
AFLERQHED	GSWGPPGGYR	LVPTLSAVHA	LLTCLASPAQ	DHGVPHDRLL	RAVDAGLTAL	120
RRLGTSDSP	DTIAVELVIP	SLEGIQHL	DPAPHRSRPA	FSQHRGSLVC	PGGLDGRITG	180
ALRSHAAAGT	PVPGKVWHAS	ETLGLSTEAA	SHLQPAQGI	GGSAATATW	LTRVAPSQQS	240
DSARRYLEEL	QHRYSQVPS	ITPITYFERA	WLLNNFAAAG	VPCEAPAALL	DSLEAALTPQ	300
GAPAGAGLPP	DADDTAAVLL	ALATHGRGR	PEVLMDYRTD	GYFQCFIGER	TPSISTNAHV	360

LETLGHVHAQ	HPQDRARYGS	AMDTASAWLL	AAQKQDGSWL	DKWHASPYYA	TVCCTQALAA	420
HASPATAPAR	QRAVRWVLAT	QRSDDGGWGLW	HSTVEETAYA	LQILAPPSGG	GNIPIVQQALT	480
RGRARLCGAL	PLTFLWHDKD	LYTPVRVVR	ARAAALYTTR	DLLLPL		527

SEQ ID NO:35

MNALSEHILS	ELRRLLEMS	DGGSVGPSVY	DTAQUALRFHG	NVTGRQDAYA	WLIAQQQADG	60
GWGSADFFLF	RHAPTWAALL	ALQRADPLPG	AADAVQTATR	FLQRQPDPIYA	HAVPEDAPIG	120
AELILPQFCG	EAAWLLGGVA	FPRHPALLPL	RQACLVLKGA	VAMLPSPGHP	LHSWEAWGTS	180
PTTACPDG	SIGISPAATA	AWRAQAVTRG	STPQVGRADA	YLOMASRATR	SGIEGVFPNV	240
WPINVFEP	SLYTLHLAGL	FAHPALAEAV	RVIVAQLEAR	LGVHGLGPAL	HFAADADDTA	300
VALCVLHLAG	RDPAVDALRH	FEIGELFVTF	PGERNASVST	NIHALHALRL	LGKPAAGASA	360
YVEANRNP	LWDNEKWHVS	WLYPTAHAVA	ALAQQKPQWR	DERALAALLQ	AQRDDGGWGA	420
GRGSTFEETA	YALFALHVMD	GSEETGRRR	IAQVVARALE	WMLARHAAHG	LPQTPLWIGK	480
ELYCPTRVVR	VAELAGLWLA	LRWRRVLAE	GAGAAP			516

SEQ ID NO:36

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cgagtattat	agtagtagat	ctcttctccg	atataatccg	ccaaaggag	aagagaagag	660
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aaagttagct	ctgttttggg	taaaaaaat	ccagtttctg	taaattatag	aataaatcaa	4500
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acctatcttg	tgatgac					4577

SEQ ID NO:37

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AKGSSLTPIV	RTDAESRRTR	WPTDDDDAEP	LVDEIRAMLT	SMSDGDIVS	AYDTAWVGLV	120
PRLDGGEGPQ	FPAAVRWIRN	NQLPDGSGWD	AALFSAYDRL	INTLACVVTL	TRWSLEPEMR	180
GRGLSFLGRN	MWKLATEDEE	SMPIGFELAF	PSLIELAKSL	GVHDFPYDHO	ALQGIYSSRE	240
IKMKRIPEV	MHTVPTSILH	SLEGMPGLDW	AKLLKLQSSD	GSFLFSPAAT	AYALMNTGDD	300
RCFSYIDRVT	KKFNGGVPNV	YPVDLFEHIW	AVDRLERLGI	SRYFQKEIEQ	CMDYVNRHWT	360
EDGICWARN	DVKEVDDTAM	AFRLRLRHGY	SVSPDVFKNF	EKDGEFFAEV	GQSNQAVTGM	420
YNLNRSQIS	FPGEDVLHRA	GAFSYEFLRR	KEAEGALRDK	WIISKDLPE	VVYTLDFPWY	480
GNLPRVEARD	YLEQYGGDD	VWIGKTLYRM	PLVNDVYLE	LARMDFNHCQ	ALHQLEWQGL	540
KRWYTENRLM	DFGVAQEDAL	RAYFLAAASV	YEPCAAERL	AWARAAILAN	AVSTHLRNSP	600
SFRERLEHSL	RCPSEETDM	SWFNSSSGSD	AVLVKAVLRL	TDSLAREAQP	IHGDPEDII	660
HKLLRSAAWE	WREKADAAD	SVNCGSSAVE	QEGSRMVHDK	QTCILLLARI	EISAGRAAGE	720
AASEDGDRRI	IQLTGSICDS	LKQKMLVSQD	PEKNEEMSH	VDELKLRIR	EFVQYLLRLG	780
EKKTGSSETR	QTFLSIVKSC	YAAHCPPHV	VDRHISRIVF	EPVSAAK		827

SEQ ID NO:38

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SEQ ID NO:39

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SQEVQHDLLP	IHEWQQLQGE	DAPQISVGSN	SNAFKEAVKS	VKTILRNLTG	GEITISAYDT	120
AWVALIDAGD	KTPAFPSAVK	WIAENQLSDG	SWGDAYLFSY	HDRLINTLAC	VVALRSWNLF	180
PHQCNKGITF	FRENIGKLED	ENDEHMPIGF	EVAFPSLLEI	ARGINIDVPY	DSPVLKDIYA	240
KKELKLRTRIP	KEIMHKIPTT	LLHSLEGMRD	LDWEKLLKLQ	SQDGSFLFSP	SSTAFAFMQT	300
RDSNACLEYL	NAVKRENGGV	PNVFPVDLFE	HIWIVDLRQR	LGISRYFEEE	IKECLDYVHR	360
YWTDNGICWA	RCSHVQDIDD	TAMAFRLLRQ	HGYQVSADVF	KNFEKEGEFF	CFVQGSNQAV	420
TGMFNLYRAS	QLAFPREEIL	KNAKEFSYNY	LLEKREREEL	IDKWIIMKDL	PGEIGFALEI	480
PWYASLPRVE	TRFYIDQYGG	ENDVWIGKTL	YRMPYVNNNG	YLELAKQDYN	NCQAQHOLEW	540
DIFQKWYEEN	RLSEWGVRRS	ELLECYYLAA	ATIFESERSH	ERMVWAKSSV	LVKAISSSFG	600
ESSDSRRSFS	DQFHEYIANA	RRSDHHFNDR	NMRLDRPGSV	QASRLAGVLI	GTLNQMSFDL	660
FMSHGRDVNN	LLYLSWGDWM	EKWLYGDEG	EGELMVKMII	LMKNNDLTNF	FTHTHFVRLA	720
EIINRICLPR	QYLKARRNDE	KEKTIKSMEK	EMGRMVELAL	SESDFTRDVS	ITFLDVAKAF	780
YYFALCGDHL	QTHISKVLFQ	KV				802

SEQ ID NO:40

MEFDEPLVDE	ARSLVQRTLQ	DYDDRYGFGT	MSCAAYDTAW	VSLVTKTVDG	RKQWLFPECF	60
EFLLTQSDA	GGWEIGNSAP	IDGILNTAAS	LLALKRHHVQT	EQIIQPHDH	KDLAGRAERA	120
AASLRAQLAA	LDVSTTEHVG	FEIIVPAMLG	PLEAEDPSLV	FDFPARKPLM	KIHDAKMSRF	180
RPEYLYGKQP	MTALHSLEAF	IGKIDFDKVR	HHRTHGSMMG	SPSSTAAYLM	HASQWGDGSE	240
AYLRHVVIKHA	AGQGTGAVPS	AFPSTHFESS	WILTTLFRAG	FSASHLACDE	LNKLVEILEG	300
SFEKEGGAIG	YAPGFQADV	DTAKTISTLA	VLGRDATPRQ	MIKVEANTH	FRTYPGERDP	360
SLTANCNALS	ALLHQPDAAM	YGSQIQKIKT	FVCDYWWKSD	GKIKDKWNTC	YLYPSVLLVE	420
VLVDLVSLLE	QGKLPDVLQ	ELQYRVAITL	FQACLRPLLD	QDAEGSWNKS	IEATAYGILI	480
LTEARRVCFE	DRLSEPLNEA	IRRGIAFADS	MSGTEAQLNY	IWIEKVSYP	ALLTKSYLLA	540
ARWAAKPLG	ASVGSSSLWTP	PREGLDKHVR	LFHQAEFRS	LPEWELRASM	IEAALFTPLL	600
RAHRLDVFP	QDVGEDKYL	VVFFFWTAAN	NRDRTYASTL	FLYDMCFIAM	LNFLDEFME	660
ATAGILFRDH	MDDLRLIHD	LLAEKTSPKS	SGRSSQGTKD	ADSGIEEDVS	MSDSASDSQD	720
RSPEYDLVFS	ALSTFTKHVL	QHPSIQSASV	WDRKLLAREM	KAYLLAHIQQ	AEDSTPLSEL	780
KDVPQKTDVT	RVSTSTTTFF	NWVRTTSADH	ISCPYSFHFV	ACHLGAALSP	KGSNGDCYPS	840
AGEKFLAAAV	CRHLATMCRM	YNDLGS AERD	SDEGNLNSLD	FPEFADSAGN	GGIEIQKAAL	900

LRLAEFERDS	YLEAFRRLOD	ESNRVHGPAG	GDEARLSRRR	MAILEFFAQQ	VDLYGQVYVI	960
RDISARIPKN	EVEKRRKLD	AFN				983

SEQ ID NO:41

MASSTLIQNR	SCGVTSSMSS	FQIFRGQPLR	FPGTRTPAAV	QCLKRRRCLR	PTESVLESSP	60
GGGSYRIVTG	PSGINPSSNG	HLQEGSLTHR	LPIMPESID	NFQSTLYVSD	IWSETLQORTE	120
CLLQVTENVQ	MNEWIEEIRM	YFRNMTLGEI	SMSPYDTAWV	ARVPALDGSH	GPQFHRSLQW	180
IIDNQLPDGD	WGEPSLFLGY	DRVCNTLACV	IALKTWGVGA	QNVERGIQFL	QSNYKMEED	240
DANHMPIGFE	IVFPAMMEDA	KALGLDLPYD	ATILQQISAE	REKKMKKIPM	AMVYKYPTTL	300
LHSLEGLHRE	VDWNKLLQLQ	SENGSFLYSP	ASTACALMYT	KDVKCFDYLN	QLLIKFDHAC	360
PNVYPVDLFE	RLWMVDRLQR	LGISRYFERE	IRDCLQYVYR	YWKDCGIGWA	SNSSVQDVDD	420
TAMAFRLLRT	HGFDVKEDCF	RQFFKDGEFF	CFAGQSSQAV	TGMFNLSRAS	QTLFPGESLL	480
KKARTFSRNF	LRTKHENNEC	FDKWIITKDL	AGEVEYNLTF	PWYASLPRLE	HRTYLDQYGI	540
DDIWIGKSLY	KMPAVTNEVF	LKLAKADFNM	CQALHKKELE	QVIKWNASCO	FRDLEFARQK	600
SVECYFAGAA	TMFEPEMVA	RLVWARCCVL	TTVLDDYFDH	GTPVEELRVF	VQAVRTWNPE	660
LINGLPEQAK	ILAFVAFMA	NTIAEEAFMA	QKRVDVHHHLK	HYWDKLITSA	LKEAEWAESG	720
YVPTFDEYME	VAEISVALEP	IVCSTLFFAG	HRLEDVLDL	YDYHLMVHLV	NRVGRILNDI	780
QGMKREASQG	KISSVQIYME	EHPSVPSEAM	AIAHLQELVD	NSMQQLTYEV	LRFTAVPKSC	840
KRIHLNMAKI	MHAIFYKDTDG	FSSLTAMTGF	VKKVLFEPVP	E		881

SEQ ID NO:42

MPGKIENGTP	KDLKTGNDFV	SAAKSLLDRA	FKSHHSYGYL	CSTSCQVYDT	AWVAMIPKTR	60
DNVKQWLFPE	CFHYLLKTA	ADGSWGSPLT	TQTAGILDTA	SAVLALLCHA	QEPLQILDVS	120
PDEMGLRIEH	GVTSLKRQLA	VWNVEDTNH	IGVEFIIPAL	LSMLEKELDV	PSFEFPCRSI	180
LERMHGKELG	HFDLEQVYVK	PSSLLHSLEA	FLGKLDLDFRL	SHHLYHGSM	ASPSTAAYL	240
IGATKWDEEA	EDYLRHVMRN	GAGHGNGGIS	GTFPTTHFEC	SWIIATLLKV	GFTLQKIDGD	300
GLRGLSTILL	EALRDENGVY	GFAPTADVD	DTAKALLALS	LVNQPVSPDI	MIKVFEGKDH	360
FTTFGSRDP	SLTSLNLVLL	SLLKQSNLSQ	YHPQILKTTL	FTCRWWGSD	HCVKDKWNLS	420
HLYPTMLLVE	AFTEVLHLID	GSELSSLFDE	SFKCKIGLSI	FQAVLRRIIL	QDNDGSRGY	480
REQTCYAILA	LQARHVCFF	THMVDRLQSC	VDRGFSWLKS	CSFHSQDLTW	TSKTAYEVGF	540
VAEAYKLAAL	QASASLEVPAA	TIGHSVTSAV	PSSDLEKYMR	LVRKTALFSP	LDEWGLMASI	600
IESSFFVPLL	QAQRVEIYPR	DNKIVDEDKY	LSIIPFTWVG	CNNRSRTFAS	NRWLYDMMYL	660
SLLGYQTDEY	MEAVAGPVFG	DVSLHQITID	KVIDNTMGNL	ARANGTVHSG	NGHQHESPNI	720
GQVEDTLTRF	TNSVLNHKDV	LNSSSSDQDT	LRREFRTFMH	AHITQIEDNS	RFSKQASSDA	780
FSSPEQSYFQ	WVNSTGGSHV	ACAYSFAFSN	CLMSANLLQG	KDAFPSTGQK	YLISVVMRHA	840
TNMCRMYNDF	GSARDNAER	NVNSIHPEF	TLCNGTSQNL	DERKERLLKI	ATYEQGYLDR	900
ALEALERQSR	DDAGDRAGSK	DMRKLKIVKL	FCDVTDLYDQ	LYVIKDLSSS	MK	952

SEQ ID NO:43

MALVNPTALF	YGTSIRTRPT	NLLNPTQKLR	PVSSSSLPSF	SSVSAILTEK	HQSNPSENNN	60
LQTHLETPFN	FDSYMLEKVN	MVNEALDASV	PLKDPIKIHE	SMRYSLLAGG	KRIRPMMCIA	120
ACEIVGGNIL	NAMPAACAVE	MIHTMSLVHD	DLPCMNDNF	RRGKPISHKV	YGEEMAVLTG	180
DALLSLSFEH	IATATKGVSK	DRIVRAIGEL	ARSVGSEGLV	AGQVVDILSE	GADVGLDHLE	240
YIHIHKTAML	LESSVVICAI	MGGGSDQIE	KLRKFARSIG	LLFQVVDIIL	DVTKSTEELG	300
KTAGKDLLTD	KTTYPKLGI	EKSREFAEKL	NKEAQEQLSG	FDRRKAAPLI	ALANYNAYRQ	360
N						361

SEQ ID NO:44

MAEQQISNLL	SMFDASHASQ	KLEITVQMD	TYHYRETTPD	SSSSEGGSL	RYDERRVSLP	60
LSHNAASPD	VSQLCFSTAM	SSELNHRWKS	QRLKVADSPY	NYILTLPSKG	IRGAFIDSLN	120
VWLEVPEDET	SVIKEVIGML	HNSSLIIDDF	QDNSPLRRGK	PSTHTVFGPA	QAINATATYVI	180
VKAIEKIQDI	VGHDALADVT	GTITTFQGG	AMDLWWTANA	IVPSIQEYLL	MVNDKTGALF	240
RLSLELLALN	SEASISDSAL	ESLSSAVSLL	GQYFQIRDDY	MNLIDNKYTD	QKGFCELDLDE	300
GKYSLLTIHA	LQTDSSDLLT	NILSMRRVQG	KLTAQKRCWF	WK		342

SEQ ID NO:45

MEKTKEKAER	ILLEPYRYLL	QLPGKQVRSK	LSQAFNHWLK	VPEDKLQIII	EVTEMLHNAS	
LLIDDIEDSS	KLRGFPVAH	SIYGVPSVIN	SANYVYFLGL	EKVLTLDHDP	AVKLFTRQLL	

ELHQGGGLDI	YWRDITYCPT	EEYKAMVLQ	KTGGLFGLAV	GLMQLFSDYK	EDLKPLLDLTL	180
GLFFQIRDDY	ANLHSKEYSE	NKSFCEDLTE	GKFSFPTIHA	IWSRPESTQV	QNILRQRTEEN	240
IDIKKYCVQY	LEDVGSFAYT	RHTLRELEAK	AYKQIEACGG	NPSLVALVKH	LSKMFTEENK	300

SEQ ID NO:46

MARFYFLNAL	LMVISLQSTT	AFTPAKLAYP	TTTTALNVAS	AETSFSLDEY	LASKIGPIES	60
ALEASVKSRI	PQTDKICESM	AYSLMAGGKR	IRPVLICIAAC	EMFGGSQDVA	MPTAVALEMI	120
HTMSLIHDDL	PSMDNDDLRR	GKPTNHVVFV	EDVAILAGDS	LLSTSFHVA	RETKGVSAEK	180
IVDVIARLKG	SVGAEGLAGG	QVMDLECEAK	PGTTLDDLKW	IHIHKTATLL	QVAVASGAVL	240
GGATPEEVAA	CELFAMNIGL	AFQVADDILD	VTASSEDLGK	TAGKDEATDK	TTYPKLLGLE	300
ESKAYARQLI	DEAKESLAPF	GDRAAPLLAI	ADFIIDRKN			339

SEQ ID NO:47

MHLAPRRVPR	GRRSPDRVP	ERQCALGRRR	GAGSTGCARA	AAGVHRRRG	GEADPSAAVH	60
RGWQAGGGTG	LPDEVVSTAA	ALEMFHAFAL	IHDDIMDDSA	TRRGSPTVHR	ALADRLGAAL	120
DPDQAGQLGV	STAILVGDLA	LTWSDELLYA	PLTPHRLAAV	LPLVTAMRAE	TVHQYLDIT	180
SARRPGTDT	LALRIARYKT	AAYTMERPLH	IGAALAGARP	ELLAGLSAYA	LPAGEAFQLA	240
DDLLGVFGDP	RRTGKPDLD	LRGGKHTVLV	ALAREHATPE	QRHTLDLTLG	TPGLDRQGAS	300
RLRCVLVATG	ARAEAEERLIT	ERRDQALTAL	NALTLPPPLA	EALARLTLGS	TAHPA	355

SEQ ID NO:48

MSYFDNYFNE	IVNSVNDIHK	SYISGDVPKL	YEASYHLFTS	GGKRLRPLIL	TISSDLFGGQ	60
RERAYYAGAA	IEVLHFTTLV	HDDIMDQDNI	RRGLPTVHVK	YGLPLAILAG	DLHAKAFQL	120
LTQALRGLPS	ETIIKAFDIF	TRSIIIISEG	QAVDMFEDR	IDIKEQEYLD	MISRKTAALF	180
SASSSIGALI	AGANDNDVRL	MSDFGTNLGI	AFQIVDDILG	LTADKELGK	PVFSDIREGK	240
KTILVIKTL	LCKEDEKKIV	LKALGNKSAS	KEELMSSADI	IKKYSLDYAY	NLAEKYKNA	300
IDSLNQVSSK	SDIPGKALKY	LAEFTIRRRK				330

SEQ ID NO:49

MVAQTFNLDT	YLSQRQQQVE	EALSAAVLPA	YPERIYEAMR	YSLLAGGKRL	RPILCLAACE	60
LAGGSVEQAM	PTACALEMIH	TMSLIHDDL	AMDNDFFRRG	KPTNHKVFGE	DIAILAGDAL	120
LAYAFEHIAS	QTRGVPPQLV	LQVIARIGHA	VAATGLVGGQ	VVDLESEGKA	ISLETLEYIH	180
SHKTGALLEA	SVVSGGILAG	ADEELLARLS	HYARDIGLAF	QIVDDILDVT	ATSEQLGKTA	240
GKDQAAAKAT	YPSLLGLEAS	RQKAEELIQS	AKEALRPYGS	QAEPLALAD	FITRRQH	297

SEQ ID NO:50

MASVTLGSWI	VVHHHHHHHP	SSILTKRSR	SCPITLTKPI	SFRSKRTVSS	SSSIVSSSVV	60
TKEDNLRQSE	PSSFDFMSYI	ITKAELVNKA	LDSAVPLREP	LKIHEAMRYS	LLAGGKRVPR	120
VLCIAACELV	GGEESTAMPA	ACAVEMIHTM	SLIHDDLPCM	DNDLRRGKP	TNHKVFGEV	180
AVLAGDALLS	FAFEHLASAT	SSDVVSPVRV	VRAVDELAKA	IGTEGLVAGQ	VVDISSEGLD	240
LNDVGLEHLE	FIHLHKTAAL	LEASAVLGAI	VGGGSDDIE	RLRKFARCI	LLFQVDDIL	300
DVTKSSKELG	KTAGKDLIAD	KLTYPKIMGL	EKSREFAEKL	NREARDQLLG	FDSKVAPLL	360
ALANYIAYRQ	N					371

SEQ ID NO:51

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ggtgaaccag	ttatcgtctc	cgaccagca	aatgctctg	cttatgaatc	agttgctgca	120
gaattgtctt	caatggtgat	cgaaaacaga	caatcgcca	tgatcgtaac	tacatcaatc	180
gctgttttga	tcggttgat	tgatcgttg	gtatggagaa	gatccggtag	tggttaattct	240
aaaagagtcg	aacctttgaa	accattagta	attaagccaa	gagaagaaga	aatagatgac	300
ggtagaaaaga	aagtacaat	atcttctcgt	accctaaactg	gtacagctga	aggttttgca	360
aaagccttag	gtgaagaagc	taaggcaaga	tacgaaaaga	ctagattcaa	gatagtcgat	420
ttggtgact	atgcccgtga	tgacgatgaa	tacgaagaaa	agttgaagaa	agaagatggt	480

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tacaaatggt	ttacagaggg	taatgatcgt	ggtgaatggt	tgaaaaactt	aaagtacggt	600
gttttcgggt	tgggtaacag	acaatacga	catttcaaca	aagttgcaaa	ggttgctgac	660
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caatgtatag	aagatgactt	tactgcctgg	agagaagctt	tggtggcctg	attagacaca	780
atcttgagag	aagaagggtg	caccgccggt	gctaccccat	atactgctgc	agtattagaa	840
tacagagttt	ccatccatga	tagtgaagac	gcaaagttta	atgatatcac	tttggccaat	900
ggtaacgggt	atacagtttt	cgatgcacaa	cacccttaca	aagctaacgt	tgacgtcaag	960
agagaattac	atacaccaga	atccgacaga	agttgtatac	acttgggaat	tgatacgcgt	1020
ggttccgggt	taaccatgaa	gttgggtgac	catgtagggt	ttttatgcca	caatttgtct	1080
gaaactggtg	atgaagcatt	gagattggtg	gatatgtccc	ctgacactta	ttttagtgtg	1140
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ggtgtagcac	ctagattgca	accaagattc	tactcaatca	gttcttcacc	taagatcgcgt	1500
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acaattgctc	aagaacaagg	ttccatggat	agtaccaaag	ctgaaggttt	cgtaaagaac	2100
ttacaaactt	ccggtagata	cttgagagat	gtctggtga			2139

SEQ ID NO:52

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caaggccata	taaacccttt	catccagttt	ggcaaacgat	taatctccaa	agggtgcaaa	120
acaacacttg	ttaccaccat	ccacacctta	aactcaaccc	taaaccacag	taaccaccacc	180
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aatattgaat	cataactaaa	ttcaagatta	ttgtttgtaa	tattctttgt	cctaaaattt	1500
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aaagagatct	taggaaagat	gatcaaacag	acaaaggctt	catctggagt	gatttggaac	660
agtttcaaa	agtttagaaga	gtctgaattg	gagactgtaa	tcagagaaat	tccagcacct	720
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caacaggaag	tttttagctca	tggcgctatt	ggggcattct	ggactcattc	cggtatggaat	1080
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SEQ ID NO:54

MDGVIDMQTI	PLRTAIAIGG	TAVALVVALY	FWFLRSYASP	SHHSNHLFPV	PEVPGVPLVG	60
NLLQLKEKKP	YMTFTKWAEM	YGPIYSIRTG	ATSMVVVSSN	EIAKEVVVTR	FPSISTRKLS	120
YALKVLTEDEK	SMVAMSDYHD	YHKTVKRHIL	TAVLGPNAQK	KFRAHRDTMM	ENVSNELHAF	180
FEKNPNQEVN	LRKIFQSQLF	GLAMKQALGK	DVESIYVKDL	ETTMKREEIF	EVLVDDPMMG	240
AIEVDWRDFE	PYLKWPVKNKS	FENIHRMYT	RREAVMKALI	QEHKKRIASG	ENLNSYIDYL	300
LSEAQTLLTDK	QLLMSLWEPI	IESSDTMTVT	TEWAMYELAK	NPNMQDRLYE	EIQSVCGSEK	360
ITEENLSQLP	YLYAVFQETL	RKHCPVPIMP	LRYVHENTVL	GGYHVPAGTE	VAINIYGCNM	420
DKKWENPEE	WNPFRFLSEK	ESMDLYKTMA	FGGKRVKAG	SLQAMVISC	GIGRLVQDFE	480
WKLKDDAED	VNTLGLTTQK	LHPLLALINP	RK			512

SEQ ID NO:55

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gttccaaaaca	agtcccttcga	aaacatcatc	catagaatgt	acactagaag	agaagctggt	840
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tacattgatt	actgtttgtc	tgaagcccaa	accttgaccg	ataagcaatt	attgatgtct	960
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SEQ ID NO:61

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SEQ ID NO:64

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gaaggtgatg	aatgggtgtc	cgtaaaactgt	tcaaaggccg	caagagatat	tgttgctaga	540
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SEQ ID NO:66

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agatcagcaa	atcaatacca	agtgtgttct	gatttcgtaa	ctttacactg	taaagagaca	2100
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SEQ ID NO:67

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ggtgacaagt	tcgaaatgat	tactgccttc	tcaagagaag	gttctaagaa	ggtttacgtc	1860
caaacacagat	tgaaggaaag	atccaaagaa	gtctccgatt	tgttgtotca	aaaggcctac	1920
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SEQ ID NO:68

MEASYLYISI	LLLLASYLFT	TQLRRKSANL	PPTVFPSIPI	IGHLYLLKKP	LYRTLAKIAA	60
KYGPILQLQL	GYRRVLVISS	PSAAEECFN	NDVIFANRPK	TLFGKIVGGT	SLGSLSYGDQ	120
WRNLRRVASI	EILSVHRLNE	FHDIRVDENR	LLIRKLRSSS	SPVTLITVFY	ALTNLVIMRM	180
ISGKRYFDSG	DRELEEBGKR	FREILDETL	LAGASNVGDY	LPILNWLGVK	SLEKKLIALQ	240
KKRDDFFQGL	IEQVRKSRGA	KVGKGRKMTI	ELLLSLQESE	PEYITDAMIR	SFVLGLLAAG	300
SDTSAGTMEW	AMSLLVNHPH	VLKKAQAEID	RVIGNNRLID	ESDIGNIPYI	GCIINETLRL	360
YPAGPLLFPH	ESSADCVISG	YNI PRGTMLI	VNQWAIHHPD	KVWDDPETFK	PERFQGLEGT	420
RDGFKLMPFG	SGRRGCPGEG	LAIRLLGMTL	GSVIQCFDWE	RVGDEMVDMT	EGLGVTLPKA	480
VPLVAKCKPR	SEMTNLLSEL					500

SEQ ID NO:69

MQSESVEAST	IDLMTAVLKD	TVIDTANASD	NGDSKMPPAL	AMMFEIRDLL	LILTTSVAVL	60
VGCFVVLVWK	RSSGKKSCKE	LEPPKIVVVK	RRLEQEVDDG	KKKVTIFFGT	QTGTAEGFAK	120
ALFEEAKARY	EKAAFKVIDL	DDYAADLDEY	AEKLLKETYA	FFFLATYGDG	EPTDNAAKFY	180
KWFTEGDEKG	VWLQKLQYGV	FGLGNRQYEH	FNKIGIVVDD	GLTEQGAKRI	VPVGLGDDDQ	240
SIEDDFSAAK	ELVWPELDDL	LRDEDDKAAA	TPYTAAIPEY	RVVFHDKPDA	FSDDHTQTNG	300
HAVHDAQHPC	RSNVAVKKEL	HTPESDRSCT	HLEFDISHTG	LSYETGDHVG	VYCENLIEVV	360
EEAGKLLGLS	TDTYFSLHID	NEDGSPPLGGP	SLQPPFPPTCT	LRKALTNYAD	LLSSPKKSTL	420
LALAAHASDP	TEADRLRFLA	SREGKDEYAE	WVVANQRSLL	EVMEAFPSAR	PPLGVFFAAV	480
APRLQPRYYS	ISSSPKMEPN	RIHVTCALVY	EKTPAGRIHK	GICSTWMKNA	VPLTESQDCS	540
WAPIFVVRTSN	FRLPIDPKVP	VIMIGPGTGL	APFRGFLQER	LALKESGTEL	GSSILFFGCR	600
NRKVDYIYEN	ELNNFVENG	LSELDVAFSR	DGPTKEYVQH	KMTQKASEIW	NMLSEGAYLY	660
VCGDAKGMAK	DVHRTLHTIV	QEQGSLDSSK	AELYVKNLQM	SGRYLRDVM		709

SEQ ID NO:70

MASITHFLQD	FQATPFATAF	AVGVSVLLIF	FFFIRGFHST	KKNEYKLP	VPVVPGLPVV	60
GNLQLLKEKK	PYKTFRLWAE	IHGPIYSIRT	GASTMVVNS	THVAKEAMVT	RFSSISTRKL	120
SKALELLTSN	KSMVATSQDN	EFHKMVKKYI	LAELLGANAQ	KRHRIRHRTL	IENVLNKLHA	180
HTKNSPLQAV	NFRKIFESL	FGLAMKQALG	YDVSLSFVEE	LGTTLSREEI	YNVLVSDMLK	240
GAIQVDRDF	FPYLKWIPIK	SFEMKIQRLA	SRRQAVMNSI	VKEQKKSIAI	GKGENCYLNY	300
LLSEAKTLTE	QKISILAWET	IIEIADTTTV	TTEWAMYELA	KNPKQDRLY	NEIQNVCGTD	360
KITEEHLTKL	PYLSAVFHET	LRKYSPLV	PLRYAHEDTQ	LGGYVVPAGT	EIAVNIYGCN	420
MDKNQWETPE	EWKPERFLDE	KYDPMDMYKT	MSFGSGKRVK	AGSLQASLIA	CTSIGRLVQE	480
FEWRLKDGVE	ENVDTLGLTT	HKLYPMQAIL	QPRN			514

SEQ ID NO:71

MASMISLLLG	FVSSFLFIF	FLKLLFFFS	RHKMSEVSR	PSVVPVGFPL	IGNLLQLKEK	60
KPHKTFKWS	ELYGPIYSIK	MGSSSLIVLN	SIETAKEAMV	SRFSSISTRK	LSNALTVLTC	120
NKSMVATSQD	DDFHKFKVRC	LLNGLLGANA	QERKRHRDA	LIENVTSKLN	AHTRNHPQEP	180
VNFRAIFEHE	LFGVALKQAF	GKDVESIYVK	ELGVTLSRDE	IFKVLVHDM	EGAIQVDRD	240
FFPYLKWIPN	NSFEARIQQK	HKRRLAVMNA	LIQDRLNQND	SESDDDCYLN	FLMSEAKTLT	300
MEQIAILVWE	TIETADTTTL	VTEWAMYEL	AKHQSVQDRL	FKEIQSVCGG	EKIKEEQLPR	360
LPYVNGVFHE	TLRKYSAPL	VPIRYAHEDT	QIGGYHIPAG	SEIAINIYGC	NMDKKRWERP	420
EEWWPERFLE	DRYESSDLHK	TMAFGAGKRV	CAGALQASLM	AGIAIGRLVQ	EFEWKLRDGE	480
EENVDTYGLT	SQKLYPLMAI	INPRRS				506

SEQ ID NO:72

MDMMGIEAVP	FATAVVLGGI	SLVVLIFIRR	FVSNRKRVS	GLPPVPDIPG	LPLIGNLLQL	60
KEKKPHKTFA	RWAETYGPIF	SIRTGASTMI	VLNSSEVAKE	AMVTRFSSIS	TRKLSNALKI	120
LTFDKCMVAT	SDYNDFHKMV	KGFILRNVLG	APAQRHRCH	RDTLIENISK	YLHAHVKTSP	180
LEPVVLKKIF	ESEIFGLALK	QALGKDIESI	YVEELGTTLS	REEIFAVLVV	DPMAGAIQVD	240
WRDFFPYLSW	IPNKSMEKMI	QRMDFRRGAL	MKALIGEQQK	RIGSGEEKNS	YIDFLLSEAT	300
TLTEKQIAML	IWETIIEISD	TTLVTSEWAM	YELAKDPNRQ	EILYREIHKV	CGSNKLTEEN	360
LSKLPYLNSV	FHETLRKYSP	APMVPVRYAH	EDTQLGGYHI	PAGSQIAINI	YGCNMNKKQW	420
ENPEEWKPER	FLDEKYDLMD	LHKTMAFGGG	KRVCAGALQA	MLIACTSIGR	FVQEFEWKLM	480
GGEENVDTV	ALTSQKLHPM	QAIKARE				508

SEQ ID NO:73

MAELDTLDIV	VLGVIFLGTV	AYFTKGLKLG	VTKDPYANGF	AAGGASKPGR	TRNIVEAMEE	60
SGKNCVVFYQ	SQTGTAEDYA	SRLAKEGKSR	FGLNTMIADL	EDYDFDNLDT	VPSDNIVMFV	120
LATYGEGETP	DNAVDFYEFI	TGEDASFNEG	NDPPLGNLNY	VAFGLGNNTY	EHYNSMVRNV	180
NKALEKLGHA	RIGEAGEGDD	GAGTMEEDFL	AWKDPMWEAL	AKKMGLEERE	AVYEPIFAIN	240
ERDDLTPEAN	EVYLGEPNKL	HLEGTAKGPF	NSHNPYIAPI	AESYELFSAK	DRNCLHMEID	300
ISGSNLKYET	GDHIAIWPTN	PGEEVNKFLD	ILLDSGKQHS	VVTVKALEPT	AKVFPNPTT	360
YDAILRYHLE	ICAPVSRQFV	STLAAFAPND	DIKAEMNRLG	SDKDYFHEKT	GPHYNYIARF	420
LASVSKGEKW	TKIPFSAFIE	GLTKLQPRYY	SISSSSLVQP	KKISITAVVE	SQQIPGRDDP	480
FRGVATNYLF	ALKQKQNGDP	NPAPFGQSYE	LTGPRNKYDG	IHPVHVHRHS	NFKLPSDPGK	540
PIIMIGPGTG	VAPFRGFVQE	RAKQARDGVE	VGKTLFFGFC	RKSTEDFMYQ	KEWQEYKEAL	600
GDKFEMITAF	SREGSKKVYV	QHRLKERSKE	VSDLLSQKAY	FYVCGDAAHM	AREVNTVLAQ	660
IIAEGRGVSE	AKGEEIVKMN	RSANQYQVCS	DFVTLHCKET	TYANSELQED	VWS	713

SEQ ID NO:74

MKVSPEFEMS	AIIKGRMDPS	NSSFESTGEV	ASVIFENREL	VAILTTSIAV	MIGCFVVLWV	60
RRAGSRKVKV	VELPKPLIVH	EPEPEVEDGK	KKVSIFFGTQ	TGTAEGFAKA	LADAEKARYE	120
KATFRVVDLD	DYAADDQYE	EKLKNEFAV	FLLATYGDGE	PTDNAARFYK	WFAEGKERGE	180
WLQNLHYAVF	GLGNRQYEH	NKIAKVADEL	LEAQGGNRLV	KVGLGDDDDQ	IEDDFSARWE	240
SLWPELDMLL	RDEDDATTVT	TPYTAAVLEY	RVVFHDSADV	AAEDKSWINA	NGHAVHDAQH	300
PFRRNVVVRK	ELHTSASDRD	CSHLEFNISG	SALNYETGDH	VGVCENLITE	TVDEALNLLG	360
LSPETYFSIY	TDNEDGTPLG	GSSLPFPFSP	CTLRTALTRY	ADLLNSPKKS	ALLALAAHAS	420
NPVEADRLRY	LASPAGKDEY	AQSVIGSQKS	LLEVMAEFPS	AKPPLGVFFA	AVAPRLQPRF	480
YSISSSPRMA	PSRIHVTCAL	VYDKMPTGRI	HKGVCSTWMK	NSVPMESKSE	CSWAPIFVRQ	540
SNFKLPAESK	VPIIMVPGTG	GLAPFRGFLO	ERLALKESGV	ELGPSILFFG	CRNRRMDYIY	600
EDELNNFVET	GALSELVIAF	SREGPTKEYV	QHKMAEKASD	IWNLISEGAY	LYVCGDAKGM	660
AKDVHRTLHT	IMQEQGSLDS	SKAESMVKNL	QMNGRYLRDV	W		701

SEQ ID NO:75

MATLLEHFQA	MPPAIPIALA	ALSWLFLFYI	KVSFFSNKSA	QAKLPPVPVV	PGLPVIGNLL	60
QLKEKKPYQT	FTRWAEYGP	IYSIRTGAST	MVVLNTTQVA	KEAMVTRYLS	ISTRKLSNAL	120
KILTADKCMV	AISYNDFFHK	MIKRYILSNV	LGPSAQKRHR	SNRDTLRANV	CSRLHSQVKV	180
SPREAVNRFR	VFEWELFGIA	LKQAFGKDIE	KPIYVEELGT	TLSRDEIFKV	LVLDIMEGAI	240
EVDWRDFPPY	LRWIPNTRME	TKIQRLYFRR	KAVMTALINE	QKKRIASGEE	INCYIDFLK	300
EGKTLTMDQI	SMLLWETVIE	TADTTMVTTE	WAMYEVAKDS	KRQDRLYQEI	QKVCSEMVT	360
EEYLSQLPYL	NAVPHETLRK	HSPAALVPLR	YAHEDTQLGG	YYIPAGTEIA	INIYGCNMDK	420
HQWESPEEWK	PERFLDPKFD	PMDLYKTMAF	GAGKRVCSGS	LQAMLIACPT	IGRLVQEFWE	480
KLRDGEENNV	DTVGLTTHKR	YPMHAILKPR	S			511

SEQ ID NO:76

MQSDSVKVP	FDLVSAMNG	KAMEKLNASE	SEDPTLPLAL	KMLVENRELL	TLFTTSFAVL	60
IGCLVFLMWR	RSSSKKLVQD	PVPQVIVVKK	KEKESEVDDG	KKKVSIFYGT	QTGTABGFAK	120
ALVEEAKVRY	EKTSFKVIDL	DDYAADDDEY	EEKLKKESLA	FFFLATYGDG	EPTDNAANFY	180
KWFTEGDDKG	EWLKKLQYGV	FGLGNRQYEH	FNKIAIVVDD	KLTEMGAKRL	VPVGLGDDDDQ	240
CIEDDFTAWK	ELVWPELDQL	LRDEDDTSVT	TPYTAAVLEY	RVVYHDKPAD	SYAEDQHTN	300
GHVVHDAQHP	SRSNVAFKKE	LHQSQSDRSC	THLEFDISHT	GLSYETGDHV	GVYSENLSSEV	360
VDEALKLLGL	SPDTYFSVHA	DKEDGTPIGG	ASLPPFPFPC	TLRDALTRYA	DVLSSPKKVA	420
LLALAAHASD	PSEADRLKFL	ASPAGKDEYA	QWIVANQRSL	LEVMSQSFPSA	KPPLGVFFAA	480
VAPRLQPRYY	SISSSPKMS	NRIHVTCALV	YETTPAGRIH	RGLCSTWMKN	AVPLTESPDC	540
SQASIFVVRTS	NFRLPVDPKV	PVIMIGPGTG	LAPFRGFLQE	RLALKESGTE	LGSSIFFFGC	600
RNRKVDIFIYE	DELNNFVETG	ALSELIVAFS	REGTAKEYVQ	HKMSQKASDI	WKLLSEGAFL	660
YVCGDAKGMA	KDVHRTLHTI	VQEQGSLDSS	KAELYVKNLQ	MSGRYLRDVW		710

SEQ ID NO:77

MSKSNSMNST	SHETLFQQLV	LGLDRMPLMD	VHWLIYVAFG	AWLCSYVIHV	LSSSSTVKVP	60
VVGYSRVFEP	TWLLRLRFVW	EGGSIIGQGY	NKFKDSIFQV	RKLGTDIVII	PPNYIDEVRK	120
LSQDKTRSVE	PFINDFAGQY	TRGMVFLQSD	LQNRVIQORL	TPKLVSLTKV	MKEELDIALT	180
KEMPDMMKND	WVEVDISSIM	VRLISRISAR	VFLGPEHCRN	QEWLTTTAEY	SESLFITGFI	240
LRVVPILRFP	FIAPLLPSYR	TLLRNVSSGR	RVIGDIIRSQ	QGDGNEDILS	WMRDAATGEE	300
KQIDNIAQRM	LILSLASLHT	TAMTMTHAMY	DLCACPEYIE	PLRDEVKSVV	GASGWDKTAL	360
NRFHKLDSFL	KESQRFNPVF	LLTFNRIYHQ	SMTLSDGTNI	PSGTRIAVPS	HAMLQDSAHV	420

PGPTPTEFD GFRYSKIRSD SNYAQKYLFS MTDSSNMAFG YGKYACPGRF YASNEMKLTL 480  
 AILLQFEFK LPDGKGRPRN ITIDSDMIPD PRARLCVRKR SLRDE 525

SEQ ID NO:78

MEDPTVLYAC LAIAVATFVW RWYRDLRSI PTVGGSDLPI LSYIGALRWT RRGREILQEG 60  
 YDGYRGSTFK IAMLDRWIVI ANGPKLADEV RRRPDEELNF MDGLGAFVQT KYTLGEAIHN 120  
 DPYHVDIIRE KLTRGLPAVL PDVIEELTLA VRQYIPTEGD EWVSVNCSKA ARDIVARASN 180  
 RVFVGLPACR NQGYLDLAI DFTLSVVKDRA IINMPELLK PIVGRVVGNA TRNVRRVAVPF 240  
 VAPLVEERRR LMEEYGEDWS EKPNDMLQWI MDEAASRDSS VKAIAERLLM VNFAAIHTSS 300  
 NTITHALYHL AEMPETLQPL REEIEPLVKE EGWTKAAMGK MWWLDSFLRE SQRYNGINIV 360  
 SLTRMADKDI TLDGTFLEPK GTLVAVPAYS THRDDAVYAD ALVDFPFRFS RMRAREGEGT 420  
 KHQFVNTSVE YVFPFHGKHA CPGRFFAANE LKAMLAYIVL NYDVKLPGDG KRPLNMYWGP 480  
 TVLPAPAGQV LFRKRQVSL 499

SEQ ID NO:79

MDAVTGLLTV PATAITIGGT AVALAVALIF WYLKSYTSAR RSQSNHLPRV PEVPGVPLLG 60  
 NLLQLKEKPK YMTFTRWAAT YGPIYSIKTG ATSMVVVSSN EIAKEALVTR FQSISTRNLS 120  
 KALKVLTADK TMVAMSDYDD YHKTVKRHIL TAVLGPNAQK KHRIHRDIMM DNISTQLHEF 180  
 VKNNPEQEEV DLRKIFQSEL FGLAMRQALG KDVESLYVED LKITMNRDEI FOVLVVDPM 240  
 GAIDVDWRDF FPYLKWVPNK KFENTIQQMY IRREAVMKSL IKEHKKRIAS GEKLSYIDY 300  
 LLSEAQTITD QQLLMSLWEP IIESDITMV TTEWAMYELA KNPKLQDRLY RDIKSVCGSE 360  
 KITTEHLSQL PYITAI FHET LRRHSPVPII PLRHVHEDTV LGGYHVPAGT ELAVNIYGCN 420  
 MDKNVWENPE EWNPERFMKE NETIDFQKTM AFGGGKRVCA GSLQALLTAS IGIGRMVQEF 480  
 EWKLKDMTQE EVNTIGLTTQ MLRPLRAIIK PRI 513

SEQ ID NO:80

atggaagtaa cagtagctag tagtgtagcc ctgagcctgg tctttattag catagtagta 60  
 agatgggcat ggagtgtggt gaattgggtg tggtttaagc cgaagaagct ggaaagattt 120  
 ttgagggagc aaggccttaa aggcaattcc tacaggtttt tatatggaga catgaaggag 180  
 aactctatcc tgctcaaac agcaagatcc aaaccctga acctctccac ctcccatgac 240  
 atagcacctc aagtcacccc tttgtcgac caaacctga aagcttacgg taagaactct 300  
 ttaattggg ttgccccat accaagggtg aacataatga atccagaaga tttgaaggac 360  
 gtcttaacaa aaaatgttga cttgttgaag ccaatatcaa acccacttat caagttgcta 420  
 gctacaggtg ttgcaatcta tgaaggtgag aaatggacta aacacagaag gattatcaac 480  
 ccaacattcc attcggagag gctaaagcgt atgttacctt cattcacca aagttgtaat 540  
 gagatggtca aggaatggga gagcttggtg tcaaaagagg gttcatcatg tgagtggat 600  
 gtctggcctt ttcttgaaaa tatgtcggca gatgtgatct cgagaacagc atttggaact 660  
 agctacaaaa aaggacagaa aatctttgaa ctcttgagag agcaagtaat atatgtaacg 720  
 aaaggcttc aaagttttga cattccagga tggaggtttc tccaactaa gatgaacaag 780  
 aggatgaaatg agattaacga agaaataaaa ggattaatca ggggtattat aattgacaga 840  
 gagcaaatca ttaaggcagg tgaagaaacc aacgatgact tattaggtgc acttatggag 900  
 tcaaaactga aggacattcg ggaacatggg aaaaacaaca aaaatgttgg gatgagtatt 960  
 gaagatgtaa ttcaggagtg taagctgttt tactttgctg ggcaagaaac cacttcagtg 1020  
 ttgctggcctt ggacaatggt ttacttgggt caaaatcaga actggcaaga tcgagcaaga 1080  
 caagaggttt tgcaagtctt tgaagcagc aagccagatt ttgatggtct agctcacctt 1140  
 aaagtogtaa coatgatttt gcttgaagtt ctctgattat acccaccagt cattgaactt 1200  
 attcgaacca ttcacaagaa aacacaactt ggaagctct cactaccaga aggagttgaa 1260  
 gtccgcttac caacactgct cattcaccat gacaaggaac tgtggggtga tgatgcaaac 1320  
 cagttcaate cagagagggt ttcggaagga gtttccaaag caacaaagaa ccgactctca 1380  
 ttcttcccct tcggagccgg tccacgcatt tgcattggac agaacttttc tatgatggaa 1440  
 gcaaagtgg ccttagcatt gatottgcaa cacttcacct ttgagcttcc tccatctcat 1500  
 gcacatgctc cttcccacg tataaccctt caaccacagt atggtgttcg tatcatttta 1560  
 catcgacgtt ag 1572

SEQ ID NO:81

atggaagtca ctgtcgctc ttctgtcgct ttatccttag tcttcatttc cattgtcgtc 60  
 agatgggctt ggtccgttgt caactgggtt tggttcaaac caagaagtt ggaagattc 120  
 ttgagagagc aaggtttgaa gggaattct tatagattct tgtacgggtga catgaaggaa 180  
 aattctattt tghtgaagca agccagatcc aaaccaatga acttgtctac ctctcatgat 240  
 attgctccac aagtactctc attcgtcgat caaacgttta aagcctacgg taagaactct 300

ttcaattggg	ttggccaat	tcctagagtt	aacatcatga	accagaaga	tttgaaggat	360
gtcttgacca	agaacgttga	cttcggttaag	ccaatttcca	accattgat	taaattgttg	420
gctactggta	ttgccattta	cgaaggtgaa	aagtggacta	agcatagaag	aatcatcaac	480
cctaccttcc	actctgaaag	attgaagaga	atgttaccat	ctttccatca	atcctgtaat	540
gaaatggtta	aggaatggga	atccttgggt	tctaaagaag	gttcttcttg	cgaattggat	600
gtttggccat	tcttggaaaa	tatgtctgct	gatgtcattt	ccagaaccgc	tttcgggtacc	660
tcctacaaga	agggccaata	gattttcgaa	ttgttgagag	agcaagttaa	ttacgttacc	720
aagggtttcc	aatccttcta	catcccaggt	tggagattct	tgccaactaa	aatgaacaag	780
cgtatgaacg	agatcaacga	agaaattaaa	ggtttgatca	gaggtattat	tatcgacaga	840
gaacaaatta	ttaaagctgg	tgaagaaacc	aacgatgatt	tgttgggtgc	ttgatggag	900
tccaacttga	aggatattag	agaacatggt	aagaacaaca	agaatgttgg	tatgtctatt	960
gaagatgta	ttcaagaatg	taagttattc	tacttcgctg	gtcaagagac	cacttctggt	1020
ttgtagcct	ggactatggt	cttgttaggt	caaaaccaaa	attggcaaga	tagagctaga	1080
caagaagttt	tgcaagtctt	cggttcttcc	aagccagact	ttgatggttt	ggcccacttg	1140
aaggttgta	ctatgatttt	gtagaagttt	ttgagattgt	accaccagt	cattgagtta	1200
atcagaacca	ttcataaaaa	gactcaattg	ggtaaattat	ctttgccaga	aggtggtgaa	1260
gtcagattac	caaccttgtt	gattcaccac	gataaggaat	tatgggggtga	cgacgtaat	1320
caatttaatc	cagaaagatt	ttccgaaggt	gtttccaagg	ctaccaaaaa	ccgtttgtcc	1380
ttcttcccat	ttgggtgctg	tccacgtatt	tgtatcggtc	aaaacttttc	catgatggaa	1440
gccaagtgg	ctttggcttt	aatcttgcaa	cacttcactt	tcgaattgtc	tccatcccat	1500
gcccacgctc	cttctcatag	aatcacttta	caaccacaat	acgggtgtcag	aatcatctta	1560
cacagaagat	aa					1572

SEQ ID NO:82

MEVTVASSVA	LSLVFISIVV	RWAWSVVNWV	WFKPKKLERF	LREQGLKGN	YRFLYGMKE	60
NSILLKQARS	KPMNLSTSHD	IAPQVTPFVD	QTVKAYGKNS	FNWVGPIPRV	NIMNPEDLKD	120
VLTKNVDFVK	PISNPLIKLL	ATGIAIYEGE	KWKHRRIIN	PTFHSERLKR	MLPSFHQSCN	180
EMVKEWESLV	SKEGSSCELD	VWPFLENMSA	DVISRTAFGT	SYKKGQKIFE	LLREQVIYVT	240
KGFQSFYIPG	WRFLPTKMNK	RMNEINEEIK	GLIRGIIIDR	EQIIKAGEET	NDDLGLALME	300
SNLKDIREHG	KNNKNVGM	EDVIQECKLF	YFAGQETTSV	LLAWTMVLLG	QNQNWQDRAR	360
QEVLVQVFGSS	KPDFDGLAHL	KVVTMILLEV	LRLYPPVIEL	IRTIHKKTQL	GKLSLPEGVE	420
VRLPILLIHH	DKELWGDND	QFNPERFSEG	VSKATNRLS	FFPFGAGPRI	CIGQNFMSME	480
AKLALALILQ	HFTFELSPSH	AHAPSHRITL	QPQYGVRIIL	HRR		523

SEQ ID NO:83

MENKTETTVR	RRRRIILFPV	PFQGHINPIL	QLANVLYSKG	FSITIFHTNF	NKPKTSNYPH	60
FTFRFILDND	PQDERISNLP	THGPLAGMRI	PIINEHGADE	LRRELELLML	ASEEDEEVSC	120
LITDALWYFA	QSVADSLNLR	RLVLMTSSLF	NFAHVSLLPQ	FDELGYLDPD	DKTRLEEQAS	180
GFPMLKVKDI	KSAYSNWQIL	KEILGKMIKQ	TKASSGVIWN	SFKELEESEL	ETVIREIPAP	240
SFLIPLPKHL	TASSSSLLDH	DRTVFQWLDQ	QPPSSVLYVS	FGSTSEVDEK	DFLEIARGLV	300
DSKQSFVLVW	RPGFVKGSTW	VEPLPDGFLG	ERGRIVKWVP	QQEVLAHGAI	GAFWTHSGWN	360
STLESVCEGV	PMIFSDFGLD	QPLNARYMSD	VLKVGVIYLN	GWERGEIANA	IRRMVMDVEEG	420
EYIRQNRVRL	KQKADVSLMK	GGSSYESLES	LVSYISSL			458

SEQ ID NO:84

MDAMATTEKK	PHVIFIPFPA	QSHIKAMLKL	AQLLHHKGLQ	ITFVNTDFIH	NQFLESSGPH	60
CLDGAAGFRF	ETIPDGVSHS	PEASIPIRES	LLRSIETNFL	DRFIDLVTKL	PDPTCIISD	120
GFLSVFTIDA	AKKLGIPVMM	YWTLAACGFM	GFYHIHSLIE	KGFAPLKDAS	YLTNGYLDTV	180
IDWVPGMEGI	RLKDFPLDWS	TDLNDKVLMF	TTEAPQRSKH	VSHHIFHTFD	ELEPSIIKTL	240
SLRYNHIYTI	GPLQLLLDQI	PEEKKQTGIT	SLHGYSLVKE	EPECFQWLQS	KEPNSVVYVN	300
FGSTTVMSLE	DMTEFGWGLA	NSNHYFLWII	RSNLVIGENA	VLPPELEEHI	KKRGFIASWC	360
SQEKVLKHP	VGGFLTHCGW	GSTIESLSAG	VPMICWPYSW	DQLTNCRYIC	KEWEVGLEM	420
TKVKRDEVKR	LVQELMGE	HKMRNKAKDW	KEKARIAIAP	NGSSSLNIDK	MVKEITVLR	480

SEQ ID NO:85

MATSDSIVDD	RKQLHVATFP	WLAFGHILPY	LQLSKLIAEK	GHKVSFLSTT	RNIQRLSSHI	60
SPLINVVQLT	LPRVQELPED	AEATTDVHPE	DIPYLKASD	GLQPEVTRFL	EQHSPDWIY	120
DYTHYWLPSI	AASLGISRAH	FSVTTPWAIA	YMGPSADAMI	NGSDGRITVE	DLTTPPKWFP	180



FPTKVCWRKH	DLARLVPIKA	PGISDGYRMG	LVLKGSDCLL	SKCYHEFGTQ	WLPLLETLHQ	240
VPVVPVGLLP	PEIPGDEKDE	TWVSIKKWLD	GKQKGSVVYV	ALGSEVLVSQ	TEVVELALGL	300
ELSGLPFVWA	YRKPFGPAKS	DSVELPDGFV	ERTDRGLVW	TSWAPQLRIL	SHEVCGFLT	360
HCGSGSIVEG	LMFGHPLIML	PIFGEIPRNE	EDGCLTKESV	ARSLRSVVVE	KEGEIYKANA	420
RELSKIYNDT	KVEKEYVSQF	VDYLEKNARA	VAIDHES			457

SEQ ID NO:86

MDSGYSSSYA	AAAGMHVVIC	PWLAFGHLLP	CLDLAQRLAS	RGHRVSFVST	PRNISRLPPV	60
RPALAPLVAF	VALPLRVEG	LPDGAESTND	VPHDRPDMVE	LHRRAPDGLA	APFSEFLGTA	120
CADWVIVDVF	HHWAAAAALE	HKVPCAMMLL	GSAHMIASIA	DRRLERAETE	SPAAAGQGRP	180
AAAPTFEVAR	MKLIIRTKGSS	GMSLAERFSL	TLRSRSLVVG	RSCVEFEPET	VPLLSTLRGK	240
PITFLGLMPP	LHEGRREDGE	DATVRWLDAQ	PAKSVVYVAL	GSEVPLGVEK	VHELALGLEL	300
AGTRFLWALR	KPTGVSDADL	LPAGFEERTR	GRGVVATRWV	PQMSILAHAA	VGAFLTHCGW	360
NSTIEGLMFG	HPLIMLPIFG	DQGPNARLIE	AKNAGLQVAR	NDGDGSDFRE	GVAAAIRAVA	420
VEEESKVFQ	AKAKKLQEI	ADMACHERYI	DGFIQQLRSY	KD		462

SEQ ID NO:87

MSSSSSSSTS	MIDLMAAIK	GEPVIVSDPA	NASAYESVAA	ELSSMLIENR	QFAMIVTTSI	60
AVLIGCIVML	VWRRSGSGNS	KRVEPLKPLV	IKPREEID	GRKKVTIFFG	TQTGTAEQFA	120
KALGEEAKAR	YEKTRFKIVD	LDDYAADDDE	YEEKLKKEDV	AFFFLATYGD	GEPTDNAARF	180
YKWFTEGNDR	GEWLKLNLYG	VFGLGNRQYE	HFNKVAKVVD	DILVEQGAQR	LVQVGLGDDD	240
QCIEDDDTAW	REALWPELDT	ILREEGDTAV	ATPYTAAVLE	YRVSIHSDSE	AKFNITLAN	300
GNGYTVFDAQ	HPYKANVAVK	RELHTPESDR	SCIHLEFDIA	GSGLTMKLG	HVGVLCDNLS	360
ETVDEALRLL	DMSPDYFSL	HAEKEDGTPI	SSSLPPFPFP	CNLRALTALRY	ACLLSSPKKS	420
ALVALAAHAS	DPTAERLKH	LASPAKDEY	SKWVVSQRS	LLEVMAEFPS	AKPPLGVFFA	480
GVAPRLQPRF	YSISSSPKIA	ETRIHVTCAL	VYEKMPGTGRI	HKGVCSTWMK	NAVPEKSEK	540
LFLGRPIFVR	QSNFKLPSDS	KVPIIMIGPG	TGLAPFRGFL	QERLALVESG	VELGPSVLFF	600
GCRNRRMDFI	YEEELQRFVE	SGALAELSVA	FSREGPTKEY	VQHKMMDKAS	DIWNMISQGA	660
YLYVCGDAKG	MARDVHRS LH	TIAQEQGSMD	STKAEGFVKN	LQTSGRYL RD	VW	712

SEQ ID NO:88

MATSDSIVDD	RKQLHVATFP	WLAFGHILPY	LQLSKLIAEK	GHKVSFLSTT	RNIQRLSSHI	60
SPLINVQILT	LPRVQELPED	AEATTDVHPE	DIPYLKASD	GLQPEVTRFL	EQHSPDWIY	120
DYTHYWLPSI	AASLGISRAH	FSVTPWAIA	YMGPSADAMI	NGSDGRITVE	DLTTPPKWFP	180
FPTKVCWRKH	DLARLVPIKA	PGISDGYRMG	MVLKGSDCLL	SKCYHEFGTQ	WLPLLETLHQ	240
VPVVPVGLLP	PEIPGDEKDE	TWVSIKKWLD	GKQKGSVVYV	ALGSEALVSQ	TEVVELALGL	300
ELSGLPFVWA	YRKPFGPAKS	DSVELPDGFV	ERTDRGLVW	TSWAPQLRIL	SHEVCGFLT	360
HCGSGSIVEG	LMFGHPLIML	PIFGDQPLNA	RLEDKQVGI	EIPRNEEDGC	LTKESVARSL	420
RSVVVEKEGE	IYKANARELS	KIYNDTKVEK	EYVSQFVDYL	EKNARAVAI	HES	473

SEQ ID NO:89

atggctactt	ctgattccat	cgttgacgat	agaaagcaat	tgcatgttgc	tacttttcca	60
tggttggctt	tcggtcatat	tttgccatac	ttgcaattgt	ccaagttgat	tgctgaaaag	120
ggtcacaaag	tttcattcct	gtctaccacc	agaaacatcc	aaagattgtc	ctctcatatc	180
tccccattga	tcaacgttgt	tcaattgact	ttgccaagag	tccaagaatt	gccagaagat	240
gctgaagcta	ctactgatgt	tcatccagaa	gatatcccct	acttgaaaaa	ggcttccgat	300
ggtttacaac	cagaagttac	tagattcctg	gaacaacatt	cccagattg	gatcatctac	360
gattatactc	attactggtt	gccatccatt	gctgcttcat	tgggtatttc	tagagcccat	420
ttctctgtta	ctactccatg	ggctattgct	tatatgggtc	catctgctga	tgctatgatt	480
aacggtctctg	atggtagaac	taccggtgaa	gatttgacta	ctccaccaa	gtggtttcca	540
tttccacaaa	aagtctgttg	gagaaaaac	gatttggcta	gattggttcc	atacaaaagt	600
ccaggatatt	ctgatggtta	cagaatgggt	atggttttga	aaggttccga	ttgcttggtg	660
tctaagtgtc	atcatgaatt	cgttactcaa	tggttgcctt	tgttgaaaac	atgcatcaa	720
gttccagttg	ttccagtagg	ttgtttgcca	ccagaaattc	caggtgacga	aaaagacgaa	780
acttgggttt	ccatcaaaaa	gtggttggat	ggtaagcaaa	agggttctgt	tgtttatgtt	840
gctttgggtt	ccgaagcttt	ggtttctcaa	accgaagtgt	ttgaattggc	tttgggtttg	900
gaattgtctg	gtttgccatt	tgtttgggct	taacagaaaac	ctaaaggtcc	agctaagtct	960
gattctgttg	aattgccaga	tggtttcgtt	gaaagaacta	gagatagagg	tttgggtttg	1020
acttcttggg	ctccacaatt	gagaattttg	tctcatgaat	ccgtctgtgg	tttcttgact	1080
catttgggtt	ctggttctat	cgttgaaggt	ttgatgtttg	gtcaccatt	gattatgttg	1140

ccaatctttg	gtgaccaacc	attgaacgct	agattattgg	aagataagca	agtcgggtatc	1200
gaaatcccaa	gaaatgaaga	agatgggtgc	ttgaccaaag	aatctgttgc	tagatctttg	1260
agatccgctg	tcggttgaaaa	agaaggtgaa	atctacaagg	ctaacgctag	agaattgtcc	1320
aagatctaca	acgataccaa	ggtcgaaaaa	gaatacgttt	cccaattcgt	tgactacttg	1380
gaaaagaatg	ctagagctgt	tgccattgat	catgaatctt	ga		1422

SEQ ID NO:90

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actttggctt	ggagagtttt	gaattgggtc	tggttaagac	caaaaaagtt	gaaagatgc	120
ttgagagaac	aaggtttgac	tggttaactct	tacagattgt	tgttcgggtga	taccaaggac	180
ttgtctaaaga	tgttggaaca	aactcaatcc	agcctatca	agttgtctac	ctctcatgat	240
attgctccaa	gagttactcc	attcttccat	agaactgtta	actccaacgg	taagaactct	300
tttgtttgga	tggttccaat	tccaagagtc	catattatga	accctgaaga	tttgaaggac	360
gctttcaaca	gacatgatga	ttccataaag	accgtcaaga	acccaattat	gaagtctcca	420
ccaccaggtg	tagttgggtat	tgaaggtgaa	caatgggcca	aacatagaaa	gattattaac	480
ccagccttcc	acttggaaaa	gttgaaggtt	atggttccaa	tcttctacca	atcctgctct	540
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SEQ ID NO:91

MEASRASCVA	LCVVVSVIVI	TLAWRVLNWV	WLRPKKLERC	LREQGLTGNS	YRLLFGDTKD	60
LSKMLEQTS	KPIKLSSTSHD	IAPRVTPFFH	RTVNSNGKNS	FVWMGPIPRV	HIMNPEDLKD	120
AFNRHDDFHK	TVKNPIMKSP	PPGIVGIEGE	QWAKHRKIIN	PAFHLEKLKG	MVPIFYQSCS	180
EMINKWESLV	SKESSELDV	WPYLENFTSD	VISRAAFGSS	YEGRKIFQL	LREEAKVYSV	240
ALRSVYIPGW	RFLPTKQNK	TKEIHNEIKG	LLKGIINKRE	EAMKAGEATK	DDLGLILMES	300
NFREIQEHGN	NKNAGMSIED	VIGECKLFYF	AGQETTSVLL	VWTMILLSQN	QDWQARAREE	360
VLKVFSGSNIP	TYEELSHLKV	VTMILLEVL	LYPSVVALPR	TTHKKTQLGK	LSLPAGVEVS	420
LPILLVHDK	ELWGEDANEF	KPERFSEGV	KATKNKFTYL	PFGGGPRICI	QONFAMVEAK	480
LALALILQHF	AFELSPSYAH	APSAVITLQP	QFGAHIILHK	R		521

SEQ ID NO:92

ASWVAVLSV	VWSMVIWAW	RVLNWWVLRP	KKLEKCLREQ	GLAGNSYRLL	FGDTKDLSKM	60
LEQTQSKPIK	LSTSHDIAPH	VTPFFHQTVN	SYGKNSFVWM	GPIPRVHIMN	PEDLKDTFNR	120
HDDFHKVVK	PIMKSLPQGI	VGIEGEQWAK	HRKIINPAFH	LEKLGMPVPI	FYRSCSEMIN	180
KWESLVSKES	SCELDVWPYL	ENFTSDVISR	AAFSSYEEG	RKIFQLLREE	AKIYTVAMRS	240
VYIPGWRFLP	TKQNKKAKEI	HNEIKGLLKG	IINKREEAMK	AGEATKDDL	GILMESNFRE	300
IQEHGNNKNA	GMSIEDVIGE	CKLFYFAGQE	TTSVLLVWTM	VLLSQNDWQ	ARAREVLQV	360
FGSNIPTYEE	LSQLKVVTMI	LLEVLRLYPS	VVALPRTHK	KTQLGKLSLP	AGVEVSLPIL	420
LVHDKELWG	EDANEFKPER	FSEGVSKATK	NQFTYFFPFG	GPRICIGQNF	AMMEAKLALS	480
LILRHFALEL	SPLYAHAPSV	TITLQPQYGA	HIILHKR			517

SEQ ID NO:93

MEASRPSCVA	LSVVLVSVIVI	AWAWRVLNWV	WLRPNKLERC	LREQGLTGNS	YRLLFGDTKE	60
ISMVVEQAQS	KPIKLSSTTHD	IAPRVIPFFH	QIVYTYGRNS	FVWMGPTPRV	TIMNPEDLKD	120

AFNKSDDEFQR	AISNPVIXSI	SQGLSSLEGE	KWAKHRKIIN	PAFHLEKLKG	MLPTFYQSCS	180
EMINKWESLV	FKEGSREMDV	WPYLENLTSD	VISRAAFGSS	YEEGRKIFQL	LREEAKFYTI	240
AARSVYIPGW	RFLPTKQNKR	MKEIHKEVRG	LLKGIINKRE	DAIKAGEAAK	GNULLGILMES	300
NFREIQEHGN	NKNAGMSIED	VIGECKLFYF	AGQETTSVLL	VWTLVLLSON	QDWQARAREE	360
VLQVFGTNIP	TYDQLSHLKV	VTMILLEVLK	LYPAVVELPR	TTYKKTQLGK	FLLPAGVEVS	420
LHIMLAHHDK	ELWGEDAKEF	KPERFSEGVK	KATKNQFTYF	PFGAGPRICI	GQNFAMLEAK	480
LALSLLQHF	TFELSPSYAH	APSVTITLHP	QFGAHFILHK	R		521

SEQ ID NO:94

CVALSIVLVLS	IVIAWAWRVL	NWVWLRPNKL	ERCLREQGLT	GNSYRLLFGD	TKEISMVVEQ	60
AQSKPIKLST	THDIAPRVIP	FSHQIVYTYG	RNSFVWMGPT	PRVTIMNPED	LKDAFNKSDE	120
FQRAISNPIV	KSISQGLSSL	EGEKWAKHRK	IINPAFHLEK	LKGMLPTFYQ	SCSEMINKWE	180
SLVFKEGSRE	MDVWPYLENL	TSDIVISRAAF	GSSYEGRKI	FQLLREEAKF	YTIAARSVYI	240
PGWRFLPTKQ	NKRKMEIHKE	VRGLLKGIIIN	KREDAIKAGE	AAKGNLLGIL	MESNFREIQE	300
HGNNKNAGMS	IEDVIGECKL	FYFAGQETTS	VLLVWTLVLL	SONQDWQARA	REEVLQVFGT	360
NIPTYDQLSH	LKVVTMILLE	VLRLYPVAVVE	LPRTTYKKTQ	LGKFLLPAGV	EVSLHIMLAH	420
HDKELWGEDA	KEFKPERFSE	GVSKATKNQF	TYFFPFGAGPR	ICIGQNFAML	EAKLALSLLI	480
QHFTFELSPS	YAHAPSVTIT	LHPQFGAHFI	LHKR			514

SEQ ID NO:95

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HLEKLGKMNVP	IFYQSCSEMI	NIWKSLSVSK	SSCELDVWPY	LENFTSDVIS	RAAFGSSYEE	120
GRKIFQLLRE	EAKVYTVAVR	SVYIPGWREL	PTKQNKKTKE	IHNEIKGLLK	GIINKREAM	180
KAGEATKDDL	LGILMESNFR	EIQEHGNNKN	AGMSIEDVIG	ECKLFYFAGQ	ETTSVLLVWT	240
MVLLSQNQDW	QARAREEVLO	VFGSNIPTYE	ELSHLKVVTM	ILLEVLRLYP	SVVALPRTTH	300
KKTQLGKLSL	PAGVEVSLPI	LLVHHDKELW	GEDANEFKPE	RFSEGVSKAT	KNQFTYFPFG	360
GGPRICIGQN	FAMMEAKLAL	SLIIQHFTFE	LSPQYSHAPS	VTITLQPQYG	AHLILHKR	418

SEQ ID NO:96

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SEQ ID NO:97

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SEQ ID NO:98

MSSNSDLVRR	LESVLGVSTG	GSVTDVSVVI	ATTSIALVIG	VLVLLWRRSS	DRSREVKQLA	60
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DYTAEDDKYG	EKLKKTMAF	FMLATYGDGE	PTDNAARFYK	WFTEGDRGV	WLEHLRYGVF	180
GLGNROYEYH	NKIAKVDDL	LVEQGAQRLV	TVGLGDDQD	IEDDFSAWKE	ALWPELDQLL	240
QDDTNTVSTP	YTAVIPEYRV	VIHDPVTSY	EDPYSNMANG	NASYDIHHC	RANVAVQKEL	300
HKPESDRSCI	HLEFDIFATG	LTYETGDHVG	VYADNCDDTV	EAAKLLGQP	LDLLFSIHTD	360
NNDGTSLGSS	LPPFFGPGCT	LRTALARYAD	LLNPPKKAAL	IALLAAHADEP	SEAERLKFLS	420
SPQKGDEYSK	WVVGSRSLV	EVMAEFPSAK	PPLGVFFAAV	VPRLQPRYYS	ISSSPRFAPH	480
RVHVTALVY	GPTPTGRIHR	GVCSFWMKNV	VPLEKSQNC	WAPIFIRQSN	FKLPADHSVP	540
IVMVGPGTGL	APFRGLQER	LALKEEGAQV	GPALLFFGCR	NRQMDFIYEV	ELNNFVEQGA	600
LSELIVAFSR	EGPSKEYVQH	KMVEKAAYMW	NLISQGGYFY	VCGDAKGMAR	DVHRTLHTIV	660
QEEKVDSTK	AESIVKQLQ	DGRYLQDV				689

SEQ ID NO:99

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SEQ ID NO:100

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VKNNPEQEEV	DLRKIFQSEL	FGLAMRQALG	KDVESLYVED	LKITMNRDEI	FQVLVVDPM	240
GAIDVDWRDF	FPYLKWPVFNK	KFENTIQQMY	IRREAVMKS	LKEHKKRIAS	GEKLSYIDY	300
LLSEAQTLTD	QQLLMSLWEP	IIESSDTTMV	TTEWAMYELA	KNPKLQDRLY	RDIKSVCGSE	360
KITEEHLSQL	PYITAI FHET	LRRHSPVPII	PLRHVHEDTV	LGGYHVPAGT	ELAVNIYGCN	420
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LVLYGSNMG	AEGTARDLAD	IAMSKGFAPQ	VATLDSHAGN	LPREGAVLIV	TASYNHPPD	600
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ADASDDFEGT	YEWEHMHWS	DVAAYFNLDI	ENSEDNKSAL	LLQFVDSAAD	MPLAKMHGAF	720
STNVVASKEL	QPGSARSTR	HLEIELPKEA	SYQEGDHLGV	IPRNYEGIVN	RVTARFGLDA	780
SQQIRLEAEE	EKLAHLPLAK	TVSVEELLQY	VELQDPVTRT	QLRAMAAKT	CPPHKVELEA	840
LLEKQAYKEQ	VLAKRLTMLE	LLEKYPACEM	EFSEFIALLP	SIRPRYSIS	SSPRVDEKQA	900
SITVSVVSGE	AWSGYGEYKG	IASNYLAELQ	EGDITCFIS	TPQSEFTLPK	DPETPLIMVG	960
PGTGVAPFRG	FVQARKQLKE	QGQSLGEAHL	YFGCRSPHED	YLYQEELENA	QSEGIITLHT	1020
AFSRMPNPK	TYVQHVMEQD	GKKLIELLDK	GAHFYICGDG	SQMAPAVEAT	LMKSYADVHQ	1080
VSEADARLWL	QQLEEKGRYA	KDVW				1104

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SEQ ID NO:102

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NAKQFVDWLD	QASADEVKGV	RYSVFGCGDK	NWATTYQKVP	AFIDEMLA AK	GAENIADRGE	660
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STNVVASKEL	QQPGSARSTR	HLEI ELPKEA	SYQEGDHLGV	IPRNYEGIVN	RVTARFGLDA	780
SQOIRLEAEE	EKLAHLPLAK	TVSVEELLQY	VELQDPVTRT	QLRAMAAKT V	CPPHKVELEA	840
LLEKQAYKEQ	VLAKRLTMLE	LLEKY PACEM	EFSEFIALLP	SIRPRYYSIS	SSPRVDEKQA	900
SITVSVVSGE	AWSGYGEYKG	IASNYLAELQ	EGDTITCFIS	TPQSEFTLPK	DPETPLIMVG	960
PGTGVAPFRG	FVQARKQLKE	QGQSLGEAHL	YFGCRSPHED	YLYQEELENA	QSEGIITLHT	1020
AFSRMNPQPK	TYVQHVMEQD	GKKLIELLDK	GAHFYICG DG	SQMAPAVEAT	LMKSYADVHO	1080
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SEQ ID NO:103

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SEQ ID NO:104

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RDEIFQVLV	DPMMGAI	WRDFFPYLKW	VPNKKFENTI	QQMYIRREAV	MKSLIKEHKK	240
RIASGEKINS	YIDYLLSEAQ	TLTDQQLMS	LWEPITIESSD	TTMVTTEWAM	YELAKNPKLQ	300
DRLYRDIKSV	CGSEKITEEH	LSQLPYITAI	FHETLRHRSP	VPIPLRHVH	EDTVLGGYHV	360
PAGTELAVNI	YGCNMDKNVW	ENPEEWNPER	FMKENETIDF	QKTMAFGGGK	RVCAGSLQAL	420
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LAAKGAENIA	DRGEADASDD	FEGTYEEWRE	HMWSDVAAYF	NLDIENSEDN	KSALLLQFVD	660
SAADMPLAKM	HGAFSTNVVA	SKELQQPGSA	RSTRHLEIEL	PKEASYQEGD	HLGVI PRNVE	720
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SEQ ID NO:105

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SEQ ID NO:106

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LVMTRFQSIST	RNLKALKKVL	TADKTMVAMS	DYDDYHKTVK	RHILTAVLGP	NAQKKHRIHR	120
DIMMNDNISTQ	LHEFVKNNPE	QEEVDLRKIF	QSEFLGLAMR	QALGKDVESL	YVEDLKITMN	180
RDEIFQVLVV	DPMMGAIDVD	WRDFFPYLKW	VPNKKFENTI	QQMYIRREAV	MKSLLIKEHKK	240
RIASGEKINS	YIDYLLSEAQ	TLTDQQLMS	LWEPIESSD	TTMVTTEWAM	YELAKNPKLQ	300
DRLYRDIKSV	CGSEKITEEH	LSQLPYITAI	FHETLRRHSP	VPIIPLRHVH	EDTVLGGYHV	360
PAGTELAVNI	YGCNMDKNVW	ENPEEWNPER	FMKENETIDF	QKTMAFGGGK	RVCAGSLQAL	420
LTASIGIGRM	VQEFWKLKD	MTQEEVNTIG	LTTQMLRPLR	AIIKPRIPSR	PSPSTEQSAK	480
KVRKKAENAH	NTPLLVLVYGS	NMGTAEGTAR	DLADIAMSKG	FAPQVATLDS	HAGNLPREGA	540
VLIVTASYNG	HPPDNAKQFV	DWLDQASADE	VKGVRYSVFG	CGDKNWATTY	QKVPAFIDEM	600
LAAKGAENIA	DRGEADASDD	FEGTYEEWRE	HMWSDVAAYF	NLDIENSEDN	KSALLLQFVD	660
SAADMPLAKM	HGAFSTNVVA	SKELQPGGSA	RSTRHLEIEL	PKEASYQEGD	HLGVI PRNVE	720
GIVNRVTARF	GLDASQQIRL	EAEEEEKLAHL	PLAKTVSVEE	LLQYVELQDP	VTRTQLRAMA	780
AKTVCPPHKV	ELEALLEKQA	YKEQVLAKRL	TMLELLEKYP	ACEMEFSEFI	ALLPSIRPRY	840
YSISSSPRVD	EKQASITVSV	VSGEAWSGYG	EYKGIASNYL	AELQEGDIT	CFISTPQSEF	900
TLPKDPETPL	IMVGPGTGVA	FRGFVQARK	QLKEQQGSLG	EAHLYFGCRS	PHEDYLYQEE	960
LENAQSEGII	TLHTAFSRMP	NRPKTYVQHV	MEQDGKLLIE	LLDKGAHFYI	CGDGSQMAPA	1020
VEATLMKSYA	DVHQVSEADA	RLWLQOLEEK	GRYAKDVA			1058

SEQ ID NO:107

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SEQ ID NO:108

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KILFADKCMV	AISDYNDFHK	MIKRYILSNV	LGPSAQKRHR	SNRDTLRANV	CSRLHSQVKN	180
SPREAVNFR	VFEWELFGIA	LKQAFGKDIE	KPIYVEELGT	TLRDEIFKV	LVLDIMEGAI	240
EVDWRDFFPY	LRWIPNTRME	TKIQRLYFRR	KAVMTALINE	QKKRIASGEE	INCYIDFLK	300
EGKTLTMDQI	SMLLWETVIE	TADTMTVTE	WAMYEVAKDS	KRQDRLYQEI	QKVCSEMVT	360
EEYLSQLPYL	NAVFHETLRK	HSPAALVPLR	YAHEDTQLGG	YYIPAGTEIA	INIYGCNMDK	420
HQWESPEEWK	PERFLDPKFD	PMDLYKTMAF	GAGKRCVAGS	LQAMLIACPT	IGRLVQEFEW	480
KLRDGEENNV	DTVGLTTHKR	YPMHAILKPR	SPSRPSFSTE	QSAKKVRKKA	ENAHNTPLL	540
LYGSNMGTAE	GTARDLADIA	MSKGFAPQVA	TLDSHAGNLP	REGAVLIVTA	SYNGHPPDNA	600
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ASDDFEGTYE	EWREHMWSDV	AAFYNLDIEN	SEDNKSALLL	QFVDSAADMP	LAKMHGAFST	720
NVVASKELQQ	PGSARSTRHL	EIELPKEASY	QEGDHLGVIP	RNYEGIVNRV	TARFGLDASQ	780
QIRLEAEEEK	LAHLPLAKTV	SVEELLQYVE	LQDPVTRTQL	RAMAAKTVC	PHKVELEALL	840
EKQAYKEQVL	AKRLTMLELL	EKYPACEMEF	SEFIALLP	RPRYYSISS	PRVDEKQASI	900
TVSVSAGEAW	SGYGEYKZIA	SNYLAEQEG	DTITCFISTP	QSEFTLPKDP	ETPLIMVPG	960
TGVAPFRGFV	QARKQLKEQG	QSLGEAHLF	GCRSPHEDYL	YQEELENAQS	EGIITLHTAF	1020
SRMPNQPKTY	VQHVMEQDQK	KLIELLDKGA	HFYICDGSQ	MAPAVEATLM	KSYADVHQVS	1080
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SEQ ID NO:109

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SEQ ID NO:110

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KILTADKCMV	AISDYNDPHK	MIKRYILSNV	LGPSAQKRHR	SNRDTLRANV	CSRLHSQVKN	180
SPREAVNFR	VFWELEFGIA	LKQAFGKDIE	KPIYVEELGT	TLRDEIFKV	LVLDIMEGAI	240
EVDWRDFFPY	LRWIPNTRME	TKIQRLYFR	KAVMTALINE	QKKRIASGEE	INCYIDFLK	300
EGKTLTMDQI	SMLLWETVIE	TADTTMVTTE	WAMYEVAKDS	KRQDRLYQEI	QKVCSEMVT	360
EEYLSQLPYL	NAVFHETLRK	HSPAALVPLR	YAHEDTQLGG	YYIPAGTEIA	INITYCNMDK	420
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KLRDGEENV	DTVGLTTHKR	YPMHAILKPR	SPSRPSPSTE	QSACKVRKKA	ENAHNTPLL	540
LYGSNMGTA	E GTARDLADIA	MSKGFAPQVA	TLD SHAGNLP	REGAVLIVTA	SYNGHPPDNA	600
KQFVDWLDQA	SADDEVKGVRY	SVFGCGDKNW	ATYQKVPF	IDEMLAAKGA	ENIADRGEAD	660
ASDDFEGTYE	EWREHMWSDV	AAYFNLDIEN	SEDNKSALLL	QFVDSAADMP	LAKMHGAFST	720
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QIRLEAEEEEK	LAHLPLAKTV	SVEELLQYVE	LQDPVTRTQL	RAMAAKTVC	PHKVELEALL	840
EKQAYKEQVL	AKRLTMELEL	EKYPACEMEF	SEPIALLPSI	RPRYYSISS	PRVDEKQASI	900

TVSVVSGEAW	SGYGEYKZIA	SNYLAELQEG	DTITCFISTP	QSEFTLPKDP	ETPLIMVGP	960
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SEQ ID NO:111

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SEQ ID NO:112

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KVLVLDIMEG	AIEVDWRDF	PYLRIWPNTR	METKIQRLYF	RRKAVMTALI	NEQKKRIASG	240
EEINCYIDFL	LKEGKTLTMD	QISMLLWETV	IETADTMVT	TEWAMYEVAK	DSKRQDRLYQ	300
EIQKVCGSEM	VTEEYLSQLP	YLNNAVPHETL	RKHSPAALVP	LRyahEDTQL	GGYYPAGTE	360
IAINIYGCNM	DKHQWESPEE	WKPERFLDPK	FDPMDLYKTM	AFGAGKRVCA	GSLQAMLIAC	420
PTIGRLVQEF	EWKLRDGEEE	NVDTVGLTTH	KRYPMHAILK	PRSPSRPSPS	TEQSAKVRK	480
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GAENIADRGE	ADASDDFEGT	YEEWREHMWS	DVAAYFNLDI	ENSEDNKSAL	LLQFVDSAAD	660
MPLAKMHGAF	STNVVASKEL	QQPGSARSTR	HLEIELPKEA	SYQEGDHLGV	IPRNYEGIVN	720
RVTARFGLDA	SQIRLEAEE	EKLAHLPLAK	TVSVEELLQY	VELQDPVTRT	QLRAMAAKTV	780
CPPHKVELEA	LLEKQAYKEQ	VLAKRLTMLE	LLEKYFACEM	EFSEFIALLP	SIRPRYYSIS	840
SSPRVDEKQA	SITVSVVSGE	AWSGYGEYKG	IASNYLAELQ	EGDTITCFIS	TPQSEETLPK	900
DPETELIMVG	PGTGVAPFRG	FVQARKQLKE	QGQSLGEAHL	YFGCRSPHED	YLYQEELENA	960
QSEGIITLHT	AFSRMPNQPK	TYVQHVMEQD	GKKLIELLDK	GAHFYICGDG	SQMAPAVEAT	1020
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SEQ ID NO:113

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SEQ ID NO:114

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NSNTWISSL	LTSGGVITAS	LYLYRIYVT	PIWPLSIQTA	SLLGFLSMV	CGLGLYGIS	240
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NVILKNAEVL	WGSFIPRGRK	KTGDFHDKLI	SILSFEKVSL	ISKPFWKFFK	NFTFSVPLSI	360
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SEQ ID NO:115

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SEQ ID NO:116

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SEQ ID NO:117

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SEQ ID NO:118

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KSALLLQFVD	SAADMPLAKM	HGAFSTNVVA	SKELQQPGSA	RSTRHLEIEL	PKEASYQEGD	240
HLGVI PRNVE	GIVNVRTARF	GLDASQQIRL	EAEEEKLAHL	PLAKTVSVEE	LLQYVELQDP	300
VTRTQLRAMA	AKTVCPPHKV	ELEALLEKQA	YKEQVLAKRL	TMLELLEKYP	ACEMEFSEFI	360
ALLPSIRPRY	YSISSSPRVD	EKQASITVSV	VSGEAWSGYG	EYKGIASNYL	AELQEGDTIT	420
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SEQ ID NO:119

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SEQ ID NO:120

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VTRTQLRAMA	AKTVCPPHKV	ELEALLEKQA	YKEQVLAKRL	TMLELLEKYP	ACEMEFSEFI	360
ALLPSIRPRY	YSISSSPRVD	EKQASITVSV	VSGEAWSGYG	EYKGIASNYL	AELQEGDTIT	420
CFISTPQSEF	TLPKDPETPL	IMVGPGTGVA	PFRGFVQARK	QLKEQQQSLG	EAHLYFGCRS	480
PHEDYLYQEE	LENAQSEGII	TLHTAFSRMP	NQPKTYVQHV	MEQDGKKLIE	LLDKGAHFYI	540
CGDGSQMAPA	VEATLMKSYA	DVHQVSEADA	RLWLQOLEEK	GRYAKDVA		588

SEQ ID NO:121

ccatcaaga

9



SEQ ID NO:122

PSR

3

**WHAT IS CLAIMED IS:**

1. A recombinant host comprising one or more of:
  - (a) a gene encoding an ent-kaurene oxidase (KO) polypeptide;
  - (b) a gene encoding a cytochrome P450 reductase (CPR) polypeptide;  
and/or
  - (c) a gene encoding an ent-kaurenoic **acid** hydroxylase (KAH) polypeptide;wherein at least one of the genes is a recombinant gene; and  
wherein the recombinant host is capable of producing a steviol glycoside precursor.
  
2. A recombinant host comprising:
  - (a) a gene encoding a geranylgeranyi diphosphate synthase (GGPPS) polypeptide;
  - (b) a gene encoding an ent-copalyl diphosphate synthase (CDPS) polypeptide;
  - (c) a gene encoding an ent-kaurene synthase (KS) polypeptide
  - (d) a gene encoding an ent-kaurene oxidase (KO) polypeptide;
  - (e) a gene encoding a cytochrome P450 reductase (CPR) polypeptide; and
  - (f) a gene encoding an ent-kaurenoic acid hydroxylase (KAH) polypeptide;wherein at least one of the **genes** is a recombinant gene; and  
wherein the recombinant host is capable of producing steviol.
  
3. The recombinant host of claims 1 or 2, wherein:
  - (a) the KO polypeptide comprises a KO polypeptide having at least 60% identity to an amino acid sequence set forth in SEQ ID NO:72 or SEQ ID NO:75; at least 65% identity to an amino acid sequence set forth in SEQ ID NO:54; at least 70% identity to an amino acid sequence set forth in SEQ ID NO: 70, SEQ ID NO:71, or

SEQ ID NO:79; at least 40% identity to an amino acid sequence set forth in SEQ ID NO:77; or at least 50% identity to an amino acid sequence set forth in SEQ ID NO:78;

- (b) the CPR polypeptide comprises a CPR polypeptide having at least 70% identity to an amino acid sequences set forth in SEQ ID NO:69, SEQ ID NO:74, SEQ ID NO:76, or SEQ ID NO:87; at least 80% identity to an amino acid sequence set forth in SEQ ID NO:73; at least 85% identity to an amino acid sequence set forth in SEQ ID NO:22; at least 65% identity to an amino acid sequence set forth in SEQ ID NO:28; or at least 50% identity to an amino acid sequence set forth in SEQ ID NO:98; and/or
- (c) the KAH polypeptide comprises a KAH polypeptide having at least 40% identity to an amino acid sequence set forth in SEQ ID NO:82; at least 50% identity to an amino acid sequence set forth in SEQ ID NO:91 ; or at least 60% identity to an amino acid sequence set forth in SEQ ID NO:68.

4 . A recombinant host comprising one or more of:

- (a) a gene encoding a KO polypeptide having at least 60% identity to an amino acid sequence set forth in SEQ ID NO:75;
- (b) a gene encoding a KAH polypeptide having at least 40% identity to an amino acid sequence set forth in SEQ ID NO:82; and/or
- (c) a gene encoding a CPR polypeptide having at least 50% identity to an amino acid sequence set forth in SEQ ID NO:98;

wherein at least one of the **genes** is a recombinant gene; and

wherein the recombinant host is capable of producing a steviol glycoside precursor.

5. A recombinant host comprising one or more of:

- (a) a gene encoding a KO polypeptide having at least 70% identity to an amino acid sequence set forth in SEQ ID NO:70;

- (b) a gene encoding a KAH polypeptide having at least 40% identity to an amino acid sequence set forth in SEQ ID NO:82; and/or
- (c) a gene encoding a CPR polypeptide having at least 50% identity to an amino acid sequence set forth in SEQ ID NO:98;

wherein at least one of the genes is a recombinant gene; and

wherein the recombinant host is capable of producing a steviol glycoside precursor.

- 6. The recombinant host of claim 4 or 5, wherein the host further comprises a gene encoding a KO polypeptide having at least 65% identity to an amino acid sequence set forth in SEQ ID NO:54.
- 7. The recombinant host of any one of claims 4-6, wherein the host further comprises a gene encoding a KAH polypeptide having at least 60% identity to an amino acid sequence set forth in SEQ ID NO:68.
- 8. The recombinant host of any one of claims 4-7, wherein the host further comprises a gene encoding a KO polypeptide having at least 70% identity to an amino acid sequence set forth in SEQ ID NO:79.
- 9. The recombinant host of any one of claims 1 or 3-8, wherein the host further comprises one or more of:
  - (a) a gene encoding a geranylgeranyl diphosphate synthase (GGPPS) polypeptide;
  - (b) a gene encoding an ent-copalyl diphosphate synthase (CDPS) polypeptide; and/or
  - (c) a gene encoding an ent-kaurene synthase (KS) polypeptide;

wherein at least one of the genes is a recombinant gene; and

wherein the recombinant host is capable of producing a steviol glycoside precursor.

10. The recombinant host of claim 9, wherein:
  - (a) the GGPPS polypeptide comprises a polypeptide having at least 70% identity to an amino acid sequence set forth in SEQ ID NO:49;
  - (b) the CDPS polypeptide comprises a polypeptide having at least 70% identity to an amino acid sequence set forth in SEQ ID NO:37; and/or
  - (c) the KS polypeptide comprises a polypeptide having at least 40% identity to an amino acid sequence set forth in SEQ ID NO:6.
11. The recombinant host of claims 1-10, wherein the host further comprises a gene encoding an endoplasmic reticulum membrane polypeptide.
12. The recombinant host of claim 11, wherein the endoplasmic reticulum membrane polypeptide comprises an Inheritance of cortical ER protein 2 (ICE2) polypeptide having at least 50% identity to the amino acid sequence set forth in SEQ ID NO:114.
13. The recombinant host of any one of claim 1-10, wherein the KO polypeptide is a fusion construct.
14. The recombinant host of claim 13, wherein the fusion construct comprises a polypeptide having at least 60% identity to an amino acid sequence set forth in SEQ ID NO:118 or SEQ ID NO:120.
15. The recombinant host of claim 13 or claim 14, wherein the fusion construct has at least 50% identity to an amino acid sequence set forth in SEQ ID NO:100, SEQ ID NO:102, SEQ ID NO:104, SEQ ID NO:106, SEQ ID NO:108, SEQ ID NO:110, or SEQ ID NO:112.

16. The recombinant host of any one of claims 1-15, wherein the host further comprises one or more of:

- (a) a gene encoding a UGT85C polypeptide;
- (b) a gene encoding a UGT76G polypeptide;
- (c) a gene encoding a UGT74G1 polypeptide;
- (d) a gene encoding a UGT91 D2 functional homolog polypeptide; and/or
- (e) a gene encoding an EUGT11 polypeptide;

wherein at least one of the genes is a recombinant gene; and

wherein the host is capable of producing a steviol glycoside.

17. The recombinant host of claim 16, wherein:

- (a) the UGT85C2 polypeptide comprises a polypeptide having at least 55% identity to an amino acid sequence set forth in SEQ ID NO:30;
- (b) the UGT76G1 polypeptide comprises a polypeptide having at least 50% identity to an amino acid sequence set forth in SEQ ID NO:83;
- (c) the UGT74G1 polypeptide comprises a polypeptide having at least 55% identity to an amino acid sequence set forth in SEQ ID NO:29;
- (d) the UGT91 D2 functional homolog polypeptide comprises a UGT91D2 polypeptide having 90% or greater identity to the amino acid sequence set forth in SEQ ID NO:84 or a UGT91D2e-b polypeptide having 90% or greater identity to the amino acid sequence set forth in SEQ ID NO:88; and/or
- (e) the EUGT11 polypeptide comprises a polypeptide having at least 65% identity to an amino acid sequence set forth in SEQ ID NO:86.

18. The recombinant host of any one of claims 1-17, wherein the recombinant host comprises a plant cell, a mammalian cell, an insect cell, a fungal cell, or a bacterial cell.

- 1<sup>19</sup>. The recombinant host of claim 18, wherein the bacterial cell comprises *Escherichia* bacteria cells, *Lactobacillus* bacteria cells, *Lactococcus* bacteria cells, *Cornebacterium* bacteria cells, *Acetobacter* bacteria cells, *Acinetobacter* bacteria cells, or *Pseudomonas* bacterial cells.
20. The recombinant host of claim 18, wherein the fungal cell comprises a yeast cell.
- 2 1<sup>1</sup> . The recombinant host of claim 20, wherein the yeast cell is a cell from *Saccharomyces cerevisiae*, *Schizosaccharomyces pombe*, *Yarrowia lipolytica*, *Candida glabrata*, *Ashbya gossypii*, *Cyberlindnera jadinii*, *Pichia pastoris*, *Kluyveromyces lactis*, *Hansenula polymorpha*, *Candida boidinii*, *Arxula adenivorans*, *Xanthophyllomyces dendrorhous*, or *Candida albicans* species.
22. The recombinant host of claim 21, wherein the yeast cell is a *Saccharomycete*.
23. The recombinant host of claim 22, wherein the yeast cell is a cell from the *Saccharomyces cerevisiae* species.
24. A method of producing a steviol glycoside or a steviol glycoside precursor, comprising:
- (a) growing the recombinant host of any one of claims 1-23 in a culture medium, under conditions in which any of the genes disclosed in any one of claims 1-23 are expressed;  
wherein the **steviol** glycoside or the steviol glycoside precursor is synthesized by said host; and/or
  - (b) optionally quantifying the steviol glycoside or the steviol glycoside precursor; and/or
  - (c) optionally isolating the steviol glycoside or the steviol glycoside precursor.

25. The method of claim 24, wherein the steviol glycoside comprises steviol-1 3-O-glucoside (13-SMG), steviol-1 ,2-bioside, steviol-1,3-bioside, steviol-19-O-glucoside (19-SMG), stevioside, 1,3-stevioside, rubusoside, Rebaudioside A (RebA), Rebaudioside B (RebB), Rebaudioside C (RebC), Rebaudioside D (RebD), Rebaudioside E (RebE), Rebaudioside F (RebF), Rebaudioside M (RebM), Rebaudioside Q (RebQ), Rebaudioside I (RebI), dulcoside A, di-glycosylated steviol, tri-glycosylated steviol, tetra-glycosylated steviol, penta-glycosylated steviol, hexa-glycosylated steviol, hepta-glycosylated steviol, or isomers thereof.
26. The steviol glycoside or the steviol glycoside precursor produced by the recombinant host of any one of claims 1-23 or the method of claim 24 or claim 25, wherein the steviol glycoside or steviol glycoside precursor accumulates to a detectable concentration when cultured under said conditions.
27. A steviol glycoside composition produced by the host of any one of claims 1-23 or the method of claim 24 or claim 25, wherein the composition has an undetectable concentration of stevia plant-derived contaminants.
28. A steviol glycoside composition produced by the host of any one of claims 1-23 or the method of claim 24 or claim 25, wherein the composition has a steviol glycoside composition enriched for RebD or RebM relative to the steviol glycoside composition of a wild-type Stevia plant.



Figure 1

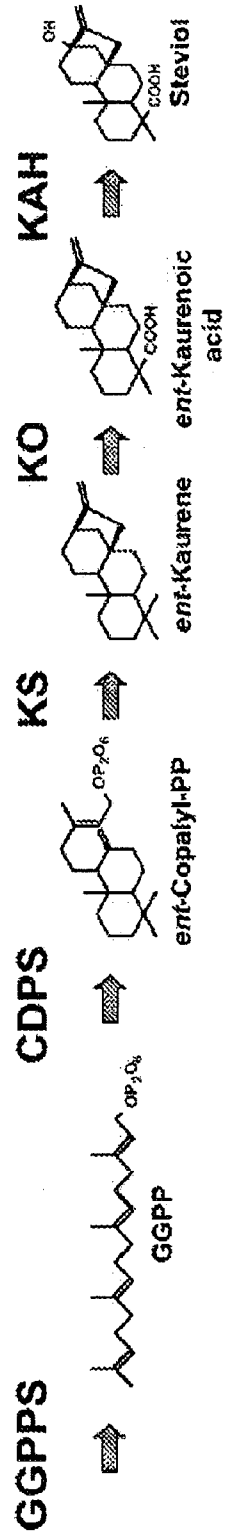




Figure 3

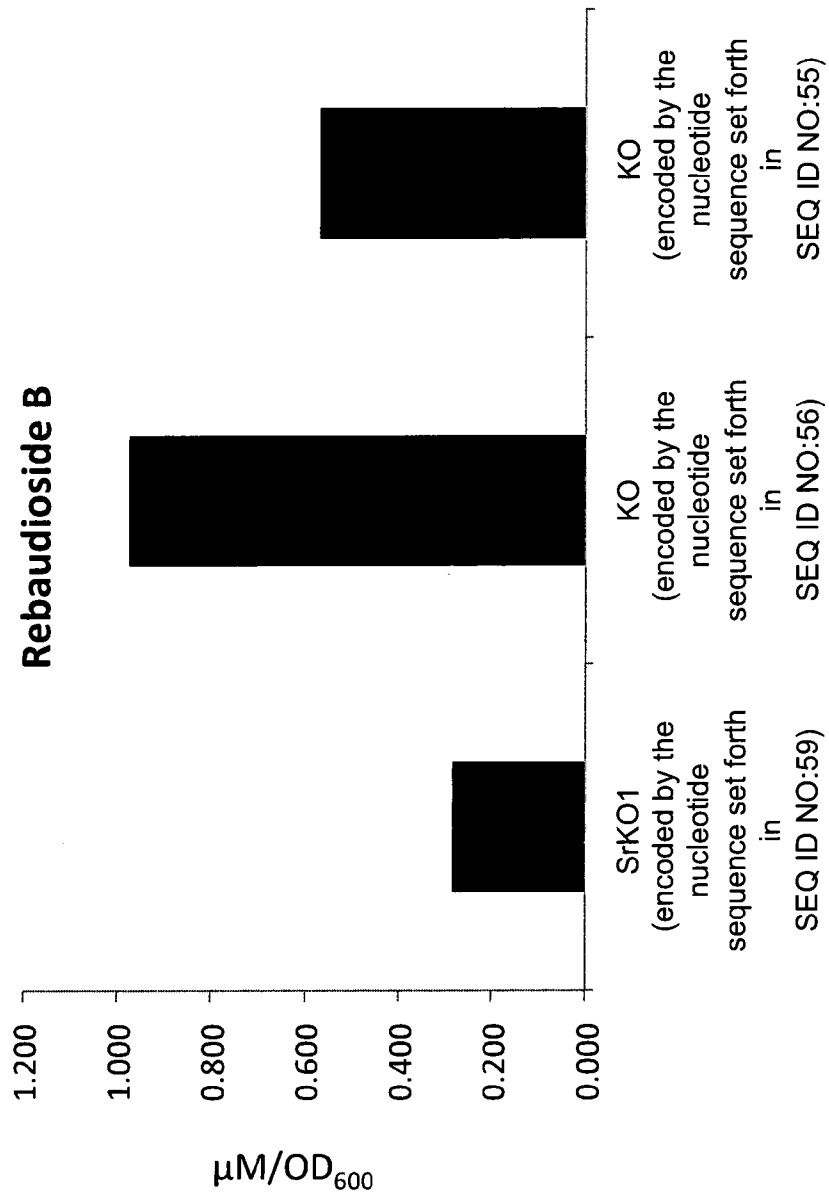
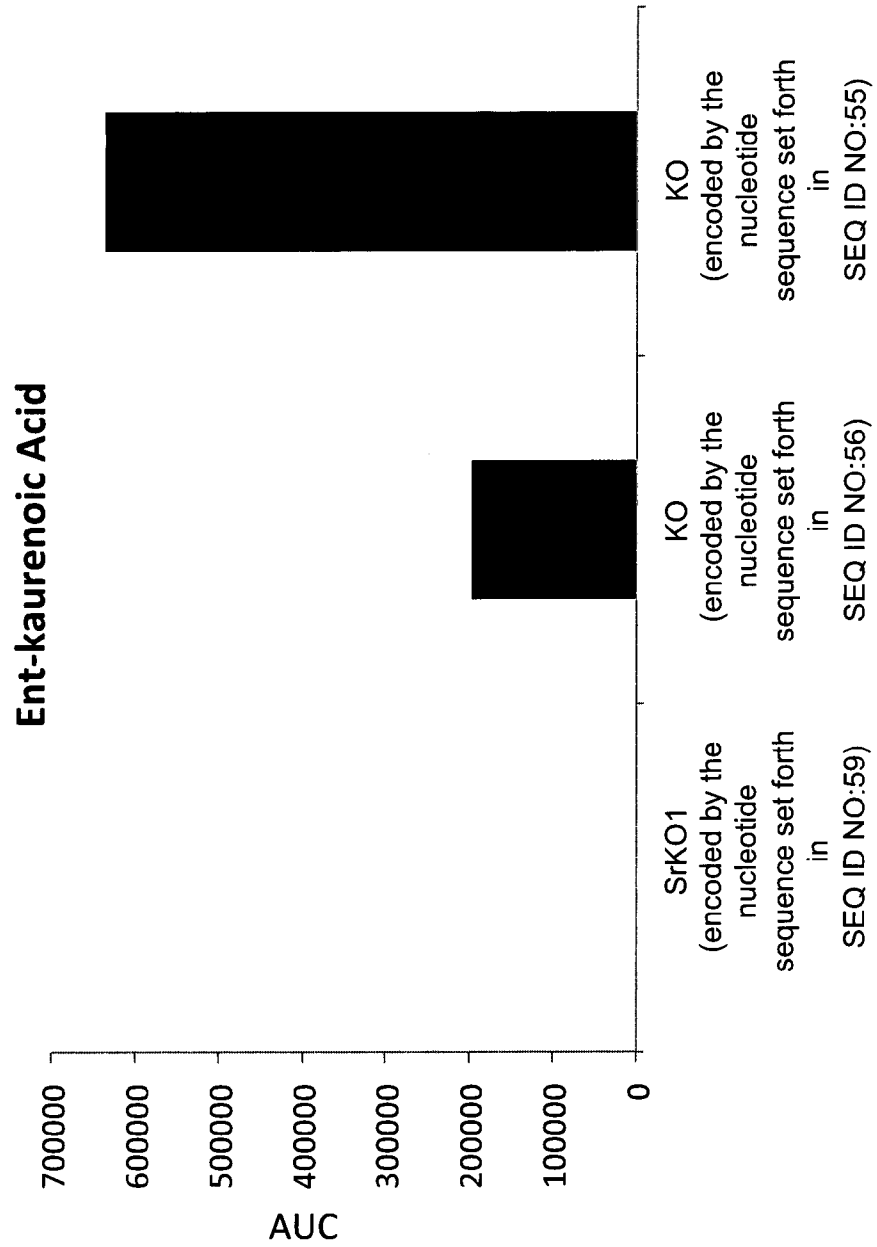


Figure 4



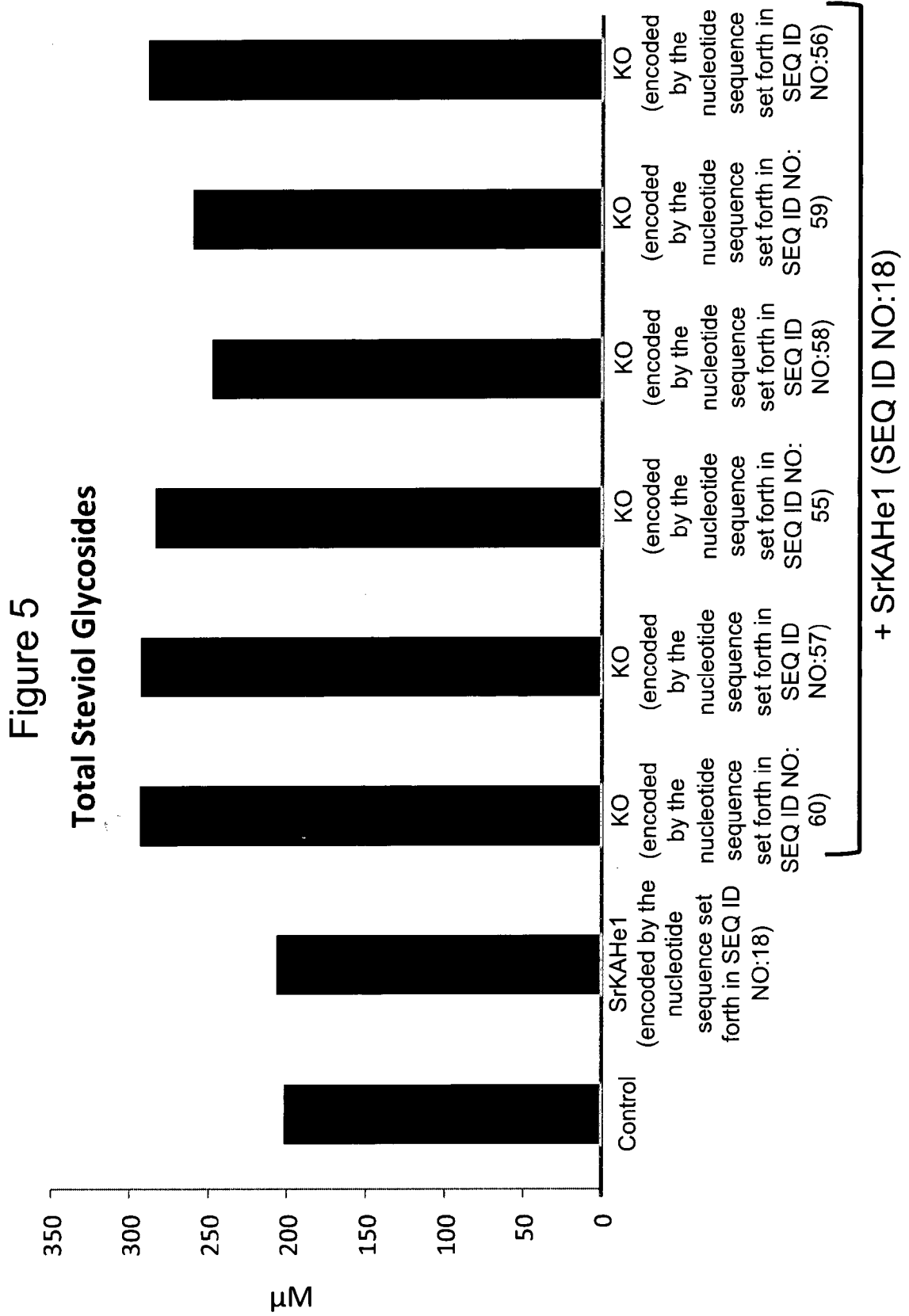


Figure 6

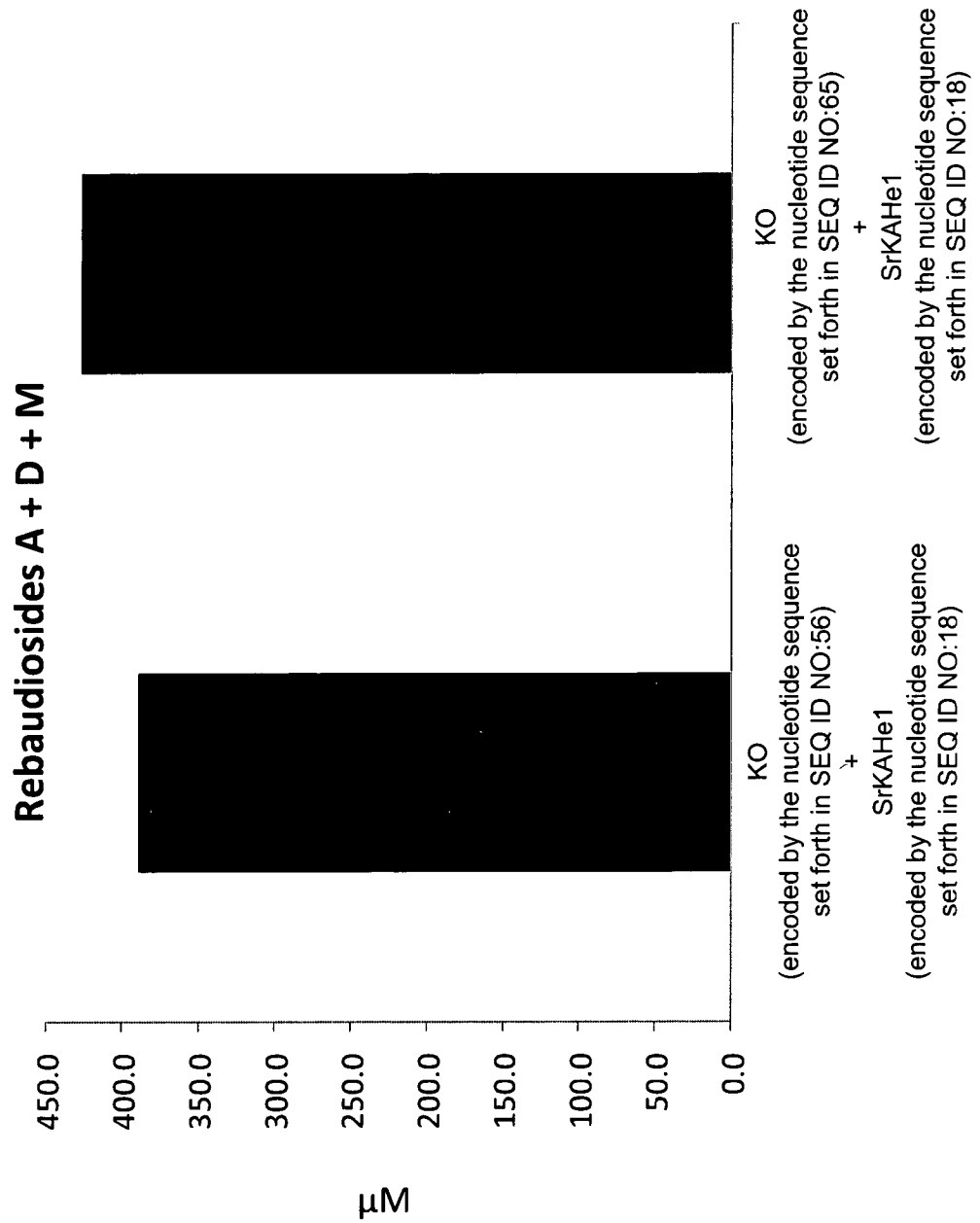


Figure 7

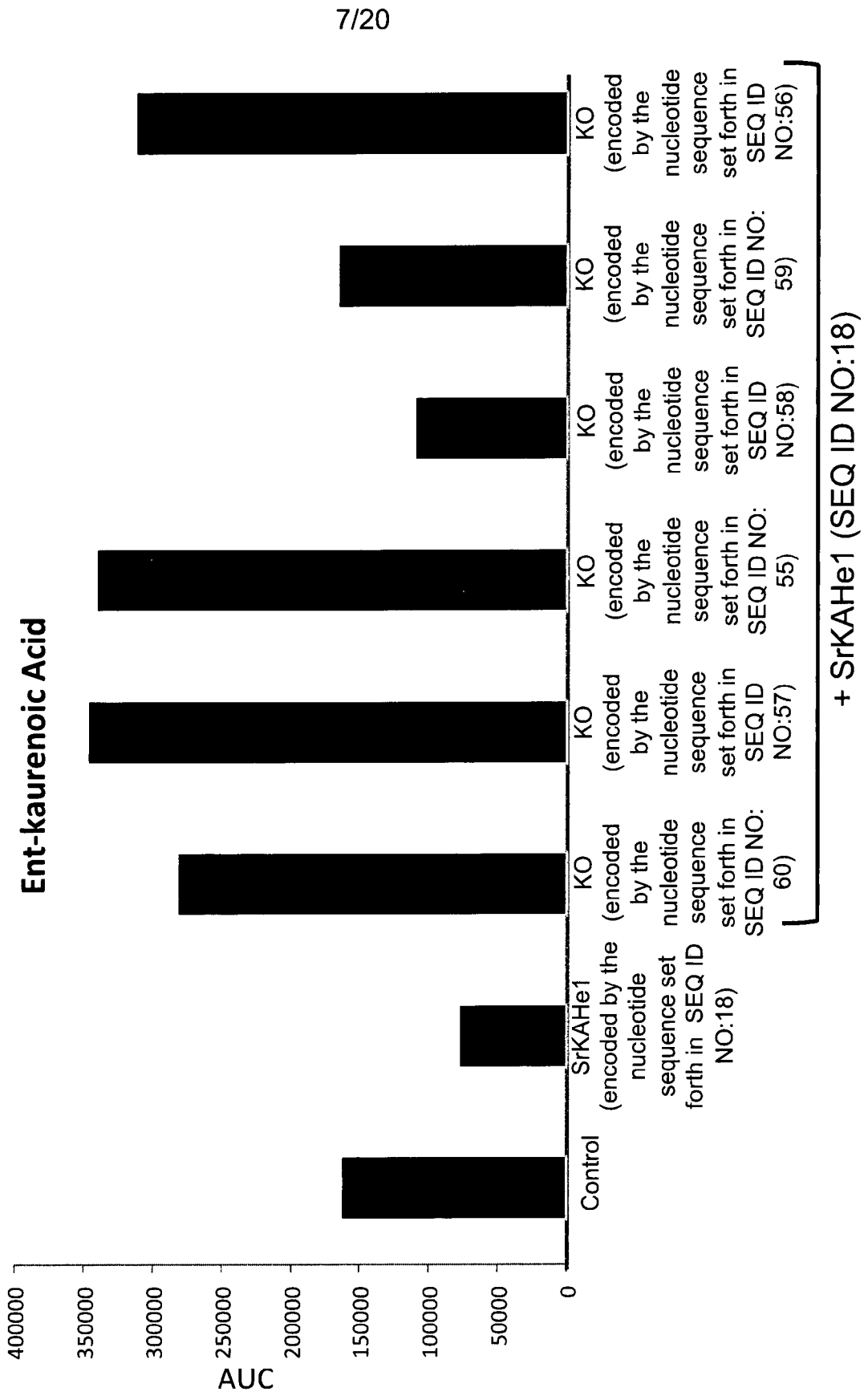
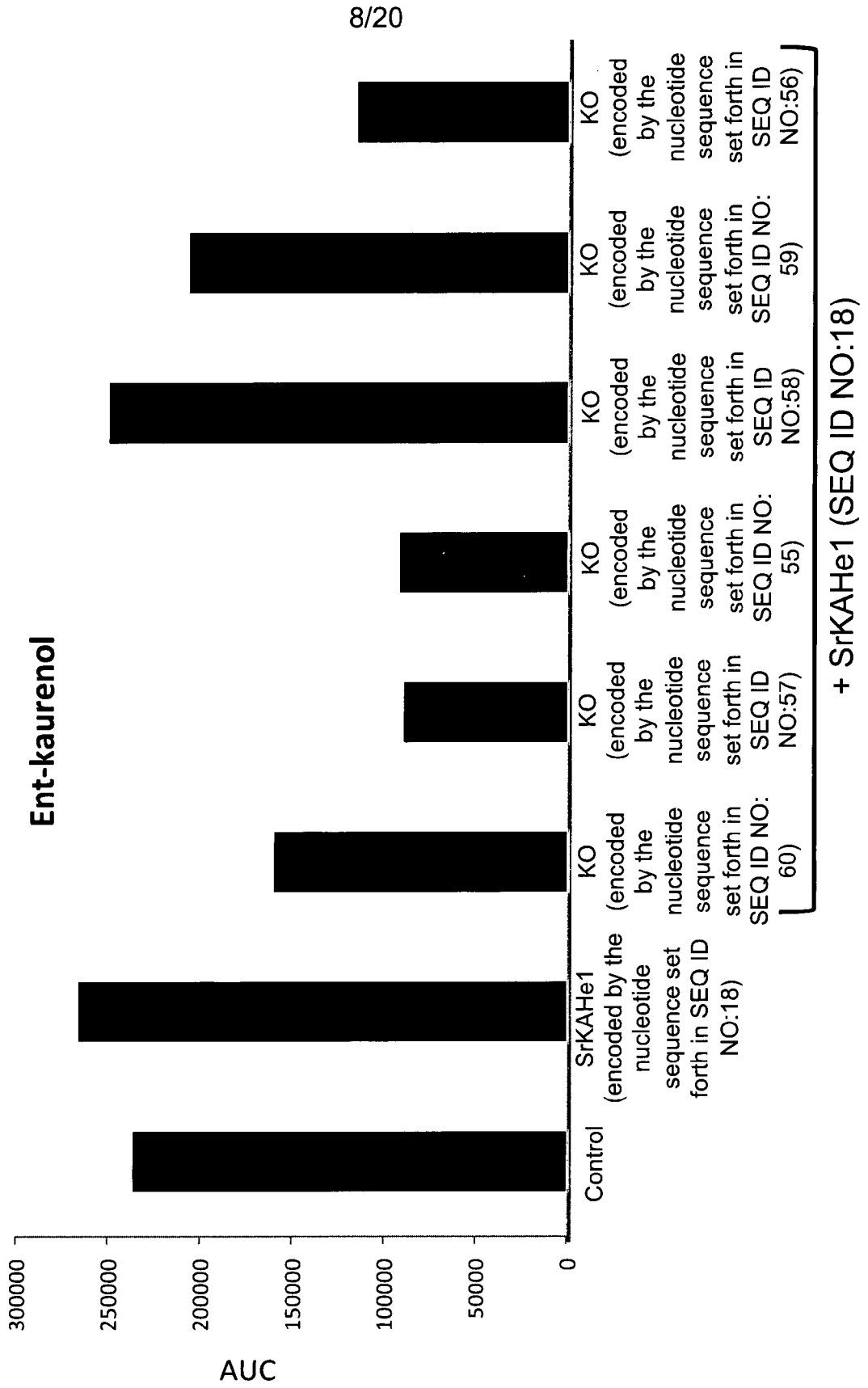


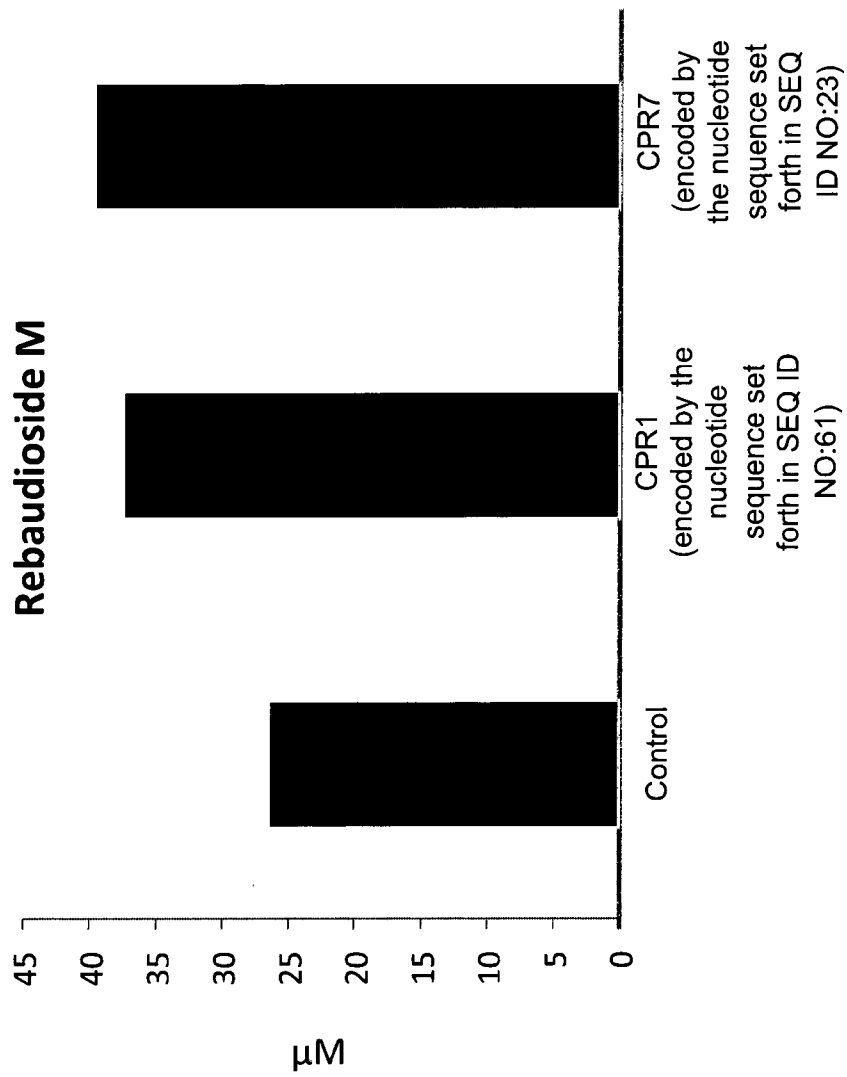
Figure 8  
Ent-kaurenol





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Figure 9



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Figure 10

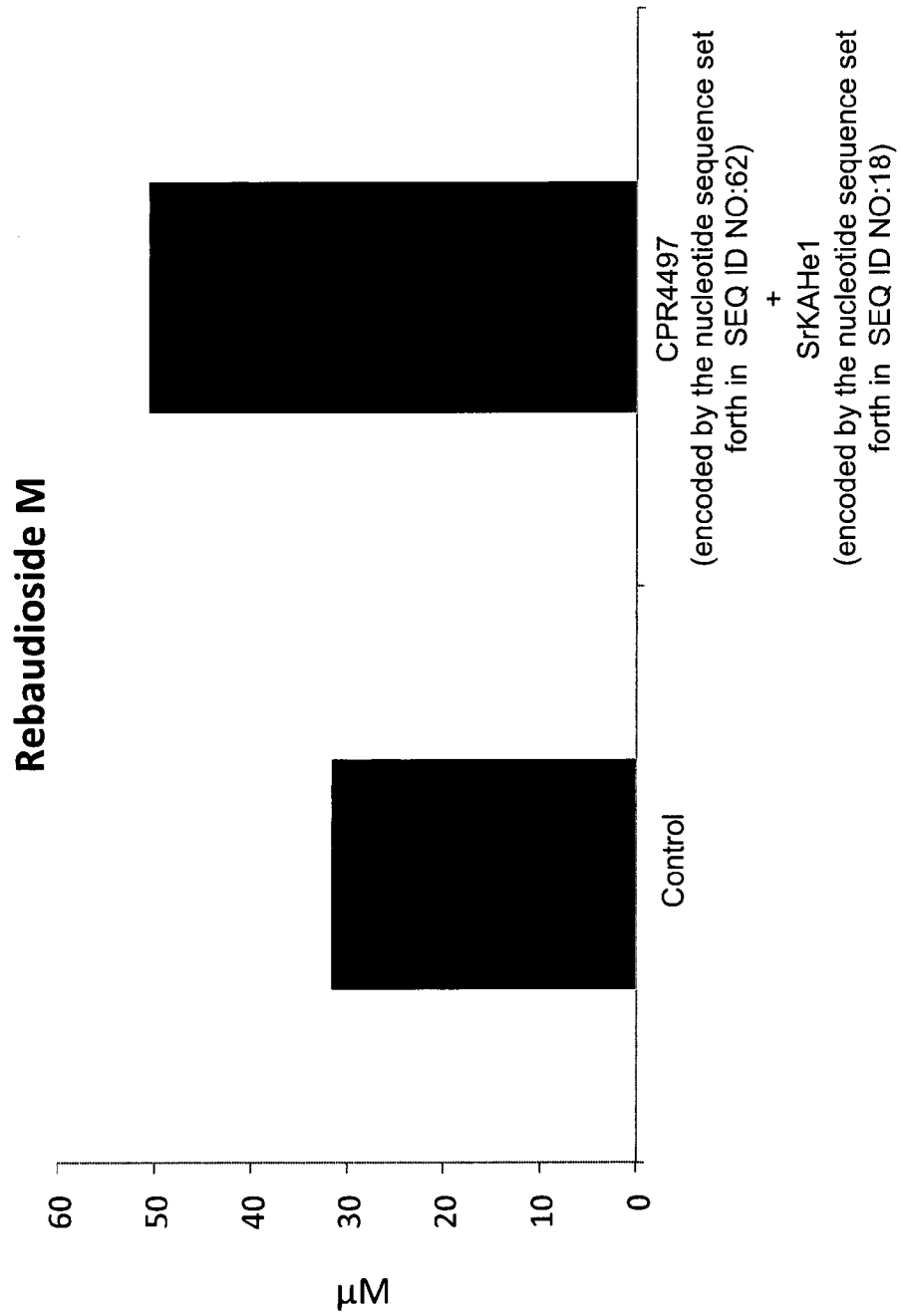


Figure 11A

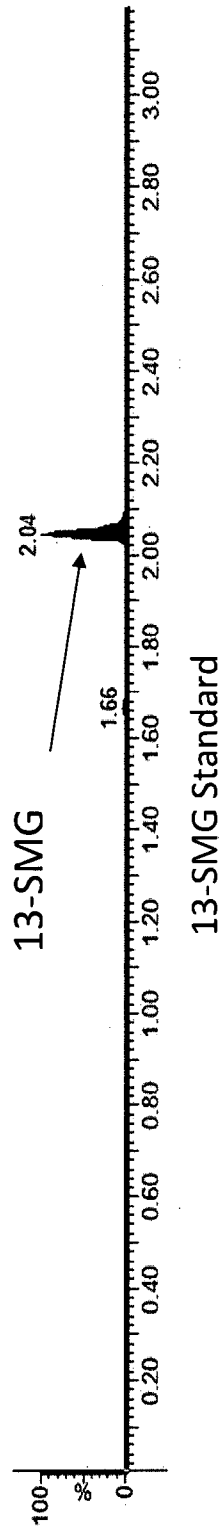
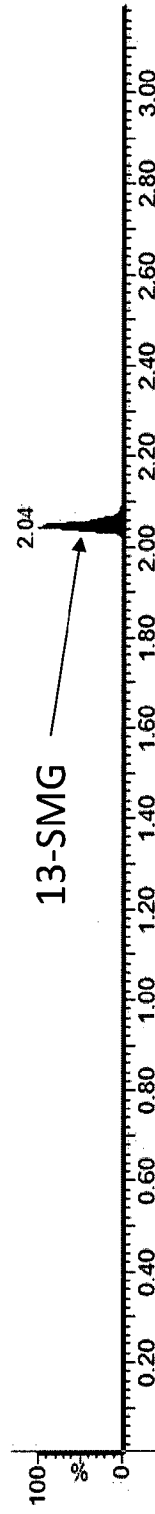
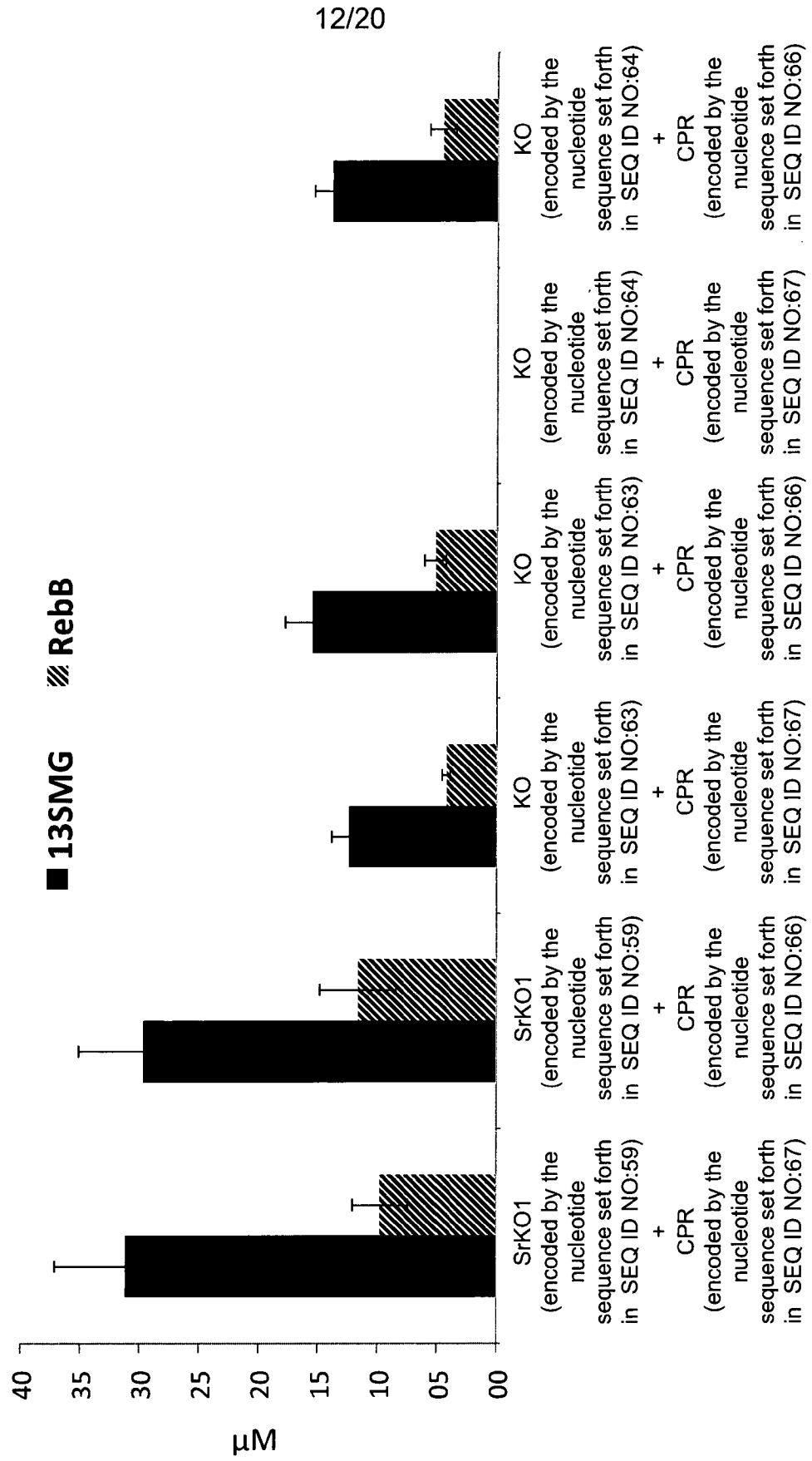


Figure 11B



*S. cerevisiae* expressing KAH encoded by nucleotide sequence set forth in SEQ ID NO:80

Figure 12



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Figure 13

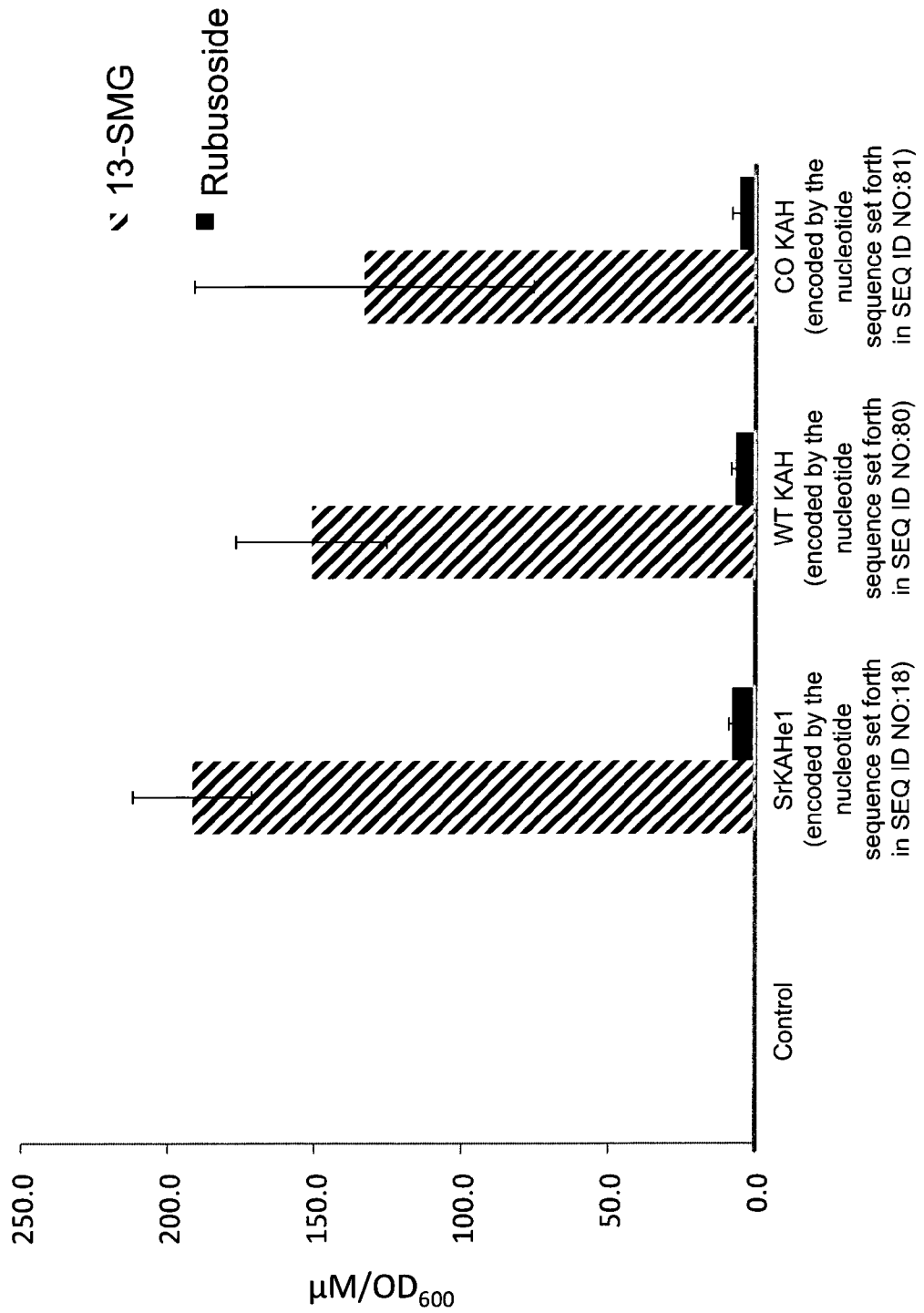


Figure 14

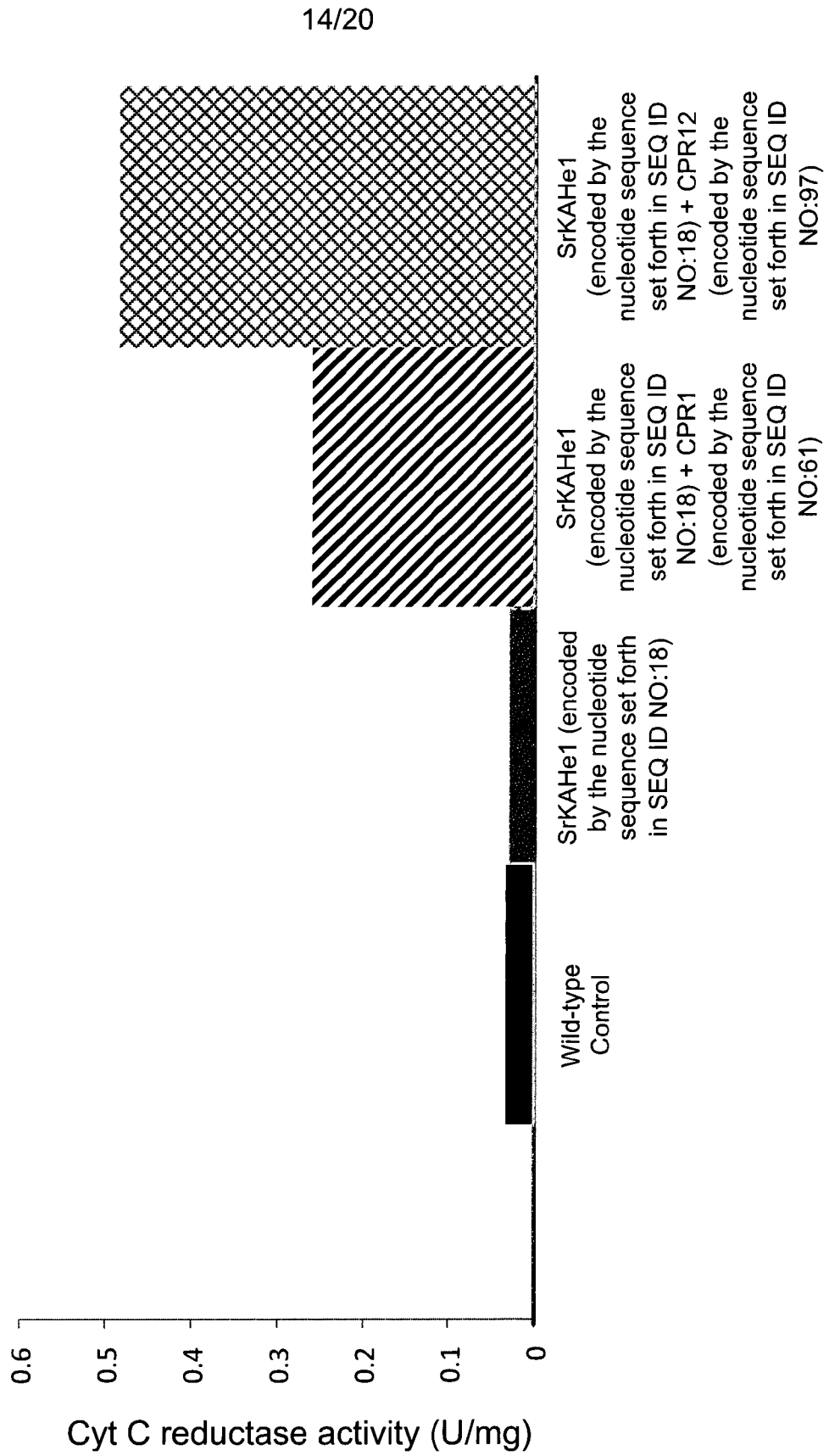


Figure 15A

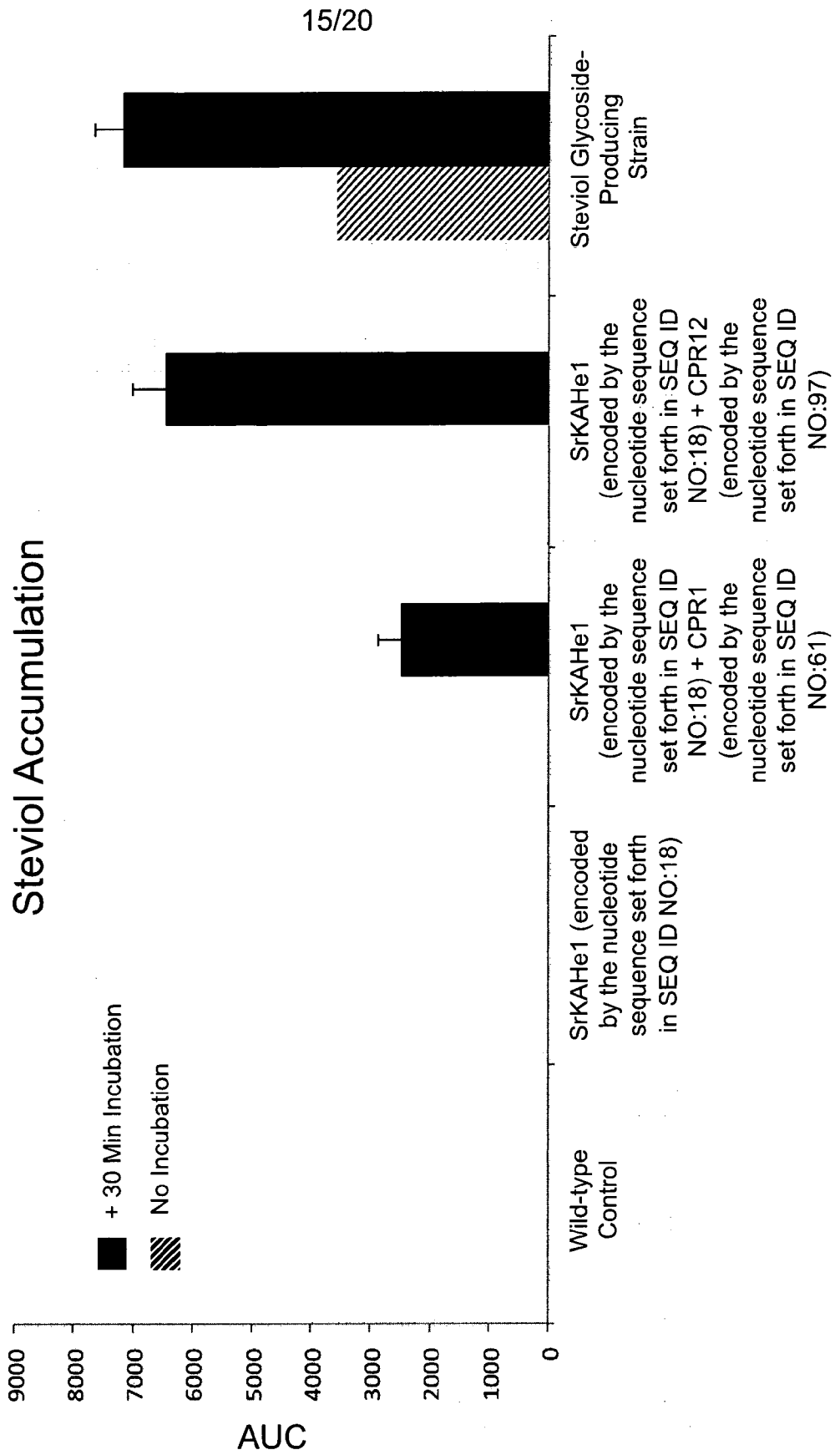


Figure 15B

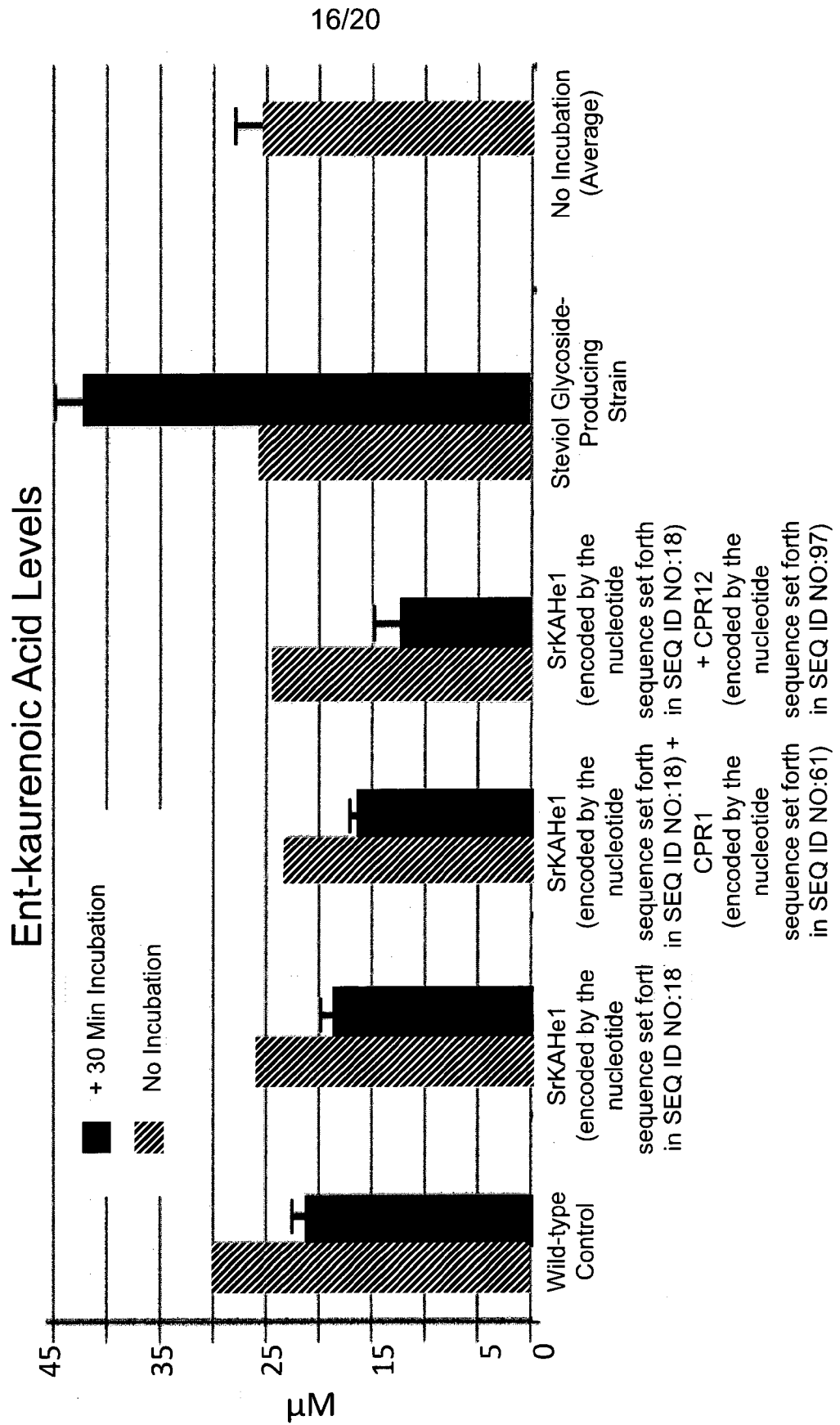




Figure 16A

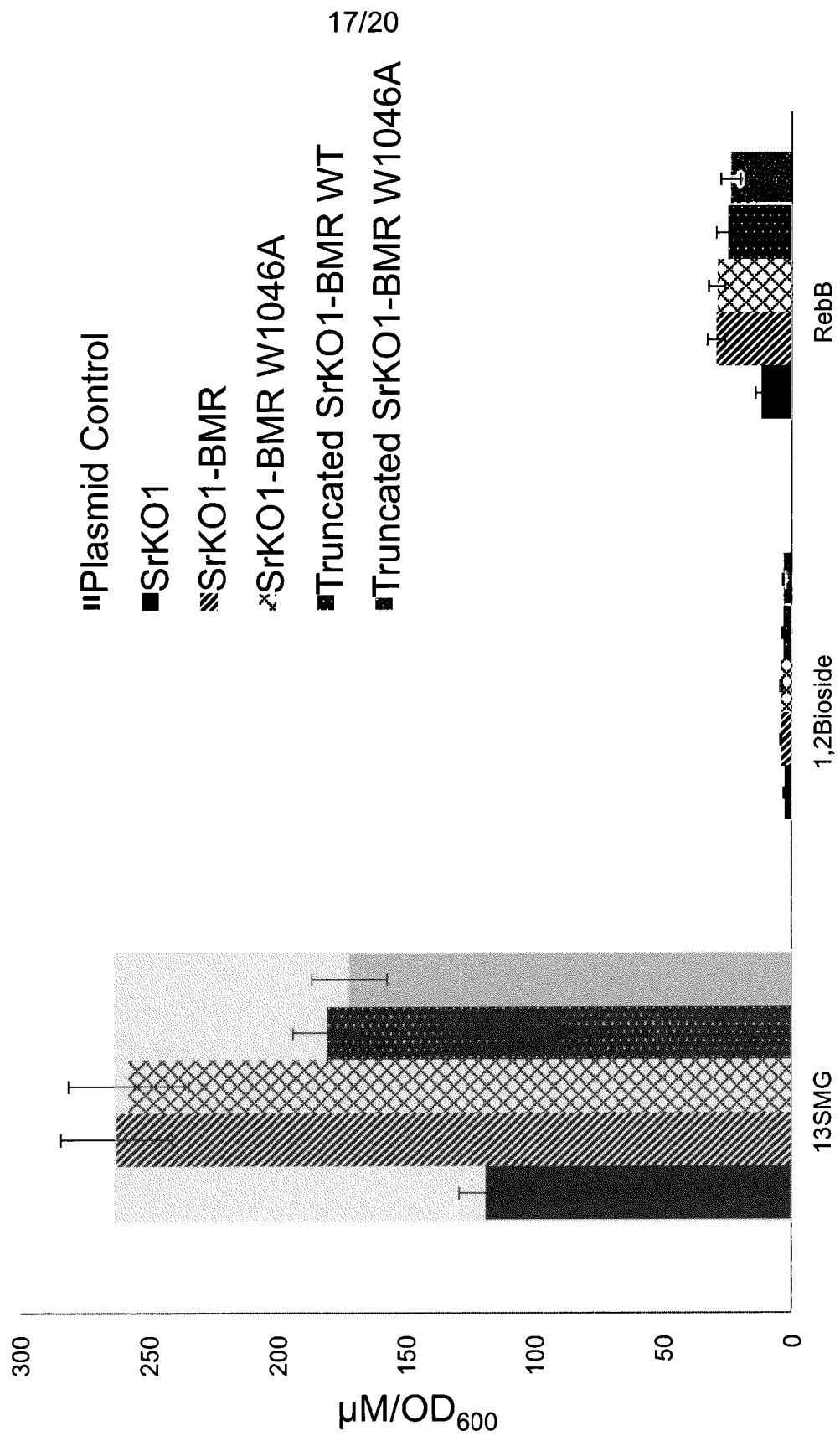


Figure 16B

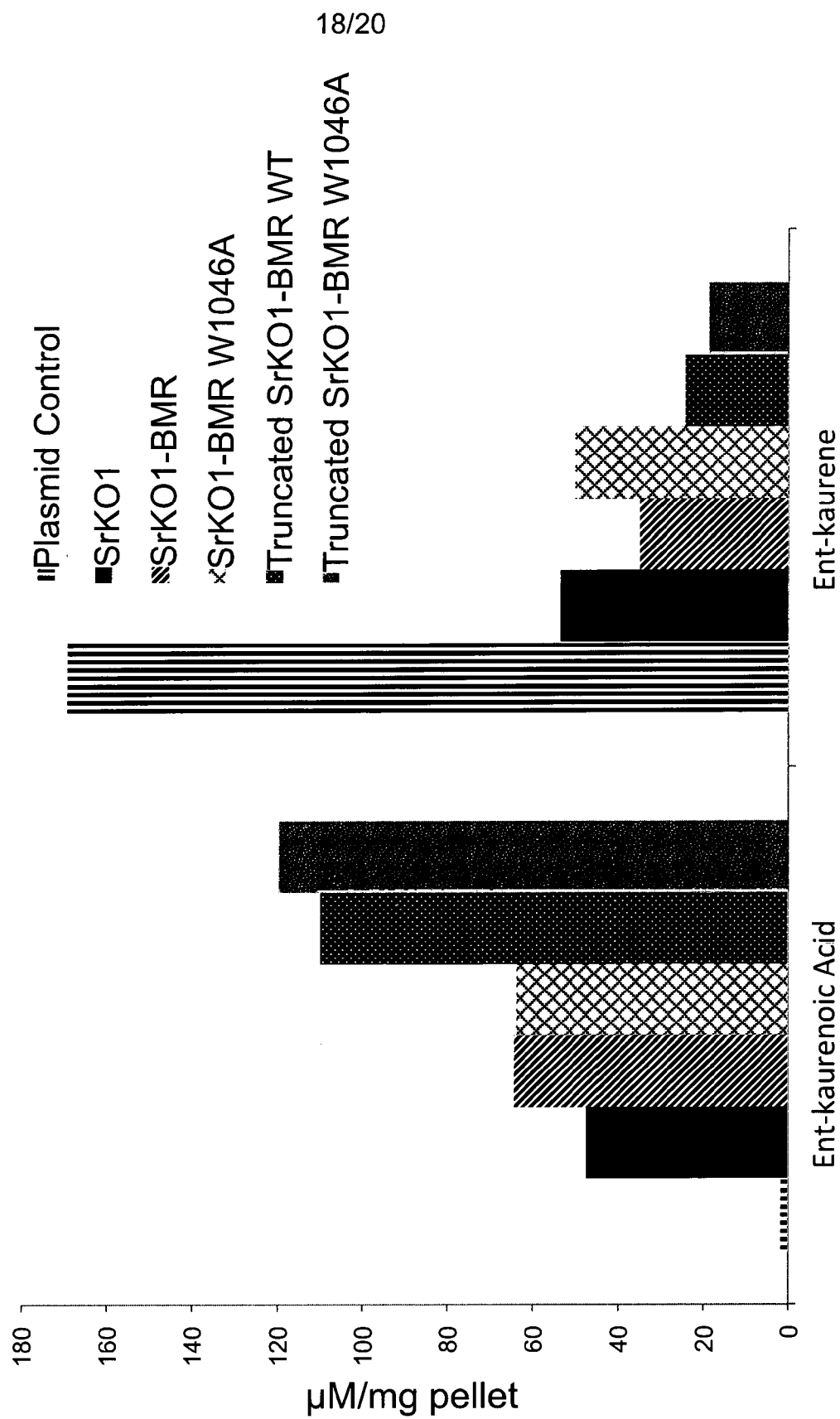
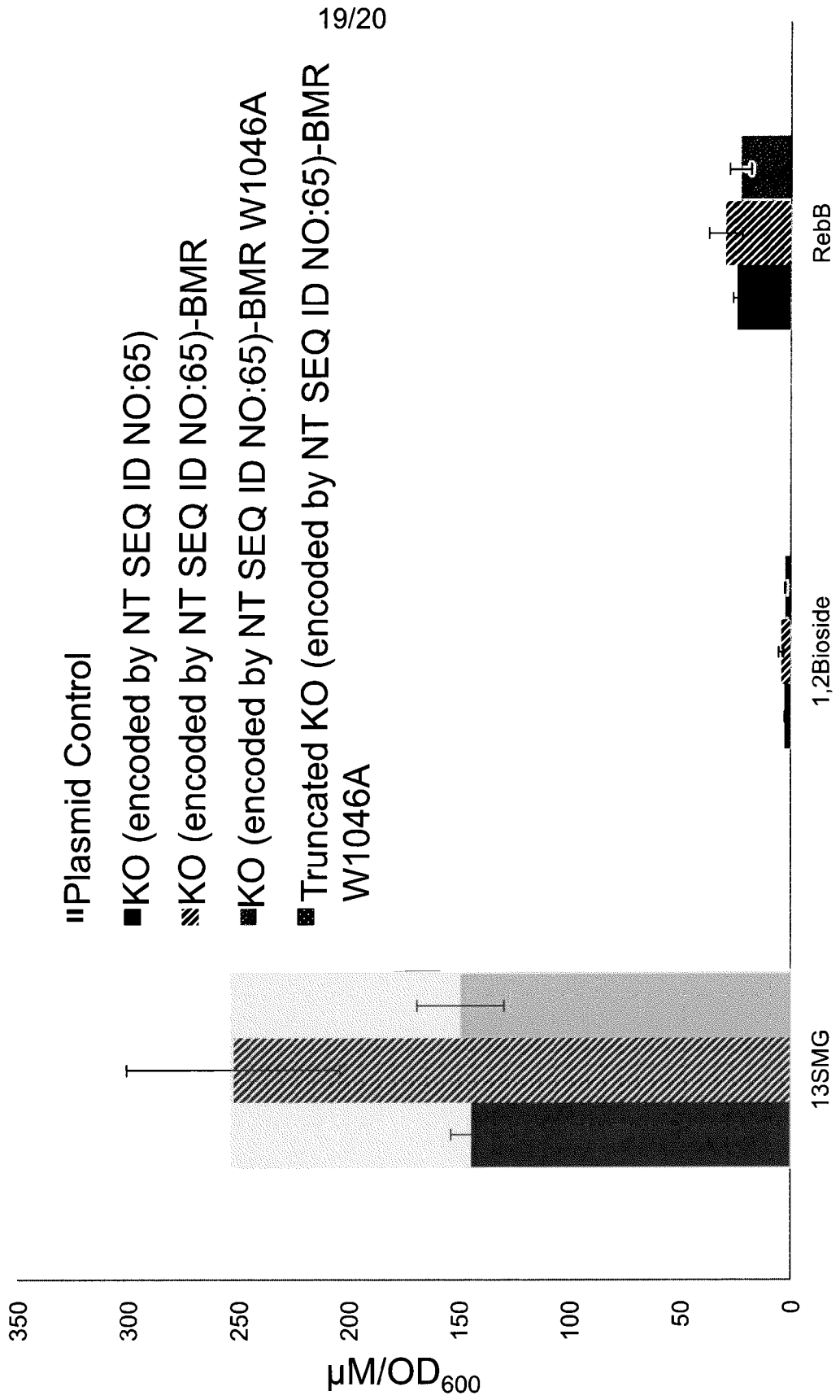


Figure 16C



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Figure 16D

