



US 20250178995A1

(19) **United States**(12) **Patent Application Publication****Lee et al.**(10) **Pub. No.: US 2025/0178995 A1**(43) **Pub. Date:****Jun. 5, 2025**(54) **COMPOSITIONS AND METHODS FOR THE PRODUCTION OF GLUCARIC ACID**(52) **U.S. Cl.**CPC *C07C 31/26* (2013.01); *B01J 31/003* (2013.01)(71) Applicant: **Solugen, Inc.**, Houston, TX (US)(72) Inventors: **Toni M. Lee**, Missouri City, TX (US); **David Weiner**, Houston, TX (US); **Konrad V. Miller**, Sugar Land (TX); **Gaurab Chakrabarti**, Houston, TX (US); **Sean Hunt**, Houston, TX (US)

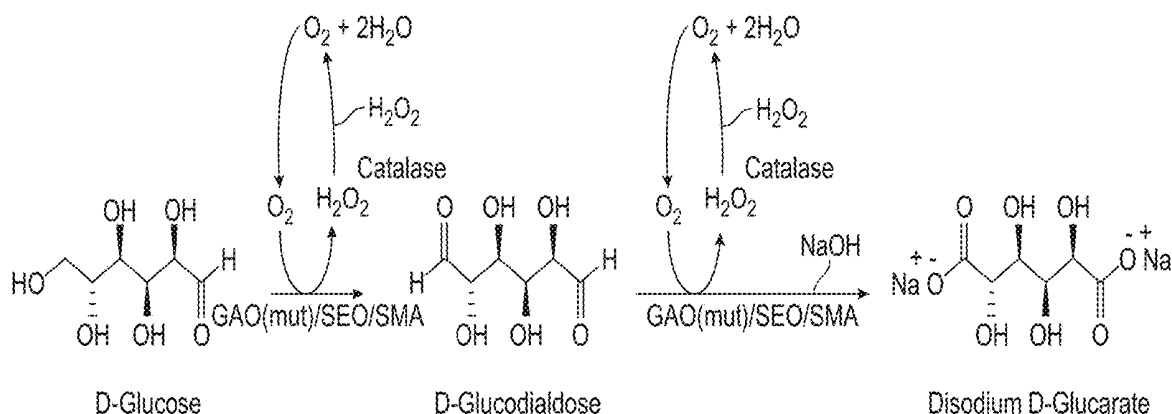
(57)

ABSTRACT

A method of preparing glucaric acid comprising contacting D-glucose and oxygen with a first catalyst composition comprising a first copper radical oxidase, a single electron oxidizer, and a small molecule activator under conditions suitable for formation of glucodialdose; and contacting glucodialdose and oxygen with a second catalyst composition comprising a second copper radical oxidase, a single electron oxidizer and a small molecule activator under conditions suitable for the formation of a product mixture comprising glucaric acid or salts thereof. A method comprising contacting a sugar with a catalyst composition comprising an oxidoreductase, a single electron oxidizer and a small molecule activator under conditions suitable for the formation of one or more oxidized sugar oxidation products comprising glucaric acid wherein the oxidoreductase comprises at least two copper radical oxidases, at least two mutated copper radical oxidases or combinations thereof.

(73) Assignee: **Solugen, Inc.**, Houston, TX (US)(21) Appl. No.: **18/965,379**(22) Filed: **Dec. 2, 2024****Related U.S. Application Data**

(60) Provisional application No. 63/604,831, filed on Nov. 30, 2023.

Publication Classification(51) **Int. Cl.***C07C 31/26* (2006.01)*B01J 31/00* (2006.01)**Specification includes a Sequence Listing.**

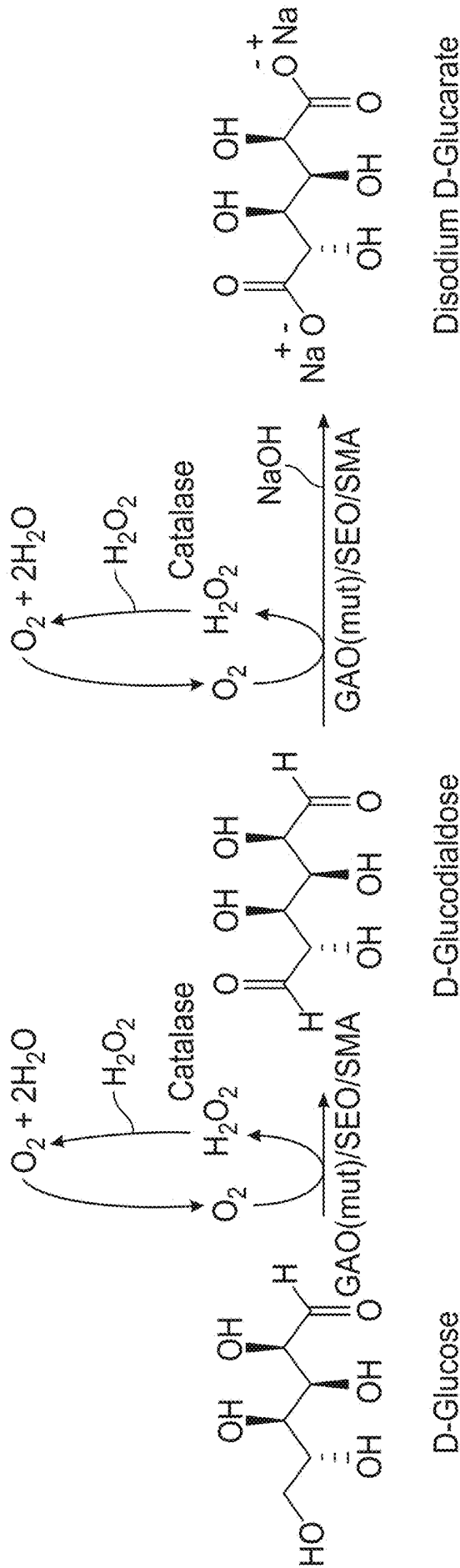


FIG. 1

COMPOSITIONS AND METHODS FOR THE PRODUCTION OF GLUCARIC ACID

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application claims priority to U.S. Provisional Patent Application No. 63/604,831 filed on Nov. 30, 2023 and entitled, "COMPOSITIONS AND METHODS FOR THE PRODUCTION OF GLUCARIC ACID," which is hereby incorporated herein by reference in its entirety for all purposes.

REFERENCE TO SEQUENCE LISTING

[0002] The instant application contains a Sequence Listing which has been submitted electronically in XML file format and is hereby incorporated by reference in its entirety. Said XML file, created on Nov. 25, 2024, is named "23ENZ008PCT_3416-18001 SEQUENCES11252024.xml" and is 94,208 bytes in size.

TECHNICAL FIELD

[0003] The present disclosure relates generally to the production of high value chemicals. More particularly, the present disclosure relates to the production of glucaric acid from reactants sourced from renewable resources.

BACKGROUND

[0004] Over the last 150 years, synthesis of inexpensive chemicals from fossilized forms of carbon (e.g. oil, coal, natural gas) has dramatically altered society through their broad applications, ranging from cosmetics to plastics. This petroleum-based carbon feedstock generates a small collection of platform chemicals from which highly efficient chemical conversions lead to the manufacture of a large variety of chemical products. However, the current approach to producing these carbon-based chemicals is inherently non-sustainable as feedstocks that required millions of years to form are being depleted. The creation of a truly sustainable chemical industry will only occur when the timescale of the feedstock formation matches the timescale of its utilization to make chemicals.

[0005] For example, the principal methodologies for producing glucaric acid commercially involve 1) nitric acid oxidation or 2) transition metal catalyzed oxidation. The drawbacks of these commercial methods include (i) the generation of a significant amount of hazardous nitrogen oxide (NO_x) gas and (ii) the processes are highly exothermic leading to controllability issues. The transition metal catalyzed (e.g., Pt or Pd) method for glucaric acid production uses glucaric acid as an intermediate in the production of bio-based adipic acid.

[0006] Other methods for the production of glucaric acid include a microbial method of producing high-purity glucaric acid using as catalysts myo-inositol-1-phosphate synthase from *S. cerevisiae*, mouse myo-inositol oxygenase, and *P. syringae* uronate dehydrogenase. Microbial methods, however, can suffer from product separation issues leading to high chemical costs, limiting usage as a commodity chemical. An ongoing need exists for novel glucaric acid production methods that lack some of the aforementioned challenges.

SUMMARY

[0007] A method of preparing glucaric acid comprising contacting D-glucose and oxygen with a first catalyst composition comprising a first copper radical oxidase, a single electron oxidizer, and a small molecule activator under conditions suitable for formation of glucodialdose; and contacting glucodialdose and oxygen with a second catalyst composition comprising a second copper radical oxidase, a single electron oxidizer and a small molecule activator under conditions suitable for the formation of a product mixture comprising glucaric acid or salts thereof.

[0008] A method comprising contacting a sugar with a catalyst composition comprising an oxidoreductase, a single electron oxidizer and a small molecule activator under conditions suitable for the formation of one or more oxidized sugar oxidation products comprising glucaric acid wherein the oxidoreductase comprises at least two copper radical oxidases, at least two mutated copper radical oxidases or combinations thereof.

[0009] Embodiments described herein comprise a combination of features and characteristics intended to address various shortcomings associated with certain prior devices, systems, and methods. The foregoing has outlined rather broadly the features and technical characteristics of the disclosed embodiments in order that the detailed description that follows may be better understood. The various characteristics and features described above, as well as others, will be readily apparent to those skilled in the art upon reading the following detailed description, and by referring to the accompanying drawings. It should be appreciated that the conception and the specific embodiments disclosed may be readily utilized as a basis for modifying or designing other structures for carrying out the same purposes as the disclosed embodiments. It should also be realized that such equivalent constructions do not depart from the spirit and scope of the principles disclosed herein.

BRIEF DESCRIPTION OF DRAWINGS

[0010] For a detailed description of various exemplary embodiments, reference will now be made to the accompanying drawings in which:

[0011] FIG. 1 schematically depicts a reaction of the present disclosure for generation of glucaric acid from D-glucose.

DETAILED DESCRIPTION

[0012] The following discussion is directed to various exemplary embodiments. However, one skilled in the art will understand that the examples disclosed herein have broad application, and that the discussion of any embodiment is meant only to be exemplary of that embodiment, and not intended to suggest that the scope of the disclosure, including the claims, is limited to that embodiment.

[0013] Certain terms are used throughout the following description and claims to refer to particular features or components. As one skilled in the art will appreciate, different persons may refer to the same feature or component by different names. This document does not intend to distinguish between components or features that differ in name but not function. The drawing FIGURES are not necessarily to scale. Certain features and components herein may be shown exaggerated in scale or in somewhat sche-

matic form and some details of conventional elements may not be shown in interest of clarity and conciseness.

[0014] Unless the context dictates the contrary, all ranges set forth herein should be interpreted as being inclusive of their endpoints, and open-ended ranges should be interpreted to include only commercially practical values. Similarly, all lists of values should be considered as inclusive of intermediate values unless the context indicates the contrary.

[0015] Where numerical ranges or limitations are expressly stated, such express ranges or limitations should be understood to include iterative ranges or limitations of like magnitude falling within the expressly stated ranges or limitations (e.g., from about 1 to about 10 includes, 2, 3, 4, etc.; greater than 0.10 includes 0.11, 0.12, 0.13, etc.).

[0016] In the following discussion and in the claims, the terms “including” and “comprising” are used in an open-ended fashion, and thus should be interpreted to mean “including, but not limited to . . .” As used herein, the phrases “consist(s) of” and “consisting of” are used to refer to exclusive components of a composition, meaning only those expressly recited components are included in the composition; whereas the phrases “consist(s) essentially of” and “consisting essentially of” are used to refer to the primary components of a composition, meaning that only small or trace amounts of components other than the expressly recited components (e.g., impurities, byproducts, etc.) may be included in the composition. For example, a composition consisting of X and Y refers to a composition that only includes X and Y, and thus, does not include any other components; and a composition consisting essentially of X and Y refers to a composition that primarily comprises X and Y, but may include small or trace amounts of components other than X and Y. In embodiments described herein, any such small or trace amounts of components other than those expressly recited following the phrase “consist(s) essentially of” or “consisting essentially of” preferably represent less than 5.0 wt % of the composition, more preferably less than 4.0 wt % of the composition, even more preferably less than 3.0 wt % of the composition, and still more preferably less than 1.0 wt % of the composition. Use of broader terms such as comprises, includes, having, etc. should be understood to provide support for narrower terms such as consisting of, consisting essentially of, comprised substantially of, etc. Use of the term “optionally” with respect to any element of a claim is intended to mean that the subject element is required, or alternatively, is not required. Both alternatives are intended to be within the scope of the claim.

[0017] Disclosed herein are chemoenzymatic methods of producing glucaric acid from a renewable resource such as a sugar. Advantageously, the disclosed chemoenzymatic method of glucaric acid production involves the use of benign conditions such as aqueous solvents and low temperatures. In one or more aspects, the disclosed compositions and methods are combined with processes for the formation of hexamethyldiamine which is subsequently utilized in the production of polyamides such as poly [imino (1,6-dioxohexamethylene) iminohexamethylene] (Nylon 66).

[0018] Advantageously, the presently disclosed methods utilize bio-based feedstocks, for example, feedstocks derived from biomass, as reactants resulting in the production of glucaric acid with high selectivity and minimized formation of toxic waste products.

[0019] In one or more aspects, a method of the present disclosure comprises contacting one or more sugars and subsequent reaction intermediates with a plurality of copper radical oxidases (CROs) under conditions suitable for the formation of glucaric acid.

[0020] In one or more aspects, the sugar comprises glucose. CROs are a class of non-flavoprotein alcohol oxidoreductases that employ molecular oxygen as a terminal electron acceptor to generate hydrogen peroxide. CROs have been labeled ‘green’ small-molecule oxidation catalysts as they lack dependence on an organic cofactor and utilize molecular oxygen as a cosubstrate.

[0021] Nonlimiting examples of CROs suitable for use in the present disclosure include glyoxal oxidases (EC 1.1.3.-, GLOX) and galactose 6-oxidases (EC 1.1.3.9, GAO). GLOXes typically function on aldehydes such as methylglyoxal to produce acids. One GLOX from *Phanerochaete chrysosporium* primarily accepts alpha-dicarbonyl and alpha-hydroxycarbonyls. In one or more aspects, the CRO is sourced from *Pycnoporus cinnabarinus*. *Pycnoporus cinnabarinus* was found to express three GLOXes, one of which has been found to function on methylglyoxal while the other two show high catalytic efficiency for glyoxylic acid.

[0022] In an aspect, the CRO is a GAO, alternatively a mutated GAO. GAOs function by oxidizing six-carbon (C6) or similar alcohols of galactose or other sugars to produce aldehydes. A particular example of a CRO is the GAO from *Fusarium graminearum*. Two additional CROs capable of oxidizing aliphatic alcohols are CgrAlcOx and CglAlcOx, which are derived from *Colletotrichum graminicola* and *C. gloeosporioides*, respectively. Another example is aryl-alcohol oxidase (CgrAAO) from *C. graminicola*.

[0023] In one or more aspects, a method of the present disclosure is depicted in FIG. 1. With reference to FIG. 1, a sugar (e.g., D-glucose) is reacted with a first CRO (designated CRO1) under conditions suitable for the formation of an intermediate, D-glucodialdose (GDA). In one or more aspects, conditions suitable for the formation of GDA include the presence of a small molecule activator (SMA) and a single electron oxidizer (SEO) such as a peroxidase; both of which facilitate the catalytic activity of the CRO1.

[0024] Without wishing to be limited by theory, the SEO performs a single electron oxidation on the SMA to generate the free radical form (SMA[•]). The SMA[•] can reverse or return a CRO (e.g., GAO) to the active state. Without wishing to be limited by theory, reversing and/or returning a CRO to the active state may comprise oxidation of one or more metals of the CRO at the active site of the enzyme to an active oxidation state.

[0025] In one or more aspects, an SMA suitable for use in this disclosure is a molecule that (i) is capable of stabilizing a free radical, (ii) can serve as a substrate for a SEO in a single electron oxidation reaction, and (iii) is capable of restoring the active state of a CRO as evidenced by detectable activity (e.g., formation of product or oxygen consumption).

[0026] Nonlimiting examples of SMAs suitable for use in the present disclosure include L-tryptophan, 2-mercaptobenzothiazole, L-histidine, methylchloroisothiazolinone, o-dianisidine, 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid (ABTS), 4-aminoantipyrine, L-tyrosine, (2,2,6,6-tetramethylpiperidin-1-yl)oxy, chloromethylisothiazolinone, 4-thiazolecarboxylic acid, Sunset yellow FCF, tartrazine, p-benzoquinone, dicoumarol, phthalimide, saccharin,

phthalic anhydride, erythrosine B, 2-aminobenzothiazole, thiabendazole, 2-hydroxybenzothiazole, phenothiazine, 6-aminobenzothiazole, indigo carmine, naphthalimide, 2-aminothiazole, thiazole, 2H-1,4-benzothiazin-3 (4H)-one, 2-oxindole, beta-lapachone, menaquinone, thiamine,

4-methyl-5-thiazoleethanol, Allura Red AC, menadione, p-cresol, Fast green FCF, Brilliant Blue FCF, methylisothiazolinone, caffeine, veratryl alcohol, fluorescein, and combinations thereof. The structures of several of these SMAs are depicted in Table 1.

TABLE 1

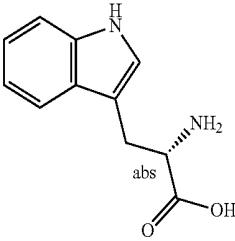
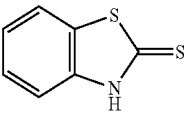
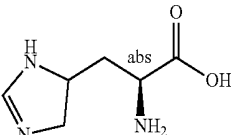
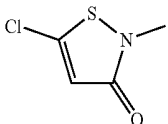
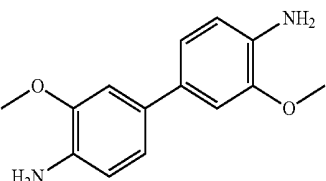
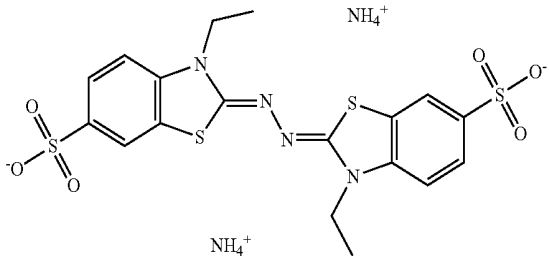
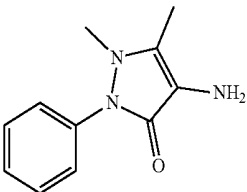
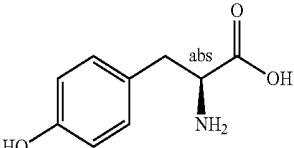
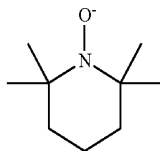
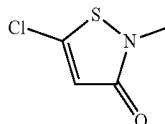
L-tryptophan	
2-mercapto-benzothiazole	
L-histidine	
Methylchloro-isothiazolinone	
o-dianisidine	
ABTS	
4-aminoantipyrine	
L-tyrosine	

TABLE 1-continued

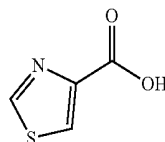
(2,2,6,6-Tetramethylpiperidin-1-yl)oxyl



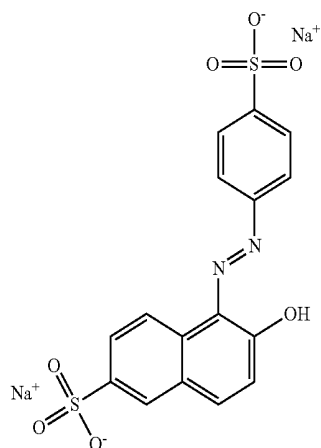
chloromethyl-isothiazolinone



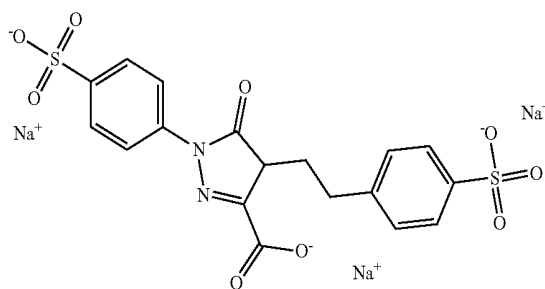
4-thiazolecarboxylic acid



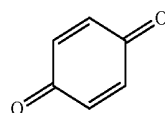
Sunset yellow FCF



Tartrazine



p-benzoquinone



dicoumarol

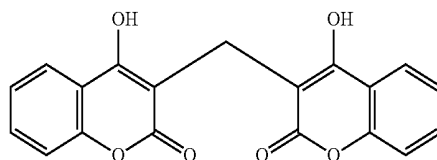


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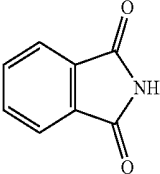
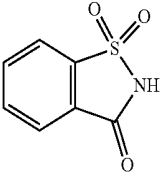
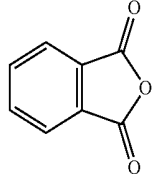
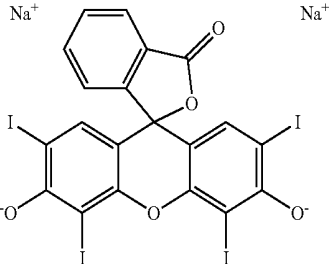
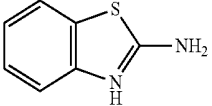
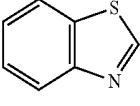
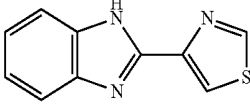
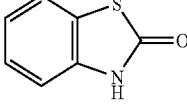
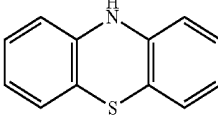
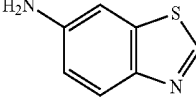
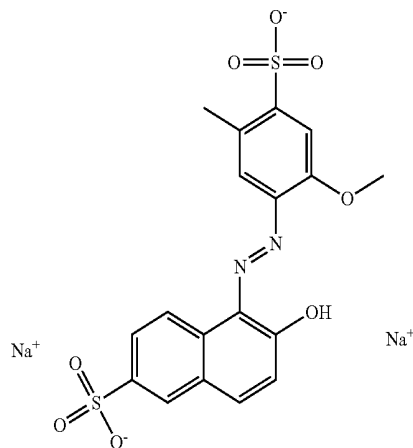
phthalimide	
saccharin	
phthalic anhydride	
Erythrosine B	
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benzothiazole	
thiabendazole	
2-Hydroxybenzothiazole	
Phenothiazine	
6-aminobenzothiazole	

TABLE 1-continued

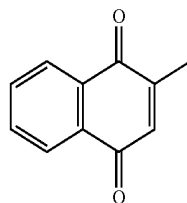
Indigo carmine	
naphthalimide	
2-aminothiazole	
thiazole	
2H-1,4-Benzothiazin-3(4H)-one	
2-oxindole	
beta-lapachone	
menaquinone	
thiamine	
4-Methyl-5-thiazoleethanol	

TABLE 1-continued

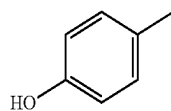
Allura Red AC



Menadione



p-cresol



Fast green FCF

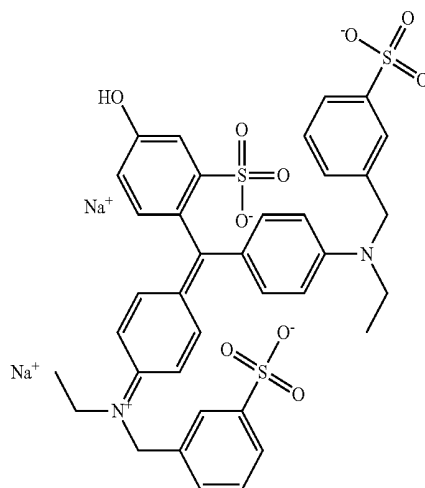
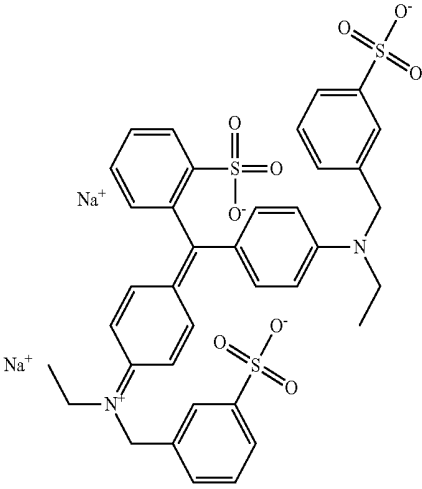
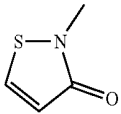
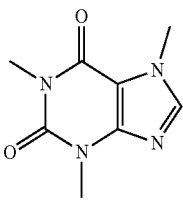
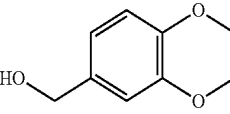
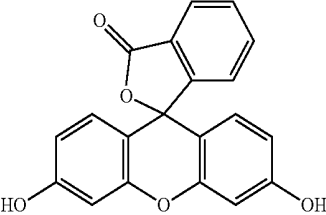


TABLE 1-continued

Brilliant Blue FCF	
Methylisothiazolinone	
caffeine	
veratryl alcohol	
fluorescein	

[0027] A SEO suitable for use in the present disclosure can be an enzyme or chemical compound (e.g., peroxidase or ferricyanide, catalase, and the like). The SEO can be any molecule that can elicit the radical state of the SMA by performing a single electron oxidation. In one or more aspects, the SEO is an enzyme. For example, the SEO is an enzyme that can use H_2O_2 as the oxidant which is advantageous as H_2O_2 is produced as a coproduct by a CRO.

[0028] In one or more aspects, the SEO is an enzyme such as a laccase, horseradish peroxidase, Dyp-type peroxidase, lactoperoxidase, chloroperoxidase, manganese peroxidase 1, ascorbate peroxidase, dye-decolorizing peroxidase, unspecific peroxygenase, dehaloperoxidase, catalase-peroxidase, lignin peroxidase, soybean seed coat peroxidase, isoforms thereof or combinations thereof.

[0029] In one or more aspects, a catalytic composition comprises a GAO, an SEO and a SMA. The catalytic

composition may comprise an GAO in an amount ranging from about 0.01 g/L to about 1 g/L, additionally or alternatively, from about 0.1 g/L to about 1 g/L, additionally or alternatively, from about 0.2 g/L to about 1 g/L, additionally or alternatively, from about 0.4 g/L to about 1 g/L, additionally or alternatively, from about 0.6 g/L to about 1 g/L, additionally or alternatively, from about 0.75 g/L to 1 g/L, additionally or alternatively, about 0.01 g/L, about 0.05 g/L, about 0.1 g/L, about 0.2 g/L, about 0.3 g/L, about 0.4 g/L, about 0.5 g/L, about 0.6 g/L, about 0.7 g/L, about 0.8 g/L, about 0.9 g/L or, additionally or alternatively, about 1 g/L. The catalytic composition may comprise an SEO in an amount ranging from about 1 mg/L to about 250 mg/L, additionally or alternatively, from about 5 mg/L to about 250 mg/L, additionally or alternatively, from about 10 mg/L to about 250 mg/L, additionally or alternatively, from about 25 mg/L to about 250 mg/L, additionally or alternatively, from

about 50 mg/L to find 250 mg/L, additionally or alternatively, from about 75 mg/L to 250 mg/L, additionally or alternatively, from about 100 mg/L to 250 mg/L, additionally or alternatively, from about 150 mg/L to 250 mg/L, additionally or alternatively, about 1 mg/L, about 5 mg/L, about 10 mg/L, about 15 mg/L, about 20 mg/L, about 25 mg/L, about 30 mg/L, about 35 mg/L, about 40 mg/L, about 45 mg/L, about 50 mg/L, about 55 mg/L, about 60 mg/L, about 65 mg/L, about 70 mg/L, about 75 mg/L, about 80 mg/L, about 85 mg/L, about 90 mg/L, about 95 mg/L, about 100 mg/L, about 105 mg/L, about 110 mg/L, about 115 mg/L, about 120 mg/L, about 125 mg/L, about 130 mg/L, about 135 mg/L, about 140 mg/L, about 145 mg/L, about 150 mg/L, about 155 mg/L, about 160 mg/L, about 165 mg/L, about 170 mg/L, about 175 mg/L, about 180 mg/L, about 185 mg/L, about 190 mg/L, about 195 mg/L, about 200 mg/L, about 205 mg/L, about 210 mg/L, about 215 mg/L, about 220 mg/L, about 225 mg/L, about 230 mg/L, about 235 mg/L, about 240 mg/L, about 245 mg/L or, additionally or alternatively, about 250 mg/L. The catalytic composition may comprise an SMA in an amount ranging from about 1 ppm to about 500 ppm, additionally or alternatively, from about 5 ppm to about 500 ppm, additionally or alternatively, from about 10 ppm to about 500 ppm, additionally or alternatively, from about 20 ppm to about 500 ppm, additionally or alternatively, or from about 40 ppm to about 400 ppm, additionally or alternatively, from about 50 ppm to about 350 ppm additionally or alternatively, from about 75 ppm to about 200 ppm additionally or alternatively, about 1 ppm, about 5 ppm, about 10 ppm, about 15 ppm, about 20 ppm, about 25 ppm, about 30 ppm, about 35 ppm, about 40 ppm, about 45 ppm, about 50 ppm, about 55 ppm, about 60 ppm, about 65 ppm, about 70 ppm, about 75 ppm, about 80 ppm, about 85 ppm, about 90 ppm, about 95 ppm, about 100 ppm, about 105 ppm, about 110 ppm, about 115 ppm, about 120 ppm, about 125 ppm, about 130 ppm, about 135 ppm, about 140 ppm, about 145 ppm, about 150 ppm, about 155 ppm, about 160 ppm, about 165 ppm, about 170 ppm, about 175 ppm, about 180 ppm, about 185 ppm, about 190 ppm, about 195 ppm, about 200 ppm, about 205 ppm, about 210 ppm, about 215 ppm, about 220 ppm, about 225 ppm, about 230 ppm, about 235 ppm, about 240 ppm, about 245 ppm, about 250 ppm, about 255 ppm, about 260 ppm, about 265 ppm, about 270 ppm, about 275 ppm, about 280 ppm, about 285 ppm, about 290 ppm, about 295 ppm, about 300 ppm, about 305 ppm, about 310 ppm, about 315 ppm, about 320 ppm, about 325 ppm, about 330 ppm, about 335 ppm, about 340 ppm, about 345 ppm, about 350 ppm, about 355 ppm, about 360 ppm, about 365 ppm, about 370 ppm, about 375 ppm, about 380 ppm, about 385 ppm, about 390 ppm, about 395 ppm, about 400 ppm, about 405 ppm, about 410 ppm, about 415 ppm, about 420 ppm, about 425 ppm, about 430 ppm, about 435 ppm, about 440 ppm, about 445 ppm, about 450 ppm, about 455 ppm, about 460 ppm, about 465 ppm, about 470 ppm, about 475 ppm, about 480 ppm, about 485 ppm, about 490 ppm, about 495 ppm, or, additionally or alternatively, about 500 ppm.

[0030] Conditions suitable for the formation of GDA may include one or more of the following reaction parameters: an amount of sugar (e.g., glucose) of from about 0.1 weight per volume percent (w/v %) to about 60 w/v %, alternatively from about 5 w/v % to about 50 w/v % or alternatively from about 10 w/v % to about 40 w/v %; a temperature ranging from about 1° C. to about 70° C., alternatively from about

5 to about 30° C. or alternatively from about 10° C. to about 25° C.; a suitable buffered media providing a pH ranging from about 5 to about 10, alternatively from about 5.5 to about 9 and a substrate amount ranging from about 6.5 to about 8.5, alternatively from about 6.5 to about 8.5 or alternatively from about 7 to about 8; an oxygen pressure of equal to or less than about 500 psi, alternatively from about 50 psi to about 250 psi or alternatively from about 70 psi to about 150 psi and a reaction time ranging from about 1 hour to about 24 hours or from about 2 hours to about 12 hours or from about 3 hours to about 6 hours.

[0031] GDA may be formed at yields that can range from about 50% to about 99%, additionally or alternatively, from about 50% to about 90%, additionally or alternatively, from about 50% to about 80%, additionally or alternatively, at least about 55%, at least about 58%, at least about 60%, at least about 62%, at least about 64%, at least about 66%, at least about 68%, at least about 70%, at least about 72%, at least about 74%, at least about 76%, at least about 78%, at least about 80%, at least about 82%, at least about 84%, at least about 86%, at least about 88%, at least about 90%, at least about 92%, at least about 94%, at least about 96%, at least about 98%, at least about or at least about 99%. The GDA intermediate formed may be used without further processing in the methods of the present disclosure. In alternative aspects, the GDA intermediate may be subjected to additional processing before (e.g., purification) prior to being utilized in other aspects of the presently disclosed methods. GDA also known as glucodialdose, L-gluco-hexodialdose, and D-gluco-hexodialdose, is a chiral intermediate with multiple industrial applications. The molecular structure corresponds with glucose, which inherently carries an aldehyde at the C1 position, oxidized to the dialdehyde at the C6 position.

[0032] In one or more aspects, a method of the present disclosure further comprises contacting the GDA formed with a second CRO (designated CRO2) under conditions suitable for the formation of glucaric acid. In one or more aspects, conditions suitable for the formation of glucaric acid include the presence of an SEO and/or SMA which may be the same as that used to facilitate the catalytic activity of CRO1. Alternatively, the SEO and/or SMA differ from those used to facilitate the catalytic activity of CRO1. In some aspects, CRO1 and CRO2 are the same enzyme, for example a GAO mutated to accept glucose as a substrate. In the alternative, CRO1 and CRO2 are different. For example, CRO 1 may be a GAO mutated to accept glucose as a substrate while CRO2 may be a GLOX mutated to oxidize GDA at the C1 and C6 positions to glucarate.

[0033] In one or more aspects, GDA is contacted with a catalytic composition comprising a GLOX, an SEO and a SMA. The catalytic composition may comprise a GLOX in an amount ranging from about 0.01 g/L to about 1 g/L, additionally or alternatively, from about 0.1 g/L to about 1 g/L, additionally or alternatively, from about 0.2 g/L to about 1 g/L, additionally or alternatively, from about 0.4 g/L to about 1 g/L, additionally or alternatively, from about 0.6 g/L to about 1 g/L, additionally or alternatively, from about 0.75 g/L to 1 g/L, additionally or alternatively, about 0.01 g/L, about 0.05 g/L, about 0.1 g/L, about 0.2 g/L, about 0.3 g/L, about 0.4 g/L, about 0.5 g/L, about 0.6 g/L, about 0.7 g/L, about 0.8 g/L, about 0.9 g/L or, additionally or alternatively, about 1 g/L. The catalytic composition may comprise an SEO in an amount ranging from about 1 mg/L to about 250

mg/L, additionally or alternatively, from about 5 mg/L to about 250 mg/L, additionally or alternatively, from about 10 mg/L to about 250 mg/L, additionally or alternatively, from about 25 mg/L to about 250 mg/L, additionally or alternatively, from about 50 mg/L to about 250 mg/L, additionally or alternatively, from about 75 mg/L to about 250 mg/L, additionally or alternatively, from about 100 mg/L to about 250 mg/L, additionally or alternatively, from about 150 mg/L to about 250 mg/L, additionally or alternatively, about 1 mg/L, about 5 mg/L, about 10 mg/L, about 15 mg/L, about 20 mg/L, about 25 mg/L, about 30 mg/L, about 35 mg/L, about 40 mg/L, about 45 mg/L, about 50 mg/L, about 55 mg/L, about 60 mg/L, about 65 mg/L, about 70 mg/L, about 75 mg/L, about 80 mg/L, about 85 mg/L, about 90 mg/L, about 95 mg/L, about 100 mg/L, about 105 mg/L, about 110 mg/L, about 115 mg/L, about 120 mg/L, about 125 mg/L, about 130 mg/L, about 135 mg/L, about 140 mg/L, about 145 mg/L, about 150 mg/L, about 155 mg/L, about 160 mg/L, about 165 mg/L, about 170 mg/L, about 175 mg/L, about 180 mg/L, about 185 mg/L, about 190 mg/L, about 195 mg/L, about 200 mg/L, about 205 mg/L, about 210 mg/L, about 215 mg/L, about 220 mg/L, about 225 mg/L, about 230 mg/L, about 235 mg/L, about 240 mg/L, about 245 mg/L or, additionally or alternatively, about 250 mg/L. The catalytic composition may comprise an SMA in an amount ranging from about 1 ppm to about 500 ppm, additionally or alternatively, from about 5 ppm to about 500 ppm, additionally or alternatively, from about 10 ppm to about 500 ppm, additionally or alternatively, from about 20 ppm to about 500 ppm, additionally or alternatively, or from about 40 ppm to about 400 ppm, additionally or alternatively, from about 50 ppm to about 350 ppm additionally or alternatively, from about 75 ppm to about 200 ppm additionally or alternatively, about 1 ppm, about 5 ppm, about 10 ppm, about 15 ppm, about 20 ppm, about 25 ppm, about 30 ppm, about 35 ppm, about 40 ppm, about 45 ppm, about 50 ppm, about 55 ppm, about 60 ppm, about 65 ppm, about 70 ppm, about 75 ppm, about 80 ppm, about 85 ppm, about 90 ppm, about 95 ppm, about 100 ppm, about 105 ppm, about 110 ppm, about 115 ppm, about 120 ppm, about 125 ppm, about 130 ppm, about 135 ppm, about 140 ppm, about 145 ppm, about 150 ppm, about 155 ppm, about 160 ppm, about 165 ppm, about 170 ppm, about 175 ppm, about 180 ppm, about 185 ppm, about 190 ppm, about 195 ppm, about 200 ppm, about 205 ppm, about 210 ppm, about 215 ppm, about 220 ppm, about 225 ppm, about 230 ppm, about 235 ppm, about 240 ppm, about 245 ppm, about 250 ppm, about 255 ppm, about 260 ppm, about 265 ppm, about 270 ppm, about 275 ppm, about 280 ppm, about 285 ppm, about 290 ppm, about 295 ppm, about 300 ppm, about 305 ppm, about 310 ppm, about 315 ppm, about 320 ppm, about 325 ppm, about 330 ppm, about 335 ppm, about 340 ppm, about 345 ppm, about 350 ppm, about 355 ppm, about 360 ppm, about 365 ppm, about 370 ppm, about 375 ppm, about 380 ppm, about 385 ppm, about 390 ppm, about 395 ppm, about 400 ppm, about 405 ppm, about 410 ppm, about 415 ppm, about 420 ppm, about 425 ppm, about 430 ppm, about 435 ppm, about 440 ppm, about 445 ppm, about 450 ppm, about 455 ppm, about 460 ppm, about 465 ppm, about 470 ppm, about 475 ppm, about 480 ppm, about 485 ppm, about 490 ppm, about 495 ppm, or, additionally or alternatively, about 500 ppm.

[0034] In one or more aspects, the reaction disclosed herein is carried out under mild conditions. Conditions suitable for the formation of glucaric acid may include one

or more of the following reaction parameters: an amount of GDA of from about 0.1 weight per volume percent (w/v %) to about 60 w/v %, alternatively from about 5 w/v % to about 50 w/v % or alternatively from about 10 w/v % to about 40 w/v %; a temperature ranging from about 5° C. to about 100° C., or from about 10° C. to about 75° C., or from about 15° C. to about 70° C., or from about 25° C. to about 50° C.; a suitable buffered media providing a pH ranging from about 4 to about 9, or from about 5.5 to about 9, or from about 6 to about 8; an oxygen pressure of equal to or less than about 500 psi, alternatively from about 50 psi to about 250 psi or alternatively from about 70 psi to about 150 psi and a reaction time ranging from about 1 hour to about 24 hours or from about 2 hours to about 12 hours or from about 3 hours to about 6 hours.

[0035] Glucaric acid may be formed at yields that can range from about 50% to about 99%, additionally or alternatively, from about 50% to about 90%, additionally or alternatively, from about 50% to about 80%, additionally or alternatively, at least about 55%, at least about 58%, at least about 60%, at least about 62%, at least about 64%, at least about 66%, at least about 68%, at least about 70%, at least about 72%, at least about 74%, at least about 76%, at least about 78%, at least about 80%, at least about 82%, at least about 84%, at least about 86%, at least about 88%, at least about 90%, at least about 92%, at least about 94%, at least about 96%, at least about 98%, at least about or at least about 99%.

[0036] In some aspects, the reaction for formation of glucaric acid further includes a caustic agent such as sodium hydroxide or another base which functions to both control pH and as a counterion for produced glucaric acid or glucarate salts.

[0037] In some aspects, the catalyst compositions disclosed herein further comprise a catalase in order to decompose the hydrogen peroxide produced during the reaction of the CRO. A catalase is used to break down two molecules of hydrogen peroxide (H₂O₂) into one molecule of oxygen (O₂) and two molecules of water (H₂O). In some aspects, oxygen generated by catalase may be captured and used in the oxidation of additional substrate (e.g., glucose or GDA).

[0038] In one or more aspects, the methods of the present disclosure produce high purity glucaric acid or salts thereof, for example, glucaric acid may have a purity of greater than about 70%, or greater than about 90% or greater than about 95%, or from about 70% to about 99%, or from about 72% to about 99%, or from about 75% to about 99%, or from about 78% to about 99%, or from about 80% to about 99%, or from about 82% to about 99%, or from about 84% to about 99%, or from about 86% to about 99%, or from about 88% to about 99%, or from about 90% to about 99%, or from about 92% to about 99%, or from about 94% to about 99%, or from about 96% to about 99%. In an aspect, the product comprises disodium glucarate, monosodium glucarate, potassium glucarate or combinations thereof in crystalline form. In an alternative aspect, the product of the present disclosure comprises a mixture of free acid in combination with salts of glucaric acid (e.g., potassium and/or sodium salt).

[0039] In one or more aspects, CRO1 which catalyzes the formation of glucodialdohexose can be any CRO or mutant thereof capable of oxidizing the C6 of glucose. In one or more aspects, CRO2 which catalyzes the formation of glucaric acid can be any CRO capable of oxidizing the C1

and C6 aldehydes of glucodialdose. In one or more aspects, a GLOX engineered to oxidize the C1 and C6 of GDA could catalyze this reaction.

[0040] In an aspect, an enzyme of the type disclosed herein is a wild type enzyme, a functional fragment thereof, or a functional variant thereof. "Fragment" as used herein is meant to include any amino acid sequence shorter than the full-length enzyme, but where the fragment maintains a catalytic activity sufficient to meet some user or process goal. Fragments may include a single contiguous sequence identical to a portion of the biocatalyst sequence. Alternatively, the fragment may have or include several different shorter segments where each segment is identical in amino acid sequence to a different portion of the amino acid sequence of the enzyme but linked via amino acids differing in sequence from the enzyme. Herein, a "functional variant" of the enzyme refers to a polypeptide which has at one or more positions of an amino acid insertion, deletion, or substitution, either conservative or non-conservative, and wherein each of these types of changes may occur alone, or in combination with one or more of the others, and/or one or more times in a given sequence but retains catalytic activity.

[0041] In the alternative or in combination with the aforementioned mutations, the enzyme may be mutated to improve the catalytic activity. Mutations may be carried out to enhance the protein or a homolog activity, increase the protein stability in the presence of substrates and products (e.g., hydrogen peroxide) and increase protein yield.

[0042] Herein, reference has been made to "sources" of enzyme. It is to be understood this refers to the biomolecule as expressed by the named organism. It is contemplated the enzyme may be obtained from the organism or a version of said enzyme (wildtype or recombinant) and provided as a suitable construct to an appropriate expression system.

[0043] In an aspect, any enzyme of the type disclosed herein may be cloned into an appropriate expression vector and used to transform cells of an expression system such as *E. coli*, *Saccharomyces* sp., *Pichia* sp., *Aspergillus* sp., *Trichoderma* sp., or *Myceliophthora* sp. A "vector" is a replicon, such as plasmid, phage, viral construct or cosmid, to which another DNA segment may be attached. Vectors are used to transduce and express a DNA segment in cells. As used herein, the terms "vector" and "construct" may include replicons such as plasmids, phage, viral constructs, cosmids, Bacterial Artificial Chromosomes (BACs), Yeast Artificial Chromosomes (YACs) Human Artificial Chromosomes (HACs) and the like into which one or more gene expression cassettes may be or are ligated. Herein, a cell has been "transformed" by an exogenous or heterologous nucleic acid or vector when such nucleic acid has been introduced inside the cell, for example, as a complex with transfection reagents or packaged in viral particles. The transforming DNA may or may not be integrated (covalently linked) into the genome of the cell.

[0044] In an aspect, the gene of an enzyme disclosed herein is provided as a recombinant sequence in a vector where the sequence is operatively linked to one or more control or regulatory sequences. "Operatively linked" expression control sequences refer to a linkage in which the expression control sequence is contiguous with the gene of interest to control the gene of interest, as well as expression control sequences that act in trans or at a distance to control the gene of interest.

[0045] The term "expression control sequence" or "regulatory sequences" are used interchangeably and are used herein refer to polynucleotide sequences which affect the expression of coding sequences to which they are operatively linked. Expression control sequences are sequences that control the transcription, post-transcriptional events, and translation of nucleic acid sequences. Expression control sequences include appropriate transcription initiation, termination, promoter, and enhancer sequences; efficient RNA processing signals such as splicing and polyadenylation signals; sequences that stabilize cytoplasmic mRNA; sequences that enhance translation efficiency (e.g., ribosome binding sites); sequences that enhance protein stability; and when desired, sequences that enhance protein secretion. The nature of such control sequences differs depending upon the host organism; in prokaryotes, such control sequences generally include promoter, ribosomal binding site, and transcription termination sequence. The term "control sequences" is intended to include, at a minimum, all components whose presence is essential for expression, and can also include additional components whose presence is advantageous, for example, leader sequences and fusion partner sequences.

[0046] The term "recombinant host cell" ("expression host cell", "expression host system", "expression system" or simply "host cell"), as used herein, is intended to refer to a cell into which a recombinant vector has been introduced. It should be understood that such terms are intended to refer not only to the particular subject cell but to the progeny of such a cell. Because certain modifications may occur in succeeding generations due to either mutation or environmental influences, such progeny may not, in fact, be identical to the parent cell, but are still included within the scope of the term "host cell" as used herein. A recombinant host cell may be an isolated cell or cell line grown in culture or may be a cell which resides in a living tissue or organism.

[0047] In one or more aspects, a CRO suitable for use in the present disclosure is a CRO comprising any of SEQ ID Nos. 1 through 39 or alternatively from about 85% to about 100% sequence identity with any of SEQ ID Nos. 1 through 39. In one or more aspects, a catalase suitable for use in the present disclosure is a catalase defined by of any of SEQ ID Nos. 40 through 44. In one or more aspects, a peroxidase suitable for use in the present disclosure is a peroxidase defined by of any of SEQ ID Nos. 45 through 61. In one or more aspects, a CRO suitable for use in the present disclosure is a CRO defined by of any of SEQ ID Nos. 62 through or alternatively from about 85% to about 100% sequence identity with any of SEQ ID Nos. 62 through 65.

[0048] In one or more aspects, a glucaric acid product of the type disclosed herein may be used in the production of into adipic acid and subsequently adipic adipodinitrile, which can then be hydrogenated to produce hexamethyldiamine.

ADDITIONAL DISCLOSURE

[0049] A first aspect which is a method of preparing glucaric acid comprising contacting D-glucose and oxygen with a first catalyst composition comprising a first copper radical oxidase, a single electron oxidizer, and a small molecule activator under conditions suitable for formation of glucodialdose; and contacting glucodialdose and oxygen with a second catalyst composition comprising a second copper radical oxidase, a single electron oxidizer and a small

molecule activator under conditions suitable for the formation of a product mixture comprising glucaric acid or salts thereof.

[0050] A second aspect which is the method of the first aspect wherein the first copper radical oxidase comprises a galactose oxidase, a galactose oxidase mutant or combinations thereof.

[0051] A third aspect which is the method of any of the first through second aspects wherein the first copper radical oxidase comprises any of SEQ ID Nos. 1 through 39.

[0052] A fourth aspect which is the method of any of the first through third aspects wherein the first copper radical oxidase has from about 85% to about 100% sequence identity with any of SEQ ID Nos. 1 through 39.

[0053] A fifth aspect which is the method of any of the first through fourth aspects wherein the second copper radical oxidase comprises a glyoxal oxidase, a glyoxal oxidase mutant or combinations thereof.

[0054] A sixth aspect which is the method of any of the first through fifth aspects wherein the second copper radical oxidase comprises any of SEQ ID Nos. 62 through 65.

[0055] A seventh aspect which is the method of any of the first through sixth aspects wherein the second copper radical oxidase has from about 85% to about 100% sequence identity with any of SEQ ID Nos. 62 through 65.

[0056] An eighth aspect which is the method of any of the first through seventh aspects wherein the small molecule activator comprises L-tryptophan, 2-mercaptobenzothiazole, L-histidine, methylchloroisoethiazolinone, o-dianisidine, 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid (ABTS)), 4-aminoantipyrine, L-tyrosine, (2,2,6,6-tetramethylpiperidin-1-yl)oxyl, chloromethylisothiazolinone, 4-thiazolecarboxylic acid, Sunset yellow FCF, tartrazine, p-benzoquinone, dicoumarol, phthalimide, saccharin, phthalic anhydride, erythrosine B, 2-aminobenzothiazole, thiabendazole, 2-hydroxybenzothiazole, phenothiazine, 6-aminobenzothiazole, indigo carmine, naphthalimide, 2-aminothiazole, thiazole, 2H-1,4-benzothiazin-3 (4H)-one, 2-oxindole, betalaphachone, menaquinone, thiamine, 4-methyl-5-thiazoleethanol, Allura Red AC, menadione, p-cresol, Fast green FCF, Brilliant Blue FCF, methylisothiazolinone, caffeine, veratryl alcohol, fluorescein, or combinations thereof.

[0057] A ninth aspect which is the method of any of the first through eighth aspects wherein the small molecule activator is present in an amount ranging from about 1 ppm to about 500 ppm.

[0058] A tenth aspect which is the method of any of the first through ninth aspects wherein the single electron oxidizer comprises laccase, horseradish peroxidase, dyp-type peroxidase, lactoperoxidase, chloroperoxidase, manganese peroxidase 1, ascorbate peroxidase, dye-decolorizing peroxidase, unspecific peroxygenase, dehaloperoxidase, catalase-peroxidase, lignin peroxidase, soybean seed coat peroxidase, isoforms thereof or combinations thereof.

[0059] An eleventh aspect which is the method of any of the first through tenth aspects wherein conditions suitable for formation of glucodialdose, conditions suitable for formation of a product mixture comprising glucaric acid or both comprise an oxygen pressure of equal to or less than about 500 psi.

[0060] A twelfth aspect which is the method of any of the first through eleventh aspects further comprising a catalase.

[0061] A thirteenth aspect which is the method of any of the first through twelfth aspects wherein the catalase is defined by any SEQ ID No. 40 through SEQ ID No. 44.

[0062] A fourteenth aspect which is the method of any of the first through thirteenth aspects further comprising introducing a caustic to the second catalyst composition, the product mixture or both.

[0063] A fifteenth aspect which is the method of any of the first through fourteenth aspects wherein the glucodialdose, glucaric acid are formed at yields ranging from about 50% to about 99%.

[0064] A sixteenth aspect which is the method of any of the first through fifteenth aspects wherein the glucaric acid or salts thereof have a purity of greater than about 70%.

[0065] A seventeenth aspect which is the method of any of the first through sixteenth aspects wherein the product mixture comprising glucaric acid further comprises one or more sugar oxidation products.

[0066] An eighteenth aspect which is the method of the seventeenth aspect wherein the one or more sugar oxidation products comprise aldonic acid, uronic acid, aldaric acid, a gluconic acid oxidation product, a gluconate, gluconic acid, glucuronic acid, glucose oxidation products, galactonic acid, galactaric acid, glutamic acid, a lactone of gluconic acid, a lactone of glucaric acid, a lactone of galactaric acid, a lactone of galactonic acid, glucodialdose, 2-ketoglucose, disaccharides, oxidized disaccharides, n-keto-acids, C2 to C6 diacids, salts thereof or combinations thereof.

[0067] A nineteenth aspect which is a method comprising contacting a sugar with a catalyst composition comprising an oxidoreductase, a single electron oxidizer and a small molecule activator under conditions suitable for the formation of one or more oxidized sugar oxidation products comprising glucaric acid wherein the oxidoreductase comprises at least two copper radical oxidases, at least two mutated copper radical oxidases or combinations thereof.

[0068] A twentieth aspect which is the method of the nineteenth aspect wherein the at least two copper radical oxidases comprises any of SEQ ID Nos. 1 through 39, any of SEQ ID Nos. 62 through 65, have 85% to about 100% sequence identity with any of SEQ ID Nos. 1 through 39 or have 85% to about 100% sequence identity with any of SEQ ID Nos. 62 through 65.

EXAMPLES

[0069] The aspects having been generally described, the following example is given as particular aspects of the disclosure and to demonstrate the practice and advantages thereof. It is understood that the example is given by way of illustration and is not intended to limit the specification or the claims in any manner.

Example 1

Generation of GAO Mutant for Producing Glucodialdose from Glucose

[0070] A GAO mutant capable of converting glucose to GDA was engineered. Following directed evolution and rational enzyme engineering, the improved GAO mutant exhibits a specific activity of 35 U mg⁻¹ on glucose.

Directed Evolution

[0071] Directed evolution of thirty sites within 10 Å of the catalytic copper was performed on a parent sequence con-

taining the following added mutations: 1) R330, Q406T, W290F to introduce less than 1 U mg⁻¹ activity on glucose to GAO, 2) C383S to lower the KM of the enzyme on galactose, and 3) Y405F and Q406E to enhance activity on a D-N-acetyl glucosamine substrate. Other mutations described in Table I were found to have neutral or deleterious effects on glucodialdose-generating activity. The new combination sequence was designated GAO-Mut1. The sequence of GAO-Mut1 contains a “MGHHHHHHSSGHIEGRHM” N-terminal his-tag and linker for expression and purification in *E. coli* and is SEQ ID NO:66.

[0072] While preferred embodiments have been shown and described, modifications thereof can be made by one skilled in the art without departing from the scope or teachings herein. The embodiments described herein are exemplary only and are not limiting. Many variations and modifications of the systems, apparatus, and processes described herein are possible and are within the scope of the disclosure. For example, the relative dimensions of various parts, the materials from which the various parts are made,

and other parameters can be varied. Accordingly, the scope of protection is not limited to the embodiments described herein, but is only limited by the claims that follow, the scope of which shall include all equivalents of the subject matter of the claims. Unless expressly stated otherwise, the steps in a method claim may be performed in any order. The recitation of identifiers such as (a), (b), (c) or (1), (2), (3) before steps in a method claim are not intended to and do not specify a particular order to the steps, but rather are used to simplify subsequent reference to such steps.

[0073] Each and every claim is incorporated into the specification as an aspect of the present disclosure. Thus, the claims are a further description and are an addition to the aspects of the present invention. The discussion of a reference herein is not an admission that it is prior art to the presently disclosed subject matter, especially any reference that may have a publication date after the priority date of this application. The disclosures of all patents, patent applications, and publications cited herein are hereby incorporated by reference, to the extent that they provide exemplary, procedural or other details supplementary to those set forth herein.

SEQUENCE LISTING

Sequence total quantity: 66

SEQ ID NO: 1 moltype = AA length = 657
FEATURE Location/Qualifiers
source 1..657
 mol_type = protein
 organism = synthetic construct

SEQUENCE: 1

MGHHHHHHSS GHIEGRHMAS APIGSAIPRN NWAVTCDSAQ SGNECNKAIK GNKDTFWHTF 60
YGANGDPKPP HTYTIDMKTT QNVNGLSVLP RQDGNQNGWI GRHEVYLLSSD GTNNGSPVAS 120
GSWFADSTTK YSNFETRPAP YVRLVAITEA NGQPWTSIAE INVVFQASSYT APQPGLGRWG 180
PTIDLPIVPA AAAIEPTSGR VLMWSSYRND AFEQSPGGIT LTSSWDPSTG IVSDRTVTVT 240
KHDMPFCPGIS MDGNGQIVVT GGNDKAKTSL YDSSSDSWIP GPDQMVARGY QSSATMSDGR 300
VFTIGGSFSG GVFEKNGEVY SPSSKTWTSL PNAKVNPMML ADKQGLYKSD NHAWLFGWKK 360
GSVVFQAGPST AMNWWYTSGS GDVKSAGKRQ SNRGVAPDAM SGNAMVYDAV KGKILTFGGS 420
PDFEDSDATT NAHIITLGEF GTSPTNVFAS NGLYFARTFH TSVVLPDGSF FITGGQRRGI 480
PFEDSTPVFT PEIYVPEQDT FYKQNPNSIV RAYHSISLLL PDGRVFNNGG GLCGDCTTNH 540
FDAQIFTPNY LYDSNGNLAT RPKITRSTQ SVKVGGRITI STDSSISKAS LIRYGTATHT 600
VNTDQRRIPL TLTNNGNSY SFQVPSDSGV ALPGYWMLFV MNSAGVPSVA STIRVTQ 657

SEQ ID NO: 2 moltype = AA length = 639
FEATURE Location/Qualifiers
source 1..639
 mol_type = protein
 organism = synthetic construct

SEQUENCE: 2

ASAPIGSAIS RNNWAVTCDS AQSNGNECKA IDGNKDTFWH TFGANGDPK PPHTYTIDMK 60
TTQNVNGLSM LPRQDGNQNG WIGRHEVYLS SDGTNNGSPV ASGSWFADST TKYSNFETRP 120
ARYVRLVAIT EANGQPWTSI AEINVQASS YTAPQPGLGR WGPTIDLPIV PAAAAIEPTS 180
GRVLMWSSYR NDAPGGSPGG ITLTSSWDPS TGIVSDRTVT VTKHDMFCPG ISMDGNGQIV 240
VTGGNDAKKT SLYDSSSDSW IPGPDQVAR GYQSSATMSD GRVFTIGGSW SGGVFEKNGE 300
VYSPSSKTWT SLPNAKVNPM LTADKQGLYR SDNHAWLFGW KKGSVFQAGP STAMNWWYTS 360
GSGDVKSAGK RQSNRGVAPD AMCGNAVMYD AVKGKILTFG GSPDYQSDA TNAHIITLG 420
EPGTSPTNVF ASNGLYFART FHTSVVLPDG STFITGGQRR GIPFEDSTPV FTPEIYVPEQ 480
DTFYKQNPNS IVRVYHSISL LLPDGRVFNQ GGLCGDCTT NHFDAQIFTP NYLYNSGNL 540
ATRPKITRST TQSVKVGGRIT ITSTDSSISK ASLIRYGTAT HTVNTDQRRIP LTLTNNNGN 600
SYSFQVPSDS GVALPGYWML FVMNSAGVPS VASTIRVTQ 639

SEQ ID NO: 3 moltype = AA length = 680
FEATURE Location/Qualifiers
source 1..680
 mol_type = protein
 organism = synthetic construct

SEQUENCE: 3

MKHFLSLALC FSSINAVAVT VPHKSGGTGS PEGSLQFLSL RASAPIGSAI SRNNWAVTCD 60
SAQSGNECKN AIDGNKDTFW HTFYGANGDP KPPHTYTIDM KTTQNVNGLS MLPRQDGNQN 120
GWIGRHEVYL SSDGTNNGSP VASGSWFADS TTKYSNFETR PARYVRLVAV TEANGQPWTS 180
IAEINVQAS SYTAPQPGLG RWGPTIDLPI VPAAAAIEPT SGRVLMWSSY RNDAPGGSPG 240

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GITLTSSWDP	STGIVSDRTV	TVTCKDMFCP	GISMDGNGQI	VVTGGNDACK	TSLYDSSSDS	300
WIPGPDQVA	RGYQSSATMS	DGRVFTIGGS	WSGGVFEKNG	EVYSPSSKTW	TSLPNAKVNP	360
MLTADKQGLY	RSDNHAWLFG	WKKGSVFQAG	PSTAMNYYT	SGSGDVKSAG	KRQSNRGVAP	420
DAMCGNAVMY	DAVKGKILTF	GGSPDYQSD	ATTNAHIITL	GEPGTSPTV	FASNGLYFAR	480
TFHTSVVLPD	GSTFITGGQR	RGIPFEDSTP	VFTPEIYVPE	QDTFYKQPNP	SIVRVYHSIS	540
LLLPDGRVFN	GGGGLCGDCT	TNHFDAQIFT	PNYLYNSNGN	LATRPKI TRT	STQSVKVGGR	600
ITISTDSSIT	KASLIRYGTGTA	THTVNTDQRR	IPLTLTNNGG	NSYSFQVPSD	SGVALPGYWM	660
LFVMSAGVP	SVASTIRVTQ					680

SEQ ID NO: 4 moltype = AA length = 680
 FEATURE Location/Qualifiers
 source 1..680
 mol_type = protein
 organism = Fusarium graminearum

SEQUENCE: 4

MKHFLSLALC	FSSINAVAVT	VPHKSGGTGS	PEGSLQFLSL	RASAPIGSAI	SRNNWAVTCD	60
SAQSGNECNK	AIDGNKDTFW	HTFYGANQDP	KPPHTYTIDM	KTTONVNGLS	MLPRQDGNQN	120
GWIGRHEVYL	SSDGTNWSGP	VASGSWFADS	TTKYSNFETR	PARYVRLVAV	TEANGQPWTS	180
IAEINVFQAS	SYTAPQPLGL	RWGPTIDLPI	VPAAAAIEPT	SGRVLMWSSY	RNDAFGGSPG	240
GITLTSSWDP	STGIVSDRTV	TVTCKDMFCP	GISMDGNGQI	VVTGGNDACK	TSLYDSSSDS	300
WIPGPDQVA	RGYQSSATMS	DGRVFTIGGS	WSGGVFEKNG	EVYSPSSKTW	TSLPNAKVNP	360
MLTADKQGLY	RSDNHAWLFG	WKKGSVFQAG	PSTAMNYYT	SGSGDVKSAG	KRQSNRGVAP	420
DAMCGNAVMY	DAVKGKILTF	GGSPDYQSD	ATTNAHIITL	GEPGTSPTV	FASNGLYFAR	480
TFHTSVVLPD	GSTFITGGQR	RGIPFEDSTP	VFTPEIYVPE	QDTFYKQPNP	SIVRVYHSIS	540
LLLPDGRVFN	GGGGLCGDCT	TNHFDAQIFT	PNYLYNSNGN	LATRPKI TRT	STQSVKVGGR	600
ITISTDSSIT	KASLIRYGTGTA	THTVNTDQRR	IPLTLTNNGG	NSYSFQVPSD	SGVALPGYWM	660
LFVMSAGVP	SVASTIRVTQ					680

SEQ ID NO: 5 moltype = AA length = 559
 FEATURE Location/Qualifiers
 source 1..559
 mol_type = protein
 organism = Phanerothecium chrysosporium

SEQUENCE: 5

MLSLAVVSL	AAATLAAPAA	SDAPGWRFDL	KPNLSGIVAL	EAIVVNSSLV	VIFDRATGQD	60
PLKINGESTW	GALWDLDTST	VRPLSVLTDS	FCASGALLSN	GTMVSMGGTP	GGTGGDVAAP	120
PGNQAIRIFE	PCASPSGDGC	TLFEDPATVH	LLEERWYPSS	VRIFDGLSMI	IGGSHVLTFF	180
YNVDPANSE	FFPSKEQTPR	PSAFLERSLP	ANLFPRAFAL	PDGTVFIVAN	NQSIIYDIEK	240
NTETILPDIP	NGVRVTNPID	GSAILLPLSP	PDFIPEVLVC	GGSTADTSLP	STSLSSQHPA	300
TSQCSRIKLT	PEGIKAGWQV	EHMLEARMP	ELVHVPNGQI	LITNGAGTGF	AALSAVADPV	360
GNSNADHPVL	TPSLYTPDAP	LGKRISNAGM	PTTTIPRMYH	STVTLTQQGN	FFIGGNPNM	420
NFTPPGTPGI	KFPSELRIET	LDPFPMFRSR	PALLTMPEKL	KFGQKVTVPI	TIPSDLKASK	480
VQVALMDLGF	SSHAFHSSAR	LVMMESSISA	DRKSLTFTAP	PNGRVFPFGP	AVVFLTIDDV	540
TSPGERVMVG	SGNPPPTLE					559

SEQ ID NO: 6 moltype = AA length = 558
 FEATURE Location/Qualifiers
 source 1..558
 mol_type = protein
 organism = Pycnoporus cinnabarinus

SEQUENCE: 6

MAPTAFSLVS	ALALASLSLA	APSAPGWSFD	LKKETSGIVA	LEAIVVSPTL	VVFFDRASDD	60
PLQINNHSWA	GALWNLESST	VRPLDVLVTS	FCASGALLSN	GTMASVGGDP	DGFVGNPAIR	120
PGNQAIRLFE	PCDSPTGDGC	TLFEDPATLH	LLEKRWYPSS	ARIFDGLSII	VGGMHEATPF	180
YNTDPALFPE	FFPRKEDTPR	PSEFLNRSPL	ANLFPRAFAL	PDGKVFVMAN	NQSIIYDIEA	240
KTERILPDVP	NNVRVTNPM	GSAILLPLSP	PDFVPEVLVC	GGSTADTIDP	SLLTSQTPAS	300
SQCSRIRLDE	EGIAKGEWEV	HMLEGRIPE	LVHLPNGQVL	IANGGRTGFA	AIASVSEPVG	360
NSNADHPVLV	PSLYTPDAPL	GRRISNVGLP	SSGIPRLYHS	SVTLTPQGNF	LIAGSNPNNR	420
TTVGPQIKFP	SEFRVQTLDP	PFMSVERPKI	LNMPKLGFP	KSFTVPIVSV	SSLARPGAKV	480
QISLMDLGF	SHAFHSSARL	VFMNGKISQD	SKSLTFTTTP	NGRVYPPGPA	TVFLTIDDV	540
SEGAWMMGSS	GNSPPTLE					558

SEQ ID NO: 7 moltype = AA length = 506
 FEATURE Location/Qualifiers
 source 1..506
 mol_type = protein
 organism = Colletotrichum graminicola

SEQUENCE: 7

MPTLRSLARN	LPAALLALAA	ACEAQNKGK	GPMVKFPVVP	VAVALVPETG	NLLVWSSGWP	60
NRWTTAGNGK	TYTSLYNVNT	GNISDAIVQN	TQHDMFCPGT	SLDADGRIIV	TGGSSAAKTS	120
VLDPKKGESS	PWTPLSNMQI	SRGYQSSCTT	SEGKIFVIGG	SFSGAGTRNG	EVYDPKANTW	180
TKLAGCPVKP	LVMQRGMFPD	SHAWLWSWKN	GSVLQAGPSK	KMNWYDTKGT	GSNTPAGLRG	240
TDEDSMCGVS	VMYDAVAGKI	PTYGGGKGYT	GVDSTNAHI	LTLGEPQAV	QVQKLANGYQ	300
NRGFANAVVM	PDGKIWVVG	MQKMWLFSDT	TPQLTPELFD	PATGSFTPTT	PHTVPRNYHS	360
TALLMADATI	WSGGGLCGA	NCKENHFDGQ	FWSPPYLFEA	DGVTPAKRPV	IQSLSDTAVR	420
AGAPITITMQ	DAGAYTFSMI	RVSATHTHTVN	TQRRIPLDG	QDGGDGKSFT	VNVPNDYGVA	480

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 IPGYMMLFAM NEAGVPCVAQ FFKVTL 506

SEQ ID NO: 8 moltype = AA length = 497
 FEATURE Location/Qualifiers
 source 1..497
 mol_type = protein
 organism = Colletotrichum gloeosporioides

SEQUENCE: 8
 EAEAQGLGQW SPLIKFPVVP VSVALLPESG NLLVWSSGWP NRWTFAGNGK TYTSLYNVQT 60
 GNVSDAVIQN TQDMFCPGT SLDAEGRIIV TGGSSAAKTS VLDPFKNGESS SWTALSNMQI 120
 SRGYQSSCTT SEGKIFVIGG SFSGAGTRNG EIYDTATNKW TKLAGCPVKP LVMQLGMFPD 180
 SHAWLWSWKN GSVLQAGPAK QMNWYDTKGT GANTPAGLRG ADQDSMCGVS VMDAVAVAGKI 240
 FTYGGGKGYT GYQSTNSNAHI LTLGEPGQQV QVQKQLONGQY NRGFANAVVM PDGKIWVVG 300
 MKQMALFSDA TPQLTPELFD PATGKFTPTA AHTVPRNYHS TALLMADGTI WSGGGGLCGA 360
 GCAANKPFDG FWSPPYLFEA DGKTPAKRPV IESLSDETVK AGAALTINMQ DEGKYTFSMI 420
 RVSATHTTVN TDQRRIPLDG QDGGDGKSFS VNMPSDYGVV IPGYMMFAM NEAGTPCVAK 480
 FFKVSLHHHH HHHHHV 497

SEQ ID NO: 9 moltype = AA length = 711
 FEATURE Location/Qualifiers
 source 1..711
 mol_type = protein
 organism = Colletotrichum graminicola

SEQUENCE: 9
 MVRSCAYKAI AAASLLSQA SAAITSCPNN ETVWETPIGV KYTLCPGSDY QNGGASLQTV 60
 RDIQSSLECA KICDSDARCN RAVYDNNKA CDVKNSTNPM QWAADDRFET IRLTNDLPEG 120
 AFISTCSFNE TSYRVPETNA EYRIPCDDTY TGVNAKVVEG VTTIQACAEI CSNTQDCRKS 180
 VFDHINNACA IKAABPATSI FWVQDKQFST IRLPENIDPA VKGKWGLIR LPVIPVAAAI 240
 VPSYPEPSRL LFFSWSNDA FSGASGMTQF GDYDFATGAI SQRTVTNTHH DMFCPGISQL 300
 EDGRILIQGG SDADTVSIYD PATNEFTRGP NMTLARGYQT SCTLNKGKVF TIGGAYSGER 360
 VGKNGEVYDP VANAWTYLPG ADFRPMLTND HEGIWREDNH AWLFGWKNGS IFQAGPSKDV 420
 HWYGIQNGT VAKAATRDDD DAMCGVWVMY DAVAGKIFSA GGSPTYDTSF ATQRAHITTI 480
 GEPNTPAEVE RVADMGFPRG FANAVVLPDG QVLVTGGQRM SLVPTNTDGI LVAELFNPET 540
 REWKQMAPMA VPRNYHSVSI LLPDATVFSG GGGMCWVQNV GDSTAGCDKT VDHSDGEIFE 600
 PPLYFNEDGS RAARPVISAI SADPIKAGAT LTFTEGVEG QGTAALIRLG SVTHSVNSDQ 660
 RRVPLNVTVS GNEYSATLPD DYGILLPGYY YLFVSTPQGT PSIAKTVHVI L 711

SEQ ID NO: 10 moltype = AA length = 611
 FEATURE Location/Qualifiers
 source 1..611
 mol_type = protein
 organism = Streptomyces lividans

SEQUENCE: 10
 STEKYHQYKI NQPEYKAANG KWEIIEFPEK YRQNTIHAAL LRTGKVLMA GSGNNQDNSD 60
 DKQYDTRIWD PVKGTIKKVP TSPDLFCTGH TQLANGNLLI AGGTKRYEKL KGDVTRKAGL 120
 MVVHNENPDK PITLPAGTKF TGKENGKTFV SKDPVLVPRV EKVFDPATGA FVRNDPGLGR 180
 IYVEAQKSGS AYETGTEDNY RVQGLSGADA RNTYQIAQKL ALDKKDFQGI RDAFEPDPA 240
 EKYIKVDPMH EARWYPTLTT LGDGKILSVS GLDDIGQLVP GKNEVYDPKT KAWTYTDKVR 300
 QFPPTYPALFL MQNGKIFYSG ANAGYGPDDV GRTPGVWDVE TNKFTKVPGM SDANMLETAN 360
 TVLLPPAQDE KYMVIQGGGK GESKLSSEKT RIADLKADDP KFDVDPSEK GTRYPQASIL 420
 PDDSVLVSGG SQDYRGRGDS NILQARLYHP DTNEFERVAD PLVGRNYHSG SILLPDGRML 480
 FFGSDSLYAD KANTKPKGFE QRIEITYPPY LYRDSRPDLS GGPQTIARGG SGTFTSRAAS 540
 TVKKVRLIRP SASTHVTVDV QRSIALDFKA DGGKLTVTVP SGKNLVQSGW YMMFVTDGEG 600
 TPSKAEWVRV P 611

SEQ ID NO: 11 moltype = AA length = 646
 FEATURE Location/Qualifiers
 source 1..646
 mol_type = protein
 organism = Actinobacteria bacterium

SEQUENCE: 11
 MSELGGGRRR ARR LAVGTVV VLALAGMNGP WVYRFSTEKY HEYAINRPEY KAENKWDII 60
 QFPKKYRQDT IHAVLLHTGK VLLIAGSGND QNFDKAKFD TRIWDPVKGT IKKVTPADL 120
 FCTGHTQLAN GNILIAGGK KYEKLKGDVK KAGGVMI LYN ENPKPITLP KGTVFVTKQ 180
 GKSFVSTDSS LIPRATKVPD KKTGKFLRNN PGYARVVVEA PKEGTYETG TEDNYTIQGL 240
 TGADERNTYG IANKLGMDKR DFEGIKNAYE FDPVAEKYIK VDPMNEARWY PTLTLLSDGK 300
 VLSVSGLDDI GQLVPGKNEI FDPKTKTWTY TTKRQFPPT PALPLMQNGK LFGVSGNAGY 360
 GPDDVGRDPG IWDVDTNAFT KLPGLSDPNM LETSATVLLA PAQEEKPMVI GGGVGVGESK 420
 STNKTRLIDL K DANPKFVDG PTLKKGTRYA QASVLPDDSV LVSGGSQDYR GRSASDIKQA 480
 RIYNPTNTF KRVADPEVGR DYHSGSILLP DGRVMFFGSN PLYADKANTK PGVFEQRIE 540
 YTPPYLYRDA RPDLSGGPPT IARGASGTF SKHASSIRKV RLIRPSSSTH VTDVDQRSIA 600
 LDYTTSGDKI TVTVPKNRNL VPSGWYMMFV DDDQGTPSKA VWVDVP 646

SEQ ID NO: 12 moltype = AA length = 666
 FEATURE Location/Qualifiers
 source 1..666

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mol_type = protein
organism = Streptomyces sp.

SEQUENCE: 12
MTRQSGRGAR  RSAGPVSRRS  GARARRSGLR  TRRFAIGTVA  VVALAGANGP  WLYRFRTTERY  60
HHYTIINTQSY  KEANGHWDPL  DIPAEFRVNA  VHAALLRTGK  VLLVAGSGNN  EKNFDAKSPR  120
SILWDPKTNA  FKDIHTPKDL  FCAGHTQLPG  GNLLIAGGTK  RYEKLGGDVT  KAGGLMIVRN  180
ENPRKPIITLP  AGTRFTGKES  GKTPI SQDPL  LVEKATKVFD  KKTGEYQRNE  AGVGRIYVEA  240
EKSGRKYETG  TEDNYRISGL  STEDTRNVYG  IAQKLALDKK  DFQGINDAYE  FDPVAEKYIP  300
VDPMDKARWY  PTLTTLKDGK  VIAVSGLDDI  GQIDVGKAEI  YDPKKKSWAP  SGIVRKFATY  360
PALPFLNDGK  LFYSGSNSGY  GLQSEGRTPG  IWDLKTNDFO  EVPGLVDADQ  METSATVRLP  420
PAQDERFMVI  GGGVGVESDK  STPKSRLVDL  QQKNPKFTEG  ASLDKGTFRP  SASLLPDDSL  480
LVAGGAGDYR  GRGGSDVLEA  RLYDAKSPTY  KRVADPAVGR  NYHSGSVLLP  DGRVMIFGSD  540
PLYMDKINTR  PGTFFQRIEI  YTPPYLYRGS  RPELTAGPKS  IKRGGTGMFT  TQHASKITSA  600
KLMRPSAVTH  VTDDTQRSIA  LELEKSADGI  TVTVPKNRAL  VPSGWYMLFV  TDAKGTPESE  660
TWVEVP

SEQ ID NO: 13      moltype = AA length = 663
FEATURE           Location/Qualifiers
source            1..663
                 mol_type = protein
                 organism = Streptomyces abyssalis

SEQUENCE: 13
MKYRPSRRTR  RTAIGAAVVL  VIAGFNGPAL  YGIASEKYHD  YKINQPEYKA  ENGHWKTQVQ  60
PEKYRINTIH  AALLHTGKVL  LVAGSGNDAK  QFNAKTFRTV  LWDPVKNTFK  NIPTPNDLFC  120
AGHTQLPDGK  LLVAGGTRRY  EKLGGDVKKA  GGLMI IHNEN  PDRAKTVKAG  TRFTGRKNGK  180
TFEAKDVLV  PRAKKTNPA  TGKVEVTPSV  ARVYVEAIKR  GKKYQTGTED  NYRLSGLRGA  240
ERRNIYGIAQ  KLSFDKKDFQ  GIKDTYEFDP  VABRYLKTDF  MDEARWYPTL  TTLEDGKVL  300
VSGLDDIGQV  VPGKNIYDP  QTKKWEFLPR  KRFFPTYPAL  HLTDDNKIFY  SGANAGYGPA  360
DKGRTPLGWN  LETNGFRKIP  GMSDPDKLET  AMSVPLPPTQ  SRKFMVLGGG  GVGESKKASE  420
KTRIVDLKDP  EPRFQDGPL  YAKARYPSSV  ILPNDKVLTT  NGSGDYRGRS  ESNILKAEIY  480
DPADNTKKPV  ADPIVGRNYH  SGALLLPDGR  VMTFGSDSLF  SDKANTKPGK  FEQQIDIYTP  540
PYLHTKGERP  ELENTGRGEQ  TVRLGGTASF  NSKDAGSIKK  MRLVRPSSFT  HVTNVEQSSV  600
ALKFKKSADG  VTKLPEPDS  LVPPGWMVMT  AVDDKGGPKS  SVWLKVPRTA  SAQAPTTPPL  660
SPP

SEQ ID NO: 14      moltype = AA length = 652
FEATURE           Location/Qualifiers
source            1..652
                 mol_type = protein
                 organism = Streptomyces indicus

SEQUENCE: 14
MTVGSTPVKH  TRRRARRIAI  GAAVVLALAG  MNGPALWRFA  SDQYHDYKIN  RPEYKADNGK  60
WDIVLEPDEY  KLMTIHAALL  HTGKVLVLAG  SGNQKNPDA  KKFDTVLWDP  VKNTYKKIPT  120
PNDLFCGTGT  QLPDGKLLVA  GGTKSYEKLL  GDVTKAGGVM  NVYNENPDEP  ITLAKGKLT  180
GKENGKTFVI  TESIVVERAK  KRFDKETGKF  LGNTPGYARV  YVEAEKRGKK  YETGTQDNYK  240
IVGLSGADAK  NTYGIAQKLA  LDKKDFQGD  DAYEFDPAE  KYIKVDPMNE  ARWYPTLTTL  300
EDGRVLALSG  LDDIGQIVPG  NRNEVYDPDT  KKWTYVKQNR  QFPTYPAIFL  LPNGKLFYSG  360
ANAGYGDDV  GREPGIWDFD  TNGFTKLGK  SDPKLMEAG  TVWLPPAQDE  KFMVVGGGV  420
GESEQSSDKT  RIIDLSEAP  EFKDGPSPDK  GTRYQSSVL  PDDSVLISGG  SEDYRGRSGS  480
NILEARMYDA  KTGEMRRVAD  PLVGRNYHSG  SLLLPDGRVV  PFGSDSLFAD  EANTKPGEPE  540
QRLEIYTPPY  LYQGAQPTLT  GGPDSIERGE  SATFKTQHAS  SIKNARLIRP  SASTHVTDDID  600
QRSIALDVKK  GADSVTVTVP  ENKNLVQPGW  YMLFVTTDQG  VPSKAQWVEI  PE  652

SEQ ID NO: 15      moltype = AA length = 645
FEATURE           Location/Qualifiers
source            1..645
                 mol_type = protein
                 organism = Streptomyces diastatochromogenes

SEQUENCE: 15
MKDQAGRRA  RRLAIGTAVV  LALAGMNGPW  LYRFGTEQYH  QYKINKPEYK  AANGHWDIVD  60
FPEEYRQDTI  HAALLHTGKV  LLVAGSGNNQ  DNFDKKKFDT  RIWDPVKGTI  KKVPTPADLF  120
CTGHTQLANG  NLLIAGGTRK  YEKLKGDVTK  AGGLMVVHNE  NPDKPMTLPA  GTKFTGKENG  180
KTFVSKDPVL  VPRAKKVDFP  KTGAFLRNDP  GLGRIYVEAQ  RSGTQYETGT  QDNYRIQGLT  240
GADARNTYGI  AQKLALDKK  FQGRDAYEF  DPVAERYIKV  DPMNEARWYP  TLTTLSDGKI  300
LSVSGLDDIG  QLVPKNEIF  DPKTKKWTYT  QKIRQFPPTY  ALFLMONGKI  FYSGANAGY  360
PDNVRVPGI  WDVGTNKFTK  LPGLSDPNMM  ETAGTVLLPP  AQDEKFMVIG  GGGVGESKLS  420
SKKTRLIDLK  AQMPHFVDP  QLEKGTYPQ  SSILPDDTVL  VSGGSEYR  RGASNILQAR  480
MYHPDSNSFT  QVADPLVGRN  YHSGSILLPD  GRVMFFGSDS  LYADKANTKP  GKFEQRIEY  540
TPPYLYRDSR  PSLSGGPQTI  ARGASGTFTS  KHASSIKKVR  LIRPSASTHV  TDVDQRSIAL  600
DFTTSGDKIT  VTPKSRNLV  QSGWYMLFVD  DDQGTPSKAQ  WVKVP  645

SEQ ID NO: 16      moltype = AA length = 658
FEATURE           Location/Qualifiers
source            1..658
                 mol_type = protein
                 organism = Streptomyces carminius

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SEQUENCE: 16
 MTYRPSRRFR RIAIASAVVA ALAGANGPAL YRFGGEAWHD HRINQPGYKA ANGHWRTVDV 60
 PERYRVNTHI AALLHTGKVL LIAGSGNDAE QFRAKTFRTV LWDPEKNTFA NIPTPKDLFC 120
 AGHTQLPDGR LLVAGGTQRY ETLEGDVEKA GGLMTVHNED PDKPVTLPAG TRFTGRANGK 180
 TFAQQAPVVV PRAGKTTAPG TGKVTVTVST ARVYVEALKK GRRHQTGTQD QYRVHGLKGD 240
 DRNRYVIAIQ KLSFDKDFQ GIRDSYEFDP VAERYVGVDP MNEARWYPTL TTLADGSLVS 300
 VSGLDEIGQV VPGRNEIYDP ETQKWSYLPE ERFPPYYPAL FLAAGGRILY TGSNAGYGPA 360
 DEGRTPGLWD LESNEFREV PGLGDPDVLET SMSVLLPPAQ EQRYMVLGGG GVGESRRATA 420
 RTGIVDLTEP EPRFTPGPDL YAPVRYPSSV ILPDDTVLTT NGSGGYRGRG GSDVLRAGLY 480
 DARTDTARTV ADPLVGRNYH SGALLLPDGR VMTFGSDSLF ADKADTRPGE FQQQIDLYSF 540
 PYLFRGKRPE LRGTGEPRTV ELGGKVTFFAS EDAARVRTAR LVRPGSPTHV TNVEQRSVAL 600
 EFERTADGLT FTLPPDDPSLV PPGYYMLNVL DGBGVPSSEV WVKVPVSPDA EPVTVAAD 658

SEQ ID NO: 17 moltype = AA length = 647
 FEATURE Location/Qualifiers
 source 1..647
 mol_type = protein
 organism = Streptomyces sp. Ru73

SEQUENCE: 17
 MARRRTRFKE RKTWLGIGAV LVLAAPNAPA ATGFAERVYH DYAINRDAYK AENGHWVDVLD 60
 VPPEFRQNTI HAALLRTGKV LLVAGSGNNV KNFDAKRFTS VLWDPVRNSF KKIHTPNDLF 120
 CTGHIHLADG KLLFAGGTRK YEKLKGDVTK AGGLMIVHNE NPNEFVRLPA GTVFTGRRNH 180
 RTFVAKDPIR IRPAKKVDFK ETGRFLRTEA SVDRIYVEAR KKGROYEAGA QDNRYVRGLT 240
 RARAHNVYGI ATKLLGLDKKD FQGIHDAYEF DPLAEKFTKV QSMHEARWYP TLVRLSDGRV 300
 LALSGLDDIG QVVPKSEIY DPKSRTWAYS GKVRQPPTYP AVFPLSDGTL FYSGSNAGYG 360
 PADVGRTPGI WHFRMTTFK LPGLDDAGLM ETSGTVLLPP AQRQKFMVVG GGGIGESAKS 420
 SKKTRLIDLTL QPHSPFHDGP SLDEGTRYPI LSVLPDDTVL VTGGSKDYRG PVAENILQAR 480
 LYDARSGMTMR RVADPQVGRN YHSGAILLPD GRVLTFGGDS LFADKAKTQT GTFEQRLLEVY 540
 TPPYLRYGAR PALSHGPTTV GRGATAVVRT KDAAAVKTAR LLHPSSTHVV TDVEQRSVAL 600
 DFTRTKSGDG IEVTLPKNRR LVPSGWYMLF VTNRQGTPTST ARWVKVP 647

SEQ ID NO: 18 moltype = AA length = 667
 FEATURE Location/Qualifiers
 source 1..667
 mol_type = protein
 organism = Streptomyces sp.

SEQUENCE: 18
 MAHRPSRRPV RILLACGAVL ALAAANAPAV YDFTSERYHA YEIDQADYKA EKGHWRTVDV 60
 PEKYRINTIH AALLNTGKVL LIAGSGNDAK QFDAKTFRTV LWDPKKNSFK EIDTPKDLFC 120
 AGHTQLPDGN LLVAGGTQRY ETLKGDVQKA GGLMIVHNEN PDKPKTLPEG TRFTGQENGK 180
 TFVSQASVLI PRAKKTVDKK TGKAKVTASS ARVYVEAPEK GEKYQTGTQD QYRVGLKGD 240
 EKQNIYGIAQ KLSFDKDFQ GIKDSFEFDP VAERYITVDP MAEARWYPTL TTLEDGKVLVS 300
 VSGLDEIGQV VPGKNEIYDP KSKWKYLPK SRFFPYYPAL FLTADSRIFV TGSNAGYGP 360
 DQGRVPLGLWD LDSNKFSEVP GLSDPDILET SMSVLLPPAQ DQKFLVLGGG GVGESPKSTD 420
 RTGIADLKAD KPRFTDGPDL YDKARYPSSV ILPDDTVLTT NGSGDYRKGK ASNVLKASLY 480
 DPGTNKFAEA AAPLVGRNYH SGGLLLPDGR VMTFGSDSLF SDKADSKPGK FEQTLEIYTP 540
 PYLHKGGERP ELRDSGDARR TVKLGASADF RSKDAASIKK MRLIRPGAFT HVTNVEQRSI 600
 AMDFKRTADG VRVTVPHNPS LVPPGWYMLT AVDDRGVPSV SVWLNVPVSK DAPHTKTAD 660
 APAPAPN 667

SEQ ID NO: 19 moltype = AA length = 654
 FEATURE Location/Qualifiers
 source 1..654
 mol_type = protein
 organism = Streptomyces sp.

SEQUENCE: 19
 MLPCCQASAA LTARNPCLPA NAHSHCSVSQ APELATLEYA SPPVSTPYHD YTIKPGYKA 60
 ANGHWDLLDV PAGYRINTIH AALLHTGKVL LVAGSGNNRK NFDKASFRSV LWDPRTGAFK 120
 NIPTPDDMFC SGHTQLPDGK LLIAGGTRRY EKLKGDVTKA GGLMIVHNEN PDRPVTLPA 180
 TRFTGRANGR TFSKDPVLV EKATKVFDRR TGRFLHNDAG LGRIYVEAEK SGARYATGAE 240
 DNYRIAGLTG SDTRDYVYGA QKIADLKKDF QGIREAFEPD PVAEKYISVD PMNEARWYPT 300
 LTTLKDGRVL ALSGLDEIQV IVPKDEIYD PRTKKWRYTG IVRFPYTPA VFLLGDGKLF 360
 YSGSNAGYGP ADVGRDPGVW DLATNSPRKI PGLGDADETE TSATVRLPPA QDERFMVIGG 420
 GGVGESDRSS AKSRLVGLKD PAPFRFDGAA LAEGTRYPGV SLLPDDSVLV TGGSSDYRGR 480
 GGSNVLQARL YDPGTDRYRR VADPAVGRNY HSGSVLLPDG RVMIFGSDSL YADRANTRPG 540
 VFEQRIEITY PPYLRYSSRP VLTGGPRSLR RGGTGGFTTA PGRAITSAKL IRPSAVTHVT 600
 DTDQRSVALG LRRTPGGITV TVPGNRALVP SGWYMLFVTD DRGTPSEGRW VEVP 654

SEQ ID NO: 20 moltype = AA length = 667
 FEATURE Location/Qualifiers
 source 1..667
 mol_type = protein
 organism = Streptomyces mobaraensis

SEQUENCE: 20
 MRLRHLDRR SGTVIGRRRT RTAVGAAVVL VLAGMNGPAI WGYASDQYHE YKINKPEYKA 60
 KNGHWDVVDV PDEYRINTIH AALLQTGKVL LVAGSGNNA NFAAKTFRTV LWDPERNTFK 120

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NVPTPKDLFC	AGHTQLPDGK	LLVAGGTQRY	EKLEGDVTRA	GGLMIVHNEN	PDKPMTFFPAG	180
TRFTGKQSGK	TFESKDAVLV	PRAKKEQGD	GKVKVTVSTA	RVYVEALEKG	QEHATGLTDN	240
YRIEGLTGDD	ARNFYGIANK	LSFDKKDFQG	IKDAFEFDPV	AERYVTVDPM	NEARWYPTLT	300
TLQDGKVLVS	SGLDEIGQVV	PGKNEVYDPK	TKKWTYLPQE	RFFPTYPALF	LTDKGGKIFYT	360
GSNAGYGPAD	KGRDPGVWDL	GNSFIPVPG	ISDPDALETS	MSVLLPPAQD	QRYMVLGGGG	420
VGEDKKSTAR	TRIVDLHTER	PRFHDGPDLY	AKARYPSSVI	LPDDTVLTTN	GGSDYRGRSA	480
SNVLKAEIYD	PKANASHRVA	DPLVGRNYHS	GALLLPDGRV	MTFGSDSLFR	DKDNTQPGVF	540
QQQIDLTYTP	YLFHKGDRPE	IRDTRRIVK	LGDKTTYRIT	SAHGVAKARL	IRPGSFTHVT	600
NIEQRSIALD	LKKEGTDRFT	VTLPKDPSLV	PPGWYMVIVV	DEEGTPSKAV	WVKVPAAEKR	660
DDEKKEE						667

SEQ ID NO: 21 moltype = AA length = 506
 FEATURE Location/Qualifiers
 source 1..506
 mol_type = protein
 organism = Colletotrichum tofieldiae

SEQUENCE: 21						
MVTVCVLR	LPAALLAF	ASEAQN	VMKFPV	VAVALPET	NMLVSSGWP	60
NRWTTAGNGK	TYTSMYDVK	GKVS	DALIQN	TQHDMFC	PGT SMDADGRIIV	TGGSSASKTS 120
VLDPKKGESS	SWAPLANMQI	SRGYQSSCTT	SEKIFLIGG	SFSGAGRRDG	EIYNPKANTW	180
TKLPGCPVKP	LVMQRGMFPD	SHAWLWSWKN	GYVLHAGPAK	QMNWYDTKGT	GANTPAGLRG	240
ADDDSMCGVS	VYDAVAGKI	PTYGGGKAYT	GVTSSNAHI	LTLGEPQAV	QVQKLONGKF	300
NRGFANAVVM	PDGIWVVG	MRQMLFSDN	TPQLTPELFD	PATGNFTPTT	PHAVPRNYHS	360
TALLMADATI	WSGGGLCGA	NCKENHFDGQ	FWSPPYLFEA	DGKTPAKRPV	IQNLSETDVK	420
AGAPITITMQ	DAGTYTFSML	RVSATHTVTN	TDQRRIPLDG	QDGGDGKDF	VNIPDDYDIA	480
VPGYMLFAM	NEAGVPCVAK	FFKVTL				506

SEQ ID NO: 22 moltype = AA length = 680
 FEATURE Location/Qualifiers
 source 1..680
 mol_type = protein
 organism = Moelleriella libera RCEF 2490

SEQUENCE: 22						
MSRFLLLAL	VAVASSAAVE	QQQQQQQPR	AIKILEHYRE	SSTFTKLF	AAPIGNEIPKN	60
AYTVTCDSYQ	PGNECALAID	GNNNTFWHTA	FSGANLPHQI	VVDLGATRNI	NGLSALPRQD	120
GNNHGFIQAH	EVAVSTDNRN	WEVVASGTWY	GGDSTTKFAN	PETRISIRYR	LKALSEAYGN	180
QWTSVAEVKV	YEAKTGPAAE	AGTGKKGPTI	NFPTIPVAGA	VDPLTQVLI	WSSYTYDNYL	240
GSSQDRVFTS	IWDPTSTGVVT	PKLVDNTNHD	MFCPGISIDG	AGRMVITGGN	SAQKTTVYQF	300
GSQTIWPGPD	MNTQRGYQAS	ATLSDGRVFT	IGGCWGGWF	EKNGEVYDPK	ASTWTSLPGA	360
LVHPLMTNDA	QGIYRADNHA	WLFGWKNGSV	FQAGPSTAMN	WYTTSGNGSV	APAGNRTSSR	420
GDAPDAMTGN	AVMYDAVNGK	ILSPGGSPSY	QDSSATNAH	IITIGSPGAP	AQSRFASNGL	480
WSPRAFHTSV	ILPDGKTFIT	GGQTYAVPFS	DDNPDLTPEM	YDPVADSFVQ	QQANTIIRVY	540
HSISLLLPDG	RVFNAGGGLC	GDCTTNHFDG	QIFTPQYLLT	RTGQLAARPA	INSVLSGRR	600
LTINTNSAIT	SAALMRYGSA	THTVNTDQRR	IPLKLTGT	NRYTADAPSD	PGILLPGYIM	660
LFVLSQGV	SVAKTVNPLV					680

SEQ ID NO: 23 moltype = AA length = 685
 FEATURE Location/Qualifiers
 source 1..685
 mol_type = protein
 organism = Purpureocillium lilacinum

SEQUENCE: 23						
MKLLGTVAAL	ALCDAASHTY	AVAVKPRPHT	AMPNLAARGT	EMSLMAAKPI	GNAINRAGWK	60
VTCDEGEEQN	ECAKAIDGDN	NTMWHATAWQN	DNPPPHIT	VDMGSAQTIN	GISVLPRQDG	120
SEHGWIARHD	VLVSDNGQTW	GDPVATGTWY	TDATAKYANF	EPRSARYVRL	VARSEAQGRP	180
WTSIAELNVY	RADGPPVPKN	GIGKWGLTLD	FPVVPVAGVV	DPLTGKVVVV	SAYENDQYEG	240
SPGGWTLTST	WDPATGEVTE	RNVTNIGHDM	FCPGVSLDAS	GRVVVTGGSN	AQKTSFYDAA	300
TEAWVPGPDM	KTRPGYQASA	TCSDGRVFTI	GGSWGGQFE	KNGEVWDEPK	NSWRALPGA	360
VKPLMTKDRG	GIYRADNHAW	LFGWRNGSVF	QAGPSTAMNW	YYTAGDGRVR	SAGQRRAPRG	420
ADPDAMCGNA	VMFDATAAGKI	LTVGGAPHYE	DADATNAHV	LTLGDAGAAP	KVVFAGNGMA	480
HPRIIFANAVV	LPDGTVFVVTG	GQQAELFKD	TPQLTPELY	DPALGAFVEQ	APNSVVRVYH	540
SMALLLPDAT	VLSGGGLCG	GACDTNHFDA	QVFSRYLFD	GEGQPARPK	IRAVASKEVH	600
AGDAIKVTTD	GPVKSAAVLR	YGSATHSVNT	DQRRVPLTLR	QDGGSYAYS	DLPRDLGVLL	660
PGYWMLFVMN	DKGVPSVAAT	VKVL				685

SEQ ID NO: 24 moltype = AA length = 671
 FEATURE Location/Qualifiers
 source 1..671
 mol_type = protein
 organism = Penicillium antarcticum

SEQUENCE: 24						
MHAKWLSGLL	LLGLSQAKDN	SQALQTAATT	NLQSAEEDPY	ATNQAPPIGE	KLSRRRWTVT	60
CDNSQDTECT	KMLDDNFDTF	WQSGNDGSIH	EVVVDLSRAE	NIQALSIVSR	RDATTGQIF	120
AHQIYVSNDK	ENWGNPVAFG	TWYNTTDEKL	AVFEPKEGQY	LRLVSLDGNA	ASISELQIYD	180
SDSPKQVSN	GWGPTIDLP	LVAVSGAVLP	GTGQVLVWSS	WAKDDYLHST	SQTLTALWDP	240
KTGNVTQRTV	QDTGDMFCA	GTSWDGNNEL	LVTGGNNDGQ	TSILDPATDT	WYPGAAMKLG	300

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RAYHASATNS	EGRVFIIGGS	WNGGTNNNRD	GEIYDPVAEE	FTGLPGARVE	PMWTNDANGG	360
YRRDASHGWF	GWKNGTVFQA	GPSKAMNWYY	TSGQGDQKPA	GNRGDADDAM	SGNAVMFDAV	420
NGKIISFGGS	PSYENTYATK	DAHLEIEGEP	GTDPKVSTAK	NPSGDGMAFA	RVFHTSVVLP	480
NGNVFVAGGQ	TYATPFNDSG	AQFTPEIYDP	ATNEFRQQGT	NSVVRVYHSI	SLLLTDRVVF	540
NGSGGLVSA	PSNHFDAQIY	TPDYLLNNDG	TLATRPTIDS	VDKRTLHPGE	KLIVKASIKL	600
TKASLIRYGS	THTMTMDQR	RVVLDSWAAN	GETYETTLPS	EAGALLPGPW	MLFILNDDGV	660
PSIAETIYIE	V					671

SEQ ID NO: 25 moltype = AA length = 673
 FEATURE Location/Qualifiers
 source 1..673
 mol_type = protein
 organism = *Penicillium flavigenum*

SEQUENCE: 25

MRPRWLTGLL	LLGLAHADND	NSVISDKQVA	NQDHSATTD	YATTQAPPRG	EKLSRRRWSV	60
TCDSQSQSGNE	CEKLLDNDLQ	SFWLSGAEGN	THEVTIDLSR	AENVHAI SVV	SRQDGTKTDGQ	120
IAAHQVPSVSG	DKNDWGEPEVA	YGTWYEGTGE	KLAIFEPKEG	QYVRLVSLGS	NVASISELNI	180
YDGADPKTVP	NGGVWGPTID	LPVVAVSGAV	IPETNEVLWV	SSWAKDDYLH	SRGYTLTAVW	240
NMNDNSVTQR	KVQETHDMF	CSGMSYDGKG	ELLVLTGGNND	KSTSIFDPAS	GNWTEGNTMI	300
ISRGYQASAT	LADGRVFIIVG	GSWNGGTNYD	KDGEIYDPNT	EEFRFLKNAL	VKPMWTDYDYN	360
SGYRRDASHGW	LFGWKNDFV	QGGPSKMNW	YYTEGDGQK	PAGTRADASD	SMSGNAVMFPD	420
AVNGKIITFG	GSPSYENSVA	TTDAYLIEID	EPGSQPKVTA	AKNPNGEEMA	YARFHTSVV	480
LPDGGVFTAG	GQSYGVFPND	SNAHLTPELY	DPKTNQFNEQ	QPNISIVRVYH	SISLLELPDGR	540
VFNGGSLGV	SAPTNHFDAQ	IYSPHYLFNQ	DGSLATRPTI	DGVVEKKLRA	GDKLTISASI	600
DVKNASLIRY	GTTTHTVNTD	QRRIALDSWT	ANGDSYETTL	PGDNGILLPG	PWMLFILNDD	660
GVPSVSQTIH	IQV					673

SEQ ID NO: 26 moltype = AA length = 684
 FEATURE Location/Qualifiers
 source 1..684
 mol_type = protein
 organism = *Fusarium sp. AF-8*

SEQUENCE: 26

MWSPILTCLL	TGLFWDEAMA	IVVPAQNSTK	VKVQMEHLSM	RATGPLGTAI	NRNNWKVTC	60
SDNPDSDNAC	QKAIDGDVNS	FWHTAWFEDT	SKDPGLPHTL	TVDMKTVKNV	NGISALPRQD	120
GTHGWIARH	DIFLSTDGKT	WGSVPVATGTW	YADGTEKFSN	PETKRARYVR	IVAITEAYDG	180
PWTSIAEFNV	YKAATYTAPK	TGIGMWGPTL	DFPVPVAGA	VDPGTGKVLV	WSSYYHDTMN	240
GSPGGMTLTS	LWDPEGTGIT	QREVFETNHD	MFCPGISMDG	TGQIVVTGGN	NAARTSTWDP	300
VKNQWVSAPD	MKI PRGYQSS	ATTNGKVFT	IGGSWGGDRA	FKPGEIFDPK	SRKWTLLPNA	360
KVEPMLTADA	QGIIFRSDNHA	WLFGWKGETV	FQAGPSSAMN	WYYTQGGKTV	KPAGKRQSSR	420
GVDPDSMCGN	AVMFDAAKGK	IVTFGGTPNY	QDSYATNAH	IITIGAGPTQ	ASVAFVSDGM	480
YYPRVFHTSV	LLPDGTVFIT	GQGEYAIPEP	DSTPQLTPEL	YIPDSDTFIK	QQPNISIVRTY	540
HSMSILLQDA	RVFNGGGGLC	GDCSTNHFDA	QIFTPSYLLT	KDGKPAARPK	IVSVSATTIK	600
VGGSITVTTG	GVVNTASLIR	YGTATHTVNT	DQRRIPLTLA	SAGKNKYTFK	VPGDSGIALP	660
GYWMLFVMNS	AGVPSVATTI	KVTT				684

SEQ ID NO: 27 moltype = AA length = 506
 FEATURE Location/Qualifiers
 source 1..506
 mol_type = protein
 organism = *Colletotrichum spinosum*

SEQUENCE: 27

MVTLCSLTRS	LPAALLALAA	LSDAQGVGQW	GPLVKFPIVP	VAVALIPESG	NMLVWSSGWP	60
NRWTNAGNGK	TFTSLYDVST	GKAGDAIVQN	TQHDMFCPGT	SLDVEGRIIV	TGGSSAAKTS	120
VLDFRKGESS	SWAPLSNMQI	SRGYQSSCTT	SEGKIFVIGG	SFSGAGTRNG	EIYDTASNKW	180
TKLAGCPVKP	LVMQRGLFQD	SHTWLWSWKN	GSVLQAGPAK	QMNWYDTKGT	GANTPAGLRG	240
ADDDAMCGVS	VMFDAVAGKI	FTYGGGKAYT	GYAASSNAHI	LTLGEPGQVQ	QVQKLANGQY	300
NRGFSNAVVL	PDGRIWVVG	MRQMQLFSDN	TPQLTPELFD	PATGKFTPTA	PHAI PRNYHS	360
TALLMADATV	WSGGGLCGA	NCKENKFDGQ	FWSPPYLFEA	DGKTPAQRPV	IDSLSDKTVR	420
AGAPLVVTMK	DEGKYTFMSL	RVSATHTVNT	TDQRRIPLDG	QDGGDGKSF	VNMPGDYGVV	480
IPGYMMFAM	NEAGVPCVAK	PFKVAL				506

SEQ ID NO: 28 moltype = AA length = 684
 FEATURE Location/Qualifiers
 source 1..684
 mol_type = protein
 organism = *Pochonia chlamydosporia 123*

SEQUENCE: 28

MELFGAIAFV	LCQLSCHVHA	VTLKQTSTTA	L VHATEQAAL	HLLAAAPVGY	RIDRANWKVT	60
CDSQEIGYEC	GKAIDGDNST	VWHTAFRADN	PKPPHNI TVD	MGSIQNVNGL	SILPRQSDSDS	120
NGVWARHQLV	VSADGQRWEH	PVATGTWYAD	STEEKFSNFEP	EQARFVRLVA	DSEVSGNPWT	180
SVAEINLYKA	GTDPAAPAKSA	SSGKWGPTLD	FPIVPVAAA	VPQSGKVLVW	SAYENDKFEG	240
SPGGYTLTST	WDPATGDVTV	RNVTNIGHDM	FCPGISMSDN	GQIVVTGGNN	AQKTSFYDAA	300
SDSWVPDPDM	TVSRGYQSSA	TCSDGRIFTI	GGSWGGQFPE	KNGEYDPST	KTKMMLSLAA	360
VKPLMLTADK	GRYRADNHGW	LFGWRNGTVF	QAGPSTAMNW	YFTGGDGGVK	SAGSRKSSRG	420
PDPDSMCGNA	VMFDATNGKI	LTVGGSPWYQ	DNDATANAH	ITVRGPGSEP	SVSFAGNGMH	480

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HARIFASSVV	LPDGSVFITG	GQQHSIPFAD	STPQFTPELY	DPEKDQFLEQ	APNSIVRVYH	540
SLALLLPDAT	VFNNGGGLCG	GSCATNHFDA	QIYSPRYLFD	HDGSPAVRPV	IKSVSRTSVK	600
PGDLSLFTTD	SAVQHASLVR	YGTATHVTNT	DQRRVPLTLD	ASGANTYAVQ	LPNDPGIMLP	660
GYWMLFVMNE	RGTPSVAASI	KAHL				684

SEQ ID NO: 29 moltype = AA length = 684
 FEATURE Location/Qualifiers
 source 1..684
 mol_type = protein
 organism = Nectria haematococca

SEQUENCE: 29

MKNLLTSLC	LGSLLDVTDA	VVIPETPGQP	PGKFKASPL	GIPIDRKGWT	VTCSNQQSGN	60
DCAKAIDGNT	ATFWQTNANT	KAPVTITVDM	KTARNVNGLS	QLPLQDNSNN	IGNWIERHSV	120
FLSTDGKNWG	SPVATGTWWT	DKTEKFSNFE	PQRARYVRLV	AIPDETHDNP	RISLAELNIY	180
AASSNSNQEP	GLGRWGPTLD	FPVVPVASAV	EPTSGQVVVV	SARAYFEPAY	RYDDFQHQNP	240
RGGLTLSAVV	DPATGVITQR	NITLTHHDMF	CPGISMDGDG	QIVVTGGNDA	KKTSLYDSDS	300
DSWITGPEMN	IARGYQSSAT	TSDGRVFTIG	GSWNGPRGGK	NGEYINPDLK	SWTLLPGALV	360
KPMLTADKEG	VYRSDNHGWL	FGWKKGTVFQ	AGPSTAMNWX	YTRGADGDVK	PAGKRQVQDI	420
VDPDSMCGNS	VMYDAVKGKI	LTFGGSPNYR	FSDSTANAHI	ITIGEPGSA	KTAFAGGGQG	480
LHPRIFHTSV	VLPDGTVFIT	GGQKHSEPFV	DSTPQLEPEM	YLPASDAFVK	QQSNSIVRVY	540
HSISLLLPDG	RVFNNGGGGLC	GTCCTNHFDA	QIFTPNYLFD	KGNLNLTRPR	ISSTSTKTKA	600
VGSTITPTTN	GPVKQGSLLR	YGTATHVTNT	DQRRIALTPT	NTGTNRYSEK	IPNDPGIALP	660
GYWMLFVLNS	AGVPSVATTI	KVTN				684

SEQ ID NO: 30 moltype = AA length = 679
 FEATURE Location/Qualifiers
 source 1..679
 mol_type = protein
 organism = Gibberella subglutinans

SEQUENCE: 30

MKSFYSLALC	LGAFNAATA	IDPEEQGQP	GKFAAAPVVG	SNPIDRKGWT	VKCSSQAPNF	60
PCGRAIDGDK	NTFWQTPYGT	TNTPPPHTIT	IDMKQTQYVS	GLQITPRQDG	NTRNWIQRHE	120
VYLSDDGTNW	GKPVAFGTYW	GDKYPWITNF	ETHPARYLRF	VALSNVNSDY	PWIAIADFQV	180
YNALKYNPPA	KGLGKGGPTI	DFPVIPVAGA	VEPVSGKVVV	WSAYRYDAFQ	GTPRGGFTL	240
TSIWDPKTNV	ISNRNVSNNH	HDMFCPGISM	DGEGQIVVTG	GNDAKKTTIL	MPDGNWVWGP	300
DMQIARGYQS	SATCSDGRVF	TIGGSWSGAR	GKNGEIDYP	RAKTWTSLPK	CLVGPMLTHD	360
KEGVYKADNH	AWLFGWKKGS	VFQAGPSTAM	NWYYTDRGTQ	GNTKAAGRTR	KNGRVDPSM	420
NGNVAMPDAV	KGKILTFGGA	TSYQQAPATA	NAHVLTIDQP	GAIQATLVG	NNGAGIHARV	480
FATSTILPDG	NVFITGGQSY	SNPPTDINAQ	LEPEMFISSE	NTFTKQQPNT	IPRTYHMSML	540
LLPDGTVFNG	GGGLCGSCKS	NHFDAQIFTP	QYLLDGNGNL	ATRPKITAVS	ATTAKVGSTI	600
TVTANSIAKS	ASLMRYGTAT	HVVNTDQRR	PLALTGAGTN	KYSFKIPNDS	GIALPGYWML	660
FVINNAGVPS	VASTIKVTV					679

SEQ ID NO: 31 moltype = AA length = 681
 FEATURE Location/Qualifiers
 source 1..681
 mol_type = protein
 organism = Gibberella subglutinans

SEQUENCE: 31

MKSFWTLAFY	LGANAVAIS	QPASKAETPE	GSLQFLSLRA	SAPIGTAINR	DKWKVTCDSQ	60
HQDECSKAI	DGDRNTFWHT	NWAAGATNDP	KPPHTITIDM	GSSQNVNGLS	VLPDQDSDH	120
GWIGRHNVFL	STDGKNWNGA	VATGTWFADI	TEKYSNFETR	PARYVRLVAV	TEANDQPWTS	180
IAEINVFNA	SYTSPQPLGL	LWGPDLDFPI	VPVAAVEPT	SGKVLVWSSY	KNDAFGGSPG	240
GVTLTSTWDP	STGVISQRTV	TVTCKHDMFCP	GISMDCGNGQV	VVTGGNNAEK	TSLYDSSSDS	300
WIPGDMKVA	RGYQSSATLS	NGRVFTIGGS	WSGGTFEKNK	EVYDPSSKTW	TSLSGALVKP	360
MLTADQQGIY	RSDNHGWLFG	WKKGSVFPQAG	PSTAMNYYT	SGKGNTKSAG	KRQSSRGTD	420
DAMCGNAVY	DAVKGKILTF	GGSPSYQDSD	ATTNAHIITI	GEPGSTPKTV	FASNGLYYPR	480
TFHTSVLIPD	GNVFTGGQQ	RGIPFADSTP	QLTPELYVFN	DDTFYKQQPN	SIVRVYHSIS	540
LLLPDGRVFN	GGGGLCGDCS	TNHFDAQIYT	PNNLYDSNGK	LATRPKITNV	SAKSAKVGK	600
ITISTDTSIK	QASLIRYGTG	THTVNTDQRR	IPLSLRSTGS	GNSYSFQVPS	DSGIALPGYW	660
MLFVMNSAGV	PSIASTLLVT	Q				681

SEQ ID NO: 32 moltype = AA length = 681
 FEATURE Location/Qualifiers
 source 1..681
 mol_type = protein
 organism = Gibberella moniliformis

SEQUENCE: 32

MKSFWTLAFY	LGGASAVAIS	QPAKSETPA	GSLQFLSLRA	SAPLGTAINR	DKWKVTCDSQ	60
HEGDECSKAI	DGDRNTFWHT	NWAAGTNDP	KPPHTITIDM	GSSQNVNGLS	VLPDQDGSNH	120
GWIGRHNVFL	STNGKNWNGA	VATGTWFADN	TEKYSNFETR	PARYVRLVAV	TEANDQAWTS	180
IAEINVFKAA	SYTSPQPLGL	RWGPDLDFPI	VPVAAVEPT	SGKVLVWSSY	KNDEFGGSPG	240
GVTLTSTWDP	STGVISQRTV	TNTCKHDMFCP	GISMDCGNGQV	VVTGGNNAEK	TSLYDSSSDS	300
WIPGDMKVA	RGYQSSATLS	NGRVFTIGGS	WSGGISEKNK	EVYDPSSKTW	TSLSGALVKP	360
MLTADQQGLY	RSDNHGWLFG	WKKGSVFPQAG	PSTAMNYYT	SGKGNTKSAG	KRQSSRGTD	420
DAMCGNAVY	DAVKGKILTF	GGSPSYQDSD	ATTNAHIITI	GEPGSTPKTV	FASNGLYYPR	480

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TFHTSVILPD	GNVFITGGQQ	RGIPFADSTP	QLTPELYVFN	DDTFYKQOPN	SIVRVYHSVS	540
LLLPDGRVFN	GGGLCGGCT	TNHFDAQIYT	PNNLYDSNGK	LATRPKITKV	SAKSVKVGK	600
ITISTDSSIK	QASLIRYGT	THTVNTDQRR	IPLSLRSTGS	GNSYSFQVPS	DSGIALPGYW	660
MLFVMNSAGV	PSVASTLLVT	Q				681

SEQ ID NO: 33 moltype = AA length = 680
 FEATURE Location/Qualifiers
 source 1..680
 mol_type = protein
 organism = *Fusarium coffeatum*

SEQUENCE: 33

MRHLLTLALC	FSSINAVAIT	KSHKAAGTGE	PEGSFQFLSL	RASAPIGTTI	SRDKWSVTCD	60
SFHDEGEGCDK	AIDGDLNFTW	HSQWNGNDP	KPPHTYTIDM	GSTQNVNGLS	VLPRQDGSSN	120
GWIGRHEVFL	SSDGKNWWSA	VATGTWYADS	TTKYSNFETR	PARVYRLVAL	TEAKGQPWTS	180
IAEINVFKAD	SYSAPRAGLG	RWGPTLDFPI	VPAAAFVEPT	SGKVVVFSSY	RNDAFGGSPG	240
GITLTSSWDP	ANNVIAERTV	TVTTHDMFCP	GISMDGAGQI	VVTGGNDACK	TSLYDSSSDS	300
WIPGPDQNA	RGYQSSATTS	DGRVFTIGGS	WSGGIFEKNG	EIYNPSSKTW	TSLSGAKVNP	360
MLTADRQGLY	RSDNHAWLFG	WKKGIVFQAG	PSTAMNYYT	SGNGDVKSAG	KRQSNRGVAT	420
DAMCGNAVMY	DAVQKILTF	GGAKDYQDTD	ATDAHIITL	GEPGTPKTV	YASNGLWYPR	480
TFHTSVVLPD	GSTFITGGQV	RGIPFEDSTP	QLTPELYVFP	DDTFYKQOPN	SIVRVYHSVS	540
LLLPDGRVFN	GGGLCGDCT	TNHFDAQVFT	PNLYDNNGN	LVTRPKITNT	STKSVKIGGR	600
VTITTDGSIQ	KASLVRYGTA	THTVNTDQRR	IPLTSLNSGR	NSYSFTVPND	SGVALPGYWM	660
LFVMSAGVP	SVSTTIRITS					680

SEQ ID NO: 34 moltype = AA length = 680
 FEATURE Location/Qualifiers
 source 1..680
 mol_type = protein
 organism = *Fusarium sporotrichioides*

SEQUENCE: 34

MKHLTLALC	FSSINAVAIT	NPHKTAGHDH	PEGSLQFLSL	RASAPIGSAI	SRNNWAVTCD	60
SAQSGNECNK	AIDGNQDFTW	HTFYGANGDP	KPPHTYTIDM	KSTQNVNGLS	MLPRQDGSRN	120
GWIGRHEVYL	STDGTMWGSF	VAAGSWFADS	TTKYSNFETR	PARVYRLVAV	TEASGQPWTS	180
IAEINVQAS	SYTAPQPLG	RWGPTIDLPI	VPAAAIVEPT	SGRVLVWSSY	RNDAFGGSPG	240
GVTLTSSWDP	SSGIVSDRTV	TVTTHDMFCP	GISMDGNGQI	VVTGGNDACK	TSLYDSSSDS	300
WIPGPDQVA	RGYQSSATMS	DGRVFTIGGS	WSGGIFEKNG	EVYSPSSKTW	TSLPNAKVN	360
MLTADKQGVY	RSDNHAWLFG	WKKGSVVFQAG	PSTAMNYYT	SGSGDVKSAG	KRQSNRGVAP	420
DAMCGNAVMY	DAVRGKILTF	GGSPDYQDSD	ATANAHIITL	GEPGSTPNTV	FASNGLYFAR	480
TFHTSVVLPD	GSTFITGGQR	RGIPFEDSTP	VFTPEVYVPE	QDTFYKQNP	SIVRVYHSIS	540
LLLPDGRVFN	GGGLCGDCT	TNHFDAQIFT	PNLYDNGN	LATRPKITRT	STQSVKVGGR	600
VTISTDSSIV	KASLIRYGTA	THTVNTDQRR	IPLTLTNNNG	NSYSFQVPSD	SGIALPGYWM	660
LFVMSAGVP	SVAATIRVTQ					680

SEQ ID NO: 35 moltype = AA length = 682
 FEATURE Location/Qualifiers
 source 1..682
 mol_type = protein
 organism = *Fusarium longipes*

SEQUENCE: 35

MKQLTLALC	FSSINAVAIN	PHGHNNKGT	GDHEGSLQFL	SLRASAPLGS	AIARENWVVT	60
CDSAQPNQNE	NKAIDGDNNT	FWHTFYGNNG	DPKPPHTYTI	DMGNNRNVNG	LSVLPQDGN	120
RNGWIGRHEV	YLDGDSNWG	SPVAYGSWFA	DSTTKYSNFE	TRPARYVRLV	ALTEASGQPW	180
TSIAEINVYQ	AGSYTAPQAG	LGRWGPTIDL	PIVPAAAAIE	PTSGRVLWMS	SYRNDAFGGS	240
PGGITLTSSW	DPSSGIVSDR	TVTVTNHDMF	CPGISMDGNG	QIVVTGGNDA	KKTSLYDSSS	300
DSWIPGPDQ	VARGYQSSAT	MSDGRVFTIG	GSWSGGIFEK	NGEVYSPSSK	TWTSPLGAKV	360
NPMLTADKQG	LYRSDNHAWL	FGWKKGSVFP	AGPSTAMNYY	YTSGSDVKS	AGRRQSNRGL	420
DPDAMCGNAV	MYDAVAGKIL	FTGGSPDYQD	SDATTNAHII	TLGEPGSTPN	TVFASNGLYF	480
PRTFHTSVVL	PDGSTFITGG	QRRGIPFEDS	TPQLTPEIYV	PEQDTFYKQN	PNSIVRVYHS	540
ISLLLPDGRV	FNGGGGLCGD	CTTNHFDAQI	FTPNLYNSN	GDLATRPKIT	RTSAQSVRVG	600
GRITISTDSS	IRRASLIRYG	TATHTVNTDQ	RRIPLTLTNN	GGNSYSFQVP	SDSGIALPGY	660
WMLFVMNSAG	VPSVATTLRV	TR				682

SEQ ID NO: 36 moltype = AA length = 645
 FEATURE Location/Qualifiers
 source 1..645
 mol_type = protein
 organism = *Streptomyces lividans*

SEQUENCE: 36

MKDRAGRRA	RFAIGTAVV	VALAGMNGPW	LYRFSTEKYH	QYKINQPEYK	AANGKWEIIE	60
FPEYRQNTI	HAALLRTGKV	LMVAGSGNNQ	DNSDDKQYDT	RIWDVPKGTI	KKVPTPSDLF	120
CTGHTQLANG	NLLIAGGTR	YEKLGKDVTK	AGGLMVVHNE	NPDKPITLPA	GTKFTGKENG	180
KTFVSKDPVL	VPRAEKVFPD	ATGAFVRNDP	GLGRIYVEAQ	KSGSAYETGT	EDNYRVQGLS	240
GADARNYTYI	AQKLALDKKD	FQGIIRDAFEP	DPVAEKYIKV	DPMHARWYP	TLTTLGDGKI	300
LSVSGLDIDG	QLVPGKNEVY	DPKTKAWTYT	DKVRQFPPTY	ALFLMONGKI	FYSGANAGYG	360
PDDVGRTPGV	WDVETNKFTK	VPGMSDANML	ETANTVLLPP	AQDEKYMVIG	GGGVGESKLS	420
SEKTRIADLK	ADDPKFDVGP	SLEKGRYPQ	ASLPPDSSVL	VSGGSQDYRG	RGDSNLLQAR	480

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LYHPDTNEFE	RVADPLVGRN	YHSGSILLPD	GRLMFFGSDS	LYADKANTKP	GKFEQRIEY	540
TPPYLYRDSR	PDLGGPQTI	ARGGSGTFTS	RAASTVKKVR	LIRPSASTHV	TDVDQRSIAL	600
DFKADGDKLT	VTVPSGKNLV	QSGWYMMFVT	DGEGTPSKAE	WVRVP		645

SEQ ID NO: 37 moltype = AA length = 730
 FEATURE Location/Qualifiers
 source 1..730
 mol_type = protein
 organism = unidentified

SEQUENCE: 37

MRHFLLPAV	AGIAGAQC	PSY	LSGEMSFTQE	QDNAGDTIEV	TEQPIDNTLY	VNDTGSYMTT	60
DFGTPISDQT	SLKAGPRGPT		LLEDFIFRQK	LQRFDERHVP	ERVVHARGAG	AYGTFKSYAD	120
WSNVTAADFL	SANDKETPMF	CRFSTVVGFR	GSVDTARDVH	GHACRFYTDE	GNVDIVGINF		180
APFFIQDAIQ	FPDLVHAIKP	MPNNEIPQAA	TAHTSAWDFE	SQQSTALHSA	LWLMSGNGIP		240
RSFRHMNGYG	VHSFRFVAAN	GTSKVVRTPW	KSQQGVASLV	WDEAQAAGK	NSDYHRQDLY		300
NAMPNGHYPK	YELQAQIMDE	ADMLRFGFDL	LDPTKLVPEE	VVPYTPLGMM	ELNANPTNYF		360
AEVEQAGFQP	GHVVPGIDFT	DDPLLQGRLE	SYLDTQLTRH	GGPNFEQIPV	NRPRKPVHNN		420
NRDGFQGOQI	PTNNWAYTPN	SMSNGYPMQA	NQTQGHGFPT	APYRYASGHL	VRQTSPTFND		480
HWSQPAMFVN	SLIPAEQQMV	VNAIVFENSK	VNSPHVRKIV	VNQLMMVMNN	LAVRVARGLG		540
LDEPSPNPTY	YTSNKTSNVG	TFGKPLLSIE	GLQVGFPLASN	SHPESIKQGG	AMAAQFSAAG		600
VDLNIIVTEAY	ADGVNTTYAL	SDAIDFDALI	IADGVQSLFA	SPALANQMNS	TATSTLYPPA		660
RPFQILVDSF	RYGKPVAAVG	SGSVALKNAG	IDSSRSGVYT	GSSETTEKIA	KEVLEGLYTF		720
RFVDRFALDE							730

SEQ ID NO: 38 moltype = AA length = 730
 FEATURE Location/Qualifiers
 source 1..730
 mol_type = protein
 organism = unidentified

SEQUENCE: 38

MRHFLLPAV	AGIAGAQC	PSY	LSGEMSFTQE	QDNAGDTIEV	TEQPIDNTLY	VNDTGSYMTT	60
DFGTPISDQT	SLKAGPRGPT		LLEDFIFRQK	LQRFDERHVP	ERVVHARGAG	AYGTFKSYAD	120
WSNVTAADFL	SANDKETPMF	CRFSTVVGFR	GSVDTARDVH	GHACRFYTDE	GNVDIVGINF		180
APFFIQDAIQ	FPDLVHAIKP	MPNNEIPQAA	TAHTSAWDFE	SQQSTALHSA	LWLMSGNGIP		240
RSFRHMNGYG	VHSFRFVAAN	GTSKVVRTPW	KSQQGVASLV	WDEAQAAGK	NSDYHRQDLY		300
NAMPNGHYPK	YELQAQIMDE	ADMLRFGFDL	LDPTKLVPEE	VVPYTPLGMM	ELNANPTNYF		360
AEVEQAGFQP	GHVVPGIDFT	DDPLLQGRLE	SYLDTQLTRH	GGPNFEQIPV	NRPRKPVHNN		420
NRDGFQGOQI	PTNNWAYTPN	SMSNGYPMQA	NQTQGHGFPT	APYRYASGHL	VRQTSPTFND		480
HWSQPAMFVN	SLIPAEQQMV	VNAIVFENSK	VNSPHVRKIV	VNQLMMVMNN	LAVRVARGLG		540
LDEPSPNPTY	YTSNKTSNVG	TFGKPLLSIE	GLQVGFPLASN	SHPESIKQGG	AMAAQFSAAG		600
VDLNIIVTEAY	ADGVNTTYAL	SDAIDFDALI	IADGVQSLFA	SPALANQMNS	TATSTLYPPA		660
RPFQILVDSF	RYGKPVAAVG	SGSVALKNAG	IDSSRSGVYT	GSSETTEKIA	KEVLEGLYTF		720
RFVDRFALDE							730

SEQ ID NO: 39 moltype = AA length = 503
 FEATURE Location/Qualifiers
 source 1..503
 mol_type = protein
 organism = *Micrococcus luteus*

SEQUENCE: 39

MEHQKTPHA	TGSTRQNGAP	AVSDRQSLTV	GSEGPVLVHD	THLLETHQHF	NRMNIPERRP		60
HAKGSGAFGE	FEVTEVSKY	TKALVFQPGT	KTETLLRFST	VAGELGSPDT	WRDVRGFALR		120
FYTEEGNYDL	VGNNTPIFFL	RDPMKFTHFI	RSQKRLPDSG	LRDATMQWDF	WTNNPESAHQ		180
VTYLMLGPRGL	PRTWREMNGY	GSHTYLWVNA	QGEKHWVKYH	FISQQGVHNL	SNDEATKIAG		240
ENADFHRQDL	FESIAGGDHP	KWDLYIQAI	YEEGKTYRFN	PFDLTKTISQ	KDYPRIKVGT		300
LTLNRNPNENH	FAQIESAAFS	PSNTRVPGIGL	SPDRMLLGRA	PAYHDAQLYR	VGAAHVQQLPV		360
NRPKNAVHNY	AFEGQMWDYH	TGDRSTYVFN	SNQDSWSDET	GPVDDGWVAD	GTLTREAQAL		420
RADDDDFGQA	GTLVREVFS	QERDDFVETV	AGALKGVRQD	VQARAFYWK	NVDATIGQRI		480
EDEVKRHEGD	GIPGVEAGGE	ARM					503

SEQ ID NO: 40 moltype = AA length = 527
 FEATURE Location/Qualifiers
 source 1..527
 mol_type = protein
 organism = *Bos taurus*

SEQUENCE: 40

MADNRDPASD	QMKHWKEQRA	AQKPDVLTG	GGNPVGDKLN	SLTVGPRGPL	LVQDVVFTDE		60
MAHFDRERIP	ERVVHAKGAG	AFGYFEVTHD	ITRYSKAKVF	EHIGKRTPIA	VRFSTVAGES		120
GSADTVRDP	GFAVKFTED	GNWDLVGNNT	PIFFIRDALL	FPSPHISQKR	NPQTHLKD		180
MVWDFWLSRP	ESLHVQVSLF	SDRGI	PDGHR	HMMNGYGSHTF	KLNVNANGEAV	YCKFHYKTDQ	240
GIKNLSVEDA	ARLAHEDPDY	GLRDLFNAIA	TGNYP	SWTLY	IQVMTFS	EAE	300
KVWPHGDYPL	IPVQKLVLRN	NPVNYFAEVE	QLAFDPSNMP	PGIEPSPDKM	LQGRLEFAYPD		360
THRHRLGPNY	LQIPVNCPYR	ARVANYQRDG	PMCMMDNQQG	APNYYPNSFS	APRHQPSALE		420
HRTHFSGDVQ	RFSANDDDN	TQVRTFYLVK	LNEEQKRLC	ENIAGHLKDA	QLFIQKAVK		480
NFSDVHPEYG	SRIQALLDKY	NEEKPKIAVH	TYVQHGSHLS	AREKANL			527

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SEQUENCE: 46
 ATPLTSLGSE QAMFHGKHQP GITTTPMQARG HLVAFDLAAG AGRKEAALL RRWSDTARRL 60
 MAGEPAGSRD TDVARDAGPS SLTVTFGFGH SFFGRGTGLEK QRPVALDPLP DFSSDHLKDN 120
 RSNGLWVQI GADDALVAFH ALRAIQRDAG AAARVRWQMN GFNRSPGATA HPMTARNLMG 180
 QVDGTRNPKP GEADFRRIF VPEEPEAGKG GPAMWANGSY VVRRIRMLL DWHEELSKA 240
 QEDVIGRRKS DGAPLGGSG ATESTEMDLE KTDGSGELVV PINAHARITR PDQNGGAAMV 300
 RRPFSYHDGF DADGVPDAGL LFVCWQADPL RGFVPVQRKL DRGDALSQFI RHEASGLFAV 360
 PGGAAEGEYV GQRLLE 376

SEQ ID NO: 47 moltype = AA length = 712
 FEATURE Location/Qualifiers
 source 1..712
 mol_type = protein
 organism = Bos taurus

SEQUENCE: 47
 MWVCLQLPVF LASVTLFEVA ASDTIAQAAS TTTISDAVSK VKIQVNKAFL DSRTLKTTL 60
 SSEAPTQQL SEYFKHAKGR TRTAIRNGQV WEESLKRRLR DTTLTNVTDL SLDLTALSW 120
 VCGGAPVPLV KCDENSPYRT ITGDCNNRRS PALGAANRAL ARWLPAEYED GLALPFGWTQ 180
 RKTRNGFRVP LAREVSNKIV GYLDEEGVLD QNRSLLFMQW GQIVDHDLDL APETELGSNE 240
 HSKTQCEEYC IQGDNCFPII FPKNDPKLKT QGKCMPPFRA GFVCPPTPYQ SLAREQINAV 300
 TSPLDASLVY GSEPSLASRL RNLSSPLGLM AVNQEAWDHG LAYLFPNNKK PSPCEPINT 360
 ARVPCFLAGD FRASEQILLA TAHTLLREH NRLARELKKL NPHWNGEKLY QEARKILGAF 420
 IQIITFRDYL PIVLGSEMOK WIPPYQGYNN SVDPRISNVF TFAFRFGHME VPSTVSRLE 480
 NYQPWGPEAE LPLHLTFEFT WRIKIDGGID PLVRGLLAKK SKLMNQDKMV TSELRNKLFQ 540
 PTHKIHFIDL AAINLQRCRD HGMPGYNSWR GFCGLSQPKT LKGLQTVLKN KILAKKMDL 600
 YKTPDNLDIW IGGNAEPMVE RGRVGPLLAC LLGRQFQQIR DGDRFVWENP GVFTKQRDS 660
 LQKVSFSRLI CDNTHITKVP LHAFAQANNYP HDFVDCSTVD KLDLSPWASR EN 712

SEQ ID NO: 48 moltype = AA length = 372
 FEATURE Location/Qualifiers
 source 1..372
 mol_type = protein
 organism = Leptoxiphium fumago

SEQUENCE: 48
 MFSKVLFPVAV AVALPHSVR QEPGSGIGYP YDNNTLPYVA PGPTDSRAPC PALNALANHG 60
 YIPHDGRAIS RETLQNAFLN HMGIANSVIE LALTNAFVVC EYVTGSDCGD SLVNLTLLE 120
 PHAFEHDSF SRKDYKQVSA NSNDFIDNRN FDAETPQTSI DVVAGKTHFD YADMNEIRLQ 180
 RESLSNELDF PGWFTESKPI QNVESGFIFA LVSDFNLPDN DENPLVRIDW WKYWFNTSEF 240
 PYHLGWHPPS PAREIEFVTS ASSAVLAASV TSTPSSLPSG AIGPGAEAVP LSFASMTMPF 300
 LLATNAPYYA QDPTLGPNDK REAAPAATTS MAVFKNPYLE AIGTQDIKNQ QAYVSSKAAA 360
 MASAMAANKA RN 372

SEQ ID NO: 49 moltype = AA length = 378
 FEATURE Location/Qualifiers
 source 1..378
 mol_type = protein
 organism = Phanerodentia chrysosporium

SEQUENCE: 49
 MAPKSLIAFV ALAAAVRAAP TAVCPDGTTRV SHAACCAFIP LAQDLQETIF QNECGEDAHE 60
 VIRLTFHDAI AISRQGPKA GGGADGSMML FPTVEPNFSA NNGIDDSVNN LIPFMQKHNT 120
 ISAADLVQFA GAVALSNCPI APRLEFLAGR PNKTIAAVDG LIPEPQDSVT KILQRFEDAG 180
 GFTPFVSVL LASHSVARAD KVDQTIDAAP FDSTPFTFDT QVFLEVLLKG VGFPGSANNT 240
 GEVASPLPLG SGSDTGEMRL QSDFALAHDP RTACIWQGFV NEQAFMAASF RAAMSKLAVL 300
 GHNRNSLIDC SDVVPVPKPA TGPQAMFPAS TGPQDLELSC PSERFPPLTT QPGASQSLIA 360
 HCPDGSMSCP GVQFNGPA 378

SEQ ID NO: 50 moltype = AA length = 430
 FEATURE Location/Qualifiers
 source 1..430
 mol_type = protein
 organism = Thermobifida fusca DyP

SEQUENCE: 50
 MTEPDTERKG SSRRGFLAGL GAAALTGAGI GMAAGEVLRP LLPDSDPAAS PKAEQRLRMA 60
 AQRADATAAP QPGISGPAPA FVHVIALDLA EEARKNPDTA RDSAAAALRS WTELAARLHE 120
 ESPHDIAGA ASAGLLPASL MVTVGIGGSL LSAIDAEDRR PDALADLPEF STDDLHPRWC 180
 GGDFMLQVGA EDPMLTAAV EELVAAAADA TAVRWSLRGF RRTAAARDP DATPRNLMGQ 240
 IDGTANPAQD HPLFDRITTA RPADNPAHAW MDGGSYLVRV RIRMLLTWR KLDVAARERV 300
 IGRRLDTGAP LGSRNEDTPV VLSARDEEGE PLIPENAHVR LASPENLGA RMFRRGYSYD 360
 QGWRDDGVRD AGLLFMAWQG DPATGFIPVQ RSLADQGDAL NRYIRHEGSA LFAVPAAREG 420
 RYLGQDLIEG 430

SEQ ID NO: 51 moltype = AA length = 403
 FEATURE Location/Qualifiers
 source 1..403
 mol_type = protein
 organism = Saccharomonospora viridis

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SEQUENCE: 51
 MKGRRFRGRR FLATGAAALG GAAALGGREL FTGDGGADGR DRPDRGDIGR ATVDFHGERQ 60
 AGVATPAQAF ATFVAFDLDL GVDREALIRW MRVWTDIER LTRGAPALTD TEPELALLPA 120
 RLTVTVGFGP GFLAAAGREE LRPSWLAPLP EFPIDRLREE FSGGDLVAQV CADDEVTVAH 180
 AVRVLTKQAR SFARPRWVQR GFRNTPGAVP EGATMRNLMG QLDGTRNLRP GPDDRLIWIW 240
 DGPEWLRGGT GMVVRRIAMN LDTWDELDRP ARELVIGRRL DNGAPLTGRH EHDEPDLEAV 300
 DERGLSVIPM FAHIRRARSQ NPDERFLRRS YNYDDPPEPG ELSNSGLVVFV TFQADIEAQF 360
 TPIQRRLAEL DSLNDWTTPI GSAVFAVPRG CRPGEYLGQP LLE 403

SEQ ID NO: 52 moltype = AA length = 299
 FEATURE Location/Qualifiers
 source 1..299
 mol_type = protein
 organism = Escherichia coli

SEQUENCE: 52
 MSQVQSGILP EHCRAAIWIE ANVKGEVDAL RAASKTFADK LATPEAKFPD AHLGAVVAFG 60
 NNTWRALSGG VGAEELKDFP GYGKGLAPTT QFDVLIHILS LRHDVNFVA QAAMEAFGDC 120
 IEVKKEIHGF RWVEERDLGS FVDGTENPAG EETRREVAVI KGDVDAGGSY VVQVWEHNL 180
 KQLNRMSVHD QEMMIGRTKE ANEEDIGDER PETSLLTRVD LKEDGKGLKI VRQSLPYGTA 240
 SGTGHLFYCA YCARLHNIEQ QLLSMFGDTD GKRDAMLRFT KPVTGGYYFA PSLDKLMAL 299

SEQ ID NO: 53 moltype = AA length = 343
 FEATURE Location/Qualifiers
 source 1..343
 mol_type = protein
 organism = Cellulomonas bogoriensis

SEQUENCE: 53
 MTPPPSFIAL VALDLASTD RASVERLLRV WTVDIERLTT GRPGLADSEP ELALVPAALT 60
 VTVGFGPGLL TAAGLRHRAP AWHLPLPPFP IDRLDPAWCD GDVVLQVCAD DRTTLAHAVR 120
 VLTKEAQLA SVRWVQRGFR RSPGISEPDG TSMRNLMGQV EGTANLDPRT DPDLLWHRDG 180
 EPGWLTGGTS MVVRRIAMNL DTWDELSRGA REATIGRTLRL TGAPLTGRAE HDEPDLEALD 240
 DHGRPVIDLE AHIRRAPRTQ REETFLRRAY NYDEAPPPGR ASDSGLLFVT YQRDVDAQFT 300
 PVQRRLDAAD LLNEWTFFVG SAVFAVPGGW SAGEYVGQRL LEG 343

SEQ ID NO: 54 moltype = AA length = 427
 FEATURE Location/Qualifiers
 source 1..427
 mol_type = protein
 organism = Thermobifida cellulositytica

SEQUENCE: 54
 MTGPEPTPAG SSRRGFLAGL GAAALTGAGL GMAAGEAVRP LLPDSSEQP DPVDAQRLDM 60
 ARRADANAA POPGISGRAP AFVHVIAFDL AEPARAEPAA AREGAATALR TWAEHAARLH 120
 ADGPEGAAASA GLLPASLMVT IGIGGSLEEA MDAADRRPDA LADLPEFATD DLRPRWCGGD 180
 LMLQVGAEDP MVLAAAVDEL VAATAPTTV RWSLRGFRRT AAAAQDPDAT PRNLMGQIDG 240
 TANPAQDHPL FTRTVTAPPA DDPAHAMMDG GSYLVVRRIR MLLDEWRRLD VPDREVRIGR 300
 HLDTGAPLGG EKETDPVLT ARDADGRLVI PEDAHVRLAN PENNLGARMV RRGYNYDEGW 360
 RDDGVRDAGL LFMAWQGNPA TGFVPVQRSL VEQGDALNRY TRHEGSALFA VPAATADRYP 420
 QDLEVEG 427

SEQ ID NO: 55 moltype = AA length = 316
 FEATURE Location/Qualifiers
 source 1..316
 mol_type = protein
 organism = Streptomyces coelicolor

SEQUENCE: 55
 MGGEVEPEP QMVLSPILSA AIFLVVTIDS GGEDTVRDLL SDVASLERAV GFRAQPDGRL 60
 SCVTGIGSEA WRDLFSGARP AGLHPPFREL GPVHRAVATP GDLLFHIRAS RLDLCFALAT 120
 EIMGRLRGAV TPQDEVHGFK YFDERDMLGF VDGTEPTGA AARRAVLVGA EDPAFAGGSY 180
 AVVQKYLHDI DAWEGLSVEA QERVIGRRKM TDVELSDDVK PADSHVALTS VTGPDGSDLE 240
 ILRDNMPFGS VGREEFGTYF IGYARTPEVT ETMLERMFLG TASAPHDRIL DFTAVTGS 300
 FFTPAADFLE DLPARP 316

SEQ ID NO: 56 moltype = AA length = 138
 FEATURE Location/Qualifiers
 source 1..138
 mol_type = protein
 organism = Amphitrite ornata

SEQUENCE: 56
 MGFKQDIATI RGDRLTYAQD IFLAFLNKYP DERRYFKNYV GKSDQELKSM AKFGDHTEKV 60
 FNLMEVADR ATDCVPLASD ANTLVQMKQH SSLTTGNFEK LFVALVEYMR ASGQSFDSQS 120
 WDRFGKNLVS ALSSAGMK 138

SEQ ID NO: 57 moltype = AA length = 261
 FEATURE Location/Qualifiers
 source 1..261
 mol_type = protein

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organism = Thermothelomyces thermophilus
SEQUENCE: 57
MRASVLPVLI AISPALAGFD TWSPPGPPYDV RAPCPMLNLT ANHGFLPHDG KDITREQTEN 60
ALFEALHINK TLASFLDFDA LTTNPKNTST FSLNDLGNHN ILEHDASLSR ADAYFGNVLQ 120
FNQTVFDETK TYWEGTIDL RMAAKARLGR IKTSQATNPT YSMSSELGDAF TYGESAAVVV 180
VLGDKESRTV KRSVWEWFFE HEQLPQHLGW KRPAASFEEE DLNSSMBEIE KYTKELEGSN 240
STSGSQKHRR RLPFRRRAHFG F 261

SEQ ID NO: 58      moltype = AA length = 504
FEATURE          Location/Qualifiers
source          1..504
                mol_type = protein
                organism = PLEUROTUS SAPIDUS

SEQUENCE: 58
MTTPAPPLDL NNIQGDILGG LPKKTETYFF FDVTNVDRFK ANMTQFIPHV KTSAGIVKDR 60
EAIKEHRQK RPGLVPMMAAV NVSPSHLGLQ KLGITDDLSD SSFTTGQRKD AEVLGDPGPK 120
NGDTFTPAWE APFLKDIHGV IFVAGDCHAS VHKKLDEIKH IFVGTSHAS ISEVTHVRGD 180
VRPGQVSAHE HFGFLDGISN PAVDQFDQNP FPGQDSIRPG FILAKENGDS RAAARPDWAK 240
DGSFLTRFYL FQMVPEFDDF LESNPVILPG LSRKEGSELL GARIVGRWKS GAPIEITPLK 300
DDPKLGADAQ RNMNFDFGDS LVRGDQTKCP FAAHIRKTYP RNDLEGPPLN ADIDNRRIIR 360
RGIQFGPEVT SQEHDKKTH HGRGLLFVCY SSSIDDPHFH IQQSWANAPN FPNNAVTSAG 420
PIPLDGVIP GFDIIGQKV GGGIRQISGT NPNDPTTNTIT LPDQDFVIRP GGEYFFSPSI 480
SALKTKFAAG VASSAPQAQA PIST 504

SEQ ID NO: 59      moltype = AA length = 480
FEATURE          Location/Qualifiers
source          1..480
                mol_type = protein
                organism = Irpex lacteus

SEQUENCE: 59
MHVKRARSTP LIGSFPQPP LPTIAQVQST SAGNDSLPEE NIQGDILVGM KDKKEKVFVF 60
HINNATAFKS VLKTYAPANI TSVATIIGPV ANQPLAFVNL AFSHAGFGAL NVTDDLQDTA 120
FSDGQFKDSP NLGDDTSTWE EAFKGTNVGD VFLIGSNDES ITAQYRDDLN AKFGDAWTIV 180
YDLDSAARPG NEKGHEHFGY LDGINSPTIP GFGTTPHGQA VVDPGIIFTG RSKDPVMNRP 240
SWALDGSFLV FRKLLQLVPE FNKYVLDNAL QNQAGNLTV EGAELLSRM FGRWKS GAPI 300
DLSPDFDDPA LGNDIERNNN FNYSHPGSDL ATDQTRCPPT AHIRKTNPRD LEGQGLFGDT 360
PHAIRAGTPY GPEVTDYEAS SNTTIDRGL AFVEYQSVIG NGFRFQQQAW ANNPRFPFSK 420
GPSIQLGLDP VIGQGSPPRET FGLDPRNASE SFTVPQVIIS NGGEYFFSPS ITAIVEKFAA 480

SEQ ID NO: 60      moltype = AA length = 250
FEATURE          Location/Qualifiers
source          1..250
                mol_type = protein
                organism = Nicotiana tabacum

SEQUENCE: 60
MGKCYPTVSE EYLKAVDKCK RKLRLIAEK NCAPLMLRLA WWSAGTYDVC SKTGGPFGTM 60
RLKAEQGHGA NNGIDIAIRL LEPIKEQFPI LSYGDFYQLA GVVAVEVTGG PDVPPHGRE 120
DKTEPPVEGR LPDATKGSFH LRDVFKQMG LSDKDIALS GGHTLGRCHK ERSGFEGPWT 180
TNPLIFDNSY FTELLSGEKE GLLQLPSDKA LLSDPAPRPL VEKYAAEDA FFADYAEABL 240
KLSELGFAEA 250

SEQ ID NO: 61      moltype = AA length = 519
FEATURE          Location/Qualifiers
source          1..519
                mol_type = protein
                organism = Trametes versicolor

SEQUENCE: 61
MGLQRFSFFV TLALVARSLA AIGPVASLVV ANAPVSPDGF LRDAIVVNGV VPSPLITGKK 60
GDRFQLNVDD TLTNHSMLKS TSIHWHGFFQ AGTNWADGPA FVNQCPIASG HSFLYDFHVP 120
DQAGTFWYHS HLSTQYCDGL RGPVVVYDPK DPHASRYDVD NESTVITLTD WYHTAARLGP 180
RFP LGADATL INGLGRSAST PTAALAVINV QHGKRYRFR LVSISCDPNYT FSIDGHNLTV 240
IEVDGINSQP LLVDSIQIFA AQRYSFV LNA NQTVGNYWVR ANPNFGTVGF AGGINSAILR 300
YQGAPVAEPT TQQTTSV IPL IETNLHPLAR MPVPGSPTPG GVDKALNLAF NFNGTNFFIN 360
NATFTPPTVP VLLQILSGAQ TAQDLLPAGS VYPLPAHSTI EITLPATALA PGAPHPFHLH 420
GHAFAVVRSR GSTTYNYNDP IFRDVSSTGT PAAGDNVTIR FQTDNPGPWF LHCHIDPHLD 480
AGFAIVFAED VADVKAANPV PKAWSDLCP I YDGLSEANQ 519

SEQ ID NO: 62      moltype = AA length = 559
FEATURE          Location/Qualifiers
source          1..559
                mol_type = protein
                organism = Phanerochaete chrysosporium

SEQUENCE: 62
MLSLAVVSL AAATLAAPAA SDAPGWRFDL KPNLSGIVAL EAIVVNSSLV VIFDRATGDQ 60
PLKINGESTW GALWDLDTST VRPLSVLTDS FCASGALLSN GTMVMGGTP GGTGGDVAAP 120
PGNQAIRIFE PCASPSGDGC TLFEDPATVH LLEERWYPS VRIFDGSLMI IGGSHVLTFF 180

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YNVDPANSFE	FFPSKEQTPR	PSAFLERSLP	ANLFPRAFAL	PDGTVFIVAN	NQSIIYDIEK	240
NTETILPDIP	NGVRVTNPID	GSAILLPLSP	PDFIPEVLVC	GGSTADTSLP	STSLSSQHPA	300
TSQCSRIKLT	PEGIKAGWQV	EHMLEARMP	ELVHVPNGQI	LITNGAGTGF	AALSAVADPV	360
GNSNADHPVL	TPSLYTPDAP	LGKRISNAGM	PTTTIPRMYH	STVTLTQQGN	FFIGGNPNM	420
NFTPPGTPGI	KFPSELRIET	LDDPFMFRSR	PALLTMPEKL	KFGQKVTVPI	TIPSDLKASK	480
VQVALMDLGF	SSHAFHSSAR	LVMMESSISA	DRKSLTFTAP	PNGRVFPPGP	AVVFLTIDDV	540
TSPGERVMMG	SGNPPPTLE					559

SEQ ID NO: 63 moltype = AA length = 928
 FEATURE Location/Qualifiers
 source 1..928
 mol_type = protein
 organism = Myceliophthora thermophila

SEQUENCE: 63

MTNAACLSYC	ASKGFPPYAGT	EYSVECFCGT	TLASSSAKVA	DSECNMPCSG	APSEPCGAGS	60
RLSLFHSSAV	TGPAANPGVN	DFTHLGCYAE	GKTGRALTYN	PGLPGADMTV	AKCTAACRAA	120
NYILAGVEYG	GECYCGNTIA	NGGAPADSGC	SMVCNGNSTE	PCGGPDRLN	YSYKNQYEP	180
ATSTTGAGST	SSSSVPSATG	LPEGWSYQGC	WIDGKQGRIL	PYQLPDSQTN	SRAACANACA	240
EAGYTVSGTE	YAVQCFGDA	IHNCGVETDE	ADCSTPCPGA	PGEKCGAGDR	LSIVSRGPPK	300
IYAPPAPIEK	IGDWEYQCA	EDNINDKRFT	FWQIFNDIM	TPEMCLDRCA	EFGYHAAGLE	360
YGQECYCGDP	ANMATHGATF	RPSEECNVVC	AGNSTAICGG	LARLTTYFWI	GTPFYSWDFP	420
QDWRAGKYEF	LVDPGNIPLI	THEITGKVS	FISKGATGPG	NETGAYEFD	ATLEFRELHI	480
KTDVFCASV	TLDPKAGRQL	NVGGWAGEAT	YGRLYWPDG	APGVPGTHDW	QENVNVLHLQ	540
AGRWYPSVLV	LTNGSVMVVG	GLIGSNDAA	PSIEILPYTG	TPPLYMDWLD	RTHPNNLYPF	600
LCILPGGGIF	VQYWNARIL	DPVTFDVTKT	LPDAPGAPND	PKGGRTYPLE	GTAVLLPQKY	660
PYTDPLGLVI	CGSTEGPGN	ALDNCVSIYP	EADEPEWQIE	RMPSPRVMT	MAPLPDGTYL	720
IANGALHGVA	GFGLVGPNL	NALLYDPSKP	LGSRI TVAAN	TTIARMYHSE	AITLLDGRVL	780
ISGSNPEDGV	NPEEYRVEVF	LPPYLLAGKP	RPTFTLENRD	WAHQGTGIPF	TLGSPARNGD	840
ITATLLGSVA	STHGNSMGAR	TLMPRVSCR	TSCTVDAPPT	ANICPPGWYQ	FFVLDGGIPA	900
VGVYVRIGGD	AGQIGNWPA	PDFSVPGV				928

SEQ ID NO: 64 moltype = AA length = 558
 FEATURE Location/Qualifiers
 source 1..558
 mol_type = protein
 organism = Pycnoporus cinnabarinus

SEQUENCE: 64

MAPTAFSLVS	ALALASLSLA	APSAPGWSFD	LKKETSGIVA	LEAIVVSP	LVVFFDRASDD	60
PLQINNHSAA	GALWNLESST	VRPLDVLTN	FCASGALLSN	GTMASVGGDP	DGFVGNPAIR	120
PGNQAIRLFE	PCDSPTGDC	TLFEDPATLH	LLEKRWYPS	ARIPDGLII	VGMHEATPF	180
YNTDPALSFE	FFPRKEDTPR	PSEFLNRS	ANLFPRVFAL	PDGKVFVMA	NQSIIYDIEA	240
KTERILPDVP	NNVRVTNPM	GSAILLPLSP	PDFVPEVLVC	GGSTDTIDP	SLLTSQTPAS	300
SQCSRIRLDE	EGIAKGEWE	HMLEGRIMPE	LVHLPNGQVL	IANGGRTGFA	AIASVSEPVG	360
NSNADHPVLV	PSLYTPDAPL	GRRISNVGLP	SSGIPRLYHS	SVTLTPQGN	LIAGSNPNR	420
TTVGPGLKFP	SEFRVQTLDP	PFMSVERPKI	LNMPPKLGFN	KSFTVPISVP	SSLARPGAKV	480
QISLMDLGF	SHAFHSSARL	VFMGKISQD	SKSLTFTTP	NGRVYPPGPA	TVFLTIDDV	540
SEGAWMMGS	GNSPPTLE					558

SEQ ID NO: 65 moltype = AA length = 559
 FEATURE Location/Qualifiers
 source 1..559
 mol_type = protein
 organism = Pycnoporus cinnabarinus

SEQUENCE: 65

MFQTLHLHF	VLVVTGRGLA	APSTPTGWQF	NLKAERSGIV	ALESIVVSP	LVVFFDRATN	60
DPLQINNHS	WGALWNLETS	TVRALDVLTN	SFCASGALLS	NGTMASIGGD	PNGFPGNPAI	120
HPGTQAIRLF	EPCDSPTGEG	CTLFEDPVTL	HLLEKRWYPS	SVRIFDGSLL	IVGGMHEETP	180
FYNTDPALS	FEFFPKESTP	RPSEFLNRS	PANLFPRVFA	LPDGKVFVMA	NNQSIIYDIE	240
ANTERILPDI	PNNVRVTNPI	DGSAILLPLS	PPDFVPEVLV	CGGTQDTID	PSLLTSQTPA	300
SSQCSRIRLD	EEGIARGWEV	EHMLEGRMMP	ELVHLPNGQV	LIANGARTGF	AAIASVSDPV	360
GGSNADHAVL	VPSLYTPDAP	LGTRISNVGL	PSSGIARVYH	SSITLTPQGN	FLIAGSNPNM	420
NSSVTAGVKF	PSEFRVQTL	PPFMFVERPK	ILSMPKCLAF	GKSFTVPIAV	PSTLAHPGAK	480
VQVSLMDLGF	SSHAFHSSAR	LVMNAKISQ	DGKSLTFTTP	PNGRVYPPGP	ATIFLTIDDV	540
TSEGAWMMG	SGNPPPTLE					559

SEQ ID NO: 66 moltype = AA length = 656
 FEATURE Location/Qualifiers
 source 1..656
 mol_type = protein
 organism = synthetic construct

SEQUENCE: 66

MGHHHHHSS	GHEGRHMAS	APIGSAIPRN	NWAVTCD	SGNECNKAID	GNKDTFWHTF	60
YGANGPKPP	HTYTIDMKTT	QNVNGLSVLP	RQDGNQNGWI	GRHEVYLS	SDD GTNWGS	120
GSWFADSTTK	YSNFETRPAR	YVRLVAITEA	NGQPWTSIAE	INVQASSYT	APQPGLGRWG	180
PTIDLPIVPA	AAAIETSGR	VLMWSSYRND	AFEGSPGGIT	LTSWDPSTG	IVSDRTVTVT	240
KHDMFCPGIS	MDGNGQIVVT	GGNDAKKTSL	YDSSSDSWIP	GPDQVARGY	QSSATMSDGR	300

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VFTIGGSFSG	GVFEKNGEVY	SPSSKTWTSL	PNAKVNPLMT	ADKQGLYKSD	NHAWLFGWKK	360
GSVFQAGPST	AMNWWYTSGS	GDVKSAGKRQ	SNRGVAPDAM	SGNAVMYDAV	KGKILTFGGS	420
PDFEDSDATT	NAHIITLGEF	GTSPTNVFAS	NGLYFARTPH	TSVVLDPDGT	FITGGQRRGI	480
PFEDSTPVFT	PEIYVPEQDT	FYKQNPNSIV	RAYHSISLLL	PDGRVFNNGG	GLCGDCTTTH	540
FDAQIFTPNY	LYDSNGNLAT	RPKITRTSTQ	SVKVGGRITI	STDSSISKAS	LIRYGTATHT	600
VNTDQRRIPL	TLTNNNGNSY	SFQVPSDSGV	ALPGYWMLFV	MNSAGVPSVA	STIRVT	656

What is claimed is:

1. A method of preparing glucaric acid, comprising: contacting D-glucose and oxygen with a first catalyst composition comprising a first copper radical oxidase, a single electron oxidizer, and a small molecule activator under conditions suitable for formation of glucodialdose; and contacting glucodialdose and oxygen with a second catalyst composition comprising a second copper radical oxidase, a single electron oxidizer and a small molecule activator under conditions suitable for the formation of a product mixture comprising glucaric acid or salts thereof.
2. The method of claim 1, wherein the first copper radical oxidase comprises a galactose oxidase, a galactose oxidase mutant or combinations thereof.
3. The method of claim 1, wherein the first copper radical oxidase comprises any of SEQ ID Nos. 1 through 39.
4. The method of claim 1, wherein the first copper radical oxidase has from about 85% to about 100% sequence identity with any of SEQ ID Nos. 1 through 39.
5. The method of claim 1, wherein the second copper radical oxidase comprises a glyoxal oxidase, a glyoxal oxidase mutant or combinations thereof.
6. The method of claim 1, wherein the second copper radical oxidase comprises any of SEQ ID Nos. 62 through 65.
7. The method of claim 1, wherein the second copper radical oxidase has from about 85% to about 100% sequence identity with any of SEQ ID Nos. 62 through 65.
8. The method of claim 1, wherein the small molecule activator comprises L-tryptophan, 2-mercaptobenzothiazole, L-histidine, methylchloroisoethiazolinone, o-dianisidine, 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulfonic acid (ABTS), 4-aminoantipyrine, L-tyrosine, (2,2,6,6-tetramethylpiperidin-1-yl)oxyl, chloromethylisothiazolinone, 4-thiazolecarboxylic acid, Sunset yellow FCF, tartrazine, p-benzoquinone, dicoumarol, phthalimide, saccharin, phthalic anhydride, erythrosine B, 2-aminobenzothiazole, thiabendazole, 2-hydroxybenzothiazole, phenothiazine, 6-aminobenzothiazole, indigo carmine, naphthalimide, 2-aminothiazole, thiazole, 2H-1,4-benzothiazin-3 (4H)-one, 2-oxindole, betalaphachone, menaquinone, thiamine, 4-methyl-5-thiazoleethanol, Allura Red AC, menadione, p-cresol, Fast green FCF, Brilliant Blue FCF, methylisothiazolinone, caffeine, veratryl alcohol, fluorescein, or combinations thereof.
9. The method of claim 1, wherein the small molecule activator is present in an amount ranging from about 1 ppm to about 500 ppm.
10. The method of claim 1, wherein the single electron oxidizer comprises laccase, horseradish peroxidase, dyp-type peroxidase, lactoperoxidase, chloroperoxidase, manga-

nese peroxidase 1, ascorbate peroxidase, dye-decolorizing peroxidase, unspecific peroxygenase, dehaloperoxidase, catalase-peroxidase, lignin peroxidase, soybean seed coat peroxidase, isoforms thereof or combinations thereof.

11. The method of claim 1, wherein conditions suitable for formation of glucodialdose, conditions suitable for formation of a product mixture comprising glucaric acid or both comprise an oxygen pressure of equal to or less than about 500 psi.

12. The method of claim 1, further comprising a catalase.

13. The method of claim 1, wherein the catalase is defined by any SEQ ID No. 40 through SEQ ID No. 44.

14. The method of claim 1, further comprising introducing a caustic to the second catalyst composition, the product mixture or both.

15. The method of claim 1, wherein the glucodialdose, glucaric acid are formed at yields ranging from about 50% to about 99%.

16. The method of claim 1, wherein the glucaric acid or salts thereof have a purity of greater than about 70%.

17. The method of claim 1, wherein the product mixture comprising glucaric acid further comprises one or more sugar oxidation products.

18. The method of claim 17, wherein the one or more sugar oxidation products comprise aldonic acid, uronic acid, aldonic acid, a gluconic acid oxidation product, a gluconate, gluconic acid, glucuronic acid, glucose oxidation products, galactonic acid, galactaric acid, glutamic acid, a lactone of gluconic acid, a lactone of glucaric acid, a lactone of galactaric acid, a lactone of galactonic acid, glucodialdose, 2-ketoglucose, disaccharides, oxidized disaccharides, n-keto-acids, C2 to C6 diacids, salts thereof or combinations thereof.

19. A method comprising:

contacting a sugar with a catalyst composition comprising an oxidoreductase, a single electron oxidizer and a small molecule activator under conditions suitable for the formation of one or more oxidized sugar oxidation products comprising glucaric acid wherein the oxidoreductase comprises at least two copper radical oxidases, at least two mutated copper radical oxidases or combinations thereof.

20. The method of claim 19, wherein the at least two copper radical oxidases comprises any of SEQ ID Nos. 1 through 39, any of SEQ ID Nos. 62 through 65, have 85% to about 100% sequence identity with any of SEQ ID Nos. 1 through 39 or have 85% to about 100% sequence identity with any of SEQ ID Nos. 62 through 65.

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